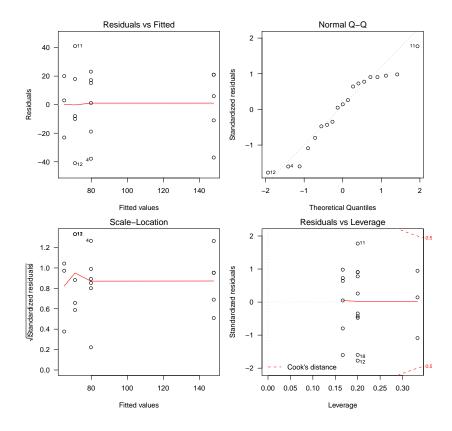
Solutions for 10.5 Exercises

1. We use a one-way ANOVA model:

```
y_{ij} = \mu_i + e_{ij}; e_{ij} \sim N(0, \sigma), i = A, B, C, D \text{ (or 1,2,3,4)}. j \text{ is different for each } i.
```

To obtain linear model diagnostics, we use:

```
> par(mfrow = c(2, 2), las = 1, mar = c(4, 4, 2, 1))
> plot(chicks.lm)
```



the las argument of the par function determines the style of axis labels; 1="always horizontal". The mar argument determines the margins, with the four numbers corresponding to (bottom, left, top, right). Type ?par for more details.

The diagnostic plots look acceptable. The variance in the four groups is similar, and the residuals appear to be consistent with a normal distribution.

```
> summary(chicks.lm)
Call:
lm(formula = weight_gain ~ feed, data = chicks)
Residuals:
  Min
           1Q Median
                         3Q
                               Max
-41.00 -14.92
                3.00 19.00 41.00
Coefficients:
            Estimate Std. Error t value Pr(>|t|)
                          14.94
                                  4.351 0.000571 ***
(Intercept)
               65.00
feedB
               14.83
                          18.30
                                  0.811 0.430247
feedC
                6.00
                          18.90
                                  0.317 0.755251
feedD
               83.00
                          18.90
                                  4.392 0.000525 ***
Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1
Residual standard error: 25.88 on 15 degrees of freedom
Multiple R-squared: 0.6758,
                                   Adjusted R-squared: 0.6109
F-statistic: 10.42 on 3 and 15 DF, p-value: 0.0005872
> anova(chicks.lm)
Analysis of Variance Table
Response: weight_gain
          Df Sum Sq Mean Sq F value
                                       Pr(>F)
           3 20937 6979.0 10.422 0.0005872 ***
feed
Residuals 15 10045
                      669.7
Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1
```

The small P-value indicates substantial evidence against the null hypothesis of equal means: $\mu_A = \mu_B = \mu_C = \mu_D$. The table for coefficients shows that β_3 is significantly different from β_0 but β_1 and β_2 are not. This means that feed B and feed C are not statistically significantly different from feed A, but feed D is. (Recall the parameterisation that R uses: feed A is the intercept, and the other three parameters are the difference between feed A and feeds B, C, and D respectively.) So to promote weight gain we would employ feed D.

```
condition 2 10.9387 5.4693 9.3466 0.008068 ** Residuals 8 4.6813 0.5852 --- Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1 1 P=0.008, so the null hypothesis of equal means is clearly rejected.
```

- (b) > qt(0.975, 8) [1] 2.306004 $8.02 6.63 \pm t_8^{0.975} \times \sqrt{0.5852 \left(\frac{1}{5} + \frac{1}{3}\right)} = 1.39 \pm 2.306 \times 0.559 = 1.39 \pm 1.29 = (0.10, 2.68).$

It appears that the distribution of the rainfall from seeded clouds has a larger mean and larger variability than the rainfall from control clouds. In both groups, the sample mean is much larger than the sample median, and the standard deviation is larger than the mean, suggesting substantial skewness in both populations.

The formal comparison of means (P = 0.051) is not significant at the 0.05 level, but it would be bad science to just ignore the difference. In a report, it would be appropriate to use words such as "some evidence of a difference".

(c) > t.test(rain ~ seeding, data = cloud, var = TRUE)
 Two Sample t-test

The equal P-values show that this analysis is equivalent to the ANOVA.

ANOVA assumes equal variances for the two groups, so the equivalent t-test needs this assumption incorporated. Note that the assumption is questionable here, and so a t-test with unequal variances would actually be more appropriate in this situation.

The standard deviations for the two groups are now very similar, so an analysis of the log transformed data would better meet the assumptions of ANOVA.

```
Two Sample t-test

data: lograin by seeding
t = -2.5443, df = 50, p-value = 0.01408
alternative hypothesis: true difference in means is not equal to 0
95 percent confidence interval:
-2.0467013 -0.2408495
sample estimates:
mean in group control mean in group seeded
3.990476
5.134252
```

The difference in means is now significant at the 0.05 level.

> t.test(lograin ~ seeding, data = cloud, var = TRUE)

(e) From this analysis, cloud seeding would be recommended. The analysis of the log transformed data (which better meets the assumptions) shows a significant increase in rainfall due to seeding.

```
4. (a) > metal.lm.1 <- lm(hardness ~ strip + tip, data = metal) > anova(metal.lm.1)
```

Analysis of Variance Table

Response: hardness

Df Sum Sq Mean Sq F value Pr(>F)

strip 4 0.843 0.210750 29.753 3.809e-06 ***

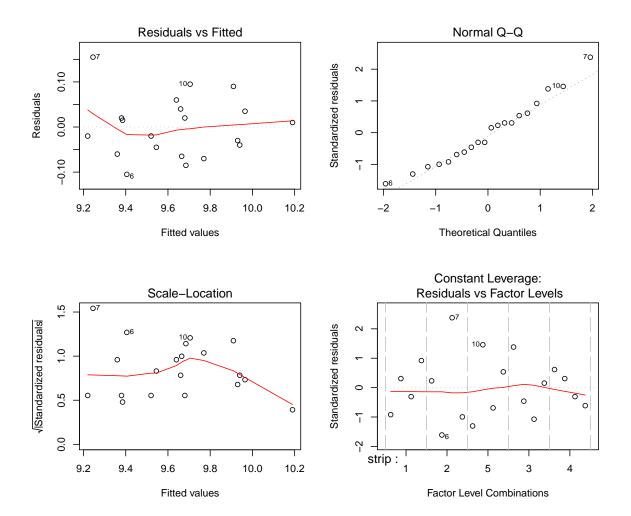
tip 3 0.460 0.153333 21.647 3.928e-05 ***

Residuals 12 0.085 0.007083

--
Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1

The very small P-value implies a highly significant difference between tips. Strip is a blocking factor, and so the small P-value is not of primary interest, except that it suggests that the blocking has been worthwhile.

- (b) > qt(0.975, 12)[1] 2.178813 $2.179 \times \sqrt{0.00708 \left(\frac{1}{5} + \frac{1}{5}\right)} = 0.12.$ 95% confidence interval = $9.88 - 9.46 \pm 0.12 = (0.30, 0.54).$



The plot of residuals vs fitted values is consistent with the assumption of constant variance, and the Q-Q plot is consistent with the assumption of normality of the errors.

(d) Pool degrees of freedom and sum of squares for strip and residuals:

Df:
$$4 + 12 = 16$$
.

Sum Sq: 0.843 + 0.085 = 0.928.

Residual Mean Sq = 0.928/16 = 0.058.

F = 0.1533/0.058 = 2.64.

This results in:

Df Sum Sq Mean Sq F value tip 3 0.460 0.153333 2.64 Residuals 16 0.928 0.058 > metal.lm.2 <- lm(hardness ~ tip, data = metal)</pre>

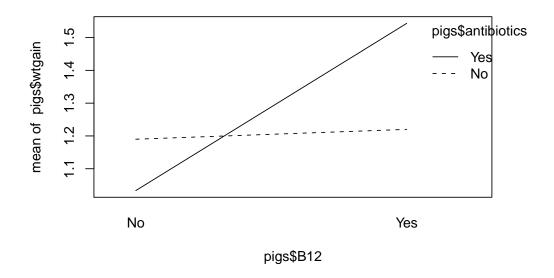
Analysis in R:

```
> anova(metal.lm.2)
       Analysis of Variance Table
       Response: hardness
                 Df Sum Sq Mean Sq F value Pr(>F)
                   3 0.460 0.15333 2.6437 0.08464 .
       Residuals 16 0.928 0.05800
       Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1
   (e) Blocking by strip in the experiment has been very worthwhile. It has substantially de-
       creased the residual MS by removing a large amount of the random variation, allowing
       much more precise inference on the factor of interest (tip). This is reflected in a much
       more significant effect of tip.
5. (a) > turnip <- read.csv("turnip.csv")
       > tapply(turnip$weight, turnip$variety, mean)
                                     С
       0.9958333 0.7866667 0.3716667 0.9660000 0.6100000 0.6503333
   (b) > turnip.lm.1 <- lm(weight ~ factor(row) + factor(col) + variety,
             data = turnip)
       > anova(turnip.lm.1)
       Analysis of Variance Table
       Response: weight
                    Df Sum Sq Mean Sq F value
                                                    Pr(>F)
       factor(row) 5 0.16445 0.03289 1.0002
                                                    0.4429
       factor(col) 5 0.08307 0.01661 0.5053
                                                    0.7688
       variety
                     5 1.67234 0.33447 10.1712 5.839e-05 ***
       Residuals
                    20 0.65768 0.03288
       Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1
       The P-value is very small, so the (null) hypothesis is decisively rejected.
   (c) Combine col and residual into new residual:
       Df = 5 + 20 = 25;
       Sum Sq = 0.0831 + 0.6577 = 0.7408;
       new residual Mean Sq = 0.7408/25 = 0.0296.
       It is not very different.
```

6. (a) Randomised block design, with litters as blocks.

library(stats)

> interaction.plot(pigs\$B12, pigs\$antibiotics, pigs\$wtgain)



There appears to be substantial interaction.

(c) > pigs.lm.1 <- lm(wtgain ~ litter + B12 * antibiotics, data = pigs)
> anova(pigs.lm.1)

Analysis of Variance Table

Response: wtgain

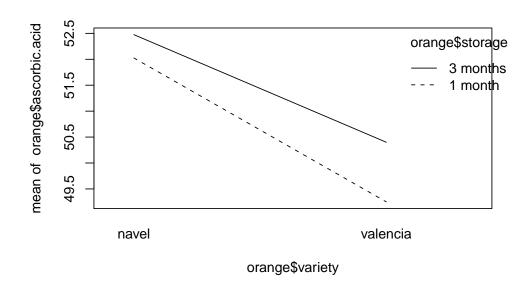
Df Sum Sq Mean Sq F value Pr(>F)
litter 2 0.008717 0.004358 1.2684 0.3471931
B12 1 0.218700 0.218700 63.6475 0.0002066 ***
antibiotics 1 0.020833 0.020833 6.0631 0.0489646 *
B12:antibiotics 1 0.172800 0.172800 50.2894 0.0003946 ***
Residuals 6 0.020617 0.003436

Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1

P < 0.001, confirming the significant interaction.

```
(d) Treatment Mean Sq = (0.2187 + 0.0208 + 0.1728)/(1 + 1 + 1) = 0.1374.
       F = 0.1374/0.003436 = 40.0, which is highly significant.
       > pigs$treatment <- factor(c(1, 2, 1, 2, 1, 2, 3, 4, 3, 4, 3, 4))
       > pigs.lm.2 <- lm(wtgain ~ litter + treatment, data = pigs)</pre>
       > anova(pigs.lm.2)
       Analysis of Variance Table
       Response: wtgain
                 Df Sum Sq Mean Sq F value
                  2 0.00872 0.004358 1.2684 0.3471931
       treatment 3 0.41233 0.137444 40.0000 0.0002319 ***
       Residuals 6 0.02062 0.003436
      Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1
7.
      SOURCE
                          DF
                                   SS
                                           MS
                                                      F
                                                                 Ρ
                                           70
      Thinning
                           2
                                  140
                                                   8.75
                                                             0.0030
      Provenance
                           4
                                 1200
                                          300
                                                  37.5
                                                            <0.0001
      Interaction
                           8
                                  430
                                           53.75
                                                   6.72
                                                             0.0008
      Error
                          15
                                  120
                                            8
      Total
                          29
                                 1890
  Calculation of P-values:
  > 1 - pf(8.75, 2, 15)
  [1] 0.003030814
  > 1 - pf(37.5, 4, 15)
  [1] 1.209653e-07
  > 1 - pf(6.72, 8, 15)
  [1] 0.000819122
8. (a) > orange <- read.csv("orange juice.csv")
       > tapply(orange$ascorbic.acid, orange[, 2:3], mean)
                 storage
                  1 month 3 months
       variety
                   52.025
                            52.475
         navel
         valencia 49.250
                            50.400
```

> interaction.plot(orange\$variety, orange\$storage, orange\$ascorbic.acid)



There appears to be not much interaction.

(b) > orange.lm <- lm(ascorbic.acid ~ variety * storage, data = orange) > anova(orange.lm)

Analysis of Variance Table

Response: ascorbic.acid

Df Sum Sq Mean Sq F value Pr(>F) 1 23.523 23.5225 11.9784 0.004709 ** variety storage 2.560 2.5600 1.3036 0.275823 0.4900 0.2495 0.626444 variety:storage 1 0.490 Residuals 12 23.565 1.9638

Signif. codes: 0 *** 0.001 ** 0.01 * 0.05 . 0.1

The interaction and the main effect of storage time are not significant. The main effect of variety is highly significant, with the mean for navels 2 to 3 mg/litre higher than for valencia.

- (c) $\hat{\sigma} = \sqrt{1.9638} = 1.40$.
- (d) Recommendation: use navel oranges if possible; it doesn't matter whether the juice is stored for 1 month or 3 months.