

MATH3014-6027 Design (and Analysis) of Experiments

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Preface

These are draft lecture notes for the modules MATH3014 and MATH6027 Design (and Analysis) of Experiments at the University of Southampton for academic year 2021-22. They are very much work in progress.

Southampton prerequisites for this module are MATH2010 or MATH6174 and STAT6123 (or equivalent modules on linear modelling).

Chapter 1

Motivation, introduction and revision

Definition 1.1. An **experiment** is the process through which data are collected to answer a scientific question (physical science, social science, actuarial science ...) by **deliberately** varying some features of the process under study in order to understand the impact of these changes on measureable responses.

In this course we consider only *intervention* experiments, in which some aspects of the process are under the experimenters' control. We do not consider *surveys* or *observational* studies.

Definition 1.2. **Design of experiments** is the topic in Statistics concerned with the selection of settings of controllable variables or factors in an experiment and their allocation to experimental units in order to maximise the effectiveness of the experiment at achieving its aim.

People have been designing experiments for as long as they have been exploring the natural world. Collecting empirical evidence is key for scientific development, as described in terms of clinical trials by xkcd:

Some notable milestones in the history of the design of experiments include:

- prior to the 20th century:
 - Francis Bacon (17th century; pioneer of the experimental methods)
 - James Lind (18th century; experiments to eliminate scurvy)
 - Charles Peirce (19th century; advocated randomised experiments and randomisation-based inference)
- 1920s: agriculture (particularly at the Rothamsted Agricultural Research Station)
- 1940s: clinical trials (Austin Bradford-Hill)
- 1950s: (manufacturing) industry (W. Edwards Deming; Genichi Taguchi)
- 1960s: psychology and economics (Vernon Smith)

- 1980s: in-silico (computer experiments)
- 2000s: online (A/B testing)

See Luca and Bazerman (2020) for further history, anecdotes and examples, especially from psychology and technology.

Figure 1.1 shows the Broadbalk agricultural field experiment at Rothamsted, one of the longest continuous running experiments in the world, which is testing the impact of different manures and fertilizers on the growth of winter wheat.



Figure 1.1: The Broadbalk experiment, Rothamsted (photograph taken 2016)

1.1 Motivation

Example 1.1. Consider an experiment to compare two treatments (e.g. drugs, diets, fertilisers, ...). We have n subjects (people, mice, plots of land, ...), each of which can be assigned one of the two treatments. A response (protein measurement, weight, yield, ...) is then measured.

Question: How many subjects should be assigned to each treatment to gain the most precise¹ inference about the difference in response from the two treatments?

Consider a linear statistical model² for the response (see MATH2010 or MATH6174/STAT6123):

¹Smallest variance.

²In this course, we will almost always start with a statistical model which we wish to use to answer our scientific question.

$$Y_j = \beta_0 + \beta_1 x_j + \varepsilon_j, \quad j = 1, \dots, n, \quad (1.1)$$

where $\varepsilon_j \sim N(0, \sigma^2)$ are independent and identically distributed errors and β_0, β_1 are unknown constants (parameters).

Let³

$$x_j = \begin{cases} -1 & \text{if treatment 1 is applied to the } j\text{th subject} \\ +1 & \text{if treatment 2 is applied to the } j\text{th subject,} \end{cases}$$

for $j = 1, \dots, n$.⁴

The difference in expected response from treatments 1 and 2 is

$$\begin{aligned} E[Y_j | x_j = +1] - E[Y_j | x_j = -1] &= \beta_0 + \beta_1 - \beta_0 + \beta_1 \\ &= 2\beta_1. \end{aligned} \quad (1.2)$$

Therefore, we require the the most precise estimator of β_1 possible. That is, we wish to make the variance of our estimator of β_1 as small as possible.

Parameters β_0 and β_1 can be estimated using least squares (see MATH2010 or MATH6174/STAT6123). For Y_1, \dots, Y_n , we can write the model down in matrix form:

$$\begin{bmatrix} Y_1 \\ \vdots \\ Y_n \end{bmatrix} = \begin{bmatrix} 1 & x_1 \\ \vdots & \vdots \\ 1 & x_n \end{bmatrix} \begin{bmatrix} \beta_0 \\ \beta_1 \end{bmatrix} + \begin{bmatrix} \varepsilon_1 \\ \vdots \\ \varepsilon_n \end{bmatrix}.$$

Or, by defining some notation:

$$\mathbf{Y} = \mathbf{X}\boldsymbol{\beta} + \boldsymbol{\varepsilon} \quad (1.3)$$

where

- \mathbf{Y} - $n \times 1$ vector of responses;
- \mathbf{X} - $n \times p$ model matrix;
- $\boldsymbol{\beta}$ - $p \times 1$ vector of parameters;
- $\boldsymbol{\varepsilon}$ - $n \times 1$ vector of errors.

The **least squares estimators**, $\hat{\boldsymbol{\beta}}$, are chosen such that the quadratic form

$$(\mathbf{Y} - \mathbf{X}\boldsymbol{\beta})^T(\mathbf{Y} - \mathbf{X}\boldsymbol{\beta})$$

³Other codings can be used: e.g. 0,1; see later in the module. It makes no difference for our current purpose.

⁴We will discuss the choice of *coding* -1, +1 later.

is minimised (recall that $E(\mathbf{Y}) = X\beta$). Therefore

$$\hat{\beta} = \operatorname{argmin}_{\beta} (\mathbf{Y}^T \mathbf{Y} + \beta^T X^T X \beta - 2\beta^T X^T \mathbf{Y}).$$

If we differentiate with respect to β ⁵,

$$\frac{\partial}{\partial \beta} = 2X^T X \beta - 2X^T \mathbf{Y},$$

and equate to 0, we get the estimators

$$\hat{\beta} = (X^T X)^{-1} X^T \mathbf{Y}. \quad (1.4)$$

These are the least squares estimators.

For Example 1.1,

$$X = \begin{bmatrix} 1 & x_1 \\ \vdots & \vdots \\ 1 & x_n \end{bmatrix}, \quad X^T X = \begin{bmatrix} n & \sum x_j \\ \sum x_j & \sum x_j^2 \end{bmatrix},$$

$$(X^T X)^{-1} = \frac{1}{n \sum x_j^2 - (\sum x_j)^2} \begin{bmatrix} \sum x_j^2 & -\sum x_j \\ -\sum x_j & n \end{bmatrix}, \quad X^T \mathbf{Y} = \begin{bmatrix} \sum Y_j \\ \sum x_j Y_j \end{bmatrix}.$$

Then,

$$\begin{aligned} \hat{\beta} = \begin{bmatrix} \hat{\beta}_0 \\ \hat{\beta}_1 \end{bmatrix} &= \frac{1}{n \sum x_j^2 - (\sum x_j)^2} \begin{bmatrix} \sum x_j^2 & -\sum x_j \\ -\sum x_j & n \end{bmatrix} \begin{bmatrix} \sum Y_j \\ \sum x_j Y_j \end{bmatrix} \\ &= \frac{1}{n \sum x_j^2 - (\sum x_j)^2} \begin{bmatrix} \sum Y_j \sum x_j^2 - \sum x_j \sum x_j Y_j \\ n \sum x_j Y_j - \sum x_j \sum Y_j \end{bmatrix}. \end{aligned} \quad (1.5)$$

We don't usually work through the algebra in such detail; the matrix form is often sufficient for theoretical and numerical calculations and software, e.g. **R**, can be used.

The precision of $\hat{\beta}$ is measured via the variance-covariance matrix, given by

$$\operatorname{Var}(\hat{\beta}) = \operatorname{Var}\{(X^T X)^{-1} X^T \mathbf{Y}\} \quad (1.6)$$

$$= (X^T X)^{-1} X^T \operatorname{Var}(\mathbf{Y}) X (X^T X)^{-1} \quad (1.7)$$

$$= (X^T X)^{-1} \sigma^2, \quad (1.8)$$

⁵Check the Matrix Cookbook for matrix calculus, amongst much else.

where $\mathbf{Y} \sim N(X\boldsymbol{\beta}, I_n\sigma^2)$, where I_n is an $n \times n$ identity matrix.

Hence, in our example,

$$\begin{aligned} \text{Var}(\hat{\boldsymbol{\beta}}) &= \frac{1}{n \sum x_j^2 - (\sum x_j)^2} \begin{bmatrix} \sum x_j^2 & -\sum x_j \\ -\sum x_j & n \end{bmatrix} \sigma^2 \\ &= \begin{bmatrix} \text{Var}(\hat{\beta}_0) & \text{Cov}(\hat{\beta}_0, \hat{\beta}_1) \\ \text{Cov}(\hat{\beta}_0, \hat{\beta}_1) & \text{Var}(\hat{\beta}_1) \end{bmatrix}. \end{aligned}$$

For estimating the difference between treatments, we are interested in

$$\begin{aligned} \text{Var}(\hat{\beta}_1) &= \frac{n}{n \sum x_j^2 - (\sum x_j)^2} \sigma^2 \\ &= \frac{n}{n^2 - (\sum x_j)^2} \sigma^2, \end{aligned}$$

as $x_j = \pm 1$, therefore $x_j^2 = 1$ for all $j = 1, \dots, n$, and hence $\sum x_j^2 = n$.

To achieve the most precise estimator, we need to minimise $\text{Var}(\hat{\beta}_1)$ or equivalently minimise $|\sum x_j|$. This goal can be achieved through the choice of x_1, \dots, x_n :

- as each x_j can only take one of two values, -1 or +1, this is equivalent to choosing the numbers of subjects assigned to treatment 1 and treatment 2;
- call these n_1 and n_2 respectively, with $n_1 + n_2 = n$

It is obvious that $\sum x_j = 0$ if and only if $n_1 = n_2$. Therefore, assuming n is even, the **optimal design** has

- $n_1 = \frac{n}{2}$ subjects assigned to treatment 1 and
- $n_2 = \frac{n}{2}$ subjects assigned to treatment 2.

For n odd, we choose $n_1 = \frac{n+1}{2}$, $n_2 = \frac{n-1}{2}$, or vice versa.

Definition 1.3. We can assess different designs using their **efficiency**:

$$\text{Eff} = \frac{\text{Var}(\hat{\beta}_1 | d^*)}{\text{Var}(\hat{\beta}_1 | d_1)} \quad (1.9)$$

where d_1 is a design we want to assess and d^* is the optimal design with smallest variance. Note that $0 \leq \text{Eff} \leq 1$.

In Figure 1.2 below, we plot this efficiency for Example 1.1, using different choices of n_1 . The total number of runs is fixed at $n = 100$, and the function **eff** calculates the efficiency from Definition 1.3 for a design with n_1 subjects assigned to treatment 1. Clearly, efficiency of 1 is achieved when $n_1 = n_2$ (equal allocation of treatments 1 and 2). If $n_1 = 0$ or $n_1 = 1$, the efficiency is zero; we cannot estimate the difference between two treatments if we only allocate subjects to one of them.

```
n <- 100  
eff <- function(n1) 1 - ((2 * n1 - n) / n)^2  
curve(eff, from = 0, to = n, ylab = "Eff", xlab = expression(n[1]))
```



Figure 1.2: Efficiencies for designs for Example 1.1 with different numbers, n_1 , of subjects assigned to treatment 1 when the total number of subjects is $n = 100$.

1.2 Aims of experimentation and some examples

Some reasons experiments are performed:

1. Treatment comparison (Chapters 2 and 3)
 - compare several treatments (and choose the best)
 - e.g. clinical trial, agricultural field trial
2. Factor screening (Chapters 4, 5 and 6)
 - many complex systems may involve a large number of (discrete) factors (controllable features)
 - which of these factors have a substantive impact?
 - (relatively) small experiments
 - e.g. industrial experiments on manufacturing processes
3. Response surface exploration (Chapter 7)

- detailed description of relationship between important (continuous) variables and response
 - typically second order polynomial regression models
 - larger experiments, often built up sequentially
 - e.g. alcohol yields in a pharmaceutical experiments
4. Optimisation (Chapter 7)
- finding settings of variables that lead to maximum or minimum response
 - typically use response surface methods and sequential “hill climbing” strategy

1.3 Some definitions

Definition 1.4. The **response** Y is the outcome measured in an experiment; e.g. yield from a chemical process. The response from the n observations are denoted Y_1, \dots, Y_n .

Definition 1.5. Factors (discrete) or **variables** (continuous) are features which can be set or controlled in an experiment; m denotes the number of factors or variables under investigation. For discrete factors, we call the possible settings of the factor its **levels**. We denote by x_{ij} the value taken by factor or variable i in the j th run of the experiment ($i = 1, \dots, m; j = 1, \dots, n$).

Definition 1.6. The **treatments** or **support points** are the *distinct* combinations of factor or variable values in the experiment.

Definition 1.7. An experimental **unit** is the basic element (material, animal, person, time unit, ...) to which a treatment can be applied to produce a response.

In Example 1.1 (comparing two treatments):

- Response Y : Measured outcome, e.g. protein level or pain score in clinical trial, yield in an agricultural field trial.
- Factor x : “treatment” applied
- Levels

treatment 1	$x = -1$
treatment 2	$x = +1$
- Treatment or support point: Two treatments or support points
- Experimental unit: Subject (person, animal, plot of land, ...).

1.4 Principles of experimentation

Three fundamental principles that need to be considered when designing an experiment are:

- replication

- randomisation
- stratification (blocking)

1.4.1 Replication

Each treatment is applied to a number of experimental units, with the j th treatment replicated r_j times. This enables the estimation of the variances of treatment effect estimators; increasing the number of replications, or replicates, decreases the variance of estimators of treatment effects. (Note: proper replication involves independent application of the treatment to different experimental units, not just taking several measurements from the same unit).

1.4.2 Randomisation

Randomisation should be applied to the allocation of treatments to units. Randomisation protects against **bias**; the effect of variables that are unknown and potentially uncontrolled or subjectivity in applying treatments. It also provides a formal basis for inference and statistical testing.

For example, in a clinical trial to compare a new drug and a control random allocation protects against

- “unmeasured and uncontrollable” features (e.g. age, sex, health)
- bias resulting from the clinician giving new drug to patients who are sicker.

Clinical trials are usually also *double-blinded*, i.e. neither the healthcare professional nor the patient knows which treatment the patient is receiving.

1.4.3 Stratification (or blocking)

We would like to use a wide variety of experimental units (e.g. people or plots of land) to ensure **coverage** of our results, i.e. validity of our conclusions across the population of interest. However, if the sample of units from the population is too heterogenous, then this will induce too much random variability, i.e. increase σ^2 in $\varepsilon_j \sim N(0, \sigma^2)$, and hence increase the variance of our parameter estimators.

We can reduce this extraneous variation by splitting our units into homogenous sets, or **blocks**, and including a blocking term in the model. The simplest blocked experiment is a **randomised complete block design**, where each block contains enough units for all treatments to be applied. Comparisons can then be made *within* each block.

Basic principle: block what you can, randomise what you cannot.

Later we will look at blocking in more detail, and the principle of **incomplete blocks**.

1.5 Revision on the linear model

Recall: $\mathbf{Y} = X\boldsymbol{\beta} + \boldsymbol{\varepsilon}$, with $\boldsymbol{\varepsilon} \sim N(\mathbf{0}, I_n\sigma^2)$. Let the j th row of X be denoted \mathbf{x}_j^T , which holds the values of the predictors, or explanatory variables, for the j th observation. Then

$$Y_j = \mathbf{x}_j^T \boldsymbol{\beta} + \varepsilon_j, \quad j = 1, \dots, n.$$

For example, quite commonly, for continuous variables

$$\mathbf{x}_j = (1, x_{1j}, x_{2j}, \dots, x_{mj})^T,$$

and so

$$\mathbf{x}_j^T \boldsymbol{\beta} = \beta_0 + \beta_1 x_{1j} + \dots + \beta_m x_{mj}.$$

The least squares estimators are given by

$$\hat{\boldsymbol{\beta}} = (X^T X)^{-1} X^T \mathbf{Y},$$

with

$$\text{Var}(\hat{\boldsymbol{\beta}}) = (X^T X)^{-1} \sigma^2.$$

1.5.1 Variance of a Prediction/Fitted Value

A prediction of the mean response at point \mathbf{x}_0 (which may or may not be in the design) is

$$\hat{Y}_0 = \mathbf{x}_0^T \hat{\boldsymbol{\beta}},$$

with

$$\begin{aligned} \text{Var}(\hat{Y}_0) &= \text{Var}(\mathbf{x}_0^T \hat{\boldsymbol{\beta}}) \\ &= \mathbf{x}_0^T \text{Var}(\hat{\boldsymbol{\beta}}) \mathbf{x}_0 \\ &= \mathbf{x}_0^T (X^T X)^{-1} \mathbf{x}_0 \sigma^2. \end{aligned}$$

For a linear model, this variance depends only on the assumed regression model and the design (through X), the point at which prediction is to be made (\mathbf{x}_0) and the value of σ^2 ; it does not depend on data \mathbf{Y} or parameters $\boldsymbol{\beta}$.

Similarly, we can find the variance-covariance matrix of the fitted values:

$$\text{Var}(\hat{Y}) = \text{Var}(X\hat{\beta}) = X(X^T X)^{-1} X^T \sigma^2.$$

1.5.2 Analysis of Variance and R^2 as Model Comparison

To assess the goodness-of-fit of a model, we can use the residual sum of squares

$$\begin{aligned} \text{RSS} &= (\mathbf{Y} - X\hat{\beta})^T (\mathbf{Y} - X\hat{\beta}) \\ &= \sum_{j=1}^n \left\{ Y_j - \mathbf{x}_j^T \hat{\beta} \right\}^2 \\ &= \sum_{j=1}^n r_j^2, \end{aligned}$$

where

$$r_j = Y_j - \mathbf{x}_j^T \hat{\beta}.$$

Often, a comparison is made to the null model

$$Y_j = \beta_0 + \varepsilon_j,$$

i.e. $Y_i \sim N(\beta_0, \sigma^2)$. The residual sum of squares for the null model is given by

$$\text{RSS}(\text{null}) = \mathbf{Y}^T \mathbf{Y} - m \bar{Y}^2,$$

as

$$\hat{\beta}_0 = \bar{Y} = \frac{1}{n} \sum_{j=1}^n Y_j.$$

How do we compare these models?

1. Ratio of residual sum of squares:

$$\begin{aligned} R^2 &= 1 - \frac{\text{RSS}}{\text{RSS}(\text{null})} \\ &= 1 - \frac{(\mathbf{Y} - X\hat{\beta})^T (\mathbf{Y} - X\hat{\beta})}{\mathbf{Y}^T \mathbf{Y} - n \bar{Y}^2}. \end{aligned}$$

The quantity $0 \leq R^2 \leq 1$ is sometimes called the **coefficient of multiple determination**:

- high R^2 implies that the model describes much of the variation in the data;
- **but** note that R^2 will always increase as p (the number of explanatory variables) increases, with $R^2 = 1$ when $p = n$;
- some software packages will report the adjusted R^2 .

$$\begin{aligned} R_a^2 &= 1 - \frac{\text{RSS}/(n-p)}{\text{RSS}(\text{null})/(n-1)} \\ &= 1 - \frac{(\mathbf{Y} - X\hat{\boldsymbol{\beta}})^T(\mathbf{Y} - X\hat{\boldsymbol{\beta}})/(n-p)}{(\mathbf{Y}^T\mathbf{Y} - n\bar{Y}^2)/(n-1)}; \end{aligned}$$

- R_a^2 does not necessarily increase with p (as we divide by degrees of freedom to adjust for complexity of the model).
2. Analysis of variance (ANOVA): An ANOVA table is compact way of presenting the results of (sequential) comparisons of nested models. You should be familiar with an ANOVA table of the following form.

Table 1.1: A standard ANOVA table.

Source	Degress of Freedom	(Sequential) Sum of Squares	Mean Square
Regression	$p - 1$	By subtraction; see (1.12)	Reg SS/ $(p - 1)$
Residual	$n - p$	$(\mathbf{Y} - X\hat{\boldsymbol{\beta}})^T(\mathbf{Y} - X\hat{\boldsymbol{\beta}})^6$	RSS/ $(n - p)$
Total	$n - 1$	$\mathbf{Y}^T\mathbf{Y} - n\bar{Y}^2$ ⁷	

In row 1 of Table 1.1 above,

$$\text{Regression SS} = \text{Total SS} - \text{RSS} = \mathbf{Y}^T\mathbf{Y} - n\bar{Y}^2 - (\mathbf{Y} - X\hat{\boldsymbol{\beta}})^T(\mathbf{Y} - X\hat{\boldsymbol{\beta}}) \quad (1.10)$$

$$= -n\bar{Y}^2 - \hat{\boldsymbol{\beta}}^T(X^T X)\hat{\boldsymbol{\beta}} + 2\hat{\boldsymbol{\beta}}^T X^T \mathbf{Y} \quad (1.11)$$

$$= \hat{\boldsymbol{\beta}}^T(X^T X)\hat{\boldsymbol{\beta}} - n\bar{Y}^2, \quad (1.12)$$

⁶Residual sum of squares for the full regression model.

⁷Residual sum of squares for the null model.

with the last line following from

$$\begin{aligned}\hat{\beta}^T X^T \mathbf{Y} &= \hat{\beta}^T (X^T X) (X^T X)^{-1} X^T \mathbf{Y} \\ &= \hat{\beta}^T (X^T X) \hat{\beta}\end{aligned}$$

This idea can be generalised to the comparison of a *sequence* of nested models - see Problem Sheet 1.

Hypothesis testing is performed using the mean square:

$$\frac{\text{Regression SS}}{p-1} = \frac{\hat{\beta}^T (X^T X) \hat{\beta} - n\bar{Y}^2}{p-1}.$$

Under $H_0 : \beta_1 = \dots = \beta_{p-1} = 0$

$$\begin{aligned}\frac{\text{Regression SS}/(p-1)}{\text{RSS}/(n-p)} &= \frac{(\hat{\beta}^T (X^T X) \hat{\beta} - n\bar{Y}^2)/(p-1)}{(\mathbf{Y} - X\hat{\beta})^T (\mathbf{Y} - X\hat{\beta})/(n-p)} \\ &\sim F_{p-1, n-p},\end{aligned}$$

an F distribution with $p-1$ and $n-p$ degrees of freedom; defined via the ratio of two independent χ^2 distributions.

Also,

$$\frac{\text{RSS}}{n-p} = \frac{(\mathbf{Y} - X\hat{\beta})^T (\mathbf{Y} - X\hat{\beta})}{n-p} = \hat{\sigma}^2$$

is an unbiased estimator for σ^2 , and

$$\frac{(n-p)}{\sigma^2} \hat{\sigma}^2 \sim \chi_{n-p}^2.$$

This is a Chi-squared distribution with $n-p$ degrees of freedom (see MATH2010 or MATH6174 notes).

1.6 Exercises

1. (Adapted from Morris, 2011) A classic and famous example of a simple hypothetical experiment was described by Fisher (1935):

A lady declares that by tasting a cup of tea made with milk she can discriminate whether the milk or the tea infusion was added first to the cup. We will consider the problem of designing an experiment by means of which this assertion can be tested. For this purpose let us first lay down a simple form of experiment with a view to studying its limitations and its characteristics, both those that are essential to the experimental method, when well developed, and those that are not essential but auxiliary.

Our experiment consists in mixing eight cups of tea, four in one way and four in the other, and presenting them to the subject for judgement in a random order. The subject has been told in advance of what the test will consist, namely that she will be asked to taste eight cups, that these shall be four of each kind, and that they shall be presented to her in a random order, that is an order not determined arbitrarily by human choice, but by the actual manipulation of the physical apparatus used in games of chance, cards, dice, roulettes, etc., or, more expeditiously, from a published collection of random sampling numbers purporting to give the actual results of such manipulation⁸. Her task is to divide the 8 cups into two sets of 4, agreeing, if possible, with the treatments received.

- a. Define the treatments in this experiment.
- b. Identify the units in this experiment.
- c. How might a “physical apparatus” from a “game of chance” be used to perform the randomisation. Explain one example.
- d. Suppose eight tea cups are available for this experiment but they are not identical. Instead they come from two sets. Four are made from heavy, thick porcelain; four from much lighter china. If each cup can only be used once, how might this fact be incorporated into the design of the experiment?

Solution

- a. There are two treatments in the experiment - the two ingredients “milk first” and “tea first”.
- b. The experimental units are the “cups of tea”, made up from the tea and milk used and also the cup itself.
- c. The simplest method here might be to select four black playing cards and four red playing cards, assign one treatment to each colour, shuffle the cards, and then draw them in order. The colour drawn indicates the treatment that should be used to make the next cup of tea. This operation would give one possible randomisation.

We could of course also use R.

⁸Now, we would use routines such as `sample` in R.

```
sample(rep(c("Milk first", "Tea first"), c(4, 4)), size = 8, replace = F)

## [1] "Milk first" "Milk first" "Milk first" "Tea first" "Tea first"
## [6] "Tea first" "Milk first" "Tea first"
```

- d. Type of cup could be considered as a blocking factor. One way of incorporating it would be to split the experiment into two (blocks), each with four cups (two milk first, two tea first). We would still wish to randomise allocation of treatments to units within blocks.

```
# block 1
sample(rep(c("Milk first", "Tea first"), c(2, 2)), size = 4, replace = F)

## [1] "Milk first" "Tea first" "Tea first" "Milk first"

# block 2
sample(rep(c("Milk first", "Tea first"), c(2, 2)), size = 4, replace = F)

## [1] "Milk first" "Tea first" "Milk first" "Tea first"
```

2. Consider the linear model

$$\mathbf{y} = X\boldsymbol{\beta} + \boldsymbol{\varepsilon},$$

with \mathbf{y} an $n \times 1$ vector of responses, X a $n \times p$ model matrix and $\boldsymbol{\varepsilon}$ a $n \times 1$ vector of independent and identically distributed random variables with constant variance σ^2 .

- a. Derive the least squares estimator $\hat{\boldsymbol{\beta}}$ for this multiple linear regression model, and show that this estimator is unbiased. Using the definition of (co)variance, show that

$$\text{Var}(\hat{\boldsymbol{\beta}}) = (X^T X)^{-1} \sigma^2.$$

- b. If $\boldsymbol{\varepsilon} \sim N(\mathbf{0}, I_n \sigma^2)$, with I_n being the $n \times n$ identity matrix, show that the maximum likelihood estimators for $\boldsymbol{\beta}$ coincide with the least squares estimators.

Solution

- a. The method of least squares minimises the sum of squared differences between the responses and the expected values, that is, minimises the expression

$$(\mathbf{y} - X\boldsymbol{\beta})^T (\mathbf{y} - X\boldsymbol{\beta}) = \mathbf{y}^T \mathbf{y} - 2\boldsymbol{\beta}^T X^T \mathbf{y} + \boldsymbol{\beta}^T X^T X \boldsymbol{\beta}.$$

Differentiating with respect to the vector $\boldsymbol{\beta}$, we obtain

$$\frac{\partial}{\partial \boldsymbol{\beta}} = -2X^T \mathbf{y} + 2X^T X \boldsymbol{\beta}.$$

Set equal to $\mathbf{0}$ and solve:

$$X^T X \hat{\beta} = X^T \mathbf{y} \Rightarrow \hat{\beta} = (X^T X)^{-1} X^T \mathbf{y}.$$

The estimator $\hat{\beta}$ is unbiased:

$$E(\hat{\beta}) = (X^T X)^{-1} X^T E(\mathbf{y}) = (X^T X)^{-1} X^T X \beta = \beta,$$

and has variance:

$$\begin{aligned} \text{Var}(\hat{\beta}) &= E \left\{ \left[\hat{\beta} - E(\hat{\beta}) \right] \left[\hat{\beta} - E(\hat{\beta}) \right]^T \right\} \\ &= E \left\{ \left[\hat{\beta} - \beta \right] \left[\hat{\beta} - \beta \right]^T \right\} \\ &= E \left\{ \hat{\beta} \hat{\beta}^T - 2\beta \hat{\beta}^T + \beta \beta^T \right\} \\ &= E \left\{ (X^T X)^{-1} X^T \mathbf{y} \mathbf{y}^T X (X^T X)^{-1} - 2\beta \mathbf{y}^T X (X^T X)^{-1} + \beta \beta^T \right\} \\ &= (X^T X)^{-1} X^T E(\mathbf{y} \mathbf{y}^T) X (X^T X)^{-1} - 2\beta E(\mathbf{y}^T) X (X^T X)^{-1} + \beta \beta^T \\ &= (X^T X)^{-1} X^T [\text{Var}(\mathbf{y}) + E(\mathbf{y}) E(\mathbf{y}^T)] X (X^T X)^{-1} - 2\beta \beta^T X^T X (X^T X)^{-1} + \beta \beta^T \\ &= (X^T X)^{-1} X^T \left[I_N \sigma^2 + X \beta \beta^T X^T \right] X (X^T X)^{-1} - \beta \beta^T \\ &= (X^T X)^{-1} \sigma^2. \end{aligned}$$

b. As $\mathbf{y} \sim N(X\beta, I_N \sigma^2)$, the likelihood is given by

$$L(\beta; \mathbf{y}) = (2\pi\sigma^2)^{-N/2} \exp \left\{ -\frac{1}{2\sigma^2} (\mathbf{y} - X\beta)^T (\mathbf{y} - X\beta) \right\}.$$

The log-likelihood is given by

$$l(\beta; \mathbf{y}) = -\frac{1}{2\sigma^2} (\mathbf{y} - X\beta)^T (\mathbf{y} - X\beta) + \text{constant}.$$

Up to a constant, this expression is $-1 \times$ the least squares equations; hence maximising the log-likelihood is equivalent to minimising the least squares equation.

3. Consider the two nested linear models

$$(i) \ Y_j = \beta_0 + \beta_1 x_{1j} + \beta_2 x_{2j} + \dots + \beta_{p_1} x_{p_1 j} + \varepsilon_j, \text{ or } \mathbf{y} = X_1 \beta_1 + \varepsilon,$$

- (ii) $Y_j = \beta_0 + \beta_1 x_{1j} + \beta_2 x_{2j} + \dots + \beta_{p_1} x_{p_1 j} + \beta_{p_1+1} x_{(p_1+1)j} + \dots + \beta_{p_2} x_{p_2 j} + \varepsilon_j$, or $\mathbf{y} = X_1 \boldsymbol{\beta}_1 + X_2 \boldsymbol{\beta}_2 + \boldsymbol{\varepsilon}$

with $\varepsilon_j \sim N(0, \sigma^2)$, and $\varepsilon_j, \varepsilon_k$ independent ($\boldsymbol{\varepsilon} \sim N(\mathbf{0}, I_n \sigma^2)$).

- Construct an ANOVA table to compare model (ii) with the null model $Y_j = \beta_0 + \varepsilon_j$.
- Extend this ANOVA table to compare models (i) and (ii) by further decomposing the regression sum of squares for model (ii).

Hint: which residual sum of squares are you interested in to compare models (i) and (ii)?

You should end up with an ANOVA table of the form

Source	Degrees of freedom	Sums of squares	Mean square
Model (i)	p_1	?	?
Model (ii)	p_2	?	?
Residual	$n - p_1 - p_2 - 1$?	?
Total	$n - 1$	$\mathbf{y}^T \mathbf{y} - n\bar{Y}^2$	

The second row of the table gives the **extra sums of squares** for the additional terms in fitting model (ii), over and above those in model (i).

- Calculate the extra sum of squares for fitting the terms in model (i), over and above those terms only in model (ii), i.e. those held in $X_2 \boldsymbol{\beta}_2$. Construct an ANOVA table containing both the extra sum of squares for the terms only in model (i) and the extra sum of squares for the terms only in model (ii). Comment on the table.

Solution

- From lectures

Source	Degrees of freedom	Sums of squares	Mean square
Regression	$p_1 + p_2$	$\hat{\boldsymbol{\beta}}^T X^T X \hat{\boldsymbol{\beta}} - n\bar{Y}^2$	$\left(\hat{\boldsymbol{\beta}}^T X^T X \hat{\boldsymbol{\beta}} - n\bar{Y}^2 \right) / (p_1 + p_2)$
Residual	$n - p_1 - p_2 - 1$	$(\mathbf{y} - X \hat{\boldsymbol{\beta}})^T (\mathbf{y} - X \hat{\boldsymbol{\beta}})$	$\text{RSS} / (n - p_1 - p_2 - 1)$
Total	$n - 1$	$\mathbf{y}^T \mathbf{y} - n\bar{Y}^2$	

where

$$\begin{aligned}
\text{RSS}(\text{null}) - \text{RSS}(\text{ii}) &= \mathbf{y}^T \mathbf{y} - n\bar{Y}^2 - (\mathbf{y} - X\hat{\beta})^T (\mathbf{y} - X\hat{\beta}) \\
&= \mathbf{y}^T \mathbf{y} - n\bar{Y}^2 - \mathbf{y}^T \mathbf{y} + 2\mathbf{y}^T X\hat{\beta} - \hat{\beta}^T X^T X\hat{\beta} \\
&= 2\hat{\beta}^T X^T X\hat{\beta} - \hat{\beta}^T X^T X\hat{\beta} - n\bar{Y}^2 \\
&= \hat{\beta}^T X^T X\hat{\beta} - n\bar{Y}^2.
\end{aligned}$$

b. To compare model (i) with the null model, we compute

$$\begin{aligned}
\text{RSS}(\text{null}) - \text{RSS}(\text{i}) &= \mathbf{y}^T \mathbf{y} - N\bar{Y}^2 - (\mathbf{y} - X_1\hat{\beta}_1)^T (\mathbf{y} - X_1\hat{\beta}_1) \\
&= \hat{\beta}_1^T X_1^T X_1\hat{\beta}_1 - n\bar{Y}^2.
\end{aligned}$$

To compare models (i) and (ii), we compare them both to the null model, and look at the difference between these comparisons:

$$\begin{aligned}
[\text{RSS}(\text{null}) - \text{RSS}(\text{ii})] - [\text{RSS}(\text{null}) - \text{RSS}(\text{i})] &= \text{RSS}(\text{i}) - \text{RSS}(\text{ii}) \\
&= (\mathbf{y} - X_1\hat{\beta}_1)^T (\mathbf{y} - X_1\hat{\beta}_1) - (\mathbf{y} - X\hat{\beta})^T (\mathbf{y} - X\hat{\beta}) \\
&= \hat{\beta}^T X^T X\hat{\beta} - \hat{\beta}_1^T X_1^T X_1\hat{\beta}_1.
\end{aligned}$$

Source	Degrees of freedom	Sums of squares	Mean square
Regression	$p_1 + p_2$	$\hat{\beta}^T X^T X\hat{\beta} - n\bar{Y}^2$	$(\hat{\beta}^T X^T X\hat{\beta} - n\bar{Y}^2) / (p_1 + p_2)$
Model (i)	p_1	$\hat{\beta}_1^T X_1^T X_1\hat{\beta}_1 - n\bar{Y}^2$	$(\hat{\beta}_1^T X_1^T X_1\hat{\beta}_1 - n\bar{Y}^2) / p_1$
Extra due to Model (ii)	p_2	$\hat{\beta}^T X^T X\hat{\beta} - \hat{\beta}_1^T X_1^T X_1\hat{\beta}_1$	$(\hat{\beta}^T X^T X\hat{\beta} - \hat{\beta}_1^T X_1^T X_1\hat{\beta}_1) / p_2$
Residual	$n - p_1 - p_2 - 1$	$(\mathbf{y} - X\hat{\beta})^T (\mathbf{y} - X\hat{\beta})$	$\text{RSS} / (n - p_1 - p_2 - 1)$
Total	$n - 1$	$\mathbf{y}^T \mathbf{y} - n\bar{Y}^2$	

By definition, the Model (i) SS and the Extra SS for Model (ii) sum to the Regression SS.

a. The extra sum of squares for the terms in model (i) over and above those

in model (ii) are obtained through comparison of the models

ia. $\mathbf{y} = X_2\boldsymbol{\beta}_2 + \boldsymbol{\varepsilon}$,

iiia. $\mathbf{y} = X_1\boldsymbol{\beta}_1 + X_2\boldsymbol{\beta}_2 + \boldsymbol{\varepsilon} = X\boldsymbol{\beta} + \boldsymbol{\varepsilon}$

Extra sum of squares for model (iia):

$$\begin{aligned} [\text{RSS}(\text{null}) - \text{RSS}(\text{iia})] - [\text{RSS}(\text{null}) - \text{RSS}(\text{ia})] &= \text{RSS}(\text{ia}) - \text{RSS}(\text{iia}) \\ &= (\mathbf{y} - X_2\hat{\boldsymbol{\beta}}_2)^T(\mathbf{y} - X_2\hat{\boldsymbol{\beta}}_2) - (\mathbf{y} - X\hat{\boldsymbol{\beta}})^T(\mathbf{y} - X\hat{\boldsymbol{\beta}}) \\ &= \hat{\boldsymbol{\beta}}^T X^T X \hat{\boldsymbol{\beta}} - \hat{\boldsymbol{\beta}}_2^T X_2^T X_2 \hat{\boldsymbol{\beta}}_2. \end{aligned}$$

Hence, an ANOVA table for the extra sums of squares is given by

Source	Degrees of freedom	Sums of squares	Mean square
Regression	$p_1 + p_2$	$\hat{\boldsymbol{\beta}} X^T X \hat{\boldsymbol{\beta}} - n\bar{Y}^2$	$\left(\hat{\boldsymbol{\beta}} X^T X \hat{\boldsymbol{\beta}} - n\bar{Y}^2 \right) / (p_1 + p_2)$
Extra Model (i)	p_1	$\hat{\boldsymbol{\beta}}^T X^T X \hat{\boldsymbol{\beta}} - \hat{\boldsymbol{\beta}}_2^T X_2^T X_2 \hat{\boldsymbol{\beta}}_2$	$\left(\hat{\boldsymbol{\beta}}^T X^T X \hat{\boldsymbol{\beta}} - \hat{\boldsymbol{\beta}}_2^T X_2^T X_2 \hat{\boldsymbol{\beta}}_2 \right) / p_1$
Extra Model (ii)	p_2	$\hat{\boldsymbol{\beta}}^T X^T X \hat{\boldsymbol{\beta}} - \hat{\boldsymbol{\beta}}_1^T X_1^T X_1 \hat{\boldsymbol{\beta}}_1$	$\left(\hat{\boldsymbol{\beta}}^T X^T X \hat{\boldsymbol{\beta}} - \hat{\boldsymbol{\beta}}_1^T X_1^T X_1 \hat{\boldsymbol{\beta}}_1 \right) / p_2$
Residual	$n - p_1 - p_2 - 1$	$(\mathbf{y} - X\hat{\boldsymbol{\beta}})^T(\mathbf{y} - X\hat{\boldsymbol{\beta}})$	$\text{RSS} / (n - p_1 - p_2 - 1)$
Total	$n - 1$	$\mathbf{y}^T \mathbf{y} - n\bar{Y}^2$	

Note that for these *adjusted* sums of squares, in general the extra sum of squares for model (i) and (ii) do not sum to the regression sum of squares. This will only be the case if the columns of X_1 and X_2 are mutually orthogonal, i.e. $X_1^T X_2 = \mathbf{0}$.

Chapter 2

Completely randomised designs

The simplest form of experiment we will consider compares t different **unstructured** treatments. By unstructured, we mean the treatments form a discrete collection, not related through the settings of other experimental features (compare with factorial experiments in Chapter 4). We also make the assumption that there are no restrictions in the randomisation of treatments to experimental units (compare with Chapter 3 on blocking). A designs for such an experiment is therefore called a **completely randomised design** (CRD).

Example 2.1. Pulp experiment (Wu and Hamada, 2009, ch. 2)

In a paper pulping mill, an experiment was run to examine differences between the reflectance (brightness; ratio of amount of light leaving a target to the amount of light striking the target) of sheets of pulp made by $t = 4$ operators. The data are given in Table 2.1 below.

```
pulp <- data.frame(operator = rep(factor(1:4), 5),  
                  repetition = rep(1:5, rep(4, 5)),  
                  reflectance = c(59.8, 59.8, 60.7, 61.0, 60.0, 60.2, 60.7, 60.8,  
                                60.8, 60.4, 60.5, 60.6, 60.8, 59.9, 60.9, 60.5, 59.8, 60.0, 60.5, 60.4, 60.5, 60.6, 60.8, 59.9, 60.9, 60.5, 59.8, 60.0, 60.5))  
  
knitr::kable(  
  tidyr::pivot_wider(pulp, names_from = operator, values_from = reflectance)[, -1],  
  col.names = paste("Operator", 1:4),  
  caption = "Pulp experiment: reflectance values (unitless) from four different operators."  
)
```

The experiment has one factor (operator) with four levels (sometimes called a one-way layout). The CRD employed has equal replication of each treatment (operator).

Table 2.1: Pulp experiment: reflectance values (unitless) from four different operators.

Operator 1	Operator 2	Operator 3	Operator 4
59.8	59.8	60.7	61.0
60.0	60.2	60.7	60.8
60.8	60.4	60.5	60.6
60.8	59.9	60.9	60.5
59.8	60.0	60.3	60.5

We can informally compare the responses from these four treatments graphically.

```
boxplot(reflectance ~ operator, data = pulp)
```

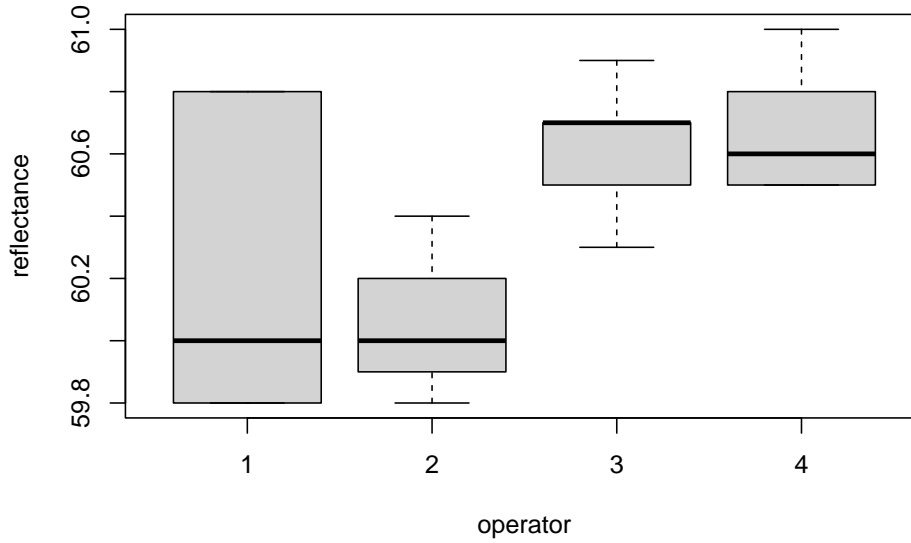


Figure 2.1: Pulp experiment: distributions of reflectance from the four operators.

Figure 2.1 shows that, relative to the variation, there may be a difference in the mean response between treatments 1 and 2, and 3 and 4. In this chapter, we will see how to make this comparison formally using linear models, and to assess how the choice of design impacts our results.

Throughout this chapter we will assume the i th treatment is applied to n_i experimental unit, with total number of runs $n = \sum_{i=1}^t n_i$ in the experiment.

2.1 A unit-treatment linear model

An appropriate, and common, model to describe data from such experiments when the response is continuous is given by

$$y_{ij} = \mu + \tau_i + \varepsilon_{ij}, \quad i = 1, \dots, t; j = 1, \dots, n_i, \quad (2.1)$$

where y_{ij} is the response from the j th application of treatment i , μ is a constant parameter, τ_i is the effect of the i th treatment, and ε_{ij} is the random individual effect from each experimental unit with $E(\varepsilon_{ij}) = 0$ and $\text{Var}(\varepsilon_{ij}) = \sigma^2$. All random errors are assumed independent and here we also assume $\varepsilon_{ij} \sim N(0, \sigma^2)$.

Model (2.1) assumes that each treatment can be randomly allocated to one of the n experimental units, and that the response observed is independent of the allocation of all the other treatments (the stable unit treatment value assumption or SUTVA).

Why is this model appropriate and commonly used? The expected response from the application of the i th treatment is

$$E(y_{ij}) = \mu + \tau_i.$$

The parameter μ can be thought of as representing the impact of many different features particular to **this** experiment but common to all units, and τ_i is the deviation due to applying treatment i . From the application of two different hypothetical experiments, A and B, the expected response from treatment i may be different due to a different overall mean. From experiment A:

$$E(y_{ij}) = \mu_A + \tau_i.$$

From experiment B:

$$E(y_{ij}) = \mu_B + \tau_i.$$

But the **difference** between treatments k and l ($k, l = 1, \dots, t$)

$$\begin{aligned} E(y_{kj}) - E(y_{lj}) &= \mu_A + \tau_k - \mu_A - \tau_l \\ &= \tau_k - \tau_l, \end{aligned}$$

is constant across different experiments. This concept of **comparison** underpins most design of experiments, and will be applied throughout this module.

2.2 The partitioned linear model

In matrix form, we can write model (2.1) as

$$\mathbf{y} = X_1\mu + X_2\boldsymbol{\tau} + \boldsymbol{\varepsilon},$$

where $X_1 = \mathbf{1}_n$, the n -vector with every entry equal to one,

$$X_2 = \bigoplus_{i=1}^t \mathbf{1}_{n_i} = \begin{bmatrix} \mathbf{1}_{n_1} & \mathbf{0}_{n_1} & \cdots & \mathbf{0}_{n_1} \\ \mathbf{0}_{n_2} & \mathbf{1}_{n_2} & \cdots & \mathbf{0}_{n_2} \\ \vdots & & \ddots & \vdots \\ \mathbf{0}_{n_t} & \mathbf{0}_{n_t} & \cdots & \mathbf{1}_{n_t} \end{bmatrix},$$

with \bigoplus denoting “direct sum”, $\mathbf{0}_{n_i}$ is the n_i -vector with every entry equal to zero, $\boldsymbol{\tau} = [\tau_1, \dots, \tau_t]^\top$ and $\boldsymbol{\varepsilon} = [\varepsilon_{11}, \dots, \varepsilon_{tn_t}]^\top$.

Why are we partitioning the model? Going back to our discussion of the role of μ and τ_i , it is clear that we are not interested in estimating μ , which represents an experiment-specific contribution to the expected mean. Our only interest is in estimating the (differences between the) τ_i . Hence, we can treat μ as a nuisance parameter.

If we define $X = [X_1 \mid X_2]$ and $\boldsymbol{\beta}^\top = [\mu \mid \boldsymbol{\tau}^\top]$, we can write the usual least squares equations

$$X^\top X \hat{\boldsymbol{\beta}} = X^\top \mathbf{y} \tag{2.2}$$

as a system of two matrix equations

$$\begin{aligned} X_1^\top X_1 \hat{\mu} + X_1^\top X_2 \hat{\boldsymbol{\tau}} &= X_1^\top \mathbf{y} \\ X_2^\top X_1 \hat{\mu} + X_2^\top X_2 \hat{\boldsymbol{\tau}} &= X_2^\top \mathbf{y}. \end{aligned}$$

Assuming $(X_1^\top X_1)^{-1}$ exists, which it does in this case, we can pre-multiply the first of these equations by $X_2^\top X_1 (X_1^\top X_1)^{-1}$ and subtract it from the second equation to obtain

$$\begin{aligned} X_2^\top [I_n - X_1 (X_1^\top X_1)^{-1} X_1^\top] X_1 \hat{\mu} + X_2^\top [I_n - X_1 (X_1^\top X_1)^{-1} X_1^\top] X_2 \hat{\boldsymbol{\tau}} \\ = X_2^\top [I_n - X_1 (X_1^\top X_1)^{-1} X_1^\top] \mathbf{y}. \end{aligned}$$

Writing $H_1 = X_1 (X_1^\top X_1)^{-1} X_1^\top$, we obtain

$$X_2^T[I_n - H_1]X_1\hat{\mu} + X_2^T[I_n - H_1]X_2\hat{\tau} = X_2^T[I_n - H_1]\mathbf{y}. \quad (2.3)$$

The matrix H_1 is a “hat” matrix for a linear model containing only the term μ , and hence $H_1X_1 = X_1$ (see MATH2010 or STAT6123). Hence the first term in (2.3) is zero, and we obtain the **reduced normal equations** for τ :

$$X_2^T[I_n - H_1]X_2\hat{\tau} = X_2^T[I_n - H_1]\mathbf{y}. \quad (2.4)$$

Note that the solutions from (2.4) are not different from the solution to $\hat{\tau}$ that would be obtained from solving (2.2); equation (2.4) is simply a re-expression, where we have eliminated the nuisance parameter μ . This fact means that we rarely need to solve (2.4) explicitly.

Recalling that for a hat matrix, $I_n - H_1$ is idempotent and symmetric (see MATH2010 or MATH6174), if we define

$$X_{2|1} = (I_n - H_1)X_2,$$

then we can rewrite equation (2.4) as

$$X_{2|1}^T X_{2|1} \hat{\tau} = X_{2|1}^T \mathbf{y}, \quad (2.5)$$

which are the normal equations for a linear model with expectation $E(\mathbf{y}) = X_{2|1}\tau$.

2.3 Reduced normal equations for the CRD

For the CRD discussed in this chapter, $X_1^T X_1 = n$, the total number of runs in the experiment¹. Hence $(X_1^T X_1)^{-1} = 1/n$ and $H_1 = \frac{1}{n}J_n$, with J_n the $n \times n$ matrix with all entries equal to 1.

The adjusted model matrix then has the form

$$\begin{aligned} X_{2|1} &= (I_n - H_1)X_2 \\ &= X_2 - \frac{1}{n}J_n X_2 \\ &= X_2 - \frac{1}{n}[n_1 \mathbf{1}_n | \cdots | n_t \mathbf{1}_n]. \end{aligned} \quad (2.6)$$

That is, every column of X_2 has been adjusted by the subtraction of the column mean from each entry². Also notice that each row of $X_{2|1}$ has a row-sum equal

¹In later chapters we will see examples where X_1 has > 1 columns, and hence $X_1^T X_1$ is a matrix.

²Often called “column centred”

to zero ($= 1 - \sum_{i=1}^t n_t/n$). Hence, $X_{2|1}$ is not of full column rank, and so the reduced normal equations do not have a unique solution³.

Although (2.5), and hence, (2.2), have no unique solution⁴, it can be shown that both $\widehat{X_{2|1}\tau}$ and $\widehat{X\beta}$ have unique solutions. Hence fitted values $\hat{\mathbf{y}} = \widehat{X\beta}$ and the residual sum of squares

$$RSS = (\mathbf{y} - \widehat{X\beta})^T (\mathbf{y} - \widehat{X\beta})$$

are both uniquely defined for any solution to (2.2). That is, every solution to the normal equations leads to the same fitted values and residual sum of squares.

In MATH2010 and STAT6123 we fitted models with categorical variables by defining a set of dummy variables and estimating a reduced model. Here, we will take a slightly different approach and study which combinations of parameters from (2.1) are estimable, and in particular which linear combinations of the treatment parameters τ_i we can estimate.

Let's study equation (2.5) in more detail. We have

$$\begin{aligned} X_{2|1}^T X_{2|1} &= X_2^T (I_n - H_1) X_2 \\ &= X_2^T X_2 - X_2^T H_1 X_2 \\ &= \text{diag}(\mathbf{n}) - \frac{1}{n} X_2^T J_n X_2 \\ &= \text{diag}(\mathbf{n}) - \frac{1}{n} \mathbf{n} \mathbf{n}^T, \end{aligned}$$

where $\mathbf{n}^T = (n_1, \dots, n_t)$. Hence, the reduced normal equations become

$$\left[\text{diag}(\mathbf{n}) - \frac{1}{n} \mathbf{n} \mathbf{n}^T \right] \hat{\boldsymbol{\tau}} = X_2^T \mathbf{y} - \frac{1}{n} X_2^T J_n \mathbf{y} \quad (2.7)$$

$$= X_2^T \mathbf{y} - \mathbf{n} \bar{y}_{..}, \quad (2.8)$$

where $\bar{y}_{..} = \frac{1}{n} \sum_{i=1}^t \sum_{j=1}^{n_i} y_{ij}$, i.e. the overall average of the observations from the experiment.

From (2.8) we obtain a system of t equations, each having the form

$$\hat{\tau}_i - \hat{\tau}_w = \bar{y}_{i.} - \bar{y}_{..}, \quad (2.9)$$

³If we recalled the material on “dummy” variables from MATH2010 or STAT6123, we would already have realised this.

⁴That is, for any two solutions $\tilde{\beta}_1$ and $\tilde{\beta}_2$, $X\tilde{\beta}_1 = X\tilde{\beta}_2$.

where $\hat{\tau}_w = \frac{1}{n} \sum_{i=1}^t n_i \hat{\tau}_i$ and $\bar{y}_{i.} = \frac{1}{n_i} \sum_{j=1}^{n_i} y_{ij}$ ($i = 1, \dots, t$).

These t equations are not independent; when multiplied by the n_i , they sum to zero due to the linear dependency between the columns of $X_{2|1}$. Hence, there is no unique solution to $\hat{\tau}$ from equation (2.8). However, we can estimate certain linear combinations of the τ_i , called *contrasts*.

2.4 Contrasts

Definition 2.1. A treatment **contrast** is a linear combination $\mathbf{c}^T \boldsymbol{\tau}$ with coefficient vector $\mathbf{c}^T = (c_1, \dots, c_t)$ such that $\mathbf{c}^T \mathbf{1} = 0$; that is, $\sum_{i=1}^t c_i = 0$.

For example, assume we have $t = 3$ treatments, then the following vectors \mathbf{c} all define contrasts:

1. $\mathbf{c}^T = (1, -1, 0)$,
2. $\mathbf{c}^T = (1, 0, -1)$,
3. $\mathbf{c}^T = (0, 1, -1)$.

In fact, they define all $\binom{3}{2} = 3$ pairwise comparisons between treatments. The following are also contrasts:

4. $\mathbf{c}^T = (2, -1, -1)$,
5. $\mathbf{c}^T = (0.5, -1, 0.5)$,

each comparing the sum, or average, of expected responses from two treatments to the expected response from the remaining treatment.

The following are not contrasts, as $\mathbf{c}^T \mathbf{1} \neq 0$:

6. $\mathbf{c}^T = (2, -1, 0)$,
7. $\mathbf{c}^T = (1, 0, 0)$,

with the final example once again demonstrating that we cannot estimate the individual τ_i .

2.5 Treatment contrast estimators in the CRD

We estimate a treatment contrast $\mathbf{c}^T \boldsymbol{\tau}$ in the CRD via linear combinations of equations (2.9):

$$\begin{aligned} \sum_{i=1}^t c_i \hat{\tau}_i - \sum_{i=1}^t c_i \hat{\tau}_w &= \sum_{i=1}^t c_i \bar{y}_{i.} - \sum_{i=1}^t c_i \bar{y}_{..} \\ \Rightarrow \sum_{i=1}^t c_i \hat{\tau}_i &= \sum_{i=1}^t c_i \bar{y}_{i.} , \end{aligned}$$

as $\sum_{i=1}^t c_i \hat{\tau}_w = \sum_{i=1}^t c_i \bar{y}_{i..} = 0$, as $\sum_{i=1}^t c_i = 0$. Hence, the unique estimator of the contrast $\mathbf{c}^T \boldsymbol{\tau}$ has the form

$$\widehat{\mathbf{c}^T \boldsymbol{\tau}} = \sum_{i=1}^t c_i \bar{y}_{i..}.$$

That is, we estimate the contrast in the treatment effects by the corresponding contrast in the treatment means.

The variance of this estimator is straightforward to obtain:

$$\begin{aligned} \text{var} \left(\widehat{\mathbf{c}^T \boldsymbol{\tau}} \right) &= \sum_{i=1}^t c_i^2 \text{var}(\bar{y}_{i..}) \\ &= \sigma^2 \sum_{i=1}^t c_i^2 / n_i, \end{aligned}$$

as, under our model assumptions, each $\bar{y}_{i..}$ is an average of independent observations with variance σ^2 . Similarly, from model (2.1) we can derive the distribution of $\widehat{\mathbf{c}^T \boldsymbol{\tau}}$ as

$$\widehat{\mathbf{c}^T \boldsymbol{\tau}} \sim N \left(\mathbf{c}^T \boldsymbol{\tau}, \sigma^2 \sum_{i=1}^t c_i^2 / n_i \right).$$

Confidence intervals and hypothesis tests for $\mathbf{c}^T \boldsymbol{\tau}$ can be constructed/conducted using this distribution, e.g.

- a $100(1 - \frac{\alpha}{2})\%$ confidence interval:

$$\mathbf{c}^T \boldsymbol{\tau} \in \sum_{i=1}^t c_i \bar{y}_{i..} \pm t_{n-t, 1-\frac{\alpha}{2}} s \sqrt{\sum_{i=1}^t c_i^2 / n_i},$$

where $t_{n-t, 1-\frac{\alpha}{2}}$ is the $1 - \frac{\alpha}{2}$ quantile of a t -distribution with $n - t$ degrees of freedom and

$$s^2 = \frac{1}{n-t} \sum_{i=1}^t \sum_{j=1}^{n_i} (y_{ij} - \bar{y}_{i..})^2 \quad (2.10)$$

is the estimate of σ^2 .

- the hypothesis $H_0 : \mathbf{c}^T \boldsymbol{\tau} = 0$ against the two-sided alternative $H_1 : \mathbf{c}^T \boldsymbol{\tau} \neq 0$ is rejected using a test of with confidence level $1 - \alpha/2$ if

Table 2.2: Pulp experiment: reflectance values (unitless) from four different operators.

Operator 1	Operator 2	Operator 3	Operator 4
59.8	59.8	60.7	61.0
60.0	60.2	60.7	60.8
60.8	60.4	60.5	60.6
60.8	59.9	60.9	60.5
59.8	60.0	60.3	60.5

$$\frac{|\sum_{i=1}^t c_i \bar{y}_i|}{s \sqrt{\sum_{i=1}^t c_i^2 / n_i}} > t_{n-t, 1-\frac{\alpha}{2}}.$$

2.6 Analysing CRDs in R

Let's return to Example 2.1.

```
knitr::kable(
  tidyr::pivot_wider(pulp, names_from = operator, values_from = reflectance)[, -1],
  col.names = paste("Operator", 1:4),
  caption = "Pulp experiment: reflectance values (unitless) from four different operators."
)
```

Clearly, we could directly calculate, and then compare, mean responses for each operator. However, there are (at least) two other ways we can proceed which use the fact we are fitting a linear model. These will be useful when we consider more complex models.

1. Using `pairwise.t.test`.

```
with(pulp,
  pairwise.t.test(reflectance, operator, p.adjust.method = 'none'))

##
## Pairwise comparisons using t tests with pooled SD
##
## data: reflectance and operator
##
##      1      2      3
## 2 0.396 -      -
## 3 0.084 0.015 -
## 4 0.049 0.008 0.775
##
```

```
## P value adjustment method: none
```

This function performs hypothesis tests for all pairwise treatment comparisons (with a default confidence level of 0.95). Here we can see that operators 1 and 4, 2 and 3, and 2 and 4 have statistically significant differences.

2. Using `lm` and the `emmeans` package.

```
pulp.lm <- lm(reflectance ~ operator, data = pulp)
anova(pulp.lm)

## Analysis of Variance Table
##
## Response: reflectance
##           Df Sum Sq Mean Sq F value Pr(>F)
## operator    3   1.34   0.447    4.2  0.023 *
## Residuals  16   1.70   0.106
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

pulp.emm <- emmeans::emmeans(pulp.lm, ~ operator)
pairs(pulp.emm, adjust = 'none')

## contrast estimate      SE df t.ratio p.value
## 1 - 2           0.18 0.206 16   0.873  0.3955
## 1 - 3          -0.38 0.206 16  -1.843  0.0839
## 1 - 4          -0.44 0.206 16  -2.134  0.0486
## 2 - 3          -0.56 0.206 16  -2.716  0.0153
## 2 - 4          -0.62 0.206 16  -3.007  0.0083
## 3 - 4          -0.06 0.206 16  -0.291  0.7748
```

Here, we have first fitted the linear model object. The `lm` function, by default, will have set up dummy variables with the first treatment (operator) as a baseline (see MATH2010 or STAT6123). We then take the intermediate step of calculating the ANOVA table for this experiment, and use an F-test to compare the model accounting for operator differences to the null model; there are differences between operators at the 5% significance level,

The choice of dummy variables in the linear model is unimportant; any set could be used⁵, as in the next line we use the `emmeans` function (from the package of the same name) to specify that we are interested in estimating contrasts in the factor `operator` (which specifies our treatments in this experiment). Finally, the `pairs` command performs hypothesis tests for all pairwise comparisons between the four treatments. The results are the same as those obtained from using `pairwise.t.test`.

⁵Recall that although μ and τ are not uniquely estimable, fitted values $\hat{y}_i = \hat{\mu} + \hat{\tau}_i$ are, and hence it does not matter which dummy variables we use in `lm`.

Our preferred approach is using method 2 (`lm` and `emmeans`), for four main reasons:

- a. The function `contrasts` in the `emmeans` package can be used to estimate arbitrary treatment contrasts (see `help("contrast-methods")`).

```
# same as `pairs` above
emmeans::contrast(pulp.emm, "pairwise", adjust = "none")
```

```
## contrast estimate SE df t.ratio p.value
## 1 - 2          0.18 0.206 16   0.873  0.3955
## 1 - 3         -0.38 0.206 16  -1.843  0.0839
## 1 - 4         -0.44 0.206 16  -2.134  0.0486
## 2 - 3         -0.56 0.206 16  -2.716  0.0153
## 2 - 4         -0.62 0.206 16  -3.007  0.0083
## 3 - 4         -0.06 0.206 16  -0.291  0.7748
```

```
# estimating single contrast c = (1, -.5, -.5)
# comparing operator 1 with operators 2 and 3
contrast1v23.emmc <- function(levs)
  data.frame('t1 v avg t2 t3' = c(1, -.5, -.5, 0))
emmeans::contrast(pulp.emm, 'contrast1v23')
```

```
## contrast      estimate SE df t.ratio p.value
## t1.v.avg.t2.t3    -0.1 0.179 16  -0.560  0.5832
```

- b. It more easily generalises to the more complicated models we will see in Chapter 3.
- c. It explicitly acknowledges that we have fitted a linear model, and so encourages us to check the model assumptions (see Exercise 3).
- d. It is straightforward to apply adjustments for multiple comparisons.

2.7 Multiple comparisons

When we perform hypothesis testing, we choose the critical region (i.e. the rule that decides if we reject H_0) to control the probability of a type I error; that is, we control the probability of incorrectly rejecting H_0 . If we need to test multiple hypotheses, e.g. to test all pairwise differences, we need to consider the overall probability of incorrectly rejecting **one or more** null hypothesis. This is called the **experiment-wise** or **family-wise** error rate.

For Example 2.1, there are $\binom{4}{2} = 6$ pairwise comparisons. Under the assumption that all tests are independent⁶, assuming each individual test has type I error 0.05, the experiment-wise type I error rate is given by:

⁶They aren't, but it simplifies the maths!

```
alpha <- 0.05
1 - (1 - alpha)^6
```

```
## [1] 0.2649
```

An experiment-wise error rate of 0.2649 is substantially greater than 0.05. Hence, we would expect to make many more type I errors than may be desirable. `xkcd` has a fun example:

```
alpha <- 0.05
1 - (1 - alpha)^20
```

```
## [1] 0.6415
```

Therefore it is usually desirable to maintain some control of the experiment-wise type I error rate. We will consider two methods.

1. The **Bonferroni method**. An upper bound on the experiment-wise type I error rate for testing k hypotheses can be shown to be

$$\begin{aligned} P(\text{wrongly reject at least one of } H_0^1, \dots, H_0^k) &= P\left(\bigcup_{i=1}^k \{\text{wrongly reject } H_0^i\}\right) \\ &\leq \sum_{i=1}^k \underbrace{P(\text{wrongly reject } H_0^i)}_{\leq \alpha} \\ &\leq k\alpha. \end{aligned}$$

Hence a *conservative*⁷ adjustment for multiple comparisons is to test each hypothesis at size α/k , i.e. for the CRD compare to the quantile $t_{n-t, 1-\frac{\alpha}{2k}}$ (or multiply each individual p-value by k).

For Example 2.1, we can test all pairwise comparisons, each at size α/k using the `adjustment` argument in `pairs`.

```
pairs(pulp.emm, adjust = 'bonferroni')
```

```
## contrast estimate SE df t.ratio p.value
## 1 - 2          0.18 0.206 16  0.873  1.0000
## 1 - 3         -0.38 0.206 16 -1.843  0.5034
## 1 - 4         -0.44 0.206 16 -2.134  0.2918
## 2 - 3         -0.56 0.206 16 -2.716  0.0915
## 2 - 4         -0.62 0.206 16 -3.007  0.0501
## 3 - 4         -0.06 0.206 16 -0.291  1.0000
##
## P value adjustment: bonferroni method for 6 tests
```

⁷So the experiment-wise type I error will actually be less than the prescribed α

Now, only one comparison is significant at an experiment-wise type I error rate of $\alpha = 0.05$ (operators 2 and 4).

2. **Tukey's method.** An alternative approach that gives an exact experiment-wise error rate of $100\alpha\%$ compares the t statistic to a critical value from the studentised range distribution⁸, given by $\frac{1}{\sqrt{2}}q_{t,n-t,1-\alpha}$ with $q_{t,n-t,1-\alpha}$ the $1 - \alpha$ quantile from the studentised range distribution (available in R as `qtukey`).

For Example 2.1:

```
pairs(pulp.emm)
```

```
## contrast estimate SE df t.ratio p.value
## 1 - 2          0.18 0.206 16  0.873 0.8185
## 1 - 3         -0.38 0.206 16 -1.843 0.2903
## 1 - 4         -0.44 0.206 16 -2.134 0.1845
## 2 - 3         -0.56 0.206 16 -2.716 0.0658
## 2 - 4         -0.62 0.206 16 -3.007 0.0377
## 3 - 4         -0.06 0.206 16 -0.291 0.9911
##
## P value adjustment: tukey method for comparing a family of 4 estimates
```

The default adjustment in the `pairs` function is the Tukey method. Comparing the p-values for each comparison using unadjusted t-tests, the Bonferroni and Tukey methods:

```
pairs.u <- pairs(pulp.emm, adjust = 'none')
pairs.b <- pairs(pulp.emm, adjust = 'bonferroni')
pairs.t <- pairs(pulp.emm)
data.frame(transform(pairs.b)[, 1:5], Bonf.p.value = transform(pairs.b)[, 6], Tukey.p.value = tra
```

```
## contrast estimate SE df t.ratio Bonf.p.value Tukey.p.value
## 1 1 - 2          0.18 0.2062 16  0.8731      1.00000      0.81854
## 2 1 - 3         -0.38 0.2062 16 -1.8433      0.50336      0.29030
## 3 1 - 4         -0.44 0.2062 16 -2.1343      0.29182      0.18448
## 4 2 - 3         -0.56 0.2062 16 -2.7164      0.09150      0.06579
## 5 2 - 4         -0.62 0.2062 16 -3.0074      0.05009      0.03767
## 6 3 - 4         -0.06 0.2062 16 -0.2910      1.00000      0.99108
## unadjust.p.value
## 1          0.395509
## 2          0.083893
## 3          0.048637
## 4          0.015251
```

⁸Given two independent samples u_1, \dots, u_l and v_1, \dots, v_m from the same distribution, the studentised range distribution is the distribution of $\frac{R}{\sqrt{2}S}$, where $R = u_{\max} - u_{\min}$ is the range of the first sample, and $S = \sqrt{\frac{1}{m-1} \sum_{i=1}^m (v_i - \bar{v})^2}$ be the sample standard deviation of the second sample.

```
## 5          0.008349
## 6          0.774758
```

Although the decision on which hypotheses to reject (comparson 2 - 4) is the same here for both methods, the p-values from the Bonferroni method are all larger, reflecting its more conservative nature.

2.8 Impact of design choices on estimation

Recall from Section 2.5 that the width of a confidence interval for a contrast $\mathbf{c}^T \boldsymbol{\tau}$ is given by $2t_{n-t, 1-\frac{\alpha}{2}} s \sqrt{\sum_{i=1}^t c_i^2/n_i}$. The expectation of the square of this quantity is given by

$$4t_{n-t, 1-\frac{\alpha}{2}}^2 \sigma^2 \sum_{i=1}^t c_i^2/n_i,$$

as $E(s^2) = \sigma^2$. It is intuitive that a good design should have small values of the square root of this quantity (divided by 2σ),

$$t_{n-t, 1-\frac{\alpha}{2}} \sqrt{\sum_{i=1}^t c_i^2/n_i},$$

which can be achieved either by increasing n , and hence reducing the size of the t -quantile, or for choice of the n_i for a fixed n , i.e. through choice of replication of each treatment.

2.8.1 Optimal treatment allocation

It is quite common that although the total number, n , of runs in the experiment may be fixed, the number n_1, n_2, \dots, n_t applied to the different treatments is under the experimenter's control. Choosing n_1, n_2 subject to $n_1 + n_2 = n$ was the first **optimal design** problem we encountered in Chapter 1.

Assume interest lies in estimating the set of p contrasts $\mathbf{c}_1^T \boldsymbol{\tau}, \dots, \mathbf{c}_p^T \boldsymbol{\tau}$, with $\mathbf{c}_l^T = (c_{l1}, \dots, c_{lt})$. One useful measure of the overall quality of the estimators of these p contrasts is the average variance, given by

$$\sigma^2 \sum_{l=1}^p \sum_{i=1}^t c_{li}^2/n_i.$$

So we will minimise this variance by allocating larger values of n_i to the treatments with correspondingly larger values of the contrast coefficients c_{li} . Therefore an approach to optimal allocation is to choose $\mathbf{n} = (n_1, \dots, n_t)^T$ so as to

$$\text{minimise } \phi(\mathbf{n}) = \sum_{l=1}^p \sum_{i=1}^t c_{li}^2/n_i \quad \text{subject to } \sum_{i=1}^t n_i = n. \quad (2.11)$$

This is a discrete optimisation problem (the n_i are integers). It is usually easier to solve the relaxed problem, where we allow continuous $0 \leq n_i \leq n$, and round the resulting solution to obtain integers. There is no guarantee that such a rounded allocation will actually be the optimal integer-valued solution, but it is usually fairly close.

To solve the continuous version of (2.11) we will use the method of Lagrange multipliers, where we define the function

$$h(\mathbf{n}, \lambda) = \phi(\mathbf{n}) + \lambda \left(\sum_{i=1}^t n_i - n \right),$$

introducing the new scalar variable λ , and solve the set of $t + 1$ equations:

$$\begin{aligned} \frac{\partial h}{\partial n_1} &= 0 \\ &\vdots \\ \frac{\partial h}{\partial n_t} &= 0 \\ \frac{\partial h}{\partial \lambda} &= 0. \end{aligned}$$

In this case, we have

$$\frac{\partial h}{\partial n_i} = - \sum_{l=1}^p c_{li}^2/n_i^2 + \lambda = 0, \quad i = 1, \dots, t, \quad (2.12)$$

and

$$\frac{\partial h}{\partial \lambda} = \sum_{i=1}^t n_i - n = 0.$$

This last equation ensures $\sum_{i=1}^t n_i = n$. From the t equations described by (2.12), we get

$$n_i \propto \sqrt{\sum_{l=1}^p c_{li}^2}$$

We don't need to explicitly solve for λ to find the normalising constant for each n_i . As we know $\sum_{i=1}^t n_i = n$, we obtain,

$$n_i = \frac{\sqrt{\sum_{l=1}^p c_{li}^2}}{\sum_{i=1}^t \sqrt{\sum_{l=1}^p c_{li}^2}} n. \quad (2.13)$$

Let's return to Example 2.1 and calculate the optimal allocations under two different sets of contrasts. First, we define an R function for calculating (2.13).

```
opt_ni <- function(C, n) {
  CtC <- t(C) %*% C
  n * sqrt(diag(CtC)) / sum(sqrt(diag(CtC)))
}
```

Checking that the function `opt_ni` matches (2.13) is left as an exercise.

Consider two sets of contrasts:

1. All pairwise comparisons between the four treatments

$$\begin{aligned} c_1 &= (-1, 1, 0, 0) \\ c_2 &= (-1, 0, 1, 0) \\ c_3 &= (-1, 0, 0, 1) \\ c_4 &= (0, -1, 1, 0) \\ c_5 &= (0, -1, 0, 1) \\ c_6 &= (0, 0, -1, 1). \end{aligned}$$

Calculating (2.13), we obtain

```
C <- matrix(
  c(
    -1, 1, 0, 0,
    -1, 0, 1, 0,
    -1, 0, 0, 1,
    0, -1, 1, 0,
    0, -1, 0, 1,
    0, 0, -1, 1),
  nrow = 6, byrow = T
)
opt_ni(C, 20)
```

```
## [1] 5 5 5 5
```

Hence confirming that equal replication of the treatments is optimal for minimising the average variance of estimators of the pairwise treatment differences.

2. If operator 4 is new to the mill, it may be desired to test their output to the average output from the other three operators, using a contrast with

coefficients $c = (1/3, 1/3, 1/3, -1)$. The allocation to minimise the variance of the corresponding estimator is given by:

```
C <- matrix(
  c(1/3, 1/3, 1/3, -1),
  nrow = 1
)
opt_ni(C, 20)

## [1] 3.333 3.333 3.333 10.000
```

So the optimal allocation splits 10 units between operators 1-3, and allocates 10 units to operator 4. There is no exact integer rounding possible, so we will use $n_1 = 4$, $n_2 = n_3 = 3$, $n_4 = 10$ and calculate the efficiency by comparing the variance of this allocation to that from the equally allocated design.

```
crd_var <- function(C, n) {
  CtC <- t(C) %*% C
  sum(diag(CtC) / n)
}
n_equal <- rep(5, 4)
n_opt <- c(4, 3, 3, 10)
crd_var(C, n_opt) / crd_var(C, n_equal)

## [1] 0.7569
```

So the efficiency of the equally allocated design for estimating this contrast is 75.69 %.

2.8.2 Overall size of the experiment

We can also consider the complementary question: suppose the proportion of runs that is to be allocated to each treatment has been fixed in advance, what size of experiment should be performed to meet the objectives? That is, given a fixed proportion, w_i , of resource to be allocated to the i th treatment, so that $n_i = nw_i$ units will be allocated to that treatment, what value of n should be chosen?

One way of thinking about this question is to consider the ratio

$$\begin{aligned} \frac{|\mathbf{c}^T \boldsymbol{\tau}|}{\sqrt{\text{Var}(\widehat{\mathbf{c}^T \boldsymbol{\tau}})}} &= \frac{|\mathbf{c}^T \boldsymbol{\tau}|}{\sqrt{\frac{\sigma^2}{n} \sum_{i=1}^t c_i^2 / w_i}} \\ &= \sqrt{n} \frac{|\mathbf{c}^T \boldsymbol{\tau}| / \sigma}{\sqrt{\sum_{i=1}^t c_i^2 / w_i}}, \end{aligned}$$

which is analogous to the test statistic for $H_0 : \mathbf{c}^T \boldsymbol{\tau} = 0$. For a given value of the signal-to-noise ratio $d = |\mathbf{c}^T \boldsymbol{\tau}| / \sigma$, we can choose n to result in a specified value of $T = |\mathbf{c}^T \boldsymbol{\tau}| / \sqrt{\widehat{\text{Var}}(\mathbf{c}^T \boldsymbol{\tau})}$:

$$n = T^2 \frac{\sum_{i=1}^t c_i^2 / w_i}{d^2}.$$

Returning to Example 2.1, assume are testing a single pairwise comparison and that we require $T = 3$, so that the null hypothesis would be comfortably rejected at the 5% level (cf 1.96 for a standard z-test). For equal allocation of the units to each treatment ($w_1 = \dots = w_4 = 1/4$) and a variety of different values of the signal-to-noise ratio d , we obtained the following optimal experiment sizes:

```
opt_n <- function(cv, prop, snr, target) target ^ 2 * c(t(cv) %*% diag( 1 / prop) %*% c
cv <- c(-1, 1, 0, 0)
w <- rep(1/4, 4)
snr <- c(0.5, 1, 1.5, 2, 2.5, 3)
cbind('Signal-to-noise' = snr, 'n' = opt_n(cv, w, snr, 3))
```

##	Signal-to-noise	n
## [1,]	0.5	288.00
## [2,]	1.0	72.00
## [3,]	1.5	32.00
## [4,]	2.0	18.00
## [5,]	2.5	11.52
## [6,]	3.0	8.00

So, for example, to achieve $T = 3$ with a signal-to-noise ratio of $d = 0.5$ requires $n = 288$ runs. As would be expected, the number of runs required to achieve this value of T decreases as the signal-to-noise ratio increases. For $d = 3$, only a very small experiment with $n = 8$ runs is needed.

2.9 Exercises

1. a. For Example 2.1, calculate the mean response for each operator and show that the treatment differences and results from hypothesis tests using the results in Section 2.5 are the same as those found in Section 2.6 using `pairwise.t.test`, and `emmeans`.
- b. Also check the results in Section 2.7 by (i) adjusting individual p-values (for Bonferroni) and (ii) using the `qtukey` command.

Solution

As a reminder, the data from the experiment is as follows.

Operator 1	Operator 2	Operator 3	Operator 4
59.8	59.8	60.7	61.0
60.0	60.2	60.7	60.8
60.8	60.4	60.5	60.6
60.8	59.9	60.9	60.5
59.8	60.0	60.3	60.5

The mean response, and variance, from each treatment is given by

operator	n_i	mean	variance
1	5	60.24	0.268
2	5	60.06	0.058
3	5	60.62	0.052
4	5	60.68	0.047

The sample variance, $s^2 = 0.106$, from (2.10). As $\sum_{i=1}^t c_i^2/n_i = \frac{2}{5}$ for contrast vectors \mathbf{c} corresponding to pairwise differences, the standard error of each pairwise difference is given by $\sqrt{\frac{2s^2}{5}} = 0.206$. Hence, we can create a table of pairwise differences, standard errors and test statistics.

contrast	estimate	SE	df	t.ratio	unadjust.p.value	Bonferroni	Tukey
1 - 2	0.18	0.206	16	0.873	0.396	1.000	0.819
1 - 3	-0.38	0.206	16	-1.843	0.084	0.503	0.290
1 - 4	-0.44	0.206	16	-2.134	0.049	0.292	0.184
2 - 3	-0.56	0.206	16	-2.716	0.015	0.092	0.066
2 - 4	-0.62	0.206	16	-3.007	0.008	0.050	0.038
3 - 4	-0.06	0.206	16	-0.291	0.775	1.000	0.991

Unadjusted p-values are obtained from the t-distribution, as twice the tail probabilities ($2 * (1 - \text{pt}(\text{abs}(\text{t.ratio}), 16))$). For Bonferroni, we simply multiply these p-values by $\binom{t}{2} = 6$, and then take the minimum of this value and 1. For the Tukey method, we use $1 - \text{ptukey}(\text{abs}(\text{t.ratio}) * \text{sqrt}(2), 4, 16)$ (see `?ptukey`).

Alternatively, to test each hypothesis at the 5% level, we can compare each t.ratio to (i) $\text{qt}(0.975, 16) = 2.12$ (unadjusted); (ii) $\text{qt}(1 - 0.025/6, 16) = 3.008$ (Bonferroni); or (iii) $\text{qtukey}(0.95, 4, 16) / \text{sqrt}(2) = 2.861$.

- (Adapted from Wu and Hamada, 2009) The bioactivity of four different drugs A , B , C and D for treating a particular illness was compared in a study and the following ANOVA table was given for the data:

Source	Degrees of freedom	Sums of squares	Mean square
Treatment	3	64.42	21.47
Residual	26	62.12	2.39
Total	29	126.54	

- i. What considerations should be made when assigning drugs to patients, and why?
- ii. Use an F -test to test at the 0.01 level the null hypothesis that the four drugs have the same bioactivity.
- iii. The average response from each treatment is as follows: $\bar{y}_A = 66.10$ ($n_A = 7$ patients), $\bar{y}_B = 65.75$ ($n_B = 8$), $\bar{y}_C = 62.63$ ($n_C = 9$), and $\bar{y}_D = 63.85$ ($n_D = 6$). Conduct hypothesis tests for all pair-wise comparisons using the Bonferroni and Tukey methods for an experiment-wise error rate of 0.05.
- iv. In fact, A and B are brand-name drugs and C and D are generic drugs. Test the null hypothesis at the 5% level that brand-name and generic drugs have the same bioactivity.

Solution

- i. Each patient should be randomly allocated to one of the drugs. This is to protect against possible bias from lurking variables, e.g. demographic variables or subjective bias from the study administrator (blinding the study can also help to protect against this).
- ii. Test statistic = (Treatment mean square)/(Residual mean square) = $21.47/2.39 = 8.98$. Under H_0 : no difference in bioactivity between the drugs, the test statistic follows an $F_{3,26}$ distribution, which has a 1% critical value of $\text{qf}(0.99, 3, 26) = 4.6366$. Hence, we can reject H_0 .
- iii. For each difference, the test statistic has the form

$$\frac{|\bar{y}_i - \bar{y}_j|}{s \sqrt{\frac{1}{n_i} + \frac{1}{n_j}}},$$

for $i, j = A, B, C, D; i \neq j$. The treatment means and repetitions are given in the question (note that not all n_i are equal). From the ANOVA table, we get $s^2 = 62.12/26 = 2.389$. The following table summarises the differences between drugs:

	$A - B$	$A - C$	$A - D$	$B - C$	$B - D$	$C - D$
Abs. difference	0.35	3.47	2.25	3.12	1.9	1.22
Test statistic	0.44	4.45	2.62	4.15	2.28	1.50

The Bonferroni critical value is $t_{26, 1-0.05/12} = 3.5069$. The Tukey critical value is $q_{4, 26, 0.95}/\sqrt{2} = 2.7433$ (available R as `qtukey(0.95, 4, 26) / sqrt(2)`). Hence under both methods, bioactivity of drugs A and C , and B and C , are significantly different.

Table 2.5: Naphthalene black experiment: yields (grams of standard colour) from six different batches of hydrochloric acid.

Batch 1	Batch 2	Batch 3	Batch 4	Batch 5	Batch 6
145	140	195	45	195	120
40	155	150	40	230	55
40	90	205	195	115	50
120	160	110	65	235	80
180	95	160	145	225	45

- iv. A suitable contrast has $\mathbf{c} = (0.5, 0.5, -0.5, -0.5)$, with $\mathbf{c}^T \boldsymbol{\tau} = (\tau_A + \tau_B)/2 - (\tau_C + \tau_D)/2$ (the difference in average treatment effects).

An estimate for this contrast is given by $(\bar{y}_A. + \bar{y}_B.)/2 - (\bar{y}_C. + \bar{y}_D.)/2$, with variance

$$\text{Var} \left(\frac{1}{2}(\bar{y}_A. + \bar{y}_B.) - \frac{1}{2}(\bar{y}_C. + \bar{y}_D.) \right) = \frac{\sigma^2}{4} \left(\frac{1}{n_A} + \frac{1}{n_B} + \frac{1}{n_C} + \frac{1}{n_D} \right).$$

Hence, a test statistic for $H_0 : \frac{1}{2}(\tau_A + \tau_B) - \frac{1}{2}(\tau_C + \tau_D) = 0$ is given by

$$\frac{\frac{1}{2}(\bar{y}_A. + \bar{y}_B.) - \frac{1}{2}(\bar{y}_C. + \bar{y}_D.)}{\sqrt{\frac{s^2}{4} \left(\frac{1}{n_A} + \frac{1}{n_B} + \frac{1}{n_C} + \frac{1}{n_D} \right)}} = \frac{2.685}{\frac{\sqrt{2.389}}{2} \sqrt{\frac{1}{7} + \frac{1}{8} + \frac{1}{9} + \frac{1}{6}}} = 4.70.$$

The critical value is $t_{26, 1-0.05/2} = 2.0555$. Hence, we can reject H_0 and conclude there is a difference between brand-name and generic drugs.

3. The below table gives data from a completely randomised design to compare six different batches of hydrochloric acid on the yield of a dye (naphthalene black 12B).

```
napblack <- data.frame(batch = rep(factor(1:6), rep(5, 6)),
  repetition = rep(1:5, 6),
  yield = c(145, 40, 40, 120, 180, 140, 155, 90, 160, 95,
            195, 150, 205, 110, 160, 45, 40, 195, 65, 145,
            195, 230, 115, 235, 225, 120, 55, 50, 80, 45)
)

knitr::kable(
  tidyr::pivot_wider(napblack, names_from = batch, values_from = yield)[, -1],
  col.names = paste("Batch", 1:6),
  caption = "Naphthalene black experiment: yields (grams of standard colour) from six different batches of hydrochloric acid"
)
```

Conduct a full analysis of this experiment, including

- exploratory data analysis;
- fitting a linear model, and conducting an F-test to compare to a model that explains variation using the six batches to the null model;
- usual linear model diagnostics;
- multiple comparisons of all pairwise differences between treatments.

Solution

- Two of the simplest ways of examining the data are to calculate basic descriptive statistics, e.g. the mean and standard deviation of the yield in each batch, and to plot the data in the different batches using a simple graphical display, e.g. a stripchart of the yields in each batch. Notice that in both `aggregate` and `stripchart` we use the formula `yield ~ batch`. This formula splits the data into groups defined by batch.

```
aggregate(yield ~ batch, data = napblack, FUN = function(x) c(mean = mean(x),
                                                                st.dev = sd(x)))
```

```
##   batch yield.mean yield.st.dev
## 1     1    105.00     63.05
## 2     2    128.00     33.28
## 3     3    164.00     37.98
## 4     4     98.00     68.70
## 5     5    200.00     50.00
## 6     6     70.00     31.02
```

```
boxplot(yield ~ batch, data = napblack)
```



Figure 2.2: Naphthalene black experiment: distributions of dye yields from the six batches.

Notice that even within any particular batch, the number of grams of standard dyestuff colour determined by the dye trial varies from observation to observation. This *within-group* variation is considered to be random or residual variation. This cannot be explained by any differences between batches. However, a second source of variation in the overall data set can be explained by variation between the batches, i.e. between the different batch means themselves. We can see from the box plots (Figure 2.2) and the mean yields in each batch that observations from batch number five appear to have given higher yields (in grams of standard colour) than those from the other batches.

- b. When we fit linear models and compare them using analysis of variance (ANOVA), it enables us to decide whether the differences that seem to be evident in these simple plots and descriptive statistics are statistically significant or whether this kind of variation could have arisen by chance, even though there are no real differences between the batches.

An ANOVA table may be used to compare a linear model including differences between the batches to the null model. The linear model we will fit is a simple unit-treatment model:

$$Y_{ij} = \mu + \tau_i + \varepsilon_{ij}, \quad i = 1, \dots, 6; j = 1, \dots, 5, \quad (2.14)$$

where Y_{ij} is the response obtained from the j th repetition of the i th batch, μ is a constant term, τ_i is the expected effect due to the observation being in the k th batch ($k = 1, \dots, 5$) and ε_{ij} are the random errors.

A test of the hypothesis that the group means are all equal is equivalent to a test that the τ_i are all equal to 0 ($H_0 : \tau_1 = \tau_2 = \dots = \tau_6 = 0$). We can use `lm` to fit model (2.14), and `anova` to test the hypothesis. Before we fit the linear model, we need to make sure `batch` has type `factor`⁹.

```
napblack$batch <- as.factor(napblack$batch)
napblack.lm <- lm(yield ~ batch, data = napblack)
anova(napblack.lm)
```

```
## Analysis of Variance Table
##
## Response: yield
##           Df Sum Sq Mean Sq F value Pr(>F)
## batch      5  56358   11272    4.6 0.0044 **
## Residuals 24  58830    2451
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

⁹Factors are variables in R which take on a limited number of different values (e.g. categorical variables). We need to define a categorical variable, like `batch` as a `factor` to ensure they are treated correctly by functions such as `lm`.

The p-value of 0.0044 indicates significant differences between at least two of the batch means. Therefore H_0 is rejected and a suitable multiple comparison test should be carried out.

- c. To perform our analysis, we have fitted a linear model. Therefore, we should use some plots of the residuals $y_{ij} - \hat{y}_{ij}$ to check the model assumptions, particularly that the errors are independently and identically normally distributed. The function `rstandard` which produces residuals which have been standardised to have variance equal to 1.

```
standres <- rstandard(napblack.lm)
fitted <- fitted(napblack.lm)
par(mfrow = c(1, 2), pty = "s")
with(napblack, {
  plot(batch, standres, xlab = "Batch", ylab = "Standardised residuals")
  plot(fitted, standres, xlab = "Fitted value", ylab = "Standardised residuals")
})
```



Figure 2.3: Residuals against batch (left) and fitted values (right) for the linear model fit to the naphthalene black data.

The plots (Figure 2.3) show no large standardised residuals (> 2 in absolute value¹⁰). While there is some evidence of unequal variation across batches, there is no obvious pattern with respect to fitted values (e.g. no “funnelling”).

We can also plot the standardised residuals against the quantiles of a standard normal distribution to assess the assumption of normality.

```
par(pty = "s")
qqnorm(standres, main = "")
```

¹⁰We would anticipate 95% of the standardised residuals to lie in $[-1.96, 1.96]$, as they will follow a standard normal distribution if the model assumptions are correct.

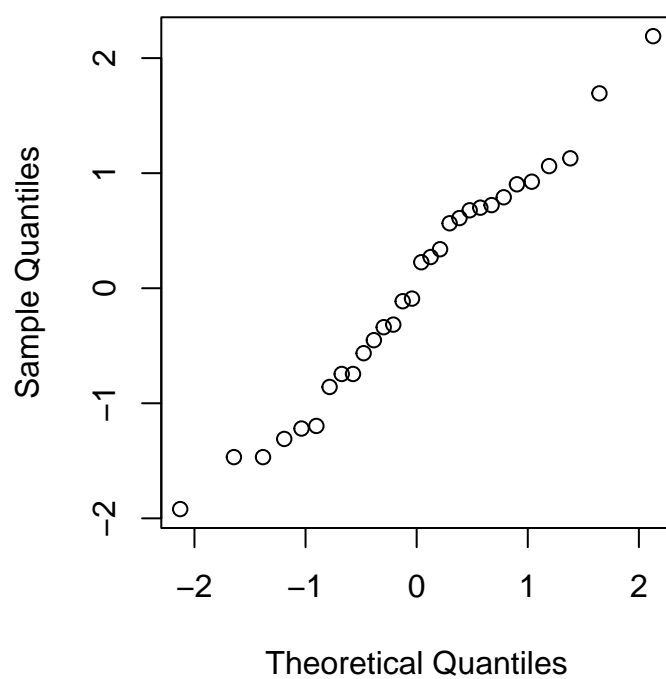


Figure 2.4: Normal probability plot for the standardised residuals for the linear model fit to the naphthalene black data.

The points lie quite well on a straight line (see Figure 2.4), suggesting the assumption of normality is valid. Overall, the residual plots look reasonable; some investigation of transformations to correct for non-constant variance could be investigated (see MATH2010/STAT6123).

- d. When a significant difference between the treatments has been indicated, the next stage is to try to determine which treatments differ. In some cases a specific difference is of interest, a control versus a new treatment for instance, in which case that difference could now be inspected. However, usually no specific differences are to be considered a priori, and *any* difference is of practical importance. A multiple comparison procedure is required to investigate all possible differences, which takes account of the number of possible differences available amongst the treatments (15 differences between the six batches here).

We will use Tukey's method for controlling the experiment-wise type I error rate, fixed here at 5%, as implemented by `emmeans`.

```
napblack.emm <- emmeans::emmeans(napblack.lm, 'batch')
pairs(napblack.emm)
```

```
## contrast estimate SE df t.ratio p.value
## 1 - 2          -23 31.3 24  -0.735  0.9755
## 1 - 3          -59 31.3 24  -1.884  0.4351
## 1 - 4           7 31.3 24   0.224  0.9999
## 1 - 5          -95 31.3 24  -3.034  0.0566
## 1 - 6           35 31.3 24   1.118  0.8692
## 2 - 3          -36 31.3 24  -1.150  0.8555
## 2 - 4           30 31.3 24   0.958  0.9266
## 2 - 5          -72 31.3 24  -2.299  0.2329
## 2 - 6           58 31.3 24   1.852  0.4535
## 3 - 4           66 31.3 24   2.108  0.3167
## 3 - 5          -36 31.3 24  -1.150  0.8555
## 3 - 6           94 31.3 24   3.002  0.0606
## 4 - 5          -102 31.3 24  -3.257  0.0348
## 4 - 6           28 31.3 24   0.894  0.9442
## 5 - 6          130 31.3 24   4.152  0.0043
##
## P value adjustment: tukey method for comparing a family of 6 estimates
```

We have two significant differences, between batches 4-5 and 5-6.

```
subset(transform(pairs(napblack.emm)), p.value < 0.05)
```

```
## contrast estimate SE df t.ratio p.value
## 13 4 - 5        -102 31.31 24  -3.257 0.034820
## 15 5 - 6         130 31.31 24   4.152 0.004295
```

4. (Adapted from Morris, 2011) Consider a completely randomised design

with $t = 5$ treatments and $n = 50$ units. The contrasts

$$\tau_2 - \tau_1, \quad \tau_3 - \tau_2, \quad \tau_4 - \tau_3, \tau_5 - \tau_4$$

are of primary interest to the experimenter.

- Find an allocation of the 50 units to the 5 treatments, i.e. find n_1, \dots, n_5 , that minimises the average variance of the corresponding contrast estimators.
- Fixing the proportions of experimental effort applied to each treatment to those found in part (a), i.e. to $w_i = n_i/50$, find the value of n required to make the ratio $T = |\mathbf{c}^T \boldsymbol{\tau}| / \sqrt{\text{var}(\widehat{\mathbf{c}^T \boldsymbol{\tau}})} = 2$ assuming a signal-to-noise ratio of 1.

Solution

- We can use the function `opt_ni` given in Section 2.8.1:

```
n <- 50
C <- matrix(
  c(
    -1, 1, 0, 0, 0,
    0, -1, 1, 0, 0,
    0, 0, -1, 1, 0,
    0, 0, 0, -1, 1
  ), nrow = 4, byrow = T
)
opt_ni(C, n)

## [1]  8.009 11.327 11.327 11.327  8.009
```

Rounding, we obtain a solution of the form $n_1 = n_5 = 8$, $n_2 = n_4 = 11$ and $n_3 = 12$. Any of n_2, n_3, n_4 may be rounded up to 12 to form a design with the same variance.

```
nv <- c(8, 11, 11, 11, 8)
crd_var(C, nv + c(0, 1, 0, 0, 0))
crd_var(C, nv + c(0, 0, 1, 0, 0))
crd_var(C, nv + c(0, 0, 0, 1, 0))

## [1] 0.7803
## [1] 0.7803
## [1] 0.7803
```

- The optimal ratios for each treatment from part (a) are $w_1 = w_5 = 0.1602$ and $w_2 = w_3 = w_4 = 0.2265$. Fixing these, we can use code from Section 2.8.2 to find the required value of n for each contrast.

```

nv <- NULL
for(i in 1:4) nv[i] <- opt_n(C[i, ], opt_ni(C, n) / n, 1, 2) # snr = 1, target = 2
nv

## [1] 42.63 35.31 35.31 42.63

```

Hence, we need $n = 43$ for to achieve $T = 2$ for the first and last contrasts, and $n = 36$ for the second and third. The differences are due to the different proportions w_i assumed for each treatment. To achieve $T = 2$ for all contrasts, we pick the larger number, $n = 43$.

Chapter 3

Blocking

The completely randomised design (CRD) works well when there is sufficient homogeneous experimental units to perform the whole experiment under the same, or very similar, conditions and there are no restrictions on the randomisation of treatments to units. The only systematic (non-random) differences in the observed responses result from differences between the treatments. While such designs are commonly and successfully used, especially in smaller experiments, their application can often be unrealistic or impractical in many settings.

A common way in which the CRD fails is a lack of sufficiently similar experimental units. If there are systematic differences between different batches, or **blocks** of units, these differences should be taken into account in both the allocation of treatments to units and the modelling of the resultant data. Otherwise, block-to-block differences may bias treatment comparisons and/or inflate our estimate of the background variability and hence reduce our ability to detect important treatment effects.

Example 3.1. Steel bar experiment (Morris, 2011, ch. 4)

Kocaoz et al. (2005) described an experiment to assess the strength of steel reinforcement bars from $t = 4$ coatings¹ (treatments). In total $n = 32$ different bars (units) were available, but the testing process meant sets of four bars were tested together. To account for potential test-specific features (e.g. environmental or operational), these different test sets were assumed to form $b = 8$ blocks of size $k = 4$. The data are shown in Table 3.1 below.

```
bar <- data.frame(coating = rep(factor(1:4), 8),  
                  block = rep(factor(1:8), rep(4, 8)),  
                  strength = c(136, 147, 138, 149, 136, 143, 122, 153, 150, 142, 131, 136,  
                               155, 148, 130, 129, 145, 149, 136, 139, 150, 149, 147, 144,
```

¹The four coatings were all made from Engineering Thermoplastic Polyurethane (ETPU); coating one was solely made from ETPU, coatings 2-4 had additional glass fibre, carbon fibre or aramid fibre added, respectively.

Table 3.1: Steel bar experiment: tensile strength values (kiliograms per square inch, ksi) from steel bars with four different coatings.

Block	Coating 1	Coating 2	Coating 3	Coating 4
1	136	147	138	149
2	136	143	122	153
3	150	142	131	136
4	155	148	130	129
5	145	149	136	139
6	150	149	147	144
7	147	150	125	140
8	148	149	118	145

```

                                147, 150, 125, 140, 148, 149, 118, 145)
                                )
knitr::kable(
  tidyr::pivot_wider(bar, names_from = coating, values_from = strength),
  col.names = c("Block", paste("Coating", 1:4)),
  caption = "Steel bar experiment: tensile strength values (kiliograms per square inch, ksi)"
)

```

Here, each block has size 4, which is equal to the number of treatments in the experiment, and each treatment is applied in each block. This is an example of a **randomised complete block design**.

We can study the data graphically, plotting by treatment and by block.

```

boxplot(strength ~ block, data = bar)
boxplot(strength ~ coating, data = bar)

```

The box plots within each plot in Figure 3.1 are comparable, as every treatment has occurred with every block the same number of times (once). For example, when we compare the box plots for treatments 1 and 3, we know each of them display one observation from each block. Therefore, differences between treatments will not be influenced by large differences between blocks. This **balance** makes our analysis more straightforward. By eye, it appears here there may be differences between both coating 3 and the other three coatings.

Example 3.2. Tyre experiment (Wu and Hamada, 2009, ch. 3)

Davies (1954), p.200, examined the effect of $t = 4$ different rubber compounds (treatments) on the lifetime of a tyre. Each tyre is only large enough to split into $k = 3$ segments whilst still containing a representative amount of each compound. When tested, each tyre is subjected to the same road conditions, and hence is treated as a block. A design with $b = 4$ blocks was used, as displayed in Table 3.2.



Figure 3.1: Steel bar experiment: distributions of tensile strength (ksi) from the eight blocks (top) and the four coatings (bottom).

Table 3.2: Tyre experiment: relative wear measurements (unitless) from tires made with four different rubber compounds.

Block	Compound 1	Compound 2	Compound 3	Compound 4
1	238	238	279	
2	196	213		308
3	254		334	367
4		312	421	412

```
tyre <- data.frame(compound = as.factor(c(1, 2, 3, 1, 2, 4, 1, 3, 4, 2, 3, 4)),
                  block = rep(factor(1:4), rep(3, 4)),
                  wear = c(238, 238, 279, 196, 213, 308, 254, 334, 367, 312, 421, 412),
                  )
options(knitr.kable.NA = '')
knitr::kable(
  tidyr::pivot_wider(tyre, names_from = compound, values_from = wear),
  col.names = c("Block", paste("Compound", 1:4)),
  caption = "Tyre experiment: relative wear measurements (unitless) from tires made with four different rubber compounds"
)
```

Here, each block has size $k = 3$, which is smaller than the number of treatments ($t = 4$). Hence, each block cannot contain an application of each treatment. This is an example of an **incomplete block design**.

Graphical exploration of the data is a little more problematic in this example. As each treatment does not occur in each block, box plots such as Figure 3.2 are not as informative. Do compounds three and four have higher average wear because they were the only compounds to both occur in blocks 3 and 4? Or do blocks 3 and 4 have a higher mean because they contain both compounds 3 and 4? The design cannot help us entirely disentangle the impact of blocks and treatments². In our modelling, we will assume variation should first be described by blocks (which are generally fixed aspects of the experiment) and then treatments (which are more directly under the experimenter's control).

```
boxplot(wear ~ block, data = tyre)
boxplot(wear ~ compound, data = tyre)
```

3.1 Unit-block-treatment model

If n_{ij} is the number of times treatment j occurs in block i , a common statistical model to describe data from a blocked experiment has the form

²This is our first example of (partial) confounding, which we will see again in Chapters 5 and 6

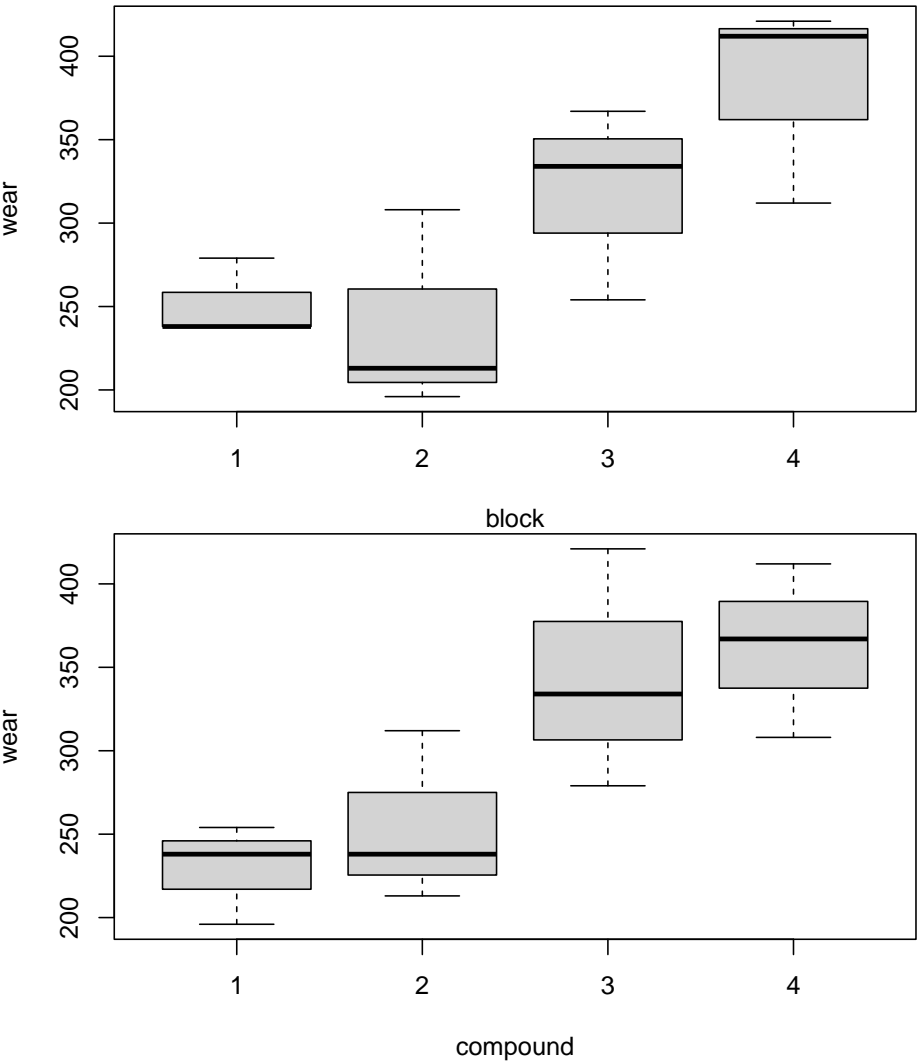


Figure 3.2: Tyre experiment: distributions of wear from the four blocks (top) and the four compounds (bottom).

$$y_{ijl} = \mu + \beta_i + \tau_j + \varepsilon_{ijl}, \quad i = 1, \dots, b; j = 1, \dots, t; l = 1, \dots, n_{ij}, \quad (3.1)$$

where y_{ijl} is the response from the l th application of the j th treatment in the i th block, μ is a constant parameter, β_i is the effect of the i th block, τ_j is the effect of treatment j , and $\varepsilon_{ijl} \sim N(0, \sigma^2)$ are once again random individual effects from each experimental unit, assumed independent. The total number of runs in the experiment is given by $n = \sum_{i=1}^b \sum_{j=1}^t n_{ij}$.

For Example 3.1, there are $t = 4$ experiments, $b = 8$ blocks and each treatment occurs once in each block, so $n_{ij} = 1$ for all i, j . In Example 3.2, there are again $t = 4$ treatments but now only $b = 4$ blocks and not every treatment occurs in every block. In fact, we have $n_{11} = n_{12} = n_{13} = 1$, $n_{14} = 0$, $n_{21} = n_{22} = n_{24} = 1$, $n_{23} = 0$, $n_{31} = n_{33} = n_{34} = 1$, $n_{32} = 0$, $n_{41} = 0$ and $n_{42} = n_{43} = n_{44} = 1$.

Writing model (3.1) in matrix form as a partitioned linear model, we obtain

$$\mathbf{y} = \mu \mathbf{1}_n + X_1 \boldsymbol{\beta} + X_2 \boldsymbol{\tau} + \boldsymbol{\varepsilon}, \quad (3.2)$$

with \mathbf{y} the n -vector of responses, X_1 and X_2 $n \times b$ and $n \times t$ model matrices for blocks and treatments, respectively, $\boldsymbol{\beta} = (\beta_1, \dots, \beta_b)^T$, $\boldsymbol{\tau} = (\tau_1, \dots, \tau_t)^T$ and $\boldsymbol{\varepsilon}$ the n -vector of errors.

In equation (3.2), assuming without loss of generality that runs of the experiment are ordered by block, the matrix X_1 has the form

$$X_1 = \bigoplus_{i=1}^b \mathbf{1}_{k_i} = \begin{bmatrix} \mathbf{1}_{k_1} & \mathbf{0}_{k_1} & \cdots & \mathbf{0}_{k_1} \\ \mathbf{0}_{k_2} & \mathbf{1}_{k_2} & \cdots & \mathbf{0}_{k_2} \\ \vdots & & \ddots & \vdots \\ \mathbf{0}_{k_b} & \mathbf{0}_{k_b} & \cdots & \mathbf{1}_{k_b} \end{bmatrix},$$

where $k_i = \sum_{j=1}^t n_{ij}$, the number of units in the i th block. The structure of matrix X_2 is harder to describe so distinctly, but each row includes a single non-zero entry, equal to one, indicating which treatment was applied in that run of the experiment. The first k_1 rows correspond to block 1, the second k_2 to block 2, and so on. We will see special cases later.

3.2 Normal equations

Writing as a partitioned model $\mathbf{y} = W\boldsymbol{\alpha} + \boldsymbol{\varepsilon}$, with $W = [\mathbf{1}|X_1|X_2]$ and $\boldsymbol{\alpha}^T = [\mu|\boldsymbol{\beta}^T|\boldsymbol{\tau}^T]$, the least squares normal equations

$$W^T W \hat{\boldsymbol{\alpha}} = W^T \mathbf{y} \quad (3.3)$$

can be written as a set of three matrix equations:

$$n\hat{\mu} + \mathbf{1}_n^T X_1 \hat{\beta} + \mathbf{1}_n^T X_2 \hat{\tau} = \mathbf{1}_n^T \mathbf{y}, \quad (3.4)$$

$$X_1^T \mathbf{1}_n \hat{\mu} + X_1^T X_1 \hat{\beta} + X_1^T X_2 \hat{\tau} = X_1^T \mathbf{y}, \quad (3.5)$$

$$X_2^T \mathbf{1}_n \hat{\mu} + X_2^T X_1 \hat{\beta} + X_2^T X_2 \hat{\tau} = X_2^T \mathbf{y}. \quad (3.6)$$

$$(3.7)$$

Above, the matrices $X_1^T X_1 = \text{diag}(k_1, \dots, k_b)$ and $X_2^T X_2 = \text{diag}(n_1, \dots, n_t)$ have simple forms as diagonal matrices with entries equal to the size of each block and the number of replications of each treatment, respectively.

The $t \times b$ matrix $N = X_2^T X_1$ is particularly important in block designs, and is called the **incidence** matrix. Each of the i th row of N indicates in which blocks the i th treatment occurs.

We can eliminate the explicit dependence on μ and β to find reduced normal equations for τ by multiplying the middle equation by $X_2^T X_1 (X_1^T X_1)^{-1}$:

$$\begin{aligned} X_2^T X_1 (X_1^T X_1)^{-1} X_1^T \mathbf{1}_n \hat{\mu} + X_2^T X_1 (X_1^T X_1)^{-1} X_1^T X_1 \hat{\beta} + X_2^T X_1 (X_1^T X_1)^{-1} X_1^T X_2 \hat{\tau} \\ = X_2^T X_1 (X_1^T X_1)^{-1} X_1^T \mathbf{1}_n \hat{\mu} + X_2^T X_1 \hat{\beta} + X_2^T X_1 (X_1^T X_1)^{-1} X_1^T X_2 \hat{\tau} \\ = X_2^T X_1 (X_1^T X_1)^{-1} X_1^T \mathbf{y} \end{aligned} \quad (3.8)$$

and subtracting from the final equation:

$$\begin{aligned} X_2^T (\mathbf{1}_n - X_1 (X_1^T X_1)^{-1} X_1^T \mathbf{1}_n) \hat{\mu} + (X_2^T X_1 - X_2^T X_1 (X_1^T X_1)^{-1} X_1^T X_1) \hat{\beta} \\ + X_2^T (I_n - X_1 (X_1^T X_1)^{-1} X_1^T) X_2 \tau \\ = X_2^T (I_n - X_1 (X_1^T X_1)^{-1} X_1^T) \mathbf{y}. \end{aligned} \quad (3.9)$$

Clearly, a zero matrix is multiplying the block effects β . Also,

$$X_1 (X_1^T X_1)^{-1} X_1^T \mathbf{1}_n = \mathbf{1}_n,$$

as

$$X_1 (X_1^T X_1)^{-1} = \bigoplus_{i=1}^b \frac{1}{k_i} \mathbf{1}_{k_i} = \begin{bmatrix} \frac{1}{k_1} \mathbf{1}_{k_1} & \mathbf{0}_{k_1} & \cdots & \mathbf{0}_{k_1} \\ \mathbf{0}_{k_2} & \frac{1}{k_2} \mathbf{1}_{k_2} & \cdots & \mathbf{0}_{k_2} \\ \vdots & \vdots & \ddots & \vdots \\ \mathbf{0}_{k_b} & \mathbf{0}_{k_b} & \cdots & \frac{1}{k_b} \mathbf{1}_{k_b} \end{bmatrix},$$

and hence

$$X_1(X_1^T X_1)^{-1} X_1^T = \bigoplus_{i=1}^b \frac{1}{k_i} J_{k_i} = \begin{bmatrix} \frac{1}{k_1} J_{k_1} & \mathbf{0}_{k_1 \times k_2} & \cdots & \mathbf{0}_{k_1 \times k_b} \\ \mathbf{0}_{k_2 \times k_1} & \frac{1}{k_2} J_{k_2} & \cdots & \mathbf{0}_{k_2 \times k_b} \\ \vdots & & \ddots & \vdots \\ \mathbf{0}_{k_b \times k_1} & \mathbf{0}_{k_b \times k_2} & \cdots & \frac{1}{k_b} J_{k_b} \end{bmatrix}.$$

Writing $H_1 = X_1(X_1^T X_1)^{-1} X_1^T$, we then get the reduced normal equations for τ :

$$X_2^T (I_n - H_1) X_2 \tau = X_2^T (I_n - H_1) \mathbf{y}. \quad (3.10)$$

We can demonstrate the form of these matrices through our two examples.

For Example 3.1:

```
one <- rep(1, 4)
X1 <- kronecker(diag(1, nrow = 8), one)
X2 <- diag(1, nrow = 4)
X2 <- do.call("rbind", replicate(8, X2, simplify = FALSE))
#incidence matrix
N <- t(X2) %*% X1
X1tX1 <- t(X1) %*% X1 # diagonal
X2tX2 <- t(X2) %*% X2 # diagonal
H1 <- X1 %*% solve(t(X1) %*% X1) %*% t(X1)
ones <- H1 %*% rep(1, 32) # H1 times vector of 1s is also a vector of 1s
A <- t(X2) %*% X2 - t(X2) %*% H1 %*% X2 # X2t(I - H1)X2
qr(A)$rank # rank 3
X2tH1 <- t(X2) %*% H1 # adjustment to y
W <- cbind(ones, X1, X2) # overall model matrix
qr(W)$rank # rank 11 (t+b - 1)
```

For Example 3.2:

```
one <- rep(1, 3)
X1 <- kronecker(diag(1, nrow = 4), one)
X2 <- matrix(
  c(1, 0, 0, 0,
    0, 1, 0, 0,
    0, 0, 1, 0,
    1, 0, 0, 0,
    0, 1, 0, 0,
    0, 0, 0, 1,
    1, 0, 0, 0,
    0, 0, 1, 0,
    0, 0, 0, 1,
```

```

      0, 1, 0, 0,
      0, 0, 1, 0,
      0, 0, 0, 1), nrow = 12, byrow = T
)
#incidence matrix
N <- t(X2) %*% X1
X1tX1 <- t(X1) %*% X1 # diagonal
X2tX2 <- t(X2) %*% X2 # diagonal
H1 <- X1 %*% solve(t(X1) %*% X1) %*% t(X1)
ones <- H1 %*% rep(1, 12) # H1 times vector of 1s is also a vector of 1s
A <- t(X2) %*% X2 - t(X2) %*% H1 %*% X2 # X2t(I - H1)X2
qr(A)$rank # rank 3
X2tH1 <- t(X2) %*% H1 # adjustment to y
W <- cbind(ones, X1, X2) # overall model matrix
qr(W)$rank # rank 7 (t+b - 1)

```

Notice that if we write $X_{2|1} = (I_n - H_1)X_2$, then the reduced normal equations become

$$X_{2|1}^T X_{2|1} \boldsymbol{\tau} = X_{2|1}^T \mathbf{y},$$

which have the same form as the CRD in Chapter 2 albeit with a different $X_{2|1}$ matrix as we are adjusting for more complex nuisance parameters.

In general, the solution of these equations will depend on the exact form of the design. For the randomised complete block design, the solution turns out to be straightforward (see Section [@ref\(#rcdb\)](#) below). By default, to fit model (3.2), the `lm` function in R applies the constraint $\tau_t = \beta_b = 0$, and removes the corresponding columns from X_1 and X_2 , to leave a W matrix with full column rank. Clearly, this solution is not unique but, as with CRDs, we will identify uniquely estimatable combinations of the model parameters (and use `emmeans` to extract these estimates from an `lm` object).

3.3 Analysis of variance

As was the case with the CRD, it can be shown that any solution to the normal equations (3.3) will produce a unique solution to $\widehat{W\alpha}$, and hence a unique analysis of variance decomposition can be obtained.

For a block experiment, the ANOVA table is comparing the full model (3.2), the model containing the block effects

$$\mathbf{y} = \mu \mathbf{1} + X_1 \boldsymbol{\beta} + \boldsymbol{\varepsilon} \quad (3.11)$$

and the null model

$$\mathbf{y} = \mu \mathbf{1} + \boldsymbol{\varepsilon}, \quad (3.12)$$

and has the form:

Source	Degrees of freedom	Sums of squares	Mean square
Blocks	$b - 1$	RSS (3.12) - RSS (3.11)	
Treatments	$t - 1$	RSS (3.11) - RSS (3.2)	[RSS (3.11) - RSS (3.2)] / $(t - 1)$
Residual	$n - b - t + 1$	RSS (3.2)	RSS (3.2) / $(n - b - t + 1)$
Total	$n - 1$	RSS (3.12)	

We test the hypothesis $H_0 : \tau_1 = \cdots = \tau_t = 0$ at the $100\alpha\%$ significance level by comparing the ratio of treatment and residual mean squares to the $1 - \alpha$ quantile of an F distribution with $t - 1$ and $n - b - t + 1$ degrees of freedom.

For Example 3.1, we obtain the following ANOVA.

```
bar.lm <- lm(strength ~ block + coating, data = bar)
anova(bar.lm)
```

```
## Analysis of Variance Table
##
## Response: strength
##           Df Sum Sq Mean Sq F value Pr(>F)
## block      7     215      31    0.55 0.7903
## coating    3    1310     437    7.75 0.0011 **
## Residuals 21    1184      56
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Clearly, the null hypothesis of no treatment effect is rejected. The `anova` function also compares the block mean square to the residual mean square to perform a test of the hypothesis $H_0 : \beta_1 = \cdots = \beta_b = 0$. This is not a hypothesis that should usually be tested. The blocks are a nuisance factor and are generally a feature of the experimental process that has not been subject to randomisation; we are not interested in testing for block-to-block differences.³

For Example 3.2, we get the ANOVA table:

```
tyre.lm <- lm(wear ~ block + compound, data = tyre)
anova(tyre.lm)
```

³R and `anova` don't, of course, know that this is a block design or that a blocking factor is being tested.

```
## Analysis of Variance Table
##
## Response: wear
##           Df Sum Sq Mean Sq F value    Pr(>F)
## block      3  39123   13041    37.2 0.00076 ***
## compound   3  20729    6910    19.7 0.00335 **
## Residuals  5   1751     350
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Again, the null hypothesis is rejected, and hence we should investigate which tyre compounds differ in their mean response.

The residual mean square for model (3.2) also provides an unbiased estimate, s^2 , of σ^2 , the variability of the ε_{ijl} , *assuming the unit-block-treatment model is correct*.

```
bar.s2 <- summary(bar.lm)$sigma^2
tyre.s2 <- summary(tyre.lm)$sigma^2
```

For Example 3.1, $s^2 = 56.3869$ and for Example 3.2, $s^2 = 350.1833$.

3.4 Randomised complete block designs

A randomised complete block design (RCBD) has each treatment replicated exactly once in each block, that is $n_{ij} = 1$ for $i = 1, \dots, b; j = 1, \dots, t$. Therefore each block has common size $k_1 = \dots = k_b = t$. The t treatments are randomised to the t units in each block. We can drop the index l from our unit-block-treatment model, as every treatment is replicated just once:

$$y_{ij} = \mu + \beta_i + \tau_j + \varepsilon_{ij}, \quad i = 1, \dots, b; j = 1, \dots, t.$$

For an RCBD, the matrix $X_{2|1}$ has the form

$$\begin{aligned} X_{2|1} &= (I_n - H_1)X_2 \\ &= X_2 - H_1X_2 \\ &= X_2 - \frac{1}{t}J_{n \times t}, \end{aligned} \tag{3.13}$$

following from the fact that

$$H_1 X_2 = X_1 (X_1^T X_1)^{-1} X_1^T X_2 \quad (3.14)$$

$$= \frac{1}{t} X_1 X_1^T X_2 \quad (3.15)$$

$$= \frac{1}{t} X_1 N^T \quad (3.16)$$

$$= \frac{1}{t} X_1 J_{b \times t} \quad (3.17)$$

$$= \frac{1}{t} J_{n \times t}, \quad (3.18)$$

as for a RCBD $X_1^T X_1 = \text{diag}(k_1, \dots, k_b) = tI_b$ and $X_2^T X_1 = N = J_{t \times b}$.

Comparing (3.13) to the form of $X_{2|1}$ for a CRD, equation (2.6), we see that for the RCBD, $X_{2|1}$ has the same form as a CRD with b replicates of each treatment (that is, $n_i = b$ for $i = 1, \dots, t$). This is a powerful result, as it tell us

- The reduced normal equations for the RCBD take the same form as for the CRD,

$$\hat{\tau}_j - \hat{\tau}_w = \bar{y}_{.j} - \bar{y}_{..},$$

with $\hat{\tau}_w = \frac{1}{t} \sum_{j=1}^t \hat{\tau}_j$, $\bar{y}_{.j} = \frac{1}{b} \sum_{i=1}^b y_{ij}$ and $\bar{y}_{..} = \frac{1}{n} \sum_{i=1}^b \sum_{j=1}^t y_{ij}$. Hence, as with a CRD, we can estimate any contrast $\mathbf{c}^T \boldsymbol{\tau}$, having $\sum_{j=1}^t c_j = 0$, with estimator

$$\widehat{\mathbf{c}^T \boldsymbol{\tau}} = \sum_{j=1}^t c_j \bar{y}_{.j}.$$

Hence, the **point estimate** for a contrast $\mathbf{c}^T \boldsymbol{\tau}$ is exactly the same as would be obtained by ignoring blocks and treating the experiment as a CRD with $n = bt$ and $n_i = b$, for $i = 1, \dots, t$.

- Inference for a contrast takes exactly the same form as for a CRD (Section 2.5), with in particular:

$$\text{var} \left(\widehat{\mathbf{c}^T \boldsymbol{\tau}} \right) = \frac{\sigma^2}{b} \sum_{j=1}^t c_j^2,$$

and

$$\widehat{\mathbf{c}^T \boldsymbol{\tau}} \sim N \left(\mathbf{c}^T \boldsymbol{\tau}, \frac{\sigma^2}{b} \sum_{j=1}^t c_j^2 \right).$$

Although these equations have the same form as for a CRD, note that σ^2 is representing different quantities in each case.

- In a CRD, σ^2 is the uncontrolled variation in the response *among all experimental units*.
- In a RCBD, σ^2 is the uncontrolled variation in the response *among all units within a common block*.

Block-to-block differences are modelled via inclusion of the block effects β_i in the model, and hence if blocking is effective, we would expect σ^2 from a RCBD to be substantially smaller than from a corresponding CRD with $n_i = b$.

Example 3.1 is a RCBD. We can estimate the contrasts

$$\tau_1 - \tau_2, \tau_1 - \tau_3, \tau_1 - \tau_4$$

between coatings⁴ using `emmeans`.

```
bar.emm <- emmeans::emmeans(bar.lm, ~ coating)
contrastv1.emmc <- function(levs)
  data.frame('t1 v t2' = c(1, -1, 0, 0), 't1 v t3' = c(1, 0, -1, 0),
    't1 v t4' = c(1, 0, 0, -1))
emmeans::contrast(bar.emm, 'contrastv1')

## contrast estimate SE df t.ratio p.value
## t1.v.t2 -1.25 3.75 21 -0.333 0.7425
## t1.v.t3 15.00 3.75 21 3.995 0.0007
## t1.v.t4 4.00 3.75 21 1.065 0.2988
##
## Results are averaged over the levels of: block
```

It is important to once again adjust for multiple comparisons. Here we can use a Bonferroni adjustment, and multiply each p-value by the number of tests (3). We obtain p-values of 1 (coating 1 versus 2), 0.002 (1 versus 3) and 0.8964 (2 versus 3). Hence, there is a significant difference between coatings 1 and 3, with $H_0 : \tau_1 = \tau_3$ rejected at the 1% significant level.

We can demonstrate the equivalence of the contrast point estimates between a RCBD and a CRD by fitting a unit-treatment model that ignores blocks:

```
bar_crd.lm <- lm(strength ~ coating, data = bar)
bar_crd.emm <- emmeans::emmeans(bar_crd.lm, ~ coating)
emmeans::contrast(bar_crd.emm, 'contrastv1')

## contrast estimate SE df t.ratio p.value
## t1.v.t2 -1.25 3.54 28 -0.354 0.7263
```

⁴These contrasts measure the difference between the coating only made from ETPU and the three coatings with added fibres.

```
## t1.v.t3      15.00 3.54 28    4.243  0.0002
## t1.v.t4      4.00 3.54 28    1.132  0.2674
crd.s2 <- summary(bar_crd.lm)$sigma^2
rcbd.s2 <- summary(bar.lm)$sigma^2
```

As expected the point estimates of the three contrasts are identical. In this case, the standard error of each contrast is actually smaller assuming a CRD without blocks, suggesting block-to-block differences were actually small here (further evidence is provided by the small block sums of squares in the ANOVA table). Here the estimate of σ from the RCB is $s_{RCB} = 7.5091$ and from the CRD is $s_{CRD} = 7.0698$, so for this example the unit-to-unit variation within and between blocks is not so different, and actually estimated to be slightly smaller in the CRD⁵.

3.5 Orthogonal blocking

The equality of the point estimates from the RCB and the CRD is a consequence of the block and treatment parameters in model (3.1) being **orthogonal**. That is, the least squares estimators for β and τ are independent in the sense that the estimators obtained from model (3.2) are the same as those obtained from the sub-models

$$\mathbf{y} = \mu \mathbf{1}_n + X_1 \beta + \varepsilon,$$

and

$$\mathbf{y} = \mu \mathbf{1}_n + X_2 \tau + \varepsilon.$$

That is, the presence or absence of the block parameters does not affect the estimator of the treatment parameters (and vice versa).

A condition for β and τ to be estimated orthogonally can be derived from normal equations (3.4) - (3.5). Firstly, we premultiply (3.4) by $\frac{1}{n} X_1^T \mathbf{1}_n$ and subtract it from (3.5):

⁵Of course, the CRD has seven more degrees of freedom for estimating σ^2 as block effects do not require estimation.

$$\begin{aligned}
& (X_1^T \mathbf{1}_n - X_1^T \mathbf{1}_n) \hat{\mu} + \left(X_1^T X_1 - \frac{1}{n} X_1^T \mathbf{1}_n \mathbf{1}_n^T X_1 \right) \hat{\beta} + \left(X_1^T X_2 - \frac{1}{n} X_1^T \mathbf{1}_n \mathbf{1}_n^T X_2 \right) \hat{\tau} \\
&= X_1^T \left(I_n - \frac{1}{n} J_n \right) X_1 \hat{\beta} + X_1^T \left(I_n - \frac{1}{n} J_n \right) X_2 \hat{\tau} \\
&= X_1^T \left(I_n - \frac{1}{n} J_n \right) \mathbf{y}. \tag{3.19}
\end{aligned}$$

Secondly, we premultiply (3.4) by $\frac{1}{n} X_2^T \mathbf{1}_n$ and subtract it from (3.6):

$$X_2^T \left(I_n - \frac{1}{n} J_n \right) X_1 \hat{\beta} + X_2^T \left(I_n - \frac{1}{n} J_n \right) X_2 \hat{\tau} = X_2^T \left(I_n - \frac{1}{n} J_n \right) \mathbf{y}. \tag{3.20}$$

For equations (3.19) and (3.20) to be independent, we require

$$X_2^T \left(I_n - \frac{1}{n} J_n \right) X_1 = \mathbf{0}_{t \times b}.$$

Hence, we obtain the following condition on the incidence matrix $N = X_2^T X_1$ for a block design to be orthogonal:

$$N = \frac{1}{n} X_2^T J_n X_1 \tag{3.21}$$

$$= \frac{1}{n} \mathbf{n} \mathbf{k}^T, \tag{3.22}$$

where $\mathbf{n}^T = (n_1, \dots, n_t)$ is the vector of treatment replications and $\mathbf{k}^T = (k_1, \dots, k_b)$ is the vector of block sizes.

The most common orthogonal block design for unstructured treatments is the RCBD, which has $n = bt$, $\mathbf{n} = b \mathbf{1}_t$, $\mathbf{k} = t \mathbf{1}_b$, and

$$N = J_{t \times b} = \frac{1}{bt} \mathbf{n} \mathbf{k}^T. \tag{3.23}$$

Hence, the condition for orthogonality is met. In an orthogonal design, such as a RCBD, all information about the treatment comparisons is contained in comparisons made within blocks. For more complex blocking structures, such as incomplete block designs, this is not the case. We shall see orthogonal blocking again in Chapter 5.

3.6 Balanced incomplete block designs

When the blocks sizes are less than the number of treatments, i.e. $k_i < t$ for all $i = 1, \dots, b$, by necessity the design is incomplete, in that not all treatments can be allocated to every block. We will restrict ourselves now to considering binary designs with common block size $k < t$. In a binary design, each treatment occurs within a block either 0 or 1 times ($n_{ij} = 0$ or $n_{ij} = 1$).

Example 3.2 is an example of an incomplete design with $k = 3 < t = 4$. For incomplete designs, it is often useful to study the *treatment concurrence matrix*, given by NN^T .

```
N <- matrix(
  c(1, 1, 1, 0,
    1, 1, 0, 1,
    1, 0, 1, 1,
    0, 1, 1, 1),
  nrow = 4, byrow = T
)
N %*% t(N)
```

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

This matrix has the number of treatment replications, n_j , on the diagonal and the off-diagonal elements are equal to the number of blocks within which each pair of treatments occurs together. We will denote as λ_{ij} the number of blocks that contain both treatment i and treatment j ($i \neq j$). For Example 3.2, $\lambda_{ij} = 2$ for all $i, j = 1, \dots, 4$; that is, each pair of treatments occurs together in two blocks.

Definition 3.1. A **balanced incomplete block design** (BIBD) is an incomplete block design with $k < t$ that meets three requirements:

1. The design is binary.
2. Each treatment is applied to a unit in the same number of blocks. It follows that the common number of units applied to each treatment must be $r = n_j = bk/t$ ($j = 1, \dots, t$), where $n = bk$. (Sometimes referred to as first-order balance).
3. Each pair of treatments is applied to two units in the same number of blocks, that is $\lambda_{ij} = \lambda$. (Sometimes referred to as second-order balance).

In fact, we can deduce that $\lambda(t-1) = r(k-1)$. To see this, focus on treatment 1. This treatment occurs in r blocks, and in each of these blocks, it occurs together with $k-1$ other treatments. But also, treatment 1 occurs

λ times with each of the other $t - 1$ treatments. Hence $\lambda(t - 1) = r(k - 1)$, or $\lambda = r(k - 1)/(t - 1)$.

The design in Example 3.2 is a BIBD with $b = 4$, $k = 3$, $t = 4$, $r = 4 \times 3/4 = 3$, $\lambda = 3 \times (3 - 1)/(4 - 1) = 2$.

3.6.1 Construction of BIBDs

BIBDs do not exist for all combinations of values of t , k and b . In particular, we must ensure

- $r = bk/t$ is integer, and
- $\lambda = r(k - 1)/(t - 1)$ is integer.

In general, we can always construct a BIBD for t treatments in $b = \binom{t}{k}$ blocks of size k , although it may not be the smallest possible BIBD. Each of the possible choices of k treatments from the total t forms one block. Such a design will have $r = \binom{t-1}{k-1}$ and $\lambda = \binom{t-2}{k-2}$. The design in Example 3.2 was constructed this way, with $b = 4$, $r = 3$ and $\lambda = 2$.

Sometimes, smaller BIBDs that satisfy the two conditions above can be constructed. Finding these designs is a combinatorial problem, and tables of designs are available in the literature⁶. A large collection of BIBDs has also been catalogued in the R package `ibd`. The function `bibd` generates BIBDs for given values of t , b , r , k and λ , or returns a message that a design is not available for those values. We can use the function to find the design used in Example 3.2.

```
tyre.bibd <- ibd::bibd(v = 4, b = 4, r = 3, k = 3, lambda = 2) # note, v is the notation for the
tyre.bibd$N # incidence matrix
```

```
##      [,1] [,2] [,3] [,4]
## [1,]    1    1    1    0
## [2,]    0    1    1    1
## [3,]    1    0    1    1
## [4,]    1    1    0    1
```

We can also use the package to find a design for bigger experiments, for example $t = 8$ treatments in $b = 14$ blocks of size $k = 4$. Here, $r = 14 \times 4/8 = 7$ and $\lambda = 7 \times 3/7 = 3$.

```
larger.bibd <- ibd::bibd(v = 8, b = 14, r = 7, k = 4, lambda = 3)
larger.bibd$N
```

```
##      [,1] [,2] [,3] [,4] [,5] [,6] [,7] [,8] [,9] [,10] [,11] [,12] [,13] [,14]
## [1,]    1    1    0    1    0    1    0    1    0    0    0    1    1    0
## [2,]    0    0    0    0    0    0    0    1    1    1    1    1    1    1
## [3,]    1    0    0    1    0    1    1    0    0    1    1    0    0    1
```

⁶See Cochran and Cox (1957) and Fisher and Yates (1963)

## [4,]	1	1	0	0	1	0	1	0	1	0	0	0	1	1
## [5,]	1	1	1	0	1	0	0	0	0	1	1	1	0	0
## [6,]	0	1	1	0	0	1	1	1	1	1	0	0	0	0
## [7,]	0	0	1	1	1	0	1	1	0	0	1	0	1	0
## [8,]	0	0	1	1	1	1	0	0	1	0	0	1	0	1

Although larger than the examples we have considered before, this design is small compared to the design that would be obtained from the naive construction above with $b = \binom{t}{k} = \binom{8}{4} = 70$ blocks.

3.6.2 Reduced normal equations

It can be shown that the reduced normal equations (3.10) for a BIBD can be written as

$$\left(I_t - \frac{1}{t}J_t\right)\hat{\tau} = \frac{k}{\lambda t} \left(X_2^T - \frac{1}{k}NX_1^T\right)\mathbf{y}. \quad (3.24)$$

Equation (3.24) defines a series of t equations of the form

$$\begin{aligned} \hat{\tau}_j - \hat{\tau}_w &= \frac{k}{\lambda t} \left(\sum_{i=1}^b n_{ij}y_{ij} - \frac{1}{k} \sum_{i=1}^b n_{ij} \sum_{j=1}^t n_{ij}y_{ij} \right) \\ &= \frac{k}{\lambda t} q_j, \end{aligned}$$

with $q_j = \sum_{i=1}^b n_{ij}y_{ij} - \frac{1}{k} \sum_{i=1}^b n_{ij} \sum_{j=1}^t n_{ij}y_{ij}$.

Notice that unlike for the RCBD, the reduced normal equations for a BIBD do not correspond to the equations for a CRD. Although the first term in q_i is the sum of the responses for the j th treatment (mirroring the CRD), the second term is no longer the overall sum (or average) of the responses. In fact, for q_j this second term is an adjusted total, just involving observations from those blocks that contain treatment j .

3.6.3 Estimation and inference

As with the CRD and RCD we can estimate contrasts in the τ_i , with estimator

$$\widehat{\mathbf{c}^T \boldsymbol{\tau}} = \frac{k}{\lambda t} \sum_{j=1}^t c_j q_j.$$

Due to the form of the reduced normal equations for the BIBD, the estimator is no longer just a linear combination of the treatment means; q_j includes a term that adjusts for the blocks in which treatment j has occurred.

The simplest way to derive the variance of this estimator is to rewrite it in the form

$$\widehat{\mathbf{c}^T \boldsymbol{\tau}} = \frac{k}{\lambda t} \mathbf{c}^T \mathbf{q},$$

with $\mathbf{q} = \left(X_2^T - \frac{1}{k} N X_1^T\right) \mathbf{y}$. Then

$$\text{var} \left(\widehat{\mathbf{c}^T \boldsymbol{\tau}} \right) = \frac{k^2}{\lambda^2 t^2} \mathbf{c}^T \text{var}(\mathbf{q}) \mathbf{c}.$$

Recalling that NN^T is the treatment coincidence matrix, the variance-covariance matrix of \mathbf{q} is given by

$$\begin{aligned} \text{var}(\mathbf{q}) &= \left(X_2^T - \frac{1}{k} N X_1^T\right) \left(X_2 - \frac{1}{k} X_1 N\right) \sigma^2 \\ &= \sigma^2 \left\{ r I_t - \frac{1}{k} N N^T \right\} \\ &= \sigma^2 \left\{ r I_t - \frac{1}{k} [(r - \lambda) I_t + \lambda J_t] \right\} \\ &= \sigma^2 \left\{ \left(\frac{r(k - 1) + \lambda}{k} \right) I_t - \frac{\lambda}{k} J_t \right\} \\ &= \sigma^2 \left\{ \left(\frac{\lambda(t - 1) + \lambda}{k} \right) I_t - \frac{\lambda}{k} J_t \right\} \\ &= \sigma^2 \left\{ \frac{\lambda t}{k} I_t - \frac{\lambda}{k} J_t \right\}. \end{aligned}$$

Hence

$$\begin{aligned} \text{var} \left(\widehat{\mathbf{c}^T \boldsymbol{\tau}} \right) &= \frac{k^2}{\lambda^2 t^2} \mathbf{c}^T \left(\frac{\lambda t}{k} I_t - \frac{\lambda}{k} J_t \right) \mathbf{c} \sigma^2 \\ &= \frac{k \sigma^2}{\lambda t} \mathbf{c}^T \mathbf{c} \\ &= \frac{k \sigma^2}{\lambda t} \sum_{j=1}^t c_j^2, \end{aligned}$$

as $\mathbf{c}^T J_t = \mathbf{0}$ as $\sum_{j=1}^t c_j = 0$. The estimator is also unbiased ($E(\widehat{\mathbf{c}^T \boldsymbol{\tau}}) = \mathbf{c}^T \boldsymbol{\tau}$), and hence the sampling distribution, upon which inference can be based, is given by

$$\widehat{\mathbf{c}^T \boldsymbol{\tau}} \sim N \left(\mathbf{c}^T \boldsymbol{\tau}, \frac{k\sigma^2}{\lambda t} \sum_{j=1}^t c_j^2 \right).$$

Returning to Example 3.2, we can use these results to estimate all pairwise differences between the four treatments. Firstly, we directly calculate the q_j from treatment and block sums, using the incidence matrix.

```
trtsum <- aggregate(wear ~ compound, data = tyre, FUN = sum)[, 2]
blocksum <- aggregate(wear ~ block, data = tyre, FUN = sum)[, 2]
q <- trtsum - N %*% blocksum / 3
C <- matrix(
  c(1, -1, 0, 0,
    1, 0, -1, 0,
    1, 0, 0, -1,
    0, 1, -1, 0,
    0, 1, 0, -1,
    0, 0, 1, -1),
  ncol = 4, byrow = T
)
k <- 3; lambda <- 2; t <- 4
pe <- k * C %*% q / (lambda * t) # point estimates
se <- sqrt(2 * k * tyre.s2 / (lambda * t)) # st error (same for each contrast)
t.ratio <- pe / se
p.value <- 1 - ptukey(abs(t.ratio) * sqrt(2), 4, 5)
data.frame(Pair = c('1v2', '1v3', '1v4', '2v3', '2v4', '3v4'),
  Estimate = pe, St.err = se, t.ratio = t.ratio,
  p.value = p.value, reject = p.value < 0.05)
```

```
##   Pair Estimate St.err t.ratio p.value reject
## 1  1v2   -4.375  16.21  -0.270 0.992273  FALSE
## 2  1v3  -76.250  16.21  -4.705 0.019509   TRUE
## 3  1v4 -100.875  16.21  -6.225 0.005912   TRUE
## 4  2v3  -71.875  16.21  -4.435 0.024757   TRUE
## 5  2v4  -96.500  16.21  -5.955 0.007188   TRUE
## 6  3v4  -24.625  16.21  -1.519 0.491534  FALSE
```

Secondly, we can use `emmeans` to generate the same output from an `lm` object.

```
tyre.emm <- emmeans::emmeans(tyre.lm, ~ compound)
pairs(tyre.emm)
```

```
##   contrast estimate    SE df t.ratio p.value
## 1 - 2        -4.37  16.2   5  -0.270  0.9923
## 1 - 3       -76.25  16.2   5  -4.705  0.0195
## 1 - 4      -100.87  16.2   5  -6.225  0.0059
## 2 - 3       -71.88  16.2   5  -4.435  0.0248
```



```
## 2 - 4      -96.50 16.2  5  -5.955  0.0072
## 3 - 4      -24.62 16.2  5  -1.519  0.4915
##
## Results are averaged over the levels of: block
## P value adjustment: tukey method for comparing a family of 4 estimates
```

For this experiment, treatments 1 and 3, 1 and 4, 2 and 3, and 2 and 4 are significantly different at an experiment-wise type I error rate of 5%.

3.7 Exercises

1. Consider the below randomised complete block design for comparing two catalysts, *A* and *B*, for a chemical reaction using six batches of material. The response is the yield (%) from the reaction.

Catalyst	Batch 1	Batch 2	Batch 3	Batch 4	Batch 5	Batch 6
A	9	19	28	22	18	8
B	10	22	30	21	23	12

- i. Write down a unit-block-treatment model for this example.
- ii. Test if there is a significant difference between catalysts at the 5% level.
- iii. Fit a unit-treatment model ignoring blocks and test again for a difference between catalysts. Comment on difference between this analysis and the one including blocks.

Solution

- i. The unit-block-treatment model for this RCBD is given by

$$y_{ij} = \mu + \beta_i + \tau_j + \varepsilon_{ij} \quad i = 1, \dots, 6; j = A, B, \quad (3.25)$$

where y_{ij} is the yield from the application of catalyst j to block i , μ is a constant parameter, β_i is the effect of block i and τ_j is the effect of treatment j . The errors follow a normal distribution $\varepsilon_{ij} \sim N(0, \sigma^2)$ with mean 0 and constant variance, and are assumed independent for different experimental units.

- ii. To test if there is a difference between catalysts, we compare model (3.25) with the model that only includes block effects:

$$y_{ij} = \mu + \beta_i + \varepsilon_{ij} \quad i = 1, \dots, 6; j = A, B. \quad (3.26)$$

The relative difference in the residual mean squares between these two models follows an F distribution under $H_0 : \tau_1 = \tau_2 = 0$, see Section 3.3.

These test statistic and associated p-value can be calculated in R using `anova`.

```
reaction <- data.frame(
  catalyst = factor(rep(c('A', 'B'), 6)),
  batch = factor(rep(1:6, rep(2, 6))),
  yield = c(9, 10, 19, 22, 28, 30, 22, 21, 18, 23, 8, 12)
)
reaction.lm <- lm(yield ~ batch + catalyst, data = reaction)
anova(reaction.lm)
```

```
## Analysis of Variance Table
##
## Response: yield
##           Df Sum Sq Mean Sq F value Pr(>F)
## batch      5    561    112.2    48.1 0.00031 ***
## catalyst   1     16     16.3     7.0 0.04566 *
## Residuals  5      12      2.3
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The p-value is (just) less than 0.05, and so we can reject H_0 (no treatment difference) at the 5% level.

iii. The unit-treatment model is given by

$$y_{ij} = \mu + \tau_j + \varepsilon_{ij}, \quad i = 1, \dots, 6; j = 1, 2,$$

where now **all** the between unit differences, including any block-to-block differences, are modelled by the unit errors ε_{ij} . We can fit this model in R.

```
reaction2.lm <- lm(yield ~ catalyst, data = reaction)
anova(reaction2.lm)
```

```
## Analysis of Variance Table
##
## Response: yield
##           Df Sum Sq Mean Sq F value Pr(>F)
## catalyst   1     16     16.3     0.29  0.6
## Residuals 10    573     57.3
```

There is no longer evidence to reject the null hypothesis of no treatment difference. This is because the residual mean square is now so much larger (57.2667 versus 2.3333). The residual mean square is also an unbiased estimate of σ^2 , and our estimate of σ^2 from the unit-treatment model is clearly much larger, as block-to-block variation has also been included.

2. Consider the data below obtained from an agricultural experiment in which six different fertilizers were given to a crop of blackcurrants in a field. The

Table 3.5: Blackcurrent experiment: yield (lbs) from six different fertilizers.

Block	Fertilizer 1	Fertilizer 2	Fertilizer 3	Fertilizer 4	Fertilizer 5	Fertilizer 6
1	14.5	13.5	11.5	13.0	15	12.5
2	12.0	10.0	11.0	13.0	12	13.5
3	9.0	9.0	14.0	13.5	8	14.0
4	6.5	8.5	10.0	7.5	7	8.0

field was divided into four equal areas of land so that the land in each area was fairly homogeneous. Each area of land was further subdivided into six plots and one of the fertilizers, chosen by a random procedure, was applied to each plot. The yields of blackcurrants obtained from each plot were recorded (in lbs) and are given in Table 3.5. In this randomised block design the treatments are the six fertilizers and the blocks are the four areas of land.

```
blackcurrent <- data.frame(fertilizer = rep(factor(1:6), 4),
  block = rep(factor(1:4), rep(6, 4)),
  yield = c(14.5, 13.5, 11.5, 13.0, 15.0, 12.5,
            12.0, 10.0, 11.0, 13.0, 12.0, 13.5,
            9.0, 9.0, 14.0, 13.5, 8.0, 14.0,
            6.5, 8.5, 10.0, 7.5, 7.0, 8.0)
)
knitr::kable(
  tidyr::pivot_wider(blackcurrent, names_from = fertilizer, values_from = yield),
  col.names = c("Block", paste("Fertilizer", 1:6)),
  caption = "Blackcurrent experiment: yield (lbs) from six different fertilizers."
)
```

Conduct a full analysis of this experiment, including

- exploratory data analysis;
- fitting an appropriate linear model, and conducting an F-test to compare a model that explains variation between the six fertilizers to the model only containing blocks;
- Linear model diagnostics;
- if appropriate, multiple comparisons of all pairwise differences between treatments.

Solution

- We start with exploratory tabular and graphical analysis.

```
aggregate(yield ~ fertilizer, data = blackcurrent,
  FUN = function(x) c(mean = mean(x), sd = sd(x)))
aggregate(yield ~ block, data = blackcurrent,
  FUN = function(x) c(mean = mean(x), sd = sd(x)))
```

```
boxplot(yield ~ fertilizer, data = blackcurrent)
boxplot(yield ~ block, data = blackcurrent)
```

```
##   fertilizer yield.mean yield.sd
## 1          1      10.500    3.488
## 2          2      10.250    2.255
## 3          3      11.625    1.702
## 4          4      11.750    2.843
## 5          5      10.500    3.697
## 6          6      12.000    2.739
##   block yield.mean yield.sd
## 1      1      13.333    1.291
## 2      2      11.917    1.281
## 3      3      11.250    2.859
## 4      4       7.917    1.242
```

There are substantial differences in average responses between blocks; differences between treatment means are smaller. These plots are only meaningful because this design is an RCBD, and each treatment occurs in each block.

The unit-block-treatment model is *additive*; that is, we assume the effect of each treatment does not vary for each block. Therefore we also need to check that there are no substantive interactions between treatments and blocks. We will do this graphically.

```
with(blackcurrent,
      interaction.plot(fertilizer, block, yield, xlab = 'Treatment', ylab = 'Yield', t
    )
```

Figure 3.4 plots the treatment means within each block. While there are clear differences between blocks, the differences between treatments do not seem to vary systematically between blocks.

- b. We now fit a linear model, and perform an F-test for the hypothesis $H_0 : \tau_1 = \tau_2 = \tau_3 = \tau_4 = \tau_5 = \tau_6 = 0$.

```
blackcurrent.lm <- lm(yield ~ block + fertilizer, data = blackcurrent)
anova(blackcurrent.lm)
```

```
## Analysis of Variance Table
##
## Response: yield
##           Df Sum Sq Mean Sq F value Pr(>F)
## block      3   94.9    31.62   8.90 0.0013 **
## fertilizer  5   11.8     2.36   0.66 0.6564
## Residuals 15   53.3     3.55
## ---
```

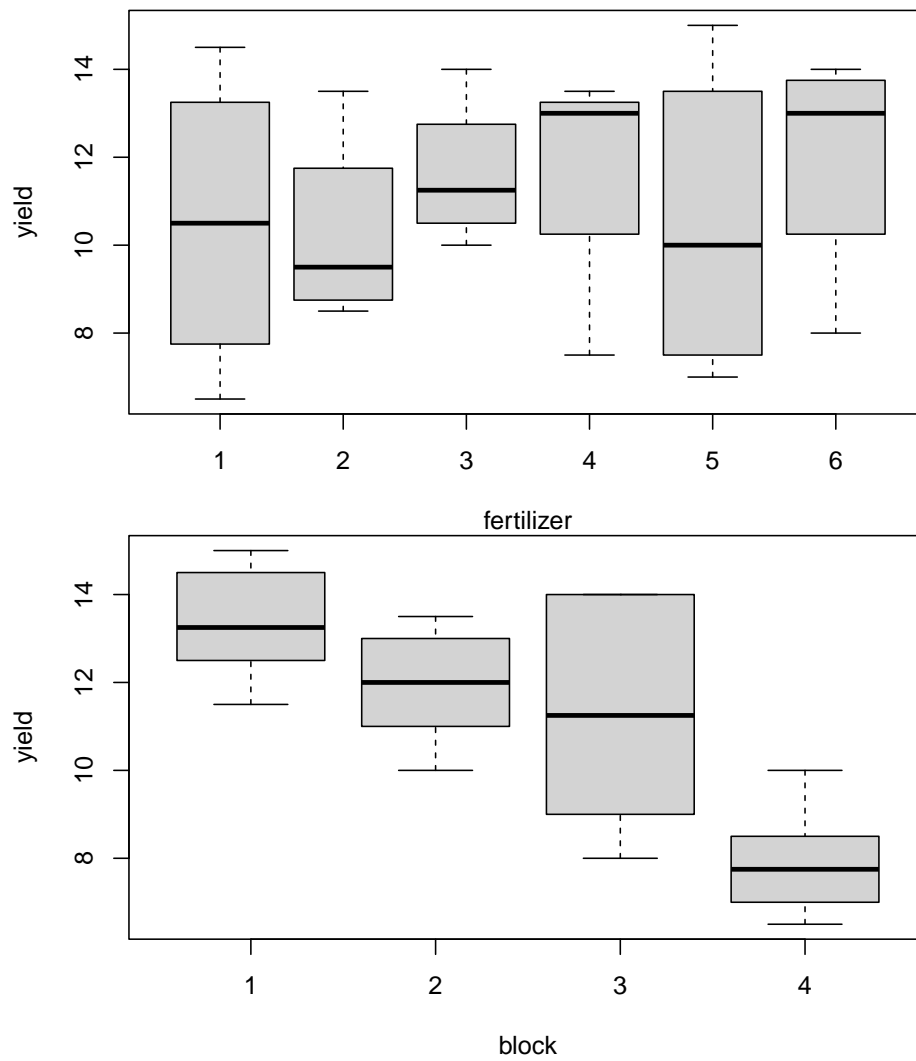


Figure 3.3: Blackcurrent experiment: yield against treatment and block.

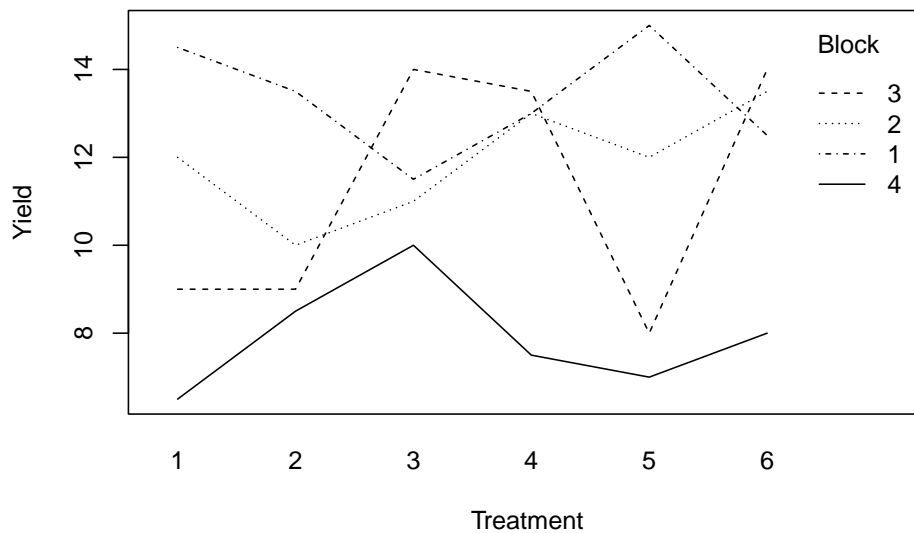


Figure 3.4: Blackcurrent experiment: treatment-block interaction plot.

```
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The “treatment” row of the ANOVA table compares the model including blocks and treatments to that only containing blocks. This comparison tests the above null hypothesis. We can see here that there is no evidence to reject H_0 (p-value = 0.6564). This outcome is not surprising, given the tabular and graphical summaries we saw above. The block sum of squares is large, but we do not perform formal hypothesis testing for blocks.

Out of curiosity, we can also assess the “efficiency” of blocking by comparing the estimate of σ^2 from the block design with the estimate that would result from ignoring the blocks, treating the experiment as a CRD and fitting a unit-treatment model.

```
blackcurrent_crd.lm <- lm(yield ~ fertilizer, data = blackcurrent)
summary(blackcurrent_crd.lm)$sig^2 / summary(blackcurrent.lm)$sig^2
```

```
## [1] 2.316
```

The estimate of σ^2 from the CRD is more the two times greater than the estimate from the block design, meaning about 100% more observations would be needed in the CRD to get the same level of precision as the RCBD.

- c. We now examine residual diagnostics, to check the assumptions of our model:
 - constant variance, with respect to the mean response, the treatment and the block

- normality of residuals
- additive treatment and block effects (already assessed in Fig 3.4).

```
standres <- rstandard(blackcurrent.lm)
fitted <- fitted(blackcurrent.lm)
par(mfrow = c(1, 3), pty = "s")
with(blackcurrent, {
  plot(fertilizer, standres, xlab = "Treatment", ylab = "Standardised residuals")
  plot(block, standres, xlab = "Block", ylab = "Standardised residuals")
  plot(fitted, standres, xlab = "Fitted value", ylab = "Standardised residuals")
})
```

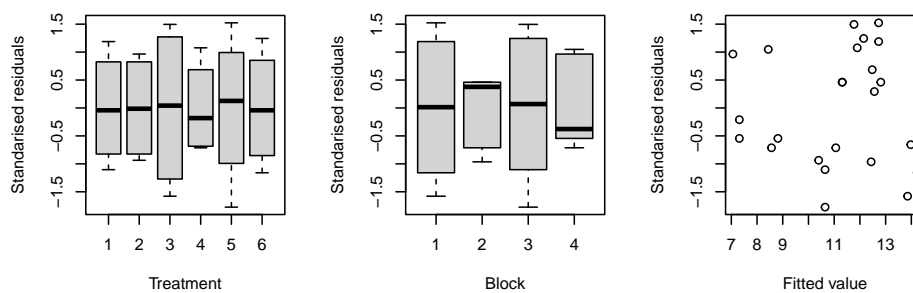


Figure 3.5: Blackcurrent experiment: Residuals against treatments (left), blocks (middle) and fitted values (right).

The plots in Figure 3.5 do not show any serious evidence of non-constant variance (maybe very slightly for blocks), and no large outliers.

```
par(pty = "s")
qqnorm(standres, main = "")
```

Figure 3.6 shows the residuals lie roughly on a straight line when plotted against theoretical normal quantiles, and hence the assumption of normally distributed errors seems reasonable.

- d. There is no evidence of a difference between treatments, so we would not normally test each pairwise difference. However, if we did, we could use the following code.

```
blackcurrent.emm <- emmeans::emmeans(blackcurrent.lm, ~ fertilizer)
pairs(blackcurrent.emm)
```

##	contrast	estimate	SE	df	t.ratio	p.value
##	1 - 2	0.250	1.33	15	0.188	1.0000
##	1 - 3	-1.125	1.33	15	-0.844	0.9541
##	1 - 4	-1.250	1.33	15	-0.938	0.9303
##	1 - 5	0.000	1.33	15	0.000	1.0000
##	1 - 6	-1.500	1.33	15	-1.125	0.8635



Figure 3.6: Blackcurrent experiment: Normal probability plot for standardised residuals.


```
## 2 - 3      -1.375 1.33 15  -1.031  0.9000
## 2 - 4      -1.500 1.33 15  -1.125  0.8635
## 2 - 5      -0.250 1.33 15  -0.188  1.0000
## 2 - 6      -1.750 1.33 15  -1.313  0.7741
## 3 - 4      -0.125 1.33 15  -0.094  1.0000
## 3 - 5       1.125 1.33 15   0.844  0.9541
## 3 - 6      -0.375 1.33 15  -0.281  0.9997
## 4 - 5       1.250 1.33 15   0.938  0.9303
## 4 - 6      -0.250 1.33 15  -0.188  1.0000
## 5 - 6      -1.500 1.33 15  -1.125  0.8635
##
## Results are averaged over the levels of: block
## P value adjustment: tukey method for comparing a family of 6 estimates
```

3. Prove that $2k/\lambda t > 2/b$, and use this result to compare the precision of a pairwise treatment comparison from a BIBD with block size k and an RCBD, both with t treatments in b blocks.

Solution

From the notes, $r = bk/t$ and hence $k = rt/b$. Therefore

$$\begin{aligned}\frac{2k}{\lambda t} &= \frac{2rt}{\lambda bt} \\ &= \frac{2r}{\lambda b} \\ &= \frac{2}{b} \times \frac{r}{\lambda} \\ &> \frac{2}{b},\end{aligned}$$

as $r/\lambda > 1$. The variance of the estimator of a pairwise comparison is

$$\text{var}_{\text{RCBD}}(\tau_i - \tau_j) = \frac{2\sigma^2}{b},$$

for a RCBD with b blocks, and

$$\text{var}_{\text{BIBD}}(\tau_i - \tau_j) = \frac{2k\sigma^2}{\lambda t},$$

for a BIBD. Hence, we can see that the ratio of these variances is

$$\frac{kb}{\lambda t} > 1,$$

Table 3.6: BIBD for $t = 5$ treatments in $b = 5$ blocks of size $k = 4$.

Block 1	1	2	3	4
Block 2	1	2	3	5
Block 3	1	2	4	5
Block 4	1	3	4	5
Block 5	2	3	4	5

and hence the variance for a BIBD is larger, i.e. incomplete blocks results in less precise treatment comparisons.

4. Construct a BIBD for $t = 5$ treatments in $b = 5$ blocks of size $k = 4$ units. What are r and λ for your design? Compare your design to a RCBD via the efficiency for estimating a pairwise treatment difference.

Solution

Here, $b = \binom{t}{k} = \binom{5}{4} = 5$, and hence we can use the “naive” construction method from the notes, where we can form one block from each possible subset of $k = 4$ treatments. This design will have $r = \binom{4}{3} = 4$ and $\lambda = \binom{3}{2} = 3$.

```
bibd <- t(combn(1:5, 4))
rownames(bibd) <- paste("Block", 1:5)
knitr::kable(bibd, caption = "BIBD for $t = 5$ treatments in $b=5$ blocks of size $k=4$")
```

From Question 3, the efficiency of this BIBD compared to a RCBD with $b = 5$ blocks of size $k = 5$ is given by $\frac{\lambda t}{kb} = 0.75$. So variances will be 25% larger under the BIBD.

5. Consider an experiment for testing the abrasion resistance of rubber-coated fabric. There are four types of material, denoted A - D. The response is the loss in weight in 0.1 milligrams (mg) over a standard period of time. The testing machine has four positions, so four samples of material can be tested at a time. Past experience suggests that there may be differences between these positions, and there may be differences between each application of the testing machine (due to changes in set-up). Therefore, we have two blocking variables, “Position” and “Application”. For this experiment, we use a **latin square** design, as follows.

```
fabric <- data.frame(material = factor(c('C', 'D', 'B', 'A', 'A', 'B', 'D', 'C',
                                         'D', 'C', 'A', 'B', 'B', 'A', 'C', 'D')),
                     position = rep(factor(1:4), 4),
                     application = rep(factor(1:4), rep(4, 4)),
                     weight = c(235, 236, 218, 268,
                                251, 241, 227, 229,
                                234, 273, 274, 226,
                                195, 270, 230, 225))
```

```

)
knitr::kable(
  tidyr::pivot_wider(fabric, id_cols = application, names_from = position,
    values_from = material),
  col.names = c("Application", paste("Position", 1:4)))

```

Application	Position 1	Position 2	Position 3	Position 4
1	C	D	B	A
2	A	B	D	C
3	D	C	A	B
4	B	A	C	D

The blocking variables are represented as the rows and columns of the square; the latin letters represent the different treatments. A latin square of order k is a $k \times k$ square of k latin letters arranged so that each letter appears exactly once in each row and column (Sudoku squares are also examples of latin squares). To perform the experiment, the levels of the blocking factors are randomly assigned to the rows and the columns, and the different treatments to the letters. A latin square design is a special case of the more general class of **row column** designs for two blocking factors.

A suitable unit-block-treatment model for a latin square design has the form

$$y_{ijk} = \mu + \beta_i + \gamma_j + \tau_k + \varepsilon_{ijk}, \quad i, j, k = 1, \dots, t,$$

with μ a constant parameter, β_i row block effects, γ_j column block effects and τ_k the treatment effects. As usual, $\varepsilon_{ijk} \sim N(0, \sigma^2)$, with errors from different units assumed independent. Note that not all combinations of i, j, k actually occur in the design; at the intersection of the i th row and j th column, only one of the t treatments is applied.

- Write down a set of normal equations for the model parameters.
- It can be shown that the reduced normal equations for the treatment parameters τ_1, \dots, τ_t have the form

$$\hat{\tau}_k - \hat{\tau}_w = \bar{y}_{..j} - \bar{y}_{...},$$

with $\hat{\tau}_w = \frac{1}{t} \sum_{k=1}^t \hat{\tau}_k$, $\bar{y}_{..k} = \frac{1}{t} \sum_{i=1}^t \sum_{j=1}^t n_{ijk} y_{ijk}$ (mean for treatment k) and $\bar{y}_{...} = \frac{1}{n} \sum_{i=1}^t \sum_{j=1}^t \sum_{k=1}^t n_{ijk} y_{ijk}$ (overall mean) where $n_{ijk} = 1$ if treatment k occurs at the intersection of row i and column j and zero otherwise, and $\sum_{i=1}^t \sum_{j=1}^t n_{ijk} = t$ for all $k = 1, \dots, t$.

Demonstrate that any contrast can therefore be estimated from this design, and derive the variance of the estimator of $\sum_{k=1}^t c_k \tau_k$.

The data for this experiment is as follows, where the entries in the table give the response for the corresponding treatment:

```
knitr::kable(
  tidyr::pivot_wider(fabric, id_cols = application, names_from = position,
    values_from = weight),
  col.names = c("Application", paste("Position", 1:4)))
```

Application	Position 1	Position 2	Position 3	Position 4
1	235	236	218	268
2	251	241	227	229
3	234	273	274	226
4	195	270	230	225

- c. For this data, test if there is a significant difference between materials. If there is, conduct multiple comparisons of all pairs at an experimentwise type I error rate of 5%.

Solution

- a. We start by writing the model in matrix form:

$$\begin{aligned}\mathbf{y} &= \mathbf{1}_n\mu + X_1\boldsymbol{\beta} + X_2\boldsymbol{\gamma} + X_3\boldsymbol{\tau} + \boldsymbol{\varepsilon} \\ &= W\boldsymbol{\alpha} + \boldsymbol{\varepsilon},\end{aligned}$$

with $W = (\mathbf{1}_n, X_1, X_2, X_3)$, X_1 , X_2 and X_3 being $n \times t$ model matrices for row blocks, column blocks and treatments, respectively, $\boldsymbol{\alpha}^T = (\mu, \boldsymbol{\beta}^T, \boldsymbol{\gamma}^T, \boldsymbol{\tau}^T)$ being a $(1+3t)$ -vector of parameters, and $\boldsymbol{\varepsilon} \sim N(\mathbf{0}_n, I_n\sigma^2)$.

The normal equations are given by $W^T W \hat{\boldsymbol{\alpha}} = W^T \mathbf{y}$, which can be expanded out to give the following four matrix equations:

$$n\hat{\mu} + \mathbf{1}_n^T X_1 \hat{\boldsymbol{\beta}} + \mathbf{1}_n^T X_2 \hat{\boldsymbol{\gamma}} + \mathbf{1}_n^T X_3 \hat{\boldsymbol{\tau}} = \mathbf{1}_n^T \mathbf{y}, \quad (3.27)$$

$$X_1^T \mathbf{1}_n \hat{\mu} + X_1^T X_1 \hat{\boldsymbol{\beta}} + X_1^T X_2 \hat{\boldsymbol{\gamma}} + X_1^T X_3 \hat{\boldsymbol{\tau}} = X_1^T \mathbf{y}, \quad (3.28)$$

$$X_2^T \mathbf{1}_n \hat{\mu} + X_2^T X_1 \hat{\boldsymbol{\beta}} + X_2^T X_2 \hat{\boldsymbol{\gamma}} + X_2^T X_3 \hat{\boldsymbol{\tau}} = X_2^T \mathbf{y}, \quad (3.29)$$

$$X_3^T \mathbf{1}_n \hat{\mu} + X_3^T X_1 \hat{\boldsymbol{\beta}} + X_3^T X_2 \hat{\boldsymbol{\gamma}} + X_3^T X_3 \hat{\boldsymbol{\tau}} = X_3^T \mathbf{y}. \quad (3.30)$$

$$(3.31)$$

- b. From the reduced normal equations given in the question, we have

$$\begin{aligned}
\sum_{k=1}^t c_k (\hat{\tau}_k - \hat{\tau}_w) &= \sum_{k=1}^t c_k \hat{\tau}_k \\
&= \sum_{k=1}^t c_k (\bar{y}_{..k} - \bar{y}_{...}) \\
&= \sum_{k=1}^t c_k \bar{y}_{..k},
\end{aligned}$$

as $\sum_{k=1}^t c_k \hat{\tau}_w = \sum_{k=1}^t c_k \bar{y}_{...} = 0$, as neither $\hat{\tau}_w$ or $\bar{y}_{...}$ depend on k . Hence,

$$\widehat{\mathbf{c}^T \boldsymbol{\tau}} = \sum_{k=1}^t c_k \bar{y}_{..k}.$$

Or notice that the reduced normal equations for this design are the same as for a CRD with each treatment replicated t times.

c. We conduct these hypothesis tests using R.

```
fabric.lm <- lm(weight ~ position + application + material, data = fabric)
anova(fabric.lm)
```

```
## Analysis of Variance Table
##
## Response: weight
##           Df Sum Sq Mean Sq F value    Pr(>F)
## position    3   1468      489    7.99 0.01617 *
## application  3    986      329    5.37 0.03901 *
## material     3   4621     1540   25.15 0.00085 ***
## Residuals    6    368        61
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The “material” line of the ANOVA table compares the models with and without the effects for material (but both models including the blocking factors). There is clearly a significant effect of material.

```
fabric.emm <- emmeans::emmeans(fabric.lm, ~ material)
fabric.pairs <- transform(pairs(fabric.emm))
dplyr::mutate(fabric.pairs, reject = p.value < 0.05)

##   contrast estimate    SE df t.ratio p.value reject
## 1    A - B     45.75 5.534  6   8.267 0.000703   TRUE
## 2    A - C     24.00 5.534  6   4.337 0.019036   TRUE
## 3    A - D     35.25 5.534  6   6.370 0.002866   TRUE
```

```
## 4      B - C    -21.75 5.534 6   -3.930 0.029477   TRUE
## 5      B - D    -10.50 5.534 6   -1.897 0.320631  FALSE
## 6      C - D     11.25 5.534 6    2.033 0.274277  FALSE
```

Using an experiment-wise error rate of 5%, we see significant differences between materials A and B, A and C, A and D, and B and C.

Chapter 4

Factorial experiments

In Chapters 2 and 3, we assumed the objective of the experiment was to investigate t **unstructured** treatments, defined only as a collection of distinct entities (drugs, advertisements, receipes, etc.). That is, there was not necessarily any explicit relationship between the treatments (although we could clearly choose which paticular comparisons between treatments were of interest via choice of contrast).

In many experiments, particularly in industry, engineering and the physical sciences, the treatments are actually defined via the choice of a **level** relating to each of a set of **factors**. We will focus on the commonly occurring case of factors at **two levels**. For example, consider the below experiment from the pharmaceutical industry.

Example 4.1. Desilylation experiment (Owen et al., 2001)

In this experiment, performed at GlaxoSmithKline, the aim was to optimise the desilylation¹ of an ether into an alcohol, which was a key step in the synthesis of a particular antibiotic. There were $t = 16$ treatments, defined via the settings of four different factors, as given in Table 4.1.

```
desilylation <- FrF2::FrF2(nruns = 16, nfactors = 4, randomize = F,
                          factor.names = list(temp = c(10, 20), time = c(19, 25),
                                                solvent = c(5, 7), reagent = c(1, 1.33)))
yield <- c(82.93, 94.04, 88.07, 93.97, 77.21, 92.99, 83.60, 94.38,
          88.68, 94.30, 93.00, 93.42, 84.86, 94.26, 88.71, 94.66)
desilylation <- data.frame(desilylation, yield = yield)
rownames(desilylation) <- paste("Trt", 1:16)
knitr::kable(desilylation,
              col.names = c("Temp (degrees C)", "Time (hours)", "Solvent (vol.)",
                           "Reagent (equiv.)", "Yield (%)"
```

¹Desilylation is a process of removing silyl, SiH₃ a silicon hydride, from a compound.

Table 4.1: Desilylation experiment: 16 treatments defined by settings of four factors, with response (yield).

	Temp (degrees C)	Time (hours)	Solvent (vol.)	Reagent (equiv.)	Yield (%)
Trt 1	10	19	5	1	82.93
Trt 2	20	19	5	1	94.04
Trt 3	10	25	5	1	88.07
Trt 4	20	25	5	1	93.97
Trt 5	10	19	7	1	77.21
Trt 6	20	19	7	1	92.99
Trt 7	10	25	7	1	83.60
Trt 8	20	25	7	1	94.38
Trt 9	10	19	5	1.33	88.68
Trt 10	20	19	5	1.33	94.30
Trt 11	10	25	5	1.33	93.00
Trt 12	20	25	5	1.33	93.42
Trt 13	10	19	7	1.33	84.86
Trt 14	20	19	7	1.33	94.26
Trt 15	10	25	7	1.33	88.71
Trt 16	20	25	7	1.33	94.66

```
caption = "Desilylation experiment: 16 treatments defined
by settings of four factors, with response (yield).")
```

Each treatment is defined by the choice of one of two levels for each of the four factors. In the R code above, we have used the function `FrF2` (from the package of the same name) to generate all $t = 2^4 = 16$ combinations of the two levels of these four factors. We come back to this function later in the chapter.

This **factorial treatment structure** lends itself to certain treatment contrasts being of natural interest.

4.1 Factorial contrasts

Throughout this chapter, we will assume there are no blocks or other restrictions on randomisation, and so we will assume a completely randomised design can be used. We start by assuming the same unit-treatment model as Chapter 2:

$$y_{ij} = \mu + \tau_i + \varepsilon_{ij}, \quad i = 1, \dots, t; j = 1, \dots, n_i, \quad (4.1)$$

where y_{ij} is the response from the j th application of treatment i , μ is a constant parameter, τ_i is the effect of the i th treatment, and ε_{ij} is the random individual

effect from each experimental unit with $\varepsilon_{ij} \sim N(0, \sigma^2)$ independent of other errors.

Now, the number of treatments $t = 2^f$, where f equals the number of factors in the experiment.

For Example 4.1, we have $t = 2^4 = 16$ and $n_i = 1$ for all $i = 1, \dots, 16$; that is, each of the 16 treatments are replicated once. In general, we shall assume common treatment replication $n_i = r \geq 1$.

If we fit model (4.1) and compute the ANOVA table, we notice a particular issue with this design.

```
desilylation <- data.frame(desilylation, trt = factor(1:16))
desilylation.lm <- lm(yield ~ trt, data = desilylation)
anova(desilylation.lm)
```

```
## Analysis of Variance Table
##
## Response: yield
##           Df Sum Sq Mean Sq F value Pr(>F)
## trt        15    427    28.5      NaN    NaN
## Residuals   0         0      NaN
```

All available degrees of freedom are being used to estimate parameters in the mean (μ and the treatment effects τ_i). There are no degrees of freedom left to estimate σ^2 . This is due to a lack of treatment replication. Without replication in the design, model (4.1) is **saturated**, with as many treatments as there are observations and an unbiased estimate of σ^2 cannot be obtained. We will return to this issue later.

4.1.1 Main effects

Studying Table 4.1, there are some comparisons between treatments which are obviously of interest. For example, comparing the average effect from the first 8 treatments with the average effect of the second 8, using

$$\mathbf{c}^T \boldsymbol{\tau} = \sum_{i=1}^t c_i \tau_i,$$

with

$$\mathbf{c}^T = (-\mathbf{1}_8^T, \mathbf{1}_8^T)/8.$$

```
desilylation.emm <- emmeans::emmeans(desilylation.lm, ~ trt)
reagent_me.emmc <- function(levs) data.frame('reagent m.e.' = rep(c(-1, 1), rep(8, 2)) / 8)
emmeans::contrast(desilylation.emm, 'reagent_me')
```

```
## contrast      estimate SE df t.ratio p.value
## reagent.m.e.      3.09 NaN  0      NaN      NaN
```

This contrast compares the average treatment effect from the 8 treatments which have **reagent** set to its low level (1 equiv.) to the average effect from the 8 treatments which have **reagent** set to its high level. This is a “fair” comparison, as both of these sets of treatments have each of the combinations of the factors **temp**, **time** and **solvent** occurring equally often (twice here). Hence, the **main effect** of **reagent** is averaged over the levels of the other three factors.

As in Chapter 2, we can estimate this treatment contrast by applying the same contrast coefficients to the treatment means,

$$\widehat{\mathbf{c}^T \boldsymbol{\tau}} = \sum_{i=1}^t c_i \bar{y}_{i.},$$

where, for this experiment, each $\bar{y}_{i.}$ is the mean of a single observation (as there is no treatment replication). We see that inference about this contrast is not possible, as no standard error can be obtained.

Definition 4.1. The **main effect** of a factor A is defined as the difference in the average response from the high and low levels of the factor

$$ME(A) = \bar{y}(A+) - \bar{y}(A-),$$

where $\bar{y}(A+)$ is the average response when factor A is set to its high level, averaged across all combinations of levels of the other factors (with $\bar{y}(A+)$ defined similarly for the low level of A).

As we have averaged the response across the levels of the other factors, the interpretation of the main effect extends beyond this experiment. That is, we can use it to infer something about the system under study. Assuming model (4.1) is correct, any variation in the main effect can only come from random error in the observations. In fact,

$$\begin{aligned} \text{var}\{ME(A)\} &= \frac{\sigma^2}{n/2} + \frac{\sigma^2}{n/2} \\ &= \frac{4\sigma^2}{n}, \end{aligned}$$

and assuming $r > 1$,

$$\hat{\sigma}^2 = \frac{1}{2^f(r-1)} \sum_{i=1}^{2^f} \sum_{j=1}^r (y_{ij} - \bar{y}_{i.})^2, \quad (4.2)$$

which is the residual mean square.

Table 4.2: Desilylation experiment: main effect contrast coefficients

	Temperature	Time	Solvent	Reagent
Trt 1	-0.125	-0.125	-0.125	-0.125
Trt 2	0.125	-0.125	-0.125	-0.125
Trt 3	-0.125	0.125	-0.125	-0.125
Trt 4	0.125	0.125	-0.125	-0.125
Trt 5	-0.125	-0.125	0.125	-0.125
Trt 6	0.125	-0.125	0.125	-0.125
Trt 7	-0.125	0.125	0.125	-0.125
Trt 8	0.125	0.125	0.125	-0.125
Trt 9	-0.125	-0.125	-0.125	0.125
Trt 10	0.125	-0.125	-0.125	0.125
Trt 11	-0.125	0.125	-0.125	0.125
Trt 12	0.125	0.125	-0.125	0.125
Trt 13	-0.125	-0.125	0.125	0.125
Trt 14	0.125	-0.125	0.125	0.125
Trt 15	-0.125	0.125	0.125	0.125
Trt 16	0.125	0.125	0.125	0.125

For Example 4.1, we can also calculate main effect estimates for the other three factors by defining appropriate contrasts in the treatments.

```
contrast.mat <- FrF2(nruns = 16, nfactors = 4, randomize = F,
                     factor.names = c("temp", "time", "solvent", "reagent"))
fac.contrasts.emmc <- function(levs)
  dplyr::mutate_all(data.frame(contrast.mat), function(x) scale(as.numeric(as.character(x)), scale = F))
main_effect_contrasts <- fac.contrasts.emmc()
rownames(main_effect_contrasts) <- paste("Trt", 1:16)
knitr::kable(main_effect_contrasts, caption = 'Desilylation experiment: main effect contrast coefficients')
```

Estimates can be obtained by applying these coefficients to the observed treatment means.

```
t(as.matrix(main_effect_contrasts)) %*% yield

##           [,1]
## temp      8.120
## time      2.567
## solvent -2.218
## reagent   3.087

emmeans::contrast(desilylation.emm, 'fac.contrasts')

## contrast estimate SE df t.ratio p.value
## temp           8.12 NaN  0      NaN      NaN
```

```
## time      2.57 NaN 0    NaN    NaN
## solvent   -2.22 NaN 0    NaN    NaN
## reagent    3.09 NaN 0    NaN    NaN
```

Main effects are often displayed graphically, using **main effect plots** which simply plot the average response for each factor level, joined by a line. The larger the main effect, the larger the slope of the line (or the bigger the difference between the averages). Figure 4.1 presents the four main effect plots for Example 4.1.

```
## calculate the means
temp_bar <- aggregate(yield ~ temp, data = desilylation, FUN = mean)
time_bar <- aggregate(yield ~ time, data = desilylation, FUN = mean)
solvent_bar <- aggregate(yield ~ solvent, data = desilylation, FUN = mean)
reagent_bar <- aggregate(yield ~ reagent, data = desilylation, FUN = mean)

## convert factors to numeric
fac_to_num <- function(x) as.numeric(as.character(x))
temp_bar$temp <- fac_to_num(temp_bar$temp)
time_bar$time <- fac_to_num(time_bar$time)
solvent_bar$solvent <- fac_to_num(solvent_bar$solvent)
reagent_bar$reagent <- fac_to_num(reagent_bar$reagent)

## main effect plots
plotmin <- min(temp_bar$yield, time_bar$yield, solvent_bar$yield, reagent_bar$yield)
plotmax <- max(temp_bar$yield, time_bar$yield, solvent_bar$yield, reagent_bar$yield)
par(cex = 2)
layout(matrix(c(1, 2, 3, 4), nrow = 2, ncol = 2, byrow = TRUE), respect = T)
plot(temp_bar, pch = 16, type = "b", ylim = c(plotmin, plotmax))
plot(time_bar, pch = 16, type = "b", ylim = c(plotmin, plotmax))
plot(solvent_bar, pch = 16, type = "b", ylim = c(plotmin, plotmax))
plot(reagent_bar, pch = 16, type = "b", ylim = c(plotmin, plotmax))
```

4.1.2 Interactions

Another contrast that could be of interest in Example 4.1 has coefficients

$$\mathbf{c}^T = (\mathbf{1}_4^T, -\mathbf{1}_8^T, \mathbf{1}_4^T)/8.$$

This contrast measures the difference between the average treatment effect from treatments 1-4, 13-16 and treatments 5-12. Checking back against Table 4.1, we see this is comparing those treatments where **solvent** and **reagent** are both set to their low (1-4) or high (13-16) level against those treatments where one of the two factors is set high and the other is set low (5-12).

Focusing on **reagent**, if the effect of this factor on the response was independent

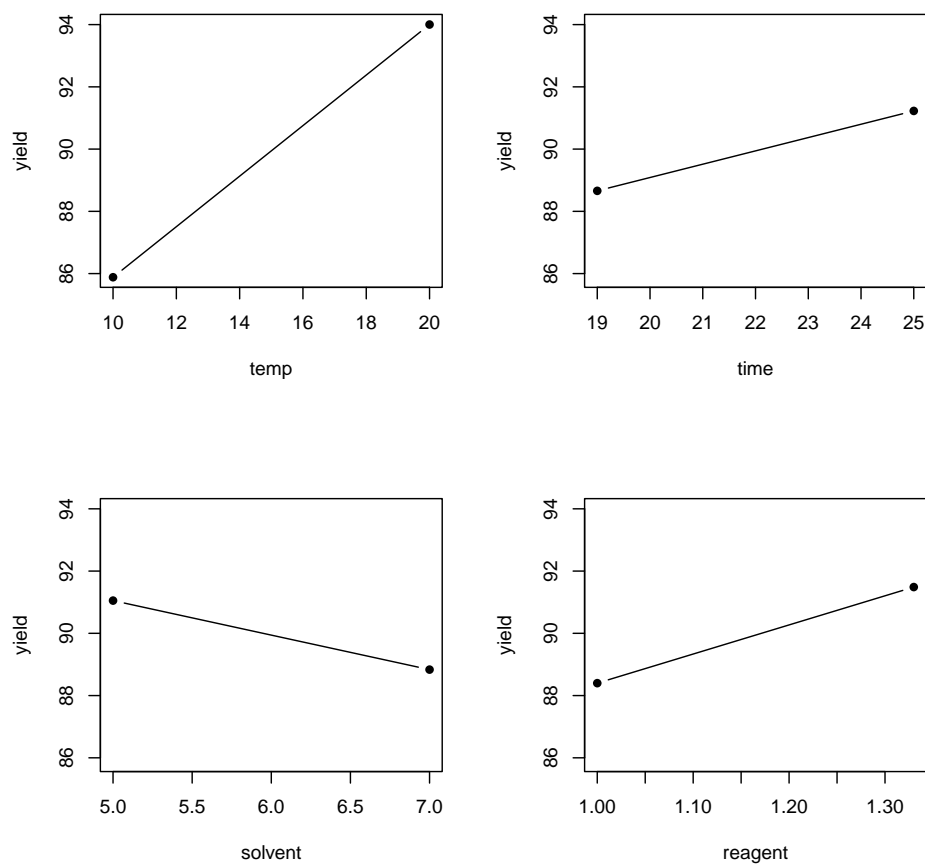


Figure 4.1: Desilylation experiment: main effect plots

of the level to which **solvent** has been set, you would expect this contrast to be zero - changing from the high to low level of **reagent** should affect the response in the same way, regardless of the setting of **solvent**. This argument can be reversed, focussing on the effect of **solvent**. Therefore, if this contrast is large, we say the two factors **interact**.

```
sol_reg_int.emmc <- function(levels)
  data.frame('reagent x solvent' = .125 * c(rep(1, 4), rep(-1, 8), rep(1, 4)))
  emmeans::contrast(desilylation.emm, 'sol_reg_int')
```

```
## contrast      estimate SE df t.ratio p.value
## reagent.x.solvent 0.49 NaN 0      NaN      NaN
```

For Example 4.1, this interaction contrast seems quite small, although of course without an estimate of the standard error we are still lacking a formal method to judge this.

It is somewhat more informative to consider the above interaction contrast as the average difference in two “sub-contrasts”

$$\mathbf{c}^T \boldsymbol{\tau} = \frac{1}{2} \left\{ \frac{1}{4} (\tau_{13} + \tau_{14} + \tau_{15} + \tau_{16} - \tau_5 - \tau_6 - \tau_7 - \tau_8) - \frac{1}{4} (\tau_9 + \tau_{10} + \tau_{11} + \tau_{12} - \tau_1 - \tau_2 - \tau_3 - \tau_4) \right\},$$

The first component in the above expression is the effect of changing **reagent** from high to low **given solvent is set to it's high level**. The second component the effect of changing **reagent** from high to low **given solvent is set to it's low level**. This leads to our definition of a two-factor interaction.

Definition 4.2. The **two-factor interaction** between factors A and B is defined as the average difference in main effect of factor A when computed at the high and low levels of factor B .

$$\begin{aligned} \text{Int}(A, B) &= \frac{1}{2} \{ \text{ME}(A \mid B+) - \text{ME}(A \mid B-) \} \\ &= \frac{1}{2} \{ \text{ME}(B \mid A+) - \text{ME}(B \mid A-) \} \\ &= \frac{1}{2} \{ \bar{y}(A+, B+) - \bar{y}(A-, B+) - \bar{y}(A+, B-) + \bar{y}(A-, B-) \}, \end{aligned}$$

where $\bar{y}(A+, B-)$ is the average response when factor A is set to its high level and factor B is set to its low level, averaged across all combinations of levels of the other factors, and other averages are defined similarly. The conditional main effects of factor A when factor B is set to its high level is defined as

$$\text{ME}(A \mid B+) = \bar{y}(A+, B+) - \bar{y}(A-, B+),$$

with similar definitions for other conditional main effects.

As the sum of the squared contrast coefficients is the same for two-factor interactions as for main effects, the variance of the contrast estimator is also the same.

$$\text{var}\{\text{Int}(A, B)\} = \frac{4\sigma^2}{n}.$$

For Example 4.1 we can calculate two-factor interactions for all $\binom{4}{2} = 6$ pairs of factors. The simplest way to calculate the contrast coefficients is as the elementwise, or Hadamard, product² of the unscaled main effect contrasts (before dividing by $n/2$).

```
fac.contrasts.int.emmc <- function(levs) {
  with(sqrt(8) * main_effect_contrasts, {
    data.frame('tem_x_tim' = temp * time,
               'tem_x_sol' = temp * solvent,
               'tem_x_rea' = temp * reagent,
               'tim_x_sol' = time * solvent,
               'tim_x_rea' = time * reagent,
               'sol_x_rea' = solvent * reagent)
  })
}
two_fi_contrasts <- fac.contrasts.int.emmc()
rownames(two_fi_contrasts) <- paste("Trt", 1:16)
knitr::kable(two_fi_contrasts, caption = 'Desilylation experiment: two-factor interaction contrasts')
```

Estimates of the interaction contrasts can again be found by considering the equivalent contrasts in the observed treatment means.

```
t(as.matrix(two_fi_contrasts)) %*% yield

##           [,1]
## tem_x_tim -2.357
## tem_x_sol  2.358
## tem_x_rea -2.773
## tim_x_sol  0.440
## tim_x_rea -0.645
## sol_x_rea  0.490

emmeans::contrast(desilylation.emm, 'fac.contrasts.int')
```

contrast	estimate	SE	df	t.ratio	p.value
tem_x_tim	-2.357	NaN	0	NaN	NaN
tem_x_sol	2.357	NaN	0	NaN	NaN
tem_x_rea	-2.772	NaN	0	NaN	NaN

²For two matrices A and B of the same dimension $m \times n$, the Hadamard product $A \odot B$ is a matrix of the same dimension with elements given by the elementwise product, $(A \odot B)_{ij} = A_{ij}B_{ij}$.

Table 4.3: Desilylation experiment: two-factor interaction contrast coefficients

	tem_x_tim	tem_x_sol	tem_x_rea	tim_x_sol	tim_x_rea	sol_x_rea
Trt 1	0.125	0.125	0.125	0.125	0.125	0.125
Trt 2	-0.125	-0.125	-0.125	0.125	0.125	0.125
Trt 3	-0.125	0.125	0.125	-0.125	-0.125	0.125
Trt 4	0.125	-0.125	-0.125	-0.125	-0.125	0.125
Trt 5	0.125	-0.125	0.125	-0.125	0.125	-0.125
Trt 6	-0.125	0.125	-0.125	-0.125	0.125	-0.125
Trt 7	-0.125	-0.125	0.125	0.125	-0.125	-0.125
Trt 8	0.125	0.125	-0.125	0.125	-0.125	-0.125
Trt 9	0.125	0.125	-0.125	0.125	-0.125	-0.125
Trt 10	-0.125	-0.125	0.125	0.125	-0.125	-0.125
Trt 11	-0.125	0.125	-0.125	-0.125	0.125	-0.125
Trt 12	0.125	-0.125	0.125	-0.125	0.125	-0.125
Trt 13	0.125	-0.125	-0.125	-0.125	-0.125	0.125
Trt 14	-0.125	0.125	0.125	-0.125	-0.125	0.125
Trt 15	-0.125	-0.125	-0.125	0.125	0.125	0.125
Trt 16	0.125	0.125	0.125	0.125	0.125	0.125

```
## tim_x_sol    0.440 NaN  0    NaN    NaN
## tim_x_rea   -0.645 NaN  0    NaN    NaN
## sol_x_rea    0.490 NaN  0    NaN    NaN
```

As with main effects, interactions are often displayed graphically using **interaction** plots, plotting average responses for each pairwise combination of factors, joined by lines.

```
plotmin <- min(desilylation$yield)
plotmax <- max(desilylation$yield)
par(cex = 2)
layout(matrix(c(1, 2, 3, 4, 5, 6), nrow = 3, ncol = 2, byrow = TRUE), respect = T)

with(desilylation, {
  interaction.plot(temp, time, yield, type = "b", pch = 16, legend = F,
                  ylim = c(plotmin, plotmax))
  legend("bottomright", legend = c("Time = 19", "Time = 25"), lty = 2:1, cex = .75)
  interaction.plot(temp, solvent, yield, type = "b", pch = 16, legend = F,
                  ylim = c(plotmin, plotmax))
  legend("bottomright", legend = c("Solvent = 5", "Solvent = 7"), lty = 2:1, cex = .75)
  interaction.plot(temp, reagent, yield, type = "b", pch = 16, legend = F,
                  ylim = c(plotmin, plotmax))
  legend("bottomright", legend = c("Reagent = 1", "Reagent = 1.33"), lty = 2:1, cex = .75)
  interaction.plot(time, solvent, yield, type = "b", pch = 16, legend = F,
                  ylim = c(plotmin, plotmax))
})
```



```

legend("bottomright", legend = c("Solvent = 5", "Solvent = 7"), lty = 2:1, cex = .75)
interaction.plot(time, reagent, yield, type = "b", pch = 16, legend = F,
                 ylim = c(plotmin, plotmax))
legend("bottomright", legend = c("Reagent = 1", "Reagent = 1.33"), lty = 2:1, cex = .75)
interaction.plot(solvent, reagent, yield, type = "b", pch = 16, legend = F,
                 ylim = c(plotmin, plotmax))
legend("bottomright", legend = c("Reagent = 1", "Reagent = 1.33"), lty = 2:1, cex = .75)
})

```

Parallel lines in an interaction plot indicate no (or very small) interaction (`time` and `solvent`, `time` and `reagent`, `solvent` and `reagent`). The three interactions with `temp` all demonstrate much more robust behaviour at the high level; changing `time`, `solvent` or `reagent` makes little difference to the response at the high level of `temp`, and much less difference than at the low level of `temp`.

If a system displays important interactions, the main effects of factors involved in those interactions should no longer be interpreted. For example, it makes little sense to discuss the main effect of `temp` when it changes so much with the level of `reagent` (from strongly positive when `reagent` is low to quite small when `reagent` is high).

Higher order interactions can be defined similarly, as average differences in lower-order effects. For example, a three-factor interaction measures how a two-factor interaction changes with the levels of a third factor.

$$\begin{aligned}
 \text{Int}(A, B, C) &= \frac{1}{2} \{ \text{Int}(A, B \mid C+) - \text{Int}(A, B \mid C-) \} \\
 &= \frac{1}{2} \{ \text{Int}(A, C \mid B+) - \text{Int}(A, C \mid B-) \} \\
 &= \frac{1}{2} \{ \text{Int}(B, C \mid A+) - \text{Int}(B, C \mid A-) \} ,
 \end{aligned}$$

where

$$\text{Int}(A, B \mid C+) = \frac{1}{2} \{ \bar{y}(A+, B+, C+) - \bar{y}(A-, B+, C+) - \bar{y}(A+, B-, C+) + \bar{y}(A-, B-, C+) \}$$

is the interaction between factors A and B using only those treatments where factor C is set to its high level. Higher order interaction contrasts can again be constructed by (multiple) hadamard products of (unscaled) main effect contrasts.

Definition 4.3. A **factorial effect** is a main effect or interaction contrast defined on a factorial experiment. For a 2^f factorial experiment with f factors, there are $2^f - 1$ factorial effects, ranging from main effects to the interaction between all f factors.



Figure 4.2: Desilylation experiment: two-factor interaction plots

For Example 4.1, we can now calculate all the factorial effects.

```
## hadamard products
unscaled_me_contrasts <- 8 * main_effect_contrasts
factorial_contrasts <- model.matrix(~.^4, unscaled_me_contrasts)[, -1] / 8

## all factorial effects - directly, as there is no treatment replication
t(factorial_contrasts) %*% yield

##
##           [,1]
## temp      8.1200
## time      2.5675
## solvent   -2.2175
## reagent    3.0875
## temp:time  -2.3575
## temp:solvent  2.3575
## temp:reagent -2.7725
## time:solvent  0.4400
## time:reagent -0.6450
## solvent:reagent  0.4900
## temp:time:solvent  0.2450
## temp:time:reagent  0.1950
## temp:solvent:reagent -0.0300
## time:solvent:reagent -0.2375
## temp:time:solvent:reagent  0.1925

## using emmeans
factorial_contrasts.emmc <- function(levs) data.frame(factorial_contrasts)
desilylation.effs <- emmeans::contrast(desilylation.emm, 'factorial_contrasts')
desilylation.effs

## contrast      estimate SE df t.ratio p.value
## temp          8.120 NaN  0      NaN      NaN
## time          2.567 NaN  0      NaN      NaN
## solvent       -2.218 NaN  0      NaN      NaN
## reagent        3.087 NaN  0      NaN      NaN
## temp.time     -2.357 NaN  0      NaN      NaN
## temp.solvent   2.358 NaN  0      NaN      NaN
## temp.reagent  -2.772 NaN  0      NaN      NaN
## time.solvent   0.440 NaN  0      NaN      NaN
## time.reagent  -0.645 NaN  0      NaN      NaN
## solvent.reagent  0.490 NaN  0      NaN      NaN
## temp.time.solvent  0.245 NaN  0      NaN      NaN
## temp.time.reagent  0.195 NaN  0      NaN      NaN
## temp.solvent.reagent -0.030 NaN  0      NaN      NaN
## time.solvent.reagent -0.238 NaN  0      NaN      NaN
## temp.time.solvent.reagent  0.193 NaN  0      NaN      NaN
```

4.2 Three principles for factorial effects

Empirical study of many experiments (Box and Meyer, 1986; Li et al., 2006) have demonstrated that the following three principles **often** hold when analysing factorial experiments.

Definition 4.4. Effect hierarchy: lower-order factorial effects are more likely to be important than higher-order effects; factorial effects of the same order are equally likely to be important.

For example, we would anticipate more large main effects from the analysis of a factorial experiment than two-factor interactions.

Effect sparsity: the number of large factorial effects is likely to be small, relative to the total number under study.

This is sometimes called the **pareto principle**.

Definition 4.5. Effect heredity: interactions are more likely to be important if at least one parent factor also has a large main effect.

These three principles will provide us with some useful guidelines when analysing, and eventually constructing, factorial experiments.

4.3 Normal effect plots for unreplicated factorial designs

The lack of an estimate for σ^2 means alternatives to formal inference methods (e.g. hypothesis tests) must be found to assess the size of factorial effects. We will discuss a method that essentially treats the identification of large factorial effects as an outlier identification problem.

Let $\hat{\theta}_j$ be the j th estimated factorial effect, with $\hat{\theta}_j = \sum_{i=1}^t c_{ij} \bar{y}_i$. for $\mathbf{c}_j^T = (c_{1j}, \dots, c_{tj})$ a vector of factorial contrast coefficients (defining a main effect or interaction). Then the estimator follows a normal distribution

$$\hat{\theta}_j \sim N\left(\theta_j, \frac{4\sigma^2}{n}\right), \quad j = 1, \dots, 2^f - 1,$$

for θ_j the true, unknown, value of the factorial effect, $j = 1, \dots, 2^f$. Further more, for $j, l = 1, \dots, 2^f - 1; j \neq l$,

$$\begin{aligned}
\text{cov}(\hat{\theta}_j, \hat{\theta}_l) &= \text{cov} \left(\sum_{i=1}^t c_{ij} \bar{y}_{i.}, \sum_{i=1}^t c_{il} \bar{y}_{i.} \right) \\
&= \sum_{i=1}^t c_{ij} c_{il} \text{var}(\bar{y}_{i.}) \\
&= \frac{\sigma^2}{r} \sum_{i=1}^t c_{ij} c_{il} \\
&= 0,
\end{aligned}$$

as $\sum_{i=1}^t c_{ij} c_{il} = 0$ for $j \neq l$. That is, the factorial contrasts are independent as the contrast coefficient vectors are orthogonal.

Hence, under the null hypothesis $H_0 : \theta_1 = \cdots = \theta_{2^f-1} = 0$ (all factorial effects are zero), the $\hat{\theta}_j$ form a sample from **independent** normally distributed random variables from the distribution

$$\hat{\theta}_j \sim N \left(0, \frac{4\sigma^2}{n} \right), \quad j = 1, \dots, 2^f - 1. \quad (4.3)$$

To assess evidence against H_0 , we can plot the ordered estimates of the factorial effects against the ordered quantiles of a standard normal distribution. Under H_0 , the points in this plot should lie on a straightline (the slope of the line will depend on the unknown σ^2). We anticipate that the majority of the effects will be small (*effect sparsity*), and hence any large effects that lie away from the line are unlikely to come from distribution (4.3) and may be significantly different from zero. We have essentially turned the problem into an outlier identification problem.

For Example 4.1, we can easily produce this plot in R. Table 4.4 gives the ordered factorial effects, which are then plotted against standard normal quantiles in Figure 4.3.

```

effs <- dplyr::arrange(transform(desilylation.effs)[,1:2], dplyr::desc(estimate))
knitr::kable(effs, caption = "Desilylation experiment: sorted estimated factorial effects")
qqnorm(sort(effs[,2]), ylab = "Factorial effects", main = "")
qqline(sort(effs[,2]))

```

In fact, it is more usual to use a **half-normal** plot to assess the size of factorial effects, where we plot the sorted absolute values of the estimated effects against the quantiles of a half-normal distribution³

³The absolute value of a normally distributed random variable follows a half-normal distribution.

Table 4.4: Desilylation experiment: sorted estimated factorial effects

contrast	estimate
temp	8.1200
reagent	3.0875
time	2.5675
temp.solvent	2.3575
solvent.reagent	0.4900
time.solvent	0.4400
temp.time.solvent	0.2450
temp.time.reagent	0.1950
temp.time.solvent.reagent	0.1925
temp.solvent.reagent	-0.0300
time.solvent.reagent	-0.2375
time.reagent	-0.6450
solvent	-2.2175
temp.time	-2.3575
temp.reagent	-2.7725

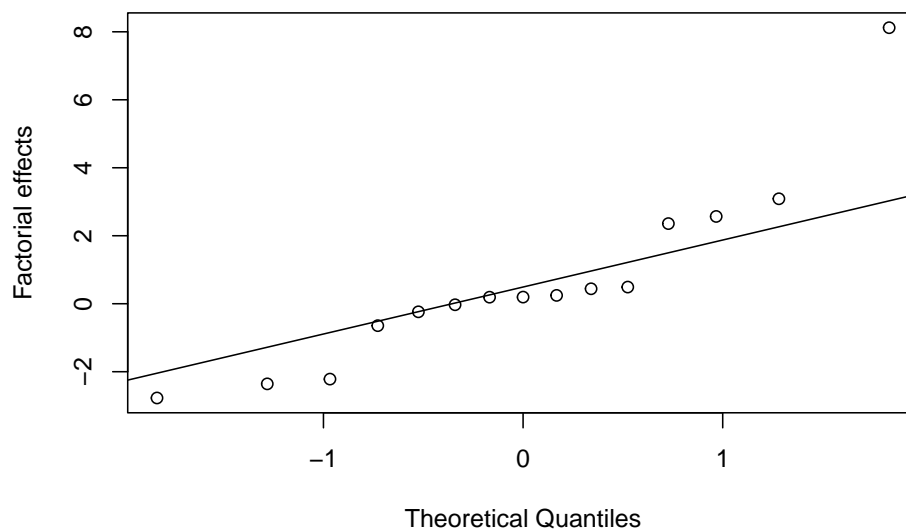


Figure 4.3: Desilylation experiment: normal effects plot

```
p <- .5 + .5 * (1:16 - .5) / 16 # probabilities we will plot against
qqplot(x = qnorm(p), y = sort(abs(effs[,2])), ylab = "Absolute factorial effects",
       xlab = "Half-normal quantiles")
```

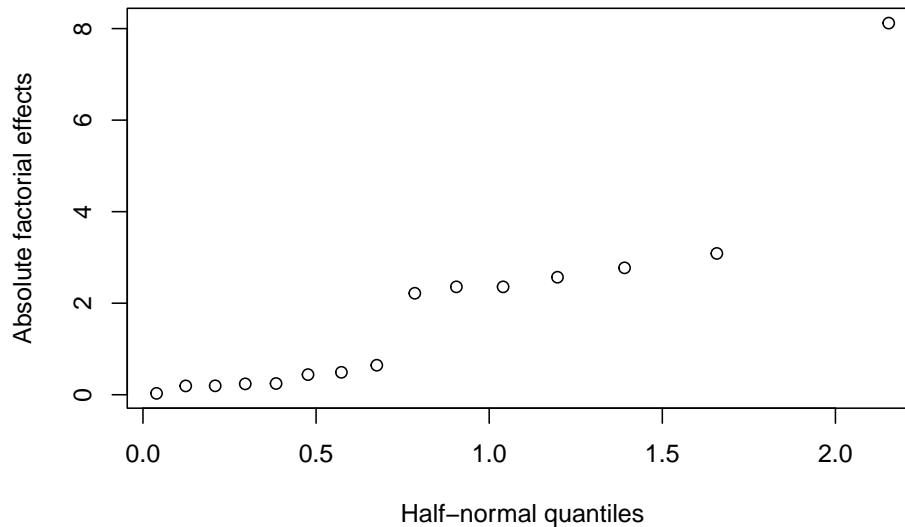


Figure 4.4: Desilylation experiment: half-normal effects plot

The advantage of a half-normal plot such as Figure 4.4 is that we only need to look at effects appearing in the top right corner (significant effects will always appear “above” a hypothetical straight line) and we do not need to worry about comparing large positive and negative values. For these reason, they are usually preferred over normal plots.

For the desilylation experiment, we can see the effects fall into three groups: one effect standing well away from the line, and almost certainly significant (`temp`, from Table ??), then a group of six effects (`reagent`, `time`, `temp.solvent`, `solvent`, `temp.time`, `temp.reagent`) which may be significant, and then a group of 8 much smaller effects.

4.3.1 Lenth’s method for approximate hypothesis testing

The assessment of normal or half-normal effect plots can be quite subjective. Lenth (1989) introduced a simple method for conducting more formal hypothesis testing in unreplicated factorial experiments.

Lenth’s method uses a **pseudo standard error** (PSE):

$$\text{PSE} = 1.5 \times \text{median}_{|\hat{\theta}_i| < 2.5s_0} |\hat{\theta}_i|,$$

where $s_0 = 1.5 \times \text{median}|\hat{\theta}_i|$ is a *consistent*⁴ estimator of the standard deviation of the $\hat{\theta}_i$ under $H_0 : \theta_1 = \dots = \theta_{2^f-1} = 0$. The PSE trims approximately 1%⁵ of the $\hat{\theta}_i$ to produce a **robust** estimator of the standard deviation, in the sense that it is not influenced by large $\hat{\theta}_i$ belonging to important effects.

For Example 4.1, we can construct the PSE as follows.

```
s0 <- 1.5 * median(abs(effs[, 2]))
trimmed <- abs(effs[, 2]) < 2.5 * s0
pse <- 1.5 * median(abs(effs[trimmed, 2]))
pse
```

```
## [1] 0.66
```

The PSE can be used to construct test statistics

$$t_{\text{PSE},i} = \frac{\hat{\theta}_i}{\text{PSE}},$$

which mimic the usual t -statistics used when an estimate of σ^2 is available. These quantities can be compared to reference distribution which was tabulated by Lenth (1989) and simulated in R using the `unrepx` package.

```
eff_est <- effs[, 2]
names(eff_est) <- effs[, 1]
lenth_tests <- unrepx::eff.test(eff_est, method = "Lenth")
knitr::kable(lenth_tests, caption = "Desilylation experiment: hypothesis tests using L
```

The function `eff.test` calculates unadjusted p-values (`p.value`) and simultaneous p-values (`simult.pval`) adjusted to account for multiple testing. Using the latter, from Table 4.5 we see that the main effects of **temp** and **reagent** are significant at the experiment-wise 5% level and, obeying *effect heredity*, their interaction (the p-value is borderline, and hovers around 0.05 depending on simulation error).

The package `unrepx` also provides the function `hnplot` to display these results graphically by adding a reference line to a half-normal plot; see Figure ???. The ME and SME lines indicate the absolute size of effects that would be required to reject $H_0 : \theta_i = 0$ at an individual or experimentwise 100% level, respectively.

```
unrepx::hnplot(eff_est, method = "Lenth", horiz = F, ID = 2.7, alpha = 0.05)
```

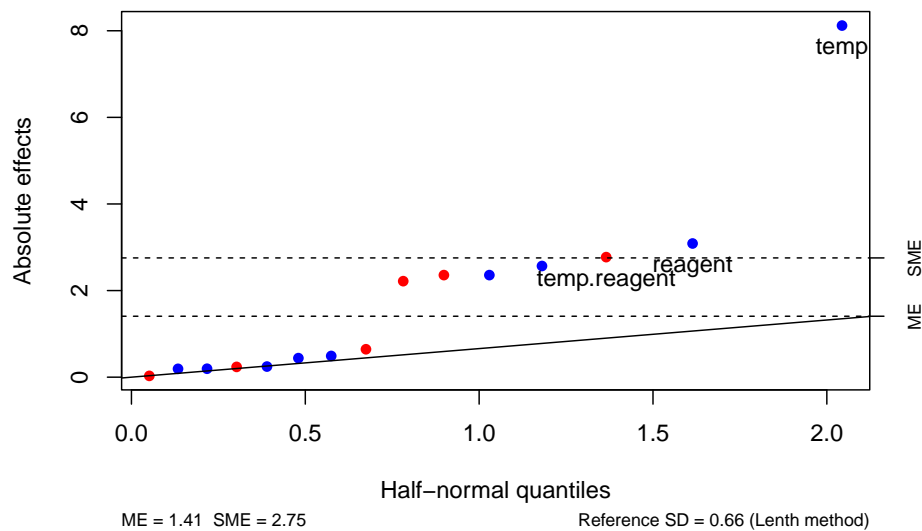
⁴Essentially, s_0 tends in probability to σ as the number of factorial effects tends to infinity.

⁵Under H_0 , the $\hat{\theta}_i$ come from a mean-zero normal distribution, and about 1% of deviates fall outside $\pm 2.57\sigma^2$.

4.3. NORMAL EFFECT PLOTS FOR UNREPLICATED FACTORIAL DESIGNS105

Table 4.5: Desilylation experiment: hypothesis tests using Lenth's method.

	effect	Lenth_PSE	t.ratio	p.value	simult.pval
temp	8.1200	0.66	12.303	0.0000	0.0007
reagent	3.0875	0.66	4.678	0.0037	0.0337
temp.reagent	-2.7725	0.66	-4.201	0.0051	0.0476
time	2.5675	0.66	3.890	0.0071	0.0682
temp.solvent	2.3575	0.66	3.572	0.0102	0.0967
temp.time	-2.3575	0.66	-3.572	0.0102	0.0967
solvent	-2.2175	0.66	-3.360	0.0125	0.1166
time.reagent	-0.6450	0.66	-0.977	0.3037	0.9936
solvent.reagent	0.4900	0.66	0.742	0.4296	1.0000
time.solvent	0.4400	0.66	0.667	0.5393	1.0000
temp.time.solvent	0.2450	0.66	0.371	0.7300	1.0000
time.solvent.reagent	-0.2375	0.66	-0.360	0.7374	1.0000
temp.time.reagent	0.1950	0.66	0.295	0.7834	1.0000
temp.time.solvent.reagent	0.1925	0.66	0.292	0.7856	1.0000
temp.solvent.reagent	-0.0300	0.66	-0.045	0.9665	1.0000



Informally, factorial effects with estimates greater than SME are thought highly likely to be significant, and effects between ME and SME are considered somewhat likely to be significant (and still worthy of further investigation if the budget allows).

4.4 Regression modelling for factorial experiments

We have identified $d = 2^f - 1$ factorial effects that we wish to estimate from our experiment. As $d < n = 2^f r$, even for an unreplicated design with $r = 1$, we can estimate these factorial effects using a full-rank linear regression model.

Let $t \times d$ matrix C hold each factorial contrast as a column. Then

$$\hat{\boldsymbol{\theta}} = C^T \bar{\mathbf{y}},$$

with $\hat{\boldsymbol{\theta}}^T = (\hat{\theta}_1, \dots, \hat{\theta}_d)$ being the vector of estimated factorial effects and $\bar{\mathbf{y}}^T = (\bar{y}_1, \dots, \bar{y}_t)$ being the vector of treatment means.

We can define an $n \times d$ expanded contrast matrix as $\tilde{C} = C \otimes \mathbf{1}_r$, where each row of \tilde{C} gives the contrast coefficients for each run of the experiment. Then,

$$\hat{\boldsymbol{\theta}} = \frac{1}{r} \tilde{C}^T \mathbf{y}.$$

To illustrate, we will imagine a hypothetical version of Example 4.1 where each treatment was repeated three times (with $y_{i1} = y_{i2} = y_{i3}$).

```
y <- kronecker(desilylation$yield, rep(1, 3)) # hypothetical response vector
C <- factorial_contrasts
Ctilde <- kronecker(C, rep(1, 3))
t(Ctilde) %*% y / 3 # to check
```

```
##      [,1]
## [1,]  8.1200
## [2,]  2.5675
## [3,] -2.2175
## [4,]  3.0875
## [5,] -2.3575
## [6,]  2.3575
## [7,] -2.7725
## [8,]  0.4400
## [9,] -0.6450
## [10,] 0.4900
## [11,] 0.2450
## [12,] 0.1950
## [13,] -0.0300
## [14,] -0.2375
## [15,] 0.1925
```

If we define a model matrix $X = \frac{2^f}{2} \tilde{C}$, then X is a $n \times d$ matrix with entries ± 1 and columns equal to unscaled factorial contrasts. Then

$$(X^T X)^{-1} X^T \mathbf{y} = \frac{1}{n} \times \frac{2^f}{2} \tilde{C}^T \mathbf{y} \quad (4.4)$$

$$= \frac{1}{2r} \tilde{C}^T \mathbf{y} \quad (4.5)$$

$$= \frac{1}{2} \hat{\boldsymbol{\theta}}. \quad (4.6)$$

$$(4.7)$$

The left-hand side of equation (4.4) is the least squares estimator $\hat{\boldsymbol{\beta}}$ from the model

$$\mathbf{y} = \mathbf{1}_n \beta_0 + X\boldsymbol{\beta} + \boldsymbol{\varepsilon},$$

where \mathbf{y} is the response vector and $\boldsymbol{\varepsilon}$ the error vector from unit-treatment model (4.1). We have simply re-expressed the mean response as $\mu + \tau_i = \beta_0 + \mathbf{x}_i^T \boldsymbol{\beta}$, where d -vector \mathbf{x}_i holds the unscaled contrast coefficients for the main effects and interactions.

We can illustrate these connections for Example 4.1.

```
X <- 8 * C
Xt <- t(X)
XtX <- Xt %*% X
2 * solve(XtX) %*% t(X) %*% desilylation$yield
```

```
##           [,1]
## temp      8.1200
## time      2.5675
## solvent   -2.2175
## reagent    3.0875
## temp:time  -2.3575
## temp:solvent  2.3575
## temp:reagent -2.7725
## time:solvent  0.4400
## time:reagent -0.6450
## solvent:reagent  0.4900
## temp:time:solvent  0.2450
## temp:time:reagent  0.1950
## temp:solvent:reagent -0.0300
## time:solvent:reagent -0.2375
## temp:time:solvent:reagent  0.1925
```

The more usual way to think about this modelling approach is as a regression model with f (quantitative⁶) variables, labelled x_1, \dots, x_{2^f-1} , scaled to lie in the

⁶When qualitative factors only have two levels, each regression term only has 1 degree of freedom, and so there is practically little difference from a quantitative variable.

Table 4.6: Desilylation example: factorial effects calculated using a regression model.

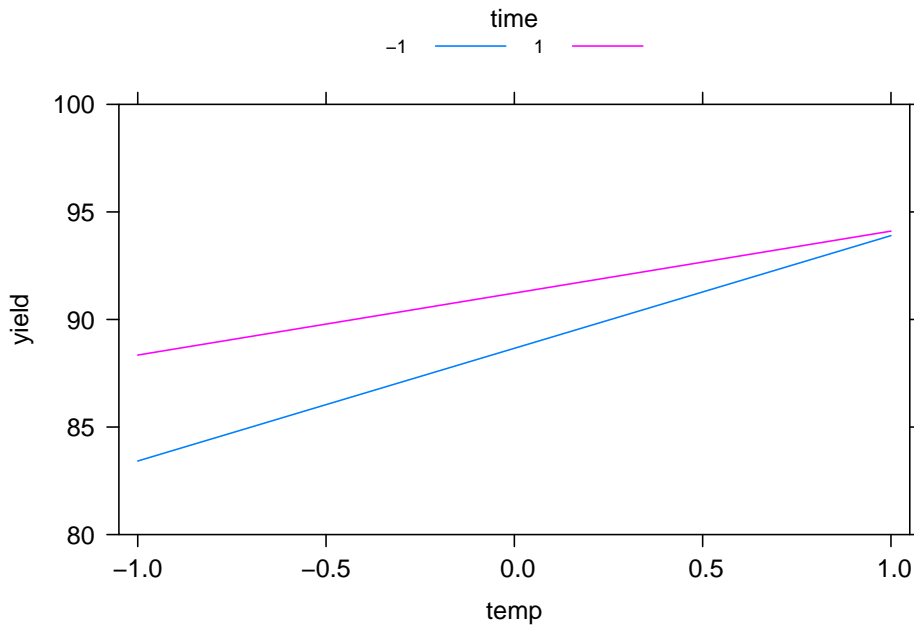
	x
temp	8.1200
time	2.5675
solvent	-2.2175
reagent	3.0875
temp:time	-2.3575
temp:solvent	2.3575
temp:reagent	-2.7725
time:solvent	0.4400
time:reagent	-0.6450
solvent:reagent	0.4900
temp:time:solvent	0.2450
temp:time:reagent	0.1950
temp:solvent:reagent	-0.0300
time:solvent:reagent	-0.2375
temp:time:solvent:reagent	0.1925

interval $[-1, 1]$ (in fact, they just take values ± 1). We can then fit a regression model in these variables, and include products of these variables to represent interactions. We usually also include the intercept term. For Example 4.1:

```
desilylation.df <- dplyr::mutate(desilylation,
                                across(.cols = temp:reagent,
                                       ~ as.numeric(as.character(.x))))
desilylation.df <- dplyr::select(desilylation.df, -c(trt))
desilylation.df <- dplyr::mutate(desilylation.df, across(.cols = temp:reagent,
                                                         ~ scales::rescale(.x, to = c(-1, 1))))
desilylation_reg.lm <- lm(yield ~ (.) ^ 4, data = desilylation.df)
knitr::kable(2 * coef(desilylation_reg.lm)[-1], caption = "Desilylation example: factor
```

A regression modelling approach is usually more straightforward to apply than defining contrasts in the unit-treatment model, and makes clearer the connection between interaction contrasts and products of main effect contrasts (automatically defined in a regression model). It also enables us to make use of the **effects** package in R to quickly produce main effect and interaction plots.

```
temp_x_time <- effects::Effect(c("temp", "time"), desilylation_reg.lm, xlevels = list(
plot(temp_x_time, main = "", rug = F, x.var = "temp", ylim = c(80, 100))
```



4.4.1 ANOVA for factorial experiments

The basic ANOVA table has the following form.

Table 4.7: The ANOVA table for a full factorial experiment

Source	Degress of Freedom	(Sequential) Sum of Squares	Mean Square
Regression	$2^f - 1$	$\sum_{j=1}^{2^f-1} n\hat{\beta}_j^2 - n\bar{y}^2$	Reg SS/ $(2^f - 1)$
Residual	$2^f(r - 1)$	$(\mathbf{Y} - X\hat{\beta})^T(\mathbf{Y} - X\hat{\beta})$	RSS/ $(2^f(r - 1))$
Total	$2^f r - 1$	$\mathbf{Y}^T \mathbf{Y} - n\bar{Y}^2$	

The regression sum of squares for a factorial experiment has a very simple form. If we include an intercept column in X , from Section 1.5.2,

$$\begin{aligned}
\text{Regression SS} &= \text{RSS}(\text{null}) - \text{RSS} \\
&= \hat{\beta}^T X^T X \hat{\beta} - n\bar{y}^2 \\
&= \sum_{j=1}^{2^f-1} n\hat{\beta}_j^2 - n\bar{y}^2,
\end{aligned}$$

as $X^T X = nI_{2^f}$. Hence, the j th factorial effect contributes $n\hat{\beta}_j^2$ to the regression sum of squares, and this quantity can be used to construct a test statistic if $r > 1$ and hence an estimate of σ^2 is available. For Example 4.1, the ANOVA table is as follows.

```
ss <- nrow(desilylation) * coef(desilylation_reg.lm)^2
regss <- sum(ss) - nrow(desilylation) * mean(desilylation$yield)^2
regss
```

```
## [1] 427.3
```

```
ss[-1]
```

##	temp	time	solvent
##	263.7376	26.3682	19.6692
##	reagent	temp:time	temp:solvent
##	38.1306	22.2312	22.2312
##	temp:reagent	time:solvent	time:reagent
##	30.7470	0.7744	1.6641
##	solvent:reagent	temp:time:solvent	temp:time:reagent
##	0.9604	0.2401	0.1521
##	temp:solvent:reagent	time:solvent:reagent	temp:time:solvent:reagent
##	0.0036	0.2256	0.1482

```
anova(desilylation_reg.lm)
```

```
## Analysis of Variance Table
```

```
##
```

```
## Response: yield
```

##	Df	Sum Sq	Mean Sq	F value	Pr(>F)
## temp	1	263.7	263.7	NaN	NaN
## time	1	26.4	26.4	NaN	NaN
## solvent	1	19.7	19.7	NaN	NaN
## reagent	1	38.1	38.1	NaN	NaN
## temp:time	1	22.2	22.2	NaN	NaN
## temp:solvent	1	22.2	22.2	NaN	NaN
## temp:reagent	1	30.7	30.7	NaN	NaN
## time:solvent	1	0.8	0.8	NaN	NaN
## time:reagent	1	1.7	1.7	NaN	NaN

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## solvent:reagent	1	1.0	1.0	NaN	NaN
## temp:time:solvent	1	0.2	0.2	NaN	NaN
## temp:time:reagent	1	0.2	0.2	NaN	NaN
## temp:solvent:reagent	1	0.0	0.0	NaN	NaN
## time:solvent:reagent	1	0.2	0.2	NaN	NaN
## temp:time:solvent:reagent	1	0.1	0.1	NaN	NaN
## Residuals	0	0.0	NaN		

Chapter 5

Blocking in factorial designs

Chapter 6

Fractional factorial designs

Chapter 7

Response surface methodology

Chapter 8

Optimal design of experiments

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