1 This is Problem 3 of Faraway (2006), Chapter 9

The ratdrink data consist of five weekly measurements of body weight for 27 rats. The first 10 rats are on a control treatment while seven rats have thyroxine added to their drinking water. Ten rats have thiouracil added to their water. Build a model for the rat weights that shows the effect of the treatment.

```
library(faraway)
data(ratdrink)
```

(a) Model the weights of the rate, incorporating the treatment effects and random effect. Use R to fit the model.

We write y_{ijk} to represent the kth rat in the jth treatment group on the ith week, where (i=1,2,3,4), (j=1,2,3), and (k=1-10 for control, k = 1-7 thyroxine, and k = 1-10 for thiouracil). μ represents the overall mean weight, α_i represents the fixed effect contribution of the jth treatment, and δ_{ij} is the interaction of weeks and treatment. The random effect u_{ik} incorporates the repeated measures of the same rat.

```
y_{ijk} = \mu + \alpha_i + \beta_j + \delta_{ij} + u_{jk} + \epsilon_{ijk}
   To fit the model in R we write:
   rat.lme <- lmer(wt ~ weeks+ treat+ weeks*treat+ (1|subject))
   The command summary (rat.lme) gives:
Linear mixed model fit by REML ['lmerMod']
Formula: wt ~ weeks + treat + weeks * treat + (1 | subject)
REML criterion at convergence: 948.4
Scaled residuals:
     Min
                1Q
                     Median
                                    3Q
                                             Max
-2.05506 -0.65511 -0.04848 0.57702 2.80847
Random effects:
 Groups
                        Variance Std.Dev.
          Name
 subject (Intercept) 71.21
                                  8.438
                        51.22
                                  7.157
 Residual
Number of obs: 135, groups: subject, 27
Fixed effects:
                        Estimate Std. Error t value
(Intercept)
                         52.8800
                                      3.1928
                                                16.56
weeks
                         26.4800
                                      0.7157
                                                37.00
treatthiouracil
                          4.7800
                                      4.5153
                                                 1.06
treatthyroxine
                         -0.7943
                                      4.9756
                                                -0.16
                                                -9.26
weeks:treatthiouracil
                        -9.3700
                                      1.0121
weeks:treatthyroxine
                          0.6629
                                      1.1153
                                                 0.59
```

Correlation of Fixed Effects:

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```
(Intr) weeks trtthr trtthy wks:trtthr weeks -0.448 treatthircl -0.707 0.317 treatthyrxn -0.642 0.288 0.454 wks:trtthrc 0.317 -0.707 -0.448 -0.203 wks:trtthyr 0.288 -0.642 -0.203 -0.448 0.454
```

(b) What is the implication of the random effect on the correlations between weights of the same rat? Is that implication reasonable? It would be nice to support your argument with data evidence.

The random effect captures the correlation between measuring the same rat on multiple weeks. Because that rat has a starting weight at week zero, subsequent measurements will be correlated with the previous weight.

We fit a completely fixed model and call summary(rat.lm) for comparison. As shown below, and as we would expect, the fixed coefficient estimates are the same without the random effect from the same rat. However, above, we were able to obtain a correlation matrix showing the correlation of the fixed effects. The correlation across weeks for the overall mean obtained was -.448, which is relatively low (highly correlated would be close to 1, or inversely correlated close to -1).

```
> rat.lm <- lm(wt ~weeks+treat+weeks*treat)</pre>
> summary(rat.lm)
Call:
lm(formula = wt ~ weeks + treat + weeks * treat)
Residuals:
   Min
             1Q Median
                             3Q
                                    Max
-23.514 -6.660
                  0.230
                          6.914
                                 28.343
Coefficients:
                      Estimate Std. Error t value Pr(>|t|)
(Intercept)
                       52.8800
                                   2.6547 19.919 < 2e-16 ***
                       26.4800
                                   1.0838 24.433 < 2e-16 ***
weeks
treatthiouracil
                        4.7800
                                   3.7544 1.273
                                                     0.205
treatthyroxine
                       -0.7943
                                   4.1371 -0.192
                                                     0.848
weeks:treatthiouracil -9.3700
                                   1.5327 -6.113 1.08e-08 ***
                        0.6629
                                   1.6890
                                            0.392
                                                     0.695
weeks:treatthyroxine
Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' 1
Residual standard error: 10.84 on 129 degrees of freedom
                                Adjusted R-squared:
Multiple R-squared: 0.9121,
F-statistic: 267.8 on 5 and 129 DF, p-value: < 2.2e-16
```

2 The article "Variability of Sliver Weights at Different Carding Stages and a Suggested Sampling Plan for Jute Processing"

by A. Lahiri (Journal of the Textile Institute, 1990) concerns the partitioning of variability in "sliver weight." (A sliver is a continuous strand of loose, untwisted wool, cotton, etc., produced along the way to making yarn.) For a particular mill, 3 (of many) machines were studied, using 5 (10 mm) pieces of sliver cut from each of 5 rolls produced on the

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machines. The weights of the (75) pieces of sliver were determined and a standard hierarchical (balanced data) ANOVA table was produced as below. (The units of weight were not given in the original article.)

Source	SS	df
Machines	1966	2
Rolls	644	12
Pieces	280	60
Total	2890	74

The model is

$$y_{ijk} = \mu + \alpha_i + u_{ij} + \epsilon_{ijk}$$

for the kth piece of the jth roll on the ith machine, where

$$\alpha_i \stackrel{iid}{\sim} N(0, \sigma_{\alpha}^2), u_{ij} \stackrel{iid}{\sim} N(0, \sigma_{u}^2), \text{and } \epsilon_i \stackrel{iid}{\sim} N(0, \sigma_{\epsilon}^2).$$

(a) Find estimates for σ^2_{α} , σ^2_{u} and σ^2_{ϵ} .

Source	SS	df	Term	MS		E(MS)
Machines	1966	2		MSA	983	$\sigma_{\epsilon}^2 + 5\sigma_u^2 + 25\sigma_{\alpha}^2$
Rolls	644	12		MSB]A	53.666667	$\sigma_{\epsilon}^2 + 5\sigma_u^2$
Pieces	280	60	error	MSE	4.6666667	σ_{ϵ}^2
Total	2890	74				

Solving for the expectations above gives the following estimates:

- $\widehat{\sigma_{\epsilon}^2} = 4.6667$
- $\widehat{\sigma_u^2} = 9.8$
- $\widehat{\sigma_{\alpha}^2} = 37.17333$

(b) Make 95% confidence intervals for each of the 3 standard deviations

 σ_{α} , σ_{u} , and σ_{ε} . Based on theses, where do you judge the largest part of the variation in measured weight to come from? You need to use the Cochran-Satterthwaite approximation for σ_{α} and σ_{u} .

For σ_{ϵ} we note that

$$\frac{SSE}{\sigma_{\epsilon}} \sim \chi_{60}^2$$
.

We write:

$$c(\sigma_{\epsilon}) = \left(\sigma_{\epsilon} : \chi_{60,.05}^2 < \frac{SSE}{\sigma_{\epsilon}^2} < \chi_{60,.95}^2\right)$$

$$c(\sigma_{\epsilon}) = \left(\sigma_{\epsilon} : \sqrt{\frac{SSE}{\chi_{60,.95}^2}} < \sigma_{\epsilon} < \sqrt{\frac{SSE}{\chi_{60,.05}^2}}\right)$$

$$c(\sigma_{\epsilon}) = (\sigma_{\epsilon} : 1.833423 < \sigma_{\epsilon} < 2.629961)$$

We use the Cochran-Satterwaithe approximation for $\sigma \alpha$ and σ_u . There is used to determine the degrees of freedom of a Chi-squared distribution which is approximately:

$$\frac{v(S^2)}{E(S^2)} \sim \chi_v^2$$

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This gives us a 1- α confidence interval for E(S²) determined by:

$$P\left(\frac{v \cdot S^2}{\chi_{v,upper}^2} < E(S^2) < \frac{v \cdot S^2}{\chi_{v,lower}^2}\right) = 1 - \alpha$$

$$\hat{v}_u = \frac{(\sigma_u^2)^2}{\frac{((MSB|A)/5)^2}{df=12} + \frac{(-MSE/5)^2}{df=60}} = 9.988675$$

$$\hat{v}_{\alpha} = \frac{(\sigma_{\alpha}^2)^2}{\frac{((-MSB|A)/25)^2}{df=12} + \frac{(MSA/25)^2}{df=2}} = 1.786695$$

Solving for our confidence intervals gives

$$c(\sigma_u) = (\sigma_u : 2.186967 < \sigma_u < 5.496051)$$

 $c(\sigma_\alpha) = (\sigma_\alpha : 3.097864 < \sigma_\alpha < 46.29789)$

Clearly, the largest contribution to the variability in the measured weight of the silver comes from the differences between machines. This is because the estimate of the standard deviation associated with $\sigma\alpha$ is the largest, and its confidence interval also ranges over the largest values.

(c) Suppose for the sake of illustration that the grand average

of all 75 weight measurements was in fact $\bar{y}_{...} = 1/75 \sum_{ijk} y_{ijk} = 35.0$.\$ Use this and information from the ANOVA table to make a 95% confidence interval for the model parameter μ .

The 95% confidence interval we are interested in is $\bar{y_{...}} \pm t_{.975,df=2} \sqrt{\frac{MSA}{3\cdot 5\cdot 5}} = (19.42305, 50.57695).$

3 Appendix: Tangled R Code

```
library (MASS); library (xtable); library (nlme)
  lvector \leftarrow function(x, dig = 2, dsply=rep("f", ncol(x)+1)) {
   x \leftarrow xtable(x, align=rep("", ncol(x)+1), display=dsply, digits=dig) # We repeat empty string 6 times
   print(x, floating=FALSE, tabular.environment="pmatrix",
     hline.after=NULL, include.rownames=FALSE, include.colnames=FALSE)
library (faraway)
data(ratdrink)
help(ratdrink)
library (lattice)
ratdrink$thecolor = "black"
ratdrink$thecolor[ratdrink$treat == "thyroxine"] = "red"
ratdrink$thecolor[ratdrink$treat == "thiouracil"] = "blue"
attach (ratdrink)
pdf("ratweights.pdf",width=7,height=5)
plot(weeks, wt, col = thecolor, main="Rat growth weights affected by additives")
legend("topleft", c("Control", "Thyroxine", "Thiouracil"), col=c("black", "red", "blue"), pch=1)
dev. off()
```

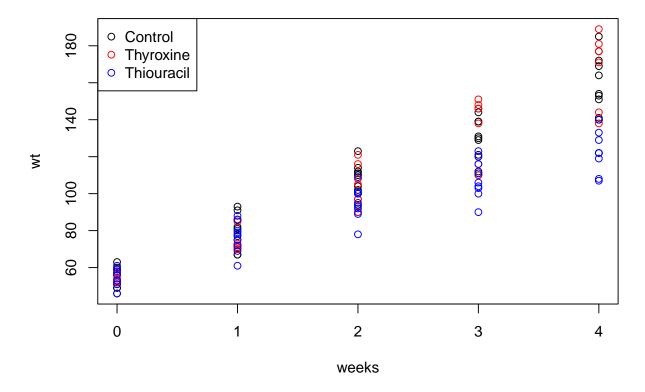
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```
#####Fit model#####
library (lme4)
rat.lme <- lmer(wt ~ weeks+ treat+ weeks*treat+ (1|subject))
summary(rat.lme)
var(ratdrink[treat=="control",]$wt)
myvars <- numeric(10);</pre>
for(i in 1:10){
    thisi <- toString(i);</pre>
    myvars[i] <- var(ratdrink[subject==thisi,]$wt)</pre>
}
mean(myvars)
typeof (weeks)
############################
sige <- 280/60
sigu \leftarrow ((644/12) - sige)/5
siga \leftarrow ((1966/2) - 5*sigu - sige)/25
sqrt (280/qchisq (.025,60))
sqrt (280/qchisq (.975,60))
MSB <- 644/12
MSA < -1966/2
MSE < -280/60
vhatu <- (sigu)^2/((((MSB/5)^2)/12)+((MSE/5)^2)/60)
vhatu
vhata \leftarrow (siga)^2/((((MSB/25)^2)/12) + (((MSA/25)^2)/2))
vhata
sqrt ((sigu*vhatu)/qchisq(.975, vhatu))
sqrt ((sigu*vhatu)/qchisq(.025,vhatu))
sqrt ((siga * vhata) / qchisq (.975, vhata))
sqrt ((siga*vhata)/qchisq(.025, vhata))
35 - qt(.975,2)*sqrt(MSA/(3*5*5))
35 + qt(.975,2)*sqrt(MSA/(3*5*5))
```

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4 Appendix: Initial evaluation of ratdrink dataset

Rat growth weights affected by additives



Plotting the ratdrink data suggested that rats that drank Thyroxine tended to have increased body weight after 5 weeks in comparison to rats drinking Thiouracil and Control. The rats that drank Thiouracil tended to have lowerbody weight than the Control and Thyroxine groups.