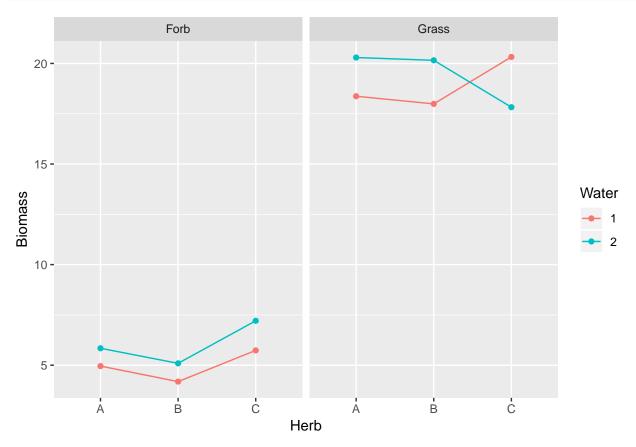
HW8 KEY

40 points total, 2 points per problem part unless otherwise noted.

```
library(dplyr)
library(ggplot2)
library(car)
library(emmeans)
InData <- read.csv("C:/hess/STAT512/HW_2019/HW8/Biomass.csv")
#str(InData)
InData$Water <- as.factor(InData$Water)</pre>
```

1. Summary Graph (4pts)



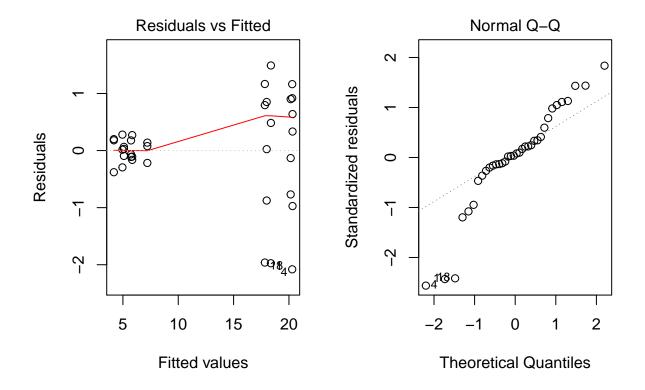
2. Three-way ANOVA table

```
options(contrasts=c("contr.sum","contr.poly"))
Model1 <- lm(Biomass ~ Type*Herb*Water, data = InData)</pre>
Anova(Model1, type = 3)
## Anova Table (Type III tests)
##
## Response: Biomass
##
                                          Pr(>F)
                   Sum Sq Df
                               F value
## (Intercept)
                  5473.8 1 5539.1323 < 2.2e-16 ***
## Type
                   1678.7 1 1698.7721 < 2.2e-16 ***
## Herb
                     5.1 2
                                2.5644 0.097872 .
## Water
                     5.9 1
                                5.9711 0.022271 *
## Type:Herb
                     5.4 2
                                2.7185 0.086259 .
## Type:Water
                     0.7 1
                                0.7092 0.408021
## Herb:Water
                     7.9 2
                                3.9973 0.031742 *
## Type:Herb:Water
                    13.1 2
                                6.6440 0.005055 **
## Residuals
                     23.7 24
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

3. Three-way Diagnostic plots

Note: Diagnostic plots not required for full credit, but shown here for completeness. Based on the plot of Resids vs Fitted, there is strong evidence of unequal variance. (Based on the QQplot, there is also some evidence against normality. But this is actually driven by the unequal variance.)

```
par(mfrow=c(1,2))
plot(Model1, which = c(1:2))
```



4. Three-way emmeans

```
emout1 <- emmeans(Model1, pairwise ~ Water | Herb*Type)
emout1$contrasts</pre>
```

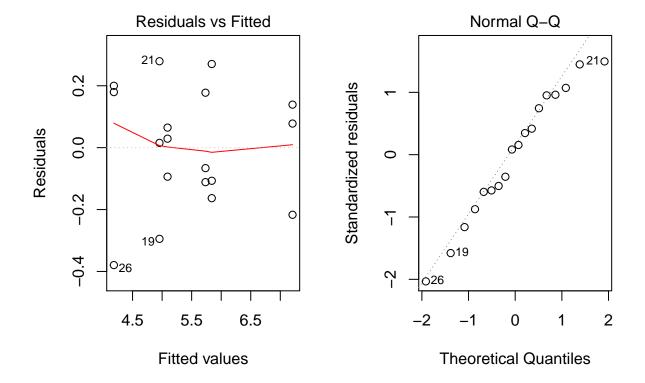
```
## Herb = A, Type = Forb:
   contrast estimate
                         SE df t.ratio p.value
              -0.885 0.812 24 -1.090 0.2865
##
## Herb = B, Type = Forb:
   contrast estimate
                         SE df t.ratio p.value
##
   1 - 2
              -0.908 0.812 24 -1.119 0.2741
## Herb = C, Type = Forb:
   contrast estimate
                         SE df t.ratio p.value
              -1.473 0.812 24 -1.815 0.0821
##
##
## Herb = A, Type = Grass:
   contrast estimate
                         SE df t.ratio p.value
              -1.924 0.812 24 -2.371 0.0261
##
##
## Herb = B, Type = Grass:
   contrast estimate
                         SE df t.ratio p.value
              -2.167 0.812 24 -2.669 0.0134
##
##
## Herb = C, Type = Grass:
## contrast estimate
                         SE df t.ratio p.value
```

```
## 1 - 2
               2.499 0.812 24 3.079 0.0051
```

5. FORB Two-way ANOVA table

```
Model2 <- lm(Biomass ~ Herb*Water, data = InData[InData$Type == "Forb",])</pre>
Anova(Model2, type =3)
## Anova Table (Type III tests)
##
## Response: Biomass
##
               Sum Sq Df
                             F value
                                        Pr(>F)
## (Intercept) 544.92 1 10432.1562 < 2.2e-16 ***
## Herb
                10.17
                       2
                             97.3895 3.820e-08 ***
                            102.1229 3.197e-07 ***
## Water
                 5.33
                       1
                       2
## Herb:Water
                 0.33
                              3.1846
                                       0.07772 .
## Residuals
                 0.63 12
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
  6. FORB diagnostic plots look much better! Some evidence of unequal variance, but not severe.
```

```
par(mfrow = c(1,2))
plot(Model2, which = c(1:2))
```



7. FORB emmeans #1 (interaction comparisons)

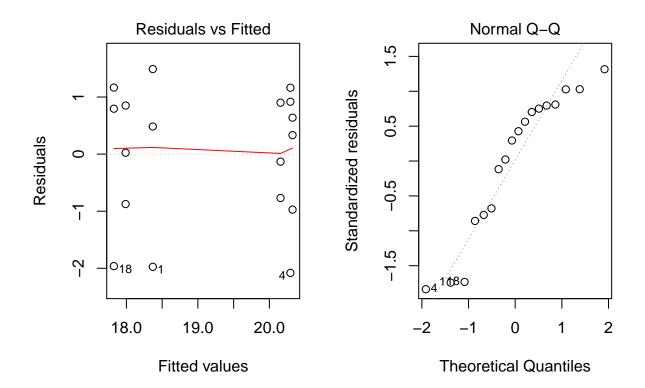
```
emout2 <- emmeans(Model2, pairwise ~ pairwise ~ Water | Herb)</pre>
emout2$contrasts
```

```
## Herb = A:
```

```
## contrast estimate
                         SE df t.ratio p.value
           -0.885 0.187 12 -4.742 0.0005
## 1 - 2
##
## Herb = B:
## contrast estimate
                         SE df t.ratio p.value
            -0.908 0.187 12 -4.868 0.0004
##
##
## Herb = C:
## contrast estimate
                         SE df t.ratio p.value
## 1 - 2
              -1.473 0.187 12 -7.894 <.0001
  8. FORB LSD #1 (interaction comparisons)
qt(0.975, df = 12)*sqrt(2*(0.63/12)/3)
## [1] 0.4076186
qt(0.975, df = 12)*0.187
## [1] 0.407438
  9. FORB emmeans #2 (main effect comparison)
emout3 <- emmeans(Model2, pairwise ~ Water)</pre>
## NOTE: Results may be misleading due to involvement in interactions
emout3$contrasts
##
   contrast estimate
                         SE df t.ratio p.value
               -1.09 0.108 12 -10.106 <.0001
## Results are averaged over the levels of: Herb
 10. FORB LSD #2 (main effect comparison)
qt(0.975, df = 12)*sqrt(2*(0.63/12)/9)
## [1] 0.2353387
qt(0.975, df = 12)*0.108
## [1] 0.2353118
 11. The power is higher for the main effect comparison (#10) because the LSD (ME) is smaller.
 12. GRASS Two-way ANOVA table
Model3 <- lm(Biomass ~ Herb*Water, data = InData[InData$Type == "Grass",])
Anova(Model3, type =3)
## Anova Table (Type III tests)
##
## Response: Biomass
               Sum Sq Df
                           F value
                                      Pr(>F)
## (Intercept) 6607.6 1 3433.9972 4.029e-16 ***
## Herb
                  0.3 2
                            0.0694
                                      0.9333
                  1.3 1
                                      0.4329
## Water
                            0.6586
## Herb:Water
                20.7 2
                            5.3786
                                      0.0215 *
## Residuals
                 23.1 12
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

13. GRASS diagnostic plots look much better! Some evidence of skew (non-normality), but not severe.

```
par(mfrow = c(1,2))
plot(Model3, which = c(1:2))
```



14. GRASS emmeans (interaction comparisons)

```
emout4 <- emmeans(Model3, pairwise ~ Water|Herb)
emout4$contrasts
## Herb = A:</pre>
```

```
##
    contrast estimate
                        SE df t.ratio p.value
##
                -1.92 1.13 12 -1.699 0.1150
##
## Herb = B:
   contrast estimate
                        SE df t.ratio p.value
##
                -2.17 1.13 12 -1.913 0.0799
##
## Herb = C:
   contrast estimate
                        SE df t.ratio p.value
##
   1 - 2
                 2.50 1.13 12 2.207 0.0476
```

- 15. No, because there is evidence of an interaction between Water and Herb. In particular, the estimated difference in mean response changes sign when comparing across Herb. When there is evidence of an interaction, it does not make sense to look at main effects.
- 16. (4pts) The estimated differences are the SAME. The SEs are DIFFERENT (Combined SE = 0.812, Forb SE = 0.187, Grass SE = 1.13). The df is DIFFERENT (combined df = 24, split df = 12).

- 17. (1) The combined analysis showed strong evidence of unequal variance. The model assumptions are better satisfied using the split analyses. (2) The combined analysis showed evidence of a three-way interaction.
- 18. (1) When we split the analysis we reduce the df (hence reducing power). (2) When we split the analysis we cannot get direct comparisons of Forb versus Grass.