Tutorial 9 ST2137-2420

Material

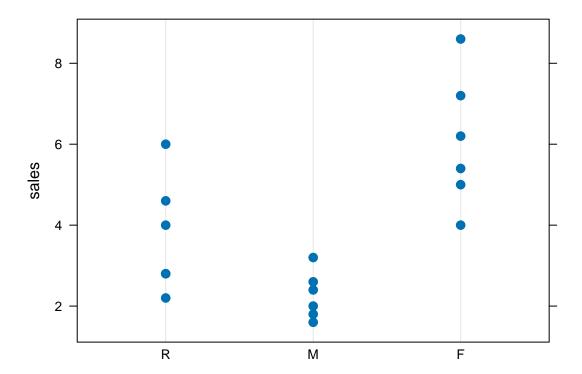
This tutorial covers the topics and concepts from chapter 8. Think of this topic as a generalisation of the approaches in chapter 7. In chapter 9, we will proceed to linear regression. Take note that 2-sample models, ANOVA models and linear regression models are all linear models.

Question 1

The retailing manager of a supermarket chain wants to determine whether product location has any effect on the sale of pet toys. Three different aisle locations are considered: front, middle, and rear. A random sample of 18 stores is selected with 6 stores randomly assigned to each aisle location. The size of the display area and price of the products are constant for all stores. At the end of a one-month trial period, the sales volumes (in thousands of dollars) of the product in each store were recorded in the file locate.txt.

- 1. Assuming that the observations are Normally distributed, use SAS to assess if there is any evidence of a significant difference in average sales among the various aisle locations, at 5% significance level.
- 2. Boxplots are typically used to assess the distribution within each group. However when we have so few observations, it is sometimes useful to plot every single point, by group. Use dotplot from the lattice package in R to create the following plot:

Sales by Aisle Location



4. In R and Python, set the reference level to be "rear". Compute the confidence interval for the differences between (i) front and rear, and (ii) middle and rear. Use a Bonferroni correction to adjust for the multiple tests so that overall, the error rate is 5%.

Solution

The SAS output indicates strong evidence against the null hypothesis. We would reject the null hypothesis and conclude the means are different.

		Dependent Varia	ble: sales		
Source	DF	Sum of Squares	Mean Square	F Value	Pr > F
Model	2	44.07111111	22.03555556	13.03	0.0005
Error	15	25.36000000	1.69066667		
Corrected Total	17	69.43111111			

In order to compare against the reference "rear", we have to set the levels properly in R. In Python, I recoded the levels so that "rear" is alphabetically first.

R code

```
anova_mod <- lm(sales ~ location, data=locate_df)
confint(anova_mod, level = 1-0.05/2)</pre>
```

1.25 % 98.75 % (Intercept) 2.4116368 5.0550298 locationM -3.3358278 0.4024944 locationF 0.4641722 4.2024944

Python code

```
import pandas as pd
import numpy as np
from scipy import stats
import statsmodels.api as sm
from statsmodels.formula.api import ols
import statsmodels.stats.multicomp as mc

locate_df = pd.read_table("data/locate.txt", delimiter="\\s+")
locate_df.replace({'F': '3-F', 'M': '2-M', 'R': '1-R'}, inplace=True)

locate_lm = ols('sales ~ C(location, Treatment)', data=locate_df).fit()
#anova_tab = sm.stats.anova_lm(locate_lm, type=3,)
#print(anova_tab)

print(locate_lm.summary(alpha=0.05/2))
```

C:\Users\stavg\penvs\p312\Lib\site-packages\scipy\stats_axis_nan_policy.py:418: UserWarning: `kurtosis
return hypotest_fun_in(*args, **kwds)

OLS Regression Results

______ sales R-squared: Dep. Variable: 0.635 Model: OLS Adj. R-squared: 0.586 Method: Least Squares F-statistic: 13.03 Fri, 11 Apr 2025 Prob (F-statistic): 0.000524 Date: 13:29:45 Log-Likelihood: Time: -28.626 AIC: No. Observations: 18 63.25 Df Residuals: 15 BIC: 65.92 2

Df Model: 2
Covariance Type: nonrobust

=======================================			========	=======		=======
	coef	std err	t	P> t	[0.0125	0.9875]
Intercept	3.7333	0.531	7.033	0.000	2.412	5.055
C(location, Treatment)[T.2-M]	-1.4667	0.751	-1.954	0.070	-3.336	0.402
C(location, Treatment)[T.3-F]	2.3333	0.751	3.108	0.007	0.464	4.202
=======================================					===	
Omnibus:	1.188	Durbin-Watson	:	1.3	186	
<pre>Prob(Omnibus):</pre>	0.552	Jarque-Bera (JB):	0.7	789	
Skew:	0.495	Prob(JB):		0.6	674	
Kurtosis:	2.732	Cond. No.		3	.73	
=======================================		=========	========	========	===	

Notes:

[1] Standard Errors assume that the covariance matrix of the errors is correctly specified.

Instead of using confint(), we can also use the formula in Section 8.4 of the textbook. We will arrive at the same answer:

```
c1 <- c(-1, 1, 0)
n_vals <- c(6,6,6)
est_coef <- coef(anova_mod)
L <- sum(c1*c(0, est_coef[2:3]))

summary_out <- anova(anova_mod)
MSW <- summary_out$^Mean Sq^[2]
df <- summary_out$Df[2]
se1 <- sqrt(MSW * sum( c1^2 / n_vals ) )</pre>
```

```
q1 <- qt(0.0125, df, 0, lower.tail = FALSE)
lower_ci <- L - q1*se1
upper_ci <- L + q1*se1
cat("The 95% CI for the diff. between the two groups is (",
    format(lower_ci, digits = 6), ", ", format(upper_ci, digits = 6), ").", sep="")</pre>
```

The 95% CI for the diff. between the two groups is (-3.33583, 0.402494).

Question 2

In earlier topics we noticed that, in the student performance dataset from $\mathtt{student-mat.csv}$, G3 scores seem to be different for different Medu groups. Remove the group corresponding to Medu=0 since there are so few observations. Use the following rule to remove outliers from $each\ group$: X_i is declared an outlier if

$$\frac{|X_i - \text{median}(X)|}{MAD(X)/0.6745} > 2.24$$

Perform the appropriate statistical test(s) to assess the following questions of interest:

- 5. Is there a significant difference between the 4 groups, at 5% significance level?
- 6. Estimate the confidence interval for a contrast comparing higher education to non-higher education (i.e. Medu = 4 vs. Medu = 1|2|3).
- 7. Use Tukey's HSD method to identify which pairs of groups are significantly different from one another at 5% family-wise error level.
- 8. Repeat the Tukey procedure with all outliers reinstated. How do the results differ?

Solution

R code

```
stud_perf <- read.table("data/student/student-mat.csv", sep=";",</pre>
                          header=TRUE)
stud_perf2 <- stud_perf[stud_perf$Medu != 0, ]</pre>
stud_perf2$Medu <- as.factor(stud_perf2$Medu)</pre>
remove_outliers <- function(d1) {</pre>
  ids \leftarrow which(abs(d1 - median(d1))/mad(d1) > 2.24)
  d1[-ids]
tmp_list <- vector(mode="list", 4)</pre>
for(ii in 1:4){
  tmp_list[[ii]] <- remove_outliers(stud_perf$G3[stud_perf$Medu == ii])</pre>
}
lens <- vapply(tmp_list, length, 2L)</pre>
stud_perf3 <- data.frame(G3 = unlist(tmp_list), Medu = as.factor(rep(1:4, times=lens)))</pre>
lm_outliers_rm <- lm(G3 ~ Medu, data=stud_perf3)</pre>
anova(lm outliers rm)
Analysis of Variance Table
Response: G3
            Df Sum Sq Mean Sq F value
             3 181.8 60.593 6.2057 0.0004073 ***
Medu
```

```
Residuals 348 3397.9 9.764
---
Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
The p-value is 0.0004073.
```

Python code

```
stud_perf = pd.read_csv("../data/student/student-mat.csv", delimiter=";")
stud_perf2 = stud_perf[stud_perf.Medu != 0]
def identify_outliers(d1):
    id_vec = np.where(.6745 * np.absolute(d1 -
                                          np.quantile(d1, 0.5))/
                                          stats.median abs deviation(d1) > 2.24)
    return id_vec
out_df_list = []
for i,df in stud_perf2.groupby('Medu'):
    to_rm = identify_outliers(df.G3)
    out df = df.drop(df.index[to rm])
    out_df_list.append(out_df)
stud_perf3 = pd.concat(out_df_list)
lm_outliers_rm = ols('G3 ~ C(Medu, Treatment)', data=stud_perf3).fit()
anova_tab = sm.stats.anova_lm(lm_outliers_rm, type=3,)
print(anova_tab)
```

```
df sum_sq mean_sq F PR(>F)
C(Medu, Treatment) 3.0 181.779541 60.593180 6.205657 0.000407
Residual 348.0 3397.936368 9.764185 NaN NaN
```

To estimate the contrast, we have to include the coefficients for all levels; I was mistaken in stating that we can leave it out the one for the reference level. Below, I have added the contrast coefficient, group size, and group effect estimate for the reference level (compared to the earlier version of the solution).

\mathbf{R} code

The 95% CI for the diff. between the two groups is (0.746, 2.13).

Python code

```
c1 = np.array([-1/3, -1/3, -1/3, 1])
n_vals = np.array([50, 87, 90, 125])

est_params = np.append([0], lm_outliers_rm.params.to_numpy()[1:])
L = np.sum(c1 * est_params)

MSW = lm_outliers_rm.mse_resid
df = lm_outliers_rm.df_resid
q1 = -stats.t.ppf(0.025, df)
se1 = np.sqrt(MSW*np.sum(c1**2 / n_vals))

lower_ci = L - q1*se1
upper_ci = L + q1*se1
print(f"""The 95% CI for the diff. between the two groups is ({lower_ci:.3f}, {upper_ci:.3f}).""")
```

The 95% CI for the diff. between the two groups is (0.746, 2.133).

SAS Code

```
proc glm data=ST2137.STUD_PERF3;
    class Medu;
    model G3=Medu / clparm;
    means Medu / hovtest=levene welch plots=none;
    lsmeans Medu / adjust=tukey pdiff alpha=.05;
    estimate 'lower_vs_higher' Medu -1 -1 -1 3 / divisor=3;
    run;
quit;
```

		The GLM I	Procedure	•			
Dependent Variable: G3							
Parameter	Estimate	Standard Error	t Value	Pr > t	95% Confidence Limits		
lower_vs_higher	1.43957088	0.35261373	4.08	<.0001	0.74604871	2.13309305	

We can see that R, Python and SAS (you will have to upload the data first) all return the same result. If we perform the estimation using the sum contrast, we should obtain exactly the same value. Here is how we can do so:

```
## Using sum contrasts:
stud_perf3$Medu2 <- stud_perf3$Medu
contrasts(stud_perf3$Medu2) <- contr.sum(4)

lm_outliers_rm_sum <- lm(G3 ~ Medu2, data=stud_perf3)
#anova(lm_outliers_rm_sum)

alpha4 <- -sum(coef(lm_outliers_rm_sum)[-1])
c1 <- c(-1/3, -1/3, -1/3, 1)
n_vals <- c(50, 87, 90, 125)
est_coef <- coef(lm_outliers_rm_sum)
L <- sum(c1*c(est_coef[2:4], alpha4))

summary_out <- anova(lm_outliers_rm)
MSW <- summary_out$ Mean Sq`[2]
df <- summary_out$Df[2]</pre>
```

```
se1 <- sqrt(MSW * sum( c1^2 / n_vals ) )
q1 <- qt(0.025, df, 0, lower.tail = FALSE)
lower_ci <- L - q1*se1
upper_ci <- L + q1*se1
cat("The 95% CI for the diff. between the two groups is (",
    format(lower_ci, digits = 3), ", ", format(upper_ci, digits = 3), ").", sep="")</pre>
```

The 95% CI for the diff. between the two groups is (0.746, 2.13).

The next portion pertains to the multiple comparison tests.

R code

```
tukey_out <- TukeyHSD(aov(lm_outliers_rm), ordered=TRUE)
tukey_out</pre>
```

Tukey multiple comparisons of means 95% family-wise confidence level factor levels have been ordered

Fit: aov(formula = lm_outliers_rm)

\$Medu

```
diff lwr upr p adj
2-1 0.8519540 -0.5795864 2.283494 0.4168866
3-1 1.0933333 -0.3294728 2.516139 0.1961010
4-1 2.0880000 0.7382076 3.437792 0.0004602
3-2 0.2413793 -0.9714333 1.454192 0.9557730
4-2 1.2360460 0.1097792 2.362313 0.0250490
4-3 0.9946667 -0.1204773 2.109811 0.0993909
```

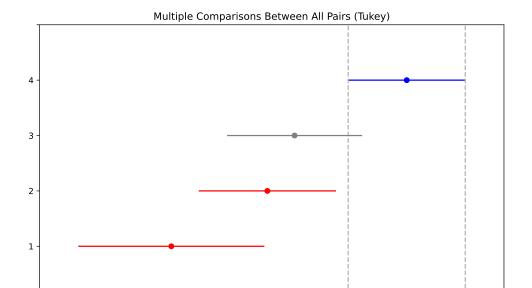
Python code

```
import statsmodels.stats.multicomp as mc

cp = mc.MultiComparison(stud_perf3.G3, stud_perf3.Medu)
tk = cp.tukeyhsd()
print(tk)
```

```
Multiple Comparison of Means - Tukey HSD, FWER=0.05
_____
group1 group2 meandiff p-adj lower upper reject
       2
   1
            0.852 0.4169 -0.5796 2.2835 False
   1
        3 1.0933 0.1961 -0.3295 2.5161 False
   1
        4 2.088 0.0005 0.7382 3.4378
        3 0.2414 0.9558 -0.9714 1.4542 False
    2
   2
            1.236 0.025 0.1098 2.3623
        4
                                    True
         4 0.9947 0.0994 -0.1205 2.1098 False
```

```
tk.plot_simultaneous(comparison_name = 4);
```



The only significant differences are between group 4 and group 1, and between group 4 and group 2.

11.0

11.5

12.0

12.5

13.0

If we were to apply the procedure on the original data, without Medu = 0:

10.5

10.0

```
lm_outliers <- lm(G3 ~ Medu, data=stud_perf2)
tukey_out2 <- TukeyHSD(aov(lm_outliers), ordered=TRUE)
tukey_out2</pre>
```

Tukey multiple comparisons of means 95% family-wise confidence level factor levels have been ordered

Fit: aov(formula = lm_outliers)

\$Medu

```
diff lwr upr p adj
2-1 1.050189 -0.8337925 2.934171 0.4762184
3-1 1.625064 -0.2727284 3.522857 0.1224640
4-1 3.085393 1.2762245 4.894561 0.0000823
3-2 0.574875 -1.0491895 2.198939 0.7977607
4-2 2.035203 0.5156448 3.554762 0.0033999
4-3 1.460328 -0.0763198 2.996977 0.0693173
```

We can see that, in terms of decisions (significant/not), the outcome does not change whether we keep the outliers in or out. However, notice that in some comparisons the p-value goes up and in some it goes down. In general, please try to make this procedure a habit - consider the analysis with and without any outliers to see how significant they were.