

## Chapter 5 in Everitt and Hothorn (2010) Analysis of Variance

Start R.

If you were not able to edit Rprofile.site, load the HSAUR2 and Rcmdr either using the commands: `library(HSAUR2);library(Rcmdr)` or from the R Console using the menu Packages > Load package ... > select HSAUR2 and Rcmdr > Ok

We will be working with the R Commander menus.

### Weight Gain in Rats

From the R Commander menus select Data > Data in packages > Read data set from an attached package... > double click on HSAUR2, select weightgain, and click ok.

To see a description, from the R commander menu select Data > Active data set > Help on active data set (if available)

Click View data set to view it.

Create a new variable (tmt) combining both source and type by entering the following command in the Script Window and Submitting it.

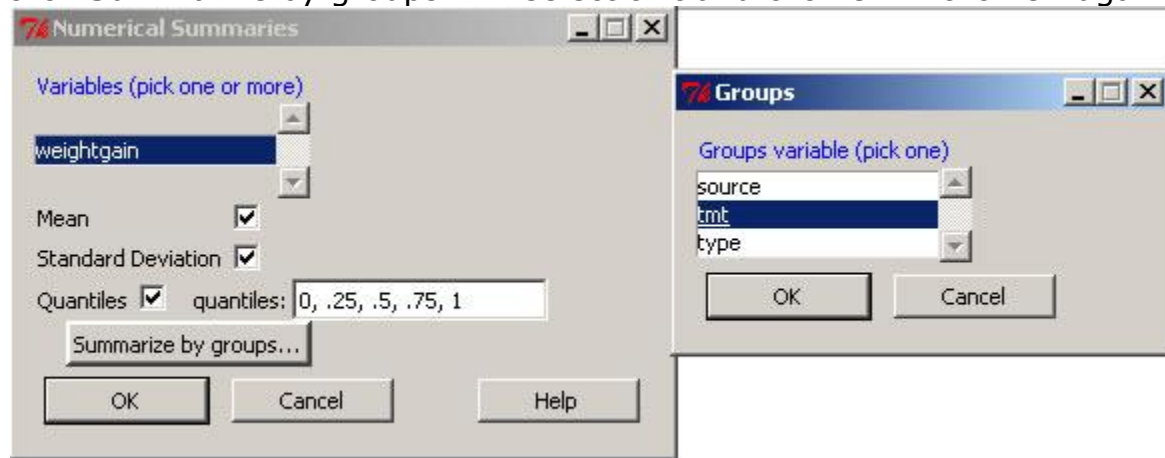
```
weightgain$tmt=factor(paste(as.character(weightgain$source) ,  
as.character(weightgain$type)))
```

You can also create a new variable from the menu Data > Manage variables in active data set > Compute new variable..., but it seems easier to just use a script command.

One of the quirks in R Commander is that you need to reset or refresh the active data set before it can recognize a new variable, so we will have to load another data set.

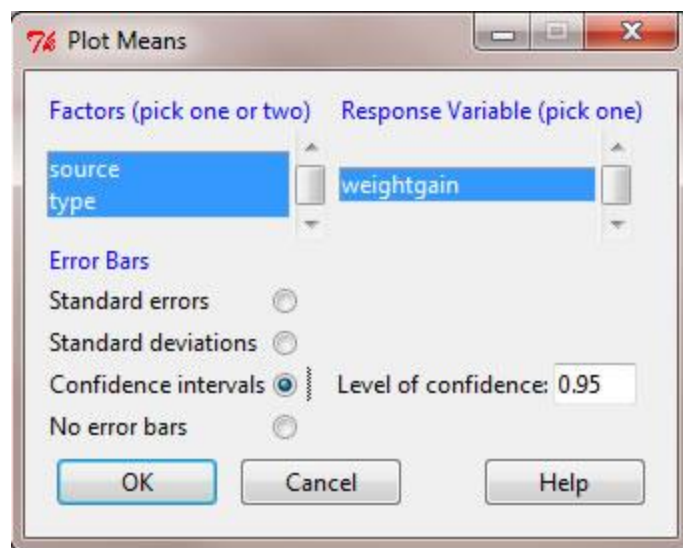
From the R Commander menus select Data > Active data set > Refresh data set.

From the menu, select Statistics > Summaries > Numerical summaries... > click Summarize by groups... > select tmt and click Ok > click Ok again.

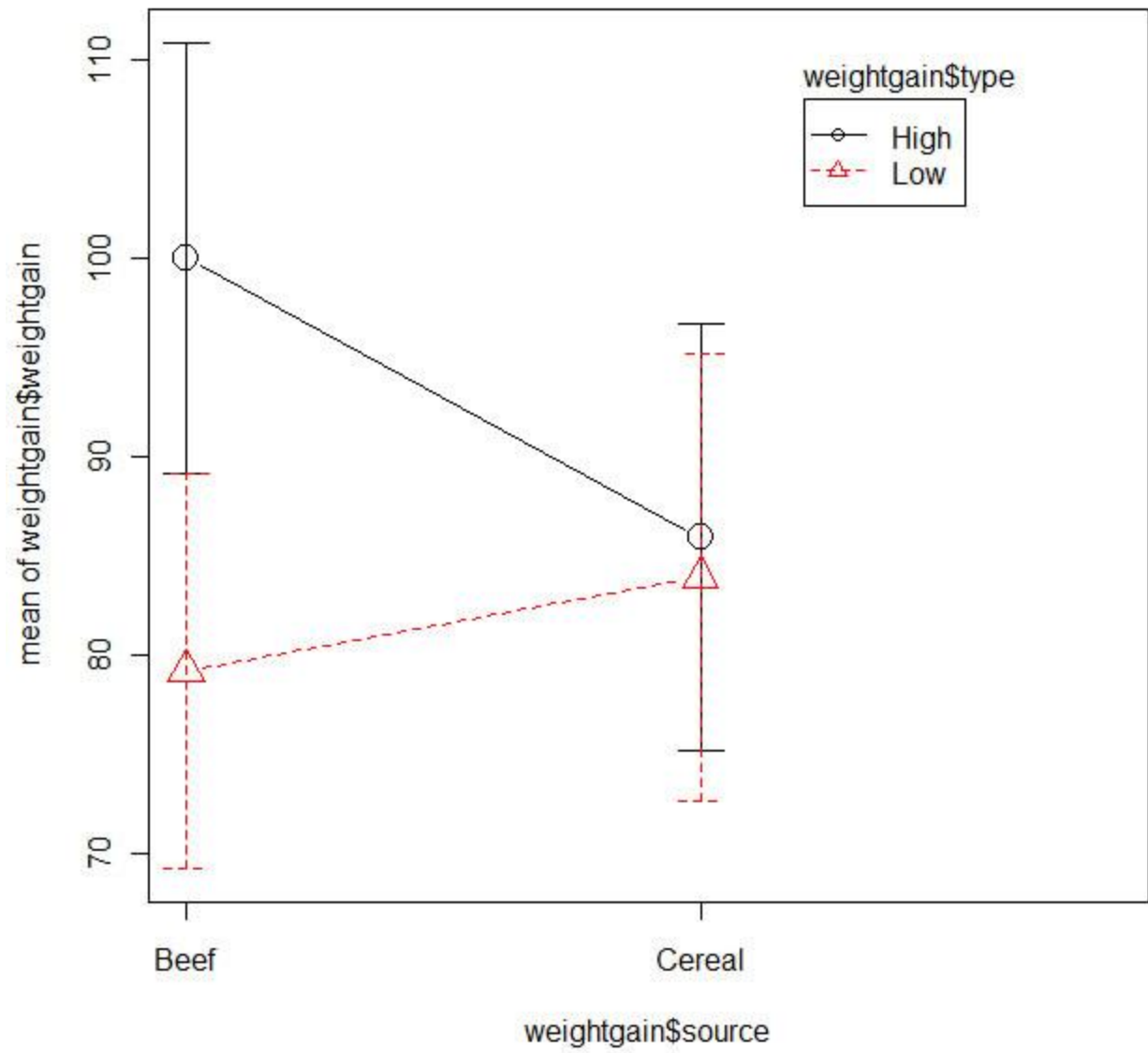


	mean	sd	0%	25%	50%	75%	100%	n
Beef High	100.0	15.13642	73	90.25	103.0	110.00	118	10
Beef Low	79.2	13.88684	51	73.00	82.0	90.00	95	10
Cereal High	85.9	15.02184	56	78.25	87.0	94.25	111	10
Cereal Low	83.9	15.70881	58	74.00	84.5	96.50	107	10

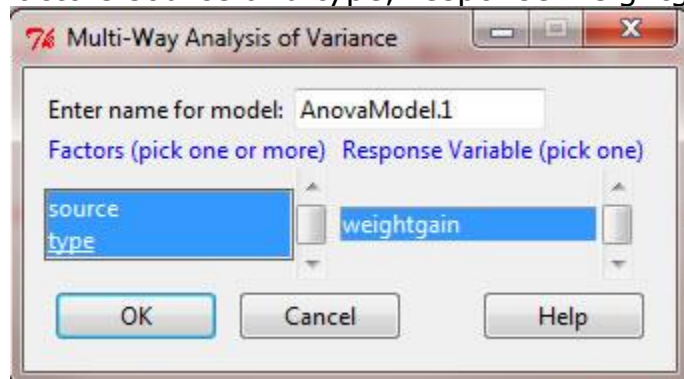
From the menu, select Graphs > Plot of means... > select factors: source and type, response: weightgain, confidence intervals and click Ok.



**Plot of Means**



From the menu, select Statistics > Means > Multi-way Anova... > select factors source and type, response weightgain and click ok.



```
< Anova (AnovaModel.1)
Anova Table (Type II tests)
```

```
Response: weightgain
```

	Sum Sq	Df	F value	Pr(>F)
source	220.9	1	0.9879	0.32688
type	1299.6	1	5.8123	0.02114 *
source:type	883.6	1	3.9518	0.05447 .
Residuals	8049.4	36		

```
---
```

```
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
< tapply(weightgain$weightgain, list(source=weightgain$source,
type=weightgain$type), mean, na.rm=TRUE) # means
```

	High	Low
source		
Beef	100.0	79.2
Cereal	85.9	83.9

```
< tapply(weightgain$weightgain, list(source=weightgain$source,
type=weightgain$type), sd, na.rm=TRUE) # std. deviations
```

	High	Low
source		
Beef	15.13642	13.88684
Cereal	15.02184	15.70881

```
< tapply(weightgain$weightgain, list(source=weightgain$source,
type=weightgain$type), function(x) sum(!is.na(x)))
```

```
+ # counts
```

	High	Low
source		
Beef	10	10
Cereal	10	10

From the menu, select Models > Summarize model.

Call:

```
lm(formula = weightgain ~ source * type, data = weightgain)
```

```
Residuals:
```

Min	1Q	Median	3Q	Max
-29.90	-9.90	2.05	10.85	25.10

Coefficients:

	Estimate	Std. Error	t value	Pr(> t )	
(Intercept)	100.000	4.729	21.148	< 2e-16	***
source[T.Cereal]	-14.100	6.687	-2.109	0.04201	*
type[T.Low]	-20.800	6.687	-3.110	0.00364	**
source[T.Cereal]:type[T.Low]	18.800	9.457	1.988	0.05447	.

---

Signif. codes: 0 '\*\*\*' 0.001 '\*\*' 0.01 '\*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 14.95 on 36 degrees of freedom

Multiple R-squared: 0.23, Adjusted R-squared: 0.1658

F-statistic: 3.584 on 3 and 36 DF, p-value: 0.02297

## Foster Feeding of Rats of Different Genotype

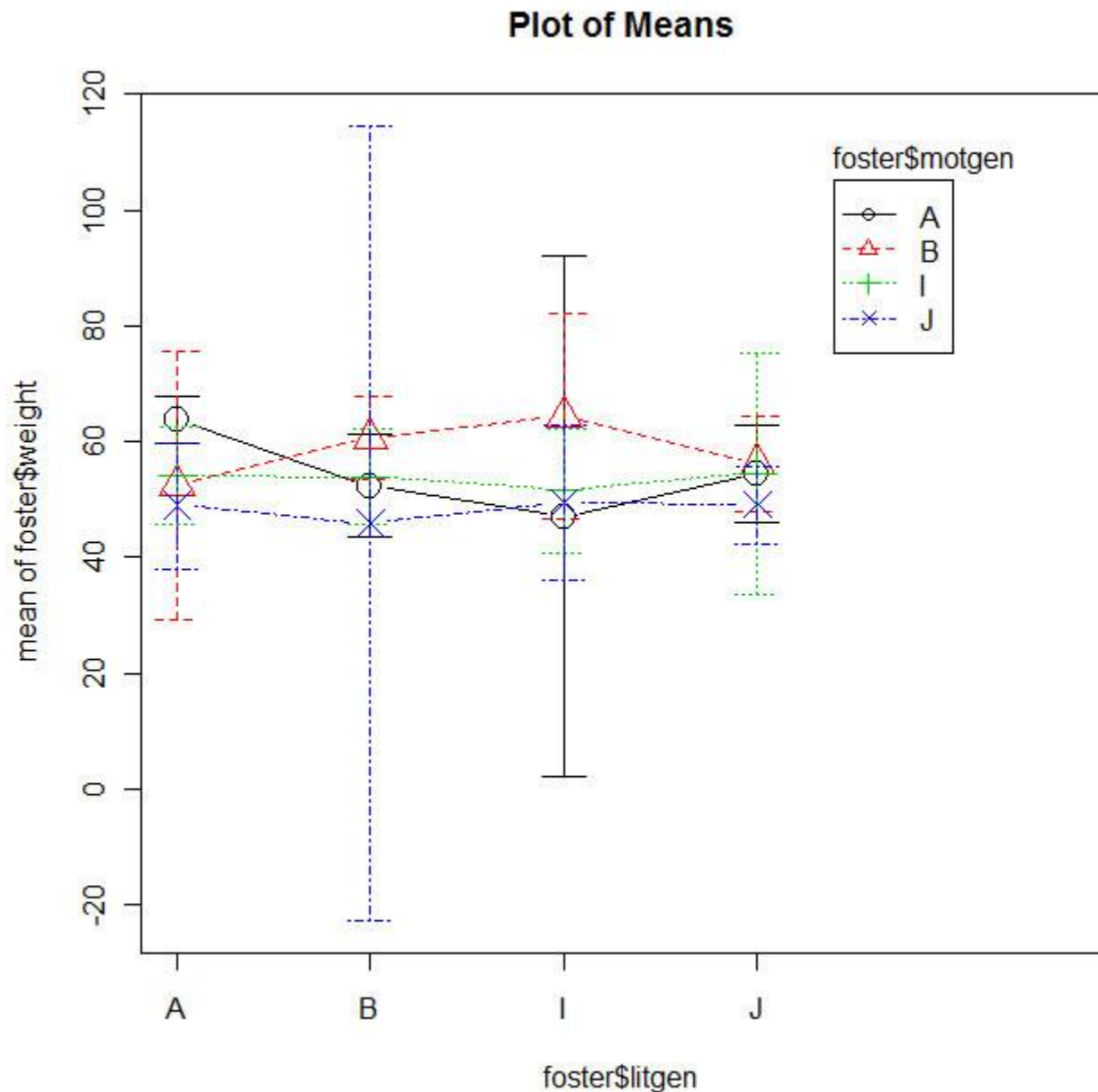
From the R Commander menus select Data > Data in packages > Read data set from an attached package... >

Double click on HSAUR2 and select foster, then click OK.

To see a description, from the R commander menu select Data > Active data set > Help on active data set (if available)

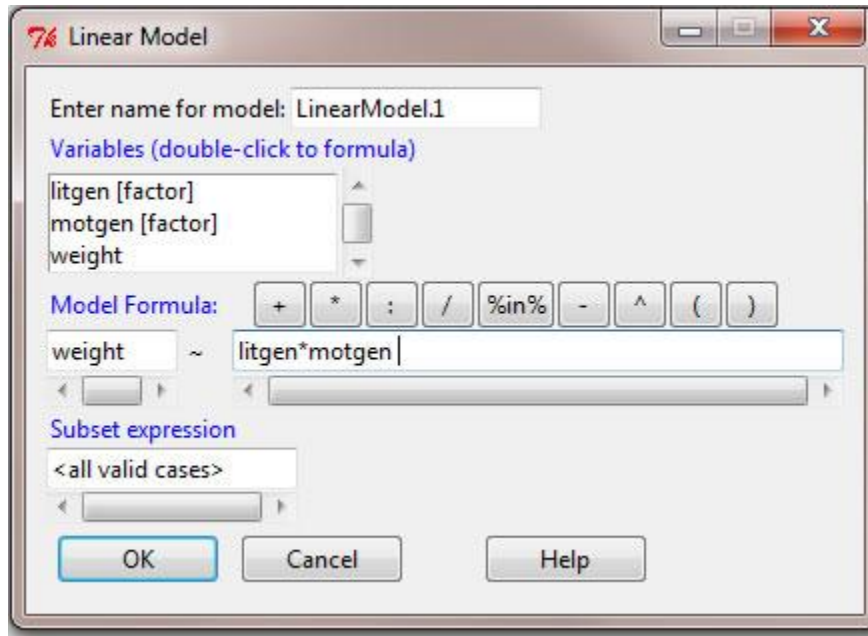
Click View data set to view it.

From the menu, select Graphs > Plot of means... > select factors: litgen and motgen, response: weight, confidence intervals and click Ok.



We could use the multi-way ANOVA menu as above, but a more flexible approach is:

From the menu, select Statistics > Fit models > Linear model... > click on weight to add it to the left side of the model equation, click on litgen, then \*, then motgen to add them to the right side of the equation, and click Ok.



Call:

```
lm(formula = weight ~ litgen * motgen, data = foster)
```

Residuals:

Min	1Q	Median	3Q	Max
-11.9000	-4.6250	0.5667	4.1000	20.9000

Coefficients:

	Estimate	Std. Error	t value	Pr(> t )	
(Intercept)	63.680	3.294	19.334	< 2e-16	***
litgen[T.B]	-11.355	4.940	-2.298	0.026241	*
litgen[T.I]	-16.580	5.378	-3.083	0.003498	**
litgen[T.J]	-9.330	4.940	-1.888	0.065417	.
motgen[T.B]	-11.280	5.378	-2.097	0.041622	*
motgen[T.I]	-9.555	4.940	-1.934	0.059413	.
motgen[T.J]	-14.720	4.658	-3.160	0.002818	**
litgen[T.B]:motgen[T.B]	19.595	7.303	2.683	0.010169	*
litgen[T.I]:motgen[T.B]	28.547	8.068	3.538	0.000948	***
litgen[T.J]:motgen[T.B]	13.030	7.783	1.674	0.101018	
litgen[T.B]:motgen[T.I]	11.155	7.178	1.554	0.127194	
litgen[T.I]:motgen[T.I]	14.055	7.303	1.925	0.060628	.
litgen[T.J]:motgen[T.I]	9.738	7.487	1.301	0.199956	
litgen[T.B]:motgen[T.J]	8.295	7.898	1.050	0.299196	
litgen[T.I]:motgen[T.J]	17.053	7.606	2.242	0.029937	*
litgen[T.J]:motgen[T.J]	9.430	6.790	1.389	0.171728	

---

Signif. codes: 0 '\*\*\*' 0.001 '\*\*' 0.01 '\*' 0.05 '.' 0.1 ' ' 1

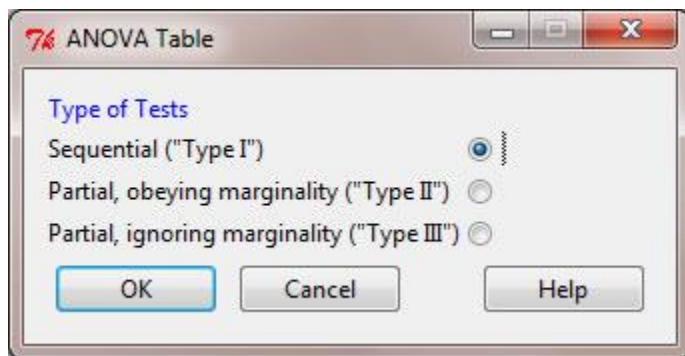
Residual standard error: 7.365 on 45 degrees of freedom

Multiple R-squared: 0.4047, Adjusted R-squared: 0.2063

F-statistic: 2.039 on 15 and 45 DF, p-value: 0.03333

From the menu, select Models > Hypothesis tests > Anova table > Type I





Repeat, selecting Type II and then Type III tests.  
Fit another model, changing the order, putting mortgen before litgen, and printout the three types of tests.

Analysis of Variance Table

Response: weight

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
litgen	3	60.16	20.052	0.3697	0.775221
motgen	3	775.08	258.360	4.7632	0.005736 **
litgen:motgen	9	824.07	91.564	1.6881	0.120053
Residuals	45	2440.82	54.240		

---

Signif. codes: 0 '\*\*\*' 0.001 '\*\*' 0.01 '\*' 0.05 '.' 0.1 ' ' 1

Anova Table (Type II tests)

Response: weight

	Sum Sq	Df	F value	Pr(>F)
litgen	63.63	3	0.3911	0.760004
motgen	775.08	3	4.7632	0.005736 **
litgen:motgen	824.07	9	1.6881	0.120053
Residuals	2440.82	45		

---

Signif. codes: 0 '\*\*\*' 0.001 '\*\*' 0.01 '\*' 0.05 '.' 0.1 ' ' 1

Anova Table (Type III tests)

Response: weight

	Sum Sq	Df	F value	Pr(>F)
(Intercept)	20275.7	1	373.8122	< 2e-16 ***
litgen	591.7	3	3.6362	0.01968 *
motgen	582.3	3	3.5782	0.02099 *
litgen:motgen	824.1	9	1.6881	0.12005
Residuals	2440.8	45		

---

Signif. codes: 0 '\*\*\*' 0.001 '\*\*' 0.01 '\*' 0.05 '.' 0.1 ' ' 1

Analysis of Variance Table

Response: weight

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
motgen	3	771.61	257.202	4.7419	0.005869 **
litgen	3	63.63	21.211	0.3911	0.760004

```

motgen:litgen  9  824.07  91.564  1.6881 0.120053
Residuals      45 2440.82  54.240
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Anova Table (Type II tests)
Response: weight
              Sum Sq Df F value    Pr(>F)
motgen       775.08  3   4.7632 0.005736 **
litgen        63.63  3   0.3911 0.760004
motgen:litgen  824.07  9   1.6881 0.120053
Residuals    2440.82 45
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Anova Table (Type III tests)
Response: weight
              Sum Sq Df  F value    Pr(>F)
(Intercept) 20275.7  1 373.8122 < 2e-16 ***
motgen       582.3   3   3.5782 0.02099 *
litgen       591.7   3   3.6362 0.01968 *
motgen:litgen  824.1  9   1.6881 0.12005
Residuals    2440.8 45
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

What is going on here?

All the treatment combinations do not have the same number of observations, so it is not possible to partition the variation into non-overlapping or orthogonal sums or squares.

Type I tests use sequential tests of litgen, motgen adjusted for litgen, and interaction.

Type II tests test each main effect adjusted for the other main effects such as litgen adjusted for motgen, motgen adjusted for litgen, and interaction. Note that these definitions of types of tests differs from the types used by SAS.

The order does not make any difference for type II and III tests.

The documentation says:

"The designations 'type-II' and 'type-III' are borrowed from SAS, but the definitions used here do not correspond precisely to those employed by SAS. Type-II tests are calculated according to the principle of marginality, testing each term after all others, except ignoring the term's higher-order relatives; so-called type-III tests violate marginality, testing each term in the model after all of the others. This definition of Type-II tests corresponds to the tests produced by SAS for analysis-of-variance models, where all of the predictors are factors, but not more generally (i.e., when there are quantitative predictors). Be very careful in formulating the model for type-III tests, or the hypotheses tested will not make sense." In almost all cases, I suggest type II tests.

Effect	Sum of Squares
litgen	60.16
litgen adjusted for morgen	63.63
morgen	771.61
morgen adjusted for litgen	775.08
interaction adjusted for litgen and morgen	824.07
intercept adjusted for everything else	20275.7
litgen adjusted for everything else	591.7
motgen adjusted for everything else	582.3
interaction adjusted for everything else	824.1

Multiple comparisons are only available using the aov(Analysis of Variance) command.

TukeyHSD will not work if RcmdrPlugin.HH is loaded.

Enter the following commands into the Script Window and Submit them.

```
m1=aov(weight ~ litgen*motgen, data=foster)
TukeyHSD(m1,"motgen")
```

Tukey multiple comparisons of means

95% family-wise confidence level

Fit: aov(formula = weight ~ litgen \* motgen, data = foster)

\$motgen

	diff	lwr	upr	p adj
B-A	3.330369	-3.859729	10.5204672	0.6078581
I-A	-1.895574	-8.841869	5.0507207	0.8853702
J-A	-6.566168	-13.627285	0.4949498	0.0767540
I-B	-5.225943	-12.416041	1.9641552	0.2266493
J-B	-9.896537	-17.197624	-2.5954489	0.0040509
J-I	-4.670593	-11.731711	2.3905240	0.3035490

## Water Hardness and Mortality

From the R Commander menus select Data > Data in packages > Read data set from an attached package... >

Double click on HSAUR2 and select water, then click OK.

To see a description, from the R commander menu select Data > Active data set > Help on active data set (if available)

Click View data set to view it.

Multivariate analysis of variance is not available from the menu, so enter the following commands in the Script Window and Submit them.

```
m1=manova(cbind(hardness,mortality) ~ location, data=water)
summary(m1, test="Hotelling-Lawley")
```

```
Df Hotelling-Lawley approx F num Df den Df      Pr(>F)
location      1          0.90021    26.106      2      58 8.217e-09 ***
Residuals 59
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

The test shows a significant difference in hardness, mortality or some combination of them.

From the menu, select Statistics > Summaries > Numerical summaries... > select variables hardness and mortality, click Summarize by groups > select location and click Ok > click Ok again.

```
Variable: hardness
      mean      sd 0%   25%   50%   75% 100%  n
North 30.40000 26.13449  6 12.50 17.0 44.00  94 35
South 69.76923 40.36068  5 40.25 75.5 99.75 138 26
```

```
Variable: mortality
      mean      sd   0%   25%   50%   75% 100%  n
North 1633.600 136.9369 1378 1557.50 1637 1718.00 1987 35
South 1376.808 140.2692 1096 1259.25 1364 1485.75 1627 26
```

## Male Egyptian Skulls

From the R Commander menus select Data > Data in packages > Read data set from an attached package... >

Double click on HSAUR2 and select skulls, then click OK.

To see a description, from the R commander menu select Data > Active data set > Help on active data set (if available)

Click View data set to view it.

From the menu, select Statistics > summaries > Numerical summaries... > select variables bh, bl, mb and nh, and click Summarize by groups > select epoch and click Ok > click Ok again.

```
Variable: bh
      mean      sd  0%   25%   50%   75% 100%  n
c4000BC 133.6000 4.469051 121 131.25 134.0 136.00 143 30
c3300BC 132.7000 4.647209 124 129.25 133.0 136.00 145 30
c1850BC 133.8000 4.978575 123 131.00 133.5 137.00 145 30
```

c200BC	132.3000	5.133729	120	130.00	132.0	135.75	142	30
cAD150	130.3333	4.971181	120	126.00	130.0	135.00	138	30

Variable: bl

	mean	sd	0%	25%	50%	75%	100%	n
c4000BC	99.16667	5.884423	89	95.00	100.0	102.75	114	30
c3300BC	99.06667	4.346488	90	97.00	98.5	101.75	107	30
c1850BC	96.03333	4.552251	87	92.25	96.0	99.75	106	30
c200BC	94.53333	4.591847	86	91.25	94.5	97.75	107	30
cAD150	93.50000	5.056576	81	91.00	94.0	97.00	103	30

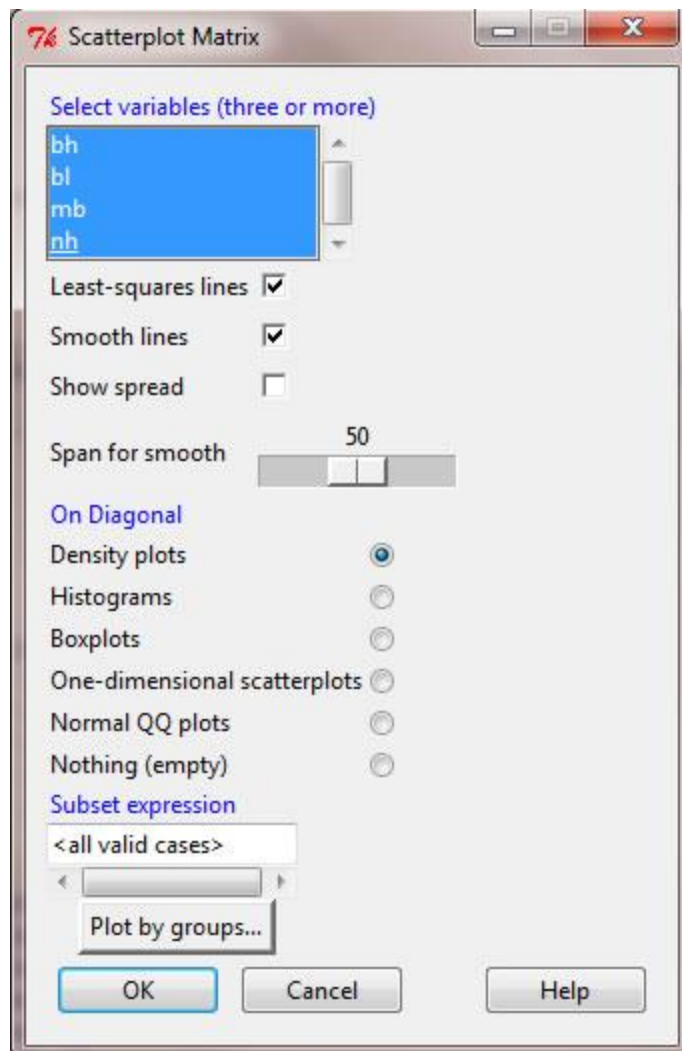
Variable: mb

	mean	sd	0%	25%	50%	75%	100%	n
c4000BC	131.3667	5.129249	119	128.00	131	134.75	141	30
c3300BC	132.3667	4.810071	123	130.00	132	134.75	148	30
c1850BC	134.4667	3.481313	126	132.25	136	137.00	140	30
c200BC	135.5000	3.919448	129	132.25	135	138.75	144	30
cAD150	136.1667	5.350368	126	132.25	137	139.00	147	30

Variable: nh

	mean	sd	0%	25%	50%	75%	100%	n
c4000BC	50.53333	2.763473	44	49.00	50.0	53.00	56	30
c3300BC	50.23333	2.955805	45	48.00	50.5	52.75	56	30
c1850BC	50.56667	3.549486	45	48.25	50.0	52.75	60	30
c200BC	51.96667	2.822121	46	50.25	52.0	53.75	60	30
cAD150	51.36667	3.718392	44	48.25	52.0	54.00	58	30

From the menu, select Graphs > Scatterplot matrix... > select variables bh, bl, mb and nh, and click Ok.



Multivariate analysis of variance is not available from the menu, so enter the following commands in the Script Window and Submit them.

```
m1=manova(cbind(mb,bh,bl,nh) ~ epoch, data=skulls)
summary(m1,test="Pillai")
summary(m1,test="Wilks")
summary(m1,test="Hotelling-Lawley")
summary(m1,test="Roy")
summary.aov(m1)
```

```
              Df  Pillai approx F num Df den Df      Pr(>F)
epoch          4 0.35331    3.512   16   580 4.675e-06 ***
Residuals 145
```

```
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
              Df  Wilks approx F num Df den Df      Pr(>F)
epoch          4 0.66359    3.9009   16 434.45 7.01e-07 ***
Residuals 145
---
```

```
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

      Df Hotelling-Lawley approx F num Df den Df      Pr(>F)
epoch      4          0.48182    4.231    16    562 8.278e-08 ***
Residuals 145
```

```
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
      Df      Roy approx F num Df den Df      Pr(>F)
epoch      4 0.4251    15.410      4    145 1.588e-10 ***
Residuals 145
```

```
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
Response mb :
      Df  Sum Sq Mean Sq F value    Pr(>F)
epoch      4   502.83  125.707   5.9546 0.0001826 ***
Residuals 145  3061.07   21.111
```

```
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
Response bh :
      Df Sum Sq Mean Sq F value    Pr(>F)
epoch      4   229.9   57.477   2.4474 0.04897 *
Residuals 145  3405.3   23.485
```

```
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
Response bl :
      Df Sum Sq Mean Sq F value    Pr(>F)
epoch      4   803.3  200.823   8.3057 4.636e-06 ***
Residuals 145  3506.0   24.179
```

```
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
Response nh :
      Df Sum Sq Mean Sq F value    Pr(>F)
epoch      4    61.2   15.300   1.507 0.2032
Residuals 145 1472.1   10.153
```

To look at pairwise multivariate tests, enter the following commands in the Script Window and Submit them.

```
summary(manova(cbind(mb,bh,bl,nh) ~ epoch, data=skulls, subset=
epoch %in% c("c4000BC", "c3300BC")))
```

```
summary(manova(cbind(mb,bh,bl,nh) ~ epoch, data=skulls, subset=
epoch %in% c("c4000BC", "c1850BC")))
```

```
summary(manova(cbind(mb,bh,bl,nh) ~ epoch, data=skulls, subset=
epoch %in% c("c4000BC", "c200BC")))
```

```
summary(manova(cbind(mb,bh,bl,nh) ~ epoch, data=skulls, subset=
epoch %in% c("c4000BC", "cAD150")))
```

```
> summary(manova(cbind(mb,bh,bl,nh) ~ epoch, data=skulls, subset= epoch %in%
c("c4000BC", "c3300BC")))
      Df  Pillai approx F num Df den Df Pr(>F)
```

```
epoch      1 0.027674  0.39135      4      55  0.814
Residuals 58
```

```
> summary(manova(cbind(mb,bh,bl,nh) ~ epoch, data=skulls, subset= epoch %in%
c("c4000BC", "c1850BC")))
```

```
      Df  Pillai approx F num Df den Df  Pr(>F)
epoch    1 0.18757   3.1744     4    55 0.02035 *
Residuals 58
```

```
---
```

```
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
> summary(manova(cbind(mb,bh,bl,nh) ~ epoch, data=skulls, subset= epoch %in%
c("c4000BC", "c200BC")))
```

```
      Df  Pillai approx F num Df den Df  Pr(>F)
epoch    1 0.30297   5.9766     4    55 0.0004564 ***
Residuals 58
```

```
---
```

```
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
> summary(manova(cbind(mb,bh,bl,nh) ~ epoch, data=skulls, subset= epoch %in%
c("c4000BC", "cAD150")))
```

```
      Df  Pillai approx F num Df den Df  Pr(>F)
epoch    1 0.36182   7.7956     4    55 4.736e-05 ***
Residuals 58
```

```
---
```

```
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
> summary(manova(cbind(mb,bh,bl,nh) ~ epoch, data=skulls, subset= epoch %in%
c("c4000BC", "c3300BC")))
```

```
      Df  Pillai approx F num Df den Df Pr(>F)
epoch    1 0.027674  0.39135     4    55  0.814
Residuals 58
```