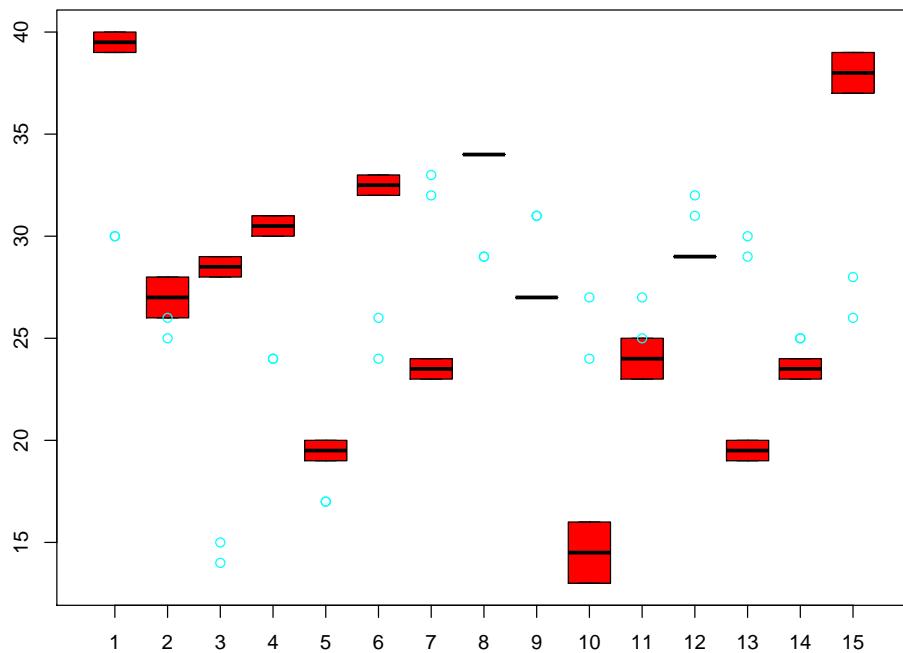


Solution to Series 5

1. a) Plot the data. Read in the data:

```
> paint <- read.table(file="http://stat.ethz.ch/Teaching/Datasets/paint.txt", header=TRUE)
> paint$SAMPLE <- as.factor(paint$SAMPLE)
> paint$BATCH <- as.factor(paint$BATCH)

> plot(paint$BATCH[paint$SAMPLE==1], paint$MOISTURE[paint$SAMPLE==1], col=2)
> points(paint$BATCH[paint$SAMPLE==2], paint$MOISTURE[paint$SAMPLE==2], col=5)
```



- b) Write down the analysis of variance model. Why are all factors random?

$$Y_{ijk} = \mu + \text{batch}_i + \text{sample}_{j(i)} + \epsilon_{ijk}$$

batch_i random effect of BATCH i , $\text{batch}_i \sim \mathcal{N}(0, \sigma_{\text{batch}}^2)$,
 $\text{sample}_{j(i)}$ random effect of SAMPLE j , $\text{sample}_{j(i)} \sim \mathcal{N}(0, \sigma_{\text{sample}}^2)$,
 ϵ_{ijk} measurement error, $\epsilon_{ijk} \sim \mathcal{N}(0, \sigma_e^2)$.

All factors are random because BATCHes are chosen at random from the population of batches and SAMPLEs are chosen from the population of all samples.

- c) Construct the skeleton analysis of variance table with degrees of freedom and expected values.

Score	df	$E(MS)$
batch	14	$\sigma_e^2 + 2\sigma_{\text{sample}}^2 + 4\sigma_{\text{batch}}^2$
batch:sample	15	$\sigma_e^2 + 2\sigma_{\text{sample}}^2$
Residuals	30	σ_e^2
Total	59	

- d) Calculate estimates for σ_{BATCH}^2 and σ_{SAMPLE}^2 . Compute the F test statistics, first by hand and then use R .

```
> mod1 <- aov(MOISTURE~BATCH/SAMPLE, data=paint)
> summary(mod1)
```

```
Df Sum Sq Mean Sq F value Pr(>F)
BATCH      14 1210.9   86.50   94.36 <2e-16 ***
BATCH:SAMPLE 15  869.7   57.98   63.26 <2e-16 ***
Residuals    30   27.5    0.92
---
Signif. codes:
0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

$$\hat{\sigma}_e^2 = 0.917$$

$$\hat{\sigma}_{sample}^2 = (57.983 - 0.917)/2 = 28.533$$

$$\hat{\sigma}_{batch}^2 = (86.495 - 57.983)/4 = 7.128$$

$$F_{\sigma_{batch}^2=0} = \frac{\hat{\sigma}_e^2 + 2\hat{\sigma}_{sample}^2 + 4\hat{\sigma}_{batch}^2}{\hat{\sigma}_e^2 + 2\hat{\sigma}_{sample}^2} = \frac{0.917 + 2 \cdot 28.533 + 4 \cdot 7.128}{0.917 + 2 \cdot 28.533} = 1.4917$$

$$F_{\sigma_{sample}^2=0} = \frac{\hat{\sigma}_e^2 + 2\hat{\sigma}_{sample}^2}{\hat{\sigma}_e^2} = 63.255$$

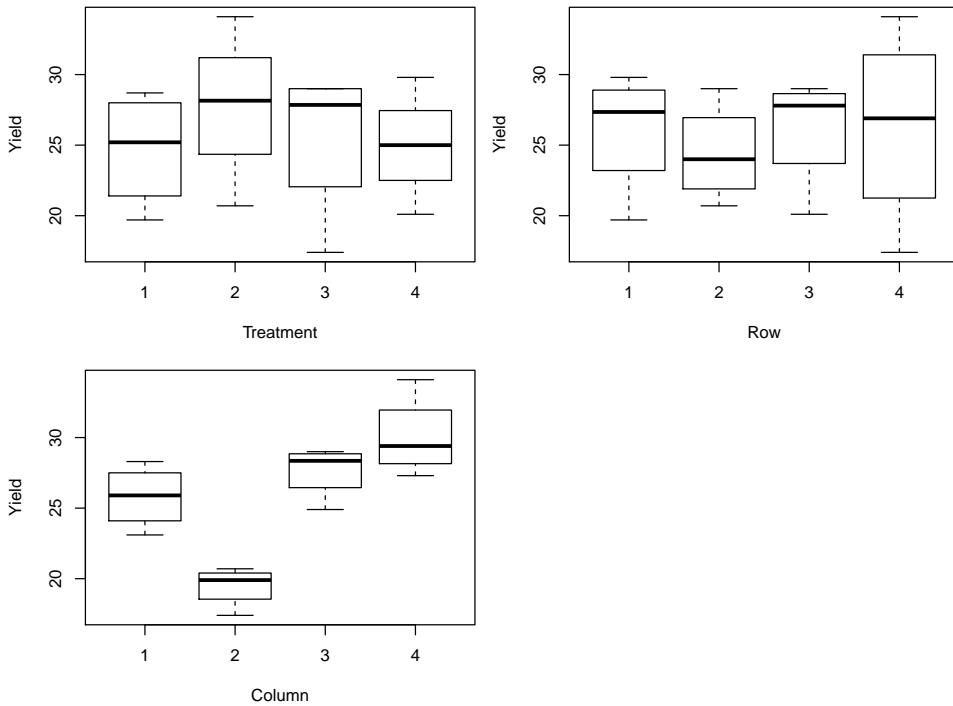
```
> mod2 <- aov(MOISTURE~BATCH+Error(SAMPLE%in% BATCH), data=paint)
> summary(mod2)

Error: SAMPLE:BATCH
      Df Sum Sq Mean Sq F value Pr(>F)
BATCH      14 1210.9   86.50   1.492  0.226
Residuals  15  869.7   57.98

Error: Within
      Df Sum Sq Mean Sq F value Pr(>F)
Residuals 30   27.5   0.9167
```

2. a) Plot the data.

```
> peanut <- read.table(file="http://stat.ethz.ch/Teaching/Datasets/Peanut.txt", header=TRUE)
> peanut$Row <- as.factor(peanut$Row)
> peanut$Column <- as.factor(peanut$Column)
> peanut$Treatment <- as.factor(peanut$Treatment)
> par(mfrow=c(2,2))
> plot(Yield ~ Treatment+Row+Column, data=peanut)
```



b) Carry out the analysis of variance and report your findings.

```
> modP1 <- aov(Yield ~ Treatment+Row+Column, data=peanut)
> summary(modP1)
```

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
Treatment	3	23.42	7.81	1.953	0.22255
Row	3	9.43	3.14	0.786	0.54394
Column	3	245.91	81.97	20.507	0.00148 **
Residuals	6	23.98	4.00		

Signif. codes:

0 ‘***’ 0.001 ‘**’ 0.01 ‘*’ 0.05 ‘.’ 0.1 ‘ ’ 1

We see that the factor Treatment is not significant.

c) Treatment 1 is the control treatment. Even though the anova F is not significant, do any of the other treatments have a significantly higher yield, if a one-sided test is used?

```
> TukeyHSD(modP1, "Treatment", conf.level=0.90)
Tukey multiple comparisons of means
90% family-wise confidence level
```

```
Fit: aov(formula = Yield ~ Treatment + Row + Column, data = peanut)
```

\$Treatment	diff	lwr	upr	p adj
2-1	3.075	-0.9887413	7.138741	0.2320297
3-1	0.825	-3.2387413	4.888741	0.9334796
4-1	0.275	-3.7887413	4.338741	0.9971031
3-2	-2.250	-6.3137413	1.813741	0.4479925
4-2	-2.800	-6.8637413	1.263741	0.2914481
4-3	-0.550	-4.6137413	3.513741	0.9782233

If we look at the first three rows of the Treatment differences table we see that all of the confidence intervals of 2-1, 3-1 and 4-1 contain 0. So we can say that none of the one-sided tests are significant. That is, none of the other treatments have a significantly higher yield compared to the control treatment number 1.

The breeder decided to replicate the experiment. He used three blocks at different locations in the field, but in each location the north-south and east-west gradients were identified. The data are in the file Peanut2.txt.

- d) Which factors are nested, which are crossed? The factors Column and Row are nested in the factor Rep which represents the three different positions on the field.
- e) Give an analysis of variance table and test whether any pairwise differences are significant.

```
> peanut2 <- read.table(file="http://stat.ethz.ch/Teaching/Datasets/Peanut2.txt", header=TRUE)
> peanut2$Row <- as.factor(peanut2$Row)
> peanut2$Column <- as.factor(peanut2$Column)
> peanut2$Treatment <- as.factor(peanut2$Treatment)
> peanut2$Rep <- as.factor(peanut2$Rep)
> modP2 <- aov(Yield ~ Treatment+Rep/(Row+Column), data=peanut2)
> summary(modP2)

Df Sum Sq Mean Sq F value    Pr(>F)
Treatment     3   95.7   31.92   3.962   0.0199 *
Rep           2   15.8    7.92   0.983   0.3888
Rep:Row       9   66.1    7.35   0.912   0.5311
Rep:Column    9  530.8   58.97   7.321 4.55e-05 ***
Residuals    24  193.3    8.06
---
Signif. codes:
0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
> TukeyHSD(modP2, "Treatment", conf.level=0.90)
Tukey multiple comparisons of means
90% family-wise confidence level

Fit: aov(formula = Yield ~ Treatment + Rep/(Row + Column), data = peanut2)

$Treatment
      diff      lwr      upr      p adj
2-1  3.36666667  0.5619194  6.17141398 0.0363778
3-1  0.50000000 -2.3047473  3.30474732 0.9724701
4-1 -0.06666667 -2.8714140  2.73808065 0.9999296
3-2 -2.86666667 -5.6714140 -0.06191935 0.0899752
4-2 -3.43333333 -6.2380806 -0.62858602 0.0320424
4-3 -0.56666667 -3.3714140  2.23808065 0.9607866

We see that Treatment 2 is significantly better than the control.
```

3. To do an anova fit you can reshape your data as follows.

```
> tennis.max=reshape(tennis[,c(1:5,9,13)], varying=c("max1", "max2", "max3"), idvar="id",
  timevar="period", v.names="max", direction="long")
> tennis.max$Treatment[tennis.max$order==1 & tennis.max$period==1]="Motrin"
> tennis.max$Treatment[tennis.max$period==2]="Washout"
> tennis.max$Treatment[tennis.max$order==1 & tennis.max$period==3]="Placebo"
> tennis.max$Treatment[tennis.max$order==2 & tennis.max$period==3]="Motrin"
> tennis.max$Treatment[tennis.max$order==2 & tennis.max$period==1]="Placebo"
> tennis.max$Treatment=factor(tennis.max$Treatment)
> tennis.max$id=factor(tennis$id)
> tennis.max$sex=factor(tennis$sex)
> tennis$sex[tennis$sex==1] <- 'male'
> tennis$sex[tennis$sex==2] <- 'female'
> tennis.max$period=factor(tennis.max$period)
```

Is the treatment with Motrin efficient?

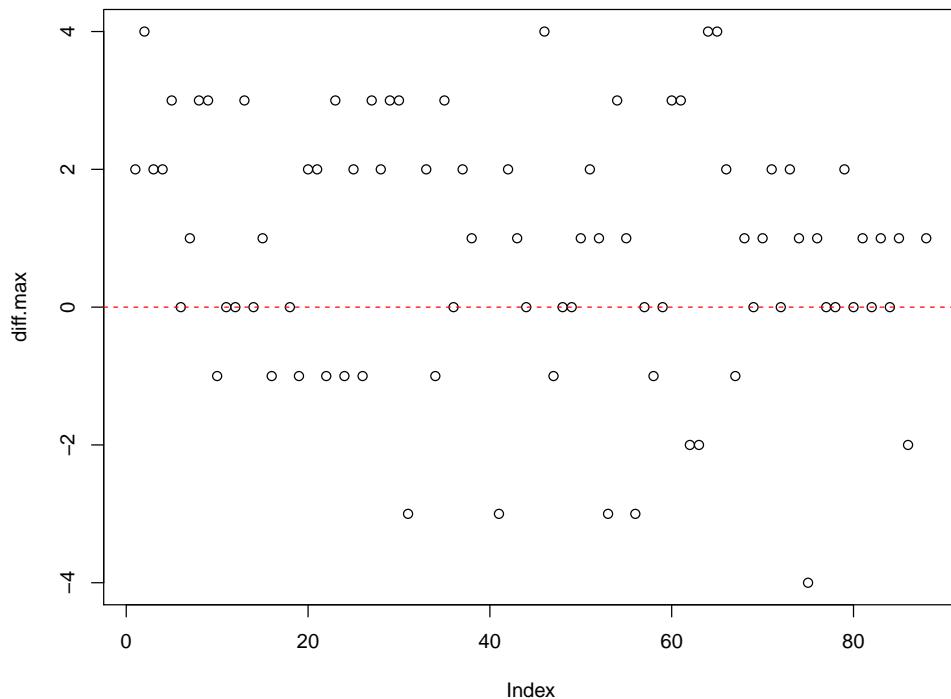
To find this out: First do a pairwise t.test and wilcox.test of the four different outcomes (max, ave, etc.). Then do an anova fit for only the maximum activity comparison taking into account id, period and treatment.

Is the washout period long enough or could there still be a carry-over effect?

Read in the data:

```
> tennis <- read.table(file="http://stat.ethz.ch/Teaching/Datasets/TENNIS.dat")
> names(tennis)=c("id", "age", "sex", "order", "max1", "twelve1", "ave1", "overall1",
  "max2", "twelve2", "ave2", "overall2", "max3", "twelve3", "ave3", "overall3")
> ## replace invalid values with NA:
> head(tennis)
> for (i in 3:16)
  tennis[,i][tennis[,i]==9 | tennis[,i]==0]=NA##identify the invalid values

> diff.max=tennis$max1-tennis$max3
> diff.max[tennis$order==2]=-diff.max[tennis$order==2]
> plot(diff.max)
> abline(h=0,col=2,lty=2)
```



```
> summary(diff.max)
```

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.	NA's
-4.0000	0.0000	1.0000	0.7927	2.0000	4.0000	6

```
> t.test(diff.max)
```

One Sample t-test

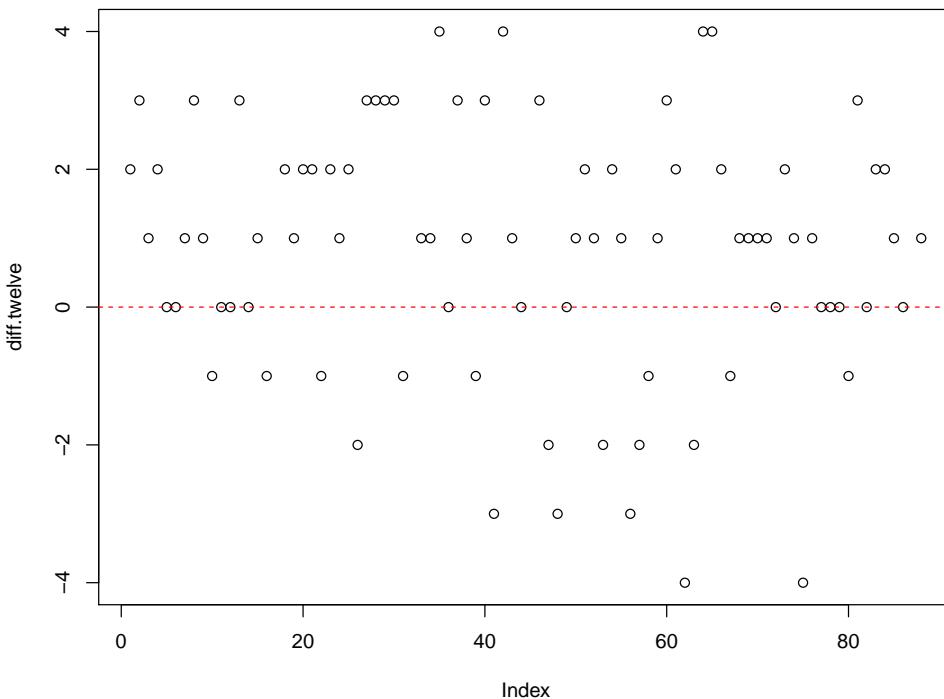
```
data: diff.max
t = 3.9354, df = 81, p-value = 0.0001746
alternative hypothesis: true mean is not equal to 0
95 percent confidence interval:
 0.3919132 1.1934527
sample estimates:
mean of x
0.7926829
```

```
> wilcox.test(diff.max)
```

Wilcoxon signed rank test with continuity correction

```
data: diff.max
V = 1578.5, p-value = 0.000271
alternative hypothesis: true location is not equal to 0

> diff.twelve=tennis$twelve1-tennis$twelve3
> diff.twelve[tennis$order==2]=-diff.twelve[tennis$order==2]
> plot(diff.twelve)
> abline(h=0,col=2,lty=2)
```



```
> summary(diff.twelve)
```

	Min.	1st Qu.	Median	Mean	3rd Qu.	Max.	NA's
	-4.0000	0.0000	1.0000	0.7976	2.0000	4.0000	4

```
> t.test(diff.twelve)
```

One Sample t-test

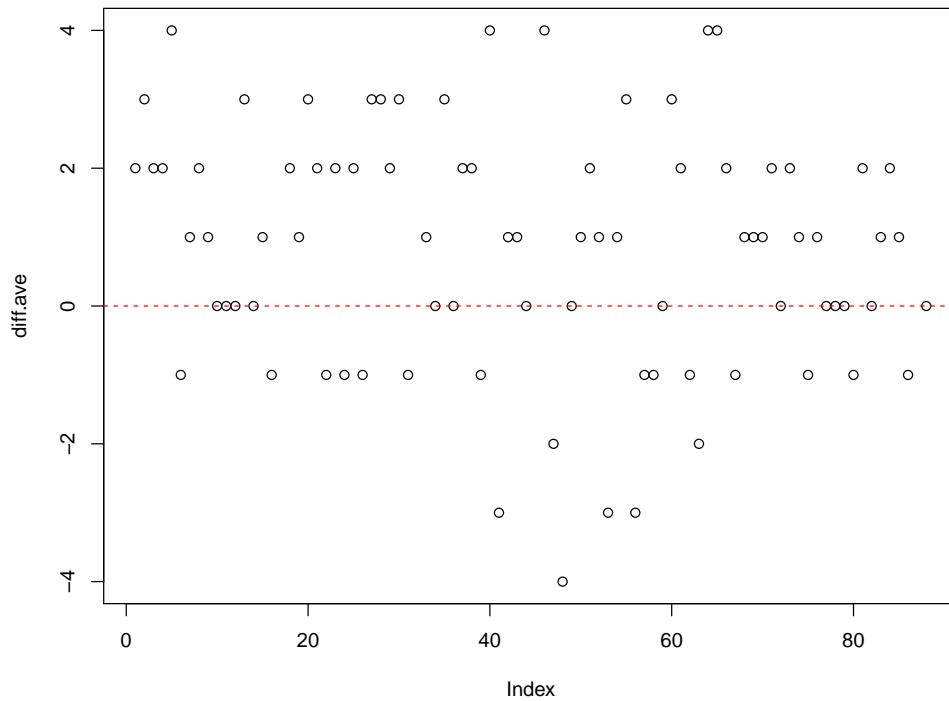
```
data: diff.twelve
t = 3.9551, df = 83, p-value = 0.0001604
alternative hypothesis: true mean is not equal to 0
95 percent confidence interval:
0.3965036 1.1987345
sample estimates:
mean of x
0.797619
```

```
> wilcox.test(diff.twelve)
```

Wilcoxon signed rank test with continuity correction

```
data: diff.twelve
V = 1855, p-value = 0.0002831
alternative hypothesis: true location is not equal to 0
```

```
> diff.ave=tennis$ave1-tennis$ave3
> diff.ave[tennis$order==2]=-diff.ave[tennis$order==2]
> plot(diff.ave)
> abline(h=0,col=2,lty=2)
```



```
> summary(diff.ave)
```

	Min.	1st Qu.	Median	Mean	3rd Qu.	Max.	NA's
	-4.0000	0.0000	1.0000	0.8214	2.0000	4.0000	4

```
> t.test(diff.ave)
```

One Sample t-test

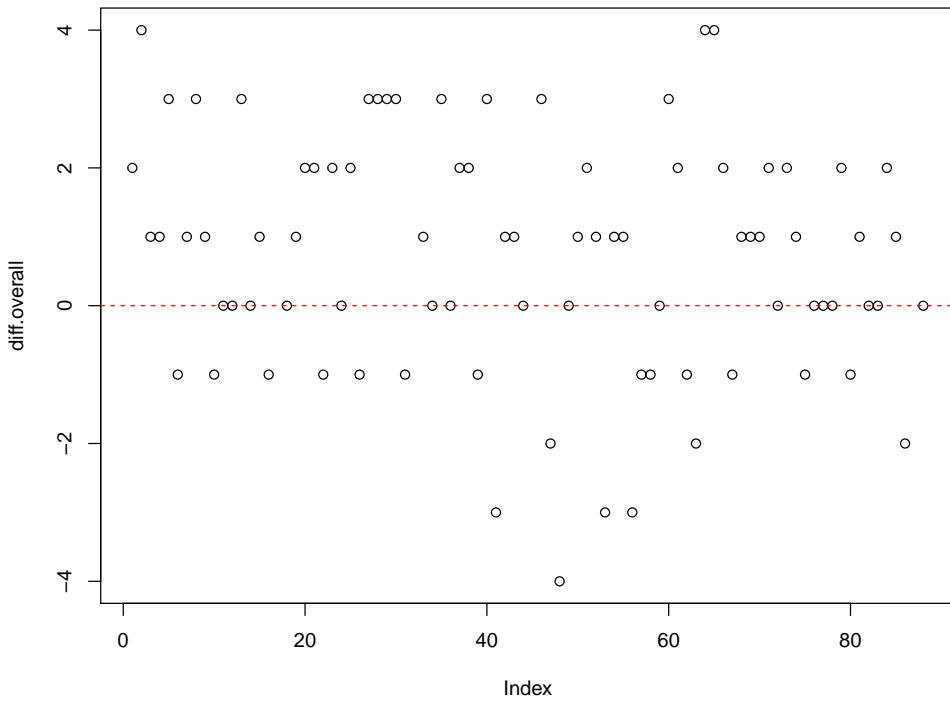
```
data: diff.ave
t = 4.2674, df = 83, p-value = 5.218e-05
alternative hypothesis: true mean is not equal to 0
95 percent confidence interval:
0.4385796 1.2042776
sample estimates:
mean of x
0.8214286
```

```
> wilcox.test(diff.ave)
```

Wilcoxon signed rank test with continuity correction

```
data: diff.ave
V = 1869, p-value = 6.026e-05
alternative hypothesis: true location is not equal to 0
```

```
> diff.overall=tennis$overall1-tennis$overall3
> diff.overall[tennis$order==2]=-diff.overall[tennis$order==2]
> plot(diff.overall)
> abline(h=0,col=2,lty=2)
```



```
> summary(diff.overall)
```

Min.	1st Qu.	Median	Mean	3rd Qu.	Max.	NA's
-4.0000	0.0000	1.0000	0.7143	2.0000	4.0000	4

```
> t.test(diff.overall)
```

One Sample t-test

```
data: diff.overall
t = 3.7938, df = 83, p-value = 0.0002808
alternative hypothesis: true mean is not equal to 0
95 percent confidence interval:
0.339812 1.088759
sample estimates:
mean of x
0.7142857
```

```
> wilcox.test(diff.overall)
```

Wilcoxon signed rank test with continuity correction

```
data: diff.overall
V = 1705.5, p-value = 0.0003242
alternative hypothesis: true location is not equal to 0
```

We see that the degree of pain while on Motrin significantly better than degree of pain on placebo for all four pain measurements.

Anova approach for max, ave, twelve and overall Example for max:

```
> tennis.max=reshape(tennis[,c(1:5,9,13)],varying=c("max1","max2","max3"),idvar="id",
+ timevar="period",v.names="max",direction="long")
> tennis.max$Treatment[tennis.max$order==1 & tennis.max$period==1]="Motrin"
```

```

> tennis.max$Treatment[tennis.max$period==2]="Washout"
> tennis.max$Treatment[tennis.max$order==1 & tennis.max$period==3]="Placebo"
> tennis.max$Treatment[tennis.max$order==2 & tennis.max$period==3]="Motrin"
> tennis.max$Treatment[tennis.max$order==2 & tennis.max$period==1]="Placebo"
> tennis.max$Treatment=factor(tennis.max$Treatment)
> tennis.max$id=factor(tennis$id)
> ##tennis.max$sex[tennis.max$sex==9]=NA##not used anyway
> tennis.max$sex=factor(tennis$sex)
> levels(tennis.max$sex)=c("male","female")
> tennis.max$period=factor(tennis.max$period)
> mod1=aov(max~id+period+Treatment,data=tennis.max[tennis.max$period!=2,])
> summary(mod1)

```

	Df	Sum Sq	Mean Sq	F value	Pr(>F)					
id	86	181.32	2.108	1.576	0.0203 *					
period	1	29.03	29.030	21.703	1.25e-05 ***					
Treatment	1	24.46	24.460	18.286	5.22e-05 ***					
Residuals	80	107.01	1.338							

Signif. codes:										
0	'***'	0.001	'**'	0.01	'*'	0.05	.	0.1	'.'	1
7	observations deleted due to missingness									

```
> model.tables(mod1,type="means")
```

Tables of means
Grand mean

3.449704

id	701	702	704	705	706	707	708	712	713	718	720	721	723	725
	4	2	5	4	3	1.5	5	3	1.5	4	5	1.5	2.5	4
rep	2	2	1	2	2	2.0	2	2	2.0	2	2	2.0	2.0	2
	727	729	732	733	734	735	736	740	741	742	744	745	746	747
	2	3	2	3.5	5	2	3.5	3.5	3.5	2.5	2	3	3.5	5
rep	2	2	2	2.0	2	1	2.0	2.0	2.0	2.0	2	2	2.0	2
	749	752	756	759	761	762	763	766	767	768	769	771	772	774
	2.5	4.5	5	2.5	4	4	4.5	3.5	4.5	3	4.5	2.5	1.5	3
rep	2.0	2.0	2	2.0	2	2	2.0	2.0	2.0	2	2.0	2.0	2.0	2
	775	776	777	779	780	781	782	783	784	785	786	789	792	793
	3.5	2.5	3.5	4	3.5	3.5	4	5	4	4.5	2	5	3.5	5
rep	2.0	2.0	2.0	2	2.0	2.0	1	2	2	2.0	1	1	2.0	2
	794	796	798	799	804	806	808	809	810	811	812	813	815	817
	2.5	4	2	4.5	2	2	5.5	3	3.5	3.5	4.5	4.5	3.5	3
rep	2.0	2	2	2.0	2	2	2.0	2	2.0	2.0	2.0	2.0	2.0	2
	821	823	824	825	826	827	831	832	836	839	840	842	843	844
	3.5	5	3.5	2.5	5	4	4	4	4	3.5	3.5	2	3.5	4
rep	2.0	2	2.0	2.0	2	2	2	2	2	2.0	2.0	2	2.0	2
	845	846	848											
	2	4	1.5											
rep	2	2	2.0											

period	1	3
	3.053	3.87
rep	87.000	82.00

Treatment

```
Motrin Placebo
 3.822  3.073
rep 85.000 84.000
```

>

Is there any carry-over effect?

If there is a lasting effect of Motrin, the treatment difference in group A is smaller than in group B, in other words the average response in group A is larger than in group B.

```
> mod2=aov(max~order,data=tennis.max[tennis.max$period!=2,])
> summary(mod2)
```

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
order	1	0.8	0.7624	0.373	0.542
Residuals	167	341.1	2.0423		
			7 observations deleted due to missingness		

This not the case.

Another possibility: Are there any differences in the washout responses?

```
> mod3=aov(max~order,data=tennis.max[tennis.max$period==2,])
> summary(mod3)
```

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
order	1	1.78	1.775	0.973	0.327
Residuals	83	151.45	1.825		
			3 observations deleted due to missingness		

No.