

05

데이터분석방법론(1)

# Analysis of Variance

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# 학습목차

- ① One-way analysis of variance
- ② Pairwise comparisons and multiple testing
- ③ Two-way analysis of variance

01

# One-way analysis of variance

# 1. Data Structure

그룹	1	...	i	...	k
1	$x_{11}$		$x_{i1}$	...	$x_{k1}$
...					
J	$x_{1j}$	...	$x_{ij}$	...	$x_{kj}$
...					
n	$x_{1n}$	...	$x_{in}$	...	$x_{kn}$
평균	$\bar{x}_1$		$\bar{x}_i$		$\bar{x}_k$

- Let  $x_{ij}$  denote observation no.  $j$  in group  $i$ , so that  $x_{35}$  is the fifth observation in group 3;  $\bar{x}_i$  is the mean for group  $i$ , and  $\bar{x}_.$  is the grand mean (average of all observations).
- Decompose  $x_{ij} - \bar{x}_. = (x_{ij} - \bar{x}_i) + (\bar{x}_i - \bar{x}_.)$
- Model  $X_{ij} = \mu + \alpha_i + \varepsilon_{ij}, \varepsilon_{ij} \sim N(0, \sigma^2)$

## 2. Decomposition of Variation

Now consider the sums of squares of the underbraced terms, known as *variation within groups*

$$SSD_W = \sum_i \sum_j (x_{ij} - \bar{x}_i)^2$$

and *variation between groups*

$$SSD_B = \sum_i \sum_j (\bar{x}_i - \bar{x}_.)^2 = \sum_i n_i (\bar{x}_i - \bar{x}_.)^2$$

It is possible to prove that

$$SSD_B + SSD_W = SSD_{\text{total}} = \sum_i \sum_j (x_{ij} - \bar{x}_.)^2$$

### 3. ANOVA table and test

- 분산분석표 (Analysis of Variance Table)

요인	제곱합	자유도	평균제곱	F값
그룹간	$SSD_B$	$k-1$	$MS_B$	$MS_B/MS_W$
그룹내	$SSD_W$	$N-k$	$MS_W$	
합	$SSD_{total}$	$N-1$		

- 검정  $H_0 : \alpha_1 = \alpha_2 = \dots = \alpha_k$  vs  $H_1 : \text{not } H_0$

검정통계량  $F_0 = MS_B/MS_W \sim F(k-1, N-k)$

## 4. One-way analysis of variance example

- “red cell folate” data from Altman (1991, p. 208).

```
> library(ISwR)
> head(red.cell.folate, 2)
  folate ventilation
1    243  N20+O2,24h
2    251  N20+O2,24h
> attach(red.cell.folate)
> summary(red.cell.folate)
   folate      ventilation
Min.   :206.0  N20+O2,24h:8
1st Qu.:249.5  N20+O2,op  :9
Median :274.0  O2,24h    :5
Mean   :283.2
3rd Qu.:305.5
Max.   :392.0
> anova(lm(folate~ventilation))
Analysis of Variance Table
Response: folate
          Df Sum Sq Mean Sq F value Pr(>F)
ventilation 2  15516  7757.9   3.7113 0.04359 *
Residuals  19  39716  2090.3
```

## 5. One-way analysis of variance example (Common mistakes)

- (Recall) the data set “juul”

```
> juul[c(6,350,450),]
      age menarche sex igf1 tanner testvol
6      0.17      NA   1  101        1      NA
350  13.20      NA   1   NA        2        8
450  16.09      NA   1  412        5       18
```

```
> anova(lm(igf1~tanner, data=juul))
Analysis of Variance Table

Response: igf1
          Df    Sum Sq Mean Sq F value    Pr(>F)
tanner     1 10985605 10985605  686.07 < 2.2e-16 ***
Residuals 790 12649728    16012
```

- Wrong !!! Since tanner is not a factor variable.
- This does not describe a grouping of data but a linear regression on the group number!  
Notice “1 DF” for the effect of tanner in the table.



## 5. One-way analysis of variance example (Correction)

- (Recall) the data set “juul” : fix

```
> juul$tanner <- factor(juul$tanner, labels=c("I","II","III","IV","V"))
```

```
> summary(juul$tanner)
```

```
 I   II  III  IV   V NA's
515 103  72  81 328 240
```

```
> anova(lm(igf1~tanner, data=juul))
```

Analysis of Variance Table

Response: igf1

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
tanner	4	12696217	3174054	228.35	< 2.2e-16 ***
Residuals	787	10939116	13900		

02

# Pairwise comparisons and multiple testing

# 1. Detecting a difference between groups

## ◉ Difference shown by the F test (ANOVA)

- If the F test shows that there is a difference between groups, the question quickly arises of where the difference lies. It becomes necessary to compare the individual groups.
- Part of this information can be found in the regression coefficients.

```
> summary(lm(folate~ventilation)) # 가변수 회귀모형
```

```
...
```

```
Coefficients:
```

	Estimate	Std. Error	t value	Pr(> t )	
(Intercept)	316.63	16.16	19.588	4.65e-14	***
ventilationN2O+O2,op	-60.18	22.22	-2.709	0.0139	*
ventilationO2,24h	-38.63	26.06	-1.482	0.1548	

```
...
```

- The interpretation of the estimates is that the intercept is the mean in the first group (N2O+O2,24h), whereas the two others describe the difference between the relevant group and the first one.

# 1. Detecting a difference between groups

```
> summary(lm(folate~ventilation)) # 가변수 회귀모형
...
Coefficients:
                Estimate Std. Error t value Pr(>|t|)
(Intercept)      316.63      16.16   19.588 4.65e-14 ***
ventilationN20+O2 -60.18      22.22   -2.709  0.0139 *
ventilationO2,24h -38.63      26.06   -1.482  0.1548
...
```

- Contrasts used by default are the so-called treatment contrasts, in which the first group is treated as a baseline and the other groups are given relative to that. Concretely, the analysis is performed as a multiple regression analysis by introducing two dummy variables, which are 1 for observations in the relevant group and 0 elsewhere.
- Among the t tests in the table, you can immediately find a test for the hypothesis that the first two groups have the same true mean ( $p = 0.0139$ , reject) and also whether the first and the third might be identical ( $p = 0.1548$ , not reject). However, a comparison of the last two groups cannot be found. This can be overcome by modifying the factor definition, but that gets tedious when there are more than a few groups.

## 2. Pairwise test (pooled SD) and Bonferroni correction

- Multiple testing (다중비교) : If we want to compare all groups, we ought to correct for multiple testing. Performing many tests will increase the probability of finding one of them to be significant; that is, the p-values tend to be exaggerated. A common adjustment method is the Bonferroni correction.
- Multiplying the p-values by the number of tests, we obtain a conservative test where the probability of a significant result is less than or equal to the formal significance level.

```
> pairwise.t.test(folate, ventilation, p.adj="bonferroni")
```

```
Pairwise comparisons using t tests with pooled SD
data: folate and ventilation
```

	N20+O2, 24h	N20+O2, op
N20+O2, op	0.042	—
O2, 24h	0.464	1.000

```
P value adjustment method: bonferroni
```

### 3. Not assuming equal variances

- Relaxing the variance assumption

- The traditional one-way ANOVA requires an assumption of equal variances for all groups. There is, however, an alternative procedure that does not require that assumption. It is (due to Welch) similar to the unequal-variances t test. This has been implemented in the `oneway.test` function:

```
> oneway.test(folate~ventilation)
```

```
One-way analysis of means (not assuming equal variances)
```

```
data: folate and ventilation
```

```
F = 2.9704, num df = 2.000, denom df = 11.065, p-value = 0.09277
```

- In this case, the p-value increased to an insignificant value, presumably related to the fact that the group that seems to differ from the two others also has the largest variance.

## 4. Pairwise test not using pooled SD

- It is also possible to perform the pairwise t tests so that they do not use a common pooled standard deviation. This is controlled by the argument `pool.sd`.

```
> pairwise.t.test(folate,ventilation,pool.sd=F)
```

Pairwise comparisons using t tests with non-pooled SD

data: folate and ventilation

	N2O+O2,24h	N2O+O2,op
N2O+O2,op	0.087	—
O2,24h	0.321	0.321

P value adjustment method: holm

- Again, it is seen that the significance disappears as we remove the constraint on the variances.

## 5. Test for same variance – Bartlett's test

### • Test for same variance in all groups

- Testing whether the distribution of a variable has the same variance in all groups can be done using Bartlett's test, although like the F test for comparing two variances, it is rather non-robust against departures from the assumption of normal distributions.

> **bartlett.test(folate~ventilation)**

Bartlett test of homogeneity of variances

data: folate by ventilation

Bartlett's K-squared = 2.0951, df = 2, p-value = **0.3508**

$$\chi^2 = \frac{(N - k) \ln(S_p^2) - \sum_{i=1}^k (n_i - 1) \ln(S_i^2)}{1 + \frac{1}{3(k-1)} \left( \sum_{i=1}^k \left( \frac{1}{n_i - 1} \right) - \frac{1}{N - k} \right)}$$

$$\text{where } N = \sum_{i=1}^k n_i \text{ and } S_p^2 = \frac{1}{N - k} \sum_{i=1}^k (n_i - 1) S_i^2$$

Reject  $H_0$  if  $\chi^2 > \chi_{\alpha, k-1}^2$

- nothing in the data contradicts the assumption of equal variances in the three groups.

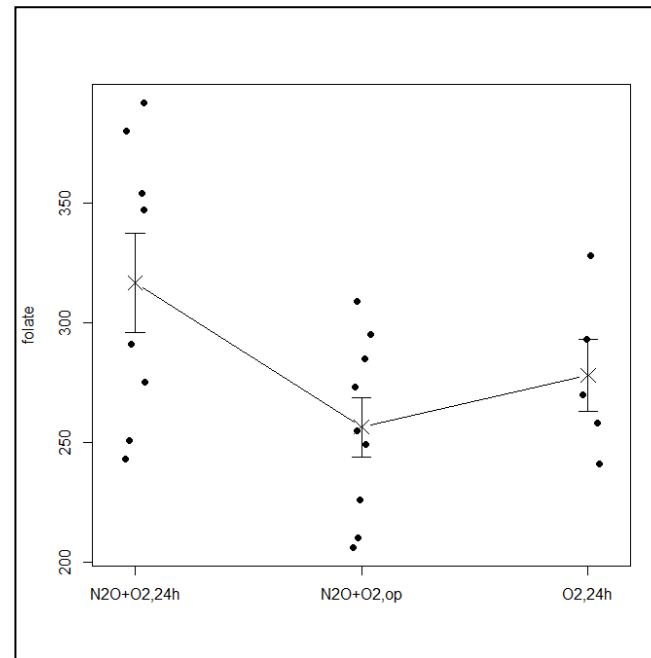


## 6. Graphical presentation

### • Stripchart

- Plot where the raw data are plotted as a stripchart and overlaid with an indication of means and SEMs.

```
> xbar <- tapply(folate, ventilation, mean)
> s <- tapply(folate, ventilation, sd)
> sem <- s/sqrt(n)
> stripchart(folate~ventilation,
method="jitter",
+ jitter=0.05, pch=16, vert=T)
> arrows(1:3,xbar+sem,1:3,xbar-sem,
+       angle=90,code=3, length=.1)
> lines(1:3,xbar,pch=4,type="b",cex=2)
```



- In many fields it appears to have become the tradition to use 1 SEM “because they are the smallest”; that is, it makes differences look more dramatic.

03

## Two-way analysis of variance

# 1. Decomposition of Variation

- Let  $x_{ij}$  denote the observation in row  $i$  and column  $j$  of the  $m \times n$  table. This is similar to the notation used for one-way analysis of variance, but notice that there is now a connection between observations with the same  $j$ , so that it makes sense to look at both row averages  $\bar{x}_{i.}$  and column averages  $\bar{x}_{.j}$ . Here we restrict ourselves to the case of a single observation per cell.
- Decompose**  $x_{ij} - \bar{x}_{..} = (\bar{x}_{i.} - \bar{x}_{..}) + (\bar{x}_{.j} - \bar{x}_{..}) + (x_{ij} - \bar{x}_{i.} - \bar{x}_{.j} + \bar{x}_{..})$

Consequently, it now makes sense to look at both *variation between rows*

$$SSD_R = n \sum_i (\bar{x}_{i.} - \bar{x}_{..})^2$$

and *variation between columns*

$$SSD_C = m \sum_j (\bar{x}_{.j} - \bar{x}_{..})^2$$

Subtracting these two from the total variation leaves the *residual variation*, which works out as

$$SSD_{\text{res}} = \sum_i \sum_j (x_{ij} - \bar{x}_{i.} - \bar{x}_{.j} + \bar{x}_{..})^2$$

## 2. Two-way analysis of variance example

- Model  $X_{ij} = \mu + \alpha_i + \beta_j + \varepsilon_{ij}, \varepsilon_{ij} \sim N(0, \sigma^2)$
- ex) heart rate after administration of enalaprilate (Altman, 1991, p. 327)

```
> library(ISwR)
> heart.rate[c(1,10,20,28),]
```

	hr	subj	time
1	96	1	0
10	92	1	30
20	108	2	60
28	92	1	120

```
heart.rate <- data.frame(hr = c(96,110,89,95,128,100,72,79,100,
                               92,106,86,78,124,98,68,75,106,
                               86,108,85,78,118,100,67,74,104,
                               92,114,83,83,118,94,71,74,102),
                        subj=gl(9,1,36),
                        time=gl(4,9,36,labels=c(0,30,60,120)))
```

## 2. Two-way analysis of variance example

```
> anova(lm(hr~subj+time, data=heart.rate))
```

Analysis of Variance Table

Response: hr

	Df	Sum Sq	Mean Sq	F value	Pr(>F)
subj	8	8966.6	1120.82	90.6391	4.863e-16 ***
time	3	151.0	50.32	4.0696	0.01802 *
Residuals	24	296.8	12.37		

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

다음시간 안내

06

# Simple Linear Regression