

Course 02402 Introduction to Statistics

Lecture 11: One-way Analysis of Variance, ANOVA

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Analysis of Variance

"ANalysis Of VAriance" (ANOVA) was introduced by R.A. Fisher 100 years ago as a systematic way of analysing groups and has since been a key part of statistics and its applications.

- Today: one factor (one-way ANOVA)
- Next week: two factors (two-way ANOVA)
- First factor is typically called *treatment*, second factor *block*.

Overview

- 1 Intro
- 2 Model and hypothesis
- 3 Computation - decomposition and the ANOVA table
- 4 Hypothesis test (F-test)
- 5 Within-group variability and relation to the 2-sample t-test
- 6 Post hoc analysis
- 7 Model control / model validation
- 8 A complete example - from the book

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One-way ANOVA - simple example

Group A	Group B	Group C
2.8	5.5	5.8
3.6	6.3	8.3
3.4	6.1	6.9
2.3	5.7	6.1

Is there a difference (in means) between the groups A, B and C?

Analysis of variance (ANOVA) can be used for the analysis, if the observations in each group can be assumed to be normally distributed.

One-way ANOVA - simple example in R

```
# Input data
y <- c(2.8, 3.6, 3.4, 2.3,
      5.5, 6.3, 6.1, 5.7,
      5.8, 8.3, 6.9, 6.1)

## Define treatment groups
treatm <- factor(c(1, 1, 1, 1,
                  2, 2, 2, 2,
                  3, 3, 3, 3))

## Plot data by treatment groups
par(mfrow = c(1,2))
plot(y ~ as.numeric(treatm), xlab = "Treatment", ylab = "y")
boxplot(y ~ treatm, xlab = "Treatment", ylab = "y")
```

Eksempel: Cow dung and antibiotics

Decomposition of cow dung: how much organic material is left?

Control	α -cypermethrin	Ivermectin	Spiramycin
2.43	3.00	3.03	2.80
2.63	3.02	2.81	2.85
2.56	2.87	3.06	2.84
2.76	2.96	3.11	2.93
2.70	2.77	2.94	
2.54	2.75	3.06	

Cow dung and antibiotics – eksempel i R

```
dung <- c(2.43, 2.63, 2.56, 2.76, 2.70, 2.54, 3.00, 3.02, 2.87, 2.96,
         2.77, 2.75, 3.03, 2.81, 3.06, 3.11, 2.94, 3.06, 2.80, 2.85,
         2.84, 2.93)
treat <- factor(c(rep("control", 6), rep("a-cyperm", 6),
                  rep("iverm", 6), rep("spiram", 4)))

plot(dung ~ as.numeric(treat), xlab = "Treatment", ylab = "y")
boxplot(dung ~ treat, xlab = "Treatment", ylab = "y")

model <- lm(dung ~ treat)
anova(model)
```

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One-way ANOVA, model

- The model may be formulated as

$$Y_{ij} = \mu + \alpha_i + \varepsilon_{ij},$$

where the ε_{ij} are assumed to be independent and identically distributed (i.i.d.) with

$$\varepsilon_{ij} \sim N(0, \sigma^2).$$

- μ : overall mean.
- α_i : effect of group (treatment) i .
- Y_{ij} : j th measurement in group i (j runs from 1 to n_i).

One-way ANOVA, hypothesis

- We want to compare the (more than 2) means $\mu + \alpha_i$ in the model

$$Y_{ij} = \mu + \alpha_i + \varepsilon_{ij}, \quad \varepsilon_{ij} \stackrel{\text{i.i.d.}}{\sim} N(0, \sigma^2).$$

- The hypothesis may be formulated as

$$H_0: \alpha_i = 0 \quad \text{for all } i$$

with alternative hypothesis

$$H_1: \alpha_i \neq 0 \quad \text{for at least one } i$$

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One-way ANOVA, decomposition and the ANOVA table

- With the model

$$Y_{ij} = \mu + \alpha_i + \varepsilon_{ij}, \quad \varepsilon_{ij} \stackrel{\text{i.i.d.}}{\sim} N(0, \sigma^2)$$

the total variation in the data can be decomposed:

$$SST = SS(Tr) + SSE.$$

- 'One-way' refers to the fact that there is only one factor in the experiment on k levels.
- The method is called analysis of variance, because the testing is carried out by comparing certain variances.

Formulas for sums of squares

- Total sum of squares ("the total variance")

$$SST = \sum_{i=1}^k \sum_{j=1}^{n_i} (y_{ij} - \bar{y})^2$$

- The sum of squares for the residuals ("residual variance after model fit")

$$SSE = \sum_{i=1}^k \sum_{j=1}^{n_i} (y_{ij} - \bar{y}_i)^2$$

- Sum of squares of treatment ("variance explained by the model")

$$SS(Tr) = \sum_{i=1}^k n_i (\bar{y}_i - \bar{y})^2$$

The ANOVA table

Source of variation	Deg. of freedom	Sums of squares	Mean sum of squares
Treatment	$k - 1$	$SS(Tr)$	$MS(Tr) = \frac{SS(Tr)}{k-1}$
Residual	$n - k$	SSE	$MSE = \frac{SSE}{n-k}$
Total	$n - 1$	SST	

```
# One-way ANOVA using anova() and lm()
anova(lm(y ~ treatm))

## Analysis of Variance Table
##
## Response: y
##          Df Sum Sq Mean Sq F value Pr(>F)
## treatm    2   30.8   15.40   26.7 0.00017 ***
## Residuals  9    5.2    0.58
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

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One-way ANOVA, F-test

- We have: (Theorem ??)

$$SST = SS(Tr) + SSE$$

- and we can find the test statistic

$$F = \frac{SS(Tr)/(k-1)}{SSE/(n-k)} = \frac{MS(Tr)}{MSE}$$

where

- k is the number of levels of the factor,
 - n is the total number of observations.
- Choose the significance level α , and compute the test statistic F .
 - Compare the test statistic to the relevant quantile of the F -distribution:

$$F \sim F_{\alpha}(k-1, n-k) \text{ (Theorem ??)}$$

The F -distribution and the F -test

```
# Remember, this is "under H0" (i.e. we compute as if H0 is true)

# Number of groups
k <- 3

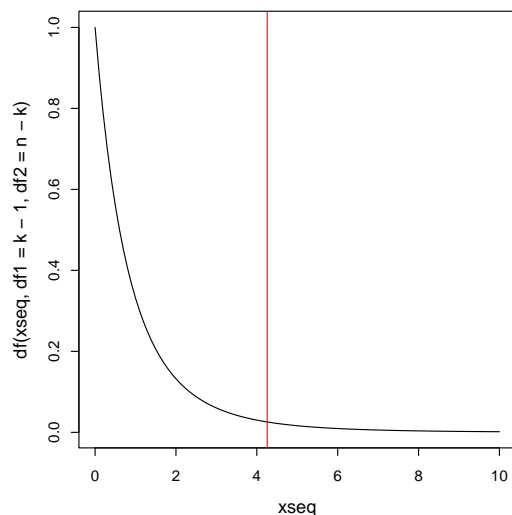
# Total number of observations
n <- 12

# Sequence for plot
xseq <- seq(0, 10, by = 0.1)

# Plot density of the F-distribution
plot(xseq, df(xseq, df1 = k-1, df2 = n-k), type = "l")

# Plot critical value for significance level 5%
cr <- qf(0.95, df1 = k-1, df2 = n-k)
abline(v = cr, col = "red")
```

An F-distribution with a critical value



The ANOVA table

Source of variation	Deg. of freedom	Sums of squares	Mean sum of squares	Test-statistic F	p -value
treatment	$k-1$	$SS(Tr)$	$MS(Tr) = \frac{SS(Tr)}{k-1}$	$F_{obs} = \frac{MS(Tr)}{MSE}$	$P(F > F_{obs})$
Residual	$n-k$	SSE	$MSE = \frac{SSE}{n-k}$		
Total	$n-1$	SST			

```
anova(lm(y ~ treatm))

## Analysis of Variance Table
##
## Response: y
##          Df Sum Sq Mean Sq F value Pr(>F)
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## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

One-way ANOVA F-test “by hand”

```

k <- 3; n <- 12 # Number of groups k, total number of observations n

# Total variation, SST
(SST <- sum( (y - mean(y))^2 ))

# Residual variance after model fit, SSE
y1 <- y[1:4]; y2 <- y[5:8]; y3 <- y[9:12]

(SSE <- sum( (y1 - mean(y1))^2 ) +
  sum( (y2 - mean(y2))^2 ) +
  sum( (y3 - mean(y3))^2 ))

# Variance explained by the model, SS(Tr)
(SSTr <- SST - SSE)

# Test statistic
(Fobs <- (SSTr/(k-1)) / (SSE/(n-k)))

# P-value
(1 - pf(Fobs, df1 = k-1, df2 = n-k))

```

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Within-group variability and relation to the 2-sample t-test (Theorem ??)

The residual sum of squares, SSE , divided by $n - k$, also called residual mean square, $MSE = SSE/(n - k)$, is the average within-group variability:

$$MSE = \frac{SSE}{n - k} = \frac{(n_1 - 1)s_1^2 + \dots + (n_k - 1)s_k^2}{n - k} \quad (1)$$

$$s_i^2 = \frac{1}{n_i - 1} \sum_{j=1}^{n_i} (y_{ij} - \bar{y}_i)^2$$

ONLY when $k = 2$: (cf. Method ??)

$$MSE = s_p^2 = \frac{(n_1 - 1)s_1^2 + (n_2 - 1)s_2^2}{n - 2}$$

$$F_{\text{obs}} = t_{\text{obs}}^2$$

where t_{obs} is the pooled t-test statistic from Methods ?? and ??.

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Post hoc confidence interval - Method ??

- A single pre-planned confidence interval for the difference between treatment i and j is found as:

$$\bar{y}_i - \bar{y}_j \pm t_{1-\alpha/2} \sqrt{\frac{SSE}{n-k} \left(\frac{1}{n_i} + \frac{1}{n_j} \right)} \quad (2)$$

where $t_{1-\alpha/2}$ is based on the t-distribution with $n - k$ degrees of freedom.

- Note the fewer degrees of freedom as more unknowns are estimated in the computation of $MSE = SSE/(n - k) = s_p^2$ (i.e. pooled variance estimate)
- If all $M = k(k - 1)/2$ combinations of pairwise confidence intervals are found use the formula M times, but each time with $\alpha_{\text{Bonferroni}} = \alpha/M$.

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Post hoc pairwise hypothesis test- Method ??

- A single pre-planned level α hypothesis test:

$$H_0 : \mu_i = \mu_j, \quad H_1 : \mu_i \neq \mu_j$$

is carried out as:

$$t_{\text{obs}} = \frac{\bar{y}_i - \bar{y}_j}{\sqrt{MSE \left(\frac{1}{n_i} + \frac{1}{n_j} \right)}} \quad (3)$$

and

$$p\text{-value} = 2P(t > |t_{\text{obs}}|)$$

where the t -distribution with $n - k$ degrees of freedom is used.

- If all $M = k(k - 1)/2$ combinations of pairwise hypothesis tests are carried out use the approach M times, but each time with significance level $\alpha_{\text{Bonferroni}} = \alpha/M$.

Variance homogeneity

Look at a box plot to check whether the variability seems different across the groups.

```
# Check assumption of homogeneous variance using, e.g.,
# a box plot.
plot(treatm, y)
```

Normal assumption

Look at a normal QQ-plot of the residuals

```
# Check normality of residuals using a normal QQ-plot
fit1 <- lm(y ~ treatm)
qqnorm(fit1$residuals)
qqline(fit1$residuals)
```

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A complete example - from the book

Introduction to Statistics

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Dokumentlegeskaber...

8.2.5 A complete worked through example: plastic types for lamps

||| **Example 8.17 Plastic types for lamps**

On a lamp two plastic screens are to be mounted. It is essential that these plastic screens have a good impact strength. Therefore an experiment is carried out for 5 different types of plastic. 6 samples in each plastic type are tested. The strengths of these items are determined. The following measurement data was found (strength in kJ/m^2):

	Type of plastic				
	I	II	III	IV	V
	44.6	52.8	53.1	51.5	48.2
	50.5	58.3	50.0	53.7	40.8
	46.3	55.4	54.4	50.5	44.5
	48.5	57.4	55.3	54.4	43.9
	45.2	58.1	50.6	47.5	45.9
	52.3	54.6	53.4	47.8	42.5

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