

Further Statistical Analysis using R

Mark Dunning, Matthew Eldridge and Sarah Vowler

Last modified: 16 Oct 2015

Contents

1	Introduction	1
2	One-Way ANOVA	1
3	Other Non-Parametric tests	4
4	Breast Cancer Incidence	6
5	Linear Regression	8

1 Introduction

In this practical, we will use several 'real-life' datasets to demonstrate some of the concepts you have seen in the lectures. We will guide you through how to analyse these datasets in R and the kinds of questions you should be asking yourself when faced with similar data. As this is not a course in using R, we will provide the R code that you will need. However, it is up to you to think carefully about the assumptions of a statistics tests and interpret the results that R gives.

The data you need for this practical are available as a <https://rawgit.com/bioinformatics-core-shared-training/intermediate-stats/master/CourseData.zip>. Please download this zip file and extract to a directory on your laptop. You will then need to tell RStudio where to find these files by setting your *working directory*

Session -> Set Working Directory -> Choose Directory

2 One-Way ANOVA

The data for this exercise are to be found in `amess.csv`. The data are the red cell folate levels in three groups of cardiac bypass patients given different levels of nitrous oxide (N_2O) and oxygen (O_2) ventilation. [There is a reference to the source of this data in Altman, Practical Statistics for Medical Research, p. 208.]

The treatments are

- i) 50% N_2O and 50% O_2 continuously for 24 hours
- ii) 50% N_2O and 50% O_2 during the operation
- iii) No N_2O but 35-50% O_2 continuously for 24 hours

1. Import the file `amess.csv` into R. Verify that the dimensions of the object that R creates are correct.

```
amess <- read.csv("amess.csv")
dim(amess)
```

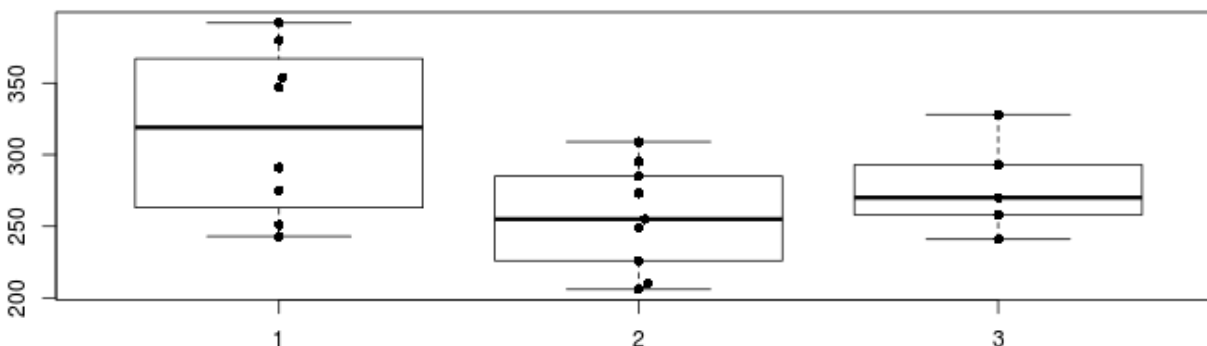
```
## [1] 22 2
```

NOTE: You can use the `View` function in RStudio to bring-up a display of the object you have created

2. Draw boxplots of the data. Does it look as though the assumptions for ANOVA are satisfied?

NOTE: It is often useful to overlay individual points on the boxplot

```
library(beeswarm)
boxplot(folate~treatmnt,data=amess)
beeswarm(folate~treatmnt,data=amess,add=TRUE,pch=16)
```



3. Perform t-tests for each of the three pair-wise comparisons. Make note of the t-test statistics obtained. Which groups, if any, differ from one another?

NOTE: If you have not performed a t test in R before, you may wish to consult the help page for `t.test`; `?t.test`.

```
t.test(folate~ treatmnt,data=amess[amess$treatmnt !=3,])
```

```
##
## Welch Two Sample t-test
##
## data: folate by treatmnt
## t = 2.4901, df = 11.579, p-value = 0.02906
## alternative hypothesis: true difference in means is not equal to 0
## 95 percent confidence interval:
## 7.310453 113.050658
## sample estimates:
## mean in group 1 mean in group 2
## 316.6250 256.4444
```

```
t.test(folate~treatmnt,data=amess[amess$treatmnt !=2,])
```

```
##
## Welch Two Sample t-test
##
## data: folate by treatmnt
## t = 1.5048, df = 10.985, p-value = 0.1606
## alternative hypothesis: true difference in means is not equal to 0
## 95 percent confidence interval:
## -17.87994 95.12994
## sample estimates:
## mean in group 1 mean in group 3
## 316.625 278.000
```

```
t.test(folate~treatmnt,data=amess[amess$treatmnt !=3,])
```

```
##
## Welch Two Sample t-test
##
## data: folate by treatmnt
## t = 2.4901, df = 11.579, p-value = 0.02906
## alternative hypothesis: true difference in means is not equal to 0
## 95 percent confidence interval:
## 7.310453 113.050658
## sample estimates:
## mean in group 1 mean in group 2
## 316.6250 256.4444
```

4. Perform an analysis of variance on the data. Why is this a more valid analysis? Interpret the output.

```
mod <- aov(folate~factor(treatmnt),data=amess)
mod
```

```
## Call:
## aov(formula = folate ~ factor(treatmnt), data = amess)
##
## Terms:
##          factor(treatmnt) Residuals
## Sum of Squares      15515.77  39716.10
## Deg. of Freedom           2       19
##
## Residual standard error: 45.72003
## Estimated effects may be unbalanced
```

```
summary(aov(mod))
```

```
##          Df Sum Sq Mean Sq F value Pr(>F)
## factor(treatmnt)  2  15516    7758   3.711 0.0436 *
## Residuals      19  39716    2090
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

5. Choose any pair of treatments and repeat the ANOVA for just these two treatments. Compare the test statistic value to the results of the t-test from question 3. What do you notice?

```
subset <- amess[amess$treatmnt != 3,]
t.test(folate~treatmnt,data=subset)
```

```
##
## Welch Two Sample t-test
##
## data: folate by treatmnt
## t = 2.4901, df = 11.579, p-value = 0.02906
## alternative hypothesis: true difference in means is not equal to 0
## 95 percent confidence interval:
## 7.310453 113.050658
## sample estimates:
## mean in group 1 mean in group 2
## 316.6250 256.4444
```

6. If the assumptions for ANOVA don't hold, then there are non-parametric alternatives available.

- If the assumption of normality doesn't hold then we might look at the Kruskal-Wallis test.

- If the assumption of equal variances doesn't hold then we might use the median test.

Perform these tests. Do they give the same answer as ANOVA? If not, why not? How important were the assumptions?

```
kruskal.test(folate~factor(treatmnt),data=amess)

##
##  Kruskal-Wallis rank sum test
##
## data:  folate by factor(treatmnt)
## Kruskal-Wallis chi-squared = 4.1852, df = 2, p-value = 0.1234

library(RVAideMemoire)
mood.medtest(folate~factor(treatmnt),data=amess)
```

```
##
##  Mood's median test
##
## data:  folate by factor(treatmnt)
## p-value = 0.2332
```

7. Use a *post-hoc* test to compute p-value for all pairwise contrasts. Unlike the t-tests performed in Question 3, the p-values returned here are adjusted for multiple testing. Two functions for doing such tests in R are TukeyHSD and pairwise.t.test. Help for both these functions is available through RStudio; ?TukeyHSD, ?pairwise.t.test

```
TukeyHSD(mod)

##  Tukey multiple comparisons of means
##    95% family-wise confidence level
##
## Fit: aov(formula = folate ~ factor(treatmnt), data = amess)
##
## $`factor(treatmnt)`
##      diff      lwr      upr      p adj
## 2-1 -60.18056 -116.61904 -3.74207 0.0354792
## 3-1 -38.62500 -104.84037 27.59037 0.3214767
## 3-2 21.55556 -43.22951 86.34062 0.6802018

pairwise.t.test(amess$folate,amess$treatmnt)

##
##  Pairwise comparisons using t tests with pooled SD
##
## data:  amess$folate and amess$treatmnt
##
##      1      2
## 2 0.042 -
## 3 0.310 0.408
##
## P value adjustment method: holm
```

3 Other Non-Parametric tests

The file `treatments.txt` records measurements from five subjects undergoing a treatment course. The data were taken from an example in *GraphPad Prism*.

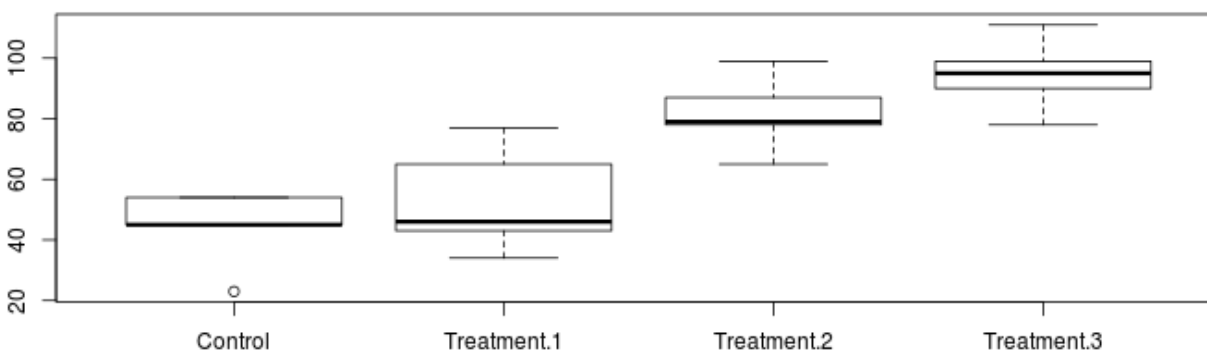
1. Import these data into R and visualise the distribution of the data

```
treatments <- read.delim("treatments.txt")
head(treatments)
```

```
##      X Control Treatment.1 Treatment.2 Treatment.3
## 1 GS      54          43          78          111
## 2 JM      23          34          65          99
## 3 HM      45          65          99          78
## 4 DR      54          77          79          90
## 5 PS      45          46          87          95
```

2. Use the Friedman test to assess the effect of treatment and interpret the result. Can you justify why this test was chosen?

```
boxplot(treatments[, -1])
```



```
friedman.test(as.matrix(treatments[, -1]))
```

```
##
## Friedman rank sum test
##
## data: as.matrix(treatments[, -1])
## Friedman chi-squared = 13.08, df = 3, p-value = 0.004467
```

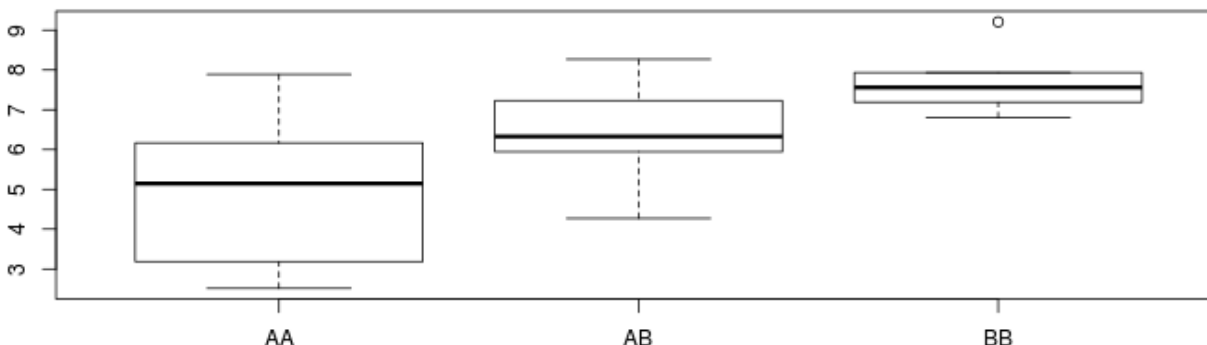
The file `genotypes.txt` contains measurements from patients in a gene expression study. For each patient in the study, their genotype for a particular gene was recorded.

3. Import the file `genotypes.txt` into R and visualise the data using a boxplot

```
gtypes <- read.delim("genotypes.txt")
gtypes
```

```
##      AA      AB      BB
## 1 2.513047 6.328862      NA
## 2 6.168767 5.607576 7.639488
## 3 3.184589 8.269598 6.795799
## 4 7.889960 4.271390 7.188640
## 5 5.146395 6.282917 7.482058
## 6      NA 7.274772 7.934725
## 7      NA 7.184816 9.208339
```

```
boxplot(gtypes)
```



NOTE: The `boxplot` function is also able to accept a data frame as input. In this case, it will plot the distribution of data in each column separately

4. Transform the data into two columns; one to indicate the genotype of the individual and the second for the gene expression level.

```
library(tidyr)
gtypes <- gather(gtypes)
gtypes$key <- rep(c(1,2,3),each=7)
```

5. Use an appropriate method to test the association between gene expression and genotype

```
library(clinfun)
jonckheere.test(gtypes$value,gtypes$key)
```

```
##
## Jonckheere-Terpstra test
##
## data:
## JT = 86, p-value = 0.007535
## alternative hypothesis: two.sided
```

4 Breast Cancer Incidence

The file `globalBreastCancerRisk.csv` gives the number of new cases of Breast Cancer (per population of 10,000) in various countries around the world, along with various health and lifestyle risk factors. These data were collected from the [gapminder](#) resource ¹. Let's suppose we are initially interested in whether the number of breast cancer cases is significantly different in different regions of the world.

1. Read these data into R

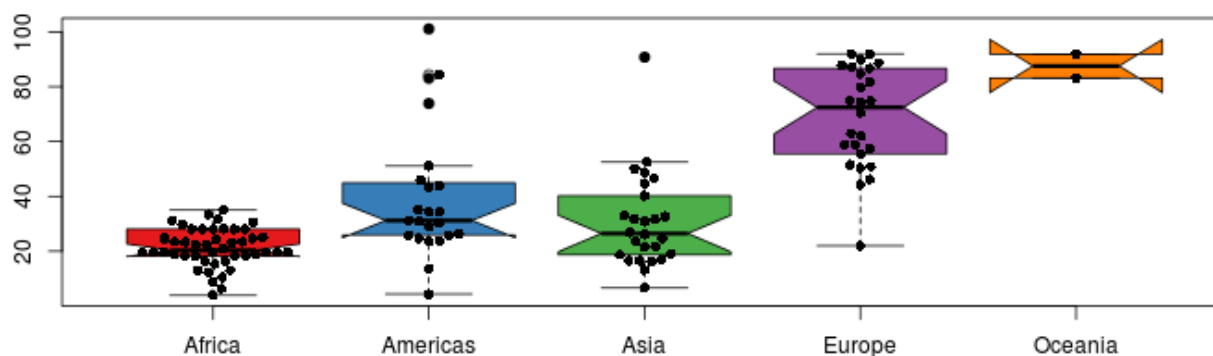
```
bcStats <- read.csv("globalBreastCancerRisk.csv")
head(bcStats)
```

¹Similar data are also detailed in a blog post at: <http://www.analyticsforfun.com/2014/06/performing-anova-test-in-r-results-and.html>

```
##      country continent year lifeExp      pop  gdpPercap
## 1 Afghanistan      Asia 2002  42.129 25268405   726.7341
## 2   Albania      Europe 2002  75.651  3508512  4604.2117
## 3   Algeria      Africa 2002  70.994 31287142  5288.0404
## 4    Angola      Africa 2002  41.003 10866106  2773.2873
## 5 Argentina Americas 2002  74.340 38331121  8797.6407
## 6  Australia Oceania 2002  80.370 19546792 30687.7547
## NewCasesOfBreastCancerIn2002 AlcoholConsumption BloodPressure
## 1                26.8                0.02         124.2085
## 2                57.4                6.68         129.0609
## 3                23.5                0.96         130.4024
## 4                23.1                5.40         129.9282
## 5                73.9               10.00         119.6538
## 6                83.2               10.02         120.5113
## BodyMassIndex Cholestrol Smoking
## 1      20.65274      4.295170      NA
## 2      25.27082      4.918646      4.0
## 3      25.69948      4.848951      0.3
## 4      22.26093      4.499115      NA
## 5      26.70460      5.143871     25.4
## 6      26.25957      5.326858     21.8
```

- Visualise the distribution of breast cancer incidence in each continent. Include a way of identifying how many observations belong to each group (continent). Are there any groups that you would consider removing?

```
library(RColorBrewer)
library(beeswarm)
boxplot(NewCasesOfBreastCancerIn2002~continent,data=bcStats,col=brewer.pal(5,"Set1"),notch=TRUE)
beeswarm(NewCasesOfBreastCancerIn2002~continent,data=bcStats,add=TRUE,pch=16)
```



```
bcStats.clean <- bcStats[bcStats$continent != "Oceania",]
```

- Would a parametric, or non-parametric, approach be suitable for this analysis? Use the result from a `bartlett.test` to support your answer. Proceed with your chosen approach to test the hypothesis that breast cancer incidence is different across the globe.

```
bartlett.test(NewCasesOfBreastCancerIn2002~continent,data=bcStats.clean)
```

```
##
```

```
## Bartlett test of homogeneity of variances
##
## data: NewCasesOfBreastCancerIn2002 by continent
## Bartlett's K-squared = 48.26, df = 3, p-value = 1.875e-10
kruskal.test(NewCasesOfBreastCancerIn2002~continent,data=bcStats.clean)
```

```
##
## Kruskal-Wallis rank sum test
##
## data: NewCasesOfBreastCancerIn2002 by continent
## Kruskal-Wallis chi-squared = 59.896, df = 3, p-value = 6.185e-13
```

4. In a manner similar to the *One-Way ANOVA* example, we can apply multiple-testing correction to all pairwise contrasts and assess which individual contrasts are significant. Which pairs of continents have significantly different breast cancer incidence rates to each other?

```
pairwise.wilcox.test(bcStats.clean$NewCasesOfBreastCancerIn2002,bcStats.clean$continent)
```

```
##
## Pairwise comparisons using Wilcoxon rank sum test
##
## data: bcStats.clean$NewCasesOfBreastCancerIn2002 and bcStats.clean$continent
##
##          Africa Americas Asia
## Americas 5.4e-05 -          -
## Asia      0.04232 0.14359 -
## Europe    8.8e-11 0.00016 7.9e-07
##
## P value adjustment method: holm
```

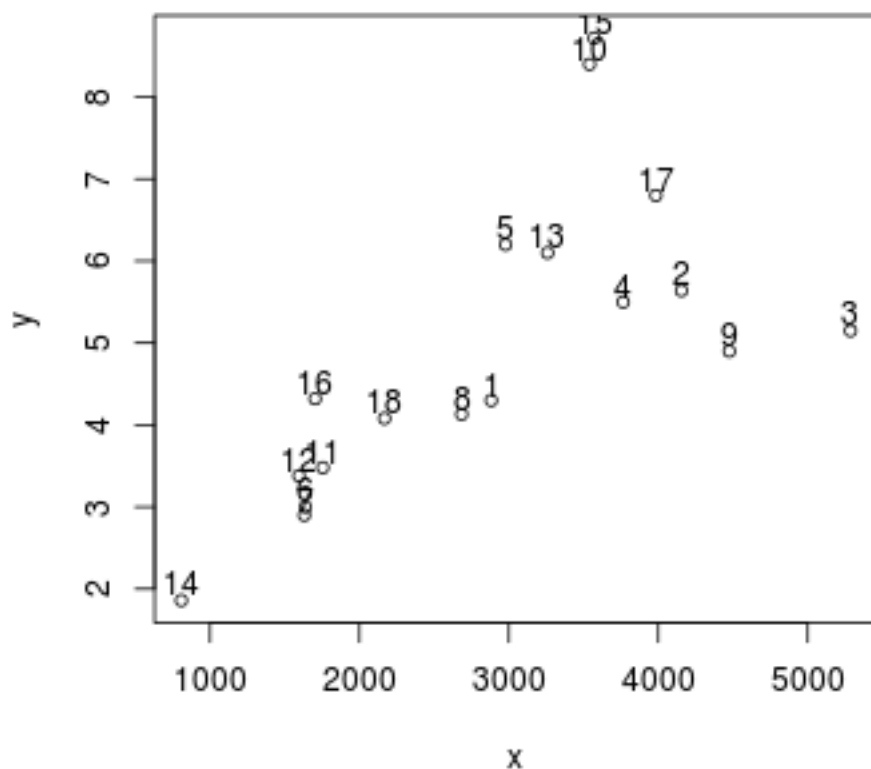
5 Linear Regression

1. Clinical Trial Data: A researcher wished to study the effect of a particular drug X on the immune system. Thus, the concentration of the drug was measured on various days, along with the white blood cell counts.
 - a) Import the clinical data from the file `ClinicalTrials.txt`
 - b) Make a scatter plot
 - c) Label each point according to the date that the measurement was made
 - d) Fit a linear model to the data and produce the diagnostic plots. Which observations fit the model least-well? Compare your answers to the scatter plot
 - e) Overlay the line-of-best fit on the scatter plot. Indicate the differences between each fitted and observed value.
 - f) Extract the R^2 value from the model summary, and print this value on the plot

```
trials <- read.delim("clinicalTrials.txt")
```

```
x <- trials$Drug.concentration
y <- trials$Cell.Count
```

```
plot(x,y,cex=0.8)
text(x,y+0.2,labels=1:length(x))
```

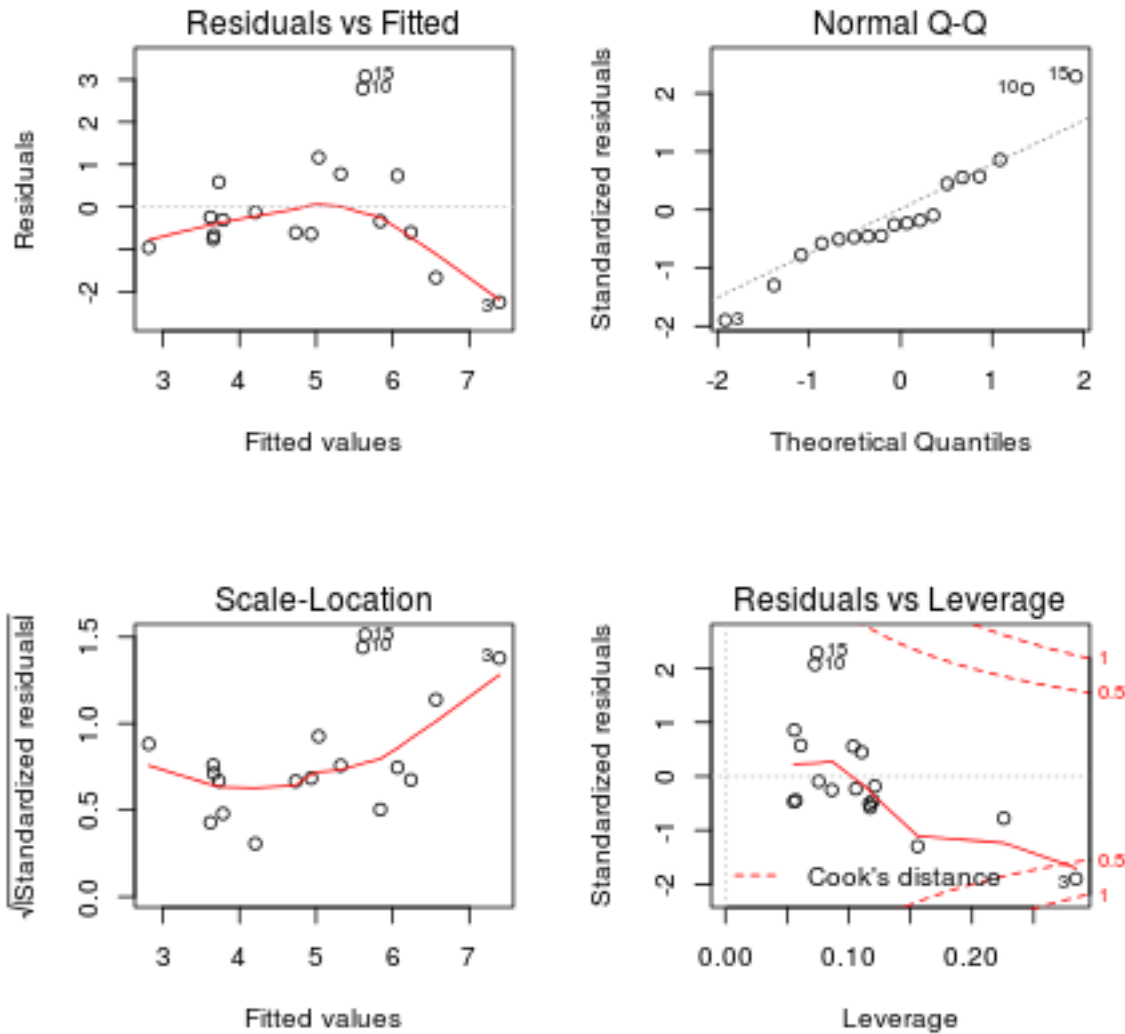
NOTE: We use the `text` function to overlay the number of each observation on-top of the scatter plot that we create initially. `text` requires a set of `x` and `y` coordinates. Here we add an offset of 0.2 to make sure the labels are not obscured by the points

```
model <- lm(y ~ x )
summary(model)
```

```
##
## Call:
## lm(formula = y ~ x)
##
## Residuals:
##      Min       1Q   Median       3Q      Max
## -2.2437 -0.6580 -0.3219  0.6999  3.0792
##
## Coefficients:
##              Estimate Std. Error t value Pr(>|t|)
## (Intercept)  1.9910174  0.8666011   2.298  0.03542 *
## x            0.0010221  0.0002781   3.675  0.00205 **
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 1.398 on 16 degrees of freedom
## Multiple R-squared:  0.4577, Adjusted R-squared:  0.4239
```

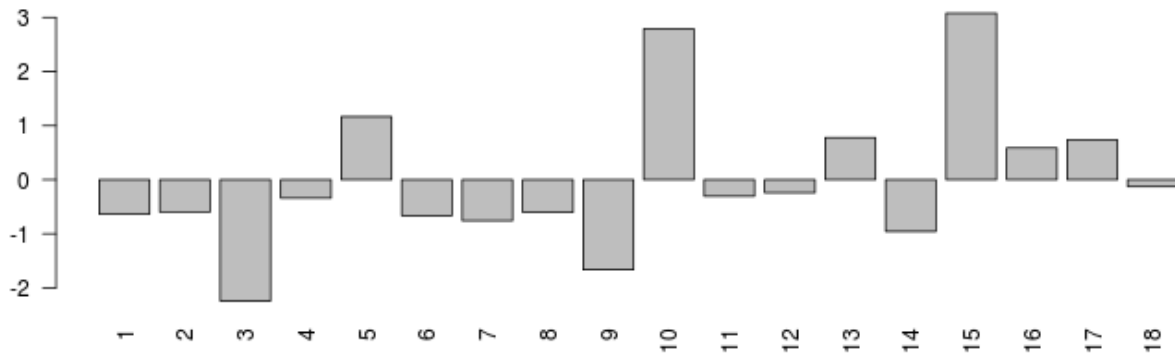
```
## F-statistic: 13.51 on 1 and 16 DF, p-value: 0.002047
```

```
par(mfrow=c(2,2))
plot(model)
```



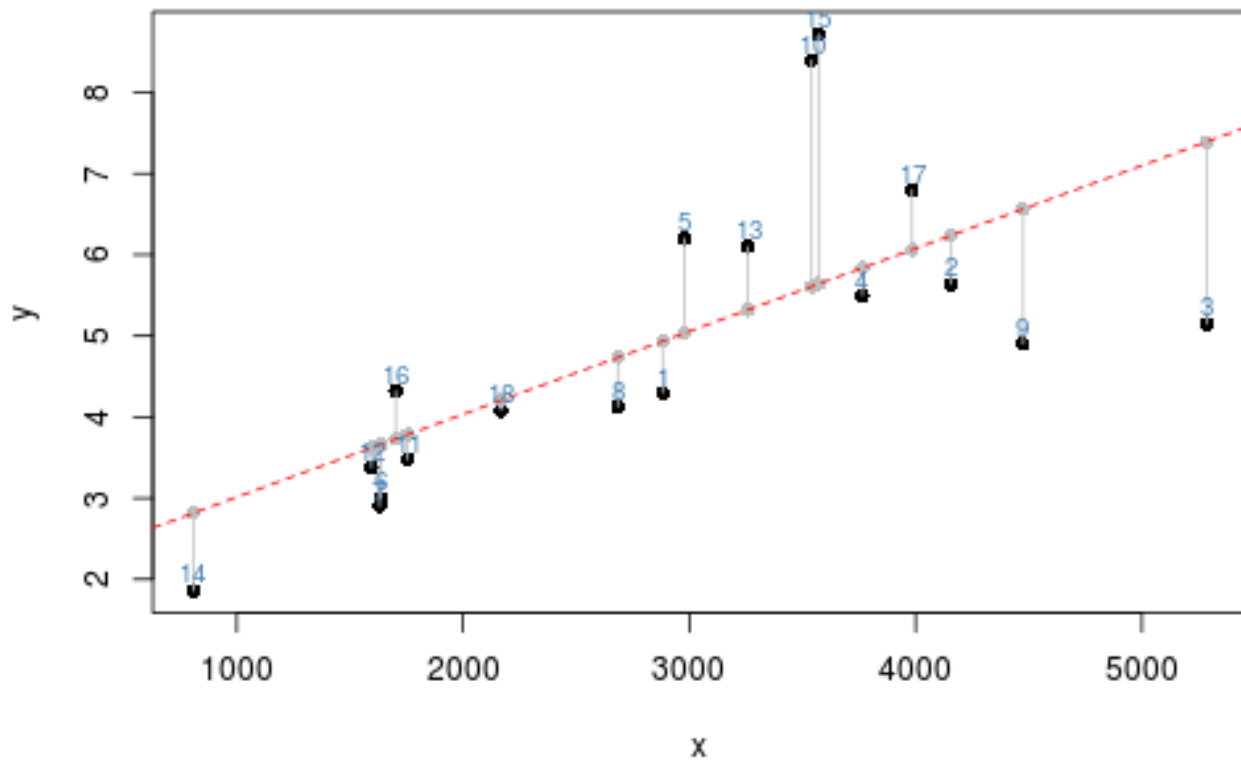
NOTE: We modify the plot layout by changing the `mfrow` parameter to allow for all the diagnostic plots to be shown in the same plotting device

```
par(mfrow=c(1,1))
barplot(model$residuals,names=1:length(x),las=2)
```



NOTE: The setting of the `las` parameter to 2 arranges the labels so they are perpendicular to the x-axis

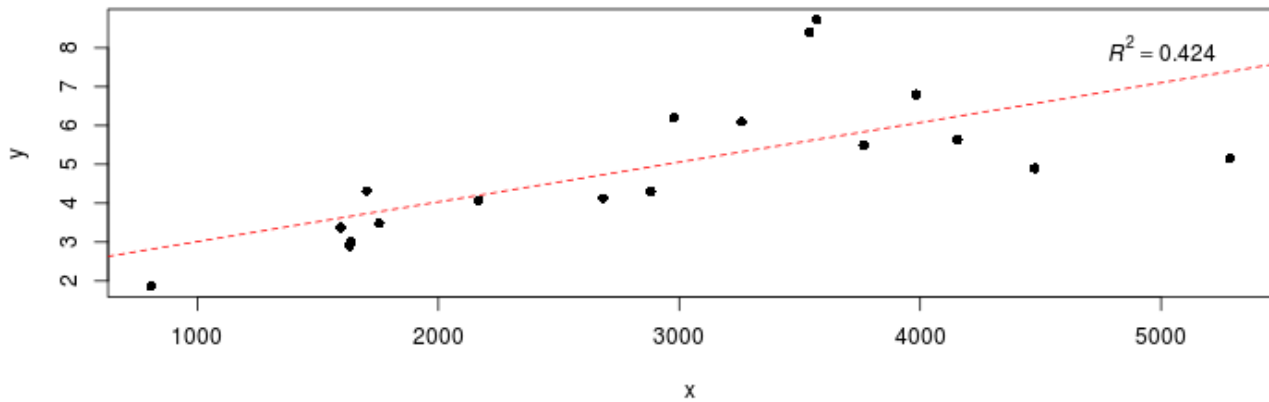
```
par(mfrow=c(1,1))
plot(x,y,pch=16)
points(x,model$fitted.values,pch=16,col="grey")
arrows(x,model$fitted.values,x,y,code=0,col="grey")
abline(model,col="red",lty=2)
text(x,y+0.2,labels=1:length(x),cex=0.8,col="steelblue")
```



NOTE: The `arrows` function was used to draw vertical lines on the plot. Each line is defined by a set of starting and ending x and y positions. In this case, we want the start and end positions to be the same and to vary the y coordinates

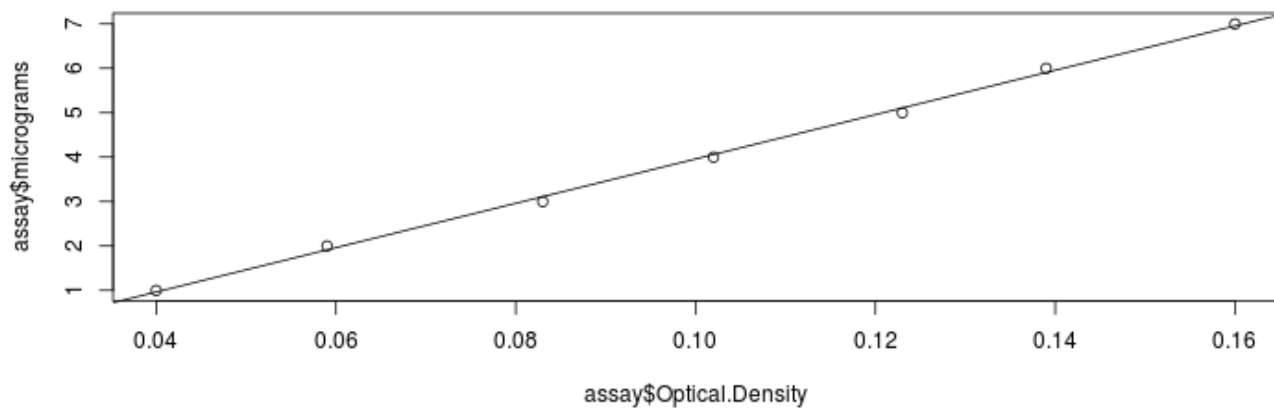
```
plot(x,y,pch=16)
abline(model,col="red",lty=2)
result <- summary(model)
r2 <- round(result$adj.r.squared,3)
my.p = result$coefficients[2,4]

mylabel = bquote(italic(R)^2 == .(format(r2, digits = 3)))
text(x = 5000, y = 8, labels = mylabel)
```



2. Interpolating Unknown values

```
assay <- read.delim("Assay.txt")
model <- lm(micrograms~Optical.Density,data=assay)
plot(assay$Optical.Density, assay$micrograms)
abline(model)
```



```
y <- assay$micrograms
x <- assay$Optical.Density

topredict <- which(is.na(y))
new <- data.frame(x = x[which(is.na(y))])
y[which(is.na(y))] <- predict(lm(y~x), new)

cols <- rep("black",length(x))
cols[topredict] <- "red"
plot(x,y,pch=16,col=cols)
```

