

Experimental Design and Data Analysis:

Assignment 4

Andrew Bedard(2566978) & Simone van Gompel(2567525)
Group 19

May 9, 2015

Exercise 1

1

To randomise the 6 combinations of conditions for the slices of bread, we may implement the following R code.

```
sample_slices = sample(1:18, 18)
```

Which simply samples 18 numbers from the range 1-18 randomly. In this way, if we have the

2

From the data bread.txt we make boxplots of hours before bread becomes mouldy versus the two factors, temperature environment and humidity:

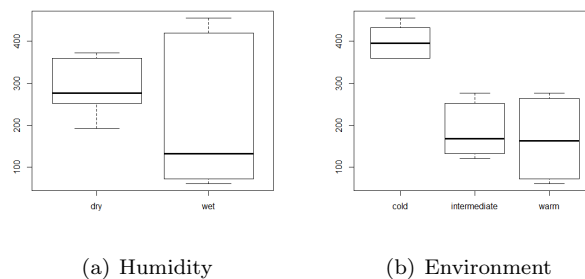


Figure 1: Boxplots of Hours with Humidity and Environment

There are a few notable features in these boxplots, mainly that the variances in the two humidity factors are significantly different, and secondly that there is zero overlap between the values obtained from the cold environment and the other two.

3

Next, by creating interaction plots, by letting one of our factors be the continuous x-axis, and plotting for individual values of the other factor, we obtain the following:

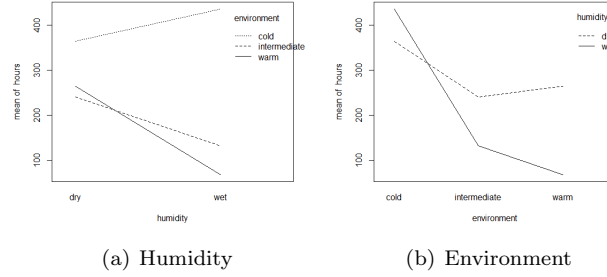


Figure 2: Interactionplots of Hours with Humidity and Environment

Notice in Fig 2:(a) the plot of cold environment is significantly different from those of the other two environments, and again in 2:(b) where the lines for wet and dry cross. Given the relatively extreme values seen in the box plots 1, it is plausible there is an interaction between environment and humidity taking place.

4

Performing an ANOVA test, to test the null hypothesis that the environment, humidity and the interaction between the two are insignificant we obtain the following:

Analysis of Variance Table

Response: hours

	Df	Sum Sq	Mean Sq	F value	Pr(>F)	
environment	2	201904	100952	233.685	2.461e-10	***
humidity	1	26912	26912	62.296	4.316e-06	***
environment: humidity	2	55984	27992	64.796	3.705e-07	***
Residuals	12	5184	432			

The results are quite compelling, we reject all the null hypothesis with all p-values $\ll 0.5$, therefore there is an effect of environment, humidity as well as an interaction between the two.

5

This interaction effect, which can be more easily seen in Fig:2, appears to be The interaction effect of both factors are significant. This means that they don't have an additive effect on each other. So if you change both the factors, it will have a negative effect on the hours.

6

The environment has the biggest numerical influence. This is not a good question, it is always better to look at the relative influence of a factor.

7

In Fig3 the QQ-plot of the residuals is shown. Here you can see that the data is roughly normally distributed. Only the most left and right data points seem to be out of place.

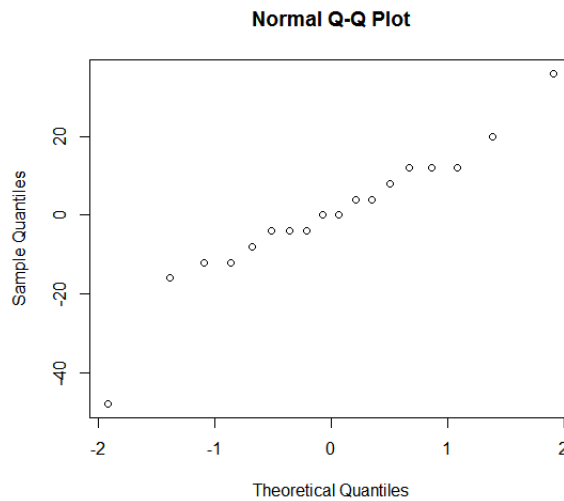


Figure 3: QQ-plot of the residuals

8

In Fig4 the fitted values versus residual values are shown. It seems that there is no heteroscedasticity present in this data.

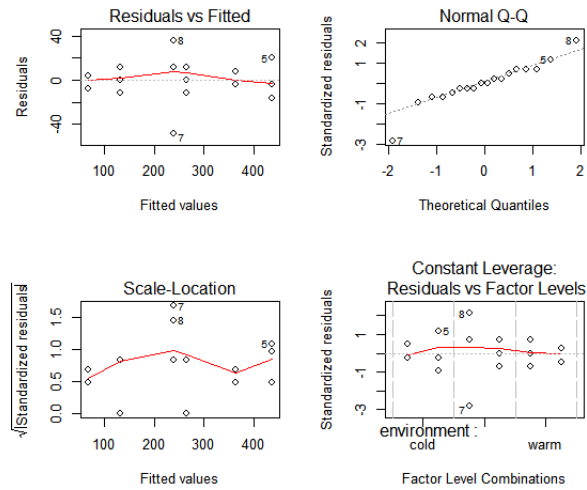


Figure 4: Fitted Values VS Residual Values

Exercise 2

- 1
- 2

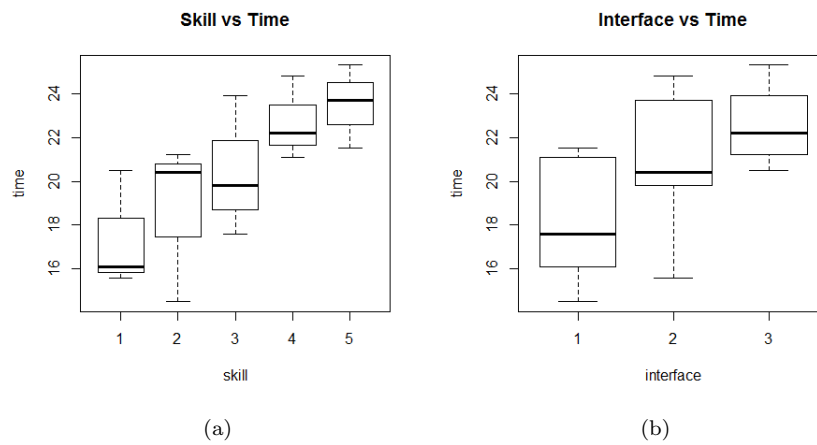


Figure 5: Box Plots of Skill and Interface vs Time

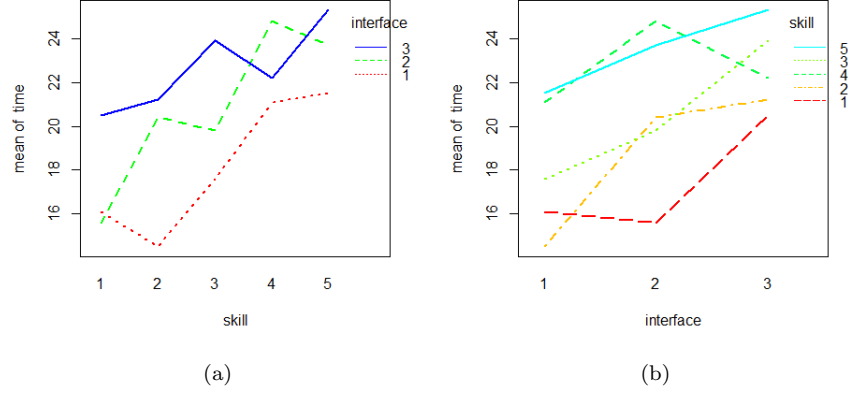


Figure 6: Interaction plots of Skill and Interface vs Time

It is difficult to conclude that there is an interaction between skill and interface as they are clearly not parallel, but they follow the same general trajectory, so this interaction may be due to noise.

3

Using the Kruskal-Wallis rank sum test, we test if the distributions of our populations in regards to the time measured for different interfaces are the same, and we obtain the following results:

Kruskal-Wallis **rank sum** test

data: **time** and **interface**

Kruskal-Wallis **chi-squared** = 4.22, **df** = 2, **p-value** = 0.1212

Thus with a p-value of 0.1212 we reject the null hypothesis that our populations are the same, therefore the search time for all interfaces is not equal.

4

5

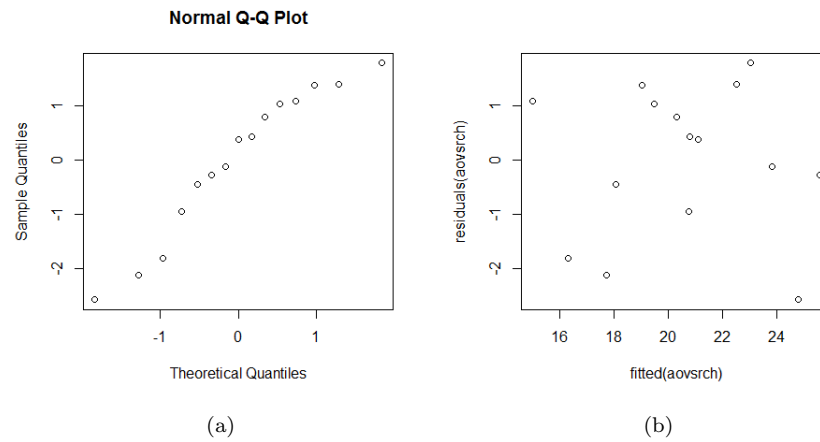


Figure 7: Diagnostic Plots

It is difficult to say for certain, there may be a slight curve in the qq-plot 7(a) but it looks approximately normal, and the fitted value plot 7(b) suggests there is no significant difference in the population variances.

6

Friedman **rank sum** test

data: **time**, interface and skill

Friedman chi-squared = 6.4, **df** = 2, p-value = 0.04076

With a p-value of 0.04076 we reject the null hypothesis, thus we conclude there is an effect of interfaces.

7

Testing the null hypothesis that the search time is the same for all interface by a ANOVA test, ignoring skill we obtain the following:

Analysis of Variance Table

Response: **time**

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
interface	2	50.465	25.233	2.8605	0.09642
Residuals	12	105.852	8.821		

With a p-value of 0.09642 we do not reject the null hypothesis, thus the time is not the same for all interfaces. This test is useful to use, if some conditions are met, as of right now, we do not have convincing evidence there is no interaction between skill and interface. It would be more useful to test along with the variable skill, and the interaction between the two as follows:

Analysis of Variance Table

Response: **time**

	Df	Sum Sq	Mean Sq	F value	Pr(>F)	
interface	1	49.729	49.729	21.4145	0.0007313	***
skill	1	78.732	78.732	33.9039	0.0001154	***
interface:skill	1	2.312	2.312	0.9956	0.3398204	
Residuals	11	25.544	2.322			

This way we are able to measure the interaction between the data sets, and determine whether there is a factor which has a greater effect on our results. From the previous table we see that with a p-value of 0.3398 we accept the null hypothesis that there is no significant interaction between skill in and interface, which is what we require for the one-way ANOVA ignoring skill to be valid.

Exercise 3

1

Analysis of Variance Table

Response: acidity

	Df	Sum Sq	Mean Sq	F value	Pr(>F)	
starter	4	44.136	11.0340	8.0835	0.001106	**
batch	1	4.855	4.8547	3.5566	0.078826	.
position	4	2.348	0.5870	0.4300	0.784786	
Residuals	15	20.475	1.3650			

2

Call:

```
lm(formula = acidity ~ starter + batch + position, data = cream)
```

Residuals:

	Min	1Q	Median	3Q	Max
	-1.7512	-0.7596	0.0132	0.8816	1.0856

Coefficients:

Estimate Std. Error t value Pr(>|t|)

(Intercept)	7.8260	0.8586	9.115	1.67e-07	***
starter2	-0.1500	0.7389	-0.203	0.84186	
starter3	-0.9800	0.7389	-1.326	0.20459	
starter4	2.8100	0.7389	3.803	0.00173	**
starter5	-0.4840	0.7389	-0.655	0.52238	
batch	0.3116	0.1652	1.886	0.07883	.
position2	-0.6180	0.7389	-0.836	0.41608	
position3	-0.0380	0.7389	-0.051	0.95966	
position4	-0.7640	0.7389	-1.034	0.31755	
position5	-0.2640	0.7389	-0.357	0.72586	

Residual standard error: 1.168 on 15 degrees of freedom
Multiple R-squared: 0.7149, Adjusted R-squared: 0.5438
F-statistic: 4.179 on 9 and 15 DF, p-value: 0.007304

3

Linear Hypotheses:

	Estimate	Std. Error	t value	Pr(> t)	
2 - 1 == 0	-0.1500	0.4673	-0.321	0.997	
3 - 1 == 0	-0.9800	0.4673	-2.097	0.282	
4 - 1 == 0	2.8100	0.4673	6.013	<0.001	***
5 - 1 == 0	-0.4840	0.4673	-1.036	0.834	
3 - 2 == 0	-0.8300	0.4673	-1.776	0.429	
4 - 2 == 0	2.9600	0.4673	6.334	<0.001	***
5 - 2 == 0	-0.3340	0.4673	-0.715	0.949	
4 - 3 == 0	3.7900	0.4673	8.110	<0.001	***
5 - 3 == 0	0.4960	0.4673	1.061	0.822	
5 - 4 == 0	-3.2940	0.4673	-7.048	<0.001	***

Starter 1 and 4 produce significantly different acidity. This is apparent first from exercise 3.2 we obtain: $\mu_1 = 7.8260$ and $\mu_4 = 7.8260 + 2.8100 = 10.636$ which are the estimates for starter 1 and 4 respectively. Furthermore the p-value for starter 1 obtained from exercise 3.2, of 1.67e-07 suggests that we reject the null hypothesis that our estimate for starter 1 is equal to that of the rest of the population, and the p-values obtained from the simultaneous method, of 0.001 suggests that we reject the null hypothesis that the estimate for the treatment effect of 4 is equal to each of the other respective treatment effects.

4

The p-value for the test $H_0 : \alpha_2 = \alpha_1$, where α_i is our estimate of the treatment effects is 0.997, whereas in exercise 3.2 our p-value for the null hypothesis $H_0 : \alpha_2 = \alpha_1$ is 0.84186. It is no coincidence that the p-value obtained from exercise 3.2, is different from that of the simultaneous calculation, this is due to the fact that when calculating the simultaneous p-values for the null hypothesis $H_0 : \alpha_i = \alpha_1$ we are doing so in such a way that the probability of rejecting

any null hypothesis in error is less than 0.5. In contrast to the method used in exercise 3.2, where for every null hypothesis our chance of making such an error is $N * 0.05$ where N is the number of possibilities to make such an error. In this way the Simultaneous p-values method gives us a higher confidence in our conclusions.

5

Linear Hypotheses:

	Estimate	lwr	upr
2 - 1 == 0	-0.1500	-1.6391	1.3391
3 - 1 == 0	-0.9800	-2.4691	0.5091
4 - 1 == 0	2.8100	1.3209	4.2991
5 - 1 == 0	-0.4840	-1.9731	1.0051
3 - 2 == 0	-0.8300	-2.3191	0.6591
4 - 2 == 0	2.9600	1.4709	4.4491
5 - 2 == 0	-0.3340	-1.8231	1.1551
4 - 3 == 0	3.7900	2.3009	5.2791
5 - 3 == 0	0.4960	-0.9931	1.9851
5 - 4 == 0	-3.2940	-4.7831	-1.8049

The confidence intervals for testing all differences $\alpha_j - \alpha_{j'}$ for $(i, i' \in 1, 2, \dots, 5)$ of the main effects for starter with simultaneous confidence level 95% are (4-1),(4-2),(4-3),(4-5),(5-4) so all those containing starter 4. This is likely due to the fact that the estimated value for starter 4, which was earlier calculated as 10.636, is so much larger than every other estimate, for example all the confidence intervals for (4-i) are greater than 0, and the one interval (5-4) is less than 0.

Exercise 4

1

Performing ANOVA test using an ordinary fixed effects model fitted with lm, the follow results were obtained:

Analysis of Variance Table

Response: milk

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
treatment	1	0.27	0.269	0.0016	0.9682
order	1	53.52	53.515	0.3268	0.5760
Residuals	15	2456.69	163.779		

Residuals:

Min	1Q	Median	3Q	Max
-16.602	-10.103	0.170	9.431	22.042

Coefficients:

	Estimate	Std. Error	t value	Pr(> t)
(Intercept)	37.1722	5.4379	6.836	5.64e-06 ***
treatmentB	-0.2444	6.0329	-0.041	0.968
orderBA	-3.4700	6.0704	-0.572	0.576

Signif. codes:

0	***	0.001	**	0.01	*	0.05	.	0.1	1
---	-----	-------	----	------	---	------	---	-----	---

Residual standard error: 12.8 on 15 degrees of freedom

Multiple R-squared: 0.02142, Adjusted R-squared: -0.1091

F-statistic: 0.1642 on 2 and 15 DF, p-value: 0.8501

We can conclude from the F-value and p-value of 0.0016 and 0.9682 respectively for the treatment effect on milk production, we do not reject the null hypothesis, therefore there is no treatment effect on milk production. Further we observe that the p-value for the null hypothesis that the individual treatments effects are all greater than 0.05, thus we accept the null hypothesis that the treatment effects are equal. This stands to reason as if we do not conclude there is any effect for all treatments, there would be no effect for individual treatments when sample sizes are equal.

2

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.
17.10	26.82	33.75	35.12	43.02	55.50

3

Analysis of Variance Table

	Df	Sum Sq	Mean Sq	F value
treatment	1	0.26889	0.26889	0.0503
order	1	0.82889	0.82889	0.1552

AIC	BIC	logLik	deviance	df.resid
125.4	129.9	-57.7	115.4	13

Scaled residuals:

Min	1Q	Median	3Q	Max
-1.31202	-0.45325	-0.01847	0.33881	1.56294

Random effects:

Groups	Name	Variance	Std.Dev.
id	(Intercept)	169.755	13.029
Residual		5.341	2.311

Number of obs: 18, groups: id, 9

Fixed **effects**:

	Estimate	Std. Error	t value
(Intercept)	37.1722	6.5881	5.642
treatmentB	-0.2444	1.0895	-0.224
orderBA	-3.4700	8.8086	-0.394

Correlation of Fixed Effects:

	(Intr)	trtmnB
treatmentB	-0.083	
orderBA	-0.743	0.000

4

Paired **t**-test

data: milk[treatment == "A"] and milk[treatment == "B"]
t = 0.2244, **df** = 8, p-value = 0.8281
alternative hypothesis: true difference in means **is not equal** to 0
95 percent confidence interval:
-2.267910 2.756799
sample estimates:
mean of the differences 0.2444444

1 R-Code

1.1 Exercise 1

```
bread = read.table("bread.txt")
hours = bread$hours
humidity = bread$humidity
environment = bread$environment
```

```
#1.1
sample_slices = sample(1:18, 18)
```

```
#1.2
boxplot(hours~humidity)
boxplot(hours~environment)
```

```
#1.3
interaction.plot(humidity, environment, hours)
interaction.plot(environment, humidity, hours)
```

```
#1.4
```

```
aovenvhum = lm(hours ~ environment+humidity, data=bread)
anova(aovenvhum)
```

```
aovenv = lm(hours ~ environment, data=bread)
anova(aovenv)
```

```
aovhum = lm(hours ~ humidity, data=bread)
anova(aovhum)
```

```
#1.5
residuals(aovhum)
qqnorm(residuals(aovhum))
qqnorm(residuals(aovenv))
qqnorm(residuals(aovenvhum))
```

1.2 Exercise 2

#Note: The interface is of greatest importance

```
search = read.table("search.txt")
```

```
time = search$time
skill = search$skill
interface = search$interface
```

```
#2.1
students = 1:15
# set the lvls of the students
skill_lvls = rep(c(1,2,3,4,5), each=3)
# sample the students
samples = sample(students, 15)
# make the block_design
block_design = matrix(c(samples[0:5],
                        samples[6:10],
                        samples[11:15])),
                    nrow = 3, ncol = 5)
skill_block_design = matrix(c(skill_lvls[samples[0:5]],
                        skill_lvls[samples[6:10]],
                        skill_lvls[samples[11:15]]),
                    nrow = 3, ncol = 5)
```

```
#2.2
```

```
boxplot(time~skill, xlab="skill", ylab="time", main="Skill_vs_Time")
boxplot(time~interface, xlab="interface", ylab="time", main="Interface_vs_Time")
```

```

interaction.plot( skill , interface , time , lwd=2, col=rainbow(3))
interaction.plot( interface , skill , time , lwd=2, col=rainbow(8))

```

```

#2.3
interface = factor(search$interface)
skill = factor(search$skill)
kruskal.test(time , interface)

```

```

#2.4

aovsrch=lm(time~ skill+interface)
summary( aovsrch)

```

```

#2.5

qqnorm(residuals( aovsrch))
plot( fitted( aovsrch) , residuals( aovsrch))

```

```

#2.6

friedman.test(time , interface , skill)

```

```

#2.7
anovinter = lm(time~ interface)
anova(anovinter)

anv = lm(time~ interface*skill , data=search)
anova(anv)

```

1.3 Exercise 3

```

library(multcomp)

cream = read.table("cream.txt" , header=T)
cream$starter = factor(cream$starter)
cream$position = factor(cream$position)

```

```

#3.1

anovacid = lm(acidity~ starter+batch+position , data=cream)
anova(anovacid)

```

```

#3.2
summary(anovacid)

```

```
#3.3
tmp = glht(creamlm, linfct=mcp(starter="Tukey"))
summary(tmp)
```

```
#3.4
confint(tmp)
```

1.4 Exercise 4

```
library(lme4)

cow = read.table("cow.txt", header=T)
cow$order = factor(cow$order)
order = cow$order
milk = cow$milk
cow$treatment = factor(cow$treatment)
treatment = cow$treatment

#4.1
cowlm = lm(milk ~ treatment + order, data=cow)
anova(cowlm)
summary(cowlm)

#4.2
summary(cow$milk)

#4.3
cowlmer = lmer(milk ~ treatment + order + (1|id), data=cow, REML=FALSE)
anova(cowlmer)
summary(cowlmer)

#4.4
attach(cow)
t.test(milk[treatment=="A"], milk[treatment=="B"], paired=TRUE)
```