Tukey HSD in R Tutorial

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1. Introduction

This document illustrates how to conduct ANOVA and a Tukey HSD test in R.

2. ANOVA

GOAL: Perform ANOVA on *chickwts*.

Important functions: setnames(), aggregate(), merge(), aov(), summary(), plotmeans(), TukeyHSD()

First, we need to load the *gplots* library into R. Loading all the libraries that we may use at the beginning is good practice, as sometimes we need libraries earlier than we initially expect.

```
library(gplots) # if not already installed, type install.packages('gplots')
```

Next, we need to load the data. The easiest way is with data(), which imports datasets pre-loaded into R.

```
data(chickwts)
```

2a. Summary Statistics

We nead the number of occurences; mean of the weights; and standard deviations for each supplement ("feed"). We know that NROW() counts the number of rows; mean() calculates the average; and sd() estimates the standard deviation. To execute these functions by groups (in this case, by feed), we can use aggregate().

```
##
          feed weight.obs weight.mean weight.sd
## 1
        casein
                  12.00000
                             323.58333
                                        64.43384
                 10.00000
## 2 horsebean
                             160.20000
                                        38.62584
## 3
       linseed
                 12.00000
                             218.75000
                                        52.23570
## 4
      meatmeal
                 11.00000
                             276.90909
                                         64.90062
## 5
                 14.00000
                             246.42857
       soybean
                                         54.12907
## 6 sunflower
                 12.00000
                             328.91667
                                         48.83638
```

The function aggregate() computes a specific function on a dataset—in this particular example, we use NROW, mean, and sd to calculate the number of observations, mena, and standard deviation, respectively, of the chick weights by type of feed. The formula $weight \sim feed$ is interpreted as "compute a function on weight BY feed."

2b. ANOVA

To estimate ANOVA, we need aov().

```
chickwts_anova <- aov(data = chickwts, formula = weight ~ feed)
summary(chickwts_anova)</pre>
```

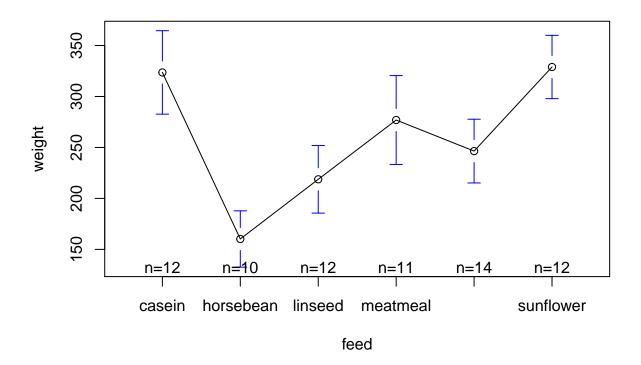
```
## Df Sum Sq Mean Sq F value Pr(>F)
## feed 5 231129 46226 15.37 5.94e-10 ***
## Residuals 65 195556 3009
## ---
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' ' 1
```

What does the p-value signify? Can we reject the null hypothesis that the feed type is irrelevant in determining a chick's weight? The p-value signifies the probability of comitting a Type I Error (i.e., a false positive). Because p < 0.05 (a conventional rule), we can reject the null hypothesis: the type of feed matters in influencing weight.

2c. Confidence Intervals

We need the *gplots* library, which we already loaded into R earlier, and *plotmeans()*.

```
with(chickwts, plotmeans(weight ~ feed))
```



```
# with() allows us to specify a dataset we want to work with.
# In this way, we don't have to type "chickwts$" repeatedly.
```

Which groups overlap? Based on the plot, horsebean overlaps with linseed; meatmeal overlaps with all groups except horsebean; linseed overlaps with soybean and horsebean; and sunflower overlaps with casein and meatmeal.

2d. Tukey HSD

We need TukeyHSD(), which we apply over our ANOVA results.

TukeyHSD(chickwts_anova)

```
##
     Tukey multiple comparisons of means
##
       95% family-wise confidence level
##
## Fit: aov(formula = weight ~ feed, data = chickwts)
##
## $feed
##
                              diff
                                           lwr
                                                      upr
## horsebean-casein
                       -163.383333 -232.346876 -94.41979 0.0000000
## linseed-casein
                       -104.833333 -170.587491 -39.07918 0.0002100
## meatmeal-casein
                        -46.674242 -113.906207 20.55772 0.3324584
                        -77.154762 -140.517054 -13.79247 0.0083653
## soybean-casein
## sunflower-casein
                          5.333333
                                    -60.420825 71.08749 0.9998902
## linseed-horsebean
                                    -10.413543 127.51354 0.1413329
                         58.550000
## meatmeal-horsebean
                        116.709091
                                     46.335105 187.08308 0.0001062
                         86.228571
## soybean-horsebean
                                     19.541684 152.91546 0.0042167
## sunflower-horsebean
                                     99.753124 237.68021 0.0000000
                        168.716667
## meatmeal-linseed
                         58.159091
                                     -9.072873 125.39106 0.1276965
## soybean-linseed
                         27.678571
                                     -35.683721 91.04086 0.7932853
## sunflower-linseed
                        110.166667
                                     44.412509 175.92082 0.0000884
## soybean-meatmeal
                        -30.480519
                                    -95.375109 34.41407 0.7391356
## sunflower-meatmeal
                         52.007576
                                    -15.224388 119.23954 0.2206962
## sunflower-soybean
                         82.488095
                                     19.125803 145.85039 0.0038845
```

Recall that each of the contrasts is a t-test. Which groups had statistically significant differences? How do you know? What can we say from these results?

3. Two-way ANOVA

GOAL: Perform Two-way ANOVA on CO2.

METHOD:

- 1. *uptake* is the response variable.
- 2. **Treatment * Type** are the factors.
- 3. aov(), TukeyHSD(), and interaction.plot() are to be used (as stated in the assignment).

Important functions: table(), aov(), summary(), plotmeans(), interaction.plot()

Load the following dataset:

```
data(CO2)
```

3a. Treatment vs. Type

Let's generate a table comparing Treatment against Type (column against row). We thus need with() to sepecify our working dataset and table() for the actual table.

```
with(CO2, table(Type, Treatment)) # we need Type as the row and Treatment as the column.
```

```
## Treatment
## Type nonchilled chilled
## Quebec 21 21
## Mississippi 21 21
```

Is the design balanced or unbalanced? Why?

3b. Main and Interaction Effects

We need a Two-way ANOVA with factors.

```
CO2_anova <- aov(uptake ~ Treatment*Type, data = CO2)
summary(CO2_anova)
```

```
##
                  Df Sum Sq Mean Sq F value
                                              Pr(>F)
## Treatment
                   1
                        988
                                988
                                    15.416 0.000182 ***
## Type
                   1
                       3366
                               3366
                                    52.509 2.38e-10 ***
## Treatment:Type
                  1
                        226
                                226
                                      3.522 0.064213 .
## Residuals
                  80
                                 64
                       5128
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
```

What are the statistically significant results? Based on what significance level?

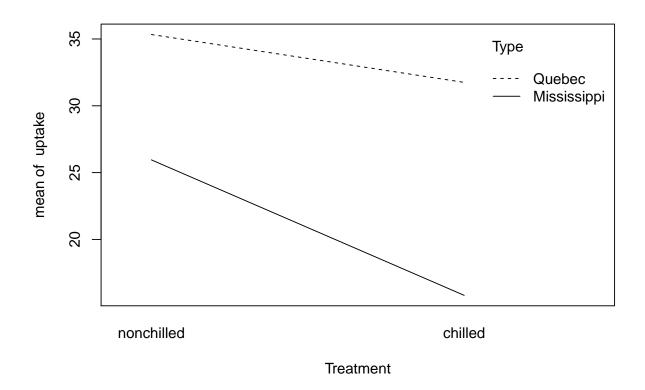
3c. Interaction Effects, continued.

We need to know the statistically significant differences between each Treatment-Type combinations. We've done something similar in Question 1d; but now we also need to have a plot of the interactions.

TukeyHSD(CO2_anova)

```
##
     Tukey multiple comparisons of means
       95% family-wise confidence level
##
##
##
  Fit: aov(formula = uptake ~ Treatment * Type, data = CO2)
##
##
  $Treatment
##
                           diff
                                       lwr
                                                 upr
                                                         p adj
  chilled-nonchilled -6.859524 -10.33624 -3.382808 0.0001817
##
## $Type
##
                           diff
                                       lwr
## Mississippi-Quebec -12.65952 -16.13624 -9.182808
##
## $`Treatment:Type`
##
                                                     diff
                                                                lwr
## chilled:Quebec-nonchilled:Quebec
                                                -3.580952 -10.06369
                                                                       2.9017869
## nonchilled:Mississippi-nonchilled:Quebec
                                                -9.380952 -15.86369
                                                                     -2.8982131
## chilled:Mississippi-nonchilled:Quebec
                                               -19.519048 -26.00179 -13.0363083
## nonchilled:Mississippi-chilled:Quebec
                                                -5.800000 -12.28274
                                                                       0.6827393
## chilled:Mississippi-chilled:Quebec
                                               -15.938095 -22.42083
                                                                     -9.4553560
## chilled:Mississippi-nonchilled:Mississippi -10.138095 -16.62083
                                                                     -3.6553560
##
                                                   p adj
## chilled:Quebec-nonchilled:Quebec
                                               0.4727714
## nonchilled:Mississippi-nonchilled:Quebec
                                               0.0015893
## chilled:Mississippi-nonchilled:Quebec
                                               0.000000
## nonchilled:Mississippi-chilled:Quebec
                                               0.0959830
## chilled:Mississippi-chilled:Quebec
                                               0.000000
## chilled:Mississippi-nonchilled:Mississippi 0.0005553
```

with(CO2, interaction.plot(Treatment, Type, uptake))



End of Document