Homework 2

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Question 1

Part a.

We can use the aov() function to solve this.

The sums of squares associated with the blocks is 1288, the sums of squares associated with nitrogen is 6360, and the sums of squares associated with departures in additivity is 21846.

Part b.

According to our test, no, blocking has not been effective. The p-value for the blocking factor V2 is not significant, indicating that there is no significant difference between the observed yields of the different blocks. We could spend the degrees of freedom associated with blocking on our residuals instead.

That being said, there may be experimentally advantageous reasons to block, for example, if the blocks are in different geographic locations and the researchers need to incorporate this into their model. That might be a case where practical significance outweighs statistical significance.

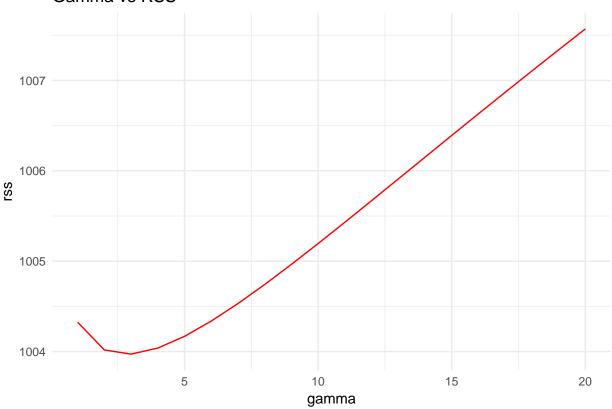
Part c.

```
nit_adjusted <- function(gamma) {
  model <- lm(V3 ~ as.factor(V2) + I(log(corn$V1 + gamma)), data = corn)
  output <- 45*(summary(model)$sigma)
  return(output)
}

gamma <- 1:20
rss <- vector(mode = "numeric", length = length(gamma))
for (g in gamma) {
  rss[g] <- nit_adjusted(g)
}

gamma_rss <- data.frame(gamma, rss)</pre>
```

Gamma vs RSS



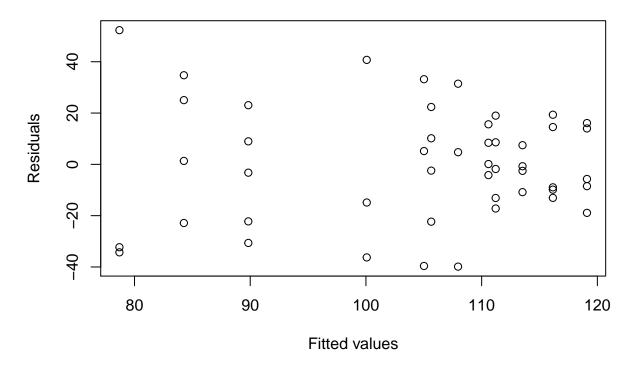
The value of γ with the smallest RSS appears to be 3. This will be our least squares estimate.

Computing a 95% Confidence interval for γ will have to involve either finding an MLE and using the asymptotic distribution of the MLE to create a Wald confidence interval **or** using some form of bootstrapping.

Part d.

```
gamma <- 3
model <- lm(V3 ~ V2 + I(log(corn$V1 + gamma)), data = corn)
plot(model$fitted.values, model$residuals,
    main = "Residual plot of logarithmic model",
    xlab = "Fitted values",
    ylab = "Residuals")</pre>
```

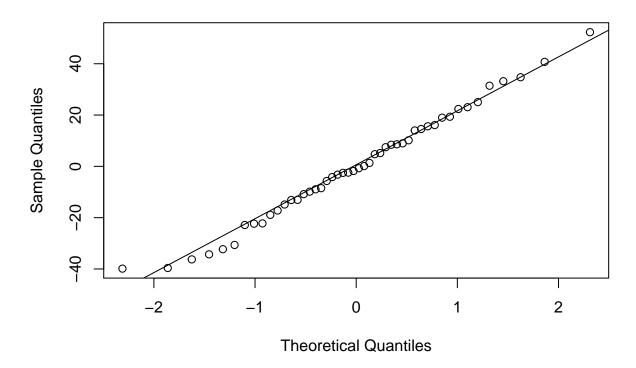
Residual plot of logarithmic model



There appears to be some heteroskedasticity here.

```
qqnorm(model$residuals)
qqline(model$residuals)
```

Normal Q-Q Plot



The errors however do appear to be roughly normal. I would suggest that this tells us the logarithmic function isn't a close fit to the actual data, but it isn't awful.

An alternative approach to checking diagnostic plots is to use a logarithmic lack of fit test where we compare the fit of the logarithmic model to an oversaturated model where each \hat{y} is just the group mean for that treatment combination (blocks x nitrogen).

```
##
## Call:
   lm(formula = V3 ~ factor(V1) * factor(V2), data = corn)
##
##
## Residuals:
##
       Min
                 1Q
                    Median
                                  3Q
                                         Max
   -39.200 -12.515
                     -0.887
                                      57.067
##
                             14.255
##
##
  Coefficients:
##
                              Estimate Std. Error t value Pr(>|t|)
## (Intercept)
                               73.9333
                                           14.1143
                                                      5.238 7.24e-06 ***
## factor(V1)50
                               22.6667
                                           19.9607
                                                               0.264
                                                      1.136
## factor(V1)100
                               30.6667
                                           19.9607
                                                      1.536
                                                                0.133
## factor(V1)150
                               32.8000
                                           19.9607
                                                      1.643
                                                               0.109
## factor(V2)1
                               19.8917
                                           18.6715
                                                      1.065
                                                                0.294
```

```
## factor(V2)2
                             11.0867
                                        17.8534
                                                  0.621
                                                           0.539
## factor(V1)50:factor(V2)1
                             -8.9167
                                        26.4055 -0.338
                                                           0.738
## factor(V1)100:factor(V2)1 -8.9167
                                        26.4055 -0.338
                                                           0.738
## factor(V1)150:factor(V2)1 -14.7500
                                        26.4055 -0.559
                                                           0.580
## factor(V1)50:factor(V2)2
                              2.6133
                                        25.2485
                                                  0.104
                                                           0.918
## factor(V1)100:factor(V2)2
                              0.8733
                                        25.2485
                                                  0.035
                                                           0.973
## factor(V1)150:factor(V2)2
                              0.6800
                                        25.2485
                                                  0.027
                                                           0.979
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 24.45 on 36 degrees of freedom
## Multiple R-squared: 0.2705, Adjusted R-squared: 0.04765
## F-statistic: 1.214 on 11 and 36 DF, p-value: 0.3134
```

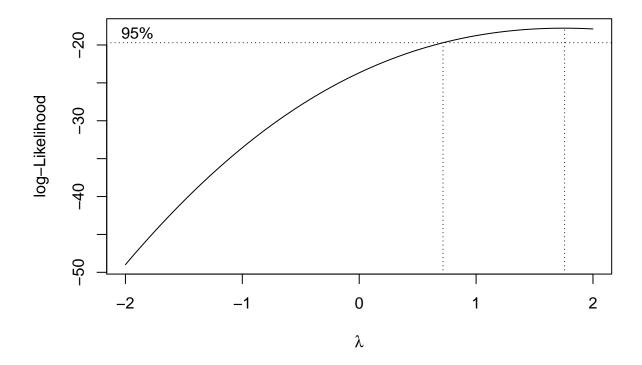
```
anova(saturated_model, model)
```

```
## Analysis of Variance Table
##
## Model 1: V3 ~ factor(V1) * factor(V2)
## Model 2: V3 ~ V2 + I(log(corn$V1 + gamma))
## Res.Df RSS Df Sum of Sq F Pr(>F)
## 1 36 21515
## 2 45 22238 -9 -722.96 0.1344 0.9984
```

This high p-value indicates that we have no strong evidence of lack of fit!

Part e.

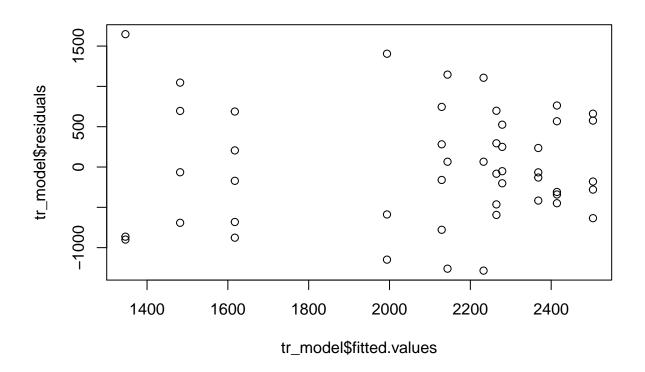
Using the boxcox function in R, we can identify the response transformation that gives us the greateset log-likelihood.

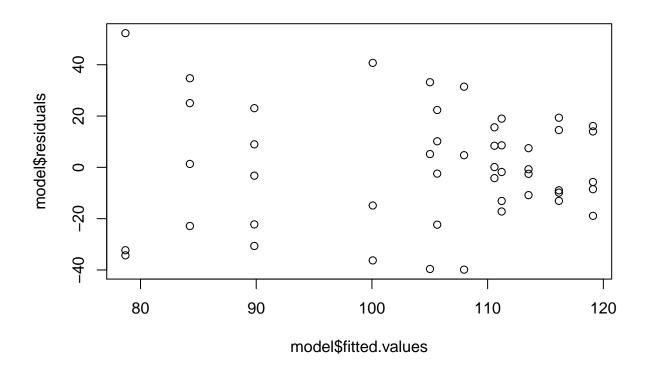


```
lambda <- with(corncox, x[y == max(y)])
lambda</pre>
```

[1] 1.757576

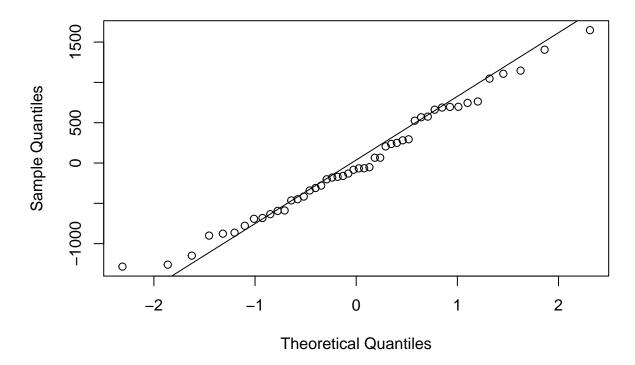
Applying this transformation to our response and refitting the model gives us the following probabilities.





```
qqnorm(tr_model$residuals)
qqline(tr_model$residuals)
```

Normal Q-Q Plot



The transformation actually made everything worse!

Part f.

Blocking was not effective, so in later experiments it might be better to not include it as an effect and have additional error degrees of freedom. The log transformation performed the best of the models we tried, noting that even then there is evidence of nonconstant variance. However, unless we are trying to make predictions, this heteroskedasticity should be fine.

Question 2

```
##
      temp weeks con
## 1
                   45
         0
                2
## 2
         0
                   47
         0
## 3
                6
                   46
## 4
         0
               8
                   46
## 5
        10
               2
                   45
## 6
        10
               4
                   43
## 7
        10
                6
                   41
## 8
        10
               8
                   37
               2
## 9
        20
                   34
## 10
        20
                4
                   28
        20
                6
                   21
## 11
## 12
        20
                8
                   16
```

The regression model here is

$$y_{t,T} = e^{-\alpha - \beta_T t + \epsilon_{t,T}}, \epsilon_{t,T} \sim$$

where $y_{t,T}$ is the concentration of ascorbic acid at time t with temperature T, β_T is the interaction effect of temperature T and storage time, and e^{α} is the intial concentration.

The score equations for this model are provided on a separate sheet of paper.

```
log_model <- glm(con ~ factor(temp):weeks,</pre>
                data = beans,
                family = gaussian(link="log"))
summary(log_model)
##
## Call:
## glm(formula = con ~ factor(temp):weeks, family = gaussian(link = "log"),
##
      data = beans)
##
## Deviance Residuals:
       Min
                       Median
                                     30
                                              Max
## -1.49820 -0.38546
                      0.08059
                                0.80866
                                          0.86028
##
## Coefficients:
##
                        Estimate Std. Error t value Pr(>|t|)
## (Intercept)
                       ## factor(temp)0:weeks -0.0007716 0.0037540 -0.206 0.842286
## factor(temp)10:weeks -0.0236121 0.0040351 -5.852 0.000382 ***
## factor(temp)20:weeks -0.1329785  0.0061597 -21.588 2.23e-08 ***
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
## (Dispersion parameter for gaussian family taken to be 1.124727)
##
##
      Null deviance: 1226.9167 on 11 degrees of freedom
## Residual deviance:
                       8.9978 on 8 degrees of freedom
## AIC: 40.599
##
```

Number of Fisher Scoring iterations: 3

Part b.

We will use these two facts to construct a confidence interval:

$$\frac{1}{2}e^{\alpha} = e^{\alpha - \beta_i h}$$
$$\frac{1}{2} = e^{-\beta_T t}$$
$$t = \frac{\ln(2)}{\beta_T}$$

and that $\frac{be\hat{t}a_i - \beta_i}{SE(\beta_i)} \sim t_8$.

We can construct a $(1 - \alpha)100\%$ confidence interval for each β_T , $T \in (0, 10, 20)$:

$$\left(\hat{\beta_T} - 2.306SE(\hat{\beta_T}), \hat{\beta_T} + 2.306SE(\hat{\beta_T})\right)$$

Noting that for each T, t is just a function of β_T , we can get $\hat{t} = \frac{\ln(2)}{\hat{\beta_T}}$ and a $(1 - \alpha)100\%$ CI as:

$$\left(\frac{ln(2)}{\hat{\beta_T} + 2.306SE(\hat{\beta_T})}, \frac{ln(2)}{\hat{\beta_T} - 2.306SE(\hat{\beta_T})}\right)$$

Plugging in the values from the model summary, we get that $\hat{t_0}=898.324,\,\hat{t_{10}}=29.356,\,\hat{t_{20}}=5.212$