

THE DESIGN AND MIXED-MODEL ANALYSIS OF EXPERIMENTS

PRACTICAL III SOLUTIONS

III.1 Prove lemma III.2

We have that $\mathbf{P} = \mathbf{X}(\mathbf{X}'\mathbf{X})^{-1}\mathbf{X}'$ so that $\mathbf{PX} = \mathbf{X}(\mathbf{X}'\mathbf{X})^{-1}\mathbf{X}'\mathbf{X} = \mathbf{X}$.

Also, $\mathbf{RX} = (\mathbf{I} - \mathbf{P})\mathbf{X} = \mathbf{X} - \mathbf{PX} = \mathbf{0}$.

III.2 Show that

$$\mathbf{P}_T = \mathbf{X}_T(\mathbf{X}_T'\mathbf{X}_T)^{-1}\mathbf{X}_T'$$

$$= \begin{bmatrix} \mathbf{1}_{n_1}(\mathbf{1}'_{n_1}\mathbf{1}_{n_1})^{-1}\mathbf{1}'_{n_1} & \mathbf{0}_{n_1 \times n_2} & \cdots & \mathbf{0}_{n_1 \times n_t} \\ \mathbf{0}_{n_2 \times n_1} & \mathbf{1}_{n_2}(\mathbf{1}'_{n_2}\mathbf{1}_{n_2})^{-1}\mathbf{1}'_{n_2} & \cdots & \mathbf{0}_{n_2 \times n_t} \\ \vdots & \vdots & \ddots & \vdots \\ \mathbf{0}_{n_t \times n_1} & \mathbf{0}_{n_t \times n_2} & \cdots & \mathbf{1}_{n_t}(\mathbf{1}'_{n_t}\mathbf{1}_{n_t})^{-1}\mathbf{1}'_{n_t} \end{bmatrix}$$

where

$$\mathbf{X}_T = \begin{bmatrix} \mathbf{1}_{n_1} & \mathbf{0}_{n_1 \times 1} & \cdots & \mathbf{0}_{n_1 \times 1} \\ \mathbf{0}_{n_2 \times 1} & \mathbf{1}_{n_2} & \cdots & \mathbf{0}_{n_2 \times 1} \\ \vdots & \vdots & \ddots & \vdots \\ \mathbf{0}_{n_t \times 1} & \mathbf{0}_{n_t \times 1} & \cdots & \mathbf{1}_{n_t} \end{bmatrix}$$

First

$$\mathbf{X}_T'\mathbf{X}_T = \begin{bmatrix} \mathbf{1}'_{n_1} & \mathbf{0}_{1 \times n_2} & \cdots & \mathbf{0}_{1 \times n_t} \\ \mathbf{0}_{1 \times n_1} & \mathbf{1}'_{n_2} & \cdots & \mathbf{0}_{1 \times n_t} \\ \vdots & \vdots & \ddots & \vdots \\ \mathbf{0}_{1 \times n_1} & \mathbf{0}_{1 \times n_2} & \cdots & \mathbf{1}'_{n_t} \end{bmatrix} \begin{bmatrix} \mathbf{1}_{n_1} & \mathbf{0}_{n_1 \times 1} & \cdots & \mathbf{0}_{n_1 \times 1} \\ \mathbf{0}_{n_2 \times 1} & \mathbf{1}_{n_2} & \cdots & \mathbf{0}_{n_2 \times 1} \\ \vdots & \vdots & \ddots & \vdots \\ \mathbf{0}_{n_t \times 1} & \mathbf{0}_{n_t \times 1} & \cdots & \mathbf{1}_{n_t} \end{bmatrix} = \begin{bmatrix} \mathbf{1}'_{n_1}\mathbf{1}_{n_1} & 0 & \cdots & 0 \\ 0 & \mathbf{1}'_{n_2}\mathbf{1}_{n_2} & \cdots & 0 \\ \vdots & \vdots & \ddots & \vdots \\ 0 & 0 & \cdots & \mathbf{1}'_{n_t}\mathbf{1}_{n_t} \end{bmatrix}$$

Since $\mathbf{1}'_{n_k}\mathbf{1}_{n_k}$ is a scalar $(\mathbf{1}'_{n_k}\mathbf{1}_{n_k})^{-1}$ exists and

$$\begin{aligned}
\mathbf{P}_T &= \mathbf{X}_T (\mathbf{X}_T' \mathbf{X}_T)^{-1} \mathbf{X}_T' \\
&= \begin{bmatrix} \mathbf{1}_{n_1} & \mathbf{0}_{n_1 \times 1} & \cdots & \mathbf{0}_{n_1 \times 1} \\ \mathbf{0}_{n_2 \times 1} & \mathbf{1}_{n_2} & \cdots & \mathbf{0}_{n_2 \times 1} \\ \vdots & \vdots & \ddots & \vdots \\ \mathbf{0}_{n_t \times 1} & \mathbf{0}_{n_t \times 1} & \cdots & \mathbf{1}_{n_t} \end{bmatrix} \begin{bmatrix} (\mathbf{1}_{n_1}' \mathbf{1}_{n_1})^{-1} & 0 & \cdots & 0 \\ 0 & (\mathbf{1}_{n_2}' \mathbf{1}_{n_2})^{-1} & \cdots & 0 \\ \vdots & \vdots & \ddots & \vdots \\ 0 & 0 & \cdots & (\mathbf{1}_{n_t}' \mathbf{1}_{n_t})^{-1} \end{bmatrix} \begin{bmatrix} \mathbf{1}_{n_1}' & \mathbf{0}_{1 \times n_2} & \cdots & \mathbf{0}_{1 \times n_t} \\ \mathbf{0}_{1 \times n_1} & \mathbf{1}_{n_2}' & \cdots & \mathbf{0}_{1 \times n_t} \\ \vdots & \vdots & \ddots & \vdots \\ \mathbf{0}_{1 \times n_1} & \mathbf{0}_{1 \times n_2} & \cdots & \mathbf{1}_{n_t}' \end{bmatrix} \\
&= \begin{bmatrix} \mathbf{1}_{n_1} (\mathbf{1}_{n_1}' \mathbf{1}_{n_1})^{-1} & \mathbf{0}_{n_1 \times 1} & \cdots & \mathbf{0}_{n_1 \times 1} \\ \mathbf{0}_{n_2 \times 1} & \mathbf{1}_{n_2} (\mathbf{1}_{n_2}' \mathbf{1}_{n_2})^{-1} & \cdots & \mathbf{0}_{n_2 \times 1} \\ \vdots & \vdots & \ddots & \vdots \\ \mathbf{0}_{n_t \times 1} & \mathbf{0}_{n_t \times 1} & \cdots & \mathbf{1}_{n_t} (\mathbf{1}_{n_t}' \mathbf{1}_{n_t})^{-1} \end{bmatrix} \begin{bmatrix} \mathbf{1}_{n_1}' & \mathbf{0}_{1 \times n_2} & \cdots & \mathbf{0}_{1 \times n_t} \\ \mathbf{0}_{1 \times n_1} & \mathbf{1}_{n_2}' & \cdots & \mathbf{0}_{1 \times n_t} \\ \vdots & \vdots & \ddots & \vdots \\ \mathbf{0}_{1 \times n_1} & \mathbf{0}_{1 \times n_2} & \cdots & \mathbf{1}_{n_t}' \end{bmatrix} \\
&= \begin{bmatrix} \mathbf{1}_{n_1} (\mathbf{1}_{n_1}' \mathbf{1}_{n_1})^{-1} \mathbf{1}_{n_1}' & \mathbf{0}_{n_1 \times n_2} & \cdots & \mathbf{0}_{n_1 \times n_t} \\ \mathbf{0}_{n_2 \times n_1} & \mathbf{1}_{n_2} (\mathbf{1}_{n_2}' \mathbf{1}_{n_2})^{-1} \mathbf{1}_{n_2}' & \cdots & \mathbf{0}_{n_2 \times n_t} \\ \vdots & \vdots & \ddots & \vdots \\ \mathbf{0}_{n_t \times n_1} & \mathbf{0}_{n_t \times n_2} & \cdots & \mathbf{1}_{n_t} (\mathbf{1}_{n_t}' \mathbf{1}_{n_t})^{-1} \mathbf{1}_{n_t}' \end{bmatrix}
\end{aligned}$$

III.3 Show that $\mathbf{P}_T \mathbf{P}_G \mathbf{X}_T \boldsymbol{\alpha} = \mathbf{1}_n \bar{\alpha}$, where $\bar{\alpha} = \left(\sum_{k=1}^t n_k \alpha_k \right) / n$ and $\mathbf{P}_T \mathbf{X}_T \boldsymbol{\alpha} = \mathbf{X}_T \boldsymbol{\alpha}$

First since $\mathbf{P}_T \mathbf{P}_G = \mathbf{P}_G$, $\mathbf{P}_T \mathbf{P}_G \mathbf{X}_T \boldsymbol{\alpha} = \mathbf{P}_G \mathbf{X}_T \boldsymbol{\alpha}$ and

$$\mathbf{P}_G \mathbf{X}_T \boldsymbol{\alpha} = \mathbf{P}_G \begin{bmatrix} \alpha_1 \mathbf{1}_{n_1} \\ \alpha_2 \mathbf{1}_{n_2} \\ \vdots \\ \alpha_t \mathbf{1}_{n_t} \end{bmatrix} = \frac{1}{n} (n_1 \alpha_1 + n_2 \alpha_2 + \dots + n_t \alpha_t) \mathbf{1}_n = \frac{\sum_{i=1}^t n_i \alpha_i}{n} \mathbf{1}_n = \bar{\alpha} \mathbf{1}_n.$$

III.4 Verify that $\mathbf{e}_G - \mathbf{t}_e = \mathbf{R}_T \mathbf{R}_G \mathbf{y}$.

Now $\mathbf{e}_G = \mathbf{R}_G \mathbf{y}$ and $\mathbf{t}_e = \mathbf{P}_T \mathbf{R}_G \mathbf{y}$

so that $\mathbf{e}_G - \mathbf{t}_e = \mathbf{R}_G \mathbf{y} - \mathbf{P}_T \mathbf{R}_G \mathbf{y} = (\mathbf{I} - \mathbf{P}_T) \mathbf{R}_G \mathbf{y} = \mathbf{R}_T \mathbf{R}_G \mathbf{y}$.

III.5 Let $E[\mathbf{Y}] = \mathbf{X}_G\mu$, $\mathbf{V}_Y = \sigma^2\mathbf{I}_n$, $R(\alpha|\mu) = \mathbf{Y}'\mathbf{P}_T\mathbf{R}_G\mathbf{Y}$ and $D(\alpha) = \mathbf{Y}'\mathbf{R}_T\mathbf{Y}$ where \mathbf{R}_G and \mathbf{R}_T are as defined in lemma III.1. Then, show that

$$E[R(\alpha|\mu)/(t-1)] = \sigma^2 \quad \text{and} \quad E[D(\alpha)/(n-t)] = \sigma^2$$

where t is the number of treatments and n is the number of observations.

For $E[R(\alpha|\mu)/(t-1)]$, we first use theorem II.11 to show that

$$\begin{aligned} E[R(\alpha|\mu)/(t-1)] &= E[\mathbf{Y}'\mathbf{P}_T\mathbf{R}_G\mathbf{Y}]/(t-1) \\ &= \left\{ \text{trace}(\mathbf{P}_T\mathbf{R}_G\sigma^2\mathbf{I}_n) + (\mathbf{X}_G\mu)' \mathbf{P}_T\mathbf{R}_G(\mathbf{X}_G\mu) \right\} / \{t-1\} \\ &= \left\{ \sigma^2 \text{trace}(\mathbf{P}_T\mathbf{R}_G) + (\mathbf{X}_G\mu)' \mathbf{P}_T\mathbf{R}_G(\mathbf{X}_G\mu) \right\} / \{t-1\} \end{aligned}$$

Now from theorem III.5, $\text{trace}(\mathbf{P}_T\mathbf{R}_G) = t-1$.

Also, by lemma III.2, $\mathbf{R}_G\mathbf{X}_G = \mathbf{0}$ and so $\mathbf{P}_T\mathbf{R}_G(\mathbf{X}_G\mu) = \mathbf{0}$.

Hence,

$$\begin{aligned} E[R(\alpha|\mu)/(t-1)] &= \left\{ \sigma^2 \text{trace}(\mathbf{P}_T\mathbf{R}_G) + (\mathbf{X}_G\mu)' \mathbf{P}_T\mathbf{R}_G(\mathbf{X}_G\mu) \right\} / \{t-1\} \\ &= \left\{ \sigma^2(t-1) + 0 \right\} / \{t-1\} \\ &= \sigma^2 \end{aligned}$$

For $E[D(\alpha)/(n-t)]$,

$$E[D(\alpha)/(n-t)] = E[\mathbf{y}'\mathbf{R}_T\mathbf{y}]/(n-t) = \left\{ \sigma^2 \text{trace}(\mathbf{R}_T) + (\mathbf{X}_G\mu)' \mathbf{R}_T(\mathbf{X}_G\mu) \right\} / \{n-t\}$$

Now from theorem III.5, $\text{trace}(\mathbf{R}_T) = n-t$.

From lemma III.1 $\mathbf{R}_T = \mathbf{R}_T\mathbf{R}_G$ and from lemma III.2 $\mathbf{R}_G\mathbf{X}_G = \mathbf{0}$ so that $\mathbf{R}_T(\mathbf{X}_G\mu) = \mathbf{R}_T\mathbf{R}_G\mathbf{X}_G\mu = \mathbf{0}$. Consequently,

$$\begin{aligned} E[D(\alpha)/(n-t)] &= \left\{ \sigma^2 \text{trace}(\mathbf{R}_T) + (\mathbf{X}_G\mu)' \mathbf{R}_T(\mathbf{X}_G\mu) \right\} / \{n-t\} \\ &= \left\{ \sigma^2(n-t) + 0 \right\} / \{n-t\} \\ &= \sigma^2 \end{aligned}$$

III.6 An investigation was conducted to examine differences between 3 brands of tyre in their braking distances (ft.) from a speed of 30 miles per hour. Altogether 9 tests were conducted with the particular brand tested at each test being chosen at random. The results are as follows:

Brand		
A	B	C
28	27	27
26	25	32
30	26	31

Perform the one-way analysis manually using mean operators (**P**).

Test	Brand	Stopping Distance y	Grand mean g	Total Test Deviations e_G	Brand effects t_e	Residual Test Deviations e_T
1	1	28	28	0	0	0
2	1	26	28	-2	0	-2
3	1	30	28	2	0	2
4	2	27	28	-1	-2	1
5	2	25	28	-3	-2	-1
6	2	26	28	-2	-2	0
7	3	27	28	-1	2	-3
8	3	32	28	4	2	2
9	3	31	28	3	2	1
Sums of squares of deviations				48	24	24
Degrees of freedom				8	2	6
Variance				6	12	4

Step 1: Set up hypotheses

$$H_0: \alpha_1 = \alpha_2 = \alpha_3 = \mu$$

H_1 : at least one pair of population brand means is different

Step 2: Calculate test statistic

The analysis of variance table for the example is:

Source	df	SSq	MSq	F
Tests	8	48		
Brands	2	24	12	3.0
Residual	6	24	4	

Step 3: Decide between hypotheses

The probability of exceeding an F of 3.0 with $v_1 = 2$ and $v_2 = 6$ is $P(F \geq 3.0) = 0.1250$. Little evidence of a difference between the brands.

- III.7** Use Genstat to produce a randomized layout for an experiment involving 21 plots to which 7 treatments are to be allocated so that each is replicated 3 times. Use the seed 413634 in producing the layout.

The layout is given at the end of the following Genstat output.

Genstat 5 Release 4.1 (PC/Windows NT) 22 March 2000 16:27:47
Copyright 1998, Lawes Agricultural Trust (Rothamsted Experimental Station)

Genstat 5 Fourth Edition - (for Windows)
Genstat 5 Procedure Library Release PL11

```
3 DELETE [redefine=yes] Plots,Treat
4 FACTOR [modify=yes;nvalues=21;levels=21] Plots
5 READ Plots; frepresentation=ordinal
```

Identifier	Values	Missing	Levels
Plots	21	0	21

```
7 FACTOR [modify=yes;nvalues=21;levels=7] Treat
8 READ Treat; frepresentation=ordinal
```

Identifier	Values	Missing	Levels
Treat	21	0	7

```
10
11 PRINT Plots,Treat
```

Plots	Treat
1	1
2	1
3	1
4	2
5	2
6	2
7	3
8	3
9	3
10	4
11	4
12	4
13	5
14	5
15	5
16	6
17	6
18	6
19	7
20	7
21	7

```
12 RANDOMIZE [BLOCKSTRUCTURE=Plots; SEED=413634] Treat
13 PDESIGN [BLOCK=Plots; TREAT=Treat]
```

*** Treatment combinations on each unit of the design ***

```
Plots
  1  6
  2  1
  3  3
  4  6
  5  7
  6  1
  7  3
  8  7
  9  2
 10  7
 11  4
 12  5
 13  1
 14  2
 15  3
 16  5
 17  6
 18  4
 19  2
 20  4
 21  5
```

Treatment factors are listed in the order: Treat

III.8 in an experiment to compare melon varieties, six plots of each of four varieties were grown. The varieties were allocated to the 24 plots in the experiment in a completely random manner. The melon yields for each plot are as follows:

Variety			
A	B	C	D
25.12	40.25	18.30	28.05
17.25	35.25	22.60	28.55
26.42	31.98	25.90	33.20
16.08	36.52	15.05	31.68
22.15	43.32	11.42	30.32
15.92	37.10	23.68	27.58

This data is available in *CRDMelon.gsh* in the directory *G:\Disciplina\Genstat*. Add the 24-level factor Plots and the factor Variety with levels A–D to this Genstat spreadsheet and make sure that the factors and variate are available in the central store.

Conduct a regression analysis to determine if there is any measurable effect of variety on the yield.

What are the treatment means?

Use the Genstat BLOCKSTRUCTURE, TREATMENTSTRUCTURE and ANOVA commands to perform an analysis of variance on this data and confirm your answers from the regression analysis.

Use Genstat to obtain a residuals-versus-fitted-values plot and a normal probability plot as a check on the assumptions underlying the analysis.

Also, carry out the LSD and Tukey's procedures to determine exactly which Varieties differed.

38 PRINT Plots,Variety,Yield

Plots	Variety	Yield
1	A	25.12
2	A	17.25
3	A	26.42
4	A	16.08
5	A	22.15
6	A	15.92
7	B	40.25
8	B	35.25
9	B	31.98
10	B	36.52
11	B	43.32
12	B	37.10
13	C	18.30
14	C	22.60
15	C	25.90
16	C	15.05
17	C	11.42
18	C	23.68
19	D	28.05
20	D	28.55
21	D	33.20
22	D	31.68
23	D	30.32
24	D	27.58

39 MODEL Yield

40 TERMS Variety

41 FIT [FPROB=y] Variety

41.....

***** Regression Analysis *****

Response variate: Yield

Fitted terms: Constant, Variety

*** Summary of analysis ***

	d.f.	s.s.	m.s.	v.r.	F pr.
Regression	3	1291.5	430.49	23.42	<.001
Residual	20	367.7	18.38		
Total	23	1659.1	72.14		

Percentage variance accounted for 74.5

Standard error of observations is estimated to be 4.29

* MESSAGE: The following units have large standardized residuals:

Unit	Response	Residual
17	11.42	-2.06

* MESSAGE: The error variance does not appear to be constant:
intermediate responses are less variable than small or large responses

*** Estimates of parameters ***

	estimate	s.e.	t(20)
Constant	20.49	1.75	11.71
Variety B	16.91	2.48	6.83
Variety C	-1.00	2.48	-0.40
Variety D	9.41	2.48	3.80

Step 1: Set up hypotheses

$$H_0: \alpha_A = \alpha_B = \alpha_C = \alpha_D = \mu$$

H_1 : at least one pair of population Variety means is different

Step 2: Calculate test statistic

The analysis of variance table for the example is:

Source	df	SSq	MSq	F	p
Plots	23	1291.5			
Treatments	3	367.7	430.49	23.49	<0.001
Residual	20	1659.1	72.14		

Step 3: Decide between hypotheses

The probability of exceeding an F of 23.49 with $v_1 = 3$ and $v_2 = 20$ is $P(F \geq 23.42) < 0.001$. The evidence suggests that there is a Variety difference.

The treatment means are 20.49, $20.49 + 16.91 = 37.40$, $20.49 - 1.00 = 19.49$ and $20.49 + 9.41 = 29.90$.

```
42 BLOCK Plots
43 TREAT Variety
44 ANOVA [FPROB=Y; PSE=LSD] Yield
```

```
44.....
```

```
***** Analysis of variance *****
```

```
Variate: Yield
```

Source of variation	d.f.	s.s.	m.s.	v.r.	F pr.
Plots stratum					
Variety	3	1291.48	430.49	23.42	<.001
Residual	20	367.65	18.38		
Total	23	1659.13			

```
* MESSAGE: the following units have large residuals.
```

```
Plots 17          -8.07    s.e. 3.91
```

```
***** Tables of means *****
```

```
Variate: Yield
```

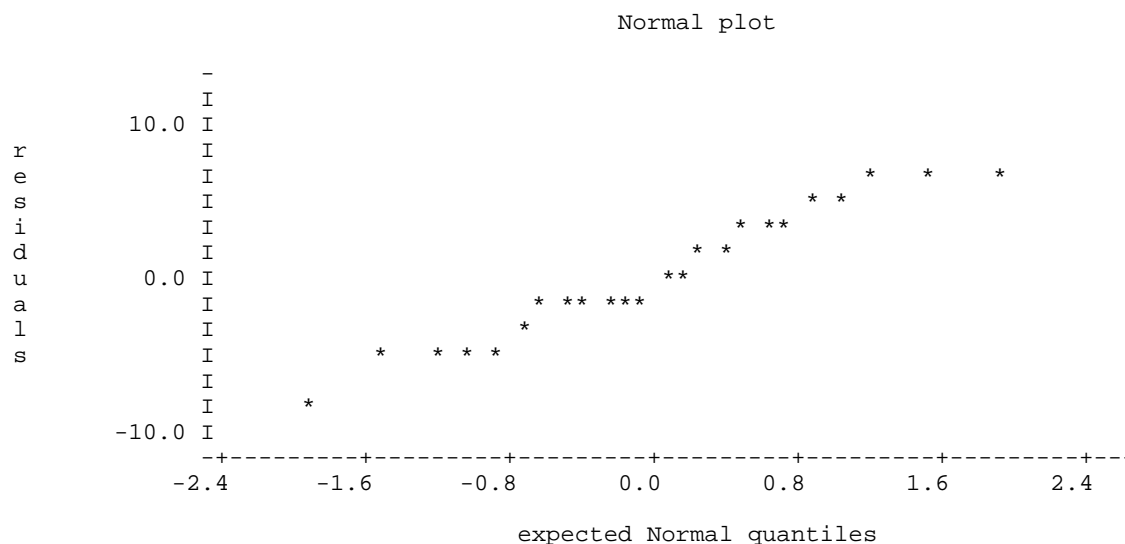
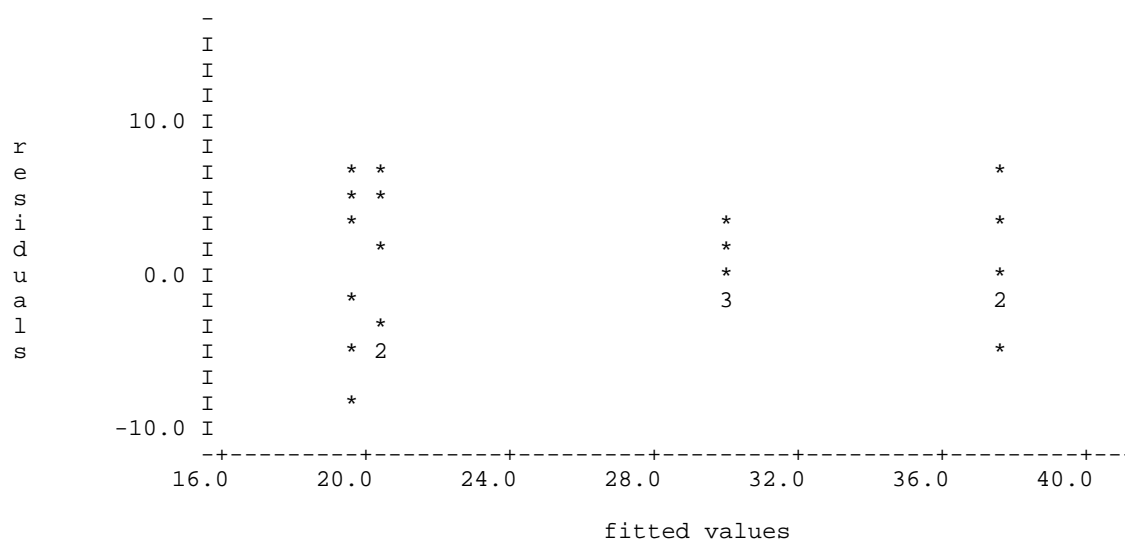
```
Grand mean  26.82
```

Variety	A	B	C	D
	20.49	37.40	19.49	29.90

*** Least significant differences of means (5% level) ***

Table	Variety
rep.	6
d.f.	20
l.s.d.	5.164

45 APLOT METHOD=fit,normal



The analysis of variance and means from the mean operator analysis are as obtained from the regression analysis.

It appears that the homogeneity of variance assumption may not be met as the size of the band for the second highest-yielding variety has less variance than the other varieties. It would be useful to use a hypothesis test to confirm this conclusion but we will not cover these in this subject. From the normal probability plot it appears that the normality assumption is met except for some outliers (an extremely small

and an extremely large residual). As we cannot check the observations in question we will do nothing further about them.

Finally, we examine the difference between the pairs of means using the LSD and Tukey's procedure. Strictly, speaking this is not appropriate as the assumptions have not been met. We do so as an exercise realizing that we cannot rely on the conclusions made.

Differences between all pairs of Variety means

Variety		C	A	D	B
	Mean	19.49	20.49	29.90	37.40
C	19.49				
A	20.49	1.00			
D	29.90	10.41	9.41		
B	37.40	17.91	16.91	7.50	
	LSD(5%)		5.164		

Variety			
C	A	D	B
19.49	20.49	29.90	37.40

Varieties C and A yielded lower than D which was in turn lower than B.

Differences between all pairs of Variety means

Variety		C	A	D	B
	Mean	19.49	20.49	29.90	37.40
C	19.49				
A	20.49	1.00			
D	29.90	10.41	9.41		
B	37.40	17.91	16.91	7.50	
	w(5%)		6.93		

$$\begin{aligned}
 w(5\%) &= \frac{3.960}{\sqrt{2}} \sqrt{18.38 \frac{2}{6}} \\
 &= 2.6879 \times 1.392 \\
 &= 6.93
 \end{aligned}$$

Variety			
C	A	D	B
19.49	20.49	29.90	37.40

Varieties C and A yielded lower than D which was in turn lower than B.

III.9 In an experiment to investigate the effect of different rates of injection on the lethal dose of ouabain, animals were injected at one of four different rates with ouabain and the lethal dose recorded. The data given in the following table:

	1	5	9	11	13	14	16	17	20	22	28	31	31
Rate of	2	3	6	22	27	27	28	28	37	40	42	50	
Injection	4	34	34	38	40	46	58	60	60	65			
(mg/kg/min)	8	51	56	62	63	70	73	76	89	92			
/1045.75													

Input the data into Genstat. Note that the replication of the rates is not equal. To set up a factor for Rates you will either have to manually enter the values of the levels or use *Stats > Design > Generate Factors in Standard Order or Spread > Column > Fill* to generate an equal number for each rate, the number generated being between the minimum and maximum replication. Then extras can be deleted and/or deficiencies rectified by adding extra cells.

Use Genstat to perform a one-way analysis of variance on this data in which a quadratic is fitted to determine if there is a relationship between the lethal dose and the rate of injection. Perform the analysis using the averaging operator method. Produce residuals-versus-fitted-values and normal probability plots to check the assumptions underlying the analysis. Obtain the equation of the fitted line and a plot of the Rate means.

Genstat 5 Release 4.1 (PC/Windows NT) 19 March 2000 22:52:34
Copyright 1998, Lawes Agricultural Trust (Rothamsted Experimental Station)

Genstat 5 Fourth Edition - (for Windows)
Genstat 5 Procedure Library Release PL11

```
3 "Data taken from File: D:/ANALYSES/LM/ONEFAC/CRDCATS.GSH"
4 DELETE [redefine=yes] Animals,Rate,Dose
5 FACTOR [modify=yes;nvalues=41;levels=41] Animals
6 READ Animals; frepresentation=ordinal
```

Identifier	Values	Missing	Levels
Animals	41	0	41

```
9 FACTOR [modify=yes;nvalues=41;levels=(1,2,4,8)] Rate
10 READ Rate; frepresentation=ordinal
```

Identifier	Values	Missing	Levels
Rate	41	0	4

```
13 VARIATE [nvalues=41] Dose
14 READ Dose
```

Identifier	Minimum	Mean	Maximum	Values	Missing
Dose	3.00	38.88	92.00	41	0

```
17
18 PRINT Animals,Rate,Dose
```

Animals	Rate	Dose
1	1.000	5.00
2	1.000	9.00
3	1.000	11.00
4	1.000	13.00

5	1.000	14.00
6	1.000	16.00
7	1.000	17.00
8	1.000	20.00
9	1.000	22.00
10	1.000	28.00
11	1.000	31.00
12	1.000	31.00
13	2.000	3.00
14	2.000	6.00
15	2.000	22.00
16	2.000	27.00
17	2.000	27.00
18	2.000	28.00
19	2.000	28.00
20	2.000	37.00
21	2.000	40.00
22	2.000	42.00
23	2.000	50.00
24	4.000	34.00
25	4.000	34.00
26	4.000	38.00
27	4.000	40.00
28	4.000	46.00
29	4.000	58.00
30	4.000	60.00
31	4.000	60.00
32	4.000	65.00
33	8.000	51.00
34	8.000	56.00
35	8.000	62.00
36	8.000	63.00
37	8.000	70.00
38	8.000	73.00
39	8.000	76.00
40	8.000	89.00
41	8.000	92.00

19 BLOCK Animals
 20 TREAT POL(Rate; 2)
 21 ANOVA [FPROB=Y] Dose

21.....

***** Analysis of variance *****

Variate: Dose

Source of variation	d.f.	s.s.	m.s.	v.r.	F pr.
Animals stratum					
Rate	3	16094.3	5364.8	35.13	<.001
Lin	1	15700.1	15700.1	102.81	<.001
Quad	1	382.6	382.6	2.51	0.122
Deviations	1	11.6	11.6	0.08	0.785
Residual	37	5650.1	152.7		
Total	40	21744.4			

***** Tables of means *****

Variate: Dose

Grand mean 38.9

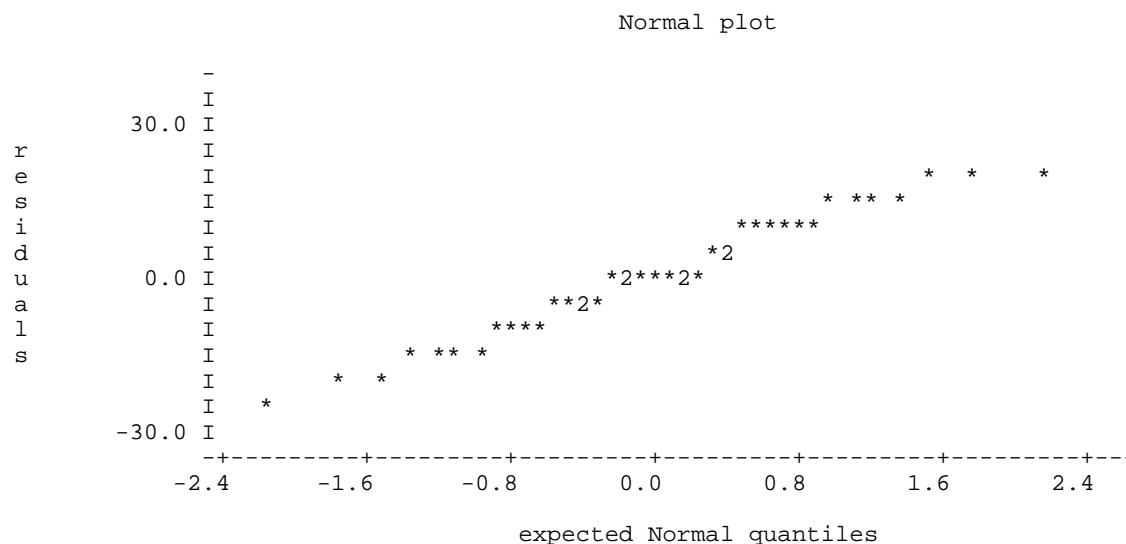
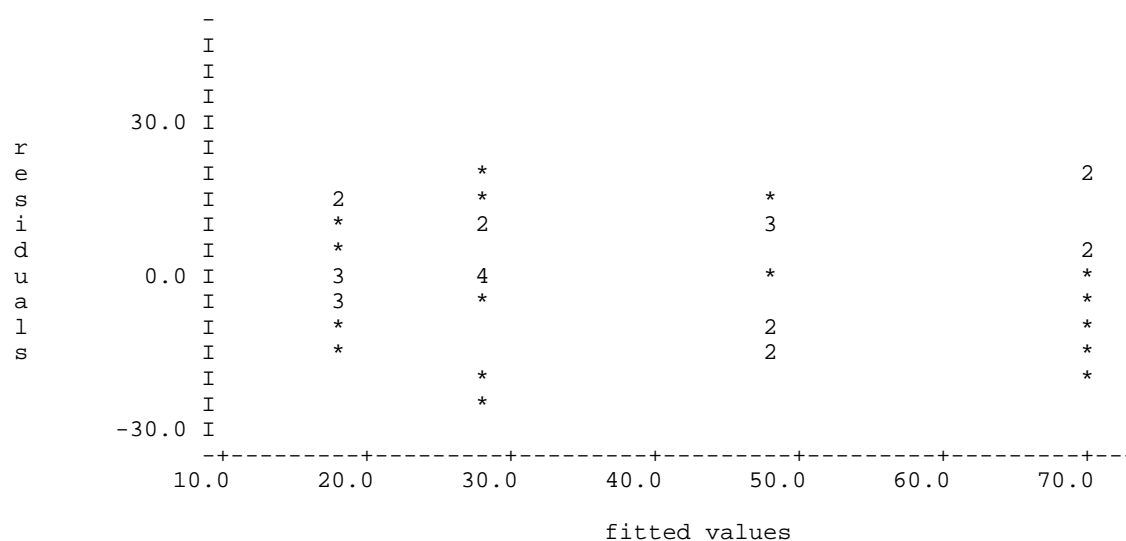
Rate	1.00	2.00	4.00	8.00
	18.1	28.2	48.3	70.2
rep.	12	11	9	9

*** Standard errors of differences of means ***

Table	Rate	
rep.	unequal	
d.f.	37	
s.e.d.	5.83	min.rep
	5.45	max-min
	5.04X	max.rep

(No comparisons in categories where s.e.d. marked with an X)

```
22  AKEEP [FIT=Fit; RES=Res]
23  APLOTT METHOD=fit,normal
```



Step 1: Set up hypotheses

- a) $H_0: \gamma_1 = 0$
 $H_1: \gamma_1 \neq 0$
- b) $H_0: \gamma_2 = 0$
 $H_1: \gamma_2 \neq 0$
- c) $H_0: \alpha_k - \mu - \gamma_1 x_k - \gamma_2 x_k^2 = 0$ for all k
 $H_1: \alpha_k - \mu - \gamma_1 x_k - \gamma_2 x_k^2 \neq 0$ for all k

Step 2: Calculate test statistics

Source	df	SSQ	MSQ	F	Prob
Animals	40	21744.4			
Rates	3	16094.3	5364.8	35.13	<0.001
Linear	1	15700.1	15700.1	102.81	<0.001
Quadratic	1	382.6	382.6	2.51	0.122
Deviations	1	11.6	11.6	0.08	0.785
Residual	37	5650.1	152.7		

Step 3: Decide between hypotheses

The Deviations and Quadratic terms are not significant. However, the Linear term is highly significant.

Note that the residual-versus-fitted-values plot looks satisfactory so that the homogeneity of variance assumption appears to be met. Also, the normal probability plot appears to be basically linear and so the normality assumption also appears to be met.

We now fit a linear equation and obtain its formula.

```

24  "
-25  **** Fit linear only ****
-26  "
27  TREAT POL(Rate; 1)
28  ANOVA [PRINT=aov] Dose

28.....

**** Analysis of variance ****

Variate: Dose

Source of variation    d.f.      s.s.      m.s.      v.r.

Animals stratum
Rate                   3       16094.3    5364.8    35.13
  Lin                   1       15700.1   15700.1   102.81
  Deviations            2         394.2    197.1     1.29
Residual               37       5650.1    152.7
Total                  40       21744.4

```

```

29  APOLYNOMIAL Rate; COEFF=CoefFs

***** Equation of the polynomial *****

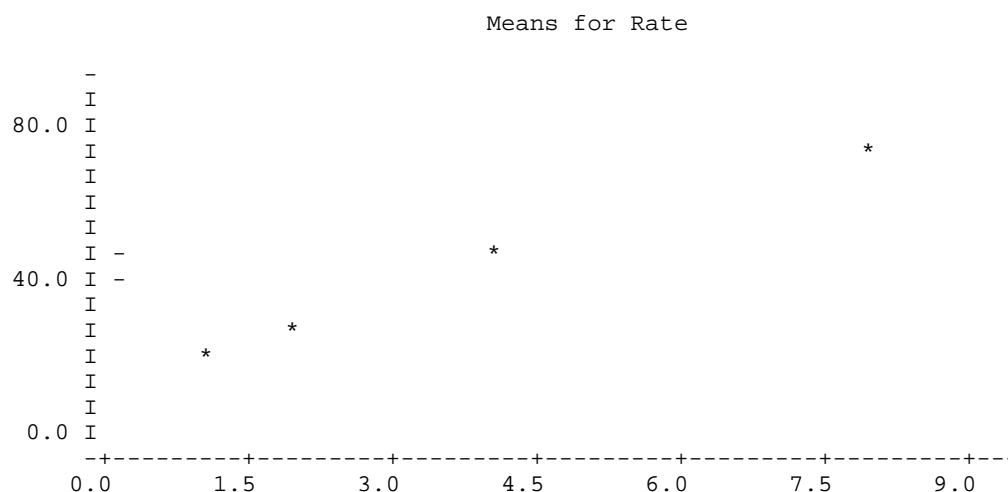
13.14 + 7.43  * Rate

30  PRINT #CoefFs

      13.14      7.433

31  AGRAPH [GRAPH=line] XFACTOR=Rate; BAR=*

```



Examination of the output reveals that the fitted equation is

$$y_i = 13.14 + 7.433 x_k.$$

III.10 An experiment was conducted to compare the pain relief afforded by three drugs. Altogether there were 39 patients that were randomly assigned one of three drug treatments to receive so that 13 patients received each drug. Each patient was given the drug assigned to them when they declared that their level of pain required relief. The number of hours relief afforded by the first administration of the drug was recorded and is given in the following table.

	D	2	6	4	13	5	8	4	6	7	6	8	12	4
Drug	T1	2	0	3	3	0	0	8	1	4	2	2	1	3
	T2	6	4	4	0	1	8	2	8	12	1	5	2	4

This data is contained in *CRDDrug.gsh* in the directory *G:\Disciplina\Genstat*. Add the 39-level factor Plots and the factor Drug with levels D, T1 and T2 to this Genstat spreadsheet and make sure that the factors and variate are available in the central store.

The drug treatment D was a form of control so it is of interest to compare this treatment with the other two. This could be achieved by using the following orthogonal contrasts:

D	Drug	
	T1	T2
2	-1	-1
0	1	-1

Use Genstat to perform a one-way analysis of variance on this data, including the fitting the orthogonal contrasts and checking the assumptions.

Genstat 5 Release 4.1 (PC/Windows NT) 20 March 2000 08:14:46
Copyright 1998, Lawes Agricultural Trust (Rothamsted Experimental Station)

Genstat 5 Fourth Edition - (for Windows)
Genstat 5 Procedure Library Release PL11

```
3 "Data taken from File: D:/ANALYSES/LM/ONEFAC/CRDDRUGALL.GSH"
4 DELETE [redefine=yes] Patients,Drug,Relief
5 FACTOR [modify=yes;nvalues=39;levels=39] Patients
6 READ Patients; frepresentation=ordinal
```

Identifier	Values	Missing	Levels
Patients	39	0	39

```
9 FACTOR [modify=yes;nvalues=39;levels=3;labels=!t('D','T1','T2')] Drug
10 READ Drug; frepresentation=ordinal
```

Identifier	Values	Missing	Levels
Drug	39	0	3

```
13 VARIATE [nvalues=39] Relief
14 READ Relief
```

Identifier	Minimum	Mean	Maximum	Values	Missing
Relief	0.000	4.385	13.000	39	0

```
17
18 PRINT Patients,Drug,Relief
```

Patients	Drug	Relief
1	D	2.000
2	D	6.000
3	D	4.000
4	D	13.000
5	D	5.000
6	D	8.000
7	D	4.000
8	D	6.000
9	D	7.000
10	D	6.000
11	D	8.000
12	D	12.000
13	D	4.000
14	T1	2.000
15	T1	0.000
16	T1	3.000
17	T1	3.000
18	T1	0.000
19	T1	0.000
20	T1	8.000
21	T1	1.000
22	T1	4.000
23	T1	2.000
24	T1	2.000
25	T1	1.000
26	T1	3.000


```

27          T2          6.000
28          T2          4.000
29          T2          4.000
30          T2          0.000
31          T2          1.000
32          T2          8.000
33          T2          2.000
34          T2          8.000
35          T2         12.000
36          T2          1.000
37          T2          5.000
38          T2          2.000
39          T2          4.000

19  "
-20  ****Orthogonal Contrasts ****
-21  "
22  TEXT ConDrug; !T('D vs rest','T1 vs T2')
23  MATRIX [ROW=ConDrug; COL=3] Contrasts; \
24          !(2,2(-1), 0,1,-1)
25  BLOCK Patients
26  TREAT REG (Drug; 2; Contrasts)
27  ANOVA [FPROB=Y; PRINT=AOV,MEANS,CONTRAST; PSE=LSD] Relief

27.....

**** Analysis of variance ****

Variate: Relief

Source of variation      d.f.        s.s.        m.s.        v.r.    F pr.

Patients stratum
Drug                     2        120.615        60.308        6.86    0.003
  D vs rest              1         90.462        90.462       10.29    0.003
  T1 vs T2               1         30.154        30.154        3.43    0.072
Residual                 36        316.615         8.795
Total                    38        437.231

**** Tables of contrasts ****

Variate: Relief

**** Patients stratum ****

*** Drug contrasts ***

D vs rest      1.08    s.e. 0.336    ss.div. 78.0
T1 vs T2      -1.08    s.e. 0.582    ss.div. 26.0

**** Tables of means ****

Variate: Relief

Grand mean  4.38

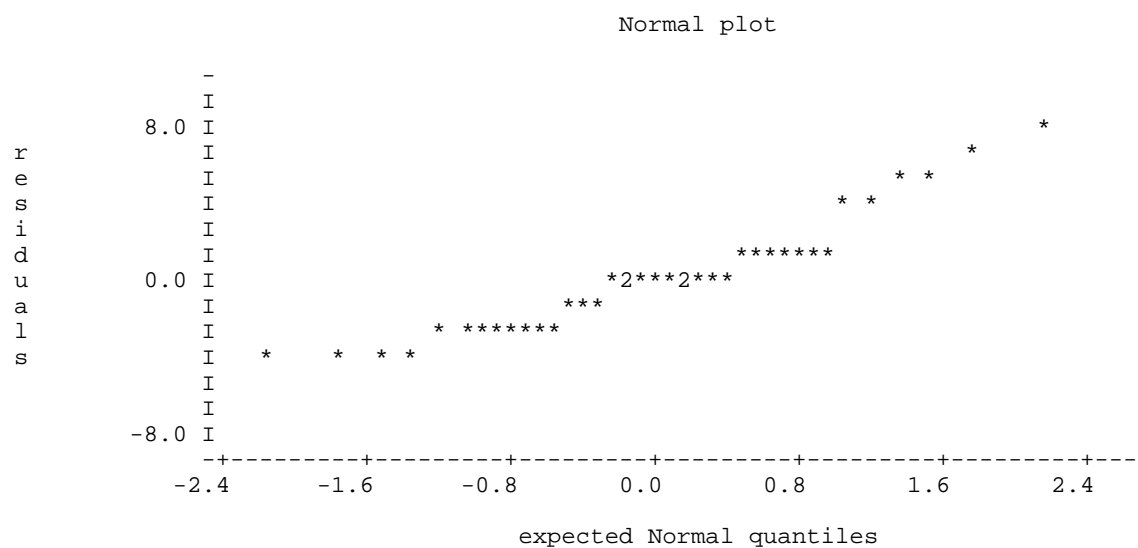
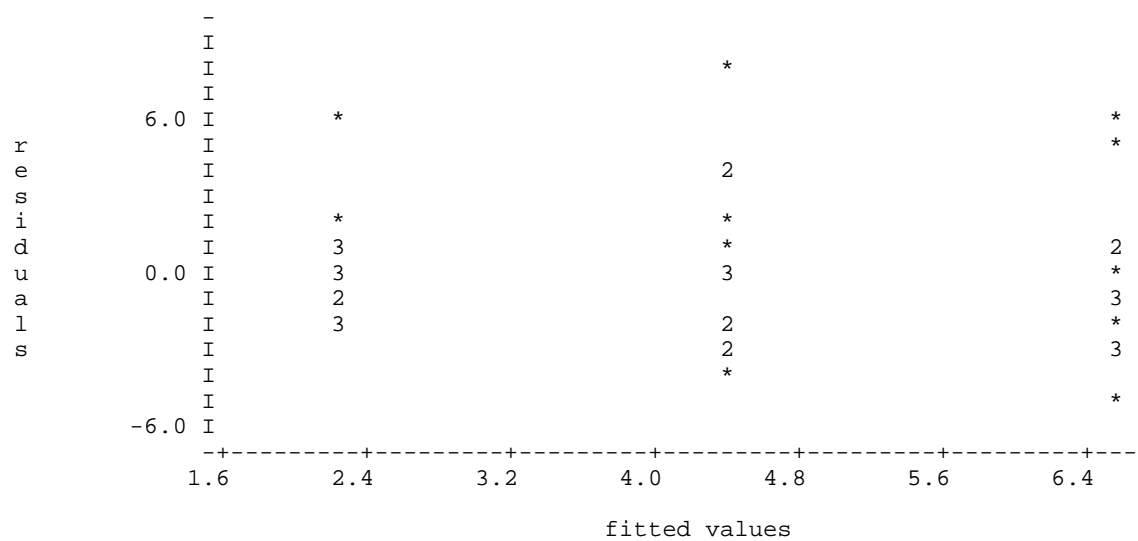
      Drug      D      T1      T2
        6.54      2.23      4.38

*** Least significant differences of means (5% level) ***

Table      Drug
rep.        13
d.f.        36
l.s.d.      2.359

```

```
28 AKEEP [FIT=Fit; RES=Res]
29 APLOT METHOD=fit,normal
```



Step 1: Set up hypotheses

For each contrast:

$H_0: \mathbf{x}'\boldsymbol{\alpha} = 0$ where \mathbf{x} is a vector of contrast coefficients

$H_1: \mathbf{x}'\boldsymbol{\alpha} \neq 0$

Step 2: Calculate test statistics

Source	df	SSQ	MSQ	F	Prob
Patients	38	437.231			
Drug	2	120.615	60.308	6.86	0.003
D vs rest	1	90.462	90.462	10.29	0.003
T1 vs T2	1	30.154	30.154	3.43	0.072
Residual	36	316.615	8.795		

Step 3: Decide between hypotheses

The relief provided by T1 and T2 is less than that provided by D; there is not a significant difference between T1 and T2.

Note that the residual-versus-fitted-values plot looks satisfactory so that the homogeneity of variance assumption appears to be met. However, the normal probability plot has some evidence of curvature in it and so it seems that the normality assumption may not be met. As ANOVA is robust to departures from normality we will continue with the analysis.

The means for the different drugs are given in the following table.

	Drug		
	D	T1	T2
Mean	6.54	2.23	4.38

Note that the estimate for the first orthogonal contrast is 1.08 and that

$$6.54 - \frac{2.23 + 4.38}{2} = 3.235 \text{ and } \frac{3.235}{3} = 1.08.$$

So the difference between D and the mean of T1 and T2 is 3.24.