

# A Second Semester Statistics Course with R

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I (Greenwood) have intentionally taken a first person perspective at times to be able to include stories from some of those interactions to try to help you avoid some of their pitfalls in your current or future usage of statistics. I would like to thank my wife, Teresa Greenwood, for allowing me the time and support to work on this. I would also like to acknowledge Dr. Gordon Bril (Luther College) who introduced me to statistics while I was an undergraduate and Dr. Snehalata Huzurbazar (University of Wyoming) that guided me to completing my Master's and Ph.D. in Statistics and still serves as a valued mentor and friend to me.

The development of this text was initially supported with funding from Montana State University's Instructional Innovation Grant Program with a grant titled Towards more active learning in STAT 217. This book was born with the goal of having a targeted presentation of topics that we cover (and few that we don't) that minimizes cost to students and incorporates the statistical software R from day one and every day after that. The software is a free, open-source platform and so is dynamically changing over time. This has necessitated frequent revisions of the text.

This is Version 3.01 of the book. It fixes a problem created with the digital links in the book that occurred during Spring 2017. Version 3.0 of the book, prepared for Fall 2016, involved edits, a couple of partially new sections, and updated R code along with a new format for how the R code is displayed to more easily distinguish it from other text. Each revision has involved a similar amount of change with Version 2.0 published in January 2015 and Version 1.0 in January 2014 after using draft chapters that were initially developed during Fall 2013.

We have made every attempt to keep costs as low as possible by making it possible for most pages to be printed in black and white. The text (in full color and with dynamic links) is also available as a free digital download from Montana State University's ScholarWorks repository at <https://scholarworks.montana.edu/xmlui/handle/1/2999>.

Enjoy your journey from introductory to intermediate statistics!

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# Chapter 1

## Preface

This book is designed primarily for use in a second semester statistics course although it can also be useful for researchers needing a quick review or ideas for using R for the methods discussed in the text. As a text primarily designed for a second statistics course, it presumes that you have had an introductory statistics course. There are now many different varieties of introductory statistics from traditional, formula-based courses (called “consensus” curriculum courses) to more modern, computing-intensity courses that use randomization ideas to try to enhance learning of basic statistical methods. We are not going to presume that you have had a particular “flavor” of introductory statistics or that you had your introductory statistics out of a particular text, just that you have had a course that tried to introduce you to the basic terminology and ideas underpinning statistical reasoning. We would expect that you are familiar with the logic (or sometimes illogic) of hypothesis testing including null and alternative hypothesis and confidence interval construction and interpretation and that you have seen all of this in a couple of basic situations. We start with a review of these ideas in one and two group situations with a quantitative response, something that you should have seen before.

This text covers a wide array of statistical tools that are connected through situation, methods used, or both. As we explore various techniques, look for the identifying characteristics of each method – what type of research questions are being addressed (relationships or group differences, for example) and what type of variables are being analyzed (quantitative or categorical). **Quantitative variables** are made up of numerical measurements that have meaningful units attached to them. **Categorical variables** take on values that are categories or labels. Additionally, you will need to carefully identify the **response** variables, where the study and variable characteristics should suggest which variables should be used as the explanatory variables that may explain variation in the response variable. Because this is an intermediate statistics course, we will start to handle more complex situations (many explanatory variables) and will provide some tools for graphical explorations to complement the more sophisticated statistical models required to handle these situations.

### 1.1 Overview of methods

After you are introduced to basic statistical ideas, a wide array of statistical methods become available. The methods explored here focus on assessing (estimating and testing for) relationships between variables, sometimes when controlling for or modifying relationships based on levels of another variable – which is where statistics gets interesting and really useful. Early statistical analyses (approximately 100 years ago) were focused on describing a single variable. Your introductory statistics course should have heavily explored methods for summarizing and doing inference in situations with one group or where you were comparing results for two groups of observations. Now, we get to consider more complicated situations – culminating in a set of tools for working with multiple explanatory variables, some of which might be categorical and related to having different groups of subjects that are being compared. Throughout the methods we will cover, it will

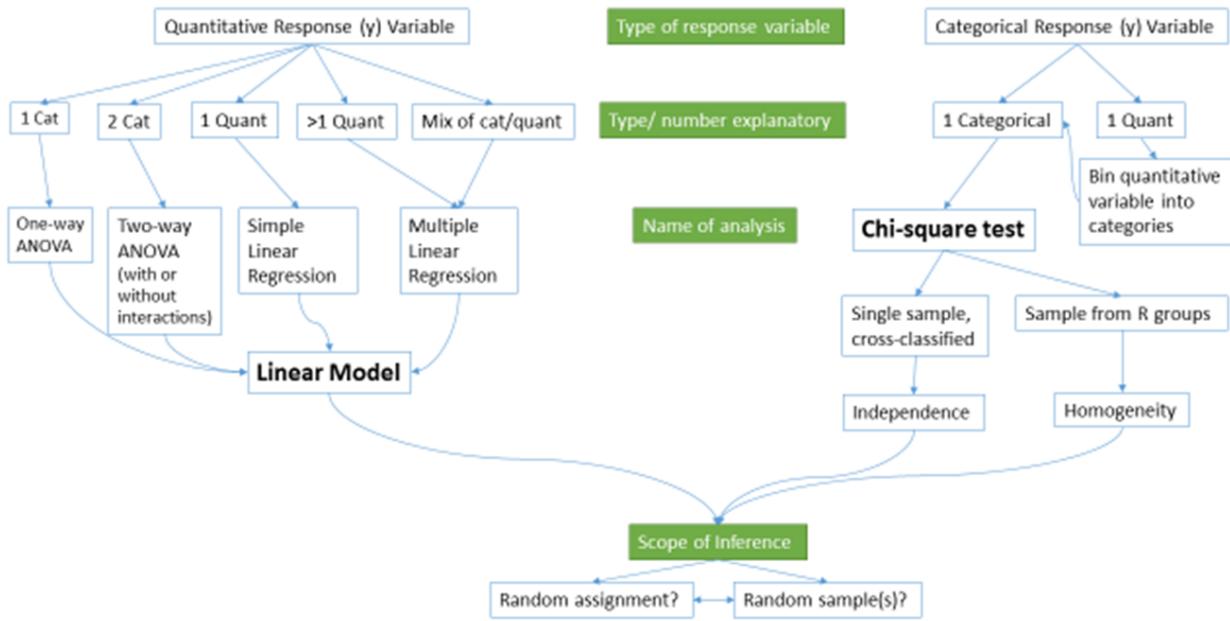


Figure 1.1: Flow chart of methods

be important to retain a focus on how the appropriate statistical analysis depends on the research question and data collection process as well as the types of variables measured.

Figure 1.1 frames the topics we will discuss. Taking a broad vision of the methods we will consider, there are basically two scenarios – one when the response is quantitative and one when the response is categorical. Examples of quantitative responses we will see later involve *suggested jail sentence* (in years) and *body fat* (percentage). Examples of categorical variables include *improvement* (none, some, or marked) in a clinical trial or whether a student has turned in copied work (never, exam or paper, or both). There are going to be some more nuanced aspects to all these analyses as the complexity of both sides of Figure 1.1 suggest, but note that near the bottom, each tree converges on a single procedure, using a **linear model** for a quantitative response variable or using a **Chi-square test** for a categorical response. After selecting the appropriate procedure and completing the necessary technical steps to get results for a given data set, the final step involves assessing the scope of inference and types of conclusions that are appropriate based on the design of the study.

We will be spending most of the semester working on methods for quantitative response variables (the left side of Figure 1.1 covered in Chapters 2, 3, 4, 6, 7, and 8) and stepping over to handle the situation with a categorical response variable (right side of Figure 1.1 that is discussed in Chapter 5). Chapter 9 contains case studies illustrating all the methods discussed previously, providing a final opportunity to explore additional examples that illustrate how finding your way through the paths in Figure 1.1 leads to the appropriate analysis.

The first topics (Chapters 1, and 2) will be more familiar as we start with single and two group situations with a quantitative response. In your previous statistics course, you should have seen methods for estimating and quantifying uncertainty for the mean of a single group and for differences in the means of two groups. Once we have briefly reviewed these methods and introduced the statistical software that we will use throughout the course, we will consider the first new statistical material in Chapter 3. It involves the situation with a quantitative response variable where there are more than 2 groups to compare – this is what we call the **One-Way ANOVA** situation. It generalizes the 2-independent sample hypothesis test to handle situations where more than 2 groups are being studied. When we learn this method, we will begin discussing model

assumptions and methods for assessing those assumptions that will be present in every analysis involving a quantitative response. The ***Two-Way ANOVA*** (Chapter 3) considers situations with two categorical explanatory variables and a quantitative response. To make this somewhat concrete, suppose we are interested in assessing differences in, say, the *yield* of wheat from a field based on the amount of *fertilizer* applied (none, low, or high) and *variety* of wheat (two types). Here, *yield* is a quantitative response variable that might be measured in bushels per acre and there are two categorical explanatory variables, *fertilizer*, with 3 levels, and *variety*, with two levels. In this material, we introduce the idea of an ***interaction*** between the two explanatory variables: the relationship between one categorical variable and the mean of the response changes depending on the levels of the other categorical variable. For example, extra fertilizer might enhance the growth of one variety and hinder the growth of another so we would say that *fertilizer* has different impacts based this interaction may or may not actually be present, we will consider two versions of the model in Two-Way ANOVAs, what are called the ***additive*** (no interaction) and the ***interaction*** models.

Following the methods for two categorical variables and a quantitative response, we explore a method for analyzing data where the response is categorical, called the ***Chi-square test*** in Chapter 5. This most closely matches the One-Way ANOVA situation with a single categorical explanatory variable, except now the response variable is categorical. For example, we will assess whether taking a drug (vs taking a *placebo*<sup>1</sup>) has an ***effect***<sup>2</sup> on the type of improvement the subjects demonstrate. There are two different scenarios for study design that impact the analysis technique and hypotheses tested in Chapter 5. If the explanatory variable reflects the group that subjects were obtained from, either through randomization of the treatment level to the subjects or by taking samples from separate populations, this is called a ***Chi-square Homogeneity Test***. It is also possible to obtain a single sample from a population and then obtain information on the levels of the explanatory variable for each subject. We will analyze these results using what is called a ***Chi-square Independence Test***. They both use the same test statistic but we use slightly different graphics and are testing different hypotheses in these two related situations. Figure 1.1 also shows that if we had a quantitative explanatory variable and a categorical response that we would need to “bin” or create categories of responses from the quantitative variable to use the Chi-square testing methods.

If the predictor and response variables are both quantitative, we start with scatterplots, correlation, and ***simple linear regression*** models (Chapters 6 and 7) – things you should have seen, at least to some degree, previously. The biggest differences here will be the depth of exploration of diagnostics and inferences for this model and discussions of transformations of variables. If there is more than one explanatory variable, then we say that we are doing ***multiple linear regression*** (Chapter 8) – the “multiple” part of the name reflects that there will be more than one explanatory variable. We use the same name if we have a mix of categorical and quantitative predictor variables but there are some new issues in setting up the models and interpreting the coefficients that we need to consider. In the situation with one categorical predictor and one quantitative predictor, we revisit the idea of an interaction. It allows us to consider situations where the estimated relationship between a quantitative predictor and the mean response varies among different levels of the categorical variable.

By the end of Chapter 9 you should be able to identify, perform using the statistical software R (R Core Team, 2017), and interpret the results from each of these methods. There is a lot to learn, but many of the tools for using R and interpreting results of the analyses accumulate and repeat during the semester. If you work hard to understand the initial methods, it will help you when the methods get more complicated. You will likely feel like you are just starting to learn how to use R at the end of the semester and for learning a new language that is actually an accomplishment. We will just be taking you on the first steps of a potentially long journey and it is up to you to decide how much further you want to go with learning the software.

All the methods you will learn require you to carefully consider how the data were collected, how that pertains to the population of interest, and how that impacts the inferences that can be made. The ***scope of inference*** from the bottom of Figure 1.1 is our shorthand term for remembering to think about two aspects

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<sup>1</sup>A *placebo* is a treatment level designed to mimic the potentially efficacious level(s) but that can have no actual effect. The *placebo effect* is the effect that thinking that an effective treatment was received has on subjects. There are other related issues in performing experiments like the *Hawthorne* or observer *effect* where subjects modify behavior because they are being observed.

<sup>2</sup>We will reserve the term “effect” for situations where we could potentially infer causal impacts on the response of the explanatory variable which occurs in situations where the levels of the explanatory variable are randomly assigned to the subjects.

of the study – *random assignment* and *random sampling*. In a given situation, you need to use the description of the study to decide if the explanatory variable was randomly assigned to study units (this allows for *causal inferences* if differences are detected) or not (so no causal statements are possible). As an example, think about two studies, one where students are randomly assigned to either get tutoring with their statistics course or not and another where the students are asked at the end of the semester whether they sought out tutoring or not. Suppose we compare the final grades in the course for the two groups (tutoring/not) and find a big difference. In the first study with random assignment, we can say the tutoring caused the differences we observed. In the second, we could only say that the tutoring was associated with differences but because students self-selected the group they ended up in, we can't say that the tutoring caused the differences. The other aspect of scope of inference concerns random sampling: If the data were obtained using a random sampling mechanism, then our inferences can be safely extended to the population that the sample was taken from. However, if we have non-random sample, our inference can only apply to the sample collected. In the previous example, the difference would be studying a random sample of students from the population of, say, Introductory Statistics students at a university vs studying a sample of students that volunteered for the research project, maybe for extra credit in the class. We could still randomly assign them to tutoring/not but the non-random sample would only lead to conclusions about those students that volunteered. The most powerful scope of inference is when there are randomly assigned levels of explanatory variables with a random sample from population – conclusions would be about causal impacts that would happen in the population.

By the end of this material, you should have some basic R skills and abilities to create basic ANOVA and Regression models, as well as to handle Chi-squared testing situations. Together, this should prepare you for future statistics courses or for other situations where you are expected to be able to identify an appropriate analysis, do the calculations for a given data set, and then effectively communicate interpretations for the methods discussed here.

## 1.2 Getting started in R

You will need to download the statistical software package called R and an enhanced interface to R called RStudio (RStudio Team, 2015). They are open source and free to download and use (and will always be that way). This means that the skills you learn now can follow you the rest of your life. R is becoming the primary language of statistics and is being adopted across academia, government, and businesses to help manage and learn from the growing volume of data being obtained. Hopefully you will get a sense of some of the power of R in this book.

The next pages will walk you through the process of getting the software downloaded and provide you with an initial experience using RStudio to do things that should look familiar even though the interface will be a new experience. Do not expect to master R quickly – it takes years (sorry!) even if you know the statistical methods being used. We will try to keep all your interactions with R code in a similar code format and that should help you in learning how to use R as we move through various methods. We will also usually provide you with example code. Everyone that learns R starts with copying other people's code and then making changes for specific applications – so expect to go back to examples from the text and focus on learning how to modify that code to work for your particular data set. Only really experienced R users “know” functions without having to check other resources. After we complete this basic introduction, Chapter 2 begins doing more sophisticated things with R, allowing us to compare quantitative responses from two groups, make some graphical displays, do hypothesis testing and create confidence intervals in a couple of different ways.

You will have two downloading activities to complete before you can do anything more than read this book<sup>3</sup>. First, you need to download R. It is the engine that will do all the computing for us, but you will only interact with it once. Go to <http://cran.rstudio.com> and click on the “**Download R for...**” button that corresponds to your operating system. On the next page, click on “**base**” and then it will take you to a

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<sup>3</sup>I recorded a video that walks through the material on the following pages that is available here: [https://camtasia.msu.montana.edu/Relay/Files/w76c139/RandRstudio\\_Final/RandRstudio\\_Final\\_-\\_20160715\\_130555\\_23.html](https://camtasia.msu.montana.edu/Relay/Files/w76c139/RandRstudio_Final/RandRstudio_Final_-_20160715_130555_23.html) in the digital version of the book.

screen to download the most current version of R that is compiled for your operating system, something like “**Download R 3.3.1 for Windows**”. Click on that link and then open the file you downloaded. You will need to select your preferred language (choose English so your instructor can help you), then hit “**Next**” until it starts to unpack and install the program (all the base settings will be fine). After you hit “**Finish**” you will not do anything further with R directly.

Second, you need to download RStudio. It is an enhanced interface that will make interacting with R less frustrating. To download RStudio, go to <http://www.rstudio.com/products/rstudio/download/> and select the correct version under “Installers for Supported Platforms” for your operating system. Download and then install RStudio using the installer. From this point forward, you should only open RStudio; it provides your interface with R. Note that both R and RStudio are updated frequently (up to four times a year) and if you downloaded either more than a few months previously, you should download the up-to-date versions, especially if something you are trying to do is not working. Sometimes code will not work in older versions of R and sometimes old code won’t work in new versions of R.<sup>4</sup>

To get started, we can complete some basic tasks in R using the RStudio interface. When you open RStudio, you will see a screen like Figure 1.2. The added annotation in this and the following screen-grabs is there to help you get initially oriented to the software interface. R is command-line software – meaning that most of the time you have to create code and then enter and execute it at a command prompt to get any results. RStudio makes the management and execution of that code more efficient than the basic version of R. In RStudio, the lower left panel is called the “console” window and is where you can type R code directly into R or where you will see the code you run and (most importantly!) where the results of your executed commands will show up. The most basic interaction with R is available once you get the cursor active at the command prompt “>” by clicking in that panel (look for a blinking vertical line). The upper left panel is for writing, saving, and running your R code. Once you have code available in this window, the “Run” button will execute the code for the line that your cursor is on or for any text that you have highlighted with your mouse. The “data management” or environment panel is in the upper right, providing information on what data sets have been loaded. It also contains the “Import Dataset” button that provides the easiest way for you to read a data set into R so you can analyze it. The lower right panel contains information on the “Packages” (additional code we will download and install to add functionality to R) that are available and is where you will see plots that you make and requests for “Help” on specific functions.

As a first interaction with R we can use it as a calculator. To do this, click near the command prompt (>) in the lower left “console” panel, type 3+4, and then hit enter. It should look like this:

```
> 3+4
[1] 7
```

You can do more interesting calculations, like finding the mean of the numbers 3, 5, 7, and 8 by adding them up and dividing by 4:

```
> (-3+5+7+8)/4
[1] 4.25
```

Note that the parentheses help R to figure out your desired order of operations. If you drop that grouping, you get a very different (and wrong!) result:

```
> -3+5+7+8/4
[1] 11
```

We could estimate the standard deviation similarly using the formula you might remember from introductory statistics, but that will only work in very limited situations. To use the real power of R this semester, we need to work with data sets that store the Basically, we need to store observations in named vectors (one dimensional arrays) that contain a list of the observations. To create a vector containing the four numbers and assign it to a variable named *variable1*, we need to create a vector using the function `c` which means “combine the items” that follow, if they are inside parentheses and have commas separating the values, as follows:

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<sup>4</sup>The need to keep the code up-to-date as R continues to evolve is one reason that this book is locally published and that this is the 3<sup>rd</sup> version in three years...

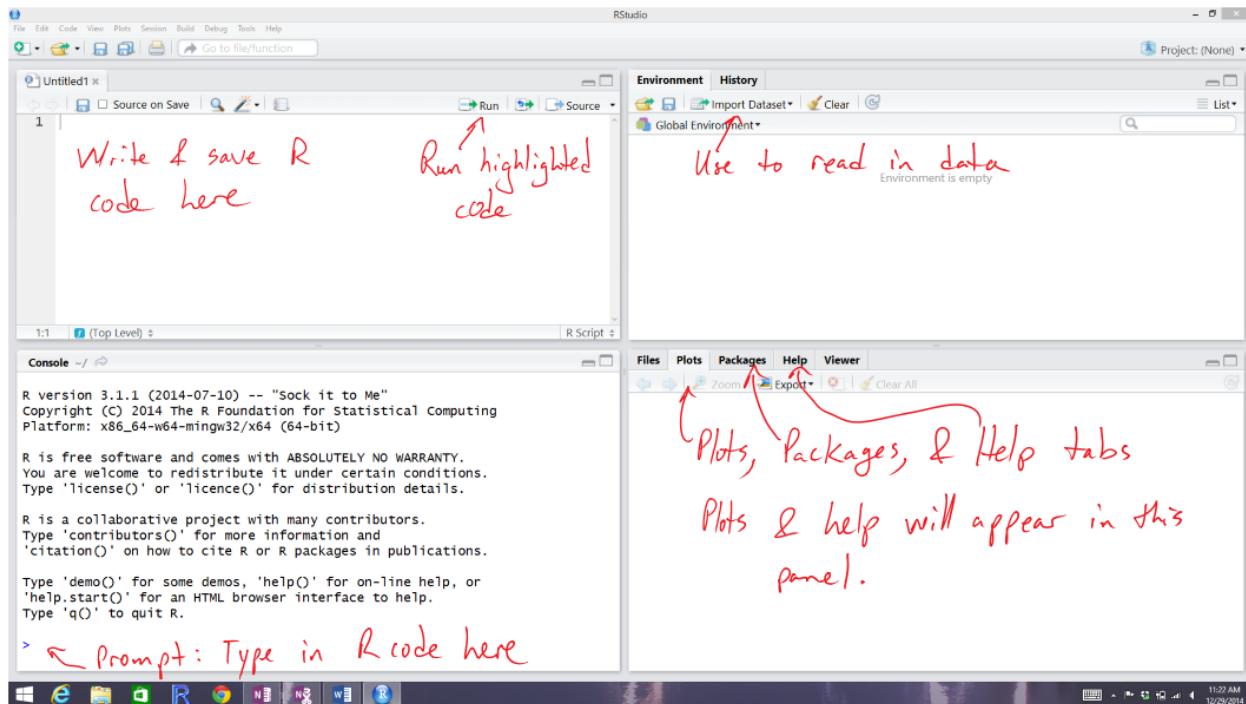


Figure 1.2: Initial RStudio layout

```
> c(-3, 5, 7, 8)
[1] -3 5 7 8
```

To get this vector stored in a variable called *variable1* we need to use the assignment operator, `<-` (read as “stored as”) that assigns the information on the right into the variable that you are creating on the left.

```
> variable1 <- c(-3, 5, 7, 8)
```

In R, the assignment operator, `<-`, is created by typing a “less than” symbol `<` followed by a “minus” sign `(-)` ever want to see what numbers are residing in an object in R, just type its name and hit *enter*. You can see how that variable contains the same information that was initially generated by `c(-3, 5, 7, 8)` but is easier to access since we just need the text for the variable name representing that vector.

```
> variable1
[1] -3 5 7 8
```

With the data stored in a variable, we can use functions such as `mean` and `sd` to find the mean and standard deviation of the observations contained in *variable1*:

```
> mean(variable1)
[1] 4.25
> sd(variable1)
[1] 4.99166
```

When dealing with real data, we will often have information about more than one variable. We could enter all observations by hand for each variable but this is prone to error and onerous for all but the smallest data sets. If you are to ever utilize the power of statistics in the evolving data-centered world, data management has to be accomplished in a more sophisticated way. While you can manage data sets quite effectively in R, it is often easiest to start with your data set in something like Microsoft Excel or OpenOffice’s Calc. You want to make sure that observations are in the rows and the names of variables are in the columns and that there is no “extra stuff” in the spreadsheet. If you have missing observations, they should be represented with blank cells. The file should be saved as a “.csv” file (stands for comma-separated values although Excel calls it “CSV (Comma Delimited)”, which basically strips off some of the junk that Excel adds to the necessary

information in the file. Excel will tell you that this is a bad idea, but it actually creates a more stable archival format and one that R can use directly<sup>5</sup>.

With data set converted to a CSV file, we need to read the data set into R. There are two ways to do this, either using the point-and-click GUI in RStudio (click the “Import Data Set” button in the upper right “Environment” panel as indicated in Figure 1.2) or modifying the

`read.csv` function to find the file of interest. To practice this, you can download an Excel (.xls) file from <http://www.math.montana.edu/courses/s217/documents/treadmill.xls> 31 males that volunteered for a study on methods for measuring fitness (Westfall and Young, 1993). In the spreadsheet, you will find a data set that starts and ends with the following information (only results for Subjects 1, 2, 30, and 31 shown here):

Sub- ject	Tread- MillOx	TreadMill- MaxPulse	RunTime	RunPulse	Rest Pulse	BodyWeight	Age
1	60.05	186	8.63	170	48	81.87	38
2	59.57	172	8.17	166	40	68.15	42
...	...	...	...	...	...	...	...
30	39.2	172	12.88	168	44	91.63	54
31	37.39	192	14.03	186	56	87.66	45

The variables contain information on the subject number (*Subject*), subjects’ treadmill oxygen consumption (*TreadMillOx*, in ml per kg per minute) and maximum pulse rate (*TreadMillMaxPulse*, in beats per minute), time to run 1.5 miles (*Run Time*, in minutes), maximum pulse during 1.5 mile run (*RunPulse*, in beats per minute), resting pulse rate (*RestPulse*, beats per minute), Body Weight (*BodyWeight*, in kg), and *Age* (in years). Open the file in Excel or equivalent software and then save it as a .csv file in a location you can find on your computer. Then go to RStudio and click on **File**, then **Import Dataset**, then **From CSV...**<sup>6</sup> Find your file and check “**Import**”. R will store the data set as an object named whatever the .csv file was named. You could use another name as well, but it is often easiest just to keep the data set name in R related to the original file name. You should see some text appear in the console (lower left panel) like in Figure 1.3. The text that is created will look something like the following – if you had stored the file in a drive labeled D:, it would be:

```
treadmill <- read.csv("D:/treadmill.csv")
```

What is put inside the “ ” will depend on the location and name of your saved .csv file. A version of the data set in what looks like a spreadsheet will appear in the upper left window due to the second line of code (`View(treadmill)`).

Just directly typing (or using) a line of code like this is actually the other way that we can read in files. If you choose to use the text-only interface, then you need to tell R where to look in your computer to find the data file. `read.csv` is a function that takes a path as an argument. To use it, specify the path to your data file, put quotes around it, and put it as the input to `read.csv(...)`. For some examples later in the book, you will be able to copy a command like this from the text and read data sets and other code directly from my the course folder, assuming you are connected to the internet.

To verify that you read the data set in correctly, it is always good to check its contents. We can view the first and last rows in the data set using the `head` and `tail` functions on the data set, which show the following results for the `treadmill` data. Note that you will sometimes need to resize the console window in RStudio to get all the columns to display in a single row which can be performed by dragging the gray bars that separate the panels.

```
head(treadmill)
```

<sup>5</sup>There are ways to read “.xls” and “.xlsx” files directly into R but to handle multiple sheets they are more complicated and not as stable across operating systems as the simpler version we recommend.

<sup>6</sup>If you are having trouble getting the file converted and read into R, copy and run the following code: `treadmill <- read.csv("http://www.math.montana.edu/courses/s217/documents/treadmill.csv", header=T)`.

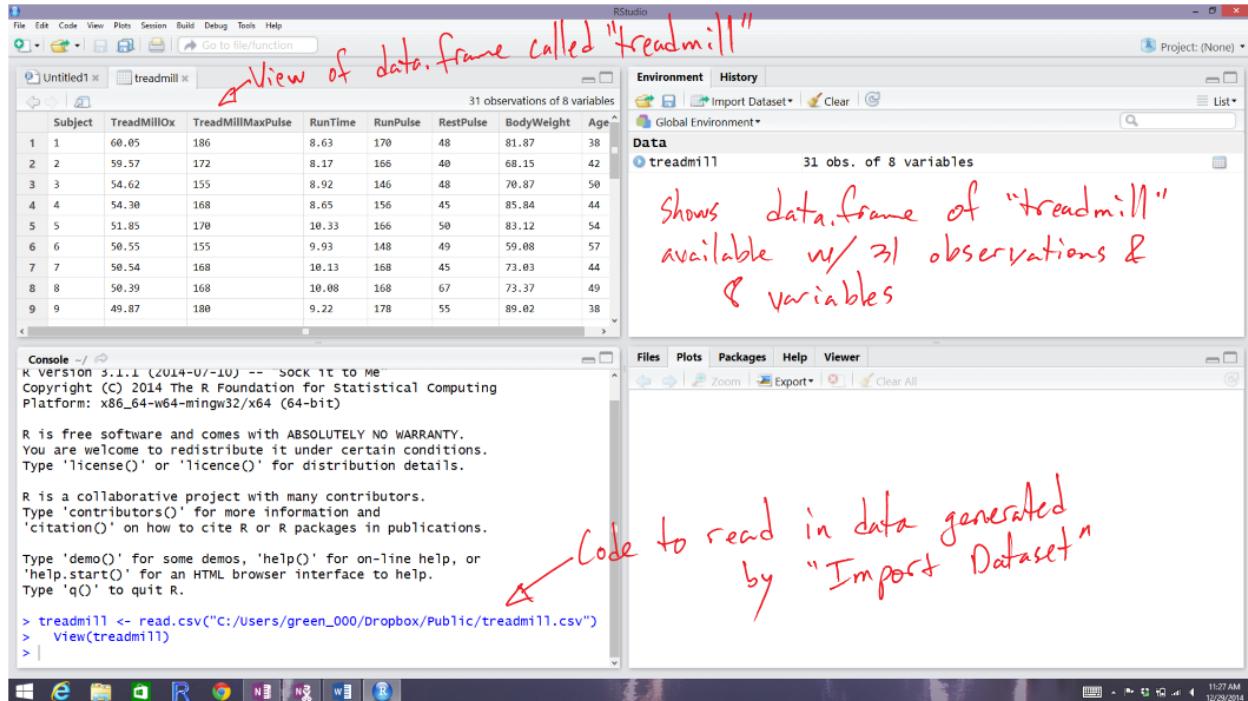


Figure 1.3: RStudio with initial data set loaded

```

##   Subject TreadMillOx TreadMillMaxPulse RunTime RunPulse RestPulse
## 1       1      60.05           186     8.63      170       48
## 2       2      59.57           172     8.17      166       40
## 3       3      54.62           155     8.92      146       48
## 4       4      54.30           168     8.65      156       45
## 5       5      51.85           170    10.33      166       50
## 6       6      50.55           155     9.93      148       49
##   BodyWeight Age
## 1     81.87 38
## 2     68.15 42
## 3     70.87 50
## 4     85.84 44
## 5     83.12 54
## 6     59.08 57
tail(treadmill)

##   Subject TreadMillOx TreadMillMaxPulse RunTime RunPulse RestPulse
## 26      26      44.61           182    11.37      178       62
## 27      27      40.84           172    10.95      168       57
## 28      28      39.44           176    13.08      174       63
## 29      29      39.41           176    12.63      174       58
## 30      30      39.20           172    12.88      168       44
## 31      31      37.39           192    14.03      186       56
##   BodyWeight Age
## 26     89.47 44
## 27     69.63 51
## 28     81.42 44
## 29     73.37 57
## 30     91.63 54
## 31     87.66 45

```

While not always required, for many of the analyses, we will tap into a large suite of additional functions available in R packages by “installing” (basically downloading) and then “loading” the packages. There are some packages that we will use frequently, starting with the `mosaic` package (Pruim et al., 2016b). To install an R package, go to the **Packages** tab in the lower right panel of RStudio. Click on the **Install** button and then type in the name of the package in the box (here type in `mosaic`). RStudio will try to auto-complete the package name you are typing which should help you make sure you got it typed correctly. This will be the first of *many* times that we will mention that R is case sensitive – in other words, `Mosaic` is different from `mosaic` in R syntax and this sort of thing applies to everything you do in R. You should only need to install each R package once on a given computer. If you ever see a message that R can’t find a package, make sure it appears in the list in the **Packages** tab and if it doesn’t, repeat the previous steps to install it.

After installing the package, we need to load it to make it active in a given work session. Go to the command prompt and type (or copy and paste) `require(mosaic)` :

```
require(mosaic)
```

You may see a warning message about versions of the package and versions of R – this is *usually* something you can ignore. Other warning messages could be more ominous for proceeding but before getting too concerned, there are couple of basic things to check. First, double check that the package is installed (see previous steps). Second, check for typographical errors in your code – especially for mis-spellings or unintended capitalization. If you are still having issues, try repeating the installation process. If that fails, find someone more used to using R to help you (for example in the Math Learning Center or by emailing your instructor)<sup>7</sup>.

To help you go from basic to intermediate R usage and especially to help with more complicated problems, you will want to learn how to manage and save your R code. The best way to do this is using the upper left panel in RStudio using what are called R Scripts, which are files that have a file extension of “.R”. To start a new “.R” file to store your code, click on **File**, then **New File**, then **R Script**. This will create a blank page to enter and edit code – then save the file as MyFileName.R in your preferred location. Saving your code will mean that you can return to where you last were working by simply re-running the saved script file. With code in the script window, you can place the cursor on a line of code or highlight a chunk of code and hit the “Run” button on the upper part of the panel. It will appear in the console with results just like what you would obtain if you typed it after the command prompt and hit enter for each line.

Figure 1.4 shows the screen with the code used in this section in the upper left panel, saved in file called Ch0.R, with the results of highlighting and executing the first section of code using the “Run” button.

### 1.3 Basic summary statistics, histograms, and boxplots using R

With RStudio running, the `mosaic` package loaded, a place to write and save code, and the `treadmill` data set loaded, we can (finally!) start to summarize the results of the study. The `treadmill` object is what R calls a **data.frame**<sup>8</sup> and contains columns corresponding to each variable in the spreadsheet. Every function in R will involve specifying the variable(s) of interest and how you want to use them. To access a particular variable (column) in a data.frame, you can use a \$ between the data.frame name and the name of the variable of interest, generically as `dataframename$variablename`. To identify the `RunTime` variable here it would be `treadmill$RunTime`. In the command line it would look like:

```
treadmill$RunTime
```

```
## [1] 8.63 8.17 8.92 8.65 10.33 9.93 10.13 10.08 9.22 8.95 10.85
## [12] 9.40 11.50 10.50 10.60 10.25 10.00 11.17 10.47 11.95 9.63 10.07
## [23] 11.08 11.63 11.12 11.37 10.95 13.08 12.63 12.88 14.03
```

---

<sup>7</sup>Most computer lab computers at Montana State University have RStudio installed and so provide another venue to try this where the software is already installed.

<sup>8</sup>Data frames in R are objects that can contain both categorical and quantitative variables on your n subjects with a name for each variable that is also the name of each column in a matrix. Each subject is a row of the data set.

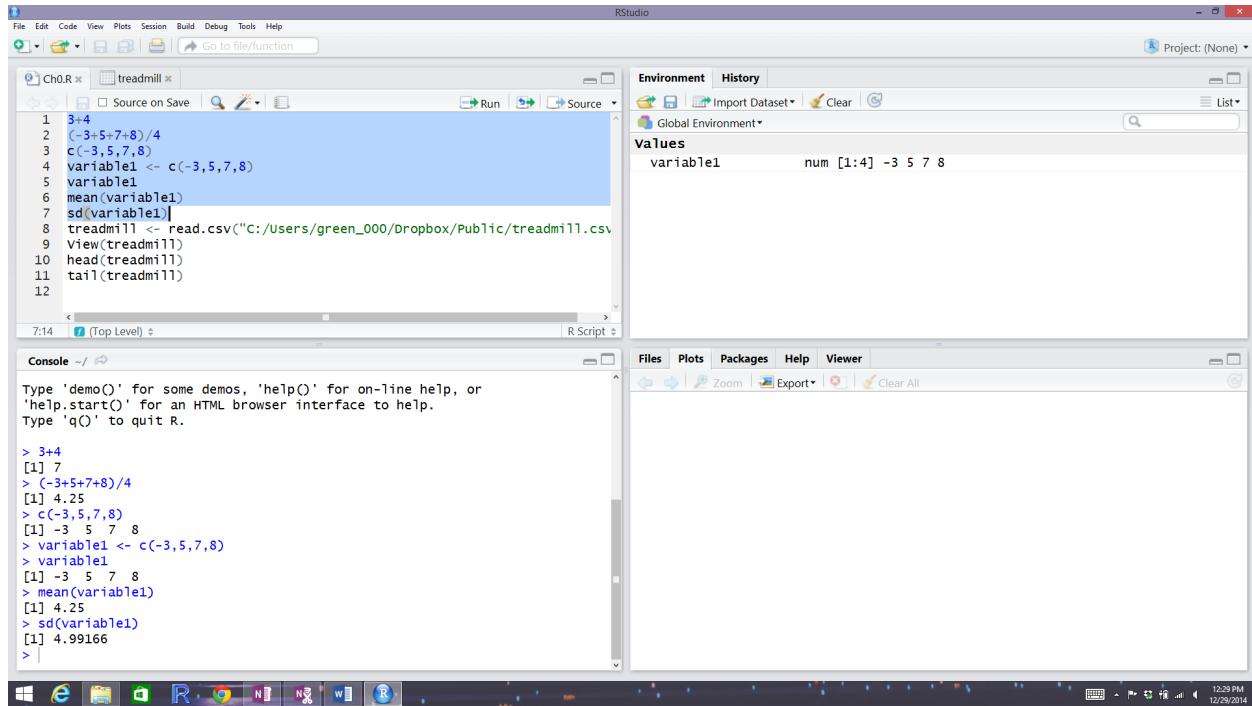


Figure 1.4: RStudio with highlighted code run

Just as in the previous section, we can generate summary statistics using functions like `mean` and `sd` by running them on a specific variable:

```
mean(treadmill$RunTime)
```

```
## [1] 10.58613
sd(treadmill$RunTime)
```

```
## [1] 1.387414
```

And now we know that the average running time for 1.5 miles for the subjects in the study was 10.6 minutes with a standard deviation (SD) of 1.39 minutes. But you should remember that the mean and SD are only appropriate summaries if the distribution is roughly *symmetric* (both sides of the distribution are approximately the same). The `mosaic` package provides a useful function called `favstats` that provides the mean and SD as well as the **5 number summary**: the minimum (`min`), the first quartile (`Q1`, the 25<sup>th</sup> percentile), the median (50<sup>th</sup> percentile), the third quartile (`Q3`, the 75<sup>th</sup> percentile), and the maximum (`max`). It also provides the number of observations (`n`) which was 31, as noted above, and a count of whether any missing values were encountered (`missing`), which was 0 here since all subjects had measurements available on this variable.

```
favstats(treadmill$RunTime)
```

```
##   min    Q1 median    Q3   max     mean      sd n missing
##  8.17  9.78  10.47 11.27 14.03 10.58613 1.387414 31       0
```

We are starting to get somewhere with understanding that the runners were somewhat fit with worst runner covering 1.5 miles in 14 minutes (the equivalent of a 9.3 minute mile) and the best running at a 5.4 minute mile pace. The limited variation in the results suggests that the sample was obtained from a restricted group with somewhat common characteristics. When you explore the ages and weights of the subjects in the Practice Problems in Section 0.5, you will get even more information about how similar all the subjects in this study were.

A graphical display of these results will help us to assess the shape of the distribution of run times – including

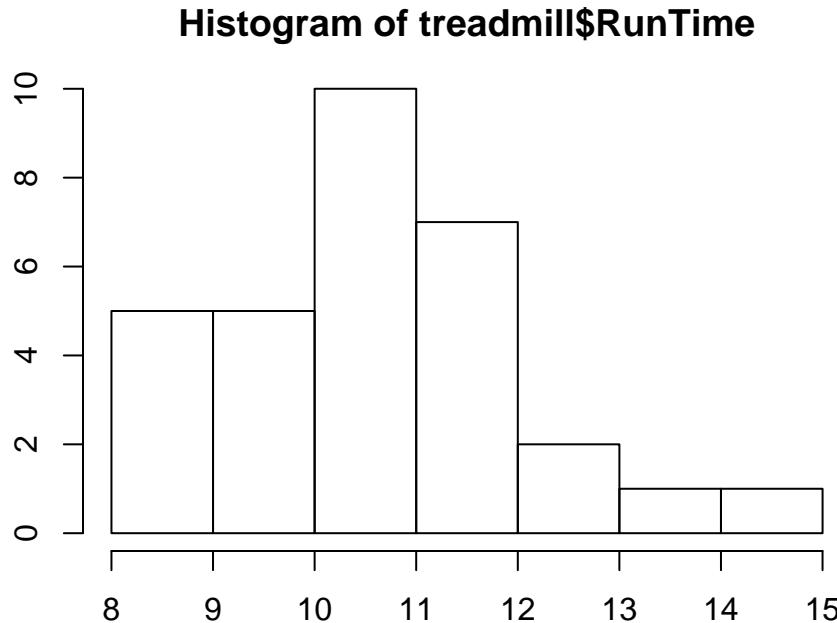


Figure 1.5: Histogram of Run Times #(minutes) of n=31 subjects in Treadmill study.

considering the potential for the presence of a *skew* (whether the right or left tail of the distribution is noticeably more spread out with left skew meaning that the left tail is more spread out than the right tail) and *outliers* (unusual observations). A *histogram* is a good place to start. Histograms display connected bars with counts of observations defining the height of bars based on a set of bins of values of the quantitative variable. We will apply the `hist` function to the `RunTime` variable, which produces Figure 1.5.

I used the **Export** button found above the plot, followed by **Copy to Clipboard** and clicking on the **Copy Plot** button. Then if you open your favorite word-processing program, you should be able to paste it into a document for writing reports that include the figures. You can see the first parts of this process in the screen grab in Figure 1.6.

You can also directly save the figures as separate files using **Save as Image** or **Save as PDF** and then insert them into your word processing documents.

The function `hist` defaults into providing a histogram on the *frequency* (count) scale. In most R functions, there are the default options that will occur if we don't make any specific choices but we can override the default options if we desire. One option we can modify here is to add labels to the bars to be able to see exactly how many observations fell into each bar. Specifically, we can turn the `labels` option "on" by making it true ("T") by adding `labels=T` to the previous call to the `hist` function, separated by a comma:

```
hist(treadmill$RunTime, labels=T)
```

Based on this histogram, it does not appear that there are any outliers in the responses since there are no bars that are separated from the other observations. However, the distribution does not look symmetric and there might be a skew to the distribution. Specifically, it appears to be *skewed right* (the right tail is longer than the left). But histograms can sometimes mask features of the data set by binning observations and it is hard to find the percentiles accurately from the plot.

When assessing outliers and skew, the *boxplot* (or *Box and Whiskers* plot) can also be helpful (Figure 1.8) to describe the shape of the distribution as it displays the 5-number summary and will also indicate observations that are "far" above the middle of the observations. R's `boxplot` function uses the standard rule to indicate an observation as a *potential outlier* if it falls more than 1.5 times the *IQR* (Inter-Quartile Range, calculated as Q3-Q1) below Q1 or above Q3. The potential outliers are plotted with circles and the

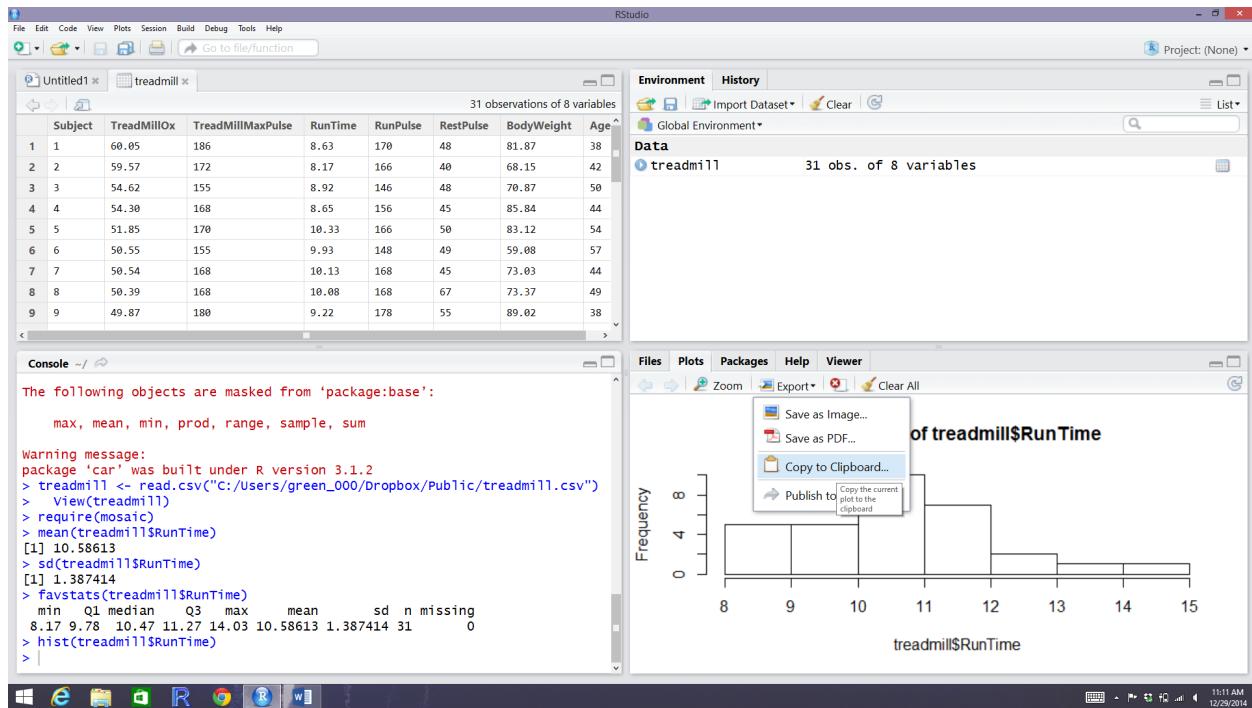


Figure 1.6: RStudio while in the process of copying the histogram

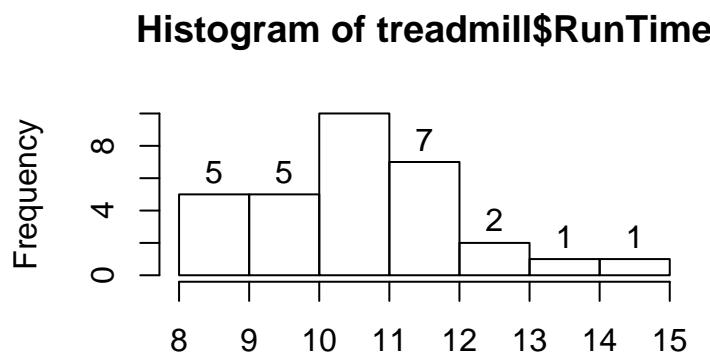


Figure 1.7: Histogram of #Run Times with counts in bars labeled.

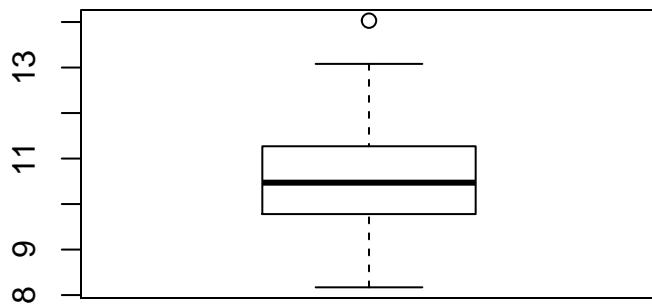


Figure 1.8: Boxplot of 1.5 mile Run Times.

*Whiskers* (lines that extend from Q1 and Q3 typically to the minimum and maximum) are shortened to only go as far as observations that are within  $1.5 \times \text{IQR}$  of the upper and lower quartiles. The *box* part of the boxplot is a box that goes from Q1 to Q3 and the median is displayed as a line somewhere inside the box<sup>9</sup>. Looking back at the summary statistics above,  $\text{Q1}=9.78$  and  $\text{Q3}=11.27$ , providing an IQR of:

```
IQR <- 11.27 - 9.78
```

One observation (the maximum value of 14.03) is indicated as a potential outlier based on this result by being larger than  $\text{Q3} + 1.5 \times \text{IQR}$ , which was 13.505:

```
11.27 + 1.5*IQR
```

```
## [1] 13.505
```

The boxplot also shows a slight indication of a right skew (skew towards larger values) with the distance from the minimum to the median being smaller than the distance from the median to the maximum. Additionally, the distance from Q1 to the median is smaller than the distance from the median to Q3. It is modest skew, but worth noting.

```
boxplot(treadmill$RunTime)
```

While the default boxplot is fine, it fails to provide good graphical labels, especially on the y-axis. Additionally, there is no title on the plot. The following code provides some enhancements to the plot by using the `ylab` and `main` options in the call to `boxplot`, with the results displayed in Figure 1.9. When we add text to plots, it will be contained within quotes and be assigned into the options `ylab` (for y-axis) or `main` (for the title) here to put it into those locations.

```
boxplot(treadmill$RunTime, ylab="1.5 Mile Run Time (minutes)",  
       main="Boxplot of the Run Times of n=31 participants")
```

Throughout the book, we will often use extra options to make figures that are easier for you to understand. There are often simpler versions of the functions that will suffice but the extra work to get better labeled figures is often worth it. I guess the point is that “a picture is worth a thousand words” but in data visualization, that is only true if the reader can understand what is being displayed. It is also important to think about the quality of the information that is being displayed, regardless of how pretty the graphic might be.

All the previous results were created by running the R code and then “grabbing” the results from either the console or by copying the figure. There is another way to use RStudio where you can have it compile the results (both output and figures) directly into a document together with the code that generated it, using

---

<sup>9</sup>The median, quartiles and whiskers sometimes occur at the same values when there are many tied observations. If you can't see all the components of the boxplot, produce the numerical summary to help you understand what happened.

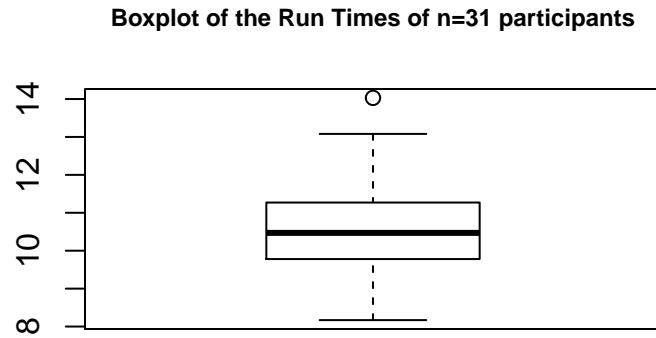


Figure 1.9: Boxplot of Run Times with improved labels.

what is called RMarkdown (<http://shiny.rstudio.com/articles/rmarkdown.html>). It adds some additional setup complexity we want to avoid for now but is what we used to do all the analyses that follow in the book. The main reason to mention this is that you will see a change in formatting of the R code and output from here forward as you will no longer see the command prompt (“>”) with the code. The output will be flagged by having two “##”’s before it. For example, the summary statistics for the *RunTime* variable from ‘favstats function would look like:

```
favstats(treadmill$RunTime)

##   min   Q1 median   Q3   max   mean      sd n missing
##  8.17 9.78 10.47 11.27 14.03 10.58613 1.387414 31       0
```

Statisticians (and other scientists) are starting to use these methods because they provide what is called “Reproducible research” (Gandrud, 2015) where all the code and output it produced are available in a single place. This allows different researchers to run and verify results or the original researchers to revisit their earlier work at a later date and recreate all their results. Scientific publications are currently encouraging researchers to work in this way and may someday require it. In this book, we focus on the R code and show the results from running it, but you may want to consider exploring these alternative options.

Finally, when you are done with your work and attempt to exit out of RStudio, it will ask you to save your workspace. You do not need to do this and would be better served not to do this. If you are in the practice of saving your workspace, you will end up with tons of data.frames that open each time you use it and it will be harder to find and manage the ones you are currently working with. If you save your R code via the script window, you can re-create any results by simply re-running that code. If you find that you have lots of “stuff” in your workspace, just run `rm(list = ls())`. It will delete all the data sets from your workspace.

## 1.4 Chapter summary

This chapter covered getting R and RStudio downloaded and some basics of working with R via RStudio. You should be able to read a data set into R and run some basic functions, all done using the RStudio interface. If you are struggling with this, you should seek additional help with these technical issues so that you are ready for more complicated statistical methods that are going to be encountered in the following chapters. For most assignments, we will give you a seed of the basic R code that you need and then you will modify it to work on your data set of interest. As mentioned previously, the way everyone learns R is by starting with some example code that does most of what you want to do and then you modify it. If you can complete the Practice Problems that follow, you are well on your way to learning to use R.

The statistical methods in this chapter were minimal and all should have been review. They involved a quick reminder of summarizing the center, spread, and shape of distributions using numerical summaries of the mean and SD and/or the min, Q1, median, Q3, and max and the histogram and boxplot as graphical summaries. We revisited the ideas of symmetry and skew. But the main point was really to get a start on using R to provide results you should be familiar with from your previous statistics experience(s).

## 1.5 Important R Code

To help you learn and use R, there is a section highlighting the most important R code used near the end of each chapter. The dark text will never change but the lighter (red) text will need to be customized to your particular application. The sub-bullet for each function will discuss the use of the function and pertinent options or packages required. You can use this as a guide to finding the function names and some hints about options that will help you to get the code to work or you can revisit the worked examples using each of the functions.

- `FILENAME<- read.csv("path to csv file/FILENAME.csv")`
  - Can be generated using “Import Dataset” button or by modifying this text.
- `DATASETNAME$VARIABLENAME`
  - To access a particular variable in a data.frame called DATASETNAME, use a \$ and then the VARIABLENAME.
- `head(DATASETNAME)`
  - Provides a list of the first few rows of the data set for all the variables in it.
- `mean(DATASETNAME$VARIABLENAME)`
  - Calculates the mean of the observations in a variable.
- `sd(DATASETNAME$VARIABLENAME)`
  - Calculates the SD of the observations in a variable.
- `favstats(DATASETNAME$VARIABLENAME)`
  - Requires the `mosaic` package to be loaded (`require(mosaic)` after installing the package).
  - Provides a suite of numerical summaries of the observations in a variable.
- `hist(DATASETNAME$VARIABLENAME)`
  - Makes a histogram.
- `boxplot(DATASETNAME$VARIABLENAME)`
  - Makes a boxplot.

## 1.6 Practice problems

In each chapter, the last section contains some questions for you to complete to make sure you understood the material. You can download the code to answer questions 0.1 to 0.5 below at <http://www.math.montana.edu/courses/s217/documents/Ch0.Rmd>. But to practice learning R, it would be most useful for you to try to accomplish the requested tasks yourself and then only refer to the provided R code if/when you struggle. These questions provide a great venue to check your learning, often to see the methods applied to another data set, and for something to discuss in study groups, with your instructor, and/or at the Math Learning Center.

1.1. Read in the treadmill data set discussed above and find the mean and SD of the Ages (*Age* variable) and Body Weights (*BodyWeight* variable). In studies involving human subjects, it is common to report a summary of characteristics of the subjects. Why does this matter? Think about how your interpretation of any study of the fitness of subjects would change if the mean age had been 20 years older or 35 years younger.

1.2. How does knowing about the distribution of results for *Age* and *BodyWeight* help you understand the results for the Run Times discussed above?

1.3. The mean and SD are most useful as summary statistics only if the distribution is relatively symmetric. Make a discuss the shape of the distribution (is it skewed right, skewed left, approximately symmetric?; are there outliers?). Approximately what range of ages does this study pertain to?

1.4. The weight responses are in kilograms and you might prefer to see them in pounds. The conversion is  $\text{lbs}=2.205\text{kgs}$ . Create a new variable in the *treadmill* data.frame called *BWlb*\* using this code:

```
treadmill$BWlb <- 2.205*treadmill$BodyWeight
```

and find the mean and SD of the new variable (*BWlb*).

1.5. Make histograms and boxplots of the original *BodyWeight* and new *BWlb* variables. Discuss aspects of the distributions that changed and those that remained the same with the transformation from kilograms to pounds.

# Chapter 2

## (R)e-Introduction to statistics

The previous material served to get us started in R and to get a quick review of some basic descriptive statistics. Now we will begin to engage some new material and exploit the power of R to do some statistical inference. Because inference is one of the hardest topics to master in statistics, we will also review some basic terminology that is required to move forward in learning more sophisticated statistical methods. To keep this “review” as short as possible, we will not consider every situation you learned in introductory statistics and instead focus exclusively on the situation where we have a quantitative response variable measured on two groups, adding a new graphic called a “bean plot” to help us see the differences in the observations in the groups.

### 2.1 Histograms, boxplots, and density curves

Part of learning statistics is learning to correctly use the terminology, some of which is used colloquially differently than it is used in formal statistical settings. The most commonly “misused” term is **data**. In statistical parlance, we want to note the plurality of data. Specifically, **datum** is a single measurement, possibly on multiple random variables, and so it is appropriate to say that “**a datum is...**”. Once we move to discussing data, we are now referring to more than one observation, again on one, or possibly more than one, random variable, and so we need to use “**data are...**” when talking about our observations. We want to distinguish our use of the term “data” from its more colloquial<sup>1</sup> usage that often involves treating it as singular. In a statistical setting “data” refers to measurements of our cases or units. When we summarize the results of a study (say providing the mean and SD), that information is not “data”. We used our data to generate that information. Sometimes we also use the term “data set” to refer to all our observations and this is a singular term to refer to the group of observations and this makes it really easy to make mistakes on the usage of this term.

It is also really important to note that **variables** have to vary – if you measure the sex of your subjects but are only measuring females, then you do not have a “variable”. You may not know if you have real variability in a “variable” until you explore the results you obtained.

The last, but probably most important, aspect of data is the context of the measurement. The “who, what, when, and where” of the collection of the observations is critical to the sort of conclusions we can make based on the results. The information on the study design provides information required to assess the scope of inference of the study. Generally, remember to think about the research questions the researchers were trying to answer and whether their study actually would answer those questions. There are no formulas to help us sort some of these things out, just critical thinking about the context of the measurements.

---

<sup>1</sup>You will more typically hear “data is” but that more often refers to information, sometimes even statistical summaries of data sets, than to observations collected as part of a study, suggesting the confusion of this term in the general public. We will explore a data set in Chapter 4 related to perceptions of this issue collected by researchers at <http://fivethirtyeight.com/>.

To make this concrete, consider the data collected from a study (Plaster, 1989) to investigate whether perceived physical attractiveness had an impact on the sentences or perceived seriousness of a crime that male jurors might give to female defendants. The researchers showed the participants in the study (men who volunteered from a prison) pictures of one of three young women. Each picture had previously been decided to be either beautiful, average, or unattractive by the researchers. Each “juror” was randomly assigned to one of three levels of this factor (which is a categorical predictor or explanatory variable) and then each rated their picture on a variety of traits such as how warm or sincere the woman appeared. Finally, they were told the women had committed a crime (also randomly assigned to either be told she committed a burglary or a swindle) and were asked to rate the seriousness of the crime and provide a suggested length of sentence. We will bypass some aspects of their research and just focus on differences in the sentence suggested among the three pictures. To get a sense of these data, let’s consider the first and last parts of the data set:

Table 2.1: First 5 and last 6 rows of the Mock Jury data set.

Subject	Attr	Crime	Years	Serious	Independent	Sincere
1	Beautiful	Burglary	10	8	9	8
2	Beautiful	Burglary	3	8	9	3
3	Beautiful	Burglary	5	5	6	3
4	Beautiful	Burglary	1	3	9	8
5	Beautiful	Burglary	7	9	5	1
...	...	...	...	...	...	...
108	Average	Swindle	3	3	5	4
109	Average	Swindle	3	2	9	9
110	Average	Swindle	2	1	8	8
111	Average	Swindle	7	4	9	1
112	Average	Swindle	6	3	5	2
113	Average	Swindle	12	9	9	1
114	Average	Swindle	8	8	1	5

When working with data, we should always start with summarizing the sample size. We will use  $n$  for the number of subjects in the sample and denote the population size (if available) with  $N$ . Here, the sample size is  $n=114$ . In this situation, we do not have a random sample from a population (these were volunteers from the population of prisoners at the particular prison) so we cannot make inferences to a larger group. But we can assess whether there is a *causal effect*<sup>2</sup>: if sufficient evidence is found to conclude that there is some difference in the responses across the treated groups, we can attribute those differences to the treatments applied, since the groups should be same otherwise due to the pictures being randomly assigned to the “jurors”. The story of the data set – that it was collected on prisoners – becomes pretty important in thinking about the ramifications of any results. Are male prisoners different from the population of college males or all residents of a state such as Montana? If so, then we should not assume that the detected differences, if detected, would also exist in some other group of male subjects. The lack of a random sample makes it impossible to assume that this set of prisoners might be like other prisoners. So there are definite limitations to the inferences in the following results. But it is still interesting to see if the pictures caused a difference in the suggested mean sentences, even though the inferences are limited to this group of prisoners. If this had been an observational study (suppose that the prisoners could select one of the three pictures), then we would have to avoid any of the “causal” language that we can consider here because the pictures were not randomly assigned to the subjects. Without random assignment, the explanatory variable of picture choice could be *confounded* with another characteristic of prisoners that was related to which picture they selected and the rating they provided. Confounding is not the only reason to avoid causal statements with non-random assignment but the inability to separate the effect of other variables (measured or unmeasured) from the differences we are observing means that our inferences in these situations need to be carefully stated.

<sup>2</sup>As noted previously, we reserve the term “effect” for situations where random assignment allows us to consider causality as the reason for the differences in the response variable among levels of the explanatory variable, but this is only the case if we find evidence against the null hypothesis of no difference in the groups.

Instead of loading this data set into R using the “Import Dataset” functionality, we can load an R package that contains the data, making for easy access to this data set. The package called `heplots` (Fox and Friendly, 2016) contains a data set called `MockJury` that contains the results of the study. We also rely the R package called `mosaic` (Pruim et al., 2016b) that was introduced previously. First (but only once), you need to install both packages, which can be done either using the Packages tab in the lower right panel of R-studio or using the `install.packages` function with quotes around the package name:

```
> install.packages("heplots")
```

After making sure that both packages are installed, we use the `require` function around the package name (no quotes now!) to load the package, something that you need to do any time you want to use features of a package.

```
require(heplots)
require(mosaic)
```

There will be some results of the loading process that may discuss loading other required packages. If the output says that it needs a package that is unavailable, then follow the same process noted above to install that package as well.

To load the data set that is available in an active package, we use the `data` function.

```
data(MockJury)
```

Now there will be a `data.frame` called `MockJury` available for us to analyze and some information about it in the Environment tab. Again, we can find out more about the data set in a couple of ways. First, we can use the `View` function to provide a spreadsheet type of display in the upper left panel. Second, we can use the `head` and `tail` functions to print out the beginning and end of the data set. Because there are so many variables, it may wrap around to show all the columns.

```
View(MockJury)
```

```
head(MockJury)
```

```
##      Attr   Crime Years Serious exciting calm independent sincere warm
## 1 Beautiful Burglary    10      8       6     9         9      8     5
## 2 Beautiful Burglary     3      8       9     5         9      3     5
## 3 Beautiful Burglary     5      5       3     4         6      3     6
## 4 Beautiful Burglary     1      3       3     6         9      8     8
## 5 Beautiful Burglary     7      9       1     1         5      1     8
## 6 Beautiful Burglary     7      9       1     5         7      5     8
##   phyattr sociable kind intelligent strong sophisticated happy ownPA
## 1         9      9     9       6     9         9      5     9
## 2         9      9     4       9     5         5      5     7
## 3         7      4     2       4     5         4      5     5
## 4         9      9     9       9     9         9      9     9
## 5         8      9     4       7     9         9      8     7
## 6         8      9     5       8     9         9      9     9
tail(MockJury)
```

```
##      Attr   Crime Years Serious exciting calm independent sincere warm
## 109 Average Swindle     3      2       7     6         9      9     6
## 110 Average Swindle     2      1       8     8         8      8     8
## 111 Average Swindle     7      4       1     6         9      1     1
## 112 Average Swindle     6      3       5     3         5      2     4
## 113 Average Swindle    12      9       1     9         9      1     1
## 114 Average Swindle     8      8       1     9         1      5     1
##   phyattr sociable kind intelligent strong sophisticated happy ownPA
## 109        4      7     6       8     6         5      7     2
## 110        8      9     9       9     9         9      9     6
## 111        1      9     4       1     1         1      1     9
## 112        1      4     9       3     3         9      5     3
## 113        1      9     1       9     9         1      9     1
```

```
## 114      1      9      1      1      9      5      1      1
```

When data sets are loaded from packages, there is often extra documentation available about the data set which can be accessed using the help function. In this case, it will bring up a screen with information about the study and each variable that was measured.

```
help(MockJury)
```

The help function is also useful with functions in R to help you understand options and, at the bottom of the help, see examples of using the function.

With many variables in a data set, it is often useful to get some quick information about all of them; the **summary** function provides useful information whether the variables are categorical or quantitative and notes if any values were missing.

```
summary(MockJury)
```

```
##          Attr        Crime       Years      Serious
## Beautiful :39    Burglary:59   Min.   : 1.000  Min.   :1.000
## Average   :38    Swindle :55    1st Qu.: 2.000  1st Qu.:3.000
## Unattractive:37                         Median : 3.000  Median :5.000
##                                         Mean   : 4.693   Mean   :5.018
##                                         3rd Qu.: 7.000  3rd Qu.:6.750
##                                         Max.   :15.000 Max.   :9.000
## exciting      calm     independent      sincere
## Min.   :1.000  Min.   :1.000  Min.   :1.000  Min.   :1.000
## 1st Qu.:3.000 1st Qu.:4.250  1st Qu.:5.000  1st Qu.:3.000
## Median :5.000  Median :6.500  Median :6.500  Median :5.000
## Mean   :4.658  Mean   :5.982  Mean   :6.132  Mean   :4.789
## 3rd Qu.:6.000 3rd Qu.:8.000  3rd Qu.:8.000  3rd Qu.:7.000
## Max.   :9.000  Max.   :9.000  Max.   :9.000  Max.   :9.000
## warm      phyattr      sociable      kind
## Min.   :1.00  Min.   :1.00  Min.   :1.000  Min.   :1.000
## 1st Qu.:2.00 1st Qu.:2.00  1st Qu.:5.000  1st Qu.:3.000
## Median :5.00  Median :5.00  Median :7.000  Median :5.000
## Mean   :4.57  Mean   :4.93  Mean   :6.132  Mean   :4.728
## 3rd Qu.:7.00 3rd Qu.:8.00  3rd Qu.:8.000  3rd Qu.:7.000
## Max.   :9.00  Max.   :9.00  Max.   :9.000  Max.   :9.000
## intelligent  strong     sophisticated     happy
## Min.   :1.000  Min.   :1.000  Min.   :1.000  Min.   :1.000
## 1st Qu.:4.000 1st Qu.:4.000  1st Qu.:3.250  1st Qu.:3.000
## Median :7.000  Median :6.000  Median :5.000  Median :5.000
## Mean   :6.096  Mean   :5.649  Mean   :5.061  Mean   :5.061
## 3rd Qu.:8.750 3rd Qu.:7.000  3rd Qu.:7.000  3rd Qu.:7.000
## Max.   :9.000  Max.   :9.000  Max.   :9.000  Max.   :9.000
## ownPA
## Min.   :1.000
## 1st Qu.:5.000
## Median :6.000
## Mean   :6.377
## 3rd Qu.:9.000
## Max.   :9.000
```

If we take a few moments to explore the output we can discover some useful aspects of the data set. The output is organized by variable, providing summary information based on the type of variable, either counts by category for categorical variables **Attr** and **Crime** mean for quantitative variables. If present, you would also get a count of missing values that are called “NAs” in R. For the first variable, called **Attr** in the data.frame and that we might find counts of the number of subjects shown each picture: 37/114 viewed the “Unattractive” picture, 38 viewed “Average”, and 39 viewed “Beautiful”. We can also see that suggested prison sentences (data.frame variable **Years**) ranged from 1 year to 15 years with a median of 3 years. It seems that all the other variables except for **Crime** (type of crime that they were told the pictured woman

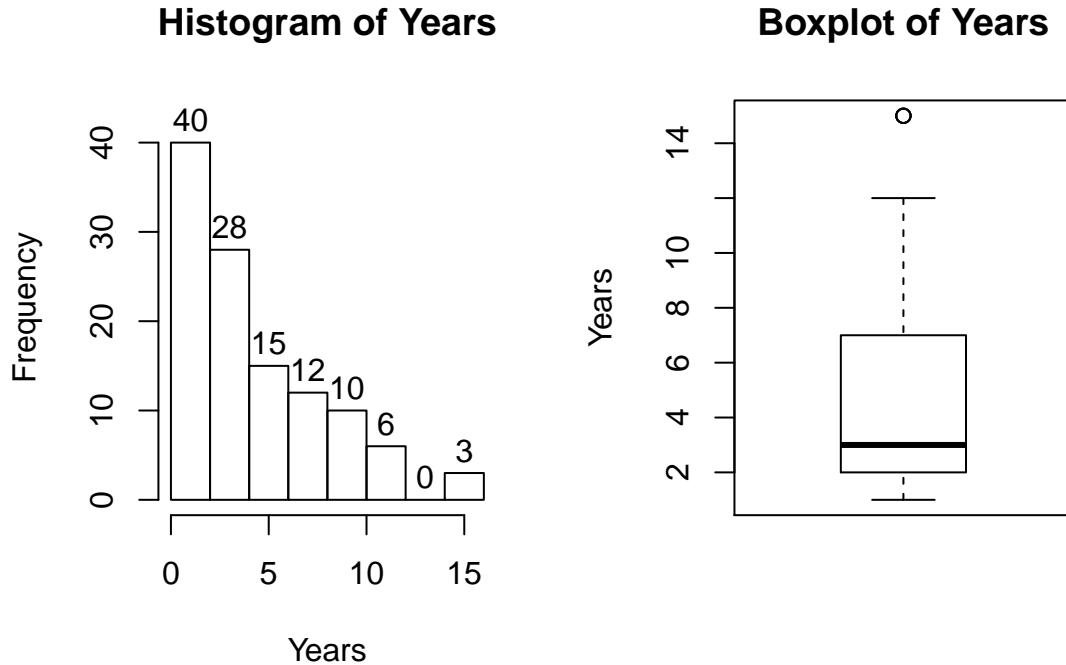


Figure 2.1: Histogram and boxplot of suggested sentences in years.

committed) contained responses between 1 and 9 based on rating scales from 1 = low to 9 = high.

To accompany the numerical summaries, histograms, and boxplots can provide some initial information on the shape of the distribution of the responses for the Figure 2.1 contains the histogram and boxplot of Years, ignoring any information on which picture the “jurors” were shown. The calls to the two plotting functions are enhanced slightly to add better labels.

```
par(mfrow=c(1,2))
hist(MockJury$Years, xlab="Years", labels=T, main="Histogram of Years")
boxplot(MockJury$Years, ylab="Years", main="Boxplot of Years")
```

The distribution appears to have a strong right skew with three observations at 15 years flagged as potential outliers. You can only tell that there are three observations and that they are at 15 by looking at both plots – the bar around 15 years in the histogram has a count of three and the boxplot only shows a single point at 15 which is actually three tied points at exactly 15 years plotted on top of each other (we call this “overplotting”). These three observations really seem to be the upper edge of the overall pattern of a strongly right skewed distribution, so even though they are flagged in the boxplot, we likely would not want to remove them from our data set. In real data sets, outliers are commonly encountered and the first step is to verify that they were not errors in recording. The next step is to study their impact on the statistical analyses performed, potentially considering reporting results with and without the influential observation(s) in the results. If the analysis is unaffected by the “unusual” observations, then it matters little whether they are dropped or not. If they do affect the results, then reporting both versions of results allows the reader to judge the impacts for themselves. It is important to remember that sometimes the outliers are the most interesting part of the data set.

Often when statisticians think of distributions, we think of the smooth underlying shape that led to the data set that is being displayed in the histogram. Instead of binning up observations and making bars in the histogram, we can estimate what is called a **density curve** as a smooth curve that represents the observed distribution of the responses. Density curves can sometimes help us see features of the data sets more clearly.

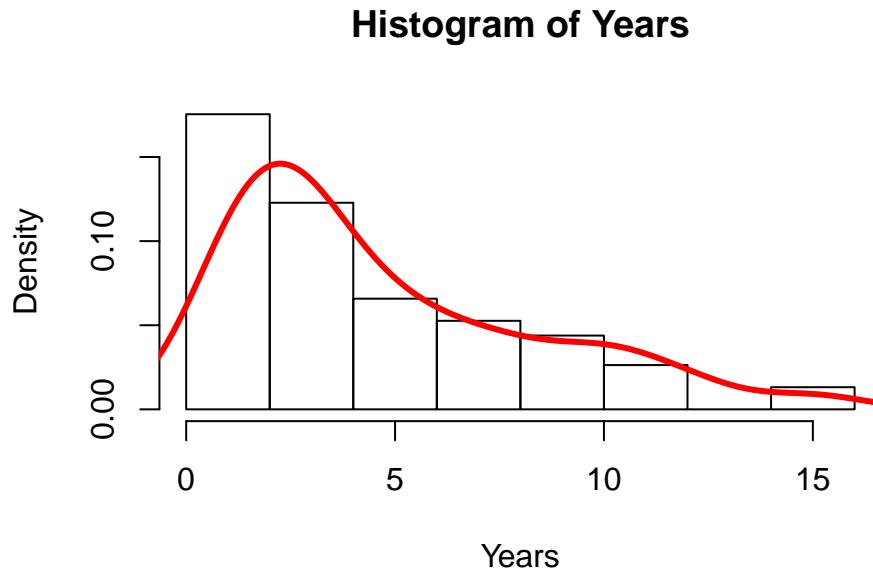


Figure 2.2: Histogram and density curve of Years data.

To understand the density curve, it is useful to initially see the histogram and density curve together. The density curve is scaled so that the total area<sup>3</sup> under the curve is 1. To make a comparable histogram, the y-axis needs to be scaled so that the histogram is also on the “density” scale which makes the bar heights required so that the proportion of the total data set in each bar is represented by the area in each bar (remember that area is height times width). So the height depends on the width of the bars and the total area across all the bars has to be 1. In the `hist` function, the `freq=F` to get density-scaled histogram bars. The density curve is added to the histogram using the R code of `lines(density())`, producing the result in Figure 2.2 with added modifications of options for `lwd` (line width) and `col` (color) to make the plot more interesting. You can see how the density curve somewhat matches the histogram bars but deals with the bumps up and down and edges a little differently. We can pick out the strong right skew using either display and will rarely make both together.

```
hist(MockJury$Years,freq=F,xlab="Years",main="Histogram of Years")
lines(density(MockJury$Years),lwd=3,col="red")
```

Histograms can be sensitive to the choice of the number of bars and even the cut-offs used to define the bins for a given number of bars. Small changes in the definition of cut-offs for the bins can have noticeable impacts on the shapes observed but this does not impact density curves. We are not going to tinker with the default choices for bars in `histogram` as they are reasonably selected, but we can add information on the original observations being included in each bar to better understand the choices that `hist` is making. In the previous display, we can add what is called a ***rug*** to the plot, where a tick mark is made on the x-axis for each observation. Because the responses were provided as whole years (1, 2, 3, ..., 15), we need to use a graphical technique called ***jittering*** to add a little noise<sup>4</sup> to each observation so all the observations at each year value do not plot as a single line. In Figure 2.3, the added tick marks on the x-axis show the approximate locations of the original observations. We can see how there are 3 observations at 15 (all were 15 and the noise added makes it possible to see them all). The limitations of the histogram arise around the 10 year sentence area

<sup>3</sup>If you've taken calculus, you will know that the curve is being constructed so that the integral from  $-\infty$  to  $\infty$  is 1. If you don't know calculus, think of a rectangle with area of 1 based on its height and width. These cover the same area but the top of the region wiggles.

<sup>4</sup>Jittering typically involves adding random variability to each observation that is uniformly distributed in a range determined based on the spacing of the function, the results will change. For more details, type `help(jitter)` in R.

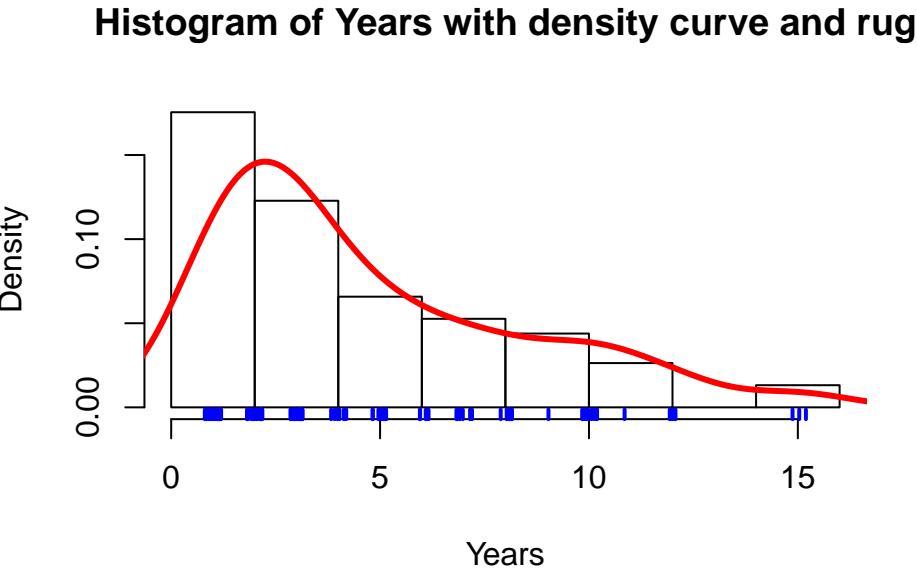


Figure 2.3: Histogram with density curve and rug plot of the jittered responses.

where there are many responses at 10 years and just one at both 9 and 11 years, but the histogram bars sort of miss this aspect of the data set. The density curve did show a small bump at 10 years. Density curves are, however, not perfect and this one shows area for sentences less than 0 years which is not possible here.

```
hist(MockJury$Years, freq=F, xlab="Years",
  main="Histogram of Years with density curve and rug")
lines(density(MockJury$Years), lwd=3, col="red")
rug(jitter(MockJury$Years), col="blue", lwd=2)
```

The graphical tools we've just discussed are going to help us move to comparing the distribution of responses across more than one group. We will have two displays that will help us make these comparisons. The simplest is *the side-by-side boxplot*, where a boxplot is displayed for each group of interest using the same y-axis scaling. In R, we can use its *formula* notation to see if the response (*Years*) differs based on the group (*Attr*) by using something like *Y~X* or, here, *Years~Attr*. We also need to tell R where to find the variables – use the last option in the command, *data=DATASETNAME*, to inform R of the data.frame to look in to find the variables. In this example, *data=MockJury*. We will use the formula and *data=...* options in almost every function we use from here forward. Figure 2.4 contains the side-by-side boxplots showing right skew for all the groups, slightly higher median and more variability for the *Unattractive* group along with some potential outliers indicated in two of the three groups.

```
boxplot(Years~Attr, data=MockJury)
```

The “~” (which is read as the *tilde* symbol, which you can find in the upper left corner of your keyboard) notation will be used in two ways this semester. The formula use in R employed previously declares that the response variable here is *Years* and the explanatory variable is *Attr*. The other use for “~” is as shorthand for “is distributed as” and is used in the context of *Y~N(0,1)*, which translates (in statistics) to defining the random variable *Y* as following a Normal distribution<sup>5</sup> with mean 0 and standard deviation of 1. In the current situation, we could ask whether the *Years* variable seems like it may follow a normal distribution, in other words, is *Years~N(0,1)*? Since the responses are right skewed with some groups having outliers, it is not reasonable to assume that the *Years* variable for any of the three groups may follow a Normal

<sup>5</sup>Remember the bell-shaped curve you encountered in introductory statistics? If not, you can see some at [https://en.wikipedia.org/wiki/Normal\\_distribution](https://en.wikipedia.org/wiki/Normal_distribution)

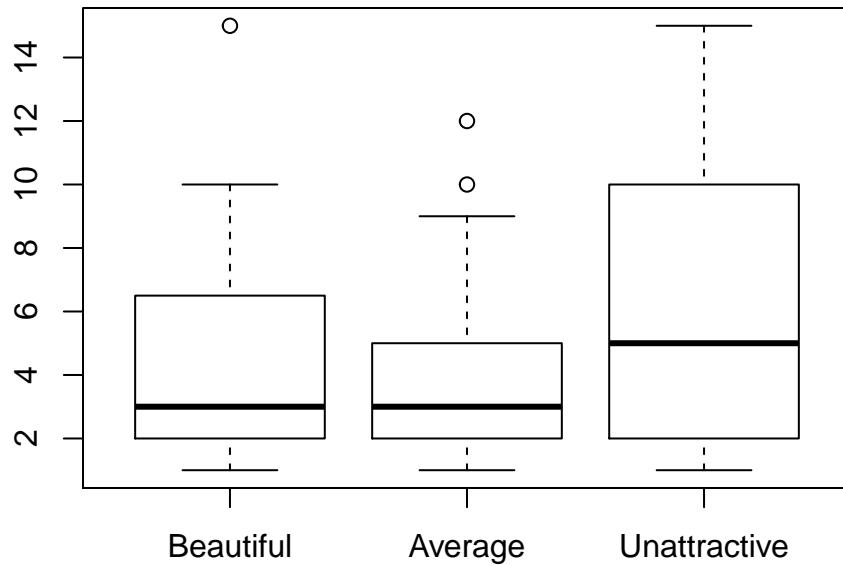


Figure 2.4: Side-by-side boxplot of Years based on picture groups.

distribution (more later on the issues this creates!). Remember that  $\mu$  and  $\sigma$  are parameters where  $\mu$  (“mu”) is our standard symbol for the ***population mean*** and that  $\sigma$  (“sigma”) is the symbol of the ***population standard deviation***.

## 2.2 Beanplots

The other graphical display for comparing multiple groups we will use is a newer display called a ***beanplot*** (Kampstra, 2008). Figure 2.5 shows an example of a beanplot that provides a side-by-side display that contains the density curves, the original observations that generated the density curve in a (jittered) rug-plot, the mean of each group, and the overall mean of the entire data set. For each group, the density curves are mirrored to aid in visual assessment of the shape of the distribution, which makes a “bean” in some cases. This mirroring also creates a shape that resembles a violin with skewed distributions so this display has also been called a “violin plot”. The innovation in the beanplot is to add bold horizontal lines at the mean for each group. It also adds a lighter dashed line for the overall mean. All together this plot shows us information on the center (mean), spread, and shape of the distributions of the responses. Our inferences typically focus on the means of the groups and this plot allows us to compare those across the groups while gaining information on the shapes of the distributions of responses in each group.

To use the **beanplot** function we need to install and load the **beanplot** package (Kampstra, 2014). The function works like the boxplot used previously except that options for `log`, `col`, and `method` need to be specified. Use these<sup>6</sup> options for any beanplots you make: `log=""`, `col="bisque"`, `method="jitter"`

```
require(beanplot)
beanplot(Years~Attr,data=MockJury,log="",col="bisque",method="jitter")
```

Figure 2.5 reinforces the strong right skew that were also detected in the boxplots previously. The three large sentences of 15 years can now be clearly identified, with one in the *Beautiful* group and two in the *Unattractive* group. The *Unattractive* group seems to have more high observations than the other groups even

<sup>6</sup>Well, you can use other colors (try “lightblue” for example), but I think bisque looks nice in these plots.

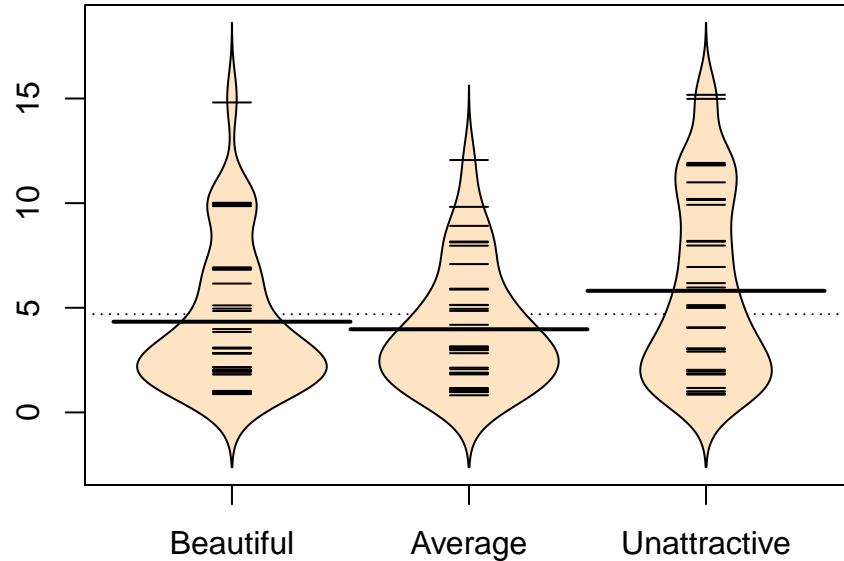


Figure 2.5: Beanplot of Years by picture group. Long, bold lines correspond to mean of each group.

though the *Beautiful* group had the largest number of observations around 10 years. The mean sentence was highest for the *Unattractive* group and the difference in the means between *Beautiful* and *Average* was small.

In this example, it appears that the mean for *Unattractive* is larger than the other two groups. But is this difference real? We will never know the answer to that question, but we can assess how likely we are to have seen a result as extreme or more extreme than our result, assuming that there is no difference in the means of the groups. And if the observed result is (extremely) unlikely to occur, then we can reject the hypothesis that the groups have the same mean and conclude that there is evidence of a real difference. To start exploring whether there are differences in the means, we need to have numerical values to compare. We can get means and standard deviations by groups easily using the same formula notation with the `mean` and `sd` functions if the `mosaic` package is loaded.

```
mean(Years ~ Attr, data = MockJury)
```

```
##      Beautiful      Average Unattractive
##      4.333333     3.973684    5.810811
sd(Years ~ Attr, data = MockJury)
```

```
##      Beautiful      Average Unattractive
##      3.405362     2.823519    4.364235
```

We can also use the `favstats` function to get those summaries and others.

```
favstats(Years ~ Attr, data = MockJury)
```

```
##          Attr min Q1 median   Q3 max     mean      sd n missing
## 1    Beautiful  1  2       3  6.5 15 4.333333 3.405362 39      0
## 2      Average  1  2       3  5.0 12 3.973684 2.823519 38      0
## 3 Unattractive  1  2       5 10.0 15 5.810811 4.364235 37      0
```

Based on these results, we can see that there is an estimated difference of almost 2 years in the mean sentence between *Average* and *Unattractive* groups. Because there are three groups being compared in this study, we will have to wait until Chapter 3 and the One-Way ANOVA test to fully assess evidence related to some difference among the three groups. For now, we are going to focus on comparing the mean *Years* between

*Average* and *Unattractive* groups – which is a **2 independent sample mean** situation and something you should have seen before. Remember that the “independent” sample part of this refers to observations that are independently observed for the two groups as opposed to the paired sample situation that you may have explored where one observation from the first group is related to an observation in the second group (repeated measures on the same person or the famous “twin” studies with one twin assigned to each group).

Here we are going to use the “simple” two independent group scenario to review some basic statistical concepts and connect two different frameworks for conducting statistical inference: randomization and parametric inference techniques. **Parametric** statistical methods involve making assumptions about the distribution of the responses and obtaining confidence intervals and/or p-values using a *named* distribution (like the z or t-distributions). Typically these results are generated using formulas and looking up areas under curves or cutoffs using a table or a computer. **Randomization**-based statistical methods use a computer to shuffle, sample, or simulate observations in ways that allow you to obtain distributions of possible results to find areas and cutoffs without resorting to using tables and named distributions. Randomization methods are what are called **nonparametric** methods that often make fewer assumptions (they are **not free of assumptions!**) and so can handle a larger set of problems more easily than parametric methods. When the assumptions involved in the parametric procedures are met by a data set, the randomization methods often provide very similar results to those provided by the parametric techniques. To be a more sophisticated statistical consumer, it is useful to have some knowledge of both of these approaches to statistical inference and the fact that they can provide similar results might deepen your understanding of both approaches.

We will start with comparing the *Average* and *Unattractive* groups to compare these two ways of doing inference. We could remove the *Beautiful* group observations in a spreadsheet program and read that new data set back into R, but it is actually pretty easy to use R to do data management once the data set is loaded. To remove the observations that came from the *Beautiful* group, we are going to generate a new variable that we will call *NotBeautiful* that is true when observations came from another group (*Average* or *Unattractive*) and false for observations from the *Beautiful* group. To do this, we will apply the **not equal** logical function (!=) to the variable *Attr*, inquiring whether it was different from the "Beautiful" level. You can see the content of the new variable in the output:

```
MockJury$NotBeautiful <- MockJury$Attr != "Beautiful"
MockJury$NotBeautiful
```

```
## [1] FALSE FALSE
## [12] FALSE TRUE
## [23] TRUE TRUE
## [34] TRUE TRUE
## [45] TRUE TRUE
## [56] TRUE TRUE
## [67] TRUE FALSE
## [78] FALSE FALSE
## [89] FALSE FALSE FALSE FALSE FALSE FALSE TRUE TRUE TRUE TRUE TRUE
## [100] TRUE TRUE
## [111] TRUE TRUE TRUE TRUE
```

This new variable is only FALSE for the *Beautiful* responses as we can see if we compare some of the results from the original and new variable:

```
head(data.frame(MockJury$Attr, MockJury$NotBeautiful))
```

```
## MockJury.Attr MockJury.NotBeautiful
## 1 Beautiful FALSE
## 2 Beautiful FALSE
## 3 Beautiful FALSE
## 4 Beautiful FALSE
## 5 Beautiful FALSE
## 6 Beautiful FALSE
```

```
tail(data.frame(MockJury$Attr, MockJury$NotBeautiful))
```

```
##      MockJury.Attr MockJury.NotBeautiful
## 109      Average           TRUE
## 110      Average           TRUE
## 111      Average           TRUE
## 112      Average           TRUE
## 113      Average           TRUE
## 114      Average           TRUE
```

To get rid of one of the groups, we need to learn a little bit about data management in R. **Brackets** (`[, ]`) are used to modify the rows or columns in a data.frame with entries before the comma operating on rows and entries after the comma on the columns. For example, if you want to see the results for the 5<sup>th</sup> subject, you can reference the 5<sup>th</sup> row of the data.frame using `[5, ]` after the data.frame name:

```
MockJury[5,]
```

```
##      Attr     Crime Years Serious exciting calm independent sincere warm
## 5 Beautiful Burglary    7      9      1      1      5      1      8
##   phyattr sociable kind intelligent strong sophisticated happy ownPA
## 5      8      9      4      7      9      9      8      7
##   NotBeautiful
## 5      FALSE
```

We could just extract the *Years* response for the 5<sup>th</sup> subject by incorporating information on the row and column of interest (*Years* is the 3<sup>rd</sup> column):

```
MockJury[5,3]
```

```
## [1] 7
```

In R, we can use logical vectors to keep any rows of the data.frame where the variable is true and drop any rows where it is false by placing the logical variable in the first element of the brackets. The reduced version of the data set should be saved with a different name such as *MockJury2* that is used here to reduce the chances of confusing it with the previous full data set:

```
MockJury2 <- MockJury[MockJury$NotBeautiful,]
```

You will always want to check that the correct observations were dropped either using `View(MockJury2)` or by doing a quick summary of the *Attr* variable in the new data.frame.

```
summary(MockJury2$Attr)
```

```
##      Beautiful      Average Unattractive
##            0             38            37
```

It ends up that R remembers the *Beautiful* category even though there are 0 observations in it now and that can cause us some problems. When we remove a group of observations, we sometimes need to clean up categorical variables to just reflect the categories that are present. The `factor` function creates categorical variables based on the levels of the variables that are observed and is useful to run here to clean up *Attr*.

```
MockJury2$Attr <- factor(MockJury2$Attr)
```

```
summary(MockJury2$Attr)
```

```
##      Average Unattractive
##            38            37
```

Now if we remake the boxplots and beanplots, they only contain results for the two groups of interest here as seen in Figure 2.6.

```
par(mfrow=c(1,2))
boxplot(Years ~ Attr, data=MockJury2)
beanplot(Years ~ Attr, data=MockJury2, log="", col="bisque", method="jitter")
```

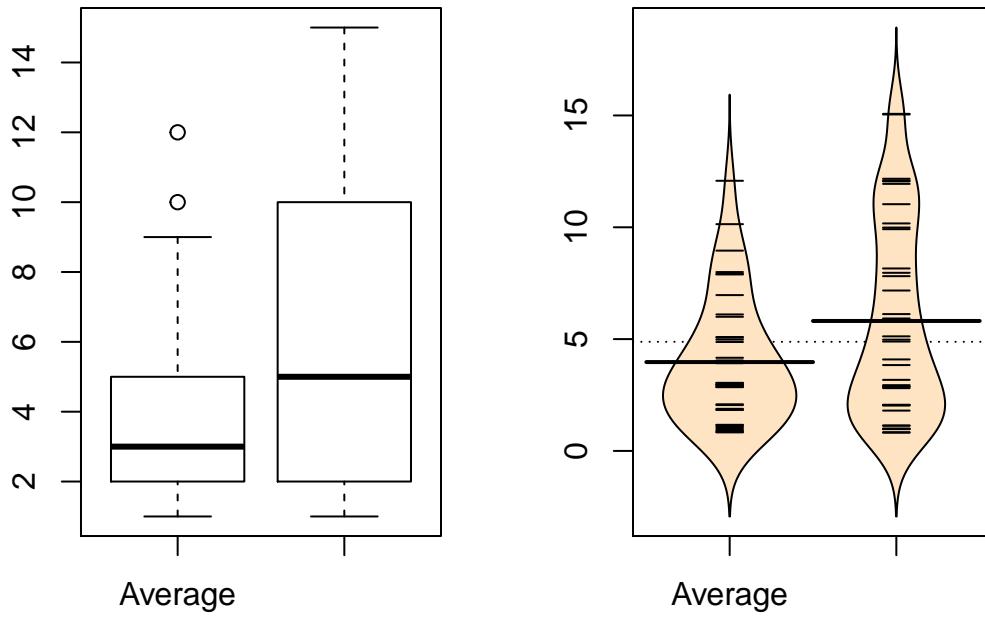


Figure 2.6: Boxplot and beanplot of the Years responses on the reduced data set.

The two-sample mean techniques you learned in your previous course all start with comparing the means the two groups. We can obtain the two means using the `mean` function or directly obtain the difference in the means using the `diffmean` function (both require the `mosaic` package). The `diffmean` function provides  $\bar{x}_{Unattractive} - \bar{x}_{Average}$  where  $\bar{x}$  (read as “x-bar”) is the sample mean of observations in the subscripted group. Note that there are two directions that you could compare the means and this function chooses to take the mean from the second group name *alphabetically* and subtract the mean from the first alphabetical group name. It is always good to check the direction of this calculation as having a difference of  $-1.84$  years versus  $1.84$  years could be important.

```
mean(Years ~ Attr, data=MockJury2)
```

```
##      Average Unattractive
##      3.973684     5.810811
diffmean(Years ~ Attr, data=MockJury2)

## diffmean
## 1.837127
```

## 2.3 Models, hypotheses, and permutations for the 2 sample mean situation

There appears to be some evidence that the *Unattractive* group is getting higher average lengths of sentences from the prisoner “jurors” than the *Average* group, but we want to make sure that the difference is real – that there is evidence to reject the assumption that the means are the same “in the population”. First, a *null*

*hypothesis*<sup>7</sup> which defines a **null model**<sup>8</sup> needs to be determined in terms of **parameters** (the true values in the population). The research question should help you determine the form of the hypotheses for the assumed population. In the 2 independent sample mean problem, the interest is in testing a null hypothesis of  $H_0 : \mu_1 = \mu_2$  versus the alternative hypothesis of  $H_A : \mu_1 \neq \mu_2$ , where  $\mu_1$  is the parameter for the true mean of the first group and  $\mu_2$  is the parameter for the true mean of the second group. The alternative hypothesis involves assuming a statistical model for the  $i^{th}$  ( $i = 1, \dots, n_j$ ) response from the  $j^{th}$  ( $j = 1, 2$ ) group,  $y_{ij}$ , that involves modeling it as  $y_{ij} = \mu_j + \varepsilon_{ij}$ , where we assume that  $\varepsilon_{ij} \sim N(0, \sigma^2)$ . For the moment, focus on the models that either assume the means are the same (null) or different (alternative), which imply:

- Null Model:  $y_{ij} = \mu + \varepsilon_{ij}$  There is **no** difference in **true** means for the two groups.
- Alternative Model:  $y_{ij} = \mu_j + \varepsilon_{ij}$  There is **a** difference in **true** means for the two groups.

Suppose we are considering the alternative model for the 4th observation ( $i = 4$ ) from the second group ( $j = 2$ ), then the model for this observation is  $y_{42} = \mu_2 + \varepsilon_{42}$ , that defines the response as coming from the true mean for the second group plus a random error term for that observation,  $\varepsilon_{42}$ . For, say, the 5th observation from the first group ( $j = 1$ ), the model is  $y_{51} = \mu_1 + \varepsilon_{51}$ . If we were working with the null model, the mean is always the same ( $\mu$ ) - the group specified does not change the mean we use for that observation.

It can be helpful to think about the null and alternative models graphically. By assuming the null hypothesis is true (means are equal) and that the random errors around the mean follow a normal distribution, we assume that the truth is as displayed in the left panel of Figure 2.7 – two normal distributions with the same mean and variability. The alternative model allows the two groups to potentially have different means, such as those displayed in the right panel of Figure 2.7 where the second group has a larger mean. Note that in this scenario, we assume that the observations all came from the same distribution except that they had different means. Depending on the statistical procedure we are using, we basically are going to assume that the observations ( $y_{ij}$ ) either were generated as samples from the null or alternative model. You can imagine drawing observations at random from the pictured distributions. For hypothesis testing, the null model is assumed to be true and then the unusualness of the actual result is assessed relative to that assumption. In hypothesis testing, we have to decide if we have enough evidence to reject the assumption that the null model (or hypothesis) is true. If we reject the null hypothesis, then we would conclude that the other model considered (the alternative model) is more reasonable. The researchers obviously would have hoped to encounter some sort of noticeable difference in the sentences provided for the different pictures and been able to find enough evidence to reject the null model where the groups “look the same”.

In statistical inference, null hypotheses (and their implied models) are set up as “straw men” with every interest in rejecting them even though we assume they are true to be able to assess the evidence against them. Consider the original study design here, the pictures were randomly assigned to the subjects. If the null hypothesis were true, then we would have no difference in the population means of the groups. And this would apply if we had done a different random assignment of the pictures to the subjects. So let’s try this: assume that the null hypothesis is true and randomly re-assign the treatments (pictures) to the observations that were obtained. In other words, keep the sentences (*Years*) the same and shuffle the group labels randomly. The technical term for this is doing a **permutation** (a random shuffling of the treatments relative to the responses). If the null is true and the means in the two groups are the same, then we should be able to re-shuffle the groups to the observed sentences (*Years*) and get results similar to those we actually observed. If the null is false and the means are really different in the two groups, then what we observed should differ from what we get under other random permutations. The differences between the two groups should be more noticeable in the observed data set than in (most) of the shuffled data sets. It helps to see an example of a permutation of the labels to understand what this means here.

In the **mosaic** package, the **shuffle** function allows us to easily perform a permutation<sup>9</sup>. Just one time, we

---

<sup>7</sup>The hypothesis of no difference that is typically generated in the hopes of being rejected in favor of the alternative hypothesis which contains the sort of difference that is of interest in the application.

<sup>8</sup>The null model is the statistical model that is implied by the chosen null hypothesis. Here, a null hypothesis of no difference translates to having a model with the same mean for both groups.

<sup>9</sup>We’ll see the **shuffle** function in a more common usage below; while the code to generate **Perm1** is provided, it isn’t something to worry about right now.

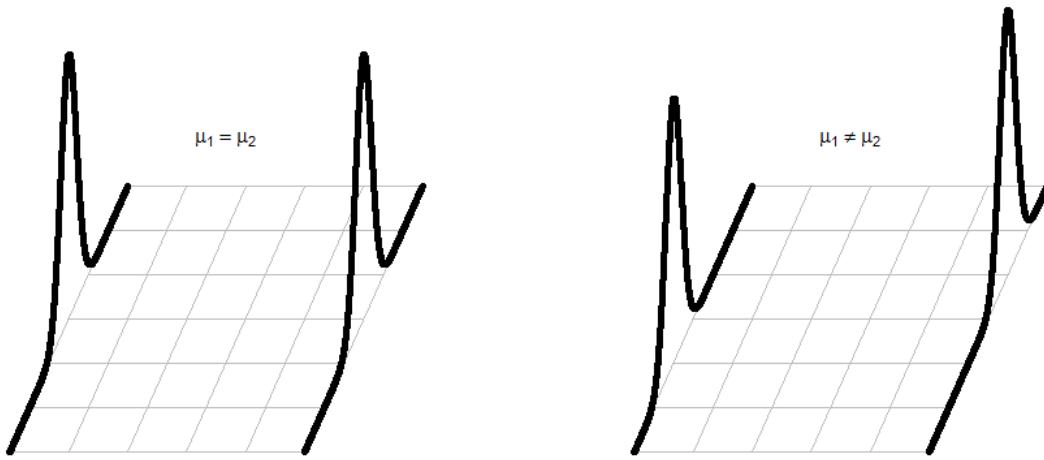


Figure 2.7: Illustration of the assumed situations under the null (left) and a single possibility that could occur if the alternative were true (right) and the true means were different.

can explore what a permutation of the treatment labels could look like in the `PermutedAttr` variable below. Note that the `Years` are held in the same place the group labels are shuffled.

```
set.seed(1234)
Perm1 <- with(MockJury2,data.frame(Years,Attr,PermutedAttr=shuffle(Attr)))
Perm1
```

```
##   Years      Attr PermutedAttr
## 1      1 Unattractive Unattractive
## 2      4 Unattractive     Average
## 3      3 Unattractive     Average
## 4      2 Unattractive     Average
## 5      8 Unattractive     Average
## 6      8 Unattractive     Average
## 7      1 Unattractive Unattractive
## 8      1 Unattractive Unattractive
## 9      5 Unattractive     Average
## 10     7 Unattractive Unattractive
## 11     1 Unattractive     Average
## 12     5 Unattractive Unattractive
## 13     2 Unattractive Unattractive
## 14    12 Unattractive     Average
## 15    10 Unattractive     Average
## 16     1 Unattractive     Average
## 17     6 Unattractive Unattractive
## 18     2 Unattractive     Average
## 19     5 Unattractive Unattractive
## 20    12 Unattractive Unattractive
## 21     6 Unattractive     Average
## 22     3 Unattractive     Average
## 23     8 Unattractive     Average
## 24     4 Unattractive Unattractive
## 25    10 Unattractive Unattractive
```

```

## 26    10 Unattractive      Average
## 27    15 Unattractive Unattractive
## 28    15 Unattractive      Average
## 29     3 Unattractive      Average
## 30     3 Unattractive      Average
## 31     3 Unattractive Unattractive
## 32    11 Unattractive      Average
## 33    12 Unattractive      Average
## 34     2 Unattractive Unattractive
## 35     1 Unattractive Unattractive
## 36     1 Unattractive Unattractive
## 37    12 Unattractive      Average
## 38     5      Average Unattractive
## 39     5      Average Unattractive
## 40     4      Average Unattractive
## 41     3      Average Unattractive
## 42     6      Average      Average
## 43     4      Average      Average
## 44     9      Average      Average
## 45     8      Average Unattractive
## 46     3      Average      Average
## 47     2      Average Unattractive
## 48    10      Average      Average
## 49     1      Average Unattractive
## 50     1      Average Unattractive
## 51     3      Average Unattractive
## 52     1      Average      Average
## 53     3      Average      Average
## 54     5      Average      Average
## 55     8      Average Unattractive
## 56     3      Average      Average
## 57     1      Average      Average
## 58     1      Average Unattractive
## 59     1      Average      Average
## 60     2      Average      Average
## 61     2      Average Unattractive
## 62     1      Average      Average
## 63     1      Average Unattractive
## 64     2      Average Unattractive
## 65     3      Average Unattractive
## 66     4      Average Unattractive
## 67     5      Average      Average
## 68     3      Average Unattractive
## 69     3      Average      Average
## 70     3      Average      Average
## 71     2      Average Unattractive
## 72     7      Average Unattractive
## 73     6      Average Unattractive
## 74    12      Average Unattractive
## 75     8      Average      Average

```

If you count up the number of subjects in each group by counting the number of times each label (Average, Unattractive) occurs, it is the same in both the `Attr` and `PermutedAttr` columns. Permutations involve randomly re-ordering the values of a variable – here the `Attr` group labels – without changing the content of the variable. This result can also be generated using what is called *sampling without replacement*: sequentially select  $n$  labels from the original variable, removing each used label and making sure that each original `Attr` label is selected once and only once. The new, randomly selected order of selected labels provides the permuted labels. Stepping through the process helps to understand how it works: after the

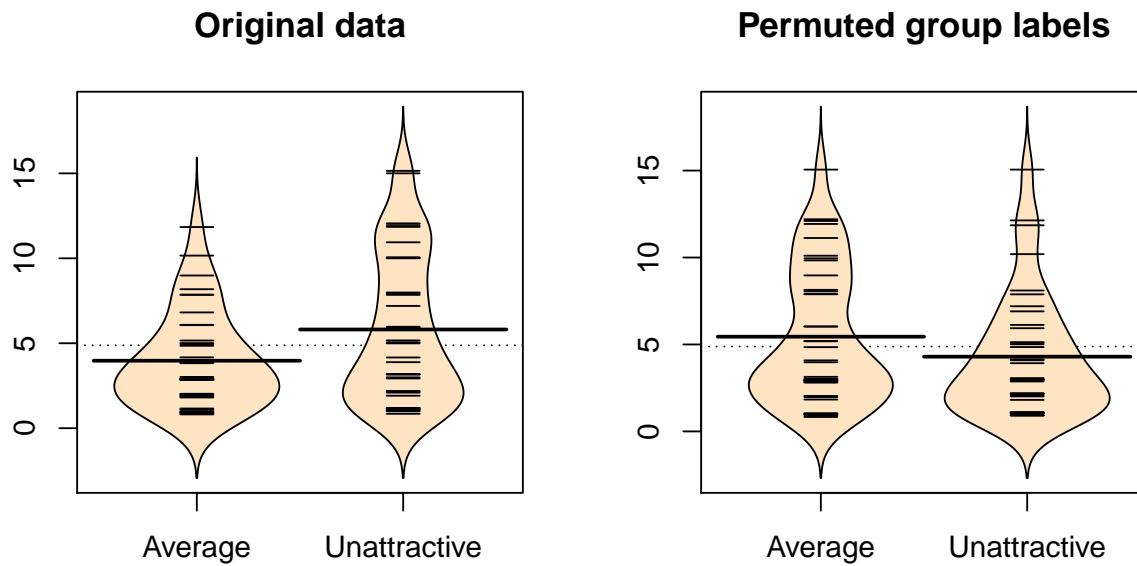


Figure 2.8: Boxplots of Years responses versus actual treatment groups and permuted groups.

initial random sample of one label, there would  $n - 1$  choices possible; on the  $n^{th}$  selection, there would only be one label remaining to select. This makes sure that all original labels are re-used but that the order is random. Sampling without replacement is like picking names out of a hat, one-at-a-time, and not putting the names back in after they are selected. It is an exhaustive process for all the original observations. ***Sampling with replacement***, in contrast, involves sampling from the specified list with each observation having an equal chance of selection for each sampled observation – in other words, observations can be selected more than once. This is like picking  $n$  names out of a hat that contains  $n$  names, except that every time a name is selected, it goes back into the hat – we'll use this technique in Section 2.8 to do what is called ***bootstrapping***. Both sampling mechanisms can be used to generate inferences but each has particular situations where they are most useful. For hypothesis testing, we will use permutations (sampling without replacement).

The comparison of the beanplots for the real data set and permuted version of the labels is what is really interesting (Figure 2.8). The original difference in the sample means of the two groups was 1.84 years (Unattractive minus Average). The sample means are the ***statistics*** that estimate the parameters for the true means of the two groups. In the permuted data set, the difference in the means is 1.15 years in the opposite direction (Average had a higher mean than Unattractive in the permuted data).

```
mean(Years ~ PermutedAttr, data=Perm1)
```

```
##      Average Unattractive
##      5.447368     4.297297
diffmean(Years ~ PermutedAttr, data=Perm1)

##  diffmean
## -1.150071
```

These results suggest that the observed difference was larger than what we got when we did a single permutation although it was only a little bit larger than a difference we could observe in permutations if we ignore the difference in directions. Conceptually, permuting observations between group labels is consistent with the null hypothesis – this is a technique to generate results that we might have gotten if the null hypothesis were true since the responses are the same in the two groups if the null is true. We just need to repeat the permutation process many times and track how unusual our observed result is relative to this distribution of potential responses if the null were true. If the observed differences are unusual relative to the

results under permutations, then there is evidence against the null hypothesis, the null hypothesis should be rejected ( Reject  $H_0$ ), and a conclusion should be made, in the direction of the alternative hypothesis, that there is evidence that the true means differ. If the observed differences are similar to (or at least not unusual relative to) what we get under random shuffling under the null model, we would have a tough time concluding that there is any real difference between the groups based on our observed data set.

## 2.4 Permutation testing for the 2 sample mean situation

In any testing situation, you must define some function of the observations that gives us a single number that addresses our question of interest. This quantity is called a ***test statistic***. These often take on complicated forms and have names like  $t$  or  $z$  statistics that relate to their parametric (named) distributions so we know where to look up ***p-values***<sup>10</sup>. In randomization settings, they can have simpler forms because we use the data set to find the distribution of the statistic and don't need to rely on a named distribution. We will label our test statistic  $T$  (for  $\mathbf{T}$  statistic) unless the test statistic has a commonly used name. Since we are interested in comparing the means of the two groups, we can define

$$T = \bar{x}_{Unattractive} - \bar{x}_{Average},$$

which coincidentally is what the `diffmean` function provided us previously. We label our ***observed test statistic*** (the one from the original data set) as

$$T_{obs} = \bar{x}_{Unattractive} - \bar{x}_{Average},$$

which happened to be 1.84 years here. We will compare this result to the results for the test statistic that we obtain from permuting the group labels. To denote permuted results, we will add a \* to the labels:

$$T^* = \bar{x}_{Unattractive} - \bar{x}_{Average^*}.$$

We then compare the  $T_{obs} = \bar{x}_{Unattractive} - \bar{x}_{Average} = 1.84$  to the distribution of results that are possible for the permuted results ( $T^*$ ) which corresponds to assuming the null hypothesis is true.

We need to consider lots of permutations to do a permutation test. In contrast to your introductory statistics course where, if you did this, it was just a click away, we are going to learn what was going on under the hood. Specifically, we need a ***for loop*** in R to be able to repeatedly generate the permuted data sets and record  $T^*$  for each one. Loops are a basic programming task that make randomization methods possible as well as potentially simplifying any repetitive computing task. To write a “for loop”, we need to choose how many times we want to do the loop (call that `B`) and decide on a counter to keep track of where we are at in the loops (call that `b`, which goes from 1 up to `B`). The simplest loop just involves printing out the index, `print(b)` at each step. This is our first use of curly braces, { and }, that are used to group the code we want to repeatedly run as we proceed through the loop. By typing the following code in the script window and then highlighting it all and hitting the run button, R will go through the loop 5 times, printing out the counter:

```
B <- 5
for (b in (1:B)){
  print(b)
}
```

Note that when you highlight and run the code, it will look about the same with “+” printed after the first line to indicate that all the code is connected when it appears in the console, looking like this:

---

<sup>10</sup>P-values are the probability of obtaining a result as extreme as or more extreme than we observed given that the null hypothesis is true.

```
> for(b in (1:B)){
+   print(b)
+}
```

When you run these three lines of code, the console will show you the following output:

```
[1] 1
[1] 2
[1] 3
[1] 4
[1] 5
```

Instead of printing the counter, we want to use the loop to repeatedly compute our test statistic across B random permutations of the observations. The `shuffle` function performs permutations of the group labels relative to responses and the `diffmean` difference in the two group means in the permuted data set. For a single permutation, the combination of shuffling `Attr` and finding the difference in the means, storing it in a variable called `Ts` is:

```
Ts <- diffmean(Years ~ shuffle(Attr), data=MockJury2)
Ts
```

```
## diffmean
## -0.616643
```

And putting this inside the `print` function allows us to find the test statistic under 5 different permutations easily:

```
B <- 5
for (b in (1:B)){
  Ts <- diffmean(Years ~ shuffle(Attr), data=MockJury2)
  print(Ts)
}

## diffmean
## -0.8300142
## diffmean
## -0.1365576
## diffmean
## -0.08321479
## diffmean
## 0.5035562
## diffmean
## 1.677098
```

Finally, we would like to store the values of the test statistic instead of just printing them out on each pass through the loop. To do this, we need to create a variable to store the results, let's call it `Tstar`. We know that we need to store B results so will create a vector of length B, which contains B elements, full of missing values (`NA`) using the `matrix` function:

```
Tstar <- matrix(NA, nrow=B)
Tstar

##      [,1]
## [1,]    NA
## [2,]    NA
## [3,]    NA
## [4,]    NA
## [5,]    NA
```

Now we can run our loop B times and store the results in `Tstar`.

```
for (b in (1:B)){
  Tstar[b] <- diffmean(Years ~ shuffle(Attr), data=MockJury2)
```

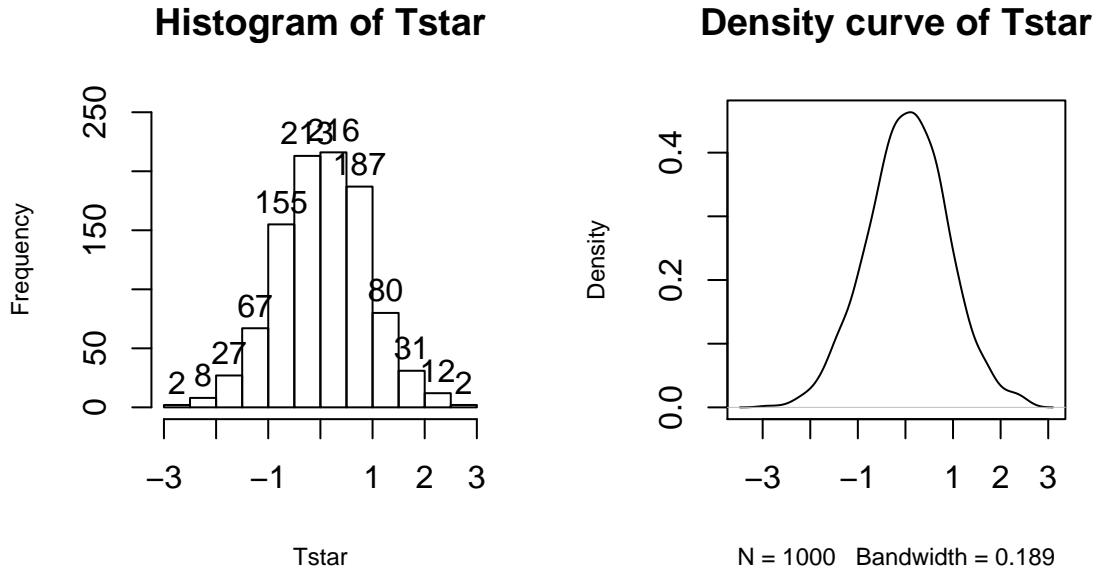


Figure 2.9: Histogram (left, with counts in bars) and density curve (right) of values of test statistic for 1,000 permutations.

```
}
Tstar

##          [,1]
## [1,] -0.08321479
## [2,]  0.23684211
## [3,] -0.24324324
## [4,] -0.61664296
## [5,]  0.66358464
```

Five permutations are still not enough to assess whether our  $T_{obs}$  of 1.84 is unusual and we need to do many permutations to get an accurate assessment of the possibilities under the null hypothesis. It is common practice to consider something like 1,000 permutations. The `Tstar` vector when we set  $B$  the permutation distribution for the selected test statistic under<sup>11</sup> the null hypothesis – what is called the **null distribution** of the statistic. The null distribution is the distribution of possible values of a statistic under the null hypothesis. We want to visualize this distribution and use it to assess how unusual our  $T_{obs}$  result of 1.84 years was relative to all the possibilities under permutations (under the null hypothesis). So we repeat the loop, now with  $B = 1000$  and generate a histogram, density curve and summary statistics of the results:

```
par(mfrow=c(1,2))
B <- 1000
Tstar <- matrix(NA, nrow=B)
for (b in (1:B)){
  Tstar[b] <- diffmean(Years ~ shuffle(Attr), data=MockJury2)
}
hist(Tstar, label=T)
plot(density(Tstar), main="Density curve of Tstar")
```

<sup>11</sup>We often say “under” in statistics and we mean “given that the following is true”.

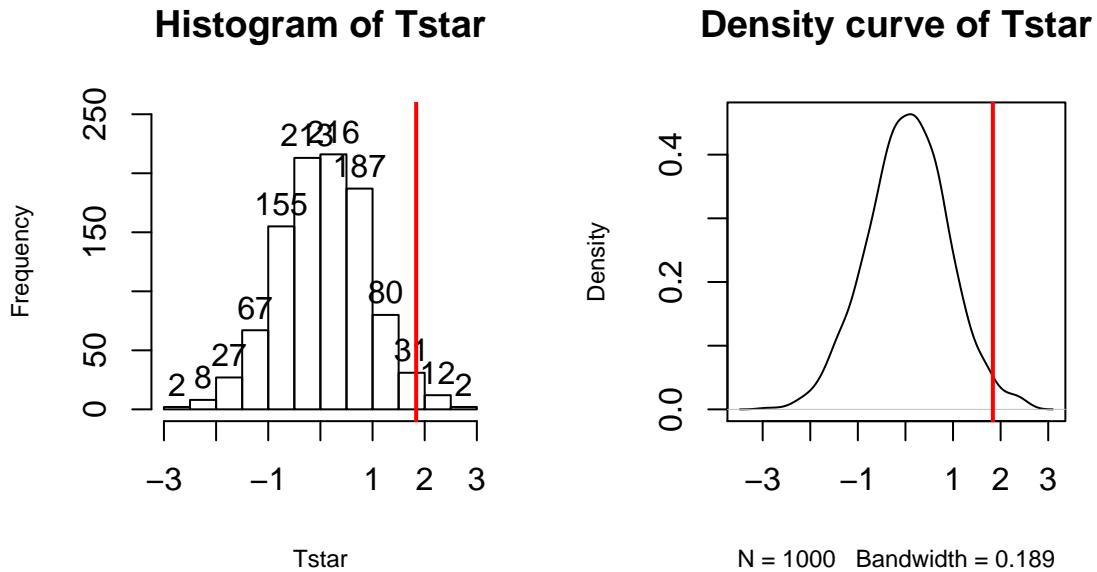


Figure 2.10: Histogram (left) and density curve (right) of values of test statistic for 1,000 permutations with bold vertical line for value of observed test statistic.

```
favstats(Tstar)
```

```
##      min      Q1     median      Q3      max      mean      sd
## -2.910384 -0.5099573 0.07681366 0.6102418 2.530583 0.04694168 0.8497364
##      n missing
##    1000      0
```

Figure 2.9 contains visualizations of  $T^*$  and the `favstats` summary provides the related numerical summaries. Our observed  $T_{obs}$  of 1.84 seems fairly unusual relative to these results with only 20  $T^*$  values over 2 based on the histogram. We need to make more specific comparisons of the permuted results versus our observed result to be able to clearly decide whether our observed result is really unusual.

To make the comparisons more concrete, first we can enhance the previous graphs by adding the value of the test statistic from the real data set, as shown in Figure 2.10, using the `abline` function.

```
par(mfrow=c(1,2))
Tobs <- 1.837
hist(Tstar, labels=T)
abline(v=Tobs, lwd=2, col="red")
plot(density(Tstar),main="Density curve of Tstar")
abline(v=Tobs, lwd=2, col="red")
```

Second, we can calculate the exact number of permuted results that were larger than what we observed. To calculate the proportion of the 1,000 values that were larger than what we observed, we will use the `pdata` function. To use this function, we need to provide the distribution of values to compare to the cut-off (`Tstar`), the cut-off point (`Tobs`), and whether we want calculate the proportion that are below (left of) or above (right of) the cut-off (`lower.tail=F` option provides the proportion of values above the cutoff of interest).

```
pdata(Tstar, Tobs, lower.tail=F)
```

```
## [1] 0.02
```

The proportion of 0.02 tells us that 20 of the 1,000 permuted results (2%) were larger than what we observed.

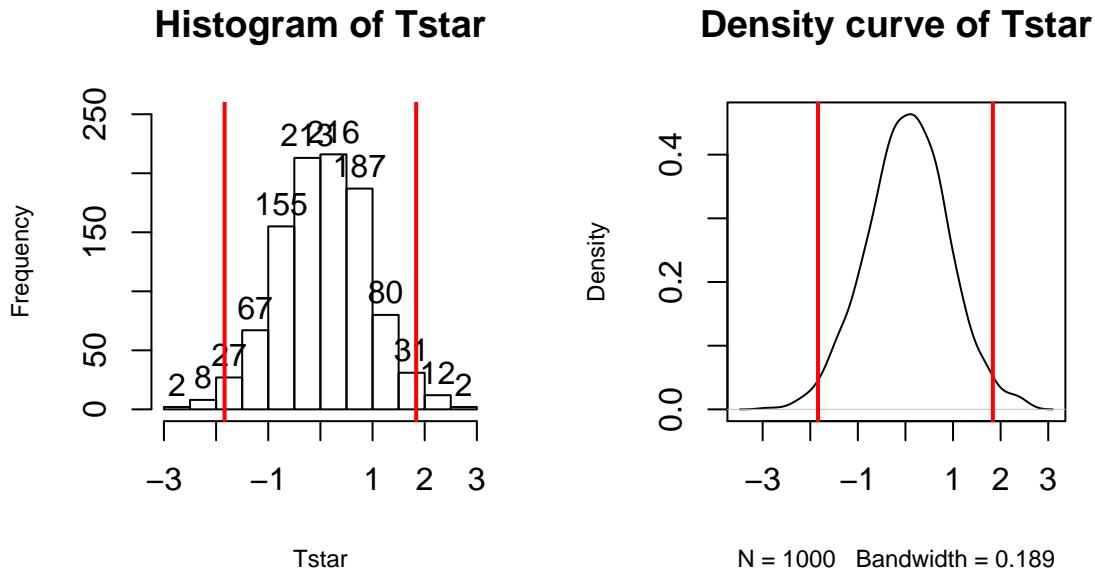


Figure 2.11: Histogram and density curve of values of test statistic for 1,000 permutations with bold lines for value of observed test statistic and its opposite value required for performing two-sided test.

This type of work is how we can generate *p-values* using permutation distributions. P-values, as you should remember, are the probability of getting a result as extreme as or more extreme than what we observed, given that the null is true. Finding only 20 permutations of 1,000 that were larger than our observed result suggests that it is hard to find a result like what we observed if there really were no difference, although it is not impossible.

When testing hypotheses for two groups, there are two types of alternative hypotheses, one-sided or two-sided. **One-sided tests** involve only considering differences in one-direction (like  $\mu_1 > \mu_2$ ) and are performed when researchers can decide *a priori*<sup>12</sup> which group should have a larger mean if there is going to be any sort of difference. In this situation, we did not know enough about the potential impacts of the pictures to know which group should be larger than the other so should do a two-sided test. It is important to remember that you can't look at the responses to decide on the hypotheses. It is often safer and more *conservative*<sup>13</sup> to start with a **two-sided alternative** ( $H_A : \mu_1 \neq \mu_2$ ). To do a 2-sided test, find the area larger than what we observed as above. We also need to add the area in the other tail (here the left tail) similar to what we observed in the right tail. Some people suggest doubling the area in one tail but we will collect information on the number that were more extreme than the same value in the other tail. In other words, we count the proportion over 1.84 and below -1.84. So we need to also find how many of the permuted results were smaller than -1.84 years to add to our previous proportion. Using `pdata -Tobs` as the cut-off and `lower.tail=T` provides this result:

```
pdata(Tstar, -Tobs, lower.tail=T)
```

```
## [1] 0.014
```

So the p-value to test our null hypothesis of no difference in the true means between the groups is  $0.02 + 0.014$ , providing a p-value of 0.034. Figure 2.11 shows both cut-offs on the histogram and density curve.

<sup>12</sup>This is a fancy way of saying “in advance”, here in advance of seeing the observations.

<sup>13</sup>Statistically, a conservative method is one that provides less chance of rejecting the null hypothesis in comparison to some other method or less than some pre-defined standard.

```
par(mfrow=c(1,2))
hist(Tstar, labels=T)
abline(v=c(-1,1)*Tobs, lwd=2, col="red")
plot(density(Tstar),main="Density curve of Tstar")
abline(v=c(-1,1)*Tobs, lwd=2, col="red")
```

In general, the ***one-sided test p-value*** is the proportion of the permuted results that are more extreme than observed in the direction of the *alternative hypothesis* (lower or upper tail, remembering that this also depends on the direction of the difference taken). For the 2-sided test, the p-value is the proportion of the permuted results that are *less than the negative version of the observed statistic and greater than the positive version of the observed statistic*. Using absolute values ( $| |$ ), we can simplify this: the ***two-sided p-value*** is the *proportion of the /permuted statistics/ that are larger than /observed statistic/*. This will always work and finds areas in both tails regardless of whether the observed statistic is positive or negative. In R, the **abs** function provides the ***absolute value*** and we can again use **pdata** to find our p-value in one line of code:

```
pdata(abs(Tstar), abs(Tobs), lower.tail=F)
```

```
## [1] 0.034
```

We will discuss the choice of ***significance level*** below, but for the moment, assume that  $\alpha$  is chosen to be our standard value of 0.05. Since the p-value is smaller than  $\alpha$ , this suggests that we can ***reject the null hypothesis*** and conclude that there is evidence of some difference in the true mean sentences given between the two types of pictures.

Before we move on, let's note some interesting features of the permutation distribution of the difference in the sample means shown in Figure 2.11.

1. It is basically centered at 0. Since we are performing permutations assuming the null model is true, we are assuming that  $\mu_1 = \mu_2$  which implies that  $\mu_1 - \mu_2 = 0$ . This also suggests that 0 should be the center of the permutation distribution and it was.
2. It is approximately normally distributed. This is due to the ***Central Limit Theorem***<sup>14</sup>, where the ***sampling distribution*** (distribution of all possible results for samples of this size) of the difference in sample means ( $\bar{x}_1 - \bar{x}_2$ ) becomes more and normally distributed as the sample sizes increase. With 38 and 37 observations in the groups, we are likely to have a relatively normal looking distribution of the difference in the sample means. This result will allow us to use a parametric method to approximate this sampling distribution under the null model if some assumptions are met, as we'll discuss below.
3. Our observed difference in the sample means (1.84 years) is a fairly unusual result relative to the rest of these results but there are some permuted data sets that produce more extreme differences in the sample means. When the observed differences are really large, we may not see any permuted results that are as extreme as what we observed. When **pdata** gives you 0, the p-value should be reported to be smaller than 0.0001 (**not 0!**) since it happened in less than 1 in 1000 tries but does occur once – in the actual data set.
4. Since our null model is not specific about the direction of the difference, considering a result like ours but in the other direction (-1.84 years) needs to be included. The observed result seems to put about the same area in both tails of the distribution but it is not exactly the same. The small difference in the tails is a useful aspect of this approach compared to the parametric method discussed below as it accounts for slight asymmetry in the sampling distribution.

Earlier, we decided that the p-value was small enough to reject the null hypothesis since it was smaller than our chosen level of significance. In this course, you will often be allowed to use your own judgment about an appropriate significance level in a particular situation (in other words, if we forget to tell you an  $\alpha$ -level, you can still make a decision using a reasonably selected significance level). Remembering that the p-value is the probability you would observe a result like you did (or more extreme), assuming the null hypothesis is true,

---

<sup>14</sup>We'll leave the discussion of the CLT to your previous stat coursework or an internet search. Remember that it has something to do with distributions looking more normal as the sample size increases.

this tells you that the smaller the p-value is, the more evidence you have against the null. The next section provides a more formal review of the hypothesis testing infrastructure, terminology, and some of things that can happen when testing hypotheses.

## 2.5 Hypothesis testing (general)

In hypothesis testing, it is formulated to answer a specific question about a population or true parameter(s) using a statistic based on a data set. In your previous statistics course, you (hopefully) considered one-sample hypotheses about population means and proportions and the two sample mean situation we are focused on here. Our hypotheses relate to trying to answer the question about whether the population mean sentences between the two groups are different, with an initial assumption of no difference.

Hypothesis testing is much like a criminal trial where you are in the role of a jury member or judge, if no jury is present. Initially, the defendant is assumed innocent. In our situation, the true means are assumed to be equal between the groups. Then evidence is presented and, as a juror, you analyze it. In statistical hypothesis testing, data are collected and analyzed. Then you have to decide if we had “enough” evidence to reject the initial assumption (“innocence” that is initially assumed). To make this decision, you want to have previously decided on the standard of evidence required to reject the initial assumption. In criminal cases, “beyond a reasonable doubt” is used. Wikipedia’s definition ([https://en.wikipedia.org/wiki/Reasonable\\_doubt](https://en.wikipedia.org/wiki/Reasonable_doubt)) suggests that this standard is that “there can still be a doubt, but only to the extent that it would not affect a reasonable person’s belief regarding whether or not the defendant is guilty”. In civil trials, a lower standard called a “preponderance of evidence” is used. Based on that defined and pre-decided (*a priori*) measure, you decide that the defendant is guilty or not guilty. In statistics, we compare our p-value to a significance level,  $\alpha$ , which is most of the time selected to be 5%. If our p-value is less than  $\alpha$ , we reject the null hypothesis. The choice of the significance level is like the variation in standards of evidence between criminal and civil trials – and in all situations everyone should know the standards required for rejecting the initial assumption before any information is “analyzed”. Once someone is found guilty, then there is the matter of sentencing which is related to the impacts (“size”) of the crime. In statistics, this is similar to the estimated size of differences and the related judgments about whether the differences are practically important or not. If the crime is proven beyond a reasonable doubt but it is a minor crime, then the sentence will be small. With the same level of evidence and a more serious crime, the sentence will be more dramatic.

There are some important aspects of the testing process to note that inform how we interpret statistical hypothesis test results. When someone is found “not guilty”, it does not mean “innocent”, it just means that there was not enough evidence to find the person guilty “beyond a reasonable doubt”. Not finding enough evidence to reject the null hypothesis does not imply that the true means are equal, just that there was not enough evidence to conclude that they were different. There are many potential reasons why we might fail to reject the null, but the most common one is that our sample size was too small (which is related to having too little evidence).

Throughout this material, we will continue to re-iterate the distinctions between parameters and statistics and want you to be clear about the distinctions between estimates based on the sample and inferences for the population or true values of the parameters of interest. Remember that statistics are summaries of the sample information and parameters are characteristics of populations (which we rarely know). In the two-sample mean situation, the sample means are always at least a little different – that is not an interesting conclusion. What is interesting is whether we have enough evidence to prove that the population or true means differ “beyond a reasonable doubt”.

The scope of any inferences is constrained based on whether there is a **random sample** (RS) and/or **random assignment** (RA). Table 2.2 contains the four possible combinations of these two characteristics of a given study. Random assignment allows for causal inferences for differences that are observed – the difference in treatment levels causes differences in the mean responses. Random sampling (or at least some sort of representative sample) allows inferences to be made to the population of interest. If we do not have RA, then causal inferences cannot be made. If we do not have a representative sample, then our inferences are limited

to the sampled subjects.

Table 2.2: Scope of inference summary.

Random Sampling/Random Assignment	Random Assignment (RA) – Yes (controlled experiment)	Random Assignment (RA) – No (observational study)
<b>Random Sampling (RS)</b> – Yes (or some method that results in a representative sample of population of interest)	Because we have RS, we can generalize inferences to the population the RS was taken from. Because we have RA we can assume the groups were equivalent on all aspects except for the treatment and can establish causal inference.	Can generalize inference to population the RS was taken from but cannot establish causal inference (no RA – cannot isolate treatment variable as only difference among groups, could be confounding variables).
<b>Random Sampling (RS)</b> – No (usually a convenience sample)	Cannot generalize inference to the population of interest because the sample was not random and could be biased – may not be “representative” of the population of interest. Can establish causal inference due to RA → the inference from this type of study applies only to the sample.	Cannot generalize inference to the population of interest because the sample was not random and could be biased – may not be “representative” of the population of interest. Cannot establish causal inference due to lack of RA of the treatment.

A simple example helps to clarify how the scope of inference can change. Suppose we are interested in studying the GPA of students. If we had taken a random sample from, say, the STAT 217 students in a given semester, our scope of inference would be the population of 217 students in that semester. If we had taken a random sample from the entire MSU population, then the inferences would be to the entire MSU population in that semester. These are similar types of problems but the two populations are very different and the group you are trying to make conclusions about should be noted carefully in your results – it does matter! If we did not have a representative sample, say the students could choose to provide this information or not, then we can only make inferences to volunteers. These volunteers might differ in systematic ways from the entire population of STAT 217 students so we cannot safely extend our inferences beyond the group that volunteered.

To consider the impacts of RA versus observational studies, we need to be comparing groups. Suppose that we are interested in differences in the mean GPAs for different sections of STAT 217 and that we take a random sample of students from each section and compare the results and find evidence of some difference. In this scenario, we can conclude that there is some difference in the population of STAT 217 students but we can't say that being in different sections caused the differences in the mean GPAs. Now suppose that we randomly assigned every 217 student to get extra training in one of three different study techniques and found evidence of differences among the training methods. We could conclude that the training methods caused the differences in these students. These conclusions would only apply to STAT 217 students and could not be generalized to a larger population of students. If we took a random sample of STAT 217 students (say only 10 from each section) and then randomly assigned them to one of three training programs and found evidence of differences, then we can say that the training programs caused the differences. We can also say that we have evidence that those differences pertain to the population of STAT 217 students. This seems similar to the scenario where all 217 students participated in the training programs except that by using random sampling, only a fraction of the population needs to actually be studied to make inferences to the entire population of interest – saving time and money.

A quick summary of the terminology of hypothesis testing is useful at this point. The **null hypothesis** ( $H_0$ ) states that there is no difference or no relationship in the population. This is the statement of no effect or no difference and the claim that we are trying to find evidence against. In this chapter,  $H_0: \mu_1 = \mu_2$ . When doing two-group problems, you always need to specify which group is 1 and which one is 2

because the order does matter. The ***alternative hypothesis*** ( $H_1$  or  $H_A$ ) states a specific difference between parameters. This is the research hypothesis and the claim about the population that we hope to demonstrate is more reasonable to conclude than the null hypothesis. In the two-group situation, we can have ***one-sided alternatives***  $H_A:\mu_1 > \mu_2$  (greater than) or  $H_A:\mu_1 < \mu_2$  (less than) or, the more common, ***two-sided alternative***  $H_A:\mu_1 \neq \mu_2$  (not equal to). We usually default to using two-sided tests because we often do not know enough to know the direction of a difference in advance, especially in more complicated situations. The ***sampling distribution under the null*** is the distribution of all possible values of a statistic under  $H_0$  is true. It is used to calculate the ***p-value***, the probability of obtaining a result as extreme or more extreme than what we observed given that the null hypothesis is true. We will find sampling distributions using ***nonparametric*** approaches (like the permutation approach used above) and ***parametric*** methods (using “named” distributions like the  $t$ ,  $F$ , and  $\chi^2$ ).

Small p-values are evidence against the null hypothesis because the observed result is unlikely due to chance if  $H_0$  is true. Large p-values provide no evidence against  $H_0$  but do not allow us to conclude that there is no difference. The ***level of significance*** is an *a priori* definition of how small the p-value needs to be to provide “enough” (sufficient) evidence against  $H_0$ . This is most useful to prevent sliding the standards after the results are found. We compare the p-value to the level of significance to decide if the p-value is small enough to constitute sufficient evidence to reject the null hypothesis. We use  $\alpha$  to denote the level of significance and most typically use 0.05 which we refer to as the 5% significance level. We compare the p-value to this level and make a decision. The two options for *decisions* are to either *reject the null hypothesis* if the p-value  $\leq \alpha$  or *fail to reject the null hypothesis* if the p-value  $> \alpha$ . When interpreting hypothesis testing results, remember that the p-value is a measure of how unlikely the observed outcome was, assuming that the null hypothesis is true. It is **NOT** the probability of the data or the probability of either hypothesis being true. The p-value, simply, is a measure of evidence against the null hypothesis.

The specific definition of  $\alpha$  is that it is the probability of rejecting  $H_0$  when  $H_0$  is true, the probability of what is called a ***Type I error***. Type I errors are also called ***false rejections***. In the two-group mean situation, a Type I error would be concluding that there is a difference in the true means between the groups when none really exists in the population. In the courtroom setting, this is like falsely finding someone guilty. We don’t want to do this very often, so we use small values of the significance level, allowing us to control the rate of Type I errors at  $\alpha$ . We also have to worry about ***Type II errors***, which are failing to reject the null hypothesis when it’s false. In a courtroom, this is the same as failing to convict a truly guilty person. This most often occurs due to a lack of evidence. You can use the Table 2.3 to help you remember all the possibilities.

Table 2.3: Table of decisions and truth scenarios in a hypothesis testing situation. But we never know the truth in a real situation.

	$H_0$ True	$H_0$ False
<b>FTR <math>H_0</math></b>	Correct decision	Type II error
<b>Reject <math>H_0</math></b>	Type I error	Correct decision

In comparing different procedures, there is an interest in studying the rate or probability of Type I and II errors. The probability of a Type I error was defined previously as  $\alpha$ , the significance level. The ***power*** of a procedure is the probability of rejecting the null hypothesis when it is false. Power is defined as

$$\text{Power} = 1 - \text{Probability}(\text{Type II error}) = \text{Probability}(\text{Reject } H_0 | H_0 \text{ is false}),$$

or, in words, the probability of detecting a difference when it actually exists. We want to use a statistical procedure that controls the Type I error rate at the pre-specified level and has high power to detect false null alternatives. Increasing the sample size is one of the most commonly used methods for increasing the power in a given situation. Sometimes we can choose among different procedures and use the power of the procedures to help us make that selection. Note that there are many ways  $H_0$  false and the power changes based on how

false the null hypothesis actually is. To make this concrete, suppose that the true mean sentences differed by either 1 or 20 years in previous example. The chances of rejecting the null hypothesis are much larger when the groups actually differ by 20 years than if they differ by just 1 year.

After making a decision (was there enough evidence to reject the null or not), we want to make the conclusions specific to the problem of interest. If we reject  $H_0$ , then we can conclude that there was sufficient evidence at the  $\alpha$ -level that the null hypothesis is wrong (and the results point in the direction of the alternative). If we fail to reject  $H_0$  (FTR  $H_0$ ), then we can conclude that there was insufficient evidence at the  $\alpha$ -level to say that the null hypothesis is wrong. We are **NOT** saying that the null is correct and we **NEVER** accept the null hypothesis. We just failed to find enough evidence to say it's wrong. If we find sufficient evidence to reject the null, then we need to revisit the method of data collection and design of the study to discuss scope of inference. Can we discuss causality (due to RA) and/or make inferences to a larger group than those in the sample (due to RS)?

To perform a hypothesis test, there are some steps to remember to complete to make sure you have thought through all the aspects of the results.

---

#### **Outline of 6+ steps to perform a Hypothesis Test**

Isolate the claim to be proved, method to use (define a test statistic T), and significance level.

1. Write the null and alternative hypotheses,
  2. Assess the “Validity Conditions” for the procedure being used (discussed below),
  3. Find the value of the appropriate test statistic,
  4. Find the p-value,
  5. Make a decision, and
  6. Write a conclusion specific to the problem, including scope of inference discussion.
- 

## **2.6 Connecting randomization (nonparametric) and parametric tests**

In developing statistical inference techniques, we need to define the test quantity of interest. To compare the means of two groups, a statistic is needed that measures their differences. In general, for comparing two groups, the choices are simple – a difference in the means often works well and is a natural choice.

There are other options such as tracking the ratio of means or possibly the difference in medians. Instead of just using the difference in the means, we also could “standardize” the difference in the means by dividing by an appropriate quantity that reflects the variation in the difference in the means. All of these are valid and can sometimes provide similar results - it ends up that there are many possibilities for testing using the randomization (nonparametric) techniques introduced previously. Parametric statistical methods focus on means because the statistical theory surrounding means is quite a bit easier (not easy, just easier) than other options but there are just a couple of test statistics that you can use and end up with named distributions to use for generating inferences. Randomization techniques allow inference for other quantities but our focus here will be on using randomization for inferences on means to see the similarities with the more traditional parametric procedures.

In two-sample mean situations, instead of working just with the difference in the means, we often calculate a test statistic that is called the ***equal variance two-independent samples t-statistic***. The test statistic is

$$t = \frac{\bar{x}_1 - \bar{x}_2}{s_p \sqrt{\frac{1}{n_1} + \frac{1}{n_2}}},$$

where  $s_1^2$  and  $s_2^2$  are the sample variances for the two groups,  $n_1$  and  $n_2$  are the sample sizes for the two groups, and the ***pooled sample standard deviation***,

$$s_p = \sqrt{\frac{(n_1 - 1)s_1^2 + (n_2 - 1)s_2^2}{n_1 + n_2 - 2}}.$$

The  $t$ -statistic keeps the important comparison between the means in the numerator that we used before and standardizes (re-scales) that difference so that  $t$  will follow a  $t$ -distribution (a parametric “named” distribution) if certain assumptions are met. But first we should see if standardizing the difference in the means had an impact on our permutation test results. Instead of using the `diffmean` function, we will use the `t.test` function (see its full use below) and have it calculate the formula for  $t$  for us. The R code “\$statistic” is basically a way of extracting just the number we want to use for  $T$  from a larger set of output the `t.test` function wants to provide you. We will see below that `t.test` switches the order of the difference (now it is *Average - Unattractive*) – always carefully check for the direction of the difference in the results. Since we are doing a two-sided test, the code resembles the permutation test code in Section 2.4 with the new  $t$ -statistic replacing the difference in the sample means that we used before.

The permutation distribution in Figure 2.12 looks similar to the previous results with slightly different  $x$ -axis scaling. The proportion of permuted results that were more extreme than the observed result was 0.031. This difference is due to a different set of random permutations being selected. If you run permutation code, you will often get slightly different results each time you run it. If you are uncomfortable with the variation in the results, you can run more than  $B = 1,000$  permutations (say 10,000) and the variability in the resulting p-values will be reduced further. Usually this uncertainty will not cause any substantive problems – but do not be surprised if your results vary from a colleagues if you are both analyzing the same data set or if you re-run your permutation code.

```
par(mfrow=c(1,2))
Tobs <- t.test(Years ~ Attr, data=MockJury2, var.equal=T)$statistic
Tobs

##      t
## -2.17023
Tstar <- matrix(NA, nrow=B)
for (b in (1:B)){
  Tstar[b] <- t.test(Years ~ shuffle(Attr), data=MockJury2, var.equal=T)$statistic
}
pdata(abs(Tstar),abs(Tobs),lower.tail=F)

##      t
## 0.031
hist(Tstar, labels=T)
abline(v=c(-1,1)*Tobs, lwd=2, col="red")
plot(density(Tstar), main="Density curve of Tstar")
abline(v=c(-1,1)*Tobs, lwd=2, col="red")
```

Permutation distribution of the  $t$ -statistic Permutation distribution of the  $t$ -statistic

The parametric version of these results is based on using what is called the ***two-independent sample t-test***. There are actually two versions of this test, one that assumes that variances are equal in the groups and one that does not. There is a rule of thumb that if the **ratio of the larger standard deviation over the smaller standard deviation is less than 2, the equal variance procedure is OK**. It ends up that this assumption is less important if the sample sizes in the groups are approximately equal and more important if the groups contain different numbers of observations. In comparing the two potential test statistics, the procedure that assumes equal variances has a complicated denominator (see the formula above for  $t$  involving  $s_p$ ) but a simple formula for **degrees of freedom (df)** for the  $t$ -distribution ( $df = n_1 + n_2 - 2$ ) that approximates the distribution of the test statistic,  $t$ , under the null hypothesis. The procedure that assumes unequal variances has a simpler test statistic and a very complicated degrees of freedom formula. The equal variance procedure is most similar to the ANOVA methods we will consider in Chapters 2 and 3 so

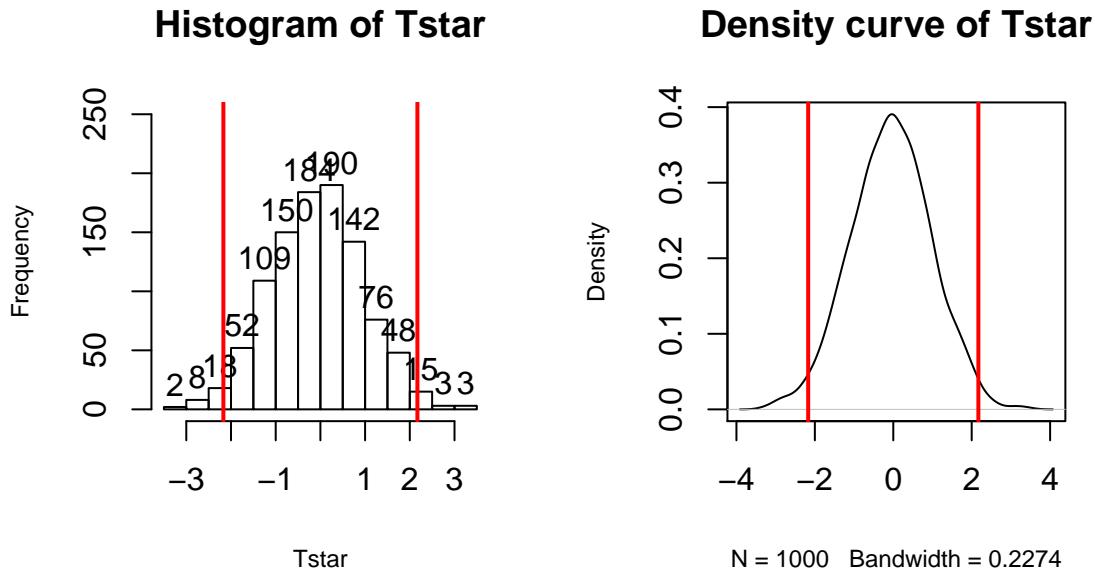


Figure 2.12: Permutation distribution of the  $t$ -statistic

that will be our focus for the two group problem. Fortunately, both of these methods are readily available in the `t.test` function in R if needed.

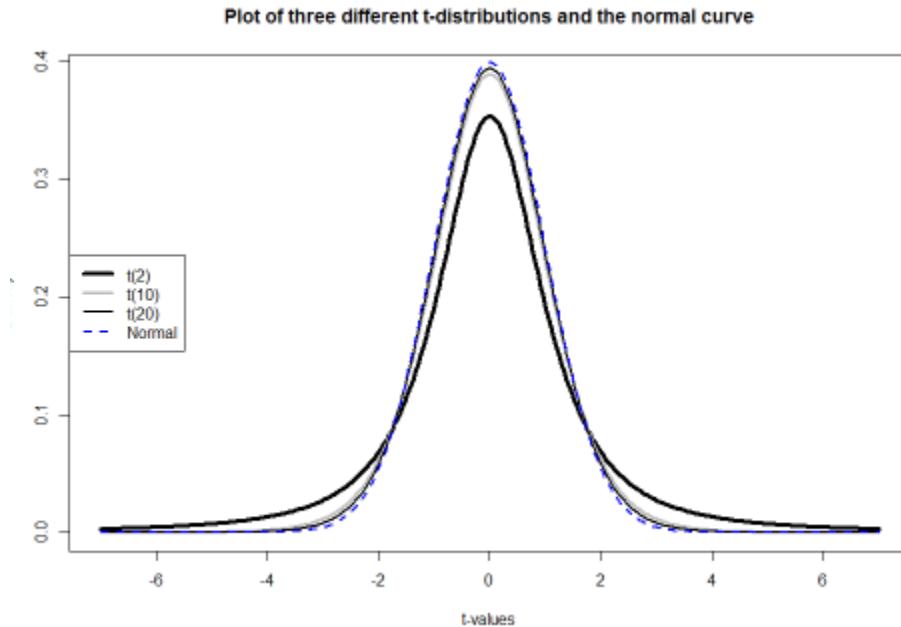
If the assumptions for the equal variance  $t$ -test are met and the null hypothesis is true, then the sampling distribution of the test statistic should follow a  $t$ -distribution with  $n_1 + n_2 - 2$  degrees of freedom. The  **$t$ -distribution** is a bell-shaped curve that is more spread out for smaller values of degrees of freedom as shown in Figure 2.13. The  $t$ -distribution looks more and more like a **standard normal distribution** ( $N(0, 1)$ ) as the degrees of freedom increase.

To get the p-value for the parametric  $t$ -test, we need to calculate the test statistic and  $df$ , then look up the areas in the tails of the  $t$ -distribution relative to the observed  $t$ -statistic. We'll learn how to use R to do this below, but for now we will allow the `t.test` function to take care of this for us. The `t.test` function uses our formula notation (`Years ~ Attr`) and then `data=...` as we saw before for making plots. To get the equal-variance test result, the `var.equal=T` option needs to be turned on. Then `t.test` provides us with lots of useful output. The three results we've been discussing are highlighted in the output below – the test statistic value (-2.17),  $df = 73$ , and the p-value, from the  $t$ -distribution with 73 degrees of freedom, of 0.033.

```
t.test(Years ~ Attr, data=MockJury2, var.equal=T)
```

```
## 
##  Two Sample t-test
##
## data: Years by Attr
## t = -2.1702, df = 73, p-value = 0.03324
## alternative hypothesis: true difference in means is not equal to 0
## 95 percent confidence interval:
## -3.5242237 -0.1500295
## sample estimates:
##      mean in group Average mean in group Unattractive
##                  3.973684                      5.810811
```

So the parametric  $t$ -test gives a p-value of 0.033 from a test statistic of -2.1702. The negative sign on the test statistic occurred because the function took *Average - Unattractive* which is the opposite direction as

Figure 2.13: Plots of  $t$  and normal distributions

`diffmean`. The p-value is very similar to the two permutation results found before. The reason for this similarity is that the permutation distribution with 73 degrees of freedom. Figure 2.14 shows how similar the two distributions happened to be here.

In your previous statistics course, you might have used an applet or a table to find p-values such as what was provided in the previous R output. When not directly provided in the output of a function, R can be used to look up p-values<sup>15</sup> from named distributions such as the  $t$ -distribution. In this case, the distribution of the test statistic under the null hypothesis is a  $t(73)$  or a  $t$  with 73 degrees of freedom. The `pt` function is used to get p-values from the  $t$ -distribution in the same manner that `pdata` could help us to find p-values from the permutation distribution. We need to provide the `df=...` and specify the tail of the distribution of interest using the `lower.tail` option along with the cutoff of interest. If we want the area to the left of -2.17:

```
pt(-2.1702, df=73, lower.tail=T)
```

```
## [1] 0.01662286
```

And we can double it to get the p-value that `t.test` provided earlier, because the  $t$ -distribution is symmetric:

```
2*pt(-2.1702, df=73, lower.tail=T)
```

```
## [1] 0.03324571
```

More generally, we could always make the test statistic positive using the absolute value, find the area to the right of it, and then double that for a two-sided test p-value:

```
2*pt(abs(-2.1702), df=73, lower.tail=T)
```

```
## [1] 1.966754
```

Permutation distributions do not need to match the named parametric distribution to work correctly, although this happened in the previous example. The parametric certain conditions to be met for the sampling distribution of the statistic to follow the named distribution and provide accurate p-values. The conditions for the equal variance t-test are:

---

<sup>15</sup>On exams, you will be asked to describe the area of interest, sketch a picture of the area of interest, and/or note the distribution you would use.

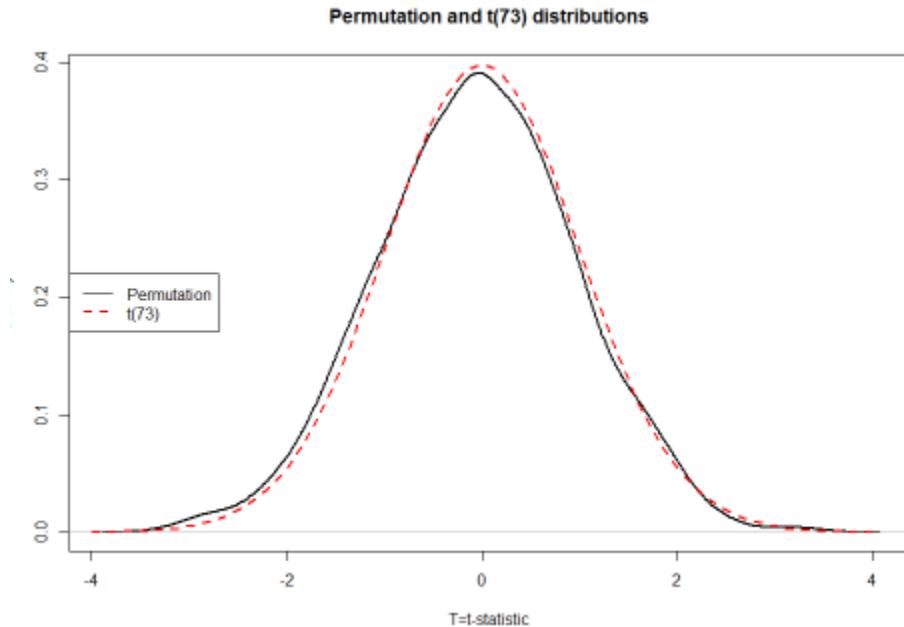


Figure 2.14: Plot of permutation and  $t$  distribution with  $df = 73$ .

1. **Independent observations:** Each observation obtained is unrelated to all other observations. To assess this, consider whether anything in the data collection might lead to clustered or related observations that are un-related to the differences in the groups. For example, was the same person measured more than once?<sup>16</sup>
2. **Equal variances** in the groups (because we used a procedure that assumes equal variances! – there is another procedure that allows you to relax this assumption if needed...). To assess this, compare the standard deviations and variability in the beanplots and see if they look noticeably different. Be particularly critical of this assessment if the sample sizes differ greatly between groups.
3. **Normal distributions** of the observations in each group. We'll learn more diagnostics later, but the boxplots and beanplots are a good place to start to help you look for skews or outliers, which were both present here. If you find skew and/or outliers, that would suggest a problem with the assumption of normality as normal distributions are symmetric and extreme observations occur very rarely.

For the permutation test, we relax the third condition and replace it with:

3. **Similar distributions between the groups:** The permutation approach allows valid inferences as long as the two groups have similar shapes and only possibly differ in their centers. In other words, the distributions need not look normal for the procedure to work well, but they do need to look similar.

In the prisoner “juror” study, we can assume that the independent observation condition is met because there is no information suggesting that the same subjects were measured more than once or that some other type of grouping in the responses was present (like the subjects were divided in groups and placed in rooms to discuss their responses prior to submitting them). The equal variance condition might be violated. The variances need not be equal as the procedure can still provide reasonable results with some violation of this assumption. The standard deviations are 2.8 vs 4.4, so this difference is not “large” according to the rule of thumb noted above. It is, however, close to being considered problematic. It would be difficult to reasonably assume that the normality condition is met here (Figure 2.6 with clear right skews in both groups and potential outliers which causes concerns for (3) for the parametric procedure. The shapes look similar for the two groups so

<sup>16</sup>In some studies, the same subject might be measured in both conditions and this violates the assumptions of this procedure.

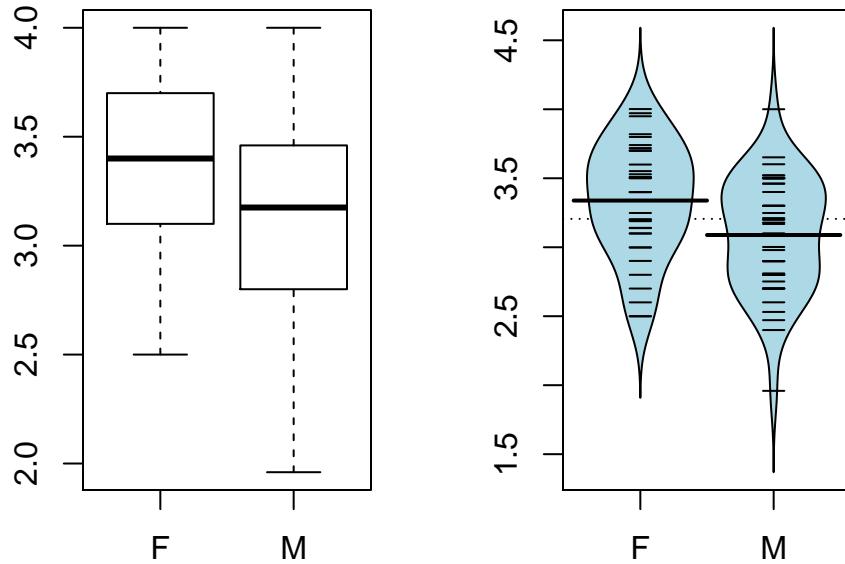


Figure 2.15: Side-by-side boxplot and beanplot of GPAs of STAT 217 students by sex.

there is less reason to be concerned with using the permutation approach based on its version of (3) above.

The permutation approach is resistant to impacts of violations of the normality assumption. It is not resistant to impact of violations of any of the other assumptions. In fact, it can be quite sensitive to unequal variances as it will detect differences in the variances of the groups instead of differences in the means. Its scope of inference is the same as the parametric approach and can lead to similarly inaccurate conclusions in the presence of non-independent observations as for the parametric approach. In this example, we discover that parametric and permutation approaches provide very similar inferences.

## 2.7 Second example of permutation tests

In every chapter, we will follow the first example used to motivate and explain the methods with a “worked” example where we focus just on the results. In a previous semester, some of the STAT 217 students ( $n=79$ ) provided information on their *Sex*, *Age*, and current cumulative *GPA*. We might be interested in whether Males and Females had different average GPAs. First, we can take a look at the difference in the responses by groups based on the output and as displayed in Figure 2.15.

```
s217 <- read.csv("http://www.math.montana.edu/courses/s217/documents/s217.csv")
```

```
require(mosaic)
```

```
require(beanplot)
```

```
mean(GPA~Sex, data=s217)
```

```
##          F          M
```

```
## 3.338378 3.088571
```

```
favstats(GPA~Sex, data=s217)
```

```
##   Sex min  Q1 median   Q3 max      mean       sd n missing
```

```
## 1   F 2.50 3.1  3.400 3.70    4 3.338378 0.4074549 37      0
```

```
## 2   M 1.96 2.8  3.175 3.46    4 3.088571 0.4151789 42      0
```

```
par(mfrow=c(1,2))
boxplot(GPA~Sex, data=s217)
beanplot(GPA~Sex, data=s217, log="", col="lightblue", method="jitter")
```

In these data, the distributions of the GPAs look to be left skewed but maybe not as dramatically as the responses were right-skewed in the previous example. The Female GPAs look to be slightly higher than for Males (0.25 GPA difference in the means) but is that a “real” difference? We need our inference tools to more fully assess these differences.

```
diffmean(GPA~Sex, data=s217)
```

```
## diffmean
## -0.2498069
```

First, we can try the parametric approach:

```
t.test(GPA~Sex, data=s217, var.equal=T)
```

```
##
##  Two Sample t-test
##
## data: GPA by Sex
## t = 2.6919, df = 77, p-value = 0.008713
## alternative hypothesis: true difference in means is not equal to 0
## 95 percent confidence interval:
##  0.06501838 0.43459552
## sample estimates:
## mean in group F mean in group M
##          3.338378      3.088571
```

So the test statistic was observed to be  $t = 2.69$  and it hopefully follows a  $t(77)$  distribution under the null hypothesis. This provides a p-value of 0.008713 that we can trust if all the conditions to use this procedure are met. Compare these results to the permutation approach, which relaxes that normality assumption, with the results that follow. In the permutation test,  $T = 2.692$  and the p-value is 0.005 which is a little smaller than the result provided by the parametric approach. The agreement of the two approaches, again, provides some re-assurance about the use of either approach.

```
Tobs <- t.test(GPA~Sex, data=s217, var.equal=T)$statistic
Tstar <- matrix(NA, nrow=B)
for (b in (1:B)){
  Tstar[b] <- t.test(GPA~shuffle(Sex), data=s217, var.equal=T)$statistic
}
pdata(abs(Tstar),abs(Tobs),lower.tail=F)
par(mfrow=c(1,2))
hist(Tstar,labels=T)
abline(v=c(-1,1)*Tobs, lwd=2, col="red")
plot(density(Tstar), main="Density curve of Tstar")
abline(v=c(-1,1)*Tobs, lwd=2, col="red")
```

Here is a full write-up of the results using all 6+ hypothesis testing steps, using the permutation results:

0. *Isolate the claim to be proved and method to use (define a test statistic  $T$ )* We want to test for a difference in the means between males and females and will use the equal-variance two-sample t-test statistic to compare them, making a decision at the 5% significance level.
1. Write the null and alternative hypotheses
  - $H_0 : \mu_{male} = \mu_{female}$ 
    - where  $\mu_{male}$  is the true mean GPA for males and  $\mu_{female}$  is true mean GPA for females.
  - $H_A : \mu_{male} \neq \mu_{female}$

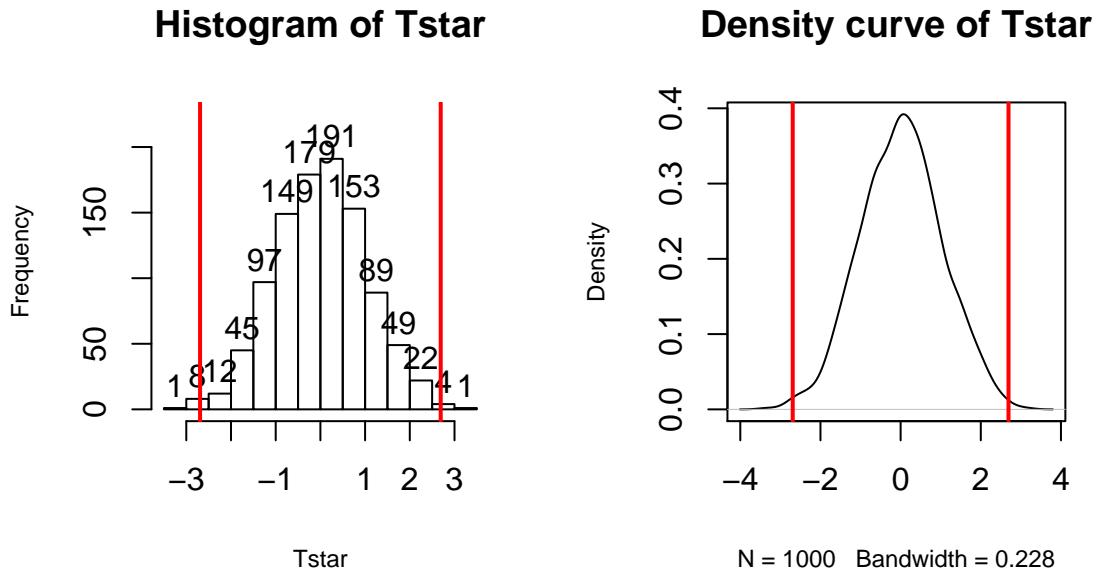


Figure 2.16: Histogram and density curve of permutation distribution of test statistic for STAT 217 GPAs.

2. Check conditions for the procedure being used

- **Independent observations condition:** It appears that this assumption is met because there is no reason to assume any clustering or grouping of responses that might create dependence in the observations. The only possible consideration is that the observations were taken from different sections and there could be some differences between the sections. However, for overall GPA this is not likely to be a big issue. The only way this could create a violation here is if certain sections tended to attract students with different GPA levels (such as the 9 am section had the best/worst GPA students...).
- **Equal variance condition :** There is a small difference in the range of the observations in the two groups but the standard deviations are very similar so there is no evidence that this condition is violated.
- **Similar distribution condition:** Based on the side-by-side boxplots and beanplots, it appears that both groups have slightly left-skewed distributions, which could be problematic for the parametric approach, but the permutation approach condition is not violated since the distributions look to have fairly similar shapes.

3. Find the value of the appropriate test statistic

- $T = 2.69$  from the previous R output.

4. Find the p-value

- p-value=0.005 from the permutation distribution results.
- This means that there is about a 0.5% chance we would observe a difference in mean GPA (female-male or male-female) of 0.25 points or more if there is in fact no difference in true mean GPA between females and males in STAT 217 in a particular semester.

### 5. Decision

- Since the p-value is “small” (*a priori* 5% significance level selected), we can reject the null hypothesis.

### 6. Conclusion and scope of inference, specific to the problem

- There is strong evidence against the null hypothesis of no difference in the true mean GPA between males and females for the STAT 217 students in this semester and so we conclude that there is evidence of a difference in the mean GPAs between males and females in STAT 217 students.
- Because this was not a randomized experiment, we can’t say that the difference in sex causes the difference in mean GPA and because it was not a random sample from a larger population, our inferences only pertain the STAT 217 students that responded to the survey in that semester.

## 2.8 Confidence intervals and bootstrapping

Randomly shuffling the treatments between the observations is like randomly sampling the treatments without replacement. In other words, we randomly sample one observation at a observations. This provides us with a technique for testing hypotheses because it provides new splits of the observations into groups that are as interesting as what we observed if the null hypothesis is assumed true. In most situations, we also want to estimate parameters of interest and provide ***confidence intervals*** for those parameters (an interval where we are   % ***confident*** that the true parameter lies). As before, there are two options we will consider – a parametric and a nonparametric approach. The nonparametric approach will be using what is called ***bootstrapping*** and draws its name from “pull yourself up by your bootstraps” where you improve your situation based on your own efforts. In statistics, we make our situation or inferences better by re-using the observations we have by assuming that the sample represents the population. Since each observation represents other similar observations in the population that we didn’t get to measure, if we ***sample with replacement*** to generate a new data set of size  $n$  from our data set (also of size  $n$ ) it mimics the process of taking our population of interest. This process also ends up giving us useful sampling distributions of statistics even when our standard normality assumption is violated, similar to what we encountered in the permutation tests. Bootstrapping is especially useful in situations where we are interested in statistics other than the mean (say we want a confidence interval for a median or a standard deviation) or when we consider functions of more than one parameter and don’t want to derive the distribution of the statistic (say the difference in two medians). In this text, bootstrapping is used to provide more trustworthy inferences when some of our assumptions (especially normality) might be violated for our parametric procedure.

To perform bootstrapping, we will use the `resample` function from the `mosaic` package. We can apply this function to a data set and get a new version of the data set by sampling new observations *with replacement* from the original one. The new, bootstrapped version of the data set (called `MockJury_BTS` below) contains a new variable called `orig.id` which is the number of the subject from the original data set. By summarizing how often each of these id’s occurred in a bootstrapped data set, we can see how the re-sampling works. The `table` function will count up how many times each observation was used in the bootstrap sample, providing a row with the id followed by a row with the count<sup>17</sup>. In the first bootstrap sample shown, the 2nd, 7th, and 9th observations were sampled one time each, the 4th observation was sampled three times, and the 1st, 3rd, 5th, and many others were not sampled at all. Bootstrap sampling thus picks some observations multiple times and to do that it has to ignore some observations.

---

<sup>17</sup>The `as.numeric` function is also used here. It really isn’t important but makes sure the output of `table` is sorted by observation number by first converting the `orig.id` variable into a numeric vector.

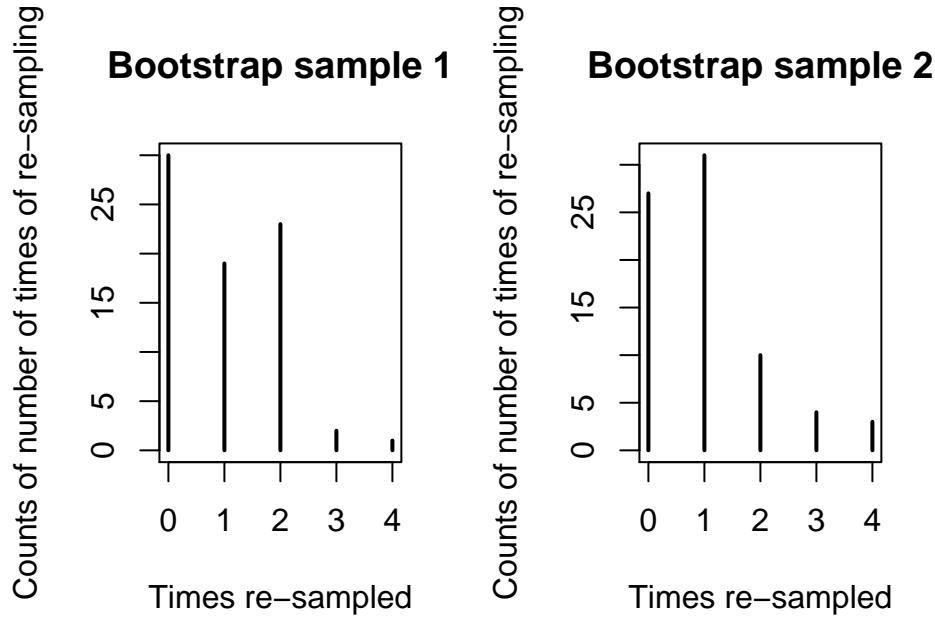


Figure 2.17: Counts of number of times of observation (or not observed for times re-sampled of 0) for two bootstrap samples.

```
MockJury_BTS <- resample(MockJury2)
table(as.numeric(MockJury_BTS$orig.id))
```

```
##
##  1  2  3  4  5  6 10 11 12 14 15 17 18 19 20 22 24 26 29 30 32 35 36 37 39
##  1  2  2  1  3  2  1  2  1  1  3  1  2  1  2  1  2  1  2  2  1  2  2  2  1  2  2
## 40 42 43 44 45 46 47 48 49 55 58 59 60 61 69 70 71 72 74 75
##  2  1  1  4  2  2  1  2  1  2  1  1  2  2  2  2  2  2  1  1  1
```

Like in permutations, one randomization isn't enough. A second bootstrap sample is also provided to help you get a sense of what it is doing to generate a data set. It did not select subject 7 but did select 2, 4, 6, and 8 two times. You can see other variations in the resulting re-sampling of subjects with the most sampled subject being the chance of selecting any observation for any slot in the new data set is  $1/75$  and the expected or mean number of appearances we expect to see for an observation is the number of tries times the probability of selection on each so  $75 * 1/75 = 1$ .

```
MockJury_BTS2 <- resample(MockJury2)
table(as.numeric(MockJury_BTS2$orig.id))
```

```
##
##  1  2  3  5  6  8 11 12 13 14 15 18 19 20 21 23 24 26 27 28 29 31 32 34 36
##  1  1  1  1  4  1  1  1  1  3  1  1  1  1  3  2  2  1  1  1  2  1  2  1  2
## 37 38 40 42 46 48 50 51 52 56 58 59 61 62 63 66 67 68 69 72 73 74 75
##  1  2  1  1  1  1  2  4  1  1  3  2  1  1  1  1  1  2  3  1  4  2
```

We can use the two results to get an idea of distribution of results in terms of number of times observations might be re-sampled when sampling with replacement and the variation in those results, as shown in Figure 2.17. We could also derive the expected counts for each number of times of re-sampling when we start with all observations having an equal chance and sampling with replacement but this isn't important for using bootstrapping methods.

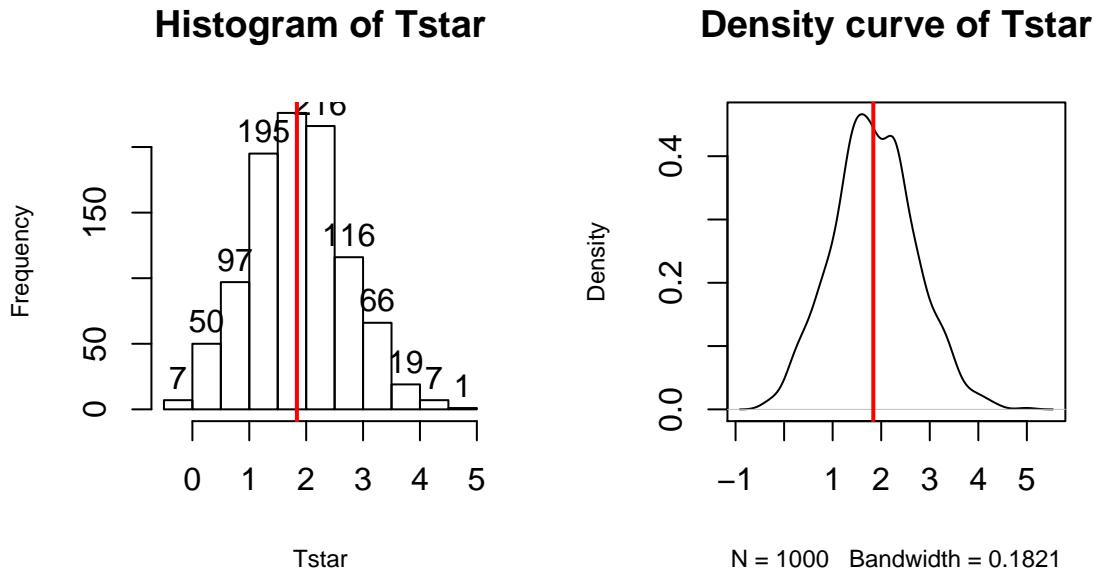


Figure 2.18: Histogram and density curve of bootstrap distributions of difference in sample mean `Years` with vertical line for the observed difference in the means of 1.84 years.

The main point of this exploration was to see that each run of the `resample` function provides a new version of the data set. Repeating this  $B$  times using another `for` loop, we will track our quantity of interest, say  $T$ , in all these new “data sets” and call those results  $T^*$ . The distribution of the bootstrapped  $T^*$  statistics will tell us about the range of results to expect for the statistic and the middle  $\underline{\hspace{2cm}}$ % of the  $T^*$ ’s provides a **bootstrap confidence interval**<sup>18</sup> for the true parameter – here the *difference in the two population means*.

To make this concrete, we can revisit our previous examples, starting with the `MockJury2` data created before and our interest in comparing the mean sentences for the *Average* and *Unattractive* picture groups. The bootstrapping code is very similar to the permutation code except that we apply the `resample` function to the entire data set as opposed to the `shuffle` function being applied to the explanatory variable.

```
par(mfrow=c(1,2))
Tobs <- diffmean(Years ~ Attr, data=MockJury2); Tobs
```

```
## diffmean
## 1.837127
B <- 1000
Tstar <- matrix(NA,nrow=B)
for (b in (1:B)){
  Tstar[b] <- diffmean(Years ~ Attr, data=resample(MockJury2))
}
favstats(Tstar)

##      min      Q1 median      Q3      max      mean        sd      n
## -0.3627312 1.305773 1.833091 2.385281 4.988756 1.854428 0.8438987 1000
## missing
## 0
```

<sup>18</sup>There are actually many ways to use this information to make a confidence interval. We are using the simplest method that is called the “percentile” method.

```
hist(Tstar, labels=T)
abline(v=Tobs, col="red", lwd=2)
plot(density(Tstar), main="Density curve of Tstar")
abline(v=Tobs, col="red", lwd=2)
```

In this situation, the observed difference in the mean sentences is 1.84 years (Unattractive-Average), which is the vertical line in Figure 2.18. The bootstrap distribution shows the results for the difference in the sample means when fake data sets are re-constructed by sampling from the data set with replacement. The bootstrap distribution is approximately centered at the observed value (difference in the sample means) and is relatively symmetric.

The permutation distribution in the same situation (Figure 2.12) had a similar shape but was centered at 0. Permutations create sampling distributions based on assuming the null hypothesis is true, which is useful for hypothesis testing. Bootstrapping creates distributions centered at the observed result, which is the sampling distribution “under the alternative” or when no null hypothesis is assumed; bootstrap distributions are useful for generating confidence intervals for the true parameter values.

To create a 95% bootstrap confidence interval for the difference in the true mean sentences ( $\mu_{Unattr} - \mu_{Avg}$ ), select the middle 95% of results from the bootstrap distribution. Specifically, find the 2.5th percentile and the 97.5th percentile (values that put 2.5 and 97.5% of the results to the left) in the bootstrap distribution, which leaves 95% in the middle for the confidence interval. To find percentiles in a distribution in R, functions are of the form `q[Name of distribution]`, with the function `qt` extracting percentiles from a *t*-distribution (examples below). From the bootstrap results, use the `qdata` function on the `Tstar` results that contain the bootstrap distribution of the statistic of interest.

```
qdata(Tstar, 0.025)
```

```
##      p quantile
## 0.0250000 0.2414232
qdata(Tstar, 0.975)

##      p quantile
## 0.9750000 3.521528
```

These results tell us that the 2.5th percentile of the bootstrap distribution is at 0.26 years and the 97.5th percentile is at 3.50 years. We can combine these results to provide a 95% confidence for  $\mu_{Unattr} - \mu_{Avg}$  that is between 0.26 and 3.50. We can interpret this as with any confidence interval, that we are 95% confident that the difference in the true mean suggested sentences (Unattractive minus Average group) is between 0.26 and 3.50 years. We can also obtain both percentiles in one line of code using:

```
quantiles <- qdata(Tstar, c(0.025,0.975))
quantiles
```

```
##      quantile      p
## 2.5%  0.2414232 0.025
## 97.5% 3.5215278 0.975
```

Figure 2.19 displays those same percentiles on the bootstrap distribution residing in `Tstar`.

```
par(mfrow=c(1,2))
hist(Tstar, labels=T)
abline(v=quantiles$quantile, col="blue", lwd=3)
plot(density(Tstar), main="Density curve of Tstar")
abline(v=quantiles$quantile, col="blue", lwd=3)
```

Although confidence intervals can exist without referencing hypotheses, we can revisit our previous  $H_0 : \mu_{Unattr} = \mu_{Avg}$ . This null hypothesis is equivalent to testing  $H_0 : \mu_{Unattr} - \mu_{Avg} = 0$ , that the difference in the true means is equal to 0 years. And the difference in the means was the scale for our confidence interval, which did not contain 0 years. We will call 0 an interesting **reference value** for the confidence interval, because here it is the value where the true means are equal to each other (have a difference of 0 years). In

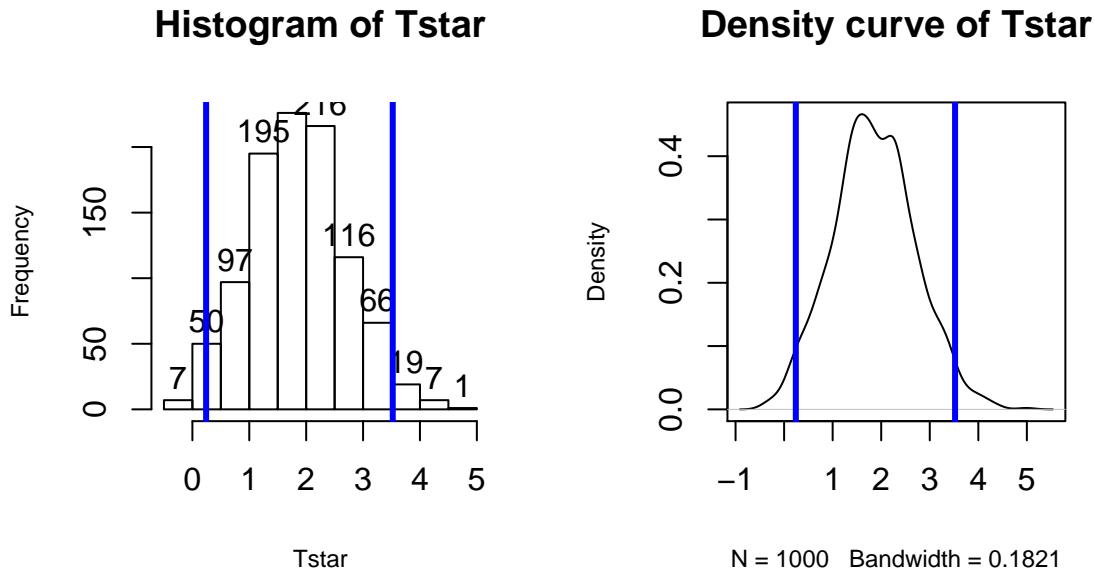


Figure 2.19: Histogram and density curve of bootstrap distribution with 95% bootstrap confidence intervals displayed (vertical lines).

general, if our confidence interval does not contain 0, then it is saying that 0 is not one of our likely values for the difference in the true means. This implies that we should reject a claim that they are equal. This provides the same inferences for the hypotheses that we considered previously using both a parametric and permutation approach.

The general summary is that we can use confidence intervals to test hypotheses by assessing whether the reference value under the null hypothesis is in the confidence interval (FTR  $H_0$ ) or outside the confidence interval (Reject  $H_0$ ). P-values are more informative about hypotheses but confidence intervals are more informative about the size of differences, so both offer useful information and, as shown here, can provide consistent conclusions about hypotheses.

As in the previous situation, we also want to consider the parametric approach for comparison purposes and to have that method available, especially to help us understand some methods where we will only consider parametric inferences in later chapters. The parametric confidence interval is called the ***equal variance, two-sample t confidence interval*** and assumes that the populations being sampled from are normally distributed and leads to using a *t*-distribution to form the interval. The output from the `t.test` function provides the parametric 95% confidence interval calculated for you:

```
t.test(Years ~ Attr, data=MockJury2, var.equal=T)
```

```
##
##  Two Sample t-test
##
## data: Years by Attr
## t = -2.1702, df = 73, p-value = 0.03324
## alternative hypothesis: true difference in means is not equal to 0
## 95 percent confidence interval:
## -3.5242237 -0.1500295
## sample estimates:
##      mean in group Average mean in group Unattractive
##                  3.973684                      5.810811
```

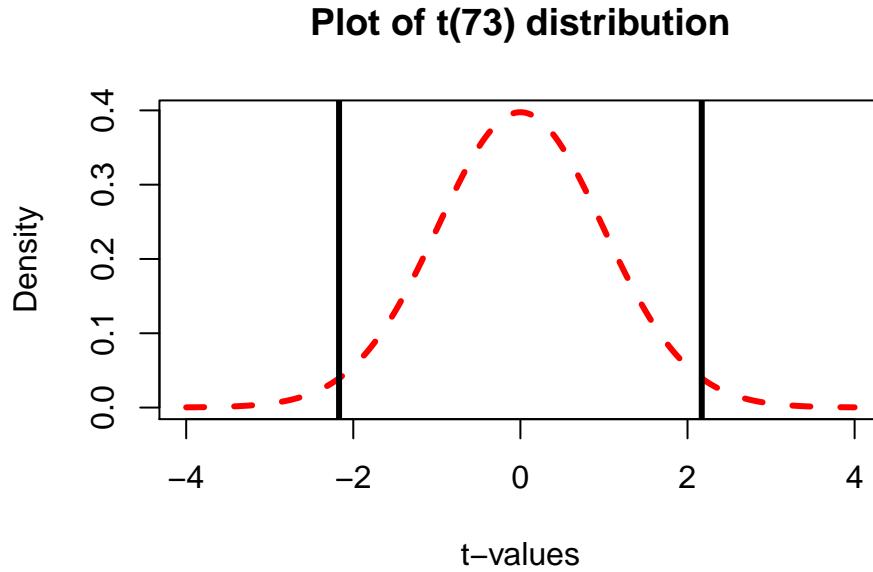


Figure 2.20: Plot of  $t(73)$  with cut-offs for putting 95% of distributions in the middle.

The `t.test` function again switched the order of the groups and provides slightly different end-points than our bootstrap confidence interval (both are made at the 95% confidence level though), which was slightly narrower. Both intervals have the same interpretation, only the methods for calculating the intervals and the assumptions differ. Specifically, the bootstrap interval can tolerate different distribution shapes other than normal and still provide intervals that work well<sup>19</sup>. The other assumptions are all the same as for the hypothesis test, where we continue to assume that we have independent observations with equal variances for the two groups.

The formula that `t.test` is using to calculate the parametric *equal-variance two-sample t confidence interval* is:

$$\bar{x}_1 - \bar{x}_2 \mp t_{df}^* s_p \sqrt{\frac{1}{n_1} + \frac{1}{n_2}}$$

In this situation, the  $df$  is again  $n_1 + n_2 - 2$  and  $s_p = \sqrt{\frac{(n_1-1)s_1^2 + (n_2-1)s_2^2}{n_1+n_2-2}}$ . The  $t_{df}^*$  is a multiplier that comes from finding the percentile from the  $t$ -distribution that puts  $C\%$  in the middle of the distribution with  $C$  being the confidence level. It is important to note that this  $t^*$  has nothing to do with the previous test statistic  $t$ . It is confusing and many of you will, at some point, happily take the result from a test statistic calculation and use it for a multiplier in a  $t$ -based confidence interval. Figure 2.20 shows the  $t$ -distribution with 73 degrees of freedom and the cut-offs that put 95% of the area in the middle.

```
par(mfrow=c(1,1))
x<-seq(from=-4,to=4,length.out=200)
plot(x,dt(x,df=73),col="red",lty=2,lwd=3,type="l",xlab="t-values",ylab="Density",
      main="Plot of t(73) distribution" )
abline(v=-2.1702,lwd=3)
abline(v=2.1702,lwd=3)
```

<sup>19</sup>When hypothesis tests “work well” they have high power to detect differences while having Type I error rates that are close to what we choose a priori. When confidence intervals “work well”, they contain the true parameter value in repeated random samples at around the selected confidence level

For 95% confidence intervals, the multiplier is going to be close to 2 and anything else is a sign of a mistake. We can use R to get the multipliers for confidence intervals using the `qt` function in a similar fashion to how `qdata` was used in the bootstrap results, except that this new value must be used in the previous confidence interval formula. This function produces values for requested percentiles, so if we want to put 95% in the middle, we place 2.5% in each tail of the distribution and need to request the 97.5th percentile. Because the *t*-distribution is always symmetric around 0, we merely need to look up the value for the 97.5th percentile and know that the multiplier for the 2.5th percentile is just  $-t^*$ . The  $t^*$  multiplier to form the confidence interval is 1.993 for a 95% confidence interval when the  $df = 73$  based on the results from `qt`:

```
qt(0.975, df=73)
```

```
## [1] 1.992997
```

Note that the 2.5th percentile is just the negative of this value due to symmetry and the real source of the minus in the minus/plus in the formula for the confidence interval.

```
qt(0.025, df=73)
```

```
## [1] -1.992997
```

We can also re-write the confidence interval formula into a slightly more general form as

$$\bar{x}_1 - \bar{x}_2 \mp t_{df}^* SE_{\bar{x}_1 - \bar{x}_2} \quad \text{OR} \quad \bar{x}_1 - \bar{x}_2 \mp ME$$

where  $SE_{\bar{x}_1 - \bar{x}_2} = s_p \sqrt{\frac{1}{n_1} + \frac{1}{n_2}}$  and  $ME = t_{df}^* SE_{\bar{x}_1 - \bar{x}_2}$ . In some situations, researchers will report the **standard error** (SE) or **margin of error** (ME) as a method of quantifying the uncertainty in a statistic. The SE is an estimate of the standard deviation of the statistic (here  $\bar{x}_1 - \bar{x}_2$ ) and the ME is an estimate of the precision of a statistic that can be used to directly form a confidence interval. The ME depends on the choice of confidence level although 95% is almost always selected.

To finish this example, R can be used to help you do calculations much like a calculator except with much more power “under the hood”. You have to make sure you are careful with using `( )` to group items and remember that the asterisk (\*) is used for multiplication in R. We need the pertinent information which is available from the `favstats` output repeated below to calculate the confidence interval “by hand” using R.

```
favstats(Years~Attr, data=MockJury2)
```

```
##          Attr min Q1 median Q3 max      mean       sd   n missing
## 1      Average  1  2      3  5 12 3.973684 2.823519 38      0
## 2 Unattractive  1  2      5 10 15 5.810811 4.364235 37      0
```

Start with typing the following command to calculate  $s_p$  and store it in a variable named `sp`:

```
sp <- sqrt(((38-1)*(2.8235^2)+(37-1)*(4.364^2))/(38+37-2))
```

```
sp
```

```
## [1] 3.665036
```

Then calculate the confidence interval that `t.test` provided using:

```
3.974-5.811+c(-1,1)*qt(.975,df=73)*sp*sqrt(1/38+1/37)
```

```
## [1] -3.5240302 -0.1499698
```

The previous code uses `c(-1, 1)` times the margin of error to subtract and add the ME to the difference in the sample means ( $3.974 - 5.811$ ), which generates the lower and then upper bounds of the confidence interval. If desired, we can also use just the last portion of the previous calculation to find the margin of error, which is 1.69 here.

```
qt(.975,df=73)*sp*sqrt(1/38+1/37)
```

```
## [1] 1.68703
```

## 2.9 Bootstrap confidence intervals for difference in GPAs

We can now apply the new confidence interval methods on the STAT 217 grade data. This time we start with the parametric 95% confidence interval “by hand” in R and then use `t.test` to verify our result. The `favstats` output provides us with the required information to calculate the confidence interval:

```
favstats(GPA~Sex,data=s217)
```

```
##   Sex min Q1 median Q3 max   mean      sd n missing
## 1   F 2.50 3.1 3.400 3.70 4 3.338378 0.4074549 37      0
## 2   M 1.96 2.8 3.175 3.46 4 3.088571 0.4151789 42      0
```

The  $df$  are  $37 + 42 - 2 = 77$ . Using the SDs from the two groups and their sample sizes, we can calculate  $s_e$ :

```
sp <- sqrt(((37-1)*(0.4075^2)+(42-1)*(0.41518^2))/(37+42-2))
```

```
sp
```

```
## [1] 0.4116072
```

The margin of error is:

```
qt(.975,df=77)*sp*sqrt(1/37+1/42)
```

```
## [1] 0.1847982
```

All together, the 95% confidence interval is:

```
3.338-3.0886+c(-1,1)*qt(.975,df=77)*sp*sqrt(1/37+1/42)
```

```
## [1] 0.0646018 0.4341982
```

So we are 95% confident that the difference in the true mean GPAs between females and males (females minus males) is between 0. 065 and 0. 434 GPA points. We get a similar<sup>20</sup> result from the `t.test` output:

```
t.test(GPA~Sex,data=s217,var.equal=T)
```

```
##
##  Two Sample t-test
##
## data:  GPA by Sex
## t = 2.6919, df = 77, p-value = 0.008713
## alternative hypothesis: true difference in means is not equal to 0
## 95 percent confidence interval:
##  0.06501838 0.43459552
## sample estimates:
## mean in group F mean in group M
##          3.338378          3.088571
```

Note that we can easily switch to 90% or 99% confidence intervals by simply changing the percentile in `qt` or changing `conf.level` in the `t.test` function. In the following two lines of code, we added octothorpes<sup>21</sup>(#) and then some text after function calls to explain what is being calculated. In computer code, octothorpes provide a way of adding comments that tell the software (here R) to ignore any text after a “#” on a given line. In the color version of the text, comments are also clearly distinguished.

```
qt(.95,df=77) # For 90% confidence and 77 df
```

```
## [1] 1.664885
```

```
qt(.995,df=77) #For 99% confidence and 77 df
```

```
## [1] 2.641198
```

<sup>20</sup>We rounded the means a little and that caused the small difference in results.

<sup>21</sup>You can correctly call octothorpes *number* symbols or, in the twitter verse, *hashtags*. For more on this symbol, see “<http://blog.dictionary.com/octothorpe/>”. I usually call them number symbols too.

```
t.test(GPA~Sex,data=s217,var.equal=T,conf.level=0.90)

##
## Two Sample t-test
##
## data: GPA by Sex
## t = 2.6919, df = 77, p-value = 0.008713
## alternative hypothesis: true difference in means is not equal to 0
## 90 percent confidence interval:
## 0.09530553 0.40430837
## sample estimates:
## mean in group F mean in group M
## 3.338378 3.088571
t.test(GPA~Sex,data=s217,var.equal=T,conf.level=0.99)

##
## Two Sample t-test
##
## data: GPA by Sex
## t = 2.6919, df = 77, p-value = 0.008713
## alternative hypothesis: true difference in means is not equal to 0
## 99 percent confidence interval:
## 0.004703598 0.494910301
## sample estimates:
## mean in group F mean in group M
## 3.338378 3.088571
```

As a review of some basic ideas with confidence intervals make sure you can answer the following questions:

1. What is the impact of increasing the confidence level in this situation?
2. What happens to the width of the confidence interval if the size of the SE increases or decreases?
3. What about increasing the sample size – should that increase or decrease the width of the interval?

All the general results you learned before about impacts to widths of CIs hold in this situation whether we are considering the parametric or bootstrap methods...

To finish this example, we will generate the comparable bootstrap 90% confidence interval using the bootstrap distribution in Figure 2.21.

```
Tobs <- diffmean(GPA ~ Sex, data=s217); Tobs
```

```
## diffmean
## -0.2498069
par(mfrow=c(1,2))
B<- 1000
Tstar<-matrix(NA,nrow=B)
for (b in (1:B)){
  Tstar[b]<-diffmean(GPA ~ Sex, data=resample(s217))
}
qdata(Tstar,.05)

##          p    quantile
## 0.0500000 -0.4032273
qdata(Tstar,.95)

##          p    quantile
## 0.9500000 -0.09521925
```

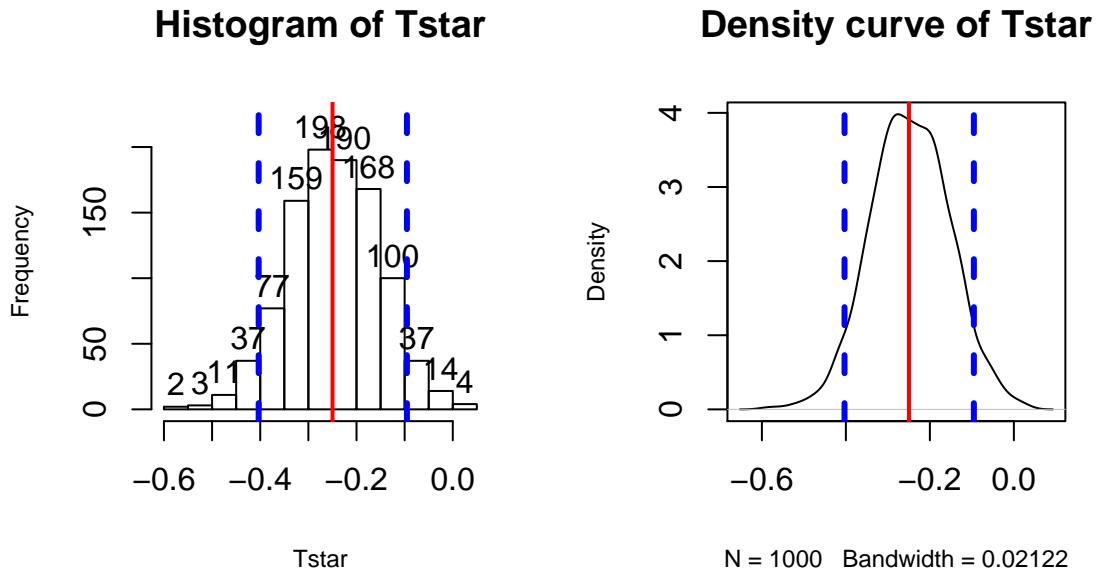


Figure 2.21: Histogram and density curve of bootstrap distribution of difference in sample mean GPAs (male minus female) with observed difference (solid vertical line) and quantiles that delineate the 90% confidence intervals (dashed vertical lines).

```
quantiles<-qdata(Tstar,c(.05,.95))
quantiles
```

```
##      quantile   p
## 5%  -0.40322729 0.05
## 95% -0.09521925 0.95
```

The output tells us that the 90% confidence interval is from -0.393 to -0.094 GPA points. The bootstrap distribution with the observed difference in the sample means and these cut-offs is displayed in Figure 2.21 using this code:

```
par(mfrow=c(1,2))
hist(Tstar,labels=T)
abline(v=Tobs,col="red",lwd=2)
abline(v=quantiles$quantile,col="blue",lwd=3,lty=2)
plot(density(Tstar),main="Density curve of Tstar")
abline(v=Tobs,col="red",lwd=2)
abline(v=quantiles$quantile,col="blue",lwd=3,lty=2)
```

In the previous output, the parametric 90% confidence interval is from 0.095 to 0.404, suggesting similar results again from the two approaches once you account for the two different orders of differencing of the groups. Based on the bootstrap CI, we can say that we are 90% confident that the difference in the true mean GPAs for STAT 217 students is between -0.393 to -0.094 GPA points (male minus females). Because sex cannot be assigned to the subjects, we cannot infer that sex is causing this difference and because this was a voluntary response sample of STAT 217 students in a given semester, we cannot infer that a difference of this size would apply to all STAT 217 students or even students in another semester.

Throughout the semester, pay attention to the distinctions between parameters and statistics, focusing on the differences between estimates based on the sample and inferences for the population of interest in the form of the parameters of interest. Remember that statistics are summaries of the sample information and

parameters are characteristics of populations (which we rarely know). And that our inferences are limited to the population that we randomly sampled from, if we randomly sampled.

## 2.10 Chapter summary

In this chapter, we reviewed basic statistical inference methods in the context of a two-sample mean problem. You were introduced to using R to do permutation testing and generate bootstrap confidence intervals as well as obtaining parametric *t*-test and confidence intervals in this same situation. You should have learned how to use a `for` loop for doing the nonparametric inferences and the `t.test` function for generating parametric inferences. In the two examples considered, the parametric and nonparametric methods provided similar results, suggesting that the assumptions were at least close to being met for the parametric procedures. When parametric and nonparametric approaches disagree, the nonparametric methods are likely to be more trustworthy since they have less restrictive assumptions but can still have problems.

When the noted conditions are not met in a hypothesis testing situation, the Type I error rates can be inflated, meaning that we reject the null hypothesis more often than we have allowed to occur by chance. Specifically, we could have a situation where our assumed 5% significance level test might actually reject the null when it is true 20% of the time. If this is occurring, we call a procedure *liberal* (it rejects too easily) and if the procedure is liberal, how could we trust a small p-value to be a “real” result and not just an artifact of violating the assumptions of the procedure? Likewise, for confidence intervals we hope that our 95% confidence level procedure, when repeated, will contain the true parameter 95% of the time. If our assumptions are violated, we might actually have an 80% confidence level procedure and it makes it hard to trust the reported results for our observed data set. Statistical inference relies on a belief in the methods underlying our inferences. If we don’t trust our assumptions, we shouldn’t trust the conclusions to perform the way we want them to. As sample sizes increase and/or violations of conditions lessen, then the procedures will perform better. In Chapter 3, we’ll learn some new tools for doing diagnostics to help us assess how much those conditions are violated.

## 2.11 Summary of important R code

The main components of R code used in this chapter follow with components to modify in red, remembering that any R packages mentioned need to be installed and loaded for this code to have a chance of working:

- `summary(DATASETNAME)`
  - Provides numerical summaries of all variables in the data set.
- `t.test(Y ~ X, data=DATASETNAME, conf.level=0.95)`
  - Provides two-sample t-test test statistic, df, p-value, and 95% confidence interval.
- `2*pt(abs(Tobs), df=DF, lower.tail=F)`
  - Finds the two-sided test p-value for an observed 2-sample t-test statistic of `Tobs`.
- `hist(DATASETNAME$Y)`
  - Makes a histogram of a variable named Y from the data set of interest.
- `boxplot(Y~X, data=DATASETNAME)`
  - Makes a boxplot of a variable named Y for groups in X from the data set.
- `beanplot(Y~X, data=DATASETNAME)`
  - Requires the `beanplot` package is loaded.
  - Makes a beanplot of a variable named Y for groups in X from the data set.

- `mean(Y~X, data=DATASETNAME); sd(Y~X, data=DATASETNAME)`
  - This usage of `mean` and `sd` requires the `mosaic` package.
  - Provides the mean and sd of responses of Y for each group described in X.
- `favstats(Y~X, data=DATASETNAME)`
  - Provides numerical summaries of Y by groups described in X.
- ```
Tobs <- t.test(Y~X, data=DATASETNAME, var.equal=T)$statistic; Tobs
B <- 1000
Tstar <- matrix(NA, nrow=B)
for (b in (1:B)){
  Tstar[b] <- t.test(Y~shuffle(X), data=DATASETNAME, var.equal=T)$statistic
}
```

  - Code to run a `for` loop to generate 1000 permuted versions of the test statistic using the `shuffle` function and keep track of the results in `Tstar`
- `pdata(Tstar, abs(Tobs, lower.tail=F)`
  - Finds the proportion of the permuted test statistics in `Tstar` that are less than  $-|Tobs|$  or greater than  $|Tobs|$ , useful for finding the two-sided test p-value.
- ```
Tobs <- diffmean(Y~X, data=DATASETNAME, var.equal=T)$statistic; Tobs
B <- 1000
Tstar <- matrix(NA, nrow=B)
for (b in (1:B)){
  Tstar[b] <- diffmean(Y~X, data=resample(DATASETNAME))
}
```

  - Code to run a `for` loop to generate 1000 bootstrapped versions of the data set using the `resample` function and keep track of the results of the statistic in `Tstar`.
- `qdata(Tstar, c(0. 025, 0. 975))`
  - Provides the values that delineate the middle 95% of the results in the bootstrap distribution (`Tstar`)

## 2.12 Practice problems

Load the `HELPrct` data set from the `mosaicData` package (Pruim et al., 2016a) (you need to install the `mosaicData` package once to be able to load it). The HELP study was a clinical trial for adult inpatients recruited from a detoxification unit. Patients with no primary care physician were randomly assigned to receive a multidisciplinary assessment and a brief motivational intervention or usual care and various outcomes were observed. Two of the variables in the data set are `sex`, a factor with levels male and female and `daysanysub` which is the time (in days) to first use of any substance post-detox. We are interested in the difference in mean number of days to first use of any substance post-detox between males and females. There are some missing responses and the following code will produce `favstats` with the missing values and then provide a data set that for complete observations by applying the `na.omit` function that removes any observations with missing values.

```
require(mosaicData)
data(HELPrct)
HELPrct <- HELPrct[, c("daysanysub", "sex")] #Just focus on two variables
HELPrct <- na.omit(HELPrct2) #Removes subjects with missing
favstats(daysanysub~sex, data=HELPrct2)
favstats(daysanysub~sex, data=HELPrct3)
```

2.1. Based on the results provided, how many observations were missing for males and females? Missing values here likely mean that the subjects didn't use any substances post-detox in the time of the study but might have at a later date – the study just didn't run long enough. This is called **censoring**. What is the problem with the numerical summaries here if the missing responses were all something larger than the largest observation?

2.2. Make a beanplot and a boxplot of `daysanysub ~ sex` using the `HELPrc3` data set created above. Compare the distributions, recommending parametric or nonparametric inferences.

2.3. Generate the permutation results and write out the 6+ steps of the hypothesis test, making sure to note the numerical value of observed test statistic you are using and include a discussion of the scope of inference.

2.4. Interpret the p-value for these results.

2.5. Generate the parametric `t.test` results, reporting the test-statistic, its distribution under the null hypothesis, and compare the p-value to those observed using the permutation approach.

2.6. Make and interpret a 95% bootstrap confidence interval for the difference in the means.

# Chapter 3

## One-Way ANOVA

### 3.1 Situation

In Chapter 2, tools for comparing the means of two groups were considered. More generally, these methods are used for a quantitative response and a categorical explanatory variable (group) which had two and only two levels. The full prisoner rating data set actually contained three groups (Figure 3.1 with *Beautiful*, *Average*, and *Unattractive* rated pictures randomly assigned to the subjects for sentence ratings. In a situation with more than two groups, we have two choices. First, we could rely on our two group comparisons, performing tests for every possible pair (*Beautiful* vs *Average*, *Beautiful* vs *Unattractive*, and *Average* vs *Unattractive*). We spent Chapter 2 doing inferences for differences between *Average* and *Unattractive*. The other two comparisons would lead us to initially end up with three p-values and no direct answer about our initial question of interest – is there some overall difference in the average sentences provided across the groups? In this chapter, we will learn a new method, called *Analysis of Variance*, or *One-Way ANOVA* since there is just one<sup>1</sup> grouping variable. After we perform our One-Way ANOVA test for overall evidence of a difference, we will revisit the comparisons similar to those considered in Chapter 2 to get more details on specific differences among *all* the pairs of groups – what we call *pair-wise comparisons*. An issue is created when you perform many tests simultaneously and we will augment our previous methods with an adjusted method for pairwise comparisons to make our results valid called *Tukey's Honest Significant Difference*.

To make this more concrete, we return to the original MockJury data, making side-by-side boxplots and beanplots (Figure 3.1 as well summarizing the sentences for the three groups using `favstats`.

```
require(heplots)
require(mosaic)
data(MockJury)
par(mfrow=c(1,2))
boxplot(Years~Attr,data=MockJury)
require(beanplot)
beanplot(Years~Attr,data=MockJury,log="",col="bisque",method="jitter")
favstats(Years~Attr,data=MockJury)

##          Attr min Q1 median   Q3 max     mean      sd n missing
## 1    Beautiful  1  2       3  6.5 15 4.333333 3.405362 39      0
## 2      Average  1  2       3  5.0 12 3.973684 2.823519 38      0
## 3 Unattractive  1  2       5 10.0 15 5.810811 4.364235 37      0
```

<sup>1</sup>In Chapter 4, methods are discussed for when there are two categorical explanatory variables that is called the Two-Way ANOVA and related ANOVA tests are used in Chapter 8 for working with extensions of these models.

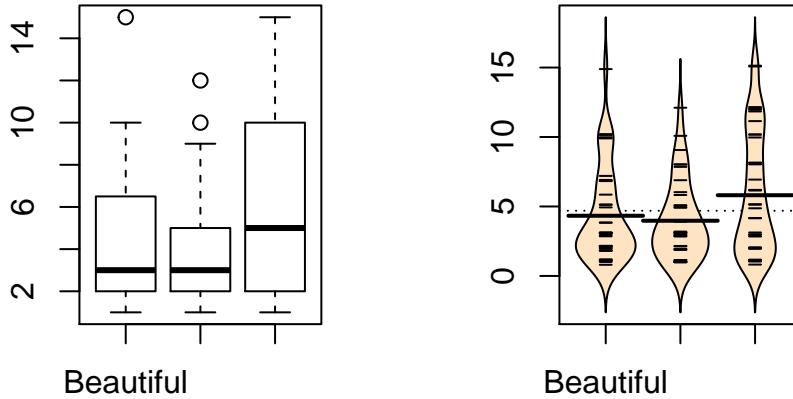


Figure 3.1: Boxplot and beanplot of the sentences (years) for the three treatment groups.

There are slight differences in the sample sizes in the three groups with 37 *Unattractive*, 38 *Average* and 39 *Beautiful* group responses, providing a data set has a total sample size of  $N = 114$ . The *Beautiful* and *Average* groups do not appear to be very different with means of 4.33 and 3.97 years. In Chapter 2, we found moderate evidence regarding the difference in *Average* and *Unattractive*. It is less clear whether we might find evidence of a difference between *Beautiful* and *Unattractive* groups since we are comparing means of 5.81 and 4.33 years. All the distributions appear to be right skewed with relatively similar shapes. The variability in *Average* and *Unattractive* groups seems like it could be slightly different leading to an overall concern of whether the variability is the same in all the groups.

## 3.2 Linear model for One-Way ANOVA (cell-means and reference-coding)

We introduced the statistical model  $y_{ij} = \mu_j + \varepsilon_j$  in Chapter 2 for the situation with  $j = 1$  or 2 to denote a situation where there were two groups and, for the model that is consistent with the alternative hypothesis, the means differed. Now we have three groups and the previous model can be extended to this new situation by allowing  $j$  to be 1, 2, or 3. Now that we have more than two groups, we need to admit that what we were doing in Chapter 2 was actually fitting what is called a *linear model*. The linear model assumes that the responses follow a normal distribution with the linear model defining the mean, all observations have the same variance, and the parameters for the mean in the model enter linearly. This last condition is hard to explain at this level of material – it is sufficient to know that there models where the parameters enter the model nonlinearly and that they are beyond the scope of this course. The result of this constraint is that we will be able to use the same general modeling framework for the methods introduced in Chapters 3, 4, 6, 7, and 8.

As in Chapter 2, we have a null hypothesis that defines a situation (and model) where all the groups have the same mean. Specifically, the *null hypothesis* in the general situation with  $J$  groups ( $J \geq 2$ ) is to have all the **true** group means equal,

$$H_0 : \mu_1 = \dots \mu_J.$$

This defines a model where all the groups have the same mean so it can be defined in terms of a single mean,  $\mu$ , for the  $i^{th}$  observation from the  $j^{th}$  group as  $y_{ij} = \mu + \varepsilon_{ij}$ . This is not the model that most researchers want to be the final description of their study as it implies no difference in the groups. There is more caution required to specify the alternative hypothesis with more than two groups. The ***alternative hypothesis*** needs to be the logical negation of this null hypothesis of all groups having equal means; to make the null hypothesis false, we only need one group to differ but more than one group could differ from the others. Essentially, there are many ways to “violate” the null hypothesis so we choose some delicate wording for the alternative hypothesis when there are more than 2 groups. Specifically, we state the alternative as

$$H_A : \text{Not all } \mu_j \text{ are equal}$$

or, in words, **at least one of the true means differs among the J groups**. You will be attracted to trying to say that all means are different in the alternative but we do not put this strict a requirement in place to reject the null hypothesis. The alternative model allows all the true group means to differ but does require that they differ with

$$\mu_j + \varepsilon_{ij}.$$

This linear model states that the response for the  $i^{th}$  observation in the  $j^{th}$  group,  $\mathbf{y}_{ij}$ , is modeled with a group  $j$  ( $j = 1, \dots, J$ ) population mean,  $\mu_j$ , and a random error for each subject in each group  $\varepsilon_{ij}$ , that we assume follows a normal distribution and that all the random errors have the same variance,  $\sigma^2$ . We can write the assumption about the random errors, often called the ***normality assumption***, as  $\varepsilon_{ij} \sim N(0, \sigma^2)$ . There is a second way to write out this model that allows extension to more complex models discussed below, so we need a name for this version of the model. The model written in terms of the  $\mu_j$ 's is called the **cell means model** and is the easier version of this model to understand.

One of the reasons we learned about beanplots is that it helps us visually consider all the aspects of this model. In the right panel of Figure 3.1, we can see the wider, bold horizontal lines that provide the estimated group means. The bigger the differences in the sample means, the more likely we are to find evidence against the null hypothesis. You can also see the null model on the plot that assumes all the groups have the same as displayed in the dashed horizontal line at 4.7 years (the R code below shows the overall mean of *Years* is 4.7). While the hypotheses focus on the means, the model also contains assumptions about the distribution of the responses – specifically that the distributions are normal and that all the groups have the same variability. As discussed previously, it appears that the distributions are right skewed and the variability might not be the same for all the groups. The boxplot provides the information about the skew and variability but since it doesn't display the means it is not directly related to the linear model and hypotheses we are considering.

`mean(MockJury$Years)`

```
## [1] 4.692982
```

There is a second way to write out the One-Way ANOVA model that provides a framework for extensions to more complex models described in Chapter 4 and beyond. The other ***parameterization*** (way of writing out or defining) of the model is called the **reference-coded model** since it writes out the model in terms of a ***baseline group*** and deviations from that baseline or reference level. The reference-coded model for the  $i^{th}$  subject in the  $j^{th}$  group is  $y_{ij} = \alpha + \tau_j + \varepsilon_{ij}$  where  $\alpha$  (alpha) is the true mean for the baseline group (first alphabetically) and the  $\tau_j$  (tau  $j$ ) are the deviations from the baseline group for group  $j$ . The deviation for the baseline group,  $\tau_1$ , is always set to 0 so there are really just deviations for groups 2 through  $J$ . The equivalence between the two models can be seen by considering the mean for the first, second, and  $J^{th}$  groups in both models:

	Cell means:	Reference-coded:
<b>Group 1 :</b>	$\mu_1$	$\alpha$
<b>Group 2 :</b>	$\mu_2$	$\alpha + \tau_2$
...	...	...
<b>Group <math>J</math> :</b>	$\mu_J$	$\alpha + \tau_J$

The hypotheses for the reference-coded model are similar to those in the cell-means coding except that they are defined in terms of the deviations,  $\tau_j$ . The null hypothesis is that there is no deviation from the baseline for any group – that all the  $\tau_j$ 's = 0,

$$H_0 : \tau_2 = \dots = \tau_J = 0.$$

The alternative hypothesis is that at least one of the deviations is not 0,

$$H_A : \text{Not all } \tau_j \text{ equal 0.}$$

In this chapter, you are welcome to use either version (unless we instruct you otherwise) but we have to use the reference-coding in subsequent chapters. The next task is to learn how to use R's linear model `lm` function to get estimates of the parameters in each model, but first a quick review of these new ideas:

### Cell Means Version

- $H_0 : \mu_1 = \dots = \mu_J$        $H_A : \text{Not all } \mu_j \text{ equal}$
- Null hypothesis in words: No difference in the true means between the groups.
- Null model  $y_{ij} = \mu_j + \varepsilon_{ij}$
- Alternative hypothesis in words: At least one of the true means differs between the groups.
- Alternative model:  $y_{ij} = \mu_j + \varepsilon_{ij}$ .

### Reference-coded Version

- $H_0 : \tau_2 \dots \tau_J = 0$        $H_A : \text{Not all } \tau_j \text{ equal}$
- Null hypothesis in words: No deviation of the true mean for any groups from the baseline group.
- Null model:  $y_{ij} = \alpha + \tau_j + \varepsilon_{ij}$
- Alternative hypothesis in words: At least one of the true deviations is different from 0 or that at least one group has a different true mean than the baseline group.
- Alternative model:  $y_{ij} = \alpha + \tau_j + \varepsilon_{ij}$

In order to estimate the models discussed above, the `lm` function is used. If you look closely in the code for the rest of the book, any model for a quantitative response will use this function, suggesting a common thread in the most commonly used statistical models. The `lm` function continues to use the same format as previous functions, `lm(Y~X, data=datasetname)`. It ends up that this code will give you the reference-coded version of the model by default (R thinks it is that important!). We want to start with the cell-means version of the model, so we have to override the standard technique and add a “-1” to the formula interface to tell R that we want to the cell-means coding. Generally, this looks like `lm(Y~X-1, data=datasetname)`. Once we fit a model in R, the `summary` function run on the model provides a useful “summary” of the model coefficients and a suite of other potentially interesting information. When fitting this version of the One-Way ANOVA model, you will find a row of output for each group relating the  $\mu_j$ 's. The output contains columns for an estimate (`Estimate`), standard error (`Std.Error`),  $t$ -value (`t value`), and p-value (`Pr(>|t|)`). We'll learn to use all the output in the following material, but for now just focus on the estimates of the parameters that the function provides that we put in bold.

```
lm1 <- lm(Years ~ Attr-1, data=MockJury)
summary(lm1)

##
## Call:
## lm(formula = Years ~ Attr - 1, data = MockJury)
##
## Residuals:
##    Min     1Q Median     3Q    Max
## -4.8108 -2.8108 -0.9737  2.1892 10.6667
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## AttrBeautiful 4.3333   0.5730   7.563 1.23e-11
## AttrAverage   3.9737   0.5805   6.845 4.41e-10
## AttrUnattractive 5.8108   0.5883   9.878 < 2e-16
##
## Residual standard error: 3.578 on 111 degrees of freedom
## Multiple R-squared:  0.6449, Adjusted R-squared:  0.6353
## F-statistic: 67.21 on 3 and 111 DF, p-value: < 2.2e-16
```

In general, we denote estimated parameters with a hat over the parameter of interest to show that it is an estimate. For the true mean of group  $j$ ,  $\mu_j$ , we estimate it with  $\hat{\mu}_j$ , which is just the sample mean for group  $j$ ,  $\bar{x}_j$ . The model suggests an estimate for each observation that we denote as  $\hat{y}_{ij}$  that we will also call a **fitted value** based on the model being considered. The three estimates are bolded in the previous output, with the same estimate used for all observations in the same group. R tries to help you to sort out which row of output corresponds to which group by appending the group name 1 with the variable name. Here, the variable name was `Attr` and the first group alphabetically was *Beautiful*, so R provides a row labeled `AttrBeautiful` with an estimate of 4.3333. The sample means from the three groups can be seen to directly match that and the other two results.

```
mean(Years ~ Attr, data=MockJury)
```

```
##   Beautiful     Average Unattractive
##   4.333333 3.973684 5.810811
```

The reference-coded version of the same model is more complicated but ends up giving the same results once we understand what it is doing. It uses a different parameterization to accomplish this so has different model output. Here is the model summary:

```
lm2 <- lm(Years ~ Attr, data=MockJury)
summary(lm2)

##
## Call:
## lm(formula = Years ~ Attr, data = MockJury)
##
## Residuals:
##    Min     1Q Median     3Q    Max
## -4.8108 -2.8108 -0.9737  2.1892 10.6667
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) 4.3333   0.5730   7.563 1.23e-11
## AttrAverage -0.3596   0.8157  -0.441  0.6601
## AttrUnattractive 1.4775   0.8212   1.799  0.0747
##
## Residual standard error: 3.578 on 111 degrees of freedom
## Multiple R-squared:  0.04754, Adjusted R-squared:  0.03038
## F-statistic: 2.77 on 2 and 111 DF, p-value: 0.067
```

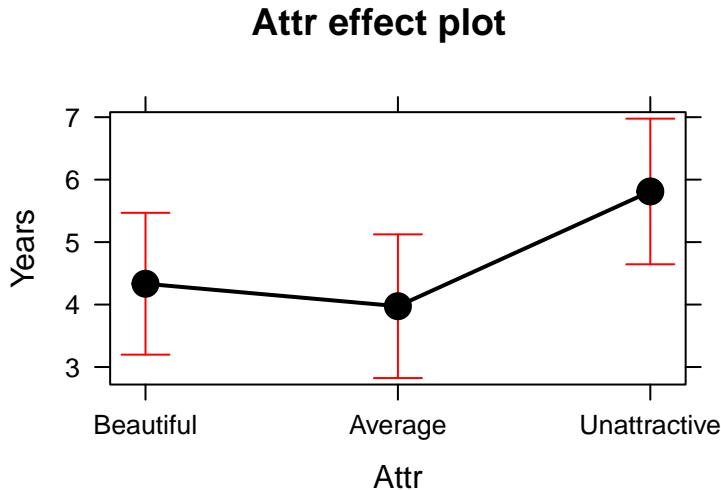


Figure 3.2: Plot of the estimated group mean sentences from the reference-coded model for the MockJury data.

The estimated model coefficients are  $\hat{\alpha} = 4.333$  years,  $\hat{\tau}_2 = -0.3596$  years,  $\hat{\tau}_3 = 1.4775$  years where R selected group 1 for *Beautiful*, 2 for *Average*, and 3 for *Unattractive*. The way you can figure out the baseline group (group 1 is *Beautiful* here) is to see which category label is *not present* in the output. **The baseline level is typically the first group label alphabetically**, but you should always check this. Based on these definitions, there are interpretations available for each coefficient. For  $\hat{\alpha} = 4.333$  years, this is an estimate of the mean sentencing time for the *Beautiful* group.  $\hat{\tau}_2 = -0.3596$  years is the deviation of the *Average* group's mean from the *Beautiful* group's mean (specifically, it is 0.36 years lower). Finally,  $\hat{\tau}_3 = 1.4775$  years tells us that the *Unattractive* group mean sentencing time is 1.48 years higher than time. These interpretations lead directly to reconstructing the estimated means for each group by combining the baseline and pertinent deviations as shown in Table 3.1.

Table 3.1: Constructing group mean estimates from the reference-coded linear model estimates.

Group	Formula	Estimates
Beautiful	$\hat{\alpha}$	<b>4.3333</b> years
Average	$\hat{\alpha} + \hat{\tau}_2$	$4.3333 - 0.3596 = \mathbf{3.974}$ years
Unattractive	$\hat{\alpha} + \hat{\tau}_3$	$4.3333 + 1.4775 = \mathbf{5.811}$ years

We can also visualize the results of our linear models using what are called ***term-plots*** or ***effect-plots*** (from the **effects** package; Fox (2003), (Fox et al., 2016)) as displayed in Figure 3.2. We don't want to use the word "effect" for these model components unless we have random assignment in the study design so we generically call these ***term-plots*** as they display terms or components from the model in hopefully useful ways to aid in model interpretation even in the presence of complicated model parameterizations. Specifically, these plots take an estimated model and show you its estimates along with 95% confidence intervals generated by the linear model. To make this plot, you need to install and load the **effects** package and then use `plot(allEffects(...))` functions together on the `lm` object called `lm2` that was estimated above. You can find the correspondence between the displayed means and the estimates that were constructed in Table 3.1.

```
require(effects)
```

```
plot(allEffects(lm2))
```

In order to assess evidence for having different means for the groups, we will compare either of the previous models (cell-means or reference-coded) to a null model based on the null hypothesis ( $H_0 : \mu_1 = \dots = \mu_J$ ) which

implies a model of  $y_{ij} = \mu_j + \varepsilon_{ij}$  in the cell-means version where  $\mu$  is a common mean for all the observations. We will call this the **mean-only** model since it only has a single mean in it. In the reference-coding version of the model, we have a null hypothesis that  $H_0 : \tau_2 = \dots = \tau_J = 0$ , so the “mean-only” model is  $y_{ij} = \alpha + \varepsilon_{ij}$  with  $\alpha$  having the same definition as  $\mu$  for the cell means model – it forces a common value for the mean for all the groups. Moving from the *reference-coded* model to the *mean-only* model is also an example of a situation where we move from a “full” model to a “reduced” model by setting some coefficients in the “full” model to 0 and, by doing this, get a simpler or “reduced” model. Simple models can be good as they are easier to groups that suggests no difference in the groups is not a very exciting result in most, but not all, situations<sup>2</sup>. In order for R to provide results for the mean-only model, we remove the grouping variable, `Attr`, from the model formula and just include a “1”. The `(Intercept)` row of the output provides the estimate for the mean-only model as a reduced model from either the cell-means or reference-coded models when we assume that the mean is the same for all groups:

```
lm3 <- lm(Years ~ 1, data=MockJury)
```

```
summary(lm3)
```

```
## $coefficients
##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) 4.692982  0.3403532 13.78857 5.765681e-26
```

This model provides an estimate of the common mean for all observations of  $4.693 = \hat{\mu} = \hat{\alpha}$  years. This value also is the dashed, horizontal line in the beanplot in Figure 3.1. Some people call this mean-only estimate the grand or overall mean.

### 3.3 One-Way ANOVA Sums of Squares, Mean Squares, and F-test

The previous discussion showed two ways of parameterizing models for the One-Way ANOVA model and getting estimates from output but still hasn’t addressed how to assess evidence related to whether the observed differences in the means among the groups is “real”. In this section, we develop what is called the **ANOVA F-test** that provides a method of aggregating the differences among the means of 2 or more groups and testing our null hypothesis of no difference in the means vs the alternative. In order to develop the test, some additional notation is needed. The sample size in each group is denoted  $n_j$  and the total sample size is  $N = \sum n_j = n_1 + n_2 + \dots + n_J$  where  $\Sigma$  (capital sigma) means “add up over whatever follows”. An estimated **residual** ( $e_{ij}$ ) is the difference between an observation,  $y_{ij}$ , and the model estimate,  $\hat{y}_{ij} = \hat{\mu}_j$ , for that observation,  $y_{ij} - \hat{y}_{ij} = e_{ij}$ . It is basically what is left over that the mean part of the model ( $\hat{\mu}_j$ ) does not explain. It is also a window into how “good” the model might be.

Consider the four different fake results for a situation with four groups ( $J = 4$ ) displayed in Figure 3.3. Which of the different results shows the most and least evidence of differences in the means? In trying to answer this, think about both how different the means are (obviously important) and how variable the results are around the mean. These situations were created to have the same means in Scenarios 1 and 2 as well as matching means in Scenarios 3 and 4. The variability around the means matches by shading (lighter or darker). In Scenarios 1 and 2, the differences in the means is smaller than in the other two results. But Scenario 2 should provide more evidence of what little difference is present than Scenario 1 because it has less variability around the means. The best situation for finding group differences here is Scenario 4 since it has the largest difference in the means and the least variability around those means. Our test statistic somehow needs to allow a comparison of the variability in the means to the overall variability to help us get results that reflect that Scenario 4 has the strongest evidence of a difference and Scenario 1 would have the least.

The statistic that allows the comparison of relative amounts of variation is called the **ANOVA F-statistic**. It is developed using **sums of squares** which are measures of total variation like are used in the numerator of the standard deviation ( $(\sum_1^N (y_i - \bar{y})^2)$ ) that took all the observations, subtracted the mean, squared the

---

<sup>2</sup>Suppose we were doing environmental monitoring and were studying asbestos levels in soils. We might be hoping that the mean-only model were reasonable to use if the groups being compared were in remediated areas and in areas known to have never been contaminated.

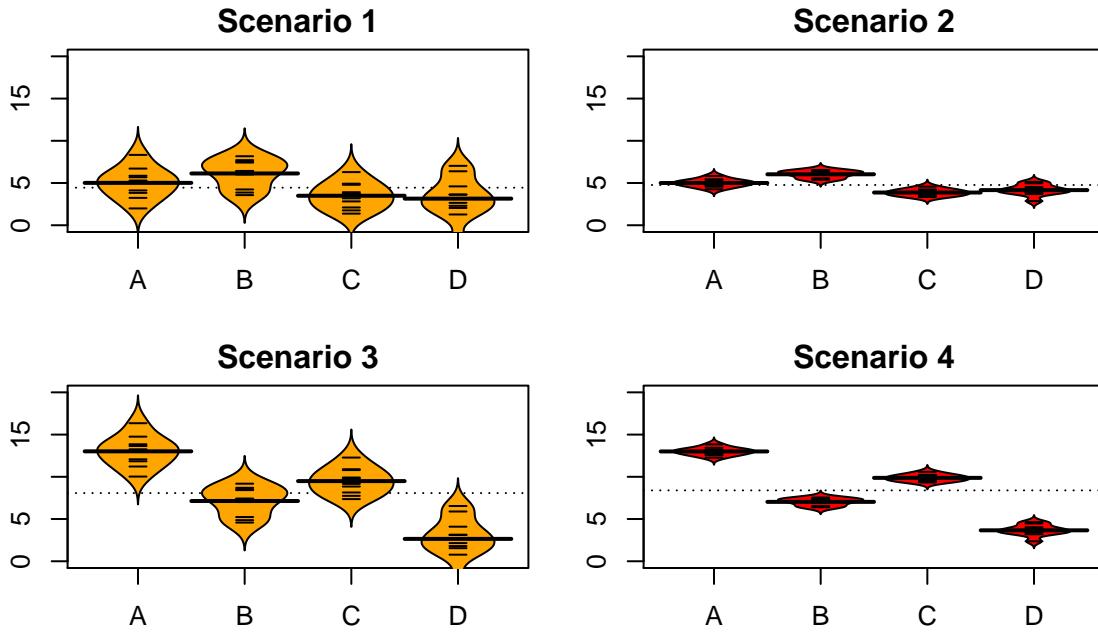


Figure 3.3: Demonstration of different amounts of difference in means relative to variability. Scenarios have same means in rows and same variance around means in columns of plot.

differences, and then added up the results over all the observations to generate a measure of total variability. With multiple groups, we will focus on decomposing that total variability (**Total Sums of Squares**) into variability among the means (we'll call this **Explanatory Variable A's Sums of Squares**) and variability in the residuals or errors (**Error Sums of Squares**). We define each of these quantities in the One-Way ANOVA situation as follows:

- $\text{SS}_{\text{Total}} = \text{Total Sums of Squares} = \sum_{j=1}^J \sum_{i=1}^{n_j} (y_{ij} - \bar{y})^2$ 
  - This is the total variation in the responses around the overall or **grand mean** ( $\bar{y}$ , the estimated mean for all the observations and available from the mean-only model).
  - By summing over all  $n_j$  observations in each group,  $\sum_{i=1}^{n_j} (\ )$ , and then adding those results up across the groups,  $\sum_{j=1}^J (\ )$ , we accumulate the variation across all  $N$  observations.
  - Note: this is the residual variation if the null model is used, so there is no further decomposition possible for that model.
  - This is also equivalent to the numerator of the sample variance,  $\sum_1^N (y_i - \bar{y})^2$  which is what you get when you ignore the information on the potential differences in the groups.
- $\text{SS}_A = \text{Explanatory Variable } A\text{'s Sums of Squares} = \sum_{j=1}^J \sum_{i=1}^{n_j} (\bar{y}_i - \bar{y})^2 = \sum_{j=1}^J n_j (\bar{y}_i - \bar{y})^2$ 
  - This is the variation in the group means around the grand mean based on the explanatory variable  $A$ .
  - Also called sums of squares for the treatment, regression, or model.
- $\text{SS}_E = \text{Error (Residual) Sums of Squares} = \sum_{j=1}^J \sum_{i=1}^{n_j} (y_{ij} - \bar{y}_i)^2 = \sum_{j=1}^J \sum_{i=1}^{n_j} (e_{ij})^2$ 
  - This is the variation in the responses around the group means.
  - Also called the sums of squares for the residuals, with the second version of the formula showing that it is just the squared residuals added up across all the observations.

The possibly surprising result given the mass of notation just presented is that the total sums of squares is **ALWAYS** equal to the sum of explanatory variable  $A$ 's sum of squares and the error sums of squares,

$$SS_{\text{Total}} = SS_A + SS_E.$$

This equality means that if the  $SS_A$  goes up, then the  $SS_E$  must go down if  $SS_{\text{Total}}$  remains the same. This result is called the *sums of squares decomposition formula*. We use these results to build our test statistic and organize this information in what is called an *ANOVA table*. The ANOVA table is generated using the `anova` function applied to the reference-coded model, `lm2`:

```
lm2<-lm(Years ~ Attr, data=MockJury)
```

```
anova(lm2)
```

```
## Analysis of Variance Table
##
## Response: Years
##             Df  Sum Sq Mean Sq F value Pr(>F)
## Attr          2   70.94  35.469   2.77  0.067
## Residuals 111 1421.32  12.805
```

Note that the ANOVA table has a row labelled `Attr`, which contains information for the grouping variable (we'll generally refer to this as explanatory variable  $A$  but here it is the picture group that was randomly assigned), and a row labelled `Residuals`, which is synonymous with "Error". The Sums of Squares (SS) are available in the `Sum Sq` column. It doesn't show a row for "Total" but the  $SS_{\text{Total}} = SS_A + SS_E = 1492.26$ .

```
70.94 + 1421.32
```

```
## [1] 1492.26
```

It may be easiest to understand the *sums of squares decomposition* by connecting it to our permutation ideas. In a permutation situation, the total variation ( $SS_{\text{Total}}$ ) cannot change – it is the same responses varying around the grand mean. However, the amount of variation attributed to variation among the means and in the residuals can change if we change which observations go with which group. In Figure 3.4 (panel a), the means, sums of squares, and 95% confidence intervals for each mean are displayed for the three treatment levels from the original prisoner rating data. Three permuted versions of the data set are summarized in panels (b), (c), and (d). The  $SS_A$  is 70.9 in the real data set and between 6.6 and 11 in the permuted data sets. If you had to pick among the plots for the one with the most evidence of a difference in the means, you hopefully would pick panel (a). This visual "unusualness" suggests that this observed result is unusual relative to the possibilities under permutations, which are, again, the possibilities tied to having the null hypothesis being true. But note that the differences here are not that great between these three permuted data sets and the real one. It is likely that at least some of you might have selected panel (d) as also looking like it shows some evidence of differences (maybe not the most?) as it also looks like it shows some evidence differences.

One way to think about  $SS_A$  is that it is a function that converts the variation in the group means into a single value. This makes it a reasonable test statistic in a permutation testing context. By comparing the observed  $SS_A = 70.9$  to the permutation results of 6.5, 9.7, and 40.5 we see that the observed result is much more extreme than the three alternate versions. In contrast to our previous test statistics where positive and negative differences were possible,  $SS_A$  is always positive with a value of 0 corresponding to no variation in the means. The larger the  $SS_A$ , the more variation there is in the means. The permutation p-value for the alternative hypothesis of **some** (not of greater or less than!) difference in the true means of the groups will involve counting the number of permuted  $SS_A^*$  results that are larger than what we observed.

```
## [1] 70.93836
```

To do a permutation test, we need to be able to calculate and extract the  $SS_A$  value. In the ANOVA table, it is in the first row and is the second number and we can use the bracket, `[, ]`, referencing to extract that number from the ANOVA table that `anova` produces with `anova(lm(Years~Attr, data=MockJury))` [1, 2]. We'll store the observed value of  $SS_A$  in `Tobs`, reusing some ideas from Chapter 2.

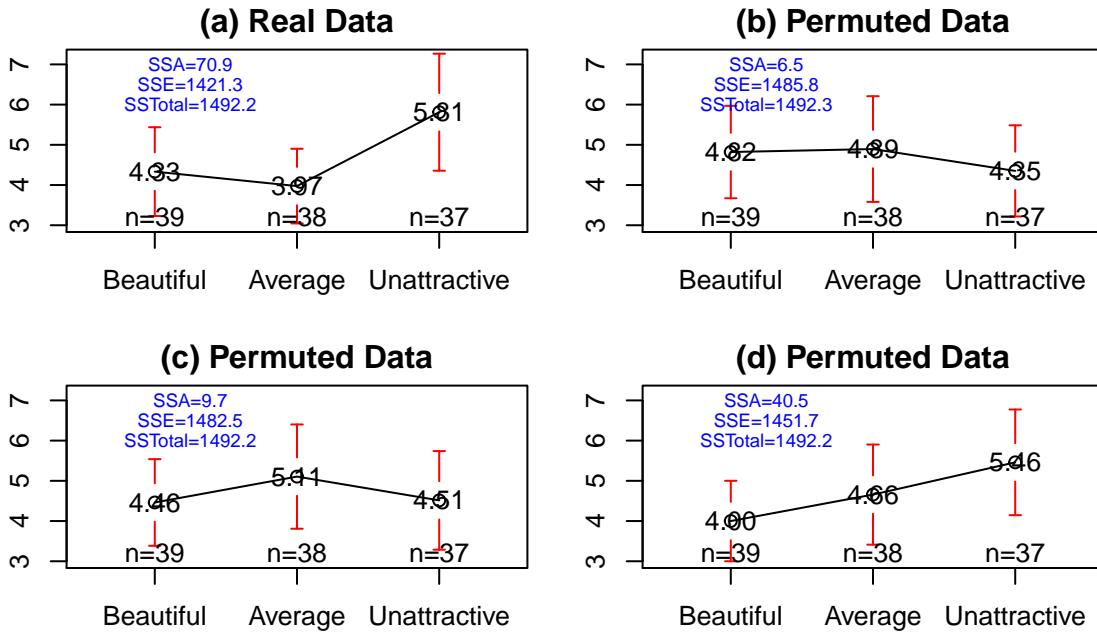


Figure 3.4: Plot of means and 95% confidence intervals for the three groups for the real data (a) and three different permutations of the treatment labels to the same responses in (b), (c), and (d). Note that SSTotal is always the same but the different amounts of variation associated with the means (SSA) or the errors (SSE) changes in permutation.

```
Tobs <- anova(lm(Years~Attr,data=MockJury))[1,2]; Tobs
## [1] 70.93836
```

The following code performs the permutations  $B=1,000$  times using the `shuffle` function, builds up a vector of results in `Tobs`, and then makes a plot of the resulting permutation distribution:

```
par(mfrow=c(1,2))
B<- 1000
Tstar<-matrix(NA,nrow=B)
for (b in (1:B)){
  Tstar[b]<-anova(lm(Years~shuffle(Attr),data=MockJury))[1,2]
}
hist(Tstar,labels=T,ylim=c(0,550))
abline(v=Tobs,col="red",lwd=3)
plot(density(Tstar),main="Density curve of Tstar")
abline(v=Tobs,col="red",lwd=3)
```

The right-skewed distribution (Figure 3.5) contains the distribution of  $SS_A$ 's under permutations (where all the groups are assumed to be equivalent under the null hypothesis). While the observed result is larger than many of the  $SS_A$ 's, there are also many permuted results that are much larger than observed. The proportion of permuted results that exceed the observed value is found using `pdata` as before, except only for the area to the right of the observed result. We know that `Tobs` will always be positive so no absolute values are required here.

```
pdata(Tstar,Tobs,lower.tail=F)
## [1] 0.072
```

This provides a permutation-based p-value of 0.072 and suggests marginal evidence against the null hypothesis

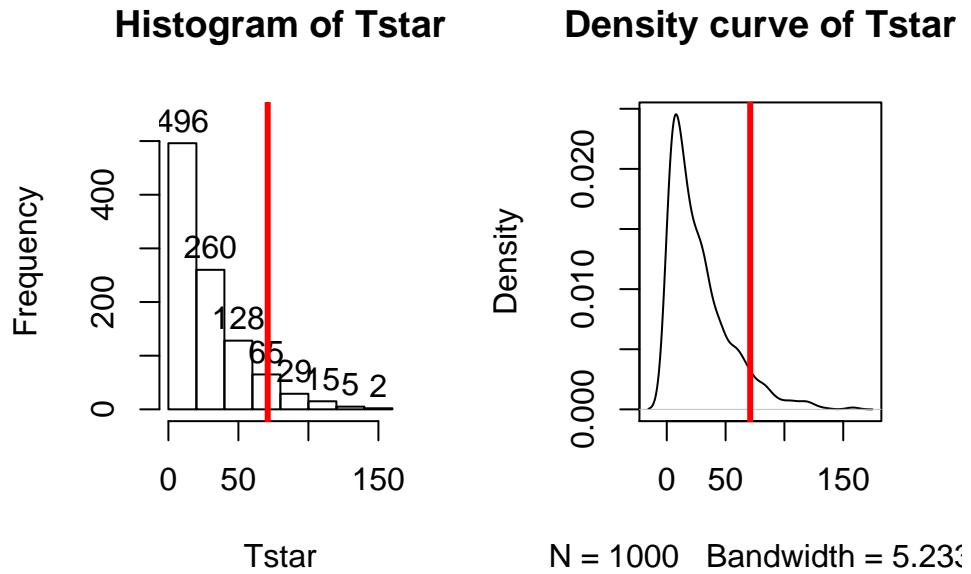


Figure 3.5: Histogram and density curve of permutation distribution of  $SS_A$  with the observed value of  $SS_A$  displayed as a bold, vertical line. The proportion of results that are larger than the observed value of  $SS_A$  provides an estimate of the p-value.

of no difference in the true means. We would interpret this p-value as saying that there is a 7.2% chance of getting a  $SS_A$  as large or larger than we observed, given that the null hypothesis is true.

It turns out that some nice parametric statistical results are available (if our assumptions are met) for the ratio of estimated variances, which are called **Mean Squares**. To turn sums of squares into mean square (variance) estimates, we divide the sums of squares by the amount of free information available. For example, remember the typical variance estimator introductory statistics,  $\sum_1^N (y_i - \bar{y})^2 / (N - 1)$ ? Your instructor spent some time trying various approaches to explaining why we have a denominator of  $N - 1$ . The most useful for our purposes moving forward is that we “lose” one piece of information to estimate the mean and there are  $N$  deviations around the single mean so we divide by  $N - 1$ . The main point is that the sums of squares were divided by something and we got an estimator for the variance, here of the observations.

Now consider  $SS_E = \sum_{j=1}^J \sum_{i=1}^{n_j} (y_i - \bar{y}_j)^2$  which still has  $N$  deviations but it varies around the  $J$  means, so the

$$\text{Mean Square Error} = MS_E = SS_E / (N - J).$$

Basically, we lose  $J$  pieces of information in this calculation because we have to estimate  $J$  means.

The similar calculation of the **Mean Square for variable A** ( $MS_A$ ) is harder to see in the formula ( $SS_A = \sum_{j=1}^J n_j (\bar{y}_j - \bar{y})^2$ ), but the same reasoning can be used to understand the denominator for forming  $MS_A$ : there are  $J$  means that vary around the grand mean so

$$MS_A = SS_A / (J - 1).$$

In summary, the two mean squares are simply:

- $MS_A = SS_A / (J - 1)$ , which estimates the variance of the group means around the grand mean.
- $MS_{\text{Error}} = SS_{\text{Error}} / (N - J)$ , which estimates the variation of the errors around the group means.

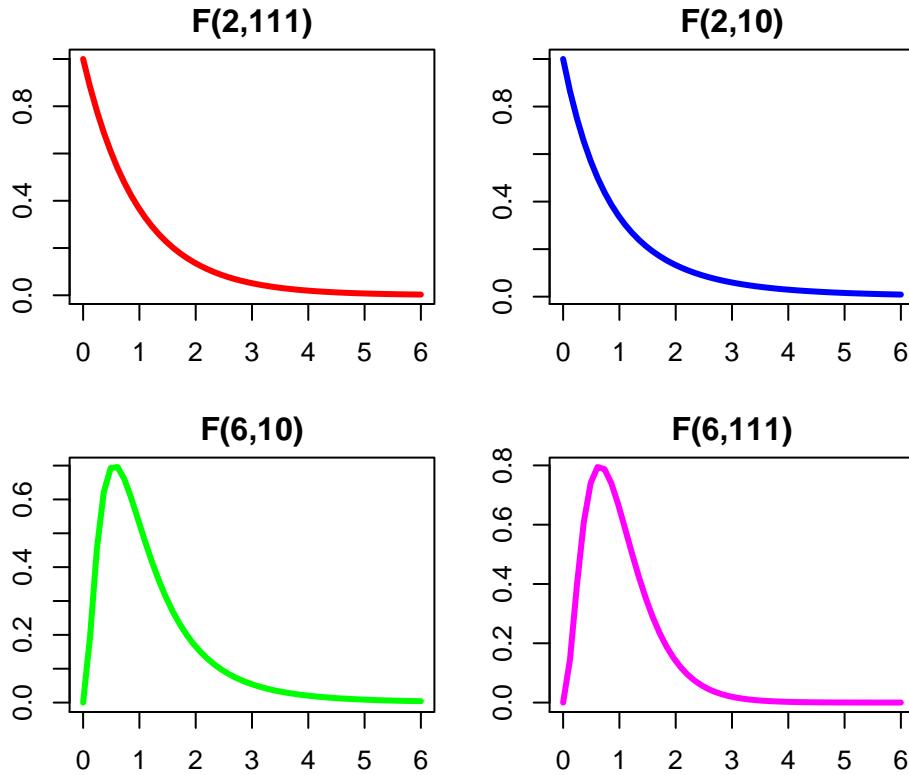


Figure 3.6: Density curves of four different  $F$ -distributions. Upper left is an  $F(2, 111)$ , upper right is  $F(2, 10)$ , lower left is  $F(6, 10)$ , and lower right is  $F(6, 111)$ . P-values are found using the areas to the right of the observed  $F$ -statistic value.

These results are put together using a ratio to define the ***ANOVA F-statistic*** (also called the ***F-ratio***) as

$$F = \text{MS}_A / \text{MS}_{\text{Error}}.$$

If the variability in the means is “similar” to the variability in the residuals, the statistic would have a value around 1. If that variability is similar then there be no evidence of a difference in the means. If the  $\text{MS}_A$  is much larger than the  $\text{MS}_E$ , the  $F$ -statistic will provide evidence against the null hypothesis. The “size” of the  $F$ -statistic is formalized by finding the p-value. The  $F$ -statistic, if assumptions discussed below are met and we assume the null hypothesis is true, follows what is called an  $F$ -distribution. The ***F-distribution*** is a right-skewed distribution whose shape is defined by what are called the ***numerator degrees of freedom*** ( $J - 1$ ) and the ***denominator degrees of freedom*** ( $N - J$ ). These names correspond to the values that we used to calculate the mean squares and where in the  $F$ -ratio each mean square was used;  $F$ -distributions are denoted by their degrees of freedom using the convention of  $F$  (*numerator df, denominator df*). Some examples of different  $F$ -distributions are displayed for you in Figure 3.6.

The characteristics of the F-distribution can be summarized as:

- Right skewed,
- Nonzero probabilities for values greater than 0,
- Its shape changes depending on the **numerator** and **denominator DF**, and
- **Always use the right-tailed area for p-values.**

Now we are ready to discuss an ANOVA table since we know about each of its components. Note the general format of the ANOVA table is<sup>3</sup>:

Table 3.2: General One-Way ANOVA table.

Source	DF	Sums of Squares	Mean Squares	F-ratio	P-value
Variable A	$J - 1$	$SS_A$	$MS_A = SS_A/(J - 1)$	$F = MS_A/MS_E$	Right tail of $F(J - 1, N - J)$
Residuals	$N - J$	$SS_E$	$MS_E = SS_E/(N - J)$		
Total	$N - 1$	$SS_{\text{Total}}$			

The table is oriented to help you reconstruct the  $F$ -ratio from each of its components. The output from R is similar although it does not provide the last row and sometimes switches the order of columns. The R version of the table for the type of picture effect (`Attr`) with  $J = 3$  levels and  $N = 114$  observations, repeated from above, is:

```
anova(lm2)
```

```
## Analysis of Variance Table
##
## Response: Years
##           Df  Sum Sq Mean Sq F value Pr(>F)
## Attr         2   70.94  35.469   2.77  0.067
## Residuals 111 1421.32 12.805
```

The p-value from the  $F$ -distribution is 0.067. We can verify this result using the observed  $F$ -statistic of 2.77 (which came from taking the ratio of the two mean squares,  $F=35.47/12.8$ ) which follows an  $F(2, 111)$  distribution if the null hypothesis is true and some other assumptions are met.

Using the `pf` function provides us with areas in the specified  $F$ -distribution with the `df1` provided to the function as the numerator `df` and `df2` as the denominator `df` and `lower.tail=F` reflecting our desire for a right tailed area.

```
pf(2.77,df1=2,df2=111,lower.tail=F)
```

```
## [1] 0.06699803
```

The result from the  $F$ -distribution using this parametric procedure is similar to the p-value obtained using permutations with the test statistic of the  $SS_A$ , which was 70.9. The  $F$ -statistic obviously is another potential test statistic to use as a test statistic in a permutation approach, now that we know about it. We should check that we get similar results from it with permutations as we did from using  $SS_A$  as a permutation test statistic. The following code generates the permutation distribution for the  $F$ -statistic (Figure 3.7) and assesses how unusual the observed  $F$ -statistic of 2.77 was in this permutation distribution. The only change in the code involves moving from extracting  $SS_A$  to extracting the  $F$ -ratio which is in the 4th column of the `anova` output:

```
Tobs <- anova(lm(Years ~ Attr, data=MockJury))[1,4]; Tobs

## [1] 2.770024
par(mfrow=c(1,2))
B<- 1000
Tstar<-matrix(NA,nrow=B)
for (b in (1:B)){
  Tstar[b]<-anova(lm(Years~shuffle(Attr), data=MockJury))[1,4]
}
```

<sup>3</sup>Make sure you can work from left to right and up and down to fill in the ANOVA table given just the necessary information to determine the other components – there is always a question like this on the exam...

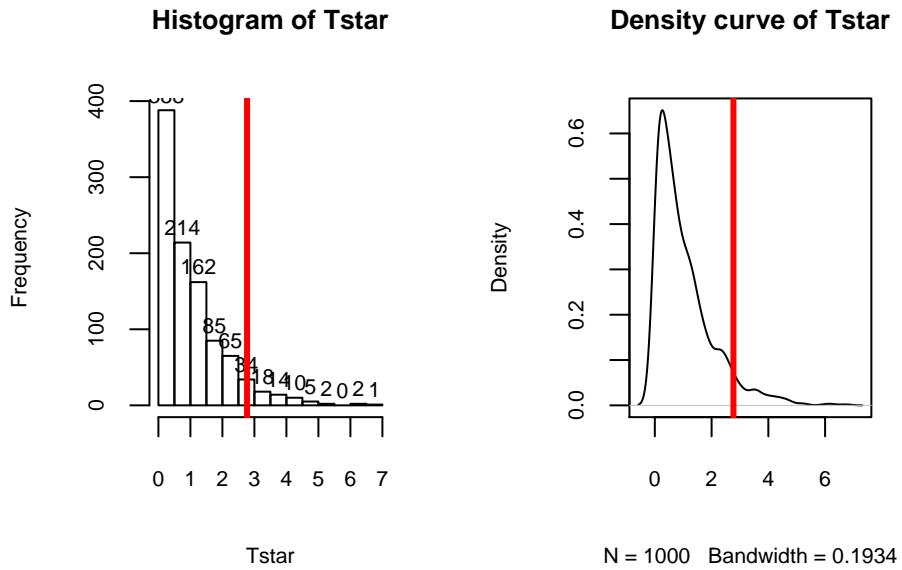


Figure 3.7: Histogram and density curve of the permutation distribution of the F-statistic with bold, vertical line for observed value of the test statistic of 2.77.

```
pdata(Tstar, Tobs, lower.tail=F)

## [1] 0.064
hist(Tstar, labels=T)
abline(v=Tobs, col="red", lwd=3)
plot(density(Tstar), main="Density curve of Tstar")
abline(v=Tobs, col="red", lwd=3)
```

The permutation-based p-value is 0.064 which, again, matches the other results closely. The first conclusion is that using a test statistic of either the  $F$ -statistic or the  $SS_A$  provide similar permutation results. However, we tend to favor using the  $F$ -statistic because it is more commonly used in reporting ANOVA results, not because it is any better in a permutation context.

It is also interesting to compare the permutation distribution for the  $F$ -statistic and the parametric  $F(2, 111)$  distribution (Figure 3.8). They do not match perfectly but are quite similar. Some the differences around 0 are due to the behavior of the method used to create the density curve and are not really a problem for the methods. The similarity in the two curves explains why both methods give similar results. In some situations, the correspondence will not be quite so close.

So how can we rectify this result ( $p\text{-value} \approx 0.06$ ) and the Chapter 2 result that detected a difference between *Average* and *Unattractive* with a  $p\text{-value} \approx 0.03$ ? I selected the two groups to compare in Chapter 2 because they were furthest apart. “Cherry-picking” the comparison that is likely to be most different creates a false sense of the real situation and inflates the Type I error rate because of the selection. If the entire suite of pairwise comparisons are considered, this result may lose some of its luster. In other words, if we consider the suite of three pair-wise differences (and the tests) implicit in comparing all of them, we may need stronger evidence in the most different pair than a  $p\text{-value}$  of 0.033 to suggest overall differences. In this situation, the *Beautiful* and *Average* groups are not that different from each other so their difference does not contribute much to the will revisit this topic and consider a method that is statistically valid for performing all possible pair-wise comparisons that is also consistent with our overall test results.

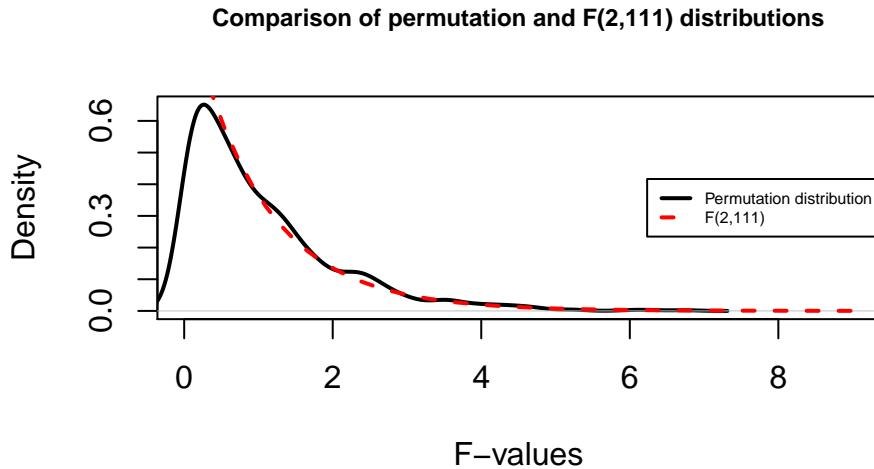


Figure 3.8: Comparison of  $F(2,111)$  (dashed line) and permutation distribution (solid line).

### 3.4 ANOVA model diagnostics including QQ-plots

The requirements for a One-Way ANOVA  $F$ -test are similar to those discussed in Chapter 2, except that there are now  $J$  groups instead of only 2. Specifically, the linear model assumes:

1. **Independent observations,**
2. **Equal variances,** and
3. **Normal distributions.**

For assessing equal variances across the groups, it is best to use plots to assess this. We can use boxplots and beanplots to compare the spreads of the groups, which were provided in Figure 3.1. The range and IQRs should be relatively similar across the groups if you do not find evidence of a problem with this assumption. You should start with noting how clear or big the violation of the assumption might be but remember that there will always be some differences in the variation among groups even if the true variability is exactly equal in the populations. In addition to our direct plotting, there are some diagnostic plots available from the `lm` function that can help us more clearly assess potential violations of the previous assumptions.

We can obtain a suite of four diagnostic plots by using the `plot` function on any linear model object that we have fit. To get all the plots together in four panels we need to add the `par(mfrow=c(2, 2))` command to tell R to make a graph with 4 panels<sup>4</sup>.

```
par(mfrow=c(2,2))
plot(lm2,pch=16)
```

There are two plots in Figure 3.9 with useful information for the equal variance assumption. The “Residuals vs Fitted” panel in the top left displays the residuals ( $e_{ij} = y_{ij} - \hat{y}_{ij}$ ) on the y-axis and the fitted values ( $\hat{y}_{ij}$ ) on the x-axis. This allows you to see if the variability of the observations differs across the groups as a function of the mean of the groups because all the observations in the same group get the same fitted value, the mean of the group. In this plot, the points seem to have fairly similar spreads at the fitted values for the three groups with fitted values of 4, 4.3, and 6. The “Scale-Location” plot in the lower left panel has the same x-axis but the y-axis contains the square-root of the absolute value of the standardized residuals. The

<sup>4</sup>We have been using this function quite a bit to make multi-panel graphs but did not show you that line of code. But you need to use this command for linear model diagnostics or you won’t get the plots we want from the model. And you really just need `plot(lm2)` but the `pch=16` option makes it easier to see some of the points in the plots.

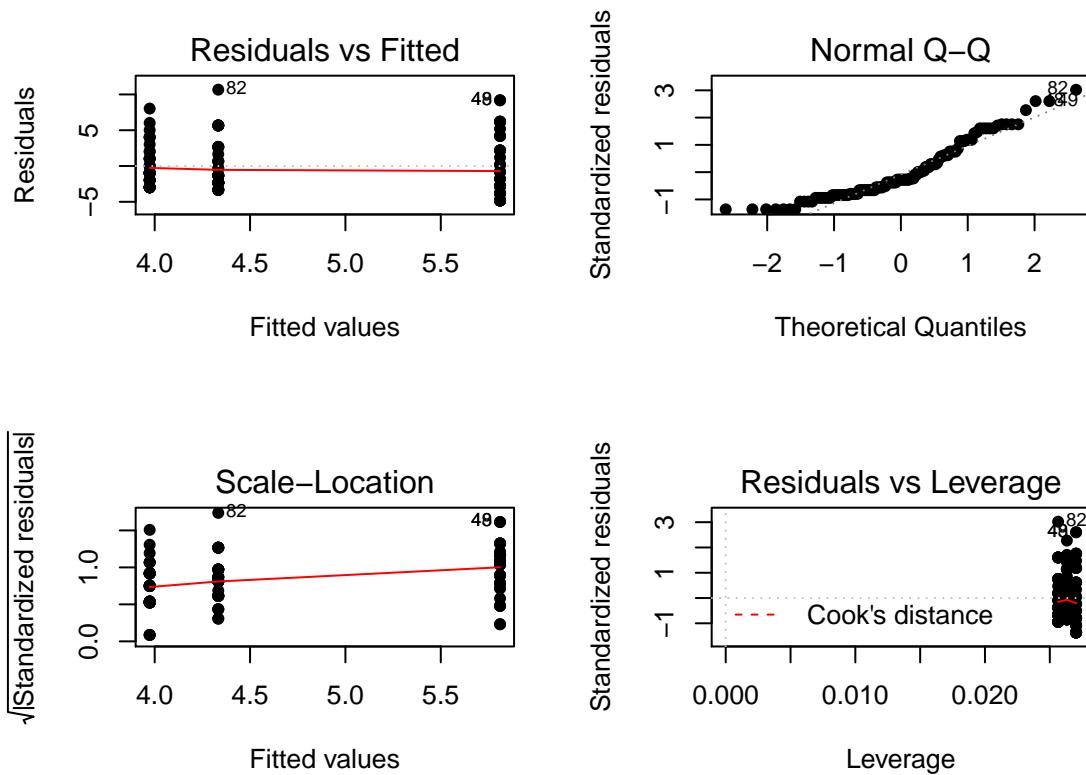


Figure 3.9: Default diagnostic plots for the linear model.

absolute value transforms all the residuals into a magnitude scale (removing direction) and the square-root helps you see differences in variability more accurately. The standardization scales them to have a variance of 1 so help you in other displays to get a sense of how many standard deviations you are away from the mean in the residual distribution. The visual assessment is similar in the two plots – you want to consider whether it appears that the groups have somewhat similar or noticeably different amounts of variability. If you see a clear funnel shape in the Residuals vs Fitted or an increase or decrease in the upper edge of points in the Scale-Location plot that may indicate a violation of the constant variance assumption. Remember that some variation across the groups is expected and is OK, but large differences in spreads are problematic for all the procedures that involve linear models. When discussing these results, you want to discuss how clearly the differences in variation are and whether that *shows a clear violation of the assumption* of equal variance for all observations. Like in hypothesis testing, you can't prove that you've met assumptions based on a plot "looking OK", but you can say that there is no clear evidence that the assumption is violated!

The linear model assumes that all the random errors ( $\varepsilon_{ij}$ ) follow a normal distribution. To gain insight into the validity of this assumption, we can explore the original observations as displayed in the beanplots, mentally subtracting off the differences in the means and focusing on the shapes of the distributions of observations in each group. These plots are especially good for assessing whether there is there a skew or outliers present in each group. If so, by definition, the normality assumption is violated. But our assumption is about the distribution of all the errors after the remove the differences in the means and so we want an overall assessment technique to understand how reasonable our assumption is overall for our model. The residuals from the entire model provide us with estimates of the random errors and if the normality assumption is met, then the residuals all-together should approximately follow a normal distribution. The **Normal Q-Q Plot** in upper right panel of Figure 3.9 is a direct visual assessment of how well our residuals match what we would expect from a normal distribution. Outliers, skew, heavy and light-tailed aspects of distributions (all

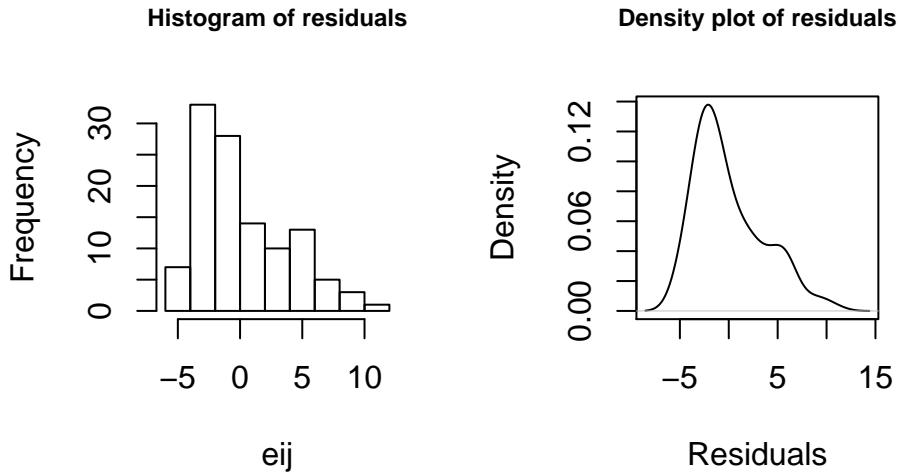


Figure 3.10: Histogram and density curve of the linear model raw residuals.

violations of normality) show up in this plot once you learn to read it – which is our next task. To make it easier to read QQ-plots, it is nice to start with just considering histograms and/or density plots of the residuals and to see how that maps into this new display. We can obtain the residuals from the linear model using the `residuals` function on any linear model object.

```
par(mfrow=c(1,2))
eij<-residuals(lm2)
hist(eij, main="Histogram of residuals", cex.main=0.75)
plot(density(eij), main="Density plot of residuals", ylab="Density",
     xlab="Residuals", cex.main=0.75)
```

Figure 3.10 shows that there is a right skew present in the residuals for the prisoner rating data model that accounted for different means in the three groups, which is consistent with the initial assessment of some right skew in the plots of observations in each group.

A Quantile-Quantile plot (**QQ-plot**) shows the “match” of an observed distribution with a theoretical distribution, almost always the normal distribution. They are also known as Quantile Comparison, Normal Probability, or Normal Q-Q plots, with the last two names being specific to comparing results to a normal distribution. In this version<sup>5</sup>, the QQ-plots display the value of observed percentiles in the residual distribution on the y-axis versus the percentiles of a theoretical normal distribution on the x-axis. If the observed distribution of the residuals matches the shape of the normal distribution, then the plotted points should follow a 1-1 relationship. If the points follow the displayed straight line then that suggests that the residuals have a similar shape to a normal distribution. Some variation is expected around the line and some patterns of deviation are worse than others for our models, so you need to go beyond saying “it does not match a normal distribution”. It is best to be specific about the type of deviation you are detecting. And to do that, we need to practice interpreting some QQ-plots.

The QQ-plot of the linear model residuals from Figure 3.9 is extracted and enhanced it a little to make Figure 3.11 so we can just focus on it. We know from looking at the histogram that this is a slightly right skewed distribution. The QQ-plot places the observed **standardized<sup>6</sup> residuals** on the y-axis and the theoretical

<sup>5</sup>Along with multiple names, there is variation of what is plotted on the x and y axes and the scaling of the values plotted, increasing the challenge of interpreting QQ-plots. We are consistent about the x and y axis choices but different functions that make these plots in R do switch the axes.

<sup>6</sup>Here this means re-scaled so that they should have similar scaling to a standard normal with mean 0 and standard deviation 1. This does not change the shape of the distribution but can make outlier identification simpler – having a standardized residual

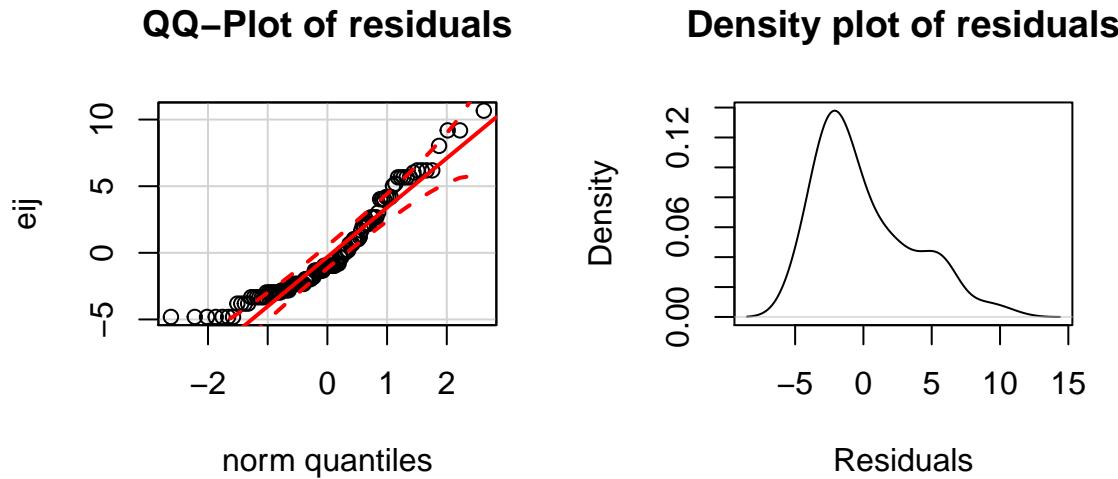


Figure 3.11: QQ-plot of residuals from linear model.

normal values on the x-axis. The most noticeable deviation from the 1-1 line is in the lower left corner of the plot. These are for the negative residuals (left tail) and there are many residuals at around the same value that are a little smaller than -1. If the distribution had followed the normal distribution here, the points would be on the 1-1 line and there would be some standardized residuals much smaller than -1.5. So we are not getting as much spread in the smaller residuals as we would expect in a normal distribution. If you go back to the histogram you can see that the smallest residuals are all stacked up and do not spread out like the left tail of a normal distribution should. In the right tail (positive) residuals, there is also a systematic lifting from the 1-1 line to larger values in the residuals than the normal would generate. For example, the point labeled as “82” (the 82nd observation in the data set) has a value of 3 in residuals but should actually be smaller (maybe 2.5) if the distribution was normal. Put together, this pattern in the QQ-plot suggests that the left tail is too compacted (too short) and the right tail is too spread out – this is the right skew we identified from the histogram and density curve!

Generally, when both tails deviate on the same side of the line (forming a sort of quadratic curve, especially in more extreme cases), that is evidence of a skew. To see some different potential shapes in QQ-plots, six different data sets are displayed in Figures 3.12 and 3.13. In each row, a QQ-plot and associated density curve are displayed. If the points are both above the 1-1 line in the lower and upper tails as in Figure 3.12(a), then the pattern is a right skew, here even more extreme than in the real data set. If the points are below the 1-1 line in both tails as in Figure 3.12(c), then the pattern is identified as a left skew. Skewed residual distributions (either direction) are problematic for models that assume normally distributed responses but not necessarily for our permutation approaches if all the groups have similar skewed shapes. The other problematic pattern is to have more spread than a normal curve as in Figure 3.12(e) and (f). This shows up with the points being below the line in the left tail (more extreme negative than expected by the normal) and the points being above the line for the right tail (more extreme positive than the normal predicts). We call these distributions **heavy-tailed** which can manifest as distributions with outliers in both tails or just a bit more spread out than a normal distribution. Heavy-tailed residual distributions can be problematic for our models as the variation is greater than what the normal distribution can account for and our methods might under-estimate the variability in the results. The opposite pattern with the left tail above the line and the right tail below the line suggests less spread (**lighter-tailed**) than a normal as in Figure 3.12(g) and (h). This pattern is relatively harmless and you can proceed with methods that assume normality safely as they will just be a little conservative.

---

more extreme than 5 or -5 would suggest a deviation from normality since we rarely see values that many standard deviations from the mean in a normal distribution. But mainly focus on the shape of the pattern in the QQ-plot.

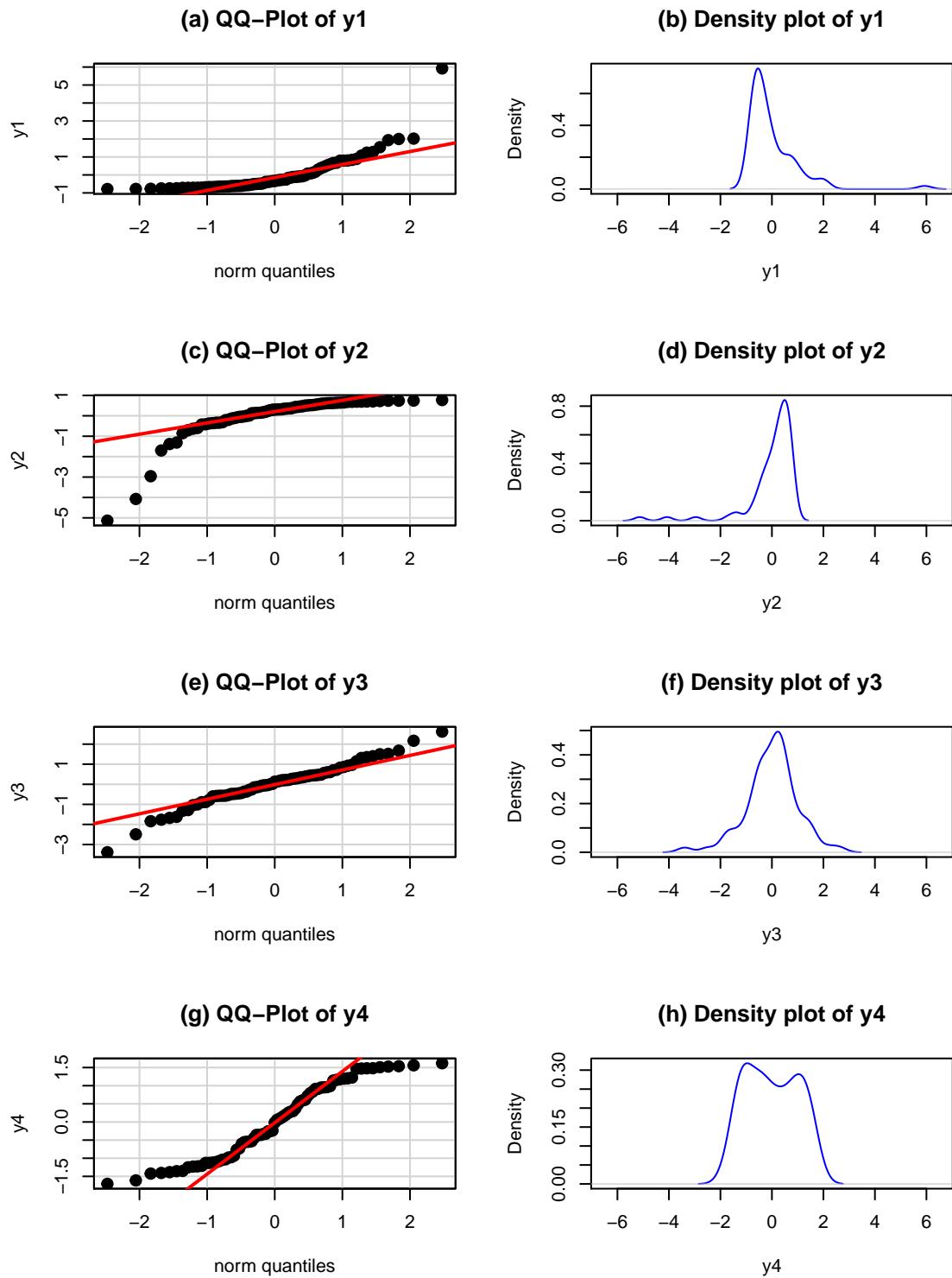


Figure 3.12: QQ-plots and density curves of four simulated distributions with different shapes.

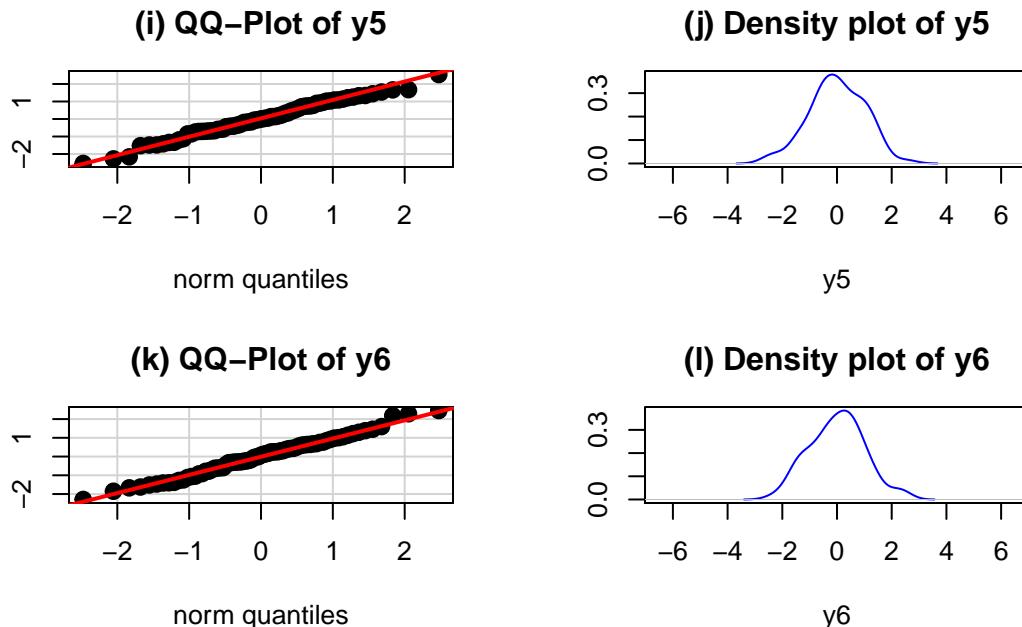


Figure 3.13: Two more simulated data sets, generated from normal distributions.

Finally, to help you calibrate expectations for data that are actually normally distributed, two data sets simulated from normal distributions are displayed in Figure 3.13. Note how neither follows the line exactly but that the overall pattern matches fairly well. **You have to allow for some variation from the line in real data sets** and focus on when there are really noticeable issues in the distribution of the residuals such as those displayed above. Again, you will never be able to prove that you have normally distributed residuals even if the residuals are all exactly on the line, but if you see QQ-plots as in Figure 3.12 you can encounter situations that provide evidence of clear violations of the normality assumption.

The last issues with assessing the assumptions in an ANOVA relates to situations where the methods are more or less *resistant*<sup>7</sup> to violations of assumptions. For reasons beyond the scope of this book, the parametric ANOVA F-test is more resistant to violations of the assumptions of the normality and equal variance assumptions if the design is balanced. A **balanced design** occurs when each group is measured the same number of times. The resistance decreases as the data set becomes less balanced, as the sample sizes in the groups are more different, so having close to balance is preferred to a more imbalanced situation if there is a choice available. There is some intuition available here – it makes some sense that you would have better results in comparing groups if the information available is similar in all the groups and none are relatively under-represented. We can check the number of observations in each group to see if they are equal or similar using the `tally` function from the `mosaic` package. This function is useful for being able to get counts of observations, especially for cross-classifying observations on two variables that is used in Chapter 5. For just a single variable, we use `tally(~x, data=...)`:

```
require(mosaic)
tally(~Attr, data=MockJury)
```

```
## Attr
##   Beautiful     Average Unattractive
##       39           38            37
```

So the sample sizes do vary among the groups and the design is technically not balanced, but it is also

---

<sup>7</sup>A resistant procedure is one that is not severely impacted by a particular violation of an assumption. For example, the median is resistant to the impact of an outlier.

very close to being balanced with only two more observations in the largest group compared to the smallest group size. This tells us that the  $F$ -test should have some resistance to violations of assumptions. This nearly balanced design, and the moderate sample size (over 37 per group is considered a good but not large sample), make the parametric and nonparametric approaches provide similar results in this data set even in the presence of the skewed residual error distribution.

### 3.5 Guinea pig tooth growth One-Way ANOVA example

A second example of the One-way ANOVA methods involves a study of length of odontoblasts (cells that are responsible for tooth growth) in 60 Guinea Pigs (measured in microns) from Crampton (1947).  $N = 60$  Guinea Pigs were obtained from a local breeder and each received one of three dosages (0.5, 1, or 2 mg/day) of Vitamin C via one of two delivery methods, Orange Juice (*OJ*) or ascorbic acid (the stuff in vitamin C capsules, called *VC* below) as the source of Vitamin C in their diets. Each guinea pig was randomly assigned to receive one of the six different treatment combinations possible (*OJ* at 0.5 mg, *OJ* at 1 mg, *OJ* at 2 mg, *VC* at 0.5 mg, *VC* at 1 mg, and *VC* at 2 mg). The animals were treated similarly otherwise and we can assume lived in separate cages and only one observation was taken for each guinea pig, so we can assume the observations are independent. We need to create a variable that combines the levels of delivery type (*OJ*, *VC*) and the dosages (0.5, 1, and 2) to use our One-Way ANOVA on the six levels. The `interaction` function can be used create a new variable that is based on combinations of the levels of other variables. Here a new variable is created in the `ToothGrowth` data.frame that we called `Treat` that provides a six-level grouping variable for our One-Way ANOVA to compare the combinations of treatments. To get a sense of the pattern of observations in the data set, the counts in `supp` (supplement type) and `dose` are provided.

```
data(ToothGrowth) #Available in Base R
require(mosaic)
tally(~supp,data=ToothGrowth) #Supplement Type (VC or OJ)

## supp
## OJ VC
## 30 30
tally(~dose,data=ToothGrowth) #Dosage level

## dose
## 0.5   1    2
##  20   20   20
#Creates a new variable Treat with 6 levels
ToothGrowth$Treat=with(ToothGrowth,interaction(supp,dose))

#New variable that combines supplement type and dosage
tally(~Treat,data=ToothGrowth)

## Treat
## OJ.0.5  VC.0.5  OJ.1    VC.1    OJ.2    VC.2
##      10      10      10      10      10      10
```

The `tally` function helps us to check for balance; this is a balanced design because the same number of guinea pigs ( $n_j = 10$  for  $j = 1, 2, \dots, 6$ ) were measured in each treatment combination.

With the variable `Treat` prepared, the first task is to visualize the results using boxplots and beanplots<sup>8</sup> (Figure 3.14) and generate some summary statistics for each group using `favstats`.

```
par(mfrow=c(1,2))
boxplot(len~Treat,data=ToothGrowth,ylab="Tooth Growth in microns")
```

---

<sup>8</sup>Note that to see all the group labels in the plot when making the figure, you have to widen the plot window before copying the figure out of R. You can resize the plot window using the small vertical and horizontal “=” signs in the grey bars that separate the different panels in RStudio.

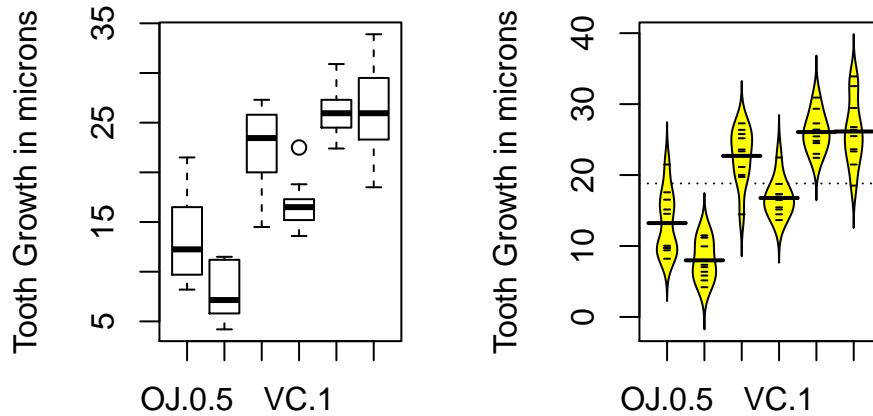


Figure 3.14: Boxplot and beanplot of tooth growth responses for the six treatment level combinations.

```
beanplot(len~Treat,data=ToothGrowth,log="",col="yellow",
        method="jitter",ylab="Tooth Growth in microns")
favstats(len~Treat,data=ToothGrowth)
```

Treat	min	Q1	median	Q3	max	mean	sd	n	missing
OJ.0.5	8.2	9.700	12.25	16.175	21.5	13.23	4.459709	10	0
VC.0.5	4.2	5.950	7.15	10.900	11.5	7.98	2.746634	10	0
OJ.1	14.5	20.300	23.45	25.650	27.3	22.70	3.910953	10	0
VC.1	13.6	15.275	16.50	17.300	22.5	16.77	2.515309	10	0
OJ.2	22.4	24.575	25.95	27.075	30.9	26.06	2.655058	10	0
VC.2	18.5	23.375	25.95	28.800	33.9	26.14	4.797731	10	0

Figure 3.14 suggests that the mean tooth growth increases with the dosage level and that OJ might lead to higher growth rates than VC except at a dosage of 2 mg/day. The variability around the means looks to be small relative to the differences among the means, so we should expect a small p-value from our  $F$ -test. The design is balanced as noted above ( $n_j = 10$  for all six groups) so the methods are somewhat resistant to impacts from non-normality and non-constant variance. There is some suggestion of non-constant variance in the plots but this will be explored further below when we can remove the difference in the means and combine all the residuals together. There might be some skew in the responses in some of the groups but there are only 10 observations per group so skew in the boxplots could be generated by impacts of very few of the observations.

Now we can apply our 6+ steps for performing a hypothesis test with these observations. The initial step is deciding on the claim to be assessed and the test statistic to use. This is a six group situation with a quantitative response, identifying it as a One-Way ANOVA where we want to test a null hypothesis that all the groups have the same population mean, at least to start. We will use a 5% significance level.

1. Hypotheses:  $H_0 : \mu_{OJ.0.5} = \mu_{VC.0.5} = \mu_{OJ.1} = \mu_{VC.1} = \mu_{OJ.2} = \mu_{VC.2}$  vs  $H_A : \text{Not all } \mu_j \text{ equal}$

- The null hypothesis could also be written in reference-coding as below since OJ.0.5 is chosen as the baseline group (discussed below).

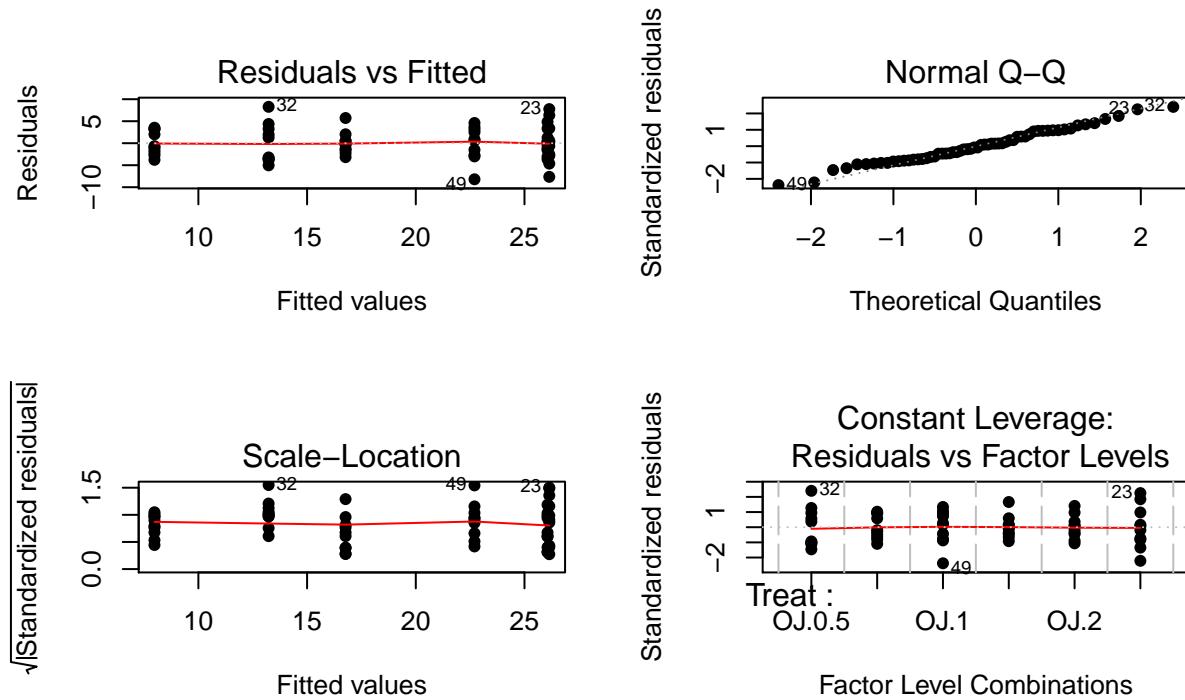


Figure 3.15: Diagnostic plots for the toothgrowth model.

- $H_0 : \tau_{VC0.5} = \tau_{OJ1} = \tau_{VC1} = \tau_{OJ2} = \tau_{VC2} = 0$
- The alternative hypothesis can be left a bit less specific:
  - $H_A : \text{Not all } \tau_j \text{ equal 0}$

## 2. Validity conditions:

- Independence:
  - This is where the separate cages note above is important. Suppose that there were cages that contained multiple animals and they competed for food or could share illness or levels of activity. The animals in one cage might be systematically different from the others and this “clustering” of observations would present a potential violation of the independence assumption. If the experiment had the animals in separate cages, there is no clear dependency in the design of the study and we can assume that there is no problem with this assumption.
- Constant variance:
  - As noted above, there is some indication of a difference in the variability among the groups in the boxplots and beanplots but the sample size was small in each group. We need to fit the linear model to get the other diagnostic plots to make an overall assessment.

```
m2<-lm(len~Treat,data=ToothGrowth)
par(mfrow=c(2,2))
plot(m2,pch=16)
```

- The Residuals vs Fitted panel in Figure 3.15 shows some difference in the spreads but the spread is not that different between the groups.
- The Scale–Location plot also shows just a little less variability in the group with the smallest fitted value but the spread of the groups looks fairly similar in this alternative scaling.

- Put together, the evidence for non-constant variance is not that strong and we can assume that there is at least not a major problem with this assumption.
- Normality of residuals:
  - The Normal Q-Q plot shows a small deviation in the lower tail but nothing that we wouldn't expect from a normal distribution. So there is no evidence of a problem with the normality assumption in the upper right panel of Figure 3.15.

### 3. Calculate the test statistic:

- The ANOVA table for our model follows, providing an  $F$ -statistic of 41.557:

```
anova(m2)
```

```
## Analysis of Variance Table
##
## Response: len
##             Df  Sum Sq Mean Sq F value    Pr(>F)
## Treat       5 2740.10 548.02 41.557 < 2.2e-16
## Residuals 54  712.11   13.19
```

### 4. Find the p-value:

- There are two options here, especially since it seems that our assumptions about variance and normality are not violated (note that we do not say “met” – we just have no clear evidence against them). The parametric and nonparametric approaches should provide similar results here.
- The parametric approach is easiest – the p-value comes from the previous ANOVA table as  $<2e-16$ . First, note that this is in scientific notation that is a compact way of saying that the p-value here is  $2.2 \times 10^{-16}$  or 0.00000000000000022. When you see  $2.2e-16$  in R output, it also means that the calculation is at the numerical precision of the computer. What R is really trying to report is that this is a very small number. **When you encounter p-values that are smaller than 0.0001, you should just report that the p-value<0.0001** Do not report that it is 0 as this gives the false impression that there is no chance of the result occurring when it is just a really small probability. This distribution (the distribution of the test statistic if the null hypothesis is true).
- The nonparametric approach is not too hard so we can compare the two approaches here as well:

```
Tobs <- anova(lm(len~Treat,data=ToothGrowth))[1,4]; Tobs
```

```
## [1] 41.55718
par(mfrow=c(1,2))
B<- 1000
Tstar<-matrix(NA,nrow=B)
for (b in (1:B)){
  Tstar[b]<-anova(lm(len~shuffle(Treat),data=ToothGrowth))[1,4]
}
pdata(Tstar,Tobs,lower.tail=F)
```

```
## [1] 0
hist(Tstar,xlim=c(0,Tobs+3))
abline(v=Tobs,col="red",lwd=3)
plot(density(Tstar),xlim=c(0,Tobs+3),main="Density curve of Tstar")
abline(v=Tobs,col="red",lwd=3)
```

- The permutation p-value was reported as 0. This should be reported as p-value<0.001 since we did 1000 permutations and found that none of the permuted  $F$ -statistics,  $F^*$ , were larger than the observed  $F$ -statistic of 41.56. The permuted results do not exceed 6 as seen in Figure 3.16,

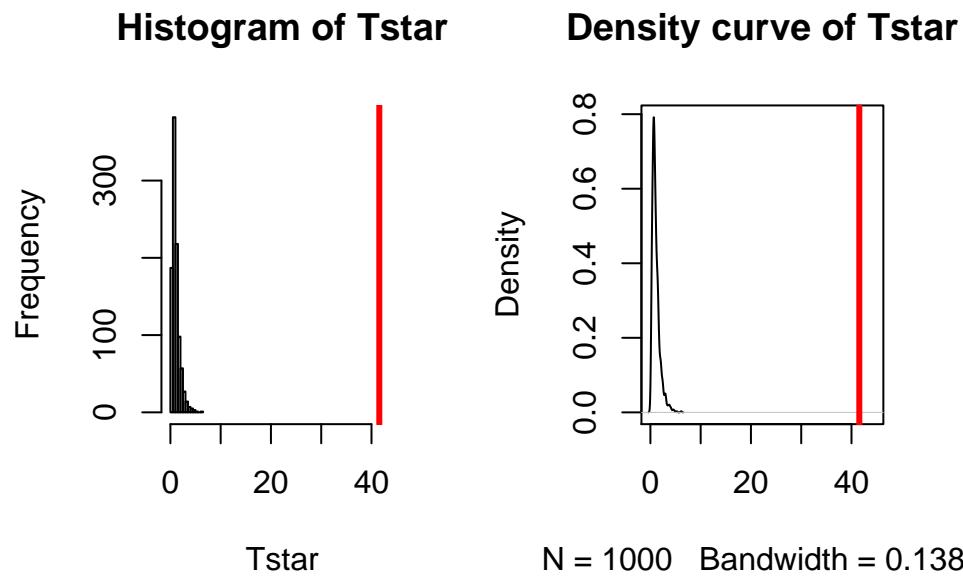


Figure 3.16: Histogram and density curve of permutation distribution for  $F$ -statistic for tooth growth data. Observed test statistic in bold, vertical line at 41.56.

so the observed result is *really unusual* relative to the null hypothesis. As suggested previously, the parametric and nonparametric approaches should be similar here and they were.

#### 5. Make a decision:

- Reject  $H_0$  since the p-value is less than 5%.

#### 6. Write a conclusion:

- There is evidence at the 5% significance level that the different treatments (combinations of OJ/VC and dosage levels) **cause some** difference in the **true** mean tooth growth for **these** guinea pigs.
  - We can make the causal statement of the treatment causing differences because the treatments were randomly assigned but these inferences only apply to these guinea pigs since they were not randomly selected from a larger population.
  - Remember that we are making inferences to the population or true means and not the sample means and want to make that clear in any conclusion. When there is not a random sample from a population it is more natural to discuss the true means since we can't extend to the population values.
  - The alternative is that there is some difference in the true means – be sure to make the wording clear that you aren't saying that all the means differ. In fact, if you look back at Figure 3.14, the means for the 2 mg dosages look almost the same so we will have a tough time arguing that all groups differ. The  $F$ -test is about finding evidence of some means. The next section will provide some additional tools to get more specific about the source of those detected differences.

Before we leave this example, we should revisit our model estimates and interpretations. The default model parameterization is into the reference-coding. Running the model `summary` function on `m2` provides the estimated coefficients:

```
summary(m2)

##
## Call:
## lm(formula = len ~ Treat, data = ToothGrowth)
##
## Residuals:
##   Min     1Q Median     3Q    Max 
## -8.20 -2.72 -0.27  2.65  8.27 
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) 13.230     1.148 11.521 3.60e-16  
## TreatVC.0.5 -5.250     1.624 -3.233 0.00209   
## TreatOJ.1    9.470     1.624  5.831 3.18e-07  
## TreatVC.1    3.540     1.624  2.180 0.03365   
## TreatOJ.2   12.830     1.624  7.900 1.43e-10  
## TreatVC.2   12.910     1.624  7.949 1.19e-10  
##
## Residual standard error: 3.631 on 54 degrees of freedom
## Multiple R-squared:  0.7937, Adjusted R-squared:  0.7746 
## F-statistic: 41.56 on 5 and 54 DF,  p-value: < 2.2e-16
```

For some practice with the reference coding used in these models, let's find the estimates for observations for a couple of the groups. To work with the parameters, you need to start with diagnosing the baseline category that was used by considering which level is not displayed in the output. The `levels` function can list the groups in a categorical variable and their coding in the data set. The first level is usually the baseline category but you should check this in the model summary as well.

```
levels(ToothGrowth$Treat)
```

```
## [1] "OJ.0.5" "VC.0.5" "OJ.1"   "VC.1"   "OJ.2"   "VC.2"
```

There is a VC.0.5 in the second row of the model summary, but there is no row for OJ.0.5 and so this must be the baseline category. That means that the fitted value or model estimate for the OJ at 0.5 mg/day group is the same as the (Intercept) row or  $\hat{\alpha}$ , estimating a mean tooth growth of 13.23 microns when the pigs get OJ at a 0.5 mg/day dosage level. You should always start with working on the baseline level in a reference-coded model. To get estimates for any other group, then you can use the (Intercept) estimate and add the deviation for the group of interest. For VC.0.5, the estimated mean tooth growth is  $\hat{\alpha} + \hat{\gamma}_2 = \hat{\alpha} + \hat{\gamma}_{VC0.5} = 13.23 + (-5.25) = 7.98$  microns. It is also potentially interesting to directly interpret the estimated difference (or deviation) between OJ0.5 (the baseline) and VC0.5 (group 2) that is  $\hat{\gamma}_{VC0.5} = -5.25$ : we estimate that the mean tooth growth in VC0.5 is 5.25 microns shorter than it is in OJ0.5. This and many other direct comparisons of groups are likely of interest to researchers involved in studying the impacts of these supplements on tooth growth and the next section will show us how to do that (correctly!).

The reference-coding is still going to feel a little uncomfortable so the comparison to the cell-means model and exploring the effect plot can help to reinforce that both models patch together the same estimated means for each group. For example, we can find our estimate of 7.98 microns for the VC0.5 group in the output and Figure 3.17. Also note that Figure 3.17 is the same whether you plot the results from `m2` or `m3`.

```
m3<-lm(len~Treat-1,data=ToothGrowth)
```

```
summary(m3)
```

```
##
## Call:
## lm(formula = len ~ Treat - 1, data = ToothGrowth)
##
## Residuals:
##   Min     1Q Median     3Q    Max 
## -8.20 -2.72 -0.27  2.65  8.27 
```

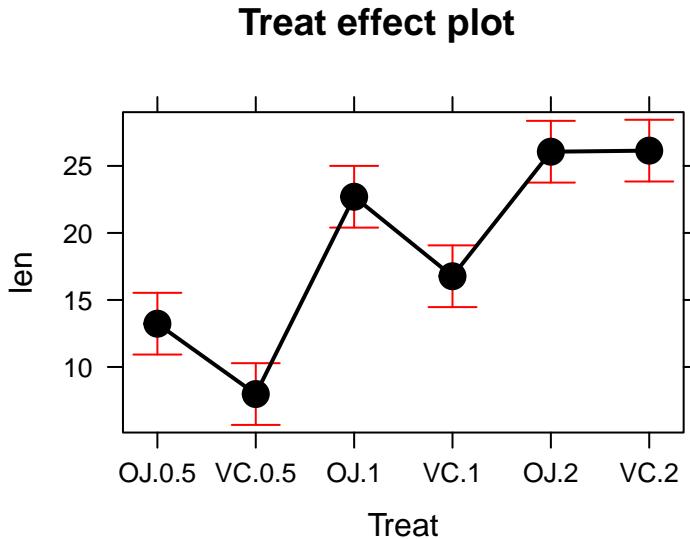


Figure 3.17: Effect plot of the One-Way ANOVA model for the toothgrowth data.

```

## 
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## TreatOJ.0.5    13.230    1.148 11.521 3.60e-16
## TreatVC.0.5     7.980    1.148  6.949 4.98e-09
## TreatOJ.1      22.700    1.148 19.767 < 2e-16
## TreatVC.1      16.770    1.148 14.604 < 2e-16
## TreatOJ.2      26.060    1.148 22.693 < 2e-16
## TreatVC.2      26.140    1.148 22.763 < 2e-16
##
## Residual standard error: 3.631 on 54 degrees of freedom
## Multiple R-squared:  0.9712, Adjusted R-squared:  0.968
## F-statistic: 303 on 6 and 54 DF,  p-value: < 2.2e-16
par(mfrow=c(1,1))
plot(allEffects(m2))

```

## 3.6 Multiple (pair-wise) comparisons using Tukey's HSD and the compact letter display

With evidence that the true means are likely not all equal, many researchers want to know which groups show evidence of differing from one another. This provides information on the source of the overall difference that was detected and detailed information on which groups differed from one another. Because this is a shot-gun/unfocused sort of approach, some people think it is an over-used procedure. Others feel that it is an important method of addressing detailed questions about group comparisons in a valid way. For example, we might want to know if OJ is dosage level and these methods will allow us to get an answer to this sort of question. It also will test for differences between the OJ,0.5 and VC,2 groups and every other pair of levels that you can construct. This method actually takes us back to the methods in Chapter 2 where we compared the means of two groups except that we need to deal with potentially many pair-wise comparisons, making an adjustment to account for that inflation in Type I errors that occurs due to many tests being performed at the same time. There are many different statistical methods to make all the pair-wise comparisons, but we will employ the most commonly used one, called *Tukey's Honest Significant Difference* (Tukey's HSD)

method<sup>9</sup>. The name suggests that not using it could lead to a dishonest answer and that it will give you an honest result. It is more that if you don't do some sort of correction for all the tests you are performing, you might find some *spurious*<sup>10</sup> results. There are other methods that could be used to do a similar correction and also provide "honest" inferences; we are just going to learn one of them.

Generally, the general challenge in this situation is that if you perform many tests at the same time, you inflate the Type I error rate. We can define the *family-wise error rate* as the probability that at least one error is made on a set of tests or, more compactly,  $\Pr(\text{At least 1 error is made})$  where  $\Pr()$  is the probability of an event occurring. The family-wise error is meant to capture the overall situation in terms of measuring the likelihood of making a mistake if we consider many tests, each with some chance of making their own mistake, and focus on how often we make at least one error when we do many tests. A quick probability calculation shows the magnitude of the problem. If we start with a 5% significance level test, then  $\Pr(\text{Type I error on one test}) = 0.05$  and the  $\Pr(\text{no errors made on one test}) = 0.95$ , by definition. This is our standard hypothesis testing situation. Now, suppose we have  $m$  independent tests, then

$$\begin{aligned}\Pr(\text{make at least 1 Type I error given all null hypotheses are true}) \\ = 1 - \Pr(\text{no errors made}) \\ = 1 - 0.95^m.\end{aligned}$$

Figure ??fig:Figure3-18) shows how the probability of having at least one false detection grows rapidly with the number of tests. The plot stops at 100 tests since it is effectively a 100% chance of at least one false detection. It might seem like doing 100 tests is a lot, but in Genetics research it is possible to consider situations where millions of tests are considered so these are real issues to be concerned about in many situations. Researchers want to make sure that when they report a "significant" result that it is really likely to be a real result and will show up as a difference in the next data set they collect.<sup>11</sup>

In pair-wise comparisons between all the pairs of means in a One-Way ANOVA, the number of tests is based on the number of pairs. We can calculate the number of tests using  $J$  choose 2,  $\binom{J}{2}$ , to get the number of unique pairs of size 2 that we can make out of  $J$  individual treatment levels. We don't need to explore the combinatorics formula for this, as the `choose` function can give us the answers:

```
choose(3, 2)
```

```
## [1] 3
choose(4, 2)
```

```
## [1] 6
choose(5, 2)
```

```
## [1] 10
choose(6, 2)
```

```
## [1] 15
```

So if you have three groups (prisoner rating study), there are 3 unique pairs to compare. For six groups, like in the guinea pig study, we have to consider 15 tests to compare all the unique pairs of groups. 15 tests seems like enough that we should be worried about inflated family-wise error rates. Fortunately, the Tukey's HSD method controls the family-wise error rate at your specified level (say 0.05) across any number of pair-wise comparisons. This means that the overall rate of at least one Type I error is controlled at the specified significance level, often 5%. To do this, each test must use a slightly more conservative cut-off than if just one test is performed and the procedure helps us figure out how much more conservative we need to be.

<sup>9</sup>When this procedure is used with unequal group sizes it is also sometimes called Tukey-Kramer's method.

<sup>10</sup>We often use "spurious" to describe falsely rejected null hypotheses, but there are also called false detections.

<sup>11</sup>Many researchers are now collecting multiple data sets to use in a single study and using one data set to identify interesting results and then using a validation or test data set that they withheld from initial analysis to verify that the first results are also present in that second data set.

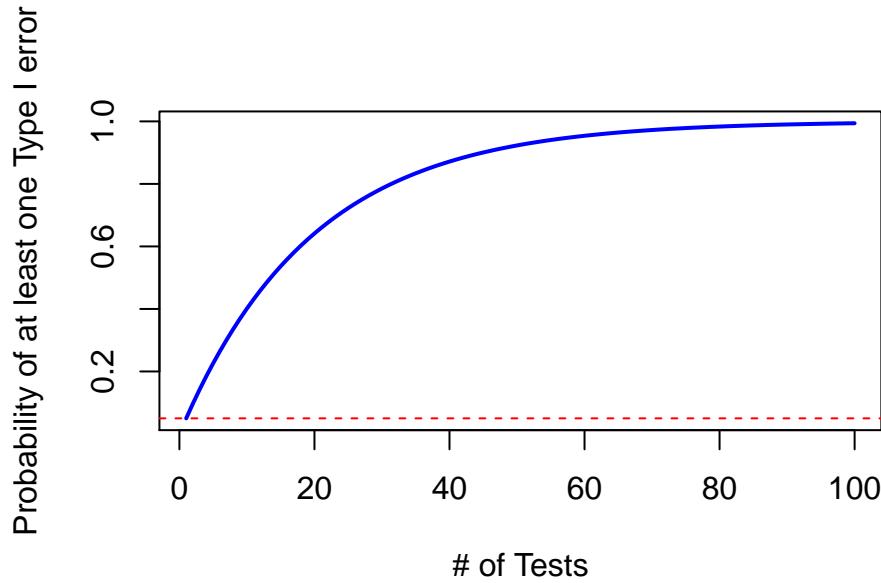


Figure 3.18: Plot of family-wise error rate as the number of tests performed increases. Dashed line indicates 0.05.

Tukey's HSD starts with focusing on the difference between the groups with the largest and smallest means ( $\bar{y}_{max} - \bar{y}_{min}$ ). If  $(\bar{y}_{max} - \bar{y}_{min}) \leq$  Margin of Error for the difference in the means, then all other pairwise differences, say  $|\bar{y}_j - \bar{y}_{j'}|$ , for two groups  $j$  and  $j'$ , will be less than or equal to that margin of error. This also means that any confidence intervals for any difference in the means will contain 0. Tukey's HSD selects a critical value so that  $(\bar{y}_{max} - \bar{y}_{min})$  will be less than the margin of error in 95% of data sets drawn from populations with a common mean. This implies that in 95% of data sets in which all the population means are the same, all confidence intervals for differences in pairs of means will contain 0. Tukey's HSD provides confidence intervals for the difference in true means between groups  $j$  and  $j'$ ,  $\mu_j - \mu_{j'}$ , for all pairs where  $j \neq j'$ , using

$$(\bar{y}_j - \bar{y}_{j'}) \mp \frac{q^*}{\sqrt{2}} \sqrt{\text{MS}_E \left( \frac{1}{n_j} + \frac{1}{n_{j'}} \right)}$$

where  $\frac{q^*}{\sqrt{2}} \sqrt{\text{MS}_E \left( \frac{1}{n_j} + \frac{1}{n_{j'}} \right)}$  is the margin of error for the intervals. The distribution used to find the multiplier,  $q^*$ , for the confidence intervals is available in the `qtukey` function and generally provides a slightly larger multiplier than the regular  $t^*$  from our two-sample  $t$ -based confidence interval discussed in Chapter 2. We will use the `confint`, `cld`, and `plot` functions applied to output from the `glht` function (all from the `multcomp` package; Hothorn and Westfall (2008), (Hothorn et al., 2016)) to easily get the required comparisons from our ANOVA model. Unfortunately, its code format is a little complicated – but there are just two places to modify the code, by including the model name and after `mcp` (stands for multiple comparisons) in the `linfct` option, you need to include the explanatory variable name as `VARIABLENAME="Tukey"`. The last part is to get the TukeyHSD multiple comparisons run on our explanatory variable. Once we obtain the intervals, we can use them to test  $H_0 : \mu_j = \mu_{j'}$  vs  $HA : \mu_j \neq \mu_{j'}$  by assessing whether 0 is in the confidence interval for each pair. If 0 is in the interval, then there is no evidence of a difference for that pair. If 0 is not in the interval, then we reject  $H_0$  and have evidence *at the specified family-wise significance level* of a difference for that pair. You will see a switch to using the word “detection” to describe rejected null hypotheses of no

difference as it can help to write up these results. The following code provides the numerical and graphical<sup>12</sup> results of applying Tukey's HSD to the linear model for the Guinea Pig data:

```
par(mfrow=c(1,1))
require(multcomp)
Tm2 <- glht(m2, linfct = mcp(Treat = "Tukey"))
confint(Tm2)

##
##   Simultaneous Confidence Intervals
##
## Multiple Comparisons of Means: Tukey Contrasts
##
##
## Fit: lm(formula = len ~ Treat, data = ToothGrowth)
##
## Quantile = 2.9545
## 95% family-wise confidence level
##
##
## Linear Hypotheses:
##                         Estimate lwr      upr
## VC.0.5 - OJ.0.5 == 0 -5.2500 -10.0482 -0.4518
## OJ.1 - OJ.0.5 == 0  9.4700  4.6718 14.2682
## VC.1 - OJ.0.5 == 0 3.5400 -1.2582  8.3382
## OJ.2 - OJ.0.5 == 0 12.8300  8.0318 17.6282
## VC.2 - OJ.0.5 == 0 12.9100  8.1118 17.7082
## OJ.1 - VC.0.5 == 0 14.7200  9.9218 19.5182
## VC.1 - VC.0.5 == 0 8.7900  3.9918 13.5882
## OJ.2 - VC.0.5 == 0 18.0800 13.2818 22.8782
## VC.2 - VC.0.5 == 0 18.1600 13.3618 22.9582
## VC.1 - OJ.1 == 0   -5.9300 -10.7282 -1.1318
## OJ.2 - OJ.1 == 0   3.3600 -1.4382  8.1582
## VC.2 - OJ.1 == 0   3.4400 -1.3582  8.2382
## OJ.2 - VC.1 == 0   9.2900  4.4918 14.0882
## VC.2 - VC.1 == 0   9.3700  4.5718 14.1682
## VC.2 - OJ.2 == 0   0.0800 -4.7182  4.8782
old.par <- par(mai=c(1,2,1,1)) #Makes room on the plot for the group names
plot(Tm2)
```

Figure 3.19 contains confidence intervals for the difference in the means for all 15 pairs of groups. For example, the first row in the plot contains the confidence interval OJ.0.5). In the numerical output, you can find that this 95% family-wise confidence interval goes from -10.05 to -0.45 microns (lwr and upr in the numerical output provide the CI endpoints). This interval does not contain 0 since its upper end point is -0.45 microns and so we can now say that there is evidence that OJ and VC have different true mean growth rates at the 0.5 mg dosage level. We can go further and say that we are 95% confident that the difference in the true mean tooth growth between VC.0.5 and OJ.0.5 (VC.0.5-OJ.0.5) is between -10.05 and -0.45 microns, after adjusting for comparing all the pairs of groups. But there are fourteen more similar intervals...

If you put all these pair-wise tests together, you can generate an overall interpretation of Tukey's HSD results that discusses sets of groups that are not detectably different from one another and those groups that were distinguished from other sets of groups. To do this, start with listing out the groups that do not appear to be different (CIs contain 0), which, here, only occurs for four of the pairs. The CIs that contain 0 are for the pairs VC.1 and OJ.0.5, OJ.2 and OJ.1, VC.2 and OJ.1, and, finally, VC.2 and OJ.2. So VC.2, OJ.1, and OJ.2 are all not detectably different from each other and VC.1 and OJ.0.5 are also not detectably

---

<sup>12</sup>The plot of results usually contains all the labels of groups but if the labels are long or there are many groups, sometimes the row labels are hard to see even with re-sizing the plot to make it taller in RStudio. The numerical output is useful as a guide to help you read the plot.

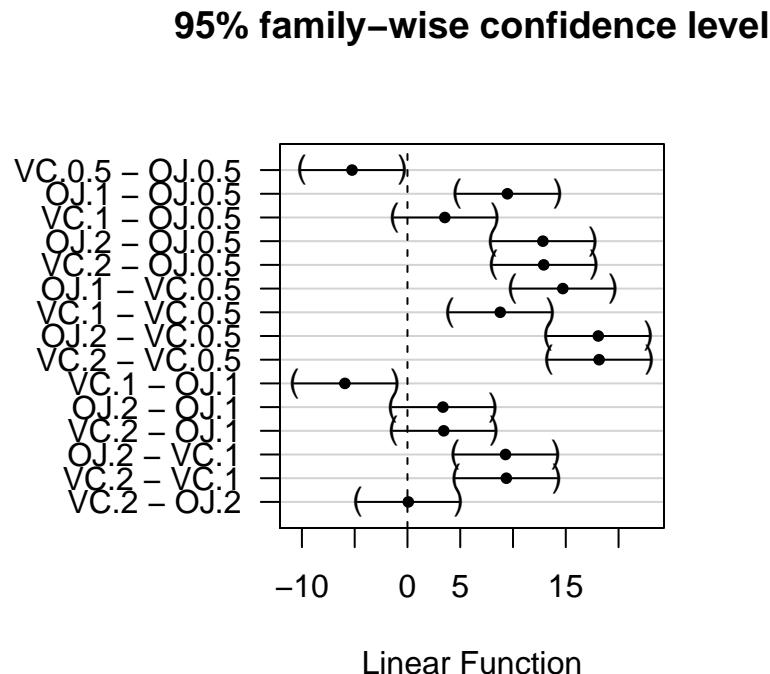


Figure 3.19: Graphical display of pair-wise comparisons from Tukey's HSD for the Guinea Pig data. Any confidence intervals that do not contain 0 provide evidence of a difference in the groups.

different. If you look carefully, VC.0.5 is detected as different from every other group. So there are basically three sets of groups that can be grouped together as “similar”: VC.2, OJ.1, and OJ.2; VC.1 and OJ.0.5; and VC.0.5. Sometimes groups overlap with some levels not being detectably different from other levels that belong to different groups and the story is not as clear as it is in this case. An example of this sort of overlap is seen in the next section.

There is a method that many researchers use to more efficiently generate and report these sorts of results that is called a ***compact letter display*** (CLD, Piepho (2004)). The `cld` function can be applied to the results from `glht` to generate the CLD that we can use to provide a “simple” summary of the sets of groups. In this discussion, we define a **set as a union of different groups that can contain one or more members** and the member of these groups are the different treatment levels.

```
cld(Tm2)
```

```
## OJ.0.5 VC.0.5  OJ.1   VC.1   OJ.2   VC.2
##    "b"    "a"    "c"    "b"    "c"    "c"
```

Groups with the same letter are not detectably different (are in the same set) and groups that are detectably different get different letters (are in different sets). Groups can have more than one letter to reflect “overlap” between the sets of groups and sometimes a set of groups contains only a single treatment level (VC.0.5 is a set of size 1). Note that if the groups have the same letter, this does not mean they are the same, just that there is **no evidence of a difference for that pair**. If we consider the previous output for the CLD, the “a” set contains VC.0.5, the “b” set contains OJ.1, OJ.2, and VC.2, and the “c” set contains OJ.0.5 and VC.1. These are exactly the groups of treatment levels that we obtained by going through all fifteen pairwise results.

One benefit of this work is that the CLD letters can be added to a beanplot to help fully report the results and understand the sorts of differences Tukey's HSD detected.

The lines with text in them are involved in placing text on the figure but are something you could do in

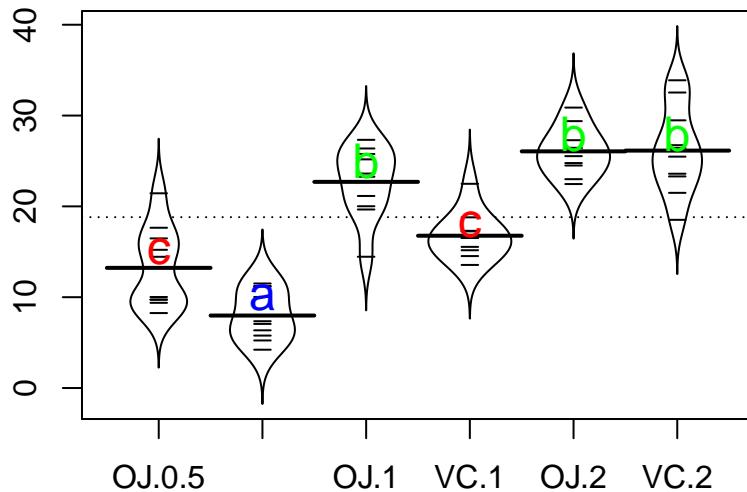


Figure 3.20: Beanplot of tooth growth by group with Tukey’s HSD compact letter display.

image editing software just as easily. Figure 3.20 enhances the discussion by showing that the “**a**” group with VC.0.5 had the lowest average tooth growth, the “**c**” group had intermediate tooth growth for treatments OJ.0.5 and VC.1, and the highest growth rates came from OJ.1, OJ.2, and VC.2. Even though VC.2 had the highest average growth rate, we are not able to prove that its true mean is any higher than the other groups labeled with “**b**”. Hopefully the ease of getting to the story of the Tukey’s HSD results from a plot like this explains why it is common to report results using these methods instead of reporting 15 confidence intervals.

There are just a couple of other details to mention on this set of methods. First, note that we interpret the set of confidence intervals simultaneously: We are 95% confident that **ALL** the intervals contain the respective differences in the true means (this is a **family-wise interpretation**). These intervals are adjusted from our regular 2 sample  $t$  intervals from Chapter 2 to allow this stronger interpretation. Specifically, they are wider. Second, if sample sizes are unequal in the groups, Tukey’s HSD is conservative and provides a family-wise error rate that is lower than the nominal (or specified) level. In other words, it fails less often than expected and the intervals provided are a little wider than needed, containing all the pairwise differences at higher than the nominal confidence level of (typically) 95%. Third, this is a parametric approach and violations of normality and constant variance will push the method in the other direction, potentially making the technique dangerously liberal. Nonparametric approaches to this problem are also possible, but will not be considered here.

### 3.7 Pair-wise comparisons for Prisoner Rating data

In our previous work with the prisoner rating data, the overall ANOVA test provided only marginal evidence of some difference in the true means across the three groups with a  $p$ -value=0.067. Tukey’s HSD does not require you to find a small from your overall  $F$ -test to employ the methods but if you apply it to situations with  $p$ -values larger than your *a priori* significance level, you are unlikely to find any pairs that are detected

as being different. Some statisticians suggest that you shouldn't employ follow-up tests such as Tukey's HSD when there is not sufficient evidence to reject the overall null hypothesis and would be able to reasonably criticize the following results. But for the sake of completeness, we can find the pair-wise comparison results at our typical 95% family-wise confidence level in this situation, with the three confidence intervals displayed in Figure 3.21.

```
lm2<-lm(Years~Attr, data=MockJury)
require(multcomp)
Tm2 <- glht(lm2, linfct = mcp(Attr = "Tukey"))
confint(Tm2)

##
##  Simultaneous Confidence Intervals
##
## Multiple Comparisons of Means: Tukey Contrasts
##
##
## Fit: lm(formula = Years ~ Attr, data = MockJury)
##
## Quantile = 2.3751
## 95% family-wise confidence level
##
##
## Linear Hypotheses:
##                               Estimate lwr      upr
## Average - Beautiful == 0     -0.3596  -2.2969  1.5776
## Unattractive - Beautiful == 0   1.4775  -0.4730  3.4280
## Unattractive - Average == 0    1.8371  -0.1258  3.8001
cld(Tm2)

##      Beautiful      Average Unattractive
##            "a"          "a"           "a"
old.par <- par(mai=c(1,2.5,1,1)) #Makes room on the plot for the group names
plot(Tm2)
```

At the family-wise 5% significance level, there are no pairs that are detectably different – they all get the same letter of “a”. Now we will produce results for the reader that thought a 10% significance was suitable for this application before seeing any of the results. We just need to change the confidence level or significance level that the CIs or tests are produced with inside the functions. For the `confint` function, the `level` option is the confidence level and for the `cld`, it is the family-wise significance level. Note that 90% confidence corresponds to a 10% significance level.

```
confint(Tm2,level=0.9)

##
##  Simultaneous Confidence Intervals
##
## Multiple Comparisons of Means: Tukey Contrasts
##
##
## Fit: lm(formula = Years ~ Attr, data = MockJury)
##
## Quantile = 2.0739
## 90% family-wise confidence level
##
##
## Linear Hypotheses:
##                               Estimate lwr      upr
## Average - Beautiful == 0     -0.3596  -2.0513  1.3320
## Unattractive - Beautiful == 0   1.4775  -0.2257  3.1806
```

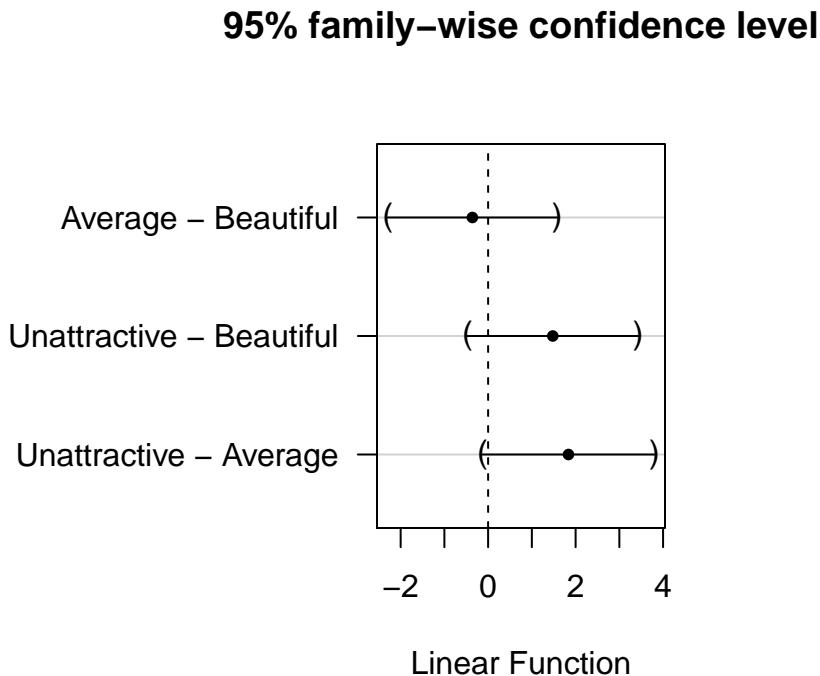


Figure 3.21: Tukey's HSD confidence interval results at the 95% family-wise confidence level.

```
## Unattractive - Average == 0      1.8371   0.1231  3.5512
cld(Tm2,level=0.1)
```

```
##      Beautiful      Average Unattractive
##      "ab"          "a"        "b"
old.par <- par(mai=c(1,2.5,1,1)) #Makes room on the plot for the group names
plot(confint(Tm2,level=.9))
```

With family-wise 10% significance and 90% confidence levels, the *Unattractive* and *Average* picture groups are detected as being different but the *Average* group is not detected as different from *Beautiful* and *Beautiful* is not detected to be different from *Unattractive*. This leaves the “overlap” of groups across the sets of groups that was noted earlier. The *Beautiful* level is not detected as being dissimilar from levels in two different sets and so gets two different letters.

The beanplot (Figure 3.23) helps to clarify some of the reasons for this set of results. The detection of a difference between *Average* and *Unattractive* just barely occurs and the mean for *Beautiful* is between the other two so it ends up not being detectably different from either one. This sort of overlap is actually a fairly common occurrence in these sorts of situations so be prepared a mixed set of letters for some levels.

```
old.par <- par(mai=c(0.5,1,1,1))
beanplot(Years~Attr,data=MockJury,log="",col="white",method="jitter")
text(c(1),c(5),"ab",col="blue",cex=1.5)
text(c(2),c(4.8),"a",col="green",cex=1.5)
text(c(3),c(6.5),"b",col="red",cex=1.5)
```

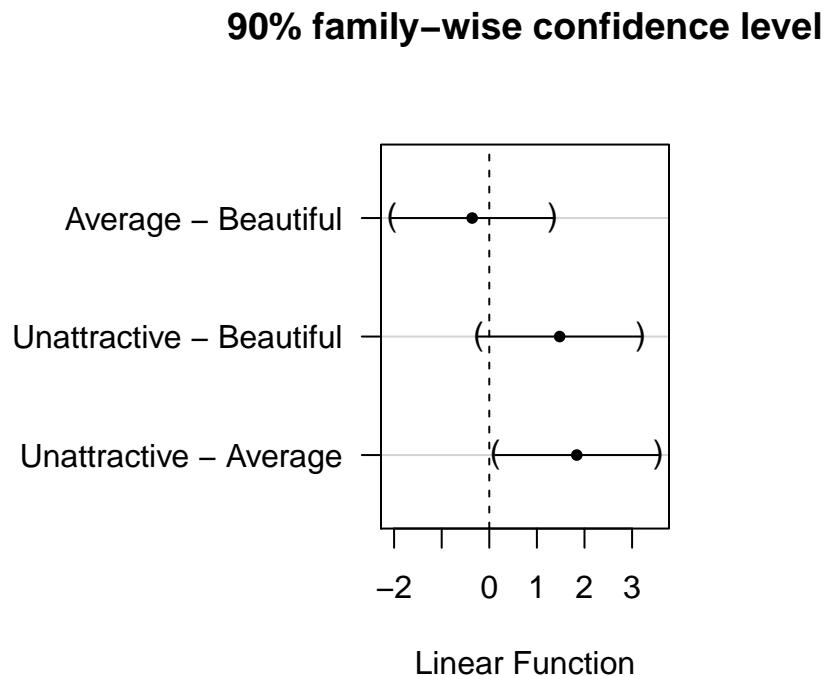


Figure 3.22: Tukey's HSD 90% family-wise confidence intervals.

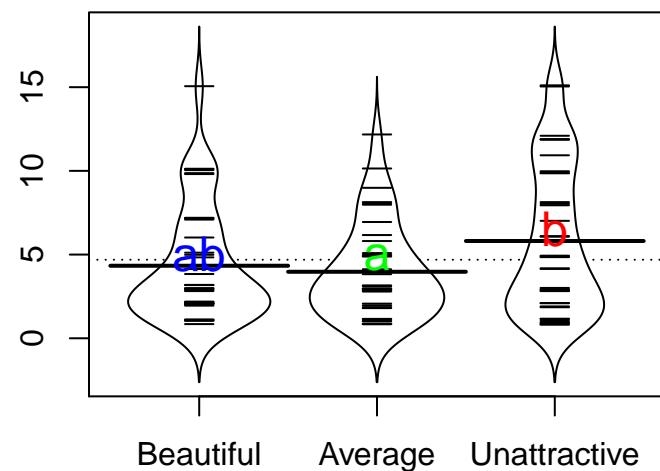


Figure 3.23: Beanplot of sentences with compact letter display results from 10% family-wise significance level Tukey's HSD. *Average* and *Unattractive* picture groups are detected as being different and are displayed as belonging to different groups. *Beautiful* picture responses are not detected as different from the other two groups.

## 3.8 Chapter Summary

In this chapter, we explored methods for comparing a quantitative response across  $J$  groups ( $J \geq 2$ ), with what is called the One-Way ANOVA procedure. The initial test is based on assessing evidence against a null hypothesis of no groups. There are two different methods for estimating these One-Way ANOVA models: the cell-means model and the reference-coded versions of the model. There are times when either model will be preferred, but for the rest of the text, the reference coding is used (sorry!). The ANOVA  $F$ -statistic, often presented with underlying information in the ANOVA table, provides a method of assessing evidence against the null hypothesis either using permutations or via the  $F$ -distribution. Pair-wise comparisons using Tukey's HSD provide a method for comparing all the groups and are a nice complement to the overall ANOVA results. A compact letter display was shown that enhanced the interpretation of Tukey's HSD result.

In the guinea pig example, we are left with some lingering questions based on these results. It appears that the effect of *dosage* changes as a function of the *delivery method* (OJ, VC) because the size of the differences between OJ and VC change for different dosages. These methods can't directly assess the question of whether the effect of delivery method is the same or not across the different dosages. In chapter 4, the two variables, *Dosage* and *Delivery method* are modeled as two separate variables so we can consider their effects both separately and together. This allows more refined hypotheses, such as *Is the effect of delivery method the same for all dosages?*, to be tested. This will introduce new models and methods for analyzing data where there are two factors as explanatory variables in a model for a quantitative response variable in what is called the Two-Way ANOVA.

## 3.9 Summary of important R code

The main components of R code used in this chapter follow with components to modify in red, remembering that any R packages mentioned need to be installed and loaded for this code to have a chance of working:

- `MODELNAME <- lm(Y~X, data=DATASETNAME)`
  - Probably the most frequently used command in R.
  - Here it is used to fit the reference-coded One-Way ANOVA model with Y as the response variable and X as the grouping variable, storing the estimated model object in MODELNAME.
- `MODELNAME <- lm(Y~X-1, data=DATASETNAME)`
  - Fits the cell means version of the One-Way ANOVA model.
- `summary(MODELNAME)`
  - Generates model summary information including the estimated model coefficients, SEs, t-tests, and p-values.
- `anova(MODELNAME)`
  - Generates the ANOVA table but **must only be run on the reference-coded version of the model**.
  - Results are incorrect if run on the cell-means model since the reduced model under the null is that the mean of all the observations is 0!
- `pf(FSTATISTIC, df1=NUMDF, df2=DENOMDF, lower.tail=F)`
  - Finds the p-value for an observed  $F$ -statistic with NUMDF and DENOMDF degrees of freedom.
- `par(mfrow=c(2,2)); plot(MODELNAME)`
  - Generates four diagnostic plots including the Residuals vs Fitted and Normal Q-Q plot.
- `plot(allEffects(MODELNAME))`

- Requires the `effects` package be loaded.
- Plots the estimated model component.
- `Tm2 <- glht(MODELNAME, linfct=mcp(X="Tukey")); confint(Tm2); plot(Tm2); cld(Tm2)`
  - Requires the `multcomp` package to be installed and loaded.
  - Can only be run on the reference-coded version of the model.
  - Generates the text output and plot for Tukey's HSD as well as the compact letter display.

## 3.10 Practice problems

For these practice problems, you will work with the cholesterol data set from the `multcomp` package that was used to generate the Tukey's HSD results. To load the data set and learn more about the study, use the following code:

```
require(multcomp)
data(cholesterol)
help(cholesterol)
```

- 3.1. Graphically explore the differences in the changes in Cholesterol levels for the five levels using boxplots and beanplots.
- 3.2. Is the design balanced?
- 3.3. Complete all 6+ steps of the hypothesis test using the parametric  $F$ -test, reporting the ANOVA table and the distribution of the test statistic under the null.
- 3.4. Discuss the scope of inference using the information that the treatment levels were randomly assigned to volunteers in the study.
- 3.5. Generate the permutation distribution and find the p-value. Compare the parametric p-value to the permutation test results.
- 3.6. Perform Tukey's HSD on the data set. Discuss the results – which pairs were detected as different and which were not? Bigger reductions in cholesterol are good, so are there any levels you would recommend or that might provide similar reductions?
- 3.7. Find and interpret the CLD and compare that to your interpretation of results from 3.6.



# Chapter 4

## Two-Way ANOVA

### 4.1 Situation

In this chapter, we extend the One-Way ANOVA to situations with two factors or categorical explanatory variables in a method that is generally called the **Two-Way ANOVA**. This allows researchers to simultaneously study more than one variable that might explain variability in the responses and explore whether the impacts of one variable change depending on the other variable. In some situations, each observation is so expensive that researchers want to use a single study to explore two different sets of research questions in the same round of data collection. For example, a company might want to study factors that affect the number of defective products per day and are interested in the impacts of two different types of training programs and three different levels of production quotas. These methods would allow engineers to compare the training programs, production quotas, and see if the training programs work differently for different production quotas. In a clinical trials context, it is well known that certain factors can change the performance of certain drugs. For example, different dosages of a drug might have different benefits or side-effects on men, versus women or children. **When the impact of one factor changes on the level of another factor**, we say that they *interact*. It is also possible for both factors to be related to differences in the mean responses and not interact. For example, suppose there is a difference in the response means between men and women and a difference among various dosages, but the effect of increasing the dosage is the same for the male and female subjects. This is an example of what is called an **additive** type of model. In general, the world is more complicated than the single factor models we considered in Chapter 3 can account for, especially in observational studies, so these models allow us to start to handle more realistic situations.

Consider the following “experiment” where we want to compare the strength of different brands of paper towels when they are wet. The response variable will be the time to failure in seconds (a continuous response variable) when a weight is placed on the towel held at the four corners. We are interested in studying the differences between brands and the impact of different amounts of water applied to the towels.

- Predictors (Explanatory Variables): **A** : Brand (2 brands of interest, named *B1* and *B2*) and **B** : Number of Drops of water (10, 20, 30 drops).
- Response: *Time* to failure (in seconds) of a towel (*y*) with a weight sitting in the middle of the towel.

### 4.2 Designing a two-way experiment and visualizing results

Ideally, we want to randomly assign the levels of each factor so that we can attribute causality to any detected effects and to reduce the chances of *confounding*. Because there are two factors, we would need to design a random assignment scheme to select the levels of both variables. For example, we could randomly select a brand and then randomly select the number of drops to apply from the levels chosen for each measurement.

Or we could decide on how many observations we want at each combination of the two factors (ideally having them all equal so the design is **balanced**) and then randomize the order of applying the different combinations of levels.

Why might it be important to randomly apply the brand and number of drops in an experiment? There are situations where the order of observations can be related to changes in the responses and we want to be able to eliminate the order of observations from being related to the levels of the factors. For example, suppose that the area where the experiment is being performed becomes wet over time and the later measurements have extra water that gets onto the paper towels and they tend to fail more quickly. If all the observations for the second brand were done later in the study, then the *order of observations* impacts could make the second brand look worse. If the order of observations is randomized, then even if there is some drift in the responses over the order of observations it should still be possible to see the differences in the randomly assigned effects. If the study incorporates repeated measurements on human subjects, randomizing the order of treatments they are exposed to can alleviate impacts of them “learning” through the study, something that we would not have to worry about with paper towels.

In observational studies, we do not have the luxury of random assignment, that is, we cannot randomly assign levels of the treatment variables to our subjects, so we cannot guarantee that the only difference between the groups are the explanatory variables. As discussed before, because we can’t control which level of the variables are assigned to the subjects, we cannot make causal inferences and have to worry about other variables being the real drivers of the results. Although we can never establish causal inference with observational studies, we can generalize our results to a larger population if we have a representative sample from our population of interest.

It is also possible that we might have studies where some of the variables are randomly assigned and others are not randomly assignable. The most common versions of this are what we sometimes call subject “demographics”, such as sex, income, race, etc. We might be performing a study where we can randomly assign treatments to these subjects but might also want to account for differences based on income level, which we can’t assign. In these cases, the scope of inference gets complicated – differences seen on randomized variables can be causally interpreted but you have to be careful to not say that the demographics caused differences. Suppose that a randomly assigned drug dosage is found to show differences in male patients but not in female patients. We could say that the dosage causes differences in males but does not in females. We are not saying that sex caused the differences but that the causal differences were modified by the sex of the subjects.

Even when we do have random assignment of treatments it is important to think about who/what is included in the sample. To get back to the paper towel example, we are probably interested in more than the sheets of the rolls we have to work with so if we could randomly select the studied paper towels from all paper towels made by each brand, our conclusions could be extended to those populations. That probably would not be practical, but trying to make sure that the towels are representative of all made by each brand by checking for defects and maybe picking towels from a few different rolls would be a good start to being able to extend inferences beyond the tested towels.

Once random assignment and random sampling is settled, the final aspect of study design involves deciding on the number of observations that should be made. The short (glib) answer is to take as many as you can afford. With more observations comes higher power to detect differences if they exist, which is a desired attribute of all studies. It is also important to make sure that you obtain multiple observations at each combination of the treatment levels, which are called **replicates**. Having replicate measurements allows estimation of the mean for each combination of the treatment levels as well as estimation and testing for an interaction. And we always prefer having balanced designs because they provide resistance to violation of some assumptions as noted in Chapter 3. A **balanced design** in a Two-Way ANOVA setting involves having the same sample size for every combination of the levels of the treatments.

With two categorical explanatory variables, there are now five possible scenarios for the truth. Different situations are created depending on whether there is an interaction between the two variables, whether both variables are important but do not interact, or whether either of the variables matter at all. Basically, there are five different possible outcomes in a randomized Two-Way ANOVA study, listed in order of increasing

model complexity:

1. Neither A or B has an effect on the responses (nothing causes differences in responses).
2. A has an effect, B does not (only A causes differences in responses).
3. B has an effect, A does not (only B causes differences in responses).
4. Both A and B have effects on response but no interaction (A and B both cause differences in responses but the impacts are additive).
5. Effect of A differs based on the levels of B, the opposite is also true (means for levels of A are different for different levels of B, or, simply, A and B interact).

To illustrate these five potential outcomes, we will consider a fake version of the paper towel example. It ended up being really messy and complicated to actually perform the experiment as we described it so these data were simulated to help us understand the Two-Way ANOVA possibilities in as simple a situation as possible. The first step is to understand what has been observed (number observations at each combination of factors) and look at some summary statistics across all the “groups”. The data set is available from the course website using:

```
pt<-read.csv("http://www.math.montana.edu/courses/s217/documents/pt.csv")
pt$drops<-factor(pt$drops)
```

The data set contains five observations per combination of treatment levels as provided by the `tally` function. To get counts for combinations of the variables, use the general formula of `tally(x1~x2, data=...)` although the order of `x1` and `x2` doesn't matter:

```
require(mosaic)
tally(brand ~ drops, data=pt)
```

```
##      drops
## brand 10 20 30
##   B1  5  5  5
##   B2  5  5  5
```

The sample sizes in each of the six treatment level combinations of `Brand` and `Drops` [( $B_1, 10$ ), ( $B_1, 20$ ), ( $B_1, 30$ ), ( $B_2, 10$ ), ( $B_2, 20$ ), ( $B_2, 30$ )] are  $n_{jk} = 5$  for  $j^{th}$  level of `Brand` ( $j = 1, 2$ ) and  $k^{th}$  level of `Drops` ( $k = 1, 2, 3$ ). The `tally` function gives us a *contingency table* with  $R = 2$  rows ( $B_1, B_2$ ) and  $C = 3$  columns (10, 20, and 30). We'll have more fun with this sort of summary of  $R$  by  $C$  tables in Chapter 5 – here it helps us see the sample size in each combination of factor levels. The `favstats` function also helps us dig into the results for all combinations of factor levels. The notation involves putting both variables after the “~” with a “+” between them. In the output, the first row contains summary information for the 5 observations for `Brand B1` and `Drops` amount 10. It also contains the sample size in the `n` column, although here it rolled into a new set of rows with the standard deviations.

```
favstats(responses ~ brand + drops, data=pt)
```

```
##   brand.drops      min       Q1     median       Q3      max      mean
## 1      B1.10 0.3892621 1.3158737 1.906436 2.050363 2.333138 1.599015
## 2      B2.10 2.3078095 2.8556961 3.001147 3.043846 3.050417 2.851783
## 3      B1.20 0.3838299 0.7737965 1.516424 1.808725 2.105380 1.317631
## 4      B2.20 1.1415868 1.9382142 2.066681 2.838412 3.001200 2.197219
## 5      B1.30 0.2387500 0.9804284 1.226804 1.555707 1.829617 1.166261
## 6      B2.30 0.5470565 1.1205102 1.284117 1.511692 2.106356 1.313946
##
##      sd n missing
## 1 0.7714970 5     0
## 2 0.3140764 5     0
## 3 0.7191978 5     0
## 4 0.7509989 5     0
## 5 0.6103657 5     0
## 6 0.5686485 5     0
```

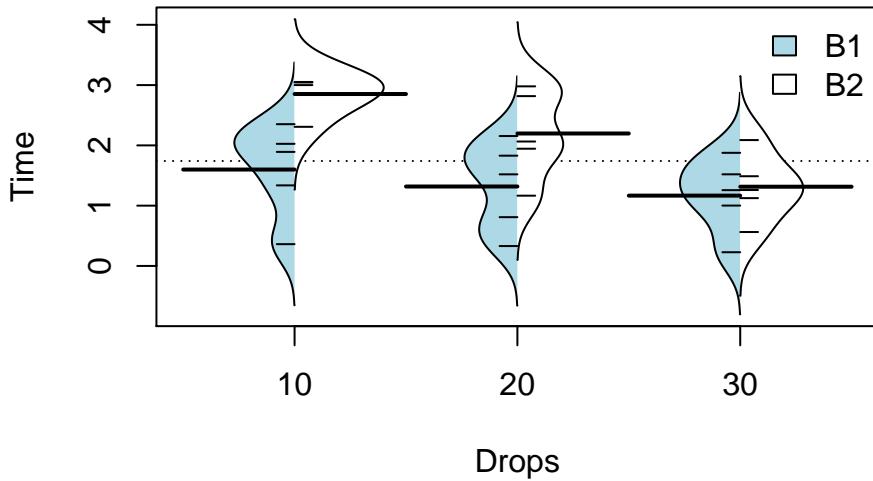


Figure 4.1: Beanplot of paper towel data by Drops (x-axis) and Brand (side of bean, shaded area for Brand B1).

The next step is to visually explore the results across the combinations of the two explanatory variables. The beanplot can be extended to handle these sorts of two-way situations only if one of the two variables is a two-level variable. This is a pretty serious constraint on this display, so we will show you the plot (Figure 4.1) but not focus on the code. The reason beanplots can only handle  $2 \times K$  designs is that the beans are split along a vertical line for the  $K$  levels of the other variable. In Figure 4.1, the Brand B1 density curves are shaded and the B2 curves are not. In reading these plots, look for differences in each level and whether those differences change across the levels of the other variable. Specifically, start with comparing the two brands for different amounts of water. Do the brands seem different? Certainly for 10 drops of water the two look different but not for 30 drops. We can also look for combinations of factors that produce the highest or lower responses in this display. It appears that the time to failure is highest in the low water drop groups but as the water levels increase, the time to failure falls and the differences in the two brands seem to decrease. The fake data seem to have relatively similar amounts of variability and distribution shapes – remembering that there are only 5 observations available for describing the shape of responses for each combination. These data were simulated using a normal distribution and constant variance if that gives you some extra confidence in assessing these model assumptions.

```
require(beanplot)
beanplot(responses ~ brand*drops, data=pt, side="b", col=list("lightblue","white"),
         xlab="Drops", ylab="Time", method="jitter", log="")
legend("topright", bty="n", c("B1","B2"), fill=c("lightblue","white"))
```

The beanplots can't handle situations where both variables have more than two levels – we need a simpler display that just focuses on the means at the combinations of the two explanatory variables. The means for each combination of levels that you can find in the `favstats` output are more usefully used in what is called an *interaction plot*. Interaction plots display the mean responses (y-axis) versus levels of one predictor variable on the x-axis, adding points and lines for each level of the other predictor variable. Because we don't like any of the available functions in R, we wrote our own function, called `intplot` that you can download using:

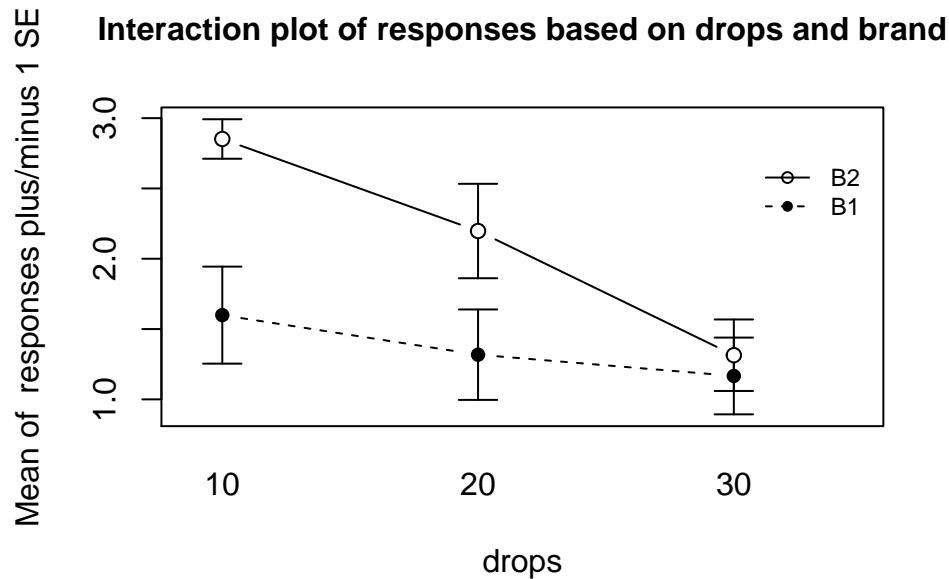


Figure 4.2: Interaction plot of the paper towel data with Drops on the x-axis.

```
source("http://www.math.montana.edu/courses/s217/documents/intplot.R")
```

The function allows a formula interface like  $Y \sim X_1 * X_2$  and provides the means  $\pm 1$  SE (vertical bars) and adds a legend to help make everything clear.

```
intplot(responses ~ brand*drops, data=pt)
```

Interaction plots can always be made two different ways by switching the order of the variables. Figure 4.2 contains **Drops** on the x-axis and Figure 4.3 has **Brand** on the x-axis. Typically putting the variable with more levels on the x-axis will make interpretation easier, but not always. Try both and decide on the one that you like best.

```
intplot(responses ~ drops*brand, data=pt)
```

The formula in this function builds on our previous notation and now we include both predictor variables with an “ $*$ ” between them. Using an asterisk between explanatory variables is one way of telling R to include an interaction between the variables. While the interaction may or may not be present, the interaction plot helps us to explore those potential differences.

There are a variety of aspects of the interaction plots to pay attention to. Initially, the question to answer is whether it appears that there is an interaction between the predictor variables. When there is an interaction, you will see **non-parallel lines** in the interaction plot. You want to look from left to right in the plot and assess whether the lines are close to parallel, relative to the amount of variability in the means. If it seems that there is clear visual evidence of non-parallel lines, then the interaction is likely worth considering (we will typically use a hypothesis test to formally assess this – see discussion below). If the lines look to be close to parallel, then there probably isn’t an interaction between the variables. Without an interaction present, that means that the differences across levels of one variable doesn’t change based on the levels of the other variable and vice-versa. This means that we can consider the **main effects** of each variable on their own<sup>1</sup>. Main effects are much like the results we found in Chapter 3 where we can compare means across levels of a single variable except that there are results for two variables to extract from the model. With the presence of

<sup>1</sup>We will use “main effects” to refer to the two explanatory variables in the additive model even if they are not randomly assigned to contrast with having those variables interacting in the model. It is the one place where we use “effects” without worrying about random assignment.

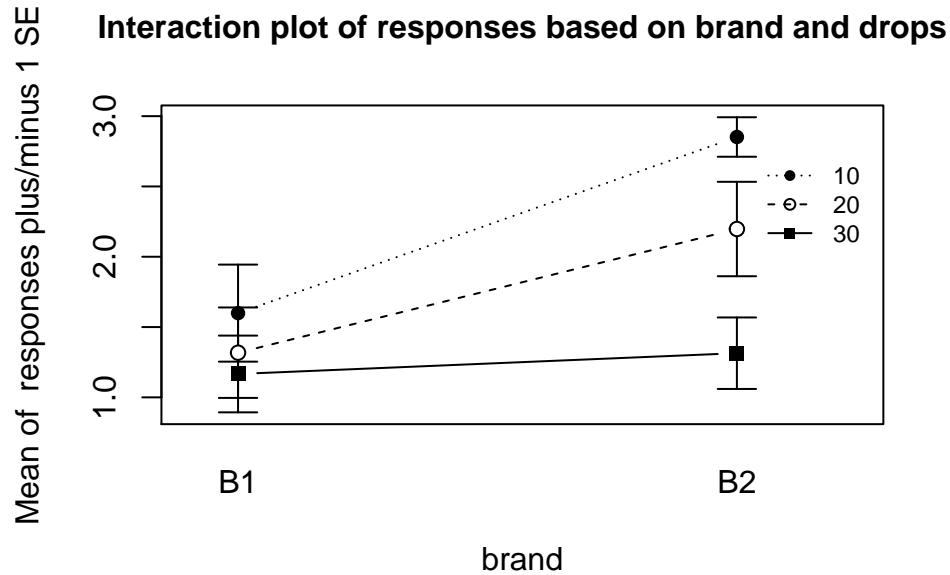


Figure 4.3: Interaction plot of paper towel data with **Brand** on the x-axis.

an interaction, it is complicated to summarize how each variable is affecting the response variable because their impacts change depending on the level of the other factor. And plots like the interaction plot provide us much useful information.

If the lines are not parallel, then focus in on comparing the levels of one variable as the other variable changes. Remember that the definition of an interaction is that the differences among levels of one variable depends on the level of the other variable being considered. “Visually” this means comparing the size of the differences in the lines from left to right. In Figures 4.2 and 4.3, the effect of amount of water changes based on the brand being considered. In Figure 4.3, the three lines represent the three water levels. The difference between the brands (left to right, *B1* to *B2*) is different depending on how much water was present. It appears that **Brand B2** lasted longer at the lower water levels but that the difference between the two brands dropped as the water levels increased. The same story appears in Figure 4.2. As the water levels increase (left to right, 10 to 20 to 30 drops), the differences between the two brands decrease. Of the two versions, Figure 4.2 is probably easier to read here. The interaction plots also are useful for identifying the best and worst mean responses for combinations of the treatment levels. For example, 10 Drops and **Brand B2** lasts longest, on average, and 30 Drops with **Brand B1** fails fastest, on average. In this situation, the lines do not appear to be parallel suggesting that further exploration of the interaction appears to be warranted.

Before we get to the hypothesis tests to formally make this assessment (you knew some sort of p-value was coming, right?), we can visualize the 5 different scenarios that could characterize the sorts of results you could observe in a Two-Way ANOVA situation. Figure 4.4 shows 4 of the 5 scenarios. In panel (a), when there are no differences from either variable (Scenario 1), it provides relatively parallel lines and basically no differences either across **Drops** levels (x-axis) or **Brand** (lines). This would result in no evidence related to a difference in brands, water levels, or any interaction between them.

Scenario 2 (Figure 4.4 panel (b)) incorporates differences based on factor A (here that is **Brand**) but no real difference based on the **Drops** or any interaction. This results in a clear shift between the little to no changes in the level of those lines across water levels. These lines are relatively parallel. We can see that **Brand B2** is better than **Brand B1** but that is all we can show with these sorts of results.

Scenario 3 (Figure 4.4 panel (c)) flips the important variable to B (**Drops**) and shows decreasing average

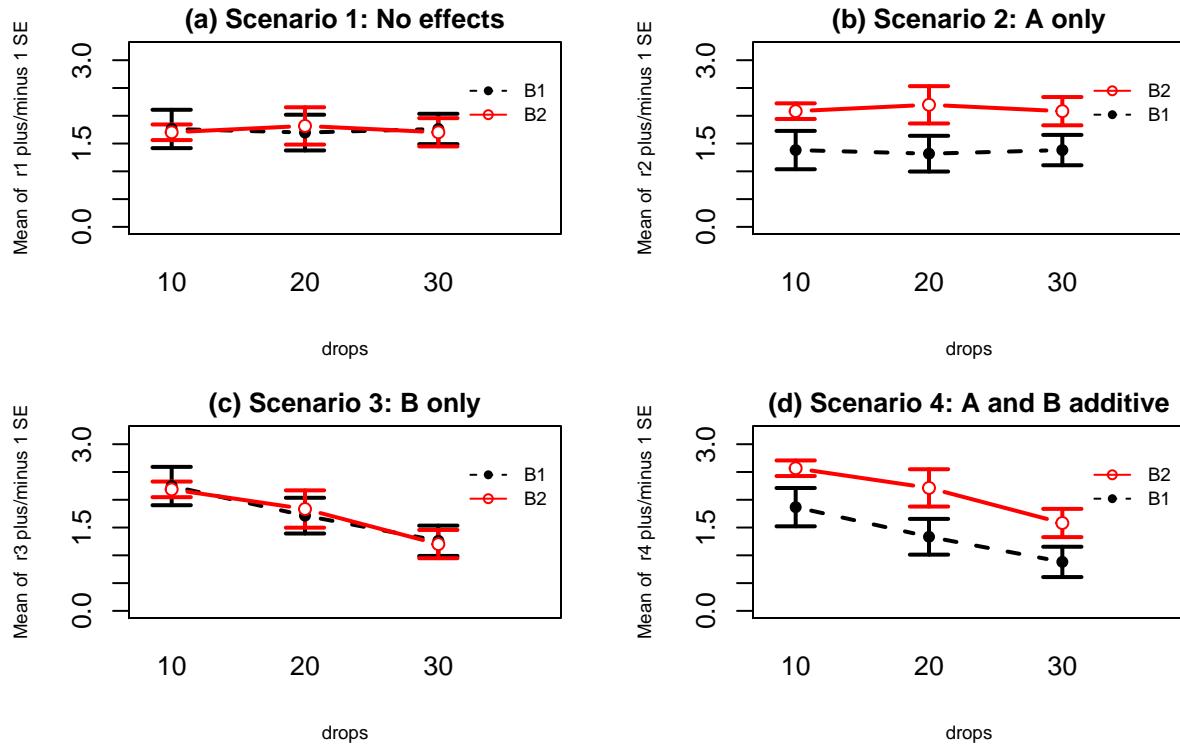


Figure 4.4: Interaction plots of four possible scenarios in the paper towel study.

times as the water levels increase. Again, the interaction panels show near parallel-ness in the lines and really just show differences among the levels of the water. In both Scenarios 2 and 3, we could use a single variable and drop the other from the model, getting back to a One-Way ANOVA model, without losing any important information.

Scenario 4 (Figure 4.4 panel (d)) incorporates effects of A and B, but they are ***additive***. That means that the effect of one variable is the same across the levels of the other variable. In this experiment, that would mean that **Drops** has the same impact on performance regardless of brand and that the brands differ but each type of difference is the same regardless of levels of the other variable. The interaction plot lines are more or less parallel but now the brands are clearly different from each other. The plot shows the decrease in performance based on increasing water levels and that **Brand B2** is better than **Brand B1**. Additive effects show the same difference in lines from left to right in the interaction plots.

Finally, Scenario 5 (Figure 4.5) involves an interaction between the two variables (**Drops** and **Brand**). There are many ways that interactions can present but the main thing is to look for clearly non-parallel lines. As noted in the previous discussion, the **Drops** effect appears to change depending on which level of **Brand** is being considered. Note that the plot here described as Scenario 5 is the same as the initial plot of the results in Figure 4.2.

The typical modeling protocol is to start with assuming that Scenario 5 is a possible description of the results, related to fitting what is called the ***interaction model***, and then attempt to simplify the model (to the ***additive model***) if warranted. We need a hypothesis test to help decide if the interaction is “real” – if there is sufficient evidence to prove that there is an interaction. We need a test because the lines will never be exactly parallel and, just like in the One-Way ANOVA situation, the amount of variation around the lines impacts the ability of the model to detect differences, in this case of an interaction.

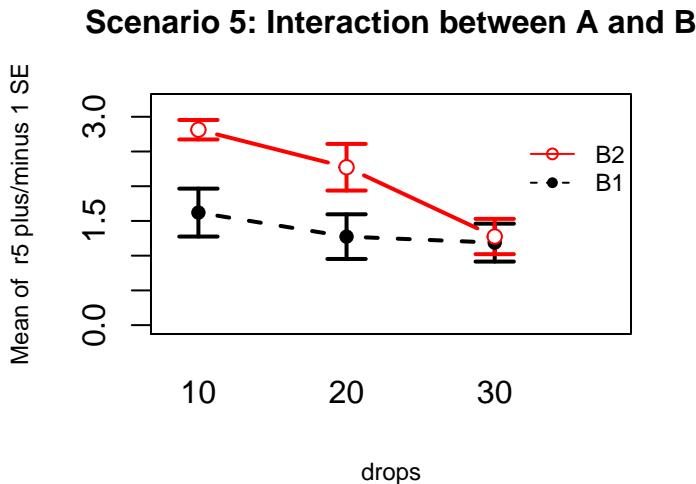


Figure 4.5: Interaction plot of Scenario 5 where it appears that an interaction is present.

### 4.3 Two-Way ANOVA models and hypothesis tests

To assess interactions with two variables, we need to fully describe models for the additive and interaction scenarios and then develop a method for assessing evidence of the need for different aspects of the models. First, we need to define the notation for these models:

- $y_{ijk}$  is the  $i^{th}$  response from the group for level  $j$  of factor A and level  $k$  of factor B
  - $j = 1, \dots, J$        $J$  is the number of levels of A
  - $k = 1, \dots, K$        $K$  is the number of levels of B
  - $i = 1, \dots, n_{jk}$        $n_{jk}$  is the sample size for level  $j$  of factor A and level  $k$  of factor B
  - $N = \sum n_{jk}$  is the total sample size (sum of the number of observations across all  $JK$  groups)

We need to extend our previous discussion of reference-coded models to develop a Two-Way ANOVA model. We start with the ***Two-Way ANOVA interaction model***:

$$y_{ijk} = \alpha + \tau_j + \gamma_k + \omega_{jk} + \varepsilon_{ijk},$$

where  $\alpha$  is the baseline group mean (for level 1 of A **and** level 1 of B),  $\tau_j$  is the deviation for the **main effect** of A from the baseline for levels  $2, \dots, J$ ,  $\gamma_k$  (gamma  $k$ ) is the deviation for the main effect of B from the baseline for levels  $2, \dots, K$ , and  $\omega_{jk}$  (omega  $jk$ ) is the adjustment for the **interaction effect** for level  $j$  of factor A and level  $k$  of factor B for  $j = 1, \dots, J$  and  $k = 1, \dots, K$ . In this model,  $\tau_1$ ,  $\gamma_1$ , and  $\omega_{11}$  are all fixed at 0. As in Chapter 3, R will choose the baseline categories alphabetically but now it is choosing a baseline for both variables and so our detective work will be doubled to sort this out.

If the interaction term is not important, based on the interaction test presented below, the  $\omega_{jk}$ 's can be dropped from the model and we get a model that corresponds to Scenario 4 above. Scenario 4 is where there are two main effects but no interaction between them. The **additive Two-Way model** is

$$y_{ijk} = \alpha + \tau_j + \gamma_k + \varepsilon_{ijk},$$

where each component is defined as in the interaction model. The difference between the interaction and additive models is setting all the  $\omega_{jk}$ 's to 0 that are present in the interaction model. When we set parameters to 0 in models it removes them from the model. Setting parameters to 0 is how we will develop our hypotheses to test for an interaction, by testing whether there is evidence enough to reject that all  $\omega_{jk}$ 's = 0.

The interaction test hypotheses are

- $H_0$ : No interaction between A and B in population  $\Leftrightarrow$  All  $\omega_{jk}$ 's = 0.
- $H_A$ : Interaction between A and B in population  $\Leftrightarrow$  At least one  $\omega_{jk} \neq 0$

To perform this test, a new ANOVA  $F$ -test is required (presented below) but there are also hypotheses relating to the main effects of A ( $\tau_j$ 's) and B ( $\gamma_k$ 's). If evidence is found to reject the null hypothesis that no interaction is present, then it is dangerous to ignore it and test for the main effects because important main effects can be masked by interactions (examples later). It is important to note that, by definition, **both variables matter if an interaction is found to be important** so the main effect tests may not be very interesting. If the interaction is found to be important based on the test and retained in the model, you should focus on the interaction model (also called the ***full model***) in order to understand and describe the form of the interaction among the variables.

If the interaction test does not return a small p-value, then we have no evidence to suggest that it is needed and it can be dropped from the model. In this situation, we would re-fit the model and focus on the results provided by the additive model – performing tests for the two additive main effects. For the first, but not last time, we encounter a model with more than one variable and test of potential interest. In models with multiple variables at similar levels (here both are main effects), we are interested in the results for each variable given that the other variable is in the model. In many situations, including more than one variable in a model changes the results for the other variable even if those variables do not interact. The reason for this is more clear in Chapter 8 and really only matters here if we have unbalanced designs, but we need to start adding a short modifier to our discussions of main effects – they are the results *conditional on* or *adjusting for* or, simply, *given*, the other variable(s) in the model. Specifically, the hypotheses for the two main effects are:

- Main effect test for A:
  - $H_0$ : No differences in means across levels of A in population, given B in the model  $\Leftrightarrow$  All  $\tau_j$ 's = 0 in additive model.
  - $H_A$ : Some difference in means across levels A in population, given B in the model  $\Leftrightarrow$  At least one  $\tau_j \neq 0$ , in additive model.
- Main effect test for B:
  - $H_0$ : No differences in means across levels of B in population, given A in the model  $\Leftrightarrow$  All  $\gamma_k$ 's = 0 in additive model.
  - $H_A$ : Some difference in means across levels B in population, given A in the model  $\Leftrightarrow$  At least one  $\gamma_k \neq 0$ , in additive model.

In order to test these effects (interaction in the interaction model and main effects in the additive model),  $F$ -tests are developed using Sums of Squares, Mean Squares, and degrees of freedom similar to those in Chapter 3. We won't worry about the details of the sums of squares formulas but you should remember the sums of squares decomposition, which still applies<sup>2</sup>. Table 4.1 summarizes the ANOVA results you will obtain for the interaction model and Table 4.2 provides the similar general results for the additive model. As we saw in Chapter 3, the degrees of freedom are the amount of information that is free to vary at a particular level and that rule generally holds here. For example, for factor A with  $J$  levels, there are  $J - 1$  parameters that are free since the baseline is fixed. The residual degrees of freedom for both models are not as easily explained but have simple formula. Note that the sum of the degrees of freedom from the main effects, (interaction if present), and error need to equal  $N - 1$ , just like in the One-Way ANOVA table.

---

<sup>2</sup>In the standard ANOVA table,  $SS_A + SS_B + SS_{AB} + SS_E = SS_{Total}$ . However, to get the tests we really desire when our designs are not balanced, a slight modification of the SS is used, using what are called Type II sums of squares and this result doesn't hold in the output you will see for additive models. This is discussed further below.

Table 4.1: Interaction Model ANOVA Table.

Source	DF	SS	MS	F-statistics
A	$J - 1$	$SS_A$	$MS_A = SS_A/df_A$	$MS_A/MS_E$
B	$K - 1$	$SS_B$	$MS_B = SS_B/df_B$	$MS_B/MS_E$
A:B (interaction)	$(J - 1)(K - 1)$	$SS_{AB}$	$MS_{AB} = SS_{AB}/df_{AB}$	$MS_{AB}/MS_E$
Error	$N - J - K + 1$	$SS_E$	$MS_E = SS_E/df_E$	
<b>Total</b>	<b>N - 1</b>	<b>SS<sub>Total</sub></b>		

Table 4.2: Additive Model ANOVA Table.

Source	DF	SS	MS	F-statistics
A	$J - 1$	$SS_A$	$MS_A = SS_A/df_A$	$MS_A/MS_E$
B	$K - 1$	$SS_B$	$MS_B = SS_B/df_B$	$MS_B/MS_E$
Error	$N - J - K + 1$	$SS_E$	$MS_E = SS_E/df_E$	
<b>Total</b>	<b>N - 1</b>	<b>SS<sub>Total</sub></b>		

The mean squares are formed by taking the sums of squares (we'll let R find those for us) and dividing by the *df* in the row. The *F*-ratios are found by taking the mean squares from the row and dividing by the mean squared error ( $MS_E$ ). They follow *F*-distributions with numerator degrees of freedom from the row and denominator degrees of freedom from the Error row (in R output this the `Residuals` row). It is possible to develop permutation tests for these methods but some technical issues arise in doing permutation tests for interaction model components so we will not use them here. This means we will have to place even more emphasis on meeting the assumptions since we only have the parametric method available.

With some basic expectations about the ANOVA tables and *F*-statistic construction in mind, we can get to actually estimating the models and exploring the results. The first example involves the fake paper towel data displayed in Figure 4.1 and 4.2. It appeared that Scenario 5 was the correct story since the lines were not parallel, but we need to know whether there is evidence to suggest that the interaction is “real” and we get that through the interaction hypothesis test. To fit the interaction model using `lm`, the general formulation is `lm(y ~ x1*x2, data=...)`. The order of the variables doesn't matter and the most important part of the model, to start with, relates to the interaction of the variables.

The ANOVA table output shows the results for the interaction model obtained by running the `anova` function on the model called `m1`. Specifically, the test that  $H_0 : \text{All } \omega_{jk}'s = 0$  has a test statistic of  $F(2, 24) = 1.92$  (in bold in the output from the row with `brands:drops`) and a p-value of 0.17. So there is insufficient evidence to reject the null hypothesis of no interaction, with a 17% chance we would observe a difference in the  $\omega_{jk}$ 's like we did or more extreme if the  $\omega_{jk}$ 's really were all 0. For the interaction model components, R presents them with a colon, `:`, between the variable names.

```
m1<-lm(responses ~ brand*drops, data=pt)
anova(m1)
```

```
## Analysis of Variance Table
##
## Response: responses
##              Df Sum Sq Mean Sq F value    Pr(>F)
## brand          1 4.3322  4.3322 10.5192 0.003458
## drops          2 4.8581  2.4290  5.8981 0.008251
## brand:drops   2 1.5801  0.7901  1.9184 0.168695
## Residuals     24 9.8840  0.4118
```

It is useful to display the estimates from this model and we can utilize `plot(allEffects(modelname))` to visualize the results for the terms in our models. If we turn on the options for `grid=T`, `multiline=T`, and

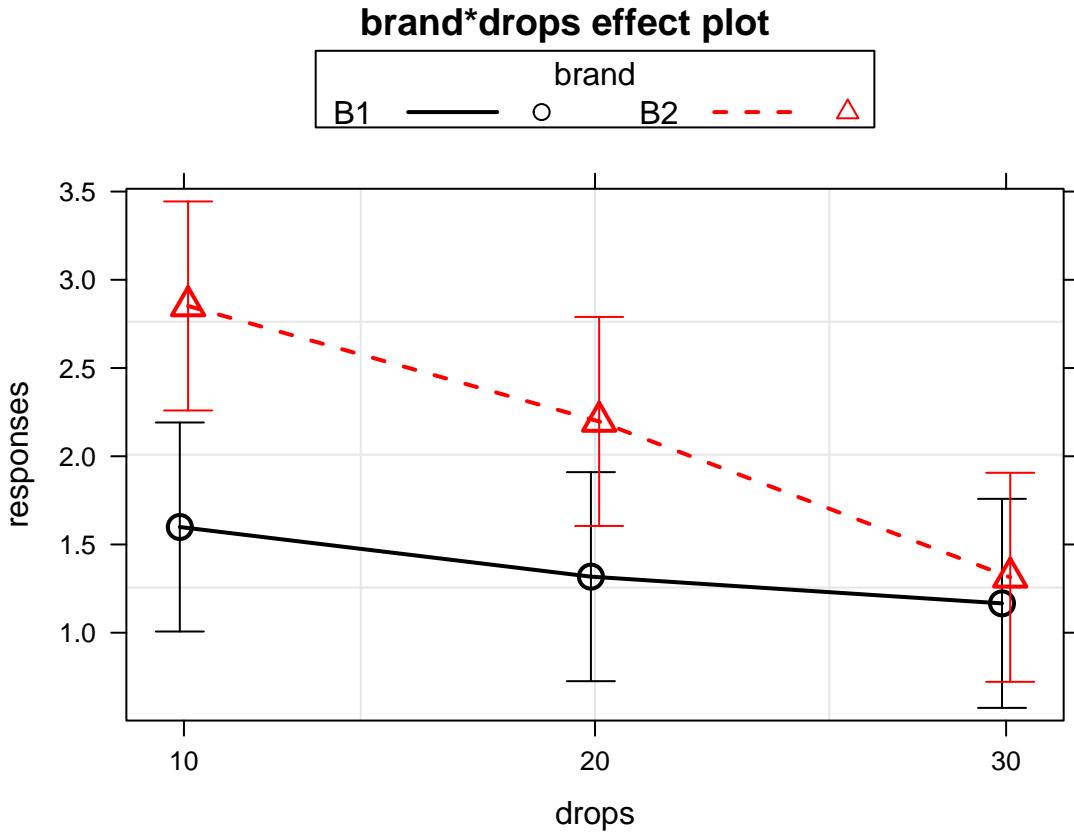


Figure 4.6: Plot of estimated results of interaction model.

ci. style="bars" we will get a more useful version of the basic “effect plot” for Two-Way ANOVA models with interaction. The results of the estimated interaction model are displayed in Figure 4.6, which looks very similar to our previous interaction plot. The only difference is that this comes from model that assumes equal variance and these plots show 95% confidence intervals for the means instead of the 1 standard error used above.

In the absence of evidence to include the interaction, the model should be simplified to the additive model and the interpretation focused on each main effect, conditional on having the other variable in the model. To fit an additive model and not include an interaction, the model formula involves a “+” instead of a “\*” between the explanatory variables.

```
m2<-lm(responses ~ brand + drops, data=pt)
anova(m2)
```

```
## Analysis of Variance Table
##
## Response: responses
##             Df  Sum Sq Mean Sq F value    Pr(>F)
## brand        1  4.3322  4.3322  9.8251 0.004236
## drops        2  4.8581  2.4290  5.5089 0.010123
## Residuals   26 11.4641  0.4409
```

The p-values for the main effects of `brand` and `drops` change slightly from the results in the interaction model due to changes in the  $MS_E$  from 0.4118 to 0.4409 (more variability is left over in the simpler model) and the  $DF_{\text{error}}$  that increases from 24 to 26. In both models, the  $SS_{\text{Total}}$  is the same (20.6544). In the interaction model,

$$\begin{aligned} \text{SS}_{\text{Total}} &= \text{SS}_{\text{brand}} + \text{SS}_{\text{drops}} + \text{SS}_{\text{brand:drops}} + \text{SS}_{\text{E}} \\ &= 4.3322 + 4.8581 + 1.5801 + 9.8840 \\ &= 20.6544 \end{aligned}$$

In the additive model, the variability that was attributed to the interaction term in the interaction model ( $\text{SS}_{\text{brand:drops}} = 1.5801$ ) is pushed into the  $\text{SS}_{\text{E}}$ , which increases from 9.884 to 11.4641. The sums of squares decomposition in the additive model is

$$\begin{aligned} \text{SS}_{\text{Total}} &= \text{SS}_{\text{brand}} + \text{SS}_{\text{drops}} + \text{SS}_{\text{E}} \\ &= 4.3322 + 4.8581 + 11.4641 \\ &= 20.6544 \end{aligned}$$

This shows that the sums of squares decomposition applies in these more complicated models as it did in the One-Way ANOVA. It also shows that if the interaction is removed from the model, that variability is lumped in with the other unexplained variability that goes in the  $\text{SS}_{\text{E}}$  in any model.

The fact that the sums of squares decomposition can be applied here is useful, except that there is a small issue with the main effect tests in the ANOVA table results that follow this decomposition when the design is not balanced. It ends up that the tests in a typical ANOVA table are only conditional on the tests higher up in the table. For example, in the additive model ANOVA table, the `Brand` test is not conditional on the `Drops` effect, but the `Drops` effect is conditional on the `Brand` effect. To fix this issue, we have to use another type of sums of squares, called **Type II sums of squares**. They will no longer always follow the rules of the sums of squares decomposition but they will test the desired hypotheses. Specifically, they provide each test conditional on any other terms at the same level of the model and match the hypotheses written out earlier in this section. To get the “correct” ANOVA results, the `car` (Fox and Weisberg (2016), Fox and Weisberg (2011)) package is required. We use the `Anova` function on our linear models from here forward to get the “right” tests in our ANOVA tables. Note how the case-sensitive nature of R code shows up in the use of the capital-A `Anova` function instead of the `anova` function used previously. In this case, because the design was balanced, the results are the same using either function. Observational studies rarely generate balanced designs (some designed studies can result in unbalanced designs) so we will generally just use the Type II version of the sums of squares. The `Anova` results using the Type II sums of squares are slightly more conservative than the results from `anova`, which are called Type I sums of squares. The sums of squares decomposition no longer can be applied, but it is a small sacrifice to get each test after adjusting for all other variables<sup>3</sup>.

```
require(car)
Anova(m2)
```

```
## Anova Table (Type II tests)
##
## Response: responses
##              Sum Sq Df F value    Pr(>F)
## brand        4.3322  1  9.8251 0.004236
## drops        4.8581  2  5.5089 0.010123
## Residuals  11.4641 26
```

The new output switches the columns around and doesn’t show you the mean squares, but gives the most critical parts of the output. Here, there is no change in results because it is balanced design with equal counts of responses in each combination of the two explanatory variables.

The additive model, when appropriate, provides simpler interpretations for each explanatory variable compared to models with interactions because the effect of one variable is the same regardless of the levels of the other variable and vice versa. There are two tools to aid in understanding the impacts of the two variables in the additive model. First, the model summary provides estimated coefficients with interpretations like those seen in Chapter 3 (deviation of group  $j$  or  $k$  from the baseline group’s mean), except with the additional wording

---

<sup>3</sup>Actually, the tests are only conditional on other main effects if Type II Sums of Squares are used for an interaction model.

of “controlling for” the other variable added to any of the discussion. Second, the term-plots now show each main effect and how the groups differ with one panel for each of the two explanatory variables in the model. These term-plots are created by holding the other variable constant at one of its levels.

```
summary(m2)
```

```
## 
## Call:
## lm(formula = responses ~ brand + drops, data = pt)
## 
## Residuals:
##      Min       1Q   Median       3Q      Max 
## -1.4561 -0.4587  0.1297  0.4434  0.9695 
## 
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) 1.8454    0.2425   7.611 4.45e-08 ***
## brandB2     0.7600    0.2425   3.134  0.00424 **  
## drops20    -0.4680    0.2970  -1.576  0.12715    
## drops30    -0.9853    0.2970  -3.318  0.00269 **  
## 
## Residual standard error: 0.664 on 26 degrees of freedom
## Multiple R-squared:  0.445, Adjusted R-squared:  0.3809 
## F-statistic: 6.948 on 3 and 26 DF,  p-value: 0.001381
```

In the model summary, the baseline combination estimated in the (Intercept) row is for **Brand B1** and **Drops 10** and estimates the mean failure time as 1.85 seconds for this combination. As before, the group labels that do not show up are the baseline but there are two variables’ baselines to identify. Now the “simple” aspects of the additive model show up. The interpretation of the **Brands B2** coefficient is as a deviation from the baseline but it applies regardless of the level of **Drops**. Any difference between **B1** and **B2** involves a shift up of 0.76 seconds in the estimated mean failure time. Similarly, going from 10 (baseline) to 20 drops results in a drop in the estimated failure mean of 0.47 seconds and going from 10 to 30 drops results in a drop of almost 1 second in the average time to failure, both estimated changes are the same regardless of the brand of paper towel being considered. Sometimes, especially in observational studies, we use the terminology “controlled for” to remind the reader that the other variable was present in the model<sup>4</sup> and also explained some of the variability in the responses. The term-plots for the additive model (Figure 4.7) help us visualize the impacts of changes brand and changing water levels, holding the other variable constant. The differences in heights in each panel correspond to the coefficients just discussed.

```
require(effects)
plot(allEffects(m2))
```

---

<sup>4</sup>In Multiple Linear Regression models in Chapter 8, the reasons for this wording will (hopefully) become clearer.

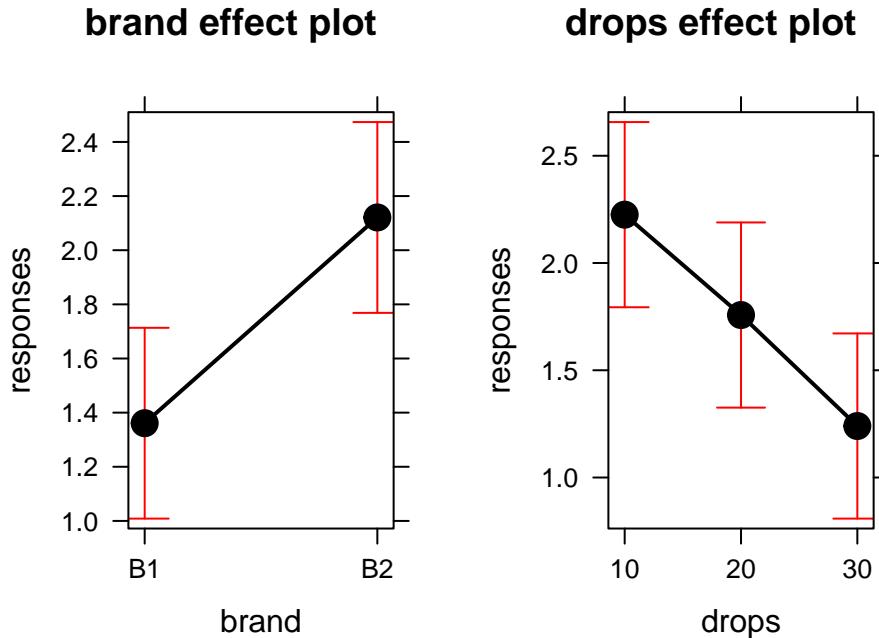


Figure 4.7: Term-plots of additive model for paper towel data. Left panel displays results for two brands and right panel for number of drops of water, each after controlling for the other.

## 4.4 Guinea pig tooth growth analysis with Two-Way ANOVA

The effects of dosage and delivery method of ascorbic acid on Guinea Pig odontoblast growth was analyzed as a One-Way ANOVA in Section 3.4 by assessing evidence of any difference in the means of any combinations of dosage method (Vit C capsule vs Orange Juice) and three dosage amounts (0.5, 1, and 2 mg/day). Now we will consider the dosage and delivery methods as two separate variables and explore their potential interaction. A beanplot and interaction plot are provided in Figure 4.8.

```
data(ToothGrowth)
par(mfrow=c(1,2))
beanplot(len ~ supp*dose, data=ToothGrowth, side="b", ylim=c(-5,40),
         main="Beanplot", col=list("white","orange"), xlab="Dosage",
         ylab="Tooth Growth")
legend("bottomright", bty="n", c("VC","OJ"), fill=c("white","orange"))
intplot(len ~ supp*dose, data=ToothGrowth, col=c(1,2),
        main="Interaction Plot", ylim=c(-5,40))
```

It appears that the effect of method changes based on the dosage as the interaction plot seems to show some evidence of non-parallel lines. Actually, it appears that the effect of delivery method is parallel for doses 0.5 and 1.0 mg/day but that the effect of delivery method changes for 2 mg/day.

We can use the ANOVA  $F$ -test for an interaction to assess whether the interaction is “real” relative to the variability in the responses. That is, is it larger than we would expect due to natural variation in the data? If yes, then it is a real effect and we should account for it. The following results fit the interaction model and provide an ANOVA table.

```
TG1 <- lm(len~supp*dose,data=ToothGrowth)
Anova(TG1)
```

```
## Anova Table (Type II tests)
##
```

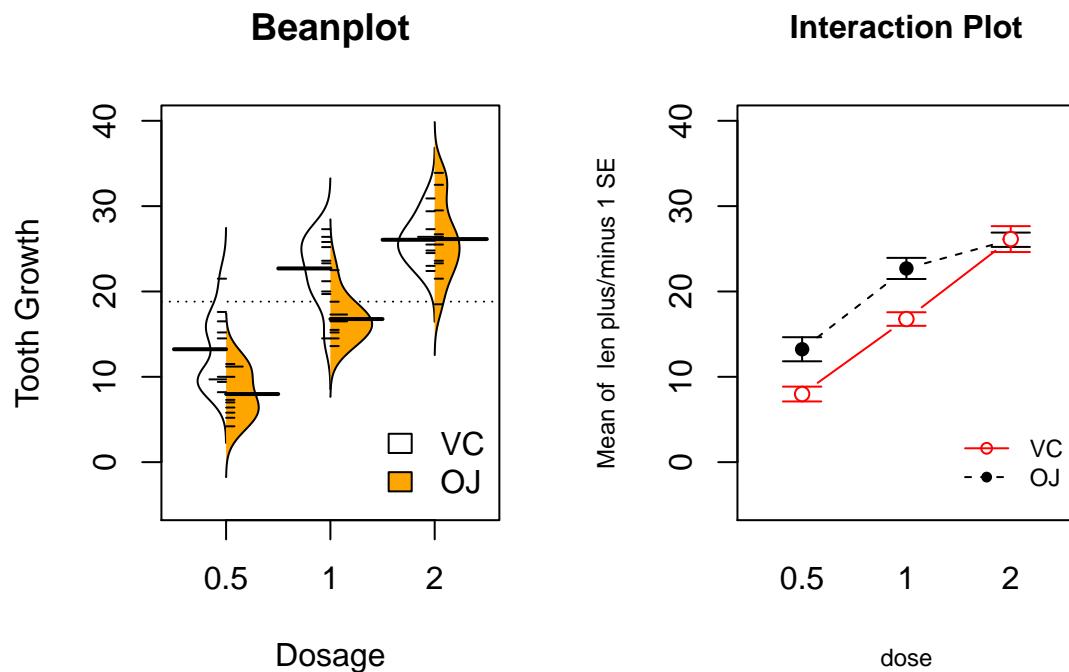


Figure 4.8: Beanplot and interaction plot of the tooth growth data set.

```
## Response: len
##           Sum Sq Df  F value    Pr(>F)
## supp      205.35  1 12.3170 0.0008936
## dose     2224.30  1 133.4151 < 2.2e-16
## supp:dose 88.92  1   5.3335 0.0246314
## Residuals 933.63 56
```

The R output is reporting an interaction test result of  $F(1, 56) = 5.3$  with a p-value of 0.025. But this should raise a red flag since the numerator degrees of freedom are not what we should expect of  $(K - 1) * (J - 1) = (2 - 1) * (3 - 1) = 2$ . This brings up an issue in R when working with categorical variables. If the levels of a categorical variable are entered numerically, R will treat them as quantitative variables and not split out the different levels of the categorical variable. To make sure that R treats categorical variables the correct way, we should use the **factor** function on any variables that are categorical but are coded numerically in the data set. The following code creates a new variable called **dosef** using the function that will help us obtain correct results from the linear model. The re-run of the ANOVA table provides the correct analysis and the expected **df** for the two rows of output involving **dosef**:

```
ToothGrowth$dosef<-factor(ToothGrowth$dose)
TG2 <- lm(len ~ supp*dosef, data=ToothGrowth)
Anova(TG2)
```

```
## Anova Table (Type II tests)
##
## Response: len
##           Sum Sq Df  F value    Pr(>F)
## supp      205.35  1 15.572 0.0002312
## dosef     2426.43  2 92.000 < 2.2e-16
## supp:dosef 108.32  2   4.107 0.0218603
## Residuals 712.11 54
```

The ANOVA  $F$ -test for an interaction between supplement type and dosage level is  $F(2, 54) = 4.107$  with a p-value of 0.022. So there appears to be enough evidence to reject the null hypothesis of no interaction between *Dosage* and *Delivery method*, supporting a changing effect on tooth growth of dosage based on the delivery method in the Guinea Pigs that were assigned.

Any similarities between this correct result and the previous WRONG result are coincidence. I (Greenwood) once attended a Master's defense where the results from a similar model were not as expected (small p-values in places they didn't expect and large p-values in places where they thought differences existed). During the presentation, the student showed some ANOVA tables and the four level categorical variable had 1 numerator  $df$  in the ANOVA table. The student passed with major revisions but had to re-run all the results and re-write all of the conclusions... So be careful to check the ANOVA results ( $df$  and for the right number of expected model coefficients) to make sure they match your expectations. This is one reason why you will be learning to fill in ANOVA tables based on information about the study so that you can be prepared to detect when your code has let you down<sup>5</sup>. It is also a great reason to explore term-plots and coefficient interpretations as that can also help diagnose errors in model construction.

Getting back to the previous results, we now have enough background information to more formally write up a focused interpretation of these results. The 6+ hypothesis testing steps in this situation would be focused on first identifying that the best analysis here is as a Two-Way ANOVA situation (these data were analyzed in Chapter 3 as a One-Way ANOVA but this version is better because it can explore whether there is an interaction between delivery method and dosage). We will use a 5% significance level and start with assessing the evidence for an interaction. If the interaction had not been dropped, we would have reported the test for the interaction, re-fit the additive model and used it to explore the main effect tests and estimates for *Dose* and *Delivery method*.

### 1. Hypotheses:

- $H_0$ : No interaction between *Delivery method* and *Dose* on odontoblast growth in population of guinea pigs  
 $\Leftrightarrow$  All  $\omega_{jk}$ 's = 0.
- $H_A$ : Interaction between *Delivery method* and *Dose* on odontoblast growth in population of guinea pigs  
 $\Leftrightarrow$  At least one  $\omega_{jk} \neq 0$ .

### 2. Validity conditions:

- Independence:
  - This assumption is presumed to be met because we don't know of a reason why the independence of the measurements of tooth growth of the guinea pigs as studied might be violated.
- Constant variance:
  - To assess this assumption, we can use the diagnostic plots in Figure 4.9.
  - In the Residuals vs Fitted and the Scale-Location plots, the differences in variability among the groups (see the different x-axis positions for each group's fitted values) is minor, so there is not strong evidence of a problem with the equal variance assumption.

```
par(mfrow=c(2,2))
plot(TG2, pch=16)
```

- Normality of residuals:

- The QQ-Plot in Figure 4.9 does not suggest a problem with this assumption.

---

<sup>5</sup>Just so you don't think that perfect R code should occur on the first try, we have all made similarly serious coding mistakes even after accumulating more than decade of experience with R. It is finding those mistakes that matters.

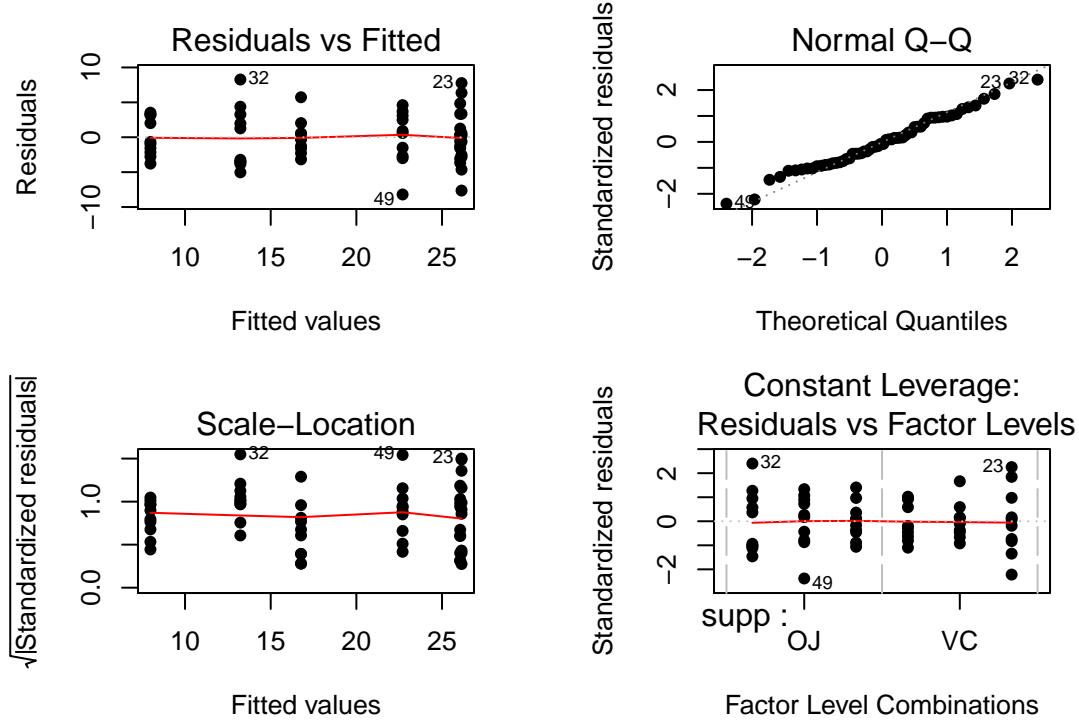


Figure 4.9: Diagnostic plots for the interaction model for Tooth Growth.

## 3. Calculate the test statistic for the Interaction test.

```
TG2 <- lm(len ~ supp*dosef, data=ToothGrowth)
Anova(TG2)
```

```
## Anova Table (Type II tests)
##
## Response: len
##             Sum Sq Df F value    Pr(>F)
## supp       205.35  1 15.572 0.0002312
## dosef      2426.43  2 92.000 < 2.2e-16
## supp:dosef 108.32  2   4.107 0.0218603
## Residuals  712.11 54
```

- The test statistic is  $F(2, 54) = 4.107$ .

## 4. Find the p-value:

- The ANOVA  $F$ -test p-value of 0.0219 for the interaction.
- To find this p-value directly in R, we can use the `pf` function.

```
pf(4.107, df1=2, df2=54, lower.tail=F)
```

```
## [1] 0.0218601
```

## 5. Make a decision:

- Reject  $H_0$  since the p-value (0.0219) is less than 0.05. With a p-value of 0.0219, there is about a 2.19% chance we would observe interaction like we did (or more extreme) if none were truly present. This provides strong evidence against the null hypothesis of no interaction between delivery method and dosage on odontoblast growth so we reject the null hypothesis of no interaction.

### 6. Write a conclusion:

- Therefore, the effects of dosage level (0.5, 1, or 2 mg/day) on population average tooth odontoblast growth rates of Guinea pigs are changed by the delivery (OJ, Vitamin C) method (and vice versa) and we should keep the interaction in the model. With the random assignment of levels but not random selection of subjects here, we could also write this as: Different dosage levels cause different changes in the odontoblast growth based on the delivery method for these guinea pigs.

In a Two-Way ANOVA, we need to go a little further to get to the final interpretations since the models are more complicated. When there is an interaction present, we should focus on the interaction plot or term-plot of the interaction model for an interpretation of the form and pattern of the interaction. If the interaction were unimportant, then the hypotheses and results should focus on the additive model results, especially the estimated model coefficients. To see why we don't spend much time with the estimated model coefficients in an interaction model, the model summary for this model is provided:

```
summary(TG2)
```

```
## 
## Call:
## lm(formula = len ~ supp * dosef, data = ToothGrowth)
## 
## Residuals:
##   Min    1Q Median    3Q   Max 
## -8.20 -2.72 -0.27  2.65  8.27 
## 
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) 13.230    1.148   11.521 3.60e-16  
## suppVC     -5.250    1.624   -3.233  0.00209   
## dosef1      9.470    1.624    5.831 3.18e-07  
## dosef2     12.830    1.624    7.900 1.43e-10  
## suppVC:dosef1 -0.680    2.297   -0.296  0.76831  
## suppVC:dosef2  5.330    2.297    2.321  0.02411  
## 
## Residual standard error: 3.631 on 54 degrees of freedom
## Multiple R-squared:  0.7937, Adjusted R-squared:  0.7746 
## F-statistic: 41.56 on 5 and 54 DF,  p-value: < 2.2e-16
```

There are two  $\omega_{jk}$ 's in the results, related to modifying the estimates for doses of 1 (-0.68) and 2 (5.33) for the Vitamin C group. If you want to re-construct the fitted values from the model that are displayed in the Figure 4.10, you have to look for any coefficients that are "turned on" for a combination of levels of interest. For example, for the OJ group (solid line in Figure 4.10), the dosage of 0.5 mg/day has an estimate of an average growth of approximately 13 mm. This is the baseline group, so the model estimate for an observation in the OJ and 0.5 mg/day dosage is simply  $\hat{y}_{i,OJ,0.5mg} = \hat{\alpha} = 13.23$  microns. For the OJ and 2 mg dosage estimate that has a value over 25 microns in the plot, the model incorporates the deviation for the 2 mg dosage:  $\hat{y}_{i,OJ,2mg} = \hat{\alpha} + \hat{\tau}_{2mg} = 13.23 + 12.83 = 26.06$  microns. For the Vitamin C group, another coefficient becomes involved from its "main effect". For the VC and 0.5 mg dosage level, the estimate is approximately 8 microns. The pertinent model components are  $\hat{y}_{i,VC,0.5mg} = \hat{\alpha} + \hat{\gamma}_{VC} = 13.23 + (-5.25) = 7.98$  microns. Finally, when we consider non-baseline results for both groups, three coefficients are required to reconstruct the results in the plot. For example, the estimate for the VC, 1 mg dosage is  $\hat{y}_{i,VC,1mg} = \hat{\alpha} + \hat{\tau}_{1mg} + \hat{\gamma}_{VC} = 13.23 + 9.47 + (-5.25) = 17.45$  microns. We usually will by-pass all this fun(!) with the coefficients in an interaction model and go from the ANOVA interaction test to focusing on the pattern of the responses in the interaction plot, but it is good to know that there are still model coefficients driving our results.

```
plot(allEffects(TG2), grid=T, multiline=T, ci.style="bars")
```

Given the presence of an important interaction, then the final step in the interpretation here is to interpret the results in the interaction plot or term-plot of the interaction model, supported by the p-value suggesting evidence of a different effect of supplement type based on the dosage level. To supplement this even more, knowing which combinations of levels differ can enhance our discussion. Tukey's HSD results (specifically the

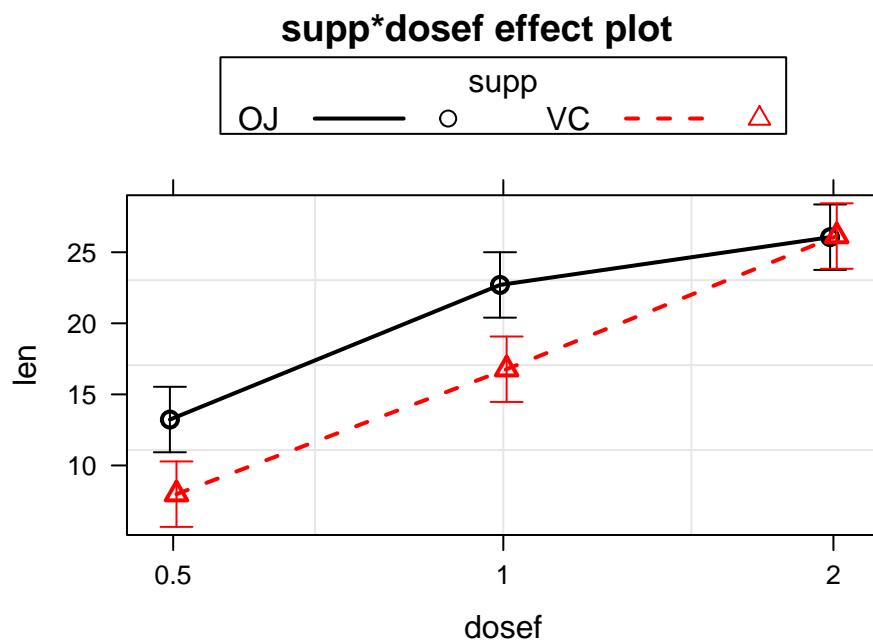


Figure 4.10: Term-plot for the estimated interaction for the Tooth Growth data.

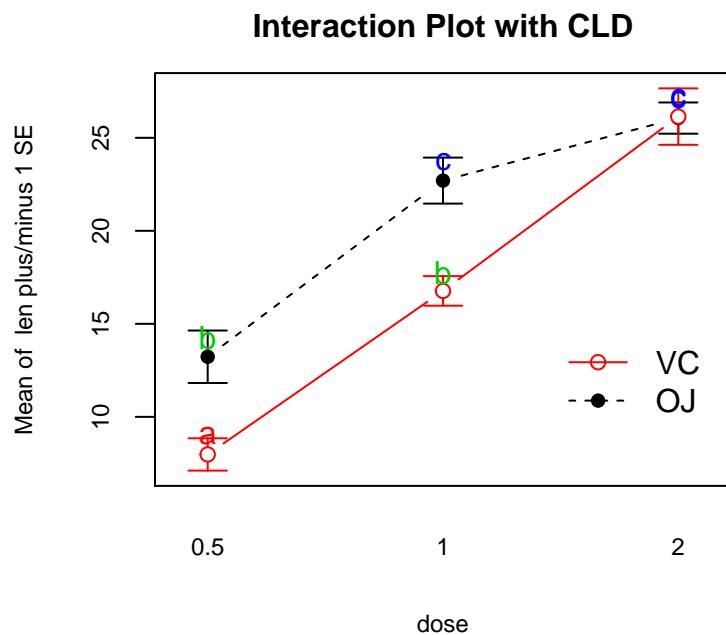


Figure 4.11: Interaction plot with added CLD from Tukey's HSD.

CLD) can be added to the original interaction plot by turning on the `cld=T` option in the `intplot` function as seen in Figure 4.11. Sometimes it is hard to see the letters and so there is also a `cldshift=...` option to move the letters up or down, here a value of 1 seemed to work.

```
intplot(len ~ supp*dose, data=ToothGrowth, col=c(1,2), cldshift=1,
       cld=T, main="Interaction Plot with CLD")
```

The interpretation of the previous hypothesis test result can be concluded with the following discussion. Generally increasing the dosage increases the amount of mean growth except for the 2 mg/day dosage level where the increase levels off in the OJ group (OJ 1 and 2 mg/day are not detectably different) and the differences between the two delivery methods disappear at the highest dosage level. But for 0.5 and 1 mg/day dosages, OJ is clearly better than VC by about 10 microns of growth on average.

## 4.5 Observational study example: The Psychology of Debt

In this section, the analysis of a survey of  $N = 464$  randomly sampled adults will be analyzed from a survey conducted by Lea and Walker (1995) and available in the `debt` data set from the `faraway` package (Faraway, 2016). The subjects responded to a variety of questions including whether they buy cigarettes (`cigbuy`: 0 if no, 1 if yes), their housing situation (`house`: 1 = rent, 2 = mortgage, and 3 = owned outright), their income group (`incomegp`: 1 = lowest, 5 = highest), and their score on a continuous scale of attitudes about debt (`prodebt`: 1 = least favorable, 5 = most favorable). `prodebt` was derived as the average of a series of questions about debt with each question measured on an *ordinal* 1 to 5 scale, with higher values corresponding to more positive responses about **going into debt** of various kinds. The ordered scale on surveys that try to elicit your opinions on topics with scales from 1 to 5 or 1 to 7 or even, sometimes, 1 to 10 is called a **Likert scale** (Likert, 1932). It is not a quantitative scale and really should be handled more carefully than taking an average of a set responses. That said, it is extremely common practice in social science research to treat ordinal responses as if they are quantitative and take the average of many of them to create a more continuous response variable like the one we are using here. If you continue your statistics explorations, you will see some better techniques for analyzing responses obtained in this fashion. That said, the scale of the response is relatively easy to understand as an amount of willingness to go into debt on a scale from 1 to 5 with higher values corresponding to more willingness to be in debt.

This data set is typical of survey data where respondents were not required to answer all questions and there are some missing responses. We will clean out any individuals that failed to respond to all questions using the `na.omit` function, which will return only subjects that responded to every question in the data set. But is this dangerous? Suppose that people did not want to provide their income levels if they were in the lowest or, maybe, highest income groups. Then we would be missing responses systematically and conclusions could be biased because of ignoring these types of subjects. This is another topic for more advanced statistical methods to try to handle but something every researcher should worry about when selected subjects do not respond at all or fail to answer some questions. Is there bias because of responses that were not observed that could invalidate all my hard won statistical conclusions? This ties back into our discussion of who was sampled. We need to think carefully about who was part of the sample but refused to participate and how that might impact our inferences.

Ignoring this potential for bias in the results for the moment, we are first interested in whether buying cigarettes/not and income groups interact in their explanation of the respondent's mean opinions on being in debt. The interaction plot (Figure 4.12) may suggest an interaction between `cigbuy` and `incomegp` from income levels 1 to 3 where the lines cross but it is not as clear as the previous examples. The interaction *F*-test helps us objectively assess evidence for that interaction. Based on the plot, there do not appear to be differences based on cigarette purchasing but there might be some differences between the income groups. If there is no interaction present, then this suggests that we might be in Scenario 2 or 3 where a single main effect of interest is present.

```
require(faraway)
data(debt)
debt$incomegp <- factor(debt$incomegp)
```

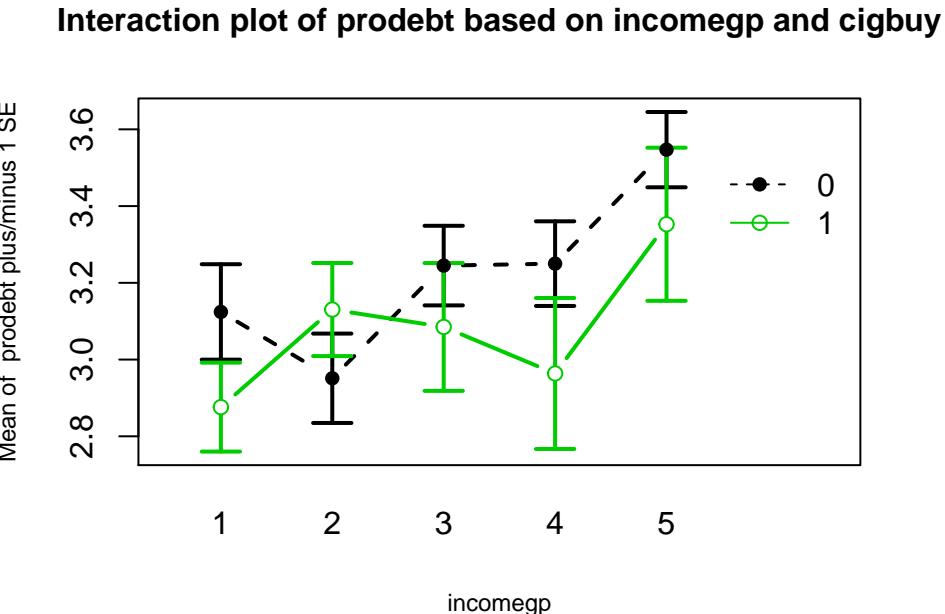


Figure 4.12: Interaction plot of prodebt by income group and buy cigarettes (0=no, 1=yes).

```
debt$cigbuy <- factor(debt$cigbuy)
debtc <- na.omit(debt)
intplot(prodebt ~ cigbuy*incomegp, data=debtc, col=c(1,3), lwd=2)
```

As in other situations, and especially with observational studies where a single large sample is analyzed, it is important to check for balance - whether all the combinations of the two predictor variables are similarly represented. Even more critically, we need to check whether all the combinations of levels of factors are measured. If a combination is not measured, then we lose the ability to estimate the mean for that combination and the ability to test for an interaction. A solution to that problem would be to collapse the categories of one of the variables, changing the definitions of the levels but if you fail to obtain information for all combinations, you can't work with the interaction model. In this situation, we barely have enough information to proceed (the smallest  $n_{jk}$  is 8 for income group 4 that buys cigarettes). We have a very unbalanced design with counts between 8 and 51 in the different combinations.

```
tally(cigbuy ~ incomegp, data=debtc)
```

```
##      incomegp
## cigbuy 1 2 3 4 5
##      0 24 40 45 47 51
##      1 23 29 18  8 19
```

The test for the interaction is always how we start our modeling in Two-Way ANOVA situations. The ANOVA table suggests that there is little evidence of interaction between the income level and buying cigarettes on the opinions of the respondents towards debt ( $F(4, 294) = 1.0003$ ,  $p\text{-value}=0.408$ ). This suggests that the initial assessment that the interaction wasn't too prominent was correct. We should move to the additive model here but first need to check the assumptions to make sure we can trust this initial test.

```
require(car)
debt1 <- lm(prodebt ~ incomegp*cigbuy, data=debtc)
Anova(debt1)
```

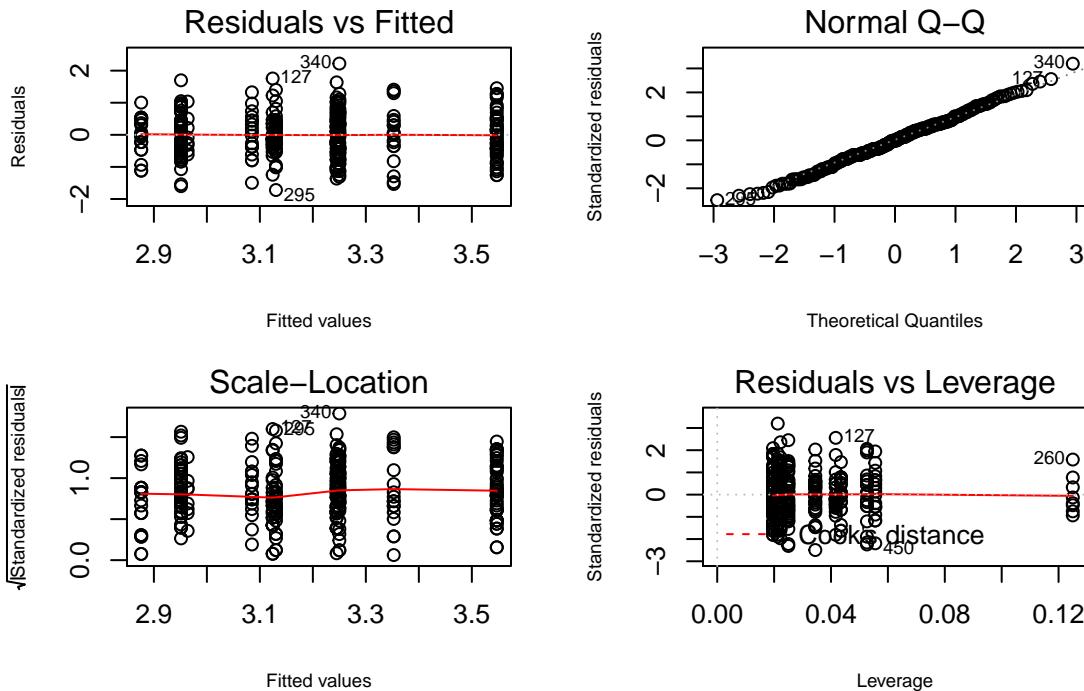


Figure 4.13: Diagnostic plot for `prodebt` by income group and buy cigarettes/not interaction model.

```
## Anova Table (Type II tests)
##
## Response: prodebt
##              Sum Sq Df F value    Pr(>F)
## incomegp      9.018  4 4.5766 0.001339
## cigbuy       0.703  1 1.4270 0.233222
## incomegp:cigbuy 1.971  4 1.0003 0.407656
## Residuals   144.835 294
```

The diagnostic plots (Figure 4.13) seem to be pretty well-behaved with no apparent violations of the normality assumption and no clear evidence of a violation of the constant variance assumption. The observations would seem to be independent because there is no indication of structure to the measurements of the survey respondents that might create dependencies. In observational studies, violations of the independence assumption might come from repeated measures of the same person or multiple measurements within the same family/household or samples that are clustered geographically. The random sampling from a population should allow inferences to a larger population except for that issue of removing partially missing responses. We also don't have much information on the population sampled, so will just leave this vague here but know that there is a population these conclusions apply to since it was random sample. All of this suggests proceeding to fitting and exploring the additive model is reasonable here. No causal inferences are possible because this is an observational study.

```
par(mfrow=c(2,2))
plot(debt1)
```

### 1. Hypotheses (Two sets apply when the additive model is the focus!):

- $H_0$ : No difference in means for `prodebt` for income groups in population, given cigarette buying in model

$\Leftrightarrow$  All  $\tau_j$ 's = 0 in additive model.

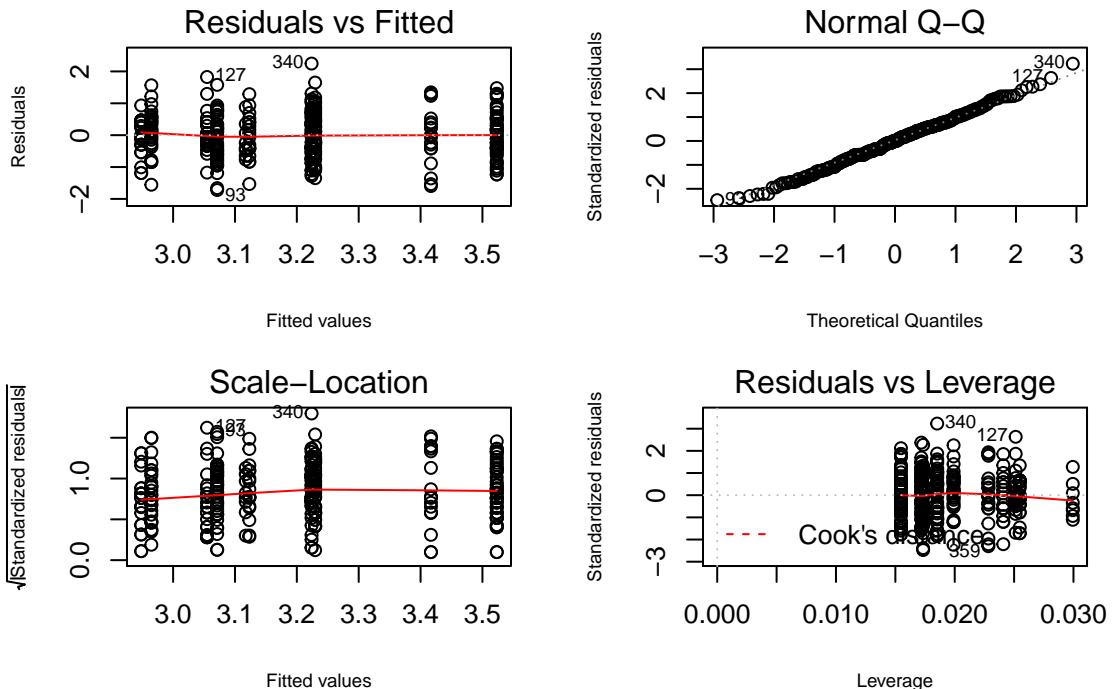


Figure 4.14: Diagnostic plot for prodebt by income group and buy cigarettes/not

- $H_A$ : Some difference in means for prodebt for income group in population, given cigarette buying in model

$\Leftrightarrow$  Not all  $\tau_j$ 's = 0 in additive model.

- $H_0$ : No difference in means for prodebt for cigarette buying/not in population, given income group in model

$\Leftrightarrow$  All  $\gamma_k$ 's = 0 in additive model.

- $H_A$ : Some difference in means for prodebt for cigarette buying/not in population, given income group in model

$\Leftrightarrow$  Not all  $\gamma_k$ 's = 0 in additive model.

## 2. Validity conditions – discussed above but with new plots for the additive:

```
debt1r<-lm(prodebt~incomegp+cigbuy, data=debtc)
plot(debt1r)
```

- Constant Variance:

– In the Residuals vs Fitted and the Scale-Location plots in Figure 4.14, the differences in variability among groups is minor and nothing suggests a violation. **If you change models, you should always revisit the diagnostic plots to make sure you didn't create problems that were not present in more complicated models.**

- Normality of residuals:

– The QQ-Plot in Figure 4.14 does not suggest a problem with this assumption.

## 3. Calculate the test statistic for the two main effect tests.

```
Anova(debt1r)
```

```
## Anova Table (Type II tests)
##
## Response: prodebt
##           Sum Sq Df F value Pr(>F)
## incomegp    9.018  4  4.5766 0.001335
## cigbuy      0.703  1  1.4270 0.233210
## Residuals 146.806 298
```

- The test statistics are  $F(4, 298) = 4.577$  and  $F(1, 298) = 1.427$ .

#### 4. Find the p-value:

- The ANOVA  $F$ -test p-values are 0.001335 for the income group variable (conditional on cigarette buy) and 0.2232 for the cigarette buy variable (conditional on income group).

#### 5. Make decisions:

- Reject  $H_0$  of no income group differences (p-value=0.0013) and fail to reject  $H_0$  of no cigarette buying differences (p-value=0.2232), each after controlling for the other variable.

#### 6. Write a conclusion:

- There was initially no evidence to support retaining the interaction of income group and cigarette buying on pro-debt feelings ( $F(4, 294) = 1.00$ , p-value = 0.408) so the interaction was dropped from the model. There is strong evidence of some difference in the mean pro-debt feelings in the population across the income groups, after adjusting for cigarette buying. There is little to no evidence of a difference in the mean pro-debt feelings in the population based on cigarette buying/not, after adjusting for income group.

So we learned that the additive model was more appropriate for these responses and that the results resemble Scenario 2 or 3 with only one main effect being important. In the additive model, the coefficients can be interpreted as shifts from the baseline after controlling for the other variable in the model. Figure 4.15 shows the increasing average comfort with being in debt as the income groups go up. Being a cigarette buyer was related to a lower comfort level with debt. But compare the y-axis scales in the two plots – the differences in the means across income groups are almost 0.5 points on a 5 point scale whereas the difference across `cigbuy`'s two levels is less than 0.15 units. The error bars for the 95% confidence intervals are of similar width but the differences in means show up clearly in the income group term-plot. This is all indirectly related to the size of the p-values for each term in the additive model but hopefully helps to build some intuition on the reason for differences.

```
plot(allEffects(debt1r))
```

The estimated coefficients can also be interesting to interpret for the additive model. Here are the model summary coefficients:

```
summary(debt1r)$coefficients
```

	Estimate	Std. Error	t value	Pr(> t )
## (Intercept)	3.05483517	0.11127284	27.4535561	1.357428e-83
## incomegp2	0.01640636	0.13288796	0.1234601	9.018260e-01
## incomegp3	0.17477182	0.13649309	1.2804444	2.013846e-01
## incomegp4	0.16900515	0.14274836	1.1839376	2.373813e-01
## incomegp5	0.46833117	0.13377661	3.5008449	5.347171e-04
## cigbuy1	-0.10640228	0.08907259	-1.1945569	2.332101e-01

In the model, the baseline group is for non-cigarette buyers (`cigbuy=0`) and income group 1 with  $\hat{\alpha} = 3.055$  points. Regardless of the `cigbuy` level, the difference between income groups 2 and 1 is estimated to be  $\hat{\tau}_2 = 0.016$ , an increase in the mean score of 0.016 points. Similarly, the difference between income groups 3 and 1 is  $\hat{\tau}_3 = 0.175$  points, regardless of cigarette smoking status. The estimated difference between cigarette buyers and non-buyers was estimated as  $\hat{\tau}_2 = -0.106$  points for any income group, remember that this variable had a moderately large p-value in this model. The additive model-based estimates for all six combinations can be found in Table 4.3.

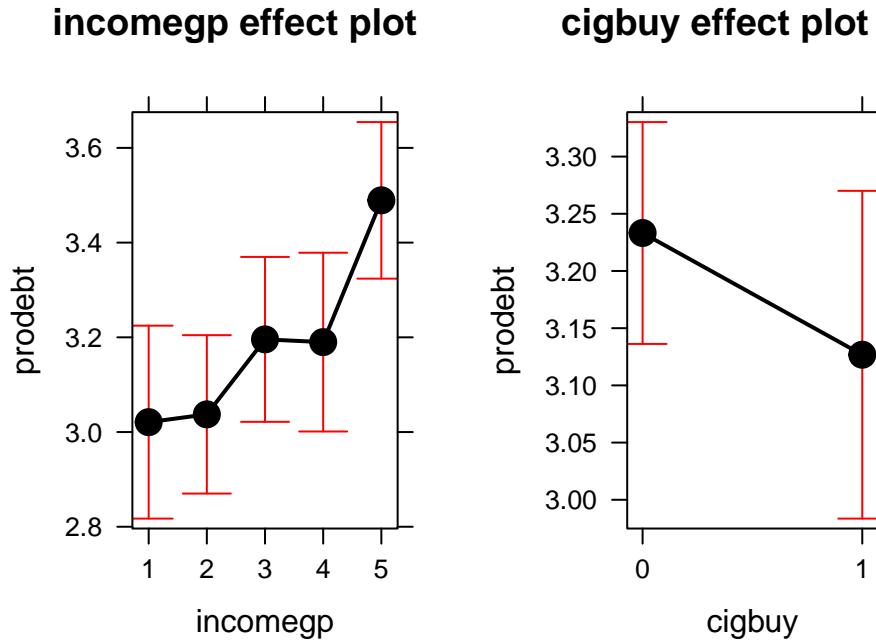


Figure 4.15: Term-plots for the `prodebt` additive model with left panel for income group and the right panel for buying cigarettes or not (1 for yes).

Table 4.3: Calculations to construct the estimates for all combinations of variables for the `prodebt` additive model.

Cig Buy	Income Group 1	Income Group 2	Income Group 3	Income Group 4	Income Group 5
0:No	$\hat{\alpha} = 3.055$	$\hat{\alpha} + \hat{\tau}_2$ $= 3.055 + 0.016$ $= 3.071$	$\hat{\alpha} + \hat{\tau}_3$ $= 3.055 + 0.175$ $= 3.230$	$\hat{\alpha} + \hat{\tau}_4$ $= 3.055 + 0.169$ $= 3.224$	$\hat{\alpha} + \hat{\tau}_5$ $= 3.055 + 0.468$ $= 3.523$
1:Yes	$\hat{\alpha} + \hat{\gamma}_2$ $= 3.055$ $- 0.106$ $= 2.949$	$\hat{\alpha} + \hat{\tau}_2 + \hat{\gamma}_2$ $= 3.055 + 0.016$ $- 0.106$ $= 2.965$	$\hat{\alpha} + \hat{\tau}_3 + \hat{\gamma}_2$ $= 3.055 + 0.175$ $- 0.106$ $= 3.124$	$\hat{\alpha} + \hat{\tau}_4 + \hat{\gamma}_2$ $= 3.055 + 0.169$ $- 0.106$ $= 3.118$	$\hat{\alpha} + \hat{\tau}_5 + \hat{\gamma}_2$ $= 3.055 + 0.468$ $- 0.106$ $= 3.417$

One final plot of the fitted values from this additive model in Figure 4.16 hopefully crystallizes the implications of an additive model and reinforces that this model creates and assumes that the differences across levels of one variable are the same regardless of the level of the other variable and that this creates parallel lines. The difference between `cigbuy` levels across all income groups is a drop in -0.106 points. The income groups have the same differences regardless of cigarette buying or not, with income group 5 much higher than the other four groups.

**In general, we proceed through the following steps in any 2-WAY ANOVA situation:**

1. Make an interaction plot.
2. Fit the interaction model; examine the test for the interaction.
3. Check the residual diagnostic plots for the interaction model (especially normality and equal variance).
  - If there is a problem with normality or equal variance, consider a “transformation” of the response as discussed in Chapter 7. This can help make the responses have similar variances or responses to be more normal, but sometimes not both.

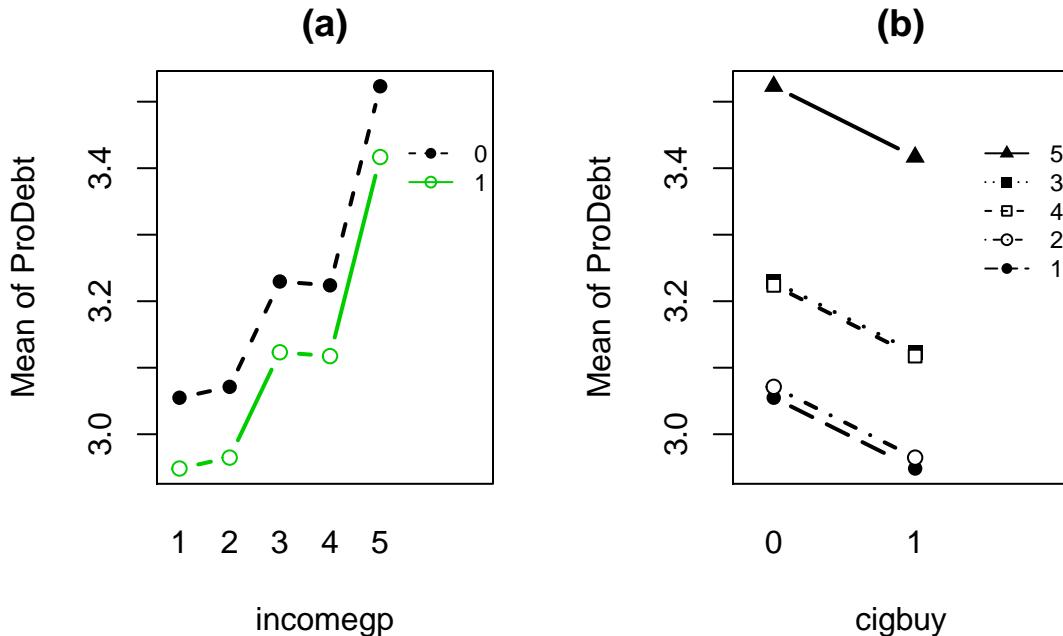


Figure 4.16: Illustration of the results from Table 4.2 showing the combined impacts of the components of the additive model for `prodebt`. Panel (a) uses income groups on the x-axis and different lines for cigarette buyers (1) or not (0). Panel (b) displays the different income groups as lines with the cigarette buying status on the x-axis.

4. If the interaction test has a small p-value, that is your main result. Focus on the interaction plot from (1) to fully understand the results, adding Tukey's HSD results to see which means of the combinations of levels are detected as being different.
5. If the interaction is not considered important, then re-fit the model without the interaction (additive model) and re-check the diagnostic plots. If the diagnostics are reasonable to proceed:
  - Focus on the results for each explanatory variable, using Type II tests especially if the design is not balanced.
  - Report the initial interaction test results and the results for the test for each variable from the model that is re-fit without the interaction.
  - Model coefficients are interesting as they are shifts from baseline for each level of each variable, controlling for the other variable – interpret those differences if the number of levels is not too great.

Whether you end up favoring an additive or interaction model, all steps of the hypothesis testing protocol should be engaged.

## 4.6 Pushing Two-Way ANOVA to the limit: Un-replicated designs

In some situations, it is too expensive to replicate combinations of treatments and only one observation at each combination of the two explanatory variables, A and B, is possible. In these situations, even though we have information about all combinations of A and B, it is no longer possible to test for an interaction. Our regular rules for degrees of freedom show that we have nothing left for the error degrees of freedom and so we have to drop the interaction and call that potential interaction variability “error”.

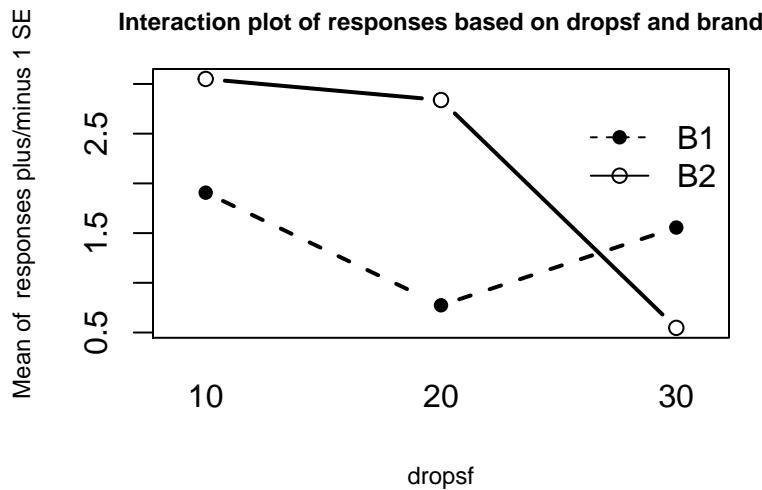


Figure 4.17: Interaction plot in paper towel data set with no replication.

We can still perform an analysis of the responses but an issue occurs with trying to estimate the interaction  $F$ -test statistic – we run out of degrees of freedom for the error. To illustrate these methods, the paper towel example is revisited except that only one response for each combination is used. Now the entire data set can be displayed:

```
ptR<-read.csv("http://www.math.montana.edu/courses/s217/documents/ptR.csv")
ptR$dropsf<-factor(ptR$drops)
ptR
```

```
##   brand drops responses dropsf
## 1   B1    10  1.9064356    10
## 2   B2    10  3.0504173    10
## 3   B1    20  0.7737965    20
## 4   B2    20  2.8384124    20
## 5   B1    30  1.5557071    30
## 6   B2    30  0.5470565    30
```

Upon first inspection the interaction plot in Figure 4.17 looks like there might be some interesting interactions present. But remember now that there is only a single observation at each combination of the brands and water levels so there is not much power to detect differences in this sort of situation and no replicates at combination that allow estimation of SEs so no bands are produced in the plot.

```
par(mfrow=c(1,1))
intplot(responses~brand*dropsf,data=ptR,lwd=2)
```

The next step would be to assess the statistical evidence related to an interaction between Brand and Drops. A problem will arise in trying to form the ANOVA table as you would see this in the console:

```
> anova(lm(responses~dropsf*brand,data=ptR))
Analysis of Variance Table
Response: responses
  Df  Sum Sq Mean Sq F value Pr(>F)
dropsf      2  2.03872 1.01936
brand       1  0.80663 0.80663
dropsf:brand 2  2.48773 1.24386
Residuals    0  0.00000
Warning message:
In anova.lm(lm(responses ~ dropsf * brand, data = ptR)) :
  ANOVA F-tests on an essentially perfect fit are unreliable
```

Warning messages in R output show up in red after you run functions that contain problems and are generally not a good thing, but can sometimes be ignored. In this case, the warning message is not needed – there are no  $F$ -statistics or p-values in the results so we know there are some issues with the results. The bolded line is key here squares of 0. Without replication, there are no degrees of freedom left to estimate the residual error. My (Greenwood's) first statistics professor, Gordon Bril at Luther College, used to refer to this as “shooting your load” by fitting too many terms in the model given the number of observations available. Maybe this is a bit graphic but hopefully will help you remember the need for replication if you want to estimate interactions – it did for me. Without replication of observations, we run out of information to estimate and test all the desired model components.

So what can we do if we can't afford replication? We can *assume* that the interaction does not exist and use those degrees of freedom and variability as the error variability. When we drop the interaction from Two-Way models, the interaction variability is added into the  $SS_E$  so this is interaction between the variables. We are not able to test for an interaction so must rely on the interaction plot to assess whether an interaction might be present. Figure 4.17 suggests there might be an interaction in these data (the two brands lines cross noticeably suggesting non-parallel lines). So in this case, assuming no interaction is present is hard to justify. But if we proceed under this dangerous assumption, tests for the main effects can be developed.

```
norep1<-lm(responses~dropsf+brand,data=ptR)
Anova(norep1)
```

```
## Anova Table (Type II tests)
##
## Response: responses
##             Sum Sq Df F value Pr(>F)
## dropsf      2.03872  2  0.8195 0.5496
## brand       0.80663  1  0.6485 0.5052
## Residuals  2.48773  2
```

In the additive model, the last row of the ANOVA table that is called the **Residuals** row is really the interaction row from the interaction model ANOVA table. Neither main effect had a small p-value (**Drops**:  $F(2, 2) = 0.82$ , p-value = 0.55 and **Brand**:  $F(1, 2) = 0.65$ , p-value = 0.51) in the additive model. To get small p-values with small sample sizes, the differences would need to be **very** large because the residual degrees of freedom have become very small. The term-plots in Figure 4.18 show that the differences among the levels are small relative to the residual variability as seen in the error bars around each point estimate.

```
plot(allEffects(norep1))
```

Hopefully by pushing the limits there are two conclusions available from this section. First, replication is important, both in being able to perform tests for interactions and for having enough power to detect differences for the main effects. Second, dropping from the interaction model to additive model, the variability explained by the interaction term is pushed into the error term, whether replication is available or not.

## 4.7 Chapter summary

In this chapter, methods for handling two different categorical predictors in the same model with a continuous response were developed. The methods build on techniques from Chapter 3 for the One-Way ANOVA and there are connections between the two models. This was most clearly seen in the guinea pig data set that was analyzed in both chapters. When two factors are available, it is better to start with the methods developed in this chapter because the interaction between the factors can, potentially, be separated from their main effects. The additive model is easier to interpret but should only be used when no evidence of an interaction is present. When an interaction is determined to be present, the main effects should not be interpreted and the interaction plot in combination with Tukey's HSD provides information on the important aspects of the results.

- If the interaction is retained in the model, there are two things you want to do with interpreting the interaction:

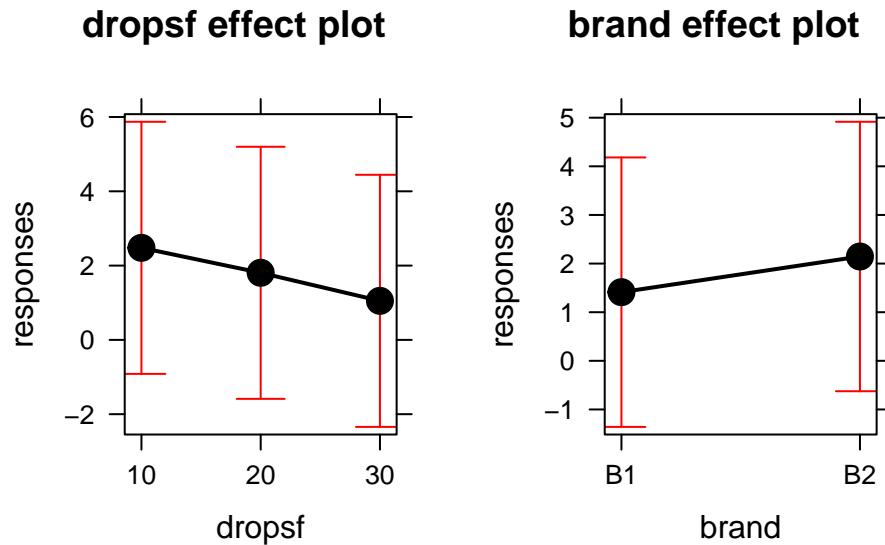


Figure 4.18: Term-plots for the additive model in paper towel data set with no replication.

1. Describe the interaction, going through the changes from left to right in the interaction plot or term-plot for each level of the other variable.
2. Suggest optimal combinations of the two variables to either get the highest or lowest possible responses.
  - a. For example, you might want to identify a dosage and delivery method for the guinea pigs to recommend and one to avoid if you want to optimize odontoblast growth.
- If there is no interaction, then the additive model provides information on each of the variables and the differences across levels of each variable are the same regardless of the levels of the other variable.
  - You can describe the deviations from baseline as in Chapter 3, but for each variable, noting that you are controlling for the variable.

Some statisticians might have different recommendations for dealing with interactions and main effects, especially in the context of evidence of an interaction. We have chosen to focus on tests for interactions to screen for “real” interactions and then interpret the interaction plots aided by the Tukey’s HSD for determining which combinations of levels are detectably different. Others might suggest exploring the main effects tests even with interactions present. In some cases, those results are interesting but in others the results can be misleading and wanted to avoid trying to tell you when that might happen. Consider two scenarios, one where the main effects have large p-values but the interaction has a small p-value and the other where the main effects and the interaction all have small p-values. The methods discussed in this chapter allow us to effectively arrive at the interpretation of the differences in the results across the combinations of the treatments due to the interaction having a small p-value. The main effects results are secondary results at best when the interaction is important because we know that impacts of one explanatory variable is changing based on the levels of the other variable.

Chapter 5 presents a bit of a different set of statistical methods that allow analyses of data sets similar to those considered in the last two chapters but with a categorical response variable. The methods are very different but are quite similar in overall goals to those in Chapter 3 where differences in responses were explored across groups. After Chapter 5, the rest of the semester will return to fitting models using the `lm` function as used here, but incorporating quantitative predictor variables and then eventually incorporating both categorical and quantitative predictor variables. The methods in Chapter 8 are actually quite similar to those considered here.

## 4.8 Important R code

The main components of R code used in this chapter follow with components to modify in red, remembering that any R packages mentioned need to be installed and loaded for this code to have a chance of working:

- `tally(A~B, data=DATASETNAME)`
  - Requires the `mosaic` package be loaded.
  - Provides the counts of observations in each combination of categorical predictor variables A and B, used to check for balance and understand sample sizes in each combination.
- `DATASETNAME$VARIABLENAME <- factor(DATASETNAME$VARIABLENAME)`
  - Use the `factor` function on any numerically coded explanatory variable where the numerical codes represent levels of a categorical variable.
- `intplot(Y~A*B, data=DATASETNAME)`
  - Download and install using:
 

```
source("http://www.math.montana.edu/courses/s217/documents/intplot.R")
```
  - Provides interaction plot.
- `INTERACTIONMODELNAME <- lm(Y~A*B, data=DATASETNAME)`
  - Fits the interaction model with main effects for A and B and an interaction between them.
  - This is the first model that should be fit in Two-Way ANOVA modeling situations.
- `ADDITIVEMODELNAME <- lm(Y~A+B, data=DATASETNAME)`
  - Fits the additive model with only main effects for A and B but no interaction between them.
  - Should only be used if the interaction has been decided to be unimportant using a test for the interaction.
- `summary(MODELNAME)`
  - Generates model summary information including the estimated model coefficients, SEs, t-tests, and p-values.
- `Anova(MODELNAME)`
  - Requires the `car` package to be loaded.
  - Generates a Type II Sums of Squares ANOVA table that is useful for both additive and interaction models, but it most important to use when working with the additive model as it provides inferences for each term conditional on the other one.
- `par(mfrow=c(2,2)); plot(MODELNAME)`
  - Generates four diagnostic plots including the Residuals vs Fitted and Normal Q-Q plot.
- `plot(allEffects(MODELNAME))`
  - Requires the `effects` package be loaded.
  - Plots the results from the estimated model.

## 4.9 Practice problems

To practice the Two-Way ANOVA, consider a data set on  $N = 861$  ACT Mathematics Usage Test scores from 1987. The test was given to a sample of high school seniors who met one of three profiles of high school mathematics course work: (a) Algebra I only; (b) two Algebra courses and Geometry; and (c) two Algebra courses, Geometry, Trigonometry, Advanced Mathematics, and Beginning Calculus. These data were generated from summary statistics for one particular form of the test as reported by Doolittle and Welch (1989). The source of this version of the data set is Ramsey and Schafer (2012) and the `Sleuth2` package (Ramsey et al., 2016). First install and then load that package.

```
require(Sleuth2)
require(mosaic)
math <- ex1320
names(math)
favstats(Score ~ Sex+Background, data=math)
```

- 4.1. Use the `favstats` summary to discuss whether the design was balanced or not.
- 4.2. Make a side-by-side beanplot and interaction plot of the results and discuss the relationship between Sex, Background, and ACT Score.
- 4.3. Write out the interaction model in terms of the Greek letters, making sure to define all the terms and don't forget the error terms in the model.
- 4.4. Fit the interaction plot and find the ANOVA table. For the test you should consider first (the interaction), write out the hypotheses, report the test statistic, p-value, distribution of the test statistic under the null, and write a conclusion related to the results of this test.
- 4.5. Re-fit the model as an additive model (why is this reasonable here?) and use `Anova` to find the Type II sums of squares ANOVA. Write out the hypothesis for the Background variable, report the test statistic, p-value, distribution of the test statistic under the null, and write a conclusion related to the results of this test.
- 4.6. Use the `effects` package to make a term-plot from the additive model from 4.5 and discuss the results. Specifically, discuss what you can conclude about the average relationship across both sexes, between Background and average ACT score?
- 4.7. Make our standard diagnostic plots and assess the assumptions using these plots. Can you assess independence using these plots? Discuss this assumption in this situation.
- 4.8. Use the estimated model coefficients to determine which of the combinations of levels provides the highest estimated average score.



# Chapter 5

## Chi-square tests

### 5.1 Situation, contingency tables, and plots

In this chapter, the focus shifts briefly from analyzing quantitative response variables to methods for handling categorical response variables. This is important because in some situations it is not possible to measure the response variable quantitatively. For example, we will analyze the results from a clinical trial where the results for the subjects were measured as one of three categories: *no improvement*, *some improvement*, and *marked improvement*. While that type of response could be treated as numerical, coded possibly as 1, 2, and 3, it would be difficult to assume that the responses such as those follow a normal distribution since they are *discrete* (not continuous, measured at whole number values only) and, more importantly, the difference between *no improvement* and *some improvement* is not necessarily the same as the difference between *some* and *marked improvement*. If it is treated numerically, then the differences are assumed to be the same unless a different coding scheme is used (say 1, 2, and 5). It is better to treat these types of responses as being in one of the three categories and use statistical methods that don't make unreasonable assumptions about what the numerical coding might mean. The study being performed here involved subjects randomly assigned to either a treatment or a placebo (control) group and we want to address research questions similar to those considered in Chapters 2 and 3 – assessing differences among two or more groups. With quantitative responses, the differences in the distributions are parameterized via the means of the groups and we used 2-sample mean or ANOVA hypotheses and tests. With categorical responses, the focus is on the probabilities of getting responses in each category and whether they differ among the groups.

We start with some useful summary techniques, both numerical and graphical, applied to some examples of studies these methods can be used to analyze. Graphical techniques provide opportunities for assessing specific patterns in variables, relationships between variables, and for generally understanding the responses obtained. There are many different types of plots and each can enhance certain features of data. We will start with a “fun” display, called a tableplot, to help us understand some aspects of the results from a double-blind randomized clinical trial investigating a treatment for rheumatoid arthritis that has the categorical response variable introduced previously. These data are available in the `Arthritis` data set available in the `vcd` package (Meyer et al., 2016). There were  $n = 84$  subjects, with some demographic information recorded along with the `Treatment` status (*Treated*, *Placebo*) and whether the patients’ arthritis symptoms `Improved` (with levels of *None*, *Some*, and *Marked*).

The `tableplot` function from the `tabplot` package (Tennekes and de Jonge, 2017) displays bars for each response in a row<sup>1</sup> based on the category of responses or as a bar with the height corresponding the value of quantitative variables. It also plots a red cell if the observations were missing on a particular variable. The plot can be obtained simply as `tableplot(DATASETNAME)`. But when using `tableplot`, we may not want to display everything in the `data.frame` and often just select some of the variables. We use `Treatment`, `Improved`,

---

<sup>1</sup>In larger data sets, multiple subjects are displayed in each row as proportions of the rows in each category.

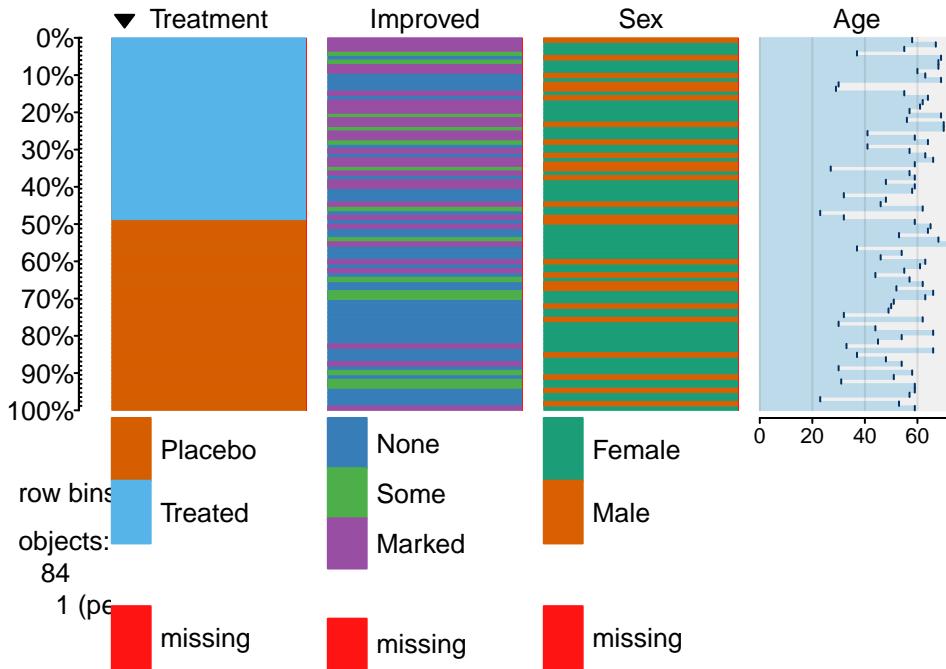


Figure 5.1: Table plot of the arthritis data set.

Sex, and Age in the `select=...` option with a `c()` and commas between the names of the variables we want to display. The first one in the list is also the one that the data are sorted based on.

```
require(vcd)
data(Arthritis) #Double-blind clinical trial with treatment and control groups
#Homogeneity example
require(tabplot)
tableplot(Arthritis, select=c(Treatment, Improved, Sex, Age))
```

The first thing we can gather from Figure 5.1 is that there are no red cells so there were no missing observations in the data set. Missing observations regularly arise in real studies when observations are not obtained for many different reasons and it is always good to check for missing data issues – this plot provides a quick visual method for doing that check. Primarily we are interested in whether the treatment led to a different pattern (or rates) of improvement responses. There seems to be more purple (*Marked*) improvement responses in the treatment group and more blue (*None*) responses in the placebo group. This sort of plot also helps us to simultaneously consider the role of other variables in the observed responses. You can see the sex of each subject in the vertical panel for `Sex` and it seems that there is a relatively reasonable mix of males and females in the treatment/placebo groups. Quantitative variables are also displayed with horizontal bars corresponding to the responses. From the panel for `Age`, we can see that the ages of subjects ranged from the 20s to 70s and that there is no clear difference in the ages between the treated and placebo groups. If, for example, all the male subjects had ended up being randomized into the treatment group, then we might have worried about whether sex and treatment were confounded and whether any differences in the responses might be due to sex instead of the treatment. The random assignment of treatment/placebo to the subjects appears to have been successful here with the ages and sexes appearing to be well-mixed among the two treatment groups. The main benefit of this sort of plot is the ability to visualize more than two categorical variables simultaneously. But now we want to focus more directly on the researchers' main question – does the treatment lead to different improvement outcomes than the placebo?

To directly assess the effects of the treatment, we want to display just the two variables of interest. **Stacked bar charts** provide a method of displaying the response patterns (in `Improved`) across the levels of a predictor variable(`Treatment`) by displaying a bar for each predictor variable level and the proportions of responses in

each category of the response in each of those groups. If the placebo is as effective as the treatment, then we would expect similar proportions of responses in each improvement category. A difference in the effectiveness would manifest in different proportions in the different improvement categories between *Treated* and *Placebo*. To get information in this direction, we start with obtaining the counts in each combination of categories using the `tally` function to generate contingency tables. *Contingency tables* with  $R$  rows and  $C$  columns (called ***R by C tables***) summarize the counts of observations in each combination of the explanatory and response variables. In these data, there are  $R = 2$  rows and  $C = 3$  columns making a  $2 \times 3$  table – note that you do not count the row and column for the “Totals” in defining the size of the table. In the table, there seems to be many more *Marked* improvement responses (21 vs 7) and fewer *None* responses (13 vs 29) in the treated group compared to the placebo group.

```
require(mosaic)
tally(~Treatment+Improved, data=Arthritis, margins=T)
```

```
##           Improved
## Treatment None Some Marked Total
##   Placebo   29    7     7    43
##   Treated   13    7    21    41
##   Total     42   14    28    84
```

Using the `tally` function with  $\sim x+y$  provides a contingency table with the  $x$  variable on the rows and the  $y$  variable on the columns, with `margins=T` as an option so we can obtain the totals along the rows, columns, and table total of  $N = 84$ . In general, contingency tables contain the counts  $n_{rc}$  in the  $r^{th}$  row and  $c^{th}$  column where  $r = 1, \dots, R$  and  $c = 1, \dots, C$ . We can also define the ***row totals*** as the sum across row  $r$  as

$$\mathbf{n}_{r\bullet} = \sum_{c=1}^C n_{rc},$$

the ***column totals*** as the sum across column  $c$  as

$$\mathbf{n}_{\bullet c} = \sum_{r=1}^R n_{rc},$$

and the ***table total*** as

$$\mathbf{N} = \sum_{r=1}^R \mathbf{n}_{r\bullet} = \sum_{c=1}^C \mathbf{n}_{\bullet c} = \sum_{r=1}^R \sum_{c=1}^C \mathbf{n}_{rc}.$$

We'll need these quantities to do some calculations in a bit. A generic contingency table with added row, column, and table totals just like the previous result from the `tally` function is provided in Table 5.1.

Table 5.1: General notation for counts in an  $R$  by  $C$  contingency table.

	Response Level 1	Response Level 2	Response Level 3	...	Response Level C	Totals
<b>Group 1</b>	$n_{11}$	$n_{12}$	$n_{13}$	...	$n_{1C}$	$\mathbf{n}_{1\bullet}$
<b>Group 2</b>	$n_{21}$	$n_{22}$	$n_{23}$	...	$n_{2C}$	$\mathbf{n}_{2\bullet}$
...	...	...	...	...	...	...
<b>Group R</b>	$n_{R1}$	$n_{R2}$	$n_{R3}$	...	$n_{RC}$	$\mathbf{n}_{R\bullet}$
<b>Totals</b>	$\mathbf{n}_{\bullet 1}$	$\mathbf{n}_{\bullet 2}$	$\mathbf{n}_{\bullet 3}$	...	$\mathbf{n}_{\bullet C}$	$\mathbf{N}$

Comparing counts from the contingency table is useful, but comparing proportions in each category is better, especially when the sample sizes in the levels of the explanatory variable differ. Switching the formula used in the `tally` function formula to  $\sim y \mid x$  and adding the `format="proportion"` option provides

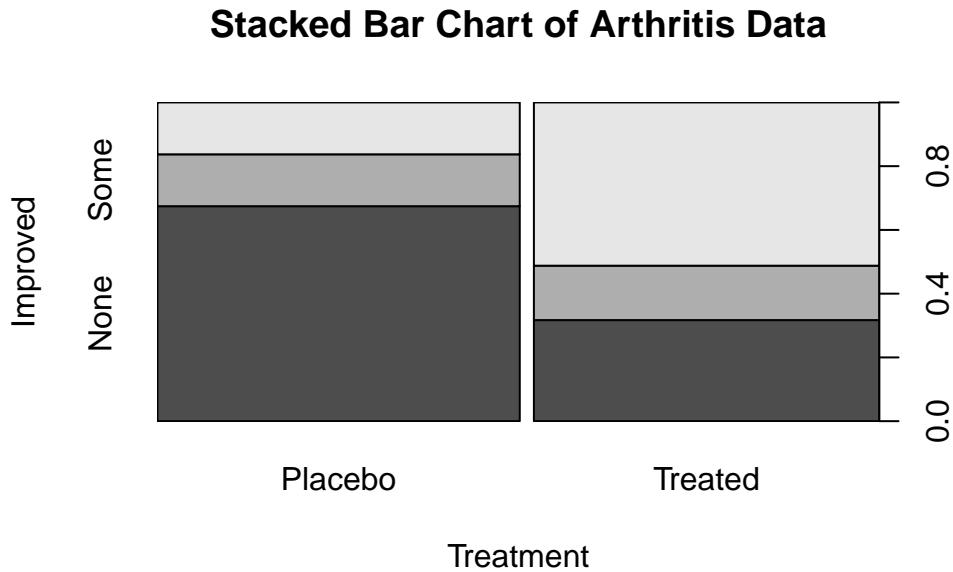


Figure 5.2: Stacked bar chart of Arthritis data. The left bar is for the Placebo group and the right bar is for the Treated group. The width of the bars is based on relative size of each group and the portion of the total height of each shaded area is the proportion of that group in each category. The darkest shading is for “none”, medium shading for “some”, and the lightest shading for “marked”, as labeled on the y-axis.

the proportions in the response categories conditional on the category of the predictor (these are called *conditional proportions* or the *conditional distribution* of, here, *Improved* on *Treatment*)<sup>2</sup>. Note that they sum to 1.0 in each level of x, *placebo* or *treated*:

```
tally(~Improved|Treatment, data=Arthritis, format="proportion", margins=T)
```

```
##          Treatment
## Improved Placebo Treated
##   None    0.6744186 0.3170732
##   Some    0.1627907 0.1707317
##  Marked   0.1627907 0.5121951
##   Total   1.0000000 1.0000000
```

Note that it switches the variables between the rows and columns from the first summary of the data but the single “Total” row makes it clear to read the proportions down the columns in this version of the table. In this application, it shows how the proportions seem to between the placebo and treatment groups. This matches the previous thoughts on this data set, but now a difference of marked improvement of 16% vs 51% is more clearly a big difference. We can also display this result using a *stacked bar-chart* that displays the same information, using the `plot` function with a y~x formula:

```
plot(Improved~Treatment, data=Arthritis,
     main="Stacked Bar Chart of Arthritis Data")
```

The stacked bar-chart in Figure 5.2 displays the previous conditional proportions for the groups, with the same relatively clear difference between the groups persisting. If you run the `plot` function with variables that are coded numerically, it will make a very different looking graph (R is smart!) so again be careful that you are instructing R to treat your variables as categorical if they really are categorical. R is powerful but can’t read your mind!

<sup>2</sup>The vertical line, “|”, in ~ y|x is available on most keyboards on the same key as “”. It is the mathematical symbol that means “conditional on” whatever follows.

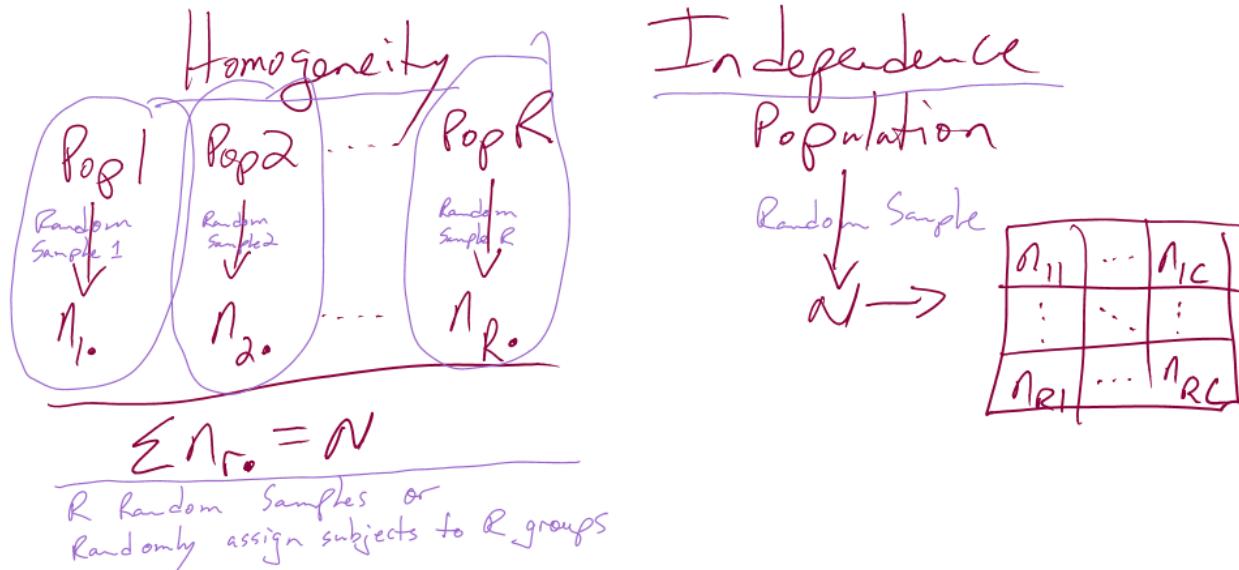


Figure 5.3: Diagram of the scenarios involved in Homogeneity and Independence tests. Homogeneity testing involves  $R$  random samples or subjects assigned to  $R$  groups. Independence testing involves a single random sample and measurements on two categorical variables.

In this chapter, we analyze data collected in two different fashions and modify the hypotheses to reflect the differences in the data collection processes, choosing either between what are called Homogeneity and Independence tests. The previous situation where levels of a treatment are randomly assigned to the subjects in a study describes the situation for what is called a **Homogeneity Test**. Homogeneity also applies when random samples are taken from each population of interest to generate the observations in each group of the explanatory variable. These sorts of situations resemble many of the examples from Chapter 3 where treatments were assigned to subjects. The other situation considered is where a single sample is collected to represent a population and then a contingency table is formed based on responses on two categorical variables. When one sample is collected and analyzed using a contingency table, the appropriate analysis is called an **Independence** or **Association test**. In this situation, it is not necessary to have variables that are clearly classified as explanatory or response although it is certainly possible. Data that often align with Independence testing are collected using surveys of subjects randomly selected from a single, large population. An example, analyzed below, involves a survey of voters and whether their party affiliation is related to who they voted for – the republican, democrat, or other candidate. There is clearly an explanatory variable of the *Party affiliation* but a single large sample was taken from the population of all likely voters so the Independence test needs to be applied. Another example where Independence is appropriate involves a study of student cheating behavior. Again, a single sample was taken from the population of students at a university and this determines that it will be an Independence test. Students responded to questions about lying to get out of turning in a paper and/or taking an exam (*none, either, or both*) and copying on an exam and/or turning in a paper written by someone else (*neither, either, or both*). In this situation, it is not clear which variable is the response or explanatory (which should explain the other) and it does not matter with the Independence testing framework. Figure 5.3 contains a diagram of the data collection processes and can help you to identify the appropriate analysis situation.

You will discover that the test statistics are the same for both methods, which can create some desire to assume that the differences in the data collection doesn't matter. In Homogeneity designs, the sample size in each group ( $n_{1\bullet}, n_{2\bullet}, \dots, n_{R\bullet}$ ) is fixed. In Independence situations, the total sample size  $N$  is fixed but all the  $n_{r\bullet}$ 's are random. These differences impact the graphs, hypotheses, and conclusions used even though the test statistics and p-values are calculated the same way – so we only need to learn one test statistic to handle the two situations, but we need to make sure we know which we're doing!

## 5.2 Homogeneity Test Hypotheses

If we define some additional notation, we can then define hypotheses that allow us to assess evidence related to whether the treatment “matters” in Homogeneity situations. This situation is similar to what we did in the One-Way ANOVA situation with quantitative responses in Chapter 3 but the parameters now relate to proportions in the response variable categories across the groups. First we can define the conditional population proportions in level  $c$  (column  $c = 1, \dots, C$ ) of group  $r$  (row  $r = 1, \dots, R$ ) as  $p_{rc}$ . Table 5.2 shows the proportions, noting that the proportions in each row sum to 1 since they are conditional on the group of interest. A *transposed* (rows and columns flipped) version of this table is produced by the `tally` function if you use the formula `~y|x`.

Table 5.2: Table of conditional proportions in the Homogeneity testing scenario.

	Response Level 1	Response Level 2	Response Level 3	...	Response Level C	Totals
Group 1	$\rho_{11}$	$\rho_{12}$	$\rho_{13}$	...	$\rho_{1C}$	<b>1.0</b>
Group 2	$\rho_{21}$	$\rho_{22}$	$\rho_{23}$	...	$\rho_{2C}$	<b>1.0</b>
...	...	...	...	...	...	...
Group R	$\rho_{R1}$	$\rho_{R2}$	$\rho_{R3}$	...	$\rho_{RC}$	<b>1.0</b>
Totals	$\rho_{\bullet 1}$	$n_{\bullet 2}$	$\rho_{\bullet 3}$	...	$\rho_{\bullet C}$	<b>1.0</b>

In the Homogeneity situation, the null hypothesis is that the distributions are the same in all the  $R$  populations. This means that the null hypothesis is:

$$\mathbf{H}_0 : \rho_{11} = \rho_{21} = \dots = \rho_{R1} \text{ and } \rho_{12} = \rho_{22} = \dots = \rho_{R2} \text{ and } \rho_{13} = \rho_{23} = \dots = \rho_{R3} \text{ and } \dots \text{ and } \rho_{1C} = \rho_{2C} = \dots = \rho_{RC}.$$

If all the groups are the same, then they all have the same conditional proportions and we can more simply write the null hypothesis as:

$$\mathbf{H}_0 : (\rho_{r1}, \rho_{r2}, \dots, \rho_{rC}) = (\rho_1, \rho_2, \dots, \rho_C) \text{ for all } r.$$

In other words, the pattern of proportions across the columns are **the same for all the R groups**. The alternative is that there is some difference in the proportions of at least one response category for at least one group. In slightly more gentle and easier to reproduce words, equivalently, we can say:

- $\mathbf{H}_0$  : The population distributions of the responses for variable  $y$  are the same across the  $R$  groups.

The alternative hypothesis is then:

- $\mathbf{H}_A$  : The population distributions of the responses for variable  $y$  are NOT ALL the same across the  $R$  groups.

To make this concrete, consider what the proportions could look like if they satisfied the null hypothesis for the *Arthritis* example, as displayed in Figure 5.4.

Note that the proportions in the different response categories do not need to be the same just that the distribution needs to be the same across the groups. To make this clear, the null hypothesis does *not* require that all three response categories (*none*, *some*, *marked*) be equally likely. It assumes that whatever the distribution of proportions is across these three levels that there is no difference in that distribution between

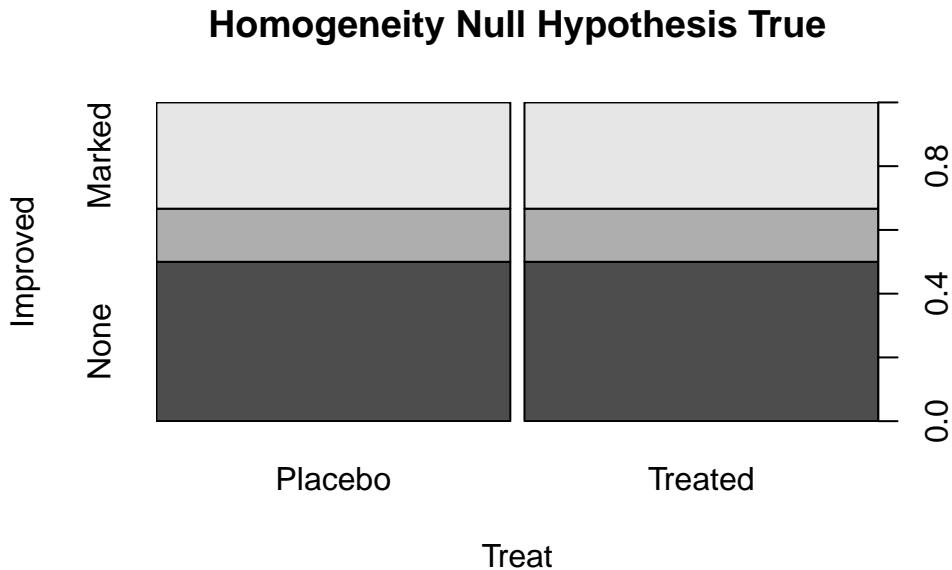


Figure 5.4: Plot of what the Arthritis proportions would look like if the null hypothesis had been true.

the explanatory variable (here treated/placebo) groups. Figure 5.4 shows an example of a situation where the null hypothesis is true and the distributions of responses across the groups look the same but the proportions for *none*, *some* and *marked* are not all equally likely. That situation satisfies the null hypothesis. Compare this plot to the one for the real data set in Figure 5.2. It looks like there might be some differences in the responses between the treated and placebo groups as that plot looks much different from this one, but we will need a test statistic and a p-value to fully address the evidence relative to the previous null hypothesis.

### 5.3 Independence Test Hypotheses

When we take a single random sample of size  $N$  and make a contingency table, our inferences relate to whether there is a relationship or *association* (that they are not independent) between the variables. This is related to whether the distributions of proportions match across rows in the table but is a more general question since we do not need to determine a variable to condition on that takes on the role of a predictor variable from the two variables of interest. In general, the hypotheses for an Independence test for variables  $x$  and  $y$  is:

- **$H_0$ : There is no relationship between  $x$  and  $y$  in the population.**
  - Or:  $H_0$ :  $x$  and  $y$  are independent in the population.
- **$H_A$ : There is a relationship between  $x$  and  $y$  in the population.**
  - Or:  $H_A$ :  $x$  and  $y$  are dependent in the population.

To illustrate a test of independence, consider an example involving data from a national random sample taken prior to the 2000 US elections from the data set `election` from the package `poLCA` (Linzer and Lewis. (2014), Linzer and Lewis (2011)). Each respondent's democratic-republican partisan identification was collected, provided in the `PARTY` variable for measurements on a seven-point scale from (1) *Strong Democrat*, (2) *Weak Democrat*, (3) *Independent-Democrat*, (4) *Independent-Independent*, (5) *Independent-Republican*, (6) *Weak Republican*, to (7) *Strong Republican*. The `VOTEF` variable that is created below will contain the candidate

that the participants voted for (the data set was originally coded with 1, 2, and 3 for the candidates and we replaced those levels with the candidate names). The contingency table shows some expected results, that individuals with strong party affiliations tend to vote for the party nominee with strong support for Gore in the democrats (PARTY = 1 and 2) and strong support for Bush in the republicans (PARTY = 6 and 7). As always, we want to support our explorations with statistical inferences, here with the potential to extend inferences to the overall population of voters. The inferences in an independence test are related to whether there is a relationship between the two variables in the population. A ***relationship*** between variables occurs when knowing the level of one variable for a person, say that they voted for Gore, informs the types of responses that you would expect for that person, here that they are likely affiliated with the Democratic Party. When there is no relationship (the null hypothesis here), knowing the level of one variable is not informative about the level of the other variable.

```
require(poLCA)
data(election) #2000 Survey
# Subset variables and remove missing values
election2 <- na.omit(election[,c("PARTY", "VOTE3")])
election2$VOTEF <- factor(election2$VOTE3)
levels(election2$VOTEF) <- c("Gore", "Bush", "Other")
levels(election2$VOTEF)

## [1] "Gore"  "Bush"  "Other"
electable <- tally(~PARTY+VOTEF, data=election2)
electable

##      VOTEF
## PARTY Gore Bush Other
##   1    238    6    2
##   2    151   18    1
##   3    113   31   13
##   4     37   37   11
##   5     21  124   12
##   6     20  121    2
##   7     3  189    1
```

The hypotheses for an Independence/Association Test here are:

- $H_0$ : There is no relationship between party affiliation and voting status in the population.
  - Or:  $H_0$ : Party affiliation and voting status are independent in the population.
- $H_A$ : There is a relationship between party affiliation and voting status in the population.
  - Or:  $H_A$ : Party affiliation and voting status are dependent in the population.

You could also write these hypotheses with the variables switched and that is also perfectly acceptable. Because these hypotheses are ambivalent about the choice of a variable as an “x” or a “y”, the summaries of results should be consistent with that idea. We should not calculate conditional proportions or make stacked bar charts since they imply a directional relationship from x to y (or results for y conditional on the levels of x) that might be hard to justify. Our summaries in these situations are the contingency table (`tally(~var1+var2, data=DATASETNAME)`) and a new graph called a ***mosaicplot*** (using the `mosaicplot` function).

Mosaic plots display a box for each cell count whose area corresponds to the proportion of the total data set that is in that cell ( $n_{rc}/N$ ). In some cases, the bars can be short or narrow if proportions of the total are small and the labels can be hard to read but the same bars or a single line exist for each category of the variables in all rows and columns. The mosaic plot makes it easy to identify the most common combination of categories. For example, in Figure 5.5 the *Gore* and *PARTY* = 1 (*Strong Democrat*) box in the top segment under column 1 of the plot has the largest area so is the highest proportion of the total. Similarly, the middle segment on the right for the *PARTY* category 7s corresponds to the *Bush* voters who were a 7 (*Strong Republican*). Knowing that the middle box in each column is for Bush voters is a little difficult as “Other”

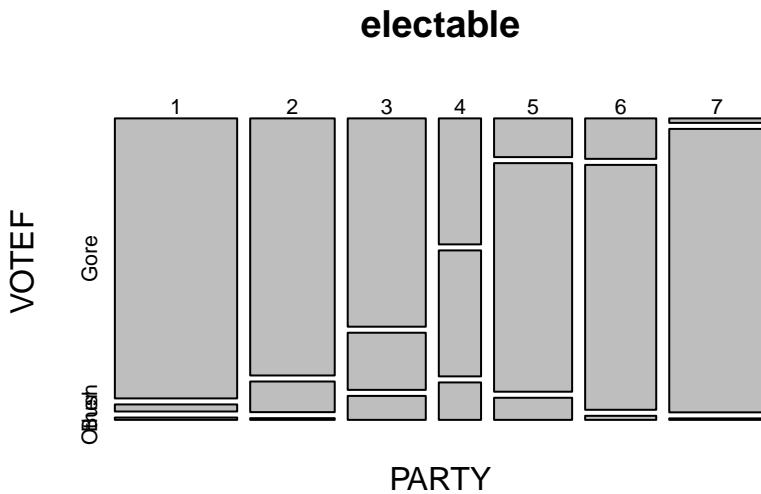


Figure 5.5: Mosaic plot of the 2000 election data comparing party affiliation and voting results.

and “Bush” overlap each other in the y-axis labeling but it is easy enough to sort out the story here if we have briefly explored the contingency table. We can also get information about the variable used to make the columns as the width of the columns is proportional to the number of subjects in each PARTY category in this plot. There were relatively few 4s (*Independent-Independent* responses) in total in the data set. Also, the *Other* category was the highest proportion of any vote-getter in the PARTY = 4 column but there were actually slightly more *Other* votes out of the total in the 3s (*Independent-Democrat*) party affiliation. Comparing the size of the 4s & *Other* segment with the 3s & *Other* segment, one should conclude that the 3s & *Other* segment is a slightly larger portion of the total data set. There is generally a gradient of decreasing/increasing voting rates for the two primary candidates across the party affiliations, but there are a few exceptions. For example, the proportion of *Gore* voters goes up slightly between the PARTY affiliations of 5s and 6s – as the voters become more strongly republican. To have evidence of a relationship, there just needs to be a pattern of variation across the plot of some sort but it does not need to follow such an easily described pattern, especially when the categorical variables do not contain natural ordering.

The mosaic plots are best made on the tables created by the `tally` function from a table that just contains the counts (**no totals**).

```
# Makes a mosaic plot where areas are related to the proportion of
# the total in the table
mosaicplot(electable)
```

In general, the results here are not too surprising as the respondents became more heavily republican, they voted for Bush and the same pattern occurs as you look at more democratic respondents. As the voters leaned towards being independent, the proportion voting for “Other” increased. So it certainly seems that there is some sort of relationship between party affiliation and voting status. As always, it is good to compare the observed results to what we would expect if the null hypothesis is true. Figure 5.6 assumes that the null hypothesis is true and shows the variation in the proportions in each category in the columns and variation in the proportions across the rows, but displays no relationship between PARTY and VOTEF. Essentially, the pattern down a column is the same for all the columns or vice-versa for the rows. The way to think of “no relationship” here would involve considering whether knowing the party level could help you predict the voting response and that is not the case in Figure 5.6 but was in certain places in Figure 5.5.

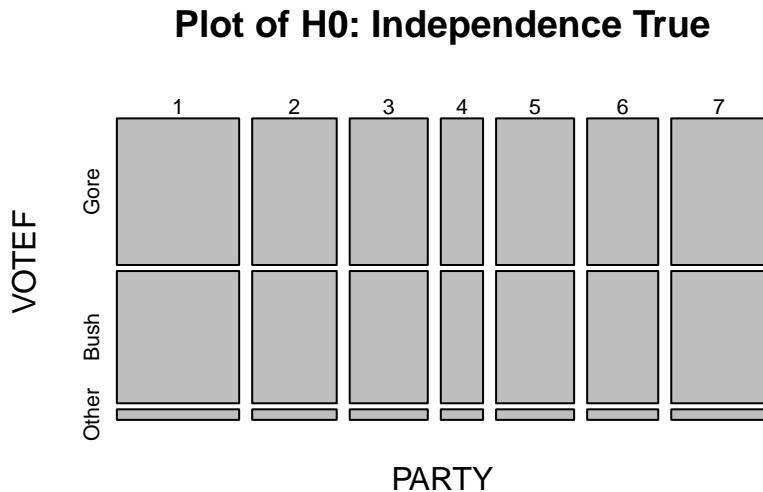


Figure 5.6: Mosaic plot of what the 2000 election data would look like if the null hypothesis of no relationship were true.

## 5.4 Models for R by C tables

This section is very short because we really do not use any “models” in this material. There are some complicated statistical models that can be employed in these situations, but they are beyond the scope of this book. What we do have in this situation is our original data summary in the form of a contingency table, graphs of the results like those seen above, a hypothesis test and p-value (presented below), and some post-test plots that we can use to understand the “source” of any evidence we found in the test.

## 5.5 Permutation tests for the $X^2$ statistic

In order to assess the evidence against our null hypotheses of no difference in distributions or no relationship between the variables, we need to define a test statistic and find its distribution under the null hypothesis. The test statistic used with both types of tests is called the  **$X^2$  statistic** (we want to call the statistic X-squared not Chi-squared). The statistic compares the observed counts in the contingency table to the ***expected counts*** under the null hypothesis, with large differences between what we observed and what we expect under the null leading to evidence against the null hypothesis. To help this statistic to follow a named parametric distribution and provide some insights into sources of interesting differences from the null hypothesis, we **standardize**<sup>3</sup> the difference between the observed and expected counts by the square-root of the expected count. The  **$X^2$  statistic** is based on the sum of squared standardized differences,

$$X^2 = \sum_{i=1}^{RC} \left( \frac{Observed_i - Expected_i}{\sqrt{Expected_i}} \right)^2,$$

---

<sup>3</sup>Standardizing involves dividing by the standard deviation of a quantity so it has a standard deviation 1 regardless of its original variability and that is what is happening here even though it doesn’t look like the standardization you are used to with continuous variables.

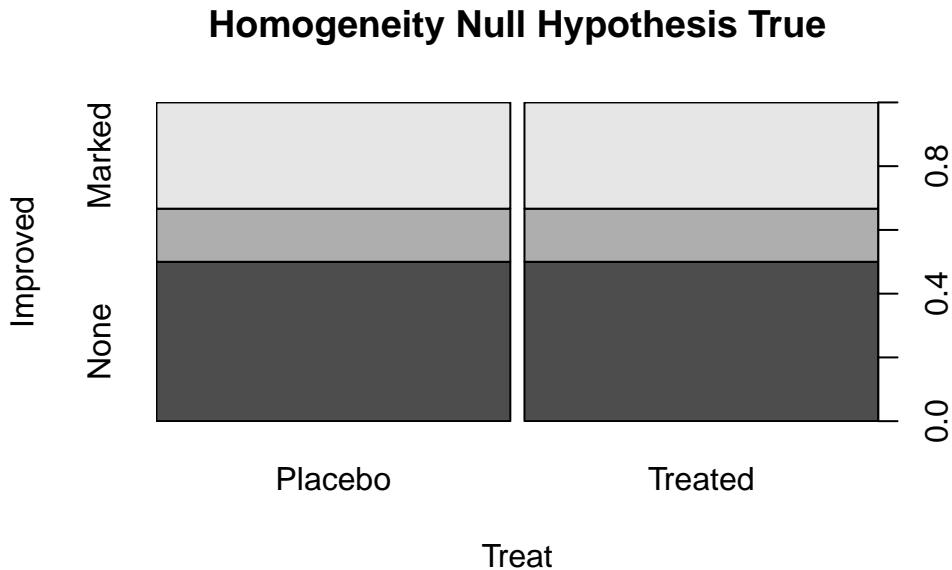


Figure 5.7: Stacked bar chart that could occur if the null hypothesis were true.

which is the sum over all ( $R$  times  $C$ ) cells in the contingency table of the square of the difference between observed and expected cell counts divided by the square root of the expected cell count. To calculate this test statistic, it is useful to start with a table of expected cell counts to go with our contingency table of observed counts. The expected cell counts are easiest to understand in the homogeneity situation but are calculated the same in either scenario.

The idea underlying finding the *expected cell counts* is to find how many observations we would expect in category  $c$  given the sample size in that group,  $n_{r\bullet}$ , if the null hypothesis is true. Under the null hypothesis across all  $R$  groups the conditional probabilities in each response category must be the same. Consider Figure 5.7 where, under the null hypothesis, the probability of *None*, *Some*, and *Marked* are the same in both treatment groups. Specifically we have  $\Pr(\text{None}) = 0.5$ ,  $\Pr(\text{Some}) = 0.167$ , and  $\Pr(\text{Marked}) = 0.333$ . With  $n_{\text{placebo}\bullet} = 43$  and  $\Pr(\text{None}) = 0.50$ , we would expect  $43 * 0.50 = 21.5$  subjects to be found in the *Placebo*, *None* combination if the null hypothesis were true. Similarly, with  $\Pr(\text{Some}) = 0.167$ , we would expect  $43 * 0.167 = 7.18$  in the *Placebo*, *Some* cell. And for the *Treated* group with  $n_{\text{Treated}\bullet} = 41$ , the expected count in the *Marked* improvement group would be  $41 * 0.333 = 13.65$ . Those conditional probabilities came from aggregating across the rows because, under the null, the row (*Treatment*) should not matter. So, the conditional probability was actually calculated as  $n_{\bullet c}/N = \text{total number of responses in category } c \text{ divided by the table total}$ . Since each expected cell count was a conditional probability times the number of observations in the row, we can re-write the expected cell count formula for row  $r$  and column  $c$  as:

$$\text{Expected cell count}_{rc} = \frac{(n_{r\bullet} * n_{\bullet c})}{N} = \frac{(\text{row } r \text{ total} * \text{column } c \text{ total})}{\text{table total}}.$$

Table 5.3 demonstrates the calculations of the expected cell counts using this formula for all 6 cells in the  $2 \times 3$  table.

Table 5.3: Demonstration of calculation of expected cell counts for Arthritis data.

	None	Some	Marked	Totals
Placebo	$\frac{n_{\text{Placebo}} * n_{\bullet\text{None}}}{N}$ = $\frac{43 * 42}{84}$ = <b>21.5</b>	$\frac{n_{\text{Placebo}} * n_{\bullet\text{Some}}}{N}$ = $\frac{43 * 14}{84}$ = <b>7.167</b>	$\frac{n_{\text{Placebo}} * n_{\bullet\text{Marked}}}{N}$ = $\frac{43 * 28}{84}$ = <b>14.33</b>	$n_{\text{Placebo}} = 43$
Treated	$\frac{n_{\text{Treated}} * n_{\bullet\text{None}}}{N}$ = $\frac{41 * 42}{84}$ = <b>20.5</b>	$\frac{n_{\text{Treated}} * n_{\bullet\text{Some}}}{N}$ = $\frac{41 * 14}{84}$ = <b>6.83</b>	$\frac{n_{\text{Treated}} * n_{\bullet\text{Marked}}}{N}$ = $\frac{41 * 28}{84}$ = <b>13.67</b>	$n_{\text{Treated}} = 41$
Totals	$n_{\bullet\text{None}} = 42$	$n_{\bullet\text{Some}} = 14$	$n_{\bullet\text{Marked}} = 28$	$N = 84$

Of course, using R can help us avoid tedium like this... The main engine for results in this chapter is the `chisq.test` function. It operates on a table of counts that has been produced **without row or column totals**. For example, `Arhtable` below contains just the observed cell counts. Applying the `chisq.test` function to `Arhtable` provides a variety of useful output. For the moment, we are just going to extract the information in the “expected” attribute of the results from running this function (using `chisq.test()$expected`). These are the expected cell counts which match the previous calculations except for some rounding in the hand-calculations.

```
Arhtable <- tally(~Treatment+Improved, data=Arthritis)
```

```
Arhtable
```

```
##           Improved
## Treatment None Some Marked
##   Placebo    29     7     7
##   Treated    13     7    21
chisq.test(Arhtable)$expected
```

```
##           Improved
## Treatment None   Some   Marked
##   Placebo 21.5 7.166667 14.33333
##   Treated 20.5 6.833333 13.66667
```

With the observed and expected cell counts in hand, we can turn our attention to calculating the test statistic. It is possible to lay out the “contributions” to the  $X^2$  statistic in a table format, allowing a simple way to finally calculate the statistic without losing any numbers. For **each cell** we need to find

$$(observed - expected / \sqrt{expected}),$$

**square them**, and then we need to add them **all** up. In the current example, there are 6 cells to add up ( $R = 2$  times  $C = 3$ ), shown in Table 5.4.

Table 5.4:  $X^2$  contributions for the Arthritis data.

	None	Some	Marked
Placebo	$\left(\frac{29-21.5}{\sqrt{21.5}}\right)^2 = 2.616$	$\left(\frac{7-7.167}{\sqrt{7.167}}\right)^2 = 0.004$	$\left(\frac{7-14.33}{\sqrt{14.33}}\right)^2 = 3.752$
Treated	$\left(\frac{13-20.5}{\sqrt{20.5}}\right)^2 = 2.744$	$\left(\frac{7-6.833}{\sqrt{6.833}}\right)^2 = 0.004$	$\left(\frac{21-13.67}{\sqrt{13.67}}\right)^2 = 3.935$

Finally, the  $X^2$  statistic here is the sum of these six results =  $2.616 + 0.004 + 3.752 + 2.744 + 0.004 + 3.935 = 13.055$

Our favorite function in this chapter, `chisq.test`, does not provide the contributions to the  $X^2$  statistic directly. It provides a related quantity called the

$$\text{standardized residual} = \left( \frac{\text{Observed}_i - \text{Expected}_i}{\sqrt{\text{Expected}_i}} \right),$$

which, when squared (in R, squaring is accomplished using `^2`), is the contribution of that particular cell to the  $X^2$  statistic that is displayed in Table 5.4.

```
(chisq.test(Arhtable)$residuals)^2
```

```
##          Improved
## Treatment      None      Some     Marked
##   Placebo 2.616279070 0.003875969 3.751937984
##   Treated 2.743902439 0.004065041 3.934959350
```

The most common error made in calculating the  $X^2$  statistic by hand involves having observed less than expected and then failing to make the  $X^2$  contribution positive for all cells (remember you are **squaring the entire quantity** in the parentheses and so the sign has to go positive!). In R, we can add up the cells using the `sum` function over the entire table of numbers:

```
sum((chisq.test(Arhtable)$residuals)^2)
```

```
## [1] 13.05502
```

Or we can let R do all this hard work for us and get straight to the good stuff:

```
chisq.test(Arhtable)
```

```
##
##  Pearson's Chi-squared test
##
## data: Arhtable
## X-squared = 13.055, df = 2, p-value = 0.001463
```

The `chisq.test` function reports a p-value by default. Before we discover how it got that result, we can rely on our permutation methods to obtain a distribution for the  $X^2$  statistic under the null hypothesis. As in Chapters 2 and 3, this will allow us to find a p-value while relaxing one of our assumptions<sup>4</sup>. In the One-WAY ANOVA in Chapter 3, we permuted the grouping variable relative to the responses, mimicking the null hypothesis that the groups are the same and so we can shuffle them around if the null is true. That same technique is useful here. If we randomly permute the grouping variable used to form the rows in the contingency table relative to the responses in the other variable and track the possibilities available for the  $X^2$  statistic under permutations, we can find the probability of getting a result as extreme as or more extreme than what we observed, our p-value. The observed statistic is the  $X^2$  calculated using the formula above. Like the  $F$ -statistic, it ends up that only results in the right tail of this distribution are desirable for finding evidence against the null hypothesis because all the values have to be positive. You can see this by observing that values of the  $X^2$  statistic close to 0 are generated when the observed values are close to the expected values and that sort of result should not be used to find evidence against the null. When the observed and expected values are “far apart”, then we should find evidence against the null. It is helpful to work through some examples to be able to understand how the  $X^2$  statistic “measures” differences between observed and expected.

To start, compare the previous observed  $X^2$  of 13.055 to the sort of results we obtain in a single permutation of the treated/placebo labels – Figure 5.8 (top left panel) shows a permuted data set that produced  $X^{2*} = 0.62$ .

---

<sup>4</sup>Here it allows us to relax a requirement that all the expected cell counts are larger than 5 for the parametric test (Section 5.6).

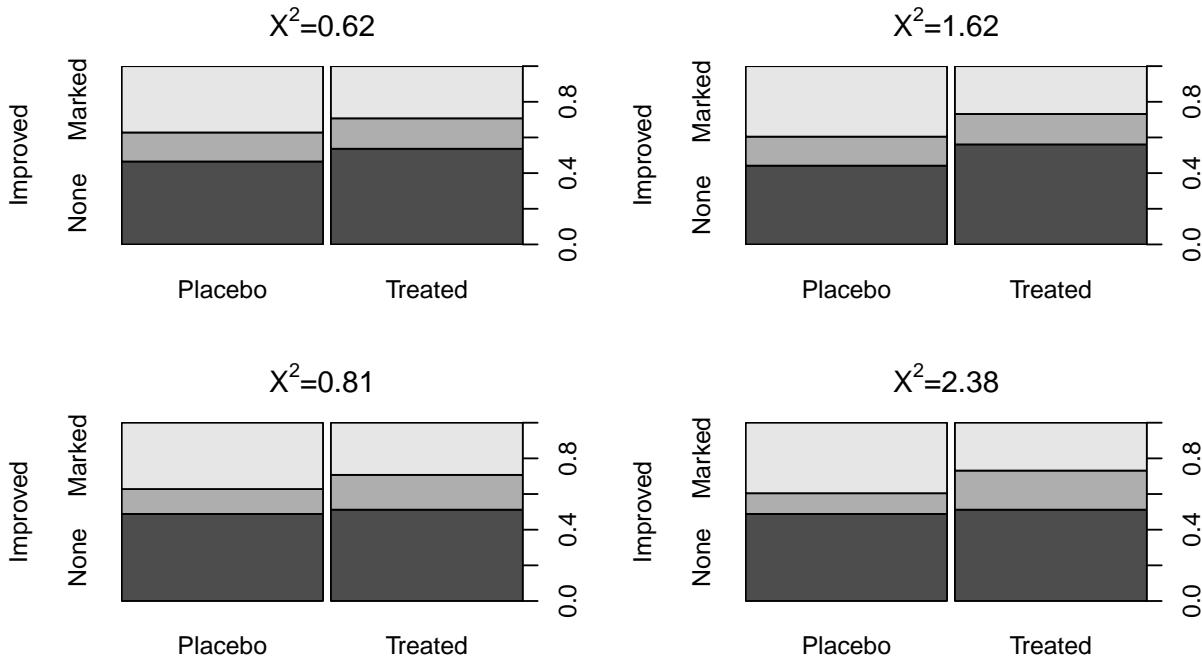


Figure 5.8: Stacked bar charts of four permuted Arthritis data sets that produced  $X^2$  between 0.62 and 2.38.

Visually, you can only see minimal differences between the treatment and placebo groups showing up in the stacked bar-chart. Three other permuted data sets are displayed in Figure 5.8 showing the variability in results in permutations but that none get close to showing the differences in the bars observed in the real data set.

```
Arthperm <- Arthritis
Arthperm$PermTreatment <- factor(shuffle(Arthperm$Treatment))

plot(Improved~PermTreatment, data=Arthperm,
     main="Stacked Bar Chart of Permuted Arthritis Data")

Arthpermtable <- tally(~PermTreatment+Improved, data=Arthperm, margins=F)
Arthpermtable
```

```
##          Improved
## PermTreatment None Some Marked
##      Placebo    20    7   16
##      Treated    22    7   12
chisq.test(Arthpermtable)

##
##  Pearson's Chi-squared test
##
##  data: Arthpermtable
##  X-squared = 0.6194, df = 2, p-value = 0.7337
```

To build the permutation-based null distribution for the  $X^2$  statistic, we need to collect up the test statistics ( $X^{2*}$ ) in many of these permuted results. The code is similar to permutation tests in Chapters 2 and 3 except that each permutation generates a new contingency table that is summarized and provided to `chisq.test` to analyze. We again extract the `$statistic`.

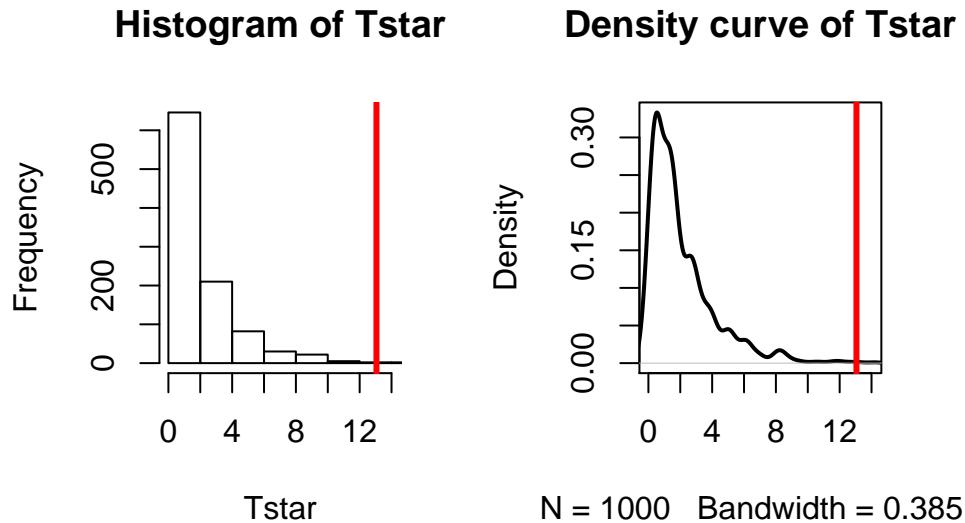


Figure 5.9: Permutation distribution for the  $X^2$  statistic for the Arthritis data with an observed  $X^2$  of 13.1 (bold, vertical line).

```

Tobs <- chisq.test(Arthtable)$statistic; Tobs

## X-squared
## 13.05502
par(mfrow=c(1,2))
B <- 1000
Tstar <- matrix(NA, nrow=B)
for (b in (1:B)){
  Tstar[b] <- chisq.test(tally(~shuffle(Treatment)+Improved,
                                data=Arthritis, margins=F))$statistic
}
pdata(Tstar, Tobs, lower.tail=F)

## X-squared
## 0.004
hist(Tstar, xlim=c(0,Tobs+1))
abline(v=Tobs, col="red", lwd=3)
plot(density(Tstar), main="Density curve of Tstar",
      xlim=c(0,Tobs+1), lwd=2)
abline(v=Tobs, col="red", lwd=3)

```

For an observed  $X^2$  statistic of 13.055, four out of 1,000 permutation results exceeded this value (pdata returned a value of 0.004). This suggests that our observed result is quite extreme relative to the null hypothesis and provides strong evidence against it.

**Validity conditions** for a permutation  $X^2$  test are:

1. Independence of observations.
2. Both variables are categorical.
3. Expected cell counts  $> 0$  (otherwise  $X^2$  is not defined).

For the permutation approach described here to provide valid inferences we need to be working with observations that are independent of one another. One way that a violation of independence can sometimes occur in this situation is when a single subject shows up in the table more than once. For example, if a single individual completes a survey more than once and those results are reported as if they came from  $N$  independent individuals. Be careful about this as it is really easy to make tables of poorly collected or non-independent observations and then consider them for these analyses. Poor data still lead to poor conclusions even if you have fancy new statistical tools to use!

## 5.6 Chi-square distribution for the $X^2$ statistic

When one additional assumption beyond the previous assumptions for the permutation test is met, it is possible to avoid permutations to find the distribution of the  $X^2$  statistic under the null hypothesis and get a p-value using what is called the ***Chi-square or  $\chi^2$ -distribution***. The name of our test statistic, X-squared, is meant to allude to the potential that this will follow a  $\chi^2$ -distribution in certain situations but may not do that all the time and we still can use the methods in Section 5.5. Along with the previous assumption regarding independence and all expected cell counts 0, we make a requirement that  $N$  (the total sample size) is “large enough” and this assumption is written in terms of the expected cell counts. If  $N$  is large, then all the expected cell counts should also be large because all those observations have to go somewhere. The problems for the  $\chi^2$ -distribution as an approximation to the distribution of the  $X^2$  statistic under the null hypothesis come when expected cell counts are below 5. And the smaller the expected cell counts become, the more problematic the  $\chi^2$ -distribution is as an approximation of the sampling distribution of the  $X^2$  statistic under the null hypothesis. **The standard rule of thumb is that all the expected cell counts need to exceed 5 for the parametric approach to be valid.** When this condition is violated, it is better to use the permutation approach. The `chisq.test` function will provide a bright red warning message to help you notice this. But it is good practice to always explore the expected cell counts using `chisq.test(...)$expected`.

```
chisq.test(Arthtable)$expected
```

```
##          Improved
## Treatment None     Some   Marked
##   Placebo 21.5 7.166667 14.33333
##   Treated 20.5 6.833333 13.66667
```

In the Arthritis data set, the sample size was sufficiently large for the  $\chi^2$ -distribution to provide an accurate p-value since the smallest expected cell count is 6.833 (it is larger than 5).

The  $\chi^2$ -distribution is a right-skewed distribution that starts at 0 as shown in Figure 5.10. Its shape changes as a function of its degrees of freedom. In the contingency table analyses, the ***degrees of freedom*** for the Chi-squared test are calculated as

$$DF = (\mathbf{R} - 1) * (\mathbf{C} - 1) = (\text{number of rows} - 1) * (\text{number of columns} - 1).$$

In the  $2 \times 3$  table above, the  $DF = (2 - 1) * (3 - 1) = 2$  leading to a Chi-square distribution with 2 *df* for the distribution of  $X^2$  under the null hypothesis. The p-value is based on the area to the right of the observed  $X^2$  value of 13.055 and the `pchisq` function provides that area as 0.00146. Note that this is very similar to the permutation result found previously for these data.

```
pchisq(13.055, df=2, lower.tail=F)
```

```
## [1] 0.001462658
```

We'll see more examples of the  $\chi^2$ -distributions in each of the examples that follow.

A small side note about sample sizes is warranted here. In contingency tables, especially those based on survey data, it is common to have large overall sample sizes ( $N$ ). With large sample sizes, it becomes easy to reject

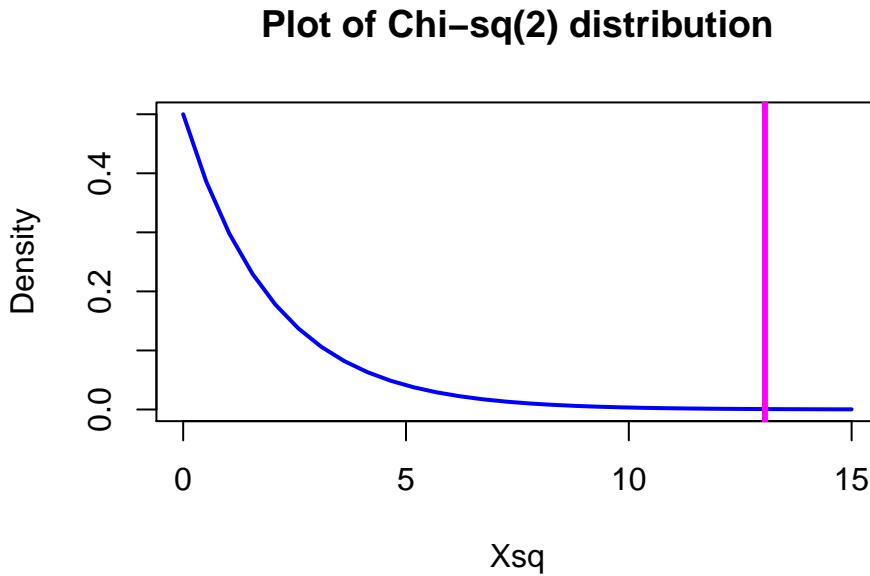


Figure 5.10:  $\chi^2$ -distribution with two degrees of freedom with 13.1 indicated with a vertical line.

the null the hypothesis, even when the “distance” from the null is relatively minor. By this we mean that it might be possible to reject the null hypothesis even if the observed proportions are a small practical distance from the situation described in the null. After obtaining a small p-value, we need to consider whether we have obtained ***practical significance*** (or maybe better described as ***practical importance***) to accompany our discussion of rejecting the null hypothesis. Whether a result is large enough to be of practical importance can only be judged by knowing something about the situation we are studying and by providing a good summary of our results to allow experts to assess the size and importance of the result. Statistical significance (what our tests address) is just a first step in any situation although many researchers are so happy to see small p-values that this is their last step.

If we revisit our observed results, re-plotted in Figure 5.11 since Figure 5.2 is many pages earlier, knowing that we have strong evidence against the null hypothesis of no difference between *Placebo* and *Treated* groups, what can we say about the effectiveness of the arthritis medication?

It seems that there is a real and important increase in the proportion of patients getting improvement (*Some* or *Marked*). If the differences “looked” smaller, even with a small p-value you might not recommend someone take the drug...

## 5.7 Examining residuals for the source of differences

Small p-values are generated by large  $X^2$  values. If we want to understand the source of a small p-value, we need to understand what made the test statistic large. To get a large  $X^2$  value, we either need many small contributions from lots of cells or a few large contributions. In most situations, there are just a few cells that show large deviations between the null hypothesis (expected cell counts) and what was observed (observed cell counts). It is possible to explore the “size” and direction of the differences between observed and expected counts to learn something about the behavior of the relationship between the variables, especially as it relates to evidence against the null hypothesis of no difference or no relationship. The ***standardized residual***,

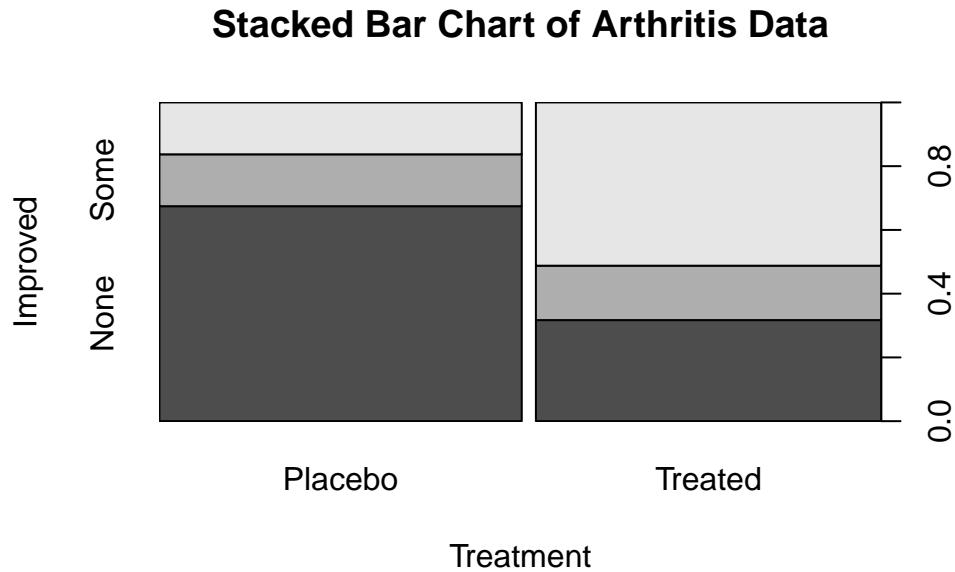


Figure 5.11: Stacked bar chart of the Arthritis data comparing *Treated* and *Placebo*.

$$\left( \frac{\text{Observed}_i - \text{Expected}_i}{\sqrt{\text{Expected}_i}} \right),$$

provides a measure of deviation of the observed from expected which retains the **direction of deviation** (whether observed was **more or less than expected** is interesting for interpretations) for each cell in the table. It is scaled much like a standard normal distribution providing a scale for “large” deviations for absolute values that are over 2 or 3. In other words, values with magnitude over 2 should be your focus in the standardized residuals, noting whether the observed counts were much more or less than expected. On the  $X^2$  scale, standardized residuals of 2 or more mean that the cells are contributing 4 or more units to the overall statistic, which is a pretty noticeable bump up in the size of the statistic. A few contributions at 4 or higher and you will likely end up with a small p-value.

There are two ways to explore standardized residuals. First, we can obtain them via the `chisq.test` and manually identify the “big ones”. Second, we can augment a mosaic plot of the table with the standardized results by turning on the `shade=T` option and have the plot help us find the big differences. This technique can be applied whether we are performing an Independence or Homogeneity test - both are evaluated with the same  $X^2$  statistic so the large standardized residuals are of interest in both situations. Both types of results are shown for the Arthritis data table:

```
##          Improved
## Treatment      None      Some     Marked
##   Placebo  1.61749160 -0.06225728 -1.93699199
##   Treated -1.65647289  0.06375767  1.98367320
chisq.test(Arthtable)$residuals
mosaicplot(Arthtable, shade=T)
```

In this data set, the standardized residuals are all less than 2 in magnitude so Figure 5.12 isn’t too helpful but this type of plot is in other examples. The largest contributions to the  $X^2$  statistic come from the *Placebo* and *Treated* groups in the *Marked* improvement cells. Those standardized residuals are -1.94 and 1.98 (both really close to 2), showing that *placebo* group had **noticeably fewer** *Marked* improvement

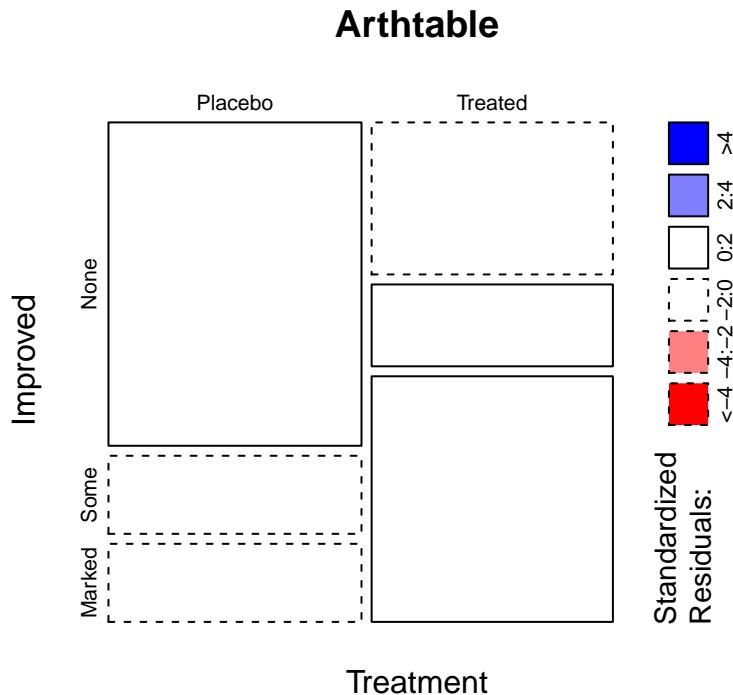


Figure 5.12: Mosaic plot of the Arthritis data with large standardized residuals indicated (actually, there were none that were indicated because all were less than 2).

results than expected and the *Treated* group had noticeably more *Marked* improvement responses than expected if the null hypothesis was true. Similarly but with smaller magnitudes, there were more *None* results than expected in the *Placebo* group and fewer *None* results than expected in the *Treated* group. The standardized residuals were very small in the two cells for the *Some* improvement category, showing that the treatment/placebo were similar in this response category and that the results were about what would be expected if the null hypothesis of no difference were true.

## 5.8 General Protocol for $X^2$ tests

In any contingency table situation, there is a general protocol to completing an analysis.

1. Identify the data collection method and whether the proper analysis is based on the Independence or Homogeneity hypotheses (Section 5.1).
2. Make a contingency table and get a general sense of response patterns. Pay attention to “small” counts, especially cells with 0 counts.
  - a. If there are many small count cells, consider combining categories on one or both variables to make a new variable with fewer categories that has larger counts per cell to have more robust inferences (See Section 5.10 for a related example).
3. Make the appropriate graphical display of results and generally describe the pattern of responses.
  - a. For Homogeneity, make a stacked bar-chart.
  - b. For Independence, make a mosaic plot.

- c. Consider a more general exploration using a tableplot if other variables were measured.
- 4. Conduct the 6+ steps of the appropriate type of hypothesis test.
  - a. Use permutations if any expected cell counts are below 5.
  - b. If all expected cell counts  $\geq 5$ , either permutation or parametric approaches are acceptable.
- 5. If sufficient evidence is found against the null to reject it, explore the standardized residuals for the “source” of the evidence against the null.
  - a. Tie the interpretation of the “large” standardized residuals and their direction (above or below expected under the null) back into the original data display. Work to find a story for the pattern of responses.

## 5.9 Political Party and Voting results: Complete Analysis

As introduced in Section 5.3, a national random sample of voters was obtained related to the 2000 Presidential Election with the party affiliations and voting results recorded for each subject. The data are available in `election` in the `polCA` package (Linzer and Lewis, 2011). It is always good to start with a bit of data exploration with a tableplot, displayed in Figure 5.13. Many of the lines of code here are just for making sure that R is treating the categorical variables that were coded numerically as categorical variables.

```
require(polCA)
data(election) #2000 Survey
require(tableplot)
election$VOTEF <- factor(election$VOTE3)
election$PARTY <- factor(election$PARTY)
election$EDUC <- factor(election$EDUC)
election$SEX <- factor(election$GENDER)
levels(election$VOTEF) <- c("Gore", "Bush", "Other")
tableplot(election, select=c(VOTEF, PARTY, EDUC, SEX))
```

In Figure 5.13, we can see many missing VOTEF responses but also some missingness in PARTY and EDUC (*Education*) status. While we don’t know too much about why people didn’t respond on the Vote question – they could have been unwilling to answer it or may not have voted. It looks like those subjects are have more of the lower education level responses (more green, brown, and blue) than in the responders to this question. There are many “middle” ratings (orange) in the party affiliation responses for the missing VOTEF responses, suggesting that independents were less likely to answer the question in the survey for whatever reason. We want to focus on those that did respond in VOTEF, so will again use `na.omit` to clean out any subjects with any missing responses on these four variables and remake this plot (Figure 5.14). The code also adds the `sort` option to the `tableplot` function call that provides an easy way to sort the data set based on other variables. It is interesting, for example, to sort the responses by *Education* level and explore the differences in other variables. These explorations are omitted here but easily available by changing the sorting column from 1 to `sort=3`.

```
election2 <- na.omit(election[,c("VOTEF", "PARTY", "EDUC", "SEX")])
tableplot(election2, select=c(VOTEF, PARTY, EDUC, SEX), sort=1)
```

Figure 5.14 shows us that there are clear differences in party affiliation based on voting for *Bush*, *Gore*, or *Other*. It is harder to see if there are differences in education level or sex based on the voting status in this plot, but, as noted above, sorting on these other variables can sometimes help to see other relationships between variables.

Focusing on the party affiliation and voting results, the appropriate analysis is with an Independence test because a single random sample was obtained from the population. The total sample size for the complete responses was  $N = 1,149$  (out of the original 1,785 subjects). Because this is an Independence test, the

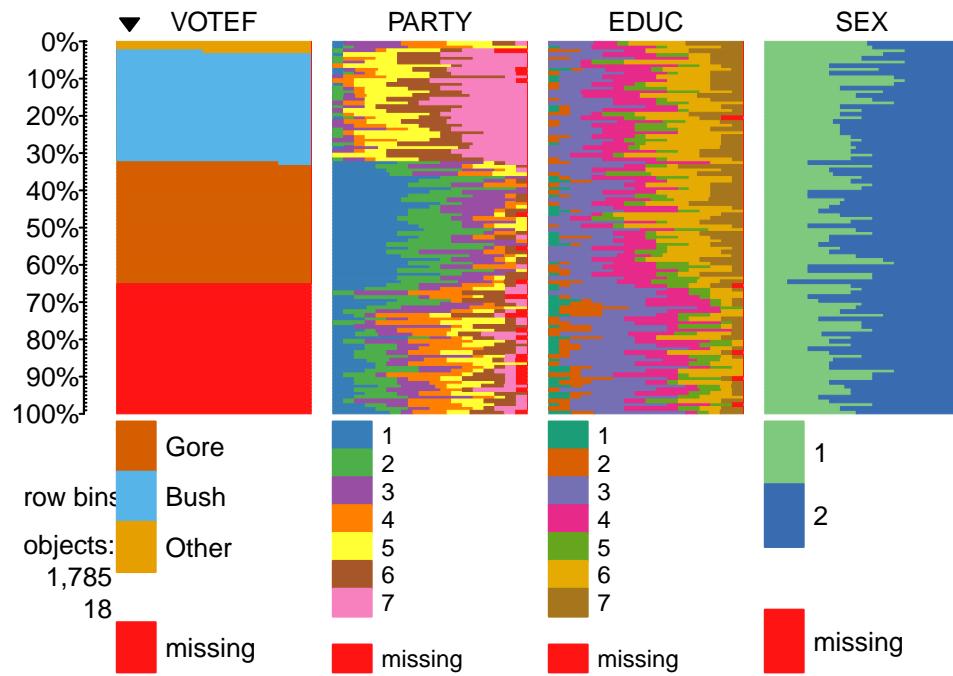


Figure 5.13: Tableplot of vote, party affiliation, education, and sex from election survey data. Note that missing observations are present in three of four variables. Education is coded from 1 to 7 with higher values related to higher educational attainment. Sex code 1 is for male and 2 is for female.

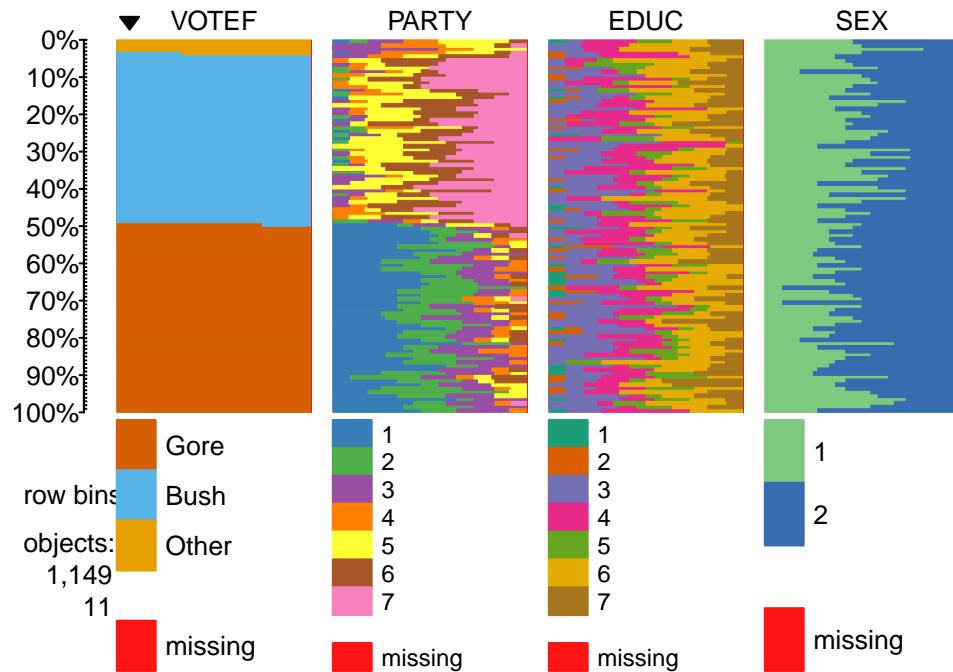


Figure 5.14: Plot of election data with subjects without any missing responses

mosaic plot is the appropriate display of the results, which was provided in Figure 5.5. We will perform this test at the 1% significance level.

```
electable <- tally(~PARTY+VOTEF, data=election2)
electable
```

```
##      VOTEF
## PARTY Gore Bush Other
##   1    238     6     2
##   2    151    18     1
##   3    113    31    13
##   4     37    36    11
##   5     21   124    12
##   6     20   121     2
##   7     3   188     1
```

There is a potential for bias in some polls because of the methods used to find and contact people. As US residents have transitioned from land-lines to cell phones, the early adopting cell phone users were often excluded from political polling. These policies are being reconsidered to adapt to the decline in residential phone lines and most polling organizations now include cell phone numbers in their list of potential respondents. This study may have some bias regarding who was considered as part of the population of interest and who was actually found that was willing to respond to their questions. We don't have much information here but biases arising from unobtainable members of populations are a potential issue in many studies, especially when questions tend toward more sensitive topics. We can make inferences here to people that were willing to respond to the request to answer the survey but should be cautious in extending it to all Americans or even voters in the year 2000. When we say "population" below, this nuanced discussion is what we mean. Because the political party is not randomly assigned to the subjects, we cannot make causal inferences for political affiliation causing different voting patterns<sup>5</sup>.

### 1. Hypotheses:

- $H_0$ : There is no relationship between the party affiliation (7 levels) and voting results (*Bush*, *Gore*, *Other*) in the population.
- $H_A$ : There is a relationship between the party affiliation (7 levels) and voting results (*Bush*, *Gore*, *Other*) in the population.

### 2. Validity conditions:

- Independence:
  - This assumption is presumed to be met since each subject is measured only once in the table. No other information suggests a potential issue since a random sample was taken from presumably a large national population.
- All expected cell counts larger than 5 to use the parametric  $\chi^2$ -distribution to find p-values:
  - We need to generate a table of expected cell counts to be able to check this condition:

```
chisq.test(electable)$expected
```

```
## Warning in chisq.test(electable): Chi-squared approximation may be
## incorrect

##      VOTEF
## PARTY      Gore      Bush     Other
##   1 124.81984 112.18799 8.992167
##   2  86.25762  77.52829 6.214099
##   3  79.66144  71.59965 5.738903
##   4  42.62141  38.30809 3.070496
```

---

<sup>5</sup>Independence tests can't be causal by their construction. Homogeneity tests could be causal or just associational, depending on how the subjects ended up in the groups.

### Plot of Chi-sq(12) distribution

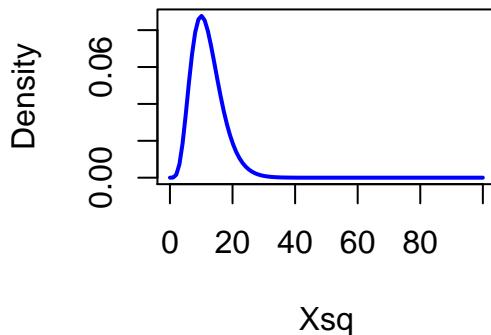


Figure 5.15: Plot of  $\chi^2$ -distribution with 12 degrees of freedom.

```
##      5  79.66144  71.59965  5.738903
##      6  72.55788  65.21497  5.227154
##      7  97.42037  87.56136  7.018277
```

- When we request the expected cell counts, R tries to help us with a warning message if the expected cell counts might be small, as in this situation.
- There is one expected cell count below 5 for *Party* = 4 who voted *Other* with an expected cell count of 3.102, so the condition is violated and the permutation approach should be used to obtain more trustworthy p-values. The conditions are met for performing a permutation test.

#### 3. Calculate the test statistic:

- This is best performed by the `chisq.test` function since there are 21 cells and many potential places for a calculation error if performed by hand.

```
chisq.test(electable)
```

```
##
## Pearson's Chi-squared test
##
## data: electable
## X-squared = 762.81, df = 12, p-value < 2.2e-16
```

- The observed  $X^2$  statistic is 762.81.

#### 4. Find the p-value:

- The parametric p-value is  $<2.2\text{e-}16$  from the R output which would be reported as  $<0.0001$ . This was based on a  $\chi^2$ -distribution with  $(7 - 1) * (3 - 1) = 12$  degrees of freedom displayed in Figure 5.15. Note that the observed test statistic of 762.81 was off the plot to the right which reflects how little area is to the right of that value in the distribution.
- If you want to repeat this calculation directly you get a similarly tiny value that R reports as  $1.5\text{e-}155$ . Again, reporting less than 0.0001 is just fine.

```
pchisq(762.81, df=12, lower.tail=F)
```

```
## [1] 1.553744e-155
```

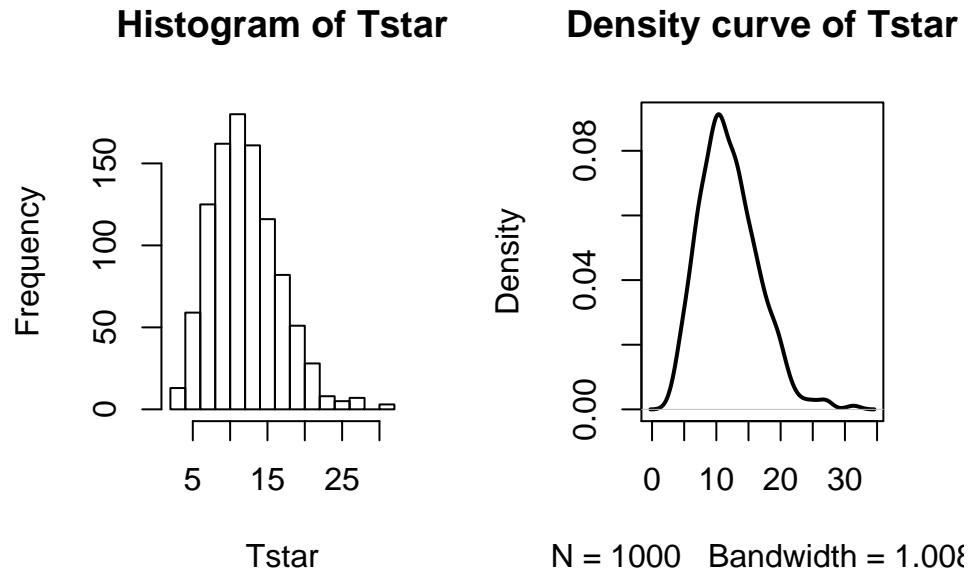


Figure 5.16: Permutation distribution of  $X^2$  for the election data. Observed value of 763 not displayed.

- But since the expected cell count condition is violated, we should use permutations as implemented in the following code to provide a trustworthy p-value:

```
Tobs <- chisq.test(electable)$statistic; Tobs

## X-squared
## 762.8095
par(mfrow=c(1,2))
B <- 1000
Tstar <- matrix(NA, nrow=B)
for (b in 1:B){
  Tstar[b] <- chisq.test(tally(~shuffle(PARTY)+VOTEF, data=election2,
                                margins=F))$statistic
}
pdata(Tstar, Tobs, lower.tail=F)

## X-squared
##          0
hist(Tstar)
abline(v=Tobs, col="red", lwd=3)
plot(density(Tstar), main="Density curve of Tstar", lwd=2)
abline(v=Tobs, col="red", lwd=3)
```

- The last results tells us that there were no permuted data sets that produced larger  $X^2$ 's than the observed  $X^2$  in 1,000 permutations, so we report that the **p-value was less than 0.001** using the permutation approach. The permutation distribution in Figure 5.16 contains no results over 40, so the observed configuration was really far from the null hypothesis of no relationship between party status and voting.

##### 5. Make a decision:

- With a p-value less than 0.001, we can say that there is almost no chance of observing a configuration

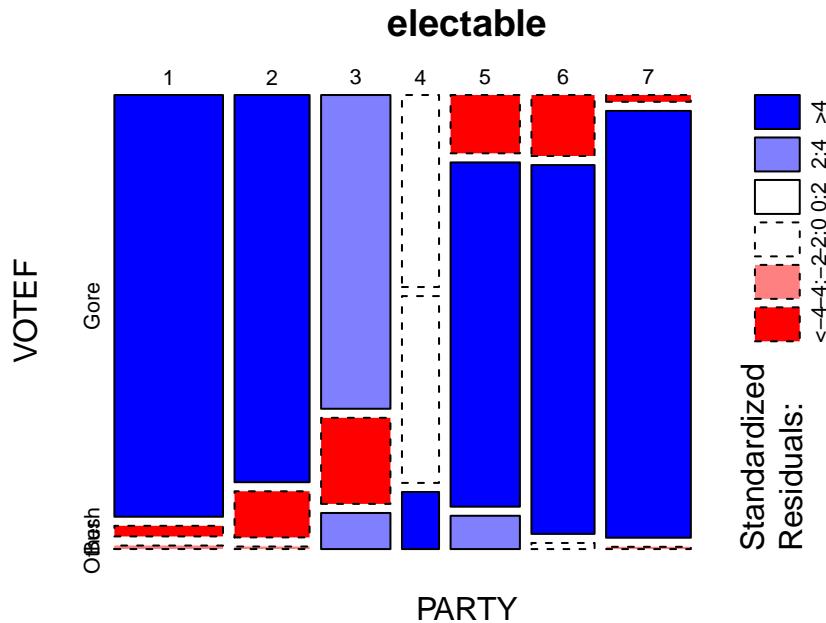


Figure 5.17: Mosaic plot with shading based on standardized residuals for the election

like ours or more extreme if the null hypothesis is true. So we should reject the null hypothesis.

#### 6. Write a conclusion:

- There is enough evidence (at 1% significance level) to conclude that there is a relationship between party affiliation and voting results in the population. The results are not causal since no random assignment was present but they do apply to the population voters in the 2000 election that were able to be contacted by those running the poll and who would be willing to answer all the questions and voted.

We can add a little more refinement to the results by exploring the standardized residuals. The numerical results are obtained using:

```
chisq.test(electable)$residuals #(Obs - expected)/sqrt(expected)
```

```
##      VOTEF
## PARTY    Gore     Bush     Other
##   1 10.1304439 -10.0254117 -2.3317373
##   2  6.9709179 -6.7607252 -2.0916557
##   3  3.7352759 -4.7980730  3.0310127
##   4 -0.8610559 -0.3729136  4.5252413
##   5 -6.5724708  6.1926811  2.6135809
##   6 -6.1701472  6.9078679 -1.4115200
##   7 -9.5662296 10.7335798 -2.2717310
```

And visually using:

```
#Adds information on the size of the residuals
mosaicplot(electable, shade=T)
```

In this example, the standardized residuals show some clear sources of the differences from the results expected if there were no relationship present. The largest contributions are found in the highest democrat category (PARTY = 1) where the standardized residual for *Gore* is 10.13 and for *Bush* is -10.03, showing much higher than expected (under  $H_0$ ) counts for *Gore* voters and much lower than expected (under  $H_0$ ) for *Bush*. Similar results in the opposite direction are found in the strong republicans (PARTY = 7). Note how the brightest

shade of blue in Figure 5.17 shows up for much higher than expected results and the brighter red for results in the other direction, where observed counts were much lower than expected. When there are many large standardized residuals, it is OK to focus on the largest results but remember that some of the intermediate deviations, or lack thereof, could also be interesting. For example, the Gore voters from PARTY = 3 had a standardized residual of 3.75 but the PARTY = 5 voters for Bush had a standardized residual of 6.17. So maybe Gore didn't have as strong of support from his center-leaning supporters as Bush was able to obtain from the same voters on the other side of the middle? A political scientist would easily obtain many more (useful) theories, especially once they understood the additional information provided by exploring the standardized residuals.

## 5.10 Is cheating and lying related in students?

A study of student behavior was performed at a university with a survey of  $N = 319$  undergraduate students (`cheating` data set from the `polCA` package originally published by Dayton (1998)). They were asked to answer four questions about their various academic frauds that involved cheating and lying. Specifically, they were asked if they had ever lied to avoid taking an exam (`LIEEXAM` with 1 for no and 2 for yes), if they had lied to avoid handing in a term paper on time (`LIEPAPER` with 2 for yes), if they had purchased a term paper to hand in as their own or obtained a copy of an exam prior to taking the exam (`FRAUD` with 2 for yes), and if they had copied answers during an exam from someone near them (`COPYEXAM` with 2 for yes). Additionally, their GPAs were obtained and put into categories: (<2.99, 3.0 to 3.25, 3.26 to 3.50, 3.51 to 3.75, and 3.76 to 4.0). These categories were coded from 1 to 5, respectively. Again, the code starts with making sure the variables are treated categorically by applying the `factor` function.

```
data(cheating) #Survey of students
cheating$LIEEXAM <- factor(cheating$LIEEXAM)
cheating$LIEPAPER <- factor(cheating$LIEPAPER)
cheating$FRAUD <- factor(cheating$FRAUD)
cheating$COPYEXAM <- factor(cheating$COPYEXAM)
cheating$GPA <- factor(cheating$GPA)
par(mfrow=c(1,1))
require(tabplot)
tableplot(cheating, sort=GPA)
```

We can explore some interesting questions about the relationships between these variables. The `tableplot` in Figure 5.18 again helps us to get a general idea of the data set and to assess some complicated aspects of the relationships between variables. For example, the rates of different unethical behaviors seem to decrease with higher GPA students (but do not completely disappear!). This data set also has a few missing GPAs that we would want to carefully consider – which sorts of students might not be willing to reveal their GPAs? It ends up that these students did not *admit* to any of the unethical behaviors... Note that we used the `sort=GPA` option in the `tableplot` function to sort the responses based on GPA to see how GPA might explain patterns of unethical behavior.

While the relationship between GPA and presence/absence of the different behaviors is of interest, we want to explore the types of behaviors. It is possible to group the lying behaviors as being a different type of unethical behavior than obtaining an exam prior to taking it, buying a paper, or copying someone else's answers. We want to explore whether there is some sort of relationship between the lying and copying behaviors – are those that engage in one type of behavior more likely to do the other? Or are they independent of each other? This is a hard story to elicit from the previous plot because there are so many variables involved.

To simplify the results, combining the two groups of variables into the four possible combinations on each has the potential to simplify the results – or at least allow exploration of additional research questions. In the table plot in Figure 5.19, you can see the four categories for each, starting with no bad behavior of either type (which is fortunately the most popular response on both variables!). For each variable, there are students who admitted to one of the two violations and some that did both. The `liar` variable has categories of *None*, *ExamLie*, *PaperLie*, and *LieBoth*. The `copier` variable has categories of *None*, *PaperCheat*, *ExamCheat*,

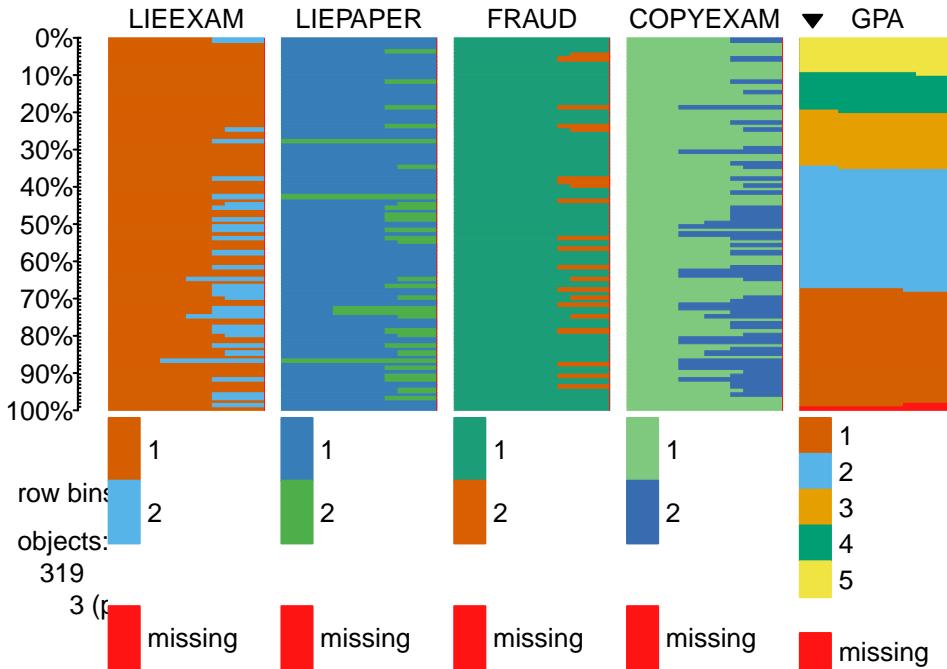


Figure 5.18: Tableplot of initial cheating and lying data set.

and *PaperExamCheat* (for doing both). The last category for copier (plotted as a peach(?) color) seems to mostly occur at the top of the plot which is where the students who had lied to get out of things reside, so maybe there is a relationship between those two types of behaviors? On the other hand, for the students who have never lied, quite a few had cheated on exams. The contingency table can help us dig further into the hypotheses related to the Chi-square test of Independence that is appropriate in this situation.

```
cheating$liar <- interaction(cheating$LIEEXAM, cheating$LIEPAPER)
levels(cheating$liar) <- c("None", "ExamLie", "PaperLie", "LieBoth")

cheating$copier <- interaction(cheating$FRAUD, cheating$COPYEXAM)
levels(cheating$copier) <- c("None", "PaperCheat", "ExamCheat", "PaperExamCheat")

cheatlietable <- tally(~liar+copier, data=cheating)
cheatlietable

##          copier
##   liar      None PaperCheat ExamCheat PaperExamCheat
##   None     207       7      46        5
##   ExamLie    10       1       3        2
##   PaperLie   13       1       4        2
##   LieBoth    11       1       4        2
tableplot(cheating, sort=liar, select=c(liar,copier))
```

Unfortunately, there were very few responses in some combinations of categories even with  $N = 319$ . For example, there was only one response each in the combinations for students that copied on papers and lied to get out of exams, papers, and both. Some other categories were pretty small as well in the groups that only had one behavior present. To get a higher number of counts in the combinations, we combined the single behavior only levels into “either” categories and left the *none* and *both* categories for each variable. This creates two new variables called *liar2* and *copier2* (tableplot in Figure 5.20). The code to create these variables and make the plot is below.

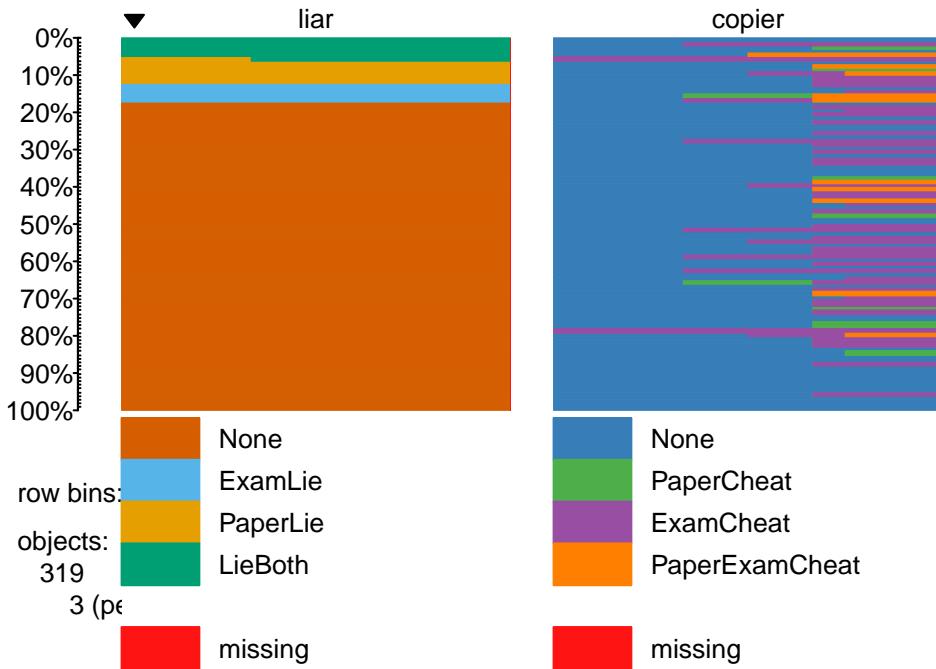


Figure 5.19: Tableplot of new variables liar and copier that allow exploration of relationships between different types of lying and cheating behaviors.

```
#Collapse the middle categories of each variable
cheating$liar2 <- cheating$liar
levels(cheating$liar2) <- c("None", "ExamorPaper", "ExamorPaper", "LieBoth")
cheating$copier2 <- cheating$copier
levels(cheating$copier2) <- c("None", "ExamorPaper", "ExamorPaper", "CopyBoth")
tableplot(cheating, sort=liar2, select=c(liar2,copier2))
cheatlietatable <- tally(~liar2+copier2, data=cheating)
cheatlietatable
```

	copier2	None	ExamorPaper	CopyBoth
## liar2				
## None	207	53	5	
## ExamorPaper	23	9	4	
## LieBoth	11	5	2	

This  $3 \times 3$  table is more manageable and has few really small cells so we will proceed with the 6+ steps of hypothesis testing applied to these data using the Independence testing methods (again a single sample was taken from the population):

### 1. Hypotheses:

- $H_0$ : Lying and copying behavior are independent in the population of students at this university.
- $H_A$ : Lying and copying behavior are dependent in the population of students at this university.

### 2. Validity conditions:

- Independence:
  - This assumption is presumed to be met since each subject is measured only once in the table. No other information suggests a potential issue but we don't have much information on how these subjects were obtained.

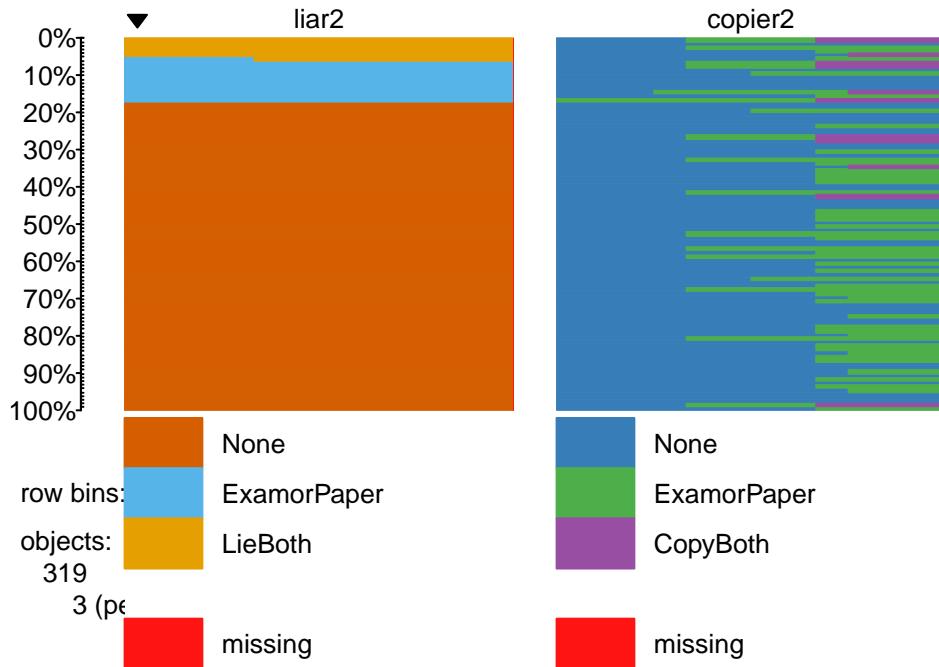


Figure 5.20: Tableplot of lying and copying variables after combining categories.

- All expected cell counts larger than 5 (required to use  $\chi^2$ -distribution to find p-values):

- We need to generate a table of expected cell counts to check this condition:

```
chisq.test(cheatlietable)$expected
```

```
##          copier2
## liar2      None ExamorPaper CopyBoth
##   None    200.20376  55.658307 9.1379310
##   ExamorPaper 27.19749   7.561129 1.2413793
##   LieBoth    13.59875   3.780564 0.6206897
```

- When we request the expected cell counts, there is a warning message (not shown).
- There are three expected cell counts below 5, so the condition is violated and a permutation approach should be used to obtain more trustworthy p-values.

### 3. Calculate the test statistic:

- Use `chisq.test` although this table is small enough to do by hand if you want the practice – see if you can find a similar answer to what the function provides:

```
chisq.test(cheatlietable)
```

```
##
##  Pearson's Chi-squared test
##
## data: cheatlietable
## X-squared = 13.238, df = 4, p-value = 0.01017
```

- The  $X^2$  statistic is 13.24.

### 4. Find the p-value:

- The parametric p-value is 0.0102 from the R output. This was based on a  $\chi^2$ -distribution with  $(3 - 1) * (3 - 1) = 4$  degrees of freedom that is displayed in Figure 5.21. Remember that this

### Plot of Chi-sq(4) distribution

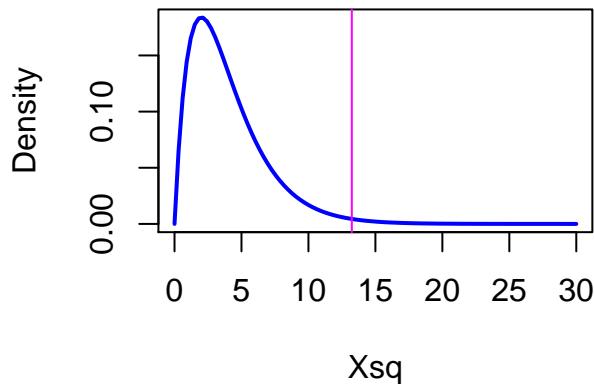


Figure 5.21: Plot of  $\chi^2$ -distribution with 4 degrees of freedom.

isn't quite the right distribution for the test statistic since our expected cell count condition was violated.

- If you want to repeat the p-value calculation directly:

```
pchisq(13.2384, df=4, lower.tail=F)
```

```
## [1] 0.01016781
```

- But since the expected cell condition is violated again, we should use permutations as implemented in the following code with the number of permutations increased to 10,000 to help get a better estimate of the p-value since it is possibly close to 0.05:

```
Tobs <- chisq.test(tally(~liar2+copier2, data=cheating))$statistic
Tobs
```

```
## X-squared
## 13.23844
par(mfrow=c(1,2))
B <- 10000 # Now performing 10,000 permutations
Tstar <- matrix(NA,nrow=B)
for (b in 1:B){
  Tstar[b] <- chisq.test(tally(~shuffle(liar2)+copier2,
                                data=cheating))$statistic
}
pdata(Tstar, Tobs, lower.tail=F)
```

```
## X-squared
##      0.017
hist(Tstar)
abline(v=Tobs, col="red", lwd=3)
plot(density(Tstar), main="Density curve of Tstar", lwd=2)
abline(v=Tobs, col="red", lwd=3)
```

- There were 170 of  $B=10,000$  permuted data sets that produced larger  $X^2$ 's than the observed as displayed in Figure 5.22, so we report that the p-value was 0.017 using the permutation approach, which was slightly larger than the result provided by the parametric method.

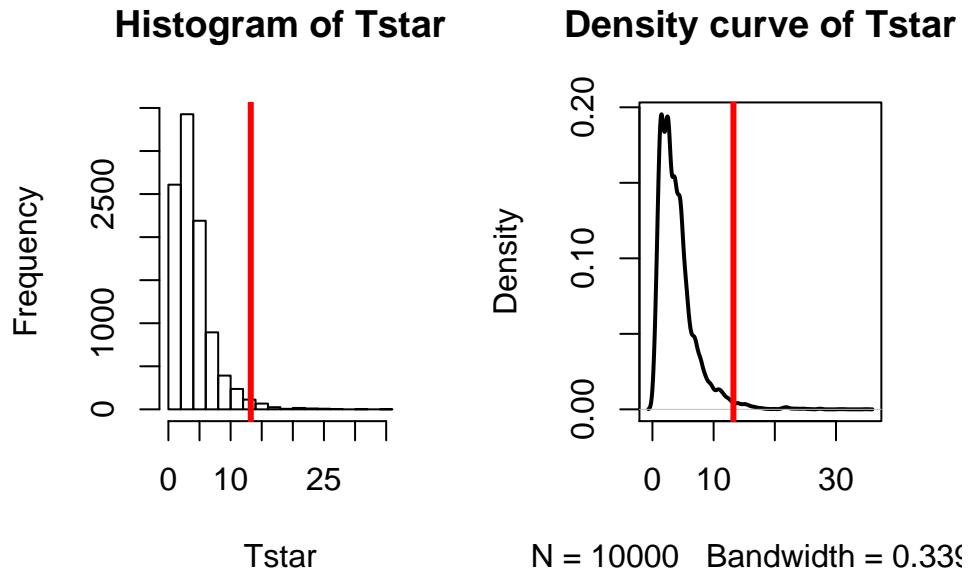


Figure 5.22: Plot of permutation distributions for cheat/lie results with observed value of 13.24 (bold, vertical line).

#### 5. Make a decision:

- With a p-value of 0.017, we can say that there is a 1.7% chance of observing a configuration like ours or more extreme if the null hypothesis is true. So we should probably reject the null hypothesis although your standards for evidence might differ – this is moderate to strong but not very strong evidence against the null hypothesis.

#### 6. Write a conclusion:

- If we reject the null hypothesis, there is enough evidence to conclude that there is a relationship between lying and copying behavior in the population of students. There is no causal inference possible here since neither variable was randomly assigned (really neither is explanatory or response here either) but we can extend the inferences to the population of students that these were selected from that would be willing to reveal their GPA (see initial discussion related to some differences in students that wouldn't answer that question).

The standardized residuals can help us more fully understand this result – the mosaic plot only had one cell shaded and so wasn't needed here.

```
chisq.test(cheatlietable)$residuals
```

```
##          copier2
## liar2      None ExamorPaper CopyBoth
##   None     0.4803220 -0.3563200 -1.3688609
## ExamorPaper -0.8048695  0.5232734  2.4759378
## LieBoth    -0.7047165  0.6271633  1.7507524
```

There is really one large standardized residual for the *ExamorPaper* liars and the *CopyBoth* copiers, with a much larger observed value than expected of 2.48. The only other medium-sized standardized residuals came from the *CopyBoth* copiers column with fewer than expected students in the *None* category and more than expected in the *LieBoth* type of lying category. So we are seeing more than expected that lied somehow and copied – we can say this suggests that the people who lie tend to copy too!

## 5.11 Analyzing a stratified random sample of California schools

In recent decades, there has been a push for quantification of school performance and tying financial punishment and rewards to growth in these metrics. One example is the API (Academic Performance Index) in California that is based mainly on student scores on standardized tests. It ranges between 200 and 1000 and year to year changes are of interest to assess “performance” of schools – calculated as one year minus the previous year (negative “growth” is also possible!). Suppose that a researcher is interested in whether the growth metric might differ between different levels of schools. Maybe it is easier or harder for elementary, middle, or high schools to attain growth. The researcher has a list of most of the schools in the state of each level that are using a data-base that the researcher has access to. In order to assess this question, the researcher takes a stratified random sample<sup>6</sup>, selecting  $n_{\text{elementary}} = 100$  schools from the population of 4421 elementary schools,  $n_{\text{middle}} = 50$  from the population of 1018 middle schools, and  $n_{\text{high}} = 50$  from the population of 755 high schools. These data are available in the `survey` package (Lumley, 2017) and the `api` and `apistrat` data sets. The `growth` (change!) in API scores for the schools between 1999 and 2000 (taken as the year 2000 score minus 1999 score) is used as the response variable. The boxplot and beanplot of the growth scores are displayed in Figure 5.23. They suggest some differences in the growth rates among the different levels. There are also a few schools flagged as being possible outliers.

```
require(survey)
data(api)
tally(~stype, data=apipop) #Population counts

## stype
##   E   H   M
## 4421 755 1018
tally(~stype, data=apistrat) #Sample counts

## stype
##   E   H   M
## 100 50 50
par(mfrow=c(1,2))
boxplot(growth~stype, data=apistrat, ylab="Growth", ylim=c(-55,160))
beanplot(growth~stype, data=apistrat, log="", col="beige",
         method="jitter", ylim=c(-55,160))
```

The One-Way ANOVA  $F$ -test, provided below, suggests evidence of some difference in the true mean growth scores among the different types of schools ( $F(2, 197) = 23.56$ ,  $p$ -value  $< 0.0001$ ). But the residuals from this model displayed in the QQ-Plot in Figure 5.24 contain a slightly long right tail, suggesting a right skewed distribution for the residuals. In a high-stakes situation such as this, reporting results with violations of the assumptions probably would not be desirable, so another approach is needed. The permutation methods would be justified here but there is another “simpler” option available using our Chi-square analysis methods.

```
m1<-lm(growth~stype, data=apistrat)
require(car)
Anova(m1)

## Anova Table (Type II tests)
##
## Response: growth
##           Sum Sq Df F value    Pr(>F)
## stype     30370  2 23.563 6.685e-10
## Residuals 126957 197
## plot(m1, which=2)
```

One way to get around the normality assumption is to use a method that does not assume the responses follow

---

<sup>6</sup>A stratified random sample involves taking a simpler random sample from each group or strata of the population. It is useful to make sure that each group is represented at a chosen level (for example the sample proportion of the total size). If a simple random sample of all schools had been taken, it is possible that a level could have no schools selected.

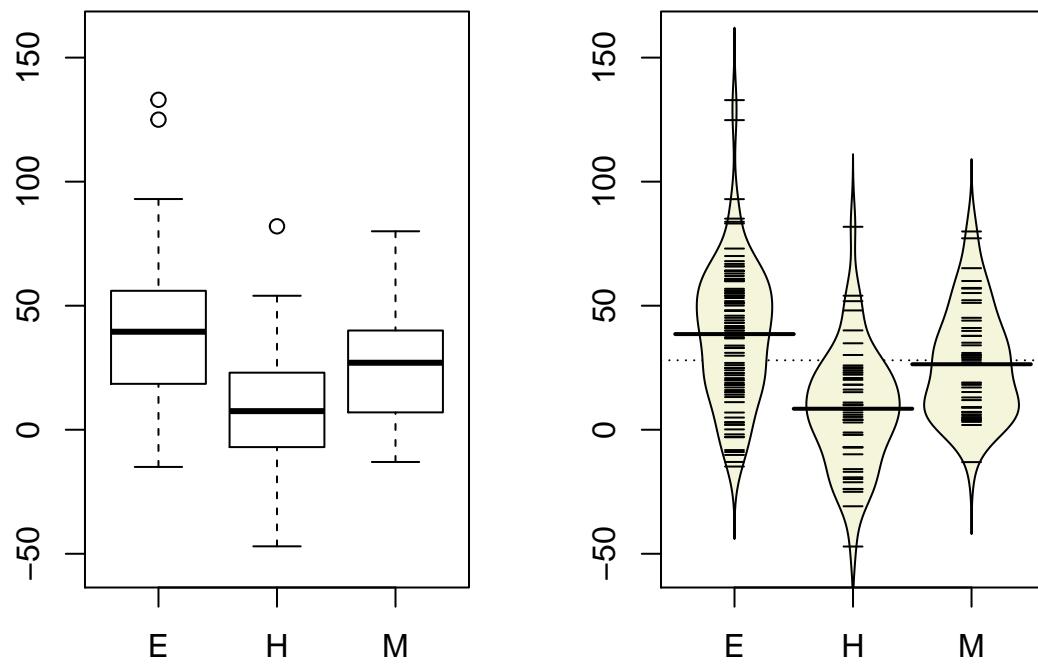


Figure 5.23: Boxplot and beanplot of the API growth scores by level of school (E for elementary, M for Middle, and H for High school).

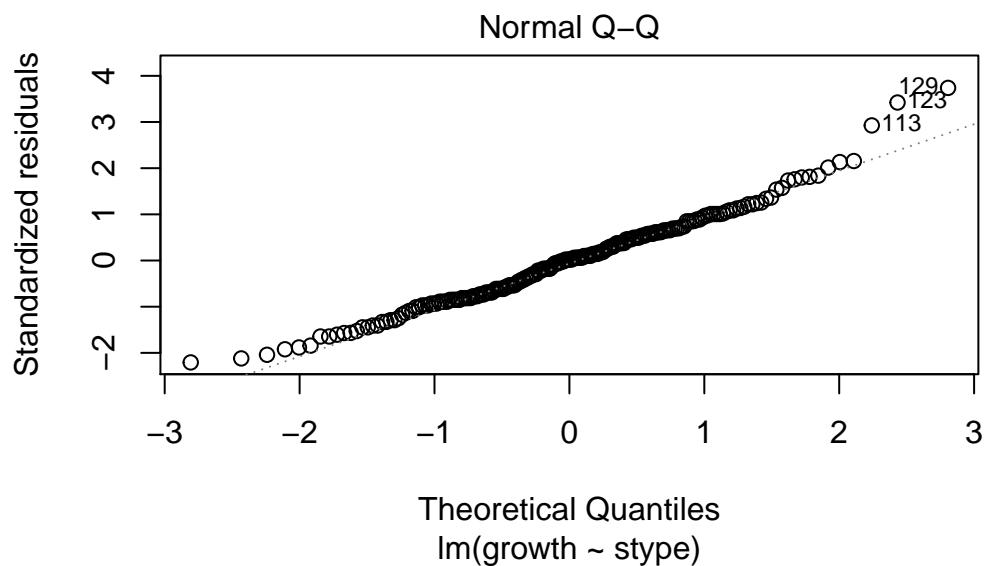


Figure 5.24: QQ-plot of standardized residuals from the One-Way ANOVA linear model.

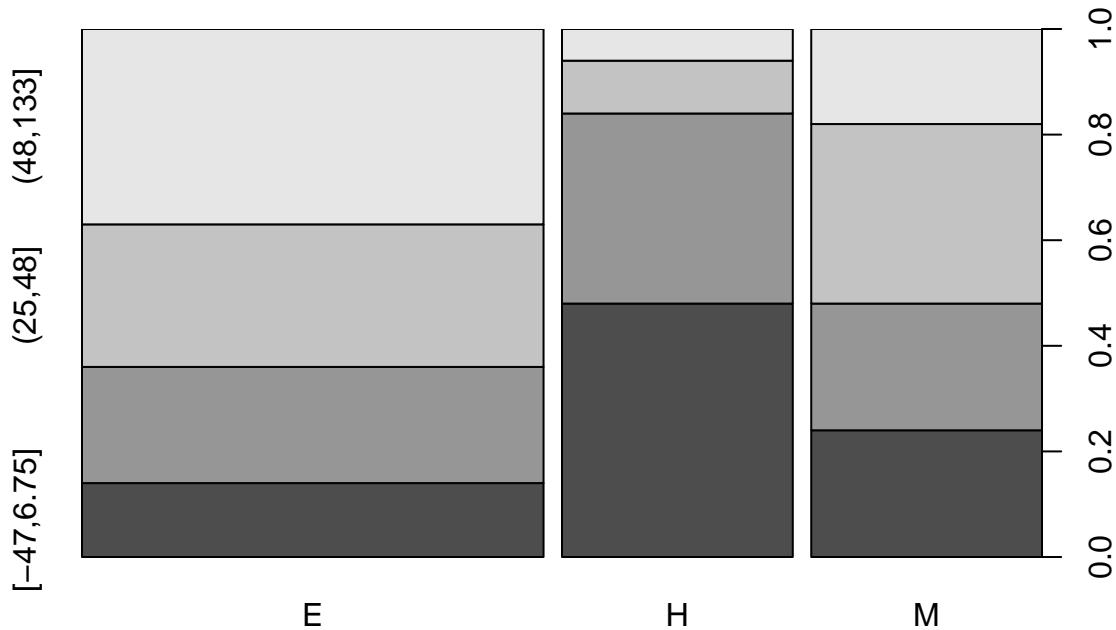


Figure 5.25: Stacked bar chart of the growth category responses by level of school.

a normal distribution. If we **bin** or cut the quantitative response variable into a set of ordered categories and apply a Chi-square test, we can proceed without concern about the lack of normality in the residuals of the ANOVA model. To create these bins, a simple idea would be to use the quartiles to generate the response variable categories, binning the quantitative responses into groups for the lowest 25%, second 25%, third 25%, and highest 25% by splitting the data at  $Q_1$ , the Median, and  $Q_3$ . In R, the **cut** function is available to turn a quantitative variable into a categorical variable. First, we can use the information from **favstats** to find the cut-points:

```
favstats(~growth, data=apistrat)
```

```
##   min    Q1 median Q3 max   mean      sd   n missing
## -47  6.75    25  48 133 27.995 28.1174 200       0
```

The **cut** function can provide the binned variable if it is provided with the end-points of the desired intervals to create new categories with those names in a new variable called **growthcut**.

```
apistrat$growthcut <- cut(apistrat$growth, breaks=c(-47,6.75,25,48,133),
                             include.lowest=T)
tally(~growthcut, data=apistrat)
```

```
## growthcut
## [-47,6.75] (6.75,25] (25,48] (48,133]
##      50        52        49        49
```

Now that we have a categorical response variable, we need to decide which sort of Chi-squared analysis to perform. The sampling design determines the correct analysis as always in these situations. The stratified random sample involved samples from each of the three populations so a Homogeneity test should be employed. In these situations, the stacked bar chart provides the appropriate summary of the data. It also shows us the labels of the categories that the **cut** function created in the new **growthcut** variable:

```
plot(growthcut~stype, data=apistrat)
```

Figure 5.25 suggests that the distributions of growth scores may not be the same across the levels of the

schools with many more high growth *Elementary* schools than in either the *Middle* or *High* school groups (the “high” growth category is labeled as (48, 133] providing the interval of growth scores placed in this category). Similarly, the proportion of the low or negative growth (category of (-47.6, 6.75] for “growth” between -47.6 and 6.75) is lowest in *Elementary* schools and highest in the *High* schools. Statisticians often work across many disciplines and so may not always have the subject area knowledge to know why these differences exist (just like you might not), but an education researcher could take this sort of information – because it is a useful summary of interesting school-level data – and generate further insights into why growth in the API metric may or may not be a good or fair measure of school performance.

Of course, we want to consider whether these results can extend to the population of all California schools. The homogeneity hypotheses for assessing the growth rate categories across the types of schools would be:

- $H_0$ : There is no difference in the distribution of growth categories across the three levels of schools in the population of California schools.
- $H_A$ : There is some difference in the distribution of growth categories across the three levels of schools in the population of California schools.

There might be an issue with the independence assumption in that schools within the same district might be more similar to one another and different between one another. Sometimes districts are accounted for in education research to account for differences in policies and demographics among the districts. We could explore this issue by finding district-level average growth rates and exploring whether those vary systematically but this is beyond the scope of the current exploration.

Checking the expected cell counts gives insight into the assumption for using the  $\chi^2$ -distribution to find the p-value:

```
growthtable <- tally(~stype+growthcut, data=apistrat)
growthtable
```

```
##      growthcut
## stype [-47,6.75] (6.75,25] (25,48] (48,133]
##   E       14       22      27      37
##   H       24       18       5       3
##   M       12       12      17       9
chisq.test(growthtable)$expected
```

```
##      growthcut
## stype [-47,6.75] (6.75,25] (25,48] (48,133]
##   E      25.0      26    24.50    24.50
##   H     12.5      13    12.25    12.25
##   M     12.5      13    12.25    12.25
```

The smallest expected count is 12.25, occurring in four different cells, so the assumptions for using the parametric approach are met.

```
chisq.test(growthtable)

##
##  Pearson's Chi-squared test
##
##  data: growthtable
##  X-squared = 38.668, df = 6, p-value = 8.315e-07
```

The observed test statistic is  $X^2 = 38.67$  and, based on a  $\chi^2(6)$  distribution, the p-value is 0.0000008. This p-value suggests that we should reject the null hypothesis. Then we can conclude that there is strong evidence of some difference in the distribution of API growth of schools among *Elementary*, *Middle* and *High School* in the population of schools in California between 1999 and 2000. We can conclude that there is evidence of some difference in the population (California schools) because the schools were randomly selected from all the California schools but because the level of schools, obviously, cannot be randomly assigned, we cannot say that level of school causes these differences.

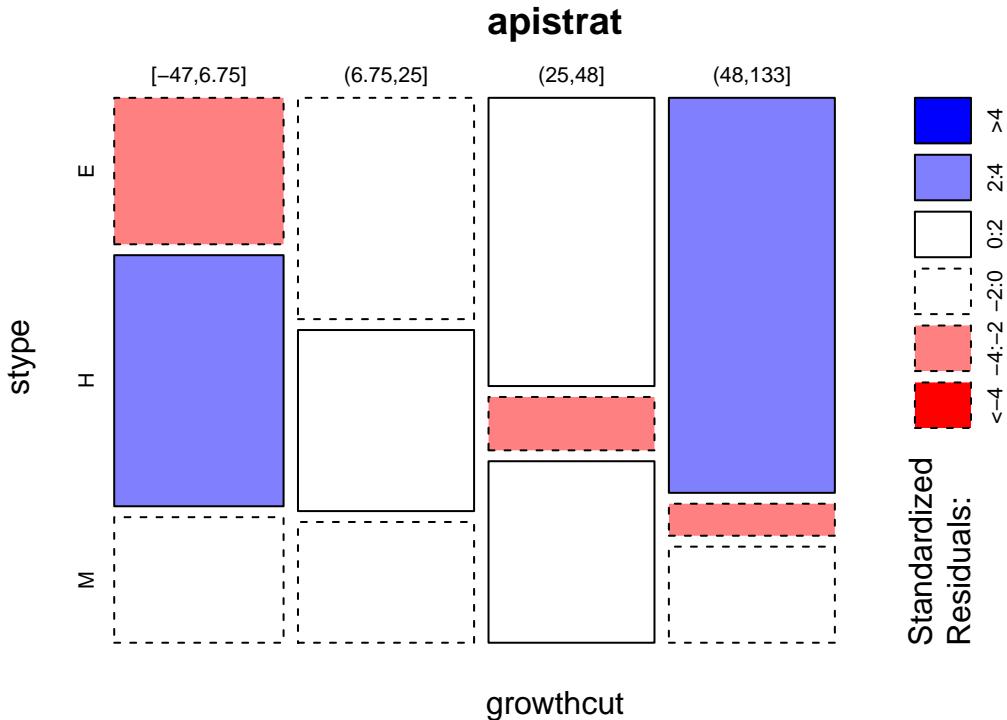


Figure 5.26: Mosaic plot of the API Growth rate categories versus level of the school with shading for size of standardized residuals.

The standardized residuals can enhance this interpretation, displayed in Figure 5.26:

```
chisq.test(growthtable)$residuals
```

```
##      growthcut
## stype [-47,6.75]  (6.75,25]    (25,48]   (48,133]
##     E -2.2000000 -0.7844645  0.5050763  2.5253814
##     H  3.2526912  1.3867505 -2.0714286 -2.6428571
##     M -0.1414214 -0.2773501  1.3571429 -0.9285714
mosaicplot(growthcut~stype,data=apistrat,shade=T)
```

The *Elementary* schools have fewer low/negative growth schools and more high growth schools than expected under the null hypothesis. The *High* schools have more low growth and fewer higher growth (over 25) schools than expected if there were no difference in patterns of response across the school levels. The *Middle* school results were closer to the results expected if there were no differences.

The binning of quantitative variables is not a first step in analyses – the quantitative version is almost always preferable. However, this analysis avoided the violation of the normality assumption that was somewhat problematic for the ANOVA and still provided useful inferences to the differences in the types of schools. When one goes from a quantitative to categorical version of a variable, one loses information (the specific details of the quantitative responses within each level created) and this almost always will result in a loss of statistical power of the procedure. In this situation, the p-value from the ANOVA was of the order  $10^{-10}$  while the Chi-square test had a p-value of order  $10^{-7}$ . This larger p-value is typical of the loss of power in going to a categorical response when more information was available. In many cases, there are no options but to use contingency table analyses. This example shows that there might be some situations where “going categorical” could be an acceptable method for handling situations where an assumption is violated.

## 5.12 Chapter summary

Chi-square tests can be generally used to perform two types of tests, the Independence and Homogeneity tests. The appropriate analysis is determined based on the data collection methodology. The parametric Chi-squared distribution for which these tests are named is appropriate when the expected cell counts are large enough (related to having a large enough overall sample). When this condition is violated, the permutation approach can provide valuable inferences in these situations in most situations.

Data displays of the stacked barchart (Homogeneity) and mosaic plots (Independence) provide a visual summary of the results that can also be found in contingency tables. You should have learned how to calculate the  $X^2$  (X-squared) test statistic based on first finding the expected cell counts. Under certain assumptions, it will follow a Chi-Squared distribution with  $(R - 1)(C - 1)$  degrees of freedom. When those assumptions are not met, it is better to use a permutation approach to find p-values. Either way, the same statistic is used to test either kind of hypothesis, independence or homogeneity. If there is evidence to reject the null hypothesis, then it is interesting to see which cells in the table contributed to the deviations from the null hypothesis. The standardized residuals provide that information. Graphing them in a mosaic plot makes for a fun display to identify the large residuals and often allows you to better understand the results. This should tie back into the original data display and contingency table where you identified initial patterns.

## 5.13 Review of Important R commands

The main components of R code used in this chapter follow with components to modify in red where  $y$  is a response variable and  $x$  is a predictor are easily identified:

- `TABLENAME <- tally(~x + y, data=DATASETNAME, margins=F)`
  - This function requires that the `mosaic` package has been loaded.
  - This provides a table of the counts in the variable called `TABLENAME`.
  - `margins=T` is used if want to display row, column, and table totals.
- `plot(y~x, data=DATASETNAME)`
  - Makes a stacked bar chart useful for homogeneity test situations.
- `mosaicplot(TABLENAME)`
  - Makes a mosaic plot useful for finding patterns in the table in independence test situations.
- `chisq.test(TABLENAME)`
  - Provides  $X^2$  and p-values based on the  $\chi^2$ -distribution with  $(R - 1)(C - 1)$  degrees of freedom.
- `chisq.test(TABLENAME)$expected`
  - Provides expected cell counts.
- `pchisq(X-SQUARED, df=(R- 1)*(C- 1), lower.tail=F)`
  - Provides p-value from  $\chi^2$ -distribution with  $(R - 1)(C - 1)$  degrees of freedom for observed test statistic.
  - See Section 5.5 for code related to finding a permutation-based p-value.
- `chisq.test(TABLENAME)$residuals^2`
  - Provides  $X^2$  contributions from each cell in table.
- `chisq.test(TABLENAME)$residuals`
  - Provides standardized residuals.

- `mosaicplot(TABLENAME, shade=T)`
  - Provides a mosaic plot with shading based on standardized residuals.

## 5.14 Practice problems

Determine which type of test is appropriate in each situation – *Independence* or *Homogeneity*?

5.1. Concerns over diseases being transmitted between birds and humans have led to many areas developing monitoring plans for the birds that are in their regions. The duck pond on campus at MSU-Bozeman is a bit like a night club for the birds that pass through Bozeman.

- i) Suppose that a researcher randomly samples 20 ducks at the duck pond on campus on 4 different occasions and records the number ducks that are healthy and number that are sick on each day. The variables in this study are the day of measurement and sick/healthy.
- ii) In another monitoring study, a researcher goes to a wetland area and collects a random sample from all birds present on a single day, classifies them by type of bird (ducks, swans, etc.) and then assesses whether each is sick or healthy. The variables in this study are type of bird and sick/healthy.

5.2. Psychologists performed an experiment on 48 male bank supervisors attending a management institute to investigate biases against women in personnel decisions. The supervisors were asked to make a decision on whether to promote a hypothetical applicant based on a personnel file. For half of them, the application file described a female candidate; for the others it described a male.

5.3. Researchers collected data on death penalty sentencing in Georgia. For 243 crimes, they categorized the crime by severity from 1 to 6 with Category 1 comprising barroom brawls, liquor-induced arguments, lovers' quarrels, and similar crimes and Category 6 including the most vicious, cruel, cold-blooded, unprovoked crimes. They also recorded the perpetrator's race. They wanted to know if there was a relationship between race and type of crime.

5.4. Epidemiologists want to see if Vitamin C helped people with colds. They would like to give some patients Vitamin C and some a placebo then compare the two groups. However, they are worried that the placebo might not be working. Since vitamin C has such a distinct taste, they are worried the participants will know which group they are in. To test if the placebo was working, they collected 200 subjects and randomly assigned half to take a placebo and the other half to take Vitamin C. 30 minutes later, they asked the subjects which supplement they received (hoping that the patients would not know which group they were assigned to).

5.5. Is zodiac sign related to GPA? 300 randomly selected students from MSU were asked their birthday and their current GPA. GPA was then categorized as < 1.50 = F, 1.51-2.50 = D, 2.51 - 3.25 = C, 3.26-3.75 = B, 3.76-4.0 = A and their birthday was used to find their zodiac sign.

5.6. In 1935, the statistician R. A. Fisher famously had a colleague claim that she could distinguish whether milk or tea was added to a cup first. Fisher presented her, in a random order, 4 cups that were filled with milk first and 4 cups that were filled with tea first.

5.7. Researchers wanted to see if people from Rural and Urban areas aged differently. They contacted 200 people from Rural areas and 200 people from Urban areas and asked the participants their age (<40, 41-50, 51-60, >60).

The FiveThirtyEight Blog often shows up with interesting data summaries that have general public appeal. Their staff includes a bunch of quants with various backgrounds. When starting their blog, they had to decide on the data is/are question that we introduced in Section 2.1. To help them think about this, they collected a nationally representative sample that contained three questions about this. Based on their survey, they concluded that

Relevant to the interests of FiveThirtyEight in particular, we also asked whether people preferred using “data” as a singular or plural noun. To those who prefer the plural, I’ll put this in your terms: The data are pretty conclusive that the vast majority of respondents think we should say “data is.” The singular crowd won by a 58 percentage-point margin, with 79 percent of respondents liking “data is” to 21 percent preferring “data are.” But only half of respondents had put any thought to the usage prior to our survey, so it seems that it’s not a pressing issue for most.

This came from a survey that contained questions about *which is the correct usage*, (`isare`), *have you thought about this issue* (`thoughtabout`) with levels Yes/No, and *do you care about this issue* (`careabout`) with four levels from *Not at all* to *A lot*. The following code loads their data set after missing responses were removed, does a little re-ordering of factor levels to help make the results easier to understand, and makes a tableplot to get a general sense of the results including information on the respondents’ gender, age, income, and education (or just look at the cover of the book).

```
csd<- read.csv("http://www.math.montana.edu/courses/s217/documents/csd.csv")
require(tabplot)
csd$careabout <- factor(csd$careabout,
                           levels=levels(csd$careabout)[c(1,4,3,2)])
csd$Education <- factor(csd$Education,
                           levels=levels(csd$Education)[c(4,3,5,1,2)])
csd$Household.Income <- factor(csd$Household.Income,
                                 levels=levels(csd$Household.Income)[c(1,4,5,6,2,3)])
tableplot(csd[,c("isare","careabout","thoughtabout","Gender",
               "Age","Household.Income","Education")])
```

5.8. If we are interested in the variables `isare` and `careabout`, what sort of test should we perform?

5.9. Make the appropriate plot of the results for the table relating those two variables relative to your answer to 5.8.

5.10. Generate the contingency table and find the expected cell counts, first “by hand” and then check them using the output. Is the parametric procedure appropriate here? Why or why not?

5.11. Report the value of the test statistic, its distribution under the null, the parametric p-value, and write a decision and conclusion, making sure to address scope of inference.

5.12. Make a mosaic plot with the standardized residuals and discuss the results. Specifically, in what way do the is/are preferences move away from the null hypothesis for people that care more about this?

We might be fighting a losing battle on this particular word usage, but since we are in the group that cares a lot about this, we are going to keep trying...



# Chapter 6

## Correlation and Simple Linear Regression

### 6.1 Relationships between two quantitative variables

The independence test in Chapter 5 provided a technique for assessing evidence of a relationship between two categorical variables. The terms ***relationship*** and ***association*** are synonyms that, in statistics, imply that values on one variable tend to occur more often with some other values of the other variable or that knowing something about the level of one variable provides information about the patterns of values on the other variable. These terms are not specific to the “form” of the relationship – any pattern (strong or weak, negative or positive, easily explained or complicated) satisfy the definition. There are two other aspects to using these terms in a statistical context. First, they are not directional – an association between  $x$  and  $y$  is the same as saying there is an association between  $y$  and  $x$ . Second, they are not causal unless the levels of the one of the variables are randomly assigned in an experimental context. We add to this terminology the idea of correlation between variables  $x$  and  $y$ . ***Correlation***, in most statistical contexts, is a measure of the specific type of relationship between the variables: the **linear relationship between two quantitative variables**<sup>1</sup>. So as we start to review these ideas from your previous statistics course, remember that associations and relationships are more general than correlations and it is possible to have no correlation where there is a strong relationship between variables. “Correlation” is used colloquially as a synonym for relationship but we will work to reserve it for its more specialized usage here to refer to the linear relationship.

Assessing and then modeling relationships between quantitative variables drives the rest of the chapters, so we should get started with some motivating examples to start to think about what relationships between quantitative variables “look like”... To motivate these methods, we will start with a study of the effects of beer consumption on blood alcohol levels (*BAC*, in grams of alcohol per deciliter of blood). A group of  $n = 16$  student volunteers at The Ohio State University drank a randomly assigned number of beers<sup>2</sup>. Thirty minutes later, a police officer measured their *BAC*. Your instincts, especially as well-educated college students with some chemistry knowledge, should inform you about the direction of this relationship – that there is a **positive relationship** between Beers and *BAC*. In other words, **higher values of one variable are associated with higher values of the other**. Similarly, lower values of one are associated with lower values of the other. In fact there are online calculators that tell you how much your *BAC* increases for each extra beer consumed (for example: <http://www.craftbeer.com/beer-studies/blood-alcohol-content-calculator>

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<sup>1</sup>There are measures of correlation between categorical variables but when statisticians say correlation they mean correlation of quantitative variables. If they are discussing correlations of other types, they will make that clear.

<sup>2</sup>Some of the details of this study have been lost, so we will assume that the subjects were randomly assigned and that a beer means a can of beer and that the beer was of regular strength. We don’t know if any of that is actually true. I have been asking the Dean of Students to repeat this study as an educational activity at MSU with no success yet.

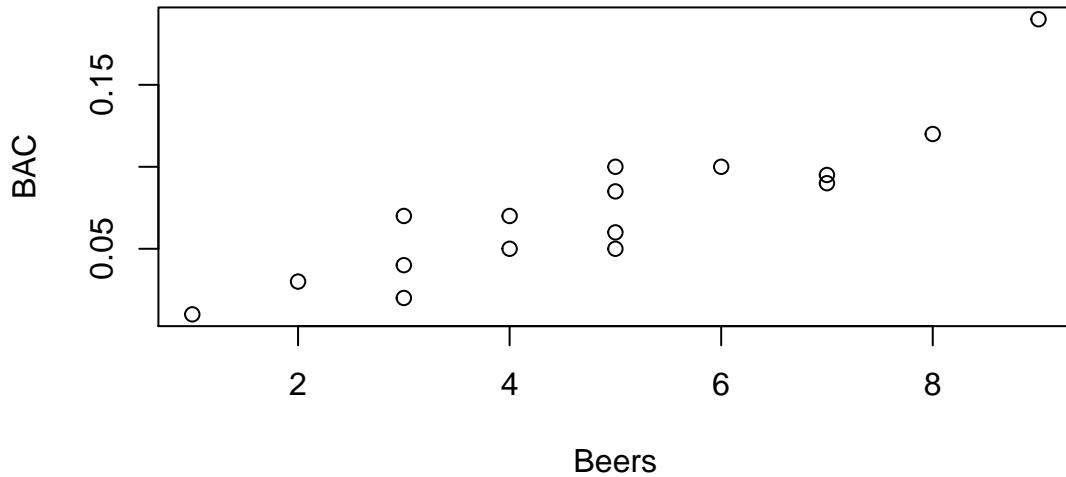


Figure 6.1: Scatterplot of *Beers* consumed versus *BAC*.

if you plug in 1 beer). The increase in  $y$  (BAC) for a 1 unit increase in  $x$  (here, 1 more beer) is an example of **slope coefficient** that is applicable if the relationship between the variables is linear and something that will be fundamental in what we will call **simple linear regression model**. In a simple linear regression model (simple means that there is only one explanatory variable) the slope is the expected change in the mean response for a one unit increase in the explanatory variable. You could also use the *BAC* calculator and the models that we are going to develop to pick a total number of beers you will consume and get a predicted *BAC*.

Before we get to the specifics of this model and how we measure correlation, we should explore the relationship between *Beers* and *BAC* in a scatterplot. Figure 6.1 shows a **scatterplot** of the results that display the expected positive relationship. Scatterplots display the response pairs for the two quantitative variables with the explanatory variable on the x-axis and the response variable on the y-axis. The relationship between *Beers* and *BAC* appears to be relatively linear but there is possibly more variability than one might expect. For example, for students consuming 5 beers, their *BACs* range from 0.05 to 0.10. If you look at the online *BAC* calculators, you will see that other factors such as weight, sex, and beer percent alcohol can impact the results. We might also be interested in previous alcohol consumption. In Chapter 8, we will learn how to estimate the relationship between *Beers* and *BAC* after correcting for those “other variables” using **multiple linear regression**, where we incorporate more than one quantitative explanatory variable into the linear model (somewhat like in the 2-Way ANOVA). Some of this variability might be hard or impossible to explain regardless of the other variables available and is considered unexplained variation and goes into the residual errors in our models, just like in the ANOVA models. To make scatterplots as in Figure 6.1, we simply use `plot(y~x, data=...)`.

```
BB <- read.csv("http://www.math.montana.edu/courses/s217/documents/beersbac.csv")
plot(BAC~Beers, data=BB)
```

There are a few general things to look for in scatterplots:

1. **Assess the direction of the relationship** – is it positive or negative?
2. **Consider the strength of the relationship**. The general idea of assessing strength visually is about how hard or easy it is to see the pattern. If it is hard to see a pattern, then it is weak. If it is easy to

see, then it is strong.

3. **Consider the linearity of the relationship.** Does it appear to curve or does it follow a relatively straight line? Curving relationships are called *curvilinear* or *nonlinear* and can be strong or weak just like linear relationships – it is all about how tightly the points follow the pattern you identify.
4. **Check for unusual observations – outliers** – by looking for points that don't follow the overall pattern. Being large in  $x$  or  $y$  doesn't mean that the point is an outlier. Being unusual relative to the overall pattern makes a point an outlier in this setting.
5. **Check for changing variability** in one variable based on values of the other variable. This will tie into a constant variance assumption later in the regression models.
6. **Finally, look for distinct groups** in the scatterplot. This might suggest that observations from two populations, say males and females, were combined but the relationship between the two quantitative variables might be different for the two groups.

Going back to Figure 6.1 it appears that there is a moderately strong linear relationship between Beers and BAC – not weak but with some variability around what appears to be a straight-line relationship. There might even be a hint of a nonlinear relationship in the higher beer values. There are no clear outliers because the observation at 9 beers seems to be following the overall pattern fairly closely. There is little evidence of non-constant variance mainly because of the limited size of the data set – we'll check this with better plots later. And there are no clearly distinct groups in this plot, mainly because the # of beers was randomly assigned. These data have one more interesting feature to be noted – that subjects managed to consume 8 or 9 beers. This seems to be a large number. I have never been able to trace this data set to the original study so it is hard to know if (1) they had this study approved by a human subjects research review board to make sure it was “safe”, (2) every subject in the study was able to consume their randomly assigned amount, and (3) whether subjects were asked to show up to the study with *BACs* of 0. We also don't know the alcohol concentration of the beer consumed or volume. But it is still a fun example to start these methods with...

In making scatterplots, there is always a choice of a variable for the x-axis and the y-axis. It is our convention to put explanatory or independent variables (the ones used to explain or predict the responses) on the x-axis. In studies where the subjects are randomly assigned to levels of a variable, this is very clearly an explanatory variable, and we can go as far as making causal inferences with it. In observational studies, it can be less clear which variable explains which. In these cases, make the most reasonable choice based on the observed variables but remember that, when the direction of relationship is unclear, you could have switched which axes and implication of which variable explains which.

## 6.2 Estimating the correlation coefficient

In terms of quantifying relationships between variables, we start with the correlation coefficient, a measure that is the same regardless of your choice of which variable is considered explanatory or response. We measure the strength and direction of linear relationships between two quantitative variables using *Pearson's r* or *Pearson's Product Moment Correlation Coefficient*. For those who really like acronyms, Wikipedia even suggests calling it the PPMCC. However, its use is so ubiquitous that the lower case *r* or just “correlation coefficient” are often sufficient to identify that you have used the PPMCC. Some of the extra distinctions arise because there are other ways of measuring correlations in other situations (for example between two categorical variables), but we will not consider them here.

The correlation coefficient, *r*, is calculated as

$$r = \frac{1}{n-1} \sum_{i=1}^n \left( \frac{x_i - \bar{x}}{s_x} \right) \left( \frac{y_i - \bar{y}}{s_y} \right),$$

where  $s_x$  and  $s_y$  are the standard deviations of  $x$  and  $y$ . This formula can also be written as

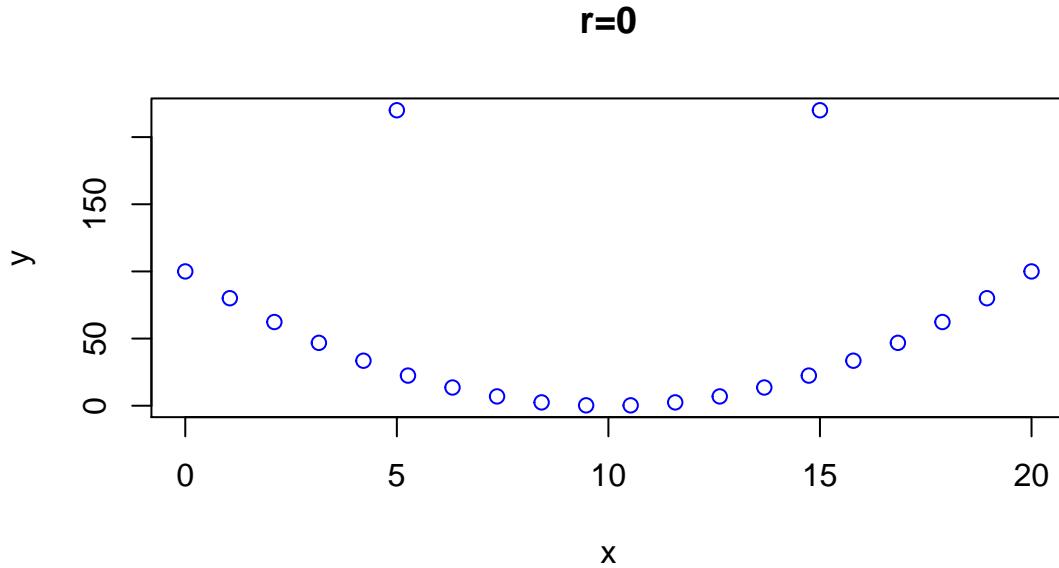


Figure 6.2: Scatterplot of an amusing relationship that has  $r = 0$ .

$$r = \frac{1}{n-1} \sum_{i=1}^n z_{x_i} z_{y_i}$$

where  $z_{x_i}$  is the z-score (observation minus mean divided by standard deviation) for the  $i^{th}$  observation on  $x$  and  $z_{y_i}$  is the z-score for the  $i^{th}$  observation on  $y$ . We won't directly use this formula, but its contents inform the behavior of  $r$ . First, because it is a sum divided by  $(n - 1)$  it is a bit like an average – it combines information across all observations and, like the mean, is sensitive to outliers. Second, it is a dimension-less measure, meaning that it has no units attached to it. It is based on z-scores which have units of standard deviations of  $x$  or  $y$  so the original units of measurement are cancelled out going into this calculation. This also means that changing the original units of measurement, say from Fahrenheit to Celsius or from miles to km for one or the other variable will have no impact on the correlation. Less obviously, the formula guarantees that  $r$  is between -1 and 1. It will attain -1 for a perfect negative linear relationship, 1 for a perfect positive linear relationship, and 0 for no linear relationship. We are being careful here to say *linear relationship* because you can have a strong nonlinear relationship with a correlation of 0. For example, consider Figure 6.2.

There are some conditions for trusting the results that the correlation coefficient provides:

1. Two quantitative variables measured.
  - This might seem silly, but categorical variables can be coded numerically and a meaningless correlation can be estimated if you are not careful what you correlate.
2. The relationship between the variables is relatively linear.
  - If the relationship is nonlinear, the correlation is meaningless and can be misleading.
3. There should be no outliers.
  - The correlation is very sensitive (technically *not resistant*) to the impacts of certain types of outliers and you should generally avoid reporting the correlation when they are present.

- One option in the presence of outliers is to report the correlation with and without outliers to see how they influence the estimated correlation.

The correlation coefficient is dimensionless but larger magnitude values (closer to -1 OR 1) mean stronger linear relationships. A rough interpretation scale based on experiences working with correlations follows, but this varies between fields and types of research and variables measured. It depends on the levels of correlation researchers become used to obtaining, so can even vary within fields. Use this scale until you develop your own experience:

- $|r| < 0.3$ : weak linear relationship,
- $0.3 < |r| < 0.7$ : moderate linear relationship,
- $0.7 < |r| < 0.9$ : strong linear relationship, and
- $0.9 < |r| < 1.0$ : very strong linear relationship.

And again note that this scale only relates to the **linear** aspect of the relationship between the variables.

When we have linear relationships between two quantitative variables,  $x$  and  $y$ , we can obtain estimated correlations from the `cor` function either using `y~x` or by running the `cor` function<sup>3</sup> on the entire data set. When you run the `cor` function on a data set it produces a **correlation matrix** which contains a matrix of correlations where you can triangulate the variables being correlated by the row and column names, noting that the correlation between a variable and itself is 1. A matrix of correlations is useful for comparing more than two variables, discussed below.

```
require(mosaic)
cor(BAC~Beers, data=BB)

## [1] 0.8943381
cor(BB)

##           Beers         BAC
## Beers  1.0000000 0.8943381
## BAC    0.8943381 1.0000000
```

Based on either version of using the function, we find that the correlation between `Beers` and `BAC` is estimated to be 0.89. This suggests a strong linear relationship between the two variables. Examples are about the only way to build up enough experience to become skillful in using the correlation coefficient. Some additional complications arise in more complicated studies as the next example demonstrates.

Gude and Haggerty (2009) explored the relationship between average summer temperature (degrees F) and area burned (natural log of hectares<sup>4</sup> =  $\log(\text{hectares})$ ) by wildfires in Montana from 1985 to 2007. The **log-transformation** is often used to reduce the impacts of really large observations on non-negative (strictly greater than 0) responses with really large observations (more on **transformations** and their impacts on regression models in Chapter 7). Based on your experiences and before analyzing the data, I'm sure you would assume that summer temperature explains the area burned by wildfires. But could it be that more fires are related to having warmer summers? That second direction is unlikely on a state-wide scale but could apply at a particular weather station that is near a fire. There is another option – some other variable is affecting both variables. For example, drier summers might be the real explanatory variable that is related to having both warm summers and lots of fires. These variables are also being measured over time making them examples of **time series**. In statistics, time series data is generally reserved for evenly spaced measurements over time, like monthly or yearly measurements. In this situation, if there are changes over time, they might be attributed to climate change. So there are really three relationships to explore with the variables measured here (remembering that the full story might require measuring even more!): log-area burned versus temperature, temperature versus year, and log-area burned versus year.

<sup>3</sup>This interface with the `cor` function only works after you load the `mosaic` package.

<sup>4</sup>The natural log ( $\log_e$  or  $\ln$ ) is used in statistics so much that the function in R `log` actually takes the natural log and if you want a  $\log_{10}$  you have to use the function `log10`. When statisticians say `log` we mean natural log.

With more than two variables, we can use the `cor` function on all the variables and end up getting a matrix of correlations or, simply, the ***correlation matrix***.

```
mtfires <- read.csv("http://www.math.montana.edu/courses/s217/documents/climateR2.csv")
# natural log transformation of area burned
mtfires$loghectacres <- log(mtfires$hectacres)

#Cuts the original hectares data so only log-scale version in data.frame
mtfiresR <- mtfires[,-3]
cor(mtfiresR)

##          Year Temperature loghectacres
## Year      1.0000000 -0.0037991   0.3617789
## Temperature -0.0037991  1.0000000   0.8135947
## loghectacres  0.3617789   0.8135947   1.0000000
```

If you triangulate the row and column labels, that cell provides the correlation between that pair of variables. For example, in the first row (`Year`) and the last column (`loghectacres`), you can find that the correlation coefficient is  $r=0.362$ . Note the symmetry in the matrix around the diagonal of 1's – this further illustrates that correlation between  $x$  and  $y$  does not depend on which variable is viewed as the “response”. The estimated correlation between `Temperature` and `Year` is -0.004 and the correlation between `loghectacres` (*log-hectares burned*) and `Temperature` is 0.81. So `Temperature` has almost no linear change over time. And there is a strong linear relationship between `loghectacres` and `Temperature`. So it appears that temperatures may be related to log-area burned but that the trend over time in both is less clear (at least the linear trends).

The correlation matrix alone is misleading – we need to explore scatterplots to check for nonlinear relationships, outliers, and clustering of observations that may be distorting the numerical measure of the linear relationship. The `pairs.panels` function from the `psych` package (Revelle, 2017) combines the numerical correlation information and scatterplots in one display. There are some options to turn off for the moment but it is an easy function to use to get lots of information in one place. As in the correlation matrix, you triangulate the variables for the pairwise relationship. The upper right diagonal of Figure 6.3 displays a correlation of 0.36 for `Year` and `loghectacres` and the lower left panel contains the scatterplot with `Year` on the x-axis and `loghectacres` on the y-axis. The correlation between `Year` and `Temperature` is really small, both in magnitude and in display, but appears to be nonlinear (it goes down between 1985 and 1995 and then goes back up), so the correlation coefficient doesn't mean much here since it just measures the overall linear relationship. We might say that this is a moderate strength (moderately “clear”) curvilinear relationship. In terms of the underlying climate process, it suggests a decrease in summer temperatures between 1985 and 1995 and then an increase in the second half of the data set.

```
require(psych)
pairs.panels(mtfiresR, ellipses=F, scale=T, smooth=F, col=0)
```

As one more example, the Australian Institute of Sport collected data on 102 male and 100 female athletes that are available in the `ais` data set from the `alr3` package (Weisberg (2011), Weisberg (2005)). They measured a variety of variables including the athlete's Hematocrit (`Hc`, units of percentage of red blood cells in the blood), Body Fat Percentage (`Bfat`, units of percentage of total body weight), and height (`Ht`, units of cm). Eventually we might be interested in predicting `Hc` based on the other variables, but for now the associations are of interest.

```
require(alr3)
data(ais)
aisR <- ais[,c("Ht", "Hc", "Bfat")]
summary(aisR)

##           Ht                  Hc                  Bfat
##  Min.   :148.9   Min.   :35.90   Min.   : 5.630
##  1st Qu.:174.0   1st Qu.:40.60   1st Qu.: 8.545
##  Median :179.7   Median :43.50   Median :11.650
##  Mean    :180.1   Mean    :43.09   Mean    :13.507
```

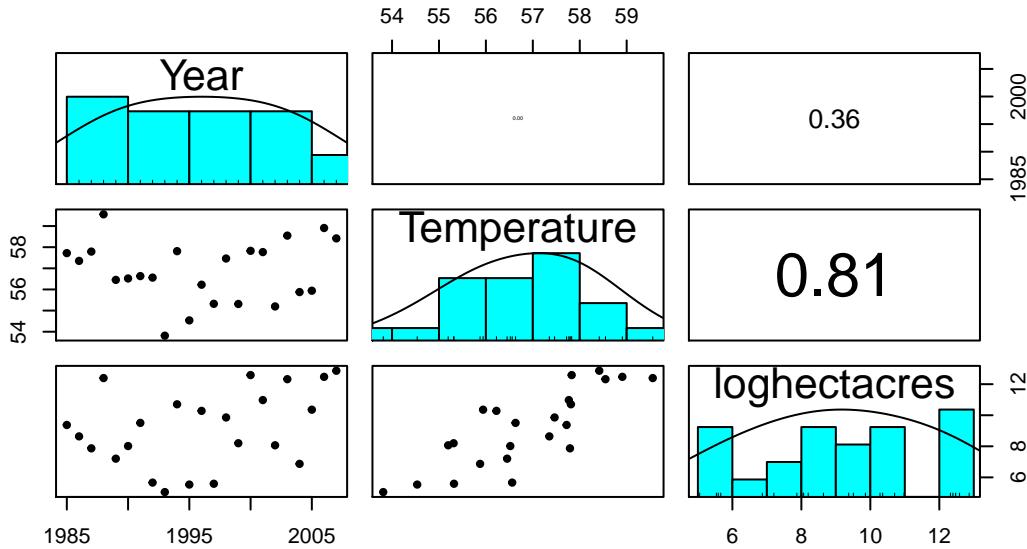


Figure 6.3: Scatterplot matrix of Montana fires data.

```

## 3rd Qu.:186.2   3rd Qu.:45.58   3rd Qu.:18.080
## Max.    :209.4   Max.    :59.70   Max.    :35.520
cor(aisR)

##          Ht         Hc         Bfat
## Ht 1.0000000 0.3711915 -0.1880217
## Hc 0.3711915 1.0000000 -0.5324491
## Bfat -0.1880217 -0.5324491 1.0000000
pairs.panels(aisR,scale=T,ellipse=F,smooth=F,col=0)

```

*Ht (Height)* and *Hc (Hematocrit)* have a moderate positive relationship that may contain a slightly nonlinearity. It also contains one clear outlier for a middle height athlete (around 175 cm) with an *Hc* of close to 60% (a result that is extremely high). One might wonder about whether this athlete has been doping or if that measurement involved a recording error. We should consider removing that observation to see how our results might change without it impacting the results. For the relationship between *Bfat (body fat)* and *Hc (hematocrit)*, that same high *Hc* value is a clear outlier. There is also a high *Bfat (body fat)* athlete (35%) with a somewhat low *Hc* value. This also might be influencing our impressions so we will remove both “unusual” values and remake the plot. The two offending observations were found for individuals numbered 56 and 166 in the data set:

```
aisR[c(56, 166),]
```

```

##      Ht     Hc     Bfat
## 56 179.8 37.6 35.52
## 166 174.9 59.7  9.56

```

We can create a reduced version of the data (*aisR2*) by removing those two rows using `[-c(56, 166),]` and then remake the plot:

```

aisR2 <- aisR[-c(56, 166),] #Removes observations in rows 56 and 166
pairs.panels(aisR2, scale=T, ellipse=F, smooth=F, col=0)

```

After removing these two unusual observations, the relationships between the variables are more obvious (Figure 6.5). There is a moderate strength, relatively linear relationship between *Height* and *Hematocrit*.

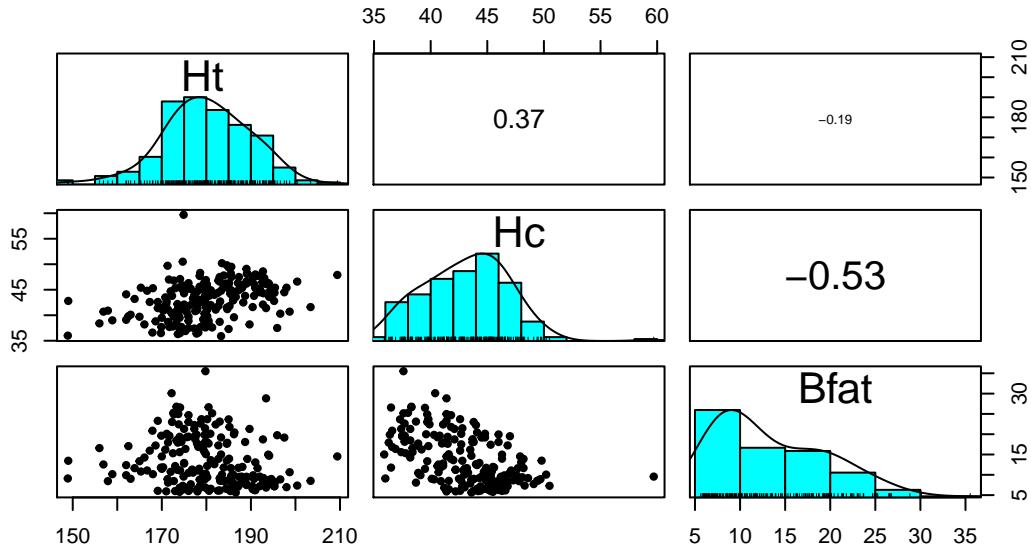


Figure 6.4: Scatterplot matrix of athlete data.

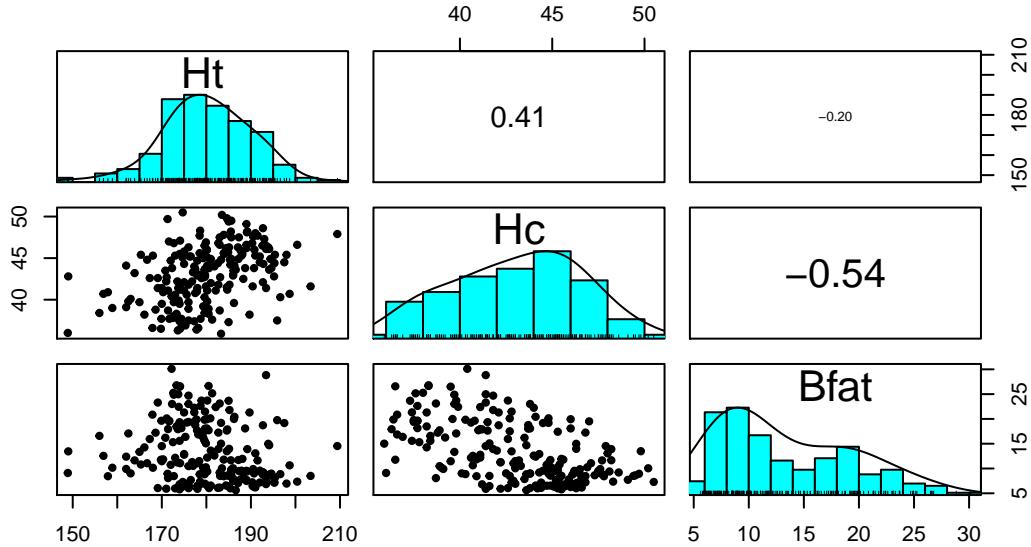


Figure 6.5: Scatterplot matrix of athlete data with two potential outliers removed.

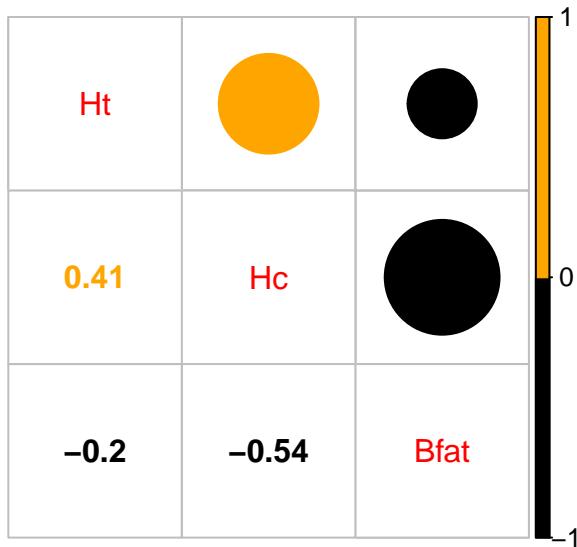


Figure 6.6: Correlation plot of the athlete data with two potential outliers removed. Lighter (orange) circle for positive correlations and black for negative correlations.

There is almost no relationship between *Height* and *Body Fat %* ( $r=-0.20$ ). There is a negative, moderate strength, somewhat curvilinear relationship between *Hematocrit* and *Body Fat %* ( $r=-0.54$ ). As hematocrit increases initially, the body fat percentage decreases but at a certain level (around 45% for Hc), the body fat percentage seems to level off. Interestingly, it ended up that removing those two outliers had only minor impacts on the estimated correlations – this will not always be the case.

Sometimes we want to just be able to focus on the correlations, assuming we trust that the correlation is a reasonable description of the results between the variables. To make it easier to see patterns of positive and negative correlations, we can employ a different version of the same display from the `corrplot` package (Wei and Simko, 2016). In this case (Figure 6.6), it tells much the same story but also allows the viewer to easily distinguish both size and direction and read off the numerical correlations if desired.

```
require(corrplot)
corrplot.mixed(cor(aisR2), col=c("black", "orange"))
```

## 6.3 Relationships between variables by groups

In assessing the relationship between variables, incorporating information from a third variable can often enhance the information gathered by either showing that the relationship between the first two variables is the same across levels of the other variable or showing that it differs. When the other variable is categorical (or just can be made categorical), it can be added to scatterplots, changing the symbols and colors for the points based on the different groups. These techniques are especially useful if the categorical variable corresponds to potentially distinct groups in the responses. In the previous example, the data set was built with male and female athletes. For some characteristics, the relationships might be the same for both sexes but for others, there are likely some physiological differences to consider.

We could continue to use the `plot` function here, but it would require additional lines of code to add these extra features. The `scatterplot` function from the `car` package (Fox and Weisberg, 2011) makes it easy to incorporate information from an additional categorical variable. We'll add to our regular formula idea

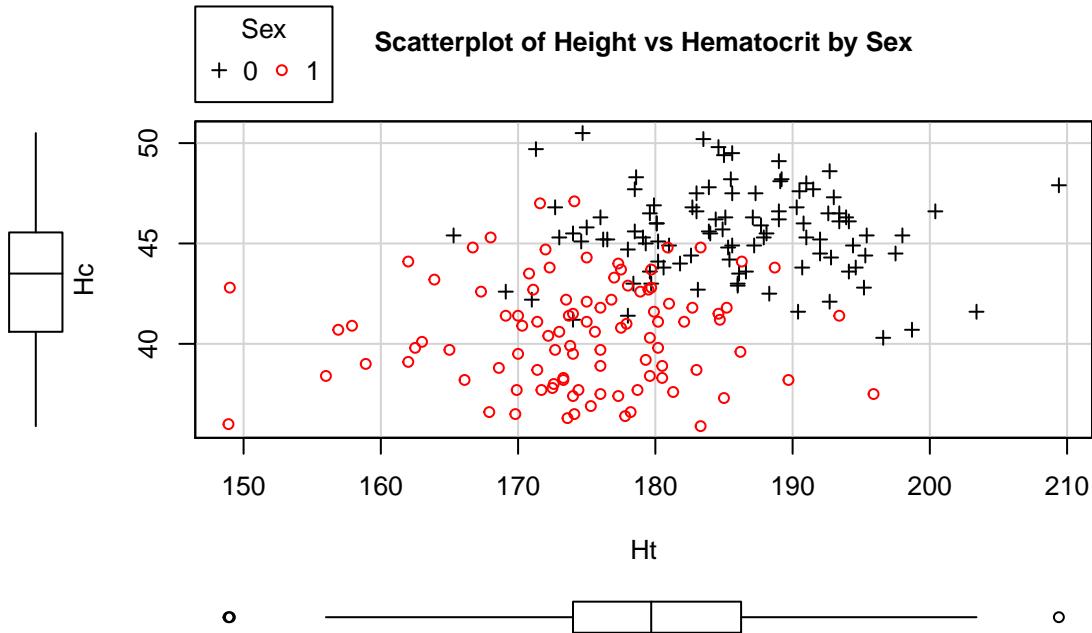


Figure 6.7: Scatterplot of athlete's height and hematocrit by sex of athletes. Males were coded as 0s and females as 1s.

( $y \sim x$ ) the vertical line “|” followed by the categorical variable  $z$ , such as  $y \sim x | z$ . As noted earlier, in statistics, “|” means “to condition on” or, here, consider the relationship between  $y$  and  $x$  by groups in  $z$ . The other options are mainly to make it easier to read the information in the plot... Using this enhanced notation, Figure 6.7 displays the *Height* and *Hematocrit* relationship with information on the sex of the athletes where sex was coded 0 for males and 1 for females.

```
aisR2 <- ais[-c(56,166),c("Ht","Hc","Bfat","Sex")]
require(car)
scatterplot(Hc~Ht|Sex, data=aisR2, pch=c(3,21), reg.line=F, smoother=F,
           boxplots="xy", main="Scatterplot of Height vs Hematocrit by Sex")
```

Adding the grouping information really changes the impressions of the relationship between *Height* and *Hematocrit* – within each sex, there is little relationship between the two variables. The overall relationship is of moderate strength and positive but the subgroup relationships are weak at best. The overall relationship is created by inappropriately combining two groups that had different means in both the  $x$  and  $y$  directions. Men have higher mean heights and hematocrit values than women and putting them together in one large group creates the misleading overall relationship.

To get the correlation coefficients by groups, we can subset the data set using a logical inquiry on the `Sex` variable in the updated `aisR2` data set, using `Sex==0` to get the male subjects only and `Sex==1` for the female subjects, then running the `cor` function on each version of the data set:

```
cor(Hc~Ht, data=aisR2[aisR2$Sex==0,]) #Males only
```

```
## [1] -0.04756589
cor(Hc~Ht, data=aisR2[aisR2$Sex==1,]) #Females only
```

```
## [1] 0.02795272
```

These results show that  $r=-0.05$  for *Height* and *Hematocrit* for *males* and  $r=0.03$  for *females*. The first suggests a very weak negative linear relationship and the second suggests a very weak positive linear relationship. The correlation when the two groups were combined (and group information conclusion here is

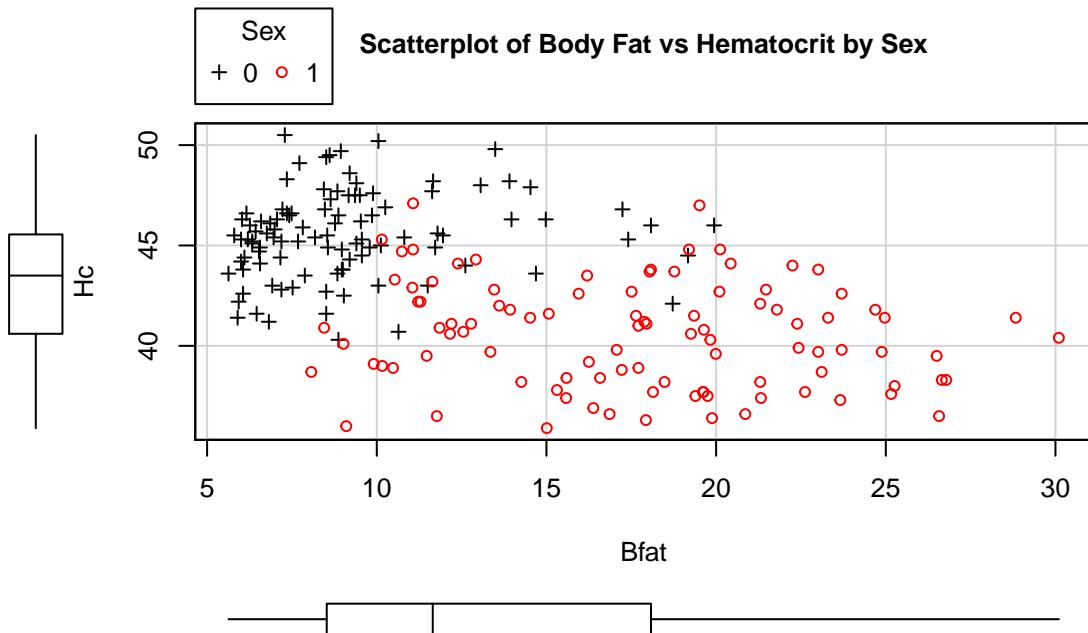


Figure 6.8: Scatterplot of athlete's body fat and hematocrit by sex of athletes. Males were coded as 0s and females as 1s.

that correlations on data sets that contain groups can be very misleading. It also emphasizes the importance of exploring for potential subgroups in the data set – these two groups were not obvious in the initial plot, but with added information the real story became clear.

For the *Body Fat vs Hematocrit* results in Figure 6.8, with an overall correlation of  $r=-0.54$ , the subgroup correlations show weaker relationships that also appear to be in different directions ( $r=0.13$  for men and  $r=-0.17$  for women). This doubly reinforces the dangers of aggregating different groups and ignoring the group information.

```
cor(Hc~Bfat, data=aisR2[aisR2$Sex==0,]) #Males only
```

```
## [1] 0.1269418
cor(Hc~Bfat, data=aisR2[aisR2$Sex==1,]) #Females only
```

```
## [1] -0.1679751
scatterplot(Hc~Bfat|Sex, data=aisR2, pch=c(3,21), reg.line=F, smoother=F,
            boxplots="xy", main="Scatterplot of Body Fat vs Hematocrit by Sex")
```

One final exploration for these data involves the *body fat* and *height* relationship displayed in Figure 6.9. This relationship shows an even greater disparity between overall and subgroup results. The overall relationship is characterized as a weak negative relationship ( $r=-0.20$ ) that is not clearly linear or nonlinear. The subgroup relationships are both clearly positive with a stronger relationship for men that might also be nonlinear (for the linear relationships  $r =0.20$  for women and  $r =0.45$  for men). Especially for male athletes, those that are taller seem to have higher body fat percentages. This might be related to the types of sports they compete in – that would be another categorical variable we could incorporate... Both groups also seem to demonstrate slightly more variability in *Body Fat* associated with taller athletes (each sort of “fans out”).

```
cor(Bfat~Ht, data=aisR2[aisR2$Sex==0,]) #Males only
```

```
## [1] 0.1954609
```

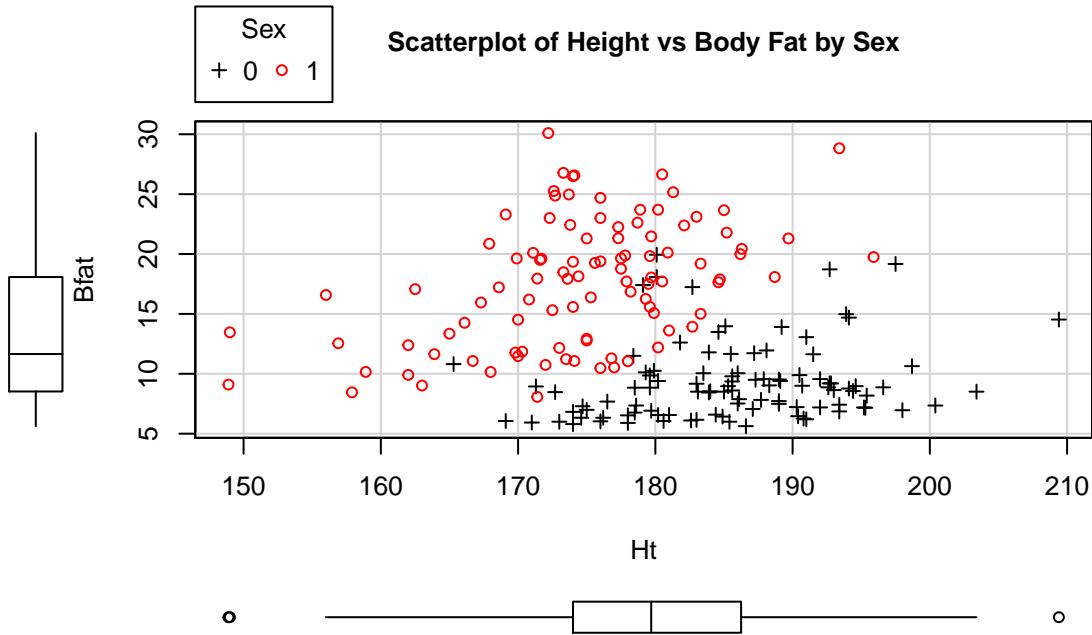


Figure 6.9: Scatterplot of athlete's body fat and height by sex.

```
cor(Bfat~Ht,data=aisR2[aisR2$Sex==1,]) #Females only
## [1] 0.4476962
scatterplot(Bfat~Ht|Sex, data=aisR2, pch=c(3,21), reg.line=F, smoother=F,
            boxplots="xy", main="Scatterplot of Height vs Body Fat by Sex")
```

In each of these situations, the sex of the athletes has the potential to cause misleading conclusions if ignored. There are two ways that this could occur – if we did not measure it then we would have no hope to account for it OR we could have measured it but not adjusted for it in our results, as I did initially. We distinguish between these two situations by defining the impacts of this additional variable as either a confounding or lurking variable:

- **Confounding variable:** affects the response variable and is related to the explanatory variable. The impacts of a confounding variable on the response variable cannot be separated from the impacts of the explanatory variable.
- **Lurking variable:** a potential confounding variable that is not measured and is not considered in the interpretation of the study.

Lurking variables show up in studies sometimes due to lack of knowledge of the system being studied or a lack of resources to measure these variables. And sometimes the two variables cannot be separated even if they are measured...

To help think about confounding and lurking variables, consider the following situation. On many highways, such as Highway 93 in Montana and north into Canada, recent construction efforts have been involved in creating safe passages for animals by adding fencing and animal crossing structures. These structures both can improve driver safety, save money from costs associated with animal-vehicle collisions, and increase connectivity of animal populations. Researchers involved in these projects are interested in which characteristics of underpasses lead to the most successful structures, mainly measured by rates of animal usage (number of times they cross under the road). Crossing structures are typically made using culverts and those tend to be cylindrical. Researchers are interested in studying the effect of height and width of crossing structures on animal usage. Unfortunately, all the tallest structures are also the widest structures. If animals prefer the

tall and wide structures, then there is no way to know if it is due to the height or width of the structure since they are confounded. If the researcher had only measured width, then they might assume that it is the important characteristic of the structures but height could be a lurking variable that really was the factor related to animal usage of the structures. This is an example where it may not be possible to design a study that prevents confounding of the two variables *height* and *width*. If the researchers could control the height and width of the structures, then they could randomly assign both variables to make sure that some narrow structures are installed that are tall and some that are short. Additionally, they would also want to have some wide structures that are short and some are tall. Careful design of studies can prevent confounding of variables if they are known in advance and it is possible to control them, but in observational studies the observed combinations of variables are uncontrollable. This is why we need to employ additional caution in interpreting results from observational studies.

## 6.4 Inference for the correlation coefficient (Optional Section)

We used bootstrapping briefly in Chapter 2 to generate nonparametric confidence intervals based on the middle 95% of the bootstrapped version of the statistic. Remember that bootstrapping involves sampling *with replacement* from the data set and creates a distribution centered near the statistic from the real data set. This also mimics sampling under the alternative as opposed to sampling under the null as in our permutation approaches. Bootstrapping is particularly useful for making confidence intervals where the distribution of the statistic may not follow a named distribution. This is the case for the correlation coefficient which we will see shortly.

The correlation is an interesting summary but it is also an estimator of a population parameter called  $\rho$  (the symbol rho), which is the ***population correlation coefficient***. When  $\rho = 1$ , we have a perfect positive linear relationship in the population; when  $\rho = -1$ , there is a perfect negative linear relationship in the population; and when  $\rho = 0$ , there is no linear relationship in the population. Therefore, to test if there is a linear relationship between two quantitative variables, we use the null hypothesis  $H_0 : \rho = 0$  (tests if the true correlation,  $\rho$ , is 0 – no linear relationship). The alternative hypothesis is that there is some (positive or negative) relationship between the variables in the population,  $H_A : \rho \neq 0$ . The distribution of the Pearson correlation coefficient can be complicated in some situations, so we will use bootstrapping methods to generate confidence intervals for  $\rho$  based on repeated random samples with replacement from the original data set. If the confidence contains 0, then we would fail to reject the null hypothesis since 0 is in the interval of our likely values for  $\rho$ . If the confidence interval does not contain 0, then we can reject the null hypothesis.

The *beers* and *BAC* example seemed to provide a strong relationship with  $r=0.89$ . As correlations approach -1 or 1, the sampling becomes more and more skewed. This certainly shows up in the bootstrap distribution that the following code produces (Figure 6.10). Remember that bootstrapping utilizes the `resample` function applied to the data set to create new realizations of the data set by re-sampling with replacement from those observations. The bold vertical line in Figure 6.10 corresponds to the estimated correlation  $r=0.89$  and the distribution contains a noticeable left skew with a few much smaller  $T^*$ 's possible in bootstrap samples. The  $C\%$  confidence interval is found based on the middle  $C\%$  of the distribution or by finding the values that put  $(100 - C)/2$  into each tail of the distribution.

```
Tobs <- cor(BAC~Beers, data=BB); Tobs
## [1] 0.8943381
par(mfrow=c(1,2))
B <- 1000
Tstar<-matrix(NA, nrow=B)
for (b in (1:B)){
  Tstar[b] <- cor(BAC~Beers, data=resample(BB))
}
quantiles <- qdata(Tstar, c(.025,.975)) #95% Confidence Interval
quantiles
```

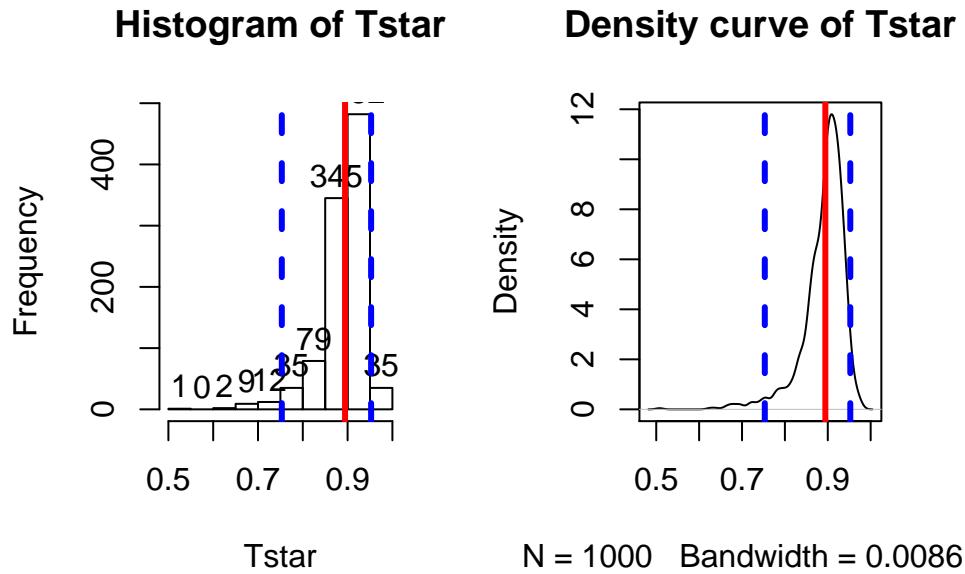


Figure 6.10: Histogram and density curve of the bootstrap distribution of the correlation coefficient with bold vertical line for observed correlation and dashed lines for 95% bootstrap confidence interval.

```
##      quantile    p
## 2.5%  0.7531517 0.025
## 97.5% 0.9524785 0.975
hist(Tstar, labels=T)
abline(v=Tobs, col="red", lwd=3)
abline(v=quantiles$quantile, col="blue", lty=2, lwd=3)

plot(density(Tstar), main="Density curve of Tstar")
abline(v=Tobs, col="red", lwd=3)
abline(v=quantiles$quantile, col="blue", lty=2, lwd=3)
```

These results tell us that the bootstrap 95% CI is from 0.75 to 0.96 – we are 95% confident that the true correlation between *Beers* and *BAC* in all OSU students like those that volunteered is between 0.75 and 0.96. Note that there are no units on the correlation coefficient.

We can also use this confidence interval to test for a linear relationship between these variables.

- $H_0 : \rho = 0$  : There is no linear relationship between *Beers* and *BAC*\* in the population.
- $H_A : \rho \neq 0$  : There is a linear relationship between *Beers* and *BAC* in the population.

The 95% confidence level corresponds to a 5% significance level test. The 95% CI is from 0.75 to 0.96, which does not contain 0, so we can reject the null hypothesis. We conclude that there is strong evidence to conclude that there is a linear relationship between *Beers* and *BAC* in OSU students. We'll revisit this example using the upcoming regression tools to explore the potential for more specific conclusions about this relationships. For these inferences to be accurate, we need to be able to trust that the sample correlation is reasonable for characterizing the relationship between these variables.

In this situation with randomly assigned levels of  $x$  and a rejected null hypothesis of no relationship, we can further conclude that changing beer consumption **causes** changes in the *BAC*. This is a much stronger conclusion than we can typically make based on correlation coefficients. Correlations and scatterplots

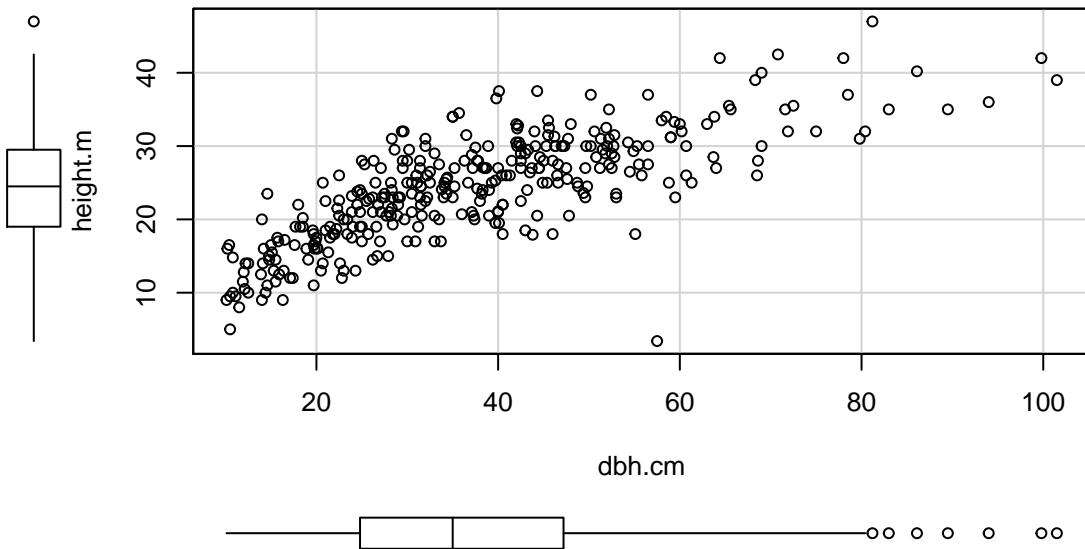


Figure 6.11: Scatterplot of tree heights (m) vs tree diameters (cm).

are enticing for infusing causal interpretations in non-causal situations. We often repeat the mantra that ***correlation is not causation*** and that generally applies – except when there is randomization involved in the study. It is rarer for researchers either to assign, or even to be able to assign, levels of quantitative variables so correlations should be viewed as non-causal unless the details of the study suggest otherwise.

## 6.5 Are tree diameters related to tree heights?

In a study at the Upper Flat Creek study area in the University of Idaho Experimental Forest, a random sample of  $n = 336$  trees was selected from the forest, with measurements recorded on Douglas Fir, Grand Fir, Western Red Cedar, and Western Larch trees. The data set called `ufc` is available from the `spuRs` package (Jones et al., 2014) and contains `dbh.cm` (tree diameter at 1.37 m from the ground, measured in cm) and `height.m` (tree height in meters). The relationship displayed in Figure 6.11 is positive, moderately strong with some curvature and increasing variability as the diameter increases. There do not appear to be groups in the data set but since this contains four different types of trees, we would want to revisit this plot by group.

```
require(spuRs) #install.packages("spuRs")
data(ufc)
scatterplot(height.m~dbh.cm, data=ufc, smooth=F, reg.line=F)
```

Of particular interest is an observation with a diameter around 58 cm and a height of less than 5 m. Observing a tree with a diameter around 60 cm is not unusual in the data set, but none of the other trees with this diameter had heights under 15 m. It turns out that the likely outlier is in observation number 168 and because it is so unusual it likely corresponds to either a damaged tree or a recording error.

```
ufc[168,]
```

```
##      plot tree species dbh.cm height.m
## 168    67     6      WL   57.5      3.4
```

With the outlier in the data set, the correlation is 0.77 and without it, the correlation increases to 0.79. The removal does not create a big change because the data set is relatively large and the *diameter* value is close

to the mean of the  $x$ 's<sup>5</sup> but it has some impact on the strength of the correlation.

```
cor(dbh.cm~height.m, data=ufc)

## [1] 0.7699552
cor(dbh.cm~height.m, data=ufc[-168,])

## [1] 0.7912053
```

If you skipped Section 6.4, you can skip the rest of this section:

With the outlier included, the bootstrap 95% confidence interval goes from 0.702 to 0.820 – we are 95% confident that the true correlation between *diameter* and *height* in the population of trees is between 0.702 and 0.820. When the outlier is dropped from the data set, the 95% bootstrap CI is 0.752 to 0.827, which shifts the lower endpoint of the interval up, reducing the width of the interval from 0.118 to 0.075. In other words, the uncertainty regarding the value of the population correlation coefficient is reduced. The reason to remove the observation is that it is unusual based on the observed pattern, which implies an error in data collection or sampling from a population other than the one used for the other observations and, if the removal is justified, it helps us refine our inferences for the population parameter. But measuring the linear relationship in these data where there is a curve violates one of our assumptions of using these methods – we'll see some other ways of detecting this issue in Section 6.10 and we'll try to “fix” this example using transformations in the Chapter 7.

```
## [1] 0.7699552

##      quantile     p
## 2.5%  0.7017769 0.025
## 97.5% 0.8203489 0.975

## [1] 0.7912053

##      quantile     p
## 2.5%  0.752252 0.025
## 97.5% 0.827492 0.975

Tobs <- cor(dbh.cm~height.m, data=ufc); Tobs
par(mfrow=c(2,1))
B <- 1000
Tstar <- matrix(NA, nrow=B)
for (b in (1:B)){
  Tstar[b] <- cor(dbh.cm~height.m, data=resample(ufc))
}
quantiles <- qdata(Tstar, c(.025,.975)) #95% Confidence Interval
quantiles
hist(Tstar, labels=T, xlim=c(0.6,0.9),
      main="Bootstrap distribution of correlation with all data")
abline(v=Tobs, col="red", lwd=3)
abline(v=quantiles$quantile, col="blue", lty=2, lwd=3)

Tobs <- cor(dbh.cm~height.m, data=ufc[-168,]); Tobs
Tstar <- matrix(NA, nrow=B)
for (b in (1:B)){
  Tstar[b] <- cor(dbh.cm~height.m, data=resample(ufc[-168,]))
}
quantiles <- qdata(Tstar, c(.025,.975)) #95% Confidence Interval
quantiles
```

---

<sup>5</sup>Observations at the edge of the  $x$ 's will be called high leverage points in Section 6.9; this point is a low leverage point because it is close to mean of the  $x$ 's.

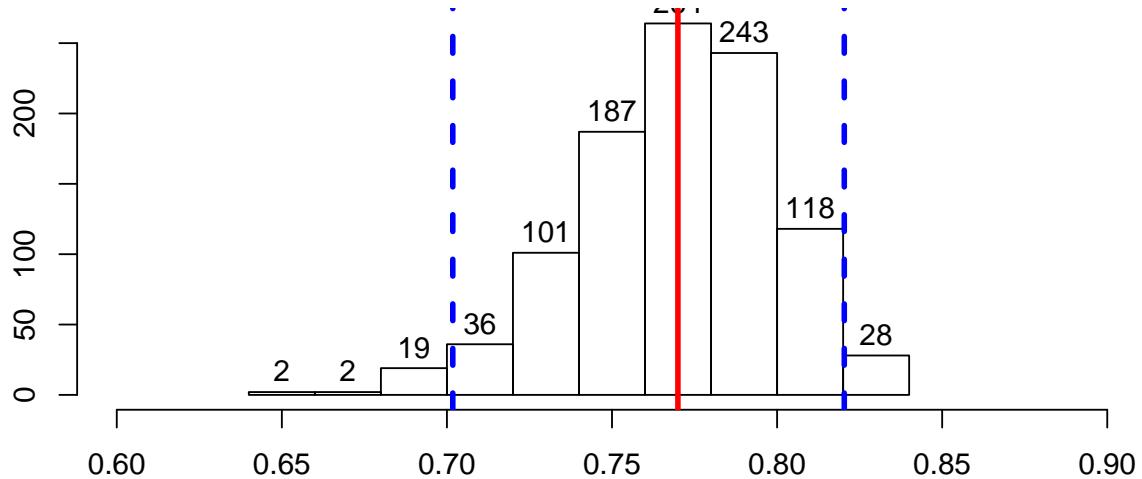
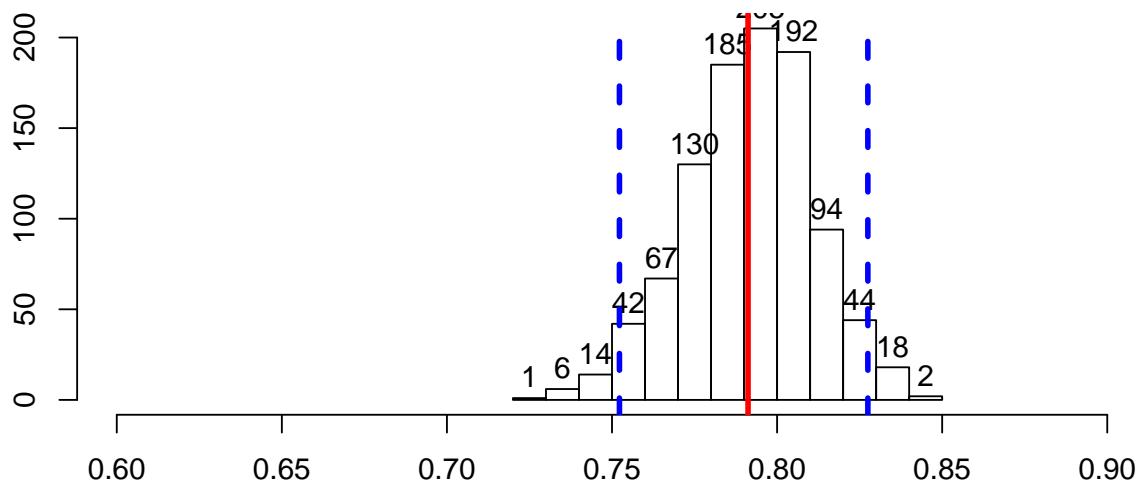
**Bootstrap distribution of correlation with all data****Bootstrap distribution of correlation without outlier**

Figure 6.12: Bootstrap distributions of the correlation coefficient for the full data set (top) and without potential outlier included (bottom) with observed correlation (bold line) and bounds for 95% confidence interval (dashed lines).



Figure 6.13: Scatterplot with estimate regression line for the *Beers* and *BAC* data.

```
hist(Tstar, labels=T, xlim=c(0.6,0.9),
      main= "Bootstrap distribution of correlation without outlier")
abline(v=Tobs, col="red", lwd=3)
abline(v=quantiles$quantile, col="blue", lty=2, lwd=3)
```

## 6.6 Describing relationships with a regression model

When the relationship appears to be relatively linear, it makes sense to estimate and then interpret a line to represent the relationship between the variables. This line is called a **regression line** and involves finding a line that best fits (explains variation in) the response variable for the given values of the explanatory variable. For regression, it matters which variable you choose for  $x$  and which you choose for  $y$  – for correlation it did not matter. This regression line describes the “effect” of  $x$  on  $y$  and also provides an equation for predicting values of  $y$  for given values of  $x$ . The *Beers* and *BAC* data provide a nice example to start our exploration of regression models. The beer consumption is a clear explanatory variable, detectable in the story because (1) it was randomly assigned to subjects and (2) basic science supports beer consumption amount being an explanatory variable for *BAC*. In some situations, this will not be so clear, but look for random assignment or scientific logic to guide your choices of variables as explanatory or response. Regression lines are actually provided by default in the `scatterplot` function with the `reg.line=T` option or just omitting `reg.line=F` from the previous versions of the code.

```
scatterplot(BAC~Beers, ylim=c(0,.2), xlim=c(0,9), data=BB,
            boxplot=F, main="Scatterplot with regression line",
            lwd=2, smooth=F)
abline(v=1:9,col="grey")
abline(h=c(0.05914,0.0771),col="blue",lty=2,lwd=2)
```

The equation for a line is  $y = a + bx$ , or maybe  $y = mx + b$ . In the version  $mx + b$  you learned that  $m$  is a slope coefficient that relates a change in  $x$  to changes in  $y$  and that  $b$  is a  $y$ -intercept (the value of  $y$  when  $x$

is 0). In Figure 6.13, some extra lines are added to help you see the defining characteristics of the line. The slope, whatever letter you use, is the change in  $y$  for a one-unit increase in  $x$ . Here, the slope is the change in BAC for a 1 beer increase in **Beers**, such as the change from 4 to 5 beers. The y-values (blue, dashed lines) for **Beers** = 4 and 5 go from 0.059 to 0.077. This means that for a 1 beer increase (+1 unit change in  $x$ ), the BAC goes up by  $0.077 - 0.059 = 0.018$  (+0.018 unit change in  $y$ ). We can also try to find the y-intercept on the graph by looking for the BAC level for 0 **Beers** consumed. The y-value (BAC) ends up being around -0.01 if you extend the regression line to **Beers**=0. You might assume that the BAC should be 0 for **Beers**=0 but the researchers did not observe any students at 0 **Beers**, so we don't really know what the BAC might be at this value. We have to use our line to ***predict*** this value. This ends up providing a prediction below 0 – an impossible value for *BAC*. If the y-intercept were positive, it would suggest that the students showed up to the experiment after having already had some drink(s).

The numbers reported were very accurate because we weren't using the plot alone to generate the values – we were using a statistical method and R to estimate the equation to describe the relationship between **Beers** and **BAC**. In statistics, we estimate “ $m$ ” and “ $b$ ”. We also write the equation starting with the y-intercept and we use slightly different notation that allows us to extend to more complicated models with more variables. Specifically, the estimated regression equation is  $\hat{y} = b_0 + b_1x$ , where

- $\hat{y}$  is the estimated value of  $y$  for a given  $x$ ,
- $b_0$  is the estimated y-intercept (predicted value of  $y$  when  $x$  is 0),
- $b_1$  is the estimated slope coefficient, and
- $x$  is the explanatory variable.

One of the differences between when you learned equations in algebra classes and our situation is that the line is not a perfect description of the relationship between  $x$  and  $y$  – it is an “on average” description and will usually leave differences between the line and the observations, which we call residuals ( $e = y - \hat{y}$ ). We worked with residuals in the ANOVA<sup>6</sup> material. The residuals describe the vertical distance in the scatterplot between our model (regression line) and the actual observed data point. The lack of a perfect fit of the line to the observations distinguishes statistical equations from those you learned in math classes. The equations work the same, but we have to modify interpretations of the coefficients to reflect this.

We also tie this estimated model to a theoretical or ***population regression model***:

$$y_i = \beta_0 + \beta_1 x_i + \varepsilon_i$$

where:

- $y_i$  is the observed response for the  $i^{th}$  observation,
- $x_i$  is the observed value of the explanatory variable for the  $i^{th}$  observation,
- $\beta_0 + \beta_1 x_i$  is the true mean function evaluated at  $x_i$
- $\beta_0$  is the true (or population) y-intercept,
- $\beta_1$  is the true (or population) slope coefficient, and
- the deviations,  $\varepsilon_i$ , are assumed to be independent and normally distributed with mean 0 and standard deviation  $\sigma$  or, more compactly,  $\varepsilon_i \sim N(0, \sigma^2)$ .

This presents another version of the linear model from Chapters 3 and 4, now with a quantitative explanatory variable instead of categorical variables. Chapter 6 focuses mostly on the estimated regression coefficients, but remember that we are doing statistics and our desire is to make inferences to a larger population. So, every estimated coefficient  $b_0$  and  $b_1$ , are approximations to theoretical coefficients,  $\beta_0$  and  $\beta_1$ . In other words,  $b_0$  and  $b_1$  are the statistics that try to estimate the true population parameters  $\beta_0$  and  $\beta_1$ , respectively.

---

<sup>6</sup>The residuals from these methods and ANOVA are the same because they all come from linear models but are completely different from the standardized residuals used in the Chi-square material in Chapter 5.

To get estimated regression coefficients, we use the `lm` function and our standard `lm(y~x, data=...)` setup. This is the same function used to estimate our ANOVA models and much of this will look familiar. In fact, the ties between ANOVA and regression are deep and fundamental but not the topic of this section. For the *Beers* and *BAC* example, the ***estimated regression coefficients*** can be found from:

```
m1 <- lm(BAC~Beers, data=BB)
m1
```

```
##
## Call:
## lm(formula = BAC ~ Beers, data = BB)
##
## Coefficients:
## (Intercept)      Beers
## -0.01270     0.01796
```

More often, we will extract these from the ***coefficient table*** produced by a model `summary`:

```
summary(m1)
```

```
##
## Call:
## lm(formula = BAC ~ Beers, data = BB)
##
## Residuals:
##    Min     1Q   Median     3Q    Max
## -0.027118 -0.017350  0.001773  0.008623  0.041027
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) -0.012701  0.012638 -1.005   0.332
## Beers       0.017964  0.002402  7.480 2.97e-06
##
## Residual standard error: 0.02044 on 14 degrees of freedom
## Multiple R-squared:  0.7998, Adjusted R-squared:  0.7855
## F-statistic: 55.94 on 1 and 14 DF, p-value: 2.969e-06
```

From either version of the output, you can find the estimated y-intercept in the `(Intercept)` part of the output and the slope coefficient in the `Beers` part of the output. So  $b_0 = -0.0127$ ,  $b_1 = 0.01796$ , and the ***estimated regression equation*** is

$$\widehat{\text{BAC}}_i = -0.0127 + 0.01796 \text{Beers}_i.$$

This is the equation that was plotted in Figure 6.13. In writing out the equation, it is good to replace  $x$  and  $y$  with the variable names to make the predictor and response variables clear. **If you prefer to write all equations with  $x$  and  $y$ , you need to define  $x$  and  $y$  or else these equations are not clearly defined.**

There is a general interpretation for the slope coefficient that you will need to master. In general, we interpret the slope coefficient as:

- **Slope interpretation (general):** For a 1 [*unit of X*] increase in  $X$ , we expect, *on average*, a  $b_1$  [*unit of Y*] change in  $Y$ .

Figure 6.14 can help you think about the different sorts of slope coefficients we might need to interpret, both providing changes in the response variable for 1 unit increases in the predictor variable.

Applied to this problem, for each additional 1 beer consumed, we expect a 0.018 gram per dL change in the *BAC on average*. Using “change” in the interpretation for what happened in the response allows you to use the same template for the interpretation even with negative slopes – be careful about saying “decrease” when the slope is negative as you can create a double-negative and end up implying an increase... Note also

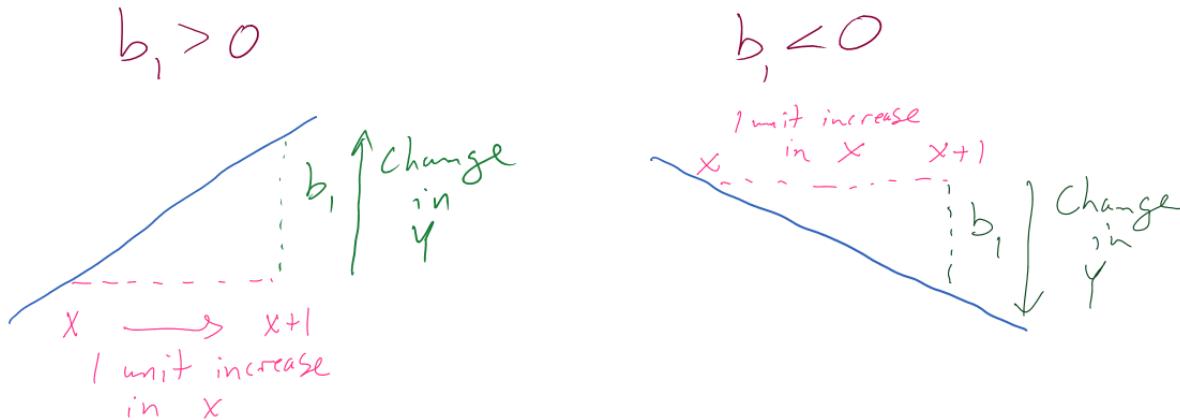


Figure 6.14: Diagram of interpretation of slope coefficients.

that you need to carefully incorporate the units of  $x$  and the units  $y$  to make the interpretation clear. For example, if the change in  $BAC$  for 1 beer increase is 0.018, then we could also modify the beer increase and then the estimated change in  $BAC$  is  $10 * 0.018 = 0.18$  g/dL. Both are correct as long as you are clear about the change in  $x$  you are talking about. Typically, we will just use the units used in the original variables and only change the scale of “change in  $x$ ” when it provides an interpretation we are particularly interested in.

Similarly, the general interpretation for a y-intercept is:

- **Y-intercept interpretation (general):** For  $X=0$  [units of  $X$ ], we expect, on average,  $b_0$  [units of  $Y$ ] in  $Y$ .

Again, applied to the  $BAC$  data set: For 0 beers for *Beers* consumed, we expect, on average, -0.012 g/dL  $BAC$ . The y-intercept interpretation is often less interesting than the slope interpretation but can be interesting in some situations. Here, it is predicting average  $BAC$  for *Beers*=0, which is a value outside the scope of the  $x$ ’s (*Beers* was observed between 1 and 9). Prediction outside the scope of the predictor values is called **extrapolation**. Extrapolation is dangerous at best and misleading at worst. That said, if you are asked to interpret the y-intercept you should still interpret it, but it is also good to note if it is outside of the region where we had observations on the explanatory variable. Another example is useful for practicing how to do these interpretations.

In the Australian Athlete data, we saw a weak negative relationship between *Body Fat* (% body weight that is fat) and *Hematocrit* (% red blood cells in the blood). The scatterplot in Figure 6.15 shows just the results for the female athletes along with the regression line which has a negative slope coefficient. The estimated regression coefficients are found using the `lm` function:

```
aisR2 <- ais[-c(56, 166), c("Ht", "Hc", "Bfat", "Sex")]
m2 <- lm(Hc ~ Bfat, data=aisR2[aisR2$Sex==1,]) #Results for Females
summary(m2)
```

```
##
## Call:
## lm(formula = Hc ~ Bfat, data = aisR2[aisR2$Sex == 1, ])
##
## Residuals:
##      Min       1Q   Median       3Q      Max
## -5.2399 -2.2132 -0.1061  1.8917  6.6453
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) 42.01378   0.93269 45.046 <2e-16
## Bfat        -0.08504   0.05067 -1.678   0.0965
```

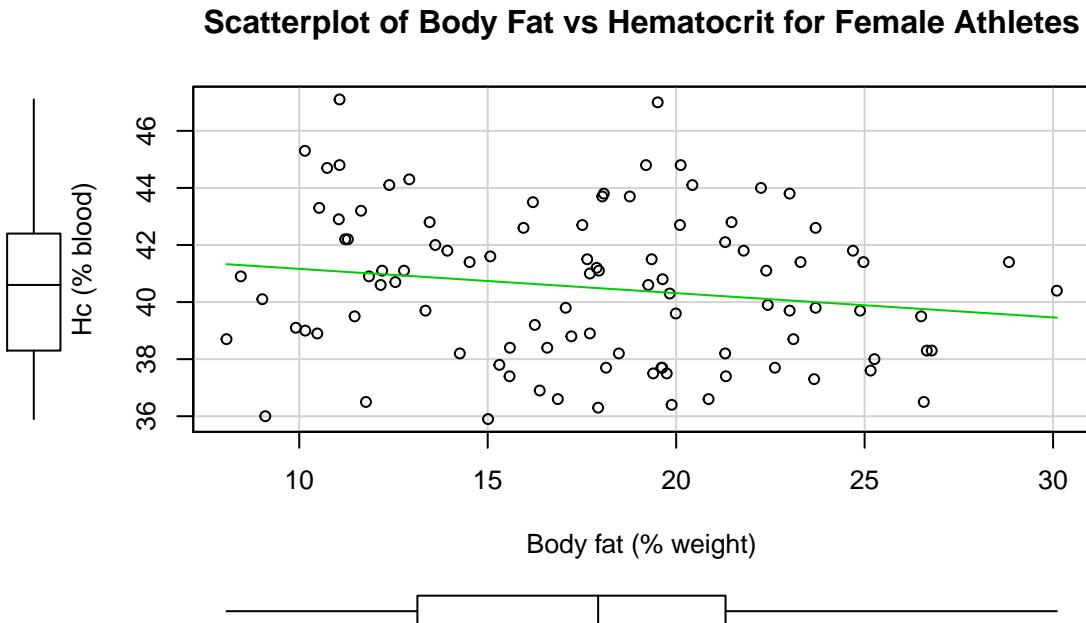


Figure 6.15: Scatterplot of Hematocrit versus Body Fat for female athletes.

```
##  
## Residual standard error: 2.598 on 97 degrees of freedom  
## Multiple R-squared:  0.02822,   Adjusted R-squared:  0.0182  
## F-statistic: 2.816 on 1 and 97 DF,  p-value: 0.09653  
scatterplot(Hc~Bfat, data=aisR2[aisR2$Sex==1,], smooth=F,  
           main="Scatterplot of Body Fat vs Hematocrit for Female Athletes",  
           ylab="Hc (% blood)", xlab="Body fat (% weight)")
```

Based on these results, the estimated regression equation is  $\widehat{Hc}_i = 42.014 - 0.085\text{BodyFat}_i$  with  $b_0 = 42.014$  and  $b_1 = -0.085$ . The slope coefficient interpretation is: For a one percent increase in body fat, we expect, on average a  $-0.085\%$  (blood) change in Hematocrit for Australian female athletes. For the y-intercept, the interpretation is: For a 0% body fat female athlete, we expect a Hematocrit of 42.014% on average. Again, this y-intercept involves extrapolation to a region of the athletes had body fat below 5% so we don't know what would happen to the hematocrit of an athlete that had no body fat except that it probably would not continue to follow a linear relationship.

## 6.7 Least Squares Estimation

The previous results used the `lm` function as a “black box” to generate the estimated coefficients. The lines produced probably look reasonable but you could imagine drawing other lines that might look equally plausible. Because we are interested in explaining variation in the response variable, we want a model that in some sense minimizes the residuals ( $e_i = y_i - \hat{y}_i$ ) to find a model that explains the responses as well as possible – has  $y_i - \hat{y}_i$  as small as possible. We can't just add these up because it would always be 0 (remember why we use the variance to measure spread from introductory statistics?). We use a similar technique in regression, we find the regression line that minimizes the squared residuals  $e_i^2 = (y_i - \hat{y}_i)^2$  over all the observations, minimizing the **Sum of Squared Residuals** =  $\Sigma e_i^2$ . Finding the estimated regression coefficients that minimize the sum of squared residuals is called the **least squares estimation** and provides us a reasonable method for finding the “best” estimated regression line of all the possible choices.

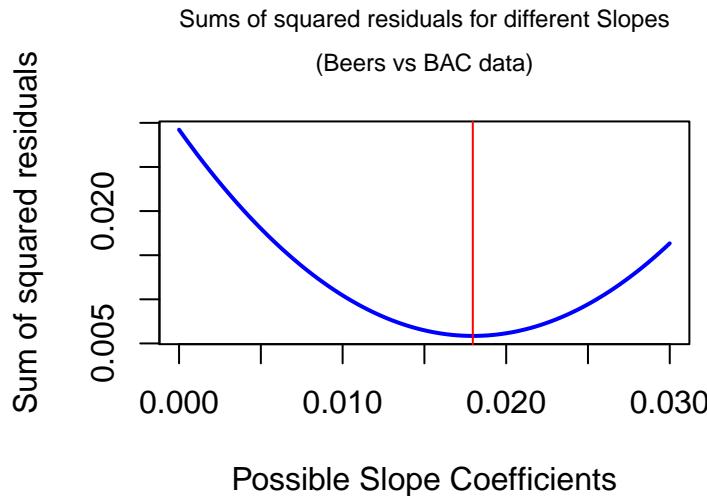


Figure 6.16: Plot of sum of squared residuals vs possible slope coefficients for *Beers* vs *BAC* data, with vertical line for the least squares estimate that minimizes the sum of squared residuals.

For the *Beers* vs *BAC* data, Figure 6.16 shows the result of a search for the optimal slope coefficient between values of 0 and 0.03. The plot shows how the sum of the squared residuals was minimized for the value that `lm` returned at 0.018. The main point of this is that if any other slope coefficient was tried, it did not do as good **on the least squares criterion** as the least squares estimates.

Sometimes it is helpful to have a go at finding the estimates yourself. If you install and load the `tigerstats` (Robinson and White, 2016) and `manipulate` (Allaire, 2014) packages in R-studio and then run `FindRegLine()`, you get a chance to try to find the optimal slope and intercept for a fake data set. Click on the “sprocket” icon in the upper left of the plot and you will see something like Figure 6.17. This interaction can help you see how the residuals are being measured in the y-direction and appreciate that `lm` takes care of this for us.

```
> require(tigerstats)
> require(manipulate)
> FindRegLine()
```

Equation of the regression line is:  
 $y = 4.34 + -0.02x$

Your final score is 13143.99  
 Thanks for playing!

It ends up that the least squares criterion does not require a search across coefficients or trial and error – there are some “simple” equations available for calculating the estimates of the y-intercept and slope:

$$b_1 = \frac{\sum_i(x_i - \bar{x})(y_i - \bar{y})}{\sum_i(x_i - \bar{x})^2} = r \frac{s_y}{s_x} \text{ and } b_0 = \bar{y} - b_1 \bar{x}.$$

You will never need to use these equations but they do inform some properties of the regression line. The slope coefficient,  $b_1$ , is based on the variability in  $x$  and  $y$  and the correlation between them. If  $r=0$ , then the slope coefficient will also be 0. The intercept is a function of the means of  $x$  and  $y$  and what the estimated slope coefficient is **If the slope coefficient,  $b_1$ , is 0, then  $b_0 = \bar{y}$**  (which is just the mean of the response variable for all observed values of  $x$  – this is a very boring model!) The slope is 0 when the correlation is 0. So when there is no linear relationship between  $x$  and  $y$  ( $r = 0$ ), the least squares regression line is a horizontal line with height  $\bar{y}$ , and the line produces the same fitted values for all  $x$  values. You can also think

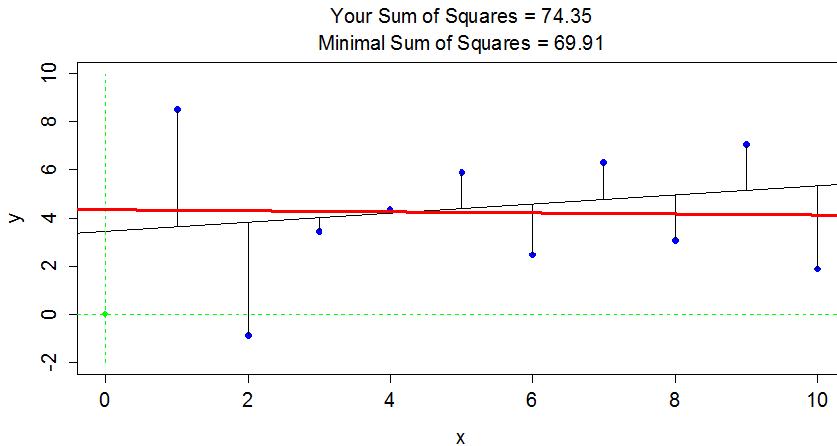


Figure 6.17: Results of running `FindRegLine()` where the user (Greenwood) didn't quite find the least squares line. The correct line is the bold (red) line and produced a smaller sum of squared residuals than the guessed thinner (black) line.

about this as when there is no relationship between  $x$  and  $y$ , the best prediction of  $y$  is the mean of the  $y$ -values and it doesn't change based on the values of  $x$ . It is less obvious in these equations, but they also imply that **the regression line ALWAYS goes through the point  $(\bar{x}, \bar{y})$** . It provides a sort of anchor point for all regression lines.

For one more example, we can revisit the Montana wildfire areas burned (log-hectares) and the average summer temperature (degrees F), which had  $r=0.81$ . The interpretations of the different parts of the regression model follow the least squares estimation provided by `lm`:

```
fire1<-lm(loghectacres~Temperature,data=mtfires)
summary(fire1)
```

```
##
## Call:
## lm(formula = loghectacres ~ Temperature, data = mtfires)
##
## Residuals:
##    Min     1Q   Median     3Q    Max 
## -3.0822 -0.9549  0.1210  1.0007  2.4728 
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) -69.7845   12.3132 -5.667 1.26e-05  
## Temperature   1.3884    0.2165  6.412 2.35e-06  
## 
## Residual standard error: 1.476 on 21 degrees of freedom
## Multiple R-squared:  0.6619, Adjusted R-squared:  0.6458 
## F-statistic: 41.12 on 1 and 21 DF, p-value: 2.347e-06
```

- Regression Equation (Completely Specified):

$$\begin{aligned} & - \widehat{\log(\text{Ha})} = -69.78 + 1.39\text{Temp} \quad \text{OR} \\ & - \hat{y} = -69.78 + 1.39x \text{ with Y:log(Ha) and X:Temperature} \end{aligned}$$

- Response Variable: Yearly *log* Hectares burned by wildfires
- Explanatory Variable: Average Summer Temperature

- Estimated y-Intercept ( $b_0$ ): -69.78
- Estimated slope ( $b_1$ ): 1.39
- Slope Interpretation: For a 1 degree Fahrenheit increase in Average Summer Temperature we would expect, **on average**, a 1.39 change in  $\log(\text{Hectares})$  burned.
- Y-intercept Interpretation: If temperature were 0 degrees F, we would expect -69.78  $\log(\text{Hectares})$  burned **on average**.

One other use of regression equations is for prediction. It is a trivial exercise (or maybe not – we'll see when you try it!) to plug an  $x$ -value of interest into the regression equation and get an estimate for  $y$  at that  $x$ . Basically, the regression lines displayed in the scatterplots show the predictions from the regression line across the range of  $x$ 's. Formally, ***prediction*** involves estimating the response for a particular value of  $x$ . We know that it won't be perfect but it is our best guess. Suppose that we are interested in predicting the log-area burned for a summer that had an average temperature of 59°F. If we plug 59°F into the regression equation,  $\widehat{\log(\text{Ha})} = -69.78 + 1.39 \bullet \text{Temp}$ , we get

$$\begin{aligned}\widehat{\log(\text{Ha})} &= -69.78 \text{ log-hectacres} + 1.39 \text{ log-hectacres}/^\circ\text{F} \bullet 59^\circ\text{F} \\ &= -69.78 \text{ log-hectacres} + 1.39 \text{ log-hectacres}/^\circ\text{F} \bullet 59^\circ\text{F} \\ &= 12.23 \text{ log-hectares}\end{aligned}$$

We did not observe any summers at exactly  $x = 59$  but did observe some nearby and this result seems relatively reasonable.

Now suppose someone asks you to use this equation for predicting Temperature = 65°F. We can run that through the equation:  $-69.78 + 1.39 * 65 = 20.57$  log-hectares but can we trust this prediction? We did not observe any summers over 60 degrees F so we are now predicting outside the scope of our observations – performing extrapolation. Having a scatterplot in hand helps us to assess the range of values where we can reasonably use the equation – between 54 and 60 degrees F.

```
scatterplot(loghectacres~Temperature, data=mtfires, smoother=T,
            main="Scatterplot with regression line for Area burned vs Temperature")
```

## 6.8 Measuring the strength of regressions: $R^2$

At the beginning of the chapter, we used the correlation coefficient to measure the strength and direction of the linear relationship. The regression line provides an even more detailed description of the direction of the linear relationship than the correlation provided; in regression we addressed the question of “for a unit change in  $x$ , what sort of change in  $y$  do we expect, on average?” whereas the correlation just addressed whether the relationship was positive or negative. However, the **regression line tells us nothing about the strength of the relationship**. Consider the three scatterplots in Figure 6.19: the left panel is the original *BAC* data and the two right panels have fake data that generated the same estimated regression model with a weaker (middle panel) and then a stronger (right panel) linear relationship between *Beers* and *BAC*. This suggests that the regression line is a useful but incomplete characterization of relationships between variables – we need a measure of strength of the relationship to go with the equation.

We could use the correlation coefficient,  $r$ , again to characterize strength but it is somewhat redundant to report a measure that contains direction information. It also will not extend to multiple regression models where we have more than one predictor variable in the same model.

In regression models, we use the **coefficient of determination** (symbol:  $\mathbf{R^2}$ ) to accompany our regression line and describe the strength of the relationship. It can either be scaled between 0 and 1 or 0 to 100% and has “units” of the proportion or percentage of the variation in  $y$  that is explained by the model that includes  $x$  (and later more than one  $x$ ). For example, and  $\mathbf{R^2}$  of 0% corresponds to explaining 0% of the variation in

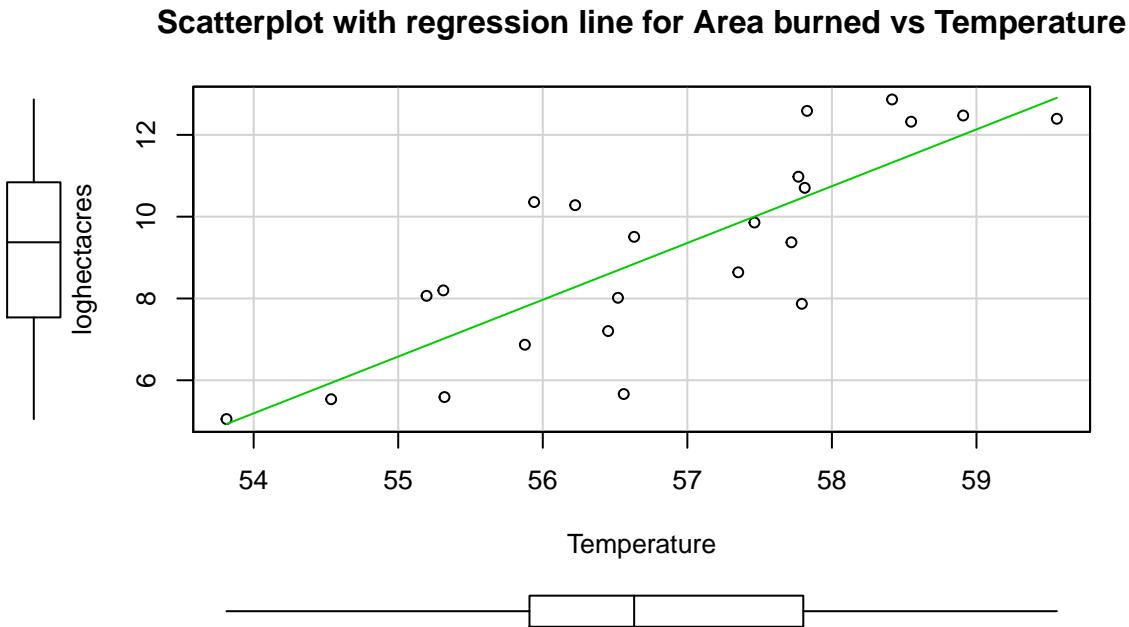


Figure 6.18: Scatterplot of log-hectares burned versus temperature with estimated regression line.

the response with our model and  $R^2 = 100\%$  means that all the variation in the response was explained by the model. In between, it provides a nice summary of how much of the total variability in the response we can account for with our model including  $x$  (and, in Chapter 8, including multiple predictor variables).

$R^2$  is calculated using the sums of squares we encountered in the ANOVA methods. We once again have some total amount of variability that is attributed to the variation based on the model fit, here we call it  $SS_{\text{regression}}$ , and the residual variability, still  $SS_{\text{error}} = \sum(y - \hat{y})^2$ . The  $SS_{\text{regression}}$  is most easily calculated as  $SS_{\text{regression}} = SS_{\text{Total}} - SS_{\text{error}}$ , the difference between the total variability and the variability not explained by the model under consideration. Using these quantities, we calculate the portion of the total variability that the model explains as

$$R^2 = \frac{SS_{\text{regression}}}{SS_{\text{Total}}} = 1 - \frac{SS_{\text{error}}}{SS_{\text{Total}}}.$$

It also ends up that the coefficient of determination for models with one predictor is the correlation coefficient ( $r$ ) squared ( $R^2 = r^2$ ). So we can quickly find coefficients of determination if we know correlations in simple linear regression models. In the real *Beers* and *BAC* data,  $r=0.8943$ . So  $R^2 = 0.79998$  or approximately 0.80. So 80% of the variation in *BAC* is explained by *Beer* consumption. That leaves 20% of the variation in the responses to be unexplained by our model. In this case much of the unexplained variation is likely attributable to differences in physical characteristics (that were not measured) but the statistical model places that unexplained variation into the category of “random errors”. We don’t actually have to find  $r$  to get coefficients of determination – the result is part of the regular summary of a regression model that we have not been reproducing. The full `lm` model summary follows with “Multiple R-squared” that is bolded below. It is reported as a proportion and it is your choice whether you want to report and interpret it as a proportion or percentage, just make that clear in how you discuss it.

```
m1<-lm(BAC~Beers,data=BB)
```

```
summary(m1)
```

```
##  
## Call:
```

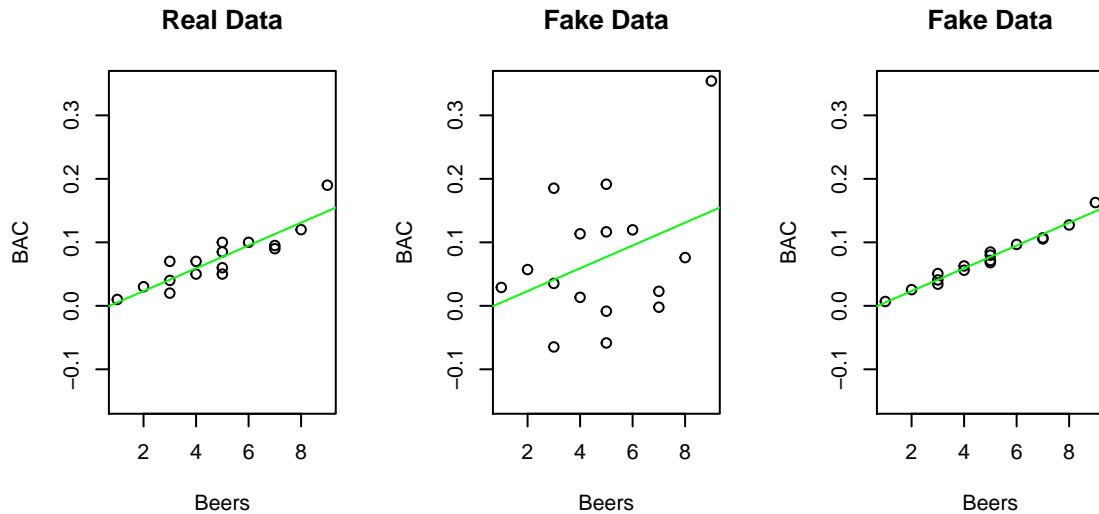


Figure 6.19: Three scatterplots with the same estimated regression line.

```
## lm(formula = BAC ~ Beers, data = BB)
##
## Residuals:
##   Min     1Q Median     3Q    Max 
## -0.027118 -0.017350  0.001773  0.008623  0.041027
## 
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) -0.012701  0.012638  -1.005   0.332    
## Beers        0.017964  0.002402   7.480 2.97e-06 ***
## 
## Residual standard error: 0.02044 on 14 degrees of freedom
## Multiple R-squared:  0.7998, Adjusted R-squared:  0.7855 
## F-statistic: 55.94 on 1 and 14 DF,  p-value: 2.969e-06
```

In this output, be careful because there is another related quantity called ***Adjusted R-squared*** that we will discuss later. This other quantity is not a measure of the strength of the relationship but will be useful.

We could also revisit the ANOVA table for this model to verify the source of the  $\mathbf{R}^2$  of 0.80 based on  $SS_{\text{regression}} = 0.02337$  and  $SS_{\text{Total}} = 0.02337 + 0.00585$ . This provides 0.80 from  $0.02337/0.02922$ .

```
anova(m1)
```

```
## Analysis of Variance Table
##
## Response: BAC
##            Df  Sum Sq Mean Sq F value Pr(>F)    
## Beers      1 0.0233753 0.0233753 55.944 2.969e-06
## Residuals 14 0.0058497 0.0004178
## SStotal <- 0.0233753 + 0.0058497
## SSregression <- 0.0233753
## SSregression/SStotal

## [1] 0.7998392
```

In Figure 6.19, we saw three examples with the same regression model, but different strengths of relationships. In the real data set  $\mathbf{R}^2 = 80\%$ . For the first fake data set (middle panel), the  $\mathbf{R}^2$  drops to 13.8% and for the

second fake data set (right panel),  $\mathbf{R}^2$  is 97.3%. As a summary,  $\mathbf{R}^2$  provides a natural scale to understand “how good” each model is at explaining the responses. We can revisit some of our previous models to get a little more practice with using this summary of strength or quality of regression models.

For the Montana fire data,  $\mathbf{R}^2 = 66.2\%$ . So the proportion of variation of log-area burned that is explained by average summer temperature is 0.662. This is “good” but also leaves quite a bit of unexplained variation in the responses. There is a long list of reasons why this explanatory variable leaves a lot of variation in the response unexplained. Note that we were careful about using the scaling of the response variable (log(area burned)) in the interpretation – this is because we would get a much different answer if area burned vs temperature was considered.

```
fire1 <- lm(loghectacres~Temperature, data=mtfires)
summary(fire1)
```

```
##
## Call:
## lm(formula = loghectacres ~ Temperature, data = mtfires)
##
## Residuals:
##      Min       1Q   Median       3Q      Max
## -3.0822 -0.9549  0.1210  1.0007  2.4728
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) -69.7845    12.3132 -5.667 1.26e-05
## Temperature  1.3884     0.2165  6.412 2.35e-06
##
## Residual standard error: 1.476 on 21 degrees of freedom
## Multiple R-squared:  0.6619, Adjusted R-squared:  0.6458
## F-statistic: 41.12 on 1 and 21 DF,  p-value: 2.347e-06
```

For the model for female Australian athletes that used *Body fat* to explain *Hematocrit*, the estimated regression model was  $\widehat{Hc}_i = 42.014 - 0.085\text{BodyFat}_i$  and  $r = -0.168$ . The coefficient of determination is  $\mathbf{R}^2 = (-0.168)^2 = 0.0282$ . So *body fat* explains 2.8% of the variation in *Hematocrit* in these women. That is not a very good regression model with over 97% of the variation in *Hematocrit* unexplained by this model. The scatterplot showed a fairly weak relationship but this provides numerical and interpretable information that drives that point home.

```
m2 <- lm(Hc~Bfat, data=aisR2[aisR2$Sex==1,]) #Results for Females
summary(m2)
```

```
##
## Call:
## lm(formula = Hc ~ Bfat, data = aisR2[aisR2$Sex == 1, ])
##
## Residuals:
##      Min       1Q   Median       3Q      Max
## -5.2399 -2.2132 -0.1061  1.8917  6.6453
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) 42.01378    0.93269 45.046   <2e-16
## Bfat        -0.08504    0.05067 -1.678    0.0965
##
## Residual standard error: 2.598 on 97 degrees of freedom
## Multiple R-squared:  0.02822, Adjusted R-squared:  0.0182
## F-statistic: 2.816 on 1 and 97 DF,  p-value: 0.09653
```

## 6.9 Outliers: leverage and influence

In the review of correlation, we loosely considered the impacts of outliers on the correlation. We removed unusual points to see both the visual changes (in the scatterplot) as well as changes in the correlation coefficient in Figures 6.4 and 6.5. In this section, we formalize these ideas in the context of impacts of unusual points on our regression equation. In regression, it is possible for a single point to have a big impact on the overall regression results but it is also possible to have a clear outlier that has little impact on the results. We call an observation *influential* if its removal causes a “big” change in the regression line, specifically in terms of impacting the slope coefficient. Points that are on the edges of the  $x$ ’s (far from the mean of the  $x$ ’s) have the potential for more impact on the line as we will see in some examples shortly.

You can think of the regression line being balanced at  $\bar{x}$  and the further from that location a point is, the more a single point can move the line. We can measure the distance of points from  $\bar{x}$  to quantify each observation’s potential for impact on the line using what is called the **leverage** of a point. Leverage is a positive numerical measure with larger values corresponding to more leverage. The scale changes depending on the sample size ( $n$ ) and the complexity of the model so all that matters is which observations have more or less relative leverage in a particular data set. The observations with  $x$ -values that provide higher leverage have increased potential to influence the estimated regression line. Along with measuring the leverage, we can also measure the influence that each point has on the regression line using **Cook’s Distance** or **Cook’s D**. It also is a positive measure with higher values suggesting more influence. The rule of thumb is that Cook’s D values over 1.0 correspond to clearly influential points, values over 0.5 have some influence and values lower than 0.5 indicate points that are not influential on the regression model slope coefficients. One part of the regular diagnostic plots we will use for regression models displays the leverages on the  $x$ -axis, the standardized residuals on the  $y$ -axis, and adds contour lines for Cook’s Distances in a panel that is labeled “Residuals vs Leverage”. This allows us to see the potential for impact of a point (leverage), how far it’s observation was from the regression line (residual), and to see a measure of that point’s influence (Cook’s D).

To extract the level of Cook’s D on the “Residuals vs Leverage” plot, look for contours to show up on the upper and lower right of the plot. They show increasing levels of influence going to the upper and lower right corners as you combine higher leverage ( $x$ -axis) and larger residuals ( $y$ -axis) – the two ingredients required to be influential on the line. The contours are displayed for Cook’s D values of 0.5 and 1.0 if there are points near or over those levels. The Cook’s D values come from a topographical surface of values that is a sort of U-shaped valley in the middle of the plot centered at  $y = 0$  with the lowest contour corresponding to Cook’s D values below 0.5 (no influence). As you move to the upper right or lower right corners, the influence increases and the edges of the valley get steeper. If you do not see any contours in the plot, then no points were even close to being influential based on Cook’s D.

To illustrate these concepts, the original *Beers* and *BAC* data are used again. In the scatter plot in Figure 6.20, two points are plotted with different characters. The point for 1 *Beer* and *BAC* of 0.010 is displayed as a “◊” and the 9 *Beer* and *BAC* 0.19 observation is displayed with a “○”. These two points are the furthest from the mean of the  $x$ ’s ( $\text{Beers} = 4.8$ ) but show two different levels of influence on the line.

The “◊” point has a leverage of 0.27 and the 9 *Beer* observation (“○”) had a leverage of 0.30. The 1 *Beer* observation was close to the pattern defined by the other points, had a small residual, and a Cook’s D value below 0.5 (it did not exceed the first of the contours). So even though it had high leverage, it was not an influential point. The 9 *Beer* observation had the highest leverage in the data set and was quite a bit above the pattern defined by the other points and ends up being an influential point with a Cook’s D over 1. We might want to consider fitting this model without that observation to get a better estimate of the effects of beer consumption on BAC or revisit our assumption that the relationship is really linear here.

To further explore influence, we will add a point to the original data set and move it around so you can see how those changes impact the results. For each scatterplot in Figure 6.21, the Residuals vs Leverage plot is displayed to its right. The original data are “●” and the original regression line is the dashed line in Figure 6.21. First, a fake observation at 11 *Beers* and 0.1 *BAC* is added, at (11, 0.1), in the top panels of the figure. This observation is clearly an outlier and heavily impacts the slope of the regression line (so is clearly influential). This added point drops the  $R^2$  from 0.80 in the original data to 0.24. The accompanying

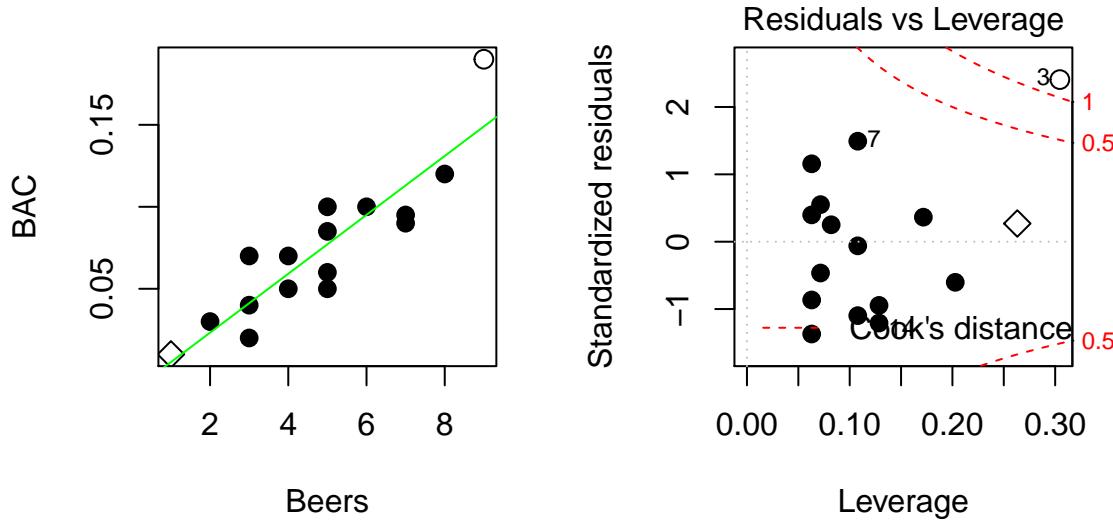


Figure 6.20: Scatterplot and Residuals vs Leverage plot for the real BAC data.

Residuals vs Leverage plot shows that this point has extremely high leverage and a Cook's D over 1 – it is a clearly influential point. However, **having high leverage does not always make points influential**. Consider the second row of plots with an added point of (11, 0.19). The regression line barely changes and  $R^2$  increases a little. This point has the same leverage as in the first example since it is the same set of  $x$ 's and the distance to the mean is unchanged. But it is not influential since its Cook's D is less than 0.5. This occurred because it followed the overall pattern of observations even though it was “far away” from the other observations in the x-direction. The last two rows of plots show what happens when low leverage outliers are encountered. Placing observations near the center of the  $x$ 's means that to be influential the points have to be very far from the pattern of the other observations. The (5, 0.19) example almost attains a Cook's D of 0.5 but has little impact on the regression line, especially the slope coefficient. It does impact the y-intercept and drops the R-squared value to 0.57. The same result occurs if the observation is noticeably lower than the other points.

When we are doing regressions, we get very worried about points “at the edges” having an undue influence on the results. When we start using multiple predictors, say if we had body weight data on these subjects as well as beer consumption, it becomes harder to “see” if the points are “far away” from the other observations and we will trust the Residuals vs Leverage plots to help us identify the influential points. These techniques work the same in the multiple regression models in Chapter 8 as they do in these simpler, single predictor regression models.

## 6.10 Residual diagnostics – setting the stage for inference

Influential points are not the only potential issue that can cause us to have concerns about our regression model. There are two levels to these considerations. The first is related to issues that directly impact the least squares regression line and cause concerns about whether a line is a reasonable representation of the relationship between the two variables. These issues for regression model estimation have been discussed previously (the same concerns in estimating correlation apply to regression models). The second level is whether the line we have will be useful for making inferences for the population that our data were collected

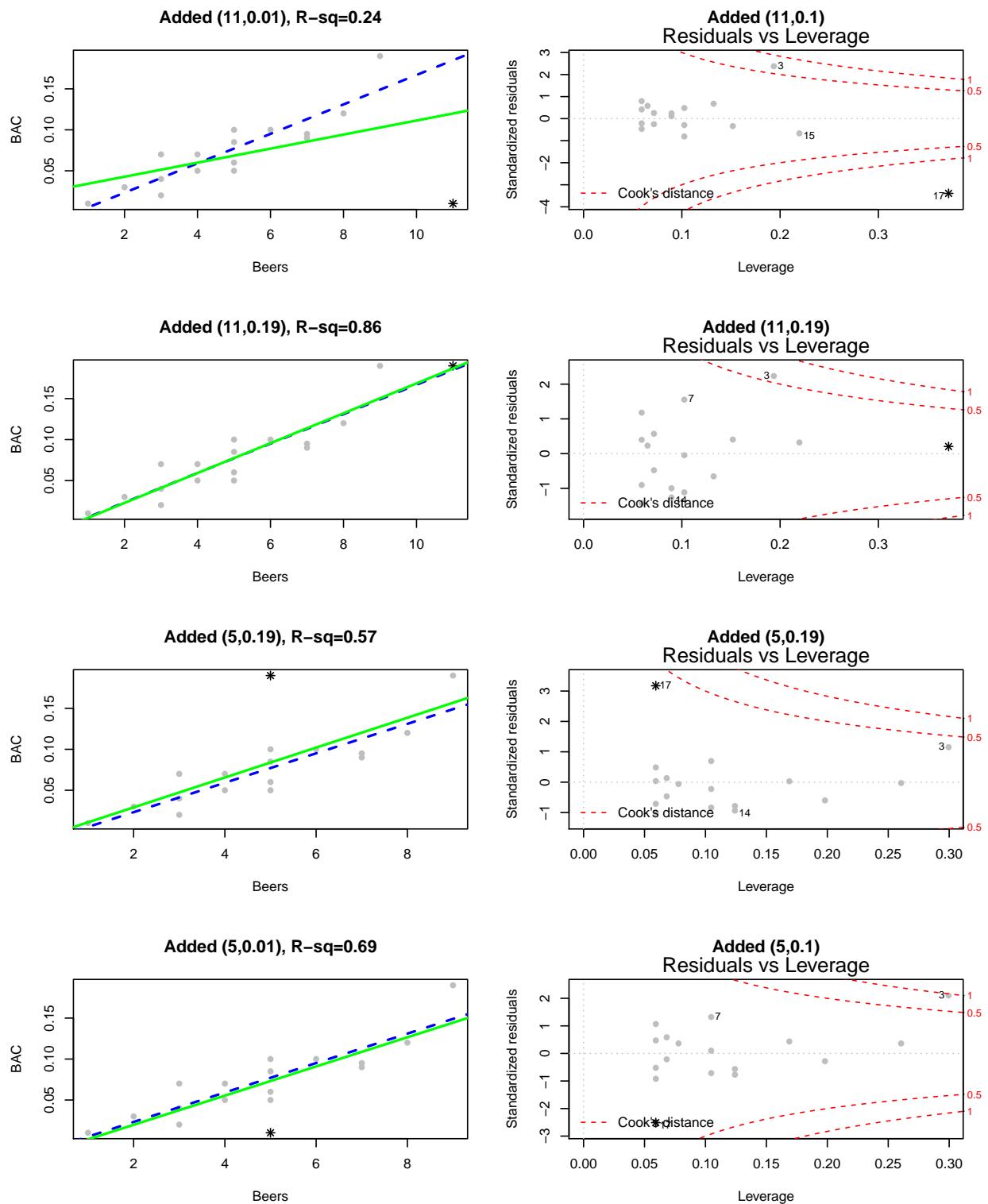


Figure 6.21: Plots exploring the impacts of moving a single additional observation.

from and whether the data follow our assumed model. Our window into problems of both types is the residuals ( $e_i = y_i - \hat{y}_i$ ). By exploring patterns in how the line “misses” the responses we can gain much information about the reasonableness of using the estimated regression line and sometimes information about how we might fix problems. The validity conditions for doing inference in a regression setting (Chapter 7) involve two sets of considerations, those that are assessed based on the data collection and measurement process and those that can be assessed using diagnostic plots. The first set is:

- **Quantitative variables condition**

- We’ll discuss using categorical predictor variables later – to use simple linear regression both the explanatory and response variables need to quantitative.

- **Independence of observations**

- As in the ANOVA models, linear regression models assume that the observations are collected in a fashion that makes them independent.
- This will be based on the “story” of the data. Consult a statistician if your data violate this assumption as there are more advanced methods that adjust for dependency in observations but they are beyond the scope of this material.

The remaining assumptions for getting valid inferences from regression models can be assessed using diagnostic plots.

- **Linearity of relationship**

- We should not report a linear regression model if the data show a curve (curvilinear relationship between  $x$  and  $y$ ).
- Examine the Residuals vs Fitted plot:
  - If the model missed a curve in the relationship, the residuals often will highlight that missed pattern and a curve will show up in this plot.
  - Try to explain or understand the pattern in what is left over. If we have a good model, there shouldn’t be much left to “explain” in the residuals (i.e., no patterns left over after accounting for  $x$ ).

- **Equal (constant) variance**

- We assume that the variation is the same for all the observations and especially that the variability does not change in the responses as a function of our predictor variables or the fitted values.
- There are two plots to help with this:
  - Examine the Residuals vs Fitted plot and look for evidence of changing spread in the residuals, being careful to try to separate curving patterns from non-constant variance (and look for situations where both are present as you can violate both conditions simultaneously).
  - Examine the “Scale-Location” plot and look for changing spread as a function of the fitted values.
    - The y-axis in this plot is the square-root of the absolute value of the standardized residual. This scale flips the negative residuals on top of the positive ones to help you better assess changing variability without being distracted by whether the residuals are above or below 0.
    - Because of the absolute value, curves in the Residuals vs Fitted plot can present as sort of looking like non-constant variance – check for nonlinearity in the residuals vs fitted before using this plot. If nonlinearity is present, just use the Residuals vs Fitted for assessing constant variance around curving pattern.

- If there are patterns of increasing or decreasing variation (often described as funnel or cone shapes), then it might be possible to use a transformation to fix this problem (more later). It is possible to have decreasing and then increasing variability in one plot and this also is a violation of this condition.

- **Normality of residuals**

- Examine the Normal QQ plot for violations of the normality assumption as in Chapters 3 and 4.
  - Specifically review the discussion of identifying skews in different directions and heavy vs light tailed distributions.
  - Skewed and heavy-tailed distributions are the main problems for our inferences, especially since both kinds of distributions can contain outliers that can wreak havoc on the estimated regression line.
  - Light-tailed distributions cause us no real inference issues except that the results are conservative so you should note when you observe these situations but feel free to proceed with using your model results.
  - Remember that clear outliers are an example of a violation of the normality assumption but some outliers may just influence the regression line and make it fit poorly for the results of the observations and this issue will be more clearly observed in the residuals vs fitted.

- **No influential points**

- Examine the Residuals vs Leverage plot as discussed in the previous section.
- Consider removing influential points and focusing on results without that point in the data set.

To assess these later assumptions, we will use the four residual diagnostic plots that R provides from `lm` fitted models. They are similar to the results from ANOVA models but the Residuals vs Leverage plot is now interesting as was discussed in Section 6.9. Now we can fully assess the potential for trusting the estimated regression models in a couple of our examples:

- **Beers vs BAC:**

- Quantitative variables condition:
  - Both variables are quantitative.
- Independence of observations:
  - We can assume that all the subjects are independent of each other. There is only one measurement per student and it is unlikely that one subject's beer consumption would impact another's BAC. Unless the students were trading blood it is isn't possible for one person's beer consumption to change someone else's BAC.

```
m1 <- lm(BAC~Beers, data=BB)
par(mfrow=c(2,2))
plot(m1, add.smooth=F, main="Beers vs BAC")
```

- Linearity, constant variance from Residuals vs Fitted:
  - We previously have identified a potentially influential outlier point in these data. Consulting the Residuals vs Fitted plot in Figure 6.22, if you trust that influential point, shows some curvature with a pattern of decreasing residuals and then an increase at the right. Or, if you do not trust that highest BAC observation, then there is a mostly linear relationship with an outlier identified. We would probably suggest that it is an outlier, should be removed from the analysis, and inferences constrained to the region of beer consumption from 1 to 8 beers since we don't know what might happen at higher values.
- Constant variance from Scale-Location:

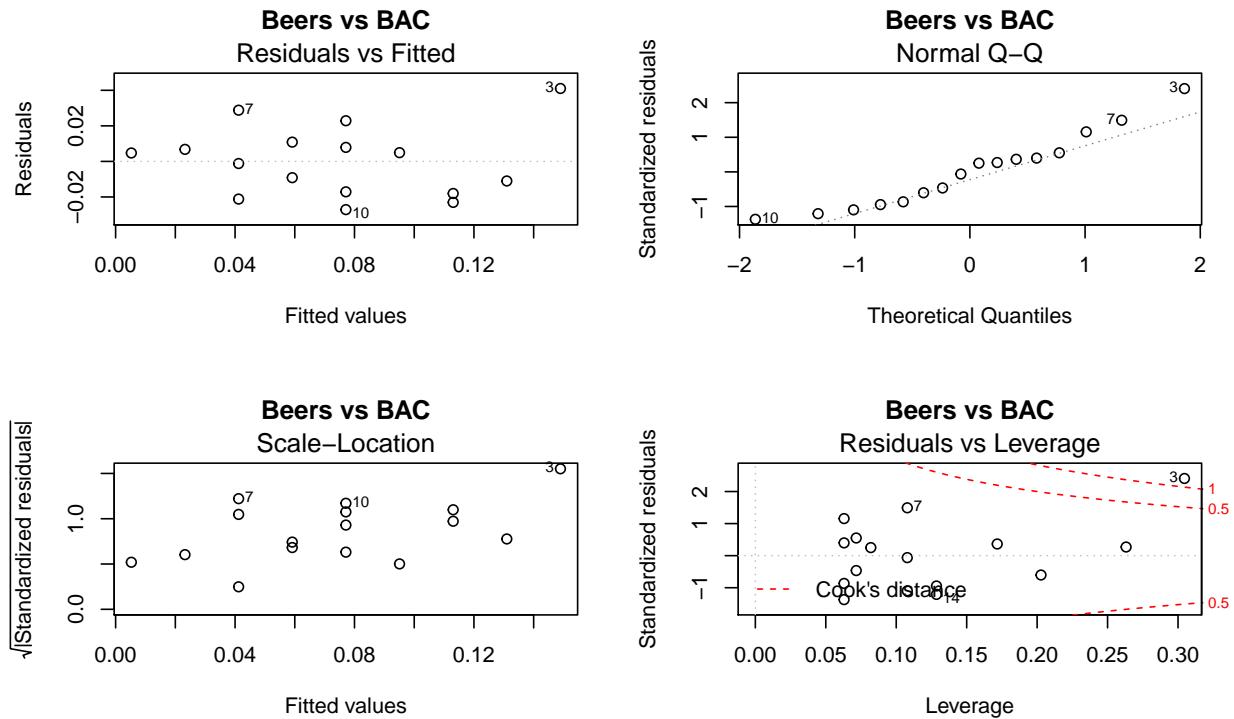


Figure 6.22: Full suite of diagnostics plots for *Beer vs BAC* data.

- There is some evidence of increasing variability in this plot as the spread of the results increases from left to right, however this is just an artifact of the pattern in the original residuals and not real evidence of non-constant variance.
- Normality from Normal QQ Plot:
  - The left tail is a little short and the right tail is a little long, suggesting a right skewed distribution in the residuals. This corresponds to having a large positive outlying value.
- Influential points from Residuals vs Leverage:
  - Previously discussed, this plot shows one influential point with a Cook's D value over 1 that is distorting the fitted model.
- **Tree height and tree diameter** (suspicious observation already removed):
  - Quantitative variables: Met
  - Independence of observations:
    - There are multiple trees that were measured in each plot. One problem might be that once a tree is established in an area, the other trees might not grow as tall. The other problem is that some sites might have better soil conditions than others. Then, all the trees in those rich soil areas might be systematically taller than the trees in other areas. Again, there are statistical methods to account for this sort of “clustering” of measurements but this technically violates the assumption that all the trees are independent of each other. So this assumption is violated, but we will proceed with that caveat on our results – the precision of our inferences might be slightly over-stated due to some potential dependency in the measurements.
  - Linearity, constant variance from Residuals vs Fitted in Figure 6.23.

- There is evidence of a curve that was missed by the linear model.
- There is also evidence of increasing variability AROUND the curve in the residuals.
- Constant variance from Scale-Location:
  - This plot actually shows relatively constant variance but this plot is misleading when curves are present in the data set. **Focus on the Residuals vs Fitted to diagnose non-constant variance in situations where a curve was missed.**
- Normality in Normal QQ plot:
  - There is no indication of any problem with the normality assumption.
- Influential points?
  - The Cook's D contours do not show up in this plot so none of the points are influential.

So the main issues with this model are the curving relationship and non-constant variance. We'll revisit this example later to see if we can find a model on transformed variables that has better diagnostics.

Reporting the following regression model that has a decent  $R^2$  of 62.6% would be misleading since it does not accurately represent the relationship between tree diameter and tree height.

```
tree1 <- lm(height.m ~ dbh.cm, data=ufc[-168,])
summary(tree1)
```

```
##
## Call:
## lm(formula = height.m ~ dbh.cm, data = ufc[-168, ])
##
## Residuals:
##    Min      1Q  Median      3Q     Max
## -12.1333 -3.1154  0.0711  2.7548 12.3076
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) 11.98364   0.57422  20.87  <2e-16
## dbh.cm      0.32939   0.01395   23.61  <2e-16
##
## Residual standard error: 4.413 on 333 degrees of freedom
## Multiple R-squared:  0.626, Adjusted R-squared:  0.6249
## F-statistic: 557.4 on 1 and 333 DF, p-value: < 2.2e-16
par(mfrow=c(2,2))
plot(tree1, add.smooth=F)
```

## 6.11 Old Faithful discharge and waiting times

A study in August 1985 considered time for Old Faithful and how that might relate to *waiting time* for the next eruption (Ripley (2017), Azzalini and Bowman (1990)). This sort of research provides the park staff a way to show tourists a predicted time to next eruption so they can quickly see it and then get back in their cars. Both variables are measured in minutes and the scatterplot in Figure 6.24 shows a moderate to strong positive and relatively linear relationship. We added a *smoothing line* (red) to this plot. Smoothing lines provide regression-like fits but are performed on local areas of the relationship and so can highlight where the relationships change and can highlight curvilinear relationships. They can also return straight lines just like the regression line if that is reasonable. The technical details of regression smoothing are not covered here but they are a useful graphical addition to help visualize nonlinearity in relationships.

In these data, there appear to be two groups of eruptions (shorter length, shorter wait and longer length, longer wait) – but we don't know enough about these data to assume that there are two groups. The

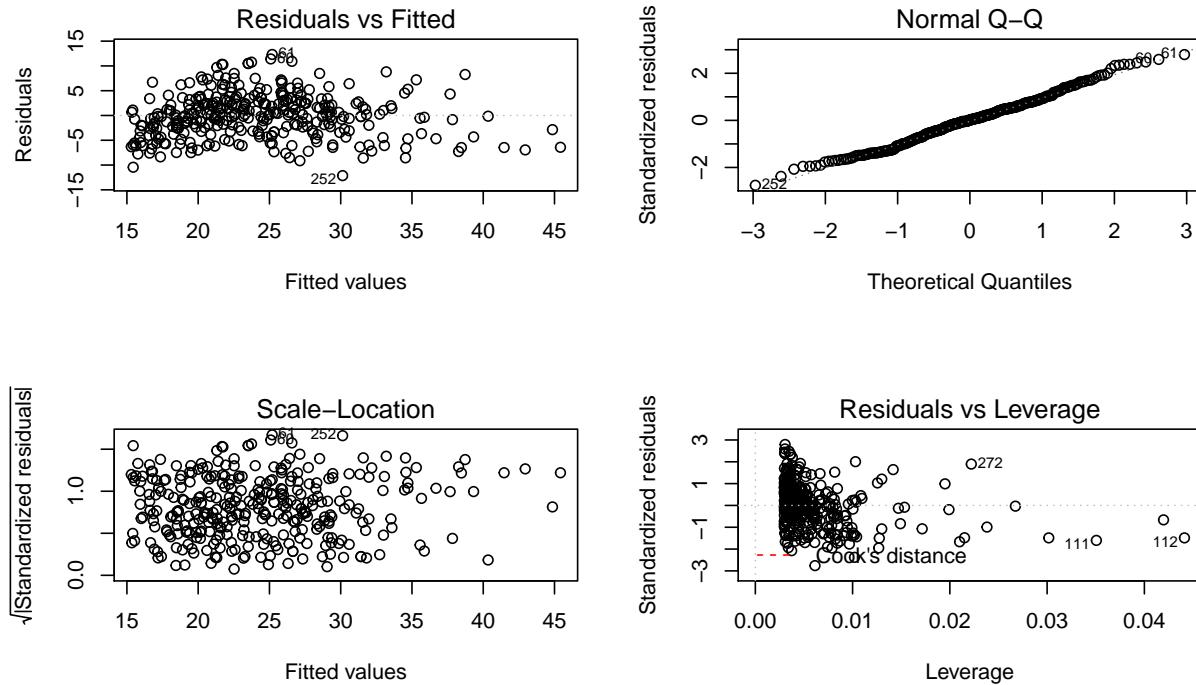


Figure 6.23: Diagnostics plots for tree height and diameter simple linear regression model.

smoothing line does help us to see if the relationship appears to change or stay the same across different values of the explanatory variable, `Duration`. The smoothing line suggests that the upper group might have a less steep slope than the lower group as it sort of levels off for observations with `Duration` of over 4 minutes. It also indicates that there is one point for an eruption under 1 minute in `Duration` that might be causing some problems. The story of these data involve some measurements during the night that were just noted as being short, medium, and long – and they were re-coded as 2, 3, or 4 minute duration eruptions. We'll see if our diagnostics detect some of these issues when we fit a simple linear regression to try to explain waiting time based on duration of prior eruption.

```
require(MASS)
data(geyser)
G2 <- data.frame(Waiting=geyser$waiting[-1],
                  Duration=geyser$duration[-299])
scatterplot(Waiting~Duration, data=G2, spread=F)
```

The first concern with these data is that the observations are likely not independent. Since they were taken consecutively, one waiting time might be related to the next waiting time – violating the independence assumption. As noted above, there might be two groups (types) of eruptions – short ones and long ones. The Normal QQ-Plot in Figure 6.25 also suggests a few observations creating a slightly long right tail. Those observations might warrant further exploration as they also show up as unusual in the Residuals vs Fitted plot. There are no highly influential points in the data set with all points having Cook's D smaller than 0.5, so these outliers are not necessarily moving the regression line around. There are two distinct groups of observations but the variability is not clearly changing so we do not have to worry about non-constant variance here. So these results might be relatively trustworthy if we assume that the same relationship holds for all levels of duration of eruptions.

```
OF1 <- lm(Waiting~Duration, data=G2)
summary(OF1)
```

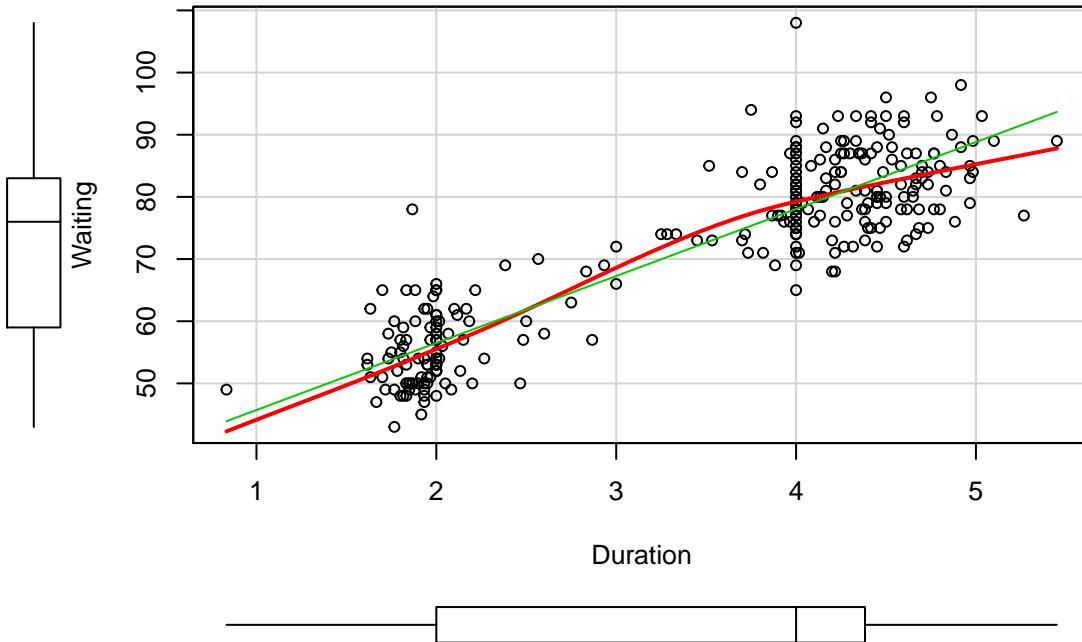


Figure 6.24: Scatterplot of Old Faithful waiting times to next eruption (minutes) and duration of prior eruption (minutes) with smoothing line (bold, red) and regression line (thin, green).

```
##  
## Call:  
## lm(formula = Waiting ~ Duration, data = G2)  
##  
## Residuals:  
##      Min       1Q   Median       3Q      Max  
## -14.6940  -4.4954  -0.0966  3.9544  29.9544  
##  
## Coefficients:  
##             Estimate Std. Error t value Pr(>|t|)  
## (Intercept) 34.9452    1.1807  29.60 <2e-16  
## Duration    10.7751    0.3235  33.31 <2e-16  
##  
## Residual standard error: 6.392 on 296 degrees of freedom  
## Multiple R-squared:  0.7894, Adjusted R-squared:  0.7887  
## F-statistic: 1110 on 1 and 296 DF, p-value: < 2.2e-16  
par(mfrow=c(2,2))  
plot(OF1)
```

The estimated regression equation is  $\widehat{\text{WaitingTime}}_i = 34.95 + 10.78 \text{ Duration}_i$ , suggesting that for a 1 minute increase in eruption Duration we would expect, on average, a 10.78 minute change in the WaitingTime. This equation might provide a useful tool for the YNP staff to predict waiting times. The  $R^2$  is fairly large: 78.9% of the variation in waiting time is explained by the duration of the previous eruption. But maybe this is more about two types of eruptions/waiting times? We could consider the relationship within the shorter and longer eruptions but since there are observations residing between the two groups, it is difficult to know where to split the explanatory variable into two groups. Maybe we really need to measure additional information that might explain why there are two groups in the responses...

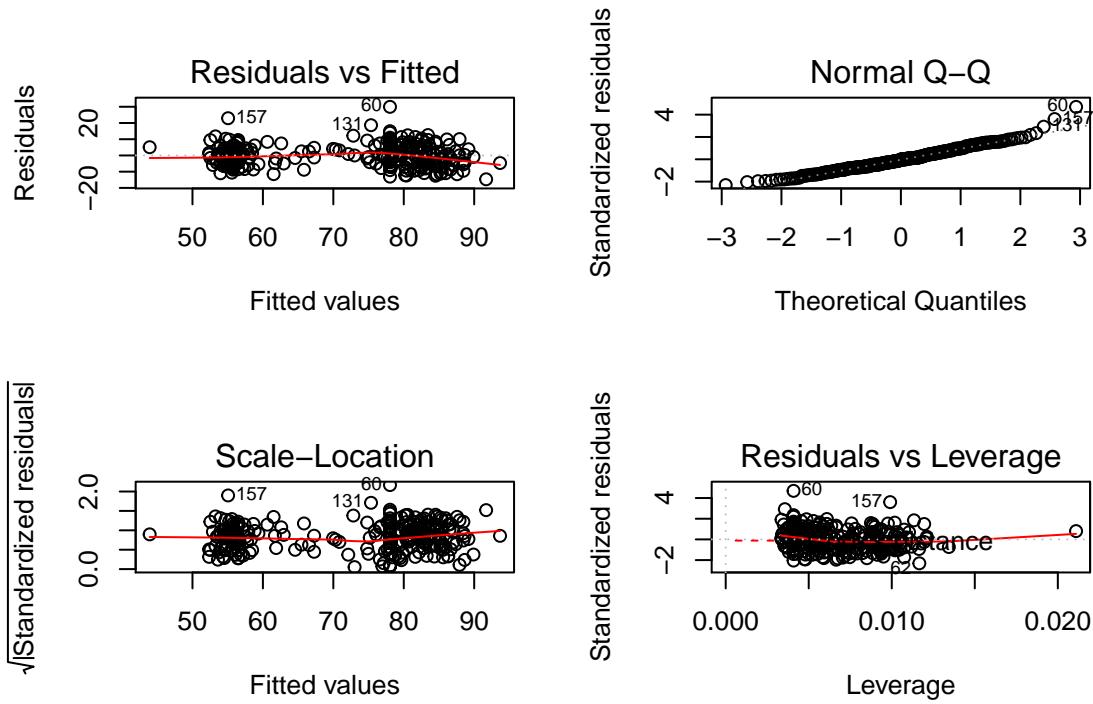


Figure 6.25: Diagnostic plots for Old Faithful waiting time model.

## 6.12 Chapter summary

The Pearson's Product Moment Correlation Coefficient) measures the strength and direction of the linear relationship between two quantitative variables. Regression models estimate the impacts of changes in  $x$  on the mean of the response variable  $y$ . Direction of the assumed relationship (which variable explains or causes the other) matters for regression models but does not matter for correlation. Regression lines only describe the pattern of the relationship; in regression, we use the coefficient of determination to describe the strength of the relationship between the variables as a percentage of the response variable that is explained by the model. If we are choosing between models, we prefer them to have higher  $R^2$  values for obvious reasons, but we will discover in Chapter 8 that maximizing the coefficient of determination is not a good way to pick a model when we have multiple candidate options.

In this chapter, a wide variety of potential problems were explored when using regression models. This included a discussion of the conditions that will be required for using the models to perform trustworthy inferences in the next two chapters. It is important to remember that correlation and regression models only measure the **linear** association between variables and that can be misleading if a nonlinear relationship is present. Similarly, influential observations can completely distort the apparent relationship between variables and should be checked for before trusting any regression output. It is also important to remember that regression lines should not be used outside the scope of the original observations – extrapolation should be checked for and avoided whenever possible or at least acknowledged.

Regression models look like they estimate the changes in  $y$  that are caused by changes in  $x$ . This is not true unless the levels of  $x$  are randomly assigned and only then we can make causal inferences. Since this is not generally true, you should initially always assume that any regression equation describes the relationship – if you observe two subjects that are 1 unit of  $x$  apart, you can expect their mean to differ by  $b_1$  – you should not, however, say that changing  $x$  causes a change in the mean of the responses. Despite all these cautions, regression models are very popular statistical methods. They provide detailed descriptions of relationships

between variables and can be extended to situations where we are interested in multiple predictor variables. They also share ties to the ANOVA models from earlier this semester. When you are running R code, you will note that all the ANOVAs and the regression models are all fit using `lm`. The assumptions and diagnostic plots are quite similar. And in the next chapter, we will see that inference techniques look similar. People still like to distinguish among the different types of situations, but the underlying *linear models* are actually exactly the same...

## 6.13 Important R code

The main components of the R code used in this chapter follow with the components to modify in red where  $y$  is a response variable,  $x$  is an explanatory variable, and the data are in `DATASETNAME`.

- `pairs.panels(DATASETNAME, ellipses=F, scale=T, smooth=F, col=0)`
  - Requires the `psych` package.
  - Makes a scatterplot matrix that also displays the correlation coefficient.
- `cor(y~x, data=DATASETNAME)`
  - Provides the estimated correlation coefficient between  $x$  and  $y$ .
- `plot(y~x, data=DATASETNAME)`
  - Provides a scatter plot.
- `scatterplot(y~x, data=DATASETNAME, smooth=F)`
  - Requires the `car` package.
  - Provides a scatter plot with a regression line.
- `MODELNAME <- lm(y~x, data=DATASETNAME)`
  - Estimates a regression model using least squares.
- `summary(MODELNAME)`
  - Provides parameter estimates and R-squared (used heavily in Chapter 7 and 8 as well).
- `par(mfrow=c(2, 2)); plot(MODELNAME)`
  - Provides four regression diagnostic plots in one plot.

## 6.14 Practice problems

These questions revisit the treadmill data set from Chapter 1. Researchers were interested in whether the run test variable could be used to replace the treadmill oxygen consumption variable that is expensive to measure. The following code loads the data set and provides a scatterplot matrix using `pairs.panels`.

```
treadmill <- read.csv("http://www.math.montana.edu/courses/s217/documents/treadmill.csv")
require(psych)
pairs.panels(treadmill, ellipses=F, smooth=F, col=0)
```

6.1. First, we should get a sense of the strength of the correlation between the variable of primary interest, `TreadMillOx`, and the other variables and consider whether outliers or nonlinearity are going to be major issues here. Which variable is it most strongly correlated with? Which variables are next most strongly correlated with this variable?

6.2. Fit the SLR using `RunTime` as explanatory variable for `TreadMillOx`. Report the estimated model.

6.3. Predict the treadmill oxygen value for a subject with a run time of 14 minutes. Repeat for a subject with a run time of 16 minutes. Is there something different about these two predictions?

6.4. Interpret the slope coefficient from the estimated model, remembering the units on the variables.

6.5. Report and interpret the y-intercept from the SLR.

6.6. Report the  $R^2$  value from the output. Show how you can find this value from the original correlation matrix result.

6.7. Produce the diagnostic plots and discuss any potential issues. What is the approximate leverage of the highest leverage observation and how large is its Cook's D? What does that tell you about its potential influence in this model?

# Chapter 7

## Simple linear regression inference

### 7.1 Model

In Chapter 6, we learned how to estimate and interpret correlations and regression equations with a single predictor variable (*simple linear regression* or SLR). We carefully explored the variety of things that could go wrong and how to check for problems in regression situations. In this chapter, that work provides the basis of performing statistical inference that mainly focuses on the population slope coefficient based on the sample slope coefficient. As a reminder, the estimated regression model is  $\hat{y}_i = b_0 + b_1 x_i$ . the population regression equation is  $y_i = \beta_0 + \beta_1 x_i + \varepsilon_i$ . Where  $\beta_0$  is the **population** (or true) **y-intercept** and  $\beta_1$  is the **population** (or true) **slope coefficient**. These are population parameters (fixed but typically unknown). This model can be re-written to think about different components and their roles. The mean of a random variable is statistically denoted as  $E(y_i)$ , the **expected value of  $y_i$** , or as  $\mu_{y_i}$  and the mean of the response variable in a simple linear model is specified by  $E(y_i) = \mu_{y_i} = \beta_0 + \beta_1 x_i$ . This uses the true regression line to define the model for the mean of the responses as a function of the value of the explanatory variable.

The other part of any statistical model is specifying a model for the variability around the mean. There are two aspects to the variability to specify here – the shape of the distribution and the spread of the distribution. This is where the normal distribution and our “normality assumption” reappears. And for normal distributions, we need to define a variance parameter,  $\sigma^2$ . Combined, the complete regression model is

$$y_i \sim N(\mu_{y_i}, \sigma^2), \text{ with } \mu_{y_i} = \beta_0 + \beta_1 x_i,$$

which can be read as “y follows a normal distribution with mean mu-y and variance sigma-squared”. This also implies that the random variability around the true mean, the errors, follow a normal distribution with mean 0 and that same variance,  $\varepsilon_i \sim N(0, \sigma^2)$ . The true deviations ( $\varepsilon_i$ ) are once again estimated by the residuals,  $e_i = y_i - \hat{y}_i = \text{observed response} - \text{predicted response}$ . We can use the residuals to estimate  $\sigma$ , which is also called the **residual standard error**,  $\hat{\sigma} = \sqrt{\sum e_i^2 / (n - 2)}$ . We will find this quantity near the end of the regression output as discussed below so the formula is not heavily used here. This provides us with the three parameters that are estimated as part of our SLR model:  $\beta_0$ ,  $\beta_1$ , and  $\sigma$ .

These definitions also formalize the assumptions implicit in the regression model:

1. The errors follow a normal distribution (**Normality assumption**).
2. The errors have the same variance (**Constant variance assumption**).
3. The observations are independent (**Independence assumption**).
4. The model for the mean is “correct” (**Linearity, No Influential points, Only one group**).

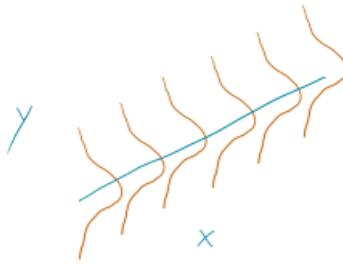


Figure 7.1: Sketch of assumed normal distributions for the responses centered at the regression line.

The diagnostics described at the end of Chapter 6 provide techniques for checking these assumptions – meeting these assumptions is fundamental to having a regression line that we trust and inferences from it that we also can trust.

To make this clearer, suppose that in the *Beers* and *BAC* study that they had randomly assigned 20 students to consume each number of beers. We would expect some variation in the *BAC* for each group of 20 at each level of *Beers* but that each group of observations will be centered at the true mean *BAC* for each number of *Beers*. The regression model assumes that the *BAC* values are normally distributed around the mean for each *Beer* level,  $BAC_i \sim N(\beta_0 + \beta_1 \text{Beers}_i, \sigma^2)$ , with the mean defined by the regression equation. We actually do not need to obtain more than one observation at each  $x$  value to make this assumption or assess it, but the plots below show you what this could look like. The sketch in Figure 7.1 attempts to show the idea of normal distributions that are centered at the true regression line, all with the same shape and variance that is an assumption of the regression model.

Figure 7.2 contains simulated realizations from a normal distribution of 20 subjects at each *Beer* level around the assumed true regression line with two different residual SEs of 0.02 and 0.06. The original BAC model has a residual SE of 0.02 but had many fewer observations at each *Beer* value.

Along with getting the idea that regression models define normal distributions in the  $y$ -direction that are centered at the regression line, you can also get a sense of how variable samples from a normal distribution can appear. Each distribution of 20 subjects at each  $x$  value came from a normal distribution but there are some of those distributions that might appear to generate small outliers and have slightly different variances. This can help us to remember to not be too particular when assessing assumptions and allow for some variability in spreads and a few observations from the tails of the distribution to occasionally arise.

In sampling from the population, we expect some amount of variability of each estimator around its true value. This variability leads to the potential variability in estimated regression lines (think of a suite of potential estimated regression lines that would be created by different random samples from the same population). Figure 7.3 contains the true regression line (bold, red) and realizations of the estimated regression line in simulated data based on results similar to the real data set.

This variability due to random sampling is something that needs to be properly accounted for to take the **single** estimated regression line to make inferences about the true line and parameters based on the sample-based estimates. The next sections develop those inferential tools.

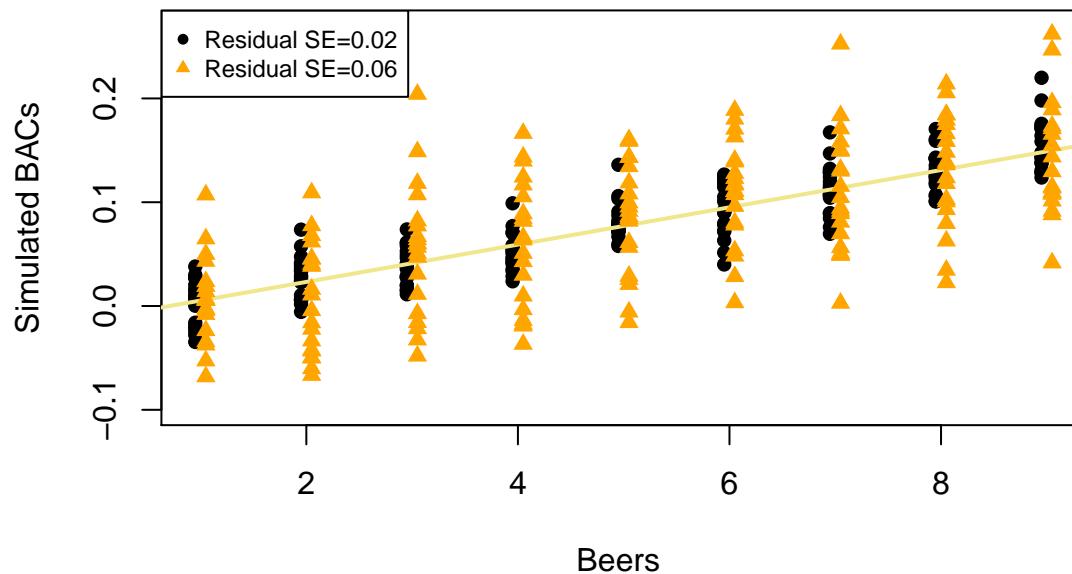


Figure 7.2: Simulated data for Beers and BAC assuming two different residual standard errors (0.02 and 0.06).

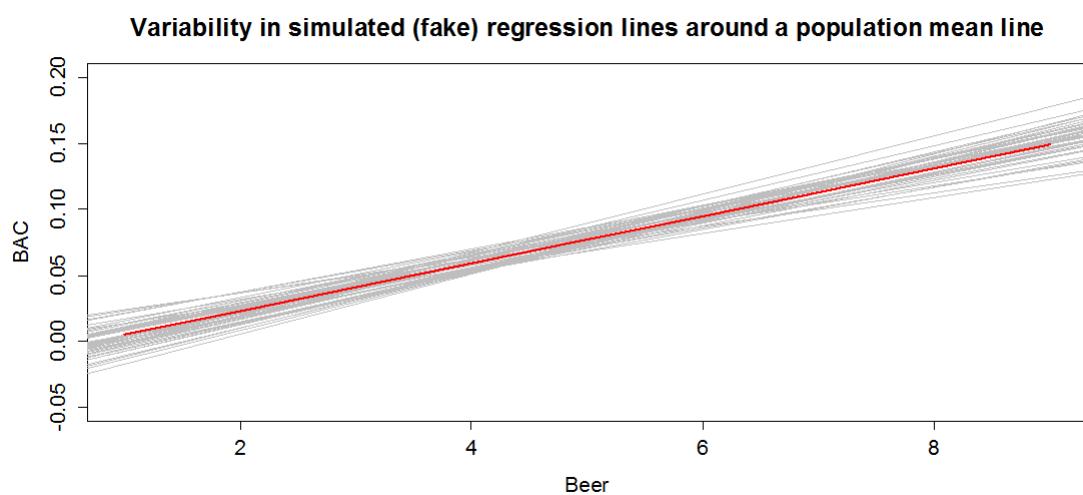


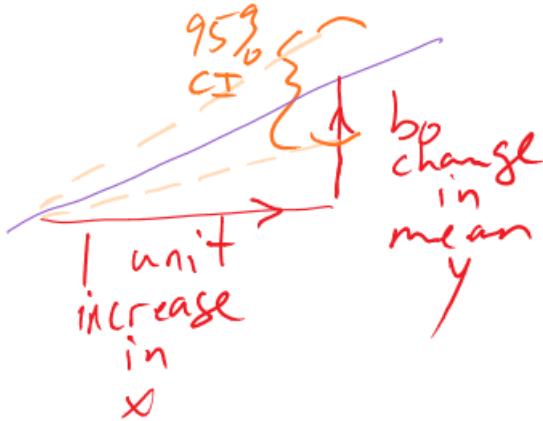
Figure 7.3: Variability in realized regression lines based on sampling variation.

## 7.2 Confidence Interval and Hypothesis tests for the slope and intercept

Our inference techniques will resemble previous material with an interest in forming confidence intervals and doing hypothesis testing, although the interpretation of confidence intervals for slope coefficients take some extra care. Remember that the general form of any parametric confidence interval is

$$\text{estimate} \mp t^* \text{SE}_{\text{estimate}},$$

so we need to obtain the appropriate standard error for regression model coefficients and the degrees of freedom to define the  $t$ -distribution to look up  $t^*$ . We will find the  $\text{SE}_{b_0}$  and  $\text{SE}_{b_1}$  in the model summary. The degrees of freedom for the  $t$ -distribution in simple linear regression are  $\text{df} = n - 2$ . Putting this together, the confidence interval for the true  $y$ -intercept,  $\beta_0$ , is  $b_0 \mp t_{n-2}^* \text{SE}_{b_0}$  although this confidence interval is rarely of interest. The confidence interval that is almost always of interest is for the true slope coefficient,  $\beta_1$ , that is  $b_1 \mp t_{n-2}^* \text{SE}_{b_1}$ . The slope confidence interval is used to do two things: (1) inference for the amount of change in the mean of  $y$  for a unit change in  $x$  in the population and (2) to potentially do hypothesis testing by checking whether 0 is in the CI or not. The sketch below illustrates the roles of the CI for the slope in terms of determining where the population slope coefficient might be – centered at the sample slope coefficient – our best guess for the true slope. This sketch informs an *interpretation of the slope coefficient confidence interval*:



For a 1 [units of  $X$ ] increase in  $X$ , we are \_\_\_\_ % confident that the true change in the mean of  $Y$  will be between LL and UL [units of  $Y$ ].

In this interpretation, LL and UL are the calculated lower and upper limits of the confidence interval. This builds on our previous interpretation of the slope coefficient, adding in the information about pinning down the true change (population change) in the mean of the response variable. The interpretation of the  $y$ -intercept CI is:

For an  $x$  of 0 [units of  $X$ ], we are 95% confident that the true mean of  $Y$  will be between LL and UL [units of  $Y$ ].

This is really only interesting if we'll see a method for generating CIs for the true mean at potentially more interesting values of  $x$  in Section 7.7. To trust the results from these confidence intervals, all the regression validity conditions need to be met (or at least close to met).

The only hypothesis test of interest in this situation is for the slope coefficient. To develop the hypotheses of interest in SLR, note the effect of having  $\beta_1 = 0$  in the mean of the regression equation,  $\mu_{y_i} = \beta_0 + \beta_1 x_i = \beta_0 + 0x_i = \beta_0$ . This is the “intercept-only” or “mean-only” model that suggests that the mean of  $y$  does not vary with different values of  $x$  as it is always  $\beta_0$ . We saw this model in the ANOVA material as the reduced model when the null hypothesis of no difference in the true means across the groups was true. Here, this is

the same as saying that there is no linear relationship between  $x$  and  $y$ , or that  $x$  is of no use in predicting  $y$ , or that we make the same prediction for  $y$  for every value of  $x$ . Thus

$$H_0 : \beta_1 = 0$$

is a test for **no linear relationship between x and y in the population**. The alternative of  $H_A : \beta_1 \neq 0$  that there is **some** linear relationship between  $x$  and  $y$  in the population, is our main test of interest in these situations. It is also possible to test greater than or less than alternatives in certain situations.

Test statistics for regression coefficients are developed, if assumptions are met, using the  $t$ -distribution with  $n - 2$  degrees of freedom. The  $t$ -test statistic is generally

$$t = \frac{b_i}{\text{SE}_{b_i}}$$

with the main interest in the test for  $\beta_1$  based on  $b_1$  for now. The p-value would be calculated using the two-tailed area from the  $t_{n-2}$  distribution calculated using the `pt` function. The p-value to test these hypotheses is also provided in the model summary as we will see below.

The greater than or less than alternatives can have interesting interpretations in certain situations. For example, the greater than alternative ( $H_A : \beta_1 > 0$ ) tests an alternative of a positive linear relationship, with the p-value extracted just from the right tail of the same  $t$ -distribution. This could be used when a researcher would only find a result “interesting” if a positive relationship is detected, such as in the study of tree height and tree diameter where a researcher might be justified in deciding to test only for a positive linear relationship. Similarly, the left-tailed alternative is also possible,  $H_A : \beta_1 < 0$ . To get one-tailed p-values from two-tailed results (the default), first check that the observed test statistic is in the direction of the alternative ( $t > 0$  for  $H_A : \beta_1 > 0$  or  $t < 0$  for  $H_A : \beta_1 < 0$ ). **If these conditions are met, then the p-value for the one-sided test from the two-sided version is found by dividing the reported p-value by 2.** If  $t > 0$  for  $H_A : \beta_1 > 0$  or  $t < 0$  for  $H_A : \beta_1 < 0$  are not met, then the p-value would be greater than 0.5 and it would be easiest to look it up directly using `pt`.

We can revisit a couple of examples for a last time with these ideas in hand to complete the analyses.

- For the *Beers*, *BAC* data, the 95% confidence for the true slope coefficient,  $\beta_1$ , is

$$\begin{aligned} b_1 \mp t^*_{n-2} \text{SE}_{b_1} &= 0.01796 \mp 2.144787 * 0.002402 \\ &= 0.01796 \mp 0.00515 \\ &\rightarrow (0.0128, 0.0231). \end{aligned}$$

You can find the components of this calculation in the model summary and from `qt(0.975, df=n-2)` which was 2.145 for the  $t^*$ -multiplier. Be careful not to use the  $t$ -value of 7.48 in the model summary to make confidence intervals – that is the test statistic used below. The related calculations are shown at the bottom of the following code:

```
m1 <- lm(BAC~Beers, data=BB)
summary(m1)
```

```
##
## Call:
## lm(formula = BAC ~ Beers, data = BB)
##
## Residuals:
##      Min       1Q   Median       3Q      Max
## -0.027118 -0.017350  0.001773  0.008623  0.041027
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
```

```

## (Intercept) -0.012701  0.012638 -1.005   0.332
## Beers        0.017964  0.002402  7.480 2.97e-06
##
## Residual standard error: 0.02044 on 14 degrees of freedom
## Multiple R-squared:  0.7998, Adjusted R-squared:  0.7855
## F-statistic: 55.94 on 1 and 14 DF,  p-value: 2.969e-06
qt(0.975, df=14) # t* multiplier for 95% CI

## [1] 2.144787
0.017964 + c(-1,1)*qt(0.975, df=14)*0.002402

## [1] 0.01281222 0.02311578
qt(0.975, df=14)*0.002402

## [1] 0.005151778

```

We can also get the confidence interval directly from the `confint` function run on our regression model, saving some calculation effort and providing both the CI for the y-intercept and the slope coefficient.

```
confint(m1)
```

```

##           2.5 %    97.5 %
## (Intercept) -0.03980535 0.01440414
## Beers       0.01281262 0.02311490

```

We interpret the 95% CI for the slope coefficient as follows: For a 1 **beer** increase in number of beers consumed, we are 95% confident that the **true** change in the **mean BAC** will be between 0.0128 and 0.0231 g/dL. While the estimated slope is our best guess of the impacts of an extra beer consumed based on our sample, this CI provides information about the likely range of potential impacts on the mean in the population. It also could be used to test the two-sided hypothesis test and would suggest that we should reject the null hypothesis since the confidence interval does not contain 0.

The width of the CI, loosely the precision of the estimated slope, is impacted by the variability of the observations around the estimated regression line, the overall sample size, and the positioning of the x-observations. Basically all those aspects relate to how “clearly” known the regression line is and that determines the estimated precision in the slope. For example, the more variability around the line that is present, the more uncertainty there is about the correct line to use (Least Squares (LS) can still find an estimated line but there are other lines that might be “close” to its optimizing choice). Similarly, more observations help us a better estimate of the mean – an idea that permeates all statistical methods. Finally, the location of x-values can impact the precision in a slope coefficient. We’ll revisit this in the context of **multi-collinearity** in the next chapter, and often we have no control of x-values, but just note that different patterns of x-values can lead to different precision of estimated slope coefficients<sup>1</sup>.

For hypothesis testing, we will almost always stick with two-sided tests in regression modeling as it is a more conservative approach and does not require us to have an expectation of a direction for relationships *a priori*. In this example, the null hypothesis for the slope coefficient is that there is no linear relationship between *Beers* and *BAC* in the population. The alternative hypothesis is that there is some linear relationship between *Beers* and *BAC* in the population. The test statistic is  $t = 0.01796/0.002402 = 7.48$  which, if assumptions hold, follows a  $t(14)$  distribution. The model summary provides the calculation of the test statistic and the two-sided test p-value of  $2.97e-6 = 0.00000297$ . So we would just report  $p\text{-value} < 0.0001$ . This suggests we should reject the null hypothesis and conclude that there is evidence at the 5% significance level of a linear relationship between *Beers* and *BAC* in the population. Because of the random assignment, we can also say that drinking beers causes changes in BAC but, because the sample was of volunteers, we cannot infer that these results would hold in the general population of OSU students or more generally.

---

<sup>1</sup>There is an area of statistical research on how to optimally choose x-values to get the most precise estimate of a slope coefficient. In observational studies we have to deal with whatever pattern of  $x$ 's we ended up with. If you can choose, generate an even spread of  $x$ 's over some range of interest similar to what was used in the *Beers vs BAC* study to provide the best distribution of values to discover the relationship across the selected range of x-values.

There are also results for the y-intercept in the output. The 95% CI is from -0.0398 to 0.0144, that the true mean *BAC* for a 0 beer consuming subject is between -0.0398 to 0.01445. This is really not a big surprise but possibly is comforting to know that these results would fail to reject the null hypothesis that the true mean *BAC* for 0 Beers is 0. Finding no evidence of a difference from 0 makes sense and makes the estimated y-intercept of -0.013 not so problematic. In other situations, the results for the y-intercept may be more illogical but this will often be because the y-intercept is extrapolating far beyond the scope of observations. The y-intercept's main function in regression models is to be at the right level for the slope to "work" to make a line that describes the responses and thus is usually of lesser interest.

As a second example, we can revisit modeling the *Hematocrit* of female Australian athletes as a function of *body fat %*. The sample size is  $n = 99$  so the *df* are 97 in the analysis. In Chapter 6, the relationship between *Hematocrit* and *body fat %* for females appeared to be a weak negative linear association. The 95% confidence interval for the slope is -0.187 to 0.0155. For a 1 % increase in body fat %, we are 95% confident that the change in the true mean Hematocrit is between -0.187 and 0.0155 % of blood. This suggests that we would fail to reject the null hypothesis of no linear relationship at the 5% significance level because this CI contains 0 – we can't reject the null that the true slope is 0. In fact the p-value is 0.0965 which is larger than 0.05 which provides a consistent conclusion with using the 95% confidence interval to perform a hypothesis test. Either way, we would conclude that there is not enough evidence at the 5% significance level to conclude that there is some linear relationship between body fat and Hematocrit in the population of female Australian athletes. If your standards were different, say if you had elected to test at the 10% significance level, you might have a different opinion about the evidence against the null hypothesis here. For this reason, we sometimes interpret this sort of marginal result as having some evidence against the null but certainly not strong evidence.

```
require(alr3)
data(ais)
aisR2<-ais[-c(56,166),c("Ht","Hc","Bfat","Sex")]
m2 <- lm(Hc~Bfat, data=aisR2[aisR2$Sex==1,]) # Results for Females
summary(m2)
```

```
##
## Call:
## lm(formula = Hc ~ Bfat, data = aisR2[aisR2$Sex == 1, ])
##
## Residuals:
##      Min       1Q   Median       3Q      Max
## -5.2399 -2.2132 -0.1061  1.8917  6.6453
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) 42.01378   0.93269 45.046  <2e-16
## Bfat        -0.08504   0.05067 -1.678   0.0965
##
## Residual standard error: 2.598 on 97 degrees of freedom
## Multiple R-squared:  0.02822,    Adjusted R-squared:  0.0182
## F-statistic: 2.816 on 1 and 97 DF,  p-value: 0.09653
confint(m2)
```

```
##           2.5 %     97.5 %
## (Intercept) 40.1626516 43.86490713
## Bfat        -0.1856071  0.01553165
```

One more worked example is provided from the Montana fire data. In this example pay particular attention to how we are handling the units of the response variable, log-hectares, and to the changes to doing inferences at the 1% significance and 99% confidence levels, and where you can find the needed results in the following output:

```
mtfires <- read.csv("http://www.math.montana.edu/courses/s217/documents/climateR2.csv")
mtfires$loghectacres <- log(mtfires$hectacres)
```

```

fire1 <- lm(loghectacres~Temperature, data=mtfires)
summary(fire1)

##
## Call:
## lm(formula = loghectacres ~ Temperature, data = mtfires)
##
## Residuals:
##      Min       1Q   Median       3Q      Max
## -3.0822 -0.9549  0.1210  1.0007  2.4728
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) -69.7845    12.3132 -5.667 1.26e-05
## Temperature  1.3884     0.2165  6.412 2.35e-06
##
## Residual standard error: 1.476 on 21 degrees of freedom
## Multiple R-squared:  0.6619, Adjusted R-squared:  0.6458
## F-statistic: 41.12 on 1 and 21 DF,  p-value: 2.347e-06
confint(fire1, level=0.99)

##
##          0.5 %    99.5 %
## (Intercept) -104.6477287 -34.921286
## Temperature    0.7753784   2.001499
qt(0.995, df=21)

```

## [1] 2.83136

- Based on the estimated regression model, we can say that if the average temperature is 0, we expect that, on average, the log-area burned would be -69.8 log-hectares.
- From the regression model summary,  $b_1 = 1.39$  with  $SE_{b_1} = 0.2165$  and  $t = 6.41$
- There were  $n = 23$  measurements taken, so  $df = n - 2 = 23 - 3 = 21$
- Suppose that we want to test for a linear relationship between temperature and log-hectares burned:

$$H_0 : \beta_1 = 0$$

- In words, the true slope coefficient between *Temperature* and *log-area burned* is 0 OR there is no linear relationship between *Temperature* and *log-area burned* in the population.

$$H_A : \beta_1 \neq 0$$

- In words, the alternative states that the true slope coefficient between *Temperature* and *log-area burned* is not 0 OR there is a linear relationship between *Temperature* and *log-area burned* in the population.

Test statistic:  $t = 1.39/0.217 = 6.41$

- Assuming the null hypothesis to be true (no linear relationship), the  $t$ -statistic follows a  $t$ -distribution with  $n - 2 = 23 - 2 = 21$  degrees of freedom.

p-value:

- From the model summary, the **p-value is  $2.35 * 10^{-6}$** 
  - Interpretation: There is less than a 0.01% chance that we would observe slope coefficient like we did or something more extreme (greater than  $1.39 \text{ log}(hectares)/{}^{\circ}\text{F}$ ) if there were in fact no linear relationship between temperature ( ${}^{\circ}\text{F}$ ) and log-area burned (log-hectares) in the population.

Decision: At the 1% significance level ( $\alpha = 0.01$ ), the p-value is less than  $\alpha$ , so reject  $H_0$ .

Conclusion: There is strong evidence to reject the null hypothesis of no linear relationship and conclude that there is, in fact, a linear relationship between Temperature and log(Hectares) burned. Since we have a time series of results, our inferences pertain to the results we could have observed for these years but not for years we did not observe – so just for the true slope for this sample of years. Because we can't randomly assign the amount of area burned, we cannot make causal inferences – there are many reasons why both the average temperature and area burned would vary together that would not involve a direct connection between them.

$$99\% \text{ CI for } \beta_1 : b_1 \mp t_{n-2}^* \text{SE}_{b_1} \rightarrow 1.39 \mp 2.831 \bullet 0.217 \rightarrow (0.78, 2.00)$$

Interpretation of 99% CI for slope coefficient:

- For a 1 degree F increase in *Temperature*, we are 99% confident that the change in the true mean log-area burned is between 0.78 and 2.00 log(Hectares).
- Another way to interpret this is:
  - For a 1 degree F increase in *Temperature*, we are 99% confident that the mean Area Burned will change by between 0.78 and 2.00 log(Hectares) **in the population**.
- Also  $R^2$  is 66.2%, which tells us that *Temperature* explains 66.2% of the variation in *log(Hectares) burned*. Or that the linear regression model built using *Temperature* explains 66.2% of the variation in yearly *log(Hectares) burned*.

## 7.3 Bozeman temperature trend

For a new example, consider the yearly average maximum temperatures in Bozeman, MT. For over 100 years, daily measurements have been taken of the minimum and maximum temperatures at hundreds of weather stations across the US. In early years, this involved manual recording of the temperatures and resetting the thermometer to track the extremes for the following day. More recently, these measures have been replaced by digital temperature recording devices that continue to track this sort of information with much less human effort and, possibly, errors. This sort of information is often aggregated to monthly or yearly averages to be able to see “on average” changes from month-to-month or year-to-year as opposed to the day-to-day variation in the temperature - something that we are all too familiar with in our part of the country (see <http://fivethirtyeight.com/features/which-city-has-the-most-unpredictable-weather/> for an interesting discussion of weather variability where Great Falls, MT had a very high rating on “unpredictability”). Often the local information is aggregated further to provide regional, hemispheric, or even global average temperatures. Climate change research involves attempting to quantify the changes over time in these sorts of records.

These data were extracted from the National Oceanic and Atmospheric Administration’s National Centers for Environmental Information’s database (<http://www.ncdc.noaa.gov/cdo-web/>) and we will focus on the yearly average of the monthly averages of the daily maximum temperature (we can call them yearly average maximum temperatures but note that it was a little more complicated than that to arrive at the response variable) in Bozeman in degrees F from 1901 to 2014.

```
bozemantemps <- read.csv("http://www.math.montana.edu/courses/s217/documents/BozemanMeanMax.csv")
summary(bozemantemps)
```

```
##      meanmax        Year
## Min.   :49.75  Min.   :1901
## 1st Qu.:53.97  1st Qu.:1930
## Median :55.43  Median :1959
## Mean   :55.34  Mean   :1958
## 3rd Qu.:57.02  3rd Qu.:1986
## Max.   :60.05  Max.   :2014
```

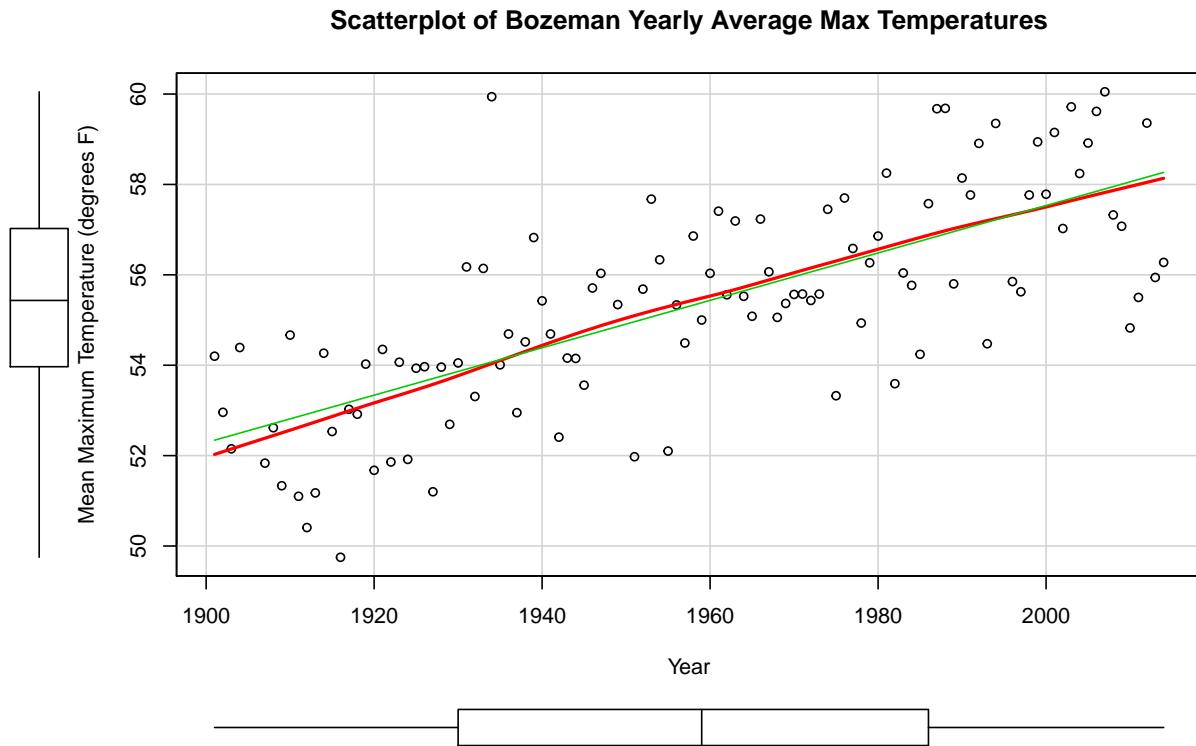


Figure 7.4: Scatterplot of average yearly maximum temperatures in Bozeman from 1900 to 2014.

```
length(bozemantemps$Year) #Some years are missing (1905, 1906, 1948, 1950, 1995)
```

```
## [1] 109
require(car)
scatterplot(meanmax~Year, data=bozemantemps,
           ylab="Mean Maximum Temperature (degrees F)", spread=F,
           main="Scatterplot of Bozeman Yearly Average Max Temperatures")
```

The scatterplot in Figure 7.4 shows the results between 1901 and 2014 based on a sample of  $n = 109$  years because four years had too many missing months to fairly include in the responses. Missing values occur for many reasons and in this case were likely just machine or human error<sup>2</sup>. These are time series data and in time series analysis we assume that the population of interest for inference is all possible realizations from the underlying process over this time frame even though we only ever get to observe one realization. In terms of climate change research, we would want to (a) assess evidence for a trend over time (hopefully assessing whether any observed trend is clearly different from a result that could have been observed by chance if there really is no change over time in the true process) and (b) quantify the size of the change over time along with the uncertainty in that estimate relative to the underlying true mean change over time. The hypothesis test for the slope answers (a) and the confidence interval for the slope addresses (b). We also should be concerned about problematic (influential) points, changing variance, and potential nonlinearity in the trend over time causing problems for the SLR inferences. The scatterplot suggests that there is a moderate or strong positive linear relationship between *temperatures* and *year* with some “wiggles” in the smoothing line at the beginning and end of the record. Smoothing lines can become quite untrustworthy at the edges of the data set, so discount the curving at the edges a bit. If the curving is real, it would suggest a less steep change before 1920, relatively linear change from 1930 to 1970, a small increase in slope through the mid-90s, and then a leveling off after that point. There also appears to be one potential large outlier in the late 1930s.

<sup>2</sup>It is actually pretty amazing that there are hundreds of locations with nearly complete daily records for over 100 years.

We'll perform all 6+ steps of the hypothesis test for the slope coefficient and add a confidence interval interpretation for this example. First, we have to decide on our significance level (5% is a typical choice), our hypotheses (the 2-sided test would be a *conservative* choice and no one that does climate change research wants to be accused of taking a *liberal* approach in their analyses<sup>3</sup>) and our test statistic,  $t = \frac{b_1}{\text{SE}_{b_1}}$ .

### 1. Hypotheses for the slope coefficient test:

$$H_0 : \beta_1 = 0 \text{ vs } H_A : \beta_1 \neq 0$$

### 2. Validity conditions:

- Quantitative variables condition

- Both Year and yearly average Temperature are quantitative variables so are suitable for an SLR analysis.

- Independence of observations

- There may be a lack of independence among years since a warm year might be followed by another warmer than average year. It would take more sophisticated models to account for this and the standard error on the slope coefficient could either get larger or smaller depending on the type of *autocorrelation* (correlation between neighboring time points or correlation with oneself at some time lag) present. This creates a caveat on these results but this model is often the first one researchers fit in these situations and often is reasonably correct even in the presence of some autocorrelation.

To assess the remaining conditions, we need to fit the regression model and use the diagnostic plots in Figure 7.5 to aid our assessment:

```
temp1 <- lm(meanmax~Year, data=bozemantemps)
par(mfrow=c(2,2))
plot(temp1, add.smooth=F)
```

- Linearity of relationship

- Examine the Residuals vs Fitted plot:

- There does not appear to be a clear curve remaining in the residuals so that initial curving in the smoothing line is not clearly showing up in the diagnostics so we should be able to proceed without worrying too much about the slight nonlinearity detected in the initial scatterplot.

- Equal (constant) variance

- Examining the Residuals vs Fitted and the “Scale-Location” plots provide little to no evidence of changing variance. The variability does decrease slightly in the middle fitted values but those changes are really minor and present no real evidence of changing variability.

- Normality of residuals

- Examining the Normal QQ-plot for violations of the normality assumption shows only one real problem in the outlier from the 32nd observation in the data set (1934) which was flagged as a large outlier in the original scatterplot. We should be careful about inferences that assume normality and contain this point in the analysis. We might consider running the analysis with it and without that point to see how much it impacts the results just to be sure it isn't creating evidence of a trend because of a violation of the normality assumption. The next check reassures us that re-running the model without this point would only result in slightly changing the SEs and not the slopes.

- No influential points:

---

<sup>3</sup>All joking aside, if researchers can find evidence of climate change using *conservative* methods (methods that reject the null hypothesis when it is true less often than stated), then their results are even harder to ignore.

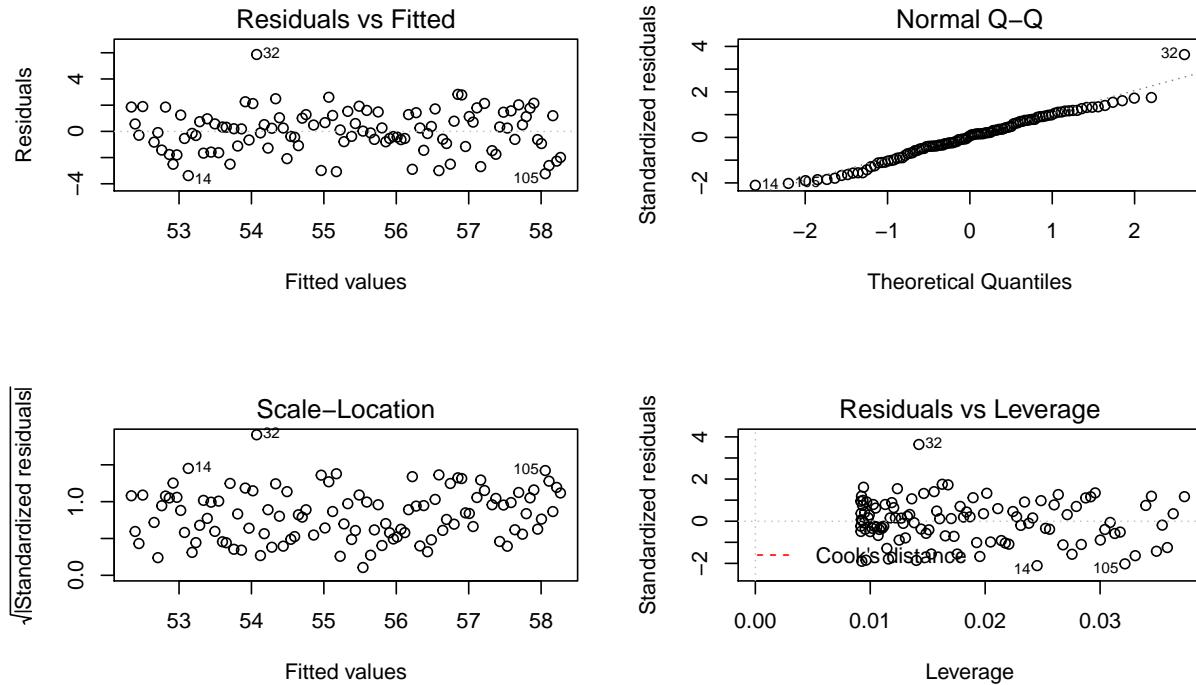


Figure 7.5: Diagnostic plots of the Bozeman yearly temperature simple linear regression model.

- There are no influential points displayed in the Residuals vs Leverage plot since the Cook's D contours are not displayed.
  - Note: by default this plot contains a smoothing line that is relatively meaningless, so ignore it if it is displayed. We suppressed it using the `add.smooth=F` option in `plot(temp1)` but if you forget to do that, just ignore the smoothers in the diagnostic plots especially in the Residuals vs Leverage plot.
- This results tells us that the outlier was not influential. If you look back at the scatterplot, it was located near the middle of the observed  $x$ 's so its potential leverage is low. You can find its leverage based on the plot to be around 0.12 when there are observations in the data set with leverages over 0.3. The high leverage points occur at the beginning and the end of the record because they are at the edges of the observed  $x$ 's and most of these points follow the overall pattern fairly well.

So the main issues are with independence of observations and one non-influential outlier that might be compromising our normality assumption a bit.

### 3. Calculate the test statistic:

$$t = 0.05244 / 0.00476 = 11.02$$

```
summary(temp1)
```

```
##
## Call:
## lm(formula = meanmax ~ Year, data = bozemantemps)
##
## Residuals:
##     Min      1Q  Median      3Q     Max 
## -3.3779 -0.9300  0.1078  1.1960  5.8698
```

```

## 
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) -47.35123   9.32184  -5.08 1.61e-06
## Year         0.05244   0.00476   11.02 < 2e-16
##
## Residual standard error: 1.624 on 107 degrees of freedom
## Multiple R-squared:  0.5315, Adjusted R-squared:  0.5271
## F-statistic: 121.4 on 1 and 107 DF,  p-value: < 2.2e-16

```

4. Find the p-value:

- From the model summary: p-value < 2e-16 or just < 0.0001
- The test statistic is assumed to follow a  $t$ -distribution with  $n - 2 = 109 - 2 = 107$  degrees of freedom. The p-value can be calculated as:

```
2*pt(11.02, df=107, lower.tail=F)
```

```
## [1] 2.498481e-19
```

- Which is then reported as < 0.0001, which means that the chances of observing a slope coefficient as extreme or more extreme than 0.052 if the null hypothesis of no linear relationship is true is less than 0.01%.

5. **Make a decision:**

Reject the null hypothesis because the p-value is less than 0.05.

6. **Write a conclusion:**

There is strong evidence against the null hypothesis of no linear relationship between *Year* and yearly mean *Temperature* so we can conclude that there is, in fact, some linear relationship between *Year* and yearly mean maximum *Temperature* in Bozeman. We can conclude that this detected trend pertains to the Bozeman area in the years 1901 to 2014 but not outside of this area or time frame. We cannot say that time caused the observed changes since it was not randomly assigned and we cannot attribute the changes to any other factors because we did not consider them. But knowing that there was a trend toward increasing temperatures is an intriguing first step in a more complete analysis of changing climate in the area.

It is also good to report the percentage of variation that the model explains: *Year* explains 54.91% of the variation in yearly average maximum *Temperature*. If this value had been very small, we might discount the previous result. Since it is moderately large, that suggests that just by using a linear trend over time we can account for quite a bit of the variation in yearly average maximum temperatures in Bozeman. Note that the percentage of variation explained would get much worse if we tried to analyze the monthly or original daily maximum temperature data.

Interpreting a confidence interval provides more useful information than the hypothesis test here – instead of just assessing evidence against the null hypothesis, we can actually provide our best guess at the true change in the mean of  $y$  for a change in  $x$ . Here, the 95% CI is (0.043, 0.062). This tells us that for a 1 year increase in change in the true mean of the yearly average maximum *Temperatures* in Bozeman is between 0.043 and 0.062 degrees F.

```
confint(temp1)
```

```

##           2.5 %      97.5 %
## (Intercept) -65.83068375 -28.87177785
## Year        0.04300681   0.06187746

```

Sometimes the scale of the x-variable makes interpretation a little difficult, so we can re-scale it to make it more interpretable. One option is to re-scale the variable and re-fit the regression model and the other (easier) option is to re-scale our interpretation. The idea here is that a 100-year change might be easier and more meaningful scale to interpret than a single year change. If we have a slope in the model of 0.052 (for a 1

year change), we can also say that a 100 year change in the mean is estimated to be  $0.052 * 100 = 0.52^{\circ}F$ . Similarly, the 95% CI for the population mean 100-year change would be from  $0.43^{\circ}F$  to  $0.62^{\circ}F$ . In 2007, the IPCC (Intergovernmental Panel on Climate Change; [http://www.ipcc.ch/publications\\_and\\_data/ar4/wg1/en/tssts-3-1-1.html](http://www.ipcc.ch/publications_and_data/ar4/wg1/en/tssts-3-1-1.html)) estimated the global temperature change from 1906 to 2005 to be  $0.74^{\circ}C$  per decade or, scaled up,  $7.4^{\circ}C$  per century ( $1.33^{\circ}F$ ). There are many reasons why our local temperature trend might differ, including that our analysis was of average maximum temperatures and the IPCC was considering the average temperature (which was not measured locally or in most places in a good way until digital instrumentation was installed) and that local trends are likely to vary around the global average change based on localized environmental conditions.

One issue that arises in local studies of climate change is that researchers often consider these sorts of tests at many locations and on many response variables (if I did the maximum temperature, why not also do the same analysis of the minimum temperature time series as well? And if I did the analysis for Bozeman, what about Butte and Helena and...?). Remember our discussion of multiple testing issues in an ANOVA context when we compared lots of groups? This issue can arise when regression modeling is repeated in many similar data sets, say different sites or different response variables or both, in one study. Moore and Greenwood (2007) considered the impacts on the assessment of evidence of trends of earlier spring onset timing in the Mountain West when the number of tests across many sites is accounted for. We found that the evidence for time trends decreases substantially but does not disappear. In a related study, Greenwood and Moore (2011) found evidence for regional trends to earlier spring onset using more sophisticated statistical models. The main point here is to **be careful when using simple statistical methods repeatedly if you are not accounting for the number of tests performed**<sup>4</sup>.

Along with the confidence interval, we can also plot the estimated model (Figure 7.6 using a term-plot from the `effects` package (Fox, 2003). This is the same function we used for visualizing results in the ANOVA models and in its basic application you just need `plot(allEffects(modelname))` but we enhanced our version a little. In regression models, we get to see the regression line along with bounds for 95% confidence intervals for the mean at every value of  $x$  that was observed (explained in next section). Note that there is also a rugplot on the x-axis showing you where values of the explanatory variable were obtained, which is useful to understanding how much information is available for different aspects of the line. Here it provides gaps for missing years of observations as sort of broken teeth in a comb.

```
require(effects)
plot(allEffects(temp1, xlevels=list(Year=bozemantemps$Year)),
     grid=T)
```

If we extended the plot for the model to `Year=0`, we could see the reason that the y-intercept in this model is  $-47.4^{\circ}F$ . This is obviously a large extrapolation for these data and provides a silly result. However, in paleoclimate data that goes back thousands of years using tree rings, ice cores, or sea sediments, the estimated mean in year 0 might be interesting and within the scope of observed values. It all depends on the application.

To make the y-intercept more interesting for our data set, we can re-scale the  $x$ 's before we fit the model to have the first year in the data set (1901) be "0". This is accomplished by calculating `Year2 = Year - 1901`.

```
bozemantemps$Year2 <- bozemantemps$Year - 1901
summary(bozemantemps$Year2)
```

```
##      Min. 1st Qu. Median   Mean 3rd Qu.   Max.
##      0.00  29.00  58.00  57.27  85.00 113.00
```

The new estimated regression equation is  $\widehat{\text{Temp}}_i = 52.34 + 0.052 * \text{Year2}_i$ . The slope and its test statistic are the same as in the previous model. The y-intercept has changed dramatically with a 95% from  $51.72^{\circ}F$  to  $52.96^{\circ}F$  for `Year2=0`. But we know that `Year2` has a 0 value for 1901 because of our subtraction. That means that this CI is for the true mean in 1901 and is now at least somewhat interesting. If you revisit Figure 7.6 you will actually see that the displayed confidence intervals provide upper and lower bounds that match this result for 1901 – the y-intercept CI matches the 95% CI for the true mean.

---

<sup>4</sup>The simplest adjustment for multiple testing is using a Bonferroni adjustment, where you multiply all the p-values by the number of tests performed. It controls the chances of at least one error to be same as your Type I error rate for one test.

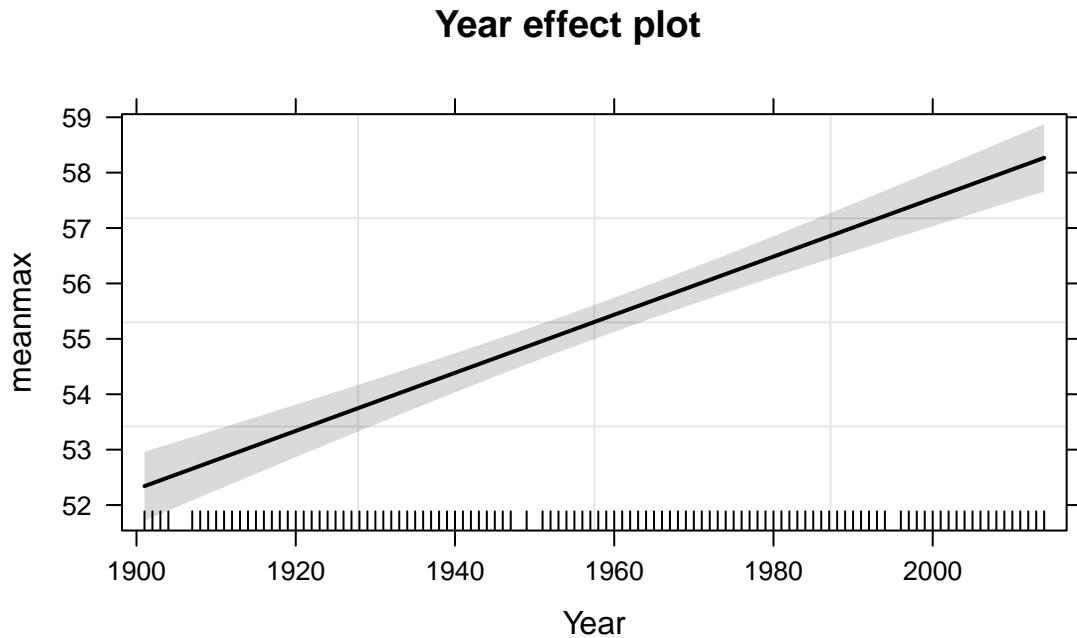


Figure 7.6: Term-plot for the Bozeman mean yearly maximum temperature linear regression model with 95% confidence interval bands for the mean in each year.

```
temp2 <- lm(meanmax~Year2, data=bozemantemps)
summary(temp2)

##
## Call:
## lm(formula = meanmax ~ Year2, data = bozemantemps)
##
## Residuals:
##      Min       1Q   Median       3Q      Max 
## -3.3779 -0.9300  0.1078  1.1960  5.8698 
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) 52.34126   0.31383 166.78 <2e-16 ***
## Year2        0.05244   0.00476 11.02 <2e-16 ***
## 
## Residual standard error: 1.624 on 107 degrees of freedom
## Multiple R-squared:  0.5315, Adjusted R-squared:  0.5271 
## F-statistic: 121.4 on 1 and 107 DF,  p-value: < 2.2e-16
confint(temp2)

##
##           2.5 %      97.5 %
## (Intercept) 51.71913822 52.96339150
## Year2        0.04300681  0.06187746
```

Ideally, we want to find a regression model that does not violate any assumptions, has a high  $R^2$  value, and a slope coefficient with a small p-value. If any of these are not the case, then we are not completely satisfied with the regression and **should be suspicious of any inference we perform**. We can sometimes resolve some of the systematic issues noted above using *transformations*, discussed in Sections 7.5 and 7.6.

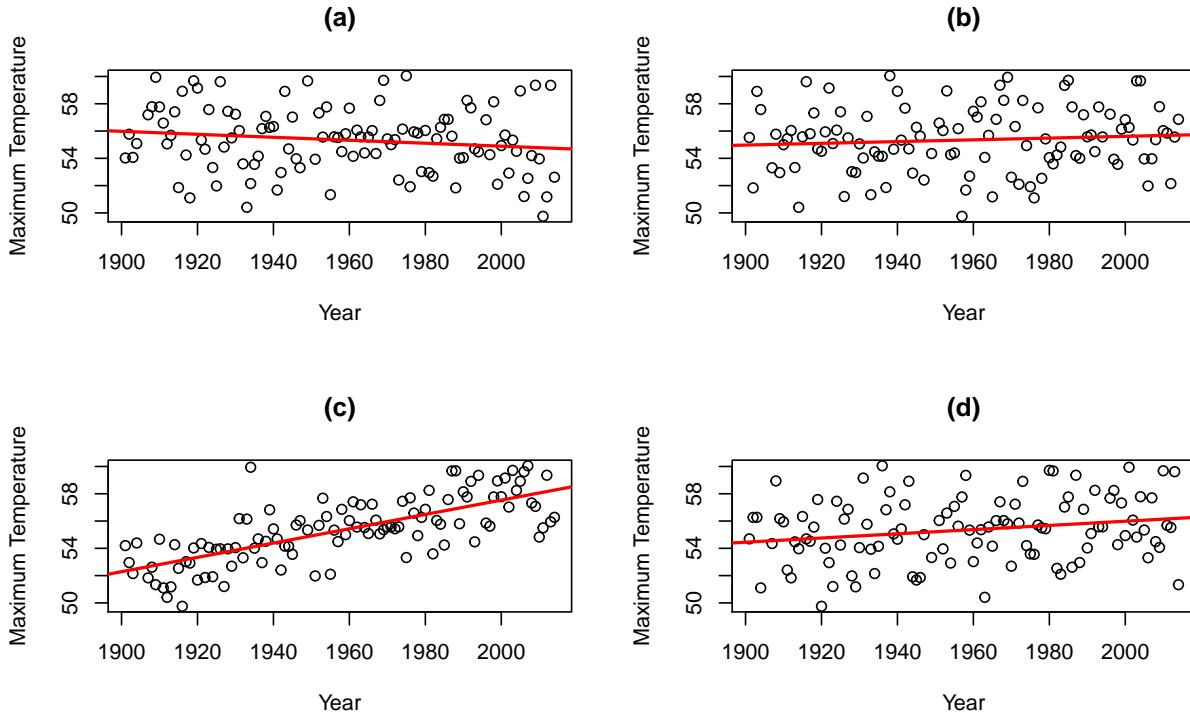


Figure 7.7: Plot of the Temperature responses versus four versions of Year, three of which are permutations of the Year variable relative to the Temperatures.

## 7.4 Randomizing inferences for the slope coefficient

Exploring permutation testing in SLR provides an opportunity to gauge the observed relationship against the sorts of relationships we would expect to see if there was no linear relationship between the variables. If the relationship is linear (not curvilinear) and the null hypothesis of  $\beta_1 = 0$  is true, then any configuration of the responses relative to the predictor variables is a good as any other. Consider the four scatterplots of the Bozeman temperature data versus `Year` and permuted versions of `Year` in Figure 7.7. First, think about which of the panels you think presents the most evidence of a linear relationship between `Year` and `Temperature`?

Hopefully you can see that panel (c) contains the most clear linear relationship among the choices. The plot in panel (c) is actually the real data set and pretty clearly presents itself as “different” from the other results. When we have small p-values, the real data set will be clearly different from the permuted results because it will be almost impossible to find a permuted data set that can attain as large a slope coefficient as was observed in the real data set<sup>5</sup>. This result ties back into our original interests in this climate change research situation – does our result look like it is different from what could have been observed just by chance if there were no linear relationship between  $x$  and  $y$ ? It seems unlikely...

Repeating this permutation process and tracking the estimated slope coefficients, as  $T^*$ , provides another method to obtain a p-value in SLR applications. This could also be performed on the  $t$ -statistic for the slope coefficient and would provide the same p-values but the sampling distribution would have a different x-axis scaling. In this situation, the observed slope of 0.052 is really far from any possible values that can be obtained using permutations as shown in Figure 7.8. The p-value would be reported as  $< 0.0001$  for the two-sided test.

<sup>5</sup>It took many permutations to get competitor plots this close to the real data set and they really aren’t that close.

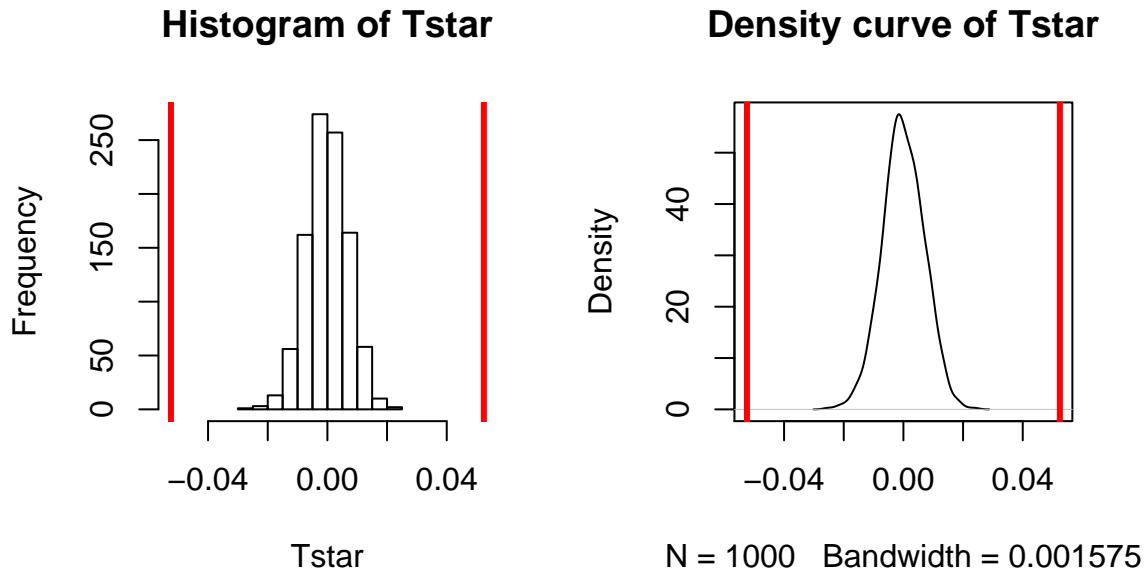


Figure 7.8: Permutation distribution of the slope coefficient in the Bozeman temperature linear regression model with bold vertical lines at  $\pm b_1 = 0.56$ .

```

Tobs <- lm(meanmax~Year, data=bozemantemps)$coef[2]
Tobs

##      Year
## 0.05244213
par(mfrow=c(1,2))
B <- 1000
Tstar <- matrix(NA,nrow=B)
for (b in (1:B)){
  Tstar[b] <- lm(meanmax~shuffle(Year), data=bozemantemps)$coef [2]
}
pdata(abs(Tstar), abs(Tobs), lower.tail=F)

## Year
## 0
hist(Tstar, xlim=c(-1,1)*Tobs)
abline(v=c(-1,1)*Tobs, col="red", lwd=3)
plot(density(Tstar), main="Density curve of Tstar", xlim=c(-1,1)*Tobs)
abline(v=c(-1,1)*Tobs, col="red", lwd=3)

```

One other interesting aspect of exploring the permuted data sets as in Figure 7.7 is that the outlier in the late 1930s “disappears” in the permuted data sets because there were many other observations that were that warm, just none that happened around that time of the century in the real data set. This reinforces the evidence for changes over time that seem to be present in these data that old unusual years don’t look unusual in more recent years.

The permutation approach can be useful in situations where the normality assumption is compromised, but there are no influential points. In these situations, we might find more trustworthy p-values using permutations but only if we are working with an initial estimated regression equation that we generally trust. I personally like the permutation approach as a way of explaining what a p-value is actually measuring –

the chance of seeing something like what we saw, or more extreme, if the null is true. And the previous scatterplots show what the “by chance” versions of this relationship might look like.

In a similar situation where we want to focus on confidence intervals for slope coefficients but are not completely comfortable with the normality assumption, it is also possible to generate bootstrap confidence intervals by sampling with replacement from the data set. This idea was introduced in Sections 2.8 and 2.9. This provides a 95% bootstrap confidence interval from 0.433 to 0.62, which almost exactly matches the parametric  $t$ -based confidence interval. The bootstrap distributions are very symmetric (Figure 7.9). The interpretation is the same and this result reinforces the other assessments that the parametric approach is not unreasonable except possibly for the independence assumption.

```
Tobs <- lm(meanmax~Year, data=bozemantemps)$coef[2]
Tobs

##          Year
## 0.05244213
par(mfrow=c(1,2))
B <- 1000
Tstar <- matrix(NA,nrow=B)
for (b in (1:B)){
  Tstar[b] <- lm(meanmax~Year, data=resample(bozemantemps))$coef[2]
}
quantiles <- qdata(Tstar, c(.025,.975))
quantiles

##      quantile     p
## 2.5%  0.04331326 0.025
## 97.5% 0.06195345 0.975
hist(Tstar,labels=T)
abline(v=Tobs,col="red",lwd=2)
abline(v=quantiles$quantile,col="blue",lwd=3)
plot(density(Tstar),main="Density curve of Tstar")
abline(v=Tobs,col="red",lwd=2)
abline(v=quantiles$quantile,col="blue",lwd=3)
```

## 7.5 Transformations part I: Linearizing relationships

When the influential point, linearity, constant variance and/or normality assumptions are violated, we cannot trust any of the inferences generated by the regression model. The violations occur on gradients from minor to really major problems. As we have seen from the examples in the last two chapters, it has been hard to find data sets that were free of all issues. Furthermore, it may seem hopeless to be able to make successful inferences in some of these situations with the previous tools. There are three potential solutions to violations of the validity conditions:

1. Consider removing an offending point or two and see if this improves the results, presenting results both with and without those points to describe their impact<sup>6</sup>,
2. Try to transform the response, explanatory, or both variables and see if you can force the data set to meet our SLR assumptions after transformation (the focus of this and the next section), or
3. Consider more advanced statistical models that can account for these issues (the focus of additional statistics courses).

---

<sup>6</sup>If the removal is of a point that is extreme in x-values, then it is appropriate to note that the results only apply to the restricted range of x-values that were actually analyzed. Our results only ever apply to the range of x-values we had available so this is a relatively minor change.

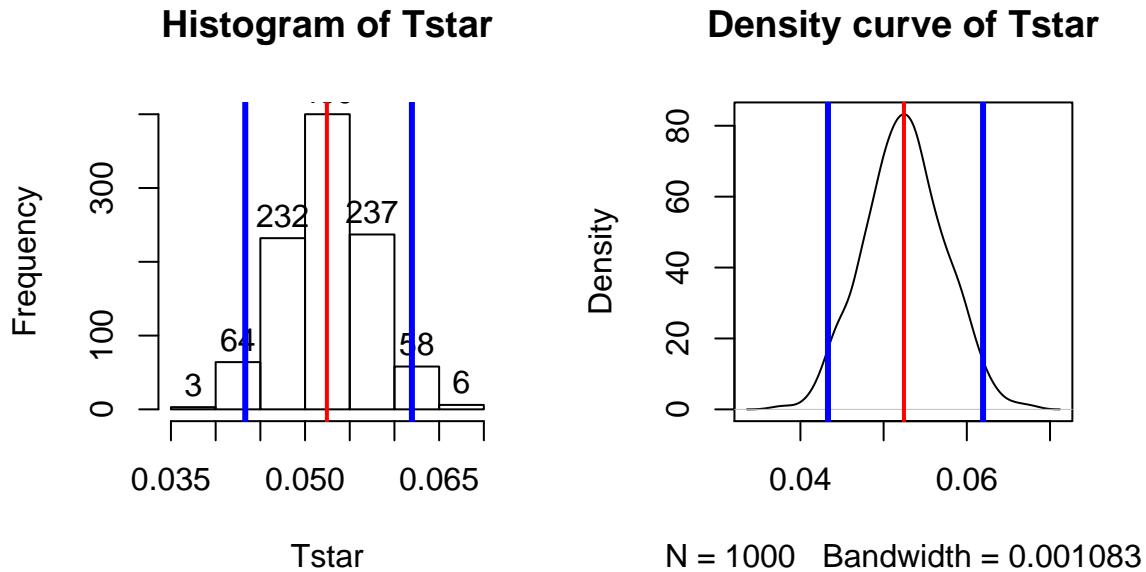


Figure 7.9: Bootstrap distribution of the slope coefficient in the Bozeman temperature linear regression model with bold vertical lines delineating 95% confidence interval and observed slope of 0.52.

**Transformations** involve applying a function to one or both variables. After applying this transformation, one hopes to have alleviated whatever issues encouraged its consideration. **Linear transformation functions**, of the form  $z_{\text{new}} = a * x + b$ , will never help us to fix assumptions in regression situations; linear transformations change the scaling of the variables but not their shape or the relationship between two variables. For example, in the Bozeman Temperature data example, we subtracted 1901 from the `Year` variable to have `Year2` start at 0 and go up to 113. We could also apply a linear transformation to change Temperature from being measured in  $^{\circ}\text{F}$  to  $^{\circ}\text{C}$  using  $^{\circ}\text{C} = [^{\circ}\text{F} - 32] * (5/9)$ . The scatterplots on both the original and transformed scales are provided in Figure 7.10. All the coefficients in the regression model and the labels on the axes change, but the “picture” is still the same. Additionally, all the inferences remain the same – and p-values are unchanged by linear transformations. So linear transformations can be “fun” but really are only useful if they make the coefficients easier to interpret. Here if you like changes in  $^{\circ}\text{C}$  for a 1 year increase, the slope coefficient is 0.029 and if you like the original change in  $^{\circ}\text{F}$  for a 1 year increase, the slope coefficient is 0.052.

```
bozemantemps$meanmaxC <- (bozemantemps$meanmax - 32)*(5/9)
temp3 <- lm(formula = meanmax ~ Year2, data=bozemantemps)
summary(temp1)
```

```
##
## Call:
## lm(formula = meanmax ~ Year, data = bozemantemps)
##
## Residuals:
##      Min       1Q   Median       3Q      Max 
## -3.3779 -0.9300  0.1078  1.1960  5.8698 
## 
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) -47.35123   9.32184  -5.08 1.61e-06  
## Year        0.05244    0.00476   11.02 < 2e-16
```

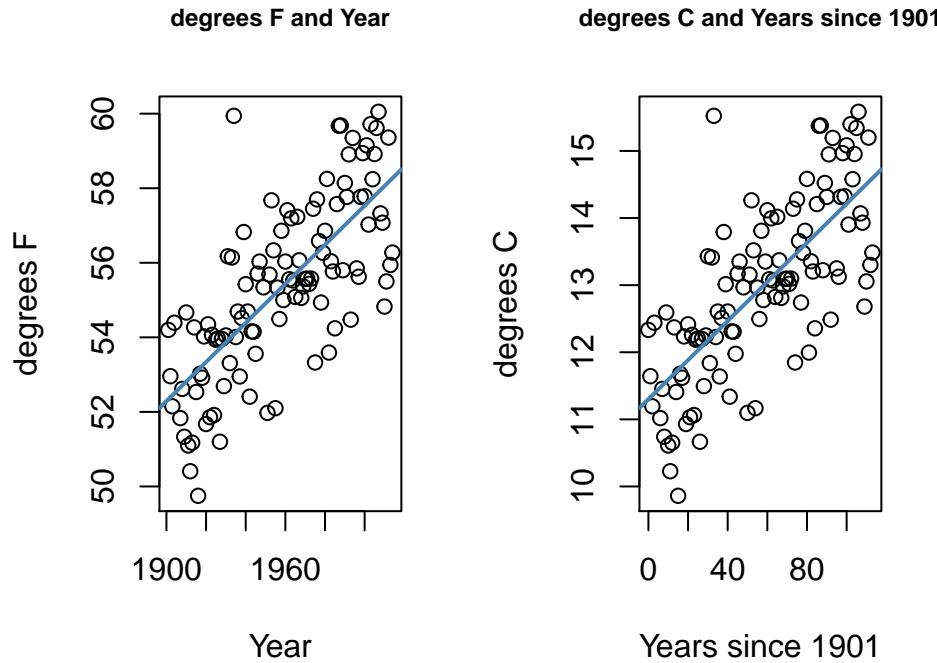


Figure 7.10: Scatterplots of *Temperature* ( $^{\circ}\text{F}$ ) versus *Year* (left) and *Temperature* ( $^{\circ}\text{C}$ ) vs *Years since 1901* (right).

```
##  
## Residual standard error: 1.624 on 107 degrees of freedom  
## Multiple R-squared:  0.5315, Adjusted R-squared:  0.5271  
## F-statistic: 121.4 on 1 and 107 DF,  p-value: < 2.2e-16  
summary(temp3)  
  
##  
## Call:  
## lm(formula = meanmaxC ~ Year2, data = bozemantemps)  
##  
## Residuals:  
##      Min       1Q   Median       3Q      Max  
## -1.8766 -0.5167  0.0599  0.6644  3.2610  
##  
## Coefficients:  
##             Estimate Std. Error t value Pr(>|t|)  
## (Intercept) 11.300703   0.174349  64.82  <2e-16  
## Year2        0.029135   0.002644   11.02  <2e-16  
##  
## Residual standard error: 0.9022 on 107 degrees of freedom  
## Multiple R-squared:  0.5315, Adjusted R-squared:  0.5271  
## F-statistic: 121.4 on 1 and 107 DF,  p-value: < 2.2e-16
```

**Nonlinear transformation functions** are where we apply something more complicated than this shift and scaling, something like  $y_{\text{new}} = f(y)$ , where  $f(\cdot)$  could be a log or power of the original variable  $y$ . When we apply these sorts of transformations, interesting things can happen to our linear models and their problems. Some examples of transformations that are at least occasionally used for transforming the response variable

are provided in Table 7.1, ranging from taking  $y$  to different powers from  $y^{-2}$  to  $y^2$ . Typical transformations used in statistical modeling exist along a gradient of powers of the response variable, defined as  $y^\lambda$  with  $\lambda$  being the power of the transformation of the response variable and  $\lambda = 0$  implying a log-transformation. Except for  $\lambda = 1$ , the transformations are all nonlinear functions of  $y$ .

Table 7.1: Ladder of powers of transformations that are often used in statistical modeling.

Power	Formula	Usage
2	$y^2$	seldom used
1	$y^1 = y$	no change
1/2	$y^{0.5} = \sqrt{y}$	counts and area responses
0	$\log(y)$ natural log of $y$	counts, normality, curves, non-constant variance
-1/2	$y^{-0.5} = 1/\sqrt{y}$	uncommon
-1	$y^{-1} = 1/y$	for ratios
-2	$y^{-2} = 1/y^2$	seldom used

There are even more transformations possible, for example  $y^{0.33}$  is sometimes useful for variables involved in measuring the volume of something. And we can also consider applying any of these transformations to the explanatory variable, and consider using them on both the response and explanatory variables at the same time. The most common application of these ideas is to transform the response variable using the log-transformation, at least as a starting point. In fact, the log-transformation is so commonly used (and miss-used), that we will just focus on its use. It is so commonplace in some fields that some researchers log-transform their data prior to even plotting their results. In other situations, such as when measuring acidity (pH), noise (decibels), or earthquake size (Richter scale), the measurements are already on logarithmic scales.

Actually, we have already analyzed data that benefited from a *log-transformation* – the *log-area burned* vs. *summer temperature* data for Montana. Figure 7.11 compares the relationship between these variables on the original hectares scale and the log-hectares scale.

```
par(mfrow=c(1,2))
plot(hectacres~Temperature, data=mtfires, main="(a)",
      ylab="Hectacres", pch=16)
plot(loghectacres~Temperature, data=mtfires, main="(b)",
      ylab="log-Hectacres", pch=16)
```

The left panel in Figure 7.11 displays a relationship that would be hard fit using SLR – it has a curve and the variance is increasing with increasing temperatures. With a log-transformation of *Hectacres*, the relationship appears to be relatively linear and have constant variance. We considered regression models for this situation previously. This shows at least one situation where a log-transformation of a response variable can linearize a relationship and reduce non-constant variance.

This transformation does not always work to “fix” curvilinear relationships, in fact in some situations it can make the relationship more nonlinear. For example, reconsider the relationship between tree diameter and tree height, which contained some curvature that we could not account for in an SLR. Figure 7.12 shows the original version of the variables and Figure 7.13 shows the same information with the response variable (height) log-transformed.

```
require(spuRs) # install.packages("spuRs")
data(ufc)
require(car)
scatterplot(height.m~dbh.cm, data=ufc[-168,], smooth=T, spread=F,
            main="Tree height vs tree diameter")
scatterplot(log(height.m)~dbh.cm, data=ufc[-168,], smooth=T, spread=F,
            main="log-Tree height vs tree diameter")
```

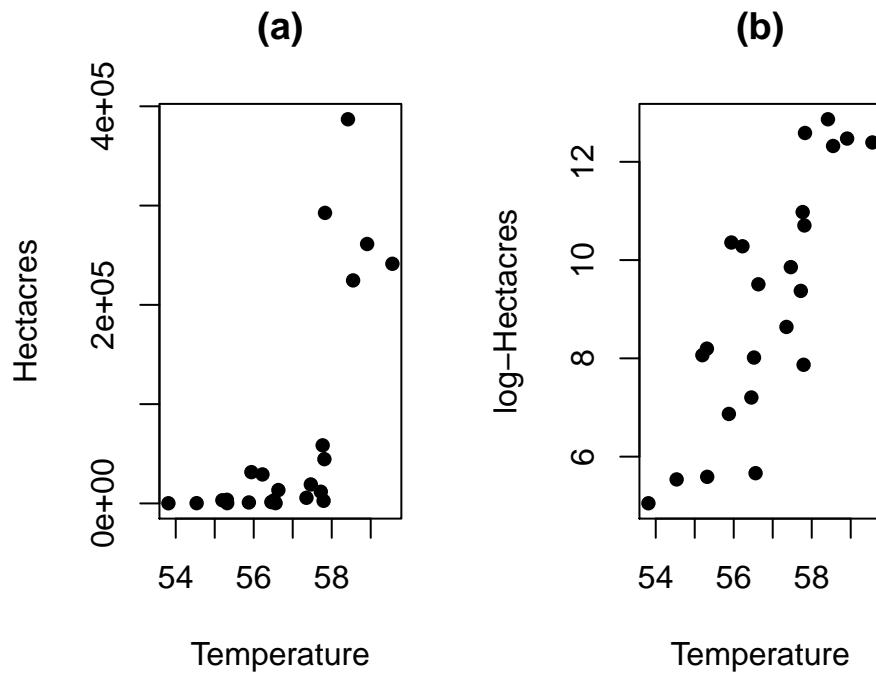


Figure 7.11: Scatterplots of Hectares (a) and log-Hectares (b) vs Temperature.

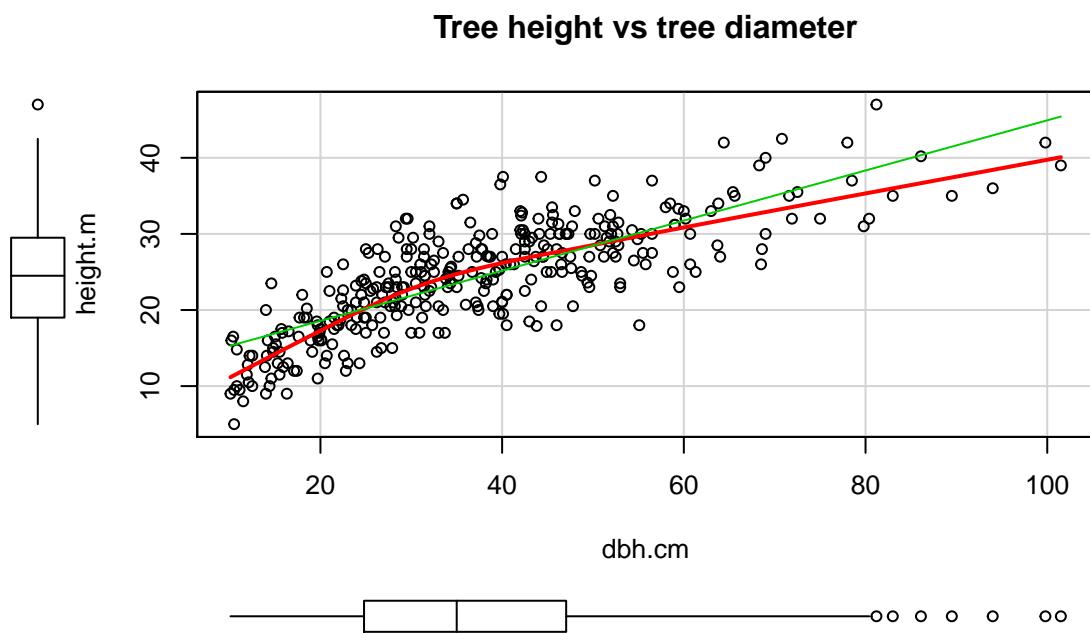


Figure 7.12: Scatterplot of tree height versus tree diameter.

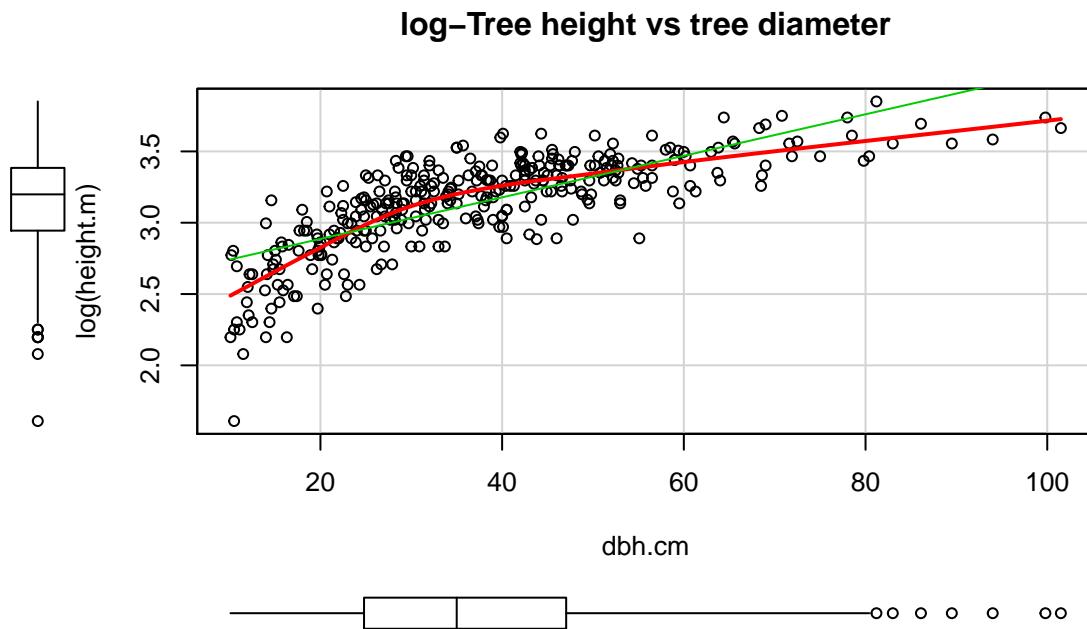


Figure 7.13: Scatterplot of  $\log(\text{tree height})$  versus tree diameter.

Figure 7.13 with the log-transformed response seems to show a more nonlinear relationship and may even have more issues with non-constant variance. This example shows that log-transforming the response variable cannot fix all problems, even though we've seen some researchers assume it can. It is OK to try a transformation but remember to always plot the results to make sure it actually helped and did not make the situation worse.

All is not lost in this situation, we can consider two other potential uses of the log-transformation and see if they can “fix” the relationship up a bit. One option is to apply the transformation to the explanatory variable ( $y \sim \log(x)$ ), displayed in Figure 7.14. If the distribution of the explanatory variable is right skewed (see the boxplot on the x-axis), then consider log-transforming the explanatory variable. This will often reduce the leverage of those most extreme observations which can be useful.

In this situation, it also seems to have been quite successful at linearizing the relationship, leaving some minor non-constant variance, but providing a big improvement from the relationship on the original scale.

The other option, especially when everything else fails, is to apply the log-transformation to both the explanatory and response variables ( $\log(y) \sim \log(x)$ ), as displayed in Figure 7.15. For this example, the transformation seems to be better than the first two options (none and only  $\log(y)$ ), but demonstrates some decreasing variability with larger  $x$  and  $y$  values. It has also created a new and different curve in the relationship. In the end, we might prefer to fit an SLR model to the tree *height* vs *log(diameter)* versions of the variables (Figure 7.14).

```
scatterplot(log(height.m) ~ log(dbh.cm), data=ufc[-168,], smooth=T,
           spread=F, main="log-Tree height vs log-tree diameter")
```

Economists also like to use  $\log(y) \sim \log(x)$  transformations. The log-log transformation tends to linearize certain relationships and has specific interpretations in terms of Economics theory. The log-log transformation shows up in many different disciplines as a way of obtaining a linear relationship on the log-log scale, but different fields discuss it differently. The following example shows a situation where transformations of both  $x$  and  $y$  are required and this double transformation seems to be quite successful in what looks like an initially hopeless situation for our linear modeling approach.

Data were collected in 1988 on the rates of infant mortality (infant deaths per 1,000 live births) and gross

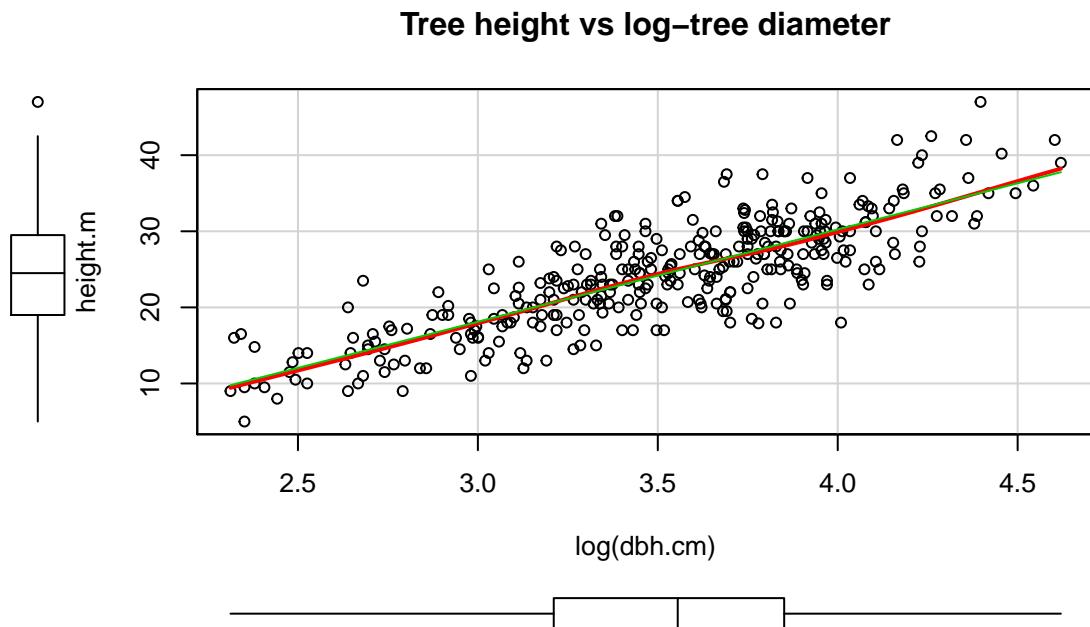


Figure 7.14: Scatterplot of tree height versus log(tree diameter).

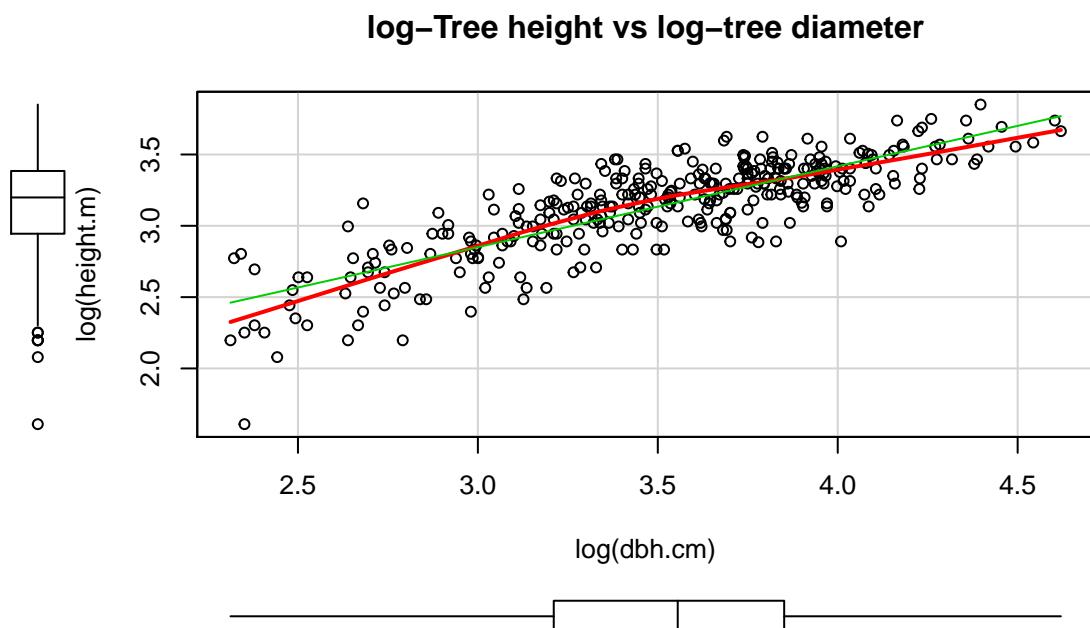


Figure 7.15: Scatterplot of log(tree height) versus log(tree diameter).

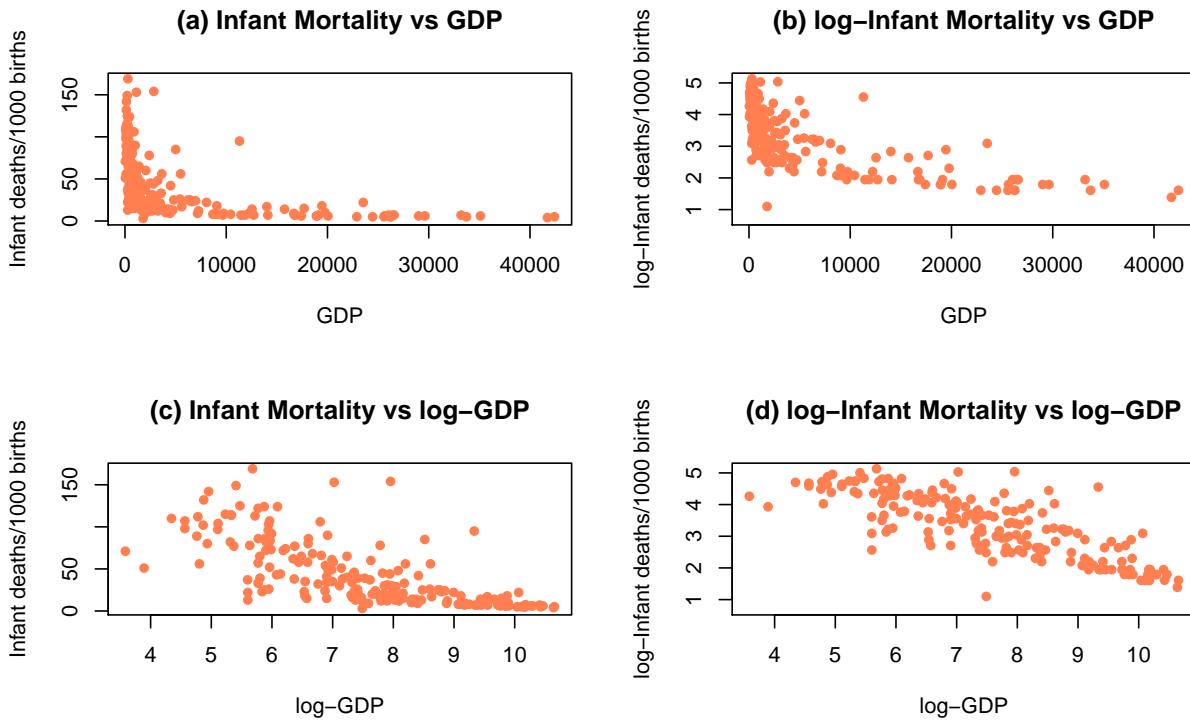


Figure 7.16: Scatterplots of Infant Mortality vs GDP under four different combinations of log-transformations.

domestic product (GDP) per capita (in 1998 US dollars) from  $n = 207$  countries. These data are available from the `car` package (Fox and Weisberg (2016), Fox (2003)) in a data set called `UN`. The four panels in Figure 7.16 show the original relationship and the impacts of log-transforming one or both variables. The only scatterplot that could potentially be modeled using SLR is the lower right panel (d) that shows the relationship between  $\log(\text{infant mortality})$  and  $\log(\text{GDP})$ . In the next section, we will fit models to some of these relationships and use our diagnostic plots to help us assess “success” of the transformations.

Almost all nonlinear transformations assume that the variables are strictly greater than 0. For example, consider what happens when we apply the `log` function to 0 or a negative value in R:

```
log(-1)
```

```
## [1] NaN
log(0)
```

```
## [1] -Inf
```

So be careful to think about the domain of the transformation function before using transformations. For example, when using the log-transformation make sure that the data values are non-zero and positive or you will get some surprises when you go to fit your regression model to a data set with NaNs (not a number) and/or  $-\infty$ 's in it.

Sometimes the log-transformations will not be entirely successful. If the relationship is **monotonic** (strictly positive or strictly negative but not both), then possibly another stop on the ladder of transformations in Table 7.1 might work. If the relationship is not monotonic, then it may be better to consider a more complex regression model that can accommodate the shape in the relationship or to split the data set in two so you can use SLR on each part.

Finally, remember that `log` in statistics and especially in R means the **natural log** ( $\ln$  or  $\log$  base  $e$ ) as you

might see it elsewhere). In these situations, applying `log10` function (which provides log base 10) to the variables would lead to very similar results, but readers may assume you used `ln` if you don't state that you used `log10`. The main thing to remember to do is to be clear when communicating the version you are using. As an example, I (Greenwood) was working with researchers on a study (Dieser and Foreman, 2010) related to impacts of environmental stresses on bacterial survival. The response variable was log-transformed counts and involved smoothed regression lines fit on this scale. I was using natural logs to fit the models and then shared the fitted values from the models and my collaborators back-transformed the results assuming that I had used `log10`. We quickly resolved our differences once we discovered them but this serves as a reminder at how important communication is in group projects – we both said we were working with log-transformations and didn't know we defaulted to different bases. Generally, in statistics, it's safe to assume that everything is log base  $e$  unless otherwise specified.

## 7.6 Transformations part II: Impacts on SLR interpretations: $\log(y)$ , $\log(x)$ , & both $\log(y)$ & $\log(x)$

The previous attempts to linearize relationships imply a desire to be able to fit SLR models. The *log*-transformations, when successful, provide the potential to apply our SLR model. There are then two options for interpretations: you can either interpret the model on the transformed scale or you can translate the SLR model on the transformed scale back to the original scale of the variables. It ends up that *log*-transformations have special interpretations on the original scales depending on whether the *log* was applied to the response variable, the explanatory variable, or both.

### Scenario 1: $\log(y)$ vs $x$ model:

First consider the  $\log(y) \sim x$  situations where the estimated model is of the form  $\widehat{\log(y)} = b_0 + b_1x$ . When only the response is *log*-transformed, some people call this a *semi-log model*. But many researchers will use this model without any special considerations, as long as it provides a situation where the SLR assumptions are reasonably well-satisfied. To understand the properties and eventually the interpretation of transformed-variables models, we need to try to “reverse” our transformation. If we exponentiate<sup>7</sup> both sides of  $\log(y) = b_0 + b_1x$ , we get:

- $y = \exp(b_0 + b_1x)$ , which can be re-written as
- $y = \exp(b_0) \exp(b_1x)$ . This is based on the rules for `exp()` where  $\exp(a + b) = \exp(a) \exp(b)$ .
- Now consider what happens if we increase  $x$  1 unit, going from  $x$  to  $x + 1$ , providing a new predicted  $y$  that we can call  $y^*$ :  $y^* = \exp(b_0) \exp[b_1(x + 1)]$
- $y^* = \underline{\exp(b_0) \exp(b_1x)}$   $\exp(b_1)$  Now note that the underlined, bold component was the  $y$ -value for  $x$ .
- $y^* = \mathbf{y} \exp(b_1)$  Replace `exp(b0) exp(b1x)` with  $\mathbf{y}$ , the value for  $x$ .

So the difference in fitted values between  $x$  and  $x + 1$  is to multiply the result for  $x$  (that was predicting  $\mathbf{y}$ ) by  $\exp(b_1)$  to get to the predicted result for  $x + 1$  (called  $y^*$ ). We can then use this result to form our  *$\log(y) \sim x$  slope interpretation*: for a 1 unit increase in  $x$ , we observe a multiplicative change of  $\exp(b_1)$  in the response. When we compute a mean on logged variables that are symmetrically distributed (this should occur if our transformation was successful) and then exponentiate the results, the proper interpretation is that the changes are happening in the **median** of the original responses. This is the only time in the course that we will switch our inferences to medians instead of means, and we don't do this because we want to, we do it because it is result of modeling on the  $\log(y)$  scale if successful.

When we are working with regression equations, slopes can either be positive or negative and our interpretations change based on this result to either result in growth ( $b_1 > 0$ ) or decay ( $b_1 < 0$ ) in the responses as the explanatory variable is increased. As an example, consider  $b_1 = 0.4$  and  $\exp(b_1) = \exp(0.4) = 1.492$ . There are a couple of ways to interpret this on the original scale of the response variable  $y$ :

---

<sup>7</sup>Note `exp(x)` is the same as  $e^{(x)}$  but easier to read in-line and `exp()` is the R function name to execute this calculation.

For  $b_1 > 0$ :

1. For a 1 unit increase in  $x$ , the multiplicative change in the median of  $y$  is 1.492.
2. We can convert this into a **percentage increase** by subtracting 1 from  $\exp(0.4)$ ,  $1.492 - 1.0 = 0.492$  and multiplying the result by 100,  $0.492 * 100 = 49.2\%$ . This is interpreted as: For a 1 unit increase in  $x$ , the median of  $y$  increases by 49.2%.

`exp(0.4)`

```
## [1] 1.491825
```

For  $b_1 < 0$ , the change on the *log*-scale is negative and that implies on the original scale that the curve decays to 0. For example, consider  $b_1 = -0.3$  and  $\exp(-0.3) = 0.741$ . Again, there are two versions of the interpretation possible:

1. For a 1 unit increase in  $x$ , the multiplicative change in the median of  $y$  is 0.741.
2. For negative slope coefficients, the percentage decrease is calculated as  $(1 - \exp(b_1)) * 100\%$ . For  $\exp(-0.3) = 0.741$ , this is  $(1 - 0.741) * 100 = 25.9\%$ . This is interpreted as: For a 1 unit increase in  $x$ , the median of  $y$  decreases by 25.9%.

We suspect that you will typically prefer interpretation #1 for both directions but it is important to be able think about the results in terms of **% change of the medians** to make the scale of change more understandable. Some examples will help us see how these ideas can be used in applications.

For the area burned data set, the estimated regression model is  $\widehat{\log(\text{hectares})} = -69.8 + 1.39 \text{ Temp}$ . On the original scale, this implies that the model is  $\text{hectares} = \exp(-69.8) \exp(1.39 \text{ Temp})$ . Figure 7.17 provides the *log(y)* scale version of the model and the model transformed to the original scale of measurement. On the log-hectares scale, the interpretation of the slope is: For a  $1^{\circ}\text{F}$  increase in summer temperature, we expect a  $1.39 \text{ log-hectares}/1^{\circ}\text{F}$  change, on average, in the log-area burned. On the original scale: A  $1^{\circ}\text{F}$  increase in temperature is related to multiplicative change in the median number of hectares burned of  $\exp(1.39) = 4.01$ . That seems like a big rate of growth but the curve does grow rapidly as shown in panel (b), especially for values over  $58^{\circ}\text{F}$  where the area burned is starting to be large. You can think of the multiplicative change here in the following way: the median number of hectares burned is 4 times higher at  $58^{\circ}\text{F}$  than at  $57^{\circ}\text{F}$  and the median area burned is 4 times larger at  $59^{\circ}\text{F}$  than at  $58^{\circ}\text{F}$ ... This can also be interpreted on a % change scale: A  $1^{\circ}\text{F}$  increase in temperature is related to a  $(4.01 - 1) * 100 = 301\%$  increase in the median number of hectares burned.

### Scenario 2: $y$ vs $\log(x)$ model:

When only the explanatory variable is log-transformed, it has a different sort of impact on the regression model interpretation. Effectively we move the percentage change onto the x-scale and modify the first part of our slope interpretation when we consider the results on the original scale. Once again, we will consider the mathematics underlying the changes in the model and then work on applying it to real situations. When the explanatory variable is logged, the estimated regression model is  $y = b_0 + b_1 \log(x)$ . This models the relationship between  $y$  and  $x$  in terms of multiplicative changes in  $x$  having an effect on the average  $y$ . To develop an interpretation on the x-scale (not  $\log(x)$ ), consider the impact of doubling  $x$ . This change will take us from the point  $(x, y = b_0 + b_1 \log(x))$  to the point  $(2x, y^* = b_0 + b_1 \log(2x))$ . Now the impact of doubling  $x$  can be simplified using the rules for logs to be:

- $y^* = b_0 + b_1 \log(2x)$
- $y^* = b_0 + b_1 \log(x) + b_1 \log(2)$  Based on the rules for logs:  $\log(2x) = \log(x) + \log(2)$ .
- $y^* = y + b_1 \log(2)$
- So if we double  $x$ , we change the **mean** of  $y$  by  $b_1 \log(2)$ .

As before, there are couple of ways to interpret these sorts of results,

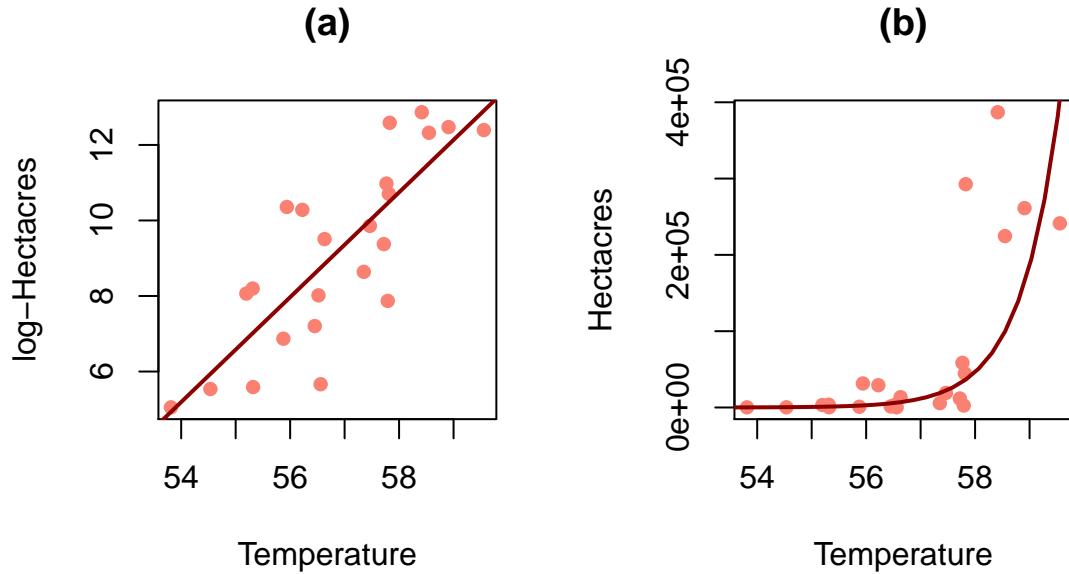


Figure 7.17: Plot of the estimated SLR (a) and implied model for the median on the original Hectares scale (b) for the area burned vs temperature data.

1. ***log-scale interpretation of  $\log(x)$  only model:*** for a 1 log-unit increase in  $x$ , we expect a  $b_1$  unit change in the mean of  $y$  or
2. ***original scale interpretation of  $\log(x)$  only model:*** for a doubling of  $x$ , we expect a  $b_1 \log(2)$  change in the mean of  $y$ . Note that both interpretations are for the mean of the  $y$ 's since we haven't changed the  $y \sim \dots$  part of the model.

While it is not a perfect model (no model is), let's consider the model for  $\text{infant mortality} \sim \log(\text{GDP})$  in order to practice the interpretation using this type of model. This model was estimated to be  $\text{infant mortality} = 168.648 - 16.6 \log(\text{GDP})$ . The first (simplest) interpretation of the slope coefficient is: For a 1 log-dollar increase in GDP per capita, we expect infant mortality to change, on average, by  $-16.6$  deaths/1000 births per log-dollar. The second interpretation is on the original GDP scale: For a doubling of GDP, we expect infant mortality to change, on average, by  $-16.6 \log(2) = -11.51$  deaths/1000 live births. Or, the mean infant mortality is reduced by 11.51 deaths per 1000 live births for each doubling of GDP. Both versions of the model are displayed in Figure 7.18 – one on the scale the SLR model was fit (panel a) and the other on the original x-scale (panel b) that matches these last interpretations.

```

data(UN)
ID1 <- lm(infant.mortality~log(gdp), data=UN)
summary(ID1)

##
## Call:
## lm(formula = infant.mortality ~ log(gdp), data = UN)
##
## Residuals:
##     Min      1Q  Median      3Q     Max 
## -62.668 -16.798 - 0.452 11.064 117.374 
## 
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) 168.648    1.375 123.48   <2e-16 ***
## log(gdp)   -16.600    0.350 -46.86   <2e-16 ***
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

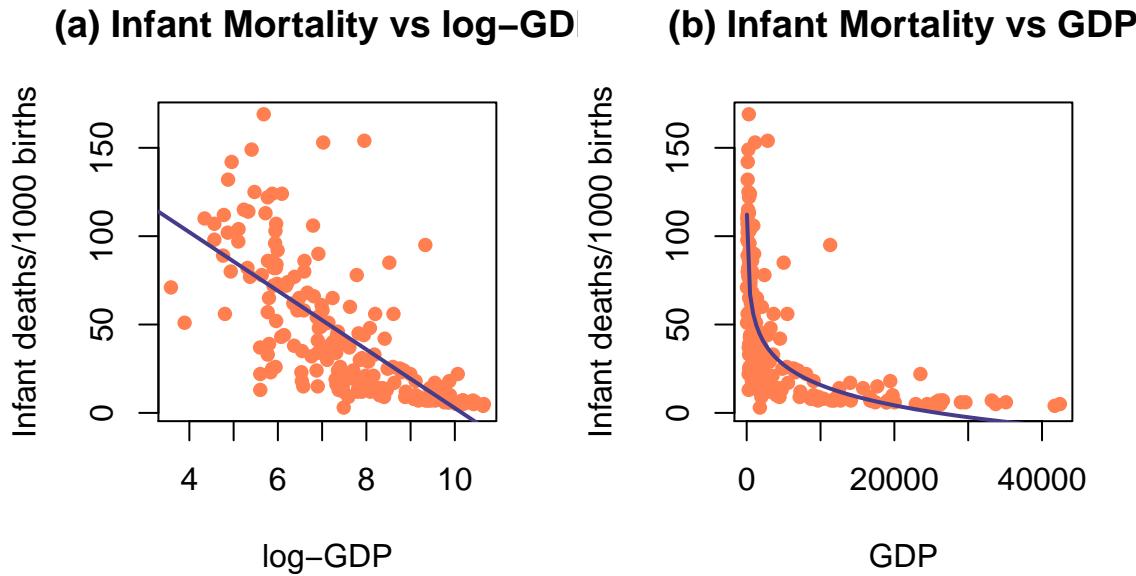


Figure 7.18: Plot of the observations and estimated SLR model (mortality~ log(GDP)) (top) and implied model (bottom) for the infant mortality data.

```

## (Intercept) 168.648      9.084   18.57 <2e-16
## log(gdp)    -16.597     1.180  -14.07 <2e-16
##
## Residual standard error: 27.09 on 191 degrees of freedom
##   (14 observations deleted due to missingness)
## Multiple R-squared:  0.5089, Adjusted R-squared:  0.5063
## F-statistic: 197.9 on 1 and 191 DF, p-value: < 2.2e-16
-16.6*log(2)

## [1] -11.50624

```

It appears that our model does not fit too well and that there might be some non-constant variance so we should check the diagnostic plots (available in Figure 7.19) before we trust any of those previous interpretations.

```

par(mfrow=c(2,2))
plot(ID1)

```

There appear to be issues with outliers and a long right tail violating the normality assumption. There is curvature and non-constant variance in the results as well. There are no influential points, but we are far from happy with this model and will be revisiting this example with the responses also transformed. Remember that the log-transformation of the response can potentially fix non-constant variance, normality, and curvature issues.

### Scenario 3: $\log(y) \sim \log(x)$ model

A final model combines log-transformations of both  $x$  and  $y$ , combining the interpretations used in the previous two situations. This model is called the ***log-log model*** and in some areas is also called the ***power law model***. The power-law model is usually written as  $y = \beta_0 x^{\beta_1} + \varepsilon$ , where  $y$  is thought to be proportional to  $x$  raised to an estimated power of  $\beta_1$  (linear if  $\beta_1 = 1$  and quadratic if  $\beta_1 = 2$ ). It is one of the models that has been used in Geomorphology to model the shape of glaciated valley elevation profiles (that classic U-shape that comes with glacier eroded mountain valleys). If you ignore the error term, it is possible to estimate the power-law model using our SLR approach. Consider the log-transformation of both sides of this

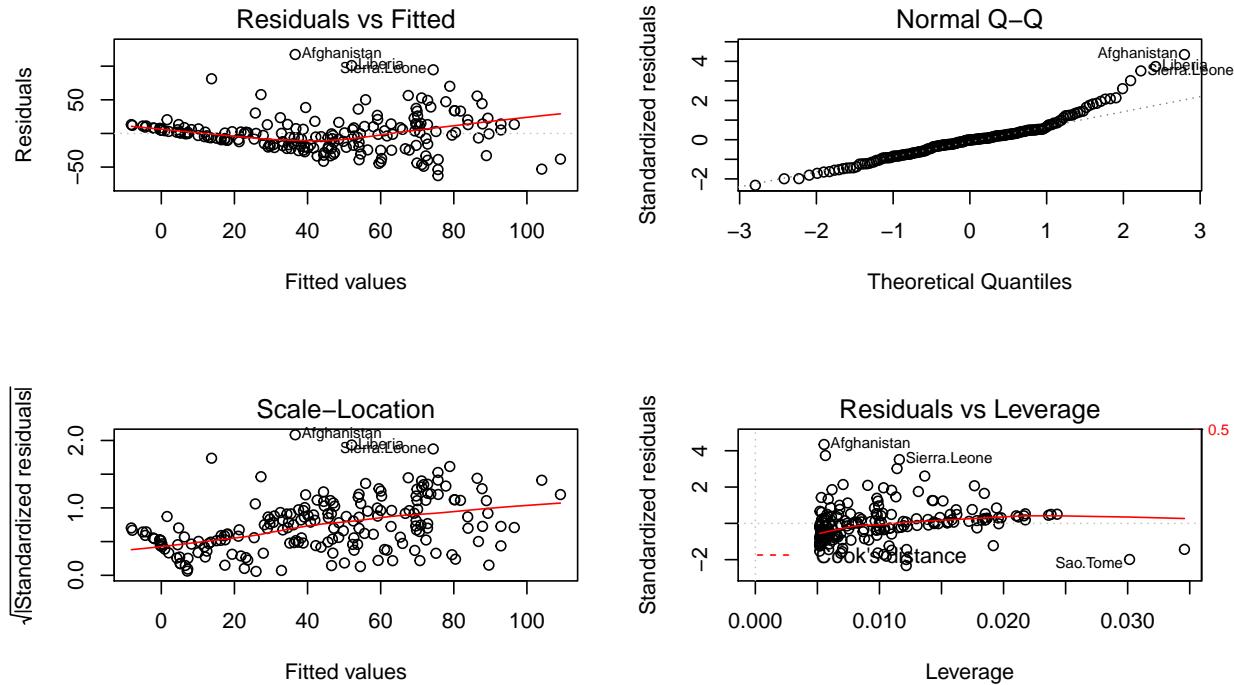


Figure 7.19: Diagnostics plots of the infant mortality model with  $\log(\text{GDP})$ .

equation starting with the power-law version:

- $\log(y) = \log(\beta_0 x^{\beta_1})$
- $\log(y) = \log(\beta_0) + \log(x^{\beta_1})$  *Based on the rules for logs:  $\log(ab) = \log(a) + \log(b)$*
- $\log(y) = \log(\beta_0) + \beta_1 \log(x)$  *Based on the rules for logs:  $\log(x^b) = b \log(x)$*

So other than  $\log(\beta_0)$  in the model, this looks just like our regular SLR model with  $x$  and  $y$  both log-transformed. The slope coefficient for  $\log(x)$  is the power coefficient in the original power law model and determines whether the relationship between the original  $x$  and  $y$  in  $y = \beta_0 x^{\beta_1}$  is linear ( $y = \beta_0 x^1$ ) or quadratic ( $y = \beta_0 x^2$ ) or even quartic ( $y = \beta_0 x^4$ ) in some really heavily glacier carved U-shaped valleys. There are some issues with “ignoring the errors” in using SLR to estimate these models (Greenwood and Humphrey, 2002) but it is still a pretty powerful result to be able to estimate the coefficients in ( $y = \beta_0 x^{\beta_1}$ ) using SLR.

We don’t typically use the previous ideas to interpret the typical log-log regression model, instead we combine our two previous interpretation techniques to generate our interpretation. We need to work out the mathematics of doubling  $x$  and the changes in  $y$  starting with the  **$\log(y) \sim \log(x)$  model** that we would get out of fitting the SLR with both variables log-transformed:

- $\log(y) = b_0 + b_1 \log(x)$
- $y = \exp(b_0 + b_1 \log(x))$  *Exponentiate both sides.*
- $y = \exp(b_0) \exp(b_1 \log(x)) = \exp(b_0) x^{b_1}$  *Rules for exponents and logs, simplifying.*

Now we can consider the impacts of doubling  $x$  on  $y$ , going from  $(x, y = \exp(b_0)x^{b_1})$  to  $(2x, y^*)$  with

- $y^* = \exp(b_0)(2x)^{b_1}$
- $y^* = \exp(b_0)2^{b_1}x^{b_1} = 2^{b_1} \exp(b_0)x^{b_1} = 2^{b_1}y$

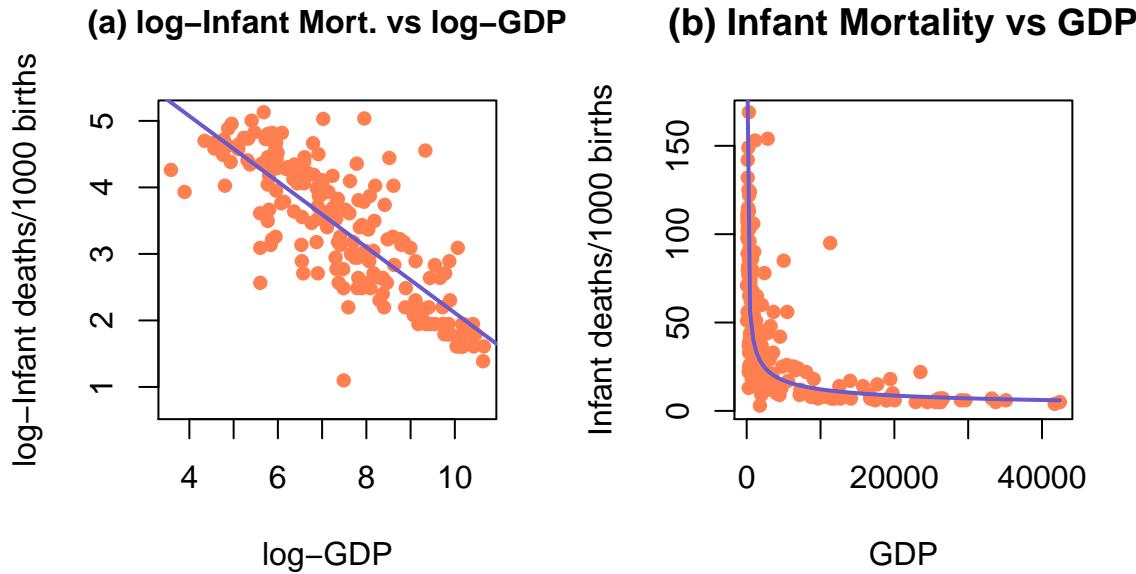


Figure 7.20: Plot of the observations and estimated SLR model  $\log(\text{mortality}) \sim \log(\text{GDP})$  (left) and implied model (right) for the infant mortality data.

So doubling  $x$  leads to a multiplicative change in the median of  $y$  of  $2^{b_1}$ .

Let's apply this idea to the GDP and infant mortality data where a  $\log(x) \sim \log(y)$  transformation actually made the resulting relationship look like it might meet the SLR assumptions. The regression line in Figure 7.20 actually looks pretty good on both the estimated log-log scale (panel a) and on the original scale (panel b) as it captures the severe nonlinearity in the relationship between the two variables.

```
ID2 <- lm(log(infant.mortality) ~ log(gdp), data=UN)
summary(ID2)
```

```
##
## Call:
## lm(formula = log(infant.mortality) ~ log(gdp), data = UN)
##
## Residuals:
##      Min       1Q   Median       3Q      Max
## -2.25334 -0.39580 -0.05258  0.35598  2.11187
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) 7.04520   0.19914  35.38  <2e-16
## log(gdp)    -0.49320   0.02586 -19.07  <2e-16
##
## Residual standard error: 0.5938 on 191 degrees of freedom
## (14 observations deleted due to missingness)
## Multiple R-squared:  0.6556, Adjusted R-squared:  0.6538
## F-statistic: 363.7 on 1 and 191 DF, p-value: < 2.2e-16
```

The estimated regression model is  $\widehat{\log(\text{infantmortality})} = 7.0452 - 0.493 \log(\text{GDP})$ . The slope coefficient can be interpreted two ways.

1. **On the log-log scale:** For a 1 log-dollar increase in  $GDP$ , we expect, on average, a change of  $-0.493$

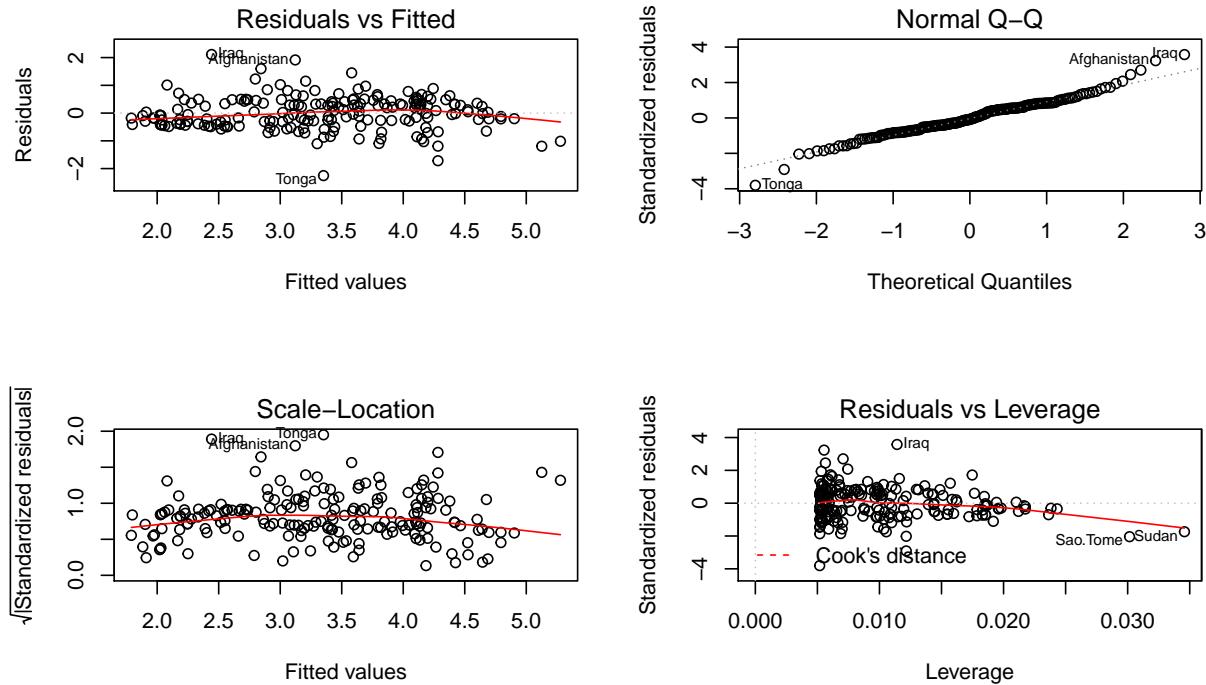


Figure 7.21: Diagnostic plots for the log-log infant mortality model.

$\log(\text{deaths}/1000 \text{ live births})$  in *infant mortality*.

2. **On the original scale:** For a doubling of  $GDP$ , we expect a  $2^{b_1} = 2^{-0.493} = 0.7105$  multiplicative change in the median *infant mortality*. That is a 28.95% decrease in the median *infant mortality* for each doubling of  $GDP$ .

The diagnostics of the log-log SLR model (Figure 7.21) show that the assumptions are fairly reasonably met although the tails of the residuals are a little heavy (more spread out than a normal distribution) and there might still be a little pattern remaining in the residuals vs fitted. There are no influential points to be concerned about in this situation.

While we will not revisit this at all except in the case-studies in Chapter 9, log-transformations can be applied to the response variable in ONE and TWO-WAY ANOVA models when we are concerned about non-constant variance and non-normality issues. The remaining methods in this chapter return to SLR and assuming that the model assumptions have been met. In fact, the methods in Section 7.7 are some of the most sensitive results to violations of the assumptions that we will explore.

## 7.7 Confidence interval for the mean and prediction intervals for a new observation

Figure 7.6 provided a term-plot of the estimated regression line and a shaded area surrounding the estimated regression equation. Those shaded areas are based on connecting the dots on 95% confidence intervals constructed for the true mean  $y$  value across all the  $x$ -values. To formalize this idea, consider a specific

value of  $x$ , and call it  $\mathbf{x}_\nu$  (pronounced **x-new**<sup>8</sup>). Then the true mean response for this **subpopulation** (a subpopulation is all observations we could obtain at  $\mathbf{x} = \mathbf{x}_\nu$ ) is given by  $E(\mathbf{Y}) = \mu_\nu = \beta_0 + \beta_1 x_\nu$ . To estimate the mean response at  $\mathbf{x}_\nu$ , we plug  $\mathbf{x}_\nu$  into the estimated regression equation:

$$\hat{\mu}_\nu = b_0 + b_1 x_\nu.$$

To form the confidence interval, we appeal to our standard formula of  $\text{estimate} \mp t^* \text{SE}_{\text{estimate}}$ . The **standard error for the estimated mean at any x-value**, denoted  $\text{SE}_{\hat{\mu}_\nu}$ , can be calculated as

$$\text{SE}_{\hat{\mu}_\nu} = \sqrt{\text{SE}_{b_1}^2 (x_\nu - \bar{x})^2 + \frac{\hat{\sigma}^2}{n}}$$

where  $\hat{\sigma}^2$  is the squared residual standard error. This formula combines the variability in the slope estimate,  $\text{SE}_{b_1}$ , scaled based on the distance of  $x_\nu$  from  $\bar{x}$  and the variability around the regression line,  $\hat{\sigma}^2$ . Fortunately, we'll use R's **predict** function to provide these results for us and avoid doing this calculation by hand most of the time. The **confidence interval for  $\mu_\nu$** , the population mean response at  $x_\nu$ , is

$$\hat{\mu}_\nu \mp t^*_{n-2} \text{SE}_{\hat{\mu}_\nu}.$$

In application, these intervals get wider the further we go from the mean of the  $x$ 's. These have interpretations that are exactly like those for the y-intercept:

- For an x-value of  $\mathbf{x}_\nu$ , we are \_\_\_\_% confident that the true mean of  $\mathbf{y}$  is between **LL** and **UL** [**units of y**].

It is also useful to remember that this interpretation applies individually to every  $x$  displayed in term-plots.

- A second type of interval in this situation takes on a more challenging task – to place an interval on where we think a new observation will fall, called a **prediction interval** (PI). This PI will need to be much wider than the CI for the mean since we need to account for both the uncertainty in the mean and the randomness in sampling a new observation from the normal distribution centered at the true mean for  $x_\nu$ . The interval is centered at the estimated regression line (where else could we center it?) with the estimate denoted as  $\hat{y}_\nu$  to help us see that this interval is for a **new y** at this x-value. The  $\text{SE}_{\hat{y}_\nu}$  incorporates the core of the previous SE calculation and adds in the variability of a new observation in  $\hat{\sigma}^2$ :

$$\text{SE}_{\hat{y}_\nu} = \sqrt{\text{SE}_{b_1}^2 (x_\nu - \bar{x})^2 + \frac{\hat{\sigma}^2}{n} + \hat{\sigma}^2} = \sqrt{\text{SE}_{\hat{\mu}_\nu}^2 + \hat{\sigma}^2}$$

The \_\_\_\_% PI is calculated as

$$\hat{y}_\nu \mp t^*_{n-2} \text{SE}_{\hat{y}_\nu}$$

and interpreted as:

- We are \_\_\_\_% sure that a new observation at  $\mathbf{x}_\nu$  will be between **LL** and **UL** [**units of y**]. Since  $\text{SE}_{\hat{y}_\nu} > \text{SE}_{\hat{\mu}_\nu}$ , the **PI will always be wider than the CI**. Additionally, if either the SE for the slope or the residual variance increases, both intervals would get wider.

---

<sup>8</sup>This silly nomenclature was adopted from De Veaux and Bock (2001) *Stats: Data and Models* text. If you find this too cheesy, you can just call it x-vee.

As in confidence intervals, we assume that a 95% PI “succeeds” – contains the new observation – in 95% of applications of the methods and fails the other 5% of the time. Remember that for any interval estimate, the true value is either in the interval or it is not and our confidence level essentially sets our failure rate! Because PIs push into the tails of the assumed distribution of the responses these methods are very sensitive to violations of assumptions so we need to be especially certain that we have met the assumptions to trust these results will work as advertised (at the *nominal* (specified) level).

There are two ways to explore CIs for the mean and PIs for a new observation. The first is to focus on a specific x-value of interest. The second is to plot the results for all  $x$ 's. To do both of these, but especially to make plots, we want to learn to use the `predict` function. It can either produce the estimate for a particular  $x_\nu$  and the  $SE_{\hat{\mu}_\nu}$  or we can get it to directly calculate the CI and PI. The first way to use it is `predict(modelname, se.fit=T)` which will provide fitted values and  $SE_{\hat{\mu}_\nu}$  for all observed  $x$ 's. We can then use the  $SE_{\hat{\mu}_\nu}$  to calculate  $SE_{\hat{y}_\nu}$  and form our own PIs. If you want CIs, run `predict(modelname, interval= "confidence")`; if you want PIs, run `predict(modelname, interval="prediction")`. If you want to do prediction at an x-value that was not in the original observations, add the option `newdata=data.frame(XVARIABLENAME_FROM_ORIGINAL_MODEL=xnu)` to the `predict` function call.

Some examples of using the `predict` function follow<sup>9</sup>. For example, it might be interesting to use the regression model to find a 95% CI and PI for the *Beers* vs *BAC* study for a student who would consume 8 beers. Four different applications of the `predict` function follow. Note that `lwr` and `upr` in the output depend on what we requested. The first use of `predict` just returns the estimated mean for 8 beers:

```
m1 <- lm(BAC~Beers, data=BB)
predict(m1, newdata=data.frame(Beers=8))
```

```
##           1
## 0.1310095
```

By turning on the `se.fit=T` option, we also get the SE for the confidence interval and degrees of freedom. Note that elements returned are labelled as `$fit`, `$se.fit`, etc.

```
predict(m1, newdata=data.frame(Beers=8), se.fit=T)
```

```
## $fit
##           1
## 0.1310095
##
## $se.fit
## [1] 0.009204354
##
## $df
## [1] 14
##
## $residual.scale
## [1] 0.02044095
```

Instead of using the components of the intervals to make them, we can also directly request the CI or PI using the `interval=...` option, as in the following two lines of code.

```
predict(m1, newdata=data.frame(Beers=8), interval="confidence")
```

```
##       fit      lwr      upr
## 1 0.1310095 0.1112681 0.1507509
predict(m1, newdata=data.frame(Beers=8), interval="prediction")
```

```
##       fit      lwr      upr
## 1 0.1310095 0.08292834 0.1790906
```

---

<sup>9</sup>I have suppressed some of the plotting code in this and the last chapter to make “pretty” pictures - which you probably are happy to not see it until you want to make a pretty plot on your own. All the code used is available upon request.

Based on these results, we are 95% confident that the true mean  $BAC$  for 8 beers consumed is between 0.111 and 0.15 grams of alcohol per dL of blood. For a new student drinking 8 beers, we are 95% sure that the observed BAC will be between 0.083 and 0.179 g/dL. You can see from these results that the PI is much wider than the CI – it has to capture a new individual's results 95% of the time which is much harder than trying to capture the mean. For completeness, we should do these same calculations by hand. The `predict(..., se.fit=T)` output provides almost all of the pieces we need to calculate the CI and PI. The `$fit` is the estimate  $\hat{\mu}_\nu = 0.131$ , the `$se.fit` is the SE for the estimate of the mean  $= SE_{\hat{\mu}_\nu} = 0.0092$ , `$df` is  $n - 2 = 16 - 2 = 14$ , and `$residual.scale` is  $\hat{\sigma} = 0.02044$ . So we just need the  $t^*$  multiplier for 95% confidence and 14  $df$ :

```
qt(.975, df=14) # t* multiplier for 95% CI or 95% PI
```

```
## [1] 2.144787
```

The 95% CI for the true mean at  $x_\nu = 8$  is then:

```
0.131 + c(-1,1)*2.1448*0.0092
```

```
## [1] 0.1112678 0.1507322
```

Which matches the previous output quite well.

The 95% PI requires the calculation of  $\sqrt{SE_{\hat{\mu}_\nu}^2 + \hat{\sigma}^2} = \sqrt{(0.0092)^2 + (0.02044)^2} = 0.0224$ .

```
sqrt(0.0092^2 + 0.02044^2)
```

```
## [1] 0.02241503
```

The 95% PI at  $x_\nu = 8$  is

```
0.131 + c(-1,1)*2.1448*0.0224
```

```
## [1] 0.08295648 0.17904352
```

These calculations are “fun” and informative but displaying these results for all x-values is a bit more informative about the performance of the two types of intervals and for results we might expect in this application. The calculations we just performed provide endpoints of both intervals at `Beers=8`. To make this plot, we need to create a sequence of `Beers` values to get other results for, say from 0 to 10, using the `seq` function. The `from` and `to` define the endpoints and the `length.out` defines the resolution of the grid of equally spaced points to create. Here, 30 points is more than enough to make the confidence and prediction intervals from 0 to 10 `Beers`.

```
beerf <- seq(from=0, to=10, length.out=30)
```

Now we can call the `predict` function at all these values stored in `beerf` to get the CIs for all these `Beers` values:

```
BBCI <- data.frame(predict(m1, newdata=data.frame(Beers=beerf),
                           interval="confidence"))
head(BBCI)
```

```
##          fit      lwr      upr
## 1 -0.0127006040 -0.039805351 0.01440414
## 2 -0.0065062033 -0.031996517 0.01898411
## 3 -0.0003118027 -0.024210653 0.02358705
## 4  0.0058825980 -0.016452670 0.02821787
## 5  0.0120769986 -0.008728854 0.03288285
## 6  0.0182713992 -0.001047321 0.03759012
```

And the PIs:

```
BBPI <- data.frame(predict(m1, newdata=data.frame(Beers=beerf),
                           interval="prediction"))
head(BBPI)
```

```
##          fit      lwr      upr
```

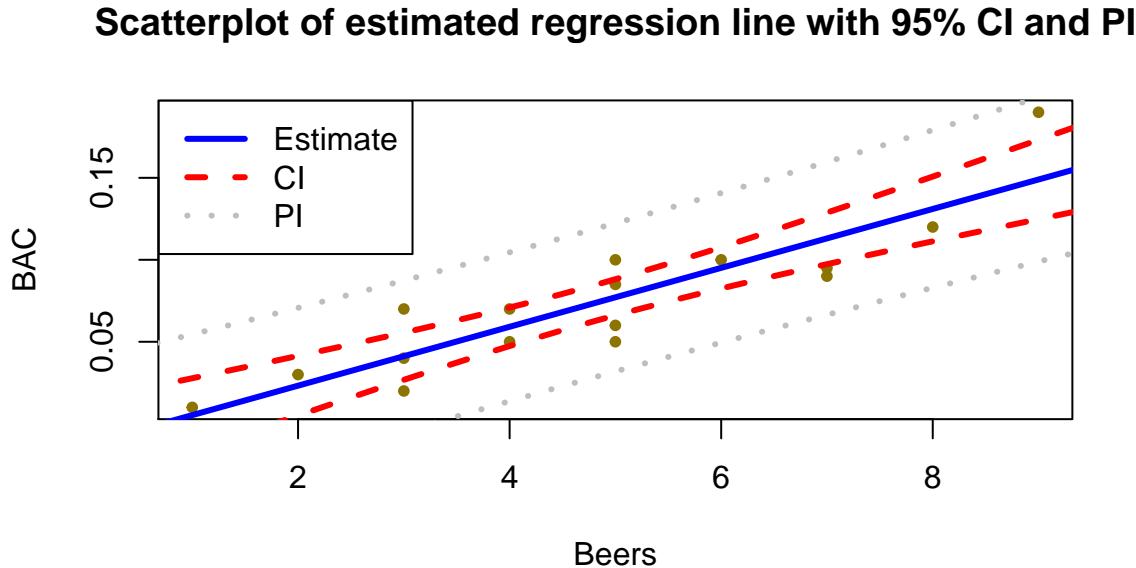


Figure 7.22: Estimated SLR for BAC data with 95% confidence (dashed lines) and 95% prediction (lighter, dotted lines) intervals.

```
## 1 -0.0127006040 -0.06424420 0.03884300
## 2 -0.0065062033 -0.05721943 0.04420702
## 3 -0.0003118027 -0.05024406 0.04962046
## 4 0.0058825980 -0.04332045 0.05508564
## 5 0.0120769986 -0.03645092 0.06060492
## 6 0.0182713992 -0.02963777 0.06618057
```

The rest of the code is just making a scatterplot and adding the five lines with a legend.

```
par(mfrow=c(1,1))
plot(BAC~Beers, data=BB, xlab="Beers", ylab="BAC", pch=20, col="gold4",
     main="Scatterplot of estimated regression line with 95% CI and PI")
lines(beerf, BBCI$fit, col="blue", lwd=3)
lines(beerf, BBCI$lwr, col="red", lty=2, lwd=3)
lines(beerf, BBCI$upr, col="red", lty=2, lwd=3)
lines(beerf, BBPI$lwr, col="grey", lty=3, lwd=3)
lines(beerf, BBPI$upr, col="grey", lty=3, lwd=3)
legend("topleft", c("Estimate","CI","PI"), lwd=3, lty=c(1,2,3),
       col = c("blue","red","grey"))
```

More importantly, note that the CI in Figure 7.22 clearly shows widening as we move further away from the mean of the  $x$ 's to the edges of the observed  $x$ -values. This reflects a decrease in knowledge of the true mean as we move away from the mean of the  $x$ 's. The PI also is widening slightly but not as clearly in this situation. The difference in widths in the two types of intervals becomes extremely clear when they are displayed together.

Similarly, the 95% CI and PIs for the Bozeman yearly average maximum temperatures in Figure 7.23 provide interesting information on the uncertainty in the estimated mean temperature over time. It is also interesting to explore how many of the observations fall within the 95% prediction intervals. The PIs are for new observations, but you can see how the PIs that were constructed to contain almost all the observations in the original data set but not all of them. In fact, only 2 of the 109 observations (1.8%) fall outside the 95% PIs.

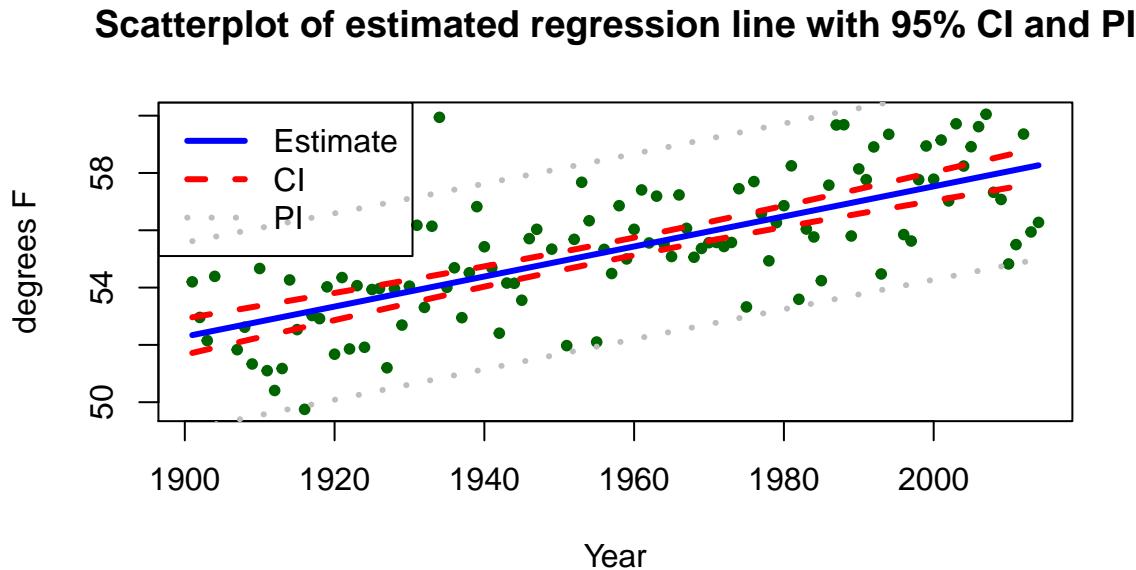


Figure 7.23: Estimated SLR for Bozeman temperature data with 95% confidence (dashed lines) and 95% prediction (lighter, dotted lines) intervals.

Since the PI needs to be concerned with unobserved new observations it makes sense that it might contain more than 95% of the observations used to make it.

We can also use these same methods to do a prediction for the year after the data set ended, 2015 and in 2050:

```
predict(temp1, newdata=data.frame(Year=2015), interval="confidence")
##           fit      lwr      upr
## 1 58.31967 57.7019 58.93744

predict(temp1, newdata=data.frame(Year=2015), interval="prediction")
##           fit      lwr      upr
## 1 58.31967 55.04146 61.59787

predict(temp1, newdata=data.frame(Year=2050), interval="confidence")
##           fit      lwr      upr
## 1 60.15514 59.23631 61.07397

predict(temp1, newdata=data.frame(Year=2050), interval="prediction")
##           fit      lwr      upr
## 1 60.15514 56.80712 63.50316
```

These results tell us that we are 95% confident that the true mean yearly average maximum temperature in 2015 is (or was?) between  $55.04^{\circ}\text{F}$  and  $61.6^{\circ}\text{F}$ . And we are 95% sure that the observed yearly average maximum temperature in 2015 will be (or would have been?) between  $59.2^{\circ}\text{F}$  and  $61.1^{\circ}\text{F}$ . Obviously, 2015 has occurred, but since the data were not published when the data set was downloaded in July 2016, we can probably best treat 2015 as a potential “future” observation. The results for 2050 are clearly for the future mean and a new observation in 2050. Note that up to 2014, no values of this response had been observed above  $60^{\circ}\text{F}$  and the predicted mean in 2050 is over  $60^{\circ}\text{F}$  if the trend persists. It is also easy to criticize the use of this model for 2050 because of its extreme amount of extrapolation.

## 7.8 Chapter summary

In this chapter, we raised our SLR modeling to a new level, considering inference techniques for relationships between two variables. The next chapter will build on these same techniques but add in additional explanatory variables for what we call ***multiple linear regression*** (MLR). For example, in the *Beers vs BAC* study, it would have been useful to control for the weight of the subjects since people of different sizes metabolize alcohol at different rates and body size might explain some of the variability. We still would want to study the effects of beer consumption but also control for the differences in subject's weights. Or if they had studied both male and female students, we might need to change the slope or intercept for each sex, allowing the relationship between *Beers* and *BAC* to change between these groups. That will also be handled using MLR techniques but result in two simple linear regression equations – one for females and one for males. And in this chapter you learned how to interpret SLR models. The next chapter will feel like it is completely new initially but it actually contains very little new material, just more complicated models that use the same concepts. There will be a couple of new issues to consider for MLR and we'll need to learn how to work with categorical variables in a regression setting – but we actually fit linear models with categorical variables in Chapters 3 and 4 so that isn't actually completely new either.

SLR is a simple (thus its name) tool for analyzing the relationship between two quantitative variables. It contains assumptions about the estimated regression line being reasonable and about the distribution of the responses around that line to do inferences for the population regression line. Our diagnostic plots help us to carefully assess those assumptions. If we cannot trust the assumptions, then the estimated line and any inferences for the population are un-trustworthy. Transformations can fix things so that we can use SLR to fit regression models. Transformations can complicate the interpretations on the original, untransformed scale but have minimal impact on the interpretations on the transformed scale. It is important to be careful with the units of the variables, especially when dealing with transformations, as this can lead to big changes in the results depending on which scale (original or transformed) the results are being interpreted on.

## 7.9 Important R code

The main components of the R code used in this chapter follow with the components to modify in red where *y* is a response variable, *x* is an explanatory variable, and the data are in **DATASETNAME**.

- `scatterplot(y~x, data=DATASETNAME, smooth=F)`
  - Requires the `car` package.
  - Provides a scatter plot with a regression line.
  - Turn on `smooth=T` to add a smoothing line to help detect nonlinear relationships.
- `MODELNAME <- lm(y~ x, data=DATASETNAME)`
  - Estimates a regression model using least squares.
- `summary(MODELNAME)`
  - Provides parameter estimates and R-squared (used heavily in Chapter 8 as well).
- `par(mfrow=c(2, 2)); plot(MODELNAME)`
  - Provides four regression diagnostic plots in one plot.
- `confint(MODELNAME, level=0.95)`
  - Provides 95% confidence intervals for the regression model coefficients.
  - Change `level` if you want other confidence levels.
- `plot(allEffects(MODELNAME))`

- Requires the `effects` package.
- Provides a term-plot of the estimated regression line with 95% confidence interval for the mean.
- `DATASETNAME$logy <- log(DATASETNAME$y)`
  - Creates a transformed variable called `logy` – change this to be more specific to your “ $y$ ” or “ $x$ ”.
- `predict(MODELNAME, se.fit=T)`
  - Provides fitted values for all observed  $x$ ’s with SEs for the mean.
- `predict(MODELNAME, newdata=data.frame(x = XNEW), interval="confidence")`
  - Provides fitted value for a specific  $x$  (`XNEW`) with CI for the mean. Replace `x` with name of explanatory variable.
- `predict(MODELNAME, newdata=data.frame(x = XNEW), interval="prediction")`
  - Provides fitted value for a specific  $x$  (`XNEW`) with PI for a new observation. Replace `x` with name of explanatory variable.
- `qt(0.975, df=n - 2)`
  - Gets the  $t^*$  multiplier for making a 95% interval with  $n - 2$  replaced by the sample size – 2.

## 7.10 Practice problems

We will continue with the treadmill data set introduced in Chapter 1 and the SLR fit in the practice problems in Chapter 6. The following code will get you back to where we stopped at the end of Chapter 6:

```
treadmill <- read.csv("http://www.math.montana.edu/courses/s217/documents/treadmill.csv",
                      header=T)
plot(TreadMillOx~RunTime, data=treadmill)
tm <- lm(TreadMillOx~RunTime, data=treadmill)
summary(tm1)
```

7.1. Use the output to test for a linear relationship between treadmill oxygen and run time, writing out all 6+ steps of the hypothesis test. Make sure to address scope of inference and interpret the p-value.

7.2. Form and interpret a 95% confidence interval for the slope coefficient “by hand” using the provided multiplier:

```
qt(0.975, df=29)
```

```
## [1] 2.04523
```

7.3. Use the `confint` function to find a similar confidence interval, checking your previous calculation.

7.4. Use the `predict` function to find fitted values, 95% confidence and 95% prediction intervals for run times of 11 and 16 minutes.

7.5. Interpret the CI and PI for the 11 minute run time.

7.6. Compare the width of either set of CIs and PIs – why are they different? For the two different predictions, why are the intervals wider for 16 minutes than for 11 minutes?

7.7. The Residuals vs Fitted plot considered in Chapter 6 should have suggested slight non-constant variance and maybe a little missed nonlinearity. Perform a log-transformation of the treadmill oxygen response variable and re-fit the SLR model. Remake the diagnostic plots and discuss whether the transformation changed any of them.

7.8 Summarize the  $\log(y) \sim x$  model and interpret the slope coefficient on the transformed and original scales, regardless of the answer to the previous question.



# Chapter 8

## Multiple linear regression

### 8.1 Going from SLR to MLR

In many situations, especially in observational studies, it is unlikely that the system is simple enough to be characterized by a single predictor variable. In experiments, if we randomly assign levels of a predictor variable we can assume that the impacts of other variables cancel out as a direct result of the random assignment. But it is possible even in these experimental situations that we can “improve” our model for the response variable by adding additional predictor variables that explain additional variation in the responses, reducing the amount of unexplained variation. This can allow more precise inferences to be generated from the model. As mentioned previously, it might be useful to know the sex or weight of the subjects in the Beers vs BAC study to account for more of the variation in the responses – this idea motivates our final topic: ***multiple linear regression* (MLR)** models. In observational studies, we can think of a suite of characteristics of observations that might be related to a response variable. For example, consider a study of yearly salaries and variables that might explain the amount people get paid. We might be most interested in seeing how incomes change based on age, but it would be hard to ignore potential differences based on sex and education level. Trying to explain incomes would likely require more than one predictor variable and we wouldn’t be able to explain all the variability in the responses just based on gender and education level, but a model using those variables might still provide some useful information about each component and about age impacts on income after we adjust (control for) sex and education. The extension to MLR allows us to incorporate multiple predictors into a regression model. Geometrically, this is a way of relating many different dimensions (number of  $x$ ’s) to what happened in a single response variable (one dimension).

We start with the same model as in SLR except now we allow  $K$  different  $x$ ’s:

$$y_i = \beta_0 + \beta_1 x_{1i} + \beta_2 x_{2i} + \dots + \beta_K x_{Ki} + \varepsilon_i$$

Note that if  $K = 1$ , we are back to SLR. In the MLR model, there are  $K$  predictors and we still have a y-intercept. The MLR model carries the same assumptions as an SLR model with a couple of slight tweaks specific to MLR (see Section 8.2 for the details on the changes to the validity conditions).

We are able to use the least squares criterion for estimating the regression coefficients in MLR, but the mathematics are beyond the scope of this course. The `lm` function takes care of finding the least squares coefficients using a very sophisticated algorithm<sup>1</sup>. The estimated regression equation it returns is:

$$\hat{y}_i = b_0 + b_1 x_{1i} + b_2 x_{2i} + \dots + b_K x_{Ki}$$

---

<sup>1</sup>If you take advanced applied mathematics courses, you can learn more about the algorithms being used by `lm`. Everyone else only cares about the algorithms when they don’t work – which is usually due to the user’s inputs in these models not the algorithm itself.

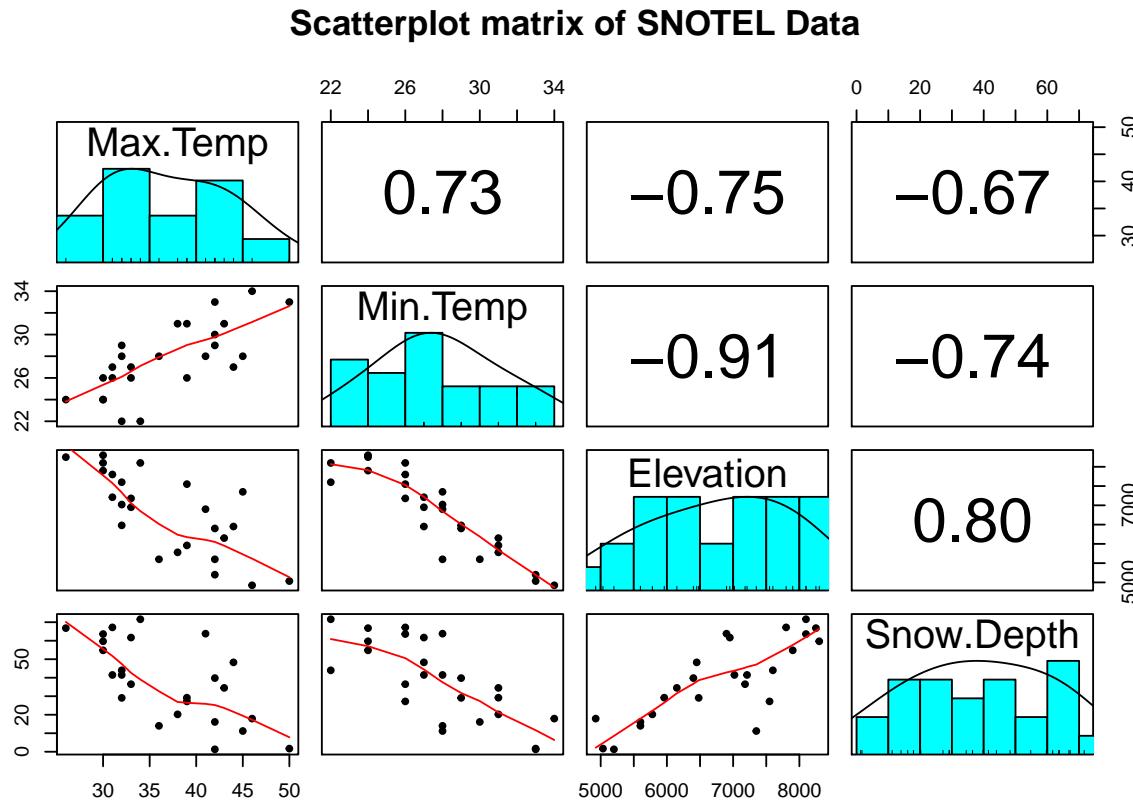


Figure 8.1: Scatterplot matrix of the SNOTEL data.

where each  $b_k$  estimates its corresponding parameter  $\beta_k$ .

An example of snow depths at some high elevation locations on a day in April provides a nice motivation for these methods. A random sample of  $n = 25$  MT locations (from the population of  $N = 85$  at the time) were obtained from the Natural Resources Conversation Service's website (<http://www.wcc.nrcs.usda.gov/snotel/Montana/montana.html>) a few years ago. Information on the snow depth (Snow.Depth) in inches, daily Minimum and Maximum Temperatures (Min.Temp and Max.Temp) in  $^{\circ}\text{F}$  and elevation of the site (Elevation) in feet. A snow science researcher (or spring back-country skier) might be interested in understanding *Snow depth* as a function of *Minimum Temperature*, *Maximum Temperature*, and *Elevation*. One might assume that colder and higher places will have more snow, but using just one of the predictor variables might leave out some important predictive information. The following code loads the data set and makes the scatterplot matrix (Figure 8.1) to allow some preliminary assessment of the pairwise relationships.

```
snotel_s <- read.csv("http://www.math.montana.edu/courses/s217/documents/snotel_s.csv")
snotel2 <- snotel_s[,c(1:2,4:6,3)] #Reorders columns for nicer pairs.panel display
require(psych)
pairs.panels(snotel2[,-c(1:2)], ellipse=F,
             main="Scatterplot matrix of SNOTEL Data")
```

It appears that there are many strong linear relationships between the variables, with *Elevation* and *Snow Depth* having the largest magnitude,  $r = 0.80$ . Higher temperatures seem to be associated with less snow - not a big surprise so far! There might be an outlier at an elevation of 7400 feet and a snow depth below 10 inches that we should explore further.

A new issue arises in attempting to build MLR models called ***multicollinearity***. Again, it is a not surprise

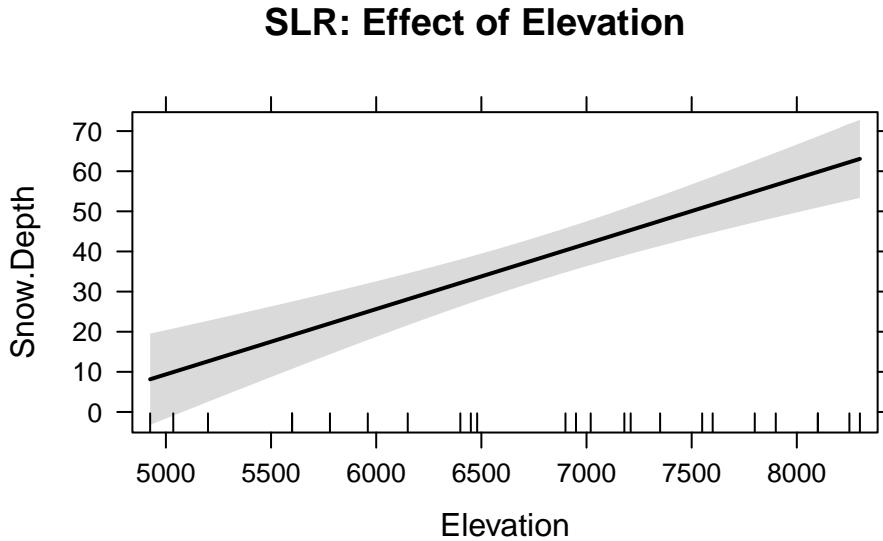


Figure 8.2: Plot of estimated SLR model for Snow Depth with Elevation as the predictor.

that temperature and elevation are correlated but that creates a problem if we try to put them both into a model to explain snow depth. Is it the elevation, temperature, or the combination of both that matters for getting and retaining more snow? **Correlation between predictor variables** is called multicollinearity and **makes estimation and interpretation of MLR models more complicated than in SLR**. Section 8.5 deals with this issue directly and discusses methods for detecting its presence. For now, remember that in MLR this issue sometimes makes it difficult to disentangle the impacts of different predictor variables on the response when the predictors share information – when they are correlated.

To get familiar with this example, we can start with fitting some potential SLR models and plotting the estimated models. Figure 8.2 contains the result for the SLR using *Elevation* and results for two temperature based models are in Figures 8.3 and 8.4. *Snow Depth* is selected as the obvious response variable both due to skier interest and potential scientific causation (snow can't change elevation but elevation could be the driver of snow deposition and retention).

Based on the model summaries provided below, the three estimated SLR models are:

$$\begin{aligned}\widehat{\text{SnowDepth}}_i &= -72.006 + 0.0163 \text{ Elevation}_i, \\ \widehat{\text{SnowDepth}}_i &= 174.096 - 4.884 \text{ MinTemp}_i, \text{ and} \\ \widehat{\text{SnowDepth}}_i &= 122.672 - 2.284 \text{ MaxTemp}_i.\end{aligned}$$

The plots of the estimated models reinforce our expected results, showing a positive change in *Snow Depth* for higher *Elevations* and negative impacts for increasing temperatures on *Snow Depth*. These plots are made across the observed range<sup>2</sup> of the predictor variable and help us to get a sense of the total impacts of predictors. For example, for elevation in Figure 8.2, the smallest observed value was 4925 feet and the largest was 8300 feet. The regression line goes from estimating a mean snow depth of 8 inches to 63 inches. That gives you some practical idea of the size of the estimated *Snow Depth* change for the changes in *Elevation* observed in the data. Putting this together, we can say that there was around a 55 inch change in predicted snow depths for a close to 3400 foot increase in elevation. This helps make the slope coefficient of 0.0163 in the model more easily understood.

<sup>2</sup>Sometimes the effects plots ignores the edge explanatory observations with the default display. Always check the original variable summaries when considering the range of observed values. The code modifications used to make these plots are a little annoying and can be avoided if you remember to check the original summary statistics or scatterplots for variable extrema.

## SLR: Effect of Min Temp

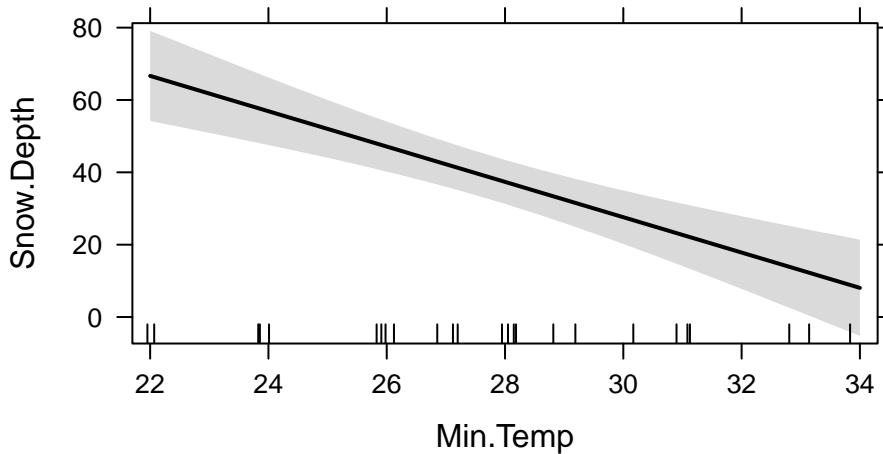


Figure 8.3: Plot of the estimated SLR model using Min Temp as predictor.

Remember that in SLR, the range of  $x$  matters just as much as the units of  $x$  in determining the practical importance and size of the slope coefficient. A value of 0.0163 looks small but is actually at the heart of a pretty good model for predicting snow depth. A one foot change of elevation is “tiny” here relative to changes in the response so the slope coefficient can be small and still amount to big changes in the predicted response across the range of values of  $x$ . If the *Elevation* had been recorded in thousands of feet, then the slope would have been  $0.0163 * 1000 = 16.3$  inches in change in mean *Snow Depth* for a 1000 foot increase in elevation.

The plots of the two estimated temperature models in Figures 8.3 and 8.4 suggest a similar change in the responses over the range of observed temperatures. Those predictors range from  $22^{\circ}\text{F}$  to  $34^{\circ}\text{F}$  (minimum temperature) and from  $26^{\circ}\text{F}$  to  $50^{\circ}\text{F}$  (maximum temperature). This tells us a  $1^{\circ}\text{F}$  increase in either temperature is a greater proportion of the observed range of each predictor than a 1 unit (foot) increase in elevation, so these two variables will generate larger apparent magnitudes of slope coefficients. But having large slope coefficients is no guarantee of a good model – in fact, the elevation model has the highest  $R^2$  value of these three models even though its slope coefficient looks tiny compared to the other models.

```
m1 <- lm(Snow.Depth~Elevation, data=snotel2)
m2 <- lm(Snow.Depth~Min.Temp, data=snotel2)
m3 <- lm(Snow.Depth~Max.Temp, data=snotel2)
require(effects)
plot(allEffects(m1, xlevels=list(Elevation=snotel2$Elevation)),
     main="SLR: Effect of Elevation")
plot(allEffects(m2, xlevels=list(Min.temp=snotel2$Min.Temp)),
     main="SLR: Effect of Min Temp")
plot(allEffects(m3, xlevels=list(Max.Temp=snotel2$Max.Temp)),
     main="SLR: Effect of Max Temp")
```

### SLR: Effect of Max Temp

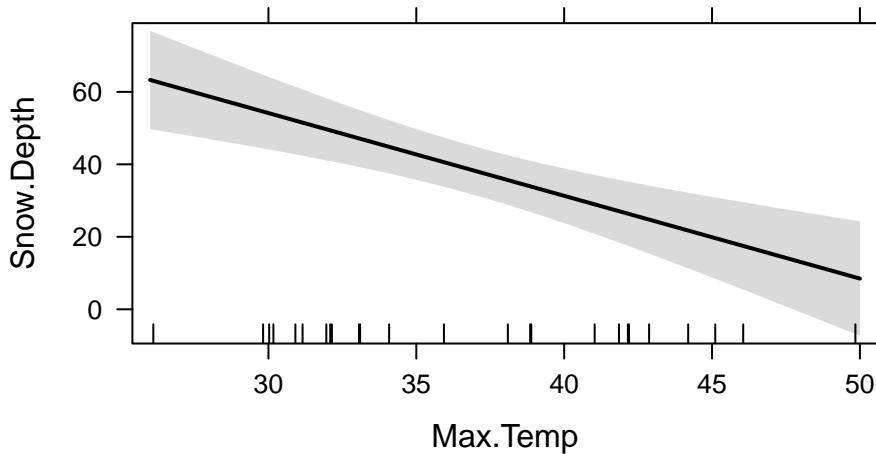


Figure 8.4: Plot of the estimated SLR model using Max Temp as predictor. Note that each of these results (from this plot and from 8.3) are from models with a single predictor variable.

```

summary(m1)

##
## Call:
## lm(formula = Snow.Depth ~ Elevation, data = snotel2)
##
## Residuals:
##    Min     1Q   Median     3Q    Max 
## -36.416 -5.135 -1.767  7.645 23.508 
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) -72.005873 17.712927 -4.065 0.000478 ***
## Elevation     0.016275  0.002579  6.311 1.93e-06 ***
## 
## Residual standard error: 13.27 on 23 degrees of freedom
## Multiple R-squared:  0.634, Adjusted R-squared:  0.618 
## F-statistic: 39.83 on 1 and 23 DF, p-value: 1.933e-06

summary(m2)

##
## Call:
## lm(formula = Snow.Depth ~ Min.Temp, data = snotel2)
##
## Residuals:
##    Min     1Q   Median     3Q    Max 
## -26.156 -11.238  2.810  9.846 26.444 
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) 174.0963   25.5628  6.811 6.04e-07 ***
## Min.Temp     -4.8836     0.9148 -5.339 2.02e-05 ***
## 
## Residual standard error: 14.65 on 23 degrees of freedom

```

```
## Multiple R-squared:  0.5534, Adjusted R-squared:  0.534
## F-statistic:  28.5 on 1 and 23 DF,  p-value: 2.022e-05
summary(m3)

##
## Call:
## lm(formula = Snow.Depth ~ Max.Temp, data = snotel2)
##
## Residuals:
##    Min     1Q   Median     3Q    Max 
## -26.447 -10.367  -4.394  10.042  34.774 
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) 122.6723   19.6380   6.247 2.25e-06  
## Max.Temp     -2.2840    0.5257  -4.345 0.000238  
## 
## Residual standard error: 16.25 on 23 degrees of freedom
## Multiple R-squared:  0.4508, Adjusted R-squared:  0.4269 
## F-statistic: 18.88 on 1 and 23 DF,  p-value: 0.0002385
```

Since all three variables look like they are potentially useful in predicting snow depth, we want to consider if an MLR model might explain more of the variability in *Snow Depth*. To fit an MLR model, we use the same general format as in other topics but with adding “+” between any additional predictors<sup>3</sup> we want to add to the model,  $y \sim x_1 + x_2 + \dots + x_k$ :

```
m4 <- lm(Snow.Depth ~ Elevation + Min.Temp + Max.Temp, data = snotel2)
summary(m4)
```

```
##
## Call:
## lm(formula = Snow.Depth ~ Elevation + Min.Temp + Max.Temp, data = snotel2)
##
## Residuals:
##    Min     1Q   Median     3Q    Max 
## -29.508 -7.679  -3.139  9.627  26.394 
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) -10.506529  99.616286  -0.105  0.9170  
## Elevation    0.012332   0.006536   1.887  0.0731  
## Min.Temp     -0.504970  2.042614  -0.247  0.8071  
## Max.Temp     -0.561892  0.673219  -0.835  0.4133  
## 
## Residual standard error: 13.6 on 21 degrees of freedom
## Multiple R-squared:  0.6485, Adjusted R-squared:  0.5983 
## F-statistic: 12.91 on 3 and 21 DF,  p-value: 5.328e-05
plot(allEffects(m4), main = "MLR model with Elev, Min & Max Temps")
```

Based on the output, the estimated MLR model is

$$\widehat{\text{SnowDepth}}_i = -10.51 + 0.0123 \text{Elevation}_i - 0.505 \text{MinTemp}_i - 0.562 \text{MaxTemp}_i.$$

The direction of the estimated slope coefficients were similar but they all changed in magnitude as compared to the respective SLRs, as seen in the estimated term-plots from the MLR model in Figure 8.5.

---

<sup>3</sup>We used this same notation in the fitting the additive Two-Way ANOVA and this is also additive in terms of these variables. Interaction models are discussed later in the chapter.

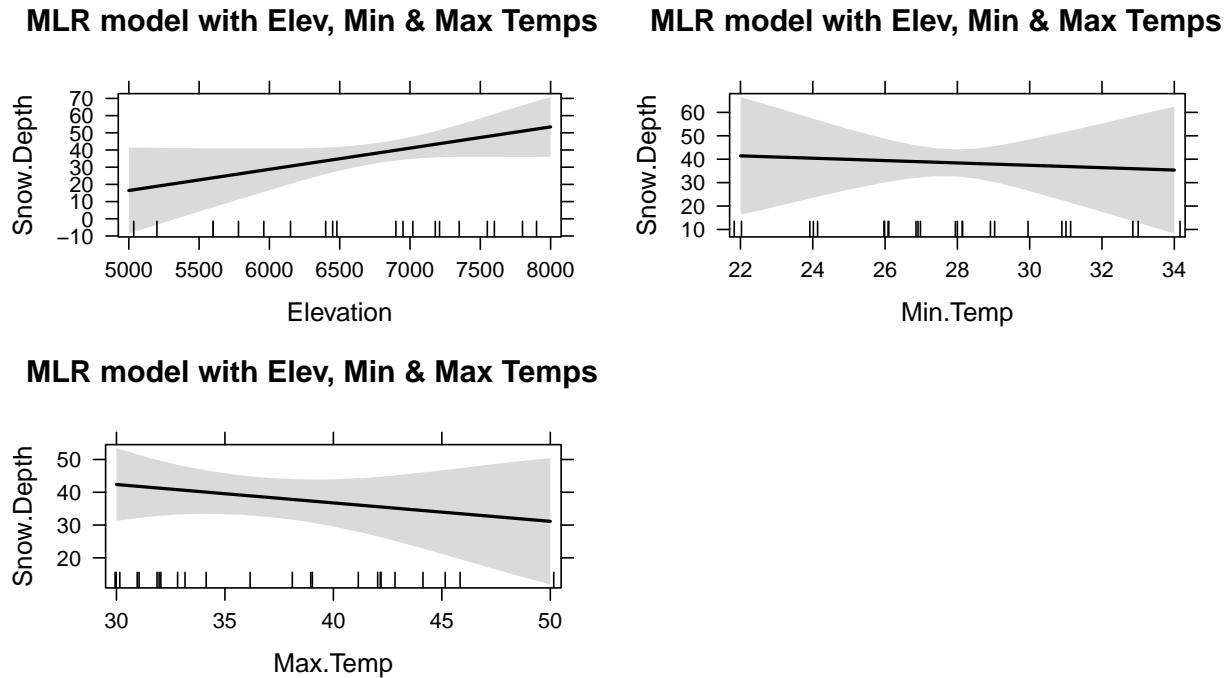


Figure 8.5: Term-plots for the MLR for Snow Depth based on Elevation, Min Temp and Max Temp. Note that the x-axis ranges are different than those used in Figures 8.2, 8.3 and 8.4 for the comparable SLR models.

There are two ways to think about the changes from individual SLR slope coefficients to the similar MLR results.

1. Each term in the MLR is the result for estimating each slope after controlling for the other two variables (and we will always use this interpretation any time we interpret MLR effects). For example, “corrected for” or “adjusted for” the variability that is explained by the temperature variables.
2. Because of multicollinearity in the predictors, the variables might share information that is useful for explaining the variability in the response variable, so the slope coefficients of each predictor get perturbed because the model cannot separate their effects on the response. This issue disappears when the predictors are uncorrelated or even just minimally correlated.

There are some ramifications of multicollinearity in MLR:

1. Adding variables to a model might lead to almost no improvement in the overall variability explained by the model.
2. Adding variables to a model can cause slope coefficients to change signs as well as magnitudes.
3. Adding variables to a model can lead to inflated standard errors for some or all of the coefficients (this is less obvious but is related to the shared information in predictors making it less clear what slope coefficient to use for each variable).
4. In extreme cases of multicollinearity, it may even be impossible to obtain any coefficient estimates.

These seem like pretty serious issues and they are but there are many, many situations where we proceed with MLR even in the presence of potentially correlated predictors. It is likely that you have heard or read about inferences from models that are dealing with this issue – for example, medical studies often report the increased risk of death from some behavior or trait after controlling for sex, age, etc. In many research articles, it is becoming common practice to report the slope for a variable most of interest with it in the model alone (SLR) and in models after adjusting for the other variables that are expected to matter. These

types of results are built with MLR or related multiple-predictor models like MLR.

## 8.2 Validity conditions in MLR

But before we get too excited about any results, we should always assess our validity conditions. For MLR, they are similar to those for SLR:

- **Quantitative variables condition:**
  - The response and all predictors need to be quantitative variables. This condition is relaxed to allow a categorical predictor in two ways in Sections 8.9 and 8.11.
- **Independence of observations:**
  - This assumption is about the responses – we must assume that they were collected in a fashion so that they can be assumed to be independent. This implies that we also have independent random errors.
  - This is not an assumption about the predictor variables!
- **Linearity of relationship (NEW VERSION FOR MLR!):**
  - Linearity is assumed between the response variable and **each** explanatory variable ( $y$  and each  $x$ ).
  - We can check this two ways:
    1. Make plots of the response versus each explanatory variable:
      - Only visual evidence of a curving relationship is a problem here. Transformations of individual explanatory variables or the response are possible.
    2. Examine the Residuals vs Fitted plot:
      - When using MLR, curves in the residuals vs. fitted values suggest a missed curving relationship with at least one predictor variable, but it will not be specific as to which one is non-linear. Revisit the scatterplots to identify the source of the issue.
- **Multicollinearity effects checked for:**
  - Issues here do not mean we cannot proceed with a given model, but it can impact our ability to trust and interpret the estimated terms.
  - Check a scatterplot or correlation matrix to assess the potential for shared information in different predictor variables.
  - Use the diagnostic measure called a ***variance inflation factor (VIF)*** discussed in Section 8.5 (we need to develop some ideas first to understand this measure).
- **Equal (constant) variance:**
  - Same as before since it pertains to the residuals.
- **Normality of residuals:**
  - Same as before since it pertains to the residuals.
- **No influential points:**
  - Leverage is now determined by how unusual a point is for multiple explanatory variables.
  - The **leverage** values in the Residuals vs Leverage plot are scaled to add up to the *degrees of freedom (df) used for the model* which is the number of explanatory variables ( $K$ ) plus 1, so  $K + 1$ .
  - The scale of leverages depends on the complexity of the model through the  $df$  and the sample size.

- The interpretation is still that the larger the leverage value, the more leverage the point has.
- The mean leverage is always  $(\text{model used } df)/n = (K+1)/n$  – so focus on the values with above average leverage.
  - For example, with  $K = 3$  and  $n = 20$ , the average leverage is  $4/20 = 1/5$ .
- High leverage points whose response does not follow the pattern defined by the other observations (now based on patterns for multiple  $x$ 's with the response) will be influential.
- Use the Residual's vs Leverage plot to identify problematic points. Explore further with Cook's D continuing to provide a measure of the influence of each observation.
  - The rules and interpretations for Cook's D are the same as in SLR (over 0.5 is possibly influential and over 1 is definitely influential).

While not a condition for use of the methods, a note about RA and RS is useful here in considering the scope of inference of any results. To make inferences about a population, we need to have a representative sample. If we have randomly assigned levels of treatment variables(s), then we can make causal inferences to subjects like those that we could have observed. And if we both have a representative sample and randomization, we can make causal inferences for the population. It is possible to randomly assign levels of variable(s) to subjects and still collect additional information from other explanatory (sometimes called **control**) variables. The causal interpretations would only be associated with the explanatory variables that were randomly assigned even though the model might contain other variables. Their interpretation still involves noting all the variables included in the model, as demonstrated below. It is even possible to include interactions between randomly assigned variables and other variables – like drug dosage and sex of the subjects. In these cases, causal inference could apply to the treatment levels but noting that the impacts differ based on the non-randomly assigned variable.

For the *Snow Depth* data, the conditions can be assessed as:

- **Quantitative variables condition:**

- These are all met.

- **Independence of observations:**

- The observations are based on a random sample of sites from the population and the sites are spread around the mountains in Montana. Many people would find it to be reasonable to assume that the sites are independent of one another but others would be worried that sites closer together in space might be more similar than they are to far-away observations (this is called **spatial correlation**). I (Greenwood) have been in a heated discussion with statistics colleagues about whether spatial dependency should be considered or if it is valid to ignore it in this sort of situation. It is certainly possible to be concerned about independence of observations here but it takes more advanced statistical methods to actually assess whether there is spatial dependency in these data and even in those models, the first task would be to fit this sort of model and then explore the results.

We need our diagnostic plots to assess the remaining assumptions. The same code as before will provide diagnostic plots. There is some extra code (`par(...)`) added to allow us to add labels to the plots to know which model is being displayed since we have so many to discuss here.

```
par(mfrow=c(2,2), oma=c(0,0,2,0))
plot(m4, sub.caption="Diagnostics for m4")
```

- **Linearity of relationship (NEW VERSION FOR MLR!):**

- Make plots of the response versus each explanatory variable:
  - In Figure 8.1, the plots of each variable versus snow depth do not clearly show any nonlinearity except for a little dip around 7000 feet in the plot vs *Elevation*.
- Examine the Residuals vs Fitted plot in Figure 8.6:

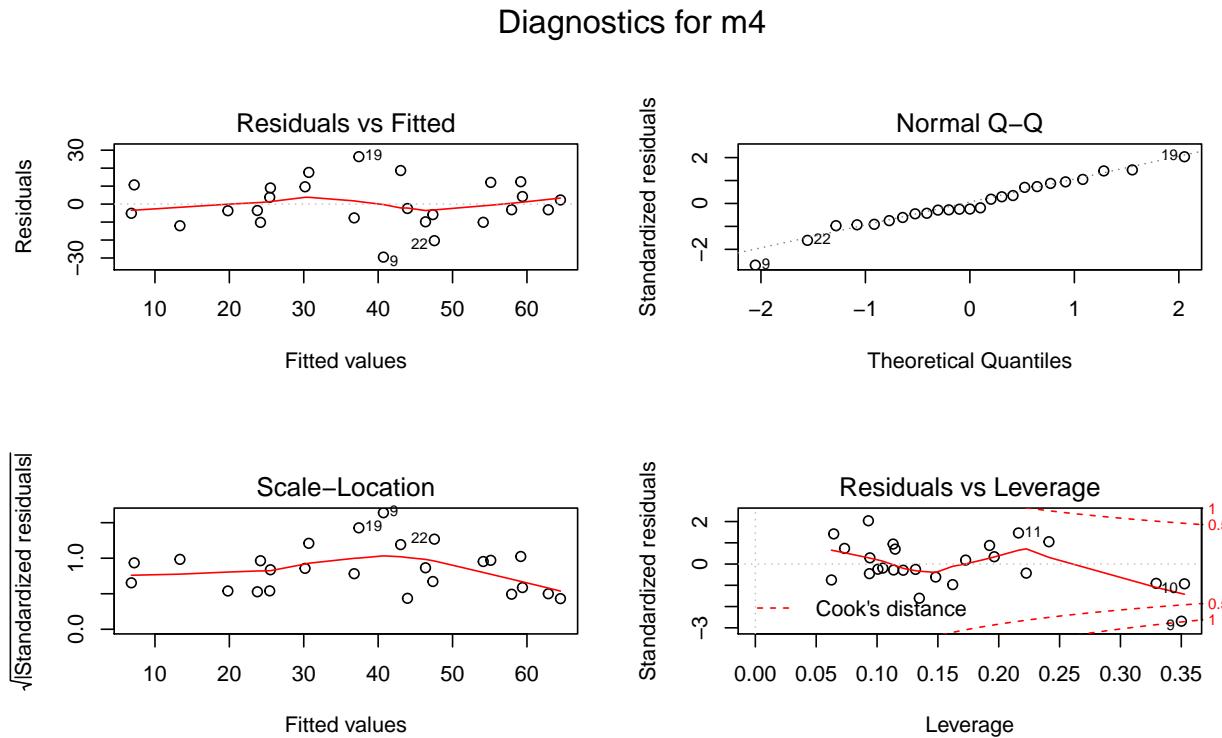


Figure 8.6: Diagnostic plots for model m4:  $\text{Snow.Depth} \sim \text{Elevation} + \text{Min.Temp} + \text{Max.Temp}$

- Generally, there is no clear curvature in the Residuals vs Fitted panel and that would be an acceptable answer. However, there is some pattern in the smoothing line that could suggest a more complicated relationship between at least one predictor and the response. This also resembles the pattern in the *Elevation vs. Snow depth* panel in Figure 8.1 so that might be the source of this “problem”. This suggests that there is the potential to do a little bit better but that it is not unreasonable to proceed on with the MLR, ignoring this little wiggle in the diagnostic plot.

- **Multicollinearity effects checked for:**

- The predictors certainly share information in this application (correlations between -0.67 and -0.91) and multicollinearity looks to be a major concern in being able to understand/separate the impacts of temperatures and elevations on snow depths.
- See Section 8.5 for more on this issue in these data.

- **Equal (constant) variance:**

- While there is a little bit more variability in the middle of the fitted values, this is more an artifact of having a smaller data set with a couple of moderate outliers that fell in the same range of fitted values and maybe a little bit of missed curvature. So there is not too much of an issue with this condition.

- **Normality of residuals:**

- The residuals are quite good in the QQ-plot, showing only a little deviation for observation 9 from a normal distribution and that deviation is extremely minor. Certainly no evidence of a violation of the normality assumption here.

- **No influential points:**

- With  $K = 3$  predictors and  $n = 25$  observations, the average leverage is  $4/25 = 0.16$ . This gives us a scale to interpret the leverage values on the x-axis of the lower right panel of our diagnostic plots.
- There are three higher leverage points (leverages over 0.3) with only one being influential (point 9) with Cook's D close to 1.
  - Note that point 10 had the same leverage but was not influential with Cook's D less than 0.5.
- We can explore both of these points to see how two observations can have the same leverage and different amounts of influence.

The two flagged points, observations 9 and 10 in the data set, are for the sites “Northeast Entrance” (to Yellowstone) and “Combination”. We can use the MLR equation to do some prediction for each observation and calculate residuals to see how far the model’s predictions are from the actual observed values for these sites. For the Northeast Entrance, the *Max.Temp* was 45, the *Min.Temp* was 28, and the *Elevation* was 7350 as you can see in this printout of just the two rows of the data set available by referencing rows 9 and 10 in the bracket from `snotel2`:

```
snotel2[c(9,10),]
```

	ID	Station	Max.Temp	Min.Temp	Elevation	Snow.Depth
## 9	18	Northeast Entrance	45	28	7350	11.2
## 10	53	Combination	36	28	5600	14.0

The estimated *Snow Depth* for the *Northeast Entrance* site (observation 9) is found using the estimated model with

$$\begin{aligned}\widehat{\text{SnowDepth}}_9 &= -10.51 + 0.0123\text{Elevation}_9 - 0.505\text{MinTemp}_9 - 0.562\text{MaxTemp}_9 \\ &= -10.51 + 0.0123 * \mathbf{7350} - 0.505 * \mathbf{28} - 0.562 * \mathbf{45} \\ &= 40.465 \text{ inches},\end{aligned}$$

but the observed snow depth was actually  $y_9 = 11.2$  inches. The observed **residual** is then

$$e_9 = y_9 - \hat{y}_9 = 11.2 - 40.465 = -29.265 \text{ inches.}$$

So the model “misses” the snow depth by over 29 inches with the model suggesting over 40 inches of snow but only 11 inches actually being present<sup>4</sup>.

```
-10.51 + 0.0123*7350 - 0.505*28 - 0.562*45
```

```
## [1] 40.465
11.2 - 40.465
```

```
## [1] -29.265
```

This point is being rated as influential (Cook's D  $\approx 1$ ) with a leverage of nearly 0.35 and a standardized residual (y-axis of Residuals vs. Leverage plot) of nearly -3. This suggests that even with this observation impacting/distorting the slope coefficients (that is what **influence** means), the model is still doing really poorly at fitting this observation. We’ll drop it and re-fit the model in a second to see how the slopes change. First, let’s compare that result to what happened for data point 10 (“Combination”) which was just as high leverage but not identified as influential.

The estimated snow depth for “Combination” is

$$\begin{aligned}\widehat{\text{SnowDepth}}_{10} &= -10.51 + 0.0123\text{Elevation}_{10} - 0.505\text{MinTemp}_{10} - 0.562\text{MaxTemp}_{10} \\ &= -10.51 + 0.0123 * \mathbf{5600} - 0.505 * \mathbf{28} - 0.562 * \mathbf{36} \\ &= 23.998 \text{ inches},\end{aligned}$$

---

<sup>4</sup>Imagine showing up to a ski area expecting a 40 inch base and there only being 11 inches. I’m sure ski areas are always more accurate than this model in their reporting of amounts of snow on the ground...

The observed snow depth here was  $y_{10} = 14.0$  inches so the observed residual is then

$$e_{10} = y_{10} - \hat{y}_{10} = 14.0 - 23.998 = -9.998 \text{ inches.}$$

This results in a standardized residual of around -1. This is still a “miss” but not as glaring as the previous result and also is not having a major impact on the model’s estimated slope coefficients based on the small Cook’s D value.

```
-10.51 + 0.0123*5600 - 0.505*28 - 0.562*36
```

```
## [1] 23.998
14 - 23.998
```

```
## [1] -9.998
```

Note that any predictions using this model presume that it is trustworthy, but the large Cook’s D on one observation suggests we should consider the model after removing that observation. We can re-run the model without the 9th observation using the data set `snote12[-9,]`.

```
m5 <- lm(Snow.Depth~Elevation+Min.Temp+Max.Temp, data=snote12[-9,])
summary(m5)
```

```
##
## Call:
## lm(formula = Snow.Depth ~ Elevation + Min.Temp + Max.Temp, data = snote12[-9,
##     ])
##
## Residuals:
##      Min       1Q   Median       3Q      Max
## -29.2918 -4.9757 -0.9146  5.4292 20.4260
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) -1.424e+02  9.210e+01 -1.546  0.13773
## Elevation    2.141e-02  6.101e-03  3.509  0.00221
## Min.Temp     6.722e-01  1.733e+00  0.388  0.70217
## Max.Temp     5.078e-01  6.486e-01  0.783  0.44283
##
## Residual standard error: 11.29 on 20 degrees of freedom
## Multiple R-squared:  0.7522, Adjusted R-squared:  0.715
## F-statistic: 20.24 on 3 and 20 DF,  p-value: 2.843e-06
plot(allEffects(m5), main="MLR model with NE Ent. Removed")
```

The estimated MLR model with  $n = 24$  after removing the influential “NE Entrance” observation is

$$\widehat{\text{SnowDepth}}_i = -142.4 + 0.0214 \text{ Elevation}_i + 0.672 \text{ MinTemp}_i + 0.508 \text{ MaxTemp}_i.$$

Something unusual has happened here: there is a positive slope for both temperature terms in Figure 8.7 that both contradicts reasonable expectations (warmer temperatures are related to higher snow levels?) and our original SLR results. So what happened? First, removing the influential point has drastically changed the slope coefficients (remember that was the definition of an influential point). Second, when there are predictors that share information, the results can be somewhat unexpected for some or all the predictors when they are all in the model together. Note that the *Elevation* term looks like what we might expect and seems to have a big impact on the predicted *Snow Depths*. So when the temperature variables are included in the model they might be functioning to explain some differences in sites that the *Elevation* term could not explain. This is where our “adjusting for” terminology comes into play. The unusual-looking slopes for the temperature effects can be explained by interpreting them as the estimated change in the response for changes in temperature **after we control for the impacts of elevation**. Suppose that *Elevation* explains most

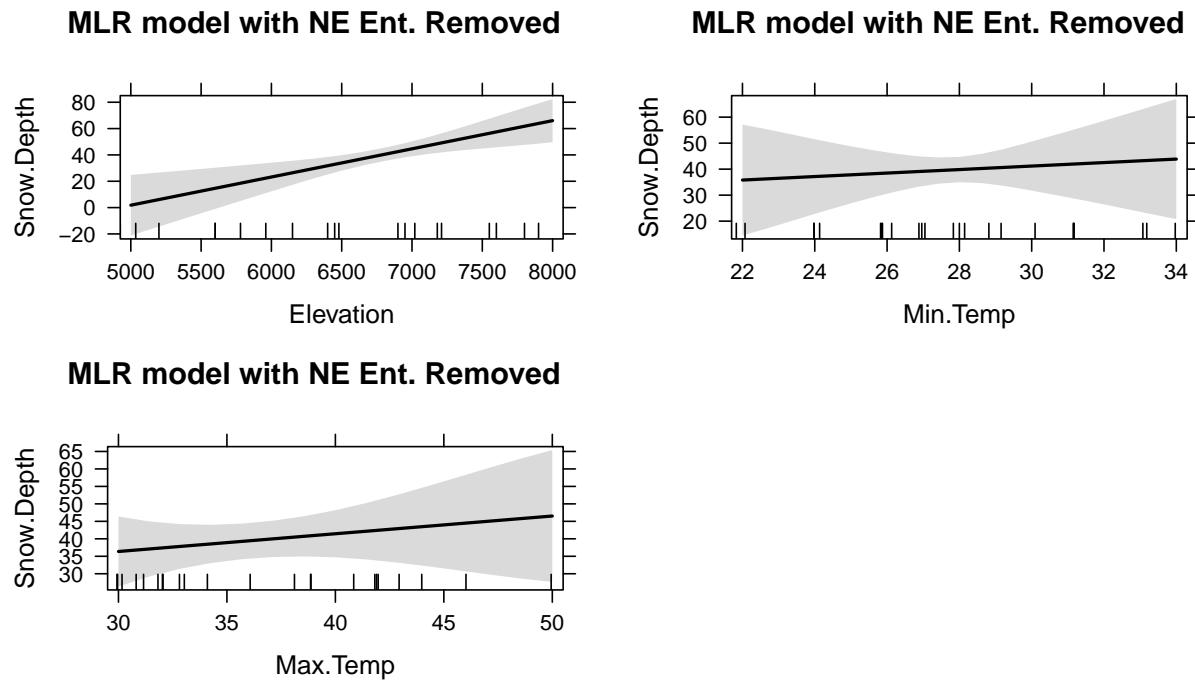


Figure 8.7: Term-plots for the MLR for Snow Depth based on Elevation, Min Temp, and Max Temp with Northeast entrance observation removed from data set ( $n=24$ ).

of the variation in *Snow Depth* except for a few sites where the elevation cannot explain all the variability and the site characteristics happen to show higher temperatures and more snow (or lower temperatures and less snow). This could be because warmer areas might have been hit by a recent snow storm while colder areas might have been missed (this is just one day and subject to spatial and temporal fluctuations in precipitation patterns). Or maybe there is another factor related to having marginally warmer temperatures that are accompanied by more snow (maybe the lower snow sites for each elevation were so steep that they couldn't hold much snow but were also relatively colder?). Thinking about it this way, the temperature model components could provide useful corrections to what *Elevation* is providing in an overall model and explain more variability than any of the variables could alone. It is also possible that the temperature variables are not needed in a model with *Elevation* in it, are just “explaining noise”, and should be removed from the model. Each of the next sections take on various aspects of these issues and eventually lead to a general set of modeling and model selection recommendations to help you work in situations as complicated as this. Exploring the results for this model assumes we trust it and, once again, we need to check diagnostics before getting too focused on any particular results from it.

The Residuals vs. Leverage diagnostic plot in Figure 8.8 for the model fit to the data set without NE Entrance (now  $n = 24$ ) reveals a new point that is somewhat influential (point 22 in the data set has Cook's D  $\approx 0.5$ ). It is for a location called “Bloody [REDACTED]<sup>5</sup> which has a leverage of nearly 0.2 and a standardized residual of nearly -3. This point did not show up as influential in the original version of the data set with the same model but it is now. It also shows up as a potential outlier. As we did before, we can explore it a bit by comparing the model predicted snow depth to the observed snow depth. The predicted snow depth for this site is

$$\widehat{\text{SnowDepth}}_{22} = -142.4 + 0.0214 * 7550 + 0.672 * 26 + 0.508 * 39 = 56.45 \text{ inches.}$$

The observed snow depth was 27.2 inches, so the estimated residual is -39.25 inches. Again, this point is

<sup>5</sup>The site name is redacted to protect the innocence of the reader. More information on this site, located in Beaverhead County, is available at <http://www.wcc.nrcs.usda.gov/nwcc/site?sitenum=355&state=mt>.

### Diagnostics for m5

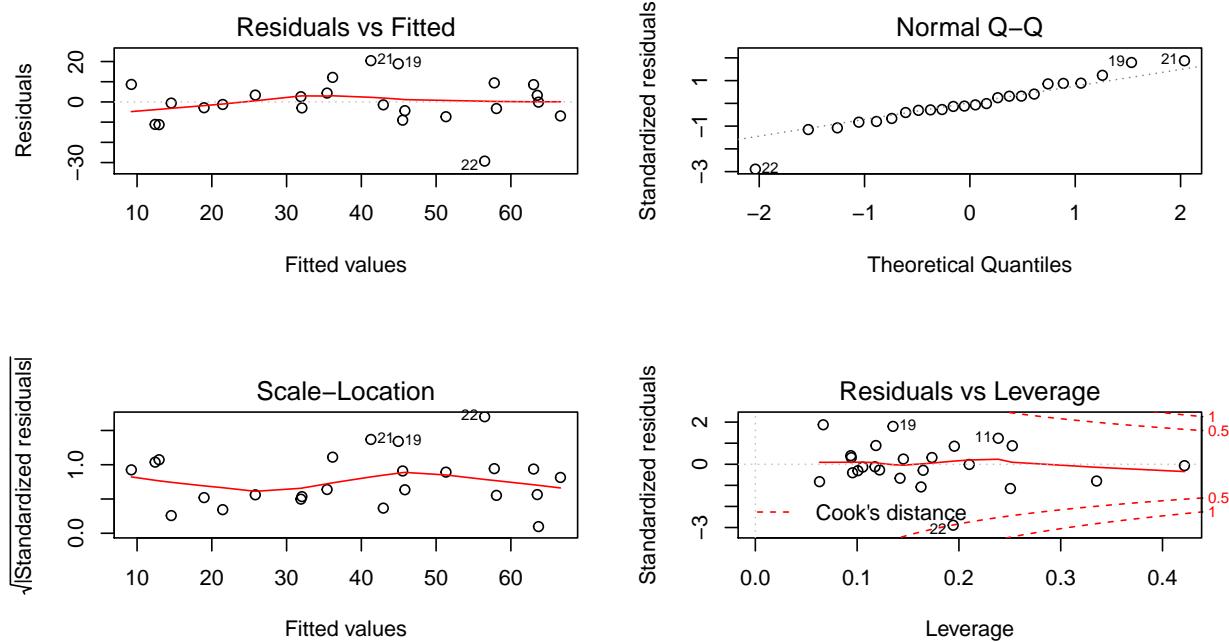


Figure 8.8: Diagnostic plots for MLR for Snow Depth based on Elevation, Min Temp and Max Temp with Northeast entrance observation removed from data set.

potentially influential and an outlier. Additionally, our model contains results that are not what we would have expected *a priori*, so it is not unreasonable to consider removing this observation to be able to work towards a model that fully trustworthy.

```
par(mfrow=c(2,2), oma=c(0,0,2,0))
plot(m5, sub.caption="Diagnostics for m5")
```

This worry-some observation is located in the 22nd row of the original data set:

```
snotel2[22,]
```

```
##   ID      Station Max.Temp Min.Temp Elevation Snow.Depth
## 22 36 Bloody [Redact.]     39       26      7550      27.2
```

With the removal of both the “Northeast Entrance” and “Bloody [REDACTED]” sites, there are  $n = 23$  observations remaining. This model (m6) seems to contain residual diagnostics (Figure 8.9) that are finally generally reasonable.

```
m6 <- lm(Snow.Depth ~ Elevation + Min.Temp + Max.Temp, data=snotel2[-c(9, 22),])
summary(m6)
```

```
##
## Call:
## lm(formula = Snow.Depth ~ Elevation + Min.Temp + Max.Temp, data = snotel2[-c(9,
## 22), ])
##
## Residuals:
##    Min      1Q  Median      3Q     Max
## -14.878 -4.486  0.024  3.996 20.728
##
```

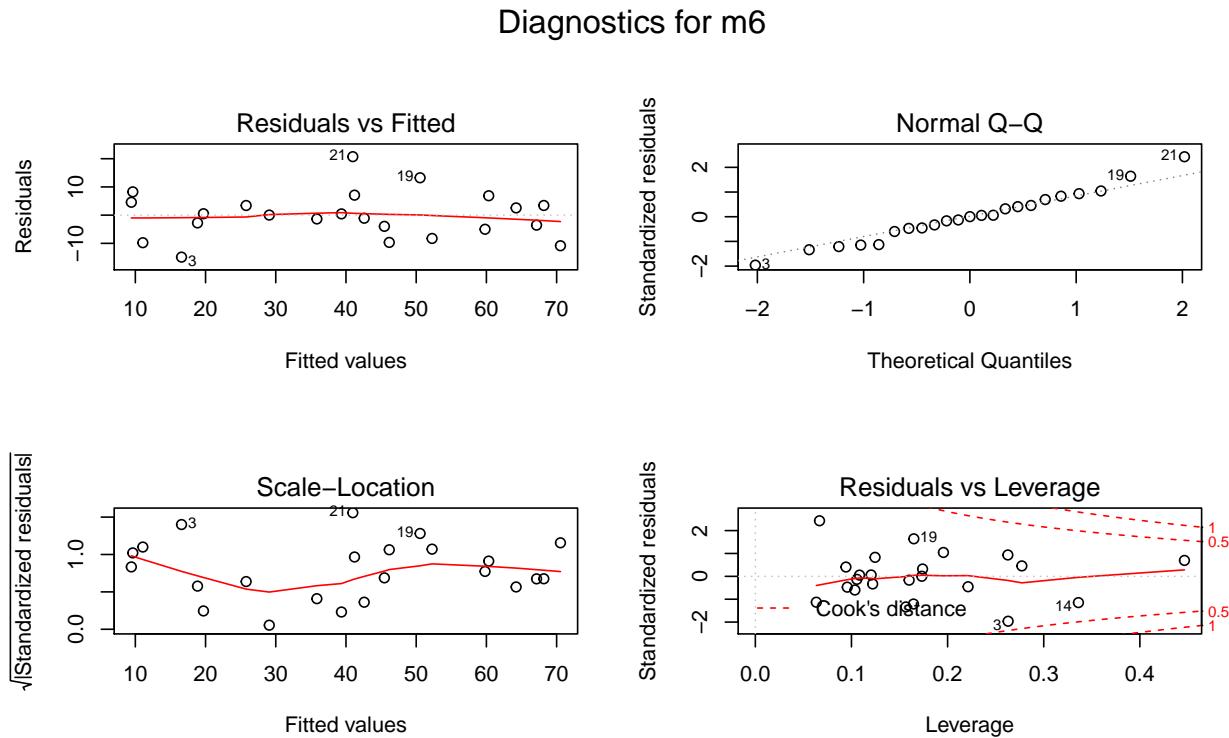


Figure 8.9: Diagnostic plots for MLR for Snow Depth based on Elevation, Min Temp and Max Temp with two observations removed ( $n = 23$ ).

```

## Coefficients:
##              Estimate Std. Error t value Pr(>|t|)
## (Intercept) -2.133e+02 7.458e+01 -2.859  0.0100
## Elevation    2.686e-02 4.997e-03  5.374 3.47e-05
## Min.Temp     9.843e-01 1.359e+00   0.724  0.4776
## Max.Temp    1.243e+00 5.452e-01   2.280  0.0343
##
## Residual standard error: 8.832 on 19 degrees of freedom
## Multiple R-squared:  0.8535, Adjusted R-squared:  0.8304
## F-statistic: 36.9 on 3 and 19 DF,  p-value: 4.003e-08
par(mfrow=c(2,2), oma=c(0,0,2,0))
plot(m6, sub.caption="Diagnostics for m6")

```

It is hard to suggest that there are any curvature issues and the slight variation in the Scale-Location plot is mostly due to few observations with fitted values around 30 happening to be well approximated by the model. The normality assumption is generally reasonable and no points seem to be overly influential on this model (finally!).

The term-plots (Figure 8.10) show that the temperature slopes are both positive although in this model *Max.Temp* seems to be more “important” than *Min.Temp*. We have ruled out individual influential points as the source of un-expected directions in slope coefficients and the more likely issue is multicollinearity – in a model that includes *Elevation*, the temperature effects may be positive, again acting with the *Elevation* term to generate the best possible predictions of the observed responses. Throughout this discussion, we have mainly focused on the slope coefficients and diagnostics. We have other tools in MLR to more quantitatively assess and compare different regression models that are considered in the next sections.

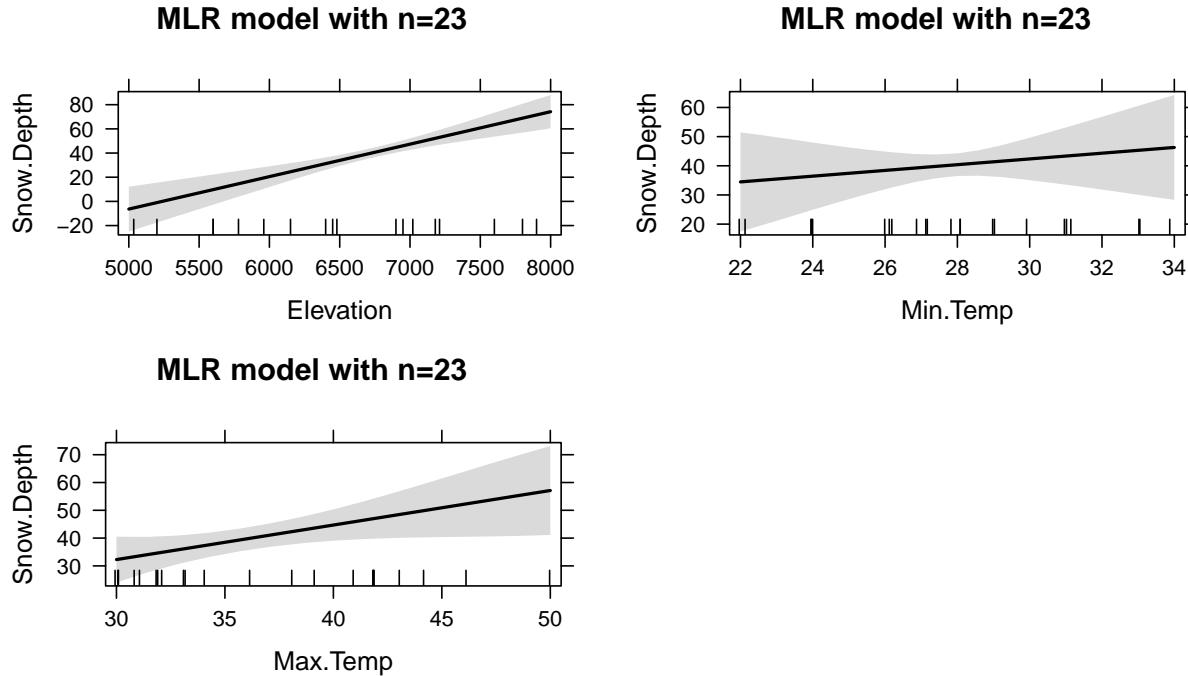


Figure 8.10: Term-plots for the MLR for Snow Depth based on Elevation, Min Temp and Max Temp with two observations removed.

```
plot(allEffects(m6), main="MLR model with n=23")
```

### 8.3 Interpretation of MLR terms

Since these results (finally) do not contain any highly influential points, we can formally discuss interpretations of the slope coefficients and how the term-plots (Figure 8.10) aid our interpretations. Term-plots in MLR are constructed by holding all the other variables at their mean and generating predictions and 95% CIs for the mean response across the levels of observed values for each predictor variable. This idea also helps us to work towards interpretations of each term in an MLR model. For example, for *Elevation*, the term-plot starts at an elevation around 5000 feet and ends at an elevation around 8000 feet. To generate that line and CIs for the mean snow depth at different elevations, the MLR model of

$$\widehat{\text{SnowDepth}}_i = -213.3 + 0.0269 \text{ Elevation}_i + 0.984 \text{ MinTemp}_i + 1.243 \text{ MaxTemp}_i$$

is used, but we need to have “something” to put in for the two temperature variables to predict *Snow Depth* for different *Elevations*. The typical convention is to hold the “other” variables at their means to generate these plots. This tactic also provides a way of interpreting each slope coefficient. Specifically, we can interpret the *Elevation* slope as: For a 1 foot increase in *Elevation*, we expect the mean *Snow Depth* to increase by 0.0269 inches, holding the minimum and maximum temperatures constant. More generally, the **slope interpretation in an MLR** is:

- For a 1 [*units of  $x_k$* ] increase in  $x_k$ , we expect the mean of  $y$  to change by  $b_k$  [*units of  $y$* ], after controlling for [list of other explanatory variables in model].

To make this more concrete, we can recreate some points in the Elevation term-plot. To do this, we first need the mean of the “other” predictors, *Min.Temp* and *Max.Temp*.

```
mean(snote12[-c(9,22),]$Min.Temp)
## [1] 27.82609
mean(snote12[-c(9,22),]$Max.Temp)
## [1] 36.3913
```

We can put these values into the MLR equation and simplify the equation, to an equation that is just in terms of just *Elevation* given that we are holding *Min.Temp* and *Max.Temp* at their means:

$$\begin{aligned}\widehat{\text{SnowDepth}}_i &= -213.3 + 0.0269 \text{ Elevation}_i + 0.984 * \mathbf{27.826} + 1.243 * \mathbf{36.391} \\ &= -213.3 + 0.0269 \text{ Elevation}_i + 27.38 + 45.23 \\ &= \mathbf{-140.69 + 0.0269 \text{ Elevation}_i}.\end{aligned}$$

So at the means on the two temperature variables, the model looks like an SLR with an estimated y-intercept of -140.69 (mean *Snow Depth* for *Elevation* of 0 if temperatures are at their means) and an estimated slope of 0.0269. Then we can plot the predicted changes in *y* across all the values of the predictor variable (*Elevation*) while holding the other variables constant. To generate the needed values to define a line, we can plug various *Elevation* values into the simplified equation:

- For an elevation of 5000 at the average temperatures, we predict a mean snow depth of  $-140.69 + 0.0269 * 5000 = -6.19$  inches.
- For an elevation of 6000 at the average temperatures, we predict a mean snow depth of  $-140.69 + 0.0269 * 6000 = 20.71$  inches.
- For an elevation of 8000 at the average temperatures, we predict a mean snow depth of  $-140.69 + 0.0269 * 8000 = 74.51$  inches.

We can plot this information (Figure 8.11) using the `plot` function to show the points we calculated and the `lines` function to add a line that connects the dots. In the `plot` function, we used the `ylim=...` option to make the scaling on the y-axis match the previous term-plot's scaling.

```
par(mfrow=c(1,1))
#Making own effect plot:
elevs <- c(5000,6000,8000)
snowdepths <- c(-6.19,20.71,74.51)
plot(snowdepths~elevs, ylim=c(-20,90), cex=2, col="blue", pch=16,
      main="Effect plot of elevation by hand")
lines(snowdepths~elevs, col="red", lwd=2)
```

Note that we only needed 2 points to define the line but need a denser grid of elevations if we want to add the 95% CIs for the true mean snow depth across the different elevations since they vary as a function of the distance from the mean of the explanatory variables.

To get the associated 95% CIs, we could return to using the `predict` function for the MLR, again holding the temperatures at their mean values. The `predict` function is sensitive and needs the same variable names as used in the original model fitting to work. First we create a “new” data set using the `seq` function to generate the desired grid of elevations and the `rep` function<sup>6</sup> to repeat the means of the temperatures for each of elevation values we need to make the plot. The code creates a specific version of the predictor variables to force the `predict` function to provide fitted values and CIs across different elevations with temperatures held constant that is stored in `newdata1`.

```
elevs <- seq(from=5000, to=8000, length.out=30)
newdata1 <- data.frame(Elevation=elevs, Min.Temp=rep(27.826,30),
                       Max.Temp=rep(36.3913,30))
newdata1
```

<sup>6</sup>The `seq` function has syntax of `seq(from=startingpoint, to=endingpoint, length.out=#ofvalues_between_start_and_end)` and the `rep` function has syntax of `rep(numbertorepeat, #oftimes)`.

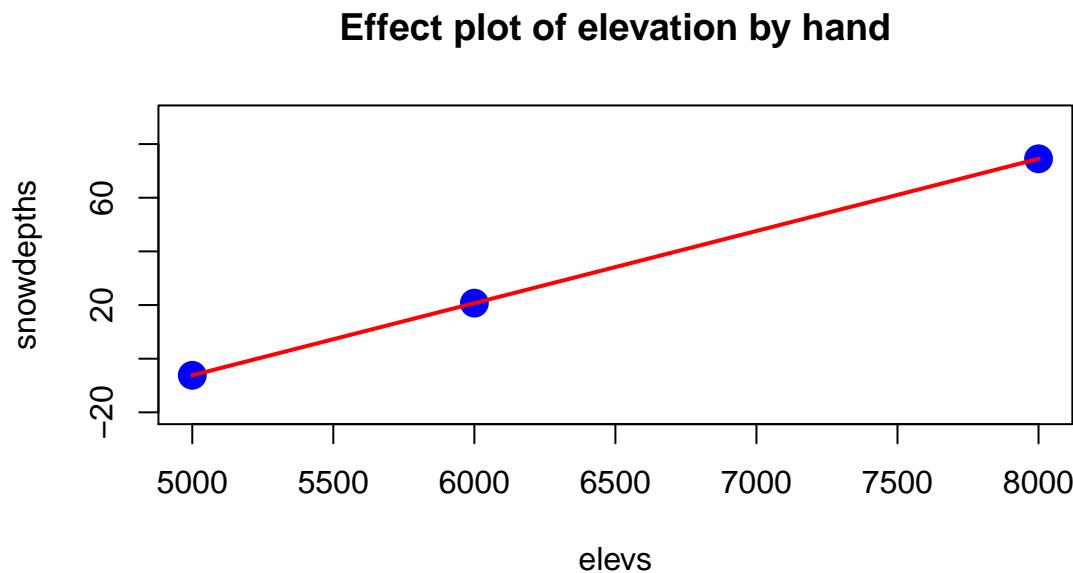


Figure 8.11: Term-plot for Elevation “by-hand”, holding temperature variables constant at their means.

```
##      Elevation Min.Temp Max.Temp
## 1    5000.000  27.826  36.3913
## 2    5103.448  27.826  36.3913
## 3    5206.897  27.826  36.3913
## 4    5310.345  27.826  36.3913
## 5    5413.793  27.826  36.3913
## 6    5517.241  27.826  36.3913
## 7    5620.690  27.826  36.3913
## 8    5724.138  27.826  36.3913
## 9    5827.586  27.826  36.3913
## 10   5931.034  27.826  36.3913
## 11   6034.483  27.826  36.3913
## 12   6137.931  27.826  36.3913
## 13   6241.379  27.826  36.3913
## 14   6344.828  27.826  36.3913
## 15   6448.276  27.826  36.3913
## 16   6551.724  27.826  36.3913
## 17   6655.172  27.826  36.3913
## 18   6758.621  27.826  36.3913
## 19   6862.069  27.826  36.3913
## 20   6965.517  27.826  36.3913
## 21   7068.966  27.826  36.3913
## 22   7172.414  27.826  36.3913
## 23   7275.862  27.826  36.3913
## 24   7379.310  27.826  36.3913
## 25   7482.759  27.826  36.3913
## 26   7586.207  27.826  36.3913
## 27   7689.655  27.826  36.3913
## 28   7793.103  27.826  36.3913
## 29   7896.552  27.826  36.3913
## 30   8000.000  27.826  36.3913
```

The predicted snow depths along with 95% confidence intervals for the mean, holding temperatures at their means, are:

```
predict(m6, newdata=newdata1, interval="confidence")

##          fit      lwr      upr
## 1 -6.3680312 -24.913607 12.17754
## 2 -3.5898846 -21.078518 13.89875
## 3 -0.8117379 -17.246692 15.62322
## 4  1.9664088 -13.418801 17.35162
## 5  4.7445555 -9.595708 19.08482
## 6  7.5227022 -5.778543 20.82395
## 7 10.3008489 -1.968814 22.57051
## 8 13.0789956  1.831433 24.32656
## 9 15.8571423  5.619359 26.09493
## 10 18.6352890  9.390924 27.87965
## 11 21.4134357 13.140233 29.68664
## 12 24.1915824 16.858439 31.52473
## 13 26.9697291 20.531902 33.40756
## 14 29.7478758 24.139153 35.35660
## 15 32.5260225 27.646326 37.40572
## 16 35.3041692 31.002236 39.60610
## 17 38.0823159 34.139812 42.02482
## 18 40.8604626 36.997617 44.72331
## 19 43.6386092 39.559231 47.71799
## 20 46.4167559 41.866745 50.96677
## 21 49.1949026 43.988619 54.40119
## 22 51.9730493 45.985587 57.96051
## 23 54.7511960 47.900244 61.60215
## 24 57.5293427 49.759987 65.29870
## 25 60.3074894 51.582137 69.03284
## 26 63.0856361 53.377796 72.79348
## 27 65.8637828 55.154251 76.57331
## 28 68.6419295 56.916422 80.36744
## 29 71.4200762 58.667725 84.17243
## 30 74.1982229 60.410585 87.98586
```

So we could do this with any model **for each predictor** variable to create term-plots, or we can just use the `allEffects` function to do this for us. This exercise is useful to complete once to understand what is being displayed in term-plots but using the `allEffects` function makes getting these plots much easier.

There are two other model components of possible interest in this model. The slope of 0.984 for *Min.Temp* suggests that for a  $1^{\circ}F$  increase in *Minimum Temperature*, we expect a 0.984 inch change in the mean *Snow Depth*, after controlling for *Elevation* and *Max.Temp* at the sites. Similarly, the slope of 1.243 for the *Max.Temp* suggests that for a  $1^{\circ}F$  increase in *Maximum Temperature*, we expect a 1.243 inch change in the mean *Snow Depth*, holding *Elevation* and *Min.Temp* constant. Note that there are a variety of ways to note that each term in an MLR is only a particular value given the other variables in the model. We can use words such as “holding the other variables constant” or “after adjusting for the other variables” or “controlling for the other variables”. The main point is to find words that reflect that this single slope coefficient might be different if we had a different overall model and the only way to interpret it is conditional on the other model components.

Term-plots have a few general uses to enhance our regular slope interpretations. They can help us see the model predicts over the range of each observed  $x$ . This can help you to get a sense of the “practical” importance of each term. Additionally, the term-plots show 95% confidence intervals for the mean response across the range of each variable, holding the other variables at their means. These intervals can be useful for assessing the precision in the estimated mean at different values of each predictor. However, note that you should not use these plots for deciding whether the term should be retained in the model – we have other tools for making that assessment. And one last note about term-plots – they do not mean that the relationships are really

linear between the predictor and response variable being displayed. The model **forces** the relationship to be linear even if that is not the real functional form. **Term-plots are not diagnostics for the model, they are summaries of the model you assumed was correct!** Any time we do linear regression, all of our inferences are contingent upon the model we chose. We know our model is not perfect, but we hope that it helps us learn something about our research question(s).

## 8.4 Comparing multiple regression models

With more than one variable, we now have many potential models that we could consider. We could include only one of the predictors, all of them, or combinations of sets of the variables. For example, maybe the model that includes *Elevation* does not “need” both *Min.Temp* and *Max.Temp*? Or maybe the model isn’t improved over an SLR with just *Elevation* as a predictor. Or maybe none of the predictors are “useful”? In this section, we discuss some general model comparison issues and a metric that can be used to pick among a suite of different models (often called a set of ***candidate models*** to reflect that they are all potentially interesting and we need to compare them and possibly pick one).

It is certainly possible the researchers may have an *a priori* reason to only consider a single model. For example, in a designed experiment where combinations of, say, three different predictors are randomly assigned, the initial model with all three predictors may be sufficient to address all the research questions of interest. One advantage in these situations is that the variable combinations can be created to prevent multicollinearity among the predictors and avoid that complication in interpretations. However, this is more the exception than the rule. Usually, there are competing predictors or questions about whether some predictors matter more than others. This type of research always introduces the potential for multicollinearity to complicate the interpretation of each predictor in the presence of others. Because of this, multiple models are often considered, where “unimportant” variables are dropped from the model. The assessment of “importance” using p-values will be discussed in Section 8.6, but for now we will consider other reasons to pick one model over another.

There are some general reasons to choose a particular model:

1. Diagnostics are better with one model compared to others.
2. One model predicts/explains the responses better than the others ( $R^2$ )).
3. *a priori* reasons to “use” a particular model, for example in a designed experiment or it includes variable(s) whose slopes need to be estimated to find interesting results (even if the variables are not “important” in the model).
4. Model selection “criteria” suggest one model is better than the others<sup>7</sup>.

It is OK to consider multiple reasons to select a model but it is dangerous to “shop” for a model across many possible models – a practice which is sometimes called ***data-dredging*** and leads to a high chance of spurious results from a single model that is usually reported based on this type of exploration. Just like in other discussions of multiple testing issues previously, if you explore many versions of a model, maybe only keeping the best ones, this is very different from picking one model (and tests) *a priori* and just exploring that result.

As in SLR, we can use the  $R^2$  (the ***coefficient of determination***) to measure the percentage of the variation in the response variable that the model explains. In MLR, it is important to remember that  $R^2$  is now an overall measure for the model and not specific to a single variable. It is comparable to other models including those fit with only a single predictor (SLR). So to meet criterion (2), we could simply find the model with the largest  $R^2$  value, finding the model that explains the most variation in the responses. Unfortunately for this idea, when you add more “stuff” to a regression model (even “unimportant” predictors), the  $R^2$  will always go up. This can be seen by considering

---

<sup>7</sup>Also see Section 8.13 for another method of picking among different models.

$$R^2 = \frac{SS_{\text{regression}}}{SS_{\text{total}}} \quad \text{where } SS_{\text{regression}} = SS_{\text{total}} - SS_{\text{error}} \quad \text{and } SS_{\text{error}} = \Sigma(y - \hat{y})^2.$$

Because adding extra variables to a linear model will only make the fitted values better, not worse, the  $SS_{\text{error}}$  will always go down if more predictors are added to the model. If  $SS_{\text{error}}$  goes down and  $SS_{\text{total}}$  is fixed, then adding extra variables will always increase  $SS_{\text{regression}}$  and, thus, increase  $R^2$ . This means that  $R^2$  is only useful for selecting models when you are picking between two models of the same size (same number of predictors). So we mainly use it as a summary of model quality once we pick a model, not a method of picking among a set of candidate models. Remember that  $R^2$  continues to have the property of being between 0 and 1 (or 0% and 100%) and that value refers to the **proportion (percentage) of variation in the response explained by the model**, whether we are using it for SLR or MLR.

However, there is an adjustment to the  $R^2$  measure that makes it useful for selecting among models. The measure is called the **adjusted  $R^2$** . The  $R^2_{\text{adjusted}}$  measure adds a penalty for adding more variables to the model, providing the potential for this measure to decrease if the extra variables do not really benefit the model. The measure is calculated as

$$R^2_{\text{adjusted}} = 1 - \frac{SS_{\text{error}}/df_{\text{error}}}{SS_{\text{total}}/(N-1)} = 1 - \frac{MS_{\text{error}}}{MS_{\text{total}}},$$

which incorporates the *degrees of freedom* for the model via the error *degrees of freedom* which go down as the model complexity increases. This adjustment means that just adding extra useless variables (variables that do not explain very much extra variation) do not increase this measure. That makes this measure useful for model selection since it can help us to stop adding unimportant variables and find a “good” model among a set of candidates. Like the regular  $R^2$ , larger values are better. The downside to  $R^2_{\text{adjusted}}$  is that it is **no longer a percentage of variation in the response that is explained by the model**; it can be less than 0 and so has no interpretable scale. It is just “larger is better”. It provides one method for building a model (different from using p-values to drop unimportant variables as discussed below), by fitting a set of candidate models containing different variables and then **picking the model with the largest  $R^2_{\text{adjusted}}$** . You will want to interpret this new measure on a percentage scale, but do not do that. It is a just a measure to help you pick a model and that is all it is!

One other caveat in model comparison is worth mentioning: make sure you are comparing models for the same responses. That may sound trivial and usually it is. But when there are missing values in the data set, especially on some explanatory variables and not others, it is important to be careful that the  $y$ 's do not change between models you are comparing. This relates to our *Snow Depth* modeling because responses were being removed due to their influential nature. We can't compare  $R^2$  or  $R^2_{\text{adjusted}}$  for  $n = 25$  to a model when  $n = 23$  – it isn't a fair comparison on either measure since they based on the total variability which is changing as the responses used change.

In the MLR (or SLR) model summaries, both the  $R^2$  and  $R^2_{\text{adjusted}}$  are available. Make sure you are able to pick out the correct one. For the reduced data set ( $n = 23$ ) *Snow Depth* models, the pertinent part of the model summary for the model with all three predictors is the last three lines:

```
m6 <- lm(Snow.Depth ~ Elevation + Min.Temp + Max.Temp, data=snote12[-c(9, 22),])
summary(m6)
```

```
##
## Call:
## lm(formula = Snow.Depth ~ Elevation + Min.Temp + Max.Temp, data = snote12[-c(9,
##      22), ])
##
## Residuals:
##     Min      1Q  Median      3Q     Max
## -14.878 -4.486  0.024  3.996 20.728
##
## Coefficients:
```

```

##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) -2.133e+02 7.458e+01 -2.859  0.0100
## Elevation    2.686e-02 4.997e-03  5.374 3.47e-05
## Min.Temp     9.843e-01 1.359e+00  0.724  0.4776
## Max.Temp    1.243e+00 5.452e-01  2.280  0.0343
##
## Residual standard error: 8.832 on 19 degrees of freedom
## Multiple R-squared:  0.8535, Adjusted R-squared:  0.8304
## F-statistic: 36.9 on 3 and 19 DF, p-value: 4.003e-08

```

There is a value for **Multiple R-Squared** of 0.8535, this is the  $R^2$  value and suggests that the model with *Elevation*, *Min* and *Max* temperatures explains 85.4% of the variation in *Snow Depth*. The  $R^2_{\text{adjusted}}$  is 0.8304 and is available further to the right labeled as **Adjusted R-Squared**. We repeated this for a suite of different models for this same  $n = 23$  data set and found the following results in Table 8.1. The top  $R^2_{\text{adjusted}}$  model is the model with *Elevation* and *Max.Temp*, which beats out the model with all three variables on  $R^2_{\text{adjusted}}$ . Note that the top  $R^2$  model is the model with three predictors, but the most complicated model will always have that characteristic.

Table 8.1: Model comparisons for Snow Depth data, sorted by model complexity.

Model	K	$R^2$	$R^2_{\text{adjusted}}$	$R^2_{\text{adjusted}}$	Rank
SD ~ Elevation	1	0.8087	0.7996		3
SD ~ Min.Temp	1	0.6283	0.6106		5
SD ~ Max.Temp	1	0.4131	0.3852		7
SD ~ Elevation + Min.Temp	2	0.8134	0.7948		4
SD ~ Elevation + Max.Temp	2	0.8495	0.8344		1
SD ~ Min.Temp + Max.Temp	2	0.6308	0.5939		6
SD ~ Elevation + Min.Temp + Max.Temp	3	0.8535	0.8304		2

The top model with *Elevation* and *Max.Temp* has an  $R^2$  of 0.8495, so we can say that the model with *Elevation* and *Maximum Temperature* explains 84.95% percent of the variation in *Snow Depth* and also that this model was selected based on the  $R^2_{\text{adjusted}}$ . One of the important features of  $R^2_{\text{adjusted}}$  available in this example – adding variables often does not always increase its value even though  $R^2$  does increase with **any** addition. In Section 8.13 we consider a competitor for this model selection criterion that may “work” a bit better and be extendable into more complicated modeling situations; that measure is called the **AIC**.

## 8.5 General recommendations for MLR interpretations and VIFs

There are some important issues to remember<sup>8</sup> when interpreting regression models that can result in common mistakes.

- **Don’t claim to “hold everything constant” for a single individual:**

Mathematically this is a correct interpretation of the MLR model but it is rarely the case that we could have this occur in real applications. Is it possible to increase the *Elevation* while holding the *Max.Temp* constant? We discussed making term-plots doing exactly this – holding the other variables

<sup>8</sup>This section was inspired by a similar section from De Veaux, Velleman, and Bock (2011).

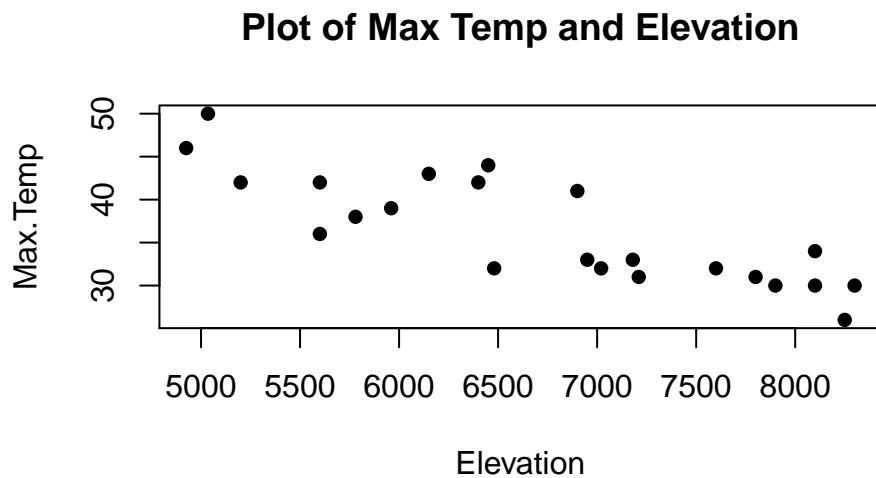


Figure 8.12: Scatterplot of observed Elevations and Maximum Temperatures for SNOTEL data.

constant at their means. If we interpret each slope coefficient in an MLR conditionally then we can craft interpretations such as: For locations that have a *Max.Temp* of, say, 45°F and *Min.Temp* of, say, 30°F, a 1 foot increase in *Elevation* tends to be associated with a 0.0268 inch increase in *Snow Depth* on average. This does not try to imply that we can actually make that sort of change but that given those other variables, the change for that variable is a certain magnitude.

- **Don't interpret the regression results causally (or casually?)...**

Unless you are analyzing the results of a designed experiment (where the levels of the explanatory variable(s) were randomly assigned) you cannot state that a change in that *x* **causes** a change in *y*, especially for a given individual. The multicollinearity in predictors makes it especially difficult to put too much emphasis on a single slope coefficient because it may be corrupted/modified by the other variables being in the model. In observational studies, there are also all the potential lurking variables that we did not measure or even confounding variables that we did measure but can't disentangle from the variable used in a particular model. While we do have a complicated mathematical model relating various *x*'s to the response, do not lose that fundamental focus on causal vs non-causal inferences based on the design of the study.

- **Be cautious about doing prediction in MLR – you might be doing extrapolation!**

It is harder to know if you are doing extrapolation in MLR since you could be in a region of the *x*'s that no observations were obtained. Suppose we want to predict the *Snow Depth* for an *Elevation* of 6000 and *Max.Temp* of 30. Is this extrapolation based on Figure 8.12? In other words, can you find any observations “nearby” in the plot of the two variables together? What about an *Elevation* of 6000 and a *Max.Temp* of 40? The first prediction is in a different proximity to observations than the second one... In situations with more than two explanatory variables it becomes even more challenging to know whether you are doing extrapolation and the problem grows as the number of dimensions to search increases... In fact, in complicated MLR models we typically do not know whether there are observations “nearby” if we are doing predictions for unobserved combinations of our predictors. Note that Figure 8.12 also reinforces our potential collinearity problem between *Elevation* and *Max.Temp* with higher elevations being strongly associated with lower temperatures.

- **Don't think that the sign of a coefficient is special...**

Adding other variables into the MLR models can cause a switch in the coefficients or change their

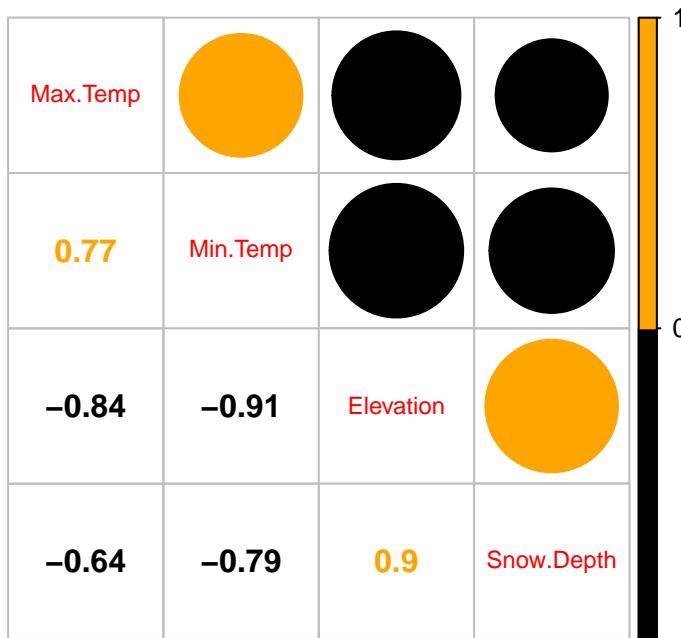


Figure 8.13: Plot of correlation matrix in the snow depth data set with influential points removed

magnitude or make them go from “important” to “unimportant” without changing the slope too much. This is related to the conditionality of the relationships being estimated in MLR and the potential for sharing of information in the predictors when it is present.

- **Multicollinearity in MLR models:**

When explanatory variables are not independent (related) to one another, then including one variable will have an impact on the other variable. Consider the correlations among the predictors in the SNOTEL data set or visually displayed in Figure 8.13:

```
##           Max.Temp Min.Temp Elevation Snow.Depth
## Max.Temp     1.00    0.77    -0.84     -0.64
## Min.Temp     0.77    1.00    -0.91     -0.79
## Elevation   -0.84   -0.91     1.00      0.90
## Snow.Depth   -0.64   -0.79     0.90      1.00
require(corrplot)
par(mfrow=c(1,1), oma=c(0,0,1,0))
corrplot.mixed(cor(snotel2[-c(9,22),3:6]), col=c("black", "orange"))
round(cor(snotel2[-c(9,22),3:6]),2)
```

The predictors all share at least moderately strong linear relationships. For example, the  $r = -0.91$  between *Min.Temp* and *Elevation* suggests that they contain very similar information and that extends to other pairs of variables as well. When variables share information, their addition to models may not improve the performance of the model and actually can make the estimated coefficients *unstable*, creating uncertainty in the correct coefficients because of the shared information. It seems that *Elevation* is related to *Snow Depth* but maybe it is because it has lower *Minimum Temperatures*? So you might wonder how we can find the “correct” slopes when they are sharing information in the response variable. The short answer is that we can’t. But we do use **Least Squares** to find coefficient estimates as we did before – except that we have to remember that these **estimates are conditional on other variables in the model** for our interpretation since they impact one another within the model. It ends up that the uncertainty of pinning those variables down in the presence of shared information leads to larger SEs for all the slopes. And that we can actually measure **how much each of the**

**SEs are inflated** because of multicollinearity with other variables in the model using what are called **Variance Inflation Factors** (or **VIFs**).

**VIFs** provide a way to assess the multicollinearity in the MLR model that is caused by including specific variables. The amount of information that is shared between a single explanatory variable and the others can be found by regressing that variable on the others and calculating  $R^2$  for that model. The code for this regression is something like: `lm(X1~X2+X3+?+XK)`, which regresses  $X_1$  on  $X_2$  through  $X_K$ . The  $1 - R^2$  from this regression is the amount of independent information in  $X_1$  that is not explained by the other variables in the model. The VIF for each variable is defined using this quantity as  $\text{VIF}_k = 1/(1 - R_k^2)$  for variable  $k$ . If there is no shared information ( $R^2 = 0$ ), then the VIF will be 1. But if the information is completely shared with other variables ( $R^2 = 1$ ), then the VIF goes to infinity ( $1/0$ ). Basically, large VIFs are bad, with the rule of thumb that values over 5 or 10 are considered “large” values indicating high multicollinearity in the model for that particular variable. We use this scale to determine if multicollinearity is a problem for a variable of interest. Additionally, the  $\sqrt{\text{VIF}_k}$  is also very interesting as it is the number of times larger than the SE for the slope for variable  $k$  is due to collinearity with other variables in the model. This is the most useful scale to understand VIFs even though the rules of thumb are on the original scale. An example will show how to easily get these results and where the results come from.

In general, the easy way to obtain VIFs is using the `vif` function from the `car` package (Fox, 2003). It has the advantage of also providing a reasonable result when we include categorical variables in models (Sections 8.9 and 8.11). We apply the `vif` function directly to a model of interest and it generates values for each explanatory variable.

```
require(car)
vif(m6)
```

```
## Elevation Min.Temp Max.Temp
## 8.164201 5.995301 3.350914
```

Not surprisingly, there is an indication of problems with multicollinearity in two of the three variables in the model with the largest issues identified for *Elevation* and *Min.Temp*. Both of their VIFs exceed 5 indicating large multicollinearity problems. On the square-root scale, the VIFs show more interpretation utility.

```
sqrt(vif(m6))
```

```
## Elevation Min.Temp Max.Temp
## 2.857307 2.448530 1.830550
```

The result for *Elevation* of 2.86 suggests that the SE for *Elevation* is 2.86 times larger than it should be because of multicollinearity with other variables in the model. Similarly, the *Min.Temp* SE is 2.45 times larger and the *Max.Temp* SE is 1.83 times larger. All of this generally suggests issues with multicollinearity in the model and that we need to be cautious in interpreting any slope coefficients from this model.

In order to see how the VIF is calculated for *Elevation*, we need to regress *Elevation* on *Min.Temp* and *Max.Temp*. Note that this model is only fit to find the percentage of variation in elevation explained by the temperature variables. It ends up being 0.8775 – so a high percentage of *Elevation* can be explained by the linear model using min and max temperatures.

```
#VIF calc:
elev1<-lm(Elevation~Min.Temp+Max.Temp,data=snotel2[-c(9,22),])
summary(elev1)

##
## Call:
## lm(formula = Elevation ~ Min.Temp + Max.Temp, data = snotel2[-c(9,
## 22), ])
##
## Residuals:
##      Min        1Q    Median        3Q       Max
## -1120.05 -142.99    14.45   186.73   624.61
##
```

```
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) 14593.21    699.77 20.854 4.85e-15
## Min.Temp     -208.82     38.94 -5.363 3.00e-05
## Max.Temp     -56.28     20.90 -2.693   0.014
##
## Residual standard error: 395.2 on 20 degrees of freedom
## Multiple R-squared:  0.8775, Adjusted R-squared:  0.8653
## F-statistic: 71.64 on 2 and 20 DF,  p-value: 7.601e-10
```

Using this result, we can calculate

$$\text{VIF}_{\text{elevation}} = \frac{1}{1 - R_{\text{elevation}}^2} = \frac{1}{1 - 0.8775} = \frac{1}{0.1225} = 8.16$$

$\text{1} - 0.8775$

```
## [1] 0.1225
1/0.1225
```

```
## [1] 8.163265
```

Note that when we observe small VIFs, that provides us with confidence that multicollinearity is not causing problems under the surface of a particular MLR model. And that we can't use the VIFs to do anything about multicollinearity in the models – it is just a diagnostic to understand the magnitude of the problem.

## 8.6 MLR Inference: Parameter inferences using the t-distribution

We have been deliberately vague about what an important variable is up to this point, and chose to focus on some bigger modeling issues. We now turn our attention to one of the most common tasks in any basic statistical model – assessing whether a particular result is more unusual than we would expect by chance. All the previous discussions of estimation in MLR models informs our interpretations of the tests. The  $t$ -tests for slope coefficients are based on our standard recipe – take the estimate, divide it by its standard error and then, assuming the statistic follows a  $t$ -distribution under the null hypothesis, find a p-value. This tests whether each true slope coefficient,  $\beta_k$ , is 0 or not, in a model that contains all the other variables. Again, sometimes we say “after adjusting for” the other  $x$ 's or “conditional on” the other  $x$ 's in the model or “after allowing for”... as in the slope coefficient interpretations above. The main point is that **you should not interpret anything related to slope coefficients in MLR without referencing the other variables that are in the model!** The tests for the slope coefficients assess  $H_0 : \beta_k = 0$ , which in words is a test that there is no linear relationship between explanatory variable  $k$  and the response variable,  $y$ , in the population, given the other variables in model. The typical alternative hypothesis is  $H_0 : \beta_k \neq 0$ . In words, the alternative hypothesis is that there is some linear relationship between explanatory variable  $k$  and the response variable,  $y$ , in population, given the other variables in the model. It is also possible to test for positive or negative slopes in the alternative, but this is rarely the first concern, especially when MLR slopes can occasionally come out in unexpected directions.

The test statistic for these hypotheses is  $t = \frac{b_k}{\text{SE}_k}$  and, if our assumptions are met, follows a  $t$ -distribution with  $n - K - 1$  df where  $K$  is the number of predictor variables in the model. We perform the test for each slope coefficient, but the test is conditional on all the other variables in the model – the order the variables are fit in does **not** change  $t$ -test results. For the *Snow Depth* example with **Elevation** and **Maximum Temperature** as predictors, the pertinent output is in the four columns of the **Coefficient table** that is the first part of the model summary we've been working with. You can find the estimated slope (**Estimate** column), the SE of the slopes (**Std. Error** column), the  $t$ -statistics (**t value** column), and the p-values (**Pr(>|t|)** column). The degrees of freedom for the  $t$ -distributions show up below the coefficients and the  $df = 20$  here. This is because  $n = 23$  and  $K = 2$ , so  $df = 23 - 2 - 1 = 20$ .

```
m5 <- lm(Snow.Depth~Elevation+Max.Temp, data=snotel2[-c(9,22),])
summary(m5)

##
## Call:
## lm(formula = Snow.Depth ~ Elevation + Max.Temp, data = snotel2[-c(9,
##   22), ])
##
## Residuals:
##    Min      1Q  Median      3Q     Max
## -14.652  -4.645   0.518   3.744  20.550
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) -1.675e+02  3.924e+01 -4.269 0.000375
## Elevation    2.407e-02  3.162e-03  7.613 2.48e-07
## Max.Temp     1.253e+00  5.385e-01  2.327 0.030556
##
## Residual standard error: 8.726 on 20 degrees of freedom
## Multiple R-squared:  0.8495, Adjusted R-squared:  0.8344
## F-statistic: 56.43 on 2 and 20 DF,  p-value: 5.979e-09
```

The hypotheses for the *Maximum Temperature* term (*Max.Temp*) are:

- $H_0 : \beta_{\text{Max.Temp}} = 0$  given that *Elevation* is in the model vs
- $H_A : \beta_{\text{Max.Temp}} \neq 0$  given that *Elevation* is in the model.

The test statistic is  $t = 2.327$  with  $df = 20$  (so under the null hypothesis the test statistic follows a  $t_{20}$ -distribution).

The output provides a p-value of 0.0306 for this test. We can also find this using `pt`:

```
2*pt(2.327, df=20, lower.tail=F)
```

```
## [1] 0.03058319
```

The decision here would probably be to reject  $H_0$ . The chance of observing a slope for *Max.Temp* as extreme or more extreme than assuming there really is no linear relationship between *Max.Temp* and *Snow Depth* (in a model with *Elevation*), is about 3%.

Conclusion: There is sufficient evidence to suggest that there is a linear relationship between *Max.Temp* and *Snow Depth*, once we account for *Elevation*, in the population of SNOTEL sites in Montana on this day. Because we cannot randomly assign the temperatures to sites, we cannot conclude that temperature causes changes in the snow depth – in fact it might even be possible for a location to have different temperatures because of different snow depths.

Similarly, we can test for *Elevation* after controlling for the *Maximum Temperature*:

$$H_0 : \beta_{\text{Elevation}} = 0 \text{ vs } H_A : \beta_{\text{Elevation}} \neq 0,$$

given that *Max.Temp* is in the model.

$t = 7.613$  ( $df = 20$ ) with a p-value of 0.00000025 or just  $< 0.00001$ .

Decision: Reject  $H_0$  and conclude that there is sufficient evidence to suggest that there is a linear relationship between *Elevation* and *Snow Depth*, once we adjust for *Max.Temp* in the population of SNOTEL sites in Montana on this day.

There is one last test that is of dubious interest in almost every situation – to test that the y-intercept ( $\beta_0$ ) in an MLR is 0. This tests if the true mean response is 0 when all the predictor variables are set to 0. I

see researchers reporting this p-value frequently and it is possibly the most useless piece of information in the regression model summary. Sometimes less educated statistics users even think this result is proof of something interesting or are disappointed when the p-value is not small. Unless you want to do some prediction and are interested in whether the mean response when all the predictors are set to 0 is different from 0, this test should not be reported or, if reported, is certainly not very interesting<sup>9</sup>. But we should at least go through the motions on this test once so you don't make the same mistakes.

$H_0 : \beta_0 = 0$  vs  $H_A : \beta_0 \neq 0$  in a model with *Elevation* and *Maximum Temperature*

$t = -4.269$ , with an assumption that the test statistic follows a  $t_{20}$ -distribution under the null hypothesis, and the p-value = 0.000375.

Decision: Reject  $H_0$

Conclusion: There is sufficient evidence to suggest that the true mean *Snow Depth* is different from 0 when the *Maximum Temperature* is 0 and the *Elevation* is 0 in the population of SNOTEL sites. To reinforce the general uselessness of this test, think about the combination of  $x$ 's – is that even physically possible in Montana (or the continental US) in April?

Remember when testing slope coefficients in MLR, that if we fail to reject the null hypothesis, it does not mean that there is no relationship or even no linear relationship between the variables, but that there is no evidence of a linear relationship **once we account for the other variables in the model**. If you do not find a small p-value for a variable, you should either be cautious when interpreting the coefficient, or not interpret it. Some model building strategies would lead to dropping the term from the model but sometimes we will have models to interpret that contain terms with larger p-values. Sometimes they are still of interest but the weight on the interpretation isn't as heavy as if the term had a small p-value – you should remember that you can't prove that coefficient is different from 0 in that model. It also may mean that you don't know too much about its specific value. Confidence intervals will help us pin down where we think the true slope coefficient might be located, given the other variables in the model.

Confidence intervals provide the dual uses of inferences for the location of the true slope and whether the true slope seems to be different from 0. The confidence intervals here have our regular format of estimate  $\mp$  margin of error. Like the previous tests, we work with  $t$ -distributions with  $n - K - 1$  degrees of freedom. Specifically the 95% confidence interval for slope coefficient  $k$  is

$$b_k \mp t_{n-K-1}^* \text{SE}_{b_k}.$$

The interpretation is the same as in SLR with the additional tag of “after controlling for the other variables in the model” for all the reasons discussed before. The general slope CI interpretation for predictor  $x_k$  in an MLR is:

- For a 1 [unit of  $x_k$ ] increase in  $x_k$ , we are 95% confident that the true mean of  $y$  changes by between **LL** and **UL** [units of Y] in the population, after adjusting for the other  $x$ 's [**list them!**].

We can either calculate these intervals as we have many times before or we can rely on the `confint` function to do this:

```
confint(m5)
```

```
##           2.5 %      97.5 %
## (Intercept) -249.37903311 -85.67576239
## Elevation     0.01747878   0.03067123
## Max.Temp     0.13001718   2.37644112
```

So for a  $1^{\circ}\text{F}$  increase in *Maximum Temperature*, we are 95% confident that the true mean *Snow Depth* will change by between 0.13 and 2.38 inches in the population, after adjusting for the *Elevation* of the sites. Similarly, for a 1 foot increase in *Elevation*, we are 95% confident that the true mean *Snow Depth* will change

---

<sup>9</sup>There are some social science models where the model is fit with the mean subtracted from each predictor so all have mean 0 and the precision of the y-intercept is interesting. But even in these models, the test the y-intercept being 0 is rarely of interest.

by between 0.0175 and 0.0307 inches in the population, after adjusting for the *Maximum Temperature* of the sites.

## 8.7 Overall F-test in Multiple Linear Regression

In the MLR summary, reported at the bottom of the output. For the model with *Elevation* and *Maximum Temperature*, the last row of the model summary is:

```
## F-statistic: 56.43 on 2 and 20 DF, p-value: 5.979e-09
```

This test is called the ***overall F-test*** in MLR and is very similar to the *F*-test in a reference-coded One-Way ANOVA model. It tests the null hypothesis that involves setting every coefficient except the y-intercept to 0 (so all the slope coefficients equal 0). We saw this reduced model in the One-Way material when we considered setting all the deviations from the baseline group to 0 under the null hypothesis. We can frame this as a comparison between a full and reduced model as follows:

- **Full Model:**  $y_i = \beta_0 + \beta_1 x_{1i} + \beta_2 x_{2i} + \cdots + \beta_K x_{Ki} + \varepsilon_i$
- **Reduced Model:**  $y_i = \beta_0 + 0x_{1i} + 0x_{2i} + \cdots + 0x_{Ki} + \varepsilon_i$

The reduced model estimates the same values for all  $y$ 's,  $\hat{y}_i = \bar{y} = b_0$  and corresponds to the null hypothesis of:

$H_0$  : No explanatory variables should be included in the model:  $\beta_1 = \beta_2 = \cdots = \beta_K = 0$ .

The full model corresponds to the alternative:

$H_A$  : At least one explanatory variable should be included in the model: Not all  $\beta_k$ 's = 0 for  $(k = 1, \dots, K)$ .

Note that  $\beta_0$  is not set to 0 in the reduced model (under the null hypothesis) – it becomes the true mean of  $y$  for all values of the  $x$ 's since all the predictors are multiplied by coefficients of 0.

The test statistic to assess these hypotheses is  $F = \text{MS}_{\text{model}}/\text{MS}_E$ , which is assumed to follow an *F*-distribution with  $K$  numerator *df* and  $n - K - 1$  denominator *df* under the null hypothesis. The output provides us with  $F(2, 20) = 56.43$  and a p-value of  $5.979 * 10^{-9}$  (p-value < 0.00001) and enough evidence to reject the null hypothesis. Thus, there is evidence that at least one of the two slope coefficients (*Max.Temp*'s or *Elevation*'s) is different from 0 in the population of SNOTEL sites in Montana on this date. While this test is a little bit interesting and a good indicator of something interesting in the model, the moment you see this result, you want to know more about each predictor variable. If neither predictor variable is important, we will discover that in the *t*-tests for each coefficient and so our general recommendation is to start there.

The overall F-test, then, is really about testing whether there is something good in the model somewhere. And that certainly is important but it is also not too informative. There is one situation where this test is really interesting, when there is only one predictor variable in the model (SLR). In that situation, this test provides exactly the same p-value as the *t*-test. *F*-tests will be important when we are mixing categorical and quantitative predictor variables in our MLR models (Section 8.12), but the overall *F*-test is of very limited utility.

## 8.8 Case Study: First year college GPA and SATs

Many universities require students to have certain test scores in order to be admitted into their institutions. They obviously must think that those scores are useful predictors of student success to use them in this way. Quality assessments of recruiting classes are also based on their test scores. The Educational Testing Service (the company behind such fun exams as the SAT and GRE) collected a data set to validate their SAT on  $n = 1000$  students from an unnamed Midwestern university; the data set is available in the *openintro*

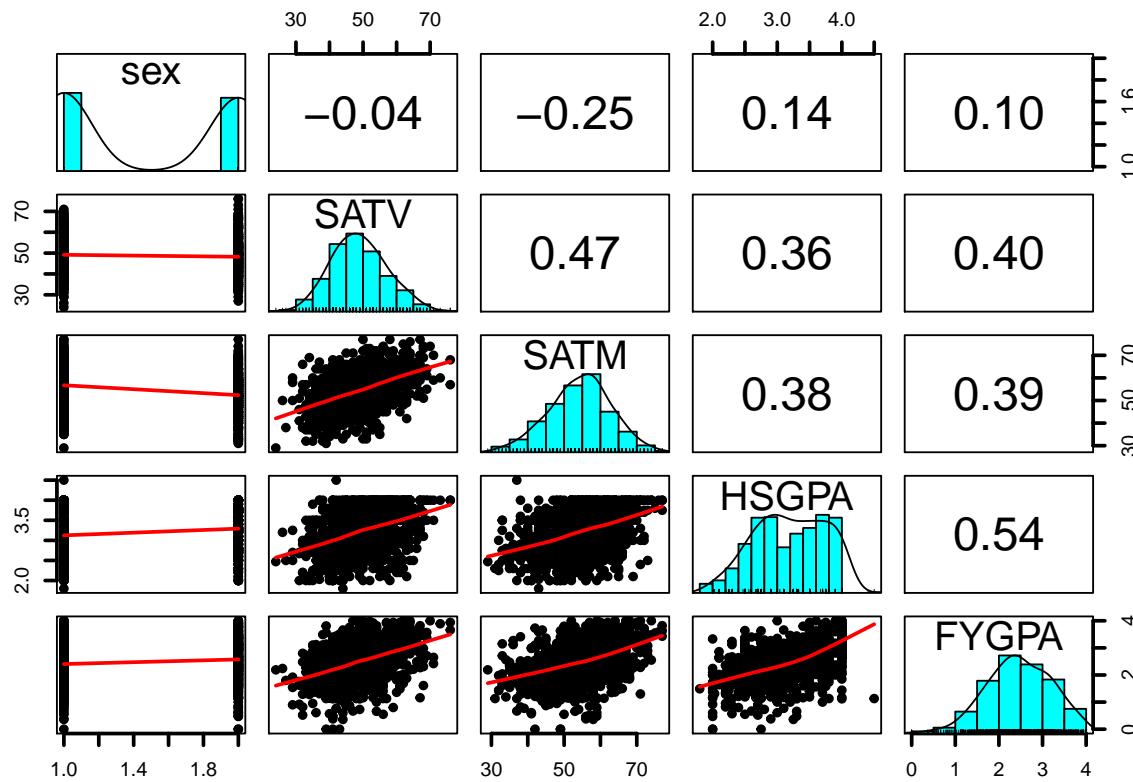


Figure 8.14: Scatterplot matrix of SAT and GPA data set.

package (Diez et al., 2012) in the `satGPA` data set. It is unclear from the documentation whether a random sample was collected, in fact it looks like it certainly wasn't a random sample of all incoming students at a large university (more later). What potential issues would arise if a company was providing a data set to show the performance of their test and it was not based on a random sample?

We will proceed assuming they used good methods in developing their test (there are sophisticated statistical models underlying the development of the SAT and GRE) and in obtaining a data set for testing out the performance of their tests that is at least representative of the students (or some types of students) at this university. They provided information on the *Sex* (`sex`) of the students (coded 1 and 2 with possibly 1 for males and 2 for females – but should this even be displayed in a plot with correlations?), *SAT Verbal* (`SATV`) and *Math* (`SATM`) percentiles (these are not the scores but the ranking percentile that each score translated to in a particular year), *High School GPA* (`HSGPA`), and *First Year* (of college) *GPA* `FYGPA`. Our interests here are in whether the two SAT percentiles are (together?) related to the first year college GPA, describing the size of their impacts and assessing the predictive potential of SAT-based measures for first year in college GPA. There are certainly other possible research questions that can be addressed with these data but this will keep us focused.

```
require(openintro)
data(satGPA)
require(psych)
pairs.panels(satGPA[,-4], ellipse=F, col="red", lwd=2)
```

There are positive relationships in Figure 8.14 among all the pre-college measures and the *college GPA* but none are above the moderate strength level. The *HSGPA* has a highest correlation with first year of college

results but its correlation is not that strong. Maybe together in a model the SAT percentiles can also be useful... Also note that plot shows an odd *HSGPA* of 4.5 that probably should be removed<sup>10</sup> if that variable is going to be used (*HSGPA* was not used in the following models so the observation remains in the data).

In MLR, the modeling process is a bit more complex and often involves more than one model, so we will often avoid the 6+ steps in testing initially and try to generate a model we can use in that more specific process. In this case, the first model of interest using the two SAT percentiles,

$$\text{FYGPA}_i = \beta_0 + \beta_{\text{SATV}} \text{SATV}_i + \beta_{\text{SATM}} \text{SATM}_i + \varepsilon_i,$$

looks like it might be worth interrogating further so we can jump straight into considering the 6+ steps involved in hypothesis testing for the two slope coefficients. We will be using a 5% significance level and *t*-based inferences, assuming that we can trust the assumptions.

Note that this is not a randomized experiment but we can assume that it is representative of the students at that single university. We would not want to extend these inferences to other universities (who might be more or less selective) or to students who did not get into this university and, especially, not to students that failed to complete the first year. The second and third constraints point to a severe limitation in this research – only students who were accepted, went to, and survived one year at this university could be studied. Lower SAT percentile students might not have been allowed in or may not have survived the first year and higher SAT students might have been attracted to other more prestigious institutions. So the scope of inference is just limited to students that were invited and chose to attend this institution and successfully completed one year of courses. It is hard to know if the SAT “works” when the inferences are so restricted in who they might apply to...

The following code fits the model of interest, provides a model summary, and the diagnostic plots, allowing us to consider the tests of interest:

```
gpa1 <- lm(FYGPA~SATV+SATM, data=satGPA)
summary(gpa1)
```

```
##
## Call:
## lm(formula = FYGPA ~ SATV + SATM, data = satGPA)
##
## Residuals:
##      Min       1Q   Median       3Q      Max
## -2.19647 -0.44777  0.02895  0.45717  1.60940
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) 0.007372  0.152292  0.048   0.961
## SATV        0.025390  0.002859  8.879  < 2e-16
## SATM        0.022395  0.002786  8.037 2.58e-15
##
## Residual standard error: 0.6582 on 997 degrees of freedom
## Multiple R-squared:  0.2122, Adjusted R-squared:  0.2106
## F-statistic: 134.2 on 2 and 997 DF,  p-value: < 2.2e-16
par(mfrow=c(2,2), oma=c(0,0,2,0))
plot(gpa1, sub.caption="Diagnostics for GPA model with SATV and SATM")
```

### 1. Hypotheses of interest:

- $H_0 : \beta_{\text{SATV}} = 0$  given  $\text{SATM}$  in the model vs  $H_0 : \beta_{\text{SATV}} \neq 0$  given  $\text{SATM}$  in the model.
- $H_0 : \beta_{\text{SATM}} = 0$  given  $\text{SATV}$  in the model vs  $H_0 : \beta_{\text{SATM}} \neq 0$  given  $\text{SATV}$  in the model.

<sup>10</sup>Either someone had a weighted GPA with bonus points, or more likely here, there was a coding error in the data set since only one observations was over 4.0 in the GPA data. Either way, we could remove it and note that our inferences for HSGPA do not extend above 4.0.

### Diagnostics for GPA model with SATV and SATM

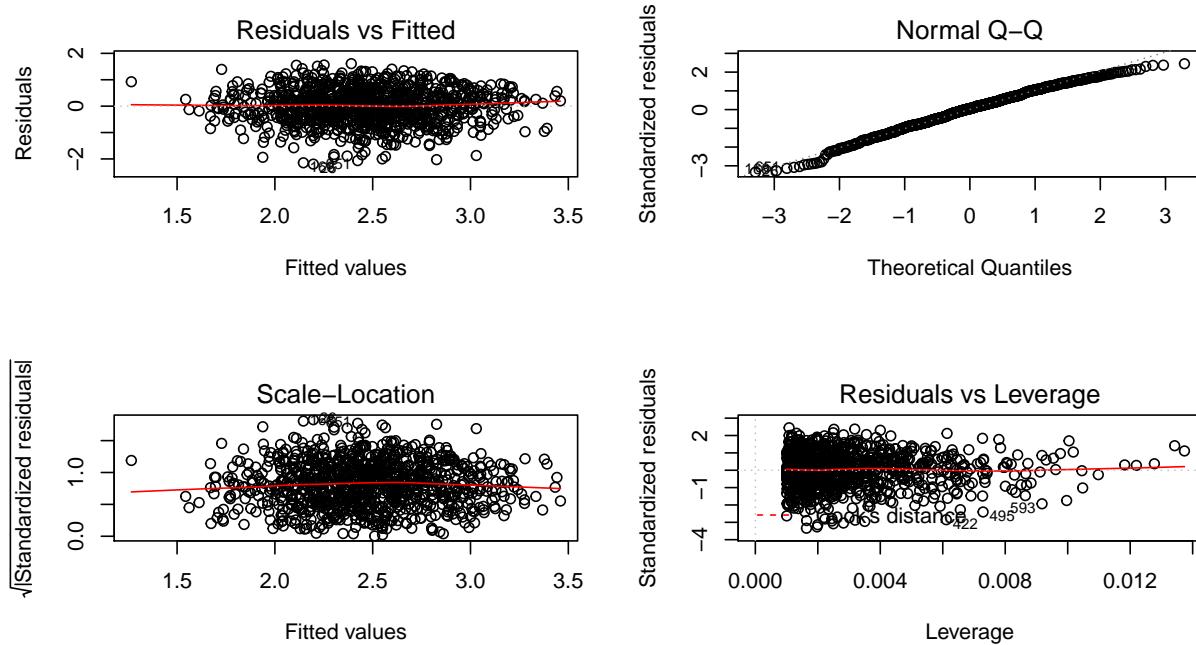


Figure 8.15: Diagnostic plots for the FYGPA ~ SATV + SATM model.

#### 2. Validity conditions:

- **Quantitative variables condition:**

- The variables used here are all quantitative. Note that the *sex* was plotted in the previous scatterplot matrix and is not quantitative – we will explore its use later.

- **Independence of observations:**

- With a sample from a single university from (we are assuming) a single year of students, there is no particular reason to assume a violation of the independence assumption.

- **Linearity of relationships:**

- The initial scatterplots (Figure 8.14) do not show any clear nonlinearities with each predictor used in this model.
- The Residuals vs Fitted and Scale-Location plots (Figure 8.15) do not show much more than a football shape, which is our desired result.

- Together, we can feel relatively comfortable with the linearity assumption.

- **Multicollinearity checked for:**

- The original scatterplots suggest that there is some collinearity between the two SAT percentiles with a correlation of 0.47. That is actually a bit lower than one might expect and suggests that each score must be measuring some independent information about different characteristics of the students.
- VIFs also do not suggest a major issue with multicollinearity in the model with the VIFs

for both variables the same at 1.278<sup>11</sup>. This suggests that both SEs are about 13% larger than they otherwise would have been due to shared information between the two predictor variables.

```
vif(gpa1)

##      SATV      SATM
## 1.278278 1.278278
sqrt(vif(gpa1))

##      SATV      SATM
## 1.13061 1.13061
```

- **Equal (constant) variance:**

- There is no clear change in variability as a function of fitted values.

- **Normality of residuals:**

- There is a minor deviation in the upper tail of the residual distribution from normality. It is not pushing towards having larger values than a normal distribution would generate so should not cause us any real problems. Note that this upper limit is likely due to using GPA as a response variable and it has an upper limit. This is an example of a potentially *censored* variable. For a continuous variable it is possible that the range of a measurement scale doesn't distinguish among subjects who differ once they pass a certain point. For example, a 4.0 high school student is going to have a high first year college GPA, on average but there is no room for variability in college GPA up, just down. For students more in the middle of the range, they can vary up or down. So in some places you can get symmetric distributions around the mean and in others you cannot. There are specific statistical models for these types of responses that are beyond our scope. In this situation, failing to account for the censoring may push some slopes toward 0 a little because we don't have responses over 4.0 to work with.

- **No influential points:**

- There are no influential points. In large data sets, the influence of any point is decreased and even high leverage and outlying points can struggle to have any impacts at all on the results.

So we are fairly comfortable with all the assumptions being at least not clearly violated and so the inferences from our model should be relatively trustworthy.

3. Calculate the test statistics:

- For *SATV*:  $t = \frac{0.02539}{0.002859} = 8.88$  with  $df = 997$ .
- For *SATM*:  $t = \frac{0.02240}{0.002786} = 8.04$  with  $df = 997$ .

4. Find the p-values:

- For *SATV*: p-value < 0.0001
- For *SATM*: p-value < 0.0001

5. Decisions:

- For *SATV*: Reject  $H_0$  because there is almost no chance of observing a test statistic as extreme or more extreme than was observed if there really were no linear relationship between *FY GPA* and *SATV*, in a model that controls for *SATM*.

---

<sup>11</sup>When there are just two predictors, the VIFs have to be the same since the proportion of information shared is the same in both directions. With more than two predictors, each variable can have a different VIF value.

- For  $SATM$ : Reject  $H_0$  because there is almost no chance of observing a test statistic as extreme or more extreme than was observed if there really were no linear relationship between  $FY GPA$  and  $SATM$ , in a model that controls for  $SATV$ .

#### 6. Conclusions and Scope of Inference:

- For  $SATV$ : There is strong evidence to reject the null hypothesis of no linear relationship between  $SATV$  and  $FY GPA$  and conclude that, in fact, there is a linear relationship between  $SATV$  percentile and the first year of college  $GPA$ , after controlling for the  $SATM$  percentile, in the population of students that completed their first year at this university.
- For  $SATM$ : There is strong evidence to reject the null hypothesis of no linear relationship between  $SATM$  and  $FY GPA$  and conclude that, in fact, there is a linear relationship between  $SATM$  percentile and the first year of college  $GPA$ , after controlling for the  $SATV$  percentile, in the population of students that completed their first year at this university.
- Note that neither inference is causal because there was no random assignment of SAT percentiles to the subjects. The inferences are also limited to students who stayed in school long enough to get a  $GPA$  from their first year of college.

We could stop there, but just reporting the test results without quantifying the size of the effects is not fully satisfying. The estimated MLR model is

$$\widehat{FY GPA}_i = 0.00737 + 0.0254SATV_i + 0.0224SATM_i.$$

So for a 1 percent increase in the  $SATV$  percentile, we expect, on average, to get a 0.0254 point change in  $GPA$ , after controlling for  $SATM$  percentile. Similarly, for a 1 percent increase in the  $SATM$  percentile, we expect, on average, to get a 0.0224 point change in  $GPA$ , after controlling for  $SATV$  percentile. While this is a correct interpretation of the slope coefficients, it is often easier to assess “practical” importance of the results by considering how much change this implies over the range of observed predictor values.

The term-plots (Figure 8.16) provide a visualization of the “size” of the differences in the response variable explained by each predictor. The  $SATV$  term-plot shows that for the range of percentiles from around the 30th percentile to the 70th percentile, the mean first year  $GPA$  is predicted to go from approximately 1.9 to 3.0. That is a pretty wide range of differences in GPAs across the range of observed percentiles. This looks like a pretty interesting and important change in the mean first year GPA across that range of different SAT percentiles. Similarly, the  $SATM$  term-plot shows that the  $SATM$  percentiles were observed to range between around the 30th percentile and 70th percentile and predict mean GPAs between 1.95 and 2.8. It seems that the SAT Verbal percentiles produce slightly more impacts in the model, holding the other variable constant, but that both are important variables. The 95% confidence intervals for the means in both plots suggest that the results are fairly precisely estimated – there is little variability around the predicted means in each plot. This is mostly a function of the sample size as opposed to the model itself explain most of the variation in the responses.

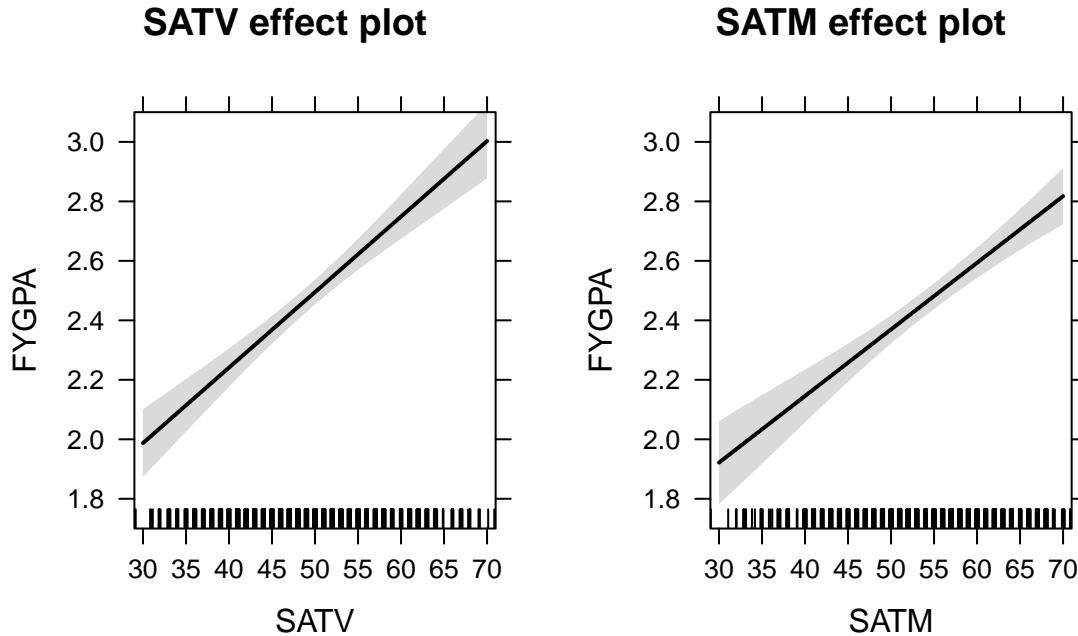
```
require(effects)
plot(allEffects(gpa1))
```

These plots also inform the types of students attending this university and successfully completing the first year of school. This seems like a good, but maybe not great, institution with few students scoring over the 75th percentile on either SAT Verbal or Math (at least that ended up in this data set). This result makes questions about their sampling mechanism re-occur as to who this data set might actually be representative of...

The confidence intervals also help us pin down the uncertainty in each estimated slope coefficient. As always, the “easy” way to get 95% confidence intervals is using the `confint` function:

```
confint(gpa1)
```

```
##              2.5 %      97.5 %
## (Intercept) -0.29147825  0.30622148
```

Figure 8.16: Term-plots for the  $FYGPA \sim SATV + SATM$  model.

```
## SATV      0.01977864 0.03100106
## SATM      0.01692690 0.02786220
```

So, for a 1 percent increase in the  $SATV$  percentile, we are 95% confident that the true mean  $FYGPA$  changes between 0.0198 and 0.031 points, in the population of students who completed this year at this institution, after controlling for  $SATM$ . The  $SATM$  result is similar with an interval from 0.0169 and 0.0279. Both of these intervals might benefit from re-scaling the interpretation to a 10 percentile increase in the predictor variable, with the change in the  $FYGPA$  for that level of increase of  $SATV$  providing an interval from 0.198 to 0.31 points and for  $SATM$  providing an interval from 0.169 to 0.279. So a boost of 10% in either exam percentile likely results in a noticeable but not huge average  $FYGPA$  increase.

One final use of these methods is to do prediction and generate prediction intervals, which could be quite informative for a student considering going to this university who has a particular set of SAT scores. For example, suppose that the student is interested in the average  $FYGPA$  to expect with  $SATV$  at the 30th percentile and  $SATM$  at the 60th percentile. The predicted mean value is

$$\begin{aligned}\hat{\mu}_{GPA_i} &= 0.00737 + 0.0254 \text{ SATV}_i + 0.0224 \text{ SATM}_i \\ &= 0.00737 + 0.0254 * 30 + 0.0224 * 60 = 2.113\end{aligned}$$

This result and the 95% confidence interval for the mean student  $GPA$  at these scores can be found using the `predict` function as:

```
predict(gpa1,newdata=data.frame(SATV=30,SATM=60))
```

```
##      1
## 2.11274
predict(gpa1,newdata=data.frame(SATV=30,SATM=60),interval="confidence")

##      fit      lwr      upr
## 1 2.11274 1.982612 2.242868
```

For students at the 30th percentile of  $SATV$  and 60th percentile of  $SATM$ , we are 95% confident that the

true mean first year GPA is between 1.98 and 2.24 points. For an individual student, we would want the 95% prediction interval:

```
predict(gpa1,newdata=data.frame(SATV=30,SATM=60),interval="prediction")
```

```
##      fit      lwr      upr
## 1 2.11274 0.8145859 3.410894
```

For a student with  $SATV=30$  and  $SATM=60$ , we are 95% sure that their first year GPA will be between 0.81 and 3.4 points. You can see that while we are very certain about the mean in this situation, there is a lot of uncertainty in the predictions for individual students. The PI is so wide as to almost not be useful.

To support this difficulty in getting a precise prediction for a new student, review the original scatterplots: there is quite a bit of vertical variability in first year *GPA*s for each level of any of the predictors. The residual SE,  $\hat{\sigma}$ , is also informative in this regard – remember that it is the standard deviation of the residuals around the regression line. It is 0.6582, so the SD of new observations around the line is 0.66 GPA points and that is pretty large on a GPA scale if we assume we would see lots of observations within 2 or 3 SDs of the mean. Figure 8.17 remakes both term-plots, holding the other predictor at its mean, and adds the 95% prediction intervals to show the difference in variability between estimating the mean and pinning down the value of a new observation. The R code is very messy and rarely needed, but hopefully this helps reinforce the differences in these two types of intervals – to make them in MLR, you have to fix all but one of the predictor variables and we usually do that by fixing the other variables at their means.

```
#Remake effects plots with 95% PIs
dv1 <- data.frame(SATV=seq(from=24,to=76,length.out=50), SATM=rep(54.4,50))
dm1 <- data.frame(SATV=rep(48.93,50), SATM=seq(from=29,to=77,length.out=50))

mv1 <- data.frame(predict(gpa1, newdata=dv1, interval="confidence"))
pv1 <- data.frame(predict(gpa1, newdata=dv1, interval="prediction"))

mm1 <- data.frame(predict(gpa1, newdata=dm1, interval="confidence"))
pm1 <- data.frame(predict(gpa1, newdata=dm1, interval="prediction"))

par(mfrow=c(1,2))

plot(dv1$SATV, mv1$fit, lwd=2, ylim=c(pv1$lwr[1],pv1$upr[50]), type="l",
      xlab="SATV Percentile", ylab="GPA", main="SATV Effect, CI and PI")
lines(dv1$SATV ,mv1$lwr, col="red", lty=2, lwd=2)
lines(dv1$SATV ,mv1$upr, col="red", lty=2, lwd=2)
lines(dv1$SATV ,pv1$lwr, col="grey", lty=3, lwd=3)
lines(dv1$SATV ,pv1$upr, col="grey", lty=3, lwd=3)
legend("topleft", c("Estimate", "CI", "PI"), lwd=3, lty=c(1,2,3),
       col = c("black", "red", "grey"))

plot(dm1$SATM, mm1$fit, lwd=2, ylim=c(pm1$lwr[1], pm1$upr[50]), type="l",
      xlab="SATM Percentile", ylab="GPA", main="SATM Effect, CI and PI")
lines(dm1$SATM ,mm1$lwr, col="red", lty=2, lwd=2)
lines(dm1$SATM ,mm1$upr, col="red", lty=2, lwd=2)
lines(dm1$SATM ,pm1$lwr, col="grey", lty=3, lwd=3)
lines(dm1$SATM ,pm1$upr, col="grey", lty=3, lwd=3)
```

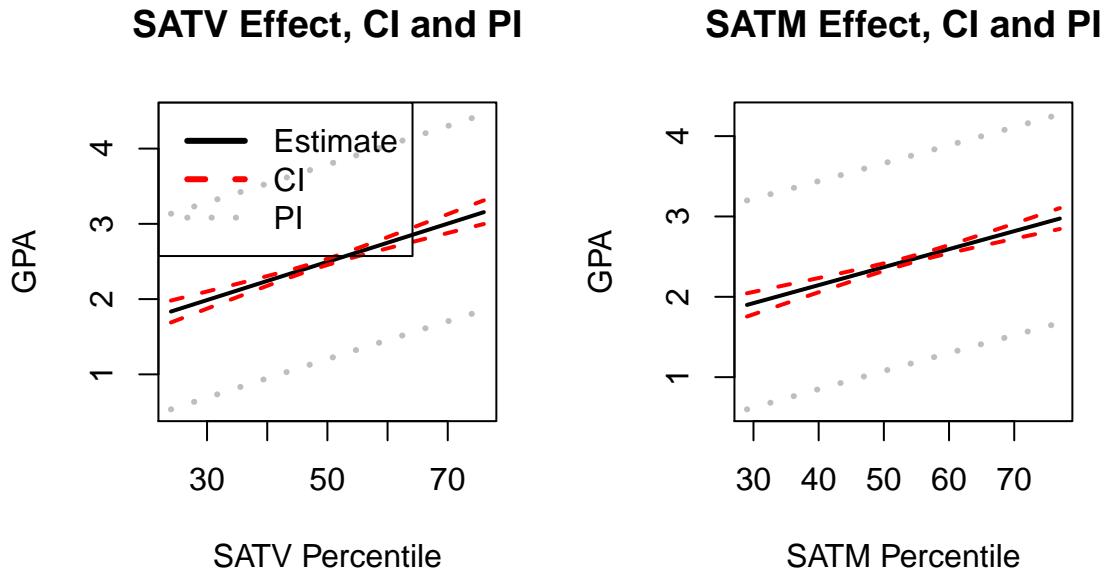


Figure 8.17: Term-plots for the  $FYGPA \sim SATV + SATM$  model with 95% PIs.

## 8.9 Different intercepts for different groups: MLR with Indicator variables

One of the implicit assumptions up to this point was that the models were being applied to a single homogeneous population. In many cases, we take a sample from a population but that group is likely a combination of individuals from different sub-populations. For example, the SAT study was interested in all students at the university but that contains the obvious sub-populations based on the sex of the students. It is dangerous to fit MLR models across subpopulations but we can also use MLR models to address more sophisticated research questions by comparing groups. We will be able to compare the intercepts (mean levels) and the slopes to see if they differ between the groups. For example, does the relationship between the *SATV* and *FY GPA* differ for male and female students? We can add the grouping information to the scatterplot of *FY GPA* vs *SATV* (Figure 8.18) and consider whether there is visual evidence of a difference in the slope and/or intercept between the two groups, with men coded as 1 and women coded as 2<sup>12</sup>.

It appears that the slope for females might be larger (steeper) in this relationship than it is for males. So increases in SAT Verbal percentiles for females might have more of an impact on the average first year GPA. We'll handle this sort of situation in Section 8.11, where we will formally consider how to change the slopes for different groups. In this section, we develop new methods needed to begin to handle these situations and explore creating models with the same slope coefficient for all groups but different y-intercepts. This material resembles what we did for the Two-Way ANOVA additive model.

These results contrast with Figure 8.19 for the relationship between first year college *GPA* and *SATM* percentile by sex of the students. The lines for the two groups appear to be mostly parallel and just seem to have different y-intercepts. We can use our MLR techniques to fit a model to the entire data set that allows for different y-intercepts. The real power of this idea is that we can then also test whether the different groups have different y-intercepts – whether the shift between the groups is “real”. In this example, it appears

<sup>12</sup>We are actually making an educated guess about what these codes mean. Other similar data sets used 1 for males but the documentation on these data is a bit sparse. We can proceed with a small potential that all conclusions regarding differences in **sex** are in the wrong direction.

to suggest that females generally have slightly higher GPAs than males, on average, but that an increase in SATM has the same impact for both groups. If this difference in y-intercepts is not “real”, then there appears to be no difference between the sexes in their relationship between SATM and GPA and we can safely continue using a model that does not differentiate the two groups. We could also just subset the data set and do two analyses, but that approach will not allow us to assess whether things are “really” different between the two groups.

```
scatterplot(FYGPAs~SATM|sex, lwd=3, data=satGPA, spread=F,
           smooth=F, main="Scatterplot of GPA vs SATM by Sex")
scatterplot(FYGPAs~SATM|sex, lwd=3, data=satGPA, spread=F,
           smooth=F, main="Scatterplot of GPA vs SATM by Sex")
```

To fit one model to a data set that contain multiple groups, we need a way of entering categorical variable information in an MLR model. Regression models require quantitative predictor variables for the  $x$ 's so we can't directly enter the sex of the students into the regression model since it contains categories. To be able to put in “numbers” as predictors, we create what are called *indicator variables*<sup>13</sup> that are made up of 0s and 1s, with the 0 reflecting one category and 1 the other, changing depending on the category of the individual in the data set. The `lm` function does this for whenever a categorical variable is used as an explanatory variable. It sets up the indicator variables using a baseline category (gets coded as a 0) and the deviation category for the other level of the variable. We can see how this works by exploring what happens when we put `SEX` into our `lm`<sup>14</sup> with SATM, after first making sure it is categorical using the `factor` function and making the group levels explicit instead of 1s and 2s.

```
satGPA$SEX <- factor(satGPA$sex)
levels(satGPA$SEX) <- c("MALE", "FEMALE")
SATSex1 <- lm(FYGPAs~SATM+SEX, data=satGPA)
summary(SATSex1)

##
## Call:
## lm(formula = FYGPAs ~ SATM + SEX, data = satGPA)
##
## Residuals:
##      Min        1Q    Median        3Q       Max
## -2.42124 -0.42363  0.01868  0.46540  1.66397
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) 0.21589   0.14858   1.453   0.147
## SATM        0.03861   0.00258  14.969 < 2e-16
## SEXFEMALE   0.31322   0.04360   7.184 1.32e-12
##
## Residual standard error: 0.6667 on 997 degrees of freedom
## Multiple R-squared:  0.1917, Adjusted R-squared:  0.1901
## F-statistic: 118.2 on 2 and 997 DF, p-value: < 2.2e-16
```

<sup>13</sup>Some people also call them *dummy variables*.

<sup>14</sup>That may not read how we intended...

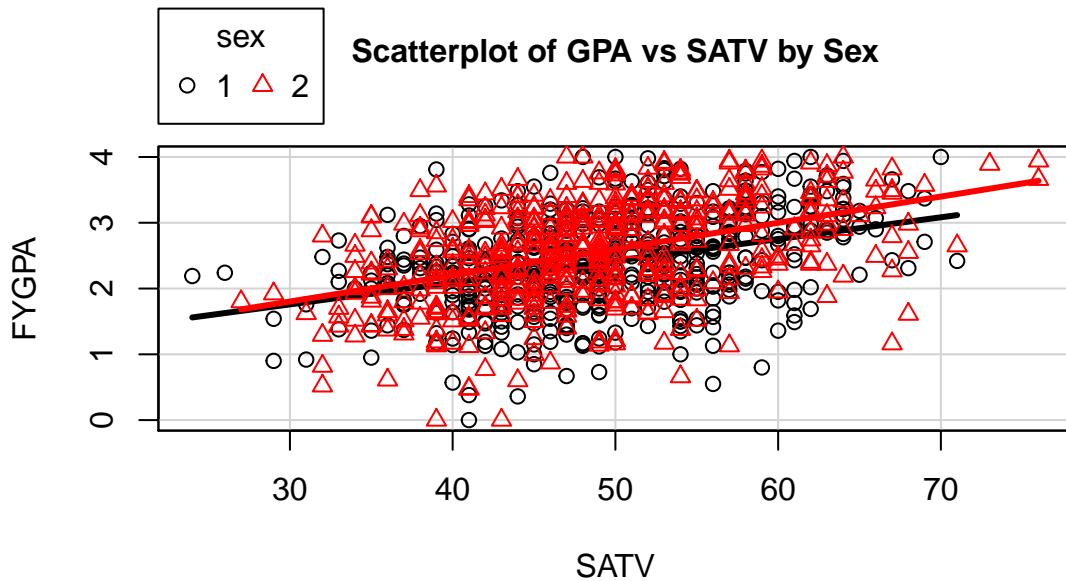


Figure 8.18: Plot of FYGPA vs SATV by Sex of students.

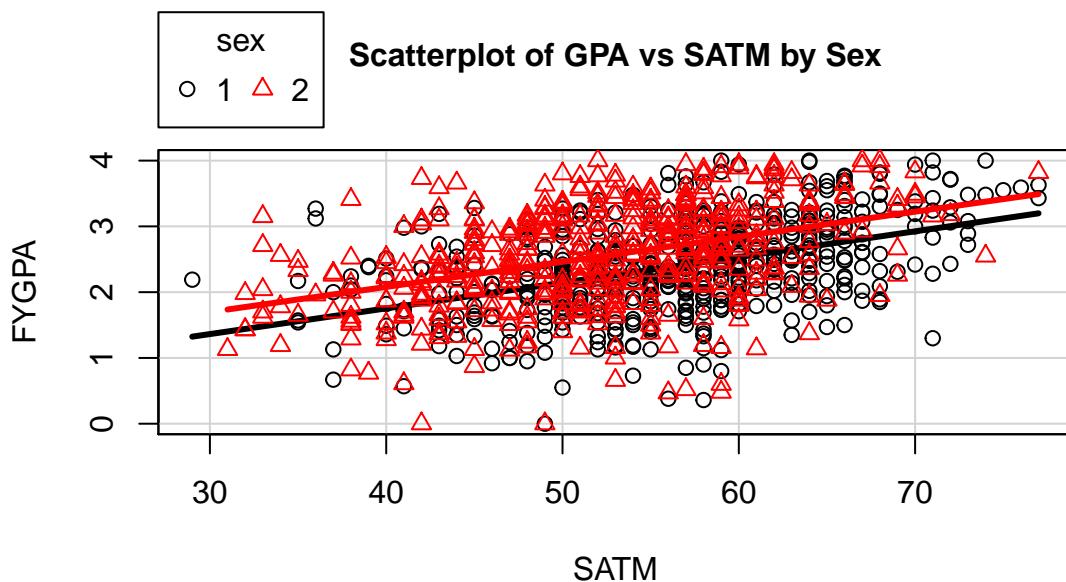


Figure 8.19: Plot of FYGPA vs SATM by Sex of students.

The `SEX` row contains information that the linear model chose `MALE` as the baseline category and `FEMALE` as the deviation category since `MALE` does not show up in the output. To see what `lm` is doing for us when we give it a two-level categorical variable, we can create our own “numerical” predictor that is 0 for *males* and 1 for *females* that we called `SEXINDICATOR`, displayed for the first 10 observations:

```
satGPA$SEXINDICATOR <- as.numeric(satGPA$SEX=="FEMALE")
head(data.frame(SEX=satGPA$SEX, SEXINDICATOR=satGPA$SEXINDICATOR), 10)
```

```
##      SEX SEXINDICATOR
## 1    MALE          0
## 2 FEMALE          1
## 3 FEMALE          1
## 4    MALE          0
## 5    MALE          0
## 6 FEMALE          1
## 7    MALE          0
## 8    MALE          0
## 9 FEMALE          1
## 10   MALE          0
```

We can define the indicator variable more generally by calling it  $I_{\text{Female},i}$  to denote that it is an indicator ( $I$ ) that takes on a value of 1 for observations in the category *Female* and 0 otherwise (*Male*) – changing based on the observation ( $i$ ). Indicator variables, once created, are quantitative variables that take on values of 0 or 1 and we can put them directly into linear models with other  $x$ 's (quantitative or categorical). If we replace the categorical `SEX` variable with our quantitative `SEXINDICATOR` and re-fit the model, we get:

```
SATSex2 <- lm(FYGPA ~ SATM + SEXINDICATOR, data=satGPA)
summary(SATSex2)
```

```
##
## Call:
## lm(formula = FYGPA ~ SATM + SEXINDICATOR, data = satGPA)
##
## Residuals:
##      Min       1Q   Median       3Q      Max 
## -2.42124 -0.42363  0.01868  0.46540  1.66397 
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) 0.21589   0.14858   1.453   0.147    
## SATM        0.03861   0.00258  14.969   < 2e-16 ***
## SEXINDICATOR 0.31322   0.04360   7.184  1.32e-12 ***
## 
## Residual standard error: 0.6667 on 997 degrees of freedom
## Multiple R-squared:  0.1917, Adjusted R-squared:  0.1901 
## F-statistic: 118.2 on 2 and 997 DF,  p-value: < 2.2e-16
```

This matches all the previous `lm` output except that we didn't get any information on the categories used since `lm` didn't know that `SEXINDICATOR` was anything different from other quantitative predictors.

Now we want to think about what this model means. We can write the estimated model as

$$\widehat{\text{FYGPA}}_i = 0.216 + 0.0386 \text{SATM}_i + 0.313I_{\text{Female},i}.$$

When we have a *male* observation, the indicator takes on a value of 0 so the 0.313 drops out of the model, leaving an SLR just in terms of `SATM`. For a *female* student, the indicator is 1 and we add 0.313 to the previous y-intercept. The following works this out step-by-step, simplifying the MLR into two SLRs:

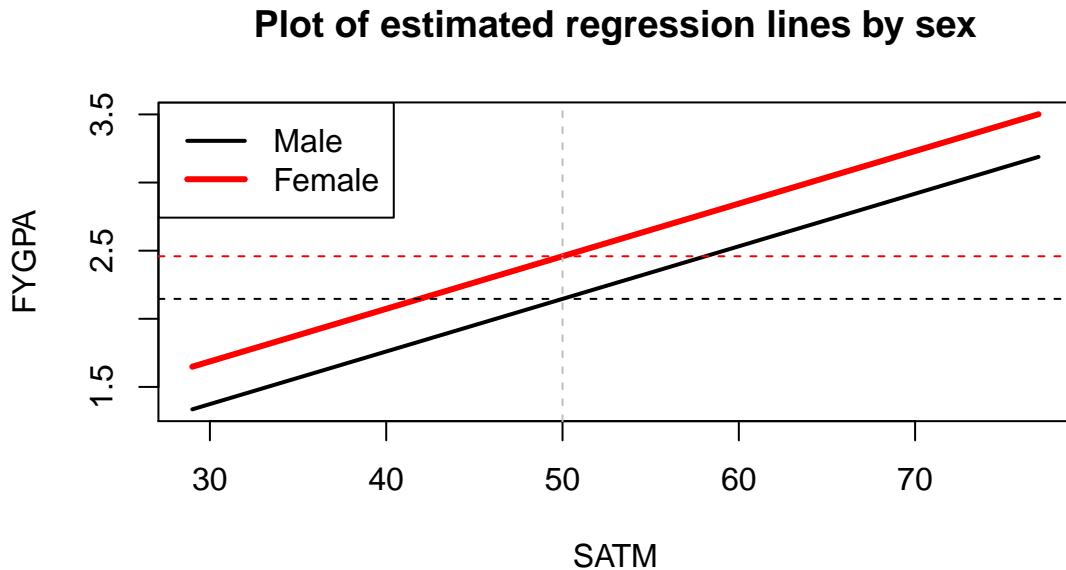


Figure 8.20: Plot of estimated model for  $FYGPA$  vs  $SATM$  by  $SEX$  of students (female line is red, bold).

- Simplified model for *Males* (plug in a 0 for  $I_{Female}$ ):

$$-\widehat{FYGPA}_i = 0.216 + 0.0386 \text{SATM}_i + 0.313 * 0 = 0.216 + 0.0386 \text{SATM}_i$$

- Simplified model for *Females* (plug in a 1 for  $I_{Female}$ ):

$$\begin{aligned} -\widehat{FYGPA}_i &= 0.216 + 0.0386 \text{SATM}_i + 0.313 * 1 \\ &= 0.216 + 0.0386 \text{SATM}_i + 0.313 \text{ (combine "like" terms to simplify the equation)} \\ &= 0.529 + 0.0386 \text{SATM}_i \end{aligned}$$

In this situation, we then end up with two SLR models that relate  $SATM$  to  $GPA$ , one model for *males* ( $\widehat{FYGPA}_i = 0.216 + 0.0386 \text{SATM}_i$ ) and one for *females* ( $\widehat{FYGPA}_i = 0.529 + 0.0386 \text{SATM}_i$ ). The only difference between these two models is in the y-intercept, with the *female* model's y-intercept shifted up from the *male* y-intercept by 0.313. And that is what adding indicator variables into models does in general<sup>15</sup> – it shifts the intercept up or down from the baseline group (here selected as *males*) to get a new intercept for the deviation group (here *females*).

To make this visually clearer, Figure 8.20 contains the regression lines that were estimated for each group. For any  $SATM$ , the difference in the groups is the 0.313 coefficient from the `SEXFEMALE` or `SEXINDICATOR` row of the model summaries. For example, at  $SATM=50$ , the difference in terms of predicted average first year GPAs between males and females is displayed as a difference between 2.15 and 2.46. This model assumes that the slope on  $SATM$  is the same for both groups except that they are allowed to have different y-intercepts, which is reasonable here because we saw approximately parallel relationships for the two groups in Figure 8.19.

Remember that `1m` selects baseline categories typically based on the alphabetical order of the levels of the categorical variable. Here, the `SEX` variable started with a coding of 1 and 2 and retained that order even with the recoding of levels that we created to give it more explicit names. Because we allow `1m` to create

<sup>15</sup>This is true for additive uses of indicator variables. In Section 8.11, we consider interactions between quantitative and categorical variables which has the effect of changing slopes and intercepts. The simplification ideas to produce estimated equations for each group are used there but we have to account for changing slopes by group too.

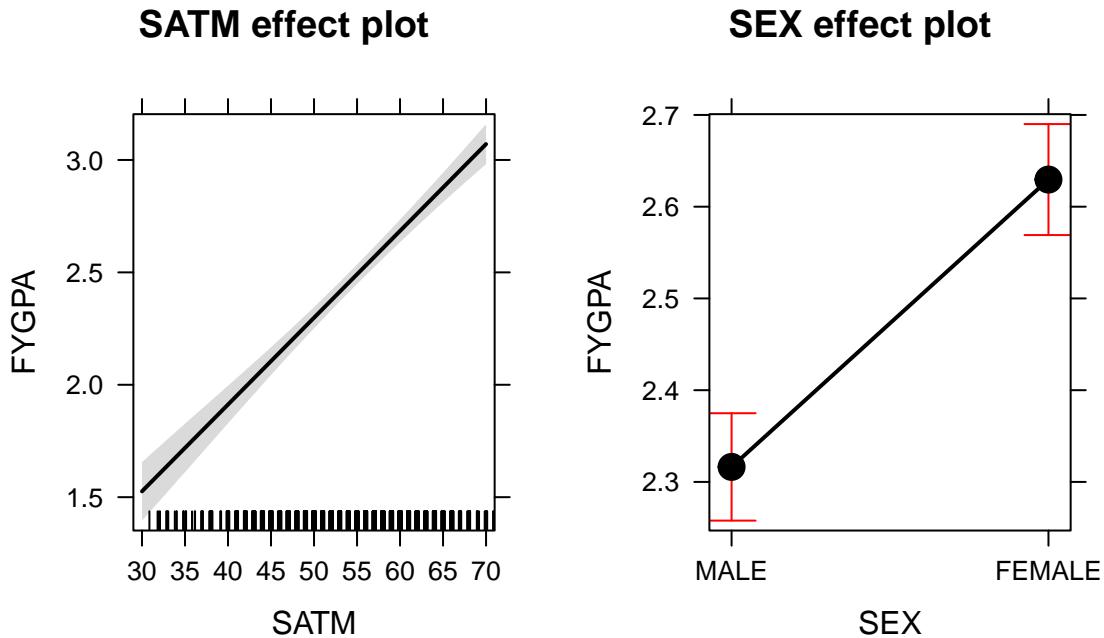


Figure 8.21: Term-plots for the estimated model for  $\text{FYGPA} \sim \text{SATM} + \text{SEX}$ .

indicator variables for us, the main thing you need to do is explore the model summary and look for the hint at the baseline level that is not displayed after the name of the categorical variable.

We can also work out the impacts of adding an indicator variable to the model in general in the theoretical model with a single quantitative predictor  $x_i$  and indicator  $I_i$ . The model starts as

$$y_i = \beta_0 + \beta_1 x_i + \beta_2 I_i + \varepsilon_i.$$

Again, there are two versions:

- For any observation  $i$  in the **baseline** category,  $I_i = 0$  and the model is  $y_i = \beta_0 + \beta_1 x_i + \varepsilon_i$ .
- For any observation  $i$  in the **non-baseline (deviation)** category,  $I_i = 1$  and the model simplifies to  $y_i = (\beta_0 + \beta_2) + \beta_1 x_i + \varepsilon_i$ .
  - This model has a y-intercept of  $\beta_0 + \beta_2$ .

The interpretation and inferences for  $\beta_1$  resemble the work with any MLR model, noting that these results are “controlled for”, “adjusted for”, or “allowing for differences based on” the categorical variable in the model. The interpretation of  $\beta_2$  is as a shift up or down in the y-intercept for the model that includes  $x_i$ . When we make term-plot in a model with a quantitative and additive categorical variable, the two reported model components match with the previous discussion – the same estimated term from the quantitative variable for all observations and a shift to reflect the different y-intercepts in the two groups. In Figure 8.21, the females are estimated to be that same 0.313 points higher on first year GPA. The males have a mean GPA slightly above 2.3 which is the predicted GPA for the average SATM percentile (remember that we have to hold the other variable at its mean to make each term-plot)<sup>16</sup>.

```
require(effects)
plot(allEffects(SATSex1))
```

<sup>16</sup>When making the SATM term-plot, the categorical variable is held at the most frequently occurring value in the data set. If you drop `ci.style="lines"` from the effect plot options, it is best to copy the figures as Bitmaps or save them as an image or they may (for some reason) lose the shaded bands.

The model summary and confidence intervals provide some potential interesting inferences in these models. Again, these are just applications of MLR methods we have already seen except that the definition of one of the variables is “different” using the indicator coding idea. For the same model, the **SEX** coefficient can be used to generate inferences for differences in the mean the groups, controlling for their *SATM* scores.

```
## SEXFEMALE    0.31322   0.04360   7.184 1.32e-12
```

Testing the null hypothesis that  $H_0 : \beta_2 = 0$  vs  $H_A : \beta_2 \neq 0$  using our regular *t*-test provides the opportunity to test for a difference in intercepts between the groups. In this situation, the test statistic is  $t = 7.184$  and, based on a  $t_{997}$ -distribution if the null is true, the p-value is  $< 0.0001$ . We can reject the null hypothesis because the chances of getting a difference in y-intercepts extreme or more extreme than what was observed is extremely small. Thus, we can conclude that there is evidence that there is a difference in the true y-intercept in a *SATM* model between *males* and *females*. The confidence interval is also informative:

```
confint(SATSex1)
```

```
##                 2.5 %      97.5 %
## (Intercept) -0.07566665  0.50744709
## SATM        0.03355273  0.04367726
## SEXFEMALE   0.22766284  0.39877160
```

We are 95% confident that the true mean GPA for females is between 0.228 and 0.399 points higher than for males, after adjusting for the *SATM* in the population of students. If we had subset the data set and fit two SLRs, we could have obtained the same simplified regression models but we never could have performed inferences for the differences between the two groups without putting all the observations together in one model and then assessing those differences with targeted coefficients. We also would not be able to get an estimate of their common slope for *SATM*, after adjusting for differences in the intercept for each group.

## 8.10 Additive MLR with more than two groups: Headache example

The same techniques can be extended to more than two groups. A study was conducted to explore sound tolerances using  $n = 98$  subjects with the data available in the **Headache** data set from the **heplots** package. Each subject was initially exposed to a tone, stopping when the tone became definitely intolerable (*DU*) and that decibel level was recorded (variable called *du1*). Then the subjects were randomly assigned to one of four treatments: *T1* (Listened again to the tone at their initial *DU* level, for the same amount of time they were able to tolerate it before); *T2* (Same as *T1*, with one additional minute of exposure); *T3* (Same as *T2*, but the subjects were explicitly instructed to use the relaxation techniques); and *Control* (these subjects experienced no further exposure to the noise tone until the final sensitivity measures were taken). Then the *DU* was measured again (variable called *du2*). One would expect that there would be a relationship between the upper tolerance levels of the subjects before and after treatment. But maybe the treatments impact that relationship? We can use our indicator approach to see if the treatments provide a shift to higher tolerances after accounting for the relationship between the two measurements. The scatterplot of the results in Figure 8.22 shows some variation in the slopes and the intercepts for the groups although the variation in intercepts seems more prominent than differences in slopes.

```
require(heplots)
data(Headache)
scatterplot(du2~du1|treatment, data=Headache, smooth=F, lwd=2,
           main="Plot of Maximum DB tolerances before and after treatment (by treatment)",
           legend.coords="topleft")
```

This data set contains a categorical variable with 4 levels. To go beyond two groups, we have to add more than one indicator variable, defining three indicators to turn on (1) or off (0) for three of the levels of the variable with the same reference level used for all the indicators. For this example, the *T1 Treatment* group

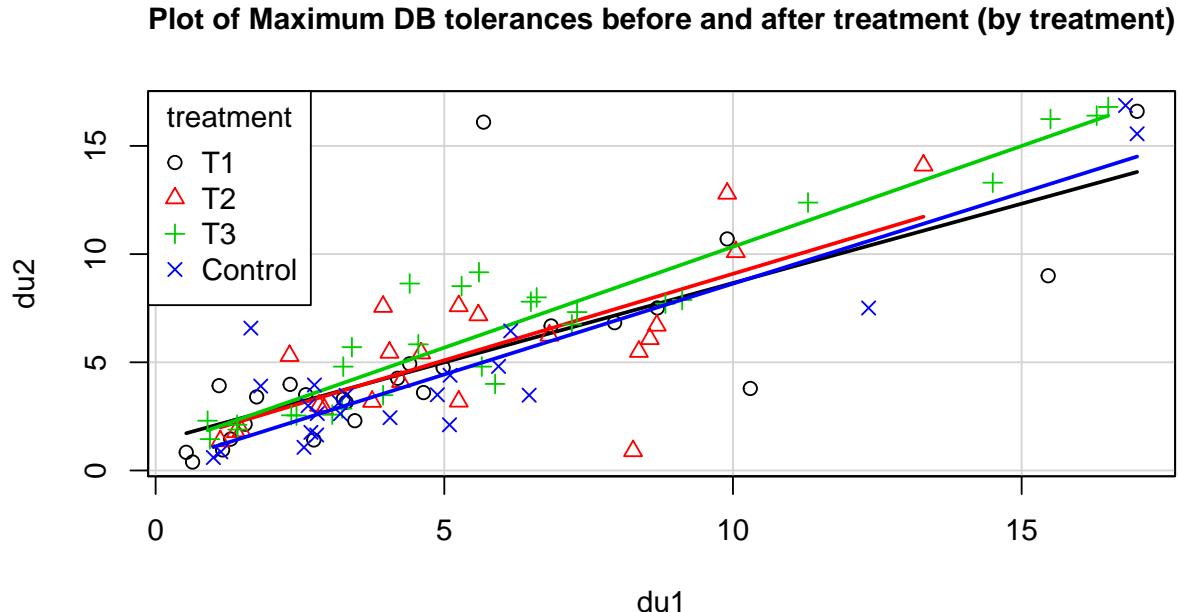


Figure 8.22: Scatterplot of post-treatment decibel tolerance ( $du_2$ ) vs pre-treatment tolerance ( $du_1$ ) by treatment level (4 groups).

is chosen as the baseline group so it sort of hides in the background while we define indicators for the other three levels. The indicators for  $T_2$ ,  $T_3$ , and *Control* levels are:

- Indicator for  $T_2$ :  $I_{T_2,i} = \begin{cases} 1 & \text{if Treatment} = T_2 \\ 0 & \text{else} \end{cases}$
- Indicator for  $T_3$ :  $I_{T_3,i} = \begin{cases} 1 & \text{if Treatment} = T_3 \\ 0 & \text{else} \end{cases}$
- Indicator for *Control*:  $I_{\text{Control},i} = \begin{cases} 1 & \text{if Treatment} = \text{Control} \\ 0 & \text{else} \end{cases}$

We can see the values of these indicators for a few observations and their original variable (`treatment`) in the following output. The bolded observations show each of the indicators being “turned on”. For  $T_1$ , all the indicators stay at 0.

Treatment	I_T2	I_T3	I_Control
<b>T3</b>	<b>0</b>	<b>1</b>	<b>0</b>
<b>T1</b>	<b>0</b>	<b>0</b>	<b>0</b>
T1	0	0	0
T3	0	1	0
T3	0	1	0
T3	0	1	0
<b>T2</b>	<b>1</b>	<b>0</b>	<b>0</b>
T1	0	0	0
T1	0	0	0
T3	0	1	0
T3	0	1	0
T2	1	0	0

Treatment	I_T2	I_T3	I_Control
T3	0	1	0
T1	0	0	0
T3	0	1	0
<b>Control</b>	<b>0</b>	<b>0</b>	<b>1</b>
T3	0	1	0

When we fit the additive model of the form  $y \sim x + \text{group}$ , the `lm` function takes the  $J$  categories and creates  $J - 1$  indicator variables. The baseline level is always handled in the intercept. The model will be of the form

$$y_i = \beta_0 + \beta_1 x_i + \beta_2 I_{\text{Level}2,i} + \beta_3 I_{\text{Level}3,i} + \cdots + \beta_{J-1} I_{\text{Level}J,i} + \varepsilon_i$$

where the  $I_{\text{CatName}_j,i}$ 's are the different indicator variables. Note that each indicator variable gets a coefficient associated with it and is “turned on” whenever the  $i^{\text{th}}$  observation is in that category. Only one of the  $I_{\text{CatName}_j,i}$ 's is a 1 for any observation, so the  $y$ -intercept will either be  $\beta_0$  for the baseline group or  $\beta_0 + \beta_j$  for  $j = 2, \dots, J$ . It is important to remember that this is an “additive” model since the effects just add and there is no interaction between the grouping variable and the quantitative predictor. To be able to trust this model, we need to check that we do not need different slope coefficients for the groups as discussed in the next section.

For these types of models, it is always good to start with a plot of the data set with regression lines for each group – assessing whether the lines look relatively parallel or not. In Figure 8.22, there are some differences in slopes – we investigate that further in the next section. For now, we can proceed with fitting the additive model with different intercepts for the four levels of `treatment` and the quantitative explanatory variable, `du1`.

```
head1 <- lm(du2 ~ du1 + treatment, data=Headache)
summary(head1)

##
## Call:
## lm(formula = du2 ~ du1 + treatment, data = Headache)
##
## Residuals:
##     Min      1Q  Median      3Q     Max 
## -6.9085 -0.9551 -0.3118  1.1141 10.5364 
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) 0.80918   0.50095  1.615   0.110    
## du1          0.83705   0.05176 16.172  <2e-16 ***
## treatmentT2  0.07692   0.62622  0.123   0.903    
## treatmentT3  0.80919   0.59271  1.365   0.175    
## treatmentControl -0.55752   0.61830 -0.902   0.370    
## 
## Residual standard error: 2.14 on 93 degrees of freedom
## Multiple R-squared:  0.7511, Adjusted R-squared:  0.7404 
## F-statistic: 70.16 on 4 and 93 DF,  p-value: < 2.2e-16
```

The complete estimated regression model is

$$\widehat{\text{du2}}_i = 0.809 + 0.837 \text{ du1}_i + 0.077 I_{\text{T2},i} + 0.809 I_{\text{T3},i} - 0.558 I_{\text{Control},i}.$$

For each group, the model simplifies to an SLR as follows:

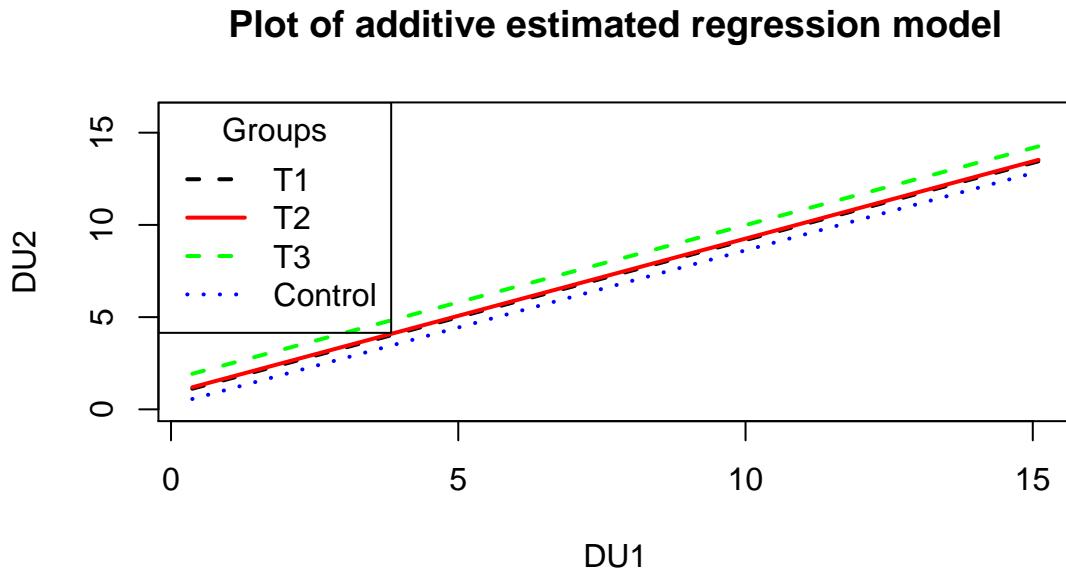


Figure 8.23: Plot of estimated noise tolerance additive model.

- For  $T1$  (baseline):

$$\begin{aligned}\widehat{\text{du2}}_i &= 0.809 + 0.837 \text{ du1}_i + 0.077I_{T2,i} + 0.809I_{T3,i} - 0.558I_{\text{Control},i} \\ &= 0.809 + 0.837 \text{ du1}_i + 0.077 * 0 + 0.809 * 0 - 0.558 * 0 \\ &= 0.809 + 0.837 \text{ du1}_i\end{aligned}$$

- For  $T2$ :

$$\begin{aligned}\widehat{\text{du2}}_i &= 0.809 + 0.837 \text{ du1}_i + 0.077I_{T2,i} + 0.809I_{T3,i} - 0.558I_{\text{Control},i} \\ &= 0.809 + 0.837 \text{ du1}_i + 0.077 * 1 + 0.809 * 0 - 0.558 * 0 \\ &= 0.809 + 0.837 \text{ du1}_i + 0.077 \\ &= 0.886 + 0.837 \text{ du1}_i\end{aligned}$$

- For  $T3$ :

$$\widehat{\text{du2}}_i = 1.618 + 0.837 \text{ du1}_i$$

- For  $\text{Control}$ :

$$\widehat{\text{du2}}_i = 0.251 + 0.837 \text{ du1}_i$$

To reinforce what this additive model is doing, Figure 8.23 displays the estimated regression lines for all four groups, showing the shifts in the  $y$ -intercepts among the groups.

The term-plot (Figure 8.24) shifted up the most relative to the others and the *Control* group seems to be noticeably lower than the others, in the model that otherwise assumes that the same relationship holds between  $\text{du1}$  and  $\text{du2}$  for all the groups. After controlling for the *Treatment* group, for a 1 decibel increase in initial tolerances, we expect, on average, to obtain a 0.84 decibel change in the second tolerance measurement. The  $R^2$  shows that this is a decent model for the responses, with this model explaining 75.1% percent of the variation in the second decibel tolerance measure. We should check the diagnostic plots and VIFs to check for

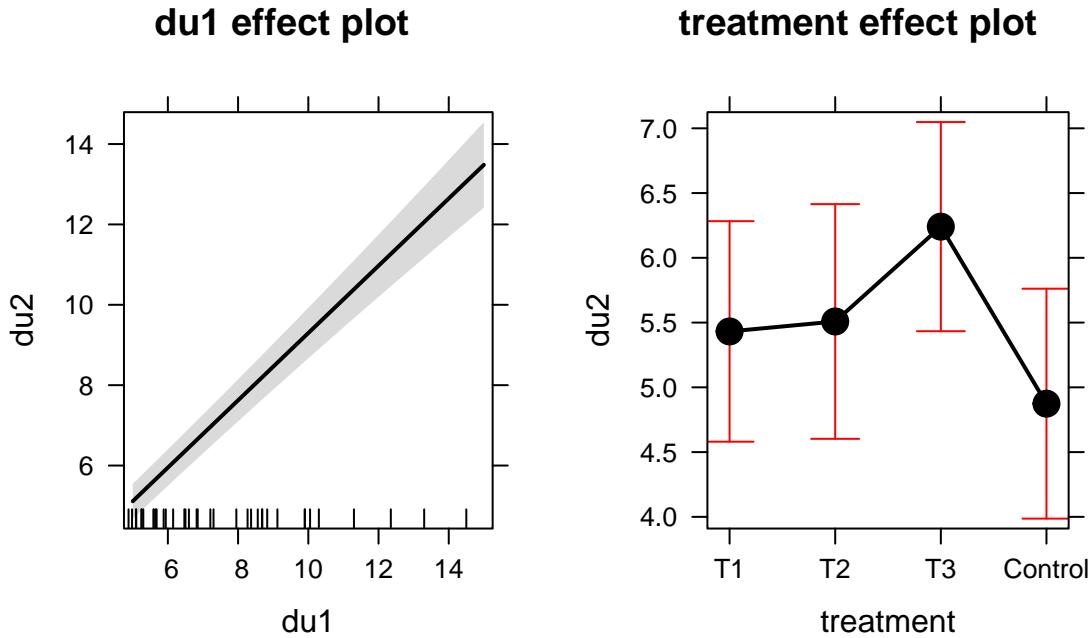


Figure 8.24: Term-plots of the additive decibel tolerance model.

any issues – all the diagnostics and assumptions are as before except that there is no assumption of linearity between the grouping variable and the responses. Additionally, sometimes we need to add group information to diagnostics to see if any patterns look different in different groups, like linearity or non-constant variance.

```
plot(allEffects(head1))
```

The diagnostic plots in Figure 8.25 provides some indications of a few observations in the tails that deviate from a normal to having slightly heavier tails but only one outlier is of real concern. There is a small indication of increasing variability as a function of the fitted values as both the Residuals vs. Fitted and Scale-Location plots show some fanning out for higher values but this is a minor issue. There are no influential points in the data set.

```
par(mfrow=c(2,2), oma=c(0,0,2,0))
plot(head1,
     sub.caption="Plot of diagnostics for additive model with du1 and treatment for du2")
```

The VIFs are different for categorical variables than for quantitative predictors in MLR. The 4 levels are combined in a measure called the ***generalized VIF (GVIF***). For GVIFs, we only focus on the inflation of the SE scale (square root for 1 df effects and raised to the power  $1/(2 * J)$  for a  $J$ -level predictor. On this scale, the interpretation is as **the multiplicative increase in the SEs due to multicollinearity with other predictors**. In this model, the SE for du1 is 1.009 times larger due to multicollinearity with other predictors and the SEs for the categorical variable are 1.003 times larger due to multicollinearity than they otherwise would have been. Neither are large so multicollinearity is not a problem in this model.

```
require(car)
vif(head1)
```

```
##          GVIF Df GVIF^(1/(2*Df))
## du1      1.01786  1      1.008891
## treatment 1.01786  3      1.002955
```

While there are inferences available in the model output, the tests for the indicator variables are not too informative since they only compare each group to the baseline. In Section 8.12, we see how to use ANOVA  $F$ -tests to help us ask general questions about including a predictor in the model. But we can compare

### Plot of diagnostics for additive model with du1 and treatment for du2

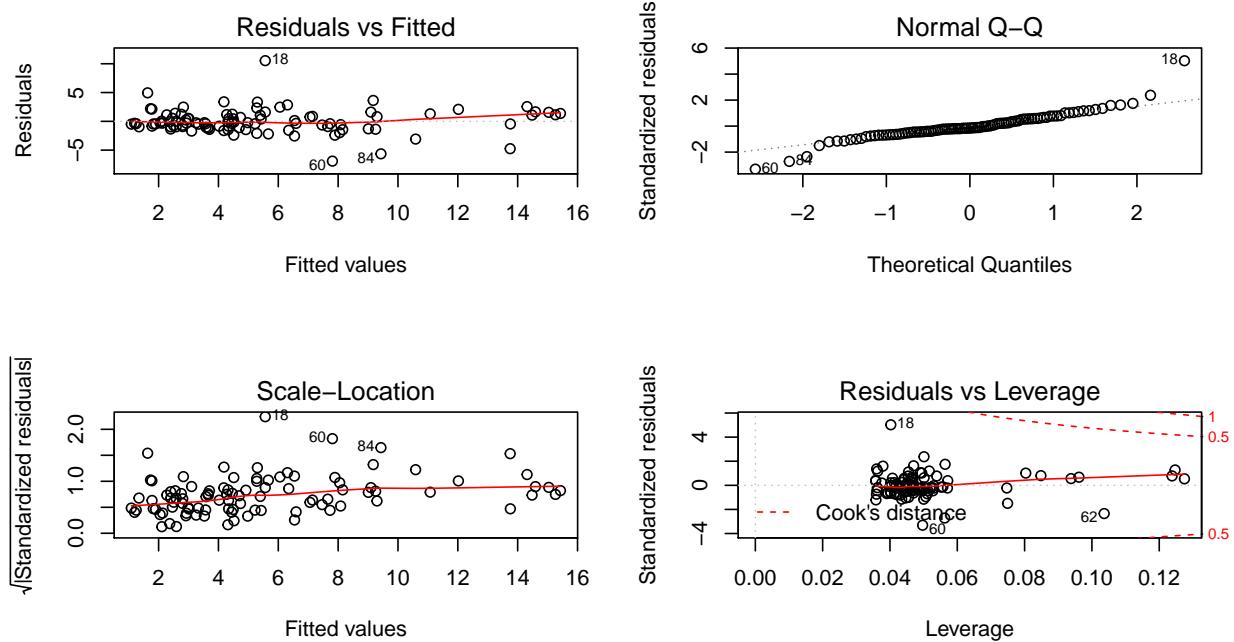


Figure 8.25: Diagnostic plots for the additive decibel tolerance model.

adjusted  $R^2$  values with and without *Treatment* to see if including the categorical variable was “worth it”:

```
head1R <- lm(du2 ~ du1, data=Headache)
summary(head1R)
```

```
##
## Call:
## lm(formula = du2 ~ du1, data = Headache)
##
## Residuals:
##      Min       1Q   Median       3Q      Max 
## -6.9887 -0.8820 -0.2765  1.1529 10.4165 
## 
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) 0.84744    0.36045   2.351   0.0208  
## du1         0.85142    0.05189  16.408  <2e-16 ***
## 
## Residual standard error: 2.165 on 96 degrees of freedom
## Multiple R-squared:  0.7371, Adjusted R-squared:  0.7344 
## F-statistic: 269.2 on 1 and 96 DF,  p-value: < 2.2e-16
```

The adjusted  $R^2$  in the model with both *Treatment* and *du1* is 0.7404 and the adjusted  $R^2$  for this reduced model with just *du1* is 0.7344, suggesting the *Treatment* is useful. The next section provides a technique to be able to work with different slopes on the quantitative predictor for each group. Comparing those results to the results for the additive model allows assessment of the assumption in this section that all the groups had the same slope coefficient for the quantitative variable.

## 8.11 Different slopes and different intercepts

Sometimes researchers are specifically interested in whether the slopes vary across groups or the regression lines in the scatterplot for the different groups may not look parallel or it may just be hard to tell visually if there really is a difference in the slopes. Unless you are **very sure** that there is not an interaction between the grouping variable and the quantitative predictor, you should start by fitting a model containing an interaction and then see if you can drop it. It may be the case that you end up with the simpler additive model from the previous sections, but you don't want to assume the same slope across groups unless you are absolutely sure that is the case. This should remind you a bit of the discussions of the additive and interacting models in the Two-way ANOVA material. The models, concerns, and techniques are very similar, but with the quantitative variable replacing one of the two categorical variables. As always, the scatterplot is a good first step to understanding whether we need the extra complexity that these models require.

A new example provides motivation for the consideration of different slopes and intercepts. A study was performed to address whether the relationship between nonverbal IQs and reading accuracy differs between dyslexic and non-dyslexic students. Two groups of students were identified one group of *dyslexic* students was identified first (19 students) and then a group of gender and age similar student matches were identified (25 students) for a total sample size of  $n = 44$ , provided in the *dyslexic3* data set from the *smdata* package (Merkle and Smithson, 2013). This type of study design is an attempt to “balance” the data from the two groups on some important characteristics to make the comparisons of the groups as fair as possible. The researchers attempted to balance the characteristics of the subjects in the two groups so that if they found different results for the two groups, they could attribute it to the main difference they used to create the groups – dyslexia or not. This design, **case-control** or **case-comparison** where each subject with a trait is matched to one or more subjects in the “control” group would hopefully reduce confounding from other factors and then allow stronger conclusions in situations where it is impossible to randomly assign treatments to subjects. We still would avoid using “causal” language but this design is about as good as you can get when you are unable to randomly assign levels to subjects.

Using these data, we can explore the relationship between nonverbal IQ scores and reading accuracy, with reading accuracy measured as a proportion correct. The fact that there is an upper limit to the response variable attained by many students will cause complications below, but we can still learn something from our attempts to analyze these data using an MLR model. The scatterplot in Figure 8.26 seems to indicate some clear differences in the *IQ* vs *reading score* relationship between the *dys=0* (non-dyslexic) and *dys=1* (dyslexic) students. Note that the IQ is standardized to have mean 0 and standard deviation of 1 which means that a 1 unit change in IQ score is a 1 SD change and that the *y*-intercept (for  $x = 0$ ) is right in the center of the plot and actually interesting.

```
require(smdata)
data("dyslexic3")
scatterplot(score~ziq|dys, xlab="Standardized nonverbal IQ scores",
            ylab="Reading score", data=dyslexic3, smooth=F,
            main="Plot of IQ vs Reading by dyslexia status")
```

To allow for both different y-intercepts and slope coefficients on the quantitative predictor, we need to include a “modification” of the slope coefficient. This is performed using an **interaction** between the two predictor variables where we allow the impacts of one variable (slopes) to change based on the levels of another variable (grouping variable). The formula notation is  $y \sim x * group$ , remembering that this also includes the **main effects** (the additive variable components) as well as the interaction coefficients as we discussed in the Two-Way ANOVA interaction model. We can start with the general model for a two-level categorical variable with an interaction, which is

$$y_i = \beta_0 + \beta_1 x_i + \beta_2 I_{\text{CatName},i} + \beta_3 I_{\text{CatName},i} x_i + \varepsilon_i,$$

where the new component involves both the indicator and the quantitative predictor variable. The  $\beta_3$  coefficient will be found in a row of output with **both** variable names in it (with the indicator level name)

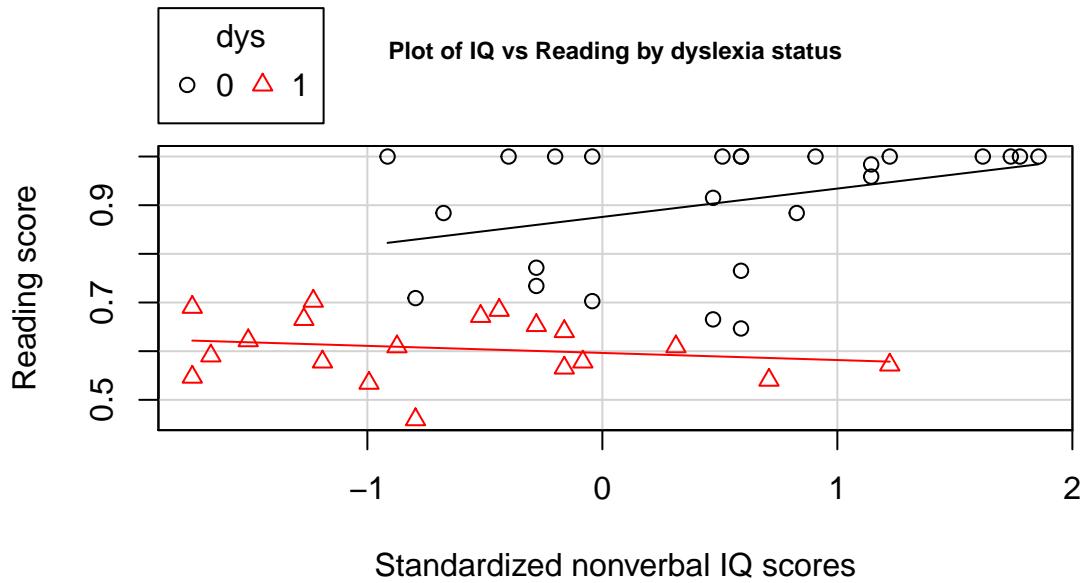


Figure 8.26: Scatterplot for reading score versus nonverbal IQ by dyslexia group.

with a colon between them (something like  $x:\text{grouplevel}$ ). As always, the best way to understand any model involving indicators is to plug in 0s or 1s for the indicator variable(s) and simplify the equations.

- For any observation in the baseline group  $I_{\text{CatName},i} = 0$ , so

$$y_i = \beta_0 + \beta_1 x_i + \beta_2 I_{\text{CatName},i} + \beta_3 I_{\text{CatName},i} x_i + \varepsilon_i$$

simplifies quickly to:

$$y_i = \beta_0 + \beta_1 x_i + \varepsilon_i.$$

- So the baseline group's model involves the initial intercept and quantitative slope coefficient.
- For any observation in the second category  $I_{\text{CatName},i} = 0$ , so

$$y_i = \beta_0 + \beta_1 x_i + \beta_2 I_{\text{CatName},i} + \beta_3 I_{\text{CatName},i} x_i + \varepsilon_i$$

is

$$y_i = \beta_0 + \beta_1 x_i + \beta_2 * 1 + \beta_3 * 1 * x_i + \varepsilon_i$$

which “simplifies” to

$$y_i = (\beta_0 + \beta_2) + (\beta_1 + \beta_3)x_i + \varepsilon_i,$$

by combining like terms.

- For the second category, the model contains a modified y-intercept, now  $\beta_0 + \beta_2$ , and a modified slope coefficient, now  $\beta_1 + \beta_3$ .

We can make this more concrete by applying this to the dyslexia data with `dys` as a categorical variable for dyslexia status of subjects (levels of 0 and 1) and `ziq` the standardized IQ. The estimated model is:

```
#Because dys was just numerically coded - makes it a factor
dyslexic3$dys <- factor(dyslexic3$dys)
dys_model <- lm(score ~ ziq * dys, data=dyslexic3)
summary(dys_model)
```

```
##
## Call:
## lm(formula = score ~ ziq * dys, data = dyslexic3)
##
## Residuals:
##      Min       1Q   Median       3Q      Max 
## -0.26362 -0.04152  0.01682  0.06790  0.17740 
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept)  0.87586   0.02391 36.628 < 2e-16  
## ziq         0.05827   0.02535  2.299  0.0268    
## dys1        -0.27951   0.03827 -7.304 7.11e-09  
## ziq:dys1    -0.07285   0.03821 -1.907  0.0638    
## 
## Residual standard error: 0.1017 on 40 degrees of freedom
## Multiple R-squared:  0.712, Adjusted R-squared:  0.6904 
## F-statistic: 32.96 on 3 and 40 DF, p-value: 6.743e-11
```

The estimated model can be written as

$$\widehat{\text{Score}}_i = 0.876 + 0.058 \text{ ZIQ}_i - 0.280 I_{\text{Level } 1,i} - \mathbf{0.073} I_{\text{Level } 1,i} \text{ ZIQ}_i$$

and simplified for the two groups as:

- For the baseline (non-dyslexic,  $I_{\text{Level } 1,i} = 0$ ) students:

$$\widehat{\text{Score}}_i = 0.876 + 0.058 \text{ ZIQ}_i$$

- For the deviation (dyslexic,  $I_{\text{Level } 1,i} = 1$ ) students:

$$\begin{aligned}\widehat{\text{Score}}_i &= 0.876 + 0.058 \text{ ZIQ}_i - 0.280 * 1 - 0.073 * 1 \text{ ZIQ}_i \\ &= (0.876 - 0.280) + (0.058 - 0.073) \text{ ZIQ}_i,\end{aligned}$$

which simplifies finally to:

$$\widehat{\text{Score}}_i = 0.596 + 0.015 \text{ ZIQ}_i$$

- So the slope switched from 0.058 in the non-dyslexic students to -0.015 in the dyslexic students. The interpretations of these coefficients are outlined below:
  - For the non-dyslexic students: For a 1 SD increase in verbal IQ score, we expect, on average, for the mean reading score to go up by 0.058 “points”.

### Plot of diagnostics for Dyslexia Interaction model

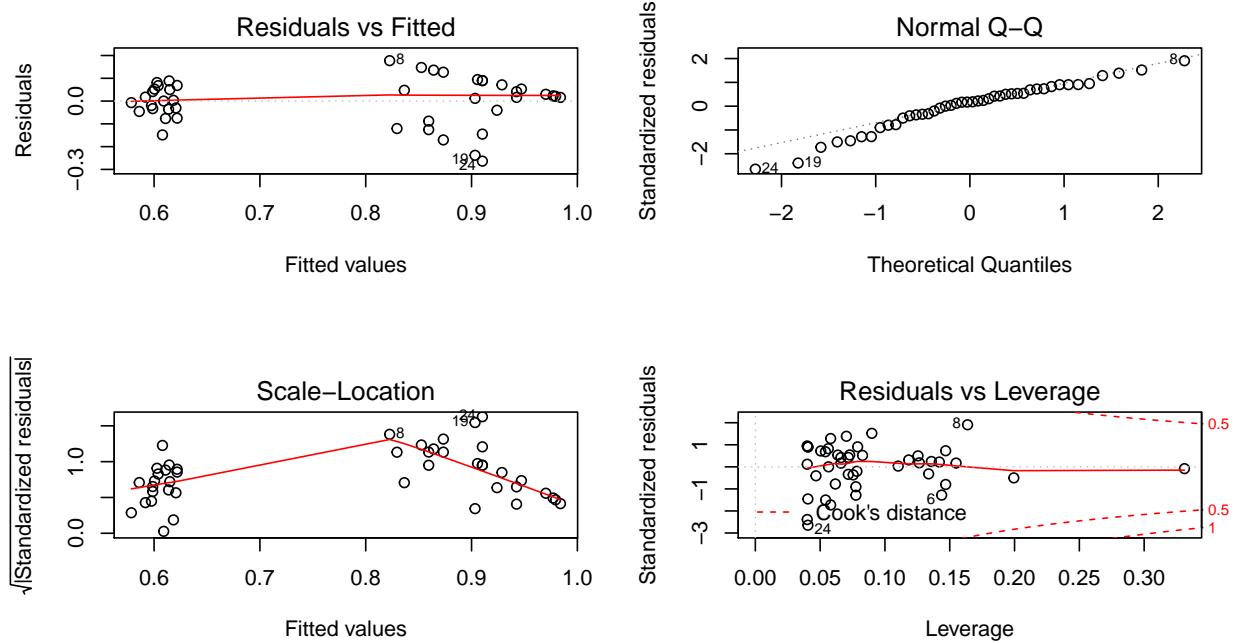


Figure 8.27: Diagnostic plots for interaction model for reading scores.

- For the dyslexic students: For a 1 SD increase in verbal IQ score, we expect, on average, for the mean reading score to change by -0.015 “points”.

So, an expected pattern of results emerges for the non-dyslexic students. Those with higher IQs tend to have higher reading accuracy; this does not mean higher IQ's cause more accurate reading because random assignment of IQ is not possible. However, for the dyslexic students, the relationship is not what one would might expect. It is slightly negative, showing that higher IQ's are related to lower reading accuracy. What we conclude from this is that we should not expect higher IQ's to show higher performance on a test like this.

Checking the assumptions is always recommended before getting focused on the inferences in the model. When fitting models with multiple groups, it is possible to see “groups” in the fitted values and that is not a problem – it is a feature of these models. You should look for issues in the residuals for each group but the residuals should overall still be normally distributed and have the same variability everywhere. It is a bit hard to see issues in Figure 8.27 because of the group differences, but note the line of residuals for the higher fitted values. This is an artifact of the upper threshold in the reading accuracy test used. As in the first year of college GPA, these observations were **censored** – their true score was outside the range of values we could observe – and so we did not really get a measure of how good these students were since a lot of their abilities were higher than the test could detect and they all binned up at the same value. The relationship in this group might be even stronger if we could really observe differences in the highest level readers. We should treat the results for the non-dyslexic group with caution even though they are clearly scoring on average higher and have a different slope than the results for the dyslexic students. The normality and influence diagnostics do not suggest any major issues other than this.

```
par(mfrow=c(2,2), oma=c(0,0,2,0))
plot(dys_model,
      sub.caption="Plot of diagnostics for Dyslexia Interaction model")
```

For these models, we have relaxed an earlier assumption that data were collected from only one group. In fact,

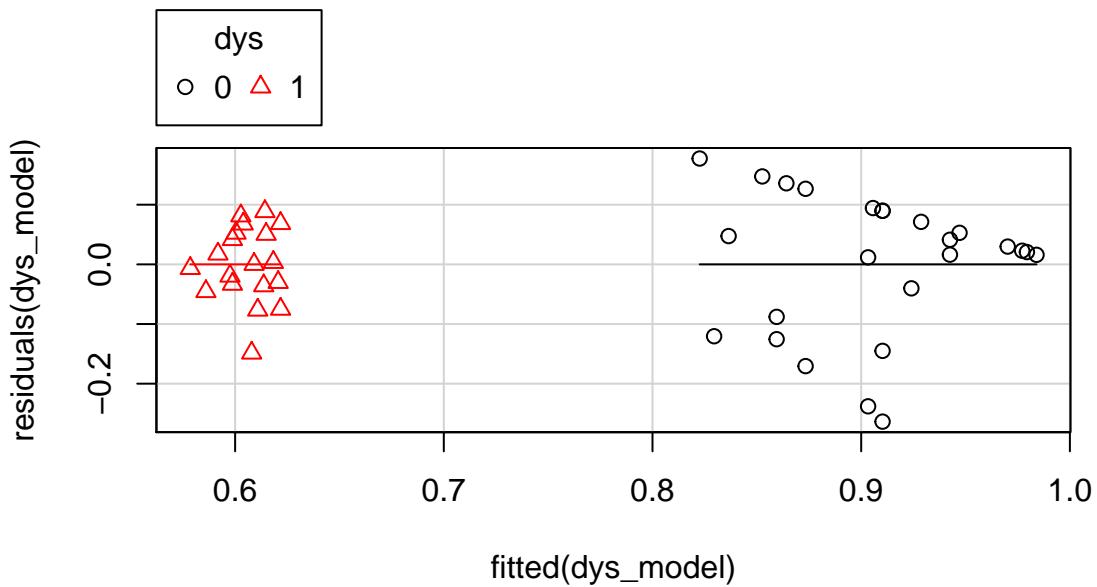


Figure 8.28: Plot of Residuals vs Fitted from interaction dyslexia data model with groups indicated.

we are doing specific research that is focused on questions about the differences between groups. However, these models still make assumptions that, within a specific group, the linearity assumption is met. They also assume that the variability in the residuals is the same for all observations. Sometimes it can be difficult to check the assumptions by looking at the overall diagnostic plots and it may be easier to go back to the original scatterplot or plot the residuals vs fitted values by group to fully assess the results. For example, Figure 8.28 shows a scatterplot of the residuals vs the quantitative explanatory variable by the groups. The variability in the residuals is a bit larger in the non-dyslexic group, possibly suggesting that variability in the reading test is higher for higher scoring individuals even though we couldn't observe all of that variability because there were so many perfect scores in this group.

```
scatterplot(residuals(dys_model)~fitted(dys_model)|dys,
           data=dyslexic3, smooth=F)
```

If we feel comfortable enough with the assumptions to trust the inferences here (this might be dangerous), then we can consider what some of the model inferences provide us within this situation. For example, the test for  $H_0 : \beta_3 = 0$  vs  $H_A : \beta_3 \neq 0$  provides an interesting comparison. Under the null hypothesis, the two groups would have the same slope so it provides an opportunity to directly consider whether the relationship (via the slope) is different between the groups in their respective populations. We find  $t = -1.907$  which, if the assumptions are met, follows a  $t(40)$ -distribution under the null hypothesis. This test statistic has a corresponding p-value of 0.0638. Depending on your standards of evidence, this might be sufficient evidence or it might not. It certainly provides some evidence of a difference in the slopes and it isn't strong evidence of that. There are serious issues (like getting the wrong idea about directions of relationships) if we ignore a potentially important interaction and some statisticians would recommend retaining interactions even if the evidence is only moderate for its inclusion in the model. For the original research question of whether the relationships differ for the two groups, we only have marginal evidence to support that result. Possibly with a larger sample size or a reading test that only a few students could get 100% on, the researchers might have detected a more pronounced difference in the slopes for the two groups.

In the presence of a categorical by quantitative interaction, term-plots can be generated that plot the results for each group on the same display. This basically provides a plot of the “simplified” SLR models for each group. In Figure 8.29 we can see noticeable differences in the slopes and intercepts. Note that testing for

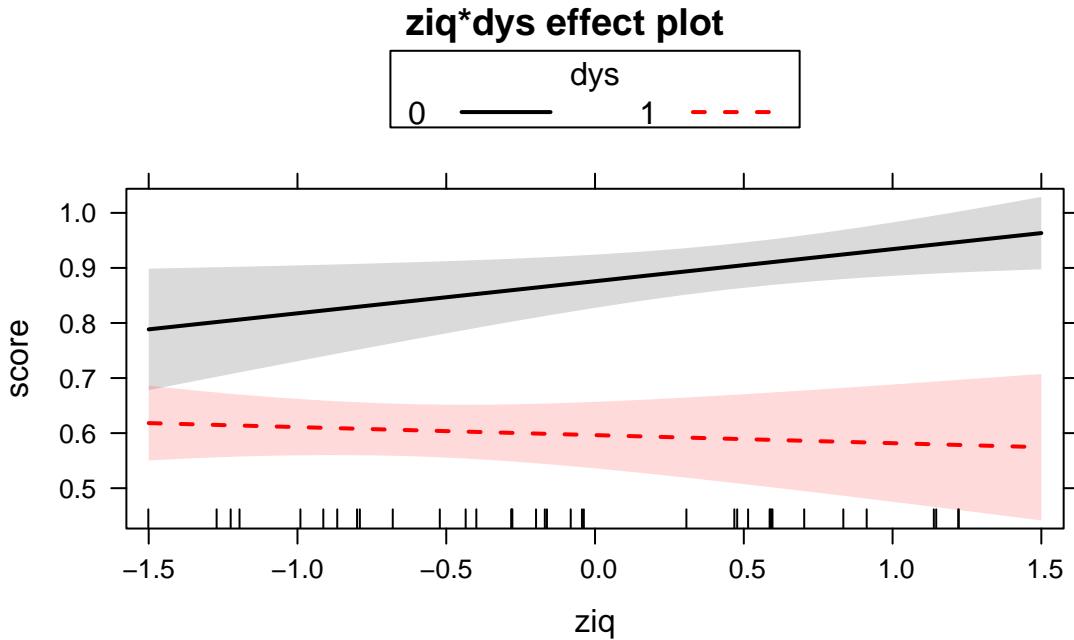


Figure 8.29: Term-plots for interaction model for reading scores.

differences in intercepts between groups is not very interesting when there are different slopes because if you change the slope, you have to change the intercept. The plot shows that there are clear differences in the means even though we don't have a test to directly assess that in this complicated of a model.

```
plot(allEffects(dys_model), ci.style="bands", multiline=T)
```

It certainly appears in the plots that IQ has a different impact on the mean score in the two groups (even though the p-value only provided marginal evidence). To reinforce the potential dangers of forcing the same slope for both groups, consider the additive model for these data. Again, this just shifts one group off the other one, but both have the same slope. The following model summary and term-plots (Figure 8.30 suggest the potentially dangerous conclusion that can come from assuming a common slope when that might not be the case.

```
dys_modelR <- lm(score ~ ziq + dys, data=dyslexic3)
summary(dys_modelR)
```

```
##
## Call:
## lm(formula = score ~ ziq + dys, data = dyslexic3)
##
## Residuals:
##    Min     1Q   Median     3Q    Max 
## -0.26062 -0.05565  0.02932  0.07577  0.13217
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) 0.89178   0.02312  38.580 < 2e-16 ***
## ziq         0.02620   0.01957   1.339   0.188    
## dys1        -0.26879   0.03905  -6.883 2.41e-08 ***
## 
## Residual standard error: 0.1049 on 41 degrees of freedom
## Multiple R-squared:  0.6858, Adjusted R-squared:  0.6705 
## F-statistic: 44.75 on 2 and 41 DF,  p-value: 4.917e-11
```

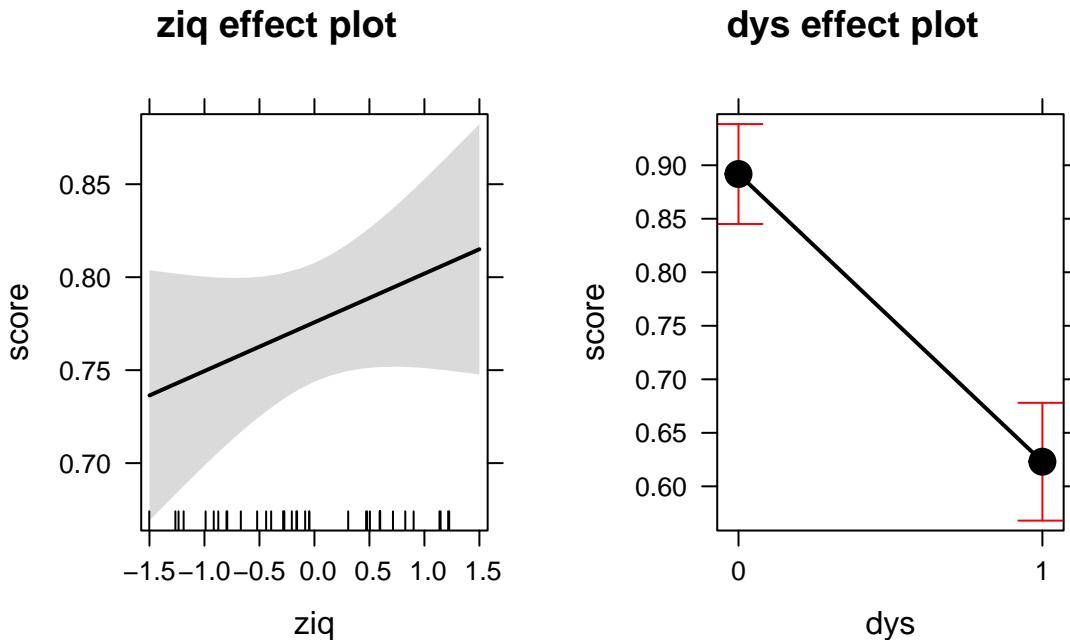


Figure 8.30: Term-plots for additive model for reading scores.

```
plot(allEffects(dys_modelR))
```

This model provides no evidence that IQ is related to reading score for all students ( $t_{41} = 1.34$ , p-value=0.188) but strong evidence of a difference in the y-intercepts ( $t_{41} = -6.88$ , p-value < 0.00001).

Since the IQ term has a large p-value, then we could drop it from the model – leaving a model that only includes the grouping variable:

```
dys_modelR2 <- lm(score~dys, data=dyslexic3)
summary(dys_modelR2)
```

```
##
## Call:
## lm(formula = score ~ dys, data = dyslexic3)
##
## Residuals:
##    Min     1Q   Median     3Q    Max 
## -0.25818 -0.04510  0.02514  0.09520  0.09694
## 
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept)  0.90480   0.02117  42.737 <2e-16 ***
## dys1        -0.29892   0.03222 -9.278  1e-11 ***
## 
## Residual standard error: 0.1059 on 42 degrees of freedom
## Multiple R-squared:  0.6721, Adjusted R-squared:  0.6643 
## F-statistic: 86.08 on 1 and 42 DF,  p-value: 1e-11
plot(allEffects(dys_modelR2))
```

These results, including the term-plot in Figure 8.31, show that there is evidence of a difference in the mean reading scores between the two groups and maybe that is all these data really say... This is the logical outcome if we decide that the interaction is not important IN THIS DATA SET. In general, if the interaction is dropped, the interaction model can be reduced to considering an additive model with the categorical

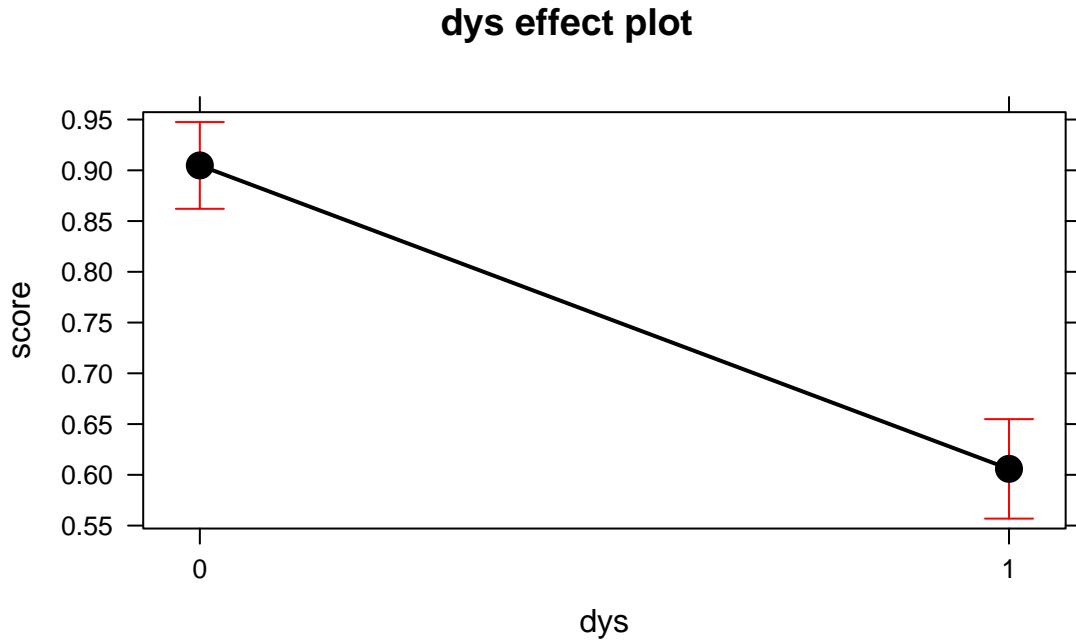


Figure 8.31: Term-plot for dyslexia status only model for reading scores.

and quantitative predictor variables. Either or both of those variables could also be considered for removal, possibly starting with the variable with the larger p-value, leaving a string of ever-simpler models possible if large p-values are continually encountered.

It is useful to note that the last model has returned us to the first model we encountered in Chapter 2 where we were just comparing the means for two groups. However, the researchers probably were not seeking to make the discovery that dyslexic students have a tougher time than non-dyslexic students on a reading test but sometimes that is all that the data support. The key part of this sequence of decisions was how much evidence you think a p-value of 0.06 contains...

For more than two categories in a categorical variable, the model contains more indicators to keep track of but uses the same ideas. We have to deal with modifying the deviation group so the task is onerous but relatively repetitive. The general model is:

$$y_i = \beta_0 + \beta_1 x_i + \beta_2 I_{\text{Level } 2,i} + \beta_3 I_{\text{Level } 3,i} + \cdots + \beta_J I_{\text{Level } J,i} + \beta_{J+1} I_{\text{Level } 2,i} x_i + \beta_{J+2} I_{\text{Level } 3,i} x_i + \cdots + \beta_{2J-1} I_{\text{Level } J,i} x_i + \varepsilon_i$$

Specific to the audible tolerance/headache data that had four groups. The model with an interaction present is

$$\text{du2}_i = \beta_0 + \beta_1 \text{du1}_i + \beta_2 I_{T2,i} + \beta_3 I_{T3,i} + \beta_4 I_{\text{Control},i} + \beta_5 I_{T2,i} \text{du1}_i + \beta_6 I_{T3,i} \text{du1}_i + \beta_7 I_{\text{Control},i} \text{du1}_i + \varepsilon_i$$

Based on the following output, the estimated general regression model is

$$\widehat{\text{du2}}_i = 1.33 + 0.733 \text{du1}_i - 0.236 I_{T2,i} - 0.316 I_{T3,i} - 1.091 I_{\text{Control},i} + 0.066 I_{T2,i} \text{du1}_i + 0.199 I_{T3,i} \text{du1}_i + 0.106 I_{\text{Control},i} \text{du1}_i$$

Then we could work out the specific equation for **each group** with replacing their indicator variable in two places with 1s and the rest of the indicators with 0. For example, for the *Control* group:

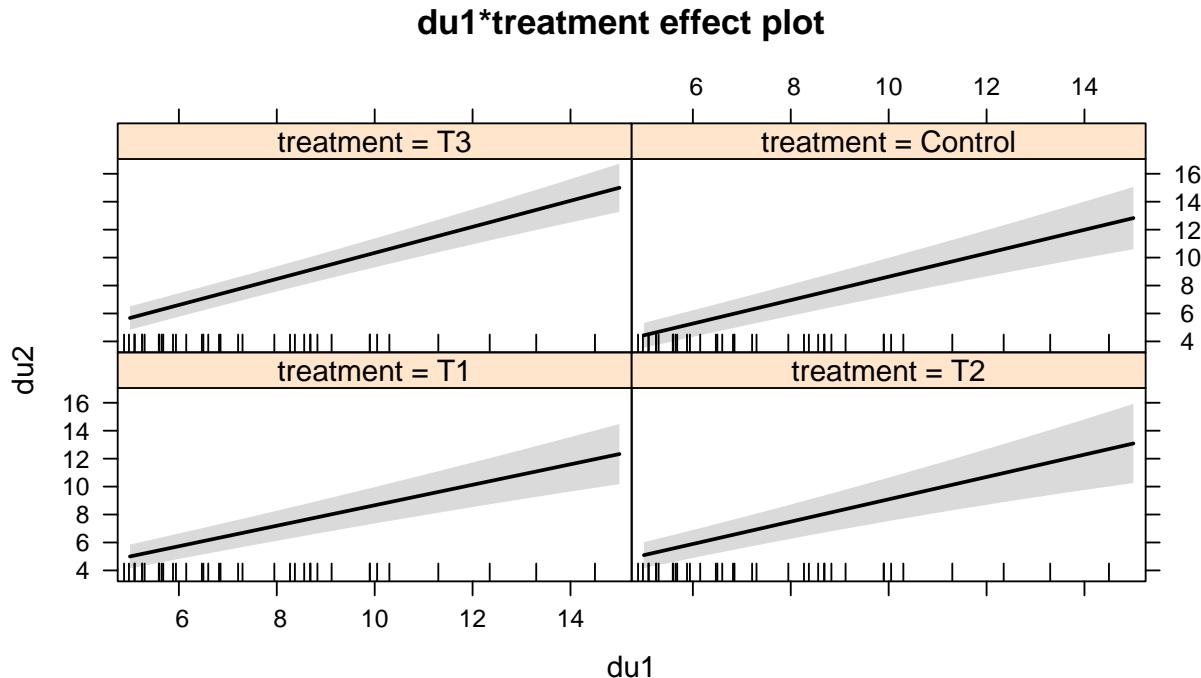


Figure 8.32: Term-plot for decibel tolerance interaction model (version 1).

$$\begin{aligned}\widehat{\text{du2}}_i &= 1.33 + 0.733 \text{ du1}_i - 0.236 * 0 - 0.316 * 0 - 1.091 * 1 \\ &\quad + 0.066 * 0 * \text{du1}_i + 0.199 * 0 * \text{du1}_i + 0.106 * 1 * \text{du1}_i \\ \widehat{\text{du2}}_i &= 1.33 + 0.733 \text{ du1}_i - 1.091 + 0.106 \text{ du1}_i \\ \widehat{\text{du2}}_i &= 0.239 + 0.839 \text{ du1}_i\end{aligned}$$

```
head2<-lm(du2~du1*treatment,data=Headache)
summary(head2)
```

```
##
## Call:
## lm(formula = du2 ~ du1 * treatment, data = Headache)
##
## Residuals:
##    Min      1Q  Median      3Q     Max 
## -6.8072 -1.0969 -0.3285  0.8192 10.6039 
## 
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) 1.33157   0.66027  2.017   0.0467  
## du1         0.73319   0.09969  7.355 8.53e-11 ***
## treatmentT2 -0.23560   1.13414 -0.208   0.8359  
## treatmentT3 -0.31613   0.95767 -0.330   0.7421  
## treatmentControl -1.09084   0.95020 -1.148   0.2540  
## du1:treatmentT2  0.06623   0.17473  0.379   0.7055  
## du1:treatmentT3  0.19904   0.13350  1.491   0.1395  
## du1:treatmentControl 0.10604   0.14326  0.740   0.4611  
## 
## Residual standard error: 2.148 on 90 degrees of freedom
## Multiple R-squared:  0.7573, Adjusted R-squared:  0.7384 
## F-statistic: 40.12 on 7 and 90 DF,  p-value: < 2.2e-16
```

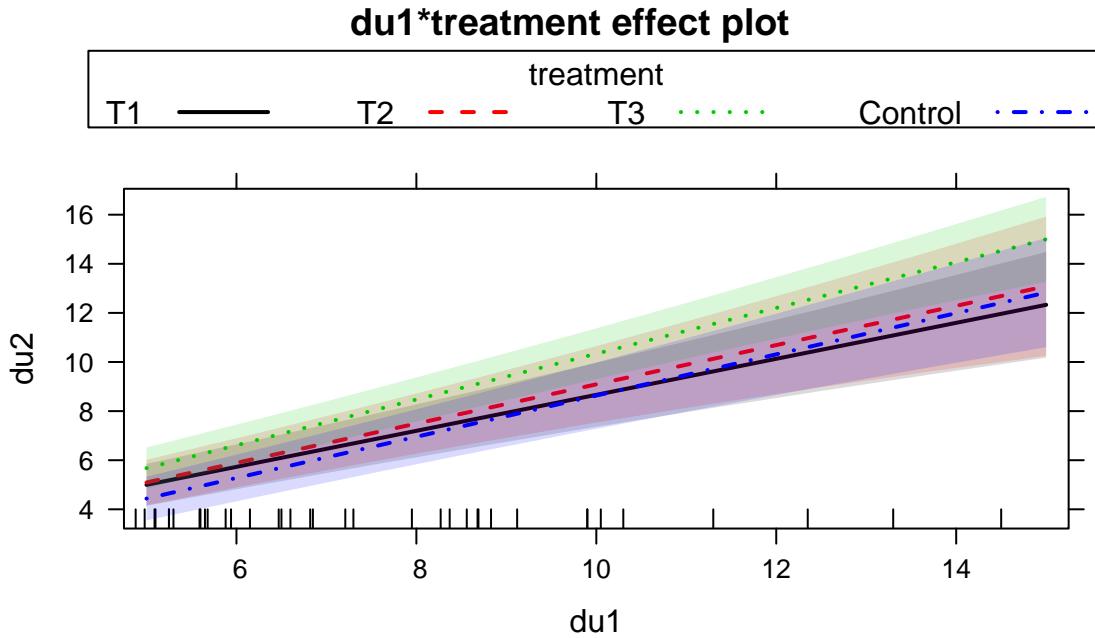


Figure 8.33: Term-plot for decibel tolerance interaction model (version 2). This plot is not printed in color because it is impossible to distinguish the four groups whether in color or black and white.

Or we can let the term-plots (Figures 8.32 and 8.33) show us all four different simplified models. Here we can see that all the slopes “look” to be pretty similar. When the interaction model is fit and the results “look” like the additive model, there is a good chance that we will be able to avoid all this complication and just use the additive model without missing anything interesting. There are two different options for displaying interaction models. Version 1 (Figure 8.32) has a different panel for each level of the categorical variable and Version 2 (Figure 8.33) puts all the lines on the same plot. In this case, neither version shows much of a difference and Version 2 overlaps so much that you can’t see all the groups.

```
plot(allEffects(head2), x.var="du1", ci.style="bands")
plot(allEffects(head2), x.var="du1", multiline=T, ci.style="bands")
```

In situations with more than 2 levels, the  $t$ -tests for the interaction or changing y-intercepts are not informative for deciding if you really need different slopes or intercepts for all the groups. They only tell you if a specific group is potentially different from the baseline group and the choice of the baseline is arbitrary. To assess whether we really need to have varying slopes or intercepts with more than two groups we need to develop  $F$ -tests for the interaction part of the model.

## 8.12 F-tests for MLR models with quantitative and categorical variables and interactions

For models with multi-category ( $J - 2$ ) categorical variables we need a method for deciding if all the extra complexity present in the additive or interaction models is necessary. We can appeal to model selection methods such as the adjusted  $R^2$  that focus on balancing model fit and complexity but interests often move to trying to decide if the differences are more extreme than we would expect by chance if there were no group differences in intercepts or slopes. Because of the multi-degree of freedom aspects of the use of indicator variables ( $J - 1$  variables for a  $J$  level categorical variable), we have to develop tests that combine and assess information across multiple “variables” – even though these indicators all pertain to a single original categorical variable. ANOVA  $F$ -tests did exactly this sort of thing in the One and Two-Way ANOVA models

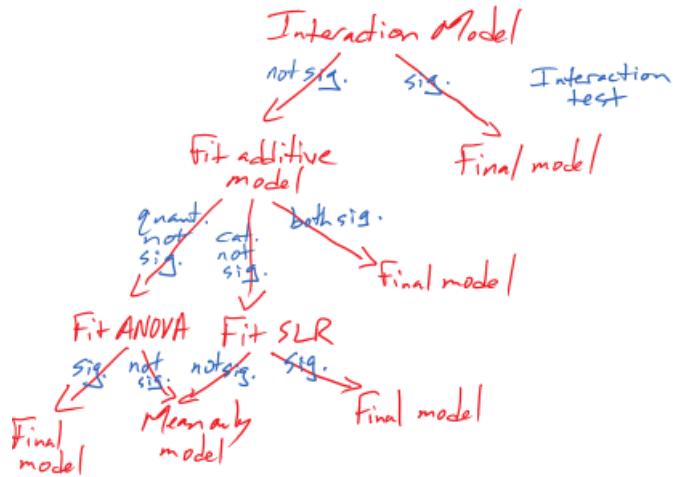


Figure 8.34: Diagram of models to consider in an interaction model.

and can do that for us here. There are two models that we perform tests in – the additive and the interaction models. We start with a discussion of the tests in an interaction setting since that provides us the **first test to consider** in most situations to assess evidence of whether the extra complexity of varying slopes is really needed. If we don't "need" the varying slopes or if the plot really does have lines for the groups that look relatively parallel, we can fit the additive model and either assess evidence of the need for different intercepts or for the quantitative predictor – either is a reasonable next step. Basically this establishes a set of **nested models** (each model is a reduced version of another more complicated model higher in the tree of models) displayed in Figure 8.34. This is based on the assumption that we would proceed through the model, dropping terms if the p-values are large ("not significant" in the diagram) to arrive at a final model.

If the initial interaction test suggests the interaction is important, then no further refinement should be considered and that model should be explored (this was the same protocol suggested in the 2-WAY ANOVA situation, the other place where we considered interactions). If the interaction is not deemed important based on the test, then the model should be re-fit using both variables in an additive model. In that additive model, both variables can be assessed conditional on the other one. If both have small p-values, then that is the final model and should be explored further. If either the categorical or quantitative variable have large p-values, then they can be dropped from the model and the model re-fit with only one variable in it, usually starting with dropping the component with the largest p-value if both are not "small". Note that if there is only a categorical variable remaining, then we would call that linear model a One-Way ANOVA (quantitative response and  $J$  group categorical explanatory) and if the only remaining variable is quantitative, then a SLR model is being fit. If that final variable has a large p-value in either model, it can be removed and all that is left to describe the responses is a mean only model. Otherwise the single variable model is the final model. Usually we will not have to delve deeply into this tree of models, but it is good to consider the potential paths that an analysis could involve before it is started.

To perform the first test (after checking that assumptions are met, of course), we can apply the `Anova` function from the `car` package to an interaction model<sup>17</sup>. It will provide three tests, one for each variable by themselves, which are not too interesting, and then the interaction test. This will result in an  $F$ -statistic that, if the assumptions are met, will follow an  $F(J - 1, n - 2J)$ -distribution under the null hypothesis. This tests the hypotheses:

- $H_0$  : The slope for  $x$  is the same for all  $J$  groups in the population vs
- $H_A$  : The slope for  $x$  in at least one group differs from the others in the population.

<sup>17</sup>We could also use the `anova` function to do this but using `Anova` throughout this material provides the answers we want in the additive model and it has no impact for the only test of interest in the interaction model since the interaction is the last component in the model.

This test is also legitimate in the case of a two-level categorical variable ( $J = 2$ ) and then follows an  $F(1, n - 4)$ -distribution under the null hypothesis. With  $J = 2$ , the p-value from this test matches the results for the  $t$ -test ( $t_{n-4}$ ) for the single slope-changing coefficient in the model summary output. The noise tolerance study, introduced in Section 8.10, provides a situation for exploring the results in detail.

With the  $J = 4$  level categorical variable (*Treatment*), the model for the second noise tolerance measurement (*du2*) as a function of the interaction between *Treatment* and initial noise tolerance (*du1*) is

$$\begin{aligned} \text{du2}_i = & \beta_0 + \beta_1 \text{du1}_i + \beta_2 I_{T2,i} + \beta_3 I_{T3,i} + \beta_4 I_{\text{Control},i} \\ & + \beta_5 I_{T2,i} \text{du1}_i + \beta_6 I_{T3,i} \text{du1}_i + \beta_7 I_{\text{Control},i} \text{du1}_i + \varepsilon_i \end{aligned}$$

We can re-write the previous hypotheses in one of two more specific ways:

- $H_0$  : The slope for *du1* is the same for all four *treatment* groups in the population OR
- $H_0 : \beta_5 = \beta_6 = \beta_7 = 0$ 
  - This defines a null hypothesis that all the deviation coefficients for getting different slopes for the different treatments are 0 in the population.
- $H_A$  : The slope for *du1* is NOT the same for all four *Treatment* groups in the population (at least one group has a different slope) OR
- $H_A$  : At least one of  $\beta_5, \beta_6, \beta_7$  is different from 0 in the population.
  - The alternative states that at least one of the deviation coefficients for getting different slopes for the different *Treatments* is not 0 in the population.

In this situation, the results for the test of these hypotheses is in the row labeled *du1:treatment* in the **Anova** output. The ANOVA table below shows a test statistic of  $F = 0.768$  with the *numerator df* of 3, coming from  $J - 1$ , and the *denominator df* of 90, coming from  $n - 2J = 98 - 2 * 4 = 90$  and also provided in the **Residuals** row in the table, leading to an  $F(3, 90)$ -distribution for the test statistic under the null hypothesis. The p-value from this distribution is 0.515, showing little to no evidence against the null hypothesis. The conclusion is that there is insufficient evidence to claim that the slope coefficient for *du1* in explaining *du2* is different for at least one of the *Treatment* groups in the population.

#### Anova(head2)

```
## Anova Table (Type II tests)
##
## Response: du2
##              Sum Sq Df  F value Pr(>F)
## du1          1197.78  1 259.5908 <2e-16
## treatment     23.90  3   1.7265 0.1672
## du1:treatment 10.63  3   0.7679 0.5150
## Residuals    415.27 90
```

Without evidence to support an interaction, we should consider both the quantitative and categorical variables in an additive model. The ANOVA table for the additive model contains two interesting tests. One test is for the quantitative variable discussed previously. The other is for the categorical variable, assessing whether different y-intercepts are needed. The additive model here is

$$\text{du2}_i = \beta_0 + \beta_1 \text{du1}_i + \beta_2 I_{T2,i} + \beta_3 I_{T3,i} + \beta_4 I_{\text{Control},i} + \varepsilon_i.$$

The hypotheses assessed in the ANOVA test for treatment are:

- $H_0$  : The y-intercept for the model with *du1* is the same for all four *Treatment* groups in the population OR
- $H_0 : \beta_2 = \beta_3 = \beta_4 = 0$

- This defines a null hypothesis that all the deviation coefficients for getting different y-intercepts for the different *Treatments* are 0 in the population.
- $H_A$  : The y-intercepts for the model with *du1* is NOT the same for all four *Treatment* groups in the population (at least one group has a different y-intercept) OR
- $H_A$  : At least one of  $\beta_2, \beta_3, \beta_4$  is different from 0 in the population.
- The alternative states that at least one of the deviation coefficients for getting different y-intercepts for the different *Treatments* is not 0 in the population.

The *F*-test for the categorical variable in an additive model follows  $F(J - 1, n - J - 1)$ -distribution under the null hypothesis. For this example, the test statistic for *Treatment* follows an  $F(3, 93)$ -distribution under the null hypothesis. The observed test statistic has a value of 1.74, generating a p-value of 0.164. So we would fail to reject the null hypothesis and conclude that there is no evidence to support the conclusion of some difference in y-intercepts between the *treatment* groups, in a model with *du1*, in the population. We could interpret this in the fashion we used initially in MLR by stating this result as: there is no evidence of a difference in the mean *du2* for the *Treatment* groups after controlling for *du1*.

```
head1 <- lm(du2~du1+treatment, data=Headache)
Anova(head1)
```

```
## Anova Table (Type II tests)
##
## Response: du2
##             Sum Sq Df  F value Pr(>F)
## du1        1197.8  1 261.5491 <2e-16
## treatment    23.9  3   1.7395 0.1643
## Residuals   425.9 93
```

In the same ANOVA table, there is a test for the *du1* effect. This tests  $H_0 : \beta_1 = 0$  vs  $H_A : \beta_1 \neq 0$  in a model with different y-intercepts estimated for the different groups. If we remove this effect from the model, all we are left with is different y-intercepts for the groups. A model just with different y-intercepts is typically called a One-Way ANOVA model. Here, there is evidence that the quantitative variable is needed in the model after controlling for the different y-intercepts for different treatments. Note that this interpretation retains the conditional wording regardless of whether the other variable had a small p-value. If you want an unconditional interpretation for a variable, then you will need to refit the model without the other variable(s) after deciding that they are not important.

## 8.13 AICs for model selection

There are a variety of techniques for selecting among a set of potential models or refining an initially fit MLR model. Hypothesis testing can be used (in the case where we have nested models) or comparisons of adjusted  $R^2$  across different potential models (it is valid for nested or non-nested model comparisons). Diagnostics should play a role in the models considered and in selecting among models that might appear to be similar on a model comparison criterion. In this section, a new model selection method is introduced that has stronger theoretical underpinnings, a slightly more interpretable scale, and, often, better performance in picking an optimal<sup>18</sup> model than the ***adjusted R***<sup>2</sup>. The measure is called the ***AIC*** (Akaike's An Information Criterion<sup>19</sup>, (Akaike, 1974)). It is extremely popular, but sometimes misused, in some fields such as Ecology and has been applied in almost every other potential application area where statistical models can be compared. Burnham

<sup>18</sup>In most situations, it would be crazy to assume that the true model for a process has been obtained so we can never pick the correct model. In fact, we won't even know if we are picking a "good" model, but just the best from a set of the candidate models. But we can study the general performance of methods using simulations where we know the true model and the AIC has some useful properties in identifying the correct model when it is in the candidate set of models. No such similar theory exists for the adjusted  $R^2$ .

<sup>19</sup>Most people now call this Akaike's (pronounced **ah-kah-ee-kay**) Information Criterion, but he used the AIC nomenclature to mean An Information Criterion – he was not so vain as to name the method after himself in the original paper that proposed it.

and Anderson (2002) have been responsible for popularizing the use of AIC for model selection, especially in Ecology. The **AIC is an estimate of the distance (or discrepancy or divergence) between a candidate model and the true model, on a log-scale**, based on a measure called the Kullback-Leibler divergence. The models that are closer (have a smaller distance) to the truth are better and we can compare how close two models are to the truth, picking the one that has a smaller distance (smaller AIC) as better. The AIC includes a component that is on the log-scale, so negative values are possible and you should not be disturbed if you are comparing large magnitude negative numbers – just pick the one with the smallest value.

The AIC is optimized (smallest) for a model that contains the optimal balance of simplicity of the model with quality of fit to the observations. Scientists are driven to different degrees by what is called the **principle of parsimony**: that **simpler explanations (models) are better if everything else is equal or even close to equal**. In this case, it would mean that if two models are similarly good on AIC, then select the simpler of the two models since it is more likely to be correct in general than the more complicated model. The AIC is calculated as  $AIC = -2\log(Likelihood) + 2m$ , where the **Likelihood** provides a measure of fit of the model (we let R calculate it for us) and gets smaller for better fitting models and  $m$  = (number of estimated  $\beta$ 's + 1). The value  $m$  is called the *model degrees of freedom* for AIC calculations and relates to how many total parameters are estimated. Note that it is a different measure of *degrees of freedom* than used in ANOVA  $F$ -tests. The main things to understand about the formula for the AIC is that as  $m$  increases, the AIC will go up and that as the fit improves, the *likelihood* will decrease (and so will the *log-likelihood*). More details of these components of the methods will be left for more advanced material – we will focus on how to use and interpret these results.

There are some other facets of this discussion to keep in mind when comparing models. More complicated models always fit better (we saw this for the  $R^2$  measure, as the proportion of variation explained always goes up if more “stuff” is put into the model even if the “stuff” isn’t useful). The AIC resembles the adjusted  $R^2$  in that it incorporates the count of the number of parameters estimated. This allows the AIC to make sure that enough extra variability is explained in the responses to justify making the model more complicated (increasing  $m$ ). The optimal model on AIC has to balance adding complexity and increasing quality of the fit. Since this measure provides an estimate of the distance or discrepancy to the “true model”, the model with the smallest value “wins” – it is top-ranked on the AIC. Note that the **top-ranked AIC model** will typically **not be the best fitting** model since the best fitting model is always the most complicated model considered. The top AIC model is the one that is estimated to be closest to the truth, where the truth is still unknown...

To help with interpreting the scale of AICs, they are often reported in a table sorted from smallest to largest values with the AIC and the “delta AIC” or, simply,  $\Delta AIC$  reported. The

$$\Delta AIC = AIC_{model} - AIC_{topModel}$$

and so provides a value of 0 for the top-ranked AIC model and a measure of how much worse on the AIC scale the other models are. A rule of thumb is that a 2 unit difference on AICs ( $\Delta AIC = 2$ ) is decent evidence of a difference in the models and more than 4 units ( $\Delta AIC > 4$ ) is a really big difference. This is more based on experience than a distinct reason or theoretical result but seems to provide reasonable results in most situations. Often researchers will consider any models within 2 AIC units of the top model ( $\Delta AIC < 2$ ) as indistinguishable on AICs and so either select the simplest model of the choices or report all the models with similar “support”, allowing the reader to explore the suite of similarly supported potential models . It is important to remember that if you search across too many models, even with the AIC to support your model comparisons, you might find a spuriously top model. Individual results that are found by exploring many tests or models have higher chances to be **spurious** and results found in this manner are difficult to **reproduce** when someone repeats a similar study<sup>20</sup>. For these reasons, there is a set of general recommendations that have been developed for using AICs:

---

<sup>20</sup>Reproducibility ideas are used in statistics first by making data and code used available to others (like all the code and data in this book) and second by trying to use methods that when others perform similar studies they will find similar results (from Physics, think of the famous cold-fusion experiments [http://en.wikipedia.org/wiki/Cold\\_fusion](http://en.wikipedia.org/wiki/Cold_fusion).

- Consider a suite of models (often pre-specified and based on prior research in the area of interest) and find the models with the top (in other words, smallest) AIC results.
  - The suite of candidate models need to contain at least some good models. Selecting the best of a set of BAD models only puts you at the top of \$%#%-mountain, which is not necessarily a good thing.
- Report a table with the models considered, sorted from smallest to largest AICs ( $\Delta\text{AICs}$  from smaller to larger) that includes a count of number of parameters estimated<sup>21</sup>, the AICs, and  $\Delta\text{AICs}$ .
- Interpret the top model or top models if a few are close on the AIC-scale to the top model.
- **DO NOT REPORT P-VALUES OR CALL TERMS “SIGNIFICANT” when selecting models using AICs.**
  - Hypothesis testing and AIC model selection are not compatible philosophies and testing in models selected by AICs invalidates the tests as they have inflated Type I error rates.
- You can describe variables as “important” or “useful” and report confidence intervals to aid in interpretation of the effects in the selected model(s) but need to avoid performing hypothesis tests.
- Remember that the selected model is not the “true” model – it is only the best model *according to AIC* among the set of models *you provided*.
- Model assumptions need to be met to use AICs. They assume that the model is specified correctly up to possibly comparing different predictor variables.

When working with AICs, there are two options. Fit the models of interest and then run the `AIC` function on each model. This can be tedious, especially when we have many possible models to consider. We can make it easy to fit all the potential candidate models that are implied by a complicated starting model by using the `dredge` function from the `MuMIn` package (Barton, 2016). The name (`dredge`) actually speaks to what ***fitting all possible models*** really engages – what is called ***data dredging***. The term is meant to refer to considering way too many models for your data set, probably finding something good from the process, but maybe identifying something spurious since you looked at so many models. Note that if you take a hypothesis testing approach where you plan to remove any terms with large p-values in this same situation, you are really considering all possible models as well because you could have removed some or all model components. Methods that consider all possible models are probably best used in exploratory analyses where you do not know if any or all effects should be important. If you have more specific research questions, then you probably should try to focus on comparisons of models that help you directly answer those questions.

The `dredge` function provides an automated method of assessing all possible simpler models based on an initial (full) model. It generates a table of AIC results,  $\Delta\text{AICs}$ , and also shows when various predictors are in or out of the model for all reduced models possible from an initial model. For quantitative predictors, the estimated slope is reported when that predictor is in the model. For categorical variables and interactions with them, it just puts a “+” in the table to let you know that the term is in the models. Note that you must run the `options(na.action = "na.fail")` code to get `dredge` to work.

To explore the AICs and compare their results to the adjusted  $R^2$  that we used before for model selection, we can relate results found in Section 8.4 and 8.1. In that situation we were considering a “full” model that included *Elevation*, *Min.Temp*, and *Max.Temp* as potential predictor variables after removing two influential points. And we considered all possible reduced models from that “full”<sup>22</sup> model. Note that the `dredge` output adds one more model that adjusted  $R^2$  can’t consider – the mean-only model that contains no predictor variables. In the following output it is the last model in the output (worst ranked on AIC). Including the mean-only model in these results helps us “prove” that there is support for having something in the model if we can clear better support for other models than this simplest possible model.

<sup>21</sup>Although often excluded, the count of parameters should include counting the residual variance as a parameter.

<sup>22</sup>We put quotes on “full” or sometimes call it “fullish” model because we could always add more to the model, like interactions or other explanatory variables. So we never have a full model but we do have our “most complicated that we are considering” model.

In reading `dredge` output as it is constructed here, the models are sorted by top to bottom AIC values (smallest AIC to largest). The column `delta` is for the  $\Delta$ AICs and shows a 0 for the first row, which is the top-ranked AIC model. Here it is for the model with *Elevation* and *Max.Temp* but not including *Min.Temp*. This was also the top ranked model from adjusted  $R^2$ , which is reproduced in the `adjRsq` column. The AIC is calculated using the previous formula based on the `df` and `logLik` columns. The `df` is the most useful column for comparing models as it helps you see how complex each model is. For example, the top model used up 4 *model df* (three  $\beta$ 's and the residual error variance) and the most complex model that included four predictor variables used up 5 *model df*.

```
require(MuMin)
options(na.action = "na.fail") #Must run this code once to use dredge
dredge(m6, rank="AIC",
       extra = c("R^2", adjRsq=function(x) summary(x)$adj.r.squared))

## Global model call: lm(formula = Snow.Depth ~ Elevation + Min.Temp + Max.Temp, data = snotel2[-c(9,
##      22), ])
## ---
## Model selection table
##   (Int)    Elv Max.Tmp Min.Temp     R^2 adjRsq df  logLik   AIC delta
## 4 -167.50 0.02408 1.2530          0.8495 0.8344  4 -80.855 169.7  0.00
## 8 -213.30 0.02686 1.2430          0.9843 0.8535  0.8304 5 -80.541 171.1  1.37
## 2 -80.41 0.01791                   0.8087 0.7996  3 -83.611 173.2  3.51
## 6 -130.70 0.02098                   1.0660 0.8134  0.7948 4 -83.322 174.6  4.93
## 5 179.60                         -5.0090 0.6283  0.6106 3 -91.249 188.5  18.79
## 7 178.60                         -0.2687 -4.6240 0.6308 0.5939 4 -91.170 190.3  20.63
## 3 119.50                         -2.1800          0.4131 0.3852  3 -96.500 199.0  29.29
## 1 40.21                           0.0000 0.0000          2 -102.630 209.3  39.55
##   weight
## 4 0.568
## 8 0.286
## 2 0.098
## 6 0.048
## 5 0.000
## 7 0.000
## 3 0.000
## 1 0.000
## Models ranked by AIC(x)
```

There are two models that are clearly favored over the others with  $\Delta$ AICs for the model with *Elevation* and *Max.Temp* of 0 and for the model with all three predictors of 1.37. The  $\Delta$ AIC for the third ranked model suggesting clear support for the top model over this. The difference between the second and third ranked models also provides relatively strong support for the more complex model over the model with just *Elevation*. And the mean-only model had a  $\Delta$ AIC of nearly 40 – suggesting extremely strong support for the top model versus using no predictors. So we have pretty clear support for models that include the *Elevation* and *Max.Temp* variables (in both top models) and some support for also including the *Min.Temp*, but the top model did not require its inclusion.

We could add further explorations of the term-plots and confidence intervals for the slopes from the top or, here, possibly top two models. We would not spend any time with p-values since we already used the AIC to assess importance of the model components and they are invalid if we model select prior to reporting them. We can quickly compare the slopes for variables that are shared in the two models since they are both quantitative variables using the output. It is interesting that the *Elevation* and *Max.Temp* slopes change little with the inclusion of *Min.Temp* in moving from the top to second ranked model (0.02408 to 0.0286 and 1.253 to 1.243).

This was an observational study and so we can't consider causal inferences here as discussed previously. Generally, the use of AICs does not preclude making causal statements but if you have randomized assignment of levels of an explanatory variable, it is more philosophically consistent to use hypothesis testing methods in

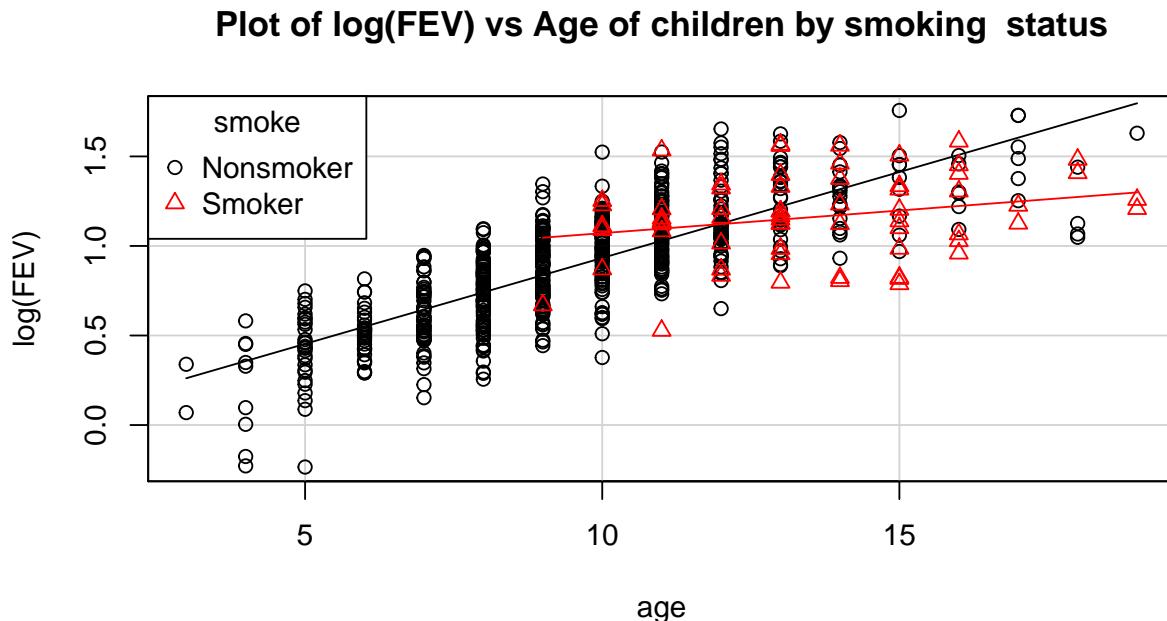


Figure 8.35: Scatterplot of  $\log(\text{FEV})$  vs Age by smoking status.

that setting. If you went to the effort to impose the levels of a treatment on the subjects, it also makes sense to see if the differences created are beyond what you might expect by chance if the treatment didn't matter.

## 8.14 Forced Expiratory Volume model selection using AICs

Researchers were interested in studying the effects of smoking by children on their lung development by measuring the forced expiratory volume (*FEV*, measured in Liters) in a representative sample of children ( $n = 654$ ) between the ages of 3 and 19; this data set is available in the *FEV* data set in the *coneproj* package (Meyer and Liao (2016), Liao and Meyer (2014)). Measurements on the *age* (in years) and *height* (in inches) as well as the *sex* and *smoking status* of the children were made. We would expect both the *age* and *height* to have positive relationships with *FEV* (lung capacity) and that smoking might decrease the lung capacity but also that older children would be more likely to smoke. So the *height* and *age* might be *confounded* with smoking status and smoking might diminish lung development for older kids – resulting in a potential interaction between *age* and *smoking*. The *sex* of the child might also matter and should be considered or at least controlled for since the response is a size-based measure. This creates the potential for including up to four variables (*age*, \* *height*, *sex*, and *smoking status*) and possibly the interaction between *age*\* and *smoking status*. Initial explorations suggested that modeling the  $\log(\text{FEV})$  would be more successful than trying to model the responses on the original scale. Figure 8.35 shows the suggestion of different slopes for the smokers than non-smokers and that there aren't very many smokers under 9 years old.

So we will start with a model that contains an *age* by *smoking* interaction and include *height* and *sex* as additive terms. We are not sure if any of these model components will be needed, so the simplest candidate model will be to remove all the predictors and just have a mean-only model (*FEV~1*). In between the mean-only and most complicated model are many different options where we can drop the interaction or drop the additive terms or drop the terms involved in the interaction if we don't need the interaction.

```
require(coneproj)
```

```
data(FEV)
```

```
FEV$sex <- factor(FEV$sex)
levels(FEV$sex) <- c("Female", "Male")
FEV$smoke <- factor(FEV$smoke)
levels(FEV$smoke) <- c("Nonsmoker", "Smoker")
require(car)
scatterplot(log(FEV)~age|smoke, data=FEV, smooth=F,
           main="Plot of log(FEV) vs Age of children by smoking status",
           legend.coords="topleft")
```

To get the needed results, start with the ***full model*** – the most complicated model you want to consider. It is good to check assumptions before considering reducing the model as they rarely get better in simpler models and the **AIC is only appropriate to use if the model assumptions are reasonably well-met**. As noted above, our full model for the  $\log(\text{FEV})$  values is specified as ‘ $\log(\text{FEV}) \sim \text{height} + \text{age} * \text{smoke} + \text{sex}$ ’.

```
fm1 <- lm(log(FEV)~height+age*smoke+sex, data=FEV)
summary(fm1)
```

```
##
## Call:
## lm(formula = log(FEV) ~ height + age * smoke + sex, data = FEV)
##
## Residuals:
##      Min        1Q    Median        3Q       Max
## -0.62926 -0.08783  0.01136  0.09658  0.40751
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)
## (Intercept) -1.919494  0.080571 -23.824 < 2e-16
## height       0.042066  0.001759  23.911 < 2e-16
## age          0.025368  0.003642   6.966 8.03e-12
## smokeSmoker  0.107884  0.113646   0.949  0.34282
## sexMale      0.030871  0.011764   2.624  0.00889
## age:smokeSmoker -0.011666  0.008465  -1.378  0.16863
##
## Residual standard error: 0.1454 on 648 degrees of freedom
## Multiple R-squared:  0.8112, Adjusted R-squared:  0.8097
## F-statistic: 556.8 on 5 and 648 DF, p-value: < 2.2e-16
par(mfrow=c(2,2), oma=c(0,0,2,0))
plot(fm1, sub.caption="Diagnostics for full FEV model")
```

There are a few outlying points noted in Figure 8.36 but they are not influential and the normality and constant variance assumptions are reasonably well met. If we select a different model(s), we would want to check its diagnostics and make sure that the results do not look noticeably worse than these do.

The AIC function can be used to generate the AIC values for a single or set of candidate models. It will also provide the model degrees of freedom used for each model. For example, suppose that we want to compare `fm1` to a model without the interaction term in the model, called `fm1R`. You need to fit both models and then apply the AIC function to them with commas between the model names:

```
fm1R <- lm(log(FEV)~height+age+smoke+sex, data=FEV)
AIC(fm1, fm1R)
```

```
##      df      AIC
## fm1    7 -658.5178
## fm1R   6 -658.6037
```

These results tell us that the `fm1R` model (the one without the interaction) is better on the AIC. Note that this model does not “fit” as well as the other model, it is just the top AIC model – the AIC results suggest that it is slightly closer to the truth than the more complicated model. But this provides only an assessment

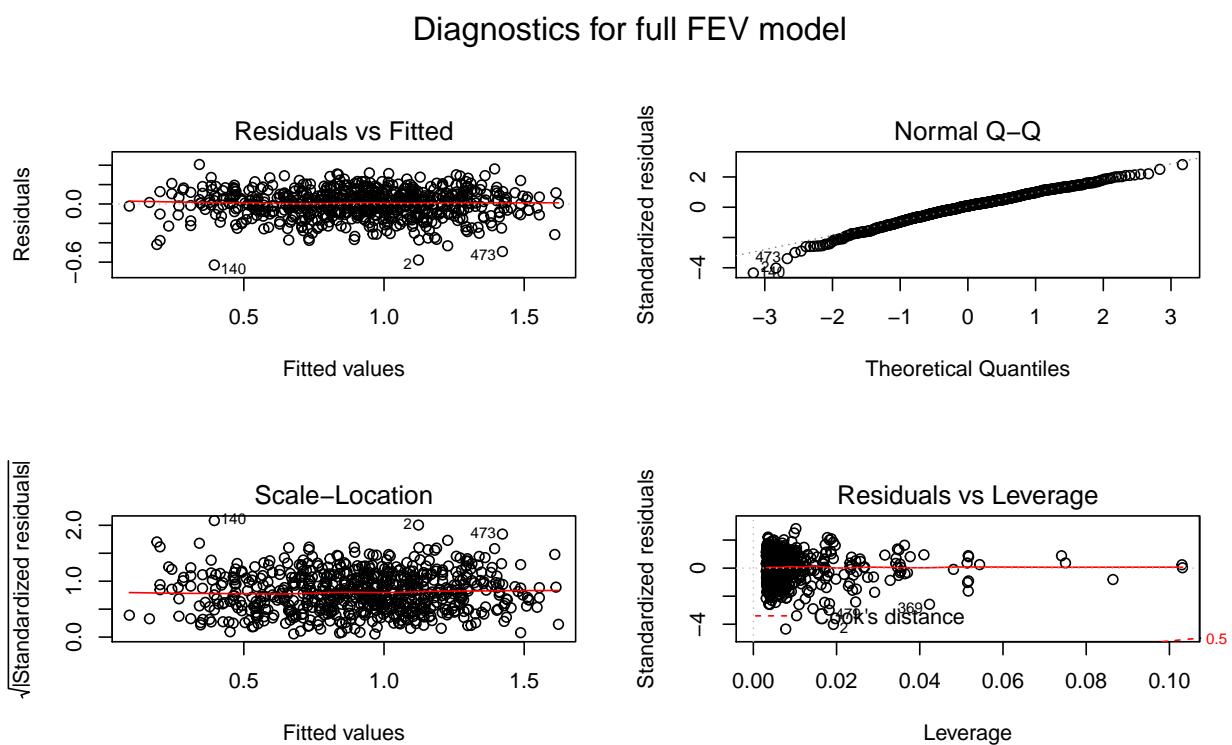


Figure 8.36: Diagnostics for the  $\log(\text{FEV})$  model that includes height, sex, and an interaction between age and smoking status (the full model).

of the difference between including or excluding the interaction between *age* and *smoking* in a model with two other predictors. We are probably also interested in whether the other terms are needed in the model. The full suite of results from dredge provide model comparisons that help us to assess the presence/absence of each model component including the interaction.

```
require(MuMin)
options(na.action = "na.fail") #Must run this code once to use dredge
dredge(fm1, rank="AIC",
       extra = c("R^2", adjRsq=function(x) summary(x)$adj.r.squared))
```

```
## Global model call: lm(formula = log(FEV) ~ height + age * smoke + sex, data = FEV)
## ---
## Model selection table
##   (Int)    age    hgh sex smk age:smk      R^2  adjRsq df  logLik
## 16 -1.944000 0.02339 0.04280  +  +     0.81060 0.80950 6 335.302
## 32 -1.919000 0.02537 0.04207  +  +     + 0.81120 0.80970 7 336.259
## 8  -1.940000 0.02120 0.04299  +           0.80920 0.80830 5 332.865
## 12 -1.974000 0.02231 0.04371          +     0.80880 0.80790 5 332.163
## 28 -1.955000 0.02388 0.04315          +     + 0.80920 0.80800 6 332.802
## 4   -1.971000 0.01982 0.04399          +           0.80710 0.80650 4 329.262
## 7   -2.265000          0.05185  +           0.79640 0.79580 4 311.594
## 3   -2.271000          0.05212          + 0.79560 0.79530 3 310.322
## 15  -2.267000          0.05190  +  +     0.79640 0.79550 5 311.602
## 11  -2.277000          0.05222  +           0.79560 0.79500 4 310.378
## 30  -0.067780 0.09493          +  +     + 0.64460 0.64240 6 129.430
## 26  -0.026590 0.09596          +           + 0.62360 0.62190 5 110.667
## 14  -0.015820 0.08963          +  +           0.62110 0.61930 5 108.465
## 6   0.004991 0.08660          +           0.61750 0.61630 4 105.363
## 10  0.022940 0.09077          +           0.60120 0.60000 4  91.790
## 2   0.050600 0.08708          +           0.59580 0.59520 3  87.342
## 13  0.822000          +  +           0.09535 0.09257 4 -176.092
## 9   0.888400          +           0.05975 0.05831 3 -188.712
## 5   0.857400          +           0.02878 0.02729 3 -199.310
## 1   0.915400          +           0.00000 0.00000 2 -208.859
##   AIC delta weight
## 16 -658.6  0.00  0.414
## 32 -658.5  0.09  0.397
## 8  -655.7  2.87  0.099
## 12 -654.3  4.28  0.049
## 28 -653.6  5.00  0.034
## 4   -650.5  8.08  0.007
## 7  -615.2  43.42 0.000
## 3  -614.6  43.96 0.000
## 15 -613.2  45.40 0.000
## 11 -612.8  45.85 0.000
## 30 -246.9  411.74 0.000
## 26 -211.3  447.27 0.000
## 14 -206.9  451.67 0.000
## 6   -202.7  455.88 0.000
## 10 -175.6  483.02 0.000
## 2  -168.7  489.92 0.000
## 13  360.2  1018.79 0.000
## 9   383.4  1042.03 0.000
## 5   404.6  1063.22 0.000
## 1   421.7  1080.32 0.000
## Models ranked by AIC(x)
```

There is a lot of information in the output and some of the needed information in the second set of rows, so we will try to point out some useful features to consider. The left columns describe the models being

estimated. For example, the first row of results is for a model with an intercept (**Int**), *age* (**age**) , *height* (**hgh**), *sex* (**sex**), and *smoking*(**smk**). For *sex* and *smoking*, there are “+”s in the output row when they are included in that model but no coefficient since they are categorical variables. There is no interaction between *age* and *smoking* in the top ranked model. The top AIC model has an  $R^2 = 0.8106$ , adjusted  $R^2$  of 0.8095, *model df*=6 (from an intercept, four slopes, and the residual variance), log-likelihood (**logLik**)=335.302, an AIC=-658.6 and  $\Delta\text{AIC}$  of 0.00. The next best model adds the interaction between *age* and *smoking*, resulting in increases in the  $R^2$ , adjusted  $R^2$ , and *model df*, but increasing the AIC by 0.09 units ( $\Delta\text{AIC} = 0.09$ ). This suggests that these two models are essentially equivalent on the AIC because the difference is so small. The simpler model is a little bit better on AIC so you could focus on it or on the slightly more complicated model – but you should probably note that the evidence is equivocal for these two models.

The comparison to other potential models shows the strength of evidence in support of all the other model components. The intercept-only model is again the last in the list with the least support on AICs with a  $\Delta\text{AIC}$  of 1080.32, suggesting it is not worth considering in comparison with the top model. **Comparing the mean-only model to our favorite model on AICs is a bit like the overall F-test we considered in Section 8.7 because it compares a model with no predictors to a complicated model.** Each model with just one predictor included is available in the table as well, with the top single predictor model based on *height* having a  $\Delta\text{AIC}$  of 43.96. So we certainly need to pursue something more complicated than SLR based on the AIC results. Closer to the top model is the third-ranked model that includes *age*, *height*, and *sex*. It has a  $\Delta\text{AIC}$  of 2.87 so we would say that these results present only marginal support for this model compared to the two top models. It is the simplest model of the top three but not close enough to be considered in detail.

This table also provides the opportunity to compare the model selection results from the adjusted  $R^2$  compared the AIC. The AIC favors the model without an interaction between *age* and *smoking* whereas the adjusted  $R^2$  favors the most complicated model considered here that included an *age* and *smoking* interaction. The AIC provides units that are more interpretable than adjusted  $R^2$  even though the scale for the AIC is a bit mysterious as **distances from the unknown true model**.

The top AIC model (and possibly the other similar models) can then be explored in more detail. You should not then focus on hypothesis testing in this model<sup>23</sup>. Confidence intervals and term-plots are useful for describing the different model components and making inferences for the estimated sizes of differences in the population. These results should not be used for deciding if effects are “significant” when the models are selected using measures like the AIC or adjusted  $R^2$ . But you can discuss how there was support or evidence to include them in the model based on AIC comparisons.

In this situation, the top model is estimated to be

$$\log(\widehat{\text{FEV}})_i = -1.94 + 0.043\text{Height}_i + 0.0234\text{Age}_i - 0.046I_{\text{Smoker},i} + 0.0293I_{\text{Male},i}$$

based on the estimated coefficients provided below. Using these results and the term-plots (Figure 8.37) we see that there are positive slopes for *Age* and *Height* on *log-FEV*, a negative coefficient for *smoking* (*Smoker*), and a positive coefficient for *sex* (*Males*), all after controlling for those other aspects of the model. We could go further with interpretations such as for the *age* term: For a 1 year increase in *age*, we expect, on average, a 0.0234 log-liter increase in *FEV*, after controlling for the *height*, *smoking status*, and *sex* of the children. We can even interpret this on the original response model using the same techniques as in Section 7.6. If we exponentiate the slope coefficient of the quantitative variable,  $\exp(0.0234) = 1.0237$ . This provides the interpretation on the original *FEV* scale, for a 1 year increase in *age*, we expect 2.4% increase in the median *FEV*, after controlling for the *height*, *smoking status*, and *sex* of the children. **The only difference from Section 7.6 when working with a log(y) model now is that we have to note that the model used to generate the slope coefficient had other components and so this estimate is after adjusting for them.**

<sup>23</sup>Hypothesis testing so permeates the use of statistics that even after using AICs many researchers are pressured to report p-values for model components. Some of this could be confusion caused when people first learned these statistical methods because when we teach you statistics we show you how to use various methods, one after another, and forget to mention that you should not use **every** method we taught you in **every** analysis.

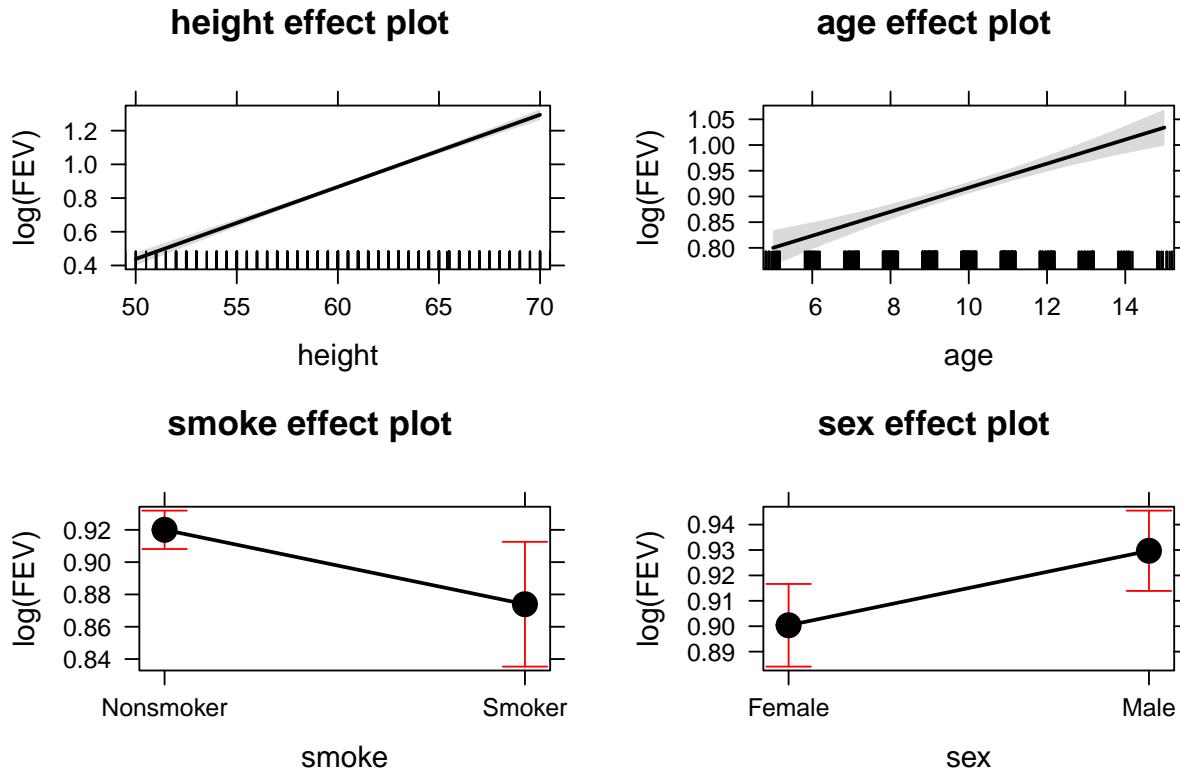


Figure 8.37: Term-plots for the top AIC model for  $\log(\text{FEV})$  that includes height, age, smoking status and sex in the model.

```
fm1R$coefficients
## (Intercept)      height         age smokeSmoker    sexMale
## -1.94399818   0.04279579   0.02338721 -0.04606754   0.02931936
confint(fm1R)
##                   2.5 %       97.5 %
## (Intercept) -2.098414941 -1.789581413
## height        0.039498923  0.046092655
## age          0.016812109  0.029962319
## smokeSmoker -0.087127344 -0.005007728
## sexMale       0.006308481  0.052330236
plot(allEffects(fm1R))
```

Like any statistical method, the AIC works better with larger sample sizes and when the assumptions are met. It also will detect important variables in models more easily when the effects of the predictor variables are strong. Along with the AIC results, it is good to report the coefficients for your top estimated model(s), confidence intervals for the coefficients and/or term-plots, and  $R^2$ . This provides a useful summary of the reasons for selecting the model(s), information on the importance of the terms within the model, and a measure of the variability explained by the model. The  $R^2$  is not used to select the model, but after selection can be a nice summary of model quality. For `fm1R`, the  $R^2 = 0.8106$  suggesting that the selected model explains 81% of the variation in  $\log\text{-FEV}$  values.

The AICs are a preferred modeling strategy in some areas such as Ecology. As with this and many other methods discussed in this book, it is sometimes as easy to find journal articles with mistakes in using statistical methods as it is to find papers doing it correctly. After completing this material, you have the potential to

have the knowledge and experience of two statistics classes and now are better trained than some researchers that frequently use these methods. This set of tools can be easily mis-applied. Try to make sure that you are thinking carefully through your problem before jumping to the statistical results. Make a graph first, think carefully about your study design and variables collected and what your models of interest might be, what assumptions might be violated based on the data collection story, and then start fitting models. Then check your assumptions and only proceed on with any inference if those conditions are reasonably well-met. The AIC provides an alternative method for selecting among different potential models and they do not need to be nested (a requirement of hypothesis testing methods used to sequentially simplify models). The automated consideration of all possible models in the `dredge` function should not be considered in all situations but can be useful in a preliminary model exploration study where no clear knowledge exists about useful models to consider.

## 8.15 Chapter summary

This chapter explored the most complicated models we're going to explore. MLR models can incorporate features of SLR and ANOVAs. The MLR's used in this chapter highlight the flexibility of the linear modeling framework to move from two-sample mean models to multi-predictor models with interactions of categorical and quantitative variables. It is useful to use the pertinent names for the simpler models, but at this point we could have called everything we are doing *fitting linear models*. The power of the linear model involves being able to add multiple predictor variables to the model and handle categorical predictors using indicator variables. All this power comes with some responsibility in that you need to know what you are trying to fit and how to interpret the results provided. We introduced each scenario working from simple to the most complicated version of the models, trying to motivate when you would encounter them, and the specific details of the interpretations of each type of model. In Chapter 9, case studies are used to review the different methods discussed with reminders of how to identify and interpret the particular methods used.

When you have to make modeling decisions, you should remember the main priorities in modeling. First, you need to find a model that can address research questions of interest. Second, find a model that is trustworthy and has assumptions that are reasonably well met. Third, report the logic and evidence that was used to identify and support the model. All too often, researchers present only a final model with little information on how they arrived at it. You should be reporting the reasons for decisions made and the evidence supporting them. For example, if you were considering an interaction model and the interaction was dropped and an additive model is re-fit and interpreted, the evidence related to the interaction test should still be reported. Similarly, if a larger MLR is considered and some variables are removed, the evidence (reason) for those removals should be provided. Because of multicollinearity in models, you should never remove more than one quantitative predictor at a time or else you could remove two variables that are important but were "hiding" when both were included in the model.

## 8.16 Important R code

There is very little "new" R code in this chapter since all these methods were either used in the ANOVA or SLR chapters. The models are more complicated but are built off of methods from previous chapters. In this code, `y` is a response variable, `x1, x2, ..., xK` are quantitative explanatory variables, `group` is a categorical variable and the data are in `DATASETNAME`.

- `scatterplot(y~x1|group, data=DATASETNAME, smooth=F)`
  - Requires the `car` package.
  - Provides a scatterplot with a regression line for each group.
- `MODELNAME <- lm(y~ x1+x2+...+xK, data=DATASETNAME)`
  - Estimates an MLR model using least squares with  $K$  quantitative predictors.

- `MODELNAME <- lm(y~ x1*group, data=DATASETNAME)`
  - Estimates an interaction model between a quantitative and categorical variable, providing different slopes and intercepts for each group.
- `MODELNAME <- lm(y~ x1+group, data=DATASETNAME)`
  - Estimates an additive model with a quantitative and categorical variable, providing different intercepts for each group.
- `summary(MODELNAME)`
  - Provides parameter estimates, overall  $F$ -test,  $R^2$ , and adjusted  $R^2$ .
- `par(mfrow=c(2, 2)); plot(MODELNAME)`
  - Provides four regression diagnostic plots in one plot.
- `confint(MODELNAME, level=0.95)`
  - Provides 95% confidence intervals for the regression model coefficients.
  - Change `level` if you want other confidence levels.
- `plot(allEffects(MODELNAME))`
  - Requires the `effects` package.
  - Provides a plot of the estimated regression lines with 95% confidence interval for the mean.
- `predict(MODELNAME, se.fit=T)`
  - Provides fitted values for all observed  $x$ 's with SEs for the mean.
- `predict(MODELNAME, newdata=data.frame(x1 = X1_NEW, x2 = X2_NEW, ..., xK = XK_NEW, interval="confidence"))`
  - Provides fitted value for specific values of the quantitative predictors with CI for the mean.
- `predict(MODELNAME, newdata=data.frame(x1 = X1_NEW, x2 = X2_NEW, ..., xK = XK_NEW, interval="prediction"))`
  - Provides fitted value for specific values of the quantitative predictors with PI for a new observation.
- `Anova(MODELNAME)`
  - Requires the `car` package.
  - Use to generate ANOVA tables and  $F$ -tests useful when categorical variables are included in either the additive or interaction models.
- `AIC(MODELNAME_1, MODELNAME_2)`
  - Use to get AIC results for two candidate models called `MODELNAME_1` and `MODELNAME_2`.
- `options(na.action = "na.fail")`  
`dredge(FULL_MODELNAME, rank="AIC", extra=c("R^2", adjRsq=function(x) summary(x)$adj.r.squared))`
  - Requires the `MuMIn` package.
  - Provides AIC and delta AIC results for all possible simpler models given a full model called `FULL_MODELNAME`.

## 8.17 Practice problems

The original research goal for the treadmill data set used for practice problems in the last two chapters was to replace the costly treadmill oxygen test with a cheap to find running time measurement but there were actually quite a few variables measured while the run time was found – maybe we can replace the treadmill test result with a combined prediction built using a few variables using the MLR techniques. The following code will get us re-started in this situation.

```
treadmill <- read.csv("http://www.math.montana.edu/courses/s217/documents/treadmill.csv",
                      header=T)
tm1 <- lm(TreadMill0x~RunTime, data=treadmill)
```

8.1. Fit the MLR that also includes the running pulse (`RunPulse`), the resting pulse (`RestPulse`), body weight (`BodyWeight`), and Age (`Age`) of the subjects using the following code. Report and interpret the  $R^2$  for this model.

8.2. Compare  $R^2$  and the adjusted  $R^2$  to the results for the SLR model that just had `RunTime` in the model. What do these results suggest?

8.3. Interpret the estimated `RunTime` slope coefficients from the SLR model and this MLR model. Explain the differences in the estimates.

8.4. Find the VIFs for this model and discuss whether there is an issue with multicollinearity noted in these results.

8.5. Report the value for the overall  $F$ -test for the MLR model and interpret the result.

8.6. Drop the variable with the largest p-value in the MLR model and re-fit it. Compare the resulting  $R^2$  and adjusted  $R^2$  values to the others found previously.

8.7. Use the `dredge` function as follows to consider some other potential reduced models and report the top two models according to adjusted  $R^2$  values. What model had the highest  $R^2$ ? Also discuss and compare the model selection results provided by the delta AICs here.

```
require(MuMIn)
options(na.action = "na.fail") #Must run this code once to use dredge
dredge(mlr1, rank="AIC",
       extra=c("R^2", adjRsq=function(x) summary(x)$adj.r.squared))
```

8.8. For one of the models, interpret the Age slope coefficient. Remember that only male subjects between 38 and 57 participated in this study. Discuss how this might have impacted the results found as compared to a more general population that could have been sampled from.

8.9. The following code creates a new three-level variable grouping the ages into low, middle, and high for those observed. The scatterplot lets you explore whether the relationship between treadmill oxygen and run time might differ across the age groups.

```
treadmill$Ageb <- cut(treadmill$Age, breaks=c(37,44.5,50.5,58))
summary(treadmill$Ageb)
require(car)
scatterplot(TreadMill0x~RunTime|Ageb, data=treadmill, smooth=F, lwd=2)
```

Based on the plot, do the lines look approximately parallel or not?

8.10. Fit the MLR that contains a `RunTime` by `Ageb` interaction – do not include any other variables. Compare the  $R^2$  and adjusted  $R^2$  results to previous models.

8.11. Find and report the results for the  $F$ -test that assesses evidence relative to the need for different slope coefficients.

8.12. Write out the overall estimated model. What level was R using as baseline? Write out the simplified model for two of the age levels.

8.13. Fit the additive model with `RunTime` and predict the mean treadmill oxygen values for subjects with run times of 11 minutes in each of the three `Ageb` groups.

8.14. Find the *F*-test results for the binned age variable in the additive model. Report and interpret those results.

# Chapter 9

## Case studies

### 9.1 Overview of material covered

At the beginning of the text, we provided a schematic of methods that you would learn about that was (probably) gibberish. Hopefully, revisiting that same diagram (Figure 9.1 will bring back memories of each of the chapters. Categorical variables create special challenges whether they are explanatory or response variables.

Every scenario with a quantitative response variable was handled using linear models. The last material on multiple linear regression modeling tied back to the One and Two-Way ANOVA models as categorical variables were added to the models. As both a review and to emphasize the connections, let's connect some of the different versions of the general linear model that we considered.

If we start with the One-Way ANOVA, the referenced-coded model was written out as:

$$y_{ij} = \alpha + \tau_j + \varepsilon_{ij}.$$

We didn't want to introduce indicator variables at that early stage of the material, but we can now write out the same model using our indicator variable approach from Chapter 8 for a  $J$ -level categorical explanatory variable using  $J - 1$  indicator variables as:

$$y_i = \beta_0 + \beta_1 I_{\text{Level } 2,i} + \beta_2 I_{\text{Level } 3,i} + \cdots + \beta_{J-1} I_{\text{Level } J,i} + \varepsilon_i.$$

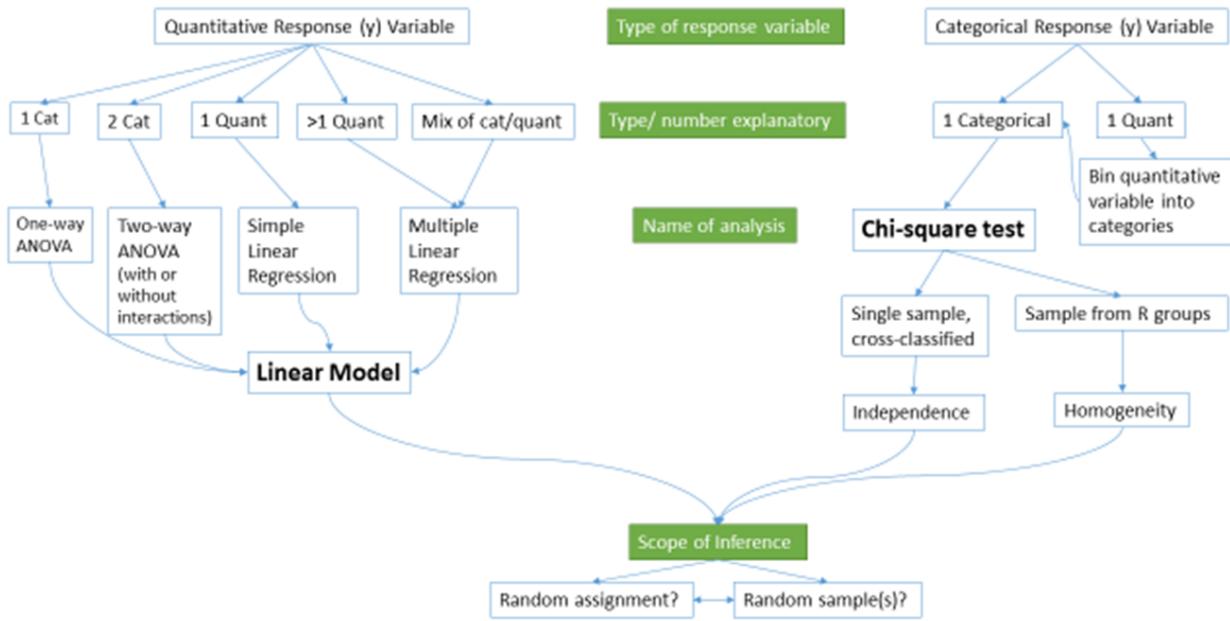


Figure 9.1: Schematic of methods covered.

We now know how the indicator variables are either 0 or 1 for each observation and only one takes in the value 1 (is “turned on”) at a time for each response. We can then equate the general notation from Chapter 8 with our specific One-Way ANOVA (Chapter 3) notation as follows:

- $\alpha = \beta_0$ :
  - The mean for the baseline category was modeled using  $\alpha$  which is the intercept term in the output that we called  $\beta_0$  in the regression models.
- For category  $j$ :
  - From the One-Way ANOVA model:
    - $\alpha + \tau_j$
  - From the regression model where the only indicator variable that is 1 is  $I_{\text{Level } j,i}$ :
    - $\beta_0 + \beta_1 I_{\text{Level } 2,i} + \beta_2 I_{\text{Level } 3,i} + \cdots + \beta_J I_{\text{Level } J,i} = \beta_0 + \beta_{j-1} * 1$
    - $= \beta_0 + \beta_{j-1}$
  - So with intercepts being equal,  $\beta_{j-1} = \tau_j$ .

The ANOVA reference-coding notation was used to focus on the coefficients that were “turned on” and their interpretation without getting bogged down in the full power (and notation) of general linear models. The same equivalence is possible to equate our work in the Two-Way ANOVA interaction model,

$$y_{ijk} = \alpha + \tau_j + \gamma_k + \omega_{jk} + \varepsilon_{ijk},$$

with the regression notation from the MLR model with an interaction:

$$\begin{aligned} y_i = & \beta_0 + \beta_1 x_i + \beta_2 I_{\text{Level } 2,i} + \beta_3 I_{\text{Level } 3,i} + \cdots + \beta_J I_{\text{Level } J,i} + \beta_{J+1} I_{\text{Level } 2,i} x_i \\ & + \beta_{J+2} I_{\text{Level } 3,i} x_i + \cdots + \beta_{2J-1} I_{\text{Level } J,i} x_i + \varepsilon_i \end{aligned}$$

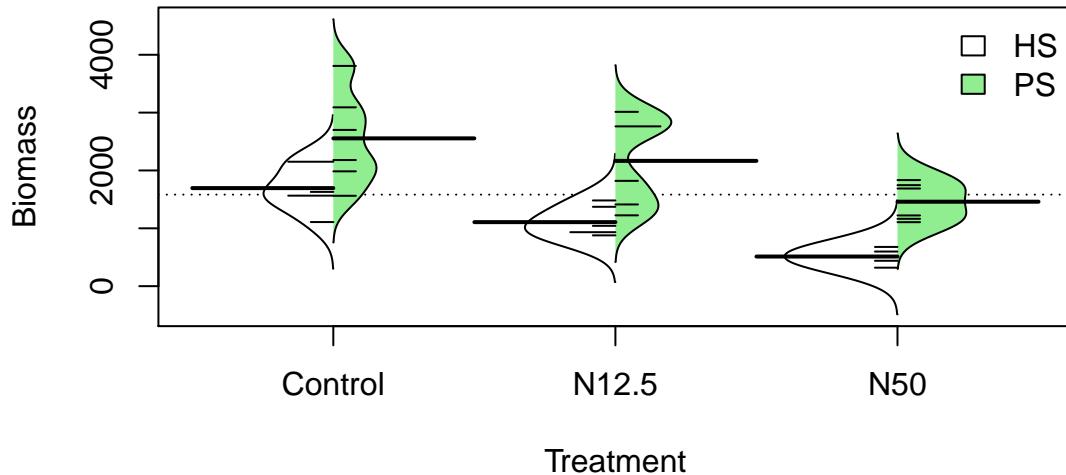


Figure 9.2: Beanplot of biomass responses by treatment and species.

If one of the categorical variables only had two levels, then we could simply replace  $x_i$  with the pertinent indicator variable and be able to equate the two versions of the notation. That said, we won't attempt that here. And if both variables have more than 2 levels, the number of coefficients to keep track of grows rapidly. The great increase in complexity of notation to fully writing out the indicator variables in the regression approach with interactions with two categorical variables is the other reason we explored the Two-Way ANOVA using a "simplified" notation system even though `lm` used the indicator approach to fit the model. The Two-Way ANOVA notation helped us distinguish which coefficients related to main effects and the interaction, something that the regression notation doesn't make clear.

In the following three sections, you will have one more chance to see applications of the methods considered here. The data sets are taken directly from published research articles, so you can see the potential utility of the methods we've been discussing for handling real problems. They are focused on biological applications because the particular journal (*Biology Letters*) that all of these were drawn from encourages authors to share their data sets, making our re-analyses possible. Use these sections to review or to re-inforce methods from earlier in the book.

## 9.2 The impact of simulated chronic nitrogen deposition on the biomass and N<sub>2</sub>-fixation activity of two boreal feather moss–cyanobacteria associations

In a 16-year experiment, Gundale and Nordin (2013) studied the impacts of Nitrogen (N) additions on the mass of two feather moss species (*Pleurozium schreberi* (PS) and *Hylocomium* (HS)) in the Svartberget Experimental Forest in Sweden. They used a randomized block design: here this means that within each of 6 blocks (pre-specified areas that were divided into three experimental units or plots of area 0.1 hectare), one of the three treatments were randomly applied. **Randomized block designs** involve randomization of levels within blocks or groups as opposed to **completely randomized designs** where each **experimental unit** (the subject or plot that will be measured) could be randomly assigned to any treatment. This is done

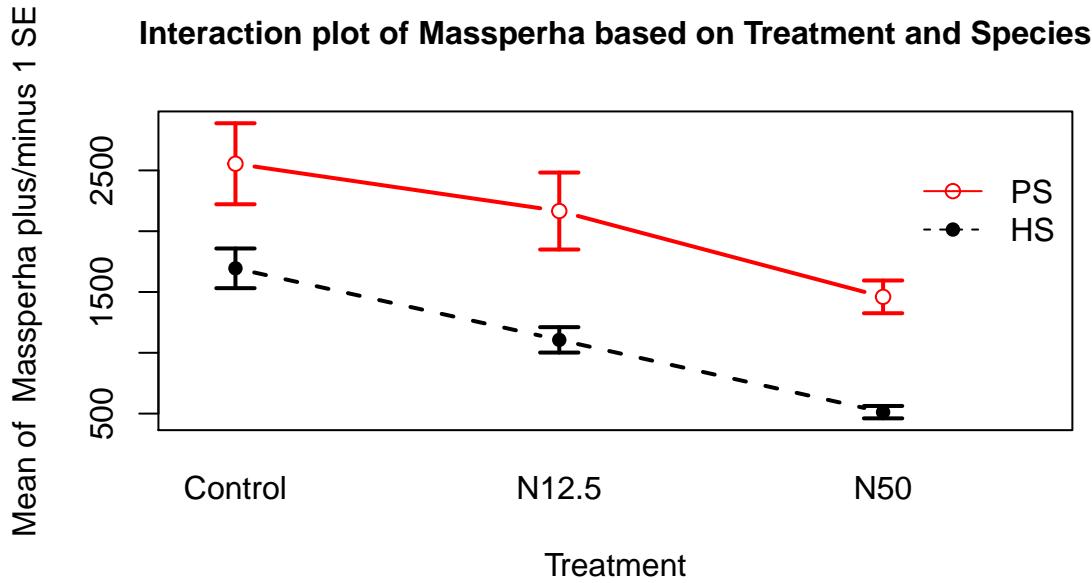


Figure 9.3: Interaction plot of biomass responses by treatment and species.

in agricultural studies to control for systematic differences across the fields by making sure each treatment level is used in each area or **block** of the field.

The three treatments involved different levels of N applied immediately after snow melt, *Control* (no additional N – just the naturally deposited amount),  $12.5 \text{ kg N ha}^{-1}\text{yr}^{-1}$  (*N12.5*), and  $50 \text{ kg N ha}^{-1}\text{yr}^{-1}$  (*N50*). The researchers were interested in whether the treatments would have differential impacts on the two species of moss growth. They measured a variety of other variables, but here we focus on the estimated *biomass* per hectare (mg/ha) of the *species* (*PS* or *HS*), both measured for each plot within each block, considering differences across the *treatments* (*Control*, *N12.5*, or *N50*). The beanplot in Figure 9.2 provides some initial information about the responses. Initially there seem to be some differences in the combinations of groups and some differences in variability in the different groups, especially with much more variability in the *control* treatment level and more variability in the *PS* responses than for the *HS* responses.

```
gdn <- read.csv("http://www.math.montana.edu/courses/s217/documents/gundalebachnordin_2.csv")
require(beanplot)
beanplot(Massperha~Species*Treatment, data=gdn, side = "b",
         col=list("white","lightgreen"), xlab="Treatment", ylab="Biomass")
legend("topright", bty="n", c("HS", "PS"), fill=c("white","lightgreen"))
```

The Two-WAY ANOVA model that contains a *species* by *treatment* interaction is of interest (so this has a quantitative response variable of *biomass* and two categorical predictors of *species* and *treatment*)<sup>1</sup>. We can make an interaction plot to focus on the observed patterns of the means across the combinations of levels as provided in Figure 9.3. The interaction plot suggests a relatively additive pattern of differences between *PS* and *HS* across the three treatment levels. However, the variability seems to be quite different based on this plot as well.

```
source("http://www.math.montana.edu/courses/s217/documents/intplot.R")
intplot(Massperha~Species*Treatment, data=gdn, col=c(1,2), lwd=2)
```

Based on the initial plots, we are going to be concerned about the equal variance assumption initially. We

<sup>1</sup>The researchers did not do this analysis so never directly addressed this research question although they did discuss it in general ways.

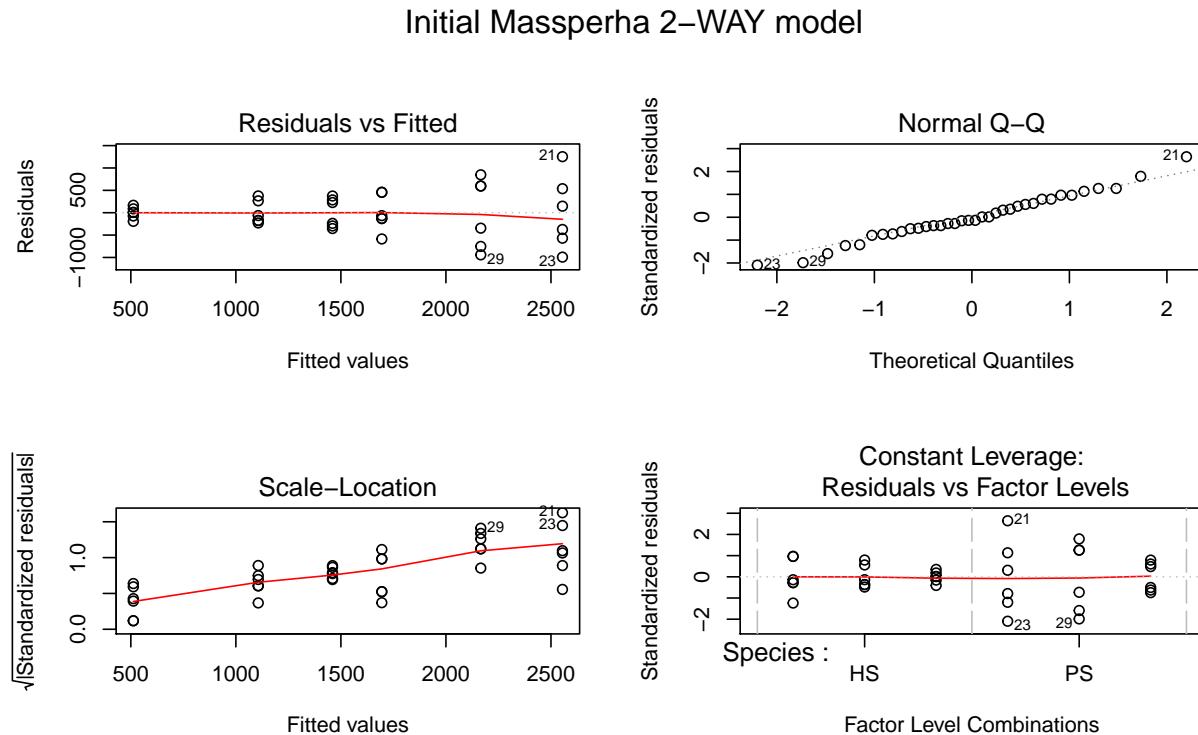


Figure 9.4: Diagnostic plots of treatment by species interaction model for Biomass.

can fit the interaction model and explore the diagnostic plots to verify that we have a problem.

```
m1 <- lm(Massperha ~ Species * Treatment, data=gdn)
summary(m1)
```

```
##
## Call:
## lm(formula = Massperha ~ Species * Treatment, data = gdn)
##
## Residuals:
##    Min     1Q Median     3Q    Max 
## -992.6 -252.2 -64.6  308.0 1252.9 
## 
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) 1694.80    211.86   8.000 6.27e-09  
## SpeciesPS    859.88    299.62   2.870 0.00745   
## TreatmentN12.5 -588.26    299.62  -1.963 0.05893   
## TreatmentN50 -1182.91    299.62  -3.948 0.00044   
## SpeciesPS:TreatmentN12.5 199.42    423.72   0.471 0.64130  
## SpeciesPS:TreatmentN50     88.29    423.72   0.208 0.83636  
## 
## Residual standard error: 519 on 30 degrees of freedom
## Multiple R-squared:  0.6661, Adjusted R-squared:  0.6104 
## F-statistic: 11.97 on 5 and 30 DF,  p-value: 2.009e-06
par(mfrow=c(2,2), oma=c(0,0,2,0))
plot(m1, sub.caption="Initial Massperha 2-WAY model")
```

There is a clear problem with non-constant variance showing up in a fanning shape<sup>2</sup> in the Residuals versus Fitted and Scale-Location plots in Figure 9.4. Interestingly, the normality assumption is not an issue so hopefully we will not worsen this result by using a transformation to try to address the non-constant variance issue. The independence assumption is violated in two ways for this model by this study design – the blocks create clusters or groups of observations and the block should be accounted for (they did this in their models by adding *block* as a categorical variable to their models). Using blocked designs and accounting for the blocks in the model will typically give more precise inferences for the effects of interest, the treatments randomized within the blocks. Additionally, **there are two measurements on each plot** within block, one for *SP* and one for *HS* and these might be related (for example, high *HS* biomass might be associated with high *SP*) so putting both observations into a model **violates the independence assumption** at a second level. It takes more advanced statistical models (called linear mixed models) to see how to fully deal with this, for now it is important to recognize the issues. The more complicated models provide similar results here and include the *treatment* by *species* interaction we are going to explore, they just add to this basic model to account for these other issues.

Remember that **before using a *log*-transformation, you always must check that the responses are strictly greater than 0::**

```
summary(gdn$Massperha)
```

```
##      Min. 1st Qu. Median Mean 3rd Qu. Max.
##    319.1 1015.1 1521.8 1582.3 2026.6 3807.6
```

The minimum is 319.1 so it is safe to apply the natural log-transformation to the response variable (*Biomass*) and repeat the previous plots:

```
gdn$logMassperha <- log(gdn$Massperha)
par(mfrow=c(1,2))
beanplot(logMassperha~Species*Treatment, data=gdn, side = "b",
          col= list("white","lightgreen"), xlab="Treatment",
          ylab="log-Biomass", main="(a)")
legend("topright", bty="n", c("HS", "PS"), fill=c("white","lightgreen"))
intplot(logMassperha~Species*Treatment, data=gdn, col=c(1,2), lwd=2, main="(b)")
```

The variability in the beanplot in Figure 9.5(a) appears to be more consistent across the groups but the lines appear to be a little less parallel in the interaction plot Figure 9.5(b) for the log-scale response. That is not problematic but suggests that we may now have an interaction present – it is hard to tell visually sometimes. Again, fitting the interaction model and exploring the diagnostics is the best way to assess the success of the transformation applied.

The log(Mass per ha) version of the response variable has little issue with changing variability present in the residuals in Figure 9.6 with much more similar variation in the residuals across the fitted values.

The normality assumption is leaning towards a slight violation with too little variability in the right tail and so maybe a little bit of a left skew. This is only a minor issue and fixes the other big issue (clear non-constant variance), so this model is at least closer to giving us trustworthy inferences than the original model. The model presents some evidence of a *Species* by *Treatment* interaction ( $F(2, 30) = 4.2$ , p-value= 0.026). This suggests that the effects on the log-biomass of the treatments differ between the two species. The mean log-biomass is lower for *HS* than *PS* with the mean log-biomass to decrease more rapidly than for *PS*. In other words, increasing nitrogen has more of an impact on the resulting log-biomass for *HS* than for *PS*. The highest mean log-biomass rates were observed under the control conditions for both species making nitrogen appear to inhibit growth of these species.

```
m2 <- lm(logMassperha~Species*Treatment, data=gdn)
summary(m2)
```

```
##
## Call:
```

---

<sup>2</sup>Instructors in this class often get asked what a problem with non-constant variance actually looks like – this is it!

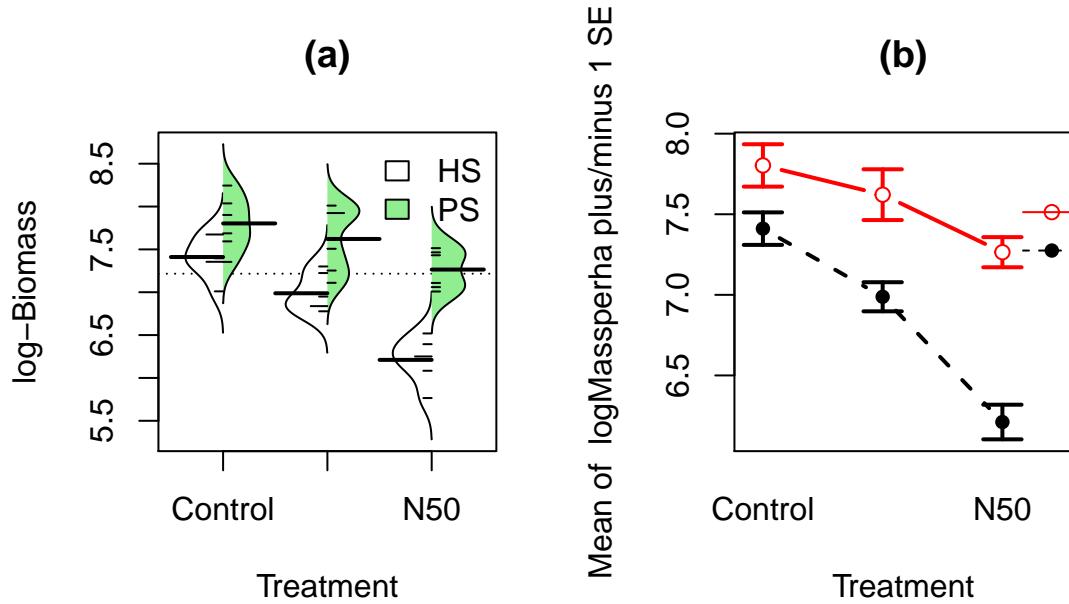


Figure 9.5: Beanplot and interaction plot of the log-Biomass responses by treatment and species.

```

## lm(formula = logMassperha ~ Species * Treatment, data = gdn)
##
## Residuals:
##    Min      1Q   Median      3Q     Max 
## -0.51138 -0.16821 -0.02663  0.23925  0.44190
## 
## Coefficients:
##                               Estimate Std. Error t value Pr(>|t|)    
## (Intercept)                 7.4108    0.1160  63.902 < 2e-16  
## SpeciesPS                  0.3921    0.1640   2.391  0.02329  
## TreatmentN12.5              -0.4228    0.1640  -2.578  0.01510  
## TreatmentN50                -1.1999    0.1640  -7.316 3.79e-08  
## SpeciesPS:TreatmentN12.5    0.2413    0.2319   1.040  0.30645  
## SpeciesPS:TreatmentN50      0.6616    0.2319   2.853  0.00778  
## 
## Residual standard error: 0.2841 on 30 degrees of freedom
## Multiple R-squared:  0.7998, Adjusted R-squared:  0.7664 
## F-statistic: 23.96 on 5 and 30 DF,  p-value: 1.204e-09
require(car)
Anova(m2)

## Anova Table (Type II tests)
## 
## Response: logMassperha
##             Sum Sq Df F value    Pr(>F)    
## Species       4.3233  1 53.577 3.755e-08  
## Treatment     4.6725  2 28.952 9.923e-08  
## Species:Treatment 0.6727  2  4.168  0.02528  
## Residuals     2.4208 30
par(mfrow=c(2,2), oma=c(0,0,2,0))
plot(m2, sub.caption="log-Massperha 2-WAY model")

```

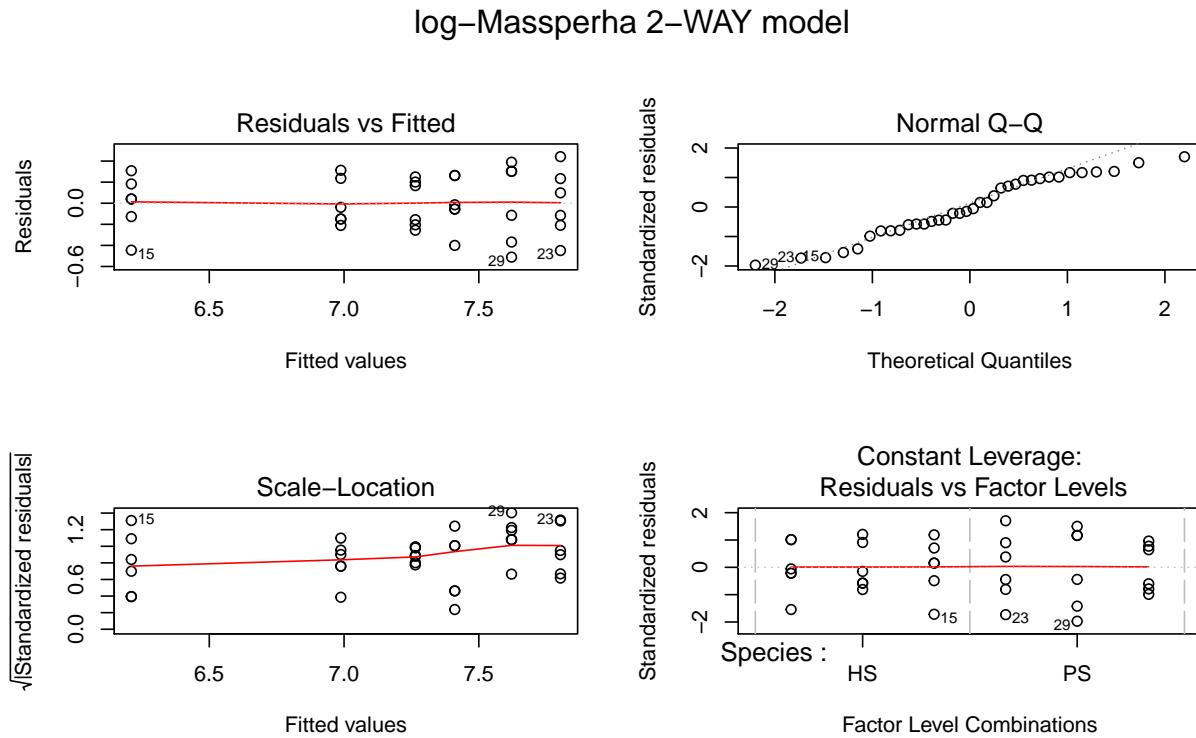


Figure 9.6: Diagnostic plots of treatment by species interaction model for log-Biomass.

The researchers actually applied a  $\log(y + 1)$  transformation to all the variables. This was used because one of their many variables had a value of 0 and so they added 1 to avoid analyzing a  $-\infty$  response. This was not needed for most of their variables because most did not attain the value of 0. Adding a small value to observations and then log-transforming is a common but completely arbitrary practice and the choice of the added value can impact the results. Sometimes considering a square-root transformation can accomplish similar benefits as the log-transform and be applied safely to responses that include 0s. Or more complicated statistical models can be used that allow 0s in responses and still account for the violations of the linear model assumptions – see a statistician or continue exploring more advanced statistical methods for ideas in this direction.

The term-plot in Figure 9.7 provides another display of the results with some information on the results for each combination of the species and treatments. Finding evidence that the treatments caused different results for the different species is a good first start. And it appears that there are some clear differences among certain combinations such as the mean for *PS-Control* is clearly larger than for *HS-N50*. The researchers were probably really interested in whether the *N12.5* results differed from *Control* for *HS* and whether the *species* differed at *Control* sites. As part of performing all pair-wise comparisons, we can assess those sorts of detailed questions. This sort of follow-up could be considered in any Two-Way ANOVA model but will be most interesting in situations where there are important interactions.

```
require(effects)
plot(allEffects(m2), multiline=T, ci.style="bars")
```

#### Follow-up Pairwise Comparisons:

Given strong evidence of an interaction, many researchers would like more details about the source of the differences. We can re-fit the model with a unique mean for each combination of the two predictor variables, fitting a One-Way ANOVA model (here with six levels) and using Tukey's HSD to provide safe inferences for differences among pairs of the true means. There are six groups corresponding to all combinations of

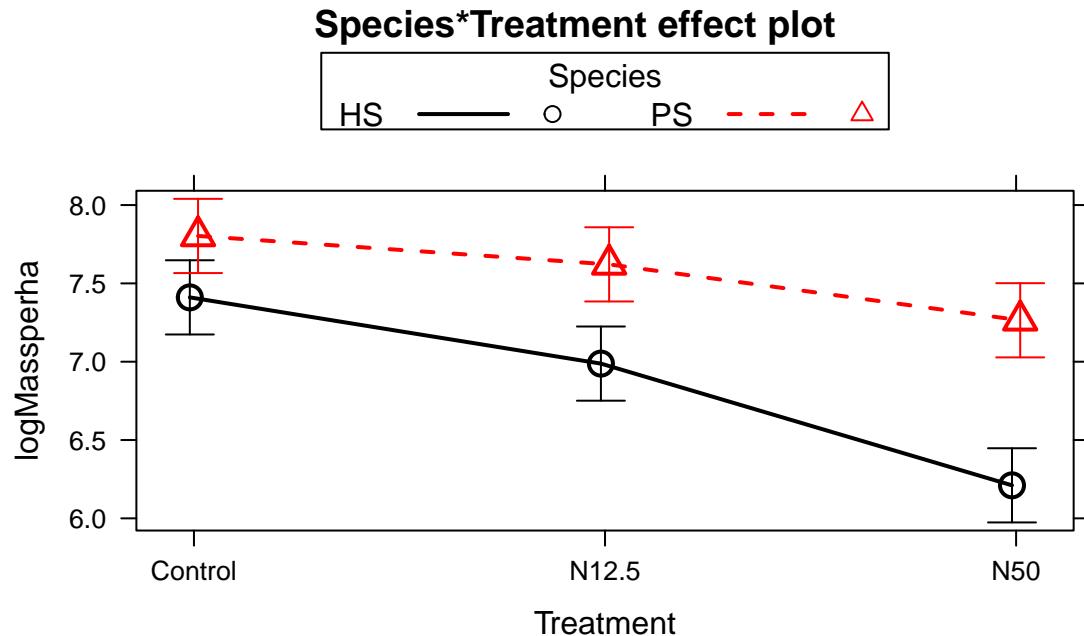


Figure 9.7: Term-plot of the interaction model for log-biomass.

*Species* (HS, PS) and treatment levels (Control, N12.5, and N50) provided in the new variable SpTrt by the interaction function with new levels of HS.Control, PS.Control, HS.N12.5, PS.N12.5, HS.N50, and PS.N50. The One-Way ANOVA F-test ( $F(5, 30) = 23.96$ , p-value < 0.0001) suggests that there is evidence of some difference in the true mean log-biomass among the six treatment combinations. Note that the One-Way ANOVA table contains the test for at least one of those means being different from the others; the interaction test above was testing a more refined hypothesis – does the effect of treatment differ between the two species? With a small p-value from the overall One-Way ANOVA test, the pair-wise comparisons should be of interest.

#Create new variable:

```
gdn$SpTrt <- interaction(gdn$Species, gdn$Treatment)
levels(gdn$SpTrt)
```

```
## [1] "HS.Control" "PS.Control" "HS.N12.5"   "PS.N12.5"   "HS.N50"
## [6] "PS.N50"
newm2 <- lm(logMassperha~SpTrt, data=gdn)
Anova(newm2)
```

```
## Anova Table (Type II tests)
##
## Response: logMassperha
##           Sum Sq Df F value    Pr(>F)
## SpTrt      9.6685  5 23.963 1.204e-09
## Residuals  2.4208 30
require(multcomp)
PWnewm2 <- glht(newm2, linfct=mcp(SpTrt="Tukey"))
confint(PWnewm2)
```

```
##
##   Simultaneous Confidence Intervals
##
##   Multiple Comparisons of Means: Tukey Contrasts
##
```

```

## 
## Fit: lm(formula = logMassperha ~ SpTrt, data = gdn)
## 
## Quantile = 3.043
## 95% family-wise confidence level
## 
## 
## Linear Hypotheses:
##                               Estimate lwr      upr
## PS.Control - HS.Control == 0  0.39210 -0.10698  0.89118
## HS.N12.5 - HS.Control == 0   -0.42277 -0.92185  0.07630
## PS.N12.5 - HS.Control == 0   0.21064 -0.28843  0.70972
## HS.N50 - HS.Control == 0    -1.19994 -1.69901 -0.70086
## PS.N50 - HS.Control == 0    -0.14620 -0.64528  0.35288
## HS.N12.5 - PS.Control == 0  -0.81487 -1.31395 -0.31580
## PS.N12.5 - PS.Control == 0  -0.18146 -0.68053  0.31762
## HS.N50 - PS.Control == 0    -1.59204 -2.09112 -1.09296
## PS.N50 - PS.Control == 0    -0.53830 -1.03738 -0.03923
## PS.N12.5 - HS.N12.5 == 0    0.63342  0.13434  1.13249
## HS.N50 - HS.N12.5 == 0     -0.77717 -1.27624 -0.27809
## PS.N50 - HS.N12.5 == 0     0.27657 -0.22250  0.77565
## HS.N50 - PS.N12.5 == 0    -1.41058 -1.90966 -0.91151
## PS.N50 - PS.N12.5 == 0    -0.35685 -0.85592  0.14223
## PS.N50 - HS.N50 == 0       1.05374  0.55466  1.55281

```

We can also generate the Compact Letter Display (CLD) to help us group up the results.

```
cld(PWnewm2)
```

```

## HS.Control PS.Control  HS.N12.5   PS.N12.5      HS.N50      PS.N50
##      "bd"        "d"      "b"      "cd"      "a"      "bc"

```

We can also add it to an interaction plot to create Figure 9.8. Researchers often use displays like this to simplify the presentation of pair-wise comparisons. Sometimes researchers add bars or stars to provide the same information about pairs that are or are not detectably different. The following code creates the plot of these results using our `intplot` function and the `cld=T` option.

```

intplot(logMassperha~Species*Treatment, cld=T, cldshift=0.15,
        cldcol=c(2,3,4,5,6,8), data=gdn, lwd=2,
        main="Interaction with CLD from Tukey's HSD on One-Way ANOVA")

```

These results suggest that *HS-N50* is detectably different from all the other groups (letter “a”). The rest of the story is more complicated since many of the sets contain overlapping groups in terms of detectable differences. Some specific aspects of those results are most interesting. The mean log-biomasses were not detectably different between the (they share a “d”). In other words, without treatment, there is no evidence of a difference in how much of the two species are present in the sites. For *N12.5* and *N50* treatments, there are detectable differences between the *Species*. These comparisons are probably of the most interest initially and suggest that the treatments have a different impact on the two species, remembering that in the control treatments, the results for the two species were not detectably different. Further explorations of the sizes of the differences that can be extracted from selected confidence intervals in the Tukey’s HSD results printed above.

### 9.3 Ants learn to rely on more informative attributes during decision-making

In Sasaki and Pratt (2013), a set of ant colonies were randomly assigned to one of two treatments to study whether the ants could be “trained” to have a preference for or against certain attributes for potential nest sites. The colonies were either randomly assigned to experience the repeated choice of two identical colony

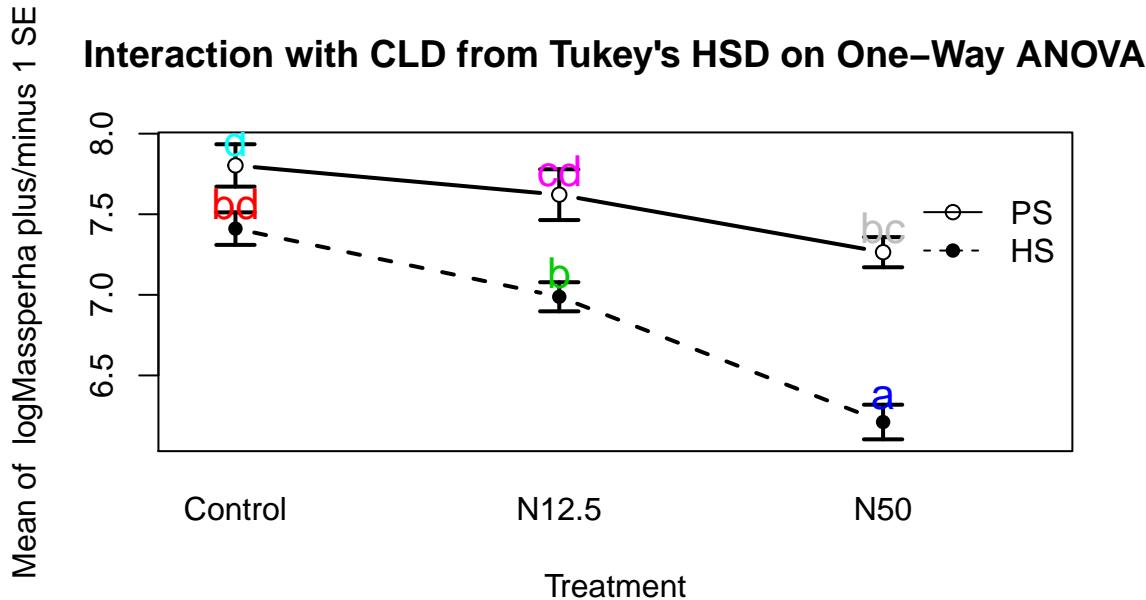


Figure 9.8: Interaction plot for log-biomass with CLD from Tukey's HSD for all pairwise comparisons.

sites except for having an inferior light or entrance size attribute. Then the ants were allowed to choose between two nests, one that had a large entrance but was dark and the other that had a small entrance but was bright. 54 of the 60 colonies that were randomly assigned to one of the two treatments completed the experiment by making a choice between the two types of sites. The data set and some processing code follows.

The first question of interest is what type of analysis is appropriate here. Once we recognize that there are two categorical variables being considered (*Treatment* group with two levels and *After* treatment choice with two levels *SmallBright* or *LargeDark*), then this is recognized as being within our Chi-square testing framework. The random assignment of colonies (the subjects here) to treatment levels tells us that the ***Chi-square Homogeneity test*** is appropriate here and that we can make causal statements if we find evidence of differences in the patterns of responses.

```
sasakipratt <- read.csv("http://www.math.montana.edu/courses/s217/documents/sasakipratt.csv")
sasakipratt$group <- factor(sasakipratt$group)
levels(sasakipratt$group) <- c("Light", "Entrance")
sasakipratt$after <- factor(sasakipratt$after)
levels(sasakipratt$after) <- c("SmallBright", "LargeDark")
sasakipratt$before <- factor(sasakipratt$before)
levels(sasakipratt$before) <- c("SmallBright", "LargeDark")
plot(after~group, data=sasakipratt)
```

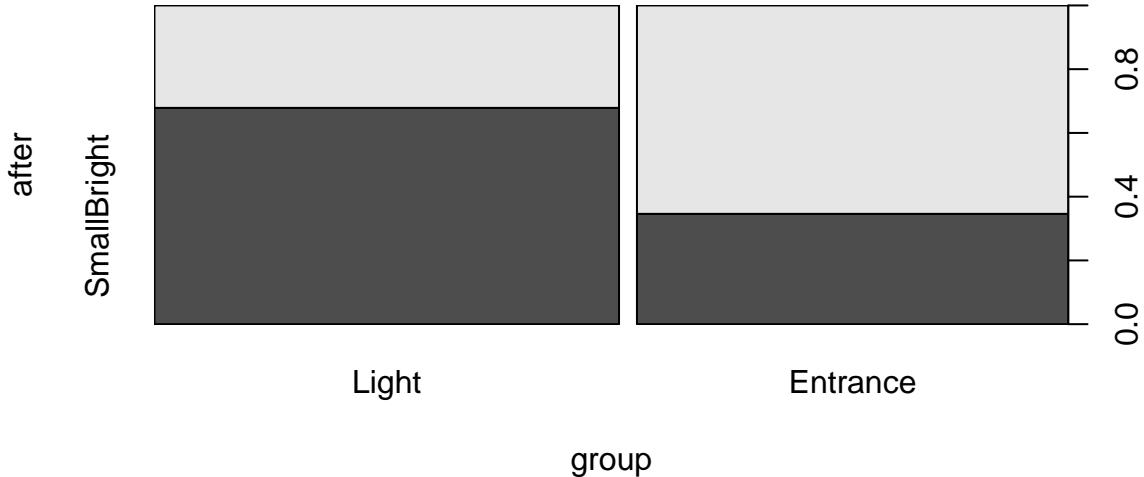


Figure 9.9: Stacked bar chart for Ant Colony results.

```

require(mosaic)
tally(~group+after, data=sasakipratt)

##           after
## group      SmallBright LargeDark
##   Light          19       9
##   Entrance        9      17
table1 <- tally(~group+after, data=sasakipratt, margins=F)

```

The null hypothesis of interest here is that there is no difference in the distribution of responses on *After* – the rates of their choice of den types – between the two treatment *groups* in the population of all ant colonies like those studied. The alternative is that there is some difference in the distributions of *After* between the *groups* in the population.

To use the Chi-square distribution to find a p-value for the  $X^2$  statistic, we need all the expected cell counts to be larger than 5, so we should check that. Note that in the following, the `correct=F` option is used to keep the function from slightly modifying the statistic used that occurs when overall sample sizes are small.

```
chisq.test(table1, correct=F)$expected
```

```

##           after
## group      SmallBright LargeDark
##   Light      14.51852  13.48148
##   Entrance    13.48148  12.51852

```

Our expected cell count condition is met, so we can proceed to explore the results of the test:

```
chisq.test(table1, correct=F)
```

```

##
## Pearson's Chi-squared test
##
## data: table1
## X-squared = 5.9671, df = 1, p-value = 0.01458

```

## 9.4. MULTI-VARIATE MODELS ARE ESSENTIAL FOR UNDERSTANDING VERTEBRATE DIVERSIFICATION IN DEEP TIME

The  $X^2$  statistic is 5.97 which, if our assumptions are met, should approximately follow a Chi-squared distribution with  $(R - 1) * (C - 1) = 1$  degrees of freedom under the null hypothesis. The p-value is 0.015, suggesting that there is good evidence against the null hypothesis. We can conclude that there is a difference in the distribution of the responses between the two treated groups in the population of all ant colonies that could have been treated. Because of the random assignment, we can say that the treatments caused differences in the colony choices. These results cannot be extended to ants beyond those being studied by these researchers because they were not randomly selected.

Further exploration of the standardized residuals can provide more insights in some situations, although here they are similar for all the cells:

```
chisq.test(table1, correct=F)$residuals
```

```
##           after
## group      SmallBright LargeDark
##   Light      1.176144 -1.220542
##   Entrance   -1.220542  1.266616
```

When all the standardized residual contributions are similar, that suggests that there are differences in all the cells from what we would expect if the null hypothesis were true. Basically, that means that what we observed is a bit larger than expected for the *Light* treatment group in the *SmallBright* choice and lower than expected in *LargeDark* – those treated ants preferred the small and bright den. And for the *Entrance* treated group, they preferred the large entrance, dark den at a higher rate than expected if the null is true and lower than expected in the small entrance, bright location.

The researchers extended this basic result a little further using a statistical model called *logistic regression*, which involves using something like a linear model but with a categorical response variable (well – it actually only works for a two-category response variable). They also had measured which of the two types of dens that each colony chose before treatment and controlled for that choice. So the actual model used in their paper contained two predictor variables – the randomized treatment received that we explored here and the prior choice of den type. The interpretation of their results related to the same treatment effect, but they were able to discuss it after adjusting for the colonies previous selection. Their conclusions were similar to those found with our simpler analysis. Logistic regression models are a special case of what are called generalized linear models and are a topic for the next level of statistics if you continue exploring.

## 9.4 Multi-variate models are essential for understanding vertebrate diversification in deep time

Benson and Mannion (2012) published a paleontology study that considered modeling the diversity of *Sauropodomorphs* across  $n = 26$  “stage-level” time bins. Diversity is measured by the count of the number of different species that have been found in a particular level of fossils. Specifically, the counts in the *Sauropodomorphs* group were obtained for a stage between *Carnian* and *Maastrichtian*, with the first three stages in the *Triassic*, the next ten in the *Jurassic*, and the last eleven concerned about variation in sampling efforts and the ability of paleontologists to find fossils across different stages creating a false impression of the changes in biodiversity over time. They first wanted to see if the species counts were related to factors such as the count of dinosaur-bearing-formations (*DBF*) and the count of dinosaur-bearing-collections (*DBC*) that have been identified for each period. The thought is that if there are more formations or collections of fossils from certain stages, the diversity might be better counted (more found of those available to find) and those stages with less information available might be under-counted. They also measured the length of each stage (*Duration*) but did not consider it in their models since they want to reflect the diversity and longer stages would likely have higher diversity.

Their main goal was to develop a model that would **control for** the effects of sampling efforts and allow them to perform inferences for whether the diversity was different between the *Triassic/Jurassic* (grouped together) and considered models that included two different versions of sampling effort variables and one

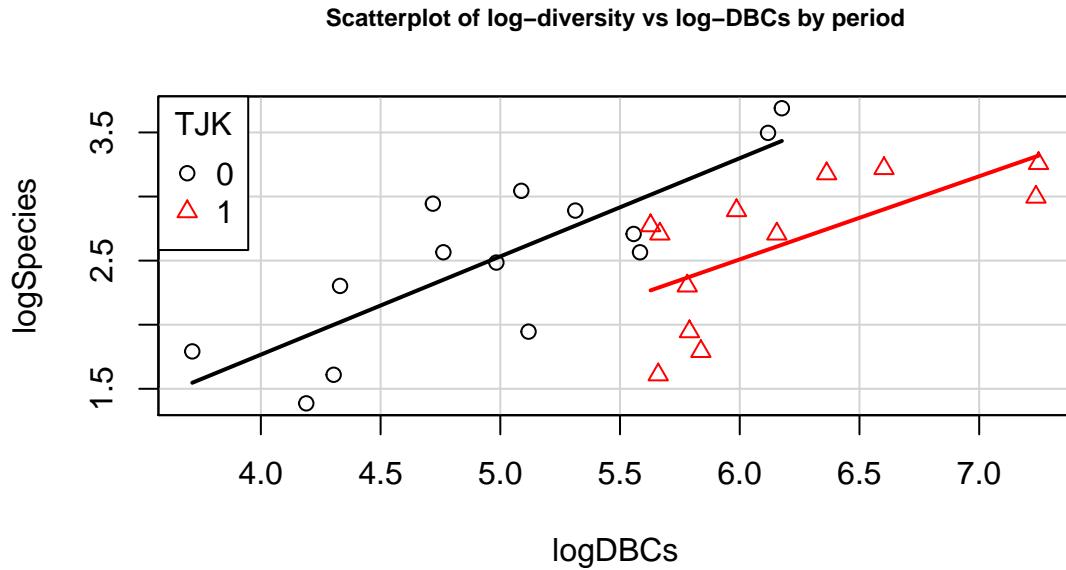


Figure 9.10: Scatterplot of log-biodiversity vs log-DBC<sub>s</sub> by TJK.

for the comparisons of periods (an indicator variable *TJK*=0 if the observation is in *Triassic* or *Jurassic* or 1 if in *Cretaceous*). They *log-e* transformed all their quantitative variables because the untransformed variables created diagnostic issues including influential points. They explored a model just based on the *DBC* predictor<sup>3</sup> and they analyzed the residuals from that model to see if the biodiversity was different in the *Cretaceous* or before, finding a “p-value  $\geq 0.0001$ ” (I think they meant  $<0.0001$ <sup>4</sup>). They were comparing the MLR models you learned to some extended regression models that incorporated a correction for correlation in the responses over time, but we can proceed with fitting some of their MLR models and using an AIC comparison similar to what they used. There are some obvious flaws in their analysis and results that we will avoid<sup>5</sup>.

First, we start with a plot of the log-diversity vs the log-dinosaur bearing collections by period. We can see fairly strong positive relationships between the log amounts of collections and species found with potentially similar slopes for the two periods but what look like different intercepts. Especially for *TJK* level 1 (*Cretaceous* period) observations, we might need to worry about a curving relationship. Note that a similar plot can also be made using the formations version of the quantitative predictor variable and that the research questions involve whether *DBF* or *DBC* are better predictor variables.

```
bm <- read.csv("http://www.math.montana.edu/courses/s217/documents/bensonmanion.csv")
bm2 <- bm[,-c(9:10)]
require(psych)
pairs.panels(bm2, ellipses=F)
bm$logSpecies <- log(bm$Species)
```

<sup>3</sup>This was not even close to their top AIC model so they made an odd choice.

<sup>4</sup>I had students read this paper in a class and one decided that this was a reasonable way to report small p-values – it is WRONG. We are interested in how small a p-value might be and saying it is over a value is never useful.

<sup>5</sup>All too often, I read journal articles that have under-utilized, under-reported, mis-applied, or mis-interpreted statistical methods and results. One of the reasons that I wanted to write this book is to help more people move from basic statistical knowledge to correct use of intermediate statistical methods and beginning to see the potential in more advanced statistical methods. It has taken me many years of being a statistician just to feel armed for battle when confronted with new applications and two stat courses are not enough to get you there, but you have to start somewhere. You are only a couple of hundred hours into your 10,000 hours required for mastery. This presentation is intended get you some solid fundamentals to build on or a few intermediate tools to use if this is your last statistics training experience.

#### 9.4. MULTI-VARIATE MODELS ARE ESSENTIAL FOR UNDERSTANDING VERTEBRATE DIVERSIFICATION IN DE

```
bm$logDBCs <- log(bm$DBCs)
bm$logDBFs <- log(bm$DBFs)
bm$TJK <- factor(bm$TJK)
require(car)
scatterplot(logSpecies~logDBCs|TJK, data=bm, smooth=F,
            main="Scatterplot of log-diversity vs log-DBCs by period", lwd=2)
```

The following results will allow us to explore models similar to theirs. One “full” model they considered is:

$$\log(\text{count})_i = \beta_0 + \beta_1 \log(\text{DBC})_i + \beta_2 I_{\text{TJK},i} + \varepsilon_i$$

which was compared to

$$\log(\text{count})_i = \beta_0 + \beta_1 \log(\text{DBF})_i + \beta_2 I_{\text{TJK},i} + \varepsilon_i$$

as well as the simpler models that each suggests:

$$\begin{aligned}\log(\text{count})_i &= \beta_0 + \beta_1 \log(\text{DBC})_i + \varepsilon_i, \\ \log(\text{count})_i &= \beta_0 + \beta_1 \log(\text{DBF})_i + \varepsilon_i, \\ \log(\text{count})_i &= \beta_0 + \beta_2 I_{\text{TJK},i} + \varepsilon_i, \text{ and} \\ \log(\text{count})_i &= \beta_0 + \varepsilon_i.\end{aligned}$$

Both versions of the models (based on *DBF* or *DBC*) start with an MLR model with a quantitative variable and two slopes. We can obtain some of the needed model selection results from the first full model using:

```
bd1 <- lm(logSpecies~logDBCs+TJK, data=bm)
require(MuMin)
options(na.action = "na.fail")
dredge(bd1, rank="AIC",
       extra=c("R^2", adjRsq=function(x) summary(x)$adj.r.squared))
```

```
## Global model call: lm(formula = logSpecies ~ logDBCs + TJK, data = bm)
## ---
## Model selection table
##   (Intrc) lgDBC TJK      R^2    adjRsq df  logLik  AIC delta weight
## 4 -1.0890 0.7243  + 0.580900 0.54440 4 -12.652 33.3  0.00  0.987
## 2  0.1988 0.4283     0.369100 0.34280 3 -17.969 41.9  8.63  0.013
## 1  2.5690          0.000000 0.00000 2 -23.956 51.9 18.61  0.000
## 3  2.5300          + 0.004823 -0.03664 3 -23.893 53.8 20.48  0.000
## Models ranked by AIC(x)
```

And from the second model:

```
bd2 <- lm(logSpecies~logDBFs+TJK, data=bm)
dredge(bd2, rank="AIC",
       extra=c("R^2", adjRsq=function(x) summary(x)$adj.r.squared))
```

```
## Global model call: lm(formula = logSpecies ~ logDBFs + TJK, data = bm)
## ---
## Model selection table
##   (Intrc) lgDBF TJK      R^2    adjRsq df  logLik  AIC delta weight
## 4 -2.4100 1.3710  + 0.519900 0.47810 4 -14.418 36.8  0.00  0.995
## 2  0.5964 0.4882     0.209800 0.17690 3 -20.895 47.8 10.95  0.004
## 1  2.5690          0.000000 0.00000 2 -23.956 51.9 15.08  0.001
## 3  2.5300          + 0.004823 -0.03664 3 -23.893 53.8 16.95  0.000
## Models ranked by AIC(x)
```

The top AIC model is  $\log(\text{count})_i = \beta_0 + \beta_1 \log(\text{DBC})_i + \beta_2 I_{\text{TJK},i} + \varepsilon_i$  with an AIC of 33.3. The next best model was  $\log(\text{count})_i = \beta_0 + \beta_1 \log(\text{DBF})_i + \beta_2 I_{\text{TJK},i} + \varepsilon_i$  with an AIC of 36.8, so 3.5 AIC units worse than the top model. We put these two runs of results together in Table 9.1, re-computing all the AICs based on the top model from the first full model considered.

Table 9.1: Model comparison table.

Model	R <sup>2</sup>	adj R <sup>2</sup>	df	logLik	AIC	ΔAIC
$\log(\text{count})_i = \beta_0 + \beta_1 \log(\text{DBC})_i + \beta_2 I_{\text{TJK},i} + \varepsilon_i$	0.5809	0.5444	4	-12.652	33.3	0
$\log(\text{count})_i = \beta_0 + \beta_1 \log(\text{DBF})_i + \beta_2 I_{\text{TJK},i} + \varepsilon_i$	0.5199	0.4781	4	-14.418	36.8	3.5
$\log(\text{count})_i = \beta_0 + \beta_1 \log(\text{DBC})_i + \varepsilon_i$	0.3691	0.3428	3	-17.969	41.9	8.6
$\log(\text{count})_i = \beta_0 + \beta_1 \log(\text{DBF})_i + \varepsilon_i$	0.2098	0.1769	3	-20.895	47.8	14.5
$\log(\text{count})_i = \beta_0 + \varepsilon_i$	0	0	2	-23.956	51.9	18.6
$\log(\text{count})_i = \beta_0 + \beta_2 I_{\text{TJK},i} + \varepsilon_i$	0.0048	-0.0366	3	-23.893	53.8	20.5

Table 9.1 suggests some interesting leads to the worst performing model on the AIC measure, ranking below a model with nothing in it (mean-only) and 20.5 AIC units worse than the top model. But the two top models distinctly benefit from the inclusion of *TJK*. This suggests that after controlling for the sampling effort, either through *DBC* or *DBF*, the differences in the stages captured by *TJK* can be more clearly observed.

So the top model in our (correct) results<sup>6</sup> suggests an effect of *log(DBC)* and different intercepts for the two periods. We can interrogate this model further but probably should do something we should have done first, check the diagnostics (Figure 9.11) and consider our model assumptions as AICs are not valid if model assumptions are not met.

```
par(mfrow=c(2,2), oma=c(0,0,2,0))
plot(bd1)
```

The constant variance and assessment of influence do not suggest any real problems. The normality assumption is possibly violated but shows lighter tails than expected from a normal distribution and so should cause few problems with inferences (we would be looking for an answer of “yes, there is a violation of the normality assumption but, for a bonus point, that problem is minor because the pattern is not the problematic type of violation because...”). The other assumption that **is violated for all our models** is that the observations are independent. Between neighboring stages in time, there would likely be some sort of relationship in the biodiversity so we should not assume that the observations are independent (this is a *timeseries* of observations). The authors acknowledged this issue but unskillfully attempted to deal with it. Because an interaction was not considered in any of the models, there also is an assumption that the results are parallel enough for the two groups. The scatterplot in Figure 9.10 suggests that using parallel lines for the two groups is probably reasonable but a full assessment really should also explore that fully to verify that there is no support for an interaction.

Ignoring the violation of the independence assumption, we are otherwise ok to explore the model **more** and see what it tells us about biodiversity of *Sauropodomorphs*. The top model is estimated to be  $\widehat{\log(\text{count})}_i = -1.089 + 0.724 \log(\text{DBC})_i - 0.75 I_{\text{TJK},i}$ . This suggests that for the early observations (*TJK*=0), the model is  $\widehat{\log(\text{count})}_i = -1.089 + 0.724 \log(\text{DBC})_i$  and for the Cretaceous period (*TJK*=1), the model is  $\widehat{\log(\text{count})}_i = -1.089 + -0.75 + 0.724 \log(\text{DBC})_i$  which simplifies to  $\widehat{\log(\text{count})}_i = -1.84 + 0.724 \log(\text{DBC})_i$ . This suggests that the sampling efforts have the same effect on all observations and having an increase in *log-DBCs* is associated with increases in the mean *log-biodiversity*. Specifically, for a 1 log-count increase in the *log-DBCs*, we expect, on average, to have a 0.724 log-count change in the mean *log-biodiversity*, after accounting for different intercepts for the two periods considered. We could also translate this to the original count scale but will leave it as is, because their real question of interest involves the differences between the periods. The change in the y-intercepts of -0.76 suggests that the Cretaceous has a lower average *log-biodiversity* by

<sup>6</sup>They also had an error in their AIC results that is difficult to explain here but was due to an un-careful usage of the results from the more advanced models that account for autocorrelation, which seems to provide the proper ranking of models (*that they ignored*) but did not provide the correct differences among models.

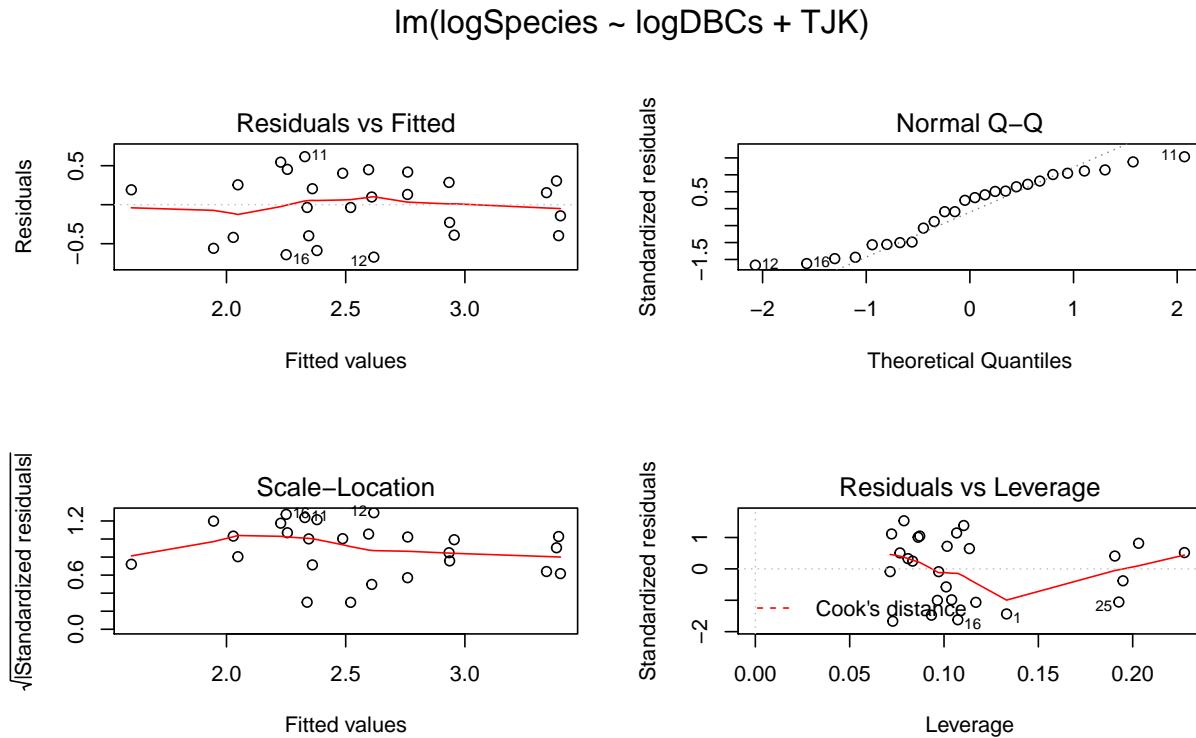


Figure 9.11: Diagnostic plots for the top AIC model.

0.75 log-count, after controlling for the log-sampling effort. This suggests that the *Cretaceous* had a lower corrected mean log-Sauropodomorph biodiversity ( $t_{23} = -3.41$ ;  $p\text{-value}=0.0024$ ) than the combined results for the Triassic and Jurassic. On the original count scale, this suggests  $\exp(-0.76) = 0.47$  times (53% drop in) the median biodiversity count per stage for Cretaceous versus the prior time period, after correcting for log-sampling effort in each stage.

```
summary(bd1)
```

```
##
## Call:
## lm(formula = logSpecies ~ logDBC}s + TJK, data = bm)
##
## Residuals:
##      Min       1Q   Median       3Q      Max 
## -0.6721 -0.3955  0.1149  0.2999  0.6158 
##
## Coefficients:
##             Estimate Std. Error t value Pr(>|t|)    
## (Intercept) -1.0887    0.6533 -1.666   0.1092    
## logDBC}s     0.7243    0.1288  5.622 1.01e-05  
## TJK1        -0.7598    0.2229 -3.409   0.0024  
## 
## Residual standard error: 0.4185 on 23 degrees of freedom
## Multiple R-squared:  0.5809, Adjusted R-squared:  0.5444 
## F-statistic: 15.94 on 2 and 23 DF,  p-value: 4.54e-05
```

Their study shows some interesting contrasts between methods. They tried to use AIC-based model selection methods across all the models but then used p-values to really make their final conclusions. This presents a philosophical inconsistency that bothers some more than others but should bother everyone. One thought is

whether they needed to use AICs at all since they wanted to use p-values? The one reason they might have preferred to use AICs is that it allows the direct comparison of

$$\log(\text{count})_i = \beta_0 + \beta_1 \log(\text{DBC})_i + \beta_2 I_{\text{TJK},i} + \varepsilon_i$$

to

$$\log(\text{count})_i = \beta_0 + \beta_1 \log(\text{DBF})_i + \beta_2 I_{\text{TJK},i} + \varepsilon_i,$$

exploring whether *DBC* or *DBF* is the “better” with *TJK* in the model. There is no hypothesis test to compare these two models because one is not ***nested*** in the other – **it is not possible to get from one model to the other by setting one or more slope coefficients to 0 so we can't test our way from one model to the other one.** The AICs suggest that the model with *DBC* and *TJK* is better than the model with *DBF* and *TJK*, so that helps us make that decision. After that step, we could rely on *t*-tests or ANOVA *F*-tests to decide whether further refinement is suggested/possible for the model with *DBC* and *TJK*. This would provide the direct inferences that they probably want and are trying to obtain from AICs along with p-values in their paper.

Finally, their results would actually be more valid if they had used a set of statistical methods designed for modeling responses that are counts of events or things, especially those whose measurements change as a function of sampling effort; models called ***Poisson rate models*** would be ideal for their application. The other aspect of the biodiversity that they measured for each stage was the duration of the stage. They never incorporated that information and it makes sense given their interests in comparing biodiversity across stages, not understanding why more or less biodiversity might occur. But other researchers might want to estimate the biodiversity after also controlling for the length of time that the stage lasted and the sampling efforts involved in detecting the biodiversity of each stage, models that are only a few steps away from those considered here. In general, this study presents some of the pitfalls of attempting to use advanced statistical methods as well as hinting at the benefits. The statistical models are the only way to access the results of interest; inaccurate usage of statistical models can provide inaccurate conclusions. They seemed to mostly get the right answers despite a suite of errors in their work.

## 9.5 General summary

As we wrap up, it is important to remember that these tools are limited by the quality of the data collected. If you are ever involved in applying these statistical models, whether in a research or industrial setting, make sure that the research questions are discussed before data collection. And before data collection is started, make sure that the methods will provide results that can address the research questions. And, finally, make sure someone involved in the project knows how to perform the appropriate statistical analysis. One way to make sure you know how to analyze a data set and, often, clarify the research questions and data collection needs, is to **make a fake data set and analyze it.** With this sort of preparation, many issues can be avoided. Remember to think about reasons why assumptions of your proposed method might be violated.

You are now **armed** and a bit **dangerous** with statistical methods. If you go to use them, remember the fundamentals and find the story in the data. After deciding on any research questions of interest, graph the data and make sure that the statistical methods will give you results that make some sense based on the graphical results. In the MLR results, it is possible that graphs will not be able to completely tell you the story, but all the other methods should follow the pictures you see. Even when (or especially when) you use sophisticated statistical methods, graphical presentations are critical to helping others understand the results. We have discussed examples that involve displaying categorical and quantitative variables and even some displays that bridge both types of variables. We hope you have enjoyed this material and been able to continue to develop your interests in statistics. You will see it in many future situations both in courses in your area of study and in real problems that need answers. You are also prepared to take more advanced statistics courses – if you want to discuss the next options, we are happy to provide additional information about the best next steps in learning statistics.

# Chapter 10

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