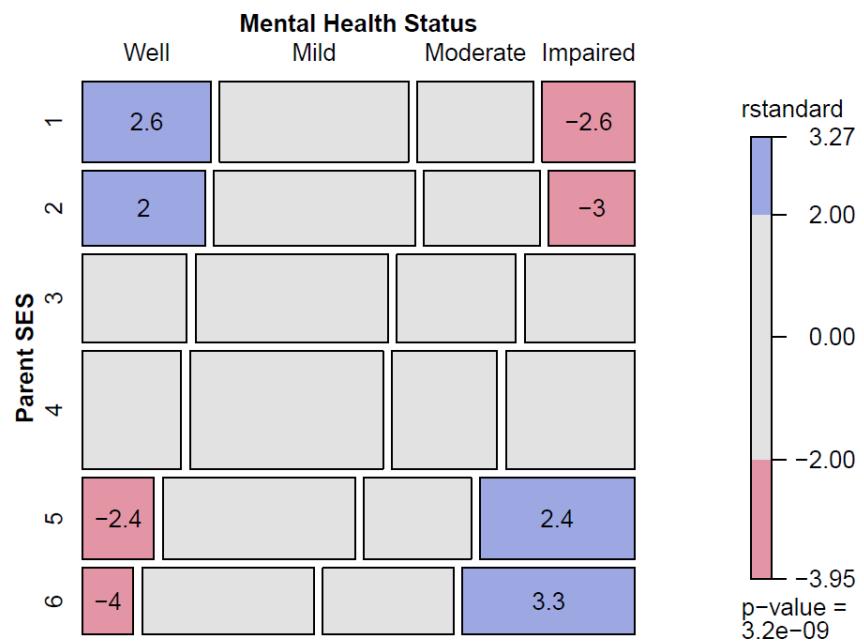


Visualizing Categorical Data with

Michael Friendly
York University

David Meyer
UAS Technikum Wien

January 13, 2015



Contents

Table of Contents	i
1 Introduction	1
1.1 Data visualization and categorical data: Overview	1
1.2 What is categorical data?	2
1.2.1 Case form vs. frequency form	3
1.2.2 Frequency data vs. count data	4
1.2.3 Univariate, bivariate, and multivariate data	4
1.2.4 Explanatory vs. Response variables	5
1.3 Strategies for categorical data analysis	5
1.3.1 Hypothesis testing approaches	6
1.3.2 Model building approaches	7
1.4 Graphical methods for categorical data	10
1.4.1 Goals and design principles for visual data display	11
1.4.2 Categorical data require different graphical methods	13
1.4.3 Effect ordering and rendering for data display	14
1.4.4 Interactive and dynamic graphics	19
1.4.5 Visualization = Graphing + Fitting + Graphing	19
1.4.6 The 80-20 rule	22
1.5 Chapter summary	24
1.6 Further reading	25
1.7 Lab exercises	25
2 Working with categorical data	27
2.1 Working with R data: vectors, matrices, arrays and data frames	27
2.1.1 Vectors	28
2.1.2 Matrices	29
2.1.3 Arrays	31
2.1.4 data frames	32
2.2 Forms of categorical data: case form, frequency form and table form	35
2.2.1 Case form	35
2.2.2 Frequency form	36
2.2.3 Table form	37
2.3 Ordered factors and reordered tables	39
2.4 Generating tables: table and xtabs	40
2.4.1 table()	40
2.4.2 xtabs()	42
2.5 Printing tables: structable and ftable	43
2.5.1 Text output	43
2.6 Subsetting data	44
2.6.1 Subsetting tables	44
2.6.2 Subsetting structables	45
2.6.3 Subsetting data frames	45
2.7 Collapsing tables	46
2.7.1 Collapsing over table factors	46
2.7.2 Collapsing table levels	48
2.8 Converting among frequency tables and data frames	49
2.8.1 Table form to frequency form	49
2.8.2 Case form to table form	51
2.8.3 Table form to case form	51
2.8.4 Publishing tables to L ^A T _E X or HTML	51
2.9 A complex example: TV viewing data	53
2.9.1 Creating data frames and arrays	54

2.10	2.9.2 Subsetting and collapsing	55
2.10	Lab exercises	56
3	Fitting and graphing discrete distributions	59
3.1	Introduction to discrete distributions	59
3.1.1	Binomial data	60
3.1.2	Poisson data	63
3.1.3	Type-token distributions	66
3.2	Characteristics of discrete distributions	68
3.2.1	The binomial distribution	69
3.2.2	The Poisson distribution	72
3.2.3	The negative binomial distribution	77
3.2.4	The geometric distribution	80
3.2.5	The logarithmic series distribution	82
3.2.6	Power series family	83
3.3	Fitting discrete distributions	83
3.3.1	R tools for discrete distributions	85
3.3.2	Plots of observed and fitted frequencies	89
3.4	Diagnosing discrete distributions: Ord plots	92
3.5	Poissonness plots and generalized distribution plots	96
3.5.1	Features of the Poissonness plot	96
3.5.2	Plot construction	96
3.5.3	The distplot function	97
3.5.4	Plots for other distributions	99
3.6	Fitting discrete distributions as generalized linear models	100
3.6.1	Covariates, overdispersion and excess zeros	103
3.7	Chapter summary	105
3.8	Lab exercises	106
4	Two-way contingency tables	111
4.1	Introduction	111
4.2	Tests of association for two-way tables	114
4.2.1	Notation and terminology	114
4.2.2	2 by 2 tables	115
4.2.3	Larger tables: Overall analysis	117
4.2.4	Tests for ordinal variables	118
4.2.5	Sample CMH Profiles	119
4.3	Stratified analysis	121
4.3.1	Assessing homogeneity of association	122
4.4	Fourfold display for 2×2 tables	123
4.4.1	Confidence rings for odds ratio	125
4.4.2	Stratified analysis for $2 \times 2 \times k$ tables	126
4.5	Sieve diagrams	131
4.5.1	Larger tables: The strucplot framework	134
4.6	Association plots	136
4.7	Observer agreement	138
4.7.1	Measuring agreement	139
4.7.2	Observer Agreement Chart	141
4.7.3	Observer bias in agreement	144
4.8	Trilinear plots	145
4.9	Chapter summary	148
4.10	Further reading	149
4.11	Lab exercises	149
5	Mosaic displays for n-way tables	153
5.1	Introduction	153
5.2	Two-way tables	154
5.3	The strucplot framework	158
5.3.1	Shading schemes	161
5.4	Three-way and larger tables	168
5.4.1	Fitting models	168
5.4.2	Sequential plots and models	172
5.4.3	Causal models	175
5.4.4	Partial association	177
5.5	Mosaic matrices for categorical data	184
5.5.1	Generalized mosaic matrices and pairs plots	190
5.6	3D mosaics	192

5.7	Visualizing the structure of loglinear models	194
5.7.1	Mutual independence	194
5.7.2	Joint independence	197
5.8	Chapter summary	198
5.9	Further reading	199
5.10	Lab exercises	199
6	Correspondence analysis	203
6.1	Introduction	203
6.2	Simple correspondence analysis	204
6.2.1	Notation and terminology	204
6.2.2	Geometric and statistical properties	205
6.2.3	R software for correspondence analysis	206
6.2.4	Corespondence analysis and mosaic displays	213
6.3	Multi-way tables: Stacking and other tricks	214
6.3.1	Interactive coding in R	215
6.3.2	Marginal tables and supplementary variables	219
6.4	Multiple correspondence analysis	222
6.4.1	Bivariate MCA	222
6.4.2	The Burt matrix	225
6.4.3	Multivariate MCA	225
6.5	Biplots for contingency tables	230
6.5.1	CA bilinear biplots	230
6.5.2	Biadditive biplots	233
6.6	Chapter summary	236
6.7	Lab exercises	236
7	Logistic Regression Models	239
7.1	Introduction	239
7.2	The logistic regression model	240
7.2.1	Fitting a logistic regression model	243
7.2.2	Model tests for simple logistic regression	245
7.2.3	Plotting a binary response	245
7.2.4	Grouped binomial data	248
7.3	Multiple logistic regression models	249
7.3.1	Conditional plots	252
7.3.2	Full-model plots	254
7.3.3	Effect plots	256
7.4	Case studies	258
7.4.1	More complex models: Model selection and visualization	269
7.5	Influence and diagnostic plots	278
7.5.1	Residuals and leverage	278
7.5.2	Influence diagnostics	279
7.5.3	Other diagnostic plots	288
7.6	Polytomous response models	296
7.6.1	Ordinal response	297
7.6.2	Visualizing results for the proportional odds model	303
7.6.3	Nested dichotomies	306
7.6.4	Generalized logit model	313
7.7	Chapter summary	318
7.8	Further reading	318
7.9	Lab exercises	318
8	Loglinear and Logit Models for Contingency Tables	323
8.1	Introduction	323
8.2	Loglinear models for frequencies	324
8.2.1	Loglinear models as ANOVA models for frequencies	324
8.2.2	Loglinear models for three-way tables	325
8.2.3	Loglinear models as GLMs for frequencies	326
8.3	Fitting and testing loglinear models	327
8.3.1	Model fitting functions	327
8.3.2	Goodness-of-fit tests	328
8.3.3	Residuals for loglinear models	330
8.3.4	Using <code>loglm()</code>	331
8.3.5	Using <code>glm()</code>	333
8.4	Equivalent logit models	336
8.5	Zero frequencies	341

8.6	Models for ordinal variables	345
8.6.1	Loglinear models for ordinal variables	346
8.6.2	Log-multiplicative (RC) models	351
8.7	Square tables	358
8.7.1	Quasi-independence, symmetry, quasi-symmetry and topological models	358
8.7.2	Ordinal square tables	365
8.8	Three-way and higher-dimensional tables	369
8.9	Multivariate responses	373
8.9.1	Bivariate, binary response models	374
8.9.2	More complex models	384
8.10	Chapter summary	395
8.11	Further reading	396
8.12	Lab exercises	396
9	Generalized linear models	399
9.1	Components of Generalized Linear Models	400
9.1.1	Variance functions	401
9.1.2	Hypothesis tests for coefficients	402
9.1.3	Goodness-of-fit tests	403
9.1.4	Comparing non-nested models	404
9.2	GLMs for count data	404
9.3	Models for overdispersed count data	412
9.3.1	The quasi-Poisson model	413
9.3.2	The negative-binomial model	414
9.3.3	Visualizing the mean-variance relation	415
9.3.4	Testing overdispersion	417
9.3.5	Visualizing goodness-of-fit	417
9.4	Models for excess zero counts	419
9.4.1	Zero-inflated models	419
9.4.2	Hurdle models	422
9.4.3	Visualizing zero counts	422
9.5	Case studies	423
9.5.1	Cod parasites	424
9.5.2	Demand for medical care by the elderly	434
9.6	Diagnostic plots for model checking	443
9.6.1	Diagnostic measures and residuals for GLMs	444
9.6.2	Quantile-quantile and half-normal plots	449
9.7	Multivariate response GLM models	452
9.8	Chapter summary	461
9.9	Further reading	462
9.10	Lab exercises	462
References	465	
Example Index	479	
Author Index	481	
Subject Index	483	

Preface

TODO: The preface has not yet been written. This is just a stub.

Audience

Overview

Acknowledgements

Chapter 1

Introduction

Categorical data consist of variables whose values comprise a set of discrete categories. Such data require different statistical and graphical methods than commonly used for quantitative data. The focus of this book is on visualization techniques and graphical methods designed to reveal patterns of relationships among categorical variables. This chapter outlines the basic orientation of the book and some key distinctions regarding the analysis and visualization of categorical data.

{ch:intro}

1.1 Data visualization and categorical data: Overview

{sec:viscat}

Beauty is truth; truth, beauty.
That is all ye know on Earth, all ye need to know.

John Keats, *Ode on a Grecian urn*

“Data visualization” can mean many things, from popular press infographics, to maps of voter turnout or party choice. Here we use this term in the narrower context of statistical analysis. As such, we refer to an approach to data analysis that focuses on *insightful* graphical display in the service of both *understanding* our data and *communicating* our results to others.

We may display the raw data, some summary statistics, or some indicators of the quality or adequacy of a fitted model. The word “insightful” suggests that the goal is (hopefully) to reveal some aspects of the data which might not be perceived, appreciated, or absorbed by other means. As in the quote from Keats, the overall aims include both beauty and truth, though each of these are only as perceived by the beholder.

Methods for visualizing quantitative data have a long history and are now widely used in both data analysis and in data presentation, and in both popular and scientific media. Graphical methods for categorical data, however, have only a more recent history, and are consequently not as widely used. The goal of this book is to show concretely how data visualization may be usefully applied to categorical data.

“Categorical” means different things in different contexts. We introduce the topic in Section 1.2 with some examples illustrating (a) types of categorical variables: binary, nominal, and ordinal, (b) data in case form vs. frequency form, (c) frequency data vs. count data, (d) univariate, bivariate, and multivariate data, and (e) the distinction between explanatory and response variables.

Statistical methods for the analysis of categorical data also fall into two quite different categories, described and illustrated in Section 1.3: (a) the simple randomization-based methods typi-

fied by the classical Pearson chi-squared (χ^2) test, Fisher's exact test, and Cochran-Mantel-Haenszel tests, and (b) the model-based methods represented by logistic regression, loglinear, and generalized linear models. In this book, Chapters 3–6 are mostly related to the randomization-based methods; Chapters 7–8 illustrate the model-based methods.

In Section 1.4 we describe some important similarities and differences between categorical data and quantitative data, and discuss the implications of these differences for visualization techniques. Section 1.4.5 outlines a strategy of data analysis focused on visualization.

In a few cases we show R code or results as illustrations here, but the fuller discussion of using R for categorical data analysis is postponed to Chapter 2.

1.2 What is categorical data?

{sec:whatis}

A **categorical variable** is one for which the possible measured or assigned values consist of a discrete set of categories, which may be *ordered* or *unordered*. Some typical examples are:

- Gender, with categories “Male”, “Female”.
- Marital status, with categories “Never married”, “Married”, “Separated”, “Divorced”, “Widowed”.
- Fielding position (in baseball), with categories “Pitcher”, “Catcher”, “1st base”, “2nd base”, . . . , “Left field”.
- Side effects (in a pharmacological study), with categories “None”, “Skin rash”, “Sleep disorder”, “Anxiety”, . . .
- Political attitude, with categories “Left”, “Center”, “Right”.
- Party preference (in Canada), with categories “NDP”, “Liberal”, “Conservative”, “Green”.
- Treatment outcome, with categories “no improvement”, “some improvement”, or “marked improvement”.
- Age, with categories “0-9”, “10-19”, “20-29”, “30-39”, . . .
- Number of children, with categories 0, 1, 2, . . .

As these examples suggest, categorical variables differ in the number of categories: we often distinguish **binary variables** (or **dichotomous variables**) such as Gender from those with more than two categories (called **polytomous variables**). For example, Table 1.1 gives data on 4,526 applicants to graduate departments at the University of California at Berkeley in 1973, classified by two binary variables, gender and admission status.

{tab:berk220}

Table 1.1: Admissions to Berkeley graduate programs

	Admitted	Rejected	Total
Males	1198	1493	2691
Females	557	1278	1835
Total	1755	2771	4526

Some categorical variables (Political attitude, Treatment outcome) may have ordered categories (and are called **ordinal**), while other (**nominal**) variables like Marital status have unordered categories.¹ For example, Table 1.2 shows a $2 \times 2 \times 3$ table of ordered outcomes (“none”, “some” or “marked” improvement) to an active treatment for rheumatoid arthritis compared to a placebo for men and women.

¹An ordinal variable may be defined as one whose categories are *unambiguously* ordered along a *single* underlying dimension. Both marital status and fielding position may be weakly ordered, but not on a single dimension, and not unambiguously.

{tab:arthritis} **Table 1.2:** Arthritis treatment data

Treatment	Sex	Improvement			Total
		None	Some	Marked	
Active	Female	6	5	16	27
	Male	7	2	5	14
Placebo	Female	19	7	6	32
	Male	10	0	1	11
Total		42	14	28	84

Finally, such variables differ in the fineness or level to which some underlying observation has been categorized for a particular purpose. From one point of view, *all* data may be considered categorical because the precision of measurement is necessarily finite, or an inherently continuous variable may be recorded only to limited precision.

But this view is not helpful for the applied researcher because it neglects the phrase “for a particular purpose”. Age, for example, might be treated as a quantitative variable in a study of native language vocabulary, or as an ordered categorical variable with decade groups (0-10, 11-20, 20-30, ...) in terms of the efficacy or side-effects of treatment for depression, or even as a binary variable (“child” vs. “adult”) in an analysis of survival following an epidemic or natural disaster. In the analysis of data using categorical methods, continuous variables are often recoded into ordered categories with a small set of categories for some purpose.²

1.2.1 Case form vs. frequency form

In many circumstances, data is recorded on each individual or experimental unit. Data in this form is called case data, or data in **case form**. The data in Table 1.2, for example, were derived from the individual data listed in the data set *Arthritis* from the *vcd* package. The following lines show the first five of $N = 84$ cases in the *Arthritis* data,

{sec:case-freq}

ID	Treatment	Sex	Age	Improved
1	57	Treated	Male	27
2	46	Treated	Male	29
3	77	Treated	Male	30
4	17	Treated	Male	32
5	36	Treated	Male	46

Whether or not the data variables, and the questions we ask, call for categorical or quantitative data analysis, when the data are in case form, we can always trace any observation back to its individual identifier or data record (for example, if the case with *ID* equal to 57 turns out to be unusual or noteworthy).

Data in **frequency form** has already been tabulated, by counting over the categories of the table variables. The same data shown as a table in Table 1.2 appear in frequency form as shown below.

Treatment	Sex	Improved	Freq
Placebo	Female	None	19
Treated	Female	None	6

²This may be a waste of information available in the original variable, and should be done for substantive reasons, not mere convenience. For example, some researchers unfamiliar with regression methods often perform a “median-split” on quantitative predictors so they can use ANOVA methods. Doing this precludes the possibility of determining if those variables have non-linear relations with the outcome.

3	Placebo	Male	None	10
4	Treated	Male	None	7
5	Placebo	Female	Some	7
6	Treated	Female	Some	5
7	Placebo	Male	Some	0
8	Treated	Male	Some	2
9	Placebo	Female	Marked	6
10	Treated	Female	Marked	16
11	Placebo	Male	Marked	1
12	Treated	Male	Marked	5

Data in frequency form may be analyzed by methods for quantitative data if there is a quantitative response variable (weighting each group by the cell frequency, with a weight variable). Otherwise, such data are generally best analyzed by methods for categorical data, where statistical models are often expressed as models for the frequency variable, in the form of an R formula like `Freq ~ ..`

In any case, an observation in a data set in frequency form refers to all cases in the cell collectively, and these cannot be identified individually. Data in case form can always be reduced to frequency form, but the reverse is rarely possible. In Chapter 2, we identify a third format, *table form*, which is the R representation of a table like Table 1.2.

1.2.2 Frequency data vs. count data

{sec:freq-count}

In many cases the observations representing the classifications of events (or variables) are recorded from *operationally independent* experimental units or individuals, typically a sample from some population. The tabulated data may be called **frequency data**. The data in Table 1.1 and Table 1.2 are both examples of frequency data because each tabulated observation comes from a different person.

However, if several events or variables are observed for the same units or individuals, those events are not operationally independent, and it is useful to use the term **count data** in this situation. These terms (following Lindsey (1995)) are by no means standard, but the distinction is often important, particularly in statistical models for categorical data.

For example, in a tabulation of the number of male children within families (Table 1.3, described in Section 1.2.3 below), the number of male children in a given family would be a *count* variable, taking values 0, 1, 2, The number of independent families with a given number of male children is a *frequency* variable. Count data also arise when we tabulate a sequence of events over time or under different circumstances in a number of individuals.

{tab:saxdata}

Table 1.3: Number of Males in 6115 Saxony Families of Size 12

Males	0	1	2	3	4	5	6	7	8	9	10	11	12
Families	3	24	104	286	670	1033	1343	1112	829	478	181	45	7

1.2.3 Univariate, bivariate, and multivariate data

{sec:uni-multi}

Another distinction concerns the number of variables: one, two or (potentially) many shown in a data set or table, or used in some analysis. Table 1.1 is an example of a bivariate (two-way) contingency table and Table 1.2 classifies the observations by three variables. Yet, we will see later that the Berkeley admissions data also recorded the department to which potential students applied (giving a three-way table), and in the arthritis data, the age of subjects was also recorded.

Any contingency table (in frequency or table form) therefore records the *marginal totals*, summed

over all variables not represented in the table. For data in case form, this means simply ignoring (or not recording) one or more variables; the “observations” remain the same. Data in frequency form, however, result in smaller tables when any variable is ignored; the “observations” are the cells of the contingency table. For example, in the *Arthritis* data, ignoring `Sex` gives the smaller 2×3 table for Treatment and Improved.

	Treatment	Improved	Freq
1	Placebo	None	29
2	Treated	None	13
3	Placebo	Some	7
4	Treated	Some	7
5	Placebo	Marked	7
6	Treated	Marked	21

In the limiting case, only one table variable may be recorded or available, giving the categorical equivalent of univariate data. For example, Table 1.3 gives data on the distribution of the number of male children in families with 12 children (discussed further in Example 3.2). These data were part of a large tabulation of the sex distribution of families in Saxony in the 19th century, but the data in Table 1.3 have only one discrete classification variable, number of males. Without further information, the only statistical questions concern the form of the distribution. We discuss methods for fitting and graphing such discrete distributions in Chapter 3. The remaining chapters relate to bivariate and multivariate data.

1.2.4 Explanatory vs. Response variables

Most statistical models make a distinction between **response variables** (or *dependent*, or *criterion* variables) and **explanatory variables** (or *independent*, or *predictor* variables).

{sec:exp-resp}

In the standard (classical) linear models for regression and analysis of variance (ANOVA), for instance, we treat one (or more) variables as responses, to be explained by the other, explanatory variables. The explanatory variables may be quantitative or categorical (e.g., factors in R). This affects only the details of how the model is specified or how coefficients are interpreted for `lm()` or `glm()`. In these classical models, the response variable (“treatment outcome”, for example), must be considered quantitative, and the model attempts to describe how the *mean* of the distribution of responses changes with the values or levels of the explanatory variables, such as age or gender.

When the response variable is categorical, however, the standard linear models do not apply, because they assume a normal (Gaussian) distribution for the model residuals. For example, in Table 1.2 the response variable is `Improvement`, and even if numerical scores were assigned to the categories “none”, “some”, “marked”, it may be unlikely that the assumptions of the classical linear models could be met.

Hence, a categorical *response* variable generally requires analysis using methods for categorical data, but categorical *explanatory* variables may be readily handled by either method.

The distinction between response and explanatory variables also becomes important in the use of loglinear models for frequency tables (described in Chapter 8), where models can be specified in a simpler way (as equivalent logit models) by focusing on the response variable.

1.3 Strategies for categorical data analysis

{sec:strategies}

Data analysis typically begins with exploratory and graphical methods designed to expose features of the data, followed by statistical analysis designed to summarize results, answer questions and draw conclusions. Statistical methods for the analysis of categorical data can be classified into two broad categories: those concerned with *hypothesis testing* per se versus those concerned with *model building*.

Done: DM: What about explorative analysis? MF: edited above

{sec:strategies-hyp}

1.3.1 Hypothesis testing approaches

In many studies, the questions of substantive interest translate readily into questions concerning hypotheses about **association** between variables, a more general idea than that of correlation (*linear* association) for quantitative variables. If a non-zero association exists, we may wish to characterize the strength of the association numerically and understand the pattern or nature of the association.

For example, in Table 1.1, a main question is: “Is there evidence of gender-bias in admission to graduate school?” Another way to frame this: “Are males more likely to be admitted?” These questions can be expressed in terms of an association between gender and admission status in a 2×2 contingency table of applicants classified by these two variables. If there is evidence for an association, we can assess its strength by a variety of measures, including the difference in proportions admitted for men and women or the ratio of the odds of admission for men compared to women, as described in Section 4.2.2.

Similarly, in Table 1.2, questions about the efficacy of the treatment for rheumatoid arthritis can be answered in terms of hypotheses about the associations among the table variables: Treatment, Sex, and the Improvement categories. Although the main concern might be focused on the overall association between Treatment and Improvement, one would also wish to know if this association is the same for men and women. A **stratified analysis** (Section 4.3) controls for the effects of background variables like Sex, and tests for **homogeneity of association** helping to determine if these associations are equal.

Questions involving tests of such hypotheses are answered most easily using a large variety of specific statistical tests, often based on randomization arguments. These include the familiar Pearson chi-squared test for two-way tables, the Cochran-Mantel-Haenszel test statistics, Fisher’s exact test, and a wide range of measures of strength of association. These tests make minimal assumptions, principally requiring that subjects or experimental units have been randomly assigned to the categories of experimental factors. The hypothesis testing approach is illustrated in Chapter 4–6, though the emphasis is on graphical methods which help to understand the nature of association between variables.

{ex:haireye0}

EXAMPLE 1.1: Hair color and eye color

The data set *HairEye* below records data on the relationship between hair color and eye color in a sample of nearly 600 students.

	Eye			
Hair	Brown	Blue	Hazel	Green
Black	68	20	15	5
Brown	119	84	54	29
Red	26	17	14	14
Blond	7	94	10	16

The standard analysis (with `chisq.test()` or `assocstats()`) gives a Pearson χ^2 of 138.3 with nine degrees of freedom, indicating substantial departure from independence. Among the measures of strength of association, the **phi coefficient**, $\phi = \sqrt{\chi^2/N} = 0.483$, and **Cramer’s V**, $V = \sqrt{\chi^2/N \min(r - 1, c - 1)} = 0.279$ indicate a substantial relationship between hair and eye color.³

	X^2	df	P(> X^2)
Likelihood Ratio	146.44	9	0
Pearson	138.29	9	0
 Phi-Coefficient : 0.483			
Contingency Coeff.:	0.435		
Cramer's V :	0.279		

³Cramer’s V varies from 0 (no association) to 1 (perfect association).

The further (and perhaps more interesting question) is how do we understand the *nature* of this association between hair and eye color? Two graphical methods related to the hypothesis testing approach are shown in Figure 1.1.

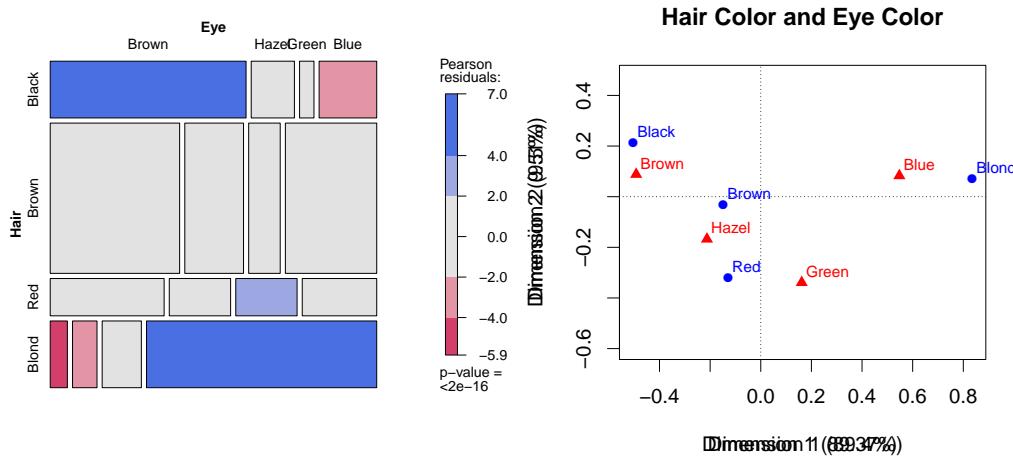


Figure 1.1: Graphical displays for the hair color and eye color data. Left: mosaic display; right: correspondence analysis plot

The left panel of Figure 1.1 is a **mosaic display** (Chapter 5), constructed so that the size of each rectangle is proportional to the observed cell frequency. The shading reflects the cell contribution to the χ^2 statistic—shades of blue when the observed frequency is substantially greater than the expected frequency under independence, shades of red when the observed frequency is substantially less, as shown in the legend.

The right panel of this figure shows the results of a correspondence analysis (Chapter 6), where the deviations of the hair color and eye color points from the origin accounts for as much of the χ^2 as possible in two dimensions.

We observe that both the hair colors and the eye colors are ordered from dark to light in the mosaic display and along Dimension 1 in the correspondence analysis plot. The deviations between observed and expected frequencies have an opposite-corner pattern in the mosaic display, except for the combination of red hair and green eyes, which also stand out as the largest values on Dimension 2 in the Correspondence analysis plot. Displays such as these provide a means to understand *how* the variables are related. \triangle

1.3.2 Model building approaches

Model-based methods provide tests of equivalent hypotheses about associations, but offer additional advantages (at the cost of additional assumptions) not provided by the simpler hypotheses-testing approaches. Among these advantages, model-based methods provide estimates, standard errors and confidence intervals for parameters, and the ability to obtain predicted (fitted) values with associated measures of precision.

We illustrate this approach here for a dichotomous response variable, where it is often convenient to construct a model relating a function of the probability, π , of one event to a linear combination of the explanatory variables. Logistic regression uses the **logit function**,

$$\text{logit}(\pi) \equiv \log_e \left(\frac{\pi}{1 - \pi} \right)$$

which may be interpreted as the *log odds* of the given event. A linear logistic model can then be expressed as

$$\text{logit}(\pi) = \beta_0 + \beta_1 x_1 + \beta_2 x_2 + \dots$$

Statistical inferences from model-based methods provide tests of hypotheses for the effects of the predictors, x_1, x_2, \dots , but they also provide estimates of parameters in the model, β_1, β_2, \dots and associated confidence intervals. Standard modeling tools allow us to graphically display the fitted response surface (with confidence or prediction intervals) and even to extrapolate these predictions beyond the given data. A particular advantage of the logit representation in the logistic regression model is that estimates of odds ratios (Section 4.2.2.1) may be obtained directly from the parameter estimates.

{ex:nasa0}

EXAMPLE 1.2: Space shuttle disaster

To illustrate the model-based approach, the graph in Figure 1.2 is based on a logistic regression model predicting the probability of a failure in one of the O-ring seals used in the 24 NASA space shuttles prior to the disastrous launch of the *Challenger* in January, 1986. The explanatory variable is the ambient temperature at the time of the flight. The sad story behind these data, and the lessons to be learned for graphical data display are related in Example 1.10.

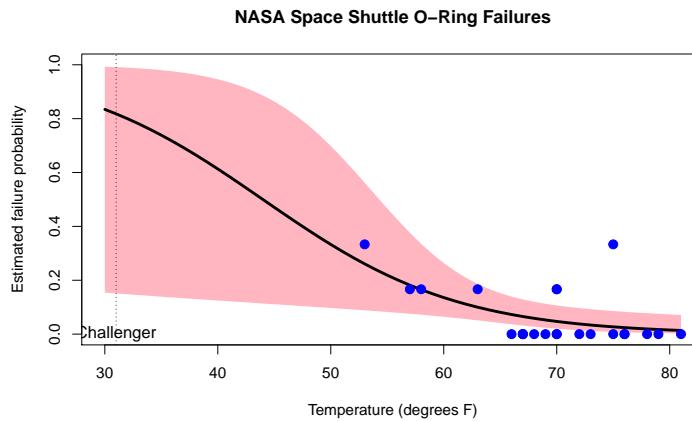


Figure 1.2: Space shuttle O-ring failure, observed and predicted probabilities. The dotted vertical line at 31° shows the prediction for the launch of the *Challenger*.
Fig:spaceshuttle0

Here, we simply note that the fitted model, shown by the solid line in Figure 1.2, corresponds to the prediction equation (with standard errors shown in parentheses),

$$\text{logit}(\text{Failure}) = 5.09 - 0.116 \text{ Temperature}_{(3.06) \quad (0.047)}$$

A hypothesis test that failure probability is unassociated with temperature is equivalent to the test that the coefficient for temperature in this model equals 0; this test has a p -value of 0.014, convincing evidence for rejection.

The parameter estimate for temperature, -0.116 , however, gives more information. Each 1° increase in temperature decreases the log odds of failure by 0.116 , with 95% confidence interval $[-0.208, -0.0235]$. The equivalent odds ratio is $\exp(-0.116) = 0.891$ [0.812, 0.977]. Equivalently, a 10° decrease in temperature corresponds to an odds ratio of a failure of $\exp(10 \times 0.116) = 3.18$, more than tripling the odds of a failure.

When the *Challenger* was launched, the temperature was only 31° . The shaded region in Figure 1.2 show 95% prediction intervals for failure probability. All previous shuttles (shown by the

points in the figure) had been launched at much warmer temperatures, so the prediction interval (the dashed vertical line) at 31° represents a considerable extrapolation beyond the available data. Nonetheless, the model building approach does provide such predictions along with measures of their uncertainty. Figure 1.2 is a graph that might have saved lives.



TODO: DM: Remove Donner party example to shorten up the intro? One example for logistic regression seems enough here.

{ex:donner0}

EXAMPLE 1.3: Donner Party

In April–May of 1846 (three years before the California gold rush), the Donner and Reed families set out for California from the American mid-west in a wagon train to seek a new life and perhaps their fortune in the new American frontier. By mid July, a large group had reached a site in present-day Wyoming; George Donner was elected to lead what was to be called the “Donner Party,” which eventually numbered 87 people in 23 wagons, along with their oxen, cattle, horses, and worldly possessions.

They were determined to reach California as quickly as possible. Lansford Hastings, a self-proclaimed trailblazer (retrospectively, of dubious distinction), proposed that the party follow him through a shorter path through the Wasatch Mountains. Their choice of “Hastings’s Cutoff” proved disastrous: Hastings had never actually crossed that route himself, and the winter of 1846 was to be one of the worst on record.

In October, 1846, heavy snow stranded them in the eastern Sierra Nevada, just to the east of a pass which bears their name today. The party made numerous attempts to seek rescue, most turned back by blizzard conditions. Relief parties in March–April 1847 rescued 40, but discovered grizzly evidence that those who survived had cannibalized those who died.

Here we briefly examine of how statistical models and graphical evidence can shed light on the question of who survived in the Donner party.

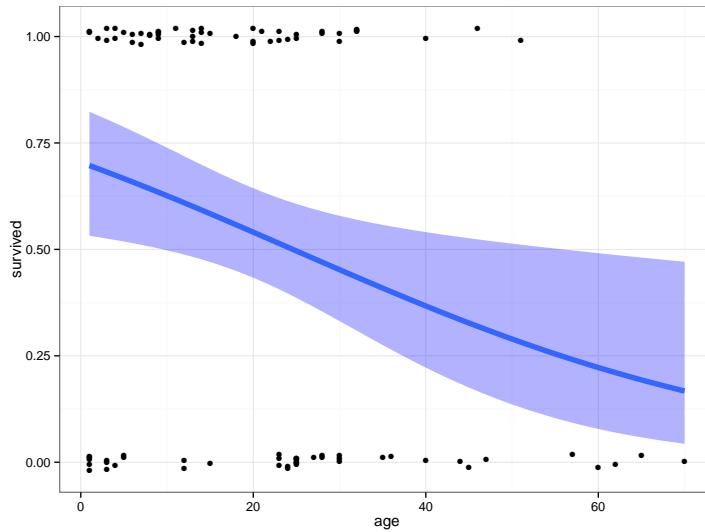


Figure 1.3: Donner party data, showing the relationship between age and survival. The blue curve and confidence band give the predicted probability of survival from a linear logistic regression model.
{fig:donner0}

Figure 1.3 is an example of what we call a *data-centric, model-based* graph of a discrete (binary)

outcome: lived (1) versus died (0). That is, it shows both the data and a statistical summary based on a fitted statistical model. The statistical model provides a smoothing of the discrete data.

The jittered points at the top and bottom of the graph show survival in relation to age of the person. You can see that there were more people who survived among the young, and more who died among the old. The blue curve in the plot shows the fitted probability of survival from a linear logistic regression model for these data with a 95% confidence band for that predictions. The prediction equation for this model can be given as:

$$\text{logit}(\text{survived}) = 0.868 - 0.0353 \text{ age}$$

(0.372) (0.015)

It implies that the log odds of survival decreases by 0.0352 with each additional year of age or by $10 \times 0.0352 = 0.352$ for an additional decade. Another way to say this is that the odds of survival is multiplied by $\exp(0.353) = .702$ with each 10 years of age, a 30% decrease.

Of course, these visual and statistical summary depends on the validity of fitted model. For contrast, Figure 1.4 shows two other model-based smoothers that relax the assumption of the linear logistic regression model. The left panel shows the result of fitting a semi-parametric model with a natural cubic spline with one more degree of freedom than the linear logistic model. The right panel shows the fitted curve for a non-parametric, loess model. Both of these hint that the relationship of survival to age is more complex than what is captured in the linear logistic regression model. We return to these data in Chapter 7.

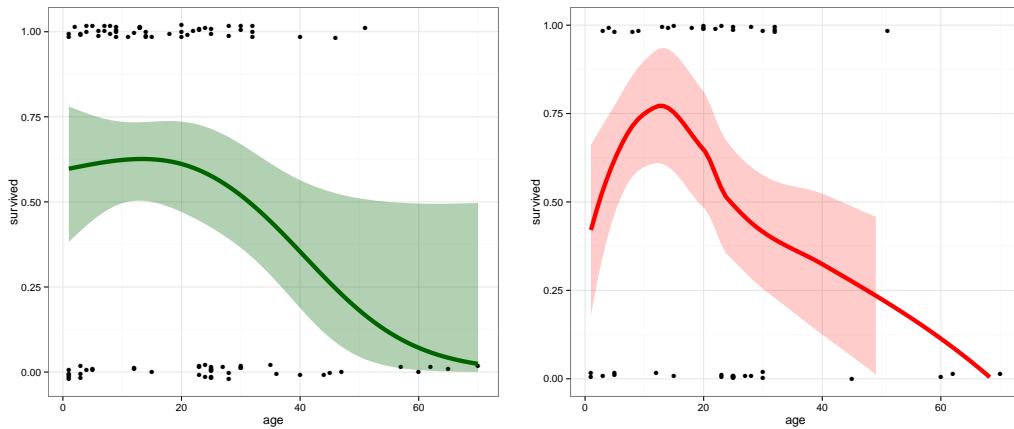


Figure 1.4: Donner party data, showing other model-based smoothers for the relationship between age and survival. Left: using a natural spline; right: using a non-parametric loess smoother.
fig:donner0-other



1.4 Graphical methods for categorical data

{sec:methods}

You can see a lot, just by looking

Yogi Berra

The graphical methods for categorical data described in this book are in some cases straightforward adaptations of more familiar visualization techniques developed for quantitative data. Graphical principles and strategies, and the relations between the visualization approach and traditional

statistical methods are described in a number of sources, including Chambers *et al.* (1983), Cleveland (1993b) and several influential books by Tufte (Tufte, 1983, 1990, 1997, 2006).

The fundamental ideas of statistical graphics as a comprehensive system of visual signs and symbols with a grammar and semantics was first proposed in Jacques Bertin's *Semiology of Graphics* (1983). These ideas were later extended to a computational theory in Wilkinson's *Grammar of Graphics* (2005), and implemented in R in Hadley Wickham's *ggplot2* package (Wickham, 2009, Wickham and Chang, 2013).

Another perspective on visual data display is presented in Section 1.4.1 focusing on the communication goals of statistical graphics. However, the discrete nature of categorical data implies that some familiar graphic methods need to be adapted, while in other cases we require a new graphic metaphor for data display. These issues are illustrated in Section 1.4.2. Section 1.4.3 discusses the principle of effect ordering for categorical variables in graphs and tables.

1.4.1 Goals and design principles for visual data display

Designing good graphics is surely an art, but as surely, it is one that ought to be informed by science. In constructing a graph, quantitative and qualitative information is encoded by visual features, such as position, size, texture, symbols and color. This translation is reversed when a person studies a graph. The representation of numerical magnitude and categorical grouping, and the apperception of patterns and their *meaning* must be extracted from the visual display.

{sec:intro-goals}

There are many views of graphs, of graphical perception, and of the roles of data visualization in discovering and communicating information. On the one hand, one may regard a graphical display as a *stimulus*—a package of information to be conveyed to an idealized observer. From this perspective certain questions are of interest: which form or graphic aspect promotes greater accuracy or speed of judgment (for a particular task or question)? What aspects lead to greatest memorability or impact? Cleveland (Cleveland and McGill, 1984, 1985, Cleveland, 1993a), Spence and Lewandowsky (Lewandowsky and Spence, 1989, Spence, 1990, Spence and Lewandowsky, 1990) have made important contributions to our understanding of these aspects of graphical display.

An alternative view regards a graphical display as an act of *communication*—like a narrative, or even a poetic text or work of art. This perspective places the greatest emphasis on the desired communication goal, and judges the effectiveness of a graphical display in how well that goal is achieved (Friendly and Kwan, 2011). Kosslyn (1985, 1989) and Tufte (1983, 1990, 1997) have articulated this perspective most clearly.

In this view, an effective graphical display, like good writing, requires an understanding of its *purpose*—what aspects of the data are to be communicated to the viewer. In writing we communicate most effectively when we know our audience and tailor the message appropriately. So too, we may construct a graph in different ways to: (a) use ourselves, (b) present at a conference or meeting of our colleagues, (c) publish in a research report, or (d) communicate to a general audience (Friendly (1991, Ch. 1), Friendly and Kwan (2011)). Figure 1.5 illustrates a basic contrast between graphs for presentation purposes, designed to appeal persuasively to a large audience (one-to-many) and the use of perhaps many graphs we might make for ourselves for exploratory data analysis (many-to-one).

Figure 1.6 shows one organization of visualization methods in terms of the *primary* use or intended communication goal, the functional *presentation goal*, and suggested corresponding *design principles*.

We illustrate these ideas and distinction in the examples below, most of which are treated again in later chapters.

{ex:arrests0}

EXAMPLE 1.4: Racial profiling: Arrests for marijuana possession

In a case study that will be examined in detail in Chapter 7 (Example 7.10), the *Toronto Star* newspaper studied a huge data base of arrest records by Toronto police for indications of possible

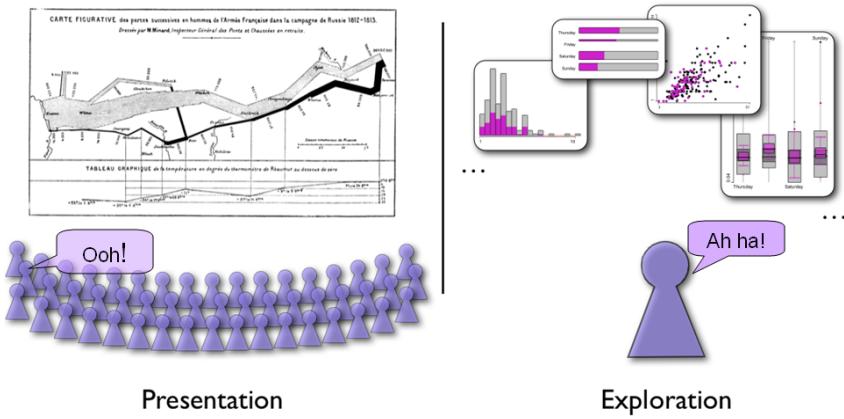


Figure 1.5: Different communication purposes require different graphs. For presentations, a single, carefully crafted graph may appeal best to a large audience; for exploratory analysis, many related images from different perspectives for a narrow audience (often you!).
Source: Adapted from a blog entry by Martin Theus, <http://www.theusrus.de/blog/presentation-vs-exploration/>.

{fig:presentation-exploration}

racial profiling, i.e., differential treatment of those arrested on the basis of skin color. They focused on the charge of simple possession of a small amount of marijuana, for which enforcement procedures allowed police discretion. An officer could release an arrestee with a summons (“Form 9”) to appear in court, or take the person to a police station for questioning (“Form 10”) or booking (“Form 11.1”) or order the person held in jail for a bail hearing (“Show cause”).

TODO: DM: Add frequency table here? Could be easier compared with mosaic plot. MF: No, not required for this example

The statistical issue was whether the data on these arrests showed evidence of differential treatment in relation to skin color, particularly in the treatment of blacks vs. whites, controlling, of course, for other factors. Statistical tests on these data (χ^2 tests, loglinear models, logistic regression) showed overwhelming evidence of differential treatment of blacks and whites. However, tables of these results do not reveal the nature of this association.

Figure 1.7 is an example of a graph designed for *analysis*—a mosaic display (Chapter 5) showing the frequencies of those arrested on this charge by skin color and release type. The size of each rectangle shows the frequency and these are shaded in relation to the association between skin color and release—blue for positive associations (more than expected under independence) to red for negative associations.

Once you know how to read such graphs, the pattern is clear: blacks were indeed more likely to be held for more severe treatment, whites were more likely to be released with a summons. But this is hardly a graph that would be clear to a general audience, and would require a good deal of explanation.

In contrast, Figure 1.8 shows a redesign of this as a *presentation graphic* prepared by the *Star* and published on December 11, 2002 in conjunction with a meeting between the newspaper and the Toronto Police Services Board to consider the issue of racial profiling. The police vehemently denied that racial profiling was taking place. The revision makes the point immediately obvious and compelling in the following ways:

- It announces the conclusion in the figure title: “Same charge, different treatment”
- The text box at the top provides the context for this conclusion
- Skin colors “Brown” and “Other”, which were of low frequency were removed, and the release categories “Form 10” and “Form 11.1” were combined as “released at station.”

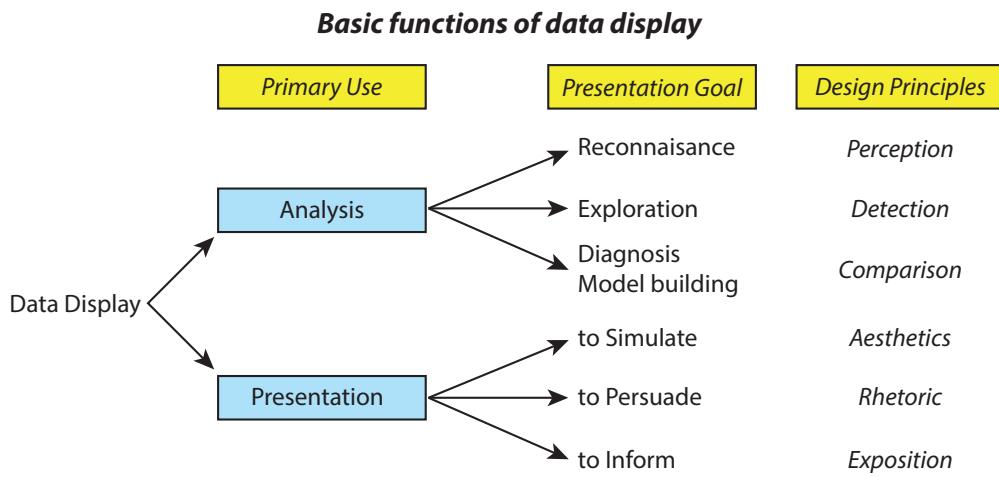


Figure 1.6: A taxonomy of the basic functions of data display by intended use, presentation goal and design principles.

{fig:datadisp}

- The graphic is still a mosaic display, however, it now shows explicitly the number of charges laid against whites and blacks and the percentage of each treatment.
- The labels for Whites and Blacks were enhanced by indicating what a reader should see for each.
- The legend for color is titled non-technically as “degree of likelihood.”

Clear communication is not achieved without effort. The revised graph required several iterations and emails between the graphic designer and the statistical consultant (the first author of this book) in the few hours available before the newspaper went to press. The main question was, “what are we trying to show here?” Starting with the original Figure 1.7 mosaic, we asked “what can we remove?” and “what can we add?” to make the message clearer.



1.4.2 Categorical data require different graphical methods

{sec:intro-catdata}

We mentioned earlier, and will see in greater detail in Chapter 7 and Chapter 8 that statistical models for discrete response data and for frequency data are close analogs of the linear regression and ANOVA models used for quantitative data. These analogies suggest that the graphical methods commonly used for quantitative data may be adapted directly to categorical data.

Happily, it turns out that many of the analysis graphs and diagnostic displays (e.g., effect plots, influence plots, added variable and partial residual plots, etc.) that have become common adjuncts in the analysis of quantitative data have been extended to generalized linear models including logistic regression (Section 7.5) and loglinear models (Section 9.6).

Unhappily, the familiar techniques for displaying raw data are often disappointing when applied to categorical data. The simple scatterplot, for example, widely used to show the relation between quantitative response and predictors, when applied to discrete variables, gives a display of the category combinations, with all identical values overplotted, and no representation of their frequency.

Instead, frequencies of categorical variables are often best represented graphically using *areas* rather than as position along a scale. Friendly (1995) describes conceptual and statistical models that give a rationale for this graphic representation. Figure 1.7 does this in the form of a modified bar chart (mosaic plot), where the widths of the horizontal bars show the proportions of whites

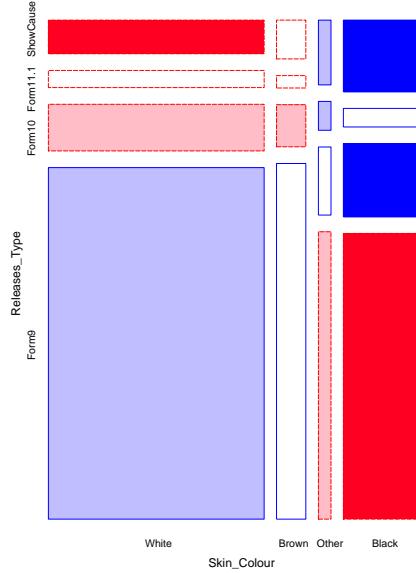


Figure 1.7: Mosaic display showing the relationship between skin color and release type for those arrested on a charge of simple possession of marijuana in Toronto, 1996-2002.

and blacks in the data, and the divisions of each group give the percents of each release type. Consequently, the areas of each bar are proportional to the frequency in the cells of this 2×3 table.

As we describe later in this book, using the visual attribute

$$\text{area} \sim \text{frequency}$$

also allows creating novel graphical displays of frequency data for special circumstances.

Figure 1.9 shows two examples. The left panel gives a *fourfold display* of the frequencies of admission and gender in the Berkeley data shown in Table 1.1. What should be seen at a glance is that males are more often admitted and females more often rejected (shaded blue); see Section 4.4 for details.

The right panel shows another specialized display, an *agreement chart* designed to show the strength of agreement in a square table for two raters (see Section 4.7.2). The example here (Example 4.17) concerns agreement of ratings of breast cancer from mammograms by two raters. The dark squares along the diagonal show exact agreement; the lighter diagonal rectangles allow 1-off agreement, and both are shown in relation to chance agreement (diagonal enclosing rectangles). What should be seen at a glance is that exact agreement is moderately strong and extremely strong if you allow the raters to differ by one rating category.

1.4.3 Effect ordering and rendering for data display

In plots of quantitative variables, standard methods (histograms, scatterplots) automatically position values along ordered scales, facilitating comparison (“which is less/more?”) and detection of patterns, trends and anomalies. However, by its nature, categorical data involves discrete variables such as education level, hair color, geographic region (state or province) or preference for a political party. With alphabetic labels for ordered categories (e.g., education: Low, Medium, High), it

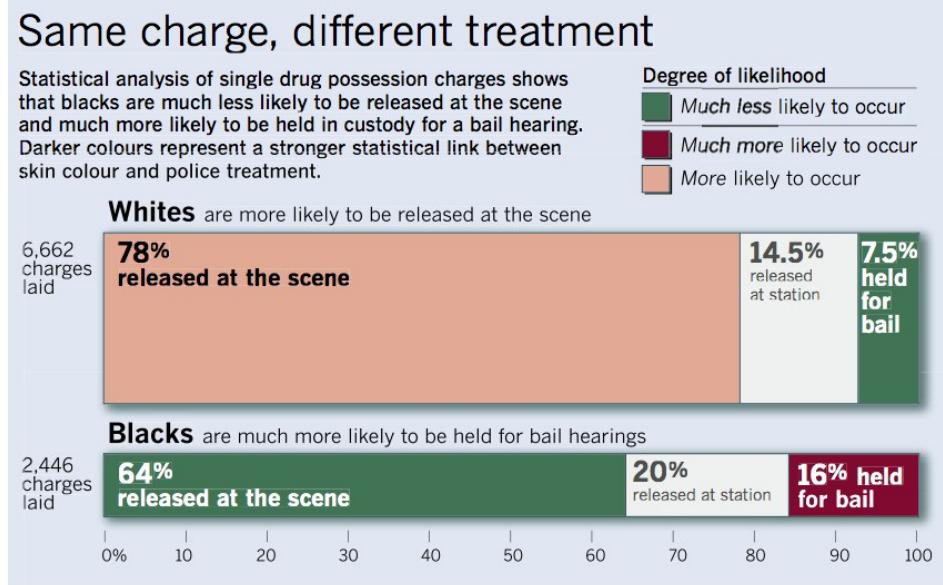


Figure 1.8: Redesign of Figure 1.7 as a presentation graphic. *Source:* Graphics department, *The Toronto Star*, December 11, 2002. Used by permission.

{fig:arrests0-star}

is unfortunately all too easy to end up with a nonsensical display with the categories ordered High, Low, Medium. Geographic regions (U.S. states) are often ordered alphabetically by default as are the names of political parties and other categorical variables. This may be useful for lookup, but for the purposes of comparison and detection, this is almost always a bad idea.

Instead, Friendly and Kwan (2003) proposed the principle of *effect-order sorting* for visual displays (tables as well as graphs):

sort the data by the effects to be seen to facilitate comparison

For quantitative data, this is often achieved by sorting the data according to means or medians of row and column factors, called *main-effect ordering*. For categorical data, graphs and tables are often most effective when the categories are arranged in an order reflecting their association, called *association ordering*.

Another important principle concerns the *rendering* of visual attributes of elements in graphical displays (Friendly, 2002). For example, categorical variables in plots (and tables) can be distinguished by any one or more of color, size, shape, or font. The examples below show the use of color to illustrate the precept:

render the data by the effects to be seen to facilitate detection

{ex:glass}

EXAMPLE 1.5: British social mobility

Bishop *et al.* (1975, p. 100) analyzed data on the occupations of 3500 British fathers and their sons from a study by Glass (1954), with five occupational categories: Professional, Managerial, Supervisory, Skilled manual and Unskilled manual.

One would expect, of course, a strong association between a son's occupation and that of his father—the apple doesn't fall very far from the tree. Mosaic plots (detailed in Chapter 5) provide a natural way to show such relationships. Figure 1.10 shows two such plots. The left panel shows the result obtained when the table variables `father` and `son` are read as factors, and therefore ordered

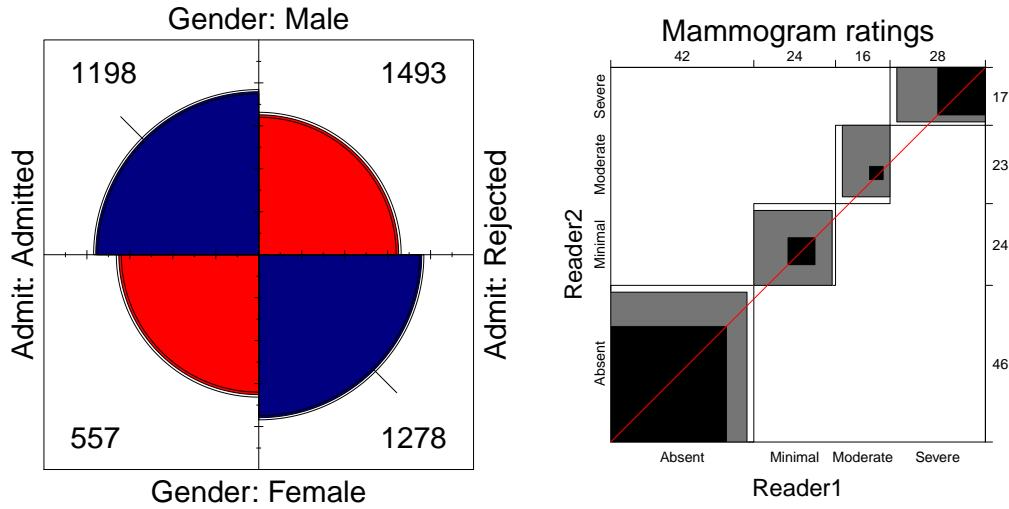


Figure 1.9: Frequencies of categorical variables shown as areas. Left: fourfold display of the relation between gender and admission in the Berkeley data; right: agreement plot for two raters assessing mammograms.

{fig:area-diagrams}

alphabetically by default. It is difficult to see any overall pattern, except for the large values in the diagonal cells (shaded blue) corresponding to equal occupational status.

In the right panel, the categories have been arranged in decreasing order of occupational status to show the association according to status. Now you can see a global pattern of shading color, where the tiles become increasingly red as one moves away from the main diagonal, reflecting a greater difference between the occupation of the father and son. The interpretation here is that most sons remain in their father's occupational class, but when they differ, there is little mobility across large steps.

In this example, `father` and `son` are clearly ordinal variables and should be treated as such in both graphs and statistical models. Correspondence analysis (Chapter 6) provides a natural way to depict association by assigning scores to the categories to optimally represent their relationships. Loglinear models provide special methods for ordinal variables (Section 8.6) and square frequency tables (Section 8.7).

△

The ideas of effect ordering and rendering with color shading to enhance perception can also be used in tabular displays, as illustrated in the next example.

{ex:barley}

EXAMPLE 1.6: Barley data

The classic `barley` dataset (in `lattice`) from Immer *et al.* (1934) gives a $10 \times 2 \times 6$ table of yields of 10 varieties of barley in two years (1931, 1932) planted at 6 different sites in Minnesota. Cleveland (1993b) and many others have used this data to illustrate graphical methods, and one surprising finding not revealed in standard tabular displays is that the data for one site (Morris) may have had the values for 1931 and 1932 switched.⁴

To focus attention on this suspicious effect in a tabular display, you can calculate the `yield`

⁴This canonical story, like many others in statistics and graphics lore turns out to be apocryphal on closer examination. Wright (2013) recently took a closer look at the original data and gives an expanded data set as `minnesota.barley.yield` in the `agridat` package. With a wider range of years (1927–1936), other local effects like weather had a greater impact than the overall year effects seen in 1931–1932, and the results for the Morris site no longer stand out as surprising.

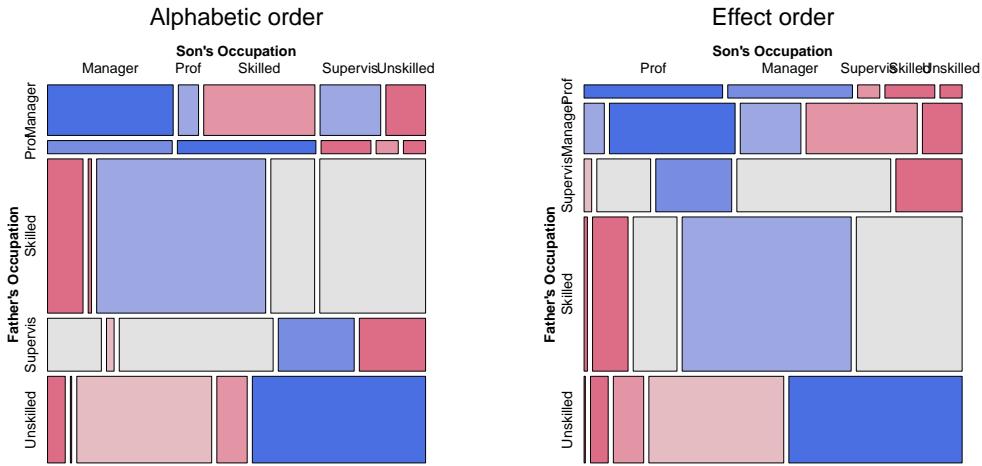


Figure 1.10: Mosaic plots for Glass' mobility table of occupational status. In these displays the area of each tile is proportional to frequency and shading color shows the departure from independence, using blue for positive, red for negative association. Left: default alphabetic ordering of categories; right: occupational categories ordered by status.

{fig:glass-mosaic}

difference $\Delta y_{ij} = y_{ij,1931} - y_{ij,1932}$. Table 1.4 shows these values in a 10×6 table with the rows and columns sorted by their means (main-effect ordering). In addition, the table cells have been colored according to the sign and magnitude of the year difference. The shading scheme uses blue for large positive values and red for large negative values, with a white background for intermediate values. The shading intensity values were determined as $|\Delta y_{ij}| > \{2, 3\} \times \hat{\sigma}(\Delta y_{ij})$.

Effect ordering and color rendering have the result of revealing a new effect, shown as a regular progression in the body of the table. The negative values for Morris now immediately stand out. In addition, the largely positive other values show a lower-triangular pattern, with the size of the yield difference increasing with both row and column means. Against this background, one other cell, for Velvet grown at Grand Rapids stands out with an anomalous negative value.

Although the use of color for graphs is now more common in some journals, color and other rendering details in tables are still difficult. The published version of Table 1.4 (Friendly and Kwan, 2003, Table 3) was forced to use only font shape (normal, italics) to distinguish positive and negative values.

△

Finally, effect ordering is also usefully applied to the variables in multivariate data sets, which by default, are often ordered in data displays according to their position in a data frame or alphabetically. **TODO: DM: Remove this section if parallel coordinates plots are removed from Sec. 5.7.**
MF: I'd like to keep this example

{ex:1.7}

EXAMPLE 1.7: Iris data

The classic *iris* data set (Anderson, 1935, Fisher, 1936b) gives the measurements in centimeters of the variables sepal length and width and petal length and width, respectively, for 50 flowers from each of 3 species of iris, *Iris setosa*, *versicolor*, and *virginica*. Such multivariate data are often displayed in **parallel coordinate plots**, using a separate vertical axis for each variable, scaled from its minimum to maximum.

The default plot, with variables shown in their data frame order is shown in the left panel of Figure 1.11, and gives rise to the epithet *spaghetti plot* for such displays because of the large number

Table 1.4: Barley data, yield differences, 1931-1932, sorted by mean difference, and shaded by value

{tab:barley2c}

Variety	Site						Mean
	Morris	Duluth	University Farm	Grand Rapids	Waseca	Crookston	
No. 475	-22	6	-5	4	6	12	0.1
Wisconsin No. 38	-18	2	1	14	1	14	2.4
Velvet	-13	4	13	-9	13	9	2.9
Peatland	-13	1	5	8	13	16	4.8
Manchuria	-7	6	0	11	15	7	5.5
Trebi	-3	3	7	9	15	5	6.1
Svansota	-9	3	8	13	9	20	7.3
No. 462	-17	6	11	5	21	18	7.4
Glabron	-6	4	6	15	17	12	8.0
No. 457	-15	11	17	13	16	11	8.8
Mean	-12.2	4.6	6.3	8.2	12.5	12.5	5.3

of line crossings. This feature arises because one variable, sepal width, has negative relations in the species means with the other variables. Simple rearrangement of the variables to put sepal width last (or first) makes the relations among the species and the variables more apparent, as shown in the right panel of Figure 1.11. This plot has also been enhanced by using *alpha-blending* (partial transparency) of thicker lines, so that the density of lines is more apparent.

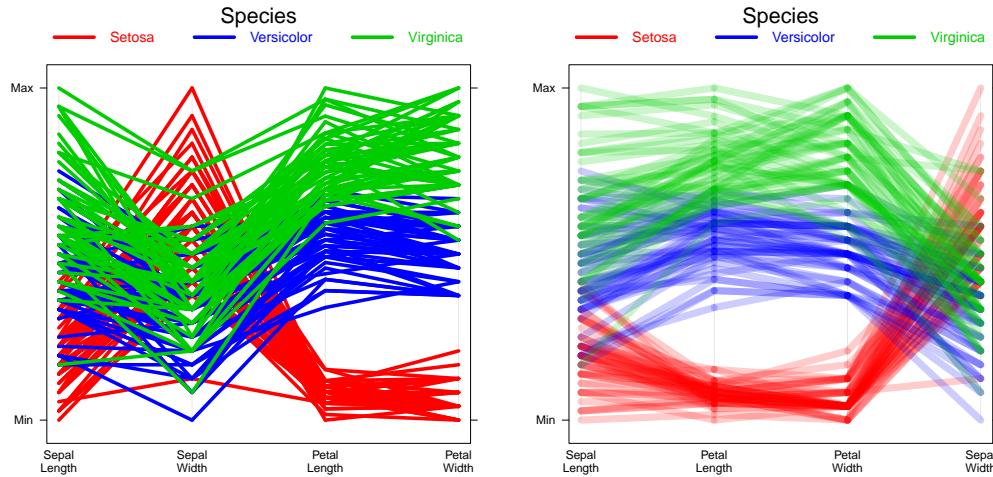


Figure 1.11: Parallel coordinates plots of the Iris data. Left: Default variable order; right: Variables ordered to make the pattern of correlations more coherent.
Fig:iris_parallel

Parallel coordinate plots for categorical data are discussed in an online supplement on the web site for the book. A general method for reordering variables in multivariate data visualizations based on cluster analysis was proposed by Hurley (2004).



1.4.4 Interactive and dynamic graphics

:intro-interactive}

Graphics displayed in print form, such as this book, are necessarily static and fixed at the time they are designed and rendered as an image. Yet, recent developments in software, web technology and media alternative to print have created the possibility to extend graphics in far more useful and interesting ways, for both presentation and analysis purposes.

Interactive graphics allow the viewer to directly manipulate the statistical and visual components of graphical display. These range from

- graphical controls (sliders, selection boxes and other widgets) to control details of an analysis (e.g., a smoothing parameter) or graph (colors and other graphic details), to
- higher-level interaction including zooming in or out, drilling down to a data subset, linking multiple displays, selecting terms in a model and so forth.

The important effect is that the analysis and/or display is immediately re-computed and updated visually.

In addition, **dynamic graphics** use animation to show a series of views, as frames in a movie. Adding time as an additional dimension allows far more possibilities, for example showing a rotating view of a 3D graph or showing smooth transitions or interpolations from one view to another.

There are now many packages in R providing interactive and dynamic plots (e.g., `rggobi`, `ipplots`) as well as capabilities to incorporate these into interactive documents, presentations and web pages (e.g., `rCharts`, `googleVis`). The `animate` package facilitates creating animated graphics and movies in a variety of formats. The RStudio editor and development environment⁵ provides its own `manipulate` package, as well as the `shiny` framework for developing interactive R web applications.

{ex:512paths}

EXAMPLE 1.8: 512 paths to the White House

Shortly before the 2012 U.S. presidential election (November 2, 2012) the *New York Times* published an interactive graphic⁶, designed by Mike Bostock And Shan Carter.⁷ showing the effect that a win for Barack Obama or Mitt Romney in the nine most highly contested states would have on the chances that either candidate would win the presidency.

With these nine states most in play there are $2^9 = 512$ possible outcomes but with different number of votes in the Electoral College. In Figure 1.12, a win for Obama in Florida and Virginia was selected, with wins for Romney in Ohio and North Carolina. Most other selections also lead to a win by Obama, but those with the most votes are made most visible at the top. An R version of this chart was created using the `rCharts` package.⁸ The design of this graphic as a **binary tree** was chosen here, but another possibility would be a **treemap** graphic (Shneiderman, 1992) or a mosaic plot.



1.4.5 Visualization = Graphing + Fitting + Graphing . . .

{sec:vis}

Look here, upon this picture, and on this.

Shakespeare, Hamlet

Statistical summaries, hypothesis tests, and the numerical parameters derived in fitted models are

⁵<http://www.rstudio.com>

⁶<http://www.nytimes.com/interactive/2012/11/02/us/politics/paths-to-the-white-house.html>

⁷see: <https://source.opennews.org/en-US/articles/nyts-512-paths-white-house/> for a description of their design process.

⁸http://timelyportfolio.github.io/rCharts_512paths/

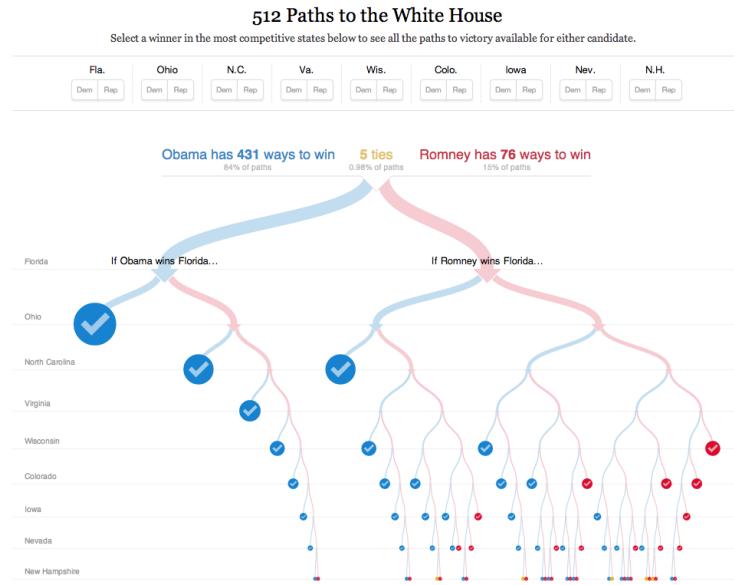


Figure 1.12: 512 paths to the White House. This interactive graphic allows the viewer to select a winner in any one or more of the nine most highly contested U.S. states and highlights the number of paths leading to a win by Obama or Romney, sorted and weighted by the number of Electoral College votes.

{fig:nyt_512paths}

designed to capture a particular feature of the data. A quick analysis of the data from Table 1.1, for example, shows that $1198/2691 = 44.5\%$ of male applicants were admitted, compared to $557/1835 = 30.4\%$ of female applicants.

Statistical tests give a Pearson χ^2 of 92.2 with 1 degree of freedom for association between admission and gender ($p < 0.001$), and various measures for the strength of association. Expressed in terms of the *odds ratio*, males were apparently 1.84 times as likely to be admitted as females, with 99% confidence bounds (1.56, 2.17). Each of these numbers expresses some part of the relationship between gender and admission in the Berkeley data. Numerical summaries such as these are each designed to compress the information in the data, focusing on some particular feature. **TODO: Use this for a lab exercise in Ch 2.**

In contrast, the visualization approach to data analysis is designed to (a) expose information and structure in the data, (b) supplement the information available from numerical summaries, and (c) suggest more adequate models. In general, the visualization approach seeks to serve the needs of both summarization and exposure.

This approach recognizes that both data analysis and graphing are *iterative* processes. You should not expect that any one model captures all features of the data, any more than we should expect that a single graph shows all that may be seen. In most cases, your initial steps should include some graphical display guided by understanding of the subject matter of the data. What you learn from a graph may then help suggest features of the data to be incorporated into a fitted model. Your desire to ensure that the fitted model is an adequate summary may then lead to additional graphs.

The precept here is that

$$\text{Visualization} = \text{Graphing} + \text{Fitting} + \text{Graphing} \dots$$

where the ellipsis indicates the often iterative nature of this process. Even for descriptive purposes,

an initial fit of salient features can be removed from the data, giving residuals (departures from a model). Displaying the residuals may then suggest additional features to account for.

Simple examples of this idea include detrending time series graphs to remove overall and seasonal effects and plots of residuals from main-effect models for ANOVA designs. For categorical data, mosaic plots (Chapter 5) display the unaccounted-for association between variables by shading, as in Figure 1.10. Additional models and plots considered in Section 8.7 can reveal additional structure in square tables beyond the obvious effect that sons tend most often to follow in their fathers' footsteps.

{ex:donner0a}

EXAMPLE 1.9: Donner party

The graphs in Figure 1.3 and Figure 1.4 suggest three different initial descriptions for survival in the Donner party. Yet they ignore all other influences, of which gender and family structure might also be important. A more complete understanding of this data can be achieved by taking these effects into account, both in fitted models and graphs. See Example 7.9 for a continuation of this story.

△

{ex:nasa}

EXAMPLE 1.10: Space shuttle disaster

The space shuttle *Challenger* mentioned in Example 1.2 exploded 73 seconds after take-off on January 28, 1986. Subsequent investigation presented to the presidential commission headed by William Rogers determined that the cause was failure of the O-ring seals used to isolate the fuel supply from burning gases. The story behind the *Challenger* disaster is perhaps the most poignant missed opportunity in the history of statistical graphics. See Tufte (1997) for a complete exposition. It may be heartbreaking to find out that some important information was there, but the graph maker missed it.

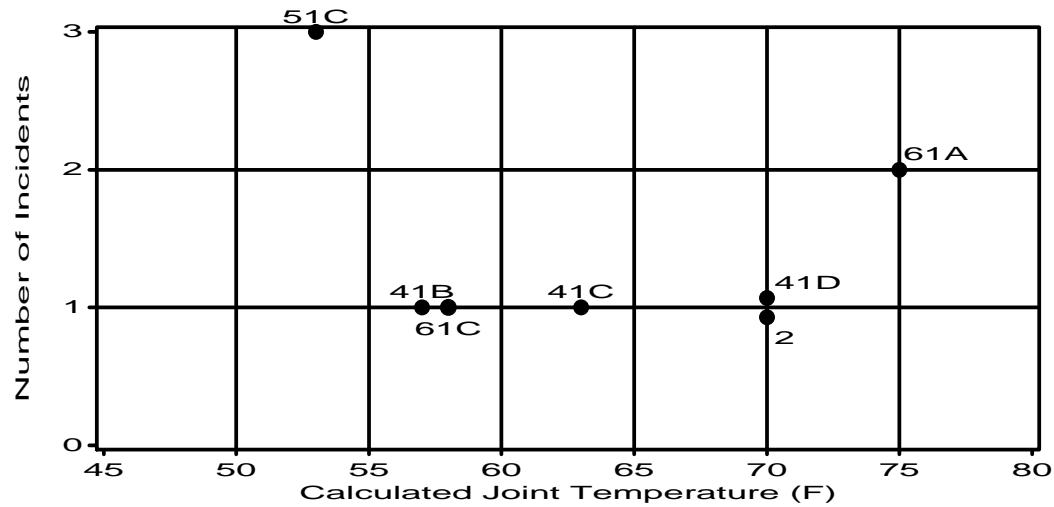


Figure 1.13: NASA Space Shuttle pre-launch graph prepared by the engineers at Morton Thiokol

{fig:nasa0}

Engineers from Morton Thiokol, manufacturers of the rocket motors, had been worried about the effects of unseasonably cold weather on the O-ring seals and recommended aborting the flight. NASA staff analysed the data, tables and charts submitted by the engineers and concluded that there was insufficient evidence to cancel the flight.

The data relating O-ring failures to temperature were depicted as in Figure 1.13, our candidate for the most misleading graph in history. There had been 23 previous launches of these rockets giving data on the number of O-rings (out of 6) that were seen to have suffered some damage or

failure. However, the engineers omitted the observations where no O-rings failed or showed signs of damage, believing that they were uninformative.

Examination of this graph seemed to indicate that there was no relation between ambient temperature and failure. Thus, the decision to launch the *Challenger* was made, in spite of the initial concerns of the Morton Thiokol engineers. Unfortunately, those observations had occurred when the launch temperature was relatively warm ($65 - 80^{\circ}\text{F}$.⁹) and were indeed informative. The coldest temperature at any previous launch was 53° ; when *Challenger* was launched on January 28, the temperature was a frigid 31° .

These data have been analyzed extensively (Dalal *et al.*, 1989, Lavine, 1991). Tufte (1997) gives a thorough and convincing visual analysis of the evidence available prior to the launch. We consider statistical analysis of these data in Chapter 7, Example 7.4.

But, what if the engineers had simply made a better graph? At the very least, that would entail (a) drawing a smoothed curve to fit the points (to show the trend) (b) removing the background grid lines (which obscure the data). Figure 1.14 shows a revised version of the same graph, highlighting the non-zero observations and adding a simple quadratic curve to allow for a possible non-linear relationship. For comparison, the excluded zero observations are also shown in grey. This plot, even showing only the non-zero points should have caused any engineer to conclude that either: (a) the data were wrong, or (b) there were excessive risks associated with both high and low temperatures. But it is well-known that brittleness of the rubber used in the O-rings is inversely proportional to Temperature cubed, so prudent interest might have focussed on the first possibility.⁹

Done: DM: Maybe remove coda: example is quite long already. MF: Made a footnote



1.4.6 The 80-20 rule

The Italian economist Vilfredo Pareto observed in 1906 that 80% of the land in Italy was owned by 20% of the population and this ratio also applied in other countries. It also applied to the yield of peas from peapods in his garden (Pareto, 1971). This idea became known as the **Pareto principle** or the **80–20 rule**. The particular 80/20 ratio is not as important as the more general idea of the uneven distribution of results and causes in a variety of areas.

Common applications are the rules of thumb that: (a) in business 80% of sales come from 20% of clients; (b) in criminology 80% of crimes are said to be committed by 20% of the population. (c) In software development, it is said that 80% of errors and (d) crashes can be eliminated by fixing the top 20% most reported bugs or that 80% of errors reside in 20% of the code.

The **Pareto chart** was designed to display the frequency distribution of a variable with a histogram or bar chart together with a cumulative line graph to highlight the most frequent category, and the **Pareto distribution** gives a mathematical form to such distributions with a parameter α (the **Pareto index**) reflecting the degree of inequality.

Applied to statistical graphics, the precept is that

20% of your effort can generate 80% of your desired result in producing a given plot.

This is good news for exploratory graphs you produce for yourself. Very often, the default settings will give a reasonable result, or you will see immediately something simple to add or change to make the plot easier to understand.

The bad news is the corollary of this rule:

⁹A coda to this story shows the role of visual explanation in practice as well (Tufte, 1997, p. 50–53). The Rogers Commission contracted the reknown theoretical physicist Richard Feynman to contribute to their investigation. He determined that the most probable cause of the shuttle failure was the lack of resiliency of the rubber O-rings at low temperature. But how could he make this point convincingly? At a televised public hearing, he took a piece of the O-ring material, squeezed it in C-clamp and plunged it into a glass of ice water. After a few minutes, he released the clamp, and the rubber did not spring back to shape. He mildly said, "... there is no resilience in this particular material when it is at a temperature of 32 degrees. I believe this has some significance for our problem" (Feynman, 1988).

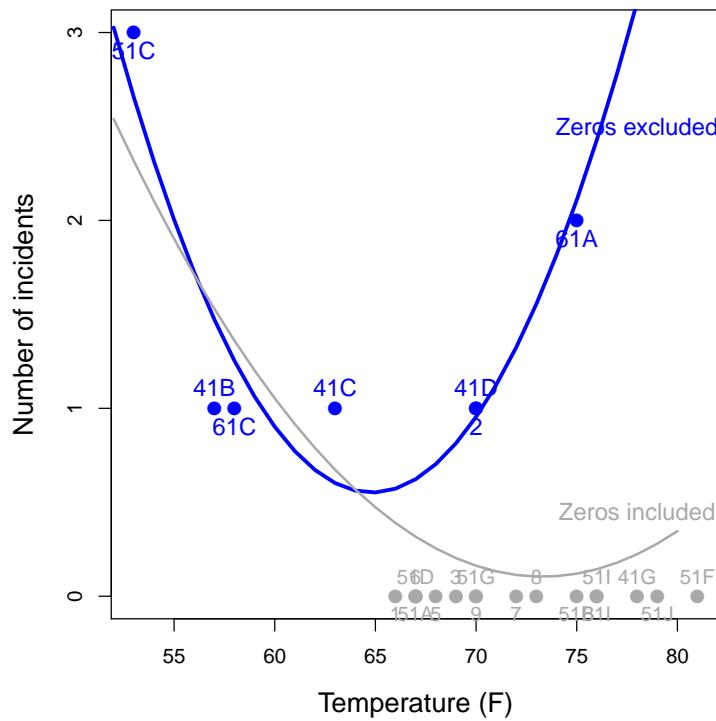


Figure 1.14: Re-drawn version of the NASA pre-launch graph, showing the locations of the excluded observations and with fitted quadratics for both sets of observations
 (fig:nasa)

80% of your effort may be required to produce the remaining 20% of a finished graph.

This is particularly important for presentation graphs, where several iterations may be necessary to get it right (or right enough) for your communication purposes. Some important details are:

graph title A presentation graphic can be more effective when it announces the main point or conclusion in the graphic title, as in Figure 1.8.

axis and value labels Axes should be labelled with meaningful variable descriptions (and perhaps the data units) rather than just plot defaults (e.g., “Temperature (degrees F)” in Figure 1.2, not `temp`). Axis values are often more of a challenge for categorical variables, where their text labels often overlap, requiring abbreviation, a smaller font or text rotation.

grouping attributes Meaningfully different subsets of the data should be rendered with distinct visual attributes such as color, shape, and line style, and sometimes with more than one.

legends and direct labels Different data groups in a graphic display shown by color, shape, etc. usually need at least a graphic legend defining the symbols and group labels. Sometimes you can do better by applying the labels directly to the graphical elements,¹⁰ as was done in Figure 1.14.

legibility A common failure in presentation graphs in journals and lectures is the use of text fonts

¹⁰For example, the `identify()` function allows points in a plot to be labeled interactively with a mouse. The `directlabels` package provides a general method for a variety of plots.

too small to be read easily. One rule of thumb is to hold the graph at arms length for a journal and put it on the floor for a lecture slide. If you can't read the labels, the font is too small.

plot annotations Beyond the basic graphic data display, additional annotations can add considerable information to interpret the context or uncertainty, as in the use of plot envelopes to show confidence bands or regions (see Figure 1.3 and Figure 1.4).

aspect ratio Line graphs (such as Figure 3.1) are often easiest to understand when the ratio of height to width is such that line segments have an average slope near 1.0 (Cleveland *et al.*, 1988). In R, you can easily manipulate a graph window manually with a mouse to observe this effect and find an aspect ratio that looks right.

Moreover, in graphs for biplots and correspondence analysis (Chapter 6), interpretation involves distances between points and angles between line segments. This requires an aspect ratio that equates the units on the axes. Careful software will do this for you,¹¹ and you should resist the temptation to re-shape the plot.

Nearly all of the graphs in this book were produced using R code in scripts saved as files. This has the advantages of reproducibility and enhancement: just re-run the code, or tweak it to improve a graph. If this is too hard, you can always use an external graphics editor (Gimp, Inkscape, Adobe Illustrator, etc.) to make improvements manually.

1.5 Chapter summary

- Categorical data differs from quantitative data because the variables take on discrete values (ordered or unordered, character or numeric) rather than continuous numerical values. Consequently, such data often appear in aggregated form representing category frequencies or in tables.
- Data analysis methods for categorical data are comprised of those concerned mainly with testing particular hypotheses versus those that fit statistical models. Model building methods have the advantages of providing parameter estimates and model-predicted values, along with measures of uncertainty (standard errors).
- Graphical methods can serve different purposes for different goals (data analysis versus presentation), and these suggest different design principles that a graphic should respect to achieve a given communication goal.
- For categorical data, some graphic forms (bar charts, line graphs, scatterplots) used for quantitative data can be readily adapted to discrete variables. However, frequency data often requires novel graphics using area and other visual attributes.
- Graphics can be far more effective when categorical variables are ordered to facilitate comparison of the effects to be seen and rendered to facilitate detection of patterns, trends or anomalies.
- The visualization approach to data analysis often entails a sequence of intertwined steps involving graphing and model fitting.
- Producing effective graphs for presentation is often hard work, requiring attention to details that support or detract from your communication goal.

¹¹For example using the graphics parameter `asp=1`, `eqsplot()` in `MASS`, or the equivalents in `lattice` (`aspect="iso"`) and `ggplot2` (`coord_equal`).

1.6 Further reading

{sec:ch01-reading}

sec:ch01-exercises1

1.7 Lab exercises

Exercise 1.1 A web page, “The top ten worst graphs,” http://www.biostat.wisc.edu/~kbroman/topten_worstgraphs/ by Karl Broman lists his picks for the worst graphs (and a table) that have appeared in the statistical and scientific literature. Each entry links to graph(s) and a brief discussion of what is wrong and how it could be improved.

- (a) Examine a number of recent issues of a scientific or statistical journal in which you have some interest. Find one or more examples of a graph or table that is a particularly bad use of display material to summarize and communicate research findings. Write a few sentences indicating how or why the display fails and how it could be improved.
- (b) Do the same task for some popular magazine or newspaper that uses data displays to supplement the text for some story. Again, write a few sentences describing why the display is bad and how it could be improved.

{lab:1.2}

Exercise 1.2 As in the previous exercise, examine the recent literature in recent issues of some journal of interest to you. Find one or more examples of a graph or table that you feel does a *good* of summarizing and communicating research findings.

- (a) Write a few sentences describing why you chose these displays.
- (b) Now take the role of a tough journal reviewer. Are there any features of the display that could be modified to make them more effective?

{lab:1.3}

Exercise 1.3 Infographics are another form of visual displays, quite different from the data graphics featured in this book, but often based on some data or analysis. Do a Google image search for the topic “Global warming” to see a rich collection.

- (a) Find and study one or two that attempt some visual explanation of causes and/or effects of global warming. Describe the main message in a sentence or two.
- (b) What visual and graphic features are used in these to convey the message?

{lab:1.4}

Exercise 1.4 The Wikipedia web page en.wikipedia.org/wiki/Portal:Global_warming gives a few data-based graphics on the topic of global warming. Read the text and study the graphs.

- (a) Write a short figure title for each that would announce the conclusion to be drawn in a presentation graphic.
- (b) Write a figure caption for each that would explain what is shown and the important graphical details for a reader to understand.

Chapter 2

Working with categorical data

Creating and manipulating categorical data sets requires some skills and techniques in R beyond those ordinarily used for quantitative data. This chapter illustrates these for the main formats for categorical data: case form, frequency form and table form.

{ch:working}

Done: DM: The chapter does not cover NA handling, although this is needed in some chapters. Maybe treat this in the also missing section on subsetting? — The note for `table` and in the subsetting section should suffice.

Categorical data can be represented as data sets in various formats: case form, frequency form, and table form. This chapter describes and illustrates the skills and techniques in R needed to input, create and manipulate R data objects to represent categorical data, and convert these from one form to another for the purposes of statistical analysis and visualization which are the subject of the remainder of the book.

As mentioned earlier, this book assumes that you have at least a basic knowledge of the R language and environment, including interacting with the R console (Rgui for Windows, R.app for Mac OS X) or some other editor/environment (e.g., R Studio), loading and using R functions in packages (e.g., `library(vcd)`) getting help for these from R (e.g., `help(matrix)`), etc. This chapter is therefore devoted to covering those topics beyond such basic skills needed in the book.¹

2.1 Working with R data: vectors, matrices, arrays and data frames

{sec:Rdata}

R has a wide variety of data structures for storing, manipulating and calculating with data. Among these, vectors, matrices, arrays and data frames are most important for the material in this book.

In R, a **vector** is a collection of values, like numbers, character strings, or logicals (TRUE, FALSE), and often correspond to a variable in some analysis. Matrices are rectangular arrays like a traditional table, composed of vectors in their columns or rows. Arrays add additional dimensions, so that, for example, a 3-way table can be represented as composed of rows, columns and layers. An important consideration is that the values in vectors, matrices and arrays must all be of the same *mode*, e.g., numbers or character strings. A **data frame** is a rectangular table, like a traditional data set in other statistical environments, and composed of rows and columns like a matrix, but allowing

¹Some excellent introductory treatments of R are: Fox and Weisberg (2011, Chapter 2), ... Tom Short's *R Reference Card*, <http://cran.us.r-project.org/doc/contrib/Short-refcard.pdf> is a handy 4-page summary of the main functions. The web sites Quick-R <http://www.statmethods.net/> and Cookbook for R <http://www.cookbook-r.com/> provide very helpful examples, organized by topics and tasks.

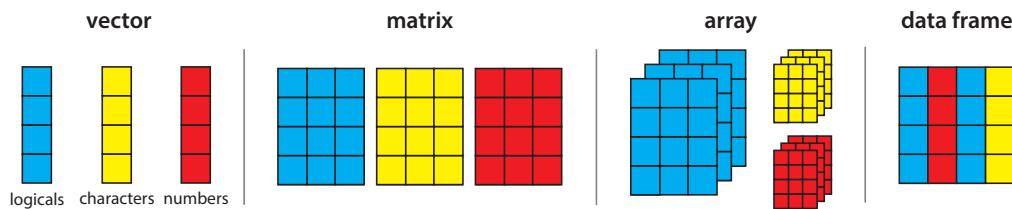


Figure 2.1: Principal data structures and data types in R. Colors represent different data types: numeric, character, logical.

{fig:datatypes}

variables (columns) of different types. These data structures and the types of data they can contain are illustrated in Figure 2.1. A more general data structure is a *list*, a generic vector which can contain any other types of objects (including lists, allowing for *recursive* data structures). A data frame is basically a list of equally-sized vectors, each representing a column of the data frame.

TODO: Delete subsections on vectors, matrices and arrays?

2.1.1 Vectors

The simplest data structure in R is a *vector*, a one-dimensional collection of elements of the same type. An easy way to create a vector is with the `c()`, which combines its arguments. The following examples create and print vectors of length 4, containing numbers, character strings and logical values respectively:

```
> c(17, 20, 15, 40)
[1] 17 20 15 40
> c("female", "male", "female", "male")
[1] "female" "male"    "female" "male"
> c(TRUE, TRUE, FALSE, FALSE)
[1] TRUE  TRUE FALSE FALSE
```

To store these values in variables, R uses the assignment operator (`<-`) or equals sign (`=`). This creates a variable named on the left-hand side. An assignment doesn't print the result, but a bare expression does, so you can assign and print by surrounding the assignment with `()`.

```
> count <- c(17, 20, 15, 40)                      # assign
> count                                         # print
[1] 17 20 15 40
> (sex <- c("female", "male", "female", "male"))   # both
[1] "female" "male"    "female" "male"
> (passed <- c(TRUE, TRUE, FALSE, FALSE))
[1] TRUE  TRUE FALSE FALSE
```

Other useful functions for creating vectors are:

- The `:` operator for generating consecutive integer sequences, e.g., `1:10` gives the integers 1 to 10. The `seq()` function is more general, taking the forms `seq(from, to)`, `seq(from, to, by=)`, and `seq(from, to, length=)` where the optional argument `by` specifies the interval between adjacent values and `length` gives the desired length of the result.
- The `rep()` function generates repeated sequences, replicating its first argument (which may be a vector) a given number of `times`, to a given `length` or each a given multiple.

```
> seq(10, 100, by = 10)      # give interval
[1] 10 20 30 40 50 60 70 80 90 100
> seq(0, 1, length = 11)     # give length
[1] 0.0 0.1 0.2 0.3 0.4 0.5 0.6 0.7 0.8 0.9 1.0
> (sex <- rep(c("female", "male"), times = 2))
[1] "female" "male"   "female" "male"
> (sex <- rep(c("female", "male"), length.out = 4)) # same
[1] "female" "male"   "female" "male"
> (passed <- rep(c(TRUE, FALSE), each = 2))
[1] TRUE  TRUE FALSE FALSE
```

2.1.2 Matrices

A **matrix** is a two-dimensional array of elements of the same type composed in a rectangular array of rows and columns. Matrices can be created by the function `matrix(values, nrow, ncol)`, which reshapes the elements in the first argument (`values`) to a matrix with `nrow` rows and `ncol` columns. By default, the elements are filled in columnwise, unless the optional argument `byrow = TRUE` is given.

```
> (matA <- matrix(1:8, nrow = 2, ncol = 4))
[,1] [,2] [,3] [,4]
[1,]    1    3    5    7
[2,]    2    4    6    8
> (matB <- matrix(1:8, nrow = 2, ncol = 4, byrow = TRUE))
[,1] [,2] [,3] [,4]
[1,]    1    2    3    4
[2,]    5    6    7    8
> (matC <- matrix(1:4, nrow = 2, ncol = 4))
[,1] [,2] [,3] [,4]
[1,]    1    3    1    3
[2,]    2    4    2    4
```

The last example illustrates that the values in the first argument are recycled as necessary to fill the given number of rows and columns.

All matrices have a `dimensions` attribute, a vector of length two giving the number of rows and columns, retrieved with the function `dim()`. Labels for the rows and columns can be assigned using

`dimnames()`,² which takes a list of two vectors for the row names and column names respectively. To see the structure of a matrix (or any other R object) and its attributes, you can use the `str()` function, as shown in the example below.

```
> dim(matA)
[1] 2 4
> str(matA)
int [1:2, 1:4] 1 2 3 4 5 6 7 8
> dimnames(matA) <- list(c("M", "F"), LETTERS[1:4])
> matA
   A B C D
M 1 3 5 7
F 2 4 6 8
> str(matA)
int [1:2, 1:4] 1 2 3 4 5 6 7 8
- attr(*, "dimnames")=List of 2
..$ : chr [1:2] "M" "F"
..$ : chr [1:4] "A" "B" "C" "D"
```

Additionally, names for the row and column *variables* themselves can also be assigned in the `dimnames` call by giving each dimension vector a name.

```
> dimnames(matA) <- list(sex = c("M", "F"), group = LETTERS[1:4])
> ## or: names(dimnames(matA)) <- c("Sex", "Group")
> matA
   group
sex A B C D
M 1 3 5 7
F 2 4 6 8
> str(matA)
int [1:2, 1:4] 1 2 3 4 5 6 7 8
- attr(*, "dimnames")=List of 2
..$ sex : chr [1:2] "M" "F"
..$ group: chr [1:4] "A" "B" "C" "D"
```

(`LETTERS` is a predefined character vector of the 26 uppercase letters). Matrices can also be created or enlarged by “binding” vectors or matrices together by rows or columns:

- `rbind(a, b, c)` creates a matrix with the vectors `a`, `b` and `c` as its rows, recycling the elements as necessary to the length of the longest one.
- `cbind(a, b, c)` creates a matrix with the vectors `a`, `b` and `c` as its columns.
- `rbind(mat, a, b, ...)` and `cbind(mat, a, b, ...)` add additional rows (columns) to a matrix `mat`, recycling or subsetting the elements in the vectors to conform with the size of the matrix.

²The `dimnames` can also be specified as an optional argument to `matrix()`.

```
> rbind(matA, c(10, 20))

  A  B  C  D
M 1  3  5  7
F 2  4  6  8
10 20 10 20

> cbind(matA, c(10, 20))

  A  B  C  D
M 1  3  5  7 10
F 2  4  6  8 20
```

Rows and columns can be swapped (transposed) using `t()`:

```
> t(matA)

      sex
group M F
      A 2
      B 4
      C 6
      D 8
```

Finally, we note that basic computations involving matrices are performed *element-wise*:

```
> 2 * matA / 100

      group
sex     A      B      C      D
M 0.02 0.06 0.10 0.14
F 0.04 0.08 0.12 0.16
```

Special operators and functions do exist for matrix operations, such as `%*%` for the matrix product.

2.1.3 Arrays

Higher-dimensional arrays are less frequently encountered in traditional data analysis, but they are of great use for categorical data, where frequency tables of three or more variables can be naturally represented as arrays, with one dimension for each table variable.

The function `array(values, dim)` takes the elements in `values` and reshapes these into an array whose dimensions are given in the vector `dim`. The number of dimensions is the length of `dim`. As with matrices, the elements are filled in with the first dimension (rows) varying most rapidly, then by the second dimension (columns) and so on for all further dimensions, which can be considered as layers. A matrix is just the special case of an array with two dimensions.

```
> (arrayA <- array(1:16, dim = c(2, 4, 2)))      # 2 rows, 4 columns, 2 layers

, , 1

[,1] [,2] [,3] [,4]
[1,]    1    3    5    7
[2,]    2    4    6    8

, , 2

[,1] [,2] [,3] [,4]
[1,]    9   11   13   15
[2,]   10   12   14   16
```

```
> str(arrayA)
int [1:2, 1:4, 1:2] 1 2 3 4 5 6 7 8 9 10 ...
> (arrayB <- array(1:16, dim = c(2, 8)))           # 2 rows, 8 columns
[,1] [,2] [,3] [,4] [,5] [,6] [,7] [,8]
[1,]    1    3    5    7    9   11   13   15
[2,]    2    4    6    8   10   12   14   16
> str(arrayB)
int [1:2, 1:8] 1 2 3 4 5 6 7 8 9 10 ...
```

In the same way that we can assign labels to the rows, columns and variables in matrices, we can assign these attributes to `dimnames` (`arrayA`), or include this information in a `dimnames=` argument to `array()`.

```
> dimnames(arrayA) <- list(sex = c("M", "F"),
+                           group = letters[1:4],
+                           time = c("Pre", "Post"))
> arrayA
, , time = Pre
group
sex a b c d
M 1 3 5 7
F 2 4 6 8
, , time = Post
group
sex a b c d
M 9 11 13 15
F 10 12 14 16
> str(arrayA)
int [1:2, 1:4, 1:2] 1 2 3 4 5 6 7 8 9 10 ...
- attr(*, "dimnames")=List of 3
..$ sex : chr [1:2] "M" "F"
..$ group: chr [1:4] "a" "b" "c" "d"
..$ time : chr [1:2] "Pre" "Post"
```

Arrays in R can contain any single type of elements—numbers, character strings, logicals. R also has a variety of functions (e.g., `table()`, `xtabs()`) for creating and manipulating "table" objects, which are specialized forms of matrices and arrays containing integer frequencies in a contingency table. These are discussed in more detail below (Section 2.4.1).

2.1.4 data frames

Data frames are the most commonly used form of data in R and more general than matrices in that they can contain columns of different types. For statistical modeling, data frames play a special role, in that many modeling functions are designed to take a data frame as a `data=` argument, and then find the variables mentioned within that data frame. Another distinguishing feature is that discrete variables (columns) like character strings ("M", "F") or integers (1, 2, 3) in data frames can be represented as *factors*, which simplifies many statistical and graphical methods.

A data frame can be created using keyboard input with the `data.frame()` function, applied

to a list of objects, `data.frame(a, b, c, ...)`, each of which can be a vector, matrix or another data frame, but typically all containing the same number of rows. This works roughly like `cbind()`, collecting the arguments as columns in the result.

The following example generates $n = 100$ random observations on three discrete factor variables, A, B, sex, and a numeric variable, age. As constructed, all of these are statistically independent, since none depends on any of the others. The function `sample()` is used here to generate n random samples from the first argument allowing replacement (`replace = TRUE`). The `rnorm()` function produces a vector of n normally distributed values with mean 30 and standard deviation 5. The call to `set.seed()` guarantees the reproducibility of the resulting data. Finally, all four variables are combined into the data frame `mydata`.

```
> set.seed(12345)    # reproducibility
> n <- 100
> A <- factor(sample(c("a1", "a2"), n, replace = TRUE))
> B <- factor(sample(c("b1", "b2"), n, replace = TRUE))
> sex <- factor(sample(c("M", "F"), n, replace = TRUE))
> age <- round(rnorm(n, mean = 30, sd = 5))
> mydata <- data.frame(A, B, sex, age)
> head(mydata, 5)

  A B sex age
1 a2 b1 F 22
2 a2 b2 F 33
3 a2 b2 M 31
4 a2 b2 F 26
5 a1 b2 F 29

> str(mydata)

'data.frame': 100 obs. of 4 variables:
 $ A : Factor w/ 2 levels "a1","a2": 2 2 2 2 1 1 1 2 2 2 ...
 $ B : Factor w/ 2 levels "b1","b2": 1 2 2 2 2 2 2 1 1 ...
 $ sex: Factor w/ 2 levels "F","M": 1 1 2 1 1 1 2 2 1 1 ...
 $ age: num 22 33 31 26 29 29 38 28 30 27 ...
```

Rows, columns and individual values in a data frame can be manipulated in the same way than a matrix. Additionally, variables can be extracted using the `$` operator:

```
> mydata[1,2]
[1] b1
Levels: b1 b2

> mydata$sex
[1] F F M F F F M M F F M F M M F F M M M F F F M M F
[31] M F M F F F F M M F F F F F F F M F M F M M F F M M M M
[61] F F F F F M M F F F M M M F F M F M M F M M M M F F F
[91] F F M M F M F M F M
Levels: F M

> ##same as: mydata[, "sex"] or mydata[, 3]
```

Values in data frames can conveniently be edited using, e.g., `fix(mydata)`, opening a simple, spreadsheet-like editor.

For real data sets, it is usually most convenient to read these into R from external files, and this is easiest using plain text (ASCII) files with one line per observation and fields separated by commas (or tabs), and with a first header line giving the variable names— called *comma-separated* or CSV

format. If your data is in the form of Excel, SAS, SPSS or other file format, you can almost always export that data to CSV format first.³

The function `read.table()` has many options to control the details of how the data are read and converted to variables in the data frame. Among these some important options are:

```
header indicates whether the first line contains variable names. The default is FALSE unless the
first line contains one fewer field than the number of columns;
sep (default: "") meaning white space, i.e., one or more spaces, tabs or newlines) specifies the
separator character between fields;
stringsAsFactors (default: TRUE) determines whether character string variables should be
converted to factors;
na.strings (default: "NA") one or more strings which are interpreted as missing data values
(NA);
```

For delimited files, `read.csv()` and `read.delim()` are convenient wrappers to `read.table()`, with default values `sep=","` and `sep=" "` respectively, and `header=TRUE`.

{ex:ch2-arth-csv}

EXAMPLE 2.1: Arthritis treatment

The file `Arthritis.csv` contains data in CSV format from Koch and Edwards (1988), representing a double-blind clinical trial investigating a new treatment for rheumatoid arthritis with 84 patients.⁴ The first (“header”) line gives the variable names. Some of the lines in the file are shown below, with ... representing omitted lines:

```
ID,Treatment,Sex,Age,Improved
57,Treated,Male,27,Some
46,Treated,Male,29,None
77,Treated,Male,30,None
17,Treated,Male,32,Marked
...
42,Placebo,Female,66,None
15,Placebo,Female,66,Some
71,Placebo,Female,68,Some
1,Placebo,Female,74,Marked
```

Done: DM: Add this to vcd or vcdExtra. — added footnote.

We read this into R using `read.table()` as shown below:

```
> path <- "ch02/Arthritis.csv" ## set path
> ## for convenience, use path <- file.choose() to retrieve a path
> ## then, use file.show(path) to inspect the data format
> Arthritis <- read.table(path, header = TRUE, sep = ",")
> str(Arthritis)

'data.frame': 84 obs. of  5 variables:
 $ ID      : int  57 46 77 17 36 23 75 39 33 55 ...
 $ Treatment: Factor w/ 2 levels "Placebo","Treated": 2 2 2 2 2 2 2 2 2 2 ...
 $ Sex     : Factor w/ 2 levels "Female","Male": 2 2 2 2 2 2 2 2 2 2 ...
 $ Age     : int  27 29 30 32 46 58 59 63 63 ...
 $ Improved: Factor w/ 3 levels "Marked","None",...: 3 2 2 1 1 1 2 1 2 2 ...
```

Note that the character variables `Treatment`, `Sex` and `Improved` were converted to factors, and the levels of those variables were ordered *alphabetically*. This often doesn’t matter much for binary variables, but here, the response variable, `Improved` has levels that should be considered *ordered*, as “`None`”, “`Some`”, “`Marked`”. We can correct this here by re-assigning

³The `foreign` package contains specialized functions to *directly* read data stored by Minitab, SAS, SPSS, Stata, Systat and other software. There are also a number of packages for reading (and writing) Excel spreadsheets directly (`gdata`, `XLConnect`, `xlsx`). The R manual, *R Data Import/Export* covers many other variations, including data in relational data bases.

⁴This data can be created using: `library(vcd); write.table(Arthritis, file = "Arthritis.csv", quote = FALSE, sep = ",")`

`Arthritis$Improved` using `ordered()`. The topic of re-ordering variables and levels in categorical data is considered in more detail in Section 2.3.

```
> levels(Arthritis$Improved)
[1] "Marked" "None"   "Some"

> Arthritis$Improved <- ordered(Arthritis$Improved,
+                               levels = c("None", "Some", "Marked"))
```

△

2.2 Forms of categorical data: case form, frequency form and table form

{sec:forms}

As we saw in Chapter 1, categorical data can be represented as ordinary data sets in case form, but the discrete nature of factors or stratifying variables allows the same information to be represented more compactly in summarized form with a frequency variable for each cell of factor combinations, or in tables. Consequently, we sometimes find data created or presented in one form (e.g., a spreadsheet data set, a two-way table of frequencies) and want to input that into R. Once we have the data in R, it is often necessary to manipulate the data into some other form for the purposes of statistical analysis, visualizing results and our own presentation. It is useful to understand the three main forms of categorical data in R and how to work with them for our purposes.

2.2.1 Case form

Categorical data in case form are simply data frames, with one or more discrete classifying variables or response variables, most conveniently represented as factors or ordered factors. In case form, the data set can also contain numeric variables (covariates or other response variables), that cannot be accommodated in other forms.

As with any data frame, X, you can access or compute with its attributes using `nrow(X)` for the number of observations, `ncol(X)` for the number of variables, `names(X)` or `colnames(X)` for the variable names and so forth.

{ex:ch2-arth}

EXAMPLE 2.2: Arthritis treatment

The `Arthritis` data is available in case form in the `vcd` package. There are two explanatory factors: Treatment and Sex. Age is a numeric covariate, and Improved is the response—an ordered factor, with levels "None" < "Some" < "Marked". Excluding Age, we would have a $2 \times 2 \times 3$ contingency table for Treatment, Sex and Improved.

```
> data(Arthritis, package = "vcd") # load the data
> names(Arthritis)      # show the variables

[1] "ID"          "Treatment"   "Sex"        "Age"        "Improved"

> str(Arthritis)      # show the structure

'data.frame': 84 obs. of  5 variables:
 $ ID    : int  57 46 77 17 36 23 75 39 33 55 ...
 $ Treatment: Factor w/ 2 levels "Placebo","Treated": 2 2 2 2 2 2 2 2 2 2 ...
 $ Sex   : Factor w/ 2 levels "Female","Male": 2 2 2 2 2 2 2 2 2 2 ...
 $ Age   : int  27 29 30 32 46 58 59 63 63 ...
 $ Improved: Ord.factor w/ 3 levels "None"><"Some"><...: 2 1 1 3 3 3 1 3 1 1 ...

> head(Arthritis, 5)    # first 5 observations, same as Arthritis[1:5,]
```

ID	Treatment	Sex	Age	Improved
1 57	Treated	Male	27	Some
2 46	Treated	Male	29	None
3 77	Treated	Male	30	None
4 17	Treated	Male	32	Marked
5 36	Treated	Male	46	Marked



2.2.2 Frequency form

Data in frequency form is also a data frame, containing one or more discrete factor variables and a frequency variable (often called `Freq` or `count`) representing the number of basic observations in that cell.

This is an alternative representation of a table form data set considered below. In frequency form, the number of cells in the equivalent table is `nrow(X)`, and the total number of observations is the sum of the frequency variable, `sum(X$Freq)`, `sum(X[, "Freq"])` or a similar expression.

EXAMPLE 2.3: General social survey

For small frequency tables, it is often convenient to enter them in frequency form using `expand.grid()` for the factors and `c()` to list the counts in a vector. The example below, from Agresti (2002) gives results for the 1991 General Social Survey, with respondents classified by sex and party identification. As a table, the data look like this:

party			
sex	dem	indep	rep
female	279	73	225
male	165	47	191

We use `expand.grid()` to create a 6×2 matrix containing the combinations of `sex` and `party` with the levels for `sex` given first, so that this varies most rapidly. Then, input the frequencies in the table by columns from left to right, and combine these two results with `data.frame()`.

```
> # Agresti (2002), table 3.11, p. 106
> tmp <- expand.grid(sex = c("female", "male"),
+                      party = c("dem", "indep", "rep"))
> tmp

  sex party
1 female   dem
2 male    dem
3 female  indep
4 male   indep
5 female   rep
6 male    rep

> GSS <- data.frame(tmp, count = c(279, 165, 73, 47, 225, 191))
> GSS

  sex party count
1 female   dem    279
2 male    dem    165
3 female  indep     73
4 male   indep     47
5 female   rep    225
6 male    rep    191

> names(GSS)
```

```
[1] "sex"    "party"  "count"
> str(GSS)

'data.frame': 6 obs. of 3 variables:
 $ sex : Factor w/ 2 levels "female", "male": 1 2 1 2 1 2
 $ party: Factor w/ 3 levels "dem", "indep", ...: 1 1 2 2 3 3
 $ count: num 279 165 73 47 225 191

> sum(GSS$count)
[1] 980
```

The last line above shows that there are 980 cases represented in the frequency table. \triangle

2.2.3 Table form

Table form data is represented as a matrix, array or table object whose elements are the frequencies in an n -way table. The number of dimensions of the table is the length, `length(dim(X))`, of its `dim` (or `dimnames`) attribute, and the sizes of the dimensions in the table are the elements of `dim(X)`. The total number of observations represented is the sum of all the frequencies, `sum(X)`.

{ex:ch2-hec}

EXAMPLE 2.4: Hair color and eye color

A classic data set on frequencies of hair color, eye color and sex is given in table form in `HairEyeColor` in the `datasets` package, reporting the frequencies of these categories for 592 students in a statistics course.

```
> data(HairEyeColor, package = "datasets")      # load the data
> str(HairEyeColor)                            # show the structure

table [1:4, 1:4, 1:2] 32 53 10 3 11 50 10 30 10 25 ...
- attr(*, "dimnames")=List of 3
..$ Hair: chr [1:4] "Black" "Brown" "Red" "Blond"
..$ Eye : chr [1:4] "Brown" "Blue" "Hazel" "Green"
..$ Sex : chr [1:2] "Male" "Female"

> dim(HairEyeColor)                          # table dimension sizes
[1] 4 4 2

> dimnames(HairEyeColor)                    # variable and level names

$Hair
[1] "Black" "Brown" "Red"     "Blond"

$Eye
[1] "Brown" "Blue"   "Hazel" "Green"

$Sex
[1] "Male"   "Female"

> sum(HairEyeColor)                        # number of cases
[1] 592
```

Three-way (and higher-way) tables can be printed in a more convenient form using `structable()` and `ftable()` as described below in Section 2.5. \triangle

Tables are often created from raw data in case form or frequency form using the functions `table()` and `xtabs()` described in Section 2.4.1. For smallish frequency tables that are already

in tabular form, you can enter the frequencies in a matrix, and then assign dimnames and other attributes.

To illustrate, we create the GSS data as a table below, entering the values in the table by rows (byrow=TRUE), as they appear in printed form.

```
> GSS.tab <- matrix(c(279, 73, 225,
+                      165, 47, 191),
+                      nrow = 2, ncol = 3, byrow = TRUE)
> dimnames(GSS.tab) <- list(sex = c("female", "male"),
+                             party = c("dem", "indep", "rep"))
> GSS.tab

      party
sex     dem indep rep
female 279    73 225
male   165    47 191
```

`GSS.tab` is a matrix, not an object of `class("table")`, and some functions are happier with tables than matrices.⁵ You should therefore coerce it to a table with `as.table()`,

```
> GSS.tab <- as.table(GSS.tab)
> str(GSS.tab)

table [1:2, 1:3] 279 165 73 47 225 191
- attr(*, "dimnames")=List of 2
..$ sex : chr [1:2] "female" "male"
..$ party: chr [1:3] "dem" "indep" "rep"
```

{ex:jobsat1}

EXAMPLE 2.5: Job satisfaction

Here is another similar example, entering data on job satisfaction classified by income and level of satisfaction from a 4×4 table given by Agresti (2002, Table 2.8, p. 57).

```
> ## A 4 x 4 table Agresti (2002, Table 2.8, p. 57) Job Satisfaction
> JobSat <- matrix(c(1, 2, 1, 0,
+                     3, 3, 6, 1,
+                     10, 10, 14, 9,
+                     6, 7, 12, 11),
+                     nrow = 4, ncol = 4)
> dimnames(JobSat) <-
+   list(income = c("< 15k", "15-25k", "25-40k", "> 40k"),
+        satisfaction = c("VeryD", "LittleD", "ModerateS", "VeryS"))
> JobSat <- as.table(JobSat)
> JobSat

      satisfaction
income  VeryD LittleD ModerateS VeryS
< 15k    1      3      10      6
15-25k   2      3      10      7
25-40k   1      6      14     12
> 40k    0      1      9      11
```

△

⁵There are quite a few functions in R with specialized methods for "table" objects. For example, `plot(GSS.tab)` gives a mosaic plot and `barchart(GSS.tab)` gives a divided bar chart.

2.3 Ordered factors and reordered tables

{sec:ordered}

As we saw above (Example 2.1), the levels of factor variables in data frames (case form or frequency form) can be re-ordered (and the variables declared as ordered factors) using `ordered()`. As well, the order of the factor values themselves can be rearranged by sorting the data frame using `sort()`.

However, in table form, the values of the table factors are ordered by their position in the table. Thus in the `JobSat` data, both `income` and `satisfaction` represent ordered factors, and the *positions* of the values in the rows and columns reflects their ordered nature, but only implicitly.

Yet, for analysis or graphing, there are occasions when you need *numeric* values for the levels of ordered factors in a table, e.g., to treat a factor as a quantitative variable. In such cases, you can simply re-assign the `dimnames` attribute of the table variables. For example, here, we assign numeric values to `income` as the middle of their ranges, and treat `satisfaction` as equally spaced with integer scores.

```
> dimnames(JobSat)$income <- c(7.5, 20, 32.5, 60)
> dimnames(JobSat)$satisfaction <- 1:4
```

A related case is when you want to preserve the character labels of table dimensions, but also allow them to be sorted in some particular order. A simple way to do this is to prefix each label with an integer index using `paste()`.

```
> dimnames(JobSat)$income <-
+   paste(1:4, dimnames(JobSat)$income, sep = ":")
> dimnames(JobSat)$satisfaction <-
+   paste(1:4, dimnames(JobSat)$satisfaction, sep = ":")
```

A different situation arises with tables where you want to *permute* the levels of one or more variables to arrange them in a more convenient order without changing their labels. For example, in the `HairEyeColor` table, hair color and eye color are ordered arbitrarily. For visualizing the data using mosaic plots and other methods described later, it turns out to be more useful to assure that both hair color and eye color are ordered from dark to light. Hair colors are actually ordered this way already: "Black", "Brown", "Red", "Blond". But eye colors are ordered as "Brown", "Blue", "Hazel", "Green". It is easiest to re-order the eye colors by indexing the columns (dimension 2) in this array to a new order, "Brown", "Hazel", "Green", "Blue", giving the indices of the old levels in the new order (here: 1,3,4,2). Again `str()` is your friend, showing the structure of the result to check that the result is what you want.

```
> data(HairEyeColor, package = "datasets")
> HEC <- HairEyeColor[, c(1, 3, 4, 2), ]
> str(HEC)

num [1:4, 1:4, 1:2] 32 53 10 3 10 25 7 5 3 15 ...
- attr(*, "dimnames")=List of 3
..$ Hair: chr [1:4] "Black" "Brown" "Red" "Blond"
..$ Eye : chr [1:4] "Brown" "Hazel" "Green" "Blue"
..$ Sex : chr [1:2] "Male" "Female"
```

Finally, there are situations where, particularly for display purposes, you want to re-order the *dimensions* of an *n*-way table, and/or change the labels for the variables or levels. This is easy when the data are in table form: `aperm()` permutes the dimensions, and assigning to `names` and `dimnames` changes variable names and level labels respectively.

```
> str(UCBAdmissions)
```

```

table [1:2, 1:2, 1:6] 512 313 89 19 353 207 17 8 120 205 ...
- attr(*, "dimnames")=List of 3
..$ Admit : chr [1:2] "Admitted" "Rejected"
..$ Gender: chr [1:2] "Male" "Female"
..$ Dept   : chr [1:6] "A" "B" "C" "D" ...
.

> UCB <- aperm(UCBAdmissions, c(2, 1, 3))
> dimnames(UCB)$Admit <- c("Yes", "No")
> names(dimnames(UCB)) <- c("Sex", "Admitted", "Department")
> str(UCB)

table [1:2, 1:2, 1:6] 512 89 313 19 353 17 207 8 120 202 ...
- attr(*, "dimnames")=List of 3
..$ Sex      : chr [1:2] "Male" "Female"
..$ Admitted : chr [1:2] "Yes" "No"
..$ Department: chr [1:6] "A" "B" "C" "D" ...
.
```

2.4 Generating tables with `table()` and `xtabs()`

{sec:table}

With data in case form or frequency form, you can generate frequency tables from factor variables in data frames using the `table()` function; for tables of proportions, use the `prop.table()` function, and for marginal frequencies (summing over some variables) use `margin.table()`. The examples below use the same case-form data frame `mydata` used earlier (Section 2.1.4).

```

> set.seed(12345)    # reproducibility
> n <- 100
> A <- factor(sample(c("a1", "a2"), n, replace = TRUE))
> B <- factor(sample(c("b1", "b2"), n, replace = TRUE))
> sex <- factor(sample(c("M", "F"), n, replace = TRUE))
> age <- round(rnorm(n, mean = 30, sd = 5))
> mydata <- data.frame(A, B, sex, age)

```

2.4.1 `table()`

{sec:table}

`table(...)` takes a list of variables interpreted as factors, or a data frame whose columns are so interpreted. It does not take a `data=` argument, so either supply the names of columns in the data frame (possibly using `with()` for convenience), or select the variables using column indexes:

```

> # 2-Way Frequency Table
> table(mydata$A, mydata$B)           # A will be rows, B will be columns

      b1  b2
a1 18 30
a2 22 30

> ## same: with(mydata, table(A, B))
> (mytab <- table(mydata[,1:2]))       # same

      B
A    b1  b2
a1 18 30
a2 22 30

```

We can use `margin.table(X, margin)` to sum a table `X` for the indices in `margin`, i.e., over the dimensions not included in `margin`. A related function is `addmargins(X, margin)`,

`FUN = sum`), which extends the dimensions of a table or array with the marginal values calculated by `FUN`.

```
> margin.table(mytab)           # sum over A & B
[1] 100

> margin.table(mytab, 1)       # A frequencies (summed over B)
A
a1 a2
48 52

> margin.table(mytab, 2)       # B frequencies (summed over A)
B
b1 b2
40 60

> addmargins(mytab)          # show all marginal totals

      B
A      b1   b2 Sum
  a1    18   30  48
  a2    22   30  52
  Sum   40   60 100
```

The function `prop.table()` expresses the table entries as a fraction of a given marginal table.

```
> prop.table(mytab)           # cell proportions

      B
A      b1   b2
  a1 0.18 0.30
  a2 0.22 0.30

> prop.table(mytab, 1)       # row proportions

      B
A      b1      b2
  a1 0.37500 0.62500
  a2 0.42308 0.57692

> prop.table(mytab, 2)       # column proportions

      B
A      b1   b2
  a1 0.45 0.50
  a2 0.55 0.50
```

`table()` can also generate multidimensional tables based on 3 or more categorical variables. In this case, use the `ftable()` or `structable()` function to print the results more attractively as a “flat” (2-way) table.

TODO: DM: is there any advantage of `ftable()` over `structable()`? I think not, so just use `structable()`?

```
> # 3-Way Frequency Table
> mytab <- table(mydata[,c("A", "B", "sex")])
> ftable(mytab)

  sex   F   M
```

A	B
a1	b1 9 9
	b2 15 15
a2	b1 12 10
	b2 19 11

`table()` ignores missing values by default, but has optional arguments `useNA` and `exclude` that can be used to control this. See `help(table)` for the details.

2.4.2 `xtabs()`

{sec:xtabs}

The `xtabs()` function allows you to create cross tabulations of data using formula style input. This typically works with case-form or frequency-form data supplied in a data frame or a matrix. The result is a contingency table in array format, whose dimensions are determined by the terms on the right side of the formula. As shown below, the `summary` method for tables produces a simple χ^2 test of independence of all factors, and indicates the number of cases and dimensions.

```
> # 3-Way Frequency Table
> mytable <- xtabs(~ A + B + sex, data = mydata)
> ftable(mytable)      # print table

      sex   F   M
A   B
a1 b1      9  9
    b2     15 15
a2 b1     12 10
    b2     19 11

> summary(mytable)    # chi-squared test of independence

Call: xtabs(formula = ~A + B + sex, data = mydata)
Number of cases in table: 100
Number of factors: 3
Test for independence of all factors:
Chisq = 1.54, df = 4, p-value = 0.82
```

When the data have already been tabulated in frequency form, include the frequency variable (usually `count` or `Freq`) on the left side of the formula, as shown in the example below for the GSS data.

```
> (GSStab <- xtabs(count ~ sex + party, data = GSS))

      party
sex      dem indep rep
female  279    73 225
male    165    47 191

> summary(GSStab)

Call: xtabs(formula = count ~ sex + party, data = GSS)
Number of cases in table: 980
Number of factors: 2
Test for independence of all factors:
Chisq = 7, df = 2, p-value = 0.03
```

For "table" objects, the `plot` method produces basic mosaic plots using the `mosaicplot()` function.

2.5 Printing tables with `structable()` and `ftable()`

{sec:structable}

2.5.1 Text output

For 3-way and larger tables, the functions `ftable()` (in the `stats` package) and `structable()` (in `vcd`) provide a convenient and flexible tabular display in a “flat” (2-way) format.

With `ftable(X, row.vars = , col.vars =)`, variables assigned to the rows and/or columns of the result can be specified as the integer numbers or character names of the variables in the array `X`. By default, the last variable is used for the columns. The formula method, in the form `ftable(colvars ~ rowvars, data)` allows a formula, where the left and right hand side of formula specify the column and row variables respectively.

```
> ftable(UCB) # default
      Department   A   B   C   D   E   F
Sex Admitted
Male Yes          512 353 120 138  53  22
      No           313 207 205 279 138 351
Female Yes         89  17 202 131  94  24
      No          19   8 391 244 299 317

> #ftable(UCB, row.vars = 1:2) # same result
> ftable(Admitted + Sex ~ Department, data = UCB) # formula method
      Admitted Yes      No
      Sex     Male Female Male Female
Department
A             512     89  313     19
B             353     17  207      8
C             120     202 205    391
D             138     131 279    244
E              53     94 138    299
F              22     24 351    317
```

The `structable()` function is similar, but more general, and uses recursive splits in the vertical or horizontal directions (similar to the construction of mosaic displays). It works with both data frames and table objects.

```
> structable(HairEyeColor) # show the table: default
      Eye Brown Blue Hazel Green
Hair Sex
Black Male     32    11    10     3
      Female    36     9     5     2
Brown Male    53    50    25    15
      Female   66    34    29    14
Red   Male    10    10     7     7
      Female   16     7     7     7
Blond Male    3    30     5     8
      Female   4    64     5     8

> structable(Hair + Sex ~ Eye, HairEyeColor) # specify col ~ row variables
      Hair Black      Brown      Red      Blond
      Sex   Male Female   Male Female  Male Female
Eye
Brown     32    36    53    66    10    16     3     4
Blue      11     9    50    34    10     7    30    64
Hazel     10     5    25    29     7     7     5     5
Green      3     2    15    14     7     7     8     8
```

It also returns an object of class "structable" for which there are a variety of special methods. For example, the transpose function `t()` interchanges rows and columns, so that a call like `t(structable(HairEyeColor))` produces the second result shown just above. There are also plot methods: for example, `plot()` produces mosaic plots.

2.6 Subsetting data

Often, the analysis of some data set is focused on a subset only. For example, the `HairEyeColor` data set introduced above tabulates frequencies of hair and eye colors for male and female students—the analysis could concentrate on one group only, or compare both groups in a stratified analysis. This section deals with extracting subsets of data in tables, structables or data frames.

2.6.1 Subsetting tables

If data are available in tabular form created with `table()` or `xtabs()`, resulting in table objects, subsetting is done via indexing, either with integers or character strings corresponding to the factor levels. The following code extracts the female data from the `HairEyeColor` data set:

```
> HairEyeColor[, , "Female"]

      Eye
Hair   Brown Blue Hazel Green
  Black    36    9     5    2
  Brown    66   34    29   14
  Red     16     7     7    7
  Blond     4   64     5    8

> ##same using index: HairEyeColor[, , 2]
```

Empty indices stand for taking all data of the corresponding dimension. The third one (Sex) is fixed at the second ("Female") level. Note that in this case, the dimensionality is reduced to a two-way table, since dimensions with only one level are dropped by default. Functions like `apply()` can iterate through all levels of one or several dimensions and apply a function to each subset. The following calculates the total amount of male and female students:

```
> apply(HairEyeColor, 3, sum)

Male Female
 279    313
```

It is of course possible to select more than one level:

```
> HairEyeColor[c("Black", "Brown"), c("Hazel", "Green"), ]

, , Sex = Male

      Eye
Hair   Hazel Green
  Black     10     3
  Brown    25    15

, , Sex = Female

      Eye
Hair   Hazel Green
  Black      5     2
  Brown    29    14
```

2.6.2 Subsetting structables

c:subsettingtables} Structables work in a similar way, but take into account the hierarchical structure imposed by the “flattened” format, and also distinguish explicitly between subsetting levels and subsetting tables. In the following example, compare the different effects of applying the [and [[operators to the structable:

```
> hec <- structable(Eye ~ Sex + Hair, data = HairEyeColor)
> hec

      Eye Brown Blue Hazel Green
Sex   Hair
Male Black     32    11    10     3
      Brown    53    50    25    15
      Red      10    10     7     7
      Blond     3    30     5     8
Female Black    36     9     5     2
      Brown   66    34    29    14
      Red     16     7     7     7
      Blond     4    64     5     8

> hec["Male",]

      Eye Brown Blue Hazel Green
Sex   Hair
Male Black     32    11    10     3
      Brown    53    50    25    15
      Red      10    10     7     7
      Blond     3    30     5     8

> hec[["Male", ]]

      Eye Brown Blue Hazel Green
Hair
Black     32    11    10     3
Brown    53    50    25    15
Red      10    10     7     7
Blond     3    30     5     8
```

The first form keeps the dimensionality, whereas the second conditions on the “Male” level and returns the corresponding subtable. The following does this twice, once for Sex, and once for Hair (restricted to the Male level):

```
> hec[ [ c("Male", "Brown"), ]]

      Eye Brown Blue Hazel Green
      53    50    25    15
```

2.6.3 Subsetting data frames

Data available in data frames (frequency or case form) can also be subsetted, either by using indexes on the rows and/or columns, or, more conveniently, by applying the `subset()` function. The following statement will extract the Treatment and Improved variables for all female patients older than 68:

{sec:subsettingdf}

```
> rows <- Arthritis$Sex == "Female" & Arthritis$Age > 68
> cols <- c("Treatment", "Improved")
> Arthritis[rows, cols]
```

```
Treatment Improved
39   Treated    None
40   Treated    Some
41   Treated    Some
84   Placebo   Marked
```

Note the use of the single & for the logical expression selecting the rows. The same result can be achieved more conveniently using the subset () function, first taking the data set, followed by an expression for selecting the rows (evaluated in the context of the data frame), and then an expression for selecting the columns:

```
> subset(Arthritis, Sex == "Female" & Age > 68, c(Treatment, Improved))

Treatment Improved
39   Treated    None
40   Treated    Some
41   Treated    Some
84   Placebo   Marked
```

Note the non-standard evaluation of c(Treatment, Improved): the meaning of c() is not “combine the two columns into a single vector”, but “select both from the data frame”. Likewise, columns can be removed using – on column names, which is not possible using standard indexing in matrices or data frames:

```
> subset(Arthritis, Sex == "Female" & Age > 68, -c(Age, ID))

Treatment   Sex Improved
39   Treated Female    None
40   Treated Female    Some
41   Treated Female    Some
84   Placebo Female   Marked
```

2.7 Collapsing tables

2.7.1 Collapsing over table factors: aggregate(), margin.table() and apply()

It sometimes happens that we have a data set with more variables or factors than we want to analyse, or else, having done some initial analyses, we decide that certain factors are not important, and so should be excluded from graphic displays by collapsing (summing) over them. For example, mosaic plots and fourfold displays are often simpler to construct from versions of the data collapsed over the factors which are not shown in the plots.

The appropriate tools to use again depend on the form in which the data are represented—a case-form data frame, a frequency-form data frame (aggregate()), or a table-form array or table object (margin.table() or apply()).

When the data are in frequency form, and we want to produce another frequency data frame, aggregate() is a handy tool, using the argument FUN = sum to sum the frequency variable over the factors *not* mentioned in the formula.

EXAMPLE 2.6: Dayton survey

The data frame *DaytonSurvey* in the *vcdExtra* package represents a 2^5 table giving the frequencies of reported use (“ever used?”) of alcohol, cigarettes and marijuana in a sample of 2276 high school seniors, also classified by sex and race.

```
> data(DaytonSurvey, package = "vcdExtra")
> str(DaytonSurvey)

'data.frame': 32 obs. of 6 variables:
 $ cigarette: Factor w/ 2 levels "Yes", "No": 1 2 1 2 1 2 1 2 1 2 ...
 $ alcohol   : Factor w/ 2 levels "Yes", "No": 1 1 2 2 1 1 2 2 1 1 ...
 $ marijuana: Factor w/ 2 levels "Yes", "No": 1 1 1 1 2 2 2 2 1 1 ...
 $ sex       : Factor w/ 2 levels "female", "male": 1 1 1 1 1 1 1 1 2 2 ...
 $ race      : Factor w/ 2 levels "white", "other": 1 1 1 1 1 1 1 1 1 1 ...
 $ Freq       : num  405 13 1 1 268 218 17 117 453 28 ...

> head(DaytonSurvey)

cigarette alcohol marijuana sex race Freq
1 Yes Yes Yes female white 405
2 No Yes Yes female white 13
3 Yes No Yes female white 1
4 No No Yes female white 1
5 Yes Yes No female white 268
6 No Yes No female white 218
```

To focus on the associations among the substances, we want to collapse over sex and race. The right-hand side of the formula used in the call to `aggregate()` gives the factors to be retained in the new frequency data frame, `Dayton_ACM_df`. The left-hand side is the frequency variable (`Freq`), and we aggregate using the `FUN = sum`.

```
> # data in frequency form: collapse over sex and race
> Dayton_ACM_df <- aggregate(Freq ~ cigarette + alcohol + marijuana,
+                               data = DaytonSurvey, FUN = sum)
> Dayton_ACM_df

cigarette alcohol marijuana Freq
1 Yes Yes Yes 911
2 No Yes Yes 44
3 Yes No Yes 3
4 No No Yes 2
5 Yes Yes No 538
6 No Yes No 456
7 Yes No No 43
8 No No No 279
```



When the data are in table form, and we want to produce another table, `apply()` with `FUN = sum` can be used in a similar way to sum the table over dimensions not mentioned in the `MARGIN` argument. `margin.table()` is just a wrapper for `apply()` using the `sum()` function.

{ex:dayton2}

EXAMPLE 2.7: Dayton survey

To illustrate, we first convert the `DaytonSurvey` to a 5-way table using `xtabs()`, giving `Dayton_tab`.

```
> # convert to table form
> Dayton_tab <- xtabs(Freq ~ cigarette + alcohol + marijuana + sex + race,
+                      data = DaytonSurvey)
> structable(cigarette + alcohol + marijuana ~ sex + race, data = Dayton_tab)

cigarette Yes          No
alcohol    Yes          No          Yes          No
marijuana Yes          No          Yes          No          Yes          No
sex        race

           cigarette Yes          No
           alcohol    Yes          No          Yes          No
           marijuana Yes          No          Yes          No          Yes          No
           sex        race
```

female white	405	268	1	17	13	218	1	117
other	23	23	0	1	2	19	0	12
male white	453	228	1	17	28	201	1	133
other	30	19	1	8	1	18	0	17

Then, use `apply()` on `Dayton_tab` to give the 3-way table `Dayton_ACM_tab` summed over sex and race. The elements in this new table are the column sums for `Dayton.tab` shown by `structable()` just above.

```
> # collapse over sex and race
> Dayton_ACM_tab <- apply(Dayton_tab, MARGIN = 1:3, FUN = sum)
> Dayton_ACM_tab <- margin.table(Dayton_tab, 1:3)    # same result
> structable(cigarette + alcohol ~ marijuana, data = Dayton_ACM_tab)

      cigarette Yes      No
      alcohol   Yes  No Yes  No
marijuana
Yes           911   3 44   2
No            538  43 456 279
```



(Note that `structable()` would do the collapsing job for us anyway.)

TODO: DM: plyr seems too much here -> remove?

Many of these operations can be performed using the `**ply()` functions in the `plyr` package. For example, with the data in a frequency form data frame, use `ddply()` to collapse over unmentioned factors, and `summarise()` as the function to be applied to each piece.

```
> Dayton_ACM_df <- ddply(DaytonSurvey, .(cigarette, alcohol, marijuana),
+                           summarise, Freq = sum(Freq))
```

2.7.2 Collapsing table levels: `collapse.table()`

A related problem arises when we have a table or array and for some purpose we want to reduce the number of levels of some factors by summing subsets of the frequencies. For example, we may have initially coded Age in 10-year intervals, and decide that, either for analysis or display purposes, we want to reduce Age to 20-year intervals. The `collapse.table()` function in `vcdExtra` was designed for this purpose.

EXAMPLE 2.8: Collapsing categories

Create a 3-way table, and collapse Age from 10-year to 20-year intervals and Education from three levels to two. To illustrate, we first generate a $2 \times 6 \times 3$ table of random counts from a Poisson distribution with mean of 100, with factors sex, age and education.

```
> # create some sample data in frequency form
> set.seed(12345)    # reproducibility
> sex <- c("Male", "Female")
> age <- c("10-19", "20-29", "30-39", "40-49", "50-59", "60-69")
> education <- c("low", "med", "high")
> dat <- expand.grid(sex = sex, age = age, education = education)
> counts <- rpois(36, 100)    # random Poisson cell frequencies
> dat <- cbind(dat, counts)
> # make it into a 3-way table
> tab1 <- xtabs(counts ~ sex + age + education, data = dat)
> structable(tab1)
```

		age	10-19	20-29	30-39	40-49	50-59	60-69
sex	education							
Male	low		105	98	123	97	95	105
	med		74	113	114	82	95	85
	high		121	116	104	103	89	100
Female	low		107	95	105	116	103	92
	med		96	88	93	118	99	108
	high		120	102	96	103	127	84

Now collapse age to 20-year intervals, and education to 2 levels. In the arguments to `collapse.table()`, levels of age and education given the same label are summed in the resulting smaller table.

```
> # collapse age to 3 levels, education to 2 levels
> tab2 <- collapse.table(tab1,
+   age = c("10-29", "10-29", "30-49", "30-49", "50-69", "50-69"),
+   education = c("<high", "<high", "high"))
> structable(tab2)

          age 10-29 30-49 50-69
sex   education
Male  <high      390    416    380
      high       237    207    189
Female <high      386    432    402
      high       222    199    211
```



TODO: DM: Add section on subsetting tables using `subset()`, and the `[` and `[[` methods for tables and structables. These skills are required in some examples.

2.8 Converting among frequency tables and data frames

As we've seen, a given contingency table can be represented equivalently in case form, frequency form and table form. However, some R functions were designed for one particular representation. Table 2.1 gives an overview on some handy tools (with sketched usage) for converting from one form to another, discussed below.

{sec:convert}

Table 2.1: Tools for converting among different forms for categorical data

{tab:convert}

From this	Case form	Frequency form	To this
Case form	—	Z <- xtabs(~ A + B) as.data.frame(Z)	table(A, B)
Frequency form	expand.dft(X)	—	xtabs(count ~ A + B)
Table form	expand.dft(X)	as.data.frame(X)	—

2.8.1 Table form to frequency form

A contingency table in table form (an object of class "table") can be converted to a data frame in frequency form with `as.data.frame()`.⁶ The resulting data frame contains columns repre-

⁶Because R is object-oriented, this is actually a short-hand for the function `as.data.frame.table()`.

senting the classifying factors and the table entries (as a column named by the `responseName` argument, defaulting to `Freq`. The function `as.data.frame()` is the inverse of `xtabs()`, which converts a data frame to a table.

EXAMPLE 2.9: General social survey

Convert the GSStab in table form to a data.frame in frequency form. By default, the frequency variable is named `Freq`, and the variables `sex` and `party` are made factors.

```
> as.data.frame(GSStab)

  sex party Freq
1 female  dem  279
2 male   dem  165
3 female indep  73
4 male   indep  47
5 female rep  225
6 male   rep  191
```



In addition, there are situations where numeric table variables are represented as factors, but you need to convert them to numerics for calculation purposes.

EXAMPLE 2.10: Death by horse kick

For example, We might want to calculate the weighted mean of `nDeaths` in the `HorseKicks` data. Using `as.data.frame()` won't work here, because the variable `nDeaths` becomes a factor.

```
> str(as.data.frame(HorseKicks))

'data.frame': 5 obs. of  2 variables:
 $ nDeaths: Factor w/ 5 levels "0","1","2","3",...: 1 2 3 4 5
 $ Freq    : int  109 65 22 3 1
```

One solution is to use `data.frame()` directly and `as.numeric()` to coerce the table names to numbers.

```
> horse.df <- data.frame(nDeaths = as.numeric(names(HorseKicks)),
+                           Freq = as.vector(HorseKicks))
> str(horse.df)

'data.frame': 5 obs. of  2 variables:
 $ nDeaths: num  0 1 2 3 4
 $ Freq    : int  109 65 22 3 1

> horse.df

  nDeaths Freq
1         0 109
2         1  65
3         2  22
4         3   3
5         4   1
```

Then, `weighted.mean()` works as we would like:

```
> weighted.mean(horse.df$nDeaths, weights=horse.df$Freq)

[1] 2
```



2.8.2 Case form to table form

{ex:Arth-convert} Going the other way, we use `table()` to convert from case form to table form.

EXAMPLE 2.11: Arthritis treatment

Convert the *Arthritis* data in case form to a 3-way table of Treatment × Sex × Improved.

We select the desired columns with their names, but could also use column numbers, e.g., `table(Arthritis[, c(2, 3, 5)])`

```
> Art.tab <- table(Arthritis[, c("Treatment", "Sex", "Improved")])
> str(Art.tab)

'table' int [1:2, 1:2, 1:3] 19 6 10 7 7 5 0 2 6 16 ...
- attr(*, "dimnames")=List of 3
..$ Treatment: chr [1:2] "Placebo" "Treated"
..$ Sex       : chr [1:2] "Female" "Male"
..$ Improved  : chr [1:3] "None" "Some" "Marked"

> ftable(Art.tab)

          Improved None Some Marked
Treatment Sex
Placebo   Female      19     7      6
           Male       10     0      1
Treated   Female      6      5     16
           Male       7      2      5
```



2.8.3 Table form to case form

There may also be times that you will need an equivalent case form data frame with factors representing the table variables rather than the frequency table. For example, the `mca()` function in package **MASS** (for multiple correspondence analysis) only operates on data in this format. The function `expand.dft()`⁷ in **vcdExtra** does this, converting a table into a case form.

{ex:Arth-convert2}

EXAMPLE 2.12: Arthritis treatment

Convert the *Arthritis* data in table form (`Art.tab`) back to a `data.frame` in case form, with factors Treatment, Sex and Improved.

```
> Art.df <- expand.dft(Art.tab)
> str(Art.df)

'data.frame': 84 obs. of  3 variables:
 $ Treatment: Factor w/ 2 levels "Placebo", "Treated": 1 1 1 1 1 1 1 1 1 1 ...
 $ Sex       : Factor w/ 2 levels "Female", "Male": 1 1 1 1 1 1 1 1 1 1 ...
 $ Improved  : Factor w/ 3 levels "Marked", "None", ...: 2 2 2 2 2 2 2 2 2 2 ...
```



2.8.4 Publishing tables to L^AT_EX or HTML

OK, you've read your data into R, done some analysis, and now want to include some tables in a L^AT_EX document or in a web page in HTML format. Formatting tables for these purposes is often tedious and error-prone.

There are a great many packages in R that provide for nicely formatted, publishable tables for a wide variety of purposes; indeed, most of the tables in this book are generated using these tools.

⁷The original code for this function was provided by Marc Schwarz on the R-Help mailing list.

See Leifeld (2013) for description of the `texreg` package and a comparison with some of the other packages.

Here, we simply illustrate the `xtable` package, which, along with capabilities for statistical model summaries, time-series data, and so forth, has a `xtable.table` method for one-way and two-way table objects.

The `HorseKicks` data is a small one-way frequency table described in Example ?? and containing the frequencies of 0, 1, 2, 3, 4 deaths per corps-year by horse-kick among soldiers in 20 corps in the Prussian army.

```
> data(HorseKicks, package = "vcd")
> HorseKicks

nDeaths
 0   1   2   3   4
109  65  22   3   1
```

By default, `xtable()` formats this in `LATEX` as a vertical table, and prints the `LATEX` markup to the `R` console. This output is shown below (without the usual `##` comment used to indicate `R` output).

```
> library(xtable)
> xtable(HorseKicks)

% latex table generated in R 3.1.2 by xtable 1.7-1 package
% Tue Jan 13 14:07:13 2015
\begin{table}[ht]
\centering
\begin{tabular}{rr}
\hline
& nDeaths \\
\hline
0 & 109 \\
1 & 65 \\
2 & 22 \\
3 & 3 \\
4 & 1 \\
\hline
\end{tabular}
\end{table}
```

When this is rendered in a `LATEX` document, the result of `xtable()` appears as shown in the table below.

```
> xtable(HorseKicks)
```

	nDeaths
0	109
1	65
2	22
3	3
4	1

The table above isn't quite right, because the column label “nDeaths” belongs to the first column, and the second column should be labeled “Freq”. To correct that, we convert the `HorseKicks` table to a data frame (see Section 2.8 for details), add the appropriate `colnames`, and use the `xtable.print` method to supply some other options.

```
> tab <- as.data.frame(HorseKicks)
> colnames(tab) <- c("nDeaths", "Freq")
> print(xtable(tab), include.rownames = FALSE, include.colnames = TRUE)
```

	nDeaths	Freq
0	109	
1	65	
2	22	
3	3	
4	1	

TODO: DM: The code looks quite complicated and the result is not satisfying (no separator between header and table) - maybe too confusing?

Finally, in Chapter 3, we display a number of similar one-way frequency tables in a transposed form to save display space. Table 3.3 is the finished version we show there. The code below uses the following techniques: (a) addmargins() is used to show the sum of all the frequency values; (b) t() transposes the data frame to have 2 rows; (c) rownames() assigns the labels we want for the rows; (d) using the caption argument provides a table caption, and a numbered table in L^AT_EX; (d) column alignment ("r" or "l") for the table columns is computed as a character string used for the align argument.

```
> horsetab <- t(as.data.frame(addmargins(HorseKicks)))
> rownames(horsetab) <- c( "Number of deaths", "Frequency" )
> horsetab <- xtable(horsetab, digits = 0,
+   caption = "von Bortkiewicz's data on deaths by horse kicks",
+   align = paste0("l|", paste(rep("r", ncol(horsetab)), collapse = " ")))
+ )
> print(horsetab, include.colnames = FALSE)
```

Number of deaths	0	1	2	3	4	Sum
Frequency	109	65	22	3	1	200

Table 2.2: von Bortkiewicz's data on deaths by horse kicks

For use in a web page, blog, or Word document, you can use type="HTML" in the call to print() for "xtable" objects. **Done:** DM: also show HTML version? — MF: added note above

2.9 A complex example: TV viewing data

If you have followed so far, congratulations! You are ready for a more complicated example that puts together a variety of the skills developed in this chapter: (a) reading raw data, (b) creating tables, (c) assigning level names to factors and (d) collapsing levels or variables for use in analysis.

For an illustration of these steps, we use the dataset tv.dat, supplied with the initial implementation of mosaic displays in R by Jay Emerson. In turn, they were derived from an early, compelling example of mosaic displays (Hartigan and Kleiner, 1984), that illustrated the method with data on a large sample of TV viewers whose behavior had been recorded for the Nielsen ratings. This data set contains sample television audience data from Nielsen Media Research for the week starting November 6, 1995.

{sec:working-complex}

The data file, `tv.dat` is stored in frequency form as a file with 825 rows and 5 columns. There is no header line in the file, so when we use `read.table()` below, the variables will be named V1 – V5. This data represents a 4-way table of size $5 \times 11 \times 5 \times 3 = 825$ where the table variables are V1 – V4, and the cell frequency is read as V5.

The table variables are:

- V1– values 1:5 correspond to the days Monday–Friday;
 - V2– values 1:11 correspond to the quarter hour times 8:00PM through 10:30PM;
 - V3– values 1:5 correspond to ABC, CBS, NBC, Fox, and non-network choices;
 - V4– values 1:3 correspond to transition states: turn the television Off, Switch channels, or Persist in viewing the current channel.

2.9.1 Creating data frames and arrays

The file `tv.dat` is stored in the `doc/extdata` directory of `vcdExtra`; it can be read as follows:

```
> tv_data <- read.table(system.file("doc", "extdata", "tv.dat", package = "vcdExtra"))
> str(tv_data)

'data.frame': 825 obs. of 5 variables:
 $ V1: int 1 2 3 4 5 1 2 3 4 5 ...
 $ V2: int 1 1 1 1 1 2 2 2 2 2 ...
 $ V3: int 1 1 1 1 1 1 1 1 1 1 ...
 $ V4: int 1 1 1 1 1 1 1 1 1 1 ...
 $ V5: int 6 18 6 2 11 6 29 25 17 29 ...

> head(tv_data, 5)

  V1 V2 V3 V4 V5
1  1  1  1  1  6
2  2  1  1  1 18
3  3  1  1  1  6
4  4  1  1  1  2
5  5  1  1  1 11
```

To read such data from a local file, just use `read.table()` in this form:

```
> tv_data <- read.table("C:/R/data/tv.dat")
```

or

```
> tv_data <- file.choose()
```

to select the path using the file chooser tool.

We could use this data in frequency form for analysis by renaming the variables, and converting the integer-coded factors V1 – V4 to R factors. The lines below use the function `within()` to avoid having to use `TV.dat$Day <- factor(TV.dat$Day)` etc., and only supplies labels for the first variable.

Alternatively, we could just reshape the frequency column (`V5` or `tv_data[, 5]`) into a 4-way array. In the lines below, we rely on the facts that the (a) the table is complete—there are no missing cells, so `nrow(tv_data)=825`; (b) the observations are ordered so that `V1` varies most rapidly and `V4` most slowly. From this, we can just extract the frequency column and reshape it into an array using the `dim` argument. The level names are assigned to `dimnames(TV)` and the variable names to `names(dimnames(TV))`.

```
> TV <- array(tv_data[, 5], dim = c(5, 11, 5, 3))
> dimnames(TV) <-
+   list(c("Mon", "Tue", "Wed", "Thu", "Fri"),
+         c("8:00", "8:15", "8:30", "8:45", "9:00", "9:15", "9:30",
+           "9:45", "10:00", "10:15", "10:30"),
+         c("ABC", "CBS", "NBC", "Fox", "Other"),
+         c("Off", "Switch", "Persist"))
> names(dimnames(TV)) <- c("Day", "Time", "Network", "State")
```

More generally (even if there are missing cells), we can use `xtabs()` to do the cross-tabulation, using `V5` as the frequency variable. Here's how to do this same operation with `xtabs()`:

```
> TV <- xtabs(V5 ~ ., data = tv_data)
> dimnames(TV) <-
+   list(Day = c("Mon", "Tue", "Wed", "Thu", "Fri"),
+         Time = c("8:00", "8:15", "8:30", "8:45", "9:00", "9:15", "9:30",
+                  "9:45", "10:00", "10:15", "10:30"),
+         Network = c("ABC", "CBS", "NBC", "Fox", "Other"),
+         State = c("Off", "Switch", "Persist"))
```

Note that in the lines above, the variable names are assigned directly as the names of the elements in the `dimnames` list.

2.9.2 Subsetting and collapsing

For many purposes, the 4-way table `TV` is too large and awkward to work with. Among the networks, `Fox` and `Other` occur infrequently, so we will remove them. We can also cut it down to a 3-way table by considering only viewers who persist with the current station.⁸

```
> TV <- TV[, 1:3, ]      # keep only ABC, CBS, NBC
> TV <- TV[, , 3]        # keep only Persist -- now a 3 way table
> structable(TV)

      Time 8:00 8:15 8:30 8:45 9:00 9:15 9:30 9:45 10:00 10:15 10:30
Day Network
Mon ABC      146  151  156   83  325  350  386  340  352  280  278
          CBS    337  293  304  233  311  251  241  164  252  265  272
          NBC    263  219  236  140  226  235  239  246  279  263  283
Tue ABC      244  181  231  205  385  283  345  192  329  351  364
          CBS    173  180  184  109  218  235  256  250  274  263  261
          NBC    315  254  280  241  370  214  195  111  188  190  210
Wed ABC      233  161  194  156  339  264  279  140  237  228  203
          CBS    158  126  207  59   98  103  122  86   109  105  110
          NBC    134  146  166  66   194  230  264  143  274  289  306
Thu ABC      174  183  197  181  187  198  211  86   110  122  117
          CBS    196  185  195  104  106  116  116  47   102  84   84
          NBC    515  463  472  477  590  473  446  349  649  705  747
Fri ABC      294  281  305  239  278  246  245  138  246  232  233
          CBS    130  144  154  81   129  153  136  126  138  136  152
          NBC    195  220  248  160  172  164  169  85   183  198  204
```

Finally, for some purposes, we might also want to collapse the 11 times into a smaller number. Here, we use `collapse.table()` (see Section 2.7.2), which was designed for this purpose.

⁸This relies on the fact that indexing an array drops dimensions of length 1 by default, using the argument `drop = TRUE`; the result is coerced to the lowest possible dimension.

```
> TV2 <- collapse.table(TV,
+                         Time = c(rep("8:00-8:59", 4),
+                                 rep("9:00-9:59", 4),
+                                 rep("10:00-10:44", 3)))
> structable(Day ~ Time + Network, TV2)

      Day   Mon   Tue   Wed   Thu   Fri
Time   Network
8:00-8:59 ABC      536  861  744  735 1119
          CBS     1167  646  550  680  509
          NBC     858 1090  512 1927  823
9:00-9:59 ABC     1401 1205 1022  682  907
          CBS     967  959  409  385  544
          NBC     946  890  831 1858  590
10:00-10:44 ABC    910 1044  668  349  711
          CBS    789  798  324  270  426
          NBC    825  588  869 2101  585
```

Congratulations! If you followed the operations described above, you are ready for the material described in the rest of the book. If not, try working through some of exercises below.

2.10 Lab exercises

Exercise 2.1 The packages `vcd` and `vcdExtra` contain many data sets with some examples of analysis and graphical display. The goal of this exercise is to familiarize yourself with these resources. You can get a brief summary of these using the function `datasets()`. Use the following to get a list of these with some characteristics and titles.

```
> ds <- datasets(package=c("vcd", "vcdExtra"))
> str(ds)

'data.frame': 70 obs. of  5 variables:
 $ Package: chr  "vcd" "vcd" "vcd" ...
 $ Item    : chr  "Arthritis" "Baseball" "BrokenMarriage" "Bundesliga" ...
 $ class   : chr  "data.frame" "data.frame" "data.frame" "data.frame" ...
 $ dim     : chr  "84x5" "322x25" "20x4" "14018x7" ...
 $ Title   : chr  "Arthritis Treatment Data" "Baseball Data" "Broken Marriage Data" "Ergebnisse der Fussball"
```

- (a) How many data sets are there altogether? How many are there in each package?
- (b) Make a tabular display of the frequencies by `Package` and `class`.
- (c) Choose one or two data sets from this list, and examine their help files (e.g., `help(Arthritis)` or `?Arthritis`). You can use, e.g., `example(Arthritis)` to run the R code for a given example.

{lab:2.2}

Exercise 2.2 The data set `UCBAdmissions` is a 3-way table of frequencies classified by Admit, Gender and Dept.

- (a) Find the total number of cases contained in this table.
- (b) For each department, find the total number of applicants.
- (c) For each department, find the overall proportion of applicants who were admitted.
- (d) Construct a tabular display of department (rows) and gender (columns), showing the proportion of applicants in each cell who were admitted.

{lab:2.3}

Exercise 2.3 The data set `DanishWelfare` in `vcd` gives a 4-way, $3 \times 4 \times 3 \times 5$ table as a data frame in frequency form, containing the variable `Freq` and four factors, `Alcohol`, `Income`, `Status` and `Urban`. The variable `Alcohol` can be considered as the response variable, and the others as possible predictors.

- Find the total number of cases represented in this table.
- In this form, the variables `Alcohol` and `Income` should arguably be considered *ordered* factors. Change them to make them ordered.
- Convert this data frame to table form, `DanishWelfare.tab`, a 4-way array containing the frequencies with appropriate variable names and level names.
- The variable `Urban` has 5 categories. Find the total frequencies in each of these. How would you collapse the table to have only two categories, `City`, `Non-city`?
- Use `structable()` or `ftable()` to produce a pleasing flattened display of the frequencies in the 4-way table. Choose the variables used as row and column variables to make it easier to compare levels of `Alcohol` across the other factors.

{lab:2.4}

Exercise 2.4 The data set `UKSoccer` in `vcd` gives the distributions of number of goals scored by the 20 teams in the 1995/96 season of the Premier League of the UK Football Association.

```
> data(UKSoccer, package="vcd")
> ftable(UKSoccer)

      Away   0   1   2   3   4
Home
0       27  29  10   8   2
1       59  53  14  12   4
2       28  32  14  12   4
3       19  14   7   4   1
4        7   8  10   2   0
```

This two-way table classifies all $20 \times 19 = 380$ games by the joint outcome (Home, Away), the number of goals scored by the Home and Away teams. The value 4 in this table actually represents 4 or more goals.

- Verify that the total number of games represented in this table is 380.
- Find the marginal total of the number of goals scored by each of the home and away teams.
- Express each of the marginal totals as proportions.
- Comment on the distribution of the numbers of home-team and away-team goals. Is there any evidence that home teams score more goals on average?

{lab:2.5}

Exercise 2.5 The one-way frequency table, `Saxony` in `vcd` records the frequencies of families with 0, 1, 2, ... 12 male children, among 6115 families with 12 children. This data set is used extensively in Chapter 3.

```
> data(Saxony, package="vcd")
> Saxony

nMales
  0    1    2    3    4    5    6    7    8    9    10   11   12
  3   24  104  286  670 1033 1343 1112  829  478  181   45    7
```

Another data set, `Geissler`, in the `vcdExtra` package, gives the complete tabulation of all combinations of `boys` and `girls` in families with a given total number of children `size`. The task here is to create an equivalent table, `Saxony12` from the `Geissler` data.

```
> data(Geissler, package="vcdExtra")
> str(Geissler)

'data.frame': 90 obs. of  4 variables:
 $ boys : int  0 0 0 0 0 0 0 0 0 ...
 $ girls: num  1 2 3 4 5 6 7 8 9 10 ...
 $ size : num  1 2 3 4 5 6 7 8 9 10 ...
 $ Freq : int  108719 42860 17395 7004 2839 1096 436 161 66 30 ...
```

- (a) Use `subset()` to create a data frame, `sax12` containing the *Geissler* observations in families with `size==12`.
- (b) Select the columns for `boys` and `Freq`.
- (c) Use `xtabs()` with a formula, `Freq ~ boys`, to create the one-way table.
- (d) Do the same steps again, to create a one-way table, `Saxony11` containing similar frequencies for families of `size==11`.

{lab:2.6}

Exercise 2.6 **Interactive coding of table factors:* Some statistical and graphical methods for contingency tables are implemented only for two-way tables, but can be extended to 3+-way tables by recoding the factors to interactive combinations along the rows and/or columns, in a way similar to what `ftable()` and `structable()` do for printed displays.

For the *UCBAdmissions* data, produce a two-way table object, `UCB.tab2`, that has the combinations of `Admit` and `Gender` as the rows, and `Dept` as its columns, to look like the result below:

	Dept					
Admit:Gender	A	B	C	D	E	F
Admitted:Female	89	17	202	131	94	24
Admitted:Male	512	353	120	138	53	22
Rejected:Female	19	8	391	244	299	317
Rejected:Male	313	207	205	279	138	351

- (a) Try this the long way: convert *UCBAdmissions* to a data frame (`as.data.frame()`), manipulate the factors (e.g., `interaction()`), then convert back to a table (`as.data.frame()`).
- (b) Try this the short way: both `ftable()` and `structable()` have `as.matrix()` methods that convert their result to a matrix.

{lab:2.7}

Exercise 2.7 The data set *VisualAcuity* in *vcd* gives $4 \times 4 \times 2$ table as a frequency data frame.

```
> data("VisualAcuity", package="vcd")
> str(VisualAcuity)

'data.frame': 32 obs. of 4 variables:
 $ Freq : num 1520 234 117 36 266 ...
 $ right : Factor w/ 4 levels "1","2","3","4": 1 2 3 4 1 2 3 4 1 2 ...
 $ left  : Factor w/ 4 levels "1","2","3","4": 1 1 1 1 2 2 2 2 3 3 ...
 $ gender: Factor w/ 2 levels "male","female": 2 2 2 2 2 2 2 2 2 2 ...
```

- (a) From this, use `xtabs()` to create two 4×4 frequency tables, one for each gender.
- (b) Use `structable()` to create a nicely organized tabular display.
- (c) Use `xtable()` to create a L^AT_EX or HTML table.

Chapter 3

Fitting and graphing discrete distributions

{ch:discrete}

Discrete data often follow various theoretical probability models. Graphic displays are used to visualize goodness of fit, to diagnose an appropriate model, and determine the impact of individual observations on estimated parameters.

Not everything that counts can be counted, and not everything that can be counted counts.

Albert Einstein

Discrete frequency distributions often involve counts of occurrences of events, such as accident fatalities, incidents of terrorism or suicide, words in passages of text, or blood cells with some characteristic. Often interest is focused on how closely such data follow a particular probability distribution, such as the binomial, Poisson, or geometric distribution, which provide the basis for generating mechanisms that might give rise to the data. Understanding and visualizing such distributions in the simplest case of an unstructured sample provides a building block for generalized linear models (Chapter 9) where they serve as one component. They also provide the basis for a variety of recent extensions of regression models for count data (Chapter 9), allowing excess counts of zeros (zero-inflated models), left- or right-truncation often encountered in statistical practice.

TODO: DM: Fix chapter ref if ch.09 gets split

This chapter describes the well-known discrete frequency distributions: the binomial, Poisson, negative binomial, geometric, and logarithmic series distributions in the simplest case of an unstructured sample. The chapter begins with simple graphical displays (line graphs and bar charts) to view the distributions of empirical data and theoretical frequencies from a specified discrete distribution.

It then describes methods for fitting data to a distribution of a given form and simple, effective graphical methods than can be used to visualize goodness of fit, to diagnose an appropriate model (e.g., does a given data set follow the Poisson or negative binomial?) and determine the impact of individual observations on estimated parameters.

3.1 Introduction to discrete distributions

{sec:discrete-intro}

Discrete data analysis is concerned with the study of the tabulation of one or more types of events, often categorized into mutually exclusive and exhaustive categories. **Binary events** having two

outcome categories include the toss of a coin (head/tails), sex of a child (male/female), survival of a patient following surgery (lived/died), and so forth. **Polytomous events** have more outcome categories, which may be *ordered* (rating of impairment: low/medium/high, by a physician) and possibly numerically-valued (number of dots (pips), 1–6 on the toss of a die) or *unordered* (political party supported: Liberal, Conservative, Greens, Socialist).

In this chapter, we focus largely on one-way frequency tables for a single numerically-valued variable. Probability models for such data provide the opportunity to describe or explain the *structure* in such data, in that they entail some data generating mechanism and provide the basis for testing scientific hypotheses, prediction of future results. If a given probability model does not fit the data, this can often be a further opportunity to extend understanding of the data or the underlying substantive theory or both.

The remainder of this section gives a few substantive examples of situations where the well-known discrete frequency distributions (binomial, Poisson, negative binomial, geometric, and logarithmic series) might reasonably apply, at least approximately. The mathematical characteristics and properties of these theoretical distributions are postponed to Section 3.2.

In many cases, the data at hand pertain to two types of variables in a one-way frequency table. There is a basic outcome variable, k , taking integer values, $k = 0, 1, \dots$, and called a **count**. For each value of k , we also have a **frequency**, n_k that the count k was observed in some sample. For example, in the study of children in families, the count variable k could be the total number of children or the number of male children; the frequency variable, n_k , would then give the number of families with that basic count k .

3.1.1 Binomial data

{sec:binom-data}

Binomial type data arise as the discrete distribution of the number of “success” events in n independent binary trials, each of which yields a success (yes/no, head/tail, lives/dies, male/female) with a constant probability p .

{ex:arbuthnot1}

Sometimes, as in Example 3.1 below, the available data record only the number of successes in n trials, with separate such observations recorded over time or space. More commonly, as in Example 3.2 and Example 3.3, we have available data on the frequency n_k of $k = 0, 1, 2, \dots n$ successes in the n trials.

EXAMPLE 3.1: Arbuthnot data

Sex ratios—births of male to female children have long been of interest in population studies and demography. Indeed, in 1710, John Arbuthnot (Arbuthnot, 1710) used data on the ratios of male to female christenings in London from 1629–1710 to carry out the first known significance test. The data for these 82 years showed that in *every* year there were more boys than girls. He calculated that the under the assumption that male and female births were equally likely, the probability of 82 years of more males than females was vanishingly small, ($\Pr \approx 4.14 \times 10^{-25}$). He used this to argue that a nearly constant birth ratio > 1 (or $\Pr(\text{Male}) > 0.5$) could be interpreted to show the guiding hand of a divine being.

Arbuthnot’s data, along with some other related variables are available in *Arbuthnot* in the *HistData* package. For now, we simply display a plot of the probability of a male birth over time. The plot in Figure 3.1 shows the proportion of males over years, with horizontal lines at $\Pr(\text{Male}) = 0.5$ and the mean, $\Pr(\text{Male}) = 0.517$. Also shown is a (loess) smoothed curve, which suggests that any deviation from a constant sex ratio is relatively small.

```
> data(Arbuthnot, package = "HistData")
> with(Arbuthnot, {
+   prob = Males / (Males + Females)
+   plot(x = Year, y = prob, type = "b",
+         ylim = c(0.5, 0.54), ylab = "Pr (Male)")
+   abline(h = 0.5, col = "red", lwd = 2)
```

```

+ abline(h = mean(prob), col = "blue")
+ text(x = 1640, y = 0.5, expression(H[0]: "Pr(Male)=0.5"), pos = 3, col = "red")
+ Arb.smooth <- loess.smooth(Year, prob)
+ lines(Arb.smooth, col = "blue", lwd = 2)
+ })

```

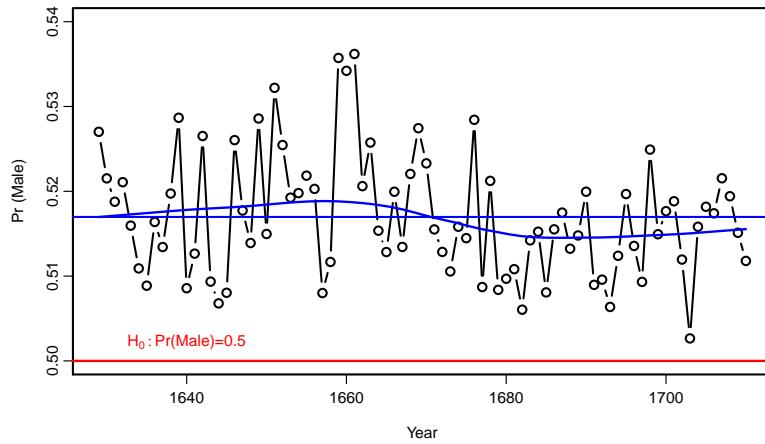


Figure 3.1: Arbuthnot's data on male/female sex ratios in London, 1629–1710, together with a (loess) smoothed curve over time and the mean $\text{Pr}(\text{Male})$

TODO: DM: use slightly simpler alternative? :

```

> prob <- with(Arbuthnot, Males / (Males + Females))
> scatter.smooth(x = Arbuthnot$Year, y = prob, type = "b",
+                   lpars = list(col = "blue", lwd = 2),
+                   xlab = "Year", ylab = "Pr(Male)", ylim = c(0.5, 0.54))
> abline(h = 0.5, col = "red", lwd = 2)
> abline(h = mean(prob), col = "blue")
> text(x = 1640, y = 0.5, expression(H[0]: "Pr(Male)=0.5"), pos = 3, col = "red")

```

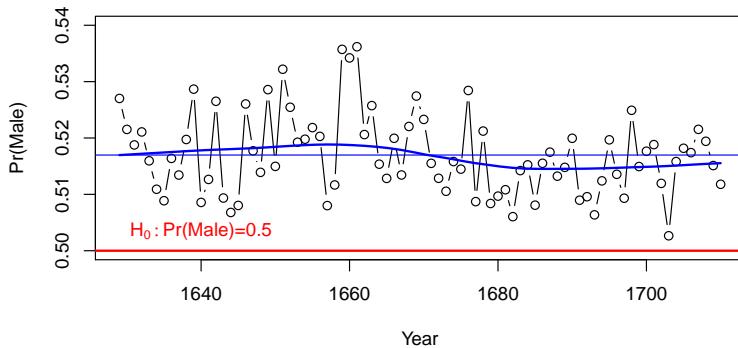


Figure 3.2: Arbuthnot's data on male/female sex ratios in London, 1629–1710, together with a (loess) smoothed curve over time and the mean $\text{Pr}(\text{Male})$

We return to this data in a later chapter where we ask whether the variation around the mean can be explained by any other considerations, or should just be considered random variation (see Exercise 7.1) \triangle

{ex:saxony1}

EXAMPLE 3.2: Families in Saxony

A related example of sex ratio data that ought to follow a binomial distribution comes from a classic study by A. Geissler (1889). Geissler listed the data on the distributions of boys and girls in families in Saxony for the period 1876–1885. In total, over four million births were recorded, and the sex distribution in the family was available because the parents had to state the sex of all their children on the birth certificate.

The complete data, classified by number of boys and number of girls (each 0–12) appear in Edwards (1958, Table 1).¹ Lindsey (1995, Table 6.2) selected only the 6115 families with 12 children, and listed the frequencies by number of males. The data are shown in table form in Table 3.1 in the standard form of a complete discrete distribution. The basic outcome variable, $k = 0, 1, \dots, 12$, is the number of male children in a family and the frequency variable, n_k is the number of families with that number of boys.

{tab:saxtab}

Table 3.1: Number of male children in 6115 Saxony families of size 12

Males (k)	0	1	2	3	4	5	6	7	8	9	10	11	12	Sum
Families (n_k)	3	24	104	286	670	1,033	1,343	1,112	829	478	181	45	7	6,115

Figure 3.3 shows a bar plot of the frequencies in Table 3.1. It can be seen that the distribution is quite symmetric. The questions of interest here are: (a) how close does the data follow a binomial distribution, with a constant $\text{Pr}(\text{Male}) = p$? (b) is there evidence to reject the hypothesis that $p = 0.5$?

```
> data(Saxony, package = "vcd")
> barplot(Saxony, xlab = "Number of males", ylab = "Number of families",
+           col = "lightblue", cex.lab = 1.5)
```

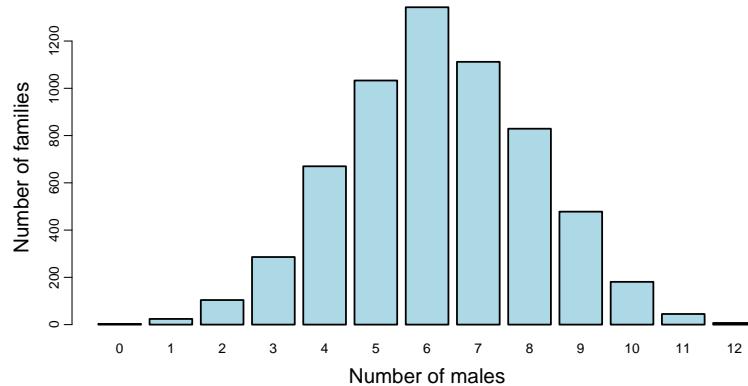


Figure 3.3: Males in Saxony families of size 12 fig:saxony-barplot

\triangle

¹Edwards (1958) notes that over these 10 years, many parents will have had several children, and their family composition is therefore recorded more than once. However, in families with a given number of children, each family can appear only once.

{ex:dice}

EXAMPLE 3.3: Weldon's dice

Common examples of binomial distributions involve tossing coins or dice, where some event outcome is considered a “success” and the number of successes (k) are tabulated in a long series of trials to give the frequency (n_k) of each basic count, k .

Perhaps the most industrious dice-tosser of all times, W. F. Raphael Weldon, an English evolutionary biologist and joint founding editor of *Biometrika* (with Francis Galton and Karl Pearson) tallied the results of throwing 12 dice 26,306 times. For his purposes, he considered the outcome of 5 or 6 pips showing on each die to be a success, and all other outcomes as failures.

Weldon reported his results in a letter to Francis Galton dated February 2, 1894, in order “to judge whether the differences between a series of group frequencies and a theoretical law . . . were more than might be attributed to the chance fluctuations of random sampling” (Kemp and Kemp, 1991). In his seminal paper, Pearson (1900) used Weldon’s data to illustrate the χ^2 goodness-of-fit test, as did Kendall and Stuart (1963, Table 5.1, p. 121).

These data are shown here as Table 3.2, in terms of the number of occurrences of a 5 or 6 in the throw of 12 dice. If the dice were all identical and perfectly fair (balanced), one would expect that $p = \Pr\{5 \text{ or } 6\} = \frac{1}{3}$ and the distribution of the number of 5 or 6 would be binomial.

A peculiar feature of these data as presented by Kendall and Stuart (not uncommon in discrete distributions) is that the frequencies of 10–12 successes are lumped together.² This grouping must be taken into account in fitting the distribution. This dataset is available as *WeldonDice* in the *vcd* package. The distribution is plotted in Figure 3.4.

Table 3.2: Frequencies of 5s or 6s in throws of 12 dice

# 5s or 6s (k)	0	1	2	3	4	5	6	7	8	9	10+	Sum
Frequency (n_k)	185	1,149	3,265	5,475	6,114	5,194	3,067	1,331	403	105	18	26,306

```
> data(WeldonDice, package = "vcd")
> dimnames(WeldonDice)$n56[11] <- "10+"
> barplot(WeldonDice, xlab = "Number of 5s and 6s", ylab = "Frequency",
+           col = "lightblue", cex.lab = 1.5)
```



3.1.2 Poisson data

Data of Poisson type arise when we observe the counts of events k within a fixed interval of time or space (length, area, volume) and tabulate their frequencies, n_k . For example, we may observe the number of radioactive particles emitted by a source per second or number of births per hour, or the number of tiger or whale sightings within some geographical regions.

In contrast to binomial data, where the counts are bounded below and above, in Poisson data the counts k are bounded below at 0, but can take integer values with no fixed upper limit. One defining characteristic for the Poisson distribution is for rare events, which occur independently with a small and constant probability, p , in small intervals, and we count the number of such occurrences.

Several examples of data of this general type are given below.

{sec:pois-data}

{ex:horsekick1}

²The unlumped entries are, for (number of 5s or 6s: frequency) — (10: 14); (11: 4), (12:0), given by Labby (2009). In this remarkable paper, Labby describes a mechanical device he constructed to repeat Weldon’s experiment physically and automate the counting of outcomes. He created electronics to roll 12 dice in a physical box, and hooked that up to a webcam to capture an image of each toss and used image processing software to record the counts.

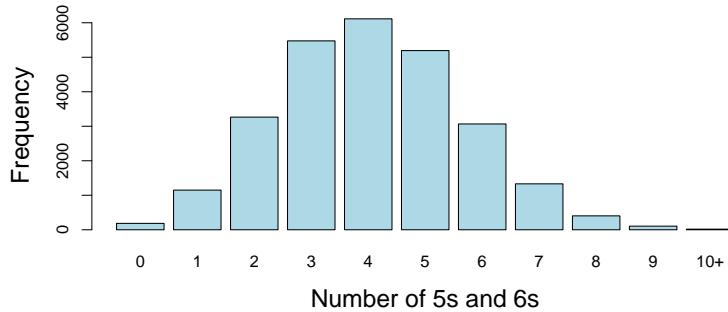


Figure 3.4: Weldon's dice data^{fig:dice}

EXAMPLE 3.4: Death by horse kick

One of the oldest and best known examples of a Poisson distribution is the data from von Bortkiewicz (1898) on deaths of soldiers in the Prussian army from kicks by horses and mules, shown in Table 3.3. Ladislaus von Bortkiewicz, an economist and statistician, tabulated the number of soldiers in each of 14 army corps in the 20 years from 1875–1894 who died after being kicked by a horse (Andrews and Herzberg, 1985, p. 18). Table 3.3 shows the data used by Fisher (1925) for 10 of these army corps, summed over 20 years, giving 200 ‘corps-year’ observations. In 109 corps-years, no deaths occurred; 65 corps-years had one death, etc.

The data set is available as *HorseKicks* in the *vcd* package. The distribution is plotted in Figure 3.5.

{tab:horsetab}

Table 3.3: von Bortkiewicz’s data on deaths by horse kicks

Number of deaths (k)	0	1	2	3	4	Sum
Frequency (n_k)	109	65	22	3	1	200

```
> data(HorseKicks, package = "vcd")
> barplot(HorseKicks, xlab = "Number of deaths", ylab = "Frequency",
+           col = "lightblue", cex.lab = 1.5)
```

{ex:madison1}



EXAMPLE 3.5: Federalist papers

In 1787–1788, Alexander Hamilton, John Jay, and James Madison wrote a series of newspaper essays to persuade the voters of New York State to ratify the U.S. Constitution. The essays were titled *The Federalist Papers* and all were signed with the pseudonym “Publius.” Of the 77 papers published, the author(s) of 65 are known, but *both* Hamilton and Madison later claimed sole authorship of the remaining 12. Mosteller and Wallace (1963, 1984) investigated the use of statistical methods to identify authors of disputed works based on the frequency distributions of certain key function words, and concluded that Madison had indeed authored the 12 disputed papers.³

³It should be noted that this is a landmark work in the development and application of statistical methods to the analysis of texts and cases of disputed authorship. In addition to *may*, they considered many such marker words, such as *any*, *by*, *from*, *upon*, and so forth. Among these, the word *upon* was the best discriminator between the works known by Hamilton (3 per 1000 words) and Madison (1/6 per 1000 words). In this work, they pioneered the use of Bayesian discriminant analysis, and the use of cross-validation to assess the stability of estimates and their conclusions.

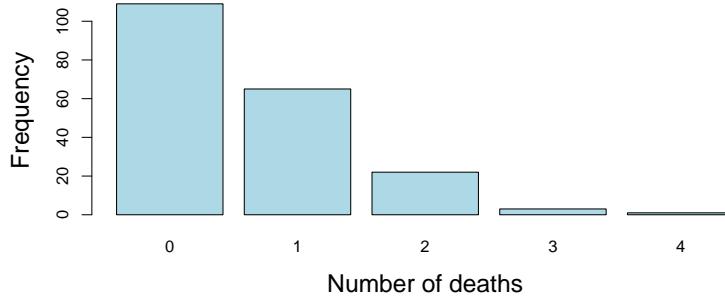


Figure 3.5: HorseKicks data fig:horsekicks

Table 3.4 shows the distribution of the occurrence of one of these “marker” words, the word *may* in 262 blocks of text (each about 200 words long) from issues of the *Federalist Papers* and other essays known to be written by James Madison. Read the table as follows: in 156 blocks, the word *may* did not occur; it occurred once in 63 blocks, etc. The distribution is plotted in Figure 3.6.

Table 3.4: Number of occurrences of the word *may* in texts written by James Madison tab:fedtab

Occurrences of <i>may</i> (k)	0	1	2	3	4	5	6	Sum
Blocks of text (n_k)	156	63	29	8	4	1	1	262

```
> data(Federalist, package = "vcd")
> barplot(Federalist,
+           xlab = "Occurrences of 'may'", ylab = "Number of blocks of text",
+           col = "lightgreen", cex.lab = 1.5)
```

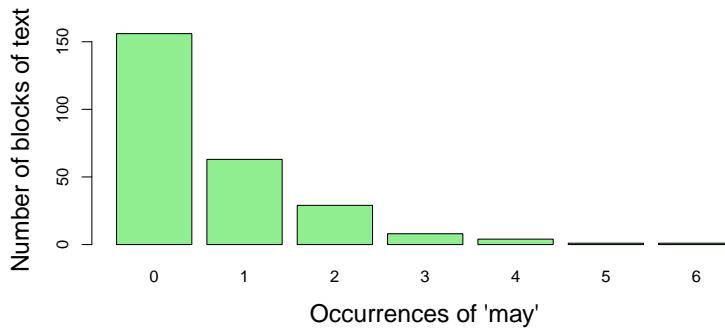


Figure 3.6: Mosteller and Wallace Federalist data fig:federalist



{ex:cyclists1}

EXAMPLE 3.6: London cycling deaths

Aberdein and Spiegelhalter (2013) observed that from November 5–13, 2013, six people were

killed while cycling in London. How unusual is this number of deaths in less than a two-week period? Was this a freak occurrence, or should Londoners petition for cycling lanes and greater road safety? To answer these question, they obtained data from the UK Department of Transport *Road Safety Data* from 2005–2012 and selected all accident fatalities of cyclists within the city of London.

It seems reasonable to assume that, in any short period of time, deaths of people riding bicycles are independent events. If, in addition, the probability of such events is constant over this time span, the Poisson distribution should describe the distribution of 0, 1, 2, 3, . . . deaths. Then, an answer to the main question can be given in terms of the probability of six (or more) deaths in a comparable period of time.

Their data, comprising 208 counts of deaths in the fortnightly periods from January 2005 to December 2012 are contained in the data set *CyclingDeaths* in *vcdExtra*. To work with the distribution, we first convert this to a one-way table.

```
> data("CyclingDeaths", package = "vcdeExtra")
> CyclingDeaths.tab <- table(CyclingDeaths$deaths)
> CyclingDeaths.tab
```

0	1	2	3
114	75	14	5

The maximum number of deaths was 3, which occurred in only 5 two-week periods. The distribution is plotted in Figure 3.7.

```
> barplot(CyclingDeaths.tab,
+           xlab = "Number of deaths", ylab = "Number of fortnights",
+           col = "pink", cex.lab = 1.5)
```

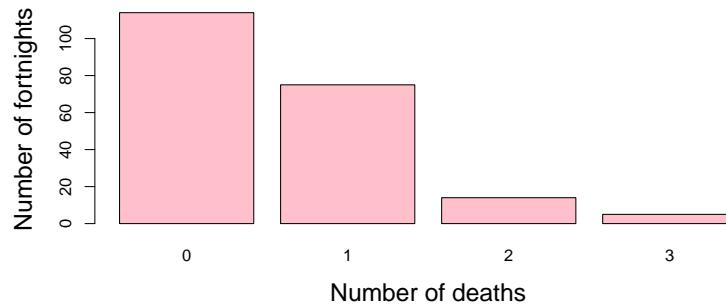


Figure 3.7: Frequencies of number of cyclist deaths in two-week periods in London, 2005–2012 fig:cyclists2

We return to this data in Example 3.10 and answer the question of how unusual six or more deaths would be in a Poisson distribution.



3.1.3 Type-token distributions

{sec:type-token} There are a variety of other types of discrete data distributions. One important class is **type-token** distributions, where the basic count k is the number of distinct types of some observed event, $k =$

$1, 2, \dots$ and the frequency, n_k , is the number of different instances observed. For example, distinct words in a book, words that subjects list as members of the semantic category “fruit”, musical notes that appear in a score, and species of animals caught in traps can be considered as types, and the occurrences of those type comprise tokens.

This class differs from the Poisson type considered above in that the frequency for value $k = 0$ is *unobserved*. Thus, questions like (a) How many words did Shakespeare know? (b) How many words in the English language are members of the “fruit” category? (c) How many wolves remain in Canada’s Northwest territories? depend on the unobserved count for $k = 0$. They cannot easily be answered without appeal to additional information or statistical theory.

{ex:butterfly}

EXAMPLE 3.7: Butterfly species in Malaya

In studies of the diversity of animal species, individuals are collected and classified by species. The distribution of the number of species (types) where $k = 1, 2, \dots$ individuals (tokens) were collected forms a kind of type-token distribution. An early example of this kind of distribution was presented by Fisher *et al.* (1943). Table 3.5 lists the number of individuals of each of 501 species of butterfly collected in Malaya. There were thus 118 species for which just a single instance was found, 74 species for which two individuals were found, down to 3 species for which 24 individuals were collected. Fisher et al. note however that the distribution was truncated at $k = 24$. Type-token distributions are often J-shaped, with a long upper tail, as we see in Figure 3.8.

Table 3.5: Number of butterfly species n_k for which k individuals were collected

{tab:buttertab}

Individuals (k)	1	2	3	4	5	6	7	8	9	10	11	12
Species (n_k)	118	74	44	24	29	22	20	19	20	15	12	14
Individuals (k)	13	14	15	16	17	18	19	20	21	22	23	24
Species (n_k)	6	12	6	9	9	6	10	10	11	5	3	3
												Sum
												501

```
> data(Butterfly, package = "vcd")
> barplot(Butterfly, xlab = "Number of individuals", ylab = "Number of species",
+           cex.lab = 1.5)
```

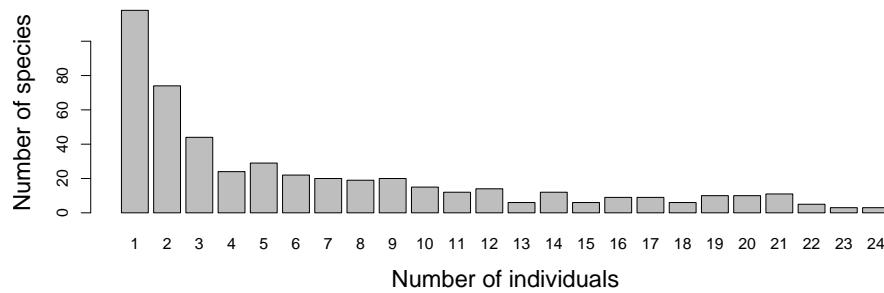


Figure 3.8: Butterfly species in Malaya^{fig:butterfly}



3.2 Characteristics of discrete distributions

This section briefly reviews the characteristics of some of the important discrete distributions encountered in practice and illustrates their use with R. An overview of these distributions is shown in Table 3.6. For more detailed information on these and other discrete distributions, Johnson *et al.* (1992) and Wimmer and Altmann (1999) present the most comprehensive treatments; Zelterman (1999, Chapter 2) gives a compact summary.

{sec:discrete-distrib}

Table 3.6: Discrete probability distributions^{tab:distns}

Discrete distribution	Probability function, $p(k)$	parameter(s)
Binomial	$\binom{n}{k} p^k (1-p)^{n-k}$	$p = \Pr(\text{success})$; $n = \# \text{ trials}$
Poisson	$e^{-\lambda} \lambda^k / k!$	$\lambda = \text{mean}$
Negative binomial	$\binom{n+k-1}{k} p^n (1-p)^k$	$p; n = \# \text{ successful trials}$
Geometric	$p(1-p)^k$	p
Logarithmic series	$\theta^k / [-k \log(1-\theta)]$	θ

For each distribution, we describe properties and generating mechanisms, and show how its parameters can be estimated and how to plot the frequency distribution. R has a wealth of functions for a wide variety of distributions. For ease of reference, their names and types for the distributions covered here are shown in Table 3.7. The naming scheme is simple and easy to remember: for each distribution, there are functions, with a prefix letter, d, p, q, r, followed by the name for that class of distribution:⁴

- d a density function,⁵ $\Pr\{X = k\} \equiv p(k)$ for the probability that the variable X takes the value k .
- p a cumulative probability function, or CDF, $F(k) = \sum_{X \leq k} p(k)$.
- q a quantile function, the inverse of the CDF, $k = F^{-1}(p)$. The quantile is defined as the smallest value x such that $F(x) \geq p$.
- r a random number generating function for that distribution.

In the R console, `help(Distributions)` gives an overview listing of the distribution functions available in the `stats` package.

Table 3.7: R functions for discrete probability distributions^{tab:distfuns}

Discrete distribution	Density (pmf) function	Cumulative (CDF)	Quantile CDF ⁻¹	Random # generator
Binomial	<code>dbinom()</code>	<code>pbinom()</code>	<code>qbinom()</code>	<code>rbinom()</code>
Poisson	<code>dpois()</code>	<code>ppois()</code>	<code>qpois()</code>	<code>rpois()</code>
Negative binomial	<code>dnbinom()</code>	<code>pnbinom()</code>	<code>qnbinom()</code>	<code>rnbinom()</code>
Geometric	<code>dgeom()</code>	<code>pgeom()</code>	<code>qgeom()</code>	<code>rgeom()</code>
Logarithmic series	<code>dlogseries()</code>	<code>plogseries()</code>	<code>qlogseries()</code>	<code>rlogseries()</code>

⁴The CRAN Task View on Probability Distributions, <http://cran.r-project.org/web/views/Distributions.html>, provides a general overview and lists a wide variety of contributed packages for specialized distributions, discrete and continuous.

⁵For discrete random variables this is usually called the probability mass function (pmf).

3.2.1 The binomial distribution

{sec:binomial}

The binomial distribution, $\text{Bin}(n, p)$, arises as the distribution of the number k of events of interest which occur in n independent trials when the probability of the event on any one trial is the constant value $p = \Pr(\text{event})$. For example, if 15% of the population has red hair, the number of red-heads in randomly sampled groups of $n = 10$ might follow a binomial distribution, $\text{Bin}(10, 0.15)$; in Weldon's dice data (Example 3.3), the probability of a 5 or 6 should be $\frac{1}{3}$ on any one trial, and the number of 5s or 6s in tosses of 12 dice would follow $\text{Bin}(12, \frac{1}{3})$.

Over n independent trials, the number of events k may range from 0 to n ; if X is a random variable with a binomial distribution, the probability that $X = k$ is given by

$$\text{Bin}(n, p) : \Pr\{X = k\} \equiv p(k) = \binom{n}{k} p^k (1 - p)^{n-k} \quad k = 0, 1, \dots, n , \quad (3.1) \quad \text{eq:binom}$$

where $\binom{n}{k} = n! / k!(n - k)!$ is the number of ways of choosing k out of n . The first three (central) moments of the binomial distribution are as follows (letting $q = 1 - p$),

$$\begin{aligned} \text{Mean}(X) &= np \\ \text{Var}(X) &= npq \\ \text{Skew}(X) &= npq(q - p) . \end{aligned}$$

It is easy to verify that the binomial distribution has its maximum variance when $p = \frac{1}{2}$. It is symmetric ($\text{Skew}(x)=0$) when $p = \frac{1}{2}$, and negatively (positively) skewed when $p < \frac{1}{2}$ ($p > \frac{1}{2}$).

If we are given data in the form of a discrete (binomial) distribution (and n is known), then the maximum likelihood estimator of p can be obtained as the weighted mean of the values k with weights n_k ,

$$\hat{p} = \frac{\bar{x}}{n} = \frac{(\sum_k k \times n_k) / \sum_k n_k}{n} ,$$

and has sampling variance $\mathcal{V}(\hat{p}) = pq/n$.

TODO: DM: either add ref to some text explaining Maximum Likelihood estimation, or maybe add a section similar to old book to the Appendix), or add a note in preface that this is assumed to be known.

3.2.1.1 Calculation and visualization

As indicated in Table 3.7 (but without listing the parameters of these functions), binomial probabilities can be calculated with `dbinom(x, n, p)`, where `x` is a vector of the number of successes in `n` trials and `p` is the probability of success on any one trial. Cumulative probabilities, summed up to a vector of quantiles, `Q` can be calculated with `pbinom(Q, n, p)`, and the quantiles (the smallest value `x` such that $F(x) \geq P$) with `qbinom(P, n, p)`. To generate `N` random observations from a binomial distribution with `n` trials and success probability `p` use `rbinom(N, n, p)`⁶.

For example, to find and plot the binomial probabilities corresponding to Weldon's tosses of 12 dice, with $k = 0, \dots, 12$ and $p = \frac{1}{3}$, we could do the following

```
> k <- seq(0, 12)
> Pk <- dbinom(k, 12, 1/3)
> plot(x = k, y = Pk, type = "h",
+       xlab = "Number of successes", ylab = "Probability",
+       lwd = 8, lend = "square")
> lines(x = k, y = Pk)
```

TODO: DM: Why not directly using a barplot?

⁶Note that the actual R function arguments differ from the ones used here.

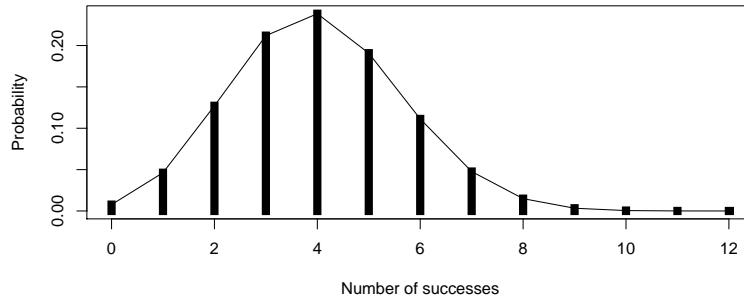


Figure 3.9: Binomial distribution for $k = 0, \dots, 12$ successes in 12 trials and $p=1/3$ fig:dbinom1

```
> k <- 0 : 12
> Pk <- dbinom(k, 12, 1/3)
> b <- barplot(Pk, names.arg = k,
+                 xlab = "Number of successes", ylab = "Probability")
> lines(x = b, y = Pk, col = "red")
```

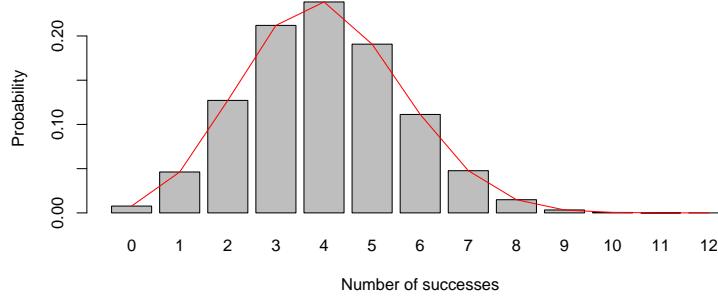


Figure 3.10: Binomial distribution for $k = 0, \dots, 12$ successes in 12 trials and $p=1/3$ fig:dbinom12

Note that in the call to `plot()`, `type = "h"` draws histogram-typed lines to the bottom of the vertical axis, and `lwd = 8` makes them wide. The call to `lines()` shows another way to plot the data, as a probability polygon. We illustrate other styles for plotting in Section 3.2.2, Example 3.11 below.

{ex:dice2}

EXAMPLE 3.8: Weldon's dice

Going a bit further, we can compare Weldon's data with the theoretical binomial distribution as shown below. Because the `WeldonDice` data collapsed the frequencies for 10–12 successes as 10+, we do the same with the binomial probabilities. The expected frequencies (`Exp`), if Weldon's dice tosses obeyed the binomial distribution, are calculated as $N \times p(k)$ for $N = 26,306$ tosses. In addition, we compute the differences of the observed (`Freq`) and expected (`Exp`) frequencies as column `Diff`, to be used for the χ^2 test for goodness of fit described later in Section 3.3, but a glance these are all negative for $k = 0, \dots, 4$ and positive thereafter.

```

> Weldon_df <- as.data.frame(WeldonDice) # convert to data frame
>
> k <- 0 : 12                         # same as seq(0, 12)
> Pk <- dbinom(k, 12, 1/3)            # binomial probabilities
> Pk <- c(Pk[1:10], sum(Pk[11:13])) # sum values for 10+
> Exp <- round(26306 * Pk)           # expected frequencies
> Diff <- Weldon_df$Freq - Exp      # raw residuals
> Chisq <- Diff^2 / Exp
> data.frame(Weldon_df, Prob = round(Pk, 5), Exp, Diff, Chisq)

  n56 Freq   Prob   Exp Diff Chisq
1    0 185 0.00771 203 -18 1.59606
2    1 1149 0.04624 1216 -67 3.69161
3    2 3265 0.12717 3345 -80 1.91330
4    3 5475 0.21195 5576 -101 1.82945
5    4 6114 0.23845 6273 -159 4.03013
6    5 5194 0.19076 5018 176 6.17298
7    6 3067 0.11127 2927 140 6.69628
8    7 1331 0.04769 1255  76 4.60239
9    8  403 0.01490 392   11 0.30867
10   9  105 0.00331  87   18 3.72414
11  10+   18 0.00054  14     4 1.14286

```

△

Finally, we can use programming features in R to calculate and plot probabilities for binomial distributions over a range of both k and p as follows, for the purposes of graphing the distributions as one or both varies. The following code uses `expand.grid()` to create a data frame `KP` containing all combinations of $k = 0:12$ and $p = c(1/6, 1/3, 1/2, 2/3)$. These values are then supplied as arguments to `dbinom()`. For the purpose of plotting, the decimal value of p is declared as a factor.

```

> KP <- expand.grid(k = 0 : 12, p = c(1/6, 1/3, 1/2, 2/3))
> bin_df <- data.frame(KP, prob = dbinom(KP$k, 12, KP$p))
> bin_df$p <- factor(bin_df$p, labels = c("1/6", "1/3", "1/2", "2/3"))
> str(bin_df)

'data.frame': 52 obs. of  3 variables:
 $ k   : int  0 1 2 3 4 5 6 7 8 9 ...
 $ p   : Factor w/ 4 levels "1/6","1/3","1/2",...: 1 1 1 1 1 1 1 1 1 ...
 $ prob: num  0.1122 0.2692 0.2961 0.1974 0.0888 ...

```

This data can be plotted using `xypplot()` in `lattice`, using the `groups` argument to make separate curves for each value of p . The following code generates Figure 3.11.

```

> library(lattice)
> mycol <- palette()[2:5]
> xypplot(prob ~ k, data = bin_df, groups = p,
+   xlab = list("Number of successes", cex = 1.25),
+   ylab = list("Probability", cex = 1.25),
+   type = "b", pch = 15:17, lwd = 2, cex = 1.25, col = mycol,
+   key = list(
+     title = "Pr(success)",
+     points = list(pch = 15 : 17, col = mycol, cex = 1.25),
+     lines = list(lwd = 2, col = mycol),
+     text = list(levels(bin_df$p)),
+     x = 0.9, y = 0.98, corner = c(x = 1, y = 1)
+   )
+ )

```

TODO: DM: Avoid lattice to reduce complexity. Either only use `ggplot2` throughout the chapter, or use base graphics here, and `ggplot2` for the remaining plots:

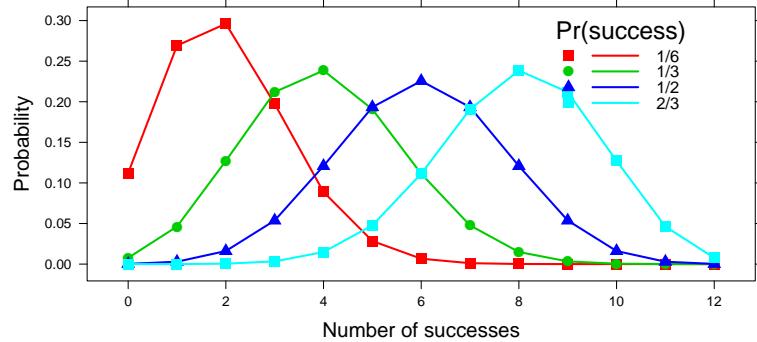


Figure 3.11: Binomial distributions for $k = 0, \dots, 12$ successes in $n = 12$ trials, and four values of p

```
> p <- c(1/6, 1/3, 1/2, 2/3)
> k <- 0 : 12
> Prob <- outer(k, p, function(k, p) dbinom(k, 12, p))
> str(Prob)

num [1:13, 1:4] 0.1122 0.2692 0.2961 0.1974 0.0888 ...
```

```
> col <- palette()[2:5]
> matplot(k, Prob,
+           type = "o", pch = 15 : 17, col = col, lty = 1,
+           xlab = "Number of Successes", ylab = "Probability")
> legend("topright", legend = c("1/6", "1/3", "1/2", "2/3"),
+         pch = 15 : 17, lty = 1, col = col, title = "Pr(Success)")
```

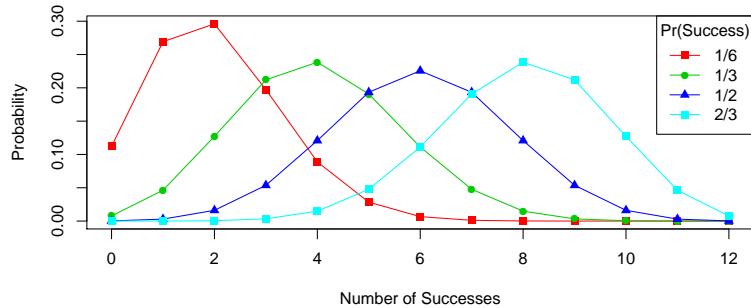


Figure 3.12: Binomial distributions for $k = 0, \dots, 12$ successes in $n = 12$ trials, and four values of p

3.2.2 The Poisson distribution

{sec:poisson}

The Poisson distribution gives the probability of an event occurring $k = 0, 1, 2, \dots$ times over a large number of independent “trials”, when the probability, p , that the event occurs on any one trial

(in time or space) is small and constant. Hence, the Poisson distribution is usually applied to the study of rare events such as highway accidents at a particular location, deaths from horse kicks, or defects in a well-controlled manufacturing process. Other applications include: the number of customers contacting a call center per unit time; the number of insurance claims per unit region or unit time; number of particles emitted from a small radioactive sample.

For the Poisson distribution, the probability function is

$$\text{Pois}(\lambda) : \Pr\{X = k\} \equiv p(k) = \frac{e^{-\lambda} \lambda^k}{k!} \quad k = 0, 1, \dots \quad (3.2) \quad \{\text{eq:poisf}\}$$

where the rate parameter, $\lambda (> 0)$, turns out to be the mean of the distribution. The first three (central) moments of the Poisson distribution are:

$$\begin{aligned} \text{Mean}(X) &= \lambda \\ \text{Var}(X) &= \lambda \\ \text{Skew}(X) &= \lambda^{-1/2} \end{aligned}$$

So, the mean and variance of the Poisson distribution are always the same, which is sometimes used to identify a distribution as Poisson. For the binomial distribution, the mean (Np) is always greater than the variance (Npq); for other distributions (negative binomial and geometric) the mean is less than the variance. The Poisson distribution is always positively skewed, but skewness decreases as λ increases.

The maximum likelihood estimator of the parameter λ in Eqn. (3.2) is just the mean of the distribution,

$$\hat{\lambda} = \bar{x} = \frac{\sum_k k n_k}{\sum_k n_k}. \quad (3.3) \quad \{\text{eq:pois-lambda}\}$$

Hence, the expected frequencies can be estimated by substituting the sample mean into Eqn. (3.2) and multiplying by the total sample size N .

There are many useful properties of the Poisson distribution. **TODO: DM: Better add some book ref? Wikipedia is not persistent ...**⁷ Among these:

- Poisson variables have a nice reproductive property: if X_1, X_2, \dots, X_m are independent Poisson variables with the same parameter λ , then their sum, $\sum X_i$ is a Poisson variate with parameter $m\lambda$; if the Poisson parameters differ, the sum is still Poisson with parameter $\sum \lambda_i$.
- For two or more independent Poisson variables, $X_1 \sim \text{Pois}(\lambda_1), X_2 \sim \text{Pois}(\lambda_2), \dots$, with rate parameters $\lambda_1, \lambda_2, \dots$, the distribution of any X_i , *conditional on their sum*, $\sum_j X_j = n$, is binomial, $\text{Bin}(n, p)$, where $p = \lambda_i / \sum_j \lambda_j$.
- As λ increases, the Poisson distribution becomes increasingly symmetric, and approaches the normal distribution $N(\lambda, \lambda)$ with mean and variance λ as $\lambda \rightarrow \infty$. The approximation is quite good with $\lambda > 20$.
- If $X \sim \text{Pois}(\lambda)$, then \sqrt{X} converges much faster to a normal distribution $N(\lambda, \frac{1}{4})$, with mean $\sqrt{\lambda}$ and constant variance $\frac{1}{4}$. Hence, the square root transformation is often recommended as a *variance stabilizing* transformation for count data when classical methods (ANOVA, regression) assuming normality are employed.

{ex:soccer}

EXAMPLE 3.9: UK Soccer scores

Table 3.8 gives the distributions of goals scored by the 20 teams in the 1995/96 season of the Premier League of the UK Football Association as presented originally by Lee (1997), and now available as the two-way table *UKSoccer* in the *vcd* package. Over a season each team plays each

⁷See: http://en.wikipedia.org/wiki/Poisson_distribution

Table 3.8: Goals scored by home and away teams in 380 games in the Premier Football League, 1995/96 season

{tab:soccer1}

Home Team Goals	Away Team Goals					Total
	0	1	2	3	4+	
0	27	29	10	8	2	76
1	59	53	14	12	4	142
2	28	32	14	12	4	90
3	19	14	7	4	1	45
4+	7	8	10	2	0	27
Total	140	136	55	38	11	380

other team exactly once, so there are a total of $20 \times 19 = 380$ games. Because there may be an advantage for the home team, the goals scored have been classified as “home team” goals and “away team” goals in the table. Of interest for this example is whether the number of goals scores by home teams and away teams follow Poisson distributions, and how this relates to the distribution of the total number of goals scored.

If we assume that in any small interval of time there is a small, constant probability that the home team or the away team may score a goal, the distributions of the goals scored by home teams (the row totals in Table 3.8) may be modeled as $\text{Pois}(\lambda_H)$ and the distribution of the goals scored by away teams (the column totals) may be modeled as $\text{Pois}(\lambda_A)$.

If the number of goals scored by the home and away teams are independent⁸, we would expect that the total number of goals scored in any game would be distributed as $\text{Pois}(\lambda_H + \lambda_A)$. These totals are shown in Table 3.9.

{tab:soccer2}

Table 3.9: Total goals scored in 380 games in the Premier Football League, 1995/95 season

Total goals	0	1	2	3	4	5	6	7
Number of games	27	88	91	73	49	31	18	3

As a preliminary check of the distributions for the home and away goals, we can determine if the means and variances are reasonably close to each other. If so, then the total goals variable should also have a mean and variance equal to the sum of those statistics for the home and away goals.

In the R code below, we first convert the two-way frequency table *UKSoccer* to a data frame in frequency form. We use *within()* to convert *Home* and *Away* to numeric variables, and calculate *Total* as their sum.

```
> data(UKSoccer, package = "vcd")
>
> soccer.df <- as.data.frame(UKSoccer, stringsAsFactors = FALSE)
> soccer.df <- within(soccer.df, {
+   Home <- as.numeric(Home)           # make numeric
+   Away <- as.numeric(Away)           # make numeric
+   Total <- Home + Away              # total goals
})
```

⁸This question is examined visually in Chapter 5 (Example 5.5) and Chapter 6 (Example 6.11), where we find that the answer is “basically, yes”.

```
+ })
> str(soccer.df)

'data.frame': 25 obs. of 4 variables:
 $ Home : num 0 1 2 3 4 0 1 2 3 4 ...
 $ Away : num 0 0 0 0 1 1 1 1 ...
 $ Freq : num 27 59 28 19 7 29 53 32 14 8 ...
 $ Total: num 0 1 2 3 4 1 2 3 4 5 ...
```

To calculate the mean and variance of these variables, first expand the data frame to 380 individual observations using `expand.dft()`. Then use `apply()` over the rows to calculate the mean and variance in each column.

```
> soccer.df <- expand.dft(soccer.df) # expand to ungrouped form
> apply(soccer.df, 2, FUN = function(x) c(mean = mean(x), var = var(x)))

   Home     Away    Total
mean 1.4868 1.0632 2.5500
var  1.3164 1.1728 2.6175
```

The means are all approximately equal to the corresponding variances. More to the point, the variance of the Total score is approximately equal to the sum of the individual variances. Note also there does appear to be an advantage for the home team, of nearly half a goal.



{ex:cyclists2}

EXAMPLE 3.10: London cycling deaths

A quick check of whether the numbers of deaths among London cyclists follows the Poisson distribution can be carried out by calculating the mean and variance. The *index of dispersion*, the ratio of the variance to the mean, is commonly used to quantify whether a set of observed frequencies is more or less dispersed than a reference (Poisson) distribution.

```
> with(CyclingDeaths, c(mean = mean(deaths),
+                         var = var(deaths),
+                         ratio = mean(deaths) / var(deaths)))

  mean      var      ratio
0.56731 0.52685 1.07679
```

Thus, there was an average of about 0.57 deaths per fortnight, or a bit more than 1 per month, and no evidence for over- or underdispersion.

We can now answer the question of whether it was an extraordinary event to observe six deaths in a two-week period, by calculating the probability of more than 5 deaths using `ppois()`.

```
> mean.deaths <- mean(CyclingDeaths$deaths)
> ppois(5, mean.deaths, lower.tail = FALSE)

[1] 2.8543e-05
```

This probability is extremely small, so we conclude that the occurrence of six deaths was a singular event. The interpretation of this result might indicate an increased risk to cycling in London, and might prompt further study of road safety.



3.2.2.1 Calculation and visualization

For the Poisson distribution, you can generate probabilities using `dpois(x, lambda)` for the numbers of events in `x` with rate parameter `lambda`. As we did earlier for the binomial distribution, we can calculate these for a collection of values of `lambda` by using `expand.grid()` to create all combinations of with the values of `x` we wish to plot.

{ex:dpois-plot}

EXAMPLE 3.11: Plotting styles for discrete distributions

In this example, we illustrate some additional styles for plotting discrete distributions, using both `lattice xyplot()` and the `ggplot2` package. The goal here is to visualize a collection of Poisson distributions for varying values of λ .

We first create the 63 combinations of $x = 0 : 20$ for three values of λ , `lambda = c(1, 4, 10)`, and use these columns as arguments to `dpois()`. Again, `lambda` is a numeric variable, but the plotting methods are easier if this variable is converted to a factor.

```
> KL <- expand.grid(k = 0 : 20, lambda = c(1, 4, 10))
> pois_df <- data.frame(KL, prob = dpois(KL$k, KL$lambda))
> pois_df$lambda = factor(pois_df$lambda)
> str(pois_df)

'data.frame': 63 obs. of  3 variables:
 $ k      : int  0 1 2 3 4 5 6 7 8 9 ...
 $ lambda: Factor w/ 3 levels "1","4","10": 1 1 1 1 1 1 1 1 1 ...
 $ prob   : num  0.3679 0.3679 0.1839 0.0613 0.0153 ...
```

Discrete distributions are often plotted as bar charts or in histogram-like form, as we did for the examples in Section 3.1, rather than the line-graph form used for the binomial distribution in Figure 3.11. With `xyplot()`, the plot style is controlled by the `type` argument, and the code below uses `type = c("h", "p")` to get *both* histogram-like lines to the origin and points. As well, the plot formula, `prob ~ k | lambda` instructs `xyplot()` to produce a multi-panel plot, conditioned on values of `lambda`. These lines produce Figure 3.13.

```
> xyplot(prob ~ k | lambda, data = pois_df,
+         type = c("h", "p"), pch = 16, lwd = 4, cex = 1.25, layout = c(3, 1),
+         xlab = list("Number of events (k)", cex = 1.25),
+         ylab = list("Probability", cex = 1.25))
```

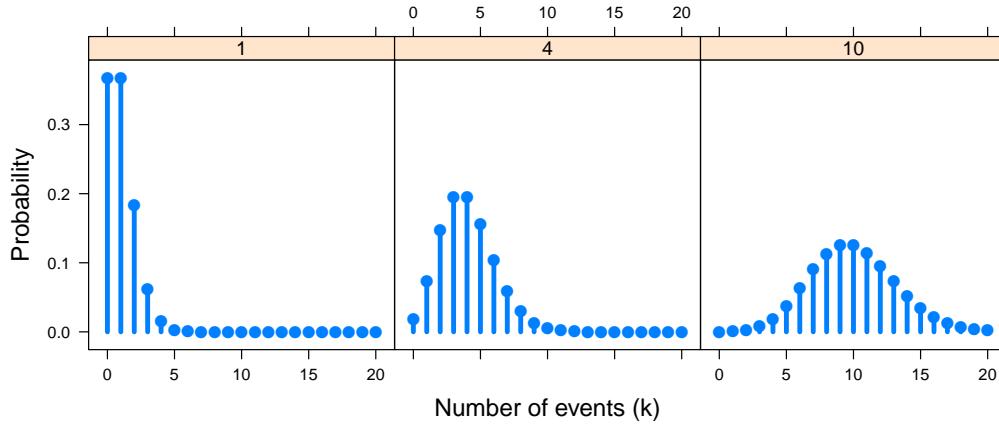


Figure 3.13: Poisson distributions for $\lambda = 1, 4, 10$, in a multi-panel display^{fig:dpois-xyplot1}

The line-graph plot style of Figure 3.11 has the advantage that it is easier to compare the separate distributions in a single plot (using the `groups` argument) than across multiple panels (using a conditioning formula). It has the disadvantages that (a) a proper legend is difficult to construct with `lattice`, and (b) is difficult to read, because you have to visually coordinate the curves in the plot with the values shown in the legend. Figure 3.14 solves both problems using the `directlabels` package.

```
> mycol <- palette() [2:4]
> plt <- xyplot(prob ~ k, data = pois_df, groups = lambda,
+   type = "b", pch = 15 : 17, lwd = 2, cex = 1.25, col = mycol,
+   xlab = list("Number of events (k)", cex = 1.25),
+   ylab = list("Probability", cex = 1.25))
>
> library(directlabels)
> direct.label=plt, list("top.points", cex = 1.5, dl.trans(y = y + 0.1)))
```

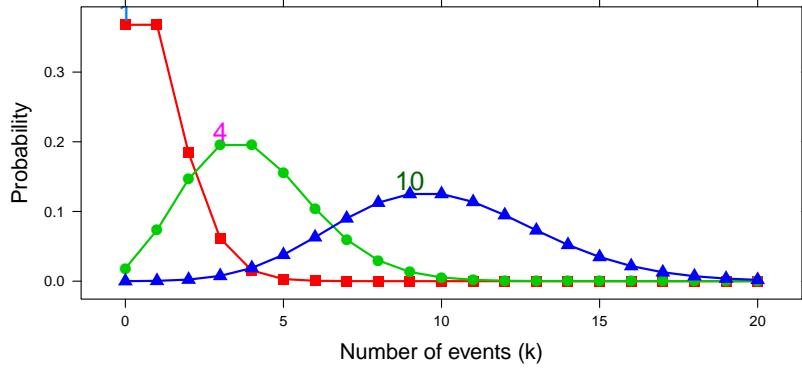


Figure 3.14: Poisson distributions for $\lambda = 1, 4, 10$, using direct labels fig:dpois-xyplot2

Note that the plot constructed by `xyplot()` is saved as a ("trellis") object, `plt`. The function `direct.label()` massages this to add the labels directly to each curve. In the second argument above, "top.points" says to locate these at the maximum value on each curve.

Finally, we illustrate the use of `ggplot2` to produce a single-panel, multi-line plot of these distributions. The basic plot uses `aes(x = k, y = prob, ...)` to produce a plot of `prob` vs. `k`, assigning color and shape attributes to the values of `lambda`.

```
> library(ggplot2)
> gplt <- ggplot(pois_df, aes(x = k, y = prob, colour = lambda, shape = lambda)) +
+   geom_line(size = 1) + geom_point(size = 3) +
+   xlab("Number of events (k)") +
+   ylab("Probability")
```

`ggplot2` allows most details of the plot to be modified using `theme()`. Here we use this to move the legend inside the plot, and enlarge the axis labels and titles.

```
> gplt + theme(legend.position = c(0.8, 0.8)) + # manually move legend
+   theme(axis.text = element_text(size = 12),
+         axis.title = element_text(size = 14, face = "bold"))
```



3.2.3 The negative binomial distribution

The negative binomial distribution is a type of waiting-time distribution, but also arises in statistical applications as a generalization of the Poisson distribution, allowing for **overdispersion** (variance > mean). See Hilbe (2011) for a comprehensive treatment of negative binomial statistical models with many applications in R.

{sec:negbin}

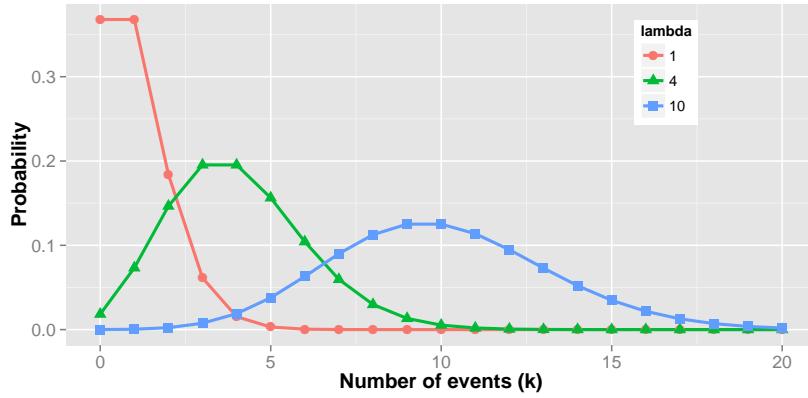


Figure 3.15: Poisson distributions for $\lambda = 1, 4, 10$, using `ggplot2` fig:dpois-ggplot2

One form of the negative binomial distribution (also called the **Pascal distribution**) arises when a series of independent Bernoulli trials is observed with constant probability p of some event, and we ask how many non-events (failures), k , it takes to observe n successful events. For example, in tossing one die repeatedly, we may consider the outcome “1” as a “success” (with $p = \frac{1}{6}$) and ask about the probability of observing $k = 0, 1, 2, \dots$ failures before getting $n = 3$ 1s.

The probability function with parameters n (a positive integer, $0 < n < \infty$) and p ($0 < p < 1$) gives the probability that k non-events (failures) are observed before the n -th event (success), and can be written⁹

$$\text{NBin}(n, p) : \Pr\{X = k\} \equiv p(k) = \binom{n+k-1}{k} p^n (1-p)^k \quad k = 0, 1, \dots, \infty \quad (3.4)$$

This formulation makes clear that a given sequence of events involves a total of $n + k$ trials of which there are n successes, with probability p^n , and k are failures, with probability $(1-p)^k$. The binomial coefficient, $\binom{n+k-1}{k}$ gives the number of ways to choose the k successes from the remaining $n + k - 1$ trials preceding the last success.

The first three central moments of the negative binomial distribution are:

$$\begin{aligned} \text{Mean}(X) &= nq/p = \mu \\ \text{Var}(X) &= nq/p^2 \\ \text{Skew}(X) &= \frac{2-p}{\sqrt{nq}}, \end{aligned}$$

where $q = 1 - p$. The variance of X is therefore greater than the mean, and the distribution is always positively skewed.

A more general form of the negative binomial distribution (the **Polya distribution**) allows n to take non-integer values and to be an unknown parameter. In this case, the combinatorial coefficient, $\binom{n+k-1}{k}$ in Eqn. (3.4) is calculated using the gamma function, $\Gamma(\bullet)$, a generalization of the factorial for non-integer values, defined so that $\Gamma(x+1) = x!$ when x is an integer.

⁹There are a variety of other parameterizations of the negative binomial distribution, but all of these can be converted to the form shown here, which is relatively standard, and consistent with R. They differ in whether the parameter n relates to the number of successes or the total number of trials, and whether the stopping criterion is defined in terms of failures or successes. See: http://en.wikipedia.org/wiki/Negative_binomial_distribution for details on these variations.

Then the probability function Eqn. (3.4) becomes

$$\Pr\{X = k\} \equiv p(k) = \frac{\Gamma(n+k)}{\Gamma(n)\Gamma(k+1)} p^n (1-p)^k \quad k = 0, 1, \dots, \infty . \quad (3.5)$$

Greenwood and Yule (1920) developed the negative binomial distribution as a model for accident proneness or susceptibility of individuals to repeated attacks of disease. They assumed that for any individual, i , the number of accidents or disease occurrences has a Poisson distribution with parameter λ_i . If individuals vary in proneness, so that the λ_i have a gamma distribution, the resulting distribution is the negative binomial.

In this form, the negative binomial distribution is frequently used as an alternative to the Poisson distribution when the assumptions of the Poisson (constant probability and independence) are not satisfied, or when the variance of the distribution is greater than the mean (overdispersion). This gives rise to an alternative parameterization in terms of the mean (μ) of the distribution and its relation to the variance. From the relation of the mean and variance to the parameters n, p given above,

$$\text{Mean}(X) = \mu = \frac{n(1-p)}{p} \implies p = \frac{n}{n+\mu} \quad (3.6)$$

$$\text{Var}(X) = \frac{n(1-p)}{p^2} \implies \text{Var}(X) = \mu + \frac{\mu^2}{n} \quad (3.7)$$

This formulation allows the variance of the distribution to exceed the mean, and in these terms, the “size” parameter n is called the **dispersion parameter**.¹⁰ Increasing this parameter corresponds to less heterogeneity, variance closer to the mean, and therefore greater applicability of the Poisson distribution.

3.2.3.1 Calculation and visualization

In R, the density (pmf), distribution (CDF), quantile and random number functions for the negative binomial distribution are a bit special, in that the parameterization can be specified using either (n, p) or (n, μ) forms, where $\mu = n(1-p)/p$. In our notation, probabilities can be calculated using `dnbino(m)` using the call `dbinom(k, n, p)` or the call `dbinom(k, n, mu=)`, as illustrated below:

```
> k <- 2
> n <- 2 : 4
> p <- 0.2
> dnbino(m)(k, n, p)

[1] 0.07680 0.03072 0.01024

> (mu <- n * (1 - p) / p)

[1] 8 12 16

> dnbino(m)(k, n, mu = mu)

[1] 0.07680 0.03072 0.01024
```

Thus, for the distribution with $k = 2$ failures and $n = 2 : 4$ successes with probability $p = 0.2$, the values $n = 2 : 4$ correspond to means $\mu = 8, 12, 16$ as shown above.

¹⁰Other terms are “shape parameter,” with reference to the mixing distribution of Poissons with varying λ , “heterogeneity parameter,” or “aggregation parameter.”

As before, we can calculate these probabilities for a range of the combinations of arguments using `expand.grid()`. In the example below, we allow three values for each of n and p and calculate all probabilities for all values of k from 0 to 20. The result, `nbin_df` is like a 3-way, $21 \times 3 \times 3$ array of prob values, but in data frame format.

```
> XN <- expand.grid(k = 0 : 20, n = c(2, 4, 6), p = c(0.2, 0.3, 0.4))
> nbin_df <- data.frame(XN, prob = dnbinom(XN$k, XN$n, XN$p))
> nbin_df$n <- factor(nbin_df$n)
> nbin_df$p <- factor(nbin_df$p)
> str(nbin_df)

'data.frame': 189 obs. of 4 variables:
 $ k    : int 0 1 2 3 4 5 6 7 8 9 ...
 $ n    : Factor w/ 3 levels "2","4","6": 1 1 1 1 1 1 1 1 1 ...
 $ p    : Factor w/ 3 levels "0.2","0.3","0.4": 1 1 1 1 1 1 1 1 1 ...
 $ prob: num 0.04 0.064 0.0768 0.0819 0.0819 ...
```

With 9 combinations of the parameters, it is most convenient to plot these in separate panels, in a 3×3 display. The formula `prob ~ k | n + p` in the call to `xyplot()` constructs plots of `prob` vs. `k` conditioned on the combinations of `n` and `p`.

```
> xyplot(prob ~ k | n + p, data = nbin_df,
+   xlab = list("Number of failures (k)", cex = 1.25),
+   ylab = list("Probability", cex = 1.25),
+   type = c("h", "p"), pch = 16, lwd = 2,
+   strip = strip.custom(strip.names = TRUE)
+ )
```

It can be readily seen that the mean increases from left to right with n , and increases from top to bottom with decreasing p . For these distributions, we can also calculate the theory-implied means, μ , across the entire distributions, $k = 0, 1, \dots \infty$, as shown below.

```
> NP <- expand.grid(n=c(2, 4, 6), p=c(0.2, 0.3, 0.4))
> NP <- within(NP, { mu = n*(1-p)/p })
> # show as matrix
> matrix(NP$mu, 3, 3, dimnames=list(n=c(2,4,6), p=(2:4)/10))

      p
n   0.2     0.3  0.4
  2    8   4.6667   3
  4   16   9.3333   6
  6   24  14.0000   9
```

TODO: DM: maybe simpler?

```
> n <- c(2, 4, 6)
> p <- c(0.2, 0.3, 0.4)
> NP <- outer(n, p, function(n, p) n * (1 - p) / p)
> dimnames(NP) <- list(n = n, p = p)
> NP

      p
n   0.2     0.3  0.4
  2    8   4.6667   3
  4   16   9.3333   6
  6   24  14.0000   9
```

3.2.4 The geometric distribution

{sec:geometricic}

The special case of the negative binomial distribution when $n = 1$ is a geometric distribution. We observe a series of independent trials and count the number of non-events (failures) preceding the

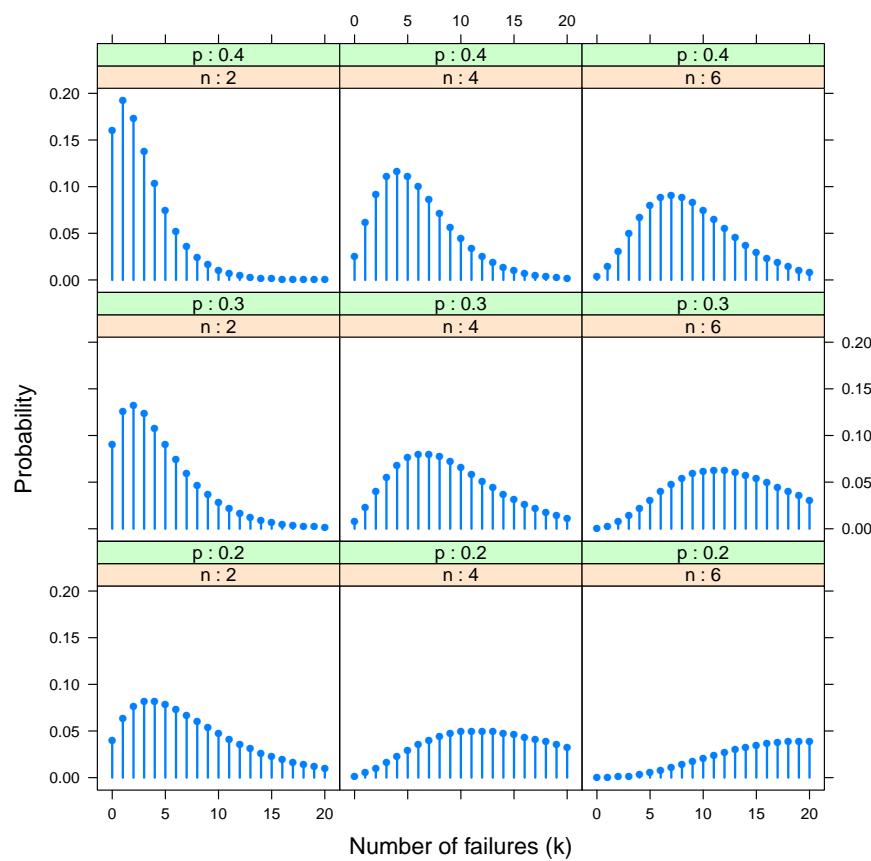


Figure 3.16: Negative binomial distributions for $n = 2, 4, 6$ and $p = 0.2, 0.3, 0.4$, using `xyplot`^{fig:dnbin3}

first successful event. The probability that there will be k failures before the first success is given by

$$\{eq:geomf\} \quad \text{Geom}(p) : \Pr\{X = k\} \equiv p(k) = p(1 - p)^k \quad k = 0, 1, \dots . \quad (3.8)$$

For this distribution the central moments are:

$$\begin{aligned} \text{Mean}(X) &= 1/p \\ \text{Var}(X) &= (1 - p)/p^2 \\ \text{Skew}(X) &= (2 - p)/\sqrt{1 - p} \end{aligned}$$

Note that estimation of the parameter p for the geometric distribution can be handled as the special case of the negative binomial by fixing $n = 1$, so no special software is needed. Going the other way, if X_1, X_2, \dots, X_n are independent geometrically distributed as $\text{Geom}(p)$, then their sum, $Y = \sum_j^n X_j$ is distributed as $\text{NBin}(p, n)$.

In R, the standard set of functions for the geometric distribution are available as `dgeom(x, prob)`, `pgeom(q, prob)`, `qgeom(p, prob)` and `rgeom(n, prob)` where `prob` represents p here. Visualization of the geometric distribution follows the pattern used earlier for other discrete distributions.

3.2.5 The logarithmic series distribution

The logarithmic series distribution is a long-tailed distribution introduced by Fisher *et al.* (1943) in connection with data on the abundance of individuals classified by species of the type shown for the distribution of butterfly species in Table 3.5.

The probability distribution function with parameter p is given by

$$\{eq:logseriesf\} \quad \text{LogSer}(p) : \Pr\{X = k\} \equiv p(k) = \frac{p^k}{-(k \log(1 - p))} = \alpha p^k/k \quad k = 1, 2, \dots, \infty , \quad (3.9)$$

where $\alpha = -1/\log(1 - p)$ and $0 < p < 1$. For this distribution, the first two central moments are:

$$\begin{aligned} \text{Mean}(X) &= \alpha \left(\frac{p}{1 - p} \right) \\ \text{Var}(X) &= -p \frac{p + \log(1 - p)}{(1 - p)^2 \log^2(1 - p)} \end{aligned}$$

Fisher derived the logarithmic series distribution by assuming that for a given species the number of individuals trapped has a Poisson distribution with parameter $\lambda = \gamma t$, where γ is a parameter of the species (susceptibility to entrapment) and t is a parameter of the trap. If different species vary so that the parameter γ has a gamma distribution, then the number of representatives of each species trapped will have a negative binomial distribution. However, the observed distribution is necessarily truncated on the left, because one cannot observe the number of species never caught (where $k = 0$). The logarithmic series distribution thus arises as a limiting form of the zero-truncated negative binomial.

Maximum likelihood estimation of the parameter p in the log-series distribution is described by Böhning (1983), extending a simpler Newton's method approximation by Birch (1963a). The `vcdExtra` package contains the set of R functions, `dlogseries(x, prob)`, `plogseries(q, prob)`, `qlogseries(p, prob)` and `rlogseries(n, prob)` where `prob` represents p here.

TODO: implement the log-series in `goodfit()` and `distplot()` so this distribution can be used in later sections.

3.2.6 Power series family

{sec:pwrseries}

We mentioned earlier that the Poisson distribution was unique among all discrete (one parameter) distributions, in that it is the only one whose mean and variance are equal (Kosambi, 1949). The relation between mean and variance of discrete distributions also provides the basis for integrating them into a general family. All of the discrete distributions described in this section are in fact special cases of a family of discrete distributions called the power series distributions by Noack (1950) and defined by

$$p(k) = a(k)\theta^k/f(\theta) \quad k = 0, 1, \dots ,$$

with parameter $\theta > 0$, where $a(k)$ is a coefficient function depending only on k and $f(\theta) = \sum_k a(k)\theta^k$ is called the series function. The definitions of these functions are shown in Table 3.10.

Table 3.10: The Power Series family of discrete distributions tab:pwrseries

Discrete Distribution	Probability function, $p(k)$	Series parameter, θ	Series function, $f(\theta)$	Series coefficient, $a(k)$
Poisson	$e^{-\lambda}\lambda^k/k!$	$\theta = \lambda$	e^θ	$1/k!$
Binomial	$\binom{n}{k} p^k (1-p)^{n-k}$	$\theta = p/(1-p)$	$(1+\theta)^n$	$\binom{n}{k}$
Negative binomial	$\binom{n+k-1}{k} p^n (1-p)^k$	$\theta = (1-p)$	$(1-\theta)^{-k}$	$\binom{n+k-1}{k}$
Geometric	$p(1-p)^k$	$\theta = (1-p)$	$(1-\theta)^{-k}$	1
Logarithmic series	$\theta^k/[-k \log(1-\theta)]$	$\theta = \theta$	$-\log(1-\theta)$	$1/k$

These relations among the discrete distribution provide the basis for graphical techniques for diagnosing the form of discrete data described later in this chapter (Section 3.5.4).

3.3 Fitting discrete distributions

In applications to discrete data such as the examples in Section 3.1, interest is often focused on how closely such data follow a particular distribution, such as the Poisson, binomial, or geometric distribution. A close fit provides for interpretation in terms of the underlying mechanism for the distribution; conversely, a bad fit can suggest the possibility for improvement by relaxing one or more of the assumptions. We examine more detailed and nuanced methods for diagnosing and testing discrete distributions in Section 3.4 and Section 3.5 below.

Fitting a discrete distribution involves three basic steps:

1. Estimating the parameter(s) of the distribution from the data, for example, p for the binomial, λ for the Poisson, n and p for the negative binomial. Typically, this is carried out by maximum likelihood methods, or a simpler method of moments, which equates sample moments (mean, variance, skewness) to those of the theoretical distribution, and solves for the parameter estimates. These methods are illustrated in Section 3.3.1.
2. From this, we can calculate the fitted probabilities, \hat{p}_k that apply for the given distribution, or equivalently, the model expected frequencies, $N\hat{p}_k$, where N is the total sample size.
3. Finally, we can calculate goodness-of-fit tests measuring the departure between the observed and fitted frequencies.

Often goodness-of-fit is examined with a classical (Pearson) **goodness-of-fit** (GOF) chi-squared test,

{sec:discrete-fit}

$$X^2 = \sum_{k=1}^K \frac{(n_k - N\hat{p}_k)^2}{N\hat{p}_k} \sim \chi^2_{(K-s-1)} , \quad (3.10) \quad \text{eq:chi2}$$

where there are K frequency classes, s parameters have been estimated from the data and \hat{p}_k is the estimated probability of each basic count, under the null hypothesis that the data follows the chosen distribution.

An alternative test statistic is the likelihood-ratio G^2 statistic,

$$G^2 = \sum_{k=1}^K n_k \log(n_k/N\hat{p}_k) , \quad (3.11)$$

when the \hat{p}_k are estimated by maximum likelihood, which also has an asymptotic $\chi^2_{(K-s-1)}$ distribution. “Asymptotic” means that these are *large sample tests*, meaning that the test statistic follows the χ^2 distribution increasingly well as $N \rightarrow \infty$. A common rule of thumb is that all *expected* frequencies should exceed one and that fewer than 20% should be less than 5.

EXAMPLE 3.12: Death by horse kick

We illustrate the basic ideas of goodness-of fit tests with the *HorseKick* data, where we expect a Poisson distribution with parameter $\lambda = \text{mean number of deaths}$. As shown in Eqn. (3.3), this is calculated as the frequency (n_k) weighted mean of the k values, here, number of deaths.

In R, such one-way frequency distributions should be converted to data frames with numeric variables. The calculation below uses `weighted.mean()` with the frequencies as weights, and finds $\lambda = 0.61$ as the mean number of deaths per corps-year.

```
> # goodness-of-fit test
> tab <- as.data.frame(HorseKicks, stringsAsFactors = FALSE)
> colnames(tab) <- c("nDeaths", "Freq")
> str(tab)

'data.frame': 5 obs. of  2 variables:
 $ nDeaths: chr  "0" "1" "2" "3" ...
 $ Freq    : int  109 65 22 3 1

> (lambda <- weighted.mean(as.numeric(tab$nDeaths), w = tab$Freq))
[1] 0.61
```

From this, we can calculate the probabilities (`phat`) of $k = 0 : 4$ deaths, and hence the expected (`exp`) frequencies in a Poisson distribution.

```
> phat <- dpois(0 : 4, lambda = lambda)
> exp <- sum(tab$Freq) * phat
> chisq <- (tab$Freq - exp)^2 / exp
>
> GOF <- data.frame(tab, phat, exp, chisq)
> GOF

  nDeaths Freq      phat      exp      chisq
1       0 109 0.5433509 108.67017 0.0010011
2       1  65 0.3314440  66.28881 0.0250573
3       2  22 0.1010904  20.21809 0.1570484
4       3   3 0.0205551   4.11101 0.3002534
5       4   1 0.0031346   0.62693 0.2220057
```

Finally, the Pearson χ^2 is just the sum of the `chisq` values and `pchisq()` is used to calculate the *p*-value of this test statistic—the probability of obtaining this χ^2 or a more extreme value if our assumption on the underlying distribution is true.

```
> sum(chisq) # chi-square value
[1] 0.70537
> pchisq(sum(chisq), df = nrow(tab) - 2, lower.tail = FALSE)
[1] 0.87194
```

The result, $\chi^2_3 = 0.70537$ shows an extremely good fit of these data to the Poisson distribution, perhaps exceptionally so.¹¹



3.3.1 R tools for discrete distributions

{sec:fitdistr}

In R, the function `fitdistr()` in the MASS is a basic work horse for fitting a variety of distributions by maximum likelihood and other methods, giving parameter estimates and standard errors. Among discrete distributions, the binomial, Poisson and geometric distributions have closed-form maximum likelihood estimates; the negative binomial distribution, parameterized by (n, μ) , is estimated iteratively by direct optimization.

These basic calculations are extended and enhanced for one-way discrete distributions in the `vcd` function `goodfit()`, which computes the fitted values of a discrete distribution (either Poisson, binomial or negative binomial) to the count data. If the parameters are not specified they are estimated either by ML or Minimum Chi-squared. `print()` and `summary()` methods for the "goodfit" objects give, respectively, a table of observed and fitted frequencies, and the Pearson and/or likelihood ratio goodness-of-fit statistics. Plotting methods for visualizing the discrepancies between observed and fitted frequencies are described and illustrated in Section 3.3.2.

{ex:saxfit}

EXAMPLE 3.13: Families in Saxony

This example uses `goodfit()` to fit the binomial to the distribution of the number of male children in families of size 12 in Saxony. Note that for the binomial, both n and p are considered as parameters, and by default n is taken as the maximum count.

```
> data(Saxony, package = "vcd")
> Sax_fit <- goodfit(Saxony, type = "binomial")
> unlist(Sax_fit$par) # estimated parameters

prob      size
0.51922 12.00000
```

So, we estimate the probability of a male in these families to be $p = 0.519$, a value that is quite close to the value found in Arbuthnot's data ($p = 0.517$).

It is useful to know that `goodfit()` returns a list structure of named components which are used by method functions for class "goodfit" objects. The `print.goodfit()` method prints the table of observed and fitted frequencies. `summary.goodfit()` calculates and prints the likelihood ratio χ^2 GOF test when the ML estimation method is used.

```
> names(Sax_fit)      # components of "goodfit" objects
[1] "observed"   "count"       "fitted"      "type"        "method"
[6] "df"          "par"
```

¹¹An exceptionally good fit occurs when the p -value for the test χ^2 statistic is so high, as to suggest that something unreasonable under random sampling might have occurred. The classic example of this is the controversy over Gregor Mendel's experiments of cross-breeding garden peas with various observed (phenotype) characteristics, where R. A. Fisher (1936a) suggested that observed frequencies of combinations like (smooth/wrinkled), (green/yellow) in a 2nd generation were uncomfortably too close to the 3 : 1 ratio predicted by genetic theory.

```
> Sax_fit          # print method

Observed and fitted values for binomial distribution
with parameters estimated by `ML'

  count observed      fitted
    0        3     0.93284
    1       24    12.08884
    2      104    71.80317
    3      286   258.47513
    4      670   628.05501
    5     1033  1085.21070
    6     1343  1367.27936
    7     1112  1265.63031
    8      829   854.24665
    9      478   410.01256
   10     181   132.83570
   11      45   26.08246
   12       7   2.34727

> summary(Sax_fit)  # summary method

Goodness-of-fit test for binomial distribution

  X^2 df P(> X^2)
Likelihood Ratio 97.007 11 6.9782e-16
```

Note that the GOF test gives a highly significant p -value (pratically zero), indicating significant lack of fit to the binomial distribution.¹² Some further analysis of this result is explored in examples below. \triangle

{ex:dicefit}

EXAMPLE 3.14: Weldon's dice

Weldon's dice data, explored in Example 3.3, are also expected to follow 9a binomial distribution, here with $p = \frac{1}{3}$. However, as given in the data set *WeldonDice*, the frequencies for counts 10–12 were grouped as “10+”. In this case, it necessary to supply the correct value of $n = 12$ as the value of the `size` parameter in the call to `goodfit()`.

```
> data(WeldonDice, package = "vcd")
> dice_fit <- goodfit(WeldonDice, type = "binomial", par = list(size = 12))
> unlist(dice_fit$par)

  prob      size
0.33769 12.00000
```

The probability of a success (a 5 or 6) is estimated as $\hat{p} = 0.3377$, not far from the theoretical value, $p = 1/3$.

```
> print(dice_fit, digits = 0)

Observed and fitted values for binomial distribution
with parameters estimated by `ML'
```

¹²A handy rule-of-thumb is to think of the ratio of χ^2/df , because, under the null hypothesis of acceptable fit, $E(\chi^2/df) = 1$, so ratios exceeding ≈ 2.5 are troubling. Here, the ratio is $97/11 = 8.8$, so the lack of fit is substantial.

```

count observed fitted
 0      185     187
 1     1149    1147
 2     3265    3216
 3     5475    5465
 4     6114    6269
 5     5194    5114
 6     3067    3042
 7     1331    1330
 8      403     424
 9      105     96
10      18     15
11      0      1
12      0      0

> summary(dice_fit)

Goodness-of-fit test for binomial distribution

X^2  df  P(> X^2)
Likelihood Ratio 11.506  9  0.2426

```

Here, we find an acceptable fit for the binomial distribution.



{ex:HKfit}

EXAMPLE 3.15: Death by horse kick

This example reproduces the calculations done “manually” in Example 3.12 above. We fit the Poisson distribution to the *HorseKicks* data by specifying `type = "poisson"` (actually, that is the default for `goodfit()`).

```

> data("HorseKicks", package = "vcd")
> HK_fit <- goodfit(HorseKicks, type = "poisson")
> HK_fit$par

$lambda
[1] 0.61

> HK_fit

Observed and fitted values for poisson distribution
with parameters estimated by `ML'

count observed     fitted
 0      109 108.67017
 1       65  66.28881
 2       22  20.21809
 3       3   4.11101
 4       1   0.62693

```

The `summary` method uses the LR test by default, so the X^2 value reported below differs slightly from the Pearson χ^2 value shown earlier.

```

> summary(HK_fit)

Goodness-of-fit test for poisson distribution

X^2  df  P(> X^2)
Likelihood Ratio 0.86822  3  0.83309

```



{ex:Fedfit}

EXAMPLE 3.16: Federalist papers

In Example 3.5 we examined the distribution of the marker word “may” in blocks of text in the *Federalist Papers* written by James Madison. A naive hypothesis is that these occurrences might follow a Poisson distribution, that is, as independent occurrences with constant probability across the 262 blocks of text. Using the same methods as above, we fit these data to the Poisson distribution:

```
> data("Federalist", package = "vcd")
> Fed_fit0 <- goodfit(Federalist, type = "poisson")
> unlist(Fed_fit0$par)

lambda
0.65649

> Fed_fit0

Observed and fitted values for poisson distribution
with parameters estimated by 'ML'

count observed      fitted
  0       156 135.891389
  1        63  89.211141
  2        29  29.283046
  3         8   6.407995
  4         4   1.051694
  5         1   0.138085
  6         1   0.015109
```

The GOF test below shows a substantial lack of fit, rejecting the assumptions of the Poisson model.

```
> summary(Fed_fit0)

Goodness-of-fit test for poisson distribution

X^2 df P(> X^2)
Likelihood Ratio 25.243 5 0.00012505
```

Mosteller and Wallace (1963) determined that the negative binomial distribution provided a better fit to these data than the Poisson. We can verify this as follows:

```
> Fed_fit1 <- goodfit(Federalist, type = "nbinomial")
> unlist(Fed_fit1$par)

size      prob
1.18633 0.64376

> summary(Fed_fit1)

Goodness-of-fit test for nbinomial distribution

X^2 df P(> X^2)
Likelihood Ratio 1.964 4  0.74238
```

Recall that the Poisson distribution assumes that the probability of a word like *may* appearing in a block of text is small and constant and that for the Poisson, $\mathcal{E}(x) = \mathcal{V}(x) = \lambda$. One interpretation of

the better fit of the negative binomial is that the use of a given word occurs with Poisson frequencies, but Madison varied its rate λ_i from one block of text to another according to a gamma distribution, allowing the variance to be greater than the mean.



3.3.2 Plots of observed and fitted frequencies

{sec:fitplot}

In the examples of the last section, we saw cases where the GOF tests showed close agreement between the observed and model-fitted frequencies, and cases where they diverged significantly, to cause rejection of a hypothesis that the data followed the specified distribution.

What is missing from such numerical summaries is any appreciation of the *details* of this statistical comparison. Plots of the observed and fitted frequencies can help to show both the shape of the theoretical distribution we have fitted and the pattern of any deviations between our data and theory.

In this section we illustrate some simple plotting tools for these purposes, using the `plot.goodfit()` method for "goodfit" objects.¹³ The left panel of Figure 3.17 shows the fit of the Poisson distribution to the Federalist papers data, using one common form of plot that is sometimes used for this purpose. In this plot, observed frequencies are shown by bars and fitted frequencies are shown by points, connected by a smooth (spline) curve.

Such a plot, however, is dominated by the largest frequencies, making it hard to assess the deviations among the smaller frequencies. To make the smaller frequencies more visible, Tukey (1977) suggest plotting the frequencies on a square-root scale, which he calls a *rootogram*. This plot is shown in the right panel of Figure 3.17.

```
> plot(Fed_fit0, scale = "raw", type = "standing")
> plot(Fed_fit0, type = "standing")
```

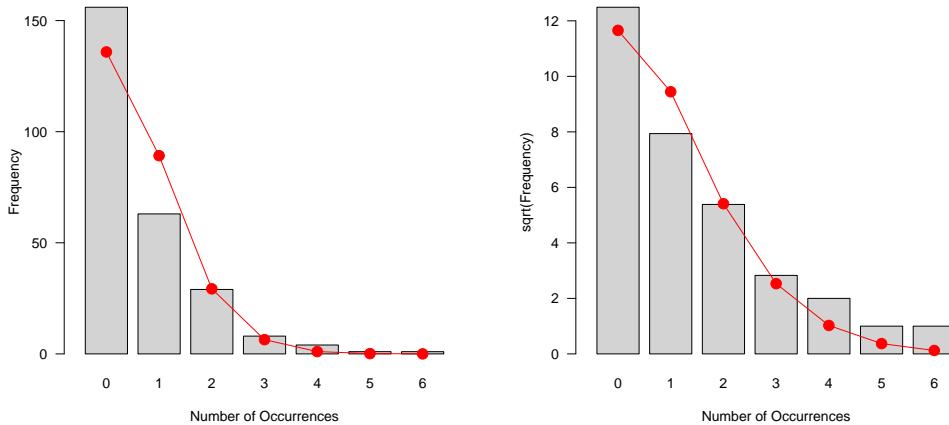


Figure 3.17: Plots for the Federalist Papers data, fitting the Poisson model. Each panel shows the observed frequencies as bars and the fitted frequencies as a smooth curve. Left: raw frequencies; right: plotted on a square root scale to emphasize smaller frequencies.^{fig:Fed0_plots1}

Additional improvements over the standard plot on the scale of raw frequencies are shown in

¹³Quantile-quantile (QQ) plots are a common alternative for the goal of comparing observed and expected values under some distribution. These plots are useful for unstructured samples, but less so when we want to also see the shape of a distribution, as is the case here.

Figure 3.18, both of which use the square root scale. The left panel moves the rootogram bars so their tops are at the expected frequencies (giving a **hanging rootogram**). This has the advantage that we can more easily judge the pattern of departures against the horizontal reference line at 0, than against the curve.

```
> plot(Fed_fit0, type = "hanging")
> plot(Fed_fit0, type = "deviation")
```

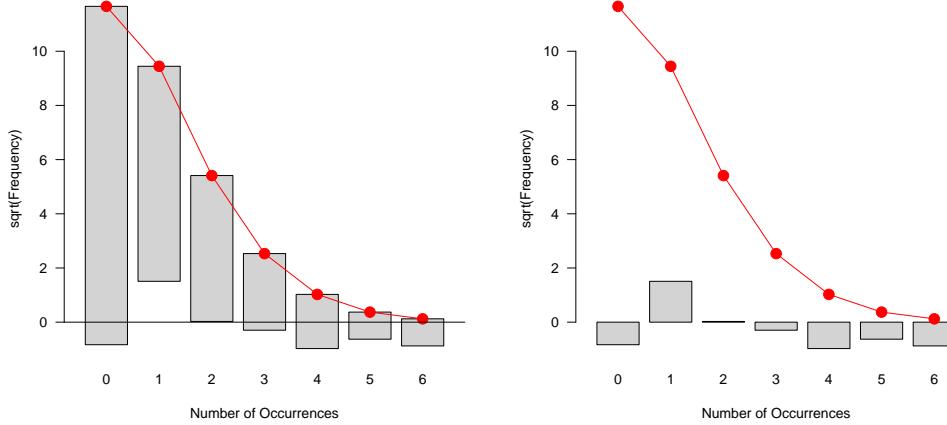


Figure 3.18: Plots for the Federalist Papers data, fitting the Poisson model. Left: hanging rootogram; right: deviation rootogram.
[Fig.Fed0_plots2]

A final variation is to emphasize the differences between the observed and fitted frequencies by drawing the bars to show the gaps between the 0 line and the (observed–expected) difference (Figure 3.18, right).

All of these plots are actually produced by the `rootogram()` function in `vcd`. The default is `type = "hanging"`, and there are many options to control the plot details.

The plots in Figure 3.17 and Figure 3.18 used the ill-fitting Poisson model on purpose to highlight how these plots show the departure between the observed and fitted frequencies. Figure 3.19 compares this with the negative binomial model, `Fed_fit1`, which we saw has a much better, and acceptable fit.

```
> plot(Fed_fit0, main = "Poisson")
> plot(Fed_fit1, main = "Negative binomial")
```

Comparing the two plots in Figure 3.19, we can see that the Poisson model underestimates the frequencies of 0 counts and the larger counts for 4–6 occurrences. The deviations for the negative binomial are small and unsystematic.

Finally, Figure 3.20 shows hanging rootograms for two atrociously bad models for the data on butterfly species in Malaya considered in Example 3.7. As we will see in Section 3.4, this long-tailed distribution is better approximated by the logarithmic series distribution, but this distribution is presently not handled by `goodfit()`.

```
> data(Butterfly, package = "vcde")
> But_fit1 <- goodfit(Butterfly, type = "poisson")
> But_fit2 <- goodfit(Butterfly, type = "nbinomial")
> plot(But_fit1, main = "Poisson")
> plot(But_fit2, main = "Negative binomial")
```

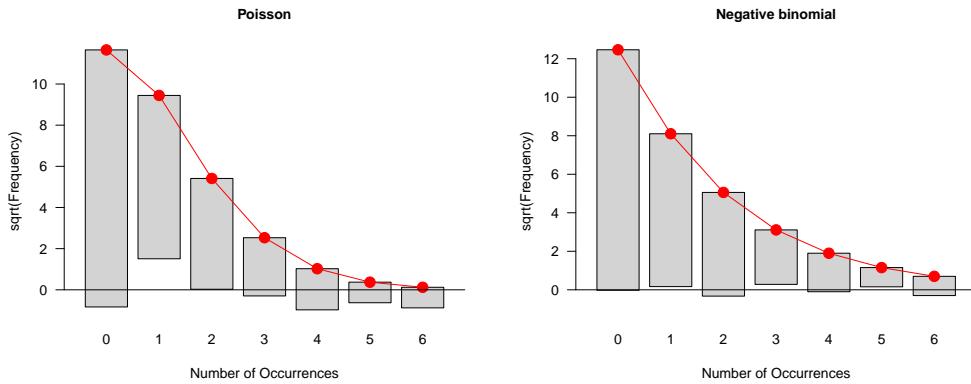


Figure 3.19: Hanging rootograms for the Federalist Papers data, comparing the Poisson and negative binomial models.
Figure 3.19: Fedo-Fedt

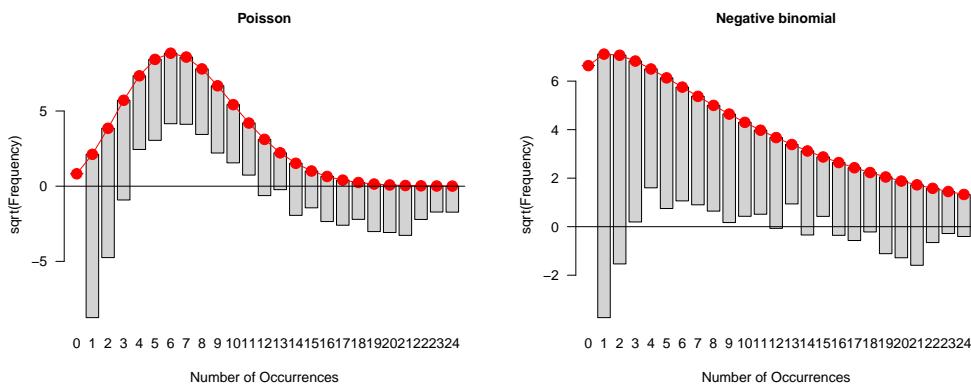


Figure 3.20: Hanging rootograms for the Butterfly data, comparing the Poisson and negative binomial models. The lack of fit for both is readily apparent.
Figure 3.20: But-But

3.4 Diagnosing discrete distributions: Ord plots

Ideally, the general form chosen for a discrete distribution should be dictated by substantive knowledge of a plausible mechanism for generating the data. When such knowledge is lacking, however, we may not know which distribution is most appropriate for some particular set of data. In these cases, the question is often turned around, so that we seek a distribution that fits well, and then try to understand the mechanism in terms of aspects of the underlying probability theory (independent trials, rare events, waiting-time to an occurrence, and so forth).

Although it is possible to fit each of several possibilities, the summary goodness-of-fit statistics can easily be influenced by one or two disparate cells, or additional (ignored or unknown) factors. One simple alternative is a plot suggested by Ord (1967) which may be used to diagnose the form of the discrete distribution.

Ord showed that a linear relationship of the form:

$$\frac{k p(k)}{p(k-1)} \equiv \frac{k n_k}{n_{k-1}} = a + b k \quad (3.12)$$

holds for each of the Poisson, binomial, negative binomial, and logarithmic series distributions, and these distributions are distinguished by the signs of the intercept, a , and slope, b , as shown in Table 3.11.

Table 3.11: Diagnostic slope and intercept for four discrete distributions. The ratios $k n_k / n_{k-1}$ plotted against k should appear as a straight line, whose slope and intercept determine the particular distribution.

Slope (b)	Intercept (a)	Distribution (parameter)	Parameter estimate
0	+	Poisson (λ)	$\lambda = a$
-	+	Binomial (n, p)	$p = b/(b-1)$
+	+	Negative binomial (n, p)	$p = 1 - b$
+	-	Log. series (θ)	$\theta = b$ $\theta = -a$

The slope, b , in Eqn. (3.12) is zero for the Poisson, negative for the binomial, and positive for the negative binomial and logarithmic series distributions; the latter two are distinguished by their intercepts. In practical applications of this idea, the details are important: how to fit the line, and how to determine if the pattern of signs are sufficient to reasonably provide a diagnosis of the distribution type.

One difficulty in applying this technique is that the number of points (distinct values of k) in the Ord plot is often small, and the sampling variances of $k n_k / n_{k-1}$ can vary enormously. A little reflection indicates that points where n_k is small should be given less weight in determining the slope of the line (and hence determining the form of the distribution). In applications it has been found that using a weighted least squares fit of $k n_k / n_{k-1}$ on k , using weights of $w_k = \sqrt{n_k - 1}$ produces reasonably good automatic diagnosis of the form of a probability distribution. Moreover, to judge whether a coefficient is positive or negative, a small tolerance is used; if none of the distributions can be classified, no parameters are estimated. Caution is advised in accepting the conclusion, because it is based on these simple heuristics.

In the `vcd` package this method is implemented in the `Ord_plot()` function. The essential ideas are illustrated using the *Butterfly* data below, which produces Figure 3.21. Note that the function returns (invisibly) the values of the intercept and slope in the weighted LS regression.

```
> ord <- Ord_plot(Butterfly,
+                   main = "Butterfly species collected in Malaya",
+                   gp = gpar(cex = 1, pch = 16))
> ord

Intercept      Slope
-0.70896     1.06082
```

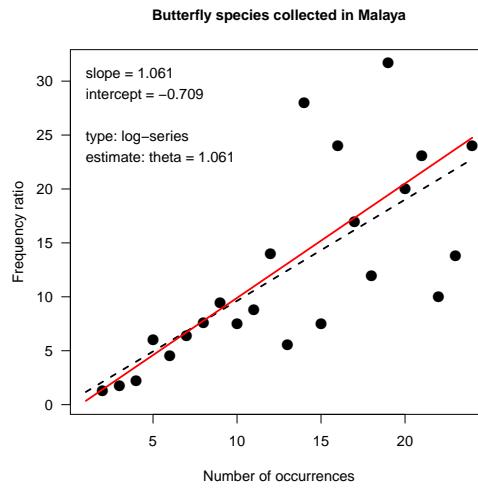


Figure 3.21: Ord plot for the `Butterfly` data. The slope and intercept in the plot correctly diagnoses the log-series distribution.^{fig:ordplot1}

In this plot, the black line shows the usual OLS regression fit of frequency, n_k on number of occurrences, k ; the red line shows the weighted least squares fit, using weights of $\sqrt{n_k} - 1$. In this case, the two lines are fairly close together, as regards their intercepts and slopes. The positive slope and negative intercept diagnoses this as a log-series distribution.

In other cases, the number of distinct points (values of k) is small, and the sampling variances of the ratios $k n_k / n_{k-1}$ can vary enormously. The following examples illustrate some other distributions and some of the details of the heuristics.

3.4.0.1 Ord plot examples

EXAMPLE 3.17: Death by horse kick

The results below show the calculations for the horse kicks data, with the frequency ratio $k n_k / n_{k-1}$ labeled y .

```
> data(HorseKicks, package = "vcd")
> nk <- as.vector(HorseKicks)
> k <- as.numeric(names(HorseKicks))
> nk1 <- c(NA, nk[-length(nk)])
> y <- k * nk / nk1
> weight <- sqrt(pmax(nk, 1) - 1)
> (ord_df <- data.frame(k, nk, nk1, y, weight))

   k  nk nk1      y  weight
1  0 109   NA    NA 10.3923
2  1   65 109 0.59633  8.0000
```

```

3 2   22   65  0.67692  4.5826
4 3     3   22  0.40909  1.4142
5 4     1     3  1.33333  0.0000

> coef(lm(y ~ k, weights = weight, data = ord_df))

(Intercept)          k
 0.656016    -0.034141

```

The weighted least squares line, with weights w_k , has a slope (-0.03) close to zero, indicating the Poisson distribution.¹⁴ The estimate $\lambda = a = .656$ compares favorably with the MLE, $\lambda = 0.610$ and the value from the Poissonness plot, shown in the following section. The call to `Ord_plot()` below produces Figure 3.22.

```

> Ord_plot(HorseKicks,
+           main = "Death by horse kicks", gp = gpar(cex = 1), pch = 16)

```

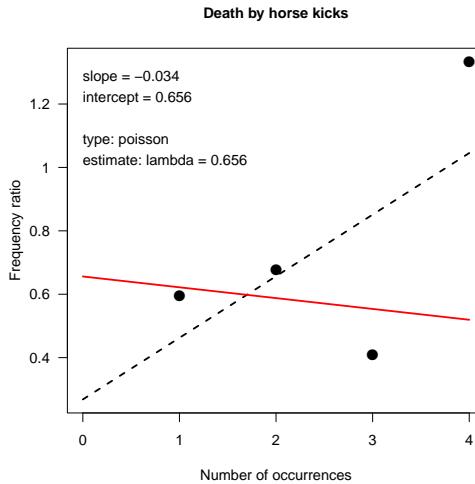


Figure 3.22: `Ord_plot` for the HorseKicks data. The plot correctly diagnoses the Poisson distribution.]

△

{ex:madison3}

EXAMPLE 3.18: Federalist papers

Figure ?? (left) shows the Ord plot for the *Federalist* data. The slope is positive, so either the negative binomial or log series are possible, according to Table 3.11. The intercept is essentially zero, which is ambiguous. However, the logarithmic series requires $b \approx -a$, so the negative binomial is a better choice. Mosteller and Wallace (1963, 1984) did in fact find a reasonably good fit to this distribution. Note that there is one apparent outlier, at $k = 6$, whose effect on the OLS line is to increase the slope and decrease the intercept. △

¹⁴The heuristic adopted in `Ord_plot()` uses a tolerance of 0.1 to decide if a coefficient is negative, zero, or positive.

```
> Ord_plot(Federalist, main = "Instances of 'may' in Federalist papers",
+           gp = gpar(cex = 1), pch = 16)
```

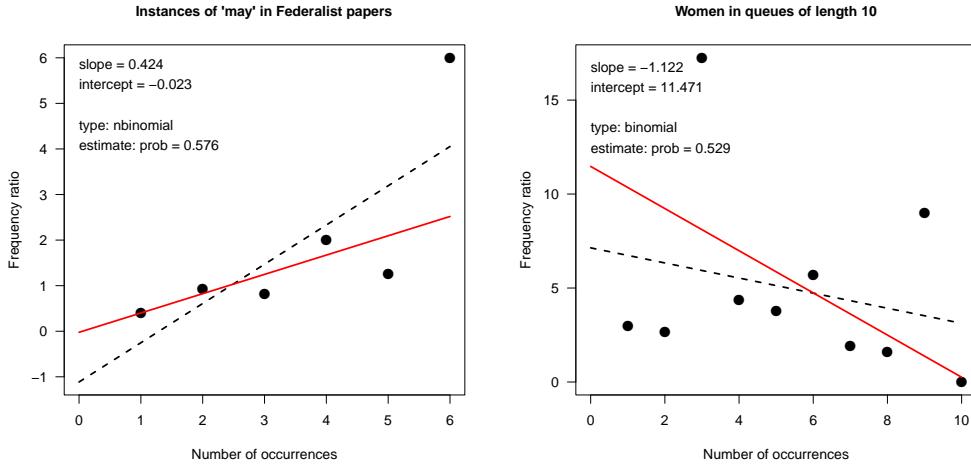


Figure 3.23: Ord plots for the Federalist (left) and WomenQueue (right) data sets.

{fig:ordplot3plot
ex:queues}

EXAMPLE 3.19: Women in queues

Jinkinson and Slater (1981), Hoaglin and Tukey (1985) give the frequency distribution of the number of females observed in 100 queues of length 10 in a London Underground station, recorded in the data set *WomenQueue* in *vcd*.

```
> data(WomenQueue, package = "vcd")
> WomenQueue

nWomen
 0   1   2   3   4   5   6   7   8   9  10
 1   3   4  23  25  19  18   5   1   1   0
```

If it is assumed that people line up independently, and that men and women are equally likely to be found in a queue (not necessarily reasonable assumptions), then the number of women out of 10 would have a (symmetric) binomial distribution with parameters $n = 10$ and $p = \frac{1}{2}$. However, there is no real reason to expect that males and females are equally likely to be found in queues in the London underground, so we may be interested in estimating p from the data and determining if a binomial distribution fits.

```
> Ord_plot(WomenQueue, main = "Women in queues of length 10",
+           gp = gpar(cex = 1), pch = 16)
```

Figure ?? (right) shows the Ord plot for these data. The negative slope and positive intercept clearly diagnose this distribution as binomial. The rough estimate of $\hat{p} = b/(1-b) = 0.53$ indicates that women are slightly more prevalent than men in these data for the London underground. \triangle

3.4.0.2 Limitations of Ord plots

Using a single simple diagnostic plot to determine one of four common discrete distributions is advantageous, but your enthusiasm should be dampened by several weaknesses:

- The Ord plot lacks resistance, since a single discrepant frequency affects the points n_k/n_{k-1} for both k and $k+1$.
- The sampling variance of $k n_k/n_{k-1}$ fluctuates widely (Hoaglin and Tukey, 1985, Jinkinson and Slater, 1981). The use of weights w_k helps, but is purely a heuristic device. The `Ord_plot()` function explicitly shows both the OLS line and the WLS line, which provides some indication of the effect of the points on the estimation of slope and intercept.

3.5 Poissonness plots and generalized distribution plots

The **Poissonness plot** (Hoaglin, 1980) is a robust plot to sensitively determine how well a one-way table of frequencies follows a Poisson distribution. It plots a quantity called a count metameter against k , designed so that the result will be points along a straight line when the data follow a Poisson distribution. When the data deviate from a Poisson, the points will be curved. Hoaglin and Tukey (1985) develop similar plots for other discrete distributions, including the binomial, negative binomial, and logarithmic series distributions. We first describe the features and construction of these plots for the Poisson distribution and then (Section 3.5.4) the extension to other distributions.

3.5.1 Features of the Poissonness plot

The Poissonness plot has the following desirable features:

- **Resistance:** a single discrepant value of n_k affects only the point at value k . (In the Ord plot it affects each of its neighbors.)
- **Comparison standard:** An approximate confidence interval can be found for each point, indicating its inherent variability and helping to judge whether each point is discrepant.
- **Influence:** Extensions of the method result in plots which show the effect of each point on the estimate of the main parameter of the distribution (λ in the Poisson).

3.5.2 Plot construction

Assume, for some fixed λ , each observed frequency, n_k equals the expected frequency, $m_k = N p_k$. Then, setting $n_k = N p_k = N e^{-\lambda} \lambda^k / k!$, and taking logs of both sides gives

$$\log(n_k) = \log N - \lambda + k \log \lambda - \log k! .$$

This can be rearranged to a linear equation in k ,

$$\phi(n_k) \equiv \log \left(\frac{k! n_k}{N} \right) = -\lambda + (\log \lambda) k . \quad (3.13)$$

The left side of Eqn. (3.13) is called the **count metameter**, and denoted $\phi(n_k)$. Hence, plotting $\phi(n_k)$ against k should give a straight line of the form $\phi(n_k) = a + bk$ with

- slope = $\log \lambda$
- intercept = $-\lambda$

when the observed frequencies follow a Poisson distribution. If the points in this plot are close enough to a straight line, then an estimate of λ may be obtained from the slope b of the line, $\hat{\lambda} = e^b$ should be reasonably close in value to the MLE of λ , $\hat{\lambda} = \bar{x}$. In this case, we might as well use the MLE as our estimate.

3.5.2.1 Leveled plot

If we have a preliminary estimate λ_0 of λ , we can use this to give a new plot where the reference line is horizontal, making comparison of the points with the line easier. In this leveled plot the vertical coordinate $\phi(n_k)$ is modified to

$$\phi'(n_k) = \phi(n_k) + \lambda_0 - k \log \lambda_0 . \quad (3.14) \quad \{\text{eq:pois-leveled}\}$$

When the data follow a Poisson distribution with parameter λ , the modified plot will have

- slope = $\log \lambda - \log \lambda_0 = \log(\lambda/\lambda_0)$
- intercept = $\lambda_0 - \lambda$

In the ideal case, where our estimate of λ_0 is close to the true λ , the line will be approximately horizontal at $\phi(n_k)' = 0$. The modified plot is particularly useful in conjunction with the confidence intervals for individual points described below.

3.5.2.2 Confidence intervals

The goal of the Poissonness plot is to determine whether the points are “sufficiently linear” to conclude that the Poisson distribution is adequate for the data. Confidence intervals for the points can help you decide, and also show the relative precision of the points in these plots.

For example, when one or two points deviate from an otherwise nearly linear relation, it is helpful to determine whether the discrepancy is consistent with chance variation. As well, we must recognize that classes with small frequencies n_k are less precise than classes with large frequencies.

Hoaglin and Tukey (1985) develop approximate confidence intervals for $\log(m_k)$ for each point in the Poissonness plot. These are calculated as

$$\phi(n_k^*) \pm h_k \quad (3.15) \quad \{\text{eq:poisCI}\}$$

where the count metameter function is calculated using a modified frequency n_k^* , defined as

$$n_k^* = \begin{cases} n_k - .8n_k - .67 & n \geq 2 \\ 1/e & n = 1 \\ \text{undefined} & n = 0 \end{cases}$$

and h_k is the half-width of the 95% confidence interval,

$$h_k = 1.96 \frac{\sqrt{1 - \hat{p}_k}}{[n_k - (.25\hat{p}_k + .47)\sqrt{n_k}]^{1/2}}$$

and $\hat{p}_k = n_k/N$.

3.5.3 The `distplot()` function

Poissonness plots (and versions for other distributions) are produced by the function `distplot()` in `vcf`. As with `Ord_plot()`, the first argument is either a vector of counts, a one-way table of frequencies of counts or a data frame or matrix with frequencies in the first column and the corresponding counts in the second column. Nearly all of the examples in this chapter use one-way tables of counts.

The `type` argument specifies the type of distribution. For `type = "poisson"`, specifying a value for `lambda = λ_0` gives the leveled version of the plot.

`{ex:horsekick4}`

EXAMPLE 3.20: Death by horse kick

The calculations for the Poissonness plot, including confidence intervals, are shown below for the *HorseKicks* data. The call to `distplot()` produces the plot in the left panel of Figure 3.24.

```
> data("HorseKicks", package="vcd")
> dp <- distplot(HorseKicks, type = "poisson",
+   xlab="Number of deaths", main="Poissonness plot: HorseKicks data")
> print(dp, digits=4)

  Counts Freq Metameter CI.center CI.width CI.lower CI.upper
1      0 109    -0.607   -0.6131   0.1305  -0.7436  -0.4827
2      1  65    -1.124   -1.1343   0.2069  -1.3412  -0.9274
3      2  22    -1.514   -1.5451   0.4169  -1.9620  -1.1281
4      3   3    -2.408   -2.6607   1.3176  -3.9783  -1.3431
5      4   1    -2.120   -3.1203   2.6887  -5.8089  -0.4316
```

In this plot, the open circles show the calculated observed values of the count $\text{Metameter} = \phi(n_k)$. The smaller filled points show the centers of the confidence intervals, $\text{CI.center} = \phi(n_k^*)$ (Eqn. (3.15)), and the dashed lines show the extent of the confidence intervals.

The fitted least squares line has a slope of -0.431 , which would indicate $\lambda = e^{-0.431} = 0.65$. This compares well with the MLE, $\lambda = \bar{x} = 0.61$.

Using $\text{lambda} = 0.61$ as below gives the leveled version shown in the right panel of Figure 3.24.

```
> # leveled version, specifying lambda
> distplot(HorseKicks, type = "poisson", lambda = 0.61,
+   xlab="Number of deaths", main="Leveled Poissonness plot")
```

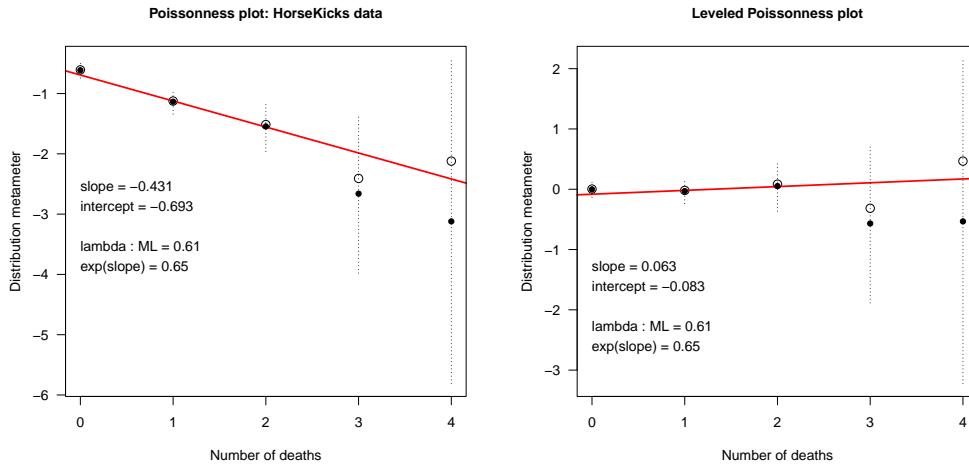


Figure 3.24: Poissonness plots for the HorseKick data. Left: standard plot; right: leveled plot. fig:distplot1

TODO: DM: In the leveled plot, the label for the slope is actually wrong, should be $\exp(\text{slope} + \log \lambda) = 0.65$

In both plots the fitted line is within the confidence intervals, indicating the adequacy of the Poisson model for these data. The widths of the intervals for $k > 2$ are graphic reminders that these observations have decreasingly low precision where the counts n_k are small.



3.5.4 Plots for other distributions

As described in Section 3.2.6, the binomial, Poisson, negative binomial, geometric, and logseries distributions are all members of the general power series family of discrete distributions. For this family, Hoaglin and Tukey (1985) develop similar plots of a count metamer against k which appear as a straight line when a data distribution follows a given family member.

The distributions which can be analyzed in this way are shown in Table 3.12, with the interpretation given to the slope and intercept in each case. For example, for the Binomial distribution, a “binomialness” plot is constructed by plotting $\log n_k^*/N(n)_k$ against k . If the points in this plot approximate a straight line, the slope is interpreted as $\log(p/(1-p))$, so the binomial parameter p may be estimated as $p = e^b/(1 + e^b)$.

Table 3.12: Plot parameters for five discrete distributions. In each case the count metamer, $\phi(n_k^*)$ is plotted against k , yielding a straight line when the data follow the given distribution.

{tab:distparms}

Distribution	Probability function, $p(k)$	Count metamer, $\phi(n_k^*)$	Theoretical Slope (b)	Theoretical Intercept (a)
Poisson	$e^{-\lambda} \lambda^k / k!$	$\log(k! n_k^* / N)$	$\log(\lambda)$	$-\lambda$
Binomial	$\binom{n}{k} p^k (1-p)^{n-k}$	$\log(n_k^* / N \binom{n}{k})$	$\log\left(\frac{p}{1-p}\right)$	$n \log(1-p)$
Negative binomial	$\binom{n+k-1}{k} p^n (1-p)^k$	$\log\left(n_k^* / N \binom{n+k-1}{k}\right)$	$\log(1-p)$	$n \log(p)$
Geometric	$p(1-p)^k$	$\log(n_k^* / N)$	$\log(1-p)$	$\log(p)$
Logarithmic series	$\theta^k / [-k \log(1-\theta)]$	$\log(k n_k^* / N)$	$\log(\theta)$	$-\log(-\log(1-\theta))$

Source: adapted from Hoaglin and Tukey (1985), Table 9-15.

Unlike the Ord plot, a different plot is required for each distribution, because the count metamer, $\phi(n_k)$, differs from distribution to distribution. Moreover, systematic deviation from a linear relationship does not indicate which distribution provides a better fit. However, the attention to robustness, and the availability of confidence intervals and influence diagnostics make this a highly useful tool for visualizing discrete distributions.

{ex:saxony-distplot}

EXAMPLE 3.21: Families in Saxony

Our analysis in Example 3.2 and Example 3.13 of the *Saxony* data showed that the distribution of male children had slightly heavier tails than the binomial, meaning the observed distribution is overdispersed. We can see this in the `goodfit()` plot shown in Figure 3.25 (left), and even more clearly in the distribution diagnostic plot produced by `distplot()` in the right panel of Figure 3.25. For a binomial distribution, we call this distribution plot a “binomialness plot”.

```
> plot(goodfit(Saxony, type="binomial", par=list(size=12)))
> distplot(Saxony, type = "binomial", size = 12,
+   xlab="Number of males")
```

The weight of evidence is thus that, as simple as the binomial might be, it is inadequate to fully explain the distribution of sex ratios in this large sample of families of 12 children. To understand this data better, it is necessary to question the assumptions of the binomial (births of males are

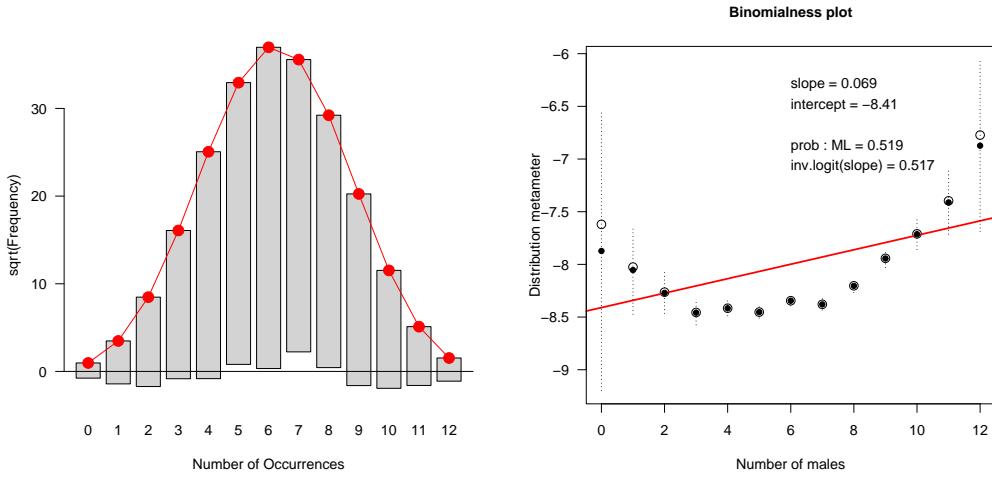


Figure 3.25: Diagnostic plots for males in Saxony families. Left: `goodfit()` plot; right: `distplot()` plot. Both plots show heavier tails than in a binomial distribution.¹⁵

independent Bernoulli trials with constant probability p) as a model for this birth distribution and/or find a more adequate model.¹⁵

`ederalist-distplot()`

EXAMPLE 3.22: Federalist papers

In Example 3.16 we carried out GOF tests for the Poisson and negative binomial models with the Federalist papers data; Figure 3.19 showed the corresponding rootogram plots. Figure 3.26 compares these two using the diagnostic plots of this section. Again the Poisson shows systematic departure from the linear relation required in the Poissonness plot, while the negative binomial model provides an acceptable fit to these data.

```
> distplot(Federalist, type = "poisson", xlab="Occurrences of 'may'")  
> distplot(Federalist, type = "nbinomial", xlab="Occurrences of 'may'")
```

△

TODO: DM: The following section assumes knowledge of GLMs, introduced in a later chapter. Either remove chapter, or move to the end after the GLM chapter.

3.6 Fitting discrete distributions as generalized linear models

{sec:fitglm}

In Section 3.2.6, we described how the common discrete distributions are all members of the general power series family. This provides the basis for the generalized distribution plots described in Section 3.5.4. Another general family of distributions—the **exponential family**—includes most of the common continuous distributions: the normal, gamma, exponential, and others, and is the basis of the class of generalized linear models (GLMs) fit by `glm()`.

¹⁵On these questions, Edwards (1958) reviews numerous other studies of these Geissler's data, and fits a so-called β -binomial model proposed by Skellam (1948), where p varies among families according to a β distribution. He concludes that there is evidence that p varies between families of the same size. One suggested explanation is that family decisions to have a further child is influenced by the balance of boys and girls among their earlier children.

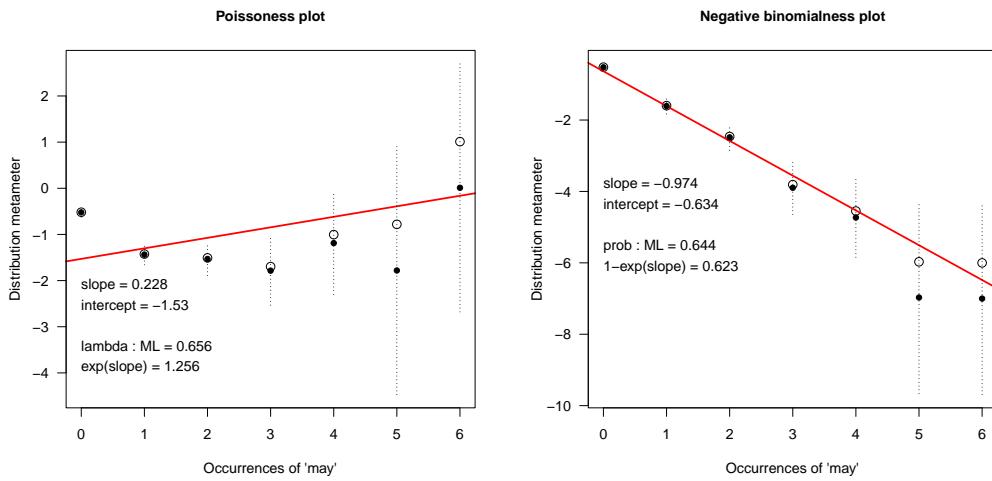


Figure 3.26: Diagnostic plots for the Federalist papers data. Left: `fig.distplots`; right: `negativebinomialness`.

A clever approach by Lindsey and Mersch (1992), Lindsey (1995, §6.1) shows how various discrete (and continuous) distributions can be fit to frequency data using generalized linear models for log frequency (which are equivalent to Poisson loglinear models). The uniform, geometric, binomial, and the Poisson distributions may all be fit easily in this way, but the idea extends to some other distributions, such as the **double binomial** distribution, that allows a separate parameter for overdispersion relative to the binomial. A clear advantage is that this method gives estimated standard errors for the distribution parameters as well as estimated confidence intervals for fitted probabilities.

The essential idea is that, for frequency data, any distribution in the exponential family may be represented by a linear model for the logarithm of the cell frequency, with a Poisson distribution for errors, otherwise known as a “Poisson loglinear regression model”. These have the form

$$\log(N\pi_k) = \text{offset} + \beta_0 + \boldsymbol{\beta}^T \mathbf{S}(k) ,$$

where N is the total frequency, π_k is the modeled probability of count k , $\mathbf{S}(k)$ is a vector of zero or more sufficient statistics for the canonical parameters of the exponential family distribution, and the offset term is a value which does not depend on the parameters.

Table 3.13 shows the sufficient statistics and offsets for several discrete distributions. See Lindsey and Mersch (1992) for further details, and definitions for the double-binomial distribution,¹⁶ and Lindsey (1995, pp. 130–133) for his analysis of the *Saxony* data using this distribution. Lindsey and Altham (1998) provide an analysis of the complete Geissler data (provided in the data set *Geissler* in *vcdExtra*) using several different models to handle overdispersion.

`{ex:saxony2}`

EXAMPLE 3.23: Families in Saxony

The binomial distribution and the double binomial can both be fit to frequency data as a Poisson regression via `glm()` using $\log(n)$ as an offset. First, we convert *Saxony* into a numeric data frame for use with `glm()`.

¹⁶In R, the double binomial distribution is implemented in the *rmutil* package, providing the standard complement of density function (`ddoublebinom()`), CDF (`pdoublebinom()`), quantiles (`qdoublebinom()`) and random generation (`rdoublebinom()`).

Table 3.13: Poisson loglinear representations for some discrete distributions

{tab:expfamily}

Distribution	Sufficient statistics	Offset
Geometric	k	
Poisson	k	$-\log(k!)$
Binomial	k	$\log \binom{n}{k}$
Double binomial	$k, k \log(k) + (n - k) \log(n - k)$	$\log \binom{n}{k}$

```
> data(Saxony, package="vcd")
> Males <- as.numeric(names(Saxony))
> Families <- as.vector(Saxony)
> Sax.df <- data.frame(Males, Families)
```

To calculate the offset for `glm()` in R, note that `choose(12, 0:12)` returns the binomial coefficients, and `lchoose(12, 0:12)` returns their logs.

```
> # fit binomial (12, p) as a glm
> Sax.bin <- glm(Families ~ Males, offset=lchoose(12, 0:12),
+                  family=poisson, data=Sax.df)
>
> # brief model summaries
> LRstats(Sax.bin)

Likelihood summary table:
      AIC BIC LR Chisq Df Pr(>Chisq)
Sax.bin 191 192     97 11    7e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

> coef(Sax.bin)

(Intercept)      Males
-0.069522     0.076898
```

As we have seen, this model fits badly. The parameter estimate for `Males`, $\beta_1 = 0.0769$ is actually estimating the logit of p , $\log p/(1-p)$, so the inverse transformation gives $\hat{p} = \frac{\exp(\beta_1)}{1+\exp(\beta_1)} = 0.5192$, as we had before.

The double binomial model can be fitted as follows. The term `YlogitY` calculates $k \log(k) + (n - k) \log(n - k)$, the second sufficient statistic for the double binomial (see Table 3.13) fitted via `glm()`.

```
> # double binomial, (12, p, psi)
> Sax.df$YlogitY <-
+   Males * log(ifelse(Males==0, 1, Males)) +
+   (12-Males) * log(ifelse(12-Males==0, 1, 12-Males))
>
> Sax.dbin <- glm(Families ~ Males + YlogitY, offset=lchoose(12, 0:12),
+                  family=poisson, data=Sax.df)
> coef(Sax.dbin)

(Intercept)      Males      YlogitY
-3.096918     0.065977     0.140205

> LRstats(Sax.bin, Sax.dbin)
```

```
Likelihood summary table:
      AIC BIC LR Chisq Df Pr(>Chisq)
Sax.bin 191 192    97.0 11     7e-16 ***
Sax.dbin 109 111   13.1 10      0.22
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

From the above, we can see that the double binomial model `Sax.dbin` with one more parameter is significantly better than the simple binomial and represents an adequate fit to the data. The table below displays the fitted values and standardized residuals for both models.

```
> results <- data.frame(Sax.df,
+                         fit.bin=fitted(Sax.bin), res.bin=rstandard(Sax.bin),
+                         fit.dbin=fitted(Sax.dbin), res.dbin=rstandard(Sax.dbin))
> print(results, digits=2)

  Males Families YlogitY fit.bin res.bin fit.dbin res.dbin
1     0        3     30  0.93    1.70     3.0    0.026
2     1       24     26 12.09    3.05    23.4    0.136
3     2      104     24 71.80    3.71   104.3   -0.036
4     3      286     23 258.48    1.87   307.8   -1.492
5     4      670     22 628.06    1.94   652.9    0.778
6     5     1033     22 1085.21   -1.87  1038.5   -0.202
7     6     1343     22 1367.28   -0.75  1264.2    2.635
8     7     1112     22 1265.63   -5.09  1185.0   -2.550
9     8      829     22  854.25   -1.03   850.1   -0.846
10    9      478     23  410.01    3.75   457.2    1.144
11   10      181     24 132.84    4.23   176.8    0.371
12   11      45      26  26.08    3.42   45.2    -0.039
13   12       7      30  2.35    2.45     6.5    0.192
```

Finally, Figure 3.27 shows the rootogram for the double binomial, which can be compared with that for the binomial model shown in Figure 3.25. We can see that the fit is now quite good, particularly in the tails. The positive coefficient for the term `YlogitY` gives additional weight in the tails.

```
> with(results, vcd::rootogram(Families, fit.dbin,
+                                xlab="Number of males"))
```



3.6.1 Covariates, overdispersion and excess zeros

All of the examples in this chapter are somewhat special, in that in each case the data consist only of a one-way frequency distribution of a basic count variable. In more general and realistic settings, there may also be one or more explanatory variables or *covariates* that influence the frequency distributions of the counts. For example, in the *Saxony* data, the number of boys in families of size 12 was aggregated over the years 1876–1885, and it is possible that any deviation from a binomial distribution could be due to variation over time or unmeasured predictors (e.g., rural vs. urban, age of parents).

This is where the generalized linear model approach introduced here (treated in detail in Chapter 9), begins to shine—because it allows such covariates to be taken into account, and then questions regarding the *form* of the distribution pertain only to the variation of the frequencies not fitted by the model. The next example illustrates what can go wrong when important predictors are omitted from the analysis.

{ex:phdpubs0}

EXAMPLE 3.24: Publications of PhD candidates

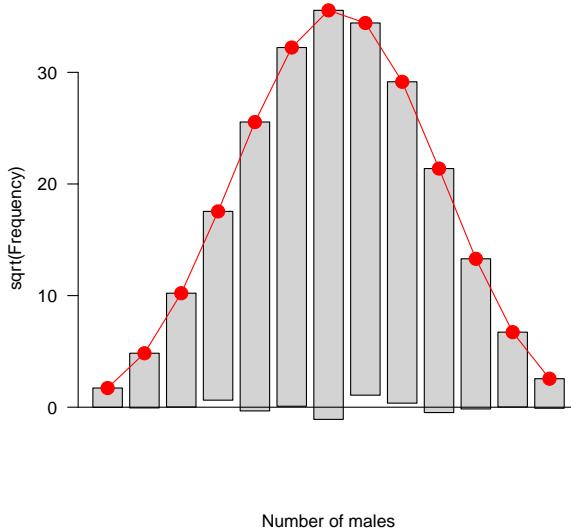


Figure 3.27: Rootogram for the double binomial model for the Saxony data. This now fits well in the tails of the distribution.
[Fig:sax-gim]

Long (1990, 1997) gave data on the number of publications by 915 doctoral candidates in biochemistry in the last three years of their PhD studies, contained in the data set *PhdPubs* in *vcdExtra*. The data set also includes information on gender, marital status, number of young children, prestige of the doctoral department and number of publications by the student's mentor. The frequency distribution of number of publications by these students is shown below.

```
> data("PhdPubs", package="vcdExtra")
> table(PhdPubs$articles)

 0   1   2   3   4   5   6   7   8   9   10  11  12  16  19 
275 246 178 84  67  27  17  12  1   2   1   1   1   2   1   1
```

The naive approach, ignoring the potential predictors is just to try fitting various probability models to this one-way distribution. Rootograms for the simpler Poisson distribution and the negative binomial that allows for overdispersion are shown in Figure 3.28.

```
> library(vcd)
> plot(goodfit(PhdPubs$articles), xlab="Number of Articles",
+       main="Poisson")
> plot(goodfit(PhdPubs$articles, type="nbinomial"), xlab="Number of Articles",
+       main="Negative binomial")
```

From these plots it is clear that the Poisson distribution doesn't fit well at all, because there is a large excess of zero counts— candidates with no publications, and most of the counts of four or more publications are larger than the Poisson model predicts. The fit of the negative binomial model in the right panel of Figure 3.28 looks much better, except that for eight or more publications, there is a systematic tendency of overfitting for 8–10 and underfitting for the observed counts of 12 or more. This lack of fit is confirmed by the formal test.

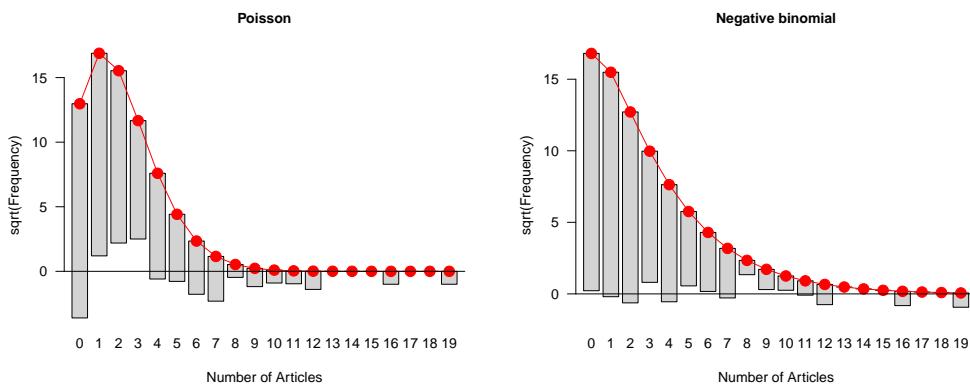


Figure 3.28: Hanging rootograms for publications by PhD candidates, comparing the Poisson and negative binomial models. The Poisson model clearly does not fit. The negative binomial is better, but still has significant lack of fit.
fig9:phdpubs-rootogram

```
> summary(goodfit(PhdPub$articles, type="nbinomial"))

Goodness-of-fit test for nbinomial distribution

X^2  df  P(> X^2)
Likelihood Ratio 31.098 12 0.0019033
```

The difficulty with this simple analysis is not only that it ignores the possible predictors of publishing by these PhD candidates, but also, by doing so, it prevents a better, more nuanced explanation of the phenomenon under study. This example is re-visited in Chapter 9, Example 9.1, where we consider generalized linear models taking potential predictors into account, as well as extended zero-inflated models allowing special consideration of zero counts. △

3.7 Chapter summary

- Discrete distributions typically involve basic *counts* of occurrences of some event occurring with varying *frequency*. The ideas and methods for one-way tables described in this chapter are building blocks for the analysis of more complex data.
- The most commonly used discrete distributions include the binomial, Poisson, negative binomial, geometric, and logarithmic series distributions. Happily, these are all members of a family called the power series distributions. Methods of fitting an observed data set to any of these distributions are described, and implemented in the `goodfit()` function.
- After fitting an observed distribution it is useful to plot the observed and fitted frequencies. Several ways of making these plots are described, and implemented in the `rootogram()` function.
- A heuristic graphical method for identifying which discrete distribution is most appropriate for a given set of data involves plotting ratios kn_k/n_{k-1} against k . These plots are constructed by the function `Ord_plot()`.
- A more robust plot for a Poisson distribution involves plotting a count metamer, $\phi(n_k)$ against

{sec:ch03-summary}

k , which gives a straight line (whose slope estimates the Poisson parameter) if the data follow a Poisson distribution. This plot provides robust confidence intervals for individual points and provides a means to assess the influence of individual points on the Poisson parameter. These plots are provided by the function `distplot()`.

- The ideas behind the Poissonness plot can be applied to the other discrete distributions.

3.8 Lab exercises

{sec:ch03-labs}

Exercise 3.1 The `Arbuthnot` data in `HistData` (Example 3.1) also contains the variable `Ratio`, giving the ratio of male to female births.

- Make a plot of `Ratio` over `Year`, similar to Figure 3.1. What features stand out? Which plot do you prefer to display the tendency for more male births?
- Plot the total number of christenings, `Males + Females` or `Total` (in 000s) over time. What unusual features do you see?

{lab:3.2}

Exercise 3.2 Use the graphical methods illustrated in Section 3.2 to plot a collection of geometric distributions for $p = 0.2, 0.4, 0.6, 0.8$, over a range of values of $k = 0, 1, \dots, 10$.

- With `xyplot()`, try the different plot formats using points connected with lines, as in Figure 3.11, or using points and lines down to the origin, as in the panels of Figure 3.13.
- Also with `xyplot()`, produce one version of a multi-line plot in a single panel that you think shows well how these distributions change with the probability p of success.
- Do the same in a multi-panel version, conditional on p .

{lab:3.3}

Exercise 3.3 Use the data set `WomenQueue` to:

- produce plots analogous to those shown in Section 3.1 (some sort of bar graph of frequencies)
- check for goodness-of-fit to the binomial distribution using the `goodfit()` methods described in Section 3.3.2.

{lab:3.4}

Exercise 3.4 Continue Example 3.13 on the distribution of male children in families in Saxony by fitting a binomial distribution, $\text{Bin}(n = 12, p = \frac{1}{2})$, specifying equal probability for boys and girls. [Hint: you need to specify both `size` and `prob` values for `goodfit()`.]

- Carry out the GOF test for this fixed binomial distribution. What is the ratio of χ^2/df ? What do you conclude?
- Test the additional lack of fit for the model $\text{Bin}(n = 12, p = \frac{1}{2})$ compared to the model $\text{Bin}(n = 12, p = \hat{p})$ where \hat{p} is estimated from the data.
- Use the `plot.goodfit()` method to visualize these two models.

{lab:3.5}

Exercise 3.5 For the `Federalist` data, the examples in Section 3.3.1 and Section 3.3.2 showed the negative binomial to provide an acceptable fit. Compare this with the simpler special case of geometric distribution, corresponding to $n = 1$.

- Use `goodfit()` to fit the geometric distribution. [Hint: use `type="nbinomial"`, but specify `size=1` as a parameter.]
- Compare the negative binomial and the geometric models statistically, by a likelihood-ratio test of the difference between these two models.
- Compare the negative binomial and the geometric models visually by hanging rootograms or other methods.

{lab:3.6}

Exercise 3.6 Mosteller and Wallace (1963, Table 2.4) give the frequencies, n_k of counts $k = 0, 1, \dots$ of other selected marker words in 247 blocks of text known to have been written by Alexander Hamilton. The data below show the occurrences of the word *upon*, that Hamilton used much more than did James Madison.

```
> count <- 0 : 5
> Freq <- c(129, 83, 20, 9, 5, 1)
```

- Read these data into R and construct a one-way table of frequencies of counts or a matrix or data frame with frequencies in the first column and the corresponding counts in the second column, suitable for use with `goodfit()`.
- Fit and plot the Poisson model for these frequencies.
- Fit and plot the negative binomial model for these frequencies.
- What do you conclude?

{lab:3.7}

Exercise 3.7 The data frame *Geissler* in the *vcdExtra* package contains the complete data from Geissler's (1889) tabulation of family sex composition in Saxony. The table below gives the number of boys in families of size 11.

boys	0	1	2	3	4	5	6	7	8	9	10	11
Freq	8	72	275	837	1,540	2,161	2,310	1,801	1,077	492	93	24

- Read these data into R
- Following Example 3.13, use `goodfit()` to fit the binomial model and plot the results. Is there an indication that the binomial does not fit these data?
- Diagnose the form of the distribution using the methods described in Section 3.4.
- Try fitting the negative binomial distribution, and use `distplot()` to diagnose whether the negative binomial is a reasonable fit.

{lab:3.8}

Exercise 3.8 The data frame *Bundesliga* gives a similar data set to that for UK soccer scores (*UKSoccer*) examined in Example 3.9, but over a wide range of years. The following lines calculate a two-way table, *BL1995*, of home-team and away-team goals for the 306 games in the year 1995.

```
> data("Bundesliga", package = "vcd")
> BL1995 <- xtabs(~ HomeGoals + AwayGoals, data = Bundesliga,
+                      subset = (Year == 1995))
> BL1995

      AwayGoals
HomeGoals 0 1 2 3 4 5 6
  0 26 16 13 5 0 1 0
  1 19 58 20 5 4 0 1
  2 27 23 20 5 1 1 1
  3 14 11 10 4 2 0 0
  4 3 5 3 0 0 0 0
  5 4 1 0 1 0 0 0
  6 1 0 0 1 0 0 0
```

- As in Example 3.9, find the one-way distributions of `HomeGoals`, `AwayGoals` and `TotalGoals` = `HomeGoals` + `AwayGoals`.
- Use `goodfit()` to fit and plot the Poisson distribution to each of these. Does the Poisson seem to provide a reasonable fit?

- (c) Use `distplot()` to assess fit of the Poisson distribution.
- (d) What circumstances of scoring goals in soccer might cause these distributions to deviate from Poisson distributions?

Exercise 3.9 ★ Repeat the exercise above, this time using the data for all years in which there was the standard number (306) of games, that is for `Year > 1965`, tabulated as shown below.

```
> BL <- xtabs(~ HomeGoals + AwayGoals, data = Bundesliga,
+             subset = (Year > 1965))
```

{lab:3.10}

Exercise 3.10 Using the data `CyclingDeaths` introduced in Example 3.6 and the one-way frequency table `CyclingDeaths.tab = table(CyclingDeaths$deaths)`,

- Make a sensible plot of the number of deaths over time. For extra credit, add a smoothed curve (e.g., using `lines(lowess(...))`).
- Test the goodness of fit of the table `CyclingDeaths.tab` to a Poisson distribution statistically using `goodfit()`.
- Continue this analysis using a `rootogram()` and `distplot()`.
- Write a one-paragraph summary of the results of these analyses and your conclusions.

{lab:3.11}

Exercise 3.11 ★ The one-way table, `Depends`, in `vcdExtra` and shown below gives the frequency distribution of the number of dependencies declared in 4,983 R packages maintained on the CRAN distribution network on January 17, 2014. That is, there were 986 packages that had no dependencies, 1,347 packages that depended on one other package, up to 2 packages that depended on 14 other packages.

TODO: Perhaps promote this table to an introductory example, leaving analysis to this exercise.

Depends	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14
# Pkgs	986	1,347	993	685	375	298	155	65	32	19	9	4	9	4	2

- (a) Make a histogram of this distribution.
- (b) Use `Ord_plot()` to see if this method can diagnose the form of the distribution.
- (c) Try to fit a reasonable distribution to describe dependencies among R packages.

{lab:3.12}

Exercise 3.12 ★ How many years does it take to get into the baseball Hall of Fame? The `Lahman` package provides a complete record of historical baseball statistics from 1871 to the present. One table, `HallOfFame`, records the history of players nominated to the Baseball Hall of Fame, and those eventually inducted. The table below, calculated in `help(HallOfFame, package="Lahman")`, records the distribution of the number of years taken (from first nomination) for the 109 players in the Hall of Fame to be inducted (1936–present). Note that `years==0` does not, and cannot, occur in this table, so the distribution is restricted to positive counts. Such distributions are called **zero-truncated distributions**. Such distributions are like the ordinary ones, but with the probability of zero being zero. Thus the other probabilities are scaled up (i.e., divided by $1 - \Pr(Y = 0)$) so they sum to 1.

years	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15
inducted	46	10	8	7	8	4	2	4	6	3	3	1	4	1	2

- (a) For the Poisson distribution, show that the zero-truncated probability function can be expressed in the form

$$\Pr\{X = k \mid k > 0\} = \frac{1}{1 - e^{-\lambda}} \times \frac{e^{-\lambda} \lambda^k}{k!} \quad k = 1, 2, \dots$$

- (b) Show that the mean is $\lambda/(1 - \exp(-\lambda))$.
(c) Enter these data into R as a one-way table, and use `goodfit()` to fit the standard Poisson distribution, as if you hadn't encountered the problem of zero truncation.

```
> remove(list=objects(pattern="\\".tab|\\".df|\\".fit"))
> .locals$ch03 <- setdiff(ls(), .globals)
> #.locals$ch03
> remove(list=.locals$ch03[sapply(.locals$ch03,function(n){!is.function(get(n))})])
```


Chapter 4

Two-way contingency tables

The analysis of two-way frequency tables concerns the association between two variables. A variety of specialized graphical displays help to visualize the pattern of association, using area of some region to represent the frequency in a cell. Some of these methods are focused on visualizing an odds ratio (for 2×2 tables), or the general pattern of association, or the agreement between row and column categories in square tables.

{ch:twoway}

4.1 Introduction

{sec:twoway-intro}

Tables are like cobwebs, like the sieve of Danaides; beautifully reticulated, orderly to look upon, but which will hold no conclusion. Tables are abstractions, and the object a most concrete one, so difficult to read the essence of.

From *Chartism* by Thomas Carlyle (1840), Chapter II, Statistics

Most methods of statistical analysis are concerned with understanding relationships or dependence among variables. With categorical variables, these relationships are often studied from data which has been summarized by a **contingency table** in table form or frequency form, giving the frequencies of observations cross-classified by two or more such variables. As Thomas Carlyle said, it is often difficult to appreciate the message conveyed in numerical tables.

This chapter is concerned with simple graphical methods for understanding the association between two categorical variables. Some examples are also presented which involve a third, **stratifying variable**, where we wish to determine if the relationship between two primary variables is the same or different for all levels of the stratifying variable. More general methods for fitting models and displaying associations for three-way and larger tables are described in Chapter 5.

In Section 4.2, We describe briefly some numerical and statistical methods for testing whether an association exists between two variables, and measures for quantifying the strength of this association. In Section 4.3 we extend these ideas to situations where the relation between two variables is of primary interest, but there are one or more background variables to be controlled.

The main emphasis, however, is on graphical methods which help to describe the *pattern* of an association between variables. Section 4.4 presents the fourfold display, designed to portray the odds ratio in 2×2 tables or a set of k such tables. **Sieve diagrams** (Section 4.5) and **association plots** (Section 4.6) are more general methods for depicting the pattern of associations in any two-way tables. When the row and column variables represent the classifications of different raters, specialized measures and visual displays for **inter-rater agreement** (Section 4.7) are particularly

{ex:berkeley1} useful. Another specialized display, a *trilinear plot* or *ternary plot*, described in Section 4.8, is designed for three-column frequency tables or compositional data. In order to make clear some of the distinctions which occur in contingency table analysis, we begin with several examples.

EXAMPLE 4.1: Berkeley admissions

Table 4.1 shows aggregate data on applicants to graduate school at Berkeley for the six largest departments in 1973 classified by admission and gender (Bickel *et al.*, 1975). See *UCBAmissions* for the complete data set. For such data we might wish to study whether there is an association between admission and gender. Are male (or female) applicants more likely to be admitted? The presence of an association might be considered as evidence of sex bias in admission practices.

Table 4.1 is an example of the simplest kind of contingency table, a 2×2 classification of individuals according to two dichotomous (binary) variables. For such a table, the question of whether there is an association between admission and gender is equivalent to asking if the proportions of males and females who are admitted to graduate school are the same, or whether the difference in proportions admitted is zero. \triangle

{tab:berk22}

Table 4.1: Admissions to Berkeley graduate programs

	Admitted	Rejected	Total	% Admit	Odds(Admit)
Males	1198	1493	2691	44.52	0.802
Females	557	1278	1835	30.35	0.437
Total	1755	2771	4526	38.78	0.633

Although the methods for quantifying association in larger tables can be used for 2×2 tables, there are specialized measures (described in Section 4.2) and graphical methods for these simpler tables.

As we mentioned in Section 1.2.4 it is often useful to make a distinction between *response*, or outcome variables, on the one hand, and possible *explanatory* or predictor variables on the other. In Table 4.1, it is natural to consider admission as the outcome, and gender as the explanatory variable. In other tables, no variable may be clearly identified as *the* outcome, or there may be several response variables, giving a multivariate problem.

{ex:haireye1}

EXAMPLE 4.2: Hair color and eye color

Table 4.2 shows data collected by Snee (1974) on the relation between hair color and eye color among 592 students in a statistics course (a two-way margin of *HairEyeColor*). Neither hair color nor eye color is considered a response in relation to the other; our interest concerns whether an association exists between them. Hair color and eye color have both been classified into four categories. Although the categories used are among the most common, they are not the only categories possible.¹ Everyday observation suggests that there probably is an association between hair color and eye color, and we will describe tests and measures of associations for larger tables in Section 4.2.3.

\triangle

{ex:mental1} If, as is suspected, hair color and eye color are associated, we would like to understand *how* they are associated. The graphical methods described later in this chapter and in Chapter 5 help reveal the pattern of associations present.

¹If students had been asked to write down their hair and eye colors, it is likely that many more than four categories of each would appear in a sample of nearly 600.

{tab:hairdat} **Table 4.2:** Hair-color eye-color data

Eye Color	Hair Color				Total
	Black	Brown	Red	Blond	
Green	5	29	14	16	64
Hazel	15	54	14	10	93
Blue	20	84	17	94	215
Brown	68	119	26	7	220
Total	108	286	71	127	592

EXAMPLE 4.3: Mental impairment and parents' SES

Srole *et al.* (1978, p. 289) gave the data in Table 4.3 on the mental health status of a sample of 1660 young New York residents in midtown Manhattan classified by their parents' socioeconomic status (SES); see *Mental* in the *vcdExtra* package. These data have also been analyzed by many authors, including Agresti (2013, §10.5.3), Goodman (1979), and Haberman (1979, p. 375).

There are five categories of SES and mental health is classified in the four categories “well”, “mild symptom formation”, “moderate symptom formation”, and “impaired”. It may be useful here to consider SES as explanatory and ask whether and how it predicts mental health status as a response.

Table 4.3: Mental impairment and parents' SES

{tab:mental-tab}

SES	Mental impairment			
	Well	Mild	Moderate	Impaired
1	64	94	58	46
2	57	94	54	40
3	57	105	65	60
4	72	141	77	94
5	36	97	54	78
6	21	71	54	71

Although there may be an overall association between these two variables, more powerful and focused tests are available when we treat these variables as *ordinal*, as we will see in Section 4.2.4.



{ex:arthritis1}

EXAMPLE 4.4: Arthritis treatment

The data in Table 4.4 compares an active treatment for rheumatoid arthritis to a placebo (Koch and Edwards, 1988), used in examples in Chapter 2 (Example 2.2). The outcome reflects whether individuals showed no improvement, some improvement, or marked improvement. Here, the outcome variable is an ordinal one, and it is probably important to determine if the relation between treatment and outcome is the same for males and females. The data set is given in case form in *Arthritis*.

This is, of course, a three-way table, with factors Treatment, Sex, and Improvement. If the relation between treatment and outcome is the same for both genders, an analysis of the Treatment by Improvement table (collapsed over sex) could be carried out. Otherwise we could perform separate analyses for men and women, or treat the combinations of treatment and sex as four levels

Table 4.4: Arthritis treatment data

{tab:arthritis}

Treatment	Sex	Improvement			Total
		None	Some	Marked	
Active	Female	6	5	16	27
	Male	7	2	5	14
Placebo	Female	19	7	6	32
	Male	10	0	1	11
Total		42	14	28	84

of a “population” variable, giving a 4×3 two-way table. These simplified approaches each ignore certain information available in an analysis of the full three-way table. \triangle

4.2 Tests of association for two-way tables

4.2.1 Notation and terminology

To establish notation, let $N = \{n_{ij}\}$ be the observed frequency table of variables A and B with r rows and c columns, as shown in Table 4.5. In what follows, a subscript is replaced by a “+” when summed over the corresponding variable, so $n_{i+} = \sum_j n_{ij}$ gives the total frequency in row i , $n_{+j} = \sum_i n_{ij}$ gives the total frequency in column j , and $n_{++} = \sum_i \sum_j n_{ij}$ is the grand total; for convenience, n_{++} is also symbolized by n .

Table 4.5: The $r \times c$ contingency table

Row Category	Column category		Total		
	1	2			
1	n_{11}	n_{12}	\cdots	n_{1c}	n_{1+}
2	n_{21}	n_{22}	\cdots	n_{2c}	n_{2+}
\vdots	\vdots	\vdots	\cdots	\vdots	\vdots
r	n_{r1}	n_{r2}	\cdots	n_{rc}	n_{r+}
Total	n_{+1}	n_{+2}	\cdots	n_{+c}	n_{++}

When each observation is randomly sampled from some population and classified on two categorical variables, A and B , we refer to the **joint distribution** of these variables, and let $\pi_{ij} = \Pr(A = i, B = j)$ denote the population probability that an observation is classified in row i , column j (or cell (ij)) in the table. Corresponding to these population joint probabilities, the cell proportions, $p_{ij} = n_{ij}/n$, give the sample joint distribution.

The row totals n_{i+} and column totals n_{+j} are called **marginal frequencies** for variables A and B respectively. These describe the distribution of each variable *ignoring* the other. For the population probabilities, the **marginal distributions** are defined analogously as the row and column totals of the joint probabilities, $\pi_{i+} = \sum_j \pi_{ij}$, and $\pi_{+j} = \sum_i \pi_{ij}$. The sample marginal proportions are, correspondingly, $p_{i+} = \sum_j p_{ij} = n_{i+}/n$, and $p_{+j} = \sum_i p_{ij} = n_{+j}/n$.

When one variable (the column variable, B , for example) is a response variable, and the other (A) is an explanatory variable, it is most often useful to examine the distribution of the response B

for each level of A separately. These define the **conditional distributions** of B , given the level of A , and are defined for the population as $\pi_{j|i} = \pi_{ij}/\pi_{i+}$.

These definitions are illustrated for the Berkeley data (Table 4.1) below, using the function `CrossTable()`.

```
> Berkeley <- margin.table(UCBAdmissions, 2:1)
> library(gmodels)
> CrossTable(Berkeley, prop.chisq=FALSE, prop.c=FALSE, format="SPSS")

Cell Contents
-----+
| Count |
| Row Percent |
| Total Percent |
-----+

Total Observations in Table: 4526

| Admit
Gender | Admitted | Rejected | Row Total |
-----+-----+-----+-----+
Male | 1198 | 1493 | 2691 |
| 44.519% | 55.481% | 59.456% |
| 26.469% | 32.987% | |
-----+-----+-----+-----+
Female | 557 | 1278 | 1835 |
| 30.354% | 69.646% | 40.544% |
| 12.307% | 28.237% | |
-----+-----+-----+-----+
Column Total | 1755 | 2771 | 4526 |
-----+-----+-----+-----+
```

The output shows the joint frequencies, n_{ij} , and joint sample percentages, $100 \times p_{ij}$, in the first row within each table cell. The second row in each cell (“Row percent”) gives the conditional percentage of admission or rejection, $100 \times p_{j|i}$ for males and females separately. The row and column labelled “Total” give the marginal frequencies, n_{i+} and n_{+j} , and marginal percentages, p_{i+} and p_{+j} .

4.2.2 2 by 2 tables

The 2×2 contingency table of applicants to Berkeley graduate programs in Table 4.1 may be regarded as an example of a **cross-sectional study**. The total of $n = 4,526$ applicants in 1973 has been classified by both gender and admission status. Here, we would probably consider the total n to be fixed, and the cell frequencies n_{ij} , $i = 1, 2; j = 1, 2$ would then represent a single **multinomial sample** for the cross-classification by two binary variables, with probabilities cell p_{ij} , $i = 1, 2; j = 1, 2$ such that

$$p_{11} + p_{12} + p_{21} + p_{22} = 1 .$$

The basic null hypothesis of interest for a multinomial sample is that of independence. Are admission and gender independent of each other?

Alternatively, if we consider admission the response variable, and gender an explanatory variable, we would treat the numbers of male and female applicants as fixed and consider the cell frequencies to represent two independent **binomial samples** for a binary response. In this case, the null hypothesis is described as that of homogeneity of the response proportions across the levels of the explanatory variable.

{sec:twoway-twobytwo}

{sec:twoway-odds}

Measures of association are used to quantify the strength of association between variables. Among the many measures of association for contingency tables, the **odds ratio** is particularly useful for 2×2 tables, and is a fundamental parameter in several graphical displays and models described later. Other measures of strength of association for 2×2 tables are described in Stokes *et al.* (2000, Chapter 2) and Agresti (1996, §2.2).

For a binary response, where the probability of a “success” is π , the **odds** of a success is defined as

$$\text{odds} = \frac{\pi}{1 - \pi} .$$

Hence, odds = 1 corresponds to $\pi = 0.5$, or success and failure equally likely. When success is more likely than failure $\pi > 0.5$, and the odds > 1 ; for instance, when $\pi = 0.75$, odds = $.75/.25 = 3$, so a success is three times as likely as a failure. When failure is more likely, $\pi < 0.5$, and the odds < 1 ; for instance, when $\pi = 0.25$, odds = $.25/.75 = \frac{1}{3}$.

The odds of success thus vary *multiplicatively* around 1. Taking logarithms gives an equivalent measure which varies *additively* around 0, called the **log odds** or **logit**:

$$\text{logit}(\pi) \equiv \log(\text{odds}) = \log\left(\frac{\pi}{1 - \pi}\right) . \quad (4.1)$$

The logit is symmetric about $\pi = 0.5$, in that $\text{logit}(\pi) = -\text{logit}(1 - \pi)$. The following lines calculate the odds and log odds for a range of probabilities. As you will see in Chapter 7, the logit transformation of a probability is fundamental in logistic regression.

```
> p <- c(0.05, .1, .25, .50, .75, .9, .95)
> odds <- p / (1-p)
> logodds <- log(odds)
> data.frame(p, odds, logodds)

      p      odds logodds
1 0.05 0.052632 -2.9444
2 0.10 0.111111 -2.1972
3 0.25 0.333333 -1.0986
4 0.50 1.000000  0.0000
5 0.75 3.000000  1.0986
6 0.90 9.000000  2.1972
7 0.95 19.000000 2.9444
```

A binary response for two groups gives a 2×2 table, with Group as the row variable, say. Let π_1 and π_2 be the success probabilities for Group 1 and Group 2. The **odds ratio**, θ , is just the ratio of the odds for the two groups:

$$\text{odds ratio} \equiv \theta = \frac{\text{odds}_1}{\text{odds}_2} = \frac{\pi_1/(1 - \pi_1)}{\pi_2/(1 - \pi_2)} .$$

Like the odds itself, the odds ratio is always non-negative, between 0 and ∞ . When $\theta = 1$, the distributions of success and failure are the same for both groups (so $\pi_1 = \pi_2$); there is no association between row and column variables, or the response is independent of group. When $\theta > 1$, Group 1 has a greater success probability; when $\theta < 1$, Group 2 has a greater success probability.

Similarly, the odds ratio may be transformed to a log scale, to give a measure which is symmetric about 0. The **log odds ratio**, symbolized by ψ , is just the difference between the logits for Groups 1 and 2:

$$\text{log odds ratio} \equiv \psi = \log(\theta) = \log\left[\frac{\pi_1/(1 - \pi_1)}{\pi_2/(1 - \pi_2)}\right] = \text{logit}(\pi_1) - \text{logit}(\pi_2) .$$

Independence corresponds to $\psi = 0$, and reversing the rows or columns of the table merely changes the sign of ψ .

For sample data, the **sample odds ratio** is the ratio of the sample odds for the two groups:

$$\hat{\theta} = \frac{p_1/(1-p_1)}{p_2/(1-p_2)} = \frac{n_{11}/n_{12}}{n_{21}/n_{22}} = \frac{n_{11}n_{22}}{n_{12}n_{21}} . \quad (4.2)$$

The sample estimate $\hat{\theta}$ in Eqn. (4.2) is the maximum likelihood estimator of the true θ . The sampling distribution of $\hat{\theta}$ is asymptotically normal as $n \rightarrow \infty$, but may be highly skewed in small to moderate samples.

Consequently, inference for the odds ratio is more conveniently carried out in terms of the log odds ratio, whose sampling distribution is more closely normal, with mean $\psi = \log(\theta)$, and asymptotic standard error (ASE)

$$\text{ASE}_{\log(\theta)} \equiv \hat{s}(\hat{\psi}) = \left\{ \frac{1}{n_{11}} + \frac{1}{n_{12}} + \frac{1}{n_{21}} + \frac{1}{n_{22}} \right\}^{1/2} = \left\{ \sum \sum n_{ij}^{-1} \right\}^{1/2} \quad (4.3) \quad \text{(eq:aselogtheta)}$$

A large-sample $100(1 - \alpha)\%$ confidence interval for $\log(\theta)$ may therefore be calculated as

$$\log(\theta) \pm z_{1-\alpha/2} \text{ASE}_{\log(\theta)} = \hat{\psi} \pm z_{1-\alpha/2} \hat{s}(\hat{\psi})$$

where $z_{1-\alpha/2}$ is the cumulative normal quantile with $1 - \alpha/2$ in the lower tail. Confidence intervals for θ itself are obtained by exponentiating the end points of the interval for $\psi = \log(\theta)$,²

$$\exp \left(\hat{\psi} \pm z_{1-\alpha/2} \hat{s}(\hat{\psi}) \right) .$$

4.2.3 Larger tables: Overall analysis

For two-way tables overall tests of association can be carried out using `assocstats()`. If the data set has more than two factors (as in the Arthritis Treatment data), the other factors will be ignored (and collapsed) if not included when the table is constructed. This simplified analysis may be misleading if the excluded factors interact with the factors used in the analysis.

{sec:twoway-overall}

{ex:arthritis2}

EXAMPLE 4.5: Arthritis treatment

Since the main interest is in the relation between Treatment and Improved, an overall analysis (which ignores Sex) can be carried out by creating a two-way table with `xtabs()` as shown below.

```
> data("Arthritis", package="vcd")
> Art.tab <- xtabs(~Treatment + Improved, data=Arthritis)
> Art.tab

Improved
Treatment None Some Marked
Placebo    29     7     7
Treated    13     7    21

> round(100*prop.table(Art.tab, margin=1), 2)

Improved
Treatment None Some Marked
Placebo 67.44 16.28 16.28
Treated 31.71 17.07 51.22
```

²Note that $\hat{\theta}$ is 0 or ∞ if any $n_{ij} = 0$. Haldane (1955) and Gart and Zweifel (1967) showed that improved estimators of θ and $\psi = \log(\theta)$ are obtained by replacing each n_{ij} by $[n_{ij} + \frac{1}{2}]$ in Eqn. (4.2) and Eqn. (4.3). This adjustment is preferred in small samples, and required if any zero cells occur. In large samples, the effect of adding 0.5 to each cell becomes negligible.

The row proportions show a clear difference in the outcome for the two groups: For those given the placebo, 67% reported no improvement; in the treated group, 51% reported marked improvement. χ^2 tests and measures of association are provided by `assocstats()` as shown below:

```
> assocstats(Art.tab)

          X^2 df  P(> X^2)
Likelihood Ratio 13.530 2 0.0011536
Pearson         13.055 2 0.0014626

Phi-Coefficient : 0.394
Contingency Coeff.: 0.367
Cramer's V       : 0.394
```

△

4.2.4 Tests for ordinal variables

For $r \times c$ tables, more sensitive tests than the test for general association (independence) are available if either or both of the row and column variables are ordinal. Generalized **Cochran-Mantel-Haenszel tests** (Landis *et al.*, 1978) which take the ordinal nature of a variable into account are provided by the `CMHtest()` in `vcdExtra`. These tests are based on assigning numerical scores to the table categories; the default (table) scores treat the levels as equally spaced. They generally have higher power when the pattern of association is determined by the order of an ordinal variable.

EXAMPLE 4.6: Mental impairment and parents' SES

We illustrate these tests using the data on mental impairment and SES introduced in Example 4.3, where both variables can be considered ordinal.

```
> data(Mental, package="vcdeExtra")
> mental.tab <- xtabs(Freq ~ ses + mental, data=Mental)
> assocstats(mental.tab)      # standard chisq tests

          X^2 df  P(> X^2)
Likelihood Ratio 47.418 15 3.1554e-05
Pearson         45.985 15 5.3458e-05

Phi-Coefficient : 0.166
Contingency Coeff.: 0.164
Cramer's V       : 0.096

> CMHtest(mental.tab)        # CMH tests

Cochran-Mantel-Haenszel Statistics for ses by mental

          AltHypothesis Chisq Df     Prob
cor      Nonzero correlation 37.2  1 1.09e-09
cmeans   Col mean scores differ 40.3  5 1.30e-07
rmeans   Row mean scores differ 40.7  3 7.70e-09
general  General association 46.0 15 5.40e-05
```

In this data set, all four tests show a highly significant association. However, the `cor` test for nonzero correlation uses only one degree of freedom, whereas the test of general association requires 15 df.

△

The four tests differ in the types of departure from independence they are sensitive to:

General Association When the row and column variables are both nominal (unordered) the only alternative hypothesis of interest is that there is *some* association between the row and column variables. The CMH test statistic is similar to the (Pearson) Chi-Square and Likelihood Ratio Chi-Square in the result from `assocstats()`; all have $(r - 1)(c - 1)$ df.

Row Mean Scores Differ If the column variable is ordinal, assigning scores to the column variable produces a mean for each row. The association between row and column variables can be expressed as a test of whether these means differ over the rows of the table, with $r - 1$ df. This is analogous to the Kruskal-Wallis non-parametric test (ANOVA based on rank scores).

Column Mean Scores Differ Same as the above, assigning scores to the row variable.

Nonzero Correlation (Linear association) When *both* row and column variables are ordinal, we could assign scores to both variables and compute the correlation (r). The CMH χ^2 is equal to $(N - 1)r^2$, where N is the total sample size. The test is most sensitive to a pattern where the row mean score changes linearly over the rows.

4.2.5 Sample CMH Profiles

Two contrived examples may make the differences among these tests more apparent. Visualizations of the patterns of association reinforces the aspects to which the tests are most sensitive, and introduces the sieve diagram described more fully in Section 4.5.

{sec:Sample}

4.2.5.1 General Association

The table below exhibits a general association between variables A and B , but no difference in row means or linear association. The row means are calculated by assigning integer scores, $b_i = i$ to the column categories. Figure 4.1(left) shows the pattern of association in this table graphically, as a sieve diagram (described in Section 4.5).

	b1	b2	b3	b4	b5	Total	Mean
a1	0	15	25	15	0	55	3.0
a2	5	20	5	20	5	55	3.0
a3	20	5	5	5	20	55	3.0
Total	25	40	35	40	25	165	3.0

This is reflected in the `CMHtest()` output shown below. **TODO: Something wrong here: does `CMHtest()` get rows/cols mixed up? Would be nice to calculate col means also.**

```
> CMHtest(cmhdemo1)

Cochran-Mantel-Haenszel Statistics

          AltHypothesis Chisq Df    Prob
cor      Nonzero correlation  0.0  1 1.00e+00
cmeans   Col mean scores differ  0.0  2 1.00e+00
rmeans   Row mean scores differ 72.2  4 7.78e-15
general  General association  91.8  8 2.01e-16
```

The chi-square values for non-zero correlation and different row mean scores are exactly zero because the row means are all equal. Only the general association test shows that A and B are associated.

```
> sieve(cmhdemo1, shade=TRUE, main="General association",
+       gp = shading_sieve(interpolate = 0, lty = c("solid", "longdash")))
> sieve(cmhdemo2, shade=TRUE, main="Linear association",
+       gp = shading_sieve(interpolate = 0, lty = c("solid", "longdash")))
```

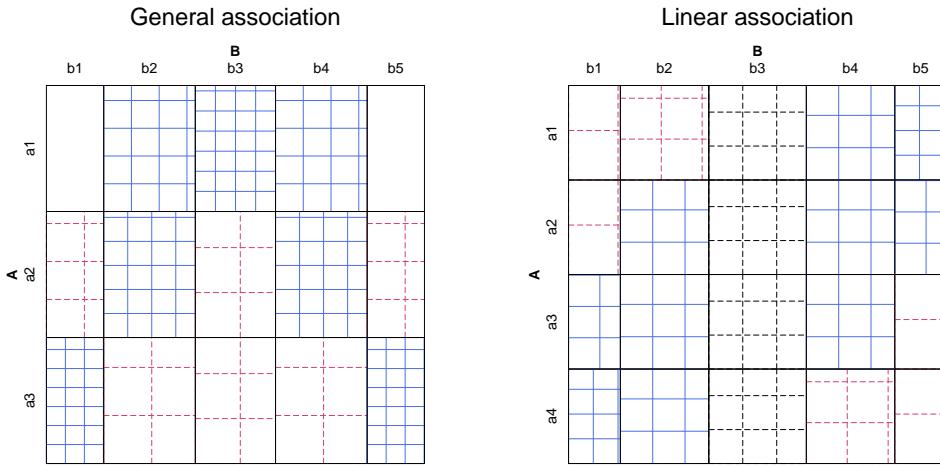


Figure 4.1: Sieve diagrams for two patterns of association: Left: General association; right: Linear association

4.2.5.2 Linear Association

The table below contains a weak, non-significant general association, but significant row mean differences and linear associations. The unstructured test of general association would therefore lead to the conclusion that no association exists, while the tests taking ordinal factors into account would conclude otherwise. Note that the largest frequencies shift towards lower levels of B as the level of variable A increases. See Figure 4.1(right) for a visual representation of this pattern.

	b1	b2	b3	b4	b5	Total	Mean
a1	2	5	8	8	8	31	3.48
a2	2	8	8	8	5	31	3.19
a3	5	8	8	8	2	31	2.81
a4	8	8	8	5	2	31	2.52
Total	17	29	32	29	17	124	3.00

Note that the χ^2 -values for the row-means and non-zero correlation tests from `CMHtest()` are very similar, but the correlation test is more highly significant since it is based on just one degree of freedom.

```
> CMHtest(cmhdemo2)

Cochran-Mantel-Haenszel Statistics

          AltHypothesis Chisq Df     Prob
cor      Nonzero correlation 10.6  1 0.00111
cmeans   Col mean scores differ 10.7  3 0.01361
rmeans   Row mean scores differ 11.4  4 0.02241
general  General association 13.4 12 0.34064
```

The difference in sensitivity and power among these tests for categorical data is analogous to the difference between general ANOVA tests and tests for linear trend (contrasts) in experimental designs with quantitative factors: The more specific test has greater power, but is sensitive to a narrower range of departures from the null hypothesis. The more focused tests for ordinal factors are a better bet when we believe that the association depends on the ordered nature of the factor levels.

4.3 Stratified analysis

An overall analysis ignores other variables (like sex), by collapsing over them. In the *Arthritis* data, it is possible that the treatment is effective only for one gender, or even that the treatment has opposite effects for men and women. If so, pooling over the ignored variable(s) can be seriously misleading.

A **stratified analysis** controls for the effects of one or more background variables. This is similar to the use of a blocking variable in an ANOVA design. Tests for association can be obtained by applying a function (`assocstats()`, `CMHtest()`) over the levels of the stratifying variables.

{sec:twoway-strat}

{ex:arthritis3}

EXAMPLE 4.7: Arthritis treatment

The statements below request a stratified analysis of the arthritis treatment data with CMH tests, controlling for sex. Essentially, the analysis is carried out separately for males and females.

The table `Art.tab2` is constructed as a three-way table, with `sex` as the last dimension.

```
> Art.tab2 <- xtabs(~Treatment + Improved + Sex, data=Arthritis)
> Art.tab2

, , Sex = Female

Improved
Treatment None Some Marked
Placebo    19     7     6
Treated     6     5    16

, , Sex = Male

Improved
Treatment None Some Marked
Placebo    10     0     1
Treated     7     2     5
```

`assocstats()` only applies to two-way tables, so we use `apply()` to run it for each level of `Sex`. `CMHtest()` is designed for such stratified tables, and uses all dimensions after the first two as strata.

```
> apply(Art.tab2, MARGIN=3, FUN=assocstats)

$Female
          X^2 df P(> X^2)
Likelihood Ratio 11.731  2 0.0028362
Pearson          11.296  2 0.0035242

Phi-Coefficient : 0.438
Contingency Coeff.: 0.401
Cramer's V       : 0.438

$Male
          X^2 df P(> X^2)
Likelihood Ratio 5.8549  2 0.053532
```

```
Pearson          4.9067  2 0.086003
Phi-Coefficient : 0.443
Contingency Coeff.: 0.405
Cramer's V       : 0.443
```

Note that even though the strength of association (ϕ -coefficient) is similar in the two groups, the χ^2 tests show significance for females, but not for males. This is true even using the more powerful CMH tests below, treating Treatment as ordinal. The reason is that there were more than twice as many females as males in this sample.

```
> CMHtest(Art.tab2)

\$`Sex:Female`
Cochran-Mantel-Haenszel Statistics for Treatment by Improved
in stratum Sex:Female

      AltHypothesis Chisq Df   Prob
cor      Nonzero correlation 10.9  1 0.000944
cmeans   Col mean scores differ 10.9  1 0.000944
rmeans   Row mean scores differ 11.1  2 0.003878
general  General association  11.1  2 0.003878

\$`Sex:Male`
Cochran-Mantel-Haenszel Statistics for Treatment by Improved
in stratum Sex:Male

      AltHypothesis Chisq Df   Prob
cor      Nonzero correlation  3.71  1 0.0540
cmeans   Col mean scores differ 3.71  1 0.0540
rmeans   Row mean scores differ 4.71  2 0.0949
general  General association  4.71  2 0.0949

> apply(Art.tab2, 3, sum)

Female    Male
      59     25
```

△

4.3.1 Assessing homogeneity of association

{sec:twoway-homog} In a stratified analysis it is often crucial to know if the association between the primary table variables is the same over all strata. For $2 \times 2 \times k$ tables this question reduces to whether the odds ratio is the same in all k strata. The `vcd` package implements Woolf's test (Woolf, 1995) in `woolf_test()` for this purpose.

For larger n -way tables, this question is equivalent to testing whether the association between the primary variables, A and B , say, is the same for all levels of the stratifying variables, C, D, \dots

In the case of a 3-way table, this can be stated as the *loglinear model* of no three-way association, $[AB][AC][BC]$. This notation (described in Section 8.2) lists only the high-order association terms in a linear model for log frequency.

EXAMPLE 4.8: Berkeley admissions

{ex:berkeley1a} Here we illustrate the use of Woolf's test for the `UCBAdmissions` data. The test is significant, indicating that the odds ratios cannot be considered equal across departments. We will see why when we visualize the data by department in the next section.

```
> woolf_test(UCBAdmissions)

Woolf-test on Homogeneity of Odds Ratios (no 3-Way
assoc.)

data: UCBAdmissions
X-squared = 17.902, df = 5, p-value = 0.003072
```

△

{ex:arthrit4}

EXAMPLE 4.9: Arthritis treatment

For the arthritis data, homogeneity means that there is no three-way Treatment * Improved * Sex association. That is, the association between treatment and outcome (`improve`) is the same for both men and women. This hypothesis can be stated as the loglinear model,

$$[\text{SexTreatment}] \quad [\text{SexImproved}] \quad [\text{TreatmentImproved}] . \quad (4.4) \quad \text{(eq:STO2)}$$

Such tests can be carried out most conveniently using `loglm()` in the **MASS** package. The model formula uses the standard R notation `()^2` to specify all terms of order 2.

```
> library(MASS)
> loglm(~ (Treatment + Improved + Sex)^2, data=Art.tab2)

Call:
loglm(formula = ~ (Treatment + Improved + Sex)^2, data = Art.tab2)

Statistics:
      X^2  df P(> X^2)
Likelihood Ratio 1.7037  2  0.42663
Pearson          1.1336  2  0.56735
```

Even though we found in the CMH analysis above that the association between Treatment and Improved was stronger for females than males, the analysis using `loglm()` is clearly non-significant, so we cannot reject homogeneity of association. △

4.4 Fourfold display for 2 x 2 tables

The **fourfold display** is a special case of a **radial diagram** (or “polar area chart”) designed for the display of 2×2 (or $2 \times 2 \times k$) tables (Fienberg, 1975, Friendly, 1994a,c). In this display the frequency n_{ij} in each cell of a fourfold table is shown by a quarter circle, whose radius is proportional to $\sqrt{n_{ij}}$, so the area is proportional to the cell count. The fourfold display is similar to a pie chart in using segments of a circle to show frequencies. It differs from a pie chart in that it keeps the angles of the segments constant and varies the radius, whereas the pie chart varies the angles and keeps the radius constant.

The main purpose of this display is to depict the sample odds ratio, $\hat{\theta} = (n_{11}/n_{12}) \div (n_{21}/n_{22})$. An association between the variables ($\theta \neq 1$) is shown by the tendency of diagonally opposite cells in one direction to differ in size from those in the opposite direction, and the display uses color or shading to show this direction. Confidence rings for the observed θ allow a visual test of the hypothesis of independence, $H_0 : \theta = 1$. They have the property that (in a standardized display) the rings for adjacent quadrants overlap *iff* the observed counts are consistent with the null hypothesis.

{sec:twoway-fourfold}

{ex:berkeley2}

EXAMPLE 4.10: Berkeley admissions

Figure 4.2(left) shows the basic, unstandardized fourfold display for the Berkeley admissions

data (Table 4.1). Here, the area of each quadrant is proportional to the cell frequency, shown numerically in each corner. The odds ratio is proportional to the product of the areas shaded dark, divided by the product of the areas shaded light. The sample odds ratio, Odds(Admit|Male) / Odds(Admit|Female) is 1.84 (see Example 4.8) indicating that males were nearly twice as likely to be admitted.

```
> fourfold(Berkeley, std="ind.max")      # unstandardized
> fourfold(Berkeley, margin=1)           # equating gender
```

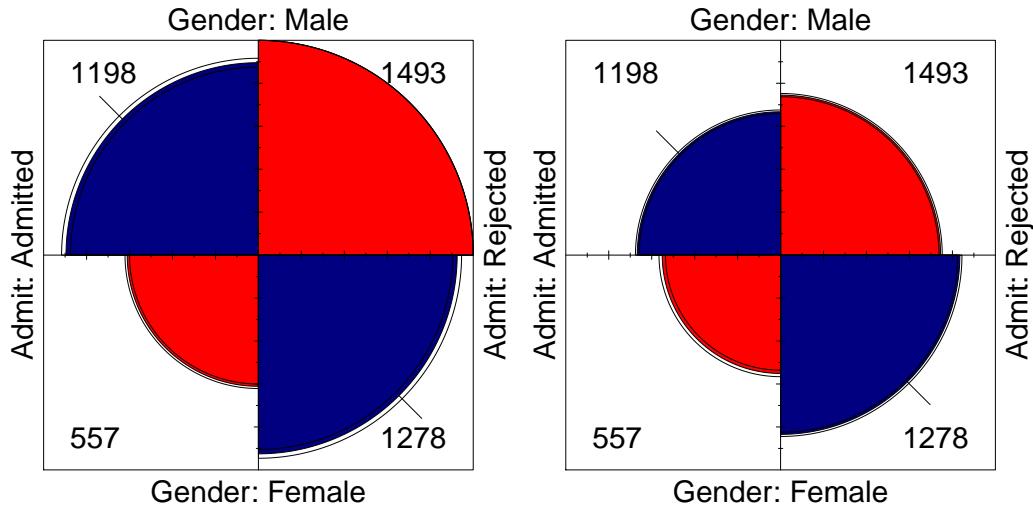


Figure 4.2: Fourfold displays for the Berkeley admission data. Left: fig:berk-fourfold unstandardized; right: equating the proportions of males and females

However, it is difficult to make these visual comparisons because there are more men than women, and because the proportions admitted and rejected are unequal. In the unstandardized display the confidence bands have no interpretation as a test of $H_0 : \theta = 1$.

{tab:berkrow}

Table 4.6: Admissions to Berkeley graduate programs, Frequencies and Row Percentages

	Frequencies		Row Percents	
	Admitted	Rejected	Admitted	Rejected
Males	1198	1493	44.52	55.48
Females	557	1278	30.35	69.65

The data in a 2×2 table can be standardized to make these visual comparisons easier. Table 4.6 shows the Berkeley data with the addition of row percentages (which equate for the number of men and women applicants) indicating the proportion of each gender accepted and rejected. We see that 44.52% of males were admitted, while only 30.35% of females were admitted. Moreover, the row percentages have the same odds ratio as the raw data: $44.52 \times 69.65 / 30.35 \times 55.48 = 1.84$. Figure 4.2(right) shows the fourfold display where the area of each quarter circle is proportional to these row percentages.

With this standardization, the confidence rings have the property that the confidence rings for each upper quadrant will overlap with those for the quadrant below it if the odds ratio does not

differ from 1.0. (Details of the calculation of confidence rings are described in the next section.) No similar statement can be made about the corresponding left and right quadrants, however, because the overall rate of admission has not been standardized.

As a final step, we can standardize the data so that *both* table margins are equal, while preserving the odds ratio. Each quarter circle is then drawn to have an area proportional to this standardized cell frequency. This makes it easier to see the association between admission and sex without being influenced by the overall admission rate or the differential tendency of males and females to apply. With this standardization, the four quadrants will align (overlap) horizontally and vertically when the odds ratio is 1, regardless of the marginal frequencies. The fully standardized display, which is usually the most useful form, is shown in Figure 4.3.

```
> fourfold(Berkeley) # standardize both margins
```

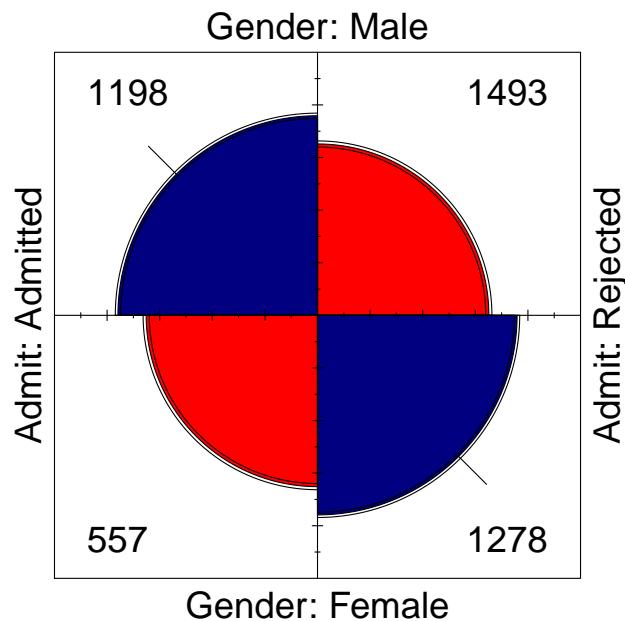


Figure 4.3: Fourfold display for Berkeley admission data with margins for gender and admission equated. The area of each quadrant shows the standardized frequency in each cell.
Fig.Berk-fourfold3



These displays also use color (blue) and diagonal tick marks to show the direction of positive association. The visual interpretation (also conveyed by area) is that males are more likely to be accepted, females more likely to be rejected.

The quadrants in Figure 4.3 do not align and the 95% confidence rings around each quadrant do not overlap, indicating that the odds ratio differs significantly from 1—putative evidence of gender bias. The very narrow width of the confidence rings gives a visual indication of the precision of the data—if we stopped here, we might feel quite confident of this conclusion.

4.4.1 Confidence rings for odds ratio

Confidence rings for the fourfold display are computed from a confidence interval for θ , whose endpoints can each be mapped into a 2×2 table. Each such table is then drawn in the same way as the data.

The interval for θ is most easily found by considering the distribution of $\hat{\psi} = \log \hat{\theta}$, whose standard error may be estimated by Eqn. (4.3). Then an approximate $1 - \alpha$ confidence interval for ψ is given by

$$\hat{\psi} \pm \hat{s}(\hat{\psi}) z_{1-\alpha/2} = \{\hat{\psi}_l, \hat{\psi}_u\},$$

as described in Section 4.2.2. The corresponding limits for the odds ratio θ are $\{\exp(\hat{\psi}_l), \exp(\hat{\psi}_u)\}$. For the data shown in Figure 4.3, $\hat{\psi} = \log \hat{\theta} = .6104$, and $\hat{s}(\hat{\psi}) = 0.0639$, so the 95%, limits for θ are $\{1.624, 2.087\}$, as shown by the calculations below. The same result is returned by `confint()` for an "oddsratio" object.

```
> summary(oddstratio(Berkeley))
   Log Odds Ratio Std. Error z value Pr(>|z|)
[1,]      0.6104     0.0639    9.55 <2e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
> exp(.6103 + c(-1, 1) * qnorm(.975) * 0.06398)
[1] 1.6240 2.0869
> confint(oddstratio(Berkeley, log=FALSE))
   lwr      upr
[1,] 1.6244 2.0867
```

Now consider how to find a 2×2 table whose frequencies correspond to the odds ratios at the limits of the confidence interval. A table standardized to equal row and column margins can be represented by the 2×2 matrix with entries

$$\begin{bmatrix} p & (1-p) \\ (1-p) & p \end{bmatrix},$$

whose odds ratio is $\theta = p^2/(1-p)^2$. Solving for p gives $p = \sqrt{\theta}/(1 + \sqrt{\theta})$. The corresponding frequencies can then be found by adjusting the standardized table to have the same row and column margins as the data. The results of these computations which generate the confidence rings in Figure 4.3 are shown in Table 4.7.

{tab:berkodds}

Table 4.7: Odds ratios and equivalent tables for 95% confidence rings for the Berkeley data.

		Odds Ratio	Standardized Table		Equivalent Frequencies	
Lower limit	1.624	0.560	0.440	1167.1	587.9	
		0.440	0.560	1523.9	1247.1	
Data	1.841	0.576	0.424	1198.0	557.0	
		0.424	0.576	1493.0	1278.0	
Upper limit	2.087	0.591	0.409	1228.4	526.6	
		0.409	0.591	1462.6	1308.4	

4.4.2 Stratified analysis for $2 \times 2 \times k$ tables

In a $2 \times 2 \times k$ table, the last dimension often corresponds to “strata” or populations, and it is typically of interest to see if the association between the first two variables is homogeneous across

c:twoway-fourstrat

strata. For such tables, simply make one fourfold panel for each stratum. The standardization of marginal frequencies is designed to allow easy visual comparison of the pattern of association when the marginal frequencies vary across two or more populations

The admissions data shown in Figure 4.2 and Figure 4.3 were actually obtained from six departments—the six largest at Berkeley (Bickel *et al.*, 1975). To determine the source of the apparent sex bias in favor of males, we make a new plot, Figure 4.4, stratified by department.

```
> # fourfold display
> UCB <- aperm(UCBAmissions, c(2, 1, 3))
> fourfold(UCB, mffrow=c(2,3))
```

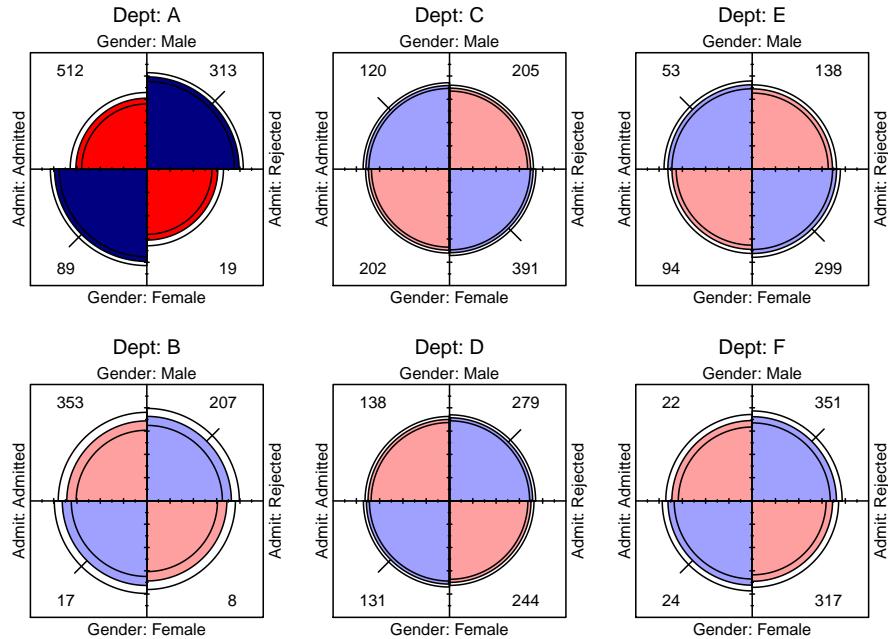


Figure 4.4: Fourfold displays for Berkeley admissions data, stratified by department. The more intense shading for Dept. A indicates a significant association.
Fig.Berk-Fourfold4

Surprisingly, Figure 4.4 shows that, for five of the six departments, the odds of admission is approximately the same for both men and women applicants. Department A appears to differ from the others, with women approximately $2.86 (= (313/19)/(512/89))$ times as likely to gain admission. This appearance is confirmed by the confidence rings, which in Figure 4.4 are joint³ 95% intervals for θ_c , $c = 1, \dots, k$.

This result, which contradicts the display for the aggregate data in Figure 4.2, is a nice example of **Simpson's paradox**⁴, and illustrates clearly why an overall analysis of a three- (or higher-) way table can be misleading. The resolution of this contradiction can be found in the large differences in admission rates among departments. Men and women apply to different departments differentially,

³For multiple-strata plots, `fourfold()` by default adjusts the significance level for multiple testing, using Holm's (1979) method provided by `p.adjust()`.

⁴Simpson's paradox (Simpson, 1951) occurs in a three-way table, $[A, B, C]$, when the marginal association between two variables, A, B collapsing over C differs in *direction* from the partial association $A, B|C = c_k$ at the separate levels of C . Strictly speaking, Simpson's paradox would require that for all departments separately the odds ratio $\theta_k < 1$ (which occurs for Departments A, B, D, and F in Figure 4.4) while in the aggregate data $\theta > 1$.

and in these data women happen to apply in larger numbers to departments that have a low acceptance rate. The aggregate results are misleading because they falsely assume men and women are equally likely to apply in each field.⁵

4.4.2.1 Visualization principles

TODO: Move this to Ch. 1 An important principle in the display of large, complex data sets is **controlled comparison**—we want to make comparisons against a clear standard, with other things held constant. The fourfold display differs from a pie chart in that it holds the angles of the segments constant and varies the radius. An important consequence is that we can quite easily compare a series of fourfold displays for different strata, since corresponding cells of the table are always in the same position. As a result, an array of fourfold displays serve the goals of comparison and detection better than an array of pie charts.

Moreover, it allows the observed frequencies to be standardized by equating either the row or column totals, while preserving the design goal for this display—the odds ratio. In Figure 4.4, for example, the proportion of men and women, and the proportion of accepted applicants were equated visually in each department. This provides a clear standard which also greatly facilitates controlled comparison.

Another principle is **visual impact**—we want the important features of the display to be easily distinguished from the less important (Tukey, 1993). Figure 4.4 distinguishes the one department for which the odds ratio differs significantly from 1 by shading intensity, even though the same information can be found by inspection of the confidence rings.

{ex:wheeze1}

EXAMPLE 4.11: Breathlessness and wheeze in coal miners

The various ways of standardizing a collection of 2×2 tables allows visualizing relations with different factors (row percentages, column percentages, strata totals) controlled. However, different kinds of graphs can speak more eloquently to other questions by focusing more directly on the odds ratio.

Agresti (2002, Table 9.8) cites data from Ashford and Sowden (1970) on the association between two pulmonary conditions, breathlessness and wheeze, in a large sample of coal miners. The miners are classified into age groups, and the question treated by Agresti is whether the association between these two symptoms is homogeneous over age. These data are available in the *CoalMiners* data in vcd, a $2 \times 2 \times 9$ frequency table. The first group, aged 20–24 has been omitted from these analyses.

```
> data("CoalMiners", package="vcd")
> CM <- CoalMiners[, 2:9]
> ftable(CM, row.vars = 3)
```

	Age	Breathlessness		Wheeze		B		NoB	
		W	NoW	W	NoW	W	NoW	W	NoW
	25–29	23	9	105	1654				
	30–34	54	19	177	1863				
	35–39	121	48	257	2357				
	40–44	169	54	273	1778				
	45–49	269	88	324	1712				
	50–54	404	117	245	1324				
	55–59	406	152	225	967				
	60–64	372	106	132	526				

The question of interest can be addressed by displaying the odds ratio in the 2×2 tables with the margins of breathlessness and wheeze equated (i.e., with the default `std='margins'` option),

⁵This explanation ignores the possibility of structural bias against women, e.g., lack of resources allocated to departments that attract women applicants.

which gives the graph shown in Figure 4.5. Although the panels for all age groups show an overwhelmingly positive association between these two symptoms, one can also (by looking carefully) see that the strength of this association declines with increasing age.

```
> fourfold(CM, mfcoll = c(2, 4))
```

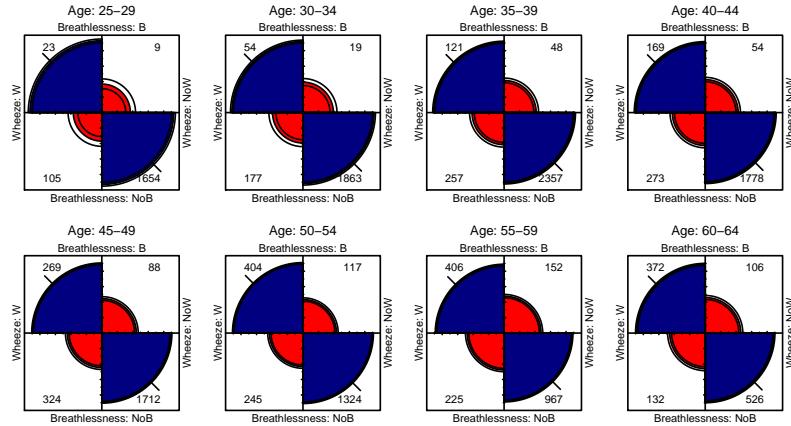


Figure 4.5: Fourfold display for CoalMiner data, both margins equated fig:coalminer1

However, note that the pattern of change over age is somewhat subtle compared to the dominant positive association within each panel. When the goal is to display how the odds ratio varies with a quantitative factor such as age, it is often better to simply calculate and plot the odds ratio directly.

The `oddsratio()` function in `vcd` calculates odds ratios for $2 \times 2 (\times k)$ tables. By default, it returns the log odds. Use the option `log=FALSE` to get the odds ratios themselves. It is easy to see that the (log) odds ratios decline with age.

```
> oddsratio(CM)

 25-29   30-34   35-39   40-44   45-49   50-54   55-59   60-64
3.6953 3.3983 3.1407 3.0147 2.7820 2.9264 2.4406 2.6380

> oddsratio(CM, log=FALSE)

 25-29   30-34   35-39   40-44   45-49   50-54   55-59   60-64
40.256 29.914 23.119 20.383 16.152 18.660 11.480 13.985
```

When the analysis goal is to understand how the odds ratio varies with a stratifying factor (which could be a quantitative variable), it is often better to plot the odds ratio directly.

The lines below use the `plot()` method for "oddsratio" objects. This produces a line graph of the log odds ratio against the stratum variable, together with confidence interval error bars. In addition, because age is a quantitative variable, we can calculate, and display the fitted relation for a linear model relating `lodds` to age. Here, we try using a quadratic model (`poly(age, 2)`) mainly to see if the trend is nonlinear.

```
> lodds <- oddsratio(CM)
> plot(lodds, lwd=2, cex=1.25, pch=16,
+       xlab = "Age Group",
+       main = "Breathlessness and Wheeze in Coal Miners")
> age <- seq(25, 60, by = 5)
> mod <- lm(lodds ~ poly(age, 2))
> lines(fitted(mod), col = "red", lwd=2)
```

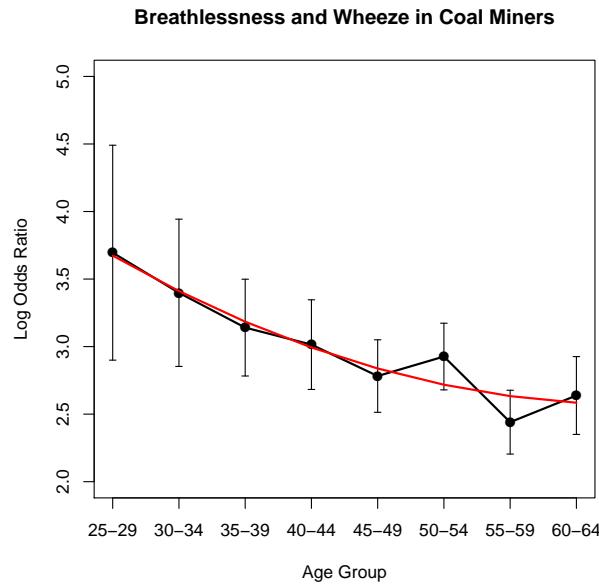


Figure 4.6: Log odds plot for the CoalMiners data. The smooth curve shows a quadratic fit to age.^{fig:coalminer3}

In Figure 4.6, it appears that the decline in the log odds ratio levels off with increasing age. One virtue of fitting the model in this way is that we can test the additional contribution of the quadratic term, which turns out to be insignificant.

```
> summary(mod)

Call:
lm(formula = lodds ~ poly(age, 2))

Residuals:
 25-29   30-34   35-39   40-44   45-49   50-54   55-59   60-64 
 0.0219 -0.0128 -0.0438  0.0213 -0.0558  0.2085 -0.1928  0.0534 

Coefficients:
            Estimate Std. Error t value Pr(>|t|)    
(Intercept) 3.0045    0.0473  63.46  1.8e-08 ***  
poly(age, 2)1 -1.0080    0.1339  -7.53  0.00066 ***  
poly(age, 2)2  0.2304    0.1339   1.72  0.14594  
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 0.134 on 5 degrees of freedom
Multiple R-squared:  0.923, Adjusted R-squared:  0.892 
F-statistic: 29.8 on 2 and 5 DF,  p-value: 0.00167
```



4.5 Sieve diagrams

{sec:twoway-sieve}

The wise ones fashioned speech with their thought, sifting it as grain is sifted through a sieve.

Buddha

For two- (and higher-) way contingency tables, the design principles of perception, detection, and comparison (see Chapter 1) suggest that we should try to show the observed frequencies in relation to what we would expect those frequencies to be under a reasonable null model—for example, the hypothesis that the row and column variables are unassociated.

To this end, several schemes for representing contingency tables graphically are based on the fact that when the row and column variables are independent, the estimated expected frequencies, m_{ij} , are products of the row and column totals (divided by the grand total).

$$m_{ij} = \frac{n_i n_j}{n_{++}} .$$

Then, each cell can be represented by a rectangle whose area shows the observed cell frequency, n_{ij} , expected frequency, m_{ij} , or deviation (residual) from independence, $n_{ij} - m_{ij}$. Visual attributes (color, shading) of the rectangles can be used to highlight the pattern of association.

For example, for any two-way table, the expected frequencies under independence can be represented by rectangles whose widths are proportional to the total frequency in each column, n_{+j} , and whose heights are proportional to the total frequency in each row, n_{i+} ; the area of each rectangle is then proportional to m_{ij} . Figure 4.7 (left) shows the expected frequencies for the hair and eye color data (Table 4.2), calculated using `independence_table()` in `vcd`.

```
> haireye <- margin.table(HairEyeColor, 1:2)
> expected = independence_table(haireye)
> round(expected, 1)

      Eye
Hair   Brown  Blue Hazel Green
  Black 40.1 39.2 17.0 11.7
  Brown 106.3 103.9 44.9 30.9
  Red   26.4 25.8 11.2  7.7
  Blond 47.2 46.1 20.0 13.7
```

Figure 4.7 (left) simply represents the model—what the frequencies would be if hair color and eye color were independent—not the data. Note, however, that the rectangles are cross-ruled so that the number of boxes in each (counting up the fractional bits) equals the expected frequency with which the cell is labeled, and moreover, the rulings are equally spaced in all cells. Hence, cross-ruling the cells to show the observed frequency would give a data display which implicitly compares observed and expected frequencies as shown in Figure 4.7 (right).

Riedwyl and Schüpbach 1983, 1994 proposed a *sieve diagram* (later called a *parquet diagram*) based on this principle. In this display the area of each rectangle is always proportional to expected frequency but observed frequency is shown by the number of squares in each rectangle, as in Figure 4.7 (right).

Hence, the difference between observed and expected frequency appears as variations in the density of shading. Cells whose observed frequency n_{ij} exceeds the expected m_{ij} appear denser than average. The pattern of positive and negative deviations from independence can be more easily seen by using color, say, red for negative deviations, and blue for positive.⁶

{ex:haireye2}

⁶Positive residuals are also shown by solid lines, negative residuals by broken lines, so that they may still be distinguished in monochrome versions.

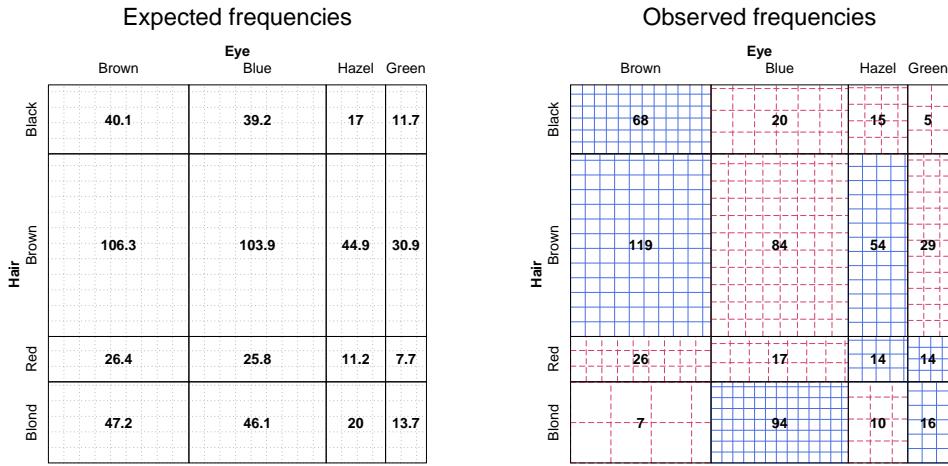


Figure 4.7: Sieve diagrams for Hair Eye color data. Left: expected frequencies shown in cells as numbers and the number of boxes; right: observed frequencies shown in cells.
Fig. 4.7 sieve

EXAMPLE 4.12: Hair color and eye color

The sieve diagram for hair color and eye color shown in Figure 4.7 (right) can be interpreted as follows: The pattern of color and shading shows the high frequency of blue-eyed blonds and people with brown eyes and dark hair. People with hazel eyes are also more likely to have red or brown hair, and those with green eyes more likely to have red or blond hair, than would be observed under independence. \triangle

{ex:vision1}

EXAMPLE 4.13: Visual acuity

In World War II, all workers in the U.K. Royal Ordnance factories were given test of visual acuity (unaided distance vision) of their left and right eyes on a 1 (high) to 4 (low) scale. The dataset *VisualAcuity* in *vcd* gives the results for 10,719 workers (3242 men, 7477 women) aged 30-39.

Figure 4.8 shows the sieve diagram for data from the larger sample of women (Kendall and Stuart (1961, Table 33.5), Bishop *et al.* (1975, p. 284)). The *VisualAcuity* data is a frequency data frame and we first convert it to table form (*VA.tab*), a $4 \times 4 \times 2$ table to re-label the variables and levels. **Done:** Make this an exercise in Ch. 2

```
> # re-assign names/dimnames
> data("VisualAcuity", package="vcd")
> VA.tab <- xtabs(Freq ~ right + left + gender, data=VisualAcuity)
> dimnames(VA.tab)[1:2] <- list(c("high", 2, 3, "low"))
> names(dimnames(VA.tab))[1:2] <- paste(c("Right", "Left"), "eye grade")
> #str(VA.tab)
```

```
> sieve(VA.tab[, , "female"], shade=TRUE)
```

The diagonal cells show the obvious: people tend to have the same visual acuity in both eyes, and there is strong lack of independence. The off diagonal cells show a more subtle pattern that suggests symmetry—the cells below the diagonal are approximately equally dense as the corresponding cells above the diagonal. Moreover, the relatively consistent pattern on the diagonals $\pm 1, \pm 2, \dots$ away

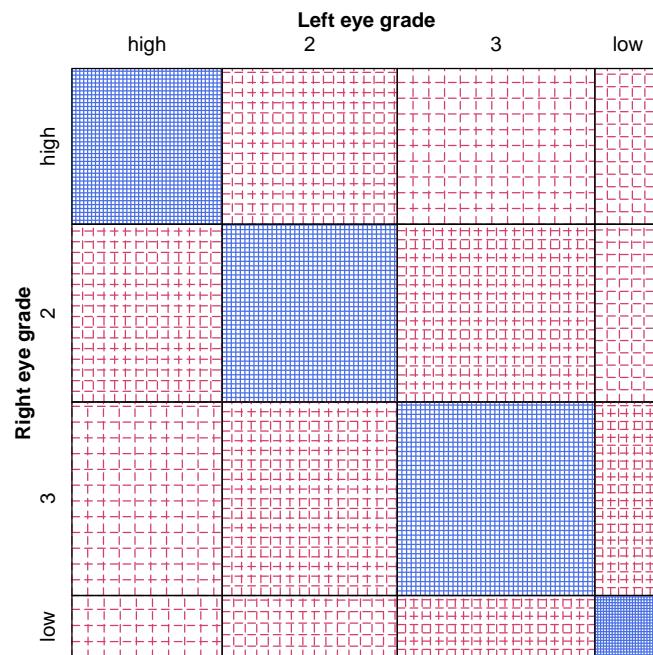


Figure 4.8: Vision classification for 7477 women in Royal Ordnance factories. The high frequencies in the diagonal cells indicate the main association, but a subtler pattern also appears in the symmetric off-diagonal cells.^{119:VA-sieve2}

from the main diagonals suggests that the association may be explained in terms of the *difference* in visual acuity between the two eyes.

These suggestions can be tested by fitting intermediate models between the null model of independence (which fits terribly) and the saturated model (which fits perfectly), as we shall see later in this book. A model of *quasi-independence*, for example (see Example 8.8 in Chapter 8) ignores the diagonal cells and tests whether independence holds for the remainder of the table. The *symmetry* model for a square table allows association, but constrains the expected frequencies above and below the main diagonal to be equal. Such models provide a way of testing *specific* explanatory models that relate to substantive hypotheses and what we observe in our visualizations. These and other models for square tables are discussed further in Section 8.7. △

4.5.1 Larger tables: The strucplot framework

The implementation of sieve diagrams in vcd is far more general than illustrated in the examples above. For one thing, the `sieve` function has a formula method, which allows one to specify the variables in the display as a model formula. For example, for the `VisualAcuity` data, a plot of the (marginal) frequencies for left and right eye grades pooling over gender can be obtained with the call below (this plot is not shown).

```
> sieve(Freq ~ right + left, data = VisualAcuity, shade=TRUE)
```

More importantly, sieve diagrams are just one example of the *strucplot framework*, a general system for visualizing n -way frequency tables in a hierarchical way. We describe this framework in more detail in Section 5.3 in context of mosaic displays. For now, we just illustrate the extension of the formula method to provide for conditioning variables. In the call below, the formula `Freq ~ right + left | gender` means to produce a separate block in the plot for the levels of gender.⁷

```
> sieve(Freq ~ right + left | gender, data = VisualAcuity, shade=TRUE)
```

In Figure 4.9, the relative sizes of the blocks for the conditioning variable (`gender`) show the much larger number of women than men in this data. Within each block, color and density of the box rules shows the association of left and right acuity, and it appears that the pattern for men is similar to that observed for women. The methods described in Section 4.3.1 can be used to test the hypothesis of homogeneity of association, and loglinear models described in Chapter 8 provide specific tests of hypotheses of *symmetry*, *quasi-independence* and other models for structured associations.

EXAMPLE 4.14: Berkeley admissions

This example illustrates some additional flexibility of sieve plots with the strucplot framework, using the Berkeley admissions data. The left panel of Figure 4.10 shows the sieve diagrams for the relation between department and admission, conditioned by gender. It can easily be seen that (a) overall, there were more male applicants than female; (b) there is a moderately similar pattern of observed > expected (blue) for males and females.

```
> # conditioned on gender
> sieve(UCBAdmissions, shade=TRUE, condvar='Gender')
> # three-way table, Department first, with cell labels
> UCB <- aperm(UCBAdmissions, c(3,1,2))
> dimnames(UCB)[[3]] <- c("M", "F") # abbreviate for display
> sieve(UCB, shade=TRUE, pop=FALSE)
> labeling_cells(text = UCB, gp_text = gpar(fontface = 2))(UCB)
```

⁷An equivalent plot, but one labeled more nicely, as in Figure 4.8 can be produced from the `VA.tab` table using `sieve(VA.tab, shade=TRUE, condvar='gender')`.

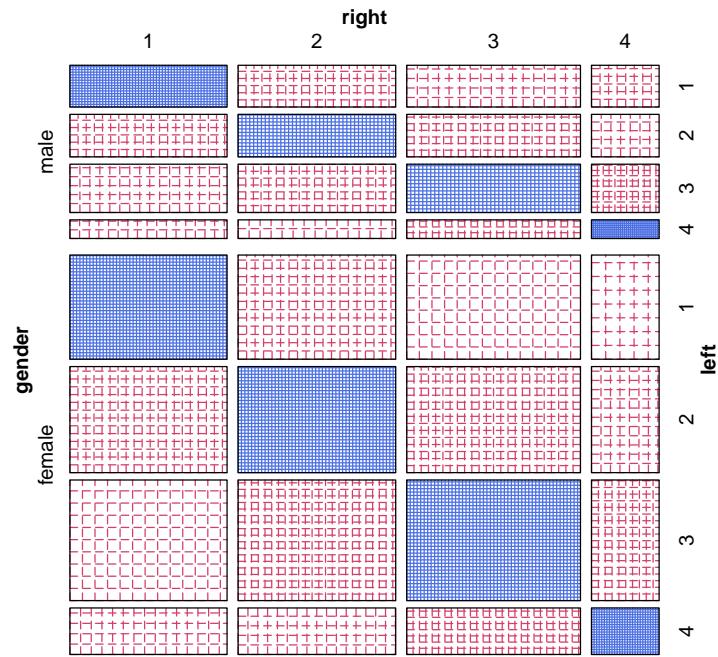


Figure 4.9: Sieve diagram for the three-way table of VisualAcuity, conditioned on gender.^{fig:VA-sieve3}

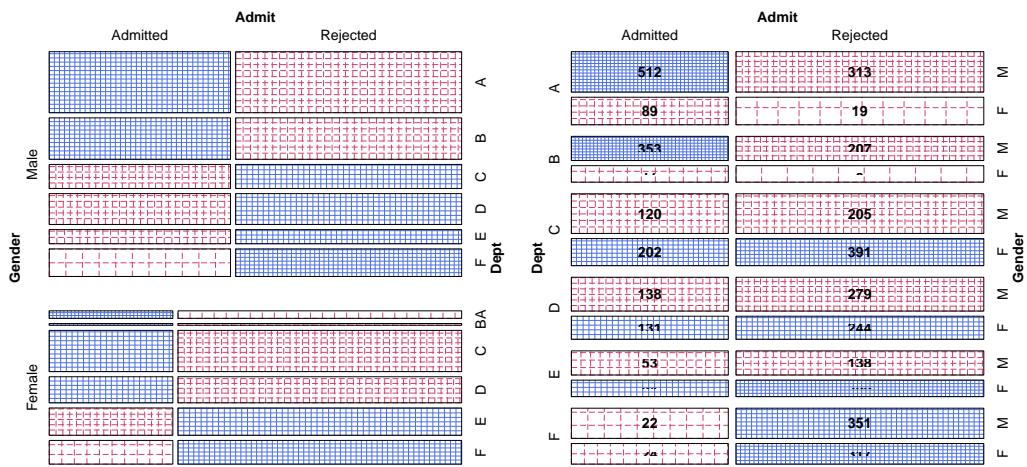


Figure 4.10: Sieve diagrams for the three-way table of the Berkeley admissions data. Left: Admit by Dept, conditioned on Gender; right: Dept re-ordered as the first splitting variable.^{fig:Berkeley-sieve}

In the right panel of Figure 4.10, the three-way table was first permuted to make Dept the first splitting variable. Each 2×2 table of Admit by Gender then appears, giving a sieve diagram version of what we showed earlier in fourfold displays (Figure 4.4). The function `labeling_cells()` is used here to write the cell frequency in each rectangle.

Finally, for tables of more than two dimensions, there is a variety of different models for “independence,” and the `strucplot` framework allows these to be specified with the `expected` argument, either as an array of numbers conforming to the `data` argument, or as a model formula for `loglm()`.

For example, a sieve diagram may be used to determine if the association between gender and department is the same across departments by fitting the model `~Admit*Gender + Dept`, which says that Dept is independent of the combinations of Admit and Gender. This is done as shown below, giving the plot in Figure 4.11.

```
> UCB2 <- aperm(UCBAdmissions, c(3,2,1))
> sieve(UCB2, shade=TRUE, expected=~Admit*Gender + Dept,
+       split_vertical=c(FALSE, TRUE, TRUE))
```

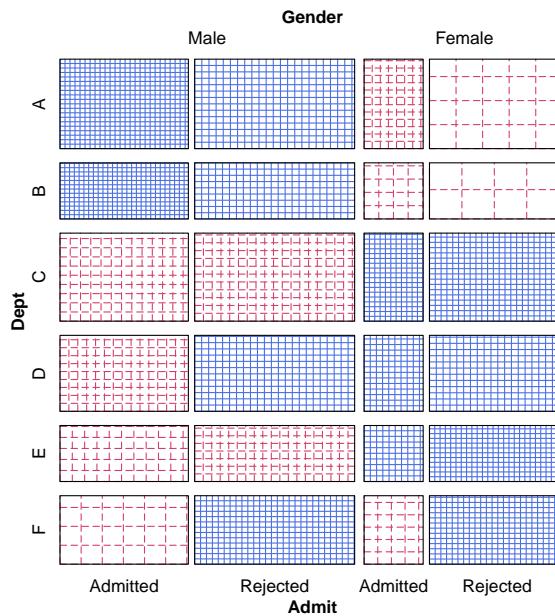


Figure 4.11: Sieve diagram for the Berkeley admissions data, fitting the model of joint independence, `Admit*Gender + Dept`
fig:berkeley-sievez

In terms of the loglinear models discussed in Chapter 5, this is equivalent to fitting the model of **joint independence**, `[AdmitGender][Dept]`. Figure 4.11 shows the greater numbers of male applicants in departments A and B (whose overall rate of admission is high) and greater numbers of female applicants in the remaining departments (where the admission rate is low).



4.6 Association plots

In the sieve diagram the foreground (rectangles) shows expected frequencies; deviations from independence are shown by color and density of shading. The **association plot** (Cohen, 1980, Friendly,

1991) puts deviations from independence in the foreground: the area of each box is made proportional to the (observed – expected) frequency.

For a two-way contingency table, the signed contribution to Pearson χ^2 for cell i, j is

$$d_{ij} = \frac{n_{ij} - m_{ij}}{\sqrt{m_{ij}}} = \text{Pearson residual}, \quad \chi^2 = \sum_{ij} (d_{ij})^2 \quad (4.5) \quad \text{eq:Pearson-residual}$$

In the association plot, each cell is shown by a rectangle, having:

- (signed) height $\sim d_{ij}$
- width $= \sqrt{m_{ij}}$.

so, the area of each cell is proportional to the raw residual, $n_{ij} - m_{ij}$. The rectangles for each row in the table are positioned relative to a baseline representing independence ($d_{ij} = 0$) shown by a dotted line. Cells with observed > expected frequency rise above the line (and are colored blue); cells that contain less than the expected frequency fall below it (and are shaded red).

```
> haireye <- margin.table(HairEyeColor, 1:2)
> assoc(haireye, shade=TRUE)
```

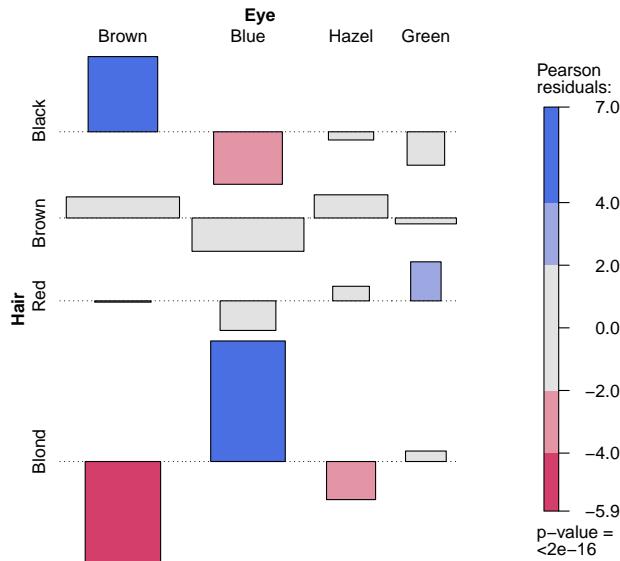


Figure 4.12: Association plot for the two-way table of hair color by eye color.^{fig:HE-assoc}

Figure 4.12 shows the association plot for the data on hair color and eye color. In constructing this plot, each rectangle is shaded according to the value of the Pearson residual Eqn. (4.5), using a simple scale shown in the legend, where residuals $|d_{ij}| > 2$ are shaded blue or red depending on their sign and residuals $|d_{ij}| > 4$ are shaded with a more saturated color.

One virtue of the association plot is that it is quite simple to interpret in terms of the pattern of positive and negative d_{ij} values. Bertin (1981) uses similar graphics to display large complex contingency tables. Like the sieve diagram, however, patterns of association are most apparent when the rows and columns of the display are ordered in a sensible way.

```
> assoc(HairEyeColor, shade=TRUE)
```

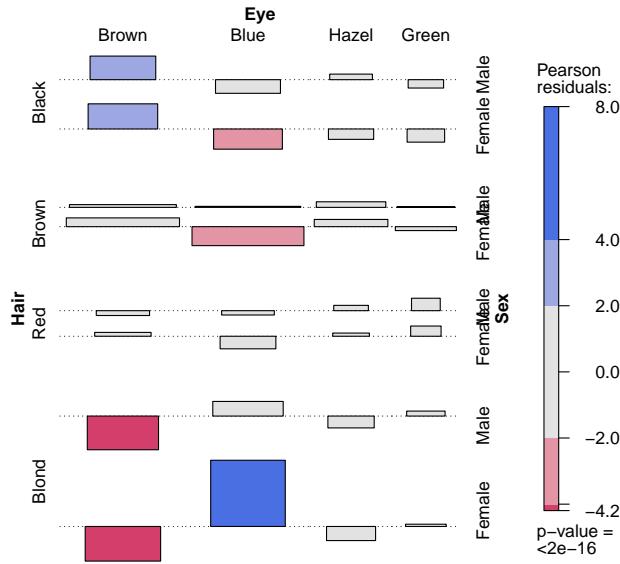


Figure 4.13: Association plot for the three-way table of hair color by eye color by sex.^{fig:HE-assoc2}

We note here that the association plot also belongs to the strucplot framework and thus extends to higher-way tables. For example, the full *HairEyeColor* table is also classified by *Sex*. The plot for the three-way table is shown in Figure 4.13. **TODO:** Perhaps combine these two figures into two panels, side by side. In this plot the third table variable (*Sex* here) is shown nested within the first two, allowing easy comparison of the profiles of hair and eye color for males and females.

4.7 Observer agreement

{sec:twoway-agree}
When the row and column variables represent different observers rating the same subjects or objects, interest is focused on **observer agreement** rather than mere association. In this case, measures and tests of agreement provide a method of assessing the reliability of a subjective classification or assessment procedure.

For example, two (or more) clinical psychologists might classify patients on a scale with categories (a) normal, (b) mildly impaired, (c) severely impaired. Or, ethologists might classify the behavior of animals in categories of cooperation, dominance and so forth, or paleologists might classify pottery fragments according to categories of antiquity or cultural groups. As these examples suggest, the rating categories are often ordered, but not always.

For two raters, a contingency table can be formed classifying all the subjects/objects rated according to the rating categories used by the two observers. In most cases, the same categories are used by both raters, so the contingency table is square, and the entries in the diagonal cells are the cases where the raters agree.

In this section we describe some measures of the strength of agreement and then a method for visualizing the pattern of agreement. But first, the following examples show some typical agreement data.

{ex:sexisfun1}

EXAMPLE 4.15: Sex is fun

The *SexualFun* table in *vcd* (Agresti (1990, Table 2.10), from Hout *et al.* (1987)) summarizes the responses of 91 married couples to a questionnaire item: “Sex is fun for me and my partner: (a) Never or occasionally, (b) fairly often, (c) very often, (d) almost always.”

```
> data("SexualFun", package="vcd")
> SexualFun
```

		Wife						
		Never	Fun	Fairly Often	Very Often	Always	fun	
Husband								
Never	Fun	7		7		2		3
Fairly Often		2		8		3		7
Very Often		1		5		4		9
Always	fun	2		8		9		14

In each row the diagonal entry is not always the largest, though it appears that the partners tend to agree more often when either responds “almost always”. \triangle

{ex:MS1}

EXAMPLE 4.16: Diagnosis of MS patients

Landis and Koch (1977) gave data on the diagnostic classification of multiple sclerosis (MS) patients by two neurologists, one from Winnipeg and one from New Orleans. There were two samples of patients, 149 from Winnipeg and 69 from New Orleans, and each neurologist classified all patients into one of four diagnostic categories: (a) Certain MS, (b) Probable MS, (c) Possible MS, (d) Doubtful, unlikely, or definitely not MS

These data are available in *MSPatients*, a $4 \times 4 \times 2$ table, as shown below. It is convenient to show the data in separate slices for the Winnipeg and New Orleans patients:

```
> MSPatients[, , "Winnipeg"]
```

		Winnipeg Neurologist			
		Certain	Probable	Possible	Doubtful
New Orleans	Neurologist				
	Certain	38	5	0	1
	Probable	33	11	3	0
	Possible	10	14	5	6
	Doubtful	3	7	3	10


```
> MSPatients[, , "New Orleans"]
```

		Winnipeg Neurologist			
		Certain	Probable	Possible	Doubtful
New Orleans	Neurologist				
	Certain	5	3	0	0
	Probable	3	11	4	0
	Possible	2	13	3	4
	Doubtful	1	2	4	14


```
> apply(MSPatients, 3, sum)      # show sample sizes
```

Winnipeg	New Orleans
149	69

In this example, note that the distribution of degree of severity of MS may differ between the two patient samples. As well, for a given sample, the two neurologists may be more or less strict about the boundaries between the rating categories. \triangle

4.7.1 Measuring agreement

In assessing the strength of *agreement* we usually have a more stringent criterion than in measuring the strength of *association*, because observers ratings can be strongly associated without strong

{sec:agreemeas}

agreement. For example, one rater could use a more stringent criterion and thus consistently rate subjects one category lower (on an ordinal scale) than another rater.

More generally, measures of agreement must take account of the marginal frequencies with which two raters use the categories. If observers tend to use the categories with different frequency, this will affect measures of agreement.

Here we describe some simple indices that summarize agreement with a single score (and associated standard errors or confidence intervals). Von Eye and Mun (2006) treat this topic from the perspective of loglinear models.

4.7.1.1 Intraclass correlation

An analysis of variance framework leads to the **intraclass correlation** as a measure of inter-rater reliability, particularly when there are more than two raters. This approach is not covered here, but various applications are described by Shrout and Fleiss (1979), and implemented in R in `ICC()` in the `psych` package.

4.7.1.2 Cohen's Kappa

Cohen's kappa (κ) (Cohen, 1960, 1968) is a commonly used measure of agreement that compares the observed agreement to agreement expected by chance if the two observer's ratings were independent. If p_{ij} is the probability that a randomly selected subject is rated in category i by the first observer and in category j by the other, then the observed agreement is the sum of the diagonal entries, $P_o = \sum_i p_{ii}$. If the ratings were independent, this probability of agreement (by chance) would be $P_c = \sum_i p_{i+} p_{+i}$. Cohen's κ is then the ratio of the difference between actual agreement and chance agreement, $P_o - P_c$, to the maximum value this difference could obtain:

$$\kappa = \frac{P_o - P_c}{1 - P_c} . \quad (\text{4.6})$$

When agreement is perfect, $\kappa = 1$; when agreement is no better than would be obtained from statistically independent ratings, $\kappa = 0$. κ could conceivably be negative, but this rarely occurs in practice. The minimum possible value depends on the marginal totals.

For large samples (n_{++}), κ has an approximate normal distribution when $H_0 : \kappa = 0$ is true and its standard error (Fleiss, 1973, Fleiss *et al.*, 1969) is given by

$$\hat{\sigma}(\kappa) = \frac{P_c + P_c^2 - \sum_i p_{i+} p_{+i} (p_{i+} + p_{+i})}{n_{++}(1 - P_c)^2} .$$

Hence, it is common to conduct a test of $H_0 : \kappa = 0$ by referring $z = \kappa / \hat{\sigma}(\kappa)$ to a unit normal distribution. The hypothesis of agreement no better than chance is rarely of much interest, however. It is preferable to estimate and report a confidence interval for κ .

4.7.1.3 Weighted Kappa

The original (unweighted) κ only counts strict agreement (the same category is assigned by both observers). A weighted version of κ (Cohen, 1968) may be used when one wishes to allow for *partial* agreement. For example, exact agreements might be given full weight, one-category difference given weight 1/2. This typically makes sense only when the categories are *ordered*, as in severity of diagnosis.

Weighted κ uses weights, $0 \leq w_{ij} \leq 1$ for each cell in the table, with $w_{ii} = 1$ for the diagonal cells. In this case P_o and P_c are defined as weighted sums

$$P_o = \sum_i \sum_j w_{ij} p_{ij}$$

$$P_c = \sum_i \sum_j w_{ij} p_{i+} p_{+j}$$

and these weighted sums are used in Eqn. (4.6).

For an $r \times r$ table, two commonly-used pattern of weights are those based on equal spacing of weights (Cicchetti and Allison, 1971) for a near-match, and *Fleiss-Cohen weights* (Fleiss and Cohen, 1972), based on an inverse-square spacing,

$$\begin{aligned} w_{ij} &= 1 - \frac{|i-j|}{r-1} && \text{equal spacing} \\ w_{ij} &= 1 - \frac{|i-j|^2}{(r-1)^2} && \text{Fleiss-Cohen} \end{aligned}$$

The Fleiss-Cohen weights attach greater importance to near disagreements, as you can see below for a 4×4 table. These weights also provide a measure equivalent to the intraclass correlation.

Integer Spacing Cicchetti Allison weights				Inverse Square Spacing Fleiss-Cohen weights			
1	2/3	1/3	0	1	8/9	5/9	0
2/3	1	2/3	1/3	8/9	1	8/9	5/9
1/3	2/3	1	2/3	5/9	8/9	1	8/9
0	1/3	2/3	1	0	5/9	8/9	1

4.7.1.4 Computing Kappa

The function `Kappa()` in `vcd` calculates unweighted and weighted Kappa. The `weights` argument can be used to specify the weighting scheme as either "Equal-Spacing" or "Fleiss-Cohen". The function returns a "Kappa" object, for which there is a `confint.Kappa()` method, providing confidence intervals. The `summary.Kappa()` method also prints the weights.

The lines below illustrate `Kappa` for the `SexualFun` data.

```
> Kappa(SexualFun)

      value     ASE     z
Unweighted 0.129 0.0686 1.89
Weighted   0.237 0.0783 3.03

> confint(Kappa(SexualFun))

Kappa        lwr      upr
Unweighted -0.0051204 0.26378
Weighted    0.0838834 0.39088
```

4.7.2 Observer Agreement Chart

The observer agreement chart proposed by Bangdiwala (1985, 1987) provides a simple graphic representation of the strength of agreement in a contingency table, and alternative measures of strength of agreement with an intuitive interpretation. More importantly, it shows the *pattern* of disagreement when agreement is less than perfect.

The agreement chart is constructed as an $n \times n$ square, where $n = n_{++}$ is the total sample size. Black squares, each of size $n_{ii} \times n_{ii}$, show observed agreement. These are positioned within k larger rectangles, each of size $n_{i+} \times n_{+i}$ as shown in the left panel of Figure 4.14. The large rectangle

{sec:twoway-Bangdiwala}

shows the maximum possible agreement, given the marginal totals. Thus, a visual impression of the strength of agreement is given by

$$B = \frac{\text{area of dark squares}}{\text{area of rectangles}} = \frac{\sum_i^k n_{ii}^2}{\sum_i^k n_{i+} n_{+i}} \quad (4.7) \quad \{\text{eq:bangb}\}$$

When there is perfect agreement, the k rectangles determined by the marginal totals are all squares, completely filled by the shaded squares reflecting the diagonal n_{ii} entries, and $B = 1$.

```
> agreementplot(SexualFun, main="Unweighted", weights=1)
> agreementplot(SexualFun, main="Weighted")
```

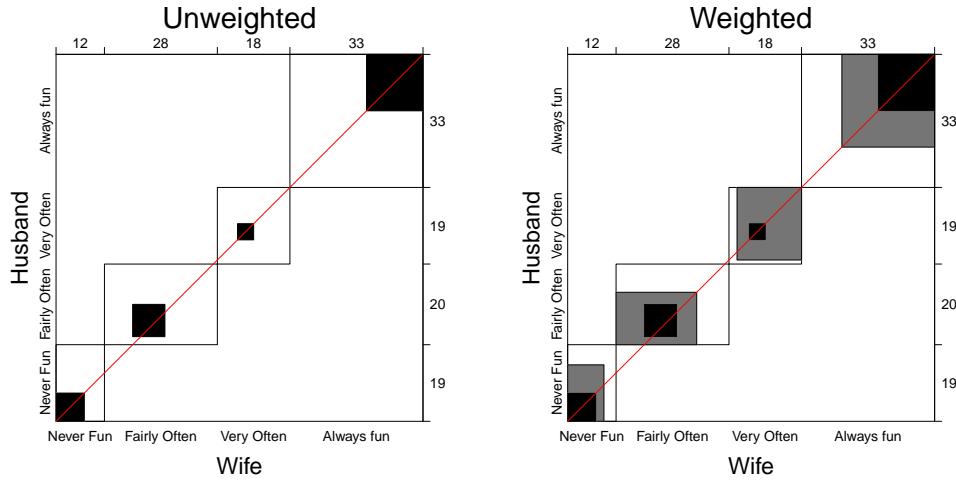


Figure 4.14: Agreement charts for husbands' and wives' sexual fun. Left: unweighted chart, showing only exact agreement; right: weighted chart, using weight $w_1 = 8/9$ for a one-step disagreement.
fig:sexfun-agree

4.7.2.1 Partial agreement

Partial agreement is allowed by including a weighted contribution from off-diagonal cells, b steps from the main diagonal. For a given cell frequency, n_{ij} , a pattern of weights, w_1, w_2, \dots, w_b is applied to the cell frequencies as shown schematically below:

$$\begin{array}{ccccccc} n_{i-b,i} & & & & w_b \\ \vdots & & & & \vdots \\ n_{i,i-b} & \cdots & n_{i,i} & \cdots & n_{i,i+b} & \Leftarrow & w_b \cdots 1 \cdots w_b \\ \vdots & & & & & & \vdots \\ n_{i-b,i} & & & & w_b & & \end{array}$$

These weights are incorporated in the agreement chart (right panel of Figure 4.14) by successively lighter shaded rectangles whose size is proportional to the sum of the cell frequencies, denoted A_{bi} , shown above. A_{1i} allows 1-step disagreements, using weights 1 and w_1 ; A_{2i} includes 2-step disagreements, etc. From this, one can define a weighted measure of agreement, B^w , analogous to

weighted κ :

$$B^w = \frac{\text{weighted sum of areas of agreement}}{\text{area of rectangles}} = 1 - \frac{\sum_i^k [n_{i+} n_{+i} - n_{ii}^2 - \sum_{b=1}^q w_b A_{bi}]}{\sum_i^k n_{i+} n_{+i}}$$

where w_b is the weight for A_{bi} , the shaded area b steps away from the main diagonal, and q is the furthest level of partial disagreement to be considered.

The function `agreementplot()` actually calculates both B and B^w and returns them invisibly as the result of the call. The results, $B = 0.146$, and $B^w = 0.498$, indicate a stronger degree of agreement when 1-step disagreements are included.

```
> B <- agreementplot(SexualFun)
> unlist(B)[1:2]

      Bangdiwala      Bangdiwala_Weighted
        0.14646           0.49817
```

{ex:mammograms}

EXAMPLE 4.17: Mammogram ratings

The *Mammograms* data in *vcdExtra* gives a 4×4 table of (probably contrived) ratings of 110 mammograms by two raters from Kundel and Polansky (2003), used to illustrate the calculation and interpretation of agreement measures in this context.⁸

```
> data("Mammograms", package="vcdExtra")
> B <- agreementplot(Mammograms, main="Mammogram ratings")
```

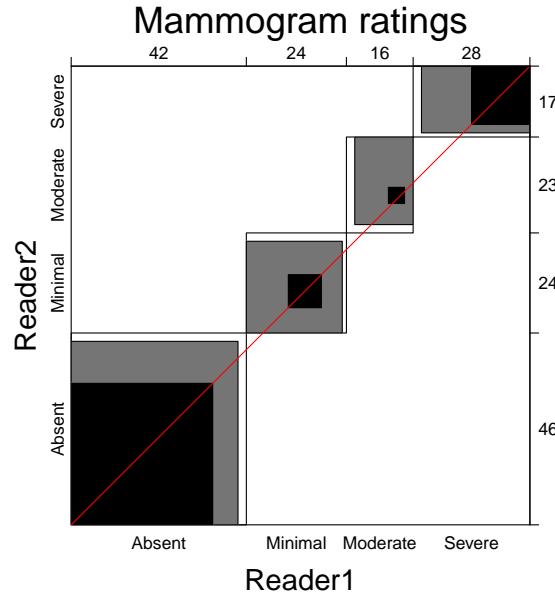


Figure 4.15: Agreement plot for the Mammograms data.^{fig:mammograms1}

The agreement plot in Figure 4.15 shows substantial agreement among the two raters, particularly when one-step disagreements are taken into account. Careful study of this graph shows that

⁸In practice, of course, rater agreement on severity of diagnosis from radiology images varies with many factors. See Antonio and Crespi (2010) for a meta-analytic study concerning agreement in breast cancer diagnosis.

the two raters more often agree exactly for the extreme categories of “Absent” and “Severe.” The amounts of unweighted and weighted agreement are shown numerically in the B and B^w statistics.

```
> unlist(B)[1:2]
```

Bangdiwala	Bangdiwala_Weighted
0.42721	0.83665

△

4.7.3 Observer bias in agreement

With an ordered scale, it may happen that one observer consistently tends to classify the objects into higher or lower categories than the other, perhaps due to using stricter thresholds for the boundaries between adjacent categories. This bias produces differences in the marginal totals, n_{i+} , and n_{+i} and decreases the maximum possible agreement. While special tests exist for **marginal homogeneity**, the observer agreement chart shows this directly by the relation of the dark squares to the diagonal line: When the marginal totals are the same, the squares fall along the diagonal. The measures of agreement, κ and B , cannot determine whether lack of agreement is due to such bias, but the agreement chart can detect this.

EXAMPLE 4.18: Diagnosis of MS patients

Agreement charts for both patient samples in the *MSPatients* data are shown in Figure 4.16. The `agreementplot()` function only handles two-way tables, so we do these separately by indexing on the last dimension (`Patients`).

```
> agreementplot(MSPatients[, "Winnipeg"], main="Winnipeg patients")
> agreementplot(MSPatients[, "New Orleans"], main="New Orleans patients")
```

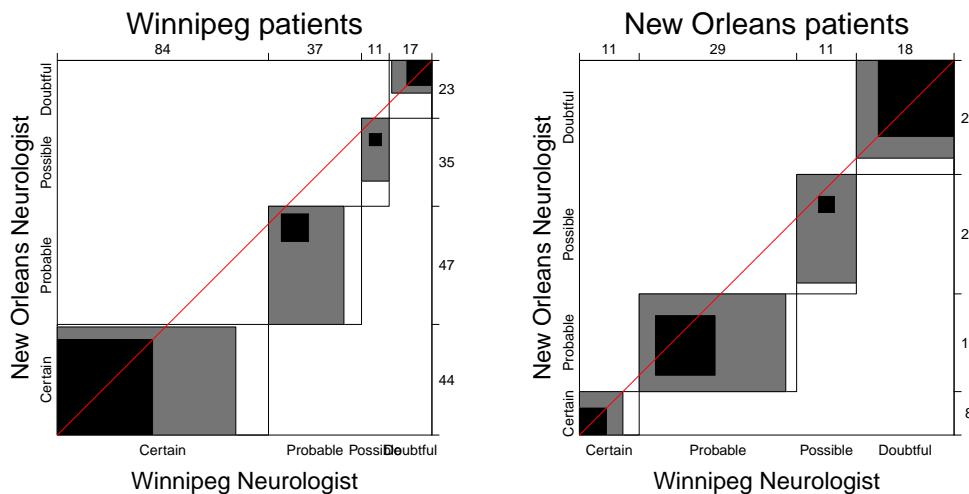


Figure 4.16: Weighted agreement charts for both patient samples in the *MSPatients* data. Departure of the middle rectangles from the diagonal indicates lack of marginal homogeneity.^{fig:MS_Agree}

It can be seen that, for both groups of patients, the rectangles for the two intermediate categories lie largely below the diagonal line (representing equality). This indicates that the Winnipeg neurologist tends to classify patients into more severe diagnostic categories. The departure from the

diagonal is greater for the Winnipeg patients, for whom the Winnipeg neurologist uses the two most severe diagnostic categories very often, as can also be seen from the marginal totals printed in the plot margins.

Nevertheless there is a reasonable amount of agreement if one-step disagreements are allowed, as can be seen in `figrefig:MS-agree` and quantified in the B^w statistics below. The agreement charts also serve to explain why the B measures for exact agreement are so much lower.

```
> agr1 <- agreementplot(MSPatients[, "Winnipeg"])
> agr2 <- agreementplot(MSPatients[, "New Orleans"])
> rbind(Winnipeg=unlist(agr1), NewOrleans=unlist(agr2)) [,1:2]

      Bangdiwala Bangdiwala_Weighted
Winnipeg      0.27210          0.73808
NewOrleans     0.28537          0.82231
```



4.8 Trilinear plots

The *trilinear plot* (also called a *ternary diagram* or *trinomial plot*) is a specialized display for a 3-column contingency table or for three variables whose relative proportions are to be displayed. Individuals may be assigned to one of three diagnostic categories, for example, or a chemical process may yield three constituents in varying proportions, or we may look at the division of votes among three parties in a parliamentary election. This display is useful, therefore, for both frequencies and proportions.

{sec:twoway-trilinear}

Trilinear plots are featured prominently in Aitchison (1986), who describes statistical models for this type of *compositional data*. Upton (1976, 1994) uses them in detailed analyses of spatial and temporal changes in British general elections. Wainer (1996) reviews a variety of other uses of trilinear plots and applies them to aid in understanding the distributions of students achievement in the National Assessment of Educational Progress, making some aesthetic improvements to the traditional form of these plots along the way.

A trilinear plot displays each observation as a point inside an equilateral triangle whose coordinate corresponds to the relative proportions in each column. The three vertices represent the three extremes when 100% occurs in one of the three columns; a point in the exact center corresponds to equal proportions of $\frac{1}{3}$ in all three columns. For instance, Figure ?? shows three points whose compositions of three variables, A, B, and C are given in the data frame DATA below.

```
> library(ggtern)

Error in library(ggtern): there is no package called 'ggtern'

> DATA <- data.frame(
+   A = c(40, 20, 10),
+   B = c(30, 60, 10),
+   C = c(30, 20, 80),
+   id = c("1", "2", "3"))
> ggtern(data = DATA,
+         mapping = aes(x=C, y=A, z=B,
+                           label=id, colour=id)) +
+   geom_point(size=2) +
+   geom_text(vjust=-.5, size=8) +
+   theme_tern_rgbw() +
+   theme(plot.margin=unit(c(0,0,0,0), "mm")) +
+   guides(size = "none")

Error in eval(expr, envir, enclos): could not find function "ggtern"
```

Note that each apex corresponds to 100% of the labeled variable, and the percentage of this variable decrease linearly along a line to the midpoint of the opposite baseline. The grid lines in the figure show the percentage value along each axis.

The construction of trilinear plots is described in detail in http://en.wikipedia.org/wiki/Ternary_plot. Briefly, let $P(a, b, c)$ represent the three components normalized so that $a + b + c = 1.0$. If the apex corresponding to Point A in Figure ?? is given (x, y) coordinates of $(x_A, y_A) = (0, 0)$, and those at apex B are $(x_B, y_B) = (100, 0)$, then the coordinates of apex C are $(x_C, y_C) = (50, 50\sqrt{3})$. The cartesian coordinates (x_P, y_P) of point P are then calculated as

$$\begin{aligned} y_P &= c y_C \\ x_P &= y_P \left(\frac{y_C - y_B}{x_C - x_B} \right) + \frac{\sqrt{3}}{2} y_C (1 - a) \end{aligned}$$

In R, trilinear plots are implemented in the `triplot()` function in the `TeachingDemos` package, and also in the `ggtern` package, an extension of the `ggplot2` framework. The latter is much more flexible, because it inherits all of the capabilities of `ggplot2` for plot annotations, faceting, and layers. In essence, the function `ggtern()` is just a wrapper for `ggplot(...)` which adds a change in the coordinate system from cartesian (x, y) coordinates to the ternary coordinate system with `coord_tern()`.

For example, the following code⁹ creates a data frame `DATA` containing 100 uniformly distributed random points. It uses `stat_density2d()` to draw contours of the densities of the points in the trilinear space.

```
> set.seed(1)
> DATA <- data.frame(x = runif(100),
+                      y = runif(100),
+                      z = runif(100))
> plot <- ggtern(data = DATA,
+                  aes(x, y, z))

Error in eval(expr, envir, enclos): could not find function "ggtern"

> plot + stat_density2d(method = "lm", fullrange = T,
+                        n = 200, geom = "polygon",
+                        aes(fill = ..level..,
+                            alpha = ..level..)) +
+   geom_point() +
+   theme_tern_rgbw() +
+   labs(title = "Uniform data with density contours") +
+   scale_fill_gradient(low = "blue", high = "red") +
+   guides(color = "none", fill = "none", alpha = "none")

Error in plot + stat_density2d(method = "lm", fullrange = T, n = 200, : non-numeric
argument to binary operator
```

{ex:lifeboat1}

EXAMPLE 4.19: Lifeboats on the *Titanic*

We examine the question of who survived and why in the sinking of the *RMS Titanic* in Section 5.4 (Example ??), where we analyze a four-way table, *Titanic*, of the 2201 people on board (1316 passengers and 885 crew), classified by Class, Sex, Age, and Survival. A related data set, *Lifeboats* in `vcd` tabulates the survivors according to the life boats on which they were loaded. This data sheds some additional light on the issue of survival and provides a nice illustration of trilinear plots.

⁹This example was taken from the `ggtern` web site, <http://ggtern.com/2013/12/12/patched-density-functions-2/>.

A bit of background: after the disaster, the British Board of Trade launched several inquiries, the most comprehensive of which resulted in the *Report on the Loss of the “Titanic” (S.S.) by Lord Mersey* (Mersey, 1912).¹⁰ The data frame *Lifeboats* in *vcd* contains the data listed on p. 38 of that report.¹¹

Of interest here is the composition of the boats by the three categories, men, women and children and crew, and according to the launching of the boats from the port or starboard side. This can be shown in a trilinear display using the following statements. The plot, shown in Figure ??, has most of the points near the top, corresponding to a high percentage of women and children. We create a variable, *id*, used to label those boats with more than 10% male passengers. In the *ggplot2* framework, plot aesthetics, such as color and shape can be mapped to variables in the data set, and here we map these both to *side* of the boat.

```
> data("Lifeboats", package="vcd")
> # label boats with more than 10% men
> Lifeboats$id <- ifelse(Lifeboats$men/Lifeboats$total > .1,
+                         as.character(Lifeboats$boat), "")
> ggtern(data = Lifeboats,
+         mapping = aes(x = women, y = men, z = crew,
+                        colour=side, shape=side, label=id)) +
+   theme_tern_rbw() +
+   theme(plot.margin=unit(c(0,0,0,0), "mm")) +
+   geom_point(aes(size=2)) +
+   labs(title = "Lifeboats on the Titanic") +
+   labs(T="Women and children") +
+   guides(size = "none") +
+   geom_smooth(method="lm", size=1.5, aes(fill=side)) +
+   geom_text(vjust=1, color="black")
```

Error in eval(expr, envir, enclos): could not find function "ggtern"

The resulting plot in Figure ??, makes it immediately apparent that many of the boats launched from the port side differ substantially from the remaining boats, whose passengers were almost entirely women and children. Boat 1 had only 20% (2 out of 10) women and children, while the percentage for boat 3 was only 50% (25 out of 50). We highlight the difference in composition of the boats launched from the two sides by adding a linear regression smooth for the relation *men* ~ *women*.

The trilinear plot scales the numbers for each observation to sum to 1.0, so differences in the total number of people on each boat cannot be seen in Figure ???. The total number reported loaded is plotted against launch time in Figure 4.17, with a separate regression line and loess smooth fit to the data for the port and starboard sides.

```
> ggplot(data = Lifeboats,
+         aes(x=launch, y=total, colour=side, label=boat)) +
+   geom_smooth(method="lm", aes(fill=side), size=1.5) +
+   geom_smooth(method="loess", aes(fill=side), se=FALSE, size=1.2) +
+   geom_point() + ylim(c(0,100)) +
+   geom_text(vjust=-.5, color="black") +
+   labs(y="Total loaded", x="Launch time")
```

¹⁰The *Titanic* was outfitted with 20 boats, half on each of the port and starboard sides, of which 14 were large lifeboats with a capacity of 65, two were emergency boats designed for 40 persons, and the remaining four were collapsible boats capable of holding 47, a total capacity of 1178 (considered adequate at that time). Two of the collapsible boats, lashed to the roof of the officers quarters, were ineffectively launched and utilized as rafts after the ship sunk. The report lists the time of launch and composition of the remaining 18 boats according to male passengers, women and children, and “men of crew”, as reported by witnesses.

¹¹The “data” lists a total of 854 in 18 boats, although only 712 were in fact saved. Mersey notes “it is obvious that these figures are quite unreliable”.

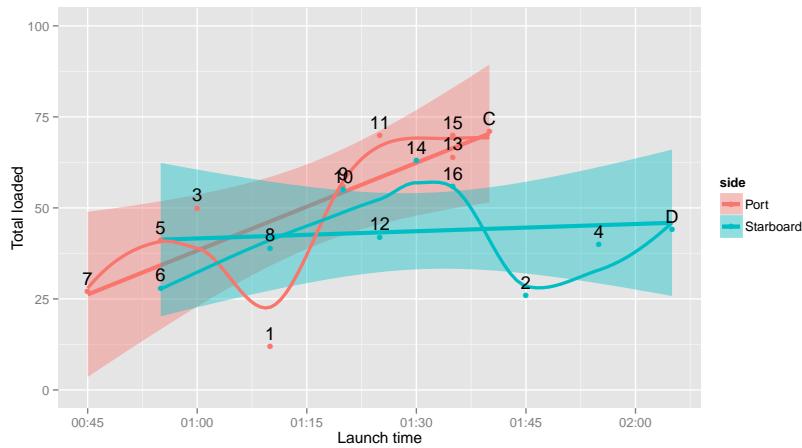


Figure 4.17: Number of people loaded on lifeboats on the Titanic vs. time of launch, by side of boat. The plot annotations show the linear regression and loess smooth.
fig:lifeboats2

From the linear regression lines in Figure 4.17, it seems that the rescue effort began in panic on the port side, with relatively small numbers loaded, and (from Figure ??), small proportions of women and children. But the loading regime on that side improved steadily over time. The procedures began more efficiently on the starboard side but the numbers loaded increased only slightly. The smoothed loess curves indicate that over time, for each side, there was still a large variability from boat to boat.



4.9 Chapter summary

- A contingency table gives the frequencies of observations cross-classified by two or more categorical variables. With such data we are typically interested in testing whether associations exist, quantifying the strength of association, and understanding the nature of the association among these variables.
- For 2×2 tables, association is easily summarized in terms of the odds ratio or its logarithm. This measure can be extended to stratified $2 \times 2 \times k$ tables, where we can also assess whether the odds ratios are equal across strata or how they vary.
- For $r \times c$ tables, measures and tests of general association between two categorical variables are most typically carried out using the Pearson's chi-square or likelihood-ratio tests provided by `assocstats()`. Stratified tests controlling for one or more background variables, and tests for ordinal categories are provided by the Cochran-Mantel-Haenszel tests given by `CMHtest()`.
- For 2×2 tables, the fourfold display provides a visualization of the association between variables in terms of the odds ratio. Confidence rings provide a visual test of whether the odds ratio differs significantly from 1. Stratified plots for $2 \times 2 \times k$ tables are also provided by `fourfold()`.
- Sieve diagrams and association plots provide other useful displays of the pattern of association in $r \times c$ tables. These also extend to higher-way tables as part of the `strucplot` framework.

- When the row and column variables represent different observers rating the same subjects, interest is focused on agreement rather than mere association. Cohen's κ is one measure of strength of agreement. The observer agreement chart provides a visual display of how the observers agree and disagree.
- Another specialized display, the trilinear plot is useful for three-column frequency tables or compositional data.

4.10 Further reading

4.11 Lab exercises

Exercise 4.1 The data set `fat`, created below, gives a 2×2 table recording the level of cholesterol in diet and the presence of symptoms of heart disease for a sample of 23 people.

```
> fat <- matrix( c(6, 4, 2, 11), 2, 2)
> dimnames(fat) <- list(diet=c("LoChol", "HiChol"),
+                           disease=c("No", "Yes"))
```

- Use `chisq.test(fat)` to test for association between diet and disease. Is there any indication that this test may not be appropriate here?
- Use a fourfold display to test this association visually. Experiment with the different options for standardizing the margins, using the `margin` argument to `fourfold()`. What evidence is shown in different displays regarding whether the odds ratio differs significantly from 1?
- `oddsratio(fat, log=FALSE)` will give you a numerical answer. How does this compare to your visual impression from fourfold displays?
- With such a small sample, Fisher's exact test may be more reliable for statistical inference. Use `fisher.test(fat)`, and compare these results to what you have observed before.
- Write a one-paragraph summary of your findings and conclusions for this data set.

Exercise 4.2 The data set `Abortion` in `vcdExtra` gives a $2 \times 2 \times 2$ table of opinions regarding abortion in relation to sex and status of the respondent. This table has the following structure:

```
> data("Abortion", package="vcdExtra")
> str(Abortion)

table [1:2, 1:2, 1:2] 171 152 138 167 79 148 112 133
- attr(*, "dimnames")=List of 3
..$ Sex : chr [1:2] "Female" "Male"
..$ Status : chr [1:2] "Lo" "Hi"
..$ Support_Abortion: chr [1:2] "Yes" "No"
```

- Taking support for abortion as the outcome variable, produce fourfold displays showing the association with sex, stratified by status.
- Do the same for the association of support for abortion with status, stratified by sex.
- For each of the problems above, use `oddsratio()` to calculate the numerical values of the odds ratio, as stratified in the question.

Exercise 4.3 The `JobSat` table on income and job satisfaction created in Example 2.5 is contained in the `vcdExtra` package.

- (a) Carry out a standard χ^2 test for association between income and job satisfaction. Is there any indication that this test might not be appropriate? Repeat this test using `simulate.p.value = TRUE` to obtain a Monte Carlo test that does not depend on large sample size. Does this change your conclusion?
- (b) Both variables are ordinal, so CMH tests may be more powerful here. Carry out that analysis. What do you conclude?

{lab:4.4}

Exercise 4.4 The `Hospital` data in `vcd` gives a 3×3 table relating the length of stay (in years) of 132 long-term schizophrenic patients in two London mental hospitals with the frequency of visits by family and friends.

- (a) Carry out a χ^2 test for association between the two variables.
- (b) Use `assocstats()` to compute association statistics. How would you describe the strength of association here?
- (c) Produce an association plot for these data, with visit frequency as the vertical variable. Describe the pattern of the relation you see here.
- (d) Both variables can be considered ordinal, so `CMHtest()` may be useful here. Carry out that analysis. Do any of the tests lead to different conclusions?

{lab:4.5}

Exercise 4.5 The two-way table `Mammograms` in `vcdExtra` gives ratings on the severity of diagnosis of 110 mammograms by two raters.

- (a) Assess the strength of agreement between the raters using Cohen's κ , both unweighted and weighted.
- (b) Use `agreementplot()` for a graphical display of agreement here.

{lab:4.6}

Exercise 4.6 Agresti and Winner (1997) gave the data in Table 4.8 on the ratings of 160 movies by the reviewers Gene Siskel and Roger Ebert for the period from April 1995 through September 1996. The rating categories were Con ("thumbs down"), Mixed and Pro ("thumbs up").

Table 4.8: Movie ratings by Siskel & Ebert, April 1995–September 1996. *Source:* Agresti and Winner (1997)

		Ebert			total
		Con	Mixed	Pro	
Siskel	Con	24	8	13	45
	Mixed	8	13	11	32
	Pro	10	9	64	83
		total	42	30	88
					160

- (a) Assess the strength of agreement between the raters using Cohen's κ , both unweighted and weighted.
- (b) Use `agreementplot()` for a graphical display of agreement here.
- (c) Assess the hypothesis that the ratings are *symmetric* around the main diagonal. Hint: Symmetry for a square table T means that $t_{ij} = t_{ji}$ for $i \neq j$. The expected frequencies under the hypothesis of symmetry are the average of the off-diagonal cells, $E = (T + T^T)/2$.

{lab:4.7}

Exercise 4.7 For the `VisualAcuity` data set:

- (a) Use the code shown in the text to create the table form, `VA.tab`.
- (b) Perform the CMH tests for this table.
- (c) Use `loglm()` method described in Section 4.3.1 to test whether the association between left and right eye acuity can be considered the same for men and women.

{lab:4.8}

Exercise 4.8 The graph in Figure 4.17 may be misleading, in that it doesn't take account of the differing capacities of the 18 life boats on the *Titanic*, given in the variable `cap` in the *Lifeboats* data.

- (a) Calculate a new variable, `pctloaded` as the percentage loaded relative to the boat capacity.
- (b) Produce a plot similar to Figure 4.17, showing the changes over time in this measure.

```
> detach(package:ggtern)
Error in detach(package:ggtern): invalid 'name' argument
> .locals$ch04 <- setdiff(ls(), .globals)
> remove(list=.locals$ch04[sapply(.locals$ch04,function(n){!is.function(get(n))})])
```


Chapter 5

Mosaic displays for n-way tables

Mosaic displays help to visualize the pattern of associations among variables in two-way and larger tables. Extensions of this technique can reveal partial associations, marginal associations, and shed light on the structure of loglinear models themselves.

{ch:mosaic}

5.1 Introduction

{sec:mosaic-intro}

Little boxes, little boxes, little boxes made of ticky-tacky;
Little boxes, little boxes, little boxes all the same.
There are red ones, and blue ones, and green ones, and yellow ones;
Little boxes, little boxes, and they all look just the same.

Pete Seeger

In Chapter 4, we described a variety of graphical techniques for visualizing the pattern of association in simple contingency tables. These methods are somewhat specialized for particular sizes and shapes of tables: 2×2 tables (fourfold display), $r \times c$ tables (sieve diagram), square tables (agreement charts), $r \times 3$ tables (trilinear plots), and so forth.

This chapter describes the *mosaic display* and related graphical methods for n -way frequency tables, designed to show various aspects of high-dimensional contingency tables in a hierarchical way. These methods portray the frequencies in an n -way contingency table by a collection of rectangular “tiles” whose size (area) is proportional to the cell frequency. In this respect, the mosaic display is similar to the sieve diagram (Section 4.5). However, mosaic plots and related methods described here:

- generalize more readily to n -way tables. One can usefully examine 3-way, 4-way and even larger tables, subject to the limitations of resolution in any graph;
- are intimately connected to loglinear models, generalized linear models and generalized nonlinear models for frequency data.
- provide a method for fitting a series of sequential loglinear models to the various marginal totals of an n -way table; and
- can be used to illustrate the relations among variables which are fitted by various loglinear models.

{sec:mosaic-twoway}

5.2 Two-way tables

The mosaic display (Friendly, 1992, 1994b, 1997, Hartigan and Kleiner, 1981, 1984) is like a grouped barchart, where the heights (or widths) of the bars show the relative frequencies of one variable, and widths (heights) of the sections in each bar show the conditional frequencies of the second variable, given the first. This gives an area-proportional visualization of the frequencies composed of tiles corresponding to the cells created by successive vertical and horizontal splits of rectangle, representing the total frequency in the table. The construction of the mosaic display, and what it reveals, are most easily understood for two-way tables.

{ex:haireye2a}

EXAMPLE 5.1: Hair color and eye color

Consider the data shown earlier in Table 4.2, showing the relation between hair color and eye color among students in a statistics course. The basic mosaic display for this 4×4 table is shown in Figure 5.1.

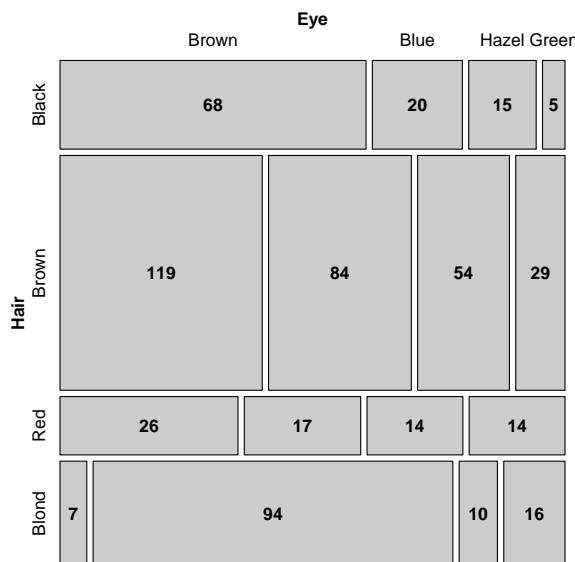


Figure 5.1: Basic mosaic display for hair color and eye color data. The area of each rectangle is proportional to the observed frequency in that cell, shown as numbers.
fig:haireye-most

```
> data(HairEyeColor, package="datasets")
> haireye <- margin.table(HairEyeColor, 1:2)
> mosaic(haireye)
```

For such a two-way table, the mosaic in Figure 5.1 is constructed by first dividing a unit square in proportion to the marginal totals of one variable, say, Hair color.

For these data, the marginal frequencies and proportions of Hair color are calculated below:

```
> (hair <- margin.table(haireye, 1))
```

Hair	Black	Brown	Red	Blond
	108	286	71	127

```
> prop.table(hair)

Hair
Black   Brown    Red   Blond
0.18243 0.48311 0.11993 0.21453
```



Figure 5.2: First step in constructing a mosaic display. Left: splitting the unit square according to frequencies of hair color; right: shading the tiles according to residuals from a model of equal marginal probabilities.
fig:haireye-mosa

These frequencies can be shown as the mosaic for the first variable (hair color), with the unit square split according to the marginal proportions as in Figure 5.2 (left). The rectangular tiles are then shaded to show the residuals (deviations) from a particular model as shown in the right panel of Figure 5.2. The details of the calculations for shading are:

- The one-way table of marginal totals can be fit to a model, in this case, the (implausible) model that all hair colors are equally probable. This model has expected frequencies $m_i = 592/4 = 148$:

```
> expected <- rep(sum(hair)/4, 4)
> names(expected) <- names(hair)
> expected

Black  Brown  Red  Blond
148    148    148    148
```

- The Pearson residuals from this model, $r_i = (n_i - m_i)/\sqrt{m_i}$, are:

```
> (residuals <- (hair - expected) / sqrt(expected))

Hair
Black   Brown    Red   Blond
-3.2880 11.3435 -6.3294 -1.7262
```

and these values are shown by color and shading as shown in the legend. The high positive value for Brown hair indicates that people with brown hair are much more frequent in this sample than the equiprobability model would predict; the large negative residual for Red hair shows that red heads are much less common. Further details of the schemes for shading are described below, but essentially we use increasing intensities of blue (red) for positive (negative) residuals.

In the next step, the rectangle for each Hair color is subdivided in proportion to the *relative* (conditional) frequencies of the second variable— Eye color, giving the following conditional row proportions:

```
> round(addmargins(prop.table(haireye, 1), 2), 3)
```

	Eye				
Hair	Brown	Blue	Hazel	Green	Sum
Black	0.630	0.185	0.139	0.046	1.000
Brown	0.416	0.294	0.189	0.101	1.000
Red	0.366	0.239	0.197	0.197	1.000
Blond	0.055	0.740	0.079	0.126	1.000

The proportions in each row determine the heights of the tiles in the second mosaic display in Figure 5.3.

```
> mosaic(haireye, shade=TRUE, suppress=0,
+         labeling=labeling_residuals, gp_text=gpar(fontface=2))
```

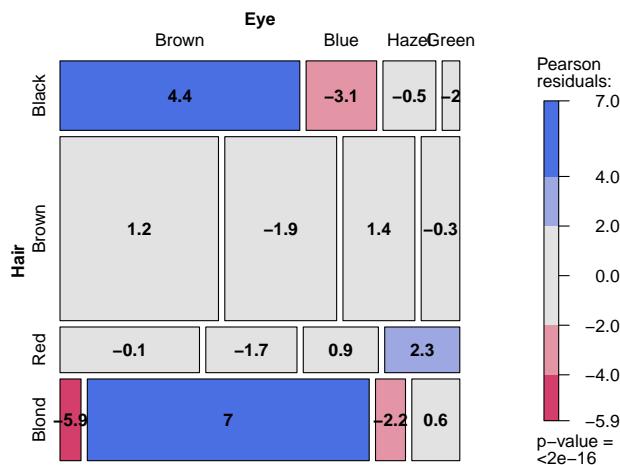


Figure 5.3: Second step in constructing the mosaic display. Each rectangle for hair color is subdivided in proportion to the relative frequencies of eye color, and the tiles are shaded in relation to residuals from the model of independence.

- Again, the cells are shaded in relation to standardized Pearson residuals, $r_{ij} = (n_{ij} - m_{ij}) / \sqrt{m_{ij}}$, from a model. For a two-way table, the model is that Hair color and Eye color are independent in the population from which this sample was drawn. These residuals are calculated as shown below using `loglm()` to fit the independence model and `residuals()`.

```
> HE.mod <- loglm(~ Hair + Eye, data=haireye)
> round(resids <- residuals(HE.mod, type="pearson"), 2)

Re-fitting to get frequencies and fitted values
```

	Eye				
Hair	Brown	Blue	Hazel	Green	
Black	4.40	-3.07	-0.48	-1.95	
Brown	1.23	-1.95	1.35	-0.35	
Red	-0.07	-1.73	0.85	2.28	
Blond	-5.85	7.05	-2.23	0.61	

- Thus, in Figure 5.3, the two tiles shaded deep blue correspond to the two cells, (Black, Brown) and (Blond, Blue), whose residuals are greater than +4, indicating much greater frequency in those cells than would be found if Hair color and Eye color were independent. The tile shaded deep red, (Blond, Brown), corresponds to the largest negative residual = -5.85, indicating this combination is extremely rare under the hypothesis of independence.
- The overall Pearson χ^2 statistic for the independence model is just the sum of squares of the residuals, with degrees of freedom $(r - 1) \times (c - 1)$.

```
> (chisq <- sum(resids^2))

[1] 138.29

> (df <- prod(dim(haireye)-1) )

[1] 9

> chisq.test(haireye)

Pearson's Chi-squared test

data: haireye
X-squared = 138.29, df = 9, p-value < 2.2e-16
```



5.2.0.1 Shading levels

A variety of schemes for shading the tiles are available in the `strucplot` framework (Section 5.3), but the simplest (and default) shading patterns for the tiles are based on the sign and magnitude of the standardized Pearson residuals, using shades of blue for positive residuals and red for negative residuals, and two threshold values for their magnitudes, $|r_{ij}| > 2$ and $|r_{ij}| > 4$.

Because the standardized residuals are approximately unit-normal $N(0, 1)$ values, this corresponds to highlighting cells whose residuals are *individually* significant at approximately the .05 and .0001 level, respectively. Other shading schemes described later provide tests of significance, but the main purpose of highlighting cells is to draw attention to the *pattern* of departures of the data from the assumed model of independence.

5.2.0.2 Interpretation and reordering

To interpret the association between Hair color and Eye color, consider the pattern of positive (blue) and negative (red) tiles in the mosaic display. We interpret positive values as showing cells whose observed frequency is substantially greater than would be found under independence; negative values indicate cells which occur less often than under independence.

The interpretation can often be enhanced by reordering the rows or columns of the two-way table so that the residuals have an *opposite corner* pattern of signs. This usually helps us interpret any systematic patterns of association in terms of the ordering of the row and column categories.

In this example, a more direct interpretation can be achieved by reordering the Eye colors as shown in Figure 5.4. Note that in this rearrangement both hair colors and eye colors are ordered from dark to light, suggesting an overall interpretation of the association between Hair color and Eye color.

```
> # re-order Eye colors from dark to light
> haireye2 <- haireye[, c("Brown", "Hazel", "Green", "Blue")]
> mosaic(haireye2, shade=TRUE)
```

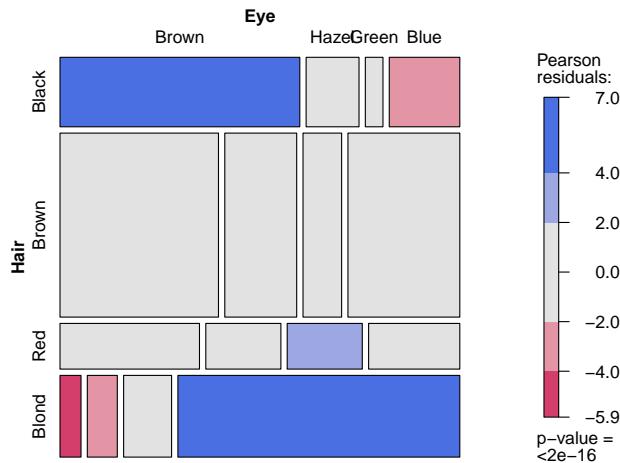


Figure 5.4: Two-way mosaic for Hair color and Eye color reordered. The Eye colors were reordered from dark to light, enhancing the interpretation.
fig:haireye-mos9

In general, the levels of a factor in mosaic displays are often best reordered by arranging them according to their scores on the first (largest) *correspondence analysis* dimension (Friendly, 1994b); see Chapter 6 for details. Friendly and Kwan (2003) use this as one example of *effect ordering* for data displays, illustrated in Chapter 1.

Thus, the mosaic in Figure 5.4 shows that the association between Hair and Eye color is essentially that:

- people with dark hair tend to have dark eyes,
- those with light hair tend to have light eyes
- people with red hair and hazel eyes do not quite fit this pattern

5.3 The strucplot framework

Mosaic displays have much in common with sieve plots and association plots described in Chapter 4 and with related graphical methods such as *doubledecker plots* described later in this chapter. The main idea is to visualize a contingency table of frequencies by “tiles” corresponding to the table cells arranged in rectangular form. For multiway tables with more than two factors, the variables are

c:mosaic-strucplot}

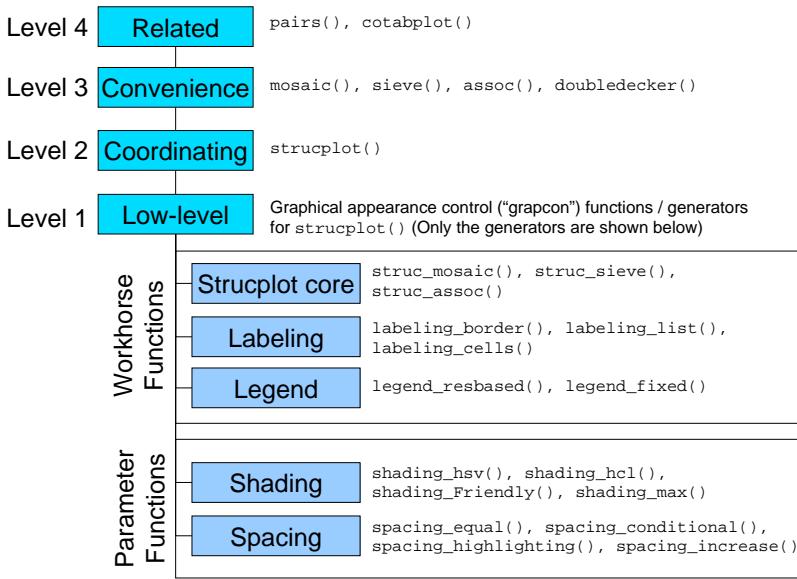


Figure 5.5: Components of the strucplot framework. High level functions use those at lower levels to provide a general system for tile-based plots of frequency tables.

{fig:struc}

nested into rows and columns using recursive conditional splits, given the table margins. The result is a “flat” representation that can be visualized in ways similar to a two-dimensional representation of a table. The `structable()` function described in Section 2.5 gives the tabular version of a strucplot. The description below follows Meyer *et al.* (2006), also included as a vignette, (accessible from R as `vignette("strucplot", pkg="vcd")`), in `vcd`.

Rather than implementing each of these methods separately, the **strucplot framework** in the `vcd` package provides a general class of methods of which these are all instances. This framework defines a class of conditional displays which allows for granular control of graphical appearance aspects, including:

- the content of the tiles, e.g., observed or expected frequencies
- the split direction for each dimension, horizontal or vertical
- the graphical parameters of the tiles’ content, e.g., color or other visual attributes
- the spacing between the tiles
- the labeling of the tiles

The strucplot framework is highly modularized: Figure 5.5 shows the hierarchical relationship between the various components. For the most part, you will use directly the convenience and related functions at the top of the diagram, but it is more convenient to describe the framework from the bottom up.

1. On the lowest level, there are several groups of workhorse and parameter functions that directly or indirectly influence the final appearance of the plot (see Table 5.1 for an overview). These are examples of *graphical appearance control* functions (called **grapcon functions**). They are created by generating functions (*grapcon generators*), allowing flexible parameterization and extensibility (Figure 5.5 only shows the generators). The generator names follow the naming convention `group_foo()`, where `group` reflects the group the generators belong to (strucplot core, labeling, legend, shading, or spacing).

Group	Grapcon generator	Description
strucplot core	struc_assoc() struc_mosaic() struc_sieve()	core function for association plots core function for mosaic plots core function for sieve plots
labeling	labeling_border() labeling_cboxed() labeling_cells() labeling_conditional() labeling_doubledecker() labeling_lboxed() labeling_left() labeling_left2() labeling_list() labeling_residuals() labeling_value()	border labels centered labels with boxes, all labels clipped, and on top and left border cell labels border labels for conditioning variables and cell labels for conditioned variables draws labels for doubledecker plot left-aligned labels with boxes left-aligned border labels left-aligned border labels, all labels on top and left border draws a list of labels under the plot show residuals in cells show values (observed, expected) in cells
shading	shading_binary() shading_Friendly() shading_hcl() shading_hsv() shading_max() shading_sieve()	visualizes the sign of the residuals implements Friendly shading (based on HSV colors) shading based on HCL colors shading based on HSV colors shading visualizing the maximum test statistic (based on HCL colors) implements Friendly shading customized for sieve plots (based on HCL colors)
spacing	spacing_conditional() spacing_dimequal() spacing_equal() spacing_highlighting() spacing_increase()	increasing spacing for conditioning variables, equal spacing for conditioned variables equal spacing for each dimension equal spacing for all dimensions increasing spacing, last dimension set to zero increasing spacing
legend	legend_fixed() legend_resbased()	creates a fixed number of bins (similar to <code>mosaicplot()</code>) suitable for an arbitrary number of bins (also for continuous shadings)

Table 5.1: Available graphical appearance control (grapcon) generators in the strucplot framework

{tab:grapcons}

- The workhorse functions (created by `struc_foo()`) are `labeling_foo()`, and `legend_foo()`. These functions directly produce graphical output (i.e., “add ink to the canvas”), for labels and legends respectively.
 - The parameter functions (created by `spacing_foo()` and `shading_foo()`) compute graphical parameters used by the others. The `grapcon` functions returned by `struc_foo()` implement the core functionality, creating the tiles and their content.
2. On the second level of the framework, a suitable combination of the low-level `grapcon` functions (or, alternatively, corresponding generating functions) is passed as “hyperparameters” to `strucplot()`. This central function sets up the graphical layout using grid viewports, and coordinates the specified core, labeling, shading, and spacing functions to produce the plot.
 3. On the third level, `vcd` provides several convenience functions such as `mosaic()`, `sieve()`, `assoc()`, and `doublededecker()` which interface to `strucplot()` through sensible parameter defaults and support for model formulae.
 4. Finally, on the fourth level, there are “related” `vcd` functions (such as `cotabplot()` and the `pairs()` methods for table objects) arranging collections of plots of the strucplot framework into more complex displays (e.g., by means of panel functions).

5.3.1 Shading schemes

Unlike other graphics functions in base R, the strucplot framework allows almost full control over the graphical parameters of all plot elements. In particular, in association plots, mosaic plots, and sieve plots, you can modify the graphical appearance of each tile individually.

{sec:mosaic-shading}

Built on top of this functionality, the framework supplies a set of shading functions choosing colors appropriate for the visualization of loglinear models. The tiles’ graphical parameters are set using the `gp` argument of the functions of the strucplot framework. This argument basically expects an object of class “`gpar`” whose components are arrays of the same shape (length and dimensionality) as the data table.

For added generality, however, you can also supply a `grapcon` function that computes such an object given a vector of residuals, or, alternatively, a *generating function* that takes certain arguments and returns such a `grapcon` function (see Table 5.1). `vcd` provides several shading functions, including support for both HSV and HCL colors, and the visualization of significance tests. **TODO:** This points to the need for a section, probably in Chapter 1, on color spaces and color schemes for categorical data graphics.

5.3.1.1 Specifying graphical parameters for strucplot displays

Strucplot displays in `vcd` are built using the `grid` graphics package. There are many graphical parameters that can be set using `gp = gpar(...)` in a call to a high-level strucplot function. Among these, the following are often most useful to control the drawing components:

<code>col</code>	Color for lines and borders.
<code>fill</code>	Color for filling rectangles, polygons, ...
<code>alpha</code>	Alpha channel for transparency of fill color.
<code>lty</code>	Line type for lines and borders.
<code>lwd</code>	Line width for lines and borders.

In addition, a number of parameters control the display of text labels in these displays:

<code>fontsize</code>	The size of text (in points)
<code>cex</code>	Multiplier applied to <code>fontsize</code>

`fontfamily` The font family
`fontface` The font face (**bold**, *italic*, ...)

See `help(gpar)` for a complete list and further details.

We illustrate this capability below using the Hair color and Eye color data as reordered in Figure 5.4. The following example produces a **Marimekko chart**, or a “poor-man’s mosaic display” as shown in the left panel of Figure 5.6. This is essentially a divided bar chart where the eye colors within each horizontal bar for the hair color group are all given the same color. In the example, the matrix `fill_colors` is constructed to conform to the `haireye2` table, using color values that approximate the eye colors.

```
> # color by hair color
> fill_colors <- c("brown4", "#acba72", "green", "lightblue")
> (fill_colors <- t(matrix(rep(fill_colors, 4), ncol=4)))

[,1]      [,2]      [,3]      [,4]
[1,] "brown4" "#acba72" "green" "lightblue"
[2,] "brown4" "#acba72" "green" "lightblue"
[3,] "brown4" "#acba72" "green" "lightblue"
[4,] "brown4" "#acba72" "green" "lightblue"

> mosaic(haireye2, gp=gpar(fill=fill_colors, col=0))
```

Note that because the hair colors and eye colors are both ordered, this shows the decreasing prevalence of light hair color amongst those with brown eyes and the increasing prevalence of light hair with blue eyes.

Alternatively, for some purposes,¹ we might like to use color to highlight the pattern of diagonal cells, and the off-diagonals 1, 2, 3 steps removed. The R function `toeplitz()` returns such a patterned matrix, and we can use this to calculate the `fill_colors` by indexing the `palette()` function. The code below produces the right panel in Figure 5.6.

```
> # toeplitz designs
> toeplitz(1:4)

[,1]  [,2]  [,3]  [,4]
[1,]    1     2     3     4
[2,]    2     1     2     3
[3,]    3     2     1     2
[4,]    4     3     2     1

> fill_colors <- palette()[1+toeplitz(1:4)]
> mosaic(haireye2, gp=gpar(fill=fill_colors, col=0))
```

More simply, to shade a mosaic according to the levels of one variable (typically a response variable), you can use the `highlighting` arguments of `mosaic()`. The first call below gives a result similar to the left panel of Figure 5.6. Alternatively, using the formula method for `mosaic()`, specify the response variable as the left-hand side.

```
> mosaic(haireye2, highlighting="Eye", highlighting_fill=fill_colors)
> mosaic(Eye ~ Hair, data=as.table(haireye2))
```

5.3.1.2 Residual-based shading

The important idea that differentiates mosaic and other strucplot displays from the “poor-man’s,” Marimekko versions (Figure 5.6) often shown in other software is that rather than just using shading

¹For example, this would be appropriate for a square table, showing agreement between row and column categories, as in Section 4.7.

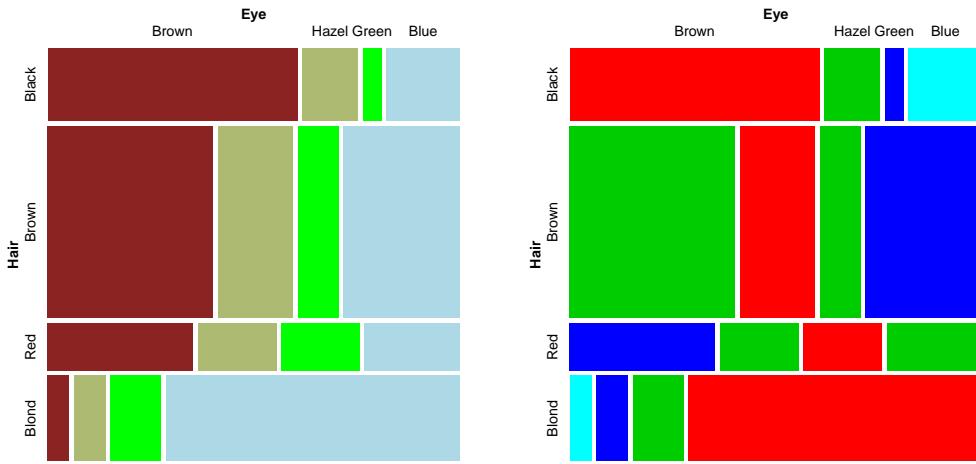


Figure 5.6: Mosaic displays for the haireye2 data, using custom colors to fill the tiles. Left: Marimekko chart, using colors to reflect the eye colors; right: Toeplitz-based colors, reflecting the diagonal strips in a square table.

{fig:HE-fill}

{ex:interp}

color to *identify* the cells, we can use these attributes to show something more—*residuals* from some model, whose pattern helps to explain the association between the table variables.

As described above, the strucplot framework includes a variety of `shading_` functions, and these can be customized with optional arguments. Zeileis *et al.* (2007) describe a general approach to residual-based shadings for area-proportional visualizations, used in the development of the strucplot framework in vcd.

EXAMPLE 5.2: Interpolation options

One simple thing to do is to modify the `interpolate` option passed to the default `shading_hcl` function, as shown in Figure 5.7.

```
> # more shading levels
> mosaic(haireye2, shade=TRUE, gp_args=list(interpolate=1:4))
>
> # continuous shading
> interp <- function(x) pmin(x/6, 1)
> mosaic(haireye2, shade=TRUE, gp_args=list(interpolate=interp))
```

{ex:shading}

For the left panel of Figure 5.7, a numeric vector is passed as `interpolate=1:4`, defining the boundaries of a step function mapping the absolute values of residuals to saturation levels in the HCL color scheme. For the right panel, a user-defined function, `interp()`, is created which maps the absolute residuals to saturation values in a continuous way (up to a maximum of 6).

Note that these two interpolation schemes produce quite similar results, differing mainly in the shading level of residuals within ± 1 and in the legend. In practice, the default discrete interpolation, using cutoffs of $\pm 2, \pm 4$ usually works quite well. \triangle

EXAMPLE 5.3: Shading functions

Alternatively, the names of shading functions can be passed as the `gp` argument, as shown below, producing Figure 5.8. Two shading function are illustrated here:

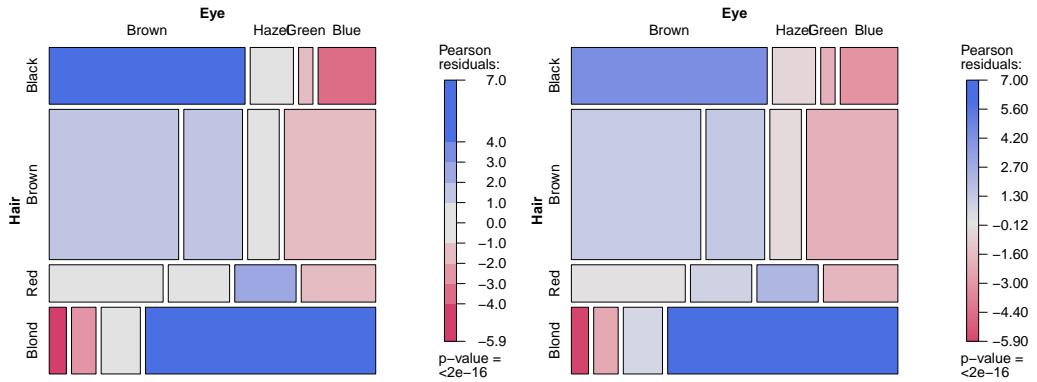


Figure 5.7: Interpolation options for shading levels in mosaic displays. Left: `shading_Friendly`; right: continuous shading.

- The left panel of Figure 5.8 uses the classical Friendly (1994b) shading scheme, `shading_Friendly` with HSV colors of blue and red and default cutoffs for absolute residuals, $\pm 2, \pm 4$, corresponding to `interpolate = c(2, 4)`. In this shading scheme, all tiles use an outline color (`coul`) corresponding to the sign of the residual. As well, the border line type (`lty`) distinguishes positive and negative residuals, which is useful if a mosaic plot is printed in black and white.
- The right panel uses the `shading_max()` function, based on the ideas of Zeileis *et al.* (2007) on residual-based shadings for area-proportional visualizations. Instead of using the cut-offs 2 and 4, it employs the critical values, M_α , for the maximum absolute Pearson residual statistic,

$$M = \max_{i,j} |r_{ij}| ,$$

by default at $\alpha = 0.10$ and 0.01 .² Only those residuals with $|r_{ij}| > M_\alpha$ are colored in the plot, using two levels for Value (“lightness”) in HSV color space. Consequently, all color in the plot signals a significant departure from independence at 90% or 99% significance level, respectively.³

```
> mosaic(haireye2, gp=shading_Friendly, legend=legend_fixed)
> set.seed(1234)
> mosaic(haireye2, gp=shading_max)
```

In this example, the difference between these two shading schemes is largely cosmetic, in that the pattern of association is similar in the two panels of Figure 5.8, and the interpretation would be the same. This is not always the case, as we will see in the next example. \triangle

²These default significance levels were chosen because this leads to displays where fully colored cells are clearly significant ($p < 0.01$), cells without color are clearly non-significant ($p > 0.1$), and cells in between can be considered to be weakly significant ($0.01 \leq p \leq 0.1$).

³This computation uses the `vcd` function `coinddep_test()` to calculate generalized tests of (conditional) independence by simulation from the marginal distribution of the input table under (conditional) independence. In these examples using `shading_max`, the function `set.seed()` is used to initialize the random number generators to a given state for reproducibility.

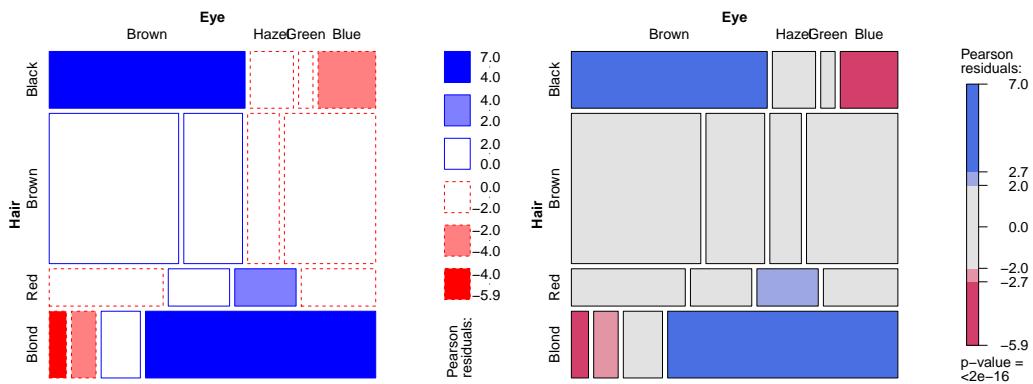


Figure 5.8: Shading functions for mosaic displays. Left: `shading_Friendly` using fixed cut-offs and the “Friendly” color scheme; right: `shading_max`, using a permutation-based test to determine significance of residuals.^{Fig:HE-shading}

```
{ex:arth-mosaic}
```

EXAMPLE 5.4: Arthritis treatment

This example uses the *Arthritis* data, illustrated earlier (Example ??), on the relation between treatment and outcome for rheumatoid arthritis. To confine this example to a two-way table, we use only the (larger) female patient group.

```
> art <- xtabs(~ Treatment + Improved, data = Arthritis,
+               subset = Sex == "Female")
> names(dimnames(art)) [2] <- "Improvement"
```

The calls to `mosaic()` below compare `shading_Friendly` and `shading_max`, giving the plots shown in Figure 5.9.

```
> mosaic(art, gp=shading_Friendly, margin = c(right = 1),
+         labeling=labeling_residuals, suppress=0, digits=2)
> set.seed(1234)
> mosaic(art, gp=shading_max, margin = c(right = 1))
```

This data set is somewhat paradoxical, in that the standard `chisq.test()` for association with these data gives a highly significant result, $\chi^2(2) = 11.3, p = 0.0035$, while the shading pattern using `shading_Friendly` in the left panel of Figure 5.9 shows all residuals within ± 2 , and thus unshaded.

On the other hand, the `shading_max` shading in the right panel of Figure 5.9 shows that significant deviations from independence occur in the four corner cells, corresponding to more of the treated group showing marked improvement, and more of the placebo group showing no improvement.

Some details behind the `shading_max` method are shown below. The Pearson residuals for this table are calculated as:

```
> residuals(loglm(~Improvement + Treatment, data=art), type="pearson")
```

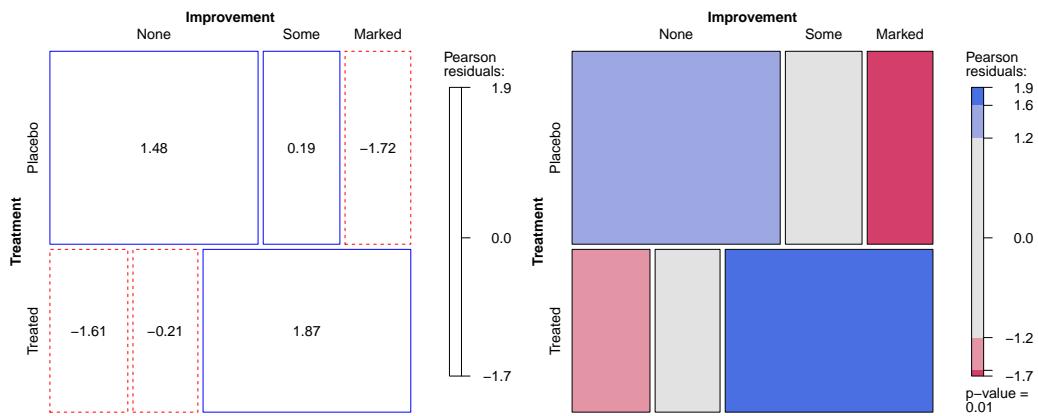


Figure 5.9: Mosaic plots for the female patients in the `Arthritis` data. Left: Fixed shading levels via `shading_Friendly`; right: shading levels determined by significant maximum residuals via `shading_max`.

```
Re-fitting to get frequencies and fitted values
Improvement
Treatment      None    Some   Marked
Placebo     1.4775  0.19267 -1.7173
Treated    -1.6085 -0.20975  1.8696
```

The `shading_max()` function then calls `coinddep_test(art)` to generate $n = 1000$ random tables with the same margins, and computes the maximum residual statistic for each. This gives a non-parametric p -value for the test of independence, $p = 0.011$ shown in the legend.

```
> set.seed(1243)
> art_max <- coinddep_test(art)
> art_max
```

```
Permutation test for conditional independence
data: art
f(x) = 1.8696, p-value = 0.011
```

Finally, the 0.90 and 0.99 quantiles of the simulation distribution are used as shading levels, passed as the value of the `interpolate` argument.

```
> art_max$qdist(c(0.90, 0.99))
 90%    99%
1.2393 1.9167
```



The converse situation can also arise in practice. An overall test for association using Pearson's χ^2 may not be significant, but the maximum residual test may highlight one or more cells worthy of greater attention, as illustrated in the following example.

{ex:soccer2}

EXAMPLE 5.5: UK Soccer scores

In Example 3.9, we examined the distribution of goals scored by the home team and the away team in 380 games in the 1995/96 season by the 20 teams in the UK Football Association, Premier League. The analysis there focused on the distribution of the total goals scored, under the assumption that the number of goals scored by the home team and the away team were independent.

Here, the rows and columns of the table *UKSoccer* are both ordered, so it is convenient and compact to carry out all the CMH tests taking ordinality into account.

```
> data("UKSoccer", package="vcd")
> CMHtest(UKSoccer)

Cochran-Mantel-Haenszel Statistics for Home by Away

      AltHypothesis Chisq Df  Prob
cor      Nonzero correlation 1.01  1 0.315
cmeans   Col mean scores differ 5.63  4 0.229
rmeans   Row mean scores differ 7.42  4 0.115
general  General association 18.65 16 0.287
```

All of these are non-significant, so that might well be the end of the story, as far as independence of goals in home and away games is concerned. Yet, one residual, $r_{42} = 3.08$ stands out, corresponding to 4 or more goals by the home team and only 2 goals by the away team, which accounts for nearly half of the $\chi^2(16) = 18.7$ for general association.

```
> set.seed(1234)
> mosaic(UKSoccer, gp=shading_max, labeling=labeling_residuals, digits=2)
```

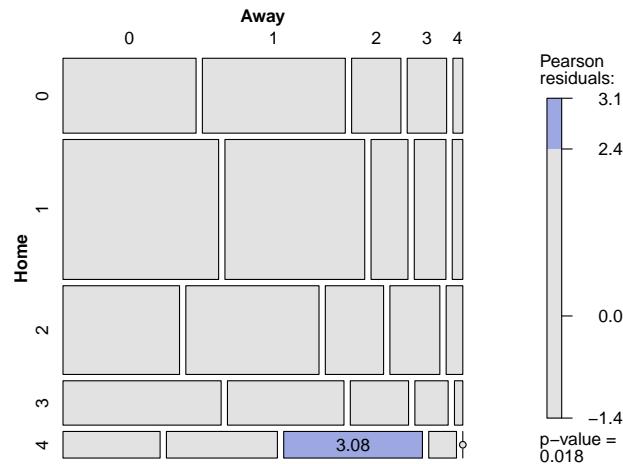


Figure 5.10: Mosaic display for UK soccer scores, highlighting one cell that stands out for further attention^{fig:UKSoccer-mosaic}

This occurrence may or may not turn out to have some explanation, but at least the mosaic plot draws it to our attention. \triangle

5.4 Three-way and larger tables

The mosaic displays and other graphical methods within the strucplot framework extend quite naturally to three-way and higher-way tables. The essential idea is that for the variables in a multiway table in a given order, each successive variable is used to subdivide the tile(s) in proportion to the relative (conditional) frequencies of that variable, given all previous variables. This process continues recursively until all table variables have been included.

{sec:mosaic-threeway}

For simplicity, we continue with the running example of Hair color and Eye color. Imagine that each cell of the two-way table for Hair and Eye color is further classified by one or more additional variables—sex and level of education, for example. Then each rectangle can be subdivided horizontally to show the proportion of males and females in that cell, and each of those horizontal portions can be subdivided vertically to show the proportions of people at each educational level in the hair-eye-sex group.

{ex:HEC1}

EXAMPLE 5.6: Hair color, eye color and sex

Figure 5.11 shows the mosaic for the three-way table, with Hair and Eye color groups divided according to the proportions of Males and Females. As explained in the next section (Section 5.4.1) there are different models for “independence” we could display. Here, we show residuals for the model of joint independence, [HairEye][Sex], which asserts that the combinations of Hair color and Eye color are independent of Sex. This model, and the corresponding mosaic plot does *not* show the (overall) association between Hair color and Eye color we explored in earlier examples (see Figure 5.3). It merely shows how where the Hair color–Eye color combinations might differ by Sex.

In the call to `mosaic()` below, the model of joint independence is specified as the argument `expected = ~ Hair*Eye + Sex`. The strucplot labeling function `labeling_residuals` is used to display the residuals in the highlighted cells.

```
> HEC <- HairEyeColor[, c("Brown", "Hazel", "Green", "Blue"), ]
> mosaic(HEC, expected = ~ Hair*Eye + Sex,
+         labeling=labeling_residuals, digits=2)
```

In Figure 5.11 it is easy to see that there is no systematic association between sex and the combinations of Hair and Eye color—except among blue-eyed blonds, where there are an overabundance of females.

The model of joint independence has a non-significant Pearson $\chi^2(15) = 19.567, p = 0.189$. Yet, the two largest residuals highlighted in the plot account for nearly half ($-2.15^2 + 2.03^2 = 8.74$) of the lack of fit, and so are worthy of attention here. An easy (probably facile) interpretation is that among the blue-eyed blonds, some of the females benefited from hair products. \triangle

5.4.1 Fitting models

When three or more variables are represented in a table, we can fit several different models of types of “independence” and display the residuals from each model. We treat these models as null or **baseline models**, which may not fit the data particularly well. The deviations of observed frequencies from expected ones, displayed by shading, will often suggest terms to be added to an explanatory model that achieves a better fit.

For a three-way table, with variables A , B and C , some of the hypothesized models which can be fit are described below and summarized in Table 5.2. Here we use `[•]` notation to list the **high-order terms** in a hierarchical loglinear model; these correspond to the margins of the table which are fitted exactly, and which translate directly into R formulas used in `loglm()` and `mosaic(..., expected=)`. **TODO:** Tweak the association diagrams here to use smaller circles, allowing longer connecting lines.

sec:mosaic-fitting}

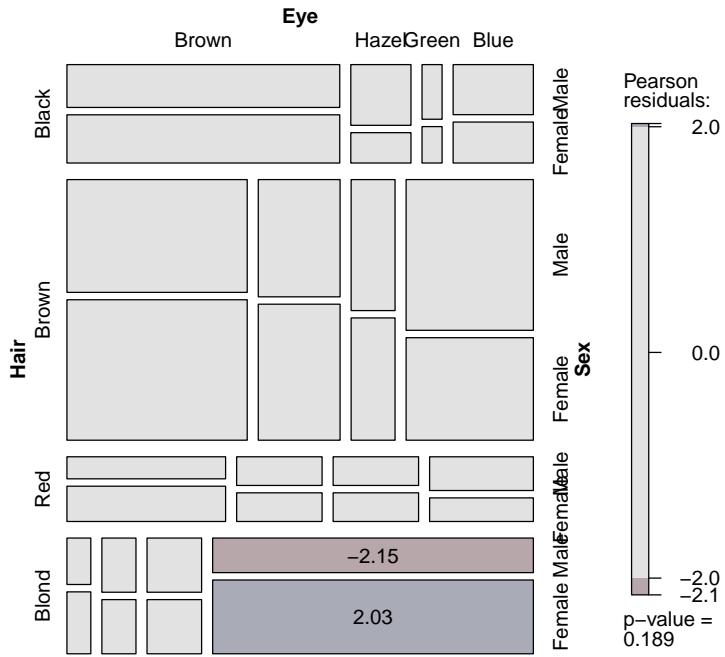


Figure 5.11: Three-way mosaic for Hair color, Eye color and Sex. Residuals from the model of joint independence, [HE][S] are shown by shading.⁴

The notation [AB][AC], for example, is shorthand for the model $\text{log}\text{lm}(\sim A*B + A*C)$ that implies

$$\{ \text{eq:AB-AC} \} \quad \log m_{ijk} = \mu + \lambda_i^A + \lambda_j^B + \lambda_k^C + \lambda_{ij}^{AB} + \lambda_{ik}^{AC}, \quad (5.1)$$

(as described in Section 8.2) and reproduces the {AB} and {AC} marginal subtables.⁴ That is, the calculated expected frequencies in these margins are always equal to the corresponding observed frequencies, $m_{i+j+} = n_{i+j+}$ and $m_{i+k+} = n_{i+k+}$.

In this table, $A \perp B$ is read, “A is independent of B.” The independence interpretation of the model Eqn. (5.1) is $B \perp C | A$, which can be read as “B is independent of C, given (conditional on) A.” Table 5.2 also depicts the relations among variables as an **association graph**, where associated variables are connected by an edge and variables that are asserted to be independent are unconnected. In mosaic-like displays, other associations present in the data will appear in the pattern of residuals.

For a three-way table, there are four general classes of independence models illustrated in Table 5.2, as described below.⁵ Not included here is the **saturated model**, [ABC], which fits the observed data exactly.

H₁: Complete independence. The model of complete (mutual) independence, symbolized $A \perp B \perp C$, with model formula $\sim A + B + C$, asserts that all joint probabilities are products of the one-way marginal probabilities:

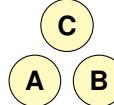
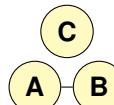
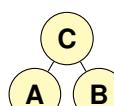
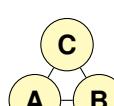
$$\pi_{ijk} = \pi_{i++} \pi_{+j+} \pi_{++k},$$

⁴The notation here uses curly braces, {•} to indicate a marginal subtable summed over all other variables.

⁵For H_2 and H_3 , permutation of the variables A, B, and C gives other members of each class.

Table 5.2: Fitted margins, model symbols and interpretations for some hypotheses for a three-way table.

{tab:hyp3way}

Hypothesis	Fitted margins	Model symbol	Independence interpretation	Association graph
H_1	$n_{i++}, n_{+j+}, n_{++k}$	$[A][B][C]$	$A \perp B \perp C$	
H_2	n_{ij+}, n_{++k}	$[AB][C]$	$(A, B) \perp C$	
H_3	n_{i+k}, n_{+jk}	$[AC][BC]$	$A \perp B C$	
H_4	$n_{ij+}, n_{i+k}, n_{+jk}$	$[AB][AC][BC]$	NA	

for all i, j, k in a three-way table. This corresponds to the log-linear model $[A][B][C]$. Fitting this model puts all higher terms, and hence all association among the variables, into the residuals.

H_2 : Joint independence. Another possibility is to fit the model in which variable C is jointly independent of variables A and B , ($\{A, B\} \perp C$), with model formula $\sim A * B + C$, where

$$\pi_{ijk} = \pi_{ij+} \pi_{++k} .$$

This corresponds to the loglinear model $[AB][C]$. Residuals from this model show the extent to which variable C is related to the combinations of variables A and B but they do not show any association between A and B , since that association is fitted exactly. For this model, variable C is also independent of A and B in the marginal $\{AC\}$ table (collapsing over B) and in the marginal $\{BC\}$.

H_3 : Conditional independence. Two variables, say A and B are conditionally independent given the third (C) if A and B are independent when we control for C , symbolized as $A \perp B | C$, and model formula $\sim A * C + B * C$. This means that conditional probabilities, $\pi_{ij|k}$ obey

$$\pi_{ij|k} = \pi_{i+k} \pi_{+jk} ,$$

where $\pi_{ij|k} = \pi_{ijk}/\pi_{ij+}$, $\pi_{i+k} = \pi_{i+k}/\pi_{i++}$, and $\pi_{+jk} = \pi_{+jk}/\pi_{++k}$. The corresponding loglinear models is denoted $[AC][BC]$. When this model is fit, the mosaic display shows the conditional associations between variables A and B , controlling for C , but does not show the associations between A and C , or B and C .

H₄: No three-way interaction. For this model, no pair is marginally or conditionally independent, so there is *no* independence interpretation. Nor is there a closed-form expression for the cell probabilities. However, the association between any two variables is the same at each level of the third variable. The corresponding loglinear model formula is [AB][AC][BC], indicating that all two-way margins are fit exactly and so only the three-way association is shown in the mosaic residuals.

TODO: Add a textbox or text describing the general scheme for translating among loglinear short-hand, R model formulas and independence interpretations.

{ex:HEC2}

EXAMPLE 5.7: Hair color, eye color and sex

We continue with the analysis of the *HairEyeColor* data from Example 5.6. Figure 5.11 showed the fit of the joint-independence model [HairEye][Sex], testing whether the joint distribution of hair color and eye color is associated with sex.

Any other model fit to this table will have the same size tiles in the mosaic since the areas depend on the observed frequencies; the residuals, and hence the shading of the tiles will differ. Figure 5.12 shows mosaics for two other models. Shading in the left panel shows residuals from the model of mutual independence, [Hair][Eye][Sex], and so includes all sources of association among these three variables. The right panel shows the conditional independence model, [HairSex][EyeSex] testing whether, given sex, hair color and eye color are independent. Note that the pattern of residuals here is similar to that in the two-way display, Figure 5.4, that collapsed over sex.

```
> abbrev <- list(abbreviate=c(FALSE, FALSE, 1))
> mosaic(HEC, expected = ~ Hair + Eye + Sex, labeling_args=abbrev,
+   main="Model: ~Hair + Eye + Sex")
> mosaic(HEC, expected = ~ Hair*Sex + Eye*Sex, labeling_args=abbrev,
+   main="Model: ~Hair*Sex + Eye*Sex")
```

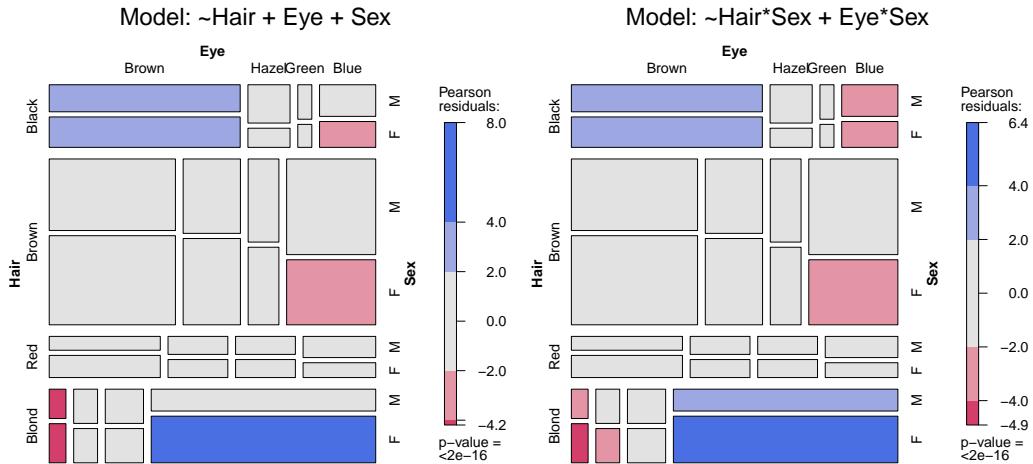


Figure 5.12: Mosaic displays for other models fit to the data on Hair Color, Eye color and Sex. Left: Mutual independence model; right: Conditional independence of Hair color and Eye color given Sex.
fig:HEC-mos2

Compared with Figure 5.11 for the joint independence model, [HairEye][Sex], it is easy to see that both of these models fit very poorly.

We consider loglinear models in more detail in Chapter 8, but for now note that these models

are fit using `loglm()` in the **MASS** package, with the model formula given in the expected argument. The details of these models can be seen by fitting these models explicitly, and the fit of several models can be summarized compactly using `LRstats()` in **vcdExtra**.

```
> library(MASS)
> mod1 <- loglm(~ Hair + Eye + Sex, data=HEC)      # mutual independence
> mod2 <- loglm(~ Hair*Sex + Eye*Sex, data=HEC)     # conditional independence
> mod3 <- loglm(~ Hair*Eye + Sex, data=HEC)         # joint independence
> LRstats(mod1, mod2, mod3)

Likelihood summary table:
    AIC BIC LR Chisq Df Pr(>Chisq)
mod1 321 333   166.3 24      <2e-16 ***
mod2 324 344   156.7 18      <2e-16 ***
mod3 193 218   19.9 15      0.18
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Alternatively, you can get the Pearson and likelihood ratio (LR) tests for a given model using `anova()`, or compare a set of models using LR tests on the *difference* in LR χ^2 from one model to the next, when a list of models is supplied to `anova()`.

```
> anova(mod1)

Call:
loglm(formula = ~Hair + Eye + Sex, data = HEC)

Statistics:
          X^2 df P(> X^2)
Likelihood Ratio 166.30 24      0
Pearson        164.92 24      0

> anova(mod1, mod2, mod3, test="chisq")

LR tests for hierarchical log-linear models

Model 1:
~Hair + Eye + Sex
Model 2:
~Hair * Sex + Eye * Sex
Model 3:
~Hair * Eye + Sex

          Deviance df Delta(Dev) Delta(df) P(> Delta(Dev))
Model 1    166.300 24
Model 2    156.678 18      9.6222      6      0.14149
Model 3    19.857 15     136.8213      3      0.00000
Saturated   0.000  0     19.8566     15      0.17750
```



5.4.2 Sequential plots and models

{sec:mosaic-seq}

As described in Section 5.2, we can think of the mosaic display for an n -way table as being constructed in stages, with the variables listed in a given order, and the unit tile decomposed recursively as each variable is entered in turn. This process turns out to have the useful property that it provides an additive (hierarchical) decomposition of the total association in a table, in a way analogous to sequential fitting with Type I sum of squares in regression models.

Typically, we just view the mosaic and fit models to the full n -way table, but it is useful to understand the connection with models for the marginal subtables, defined by summing over all

variables not yet entered. For example for a three-way table with variables, A, B, C , the marginal subtables $\{A\}$ and $\{AB\}$ are calculated in the process of constructing the three-way mosaic. The $\{A\}$ marginal table can be fit to a model where the categories of variable A are equiprobable as shown in Figure 5.2 (or some other discrete distribution); the independence model can be fit to the $\{AB\}$ subtable as in Figure 5.2 and so forth.

This connection can be seen in the following formula that decomposes the joint cell probability in an n -way table with variables v_1, v_2, \dots, v_n as a sequential product of conditional probabilities,

$$p_{ijkl\dots} = \underbrace{p_i \times p_j|_i \times p_k|_{ij}}_{\{v_1 v_2 v_3\}} \times p_{\ell|ijk} \times \dots \times p_{n|ijk\dots} \quad (5.2) \quad \{\text{eq:seqprod}\}$$

In Eqn. (5.2), the first term corresponds to the one-way mosaic for v_1 , the first two terms to the mosaic for v_1 and v_2 , the first three terms to the mosaic for v_1 , v_2 and v_3 , and so forth.

It can be shown (Friendly, 1994b) that this sequential product of probabilities corresponds to a set of sequential models of *joint independence*, whose likelihood ratio G^2 statistics provide an additive decomposition of the total association, $G^2_{[v_1][v_2]\dots[v_n]}$ for the mutual independence model in the full table:

$$G_{[v_1][v_2]\dots[v_n]}^2 = G_{[v_1][v_2]}^2 + G_{[v_1v_2][v_3]}^2 + G_{[v_1v_2v_3][v_4]}^2 + \dots + G_{[v_1\dots v_{n-1}][v_n]}^2 \quad (5.3) \quad \{\text{eq:seqgsq}\}$$

For example, for the hair-eye data, the mosaic displays for the [Hair] [Eye] marginal table (Figure 5.4) and the [HairEye] [Sex] table (Figure 5.11) can be viewed as representing the partition of G^2 shown as a table below:

Model	Model symbol	df	G^2
Marginal	[Hair] [Eye]	9	146.44
Joint	[Hair, Eye] [Sex]	15	19.86
Mutual	[Hair] [Eye] [Sex]	24	166.30

The decomposition in this table reflecting Eqn. (5.3) is shown as a visual equation in Figure 5.13. You can see from the shading how the two sequential submodels contribute to overall association in the model of mutual independence.

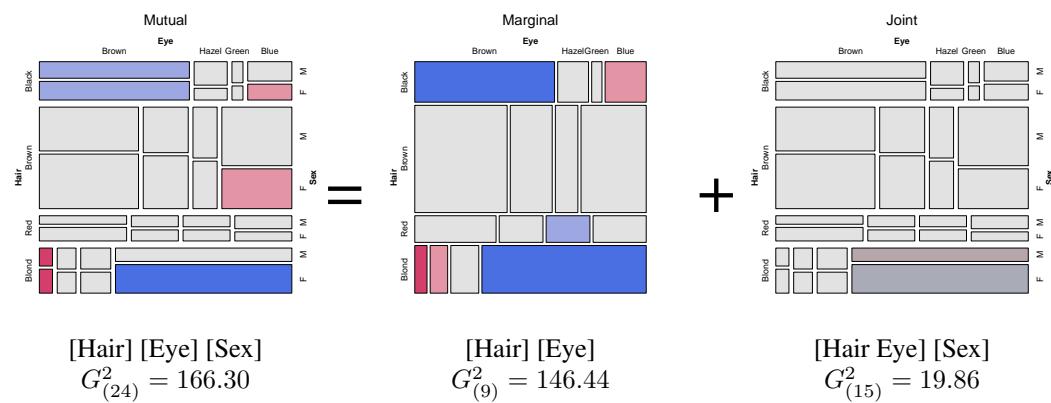


Figure 5.13: Visual representation of the decomposition of the G^2 for mutual independence (total) as the sum of marginal and joint independence.

{ fig:HEC-seq }

Although sequential models of joint independence have the nice additive property illustrated above, other classes of sequential models are possible, and sometimes of substantive interest. The

main types of these models are illustrated in Table 5.3 for 3-, 4-, and 5-way tables, with variables A, B, ... E. In all cases, the natural model for the one-way margin is the equiprobability model, and that for the two-way margin is [A][B].

Table 5.3: Classes of sequential models for n -way tables

{tab:seqmodels}

function	3-way	4-way	5-way
mutual	[A] [B] [C]	[A] [B] [C] [D]	[A] [B] [C] [D] [E]
joint	[AB] [C]	[ABC] [D]	[ABCE] [E]
joint (with=1)	[A] [BC]	[A] [BCD]	[A] [BCDE]
conditional	[AC] [BC]	[AD] [BD] [CD]	[AE] [BE] [CE] [DE]
conditional (with=1)	[AB] [AC]	[AB] [AC] [AD]	[AB] [AC] [AD] [AE]
markov (order=1)	[AB] [BC]	[AB] [BC] [CD]	[AB] [BC] [CD] [DE]
markov (order=2)	[A] [B] [C]	[ABC] [BCD]	[ABC] [BCD] [CDE]
saturated	[ABC]	[ABCD]	[ABCDE]

The `vcdExtra` package provides a collection of convenience functions that generate the loglinear model formulae symbolically, as indicated in the **function** column. The functions `mutual()`, `joint()`, `conditional()`, `markov()` and so forth simply generate a list of terms suitable for a model formula for `loglin()`. See `help(loglin-utilities)` for further details.

Wrapper functions `loglin2string()` and `loglin2formula()` convert these to character strings or model formulae respectively, for use with `loglm()` and `mosaic()`-related functions in `vcdExtra`. Some examples are shown below.

```
> for(nf in 2:5) {
+   print(loglin2string(joint(nf, factors=LETTERS[1:5])))
+ }

[1] "[A] [B]"
[1] "[A,B] [C]"
[1] "[A,B,C] [D]"
[1] "[A,B,C,D] [E]"

> for(nf in 2:5) {
+   print(loglin2string(conditional(nf, factors=LETTERS[1:5]), sep=""))
+ }

[1] "[A] [B]"
[1] "[AC] [BC]"
[1] "[AD] [BD] [CD]"
[1] "[AE] [BE] [CE] [DE]"

> for(nf in 2:5) {
+   print(loglin2formula(conditional(nf, factors=LETTERS[1:5])))
+ }

~A + B
~A:C + B:C
~A:D + B:D + C:D
~A:E + B:E + C:E + D:E
```

Applied to data, these functions take a `table` argument, and deliver the string or formula representation of a type of model for that table:

```
> loglin2formula(joint(3, table=HEC))
~Hair:Eye + Sex
> loglin2string(joint(3, table=HEC))
[1] "[Hair, Eye] [Sex]"
```

Their main use, however, is within higher-level functions, such as `seq_loglm()`, which fit the collection of sequential models of a given type.

```
> HEC.mods <- seq_loglm(HEC, type="joint")
> LRstats(HEC.mods)

Likelihood summary table:
      AIC BIC LR Chisq Df Pr(>Chisq)
model.1 194 194    165.6 3     <2e-16 ***
model.2 241 246    146.4 9     <2e-16 ***
model.3 193 218     19.9 15      0.18
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

In this section we have described a variety of models which can be fit to higher-way tables, some relations among those models, and the aspects of lack-of-fit which are revealed in the mosaic displays. The following examples illustrate the process of model fitting, using the mosaic as an interpretive guide to the nature of associations among the variables. In general, we start with a minimal baseline model.⁶ The pattern of residuals in the mosaic will suggest associations to be added to an adequate explanatory model. As the model achieves better fit to the data, the degree of shading decreases, so we may think of the process of model fitting as “cleaning the mosaic.”

5.4.3 Causal models

The sequence of models of joint independence has another interpretation when the ordering of the variables is based on a set of ordered hypotheses involving causal relationships among variables (Goodman (1973), Fienberg (1980, §7.2)). Suppose, for example, that the causal ordering of four variables is $A \rightarrow B \rightarrow C \rightarrow D$, where the arrow means “is antecedent to.” Goodman suggests that the conditional joint probabilities of B , C , and D given A can be characterized by a set of recursive logit models which treat (a) B as a response to A , (b) C as a response to A and B jointly, (c) and D as a response to A , B and C . These are equivalent to the loglinear models which we fit as the sequential baseline models of joint independence, namely $[A][B]$, $[AB][C]$, and $[ABC][D]$. The combination of these models with the marginal probabilities of A gives a characterization of the joint probabilities of all four variables, as in Eqn. (5.2). In application, residuals from each submodel show the associations that remain unexplained.

{sec:causal}

{ex:marital1}

EXAMPLE 5.8: Marital status and pre- and extramarital sex

A study of divorce patterns in relation to premarital and extramarital sex by Thornes and Collard (1979) reported the 2^4 table shown below, and included in `vcd` as `PreSex`.

```
> data("PreSex", package="vcd")
> structable(Gender+PremaritalSex+ExtramaritalSex ~ MaritalStatus, PreSex)
```

⁶When one variable, R is a response, this normally is the model of joint independence, $[E_1 E_2 \dots] [R]$, where E_1, E_2, \dots are the explanatory variables. Better-fitting models will often include associations of the form $[E_i R]$, $[E_i E_j R] \dots$

	Gender	Women			Men		
	PremaritalSex	Yes	No		Yes	No	
	ExtramaritalSex	Yes	No	Yes	No	Yes	No
MaritalStatus							
Divorced		17	54	36	214	28	60
Married		4	25	4	322	11	42
						4	130

These data were analysed by Agresti (2013, §6.1.7) and by Friendly (1994b, 2000), from which this account draws. A sample of about 500 people who had petitioned for divorce, and a similar number of married people were asked two questions regarding their pre- and extramarital sexual experience: (1) “Before you married your (former) husband/wife, had you ever made love with anyone else?,” (2) “During your (former) marriage (did you) have you had any affairs or brief sexual encounters with another man/woman?” The table variables are thus gender (G), reported premarital (P) and extramarital (E) sex, and current marital status (M).

In this analysis we consider the variables in the order G , P , E , and M , and first reorder the table variables for convenience.

```
> PreSex <- aperm(PreSex, 4:1)    # order variables G, P, E, M
```

That is, the first stage treats P as a response to G and examines the [Gender][Pre] mosaic to assess whether gender has an effect on premarital sex. The second stage treats E as a response to G and P jointly; the mosaic for [Gender, Pre] [Extra] shows whether extramarital sex is related to either gender or premarital sex. These are shown in Figure 5.14.

```
> # (Gender Pre)
> mosaic(margin.table(PreSex, 1:2), shade=TRUE,
+         main = "Gender and Premarital Sex")
>
> ## (Gender Pre) (Extra)
> mosaic(margin.table(PreSex, 1:3),
+         expected = ~Gender * PremaritalSex + ExtramaritalSex ,
+         main = "Gender*Pre + ExtramaritalSex")
```

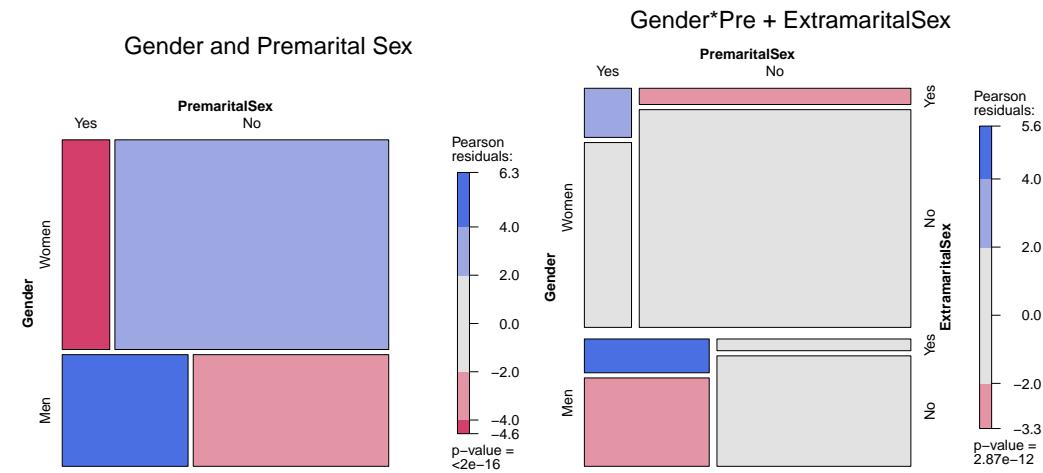


Figure 5.14: Mosaic displays for the first two marginal tables in the PreSex data. Left: `margin.table(PreSex, 1:2)`; Right: `margin.table(PreSex, 1:3)`

Finally, the mosaic for [Gender, Pre, Extra] [Marital] is examined for evidence of the dependence of marital status on the three previous variables jointly. As noted above, these models are equivalent to the recursive logit models whose path diagram is $G \rightarrow P \rightarrow E \rightarrow M$.⁷ The G^2 values for these models shown below provide a decomposition of the G^2 for the model of complete independence fit to the full table.

Model	df	G^2
[G] [P]	1	75.259
[GP] [E]	3	48.929
[GPE] [M]	7	107.956
[G] [P] [E] [M]	11	232.142

The [Gender] [Pre] mosaic in the left panel of Figure 5.14 shows that men are much more likely to report premarital sex than are women; the sample odds ratio is 3.7. We also see that women are about twice as prevalent as men in this sample. The mosaic for the model of joint independence, [Gender Pre] [Extra] in the right panel of Figure 5.14 shows that extramarital sex depends on gender and premarital sex jointly. From the pattern of residuals in Figure 5.14 we see that men and women who have reported premarital sex are far more likely to report extramarital sex than those who have not. In this three-way marginal table, the conditional odds ratio of extramarital sex given premarital sex is nearly the same for both genders (3.61 for men and 3.56 for women). Thus, extramarital sex depends on premarital sex, but not on gender.

```
> oddsratio(margin.table(PreSex, 1:3), stratum=1, log=FALSE)
Women      Men
3.562 3.605

> ## (Gender Pre Extra) (Marital)
> mosaic(PreSex,
+         expected = ~Gender*PremaritalSex*ExtramaritalSex
+                     + MaritalStatus,
+         main = "Gender*Pre*Extra + MaritalStatus")
> ## (GPE) (PEM)
> mosaic(PreSex,
+         expected = ~ Gender * PremaritalSex * ExtramaritalSex
+                     + MaritalStatus * PremaritalSex * ExtramaritalSex,
+         main = "G*P*E + P*E*M")
```

△

TODO: Complete this section with other examples: Titanic

5.4.4 Partial association

In a three-way (or larger) table it may be that two variables, say A and B , are associated at some levels of the third variable, C , but not at other levels of C . More generally, we may wish to explore whether and how the association among two (or more) variables in a contingency table varies over the levels of the remaining variables. The term **partial association** refers to the association among some variables within the levels of the other variables.

Partial association represents a useful “divide and conquer” statistical strategy: it allows you to refine the question you want to answer for complex relations by breaking it down to smaller, easier

{sec:mospart}

⁷Agresti (2013, Figure 6.1) considers a slightly more complex, but more realistic model in which premarital sex affects both the propensity to have extramarital sex and subsequent marital status.

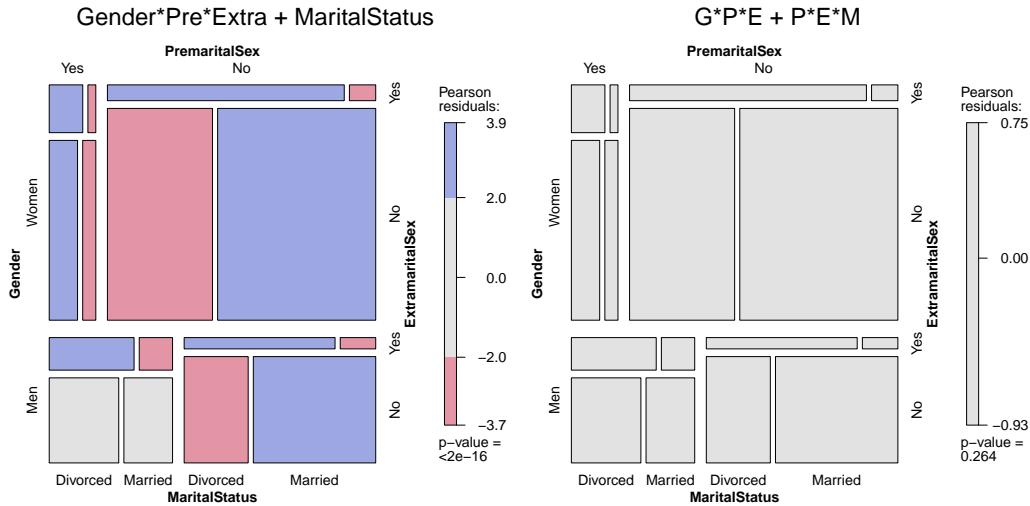


Figure 5.15: Four-way mosaics for the PreSex data. The left panel fits the model [GPE][M]. The pattern of residuals suggests other associations with marital status. The right panel fits the model [GPE][PEM].
Fig. preses

questions.⁸ It is a statistically happy fact that an answer to the larger, more complex question can be expressed as an algebraic sum of the answers to the smaller questions, just as was the case with sequential models of joint independence.

For concreteness, consider the case where you want to understand the relationship between *attitude* toward corporal punishment of children by parents or teachers (Never, Moderate use OK) and *memory* that the respondent had experiences corporal punishment as a child (Yes, No). But you also have measured other variables on the respondents, including their level of *education* and *age* category. In this case, the question of association among all the table variables may be complex, but we can answer a highly relevant, specialized question precisely, “is there an association between attitude and memory, *controlling for education and age?*” The answer to this question can be thought of as the sum of the answers to the simpler question of association between attitude and memory across the education, age categories.

A simpler version of this idea is considered first below (Example 5.9): among workers who were laid off due to either the closure of a plant or business vs. replacement by another worker, the (conditional) relationship of employment status (new job vs. still unemployed) and duration of unemployment can be studied as a sum of the associations between these focal variables over the separate tables for cause of layoff.

To make this idea precise, consider for example the model of conditional independence, $A \perp\!\!\!\perp C | B$ for a three-way table. This model asserts that A and B are independent within *each* level of C . Denote the hypothesis that A and B are independent at level $C(k)$ by $A \perp\!\!\!\perp B | C(k)$, $k = 1, \dots, K$. Then one can show (Andersen, 1991) that

$$\{eq:partial1\} G^2_{A \perp\!\!\!\perp B | C} = \sum_k^K G^2_{A \perp\!\!\!\perp B | C(k)} \quad (5.4)$$

That is, the overall likelihood ratio G^2 for the conditional independence model with $(I-1)(J-1)K$ degrees of freedom is the sum of the values for the ordinary association between A and B over the

⁸This is an analog, for categorical data, of the ANOVA strategy for “probing interactions” by testing *simple effects* at the levels of one or more of the factors involved in a two- or higher-way interaction.

levels of C (each with $(I - 1)(J - 1)$ degrees of freedom). The same additive relationship holds for the Pearson χ^2 statistics: $\chi^2_{A \perp B | C} = \sum_k^K \chi^2_{A \perp B | C(k)}$.

Thus, (a) the overall G^2 (χ^2) may be decomposed into portions attributable to the AB association in the layers of C , and (b) the collection of mosaic displays for the dependence of A and B for each of the levels of C provides a natural visualization of this decomposition. These provide an analog, for categorical data, of the conditioning plot, or *coplot*, that Cleveland (1993b) has shown to be an effective display for quantitative data. See Friendly (1999a) for further details.

Mosaic and other displays in the strucplot framework for partial association can be produced in several different ways. One way is to use a model formula in the call to `mosaic()` which lists the conditioning variables after the "`|`" (given) symbol, as in `~ Memory + Attitude | Age + Education`. Another way is to use `cotabplot()`. This takes the same kind of conditioning model formula, but presents each panel for the conditioning variables in a separate frame within a trellis-like grid.⁹

{ex:employ}

EXAMPLE 5.9: Employment status data

Data from a 1974 Danish study of 1314 employees who had been laid off are given in the data table *Employment* in *vcd* (from Andersen (1991, Table 5.12)). The workers are classified by: (a) their employment status, on January 1, 1975 ("NewJob" or still "Unemployed"), (b) the length of their employment at the time of layoff, (c) the cause of their layoff ("Closure", etc. or "Replaced").

```
> data("Employment", package = "vcd")
> structable(Employment)
```

		EmploymentLength <1Mo 1-3Mo 3-12Mo 1-2Yr 2-5Yr >5Yr					
EmploymentStatus	LayoffCause						
		Closure	Replaced	Closure	Replaced	Closure	Replaced
NewJob	Closure	8	35	70	62	56	38
	Replaced	40	85	181	85	118	56
Unemployed	Closure	10	42	86	80	67	35
	Replaced	24	42	41	16	27	10

In this example, it is natural to regard *EmploymentStatus* (variable A) as the response variable, and *EmploymentLength* (B) and *LayoffCause* (C) as predictors. In this case, the minimal baseline model is the joint independence model, $[A][BC]$, which asserts that employment status is independent of both length and cause. This model fits quite poorly, as shown in the output from `loglm()` below.

```
> loglm(~ EmploymentStatus + EmploymentLength*LayoffCause, data=Employment)

Call:
loglm(formula = ~EmploymentStatus + EmploymentLength * LayoffCause,
      data = Employment)

Statistics:
          X^2  df P(> X^2)
Likelihood Ratio 172.28 11     0
Pearson         165.70 11     0
```

The residuals, shown in Figure 5.16, indicate an opposite pattern for the two categories of *LayoffCause*: those who were laid off as a result of a closure are more likely to be unemployed, regardless of length of time they were employed. Workers who were replaced, however, apparently are more likely to be employed, particularly if they were employed for 3 months or more.

⁹Depending on your perspective, this has the advantage of adjusting for the total frequency in each conditional panel, or the disadvantage of ignoring these differences.

```
> # baseline model [A][BC]
> mosaic(Employment, shade=TRUE,
+         expected = ~ EmploymentStatus + EmploymentLength*LayoffCause,
+         main = "EmploymentStatus + Length * Cause")
```

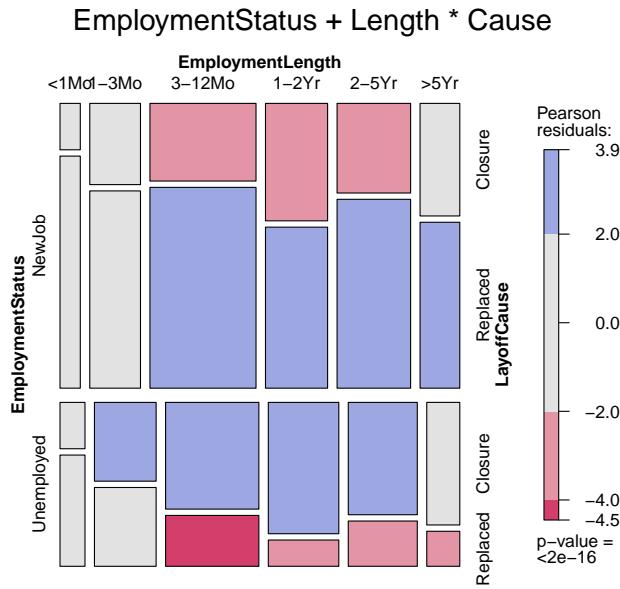


Figure 5.16: Mosaic display for the employment status data, fitting the baseline model of joint independence.
fig:employ_mos1

Beyond this baseline model, it is substantively more meaningful to consider the conditional independence model, $A \perp B | C$, (or $[AC][BC]$ in shorthand notation), which asserts that employment status is independent of length of employment, given the cause of layoff. We fit this model as shown below:

```
> loglm(~ EmploymentStatus*LayoffCause + EmploymentLength*LayoffCause,
+        data=Employment)

Call:
loglm(formula = ~EmploymentStatus * LayoffCause + EmploymentLength *
    LayoffCause, data = Employment)

Statistics:
      X^2   df   P(> X^2)
Likelihood Ratio 24.630 10 0.0060927
Pearson          26.072 10 0.0036445
```

This model fits far better ($G^2(10) = 24.63$), but the lack of fit is still significant. The residuals, shown in Figure 5.17, still suggest that the pattern of association between employment and length is different for replaced workers and those laid off due to closure of their workplace.

```
> mosaic(Employment, shade=TRUE, gp_args=list(interpolate=1:4),
+         expected = ~ EmploymentStatus*LayoffCause + EmploymentLength*LayoffCause,
+         main = "EmploymentStatus * Cause + Length * Cause")
```

To explain this result better, we can fit separate models for the *partial* relationship between

EmploymentStatus * Cause + Length * Cause

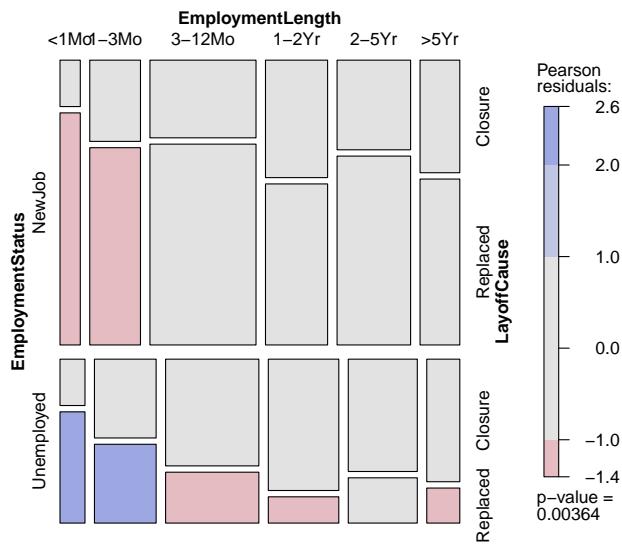


Figure 5.17: Mosaic display for the employment status data, fitting the model of conditional independence, [AC][BC].

EmploymentStatus and EmploymentLength for the two levels of LayoffCause. In R, with the *Employ* data as in table form, this is easily done using `apply()` over the LayoffCause margin, giving a list containing the two `loglm()` models.

```
> mods.list <- apply(Employment, "LayoffCause",
+                      function(x) loglm(~EmploymentStatus + EmploymentLength, data=x))
> mods.list

$Lclosure
Call:
loglm(formula = ~EmploymentStatus + EmploymentLength, data = x)

Statistics:
      X^2  df P(> X^2)
Likelihood Ratio 1.4786  5  0.91553
Pearson          1.4835  5  0.91497

$Replaced
Call:
loglm(formula = ~EmploymentStatus + EmploymentLength, data = x)

Statistics:
      X^2  df P(> X^2)
Likelihood Ratio 23.151  5  0.00031578
Pearson          24.589  5  0.00016727
```

Extracting the model fit statistics for these partial models and adding the fit statistics for the overall model of conditional independence, [AC][BC] gives the table below, illustrating the additive property of G^2 , (Eqn. (5.4)) and χ^2 .

Model	df	G^2	χ^2
$A \perp B C_1$	5	1.49	1.48
$A \perp B C_2$	5	23.15	24.59
$A \perp B C$	10	24.63	26.07

One simple way to visualize these results is to call `mosaic()` separately for each of the layers corresponding to `LayoffCause`. The result is shown in Figure 5.18.

```
> mosaic(Employment[, "Closure"], shade=TRUE, gp_args=list(interpolate=1:4),
+         margin = c(right = 1), main = "Layoff: Closure")
> mosaic(Employment[, "Replaced"], shade=TRUE, gp_args=list(interpolate=1:4),
+         margin = c(right = 1), main = "Layoff: Replaced")
```

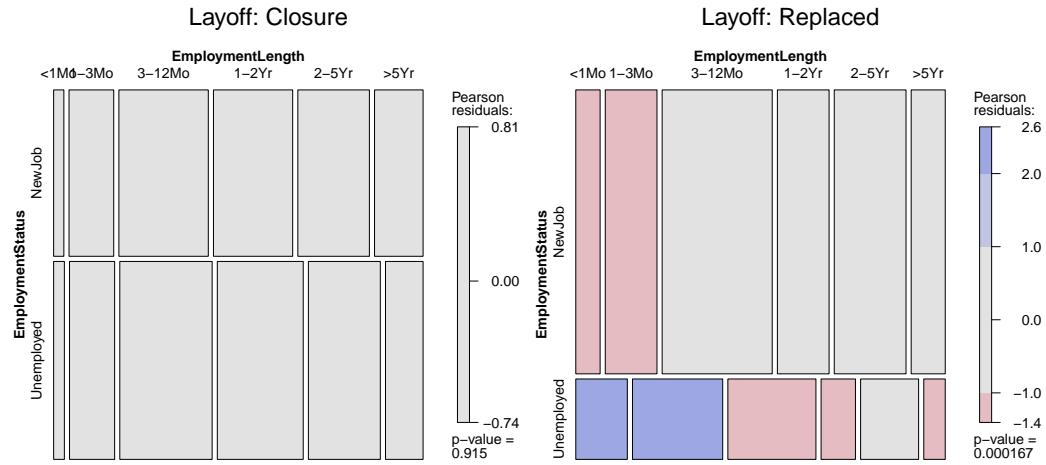


Figure 5.18: Mosaic displays for the employment status data, with separate panels for cause of layoff.
fig.employ_moss

The simple summary from this example is that for workers laid off due to closure of their company, length of previous employment is unrelated to whether or not they are re-employed. However, for workers who were replaced, there is a systematic pattern: those who had been employed for three months or less are likely to remain unemployed, while those with longer job tenure are somewhat more likely to have found a new job. \triangle

The statistical methods and R techniques described above for three-way tables extend naturally to higher-way tables, as can be seen in the next example.

EXAMPLE 5.10: Corporal punishment data

Here we use the `Punishment` data from `vcd` which contains the results of a study by the Gallup Institute in Denmark in 1979 about the attitude of a random sample of 1,456 persons towards corporal punishment of children (Andersen, 1991, pp. 207-208). As shown below, this data set is a frequency data frame representing a $2 \times 2 \times 3 \times 3$ table, with table variables (a) attitude toward use of corporal punishment (approve of “moderate” use or “no” approval) (b) memory of whether the respondent had experienced corporal punishment as a child (yes/no); (c) education level of respondent (elementary, secondary, high); (d) age category of respondent.

```
> data("Punishment", package = "vcd")
> str(Punishment)
```

```
'data.frame': 36 obs. of 5 variables:
$ Freq      : num 1 3 20 2 8 4 2 6 1 26 ...
$ attitude   : Factor w/ 2 levels "no","moderate": 1 1 1 1 1 1 1 1 1 1 ...
$ memory     : Factor w/ 2 levels "yes","no": 1 1 1 1 1 1 1 1 1 2 ...
$ education  : Factor w/ 3 levels "elementary","secondary",...: 1 1 1 2 2 2 3 3 3 1 ...
$ age        : Factor w/ 3 levels "15-24","25-39",...: 1 2 3 1 2 3 1 2 3 1 ...
```

Of main interest here is the association between attitude toward corporal punishment as an adult (A) and memory of corporal punishment as a child (B), controlling for age (C) and education (D); that is, the model $A \perp B | (C, D)$, or $[ACD][BCD]$ in shorthand notation.

As noted above, this conditional independence hypothesis can be decomposed into the 3×3 partial tests of $A \perp B | (C_k, D_\ell)$.

These tests and the associated graphics are somewhat easier to carry out with the data in table form (`pun`) constructed below. While we're at it, we recode the variable names and factor levels for nicer graphical displays.

```
> pun <- xtabs(Freq ~ memory + attitude + age + education, data = Punishment)
> dimnames(pun) <- list(
+   Memory = c("yes", "no"),
+   Attitude = c("no", "moderate"),
+   Age = c("15-24", "25-39", "40+"),
+   Education = c("Elementary", "Secondary", "High"))
```

Then, the overall test of conditional independence can be carried using `loglm()` out as

```
> (mod.cond <- loglm(~ Memory*Age*Education + Attitude*Age*Education,
+                      data = pun))

Call:
loglm(formula = ~Memory * Age * Education + Attitude * Age *
       Education, data = pun)

Statistics:
          X^2    df    P(> X^2)
Likelihood Ratio 39.679  9 8.6851e-06
Pearson          34.604  9 6.9964e-05
```

Alternatively, `coindep_test()` in `vcd` provides tests of conditional independence of two variables in a contingency table by simulation from the marginal permutation distribution of the input table. The version reporting a Pearson χ^2 statistic is given by

```
> set.seed(1071)
> coindep_test(pun, margin=c("Age", "Education"),
+               indepfun = function(x) sum(x^2), aggfun=sum)

Permutation test for conditional independence

data: pun
f(x) = 34.604, p-value < 2.2e-16
```

These tests all show substantial association between attitude and memory of corporal punishment. How can we understand and explain this?

As in Example 5.9, we can partition the overall G^2 or χ^2 to show the contributions to this association from the combinations of age and education. The call to `apply()` below returns a 3×3 matrix, each of whose elements is the list of results returned by `loglm()`. The Pearson χ^2 statistics for each subtable can be extracted using `sapply()` as shown below.

```
> mods.list <- apply(pun, c("Age", "Education"),
+                      function(x) loglm(~Memory + Attitude, data=x))
> XSQ <- matrix( sapply(mods.list, function(x)x$pearson), 3, 3)
> dimnames(XSQ) <- dimnames(mods.list)
> addmargins(XSQ)

      Education
Age   Elementary Secondary High Sum
15-24     3.5907  0.078782 0.091429 3.7609
25-39     8.5844  0.934669 0.480000 9.9990
40+      11.6256  6.094854 3.123714 20.8442
Sum      23.8006  7.108305 3.695143 34.6041
```

One visual analog of this table of χ^2 statistics is a `cotabplot()` of the (conditional) association of attitude and memory over the age and education cells, shown in Figure 5.19. `cotabplot()` is very general, allowing a variety of functions of the residuals to be used for shading (Zeileis *et al.*, 2007). Here we use the (Pearson) sum of squares statistic, $\sum_{k,\ell} \chi_{k,\ell}^2$.

```
> set.seed(1071)
> pun_cotab <- cotab_coinddep(pun, condvars = 3:4, type = "mosaic",
+                                varnames = FALSE, margins = c(2, 1, 1, 2),
+                                test = "sumchisq", interpolate = 1:2)
> cotabplot(~ Memory + Attitude | Age + Education, data =
+             pun, panel = pun_cotab)

Error in grid.Call.graphics(L_downvppath, name$path, name$name, strict): Viewport
'panel:Age=40+,Education=Elementary|base' was not found
```

Alternatively, the pattern of conditional association can be shown somewhat more directly in a conditional mosaic plot (Figure 5.20), using the same model formula to condition on age and education. This simply organizes the display to split on the conditioning variables first, with larger spacings.

```
> mosaic(~ Memory + Attitude | Age + Education, data = pun,
+         shade=TRUE, gp_args=list(interpolate=1:4))
```

Both Figure 5.19 and Figure 5.20 reveal that the association between attitude and memory becomes stronger with increasing age among those with the lowest education (first column). Among those in the highest age group (bottom row), the strength of association *decreases* with increasing education. These two displays differ in that in the `cotabplot()` of Figure 5.19 the marginal frequencies of age and education are not shown, whereas in the `mosaic()` of Figure 5.20 they determine the relative sizes of the tiles for the combinations of age and education.

The divide-and-conquer strategy of partial association using statistical tests and visual displays now provides a simple, coherent explanation for this table: memory of experienced violence as a child tends to engender a more favorable attitude toward corporal punishment as an adult, but this association varies directly with both age and education. \triangle

5.5 Mosaic matrices for categorical data

{sec:mosmat}

One reason for the wide usefulness of graphs of quantitative data has been the development of effective, general techniques for dealing with high-dimensional data set. The *scatterplot matrix* shows all pairwise (marginal) views of a set of variables in a coherent display, whose design goal is to show the interdependence among the collection of variables as a whole. It combines multiple views of the data into a single display which allows detection of patterns which could not readily be discerned from a series of separate graphs. In effect, a multivariate data set in p dimensions

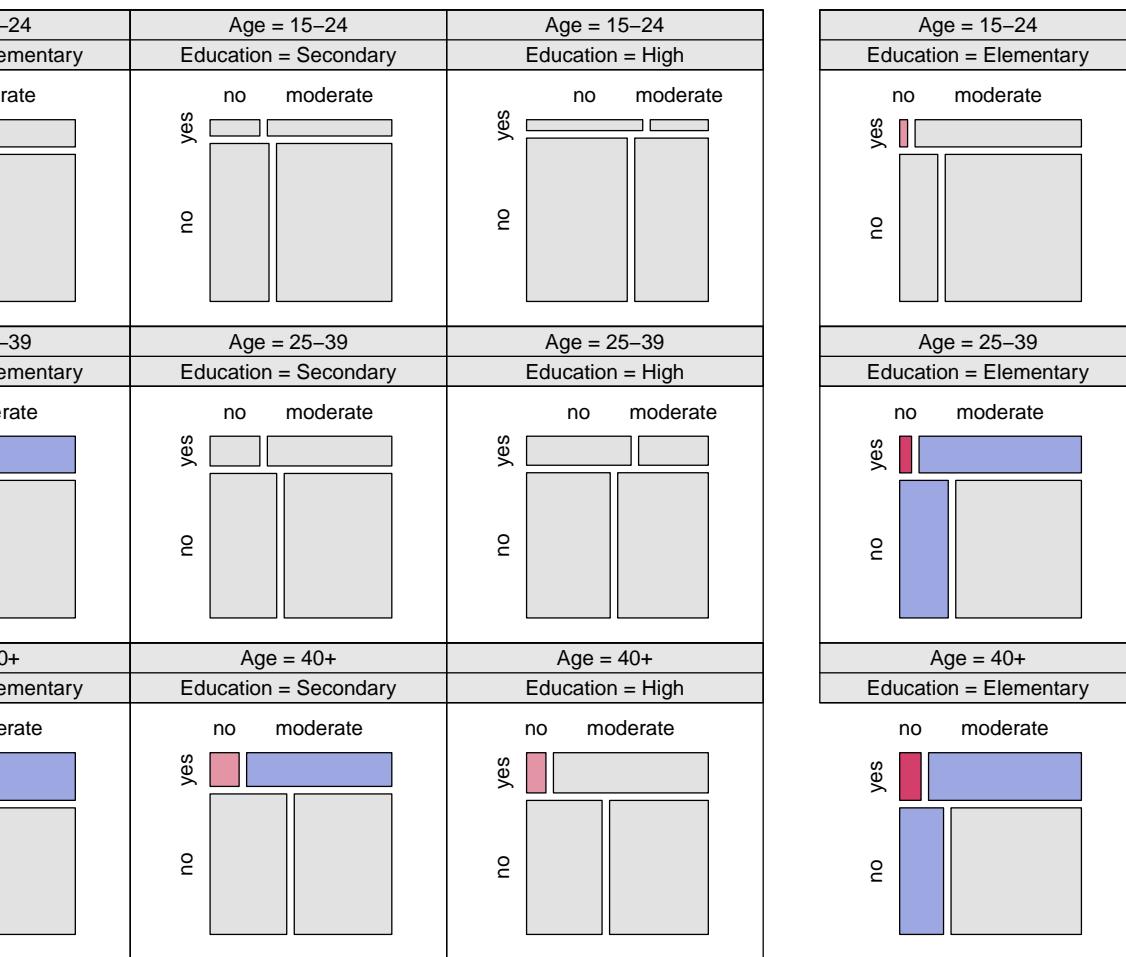


Figure 5.19: Conditional mosaic plot of the Punishment data for the model of conditional independence of attitude and memory, given age and education. Shading of tiles is based on the sum of squares statistic.
fig:punish-cond

(variables) is shown as a collection of $p(p - 1)$ two-dimensional scatterplots, each of which is the projection of the cloud of points on two of the variable axes. These ideas can be readily extended to categorical data.

A multiway contingency table of p categorical variables, A, B, C, \dots , contains the interdependence among the collection of variables as a whole. The saturated loglinear model, $[ABC\dots]$ fits this interdependence perfectly, but is often too complex to describe or understand.

By summing the table over all variables except two, A and B , say, we obtain a two-variable (marginal) table, showing the bivariate relationship between A and B , which is also a projection of the p -variable relation into the space of two (categorical) variables. If we do this for all $p(p - 1)$ unordered pairs of categorical variables and display each two-variable table as a mosaic, we have a categorical analog of the scatterplot matrix, called a **mosaic matrix**. Like the scatterplot matrix, the mosaic matrix can accommodate any number of variables in principle, but in practice is limited by the resolution of our display to three or four variables.

In R, the main implementation of this idea is in the generic function `pairs()`. The `vcd` pack-

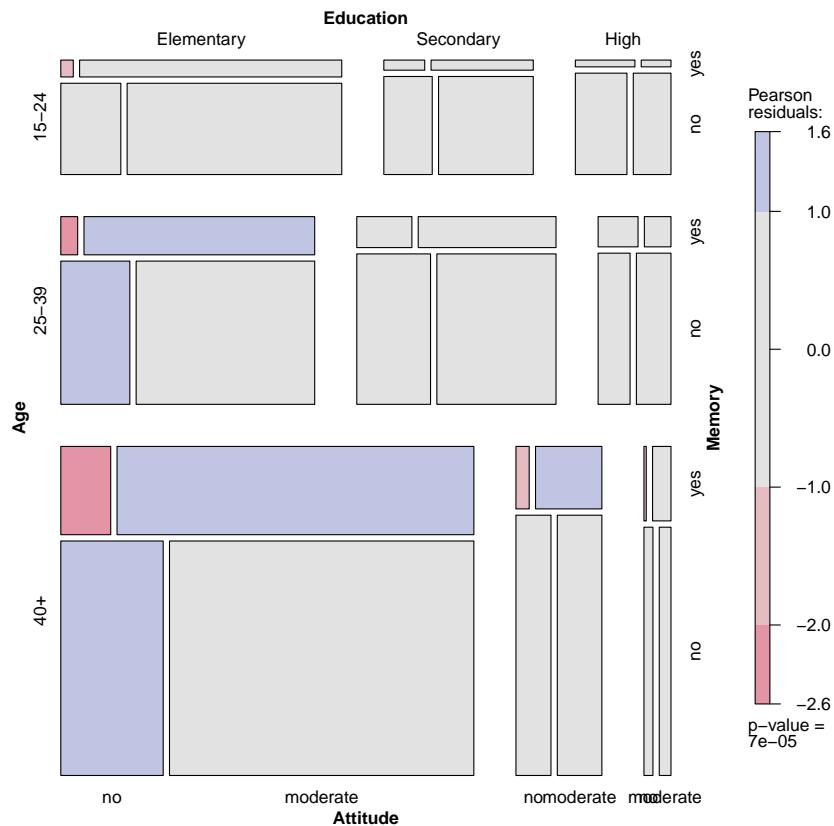


Figure 5.20: Conditional mosaic plot of the Punishment data for the model of conditional independence of attitude and memory, given age and education. This plot explicitly shows the total frequencies in the cells of age and education by the areas of the main blocks for these variables.
Fig:punish-cond2

age extends this to mosaic matrices with methods for "table" and "structable" objects. The gpairs package provides a **generalized pairs plot**, with appropriate graphics for a mixture of quantitative and categorical variables.

{ex:bartlett}

EXAMPLE 5.11: Bartlett data on plum root cuttings

The simplest example of what you can see in a mosaic matrix is provided by the $2 \times 2 \times 2$ table used by Bartlett (1935) to illustrate a method for testing for no three-way interaction in a contingency table (hypothesis H_4 in Table 5.2).

The data set *Bartlett* in *vcdExtra* gives the result of an agricultural experiment to investigate the survival of plum root cuttings (*Alive*) in relation to two factors: *Time* of planting and the *Length* of the cutting. In this experiment, 240 cuttings were planted for each of the 2×2 combinations of these factors, and their survival (*Alive*, *Dead*) was later recorded.

```
> pairs(Bartlett, gp=shading_Friendly)
```

The mosaic matrix for these data, showing all twoway marginal relations, is shown in Figure 5.21. It can immediately be seen that *Time* and *Length* are independent by the design of the experiment; we use *gp=shading_Friendly* here to emphasize this.

The top row and left column show the relation of survival to each of time of planting and cutting length. It is easily seen that greater survival is associated with cuttings taken now (vs. spring) and

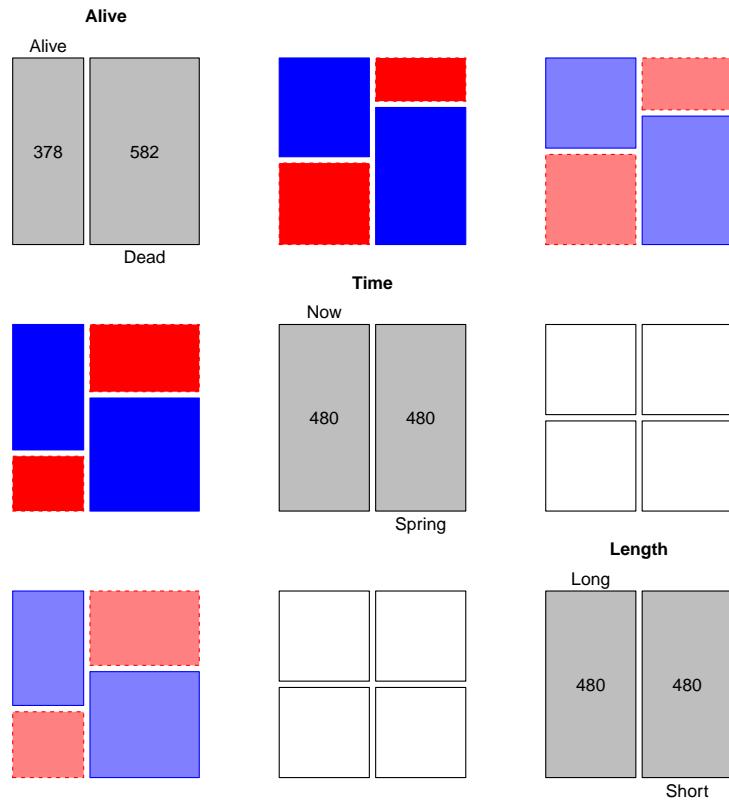


Figure 5.21: Mosaic pairs plot for the Bartlett data. Each panel shows the bivariate marginal relation between the row and column variables.
Fig.bartlett_pairs

those cut long (vs. short), and the degree of association is stronger for planting time than for cutting length. \triangle

{ex:marital2}

EXAMPLE 5.12: Marital status and pre- and extramarital sex

In Example 5.8 we examined a series of models relating marital status to reported premarital and extramarital sexual activity and gender in the *PreSex* data. Figure 5.22 shows the mosaic matrix for these data. The diagonal panels show the labels for the category levels as well as the one-way marginal totals.

```
> data("PreSex", package="vcd")
> pairs(PreSex, gp=shading_Friendly, gp_args=list(interpolate=1:4), space=0.25)
```

If we view gender, premarital sex and extramarital sex as explanatory, and marital status (Divorced vs. still Married) as the response, then the mosaics in row 1 (and in column 1)¹⁰ shows how marital status depends on each predictor marginally. The remaining panels show the relations within the set of explanatory variables.

Thus, we see in row 1, column 4, that marital status is independent of gender (all residuals equal zero, here), by design of the data collection. In the (1, 3) panel, we see that reported premarital sex is

¹⁰Rows and columns in a mosaic matrix are identified as in a table or numerical matrix, with row 1, column 1 in the upper left corner.

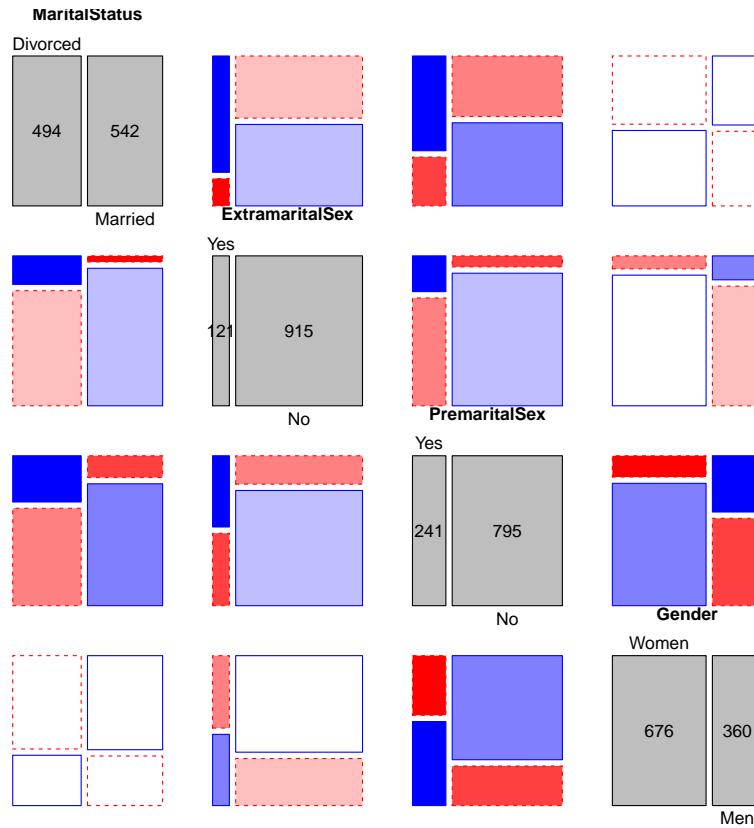


Figure 5.22: Mosaic pairs plot for the PreSex data. Each panel shows the bivariate marginal relation between the row and column variables.

more often followed by divorce, while non-report is more prevalent among those still married. The (1, 2) panel shows a similar, but stronger relation between extramarital sex and marriage stability. These effects pertain to the associations of P and E with marital status (M)—the terms [PM] and [EM] in the loglinear model. We saw earlier that an interaction of P and E (the term [PEM]) is required to fully account for these data. This effect is not displayed in Figure 5.22.

Among the background variables (the loglinear term [GPE]), the (2, 3) panel shows a strong relation between premarital sex and subsequent extramarital sex, while the (2, 4) and (3, 4) panels show that men are far more likely to report premarital sex than women in this sample, and also more likely to report extramarital sex.

Even though the mosaic matrix shows only pairwise, bivariate associations, it provides an integrated view of all of these together in a single display.

△

{ex:berkeley4}

EXAMPLE 5.13: Berkeley admissions

In Chapter 4 we examined the relations among the variables Admit, Gender and Department in the Berkeley admissions data (Example 4.1, Example 4.10, Example 4.14) using fourfold displays (Figure 4.3 and Figure 4.4) and sieve diagrams (Figure 4.10). These displays showed either a marginal relation (e.g., Admit, Gender) or the full three-way table.

In contrast, Figure 5.23 shows all pairwise marginal relations among these variables, produced

using `pairs()`. Some additional arguments are used to control the details of labels for the diagonal and off-diagonal panels.

```
> largs <- list(labeling = labeling_border(varnames = FALSE,
+                                         labels = c(T, T, F, T), alternate_labels = FALSE))
> dargs <- list(gp_varnames = gpar(fontsize = 20), offset_varnames = -1,
+                 labeling = labeling_border(alternate_labels = FALSE))
> pairs(UCBAdmissions, shade = TRUE, space = 0.25,
+        diag_panel_args = dargs,
+        upper_panel_args = largs, lower_panel_args = largs)
```

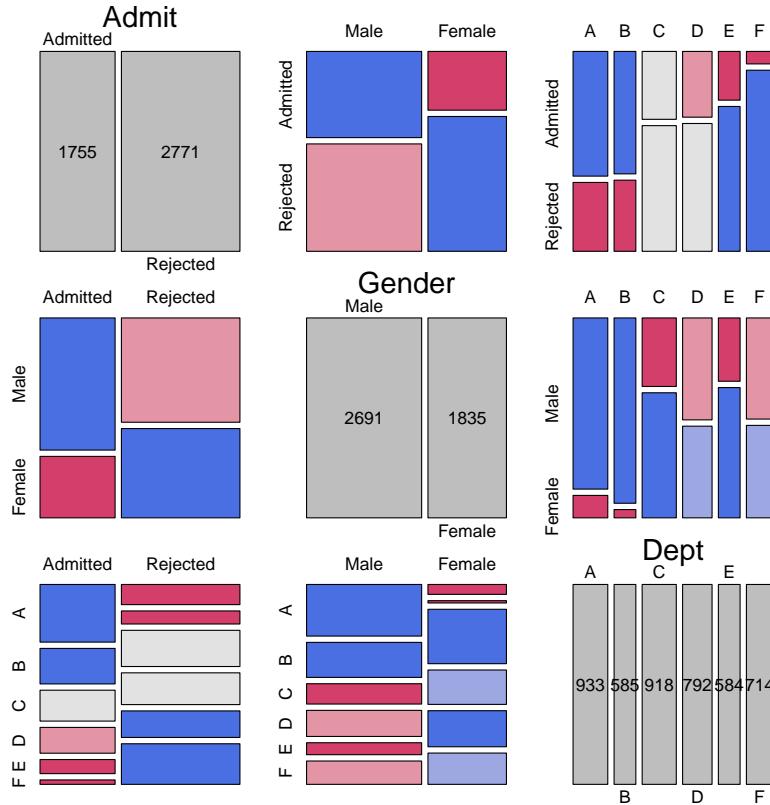


Figure 5.23: Mosaic matrix of the UCBAdmissions data showing bivariate marginal relations^{fig:berk-pairs1}

The panel in row 2, column 1 shows that Admission and Gender are strongly associated marginally, as we saw in Figure 4.3, and overall, males are more often admitted. The diagonally-opposite panel (row 1, column 2) shows the same relation, splitting first by gender.¹¹

The panels in the third column (and third row) provide the explanation for the paradoxical result (see Figure 4.4) that, within all but department A, the likelihood of admission is equal for men and women, yet, overall, there appears to be a bias in favor of admitting men (see Figure ??). The (1,3) and (3, 1) panels show the marginal relation between Admission and Department, that is, how

¹¹Note that this is different than just the transpose or interchange of horizontal and vertical dimensions as in a scatterplot matrix, because the mosaic display splits the total frequency first by the horizontal variable and then (conditionally) by the vertical variable. The areas of all corresponding tiles are the same in each diagonally opposite pair, however, as are the residuals shown by color and shading.

admission rate varies across departments. Departments A and B have the greatest overall admission rate, departments E and F the least. The (2, 3) and (3,2) panels show how men and women apply differentially to the various departments. It can be seen that men apply in much greater numbers to departments A and B, with higher admission rates, while women apply in greater numbers to the departments C–F, with the lowest overall rate of admission.



5.5.1 Generalized mosaic matrices and pairs plots

{sec:condmat}

We need not show only the marginal relation between each pair of variables in a mosaic matrix. (Friendly, 1999b) describes the extension of this idea to conditional, partial, and other views of a contingency table.

In `pairs.table()`, different *panel functions* can be used to specify what is displayed in the upper, lower and diagonal panels. For the off-diagonal panels, a `type` argument can be used to plot mosaics showing various kinds of independence relations:

```
type="pairwise" Shows bivariate marginal relations, collapsed over all other variables.  
type="total" Shows mosaic plots for mutual independence.  
type="conditional" Shows mosaic plots for conditional independence given all other variables.  
type="joint" Shows mosaic plots for joint independence of all pairs of variables from the others.
```

{ex:berkeley4b}

EXAMPLE 5.14: Berkeley admissions

Figure 5.24 shows the generalized mosaic matrix for the *UCBAdmissons* data, using 3-way mosaics for all the off-diagonal cells. The observed frequencies, of course, are the same in all these cells. However, in the lower panels, the tiles are shaded according to models of joint independence, while in the upper panels, they are shaded according to models of mutual independence.

```
> pairs(UCBAdmissons,  
+        lower_panel = pairs_mosaic(type = "joint", shade=TRUE),  
+        upper_panel = pairs_mosaic(type = "total", shade=TRUE),  
+        space=0.2)
```

TODO: Replace this with a figure using `type = "conditional"`, once we can get this to work.

In this example, it is more useful to fit and display the models of conditional independence for each pair of row, column variables given the remaining one, as shown in Figure 5.25.

```
> pairs(UCBAdmissons,  
+        lower_panel = pairs_mosaic(type = "conditional", shade=TRUE),  
+        upper_panel = pairs_mosaic(type = "conditional", shade=TRUE),  
+        space=0.2)
```

Thus, the shading in the (1,2) and (2,1) panels show the fit of the model [Admit, Dept] [Gender, Dept], which asserts that Admission and Gender are independent, given (controlling for) Department. Except for Department A, this model fits quite well, again indicating lack of gender bias. The (1,3) and (3,1) panels show the relation between admission and department controlling for gender, highlighting the differential admission rates across departments. **TODO: This isn't quite right!**



Beyond this, the framework of pairs plots can be further generalized to *mixtures* of quantitative and categorical variables, as first described in Friendly (2003) and then in a wider context by

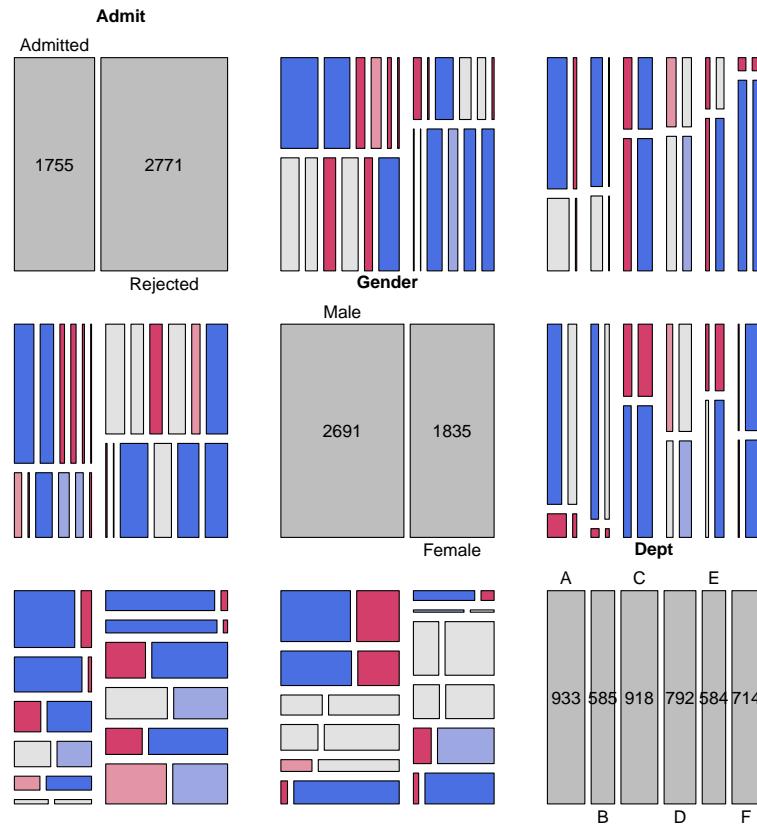


Figure 5.24: Generalized mosaic matrix of the UCBAdmissions data. The above-diagonal plots fit models of joint independence; below-diagonal plots fit models of mutual independence.
Figure 5.24 is a 3x3 grid of mosaic plots. The top row shows 'Admit' (Admitted vs. Rejected) by 'Gender' (Male vs. Female). The middle row shows 'Male' vs. 'Female' by 'Dept' (A, C, E, F). The bottom row shows 'Dept' (B, D, E, F) by 'Dept' (A, C, E, F). The plots are stacked bar charts where the height of each segment represents the count for that category combination.

Emerson *et al.* (2013), Friendly (2013). The essential idea is to consider the combination of two variables, each of which can be either categorical (**C**) or quantitative (**Q**), and various ways to *render* that combination in a graphical display:

CC: mosaic display, sieve diagram, doubledecker plot, faceted or divided bar chart;

CQ: side-by-side boxplots, stripcharts, faceted histograms, aligned density plots;

QQ: scatterplot, corrgram, data ellipses, etc.

In R some of these possibilities are provided in the **gpairs** package (using **grid** graphics and the **vcd** strucplot framework), and the **GGally** package (an extension to **ggplot2**).
(ex:arthritis-gpairs}

EXAMPLE 5.15: Arthritis treatment

We illustrate these ideas with the *Arthritis* data using the **gpairs** package in Figure ???. In this data, the variables Treatment, Sex and Improved are categorical, and Age is quantitative. The call to **gpairs()** below reorders the variables to put the response variable Improved in row 1, column 1. Various options can be passed to **mosaic()** using the **mosaic.pars** argument.

```
> library(gpairs)
Error in library(gpairs): there is no package called 'gpairs'
> data("Arthritis", package="vcd")
```

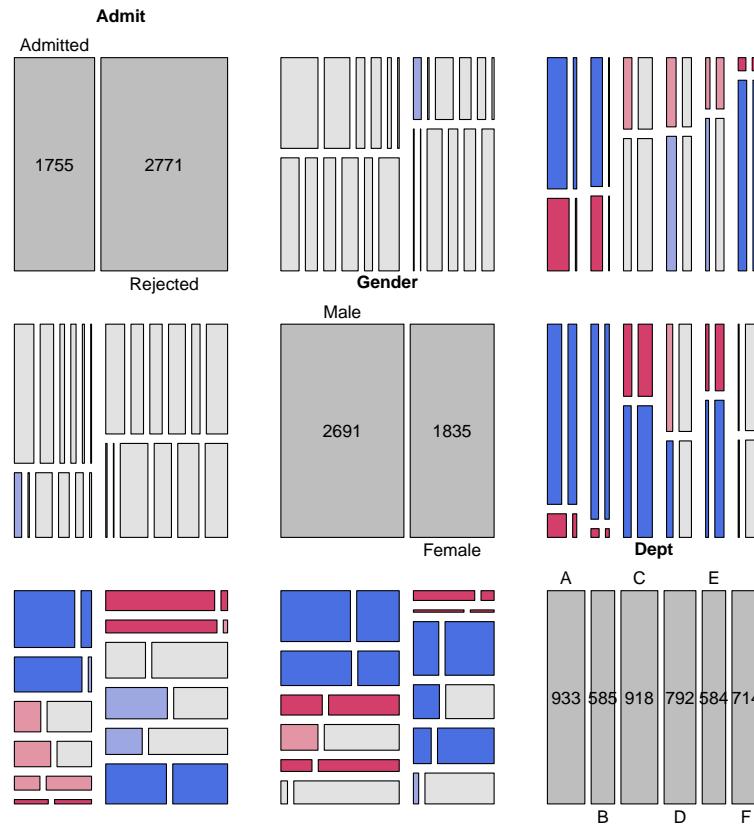


Figure 5.25: Generalized mosaic matrix of the UCBAdmissions data. The above-diagonal plots fit models of joint independence; below-diagonal plots fit models of mutual independence.

```
> gpairs(Arthritis[,c(5,2,3,4)],
+         diag.pars=list(fontsize = 20),
+         mosaic.pars=list(gp=shading_Friendly,
+                           gp_args=list(interpolate=1:4)))

Error in eval(expr, envir, enclos): could not find function "gpairs"
```

`gpairs()` provides a variety of options for the **CQ** and **QQ** combinations, as well as the diagonal cells, but only the defaults are used here. The bottom row, corresponding to `Age` uses boxplots to show the distributions of age for each of the categorical variables. The last column shows these same variables as stripcharts (or “barcodes”), which show all the individual observations. In the (1,4) and (4,1) panels, it can be seen that younger patients are more likely to report no improvement. The other panels in the first row (and column) show that improvement is more likely in the treated condition and greater among women than men. △

5.6 3D mosaics

{sec:3D}

Mosaic-like displays use the idea of recursive partitioning of a unit square to portray the frequencies in an n -way table by the area of rectangular tiles with (x, y) coordinates. The same idea extends

naturally to a 3D graphic. This starts with a unit cube, which is successively subdivided into 3D cuboids along (x, y, z) dimensions, and the frequency in a table cell is then represented by volume.

As in the 2D versions, each cuboid can be shaded to represent some other feature of the data, typically the residual from some model of independence. In principle, the display can accommodate more than 3 variables by using a sequence of split directions along the (x, y, z) axes.

One difficulty in implementing this method is that, short of using a 3D printer, the canvas for a 3D plot on a screen or printer is still projected on a two-dimensional surface, and graphical elements (volumes, lines, text) toward the front of the view will obscure those in the back. In R, a major advance in 3D graphics is available in the `rgl` package, that mitigates these problems by: (a) providing an interactive graphic window that can be zoomed and rotated manually with the mouse; (b) allowing dynamic graphics under program control, for example to animate a plot or make a movie; (c) providing control of the details of 3D rendering, including transparency of shapes, surface shading, lighting and perspective.

The `vcdExtra` package implements 3D mosaics using `rgl` graphics. `mosaic3d()` provides methods for "loglm" as well as "table" (or "structable") objects. At the time of writing, only some features of 2D mosaics are available.

{ex:bartlett-3d}

EXAMPLE 5.16: Bartlett data on plum root cuttings

In Example 5.11 we showed the mosaic matrix for the *Bartlett*, fitting the model of mutual independence to show all associations among the table variables, *Alive*, *Time* of planting and *Length* of cutting. Figure 5.26 shows the 3D version, produced using `mosaic3d()`:

```
> mosaic3d(Bartlett)
```

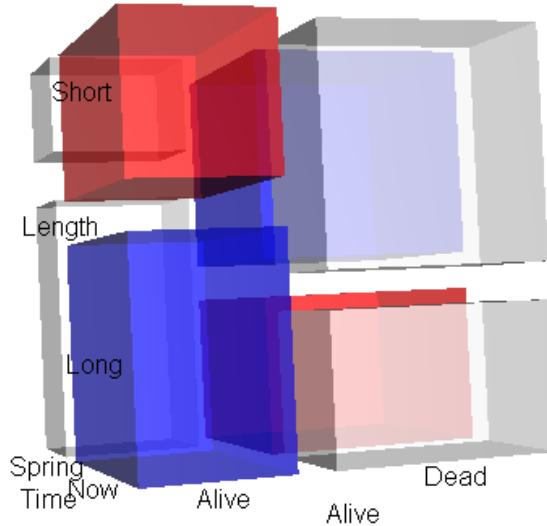


Figure 5.26: 3D mosaic plot of the Bartlett data, according to the model of mutual independence

{fig:mos3d-bartlett}

In the view of this figure, it can be seen that cuttings are more likely to be alive when planted Now and when cut Long. These relations can more easily be appreciated by rotating the 3D display.



Done: Deleted section on parallel coordinate plots. Need to fixup Example 1.7.

5.7 Visualizing the structure of loglinear models

For quantitative response data, it is easy to visualize a fitted model—for linear regression, this is just a plot of the fitted line; for multiple regression or non-linear regression with two predictors, this is a plot of the fitted response surface. For a categorical response variable, an analog of such plots is provided by effect plots, described later in this book.

{sec:mosaic-struc}

For contingency table data, mosaic displays can be used in a similar manner to illuminate the relations among variables in a contingency table represented in various loglinear models, a point described by Theus and Lauer (1999). In fact, each of the model types depicted in Table 5.2 has a characteristic shape and structure in a mosaic display. This, in turn, leads to a clearer understanding of the structure which appears in real data when a given model fits, the relations among the models, and the use of mosaic displays. The essential idea is a simple extension of what we do for more traditional models: show the *expected* (fitted) frequencies under a given model rather than observed frequencies in a mosaic-like display.

To illustrate, we use some artificial data on the relations among age, sex and symptoms of some disease shown in the $2 \times 2 \times 2$ table `struc` below.

```
> struc <- array(c(6, 10, 312, 44,
+                   37, 31, 192, 76),
+                   dim = c(2, 2, 2),
+                   dimnames = list(Age=c("Young", "Old"),
+                                   Sex=c("F", "M"),
+                                   Disease=c("No", "Yes")))
+ )
> struc <- as.table(struc)
> structable(Sex+Age ~ Disease, struc)

      Sex          F          M
      Age Young Old Young Old
Disease
No            6   10   312   44
Yes           37   31   192   76
```

First, note that there are substantial associations in this table, as shown in Figure 5.27, fitting the (default) mutual independence model.

```
> mosaic(struc, shade=TRUE)
```

The first split by Age shows strong partial associations between Sex and Disease for both young and old. However the residuals have an opposite pattern for young and old, suggesting a more complex relationship among these variables.

In this section we are asking a different question: what would mosaic displays look like if the data were in accord with simpler models? One way to do this is simply to use the expected frequencies to construct the tiles, as in sieve diagrams. The result, in Figure 5.28, shows that the tiles for sex and disease align for each of the age groups, but it is harder to see the relations among all three variables in this plot.

```
> mosaic(struc, type="expected")
```

We can visualize the model-implied relations among all variables together more easily using mosaic matrices.

5.7.1 Mutual independence

For example, to show the structure of a table which exactly fits the model of mutual independence, H_1 , use the `loglm()` to find the fitted values, `fit`, as shown below. The function `fitted()` extracts these from the "`loglm`" object.

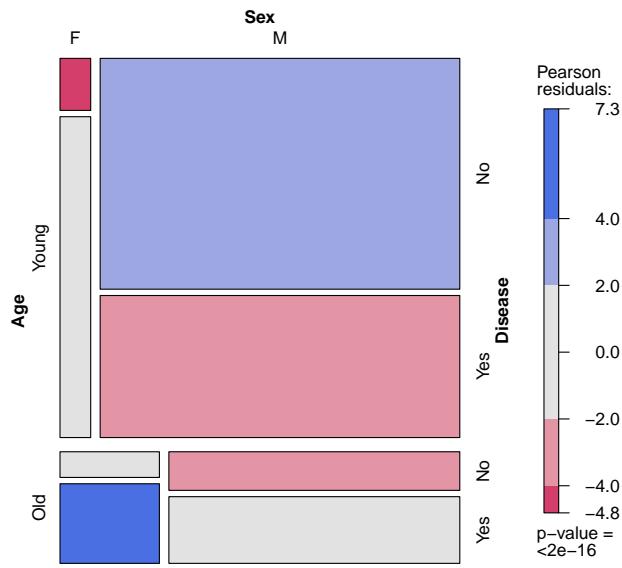


Figure 5.27: Mosaic display for the data on age, sex and disease. Observed frequencies are shown in the plot, and residuals reflect departure from the model of mutual independence.
fig:struct-mos1

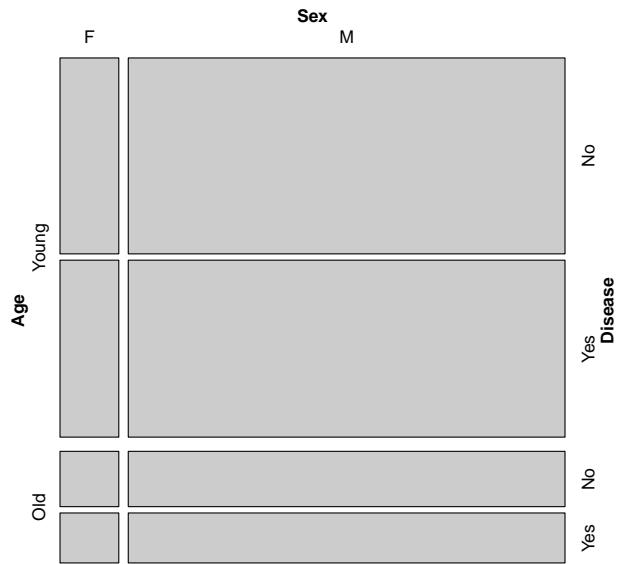


Figure 5.28: Mosaic display for the data on age, sex and disease, using expected frequencies under mutual independence. Left:
fig:struct-mos2

```
> mutual <- loglm(~Age+Sex+Disease, data=struk, fitted=TRUE)
> fit <- as.table(fitted(mutual))
> structable(Sex+Age ~ Disease, fit)
```

	Sex	F	M		
	Age	Young	Old	Young	Old
Disease	No	34.0991	10.0365	253.3077	74.5567
	Yes	30.7992	9.0652	228.7940	67.3416

These fitted frequencies then have the same one-way margins as the data in *struk*, but have no two-way or higher associations. Then, *pairs()* for this table, using *type="total"* shows the three-way mosaic for each pair of variables, giving the result in Figure 5.28. We use *gp=shading_Friendly* to explicitly indicate the zero residuals in the display.

```
> pairs(fit, gp=shading_Friendly, type="total")
```

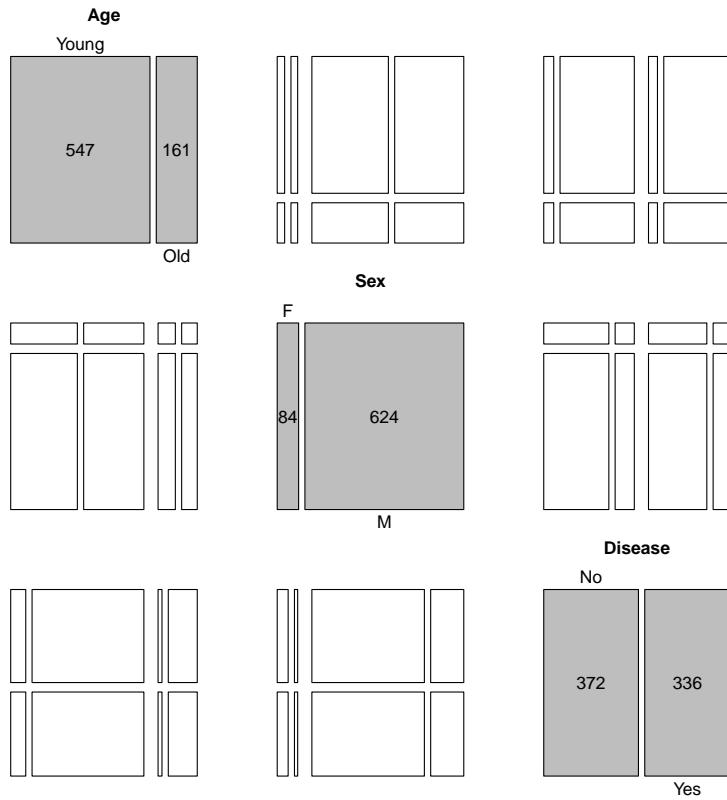


Figure 5.29: Mosaic matrix for fitted values under mutual independence. In all panels the joint frequencies conform to the one-way margins.^{fig:struk-moss}

In this figure the same data are shown in all the off-diagonal panels and the mutual independence model was fitted in each case, but with the table variables permuted. All residuals are exactly zero in all cells, by construction. We see that in each view, the four large tiles, corresponding to the first two variables align, indicating that these two variable are marginally independent. For example, in the (1,2) panel, age and sex are independent, collapsed over disease.

Moreover, comparing the top half to the bottom half in any panel we see that the divisions by the third variable are the same for both levels of the second variable. In the (1, 2) panel, for example, age and disease are independent for both males and females. This means that age and sex are conditionally independent given disease ($\text{age} \perp \text{sex} \mid \text{disease}$).

Because this holds in all six panels, we see that mutual independence implies that *all pairs* of variables are conditionally independent, given the remaining one, $(X \perp Y \mid Z)$ for all permutations of variables. A similar argument can be used to show that joint independence also holds, i.e., $((X, Y) \perp Z)$ for all permutations of variables.

Alternatively, you can also visualize these relationships interactively in a 3D mosaic using `mosaic3d()` that allows you to rotate the mosaic to see all views. In Figure 5.30, all of the 3D tiles are unshaded and you can see that the 3D unit cube has been sliced according to the marginal frequencies.

```
> mosaic3d(fit)
```

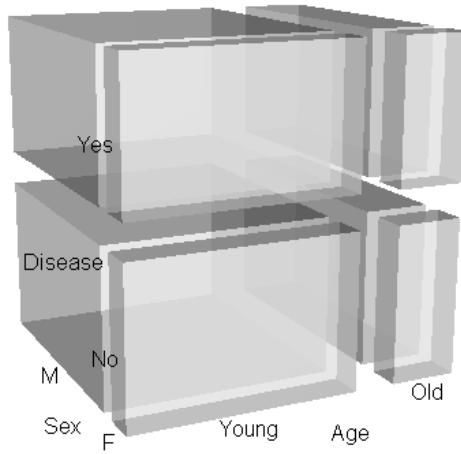


Figure 5.30: 3D mosaic plot of frequencies according to the model of mutual independence

{fig:struct-mos3d1}

5.7.2 Joint independence

The model of joint independence, $H_2 : (A, B) \perp C$, or equivalently, the loglinear model $[AB][C]$ may be visualized similarly by a mosaic matrix in which the data are replaced by fitted values under this model. We illustrate this for the model $[\text{Age Sex}][\text{Disease}]$, calculating the fitted values in a similar way as before.

```
> joint <- loglm(~Age*Sex + Disease, data=struc, fitted=TRUE)
> fit <- as.table(fitted(joint))
> structable(Sex+Age ~ Disease, fit)

      Sex          F          M
      Age   Young   Old   Young   Old
Disease
No        22.593 21.542 264.814 63.051
Yes       20.407 19.458 239.186 56.949
```

The `pairs.table()` plot, now using simpler pairwise plots (`type="pairwise"`), is shown in Figure 5.31.

```
> pairs(fit, gp=shading_Friendly)
```

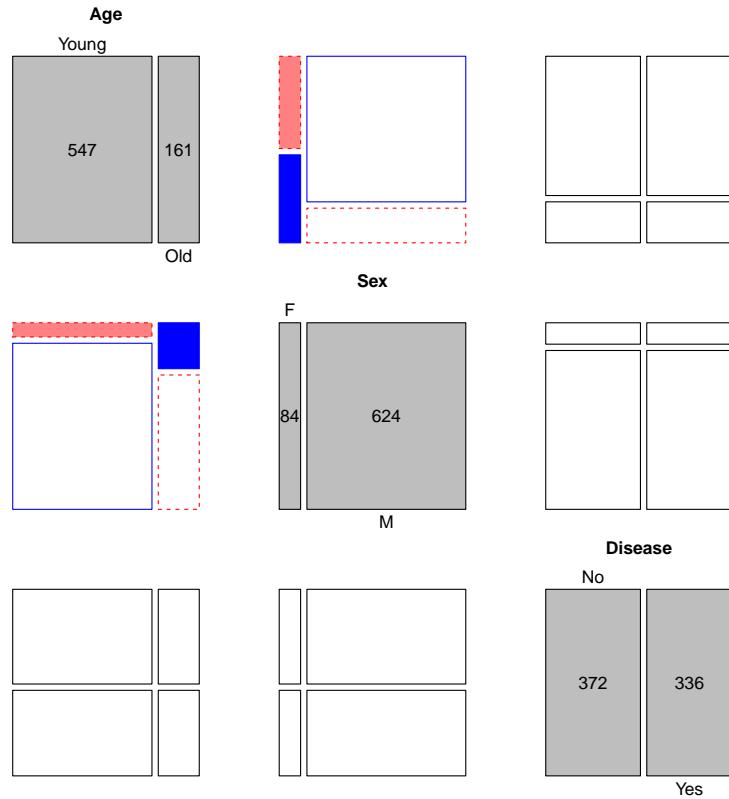


Figure 5.31: Mosaic matrix for fitted values under joint independence.
fig:struc-mos4

This shows, in row 3 and column 3, the anticipated independence of both age and sex with disease, collapsing over the remaining variable. The (1,2) and (2,1) panels show that age and sex are still associated when disease is ignored.

5.8 Chapter summary

- The mosaic display depicts the frequencies in a contingency table by a collection of rectangular “tiles” whose area is proportional to the cell frequency. The residual from a specified model is portrayed by shading the tile to show the sign and magnitude of the deviation from the model.
- For two-way tables, the tiles for the second variable align at each level of the first variable when the two variables are independent (see Figure 5.10).
- The perception and understanding of *patterns of association* (deviations from independence) are enhanced by reordering the rows or columns to give the shading of the residuals a more coherent pattern. An opposite-corner pattern “explains” the association in terms of the ordering of the factor levels.

- For three-way and larger tables, a variety of models can be fit and visualized. Starting with a minimal baseline model, the pattern of residuals will often suggest additional terms which must be added to “clean the mosaic.”
- It is often useful to examine the *sequential* mosaic displays for the marginal subtables with the variables in a given order. Sequential models of joint independence provide a breakdown of the total association in the full table, and are particularly appropriate when the last variable is a response.
- Partial association, which refers to the associations among a subset of variables, within the levels of other variables, may be easily studied by constructing separate mosaics for the subset variables for the levels of the other, “given” variables. These displays provide a breakdown of a model of conditional association for the whole table, and serve as an analog of coplots for quantitative data.
- Mosaic matrices, consisting of all pairwise plots of an n -way table, provide a way to visualize all marginal, joint, or conditional relations simultaneously. Parallel set plots provide another method to visualize n -way tables.
- The structural relations among model terms in various loglinear models themselves can also be visualized by mosaic matrices showing the expected, rather than observed, frequencies under different models.

5.9 Further reading

5.10 Lab exercises

Exercise 5.1 The data set *criminal* in the package *logmult* gives the 4×5 table below of the number of men aged 15–19 charged with a criminal case for whom charges were dropped in Denmark from 1955–1958.

```
> data("criminal", package="logmult")
Error in find.package(package, lib.loc, verbose = verbose): there is no package
called 'logmult'
> criminal
Error in eval(expr, envir, enclos): object 'criminal' not found
```

- (a) Use *loglm()* to test whether there is an association between Year and Age. Is there evidence that dropping of charges in relation to age changed over the years recorded here?
- (b) Use *mosaic()* with the option *shade=TRUE* to display the pattern of signs and magnitudes of the residuals. Compare this with the result of *mosaic()* using “Friendly shading,” from the option *gp=shading_Friendly*. Describe verbally what you see in each regarding the pattern of association in this table.

Exercise 5.2 The *Lahman* package contains comprehensive data on baseball statistics for Major League Baseball from 1871 through 2012. For all players, the *Master* table records the handedness of players, in terms of throwing (L, R) and batting (B, L, R), where B indicates “both.” The table below was generated using the following code:

{sec:mosaic-reading}

{lab:5.1}

{lab:5.2}

```
> library(Lahman)
> data("Master", package="Lahman")
> basehands <- with(Master, table(throws, bats))
```

Throws	Bats		
	B	L	R
L	177	2640	527
R	924	1962	10442

- Use the code above, or else enter these data into a frequency table in R.
- Construct mosaic displays showing the relation of batting and throwing handedness, split first by batting and by throwing.
- From these displays, what can be said about players who throw with their left or right hands in terms of their batting handedness?

{lab:5.3}

Exercise 5.3 ★ A related analysis concerns differences in throwing handedness among baseball players according to the fielding position they play. The following code calculates a such a frequency table.

```
> library(Lahman)

Error in library(Lahman): there is no package called 'Lahman'

> MasterFielding <- data.frame(merge(Master, Fielding, by="playerID"))

Error in merge(Master, Fielding, by = "playerID"): object 'Master' not found

> throwPOS <- with(MasterFielding, table(POS, throws))

Error in with(MasterFielding, table(POS, throws)): object 'MasterFielding' not found
```

- Make a mosaic display of throwing hand vs. fielding position.
- Calculate the percentage of players throwing left-handed by position. Make a sensible graph of this data.
- Re-do the mosaic display with the positions sorted by percentage of left-handers.
- Is there anything you can say about positions that have very few left-handed players?

{lab:5.4}

Exercise 5.4 For the *Bartlett* data described in Example 5.11, fit the model of no three-way association, H_4 in Table 5.2.

- Summarize the goodness of fit for this model, and compare to simpler models that omit one or more of the two-way terms.
- Use a mosaic-like display to show the lack of fit for this model.

{lab:5.5}

Exercise 5.5 Red core disease, caused by a fungus, is not something you want if you are a strawberry. The data set *jansen.strawberry* from the *agridat* package gives a frequency data frame of counts of damage from this fungus from a field experiment reported by Jansen (1990). See the help file for details. The following lines create a a $3 \times 4 \times 3$ table of crossings of 3 male parents with 4 (different) female parents, recording the number of plants in four blocks of 9 or 10 plants each showing red core disease in three ordered categories, C1, C2 or C3.

```
> data("jansen.strawberry", package="agridat")
>
> dat <- jansen.strawberry
> dat <- transform(dat, category=ordered(category, levels=c('C1', 'C2', 'C3')))
> levels(dat$male) <- paste0("M", 1:3)
> levels(dat$female) <- paste0("F", 1:4)
>
> jansen.tab <- xtabs(count~male + female + category, data=dat)
> names(dimnames(jansen.tab)) <- c("Male parent", "Female parent",
+                                     "Disease category")
> ftable(jansen.tab)
```

- (a) Use pairs(jansen.tab, shade=TRUE) to display the pairwise associations among the three variables. Describe how disease category appears to vary with male and female parent? Why is there no apparent association between male and female parent?
- (b) As illustrated in Figure 5.6, use mosaic to prepare a 3-way mosaic plot with the tiles colored in increasing shades of some color according to disease category. Describe the pattern of category C3 in relation to male and female parent. (Hint: the highlighting arguments are useful here.)
- (c) With category as the response variable, the minimal model for association is $[MF][C]$, or $\sim 1*2 + 3$. Fit this model using loglm() and display the residuals from this model with mosaic(). Describe the pattern of lack of fit of this model.

{lab:5.6}

Exercise 5.6 The data set *caith* in MASS gives another classic 4×5 table tabulating hair color and eye color, this for people in Caithness, Scotland, originally from Fisher (1940). The data is stored as a data frame of cell frequencies, whose rows are eye colors and whose columns are hair colors.

```
> data("caith", package="MASS")
> caith

      fair red medium dark black
blue    326  38    241   110     3
light   688 116    584   188     4
medium  343  84    909   412    26
dark    98   48    403   681    85
```

- (a) The loglm() and mosaic() functions don't understand data in this format, so use Caith <- as.matrix(caith) to convert to array form. Examine the result, and use names(dimnames(Caith)) <- c() to assign appropriate names to the row and column dimensions.
- (b) Fit the model of independence to the resulting matrix using loglm().
- (c) Calculate and display the residuals for this model.
- (d) Create a mosaic display for this data.

{lab:5.7}

Exercise 5.7 The *HairEyePlace* data in vcdExtra, gives similar data on hair color and eye color, for both Caithness and Aberdeen as a $4 \times 5 \times 2$ table.

- (a) Prepare separate mosaic displays, one for each of Caithness and Aberdeen. Comment on any difference in the pattern of residuals.
- (b) Construct condition mosaic plots, using the formula $\sim \text{Hair} + \text{Eye} | \text{Place}$ and both mosaic() and cobabplot(). It is probably more useful here to suppress the legend in these plots. Comment on the difference in what is shown in the two displays.

{lab:mosaic-accident}

Exercise 5.8 Bertin (1983, p. 30–31) used a 4-way table of frequencies of traffic accident victims

in France in 1958 to illustrate his scheme for classifying data sets by numerous variables, each of which could have various types and could be assigned to various visual attributes. His data are contained in *Accident* in *vcdExtra*, a frequency data frame representing his $5 \times 2 \times 4 \times 2$ table of the variables age, result (died or injured), mode of transportation and gender.

```
> data("Accident", package="vcdExtra")
> str(Accident)

'data.frame': 80 obs. of 5 variables:
 $ age   : Ord.factor w/ 5 levels "0-9"<"10-19"<...: 5 5 5 5 5 5 5 5 5 ...
 $ result: Factor w/ 2 levels "Died","Injured": 1 1 1 1 1 1 1 2 2 ...
 $ mode   : Factor w/ 4 levels "4-Wheeled","Bicycle",...: 4 4 2 2 3 3 1 1 4 4 ...
 $ gender : Factor w/ 2 levels "Female","Male": 2 1 2 1 2 1 2 1 2 1 ...
 $ Freq   : int 704 378 396 56 742 78 513 253 5206 5449 ...
```

- (a) Use `loglm()` to fit the model of mutual independence, $\text{Freq} \sim \text{age} + \text{mode} + \text{gender} + \text{result}$ to this data.
- (b) Use `mosaic()` to produce an interpretable mosaic plot of the associations among all variables under the model of mutual independence. Try different orders of the variables in the mosaic. (*Hint:* the abbreviate component of the `labeling_args` argument to `mosaic()` will be useful to avoid some overlap of the category labels.)
- (c) Treat `result` ("Died" vs. "Injured") as the response variable, and fit the model $\text{Freq} \sim \text{age} * \text{mode} * \text{gender} + \text{result}$ that asserts independence of `result` from all others jointly.
- (d) Construct a mosaic display for the residual associations in this model. Which combinations of the predictor factors are more likely to result in death?

```
> #detach(package:ggtern)
> .locals$ch05 <- setdiff(ls(), .globals)
> #.locals$ch05
> remove(list=.locals$ch05[sapply(.locals$ch05, function(n) {!is.function(get(n))})])
```

Chapter 6

Correspondence analysis

Correspondence analysis provides visualizations of associations in a two-way contingency table in a small number of dimensions. Multiple correspondence analysis extends this technique to n -way tables. Other graphical methods, including mosaic matrices and biplots provide complementary views of loglinear models for two-way and n -way contingency tables.

{ch:corresp}

6.1 Introduction

Whenever a large sample of chaotic elements is taken in hand and marshalled in the order of their magnitude, an unsuspected and most beautiful form of regularity proves to have been latent all along.

Sir Francis Galton (1822–1911)

Correspondence analysis (CA) is an exploratory technique which displays the row and column categories in a two-way contingency table as points in a graph, so that the positions of the points represent the associations in the table. Mathematically, correspondence analysis is related to the *biplot*, to *canonical correlation*, and to *principal component analysis*.

This technique finds scores for the row and column categories on a small number of dimensions which account for the greatest proportion of the χ^2 for association between the row and column categories, just as principal components account for maximum variance of quantitative variables. But CA does more—the scores provide a quantification of the categories, and have the property that they maximize the correlation between the row and column variables. For graphical display two or three dimensions are typically used to give a reduced rank approximation to the data.

Correspondence analysis has a very large, multi-national literature and was rediscovered several times in different fields and different countries. The method, in slightly different forms, is also discussed under the names *dual scaling*, *optimal scaling*, *reciprocal averaging*, *homogeneity analysis*, and *canonical analysis of categorical data*.

See Greenacre (1984) and Greenacre (2007) for an accessible introduction to CA methodology, or Gifi (1981), Lebart *et al.* (1984) for a detailed treatment of the method and its applications from the Dutch and French perspectives. Greenacre and Hastie (1987) provide an excellent discussion of the geometric interpretation, while van der Heijden and de Leeuw (1985) and van der Heijden *et al.* (1989) develop some of the relations between correspondence analysis and log-linear methods for three-way and larger tables. Correspondence analysis is usually carried out in an exploratory,

graphical way. Goodman (1981, 1985, 1986) has developed related inferential models, the *RC* model and the canonical correlation model, with close links to CA.

One simple development of CA is as follows: For a two-way table the scores for the row categories, namely $\mathbf{X} = \{x_{im}\}$, and column categories, $\mathbf{Y} = \{y_{jm}\}$, on dimension $m = 1, \dots, M$ are derived from a (generalized) **singular value decomposition** of (Pearson) residuals from independence, expressed as d_{ij}/\sqrt{n} , to account for the largest proportion of the χ^2 in a small number of dimensions. This decomposition may be expressed as

$$\frac{d_{ij}}{\sqrt{n}} = \frac{n_{ij} - m_{ij}}{\sqrt{n m_{ij}}} = \mathbf{X} \mathbf{D}_\lambda \mathbf{Y}^\top = \sum_{m=1}^M \lambda_m x_{im} y_{jm}, \quad (6.1)$$

where m_{ij} is the expected frequency and where \mathbf{D}_λ is a diagonal matrix with elements $\lambda_1 \geq \lambda_2 \geq \dots \geq \lambda_M$, and $M = \min(I - 1, J - 1)$. In M dimensions, the decomposition Eqn. (6.1) is exact. For example, an $I \times 3$ table can be depicted exactly in two dimensions when $I \geq 3$. The useful result for visualization purposes is that a rank- d approximation in d dimensions is obtained from the first d terms on the right side of Eqn. (6.1). The proportion of the Pearson χ^2 accounted for by this approximation is

$$n \sum_m^d \lambda_m^2 / \chi^2.$$

The quantity $\chi^2/n = \sum_i \sum_j d_{ij}^2/n$ is called the total **inertia** and is identical to the measure of association known as Pearson's mean-square contingency, the square of the ϕ coefficient.

Thus, correspondence analysis is designed to show how the data deviate from expectation when the row and column variables are independent, as in the sieve diagram, association plot and mosaic display. However, the sieve, association and mosaic plots depict every *cell* in the table, and for large tables it may be difficult to see patterns. Correspondence analysis shows only row and column *categories* as points in the two (or three) dimensions which account for the greatest proportion of deviation from independence. The pattern of the associations can then be inferred from the positions of the row and column points.

6.2 Simple correspondence analysis

6.2.1 Notation and terminology

Because Correspondence analysis grew up in so many homes, the notation, formulae and terms used to describe the method vary considerably. The notation used here generally follows Greenacre (1984, 1997, 2007).

The descriptions here employ the following matrix and vector definitions:

- $\mathbf{N} = \{n_{ij}\}$ is the $I \times J$ contingency table with row and column totals n_{i+} and n_{+j} , respectively. The grand total n_{++} is also denoted by n for simplicity.
- $\mathbf{P} = \{p_{ij}\} = \mathbf{N}/n$ is the matrix of joint cell proportions, called the **correspondence matrix**.
- $\mathbf{r} = \sum_j p_{ij} = \mathbf{P}\mathbf{1}$ is the row margin of \mathbf{P} ; $\mathbf{c} = \sum_i p_{ij} = \mathbf{P}^\top \mathbf{1}$ is the column margin. \mathbf{r} and \mathbf{c} are called the *row masses* and *column masses*.
- \mathbf{D}_r and \mathbf{D}_c are diagonal matrices with \mathbf{r} and \mathbf{c} on their diagonals, used as weights.
- $\mathbf{R} = \mathbf{D}_r^{-1} \mathbf{P} = \{n_{ij}/n_{+j}\}$ is the matrix of row conditional probabilities, called *row profiles*. Similarly, $\mathbf{C} = \mathbf{D}_c^{-1} \mathbf{P}^\top = \{n_{ij}/n_{i+}\}$ is the matrix of column conditional probabilities or *column profiles*.

Two types of coordinates, \mathbf{X} , \mathbf{Y} for the row and column categories are defined, based on the generalized singular value decomposition of \mathbf{P} ,

$$\mathbf{P} = \mathbf{A}\mathbf{D}_\lambda\mathbf{B}^\top$$

where \mathbf{D}_λ is the diagonal matrix of singular values $\lambda_1 \geq \lambda_2 \geq \dots \geq \lambda_M$, \mathbf{A} is the $I \times M$ matrix of left singular vectors, normalized so that $\mathbf{A}\mathbf{D}_r^{-1}\mathbf{A}^\top = \mathbf{I}$, and \mathbf{B} is the $J \times M$ matrix of right singular vectors, normalized so that $\mathbf{B}\mathbf{D}_c^{-1}\mathbf{B}^\top = \mathbf{I}$. Thus the columns of \mathbf{A} and \mathbf{B} are orthogonal in the weighted metrics defined by the row and column margins, \mathbf{D}_r^{-1} and \mathbf{D}_c^{-1} , respectively.

principal coordinates: The coordinates of the row (\mathbf{F}) and column (\mathbf{G}) profiles with respect to their own principal axes are defined so that the inertia along each axis is the corresponding singular value, λ_i ,

$$\mathbf{F} = \mathbf{D}_r^{-1}\mathbf{A}\mathbf{D}_\lambda \quad \text{scaled so that} \quad \mathbf{F}^\top\mathbf{D}_r\mathbf{F} = \mathbf{D}_\lambda \quad (6.2) \quad \{\text{eq:pcoord1}\}$$

$$\mathbf{G} = \mathbf{D}_c^{-1}\mathbf{B}\mathbf{D}_\lambda \quad \text{scaled so that} \quad \mathbf{G}^\top\mathbf{D}_c\mathbf{G} = \mathbf{D}_\lambda \quad (6.3) \quad \{\text{eq:pcoord2}\}$$

The joint plot in principal coordinates, \mathbf{F} and \mathbf{G} , is called the **symmetric map** because both row and column profiles are overlaid in the same coordinate system.

standard coordinates: The standard coordinates (Φ , Γ) are a rescaling of the principal coordinates to unit inertia along each axis,

$$\Phi = \mathbf{D}_r^{-1}\mathbf{A} \quad \text{scaled so that} \quad \Phi^\top\mathbf{D}_r\Phi = \mathbf{I} \quad (6.4) \quad \{\text{eq:scoord1}\}$$

$$\Gamma = \mathbf{D}_c^{-1}\mathbf{B} \quad \text{scaled so that} \quad \Gamma^\top\mathbf{D}_c\Gamma = \mathbf{I} \quad (6.5) \quad \{\text{eq:scoord2}\}$$

These differ from the principal coordinates in Eqn. (6.2) and Eqn. (6.3) simply by the absence of the scaling factors, \mathbf{D}_λ . An **asymmetric map** shows one set of points (say, the rows) in principal coordinates and the other set in standard coordinates.

Thus, the weighted average of the squared principal coordinates for the rows or columns on a principal axis equals the squared singular value, λ for that axis, whereas the weighted average of the squared standard coordinates equals 1. The relative positions of the row or column points along any axis is the same under either scaling, but the distances between points differ, because the axes are weighted differentially in the two scalings.

6.2.2 Geometric and statistical properties

We summarize here some geometric and statistical properties of the Correspondence analysis solutions which are useful in interpretation.

{sec:ca-properties}

nested solutions: Because they use successive terms of the SVD Eqn. (6.1), correspondence analysis solutions are *nested*, meaning that the first two dimensions of a three-dimensional solution will be identical to the two-dimensional solution.

centroids at the origin: In both principal coordinates and standard coordinates the points representing the row and column profiles have their centroids (weighted averages) at the origin. Thus, in CA plots, the origin represents the (weighted) average row profile and column profile.

reciprocal averages: CA assigns scores to the row and column categories such that the column scores are proportional to the weighted averages of the row scores, and vice-versa.

chi-square distances: In principal coordinates, the row coordinates may be shown equal to the row profiles $D_r^{-1}P$, rescaled inversely by the square-root of the column masses, $D_c^{-1/2}$. Distances between two row profiles, R_i and $R_{i'}$ are most sensibly defined as χ^2 distances, where the squared difference $[R_{ij} - R_{i'j}]^2$ is inversely weighted by the column frequency, to account for the different relative frequency of the column categories. The rescaling by $D_c^{-1/2}$ transforms this weighted χ^2 metric into ordinary Euclidean distance. The same is true of the column principal coordinates.

interpretation of distances: In principal coordinates, the distance between two row points may be interpreted as described above, and so may the distance between two column points. The distance between a row and column point, however, does not have a clear distance interpretation.

residuals from independence: The distance between a row and column point do have a rough interpretation in terms of residuals or the difference between observed and expected frequencies, $n_{ij} - m_{ij}$. Two row (or column) points deviate from the origin (the average profile) when their profile frequencies have similar values. A row point appears in a similar direction away from the origin as a column point when $n_{ij} - m_{ij} > 0$, and in an opposite different direction from that column point when the residual is negative.

Because of these differences in interpretations of distances, there are different possibilities for graphical display. A joint display of principal coordinates for the rows and standard coordinates for the columns (or vice-versa), sometimes called an *asymmetric map* is suggested by Greenacre and Hastie (1987) and by Greenacre (1989) as the plot with the most coherent geometric interpretation (for the points in principal coordinates) and is sometimes used in the French literature.

Another common joint display is the *symmetric map* of the principal coordinates in the same plot. This is the default in the `ca` package described below. In the authors' opinion, this produces better graphical displays, because both sets of coordinates are scaled with the same weights for each axis. Symmetric plots are used exclusively in this book, but that should not imply that these plots are universally preferred. Another popular choice is to avoid the possibility of misinterpretation by making separate plots of the row and column coordinates.

6.2.3 R software for correspondence analysis

{sec:ca-R}

Correspondence analysis methods for computation and plotting are available in a number of R packages including:

MASS: `corresp()`; the plot method calls `biplot()` for a 2 factor solution, using a symmetric biplot factorization that scales the row and column points by the square roots of the singular values. There is also a `mca()` function for multiple correspondence analysis.

ca: `ca()`; provides 2D plots via the `plot.ca()` method and interactive (`rgl`) 3D plots via `plot3d.ca()`. This package is the most comprehensive in terms of plotting options for various coordinate types, plotting supplementary points. It also provides `mjca()` for multiple and joint correspondence analysis of higher-way tables.

FactoMineR: `CA()`; provides a wide variety of measures for the quality of the CA representation and many options for graphical display

These methods also differ in terms of the types of input they accept. For example, `MASS::corresp` handles matrices, data frames and "xtabs" objects, but not "table" objects. `ca()` is the most general, with methods for two-way tables, matrices, data frames, and "xtabs" objects. In the following, we largely use the `ca` package.

EXAMPLE 6.1: Hair color and eye color

The script below uses the two-way table `haireye` from the `HairEyeColor` data, collapsed

{ex:haireye3}

over Sex. In this table, Hair colors form the rows, and Eye colors form the columns. By default, ca() produces a 2-dimensional solution. In this example, the complete, exact solution would have $M = \min((I - 1), (J - 1)) = 3$ dimensions, and you could obtain this using the argument nd=3 in the call to ca().

```
> haireye <- margin.table(HairEyeColor, 1:2)
> library(ca)
> (haireye.ca <- ca(haireye))

Principal inertias (eigenvalues):
      1          2          3
Value 0.208773 0.022227 0.002598
Percentage 89.37% 9.52% 1.11%

Rows:
      Black     Brown     Red   Blond
Mass    0.18243 0.48311 0.1199 0.2145
ChiDist 0.55119 0.15946 0.3548 0.8384
Inertia 0.05543 0.01228 0.0151 0.1508
Dim. 1 -1.10428 -0.32446 -0.2835 1.8282
Dim. 2  1.44092 -0.21911 -2.1440 0.4667

Columns:
      Brown     Blue     Hazel    Green
Mass    0.37162 0.3632 0.15710 0.10811
ChiDist 0.50049 0.5537 0.28865 0.38573
Inertia 0.09309 0.1113 0.01309 0.01608
Dim. 1 -1.07713 1.1981 -0.46529 0.35401
Dim. 2  0.59242 0.5564 -1.12278 -2.27412
```

In the printed output, the table labeled “Principal inertias (eigenvalues)” indicates that nearly 99% of the Pearson χ^2 for association is accounted for by two dimensions, with most of that attributed to the first dimension.

The summary method for “ca” objects gives a more nicely formatted display, showing a *scree plot* of the eigenvalues, a portion of which is shown below.

```
> summary(haireye.ca)

Principal inertias (eigenvalues):

  dim   value     %   cum%   scree plot
  1    0.208773 89.4 89.4 ****
  2    0.022227  9.5 98.9 **
  3    0.002598   1.1 100.0
  -----
  Total: 0.233598 100.0
  ...
```

The result returned by ca() can be plotted using the plot.ca() method. However, it is useful to understand that ca() returns the CA solution in terms of *standard coordinates*, Φ (rowcoord) and Γ (colcoord). We illustrate Eqn. (6.4) and Eqn. (6.5) using the components of the “ca” object haireye.ca.

```
> # standard coordinates Phi (Eqn 6.4) and Gamma (Eqn 6.5)
> (Phi <- haireye.ca$rowcoord)
```

```

      Dim1      Dim2      Dim3
Black -1.10428  1.44092 -1.08895
Brown -0.32446 -0.21911  0.95742
Red   -0.28347 -2.14401 -1.63122
Blond  1.82823  0.46671 -0.31809

> (Gamma <- haireye.ca$colcoord)

      Dim1      Dim2      Dim3
Brown -1.07713  0.59242 -0.423960
Blue   1.19806  0.55642  0.092387
Hazel -0.46529 -1.12278  1.971918
Green  0.35401 -2.27412 -1.718443

> # demonstrate orthogonality of std coordinates
> Dr <- diag(haireye.ca$rowmass)
> zapsmall(t(Phi) %*% Dr %*% Phi)

      Dim1 Dim2 Dim3
Dim1    1    0    0
Dim2    0    1    0
Dim3    0    0    1

> Dc <- diag(haireye.ca$colmass)
> zapsmall(t(Gamma) %*% Dc %*% Gamma)

      Dim1 Dim2 Dim3
Dim1    1    0    0
Dim2    0    1    0
Dim3    0    0    1

```

These standard coordinates are transformed internally within the plot function according to the map argument, which defaults to map="symmetric", giving principal coordinates. The following call to `plot.ca()` produces Figure 6.1.

```

> op <- par(cex=1.4, mar=c(5, 4, 1, 2)+.1)
> res <- plot(haireye.ca)
> par(op)

```

For use in further customizing such plots (as we will see in the next example), the function `plot.ca()` returns (invisibly)¹ the coordinates for the row and column points actually plotted, which we saved above as `res`:

```

> res

$rows
      Dim1      Dim2
Black -0.50456  0.214820
Brown -0.14825 -0.032666
Red   -0.12952 -0.319642
Blond  0.83535  0.069579

$cols
      Dim1      Dim2
Brown -0.49216  0.088322
Blue   0.54741  0.082954
Hazel -0.21260 -0.167391
Green  0.16175 -0.339040

```

It is important to understand that in CA plots (and related biplots, Section 6.5), the interpretation

¹This uses features incorporated in the `ca` package, version 0.54+.

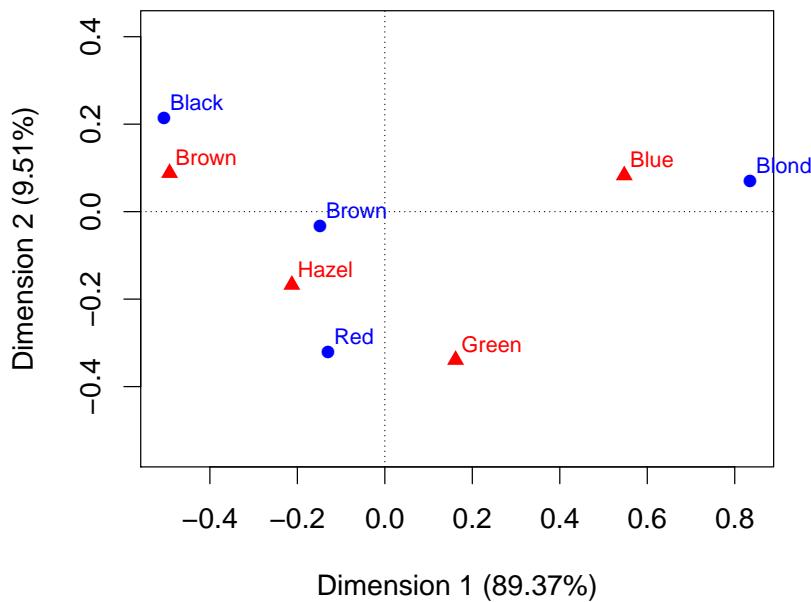


Figure 6.1: Correspondence analysis solution for the Hair color and Eye color data fig:ca-haireye-plot

of distances between points (and angles between vectors) is meaningful. In order to achieve this, the axes in such plots must be *equated*, meaning that the two axes are scaled so that the number of data units per inch are the same for both the horizontal and vertical axes, or an *aspect ratio* = 1.²

The interpretation of the CA plot in Figure 6.1 is then as follows:

- Dimension 1, accounting for nearly 90% of the association between hair and eye color corresponds to dark (left) vs. light (right) on both variables.
- Dimension 2 largely contrasts red hair and green eyes with the remaining categories, accounting for an additional 9.5% of the Pearson χ^2 .
- With equated axes, and a symmetric map, the distances between row points and column points are meaningful. Along Dimension 1, the eye colors could be considered roughly equally spaced, but for the hair colors, Blond is quite different in terms of its frequency profile.

△

{ex:mental3}

EXAMPLE 6.2: Mental impairment and parents' SES

In Example 4.3 we introduced the data set *Mental*, relating mental health status to parents' SES. As in Example 4.6, we convert this to a two-way table, *mental.tab* to conduct a correspondence analysis.

```
> data("Mental", package="vcdExtra")
> mental.tab <- xtabs(Freq ~ ses + mental, data=Mental)
```

We calculate the CA solution, and save the result in *mental.ca*:

²In base R graphics, this is achieved with the `plot()` option `asp=1`.

```
> mental.ca <- ca(mental.tab)
> summary(mental.ca)

Principal inertias (eigenvalues):

  dim   value      %   cum%   scree plot
  1    0.026025  93.9  93.9  *****
  2    0.001379   5.0  98.9   *
  3    0.000298   1.1 100.0
  -----
Total: 0.027702 100.0
...
```

The scree plot produced by `summary(mental.ca)` shows that the association between mental health and parents' SES is almost entirely 1-dimensional, with 94% of the χ^2 (45.98, with 15 df) accounted for by Dimension 1.

We then plot the solution as shown below, giving Figure 6.2. For this example, it is useful to connect the row points and the column points by lines, to emphasize the pattern of these ordered variables.

```
> op <- par(cex=1.3, mar=c(5,4,1,1)+.1)
> res <- plot(mental.ca, ylim=c(-.2, .2))
> lines(res$rows, col="blue", lty=3)
> lines(res$cols, col="red", lty=4)
> par(op)
```

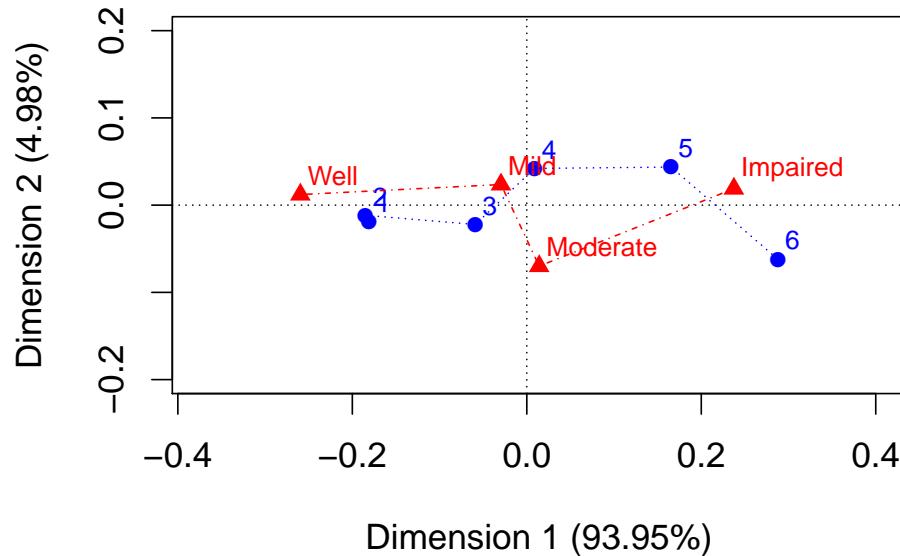


Figure 6.2: Correspondence analysis solution for the Mental health data fig:ca-mental-plot

The plot of the CA scores in Figure 6.2 shows that diagnostic mental health categories are well-aligned with Dimension 1. The mental health scores are approximately equally spaced, except that the two intermediate categories are a bit closer on this dimension than the extremes. The SES categories are also aligned with Dimension 1, and approximately equally spaced, with the exception

of the highest two SES categories, whose profiles are extremely similar, suggesting that these two categories could be collapsed.

Because both row and column categories have the same pattern on Dimension 1, we may interpret the plot as showing that the profiles of both variables are ordered, and their relation can be explained as a positive association between high parents' SES and higher mental health status of children. A mosaic display of these data (Exercise 6.5) would show a characteristic opposite corner pattern of association.

From a modeling perspective, we might ask how strong is the evidence for the spacing of categories noted above. For example, we might ask whether assigning integer scores to the levels of SES and mental impairment provides a simpler, but satisfactory account of their association. Questions of this type can be explored in connection with loglinear models in Chapter 8.



{ex:victims2}

EXAMPLE 6.3: Repeat victimization

The data set *RepVict* in the *vcd* package gives a 8×8 table (from Fienberg (1980, Table 2-8)) on repeat victimization for various crimes among respondents to a U.S. National Crime Survey. A special feature of this data set is that row and column categories reflect the *same* crimes, so substantial association is expected. Here we examine correspondence analysis results in a bit more detail and also illustrate how to customize the displays created by `plot(ca(...))`.

```
> data("RepVict", package="vcd")
> victim.ca <- ca(RepVict)
> summary(victim.ca)

Principal inertias (eigenvalues):

  dim   value     %   cum%   scree plot
  1    0.065456 33.8 33.8 *****
  2    0.059270 30.6 64.5 *****
  3    0.029592 15.3 79.8 *****
  4    0.016564  8.6 88.3 ****
  5    0.011140  5.8 94.1 ***
  6    0.007587  3.9 98.0 **
  7    0.003866  2.0 100.0
  -----
  Total: 0.193474 100.0
  ...
  
```

The results above show that, for this 8×8 table, 7 dimensions are required for an exact solution, of which the first two account for 64.5% of the Pearson χ^2 . The lines below illustrate that the Pearson χ^2 is n times the sum of the squared singular values, $n \sum \lambda_i^2$.

```
> chisq.test(RepVict)

Pearson's Chi-squared test

data: RepVict
X-squared = 11131, df = 49, p-value < 2.2e-16

> (chisq <- sum(RepVict) * sum(victim.ca$sv^2))

[1] 11131
```

The default plot produced by `plot.ca(victim.ca)` plots both points and labels for the row

and column categories. However, what we want to emphasize here is the relation between the *same* crimes on the first and second occurrence.

To do this, we label each crime just once (using `labels=c(2, 0)`) and connect the two points for each crime by a line, using `segments()`, as shown in Figure 6.3. The addition of a `legend()` makes the plot more easily readable.

```
> op <- par(cex=1.3, mar=c(4, 4, 1, 1)+.1)
> res <- plot(victim.ca, labels=c(2, 0))
> segments(res$rows[,1], res$rows[,2], res$cols[,1], res$cols[,2])
> legend("topleft", legend=c("First", "Second"), title="Occurrence",
+        col=c("blue", "red"), pch=16:17, bg="gray90")
> par(op)
```

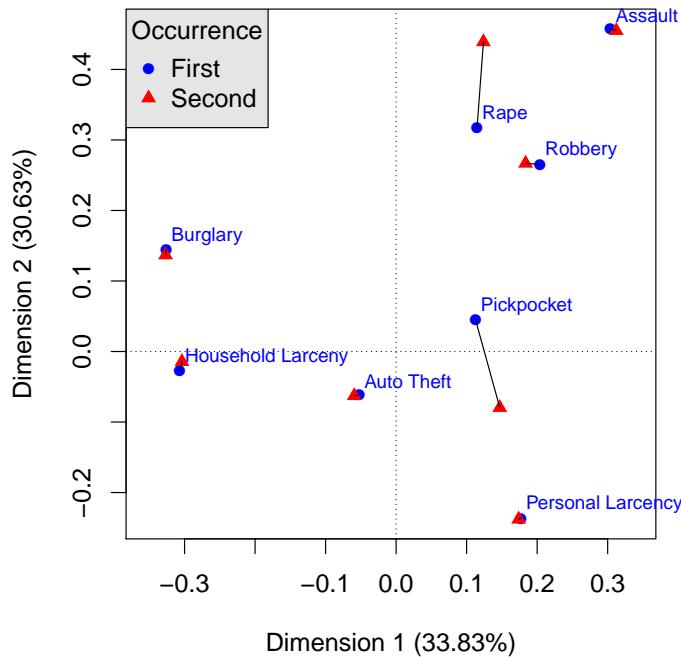


Figure 6.3: 2D CA solution for the repeat victimization data. Lines connect the category points for first and second occurrence to highlight these relations.
fig.ca-victims-plot

In Figure 6.3 it may be seen that most of the points are extremely close for the first and second occurrence of a crime, indicating that the row profile for a crime is very similar to its corresponding column profile, with Rape and Pick Pocket as exceptions.

In fact, if the table was symmetric, the row and column points in Figure 6.3 would be identical, as can be easily demonstrated by analyzing a symmetric version.

```
> RVsym <- (RepVict + t(RepVict))/2
> RVsym.ca <- ca(RVsym)
> res <- plot(RVsym.ca)
> all.equal(res$rows, res$cols)

[1] TRUE
```

The first dimension appears to contrast crimes against the person (right) with crimes against property (left), and it may be that the second dimension represents degree of violence associated

with each crime. The latter interpretation is consistent with the movement of Rape towards a higher position and Pickpocket towards a lower one on this dimension.



6.2.4 Correspondence analysis and mosaic displays

For a two-way table, CA and mosaic displays give complementary views of the pattern of association between the row and column variables, but both are based on the (Pearson) residuals from independence. CA shows the row and column categories as points in a 2D (or 3D) space accounting for the largest proportion of the Pearson χ^2 , while mosaics show the association by the pattern of shading in the mosaic tiles. It is useful to compare them directly to see how associations can be interpreted from these graphs.

{ex:TV2}

EXAMPLE 6.4: TV viewing data

The data on television viewership from Hartigan and Kleiner (1984) was used as an example of manipulating complex categorical data in Section 2.9 and shown as a three-way mosaic plot in Figure ???. From that figure, the main association concerned how viewership across days of the week varied by TV network, so we first collapse the *TV* data to a 5×3 two-way table.

```
> data("TV", package="vcdExtra")
> TV2 <- margin.table(TV, c(1, 3))
> TV2

      Network
Day      ABC   CBS   NBC
Monday    2847 2923 2629
Tuesday   3110 2403 2568
Wednesday 2434 1283 2212
Thursday  1766 1335 5886
Friday    2737 1479 1998
```

In this case, the 2D CA solution is exact, meaning that two dimensions account for 100% of the association.

```
> TV.ca <- ca(TV2)
> TV.ca

Principal inertias (eigenvalues):
      1       2
Value 0.081934 0.010513
Percentage 88.63% 11.37%
...
```

The plot of this solution is shown in the left panel of Figure 6.4, using lines from the origin to the category points for the networks.

```
> res <- plot(TV.ca)
> segments(0, 0, res$cols[,1], res$cols[,2], col="red", lwd=2)
```

An analogous mosaic display, informed by the CA solution, is shown in the right panel of Figure 6.4. Here, the days of the week are reordered according to their positions on the first CA dimension, another example of effect ordering.

```
> days.order <- order(TV.ca$rowcoord[,1])
> mosaic(t(TV2[days.order,]), shade=TRUE, legend=FALSE,
+         labeling=labeling_residuals, suppress=0)
```

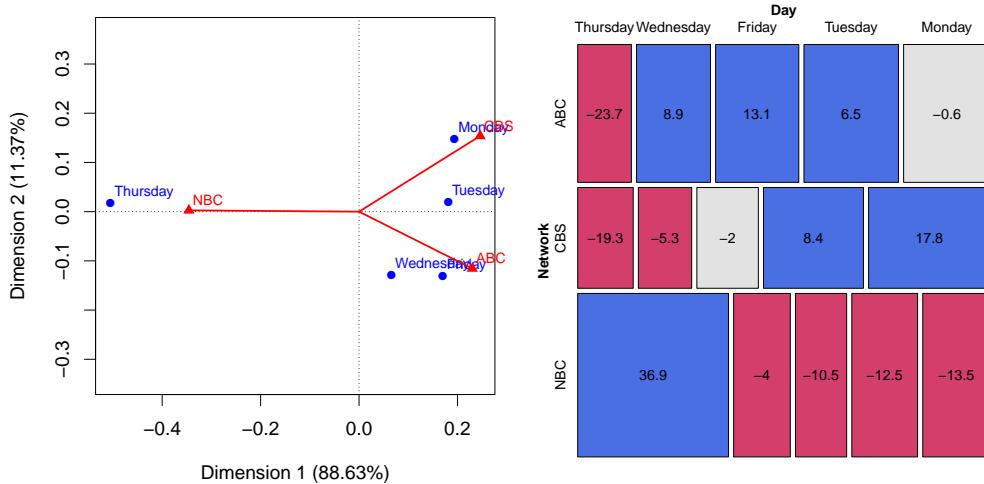


Figure 6.4: CA plot and mosaic display for the TV viewing data. The days of the week in the mosaic plot were permuted according to their order in the CA solution.

{fig:TV-mosaic-ca}

In the CA plot, you can see that the dominant dimension separates viewing on Thursday, with the largest share of viewers watching NBC, from the other weekdays. In the mosaic plot, Thursday stands out as the only day with a higher than expected frequency for NBC, and this is the largest residual in the entire table. The second dimension in the CA plot separates CBS, with its' greatest proportion of viewers on Monday, from ABC, with greater viewership on Wednesday and Friday.

? Fig. 2 gives a table listing the shows in each half-hour time slot. Could the overall popularity of NBC on Thursday be due to *Friends* or *Seinfeld*? An answer to this and similar questions requires analysis of the three-way table (Exercise 6.7) and model-based methods for polytomous outcome variables described in Section 7.6.4.

△

6.3 Multi-way tables: Stacking and other tricks

A three- or higher-way table can be analyzed by correspondence analysis in several ways. Multiple correspondence analysis (MCA), described in Section 6.4, is an extension of simple correspondence analysis which analyzes simultaneously all possible two-way tables contained within a multiway table. Another approach, described here, is called *stacking* or *interactive coding*. This is a bit of a trick, to force a multiway table into a two-way table for a standard correspondence analysis, but is a useful one.

A three-way table, of size $I \times J \times K$ can be sliced into I two-way tables, each $J \times K$. If the slices are concatenated vertically, the result is one two-way table, of size $(I \times J) \times K$, as illustrated in Figure 6.5. In effect, the first two variables are treated as a single composite variable with IJ levels, which represents the main effects and interaction between the original variables that were combined. Van der Heijden and de Leeuw (1985) discuss this use of correspondence analysis for multi-way tables and show how *each* way of slicing and stacking a contingency table corresponds to the analysis of a specified loglinear model. Like the mosaic display, this provides another way to visualize the relations in a loglinear model.

In particular, for the three-way table with variables A, B, C that is reshaped as a table of size $(I \times J) \times K$, the correspondence analysis solution analyzes residuals from the log-linear model

{sec:ca-multiway}

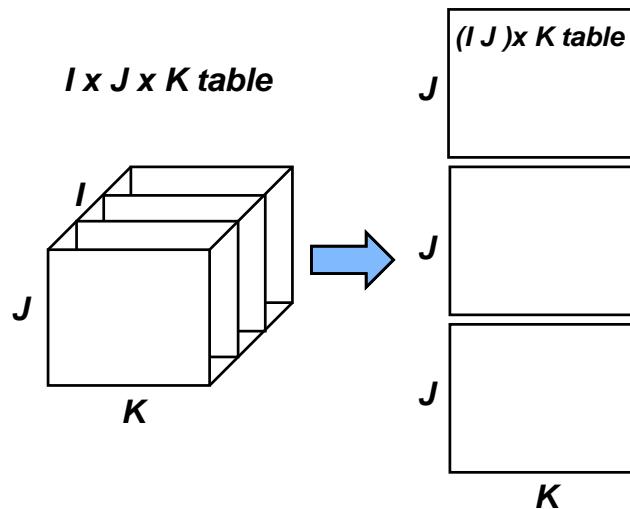


Figure 6.5: Stacking approach for a three-way table. Two of the table variables are combined interactively to form the rows of a two-way table.

{fig:stacking}

$[AB][C]$. That is, for such a table, the $I \times J$ rows represent the joint combinations of variables A and B. The expected frequencies under independence for this table are

$$m_{[ij]k} = \frac{n_{ij} + n_{++k}}{n} \quad (6.6) \quad \text{(eq:mij-k)}$$

which are the ML estimates of expected frequencies for the log-linear model $[AB][C]$. The χ^2 that is decomposed by correspondence analysis is the Pearson χ^2 for this log-linear model. When the table is stacked as $I \times (J \times K)$ or $J \times (I \times K)$, correspondence analysis decomposes the residuals from the log-linear models $[A][BC]$ and $[B][AC]$, respectively, as shown in Table 6.1. In this approach, only the associations in separate $[]$ terms are analysed and displayed in the correspondence analysis maps. Van der Heijden and de Leeuw (1985) show how a generalized form of correspondence analysis can be interpreted as decomposing the difference between two specific loglinear models, so their approach is more general than is illustrated here.

Table 6.1: Each way of stacking a three-way table corresponds to a loglinear model

{tab:stacking}

Stacking structure	Loglinear model
$(I \times J) \times K$	$[AB][C]$
$I \times (J \times K)$	$[A][BC]$
$J \times (I \times K)$	$[B][AC]$

6.3.1 Interactive coding in R

In the general case of an n -way table, the stacking approach is similar to that used by `ftable()` and `structable()` in `vcd` as described in Section 2.5 to flatten multiway tables to a two-way, printable form, where some variables are assigned to the rows and the others to the columns. Both `ftable()` and `structable()` have `as.matrix()` methods³ that convert their result into a matrix suitable as input to `ca()`.

{sec:ca-interactiveR}

³This requires at least R version 3.1.0 or vcd 1.3-2 or later.

With data in the form of a frequency data frame, you can easily create interactive coding using `interaction()` or simply use `paste()` to join the levels of stacked variables together.

To illustrate, create a 4-way table of random Poisson counts (with constant mean, $\lambda = 15$) of types of Pet, classified by Age, Color and Sex.

```
> set.seed(1234)
> dim <- c(3, 2, 2, 2)
> tab <- array(rpois(prod(dim), 15), dim=dim)
> dimnames(tab) <- list(Pet=c("dog", "cat", "bird"),
+                         Age=c("young", "old"),
+                         Color=c("black", "white"),
+                         Sex=c("male", "female"))
```

You can use `ftable()` to print this, with a formula that assigns Pet and Age to the columns and Color and Sex to the rows.

```
> ftable(Pet + Age ~ Color + Sex, tab)

      Pet    dog      cat      bird
      Age young old young old young old
Color Sex
black male       10   12     16   16     16   12
      female      8    12     13   15     11   13
white male       18   11     12   18     13   20
      female      13   13     16   15     12   15
```

Then, `as.matrix()` creates a matrix with the levels of the stacked variables combined with some separator character. Using `ca(pet.mat)` would then calculate the CA solution for the stacked table, analyzing only the associations in the loglinear model [PetAge][ColorSex].⁴

```
> (pet.mat <- as.matrix(ftable(Pet + Age ~ Color + Sex, tab), sep='.'))

      Pet.Age
Color.Sex      dog.young dog.old cat.young cat.old bird.young bird.old
      black.male    10        12      16        16      16        12
      black.female   8        12      13        15      11        13
      white.male    18        11      12        18      13        20
      white.female   13        13      16        15      12        15
```

With data in a frequency data frame, a similar result (as a frequency table), can be obtained using `interaction()` as shown below. The result of `xtabs()` looks the same as `pet.mat`.

```
> tab.df <- as.data.frame(as.table(tab))
> tab.df <- within(tab.df,
+   {Pet.Age = interaction(Pet, Age)
+   Color.Sex = interaction(Color, Sex)
+ })
> xtabs(Freq ~ Color.Sex + Pet.Age, data=tab.df)
{ex:suicide1}
```

EXAMPLE 6.5: Suicide rates in Germany

To illustrate the use of correspondence analysis for the analysis for three-way tables, we use data on suicide rates in West Germany classified by sex, age, and method of suicide used. The data, from Heuer (1979, Table 1) have been discussed by Friendly (1991, 1994b), van der Heijden and de Leeuw (1985) and others.

The original $2 \times 17 \times 9$ table contains 17 age groups from 10 to 90 in 5-year steps and 9 categories of suicide method, contained in the frequency data frame `Suicide` in `vcd`, with table variables `sex`, `age` and `method`. To avoid extremely small cell counts and cluttered displays, this

⁴The result would not be at all interesting here. Why?

example uses a reduced table in which age groups are combined in the variable `age.group`, a factor with 15 year intervals except for the last interval, which includes ages 70–90; the methods “toxic gas” and “cooking gas” were collapsed (in the variable `method2`) giving the $2 \times 5 \times 8$ table shown in the output below. These changes do not affect the general nature of the data or conclusions drawn from them.

In this example, we decided to stack the combinations of age and sex, giving an analysis of the loglinear model [`AgeSex`][`Method`], to show how the age-sex categories relate to method of suicide.

In the case of a frequency data frame, it is quite simple to join two or more factors to form the rows of a new two-way table. Here we use `paste()` to form a new, composite factor, called `age_sex` here, abbreviating `sex` for display purposes.

```
> data("Suicide", package="vcd")
> # interactive coding of sex and age.group
> Suicide <- within(Suicide, {
+   age_sex <- paste(age.group, toupper(substr(sex,1,1)))
+ })
```

Then, use `xtabs()` to construct the two-way table `suicide.tab`:

```
> suicide.tab <- xtabs(Freq ~ age_sex + method2, data=Suicide)
> suicide.tab
```

age_sex		method2							
		poison	gas	hang	drown	gun	knife	jump	other
10-20	F	921	40	212	30	25	11	131	100
10-20	M	1160	335	1524	67	512	47	189	464
25-35	F	1672	113	575	139	64	41	276	263
25-35	M	2823	883	2751	213	852	139	366	775
40-50	F	2224	91	1481	354	52	80	327	305
40-50	M	2465	625	3936	247	875	183	244	534
55-65	F	2283	45	2014	679	29	103	388	296
55-65	M	1531	201	3581	207	477	154	273	294
70-90	F	1548	29	1355	501	3	74	383	106
70-90	M	938	45	2948	212	229	105	268	147

The results of the correspondence analysis of this table are shown below:

```
> suicide.ca <- ca(suicide.tab)
> summary(suicide.ca)

Principal inertias (eigenvalues):
```

dim	value	%	cum%	scree	plot
1	0.096151	57.2	57.2	*****	*****
2	0.059692	35.5	92.6	*****	*****
3	0.008183	4.9	97.5	**	
4	0.002158	1.3	98.8	*	
5	0.001399	0.8	99.6		
6	0.000557	0.3	100.0		
7	6.7e-050	0.0	100.0		
Total:	0.168207	100.0			
...					

It can be seen that 92.6% of the χ^2 for this model is accounted for in the first two dimensions. Plotting these gives the display shown in Figure 6.6.

```
> plot(suicide.ca)
```

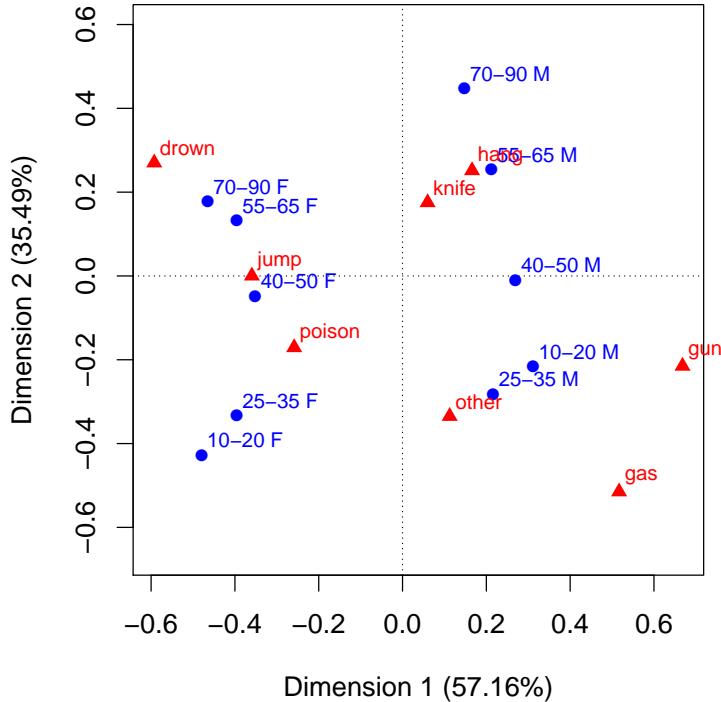


Figure 6.6: 2D CA solution for the stacked [AgeSex][Method] table of the suicide data fig:ca-suicide-plot

Dimension 1 in the plot separates males (right) and females (left), indicating a large difference between suicide profiles of males and females with respect to methods of suicide. The second dimension is mostly ordered by age with younger groups at the bottom and older groups at the top. Note also that the positions of the age groups are roughly parallel for the two sexes. Such a pattern indicates that sex and age do not interact in this analysis.

The relation between the age-sex groups and methods of suicide can be approximately interpreted in terms of similar distance and direction from the origin, which represents the marginal row and column profiles. Young males are more likely to commit suicide by gas or a gun, older males by hanging, while young females are more likely to ingest some toxic agent and older females by jumping or drowning. △

{ex:suicide2}

EXAMPLE 6.6: Suicide rates in Germany: mosaic plot

For comparison, it is useful to see how to construct a mosaic display showing the same associations for the loglinear model $[AS][M]$ as in the correspondence analysis plot. To do this, we first construct the three-way table, `suicide.tab3`,

```
> suicide.tab3 <- xtabs(Freq ~ sex + age.group + method2, data=Suicide)
```

As discussed in Chapter 5, mosaic plots are sensitive both to the order of variables used in successive splits, and to the order of levels within variables and are most effective when these orders are chosen to reflect the some meaningful ordering.

In the present example, `method2` is an unordered table factor, but Figure 6.6 shows that the

methods of suicide vary systematically with both sex and age, corresponding to dimensions 1 and 2 respectively. Here we choose to reorder the table according to the coordinates on Dimension 1. We also delete the low-frequency "other" category to simplify the display.

```
> # methods, ordered as in the table
> suicide.ca$colnames

[1] "poison"   "gas"      "hang"     "drown"    "gun"      "knife"
[7] "jump"     "other"

> # order of methods on CA scores for Dim 1
> suicide.ca$colnames[order(suicide.ca$colcoord[,1])]

[1] "drown"    "jump"     "poison"   "knife"    "other"    "hang"
[7] "gas"      "gun"

> # reorder methods by CA scores on Dim 1
> suicide.tab3 <- suicide.tab3[, , order(suicide.ca$colcoord[,1])]
> # delete "other"
> suicide.tab3 <- suicide.tab3[, , -5]
> ftable(suicide.tab3)

method2 drown jump poison knife hang gas gun
sex   age.group
male  10-20          67 189 1160 47 1524 335 512
      25-35          213 366 2823 139 2751 883 852
      40-50          247 244 2465 183 3936 625 875
      55-65          207 273 1531 154 3581 201 477
      70-90          212 268 938 105 2948 45 229
female 10-20          30 131 921 11 212 40 25
      25-35          139 276 1672 41 575 113 64
      40-50          354 327 2224 80 1481 91 52
      55-65          679 388 2283 103 2014 45 29
      70-90          501 383 1548 74 1355 29 3
```

To construct the mosaic display for the same model analysed by correspondence analysis, we use the argument `expected=~age.group*sex + method2` to supply the model formula. For this large table, it is useful to tweak the labels for the `method2` variable to reduce overplotting; the `labeling_args` argument provides many options for customizing `strucplot` displays.

```
> library(vcdExtra)
> mosaic(suicide.tab3, shade=TRUE, legend=FALSE,
+         expected=~age.group*sex + method2,
+         labeling_args=list(abbreviate_labs=c(FALSE, FALSE, 5)),
+         rot_labels = c(0, 0, 0, 90))
```

This figure (Figure 6.7) again shows the prevalence of gun and gas among younger males and decreasing with age, whereas use of hang increases with age. For females, these three methods are used less frequently, whereas poison, jump, and drown occur more often. You can also see that for females the excess prevalence of these high frequency methods varies somewhat less with age than it does for males.



6.3.2 Marginal tables and supplementary variables

An n -way table in frequency form or case form is automatically collapsed over factors which are not listed in the call to `xtabs()` when creating the table input for `ca()`. The analysis gives a **marginal model** for the categorical variables which *are* listed.

{ca:marginal}

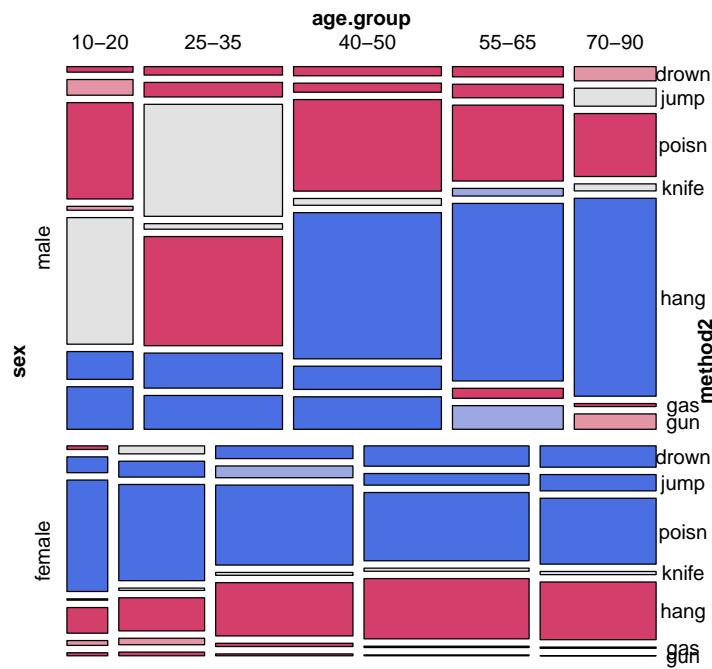


Figure 6.7: Mosaic display showing deviations from the model [AgeSex][Method] for the suicide data

The positions of the categories of the omitted variables may nevertheless be recovered, by treating them as *supplementary variables*, given as additional rows or columns in the two-way table. A supplementary variable is ignored in finding the CA solution, but its categories are then projected into that space. This is another useful trick to extend traditional CA to higher-way tables.

To illustrate, the code below list only the `age` and `method2` variables, and hence produces an analysis collapsed over `sex`. This ignores not only the effect of sex itself, but also all associations of `age` and `method` with `sex`, which are substantial. We don't show the `ca()` result or the plot yet.

```
> # two way, ignoring sex
> suicide.tab2 <- xtabs(Freq ~ age.group + method2, data=Suicide)
> suicide.tab2

      method2
age.group poison  gas  hang drown  gun  knife  jump other
  10-20     2081 375 1736    97 537   58   320   564
  25-35     4495 996 3326   352 916   180   642  1038
  40-50     4689 716 5417   601 927   263   571   839
  55-65     3814 246 5595   886 506   257   661   590
  70-90     2486  74 4303   713 232   179   651   253

> suicide.ca2 <- ca(suicide.tab2)
```

To treat the levels of `sex` as supplementary points, we calculate the two-way table of `sex` and `method`, and append this to the `suicide.tab2` as additional rows:

```
> # relation of sex and method
> suicide.sup <- xtabs(Freq ~ sex + method2, data=Suicide)
> suicide.tab2s <- rbind(suicide.tab2, suicide.sup)
```

In the call to `ca()`, we then indicate these last two rows as supplementary:

```
> suicide.ca2s <- ca(suicide.tab2s, suprow=6:7)
> summary(suicide.ca2s)
```

Principal inertias (eigenvalues):

dim	value	%	cum%	scree	plot
1	0.060429	93.9	93.9	*****	*****
2	0.002090	3.2	97.1	*	
3	0.001479	2.3	99.4		
4	0.000356	0.6	100.0		
Total:	0.064354	100.0			

...

This CA analysis has the same total Pearson chi-square, $\chi^2(28) = 3422.5$ as the result of `chisq.test(suicide.tab2)`. However, the scree plot display above shows that the association between age and method is essentially one-dimensional, but note also that dimension 1 ("age-method") in this analysis has nearly the same inertia (0.0604) as the second dimension (0.0596) in the analysis of the stacked table. We plot the CA results as shown below (see Figure 6.8), and add a line connecting the supplementary points for sex.

```
> op <- par(cex=1.3, mar=c(4, 4, 1, 1)+.1)
> res <- plot(suicide.ca2s, pch=c(16, 15, 17, 24))
> lines(res$rows[6:7,])
> par(op)
```

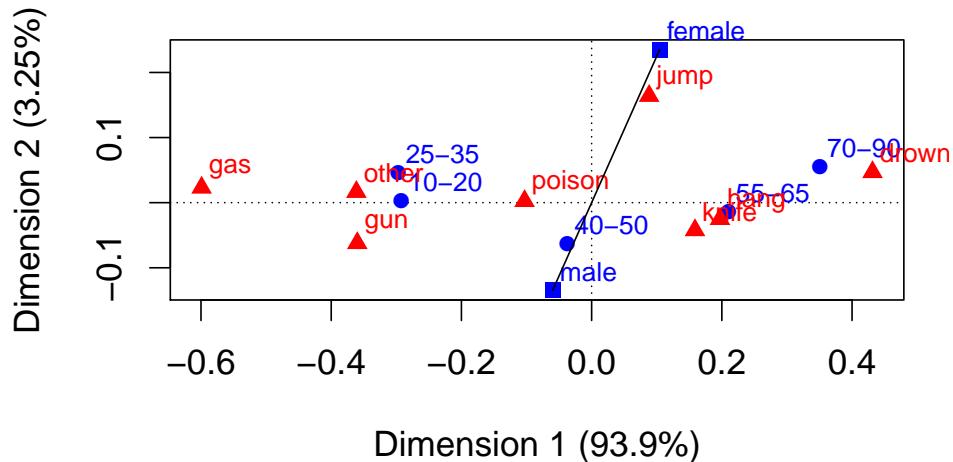


Figure 6.8: 2D CA solution for the [Age] [Method] marginal table. Category points for Sex are shown as supplementary points

Comparing this graph with Figure 6.6, you can see that ignoring sex has collapsed the differences between males and females which were the dominant feature of the analysis including sex. The dominant feature in Figure 6.8 is the Dimension 1 ordering of both age and method. However, as in

Figure 6.6, the supplementary points for sex point toward the methods that are more prevalent for females and males.

6.4 Multiple correspondence analysis

{sec:mca}

Multiple correspondence analysis (MCA) is designed to display the relationships of the categories of two or more discrete variables, but it is best used for multiway tables where the extensions of classical CA described in Section 6.3 do not suffice. Again, this is motivated by the desire to provide an *optimal scaling* of categorical variables, giving scores for the discrete variables in an n -way table with desirable properties and which can be plotted to visualize the relations among the category points.

The most typical development of MCA starts by defining indicator (“dummy”) variables for each category and reexpresses the n -way contingency table in the form of a cases by variables indicator matrix, Z . Simple correspondence analysis for a two-way table can, in fact, be derived as the canonical correlation analysis of the indicator matrix.

Unfortunately, the generalization to more than two variables follows a somewhat different path, so that simple CA does not turn out to be precisely a special case of MCA in some respects, particularly in the decomposition of an interpretable χ^2 over the dimensions in the visual representation.

Nevertheless, MCA does provide a useful graphic portrayal of the *bivariate* relations among any number of categorical variables, and has close relations to the mosaic matrix (Section 5.5). If its limitations are understood, it is helpful in understanding large, multivariate categorical data sets, in a similar way to the use of scatterplot matrices and dimension-reduction techniques (e.g., principal component analysis) for quantitative data.

6.4.1 Bivariate MCA

{sec:mca-bi}

For the hair color, eye color data, the indicator matrix Z has 592 rows and $4 + 4 = 8$ columns. The columns refer to the eight categories of hair color and eye color and the rows to the 592 students in Snee’s 1974 sample.

For simplicity, we show the calculation of the indicator matrix below in frequency form, using `model.matrix()` to compute the dummy (0/1) variables for the levels of hair color (`Hair1–Hair4`) and eye color (`Eye1–Eye4`).

```
> haireye.df <- cbind(
+   as.data.frame(haireye),
+   model.matrix(Freq ~ Hair + Eye, data=haireye,
+   contrasts.arg=list(Hair=diag(4), Eye=diag(4)))[, -1]
+ )
> haireye.df

  Hair   Eye Freq Hair1 Hair2 Hair3 Hair4 Eye1 Eye2 Eye3 Eye4
1 Black Brown  68     1     0     0     0     1     0     0     0
2 Brown Brown 119     0     1     0     0     0     1     0     0     0
3 Red  Brown  26     0     0     1     0     0     1     0     0     0
4 Blond Brown  7      0     0     0     1     1     0     0     0     0
5 Black Blue  20     1     0     0     0     0     0     1     0     0
6 Brown Blue  84     0     1     0     0     0     0     1     0     0
7 Red  Blue  17     0     0     1     0     0     0     1     0     0
8 Blond Blue  94     0     0     0     1     0     1     0     0     0
9 Black Hazel 15     1     0     0     0     0     0     0     1     0
10 Brown Hazel 54     0     1     0     0     0     0     0     1     0
11 Red  Hazel 14     0     0     1     0     0     0     0     1     0
12 Blond Hazel 10     0     0     0     1     0     0     0     1     0
13 Black Green  5      1     0     0     0     0     0     0     0     1
14 Brown Green 29     0     1     0     0     0     0     0     0     1
15 Red  Green 14     0     0     1     0     0     0     0     0     1
16 Blond Green 16     0     0     0     1     0     0     0     0     1
```

Thus, the first row in `haireye.df` represents the 68 individuals having black hair (`Hair1=1`) and brown eyes (`Eye1=1`). The indicator matrix Z is then computed by replicating the rows in `haireye.df` according to the `Freq` value, using the function `expand.dft`. The result has 592 rows and 8 columns.

```
> Z <- expand.dft(haireye.df) [,-(1:2)]
> vnames <- c(levels(haireye.df$Hair), levels(haireye.df$Eye))
> colnames(Z) <- vnames
> dim(Z)

[1] 592    8
```

Note that if the indicator matrix is partitioned as $Z = [Z_1, Z_2]$, corresponding to the two sets of categories, then the contingency table is given by $N = Z_1^T Z_2$.

```
> (N <- t(as.matrix(Z[, 1:4])) %*% as.matrix(Z[, 5:8]))

  Brown Blue Hazel Green
Black   68   20    15     5
Brown  119   84    54    29
Red     26   17    14    14
Blond    7   94    10    16
```

With this setup, MCA can be described as the application of the simple correspondence analysis algorithm to the indicator matrix Z . This analysis would yield scores for the rows of Z (the cases), usually not of direct interest and for the columns (the categories of both variables). As in simple CA, each row point is the weighted average of the scores for the column categories, and each column point is the weighted average of the scores for the row observations.⁵

Consequently, the point for any category is the centroid of all the observations with a response in that category, and all observations with the same response pattern coincide. As well, the origin reflects the weighted average of the categories for *each* variable. As a result, category points with low marginal frequencies will be located further away from the origin, while categories with high marginal frequencies will be closer to the origin. For a binary variable, the two category points will appear on a line through the origin, with distances inversely proportional to their marginal frequencies.

{ex:haireye4}

EXAMPLE 6.7: Hair color and eye color

For expository purposes, we illustrate the analysis of the indicator matrix below for the hair color, eye color data using `ca()`, rather than the function `mjca()` which is designed for a more general approach to MCA.

```
> Z.ca <- ca(Z)
> res <- plot(Z.ca, what=c("none", "all"))
```

In the call to `plot.ca`, the argument `what` is used to suppress the display of the row points for the cases. The plot shown in Figure 6.9 is an enhanced version of this basic plot.

	Dim1	Dim2	factor	levels
1	-0.94250	1.09220	Hair	Black
2	-0.27693	-0.16608	Hair	Brown
3	-0.24194	-1.62513	Hair	Red
4	1.56039	0.35376	Hair	Blond
5	-0.91933	0.44905	Eye	Brown

⁵Note that, in principle, this use of an indicator matrix could be extended to three (or more) variables. That extension is more easily described using an equivalent form, the **Burt matrix**, described in Section 6.4.2.

6	1.02254	0.42176	Eye	Blue
7	-0.39712	-0.85105	Eye	Hazel
8	0.30215	-1.72375	Eye	Green

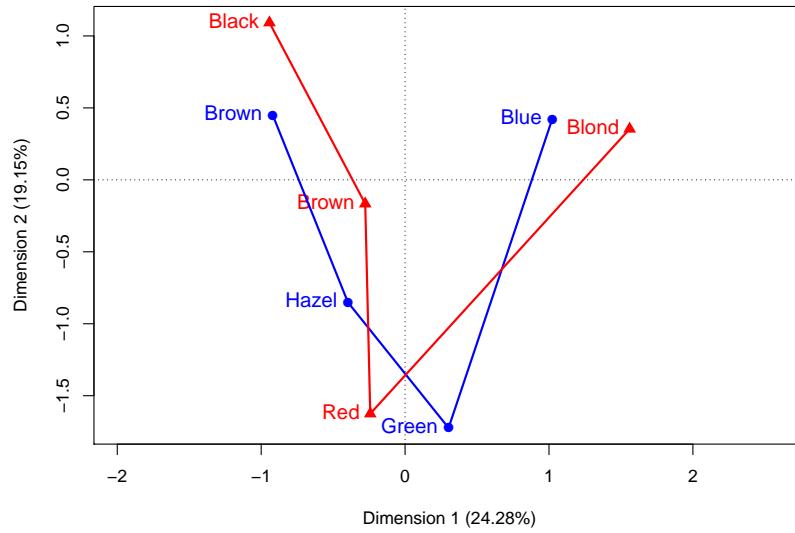


Figure 6.9: Correspondence analysis of the indicator matrix Z for the hair color, eye color data.
The category points are joined separately by lines for the hair color and eye color categories.
Fig:mca-haireye1

Comparing Figure 6.9 with Figure 6.1, we see that the general pattern of the hair color and eye color categories is the same in the analysis of the contingency table (Figure 6.1) and the analysis of the indicator matrix (Figure 6.9), except that the axes are scaled differently—the display has been stretched along the second (vertical) dimension. The interpretation is the same: Dimension 1 reflects a dark–light ordering of both hair and eye colors, and Dimension 2 reflects something that largely distinguishes red hair and green eyes from the other categories.

Indeed, it can be shown (Greenacre, 1984, 2007) that the two displays are identical, except for changes in scales along the axes. There is no difference at all between the displays in standard coordinates. Greenacre (1984, pp. 130–134) describes the precise relations between the geometries of the two analyses.



Aside from the largely cosmetic difference in relative scaling of the axes, a major difference between analysis of the contingency table and analysis of the indicator matrix is in the decomposition of principal inertia and corresponding χ^2 contributions for the dimensions. The plot axes in Figure 6.9 indicate 24.3% and 19.2% for the contributions of the two dimensions, whereas Figure 6.1 shows 89.4% and 9.5%. This difference is the basis for the more general development of MCA methods and is reflected in the `mca.ja()` function illustrated later in this chapter. But first, we describe a second approach to extending simple CA to the multivariate case based on the **Burt matrix**.

6.4.2 The Burt matrix

{sec:mca-burt}

The same solution for the category points as in the analysis of the indicator matrix may be obtained more simply from the so-called **Burt matrix** (Burt, 1950),

$$\mathbf{B} = \mathbf{Z}^T \mathbf{Z} = \begin{bmatrix} \mathbf{N}_1 & \mathbf{N} \\ \mathbf{N}^T & \mathbf{N}_2 \end{bmatrix},$$

where \mathbf{N}_1 and \mathbf{N}_2 are diagonal matrices containing the marginal frequencies of the two variables (the column sums of \mathbf{Z}_1 and \mathbf{Z}_2). In this representation, the contingency table of the two variables, \mathbf{N} appears in the off-diagonal block, \mathbf{N} in this equation. This calculation is shown below.

```
> Burt <- t(as.matrix(Z)) %*% as.matrix(Z)
> rownames(Burt) <- colnames(Burt) <- vnames
> Burt
```

	Black	Brown	Red	Blond	Brown	Blue	Hazel	Green
Black	108	0	0	0	68	20	15	5
Brown	0	286	0	0	119	84	54	29
Red	0	0	71	0	26	17	14	14
Blond	0	0	0	127	7	94	10	16
Brown	68	119	26	7	220	0	0	0
Blue	20	84	17	94	0	215	0	0
Hazel	15	54	14	10	0	0	93	0
Green	5	29	14	16	0	0	0	64

The standard coordinates from an analysis of the Burt matrix \mathbf{B} are identical to those of \mathbf{Z} . (However, the singular values of \mathbf{B} are the squares of those of \mathbf{Z} .) Then, the following code, using Burt produces the same display of the category points for hair color and eye color as shown for the indicator matrix Z in Figure 6.9.

```
> Burt.ca <- ca(Burt)
> plot(Burt.ca)
```

6.4.3 Multivariate MCA

{sec:mca-multi}

The coding of categorical variables in an indicator matrix and the relationship to the Burt matrix provides a direct and natural way to extend this analysis to more than two variables. If there are Q categorical variables, and variable q has J_q categories, then the Q -way contingency table, of size $J = \prod_{q=1}^Q J_q = J_1 \times J_2 \times \cdots \times J_Q$, with a total of $n = n_{++\dots}$ observations may be represented by the partitioned $(n \times J)$ indicator matrix $[\mathbf{Z}_1 \ \mathbf{Z}_2 \ \dots \ \mathbf{Z}_Q]$.

Then the Burt matrix is the symmetric partitioned matrix

$$\mathbf{B} = \mathbf{Z}^T \mathbf{Z} = \begin{bmatrix} \mathbf{N}_1 & \mathbf{N}_{[12]} & \cdots & \mathbf{N}_{[1Q]} \\ \mathbf{N}_{[21]} & \mathbf{N}_2 & \cdots & \mathbf{N}_{[2Q]} \\ \vdots & \vdots & \ddots & \vdots \\ \mathbf{N}_{[Q1]} & \mathbf{N}_{[Q2]} & \cdots & \mathbf{N}_Q \end{bmatrix}, \quad (6.7) \quad \text{eq:burt}$$

where again the diagonal blocks \mathbf{N}_i contain the one-way marginal frequencies. The off-diagonal blocks $\mathbf{N}_{[ij]}$ contain the bivariate marginal contingency tables for each pair (i, j) of variables.

Classical MCA (see, e.g., Greenacre (1984), Gower and Hand (1996)) can then be defined as a singular value decomposition of the matrix \mathbf{B} which produces scores for the categories of *all* variables so that the greatest proportion of the bivariate, pairwise associations in all off-diagonal blocks is accounted for in a small number of dimensions.

In this respect, MCA resembles multivariate methods for quantitative data based on the joint bivariate correlation or covariance matrix (Σ) and there is some justification to regard the Burt matrix as the categorical analog of Σ .⁶

There is a close connection between this analysis and the bivariate mosaic matrix (Section 5.5): The mosaic matrix displays the residuals from independence for each pair of variables, and thus provides a visual representation of the Burt matrix. The one-way margins shown (by default) in the diagonal cells reflect the diagonal matrices N_i in Eqn. (6.7). The total amount of shading in all the individual mosaics portrays the total pairwise associations decomposed by MCA. See Friendly (1999a) for further details.

For interpretation of MCA plots, we note the following relations (Greenacre, 1984, §5.2):⁷

- The inertia contributed by a given variable increases with the number of response categories.
- The centroid of the categories for each discrete variable is at the origin of the display.
- For a particular variable, the inertia contributed by a given category increases as the marginal frequency in that category *decreases*. Low frequency points therefore appear further from the origin.
- The category points for a binary variable lie on a line through the origin. The distance of each point to the origin is inversely related to the marginal frequency.

{ex:marital3}

EXAMPLE 6.8: Marital status and pre- and extramarital sex

The data on the relation between marital status and reported premarital and extramarital sex was explored earlier using mosaic displays in Example 5.8 and Example 5.12.

Using the `ca` package, an MCA analysis of the `PreSex` data is carried out using `mjca()`. This function typically takes a data frame in *case form* containing the factor variables, but converts a table to this form. This example analyzes the Burt matrix calculated from the `PreSex` data, specified as `lambda="Burt"`

```
> data("PreSex", package="vcd")
> PreSex <- aperm(PreSex, 4:1) # order variables G, P, E, M
> presex.mca <- mjca(PreSex, lambda="Burt")
> summary(presex.mca)
```

Principal inertias (eigenvalues) :

dim	value	%	cum%	scree	plot
1	0.149930	53.6	53.6	*****	*****
2	0.067201	24.0	77.6	*****	*****
3	0.035396	12.6	90.2	**	**
4	0.027365	9.8	100.0		
<hr/>					
Total:	0.279892	100.0			

The output from `summary()` seems to show that 77.6% of the total inertia is accounted for in two dimensions. A basic, default plot of the MCA solution is provided by the `plot()` method for "mjca" objects.

```
> plot(presex.mca)
```

This plotting method is not very flexible in terms of control of graphical parameters or the ability

⁶For multivariate normal data, however, the mean vector and covariance matrix are sufficient statistics, so all higher-way relations are captured in the covariance matrix. This is not true of the Burt matrix. Moreover, the covariance matrix is typically expressed in terms of mean-centered variables, while the Burt matrix involves the marginal frequencies. A more accurate statement is that the uncentered covariance matrix is analogous to the Burt matrix.

⁷This book, now out of print, is available for free download at <http://www.carme-n.org/>

to add additional annotations (labels, lines, legend) to ease interpretation. Instead, we use the plot method to create an empty plot (with no points or labels), and return the calculated plot coordinates (`res`) for the categories. A bit of processing of the coordinates provides the customized display shown in Figure 6.10.

```
> # plot, but don't use point labels or points
> res <- plot(presex.mca, labels=0, pch='.', cex.lab=1.2)
>
> # extract factor names and levels
> coords <- data.frame(res$cols, presex.mca$ factors)
> cols <- c("blue", "red", "brown", "black")
> nlev <- presex.mca$levels.n
>
> points(coords[,1:2], pch=rep(16:19, nlev), col=rep(cols, nlev), cex=1.2)
> text(coords[,1:2], label=coords$level, col=rep(cols, nlev), pos=3,
+ + cex=1.2, xpd=TRUE)
> lines(Dim2 ~ Dim1, data=coords, subset=factor=="Gender",
+ + lty=1, lwd=2, col="blue")
> lines(Dim2 ~ Dim1, data=coords, subset=factor=="PremaritalSex",
+ + lty=1, lwd=2, col="red")
> lines(Dim2 ~ Dim1, data=coords, subset=factor=="ExtraMaritalSex",
+ + lty=1, lwd=2, col="brown")
> lines(Dim2 ~ Dim1, data=coords, subset=factor=="MaritalStatus",
+ + lty=1, lwd=3, col="black")
>
> legend("bottomright", legend=c("Gender", "PreSex", "ExtraSex", "Marital"),
+ + title="Factor", title.col="black",
+ + col=cols, text.col=cols, pch=16:19,
+ + bg="gray95", cex=1.2)
```

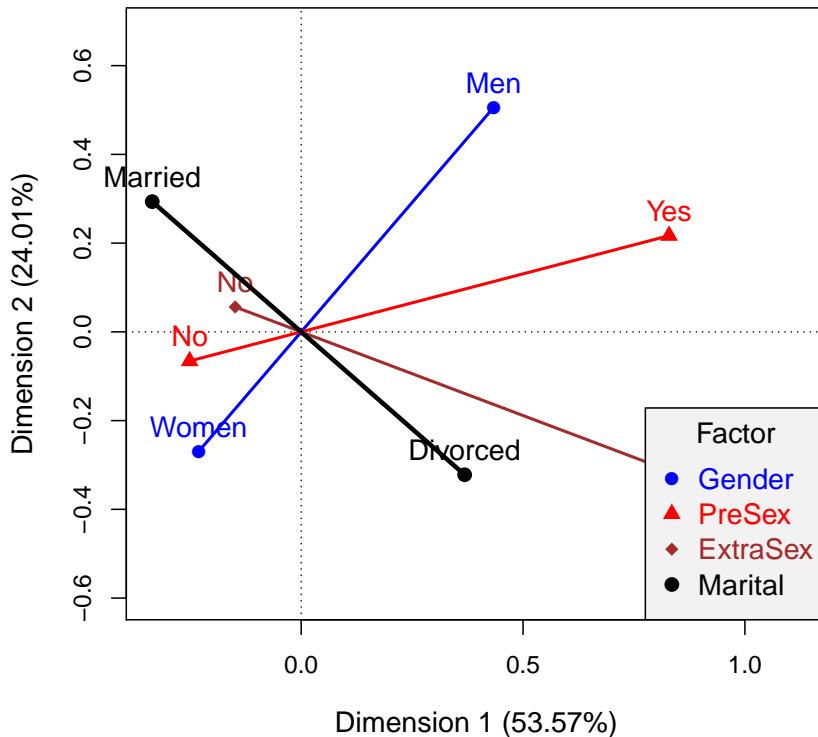


Figure 6.10: MCA plot of the Burt matrix for the PreSex data. The category points are joined separately by lines for the factor variables.
Fig:presex-mca-plot

As indicated above, the category points for each factor appear on lines through the origin, with

distances inversely proportional to their marginal frequencies. For example, the categories for No premarital and extramarital sex are much larger than the corresponding Yes categories, so the former are positioned closer to the origin. In contrast, the categories of gender and marital status are more nearly equal marginally.

Another aspect of interpretation of Figure 6.10 concerns the alignment of the lines for different factors. The positions of the category points on Dimension 1 suggest that Women are less likely to have had pre-marital and extra-marital sex and that still being married is associated with the absence of pre- and extra-marital sex. As well, the lines for gender and marital status are nearly at right angles, suggesting that these variables are unassociated. This interpretation is more or less correct, but it is only approximate in this MCA scaling of the coordinate axes. An alternative scaling, based on a *biplot* representation is described in Section 6.5.

If you compare the MCA result in Figure 6.10 with the mosaic matrix in Figure 5.22, you will see that they are both showing the bivariate pairwise associations among these variables, but in different ways. The mosaic plots show the details of marginal and joint frequencies together with residuals from independence for each 2×2 marginal subtable. The MCA plot using the Burt matrix summarizes each category point in terms of a 2D representation of contributions to total inertia (association). \triangle

6.4.3.1 Inertia decomposition

The transition from simple CA to MCA is straight-forward in terms of the category scores derived from the indicator matrix Z or the Burt matrix, B . It is less so in terms of the calculation of total inertia, and therefore in the chi-square values and corresponding percentages of association accounted for in some number of dimensions.

In simple CA, the total inertia is χ^2/n , and it therefore makes sense to talk of percentage of association accounted for by each dimension. But in MCA of the indicator matrix the total inertia, $\sum \lambda$, is simply $(J - Q)/Q$, because the inertia of each subtable, Z_i is equal to its dimensionality, $J_i - 1$, and the total inertia of an indicator matrix is the average of the inertias of its subtables. Consequently, the average inertia per dimension is $1/Q$, and it is common to interpret only those dimensions that exceed this average (analogous to the use of 1 as a threshold for eigenvalues in principal components analysis).

To more adequately reflect the percentage of association in MCA, Greenacre (1990) (see also Greenacre (2007, Chapter 19) for details), revising an earlier proposal by Benzécri (1977), suggested the calculation of *adjusted inertia*, which ignores the contributions of the diagonal blocks in the Burt matrix,

$$\{\text{eq:benzecri}\} (\lambda_i^*)^2 = \left[\frac{Q}{Q-1} \left(\lambda_i^Z - \frac{1}{Q} \right) \right]^2 \quad (6.8)$$

as the principal inertia due to the dimensions with $(\lambda^Z)^2 > 1/Q$. This adjustment expresses the contribution of each dimension as $(\lambda_i^*)^2 / \sum(\lambda_i^*)^2$, with the summation over only dimensions with $(\lambda^Z)^2 > 1/Q$.

A related method, also handled by `mjca()`, is *joint correspondence analysis* (Greenacre, 1988, ?), an iterative method that replaces the diagonal blocks of the Burt matrix with values that minimize their impact on inertia. Unlike MCA, solutions in JCA are not nested, however.

EXAMPLE 6.9: Survival on the Titanic

An MCA analysis of the *Titanic* data is carried out using `mjca()` as shown below.

```
> titanic.mca <- mjca(Titanic)
```

`mjca()` allows different scaling methods for the contributions to inertia of the different dimensions. The default (`lambda="adjusted"`), used here, is the adjusted inertias as in Eqn. (6.8).

```
> summary(titanic.mca)

Principal inertias (eigenvalues):

  dim      value      %   cum%   scree plot
  1      0.067655  76.8  76.8 ****
  2      0.005386   6.1  82.9 **
  3      00000000   0.0  82.9
  -----
  Total: 0.088118
```

Using similar code to that used in Example 6.8, Figure 6.11 shows an enhanced version of the default plot that connects the category points for each factor by lines using the result returned by the `plot()` function.

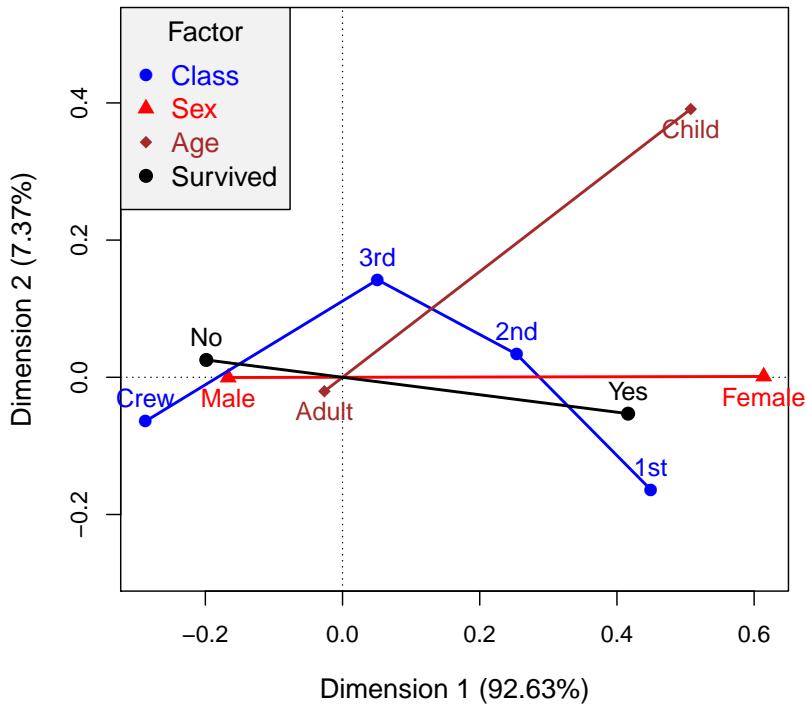


Figure 6.11: MCA plot of the Titanic data. The category points are joined separately by lines for the factor variables.
[Fig:titanic-mca-plot]

In this plot, the points for each factor have the property that the sum of coordinates on each dimension, weighted inversely by the marginal proportions, equals zero. Thus high frequency categories (e.g., Adult and Male) are close to the origin.

The first dimension is perfectly aligned with gender, and also strongly aligned with Survival. The second dimension pertains mainly to Class and Age effects. Considering those points which differ from the origin most similarly (in distance and direction) to the point for Survived, gives the

interpretation that survival was associated with being female or upper class or (to a lesser degree) being a child.



6.5 Biplots for contingency tables

{sec:biplot}

Like correspondence analysis, the *biplot* (Bradu and Gabriel, 1978, Gabriel, 1971, 1980, 1981, Gower *et al.*, 2011) is a visualization method which uses the SVD to display a matrix in a low-dimensional (usually 2-dimensional) space. They differ in the relationships in the data that are portrayed, however:

- In correspondence analysis the (weighted, χ^2) *distances* between row points and distances between column points are designed to reflect *differences* between the row profiles and column profiles.
- In the biplot, on the other hand, row and column points are represented by *vectors* from the origin such that the projection (inner product) of the vector a_i for row i on b_j for column j approximates the data element y_{ij} ,

{eq:biplot1}

$$\mathbf{Y} \approx \mathbf{AB}^\top \iff y_{ij} \approx a_i^\top b_j . \quad (6.9)$$

Geometrically, Eqn. (6.9) may be described as approximating the data value y_{ij} by the projection of the end point of vector a_i on b_j (and vice-versa), as shown in Figure 6.12.

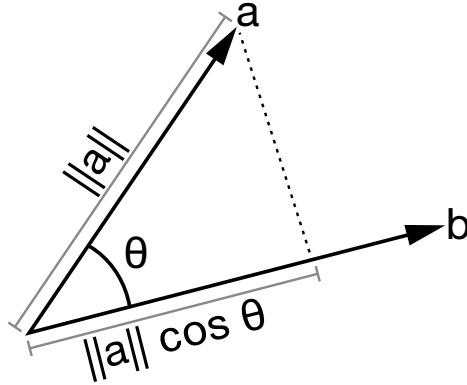


Figure 6.12: The scalar product of vectors of two points from the origin is the length of the projection of one vector on the other.

6.5.1 CA bilinear biplots

As in CA, there are a number of different representations of coordinates for row and column points for a contingency table within a biplot framework. One set of connections between CA and the biplot can be seen through the *reconstitution formula*, giving the decomposition of the correspondence matrix $\mathbf{P} = N/n$ in terms of the standard coordinates Φ and Γ defined in Eqn. (6.4) and Eqn. (6.5) as:

{fig:Scalarproduct}

$$p_{ij} = r_i c_j \left(1 + \sum_{m=1}^M \sqrt{\lambda_m} \phi_{im} \gamma_{jm} \right) \quad (6.10)$$

or, in matrix terms,

$$\mathbf{P} = \mathbf{D}_r (\mathbf{1}\mathbf{1}^\top + \boldsymbol{\Phi} \mathbf{D}_\lambda^{1/2} \boldsymbol{\Gamma}^\top) \mathbf{D}_c \quad (6.11)$$

The CA solution approximates this by a sum over $d \ll M$ dimensions, or by using only the first d (usually 2) columns of $\boldsymbol{\Phi}$ and $\boldsymbol{\Gamma}$.

Eqn. (6.10) can be re-written in biplot scalar form as

$$\left(\frac{p_{ij}}{r_i c_j} \right) - 1 \approx \sum_{m=1}^d (\sqrt{\lambda_m} \phi_{im}) \gamma_{jm} = \sum_{m=1}^d f_{im} \gamma_{jm} \quad (6.12) \quad \text{(eq:rowprincipal)}$$

where $f_{im} = (\sqrt{\lambda_m} \phi_{im})$ gives the principal coordinates of the row points. The left-hand side of Eqn. (6.12) contains the **contingency ratios**, $p_{ij}/r_i c_j$ of the observed cell probabilities to their expected values under independence. This shows that an **asymmetric CA plot** of row principal coordinates \mathbf{F} and the column standard coordinates $\boldsymbol{\Gamma}$ is a biplot that approximates the deviations of the contingency ratios from their values under independence.

In the **ca** package, this plot is obtained by specifying `map="rowprincipal"` in the call to `plot()`, or `map="colprincipal"` to plot the column points in principal coordinates. It is typical in such biplots to display one set of coordinates as points and the other as vectors from the origin, as controlled by the `arrows` argument, so that one can interpret the data values represented as approximated by the projections of the points on the vectors.

Two other types asymmetric “maps” are also defined with different scalings that turn out to have better visual properties in terms of representing the relations between the row and column categories, particularly when the strength of association (inertia) in the data is low.

- The option `map="rowgab"` (or `map="colgab"`) gives a biplot form proposed by Gabriel and Odoroff (1990) with the rows (columns) shown in principal coordinates and the columns (rows) in standard coordinates multiplied by the mass c_j (r_i) of the corresponding point.
- The **contribution biplot** for CA (Greenacre, 2013), with the option `map="rowgreen"` (or `map="colgreen"`) provides a reconstruction of the standardized residuals from independence, using the points in standard coordinates multiplied by the square root of the corresponding masses. This has the nice visual property of showing more directly the contributions of the vectors to the low-dimensional solution.

(ex:suicide3)

EXAMPLE 6.10: Suicide rates in Germany: biplot

To illustrate the biplot representation, we continue with the data on suicide rates in Germany from Example 6.5 using the stacked table `suicide.tab` comprised of the age–sex combinations as rows and methods of suicide as columns.

```
> suicide.tab <- xtabs(Freq ~ age_sex + method2, data=Suicide)
> suicide.ca <- ca(suicide.tab)
```

Using this result, `suicide.ca`, in the call to `plot()` below, we use `map="colgreen"` and vectors represent the methods of suicide, as shown in Figure 6.13.

```
> plot(suicide.ca, map="colgreen", arrows=c(FALSE, TRUE))
```

The interpretation of the row points for the age–sex categories is similar to what we saw earlier in Figure 6.6. But now, the vectors for the suicide categories reflect the contributions of those

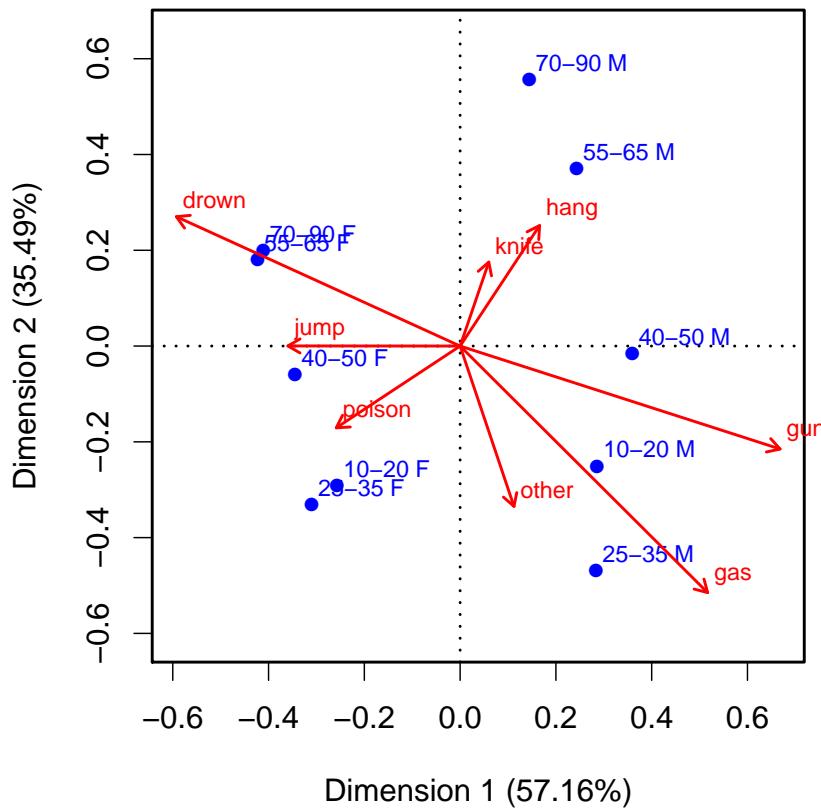


Figure 6.13: CA biplot of the suicide data using the contribution biplot scaling. Associations between the age-sex categories and the suicide methods can be read as the projections of the points on the vectors. The lengths of the vectors for the suicide categories reflect their contributions to this representation in a 2D plot. ^{Fig.ca-suicide-biplot}

methods to the representation of association. Thus, the methods `drown`, `gun` and `gas` have large contributions, while `knife`, `hang`, and `poison` are relatively small. Moreover, the projections of the points for the age-sex combinations on the method vectors reflect the standardized residuals from independence.

The most comprehensive modern treatment of biplot methodology is the book *Understanding Biplots* (Gower *et al.*, 2011). Together with the book, they provide an R package, `UBbipl`, that is capable of producing an astounding variety of high-quality plots. Unfortunately, that package is only available on their publisher's web site⁸ and you need the book to be able to use it because all the documentation is in the book. Nevertheless, we illustrate the use of the `cabipl()` function to produce the version of the CA biplot shown in Figure 6.14.

```
> library(UBbipl)
> cabipl(as.matrix(suicide.tab),
+ axis.col = gray(.4), ax.name.size=1,
+ ca.variant = "PearsonResA",
+ markers = FALSE,
+ row.points.size = 1.5,
+ row.points.col = rep(c("red", "blue"), 4),
+ plot.col.points = FALSE,
```

⁸<http://www.wiley.com/legacy/wileychi/gower/material.html>

```
+     marker.col = "black", marker.size=0.8,
+     offset = c(2, 2, 0.5, 0.5),
+     offset.m = rep(-0.2, 14),
+     output=NULL)
```

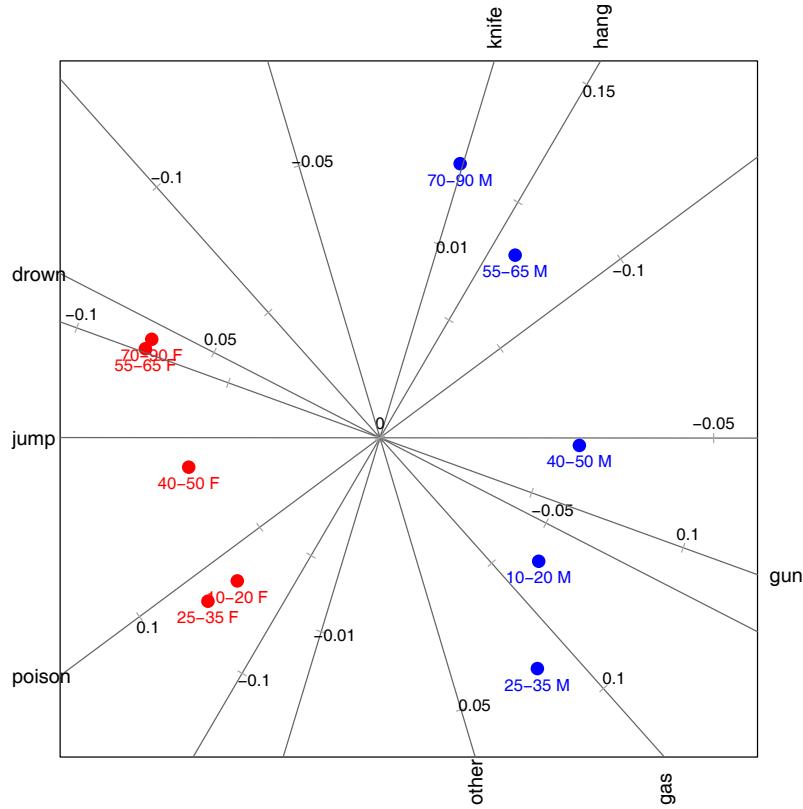


Figure 6.14: CA biplot of the suicide data, showing calibrated axes for the suicide methods.

{fig:cabipl-suicide}

This plot uses `ca.variant = "PearsonResA"` to specify that the biplot is to approximate the standardized Pearson residuals by the inner product of each row point on the vector for the column point for the suicide methods, as also in Figure 6.13. However, Figure 6.14 represents the methods calibrated axis lines, designed to be read as scales for the projections of the row points (age–sex) on the methods. The `UBbipl` package has a huge number of options for controlling the details of the biplot display. See (Gower *et al.*, 2011, Ch. 2) for all the details.



6.5.2 Biadditive biplots

A different use of biplots for contingency tables stems from the close analogy between additive relations for a quantitative response when there is no interaction between factors, and the multiplicative relations for a contingency table when there is no association.

For quantitative data Bradu and Gabriel (1978) show how the biplot can be used to diagnose additive relations among rows and columns. For example, when a two-way table is well-described

by a two-factor ANOVA model with no interaction,

$$y_{ij} = \mu + \alpha_i + \beta_j + \epsilon_{ij} \iff \mathbf{Y} \approx \mathbf{a}\mathbf{1}^\top + \mathbf{1}\mathbf{b}^\top$$

then, the row points, \mathbf{a}_i , and the column points, \mathbf{b}_j , will fall on two straight lines at right angles to each other in the biplot. For a contingency table, the multiplicative relations among frequencies under independence become additive relations in terms of log frequency, and Gabriel *et al.* (1997) illustrate how biplots of log frequency can be used to explore associations in two-way and three-way tables.

That is, For a two-way table, independence, $A \perp B$, implies that ratios of frequencies should be proportional for any two rows, i, i' and any two columns, j, j' . Equivalently, this means that the log odds ratio for all such sets of four cells should be zero:

$$A \perp B \iff \log \theta_{ii',jj'} = \log \left(\frac{n_{ij}n_{i'j'}}{n_{i'j}n_{ij'}} \right) = 0$$

Now, if the log frequencies have been centered by subtracting the grand mean, Gabriel *et al.* (1997) show that $\log \theta_{ii',jj'}$ is approximated in the biplot (of $\log(n_{ij}) - \bar{\log}(n_{ij})$)

$$\log \theta_{ii',jj'} \approx \mathbf{a}_i^\top \mathbf{b}_j - \mathbf{a}_{i'}^\top \mathbf{b}_j - \mathbf{a}_i^\top \mathbf{b}_{j'} + \mathbf{a}_{i'}^\top \mathbf{b}_{j'} = (\mathbf{a}_i - \mathbf{a}_{i'})^\top (\mathbf{b}_j - \mathbf{b}_{j'})$$

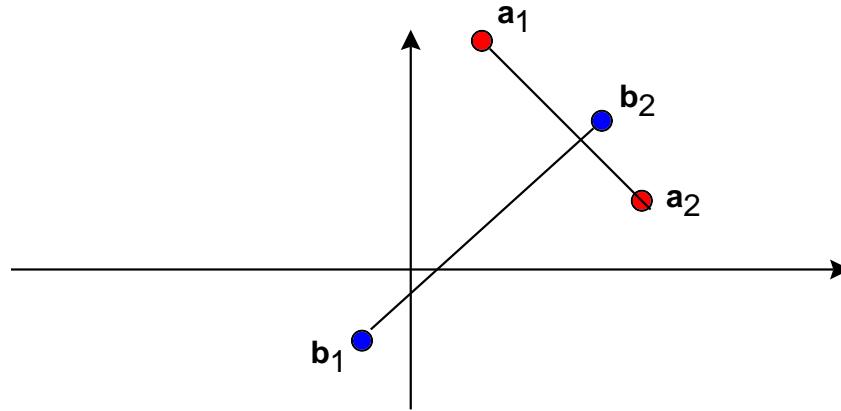


Figure 6.15: Independence implies orthogonal vector differences in a biplot of log frequency. The line joining \mathbf{a}_1 to \mathbf{a}_2 represents $(\mathbf{a}_1 - \mathbf{a}_2)$. This line is perpendicular to the line $(\mathbf{b}_1 - \mathbf{b}_2)$ under independence.

{fig:bidemo}

Therefore, this biplot criterion for independence in a two-way table is whether $(\mathbf{a}_i - \mathbf{a}_{i'})^\top (\mathbf{b}_j - \mathbf{b}_{j'}) \approx 0$ for all pairs of rows, i, i' , and all pairs of columns, j, j' . But $(\mathbf{a}_i - \mathbf{a}_{i'})$ is the vector connecting \mathbf{a}_i to $\mathbf{a}_{i'}$ and $(\mathbf{b}_j - \mathbf{b}_{j'})$ is the vector connecting \mathbf{b}_j to $\mathbf{b}_{j'}$, as shown in Figure 6.15, and the inner product of any two vectors equals zero iff they are orthogonal. Hence, this criterion implies that all lines connecting pairs of row points are orthogonal to lines connecting pairs of column points, as illustrated in Figure 6.15.

{ex:soccer3}

EXAMPLE 6.11: UK Soccer scores

We examined the data on UK Soccer scores in Example 5.5 and saw that the number of goals scored by the home and away teams were largely independent (see Figure 5.10). This data set provides a good test of the ability of the biplot to diagnose independence.

```
> data("UKSoccer", package="vcd")
> dimnames(UKSoccer) <- list(Home=paste0("H", 0:4),
+                               Away=paste0("A", 0:4))
```

Basic biplots in R are provided by `biplot()` that works mainly with the result calculated by `prcomp()` or `princomp()`. Here, we use `prcomp()` on the log frequencies in the `UKSoccer` table, adding 1, because there is one cell with zero frequency.

```
> soccer.pca <- prcomp(log(UKSoccer+1), center=TRUE, scale.=FALSE)
```

The result is plotted using a customized plot based on `biplot()` as shown in Figure 6.16.

```
> biplot(soccer.pca, scale=0, var.axes=FALSE,
+         col=c("blue", "red"), cex=1.2, cex.lab=1.2,
+         xlab="Dimension 1", ylab="Dimension 2")
```

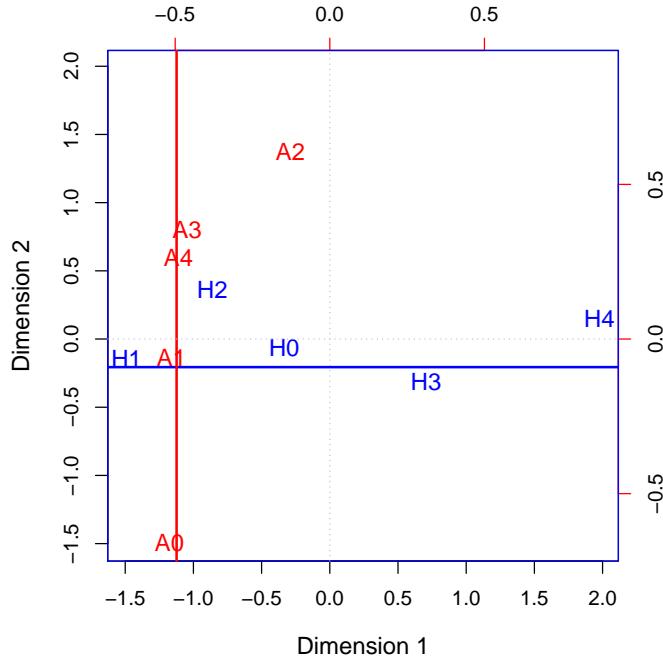


Figure 6.16: Biplot for the biadditive representation of independence for the UK Soccer scores. The row and column categories are independent in this plot when they appear as points on approximately orthogonal lines.¹

To supplement this plot and illustrate the orthogonality of row and column category points under independence, we added horizontal and vertical lines as calculated below, using the results returned by `prcomp()`. The initial version of this plot showed that two points, A2 and H2 did not align with the others, so these were excluded from the calculations.

```
> # get the row and column scores
> rscores <- soccer.pca$x[,1:2]
> cscores <- soccer.pca$rotation[,1:2]
```

```
> # means, excluding A2 and H2
> rmean <- colMeans(rscores[-3,]) [2]
> cmean <- colMeans(cscores[-3,]) [1]
>
> abline(h=rmean, col="blue", lwd=2)
> abline(v=cmean, col="red", lwd=2)
> abline(h=0, lty=3, col="gray")
> abline(v=0, lty=3, col="gray")
```

You can see that all the A points (except for A2) and all the H points (except for H2) lie along straight lines, and these lines are indeed at right angles, signifying independence. The fact that these straight lines are parallel to the coordinate axes is incidental, and unrelated to the independence interpretation.



6.6 Chapter summary

{sec:ca-summary}

- Correspondence analysis is an exploratory technique, designed to show the row and column categories in a two- (or three-) dimensional space. These graphical displays, and various extensions, provide ways to interpret the patterns of association and explore visually the adequacy of certain loglinear models.
- The scores assigned to the categories of each variable are optimal in several equivalent ways. Among other properties, they maximize the (canonical) correlations between the quantified variables (weighted by cell frequencies), and make the regressions of each variable on the other most nearly linear, for each CA dimension.
- Multi-way tables may be analyzed in several ways. In the “stacking” approach, two or more variables may be combined interactively in the rows and/or columns of an n -way table. Simple CA of the restructured table reveals associations between the row and column categories of the restructured table, but hides associations between the variables combined interactively. Each way of stacking corresponds to a particular loglinear model for the full table.
- Multiple correspondence analysis is a generalization of CA to two or more variables based on representing the data as an indicator matrix, or the Burt matrix. The usual MCA provides an analysis of the joint, bivariate relations between all pairs of variables.
- The biplot is a related technique for visualizing the elements of a data array by points or vectors in a joint display of their row and column categories. A standard CA biplot represents the contributions to lack of independence as the projection of the points for rows (or columns) on vectors for the other categories.
- Another application of the biplot to contingency table data is described, based on analysis of log frequency. This analysis also serves to diagnose patterns of independence and partial independence in two-way and larger tables.

6.7 Lab exercises

{sec:lab:6lab}

Exercise 6.1 The *JobSat* data in *vcdExtra* gives a 4×4 table recording job satisfaction in relation to income.

- Carry out a simple correspondence analysis on this table. How much of the inertia is accounted for by a one-dimensional solution? How much by a two-dimensional solution?

- (b) Plot the 2D CA solution. To what extent can you consider the association between job satisfaction and income “explained” by the ordinal nature of these variables?

{lab:6.2}

Exercise 6.2 Refer to Exercise 1 in Chapter 5. Carry out a simple correspondence analysis on the 4×5 table *criminal* from the *logmult* package.

- (a) What percentages of the Pearson χ^2 for association are explained by the various dimensions?
 (b) Plot the 2D correspondence analysis solution. Describe the pattern of association between year and age.

{lab:6.3}

Exercise 6.3 The data set *caith* in *MASS* gives a classic table tabulating hair color and eye color of people in Caithness, Scotland, originally from Fisher (1940).

- (a) Carry out a simple correspondence analysis on this table. How many dimensions seem necessary to account for most of the association in the table?
 (b) Plot the 2D solution. The interpretation of the first dimension should be obvious; is there any interpretation for the second dimension?

{lab:6.4}

Exercise 6.4 The same data, plus a similar table for Aberdeen, are given as a three-way table as *HairEyePlace* in *vcdExtra*.

- (a) Carry out similar correspondence analysis analysis to the last exercise for the data from Aberdeen. Comment on any differences in the placement of the category points.
 (b) Analyze the three-way table, stacked to code hair color and place interactively, i.e., for the loglinear model [*HairPlace*][*Eye*]. What does this show?

{lab:6.5}

Exercise 6.5 For the mental health data analyzed in Example 6.2, construct a shaded sieve diagram and mosaic plot. Compare these with the correspondence analysis plot shown in Figure 6.2. What features of the data and the association between SES and mental health status are shown in each?

{lab:6.6}

Exercise 6.6 Simulated data is often useful to help understand the connections between data, analysis methods and associated graphic displays. Section 6.3.1 illustrated interactive coding in R, using a simulated 4-way table of counts of pets, classified by age, color and sex, but with no associations because the counts had a constant Poisson mean, $\lambda = 15$.

- (a) Re-do this example, but in the call to *rpois()*, specify a non-negative vector of Poisson means to create some associations among the table factors.
 (b) Use CA methods to determine if and how the structure you created in the data appears in the results.

{lab:EV3}

Exercise 6.7 The *TV* data was analyzed using CA in Example 6.4, ignoring the variable *Time*. Carry out analyses of the 3-way table, reducing the number of levels of *Time* to three hourly intervals as shown below.

```
> data("TV", package="vcdExtra")
> # reduce number of levels of Time
> TV.df <- as.data.frame.table(TV)
> levels(TV.df$Time) <- rep(c("8", "9", "10"), c(4, 4, 3))
> TV3 <- xtabs(Freq ~ Day + Time + Network, TV.df)
> structable(Day ~ Network + Time, TV3)
```

		Day	Monday	Tuesday	Wednesday	Thursday	Friday
Network	Time						
ABC	8		536	861	744	735	1119
	9		1401	1205	1022	682	907
	10		910	1044	668	349	711

CBS	8	1167	646	550	680	509
	9	967	959	409	385	544
	10	789	798	324	270	426
NBC	8	858	1090	512	1927	823
	9	946	890	831	1858	590
	10	825	588	869	2101	585

- (a) Use the stacking approach (Section 6.3) to perform a CA of the table with `Network` and `Time` coded interactively. You can create this using the `as.matrix()` method for a "structable" object.

```
> TV3S <- as.matrix(structable(Day ~ Network + Time, TV3), sep=":")
```

- (b) What loglinear model is analyzed by this approach?
 (c) Plot the 2D solution. Compare this to the CA plot of the two-way table in Figure 6.4.
 (d) Carry out an MCA analysis using `mjca()` of the three-way table `TV3`. Plot the 2D solution, and compare this with both the CA plot and the solution for the stacked three-way table.

{lab6ipbese8}

Exercise 6.8 Refer to the MCA analysis of the `PreSex` data in Example 6.8. Use the stacking approach to analyze the stacked table with the combinations of premarital and extramarital sex in the rows and the combinations of gender and marital status in the columns. As suggested in the exercise above, you can use `as.matrix(structable())` to create the stacked table

- (a) What loglinear model is analyzed by this approach? Which associations are included and which are excluded in this analysis?
 (b) Plot the 2D CA solution for this analysis. You might want to draw lines connecting some of the row points or column points to aid in interpretation.
 (c) How does this analysis differ from the MCA analysis shown in Figure 6.10?

```
> #remove(list=objects(pattern="\\".tab/\\".df/\\".fit"))
> .locals$ch06 <- setdiff(ls(), .globals)
> #.locals$ch06
> remove(list=.locals$ch06[sapply(.locals$ch06,function(n){!is.function(get(n))})])
```

Chapter 7

Logistic Regression Models

This chapter introduces the modeling framework for categorical data in the simple situation where we have a categorical response variable, often binary, and one or more explanatory variables. A fitted model provides both statistical inference and prediction, accompanied by measures of uncertainty. Data visualization methods for discrete response data must often rely on smoothing techniques, including both direct, non-parametric smoothing and the implicit smoothing that results from a fitted parametric model. Diagnostic plots help us to detect influential observations which may distort our results.

{ch:logistic}

7.1 Introduction

{sec:logist-intro}

All models are wrong, but some are useful

George E. P. Box, (Box and Draper, 1987, p. 424)

Chapters 4–6 have been concerned primarily with simple, exploratory methods for studying the relations among categorical variables and with testing hypotheses about their associations through non-parametric tests and with overall goodness-of-fit statistics.

This chapter begins our study of model-based methods for the analysis of discrete data. These models differ from those we have examined earlier primarily in that they consider *explicitly* an assumed probability distribution for the observations, and make clear distinctions between the systematic component, which is explained by the model, and the random component, which is not. More importantly, the model-based approach allows a compact summary of categorical data in terms of a (hopefully) small number of parameters accompanied by measures of uncertainty (standard errors), and the ability to estimate predicted values over the range of explanatory variables.

This model-fitting approach has several advantages: (a) Inferences for the model parameters include both hypothesis tests and confidence intervals. (b) The former help us to assess which explanatory variables affect the outcome; the size of the estimated parameters and the widths of their confidence intervals help us to assess the strength and importance of these effects. (c) There are a variety of methods for model selection, designed to help determine a favorable trade-off between goodness-of-fit and parsimony. (d) Finally, the predicted values obtained from the model effectively smooth the discrete responses, allow predictions for unobserved values of the explanatory variables, and provide important means to interpret the fitted relationship graphically.

Figure 7.1 provides a visual overview of the steps for fitting and graphing with model-based methods in R. (a) A modeling function such as `glm()` is applied to an input data frame. The result

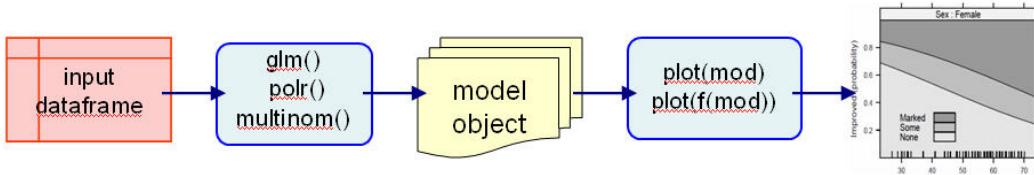


Figure 7.1: Overview of fitting and graphing for model-based methods in R.

{fig:goverview}

is a **model object** containing all the information from the fitting process. (b) As is standard in R, `print()` and `summary()` methods give, respectively, basic and detailed printed output. (c) Many modeling functions have `plot()` methods that produce different types of summary and diagnostic plots. (d) For visualizing the fitted model, most model methods provide a `predict()` method that can be used to plot the fitted values from the model over the ranges of the predictors. Such plots can be customized by the addition of points (showing the observations), lines, confidence bands, and so forth.

In this chapter we consider models for a **binary response**, such as “success” or “failure”, or the number of “successes” in a fixed number of “trials”, where we might reasonably assume a binomial distribution for the random component. These methods extend readily to a **polytomous response** with more than two outcome categories, such as improvement in therapy, with categories “none,” “some” and “marked.”

These models can be seen as simple extensions of familiar ANOVA and regression models for quantitative data. They are also important special cases of a more general approach, the **generalized linear model** that subsumes a wide variety of families of techniques within a single, unified framework. However, rather than starting at the top with the fully general version, this chapter details the important special cases of models for discrete outcomes, beginning with binary responses.

This chapter proceeds as follows: in Section 7.2 we introduce the simple logistic regression model for a binary response and a single quantitative predictor. This model extends directly to models for grouped, binomial data (Section 7.2.4) and to models with any number of regressors (Section 7.3), which can be quantitative, discrete factors and more general forms. For interpreting and understanding the results of a fitted model, we emphasize plotting predicted probabilities and predicted log odds in various ways, for which effect plots (Section 7.3.3) are particularly useful for complex models. Individual observations sometimes exert great influence on a fitted model. Some measures of influence and diagnostic plots are illustrated in Section 7.5. In Section 7.6, we develop several approaches to modelling a multi-category (polytomous) response.

7.2 The logistic regression model

The logistic regression model describes the relationship between a discrete outcome variable, the “response”, and a set of explanatory variables. The response variable is often **dichotomous**, although extensions to the model permit multi-category, **polytomous** outcomes, discussed in Section 7.6. The explanatory variables may be continuous or (with factor variables) discrete.

For a binary response, Y , and a continuous explanatory variable, X , we may be interested in modeling the probability of a successful outcome, which we denote $\pi(x) \equiv \Pr(Y = 1 | X = x)$. That is, at a given value $X = x$, you can imagine that there is a binomial distribution of the responses, $\text{Bin}(\pi(x), n_x)$.

The simplest naive model, called the **linear probability model**, supposes that this probability, $\pi(x)$ varies linearly with the value of x ,

$$\{eq:logit0\} \quad E(Y | x) = \pi(x) = \alpha + \beta x , \quad (7.1)$$

where the notation $E(Y | x)$ indicates that the probability $\pi(x)$ represents the population conditional average of the 1s and 0s for all observations with a fixed value of x . For binary observations, this is simply the proportion of 1s.

Figure 7.2 illustrates the basic setup for modeling a binary outcome using the *Arthritis* data, and described more fully in Example 7.1–Example 7.3. The 0/1 observations are shown as (jittered) points. The predicted values under the linear probability model Eqn. (7.1) are shown as the black line. As you can see, this model cannot be right, because it predicts a probability less than 0 for small values of Age, and would also predict probabilities greater than 1 for larger values of Age.

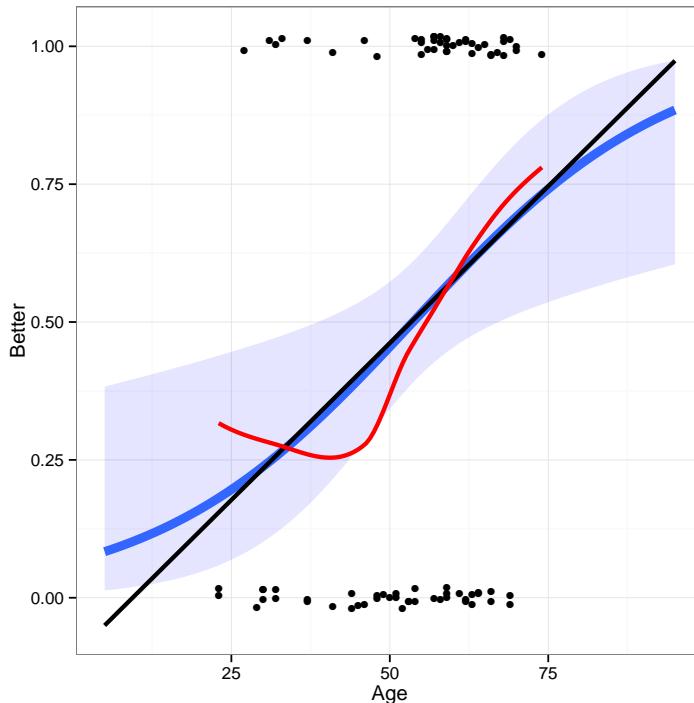


Figure 7.2: Arthritis treatment data, for the relationship of the binary response “Better” to Age. The blue curve and shaded confidence band show a fitted logistic regression to the observations shown as jittered points. The black line shows a simple linear regression and the red curve shows a non-parametric (loess) smoothed curve.

{fig:arthritis-age}

The linear probability model is also wrong because it assumes that the distribution of residuals, $Y_i - \hat{\pi}(x_i)$ is normal, with mean 0 and constant variance. However, because Y is dichotomous, the residuals are also dichotomous, and have variance $\pi(x_i)(1 - \pi(x_i))$, which is maximal for $\pi = 0.5$ and decreases as π goes toward 0 or 1.

One way around the difficulty of needing to constrain the predicted values to the interval $[0, 1]$ is to re-specify the model so that a *transformation* of π has a linear relation to x , and that transformation keeps $\hat{\pi}$ between 0 and 1 for all x . This idea, of modeling a transformation of the response that has desired statistical properties is one of the fundamental ones that led to the development of **generalized linear models**, which we treat more fully later in Chapter 9.

A particularly convenient choice of the transformation gives the **linear logistic regression model**

(or **linear logit model**¹) which posits a linear relation between the **log odds** (or **logit**) of this probability and x ,

$$\text{logit}[\pi(x)] \equiv \log\left(\frac{\pi(x)}{1 - \pi(x)}\right) = \alpha + \beta x . \quad (7.2)$$

When $\beta > 0$, $\pi(x)$ and the log odds increase as X increases; when $\beta < 0$ they decrease with X .

This model can also be expressed as a model for the probabilities $\pi(x)$ in terms of the *inverse* of the logit transformation used in Eqn. (7.2),

$$\pi(x) = \text{logit}^{-1}[\pi(x)] = \frac{1}{1 + \exp[-(\alpha + \beta x)]} \quad (7.3)$$

This transformation uses the cumulative distribution function of the logistic distribution, $\Lambda(p) = \frac{1}{1 + \exp(-p)}$, giving rise to the term *logistic regression*.²

From Eqn. (7.2) we see that the odds of a success response can be expressed as

$$\text{odds}(Y = 1) \equiv \frac{\pi(x)}{1 - \pi(x)} = \exp(\alpha + \beta x) = e^\alpha (e^\beta)^x , \quad (7.4)$$

which is a multiplicative model for the odds. So, under the logistic model,

- β is the change in the log odds associated with a unit increase in x . The odds are multiplied by e^β for each unit increase in x .
- α is log odds at $x = 0$; e^α is the odds of a favorable response at this x -value (which may not have a reasonable interpretation if $X = 0$ is far from the range of the data).

It is easy to explore the relationships among probabilities, odds and log odds using R as we show below, using the function `fractions()` in MASS to print the odds corresponding to probability p as a fraction.

```
> library(MASS)
> p <- c(.05, .10, .25, .50, .75, .90, .95)
> data.frame(p,
+             odds=as.character(fractions(p/(1-p))),
+             logit=log(p/(1-p)))
```

	p	odds	logit
1	0.05	1/19	-2.9444
2	0.10	1/9	-2.1972
3	0.25	1/3	-1.0986
4	0.50	1	0.0000
5	0.75	3	1.0986
6	0.90	9	2.1972
7	0.95	19	2.9444

Thus, a probability of $\pi = 0.25$ represents an odds of 1 to 3, or 1/3, while a probability of $\pi = 0.75$ represents an odds of 3 to 1, or 3. The logits are symmetric around 0, so $\text{logit}(0.25) = -\text{logit}(0.75)$.

Another simple way to interpret the parameter β in the logistic regression model is to consider the relationship between the probability $\pi(x)$ and x . From Eqn. (7.3) it can be shown that the fitted

¹Some writers use the term *logit model* to refer to those using only categorical predictors; we use the terms logistic regression and logit regression interchangeably.

²Any other cumulative probability transformation serves the purpose of constraining the probabilities to the interval [0, 1]. The cumulative normal transformation $\pi(x) = \Phi(\alpha + \beta x)$ gives the **linear probit regression** model. We don't treat probit models here because: (a) The logistic and probit models give results so similar that it is hard to distinguish them in practice; (b) The logistic model is simpler to interpret as a linear model for the log odds or multiplicative model for the odds.

curve (the blue line in Figure 7.2) has slope equal to $\beta\pi(1 - \pi)$. This has a maximum value of $\beta/4$ when $\pi = \frac{1}{2}$, so taking $\beta/4$ gives a quick estimate of the maximum effect of x on the probability scale.

In Figure 7.2 and other plots later in this chapter we try to show the binary responses (as jittered points or a rug plot) to help you appreciate how the fitted logistic curve arises from their distribution across the range a predictor. For didactic purposes this can be seen more readily by plotting the conditional distributions of $x|y = \{0, 1\}$ as a histogram, boxplot or density plot. The function `logi.hist.plot()` in the `probio` package is a nice implementation of this idea (de la Cruz Rot, 2005). The call below produces Figure 7.3, and it is easy to see how increasing age produces a greater probability of a Better response.

```
> with(Arthritis,
+       logi.hist.plot(Age, Better, type="hist", counts=TRUE,
+                      ylabel="Probability (Better)", xlab="Age",
+                      col.cur="blue", col.hist="lightblue", col.box="lightblue")
+     )
```

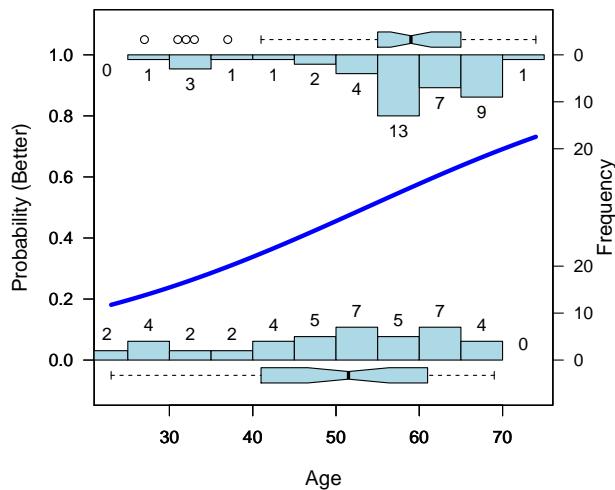


Figure 7.3: Plot of the Arthritis treatment data, showing the conditional distributions of the 0/1 observations of the Better response by histograms and boxplots.
fig:arthritis-logi-hist

7.2.1 Fitting a logistic regression model

Logistic regression models are the special case of generalized linear models fit in R using `glm()` for a binary response using `family=binomial`. We first illustrate how simple models can be fit and interpreted.

```
{sec:logist-fitting}
{ex:arthritis6}
```

EXAMPLE 7.1: Arthritis treatment

In Chapter 4 we examined the data on treatment for rheumatoid arthritis in relation to two categorical predictors, sex of patient and treatment. In addition, the `Arthritis` data gives the age of each patient in this study, and we focus here on the relationship between Age and the outcome, Improved. This response variable has three categories (none, some, or marked improvement), but for now we consider whether the patient showed any improvement at all, defining the event Better to be some or marked improvement.

```
> data("Arthritis", package="vcd")
> Arthritis$Better <- as.numeric(Arthritis$Improved > "None")
```

The logistic regression model is fit using `glm()` as shown below, specifying `family=binomial` for a binary response.

```
> arth.logistic <- glm(Better ~ Age, data=Arthritis, family=binomial)
```

As usual for R modeling functions, the `print()` method for "`glm`" objects gives brief printed output, while the `summary()` method is more verbose, and includes standard errors and hypothesis tests for the model coefficients. To save some space, it is convenient to use the generic function `coeftest()` from the `lmtest` package. Then, we can use this instead of the more detailed `summary()`:

```
> library(lmtest)
> coeftest(arth.logistic)

z test of coefficients:

            Estimate Std. Error z value Pr(>|z|)
(Intercept) -2.6421    1.0732   -2.46   0.014 *
Age          0.0492    0.0194    2.54   0.011 *
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

In the output above, the parameter estimates are $\alpha = -2.642$, and $\beta = 0.0492$. So, the estimated odds of a better response are multiplied by $e^\beta = \exp(0.0492) = 1.05$ for each one year increase in age. Equivalently, you can think of this as a 5% increase per year (using $100(e^\beta - 1)$ to convert). Over 10 years, the odds are multiplied by $\exp(10 \times 0.0492) = 1.64$, a 64% increase, a substantial effect in the range for these data. You can do these calculations in R using the `coef()` method for the "`glm`" object.

```
> exp(coef(arth.logistic))

(Intercept)           Age
 0.071214        1.050482

> exp(10*coef(arth.logistic)[2])

Age
1.6364
```

For comparison with the logistic model, we could fit the linear probability model Eqn. (7.1) using either `lm()` or `glm()` with the default `family=gaussian` argument.

```
> arth.lm <- glm(Better ~ Age, data=Arthritis)
> coef(arth.lm)

(Intercept)           Age
-0.107170        0.011379
```

The coefficient for age can be interpreted to indicate that the probability of a better response increases by 0.011 for each one year increase in age. You can compare this with the $\beta/4$ rule of thumb, that gives $0.0492/4 = 0.0123$. Even though the linear probability model is inappropriate theoretically, you can see in Figure 7.2 (the black line) that it gives similar predicted probabilities to those of the logistic model between age 25–75, where most of the data points are located.



7.2.2 Model tests for simple logistic regression

{sec:logist-tests} There are two main types of hypothesis tests one might want to perform for a logistic regression model. We postpone general discussion of this topic until Section 7.3, but introduce the main ideas here using the analysis of the *Arthritis* data.

- The most basic test answers the question “How much better is the fitted model, $\text{logit}(\pi) = \alpha + \beta x$ than the null model $\text{logit}(\pi) = \alpha$ that includes only the regression intercept?” One answer to this question is given by the (Wald) test of the coefficient for age testing the hypothesis $H_0 : \beta = 0$ that appeared in the output from `summary(arth.logistic)` shown above. The more direct test compares the deviance of the fitted model to the deviance of the null model, and can be obtained using the `anova()` function:

```
> anova(arth.logistic, test="Chisq")

Analysis of Deviance Table

Model: binomial, link: logit

Response: Better

Terms added sequentially (first to last)

Df Deviance Resid. Df Resid. Dev Pr(>Chi)
NULL          83      116
Age     1    7.29      82      109   0.007 **
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

- A second question is “How bad is this model, compared to a model (the **saturated model**) that fits the data perfectly?” This is a test of the size of the residual deviance, that is given by the function `LRstats()` in `vcdExtra`.

```
> LRstats(arth.logistic)

Likelihood summary table:
           AIC BIC LR Chisq Df Pr(>Chisq)
arth.logistic 113 118    109 82      0.024 *
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The summary of these tests is that linear logistic model Eqn. (7.2) fits significantly better than the null model, but that model also shows significant lack of fit.

7.2.3 Plotting a binary response

It is often difficult to understand how a binary response can give rise to a smooth, continuous relation between the predicted response, usually the probability of an event, and a continuous explanatory variable. Beyond this, plots of the data together with fitted models help you to interpret what these models imply.

We illustrate two approaches below using the *Arthritis* data shown in Figure 7.2, first using R base graphics, and then with the `ggplot2` package that makes such graphs somewhat easier to do.

That plot, which was designed for didactic purposes, has the following features:

- It shows the *data*, that is, the 0/1 observations of the `Better` response in relation to age. To do this effectively and avoid over-plotting, the binary responses are jittered.

- It plots the predicted (fitted) logistic regression relationship on the scale of probability, together with a 95% confidence band.
- It also plots the predicted probabilities from the linear probability model.
- A smoothed, non-parametric regression curve for the binary observations is also added to the plot to give some indication of possible non-linearity in the relationship of Better to age.

{ex:arthrit7}

EXAMPLE 7.2: Arthritis treatment: Plotting logistic regression with base graphics

Here we explain how plots similar to Figure 7.2 can be constructed using R base graphics. We describe the steps needed to calculate predicted values and confidence bands and how to add these to a basic plot. These ideas are the basis for the higher-level and more convenient plotting methods illustrated later in this chapter. The steps detailed below give the plot shown in Figure 7.4.

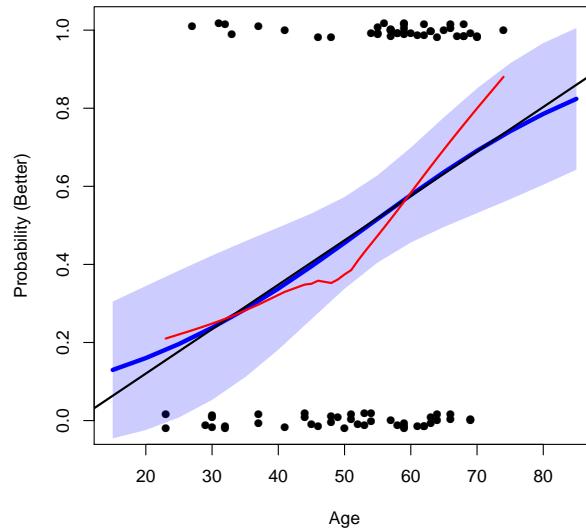


Figure 7.4: A version of plot of the Arthritis treatment data (Figure 7.2) produced with R base graphics, showing logistic, linear regression and lowess fits.^{fig:arthritis-agez}

First, we set up the basic plot of the jittered values of Better vs. Age, setting `xlim` to a larger range than that in the data, only to emphasize where the logistic and linear probability models diverge.

```
> plot(jitter(Better, .1) ~ Age, data=Arthritis,
+       xlim = c(15, 85), pch=16,
+       ylab="Probability (Better)")
```

The fitted logistic curve can be obtained using the `predict()` method for the "glm" object `arth.logistic`. For this example, we wanted to get fitted values for the range of Age from 15–85, which is specified in the `newdata` argument.³ The argument `type="response"` gives fitted values of the probabilities. (The default, `type="link"` would give predicted logits.) Standard errors of the fitted values are not calculated by default, so we set `se.fit=TRUE`.

³Omitting the `newdata` argument would give predicted values using the linear predictors in the data used for the fitted model. Some care needs to be taken if the predictor(s) contain missing values.

```
> xvalues <- seq(15, 85, 5)
> pred.logistic <- predict(arth.logistic,
+                           newdata=data.frame(Age=xvalues),
+                           type="response", se.fit=TRUE)
```

When `se.fit=TRUE`, the `predict()` function returns its result in a list, with components `fit` for the fitted values and `se.fit` for the standard errors. From these, we can calculate 95% pointwise prediction intervals using the standard normal approximation.

```
> upper <- pred.logistic$fit + 1.96 * pred.logistic$se.fit
> lower <- pred.logistic$fit - 1.96 * pred.logistic$se.fit
```

We can then plot the confidence band using `polygon()` and the fitted logistic curve using `lines()`. A graphics trick is used here to use a transparent color for the confidence band using `rgb(r, g, b, alpha)`, where `alpha` is the transparency value.

```
> polygon(c(xvalues, rev(xvalues)),
+          c(upper, rev(lower)),
+          col=rgb(0, 0, 1, .2), border=NA)
> lines(xvalues, pred.logistic$fit, lwd=4, col="blue")
```

This method, using `predict()` for calculations and `polygon()` and `lines()` for plotting can be used to display the predicted relationships and confidence bands under other models. Here, we simply used `abline()` to plot the fitted line for the linear probability model `arth.lm` and `lowess()` to calculate a smoothed, non-parametric curve.

```
> abline(arth.lm, lwd=2)
> lines(lowess(Arthritis$Age, Arthritis$Better, f=.9), col="red", lwd=2)
```



{ex:arthrit8}

EXAMPLE 7.3: Arthritis treatment: Plotting logistic regression with ggplot2

Model-based plots such as Figure 7.2 are relatively more straight-forward to produce using `ggplot2`. The basic steps here are to:

- set up the plot frame with `ggplot()` using Age and Better as (x, y) coordinates;
- use `geom_point()` to plot the observations, whose positions are jittered with `position_jitter()`;
- use `stat_smooth()` with `method = "glm"` and `family = binomial` to plot the predicted probability curve and confidence band. By default, `stat_smooth()` calculates and plots 95% confidence bands on the response (probability) scale.

```
> library(ggplot2)
> # basic logistic regression plot
> gg <- ggplot(Arthritis, aes(x=Age, y=Better)) +
+   xlim(5, 95) + theme_bw() +
+   geom_point(position = position_jitter(height = 0.02, width = 0)) +
+   stat_smooth(method = "glm", family = binomial, alpha = 0.1, fill="blue",
+               size=2.5, fullrange=TRUE)
```

Finally, we can add other smoothers to the plot, literally by using `+` to add these to the "ggplot" object.

```
> # add linear model and loess smoothers
> gg <- gg + stat_smooth(method = "lm", se=FALSE,
+                         size=1.2, color="black", fullrange=TRUE)
> gg <- gg + stat_smooth(method = "loess", se=FALSE,
+                         span=0.95, colour="red", size=1.2)
> gg # show the plot
```



7.2.4 Grouped binomial data

A related case occurs with grouped data, where rather than binary observations, $y_i \in \{0, 1\}$ in case form, the data is given in what is called *events/trials form* that records the number of successes, y_i that occurred in n_i trials associated with each setting of the explanatory variable(s) x_i .⁴ Case form, with binary observations is the special case where $n_i = 1$.

Data in events/trials form often arises from contingency table data with a binary response. For example in the *UCBAdmissions* data, the response variable `Admit` with levels "Admitted", "Rejected" could be treated in this way using the number of applicants as the number of trials.

As before, we can consider y_i/n_i to estimate the probability of success, π_i and the distribution of Y to be binomial, $\text{Bin}(\pi_i, n_i)$ at each x_i .

In practical applications, there are two main differences between the cases of ungrouped, case form data and grouped, event/trials form.

- In fitting models using `glm()`, the model formula, `response ~ terms`, can be given using a response consisting of a two-column matrix, whose columns contain the numbers of successes y_i and failures $n_i - y_i$. Alternatively, the response can be given as the proportion of successes, y_i/n_i , but then it is necessary to specify the number of trials as a weight.
- In plotting the fitted model on the scale of probability, you usually have to explicitly plot the fraction of successes, y_i/n_i .

{ex:nasa-temp}

EXAMPLE 7.4: Space shuttle disaster

In Example 1.2 and Example 1.10 we described the background behind the post-mortem examination of the evidence relating to the disastrous launch of the space shuttle *Challenger* on January 28, 1986. Here we consider a simple, but proper analysis of the data available at the time of launch. We also use this example to illustrate some details of the fitting and plotting of grouped binomial data. As well, we describe some of the possibilities for dealing with missing data.

The data set *SpaceShuttle* in *vcd* contains data on the failures of the O-rings in 24 NASA launches preceding the launch of *Challenger*, as given by Dalal *et al.* (1989) and Tufte (1997) also analysed by Lavine (1991).

Each launch used two booster rockets with a total of six O-rings, and the data set records as `nFailures` the number of these that were considered damaged after the rockets were recovered at sea. In one launch (flight # 4), the rocket was lost at sea, so the relevant response variables are missing.

In this example, we focus on the variable `nFailures` as a binomial with $n_i = 6$ trials. The missing data for flight 4 can be handled in several ways in the call to `glm()`

```
> data("SpaceShuttle", package="vcd")
> shuttle.mod <- glm(cbind(nFailures, 6 - nFailures) ~ Temperature,
+   data = SpaceShuttle, na.action = na.exclude,
+   family = binomial)
```

Alternatively, we can add an explicit `trials` variable, represent the response as the proportion `nFailures/trials`, and use `weight = trials` to indicate the total number of observations.

```
> SpaceShuttle$trials <- 6
> shuttle.modw <- glm(nFailures/trials ~ Temperature, weight = trials,
+   data = SpaceShuttle, na.action = na.exclude,
+   family = binomial)
```

⁴Alternatively, the data may record the number of successes, y_i , and number of failures, $n_i - y_i$.

{sec:logist-grouped}

These two approaches give identical results for all practical purposes:

```
> all.equal(coef(shuttle.mod), coef(shuttle.modw))
[1] TRUE
```

As before, we can test whether temperature significantly improves prediction of failure probability using `anova()`:

```
> # testing, vs. null model
> anova(shuttle.mod, test="Chisq")

Analysis of Deviance Table

Model: binomial, link: logit

Response: cbind(nFailures, 6 - nFailures)

Terms added sequentially (first to last)

          Df Deviance Resid. Df Resid. Dev Pr(>Chi)
NULL           22      24.2
Temperature    1       6.14      21      18.1   0.013 *
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The code below gives a `ggplot2` version in Figure 7.5 of the plot we showed earlier in Example 1.2 (Figure 1.2). The relevant details here are:

- We specify `y = nFailures / trials` to calculate the failure probabilities.
- Points are jittered in the call to `geom_point()` to prevent overplotting.
- In the call to `geom_smooth()`, we need to use `weight = trials`, just as in the call to `glm()` above.
- `fullrange = TRUE` makes the fitted regression curve and confidence band extend across the entire plot

```
> library(ggplot2)
> ggplot(SpaceShuttle, aes(x = Temperature, y = nFailures / trials)) +
+   xlim(30, 81) + theme_bw() +
+   xlab("Temperature (F)") +
+   ylab("O-Ring Failure Probability") +
+   geom_point(position=position_jitter(width=0, height=0.01),
+             aes(size = 2)) +
+   theme(legend.position="none") +
+   geom_smooth(method = "glm", family = binomial, fill="blue",
+             aes(weight = trials), fullrange = TRUE, alpha=0.2, size=2)
```



7.3 Multiple logistic regression models

As is the case in classical regression, generalizing the simple logistic regression to an arbitrary number of explanatory variables is quite straightforward. We let $\boldsymbol{x}_i = (x_{i1}, x_{i2}, \dots, x_{ip})$ denote the vector of p explanatory variables for case or cluster i . Then the general logistic regression model can be expressed as

$$\begin{aligned} \text{logit}(\pi_i) \equiv \log \frac{\pi_i}{1 - \pi_i} &= \alpha + \boldsymbol{x}_i^\top \boldsymbol{\beta} \\ &= \alpha + \beta_1 x_{i1} + \beta_2 x_{i2} + \cdots + \beta_p x_{ip}. \end{aligned} \tag{7.5}$$

{sec:logist-mult}

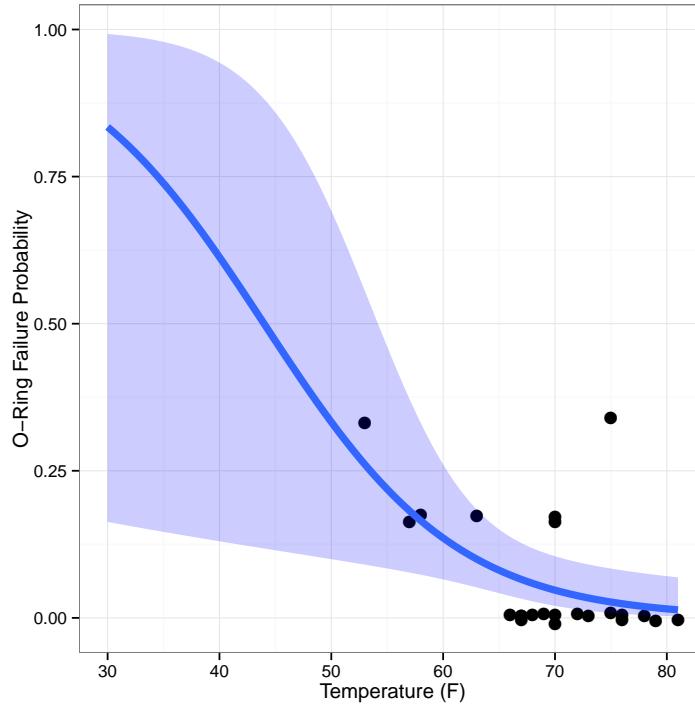


Figure 7.5: Space shuttle data, with fitted logistic regression model

{fig:nasa-temp-ggplot}

Equivalently, we can represent this model in terms of probabilities as the logistic transformation of the **linear predictor**, $\eta_i = \alpha + \mathbf{x}_i^\top \boldsymbol{\beta}$,

$$\begin{aligned}\pi_i = \Lambda(\eta_i) &= \Lambda(\alpha + \mathbf{x}_i^\top \boldsymbol{\beta}) \\ &= \frac{1}{1 + \exp(\alpha + \beta_1 x_{i1} + \beta_2 x_{i2} + \cdots + \beta_p x_{ip})}.\end{aligned}\quad (7.6) \quad \text{(eq:logistm1)}$$

The xs can include any of the following sorts of regressors, as in the general linear model:

- **quantitative** variables (e.g., age, income)
- **polynomial** powers of quantitative variables (e.g., age, age^2 , age^3)
- **transformations** of quantitative variables (e.g., log salary)
- factors, represented as **dummy** variables for qualitative predictors (e.g., P_1, P_2, P_3 for four political party affiliations)
- **interaction** terms (e.g., $sex \times age$, or $age \times income$)

{ex:arthrit-mult}

EXAMPLE 7.5: Arthritis treatment

We continue with the analysis of the *Arthritis* data, fitting a model containing the main effects of Age, Sex and Treatment, with Better as the response. This model has the form

$$\text{logit}(\pi_i) = \alpha + \beta_1 x_{i1} + \beta_2 x_{i2} + \beta_3 x_{i3}$$

where x_1 is Age and x_2 and x_3 are the factors representing Sex and Treatment, respectively. Using the default (0/1) dummy coding that R uses (“treatment” contrasts against the lowest factor

level),⁵ they are defined as:

$$x_2 = \begin{cases} 0 & \text{if Female} \\ 1 & \text{if Male} \end{cases} \quad x_3 = \begin{cases} 0 & \text{if Placebo} \\ 1 & \text{if Treatment} \end{cases}$$

In this model,

- α doesn't have a sensible interpretation here, but formally it would be the log odds of improvement for a person at age $x_1 = 0$ in the baseline or reference group with $x_2 = 0$ and $x_3 = 0$ —females receiving the placebo. To make the intercept interpretable, we will fit the model centering age near the mean, by using $x_1 - 50$ as the first regressor.
- β_1 is the increment in log odds of improvement for each one-year increase in age.
- β_2 is the increment in log odds for male as compared to female. Therefore, e^{β_2} gives the odds of improvement for males relative to females.
- β_3 is the increment in log odds for being in the treated group. e^{β_3} gives the odds of improvement for the active treatment group relative to placebo.

We fit the model as follows. In `glm()` model formulas, “–” has a special meaning, so we use the identity function, `I(Age-50)` to center age.

```
> arth.logistic2 <- glm(Better ~ I(Age-50) + Sex + Treatment,
+                         data=Arthritis,
+                         family=binomial)
```

The parameters defined here are *incremental effects*. The intercept corresponds to a baseline group (50 year-old females given the placebo); the other parameters are incremental effects for the other groups compared to the baseline group. Thus, when α , β_1 , β_2 and β_3 have been estimated, the fitted logits and predicted odds at Age==50 are:

Sex	Treatment	Logit	Odds Improved
Female	Placebo	α	e^α
Female	Treated	$\alpha + \beta_2$	$e^{\alpha+\beta_2}$
Male	Placebo	$\alpha + \beta_1$	$e^{\alpha+\beta_1}$
Male	Treated	$\alpha + \beta_1 + \beta_2$	$e^{\alpha+\beta_1+\beta_2}$

We first focus on the interpretation of the coefficients estimated for this model shown below.

```
> coeftest(arth.logistic2)

z test of coefficients:

            Estimate Std. Error z value Pr(>|z|)
(Intercept) -0.5781    0.3674  -1.57   0.116
I(Age - 50)  0.0487    0.0207   2.36   0.018 *
SexMale     -1.4878    0.5948  -2.50   0.012 *
TreatmentTreated 1.7598    0.5365   3.28   0.001 **

Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

⁵For factor variables with the default treatment contrasts, you can change the reference level using `relevel()`. In this example, you could make male the baseline category using `Arthritis$Sex <- relevel(Arthritis$Sex, ref = "Male")`.

To interpret these in terms of odds ratios and also find confidence intervals, just use `exp()` and `confint()`.

```
> exp(cbind(OddsRatio=coef(arth.logistic2),
+             confint(arth.logistic2)))

```

	OddsRatio	2.5 %	97.5 %
(Intercept)	0.5609	0.26475	1.1323
I(Age - 50)	1.0500	1.01000	1.0963
SexMale	0.2259	0.06524	0.6891
TreatmentTreated	5.8113	2.11870	17.7266

Here,

- $\alpha = -0.578$: At age 50, females given the placebo have an odds of improvement of $\exp(-0.578) = 0.56$.
- $\beta_1 = 0.0487$: Each year of age multiplies the odds of improvement by $\exp(0.0487) = 1.05$, or a 5% increase.
- $\beta_2 = -1.49$: Males are only $\exp(-1.49) = 0.26$ times as likely to show improvement relative to females. Equivalently, you could say that females are $\exp(1.49) = 4.437$ times more likely than males to improve.
- $\beta_3 = 1.76$: People given the active treatment are $\exp(1.76) = 5.8$ times more likely to show improvement.

As you can see, the interpretation of coefficients in multiple logistic models is straightforward, though a bit cumbersome. This becomes more difficult in larger models, particularly when there are interactions. In these cases, you can understand (and explain) a fitted model more easily through plots of predicted values, either on the scale of response probability or on the logit scale of the linear predictor. We describe these methods in Section 7.3.1–Section 7.3.3 below.



7.3.1 Conditional plots

The simplest kind of plots display the data together with a representation of the fitted relationship (predicted values, confidence bands) separately for subsets of the data defined by one or more of the predictors. Such plots can show the predicted values for the response variable on the ordinate against one chosen predictor on the abscissa, and can use multiple curves and multiple panels to represent other predictors.

However, these plots are *conditional plots*, meaning that the data shown in each panel and used in each fitted curve are limited to the subset of the observations defined by the curve and panel variables. As well, predictors that are not shown in a given plot are effectively ignored (or marginalized), as was the case in Figure 7.2 that showed only the effect of age in the *Arthritis* data.

EXAMPLE 7.6: Arthritis treatment: conditional plots

For the *Arthritis* data, a basic conditional plot of `Better` vs. `Age`, showing the observations as jittered points (with `geom_point()`) and the fitted logistic curves (with `stat_smooth()` using `method="glm"`) can be produced with `ggplot2` as shown below, giving Figure 7.6.

```
> library(ggplot2)
> gg <- ggplot(Arthritis, aes(Age, Better, color=Treatment)) +
+   xlim(5, 95) + theme_bw() +
+   geom_point(position = position_jitter(height = 0.02, width = 0)) +
+   stat_smooth(method = "glm", family = binomial, alpha = 0.2,
+             aes(fill=Treatment), size=2.5, fullrange=TRUE)
> gg    # show the plot
```

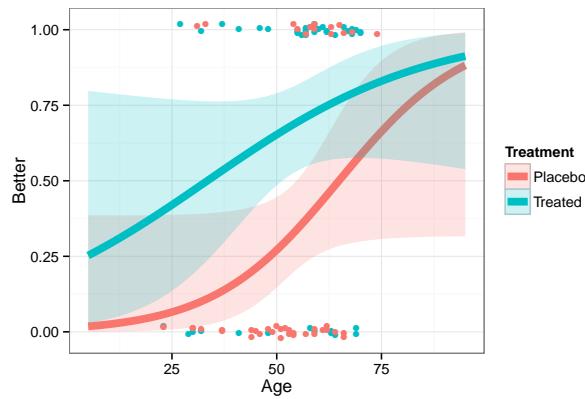


Figure 7.6: Conditional plot of Arthritis data showing separate points and fitted curves stratified by Treatment. A separate fitted curve is shown for the two treatment conditions, ignoring Sex.^{fig:arth-cond1}

In this call to `ggplot()`, specifying `color=Treatment` gives different point and line colors, but also automatically stratifies the fitted curves using the levels of this variable.

With such a plot, it is easy to add further stratifying variables in the data using `facets` to produce separate panels (functions `facet_wrap()` or `facet_grid()`, with different options to control the details). The following line further stratifies by Sex, producing Figure 7.7.

```
> gg + facet_wrap(~ Sex)
```

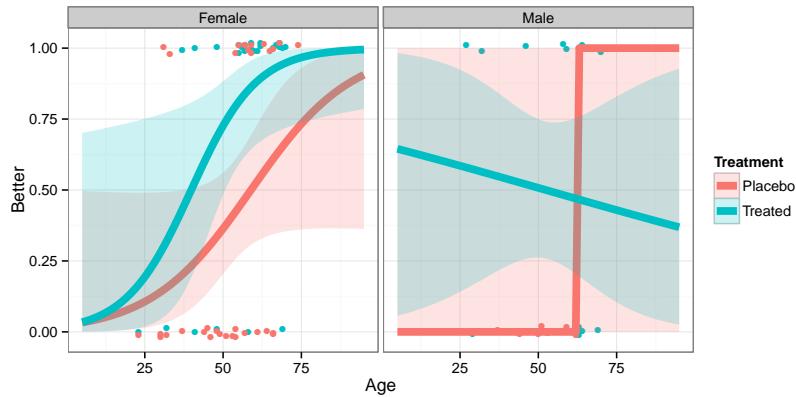


Figure 7.7: Conditional plot of Arthritis data, stratified by Treatment and Sex. The unusual patterns in the panel for Males signals a problem with this data.^{fig:arth-cond2}

However, you can see from this plot how this method breaks down when the sample size is small in some of the groups defined by the stratifying factors. The panel for males shows a paradoxical negative relation with age for the treated group and a step function for the placebo group. The explanation for this is shown in the two-way frequency table of the sex and treatment combinations:

```
> addmargins(xtabs(~Sex + Treatment, data=Arthritis), 2)
Treatment
```

Sex	Placebo	Treated	Sum
Female	32	27	59
Male	11	14	25

Less than 1/3 of the sample were males, and of these only 11 were in the placebo group. `glm()` cannot estimate the fitted relationship against `Age` here—the slope coefficient is infinite, and the fitted probabilities are all either 0 or 1.⁶

△

7.3.2 Full-model plots

c:list=fullplots}
 For a model with two or more explanatory variables, *full-model plots* display the fitted response surface for all predictors together, rather than stratified by conditioning variables. Such plots show the predicted values for the response variable on the ordinate against one chosen predictor on the abscissa, and can use multiple curves and multiple panels to represent other predictors.

A simple R trick⁷ makes this method far easier and more general than the naive plotting method used in Example 7.2. The trick is simply to combine the columns in the original data frame with the result of the `predict()` method for the fitted model and plot the calculated `fit` value, together with confidence bands (if you use `se.fit=TRUE`).

EXAMPLE 7.7: Arthritis treatment: full-model plots

```
> arth.fit2 <- cbind(Arthritis,
+                      predict(arth.logistic2, se.fit = TRUE))
```

The fitted values here are on the logit scale, which means that it takes one more trick to show the data points on the plot. We simply define a new variable, `obs` with convenient logit values corresponding the `Better` values of 0 and 1.

```
> arth.fit2$obs <- c(-4, 4)[1+arth.fit2$Better]
```

We can then plot the fitted logit against `Age` using `x=Age`, `y=fit` from the data frame containing the fitted values. The call to `ggplot()` below produces Figure 7.8. Here, we used `color=Treatment` to produce separate points, lines and confidence bands colored by `Treatment`. Confidence bands in the plot are constructed using `geom_ribbon()`.

```
> ggplot(arth.fit2, aes(x=Age, y=fit, color=Treatment)) +
+   geom_line(size = 2) + theme_bw() +
+   geom_ribbon(aes(ymax = fit - 1.96 * se.fit,
+                  ymax = fit + 1.96 * se.fit,
+                  fill = Treatment), alpha = 0.2,
+                  color = "transparent") +
+   labs(x = "Age", y = "Log odds (Better)") +
+   geom_point(aes(y=obs),
+              position=position_jitter(height=0.25, width=0)) +
+   facet_wrap(~ Sex)
```

This plot method has several nice features:

- Plotting on the logit scale shows the additive, linear effects of all predictors.
- It provides a visual representation of the information contained in the table of coefficients. Note, however, that the choice to display `Treatment` within each panel makes it easier to judge the size of this effect, compared to the effect of `Sex` which must be judged across the panels.

⁶This is called **complete separation**, and occurs whenever the responses have no overlap on the predictor variable(s) used in fitting the logistic regression model.

⁷Thanks to Dennis Murphy for suggesting this method.

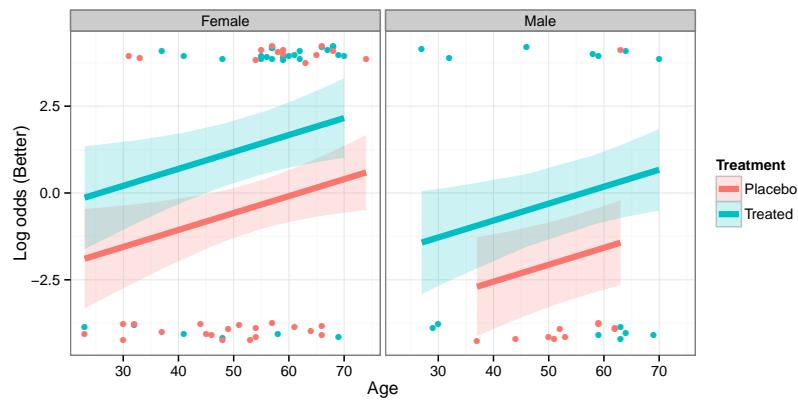


Figure 7.8: Full-model plot of Arthritis data, showing fitted logits by Treatment and Sex.^{fig:arth-full1}

- It shows the data as points, and the fitted lines and confidence bands are restricted to the range of the data in each. You can easily see the reason for the unusual pattern in the conditional plot for Males shown in Figure 7.7.
- It generalizes directly to any fitted model, because the same plotting code can be used once the model predicted values have been calculated.
- Additional predictors, either factors or quantitative variables can easily be accommodated by including them in the `facet_wrap()` call. For example, if the patients were also categorized by education and this had been included in the model, `facet_wrap(~ Sex + Education)` would produce separate panels for the combinations of these two variables.

While plots on the logit scale have a simpler form, many people find it easier to think about such relationships in terms of probabilities, as we have done in earlier plots in this chapter. You can do the same for full-model plots with a simple extension of this method. All you need to do is to transform the `fit` and end points of the confidence band back to the scale of probabilities. The function `plogis()` does this for the logistic distribution.

```
> arth.fit2 <- within(arth.fit2, {
+   prob   <- plogis(fit)
+   lower  <- plogis(fit - 1.96 * se.fit)
+   upper  <- plogis(fit + 1.96 * se.fit)
+ })
```

The plot step is then similar to what we used above (but with `prob`, `lower` and `upper`), producing Figure 7.9.

```
> ggplot( arth.fit2, aes(x=Age, y=prob, color=Treatment)) +
+   geom_line(size = 2) + theme_bw() +
+   geom_ribbon(aes(ymin = lower,
+                  ymax = upper,
+                  fill = Treatment), alpha = 0.2,
+                  color = "transparent") +
+   labs(x = "Age", y = "Probability (Better)") +
+   geom_point(aes(y=Better),
+              position=position_jitter(height=0.02, width=0)) +
+   facet_wrap(~ Sex)
```



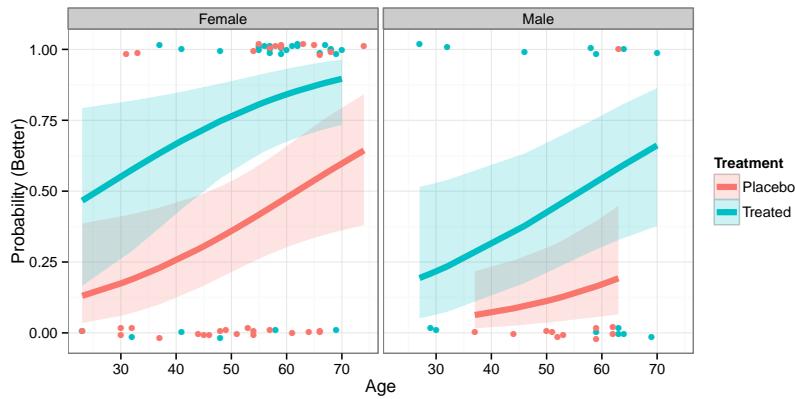


Figure 7.9: Full-model plot of Arthritis data, showing fitted probabilities by Treatment and Sex.^{fig:arth-full2}

{sec:logist-effplots}

7.3.3 Effect plots

For more than two variables, full-model plots of the fitted response surface can be cumbersome, particularly when the model contains interactions or when the main substantive interest is focused on a given main effect or interaction, controlling for all other explanatory variables. The method of *effect displays* (tables and graphs), developed by John Fox (1987, 2003) and implemented in the *effects* package, is a useful solution to these problems.

The idea of effect plots is quite simple but very general:⁸ consider a particular subset of predictors (*focal predictors*) we wish to visualize in a given linear model or generalized linear model. The essence is to calculate fitted values (and standard errors) for the model terms involving these variables and all low-order relatives (e.g., main effects that are marginal to an interaction), as these variables are allowed to vary over their range. All other variables are “controlled” by being fixed at typical values. For example a quantitative covariate could be fixed at its mean or median; a factor could be fixed at equal proportions of its levels or its proportions in the data. The result, when plotted, shows all effects of the focal predictors and their low-order relatives, but with all other variables not included controlled or adjusted for.

More formally, assume we have fit a model with a linear predictor $\eta_i = \alpha + \mathbf{x}_i^\top \boldsymbol{\beta}$ (on the logit scale, for logistic regression). Letting $\beta_0 = \alpha$ and $\mathbf{x}_0 = \mathbf{1}$, we can rewrite this in matrix form as $\boldsymbol{\eta} = \mathbf{X}\boldsymbol{\beta}$ where \mathbf{X} is the model matrix constructed by the modeling function, such as `glm()`. Fitting the model gives the estimated coefficients $\hat{\boldsymbol{b}}$ and its estimated covariance matrix $\hat{\mathcal{V}}(\hat{\boldsymbol{b}})$.

The `Effect()` function constructs an analogous *score model matrix*, \mathbf{X}^* , where the focal variables have been varied over their range, and all other variables represented as constant, typical values. Using this as input (the `newdata` argument) to the `predict()` function then gives the fitted values $\boldsymbol{\eta}^* = \mathbf{X}^*\hat{\boldsymbol{b}}$. Standard errors used for confidence intervals are calculated by `predict()` (when `se.fit=TRUE`) as the square roots of $\text{diag}(\mathbf{X}^*\hat{\mathcal{V}}(\hat{\boldsymbol{b}})\mathbf{X}^{*\top})$. Note that these ideas work not only for `glm()` models, but potentially for any modeling function that has a `predict()` and `vcov()` method.⁹

These results are calculated on the scale of the linear predictor $\boldsymbol{\eta}$ (logits, for logistic regression)

⁸Less general expression of these ideas include the use of *adjusted means* in analysis of covariance, and *least squares means* or *population marginal means* (Searle *et al.*, 1980) in analysis of variance; for example, see the `lsmeans` package for classical linear models.

⁹For example, the `effects` package presently provides methods for models fit by `lm()` (including multivariate linear response models), `glm()`, `gls()`, multinomial (`multinom()` in the `nmf` package) and proportional odds models (`polr()` in `MASS`), polytomous latent class models (`polCA` package), as well as a variety of multi-level and mixed-effects linear models fit with `lmer()` from the `lme4` package, or with `lme()` from the `nlme` package.

when the `type` argument to `predict()` is `type="link"` or on the response scale (probabilities, here) when `type="response"`. The latter makes use of the inverse transformation, Eqn. (7.6).

There are two main calculation functions in the `effects` package:

- `Effect()` takes a character vector of the names of a subset of focal predictors and constructs the score matrix X^* by varying these over their ranges, while holding all other predictors constant at “typical” values. There are many options that control these calculations. For example, `xlevels` can be used to specify the values of the focal predictors; `typical` or `given.values` respectively can be used to specify either a function (`mean`, `median`) or a list of specific typical values used for the variables that are controlled. The result is an object of class “`eff`”, for which there are `print()`, `summary()` and (most importantly) `plot()` methods. See `help(Effect)` for a complete description.
- `allEffects()` takes a model object, and calculates the effects for each high-order term in the model (including their low-order) relatives. Similar optional arguments control the details of the computation. The result is an object of class “`efflist`”.

In addition, the plotting methods for “`eff`” and “`efflist`” objects offer numerous options to control the plot details, only a few of which are used in the examples below. For logistic regression models, they also solve the problem of the trade-off between plots on the logit scale, that have a simple representation in terms of additive effects, and plots on the probability scale that are usually simpler to understand. By default, the fitted model effects are plotted on the logit scale, but the response y axis is labeled with the corresponding probability values.

{ex:arthrit-eff}

EXAMPLE 7.8: Arthritis treatment

Here we illustrate the use of the `effects` package with the simple main effects model which was fit in Example 7.5. `allEffects()` is used to calculate the predicted probabilities of the Better response for Age and the two factors, Sex and Treatment.

```
> library(effects)

Error: package 'effects' was built before R 3.0.0: please re-install it
> arth.eff2 <- allEffects(arth.logistic2)

Error in eval(expr, envir, enclos): could not find function "allEffects"
> names(arth.eff2)

Error in eval(expr, envir, enclos): object 'arth.eff2' not found
```

The result, `arth.eff2` is a list containing the fitted values (response probabilities, by default) for each of the model terms. No `xlevels` argument was specified, so by default the function calculated the effects for Age at a reasonable selection of equally-spaced values:

```
> arth.eff2[[1]]

Error in eval(expr, envir, enclos): object 'arth.eff2' not found
```

The default plot method for the “`efflist`” object produces one plot for each high-order term, which are just the main effect in this model. The call below produces Figure ??.

```
> plot(arth.eff2, rows=1, cols=3)

Error in plot(arth.eff2, rows = 1, cols = 3): object 'arth.eff2' not found
```

You can also produce full-model plots quite easily by using all predictors in the model in a call to `Effect()`.

```
> arth.full <- Effect(c("Age", "Treatment", "Sex"), arth.logistic2)
Error in eval(expr, envir, enclos): could not find function "Effect"
```

Then plotting the result, with some options, gives the plot shown in Figure ??.

```
> plot(arth.full, multiline=TRUE, ci.style="bands",
+       colors = c("red", "blue"), lwd=3,
+       ticks=list(at=c(.05, .1, .25, .5, .75, .9, .95)),
+       key.args=list(x=.52, y=.92), grid=TRUE)

Error in plot(arth.full, multiline = TRUE, ci.style = "bands", colors = c("red",
: object 'arth.full' not found
```

Alternatively, we can plot these results directly on the scale of probabilities, as shown in Figure ??.

```
> plot(arth.full, multiline=TRUE, ci.style="bands", rescale.axis=FALSE,
+       colors = c("red", "blue"), lwd=3,
+       key.args=list(x=.52, y=.92), grid=TRUE)

Error in plot(arth.full, multiline = TRUE, ci.style = "bands", rescale.axis =
FALSE, : object 'arth.full' not found
```



7.4 Case studies

The examples below take up some issues of data analysis, model building and visualization in the context of multiple logistic regression models. We focus on the combination of exploratory plots to see the data, modeling steps and graphs to interpret a given model.

EXAMPLE 7.9: Donner Party

In Chapter 1, Example 1.3, we described the background behind the sad story of the Donner Party, perhaps the most famous tragedy in the history of the westward settlement in the United States. In brief, the party was stranded on the eastern side of the Sierra Nevada mountains by heavy snow in late October, 1846, and by the time the last survivor was rescued in April, 1847, nearly half of the members had died from famine and exposure to extreme cold. Figure 1.3 showed that survival decreased strongly with age.

Here we consider a more detailed analysis of these data, which are contained in the data set `Donner` in `vcdExtra`. This data set lists 90 people in the Donner Party by name, together with age, sex, survived (0/1) and the date of death for those who died.¹⁰

```
> data("Donner", package="vcdExtra")    # load the data
> library(car)                      # for some() and Anova()
> some(Donner, 8)
```

¹⁰Most historical sources count the number in the Donner Party at 87 or 89. An exact accounting of the members of the Donner Party is difficult, because: (a) several people joined the party in mid-route, at Fort Bridger and in the Wasatch Mountains; (b) several rode ahead to search for supplies and one (Charles Stanton) brought two more with him (Luis and Salvador); (c) five people died before reaching the Sierra Nevada mountains. It incorporates updated information from Kristin Johnson's listing, <http://user.xmission.com/~octa/DonnerParty/Roster.htm>.

	family	age	sex	survived	death
Breen, Peter	Breen	3	Male	1	<NA>
Donner, Jacob	Donner	65	Male	0	1846-12-21
Foster, Jeremiah	MurFosPik	1	Male	0	1847-03-13
Graves, Nancy	Graves	9	Female	1	<NA>
McCutchen, Harriet	McCutchens	1	Female	0	1847-02-02
Reed, James	Reed	46	Male	1	<NA>
Reinhardt, Joseph	Other	30	Male	0	1846-12-21
Wolfinger, Doris	FosdWolf	20	Female	1	<NA>

The main purpose of this example is to try to understand, through graphs and models, how survival was related to age and sex. However, first, we do some data preparation and exploration. The response variable, `survived` is a 0/1 integer, and it is more convenient for some purposes to make it a factor.

```
> Donner$survived <- factor(Donner$survived, labels=c("no", "yes"))
```

Some historical accounts (Grayson, 1990) link survival in the Donner Party to kinship or family groups, so we take a quick look at this factor here. The variable `family` reflects a recoding of the last names of individuals to reduce the number of factor levels. The main families in the Donner party were: Donner, Graves, Breen and Reed. The families of Murphy, Foster and Pike are grouped as "MurFosPik", those of Fosdick and Wolfinger are coded as "FosdWolf", and all others as "Other".

```
> xtabs(~family, data=Donner)

family
Breen     Donner      Eddy   FosdWolf      Graves   Keseberg
    9        14          4        4          10          4
McCutchen MurFosPik   Other      Reed
    3        12         23        7
```

For the present purposes, we reduce these 10 family groups further, collapsing some of the small families into "Other", and reordering the levels. Assigning new values to the `levels()` of a factor is a convenient trick for recoding factor variables.

```
> # collapse small families into "Other"
> fam <- Donner$family
> levels(fam)[c(3,4,6,7,9)] <- "Other"
>
> # reorder, putting Other last
> fam = factor(fam, levels(fam)[c(1, 2, 4:6, 3)])
> Donner$family <- fam
> xtabs(~family, data=Donner)

family
Breen     Donner      Graves   MurFosPik      Reed      Other
    9        14          10         12          7         38
```

`xtabs()` then shows the counts of survival by these family groups:

```
> xtabs(~survived+family, data=Donner)

family
survived Breen Donner Graves MurFosPik Reed Other
  no      0     7     3       6     1    25
  yes     9     7     7       6     6    13
```

Plotting this distribution of survival by family with a formula gives a *spineplot*, a special case

of the mosaic plot, or a generalization of a stacked bar plot, shown in Figure 7.10. The widths of the bars are proportional to family size, and the shading highlights in light blue the proportion who survived in each family.

```
> plot(survived ~ family, data=Donner, col=c("pink", "lightblue"))
```

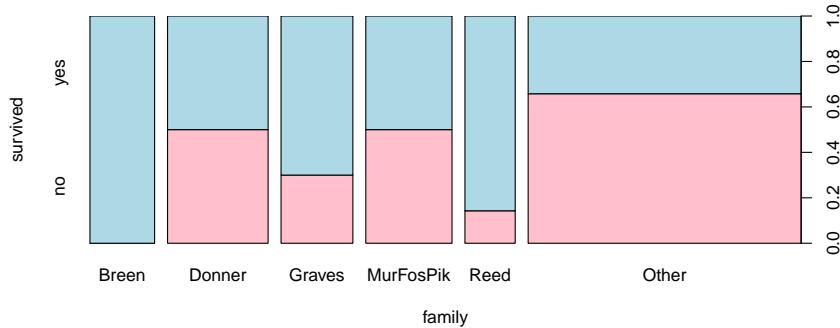


Figure 7.10: Spineplot of survival in the Donner Party by family.^{fig:donner1-spineplot}

A generalized pairs plot (Section 5.5.1), shown in Figure ?? gives a visual overview of the data. The diagonal panels here show the marginal distributions of the variables as bar plots, and highlight the skewed distribution of age and the greater number of males than females in the party. The boxplots and barcode plots for survived and age show that those who survived were generally younger than those who perished.

```
> library(gpairs)

Error in library(gpairs): there is no package called 'gpairs'

> library(vcd)
> gpairs(Donner[,c(4,2,3,1)],
+   diag.pars=list(fontsize=20, hist.color="gray"),
+   mosaic.pars=list(gp=shading_Friendly), outer.rot=c(45,45)
+ )

Error in eval(expr, envir, enclos): could not find function "gpairs"
```

From an exploratory perspective, we now proceed to examine the relationship of survival to age and sex, beginning with the kind of conditional plots we illustrated earlier (in Example 7.6). Figure 7.11 shows a plot of survived, converted back to a 0/1 variable as required by `ggplot()`, together with the binary responses as points and the fitted logistic regressions separately for males and females.

```
> ggplot(Donner, aes(age, as.numeric(survived=="yes"), color = sex)) +
+   theme_bw() + ylab("Survived") +
+   geom_point(position = position_jitter(height = 0.02, width = 0)) +
+   stat_smooth(method = "glm", family = binomial, formula = y ~ x,
+               alpha = 0.2, size=2, aes(fill = sex))
```

It is easy to see that survival among women was greater than for men, perhaps narrowing the gap among the older people, but the data gets thin towards the upper range of age.

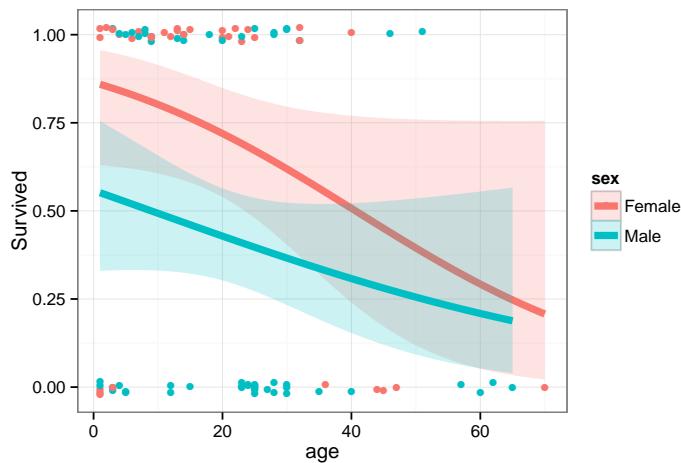


Figure 7.11: Conditional plot of the Donner data, showing the relationship of survival to age and sex. The smoothed curves and confidence bands show the result of fitting separate linear logistic regressions on age for males and females.¹¹

The curves plotted in Figure 7.11 assume a linear relationship between the log odds of survival and age (expressed as `formula = y ~ x` in the call to `stat_smooth()`). One simple way to check whether the relationship between survival and age is non-linear is to re-do this plot, but now allow a quadratic relationship with age, using `formula = y ~ poly(x, 2)`. The result is shown in the left panel of Figure 7.12.

```
> gg <- ggplot(Donner, aes(age, as.numeric(survived=="yes"), color = sex)) +
+   theme_bw() + ylab("Survived") +
+   geom_point(position = position_jitter(height = 0.02, width = 0))
>
> gg + stat_smooth(method = "glm", family = binomial, formula = y ~ poly(x, 2),
+                    alpha = 0.2, size=2, aes(fill = sex))
>
> gg + stat_smooth(method = "loess", span=0.9, alpha = 0.2, size=2,
+                    aes(fill = sex)) + coord_cartesian(ylim=c(-.05,1.05))
```

This plot is quite surprising. It suggests quite different regimes relating to survival for men and women. Among men, survival probability decreases steadily with age, at least after age 20. For women, those in the age range 10–35 were very likely to have lived, while those over 40 were almost all predicted to perish.

Another simple technique is to fit a non-parametric loess smooth, as shown in the right panel of Figure 7.12.¹¹ The curve for females is similar to that of the quadratic fit in the left panel, but the curve for males suggests that survival also has a peak around the teenage years. One lesson to be drawn from these graphs is that a linear logistic regression, such as shown in Figure 7.12 may tell only part of the story, and, for a binary response it is not easy to discern whether the true relationship is linear. If it really is, all these graphs would look much more similar. As well, we usually obtain a more realistic smoothing of the data using full-model plots or effect plots.

The suggestions from these exploratory graphs can be used to define and test some models for survival in the Donner Party. The substantive questions of interest are:

¹¹A technical problem with the use of the loess smoother for binary data is that it can produce fitted values outside the [0–1] interval, as happens in the right panel of this figure. Kernel smoothers, such as the KernSmooth package avoid this problem, but are not available through ggplot2.

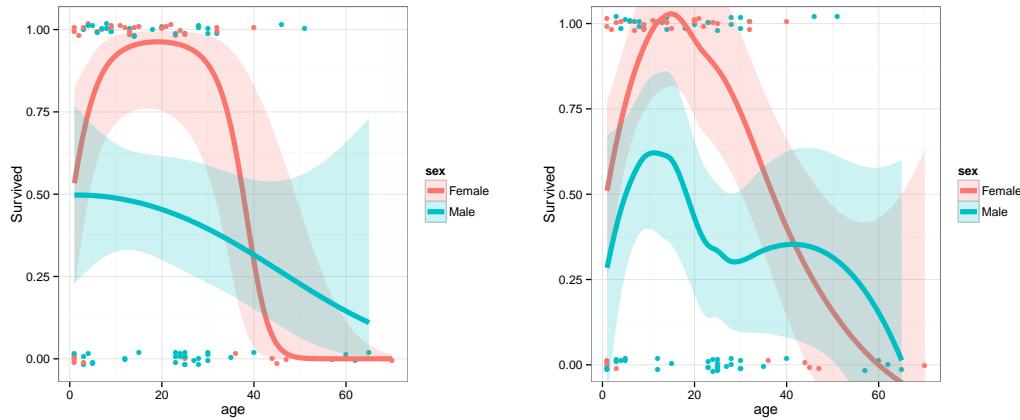


Figure 7.12: Conditionals plot of the Donner data, showing the relationship of survival to age and sex. Left: The smoothed curves and confidence bands show the result of fitting separate quadratic logistic regressions on age for males and females. Right: Separate loess smooths are fit to the data for males and females
fig:donner-conds

- Is relationship the same for men and women? This is, is it necessary to allow for an interaction of age with sex, or separate fitted curves for men and women?
- Is the relationship between survival and age well-represented in a linear logistic regression model?

The first question is the easiest to deal with: we can simply fit a model allowing an interaction of age (or some function of age) and sex,

```
survived ~ age * sex
survived ~ f(age) * sex
```

and compare the goodness of fit with the analogous additive, main-effects models.

From a modeling perspective, there is a wide variety of approaches for testing for non-linear relationships. We only scratch the surface here, and only for a single quantitative predictor, x , such as age in this example. One simple approach, illustrated in Figure 7.12 is to allow a quadratic (or higher-power, e.g., cubic) function to describe the relationship between the log odds and x ,

$$\begin{aligned}\text{logit}(\pi_i) &= \alpha + \beta_1 x_i + \beta_2 x_i^2 \\ \text{logit}(\pi_i) &= \alpha + \beta_1 x_i + \beta_2 x_i^2 + \beta_3 x_i^3 \\ &\dots\end{aligned}$$

In R, these model terms can be fit using `poly(x, 2)`, `poly(x, 3)` ..., which generate orthogonal polynomials for the powers of x . A simple way to test for non-linearity is a likelihood ratio test comparing the more complex model to the linear one. This method is often sufficient for a hypothesis test, and, if the relationship truly is linear, the fitted logits and probabilities will not differ greatly from what they would be under a linear model. A difficulty with this approach is that polynomial models are often unrealistic, particularly for data that approach an asymptote.

Another simple approach is to use a **regression spline**, that fits the relationship with x in terms of a set of piecewise polynomials, usually cubic, joined at a collection of points, called *knots* so that the overall fitted relationship is smooth and continuous. See Fox (2008, §17.2) for a cogent, brief description of these methods.

One particularly convenient method is a **natural spline**, implemented in the **splines** package in the `ns()` function. This method constrains the fitted cubic spline to be linear at lower and upper limits of x , and, for k knots, fits $df = k + 1$ parameters not counting the intercept. The k knots can be conveniently chosen as k cutpoints in the percentiles of the distribution of x . For example, with $k = 1$, the knot would be placed at the median, or 50th percentile; with $k = 3$, the knots would be placed at the quartiles of the distribution of x ; $k = 0$ corresponds to no knots, i.e., a simple linear regression.

In the `ns()` function, you can specify the locations of knots or the number of knots with the `knots` argument, but it is conceptually simpler to specify the number of degrees of freedom used in the spline fit. Thus, `ns(x, 2)` and `poly(x, 2)` both specify a term in x of the same complexity, the former a natural spline with $k = 1$ knot and the latter a quadratic function in x .

We illustrate these ideas in the remainder of this example, fitting a 2×2 collection of models to the `Donner` data corresponding to: (a) whether or not age and sex effects are additive; (b) whether the effect is linear on the logit scale or non-linear (quadratic, here). A brief summary of each model is given using the `Anova()` in the `car` package, providing Type II tests of each effect. As usual, `summary()` would give more detailed output, including tests for individual coefficients. First, we fit the linear models, without and with an interaction term:

```
> donner.mod1 <- glm(survived ~ age + sex,
+                      data=Donner, family=binomial)
> Anova(donner.mod1)

Analysis of Deviance Table (Type II tests)

Response: survived
          LR Chisq Df Pr(>Chisq)
age        5.52   1    0.0188 *
sex        6.73   1    0.0095 **
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

> donner.mod2 <- glm(survived ~ age * sex,
+                      data=Donner, family=binomial)
> Anova(donner.mod2)

Analysis of Deviance Table (Type II tests)

Response: survived
          LR Chisq Df Pr(>Chisq)
age        5.52   1    0.0188 *
sex        6.73   1    0.0095 **
age:sex    0.40   1    0.5269
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The main effects of `age` and `sex` are both significant here, but the interaction term, `age:sex` is not in model `donner.mod2`. Note that the terms tested by `Anova()` in `donner.mod1` are a redundant subset of those in `donner.mod2`.

Next, we fit non-linear models, representing the linear and non-linear trends in `age` by `poly(age, 2)`.¹² The `Anova()` results for terms in both models are contained in the output from `Anova(donner.mod4)`.

```
> donner.mod3 <- glm(survived ~ poly(age, 2) + sex,
+                      data=Donner, family=binomial)
> donner.mod4 <- glm(survived ~ poly(age, 2) * sex,
+                      data=Donner, family=binomial)
> Anova(donner.mod4)
```

¹²Alternatively, we could use the term `ns(age, 2)` or higher-degree polynomials or natural splines with more knots, but we don't do this here.

```
Analysis of Deviance Table (Type II tests)

Response: survived
          LR Chisq Df Pr(>Chisq)
poly(age, 2)      9.91  2     0.0070 **
sex                 8.09  1     0.0044 **
poly(age, 2):sex   8.93  2     0.0115 *
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Now, in model `donner.mod4`, the interaction term `poly(age, 2):sex` is significant, indicating that the fitted quadratics for males and females differ in “shape,” meaning either their linear (slope) or quadratic (curvature) components.

These four models address the questions posed earlier. A compact summary of these models, giving the likelihood ratio tests of goodness of fit, together with AIC and BIC statistics are shown below, using the `LRstats()` method in `vcdExtra` for a list of "glm" models. **TODO: LRstats gives the wrong pvalues**

```
> library(vcdExtra)
> LRstats(donner.mod1, donner.mod2, donner.mod3, donner.mod4)

Likelihood summary table:
      AIC BIC LR Chisq Df Pr(>Chisq)
donner.mod1 117 125    111.1 87     0.042 *
donner.mod2 119 129    110.7 86     0.038 *
donner.mod3 115 125    106.7 86     0.064 .
donner.mod4 110 125     97.8 84     0.144
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

By AIC and BIC, `donner.mod4` is best, and it is also the only model with a non-significant LR χ^2 (residual deviance). Because these models comprise a 2×2 set of hypotheses, it is easier to compare models by extracting the LR statistics and arranging these in a table, together with the their row and column differences. The entries in the table below are calculated as follows.

```
> mods <- list(donner.mod1, donner.mod2, donner.mod3, donner.mod4)
> LR <- sapply(mods, function(x) x$deviance)
> LR <- matrix(LR, 2, 2)
> rownames(LR) <- c("additive", "non-add")
> colnames(LR) <- c("linear", "non-lin")
> LR <- cbind(LR, diff= LR[,1]-LR[,2])
> LR <- rbind(LR, diff= c(LR[1,1:2]-LR[2,1:2], NA))
```

	linear	non-linear	$\Delta\chi^2$	p-value
additive	111.128	106.731	4.396	0.036
non-additive	110.727	97.799	12.928	0.000
$\Delta\chi^2$	0.400	8.932		
p-value	0.527	0.003		

Thus, the answer to our questions seems to be that: (a) there is evidence that the relationship of survival to age differs for men and women in the Donner Party; (b) these relationships are not well-described by a linear logistic regression.

For simplicity, we used a quadratic effect, `poly(age, 2)`, to test for non-linearity here. An alternative test of the same complexity could use a regression spline, `ns(age, 2)`, also with 2 degrees of freedom for the main effect and interaction, or allow more knots. To illustrate, we fit two natural spline modes models with 2 and 4 df, and compare these with the quadratic model (`donner.mod4`), all of which include the interaction of age and sex.

```

> library(splines)
> donner.mod5 <- glm(survived ~ ns(age, 2) * sex, data=Donner,
+                      family=binomial)
> Anova(donner.mod5)

Analysis of Deviance Table (Type II tests)

Response: survived
          LR Chisq Df Pr(>Chisq)
ns(age, 2)      9.28   2     0.0097 **
sex            7.98   1     0.0047 **
ns(age, 2):sex  8.71   2     0.0129 *
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

> donner.mod6 <- glm(survived ~ ns(age, 4) * sex, data=Donner,
+                      family=binomial)
> Anova(donner.mod6)

Analysis of Deviance Table (Type II tests)

Response: survived
          LR Chisq Df Pr(>Chisq)
ns(age, 4)      22.05   4     0.0002 ***
sex            10.49   1     0.0012 **
ns(age, 4):sex  8.54   4     0.0737 .
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

> LRstats(donner.mod4, donner.mod5, donner.mod6)

Likelihood summary table:
      AIC BIC LR Chisq Df Pr(>Chisq)
donner.mod4 110 125    97.8 84     0.14
donner.mod5 111 126    98.7 84     0.13
donner.mod6 106 131    86.1 80     0.30

```

With four more parameters, `donner.mod6` fits better and has a smaller AIC.

We conclude this example with an effect plot for the spline model `donner.mod6` shown in Figure ???. The complexity of the fitted relationships for men and women is intermediate between the two conditional plots shown in Figure 7.12. (However, note that the fitted effects are plotted on the logit scale in Figure ?? and labeled with the corresponding probabilities, whereas the conditional plots are plotted directly on the probability scale.)

```

> library(effects)

Error: package 'effects' was built before R 3.0.0: please re-install it

> donner.eff6 <- allEffects(donner.mod6, xlevels=list(age=seq(0,50,5)))

Error in eval(expr, envir, enclos): could not find function "allEffects"

> plot(donner.eff6, ticks=list(at=c(0.001, 0.01, 0.05, 0.1, 0.25,
+                                     0.5, 0.75, 0.9, 0.95, 0.99, 0.999)))
Error in plot(donner.eff6, ticks = list(at = c(0.001, 0.01, 0.05, 0.1, : object
'donner.eff6' not found

```

This plot confirms that for women in the Donner Party, survival was greatest for those aged 10-30. Survival among men was overall much less and there is a hint of greater survival for men aged 10-15.

Of course, this statistical analysis does not provide explanations for these effects, and it ignores the personal details of the Donner Party members and the individual causes and circumstances of death, which are generally well-documented in the historical record (Johnson, 1996). See <http://user.xmission.com/~octa/DonnerParty/> for a comprehensive collection of historical sources.

Grayson (1990) attributes the greater survival of women of intermediate age to demographic arguments that women are overall better able to withstand conditions of famine and extreme cold, and high age-specific mortality rates among the youngest and oldest members of human societies. He also concludes (without much analysis) that members with larger social and kinship networks would be more likely to survive. △

{ex:arrests}

EXAMPLE 7.10: Racial profiling: Arrests for marijuana possession

In the summer of 2002, the *Toronto Star* newspaper launched an investigation on the topic of possible racial profiling by the Toronto police service. Through freedom of information requests, they obtained a data base of over 600,000 arrest records on all potential charges in the period from 1996–2002, the largest data bases on crime arrests and disposition ever assembled in Canada. An initial presentation of this study was given in Example 1.4.

In order to examine the issue of racial profiling (different treatment as a function of race) they excluded all charges such as assault, robbery, speeding and driving under the influence, where the police have no discretion regarding the laying of a charge. They focused instead on a subset of arrests, where the police had various options.

Among these, for people arrested for a single charge of simple possession of a small amount of marijuana, police have the option of releasing the arrestee, with a summons (“Form 9”) to appear in court (similar to a parking ticket), or else the person could be given harsher treatment—brought to a police station or held in jail for a bail hearing (“Show cause”). The main question for the *Toronto Star* was whether the subject’s skin color had any influence on the likelihood that the person would be released with a summons.¹³

Their results, published in a week-long series of articles in December 2002, concluded that there was strong evidence that black and white subjects were treated differently. For example, the analysis showed that blacks were 1.5 times more likely than whites to be given harsher treatment than release with a summons; if the subject was taken to the police station, a black was 1.6 times more likely to be held in jail for a bail hearing. An important part of the analysis and the public debate that ensued was to show that other variables that might account for these differences had been controlled or adjusted for.¹⁴

The data set *Arrests* in the *effects* package gives a simplified version of the *Star* database, containing records for 5226 cases of arrest on the charge of simple possession of marijuana analyzed by the newspaper. The response variable here is *released* (Yes/No) and the main predictor of interest is skin color of the person arrested, *colour* (Black/White).¹⁵ A random subset of the data set is shown below.

```
> library(effects)
Error: package 'effects' was built before R 3.0.0: please re-install it
```

¹³Another discretionary charge they investigated was police stops for non-moving violations under the Ontario *Highway Traffic Act*, such as being pulled over for a faulty muffler or having an expired license plate renewal sticker. A disproportionate rate of charges against blacks is sometimes referred to as “driving while black” (DWB). This investigation found that the number of blacks so charged, but particularly young black males, far out-weighed their representation in the population.

¹⁴The Toronto Police Service launched a class-action libel law suit against the *Toronto Star* and the first author of this book, who served as their statistical consultant, claiming damages of \$5,000 for every serving police officer in the city, a total of over 20 million dollars. The suit was thrown out of court, and the Toronto police took efforts to enhance training programs to combat the perception of racial profiling.

¹⁵The original data set also contained the categories Brown and Other, but these appeared with small frequencies.

```
> data("Arrests", package="effects")
> Arrests[sample(nrow(Arrests), 6),]

  released colour year age sex employed citizen checks
3768     Yes  Black 2000  23 Male      No     Yes     4
4576     Yes  Black 2001  17 Male     Yes     Yes     0
3976      No  White 2002  20 Male      No     Yes     3
4629     Yes  White 2000  18 Male     Yes     Yes     1
2384      No  Black 2000  19 Male     Yes     Yes     3
869      Yes  White 2001  15 Male     Yes     Yes     1
```

Other available predictors, to be used as control variables included the year of the arrest, age and sex of the person, and binary indicators of whether the person was employed and a citizen of Canada. In addition, when someone is stopped by police, his/her name is checked in six police data bases that record previous arrests, convictions, whether on parole, etc. The variable `checks` records the number, 0–6, in which the person's name appeared.

A variety of logistic models were fit to these data including all possible main effects and some two-way interactions. To allow for possible non-linear effects of `year`, this variable was treated as a factor rather than as a (linear) numeric variable, but the effects of `age` and `checks` were reasonably linear on the logit scale. A reasonable model included the interactions of `colour` with both `year` and `age`, as fit below:

```
> Arrests$year <- as.factor(Arrests$year)
> arrests.mod <- glm(released ~ employed + citizen + checks
+                      + colour*year + colour*age,
+                      family=binomial, data=Arrests)
```

For such models, significance tests for the model terms are best carried out using the `Anova()` function in the `car` package that uses Type II tests ...

```
> library(car)
> Anova(arrests.mod)

Analysis of Deviance Table (Type II tests)

Response: released
          LR Chisq Df Pr(>Chisq)
employed    72.7   1    < 2e-16 ***
citizen     25.8   1    3.8e-07 ***
checks      205.2  1    < 2e-16 ***
colour       19.6   1    9.7e-06 ***
year         6.1    5    0.29785
age          0.5    1    0.49827
colour:year  21.7   5    0.00059 ***
colour:age   13.9   1    0.00019 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The difficulty in interpreting these results from tables of coefficients can be seen in the output below:

```
> coeftest(arrests.mod)

z test of coefficients:
                                         Estimate Std. Error z value Pr(>|z|)
(Intercept)                  0.34443   0.31007   1.11  0.26665
employedYes                 0.73506   0.08477   8.67 < 2e-16 ***
citizenYes                  0.58598   0.11377   5.15  2.6e-07 ***
checks                     -0.36664   0.02603  -14.08 < 2e-16 ***
```

```

colourWhite      1.21252   0.34978   3.47  0.00053 ***
year1998       -0.43118   0.26036  -1.66  0.09770 .
year1999       -0.09443   0.26154  -0.36  0.71805
year2000       -0.01090   0.25921  -0.04  0.96647
year2001        0.24306   0.26302   0.92  0.35541
year2002        0.21295   0.35328   0.60  0.54664
age            0.02873   0.00862   3.33  0.00086 ***
colourWhite:year1998  0.65196   0.31349   2.08  0.03756 *
colourWhite:year1999  0.15595   0.30704   0.51  0.61152
colourWhite:year2000  0.29575   0.30620   0.97  0.33411
colourWhite:year2001 -0.38054   0.30405  -1.25  0.21073
colourWhite:year2002 -0.61732   0.41926  -1.47  0.14091
colourWhite:age      -0.03737   0.01020  -3.66  0.00025 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

By direct calculation (e.g., using `exp(coef(arrests.mod))`) you can find that the odds of a quick release was $\exp(0.735) = 2.08$ times greater for someone employed, $\exp(0.586) = 1.80$ times more likely for a Canadian citizen and $\exp(1.21) = 3.36$ times more likely for a white than a black person. It is much more difficult to interpret the interaction terms.

The primary question for the newspaper concerned the overall difference between the the treatment of blacks and whites– the main effect of `colour`. We plot this as shown below, giving the plot shown in Figure ???. This supports the claim by the *Star* because the 95% confidence limits for blacks and whites do not overlap, and all other relevant predictors that could account for this effect have been controlled or adjusted for.

```

> plot(Effect("colour", arrests.mod),
+       lwd=3, ci.style="bands", main="",
+       xlab = list("Skin color of arrestee", cex=1.25),
+       ylab = list("Probability(released)", cex=1.25)
+     )

Error in plot(Effect("colour", arrests.mod), lwd = 3, ci.style = "bands", :
could not find function "Effect"

```

Of course, one should be very wary of interpreting main effects when there are important interactions, and the story turned out to be far more nuanced than was reported in the newspaper. In particular, the interactions of color with with age and year provided a more complete account. Effect plots for these interactions are shown in Figure ???.

```

> # colour x age interaction
> plot(Effect(c("colour", "age"), arrests.mod),
+       lwd=3, multiline=TRUE,
+       xlab=list("Age", cex=1.25),
+       ylab=list("Probability(released)", cex=1.25),
+       key.args=list(x=.05, y=.99, cex=1.2)
+     )

Error in plot(Effect(c("colour", "age"), arrests.mod), lwd = 3, multiline =
TRUE, :  could not find function "Effect"

> # colour x year interaction
> plot(Effect(c("colour", "year"), arrests.mod),
+       lwd=3, multiline=TRUE,
+       xlab=list("Year", cex=1.25),
+       ylab=list("Probability(released)", cex=1.25),
+       key.args=list(x=.7, y=.99, cex=1.2)
+     )

Error in plot(Effect(c("colour", "year"), arrests.mod), lwd = 3, multiline
= TRUE, :  could not find function "Effect"

```

From the left panel in Figure ??, it is immediately apparent that the effect of age was in opposite directions for blacks and whites: Young blacks were indeed treated more severely than young whites; however for older people, blacks were treated less harshly than whites, controlling for all other predictors.

The right panel of Figure ?? shows the changes over time in the treatment of blacks and whites. It can be seen that up to the year 2000 there was strong evidence for differential treatment on these charges, again controlling for other predictors. There was also evidence to support the claim by the police that in the year 2001 they began training of officers to reduce racial effects in treatment.

Finally, the `effects` package provides a convenience function, `allEffects()`, that calculates the effects for all high-order terms in a given model. The `plot()` method for the "efflist" object can be used to plot individual terms selectively from a graphic menu, or plot all terms together in one comprehensive display using `ask=FALSE`.

```
> arrests.effects <- allEffects(arrests.mod,
+                               xlevels=list(age=seq(15, 45, 5)))
Error in eval(expr, envir, enclos): could not find function "allEffects"
> plot(arrests.effects,
+       ylab="Probability(released)", ci.style="bands", ask=FALSE)
Error in plot(arrests.effects, ylab = "Probability(released)", ci.style = "bands",
: object 'arrests.effects' not found
```

The result, shown in Figure ?? is a relatively compact and understandable summary of the `arrests.mod` model: (a) people were more likely to be released if they were employed and citizens. (b) each additional police check decreased the likelihood of release with a summons. (c) the effect of skin color varied with age and year of arrest, in ways that tell a far more nuanced story than reported in the newspaper.

Finally, another feature of this plot bears mention: by default, the scales for each effect plot are determined separately for each effect, to maximize use of the plot region. However, you have to read the Y scale values to judge the relative sizes of these effects. An alternative plot, using the *same* scale in each subplot¹⁶ would show the relative sizes of these effects.

△

7.4.1 More complex models: Model selection and visualization

Models with more predictors or more complex terms (interactions, non-linear terms) present additional challenges for model fitting, summarization, and visualization and interpretation. A very complicated model, with many terms and interactions may fit the data at hand quite well. However, because goodness-of-fit is optimized in the sample, terms that appear significant are less likely to be important in a future sample, and we need to worry about inflation of Type I error rates that accompany multiple significance tests. As well, it becomes increasingly difficult to visualize and understand a fitted model as the model becomes increasingly complex.

On the other hand, a very simple model may omit important predictors, interactions, or non-linear relationships with the response and give an illusion of a comfortable interpretation.

TODO: Complete this brief introduction to model selection and define AIC/BIC,

{sec:complex}

{ex:icul}

EXAMPLE 7.11: Death in the ICU

In this example we examine briefly some aspects of logistic regression related to model selection and graphical display with a large collection of potential predictors, including both quantitative and

¹⁶With the `effects` package, you can set the `ylim` argument to equate the vertical range for all plots, but this should be done on the logit scale. For this plot, `ylim = plogis(c(0.5, 1))` would work.

discrete variables. We use data from a classic study by Lemeshow *et al.* (1988) of patients admitted to an intensive care unit at Baystate Medical Center in Springfield, Massachusetts. The major goal of this study was to develop a model to predict the probability of survival (until hospital discharge) of these patients and to study the risk factors associated with ICU mortality. The data, contained in the data set *ICU* in *vcdExtra*, gives the results for a sample of 200 patients that was presented in Hosmer *et al.* (2013) (and earlier editions).

The *ICU* contains 22 variables of which the first, *died* is a factor. Among the predictors, two variables (*race*, *coma*) were represented initially as 3-level factors, but then recoded to binary variables (*white*, *uncons*).

```
> data("ICU", package="vcdeExtra")
> names(ICU)

[1] "died"      "age"       "sex"        "race"       "service"
[6] "cancer"     "renal"      "infect"     "cpr"        "systolic"
[11] "hrrate"    "previcu"   "admit"     "fracture"   "po2"
[16] "ph"         "pco"       "bic"        "creatin"   "coma"
[21] "white"      "uncons"    # remove redundant race, coma
```

Removing the 3-level versions leaves 19 predictors, of which three (age, heart rate, systolic blood pressure) are quantitative and the remainder are either binary (*service*, *cancer*) or had previously been dichotomized ($\text{ph} < 7.25$).

As an initial step, and a basis for comparison, we fit the full model containing all 19 predictors.

```
> icu.full <- glm(died ~ ., data=ICU, family=binomial)
> summary(icu.full)

Call:
glm(formula = died ~ ., family = binomial, data = ICU)

Deviance Residuals:
    Min      1Q  Median      3Q      Max 
-1.8040 -0.5606 -0.2044 -0.0863  2.9773 

Coefficients:
            Estimate Std. Error z value Pr(>|z|)    
(Intercept) -6.72670  2.38551 -2.82   0.0048 **  
age          0.05639  0.01862  3.03   0.0025 **  
sexMale      0.63973  0.53139  1.20   0.2286    
serviceSurgical -0.67352  0.60190 -1.12   0.2631    
cancerYes    3.10705  1.04585  2.97   0.0030 **  
renalYes     -0.03571  0.80165 -0.04   0.9645    
infectYes    -0.20493  0.55319 -0.37   0.7110    
cprYes        1.05348  1.00661  1.05   0.2953    
systolic      -0.01547  0.00850 -1.82   0.0686 .    
hrrate        -0.00277  0.00961 -0.29   0.7732    
previcuYes   1.13194  0.67145  1.69   0.0918 .    
admitEmergency 3.07958  1.08158  2.85   0.0044 **  
fractureYes   1.41140  1.02971  1.37   0.1705    
po2<=60       0.07382  0.85704  0.09   0.9314    
ph<7.25       2.35408  1.20880  1.95   0.0515 .    
pco>45       -3.01844  1.25345 -2.41   0.0160 *    
bic<18        -0.70928  0.90978 -0.78   0.4356    
creatin>2    0.29514  1.11693  0.26   0.7916    
whiteNon-white 0.56573  0.92683  0.61   0.5416    
unconsYes     5.23229  1.22630  4.27   2e-05 *** 
---

```

```
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
(Dispersion parameter for binomial family taken to be 1)

Null deviance: 200.16 on 199 degrees of freedom
Residual deviance: 120.78 on 180 degrees of freedom
AIC: 160.8

Number of Fisher Scoring iterations: 6
```

You can see that a few predictors are individually significant, but many are not.

However, it is useful to carry out a simultaneous global test of $H_0 : \beta = 0$ that *all* regression coefficients are zero. If this test is not significant, it makes little sense to use selection methods to choose individually significant predictors. For convenience, we define a simple function, `LRtest()`, to calculate the likelihood ratio test from the model components.

```
> LRtest <- function(model)
+   c(LRchisq=(model$null.deviance - model$deviance),
+     df=(model$df.null - model$df.residual))
>
> (LR <- LRtest(icu.full))

LRchisq      df
79.383 19.000

> (pvalue=1-pchisq(LR[1],LR[2]))

LRchisq
2.3754e-09
```

At this point, it is tempting to examine the output from `summary(icu.full)` shown above and eliminate those predictors which fail significance at some specified level such as the conventional $\alpha = 0.05$. This is generally a bad idea for many reasons.¹⁷

A marginally better approach is to remove non-significant variables whose coefficients have signs that don't make sense from the substance of the problem. For example, in the full model, both `renal` (history of chronic renal failure) and `infect` (infection probable at ICU admission) have negative signs, meaning that their presence *decreases* the odds of death. We remove those variables using `update()`; as expected they make little difference.

```
> icu.full1 <- update(icu.full, . ~ . - renal - fracture)
> anova(icu.full1, icu.full, test="Chisq")

Analysis of Deviance Table

Model 1: died ~ age + sex + service + cancer + infect + cpr + systolic +
          hrrate + previcu + admit + po2 + ph + pco + bic + creatin +
          white + uncons
Model 2: died ~ age + sex + service + cancer + renal + infect + cpr +
          systolic + hrrate + previcu + admit + fracture + po2 + ph +
          pco + bic + creatin + white + uncons
Resid. Df Resid. Dev Df Deviance Pr(>Chi)
1       182      122
2       180      121  2      1.7      0.43
```

Before proceeding to consider model selection, it is useful to get a better visual overview of the current model than is available from a table of coefficients and significance tests. Some very useful `print()`, `summary()` and `plot()` methods are available in the `rsm` package. Unfortunately,

¹⁷It ignores the facts of (a) an arbitrary cutoff value for significance, (b) the strong likelihood that chance features of the data or outliers influence the result, (c) problems of collinearity, etc. See Harrell (2001, §4.3) for a useful discussion of these issues.

these require that the logistic model is fitted with `lrm()` in that package rather than with `glm()`. We pause here to refit the same model as `icu.full1` in order to show a plot of odds ratios for the terms in this model.

```
> library(rms)

Error in library(rms): there is no package called 'rms'

> dd <- datadist(ICU[, -1])

Error in eval(expr, envir, enclos): could not find function "datadist"

> options(datadist="dd")
> icu.lrml <- lrm(died ~ ., data=ICU)

Error in eval(expr, envir, enclos): could not find function "lrm"

> icu.lrml <- update(icu.lrml, . ~ . - renal - fracture)

Error in update(icu.lrml, . ~ . - renal - fracture): object 'icu.lrml' not
found
```

The `summary()` method for "rms" objects produces a much more detailed descriptive summary of a fitted model, and the `plot()` method for that summary object gives a sensible plot of the odds ratios for the model terms together with confidence intervals, at levels (0.9, 0.95, 0.99) by default. The following lines produce Figure 7.13.

```
> sum.lrml <- summary(icu.lrml)
> plot(sum.lrml, log=TRUE, main="Odds ratio for 'died'", cex=1.25,
+       col = rgb(0.1, 0.1, 0.8, alpha = c(0.3, 0.5, 0.8)))
```

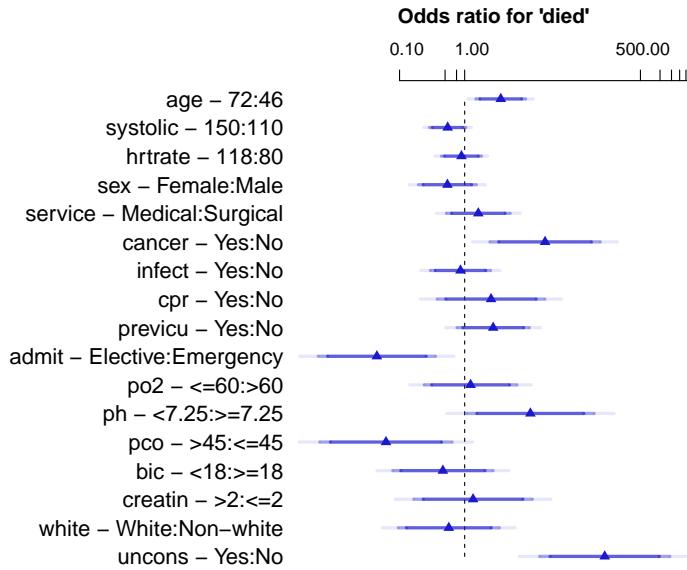


Figure 7.13: Odds ratios for the terms in the model for the ICU data. Each line shows the odds ratio for a term, together with lines for 90, 95 and 99% confidence intervals in progressively darker shades.

In this plot, continuous variables are shown at the top, followed by the discrete predictors. In

each line, the range or levels of the predictors are given in the form $a : b$, such that the value a corresponds to the numerator of the odds ratio plotted. Confidence intervals that don't overlap the vertical line for odds ratio = 1 are significant, but this graph shows those at several confidence levels, allowing you to decide what is "significant" visually. As well, the widths of those intervals convey the precision of these estimates.

Among several stepwise selection methods in R for "glm" models, `stepAIC()` in the MASS package implements a reasonable collection of methods for forward, backward and stepwise selection using penalized AIC-like criteria that balance goodness of fit against parsimony. The method takes an argument, `scope`, which is a list of two model formulae; `upper` defines the largest (most complex) model to consider and `lower` defines the smallest (simplest) model, e.g., `lower = ~ 1` is the intercept-only model.

By default, the function produces verbose printed output showing the details of each step, but we suppress that here to save space. It returns the final model as its result, along with an `anova` component that summarises the deviance and AIC from each step.

```
> library(MASS)
> icu.step1 <- stepAIC(icu.full1, trace = FALSE)
> icu.step1$anova

Stepwise Model Path
Analysis of Deviance Table

Initial Model:
died ~ age + sex + service + cancer + infect + cpr + systolic +
     hrrate + previcu + admit + po2 + ph + pco + bic + creatin +
     white + uncons

Final Model:
died ~ age + cancer + systolic + admit + ph + pco + uncons

      Step Df Deviance Resid. Df Resid. Dev      AIC
1              - po2   1  0.062446    182  122.48 158.48
2 - creatin   1  0.059080    183  122.54 156.54
3 - hrrate   1  0.072371    184  122.60 154.60
4 - infect   1  0.122772    185  122.67 152.67
5 - white    1  0.334999    186  122.79 150.79
6 - service   1  0.671313    187  123.13 149.13
7 - bic      1  0.377521    188  123.80 147.80
8 - cpr      1  1.148260    189  124.18 146.18
9 - sex      1  1.543523    190  125.33 145.33
10 - previcu  1  1.569976   191  126.87 144.87
11 -          1               192  128.44 144.44
```

Alternatively, we can use the BIC criterion, by specifying `k=log(n)`, which generally will select a smaller model when the sample size is reasonably large.

```
> icu.step2 <- stepAIC(icu.full, trace = FALSE, k=log(200))
> icu.step2$anova

Stepwise Model Path
Analysis of Deviance Table

Initial Model:
died ~ age + sex + service + cancer + renal + infect + cpr +
     systolic + hrrate + previcu + admit + fracture + po2 + ph +
     pco + bic + creatin + white + uncons

Final Model:
died ~ age + cancer + admit + uncons
```

	Step	Df	Deviance	Resid.	Df	Resid.	Dev	AIC
1				180		120.78	226.74	
2	- renal	1	0.0019881	181		120.78	221.45	
3	- po2	1	0.0067968	182		120.79	216.16	
4	- creatin	1	0.0621463	183		120.85	210.92	
5	- hrrate	1	0.0658870	184		120.92	205.69	
6	- infect	1	0.2033221	185		121.12	200.59	
7	- white	1	0.3673180	186		121.49	195.66	
8	- bic	1	0.6002993	187		122.09	190.96	
9	- service	1	0.7676303	188		122.85	186.43	
10	- fracture	1	1.3245086	189		124.18	182.46	
11	- cpr	1	1.1482598	190		125.33	178.31	
12	- sex	1	1.5435228	191		126.87	174.55	
13	- previcu	1	1.5699762	192		128.44	170.83	
14	- ph	1	4.4412370	193		132.88	169.97	
15	- pco	1	2.7302934	194		135.61	167.40	
16	- systolic	1	3.5231028	195		139.13	165.63	

This model differs from model `icu.step1` selected using AIC in the last three steps, that also removed `ph`, `pco` and `systolic`.

```
> coefest(icu.step2)

z test of coefficients:

Estimate Std. Error z value Pr(>|z|)
(Intercept) -6.8698    1.3188   -5.21  1.9e-07 ***
age          0.0372    0.0128    2.91  0.00360 **
cancerYes    2.0971    0.8385    2.50  0.01238 *
admitEmergency 3.1022    0.9186    3.38  0.00073 ***
unconsYes    3.7055    0.8765    4.23  2.4e-05 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

These two models are nested, so we can compare them directly using a likelihood ratio test from `anova()`.

```
> anova(icu.step2, icu.step1, test="Chisq")

Analysis of Deviance Table

Model 1: died ~ age + cancer + admit + uncons
Model 2: died ~ age + cancer + systolic + admit + ph + pco + uncons
  Resid. Df Resid. Dev Df Deviance Pr(>Chi)
1        195     139
2        192     128  3      10.7    0.013 *
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The larger model is significantly better by this test, but the smaller model is simpler to interpret. We retain these both as “candidate models” to be explored further, but for ease in this example, we do so using the smaller model, `icu.step2`.

Another important step is to check for non-linearity of quantitative predictors such as `age` and interactions among the predictors. This is easy to do using `update()` and `anova()` as shown below. First, allow a non-linear term in `age`, and all two-way interactions of the binary predictors.

```
> icu.glm3 <- update(icu.step2, . ~ . -age + ns(age, 3) + (cancer+admit+uncons)^2)
> anova(icu.step2, icu.glm3, test="Chisq")

Analysis of Deviance Table

Model 1: died ~ age + cancer + admit + uncons
Model 2: died ~ cancer + admit + uncons + ns(age, 3) + cancer:admit +
          cancer:uncons + admit:uncons
Resid. Df Resid. Dev Df Deviance Pr(>Chi)
1       195      139
2       191      135   4     3.73    0.44
```

Next, we can check for interactions with age:

```
> icu.glm4 <- update(icu.step2, . ~ . + age*(cancer+admit+uncons))
> anova(icu.step2, icu.glm4, test="Chisq")

Analysis of Deviance Table

Model 1: died ~ age + cancer + admit + uncons
Model 2: died ~ age + cancer + admit + uncons + age:cancer + age:admit +
          age:uncons
Resid. Df Resid. Dev Df Deviance Pr(>Chi)
1       195      139
2       192      134   3     5.37    0.15
```

None of these additional terms have much effect.

So, we will tentatively adopt the simple main effects model, `icu.step2`, and consider how to visualize and interpret this result. One interesting display is a **nomogram** that shows how values on the various predictors translate into a predicted value of the log odds, and the relative strengths of their effects on this prediction. This kind of plot is shown in Figure 7.14, produced using `nomogram()` in the `rms` package as follows. It only works with models fit using `lrm()`, so we have to refit this model.

```
> icu.lrm2 <- lrm(died ~ age + cancer + admit + uncons, data=ICU)
> plot(nomogram(icu.lrm2), cex.var=1.2, lplabel="Log odds death")
```

In this nomogram, each predictor is scaled according to the size of its effect on a common scale of 0–100 “points.” A representative observation is shown by the marked points, corresponding to a person of age 60, without cancer, who was admitted to emergency and was unconscious at that time. Adding the points associated with each variable value gives the result shown on the scale of total points. For this observation, the result is $50 + 0 + 84 + 100 = 234$, for which the scale of log odds at the bottom gives a predicted logit of 2.2, or a predicted probability of death of $1/(1 + \exp(-2.2)) = 0.90$.

This leaves us with the problem of how to visualize the fitted model compactly and comprehensively. Full-model plots and effect plots, as we have used them, are somewhat unwieldy with four or more predictors if we want to view all effects simultaneously because it is more difficult to make comparisons across multiple panels (particularly if the vertical scales differ).

One way to reduce the visual complexity of such graphs is to combine some predictors that would otherwise be shown in separate panels into a recoding that can be shown as multiple curves for their combinations in fewer panels. In general, this can be done by combining some predictors interactively; for example with sex and education as factors, their combinations, M:Hi, M:Lo, etc. could be used to define a new variable, `group` used as the curves in one plot, rather than separate panels.

In this case, because age is continuous, it makes sense to plot fitted values against age. With `cancer`, `admit` and `uncons` as binary factors associated with risk of death, it is also sensible

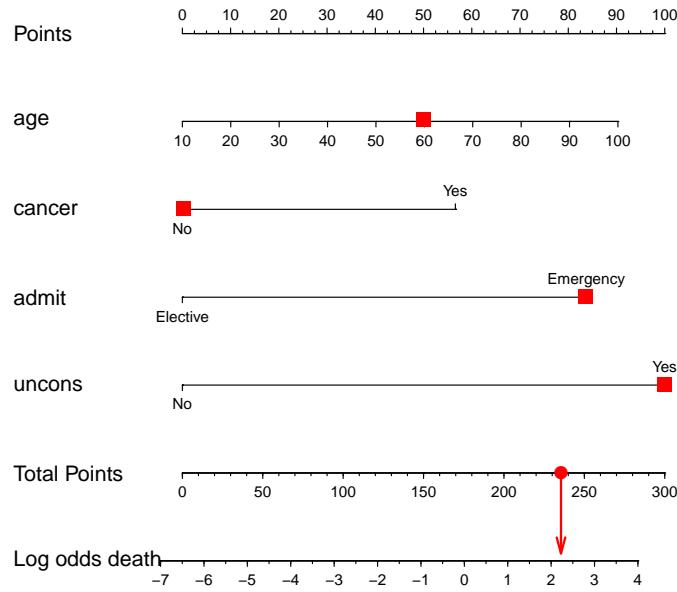


Figure 7.14: Nomogram for predicted values in the simple main effects model for the ICU data. Each predictor is scaled in relation to its effect on the outcome in terms of “points”, 0–100. Adding the points for a given case gives total points that have a direct translation to log odds. The marked points show the prediction for someone of age 60, admitted to the emergency ward and unconscious.

{fig:icu-nomogram}

to combine them all into a single variable, `risks`, indicating which one or more risk factors are present for each case. We first convert each variable to an abbreviation for the risk, if present, or "", and paste these together.

```
> # combine categorical risk factors to a single string
> risks <- ICU[, c("cancer", "admit", "uncons")]
> risks[1] <- ifelse(risks[1]=="Yes", "Cancer", "")
> risks[2] <- ifelse(risks[2]=="Emergency", "Emerg", "")
> risks[3] <- ifelse(risks[3]=="Yes", "Uncons", "")
> risks <- apply(risks, 1, paste, collapse="")
> risks[risks==""] <- "(none)"
```

	Cancer	CancerEmerg	Emerg	EmergUncons	(none)
Cancer	15	5	128	14	37
Uncons					1

The frequency counts of the risk combinations show that admission to the emergency ward alone was most frequent, and only one patient had unconsciousness as the only risk.

As done before, we can then get the fitted logit values for the chosen model, and combine these with the data and the `risks` variable.

```
> icu.glm2 <- glm(died ~ age + cancer + admit + uncons,
+                     data=ICU, family=binomial)
> icu.fit <- cbind(ICU, predict(icu.glm2, se=TRUE), risks)
```

In the plot step below, we use `geom_ribbon()` to plot a one standard error confidence band around the fitted logits. `color=risks` gives separate curves for each level of the `risks` factor.

```
> gg <- ggplot( icu.fit, aes(x=age, y=fit, color=risks)) +
+   geom_line(size = 1.2) + theme_bw() +
+   geom_ribbon(aes(ymin = fit - se.fit,
+                   ymax = fit + se.fit,
+                   fill = risks),
+               alpha = 0.2,
+               color = "transparent") +
+   theme_bw() +
+   labs(x = "Age", y = "Log odds (died)") +
+   geom_point(size=2)
```

By default, `ggplot()` uses a legend to display the labels for the curve variable, but the graph is more readable using `directlabels`, giving the plot shown in Figure 7.15.

```
> library(directlabels)
> direct.label(gg+xlim(10,100), last.points)
```

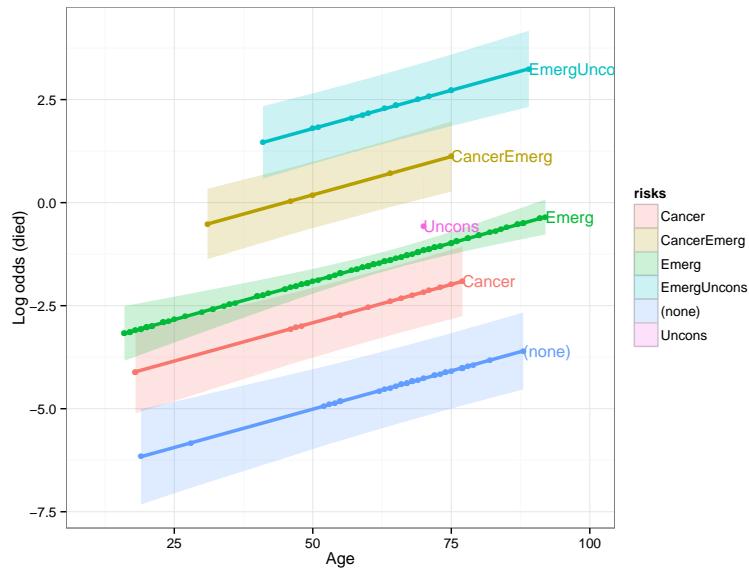


Figure 7.15: Fitted log odds of death in the ICU data. Each line shows the relationship with age, for patients having various combinations of risk factors.
Figure 7.15: Fitted log odds of death in the ICU data. Each line shows the relationship with age, for patients having various combinations of risk factors.

From this graph, it is apparent that the log odds of mortality increases with age in all cases. Relative to the line labeled `(none)`, mortality is higher when any of these risk factors are present, particularly when the patient is admitted to Emergency; it is highest when the patient is also unconscious at admission. The vertical gaps between lines that share a common risk (e.g., `Cancer`, `CancerEmerg`) indicate the additional increment from one more risk. Finally, the plotted points show the number and age distribution of these various combinations.

Before concluding that this model provides an adequate description of the data, we should examine whether any individual cases are unduly influencing the predicted results, and more importantly, the choice of variables in the model. We examine this question in Section 7.5 where we return to these data (Example 7.13).

```
Error in detach(package:rms): invalid 'name' argument
Error in detach(package:Hmisc): invalid 'name' argument
```



7.5 Influence and diagnostic plots

In ordinary least squares (OLS) regression, measures of *influence* (leverage, Cook's D, DFBETAs, etc.) and associated plots help you to determine whether individual cases (or cells in grouped data) have undue impact on the fitted regression model and the coefficients of individual predictors. Analogs of most of these measures have been suggested for logistic regression and generalized linear models. Pregibon (1981) provided the theoretical basis for these methods, exploiting the relationship between logistic models and weighted least squares. Some additional problems occur in practical applications to logistic regression because the response is discrete, and because the leave-one-out diagnostics are more difficult to compute, but the ideas are essentially the same.

{sec:logist-infl}

7.5.1 Residuals and leverage

As in ordinary least squares regression, the influence (actual impact) of an observation in logistic models depends multiplicatively on its residual (disagreement between y_i and \hat{y}_i) and its leverage (how unusual x_i is in the space of the explanatory variables). A conceptual formula is

$$\text{Influence} = \text{Leverage} \times \text{Residual}$$

This multiplicative definition implies that a case is influential to the extent that it is both poorly fit and has unusual values of the predictors.

7.5.1.1 Residuals

In logistic regression, the simple raw residual is just $e_i \equiv y_i - \hat{p}_i$, where $\hat{p}_i = 1/[1 + \exp(-\mathbf{x}_i^\top \mathbf{b})]$.

The Pearson and deviance residuals are more useful for identifying poorly fitted observations, and are components of overall goodness-of-fit statistics. The (raw) **Pearson residual** is defined as

$$r_i \equiv \frac{e_i}{\sqrt{\hat{p}_i(1 - \hat{p}_i)}} \quad (7.7)$$

and the Pearson chi-square is therefore $\chi^2 = \sum r_i^2$. The **deviance residual** is

$$g_i \equiv \pm -2[y_i \log \hat{p}_i + (1 - y_i) \log(1 - \hat{p}_i)]^{1/2} \quad (7.8)$$

where the sign of g_i is the same as that of e_i . Likewise, the sum of squares of the deviance residuals gives the overall deviance, $G^2 = -2 \log \mathcal{L}(\mathbf{b}) = \sum g_i^2$.

When y_i is a binomial count based on n_i trials (grouped data), the Pearson residuals Eqn. (7.7) then become

$$r_i \equiv \frac{y_i - n_i \hat{p}_i}{\sqrt{n_i \hat{p}_i(1 - \hat{p}_i)}}$$

with similar modifications made to Eqn. (7.8).

In R, `residuals()` is the generic function for obtaining (raw) residuals from a model fitted with `glm()` (or `lm()`). However **standardized residuals**, given by `rstandard()`, and **studentized residuals**, provided by `rstudent()` are often more useful because they rescale the residuals to have unit variance. They use, respectively, an overall estimate, $\hat{\sigma}^2$ of error variance, and the leave-one-out estimate, $\hat{\sigma}_{(-i)}^2$, omitting the i th observation; the studentized version is usually to be preferred in model diagnostics because it also accounts for the impact of the observation on residual variance.

7.5.1.2 Leverage

Leverage measures the *potential* impact of an individual case on the results, which is directly proportional to how far an individual case is from the centroid in the space of the predictors. Leverage is defined as the diagonal elements, h_{ii} , of the “Hat” matrix, \mathbf{H} ,

$$\mathbf{H} = \mathbf{X}^* (\mathbf{X}^{*\top} \mathbf{X}^*)^{-1} \mathbf{X}^{*\top}$$

where $\mathbf{X}^* = \mathbf{V}^{1/2} \mathbf{X}$, and $\mathbf{V} = \text{diag}[\hat{p}(1 - \hat{p})]$. As in OLS, leverage values are between 0 and 1, and a leverage value, $h_{ii} > \{2 \text{ or } 3\}k/n$ is considered “large”; here, $k = p + 1$ is the number of coefficients including the intercept and n is the number of cases. In OLS, however, the hat values depend only on the X s, whereas in logistic regression, they also depend on the dependent variable values and the fitted probabilities (through \mathbf{V}). As a result, an observation may be extremely unusual on the predictors, yet not have a large hat value, if the fitted probability is near 0 or 1. The function `hatvalues()` calculates these values for a fitted "glm" model object.

7.5.2 Influence diagnostics

Influence measures assess the effect that deleting an observation has on the regression parameters, fitted values, or the goodness-of-fit statistics. In OLS, these measures can be computed exactly from a single regression. In logistic regression, the exact effect of deletion requires refitting the model with each observation deleted in turn, a time-intensive computation. Consequently, Pregibon (1981) showed how analogous deletion diagnostics may be approximated by performing one additional step of the iterative procedure. Most modern implementations of these methods for generalized linear models follow Williams (1987).

The simplest measure of influence of observation i is the standardized change in the coefficient for each variable due to omitting that observation, termed **DFBETAs**. From the relation (Pregibon, 1981, p. 716)

$$\mathbf{b} - \mathbf{b}_{(-i)} = (\mathbf{X}^\top \mathbf{V} \mathbf{X})^{-1} \mathbf{x}_i (y_i - \hat{p}_i) / (1 - h_{ii}) ,$$

the estimated standardized change in the coefficient for variable j is

$$\text{DFBETA}_{ij} \equiv \frac{b_{(-i)j} - b_j}{\hat{\sigma}(b_j)} , \quad (7.9) \quad \{\text{eq:dfbeta}\}$$

where $\hat{\sigma}(b_j)$ is the estimated standard error of b_j . With k regressors, there are $k + 1$ sets of DFBETAs, which makes their examination burdensome. Graphical displays ease this burden, as do various summary measures considered below.

The most widely used summary of the overall influence of observation i on the estimated regression coefficients is **Cook's distance**, which measures the average squared distance between \mathbf{b} for all the data and $\mathbf{b}_{(-i)}$ estimated without observation i . It is defined as

$$C_i \equiv (\mathbf{b} - \mathbf{b}_{(-i)})^\top \mathbf{X}^\top \mathbf{V} \mathbf{X} (\mathbf{b} - \mathbf{b}_{(-i)}) / k \hat{\sigma}^2 .$$

However, Pregibon (1981) showed that C_i could be calculated simply as

$$C_i = \frac{r_i^2 h_{ii}}{k(1 - h_{ii})^2} , \quad (7.10) \quad \{\text{eq:cookd2}\}$$

where $r_i = y_i - \hat{p}_i / \sqrt{v_{ii}(1 - h_{ii})}$ is the i th standardized Pearson residual and v_{ii} is the i th diagonal element of \mathbf{V} . Rules of thumb for noticeably “large” values of Cook’s D are only rough indicators, and designed so that only “noteworthy” observations are nominated as unusually influential. One common cutoff for an observation to be treated as influential is $C_i > 1$. Others refer the values of C_i to a χ^2_k or $F_{k,n-k}$ distribution.

Another commonly used summary statistic of overall influence is the **DFFITS** statistic, a standardized measure of the difference between the predicted value \hat{y}_i using all the data and the predicted value $\hat{y}_{(-i)}$ calculated omitting the i th observation.

$$\text{DFFITS}_i = \frac{\hat{y}_i - \hat{y}_{(-i)}}{\hat{\sigma}_{(-i)} \sqrt{h_{ii}}} ,$$

where $\hat{\sigma}_{(-i)}$ is the estimated standard error with the i th observation deleted. For computation, DFFITS can be expressed in terms of the standardized Pearson residual and leverage as

$$\text{DFFITS}_i = r_i \sqrt{\frac{h_{ii}}{(1-h_{ii})} \frac{v_{ii}}{v_{(-ii)}}} . \quad (7.11)$$

From Eqn. (7.10) and Eqn. (7.11) it can be shown that Cook's distance is nearly the square of DFFITS divided by k ,

$$C_i = \frac{v_{(-ii)}^2}{v_{ii}^2} \frac{\text{DFFITS}_i^2}{k} . \quad (7.12)$$

Noteworthy values of DFFITS are often nominated by the rule-of-thumb $\text{DFFITS}_i > 2$ or $3\sqrt{k/n - k}$.

In R, these influence measures are calculated for a fitted "glm" model using `cooks.distance()` and `dffits()`. The `dfbeta()` function calculates and returns the matrix of all standardized changes in the model coefficients (Eqn. (7.9)) due to omitting each observation in turn.¹⁸ A convenience function, `influence.measures()` gives a tabular display showing the DFBETA_{ij} for each model variable, DFFITS, Cook's distances and the diagonal elements of the hat matrix. Cases which are influential with respect to any of these measures are marked with an asterisk.¹⁹

Beyond printed output of these numerical summaries, plots of these measures can shed light on potential problems due to influential or other noteworthy cases. By highlighting them, such plots provide the opportunity to determine if and how any of these affect your conclusions, or to take some corrective action.

A basic collection of diagnostic plots is provided by the `plot()` method for a "glm" model object. The `car` package contains a variety of other functions for model diagnostic plots. We illustrate some of these in the examples below.

EXAMPLE 7.12: Donner Party

This example re-visits the data on the Donner Party examined in Example 7.9. For illustrative purposes, we consider the influence measures and diagnostic plots for one specific model, the model `donner.mod3`, that included a quadratic effect of age and a main effect of sex, but no interaction.

The simplest overview of the adequacy of a fitted model is provided by the `plot()` method for a "glm" (or "lm") object. This function can produce up to six different plots that can be plotted individually or selected (using the argument `which`) and composed into a single overview figure using `par(mfrow=c(rows,cols))` as shown below.

It is useful to see the entire collection because, by default, only four are plotted (`which=c(1:3, 5)`) and this selection (sometimes called the **regression quartet** of diagnostic plots) is tuned more to classical linear models for quantitative data. Important feature of these plots are that (a) plot annotations are added to each showing trends or expected behaviour under the assumptions of a fitted model; (b) noteworthy observations are labeled individually.

¹⁸TODO: Not quite true: `dfbeta()` doesn't have a "glm" method. Either omit this or write a `dfbeta.glm()` method that gives the same results as `influence.measures()`.

¹⁹See `help(influence.measures)` for the description of all of these functions for residuals, leverage and influence diagnostics in generalized linear models.

```
> caption = list("(1) Residuals vs Fitted",
+                 "(2) Normal Q-Q",
+                 "(3) Scale-Location",
+                 "(4) Cook's distance",
+                 "(5) Residuals vs Leverage",
+                 expression("(6) Cook's dist vs Leverage   "
+                           * h[ii] / (1 - h[ii])))
> op <- par(mfrow=c(3,2), mar=c(4,4,2,1)+.1, cex.lab=1.2, cex=1)
> plot(donner.mod3, which=1:6, caption=caption)
> par(op)
```

The six plots, corresponding to the values of `which`, shown in Figure 7.16 for the `donner.mod3` model are:

1. a plot of residuals against fitted values. In a classical linear model, this plot should appear unstructured (random around the zero line), but for logistic regression there will always be two sequences of points, corresponding to the 0/1 observations.
2. a normal Q-Q plot of ordered residuals vs. the corresponding quantiles of the gaussian distribution. In a classical linear model, all points should follow the dotted reference line, but this will rarely hold for logistic regression models.
3. a Scale-Location plot of $\sqrt{|\text{residuals}|}$ against fitted values, with a loess smoothed curve showing the trend for variance of the residual to change with the predicted value. This is useful to detect non-constant residual variance in classical models, but in logistic regression, you will almost always see a U-shaped pattern corresponding to the fact that the variance around the fitted value is a function of $\sqrt{\hat{p}_i(1 - \hat{p}_i)}$.
4. an index plot of Cook's distances versus observation numbers, with the largest `id.n` values labeled.
5. a plot of residuals against leverages, showing contours of Cook's distances. Among all of these plots, this is probably the most useful for assessment of influence in both classical and generalized linear models. The function `influencePlot()` in `car` provides a better version of this plot, using the size of a bubble symbol to also show Cook's distance directly.
6. a plot of Cook's distances against leverage/(1-leverage). In this plot contours of standardized residuals that are equal in magnitude are lines through the origin, and labeled with their absolute values. Consequently, more influential observations appear toward the top.

In all these plots, three observations are labeled as noteworthy, by one criterion or another with a default number given by `id.n=3`. Plotting just the residual-leverage graph (`which=5`) with some additional annotations to show the conventional cutoff values gives Figure 7.17.

```
> op <- par(mar=c(5, 4, 4, 2)+.1)
> plot(donner.mod3, which=5, cex.id=1, cook.levels=c(0.25, 0.5), id.n=3)
> abline(h=c(-2, 2), col="gray")
> k <- length(coef(donner.mod3))
> n <- nrow(Donner)
> abline(v=c(2, 3)*k/n, col="gray")
> text(x=c(2, 3)*k/n, y=-2.3, c("2k/n", "3k/n"))
> par(op)
```

Details of all the diagnostic measures for a given model including the DFBETAs for individual coefficients can be obtained using `influence.measures`. This can be useful for custom plots not provided elsewhere (see Example 7.13).

```
> infl <- influence.measures(donner.mod3)
> names(infl)
[1] "infmat" "is.inf" "call"
```

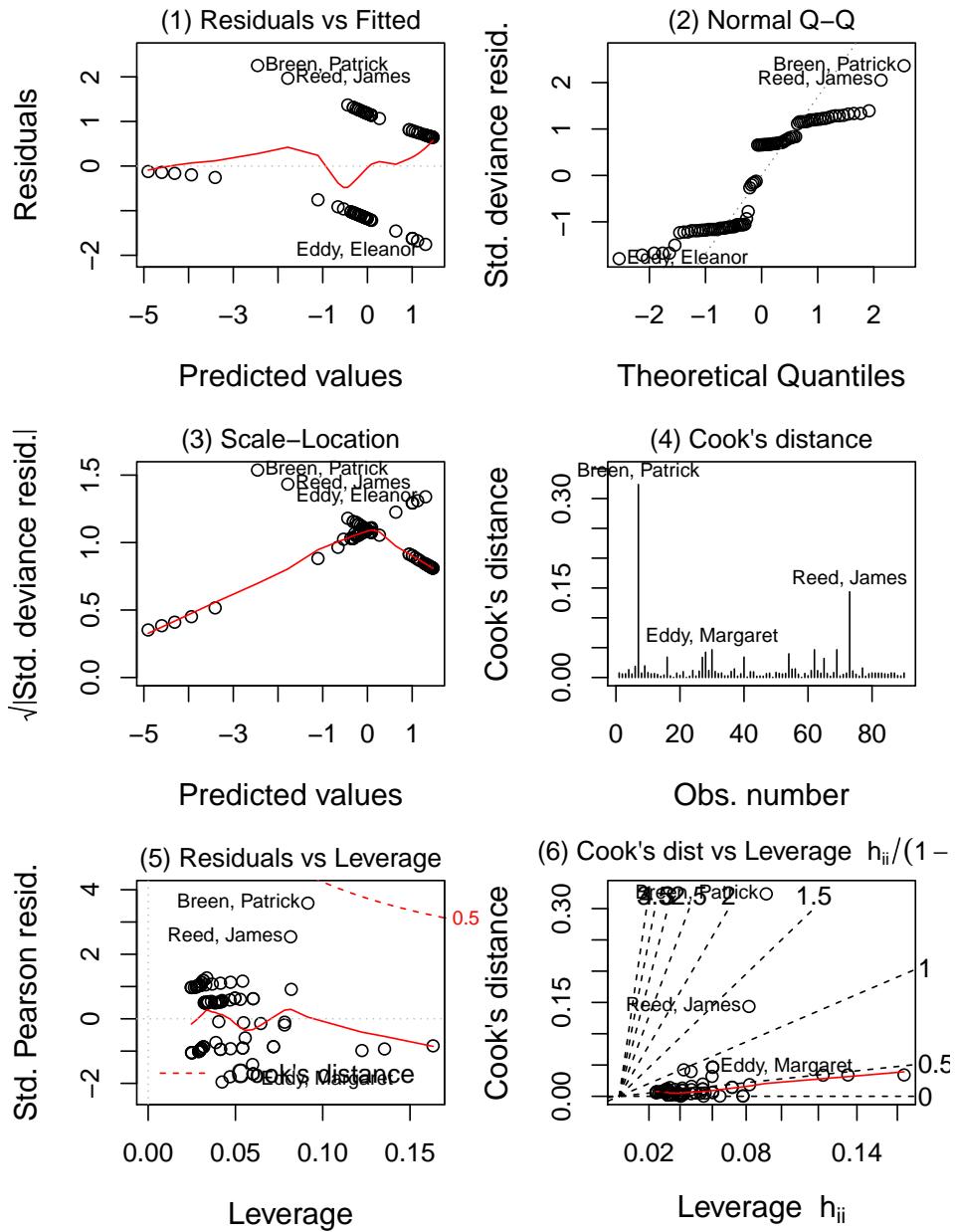


Figure 7.16: Diagnostic plots for a `glm` object, using the fitted model `donner.mod3` for the Donner Party data. Each plot shows some additional annotations or smoothed curves and labels observations considered noteworthy in terms of influence.
19:donner2\$plot

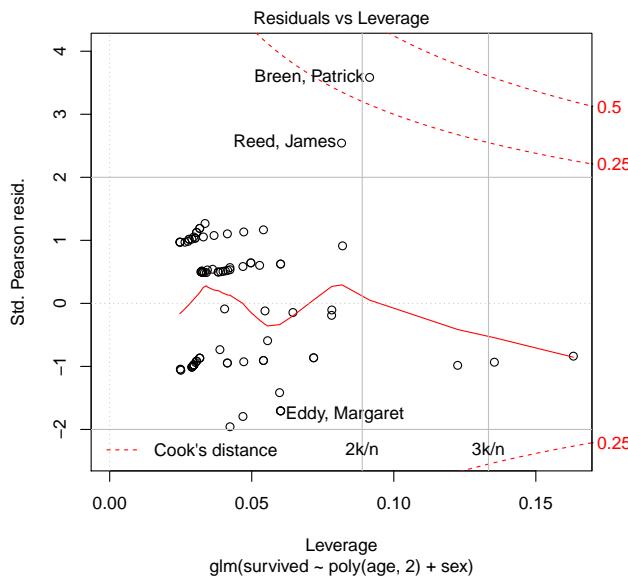


Figure 7.17: Residual vs. Leverage plot for the Donner data model. Horizontal and vertical reference lines show typical cutoff values for noteworthy residuals and leverage.
fig:donner-plots

The `summary()` method for the "infl" object prints those observations considered noteworthy on one or more of these statistics, as indicated by a "*" next to the value.

```
> summary(infl)

Potentially influential observations of
  glm(formula = survived ~ poly(age, 2) + sex, family = binomial,
      data = Donner) :

          dfb.1_ dfb.p(,2)_1 dfb.p(,2)_2 dfb.sxM1 dffit   cov.r   cook.d  hat
Breen, Patrick     0.08    0.65       0.56     0.23   0.69_*  0.93   0.32   0.09
Donner, Elizabeth -0.26   -0.34      -0.22     0.12   -0.40   1.15_*  0.03   0.14_*
Graves, Elizabeth C. -0.24   -0.37      -0.26     0.10   -0.42   1.20_*  0.03   0.16_*
```

The function `influencePlot()` in the `car` package gives a similar plot, but uses the size (area) of the plotting symbol to also show the value of Cook's D as shown in Figure 7.18. Like other diagnostic plots in `car`, it is considerably more general than illustrated here, because it allows for different `id.methods` to label noteworthy points, including `id.method="identify"` for interactive point identification by clicking with the mouse. The `id.n` argument works differently than with `plot()`, because it selects the most extreme `id.n` observations on *each* of the studentized residual, hat value and Cook's D, and labels all of these.

```
> library(car)
> res <- influencePlot(donner.mod3, id.col="blue", scale=8, id.n=2)
> text(x=c(2, 3)*k/n, y=-1.8, c("2k/n", "3k/n"))
```

Conveniently, `influencePlot()` returns a data frame containing the influence statistics for the points identified in the plot (`res` in the call above). We can combine this with the data values to help learn why these points are considered influential.

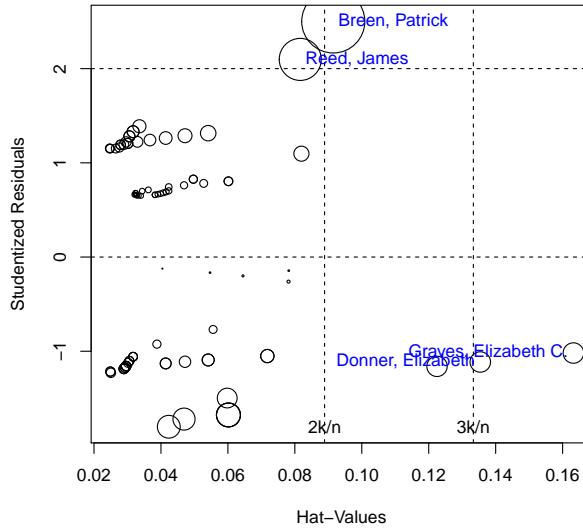


Figure 7.18: Influence plot (residual vs. leverage) for the Donner data model, showing Cook's D as the size of the bubble symbol. Horizontal and vertical reference lines show typical cutoff values for noteworthy residuals and leverage.^{fig:donnerz-inflplot}

```
> # show data together with diagnostics for influential cases
> idx <- which(rownames(Donner) %in% rownames(res))
> cbind(Donner[idx, 2:4], res)

      age sex survived StudRes     Hat   CookD
Breen, Patrick    51 Male     yes  2.501 0.09148 0.5688
Donner, Elizabeth 45 Female   no -1.114 0.13541 0.1846
Graves, Elizabeth C. 47 Female   no -1.019 0.16322 0.1849
Reed, James       46 Male     yes  2.098 0.08162 0.3790
```

We can see that Patrick Breen and James Reed²⁰ are unusual because they were both older men who survived, and have large positive residuals; Breen is the most influential by Cook's D, but this value is not excessively large. The two women were among the older women who died. They are selected here because they have the largest hat values, meaning they are unusual in terms of the distribution of age and sex, but they are not particularly influential in terms of Cook's D.

A related graphical display is the collection of index plots provided by `influenceIndexPlot()` in `car`, which plots various influence diagnostics against the observation numbers in the data. The `id.n` argument here works to label that number of the most extreme observations *individually* for each measure plotted. The following call produces Figure 7.19.

```
> influenceIndexPlot(donner.mod3, vars=c("Cook", "Studentized", "hat"),
+ id.n=4)
```

In our opinion, *separate* index plots are often less useful than combined plots such as the leverage-influence plot that shows residuals, leverage and Cook's D together. However, the `car`

²⁰Breen and Reed, both born in Ireland, were the leaders of their family groups. Among others, both kept detailed diaries of their experiences, from which most of the historical record derives. Reed was also the leader of two relief parties sent out to find rescue or supplies over the high Sierra mountains, so it is all the more remarkable that he survived.

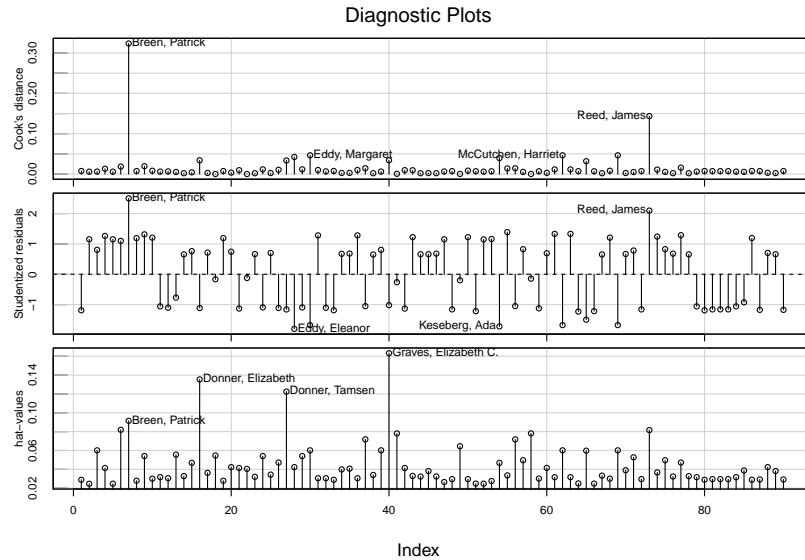


Figure 7.19: Index plots of influence measures for the *Donner* data model. The four most extreme observations on each measure are labeled.
fig:donner2-indexplot

version in Figure 7.19 does that too, and allows us to consider how unusual the labeled observations are both individually and in combination.



{ex:icu2}

EXAMPLE 7.13: Death in the ICU

In Example 7.11 we examined several models to account for death in the *ICU* data set. We continue this analysis here, with a focus on the simple main effects model, *icu.glm2*, for which the fitted logits were shown in Figure 7.15. For ease of reference, we restate that model here:

```
> icu.glm2 <- glm(died ~ age + cancer + admit + uncons,
+                     data=ICU, family=binomial)
```

The plot of residual vs. leverage for this model is shown in Figure 7.20.

```
> library(car)
> res <- influencePlot(icu.glm2, id.col="red", scale=8, id.cex=1.5, id.n=3)
```

Details for the cases identified in the figure are shown below, again using *rownames*(*res*) to select the relevant observations from the *ICU* data.

```
> idx <- which(rownames(ICU) %in% rownames(res))
> cbind(ICU[idx, c("died", "age", "cancer", "admit", "uncons")], res)

  died age cancer      admit uncons StudRes      Hat CookD
84   No  59      No Emergency    Yes -2.258 0.06781 0.3626
371  No  46      Yes Emergency   No -1.277 0.16408 0.2210
766  No  31      Yes Emergency   No -1.028 0.17062 0.1719
881  No  89      No Emergency   Yes -2.718 0.03081 0.4106
127  Yes 19      No Emergency   No  2.565 0.01679 0.2724
208  Yes 70      No Elective    Yes  1.662 0.29537 0.4568
380  Yes 20      No Emergency   No  2.548 0.01672 0.2668
```

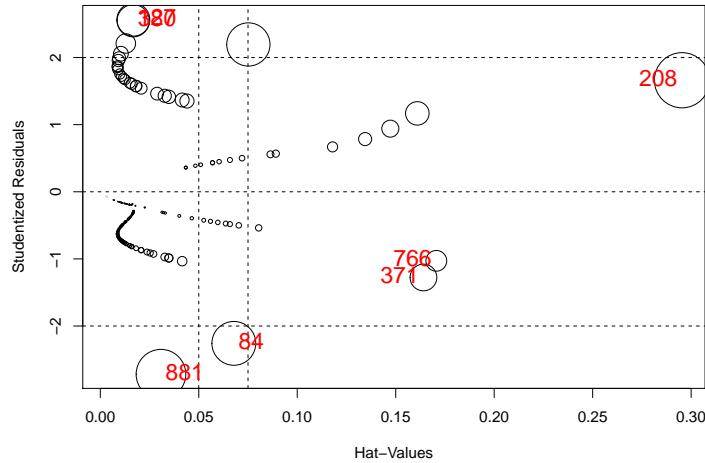


Figure 7.20: Influence plot for the main effects model for the ICU data fig:icu2-inflplot

None of the cases are particularly influential on the model coefficients overall: the largest Cook's D is only 0.45 for case 208. This observation also has the largest hat value. It is unusual on the predictors in this sample: a 70 year old man without cancer, admitted on an elective basis, who nonetheless died. However, this case is also highly unusual in having been unconscious on admission for an elective procedure, and signals that there might have been a coding error or other anomaly for this observation.

Another noteworthy observation identified here is case 881, an 89 year old male, admitted unconscious as an emergency; this case is poorly predicted because he survived. Similarly, two other cases (127, 380) with large studentized residuals are poorly predicted because they died, although they were young, did not have cancer, and conscious at admission. However, these cases have relatively small Cook's D values. From this evidence we might conclude that, case 208 bears further scrutiny, but none of these cases greatly affects the model, its coefficients, or interpretation.

For comparison with Figure 7.20, the related index plot of these measures is shown in Figure 7.21.

```
> influenceIndexPlot(icu.glm2, vars=c("Cook", "Studentized", "hat"), id.n=4)
```

Cook's D and DFFITS are *overall* measures of the total influence that cases have on the regression coefficients and fitted values respectively. It might be that some cases have a large impact on some individual regression coefficients, but don't appear particularly unusual in these aggregate measures.

One way to study this is to make plots of the $DFBETA_{ij}$ statistics. Such plots are not available (as far as we know) in R packages, but it is not hard to construct them from the result returned by `influence.measures()`. To do this, we select the appropriate columns from the `infmat` component returned by that function.

```
> infl <- influence.measures(icu.glm2)
> dfbetas <- data.frame(infl$infmat[,2:5])
> colnames(dfbetas) <- c("dfb.age", "dfb.cancer", "dfb.admit", "dfb.uncons")
> head(dfbetas)

dfb.age dfb.cancer dfb.admit dfb.uncons
```

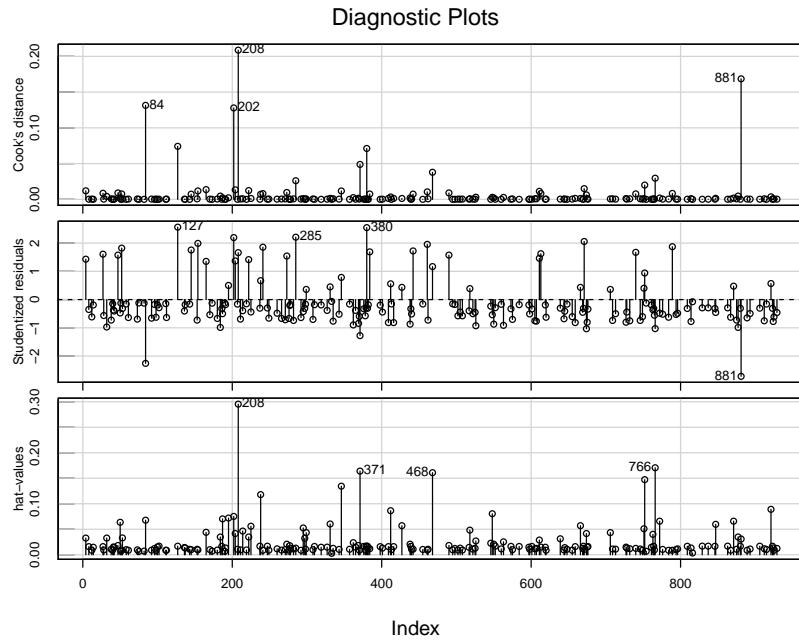


Figure 7.21: Index plots of influence measures for the `ICU` data model. The four most extreme observations on each measure are labeled.
fig:icu-infl-index

8	0.047340	0.013418	0.004067	0.009254
12	0.018988	0.018412	-0.004174	0.018106
14	-0.001051	0.014882	0.026278	0.005555
28	0.031562	0.018424	-0.001511	0.016640
32	-0.164084	0.003788	-0.036505	0.023488
38	-0.021525	0.016539	-0.011937	0.020803

To illustrate this idea, plotting an individual column of `dfbetas` using `type = "h"` gives an index plot against the observation number. This is shown in Figure 7.22 for the impact on the coefficient for age. The lines and points are colored blue or red according to whether the patient lived or died. Observations for which the $|DFBETA_{age}| > 0.2$ (an arbitrary value) are labeled.

```
> cols $\leftarrow$ ifelse (ICU$died=="Yes", "red", "blue")
> op  $\leftarrow$  par(mar=c(5,5,1,1)+.1)
> plot(dfbetas[,1], type = "h", col=cols,
+       xlab="Observation index",
+       ylab=expression(Delta * beta[Age]),
+       cex.lab=1.3)
> points(dfbetas[,1], col=cols)
> # label some points
> big  $\leftarrow$  abs(dfbetas[,1])  $>$  .25
> idx  $\leftarrow$  1:nrow(dfbetas)
> text(idx[big], dfbetas[big,1], label=rownames(dfbetas)[big],
+       cex=0.9, pos=ifelse(dfbetas[big,1]>0, 3, 1),
+       xpd=TRUE)
> abline(h=c(-.25, 0, .25), col="gray")
> par(op)
```

None of the labeled points here are a cause for concern, since the standardized DFBETAs are all

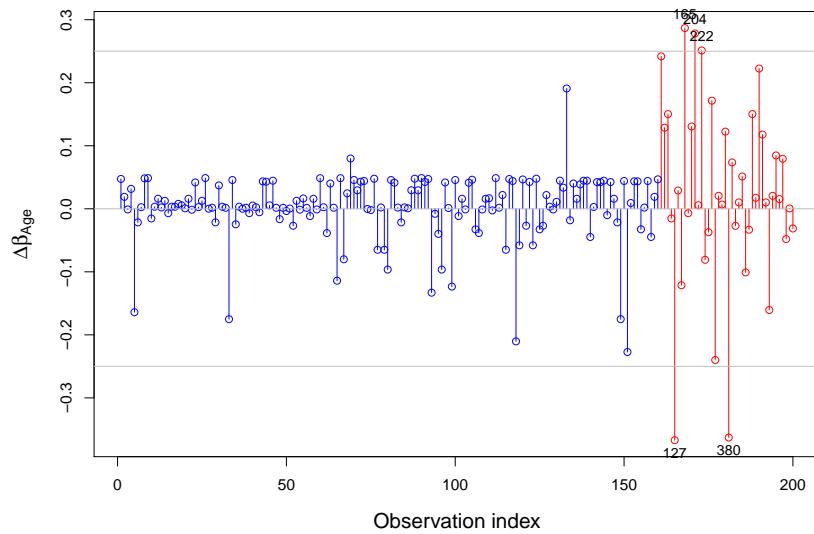


Figure 7.22: Index plot for DFBETA (Age) in the ICU data model. The observations are colored blue or red according to whether the patient lived or died.^{19:icuz-dbeta}

relatively small. However, the plot shows that patients who died have generally larger impacts on this coefficient.

An alternative to individual index plots is a scatterplot matrix, that shows the pairwise changes in the regression coefficients for the various predictors. Here we use `scatterplotMatrix()` from `car` that offers features for additional plot annotations, including identifying the most unusual points in each pairwise plot. In each off-diagonal panel, a 95% data ellipse and linear regression line helps to show the marginal relationship between the two measures and highlight why the labeled points are atypical in each plot.²¹

```
> scatterplotMatrix(dfbetas, smooth=FALSE, id.n=2,
+   ellipse=TRUE, levels=0.95, robust=FALSE,
+   diagonal="histogram",
+   groups=ICU$died, col=c("blue", "red"))
```



7.5.3 Other diagnostic plots

The graphical methods described in this section are relatively straight-forward indicators of the adequacy of a particular model, with a specified set of predictors, each expressed in a given way. More sophisticated methods have also been proposed, which focus on the need to include a particular predictor and whether its relationship is linear. These include the **component-plus-residual plot**, the **added-variable plot**, and the **constructed variable plot**, which are all analogous to techniques developed in OLS.

²¹This plot uses the `id.method="mahal"` method to label the most extreme observations according to the Mahalanobis distance of each point from the centroid in the plot.

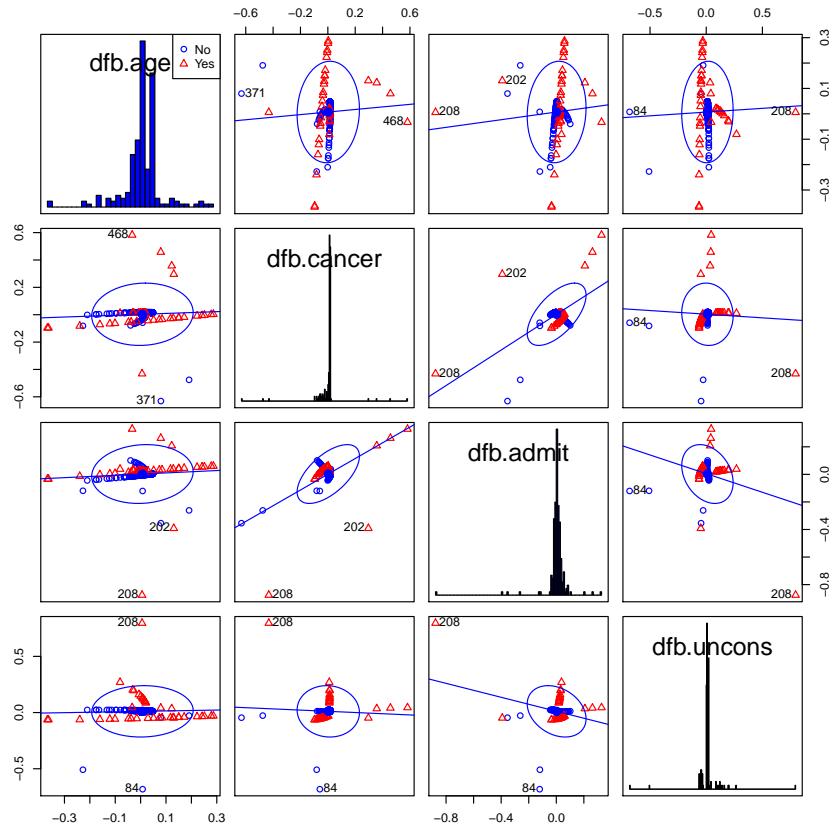


Figure 7.23: Scatterplot matrix for DFBETAs from the model for the ICU data. Those who lived or died are shown with blue circles and red triangles, respectively. The diagonal panels show histograms of each variable.
Fig.1icuz-abscatmat

7.5.3.1 Component-plus-residual plots

The **component-plus-residual plot** (also called a **partial residual plot**) proposed originally by Larsen and McCleary (1972) is designed to show whether a given quantitative predictor, x_j , included linearly in the model, actually shows a nonlinear relation, requiring transformation. The essential idea is to move the linear term for x_j back into the residual, by calculating the *partial residuals*,

$$r_j^* = r + \beta_j x_j$$

Then, a plot of r_j^* against x_j will have the same slope, β_j , as the full model including it among other predictors. However, any non-linear trend will be shown in the pattern of the points, usually aided by a smoothed non-parametric curve.

As adapted to logistic regression by Landwehr *et al.* (1984), the partial residual for variable x_j is defined as

$$r_j^* = V^{-1}r + \beta_j x_j$$

The partial residual plot is then a plot of r_j^* against x_j , possibly with the addition of a smoothed lowess curve (Fowlkes, 1987) and a linear regression line to aid interpretation. The linear regression of the partial residuals on x_j has the same slope, β_j , as in the full model.

If x_j affects the binary response linearly, the plot should be approximately linear with a slope approximately equal to β_j . A nonlinear plot suggests that x_j needs to be transformed, and the shape of the relation gives a rough guide to the required transformation. For example, a parabolic shape would suggest a term in x_j^2 . These plots complement the conditional data plots described earlier (Section 7.3.1), and are most useful when there are several quantitative predictors, so that it is more convenient and sensible to examine their relationships individually.

The **car** package implements these plots in the `crPlots()` and `crPlot()` functions. They also work for models with factor predictors (using parallel boxplots for the factor levels) but not for those with interaction terms.

EXAMPLE 7.14: Donner Party

In Example 7.12, we fit several models for the Donner Party data, and we recall two here to illustrate component-plus-residual plots. Both assert additive effects of age and sex, but the model `donner.mod3` allows a quadratic effect of age.

```
> donner.mod1 <- glm(survived ~ age + sex, data=Donner, family=binomial)
> donner.mod3 <- glm(survived ~ poly(age, 2) + sex, data=Donner, family=binomial)
```

Had we not made exploratory plots earlier (Example 7.12), and naively fit only the linear model in age, `donner.mod1`, we could use `crPlots()` to check for a non-linear relationship of survival with age as follows, giving Figure 7.24.

```
> crPlots(donner.mod1, ~age, id.n=2)
```

The smoothed loess curve in this plot closely resembles the trend we saw in the conditional plot for age by sex (Figure 7.12), suggesting the need to include a non-linear term for age. The points identified in this plot, by default, are those with either the most extreme x values (giving them high leverage) or the largest absolute Pearson residuals in the full model. The four structured bands of points in the plot correspond to the combinations of sex and survival.

For comparison, you can see the result of allowing for a non-linear relationship in age in a partial residual plot for the model `donner.mod3` that includes the effect `poly(age, 2)` for age. Note that the syntax of the `crPlots()` function requires that you specify a *term* in the model, rather than just a predictor variable.

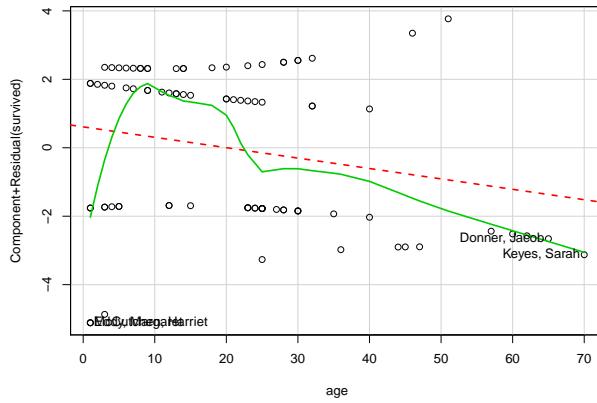


Figure 7.24: Component-plus-residual plot for the simple additive linear model, `donner.mod1`. The dashed red line shows the slope of age in the full model; the smoothed green curve shows a loess fit with span = 0.5.^{fig:donner-cr1}

```
> crPlots(donner.mod3, ~poly(age, 2), id.n=2)
```

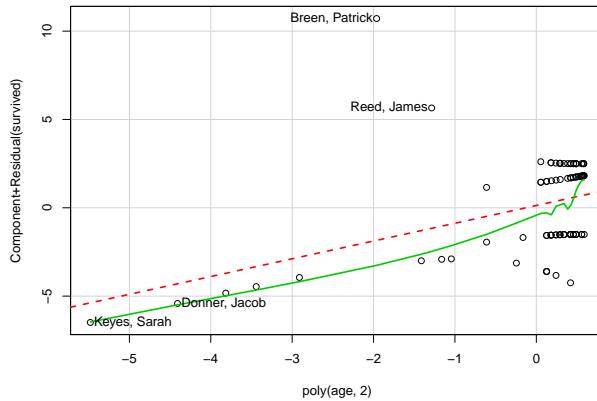


Figure 7.25: Component-plus-residual plot for the non-linear additive model, `donner.mod3`^{fig:donner-cr2}

Except possibly at the extreme right, this plot (Figure 7.25) shows no indication of a non-linear relationship.



7.5.3.2 Added-variable plots

Added-variable plots (Cook and Weisberg, 1999, Wang, 1985) (also called *partial-regression plots*) are another important tool for diagnosing problems in logistic regression and other linear or generalized linear models. These are essentially plots, for each x_i , of an adjusted response, $y_i^* = y | \text{others}_i$, against an adjusted predictor, $x_i^* = x_i | \text{others}_i$, where $\text{others}_i = X \setminus x_i \equiv X^{(-i)}$

indicates all other predictors excluding x_i . As such, they show the *conditional* relationship between the response and the predictor x_i , controlling for, or adjusting for, all other predictors. Here, y_i^* and x_i^* represent respectively the residuals from the regressions of y and x_i on all the other x s excluding x_i .

It might seem from this description that each added-variable plot requires two additional auxiliary logistic regressions to calculate the residuals y_i^* and x_i^* . However, Wang (1985) showed that the added-variable plot may be constructed by following the logistic regression for the model $y \sim \mathbf{X}^{(-i)}$ with one weighted least squares regression of x_i on $\mathbf{X}^{(-i)}$ to find the residual part, x_i^* , of x not predicted by the other regressors.

Let r be the vector of Pearson residuals from the initial logistic fit of y on the variables in $\mathbf{X}^{(-i)}$, and let \mathbf{H} and $\mathbf{V} = \text{diag}[\hat{p}(1 - \hat{p})]$ be the hat matrix and V matrix from this analysis. Then, the added-variable plot is a scatterplot of the residuals r against the x_i -residuals,

$$x_i^* = (\mathbf{I} - \mathbf{H})\mathbf{V}^{1/2}\mathbf{x} .$$

There are several important uses of added-variable plots:

First, *marginal* plots of the response variable y against the predictor variables x_i can conceal or misrepresent the relationships in a model including several predictors together due to correlations or associations among the predictors. This problem is compounded by the fact that graphical methods for discrete responses (boxplots, mosaic plots) cannot easily show influential observations or non-linear relationships. Added-variable plots solve this problem by plotting the residuals, $y_i^* = y | \text{others}_i$, which are less discrete than the marginal responses in y .

Second, the numerical measures and graphical methods for detecting influential observations described earlier in this section are based on the idea of *single-case deletion*, comparing coefficients or fitted values for the full data, with those that result from deleting each case in turn. Yet, it is well-known (Lawrance, 1995), that sets of two (or more) observations can have *joint influence*, that greatly exceeds their individual influential. Similarly, the influence of one discrepant point can be offset by another influential point in an opposite direction, a phenomenon called *masking*. The main cases of joint influence are illustrated in Figure 7.26. Added-variable plots, showing the partial regression for one predictor controlling all others can make such cases visually apparent.

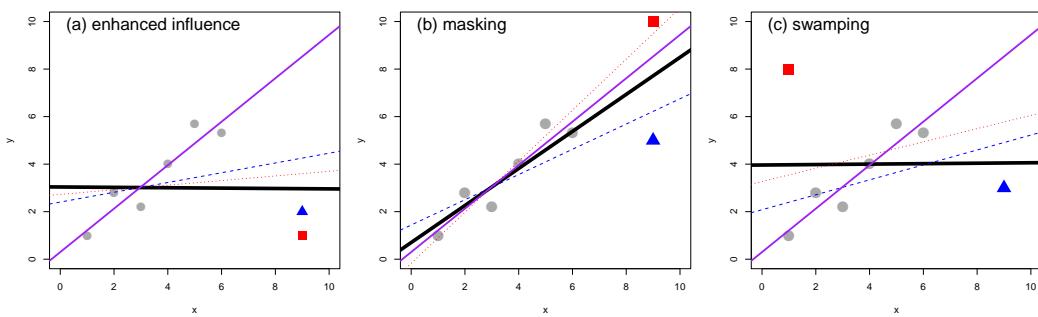


Figure 7.26: Jointly influential points in regression models. In each panel, the thick black line shows the regression of y on x using all the data points. The solid purple line shows the regression deleting *both* the red and blue points and the broken and dotted lines show the regression retaining only the point in its color in addition to the constant gray points. (a) Two points whose joint influence enhance each other; (b) two points where the influence of one is masked by that of the other; (c) two points whose combined influence greatly exceeds the effect of either one individually.

{fig:joint}

Finally, given a tentative model using predictors \mathbf{x} , the added-variable plot for another regressor, z can provide a useful visual assessment of its additional contribution. An overall test could be based

on the difference in G^2 for the enlarged model $\text{logit}(\mathbf{p}) = \mathbf{X}\boldsymbol{\beta} + \gamma z$, compared to the reduced model $\text{logit}(\mathbf{p}) = \mathbf{X}\boldsymbol{\beta}$. But the added-variable plot shows whether the evidence for including z is spread throughout the sample or confined to a small subset of observations. The regressor z may be a new explanatory variable, or a higher-order term for variable(s) already in the model.

The `car` package implements these plots with the function `avPlot()` for a single term and `avPlots()` for all terms in a linear or generalized linear model, as shown in the example(s) below. See <http://www.datavis.ca/gallery/animation/duncanAV/> for an animated graphic showing the transition between a marginal plot of the relationship of y to x and the added-variable plot of y^* to x^* for the case of multiple linear regression with a quantitative response.

{ex:donner4}

EXAMPLE 7.15: Donner Party

The simple additive model `donner.mod1` for the Donner Party data can be used to illustrate some features of added-variable plots. In the call to `avPlots()` below, we use color to distinguish those who survived vs. died, shape to distinguish men from women.

```
> col <- ifelse(Donner$survived=="yes", "blue", "red")
> pch <- ifelse(Donner$sex=="Male", 16, 17)
> avPlots(donner.mod1, id.n=2, col=col, pch=pch, col.lines="darkgreen")
```

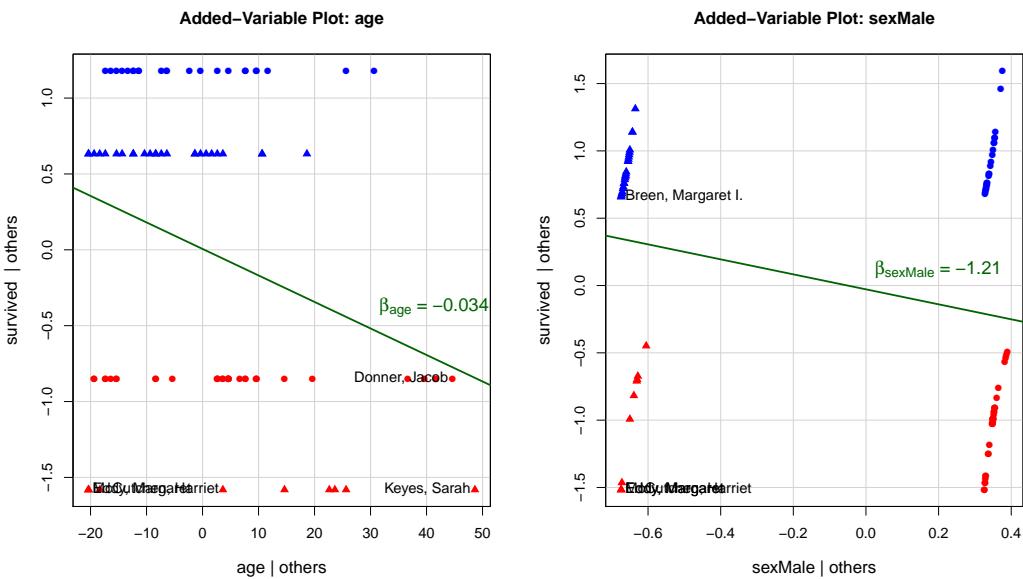


Figure 7.27: Added-variable plots for age (left) and sex (right) in the Donner Party main effects model. Those who survived are shown in blue; those who died in red. Men are plotted with filled circles; women with filled triangles. [Fig:donner4-avp]

These plots have the following properties:

1. The slope in the simple regression of y_i^* on x_i^* is the same as the partial coefficient β_i in the full multiple regression model including both predictors here (or all predictors in general).
2. The residuals from this regression line are the same as the residuals in the full model.
3. Because the response, `survived`, is binary, the vertical axis y_{age}^* in the left panel for `age` is the part of the logit for survival that cannot be predicted from `sex`. Similarly, the vertical axis

in the right panel is the part of survival that cannot be predicted from age. This property allows the clusters of points corresponding to discrete variables to be seen more readily, particularly if they are distinguished by visual attributes such as color and shape, as in Figure 7.27.



{ex:icu3}

EXAMPLE 7.16: Death in the ICU

We illustrate some of the uses of added-variable plots using the main effects model, `icu.glm2`, predicting death in the ICU from the variables `age`, `cancer`, `admit` and `uncons`.

To see why marginal plots of the discrete response against each predictor are often unrevealing for the purpose of model assessment, consider the collection of plots in Figure 7.28 showing the default plots (spineplots) for the factor response, `died` against each predictor. These show the marginal distribution of each predictor by the widths of the bars, and highlight the proportion who died by color. Such plots are useful for some purposes, but not for assessing the adequacy of the fitted model.

```
> op <- par(mfrow=c(2,2), mar=c(4,4,1,2.5)+.1, cex.lab=1.4)
> plot(died ~ age, data=ICU, col=c("lightblue", "red"))
> plot(died ~ cancer, data=ICU, col=c("lightblue", "red"))
> plot(died ~ admit, data=ICU, col=c("lightblue", "red"))
> plot(died ~ uncons, data=ICU, col=c("lightblue", "red"))
> par(op)
```

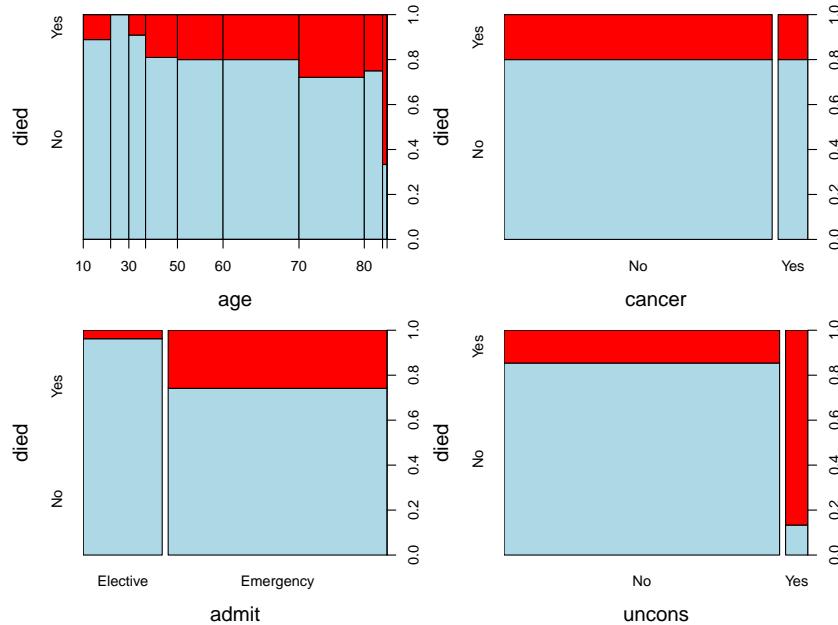


Figure 7.28: Marginal plots of the response `died` against each of the predictors in the model `icu.glm2` for the `ICU` data

The added-variable plot for this model is shown in Figure 7.29. In each plot, the solid red line shows the partial slope, β_j for the focal predictor, controlling for all others.

```
> pch <- ifelse(ICU$died=="No", 1, 2)
> avPlots(icu.glm2, id.n=2, pch=pch, cex.lab=1.3)
```

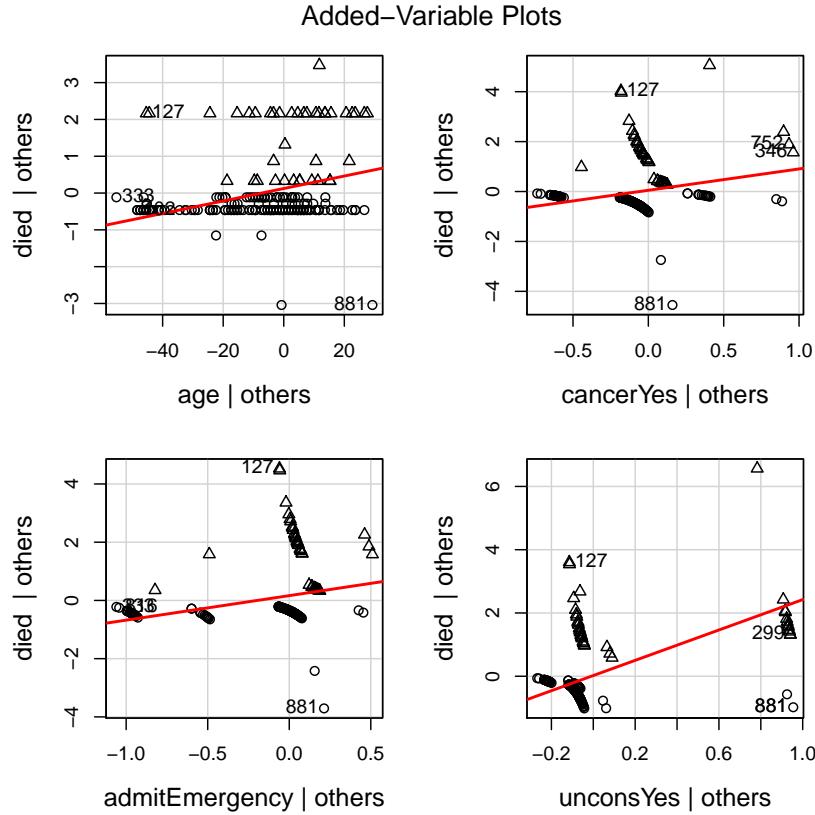


Figure 7.29: Added-variable plots for the predictors in the model for the ICU data. Those who died and survived are shown by triangles (Δ) and circles (\circ) respectively.
FIG:ICU3-avp1

The labeled points in each panel use the default `id.method` for `avPlots()`, selecting those with either large absolute model residuals or extreme x_i^* residuals, given all other predictors. Cases 127 and 881, identified earlier as influential stand out in all these plots.

Next, we illustrate the use of added-variable plots for checking the effect of influential observations on the decision to include an additional predictor in some given model. In the analysis of the *ICU* data using model selection methods, the variable `systolic` (systolic blood pressure at admission) was nominated by several different procedures. Here we take a closer look at the evidence for inclusion of this variable in a predictive model. We fit a new model adding `systolic` to the others and test the improvement with a likelihood ratio test:

```
> icu.glm2a <- glm(died ~ age + cancer + admit + uncons + systolic,
+                      data=ICU, family=binomial)
> anova(icu.glm2, icu.glm2a, test="Chisq")
```

Analysis of Deviance Table

```
Model 1: died ~ age + cancer + admit + uncons
Model 2: died ~ age + cancer + admit + uncons + systolic
  Resid. Df Resid. Dev Df Deviance Pr(>Chi)
```

```

1      195      139
2      194      136  1     3.52    0.061 .
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

So, the addition of systolic blood pressure is nearly significant at the conventional $\alpha = 0.05$ level. The added-variable plot for this variable in Figure 7.30 shows the strength of evidence for its contribution, above and beyond the other variables in the model, as well as the partial leverage and influence of individual points.

```
> avPlot(icu.glm2a, "systolic", id.n=3, pch=pch)
```

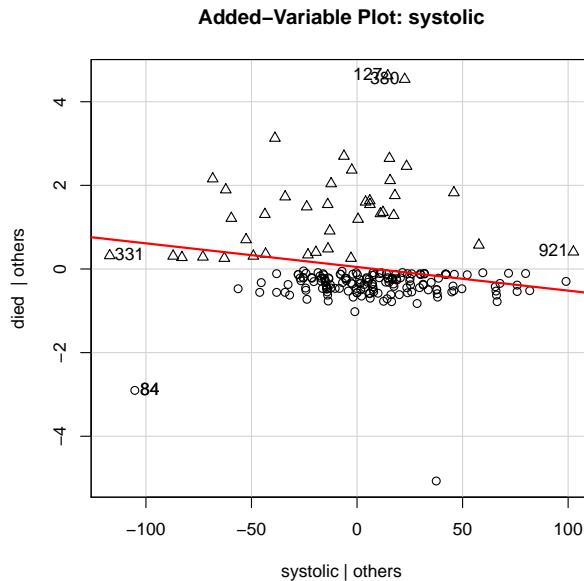


Figure 7.30: added-variable plot for the effect of adding systolic blood pressure to the main effects model for the ICU data.^{fig:icu3-avp2}

In this plot, cases 331 and 921 have high partial leverage, but they are not influential. Case 84, however, has high leverage and a large residual, so is possibly influential on the evidence for inclusion of `systolic` in the model. Note also that the partial regression line in this plot nicely separates nearly all the patients who died from those who survived.



TODO: This chapter is quite long. Could start a new chapter here, but that would require re-numbering file names, or introduce Ch07a.

7.6 Polytomous response models

{sec:logist-poly} Polytomous response data arise when the outcome variable, Y , takes on $m > 2$ discrete values. For example, (a) patients may record that their improvement after treatment is “none,” “some” or “marked;” (b) high school students may choose a general, vocational or academic program; (c) women’s labor force participation may be recorded in a survey as not working outside the home,

working part-time, or working full-time; (d) Canadian voters may express a preference for the Conservative, Liberal, NDP, Green party. These response categories may be considered ordered or simply nominal.

In this situation, there are several different ways to model the response probabilities. Let $\pi_{ij} \equiv \pi_j(\mathbf{x}_i)$ be the probability of response j for case or group i , given the predictors \mathbf{x}_i . Because $\sum_j \pi_{ij} = 1$, only $m - 1$ of these probabilities are independent. The essential idea here is to construct a model for the polytomous (or multinomial) response composed of $m - 1$ logit comparisons among the response categories in a similar way to how factors are treated in the predictor variables.

The simplest approach uses the ***proportional odds model***, described in Section 7.6.1. This model applies only when the response is ordinal (as in improvement after therapy) *and* an additional assumption (the proportional odds assumption) holds. This model can be fit using `polr()` in the MASS package, `lrm()` in the rms package, and `vglm()` in VGAM.

However, if the response is purely nominal (e.g., vote Conservative, Liberal, NDP, Green), or if the proportional odds assumption is untenable, another particularly simple strategy is to fit separate models to a set of $m - 1$ ***nested dichotomies*** derived from the polytomous response (described in Section 7.6.3). This method allows you to resolve the differences among the m response categories into independent statistical questions (similar to orthogonal contrasts in ANOVA). For example, for women's labor force participation, it might be substantively interesting to contrast not working vs. (part-time and full-time) and then part-time vs. full-time for women who are working. You fit such nested dichotomies by running the $m - 1$ binary logit models and combining the statistical results.

The most general approach is the ***generalized logit model***, also called the ***multinomial logit model***. This model fits *simultaneously* the $m - 1$ simple logit models against a baseline or reference category, for example, the last category, m . With a 3-category response, there are two generalized logits, $L_{i1} = \log(\pi_{i1}/\pi_{i3})$ and $L_{i2} = \log(\pi_{i2}/\pi_{i3})$, contrasting response categories 1 and 2 against category 3. In this approach, it doesn't matter which response category is chosen as the baseline, because all pairwise comparisons can be recovered from whatever is estimated. This model is conveniently fitted using `multinom()` in nnet.

7.6.1 Ordinal Response: Proportional Odds Model

For an ordered response Y , with categories $j = 1, 2, \dots, m$, the ordinal nature of the response can be taken into account by forming logits based on the $m - 1$ adjacent category cutpoints between successive categories. That is, if the cumulative probabilities are

$$\Pr(Y \leq j | \mathbf{x}) = \pi_1(\mathbf{x}) + \pi_2(\mathbf{x}) + \dots + \pi_j(\mathbf{x}) ,$$

then the ***cumulative logit*** for category j is defined as

$$L_j \equiv \text{logit}[\Pr(Y \leq j | \mathbf{x})] = \log \frac{\Pr(Y \leq j | \mathbf{x})}{\Pr(Y > j | \mathbf{x})} = \log \frac{\Pr(Y \leq j | \mathbf{x})}{1 - \Pr(Y \leq j | \mathbf{x})} \quad (7.13) \quad \text{(eq:cumlogit)}$$

for $j = 1, 2, \dots, m - 1$.

In our running example of responses to arthritis treatment, the actual response variable is `Improved`, with ordered levels "None" < "Some" < "Marked". In this case, the cumulative logits would be defined as

$$L_1 = \log \frac{\pi_1(\mathbf{x})}{\pi_2(\mathbf{x}) + \pi_3(\mathbf{x})} = \text{logit} (\text{None vs. [Some or Marked]})$$

$$L_2 = \log \frac{\pi_1(\mathbf{x}) + \pi_2(\mathbf{x})}{\pi_3(\mathbf{x})} = \text{logit} ([\text{None or Some}] \text{ vs. Marked}) ,$$

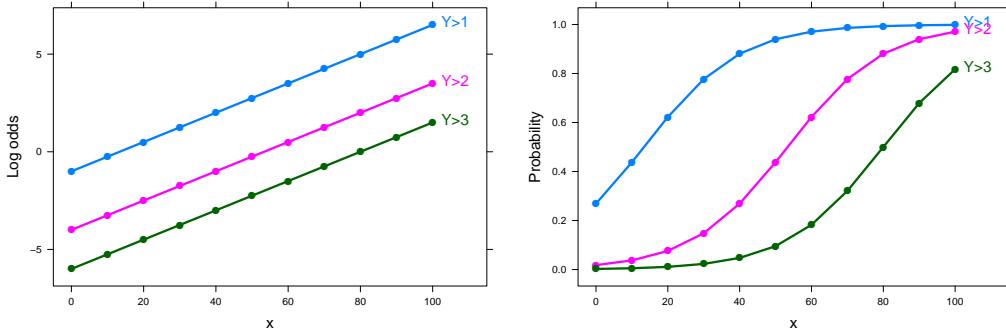


Figure 7.31: Proportional odds model for an ordinal response. The model assumes equal slopes for the cumulative response logits. Left: logit scale; right: probability scale.
 {fig:podds}

where \mathbf{x} represents the predictors (sex, treatment and age).

The **proportional odds model** (PO) (McCullagh, 1980) proposes a simple and parsimonious account of these effects, where the predictors in (\mathbf{x}) are constrained to have the same slopes for all cumulative logits,

$$\{eq:propodds\} \quad L_j = \alpha_j + \mathbf{x}^\top \boldsymbol{\beta} \quad j = 1, \dots, m-1. \quad (7.14)$$

That is, the effect of the predictor x_i is the same, β_i , for all cumulative logits. The cumulative logits differ only in their intercepts. In this formulation, the $\{\alpha_j\}$ increase with j , because $\Pr(Y \leq j | \mathbf{x})$ increases with j for fixed \mathbf{x} .²² Figure 7.31 portrays the PO model for a single quantitative predictor x with $m = 4$ response categories.

The name “proportional odds” stems from the fact that under Eqn. (7.14), for fixed \mathbf{x} , the cumulative log odds (logits) for categories j and j' are constant, $(\alpha_j - \alpha_{j'})$, so the odds, $\exp(\alpha_j - \alpha_{j'})$ have a constant ratio, or are proportional. Similarly, the ratio of the cumulative odds of making a response $Y \leq j$ at values of the predictors $\mathbf{x} = \mathbf{x}_1$ are $\exp((\mathbf{x}_1 - \mathbf{x}_2)^\top \boldsymbol{\beta})$ times the odds of this response at $\mathbf{x} = \mathbf{x}_2$, so the log cumulative odds ratio is proportional to the difference between \mathbf{x}_1 and \mathbf{x}_2 .

7.6.1.1 Latent variable interpretation

For a binary response, an alternative motivation for logistic regression regards the relation of the observed Y as arising from a continuous, unobserved, (latent) response variable, ξ representing the propensity for a “success” (1) rather than “failure” (0). The latent response is assumed to be linearly related to the predictors \mathbf{x} according to

$$\{eq:latent\} \quad \xi_i = \alpha + \mathbf{x}_i^\top \boldsymbol{\beta} + \epsilon_i = \alpha + \beta_1 x_{i1} + \dots + \beta_p x_{ip} + \epsilon_i \quad (7.15)$$

However, we can only observe $Y_i = 1$ when ξ_i passes some threshold, that with some convenient scaling can be taken as $\xi_i > 0 \implies Y_i = 1$.²³

²²Some authors and some software describe the PO model in terms of logit [$\Pr(Y > j | \mathbf{x})$], so the signs and order of the intercepts, α_j are reversed.

²³The latent variable derivation of logistic regression (and the related probit model) was fundamental in the history of statistical methods for discrete response outcomes. An early example was Thurstone's (1927) *Law of comparative judgment* designed to account for psychological preference by assuming an underlying latent continuum of “hedonic values.” Similarly, the probit model arose from dose-response studies in toxicology (??) where the number killed by some chemical agent was related to its’ type, dose or concentration. The idea of a latent variable was also at the heart of the development of factor

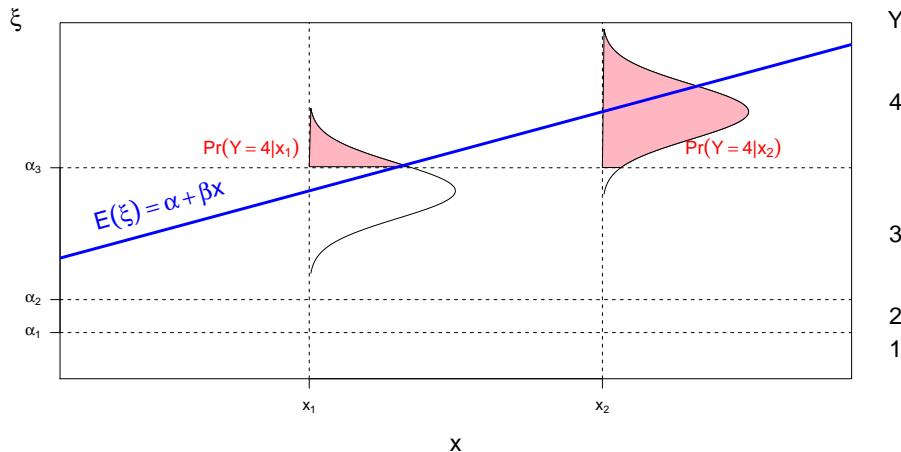


Figure 7.32: Latent variable representation of the proportional odds model for $m = 4$ response categories and a single quantitative predictor, x . *Source:* Adapted from Fox (2008, Fig 14.10), using code provided by John Fox.

{fig:latent}

The latent variable motivation extends directly to an ordinal response under the PO model. We now assume that there is a set of $m - 1$ thresholds, $\alpha_1 < \alpha_2 < \dots < \alpha_{m-1}$ for the latent variable ξ_i in Eqn. (7.15) and we observe

$$Y_i = j \quad \text{if} \quad \alpha_{j-1} < \xi_i \leq \alpha_j ,$$

with appropriate modifications to the inequalities at the end points.

This is illustrated in Figure 7.32 for a response with $m = 4$ ordered categories and a single quantitative predictor, x . The observable response Y categories are shown on the right vertical axis, and the corresponding latent continuous variable ξ on the left axis together with the thresholds $\alpha_1, \alpha_2, \alpha_3$. The (conditional) logistic distribution of ξ is shown at two values of x , and the shaded areas under the curve give the conditional probabilities $\Pr(Y = 4 | x_i)$ for the two values x_1 and x_2 .

7.6.1.2 Fitting the proportional odds model

As mentioned earlier, there are a number of different R packages that provide facilities for fitting the PO model. These have somewhat different capabilities for reporting results, testing hypotheses and plotting, so we generally use `polr()` in the MASS package, except where other packages offer greater convenience.

Unless the response variable has numeric values, it is important to ensure that it has been defined as an *ordered* factor (using `ordered()`). In the *Arthritis* data, the response, `Improved` was setup this way, as we can check by printing some of the values.²⁴

```
> head(Arthritis$Improved, 8)
[1] Some    None    None    Marked  Marked  Marked  None    Marked
Levels: None < Some < Marked
```

analysis **TODO:** citation? and latent class analysis (??) was developed to treat the problem of classifying individuals into discrete latent classes from fallible measurements. See ? for a useful overview of latent variable models in the social sciences.

²⁴As an unordered factor, the levels would be treated as ordered alphabetically, i.e., Marked, None, Some.

We fit the main effects model for the ordinal response using `polr()` as shown below. We also specify `Hess=TRUE` to have the function return the observed information matrix (called the Hessian), that is used in other operations to calculate standard errors.

```
> arth.polr <- polr(Improved ~ Sex + Treatment + Age,
+                      data=Arthritis, Hess=TRUE)
> summary(arth.polr)

Call:
polr(formula = Improved ~ Sex + Treatment + Age, data = Arthritis,
      Hess = TRUE)

Coefficients:
            Value Std. Error t value
SexMale     -1.2517   0.5464  -2.29
TreatmentTreated 1.7453   0.4759   3.67
Age         0.0382   0.0184   2.07

Intercepts:
            Value Std. Error t value
None|Some    2.532  1.057    2.395
Some|Marked  3.431  1.091    3.144

Residual Deviance: 145.46
AIC: 155.46
```

The output from the `summary()` method, shown above, gives the estimated coefficients (β) and intercepts (α_j) labeled by the cutpoint on the ordinal response. It provides standard errors and t -values ($\beta_i/SE(\beta_i)$), but no significance tests or p -values.

```
> library(car)
> Anova(arth.polr)

Analysis of Deviance Table (Type II tests)

Response: Improved
          LR Chisq Df Pr(>Chisq)
Sex           5.69   1   0.01708 *
Treatment     14.71   1   0.00013 ***
Age           4.57   1   0.03251 *
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

7.6.1.3 Testing the proportional odds assumption

The simplicity of the PO model is achieved only when the proportional odds model holds for a given data set. In essence, a test of this assumption involves a contrast between the PO model and a generalized logit NPO model that allows different effects (slopes) of the predictors across the response categories:

$$\{eq:po\} \quad \text{PO : } L_j = \alpha_j + \mathbf{x}^T \boldsymbol{\beta} \quad j = 1, \dots, m-1 \quad (7.16)$$

$$\{eq:npo\} \quad \text{NPO : } L_j = \alpha_j + \mathbf{x}^T \boldsymbol{\beta}_j \quad j = 1, \dots, m-1 \quad (7.17)$$

The most general test involves fitting both models and testing the difference in the residual deviance by a likelihood ratio test or using some other measure (such as AIC) for model comparison. The PO model (Eqn. (7.16)) has $(m-1) + p$ parameters, while the NPO model (Eqn. (7.17)) has $(m-1)(1+p) = m(1+p)$ parameters, which may be difficult to fit if this is large relative to the number of observations. An intermediate model, the *partial proportional odds model* (Peterson

and Harrell, 1990) allows one subset of predictors, \mathbf{x}_{po} , to satisfy the proportional odds assumption (equal slopes), while the remaining predictors \mathbf{x}_{npo} have slopes varying with the response level:

$$\text{PPO : } L_j = \alpha_j + \mathbf{x}_{po}^\top \boldsymbol{\beta} + \mathbf{x}_{npo}^\top \boldsymbol{\beta}_j \quad j = 1, \dots, m-1 . \quad (7.18)$$

In R, the PO and NPO models can be readily contrasted by fitting them both using `vglm()` in the VGAM package. This defines the cumulative family of models and allows a `parallel` option. With `parallel=TRUE`, this is equivalent to the `polr()` model, except that the signs of the coefficients are reversed.

```
> library(VGAM)
> arth.po <- vglm(Improved ~ Sex + Treatment + Age, data=Arthritis,
+                     family = cumulative(parallel=TRUE))
> arth.po

Call:
vglm(formula = Improved ~ Sex + Treatment + Age, family = cumulative(parallel = TRUE),
      data = Arthritis)

Coefficients:
(Intercept):1      (Intercept):2          SexMale
              2.531990        3.430988       1.251671
TreatmentTreated           Age
              -1.745304       -0.038163

Degrees of Freedom: 168 Total; 163 Residual
Residual deviance: 145.46
Log-likelihood: -72.729
```

The more general NPO model can be fit using `parallel=FALSE`.

```
> arth.npo <- vglm(Improved ~ Sex + Treatment + Age, data=Arthritis,
+                     family = cumulative(parallel=FALSE))
> arth.npo

Call:
vglm(formula = Improved ~ Sex + Treatment + Age, family = cumulative(parallel = FALSE),
      data = Arthritis)

Coefficients:
(Intercept):1      (Intercept):2          SexMale:1
              2.618539        3.431175       1.509827
SexMale:2 TreatmentTreated:1 TreatmentTreated:2
              0.866434       -1.836929      -1.704011
Age:1           Age:2
              -0.040866       -0.037294

Degrees of Freedom: 168 Total; 160 Residual
Residual deviance: 143.57
Log-likelihood: -71.787
```

The VGAM package defines a `coef()` method that can print the coefficients in a more readable matrix form giving the category cutpoints:

```
> coef(arth.po, matrix=TRUE)

            logit(P[Y<=1])  logit(P[Y<=2])
(Intercept)      2.531990      3.430988
SexMale         1.251671      1.251671
TreatmentTreated -1.745304     -1.745304
Age             -0.038163     -0.038163
```

```
> coef(arth.npo, matrix=TRUE)

      logit(P[Y<=1]) logit(P[Y<=2])
(Intercept)        2.618539     3.431175
SexMale           1.509827     0.866434
TreatmentTreated -1.836929    -1.704011
Age                -0.040866   -0.037294
```

In most cases, nested models can be tested using an `anova()` method, but the VGAM package has not implemented this for "vglm" objects. Instead, it provides an analogous function, `lrtest()`:

```
> VGAM:::lrtest(arth.npo, arth.po)

Likelihood ratio test

Model 1: Improved ~ Sex + Treatment + Age
Model 2: Improved ~ Sex + Treatment + Age
#Df LogLik Df Chisq Pr(>Chisq)
1 160    -71.8
2 163    -72.7  3  1.88      0.6
```

The LR test can be also calculated as “manually” shown below using the difference in residual deviance for the two models.

```
> tab <- cbind(
+   Deviance = c(deviance(arth.npo), deviance(arth.po)),
+   df = c(df.residual(arth.npo), df.residual(arth.po)))
+ )
> tab <- rbind(tab, diff(tab))
> rownames(tab) <- c("GenLogit", "PropOdds", "LR test")
> tab <- cbind(tab, pvalue=1-pchisq(tab[,1], tab[,2]))
> tab

      Deviance   df   pvalue
GenLogit 143.5741 160 0.81966
PropOdds 145.4579 163 0.83435
LR test    1.8838   3 0.59686
```

The `vglm()` can also fit partial proportional odds models, by specifying a formula giving the terms for which the PO assumption should be taken as TRUE or FALSE. Here we illustrate this using `parallel=FALSE ~ Sex`, to fit separate slopes for males and females, but parallel lines for the other predictors. The same model would be fit using `parallel=TRUE ~ Treatment + Age`.

```
> arth.ppo <- vglm(Improved ~ Sex + Treatment + Age, data=Arthritis,
+   family = cumulative(parallel=FALSE ~ Sex))
> coef(arth.ppo, matrix=TRUE)

      logit(P[Y<=1]) logit(P[Y<=2])
(Intercept)        2.542452     3.615561
SexMale           1.483336     0.867362
TreatmentTreated -1.775742    -1.775742
Age                -0.039622   -0.039622
```

7.6.1.4 Graphical assessment of proportional odds

There are several graphical methods for visual assessment of the proportional odds assumption. These are all *marginal* methods, in that they treat the predictors one at a time. However, that

provides one means to determine if a partial proportional odds model might be more appropriate. Harrell's 2001, Ch. 13-14 *Regression Modeling Strategies* and the corresponding `rms` package provide an authoritative treatment and methods in R.

One simple idea is to plot the conditional mean or expected value $E(X | Y)$ of a given predictor, X , at each level of the ordered response Y . If the response behaves ordinally in relation to X , these means should be strictly increasing or decreasing with Y . For comparison, one can also plot the estimated conditional means $\hat{E}(X | Y = j)$ under the fitted PO model X as the only predictor. If the PO assumption holds for this X , the model-mean curve should be close to the data mean curve.

```
> library(rms)

Error in library(rms): there is no package called 'rms'

> arth.po2 <- lrm(Improved ~ Sex + Treatment + Age, data=Arthritis)

Error in eval(expr, envir, enclos): could not find function "lrm"

> arth.po2

Error in eval(expr, envir, enclos): object 'arth.po2' not found
```

The plot of conditional X means is produced using the `plot.xmean.ordinally()` as shown below. It produces one marginal panel for each predictor in the model. For categorical predictors, it plots only the overall most frequent category. The resulting plot is shown in Figure ??.

```
> op <- par(mfrow=c(1,3))
> plot.xmean.ordinally(Improved ~ Sex + Treatment + Age, data=Arthritis,
+                      lwd=2, pch=16, subn=FALSE)

Error in eval(expr, envir, enclos): could not find function "plot.xmean.ordinally"

> par(op)
```

In Figure ??, there is some evidence that the effect of `Sex` is non-monotonic and the means differ from their model-implied values under the PO assumption. The effect of `Treatment` looks good by this method, and the effect of `Age` hints that the upper two categories may not be well-distinguished as an ordinal response.

Of course, this example has only a modest total sample size, and this method only examines the marginal effects of the predictors. Nevertheless, it is a useful supplement to the statistical methods described earlier.

7.6.2 Visualizing results for the proportional odds model

Results from the PO model (and other models for polytomous responses) can be graphed using the same ideas and methods shown earlier for a binary or binomial response. In particular, full-model plots (described earlier in Section 7.3.2) and effect plots (Section 7.3.3) are still very helpful.

But now there is the additional complication that the response variable has $m > 2$ levels and so needs to be represented by $m - 1$ curves or panels in addition to those related to the predictor variables.

7.6.2.1 Full-model plots

For full-model plots, we continue the idea of appending the fitted response probabilities (or logits) to the data frame and plotting these in relation to the predictors. The `predict()` method returns the highest probability category label by default (with `type="class"`), so to get the fitted probabilities you have to ask for `type="probs"`, as shown below.

{sec:vis-propodds}

{sec:po-fullplots}

```
> arth.fitp <- cbind(Arthritis,
+                      predict(arth.polr, type="probs"))
> head(arth.fitp)

  ID Treatment Sex Age Improved Better    None    Some   Marked
1 57   Treated Male 27     Some      1 0.73262 0.13806 0.12932
2 46   Treated Male 29     None      0 0.71740 0.14443 0.13816
3 77   Treated Male 30     None      0 0.70960 0.14763 0.14277
4 17   Treated Male 32     Marked    1 0.69363 0.15400 0.15237
5 36   Treated Male 46     Marked    1 0.57025 0.19504 0.23471
6 23   Treated Male 58     Marked    1 0.45634 0.21713 0.32653
```

For plotting, it is most convenient to reshape these from wide to long format using `melt()` in the `reshape2` package. The response category is named `Level`.

```
> library(reshape2)
> plotdat <- melt(arth.fitp,
+                   id.vars = c("Sex", "Treatment", "Age", "Improved"),
+                   measure.vars=c("None", "Some", "Marked"),
+                   variable.name = "Level",
+                   value.name = "Probability")
> ## view first few rows
> head(plotdat)

  Sex Treatment Age Improved Level Probability
1 Male   Treated 27     Some   None 0.73262
2 Male   Treated 29     None   None 0.71740
3 Male   Treated 30     None   None 0.70960
4 Male   Treated 32     Marked  None 0.69363
5 Male   Treated 46     Marked  None 0.57025
6 Male   Treated 58     Marked  None 0.45634
```

We can now plot `Probability` against `Age`, using `Level` to assign different colors to the lines for the response categories. `facet_grid()` is used to split the plot into separate panels by `Sex` and `Treatment`. In this example, the `directlabels` package is also used replace the default legend created by `ggplot()` with category labels on the curves themselves, which is easier to read.

```
> library(ggplot2)
> library(directlabels)
> gg <- ggplot(plotdat, aes(x = Age, y = Probability, colour = Level)) +
+   geom_line(size=2.5) + theme_bw() + xlim(10, 80) +
+   geom_point(color="black", size=1.5) +
+   facet_grid(Sex ~ Treatment,
+              labeller = function(x, y) sprintf("%s = %s", x, y))
+ )
> direct.label(gg)
```

Although we now have three response curves in each panel, this plot is relatively easy to understand: (a) In each panel, the probability of no improvement decreases with age, while that for marked improvement increases. (b) It is easy to compare the placebo and treated groups in each row, showing that no improvement decreases, while marked improvement increases with the active treatment. (On the other hand, this layout makes it harder to compare panels vertically for males and females in each condition.) (c) The points show where the observations are located in each panel; so, we can see that the data is quite thin for males given the placebo.²⁵

²⁵One way to improve (pun intended) this graph would be to show the points on the lines only for the actual level of `Improved` for each observation.

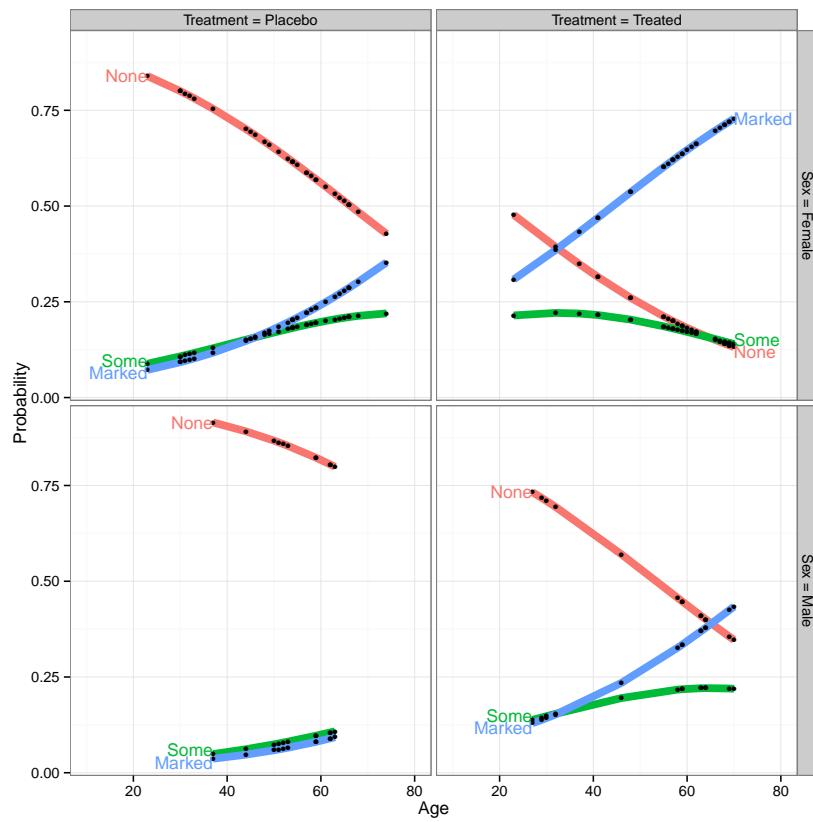


Figure 7.33: Predicted probabilities for the proportional odds model fit to the Arthritis data fig:arth-polr1

7.6.2.2 Effect plots

{sec:po-effplots} For PO models fit using `polr()`, the `effects` package provides two different styles for plotting a given effect. By default, curves are plotted in separate panels for the different response levels of a given effect, together with confidence bands for predicted probabilities. This form provides confidence bands and rug plots for the observations, but the default vertical arrangement of the panels makes it harder to compare the trends for the different response levels. The alternative *stacked* format shows the changes in response level more directly, but doesn't provide confidence bands.

Figure ?? shows these two styles for the main effect of Age in the proportional odds model, `arth.polr` fit earlier.

```
> plot(Effect("Age", arth.polr))

Error in plot(Effect("Age", arth.polr)): error in evaluating the argument
'x' in selecting a method for function 'plot': Error: could not find function
"Effect"

> plot(Effect("Age", arth.polr), style='stacked',
+       key.args=list(x=.55, y=.9))

Error in plot(Effect("Age", arth.polr), style = "stacked", key.args = list(x
= 0.55, : error in evaluating the argument 'x' in selecting a method for function
'plot': Error: could not find function "Effect"
```

Even though this model includes only main effects, you can still plot the higher-order effects for more focal predictors in a coherent display. Figure ?? shows the predicted probabilities for all three predictors together. Again, visual comparison is easier horizontally for placebo versus treated groups, but you can also see that the prevalence of marked improvement is greater for females than for males.

```
> plot(Effect(c("Treatment", "Sex", "Age"), arth.polr),
+       style="stacked", key.args=list(x=.8, y=.9))

Error in plot(Effect(c("Treatment", "Sex", "Age"), arth.polr), style = "stacked",
: error in evaluating the argument 'x' in selecting a method for function
'plot': Error: could not find function "Effect"
```

Finally, the latent variable interpretation of the PO model provides for simpler plots on the logit scale. Figure ?? shows this plot for the effects of Treatment and Age (collapsed over Sex) produced with the argument `latent=TRUE` to `Effect()`. In this plot, there is a single line in each panel for the effect (slope) of Age on the log odds. The dashed horizontal lines give the thresholds between the adjacent response categories corresponding to the intercepts.

```
> plot(Effect(c("Treatment", "Age"), arth.polr, latent=TRUE), lwd=3)

Error in plot(Effect(c("Treatment", "Age"), arth.polr, latent = TRUE), : error
in evaluating the argument 'x' in selecting a method for function 'plot': Error:
could not find function "Effect"
```

7.6.3 Nested dichotomies

{sec:nested} The method of *nested dichotomies* provides another simple way to analyse a polytomous response in the framework of logistic regression (or other generalized linear models). This method does not require an ordinal response or special software. Instead, it uses the familiar binary logistic model

and fits $m - 1$ separate models for each of a hierarchically nested set of comparisons among the response categories.

Taken together, this set of models for the dichotomies comprises a complete model for the polytomous response. As well, these models are statistically independent, so test statistics such as G^2 or Wald tests can be added to give overall tests for the full polytomy.

For example, the response categories $Y = \{1,2,3,4\}$ could be divided first as $\{1,2\}$ vs. $\{3,4\}$, as shown in the left side of Figure 7.34. Then these two dichotomies could be divided as $\{1\}$ vs. $\{2\}$, and $\{3\}$ vs. $\{4\}$. Alternatively, these response categories could be divided as shown in the right side of Figure 7.34: first, $\{1\}$ vs. $\{2,3,4\}$, then $\{2\}$ vs $\{3,4\}$, and finally $\{3\}$ vs. $\{4\}$.

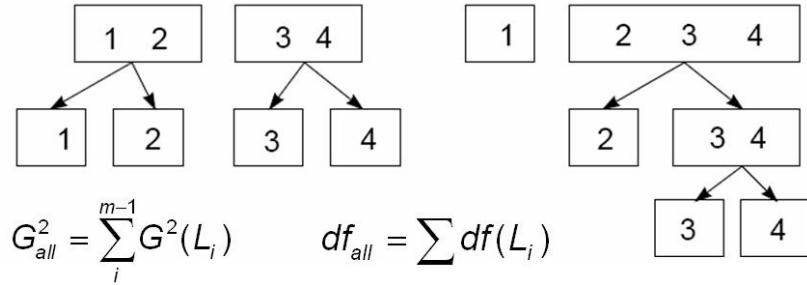


Figure 7.34: Nested dichotomies. The boxes show two different ways a four-category response can be represented as three nested dichotomies. Adapted from Fox (2008).

{fig:nested2}

Such models make the most sense when there are substantive reasons for considering the response categories in terms of such dichotomies. Two examples are shown in Figure 7.35.

- For the *Arthritis* data, it is sensible to consider one dichotomy (“better”), with logit L_1 , between the categories of “None” compared to “Some” or “Marked”. A second dichotomy, with logit L_2 , would then distinguish between the some and marked response categories.
- For a second case where patients are classified into $m = 4$ psychiatric diagnostic categories, the first dichotomy, with logit L_1 distinguishes those considered normal from all others given a clinical diagnosis. Two other dichotomies are defined to further divide the non-normal categories.

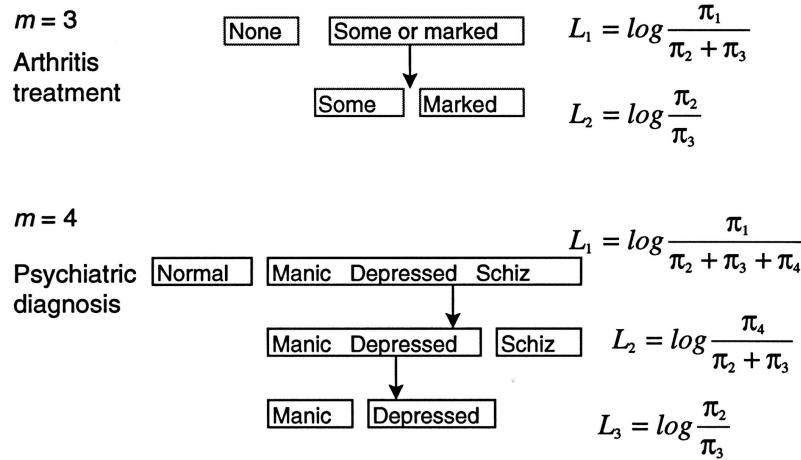
Then, consider the separate logit models for these $m - 1$ dichotomies, with different intercepts α_j and slopes β_j for each dichotomy,

$$\begin{aligned} L_1 &= \alpha_1 + \mathbf{x}^\top \boldsymbol{\beta}_1 \\ L_2 &= \alpha_2 + \mathbf{x}^\top \boldsymbol{\beta}_2 \\ &\vdots = \vdots \\ L_{m-1} &= \alpha_{m-1} + \mathbf{x}^\top \boldsymbol{\beta}_{m-1} \end{aligned}$$

{ex:wlfpart1}

EXAMPLE 7.17: Women’s labor force participation

The data set *Womenlf* in the *car* package gives the result of a 1977 Canadian survey. It contains data for 263 married women of age 21–30 who indicated their working status (outside the home) as not working, working part time or working full time, together with their husband’s income and a binary indicator of whether they had one or more young children in their household. (Another variable, region of Canada, had no effects in these analyses, and is not examined here.) This example follows Fox and Weisberg (2011, §5.8).

**Figure 7.35:** Examples of nested dichotomies and the corresponding logits

{fig:nested1}

```
> library(car) # for data and Anova()
> data("Womenlf", package="car")
> some(Womenlf)

      partic hincome children region
43  parttime      28    absent Ontario
50  fulltime       15    absent Prairie
82  parttime      15   present Ontario
100 not.work      15   present Ontario
105 parttime      12    absent     BC
184 fulltime      18    absent Ontario
196 fulltime       5   present Prairie
206 not.work      19   present Quebec
214 not.work      15   present Quebec
239 not.work      13   present Quebec
```

In this example, it makes sense to consider a first dichotomy (working) between women who are not working, vs. those who are (full time or part time). A second dichotomy (fulltime) contrasts full time work vs. part time work, among those women who are working at least part time. These two binary variables are created in the data frame using the `recode()` function from the `car` package.

```
> # create dichotomies
> Womenlf <- within(Womenlf, {
+   working <- recode(partic, " 'not.work' = 'no'; else = 'yes' ")
+   fulltime <- recode(partic,
+     "'fulltime' = 'yes'; 'parttime' = 'no'; 'not.work' = NA"))
> some(Womenlf)

      partic hincome children region fulltime working
41  not.work      9   present      BC    <NA>      no
43  parttime     28    absent  Ontario      no     yes
69  not.work     19   present  Ontario    <NA>      no
78  not.work      7   present  Atlantic    <NA>      no
91  not.work     35    absent  Ontario    <NA>      no
194 not.work    17   present  Prairie    <NA>      no
212 not.work    13   present  Quebec    <NA>      no
226 not.work    15   present  Quebec    <NA>      no
244 fulltime      6   present  Quebec      yes     yes
255 fulltime    11    absent  Quebec      yes     yes
```

The tables below show how the response `partic` relates to the recoded binary variables, `working` and `fulltime`. Note that the `fulltime` variable is recoded to NA for women who are not working.

```
> with(Womenlf, table(partic, working))

      working
partic      no yes
  fulltime    0   66
  not.work 155   0
  parttime    0   42

> with(Womenlf, table(partic, fulltime, useNA="ifany"))

      fulltime
partic      no yes <NA>
  fulltime    0   66     0
  not.work    0     0 155
  parttime    42     0     0
```

We proceed to fit two separate binary logistic regression models for the derived dichotomous variables. For the `working` dichotomy, we get the following results:

```
> mod.working <- glm(working ~ hincome + children, family=binomial,
+                      data=Womenlf)
> summary(mod.working)

Call:
glm(formula = working ~ hincome + children, family = binomial,
     data = Womenlf)

Deviance Residuals:
    Min      1Q  Median      3Q      Max 
-1.677  -0.865  -0.777   0.929   1.997 

Coefficients:
            Estimate Std. Error z value Pr(>|z|)    
(Intercept)  1.3358    0.3838   3.48   0.0005 ***  
hincome     -0.0423    0.0198  -2.14   0.0324 *    
childrenpresent -1.5756    0.2923  -5.39   7e-08 ***  
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for binomial family taken to be 1)

Null deviance: 356.15 on 262 degrees of freedom
Residual deviance: 319.73 on 260 degrees of freedom
AIC: 325.7

Number of Fisher Scoring iterations: 4
```

And, similarly for the `fulltime` dichotomy:

```
> mod.fulltime <- glm(fulltime ~ hincome + children, family=binomial,
+                      data=Womenlf)
> summary(mod.fulltime)

Call:
glm(formula = fulltime ~ hincome + children, family = binomial,
     data = Womenlf)
```

```

Deviance Residuals:
    Min      1Q  Median      3Q     Max
-2.405 -0.868   0.395   0.621   1.764

Coefficients:
            Estimate Std. Error z value Pr(>|z|)
(Intercept) 3.4778    0.7671   4.53  5.8e-06 ***
hincome     -0.1073    0.0392  -2.74  0.0061 **
childrenpresent -2.6515    0.5411  -4.90  9.6e-07 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for binomial family taken to be 1)

Null deviance: 144.34 on 107 degrees of freedom
Residual deviance: 104.49 on 105 degrees of freedom
(155 observations deleted due to missingness)
AIC: 110.5

Number of Fisher Scoring iterations: 5

```

Although these were fit separately, we can view this as a combined model for the three-level response, with the following coefficients:

```

> cbind(working=coef(mod.working), fulltime=coef(mod.fulltime))

           working fulltime
(Intercept) 1.335830 3.47777
hincome     -0.042308 -0.10727
childrenpresent -1.575648 -2.65146

```

Writing these out as equations for the logits, we have:

$$L_1 = \log \frac{\Pr(\text{working})}{\Pr(\text{notworking})} = 1.336 - 0.042 \text{hincome} - 1.576 \text{children} \quad (7.19)$$

$$L_2 = \log \frac{\Pr(\text{fulltime})}{\Pr(\text{parttime})} = 3.478 - 0.1072 \text{hincome} - 2.652 \text{children} \quad (7.20)$$

For both dichotomies, increasing income of the husband and the presence of young children decrease the log odds of a greater level of work. However, for those women who are working the effects of husband's income and and children are greater on the choice between full time and part time work than they are for all women on the choice between working and not working.

As we mentioned above, the use of nested dichotomies implies that the models fit to the separate dichotomies are statistically independent. Thus, we can additively combine χ^2 statistics and degrees of freedom to give overall tests for the polytomous response.

For example, here we define a function, `LRtest()` to calculate the likelihood ratio test of the hypothesis $H_0 : \beta = 0$ for all predictors simultaneously. We then use this to display these tests for each sub-model, as well as the combined model based on the sums of the test statistic and degrees of freedom.

```

> LRtest <- function(model)
+   c(LRchisq=(model$null.deviance - model$deviance),
+       df=(model$df.null - model$df.residual))
> tab <- rbind(working=LRtest(mod.working),
+               fulltime=LRtest(mod.fulltime))
> tab <- rbind(tab, All = colSums(tab))
> tab <- cbind(tab, pvalue = 1- pchisq(tab[,1], tab[,2]))
> tab

```

	LRchisq	df	pvalue
working	36.418	2	1.2355e-08
fulltime	39.847	2	2.2252e-09
All	76.265	4	1.1102e-15

Similarly, you can carry out tests of individual predictors, $H_0 : \beta_i = \mathbf{0}$ for the polytomy by adding the separate χ^2 s from `Anova()`.

```
> Anova(mod.working)

Analysis of Deviance Table (Type II tests)

Response: working
      LR Chisq Df Pr(>Chisq)
hincome   4.82637  1  0.028028 *
children 31.32288  1 2.1849e-08 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

> Anova(mod.fulltime)

Analysis of Deviance Table (Type II tests)

Response: fulltime
      LR Chisq Df Pr(>Chisq)
hincome   8.9813  1  0.0027275 **
children 32.1363  1 1.4373e-08 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

For example, the test for husband's income gives $\chi^2 = 4.826 + 8.981 = 13.807$ with 2 df.

As before, you can plot the fitted values from such models, either on the logit scale (for the separate logit equations) or in terms of probabilities for the various responses. The general idea is the same: obtain the fitted values from `predict()` using data frame containing the values of the predictors. However, now we have to combine these for each of the sub-models.

We calculate these values below, on both the logit scale and the response scale of probabilities. The `newdata` argument to `predict()` is constructed as the combinations of values for `hincome` and `children`.²⁶

```
> predictors <- expand.grid(hincome=1:50,
+                               children=c('absent', 'present'))
> fit <- data.frame(predictors,
+                      p.working = predict(mod.working, predictors, type='response'),
+                      p.fulltime = predict(mod.fulltime, predictors, type='response'),
+                      l.working = predict(mod.working, predictors, type='link'),
+                      l.fulltime = predict(mod.fulltime, predictors, type='link'))
+ )
> print(some(fit, 5), digits=3)

  hincome children p.working p.fulltime l.working l.fulltime
1        1    absent     0.7847     0.9668     1.294    3.3705
27       27   absent     0.5482     0.6414     0.194    0.5815
32       32   absent     0.4955     0.5113    -0.018    0.0452
77       27  present     0.2007     0.1121    -1.382   -2.0699
98       48  present     0.0936     0.0131    -2.271   -4.3225
```

One wrinkle here is that the probabilities for working full time and part time are conditional on working. We calculate the unconditional probabilities as shown below and choose to display the probability of *not* working as the complement of working.

²⁶Alternatively, using the predictor values in the `Womenlf` data would give the fitted values for the cases in the data, and allow a more data-centric plot as shown in Figure 7.33.

```
> fit <- within(fit, {
+   full <- p.working * p.fulltime
+   part <- p.working * (1 - p.fulltime)
+   not <- 1 - p.working
+ })
```

Plotting these fitted values using `ggplot2` would require reshaping the `fit` data frame from wide to long format. Instead, we use R base graphics to produce plots of the probabilities and log odds. This method doesn't automatically give plots in separate panels, so a `for`-loop is used to generate panels for the levels of `children`. We set up an empty plot frame (`type="n"`) for each panel and then use `lines()` to plot the fitted probabilities. Using `par(mfrow=c(1, 2))` places these plots in two side-by-side panels in a single display. The lines below give the plot shown in Figure 7.36.

```
> op <- par(mfrow=c(1, 2), mar=c(5, 4, 4, 1)+.1)
> Hinc <- 1:max(fit$hincome)
> for ( kids in c("absent", "present") ) {
+   dat <- subset(fit, children==kids)
+   plot( range(Hinc), c(0,1), type="n", cex.lab=1.25,
+         xlab="Husband's Income", ylab='Fitted Probability',
+         main = paste("Children", kids))
+   lines(Hinc, dat$not, lwd=3, col="black", lty=1)
+   lines(Hinc, dat$part, lwd=3, col="blue", lty=2)
+   lines(Hinc, dat$full, lwd=3, col="red", lty=3)
+   if (kids=="absent") {
+     legend("topright", lty=1:3, lwd=3, col=c("black", "blue", "red"),
+           legend=c('not working', 'part-time', 'full-time'))
+   }
+ }
> par(op)
```

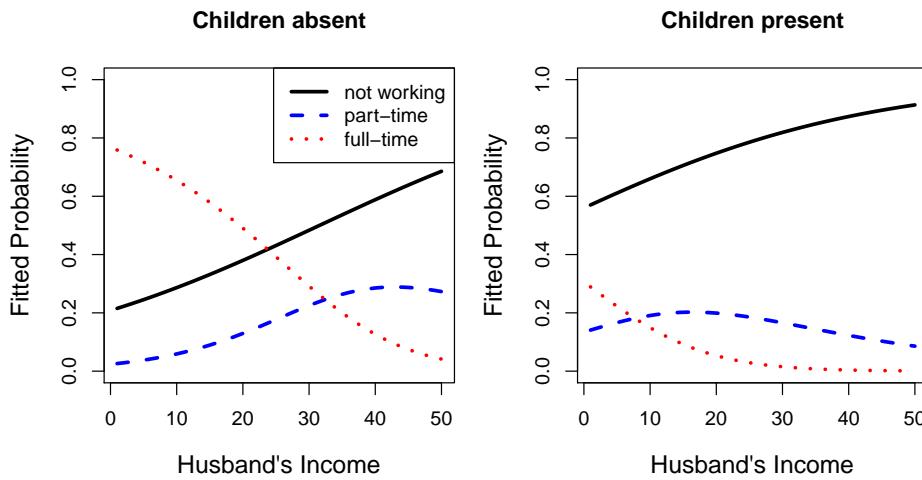


Figure 7.36: Fitted probabilities from the models for nested dichotomies fit to the data on women's labor force participation.^{fig-wif-fitted-prob}

We can see how that the decision not to work outside the home increases strongly with husband's income, and is higher when there are children present. As well, among working women, the decision to work full time as opposed to part time decreases strongly with husband's income, and is less likely with young children.

Similarly, we plot the fitted logits for the two dichotomies in `l.working` and `l.fulltime` as shown below, giving Figure 7.37.

```
> op <- par(mfrow=c(1,2), mar=c(5,4,1,1)+.1)
> for ( kids in c("absent", "present") ) {
+   dat <- subset(fit, children==kids)
+   plot( range(Hinc), c(-4,5), type="n", cex.lab=1.25,
+         xlab="Husband's Income", ylab='Fitted log odds')
+   lines(Hinc, dat$l.working, lwd=3, col="black", lty=1)
+   lines(Hinc, dat$l.fulltime, lwd=3, col="blue", lty=2)
+   text(25, 4.5, paste("Children", kids), cex=1.4)
+   if (kids=="absent") {
+     legend("bottomleft", lty=1:3, lwd=3, col=c("black", "blue"),
+           legend=c('working', 'full-time'))
+   }
+ }
> par(op)
```

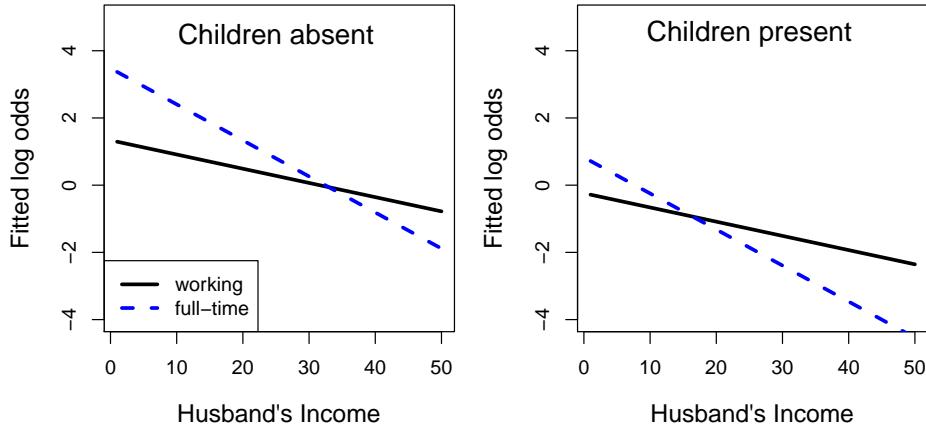


Figure 7.37: Fitted log odds from the models for nested dichotomies fit to the data on women's labor force participation.
fig:wif-fitted-logit

This is essentially a graph of the fitted equations for L_1 and L_2 shown in Eqn. (7.19). It shows how the choice of full time work as opposed to part time depends more strongly on husband's income among women who are working than does the choice of working at all among all women. It also illustrates why the proportional odds assumption would not be reasonable for this data: that would require equal slopes for the two lines within each panel.



7.6.4 Generalized logit model

The generalized logit (or multinomial logit) approach models the probabilities of the m response categories directly as a set of $m - 1$ logits. These compare each of the first $m - 1$ categories to the last category, which serves as the baseline.²⁷ The logits for any other pair of categories can be retrieved from the $m - 1$ fitted ones.

{sec:genlogit}

²⁷When the response is a factor, any category can be selected as the baseline level using `relevel()`.

When there are p predictors, x_1, x_2, \dots, x_p , which may be quantitative or categorical, the generalized logit model expresses the logits as

$$\begin{aligned} L_{jm} \equiv \log \frac{\pi_{ij}}{\pi_{im}} &= \beta_{0j} + \beta_{1j} x_{i1} + \beta_{2j} x_{i2} + \cdots + \beta_{kj} x_{ip} \quad j = 1, \dots, m-1 \\ &= \mathbf{x}_i^\top \boldsymbol{\beta}_j \end{aligned} \quad (7.21) \quad \text{eq:glogit1}$$

Thus, there is one set of fitted coefficients, $\boldsymbol{\beta}_j$ for each response category except the last. Each coefficient, β_{hj} , gives the effect, for a unit change in the predictor x_h , on the log odds that an observation had a response in category $Y = j$, as opposed to category $Y = m$.

The probabilities themselves can be expressed as

$$\begin{aligned} \pi_{ij} &= \frac{\exp(\mathbf{x}_i^\top \boldsymbol{\beta}_j)}{1 + \sum_{\ell=1}^{m-1} \exp(\mathbf{x}_i^\top \boldsymbol{\beta}_\ell)} \quad j = 1, 2, \dots, m-1 \\ \pi_{im} &= 1 - \sum_{i=1}^{m-1} \pi_{ij} \quad \text{for } Y = m \end{aligned}$$

Parameters in the $m-1$ equations Eqn. (7.21) can be used to determine the probabilities or the predicted log odds for any pair of response categories by subtraction. For instance, for an arbitrary pair of categories, a and b , and two predictors, x_1 and x_2 ,

$$\begin{aligned} L_{ab} &= \log \frac{\pi_{ia}/\pi_{im}}{\pi_{ib}/\pi_{im}} \\ &= \log \frac{\pi_{ia}}{\pi_{im}} - \log \frac{\pi_{ib}}{\pi_{im}} \\ &= (\beta_{0a} - \beta_{0b}) + (\beta_{1a} - \beta_{1b})x_{i1} + (\beta_{2a} - \beta_{2b})x_{i2} \end{aligned}$$

For example, the coefficient for x_{i1} in L_{ab} is just $(\beta_{1a} - \beta_{1b})$. Similarly, the predicted logit for any pair of categories can be calculated as

$$\hat{L}_{ab} = \hat{L}_{am} - \hat{L}_{bm} .$$

The generalized logit model can be fit most conveniently in R using the function `multinom()` in the `nnet` package and the `effects` package has a set of methods for "multinom" models. These models can also be fit using `VGAM` and the `mlogit` package.

EXAMPLE 7.18: Women's labor force participation

To illustrate this method, we fit the generalized logit model to the women's labor force participation data as explained below. The response, `partic` is a character factor, and, by default `multinom()` treats these in alphabetical order and uses the *first* level as the baseline category.

```
> levels(Womenlf$partic)
[1] "fulltime" "not.work" "parttime"
```

Although the multinomial model does not depend on the baseline category, it makes interpretation easier to choose "not.work" as the reference level, which we do with `relevel()`.²⁸

²⁸Alternatively, we could declare `partic` an *ordered* factor, using `ordered()`.

```
> # choose not working as baseline category
> Womenlf$partic <- relevel(Womenlf$partic, ref="not.work")
```

We fit the main effects model for husband's income and children as follows. As we did with `polr()` (Section 7.6.1), specifying `Hess=TRUE` saves the Hessian and facilitates calculation of standard errors and hypothesis tests.

```
> library(nnet)
> wlf.multinom <- multinom(partic ~ hincome + children,
+                               data=Womenlf, Hess=TRUE)

# weights: 12 (6 variable)
initial value 288.935032
iter 10 value 211.454772
final value 211.440963
converged
```

The `summary()` method for "multinom" objects doesn't calculate test statistics for the estimated coefficients by default. The option `Wald=TRUE` produces Wald z -test statistics, calculated as $z = \beta/SE(\beta)$.

```
> summary(wlf.multinom, Wald=TRUE)

Call:
multinom(formula = partic ~ hincome + children, data = Womenlf,
Hess = TRUE)

Coefficients:
            (Intercept) hincome childrenpresent
fulltime      1.9828 -0.0972321     -2.558605
parttime     -1.4323  0.0068938      0.021456

Std. Errors:
            (Intercept) hincome childrenpresent
fulltime      0.48418  0.028096      0.36220
parttime      0.59246  0.023455      0.46904

Value/SE (Wald statistics):
            (Intercept) hincome childrenpresent
fulltime      4.0953 -3.46071     -7.064070
parttime     -2.4176  0.29392      0.045744

Residual Deviance: 422.88
AIC: 434.88
```

Notice that the coefficients, their standard errors and the Wald test z values are printed in separate tables. The first line in each table pertains to the logit comparing full time work with the not working reference level; the second line compares part time work against not working.

For those who like p -values for significance tests, you can calculate these from the results returned by the `summary()` method in the `Wald.ratios` component, using the standard normal asymptotic approximation:

```
> stats <- summary(wlf.multinom, Wald=TRUE)
> z <- stats$Wald.ratios
> p <- 2 * (1 - pnorm(abs(z)))
> zapsmall(p)

            (Intercept) hincome childrenpresent
fulltime      0.00004  0.00054      0.00000
parttime      0.01562  0.76882      0.96351
```

The interpretation of these tests is that both husband's income and presence of children have highly significant effects on the comparison of working full time as opposed to not working, while neither of these predictors are significant for the comparison of working part time vs. not working.

So far, we have assumed that the effects of husband's income and presence of young children are additive on the log odds scale. We can test this assumption by allowing an interaction of those effects and testing it for significance.

```
> wlf.multinom2 <- multinom(partic ~ hincome * children,
+                               data=Womenlf, Hess=TRUE)

# weights: 15 (8 variable)
initial value 288.935032
iter 10 value 210.797079
final value 210.714841
converged

> Anova(wlf.multinom2)

Analysis of Deviance Table (Type II tests)

Response: partic
          LR Chisq Df Pr(>Chisq)
hincome      15.2   2    0.00051 ***
children     63.6   2    1.6e-14 ***
hincome:children  1.5   2    0.48378
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The test for the interaction term, `hincome:children` is not significant, so we can abandon this model.

Full model plots of the fitted values can be plotted as shown earlier in Example 7.17: obtain the fitted values over a grid of the predictors and plot these.

```
> predictors <- expand.grid(hincome=1:50,
+                             children=c('absent', 'present'))
> fit <- data.frame(predictors,
+                      predict(wlf.multinom, predictors, type='probs'))
```

Plotting these fitted values gives the plot shown in Figure 7.38.

```
> op <- par(mfrow=c(1, 2), mar=c(5, 4, 4, 1)+.1)
> Hinc <- 1:max(fit$hincome)
> for ( kids in c("absent", "present") ) {
+   dat <- subset(fit, children==kids)
+   plot( range(Hinc), c(0,1), type="n", cex.lab=1.25,
+         xlab="Husband's Income", ylab='Fitted Probability',
+         main = paste("Children", kids))
+   lines(Hinc, dat$not.work, lwd=3, col="black", lty=1)
+   lines(Hinc, dat$parttime, lwd=3, col="blue", lty=2)
+   lines(Hinc, dat$fulltime, lwd=3, col="red", lty=3)
+   if (kids=="absent") {
+     legend("topright", lty=1:3, lwd=3, col=c("black", "blue", "red"),
+           legend=c('not working', 'part-time', 'full-time'))
+   }
+ }
> par(op)
```

The results shown in this plot are roughly similar to those obtained from the nested dichotomy models, graphed in Figure 7.36. However, the predicted probabilities of not working under the

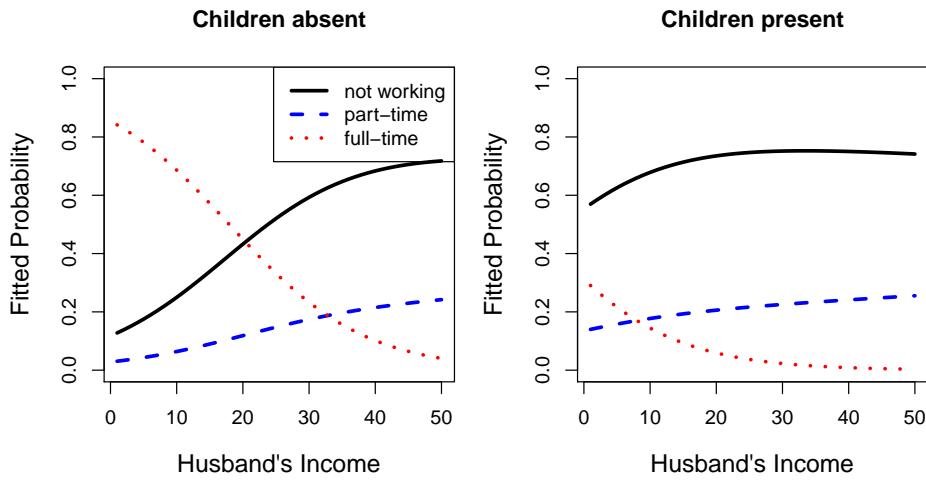


Figure 7.38: Fitted probabilities from the generalized logit model fit to the data on women's labor force participation.

generalized logit model rise more steeply with husband's income for women with no children and level off sooner for women with young children.

The `effects` package has special methods for "multinom" models. It treats the response levels in the order given by `levels()`, so before plotting we use `ordered()` to arrange levels in their natural order. The `update()` method provides a simple way to get a new fitted model; in the call, the model formula `. ~ .` means to fit the same model as before, i.e., `partic ~ hincome + children`.

```
> levels(Womenlf$partic)
[1] "not.work" "fulltime" "parttime"

> Womenlf$partic <- ordered(Womenlf$partic,
+                               levels=c('not.work', 'parttime', 'fulltime'))
> wlf.multinom <- update(wlf.multinom, . ~ .)

# weights: 12 (6 variable)
initial value 288.935032
iter 10 value 211.454772
final value 211.440963
converged
```

As illustrated earlier, you can use `plot(allEffects(model), ...)` to plot all the high-order terms in the model, either with separate curves for each response level (`style="lines"`) or as cumulative filled polygons (`style="stacked"`). Here, we simply plot the effects for the combinations of husband's income and children in stacked style, giving a plot (Figure ??) that is analogous to the full-model plot shown in Figure 7.38.

```
> plot(Effect(c("hincome", "children"), wlf.multinom),
+       style="stacked", key.args=list(x=.05, y=.9))

Error in plot(Effect(c("hincome", "children"), wlf.multinom), style = "stacked",
: error in evaluating the argument 'x' in selecting a method for function
'plot': Error: could not find function "Effect"
```



7.7 Chapter summary

{sec:ch07-summary}

- Model-based methods for categorical data provide confidence intervals for parameters and predicted values for observed and unobserved values of the explanatory variables. Graphical displays of predicted values help us to interpret the fitted relations by smoothing a discrete response.
- The logistic regression model (Section 7.2) describes the relationship between a categorical response variable, usually dichotomous, and a set of one or more quantitative or discrete explanatory variables (Section 7.3). It is conceptually convenient to specify this model as a linear model predicting the log odds (or logit) of the probability of a success from the explanatory variables.
- The relationship between a discrete response and a quantitative predictor may be explored graphically by plotting the binary observations against the predictor with some smoothed curve(s), either parametric or non-parametric, possibly stratified by other predictors.
- For both quantitative and discrete predictors, the results of a logistic regression are most easily interpreted from full-model plots of the fitted values against the predictors, either on the scale of predicted probabilities or log odds (Section 7.3.2). In these plots, confidence intervals provide a visual indication of the precision of the predicted results.
- When there are multiple predictors and/or higher-order interaction terms, effect plots (Section 7.3.3) provide an important method for constructing simplified displays, focusing on the higher-order terms in a given model.
- Influence diagnostics (Section 7.5) assess the impact of individual cases or groups on the fitted model, predicted values, and the coefficients of individual predictors. Among other displays, plots of residuals against leverage showing Cook's D are often most useful.
- Other diagnostic plots (Section 7.5.3) include component-plus-residual plots, that are useful for detecting non-linear relationships for a quantitative predictor, and added-variable plots, that show the partial relations of the response to a given predictor, controlling or adjusting for all other predictors.
- Polytomous responses may be handled in several ways as extensions of binary logistic regression (Section 7.6): (a) The *proportional odds model* (Section 7.6.1) is simple and convenient, but its validity depends on an assumption of equal slopes for adjacent-category logits. (b) *Nested dichotomies* (Section 7.6.3) among the response categories give a set of models which may be regarded as a single, combined model for the polytomous response. (c) *Generalized logit models* (Section 7.6.4) may be used to construct models comparing any pair of categories.

7.8 Further reading

7.9 Lab exercises

Exercise 7.1 Arbuthnot's data on the sex ratio of births in London was examined in Example 3.1. Use a binomial logistic regression model to assess whether the proportion of male births varied with the variables Year, Plague and Mortality in the *Arbuthnot* data set. Produce effect plots for the terms in this model. What do you conclude?

{sec:ch07-reading}

{sec:ch07-exercises}

{lab:7.1}

{lab:7.2}

Exercise 7.2 For the Donner Party data in *Donner*, examine Grayson's 1990 claim that survival in the Donner Party was also mediated by the size of the family unit. This takes some care, because the `family` variable in the *Donner* data is a simplified grouping based on the person's name and known alliances among families from the historical record. Use the following code to compute a `family.size` variable from each individual's last name:

```
> data("Donner", package="vcdExtra")
> Donner$survived <- factor(Donner$survived, labels=c("no", "yes"))
> # use last name for family
> lname <- strsplit(rownames(Donner), ", ")
> lname <- sapply(lname, function(x) x[[1]])
> Donner$family.size <- as.vector(table(lname))
```

- (a) Choose one of the models (`donner.mod4`, `donner.mod6`) from Example 7.9 that include the interaction of age and sex and non-linear terms in age. Fit a new model that adds a main effect of `family.size`. What do you conclude about Grayson's claim?
- (b) Produce an effect plot for this model.
- (c) Continue, by examining whether the effect of family size can be taken as linear, or whether a non-linear term should be added.

{lab:7.3}

Exercise 7.3 Use component+residual plots (Section 7.5.3) to examine the additive model for the *ICU* data given by

```
> icu.glm2 <- glm(died ~ age + cancer + admit + uncons,
+                   data=ICU, family=binomial)
```

- (a) What do you conclude about the linearity of the (partial) relationship between age and death in this model?
- (b) An alternative strategy is to allow some non-linear relation for age in the model using a quadratic (or cubic) term like `poly(age, 2)` (or `poly(age, 3)`) in the model formula. Do these models provide evidence for a non-linear effect of age on death in the ICU?

{lab:7.4}

Exercise 7.4 Explore the use of other marginal and conditional plots to display the relationships among the variables predicting death in the ICU in the model `icu.glm2`. For example, you might begin with a marginal `gpairs()` plot showing all bivariate marginal relations, something like this:

```
> library(gpairs)
> gpairs(ICU[,c("died", "age", "cancer", "admit", "uncons")],
+        diag.pars=list(fontsize=16, hist.color="lightgray"),
+        mosaic.pars=list(gp=shading_Friendly,
+                         gp_args=list(interpolate=1:4)))
```

{lab:7.5}

Exercise 7.5 For the women's labor force participation data (*Womenlf*) the response variable, `partic`, can be treated as ordinal by using

```
> Womenlf$partic <- ordered(Womenlf$partic,
+                               levels=c('not.work', 'parttime', 'fulltime'))
```

Use the methods in Section 7.6.1 to test whether the proportional odds model holds for these data.

{lab:7.6}

Exercise 7.6 The data set *housing* in the MASS package gives a $3 \times 3 \times 4 \times 2$ table in frequency form relating (a) satisfaction (Sat) of residents with their housing (High, Medium, Low), (b) perceived degree of influence (Infl) they have on the management of the property (High, Medium, Low), (c) Type of rental (Tower, Atrium, Apartment, Terrace), and (d) contact (Cont) residents have with other residents (Low, High). Consider satisfaction as the ordinal response variable.

- Fit the proportional odds model with additive (main) effects of housing type, influence in management and contact with neighbors to this data. (Hint: Using `polr()`, with the data in frequency form, you need to use the `weights` argument to supply the `Freq` variable.)
- Investigate whether any of the two-factor interactions among Infl, Type and Cont add substantially to goodness of fit of this model. (Hint: use `stepAIC()`, with the scope formula $\sim .^2$ and `direction="forward"`.)
- For your chosen model from the previous step, use the methods of Section 7.6.2 to plot the probabilities of the categories of satisfaction.
- Write a brief summary these analyses, interpreting *how* satisfaction with housing depends on the predictor variables.

{lab:7.7}

Exercise 7.7 The data *TV* on television viewing was analyzed using correspondence analysis in Example 6.4, ignoring the variable *Time* and extended in Exercise 6.7. Treating *Network* as a three-level response variable, fit a generalized logit model (Section 7.6.4) to explain the variation in viewing in relation to *Day* and *Time*. The *TV* data is a three-way table, so you will need to convert it to a frequency data frame first.

```
> data("TV", package="vcdExtra")
> TV.df <- as.data.frame.table(TV)
```

- Fit the main-effects model, *Network* ~ *Day* + *Time* with `multinom()`. Note that you will have to supply the `weights` argument because each row of *TV.df* represents the number of viewers in the `Freq` variable.
- Prepare an effects plot for the fitted probabilities in this model.
- Interpret these results in comparison to the correspondence analysis analysis in Example 6.4.

{lab:7.8}

Exercise 7.8 Refer to Exercise 5.8 for a description of the *Accident* data. The interest here is to model the probability that an accident resulted in death rather than injury from the predictors *age*, *mode* and *gender*. With `glm()`, and the data in the form of a frequency table, you can use the argument `weight=Freq` to take cell frequency into account.

- Fit the main effects model, `result=="Died" ~ age + mode + gender`. Use `car::Anova()` to assess the model terms.
- Fit the model that allows all two-way interactions. Use `anova()` to test whether this model is significantly better than the main effects model.
- Fit the model that also allows the three-way interaction of all factors. Does this offer any improvement over the two-way model?
- Interpret the results of the analysis using effect plots for the two-way model, separately for each of the model terms. Describe verbally the nature of the `age*gender` effect. Which mode of transportation leads to greatest risk of death?

```
> .locals$ch07 <- setdiff(ls(), .globals)
> #.locals$ch07
> remove(list=.locals$ch07[sapply(.locals$ch07, function(n) {!is.function(get(n))})])
> detach(package:gpairs)
```

```
Error in detach(package:gpairs): invalid 'name' argument
> detach(package:rms)

Error in detach(package:rms): invalid 'name' argument
> detach(package:VGAM)
> .pkgs$ch07 <- setdiff(.packages(), .pkgs$ch07)
> .pkgs$ch07

[1] "nnet"          "reshape2"       "stats4"        "proto"
[5] "car"           "lmtree"         "zoo"           "MASS"
[9] "gmodels"       "directlabels"  "quadprog"      "xtable"
[13] "vcdExtra"      "gnm"            "lattice"       "splines"
[17] "ggplot2"       "ca"             "vcd"           "grid"
```


Chapter 8

Loglinear and Logit Models for Contingency Tables

Loglinear models comprise another special case of generalized linear models designed for contingency tables of frequencies. They are most easily interpreted through visualizations, including mosaic displays and plots of associated logit models. Special cases arise for ordered categorial variables and square tables that allow more parsimonious models for associations.

{ch:loglin}

8.1 Introduction

{sec:loglin-intro}

Tables are like cobwebs, like the sieve of Danaides; beautifully reticulated, orderly to look upon, but which will hold no conclusion. Tables are abstractions, and the object a most concrete one, so difficult to read the essence of.

From *Chartism* by Thomas Carlyle (1840), Chapter II, Statistics

The chapter continues the modeling framework begun in Chapter 7, and takes up the case of loglinear models for contingency tables of frequencies, when all variables are discrete, another special case of generalized linear models. These models provide a comprehensive scheme to describe and understand the associations among two or more categorical variables. Whereas logistic regression models focus on the prediction of one response factor, loglinear models treat all variables symmetrically, and attempt to model all important associations among them.

In this sense, loglinear models are analogous to a correlation analysis of continuous variables, where the goal is to determine the patterns of dependence and independence among a set of variables. When one variable is a response and the others are explanatory, certain loglinear models are equivalent to logistic models for that response. Such models are also particularly useful when there are two or more response variables, a case that would require a multivariate version of the generalized linear model, for which the current theory and implementations are thin at best.

Chapter 5 and Chapter 6 introduced some basic aspects of loglinear models in connection with mosaic displays and correspondence analysis. In this chapter, the focus is on fitting and interpreting loglinear models. The usual analyses, with `loglm()` and `glm()` present the results in terms of tables of parameter estimates. Particularly for larger tables, it becomes difficult to understand the nature of these associations from tables of parameter estimates. Instead, we emphasize plots of

observed and predicted frequencies, probabilities or log odds (when there are one or more response variables), as well as mosaic and other displays for interpreting a given model. We also illustrate how mosaic displays and correspondence analysis plots may be used in a complementary way to the usual numerical summaries, to provide additional insights into the data.

Section 8.2 gives a brief overview of loglinear models in relation to the more familiar ANOVA and regression models for quantitative data. Methods and software for fitting these models are discussed in Section 8.3. When one variable is a response, logit models for that response provide a simpler, but equivalent means for interpreting and graphing results of loglinear models, as we describe in Section 8.4. Another class of simplified models (Section 8.6) occurs when one or more of the explanatory variables are ordinal, and discrete levels might be replaced by numerical values. Models for square tables (Section 8.7), with the same row and column categories comprise another special case giving simpler descriptions than the saturated model of general association. These important special cases are extended to three-way and higher-dimensional tables in Section 8.8. Finally, Section 8.9 describes some methods for dealing with situations where there are several response variables, and it is useful to understand both the marginal relations of the responses with the predictors as well as how their association varies with the predictors

8.2 Loglinear models for frequencies

{sec:loglin-counts}

Loglinear models have been developed from two formally distinct, but related perspectives. The first is a discrete analog of familiar ANOVA models for quantitative data, where the multiplicative relations among joint and marginal probabilities are transformed into an additive one by transforming the counts to logarithms. The second is an analog of regression models, where the log of the cell frequency is modeled as a linear function of discrete predictors, with a random component often taken as the Poisson distribution and called **Poisson regression**; this approach is treated in more detail as generalized linear models for count data in Chapter 9.

8.2.1 Loglinear models as ANOVA models for frequencies

For two discrete variables, A and B , suppose we have a multinomial sample of n_{ij} observations in each cell i, j of an $I \times J$ contingency table. To ease notation, we replace a subscript by $+$ to represent summation over that dimension, so that $n_{i+} = \sum_j n_{ij}$, $n_{+j} = \sum_i n_{ij}$, and $n_{++} = \sum_{ij} n_{ij}$.

Let π_{ij} be the joint probabilities in the table, and let $m_{ij} = n_{++}\pi_{ij}$ be the expected cell frequencies under any model. Conditional on the observed total count, n_{++} , each count has a Poisson distribution, with mean m_{ij} . Any loglinear model may be expressed as a linear model for the log m_{ij} . For example, the hypothesis of independence means that the expected frequencies, m_{ij} , obey

$$m_{ij} = \frac{m_{i+} m_{+j}}{m_{++}} .$$

This multiplicative model can be transformed to an additive (linear) model by taking logarithms of both sides:

$$\log(m_{ij}) = \log(m_{i+}) + \log(m_{+j}) - \log(m_{++}) ,$$

which is usually expressed in an equivalent form in terms of model parameters,

{eq:1main}

$$\log(m_{ij}) = \mu + \lambda_i^A + \lambda_j^B \quad (8.1)$$

where μ is a function of the total sample size, λ_i^A is the “main effect” for variable A, $\lambda_i^A = \log \pi_{i+} - \sum_k (\log \pi_{k+})/I$, and λ_j^B is the “main effect” for variable B, $\lambda_j^B = \log \pi_{+j} - \sum_k (\log \pi_{+k})/J$. Model Eqn. (8.1) is called the **loglinear independence model** for a two-way table.

In this model, there are $I + J$ parameters, but only $(I - 1) + (J - 1)$ are separately estimable. Hence, the typical ANOVA sum-to-zero restrictions are usually applied to the parameters:

$$\sum_i^I \lambda_i^A = \sum_j^J \lambda_j^B = 0 .$$

These “main effects” in loglinear models pertain to differences among the marginal probabilities of a variable (which are usually not of direct interest).

Other restrictions to make the parameters identifiable are also used. Setting the first values, λ_1^A and λ_1^B to zero (the default in `glm()`), defines $\lambda_i^A = \log \pi_{i+} - \log \pi_{1+}$, and $\lambda_j^B = \log \pi_{+j} - \log \pi_{+1}$, as deviations from the first, reference category, but these parameterizations are otherwise identical. For modeling functions in R (`lm()`, `glm()`, etc.) the reference category parameterization is obtained using `contr.treatment()`, while the sum-to-zero constraints are obtained with `contr.sum()`.

Model Eqn. (8.1) asserts that the row and column variables are independent. For a two-way table, a model that allows an arbitrary association between the variables is the **saturated model**, including an additional term, λ_{ij}^{AB} :

$$\log(m_{ij}) = \mu + \lambda_i^A + \lambda_j^B + \lambda_{ij}^{AB} , \quad (8.2) \quad \text{(eq:lsat)}$$

where again, restrictions must be imposed for estimation:

$$\sum_i^I \lambda_i^A = 0, \quad \sum_j^J \lambda_j^B = 0, \quad \sum_i^I \lambda_{ij}^{AB} = \sum_j^J \lambda_{ij}^{AB} = 0 . \quad (8.3) \quad \text{(eq:lrestrict)}$$

There are thus $I - 1$ linearly independent λ_i^A row parameters, $J - 1$ linearly independent λ_j^B column parameters, and $(I - 1)(J - 1)$ linearly independent λ_{ij}^{AB} association parameters. This model is called the **saturated model** because the number of parameters in $\mu, \lambda_i^A, \lambda_j^B$, and λ_{ij}^{AB} is equal to the number of frequencies in the two-way table,

$$\frac{1}{(\mu)} + \frac{I - 1}{(\lambda_i^A)} + \frac{J - 1}{(\lambda_j^B)} + \frac{(I - 1)(J - 1)}{(\lambda_{ij}^{AB})} = IJ$$

The association parameters λ_{ij}^{AB} express the departures from independence, so large absolute values pertain to cells that differ from the independence model.

Except for the difference in notation, model Eqn. (8.2) is formally the same as a two-factor ANOVA model with an interaction, typically expressed as $E(y_{ij}) = \mu + \alpha_i + \beta_j + (\alpha\beta)_{ij}$. Hence, associations between variables in loglinear models are analogous to interactions in ANOVA models. The use of superscripted symbols, $\lambda_i^A, \lambda_j^B, \lambda_{ij}^{AB}$ rather than separate Greek letters is a convention in loglinear models, and useful mainly for multiway tables.

Models such as Eqn. (8.1) and Eqn. (8.2) are examples of **hierarchical models**. This means that the model must contain all lower-order terms contained within any high-order term in the model. Thus, the saturated model, Eqn. (8.2) contains λ_{ij}^{AB} , and therefore *must* contain λ_i^A and λ_j^B . As a result, hierarchical models may be identified by the shorthand notation which lists only the high-order terms: model Eqn. (8.2) is denoted $[AB]$, while model Eqn. (8.1) is $[A][B]$.

8.2.2 Loglinear models for three-way tables

Loglinear models for three-way contingency tables were described briefly in Section 5.4.1. Each type of model allows associations among different sets of variables and each has a different independence interpretation, as illustrated in Table 5.2.

{sec:loglin-3way}

For a three-way table, the saturated model, denoted $[ABC]$ is

$$\log m_{ijk} = \mu + \lambda_i^A + \lambda_j^B + \lambda_k^C + \lambda_{ij}^{AB} + \lambda_{ik}^{AC} + \lambda_{jk}^{BC} + \lambda_{ijk}^{ABC}. \quad (8.4) \quad \{eq:lsat3\}$$

This model allows all variables to be associated; Eqn. (8.4) fits the data perfectly because the number of independent parameters equals the number of table cells. Two-way terms, such as λ_{ij}^{AB} pertain to the *conditional association* between pairs of factors, controlling for the remaining variable. The presence of the three-way term, λ_{ijk}^{ABC} , means that the partial association (conditional odds ratio) between any pair varies over the levels of the third variable.

Omitting the three-way term in Model Eqn. (8.4) gives the model $[AB][AC][BC]$,

$$\{eq:lno3way\} \quad \log m_{ijk} = \mu + \lambda_i^A + \lambda_j^B + \lambda_k^C + \lambda_{ij}^{AB} + \lambda_{ik}^{AC} + \lambda_{jk}^{BC}, \quad (8.5)$$

in which all pairs are conditionally dependent given the remaining one. For any pair, the conditional odds ratios are the *same* at all levels of the remaining variable, so this model is often called the **homogeneous association model**.

The interpretation of terms in this model may be illustrated using the Berkeley admissions data (Example 4.10 and Example 4.14), for which the factors are Admit, Gender, and Department, in a $2 \times 2 \times 6$ table. In the homogeneous association model,

$$\{eq:berk1\} \quad \log m_{ijk} = \mu + \lambda_i^A + \lambda_j^D + \lambda_k^G + \lambda_{ij}^{AD} + \lambda_{ik}^{AG} + \lambda_{jk}^{DG}, \quad (8.6)$$

the λ -parameters have the following interpretations:

- The main effects, λ_i^A , λ_j^D and λ_k^G pertain to differences in the one-way marginal probabilities. Thus λ_j^D relates to differences in the total number of applicants to these departments, while λ_k^G relates to the differences in the overall numbers of men and women applicants.
- λ_{ij}^{AD} describes the conditional association between admission and department, that is different admission rates across departments (controlling for gender).
- λ_{ik}^{AG} relates to the conditional association between admission and gender, controlling for department. This term, if significant, might be interpreted as indicating gender-bias in admissions.
- λ_{jk}^{DG} , the association between department and gender, indicates whether males and females apply differentially across departments.

As we discussed earlier (Section 5.4), loglinear models for three-way (and larger) tables often have an interpretation in terms of various types of independence relations illustrated in Table 5.2. The model Eqn. (8.5) has no such interpretation, however the smaller model $[AC][BC]$ can be interpreted as asserting that A and B are (conditionally) independent controlling for C ; this independence interpretation is symbolized as $A \perp B | C$. Similarly, the model $[AB][C]$ asserts that A and B are jointly independent of C : $(A, B) \perp C$, while the model $[A][B][C]$ is the model of mutual (complete) independence, $A \perp B \perp C$.

8.2.3 Loglinear models as GLMs for frequencies

{sec:loglin-glms} In the GLM approach, a loglinear model may be cast in the form of a regression model for $\log \mathbf{m}$, where the table cells are reshaped to a column vector. One advantage is that models for tables of any size and structure may be expressed in a compact form.

For a contingency table of variables A, B, C, \dots , with $N = I \times J \times K \times \dots$ cells, let \mathbf{n} denote a column vector of the observed counts arranged in standard order, and let \mathbf{m} denote a similar vector of the expected frequencies under some model. Then any loglinear model may be expressed in the form

$$\log \mathbf{m} = \mathbf{X}\boldsymbol{\beta},$$

where \mathbf{X} is a known design or **model matrix** and $\boldsymbol{\beta}$ is a column vector containing the unknown λ parameters.

For example, for a 2×2 table, the saturated model Eqn. (8.2) with the usual zero-sum constraints Eqn. (8.3) can be represented as

$$\log \begin{pmatrix} m_{11} \\ m_{12} \\ m_{21} \\ m_{22} \end{pmatrix} = \begin{bmatrix} 1 & 1 & 1 & 1 \\ 1 & 1 & -1 & -1 \\ 1 & -1 & 1 & -1 \\ 1 & -1 & -1 & 1 \end{bmatrix} \begin{pmatrix} \mu \\ \lambda_1^A \\ \lambda_1^B \\ \lambda_{11}^{AB} \end{pmatrix}$$

Note that only the linearly independent parameters are represented here. $\lambda_2^A = -\lambda_1^A$, because $\lambda_1^A + \lambda_2^A = 0$, and $\lambda_2^B = -\lambda_1^B$, because $\lambda_1^B + \lambda_2^B = 0$, and so forth.

An additional substantial advantage of the GLM formulation is that it makes it easier to express models with ordinal or quantitative variables. `glm()`, with a model formula of the form `Freq ~ .` involving factors A, B, \dots and quantitative variables x_1, x_2, \dots , constructs the model matrix \mathbf{X} from the terms given in the formula. A factor with K levels gives rise to $K - 1$ columns for its main effect and sets of $K - 1$ columns in each interaction effect. A quantitative predictor, say x_1 (with a linear effect) creates a single column with its values and interactions with other terms are calculated at the products of the columns for the main effects.

The parameterization for factors is controlled by the contrasts assigned to a given factor (if any), or by the general `contrasts` option, that gives the contrast functions used for unordered and ordered factors:

```
> options("contrasts")
$contrasts
  unordered          ordered
"contr.treatment"   "contr.poly"
```

This says that, by default, unordered factors use the baseline (first) reference-level parameterization, while ordered factors are given a parameterization based on orthogonal polynomials, allowing linear, quadratic, ... effects, assuming integer-spacing of the factor levels.

8.3 Fitting and testing loglinear models

For a given table, possible loglinear models range from the baseline model of mutual independence, $[A][B][C][\dots]$ to the saturated model, $[ABC\dots]$ that fits the observed frequencies perfectly, but offers no simpler description or interpretation than the data itself.

Fitting a loglinear model is usually a process of deciding which association terms are large enough (“significantly different from zero”) to warrant inclusion in a model to explain the observed frequencies. Terms which are excluded from the model go into the residual or error term, which reflects the overall badness-of-fit of the model. The usual goal of loglinear modeling is to find a small model (few association terms) which nonetheless achieves a reasonable fit (small residuals).

8.3.1 Model fitting functions

In R, the most basic function for fitting loglinear models is `loglin()` in the `stats` package. This uses the classical iterative proportional fitting (IPF) algorithm described in Haberman (1972) and Fienberg (1980, §3.4). It is designed to work with the frequency data in table form, and a model specified in terms of the (high-order) table margins to be fitted. For example, the model Eqn. (8.5) of homogenous association for a three-way table is specified as

{sec:loglin-fitting}

{sec:loglin-functions}

```
> loglin(mytable, margin=list(c(1, 2), c(1, 3), c(2, 3)))
```

The function `loglm()` in MASS provides a more convenient front-end to `loglin()` to allow loglinear models to be specified using a model formula. With table variables A, B and C, the same model can be fit using `loglm()` as

```
> loglm(~ (A + B + C)^2, data=mytable)
```

When the data is a frequency data frame with frequencies in `Freq`, for example, the result of `mydf <- as.data.frame(mytable)`, you can also use a two-sided formula:

```
> loglm(Freq ~ (A + B + C)^2, data=mydf)
```

As implied in Section 8.2.3, loglinear models can also be fit using `glm()`, using `family=poisson` which constructs the model for `log(Freq)`. The same model is fit with `glm()` as:

```
> glm(Freq ~ (A + B + C)^2, data=mydf, family=poisson)
```

While all of these fit equivalent models, the details of the printed output, model objects, and available methods differ, as indicated in some of the examples that follow.

It should be noted that both the `loglin()`/`loglm()` methods based on iterative proportional fitting, and the `glm()` approach using the Poisson model for log frequency give maximum likelihood estimates, \hat{m} , of the expected frequencies, as long as all observed frequencies n are *all* positive. Some special considerations when there cells with zero frequencies are described in Section 8.5.

8.3.2 Goodness-of-fit tests

For an n -way table, global goodness-of-fit tests for a loglinear model attempt to answer the question “How well does the model reproduce the observed frequencies?” That is, how close are the fitted frequencies estimated under the model to those of the saturated model or the data?

To avoid multiple subscripts for an n -way table, let $\mathbf{n} = n_1, n_2, \dots, n_N$ denote the observed frequencies in a table with N cells, and corresponding fitted frequencies $\hat{\mathbf{m}} = \hat{m}_1, \hat{m}_2, \dots, \hat{m}_N$ according to a particular loglinear model. The standard goodness-of-fit statistics are sums over the cells of measures of the difference between the \mathbf{n} and $\hat{\mathbf{m}}$.

The most commonly used are the familiar Pearson chi-square,

$$\text{(eq:pchi)} \quad X^2 = \sum_i^N \frac{(n_i - \hat{m}_i)^2}{\hat{m}_i} , \quad (8.7)$$

and the likelihood-ratio G^2 or **deviance** statistic,

$$\text{(eq:pgsq)} \quad G^2 = 2 \sum_i^N n_i \log \left(\frac{n_i}{\hat{m}_i} \right) . \quad (8.8)$$

Both of these statistics have asymptotic χ^2 distributions (as $\sum \mathbf{n} \rightarrow \infty$), reasonably well-approximated when all expected frequencies are large.¹ The (residual) degrees of freedom are the number of cells

¹Except in bizarre or borderline cases, these tests provide the same conclusions when expected frequencies are at least moderate (all $\hat{m} > 5$). However, G^2 approaches the theoretical chi-squared distribution more slowly than does χ^2 , and the approximation may be poor when the average cell frequency is less than 5.

(N) minus the number of estimated parameters. The likelihood-ratio test can also be expressed as twice the difference in log-likelihoods under saturated and fitted models,

$$G^2 = 2 \log \left[\frac{\mathcal{L}(\mathbf{n}; \mathbf{n})}{\mathcal{L}(\widehat{\mathbf{m}}; \mathbf{n})} \right] = 2[\log \mathcal{L}(\mathbf{n}; \mathbf{n}) - \log \mathcal{L}(\widehat{\mathbf{m}}; \mathbf{n})] ,$$

where $\mathcal{L}(\mathbf{n}; \mathbf{n})$ is the likelihood for the saturated model and $\mathcal{L}(\widehat{\mathbf{m}}; \mathbf{n})$ is the corresponding maximized likelihood for the fitted model.

In practice such global tests are less useful for comparing competing models. You may find that several different models have an acceptable fit or, sadly, that none do (usually because you are “blessed” with a large sample size). It is then helpful to compare competing models *directly*, and two strategies are particularly useful in these cases.

First, the likelihood-ratio G^2 statistic has the property in that one can compare two **nested models** by their difference in G^2 statistics, which has a χ^2 distribution on the difference in degrees of freedom. Two models, M_1 and M_2 , are nested when one, say, M_2 , is a special case of the other. That is, model M_2 (with ν_2 residual df) contains a subset of the parameters of M_1 (with ν_1 residual df), the remaining ones being effectively set to zero. Model M_2 is therefore more restrictive and cannot fit the data better than the more general model M_1 , i.e., $G^2(M_2) \geq G^2(M_1)$. The least restrictive of all models, with $G^2 = 0$ and $\nu = 0$ df is the saturated model for which $\widehat{\mathbf{m}} = \mathbf{n}$.

Assuming that the less restrictive model M_1 fits, the difference in G^2 ,

$$\Delta G^2 \equiv G^2(M_2 | M_1) = G^2(M_2) - G^2(M_1) \quad (8.9) \quad \text{(eq:gsqlnest1)}$$

$$= 2 \sum_i n_i \log(\widehat{m}_{i1}/\widehat{m}_{i2}) \quad (8.10) \quad \text{(eq:gsqlnest2)}$$

has a chi-squared distribution with $\text{df} = \nu_2 - \nu_1$. The last equality Eqn. (8.10) follows from substituting in Eqn. (8.8).

Rearranging terms in Eqn. (8.9), we see that we can partition the $G^2(M_2)$ into two terms,

$$G^2(M_2) = G^2(M_1) + G^2(M_2 | M_1) .$$

The first term measures the difference between the data and the more general model M_1 . If this model fits, the second term measures the additional lack of fit imposed by the more restrictive model. In addition to providing a more focused test, $G^2(M_2 | M_1)$ also follows the chi-squared distribution more closely when some $\{m_i\}$ are small (Agresti, 2013, §10.6.3).

Alternatively, a second strategy uses other measures that combine goodness-of-fit with model parsimony and may also be used to compare non-nested models. The statistics described below are all cast in the form of badness-of-fit relative to degrees of freedom, so that smaller values reflect “better” models.

The simplest idea (Goodman, 1971) is to use G^2/df (or χ^2/df), which has an asymptotic expected value of 1 for a good-fitting model. This type of measure is not routinely reported by R software, but is easy to calculate from output.

The **Akaike Information Criterion** (AIC) statistic (Akaike, 1973) is a very general criterion for model selection with maximum likelihood estimation, based on the idea of maximizing the information provided by a fitted model. AIC is defined generally as

$$\text{AIC} = -2 \log \mathcal{L} + 2k$$

where $\log \mathcal{L}$ is the maximized log likelihood and k is the number of parameters estimated in the model. Better models correspond to *smaller* AIC. For loglinear models, minimizing AIC is equivalent to minimizing

$$\text{AIC}^* = G^2 - 2\nu ,$$

where ν is the residual df, but the values of AIC and AIC* differ by an arbitrary constant. This form is easier to calculate by hand from the output of any modeling function if AIC is not reported, or an AIC() method is not available.

A third statistic of this type is the ***Bayesian Information Criterion*** (BIC) due to Schwartz (1978) and Raftery (1986),

$$\text{BIC} = G^2 - \log(n) \nu ,$$

where n is the total sample size. Both AIC and BIC penalize the fit statistic for increasing number of parameters. BIC also penalizes the fit directly with (log) sample size, and so expresses a preference for less complex models than AIC as the sample size increases.

8.3.3 Residuals for loglinear models

Test statistics such as G^2 can determine whether a model has significant lack of fit, and model comparison tests using $\Delta G^2 = G^2(M_2 | M_1)$ can assess whether the extra term(s) in model M_1 significantly improves the model fit. Beyond these tests, the pattern of residuals for individual cells offers important clues regarding the nature of lack of fit and can help suggests associations that could be accounted for better.

As with logistic regression models (Section 7.5.1), several types of residuals are available for loglinear models. For cell i in the vector form of the contingency table, the ***raw residual*** is simply the difference between the observed and fitted frequencies, $e_i = n_i - \hat{m}_i$.

The ***Pearson residual*** is the square root of the contribution of the cell to the Pearson χ^2 ,

$$r_i = \frac{n_i - \hat{m}_i}{\sqrt{\hat{m}_i}} \quad (8.11)$$

Similarly, the ***deviance residual*** can be defined as

$$g_i = \text{sign}(n_i - \hat{m}_i) \sqrt{2n_i \log(n_i/\hat{m}_i) - 2(n_i - \hat{m}_i)} \quad (8.12)$$

Both of these attempt to standardize the distribution of the residuals to a standard normal, $N(0, 1)$ form. However, as pointed out by Haberman (1973), the asymptotic variance of these is less than one (with average value df/N) but, worse—the variance decreases with \hat{m}_i . That is, residuals for cells with small expected frequencies have larger sampling variance, as might be expected.

Consequently, Haberman suggested dividing the Pearson residual by its estimated standard error, giving what are often called ***adjusted residuals***. When loglinear models are fit using the GLM approach, the adjustment may be calculated using the leverage (“hat value”), h_i to give appropriately standardized residuals,

$$\begin{aligned} r_i^* &= r_i / \sqrt{1 - h_i} \\ g_i^* &= g_i / \sqrt{1 - h_i} \end{aligned}$$

These standardized versions are generally preferable, particularly for visualizing model lack of fit using mosaic displays. The reason for preferring adjusted residuals is illustrated in Figure 8.1, a plot of the factors, $\sqrt{1 - h_i}$, determining the standard errors of the residuals against the fitted values, \hat{m}_i , in the model for the *UCBAmissions* data described in Example 8.2 below. The values shown in this plot are calculated as:

```
> berkeley <- as.data.frame(UCBAmissions)
> berk.glm1 <- glm(Freq ~ Dept * (Gender+Admit), data=berkeley, family="poisson")
> fit <- fitted(berk.glm1)
> hat <- hatvalues(berk.glm1)
> stderr <- sqrt(1-hat)
```

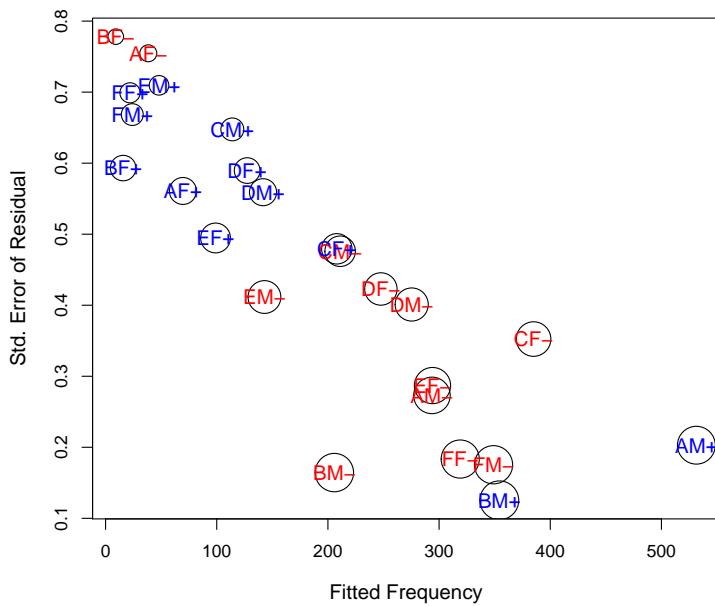


Figure 8.1: Standard errors of residuals, $\sqrt{1 - h_i}$ decrease with expected frequencies. This plot shows why ordinary Pearson and deviance residuals may be misleading. The symbol size in the plot is proportional to leverage, h_i . Labels abbreviate Department, Gender and Admit, colored by Admit.
[Fig. 8.1 is a scatter plot titled "Std. Error of Residual" on the y-axis (ranging from 0.1 to 0.8) versus "Fitted Frequency" on the x-axis (ranging from 0 to 500). The data points are labeled with three-letter abbreviations representing combinations of Department (D), Gender (G), and Admit status (A). Blue labels indicate Admit=Yes, while red labels indicate Admit=No. The size of the circles is proportional to leverage, h_i . A general downward trend is visible, where points with higher fitted frequencies tend to have lower standard errors of residuals.]

In R, raw, Pearson and deviance residuals may be obtained using `residuals(model, type=)`, where `type` is one of "raw", "pearson" and "deviance". Standardized (adjusted) residuals can be calculated using `rstandard(model, type=)`, for `type="pearson"` and `type="deviance"` versions.

8.3.4 Using `loglm()`

Here we illustrate the basics of fitting loglinear models using `loglm()`. As indicated in Section 8.3.1, the model to be fitted is specified by a model formula involving the table variables. The MASS package provides a `coef()` method for "loglm" objects that extracts the estimated parameters and a `residuals()` method that calculates various types of residuals according to a `type` argument, one of "deviance", "pearson", "response". `vcd` and `vcdExtra` provide a variety of plotting methods, including `assoc()`, `sieve()`, `mosaic()` and `mosaic3d()` for "loglm" objects.

{loglin-loglin}

{ex:berkeley5}

EXAMPLE 8.1: Berkeley admissions

The `UCBAdmissions` on admissions to the six largest graduate departments at U.C. Berkeley was examined using graphical methods in Chapter 4 (Example 4.14) and in Chapter 5 (Example 5.13). We can fit and compare several loglinear models as shown below.

The model of mutual independence, $[A][D][G]$, is not substantively reasonable here, because the association of `Dept` and `Gender` should be taken into account to control for these variables, but we show it here to illustrate the form of the printed output, giving the Pearson χ^2 and likelihood-ratio G^2 tests of goodness of fit, as well as some optional arguments for saving additional components in the result.

```
> data("UCBAdmissions")
> library(MASS)
> berk.loglm0 <- loglm(~ Dept + Gender + Admit, data=UCBAdmissions,
+                         param=TRUE, fitted=TRUE)
> berk.loglm0

Call:
loglm(formula = ~Dept + Gender + Admit, data = UCBAdmissions,
      param = TRUE, fitted = TRUE)

Statistics:
          X^2   df  P(> X^2)
Likelihood Ratio 2097.7 16      0
Pearson         2000.3 16      0
```

The argument `param=TRUE` stores the estimated parameters in the loglinear model and `fitted=TRUE` stores the fitted frequencies \hat{m}_{ijk} . The fitted frequencies can be extracted from the model object using `fitted()`.

```
> structable(Dept ~ Admit+Gender, fitted(berk.loglm0))

          Dept      A      B      C      D      E      F
Admit   Gender
Admitted Male    215.10 134.87 211.64 182.59 134.64 164.61
        Female   146.68  91.97 144.32 124.51  91.81 112.25
Rejected Male   339.63 212.95 334.17 288.30 212.59 259.91
        Female   231.59 145.21 227.87 196.59 144.96 177.23
```

Similarly, you can extract the estimated parameters with `coef(berk.loglm0)`, and the Pearson residuals with `residuals(berk.loglm0, type="pearson")`.

Next, consider the model of conditional independence of gender and admission given department, $[AD][GD]$ that allows associations of admission with department and gender with department.

```
> # conditional independence in UCB admissions data
> berk.loglm1 <- loglm(~ Dept * (Gender + Admit), data=UCBAdmissions)
> berk.loglm1

Call:
loglm(formula = ~Dept * (Gender + Admit), data = UCBAdmissions)

Statistics:
          X^2   df  P(> X^2)
Likelihood Ratio 21.736  6 0.0013520
Pearson         19.938  6 0.0028402
```

Finally for this example, the model of homogeneous association, $[AD][AG][GD]$ can be fit as follows.²

```
> berk.loglm2 <- loglm(~(Admit + Dept + Gender)^2, data=UCBAdmissions)
> berk.loglm2

Call:
loglm(formula = ~(Admit + Dept + Gender)^2, data = UCBAdmissions)

Statistics:
          X^2   df  P(> X^2)
Likelihood Ratio 20.204  5 0.0011441
Pearson         18.823  5 0.0020740
```

²It is useful to note here that the added term $[AG]$ allows a general association of admission with gender (controlling for department). A significance test for this term, or for model `berk.loglm2` against `berk.loglm1` is a proper test for the assertion of gender bias in admissions.

Neither of these models fits particularly well, as judged by the goodness-of-fit Pearson χ^2 and likelihood-ratio G^2 test against the saturated model. The `anova()` method for a nested collection of "loglm" models gives a series of likelihood-ratio tests of the difference, ΔG^2 between each sequential pair of models according to Eqn. (8.9).

```
> anova(berk.loglm0, berk.loglm1, berk.loglm2, test="Chisq")
LR tests for hierarchical log-linear models

Model 1:
~Dept + Gender + Admit
Model 2:
~Dept * (Gender + Admit)
Model 3:
~(Admit + Dept + Gender)^2

          Deviance df Delta(Dev)  Delta(df) P(> Delta(Dev)
Model 1    2097.671 16      21.736       6   2075.9357      10      0.000000
Model 2    21.736   6      20.204       5     1.5312       1      0.21593
Model 3    20.204   5     Saturated   0.000       0     20.2043       5      0.00114
```

The conclusion from these results is that the model `berk.loglm1` is not much worse than model `berk.loglm2`, but there is still significant lack-of-fit. The next example, using `glm()`, shows how to visualize the lack of fit and account for it.



8.3.5 Using `glm()`

Loglinear models fit with `glm()` require the data in a data frame in frequency form, for example as produced by `as.data.frame()` from a table. The model formula expresses the model for the frequency variable, and uses `family=poisson` to specify the error distribution. More general distributions for frequency data are discussed in Chapter 9.

{sec:loglin-glm}
{ex:berkeley6}

EXAMPLE 8.2: Berkeley admissions

For the $2 \times 2 \times 6$ *UCBAdmissions* table, first transform this to a frequency data frame:

```
> berkeley <- as.data.frame(UCBAdmissions)
> head(berkeley)

  Admit Gender Dept Freq
1 Admitted Male    A  512
2 Rejected  Male    A  313
3 Admitted Female  A   89
4 Rejected  Female  A   19
5 Admitted Male    B  353
6 Rejected  Male    B  207
```

Then, the model of conditional independence corresponding to `berk.loglm1` can be fit using `glm()` as shown below.

```
> berk.glm1 <- glm(Freq ~ Dept * (Gender+Admit),
+                      data=berkeley, family="poisson")
```

Similarly, the all two-way model of homogeneous association is fit using

```
> berk.glm2 <- glm(Freq ~ (Dept + Gender + Admit)^2,
+                         data=berkeley, family="poisson")
```

These models are equivalent to those fit using `loglm()` in Example 8.1. We get the same residual G^2 as before, and the likelihood-ratio test of ΔG^2 given by `anova()` gives the same result, that the model `berk.glm2` offers no significant improvement over model `berk.glm1`.

```
> anova(berk.glm1, berk.glm2, test="Chisq")

Analysis of Deviance Table

Model 1: Freq ~ Dept * (Gender + Admit)
Model 2: Freq ~ (Dept + Gender + Admit)^2
  Resid. Df Resid. Dev Df Deviance Pr(>Chi)
1          6    21.7
2          5    20.2  1      1.53     0.22
```

Among other advantages of using `glm()` as opposed to `loglm()` is that an `anova()` method is available for *individual "glm"* models, giving significance tests of the contributions of each *term* in the model, as opposed to the tests for individual coefficients provided by `summary()`.³

```
> anova(berk.glm1, test="Chisq")

Analysis of Deviance Table

Model: poisson, link: log

Response: Freq

Terms added sequentially (first to last)

  Df Deviance Resid. Df Resid. Dev Pr(>Chi)
NULL              23    2650
Dept      5       160     18    2491   <2e-16 ***
Gender    1       163     17    2328   <2e-16 ***
Admit     1       230     16    2098   <2e-16 ***
Dept:Gender 5      1221     11    877   <2e-16 ***
Dept:Admit  5       855      6    22   <2e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

We proceed to consider what is wrong with these models and how they can be improved. A mosaic display can help diagnose the reason(s) for lack of fit of these models. We focus here on the model $[AD][GD]$ that allows an association between gender and department (i.e., men and women apply at different rates to departments).

The `mosaic()` method for "glm" objects in `vcdExtra` provides a `residuals_type` argument, allowing `residuals_type="rstandard"` for standardized residuals. The `formula` argument here pertains to the order of the variables in the mosaic, not a model formula.

```
> library(vcdExtra)
> mosaic(berk.glm1, shade=TRUE, formula=~Admit+Dept+Gender,
+         residuals_type="rstandard", labeling=labeling_residuals,
+         main="Model: [AdmitDept] [GenderDept]")
```

³Unfortunately, in the historical development of R, the `anova()` methods for linear and generalized linear models provide only *sequential* ("Type I") tests that are computationally easy, but useful only under special circumstances. The `car` package provides an analogous `Anova()` method that gives more generally useful *partial* ("Type II") tests for the additional contribution of each term beyond the others, taking marginal relations into account.

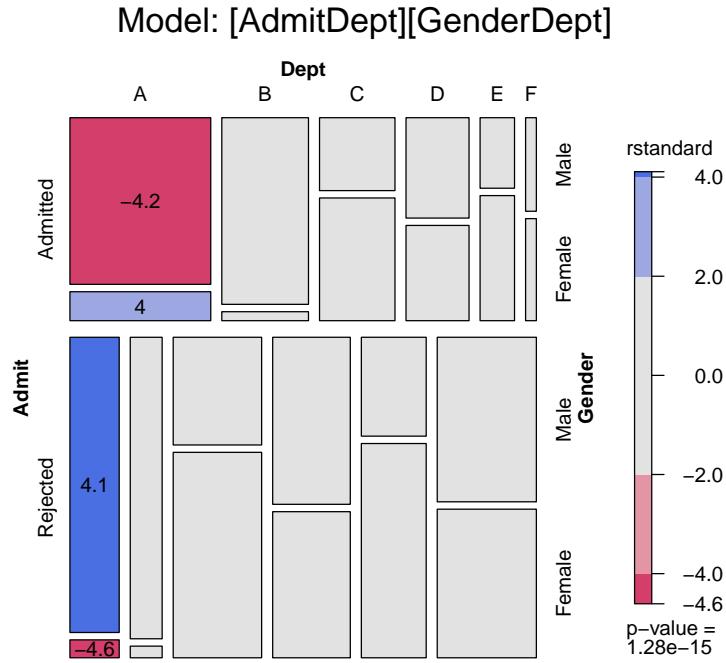


Figure 8.2: Mosaic display for the model $[AD][GD]$, showing standardized residuals for the cell contributions to G^2 | [fig:berk-glm1-mosaic](#)

The mosaic display, shown in Figure 8.2, indicates that this model fits well (residuals are small) except in Department A. This suggests a model which allows an association between Admission and Gender in Department A only,

$$\log m_{ijk} = \mu + \lambda_i^A + \lambda_j^D + \lambda_k^G + \lambda_{ij}^{AD} + \lambda_{jk}^{DG} + I(j=1)\lambda_{ik}^{AG}, \quad (8.13) \quad \{eq:berk2\}$$

where the indicator function $I(j=1)$ equals 1 for Department A ($j=1$) and is zero otherwise. This model asserts that Admission and Gender are conditionally independent, given Department, except in Department A. It has one more parameter than the conditional independence model, $[AD][GD]$, and forces perfect fit in the four cells for Department A.

Model Eqn. (8.13) may be fit with `glm()` by constructing a variable equal to the interaction of gender and admit with a dummy variable having the value 1 for Department A and 0 for other departments.

```
> berkeley <- within(berkeley,
+                         dept1AG <- (Dept=='A') * (Gender=='Female') * (Admit=='Admitted'))
> head(berkeley)

   Admit Gender Dept Freq dept1AG
1 Admitted   Male    A  512      0
2 Rejected   Male    A  313      0
3 Admitted Female   A   89      1
4 Rejected Female   A   19      0
5 Admitted   Male   B  353      0
6 Rejected   Male   B  207      0
```

Fitting this model with the extra term `dept1AG` gives `berk.glm3`

```
> berk.glm3 <- glm(Freq ~ Dept * (Gender+Admit) + dept1AG,
+                         data=berkeley, family="poisson")
```

This model does indeed fit well, and represents a substantial improvement over model `berk.glm1`:

```
> vcdExtra::LRstats(berk.glm3)

Likelihood summary table:
      AIC BIC LR Chisq Df Pr(>Chisq)
berk.glm3 200 222     2.68   5      0.75

> anova(berk.glm1, berk.glm3, test="Chisq")

Analysis of Deviance Table

Model 1: Freq ~ Dept * (Gender + Admit)
Model 2: Freq ~ Dept * (Gender + Admit) + dept1AG
  Resid. Df Resid. Dev Df Deviance Pr(>Chi)
1          6    21.74
2          5     2.68   1     19.1  1.3e-05 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The parameter estimate for the `dept1AG` term, $\hat{\lambda}_{ik}^{AG} = 1.052$ may be interpreted as the log odds ratio of admission for females as compared to males in Dept. A. The odds ratio is $\exp(1.052) = 2.86$, the same as the value calculated from the raw data (see Section 4.4.2).

```
> coef(berk.glm3)[["dept1AG"]]
[1] 1.0521
> exp(coef(berk.glm3)[["dept1AG"]])
[1] 2.8636
```

Finally, Figure 8.3 shows the mosaic for this revised model. The absence of shading indicates a well-fitting model.

```
> mosaic(berk.glm3, shade=TRUE, formula=~Admit+Dept+Gender,
+           residuals_type="rstandard", labeling=labeling_residuals,
+           main="Model: [DeptGender] [DeptAdmit] + DeptA* [GA]")
```



8.4 Equivalent logit models

{sec:loglin-logit} Because loglinear models are formulated as models for the log (expected) frequency, they make no distinction between response and explanatory variables. In effect, they treat all variables as responses and describe their associations.

Logit (logistic regression) models, on the other hand, describe how the log odds for one variable depends on other, explanatory variables. There is a close connection between the two: When there is a response variable, each logit model for that response is equivalent to a loglinear model.

This relationship often provides a simpler way to formulate and test the model, and to plot and interpret the fitted results. Even when there is no response variable, the logit representation for one variable helps to interpret a loglinear model in terms of odds ratios. The price paid for this simplicity is that associations among the explanatory variables are not expressed in the model.

Model: [DeptGender][DeptAdmit] + DeptA*[GA]

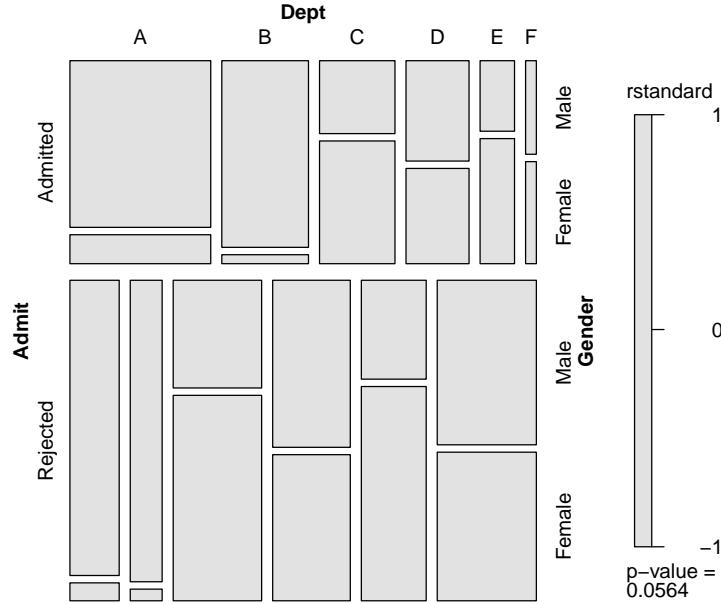


Figure 8.3: Mosaic display for the model `berk.glm3`, allowing an association of gender and admission in Department A. This model now fits the data well.
Fig:berk.glm3_mosaic

Consider, for example, the model of homogeneous association, $[AB][AC][BC]$, Eqn. (8.5) for a three-way table, and let variable C be a binary response. Under this model, the logit for variable C is

$$\begin{aligned} L_{ij} = \log \left(\frac{\pi_{ij|1}}{\pi_{ij|2}} \right) &= \log \left(\frac{m_{ij1}}{m_{ij2}} \right) \\ &= \log(m_{ij1}) - \log(m_{ij2}) . \end{aligned}$$

Substituting from Eqn. (8.5), all terms which do not involve variable C cancel, and we are left with

$$\begin{aligned} L_{ij} = \log(m_{ij1}/m_{ij2}) &= (\lambda_1^C - \lambda_2^C) + (\lambda_{i1}^{AC} - \lambda_{i2}^{AC}) + (\lambda_{j1}^{BC} - \lambda_{j2}^{BC}) \\ &= 2\lambda_1^C + 2\lambda_{i1}^{AC} + 2\lambda_{j1}^{BC} , \end{aligned} \tag{8.14} \quad \text{(eq:logitab1)}$$

because all λ terms sum to zero. We are interested in how these logits depend on A and B , so we can simplify the notation by replacing the λ parameters with more familiar ones, $\alpha = 2\lambda_1^C$, $\beta_i^A = 2\lambda_{i1}^{AC}$, etc., which express this relation more directly,

$$L_{ij} = \alpha + \beta_i^A + \beta_j^B . \tag{8.15} \quad \text{(eq:logitab2)}$$

In the logit model Eqn. (8.15), the response, C , is affected by both A and B , which have additive effects on the log odds of response category C_1 compared to C_2 . The terms β_i^A and β_j^B correspond directly to $[AC]$ and $[BC]$ in the loglinear model Eqn. (8.5). The association among the explanatory variables, $[AB]$ is assumed in the logit model, but this model provides no explicit representation of that association. The logit model Eqn. (8.14) is equivalent to the loglinear model $[AB][AC][BC]$ in goodness-of-fit and fitted values, and parameters in the two models correspond directly.

Table 8.1: Equivalent loglinear and logit models for a three-way table, with C as a binary response variable.

{tab:loglin-logit}

Loglinear model	Logit model	Logit formula
$[AB][C]$	α	$C \sim 1$
$[AB][AC]$	$\alpha + \beta_i^A$	$C \sim A$
$[AB][BC]$	$\alpha + \beta_j^B$	$C \sim B$
$[AB][AC][BC]$	$\alpha + \beta_i^A + \beta_j^B$	$C \sim A + B$
$[ABC]$	$\alpha + \beta_i^A + \beta_j^B + \beta_{ij}^{AB}$	$C \sim A * B$

Table 8.1 shows the equivalent relationships between all loglinear and logit models for a three-way table when variable C is a binary response. Each model necessarily includes the $[AB]$ association involving the predictor variables. The most basic model, $[AB][C]$, is the intercept-only model, asserting constant odds for variable C . The saturated loglinear model $[ABC]$, allows an interaction in the effects of A and B on C , meaning that the AC association or odds ratio varies with B .

More generally, when there is a binary response variable, say R , and one or more explanatory variables, A, B, C, \dots , any logit model for R has an equivalent loglinear form. Every term in the logit model, such as β_{ik}^{AC} , corresponds to an association of those factors with R , that is, $[ACR]$ in the equivalent loglinear model.

The equivalent loglinear model must also include all associations among the explanatory factors, the term $[ABC \dots]$. Conversely, any loglinear model which includes all associations among the explanatory variables has an equivalent logit form. When the response factor has more than two categories, models for generalized logits (Section 7.6.4) also have an equivalent loglinear form.

{ex:berkeley7}

EXAMPLE 8.3: Berkeley admissions

The homogeneous association model, $[AD][AG][DG]$ did not fit the *UCBadmissions* data very well, and we saw that the term $[AG]$ was unnecessary. Nevertheless, it is instructive to consider the equivalent logit model. We illustrate the features of the logit model which lead to the same conclusions and simplified interpretation from graphical displays.

Because Admission is a binary response variable, model Eqn. (8.6) is equivalent to the logit model,

$$\text{eq:berk3} \quad L_{ij} = \log \left(\frac{m_{\text{Admit}(ij)}}{m_{\text{Reject}(ij)}} \right) = \alpha + \beta_i^{\text{Dept}} + \beta_j^{\text{Gender}} . \quad (8.16)$$

That is, the logit model Eqn. (8.16) asserts that department and gender have additive effects on the log odds of admission. A significance test for the term β_j^{Gender} here is equivalent to the test of the $[AG]$ term for gender bias in the loglinear model. The observed log odds of admission here can be calculated as:

```
> (obs <- log(UCBAdmissions[1,,] / UCBAdmissions[2,,]))
```

		Dept	A	B	C	D	E	F
Gender	Male	0.4921	0.5337	-0.5355	-0.704	-0.957	-2.770	
Female	1.5442	0.7538	-0.6604	-0.622	-1.157	-2.581		

With the data in the form of the frequency data frame *berkeley* we used in Example 8.2, the logit model Eqn. (8.16) can be fit using `glm()` as shown below. In the model formula, the binary response is `Admit=="Admitted"`. The `weights` argument gives the frequency, `Freq` in each table cell.⁴

⁴Using weights gives the same fitted values, but not the same LR tests for model fit.

```

> berk.logit2 <- glm(Admit=="Admitted" ~ Dept + Gender,
+                         data=berkeley, weights=Freq, family="binomial")
> summary(berk.logit2)

Call:
glm(formula = Admit == "Admitted" ~ Dept + Gender, family = "binomial",
      data = berkeley, weights = Freq)

Deviance Residuals:
    Min      1Q   Median      3Q     Max 
-25.342 -13.058 -0.163  16.017  21.320 

Coefficients:
            Estimate Std. Error z value Pr(>|z|)    
(Intercept)  0.5821    0.0690   8.44   <2e-16 ***  
DeptB       -0.0434    0.1098  -0.40    0.69    
DeptC       -1.2626    0.1066  -11.84  <2e-16 ***  
DeptD       -1.2946    0.1058  -12.23  <2e-16 ***  
DeptE       -1.7393    0.1261  -13.79  <2e-16 ***  
DeptF       -3.3065    0.1700  -19.45  <2e-16 ***  
GenderFemale 0.0999    0.0808   1.24    0.22    
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for binomial family taken to be 1)

Null deviance: 6044.3 on 23 degrees of freedom
Residual deviance: 5187.5 on 17 degrees of freedom
AIC: 5201

Number of Fisher Scoring iterations: 6

```

As in logistic regression models, parameter estimates may be interpreted as increments in the log odds, or $\exp(\beta)$ may be interpreted as the multiple of the odds associated with the explanatory categories. Because **glm()** uses a baseline category parameterization (by default) the coefficients of the first category of Dept and Gender are set to zero. You can see from the **summary()** output that the coefficients for the departments decline steadily from A–F.⁵ The coefficient $\beta_F^{\text{Gender}} = 0.0999$ for females indicates that, overall, women were $\exp(0.0999) = 1.105$ times as likely as male applicants to be admitted to graduate school at U.C. Berkeley, a 10% advantage.

Similarly, the logit model equivalent of the loglinear model Eqn. (8.13) **berk.glm3** containing the extra 1 df term for an effect of gender in Department A is

$$L_{ij} = \alpha + \beta_i^{\text{Dept}} + I(j=1)\beta^{\text{Gender}}. \quad (8.17) \quad \text{(eq:berk4)}$$

This model can be fit as follows:

```

> berkeley <- within(berkeley,
+                         dept1AG <- (Dept=='A') * (Gender=='Female'))
> berk.logit3 <- glm(Admit=="Admitted" ~ Dept + Gender + dept1AG,
+                         data=berkeley, weights=Freq, family="binomial")

```

In contrast to the tests for individual coefficients, the **Anova()** method in the **car** package gives likelihood-ratio tests of the terms in a model. As mentioned earlier, this provides *partial* (“Type II”) tests for the additional contribution of each term beyond all others.

⁵In fact, the departments were labeled A–F in decreasing order of rate of admission.

```
> library(car)
> Anova(berk.logit2)

Analysis of Deviance Table (Type II tests)

Response: Admit == "Admitted"
          LR Chisq Df Pr(>Chisq)
Dept      763.4  5    <2e-16 ***
Gender     1.5   1    0.216
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

> Anova(berk.logit3)

Analysis of Deviance Table (Type II tests)

Response: Admit == "Admitted"
          LR Chisq Df Pr(>Chisq)
Dept      646.7  5    < 2e-16 ***
Gender     0.1   1    0.724
dept1AG   17.6   1    2.66e-05 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Plotting logit models

Logit models are easier to interpret than the corresponding loglinear models because there are fewer parameters, and because these parameters pertain to the odds of a response category rather than to cell frequency. Nevertheless, interpretation is often easier still from a graph than from the parameter values.

The simple interpretation of these logit models can be seen by plotting the logits for a given model. To do that, it is necessary to construct a data frame containing the observed (`obs`) and fitted (`fit`) for the combinations of gender and department.

```
> pred2 <- cbind(berkeley[,1:3], fit=predict(berk.logit2))
> pred2 <- cbind(subset(pred2, Admit=="Admitted"), obs=as.vector(obs) )
> head(pred2)

  Admit Gender Dept      fit      obs
1 Admitted Male     A  0.58205  0.49212
3 Admitted Female   A  0.68192  1.54420
5 Admitted Male     B  0.53865  0.53375
7 Admitted Female   B  0.63852  0.75377
9 Admitted Male     C -0.68055 -0.53552
11 Admitted Female  C -0.58068 -0.66044
```

In this form, these results can be plotted as a line plot of the fitted logits vs. department, with separate curves for males and females, and adding points to show the observed values. Here, we use `ggplot2` as shown below, with the `aes()` arguments `group=Gender`, `color=Gender`. This produces the left panel in Figure 8.4. The same steps for the model `berk.logit3` gives the right panel in this figure. The observed logits, of course, are the same in both plots.

```
> library(ggplot2)
> ggplot(pred2, aes(x=Dept, y=fit, group=Gender, color=Gender)) +
+   geom_line(size=1.2) +
+   geom_point(aes(x=Dept, y=obs, group=Gender, color=Gender), size=4) +
+   ylab("Log odds (Admitted)") + theme_bw() +
+   theme(legend.position=c(.8, .9),
+         legend.title=element_text(size=14),
```

```
+     legend.text=element_text(size=14))
```

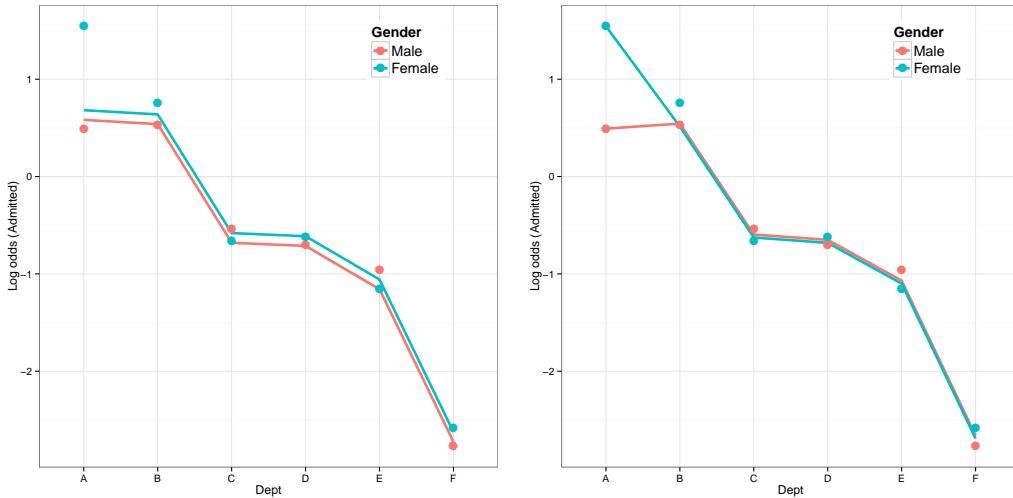


Figure 8.4: Observed (points) and fitted (lines) log odds of admissions in the logit models for the *UCBAdmissions* data. Left: the logit model Eqn. (8.16) corresponding to the loglinear model [AD] [AG] [DG]. Right: the logit model Eqn. (8.17), allowing only a 1 df term for Department A.

{fig:berk-logit}

The effects seen in our earlier analyses (Examples 5.13, 5.14 and 8.2) may all be observed in these plots. In the left panel of Figure 8.4, corresponding to the loglinear model [AD][AG][DG], the effect of gender, β_j^{Gender} , in the equivalent logit model is shown by the constant separation between the two curves. From the plot we see that this effect is very small (and nonsignificant). In the right panel, corresponding to the logit model Eqn. (8.17), there is no effect of gender on admission, except in department A, where the extra parameter allows perfect fit.

△

8.5 Zero frequencies

{sec:loglin-zeros}

Cells with frequencies of zero create problems for loglinear and logit models. For loglinear models, most of the derivations of expected frequencies by maximum likelihood and other quantities that depend on these (e.g., G^2 tests) assume that all $n_{ijk\dots} > 0$. In analogous logit models, the observed log odds (e.g., for a three-way table), $\log(n_{ij1}/n_{ij2})$, will be undefined if either frequency is zero.

Zero frequencies may occur in contingency tables for two different reasons:

- **structural zeros** (also called *fixed zeros*) will occur when it is impossible to observe values for some combinations of the variables. For these cases we should have $\hat{m}_i = 0$ wherever $n_i = 0$. For example, suppose we have three different methods of contacting people at risk for some obscure genetically inherited disease: newspaper advertisement, telephone campaign, and radio appeal. If each person contacted in any way is classified dichotomously by the three methods of contact, there can never be a non-zero frequency in the ‘No-No-No’ cell.⁶ Similarly, in a tabulation of seniors by gender and health concerns, there can never be males citing menopause

⁶Yet, if we fit an unsaturated model, expected frequencies may be estimated for all cells, and provide a means to estimate the total number at risk in the population. See Lindsey (1995, Section 5.4).

or females citing prostate cancer. Square tables, such as wins and losses for sporting teams often have structural zeros in the main diagonal.

- **sampling zeros** (also called *random zeros*) occur when the total size of the sample is not large enough in relation to the probabilities in each of the cells to assure that someone will be observed in every cell. Here, it is permissible to have $\hat{m}_i > 0$ when $n_i = 0$. This problem increases with the number of table variables. For example, in a European survey of religious affiliation, gender and occupation, we may not happen to observe any female Muslim vineyard-workers in France, although such individuals surely exist in the population. Even when zero frequencies do not occur, tables with many cells relative to the total frequency tend to produce small expected frequencies in at least some cells, which tends to make the G^2 statistics for model fit and likelihood-ratio statistics for individual terms unreliable.

Following Birch (1963b), Haberman (1974) and many others (e.g., Bishop *et al.*, 1975) identified conditions under which the maximum likelihood estimate for a given loglinear model does not exist, meaning that the algorithms used in `loglin()` and `glm()` do not converge to a solution. The problem depends on the number and locations of the zero cells, but not on the size of the frequencies in the remaining cells. Fienberg and Rinaldo (2007) give a historical overview of the problem and current approaches and Agresti (2013, §10.6) gives a compact summary.

In R, the mechanism to handle structural zeros in the IPF approach of `loglin()` and `loglm()` is to supply the argument `start`, giving a table conforming to the data, containing values of 0 in the locations of the zero cells, and non-zero elsewhere.⁷ In the `glm()` approach, the argument `subset=Freq > 0` can be used to remove the cells with zero frequencies from the data, or else, zero frequencies can be set to `NA`. This usually provides the correct degrees of freedom, however some estimated coefficients may be infinite.

For a complete table, the residual degrees of freedom are determined as

$$df = \# \text{ of cells} - \# \text{ of fitted parameters}$$

For tables with structural zeros, an analogous general formula is

$$\{eq:dfzeros\} \quad df = (\# \text{ cells} - \# \text{ of parameters}) - (\# \text{ zero cells} - \# \text{ of NA parameters}) \quad (8.18)$$

where NA parameters refers to parameters that cannot be estimated due to zero marginal totals in the model formula.

In contrast, sampling zeros are often handled by some modification of the data frequencies to ensure all non-zero cells. Some suggestions are:

- Add a small positive quantity (0.5 is often recommended) to *every* cell in the contingency table (Goodman, 1970), as is often done in calculating empirical log odds (Example 8.13); this simple approach over-smooths the data for unsaturated models, and should be deprecated, although widely used in practice.
- Replace sampling zeros by some small number, typically 10^{-10} or smaller (Agresti, 1990).
- Add a small quantity, like 0.1, to *all* zero cells, sampling or structural (Evers and Namboordiri., 1977).

In complex, sparse tables, a sensitivity analysis, comparing different approaches can help determine if the substantive conclusions vary with the approach to zero cells.

⁷If structural zeros are present, the calculation of degrees of freedom may not be correct. `loglm()` deducts one degree of freedom for each structural zero, but cannot make allowance for patterns of zeros based on the fitted margins that lead to gains in degrees of freedom due to smaller dimension in the parameter space. `loglin()` makes no such correction.

EXAMPLE 8.4: Health concerns of teenagers

Fienberg (1980, Table 8-3) presented a classic example of structural zeros in the analysis of the $4 \times 2 \times 2$ table shown in Table 8.2. The data come from a survey of health concerns among teenagers, originally from Brunswick (1971). Among the health concerns, the two zero entries for menstrual problems among males are clearly structural zeros and there therefore one structural zero in the concern by gender marginal table. As usual, we abbreviate the table variables concern, age, gender by their initial letters, C, A, G below.

Table 8.2: Results from a survey of teenagers, regarding their health concerns. Two cells with structural zeros are highlighted. *Source:* Fienberg (1980, Table 8-3)

{tab:health}

Health Concerns	Gender: Age:	Male		Female	
		12-15	16-17	12-15	16-17
sex, reproduction		4	2	9	7
menstrual problems		0	0	4	8
how healthy I am		42	7	19	10
nothing		57	20	71	21

The *Health* data is created as a frequency data frame as follows.

```
> Health <- expand.grid(concerns = c("sex", "menstrual",
+                                         "healthy", "nothing"),
+                         age      = c("12-15", "16-17"),
+                         gender   = c("M", "F"))
> Health$Freq <- c(4, 0, 42, 57, 2, 0, 7, 20,
+                  9, 4, 19, 71, 7, 8, 10, 21)
```

In this form, we first use `glm()` to fit two small models, neither of which involves the $\{CG\}$ margin. Model `health.glm0` is the model of mutual independence, $[C][A][G]$. Model `health.glm1` is the model of joint independence, $[C][AG]$, allowing an association between age and gender, but neither with concern. As noted above, the argument `subset=(Freq>0)` is used to eliminate the structural zero cells.

```
> health.glm0 <- glm(Freq ~ concerns + age + gender, data=Health,
+                      subset=(Freq>0), family=poisson)
> health.glm1 <- glm(Freq ~ concerns + age * gender, data=Health,
+                      subset=(Freq>0), family=poisson)
```

Neither of these fits the data well. To conserve space, we show only the results of the G^2 tests for model fit.

```
> vcdExtra::LRstats(health.glm0, health.glm1)

Likelihood summary table:
          AIC BIC LR Chisq Df Pr(>Chisq)
health.glm0 100.7 105    27.7  8    0.00053 ***
health.glm1  99.9 104    24.9  7    0.00080 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

To see why, Figure 8.5 shows the mosaic display for model `health.glm1`, $[C][AG]$. Note that `mosaic()` takes care to make cells of zero frequency more visible by marking them with a small “o”, as these have an area of zero.

```
> mosaic(health.glm1, ~concerns+age+gender, residuals_type="rstandard")
```

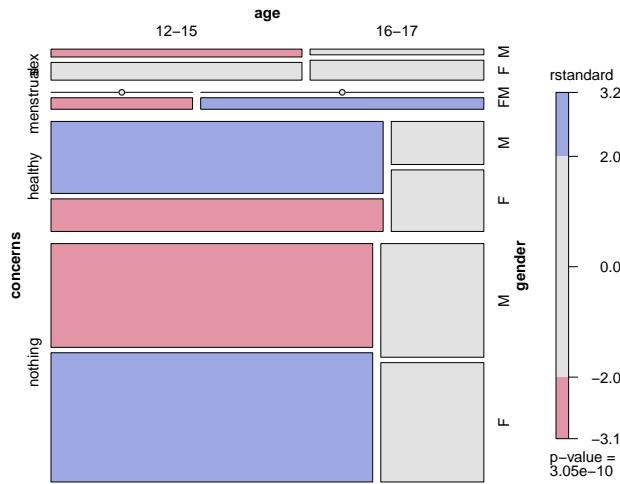


Figure 8.5: Mosaic display for the Health data, model `health.glm1` fig:health-mosaic

This suggests that there are important associations at least between concern and gender ($[CG]$) and between concern and age ($[CA]$). These are incorporated into the next model:

```
> health.glm2 <- glm(Freq ~ concerns*gender + concerns*age, data=Health,
+                       subset=(Freq>0), family=poisson)
> vcdExtra::LRstats(health.glm2)

Likelihood summary table:
      AIC    BIC   LR Chisq Df Pr(>Chisq)
health.glm2 87.7 94.7     4.66  3        0.2
```

The degrees of freedom are correct here. Eqn. (8.18), with 2 zero cells and 1 NA parameter due to the zero in the $\{CG\}$ margin gives $df = (16 - 12) - (2 - 1) = 3$. The loss of one estimable parameter can be seen in the output from `summary`.

```
> summary(health.glm2)

Call:
glm(formula = Freq ~ concerns * gender + concerns * age, family = poisson,
     data = Health, subset = (Freq > 0))

Deviance Residuals:
    1      3      4      5      7      8      9      10     11     12 
  0.236  0.585 -0.173 -0.300 -1.202  0.302 -0.149  0.000 -0.795  0.158 
    13     14     15     16 
  0.176  0.000  1.348 -0.282 

Coefficients: (1 not defined because of singularities)
              Estimate Std. Error z value Pr(>|z|)    
(Intercept)  1.266     0.445   2.84   0.0045 **  
concernsmenstrual -0.860     0.586  -1.47   0.1425    
concernshealthy   2.380     0.471   5.05  4.4e-07 ***  
concernsnothing   2.800     0.462   6.07  1.3e-09 ***  
genderF          0.981     0.479   2.05   0.0405 *   
age16-17         -0.368     0.434  -0.85   0.3964    
concernsmenstrual:genderF NA        NA      NA      NA      
```

```

concernshealthy:genderF    -1.505     0.533   -2.82   0.0047 **
concernsnothing:genderF   -0.803     0.503   -1.60   0.1105
concernsmenstrual:age16-17 1.061     0.750   1.41    0.1574
concernshealthy:age16-17   -0.910     0.513   -1.77   0.0761 .
concernsnothing:age16-17   -0.771     0.469   -1.64   0.1005
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for poisson family taken to be 1)

Null deviance: 252.4670  on 13  degrees of freedom
Residual deviance: 4.6611  on  3  degrees of freedom
AIC: 87.66

Number of Fisher Scoring iterations: 4

```

In contrast, `loglm()` reports the degrees of freedom incorrectly for models containing zeros in any fitted margin. For use with `loglm()`, we convert it to a $4 \times 2 \times$ table.

```
> health.tab <- xtabs(Freq ~ concerns + age + gender, data = Health)
```

The same three models are fitted with `loglm()` as shown below. The locations of the positive frequencies are marked in the array `nonzeros` and supplied as the value of the `start` argument.

```

> nonzeros <- ifelse(health.tab>0, 1, 0)
> health.loglm0 <- loglm(~ concerns + age + gender,
+   data = health.tab, start = nonzeros)
> health.loglm1 <- loglm(~ concerns + age * gender,
+   data = health.tab, start = nonzeros)
> # df is wrong
> health.loglm2 <- loglm(~ concerns*gender + concerns*age,
+   data = health.tab, start = nonzeros)
> LRstats(health.loglm0, health.loglm1, health.loglm2)

Likelihood summary table:
      AIC BIC LR Chisq Df Pr(>Chisq)
health.loglm0 104.7 111   27.74  8   0.00053 ***
health.loglm1 103.9 111   24.89  7   0.00080 ***
health.loglm2  93.7 104    4.66  2   0.09724 .
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

The results agree with those of `glm()`, except for the degrees of freedom for the last model.



8.6 Models for ordinal variables

Standard loglinear models treat all classification variables as nominal, unordered factors. In these models, all statistical tests are identical and parameter estimates are equivalent if the categories of any of the table variable are reordered. Yet we have seen that the ordering of categories often provides important information about the nature of associations and we showed (Section 4.2.4) that non-parametric tests which take into account the ordered nature of a factor are more powerful.

(sec:loglin-ordinal)

Correspondence analysis plots (Chapter 6) make it easy to see the relationships between ordinal variables, because the method assigns quantitative scores to the table variables which maximally account for their association. As we saw for the hair-eye color data (Figure 6.1) and the mental impairment data (Figure 6.2), an association can be interpreted in terms of ordered categories when the points for two factors are ordered similarly, usually along the first CA dimension.

Similarly, in a mosaic display, an ordered associative effect is seen when the residuals have an

opposite-corner pattern of positive and negative signs and magnitudes (e.g., for the hair-eye color data, Figure 5.4). In these cases loglinear and logit models which use the ordered nature of the factors offer several advantages.

- Because they are more focused, tests which use the ordinal structure of the table variables are more powerful when the association varies systematically with the ordered values of a factor.
- Because they consume fewer degrees of freedom, we can fit unsaturated models where the corresponding model for nominal factors would be saturated. In a two-way table, for example, a variety of models for ordinal factors may be proposed which are intermediate between the independence model and the saturated model.
- Parameter estimates from these models are fewer in number, are easier to interpret, and quantify the nature of effects better than corresponding quantities in models for nominal factors. Estimating fewer parameters typically gives smaller standard errors.

These advantages are analogous to the use of tests for trends or polynomial contrasts in ANOVA models. More importantly, in some research areas in the social sciences (where categorical data is commonplace), models for ordinal variables have proved crucial in theory construction and debates, giving more precise tests of hypotheses than available from less focused or descriptive methods (Agresti, 1984).

8.6.1 Loglinear models for ordinal variables

For a two-way table, when either the row variable or the column variable, or both, are ordinal, one simplification comes from assigning ordered scores, $\mathbf{a} = \{a_i\}, a_1 \leq a_2 \leq \dots a_I$, and/or $\mathbf{b} = \{b_j\}, b_1 \leq b_2 \leq \dots b_J$ to the categories so that the ordinal relations are necessarily included in the model. Typically, equally spaced scores are used, for example, integer scores, $\{a_i\} = i$, or the zero-sum equivalent, $\{a_i\} = i - (I + 1)/2$ (e.g., $\{a_i\} = \{-1, 0, 1\}$ for $I = 3$).

Using such scores gives simple interpretations of the association parameters in terms of *local odds ratios* for adjacent 2×2 subtables,

$$\theta_{ij} = \frac{m_{ij} m_{i+1,j+1}}{m_{i,j+1} m_{i+1,j}}, \quad (8.19)$$

which is the odds ratio for pairs of adjacent rows and adjacent columns.

When both variables are assigned scores, this gives the ***linear-by-linear model*** ($L \times L$)

$$\log(m_{ij}) = \mu + \lambda_i^A + \lambda_j^B + \gamma a_i b_j. \quad (8.20)$$

Because the scores \mathbf{a} and \mathbf{b} are fixed, this model has only one extra parameter, γ , compared to the independence model, which is the special case, $\gamma = 0$. In contrast, the saturated model, allowing general association λ_{ij}^{AB} uses $(I - 1)(J - 1)$ additional parameters.

The terms $\gamma a_i b_j$ in Eqn. (8.20) describe a pattern of association where deviations from independence increase linearly with a_i and b_j in opposite directions towards the opposite corners of the table, as we have often observed in mosaic displays.

In the linear-by-linear association model, the local log odds ratios are

$$\log(\theta_{ij}) = \gamma(a_{i+1} - a_i)(b_{j+1} - b_j),$$

which reduces to

$$\log(\theta_{ij}) = \gamma$$

for integer-spaced scores, so γ is the common local log odds ratio. As a result, the linear-by-linear model is sometimes called the ***uniform association model*** (Goodman, 1979).

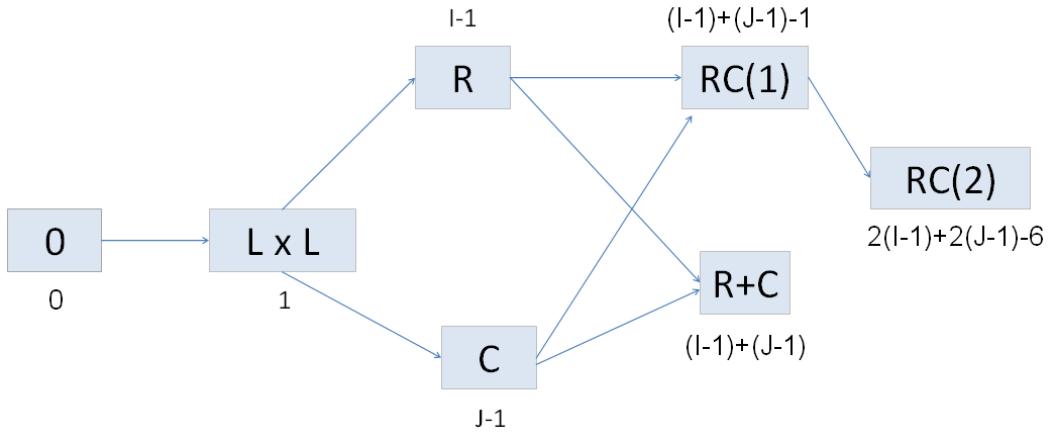


Figure 8.6: Nesting relationships among some association models for an $I \times J$ table specifying the association parameters, λ_{ij}^{AB} . Model **0** is the independence model. Formulas near the boxes give the number of identifiable association parameters. Arrows point from one nested model to another that is a more general version.

{fig:assoc-models}

Generalizations of the linear-by-linear model result when only one variable is assigned scores. In the **row effects model** (R), the row variable, A , is treated as nominal, while the column variable, B , is assigned ordered scores $\{b_j\}$. The loglinear model is then

$$\log(m_{ij}) = \mu + \lambda_i^A + \lambda_j^B + \alpha_i b_j , \quad (8.21)$$

where the α_i parameters are the **row effects**. An additional constraint, $\sum_i \alpha_i = 0$ or $\alpha_1 = 0$ is imposed, so that model Eqn. (8.21) has only $(I - 1)$ more parameters than the independence model. The linear-by-linear model is the special case where the row effects are equally spaced, and the independence model is the special case where all $\alpha_i = 0$.

The row-effects model Eqn. (8.21) also has a simple odds ratio interpretation. The local log odds ratio for adjacent pairs of rows and columns is

$$\log(\theta_{ij}) = \alpha_{i+1} - \alpha_i ,$$

which is constant for all pairs of adjacent columns. Plots of the local log odds ratio against i would appear as a set of parallel curves.

In the analogous **column effects model** (C), $(J - 1)$ linearly independent column effect parameters β_j are estimated for the column variable, while fixed scores $\{a_i\}$ are assigned to the row variable. It is also possible to fit a **row plus column effects model** (R+C), that assigns specified scores to both the rows and column variables.

Nesting relationships among these models and others described in Section 8.6.2 are shown in Figure 8.6. Any set of models connected by a path can be directly compared with likelihood-ratio tests of the form $G^2(M_2|M_1)$.

In R, the $L \times L$, row effects and column effects models can all be fit using `glm()` simply by replacing the appropriate table factor variable(s) with their `as.numeric()` equivalents.

{ex:mental4}

EXAMPLE 8.5: Mental impairment and parents' SES

The `Mental` data on the mental health status of young New York residents in relation to their parents' socioeconomic status was examined in Example 4.6 using CMH tests for ordinal association and in Example 6.2 using correspondence analysis. Figure 6.2 showed that nearly all of the

association in the table was accounted for by a single dimension along which both factors were ordered, consistent with the view that mental health increased in relation to parents' SES.

Because these models provide their interpretations in terms of local odds ratios, Eqn. (8.19), it is helpful to see these values for the observed data, corresponding to the saturated model. The values $\log(\theta_{ij})$ are calculated by `loddsratio()` in `vcdExtra`, with the data in table form.

```
> (mental.tab <- xtabs(Freq ~ mental+ses, data=Mental))

      ses
mental    1   2   3   4   5   6
  Well    64  57  57  72  36  21
  Mild    94  94 105 141  97  71
  Moderate 58  54  65  77  54  54
  Impaired 46  40  60  94  78  71

> loddsratio(mental.tab)

log odds ratios for mental and ses

      ses
mental          1:2     2:3     3:4     4:5     5:6
  Well:Mild      0.1158  0.1107  0.0612  0.3191  0.227
  Mild:Moderate -0.0715  0.0747 -0.1254  0.0192  0.312
  Moderate:Impaired -0.0683  0.2201  0.2795  0.1682 -0.094
```

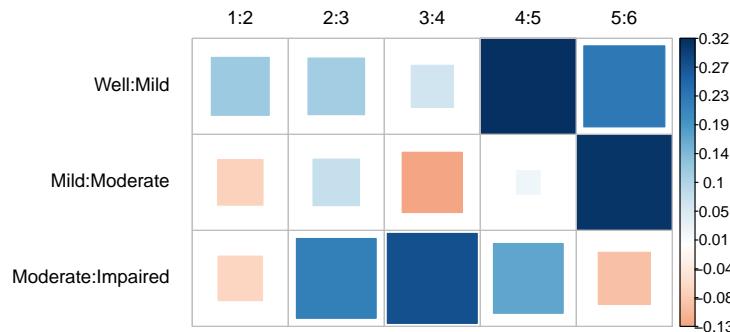


Figure 8.7: Shaded-square plot of the local odds ratios in the *Mental* data.

A simple plot of these values, using area- and color-proportional shaded squares is shown in Figure 8.7. This plot is drawn using the `corrplot` package. It is easy to see that most of the local odds ratios are mildly positive.

```
> M <- as.matrix(loddsratio(mental.tab))
> library(corrplot)
> corrplot(M, method="square", is.corr=FALSE,
+           tl.col="black", tl.srt=0, tl.offset=1)
```

For comparison with the $L \times L$ model fitted below, the mean local log odds ratio is 0.103.

```
> mean(loddsratio(mental.tab)$coefficients)
[1] 0.10323
```

As a baseline, we first fit the independence model (testing $H_0 : \log(\theta_{ij}) = 0$) with `glm()`. As expected, this model fits quite badly, with $G^2(15) = 47.418$.

```
> indep <- glm(Freq ~ mental + ses,
+   family = poisson, data = Mental)
> vcdExtra::LRstats(indep)

Likelihood summary table:
  AIC BIC LR Chisq Df Pr(>Chisq)
indep 210 220     47.4 15    3.2e-05 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The mosaic display of standardized residuals from this model is shown in Figure 8.8. The argument `labeling=labeling_residuals` is used to show the numerical values in the cells with absolute values greater than `suppress=1`.

```
> long.labels <- list(set_varnames = c(mental="Mental Health Status",
+   ses="Parent SES"))
> mosaic(indep,
+   gp=shading_Friendly,
+   residuals_type="rstandard",
+   labeling_args = long.labels,
+   labeling=labeling_residuals, suppress=1,
+   main="Mental health data: Independence")
```



Figure 8.8: Mosaic display of the independence model for the mental health data.^{fig:mental-indep}

This figure shows the classic opposite-corner pattern of the signs and magnitudes of the residuals that would arise if the association between mental health and SES was could be explained by the ordinal relation of these factors using one of the $L \times L$, R or C models.

To fit such ordinal models, you can use `as.numeric()` on a factor variable to assign integer scores, or assign other values if integer spacing is not appropriate.

```
> Csore <- as.numeric(Mental$ses)
> Rsore <- as.numeric(Mental$mental)
```

Then, the $L \times L$, R and C models can be fit as follows, where beyond the main effects of mental and ses, their association is represented as the interaction of the numeric score(s) or factor(s), as appropriate in each case.

```
> linlin <- glm(Freq ~ mental + ses + Rsore:Csore,
+                      family = poisson, data = Mental)
> roweff <- glm(Freq ~ mental + ses + mental:Csore,
+                      family = poisson, data = Mental)
> coleff <- glm(Freq ~ mental + ses + Rsore:ses,
+                      family = poisson, data = Mental)
```

Goodness-of-fit tests for these models are shown below. They show that all of the $L \times L$, R and C models are acceptable in terms of the likelihood-ratio G^2 . The $L \times L$ model, with only one more parameter than the independence model is judged the best by both AIC and BIC.

```
> vcdExtra:::LRstats(indep, linlin, roweff, coleff)

Likelihood summary table:
      AIC   BIC  LR Chisq Df Pr(>Chisq)
indep 209.6 220.2    47.42 15   3.16e-05 ***
linlin 174.1 185.8     9.90 14    0.770
roweff 174.4 188.6     6.28 12    0.901
coleff 179.0 195.5     6.83 10    0.741
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

In cases where such overall tests are unclear, you can carry out tests of nested sets of models using **anova()**, giving tests of ΔG^2 .

```
> anova(indep, linlin, roweff, test="Chisq")

Analysis of Deviance Table

Model 1: Freq ~ mental + ses
Model 2: Freq ~ mental + ses + Rsore:Csore
Model 3: Freq ~ mental + ses + mental:Csore
  Resid. Df Resid. Dev Df Deviance Pr(>Chi)
1       15      47.4
2       14      9.9  1     37.5    9e-10 ***
3       12      6.3  2     3.6     0.16
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

> anova(indep, linlin, coleff, test="Chisq")

Analysis of Deviance Table

Model 1: Freq ~ mental + ses
Model 2: Freq ~ mental + ses + Rsore:Csore
Model 3: Freq ~ mental + ses + Rsore:ses
  Resid. Df Resid. Dev Df Deviance Pr(>Chi)
1       15      47.4
2       14      9.9  1     37.5    9e-10 ***
3       10      6.8  4     3.1     0.55
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Under the $L \times L$ model, the estimate of the coefficient of **Rsore:Csore** is $\hat{\gamma} = 0.0907$ (s.e.=0.015) with unit-spaced scores, as shown below.

```
> # interpret linlin association parameter
> coef(linlin)[["Rscore:Cscore"]]

[1] 0.090687

> exp(coef(linlin)[["Rscore:Cscore"]])

[1] 1.0949
```

This corresponds to a local odds ratio, $\hat{\theta}_{ij} = \exp(0.0907) = 1.095$. This single number describes the association succinctly: each step down the socioeconomic scale increases the odds of being classified one step poorer in mental health by 9.5%.



8.6.2 Log-multiplicative (RC) models

{sec:RCmodels}

The association models described above are all more parsimonious and easier to interpret than the saturated model. However, they depend on assigning fixed and possibly arbitrary scores to the variable categories. A generalization of the $L \times L$ model that treats *both* row and column scores as parameters is the **row-and-column effects model** (RC(1)) suggested by Goodman (1979),

$$\log(m_{ij}) = \mu + \lambda_i^A + \lambda_j^B + \gamma \alpha_i \beta_j , \quad (8.22) \quad \text{(eq:RC1)}$$

where γ , α and β comprise additional parameters to be estimated beyond the independence model.⁸ This model has a close connection with correspondence analysis (Goodman, 1985), where the estimated scores α and β are analogous to correspondence analysis scores on a first dimension.⁹ γ , called the *intrinsic association coefficient* is analogous to the same parameter in the $L \times L$ model.

For identifiability and interpretation it is necessary to impose some normalization constraints on the α and β . An *unweighted, unit standardized* solution forces $\sum_i \alpha_i = \sum_j \beta_j = 0$ and $\sum_i \alpha_i^2 = \sum_j \beta_j^2 = 1$. Alternatively, and more akin to correspondence analysis solutions, the *marginally weighted* solution uses the marginal probabilities π_{i+} of the row variable and π_{+j} of the columns as weights.

$$\begin{aligned} \sum_i \alpha_i \pi_{i+} &= \sum_j \beta_j \pi_{+j} = 0 \\ \sum_i \alpha_i^2 \pi_{i+} &= \sum_j \beta_j^2 \pi_{+j} = 1 \end{aligned} \quad (8.23) \quad \text{(eq:RC-constraints)}$$

Goodman (1986) generalized this to multiple bilinear terms of the form $\gamma_k \alpha_{ik} \beta_{jk}$, with M terms (the RC(M) model) and showed that *all* associations in the saturated model could be expressed exactly as

$$\lambda_{ij}^{AB} = \sum_{k=1}^M \gamma_k \alpha_{ik} \beta_{jk} \quad M = \min(I - 1, J - 1) . \quad (8.24) \quad \text{(eq:RCm)}$$

In practice, models with fewer terms usually suffice. For example, an RC(2) model with two multiplicative terms is analogous to a two-dimensional correspondence analysis solution. In addition to the normalization constraints for the RC(1) model, parameters in an RC(M) model must satisfy

⁸In contrast to the R, C and R+C models, RC models do not assume that the categories are appropriately ordered because the category scores are estimated from the data.

⁹However, when estimated by maximum likelihood, the RC(1) model allows likelihood-ratio tests of parameters and model fit, AIC and BIC statistics, and methods for estimating standard errors of the parameters. Such model-based methods are not available for correspondence analysis.

the additional constraints that the (possibly weighted) scores for distinct dimensions are orthogonal (uncorrelated), similar to correspondence analysis solutions.

The RC model is *not* a loglinear model because it contains a multiplicative term in the parameters. This model and a wide variety of other nonlinear models for categorical data can be fit using `gnm()` in the `gnm` package. This provides the basic machinery for extending `glm()` models to nonlinear terms, quite generally. The function `rc()` in the `logmult` package uses `gnm()` for fitting, and offers greater convenience in normalizing the category scores, calculating standard errors and plotting.

{ex:mental5}

EXAMPLE 8.6: Mental impairment and parents' SES

The `gnm` package provides a number of functions that can be used in model formulas for nonlinear association terms. Among these, `Mult()` expresses a multiplicative association in terms of two (or more) factors. The RC(1) model for factors A, B uses `Mult(A, B)` for the association term in Eqn. (8.22). Multiple multiplicative RC terms, as in Eqn. (8.24) can be expressed using `instances(Mult(A, B), m)`.

To illustrate, we fit the RC(1) and RC(2) models to the `Mental` data using `gnm()`. In this table, both factors are ordered, but we don't want to use the default polynomial contrasts, so we set their contrast attributes to treatment.

```
> library(gnm)
> Mental$mental <- C(Mental$mental, treatment)
> Mental$ses <- C(Mental$ses, treatment)
> RC1 <- gnm(Freq ~ mental + ses + Mult(mental, ses),
+               family = poisson, data = Mental, verbose=FALSE)
> RC2 <- gnm(Freq ~ mental + ses + instances(Mult(mental, ses), 2),
+               family = poisson, data = Mental, verbose=FALSE)
```

For comparison with the loglinear association models fit in Example 8.5 we show the G^2 goodness of fit tests for all these models. The ordinal loglinear models and the RC models all fit well, with the $L \times L$ model preferred on the basis of parsimony by AIC and BIC.

```
> vcdExtra::LRstats(indep, linlin, roweff, colegg, RC1, RC2)

Likelihood summary table:
      AIC    BIC   LR Chisq Df Pr(>Chisq)
indep 209.6 220.2   47.42 15   3.16e-05 ***
linlin 174.1 185.8   9.90 14     0.770
roweff 174.4 188.6   6.28 12     0.901
colegg 179.0 195.5   6.83 10     0.741
RC1    179.7 198.6   3.57  8     0.894
RC2    186.7 211.4   0.52  3     0.914
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The substantive difference between the $L \times L$ model and the RC(1) model is whether the categories of mental health status and SES can be interpreted as equally spaced along some latent continua, versus the alternative that category spacing is unequal. We can test this directly using the likelihood-ratio test, $G^2(L \times L | RC(1))$. Similarly, model `RC1` is nested within model `RC2`, so $G^2(RC(1) | RC(2))$ gives a direct test of the need for a second dimension.

```
> anova(linlin, RC1, RC2, test="Chisq")

Analysis of Deviance Table

Model 1: Freq ~ mental + ses + Rscore:Cscore
Model 2: Freq ~ mental + ses + Mult(mental, ses)
Model 3: Freq ~ mental + ses + Mult(mental, ses, inst = 1) + Mult(mental,
```

	ses, inst = 2)	Resid. Df	Resid. Dev	Df	Deviance	Pr(>Chi)
1		14	9.90			
2		8	3.57	6	6.32	0.39
3		3	0.52	5	3.05	0.69

We see that estimated scores for the categories in the model RC1 do not provide a significantly better fit, and there is even less evidence for a second dimension of category parameters in the RC2 model.

Nevertheless, for cases where RC models *do* provide some advantage, it is useful to know how to visualize the estimated category parameters. The key to this is the function `getContrasts()` which computes contrasts or scaled contrasts for a set of (non-eliminated) parameters from a "gnm" model, together with standard errors for the estimated contrasts following the methods of Firth (2003), Firth and Menezes (2004). The details are explained in `help(getContrasts)` and in `vignette("gnmOverview")` that comes with the `gnm` package.

The coefficients in the marginally-weighted solution Eqn. (8.23) can be obtained as follows.

```
> rowProbs <- with(Mental, tapply(Freq, mental, sum) / sum(Freq))
> colProbs <- with(Mental, tapply(Freq, ses, sum) / sum(Freq))
> mu <- getContrasts(RC1, pickCoef(RC1, "[.]mental"),
+                      ref = rowProbs, scaleWeights = rowProbs)
> nu <- getContrasts(RC1, pickCoef(RC1, "[.]ses"),
+                      ref = colProbs, scaleWeights = colProbs)
```

In our notation, the coefficients α and β can be extracted as the `qvframe` component of the `"qv"` object returned by `getContrasts()`.

```
> (alpha <- mu$qvframe)

Estimate Std. Error
Mult(., ses).mentalWell      1.67378   0.19043
Mult(., ses).mentalMild      0.14009   0.20018
Mult(., ses).mentalModerate -0.13669   0.27948
Mult(., ses).mentalImpaired -1.41055   0.17418

> (beta <- nu$qvframe)

Estimate Std. Error
Mult(mental, .).ses1    1.111361   0.29921
Mult(mental, .).ses2    1.120459   0.31422
Mult(mental, .).ses3    0.370752   0.31915
Mult(mental, .).ses4   -0.027006   0.27328
Mult(mental, .).ses5   -1.009480   0.31470
Mult(mental, .).ses6   -1.816647   0.28095
```

For plotting this RC(1) solution for the scaled category scores together with their estimated standard errors, a `dotchart()`, shown in Figure 8.9 provides a reasonable visualization.

To create this plot, first combine the row and column scores in a data frame, and add columns `lower`, `upper` corresponding to ± 1 standard error (or some other multiple).

```
> scores <- rbind(alpha, beta)
> scores <- cbind(scores,
+                   factor=c(rep("mental", 4), rep("ses", 6)))
> rownames(scores) <- c(levels(Mental$mental), levels(Mental$ses))
> scores$lower <- scores[,1]-scores[,2]
> scores$upper <- scores[,1]+scores[,2]
> scores
```

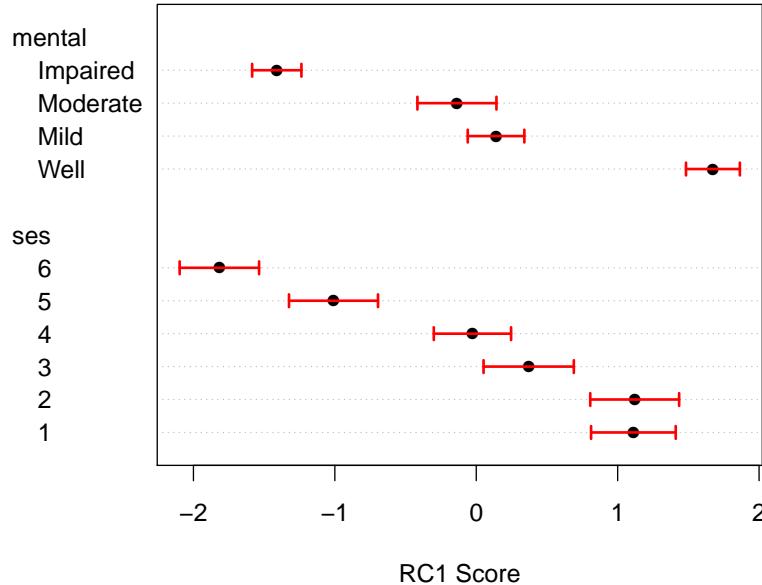


Figure 8.9: Dotchart of the scaled category scores for the RC(1) model fit the mental health data. Error bars show ± 1 standard error.

{fig:mental-RC1}

	Estimate	Std.	Error	factor	lower	upper
Well	1.674	0.190	mental	1.4834	1.864	
Mild	0.140	0.200	mental	-0.0601	0.340	
Moderate	-0.137	0.279	mental	-0.4162	0.143	
Impaired	-1.411	0.174	mental	-1.5847	-1.236	
1	1.111	0.299	ses	0.8121	1.411	
2	1.120	0.314	ses	0.8062	1.435	
3	0.371	0.319	ses	0.0516	0.690	
4	-0.027	0.273	ses	-0.3003	0.246	
5	-1.009	0.315	ses	-1.3242	-0.695	
6	-1.817	0.281	ses	-2.0976	-1.536	

The dotchart shown in Figure 8.9 is then a plot of Estimate, grouped by factor, with arrows showing the range of lower to upper for each parameter.

```
> with(scores, {
+   dotchart(Estimate, groups=factor, labels=rownames(scores),
+             cex=1.2, pch=16, xlab="RC1 Score",
+             xlim=c(min(lower), max.upper)))
+   arrows(lower, c(8+(1:4), 1:6), upper, c(8+(1:4), 1:6),
+          col="red", angle=90, length=.05, code=3, lwd=2)
+ })
```

In this plot, the main substantive difference from the $L \times L$ model is in the spacing of the lowest two categories of ses and the middle two categories of mental which are not seen to differ in the RC1 model.

The coefficients in the RC2 model can also be plotted (in a 2D plot) by extracting the coefficients from the "gnm" object and reshaping them to 2-column matrices. The function `pickCoef()` is handy here to get the indices of a subset of parameters by matching a pattern in their names. **TODO:** Maybe delete some of this, in favor of using `logmult`.

```

> alpha <- coef(RC2) [pickCoef(RC2, "[.]mental")]
> alpha <- matrix(alpha, ncol=2)
> rownames(alpha) <- levels(Mental$mental)
> colnames(alpha) <- c("Dim1", "Dim2")
> alpha

      Dim1      Dim2
Well     -0.747969  0.084693
Mild     -0.023971 -0.057400
Moderate 0.345882 -0.464567
Impaired 0.562678  0.017106

> beta <- coef(RC2) [pickCoef(RC2, "[.]ses")]
> beta <- matrix(beta, ncol=2)
> rownames(beta) <- levels(Mental$ses)
> colnames(beta) <- c("Dim1", "Dim2")
> beta

      Dim1      Dim2
1 -0.465521 -0.36373
2 -0.477669 -0.43232
3 -0.159825 -0.16501
4  0.031569  0.33016
5  0.414625  0.51511
6  0.753808  0.16155

```

The simple, unweighted scaling to mean 0, variance 1 can be obtained with **scale()**:

```

> alpha <- scale(alpha)
> beta <- scale(beta)

```

Alternatively, the marginal-weighted scaling of Eqn. (8.23) is obtained by centering at the weighted mean and dividing by the weighted sum of squares. We use this scaling here.

```

> alpha <- apply(alpha, 2, function(x) x - sum(x*rowProbs))
> alpha <- apply(alpha, 2, function(x) x/sqrt(sum(x^2 * rowProbs)))
> beta <- apply(beta, 2, function(x) x - sum(x*colProbs))
> beta <- apply(beta, 2, function(x) x/sqrt(sum(x^2 * colProbs)))

```

To plot these category scores, first combine them into a single data frame,

```

> scores <- data.frame(rbind(alpha,beta))
> scores$factor <- c(rep("mental", 4), rep("ses", 6))
> scores$probs <- c(rowProbs, colProbs)
> scores

      Dim1      Dim2 factor probs
Well     -1.8058  0.946 mental 0.185
Mild     -0.1882  0.228 mental 0.363
Moderate 0.6381 -1.830 mental 0.218
Impaired 1.1225  0.604 mental 0.234
1       -1.1162 -1.121   ses 0.158
2       -1.1453 -1.317   ses 0.148
3       -0.3839 -0.555   ses 0.173
4        0.0746  0.855   ses 0.231
5        0.9923  1.382   ses 0.160
6       1.8048  0.375   ses 0.131

```

Then, we use **xyplot()** to plot the scores on Dim2 against Dim1, with separate lines and colors for the two factors. The resulting plot is shown in Figure 8.10.

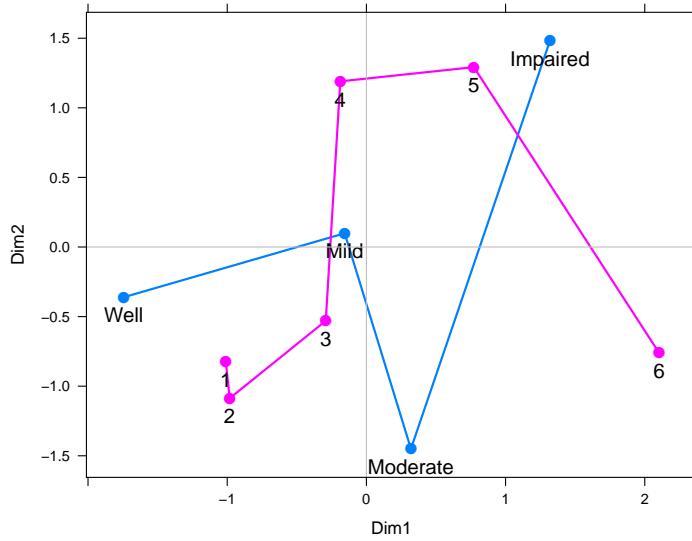


Figure 8.10: Scaled category scores for the RC(2) model fit the mental health data.

{fig:mental-RC2}

```
> library(lattice)
> xyplot(Dim2 ~ Dim1, groups=factor, data=scores, type="b",
+         cex=1.3, pch=16, lwd=2, aspect="iso",
+         panel=function(x, y, ...) {
+           panel.xyplot(x, y, ...)
+           panel.text(x=x, y=y, labels=rownames(scores), pos=1, cex=1.2)
+           panel.abline(h=0, col="gray")
+           panel.abline(v=0, col="gray")
+         }
+       )
```

The patterns of the row and column category scores here are quite similar to the 2D correspondence analysis solution shown in Figure 6.2. The main difference is in the relative scaling of the axes. In Figure 8.10, the variances of the two dimensions are equated; in the correspondence analysis plot, the axes are scaled in relation to their contributions to Pearson χ^2 , allowing an interpretation of distance between points in terms of χ^2 -distance.



8.6.2.1 Using logmult

From the previous example, you can see that it takes a fair bit of work to extract the coefficients from "gnm" objects and carry out the scaling necessary for informative plots. Much of this effort is now performed by the **logmult** package with several convenience functions that do the heavy lifting.

`rc()` fits the class of RC(M) models, allowing an argument `nd` to specify the number of dimensions, and also providing for standard errors estimated using jackknife and bootstrap methods (Milan and Whittaker, 1995), which are computationally intensive. For square tables, a `symmetric` argument constrains the row and column scores to be equal, and a `diagonal` option fits parameters for each diagonal cell, providing for models of quasi-independence and quasi-symmetry (see Section 8.7).

It returns an object of class "rc" with the components of the "gnm" object. An `assoc` component is also returned, containing the normalized association parameters for the categories.

`rcl()` fits extensions of RC models to tables with multiple layers, called RC(M)-L models by Wong (2010).

`plot.rc()` is a plot method for visualizing scores for RC(M) models in two selected dimensions.

Among other options, it can plot confidence ellipses for the category scores, using the estimated covariance matrix (assuming a normal distribution of the category scores). The plot method returns (invisibly) the coordinates of the scores as plotted, facilitating additional plot annotation.

{ex:mental6}

EXAMPLE 8.7: Mental impairment and parents' SES

Here we use `rc()` to estimate the RC(1) and RC(2) models for the `Mental` data. In contrast to `gnm()`, which has a formula interface for a `data` argument, `rc()` requires the input in the form of a two-way table, given here as `mental.tab`.

```
> library(logmult)
> rcl <- rc(mental.tab, verbose=FALSE, weighting="marginal",
+            se="jackknife")
> rc2 <- rc(mental.tab, verbose=FALSE, weighting="marginal", nd=2,
+            se="jackknife")
```

The option `weighting="marginal"` gives the marginally-weighted solution and `se="jackknife"` estimates the covariance matrix using the leave-one-out jackknife.¹⁰

A plot of the scaled category scores similar to Figure 8.10, with 1 standard error confidence ellipses (making them comparable to the 1D solution shown in Figure 8.9) but no connecting lines can then be easily produced with the `plot()` method for "rc" objects.

```
> coords <- plot(rc2, conf.ellipses=0.68, cex=1.5, rev.axes=c(TRUE, FALSE))
```

The orientation of the axes is arbitrary in RC(M) models, so the horizontal axis is reversed here to conform with Figure 8.10.

This produces (in Figure 8.11) a symmetric biplot in which the scaled coordinates of points for rows (α_{ik}) and columns (β_{jk}) on both axes are the product of normalized scores and the square root of the intrinsic association coefficient (γ_k) corresponding to each dimension.

Such plots can be customized using the category coordinates (`coords`) returned by the `plot()` method. As in other biplots, joining the row and column points by lines (sorted by the first dimension) makes it easier to see their relationships across the two dimensions. The following code draws the lines shown in Figure 8.11.

```
> scores <- rbind(coords$row, coords$col)
> lines(scores[1:4,], col="blue", lwd=2)
> lines(scores[-(1:4),], col="red", lwd=2)
```

We saw earlier that there was not strong evidence supporting the need for a second RC dimension to describe the relationship between mental health and SES. This is apparent in the sizes of the confidence ellipses, which overlap much more along Dimension 2 than Dimension 1. \triangle

¹⁰Becker and Clogg (1989) recommend using unweighted solutions, `weighting="none"` (they call them "uniformly weighted") to preserve independence of inferences about association and marginal effects and estimates of the intrinsic association parameters, γ_k . That choice makes very little difference in the plots for this example, but the γ_k parameters are affected considerably.

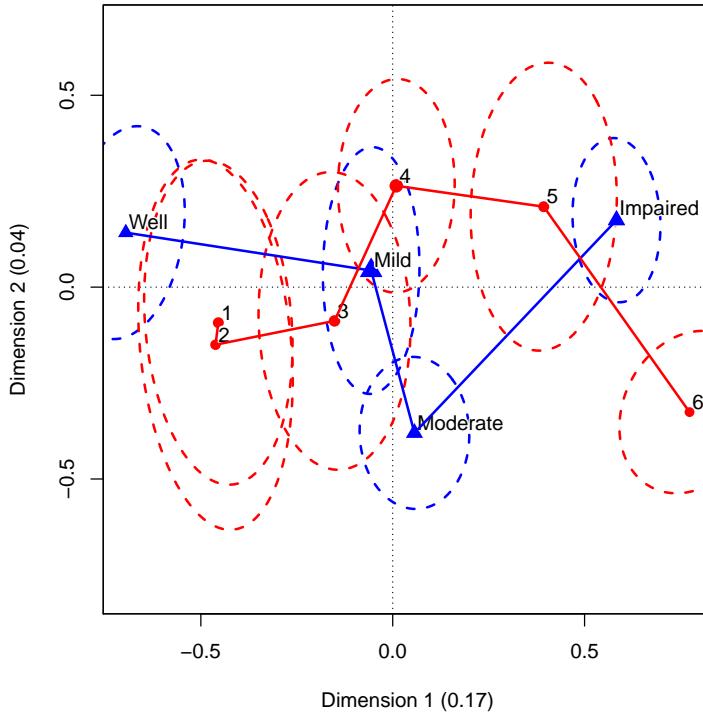


Figure 8.11: Scaled category scores for the RC(2) model fit and plotted using the `logmult` package. The 68% confidence ellipses correspond to bivariate ± 1 confidence intervals for the category parameters.

{fig:mental-logmult-rc}

8.7 Square tables

Square tables, where the row and column variables have the same categories comprise an important special case for loglinear models that can account for associations more parsimoniously than the saturated model. Some examples are the data on visual acuity in Example 4.13, categorical ratings of therapy clients by two observers, and mobility tables, tracking the occupational categories between generations in the same families or migration tables, giving movement of people between regions. The latter topics has been important in sociological and geographic research and has spurred the development of a wide range of specialized loglinear models for this purpose.

{sec:loglin-square}

8.7.1 Quasi-independence, symmetry, quasi-symmetry and topological models

{sec:sq-quasi}

In many square tables, such as the `Vision` data, independence is not a credible hypothesis because the diagonal cells, representing equal values of the row and column variables tend to be very large and often contribute most of the lack of fit. A substantively more interesting hypothesis is whether the table exhibits independence, ignoring the diagonal cells. This leads to what is called the **quasi-independence model**, that specifies independence only in the off-diagonal cells.

For a two-way table, quasi-independence can be expressed as

$$\pi_{ij} = \pi_{i+}\pi_{+j} \quad \text{for } i \neq j$$

or in loglinear form as

$$\log m_{ij} = \mu + \lambda_i^A + \lambda_j^B + \delta_i I(i = j) .$$

This model effectively adds one parameter, δ_i , for each main diagonal cell which fits those frequencies perfectly.

Another hypothesis of substantive interest for square tables, particularly those concerning occupational and geographical mobility is that the joint distribution of row and column variables is symmetric, that is, $\pi_{ij} = \pi_{ji}$ for all $i \neq j$. For example, this **symmetry model** (S) asserts that sons are as likely to move from their father's occupation i to another, j , as the reverse. This form of symmetry is quite strong, because it also implies **marginal homogeneity** (MH), that the marginal probabilities of the row and column variables are equal, $\pi_{i+} = \sum_j \pi_{ij} = \sum_j \pi_{ji} = \pi_{+i}$ for all i .

To separate marginal homogeneity from symmetry of the association terms per se, the model of **quasi-symmetry** (QS) uses the standard main-effect terms in the loglinear model,

$$\log m_{ij} = \mu + \lambda_i^A + \lambda_j^B + \lambda_{ij} , \quad (8.25) \quad \{eq:quasi-symm\}$$

where $\lambda_{ij} = \lambda_{ji}$. It can be shown (Caussinus, 1966) that

$$\begin{aligned} \text{symmetry} &= \text{quasi-symmetry} + \text{marginal homogeneity} \\ G^2(S) &= G^2(QS) + G^2(MH) \end{aligned}$$

where $G^2(MH)$ is defined by the likelihood-ratio test of the difference between the S and QS models,

$$G^2(MH) \equiv G^2(S | QS) = G^2(S) - G^2(QS) . \quad (8.26) \quad \{eq:mh\}$$

The **gnm** package provides several model building convenience functions that facilitate fitting these and related models:

- `Diag(row, col, ...)` constructs a diagonals association factor for two (or more) factors with integer levels where the original factors are equal, and " ." otherwise.
- `Symm(row, col, ...)` constructs an association factor giving equal levels to sets of symmetric cells. The QS model is specified using `Diag() + Symm()`.
- `Topo(row, col, ..., spec)` creates an association factor for two or more factors, as specified by an array of levels, which may be arbitrarily structured. Both `Diag()` and `Symm()` factors are special cases of `Topo()`.

The factor levels representing these association effects for a 4×4 table are shown below by their unique values in each array.

$$\text{Diag}_{4 \times 4} = \begin{bmatrix} 1 & . & . & . \\ . & 2 & . & . \\ . & . & 3 & . \\ . & . & . & 4 \end{bmatrix} \quad \text{Symm}_{4 \times 4} = \begin{bmatrix} 11 & 12 & 13 & 14 \\ 12 & 22 & 23 & 24 \\ 13 & 23 & 33 & 34 \\ 14 & 24 & 34 & 44 \end{bmatrix} \quad \text{Topo}_{4 \times 4} = \begin{bmatrix} 2 & 3 & 4 & 4 \\ 3 & 3 & 4 & 4 \\ 4 & 4 & 5 & 5 \\ 4 & 4 & 5 & 1 \end{bmatrix}$$

{ex:vision-glm}

EXAMPLE 8.8: Visual acuity

Example 4.13 presented the data on tests of visual acuity in the left and right eyes of a large sample of women working in the Royal Ordnance factories in World War II. A sieve diagram (Figure 4.8) showed that, as expected, most women had the same acuity in both eyes, but the off-diagonal cells had a pattern suggesting some form of symmetry.

The data set `VisualAcuity` contains data for both men and women in frequency form and for this example we subset this to include only the 4×4 table for women.

```
> data("VisualAcuity", package="vcd")
> women <- subset(VisualAcuity, gender=="female", select=-gender)
```

The four basic models of independence, quasi-independence, symmetry and quasi-symmetry for square tables are fit as shown below. We use `update()` to highlight the relations among these models in two pairs.

```
> #library(vcdExtra)
> indep <- glm(Freq ~ right + left, data = women, family = poisson)
> quasi <- update(indep, . ~ . + Diag(right, left))
>
> symm <- glm(Freq ~ Symm(right, left), data = women, family = poisson)
> qsymm <- update(symm, . ~ right + left + .)
```

The brief summary of goodness of fit of these models below shows that the QS model fits reasonably well, but none of the others do by likelihood-ratio tests or AIC or BIC.

```
> vcdExtra::LRstats(indep, quasi, symm, qsymm)

Likelihood summary table:
  AIC  BIC LR Chisq Df Pr(>Chisq)
indep 6803 6808     6672  9      <2e-16 ***
quasi  338   347      199  5      <2e-16 ***
symm   157   164      19   6      0.0038 **
qsymm  151   161       7   3      0.0638 .
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Beyond just saying that the QS model fits best, the reasons *why* it does can be seen in mosaic displays. Figure 8.12 compares the mosaics for the models of quasi-independence (accounting only for the diagonal cells) and quasi-symmetry (also accounting for symmetry). It can be seen in the left panel that the non-diagonal associations are largely symmetric, and also that when they differ, visual acuity in the two eyes are most likely to differ by only one eye grade.

```
> labs <- c("High", "2", "3", "Low")
> largs <- list(set_varnames = c(right="Right eye grade",
+                                    left="Left eye grade"),
+                  set_labels=list(right=labs, left=labs))
> mosaic(quasi, ~right + left, residuals_type="rstandard",
+          gp=shading_Friendly,
+          labeling_args=largs,
+          main="Quasi-Independence (women)")
> mosaic(qsymm, ~right + left, residuals_type="rstandard",
+          gp=shading_Friendly,
+          labeling_args=largs,
+          main="Quasi-Symmetry (women)")
```

Finally, as usual, `anova()` can be used to carry out specific tests of nested models. For example, the test of marginal homogeneity Eqn. (8.26) compares models S and QS and shows here that the marginal probabilities for the left and right eyes differ.

```
> anova(symm, qsymm, test="Chisq")

Analysis of Deviance Table

Model 1: Freq ~ Symm(right, left)
Model 2: Freq ~ right + left + Symm(right, left)
  Resid. Df Resid. Dev Df Deviance Pr(>Chi)
1           6    19.25
2           3     7.27  3        12    0.0075 **
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

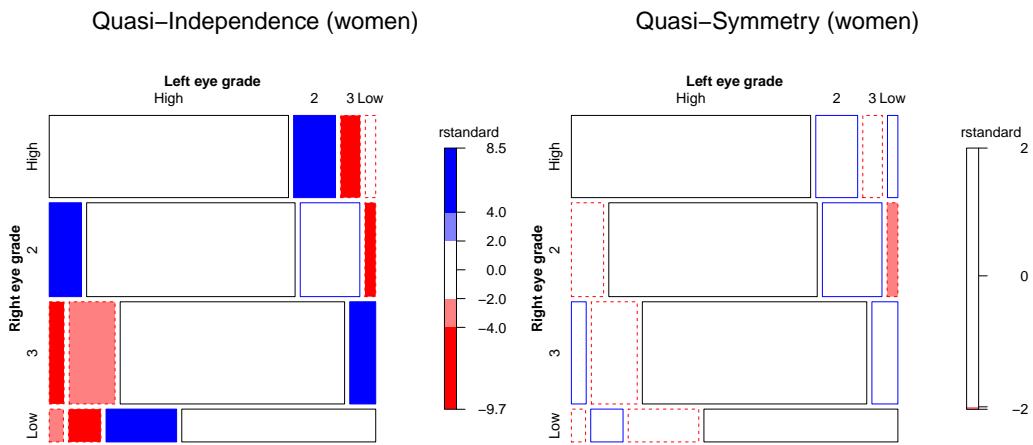


Figure 8.12: Mosaic displays comparing the models of quasi-independence and quasi-symmetry for visual acuity in women.

△
{ex:hauser1}

EXAMPLE 8.9: Hauser's occupational mobility table

The data `Hauser79` in `vcdExtra`, from Hauser (1979), gives a 5×5 table in frequency form cross-classifying 19,912 individuals in the United States by father's occupation and son's first occupation. The occupational categories are represented by abbreviations, of Upper Non-Manual (`UpNM`), Lower Non-Manual (`LoNM`), Upper Manual (`UpM`), Lower Manual (`LoM`) and Farm. These data were also analysed by Powers and Xie (2008).

```
> data("Hauser79", package="vcdExtra")
> structable(~Father+Son, data=Hauser79)
```

	Son	UpNM	LoNM	UpM	LoM	Farm
Father						
UpNM	1414	521	302	643	40	
LoNM	724	524	254	703	48	
UpM	798	648	856	1676	108	
LoM	756	914	771	3325	237	
Farm	409	357	441	1611	1832	

Before fitting any models, it is useful to calculate and plot the observed local log odds ratios, as we did in Example 8.5 to see the patterns in the data that need to be accounted for. These are calculated using `loddsratio()`.

```
> hauser.tab <- xtabs(Freq ~ Father+Son, data=Hauser79)
> (lor.hauser <- loddsratio(hauser.tab))

log odds ratios for Father and Son

          Son
Father      UpNM:LoNM LoNM:UpM UpM:LoM LoM:Farm
  UpNM:LoNM    0.67513 -0.17883  0.26230  0.093109
  LoNM:UpM     0.11508  1.00254 -0.34613 -0.057878
  UpM:LoM      0.39801 -0.44852  0.78964  0.100869
  LoM:Farm     -0.32577  0.38145 -0.16597  2.769718
```

This 4×4 table is graphed using `matplot()`, giving Figure 8.13.

```
> matplot(as.matrix(lor.hauser), type='b', lwd=2,
+         ylab='Local log odds ratio',
+         xlab="Son's status comparisons",
+         xaxt='n', cex.lab=1.2,
+         xlim=c(1,4.5), ylim=c(-.5, 3)
+       )
> abline(h=0, col='gray') # independence
> abline(h=mean(lor.hauser$coefficients)) # mean
> axis(side=1, at=1:4, labels=colnames(lor.hauser))
> text(4, as.matrix(lor.hauser)[4,], rownames(lor.hauser),
+       pos=4, col=1:4, xpd=TRUE, cex=1.2)
> text(4, 3, "Father's status", cex=1.2)
```

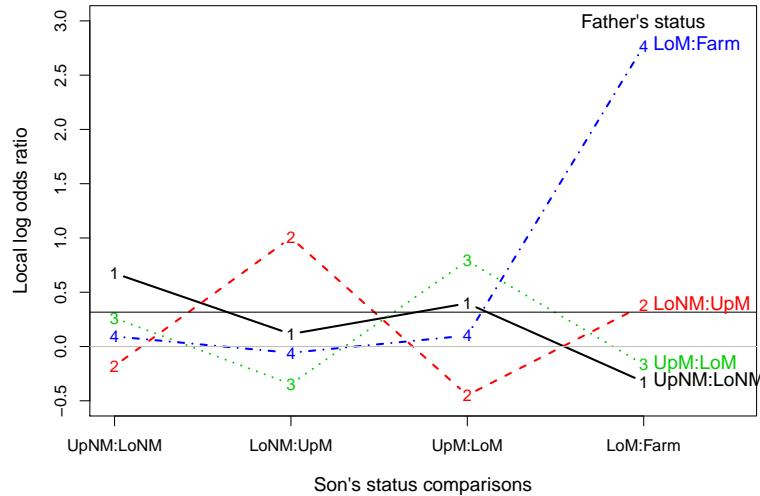


Figure 8.13: Plot of observed local log odds ratios in the Hauser79 data. The gray horizontal line at zero shows local independence; the black horizontal line shows the mean.
fig.hauser-lor-plot

Amongst the features here, you can see that there is a tendency for the odds ratio contrasting fathers in the non-manual categories (`UpNM:LoNM`) to decline with the adjacent comparisons of their sons' occupations. As well, the 2×2 table for fathers and sons in the `LoM:Farm` stands out as deserving some attention. These observed features will be smoothed by fitting models, as described below. For additional interpretation, you can always construct similar plots of the log odds ratios using the `fitted()` values from any of the models described below.

We begin by fitting the independence model and the quasi-independence model, where the diagonal parameters in the latter are specified as `Diag(Father, Son)`. As expected, given the large frequencies in the diagonal cells, the quasi-independence model is a considerable improvement, but the fit is still very poor.

```
> hauser.indep <- gnm(Freq ~ Father + Son, data=Hauser79, family=poisson)
> hauser.quasi <- update(hauser.indep, ~ . + Diag(Father, Son))
> vcdExtra::LRstats(hauser.indep, hauser.quasi)

Likelihood summary table:
      AIC    BIC   LR Chisq Df Pr(>Chisq)
```

```
hauser.indep 6391 6402      6170 16      <2e-16 ***
hauser.quasi   914   931      683 11      <2e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The pattern of associations can be seen in the mosaic displays for both models, shown in Figure 8.14.

```
> mosaic(hauser.indep, ~Father+Son, main="Independence model",
+         gp=shading_Friendly)
> mosaic(hauser.quasi, ~Father+Son, main="Quasi-independence model",
+         gp=shading_Friendly)
```

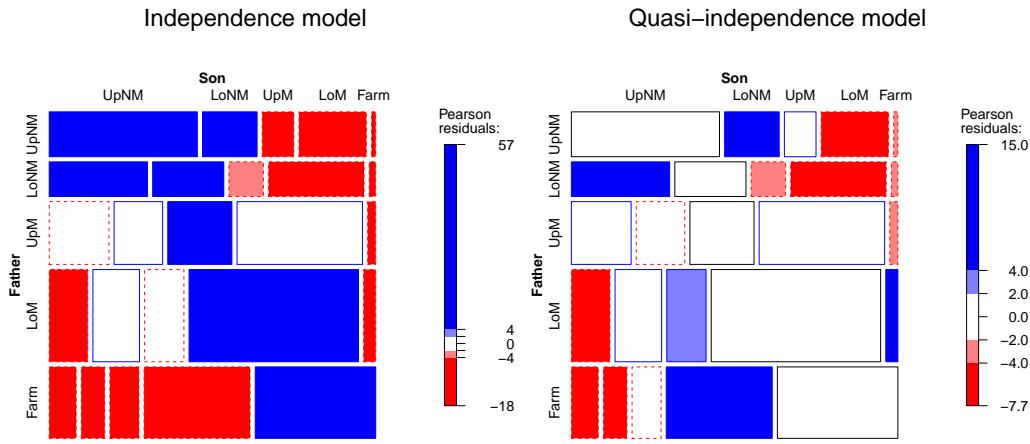


Figure 8.14: Mosaic displays for the Hauser79 data. Left: independence model; right: quasi-independence model.
fig:hauser-mosaici

The mosaic for quasi-independence shows an approximately symmetric pattern of residuals, so we proceed to add `Symm(Father, Son)` to the model to specify quasi-symmetry.

```
> hauser.qsymm <- update(hauser.indep,
+                         ~ . + Diag(Father,Son) + Symm(Father,Son))
> vcdExtra::LRstats(hauser.qsymm)

Likelihood summary table:
      AIC BIC LR Chisq Df Pr(>Chisq)
hauser.qsymm 268 291     27.4 6    0.00012 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

This model represents a huge improvement in goodness of fit. With such a large sample size, it might be considered an acceptable fit. The remaining lack of fit is shown in the mosaic for this model, Figure 8.15.

```
> mosaic(hauser.qsymm, ~Father+Son, main="Quasi-symmetry model",
+         gp=shading_Friendly, residuals_type="rstandard")
```

The cells with the largest lack of symmetry (using standardized residuals) are those for the upper

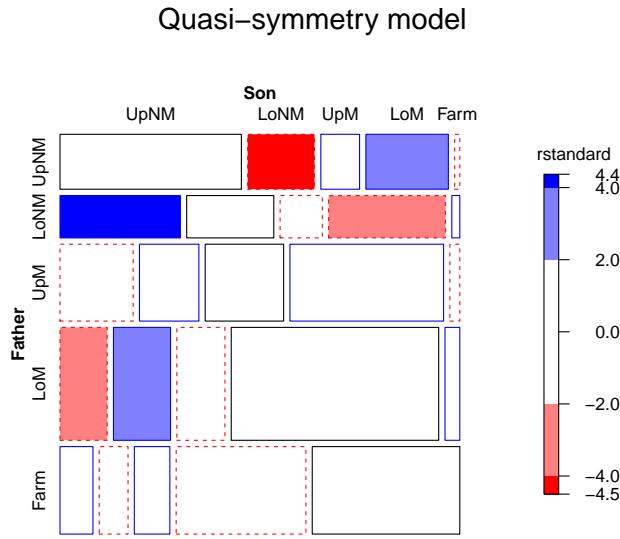


Figure 8.15: Mosaic display for the model of quasi-symmetry fit to the Hauser79 data.^{fig:hauser-mosaic2}

and lower non-manual occupations, where the son of an upper manual worker is less likely to move to lower non-manual work than the reverse.

For cases like this involving structured associations in square tables, Hauser (1979) developed the more general idea of grouping the row and column categories into levels of an association factor based on similar values of residuals or local odds ratios observed from the independence model. Such models are called *topological models* or *levels models*, which are implemented in the `Topo()`.

To illustrate, Hauser suggested the following matrix of levels to account for the pattern of associations seen in Figure 8.14. The coding here takes the diagonal cell for the Farm category as the reference cell. Four other parameters are assigned by the numbers 2–5 to account for lack of independence.

```
> levels <- matrix(c(
+   2, 4, 5, 5, 5,
+   3, 4, 5, 5, 5,
+   5, 5, 5, 5, 5,
+   5, 5, 5, 4, 4,
+   5, 5, 5, 4, 1
+ ), 5, 5, byrow=TRUE)
```

This model is fit using `Topo()` as shown below. It also provides a huge improvement over the independence model, with 4 additional parameters.

```
> hauser.topo <- update(hauser.indep, ~ . + Topo(Father, Son, spec=levels))
> vcdExtra::LRstats(hauser.topo)

Likelihood summary table:
      AIC BIC LR Chisq Df Pr(>Chisq)
hauser.topo 295 311    66.6 12    1.4e-09 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

As with other models fit using `gnm()`, you can extract the coefficients for particular terms using `pickCoef()`.

```
> as.vector( (coef(hauser.topo) [pickCoef(hauser.topo, "Topo")]))  
[1] -1.8128 -2.4973 -2.8035 -3.4026
```

The models fit in this example are summarized below. Note that AIC prefers the quasi-symmetry model, `hauser.quasi`, while, because of the large sample size, BIC prefers the topological model, `hauser.topo`.

```
> vcdExtra::LRstats(hauser.indep, hauser.quasi, hauser.qsymm, hauser.topo)  
  
Likelihood summary table:  
      AIC   BIC  LR Chisq Df Pr(>Chisq)  
hauser.indep 6391 6402     6170 16    < 2e-16 ***  
hauser.quasi  914  931     683 11    < 2e-16 ***  
hauser.qsymm  268  291     27  6    0.00012 ***  
hauser.topo   295  311     67 12    1.4e-09 ***  
---  
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

△

8.7.2 Ordinal square tables

The theory presented in Section 8.7.1 treats the row and column variables as nominal. In many instances, such as Example 8.9, the variable categories are also ordered, yet these models do not exploit their ordinal nature. In such cases, the models such as uniform association ($L \times L$), row effects, RC and others discussed in Section 8.6 can be combined with terms for quasi-independence and symmetry of the remaining associations.

For example, the $L \times L$ model Eqn. (8.20) of uniform association applies directly to square tables, and, for square tables, can also be amended to include a diagonals term, `Diag()`, giving a model of *quasi-uniform association*. In this model, all adjacent 2×2 sub-tables not involving diagonal cells have a common local odds ratio.

A related model is the *crossings model* (Goodman, 1972). This hypothesizes that there are different difficulty parameters for crossing from one category to the next, and that the associations between categories decreases with their separation. In the crossings model for an $I \times I$ table, there are $I - 1$ crossings parameters, $\nu_1, \nu_2, \dots, \nu_{I-1}$. The association parameters, λ_{ij}^{AB} have the form of the product of the intervening ν parameters,

$$\lambda_{ij}^{AB} = \begin{cases} \prod_{k=j}^{k=i-1} \nu_k & : i > j \\ \prod_{k=j-1}^{k=i} \nu_k & : i < j \end{cases}$$

This model can also be cast in *quasi* form, by addition of a `Diag` term to fit the main diagonal cells. See Powers and Xie (2008, §4.4.7) for further details of this model. The `Crossings()` function in `vcdExtra` implements such crossings terms.

{sec:sg-ordinal}

{ex:hauser2}

EXAMPLE 8.10: Hauser's occupational mobility table

Without much comment or detail, for reference we first fit some of the ordinal models to the `Hauser79` data: Uniform association ($L \times L$), row effects, and the `RC(1)` model.

```
> Fscore <- as.numeric(Hauser79$Father)      # numeric scores
> Sscore <- as.numeric(Hauser79$Son)        # numeric scores
>
> # uniform association
> hauser.UA <- update(hauser.indep, ~ . + Fscore*Sscore)
> # row effects model
> hauser.roweff <- update(hauser.indep, ~ . + Father*Sscore)
> # RC model
> hauser.RC <- update(hauser.indep, ~ . + Mult(Father, Son), verbose=FALSE)
```

All of these fit very poorly, yet they are all substantial improvements over the independence model.

```
> vcdExtra::LRstats(hauser.indep, hauser.UA, hauser.roweff, hauser.RC)

Likelihood summary table:
  AIC BIC LR Chisq Df Pr(>Chisq)
hauser.indep 6391 6402     6170 16    <2e-16 ***
hauser.UA    2503 2516     2281 15    <2e-16 ***
hauser.roweff 2309 2325     2080 12    <2e-16 ***
hauser.RC     920  940      685  9    <2e-16 ***

---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The $L \times L$ model, hauser.UA might be improved by ignoring the diagonals, and, indeed it is.

```
> hauser.UAdiag <- update(hauser.UA, ~ . + Diag(Father, Son))
> anova(hauser.UA, hauser.UAdiag, test="Chisq")

Error in anova.gnm(hauser.UA, hauser.UAdiag, test = "Chisq"): could not find
function "anova.glmlist"
```

In this model, the estimated common local log odds ratio—the coefficient for the linear-by-linear term Fscore:Sscore is

```
> coef(hauser.UAdiag)[["Fscore:Sscore"]]
[1] 0.1584
```

For comparisons not involving the diagonal cells, each step down the scale of occupational categories for the father multiplies the odds that the son will also be in one lower category by $\exp(0.158) = 1.172$, an increase of 17%.

The crossings model, with and without the diagonal cells can be fit as follows:

```
> hauser.CR <- update(hauser.indep, ~ . + Crossings(Father, Son))
> hauser.CRdiag <- update(hauser.CR, ~ . + Diag(Father, Son))
> vcdExtra::LRstats(hauser.CR, hauser.CRdiag)

Likelihood summary table:
  AIC BIC LR Chisq Df Pr(>Chisq)
hauser.CR     319 334     89.9 12    5.1e-14 ***
hauser.CRdiag 299 318     64.2  9    2.0e-10 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The quasi-crossings model hauser.CRdiag has a reasonable G^2 fit statistic, and its interpretation and lack of fit is worth exploring further. The crossings coefficients ν can be extracted as follows.

```
> nu <- coef(hauser.CRdiag) [pickCoef(hauser.CRdiag, "Crossings")]
> names(nu) <- gsub("Crossings(Father, Son)C", "nu", names(nu), fixed=TRUE)
> nu
  nu1      nu2      nu3      nu4
-0.42275 -0.38768 -0.27500 -1.40244
```

They indicate the steps between adjacent categories in terms of the barriers for a son moving to a lower occupational category. The numerically largest gap separates the lower non-manual category from farming.

In contrast to the UAdiag model, the quasi-crossing model with diagonal terms implies that all 2×2 off-diagonal sub-tables are independent, i.e., the local odds ratios are all equal to 1.0. The reasons for lack of fit of this model can be seen in the corresponding mosaic display, shown in Figure 8.16

```
> mosaic(hauser.CRdiag, ~Father+Son,
+       gp=shading_Friendly, residuals_type="rstandard",
+       main="Crossings() + Diag()")
```

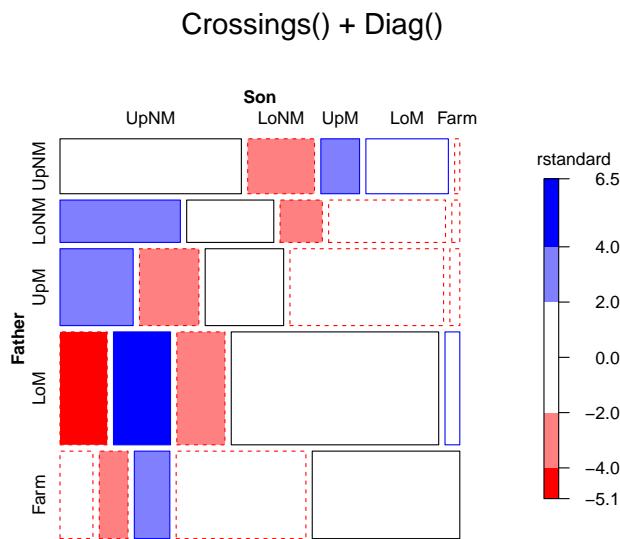


Figure 8.16: Mosaic display for the quasi-crossings model fit to the Hauser79 data.^{fig:hauser-mosaic3}

It can be seen that lack of fit for this model is largely concentrated in the lower triangle, where the father's occupation is lower than that of his son.

In this example and the last, we have fit quite a few different models to the Hauser (1979) data. In presentations, articles and books it is common to summarize such a collection in a table, sorted by G^2 , degrees of freedom, AIC or BIC, to show their ordering along some metric. For instance, here we collect all the models fit in Example 8.9 and this example in a **glmlist()** and sort in decreasing order of BIC to show model fit by this measure.

```
> modlist <- glmlist(hauser.indep, hauser.roweff, hauser.UA, hauser.UAdiag,
+                      hauser.quasi, hauser.qsymm, hauser.topo,
```

```

+           hauser.RC, hauser.CR, hauser.CRdiag)
> LRstats(modlist, sortby="BIC")

Likelihood summary table:
      AIC  BIC  LR Chisq Df Pr(>Chisq)
hauser.indep 6391 6402      6170 16   < 2e-16 ***
hauser.UA    2503 2516      2281 15   < 2e-16 ***
hauser.roweff 2309 2325      2080 12   < 2e-16 ***
hauser.RC    920   940      685  9   < 2e-16 ***
hauser.quasi  914   931      683 11   < 2e-16 ***
hauser.CR     319   334      90 12   5.1e-14 ***
hauser.UAdiag 306   324      73 10   1.2e-11 ***
hauser.CRdiag 299   318      64  9   2.0e-10 ***
hauser.topo   295   311      67 12   1.4e-09 ***
hauser.qsymm  268   291      27  6   0.00012 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

When there are more than just a few models, a more useful display is a **model comparison plot** of measures like G^2/df , AIC or BIC against degrees of freedom. For example, Figure 8.17 plots BIC against Df from the result of `LRstats()`. Because interest is focused on the smallest values of BIC and these values span a large range, BIC is shown on the log scale using `log="y"`.

```

> sumry <- LRstats(modlist)
> mods <- substring(rownames(sumry), 8)
> with(sumry, {
+   plot(Df, BIC, cex=1.3, pch=19,
+         xlab='Degrees of freedom', ylab='BIC (log scale)',
+         log="y", cex.lab=1.2)
+   pos <- ifelse(mods=="UAdiag", 1, 3)
+   text(Df, BIC+55, mods, pos=pos, col='red', xpd=TRUE, , cex=1.2)
+ })

```

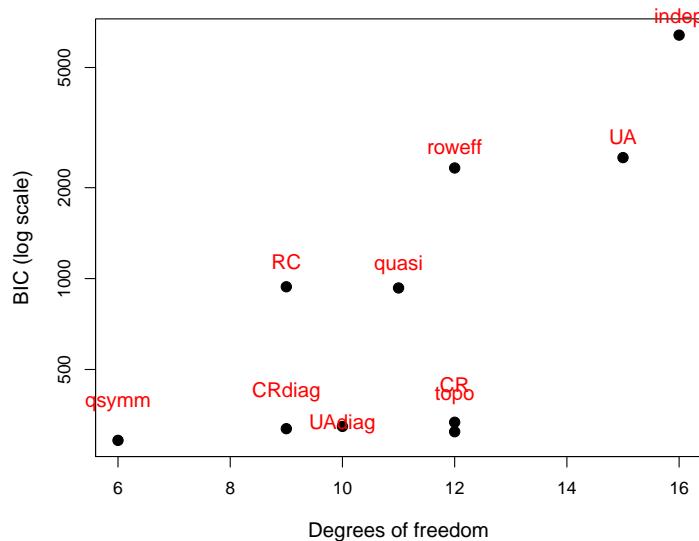


Figure 8.17: Model comparison plot for the models fit to the Hauser79 data fig:hauser-sumry-plot

Compared with the sorted tabular display shown above, such a plot sorts the models *both* by a measure of fit and by model complexity (degrees of freedom). Figure 8.17 shows that the quasi-symmetry model is best by BIC, but also shows that the next four best models by this measure are quite similar in terms of BIC. Similar plots for AIC and G^2/df show that the model of quasi-symmetry is favored by these measures.

△

8.8 Three-way and higher-dimensional tables

{sec:loglin-3wayord}

The models and methods for ordinal factors and square tables described in Section 8.6 and Section 8.7 extend readily to multidimensional tables with these properties for some of the factors. In three-way tables, these models provide a more parsimonious account than the saturated model, $[ABC]$, and also allow simpler models than the general model of homogeneous association, $[AB][AC][BC]$ using scores for ordinal factors or terms for symmetry and diagonal factors in square layers.

For example, consider the case where all three factors are ordinal and the model of homogeneous association $[AB][AC][BC]$ fits poorly. In this case we can generalize the model of uniform association by assigning scores a , b and c and model the three-way association, λ_{ijk}^{ABC} as

$$\lambda_{ijk}^{ABC} = \gamma a_i b_j c_k$$

with only one more parameter. This gives the model of ***uniform interaction*** (or *homogeneous uniform association*)

$$\log(m_{ijk}) = \mu + \lambda_i^A + \lambda_j^B + \lambda_k^C + \lambda_{ij}^{AB} + \lambda_{ik}^{AC} + \lambda_{jk}^{BC} + \gamma a_i b_j c_k . \quad (8.27) \quad \text{(eq:uni-inter)}$$

This model posits that (with equally spaced scores) all local odds ratios θ_{ijk} in adjacent rows, columns and layers are constant,

$$\log(\theta_{ijk}) = \gamma \quad \forall i, j, k$$

The homogeneous association model is the special case of $\log \theta_{ijk} = \gamma = 0$.

A less restricted model of ***heterogeneous uniform association*** retains the linear-by-linear form of association for factors A and B , but allows the strength of this association to vary over layers, C , representing λ_{ijk}^{ABC} as

$$\lambda_{ijk}^{ABC} = (\gamma + \gamma_k) a_i b_j$$

with the constraint $\sum_k \gamma_k = 0$. This model is equivalent to fitting separate models of uniform association at each level k of factor C and gives estimates of the conditional local log odds ratios, $\log \theta_{ij(k)} = \gamma + \gamma_k$.

Following the development in Section 8.6 there is a large class of other models for ordinal factors (see Figure 8.6), where not all factors are assigned scores. For three-way tables, these can be represented in homogeneous form when the two-way association of A and B is the same for all levels of C , or in a heterogeneous form, when it varies over C .

Similarly, the models for square tables described in Section 8.7 extend to three-way tables with several layers (strata), allowing both homogeneous and heterogeneous terms for diagonals and symmetry describing the AB association over levels of C .

{ex:vision-glm2}

EXAMPLE 8.11: Visual acuity

We continue the analysis of the *VisualAcuity* data, but now consider the three-way, $4 \times 4 \times 2$ table comprising both men and women. The main questions here are whether the pattern of quasi-symmetry observed in the analysis for women also pertains to men and whether there is heterogeneity of the association between right, left acuity across gender.

A useful first step for n -dimensional tables is to consider the models composed of all 1-way, 2-way, ... n -way terms as a quick overview. The function `Kway()` in `vcdExtra` package does this automatically, returning a "glmlist" object containing the fitted models.¹¹

```
> vis.kway <- Kway(Freq ~ right + left + gender, data=VisualAcuity)
> vcdExtra::LRstats(vis.kway)

Likelihood summary table:
      AIC    BIC LR Chisq Df Pr(>Chisq)
kway.0 13857 13858     13631 31    < 2e-16 ***
kway.1  9925  9937      9686 24    < 2e-16 ***
kway.2   298   332       28  9    0.00079 ***
kway.3   287   334       0  0    < 2e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

This shows that the model of homogeneous association `kway.2` ($[RL][RG][LG]$) does not fit well, but it doesn't account for diagonal agreement or symmetry to simplify the associations.

As a basis for comparison, we first fit the simple models of quasi-independence and quasi-symmetry that do not involve `gender`, asserting the same pattern of diagonal and off-diagonal cells for males and females.

```
> vis.indep <- glm(Freq ~ right + left + gender, data = VisualAcuity,
+                      family=poisson)
> vis.quasi <- update(vis.indep, . ~ . + Diag(right, left))
> vis.qsymm <- update(vis.indep, . ~ . + Diag(right, left) + Symm(right, left))
>
> LRstats(vis.indep, vis.quasi, vis.qsymm)

Likelihood summary table:
      AIC    BIC LR Chisq Df Pr(>Chisq)
vis.indep 9925  9937     9686 24    <2e-16 ***
vis.quasi  696   714      449 20    <2e-16 ***
vis.qsymm  435   456      184 18    <2e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The model of homogeneous quasi-symmetry fits quite badly, even worse than the all two-way association model. We can see why in the mosaic for this model, shown in Figure 8.18.

```
> mosaic(vis.qsymm, ~ gender + right + left, condvars="gender",
+          residuals_type="rstandard", gp=shading_Friendly,
+          labeling_args=largs,
+          main="Homogeneous quasi-symmetry")
```

It can be seen in Figure 8.18 that the pattern of residuals for men and women are nearly completely opposite in the upper and lower portions of the plot: men have positive residuals in the same right, left cells where women have negative residuals, and vice-versa. In particular, the diagonal cells of both tables have large absolute residuals, because the term `Diag(right, left)` fits a common set of diagonals for both men and women.

We can correct for this by allowing separate diagonal and symmetry terms, given as interactions of `gender` with `Diag()` and `Symm()`.

```
> vis.hetdiag <- update(vis.indep, . ~ . + gender*Diag(right, left) +
+                         Symm(right, left))
> vis.hetqsymm <- update(vis.indep, . ~ . + gender*Diag(right, left) +
```

¹¹For completeness, this also fits the 0-way model, corresponding to $\log m_{ijk\dots} = \mu$, or the model formula `Freq ~ 1`.

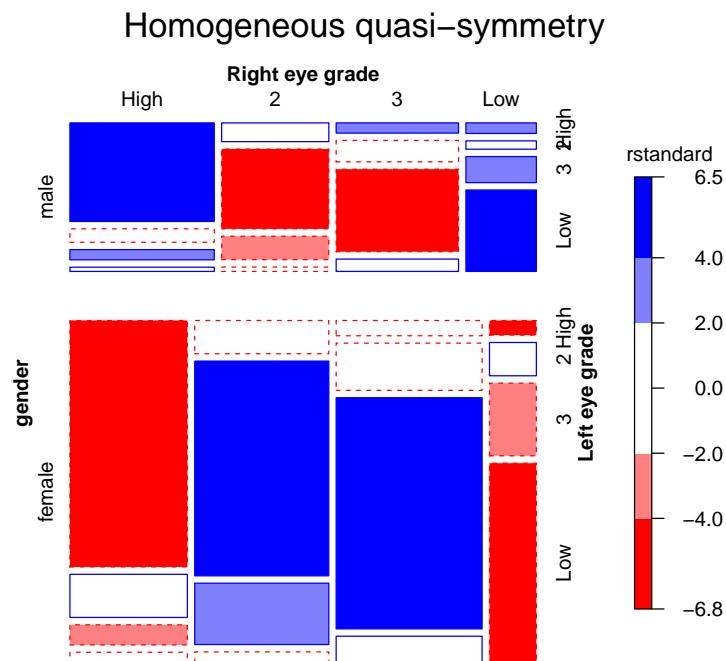


Figure 8.18: Mosaic display for the model of homogeneous quasi-symmetry fit to the VisualAcuity data.
fig:visionz-quasy

```

+           gender*Symm(right, left))
> #vis.hetmodels <- glmlist(vis.qsymm, vis.hetdiag, vis.hetqsymm)
> LRestats(vis.qsymm, vis.hetdiag, vis.hetqsymm)

Likelihood summary table:
      AIC BIC LR Chisq Df Pr(>Chisq)
vis.qsymm   435 456   183.7 18     < 2e-16 ***
vis.hetdiag  312 338    52.3 14     2.5e-06 ***
vis.hetqsymm 287 321    17.7  9      0.038 *
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

Note that the model `vis.hetqsymm` fits better than the model `vis.hetdiag` in absolute terms and by AIC, but the latter, with fewer parameters, fits better by BIC. The mosaic for the model `vis.hetqsymm` is shown in Figure 8.19.

```

> mosaic(vis.hetqsymm, ~ gender + right + left, condvars="gender",
+         residuals_type="rstandard", gp=shading_Friendly,
+         labeling_args=largs,
+         main="Heterogeneous quasi-symmetry")

```

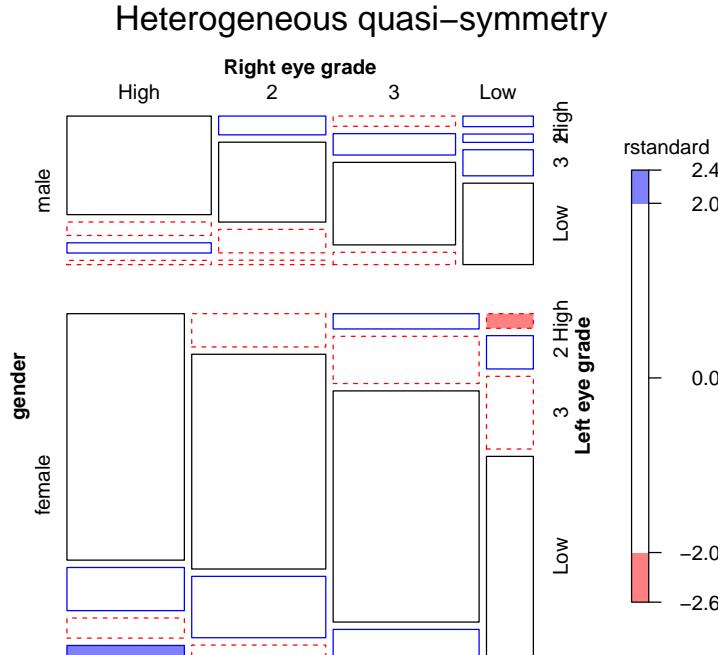


Figure 8.19: Mosaic display for the model of heterogeneous quasi-symmetry fit to the VisualAcuity data.
fig:visionz-hetqsymm

As in the two-way case, this model now fits the diagonal cells in each table exactly, effectively ignoring this part of the association between right and left eye acuity. All remaining residuals are relatively small in magnitude, except for the two opposite off-diagonal cells (Low, High) and (High, Low) in the table for women.

The substantive interpretation of this example is that visual acuity is largely the same (diagonal cells) in the right and left eyes of both men and women. Ignoring the diagonal cells, when visual

acuity differs, both men and women exhibit approximately symmetric associations. However, deviations from symmetry (Figure 8.18) are such that men are slightly more likely to have a lower grade in the right eye, while women are slightly more likely to have a higher grade in the right eye.



8.9 Multivariate responses

{sec:loglin-multiv}

In many studies, there may be *several* categorical responses observed along with one or more explanatory variables. In a clinical trial, for example, the efficacy of a drug might be the primary response, but the occurrence of side-effects might give rise to additional response variables of substantive interest. Or, in a study of occupational health, the occurrence of two or more distinct symptoms might be treated as response variables.

If there are *no* explanatory variables, then the problem is simply to understand the joint distribution of the response categories, and the loglinear models and graphical displays described earlier are sufficient. Otherwise, in these cases we usually wish to understand how the various responses are affected by the explanatory variables. Moreover, it may also be important to understand how the association between the categorical responses depends on the explanatory variables. That is, we would like to study how *both* the marginal distributions of the responses, and their joint distribution depends on the predictors. In the occupational health example, the goal might be to understand both how the prevalence of several symptoms varies with one or more predictors, and how the association (loosely, “correlation”) among those symptoms varies with those predictors.

Although the general loglinear model is often used in these situations, there are special reparameterizations that may be used to separate the *marginal* dependence of each response on the explanatory variables from the relationship of the *association* among the responses on the explanatory variables.

Let us say that categorical responses, Y_1, Y_2, \dots have been observed, together with possible explanatory variables, X_1, X_2, \dots , and let $\pi_{ij\dots}$ be the joint probability of all the responses and explanatory variables; we also use \boldsymbol{x} to refer to the values of X_1, X_2, \dots

Note that the minimal model of independence of all responses from each other and from the explanatory variables is the loglinear model $[Y_1][Y_2] \cdots [X_1 X_2 \cdots]$ (i.e., all associations among the X_i must be included). A no-effect model, in which the responses do not depend on the explanatory variables, but may be associated among themselves is $[Y_1 Y_2 \cdots][X_1 X_2 \cdots]$. However, these models do not separate the individual (marginal) effects of X_1, X_2, \dots on each Y_i from their associative effects on the joint relationships among the Y_i .

There are three useful general approaches which *do* separate these effects:

1. Model the marginal dependence of each response, Y_i separately on X_1, X_2, \dots , and, in addition, model the interdependence among the responses, Y_1, Y_2, \dots ¹²
2. Model the joint dependence of all responses on X_1, X_2, \dots , but parameterized so that marginal and associative effects are delineated.
3. Construct simultaneous models, estimated together, for the marginal and joint dependence of the responses on the explanatory variables.

The first approach is the simplest, an informative starting place, and is satisfactory in the (often unlikely) case that the responses are not associated, or if the associations among responses do not vary much over the explanatory variables (i.e., no terms like $[Y_1 Y_2 X_j]$ are required). In the clinical

¹²For quantitative responses, this is roughly analogous to fitting univariate response models for each Y_i , followed by something like a principal component analysis of the relationships among the Y_i . But in this case, the multivariate linear model, $\mathbf{Y} = \mathbf{X}\mathbf{B} + \mathbf{E}$ provides a general solution.

trial example, we would construct separate loglinear or logit models for efficacy of the drug, and for occurrence of side-effects, and supplement these analyses with mosaic or other displays showing the relations between efficacy and side-effects and a model for their joint association. If those who improve with the drug also show more serious side effects, the worth of the treatment would be questioned. A limitation of this method is that it does not provide an overall model comprising these effects.

In the second approach, the joint probabilities, $\pi_{ij\dots}$, are recast to give separate information regarding the dependence of the univariate marginal probabilities $\pi_{i\bullet}, \pi_{\bullet j}, \dots$, on the explanatory variables and the dependence of the intra-response associations on the explanatory variables. The VGAM package provides several versions of this approach with the function `vglm()` (for *vector generalized linear model*).

The third approach, developed, for example, by Lang and Agresti (1994), is the most general, and provides a scheme to represent a model $\mathcal{J}(\bullet)$ for the joint distributions of the X, Y variables together with a model $\mathcal{M}(\bullet)$ for their first-order marginal distributions. The joint models are typically loglinear models, ranging from the mutual independence model, $\mathcal{J}(I) = [Y_1][Y_2]\dots[X_1][X_2]\dots$ to the saturated model, $\mathcal{J}(S) = [Y_1 Y_2 \dots X_1 X_2 \dots]$, while the marginal models are logit models for the response variables. The combined model, denoted $\mathcal{J}(\bullet) \cap \mathcal{M}(\bullet)$, is estimated simultaneously by maximum likelihood. This approach is implemented in R in the `hmmm` package (hierarchical multinomial marginal models). However, model specification in this implementation is complicated, and it will not be considered further here.

8.9.1 Bivariate, binary response models

We focus here on two related models reflecting the second approach, as discussed by McCullagh and Nelder (1989, Section 6.5). We consider here only the case of two binary responses, though the general approach can be applied to $R > 2$ responses Y_1, Y_2, \dots, Y_R , and these may be polytomous or ordinal.

Let \boldsymbol{x} refer to the values of all the explanatory variables and let $\pi_{ij}(\boldsymbol{x})$ be the joint probabilities in cell $Y_1 = i, Y_2 = j$. The essential idea of the **bivariate logistic model** arises from a linear transformation of the cell probabilities $\boldsymbol{\pi}$ to interpretable functions of the marginal probabilities (logits) and their association (odds ratio), a mapping of $\boldsymbol{\pi} \rightarrow \boldsymbol{\eta}$,

$$\begin{aligned}\eta_1 &= \text{logit}(\pi_{1\bullet}) \\ \eta_2 &= \text{logit}(\pi_{\bullet 1}) \\ \eta_{12} &= \frac{\pi_{11} \pi_{22}}{\pi_{12} \pi_{21}}\end{aligned}\tag{8.28}$$

The predictors in \boldsymbol{x} are then taken into account by considering models that relate $\boldsymbol{\pi}$ to \boldsymbol{x} through $\boldsymbol{\eta}$,

$$\begin{aligned}\eta_1 &= \boldsymbol{x}_1^\top \boldsymbol{\beta}_1 \\ \eta_2 &= \boldsymbol{x}_2^\top \boldsymbol{\beta}_2 \\ \eta_{12} &= \boldsymbol{x}_{12}^\top \boldsymbol{\beta}_{12}\end{aligned}\tag{8.29}$$

where $\boldsymbol{x}_1, \boldsymbol{x}_2$ and \boldsymbol{x}_{12} are subsets of the predictors in \boldsymbol{x} for each sub-model, and $\boldsymbol{\beta}_1, \boldsymbol{\beta}_2$ and $\boldsymbol{\beta}_{12}$ are the corresponding parameters to be estimated.

McCullagh and Nelder (1989) arrive at this joint bivariate model in two steps. First, transform the cell probabilities $\boldsymbol{\pi}$ to a vector of probabilities $\boldsymbol{\gamma}$ which also includes the univariate margins, given by

$$\boldsymbol{\gamma} = \boldsymbol{L}\boldsymbol{\pi}\tag{8.30}$$

where \mathbf{L} is a matrix of 0s and 1s of the form of a factorial design matrix. In the 2×2 case,

$$\gamma = \begin{pmatrix} \pi_{1\bullet} \\ \pi_{2\bullet} \\ \pi_{\bullet 1} \\ \pi_{\bullet 2} \\ \pi_{11} \\ \pi_{12} \\ \pi_{21} \\ \pi_{22} \end{pmatrix} = \begin{bmatrix} 1 & 1 & 0 & 0 \\ 0 & 0 & 1 & 1 \\ 1 & 0 & 1 & 0 \\ 0 & 1 & 0 & 1 \\ 1 & 0 & 0 & 0 \\ 0 & 1 & 0 & 0 \\ 0 & 0 & 1 & 0 \\ 0 & 0 & 0 & 1 \end{bmatrix} \begin{pmatrix} \pi_{11} \\ \pi_{12} \\ \pi_{21} \\ \pi_{22} \end{pmatrix}. \quad (8.31)$$

There are of course only three linearly independent probabilities, because $\sum \sum \pi_{ij} = 1$. In the second step, the bivariate logistic model is formulated in terms of factorial contrasts on the elements of γ which express separate models for the two logits and the log odds. The model is expressed as

$$\eta = \mathbf{C} \log \gamma = \mathbf{C} \log \mathbf{L}\pi, \quad (8.32) \quad \text{eq:eta1}$$

where \mathbf{C} is a matrix of contrasts. In the 2×2 case, the usual contrasts may be defined by

$$\eta = \begin{pmatrix} \eta_1 \\ \eta_2 \\ \eta_{12} \end{pmatrix} = \begin{pmatrix} \text{logit } \pi_{1\bullet} \\ \text{logit } \pi_{\bullet 1} \\ \theta_{12} \end{pmatrix} = \begin{bmatrix} 1 & -1 & 0 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & 1 & -1 & 0 & 0 & 0 & 0 \\ 0 & 0 & 0 & 0 & 1 & -1 & -1 & 1 \end{bmatrix} \begin{pmatrix} \pi_{1\bullet} \\ \pi_{2\bullet} \\ \pi_{\bullet 1} \\ \pi_{\bullet 2} \\ \pi_{11} \\ \pi_{12} \\ \pi_{21} \\ \pi_{22} \end{pmatrix} \quad (8.33) \quad \text{eq:eta2}$$

Thus, we are modeling the marginal odds of each response, together with the log odds ratio θ_{12} simultaneously.

Specific models are then formulated for the dependence of $\eta_1(\mathbf{x}), \eta_2(\mathbf{x})$ and $\eta_{12}(\mathbf{x})$ on some or all of the explanatory variables. For example, with one quantitative explanatory variable, x , the model

$$\begin{pmatrix} \eta_1 \\ \eta_2 \\ \eta_{12} \end{pmatrix} = \begin{pmatrix} \alpha_1 + \beta_1 x \\ \alpha_2 + \beta_2 x \\ \theta \end{pmatrix} \quad (8.34) \quad \text{eq:bilogit1}$$

asserts that the log odds of each response changes linearly with x , while the odds ratio between the responses remains constant. In the general form given by McCullagh and Nelder (1989) the submodels in Eqn. (8.34) may each depend on the explanatory variables in different ways. For example, the logits could both depend quadratically on x , while an intercept-only model could be posited for the log odds ratio.

The second model is the **bivariate loglinear model**, the special case obtained by taking $\mathbf{L} = \mathbf{I}$ in Eqn. (8.30) and Eqn. (8.32) so that $\gamma = \pi$. Then a loglinear model of the form

$$\eta(\mathbf{x}) = \mathbf{C} \log \pi$$

expresses contrasts among log probabilities as linear functions of the explanatory variables. For the 2×2 case, we take the contrasts \mathbf{C} as shown below

$$\eta = \begin{pmatrix} l_1 \\ l_2 \\ \eta_{12} \end{pmatrix} = \begin{bmatrix} 1 & 1 & -1 & -1 \\ 1 & -1 & 1 & -1 \\ 1 & -1 & 1 & -1 \end{bmatrix} \begin{pmatrix} \log \pi_{11} \\ \log \pi_{12} \\ \log \pi_{21} \\ \log \pi_{22} \end{pmatrix} \quad (8.35) \quad \text{eq:eta3}$$

and models for the dependence of $l_1(\mathbf{x})$, $l_2(\mathbf{x})$ and $\eta_{12}(\mathbf{x})$ are expressed in the same way as in Eqn. (8.34). The estimates of the odds ratio, η_{12} are the same under both models. The marginal functions are parameterized differently, however, but lead to similar predicted probabilities.

In R, bivariate logistic models of the form Eqn. (8.28) and Eqn. (8.29) can be fit using `vglm()` with the `binom2.or()` family in the `VGAM` package.¹³ The fitting and graphing of these models is illustrated in the next example.

EXAMPLE 8.12: Breathlessness and wheeze in coal miners

In Example 4.11 we examined the association between the occurrence of two pulmonary conditions, breathlessness and wheeze, among coal miners classified by age (Ashford and Sowden, 1970). Figure 4.5 showed fourfold displays focused on the odds ratio for the co-occurrence of these symptoms, and Figure 4.6 plotted these odds ratios against age directly. Here, we consider models which examine the changes in prevalence of the two symptoms over age, together with the changes in their association.

Plotting bivariate response data

As a starting point and overview of what is necessary for bivariate response models, we calculate the empirical log odds for breathlessness and for wheeze, and the log odds ratio for their association in each 2×2 table. The log odds ratios are the same values plotted in Figure 4.6 (but the youngest age group was not included in the earlier analysis).

The `CoalMiners` data is $2 \times 2 \times 9$ table. For convenience in this analysis (and for use with `VGAM`) we convert it to a 4×9 data frame, and relabel the columns to use the combinations of ("B", "b") and ("W", "w") to represent the conditions of breathlessness and wheeze, where the upper case letter indicates presence of the condition. A variable `age` is also created, using the midpoints of the age categories.

```
> data("CoalMiners", package="vcd")
> coalminers <- data.frame(t(matrix(aperm(CoalMiners, c(2,1,3)),
+                                         4, 9)))
> colnames(coalminers) <- c("BW", "Bw", "bW", "bw")
> coalminers$age <- c(22, 27, 32, 37, 42, 47, 52, 57, 62)
> coalminers

   BW   Bw   bW   bw age
1  9    7  95 1841  22
2 23    9 105 1654  27
3 54   19 177 1863  32
4 121   48 257 2357  37
5 169   54 273 1778  42
6 269   88 324 1712  47
7 404  117 245 1324  52
8 406  152 225  967  57
9 372  106 132  526  62
```

With the data in this form, a simple function `blogits()` in `vcdExtra` calculates the logits and log odds ratios corresponding to Eqn. (8.28). The `add` argument accommodates cases where there are very small, or 0 frequencies in some cells, and it is common to add a small constant, such as 0.5 to each cell in calculating *empirical logits*. This function is used to calculate the empirical logits and log odds as follows:

```
> logitsCM <- vcdExtra::blogits(coalminers[,1:4], add=0.5)
> colnames(logitsCM)[1:2] <- c("logitB", "logitW")
> logitsCM
```

¹³This package also provides for bivariate and trivariate loglinear models with `loglinb2()` and `loglinb3()`.

```

logitB    logitW    logOR
[1,] -4.73568 -2.86844 3.1956
[2,] -3.97656 -2.55717 3.6583
[3,] -3.31713 -2.09388 3.3790
[4,] -2.73322 -1.84818 3.1327
[5,] -2.21492 -1.42014 3.0069
[6,] -1.73870 -1.10922 2.7770
[7,] -1.10116 -0.79681 2.9217
[8,] -0.75808 -0.57219 2.4368
[9,] -0.31902 -0.22591 2.6318

```

We plot these as shown below, using `matplot()`, which is convenient for plotting multiple columns against a given horizontal variable, `age` here.¹⁴ For ease of interpretation of the log odds, we also use right vertical axis showing the equivalent probabilities for breathlessness and wheeze.

```

> col <- c("blue", "red", "black")
> pch <- c(15, 17, 16)
> age <- coalminers$age
>
> op <- par(mar=c(4, 4, 1, 4)+.2)
> matplot(age, logitsCM, type="p",
+   col=col, pch=pch, cex=1.2, cex.lab=1.25,
+   xlab="Age", ylab="Log Odds or Odds Ratio")
> abline(lm(logitsCM[,1] ~ age), col=col[1], lwd=2)
> abline(lm(logitsCM[,2] ~ age), col=col[2], lwd=2)
> abline(lm(logitsCM[,3] ~ age), col=col[3], lwd=2)
>
> # right probability axis
> probs <- c(.01, .05, .10, .25, .5)
> axis(4, at=qlogis(probs), labels=probs)
> mtext("Probability", side=4, cex=1.2, at=-2, line=2.5)
> # curve labels
> text(age[2], logitsCM[2,1]+.5, "Breathlessness", col=col[1], pos=NULL, cex=1.2)
> text(age[2], logitsCM[2,2]+.5, "Wheeze", col=col[2], pos=NULL, cex=1.2)
> text(age[2], logitsCM[2,3]-.5, "log OR\n(B|W) / (B|w)", col=col[3], pos=1, cex=1.2)
> par(op)

```

In Figure 8.20 we see that both symptoms, while quite rare among young miners, increase steadily with age (or years working in the mine). By age 60, the probability is nearly 0.5 of having either condition. There is a hint of curvilinearity, particularly in the logit for breathlessness. The decline in the odds ratio with age may reflect selection, as miners who had retired for health or other reasons were excluded from the study.

Fitting `glm` models

Next, we illustrate what can easily be achieved using the standard `glm()` approach for loglinear models and why the bivariate models we described are more useful in this situation. `glm()` requires a data frame as input, so first reshape `CoalMiners` to a frequency data frame. For convenience, we simplify the variable names to `B` and `W`.

```

> CM <- as.data.frame(CoalMiners)
> colnames(CM)[1:2] <- c("B", "W")
> str(CM)

'data.frame': 36 obs. of 4 variables:

```

¹⁴It is actually a small graphical misdemeanor to plot logits and odds ratios on the same vertical axis because they are not strictly commensurable. We plead guilty with the explanation that this graph shows what we want to see here and does not distort the data.

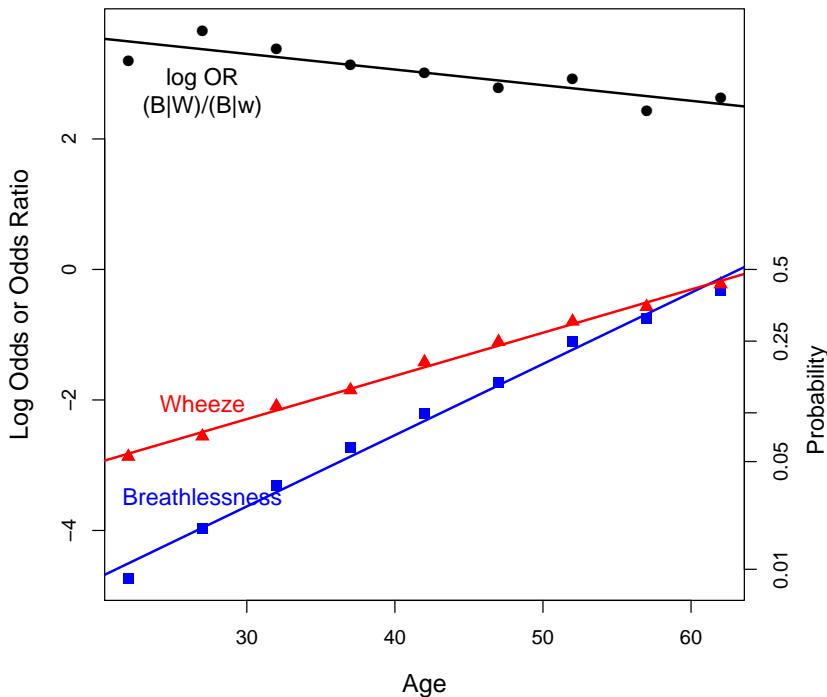


Figure 8.20: Empirical logits and log odds ratio for breathlessness and wheeze in the CoalMiners data. The lines show separate linear regressions for each function. The right vertical axis shows equivalent probabilities for the logits.
[fig:cm-biologits]

```
$ B    : Factor w/ 2 levels "B", "NoB": 1 2 1 2 1 2 1 2 1 2 ...
$ W    : Factor w/ 2 levels "W", "NoW": 1 1 2 2 1 1 2 2 1 1 ...
$ Age  : Factor w/ 9 levels "20-24", "25-29", ...: 1 1 1 1 2 2 2 3 3 ...
$ Freq: num  9 95 7 1841 23 ...
```

As a point of comparison, we fit the mutual independence model, $[B][W][\text{Age}]$ and the baseline model for associated responses, $[BW][\text{Age}]$ which asserts that the association between B and W is independent of Age.

```
> cm.glm0 <- glm(Freq ~ B + W + Age, data=CM, family=poisson)
> cm.glm1 <- glm(Freq ~ B * W + Age, data=CM, family=poisson)
> vcdExtra::LRstats(cm.glm0, cm.glm1)

Likelihood summary table:
      AIC  BIC LR Chisq Df Pr(>Chisq)
cm.glm0 7217 7234     6939 25      <2e-16 ***
cm.glm1 2981 3000     2702 24      <2e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The baseline model `cm.glm1` fits very badly. We can see the pattern of the residual association in a mosaic display for this model shown in Figure 8.21. The formula argument here specifies the order of the variables in the mosaic.

```
> vnames <- list(set_varnames = c(B="Breathlessness", W="Wheeze"))
> lnames <- list(B=c("B", "b"), W = c("W", "w"))
> mosaic(cm.glm1, ~ Age + B + W,
```

```
+ labeling_args=vnames, set_labels=lnames)
```

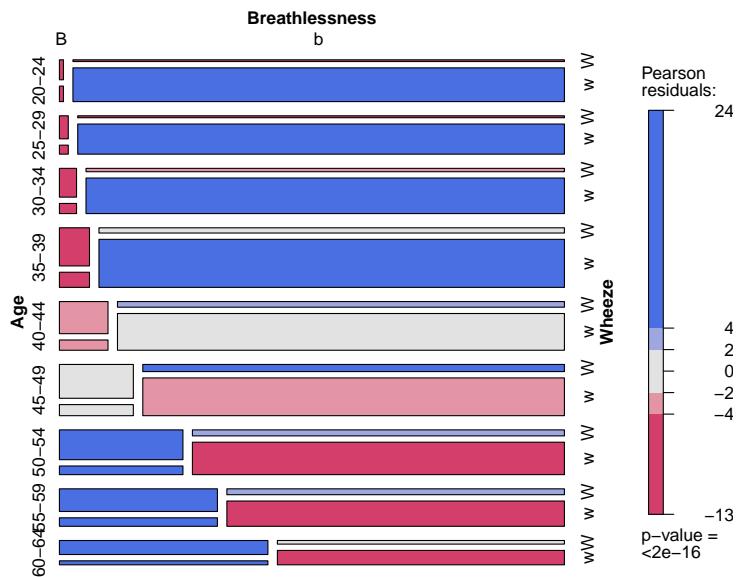


Figure 8.21: Mosaic display for the baseline model, [BW][Age], fit to the CoalMiners data fig:cm-mosaic1

As structured here, it is easy to see the increase in the prevalence of breathlessness and wheeze with age and the changing pattern of their association with age.

From Figure 8.20 and Figure 8.21, it is apparent that both breathlessness and wheeze increase with age, so we can model this by adding terms [B Age][W Age] to the baseline model. This is the no-three-way interaction model, which could also be specified as `Freq ~ (B + W + Age)^2`.

```
> cm.glm2 <- glm(Freq ~ B * W + (B + W) * Age, data=CM, family=poisson)
> vcdExtra::LRstats(cm.glm1, cm.glm2)

Likelihood summary table:
      AIC    BIC LR Chisq Df Pr(>Chisq)
cm.glm1 2981 3000   2702 24      <2e-16 ***
cm.glm2  338   383     27   8      8e-04 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The improvement in fit is substantial, and all terms are highly significant, yet, the residual $G^2(8)$ indicates there is still lack of fit.

```
> library(car)
> Anova(cm.glm2)

Analysis of Deviance Table (Type II tests)

Response: Freq
          LR Chisq Df Pr(>Chisq)
B        11026   1      <2e-16 ***
W        7038   1      <2e-16 ***
```

```

Age      887  8    <2e-16 ***
B:W     3025  1    <2e-16 ***
B:Age    1130  8    <2e-16 ***
W:Age    333   8    <2e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

One way to improve the model using the `glm()` framework is to make use of Age as a quantitative variable and add a term to allow the odds ratio for the [BW] association to vary linearly with age. Here, we construct the variable `age` using the midpoints of the Age intervals.

```
> CM$age <- rep(seq(22, 62, 5), each=4)
```

In the `glm()` approach, the odds ratio cannot be modeled directly, but we can use the following trick: For each 2×2 subtable, the odds ratio can be parameterized in terms of the frequency in any one cell, say, n_{11k} , given that the marginal total n_{++k} is included in the model. We do this by adding a new interaction variable, `ageOR` having the value of `age` for the $(1, 1, k)$ cells and 0 otherwise.

```

> CM$ageOR <- (CM$B=="B") * (CM$W=="W") * CM$age
> cm.glm3 <- update(cm.glm2, . ~ . + ageOR)
> vcdExtra::LRstats(cm.glm0, cm.glm1, cm.glm2, cm.glm3)

Likelihood summary table:
      AIC  BIC  LR Chisq Df Pr(>Chisq)
cm.glm0 7217 7234      6939 25    <2e-16 ***
cm.glm1 2981 3000      2702 24    <2e-16 ***
cm.glm2 338   383       27   8    0.0008 ***
cm.glm3 320   366       7   7    0.4498
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

The model `cm.glm3`, with one more parameter, now fits reasonably well, having residual $G^2(7) = 6.80$. The likelihood ratio test of model `cm.glm3` against `cm.glm2`, which assumes equal odds ratios over age, can be regarded as a test of the hypothesis of homogeneity of odds ratios, against the alternative that the [BW] association changes linearly with age. The `glm()` models fit in this example are summarized above. As usual, `anova()` can be used to compare competing nested models.

```

> anova(cm.glm2, cm.glm3, test="Chisq")

Analysis of Deviance Table

Model 1: Freq ~ B * W + (B + W) * Age
Model 2: Freq ~ B + W + Age + ageOR + B:W + B:Age + W:Age
  Resid. Df Resid. Dev Df Deviance Pr(>Chi)
1          8      26.7
2          7      6.8   1      19.9  8.2e-06 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

This analysis, while useful, also shows the limitations of the `glm()` approach: (a) It doesn't easily allow us to represent and test the substantively interesting hypotheses regarding *how* the prevalence of the binary responses, `B` and `W` vary with `Age`, such as seen in Figure 8.20. (b) It doesn't represent the odds ratio for the [BW] association directly, but only through the coding trick we used here. Thus, it is difficult to interpret the coefficient for `ageOR = -0.02613` in a substantively meaningful way, except that it shows that the odds ratio is decreasing.¹⁵

¹⁵Actually, the interpretability of the coefficient for the log odds ratio can be enhanced here by centering `age`, and representing its units in steps of 5 years, as we do below.

Fitting vglm models

The `vglm()` function in the VGAM package provides a very general implementation of these and other models for discrete multivariate responses. The family function, `binom2.or()` for binary logistic models allows some or all of the logits or odds ratio submodels to be constrained to be intercept-only (e.g., as in Eqn. (8.34)) and the two marginal distributions can be constrained to be equal.

Quantitative predictors (such as age, here), can be modeled linearly or nonlinearly, using `poly()` for a parametric fit, or smooth regression splines, as provided by the functions `ns()`, `bs()` and others in model formulas. In this illustration, we fit bivariate linear and quadratic models in age.

`vglm()` takes its input data in the wide form we called `coalminers` at the beginning of this example. We could use the 9-level factor, `Age` as we did with `glm()`, but we plan to use `age` as a numeric variable in all three submodels. The coefficients in these models will be more easily interpreted if we center `age` and express it as `agec` in units of five years, as shown below.

```
> coalminers <- transform(coalminers, agec=(age-42)/5)
> coalminers$Age <- dimnames(CoalMiners)[[3]]
> coalminers

   BW   Bw   bW   bw age agec   Age
1   9    7   95 1841  22   -4 20-24
2  23    9  105 1654  27   -3 25-29
3  54   19  177 1863  32   -2 30-34
4 121   48  257 2357  37   -1 35-39
5 169   54  273 1778  42    0 40-44
6 269   88  324 1712  47    1 45-49
7 404  117  245 1324  52    2 50-54
8 406  152  225  967  57    3 55-59
9 372  106  132  526  62    4 60-64
```

`vglm()` takes the 2×2 response frequencies as a 4-column matrix on the right hand side of the model formula. However, denoting the responses of failure and success by 0 and 1 respectively, it takes these in the order $y_{00}, y_{01}, y_{10}, y_{11}$. We specify the order below so that the logits are calculated for the occurrence of breathlessness or wheeze, rather than their absence.

```
> library(VGAM)
> #          00  01  10  11
> cm.vglm1 <- vglm(cbind(bw, bW, Bw, BW) ~ agec,
+                      binom2.or(zero=NULL), data=coalminers)
> cm.vglm1

Call:
vglm(formula = cbind(bw, bW, Bw, BW) ~ agec, family = binom2.or(zero = NULL),
      data = coalminers)

Coefficients:
(Intercept):1 (Intercept):2 (Intercept):3           agec:1
-2.26247      -1.48776      3.02191      0.51451
  agec:2        agec:3
  0.32545     -0.13136

Degrees of Freedom: 27 Total; 21 Residual
Residual deviance: 30.394
Log-likelihood: -100.53
```

In this call, the argument `zero=NULL` indicates that none of the linear predictors, $\eta_1, \eta_2, \eta_{12}$ are modeled as constants.¹⁶

¹⁶The default, `zero=3` gives the model shown in Eqn. (8.34), with the odds ratio constant.

At this writing, there is no `anova()` method for the "vgam" objects produced by `vglm()`, but we can test the residual deviance of the model (against the saturated model) as follows, showing that this model has an acceptable fit.

```
> (G2 <- deviance(cm.vglm1))
[1] 30.394
> # test residual deviance
> 1-pchisq(deviance(cm.vglm1), cm.vglm1@df.residual)
[1] 0.084355
```

The estimated coefficients in this model are usefully shown as below, using the argument `matrix=TRUE` in `coef()`. Using `exp()` on the result gives values of odds that can be easily interpreted:

```
> coef(cm.vglm1, matrix=TRUE)
      logit(mu1)  logit(mu2)  log(oratio)
(Intercept) -2.26247   -1.48776    3.02191
agec         0.51451    0.32545   -0.13136
> exp(coef(cm.vglm1, matrix=TRUE))
      logit(mu1)  logit(mu2)  log(oratio)
(Intercept)  0.10409    0.22588   20.5304
agec        1.67282    1.38465   0.8769
```

Thus, the odds of a miner showing breathlessness are multiplied by 1.67, a 67% increase, for each 5 years increase in age; similarly, the odds of wheeze are multiplied by 1.38, a 38% increase. The odds ratio for the association between the two symptoms are multiplied by 0.88, a 12% decrease over each 5 year interval.

The VGAM package has no special plot methods for "vglm" objects, but it is not hard to construct these using the methods we showed earlier in this example. First, we can obtain the fitted probabilities for the 4 response combinations using `fitted()` and the corresponding observed probabilities using `depvar()`.

```
> age <- coalminers$age
> P <- fitted(cm.vglm1)
> colnames(P) <- c("bw", "bW", "Bw", "BW")
> head(P)
      bw       bW       Bw       BW
1 0.93747 0.049409 0.0046356 0.0084831
2 0.91461 0.063636 0.0069757 0.0147776
3 0.88411 0.080029 0.0104965 0.0253679
4 0.84394 0.097484 0.0158138 0.0427671
5 0.79188 0.113839 0.0238598 0.0704196
6 0.72578 0.125910 0.0359684 0.1123366
> Y <- depvar(cm.vglm1)
```

In the left panel of Figure 8.22, we plot the fitted probabilities in the matrix `P` using `matplot()` and the observed probabilities in `Y` using `matpoints()`.

```
> col <- c("red", "blue", "red", "blue")
> pch <- c(1,2,16,17)
>
> op <- par(mar=c(5, 4, 1, 1)+.1)
```

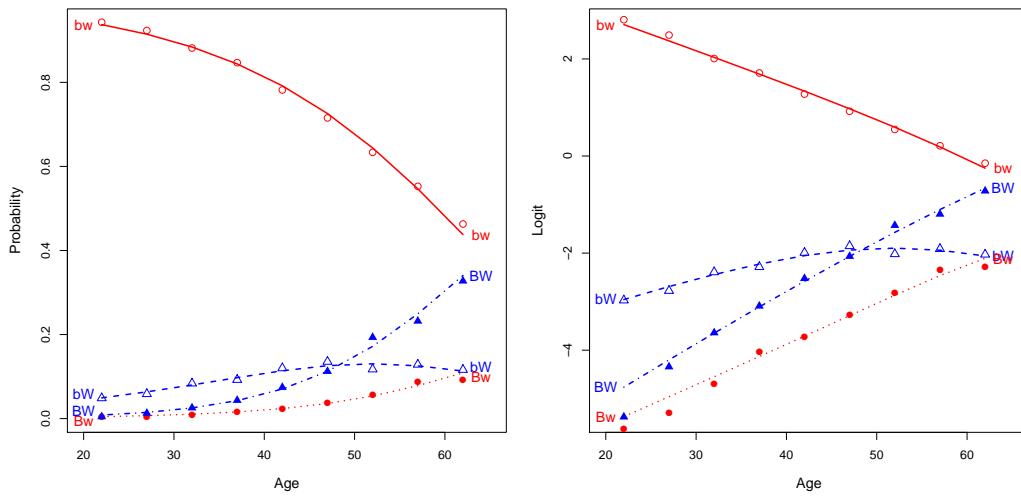


Figure 8.22: Observed and fitted values for the combinations of breathlessness and wheeze in the binary logistic regression model `cm.vglm1`. Left: probabilities; right: on the log odds scale.

```
> matplot(age, P, type="l",
+ col=col,
+ lwd=2, cex=1.2, cex.lab=1.2,
+ xlab="Age", ylab="Probability",
+ xlim=c(20,65))
> matpoints(age, Y,
+ pch=pch, cex=1.2, col=col)
> # legend
> text(64, P[9,]+ c(0,.01, -.01, 0), labels=colnames(P), col=col, cex=1.2)
> text(20, P[1,]+ c(0,.01, -.01, .01), labels=colnames(P), col=col, cex=1.2)
> par(op)
```

The right panel of Figure 8.22 shows these on the log odds scale, produced using the same code as above, applied to the probabilities transformed using `qlogis()`, the quantile function for the logistic distribution.

```
> lP <- qlogis(P)
> lY <- qlogis(Y)
```

In Figure 8.20 we plotted the empirical logits and log odds using the function `bilogits()` to transform frequencies to these values. An essentially identical plot can be produced by transforming the fitted and observed probabilities, as calculated below.

```
> # bilogits, but for B and W
> logitsP <- bilogits(P[,4:1])
> logitsY <- bilogits(Y[,4:1])
```

To test for nonlinearity in the prevalence of the symptoms or their odds ratio with age, we can fit a similar model using `poly()` or a smoothing spline, such as `ns()`. We illustrate this here using a bivariate model allowing quadratic effects of age on all three components.

```
> cm.vglm2 <- vglm(cbind(bw, bW, Bw, BW) ~ poly(agec,2),
+ binom2.or(zero=NULL), data=coalminers)
```

This model has a residual $G^2 = 16.963$ with 18 df. Compared to the linear model `cm.vglm1`, this represents a significant improvement in goodness of fit.

```
> (LR <- deviance(cm.vglm1) - deviance(cm.vglm2))
[1] 13.43
> 1 - pchisq(LR, cm.vglm1@df.residual - cm.vglm2@df.residual)
[1] 0.0037925
```

A plot of the fitted logits and log odds ratios under this model is shown in Figure 8.23. You can interpret this plot as showing that the statistical evidence for the quadratic model indicates some slight tendency for the prevalence of breathlessness and wheeze levels off slightly with age, particularly the former.

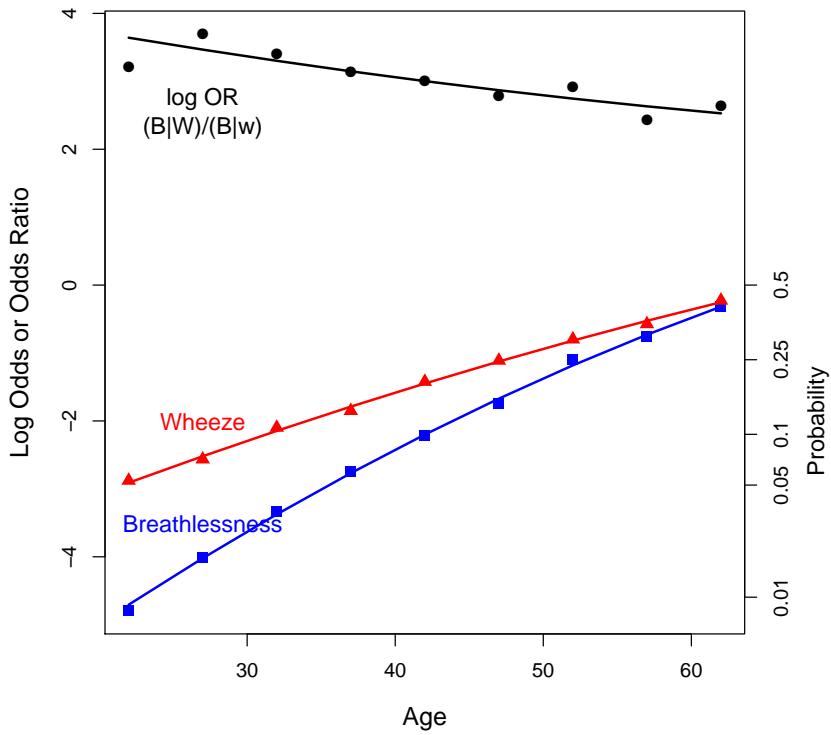


Figure 8.23: Observed (points) and fitted (lines) logits and log odds ratios for the quadratic binary logistic regression model `cm.vglm2`.



8.9.2 More complex models

When there is more than one explanatory variable and several responses, the methods described above using `glm()` and `vglm()` still apply. However, it is useful to begin with a more thorough visual examination of the relations within and between these sets. Some useful graphical displays include:

- mosaic displays showing the marginal relations among the response variables and of the explanatory variables, each collapsed over the other set;
- conditional mosaics or fourfold displays of the associations among the responses, stratified by one or more of the explanatory variables;
- plots of empirical logits and log odds ratios, as in Figure 8.20 or model-based plots, such as Figure 8.23, showing a model-smoothed summary.

These displays can, and should, inform our search for an adequate descriptive or explanatory model. Some of these ideas are illustrated in the following example.

{ex:toxaemia}

EXAMPLE 8.13: Toxaemic symptoms in pregnancy

Brown *et al.* (1983) gave the data used here on two signs of *toxaemia*, an abnormal condition during pregnancy characterized by high blood pressure (hypertension) and high levels of protein in the urine. If untreated, both the mother and baby are at risk of complications or death. The data frame *Toxaemia* in *vcdExtra* represents 13,384 expectant mothers in Bradford, England in their first pregnancy, who were also classified according to social class and the number of cigarettes smoked per day.

There are thus two response variables, and two explanatory variables in this data set in frequency form. For convenience, we also convert it to a $2 \times 2 \times 5 \times 3$ table.

```
> data("Toxaemia", package="vcdeExtra")
> str(Toxaemia)

'data.frame': 60 obs. of 5 variables:
 $ class: Factor w/ 5 levels "1","2","3","4",...: 1 1 1 1 1 1 1 1 1 ...
 $ smoke: Factor w/ 3 levels "0","1-19","20+": 1 1 1 2 2 2 3 3 ...
 $ hyper: Factor w/ 2 levels "Low","High": 2 2 1 1 2 2 1 1 2 2 ...
 $ urea : Factor w/ 2 levels "Low","High": 2 1 2 1 2 1 2 1 2 1 ...
 $ Freq : int 28 82 21 286 5 24 5 71 1 3 ...

> tox.tab <- xtabs(Freq~class + smoke + hyper + urea, Toxaemia)
> ftable(tox.tab, row.vars=1)

      smoke    0          1-19          20+
      hyper Low High   Low High Low High Low High Low High
      urea  Low High  Low High Low High Low High Low High
class
1       286  21  82  28  71  5  24  5  13  0  3  1
2       785  34 266  50 284 17  92 13  34  3 15  0
3      3160 164 1101 278 2300 142 492 120 383 32 92 16
4       656  52 213  63 649 46 129 35 163 12 40  7
5       245  23  78  20 321 34  74 22  65  4 14  7
```

The questions of main interest are how the occurrence of each symptom varies with social class and smoking, and how the association between these symptoms varies. It is useful, however, to examine first the marginal relationship between the two responses, and between the two predictors. The calls to *mosaic()* below produce the two panels in Figure 8.24.

```
> mosaic(~smoke + class, data=tox.tab, shade=TRUE,
+         main="Predictors", legend=FALSE)
> mosaic(~hyper + urea, data=tox.tab, shade=TRUE,
+         main="Responses", legend=FALSE)
```

We see in Figure 8.24 that the majority of the mothers are in the third social class, and that smoking is negatively related to social class, with the highest levels of smoking in classes 4 and 5. (Social class 1 is the highest in status here.) More than 50% are non-smokers. Within the responses, the great majority of women exhibit neither symptom, but showing one symptom makes it much more likely to show the other. Marginally, hypertension is somewhat more prevalent than protein urea.

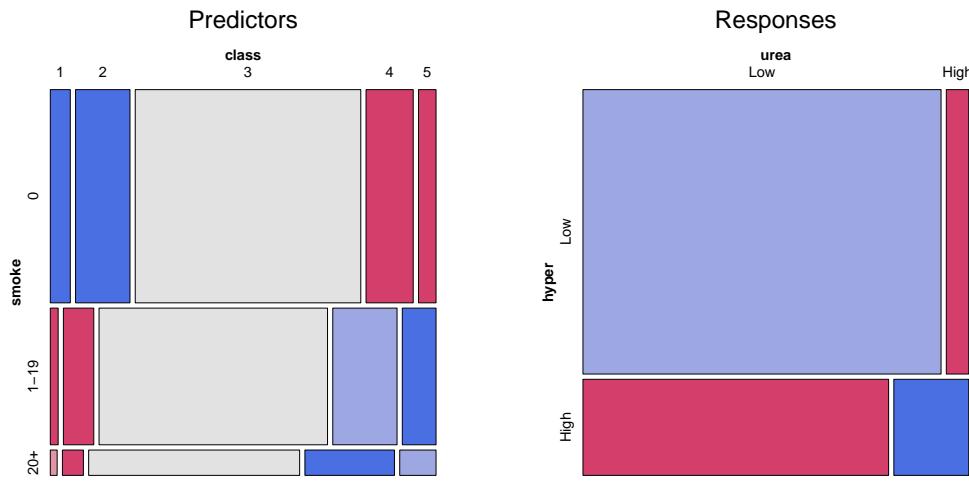


Figure 8.24: Mosaic displays for Toxaemia data: Predictor and response associations fig:tox-mosaic1

We next examine how the association between responses varies with social class and with smoking. Figure 8.25 shows a collection of conditional mosaic plots using `cotabplot()` of the association between hypertension and urea, for each level of smoking, collapsed over social class.

```
> cotabplot(~hyper + urea | smoke, tox.tab, shade=TRUE,
+           legend=FALSE, layout=c(1, 3))
```

```
Error in grid.Call.graphics(L_downvppath, name$path, name$name, strict): Viewport
'panel:smoke=20+|base' was not found
```

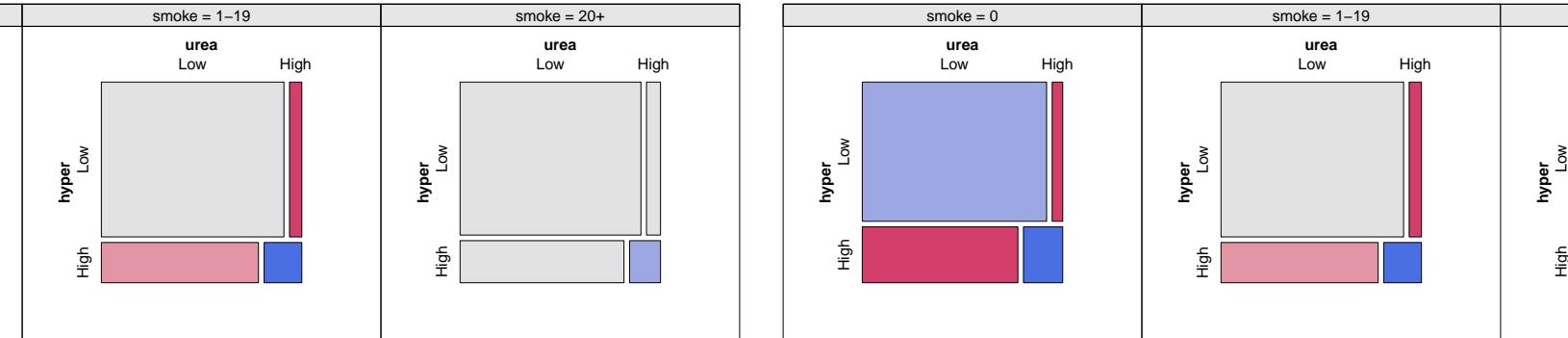


Figure 8.25: Toxaemia data: Response association conditioned on smoking level fig:tox-mosaic2

Figure 8.26 is similar, but stratified by social class. The marginal frequencies of the conditioning variable is not represented in these plots. (For example, as can be seen in Figure 8.24, the greatest number of women are in class 3.)

```
> cotabplot(~hyper + urea | class, tox.tab, shade=TRUE,
+           legend=FALSE, layout=c(1,5))

, , class = 1

      urea
hyper   Low High
  Low    370   26
  High   109   34

, , class = 2

      urea
hyper   Low High
  Low   1103   54
  High   373   63

, , class = 3

      urea
hyper   Low High
  Low   5843   338
  High  1685   414

, , class = 4

      urea
hyper   Low High
  Low   1468   110
  High   382   105

, , class = 5

      urea
hyper   Low High
  Low    631   61
  High   166   49
```

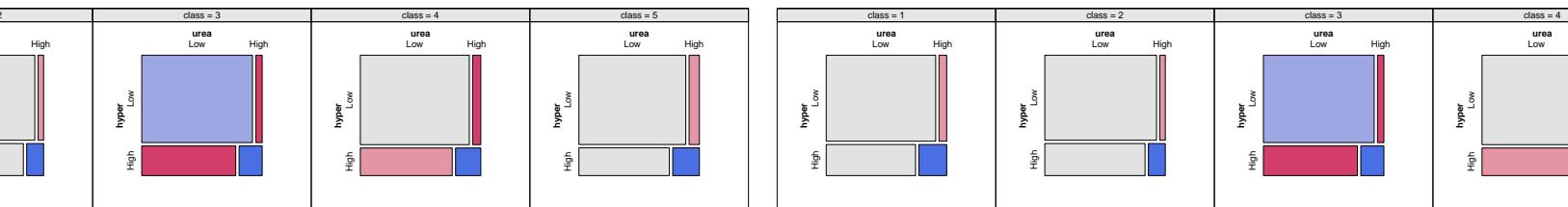


Figure 8.26: Toxaemia data: Response association conditioned on social class fig:tox-mosaic3

Ignoring social class, the association between hypertension and protein urea decreases with smoking. Ignoring smoking, the association is greatest in social class 3. However, these displays don't show directly how the two symptoms are associated in the combinations of social class and smoking. The fourfold display in Figure 8.27, does that.

```
> fourfold(aperm(tox.tab), fontsize=16)
```

It can be seen in Figure 8.27 that the odds ratio appears to increase with both smoking and social class number and these two symptoms are positively associated in nearly all cases. In only two cases the odds ratio is not significantly different from 1: mothers in classes 1 and 2, who smoke more than 20 cigarettes a day, but the frequency in this cell is quite small.

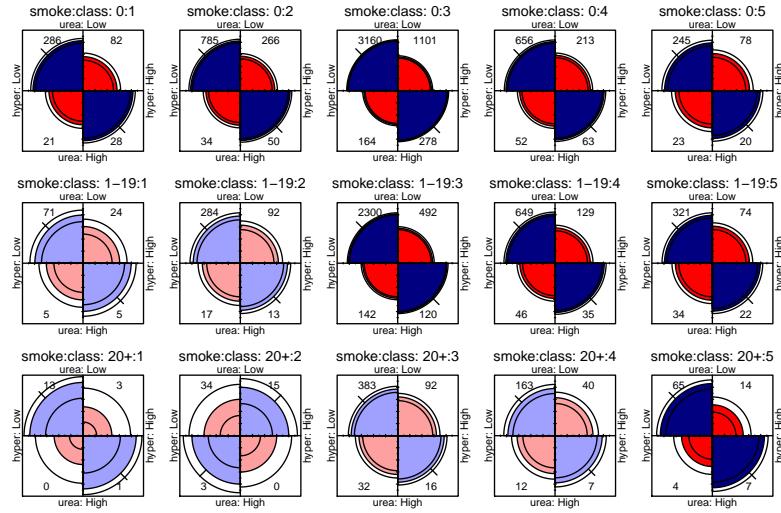


Figure 8.27: Fourfold display for Toxaemia data. Smoking levels vary in the rows and social class in the columns.

(fig:tox-fourfold)

```
> t(apply(tox.tab, MARGIN=1:2, FUN=sum))
```

	class				
smoke	1	2	3	4	5
0	417	1135	4703	984	366
1-19	105	406	3054	859	451
20+	17	52	523	222	90

From these plots, it is useful to examine the association between hypertension and urea more directly, by calculating and plotting the odds ratios. For a $2 \times 2 \times K \times L \times \dots$ table, the function `oddsratio()` in `vcd` calculates these for each 2×2 subtable, and returns an array of dimension $K \times L \times \dots$, together with similar array of standard errors.

```
> LOR <- oddsratio(aperm(tox.tab))
> LOR
```

	1	2	3	4	5
0	1.5370	1.46785	1.5821	1.31676	1.0048
1-19	1.0846	0.85892	1.3739	1.34233	1.0321
20+	2.4485	-1.14579	0.7331	0.86587	2.0949

The `plot()` method for the resulting "logoddsratio" object only handles a single stratum dimension, but in the present case it is easy to plot the result using `matplot()` as we did earlier. The lines below produce Figure 8.28.

```
> matplot(t(LOR), type="b",
+           cex=1.5, pch=15:17, cex.lab=1.5, lwd=2, lty=1,
+           ylab='log odds ratio: Urea | Hypertension',
+           xlab='Social class of mother',
+           xlim=c(1,5.5), col=c("blue", "black", "red"))
+ )
> abline(h=0, col='gray')
> text(5.2, LOR[,5]+c(-.05,.05,0), labels=rownames(LOR), cex=1.25)
> text(5.2, max(LOR[,5])+.2, "Smoking", cex=1.4)
```

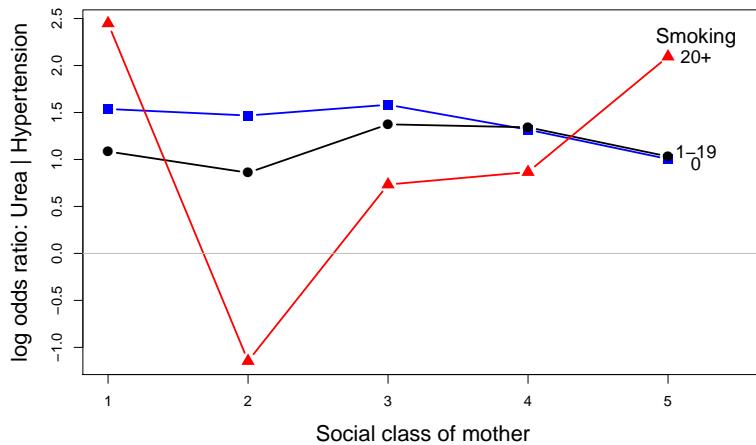


Figure 8.28: Log odds ratios for protein urea given hypertension, by social class and level of maternal smoking¹⁶

The association between the response symptoms, shown in Figure 8.28 is clearer, once we take the variation in sample sizes into account. Except for the heavy smokers, particularly in social classes 1 and 2, the log odds ratio appears to range only between 1–1.5, meaning that, given one symptom, the odds of also having the other range between $\exp(1) = 2.72$ and $\exp(1.5) = 4.48$.

This initial overview of the data is completed by calculating and plotting the log odds for each symptom within each class-smoke population. This could be done in the same way as in Example 8.12, (except that there are now two explanatory factors). The steps used there were: (a) Reshape the $2 \times 2 \times K \cdots$ table to a matrix with four columns corresponding to the binary response combinations. (b) Calculate the logits (and log odds ratio) using `blogits()`. **TODO: Use this as an exercise.**

Here, it is more useful to make separate plots for each of the logits, and we illustrate a more general approach that applies to two or more binary responses, with two or more predictor variables. The essential idea is to fit a separate logit model for each response separately, using the *highest-order interaction* of all predictors (the saturated model). The fitted logits in these models then match those in the data.

```
> tox.hyper <- glm(hyper=='High' ~ class*smoke, weights=Freq,
+                      data=Toxaemia, family=binomial)
> tox.urea <- glm(urea=='High' ~ class*smoke, weights=Freq,
+                      data=Toxaemia, family=binomial)
```

It is then simple to plot these results using the `effects` package as shown in Figure ???. Each plot shows the logit for the response measure against class, with separate curves for the levels of smoking.¹⁷

```
> library(effects)
Error: package 'effects' was built before R 3.0.0: please re-install it
> plot(allEffects(tox.hyper),
```

¹⁷As is usual for effect plots of binary response `glm()` models, the vertical axis is plotted on the scale of log odds, but labeled in terms of probabilities.

```

+   ylab = "Probability (hypertension)",
+   xlab = "Social class of mother",
+   main = "Hypertension: class*smoke effect plot",
+   colors = c("blue", "black", "red"),
+   lwd=3, multiline=TRUE,
+   key.args=list(x=0.05, y=0.2, cex=1.2)
+ )

Error in plot(allEffects(tox.hyper), ylab = "Probability (hypertension)", :
error in evaluating the argument 'x' in selecting a method for function 'plot':
Error: could not find function "allEffects"

> plot(allEffects(tox.urea),
+       ylab = "Probability (Urea)",
+       xlab = "Social class of mother",
+       main = "Urea: class*smoke effect plot",
+       colors = c("blue", "black", "red"),
+       lwd=3, multiline=TRUE,
+       key.args=list(x=0.65, y=0.2, cex=1.2)
+ )

Error in plot(allEffects(tox.urea), ylab = "Probability (Urea)", xlab = "Social
class of mother", : error in evaluating the argument 'x' in selecting a method
for function 'plot': Error: could not find function "allEffects"

```

From Figure ??, it can be seen that the prevalence of these symptoms has a possibly complex relation to social class and smoking. However, the mosaic for these predictors in Figure 8.24 has shown us that several of the class-smoking categories are quite small (particularly heavy smokers in Classes 1 and 2) so the response effects for these classes will be poorly estimated. Taking this into account, we suspect that protein urea varies with social class, but not with smoking, while the prevalence of hypertension may truly vary with neither, just one, or both of these predictors.

Fitting models

The plots shown so far in this example are all essentially *data-based*, in that they use the observed frequencies or transformations of them and don't allow for a simpler view, based on a reasonable model. That is, abbreviating the table variables by their initial letters, the plots in Figure 8.28 and Figure ?? are plots of the saturated model, [CSHU] that fits perfectly, but with the data transformed for each 2×2 subtable to the log odds ratio and the two log odds for *hyper* and *urea*.

The bivariate logistic model fit by *vglm()* still applies when there are two or more predictors; however, like other multivariate response models, it doesn't easily allow the logits to depend on *different* predictor terms. To illustrate this, we first transform the *Toxaemia* to a 15×4 data frame in the form required by *vglm()*.

```

> tox.tab <- xtabs(Freq~class + smoke + hyper + urea, Toxaemia)
> toxaemia <- t(matrix(aperm(tox.tab), 4, 15))
> colnames(toxaemia) <- c("hu", "hU", "Hu", "HU")
> rowlabs <- expand.grid(smoke=c("0", "1-19", "20+"), class=factor(1:5))
> toxaemia <- cbind(toxaemia, rowlabs)
> head(toxaemia)

  hu hU Hu HU smoke class
1 286 21 82 28    0    1
2  71  5 24  5   1-19    1
3 13  0  3  1   20+    1
4 785 34 266 50    0    2
5 284 17 92 13   1-19    2
6 34  3 15  0   20+    2

```

In the model specification for `vglm()`, the `zero` argument in `binom.or()` allows any one or more of the two log odds and log odds ratio to be fit as a constant (intercept-only) in Eqn. (8.29). However, in that equation, the predictors x_1, x_2, x_{12} , must be the *same* in all three submodels. For example, the model `tox.vgml1` below uses main effects of `class` and `smoke` in both models for the logits, and `zero=3` for a constant log odds ratio.

```
> tox.vgml1 <- vglm(cbind(hu, hU, Hu, Hu) ~ class + smoke,
+                      binom2.or(zero=3), data=toxaemia)
> coef(tox.vgml1, matrix=TRUE)

      logit(mu1) logit(mu2) log(oratio)
(Intercept) -0.50853648 -1.2214518   2.7808
class2       0.18156457  0.0382046   0.0000
class3       0.06332765 -0.0087552   0.0000
class4      -0.02227055 -0.0031541   0.0000
class5      -0.00077172  0.0821863   0.0000
smoke1-19   -0.41298650 -0.2198673   0.0000
smoke20+    -0.30562472 -0.1245019   0.0000
```

Instead, when there are no quantitative predictors, and when the odds ratio is relatively constant (as here) it is easier to fit ordinary loglinear models than to use the bivariate logit formulation of the previous example. These allow the responses H and U to depend on the class-smoking combinations separately, by including the terms $[CSH]$ or $[CSU]$, respectively.

The minimal, null model, $[CS][H][U]$ fits the marginal association of the numbers in each class-smoking category, but asserts that the responses, H and U are independent, which we have already seen is contradicted by the data. We take $[CS][HU]$ as the baseline model (Model 1), asserting no relation between response and predictor variables, but associations within each set are allowed. These models are fit as shown below.

```
> # null model
> tox.glm0 <- glm(Freq ~ class*smoke + hyper + urea,
+                     data=Toxaemia, family=poisson)
> # baseline model: no association between predictors and responses
> tox.gml1 <- glm(Freq ~ class*smoke + hyper*urea,
+                     data=Toxaemia, family=poisson)
```

We proceed to fit a collection of other models, adding terms to allow more associations between the responses and predictors. Summary measures of goodness of fit and parsimony are shown in Table 8.3.

Table 8.3: Loglinear models, `tox.glm*`, fit to the Toxaemia data

{tab:toxmod}

Model	Terms	df	G^2	p-value	G^2/df	AIC	BIC	R^2
0	CS H U	43	672.85	0.0000	15.65	586.85	264.27	.
1	CS HU	42	179.03	0.0000	4.26	95.03	-220.04	0.000
2	CS HU SH CU	36	46.12	0.1203	1.28	-25.88	-295.94	0.742
3	CS CH CU HU SH CU	30	40.47	0.0960	1.35	-19.53	-244.58	0.774
4	CSH CU HU	24	26.00	0.3529	1.08	-22.00	-202.04	0.855
5	CSH CU SU HU	22	25.84	0.2588	1.17	-18.16	-183.20	0.856
6	CSH CSU HU	14	22.29	0.0729	1.59	-5.71	-110.74	0.875
7	CSH CSU SHU	12	15.65	0.2079	1.30	-8.35	-98.37	0.913
8	CSH CSU CHU SHU	8	12.68	0.1233	1.59	-3.32	-63.33	0.929
9	CSHU	0	0.00	0	0	0.00	0.00	1.000

```
> tox.glm2 <- update(tox.glm1, . ~ . + smoke*hyper + class*urea)
>
> tox.glm3 <- glm(Freq ~ (class + smoke + hyper + urea)^2,
+                     data=Toxaemia, family=poisson)
>
> tox.glm4 <- glm(Freq ~ class*smoke*hyper + hyper*urea + class*urea,
+                     data=Toxaemia, family=poisson)
>
> tox.glm5 <- update(tox.glm4, . ~ . + smoke*urea)
>
> tox.glm6 <- update(tox.glm4, . ~ . + class*smoke*urea)
>
> tox.glm7 <- update(tox.glm6, . ~ . + smoke*hyper*urea)
>
> tox.glm8 <- glm(Freq ~ (class + smoke + hyper + urea)^3,
+                     data=Toxaemia, family=poisson)
>
> tox.glm9 <- glm(Freq ~ (class + smoke + hyper + urea)^4,
+                     data=Toxaemia, family=poisson)
```

Model 2 adds the simple dependence of hypertension on smoking ($[SH]$) and that of urea on class ($[CU]$). Model 3 includes all two-way terms. In Model 4, hypertension is allowed to depend on both class and smoking jointly ($[CSH]$). In Model 5 an additional dependence of urea on smoking ($[SU]$) is included, while in Model 6 urea depends on class and smoking jointly ($[CSU]$).

None of these models contain three-way terms involving both H and U , so these models assume that the log odds ratio for hypertension given urea is constant over the explanatory variables. Recalling the conditional mosaics (Figure 8.25 and Figure 8.26), Models 7 and 8 add terms which allow the odds ratio to vary, first with smoking ($[SHU]$), then with class ($[CHU]$) as well. Finally, Model 9 is the saturated model, that fits perfectly.

How do we choose among these models? Model 2 is the smallest model whose deviance is non-significant. Models 4 and 5 both have a smaller ratio of G^2/df . For comparing nested models, we can also examine the change in deviance as terms are added (or dropped). Thus, going from Model 2 to Model 3 decreases the deviance by 5.65 on 6 df, while the step from Model 3 to Model 4 gives a decrease of 14.47, also on 6 df. These tests can be performed using `lrtest()` in the `lmtest` package, shown below for models `tox.glm1`-`tox.glm5`.

```
> library(lmtest)
> lmtest::lrtest(tox.glm1, tox.glm2, tox.glm3, tox.glm4, tox.glm5)

Likelihood ratio test

Model 1: Freq ~ class * smoke + hyper * urea
Model 2: Freq ~ class + smoke + hyper + urea + class:smoke + hyper:urea +
           smoke:hyper + class:urea
Model 3: Freq ~ (class + smoke + hyper + urea)^2
Model 4: Freq ~ class * smoke * hyper * urea + class * urea
Model 5: Freq ~ class + smoke + hyper + urea + class:smoke + class:hyper +
           smoke:hyper + hyper:urea + class:urea + smoke:urea + class:smoke:hyper
#Df LogLik Df Chisq Pr(>Chisq)
1 18    -260
2 24    -194   6 132.91      <2e-16 ***
3 30    -191   6   5.65      0.464
4 36    -184   6   14.47      0.025 *
5 38    -184   2    0.17      0.920
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The AIC and BIC statistics, balancing parsimony and goodness-of-fit, have their minimum value for Model 2, which we adopt here for this example.

Plotting model results

Whatever model is chosen, as a final step, it is important to determine what that model implies about the original research questions. Because our focus here is on the prevalence of each symptom, and

their association, it is helpful to graph the fitted logits and log odds ratios implied by the model, as was done in Figure 8.22 and Figure 8.23.

The presentation goal here is to produce plots showing the observed logits and log odds ratios as in Figure ?? and Figure 8.28, supplemented by lines showing these values according to the fitted model. In Example 8.12 we fit the bivariate logit model, for which the response functions were the desired logits and log odds. Here, where we have fit ordinary loglinear models, the observed and fitted logits can be calculated from the observed and fitted frequencies. The calculations require a bit of R calisthenics to arrange these into forms suitable for plotting.

As we did earlier, we first reshape the *Toxaemia* to wide format, as a 15×4 table of observed frequencies. Because there are now two predictor variables, we take care to include the levels of *smoke* and *class* as additional columns.

```
> # reshape to 15 x 4 table of frequencies
> tox.tab <- xtabs(Freq~class + smoke + hyper + urea, Toxaemia)
> toxaemia <- t(matrix(aperm(tox.tab), 4, 15))
> colnames(toxaemia) <- c("hu", "hU", "Hu", "HU")
> rowlabs <- expand.grid(smoke=c("0", "1-19", "20+"), class=factor(1:5))
> toxaemia <- cbind(toxaemia, rowlabs)
```

Applying `blogits()`, we get the observed logits and log odds ratios in `logitsTox`.

```
> # observed logits and log odds ratios
> logitsTox <- blogits(toxaemia[, 4:1], add=0.5)
> colnames(logitsTox)[1:2] <- c("logitH", "logitU")
> logitsTox <- cbind(logitsTox, rowlabs)
> head(logitsTox)

  logitH logitU    logOR smoke class
1 -1.02057 -1.9988  1.52679     0     1
2 -0.94261 -2.1665  1.07102   1-19    1
3 -1.02962 -2.1401  2.44854   20+    1
4 -0.95040 -2.5158  1.46196     0     2
5 -1.04699 -2.4983  0.86401   1-19    2
6 -0.86500 -2.5257 -1.14579   20+    2
```

The fitted frequencies are extracted using `predict(tox.glm2, type="response")` and then manipulated in a similar way to give `logitsFit`.

```
> # fitted frequencies, as a 15 x 4 table
> Fit <- t(matrix(predict(tox.glm2, type="response"), 4, 15))
> colnames(Fit) <- c("H\u00d5", "Hu", "hU", "hu")
> Fit <- cbind(Fit, rowlabs)
> logitsFit <- blogits(Fit[, 1:4], add=0.5)
> colnames(logitsFit)[1:2] <- c("logitH", "logitU")
> logitsFit <- cbind(logitsFit, rowlabs)
```

In tabular form, you can examine any of these components, for example, the log odds ratios from the fitted values shown below.

```
> matrix(logitsFit$logOR, 3, 5,
+         dimnames=list(smoke=c("0", "1-19", "20+"), class=1:5))

      class
smoke    1     2     3     4     5
  0    1.3588 1.3638 1.3675 1.3643 1.3582
  1-19 1.3582 1.3678 1.3683 1.3674 1.3658
  20+ 1.2799 1.3471 1.3662 1.3622 1.3511
```

Finally, we can plot the observed values in `logitsTox` (as points) and the fitted values under

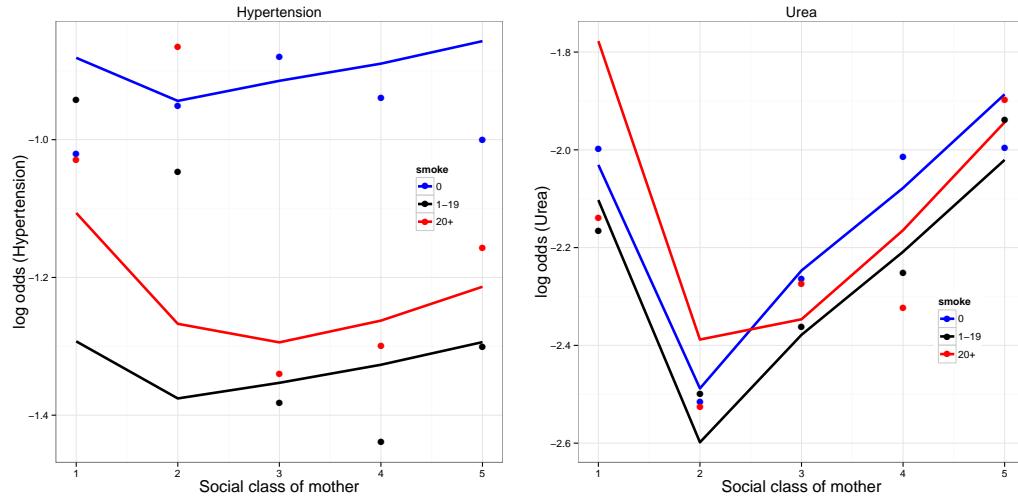


Figure 8.29: Observed (points) and fitted (lines) logits for the *Toxemia* data under Model 2.

{fig:tox-glm-logits1}

Model 2 in `logitsFit` (as lines), separately for the `logitH`, `logitU`, and `logOR` components. The code below uses `ggplot2` for the log odds of hypertension, and is repeated for urea and the log odds ratio. These graphs are shown in Figure 8.29 and Figure 8.30.

```
> ggplot(logitsFit, aes(x=as.numeric(class), y=logitH, color=smoke)) +
+   theme_bw() +
+   geom_line(size=1.2) +
+   scale_color_manual(values=c("blue", "black", "red")) +
+   ylab("log odds (Hypertension)") +
+   xlab("Social class of mother") +
+   ggtitle("Hypertension") +
+   theme(axis.title=element_text(size=16)) +
+   geom_point(data=logitsTox,
+             aes(x=as.numeric(class), y=logitH, color=smoke), size=3) +
+   theme(legend.position=c(0.85, .6))
```

According to this model, Figure 8.30 shows that the fitted log odds ratio is in fact nearly constant, while Figure 8.29 shows that the log odds for hypertension depends mainly on smoking (with a large difference of the non-smoking mothers from the rest) and that for protein urea depends mainly on social class.¹⁸

Yet, the great variability of the observed points around the fitted curves indicates that these relationships are not well-determined. Adding error bars showing the standard error around each fitted point would indicate that the data conforms as closely to the model as can be expected, given the widely different sample sizes. However, this would make the plots more complex, and so was omitted here. In addition to showing the pattern of the results according to the fitted model, such graphs also help us to appreciate the model's limitations.

△

¹⁸Some possible enhancements to these graphs include (a) plotting on the scale of probabilities or including a right vertical axis showing corresponding probabilities; (b) using the same vertical axis limits for the two graphs for direct comparison.

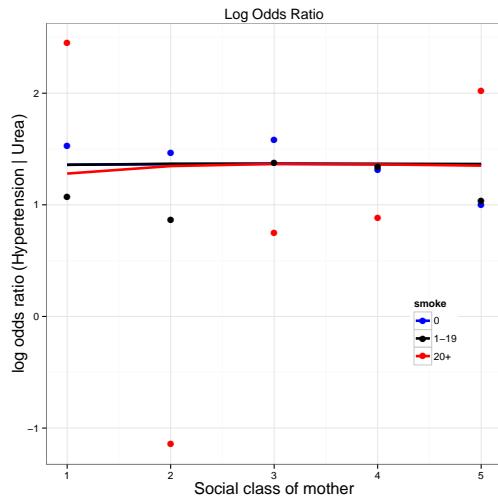


Figure 8.30: Observed (points) and fitted (lines) log odds ratios for the *Toxemia* data under Model 2.

8.10 Chapter summary

- Loglinear models provide a comprehensive scheme to describe and understand the associations among two or more categorical variables. It is helpful to think of these as discrete analogs of ANOVA models, or of regression models, where the log of cell frequency is modelled as a linear function of predictors.
- Loglinear models typically make no distinction between response and explanatory variables. When one variable *is* a response, however, any logit model for that response has an equivalent loglinear model. The logit form is usually simpler to formulate and test, and plots of the observed and fitted logits are easier to interpret.
- Models for square tables, with the same row and column categories are an important special case. For these and other structured tables, a variety of techniques provide the opportunity to fit models more descriptive than the independence model and more parsimonious than the saturated model.
- Standard loglinear models treat all variables as unordered factors. When one or more factors are ordinal, however, loglinear and logit models may be simplified by assigning quantitative scores to the levels of an ordered factor. Such models are often more sensitive and have greater power because they are more focused.
- In all these cases, the interplay between graphing and fitting is important in arriving at an understanding of the relationships among variables and an adequate descriptive model which is faithful to the details of the data.
- When there are several categorical responses, along with one or more explanatory variables, some special forms of loglinear and logit models may be used to separate the marginal dependence of each response on the explanatory variables from the interdependence among the responses.

8.11 Further reading

8.12 Lab exercises

Exercise 8.1 Example 8.8 presented an analysis of the data on visual acuity for the subset of women in the *VisualAcuity* data. Carry out a parallel analysis of the models fit there for the men in this data set, given by:

```
> data("VisualAcuity", package="vcd")
> men <- subset(VisualAcuity, gender=="male", select=-gender)
```

{lab:8.2}

Exercise 8.2 Table 8.4 gives a 4×4 table of opinions about premarital sex and whether methods of birth control should be made available to teenagers aged 14–16 from the 1991 General Social Survey (Agresti, 2013, Table 10.3). Both variables are ordinal, and their grades are represented by the case of the row and column labels.

Table 8.4: Opinions about premarital sex and availability of teenage birth control. *Source:* Agresti (2013, Table 10.3)

Premarital sex	Birth control			AGREE
	DISAGREE	disagree	agree	
WRONG	81	68	60	38
Wrong	24	26	29	14
wrong	18	41	74	42
OK	36	57	161	157

- (a) Fit the independence model to these data using `loglm()` or `glm()`.
- (b) Make a mosaic display showing departure from independence and describe verbally the pattern of association.
- (c) Treating the categories as equally spaced, fit the $L \times L$ model of uniform association, as in Section 8.6. Test the difference against the independence model with a likelihood-ratio test.
- (d) Fit the RC(1) model with `gnm()`, and test the difference of this against the model of uniform association.
- (e) Write a brief summary of these results, including plots useful for explaining the relationships in this data set.

{lab:8.3}

Exercise 8.3 The data set *gss8590* in *logmult* gives a $4 \times 5 \times 4$ table of education levels and occupational categories for the four combinations of gender and race from the General Social Surveys, 1985–1990 as reported by Wong (2001, Table 2). Wong (2010, Table 2.3B) later used the subset pertaining to women to illustrate RC(2) models. This data is created below as *Women.tab*, correcting an inconsistency to conform with the 2010 table.

```
> data(gss8590, package="logmult")

Error in find.package(package, lib.loc, verbose = verbose): there is no package
called 'logmult'

> Women.tab <- margin.table(gss8590[, , c("White Women", "Black Women")], 1:2)

Error in margin.table(gss8590[, , c("White Women", "Black Women")], 1:2): object
'gss8590' not found

> Women.tab[2, 4] <- 49
```

{sec:loglin-reading}

{lab:8.11-loglin-lab}

```
Error in Women.tab[2, 4] <- 49: object 'Women.tab' not found
> colnames(Women.tab) [5] <- "Farm"
Error in colnames(Women.tab) [5] <- "Farm": object 'Women.tab' not found
```

- (a) Fit the independence model, and also the RC(1) and RC(2) models using `rc()` with marginal weights, as illustrated in Example 8.7. Summarize these statistical tests in a table.
- (b) Plot the solution for the RC(2) model with 68% confidence ellipses. What verbal labels would you use for the two dimensions?
- (c) Is there any indication that a simpler model, using integer scores for the row (Education) or column (Occupation) categories or both might suffice? If so, fit the analogous column effects, row effects or $L \times L$ model, and compare with the models fit in part (a).

```
> #detach(package:corrplot)
> detach(package:VGAM)
> #detach(package:logmult)
> #remove(list=objects(pattern="\\.tab/\\.df/\\.fit"))
> .locals$ch08 <- setdiff(ls(), .globals)
> #.locals$ch08
> #remove(list=.locals$ch08[sapply(.locals$ch03, function(n) !is.function(get(n)) )])
```


Chapter 9

Generalized linear models

{ch:glm}

Generalized linear models extend the familiar linear models of regression and ANOVA to include counted data, frequencies, and other data for which the assumptions of independent, normal errors are not reasonable. We rely on the analogies between ordinary and generalized linear models (GLMs) to develop visualization methods to explore the data, display the fitted relationships and check model assumptions. The main focus of this chapter is on models for count data.

In one word, to draw the rule from experience, one must generalize; this is a necessity that imposes itself on the most circumspect observer.

Henri Poincaré, *The Value of Science: Essential Writings of Henri Poincaré*

In the modern history of statistics, most developments occur incrementally, with small additions to existing models and theory that extend their range and applicability to new problems and data. Occasionally, there is a major synthesis that unites a wide class of existing methods in a general framework and provides opportunities for far greater growth.

A prime example is the theory of generalized linear models, introduced originally by Nelder and Wedderburn (1972), that extended the familiar (classical) linear models for regression and ANOVA to include related models, such as logistic regression and logit models (described in Chapter 7) and loglinear models (described in Chapter 8) and other variations as “families” within a single general system.

This approach has proved attractive because it: (a) integrates many familiar statistical models in a general theory where they are just special cases; (b) provides the basis for extending these and developing new models within the same or similar framework; (c) simplifies the implementation of these models in software, since the same algorithm can be used for estimation, inference and assessing model adequacy for all generalized linear models.

Section 9.1 gives a brief sketch of the GLM framework. The focus of this book is on visualization methods for categorical data, and the two important topics concern models and methods for binomial response data and for count data. The first of these, was described extensively in Chapter 7, with extensions to multinomial data (Section 7.6) and there is little to add here, except for changes in notation.

GLM models for count data, however, provide the opportunity to extend the scope of these methods beyond what was covered in Chapter 8, and this topic is introduced in Section 9.2. The GLM framework also provides the opportunity to deal with common problems of overdispersion (Section 9.3) and an overabundance of zero counts (Section 9.3), giving some new models and

visualization methods that help to understand such data in greater detail. Section 9.6 illustrates other graphical methods for diagnostic model checking, some of which were introduced in earlier chapters. Finally, Section 9.7 outlines some simple extensions of these models to handle multivariate responses.

9.1 Components of Generalized Linear Models

{glm:components}

The motivation for the **generalized linear model** (GLM) and its structure are most easily seen by considering the classical linear model,

$$y_i = \mathbf{x}_i^\top \boldsymbol{\beta} + \epsilon_i$$

where y_i is the response variable for case $i, i = 1, \dots, n$, \mathbf{x}_i is the vector of explanatory variables or regressors, $\boldsymbol{\beta}$ is the vector of model parameters, and the ϵ_i are random errors. In the classical linear model, the ϵ_i are assumed to (a) have constant variance, σ_ϵ^2 , (b) follow a normal (Gaussian) distribution (conditional on \mathbf{x}_i), (c) be independent across observations.

Thus, Nelder and Wedderburn (1972) generalized this Gaussian linear model to consist of the following three components, by relaxing assumptions (a) and (b) above:¹

random component The conditional distribution of the $y_i | \mathbf{x}_i$, with mean $\mathcal{E}(y_i) = \mu_i$. Under classical assumptions, this is independent, normal with constant variance σ^2 , i.e., $y_i \stackrel{\text{iid}}{\sim} N(\mu_i, \sigma^2)$. In the GLM, the probability distribution of the y_i can be any member of the **exponential family**, including the normal, Poisson, binomial, gamma and others. Subsequent work has extended this framework to include multinomial distributions and some non-exponential families such as the negative binomial distribution.

systematic component The idea that the predicted value of y_i itself is a linear combination of the regressors is replaced by that of a **linear predictor**, η , that captures this aspect of linear models,

$$\eta_i = \mathbf{x}_i^\top \boldsymbol{\beta}$$

link function The connection between the mean of the response, μ_i , and the linear predictor, η_i , is specified by the **link function**, $g(\bullet)$, giving

$$g(\mu_i) = \eta_i = \mathbf{x}_i^\top \boldsymbol{\beta}$$

The link function $g(\bullet)$ must be both *smooth* and *monotonic*, meaning that it is one-to-one, so an inverse transformation, $g^{-1}(\bullet)$ exists,

$$\mu_i = g^{-1}(\eta_i) = g^{-1}(\mathbf{x}_i^\top \boldsymbol{\beta})$$

which allows us to obtain and plot the predicted values on their original scale. The link function captures the familiar idea that linear models are often estimated with a transformation of the response, such as $\log(y_i)$ for a frequency variable or $\text{logit}(y_i)$ for a binomial variable. The inverse function $g^{-1}(\bullet)$ is also called the **mean function**.

Some commonly used link functions are shown in Table 9.1. Some of these link functions have restrictions on the range of y_i to which they can be applied. For example, the square-root and log links apply only to non-negative and positive values respectively. The last four link functions in this table are for binomial data, where y_i represents the observed proportion of successes in n_i independent trials, and thus the mean μ_i represents the probability of success (symbolized by π_i in Chapter 7). Binary data are the special case where $n_i = 1$.

¹The remaining assumption of independent observations is relaxed in **generalized linear mixed models** (GLMMs), in which random effects to account for non-independence are added to the linear predictor. This allows the modeling of correlated (responses of family members), clustered (residents in different communities) or hierarchical data (patients within hospitals within regions). See: McCulloch and Neuhaus (2005) ... **TODO: other references?**

{tab:link-funcs}

Table 9.1: Common link functions and their inverses used in generalized linear models

Link name	Function: $\eta_i = g(\mu_i)$	Inverse: $\mu_i = g^{-1}(\eta_i)$
identity	μ_i	η_i
square-root	$\sqrt{\mu_i}$	η_i^2
log	$\log_e(\mu_i)$	$\exp(\eta_i)$
inverse	μ_i^{-1}	η_i^{-1}
inverse-square	μ_i^{-2}	$\eta_i^{-1/2}$
logit	$\log_e \frac{\mu_i}{1-\mu_i}$	$\frac{1}{1+\exp(-\eta_i)}$
probit	$\Phi^{-1}(\mu_i)$	$\Phi^{-1}(\eta_i)$
log-log	$-\log_e[-\log_e(\mu_i)]$	$\exp[-\exp(-\eta_i)]$
comp. log-log	$\log_e[-\log_e(1-\mu_i)]$	$1 - \exp[-\exp(\eta_i)]$

9.1.1 Variance functions

The GLM has the additional property that, for distributions in the exponential family, the conditional variance of $y_i | \eta_i$ is a known function, $\mathcal{V}(\mu_i)$ of the mean and possibly one other parameter called the **scale parameter** or **dispersion parameter**, ϕ . Some commonly used distributions in the exponential family and their variance functions are shown in Table 9.2.

Table 9.2: Common distributions in the exponential family used with generalized linear models and their canonical link and variance functions

{tab:exp-families}

Family	Notation	Canonical link	Range of y	Variance function, $\mathcal{V}(\mu \eta)$
Gaussian	$N(\mu, \sigma^2)$	identity: μ	$(-\infty, +\infty)$	ϕ
Poisson	$\text{Pois}(\mu)$	$\log_e(\mu)$	$0, 1, \dots, \infty$	μ
Negative-Binomial	$\text{NBin}(\mu, \theta)$	$\log_e(\mu)$	$0, 1, \dots, \infty$	$\mu + \mu^2/\theta$
Binomial	$\text{Bin}(n, \mu)/n$	logit(μ)	$\{0, 1, \dots, n\}/n$	$\mu(1 - \mu)/n$
Gamma	$G(\mu, \nu)$	μ^{-1}	$(0, +\infty)$	$\phi\mu^2$
Inverse-Gaussian	$IG(\mu, \nu)$	μ^2	$(0, +\infty)$	$\phi\mu^3$

- In the classical Gaussian linear model, the conditional variance is constant, $\phi = \sigma_\epsilon^2$.
- In the Poisson family, $\mathcal{V}(\mu_i) = \mu_i$ and the dispersion parameter is fixed at $\phi = 1$. In practice, it is common for count data to exhibit **overdispersion**, meaning that $\mathcal{V}(\mu_i) > \mu_i$. One way to correct for this is to extend the GLM to allow the dispersion parameter to be estimated from the data, giving what is called the **quasi-Poisson** family, with $\mathcal{V}(\mu_i) = \hat{\phi}\mu_i$.
- Similarly, for binomial data, the variance function is $\mathcal{V}(\mu_i) = \mu_i(1 - \mu_i)/n_i$, with ϕ fixed at 1. Overdispersion often results from failures of the assumptions of the binomial model: supposedly independent observations may be correlated or clustered and the probability of success may not be constant, or vary with unmeasured or unmodeled variables.
- The gamma and inverse-Gaussian families are distributions useful for modeling a continuous and positive response variable with no upper bound (e.g., reaction time). They both have the

property that conditional variance increases with the mean, and for the inverse-Gaussian, variance increases at a faster rate. Their dispersion parameters ϕ are simple functions of their intrinsic “shape” parameters, indicated as ν in the table.

The important points from this discussion are that the GLM together with the exponential family of distributions:

- provide for simple, linear relations between the response and the predictors via the link function and the linear predictor.
- allows a very flexible relationship between the mean and conditional variance to be specified in terms of a set of known families.
- incorporates a dispersion parameter ϕ that in some cases can be estimated or tested for departure from that entailed in a given family.
- has allowed further extensions of this framework outside the exponential family, ranging from simple adjustments for statistical inference (“quasi” families, adjusted “sandwich” covariances) to separate modeling of the variance relation to the predictors.

Further details of generalized linear models are beyond the scope of this book, but the interested reader should consult Fox (2008, §15.3) and Agresti (2013, Ch. 4) for a comprehensive treatment.

9.1.2 Hypothesis tests for coefficients

GLMs are fit using maximum likelihood estimation, and implemented in software using an iterative algorithm known as *iteratively weighted least squares* that generalizes the least squares method for classical linear models. This provides estimates $\hat{\beta}$ of the model coefficients for the predictors in x , as well as an estimated asymptotic (large sample) variance matrix of $\hat{\beta}$, given by

$$\mathcal{V}(\hat{\beta}) = \phi(\mathbf{X}^\top \mathbf{W} \mathbf{X}) \quad (9.1)$$

where \mathbf{W} is a diagonal matrix of weights computed in the final iteration. In the standard Poisson GLM, the weight matrix is $\mathbf{W} = \text{diag}(\hat{\mu})$ and $\phi = 1$ is assumed.

Asymptotic standard errors, $\text{se}(\hat{\beta}_j)$, for the coefficients are then the square roots of the diagonal elements of $\mathcal{V}(\hat{\beta})$, and tests of hypotheses regarding an individual coefficient, e.g., $H_0 : \beta_j = 0$, can be carried out using the Wald test statistic, $z_j = \hat{\beta}_j / \text{se}(\hat{\beta}_j)$. When the null hypothesis is true, z_j has a standard normal $\mathcal{N}(0, 1)$ distribution, providing p -values for significance tests.²

More generally, we can test any *linear hypothesis*, of the form $H_0 : \mathbf{L}\beta = \mathbf{c}$, where \mathbf{L} is a constant hypothesis matrix of size $h \times p$ giving h linear combinations of the coefficients, to be tested for equality with the constants in \mathbf{c} , typically taken as $\mathbf{c} = \mathbf{0}$. The test statistic is the Wald chi-square,

$$Z^2 = (\mathbf{L}\hat{\beta} - \mathbf{c})^\top [\mathbf{L}\mathcal{V}(\hat{\beta})\mathbf{L}^\top]^{-1} (\mathbf{L}\hat{\beta} - \mathbf{c}) \quad (9.2)$$

which has a χ^2 distribution on h degrees of freedom.³

For example, to test the hypothesis that all of $\beta_1 = \beta_2 = \beta_3 = 0$ in a model with three predictors, you can use

$$\mathbf{L} = \begin{bmatrix} 0 & 1 & 0 & 0 \\ 0 & 0 & 1 & 0 \\ 0 & 0 & 0 & 1 \end{bmatrix} = \begin{bmatrix} \mathbf{0} & \mathbf{I} \end{bmatrix}, \quad \mathbf{c} = \begin{pmatrix} 0 \\ 0 \\ 0 \end{pmatrix}$$

²Wald tests are sometimes carried out using z^2 , which has an equivalent χ^2_1 distribution with 1 degree of freedom.

³When a dispersion parameter ϕ has been estimated from the data, it is common to use an F -test, using the statistic $F = Z^2/h$, with h and $n - p$ degrees of freedom.

Similarly, to test the hypothesis that $\beta_1 = \beta_2$ in the same model, you can use $L = [0, 1, -1, 0]$ and $c = [0]$.⁴

In R, such tests are most conveniently carried out using `linearHypothesis()` in the `car` package. The hypothesis matrix L can be supplied as a numeric matrix, or more conveniently, the hypothesis can be specified symbolically as a character vector of the names of the coefficients involved in each row of L . For example, the first hypothesis test above could be specified using the vector `c("x1=0", "x2=0", "x3=0")` and the test of equality as `"x1-x2=0"`.

9.1.3 Goodness-of-fit tests

{sec:glm-goodfit}

The basic ideas for testing goodness-of-fit were discussed in Section 8.3.2 in connection with log-linear models for contingency tables. As before, these assess the overall performance of a model in reproducing the data. The commonly used measures include the Pearson chi-square and likelihood-ratio deviance statistics, which can be seen as weighted sums of residuals. We re-state these test statistics here in the wider context of the GLM.

Let $y_i, i = 1, 2, \dots, n$ be the response and $\hat{\mu}_i = g^{-1}(\mathbf{x}_i^T \hat{\beta})$ the fitted mean using the estimated coefficients, having estimated variance $\hat{\omega}_i = V(\hat{\mu}_i | \eta_i)$ as in Table 9.2. Then the normalized squared residual for observation i is $(y_i - \hat{\mu}_i)^2 / \hat{\omega}_i$, and the Pearson statistic is

$$X_P^2 = \sum_{i=1}^n \frac{(y_i - \hat{\mu}_i)^2}{\hat{\omega}_i}. \quad (9.3) \quad \text{(eq:pearson)}$$

In the GLM for count data, the main focus of this chapter, the Poisson family sets $\omega = \mu$ with the dispersion parameter fixed at $\phi = 1$.

The **residual deviance** statistic, as in logistic regression and loglinear models is defined as twice the difference between the maximum possible log-likelihood for the *saturated model* that fits perfectly and maximized log-likelihood for the fitted model. The deviance can be defined as

$$D(\mathbf{y}, \hat{\boldsymbol{\mu}}) \equiv 2[\log_e \mathcal{L}(\mathbf{y}; \mathbf{y}) - \log_e \mathcal{L}(\mathbf{y}; \hat{\boldsymbol{\mu}})]$$

For classical linear models under normality, the deviance is simply the residual sum of squares, $\sum_i^n (y_i - \hat{\mu}_i)$. This has led to the deviance being taken in the GLM framework as a generalization of the sum of squares used in ANOVA, and hence, an analogous **analysis of deviance** to carry out tests for individual terms in GLMs, or to compare nested models.

In R, `anova(mod)` for the "glm" object `mod` gives *sequential* ("Type I") tests of successive terms in a model, while `Anova()` in the `car` package gives the more generally useful "Type II" (and "Type III") *partial* tests, that assess the additional contribution of each term above all others, taking marginality into account.

For Poisson models with a log link giving $\boldsymbol{\mu} = \exp(\mathbf{x}^T \boldsymbol{\beta})$, the deviance takes the form⁵

$$D(\mathbf{y}, \hat{\boldsymbol{\mu}}) = 2 \sum_{i=1}^n \left[y_i \log_e \left(\frac{y_i}{\hat{\mu}_i} \right) - (y_i - \hat{\mu}_i) \right] \quad (9.4) \quad \text{(eq:pois-deviance)}$$

For a GLM with p parameters, both the Pearson and residual deviance statistics follow approximate χ^2_{n-p} distributions with $n - p$ degrees of freedom.

⁴Such a test is only sensible if the predictors \mathbf{x}_1 and \mathbf{x}_2 are on the same scale, so their coefficients are commensurable.

⁵In the context of the loglinear models discussed in Section 8.3.2, this is also referred to as the likelihood-ratio G^2 statistic.

9.1.4 Comparing non-nested models

The flexibility of the GLM and its extensions allows us to fit models to the same data using different families and different link functions, and to fit models that allow for overdispersion (Section 9.3) or that make special provisions for zero counts (Section 9.4). One price paid for this additional versatility is that standard LR tests and F tests (such as provided by `anova()` and `linearHypothesis()` in the `car` package) do not apply to models that are not nested, that is, where one model cannot be represented as a restricted, special case of another.

For models estimated by maximum likelihood, one general route to comparing non-nested models is through the AIC information criterion proposed initially by Akaike (1973) and the related BIC criterion (Schwartz, 1978) based on the fitted log-likelihood function.

$$\text{AIC} = -2 \log_e \mathcal{L} + 2k \quad (9.5)$$

$$\text{BIC} = -2 \log_e \mathcal{L} + \log_e(n)k \quad (9.6)$$

As noted in Section 8.3.2, these both penalize models with larger k , the number of parameters in the model, with BIC adding a greater penalty with larger sample size. However, because they are based only on the maximized log-likelihood, they are agnostic as to whether models are nested or not, and give comparable results (lower is better) provided the same observations have been used in all models.

In R, these results are given for a collection of models by the generic functions `AIC()` and `BIC()`; these can be calculated for any model for which `logLik()` and (for BIC) `nobs()` methods exist. The `vcdExtra` function `LRstats()` is a convenient wrapper for these methods.

AIC and BIC do not give significance tests for assessing whether one model is significantly “better” than another. One test that *does* this was proposed by Vuong (1989), unsurprisingly called **Vuong’s test**. The test is based on comparing the predicted probabilities or the pointwise log-likelihoods of the two models, and tests the null hypothesis that each is equally close to the saturated model, against the alternative that one model is closer.

For two such models, let $f_1(y_i | \mathbf{x}_i, \boldsymbol{\theta}_1)$ be the density function under model 1, with parameters $\boldsymbol{\theta}_1$ and similarly $f_2(y_i | \mathbf{x}_i, \boldsymbol{\theta}_2)$ under model 2 with parameters $\boldsymbol{\theta}_2$, where $f_1(\bullet)$ and $f_2(\bullet)$ need not be the same. Vuong’s test compares these based on the observation-wise log-likelihood ratios,

$$\ell_i = \log_e \left(\frac{f_1(y_i | \mathbf{x}_i, \hat{\boldsymbol{\theta}}_1)}{f_2(y_i | \mathbf{x}_i, \hat{\boldsymbol{\theta}}_2)} \right)$$

The test statistic is

$$V = \frac{\bar{\ell} - \text{penalty}}{\sqrt{n}s_\ell}$$

where $\bar{\ell}$ is the mean of the ℓ_i , s_ℓ is their variance, and penalty is an adjustment for model parsimony, typically taken as $\log(n)(k_1 - k_2)/2$ when model 1 has k_1 parameters in $\boldsymbol{\theta}_1$ and model 2 has k_2 parameters in $\boldsymbol{\theta}_2$.

The test statistic V has an asymptotic normal $N(0, 1)$ distribution, and is directional, with large positive values favoring model 1, and large negative values favoring model 2. This test is implemented as the `vuong()` function in the `pscl` package.

9.2 GLMs for count data

{sec:glm-count}

The prototypical GLM for count data, where the response y_i takes on non-negative values $0, 1, 2, \dots$, uses the Poisson family with the log link. We used this model extensively throughout all of Chapter 8. There the focus was on the special case of the loglinear model applied largely to contingency

{sec:glm-nonnest}

tables, where the loglinear model could be seen as a fairly direct extension of ANOVA models for a quantitative response applied to the log of cell frequency.

The advantage there was that models for two-way, three-way and by implication n -way tables could be discussed and illustrated using notation and graphs that separated the parameters and effects for one-way terms (“main effects”), two-way terms (“simple associations”) and higher-way terms (“conditional associations”).

The disadvantage is that these models as formulated there do not easily accommodate general quantitative predictors and were limited to the log link and the Poisson family. For example, the models discussed in Section 8.6 for ordinal variables allow one or more table factors to be assigned quantitative scores or have such scores estimated from the data, as in RC() models (Section 8.6.2). Yet, the contingency table approach for loglinear models breaks down if there are continuous predictors, and count data often exhibits features that make the equivalent Poisson regression model unsuitable or incomplete. We consider some extended models here.

{ex:phdpubs1}

EXAMPLE 9.1: Publications of PhD candidates

In Example 3.24 we considered the distribution of the number of publications by PhD candidates in their last three years of study, but without taking any available predictors into account. For these data, a simple calculation shows why the Poisson distribution is unsuitable (for the marginal distribution), because the variance is 2.19 times the mean.

```
> data("PhdPub", package="vcdExtra")
> with(PhdPub, c(mean=mean(articles), var=var(articles),
+ ratio=var(articles)/mean(articles)))
mean      var      ratio
1.6929  3.7097  2.1914
```

The earlier example showed rootograms (in Figure 3.28) of the number of articles, but here it is useful to consider some more basic exploratory displays. A basic barplot of the frequency distribution of number of articles published is shown in the left panel of Figure 9.1. A quick look indicates that the distribution is highly skewed and there is a large number of counts of zero.

Another problem is that the frequencies of 0–2 articles account for over 75% of the total, so that the frequencies of the larger counts get lost in the display. The rootogram corrects for this by plotting frequency on the square-root scale. However, because we are contemplating a model with a log link, the same goal can be achieved by plotting log of frequency, as shown in the right panel of Figure 9.1. To accommodate the zero frequencies, the plot shows $\log(\text{Frequency}+1)$, avoiding errors from $\log(0)$. It can be seen that log frequency decreases steadily up to 7 articles and then levels off approximately.

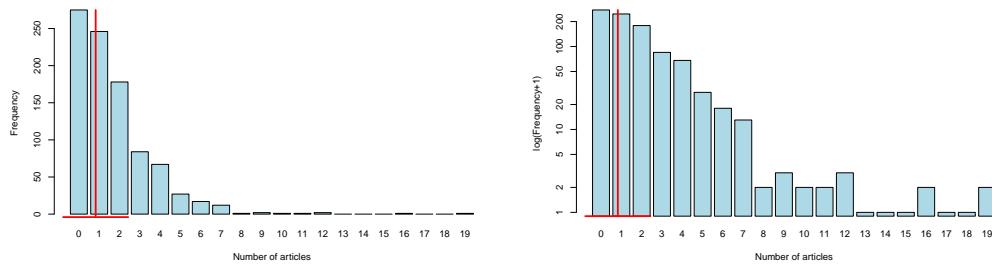


Figure 9.1: Barplots showing the frequency distribution of number of publications by PhD candidates. Left: raw scale; Right: a log scale makes the smaller counts more visible. The vertical red lines show the mean and horizontal lines show $\text{mean} \pm 1$ standard deviation.

{fig:phdpubs-barplot}

These plots are produced as shown below. The frequency distribution of articles can be tabulated by `table()`, but there is a subtle wrinkle here: By default, `table()` excludes the values of `articles` that do not occur in the data (zero frequencies). To include all values in the entire range, it is necessary to treat `articles` as a factor with levels 0:19.

```
> art.fac <- factor(PhdPubs$articles, levels=0:19) # include zero frequencies
> art.tab <- table(art.fac)
> art.tab

art.fac
 0   1   2   3   4   5   6   7   8   9   10  11  12  13  14  15  16  17  18  19 
275 246 178 84 67 27 17 12 1  2  1  1  2  13 0  0  1  0  1  1  0  1 
```

Then, the basic plot on the frequency scale is created using `barplot()`, and some annotations showing the mean and a one standard deviation interval can be added using standard plotting tools.

```
> barplot(art.tab, xlab="Number of articles", ylab="Frequency",
+           col="lightblue")
> abline(v=mean(PhdPubs$articles), col="red", lwd=3)
> ci <- mean(PhdPubs$articles)+c(-1,1) * sqrt(var(PhdPubs$articles))
> lines(x=ci, y=c(-4, -4), col="red", lwd=3, xpd=TRUE)
```

Similarly, the plot on the log scale in the right panel of Figure 9.1 is produced with `barplot()`, but using `art.tab+1` to start frequency at one and `log="y"` to scale the vertical axis to log.

```
> barplot(art.tab+1, ylab="log(Frequency+1)", xlab="Number of articles",
+           col="lightblue", log="y")
```

Other useful exploratory plots for count data include boxplots of the response (on a log scale) and scatterplots against continuous predictors, where jittering the response is often necessary to avoid overplotting and a smooth nonparametric curve can show possible non-linearity. The `log="y"` option is again handy, and the formula method allows adding a start value to the response. Figure 9.2 illustrates these ideas, for the factor `married` and the covariate `mentor`.

```
> boxplot(articles+1 ~ married, data=PhdPubs, log="y", varwidth=TRUE,
+           ylab="log(articles+1)", xlab="married", cex.lab=1.25)
> plot(jitter(articles+1) ~ mentor, data=PhdPubs, log="y",
+       ylab="log(articles+1)", cex.lab=1.25)
> lines(lowess(PhdPubs$mentor, PhdPubs$articles+1), col="blue", lwd=3)
```

It can be seen that the distribution of articles for married and non-married are quite similar, except that for the married students there are quite a few observations with a large number of publications. The relationship between `log(articles)` and `mentor` publications seems largely linear except possibly at the very low end. The large number of zero counts at the lower left corner stands out; this would not be seen without jittering.

Plots similar to those in Figure 9.2 can also be produced using `ggplot2` with greater flexibility, but perhaps greater effort to get the details right. One key feature is the use of `scale_y_log10()` to plot the response, and all other features on a log scale. The following code gives a plot similar to the right panel of Figure 9.2, but also plots a confidence band around the smoothed curve, and adds a linear regression line of `log(articles)` on `mentor` publications. This plot is not shown here, but it is a good exercise to reproduce it for yourself.

```
> ggplot(PhdPubs, aes(mentor, articles+1)) +
+   geom_jitter(position=position_jitter(h=0.05)) +
+   stat_smooth(method="loess", size=2, fill="blue", alpha=0.25) +
+   stat_smooth(method="lm", color="red", size=1.25, se=FALSE) +
+   scale_y_log10(breaks=c(1,2,5,10,20)) +
+   labs(y = "log (articles+1)", x="Mentor publications")
```

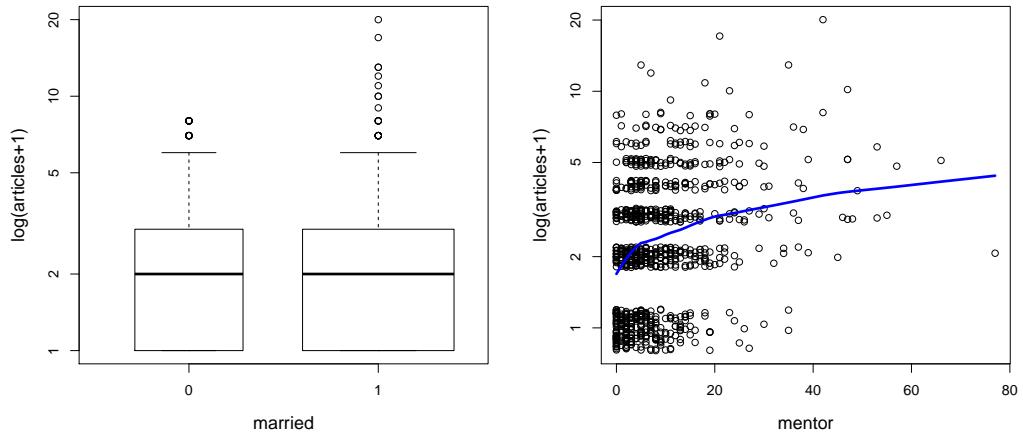


Figure 9.2: Exploratory plots for the number of articles in the PhdPubs data. Left: boxplots for married (1) vs. non-married (0); right: jittered scatterplot vs. mentor publications with a lowess smoothed curve.

To start analysis, we fit the Poisson model using all predictors—female, married, kid5, phdprestige, and mentor. As recorded in *PhdPubs*, female and married are both dummy (0/1) variables, and it slightly more convenient for plotting purposes to make them factors.

```
> PhdPubs <- within(PhdPubs, {
+   female <- factor(female)
+   married <- factor(married)
+ })
```

The model is fit as shown below and summarized using `summary()`, but with abbreviated output. **TODO:** `output.lines=9:24` works in *chapter.Rnw*, but not in *book.Rnw*. Why???

```
> phd.pois <- glm(articles ~ ., data=PhdPubs, family=poisson)
> summary(phd.pois)

Call:
glm(formula = articles ~ ., family = poisson, data = PhdPubs)

Deviance Residuals:
    Min      1Q  Median      3Q     Max 
-3.488  -1.538  -0.365   0.577   5.483 

Coefficients:
            Estimate Std. Error z value Pr(>|z|)    
(Intercept) 0.26562   0.09962   2.67   0.0077 **  
female1     -0.22442   0.05458  -4.11   3.9e-05 *** 
married1     0.15732   0.06125   2.57   0.0102 *    
kid5        -0.18491   0.04012  -4.61   4.0e-06 *** 
phdprestige  0.02538   0.02527   1.00   0.3153    
mentor       0.02523   0.00203  12.43 < 2e-16 *** 
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for poisson family taken to be 1)
```

```
Null deviance: 1817.4 on 914 degrees of freedom
Residual deviance: 1633.6 on 909 degrees of freedom
AIC: 3313

Number of Fisher Scoring iterations: 5
```

Significance tests for the individual coefficients show that all are significant, except for `phdprestige`. We ignore this here, and continue to interpret and extend the full main effects model.⁶

The estimated coefficients β for the predictors are shown below. Recall that using the log link means, for example, that being married increases the log of the expected number of articles published by 0.157, holding all other predictors constant. Each additional child of age 5 or less decreases this by 0.185.

```
> round(cbind(beta=coef(phd.pois),
+               expbeta=exp(coef(phd.pois)),
+               pct=100*(exp(coef(phd.pois))-1)), 3)

      beta expbeta     pct
(Intercept) 0.266   1.304 30.425
female1     -0.224   0.799 -20.102
married1     0.157   1.170 17.037
kid5        -0.185   0.831 -16.882
phdprestige  0.025   1.026  2.570
mentor       0.025   1.026  2.555
```

It is somewhat easier to interpret the exponentiated coefficients, $\exp(\beta)$ as multiplicative effects on the expected number of articles and convert these to percentage change, again holding other predictors constant. For example, expected publications by married candidates are 1.17 times that of non-married, a 17% increase, while each additional child multiplies articles by 0.831, a 16.88% decrease.

Alternatively, we recommend visual displays for model interpretation, and effect plots do well in most cases, as shown in Figure ???. For a Poisson GLM, an important feature is that the response is plotted on the log scale, so that effects in the model appear as linear functions, while the values of the response (number of articles) are labeled on their original scale, facilitating interpretation. The confidence bands and error bars give 95% confidence intervals around the fitted effects.

```
> library(effects)

Error: package 'effects' was built before R 3.0.0: please re-install it

> plot(allEffects(phd.pois), band.colors="blue", lwd=3,
+       ylab="Number of articles", main="")

Error in plot(allEffects(phd.pois), band.colors = "blue", lwd = 3, ylab = "Number of articles", : error in evaluating the argument 'x' in selecting a method for function 'plot': Error: could not find function "allEffects"
```

In Figure ?? we can see the decrease in published articles with number of young children, but also that the confidence band gets wider with increasing children. The predicted effect here of number of publications by the student's mentor is more dramatic, particularly for those whose mentor were truly prolific.

You should note that the panels for the predictors in Figure ?? are scaled individually for the range of the fitted main effects. This is often a sensible default and all predictors except `mentor`

⁶It is usually less harmful to include a non-significant predictor, (which in any case may be a variable useful to control, as `phdprestige` here), than to omit a potentially important predictor, or worse—to fail to account for an important interaction.

give a similar range here. To make all of these plots strictly comparable, provide a `ylim` argument, giving the range of the response on the log scale, as below (but not shown here).

```
> plot(allEffects(phd.pois), band.colors="blue", ylim=c(0,log(10)))
```

All of the above is useful, but still leaves aside the question of how well the Poisson model fits the data. The output from `summary(phd.pois)` above showed that the Poisson model fits quite badly. The residual deviance of 1633.6 with 909 degrees of freedom is highly significant.



{ex:crabs1}

EXAMPLE 9.2: Mating of horseshoe crabs

Brockmann (1996) studied the mating behavior of female horseshoe crabs in the Gulf of Mexico. In the mating season, crabs arrive on the beach in female/male pairs to lay and fertilize eggs. However, unattached males, called “satellites,” also come to the beach, crowd around the nesting couples and compete with attached males for fertilizations, contributing to reproductive success. Some females are ignored by satellite males, and some attract more satellites than others, and the question is: what factors contribute to the number of satellites for each female? Or, perhaps better, how do unattached males choose among available females? This is another example in which zero counts may require special treatment.

The data, given in `CrabSatellites` in the `countreg` package, give the response variable, `satellites` for 173 females. Possible predictors are the female’s `color` and `spine` condition, given as ordered factors, as well as her `weight` and `carapace` (shell) width.

```
> data("CrabSatellites", package = "countreg")
> str(CrabSatellites)

'data.frame': 173 obs. of 5 variables:
 $ color      : Ord.factor w/ 4 levels "lightmedium"<...: 2 3 3 4 2 1 4 2 2 2 ...
 $ spine       : Ord.factor w/ 3 levels "bothgood"<"onebroken"<...: 3 3 3 2 3 2 3 3 1 3 ...
 $ width       : num  28.3 26 25.6 21 29 25 26.2 24.9 25.7 27.5 ...
 $ weight      : num  3.05 2.6 2.15 1.85 3 2.3 1.3 2.1 2 3.15 ...
 $ satellites: int  8 4 0 0 1 3 0 0 8 6 ...
```

Agresti (2013, §4.3) analyses the number of satellites using count data GLMs, and in his Chapter 5, describes separate logistic regression models for the binary outcome of one or more satellites vs. none. Later in this chapter (Section 9.4) we consider hurdle and zero-inflated models for count data. These have the advantage of modeling the zero counts together with a model for the positive counts.

A useful overview plot of the data is shown using `gpairs()` in Figure ???. You can see that the distribution of `satellites` is quite positively skewed, with many zero counts. `width` and `weight` are highly correlated (0.89), and both relate to the size of the female. Their scatterplots in the first row show that larger females attract more satellites. The categorical ordered factors `spine` condition and `color` are strongly associated, with the lightest colored crabs having the best conditions.

```
> library(vcd)
> library(gpairs)

Error in library(gpairs): there is no package called 'gpairs'

> gpairs(CrabSatellites[,5:1],
+         diag.pars = list(fontsize=16),
+         mosaic.pars = list(gp=shading_Friendly, gp_args=list(interpolate=1:4)))

Error in eval(expr, envir, enclos): could not find function "gpairs"
```

Figure 9.3 shows the scatterplots of `satellites` against `width` and `weight` together with smoothed lowess curves.

```
> plot(jitter(satellites) ~ width, data=CrabSatellites,
+       ylab="Number of satellites (jittered)", xlab="Carapace width",
+       cex.lab=1.25)
> with(CrabSatellites, lines(lowess(width, satellites), col="red", lwd=2))
> plot(jitter(satellites) ~ weight, data=CrabSatellites,
+       ylab="Number of satellites (jittered)", xlab="Weight",
+       cex.lab=1.25)
> with(CrabSatellites, lines(lowess(weight, satellites), col="red", lwd=2))
```

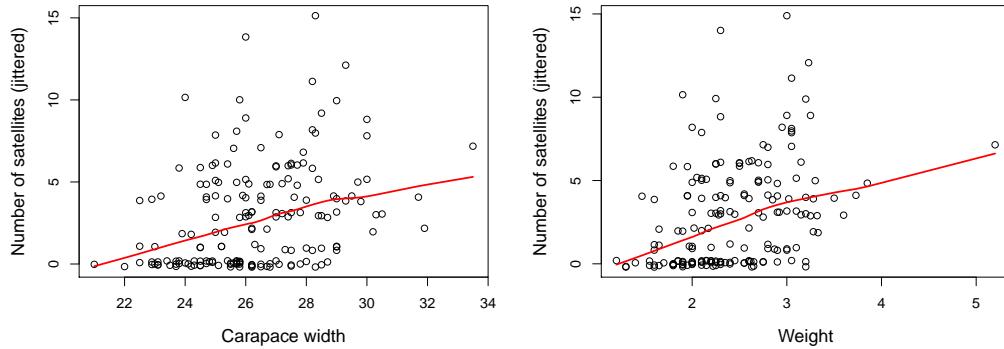


Figure 9.3: Scatterplots of number of satellites vs. width and weight, with lowess smooths.
fig:crabs1-scats

Both variables show approximately linear relations to the mean number of satellites, so it would not be unreasonable to fit models using the identity link ($\mu \sim x$) rather than the log link ($\mu \sim \log(x)$) with the Poisson family GLM.

In these plots, we reduce the problem of overplotting of the discrete response by jittering, but an alternative technique is to transform a numeric count or continuous predictor to a factor (for visualization purposes only), thereby giving boxplots. A convenience function for this purpose, `cutfac()` is defined in `vcdExtra`. It acts like `cut()`, but gives nicer labels for the factor levels and by default chooses convenient breaks among the values based on deciles. Using this, the plots in Figure 9.3 can be re-drawn as boxplots, giving Figure 9.4.

```
> plot(satellites ~ cutfac(width), data=CrabSatellites,
+       ylab="Number of satellites", xlab="Carapace width (deciles)")
> plot(satellites ~ cutfac(weight), data=CrabSatellites,
+       ylab="Number of satellites", xlab="Weight (deciles)")
```

With this visual overview, we proceed to an initial Poisson GLM model, using all predictors. Note that `color` and `spine` are ordered factors, so `glm()` represents them as polynomial contrasts, as if they were coded numerically. **TODO: output.lines=9:26 works in chapter.Rnw, but not in book.Rnw. Why???**

```
> crabs.pois <- glm(satellites ~ ., data=CrabSatellites, family=poisson)
> summary(crabs.pois)

Call:
glm(formula = satellites ~ ., family = poisson, data = CrabSatellites)

Deviance Residuals:
    Min      1Q  Median      3Q     Max 
      -1      -1      -1      -1      -1 

Coefficients:
            Estimate Std. Error z value Pr(>|z|)    
(Intercept)  1.3825    0.0982  14.00   <2e-16 ***
width       -0.0320    0.0080  -4.00   1.1e-09 ***
weight      -0.0180    0.0080  -2.25    0.0250 *  
colorL      -0.0001    0.0001  -0.01    0.9900    
colorR      -0.0001    0.0001  -0.01    0.9900    
spineL      -0.0001    0.0001  -0.01    0.9900    
spineR      -0.0001    0.0001  -0.01    0.9900    
---
Signif. codes:  '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 1.382 on 11 degrees of freedom

```

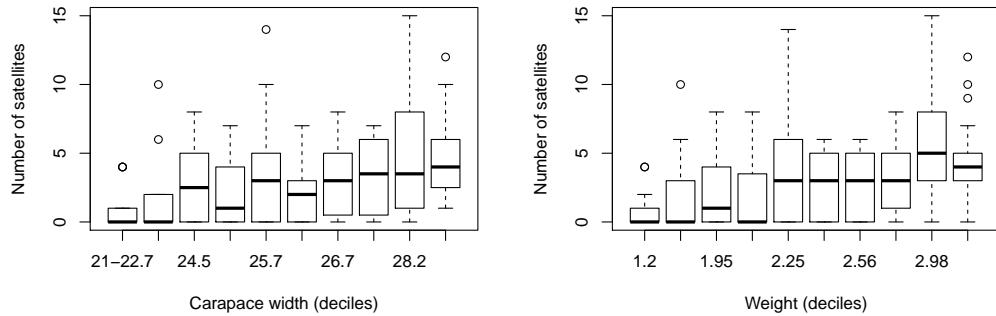


Figure 9.4: Boxplots of number of satellites vs. width and weight.
fig:crabs1-boxplots

```

-3.029 -1.863 -0.599  0.933  4.945

Coefficients:
            Estimate Std. Error z value Pr(>|z|)
(Intercept) -0.7057   0.9344  -0.76  0.4501
color.L      -0.4120   0.1567  -2.63  0.0085 **
color.Q       0.1237   0.1231   1.00  0.3150
color.C       0.0481   0.0914   0.53  0.5983
spine.L      0.0618   0.0848   0.73  0.4660
spine.Q      0.1585   0.1609   0.99  0.3244
width        0.0165   0.0489   0.34  0.7358
weight       0.4971   0.1663   2.99  0.0028 **

Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for poisson family taken to be 1)

Null deviance: 632.79 on 172 degrees of freedom
Residual deviance: 549.56 on 165 degrees of freedom
AIC: 920.9

Number of Fisher Scoring iterations: 6

```

The Wald tests for the coefficients show that only the linear effect of color and the effect of width are significant. Effect plots, in Figure ??, show the nature of these effects—lighter colored females attract more satellites, as do wider and heavier females.

```

> plot(allEffects(crabs.pois), main="")
Error in plot(allEffects(crabs.pois), main = ""): error in evaluating the
argument 'x' in selecting a method for function 'plot': Error: could not
find function "allEffects"

```

A simpler model can be constructed using `color` as a numeric variable, and either `width` or `weight` to represent female size. We choose `weight` here.⁷

⁷Agresti (2013, §4.3) and others who have analyzed this example uses carapace width as the main quantitative predictor, possibly because width might be more biologically salient to the single males than weight. This is a case where two highly correlated predictors are each strongly related to the outcome, yet partial tests (controlling for all others) may prefer one over the other.

```

> CrabSatellites1 <- transform(CrabSatellites,
+   color = as.numeric(color))
>
> crabs.pois1 <- glm(satellites ~ weight + color, data=CrabSatellites1,
+   family=poisson)
> summary(crabs.pois1)

Call:
glm(formula = satellites ~ weight + color, family = poisson,
     data = CrabSatellites1)

Deviance Residuals:
    Min      1Q  Median      3Q      Max 
-2.978  -1.916  -0.547   0.918   4.834 

Coefficients:
            Estimate Std. Error z value Pr(>|z|)    
(Intercept)  0.0888    0.2544    0.35   0.727    
weight       0.5458    0.0675    8.09  6e-16 ***  
color        -0.1728    0.0615   -2.81   0.005 **  
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for poisson family taken to be 1)

Null deviance: 632.79 on 172 degrees of freedom
Residual deviance: 552.77 on 170 degrees of freedom
AIC: 914.1

Number of Fisher Scoring iterations: 6

```

From the statistical and graphical analysis so far, the answer to the question posed in this example is clear: unattached male horseshoe crabs prefer light-colored big, fat mamas!

Yet, neither of these models fit well, as can be seen from their residual deviances and likelihood-ratio tests.

```

> vcdExtra::LRstats(crabs.pois, crabs.pois1)

Likelihood summary table:
          AIC BIC LR Chisq Df Pr(>Chisq)
crabs.pois 921 946      550 165    <2e-16 ***
crabs.pois1 914 924      553 170    <2e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

Perhaps there is something else to be learned here.



9.3 Models for overdispersed count data

{sec:glm-overdisp}

In practice, the Poisson model is often very useful for describing the relationship between the mean μ_i and the linear predictors, but typically underestimates the variance in the data. The consequence is that the Poisson standard errors are too small, rendering the Wald tests of coefficients, $z_j = \hat{\beta}_j/\text{se}(\hat{\beta}_j)$ (and other hypothesis test statistics) too large, and thus overly liberal.

In applications of the GLM, overdispersion is usually assessed by the likelihood-ratio test of the deviance (or the Pearson statistic) given in Section 9.1.3, but there is a subtle problem here. Lack of fit in a GLM for count data can result either from a mis-specified model for the systematic

component (omitted or unmeasured predictors, non-linear relations, etc.) or from failure of the Poisson mean = variance assumption. Thus, use of these methods requires some high degree of confidence that the systematic part of the model has been correctly specified, so that any lack of fit can be attributed to overdispersion.

One way of dealing with this is to base inference on so-called *sandwich* covariance estimators that are robust against some types of model mis-specification. In R, this is provided by the `sandwich()` function in the `sandwich` package, and can be used with `coeftest(model, vcov=sandwich)` to give overdispersion-corrected hypothesis tests. Alternatively, the Poisson model variance assumption can be relaxed in the quasi-Poisson model and the negative-binomial model as discussed below.

9.3.1 The quasi-Poisson model

One obvious solution to the problem of overdispersion for count data is the relaxed assumption that the conditional variance is merely *proportional* to the mean,

$$\mathcal{V}(y_i|\eta_i) = \phi\mu_i$$

Overdispersion is the common case of $\phi > 1$, implying that the conditional variance increases faster than the mean, but the opposite case of underdispersion, $\phi < 1$ is also possible, though relatively rare in practice. This strategy entails estimating the dispersion parameter ϕ from the data, and gives the **quasi-Poisson model** for count data.

One possible estimate is the residual deviance divided by degrees of freedom. However, it is more common to use the Pearson statistics, that gives a method-of-moments estimate with improved statistical properties.

$$\hat{\phi} = \frac{X_P^2}{n-p} = \sum_{i=1}^n \frac{(y_i - \hat{\mu}_i)^2}{\hat{\mu}_i} / (n-p)$$

It turns out that this model gives the same coefficient estimates as the standard Poisson GLM, but inference is adjusted for over/under dispersion. In particular, following Eqn. (9.1) the standard errors of the model coefficients are multiplied by $\hat{\phi}^{1/2}$ and so are inflated when overdispersion is present. In R, the quasi-Poisson model with this estimated dispersion parameter is fitted with the `glm()` function, by setting `family=quasipoisson`.

{sec:glm-quasi}

{ex:phdpubs2}

EXAMPLE 9.3: Publications of PhD candidates

For the `PhdPub` data, the deviance and Pearson estimates of dispersion ϕ can be calculated using the results of the Poisson model saved in the `phd.pois` object. The Pearson estimate, 1.83, indicates that standard errors of coefficients in this model should be multiplied by $\sqrt{1.83} = 1.35$, a 35% increase, to correct for overdispersion.

```
> with(phd.pois, deviance/df.residual)
[1] 1.7971
> sum(residuals(phd.pois, type = "pearson")^2)/phd.pois$df.residual
[1] 1.8304
```

The quasi-Poisson model is then fitted using `glm()` as:

```
> phd.qpois <- glm(articles ~ ., data=PhdPub, family=quasipoisson)
```

For use in other computation, the dispersion parameter estimate $\hat{\phi}$ can be obtained as the `dispersion` value of the `summary()` method for a quasi-Poisson model.

```
> (phi <- summary(phd.qpois)$dispersion)
[1] 1.8304
```

Note that this value can be compared to the variance/mean ratio of 2.91 calculated for the marginal distribution in Example 9.1; there is considerable improvement taking the predictors into account.



9.3.2 The negative-binomial model

{sec:glm-negbin}

The negative-binomial (NB) model for count data was introduced in Section 3.2.3 as a different generalization of the Poisson model that allows for overdispersion. In the context of the GLM, this can be developed as the extended form where the distribution of $y_i | \boldsymbol{x}_i$ where the mean μ_i for fixed \boldsymbol{x}_i can vary across observations i according to a gamma distribution with mean μ_i and a constant shape parameter, θ , reflecting the additional variation due to heterogeneity.

For a fixed value of θ , the negative-binomial is another special case of the GLM. The expected value of the response is again $E(y_i) = \mu_i$, but the variance function is $V(y_i) = \mu_i + \mu_i^2/\theta$, so the variance of y increases more rapidly than that of the Poisson distribution. Some authors (e.g., Agresti (2013), Hilbe (2014)) prefer to parameterize the variance function in terms of $\alpha = 1/\theta$, giving

$$V(y_i) = \mu_i + \mu_i^2/\theta = \mu_i + \alpha\mu_i^2 ,$$

so that α is a kind of dispersion parameter. Note that as $\alpha \rightarrow 0$, $V(y_i) \rightarrow \mu_i$ and the negative-binomial converges to the Poisson.

The MASS package provides the family function `negative.binomial(theta)` that can be used directly with `glm()` provided that the argument `theta` is specified. One example would be the related geometric distribution (Section 3.2.4), that is the special case of $\theta = 1$. This can be fitted in R by setting `family=negative.binomial(theta=1)` in the call to `glm()`.

Most often, θ is unknown and must be estimated from the data. In this case, the negative-binomial model is not a special case of the GLM, but it is possible to obtain maximum likelihood estimates of both β and θ , by iteratively estimating β for fixed θ and vice-versa. This method is implemented in the `glm.nb()` in the package MASS.

EXAMPLE 9.4: Mating of horseshoe crabs

For example, for the `CrabSatellites` data, we can fit the general negative-binomial model with θ free.

```
> library(MASS)
> crabs.nbin <- glm.nb(satellites ~ weight + color, data=CrabSatellites1)
> crabs.nbin$theta
[1] 0.95562
```

The estimated value $\hat{\theta}$ returned by `glm.nb()` is not very far from 1. Hence, we might also consider fixing $\theta = 1$, as illustrated below.

```
> crabs.nbin1 <- glm(satellites ~ weight + color, data=CrabSatellites1,
+ family=negative.binomial(1))
```



9.3.3 Visualizing the mean–variance relation

The quasi-Poisson and negative-binomial models have different variance functions, and one way to visualize which provides a better fit to the data is to group the data according to the fitted value of the linear predictor, calculate the mean and variance for each group, and then plot the variances against the means. A smoothed curve will then approximate the *empirical* mean–variance relationship. To this, we can add curves showing the mean–variance function implied by various models.⁸

{ex:phdpubs3}

EXAMPLE 9.5: Publications of PhD candidates

For the *PhdPub*s data, the fitted values are obtained with `fitted()` for the Poisson and negative binomial models. Either set can be used to categorize the observations into groups for the purpose of calculating means and variances of the response.

```
> fit.pois <- fitted(phd.pois, type="response")
> fit.nbin <- fitted(phd.nbin, type="response")
```

Here we use a simpler version of the `cutfac()` function to group a numeric variable into quantile-based groups. `cutq()` also uses deciles by default, and just uses simple integer values for the factor labels.

```
> cutq <- function(x, q = 10) {
+   quantile <- cut(x, breaks = quantile(x, probs = 0:q/q),
+                     include.lowest = TRUE, labels = 1:q)
+   quantile
+ }
```

Using this, we create a variable `group` giving 20 quantile groups of the fitted values, and then use `aggregate()` to find the mean and variance of the number of articles in each group.

```
> group <- cutq(fit.nbin, q=20)
> qdat <- aggregate(PhdPub$articles,
+                      list(group),
+                      FUN = function(x) c(mean=mean(x), var=var(x)))
> qdat <- data.frame(qdat$x)
> qdat <- qdat[order(qdat$mean),]
```

We can then calculate the theoretical variances implied by the quasi-Poisson and negative-binomial models:

```
> phi <- summary(phd.qpois)$dispersion
> qdat$qvar <- phi * qdat$mean
> qdat$nbvar <- qdat$mean + (qdat$mean^2) / phd.nbin$theta
> head(qdat)

  mean      var    qvar    nbvar
1 0.61224 0.78401 1.1206 0.7776
2 1.14894 1.78168 2.1030 1.7312
6 1.22917 1.71232 2.2499 1.8956
8 1.24444 2.46162 2.2778 1.9276
4 1.26087 1.70821 2.3079 1.9622
5 1.33333 2.81818 2.4405 2.1175
```

The plot, shown in Figure 9.5, then simply plots the points and uses `lines()` to plot the model-implied variances.

⁸This idea and the example that follows was suggested by Germán Rodrigues in a Stata example given at <http://data.princeton.edu/wws509/stata/overdispersion.html>.

```
> with(qdat, {
+   plot(var ~ mean, xlab="Mean number of articles", ylab="Variance",
+       pch=16, cex=1.2, cex.lab=1.2)
+   abline(h=mean(PhdPubs$articles), col=gray(.40), lty="dotted")
+   lines(mean, qvar, col="red", lwd=2)
+   lines(mean, nbvar, col="blue", lwd=2)
+   lines(lowess(mean, var), lwd=2, lty="dashed")
+   text(3, mean(PhdPubs$articles), "Poisson", col=gray(.40))
+   text(3, 5, "quasi-Poisson", col="red")
+   text(3, 6.7, "negbin", col="blue")
+   text(3, 8.5, "lowess")
+ })
```

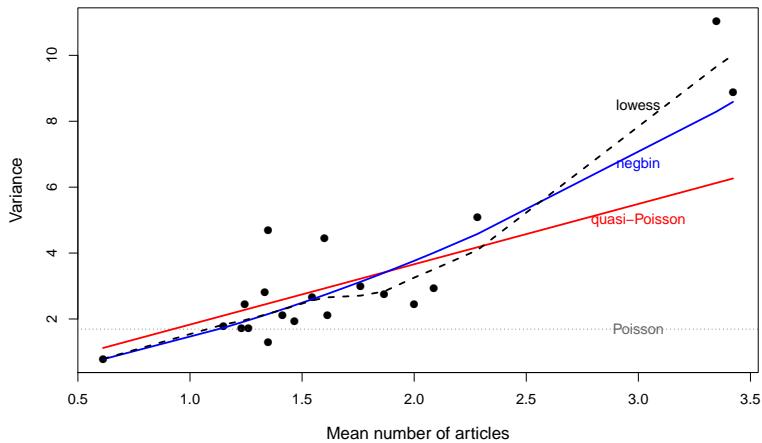


Figure 9.5: Mean–variance functions for the PhdPubs data. Points show the observed means and variances for 20 quantile groups based on the fitted values in the negative-binomial model. The labeled lines and curves show the variance functions implied by various models.
fig:phd-mean-var-plot

We can see from this plot that the variances implied by the quasi-Poisson and negative-binomial models are in reasonable accord with the data and with each other up to a mean of about 2.5. They diverge substantially at the upper end, for the 20–30% of the most productive candidates, where the quadratic variance function of the negative-binomial provides a better fit.

Finally, we can also compare the standard errors of coefficients for the various methods designed to correct for overdispersion. These are extracted as the diagonal elements of the `vcov()` and `sandwich()` methods from the model objects.

```
> library(sandwich)
> phd.SE <- sqrt(cbind(
+   pois=diag(vcov(phd.pois)),
+   sand=diag(sandwich(phd.pois)),
+   qpois=diag(vcov(phd.qpois)),
+   nbin=diag(vcov(phd.nbin))))
> round(phd.SE, 4)
```

	pois	sand	qpois	nbin
(Intercept)	0.0996	0.1382	0.1348	0.1327
female1	0.0546	0.0714	0.0738	0.0726
married1	0.0613	0.0823	0.0829	0.0819
kid5	0.0401	0.0560	0.0543	0.0528

```
phdprestige 0.0253 0.0392 0.0342 0.0343
mentor      0.0020 0.0039 0.0027 0.0032
```

For this example, the sandwich, quasi-Poisson and negative-binomial methods give similar results, all about 40% larger on average than those from the Poisson model. \triangle

9.3.4 Testing overdispersion

The forms of overdispersion seen in these examples and in Figure 9.5 give rise to a statistical test (Cameron and Trivedi 1990; Cameron and Trivedi 1998, §3.4) for the null hypothesis of Poisson variation, $H_0 : \mathcal{V}(y) = \mu$ against an alternative that the variance has a particular form depending on the mean,

$$\mathcal{V}(y) = \mu + \alpha \times f(\mu) ,$$

where $f(\mu)$ is a given transformation function of the mean.

Overdispersion corresponds to $\alpha > 0$ and underdispersion to $\alpha < 0$. The coefficient α can be estimated by an auxiliary OLS regression (without an intercept, i.e., of the form

```
lm(var ~ -1 + f(mean))
```

and tested with the corresponding t (or z) statistic, which is asymptotically standard normal under the null hypothesis.

Common specifications of the transformation function are $f(\mu) = \mu$ and $f(\mu) = \mu^2$. The first corresponds to a NB model with a linear variance function (called NB1 by various authors) or a quasi-Poisson model with dispersion parameter ϕ , i.e.,

$$\mathcal{V}(y) = (1 + \alpha)\mu = \phi\mu .$$

The second is the more traditional form with quadratic variance function described in Section 9.3.2 (called NB2 by some authors).

These tests are carried out using the `dispersiontest()` function in the `AER` package. The first argument is a Poisson GLM model; the second specifies the alternative hypothesis, either as an integer power of μ or a function of the mean.

```
> library(AER)

Error in library(AER) : there is no package called 'AER'

> dispersiontest(phd.pois)

Error in eval(expr, envir, enclos) : could not find function "dispersiontest"

> dispersiontest(phd.pois, 2)

Error in eval(expr, envir, enclos) : could not find function "dispersiontest"
```

These tests use a specified alternative hypothesis, so there is no way to compare directly which of the NB1 or NB2 models is better or worse, except by using methods such as AIC or BIC described in Section 9.1.4.

9.3.5 Visualizing goodness-of-fit

Even with correction for overdispersion, goodness-of-fit tests provide only an overall summary of model fit. Some specialized tests for particular forms of overdispersion are also available (e.g., see Cameron and Trivedi (1998, Chapter 5)), but these only identify general problems and cannot provide detailed indications of the possible source of these problems.

{sec:glm-disptest}

{sec:glm-visfit}

In Chapter 3, we illustrated the use of rootograms for visualizing goodness-of-fit to a wide variety discrete distributions using the `plot()` method for class "goodfit" objects with the `vcd` package. However, those methods were developed for one-way discrete distributions without explanatory variables.

Kleiber and Zeileis (2014) have generalized this idea to the wider class of GLM-related count regression models considered here. The `countreg` package provides a new implementation of `rootogram()` with methods for all of these models (and others not mentioned). We illustrate these plots for the models considered to this point, and then extend this use for models allowing for excess zero counts in Section 9.4.

EXAMPLE 9.6: Publications of PhD candidates

For the `PhdPubs` data, Figure 9.6 shows hanging rootograms for the Poisson and negative-binomial models produced using `countreg::rootogram`⁹ on the fitted model objects. We are looking both for general patterns of under/over fit, as well as counts that stand out as poorly fitted against the background.

```
> library(countreg)
> countreg::rootogram(phd.pois, max=12, main="PhDPubs: Poisson")
> countreg::rootogram(phd.nbin, max=12, main="PhDPubs: Negative-Binomial")
```

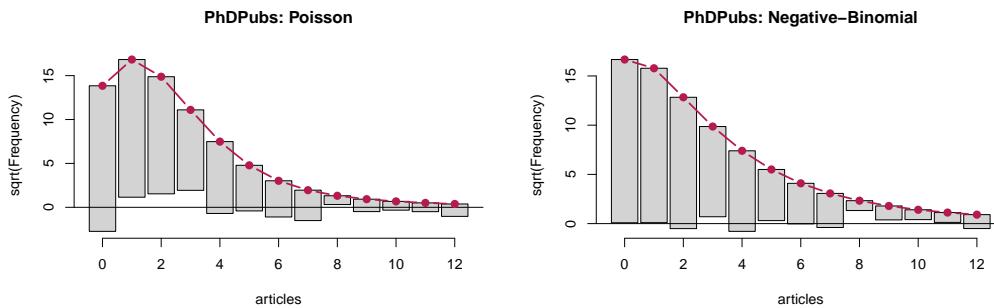


Figure 9.6: Hanging rootograms for the `PhdPubs` data.^{fig:phdpubs4-rootogram}

The Poisson model shows a systematic, wave-like pattern with excess zeros, too few observed frequencies for counts of 1–3, but generally greater frequencies for counts of 4 or more. The negative-binomial model clearly fits much better, though there is a peculiar tendency among the smaller frequencies for 8 or more articles. △

```
{ex:crabs2}
```

EXAMPLE 9.7: Mating of horseshoe crabs

Figure 9.7 shows similar plots for the same two models fit to the number of crab satellites. The fit of the Poisson model clearly reveals the excess of zero male satellites. For the negative-binomial, the rootogram no longer exhibits same wave-like pattern, however, the underfitting of the count for 0 and overfitting for counts 1–2 is characteristic of data with excess zeros.

```
> countreg::rootogram(crabs.pois, max=15, main="CrabSatellites: Poisson")
> countreg::rootogram(crabs.nbin, max=15, main="CrabSatellites: Negative-Binomial")
```

△

⁹At the time of this writing, `rootogram` in `countreg` conflicts with the version in `vcd`, so we qualify the use here with the package name.

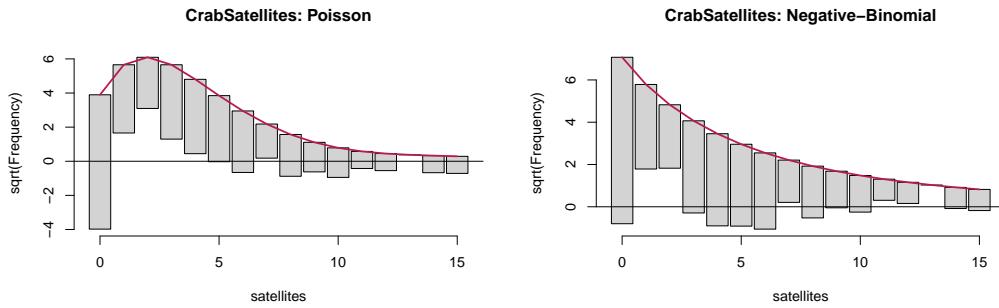


Figure 9.7: Hanging rootograms for the CrabSatellites data.
fig:crabs2-rootogram

9.4 Models for excess zero counts

{ sec:glm-zeros }

In addition to overdispersion, many sets of empirical data exhibit a greater prevalence of zero counts than can be accommodated by the Poisson or negative-binomial models. We saw this in the *PhdPub*s data set, where there were many candidates who had not published at all, and in the *CrabSatellites* data where a large number of females attracted no unattached males. Other examples abound in many different fields: studies of the use of health care services often find that many people never visit a hospital in some time frame; similarly, the distribution of insurance claims often shows large numbers who make no claims (Yip and Yau, 2005) because of under-reporting of small claims, policy deductible provisions and desire to avoid premium increases.

Beyond simply identifying this as a problem of lack-of-fit, understanding the reasons for excess zero counts can make a contribution to a more complete explanation of the phenomenon of interest, and this requires both new statistical models and visualization techniques illustrated in this section.

In the first example, Long (1997) argued that the PhD candidates might fall into two distinct groups: “publishers” (perhaps striving for an academic career) and “non-publishers” (seeking other career paths). Of the 275 observations having `articles==0`, some might not have published due to chance or unmeasured factors. One reasonable form of explanation is that the observed zero counts reflect a mixture of the two latent classes—those who simply have not yet published and those who will likely never publish. A statistical formulation of this idea leads to the class of *zero-inflated* models described below.

A different form of explanation is that there may be some special circumstance or “hurdle” required to achieve a positive count, like publishing the master’s thesis (such as being driven internally by a personality trait or externally by pressure from a mentor). This idea leads to the class of *hurdle* models that entertain and fit (simultaneously) two separate models: one for the occurrence of the zero counts, and one for the positive counts. These two approaches are illustrated in Figure 9.8

9.4.1 Zero-inflated models

{ sec:glm-zip }

Zero-inflated models, introduced by Lambert (1992) as the *zero-inflated Poisson* (ZIP) model, provide an attractive solution to the problem of dealing with an overabundance of zero counts. It postulates that the observed counts arise from a mixture of two latent classes of observations: some structural zeros for whom y_i will always be 0, and the rest, sometimes giving random zeros. The ZIP model is comprised of two components:

- A model for the binary event of membership in the unobserved (latent) class of those for whom the count is necessarily zero (e.g., “non-publishers”). This is typically taken as a logistic regression for the probability π_i that observation i is in this class, with predictors z_1, z_2, \dots, z_q ,

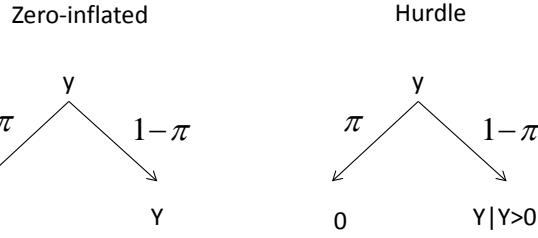


Figure 9.8: Models for excess zeros. The observed response y is derived from a latent or parent distribution for Y yielding zero counts with probability π .

{fig:ExcessZeros}

giving

$$\text{logit}(\pi_i) = \mathbf{z}_i^\top \boldsymbol{\gamma} = \gamma_0 + \gamma_1 z_{i1} + \gamma_2 z_{i2} + \cdots + \gamma_q z_{iq} . \quad (9.7) \quad \text{(eq:zip-logit)}$$

- A Poisson model for the other class (e.g., “publishers”), for whom the observed count may 0 or positive. This model typically uses the usual log link to predict the mean, using predictors $\mathbf{x}_1, \mathbf{x}_2, \dots, \mathbf{x}_p$, so

$$\log_e \mu(\mathbf{x}_i) = \mathbf{x}_i^\top \boldsymbol{\beta} = \beta_0 + \beta_1 x_{i1} + \beta_2 x_{i2} + \cdots + \beta_q x_{ip} . \quad (9.8) \quad \text{(eq:zip-pois)}$$

In application, it is permissible and not uncommon to use the same set of predictors $x = z$ in both submodels, but the notation indicates that this is not required. Some simple special cases arise when the model for the always zero latent class is an intercept-only model, $\text{logit}(\pi_i) = \gamma_0$, implying the same probability for all individuals, and (less commonly) when the Poisson mean model is intercept-only with no predictors but there might be excess zero counts.

With this setup, one can show that the probability of observing counts of $y_i = 0$ and $y_i > 0$ are

$$\begin{aligned} \Pr(y_i = 0 | \mathbf{x}, \mathbf{z}) &= \pi_i + (1 - \pi_i)e^{-\mu_i} \\ \Pr(y_i | \mathbf{x}, \mathbf{z}) &= (1 - \pi_i) \times \left[\frac{\mu_i^{y_i} e^{-\mu_i}}{y_i!} \right] , \quad y_i \geq 0 \end{aligned} \quad (9.9)$$

where the term in brackets in the second equation is the Poisson probability $\Pr(y = y_i)$ with rate parameter $\text{Pois}(\mu_i)$. In these equations, $\pi_i = \text{logit}^{-1}(\mathbf{z}_i^\top \boldsymbol{\gamma})$ depends on the \mathbf{z} through Eqn. (9.7), and $\mu_i = \exp(\mathbf{x}_i^\top \boldsymbol{\beta})$ depends on the \mathbf{x} through Eqn. (9.8).

The conditional expectation and variance of y_i then have the forms

$$\begin{aligned} \mathcal{E}(y_i) &= (1 - \pi_i) \mu_i \\ \mathcal{V}(y_i) &= (1 - \pi_i) \mu_i (1 + \mu_i \pi_i) . \end{aligned}$$

Thus, when $\pi_i > 0$, the mean of y is always less than μ_i , and the variance of y is greater than its mean by a dispersion factor of $(1 + \mu_i \pi_i)$.

There is nothing special about the use of the Poisson distribution here. The model for the count variable could also be taken as the negative-binomial, giving a *zero-inflated negative-binomial* (ZINB) model using $\text{NBin}(\mu, \theta)$ or a *zero-inflated geometric* model using $\text{NBin}(\mu, \theta = 1)$.

EXAMPLE 9.8: Simulating zero-inflated data

A simple way of understanding the effects of zero-inflation on count data is to simulate data from their distribution and plot it. For the standard Poisson and negative-binomial, random values can be generated using `rpois()` and `rnegbin()` (in **MASS**), respectively. Their zero-inflated counterparts are implemented in the **VGAM** package as `rzipois()` and `rzinegbin()`.

To illustrate this use, we generate two random data sets using `rziropois()` having constant mean $\mu = 3$. The first is a standard Poisson ($\pi = 0$), while the second has a constant probability $\pi = 0.3$ of an excess zero.

```
> library(VGAM)
> set.seed(1234)
> data1 <- rziropois(200, 3, 0)
> data2 <- rziropois(200, 3, .3)
```

Barplots of the frequencies in these data sets are shown in Figure 9.9. The sample mean in `data1` is 2.925, quite close to $\mu = 3$. In the zero-inflated `data2`, the mean is only 2.25 due to the excess zeros.

```
> tdata1 <- table(data1)
> barplot(tdata1, xlab="Count", ylab="Frequency",
+           main="Poisson(3)")
> tdata2 <- table(data2)
> barplot(tdata2, xlab="Count", ylab="Frequency",
+           main=expression("ZI Poisson(3, " * pi * " = .3)"))
```

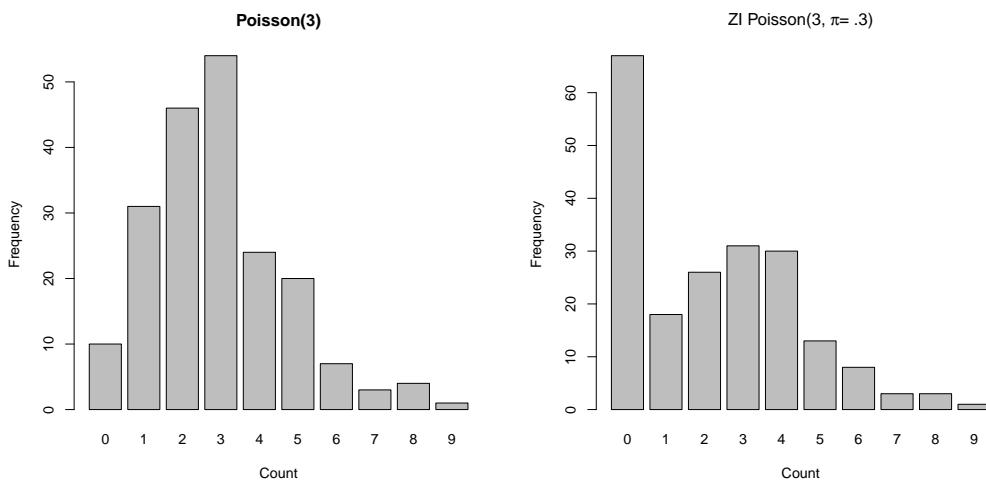


Figure 9.9: Bar plots of simulated data from Poisson and zero-inflated Poisson distributions fig:ziropois-plot



There are several packages in R capable of fitting zero-inflated models. The most mature and complete of these is `zeroinfl()` in the `countreg` package (a successor to the `pscl` package). The function `zeroinfl()` is modeled after `glm()`, but provides an extended syntax for the model formula.

If the `formula` argument is supplied in the form $y \sim x_1 + x_2 + \dots$, it not only describes the count regression of y on x_1, x_2, \dots , but also implies that the *same* set of regressors, $z_j = x_j$, is used for the zero count binary submodel. The extended syntax uses the notation $y \sim x_1 + x_2 + \dots \mid z_1 + z_2 + \dots$ to specify the x variables separately, conditional on (\mid) the always-zero count model $y \sim z_1 + z_2 + \dots$. The model for the not-always-zero class can be specified using the `dist` argument, with possible values "poisson", "negbin" and "geometric".

9.4.2 Hurdle models

A different class of models capable of accounting for excess zero counts is the ***hurdle model*** (also called the ***zero-altered model***) proposed initially by Cragg (1971) and developed further by Mullahy (1986). This model also uses a separate logistic regression submodel to distinguish counts of $y = 0$ from larger counts, $y > 0$. The submodel for the positive counts is expressed as a (left) *truncated* Poisson or negative-binomial model, excluding the zero counts. As an example, consider a study of behavioral health in which one outcome is the number of cigarettes smoked in one month. All the zero counts will come from non-smokers and smokers will nearly always smoke a positive number.

This differs from the set of ZIP models in that classes of $y = 0$ and $y > 0$ are now considered fully-observed, rather than latent. Conceptually, there is one process and submodel accounting for the zero counts and a separate process accounting for the positive counts, once the “hurdle” of $y = 0$ has been passed. In other words, for ZIP models, the first process generates only extra zeros beyond those of the regular Poisson distribution. For hurdle models, the first process generates all of the zeros. The probability equations corresponding to Eqn. (9.9) are:

$$\begin{aligned}\Pr(y_i = 0 | \mathbf{x}, \mathbf{z}) &= \pi_i \\ \Pr(y_i | \mathbf{x}, \mathbf{z}) &= \frac{(1 - \pi_i)}{1 - e^{-\mu_i}} \times \left[\frac{\mu_i^{y_i} e^{-\mu_i}}{y_i!} \right], \quad y_i \geq 0\end{aligned}\tag{9.10}$$

The hurdle model can be fitted in R using the `hurdle()` function from the `countreg` package. The syntax for the model formula is the same extended form provided by `zeroinfl()`, where $y \sim x_1 + x_2$ uses the same regressors for the zero and positive count submodels, while $y \sim x_1 + x_2 | z_1 + z_2$ uses $y \sim z_1 + z_2$ for the zero hurdle model. Similarly, the count distribution can be given as "poisson", "negbin" or "geometric" with the `dist` argument. For `hurdle()`, the distribution for zero model can be specified with a `zero.dist` argument. The default is "binomial" (with a logit link), but other right-censored distributions can also be specified.

9.4.3 Visualizing zero counts

Both the zero-inflated and hurdle models treat the zero counts $y = 0$ specially with separate submodels, so the binary event of $y = 0$ vs. $y > 0$ can be visualized using any of the techniques illustrated in Chapter 7. See Section 7.2.3, Section 7.3.1 and Section 7.3.2 for some examples that plot both the binary observations and a model summary or smoothed curve to show the relationships with one or more regressors. To apply these ideas in the current context, simply define or plot a logical variable corresponding to the expression `y==0`, giving values of TRUE or FALSE.

A different, and simpler idea is illustrated here using what is called a ***spine plot*** Hummel (1996) when a predictor x is a discrete factor or ***spinogram*** when x is continuous. Both are forms of mosaic plots with special formatting of spacing and shading, and in this context they plot $\Pr(y = 0|x)$ against $\Pr(x)$; when x is numerical, it is first made discrete, as in a histogram. Then, in the spine plot or spinogram, the widths of the bars correspond to the relative frequencies of x and heights of the bars correspond to the conditional relative frequencies of $y = 0$ in every x group. In R, spine plots are implemented in the function `spineplot()`, however, this is what you get by default if you use `plot(y==0 ~ x)` to plot the binary factor against any regressor x .

A related graphical method is the ***conditional density plot*** (Hofmann and Theus, 2005). The conditional probabilities $\Pr(y = 0|x)$ are derived using a smoothing approach (via `density()`) over x rather than by making x discrete. These plots are provided by `cdplot()` in the `graphics` package and a similar `cd_plot()` in `vcd`. The smoothing method for the density estimate is controlled by a `bw` (bandwidth) method and other arguments.

EXAMPLE 9.9: Mating of horseshoe crabs

For the *CrabSatellites* data, we can examine the relationship of the zero counts (females who attract no unattached male satellites) to the predictors using spinograms or conditional density plots. Here, we consider `weight` and `color` (treated numerically) as predictors. **TODO:** Fixup use of color in Example 9.2 so it doesn't cause a problem here

Spinograms for the occurrence of zero satellites against `weight` and `color` are shown in Figure 9.10, where we have used quantiles of those distributions to define the breaks on the horizontal axis. Using `ylevels=2:1` reverses the order of the vertical categories. You can easily see that the zeros decrease steadily with `weight` and increase with darkness.

```
> op <- par(cex.lab=1.2, mfrow = c(1, 2))
> plot(factor(satellites == 0) ~ weight, data = CrabSatellites,
+       breaks = quantile(weight, probs=seq(0,1,.2)), ylevels=2:1,
+       ylab="No satellites")
> plot(factor(satellites == 0) ~ color, data = CrabSatellites,
+       breaks = quantile(color, probs=seq(0,1,.33)), ylevels=2:1,
+       ylab="No satellites")
> par(op)
```

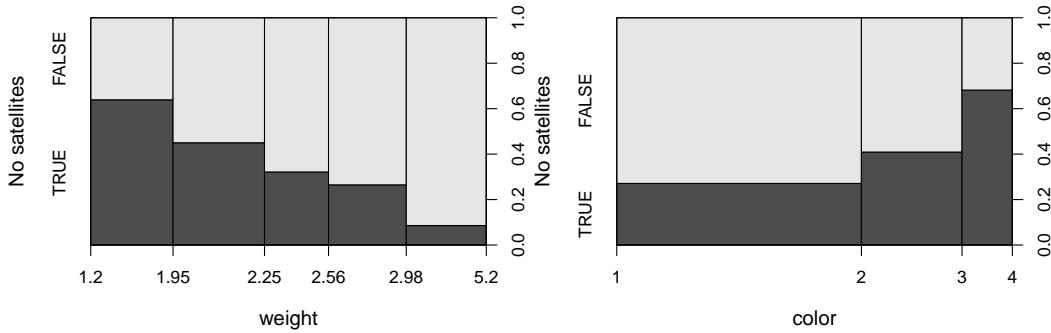


Figure 9.10: Spinograms for the *CrabSatellites* data. The variables `weight` (left) and `color` (right) have been made discrete using quantiles of their distributions.
fig:crabs-zero-spinogram

Similar plots in the form of conditional density plots are shown in Figure 9.11, with a similar interpretation.

```
> op <- par(cex.lab=1.2, mfrow = c(1, 2))
> cddplot(factor(satellites == 0) ~ weight, data = CrabSatellites,
+          ylevels=2:1, ylab="No satellites")
> cddplot(factor(satellites == 0) ~ color, data = CrabSatellites,
+          ylevels=2:1, , ylab="No satellites")
> par(op)
```



9.5 Case studies

In this section, we introduce two extended examples, designed to illustrate aspects of exploratory analysis, visualization, model fitting, and interpretation for count data GLMs. The first (Section 9.5.1) concerns another well-known data set from ethology, where (a) excess zeros require

{sec:glm-casesstudies}

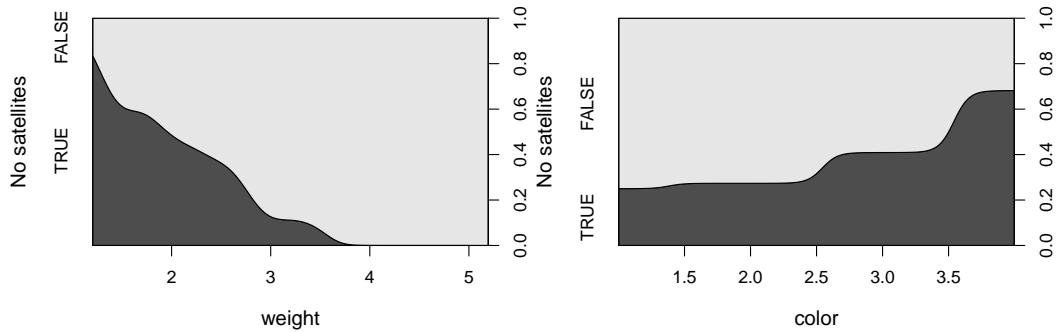


Figure 9.11: Conditional density plots for the CrabSatellites data. The region shaded below shows the conditional probability density estimate for a count of zero.^{fig:crabs-zero-caplot}

special treatment, (b) the occurrence of zero counts has substantive meaning, and (c) an interaction between two factors is important.

The second case study (Section 9.5.2) uses a larger, also well-known data set from health economics, with more predictors and more potential interactions. The emphasis shifts here from fitting and comparing models with different distributional forms and link functions to selecting terms for an adequate descriptive and explanatory model. Another feature of these examples is that the relatively large sample size in this data supports a wider range of model complexity than is available in smaller samples.

9.5.1 Cod parasites

The cod fishery is extremely important to the economy of Norway, so anything that affects the health of the cod population and its ecosystem can have severe consequences. The red king crab *Paralithodes camtschaticus* was deliberately introduced by Russian scientists to the Barents Sea in the 1960s and 1970s from its native area in the North Pacific. The carapace of these crabs is used by the leech *Johanssonia arctica* to deposit its eggs. This leech in turn is a vector for the blood parasite *Trypanosoma murmanensis* that can infect marine fish, including cod.

Hemmingsen *et al.* (2005) examined cod for trypanosome infections during annual cruises along the coast of Finnmark in North Norway over three successive years and in four different areas (A1: Sørøya; A2: Magerøya; A3: Tanafjord; A4: Varangerfjord). They show that trypanosome infections are strongest in the area Varangerfjord where the density of red king crabs is highest. Thus, there is evidence that the introduction of the foreign red king crabs had an indirect detrimental effect on the health of the native cod population. This situation stands out because it is not an introduced *parasite* that is dangerous for a native host, but rather an introduced *host* that promotes transmission of two endemic parasites. They call the connections among these factors “an unholy trinity.”¹⁰

EXAMPLE 9.10: Cod parasites

The data from Hemmingsen *et al.* (2005) is contained in *CodParasites* in the *countreg* package. It gives the results for 1254 cod caught by one ship in annual autumn cruises from 1999–2001. The main response variable, *intensity*, records the counted number of *Trypanosoma*

{fn:russian}

¹⁰ The four areas A1–A4 are arranged from east to west, with Varangerfjord (A4) closest to the Russian Kola Peninsula where the red king crabs initially migrated. A more specific test of the “Russian hypothesis” could be developed by treating area as an ordered factor and testing the linear component. We leave this analysis to an exercise for the reader.

parasites found in blood samples from these fish. To distinguish between infected vs. non-infected fish, a secondary response, prevalence is also recorded, corresponding to the expression

```
> CodParasites$prevalence <- ifelse(CodParasites$intensity == 0, "no", "yes")
```

Thus, intensity is the basic count response variable, and prevalence reflects the zero count that would be assessed in zero-inflated and hurdle models. In substantive terms, in a hurdle model, prevalence corresponds to whether a fish is infected or not; once infected, intensity gives the degree of infection. In a zero-inflated model, infected could be considered a latent variable; there are extra zeros from non-infected fish, but some infected fish are measured as “normal” zeros.

Hemmingsen *et al.* (2005) consider only three explanatory predictors: area, year (both factors) and length of the fish.¹¹ A quick numerical summary of the univariate properties of these variables is shown below. The intensity values are indeed extremely skewed, with a median of 0 and a maximum of 257. However, there are some missing values (NAs) among the response variables and a few in the length variable.

```
> data("CodParasites", package = "countreg")
> summary(CodParasites[, c(1:4, 7)])
```

	intensity	prevalence	area	year	length
Min.	0.00	no :654	soroya :272	1999:567	Min. : 17.0
1st Qu.	0.00	yes :543	mageroya :255	2000:230	1st Qu.: 44.0
Median	0.00	NA's: 57	tanafjord :415	2001:457	Median : 54.0
Mean	6.18		varangerfjord:312		Mean : 53.4
3rd Qu.	4.00				3rd Qu.: 62.0
Max.	257.00				Max. :101.0
NA's	57				NA's : 6

Even better, a quick univariate and bivariate summary of these variables can be shown in a generalized pairs plot (Figure ??).

```
> library(vcd)
> library(gpairs)

Error in library(gpairs): there is no package called 'gpairs'

> gpairs(CodParasites[, c(1:4, 7)],
+         diag.pars=list(fontsize=16),
+         mosaic.pars=list(gp=shading_Friendly))

Error in eval(expr, envir, enclos): could not find function "gpairs"
```

In this plot, among the categorical variables, prevalence is strongly associated with area, but also with year. As well there seems to be an association between area and year, meaning the number of cod samples collected in difference areas varied over time. In the univariate plots on the diagonal, intensity stands out as extremely skewed, and the distribution of length appears reasonably symmetric.

Before fitting any models, some more detailed exploratory plots are helpful for understanding the relationship of both prevalence and intensity to the predictors. The general idea is to make separate plots of prevalence and intensity and to try to show both the data and some simple summaries. In their Table 1, Hemmingsen *et al.* (2005) counted the missing observations as infected and we do the same to get a similar contingency table.

```
> cp.tab <- xtabs(~ area + year + factor(is.na(prevalence) |
+                  prevalence == "yes"),
+                  data = CodParasites)
> dimnames(cp.tab)[3] <- list(c("No", "Yes"))
> names(dimnames(cp.tab))[3] <- "prevalence"
```

¹¹Other potential predictors include weight, sex, age, and developmental stage, as well as the depth at which the fish were caught.

For the factors `area` and `year`, we can visualize prevalence as before (Example 9.9) using spineplots, but, for two (or more) factors, doubledecker and mosaic plots are better because they are more flexible and keep the factors distinct. The doubledecker plot (Figure 9.12) highlights the infected fish, and shows that prevalence is indeed highest in all years in Varangerfjord.

```
> doubledecker(prevalence ~ area + year, data=cp.tab)
```

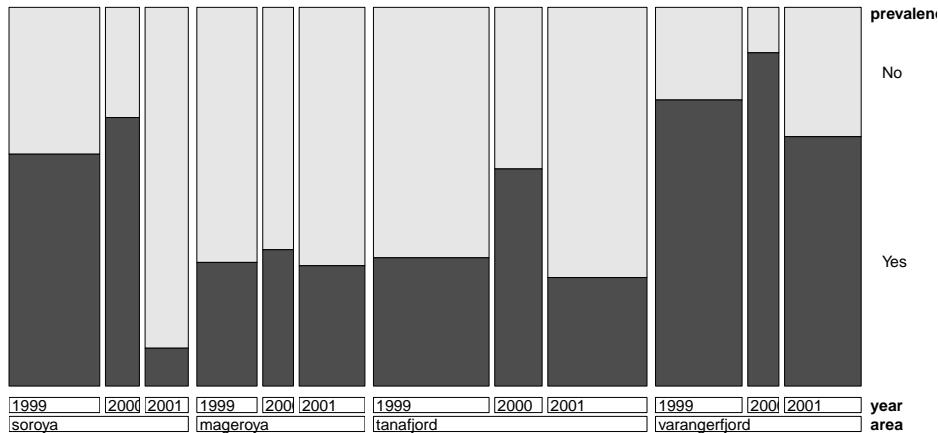


Figure 9.12: Doubledecker plot for prevalence against area and year in the CodParasites data. The cases of infected fish are highlighted.
Fig:cod1-doubledecker

A similar plot, in the doubledecker format, can be drawn as a mosaic plot, but now shading the tiles according a model for the expected counts. It makes sense here to consider the null loglinear model for prevalence as a response, independent of the combinations of area and year. This plot (Figure 9.13) shows further that prevalence differs substantially over the area-year combinations, so we should expect an interaction in the model for zero counts. As well, Varangerfjord stands out as having consistently greater prevalence in all years than expected under this model.

```
> mosaic(~area + year + prevalence, data=cp.tab,
+         split_vertical=c(TRUE, TRUE, FALSE),
+         labeling=labeling_doubledecker, spacing=spacing_highlighting,
+         expected = ~year:area + prevalence)
```

The effect of fish length on prevalence can be most easily seen by treating the factor as a numeric (0/1) variable and smoothing, as shown in Figure 9.14. The loess smoothed curve shows an apparent U-shaped relationship, however the plotted observations and the confidence bands make clear that there is very little data in the extremes of length.

```
> library(ggplot2)
> ggplot(CodParasites, aes(x=length, y=as.numeric(prevalence)-1)) +
+   geom_jitter(position=position_jitter(height=.05), alpha=0.25) +
+   geom_rug(position='jitter', sides='b') +
+   stat_smooth(method="loess", color="red", fill="red", size=1.5) +
+   theme_bw() + labs(y='prevalence')
```

For the positive counts of intensity, boxplots by area and year show the distributions of parasites, and it is again useful to display these on a log scale. In Figure 9.15, we have used `ggplot2`, with `geom_boxplot()` and `geom_jitter()` to also plot the individual observations.

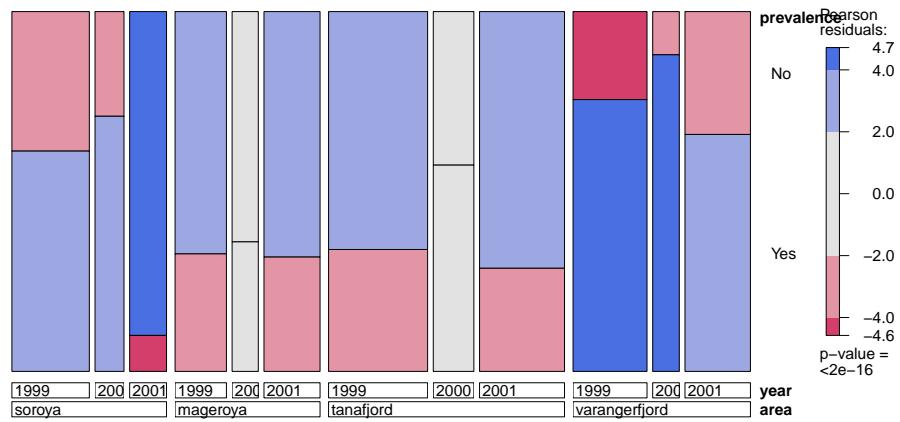


Figure 9.13: Mosaic plot for prevalence against area and year in the CodParasites data, in the doubledecker format. Shading reflects departure from a model in which prevalence is independent of area and year jointly.
fig:cod1-mosaic

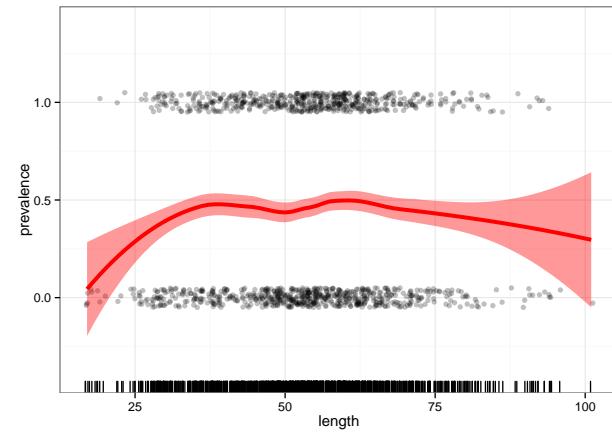


Figure 9.14: Jittered scatterplot of prevalence against length of fish, with loess smooth.
fig:cld-length-prevalence

Note that `facet_grid()` makes it easy to organize the display with separate panels for each area, a technique that could extend to additional factors.

```
> # plot only positive values of intensity
> CPpos <- subset(CodParasites, intensity>0)
> ggplot(CPpos, aes(x=year, y=intensity)) +
+   geom_boxplot(outlier.size=3, notch=TRUE, aes(fill=year), alpha=0.2) +
+   geom_jitter(position=position_jitter(width=0.1), alpha=0.25) +
+   facet_grid(.~area) +
+   scale_y_log10(breaks=c(1,2,5,10,20,50,100, 200)) +
+   theme_bw() + theme(legend.position="none") +
+   labs(y='intensity (log scale)')
```

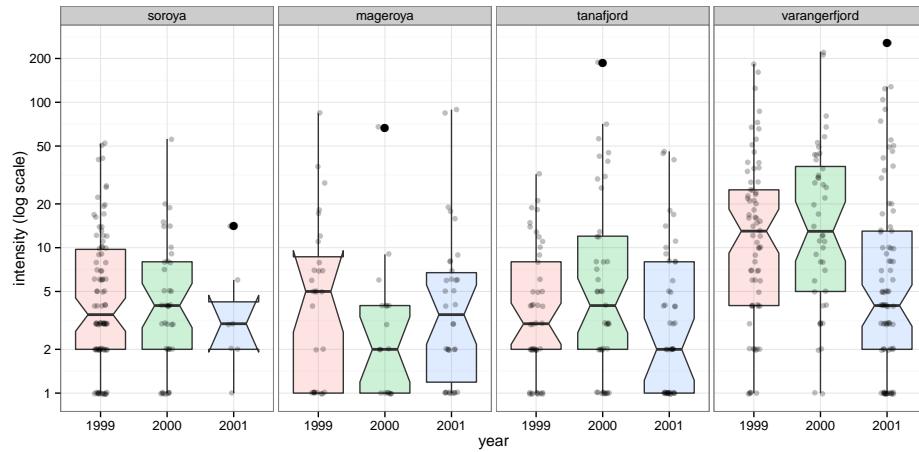


Figure 9.15: Notched boxplots for log (intensity) of parasites by area and year in the CodParasites data. Significant differences in the medians are signaled when the notches of two groups do not overlap.
Fig:cod1_boxplot

Most of these distributions are positively skewed and there are a few high outliers, but probably not more than would be expected in a sample of this size. The positive counts (degree of infection) are also higher in all years in Varangerfjord than other areas. You can also see that the intensity values were generally lower in 2001 than other years.

For the effect of length of fish, we want to know if log (intensity) is reasonably linear on length. A jittered scatterplot produced with `ggplot2` is shown in Figure 9.16. The smoothed loess curve together with the linear regression line show no indication of non-linearity.

```
> ggplot(CPpos, aes(x=length, y=intensity)) +
+   geom_jitter(position=position_jitter(height=.1), alpha=0.25) +
+   geom_rug(position='jitter', sides='b') +
+   scale_y_log10(breaks=c(1,2,5,10,20,50,100, 200)) +
+   stat_smooth(method="loess", color="red", fill="red", size=2) +
+   stat_smooth(method="lm", size=1.5) + theme_bw()
```



9.5.1.1 Fitting models

The simple summary of these exploratory analyses is that both the zero component (prevalence) and non-zero component (intensity) involve an interaction of area and year and at least intensity depends on length. We proceed to fit some count data models.

{ex:cod2}

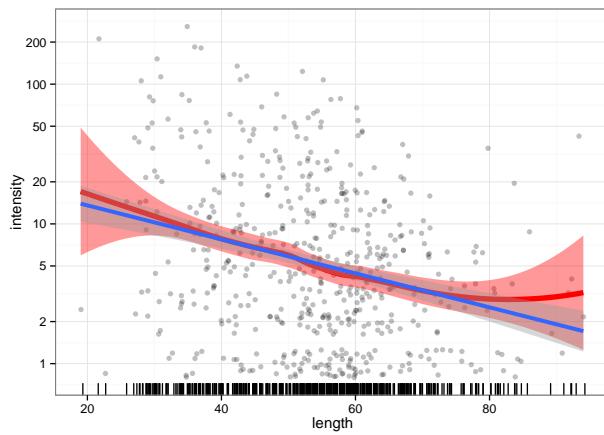


Figure 9.16: Jittered scatterplot of $\log(\text{intensity})$ for the positive counts against length of fish, with loess smooth and linear regression line.
fig:cod-length-scatt

EXAMPLE 9.11: Cod parasites

For a baseline reference, we first fit the standard Poisson and negative-binomial models, not allowing for excess zeros.

```
> library(MASS); library(countreg)
> cp_p <- glm(intensity ~ length + area * year,
+               data = CodParasites, family = poisson)
> cp_nb <- glm.nb(intensity ~ length + area * year,
+                   data = CodParasites)
```

Next, we fit analogous hurdle and zero-inflated models, in each case allowing the non-zero count component to be either Poisson or negative-binomial. The zero components are fit as logistic regressions with the same predictors and the logit link.

```
> cp_hp <- hurdle(intensity ~ length + area * year,
+                    data = CodParasites, dist = "poisson")
> cp_hnb <- hurdle(intensity ~ length + area * year,
+                     data = CodParasites, dist = "negbin")
> cp_zip <- zeroinfl(intensity ~ length + area * year,
+                      data = CodParasites, dist = "poisson")
> cp_znb <- zeroinfl(intensity ~ length + area * year,
+                      data = CodParasites, dist = "negbin")
```

Following Section 9.3.5, we can compare the fit of these models using rootograms. The details of fit of these six models are shown in Figure 9.17.

```
> op <- par(mfrow = c(3, 2))
> countreg::rootogram(cp_p, max = 50, main = "Poisson")
> countreg::rootogram(cp_nb, max = 50, main = "Negative Binomial")
> countreg::rootogram(cp_hp, max = 50, main = "Hurdle Poisson")
> countreg::rootogram(cp_hnb, max = 50, main = "Hurdle Negative Binomial")
> countreg::rootogram(cp_zip, max = 50, main = "Zero-inflated Poisson")
> countreg::rootogram(cp_znb, max = 50, main = "Zero-inflated Negative Binomial")
> par(op)
```

The basic Poisson model of course fits terribly due to the excess zero counts. The hurdle Poisson and zero-inflated Poisson fit the zero counts perfectly, but at the expense of underfitting the counts

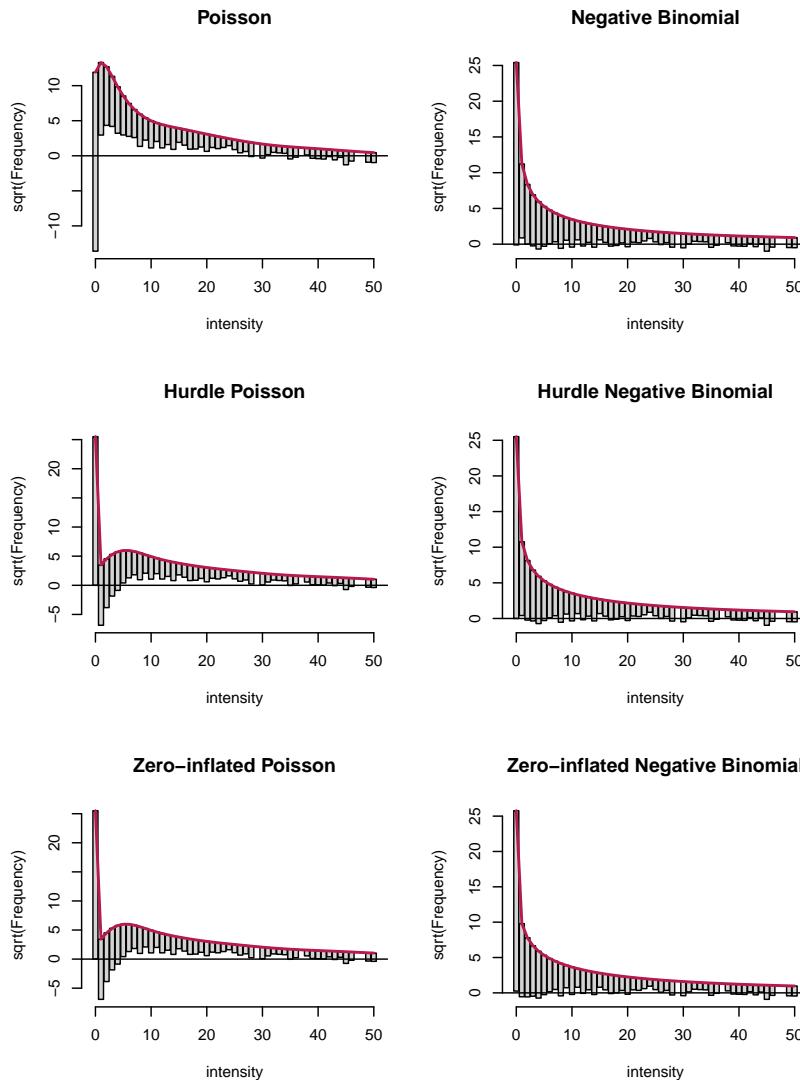


Figure 9.17: Rootograms for six models fit to the CodParasites data fig:cob2-rootograms

for low intensity values. All of the negative binomial models show a reasonable fit (at the scale shown in this plot), and none show a systematic pattern of under/overfitting.

These models are all in different GLM and extended-GLM families, and there are no `anova()` methods for hurdle and zero-inflated models. Each pair of Poisson and negative-binomial models are a nested set, because the Poisson is a special case of the negative-binomial where $\theta \rightarrow \infty$, and so can be compared using likelihood-ratio tests available with `lrtest()` from `lmtest`. However, this cannot be used to compare models of different class, such as a hurdle model vs. a zero-inflated model. (In Figure 9.17, each pair in the same row are nested models, while all other pairs are non-nested.) Yet, they all have `logLik()` methods to calculate their log likelihood, and so `AIC()` and `BIC()` can be used.

```
> vcdExtra::LRstats(cp_p, cp_nb, cp_hp, cp_hnb, cp_zip, cp_znb, sortby="BIC")
Likelihood summary table:
      AIC    BIC   LR Chisq   Df Pr(>Chisq)
cp_p    20378 20444    20352 1178     <2e-16 ***
cp_hp   13688 13820    13636 1165     <2e-16 ***
cp_zip  13687 13819    13635 1165     <2e-16 ***
cp_nb    5031  5102     5003 1178     <2e-16 ***
cp_znb   4955  5092     4901 1164     <2e-16 ***
cp_hnb   4937  5074     4883 1164     <2e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

These show that all the Poisson models fit quite badly, and among the negative-binomial models, the hurdle version, `cp_hnb`, is preferred by both AIC and BIC. If you want to carry out formal tests, `lrtest()` can be used to compare a given Poisson model to its negative-binomial counterpart, which are nested. For example, the test below compares the hurdle Poisson to the hurdle negative-binomial and confirms that the latter is a significant improvement.

```
> library(lmtest)
> lrtest(cp_hp, cp_hnb)

Likelihood ratio test

Model 1: intensity ~ length + area * year
Model 2: intensity ~ length + area * year
#Df LogLik Df Chisq Pr(>Chisq)
1 26    -6818
2 27   -2442  1   8752     <2e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Of greater interest is the difference among the negative-binomial models, that are not nested. As described in Section 9.1.4, these can be compared using Young's test.

```
> library(pscl)

Error in library(pscl): there is no package called 'pscl'

> vuong(cp_nb, cp_hnb)      # nb vs. hurdle nb

Error in eval(expr, envir, enclos): could not find function "vuong"

> vuong(cp_hnb, cp_znb)     # hurdle nb vs znb

Error in eval(expr, envir, enclos): could not find function "vuong"
```

The negative-binomial model is considered to be a closer fit than the hurdle version (because

it is more parsimonious), while the hurdle NB model has a significantly better fit than the zero-inflated NB model. For this example, we continue to work with the hurdle NB model. The tests for individual coefficients in this model are shown below.

```
> summary(cp_hnb)

Call:
hurdle(formula = intensity ~ length + area * year, data = CodParasites,
       dist = "negbin")

Pearson residuals:
    Min      1Q Median      3Q     Max 
-0.696 -0.407 -0.336 -0.108 11.114 

Count model coefficients (truncated negbin with log link):
            Estimate Std. Error z value Pr(>|z|)    
(Intercept)  3.37580   0.39947   8.45 < 2e-16 ***
length      -0.03748   0.00587  -6.38  1.7e-10 ***
areamageroya 0.37898   0.38105   0.99  0.3199    
areatanafjord -0.50480   0.31238  -1.62  0.1061    
areavarangerfjord 0.89159   0.29161   3.06  0.0022 **  
year2000    -0.03957   0.32857  -0.12  0.9041    
year2001     -0.75388   0.68925  -1.09  0.2741    
areamageroya:year2000 -0.63981   0.61667  -1.04  0.2995    
areatanafjord:year2000 1.19387   0.49479   2.41  0.0158 *  
areavarangerfjord:year2000 0.51074   0.47719   1.07  0.2845    
areamageroya:year2001  0.70444   0.82036   0.86  0.3905    
areatanafjord:year2001  0.90824   0.77685   1.17  0.2424    
areavarangerfjord:year2001 0.59838   0.74738   0.80  0.4233    
Log(theta)      -1.49866   0.23904  -6.27  3.6e-10 ***
Zero hurdle model coefficients (binomial with logit link):
            Estimate Std. Error z value Pr(>|z|)    
(Intercept)  0.08526   0.29505   0.29  0.773    
length      0.00693   0.00465   1.49  0.136    
areamageroya -1.32137  0.28526  -4.63  3.6e-06 ***
areatanafjord -1.44918  0.24388  -5.94  2.8e-09 ***
areavarangerfjord 0.30073  0.27111   1.11  0.267    
year2000    0.39507  0.34382   1.15  0.251    
year2001     -2.65201  0.43340  -6.12  9.4e-10 ***
areamageroya:year2000 -0.08034  0.50797  -0.16  0.874    
areatanafjord:year2000 0.87058  0.45027   1.93  0.053 .  
areavarangerfjord:year2000 0.86462  0.59239   1.46  0.144    
areamageroya:year2001  2.73749  0.53291   5.14  2.8e-07 *** 
areatanafjord:year2001  2.71899  0.49949   5.44  5.2e-08 *** 
areavarangerfjord:year2001 2.54144  0.51825   4.90  9.4e-07 *** 
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Theta: count = 0.223
Number of iterations in BFGS optimization: 25
Log-likelihood: -2.44e+03 on 27 Df
```

From the above and from Figure 9.14, it appears that `length` is not important as a linear effect in the submodel for prevalence. A revised model excludes this from the zero formula.

```
> cp_hnb1 <- hurdle(intensity ~ length + area * year | area*year,
+                      data = CodParasites, dist = "negbin")
```

A likelihood-ratio test shows no advantage for the smaller model, however Vuong's test leads to the conclusion that this reduced model is preferable:

```
> lrtest(cp_hnb, cp_hnb1)

Likelihood ratio test

Model 1: intensity ~ length + area * year
Model 2: intensity ~ length + area * year | area * year
#Df LogLik Df Chisq Pr(>Chisq)
```

```

1 27 -2442
2 26 -2443 -1 2.23      0.14

> vuong(cp_hnb, cp_hnbl)

Error in eval(expr, envir, enclos): could not find function "vuong"

Error in detach(package:pscl): invalid 'name' argument

```



9.5.1.2 Model interpretation: Effect plots

Interpreting these models from their coefficients is very difficult because an interaction is present and there are separate submodels for the zero and count components. This task is much easier with effects plots. The `effects` package has methods for any GLM, but cannot handle the extended forms of the zero-inflated and hurdle models.

When the same predictors are used in both submodels, and a standard GLM such as the negative-binomial provides a reasonable fit, you can use the standard `effects` functions to visualize the (total) expected count, which for the zeros would include both the extra zeros and those that derive from the count submodel. For visual interpretation, these will be sufficiently similar, even though the hurdle and zero-inflated models differ with respect to explaining overdispersion and/or excess zeros.

Alternatively, if you want to visualize and interpret the zero and nonzero components separately, perhaps with different predictors, you can fit the implied submodels separately, and then use `effects` functions for the effects in each. These ideas are illustrated in the next example.

{ex:cod3}

EXAMPLE 9.12: Cod parasites

The expected counts for intensity, including both zero and positive counts can be plotted using `effects` for the `cp_nb` NB model. Figure 9.17 gives some confidence that the fitted values are similar to those in the hurdle and zero-inflated versions.

We use `allEffects()` to calculate the effects for the high-order terms—the main effect of `length` and the interaction of `area` and `year`. These could be plotted together by plotting the resulting `eff.nb` object, but we plot them separately to control the plot details. In these plots, the argument `rescale=FALSE` gives plots on the response scale, and we use `ylim` to equate the ranges to make the plots directly comparable. The code below produces Figure ??.

```

> library(effects)

Error: package 'effects' was built before R 3.0.0: please re-install it

> eff.nb <- allEffects(cp_nb)

Error in eval(expr, envir, enclos): could not find function "allEffects"

> plot(eff.nb[1], rescale=FALSE, ylim=c(0, 30),
+       main="NB model: length effect")

Error in plot(eff.nb[1], rescale = FALSE, ylim = c(0, 30), main = "NB model: length effect"): error in evaluating the argument 'x' in selecting a method for function 'plot': Error: object 'eff.nb' not found

> plot(eff.nb[2], rescale=FALSE, ylim=c(0, 30),
+       multiline=TRUE, ci.style='bars',
+       key.args=list(x=.05, y=.95),
+       colors=c("black", "red", "blue"),
+       symbols=15:17, cex=2,
+       main="NB model: area*year effect")

```

```
Error in plot(eff.nb[2], rescale = FALSE, ylim = c(0, 30), multiline = TRUE,
: error in evaluating the argument 'x' in selecting a method for function
'plot': Error: object 'eff.nb' not found
```

This helps to interpret the nature of the area by year effect. The pattern of mean expected intensity of cod parasites is similar in 1999 and 2001, except for the Sørøya area. The results in year 2000 differ mainly in greater intensity in Tanafjord and Varangerfjord. Varangerfjord shows larger infection counts overall, but particularly in year 2000. The effect plot for length on this scale is roughly comparable to the variation in areas and years.

In this example, the submodels for zero and positive counts have substantively different interpretations. To visualize the fitted effects in these submodels using `effects`, first fit the equivalent submodels separately using GLM methods. The following models for prevalence, using the binomial family, and the positive counts for intensity, using `glm.nb()`, give similar fitted results to those obtained from the hurdle negative-binomial model, `cp_hnb` discussed earlier.

```
> cp_zero <- glm(prevalence ~ length + area * year,
+                   data = CodParasites, family=binomial)
> cp_nzero <- glm.nb(intensity ~ length + area * year,
+                      data = CodParasites, subset=intensity>0)
```

We could construct effect plots for each of these submodels, but interest here is largely on the binomial model for the zero counts, `cp_zero`. Effect plots for the terms in this model are shown in Figure ???. Again, we set the `ylim` values to equate the vertical ranges to make the plots comparable.

```
> eff.zero <- allEffects(cp_zero)

Error in eval(expr, envir, enclos): could not find function "allEffects"

> plot(eff.zero[1], ylim=c(-2.5, 2.5),
+       main="Hurdle zero model: length effect")

Error in plot(eff.zero[1], ylim = c(-2.5, 2.5), main = "Hurdle zero model:
length effect"): error in evaluating the argument 'x' in selecting a method
for function 'plot': Error: object 'eff.zero' not found

> plot(eff.zero[2], ylim=c(-2.5, 2.5),
+       multiline=TRUE,
+       key.args=list(x=.05, y=.95),
+       colors=c("black", "red", "blue"),
+       symbols=15:17, cex=2,
+       main="Hurdle zero model: area*year effect")

Error in plot(eff.zero[2], ylim = c(-2.5, 2.5), multiline = TRUE, key.args
= list(x = 0.05, : error in evaluating the argument 'x' in selecting a method
for function 'plot': Error: object 'eff.zero' not found
```

The effect of length on prevalence is slightly increasing, but we saw earlier that this is not significant. For the area-year interaction, the three curves have similar shapes, except for the aberrant value for Sørøya in 2001 and the closeness of the values at Magerøya in all years. Overall, prevalence was highest in 2000, and also in the Varangerfjord samples.



9.5.2 Demand for medical care by the elderly

{sec:glm-case-nmes} A large cross-sectional study was carried out by the U.S. National Medical Expenditure Survey (NMES) in 1987–1988 to assess the demand for medical care, as measured by the number of

physician/non-physician office visits and the number of hospital outpatient visits to a physician/non-physician. The survey was based upon a representative, national probability sample of the civilian non-institutionalized population and individuals admitted to long-term care facilities during 1987. A subsample of 4,406 individuals ages 66 and over, all of whom are covered by Medicare is contained in the *NMES1988* data set in the *AER* package. These data were previously analyzed by Deb and Trivedi (1997) and Zeileis *et al.* (2008), from which this account borrows. The objective of the study and these analyses is to create a descriptive, and hopefully predictive, model for the demand for medical care in this elderly population.

{ex:nmes1}

EXAMPLE 9.13: Demand for medical care

The potential response variables in the *NMES1988* data set form a 2×2 set of the combinations of *place of visit* (office vs. hospital) and (physician vs. non-physician) *practitioner*. Here, we focus on the highest total frequency variable *visits*, recording office visits to a physician. There are quite a few potential predictors, but here we consider only the following:

- *hospital*: number of hospital stays¹²
- *health*: a factor indicating self-perceived health status, with categories "poor", "average" (reference category), "excellent"
- *chronic*: number of chronic conditions
- *gender*
- *school*: number of years of education
- *insurance*: a factor. Is the individual covered by private insurance?

For convenience, these variables are extracted to a reduced data set, *nmes*.

```
> data("NMES1988", package="AER")

Error in find.package(package, lib.loc, verbose = verbose): there is no package
called 'AER'

> nmes <- NMES1988[, c(1, 6:8, 13, 15, 18)]

Error in eval(expr, envir, enclos): object 'NMES1988' not found
```

A quick overview of the response variable, *visits* is shown as simple (unbinned) histograms on the frequency and log(frequency) scales in Figure ???. The zero counts are not as extreme as we have seen in other examples. On the log scale, there is a small, but noticeable spike at 0, followed by a progressive, nearly linear decline, up to about 30 visits.

```
> plot(table(nmes$visits),
+       xlab="Physician office visits", ylab="Frequency")

Error in plot(table(nmes$visits), xlab = "Physician office visits", ylab =
"Frequency"): error in evaluating the argument 'x' in selecting a method for
function 'plot': Error in table(nmes$visits) : object 'nmes' not found

> plot(log(table(nmes$visits)),
+       xlab="Physician office visits", ylab="log(Frequency)")

Error in plot(log(table(nmes$visits)), xlab = "Physician office visits", :
error in evaluating the argument 'x' in selecting a method for function 'plot':
Error in table(nmes$visits) : object 'nmes' not found
```

¹²It is arguable that number of hospitalizations should be regarded as a dependent variable, reflecting another aspect of demand for medical care, rather than as a predictor. We include it here as a predictor to control for its relationship to the outcome *visits*.

However as a benchmark, without taking any predictors into account, there is very substantial overdispersion relative to a Poisson distribution, the variance being nearly 8 times the mean.

```
> with(nmes, c(mean=mean(visits),
+             var=var(visits),
+             ratio=var(visits)/mean(visits)))
```

```
Error in with(nmes, c(mean = mean(visits), var = var(visits), ratio = var(visits)/mean(visits)))
object 'nmes' not found
```

As before, it is useful to precede formal analysis with a variety of exploratory plots. Figure ?? shows a few of these as boxplots, using `cutfac()` to make predictors discrete, and plotting `visits` on a log scale, started at 1. All of these show the expected relationships, e.g., number of office visits increases with numbers of chronic conditions and hospital stays, but decreases with better perceived health status.

```
> op <- par(mfrow=c(1, 3), cex.lab=1.4)
> plot(log(visits+1) ~ cutfac(chronic), data = nmes,
+       ylab = "Physician office visits (log scale)",
+       xlab = "Number of chronic conditions", main = "chronic")
```

```
Error in eval(expr, envir, enclos): object 'nmes' not found
```

```
> plot(log(visits+1) ~ health, data = nmes, varwidth = TRUE,
+       ylab = "Physician office visits (log scale)",
+       xlab = "Self-perceived health status", main = "health")
```

```
Error in eval(expr, envir, enclos): object 'nmes' not found
```

```
> plot(log(visits+1) ~ cutfac(hospital, c(0:2, 8)), data = nmes,
+       ylab = "Physician office visits (log scale)",
+       xlab = "Number of hospital stays", main = "hospital")
```

```
Error in eval(expr, envir, enclos): object 'nmes' not found
```

```
> par(op)
```

Similar plots for insurance and gender show that those with private insurance have more office visits and women slightly more than men.

The relationship with number of years of education could be shown in boxplots by the use of `cutfac(school)`, or with `spineplot()` by making both variables discrete. However, it is more informative (shows the data) to depict this in a smoothed and jittered scatterplot, as in Figure ??.

```
> library(ggplot2)
> ggplot(nmes, aes(x=school, y=visits+1)) +
+   geom_jitter(alpha=0.25) +
+   stat_smooth(method="loess", color="red", fill="red", size=1.5, alpha=0.3) +
+   labs(x="Number of years of education", y="log(Physician office visits+1)") +
+   scale_y_log10(breaks=c(1,2,5,10,20,50,100)) + theme_bw()
```

```
Error in ggplot(nmes, aes(x = school, y = visits + 1)): object 'nmes' not
found
```

As you might expect, there is a small but steady increase in mean office visits with years of education. It is somewhat surprising that there are quite a few individuals with 0 years of education; jittering also shows the greater density of observations at 8 and 12 years.

As in previous examples, a variety of other exploratory plots would be helpful in understanding the relationships among these variables *jointly*, particularly how office visits depends on combinations of two (or more) predictors. Some natural candidates would include mosaic and doubledecker

plots (using `cutfac(visits)`), e.g., as in Figure 9.13, and conditional or faceted versions of the boxplots shown in Figure ??, each stratified by one (or more) additional predictors. These activities are left as exercises for the reader.



9.5.2.1 Fitting models

Most previous analyses of these data have focused on exploring and comparing different types of count data regression models. Deb and Trivedi (1997) compared the adequacy of fit of the negative-binomial, a hurdle NB, and models using finite mixtures of NB models. Zeileis *et al.* (2008) used this data to illustrate hurdle and zero-inflated models using the `countreg` package, while Cameron and Trivedi (1998, 2013) explored a variety of competing models, including 1- and 2-parameter NB models and C -component finite mixture models that can be thought of as generalizations of the 2-component models described in Section 9.4.

In most cases, the full set of available predictors was used, and models were compared using the standard methods for model selection: likelihood-ratio tests for nested models, AIC, BIC and so forth. An exception is Kleiber and Zeileis (2014), who used a reduced set of predictors similar to those employed here, and illustrated the use of rootograms and plots of predicted values for visualizing and comparing fitted models.

This is where model comparison and selection for count data models (and other GLMs) adds another layer of complexity beyond what needs to be considered for classical (Gaussian) linear models, standard logistic regression models and the special case of loglinear models treated earlier. Thus, when we consider and compare different distribution types or link functions, we have to be reasonably confident that the systematic part of the model has been correctly specified (as we noted in Section 9.3), and is the *same* in all competing models, so that any differences can be attributed to the distribution type. However, lack-of-fit may still arise because the systematic part of the model is incorrect.

In short, we cannot easily compare apples to oranges (different distributions with different regressors), but we also have to make sure we have a good apple to begin with. The important questions are:

- Have all important predictors and control variables have been included in the model?
- Are quantitative predictors represented on the correct scale (via transformations or non-linear terms) so their effects are reasonably additive for the linear predictor?
- Are there important interactions among the explanatory variables?

{ex:nmes2}

EXAMPLE 9.14: Demand for medical care

In this example, we start with the all main-effects model of the predictors in the `nmes` data, similar to that considered by Zeileis *et al.* (2008). We first fit the basic Poisson and NB models, as points of reference.

```
> nmes.pois <- glm(visits ~ ., data = nmes, family = poisson)
Error in is.data.frame(data): object 'nmes' not found
> nmes.nbin <- glm.nb(visits ~ ., data = nmes)
Error in is.data.frame(data): object 'nmes' not found
```

A quick check with `lmtest()` shows that the NB model is clearly superior to the standard Poisson regression model as we expect (and also to the quasi-Poisson).

```
> library(lmtest)
> lrtest(nmes.pois, nmes.nbin)

Error in lrtest(nmes.pois, nmes.nbin): object 'nmes.pois' not found
```

The model summary for the NB model below shows the coefficients area all significant. Moreover, the signs of the coefficients are all as we would expect from our exploratory plots. For example, log(visits) increases with number of hospital stays, chronic conditions and education, and is greater for females and those with private health insurance. So, what's not to like?

```
> summary(nmes.nbin)

Error in summary(nmes.nbin): error in evaluating the argument 'object' in
selecting a method for function 'summary': Error: object 'nmes.nbin' not
found
```

This all-main-effects model is relatively simple to interpret, but a more important question is whether it adequately explains the relations of the predictors to the outcome, visits.

Significant interactions among the predictors could substantially change the interpretation of the model, and in the end, could affect policy recommendations based on this analysis. This question turns out to be far more interesting and important than the subtle differences among models for handling overdispersion and zero counts.

One simple way to consider whether there are important interactions among the predictors that better explain patient visits is to get simple tests of the additional contribution of each two-way (or higher-way) interaction using the add1() function. The formula argument in the call below specifies to test the addition of all two-way terms.

```
> add1(nmes.nbin, . ~ .^2, test="Chisq")

Error in add1(nmes.nbin, . ~ .^2, test = "Chisq"): object 'nmes.nbin' not
found
```

From this, we decide to add all two-way interactions among health, hosp and numchron, and also the two-way interaction health:school. Other significant interactions could also be explored, but we don't do this here.

```
> nmes.nbin2 <- update(nmes.nbin,
+                         . ~ . + (health+chronic+hospital)^2
+                         + health:school)

Error in update(nmes.nbin, . ~ . + (health + chronic + hospital)^2 + health:school):
error in evaluating the argument 'object' in selecting a method for function
'update': Error: object 'nmes.nbin' not found
```

This model clearly fits much better than the main effects model, as shown by a likelihood ratio test. The same conclusion would result from anova().

```
> lrtest(nmes.nbin, nmes.nbin2)

Error in lrtest(nmes.nbin, nmes.nbin2): object 'nmes.nbin' not found
```



9.5.2.2 Model interpretation: Effect plots

Complex models with more than a few predictors are difficult to understand and explain, even more so when there are interactions among the predictors. As we have noted previously, effect plots (Fox, 1987, Fox and Andersen, 2006) provide a ready solution.

They have the advantage that each plot shows the correct *partial* relation between the response and the variables in the term shown, controlling (adjusting) for all other variables in the model, as opposed to *marginal* plots that ignore all other variables. From these, it is possible to read an interpretation of a given model effect directly from the effect plot graphs, knowing that all variables not shown in a given graph have been controlled (adjusted for) by setting them equal to average or typical values.

A disadvantage is that these plots show only the predicted (fitted) effects under the *given model* (and not the *data*). If relationships of the response to predictors are nonlinear, or important interactions are not included in the model, you won't see this in an effect plot. We illustrate this point using the results of the main effect NB model, nmes.nbin, as shown in Figure ??.

{ex:nmes2a}

EXAMPLE 9.15: Demand for medical care

```
> library(effects)

Error: package 'effects' was built before R 3.0.0: please re-install it

> plot(allEffects(nmes.nbin), ylab = "Office visits")

Error in plot(allEffects(nmes.nbin), ylab = "Office visits"): error in evaluating
the argument 'x' in selecting a method for function 'plot': Error: could
not find function "allEffects"
```

All of these panels show the expected relations of the predictors to the visits response, and the confidence bands and error bars provide visual tests of the sizes of differences. But they don't tell the full story, because the presence of an important interaction (such as health:chronic) means that the effect of one predictor (health) differs over the values of the other (chronic).

We can see this clearly in effect plots for the model nmes.nbin2 with interactions. For display purposes, it is convenient here to calculate the fitted effects for model terms over a smaller but representative subset of the levels of the integer-valued predictors, using the `xlevels`= argument to `allEffects()`.

```
> eff_nbin2 <- allEffects(nmes.nbin2,
+   xlevels=list(hospital=c(0:3, 6, 8), chronic=c(0:3, 6, 8), school=seq(0, 20, 5)))

Error in eval(expr, envir, enclos): could not find function "allEffects"
```

The result of `allEffects()`, `eff_nbin2`, is a "efflist" object, a list of effects for each *high-order term* in the model. Note that only the terms `gender` and `insurance`, not involved in any interaction, appear as main effects here.

```
> names(eff_nbin2)

Error in eval(expr, envir, enclos): object 'eff_nbin2' not found
```

Plotting the entire "efflist" object gives a collection of plots, one for each high-order term, as in Figure ??, and is handy for a first look. However, the `plot()` methods for effects objects offer greater flexibility when you plot terms individually using additional options. For example, Figure ?? plots the effect for the interaction of `health` and number of chronic conditions with a few optional arguments. See `help(plot.eff, package="effects")` for the available options.

```
> plot(eff_nbin2, "health:chronic", layout=c(3,1),
+       ylab = "Office visits", colors="blue")

Error in plot(eff_nbin2, "health:chronic", layout = c(3, 1), ylab = "Office
visits", : error in evaluating the argument 'x' in selecting a method for
function 'plot': Error: object 'eff_nbin2' not found
```

The default style shown in Figure ?? is a conditional or faceted plot, graphing the response against the X variable with the greatest number of levels, with separate panels for the levels of the other predictor. Alternatively, the effects for a given term can be shown overlaid in a single plot, using the multiline=TRUE argument, as shown in Figure ?? for the two interactions involving health status. Not only is this style more compact, but it also makes direct comparison of the trends for the other variable easier.

```
> plot(eff_nbin2, "health:chronic", multiline=TRUE, ci.style="bands",
+       ylab = "Office visits", xlab="# Chronic conditions",
+       key.args = list(x = 0.05, y = .80, corner = c(0, 0)))

Error in plot(eff_nbin2, "health:chronic", multiline = TRUE, ci.style = "bands",
: error in evaluating the argument 'x' in selecting a method for function
'plot': Error: object 'eff_nbin2' not found

> plot(eff_nbin2, "hospital:health", multiline=TRUE, ci.style="bands",
+       ylab = "Office visits", xlab="Hospital stays",
+       key.args = list(x = 0.05, y = .80, corner = c(0, 0)))

Error in plot(eff_nbin2, "hospital:health", multiline = TRUE, ci.style = "bands",
: error in evaluating the argument 'x' in selecting a method for function
'plot': Error: object 'eff_nbin2' not found
```

From both Figure ?? and the left panel of and Figure ??, it can be seen that for people with poor health status, the relationship of chronic conditions to office visits is relatively flat. For those who view their health status as excellent, their use of office visits is much more strongly related to their number of chronic conditions.

The interaction of perceived health status with number of hospital stays (right panel of Figure ??) shows that the difference in office visits according to health status is mainly important only for those with 0 or 1 hospital stays.

The remaining two interaction effects are plotted in Figure ???. The interaction of hospital stays and number of chronic conditions (left panel of Figure ??) has a clearly interpretable pattern: for those with few chronic conditions, there is a strong positive relationship between hospital stays and office visits. As the number of chronic conditions increases, the relation with hospital stays decreases in slope.

```
> plot(eff_nbin2, "hospital:chronic", multiline=TRUE, ci.style="bands",
+       ylab = "Office visits", xlab="Hospital stays",
+       key.args = list(x = 0.05, y = .70, corner = c(0, 0)))

Error in plot(eff_nbin2, "hospital:chronic", multiline = TRUE, ci.style = "bands",
: error in evaluating the argument 'x' in selecting a method for function
'plot': Error: object 'eff_nbin2' not found

> plot(eff_nbin2, "health:school", multiline=TRUE, ci.style="bands",
+       ylab = "Office visits", xlab="Years of education",
+       key.args = list(x = 0.65, y = .1, corner = c(0, 0)))

Error in plot(eff_nbin2, "health:school", multiline = TRUE, ci.style = "bands",
: error in evaluating the argument 'x' in selecting a method for function
'plot': Error: object 'eff_nbin2' not found
```

Finally, the interaction of `health:school` is shown in the right panel of Figure ???. It can be readily seen that for those of poor health, office visits are uniformly high, and have no relation to years of education. Among those of average or excellent health, office visits increase with years of education in roughly similar ways. \triangle

9.5.2.3 More model wrinkles: Nonlinear terms

Effect plots such as those above are much easier to interpret than tables of fitted coefficients. However, we emphasize that these only reflect the *fitted model*. It might be that the effects of both `hospital` and `chronic` are nonlinear (on the scale of `log(visits)`). In assessing this question, we increase the complexity of model and try to balance parsimony against goodness-of-fit, but also assure that the model retains a sensible interpretation.

{ex:nmes3}

EXAMPLE 9.16: Demand for medical care

The simplest approach is to use `poly(hosp, 2)` and/or `poly(numchron, 2)` to add possible quadratic (or higher power) relations to the model `nmes.nbin2` containing interactions studied above. A slightly more complex model could use `poly(hosp, numchron, degree=2)` for a response-surface model in these variables. A significantly improved fit of such a model is evidence for nonlinearity of the effects of these predictors. This is easily done using `update()`:

```
> nmes.nbin3 <- update(nmes.nbin2, . ~ . + I(chronic^2) + I(hospital^2))

Error in update(nmes.nbin2, . ~ . + I(chronic^2) + I(hospital^2)): error
in evaluating the argument 'object' in selecting a method for function 'update':
Error: object 'nmes.nbin2' not found
```

This model is equivalent to the long-form version below:

```
> nmes.nbin3 <- glm.nb(visits ~ poly(hospital, 2) + poly(chronic, 2) +
+                         insurance + school + gender +
+                         (health+chronic+hospital)^2 + health:school, data = nmes)
```

Comparing these models using `anova()`, we see that there is a substantial improvement in the model fit by including these nonlinear terms. The quadratic model also fits best by AIC and BIC.

```
> anova(nmes.nbin, nmes.nbin2, nmes.nbin3)

Error in anova(nmes.nbin, nmes.nbin2, nmes.nbin3): object 'nmes.nbin' not
found

> vcdExtra::LRstats(nmes.nbin, nmes.nbin2, nmes.nbin3)

Error in vcdExtra::LRstats(nmes.nbin, nmes.nbin2, nmes.nbin3): object 'nmes.nbin'
not found
```

However, effect plots for this model quickly reveal a *substantive* limitation of this approach using polynomial terms. Figure ?? shows one such plot for the interaction of `health` and number of `chronic` conditions that you should compare with Figure ??.

```
> eff_nbin3 <- allEffects(nmes.nbin3,
+   xlevels=list(hospital=c(0:3, 6, 8), chronic=c(0:3, 6, 8), school=seq(0, 20, 5)))

Error in eval(expr, envir, enclos): could not find function "allEffects"

> plot(eff_nbin3, "health:chronic", layout=c(3,1))

Error in plot(eff_nbin3, "health:chronic", layout = c(3, 1)): error in evaluating
the argument 'x' in selecting a method for function 'plot': Error: object
'eff_nbin3' not found
```

The quadratic fits for each level of health in Figure ?? imply that office visits increase with chronic conditions up to a point and then decrease—with a quadratic, what goes up must come down, the same way it went up! This makes no sense here, particularly for those with poor health status. As well, the confidence bands in this figure are uncomfortably wide, particularly at higher levels of chronic conditions, compared to those in Figure ???. The quadratic model is thus preferable statistically and descriptively, but serves less well for explanatory, substantive and predictive goals.

An alternative approach nonlinearity is to use regression splines (as in Example 7.9) or a **generalized additive model** (Hastie and Tibshirani, 1990) for these terms. The latter specifies the linear predictor as a sum of smooth functions,

$$g(\mathcal{E}(y)) = \beta_0 + f_1(x_1) + f_2(x_2) + \cdots + f_m(x_m) .$$

where each $f_j(x_j)$ may be a function with a specified parametric form (for example a polynomial) or may be specified non-parametrically, simply as “smooth functions”, to be estimated by non-parametric means.

In R, a very general implementation of the generalized additive model (GAM) is provided by `gam()` in the `mgcv` package and described in detail by Wood (2006). Particular features of the package are facilities for automatic smoothness selection (Wood, 2004), and the provision of a variety of smooths of more than one variable. This example just scratches the surface of GAM methodology.

In the context of the NB model we are considering here, the analog of model `nmes.nbin3` fitted using `gam()` is `nmes.gamnb` shown below. The negative-binomial distribution can be specified using `family=nb()` when the parameter θ is also estimated from the data (as with `glm.nb()`), or `family=negbin(theta)` when θ is taken as fixed, for example using the value `theta=1.24` available from models `nmes.nbin2`, and `nmes.nbin3`.

```
> library(mgcv)
> nmes.gamnb <- gam(visits ~ s(hospital, k=3) + s(chronic, k=3) +
+                      insurance + school + gender +
+                      (health+chronic+hospital)^2 + health:school,
+                      family=nb(), data = nmes)

Error in is.data.frame(data): object 'nmes' not found
```

The key feature here is the specification of the smooth terms for `s(hospital, k=3)` and `s(chronic, k=3)`, where `k=3` specifies the dimension of the basis used to represent the smooth term. There are many other possibilities with `gam()`, but these are beyond the scope of this example.

We could again visualize the predicted values from this model using effect plots. However a different approach is to visualize the *fitted surface* in 3D, using a range of values for two of the predictors, and controlling for the others.

The `rsm` package provides extensions of the standard `contour()`, `image()` and `persp()` functions for this purpose. The package provides S3 methods (e.g., `persp.lm()`) for "lm" objects, or classes (such as "negbin" and "glm") that inherit methods from `lm`. The calculation of fitted values in these plots use the applicable `predict()` method for the model object. As in effect plots, the remaining predictors are controlled at their average values (or other values specified in the `at` argument).

Two such plots are shown in Figure ???. The left panel shows the interaction of hospital stays and chronic conditions, included in the model with smoothed terms for their main effects. The right panel shows the joint effects of years of education and chronic conditions on office visits, but there is no interaction of these variables in the GAM model `nmes.gamnb`. These plots use `rainbow()` colors to depict the predicted values of office visits. Contours of these values are projected into the bottom or top plane with corresponding color coding.¹³

¹³The vignette `vignette("rsm-plots", package="rsm")` illustrates some of these options.

```

> library(rsm)

Error in library(rsm): there is no package called 'rsm'

> persp(nmes.gamnb, hospital ~ chronic, zlab="log Office visits",
+   col=rainbow(30), contour=list(col="colors", lwd=2),
+   at=list(school=10, health='average'), theta=-60)

Error in persp(nmes.gamnb, hospital ~ chronic, zlab = "log Office visits",
: object 'nmes.gamnb' not found

> persp(nmes.gamnb, school ~ chronic, zlab="log Office visits",
+   col=rainbow(30), contour=list(col="colors", lwd=2, z="top"),
+   at=list(hospital=0.3, health='average'), theta=-60)

Error in persp(nmes.gamnb, school ~ chronic, zlab = "log Office visits", :
object 'nmes.gamnb' not found

```

A simple, credible interpretation of the plot in the left panel is that office visits rise steeply initially with both hospital stays and number of chronic conditions, and then levels off. For those with no chronic conditions, the effect of hospital stays rises to a higher level compared with the effect of chronic conditions among those who have had no hospital stays. However, as we have seen before, the data is quite thin at the upper end of these predictors, and this plot does not show model uncertainty.

The right panel of Figure ?? illustrates the form of model predictions for a term where one variable (`chronic`) is treated as possibly nonlinear using a smooth `s()` effect, the other is treated as linear (`school`), and no interaction between these is included in the model. At each fixed value of `chronic`, increasing education results in greater office visits. At each fixed value of `school`, the number of chronic conditions shows a steep increase in office visits initially, leveling off toward higher levels, but these all have the same predicted shape.

△

9.6 Diagnostic plots for model checking

(sec:glm-diag)

Models, of course, are never true, but fortunately it is only necessary that they be useful.

G. E. P. Box, *Some Problems of Statistics of Everyday Life*, 1979, p. 2

Most of the model diagnostic methods for classical linear models extend in a relatively direct way to GLMs. These include (a) plots of residuals of various types, (b) diagnostic measures and plots of leverage and influence, as well as some (c) more specialized plots (component-plus-residual plots, added-variable plots) designed to show the specific contribution of a given predictor among others in a linear model. These methods were described in Section 7.5 in the context of logistic regression, and most of that discussion is applicable here in wider GLM class.

One additional complication here is that in any GLM we are specifying: (a) the distribution of the random component, which for count data models may also involve a dispersion parameter or other additional parameters; (b) the form of the linear predictor, $\eta = \mathbf{x}^T \boldsymbol{\beta} = \beta_0 + \beta_1 x_1 + \dots$, where all important regressors have been included, and on the right scale; (c) the correct link function, $g(\mu) = \eta$ transforming the conditional mean of the response y to the predictor variables where they have linear relationships.

Thus, there are a lot of things that could go wrong, but the famous quote from George Box should remind us that all models are approximate, and the goal for model diagnosis should be an adequate model, useful for description, estimation or prediction as the case may be. What is most

important is that our models cannot be misleadingly wrong, that is they should not affect substantive conclusions or interpretation.

9.6.1 Diagnostic measures and residuals for GLMs

Estimation of GLMs by maximum likelihood uses an iterative weighted least squares (IWLS) algorithm, and many of the diagnostic measures for these models are close counterparts of their forms for classical linear models. Roughly speaking, these follow from replacing \mathbf{y} and $\hat{\mathbf{y}}$ in least squares diagnostics by a “working response” and $\hat{\eta}$, replacing the residual variance $\hat{\sigma}^2$ by $\hat{\phi}$, and using a weighted form of the Hat matrix.

9.6.1.1 Leverage

Hat values, h_i , measuring leverage or the potential of an observation to affect the fitted model are defined as the diagonal elements of the hat matrix \mathbf{H} , using the weight matrix \mathbf{W} from the final IWLS iteration. This has the same form as in a weighted least squares regression using a fixed \mathbf{W} matrix:

$$\mathbf{H} = \mathbf{W}^{1/2} \mathbf{X} (\mathbf{X}^\top \mathbf{W} \mathbf{X})^{-1} \mathbf{X}^\top \mathbf{W}^{1/2} .$$

In contrast to OLS, the weights depend on the \mathbf{y} values as well as the \mathbf{X} values, so high leverage observations do not necessarily reflect only unusualness in the space of the predictors.

9.6.1.2 Residuals

Several types of residuals can be defined starting from the goodness-of-fit measures discussed in Section 9.1.3. The **raw residual** or **response residual** is simply the difference $y_i - \hat{\mu}_i$ between the observed response y_i and the estimated mean, $\hat{\mu} = g^{-1}(\hat{\eta}_i) = g^{-1}(\mathbf{x}_i^\top \hat{\beta})$.

From this, the **Pearson residual** is defined as

$$r_i^P = \frac{y_i - \hat{\mu}_i}{\sqrt{\hat{\mathcal{V}}(y_i)}} \quad (9.11)$$

and the **deviance residual** is defined as the signed square root of the contribution of observation i to the deviance in Eqn. (9.4).

$$r_i^D = \text{sign}(y_i - \hat{\mu}_i) \sqrt{d_i} \quad (9.12)$$

The Pearson and deviance residuals do not account for dispersion or for differential leverage (which makes their variance smaller), so **standardized residuals** (sometimes called *scaled* residuals) can be calculated as

$$\tilde{r}_i^P = \frac{r_i^P}{\sqrt{\hat{\phi}(1 - h_i)}} \quad (9.13)$$

$$\tilde{r}_i^D = \frac{r_i^D}{\sqrt{\hat{\phi}(1 - h_i)}} \quad (9.14)$$

These have approximate standard normal $\mathcal{N}(0, 1)$ distributions, and will generally have quite similar values (except for small values in $\hat{\mu}$). Consequently, convenient thresholds like $|\tilde{r}_i| > 2$ or $|\tilde{r}_i| > 4$ are useful for identifying unusually large residuals.

Finally, the **studentized residual** (or *deletion* residual) gives the standardized residual that would

result omitting each observation in turn and calculating the change in the deviance. Calculating these exactly would require refitting the model n times, but an approximation is

$$\tilde{r}_i^S = \text{sign}(y_i - \hat{\mu}_i) \sqrt{(\tilde{r}_i^D)^2 + (\tilde{r}_i^P)^2 h_i / (1 - h_i)} . \quad (9.15)$$

From the theory of classical linear models, these provide formal outlier tests for individual observations (Fox, 2008, §11.3) as a *mean-shift* outlier model that dedicates an additional parameter to fit observation i exactly. To correct for multiple testing and a focus on the largest absolute residuals, it is common to apply a Bonferroni adjustment to the p -values of these tests, multiplying them by n .

For a class "glm" object, the function `residuals(object, type)` returns the unstandardized residuals for `type="pearson"` or `type="deviance"`.¹⁴ The standardized versions are obtained using `rstandard()`, again with a `type` argument for the Pearson or deviance flavor. `rstudent()` calculates the studentized deletion residuals.

9.6.1.3 Influence

As discussed in Section 7.5 in the context of logistic regression, influence measures attempt to evaluate the effect that an observation exerts on the parameters, fitted values or goodness-of-fit statistics by comparing a statistic calculated for all the data with the value obtained omitting each observation in turn. Again, approximations are used to estimate these effects without laboriously refitting the model n times.

Overall measures of influence include

- Cook's distance (Eqn. (7.10)), a squared measure of the difference $\hat{\beta} - \hat{\beta}_{(-i)}$ in all p coefficients in the model. The approximation used in `cooks.distance()` is

$$C_i = \frac{\tilde{r}_i h_i}{\hat{\phi} p (1 - h_i)} .$$

This follows Williams (1987), but scales the result by the estimated dispersion $\hat{\phi}$ as an approximate $F_{p,n-p}$ statistic rather than χ_p^2 .

- DFFITS, the standardized signed measure of the difference of the fitted value $\hat{\mu}_i$ using all the data and the value $\hat{\mu}_{(-i)}$ omitting observation i .

{ex:phdpubs5}

EXAMPLE 9.17: Publications of PhD candidates

For models that inherit methods from the "glm" class (including NB models fit using `glm.nb()`), the simplest initial diagnostic plots are provided by the `plot()` method. Figure 9.18 shows the default *regression quartet* of plots for the negative-binomial model `phd.nbin` examined in earlier examples. By default, the `id.n=3` most noteworthy observations are labeled with their row names from the original data set.

```
> op <- par(mfrow=c(2,2), mar=c(4,4,2,1)+.1, cex.lab=1.2)
> plot(phd.nbin)
> par(op)
```

The plot of residuals against predicted values in the upper left panel of Figure 9.18 should show no overall systematic trend for a well-fitting model. The smoothed loess curve in red suggests that this is not the case.

Several functions in the `car` package make these plots more flexibly and with greater control of

¹⁴Other types include raw response residuals (`type="response"`), working residuals (`type="working"`) and partial residuals (`type="partial"`).

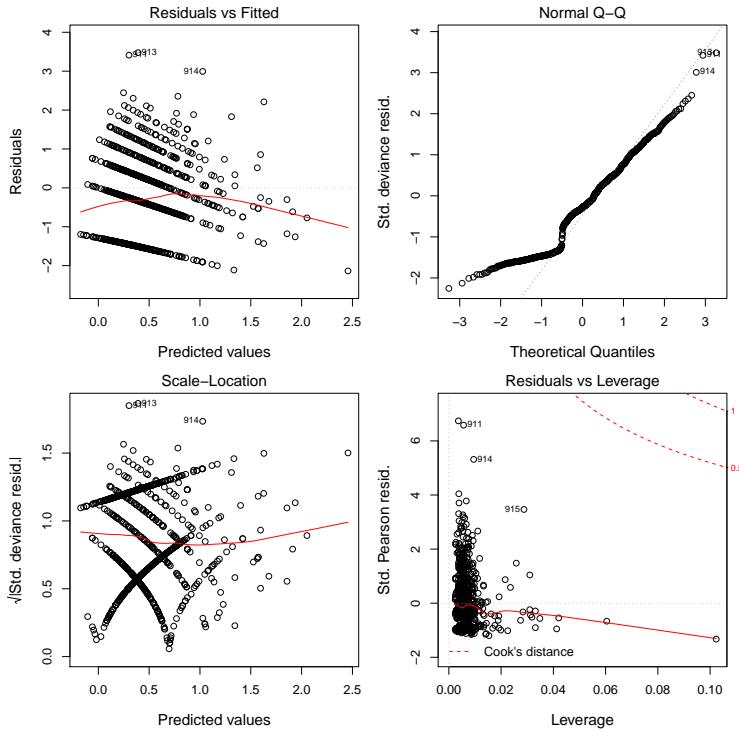


Figure 9.18: Default diagnostic plots for the negative-binomial model fit to the PhdPubs data.
fig:phdpubs5-plot

the details. Figure 9.19 shows the plot of residuals against predicted values two ways. The right panel explains the peculiar pattern of diagonal band of points. These correspond to the different discrete values of the response variable, number of articles published.

```
> library(car)
> residualPlot(phd.nbin, type="rstandard", col.smooth="red", id.n=3)
> residualPlot(phd.nbin, type="rstandard",
+               groups=PhdPubs$articles, key=FALSE, linear=FALSE, smoother=NULL)
```

Other useful plots show the residuals against each predictor. For a good-fitting model, the average residual should not vary systematically with the predictor. As shown in Figure 9.20, `residualPlot()` draws a lowess smooth, and also computes a curvature test for each of the plots by adding a quadratic term and testing the quadratic to be zero.

```
> residualPlot(phd.nbin, "mentor", type="rstudent",
+               quadratic=TRUE, col.smooth="red", col.quad="blue", id.n=3)
> residualPlot(phd.nbin, "phdprestige", type="rstudent",
+               quadratic=TRUE, col.smooth="red", col.quad="blue", id.n=3)
```

In the plot at the left for number of articles by the student's mentor, the curvature is quite pronounced: at high values of `mentor`, nearly all of the residuals are negative, these students publishing fewer articles than would be expected. This would indicate a problem in the scale for `mentor` if there were more observations at the high end; but only about 1.5% points occur for `mentor > 45`, so this can be discounted.

Figure 9.21 gives a better version of the influence plot shown in the lower right panel of Figure 9.18. This plots studentized (deletion) residuals against leverage, showing the value of Cook's distance by the area of the bubble symbol.

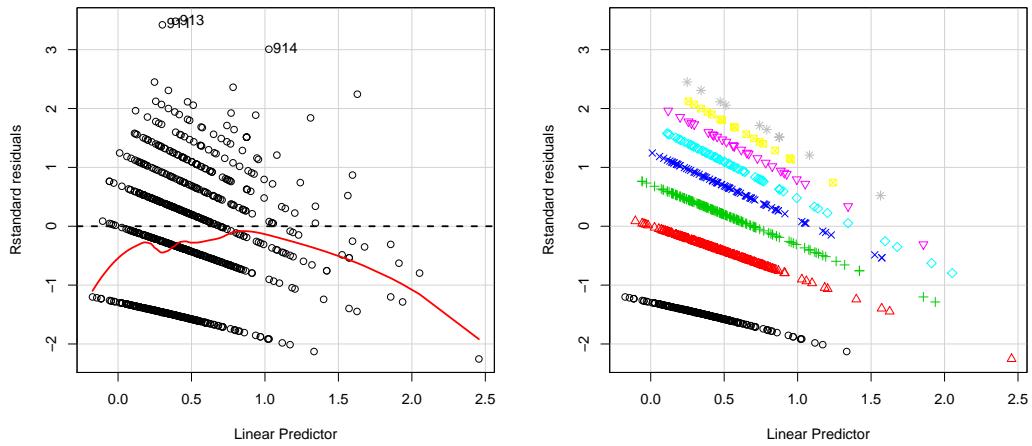


Figure 9.19: Plots of residuals against the linear predictor using `residualPlot()`. The right panel shows that the diagonal bands correspond to different values of the discrete response.
fig:phdpubs5-resplot1

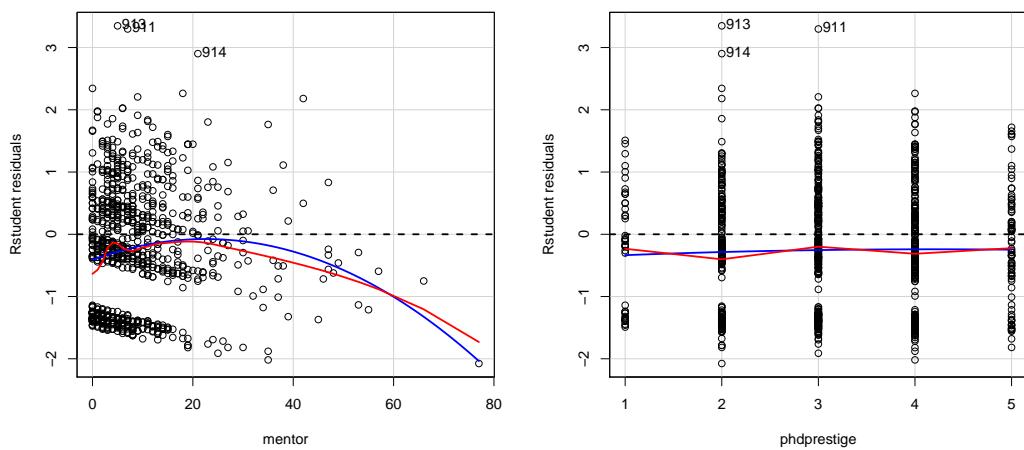


Figure 9.20: Plots of residuals against two predictors in the `phd.nbin` model. Such plots should show no evidence of a systematic trend for a good-fitting model.
fig:phdpubs5-resplot2

```
> influencePlot(phd.nbin)

  StudRes      Hat   CookD
328 -2.0762 0.1023449 0.18325
913  3.3488 0.0036473 0.16652
915  2.1810 0.0287496 0.24345
```

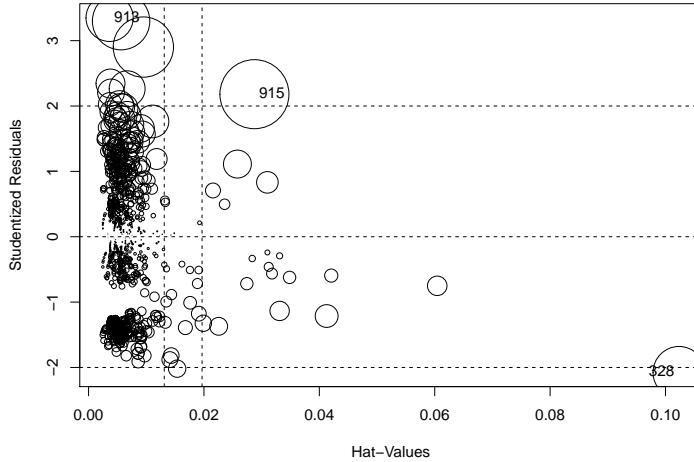


Figure 9.21: Influence plot showing leverage, studentized residuals and Cook's distances for the negative-binomial model fit to the PhdPubs data. Conventional cutoffs for studentized residuals are shown by dashed horizontal lines at ± 2 ; vertical lines show 2 and 3 times the average hat-value.¹⁵

Fig:phdpubs5-influenceplot

Several observations are considered noteworthy, because of one or more of large absolute residual, large leverage or large Cook's distance. `influencePlot()` uses different default rules for point labeling than does the `plot()` method, but provides many options to control the details. Observation 328 stands out as having the largest leverage and a large negative residual; case 913 has the largest absolute residual, but is less influential than case 915.¹⁵

The `outlierTest()` function in `car` gives a formal test of significance of the largest absolute studentized residuals, with a Bonferroni-adjusted p -value accounting for choosing the largest values among n such tests. Individually, case 913 is extreme, but it is not at all extreme among $n = 915$ such tests, each using $\alpha = .05$.

```
> outlierTest(phd.nbin)

No Studentized residuals with Bonferonni p < 0.05
Largest |rstudent|:
  rstudent unadjusted p-value Bonferonni p
913    3.3488          0.00084491     0.77309
```

This example started with the negative-binomial model, the best-fitting from the previous examples. It highlighted a few features of the data not seen previously and worth considering, but doesn't seriously challenge the substantive interpretation of the model. This is what we hope for from model diagnostic plots.

¹⁵The higher case numbers appear in these plots and diagnostics because the data set `PhdPubs` had been sorted by the response, `articles`.



9.6.2 Quantile-quantile and half-normal plots

As we noted above, in theory the standardized and studentized Pearson and deviance residuals have approximate standard normal $\mathcal{N}(0, 1)$ distributions (in large samples) when the fitted model is correct. This suggests a plot of the sorted residuals, $r_{(i)}$, against the corresponding expected values, $z_{(i)}$ an equal-sized sample of size n would have in a normal distribution.¹⁶

If the distribution of the residuals is approximately normal, the points $(r_{(i)}, z_{(i)})$ should lie along a line with unit slope through the origin; systematic or individual departure from this line signals a potential violation of assumptions. The expected values are typically calculated as $z_{(i)} = \Phi^{-1}\{(i - \frac{3}{8})/(n + \frac{1}{4})\}$, where $\Phi^{-1}(\bullet)$ is the inverse normal, or normal quantile function, `qnorm()` in R.

Such plots, called **normal quantile plots** or **normal QQ plots**, are commonly used for GLMs with a quantitative response variable. The upper right panel of Figure 9.18 illustrates the form of such plots produced by `plot()` for a "glm" object.

One difficulty with the default plots is that it is hard to tell to what extent the points deviate from the unit line because there is no visual reference for the line or envelope to indicate expected variability about that line. This problem is easily remedied using `qqPlot()` from `car`.

Figure 9.22 shows the result for the model `phd.nbin`. The envelope lines used here are at the quartiles of the expected normal distribution. They suggest a terrible fit, but, surprisingly, the largest three residuals are within the envelope.

```
> qqPlot(rstudent(phd.nbin), id.n=3,
+         xlab="Normal quantiles", ylab="Studentized residuals")
913 911 914
915 914 913
```

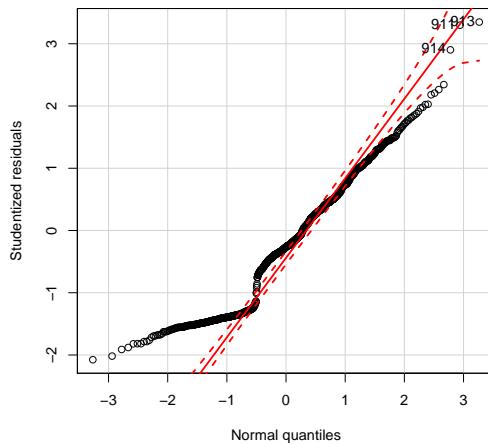


Figure 9.22: Normal QQ plot of the studentized residuals from the NB model for the `PhdPubs` data. The normal-theory reference line and confidence envelope are misleading here! Fig:phdpubs6-qqplot

¹⁶The subscripted notation $r_{(i)}$, and $z_{(i)}$ here denotes an *order statistic*, the i^{th} largest value in a set arranged in increasing order.

For GLMs with discrete responses, such plots are often disappointing, even with a reasonably good-fitting model, because: (a) possible outliers can appear at both the lower and upper ends of the distribution of residuals; (b) the theoretical normal distribution used to derive the envelope may not be well approximated in a given model.

Atkinson (1981, 1987) suggested a more robust and useful version of these QQ plots: half normal plots, with simulated confidence envelopes. The essential ideas are:

- Model departures and outliers are often easier to see for discrete data when the *absolute values* of residuals are plotted, because large positive and negative values are sorted together. This gives the **half-normal plot**, in which the absolute values of residuals, arranged in increasing order, $|r|_{(i)}$, are plotted against $|z|_{(i)} = \Phi^{-1}\{(n + i - \frac{1}{8})/(2n + \frac{1}{2})\}$. All outliers will then appear in the upper right of such a plot, as points separated from the trend of the remaining cells.
- The normal-theory reference line, $|r|_{(i)} = |z|_{(i)}$ and the normal-theory confidence envelope can be replaced by simulating residuals from the assumed distribution, that need not be normal. The reference line is taken as the mean of S simulations and the envelope with $1 - \alpha$ coverage is taken as the $(\alpha/2, 1 - \alpha/2)$ quantiles of their values.
- Specifically, for a GLM, S sets of random observations $y_j, j = 1, 2, \dots, S$ are generated from the fitted model, each with mean $\hat{\mu}$, the fitted values under from the model and with the *same* distribution. In R, this is readily accomplished using the generic `simulate()` function; the random variation around $\hat{\mu}$ uses `rnorm()`, `rpois()`, `rnegbin()`, etc., as appropriate for the family of the model.
- The same model is then fit to each simulated y_j , giving a new set of residuals for each simulation. Sorting their absolute values then gives the simulation distribution used as reference for the observed residuals.

At the time of writing there is no fully general implementation of these plots in R, but the technique is not too difficult and is sufficiently useful to illustrate here.

EXAMPLE 9.18: Publications of PhD candidates

First, calculate the sorted absolute values of the residuals $|r|_{(i)}$ and their expected normal values, $|z|_{(i)}$. The basic plot will be `plot(expected, observed)`.

```
> observed <- sort(abs(rstudent(phd.nbin)))
> n <- length(observed)
> expected <- qnorm((1:n + n - 1/8)/(2*n + 1/2))
```

Then, use `simulate()` to generate $S = 100$ simulated response vectors around the fitted values in the model. Here this uses the negative-binomial random number generator (`rnegbin()`) with the same dispersion value ($\hat{\theta} = 2.267$) estimated in the model. The result, called `sims` here, is a data frame of $n = 915$ rows and $S = 100$ columns, named `sim_1, sim_2, ...`

```
> S <- 100
> sims <- simulate(phd.nbin, nsim=S)
> simdat <- cbind(PhdPubs, sims)
```

The next step is computationally intensive, because we have to fit the NB model $S = 100$ times and a little bit tricky, because we need to use the same model formula as the original, but with the simulated y . We first define a function `resids` to do this for a given y , and then use a loop to calculate them all. To save computing time, the coefficients from the `phd.nbin` model are used as starting values.

```
> # calculate residuals for one simulated data set
> resids <- function(y)
+   rstudent(glm.nb(y ~ female + married + kid5 + phdprestige + mentor,
+     data=simdat, start=coef(phd.nbin)))
> # do them all ...
> simres <- matrix(0, nrow(simdat), S)
> for(i in 1:S) {
+   simres[,i] <- sort(abs(resids(dat[,paste("sim", i, sep="_")]))))
+ }
```

We can then use `apply()` to compute the summary measures defining the center and limits for the simulated confidence interval.

```
> envelope <- 0.95
> mean <- apply(simres, 1, mean)
> lower <- apply(simres, 1, quantile, prob=(1 - envelope)/2)
> upper <- apply(simres, 1, quantile, prob=(1 + envelope)/2)
```

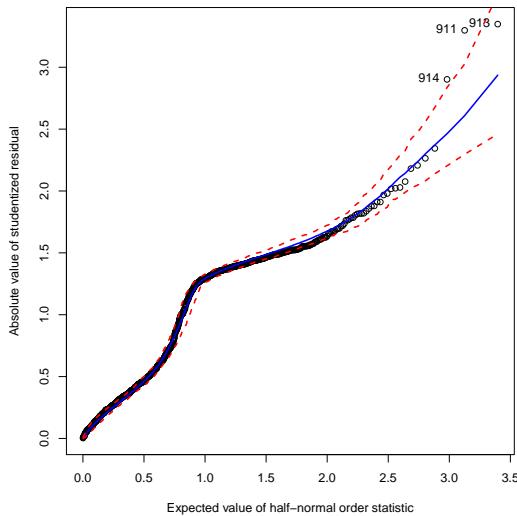


Figure 9.23: Half-normal QQ plot of studentized residuals for the NB model fit to the PhdPubs data. The reference line and confidence envelope reflect the mean and (2.5%, 97.5%) quantiles of the simulation distribution under the negative-binomial model for the same data.

(fig:phd-halfnorm)

Finally, plot the observed against expected absolute residuals as points, and add the lines for the confidence envelope, producing Figure 9.23.

```
> plot(expected, observed,
+       xlab='Expected value of half-normal order statistic',
+       ylab='Absolute value of studentized residual')
> lines(expected, mean, lty=1, lwd=2, col="blue")
> lines(expected, lower, lty=2, lwd=2, col="red")
> lines(expected, upper, lty=2, lwd=2, col="red")
> identify(expected, observed, labels=names(observed), n=3)
```

The shape of the QQ plot in Figure 9.22 shows a peculiar bend at low values and the half-normal version in Figure 9.23 has a peculiar hump in the middle. What could be the cause?

Figure 9.24 shows two additional plots of the studentized residuals that give a clear answer. The density plot at the left shows a strongly bimodal distribution of the residuals. An additional plot at

the right of residuals against the log(response) confirms the guess that the lower mode corresponds to those students who published no articles—excess zeros again!

```
> # examine distribution of residuals
> res <- rstudent(phd.nbin)
> plot(density(res), lwd=2, col="blue",
+       main="Density of studentized residuals")
> rug(res)
>
> # why the bimodality?
> plot(jitter(log(PhdPubs$articles+1), factor=1.5), res,
+       xlab="log (articles+1)", ylab="Studentized residual")
```

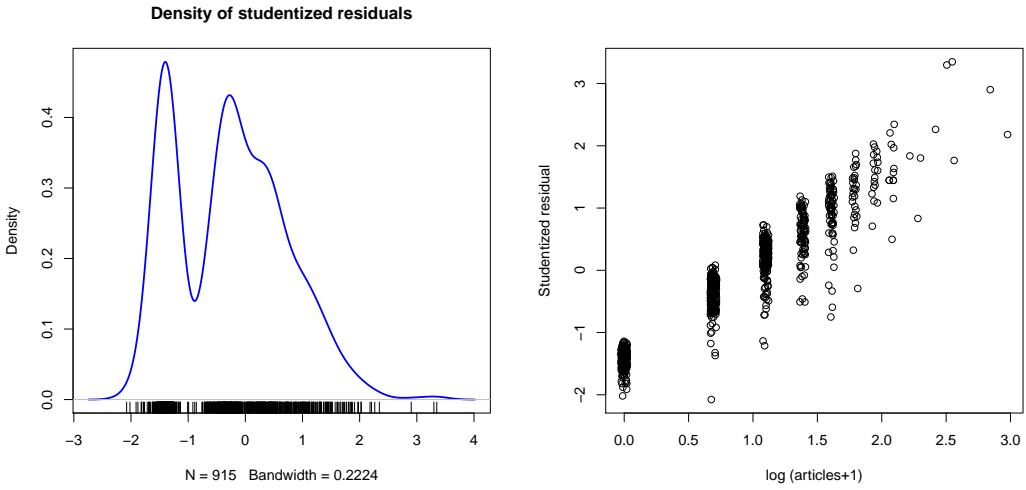


Figure 9.24: fig:phdpubs-res-plots Further plots of studentized residuals. Left: density plot; right: residuals against $\log(\text{articles}+1)$

Now we have something to worry about that *could* affect substantive interpretation or conclusions from this analysis using the NB model, but not accounting for excess zeros. If we believe, following Long (1997), that there is a separate latent class of students who don't publish, it would be sensible to fit a zero-inflated NB model, perhaps with a different subset of predictors for the zero component. The alternative theory of a “hurdle” to a first publication suggests fitting a hurdle model. We leave these as exercises for the reader.



9.7 Multivariate response GLM models

{sec:glm-multiv}

Far better an approximate answer to the right question, which is often vague, than an exact answer to the wrong question, which can always be made precise.

John W. Tukey (1962), *The future of data analysis*

As noted in Section 8.9, in many studies, there may be several response variables along with one or more explanatory variables, and it is useful to try to model some properties of their joint distribution as well as their separate dependence on the predictors. In the current chapter, the case

study (Section 9.5.2) of demand for medical care by the elderly provides a relevant example. There are actually four indicators of medical care, a 2×2 set of (office vs. hospital) place and (physician vs. non-physician) practitioner. That case study analysed only the office visits by physicians.

This section describes a few steps in this direction. To provide some context, we begin with a capsule overview of classical multivariate response models.

In the case of classical linear models with Gaussian error distributions, the model for a univariate response, $\mathbf{y} = \mathbf{X}\boldsymbol{\beta} + \boldsymbol{\epsilon}$, with $\boldsymbol{\epsilon} \sim \mathcal{N}(\mathbf{0}, \Sigma)$ extends quite readily to the ***multivariate linear model*** (MLM) for q response variables, $\mathbf{Y} = \{\mathbf{y}_1, \mathbf{y}_2, \dots, \mathbf{y}_q\}$. The MLM has the form

$$\underset{(n \times q)}{\mathbf{Y}} = \underset{(n \times p)(p \times q)}{\mathbf{X}} \underset{(n \times q)}{\boldsymbol{\beta}} + \underset{(n \times q)}{\mathbf{E}} \quad (9.16) \quad \text{eq:m1m}$$

where \mathbf{Y} is a matrix of n observations on q response variables; \mathbf{X} is a model matrix with columns for p regressors, typically including an initial column of 1s for the regression constant; $\boldsymbol{\beta}$ is a matrix of regression coefficients, one column for each response variable; and \mathbf{E} is a matrix of errors.

It is important to note that:

- The maximum likelihood estimator of $\boldsymbol{\beta}$ in the MLM is equivalent to the result of fitting q separate univariate models for the individual responses and joining the coefficients columwise, giving

$$\hat{\boldsymbol{\beta}} = \{\hat{\beta}_1, \hat{\beta}_2, \dots, \hat{\beta}_q\} = (\mathbf{X}^\top \mathbf{X})^{-1} \mathbf{X}^\top \mathbf{Y}$$

- Procedures for statistical inference (hypothesis tests, confidence intervals), however, take account of the correlations among the responses. Multivariate tests can therefore be more powerful than separate univariate tests under some conditions.
- A unique feature of the MLM stems from the assumption of multivariate normality of the errors, so that each row, $\boldsymbol{\epsilon}_i^\top$ of \mathbf{E} is assumed to be distributed independently, $\boldsymbol{\epsilon}_i^\top \sim \mathcal{N}_q(\mathbf{0}, \Sigma)$, where $\Sigma_{q \times q}$ is the error covariance matrix, constant across observations, like σ^2 in univariate models. Then, the conditional distributions of $\mathbf{y}_j | \mathbf{X}$ are all univariate normal, all bivariate distributions, $\mathbf{y}_j, \mathbf{y}_k | \mathbf{X}$ are bivariate normal, and any linear combination of the conditional ys is univariate normal.
- Consequently, all relationships among the ys can be summarized by correlations and relationships between the ys and xs by linear regressions. These can be visualized using ***data ellipses*** (Friendly *et al.*, 2013) and hypothesis tests in the MLM can be visualized by ellipses using ***hypothesis-error plots*** (Friendly, 2007, Fox *et al.*, 2009).

This generality of the MLM is lost, however, when we move to multivariate response models in the non-Gaussian case. For binomial responses, Section 8.9 described several approaches toward a multivariate logistic regression model that attempt to separate the marginal dependence of each y on the xs from the relationship of the association among the ys on the xs . The bivariate logistic model for (y_1, y_2) for example, was parameterized (see Eqn. (8.33)) in terms of submodels for a logit for each response, $\eta_1 = \mathbf{x}^\top \boldsymbol{\beta}_1$, $\eta_2 = \mathbf{x}^\top \boldsymbol{\beta}_2$ and a submodel for the odds ratio, $\theta_{12} = \mathbf{x}^\top \boldsymbol{\beta}_{12}$.

The situation becomes more difficult for multivariate count data responses, because parametric approaches to their joint distribution (e.g., a multivariate Poisson distribution) given a set of explanatory variables are computationally and analytically intractable. Cameron and Trivedi (2013, Chapter 8) provide a detailed description of the problems and some solutions for the bivariate case, including bivariate Poisson, negative-binomial and hurdle models.

Consequently, only a few special cases have been worked out theoretically, and mostly for the bivariate case. For example, King (1989) described a seemingly unrelated bivariate Poisson model for two correlated count variables. This models the separate linear predictors for y_1 and y_2 as

$$\begin{aligned} g(\mu_1) &= \mathbf{x}_1^\top \boldsymbol{\beta}_1 \\ g(\mu_2) &= \mathbf{x}_2^\top \boldsymbol{\beta}_2 , \end{aligned}$$

with the covariance between y_1 and y_2 represented as ξ . As in the MLM, the coefficients have the same point estimates as in equation-by-equation Poisson models. However, there is a gain in efficiency (reduced standard errors) resulting from a bivariate full-information maximum likelihood solution, and efficiency increases with the covariance ξ between the two count variables.

As a result, for lack of a fully general model for multivariate count data, one simple approach is to employ a method for simultaneous estimation of the equation-by-equation coefficients, accepting some loss of efficiency. This allows for hypothesis tests that may not be most powerful, but provide approximate answers to more interesting questions. We can supplement this with separate analysis of the dependencies among the responses, and how these vary with the explanatory variables.

In R, the VGAM package is the most general available package for analysis of multivariate response GLMs. For multivariate count data, it provides for both Poisson and negative-binomial models. For NB models, the dispersion parameters $\theta_j = \alpha_j^{-1}$ can be allowed to vary with the predictors via a GLM of the form $\log \theta_j = \mathbf{x}^\top \boldsymbol{\gamma}_j$ or can be constrained to be “intercept-only,” $\log \theta_j = \gamma_{0j}$, giving separate global dispersion estimates for each response. In the latter case, the resulting coefficients are the same as fitting a separate model for each response using `glm.nb()`.

{ex:nmes4}

EXAMPLE 9.19: Demand for medical care

In the examples in Section 9.5.2 we considered a variety of models for the number of office visits to physicians (`visits`) as the primary outcome variable in the study of demand for medical care by the elderly. We noted that other indicators of demand included office visits to non-physicians and hospital visits to both physicians and non-physicians. A more complete analysis of this data would consider all four response indicators together.

A special feature of this example is that the four response variables constitute a 2×2 set of the combinations of *place of visit* (office vs. hospital) and (physician vs. non-physician) *practitioner*. These are all counts, and could be transformed to two binary responses according to place and practitioner. Instead, we treat them individually here.

We start by selecting the variables to consider from the *NMES1988* data, giving a new working data set `nmes2`.

```
> data("NMES1988", package="AER")

Error in find.package(package, lib.loc, verbose = verbose): there is no package
called 'AER'

> nmes2 <- NMES1988[, c(1:4, 6:8, 13, 15, 18)]

Error in eval(expr, envir, enclos): object 'NMES1988' not found

> names(nmes2)[1:4]      # responses

Error in eval(expr, envir, enclos): object 'nmes2' not found

> names(nmes2)[-c(1:4)]   # predictors

Error in eval(expr, envir, enclos): object 'nmes2' not found
```

9.7.0.1 Analyzing correlations: HE plots

For purely descriptive purposes, a useful starting point is often an analysis of the $\log(y)$ on the predictor variables using the classical MLM, a rough analog of a multivariate Poisson regression with a log link. Inferential statistics will be biased, but we can use the result to visualize the pairwise linear relations that exist among all responses and all predictors compactly using hypothesis-error (HE) plots (Friendly, 2007).

Zero counts cause problems because the log of zero is undefined, so we add 1 to each y_{ij} in the call to `lm()`. The result is an object of class "mlm".

```
> clog <- function(x) log(x+1)
> nmes.mlm <- lm(clog(cbind(visits, nvisits, ovisits, novisits)) ~ .,
+                   data=nmes2)

Error in is.data.frame(data): object 'nmes2' not found
```

An HE plot provides a visualization of the covariances of effects for the linear hypothesis (H) for each term in a MLM in relation to error covariances (E) using data ellipsoids in the space of dimension q , the number of response variables. The size of each H ellipsoid in relation to the E ellipsoid indicates the strength of the linear relations between the responses and the individual predictors.¹⁷ The orientation of each H ellipsoid shows the direction of the correlations for that term with the response variables. For 1 degree of freedom terms (a covariate or factor with two levels), the corresponding H ellipsoid collapses to a line.

The `heplots` package contains functions for 2D plots (`heplot()`) of pairs of y variables, 3D plots (`heplot3d()`), and all pairwise plots (`pairs()`). We illustrate this here using `pairs()` for the MLM model, giving the plot shown in Figure ??.

```
> library(heplots)

Error in library(heplots): there is no package called 'heplots'

> vlabels <- c("Physician\noffice visits", "Non-physician\n office visits",
+             "Physician\nhospital visits", "Non-physician\nhospital visits")
> pairs(nmes.mlm, factor.means="health", fill=TRUE, var.labels=vlabels)

Error in pairs(nmes.mlm, factor.means = "health", fill = TRUE, var.labels =
vlabels): object 'nmes.mlm' not found
```

The top row in Figure ?? shows the relationship of physician office visits to the other types of medical services. It can be seen that chronic conditions and hospital stays are positively correlated with both responses, as they also are in all other pairwise plots. Having private health insurance is positively related to some of these outcomes, and negatively to others. Except for difficulties with overlapping labels and the obvious violation of statistical assumptions of the MLM here, such plots give reasonably useful overview of the relationships among the y and x variables.

9.7.0.2 Analyzing associations: Odds ratios and fourfold plots

In the analysis below, we first attempt to understand the association among these response variables and how these associations relate to the explanatory variables. It is natural to think of this in terms of the (log) odds ratio of a visit to a physician vs. a non-physician, given that the place is in an office as opposed to a hospital. Following this, we consider some multivariate negative binomial models relating these counts to the explanatory variables.

In order to treat the four response variables as a single response (`visit`), distinguished by `type`, it is necessary to reshape the data from a wide format to a long format with four rows for each input observation.

```
> vars <- colnames(nmes2)[1:4]

Error in is.data.frame(x): object 'nmes2' not found
```

¹⁷When the errors, E in Eqn. (9.16) are approximately multivariate normal, the H ellipsoid provides a visual test of significance: the H ellipsoid projects outside the E ellipsoid if and only if Roy's test is significant at a chosen α level.

```
> nmes.long <- reshape(nmes2,
+   varying = vars,
+   v.names = "visit",
+   timevar = "type",
+   times = vars,
+   direction = "long",
+   new.row.names = 1:(4*nrow(nmes2)))
```

Error in reshape(nmes2, varying = vars, v.names = "visit", timevar = "type", : object 'vars' not found

Then, the `type` variable can be used to create two new variables, `practitioner` and `place` corresponding to the distinctions among visits. While we are at it, we create factors for the two of the predictors.

```
> nmes.long <- nmes.long[order(nmes.long$id),]
```

Error in eval(expr, envir, enclos): object 'nmes.long' not found

```
> nmes.long <- transform(nmes.long,
+   practitioner = ifelse(type %in% c("visits", "ovisits"),
+     "physician", "nonphysician"),
+   place = ifelse(type %in% c("visits", "nvisits"), "office", "hospital"),
+   hospf = cutfac(hospital, c(0:2, 8)),
+   chronicf = cutfac(chronic))
```

Error in transform(nmes.long, practitioner = ifelse(type %in% c("visits", : object 'nmes.long' not found

Then, we can use `xtabs()` to create a frequency table of `practitioner` and `place` classified by any one or more of these factors. For example, the total number of visits of the four types is given by

```
> xtabs(visit ~ practitioner + place, data=nmes.long)
```

Error in terms.formula(formula, data = data): object 'nmes.long' not found

From this, we can calculate the odds ratio and visualize the association with a fourfold or mosaic plot. More generally, by including more factors in the call to `xtabs()`, we can calculate and visualize how the *conditional* association varies with these factors. For example, Figure ?? shows fourfold plots conditioned by health status. It can be seen that there is a strong positive association, except for those with excellent health: people are more likely to see a physician in an office visit, and a non-physician in a hospital visit. The corresponding log odds ratios are shown numerically using `loddsratio()`.

```
> library(vcdExtra)
> fourfold(xtabs(visit ~ practitioner + place + health, data=nmes.long),
+   mfrw=c(1,3))
```

Error in terms.formula(formula, data = data): object 'nmes.long' not found

```
> loddsratio(xtabs(visit ~ practitioner + place + health, data=nmes.long))
```

Error in terms.formula(formula, data = data): object 'nmes.long' not found

Going further, we can condition by more factors. Figure ?? shows the fourfold plots conditioned by the number of chronic conditions (in the rows) and the combinations of gender and private insurance (columns).

```
> tab <- xtabs(visit ~ practitioner + place + gender + insurance + chronicf,
+               data=nmes.long)

Error in terms.formula(formula, data = data): object 'nmes.long' not found

> fourfold(tab, mfcoll=c(4,4))

Error in fourfold(tab, mfcoll = c(4, 4)): object 'tab' not found
```

The systematic patterns seen here are worth exploring further by graphing the log odds ratios directly. The call `as.data.frame(loddsratio(tab))` converts the result of `loddsratio(tab)` to a data frame with factors for these variables and variables LOR and ASE containing the estimated log odds ratio ($\hat{\vartheta}$) and its asymptotic standard error ($ASE(\hat{\vartheta})$). Figure ?? shows the plot of these values as line graphs with associated 95% error bars produced using `ggplot2`.

```
> lodds.df <- as.data.frame(loddsratio(tab))

Error in loddsratio(tab): object 'tab' not found

> library(ggplot2)
> ggplot(lodds.df, aes(x=chronicf, y=LOR,
+                         ymin=LOR-1.96*ASE, ymax=LOR+1.96*ASE,
+                         group=insurance, color=insurance)) +
+   geom_line(size=1.2) + geom_point(size=3) +
+   geom_linerange(size=1.2) +
+   geom_errorbar(width=0.2) +
+   geom_hline(yintercept=0) +
+   facet_grid(. ~ gender, labeller=label_both) +
+   labs(x="Number of chronic conditions",
+        y="log odds ratio (physician|place)") +
+   theme_bw()

Error in ggplot(lodds.df, aes(x = chronicf, y = LOR, ymin = LOR - 1.96 * : object 'lodds.df' not found
```

It can be seen that for those with private insurance, the log odds ratios are uniformly positive, but males and females exhibit a somewhat different pattern over number of chronic conditions. Among those with no private insurance, the log odds ratios generally increase over number of chronic conditions, except for females with 3 or more such conditions.

Beyond this descriptive analysis, you can test hypotheses about the effects of the predictors on the log odds ratios using a simple ANOVA model. Under the null hypothesis, $H_0 : \vartheta_{ijk\dots} = 0$, the $\hat{\vartheta}$ are each distributed normally, $\mathcal{N}(0, ASE(\hat{\vartheta}))$, so a weighted ANOVA can be used to test for differences according to the predictors. This analysis gives the results below.

```
> lodds.mod <- lm(LOR ~ (gender + insurance + chronicf)^2,
+                   weights=1/ASE^2, data=lodds.df)

Error in is.data.frame(data): object 'lodds.df' not found

> anova(lodds.mod)

Error in anova(lodds.mod): object 'lodds.mod' not found
```

As might be expected from the graph in Figure ??, having private insurance is a primary determinant of the decision to seek an office visit with a physician, but this effect interacts slightly according to number of chronic conditions and gender.



9.7.0.3 Fitting and testing multivariate count data models

With a multivariate response, `vglm()` in the `VGAM` package estimates the separate coefficients for each response jointly. A special feature of this formulation is that constraints can be imposed to force the coefficients for a given term in a model to be the same for all responses. A likelihood-ratio test against the unconstrained model can then be used to test for differences in the effects of predictors across the response variables.

This is achieved by formulating the linear predictor as a sum of terms,

$$\eta(\mathbf{x}) = \sum_{k=1}^p \mathbf{H}_k \boldsymbol{\beta}_k \mathbf{x}_k$$

where $\mathbf{H}_1, \dots, \mathbf{H}_p$ are known full-rank constraint matrices. With no constraints the \mathbf{H}_k are identity matrices \mathbf{I}_q for all terms. With `vglm()`, the constraint matrices for a given model are returned using `constraints()`, and can be set for a new, restricted model using the `constraints` argument. To constrain the coefficients for a term k to be equal for all responses, use $\mathbf{H}_k = \mathbf{1}_q$, a unit vector.

More general Wald tests of hypotheses can be carried out without refitting using `linearHypothesis()` in the `car` package. These include (a) joint tests that a subset of predictors for a given response have null effects; (b) across-response tests of equality of coefficients for one or more model terms.

{ex:nmes5}

EXAMPLE 9.20: Demand for medical care

In the examples in Section 9.5.2, we described a series of increasingly complex models for physician office visits, including interactions and non-linear terms. The multivariate case is computationally more intensive, and estimation can break down in complex models. We can illustrate the main ideas here using the multivariate analog of the simple main effects model discussed in Example 9.14.

Using `vglm()`, the response variables are specified as the matrix form \mathbf{Y} using `cbind()` on the left-hand side of the model formula. The right-hand side, `~ .` here specifies all other variables as predictors. `family = negbinomial` uses the NB model for each y_j , with an intercept-only model for the dispersion parameters by default.

```
> nmes2.nbin <- vglm(cbind(visits, nvisits, ovisits, novisits) ~ .,
+                      data = nmes2, family = negbinomial)

Error in is.data.frame(data): object 'nmes2' not found
```

The estimated parameters from this model are returned by the `coef()` method as pairs of columns labeled `log(mu)`, `logsize` for each response. For example, the parameters for the `visits` response are in the first two columns, and are the same as those estimated for the model `nmes.nbin` using `glm.nb()`.

```
> # coefficients for visits
> coef(nmes2.nbin, matrix=TRUE) [,c(1,2)]

Error in coef(nmes2.nbin, matrix = TRUE): error in evaluating the argument
'object' in selecting a method for function 'coef': Error: object 'nmes2.nbin'
not found

> # theta for visits
> exp(coef(nmes2.nbin, matrix=TRUE) [1,2])

Error in coef(nmes2.nbin, matrix = TRUE): error in evaluating the argument
'object' in selecting a method for function 'coef': Error: object 'nmes2.nbin'
not found
```

The `log(mu)` coefficients for all four response variables are shown below.

```
> coef(nmes2.nbin, matrix=TRUE) [,c(1,3,5,7)]  
  
Error in coef(nmes2.nbin, matrix = TRUE): error in evaluating the argument  
'object' in selecting a method for function 'coef': Error: object 'nmes2.nbin'  
not found
```

We notice that the coefficients for hospital and chronic have values with the same signs for all four responses. If it is desired to test the hypothesis that their coefficients are all the same for each of these predictors, first extract the H matrices for the unconstrained model using `constraints()`.

```
> clist <- constraints(nmes2.nbin, type = "term")  
  
Error in constraints(nmes2.nbin, type = "term"): error in evaluating the argument  
'object' in selecting a method for function 'constraints': Error: object  
'nmes2.nbin' not found  
  
> clist$hospital[c(1,3,5,7),]  
  
Error in eval(expr, envir, enclos): object 'clist' not found
```

Then, reset the constraints for these terms to be unit vectors, forcing them to be all equal.

```
> clist2 <- clist  
  
Error in eval(expr, envir, enclos): object 'clist' not found  
  
> clist2$hospital <- cbind(rowSums(clist$hospital))  
  
Error in is.data.frame(x): object 'clist' not found  
  
> clist2$chronic <- cbind(rowSums(clist$chronic))  
  
Error in is.data.frame(x): object 'clist' not found  
  
> clist2$hospital[c(1,3,5,7), 1, drop=FALSE]  
  
Error in eval(expr, envir, enclos): object 'clist2' not found
```

Now, fit the same model as before, but using the constraints in `clist2`.

```
> nmes2.nbin2 <- vglm(cbind(visits, nvisits, ovisits, novisits) ~ ., data = nmes2,  
+                         constraints = clist2,  
+                         family = negbinomial(zero = NULL))  
  
Error in is.data.frame(data): object 'nmes2' not found
```

The coefficients for the constrained model are shown below. As you can see, the coefficients for hospital and chronic have the same estimates for all four responses.

```
> coef(nmes2.nbin2, matrix=TRUE) [,c(1,3,5,7)]  
  
Error in coef(nmes2.nbin2, matrix = TRUE): error in evaluating the argument  
'object' in selecting a method for function 'coef': Error: object 'nmes2.nbin2'  
not found
```

A likelihood-ratio test prefers the reduced model with equal coefficients for these two predictors. The degrees of freedom for this test (6) is the number of constrained parameters in the smaller model.

```
> lrtest(nmes2.nbin, nmes2.nbin2)

Error in lrtest(nmes2.nbin, nmes2.nbin2): error in evaluating the argument
'object' in selecting a method for function 'lrtest': Error: object 'nmes2.nbin'
not found
```

Alternatively, these tests can be performed as tests of linear hypotheses (see Section 9.1.2) on the coefficients B from the original model without refitting. Using `linearHypothesis()`, a hypothesis matrix L specifying equality of the coefficients for a given predictor can be easily generated using a character vector of the coefficient names.

```
> lh <- paste("hospital:", 1:3, " = ", "hospital:", 2:4, sep="")
> lh
[1] "hospital:1 = hospital:2" "hospital:2 = hospital:3"
[3] "hospital:3 = hospital:4"
```

Using `lh` as the `linear.hypothesis` argument then gives the following result for the coefficients of `hospital`, rejecting the hypothesis that they are all equal across response variables.

```
> car::linearHypothesis(nmes2.nbin, lh)

Error in car::linearHypothesis(nmes2.nbin, lh): object 'nmes2.nbin' not found
```

△

To pursue this analysis further, you could investigate whether any interactions of these effects were interesting and important as in Example 9.14, but now for the multivariate response variables.

To interpret a given model visually, you could use effect plots for the terms predicting each of the responses, as in Example 9.15. The `effects` package cannot handle models fit with `VGAM` directly, but you can use `glm()` or `glm.nb()` to fit the equivalent submodels for each response separately, and then use the `plot(Effect())` methods to display the effects for interesting terms. Figure 9.25 shows one such plot, for the effects of health status on each of the four response variables.

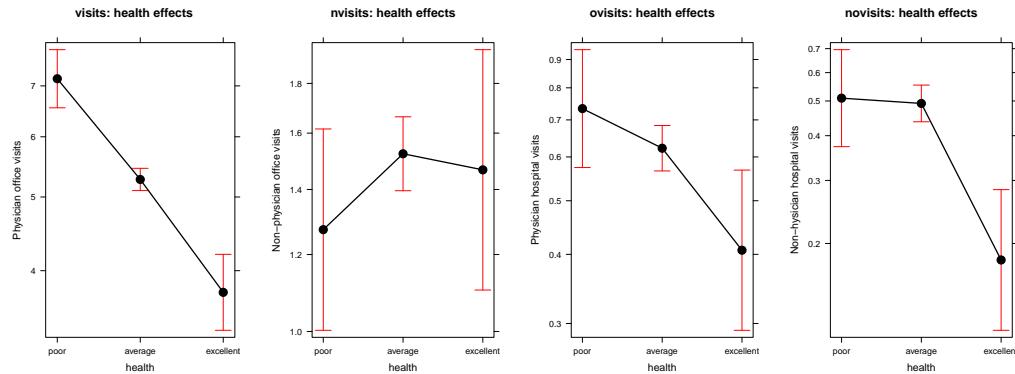


Figure 9.25: Effect plots for the effects of health status on the four response variables in the `nmes2` data.

9.8 Chapter summary

- The generalized linear model extends the familiar classical linear models for regression and ANOVA to encompass models for discrete responses and continuous responses for which the assumption of normality of errors is untenable.
- It does this by retaining the idea of a *linear predictor*— a linear function of the regressors, $\eta_i = \mathbf{x}^T \boldsymbol{\beta}$, but then allowing:
 - a *link function*, $g(\bullet)$ connecting the linear predictor η_i to the mean, $\mu_i = \mathcal{E}(y_i)$, of the response variable, so that $g(\mu_i) = \eta_i$. The link function formalizes the more traditional approach of analyzing an ad-hoc transformation of y , such as $\log(y)$, \sqrt{y} , y^2 , or Box-Cox (Box and Cox, 1964) transformations y^λ to determine an empirical optimal power transformation.
 - a *random component*, specifying the conditional distribution of $y_i | \mathbf{x}_i$ as any member of the exponential family, including the normal, binomial, Poisson, gamma and other distributions.
- For the analysis of discrete response variables, and count data in particular, a key feature of the GLM is recognition of a *variance function* for the conditional variance of y_i , not forced to be constant, but rather allowed to depend on the mean μ_i and possibly a dispersion parameter, ϕ .
- From this background, we focus on GLMs for discrete count data response variables that extend considerably the loglinear models for contingency tables treated in Chapter 8. The Poisson distribution with a log link function is an equivalent starting point, however, count data GLMs often exhibit overdispersion in relation to the Poisson assumption that the conditional variance is the same as the mean, $\mathcal{V}(y_i | \eta_i) = \mu_i$.
 - One simple approach to this problem is the quasi-Poisson model, that estimates the dispersion parameter ϕ from the data, and uses this to correct standard errors and inferential tests.
 - Another is the wider class of negative-binomial models that allow a more flexible mean-variance function such as $\mathcal{V}(y_i | \eta_i) = \mu_i + \alpha\mu_i^2$.
- In practical application, many sets of empirical count data also exhibit a greater prevalence of zero counts than can be fit well using (quasi-) Poisson or negative-binomial models. Two simple extensions beyond the GLM class are
 - zero-inflated models, that posit a latent class of observations that always yield $y_i = 0$ counts, among the rest that have a Poisson or negative-binomial distribution including some zeros;
 - hurdle (or zero-altered) models, with one submodel for the zero counts and a separate submodel for the positive counts.
- Data analysis and visualization of count data therefore requires flexible tools and graphical methods. Some useful exploratory methods include jittered scatterplots and boxplots of $\log(y)$ against predictors enhanced by smoothed curves and trend lines, spine plots and conditional density plots. Rootograms are quite helpful in visualizing the goodness-of-fit of count data models.
- Effect plots provide a convenient visual display of the high-order terms in a possibly complex GLM. They show the fitted values of the linear predictor $\hat{\eta}^* = \mathbf{X}^* \hat{\boldsymbol{\beta}}$, using a score matrix \mathbf{X}^* that varies the predictors in a given term over their range while holding all other predictors constant. It is important to recognize, however, that like any model summary these show only the fitted effects under a given model, not the data.

- Model diagnostic measures (leverage, residuals, Cook's distance, etc.) and plots of these provide important ancillary information about the adequacy of a given model as a summary of relationships in the data. These help to detect problems of violations of assumptions, unusual or influential observations or patterns that suggest that an important feature has not been accounted for.
- For multivariate response count data, there is no fully general theory as there is for the MLM with multivariate normality assumed for the errors. Nevertheless, there is a lot one can do to analyse such data combining the ideas of estimation for the separate responses with analysis of dependencies among the responses, conditioned by the explanatory variables.

9.9 Further reading

9.10 Lab exercises

{lab:9.1}

Exercise 9.1 ? studied the mating behavior of elephants over 8 years in Amboseli National Park, Kenya. A focal aspect of the study concerned the mating success of males in relation to age, since larger males tend to be more successful in mating. Her data were used by ?, Chapter 22 as a case study, and are contained in the `Sleuth2` package (Ramsey *et al.*, 2012) as `case2201`.

For convenience, rename this to `elephants`, and study the relation between `Age` (at the beginning of the study) and number of successful `Matings` for the 41 adult male elephants observed over the course of this study, ranging in age from 27–52.

```
> data("case2201", package="Sleuth2")

Error in find.package(package, lib.loc, verbose = verbose): there is no package
called 'Sleuth2'

> elephants <- case2201

Error in eval(expr, envir, enclos): object 'case2201' not found

> str(elephants)

Error in str(elephants): object 'elephants' not found
```

- Create some exploratory plots of `Matings` against `Age` in the styles illustrated in this chapter. To do this successfully, you will have to account for the fact that `Matings` has a range of only 0–9, and use some smoothing methods to show the trend.
- Repeat (a) above, but now plotting $\log(\text{Matings}+1)$ against `Age` to approximate a Poisson regression with a log link and avoid problems with the zero counts.
- Fit a linear Poisson regression model for `Matings` against `Age`. Interpret the fitted model *verbally* from a graph of predicted number of matings and/or from the model coefficients. (*Hint:* Using `Age-27` will make the intercept directly interpretable.)
- Check for non-linearity in the relationship by using the term `poly(Age, 2)` in a new model. What do you conclude?
- Assess whether there is any evidence of overdispersion in these data by fitting analogous quasi-Poisson and negative-binomial models.

{lab:9.2}

Exercise 9.2 The data set `quine` in MASS gives data on absenteeism from schools in rural New South Wales, Australia. 146 children were classified by ethnic background (`Eth`), age (`Age`, a factor), Sex and Learner status (`Lrn`) and the number of days absent (`Days`) from school in a particular school year was recorded.

- Fit the all main-effects model in the Poisson family and examine the tests of these effects using `summary()` and `car::Anova()`. Are there any terms that should be dropped according to these tests?
- Re-fit this model as a quasi-Poisson model. Is there evidence of overdispersion? Test for overdispersion formally, using `dispersiontest()` from `AER`.
- Carry out the same significance tests and explain why the results differ from those for the Poisson model.

{lab:9.3}

Exercise 9.3 Male double-crested cormorants use advertising behavior to attract females for breeding. The `Cormorants` data set in `vcdExtra` gives some results from a study by Meagan Mc Rae (?) on counts of advertising males observed two or three times a week at six stations in a tree-nesting colony for an entire breeding season. The number of advertising birds was counted and these observations were classified by characteristics of the trees and nests. The goal was to determine how this behavior varies temporally over the season and spatially over observation stations, as well as with characteristics of nesting sites. The response variable is `count` and other predictors are shown below. See `help(Cormorants, package="vcdExtra")` for further details.

```
> data("Cormorants", package="vcdExtra")
> car::some(Cormorants)
```

	category	week	station	nest	height	density	tree_health	count
17	Pre	1	C3	no	mid	few	dead	8
40	Pre	1	B1	full	mid	few	healthy	1
92	Pre	2	B1	no	high	few	healthy	4
118	Pre	3	C2	no	mid	few	dead	3
130	Pre	3	C4	no	high	few	dead	6
160	Incubation	4	C2	partial	high	few	healthy	1
169	Incubation	4	C3	partial	high	few	dead	2
190	Incubation	4	B1	full	high	few	healthy	2
191	Incubation	4	B2	no	low	few	healthy	5
300	Incubation	8	B2	no	high	few	healthy	4

- Using the methods illustrated in this chapter, make some exploratory plots of the number of advertising birds against week in the breeding season, perhaps stratified by another predictor, like tree height, nest condition, or observation station. To see anything reasonable, you should plot `count` on a log (or square root) scale, jitter the points, and add smoothed curves. The variable `category` breaks the weeks into portions of the breeding season, so adding vertical lines separating those will be helpful for interpretation.
- Fit a main-effects Poisson GLM to these data and test the terms using `Anova()` from the `car` package.
- Interpret this model using an effects plot.
- Investigate whether the effect of `week` should be treated as linear in the model. You could try using a polynomial term like `poly(week, degree)` or perhaps better, using a natural spline term like `ns(week, df)` from the `splines` package.
- Test this model for overdispersion, using either a `quasipoisson` family or `dispersiontest()` in `AER`.

{lab:9.4}

Exercise 9.4 For the `CodParasites` data, recode the `area` variable as an ordered factor as suggested in footnote 10. Test the hypotheses that prevalence and intensity of cod parasites is linearly related to `area`.

{lab:9.5}

Exercise 9.5 In Example 9.10, we ignored other potential predictors in the `CodParasites` data: `depth`, `weight`, `length`, `sex`, `stage`, and `age`. Use some of the graphical methods shown in this case study to assess whether any of these are related to prevalence and intensity.

{lab:9.6}

Exercise 9.6 The analysis of the `PhdPubs` data in the examples in this chapter were purposely left incomplete, going only as far as the negative binomial model.

- {lab:9.7}
- (a) Fit the zero-inflated and hurdle models to this data set, considering whether the count component should be Poisson or negative-binomial, and whether the zero model should use all predictors or only a subset. Describe your conclusions from this analysis in a few sentences.
 - (b) Using the methods illustrated in this chapter, create some graphs summarizing the predicted counts and probabilities of zero counts for one of these models.
 - (c) For your chosen model, use some of the diagnostic plots of residuals and other measures shown in Section 9.6 to determine if your model solves any of the problems noted in Example 9.17 and Example 9.18, and whether there are any problems that remain.

Exercise 9.7 In Example 9.19 we used a simple analysis of $\log(y + 1)$ for the multivariate responses in the *NMES1988* data using a classical MLM (Eqn. (9.16)) as a rough approximation of a multivariate Poisson model. The HE plot in Figure ?? was given as a visual summary, but did not show the data. Examine why the MLM is not appropriate statistically for these data, as follows:

- (a) Calculate residuals for the model `nmes.mlm` using

```
> resids <- residuals(nmes.mlm, type = "deviance")
Error in residuals(nmes.mlm, type = "deviance"): error in evaluating the
argument 'object' in selecting a method for function 'residuals': Error:
object 'nmes.mlm' not found
```

- (b) Make univariate density plots of these residuals to show their univariate distributions. These should be approximately normal under the MLM. What do you conclude?
- (c) Make some bivariate plots of these residuals. Under the MLM, each should be bivariate normal with elliptical contours and linear regressions. Add 2D density contours (`kde2d()`, or `geom_density2d()` in `ggplot2`) and some smoothed curve. What do you conclude?

```
> # detach(package:ggtern) ## detach any masking packages
> .locals$ch09 <- setdiff(ls(), .globals)
> #.locals$ch09
> remove(list=.locals$ch09[sapply(.locals$ch09, function(n) {!is.function(get(n))}))])
```

References

- Aberdein, J. and Spiegelhalter, D. (2013). Have London's roads become more dangerous for cyclists? *Significance*, 10(6), 46–48.
- Agresti, A. (1984). *Analysis of Ordinal Categorical Data*. New York: Wiley.
- Agresti, A. (1990). *Categorical Data Analysis*. New York: Wiley-Interscience.
- Agresti, A. (1996). *An Introduction to Categorical Data Analysis*. New York: Wiley Interscience.
- Agresti, A. (2002). *Categorical Data Analysis*. Wiley Series in Probability and Statistics. New York: Wiley-Interscience [John Wiley & Sons], 2nd edn.
- Agresti, A. (2013). *Categorical Data Analysis*. Wiley Series in Probability and Statistics. New York: Wiley-Interscience [John Wiley & Sons], 3rd edn.
- Agresti, A. and Winner, L. (1997). Evaluating agreement and disagreement among movie reviewers. *CHANCE*, 10(2), 10–14.
- Aitchison, J. (1986). *The Statistical Analysis of Compositional Data*. London: Chapman and Hall.
- Akaike, H. (1973). Information theory and an extension of the maximum likelihood principal. In B. N. Petrov and F. Czaki, eds., *Proceedings of the 2nd International Symposium on Information*. Budapest: Akademiai Kiado.
- Andersen, E. B. (1991). *Statistical Analysis of Categorical Data*. Berlin: Springer-Verlag, 2nd edn.
- Anderson, E. (1935). The irises of the Gaspé peninsula. *Bulletin of the American Iris Society*, 35, 2–5.
- Andrews, D. F. and Herzberg, A. M. (1985). *Data: A Collection of Problems from Many Fields for the Student and Research Worker*. New York, NY: Springer-Verlag.
- Antonio, A. L. M. and Crespi, C. M. (2010). Predictors of interobserver agreement in breast imaging using the breast imaging reporting and data system. *Breast Cancer Research and Treatment*, 120(3), 539–546.
- Arbuthnot, J. (1710). An argument for devine providence, taken from the constant regularity observ'd in the births of both sexes. *Philosophical Transactions*, 27, 186–190. Published in 1711.
- Ashford, J. R. and Sowden, R. D. (1970). Multivariate probit analysis. *Biometrics*, 26, 535–546.
- Atkinson, A. C. (1981). Two graphical displays for outlying and influential observations in regression. *Biometrika*, 68, 13–20.
- Atkinson, A. C. (1987). *Plots, Transformations and Regression: An Introduction to Graphical Methods of Diagnostic Regression Analysis*. New York: Oxford University Press.

- Bangdiwala, S. I. (1985). A graphical test for observer agreement. In *Proceeding of the International Statistics Institute*, vol. 1, (pp. 307–308). Amsterdam: ISI.
- Bangdiwala, S. I. (1987). Using SAS software graphical procedures for the observer agreement chart. *Proceedings of the SAS User's Group International Conference*, 12, 1083–1088.
- Bartlett, M. S. (1935). Contingency table interactions. *Journal of the Royal Statistical Society, Supplement*, 2, 248–252.
- Becker, M. P. and Clogg, C. C. (1989). Analysis of sets of two-way contingency tables using association models. *Journal of the American Statistical Association*, 84(405), 142–151.
- Benzécri, J.-P. (1977). Sur l'analyse des tableaux binaires associés à une correspondance multiple. *Cahiers de l'Analyse des Données*, 2, 55–71.
- Bertin, J. (1981). *Graphics and Graphic Information-processing*. New York: de Gruyter. (trans. W. Berg and P. Scott).
- Bertin, J. (1983). *Semiology of Graphics*. Madison, WI: University of Wisconsin Press. (trans. W. Berg).
- Bickel, P. J., Hammel, J. W., and O'Connell, J. W. (1975). Sex bias in graduate admissions: Data from Berkeley. *Science*, 187, 398–403.
- Birch, M. W. (1963a). An algorithm for the logarithmic series distributions. *Biometrics*, 19, 651–652.
- Birch, M. W. (1963b). Maximum likelihood in three-way contingency tables. *Journal of the Royal Statistical Society, Series B*, 25, 220–233.
- Bishop, Y. M. M., Fienberg, S. E., and Holland, P. W. (1975). *Discrete Multivariate Analysis: Theory and Practice*. Cambridge, MA: MIT Press.
- Böhning, D. (1983). Maximum likelihood estimation of the logarithmic series distribution. *Statistische Hefte (Statistical Papers)*, 24(1), 121–140.
- Box, G. E. P. (1979). Some problems of statistics and everyday life. *Journal of the American Statistical Association*, 74(365), 1–4.
- Box, G. E. P. and Cox, D. R. (1964). An analysis of transformations (with discussion). *Journal of the Royal Statistical Society, Series B*, 26, 211–252.
- Box, G. E. P. and Draper, N. R. (1987). *Empirical Model Building and Response Surfaces*. New York, NY: John Wiley & Sons.
- Bradu, D. and Gabriel, R. K. (1978). The biplot as a diagnostic tool for models of two-way tables. *Technometrics*, 20, 47–68.
- Brockmann, H. J. (1996). Satellite male groups in horseshoe crabs, *Limulus polyphemus*. *Ethology*, 102(1), 1–21.
- Brown, P. J., Stone, J., and Ord-Smith, C. (1983). Toxaemic signs during pregnancy. *Journal of the Royal Statistical Society, Series C (Applied Statistics)*, 32, 69–72.
- Brunswick, A. F. (1971). Adolescent health, sex, and fertility. *American Journal of Public Health*, 61(4), 711–729.

- Burt, C. (1950). The factorial analysis of qualitative data. *British Journal of Statistical Psychology*, 3, 166–185.
- Cameron, A. and Trivedi, P. (1990). Regression-based tests for overdispersion in the poisson model. *Journal of Econometrics*, 46, 347–364.
- Cameron, A. C. and Trivedi, P. K. (1998). *Regression analysis of count data*. Econometric society monographs. Cambridge (U.K.), New York: Cambridge University Press.
- Cameron, A. C. and Trivedi, P. K. (2013). *Regression analysis of count data*. Econometric society monographs. Cambridge (U.K.), New York: Cambridge University Press, 2nd edn.
- Carlyle, T. (1840). *Chartism*. London: J. Fraser.
- Caussinus, H. (1966). Contribution à l'analyse statistique des tableaux de corrélation. *Annales de la Faculté des Sciences de l'Université de Toulouse*, 39 (année 1965), 77–183.
- Chambers, J. M., Cleveland, W. S., Kleiner, B., and Tukey, P. A. (1983). *Graphical Methods for Data Analysis*. Belmont, CA: Wadsworth.
- Cicchetti, D. V. and Allison, T. (1971). A new procedure for assessing reliability of scoring EEG sleep recordings. *American Journal of EEG Technology*, 11, 101–109.
- Cleveland, W. S. (1993a). A model for studying display methods of statistical graphics. *Journal of Computational and Graphical Statistics*, 2, 323–343.
- Cleveland, W. S. (1993b). *Visualizing Data*. Summit, NJ: Hobart Press.
- Cleveland, W. S., McGill, M. E., and McGill, R. (1988). The shape parameter of a two-variable graph. *Journal of the American Statistical Association*, 83, 289–300.
- Cleveland, W. S. and McGill, R. (1984). Graphical perception: Theory, experimentation and application to the development of graphical methods. *Journal of the American Statistical Association*, 79, 531–554.
- Cleveland, W. S. and McGill, R. (1985). Graphical perception and graphical methods for analyzing scientific data. *Science*, 229, 828–833.
- Cohen, A. (1980). On the graphical display of the significant components in a two-way contingency table. *Communications in Statistics— Theory and Methods*, A9, 1025–1041.
- Cohen, J. (1960). A coefficient of agreement for nominal scales. *Educational and Psychological Measurement*, 20, 37–46.
- Cohen, J. (1968). Weighted kappa: Nominal scale agreement with provision for scaled disagreement or partial credit. *Psychological Bulletin*, 70, 213–220.
- Cook, R. D. and Weisberg, S. (1999). *Applied Regression Including Computing and Graphics*. New York: Wiley.
- Cragg, J. G. (1971). Some statistical models for limited dependent variables with application to the demand for durable goods. *Econometrica*, 39, 829–844.
- Dalal, S., Fowlkes, E. B., and Hoadley, B. (1989). Risk analysis of the space shuttle: Pre-Challenger prediction of failure. *Journal of the American Statistical Association*, 84(408), 945–957.
- de la Cruz Rot, M. (2005). Improving the presentation of results of logistic regression with r. *Bulletin of the Ecological Society of America*, 86, 41–48.

- Deb, P. and Trivedi, P. K. (1997). Demand for medical care by the elderly: A finite mixture approach. *Journal of Applied Econometrics*, 12, 313–336.
- Edwards, A. W. F. (1958). An analysis of geissler's data on the human sex ratio. *Annals of Human Genetics*, 23(1), 6–15.
- Emerson, J. W., Green, W. A., Schloerke, B., Crowley, J., Cook, D., Hofmann, H., and Wickham, H. (2013). The generalized pairs plot. *Journal of Computational and Graphical Statistics*, 22(1), 79–91.
- Evers, M. and Namboordiri., N. K. (1977). A Monte Carlo assessment of the stability of log-linear estimates in small samples. In *Proceedings of the Social Statistics Section*. Alexandria, VA: American Statistical Association.
- Feynman, R. P. (1988). *What Do You Care What Other People Think? Further Adventures of a Curious Character*. New York: W. W. Norton.
- Fienberg, S. E. (1975). Perspective Canada as a social report. *Social Indicators Research*, 2, 153–174.
- Fienberg, S. E. (1980). *The Analysis of Cross-Classified Categorical Data*. Cambridge, MA: MIT Press, 2nd edn.
- Fienberg, S. E. and Rinaldo, A. (2007). Three centuries of categorical data analysis: Log-linear models and maximum likelihood estimation. *Journal of Statistical Planning and Inference*, 137(11), 3430–3445.
- Firth, D. (2003). Overcoming the reference category problem in the presentation of statistical models. *Sociological Methodology*, 33, 1–18.
- Firth, D. and Menezes, R. X. d. (2004). Quasi-variances. *Biometrika*, 91, 65–80.
- Fisher, R. A. (1925). *Statistical Methods for Research Workers*. London: Oliver & Boyd.
- Fisher, R. A. (1936a). Has Mendel's work been rediscovered? *Annals of Science*, 1, 115–137.
- Fisher, R. A. (1936b). The use of multiple measurements in taxonomic problems. *Annals of Eugenics*, 8, 379–388.
- Fisher, R. A. (1940). The precision of discriminant functions. *Annals of Eugenics*, 10, 422–429.
- Fisher, R. A., Corbet, A. S., and Williams, C. B. (1943). The relation between the number of species and the number of individuals. *Journal of Animal Ecology*, 12, 42.
- Fleiss, J. L. (1973). *Statistical Methods for Rates and Proportions*. New York: John Wiley and Sons.
- Fleiss, J. L. and Cohen, J. (1972). The equivalence of weighted kappa and the intraclass correlation coefficient as measures of reliability. *Educational and Psychological Measurement*, 33, 613–619.
- Fleiss, J. L., Cohen, J., and Everitt, B. S. (1969). Large sample standard errors of kappa and weighted kappa. *Psychological Bulletin*, 72, 332–327.
- Fowlkes, E. B. (1987). Some diagnostics for binary logistic regression via smoothing. *Biometrika*, 74(3), 503–5152.
- Fox, J. (1987). Effect displays for generalized linear models. In C. C. Clogg, ed., *Sociological Methodology*, 1987, (pp. 347–361). San Francisco: Jossey-Bass.

- Fox, J. (2003). Effect displays in R for generalized linear models. *Journal of Statistical Software*, 8(15), 1–27.
- Fox, J. (2008). *Applied Regression Analysis and Generalized Linear Models*. Thousand Oaks, CA: Sage, 2nd edn.
- Fox, J. and Andersen, R. (2006). Effect displays for multinomial and proportional-odds logit models. *Sociological Methodology*, 36, 225–255.
- Fox, J., Friendly, M., and Monette, G. (2009). Visualizing hypothesis tests in multivariate linear models: The *heplots* package for R. *Computational Statistics*, 24(2), 233–246. (Published online: 15 May 2008).
- Fox, J. and Weisberg, S. (2011). *An R Companion to Applied Regression*. Thousand Oaks CA: Sage, 2nd edn.
- Friendly, M. (1991). *SAS System for Statistical Graphics*. Cary, NC: SAS Institute, 1st edn.
- Friendly, M. (1992). Mosaic displays for loglinear models. In ASA, *Proceedings of the Statistical Graphics Section*, (pp. 61–68). Alexandria, VA.
- Friendly, M. (1994a). A fourfold display for 2 by 2 by K tables. Tech. Rep. 217, York University, Psychology Dept.
- Friendly, M. (1994b). Mosaic displays for multi-way contingency tables. *Journal of the American Statistical Association*, 89, 190–200.
- Friendly, M. (1994c). SAS/IML graphics for fourfold displays. *Observations*, 3(4), 47–56.
- Friendly, M. (1995). Conceptual and visual models for categorical data. *The American Statistician*, 49, 153–160.
- Friendly, M. (1997). Conceptual models for visualizing contingency table data. In M. Greenacre and J. Blasius, eds., *Visualization of Categorical Data*, chap. 2, (pp. 17–35). San Diego, CA: Academic Press.
- Friendly, M. (1999a). Extending mosaic displays: Marginal, conditional, and partial views of categorical data. *Journal of Computational and Graphical Statistics*, 8(3), 373–395.
- Friendly, M. (1999b). Extending mosaic displays: Marginal, conditional, and partial views of categorical data. *Journal of Computational and Graphical Statistics*, 8(3), 373–395.
- Friendly, M. (2000). *Visualizing Categorical Data*. Cary, NC: SAS Institute.
- Friendly, M. (2002). Corrrgrams: Exploratory displays for correlation matrices. *The American Statistician*, 56(4), 316–324.
- Friendly, M. (2003). Visions of the past, present and future of statistical graphics: An ideo-graphic view. American Psychological Association. Toronto, ON, URL: <http://datavis.ca/papers/apa-2x2.pdf>.
- Friendly, M. (2007). HE plots for multivariate general linear models. *Journal of Computational and Graphical Statistics*, 16(2), 421–444.
- Friendly, M. (2013). Comment on the generalized pairs plot. *Journal of Computational and Graphical Statistics*, 22(1), 290–291.

- Friendly, M. and Kwan, E. (2003). Effect ordering for data displays. *Computational Statistics and Data Analysis*, 43(4), 509–539.
- Friendly, M. and Kwan, E. (2011). Comment (graph people versus table people). *Journal of Computational and Graphical Statistics*, 20(1), 18–27.
- Friendly, M., Monette, G., and Fox, J. (2013). Elliptical insights: Understanding statistical methods through elliptical geometry. *Statistical Science*, 28(1), 1–39.
- Gabriel, K. R. (1971). The biplot graphic display of matrices with application to principal components analysis. *Biometrics*, 58(3), 453–467.
- Gabriel, K. R. (1980). Biplot. In N. L. Johnson and S. Kotz, eds., *Encyclopedia of Statistical Sciences*, vol. 1, (pp. 263–271). New York: John Wiley and Sons.
- Gabriel, K. R. (1981). Biplot display of multivariate matrices for inspection of data and diagnosis. In V. Barnett, ed., *Interpreting Multivariate Data*, chap. 8, (pp. 147–173). London: John Wiley and Sons.
- Gabriel, K. R., Galindo, M. P., and Vincente-Villardón, J. L. (1997). Use of biplots to diagnose independence models in three-way contingency tables. In M. Greenacre and J. Blasius, eds., *Visualization of Categorical Data*, chap. 27, (pp. 391–404). San Diego, CA: Academic Press.
- Gabriel, K. R. and Odoroff, C. L. (1990). Biplots in biomedical research. *Statistics in Medicine*, 9, 469–485.
- Gart, J. J. and Zweifel, J. R. (1967). On the bias of various estimators of the logit and its variance with applications to quantal bioassay. *Biometrika*, 54, 181–187.
- Geissler, A. (1889). Beitrage zur frage des geschlechts verhaltnisses der geborenen. *Z. K. Sachsischen Statistischen Bureaus*, 35(1), n.p.
- Gifi, A. (1981). *Nonlinear Multivariate Analysis*. The Netherlands: Department of Data Theory, University of Leiden.
- Glass, D. V. (1954). *Social Mobility in Britain*. Glencoe, IL: The Free Press.
- Goodman, L. A. (1970). The multivariate analysis of qualitative data: Interactions among multiple classifications. *Journal of the American Statistical Association*, 65, 226–256.
- Goodman, L. A. (1971). The analysis of multidimensional contingency tables: Stepwise procedures and direct estimates for building models for multiple classifications. *Technometrics*, 13, 33–61.
- Goodman, L. A. (1972). Some multiplicative models for the analysis of cross classified data. In *Proceedings of the sixth Berkeley Symposium on Mathematical Statistics and Probability*, (pp. 649–696). Berkeley, CA: University of California.
- Goodman, L. A. (1973). The analysis of multidimensional contingency tables when some variables are posterior to others: A modified path analysis approach. *Biometrika*, 60, 179–192.
- Goodman, L. A. (1979). Simple models for the analysis of association in cross-classifications having ordered categories. *Journal of the American Statistical Association*, 74, 537–552.
- Goodman, L. A. (1981). Association models and canonical correlation in the analysis of cross-classifications having ordered categories. *Journal of the American Statistical Association*, 76(374), 320–334.

- Goodman, L. A. (1985). The analysis of cross-classified data having ordered and/or unordered categories: Association models, correlation models, and asymmetry models for contingency tables with or without missing entries. *Annals of Statistics*, 13(1), 10–69.
- Goodman, L. A. (1986). Some useful extensions of the usual correspondence analysis approach and the usual log-linear models approach in the analysis of contingency tables. *International Statistical Review*, 54(3), 243–309. With a discussion and reply by the author.
- Gower, J., Lubbe, S., and Roux, N. (2011). *Understanding Biplots*. Wiley.
- Gower, J. C. and Hand, D. J. (1996). *Biplots*. London: Chapman & Hall.
- Grayson, D. K. (1990). Donner party deaths: A demographic assessment. *Journal of Anthropological Research*, 46(3), 223–242.
- Greenacre, M. (1984). *Theory and Applications of Correspondence Analysis*. London: Academic Press.
- Greenacre, M. (1988). Correspondence analysis of multivariate categorical data by weighted least squares. *Biometrika*, 75, 457–467.
- Greenacre, M. (1989). The Carroll-Green-Schaffer scaling in correspondence analysis: A theoretical and empirical appraisal. *Journal of Marketing Research*, 26, 358–365.
- Greenacre, M. (1990). Some limitations of multiple correspondence analysis. *Computational Statistics Quarterly*, 3, 249–256.
- Greenacre, M. (1997). Diagnostics for joint displays in correspondence analysis. In J. Blasius and M. Greenacre, eds., *Visualization of Categorical Data*, (pp. 221–238). Academic Press.
- Greenacre, M. (2013). Contribution biplots. *Journal of Computational and Graphical Statistics*, 22(1), 107–122.
- Greenacre, M. and Hastie, T. (1987). The geometric interpretation of correspondence analysis. *Journal of the American Statistical Association*, 82, 437–447.
- Greenacre, M. J. (2007). *Correspondence analysis in practice*. Boca Raton: Chapman & Hall/CRC.
- Greenwood, M. and Yule, G. U. (1920). An inquiry into the nature of frequency distributions of multiple happenings, with particular reference to the occurrence of multiple attacks of disease or repeated accidents. *Journal of the Royal Statistical Society, Series A*, 83, 255–279.
- Haberman, S. J. (1972). Statistical algorithms: Algorithm AS 51: Log-linear fit for contingency tables. *Applied Statistics*, 21(2), 218–225.
- Haberman, S. J. (1973). The analysis of residuals in cross-classified tables. *Biometrics*, 29, 205–220.
- Haberman, S. J. (1974). *The Analysis of Frequency Data*. Chicago: University of Chicago Press.
- Haberman, S. J. (1979). *The Analysis of Qualitative Data: New Developments*, vol. II. New York: Academic Press.
- Haldane, J. B. S. (1955). The estimation and significance of the logarithm of a ratio of frequencies. *Annals of Human Genetics*, 20, 309–311.
- Harrell, Jr, F. E. (2001). *Regression Modeling Strategies: With Applications to Linear Models, Logistic Regression, and Survival Analysis*. Graduate Texts in Mathematics. New York: Springer.

- Hartigan, J. A. and Kleiner, B. (1981). Mosaics for contingency tables. In W. F. Eddy, ed., *Computer Science and Statistics: Proceedings of the 13th Symposium on the Interface*, (pp. 268–273). New York, NY: Springer-Verlag.
- Hartigan, J. A. and Kleiner, B. (1984). A mosaic of television ratings. *The American Statistician*, 38, 32–35.
- Hastie, T. J. and Tibshirani, R. J. (1990). *Generalized Additive Models*. London: Chapman & Hall.
- Hauser, R. M. (1979). Some exploratory methods for modeling mobility tables and other cross-classified data. In K. F. Schuessler, ed., *Sociological Methodology 1980*, (pp. 413–458). San Francisco: Jossey-Bass.
- Hemmingsen, W., Jansen, P. A., and Mackenzie, K. (2005). Crabs, leeches and trypanosomes: an unholy trinity? *Marine Pollution Bulletin*, 50(3), 336–339.
- Heuer, J. (1979). *Selbstmord Bei Kinder Und Jugendlichen*. Stuttgart: Ernst Klett Verlag. [Suicide by children and youth.]
- Hilbe, J. (2011). *Negative Binomial Regression*. Cambridge University Press, 2nd edn.
- Hilbe, J. M. (2014). *Modeling Count Data*. New York, NY: Cambridge University Press.
- Hoaglin, D. C. (1980). A poissonness plot. *The American Statistician*, 34, 146–149.
- Hoaglin, D. C. and Tukey, J. W. (1985). Checking the shape of discrete distributions. In D. C. Hoaglin, F. Mosteller, and J. W. Tukey, eds., *Exploring Data Tables, Trends and Shapes*, chap. 9. New York: John Wiley and Sons.
- Hofmann, H. and Theus, M. (2005). Interactive graphics for visualizing conditional distributions. Unpublished manuscript.
- Holm, S. (1979). A simple sequentially rejective multiple test procedure. *Scandinavian Journal of Statistics*, 6(2), 65–70.
- Hosmer, Jr, D. W., Lemeshow, S., and Sturdivant, R. X. (2013). *Applied Logistic Regression*. New York: John Wiley and Sons, 3rd edn.
- Hout, M., Duncan, O. D., and Sobel, M. E. (1987). Association and heterogeneity: Structural models of similarities and differences. *Sociological Methodology*, 17, 145–184.
- Hummel, J. (1996). Linked bar charts: Analyzing categorical data graphically. *Computational Statistics*, 11, 23–33.
- Hurley, C. B. (2004). Clustering visualizations of multidimensional data. *Journal of Computational and Graphical Statistics*, 13, 788–806.
- Immer, F. R., Hayes, H., and Powers, L. R. (1934). Statistical determination of barley varietal adaptation. *Journal of the American Society of Agronomy*, 26, 403–419.
- Jansen, J. (1990). On the statistical analysis of ordinal data when extravariation is present. *Journal of the Royal Statistical Society. Series C (Applied Statistics)*, 39(1), 75–84.
- Jinkinson, R. A. and Slater, M. (1981). Critical discussion of a graphical method for identifying discrete distributions. *The Statistician*, 30, 239–248.
- Johnson, K. (1996). *Unfortunate Emigrants: Narratives of the Donner Party*. Logan, UT: Utah State University Press.

- Johnson, N. L., Kotz, S., and Kemp, A. W. (1992). *Univariate Discrete Distributions*. New York, NY: John Wiley and Sons, 2nd edn.
- Kemp, A. W. and Kemp, C. D. (1991). Weldon's dice data revisited. *The American Statistician*, 45, 216–222.
- Kendall, M. G. and Stuart, A. (1961). *The Advanced Theory of Statistics*, vol. 2. London: Griffin.
- Kendall, M. G. and Stuart, A. (1963). *The Advanced Theory of Statistics*, vol. 1. London: Griffin.
- King, G. (1989). A seemingly unrelated Poisson regression model. *Sociological Methods and Research*, 17(3), 235–255.
- Kleiber, C. and Zeileis, A. (2014). Visualizing count data regressions using rootograms. Working papers, Faculty of Economics and Statistics, University of Innsbruck.
- Koch, G. and Edwards, S. (1988). Clinical efficiency trials with categorical data. In K. E. Peace, ed., *Biopharmaceutical Statistics for Drug Development*, (pp. 403–451). New York: Marcel Dekker.
- Kosambi, D. D. (1949). Characteristic properties of series distributions. *Proceedings of the National Institute of Science of India*, 15, 109–113.
- Kosslyn, S. M. (1985). Graphics and human information processing: A review of five books. *Journal of the American Statistical Association*, 80, 499–512.
- Kosslyn, S. M. (1989). Understanding charts and graphs. *Applied Cognitive Psychology*, 3, 185–225.
- Kundel, H. L. and Polansky, M. (2003). Measurement of observer agreement. *Radiology*, 228(2), 303–308.
- Labby, Z. (2009). Weldon's dice, automated. *Chance*, 22(4), 6–13.
- Lambert, D. (1992). Zero-inflated poisson regression, with an application to defects in manufacturing. *Technometrics*, 34, 1–14.
- Landis, J. R. and Koch, G. G. (1977). The measurement of observer agreement for categorical data. *Biometrics*, 33, 159–174.
- Landis, R. J., Heyman, E. R., and Koch, G. G. (1978). Average partial association in three-way contingency tables: A review and discussion of alternative tests,. *International Statistical Review*, 46, 237–254.
- Landwehr, J. M., Pregibon, D., and Shoemaker, A. C. (1984). Graphical methods for assessing logistic regression models. *Journal of the American Statistical Association*, 79, 61–71.
- Lang, J. B. and Agresti, A. (1994). Simultaneously modeling joint and marginal distributions of multivariate categorical responses. *Journal of the American Statistical Association*, 89(426), 625–632.
- Larsen, W. A. and McCleary, S. J. (1972). The use of partial residual plots in regression analysis. *Technometrics*, 14, 781–790.
- Lavine, M. (1991). Problems in extrapolation illustrated with space shuttle O-ring data. *Journal of the American Statistical Association*, 86, 912–922.
- Lawrance, A. J. (1995). Deletion influence and masking in regression. *Journal of the Royal Statistical Society. Series B (Methodological)*, 57(1), 181–189.

- Lebart, L., Morineau, A., and Warwick, K. M. (1984). *Multivariate Descriptive Statistical Analysis: Correspondence Analysis and Related Techniques for Large Matrices*. New York: John Wiley and Sons.
- Lee, A. J. (1997). Modelling scores in the Premier League: Is Manchester United really the best? *Chance*, 10(1), 15–19.
- Leifeld, P. (2013). texreg: Conversion of statistical model output in r to latex and html tables. *Journal of Statistical Software*, 55(8), 1–24.
- Lemeshow, S., Avrunin, D., and Pastides, J. S. (1988). Predicting the outcome of intensive care unit patients. *Journal of the American Statistical Association*, 83, 348–356.
- Lewandowsky, S. and Spence, I. (1989). The perception of statistical graphs. *Sociological Methods & Research*, 18, 200–242.
- Lindsey, J. K. (1995). *Modelling Frequency and Count Data*. Oxford, UK: Oxford University Press.
- Lindsey, J. K. and Altham, P. M. E. (1998). Analysis of the human sex ratio by using overdispersion models. *Journal of the Royal Statistical Society: Series C (Applied Statistics)*, 47(1), 149–157.
- Lindsey, J. K. and Mersch, G. (1992). Fitting and comparing probability distributions with log linear models. *Computational Statistics and Data Analysis*, 13, 373–384.
- Long, J. S. (1990). The origins of sex differences in science. *Social Forces*, 68(4), 1297–1316.
- Long, J. S. (1997). *Regression Models for Categorical and Limited Dependent Variables*. Thousand Oaks, CA: Sage Publications.
- McCullagh, P. (1980). Regression models for ordinal data. *Journal of the Royal Statistical Society, Series B*, 42, 109–142.
- McCullagh, P. and Nelder, J. A. (1989). *Generalized Linear Models*. London: Chapman and Hall.
- McCulloch, C. E. and Neuhaus, J. M. (2005). Generalized linear mixed models. In *Encyclopedia of Biostatistics*. John Wiley & Sons, Ltd.
- Mersey, L. (1912). Report on the loss of the “Titanic” (S. S.). Parliamentary command paper 6352.
- Meyer, D., Zeileis, A., and Hornik, K. (2006). The strucplot framework: Visualizing multi-way contingency tables with vcd. *Journal of Statistical Software*, 17(3), 1–48.
- Milan, L. and Whittaker, J. (1995). Application of the parametric bootstrap to models that incorporate a singular value decomposition. *Journal of the Royal Statistical Society. Series C (Applied Statistics)*, 44(1), 31–49.
- Mosteller, F. and Wallace, D. L. (1963). Inference in an authorship problem. *Journal of the American Statistical Association*, 58(302), 275–309.
- Mosteller, F. and Wallace, D. L. (1984). *Applied Bayesian and Classical Inference: The Case of the Federalist Papers*. New York, NY: Springer-Verlag.
- Mullahy, J. (1986). Specification and testing of some modified count data models. *Journal of Econometrics*, 33, 341–365.
- Nelder, J. A. and Wedderburn, R. W. M. (1972). Generalized linear models. *Journal of the Royal Statistical Society, Series A*, 135, 370–384.

- Noack, A. (1950). A class of random variables with discrete distributions. *Annals of Mathematical Statistics*, 21, 127–132.
- Ord, J. K. (1967). Graphical methods for a class of discrete distributions. *Journal of the Royal Statistical Society, Series A*, 130, 232–238.
- Pareto, V. (1971). *Manuale di economia politica* (“Manual of political economy”). New York: A.M. Kelley. Translated by Ann S. Schwier. Edited by Ann S. Schwier and Alfred N. Page.
- Pearson, K. (1900). On the criterion that a given system of deviations from the probable in the case of a correlated system of variables is such that it can be reasonably supposed to have arisen by random sampling. *Philosophical Magazine*, 50(5th Series), 157–175.
- Peterson, B. and Harrell, F. (1990). Partial proportional odds models for ordinal response variables. *Applied Statistics*, 39, 205–217.
- Powers, D. A. and Xie, Y. (2008). *Statistical Methods for Categorical Data Analysis*. Bingley, UK: Emerald, 2nd edn.
- Pregibon, D. (1981). Logistic regression diagnostics. *Annals of Statistics*, 9, 705–724.
- Raftery, A. E. (1986). Choosing models for cross-classifications. *American Sociological Review*, 51, 146–146.
- Ramsey, F. L., Schafer, D. W., Sifneos, J., and Turlach, B. A. (2012). *Sleuth2: Data sets from Ramsey and Schafer's "Statistical Sleuth (2nd ed)"*. R package version 1.0-7.
- Riedwyl, H. and Schüpbach, M. (1983). Siebdiagramme: Graphische darstellung von kontingenztafeln. Tech. Rep. 12, Institute for Mathematical Statistics, University of Bern, Bern, Switzerland.
- Riedwyl, H. and Schüpbach, M. (1994). Parquet diagram to plot contingency tables. In F. Faulbaum, ed., *Softstat '93: Advances In Statistical Software*, (pp. 293–299). New York: Gustav Fischer.
- Schwartz, G. (1978). Estimating the dimensions of a model. *Annals of Statistics*, 6, 461–464.
- Searle, S. R., Speed, F. M., and Milliken, G. A. (1980). Population marginal means in the linear model: An alternative to least squares means. *The American Statistician*, 34(4), 216–221.
- Shneiderman, B. (1992). Tree visualization with treemaps: A 2-D space-filling approach. *ACM Transactions on Graphics*, 11(1), 92–99.
- Shrout, P. E. and Fleiss, J. L. (1979). Intraclass correlations: Uses in assessing rater reliability. *Psychological Bulletin*, 86, 420–428.
- Simpson, E. H. (1951). The interpretation of interaction in contingency tables. *Journal of the Royal Statistical Society, Series B*, 30, 238–241.
- Skellam, J. G. (1948). A probability distribution derived from the binomial distribution by regarding the probability of success as variable between the sets of trials. *Journal of the Royal Statistical Society. Series B (Methodological)*, 10(2), 257–261.
- Snee, R. D. (1974). Graphical display of two-way contingency tables. *The American Statistician*, 28, 9–12.
- Spence, I. (1990). Visual psychophysics of simple graphical elements. *Journal of Experimental Psychology: Human Perception and Performance*, 16, 683–692.

- Spence, I. and Lewandowsky, S. (1990). Graphical perception. In J. Fox and J. S. Long, eds., *Modern Methods of Data Analysis*, chap. 1, (pp. 13–57). Sage Publications, Inc.
- Srole, L., Langner, T. S., Michael, S. T., Kirkpatrick, P., Opler, M. K., and Rennie, T. A. C. (1978). *Mental Health in the Metropolis: The Midtown Manhattan Study*. New York: NYU Press.
- Stokes, M. E., Davis, C. S., and Koch, G. G. (2000). *Categorical Data Analysis Using the SAS System*. Cary, NC: SAS Institute, 2nd edn.
- Theus, M. and Lauer, S. R. W. (1999). Visualizing loglinear models. *Journal of Computational and Graphical Statistics*, 8(3), 396–412.
- Thornes, B. and Collard, J. (1979). *Who Divorces?* London: Routledge & Kegan.
- Thurstone, L. L. (1927). A law of comparative judgment. *Psychological Review*, 34, 278–286.
- Tufte, E. (2006). *Beautiful Evidence*. Cheshire, CT: Graphics Press.
- Tufte, E. R. (1983). *The Visual Display of Quantitative Information*. Cheshire, CT: Graphics Press.
- Tufte, E. R. (1990). *Envisioning Information*. Cheshire, CT: Graphics Press.
- Tufte, E. R. (1997). *Visual Explanations: Images and Quantities, Evidence and Narrative*. Cheshire, CT: Graphics Press.
- Tukey, J. W. (1962). The future of data analysis. *Annals of Mathematical Statistics*, 33, 1–67 and 81.
- Tukey, J. W. (1977). *Exploratory Data Analysis*. Reading, MA: Addison Wesley.
- Tukey, J. W. (1993). Graphic comparisons of several linked aspects: Alternative and suggested principles. *Journal of Computational and Graphical Statistics*, 2(1), 1–33.
- Upton, G. J. G. (1976). The diagrammatic representation of three-party contests. *Political Studies*, 24, 448–454.
- Upton, G. J. G. (1994). Picturing the 1992 British general election. *Journal of the Royal Statistical Society, Series A*, 157(Part 2), 231–252.
- van der Heijden, P. G. M., de Falguerolles, A., and de Leeuw, J. (1989). A combined approach to contingency table analysis using correspondence analysis and log-linear analysis. *Applied Statistics*, 38(2), 249–292.
- van der Heijden, P. G. M. and de Leeuw, J. (1985). Correspondence analysis used complementary to loglinear analysis. *Psychometrika*, 50, 429–447.
- von Bortkiewicz, L. (1898). *Das Gesetz der Kleinen Zahlen*. Leipzig: Teubner.
- Von Eye, A. and Mun, E. (2006). *Analyzing Rater Agreement: Manifest Variable Methods*. Taylor & Francis.
- Vuong, Q. H. (1989). Likelihood ratio tests for model selection and non-nested hypotheses. *Econometrica*, 57(2), pp. 307–333.
- Wainer, H. (1996). Using trilinear plots for NAEP state data. *Journal of Educational Measurement*, 33(1), 41–55.
- Wang, P. C. (1985). Adding a variable in generalized linear models. *Technometrics*, 27, 273–276.

- Wickham, H. (2009). *ggplot2: Elegant Graphics for Data Analysis*. Springer New York.
- Wickham, H. and Chang, W. (2013). *ggplot2: An implementation of the Grammar of Graphics*. R package version 0.9.3.1.
- Wilkinson, L. (2005). *The Grammar of Graphics*. New York: Springer, 2nd edn.
- Williams, D. A. (1987). Generalized linear model diagnostics using the deviance and single case deletions. *Applied Statistics*, 36, 181–191.
- Wimmer, G. and Altmann, G. (1999). *Thesaurus of univariate discrete probability distributions*. Stamm.
- Wong, R. S.-K. (2001). Multidimensional association models: A multilinear approach. *Sociological Methods and Research*, 30(2), 197–240.
- Wong, R. S.-K. (2010). *Association Models*. Quantitative Applications in the Social Sciences. Los Angeles: SAGE Publications.
- Wood, S. N. (2004). Stable and efficient multiple smoothing parameter estimation for generalized additive models. *Journal of the American Statistical Association*, 99(467), 673–686.
- Wood, S. N. (2006). *Generalized Additive Models: An Introduction with R*. Chapman and Hall/CRC Press.
- Woolf, B. (1995). On estimating the relation between blood group and disease. *Annals of Human Genetics*, 19, 251–253.
- Wright, K. (2013). Revisiting immer's barley data. *The American Statistician*, 67(3), 129–133.
- Yip, K. C. and Yau, K. K. (2005). On modeling claim frequency data in general insurance with extra zeros. *Insurance: Mathematics and Economics*, 36(2), 153–163.
- Zeileis, A., Kleiber, C., and Jackman, S. (2008). Regression models for count data in R. *Journal of Statistical Software*, 27(8).
- Zeileis, A., Meyer, D., and Hornik, K. (2007). Residual-based shadings for visualizing (conditional) independence. *Journal of Computational and Graphical Statistics*, 16(3), 507–525.
- Zelterman, D. (1999). *Models for Discrete Data*. New York: Oxford University Press.

Example Index

- 512 paths to the White House, 20
- Arbuthnot data, 58–59
- Arthritis treatment, 2, 33–35, 49–50, 109–110, 113–114, 117–119, 161–162, 187–188, 239–240, 246–248, 253–254
logistic regression, 239
- Arthritis treatment: conditional plots, 248–250
- Arthritis treatment: full-model plots, 250–251
- Arthritis treatment: Plotting logistic regression with base graphics, 242–243
- Arthritis treatment: Plotting logistic regression with ggplot2, 243
- Barley data, 17–18
- Bartlett data on plum root cuttings, 182, 189
- Berkeley admissions, 2, 108, 118–121, 130–132, 184–186, 329–334, 336–339
- Breathlessness and wheeze in coal miners, 124–126, 374–382
- British social mobility, 16
- Butterfly species in Malaya, 64
- Cod parasites, 422–432
- Collapsing categories, 47–48
- Corporal punishment data, 178–180
- Dayton survey, 45–47
- Death by horse kick, 49, 61–62, 80–81, 83, 89–90, 94
- Death in the ICU, 268–275, 283–287, 292–294
- Demand for medical care, 433–444, 455–464
- Diagnosis of MS patients, 135, 140–141
- Donner Party, 9–10, 255–263, 279–283, 288–289, 291–292
- Donner party, 21
- Employment status data, 175–178
- Families in Saxony, 4, 5, 59–60, 81–82, 95–99
- Federalist papers, 62, 84–85, 90, 96
- General social survey, 35, 48–49
- Hair color and eye color, 6–7, 36, 108, 128, 150–154, 202–205, 219–220
- Hair color, eye color and sex, 164, 167–168
- Hauser's occupational mobility table, 359–366
- Health concerns of teenagers, 341–343
- Interpolation options, 159
- Iris data, 18–19
- Job satisfaction, 37
- Lifeboats on the *Titanic*, 143–145
- London cycling deaths, 62–63, 71–72
- Mammogram ratings, 139–140
- Marital status and pre- and extramarital sex, 171–173, 183–184, 222–223
- Mating of horseshoe crabs, 405–412, 416, 420
- Mental impairment and parents' SES, 109, 114, 205–207, 345–355
- Plotting styles for discrete distributions, 72–74
- Publications of PhD candidates, 99–101, 401–405, 410–415, 447–450, 452–453
- Racial profiling: Arrests for marijuana possession, 12–14, 263–267
- Repeat victimization, 207–209
- Sex is fun, 135
- Shading functions, 159–160
- Simulating zero-inflated data, 418
- Space shuttle disaster, 8–9, 21–23, 244–245
- Suicide rates in Germany, 212–214
- Suicide rates in Germany: biplot, 227–229

- Suicide rates in Germany: mosaic plot, 214–215
Survival on the *Titanic*, 224–225
Toxaemic symptoms in pregnancy, 383–392
TV viewing data, 50–53, 209–210
UK Soccer scores, 70–71, 163, 231–232
Visual acuity, 128–130, 357–359, 368–371
Weldon's dice, 60, 67–68, 82–83
Women in queues, 91
Women's labor force participation, 306–316

Author Index

- Aberdein, J. 62
Agresti, A. 35, 37, 109, 112, 124, 135, 147, 172, 173, 327, 340, 344, 372, 393, 394, 398, 405, 408, 411
Aitchison, J. 141
Akaike, H. 327, 400
Allison, T. 137
Altham, P. M. E. 97
Altmann, G. 65
Andersen, E. B. 174, 175, 178
Andersen, R. 438
Anderson, E. 18
Andrews, D. F. 61
Antonio, A. L. M. 139
Arbuthnott, J. 58
Ashford, J. R. 124, 374
Atkinson, A. C. 451
Avrunin, D. 268

Bangdiwala, S. I. 137
Bartlett, M. S. 182
Becker, M. P. 355
Benzécri, J.-P. 224
Bertin, J. 11, 133, 198
Bickel, P. J. 108, 123
Birch, M. W. 79, 340
Bishop, Y. M. M. 16, 128, 340
Bliss, C. I. 296
Böhning, D. 79
Bollen, K. A. 297
Box, G. E. P. 235, 445, 465
Bradu, D. 225, 230
Brockmann, H. J. 405
Brown, P. J. 383
Brunswick, A. F. 341
Burt, C. 221

Cameron, A. 414
Cameron, A. C. 414, 415, 436, 455
Carlyle, T. 107
Caussinus, H. 357
Chambers, J. M. 11
Chang, W. 11

Cicchetti, D. V. 137
Cleveland, W. S. 11, 17, 24, 175
Clogg, C. C. 355
Cohen, A. 133
Cohen, J. 136, 137
Collard, J. 171
Cook, D. 186
Cook, R. D. 289
Corbet, A. S. 64, 78
Cox, D. R. 465
Cragg, J. G. 419
Crespi, C. M. 139
Crowley, J. 186

Dalal, S. 22, 244
Davis, C. S. 112
de Falguerolles, A. 199
de la Cruz Rot, M. 239
de Leeuw, J. 199, 210–212
Deb, P. 433, 436
Draper, N. R. 235
Duncan, O. D. 135

Edwards, A. W. F. 59, 96
Edwards, S. 33, 109
Emerson, J. W. 186, 210
Everitt, B. S. 136
Evers, M. 340

Feynman, R. P. 23
Fienberg, S. E. 16, 119, 128, 171, 207, 325, 340, 341
Finney, D. J. 296
Firth, D. 351
Fisher, R. A. 18, 61, 64, 78, 81, 197, 233
Fleiss, J. L. 136, 137
Fowlkes, E. B. 22, 244, 288
Fox, J. 27, 252, 260, 297, 306, 307, 398, 438, 446, 455
Friendly, M. 11, 15, 16, 18, 119, 133, 150, 154, 160, 169, 172, 175, 186, 212, 222, 455, 456
Gabriel, K. R. 225, 227, 230
Gabriel, R. K. 225, 230

- Galindo, M. P. 230
 Gart, J. J. 113
 Geissler, A. 59, 103
 Gifi, A. 199
 Glass, D. V. 16
 Goodman, L. A. 363
 Gower, J. 225, 228, 229
 Gower, J. C. 221
 Grayson, D. K. 256, 263, 318
 Green, W. A. 186
 Greenacre, M. 199, 200, 202, 220–222, 224, 227
 Greenacre, M. J. 199, 200, 220, 224
 Greenwood, M. 75
 Haberman, S. J. 109, 325, 328, 340
 Haldane, J. B. S. 113
 Hammel, J. W. 108, 123
 Hand, D. J. 221
 Harrell, F. 299
 Harrell, F. E., Jr 270, 301
 Hartigan, J. A. 50, 150, 209
 Hastie, T. 199, 202
 Hastie, T. J. 443
 Hauser, R. M. 359, 362, 365
 Hayes, H. 17
 Hemmingsen, W. 422, 424
 Herzberg, A. M. 61
 Heuer, J. 212
 Heyman, E. R. 114
 Hilbe, J. 74
 Hilbe, J. M. 411
 Hoadley, B. 22, 244
 Hoaglin, D. C. 91–93, 95
 Hofmann, H. 186, 420
 Holland, P. W. 16, 128, 340
 Holm, S. 123
 Hornik, K. 155, 159, 160, 180
 Hosmer, D. W., Jr 268
 Hout, M. 135
 Hummel, J. 420
 Hurley, C. B. 19
 Immer, F. R. 17
 Jackman, S. 433, 436
 Jansen, J. 197
 Jansen, P. A. 422, 424
 Jinkinson, R. A. 91, 92
 Johnson, K. 263
 Johnson, N. L. 65
 Kemp, A. W. 60, 65
 Kemp, C. D. 60
 Kendall, M. G. 60, 128
 King, G. 455
 Kirkpatrick, P. 109
 Kleiber, C. 415, 433, 436
 Kleiner, B. 11, 50, 150, 209
 Koch, G. 33, 109
 Koch, G. G. 112, 114, 135
 Kosambi, D. D. 79
 Kosslyn, S. M. 11
 Kotz, S. 65
 Kundel, H. L. 139
 Kwan, E. 11, 16, 18, 154
 Labby, Z. 60
 Lambert, D. 417
 Landis, J. R. 135
 Landis, R. J. 114
 Landwehr, J. M. 288
 Lang, J. B. 372
 Langner, T. S. 109
 Larsen, W. A. 288
 Lauer, S. R. W. 190
 Lavine, M. 22, 244
 Lawrance, A. J. 290
 Lazarsfeld, P. F. 297
 Lebart, L. 199
 Lee, A. J. 70
 Leifeld, P. 43
 Lemeshow, S. 268
 Lewandowsky, S. 11
 Lindsey, J. K. 4, 59, 96, 97, 339
 Long, J. S. 99, 417, 453
 Lubbe, S. 225, 228, 229
 Mackenzie, K. 422, 424
 McRae, M. 467
 McCleary, S. J. 288
 McCullagh, P. 296, 372, 373
 McCulloch, C. E. 396
 McGill, M. E. 24
 McGill, R. 11, 24
 Menezes, R. X. d. 351
 Mersch, G. 96, 97
 Mersey, L. 143
 Meyer, D. 155, 159, 160, 180
 Michael, S. T. 109
 Milan, L. 354
 Milliken, G. A. 252
 Monette, G. 455
 Morineau, A. 199
 Mosteller, F. 62, 84, 90, 103
 Mullahy, J. 419
 Mun, E. 136
 Namboordiri, N. K. 340
 Nelder, J. A. 372, 373, 395, 396
 Neuhaus, J. M. 396
 Noack, A. 79

- O'Connell, J. W. 108, 123
Odoroff, C. L. 227
Opler, M. K. 109
Ord, J. K. 88
Ord-Smith, C. 383

Pareto, V. 23
Pastides, J. S. 268
Pearson, K. 60
Peterson, B. 299
Polansky, M. 139
Poole, J. H. 466
Powers, D. A. 359, 363
Powers, L. R. 17
Pregibon, D. 276–278, 288

Raftery, A. E. 328
Ramsey, F. L. 466
Rennie, T. A. C. 109
Riedwyl, H. 127
Rinaldo, A. 340
Roux, N. 225, 228, 229

Schafer, D. W. 466
Schloerke, B. 186
Schüpbach, M. 127
Schwartz, G. 328, 400
Searle, S. R. 252
Shneiderman, B. 20
Shoemaker, A. C. 288
Shrout, P. E. 136
Sifneos, J. 466
Simpson, E. H. 123
Skellam, J. G. 96
Slater, M. 91, 92
Snee, R. D. 108, 218
Sobel, M. E. 135
Sowden, R. D. 124, 374
Speed, F. M. 252
Spence, I. 11
Spiegelhalter, D. 62
Srole, L. 109
Stokes, M. E. 112
Stone, J. 383
Stuart, A. 60, 128
Sturdivant, R. X. 268

Theus, M. 190, 420
Thornes, B. 171
Thurstone, L. L. 296
Tibshirani, R. J. 443
Trivedi, P. 414
Trivedi, P. K. 414, 415, 433, 436, 455
Tufte, E. 11
Tufte, E. R. 11, 22, 23, 244
Tukey, J. W. 85, 91–93, 95, 124, 454
Tukey, P. A. 11
Turlach, B. A. 466

Upton, G. J. G. 141

van der Heijden, P. G. M. 199, 210–212
Vincente-Villardón, J. L. 230
von Bortkiewicz, L. 61
Von Eye, A. 136
Vuong, Q. H. 400

Wainer, H. 141
Wallace, D. L. 62, 84, 90, 103
Wang, P. C. 289, 290
Warwick, K. M. 199
Wedderburn, R. W. M. 395, 396
Weisberg, S. 27, 289, 307
Whittaker, J. 354
Wickham, H. 11, 186
Wilkinson, L. 11
Williams, C. B. 64, 78
Williams, D. A. 277, 447
Wimmer, G. 65
Winner, L. 147
Wong, R. S.-K. 355, 394
Wood, S. N. 443
Woolf, B. 118
Wright, K. 17

Xie, Y. 359, 363

Yau, K. K. 416
Yip, K. C. 416
Yule, G. U. 75

Zeileis, A. 155, 159, 160, 180, 415, 433, 436
Zelterman, D. 65
Zweiful, J. R. 113

Index

- β -binomial, 100
- ϕ coefficient, 204
- 512 paths to the White House, 19
- 80–20 rule, 22
- `abline()`, 247
- `add1()`, 438
- added-variable plot, 288
- `addmargins()`, 41, 53
- adjusted inertia, 228
- adjusted means, 256
- adjusted residual, 330
- AER** package, 417, 435, 463
- `aes()`, 340
- `aggregate()`, 46, 47, 415
- agreement
 - Cohen’s κ , 140
 - intraclass correlation, 140
 - observer agreement chart, 141
 - partial, 142
- agreement chart, 14
- `agreementplot()`, 143, 144, 150
- agridat** package, 16, 200
- AIC, 329
- `AIC()`, 330, 404, 431
- Akaike Information Criterion, 329
- `allEffects()`, 257, 269, 433, 439
- alpha-blending, 18
- analysis of deviance, 403
- animate** package, 19
- `Anova()`, 263, 267, 311, 320, 334, 339, 403, 463
- `anova()`, 172, 245, 249, 274, 302, 320, 333, 334, 350, 360, 380, 382, 404, 431, 438, 441
- `aperm()`, 39
- `apply()`, 44, 46–48, 75, 121, 181, 183, 451
- Arbuthnot data, 60–62
- `array()`, 32
- Arthritis treatment, 2, 34–36, 51, 113–114, 117–118, 121–123, 165–166, 191–192, 243–244, 250–252, 257–258
- logistic regression, 243
- Arthritis treatment: conditional plots, 252–254
- Arthritis treatment: full-model plots, 254–255
- Arthritis treatment: Plotting logistic regression with base graphics, 246–247
- Arthritis treatment: Plotting logistic regression with **ggplot2**, 247
- `as.data.frame()`, 49, 50, 58, 333
- `as.data.frame.table()`, 49
- `as.matrix()`, 58, 215, 216, 238
- `as.numeric()`, 50, 347, 349
- `as.table()`, 38
- aspect ratio, 209
- `assoc()`, 161, 331
- association, 6
- association graph, 169
- association ordering, 15
- association plot, 136
- association plots, 111
- `assocstats()`, 6, 117–119, 121, 148, 150
- asymmetric CA plot, 231
- asymmetric map, 205, 206
- `avPlot()`, 293
- `avPlots()`, 293, 295
- axes
 - equating, 209
- Barley data, 16–17
- `barplot()`, 406
- Bartlett data on plum root cuttings, 186–187, 193
- baseline models, 168
- Bayesian Information Criterion, 330
- Berkeley admissions, 2, 112, 122–125, 134–136, 188–190, 331–336, 338–341
- BIC, 330
- `BIC()`, 404, 431
- Binary events, 59

- binary response, 240
binary tree, 19
binary variables, 2
`binom.or()`, 391
`binom2.or()`, 376, 381
binomial distribution, 69–72, 95, 96
binomial samples, 115
biplot, 203, 228, 230–236, 357
`biplot()`, 206, 235
bivariate logistic model, 374
bivariate loglinear model, 375
`blogits()`, 376, 383, 389, 393
Breathlessness and wheeze in coal miners, 128–130, 376–384
British social mobility, 15–16
`bs()`, 381
Burt matrix, 223–225
Butterfly species in Malaya, 67
- `c()`, 28, 36, 46
`CA()`, 206
ca package, 206, 208, 226, 231
`ca()`, 206, 207, 215, 219–221, 223
`cabipl()`, 232
canonical analysis of categorical data, 203
canonical correlation, 203
car package, 263, 267, 280, 281, 283, 284, 288, 290, 293, 307, 308, 320, 334, 339, 403, 404, 445, 448, 449, 458, 463
case form, 3
categorical variable, 2
`cbind()`, 33, 458
`cd_plot()`, 422
`cdplot()`, 422
`chisq.test()`, 6, 165
`CMHtest()`, 118–121, 148, 150
Cochran-Mantel-Haenszel tests, 118
general association, 119
linear association, 119, 120
row means differ, 119
Cod parasites, 424–434
`coef()`, 244, 301, 331, 382, 458
`coeftest()`, 244
Cohen's κ , 140
`coindep_test()`, 164, 183
`collapse.table()`, 48, 49, 55
Collapsing categories, 48–49
column effects model, 347
complete separation, 254
component-plus-residual plot, 288, 290
- compositional data, 145
`conditional()`, 174
conditional density plot, 422
conditional distributions, 115
conditional plots, 252
`confint()`, 126, 252
`confint.Kappa()`, 141
`constraints()`, 458, 459
constructed variable plot, 288
contingency ratios, 231
contingency table, 111
`contour()`, 442
`contr.sum()`, 325
`contr.treatment()`, 325
controlled comparison, 128
Cook's distance, 279
`cooks.distance()`, 280, 445
`coplot`, 179
Corporal punishment data, 182–184
`corresp()`, 206
correspondence analysis, 323
asymmetric map, 206
interactive coding, 214
principal coordinates, 205
properties, 205–206
stacking, 214–219
standard coordinates, 205
supplementary variables, 219
symmetric map, 206
two-way tables, 204–213
correspondence matrix, 204
corrplot package, 348
`cotabplot()`, 161, 179, 184, 201, 386
count, 60
count data, 4
count metamer, 96
countreg package, 409, 418, 421, 422, 424, 437
covariate, 103
Cramer's V, 6
cross-sectional study, 115
`Crossings()`, 365
crossings model, 365
`CrossTable()`, 115
`crPlot()`, 290
`crPlots()`, 290
cumulative logit, 297
`cut()`, 410
`cutfac()`, 410, 415, 436
`cutq()`, 415

- data ellipse, 453
 data frame, 27
 data sets
 Abortion, 149
 Accident, 202, 320
 Arbuthnot, 60, 106, 318
 Arrests, 266
 Arthritis, 3, 5, 35, 51, 113, 121, 165,
 191, 241, 243, 245, 250, 252, 299,
 307
 arthritis treatment, 117, 121
 barley, 16
 Bartlett, 186, 193, 200
 Berkeley admissions, 127
 Bundesliga, 107
 Butterfly, 92
 caith, 201, 237
 case2201, 462
 CoalMiners, 128, 376, 377
 CodParasites, 424, 463
 Cormorants, 463
 CrabSatellites, 409, 414, 419, 423
 criminal, 199, 237
 CyclingDeaths, 66, 108
 DanishWelfare, 56
 DaytonSurvey, 46, 47
 death by horse kick, 94, 98
 Depends, 108
 Donner, 258, 263, 319
 Employment, 179, 181
 Federalist, 94, 106
 Federalist papers, 94
 Geissler, 57, 58, 101, 107
 gss8590, 396
 HairEye, 6
 HairEyeColor, 37, 39, 44, 112, 138, 171,
 206
 HairEyePlace, 201, 237
 HallOfFame, 108
 Hauser79, 361, 365
 Health, 343
 HorseKick, 84
 HorseKicks, 50, 52, 64, 87, 98
 Hospital, 150
 housing, 320
 ICU, 270, 285, 294, 295, 319
 iris, 17
 jansen.strawberry, 200
 JobSat, 39, 149, 236
 Lifeboats, 146, 147, 151
 Mammograms, 143, 150
 Master, 199
 Mental, 113, 209, 347, 348, 352, 357
 minnesota.barley.yield, 16
 MSPatients, 139, 144
 multiple sclerosis diagnosis, 145
 NMES1988, 435, 454, 464
 PhdPub, 104, 407, 413, 415, 418, 419,
 448, 463
 PreSex, 175, 187, 226, 238
 Punishment, 182
 quine, 462
 RepVict, 211
 Saxony, 57, 99, 101, 103
 SexualFun, 139, 141
 Space shuttle disaster, 8–9
 SpaceShuttle, 248
 struc, 196
 Suicide, 216
 Titanic, 146, 228
 Toxaemia, 385, 390
 Toxemia, 393–395
 TV, 213, 237, 320
 UCBADmissions, 56
 UCBAdmissions, 58, 112, 122, 190, 248,
 330, 331, 333, 341
 UCBadmissions, 338
 UKSoccer, 57, 73, 74, 107, 167, 235
 Vision, 358
 visual acuity, 134
 VisualAcuity, 58, 132, 134, 150, 359,
 369, 396
 WeldonDice, 63, 70, 86
 Womenlf, 307, 311, 319
 WomenQueue, 95, 106
 data.frame(), 32, 36, 50
 datasets package, 37
 datasets(), 56
 Dayton survey, 46–48
 dbinom(), 68, 71
 ddoublebinom(), 101
 ddply(), 48
 Death by horse kick, 50, 64, 84–85, 87, 93–
 94, 98
 Death in the ICU, 269–277, 285–288, 294–
 296
 Demand for medical care, 435–443, 454–460
 density(), 422
 depvar(), 382
 deviance, 328
 deviance residual, 278, 330, 444
 DFBETA, 279

- `dfbeta()`, 280
`dfbeta.glm()`, 280
`DFFITS`, 280
`dffits()`, 280
`dgeom()`, 68
`Diag()`, 359, 365, 370
Diagnosis of MS patients, 139, 144–145
dichotomous, 240
dichotomous variables, 2
`dim()`, 29
`dimnames()`, 30
`direct.label()`, 77
directlabels package, 23, 76, 277, 304
dispersion parameter, 79, 401
`dispersiontest()`, 417, 463
`distplot()`, 82, 97–100, 106–108
`dlogseries()`, 68
`dnbinom()`, 68, 79
Donner Party, 9–10, 258–266, 280–285, 290–291, 293–294
Donner party, 21
`dotchart()`, 353
double binomial, 101
`doubledecker()`, 161
doubledecker plots, 158
`dpois()`, 68, 76
dual scaling, 203, *see* correspondence analysis
dynamic graphics, 19
- Effect** (), 256–258, 306
effect displays, 256
effect ordering, 158, 213
effect plot, 194
effect-order sorting, 15
effects package, 256, 257, 266, 269, 306, 314, 317, 389, 433, 434, 460
empirical logits, 376
Employment status data, 179–182
`eqsplot()`, 24
events/trials form, 248
`exp()`, 252, 382
`expand.dft()`, 51, 75
`expand.grid()`, 36, 71, 75, 80
explanatory, 112
explanatory variables, 5
exponential family, 100, 400
- `facet_grid()`, 253, 304, 428
`facet_wrap()`, 253, 255
facets, 253
- FactoMineR** package, 206
`factor`, 32
factor analysis, 298
Families in Saxony, 4, 5, 62, 85–86, 99–103
Federalist papers, 64–65, 88–89, 94, 100
`fitdistr()`, 85
`fitted()`, 194, 332, 362, 382, 415
foreign package, 34
`fourfold()`, 127, 148, 149
fourfold display
 confidence rings, 125–126
fourfold display, 14, 123–130
`fractions()`, 242
frequency, 60
frequency data, 4
frequency form, 3
`ftable()`, 37, 41, 43, 57, 58, 215, 216
- `gam()`, 442
gdata package, 34
General social survey, 36–37, 50
generalized additive model, 442
generalized linear mixed models, 400
generalized linear model, 240, 241, 400
generalized logit model, 297
generalized pairs plot, 186
`geom_boxplot()`, 426
`geom_density2d()`, 464
`geom_jitter()`, 426
`geom_point()`, 247, 249, 252
`geom_ribbon()`, 254, 276
`geom_smooth()`, 249
geometric distribution, 80–82
`getContrasts()`, 353
GGally package, 191
`ggplot()`, 247, 253, 254, 260, 277, 304
ggplot2 package, 11, 24, 76, 77, 146, 191, 245, 247, 249, 252, 261, 312, 340, 394, 406, 426, 428, 457, 464
- ggttern** package, 146
`ggttern()`, 146
`glm()`, 5, 100–102, 239, 243, 244, 248, 249, 251, 254, 256, 272, 278, 320, 323, 325, 327, 328, 333–335, 338, 339, 342, 343, 345, 347, 348, 352, 377, 380, 381, 384, 389, 396, 410, 413, 414, 421, 460
`glm.nb()`, 414, 434, 442, 445, 454, 458, 460
`glmlist()`, 367
`gls()`, 256

gnm package, 352, 353, 359
gnm(), 352, 357, 365, 396
goodfit(), 82, 85–87, 90, 99, 100, 105–109
 goodness-of-fit, 83
googleVis package, 19
gpairs package, 186, 191
gpairs(), 191, 192, 319, 409
grapcon functions, 159
graphics
 interactive, 19
graphics package, 422
grid package, 161, 191

Hair color and eye color, 6–7, 37, 112, 132, 154–158, 206–209, 223–224
 Hair color, eye color and sex, 168, 171–172
 half-normal plot, 450
 hanging rootogram, 90
hatvalues(), 279
 Hauser’s occupational mobility table, 361–369
 Health concerns of teenagers, 343–345
heplot(), 455
heplot3d(), 455
heplots package, 455
 heterogeneous uniform association, 369
 hierarchical models, 325
 high-order terms, 168
HistData package, 60, 106
hmmm package, 374
 homogeneity analysis, 203, *see* correspondence analysis
 homogeneity of association, 6
 homogeneous association model, 326
hurdle(), 422
 hurdle model, 422
 hypothesis-error plot, 453

ICC(), 140
identify(), 23
image(), 442
independence_table(), 131
 index of dispersion, 75
 indicator matrix, 222
 inertia, 204
 influence, 278
influence.measures(), 280, 286
influenceIndexPlot(), 284
influencePlot(), 281, 283, 448
 inter-rater agreement, 111, *see* agreement
interaction(), 58, 216

interactive coding, 214
 interactive graphics, 19
interp(), 163
 Interpolation options, 163
 intraclass correlation, 140
iplots package, 19
 Iris data, 17–18

Job satisfaction, 38
joint(), 174
 joint correspondence analysis, 228
 joint distribution, 114
 joint independence, 136
 joint influence, 292

Kappa(), 141
kde2d(), 464
KernSmooth package, 261
 Kruskal-Wallis test, 119
Kway(), 370

labeling_border(), 160
labeling_cboxed(), 160
labeling_cells(), 160
labeling_conditional(), 160
labeling_doubledecker(), 160
labeling_lboxed(), 160
labeling_left(), 160
labeling_left2(), 160
labeling_list(), 160
labeling_residuals(), 160
labeling_value(), 160
Lahman package, 108, 199
 latent class analysis, 299
lattice package, 16, 24, 71, 76
 least squares means, 256
legend(), 212
legend_fixed(), 160
legend_resbased(), 160
levels(), 259, 317
 levels model, 364
 Lifeboats on the *Titanic*, 146–148
 likelihood ratio test, 119
 linear hypothesis, 402
 linear logistic regression model, 241
 linear logit model, 242
 linear predictor, 250, 400
 linear probability model, 240
 linear probit regression, 242
 linear-by-linear model, 346
linearHypothesis(), 403, 404, 458, 460

- `lines()`, 70, 247, 312, 415
`link function`, 400
`lme()`, 5, 244, 256, 278, 325, 455
`lme4 package`, 256
`lmer()`, 256
`lmtest package`, 244, 392, 431
`lmtest()`, 437
`loddsratio()`, 348, 361, 456
`log odds`, 8, 116, 242
`log odds ratio`, 116, 234
 - local, 346`logarithmic series distribution`, 82, 93, 94
`logi.hist.plot()`, 243
`logistic regression`
 - influence diagnostics, 279–280
 - leverage, 279
 - residuals, 278`logit`, 116, 242
`logit function`, 7
`logLik()`, 404, 431
`loglin()`, 174, 327, 328, 342
`loglin2formula()`, 174
`loglin2string()`, 174
`loglinb2()`, 376
`loglinear independence model`, 324
`loglinear model`, 122
`loglm()`, 123, 136, 150, 156, 168, 172, 174, 179, 181, 183, 194, 199, 201, 202, 323, 328, 331, 334, 342, 345, 396
`logmult package`, 199, 237, 352, 354, 356, 358, 396
`London cycling deaths`, 65–66, 75
`lowess()`, 247
`lrm()`, 272, 275, 297
`LRstats()`, 172, 245, 264, 368, 404
`LRtest()`, 271, 310
`lrtest()`, 302, 392, 431
`lsmeans package`, 256
- `main-effect ordering`, 15
`Mammogram ratings`, 143–144
`manipulate package`, 19
`margin.table()`, 40, 41, 46, 47
`marginal distributions`, 114
`marginal frequencies`, 114
`marginal homogeneity`, 140, 144, 359
`marginal model`, 219
`Marimekko chart`, 162
`Marital status and pre- and extramarital sex`, 175–177, 187–188, 226–228
- `markov()`, 174
`masking`, 292
`MASS package`, 24, 51, 85, 123, 172, 201, 206, 237, 242, 256, 273, 297, 299, 320, 328, 331, 414, 420, 462
`Mating of horseshoe crabs`, 409–412, 414, 418, 422–423
`matplot()`, 362, 377, 382, 388
`matpoints()`, 382
`matrix`, 29
`matrix()`, 30
`mca()`, 51, 206
`mcja()`, 224
`mean function`, 400
`mean-square contingency coefficient`, 204
`melt()`, 304
`Mental impairment and parents' SES`, 113, 118, 209–211, 347–357
`mgcv package`, 442
`mjca()`, 206, 223, 226, 228, 238
`mlogit package`, 314
`model comparison plot`, 368
`model matrix`, 327
`model object`, 240
`model.matrix()`, 222
`mosaic()`, 161, 162, 165, 168, 174, 179, 182, 184, 191, 199, 201, 202, 331, 334, 343, 385
`mosaic display`, 7, 153, 323
`mosaic matrix`, 185, 222
`mosaic3d()`, 193, 197, 331
`mosaicplot()`, 42, 160
`Mult()`, 352
`multinom()`, 256, 297, 314, 320
`multinomial logit model`, 297
`multinomial sample`, 115
`multiple correspondence analysis`
 - bivariate, 222–224`multivariate linear model`, 453
`mutual()`, 174
- `natural spline`, 263
`negative binomial distribution`, 77–80, 94, 96
`nested dichotomies`, 297, 306–313
`nested models`, 329
`nlme package`, 256
`nnet package`, 256, 297, 314
`nobs()`, 404
`nominal`, 2
`nomogram`, 275
`nomogram()`, 275

normal QQ plots, 449
 normal quantile plots, 449
`ns()`, 263, 381, 383
 observer agreement, 138
 observer agreement chart, 141
 odds, 116
 odds ratio, 20, 116–117, 122
`oddsratio()`, 129, 149, 388
 optimal scaling, 203, *see* correspondence analysis
`Ord` plot, 92–96
`Ord_plot()`, 92, 94, 96, 97, 105, 108
`ordered()`, 35, 39, 299, 314, 317
 ordinal, 2
`outlierTest()`, 448
 overdispersion, 77, 401
`p.adjust()`, 127
 package
 AER, 417, 435, 463
 agridat, 16, 200
 animate, 19
 ca, 206, 208, 226, 231
 car, 263, 267, 280, 281, 283, 284, 288,
 290, 293, 307, 308, 320, 334, 339,
 403, 404, 445, 448, 449, 458, 463
 corrplot, 348
 countreg, 409, 418, 421, 422, 424, 437
 datasets, 37
 directlabels, 23, 76, 277, 304
 effects, 256, 257, 266, 269, 306, 314,
 317, 389, 433, 434, 460
 FactoMineR, 206
 foreign, 34
 gdata, 34
 GGally, 191
 ggplot2, 11, 24, 76, 77, 146, 191, 245,
 247, 249, 252, 261, 312, 340, 394,
 406, 426, 428, 457, 464
 ggtern, 146
 gnm, 352, 353, 359
 googleVis, 19
 gpairs, 186, 191
 graphics, 422
 grid, 161, 191
 heplots, 455
 HistData, 60, 106
 hmmm, 374
 iplots, 19
 KernSmooth, 261

Lahman, 108, 199
 lattice, 16, 24, 71, 76
 lme4, 256
 lmtest, 244, 392, 431
`logmult`, 199, 237, 352, 354, 356, 358,
 396
 lsmeans, 256
 manipulate, 19
 MASS, 24, 51, 85, 123, 172, 201, 206,
 237, 242, 256, 273, 297, 299, 320,
 328, 331, 414, 420, 462
 mgcv, 442
 mlogit, 314
 nlme, 256
 nnet, 256, 297, 314
 plyr, 48
 poLCA, 256
 probio, 243
 pscl, 404, 421
 psych, 140
 rCharts, 19
 reshape2, 304
 rggobi, 19
 rgl, 193, 206
 rms, 275, 297, 303
 rmutil, 101
 rsm, 271, 442
 sandwich, 413
 shiny, 19
 Sleuth2, 462
 splines, 263, 463
 stats, 43, 68, 327
 TeachingDemos, 146
 texreg, 52
 UBbipl, 232, 233
 vcd, 3, 35, 43, 56–58, 63, 64, 73, 85, 90,
 92, 95, 97, 122, 128, 129, 131, 132,
 134, 139, 141, 146, 147, 150, 159,
 161, 163, 164, 175, 179, 182, 183,
 185, 191, 211, 215, 216, 248, 331,
 388, 418, 422, 474
 vcdExtra, 46, 48, 51, 54, 56, 57, 66, 82,
 101, 104, 107, 108, 113, 118, 143,
 149, 150, 172, 174, 186, 193, 201,
 202, 236, 237, 245, 258, 264, 270,
 331, 334, 348, 361, 365, 370, 376,
 385, 404, 410, 463
 VGAM, 297, 301, 302, 314, 374, 376,
 381, 382, 420, 454, 458, 460
 XLConnect, 34
 xlsx, 34

- xtable**, 52
pairs(), 161, 185, 189, 196, 455
pairs.table(), 190, 198
palette(), 162
panel functions, 190
parallel coordinate plot, 17
Pareto chart, 22
Pareto distribution, 22
Pareto principle, 22
parquet diagram, 131, *see* sieve diagram
partial association, 177
partial proportional odds model, 300
partial residual plot, 290
partial-regression plot, 291
Pascal distribution, 78
paste(), 39, 216, 217
pbinom(), 68
pchisq(), 84
pdoublebinom(), 101
Pearson residual, 278, 330, 444
persp(), 442
persp.lm(), 442
pgeom(), 68
phi coefficient, 6
phi (ϕ) coefficient, 204
pickCoef(), 354, 365
plogis(), 255
plogseries(), 68
plot(), 44, 70, 129, 209, 226, 229, 231, 240, 257, 269, 271, 272, 280, 283, 357, 388, 418, 439, 445, 448, 449
plot.ca(), 206–208
plot.goodfit(), 89
plot.gootfit(), 106
plot.rc(), 357
plot.xmean.ordinaly(), 303
plot3d.ca(), 206
Plotting styles for discrete distributions, 76–77
plyr package, 48
pnbnom(), 68
Poisson distribution, 72–77, 94, 96
Poisson regression, 324
Poissonness plot, 96–98
poLCA package, 256
polr(), 256, 297, 299–301, 306, 315, 320
poly(), 381, 383
Polya distribution, 78
polygon(), 247
polytomous, 240
Polytomous events, 60
polytomous response, 240
polytomous variables, 2
population marginal means, 256
position_jitter(), 247
power series distributions, 83
ppois(), 68, 75
prcomp(), 235
predict(), 240, 246, 247, 254, 256, 257, 303, 311, 442
principal component analysis, 203, 222
principal coordinates, 205
princomp(), 235
print(), 53, 85, 240, 244, 257, 271
print.goodfit(), 85
probio package, 243
prop.table(), 40, 41
proportional odds model, 297–306
pscl package, 404, 421
psych package, 140
Publications of PhD candidates, 103–105, 405–409, 413–418, 445–448, 450–452
qbinom(), 68
qdoublebinom(), 101
qgeom(), 68
qlogis(), 383
qlogseries(), 68
qnbinom(), 68
qnorm(), 449
qpois(), 68
qqPlot(), 449
quasi-independence, 134
quasi-independence model, 358
quasi-Poisson, 401
quasi-Poisson model, 413
quasi-symmetry, 359
Racial profiling: Arrests for marijuana possession, 11–13, 266–269
radial diagram, 123
rainbow(), 442
raw residual, 330, 444
rbinom(), 68
rc(), 352, 356, 357, 397
rCharts package, 19
rcL(), 357
rdoublebinom(), 101
read.csv(), 34
read.delim(), 34
read.table(), 34, 54

- reciprocal averaging, 203, *see* correspondence analysis
`recode()`, 308
 regression quartet, 280
 regression spline, 262
`relevel()`, 251, 313, 314
 rendering, 15
`rep()`, 29
 Repeat victimization, 211–213
reshape2 package, 304
 residual deviance, 403
`residualPlot()`, 446
 residuals
 standardized, 278
 studentized, 278
`residuals()`, 156, 278, 331
 response, 112
 response residual, 444
 response variables, 5
`rgeom()`, 68
rggobi package, 19
rgl package, 193, 206
`rlogseries()`, 68
rms package, 275, 297, 303
rmutil package, 101
`rnbnom()`, 68
`rnegbin()`, 420, 450
`rnorm()`, 450
`rnorm()()`, 33
 rootogram, 89
`rootogram()`, 90, 105, 108, 418
 row effects model, 347
 row plus column effects model, 347
 row-and-column effects model, 351
`rownames()`, 53
`rpois()`, 68, 237, 420, 450
rsm package, 271, 442
`rstandard()`, 278, 445
`rstudent()`, 278, 445
`rzinegbin()`, 420
`rzipois()`, 420, 421
- `s()`, 443
`sample()`, 33
 sample odds ratio, 117
 sampling zeros, 342
sandwich package, 413
`sandwich()`, 413, 416
`sapply()`, 183
 saturated model, 169, 245, 325
`scale()`, 355
 scale parameter, 401
`scale_y_log10()`, 406
 scatterplot matrix, 184, 222
`scatterplotMatrix()`, 288
 scree plot, 207
`segments()`, 212
`seq()`, 29
`seq_loglm()`, 175
`set.seed()`, 33, 164
 Sex is fun, 139
 Shading functions, 163–164
`shading_binary()`, 160
`shading_Friendly()`, 160
`shading_hcl()`, 160
`shading_hsv()`, 160
`shading_max()`, 160, 164, 166
`shading_sieve()`, 160
shiny package, 19
`sieve()`, 161, 331
 sieve diagram, 131–136
 sieve diagrams, 111
 simple effects, 178
 Simpson’s paradox, 127
`simulate()`, 450
 Simulating zero-inflated data, 420–421
 singular value decomposition, 204, 205
Sleuth2 package, 462
`sort()`, 39
 Space shuttle disaster, 8–9, 21–22, 248–249
`spacing_conditional()`, 160
`spacing_dimequal()`, 160
`spacing_equal()`, 160
`spacing_highlighting()`, 160
`spacing_increase()`, 160
 spaghetti plot, *see* parallel coordinates plot
 spine plot, 422
`spineplot`, 259
`spineplot()`, 422, 436
`spinogram`, 422
splines package, 263, 463
 stacking, 214
 standard coordinates, 205
 standardized residuals, 278, 444
`stat_smooth()`, 247, 252, 261
stats package, 43, 68, 327
`stepAIC()`, 273, 320
`str()`, 30, 39
 stratified analysis, 6, 121
 stratifying variable, 111
`struc_assoc()`, 160
`struc_mosaic()`, 160

- `struc_sieve()`, 160
`strucplot()`, 161
strucplot framework, 134, 159
`structable()`, 37, 41, 43, 48, 57, 58, 159, 215
structural zeros, 341
studentized residual, 444
studentized residuals, 278
`subset()`, 45, 46, 49, 58
Suicide rates in Germany, 216–218
Suicide rates in Germany: biplot, 231–233
Suicide rates in Germany: mosaic plot, 218–219
`sum()`, 47
`summarise()`, 48
`summary()`, 85, 226, 240, 244, 257, 263, 271, 272, 283, 300, 315, 334, 339, 407, 413, 463
`summary.goodfit()`, 85
`summary.Kappa()`, 141
supplementary variables, 220
Survival on the *Titanic*, 228–230
`Symm()`, 359, 370
symmetric map, 205, 206
symmetry, 134
symmetry model, 359
- `t()`, 31, 44, 53
`table()`, 32, 37, 40–42, 44, 51, 406
table form, 4
TeachingDemos package, 146
ternary plot, 112
`texreg` package, 52
`theme()`, 77
`toeplitz()`, 162
`Topo()`, 359, 364
topological model, 364
Toxaemic symptoms in pregnancy, 385–394
treemap, 19
trilinear plot, 112, 145
`triplot()`, 146
TV viewing data, 53–214
type-token, 66
- `UBbipl` package, 232, 233
UK Soccer scores, 73–75, 167, 234–236
uniform association model, 346
uniform interaction, 369
`update()`, 271, 274, 317, 360, 441
- variable
- response \hat{z} , 5
`vcd` package, 3, 35, 43, 56–58, 63, 64, 73, 85, 90, 92, 95, 97, 122, 128, 129, 131, 132, 134, 139, 141, 146, 147, 150, 159, 161, 163, 164, 175, 179, 182, 183, 185, 191, 211, 215, 216, 248, 331, 388, 418, 422, 474
`vcdExtra` package, 46, 48, 51, 54, 56, 57, 66, 82, 101, 104, 107, 108, 113, 118, 143, 149, 150, 172, 174, 186, 193, 201, 202, 236, 237, 245, 258, 264, 270, 331, 334, 348, 361, 365, 370, 376, 385, 404, 410, 463
`vcov()`, 256, 416
vector, 27, 28
`VGAM` package, 297, 301, 302, 314, 374, 376, 381, 382, 420, 454, 458, 460
`vglm()`, 297, 301, 302, 374, 376, 381, 382, 384, 390, 391, 458
Visual acuity, 132–134, 359–361, 369–373
visual impact, 128
`vuong()`, 404
Vuong’s test, 404
- `weighted.mean()`, 50, 84
Weldon’s dice, 63, 70–71, 86–87
`with()`, 40
`within()`, 54, 74
Women in queues, 95
Women’s labor force participation, 307–318
- `XLConnect` package, 34
`xlsx` package, 34
`xtable` package, 52
`xtable()`, 52, 58
`xtabs()`, 32, 37, 40, 42, 44, 47, 50, 55, 58, 117, 216, 217, 219, 259, 456
`xyplot()`, 71, 76, 77, 80, 106, 355
- zero-altered model, 422
zero-inflated Poisson, 419
zero-truncated distribution, 108
`zeroinfl()`, 421, 422

This document was produced using:

```
> print(sessionInfo(), locale = FALSE)

R version 3.1.2 (2014-10-31)
Platform: x86_64-pc-linux-gnu (64-bit)

attached base packages:
[1] stats4      splines     grid        stats       graphics   grDevices
[7] utils       datasets    methods    base

other attached packages:
[1] VGAM_0.9-3           mgcv_1.8-3
[3] nlme_3.1-119         countreg_0.1-1
[5] sandwich_2.3-0       nnet_7.3-7
[7] reshape2_1.2.2       proto_0.3-10
[9] car_2.0-22          lmtest_0.9-32
[11] zoo_1.7-11          MASS_7.3-37
[13] gmodels_2.15.4.1    directlabels_2013.6.15
[15] quadprog_1.5-5      xtable_1.7-1
[17] vcdExtra_0.6-5      gnm_1.0-6
[19] lattice_0.20-29    ggplot2_0.9.3.1
[21] ca_0.56             vcd_1.3-3
[23] knitr_1.8

loaded via a namespace (and not attached):
[1] codetools_0.2-8    colorspace_1.2-4  dichromat_2.0-0
[4] digest_0.6.4       evaluate_0.5.5   formatR_1.0
[7] gdata_2.13.3       gtable_0.1.2    gtools_3.4.1
[10] highr_0.4          labeling_0.2   Matrix_1.1-4
[13] munsell_0.4.2      plyr_1.8       qvcalc_0.8-8
[16] RColorBrewer_1.0-5 relimp_1.0-3  scales_0.2.3
[19] stringr_0.6.2      tcltk_3.1.2   tools_3.1.2
```