01NAEX - Lecture 10 Introduction to Mixed Linear Models

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Introduction to Mixed Linear Models

Throughout the lecture we have assumed that the factors in an experiment were **fixed factors** and therefore the statistical inference made about these factors are confined to the specific levels studied.

In some situations, the factor levels are chosen at random from a larger population of possible levels, and we wish to draw a conclusions about the entire population of levels, not just those that were used in the experimental design. In this situation, the factor is said to be a **random factor**.

In this lesson we focus on random effects and on mixed effects model. Nested and split-plot designs, two situations where random factors are frequently encountered in practice, will be presented in following lectures.

Introduction in to Mixed Linear Models

What is mixed models?

Mixed models model the covariance structure of data and they are generalization of linear models where observations are not independent.

What are the most common mixed models?

- Random effects models: certain effects arise from a distribution and add additional source of variation
- Random coefficients: used for repeated measures to model relationship with time, estimates are correlated.

Mixed Effects models

Mixed-effects models describe the relationship between a response variable and one or more covariates recorded with it.

- ► Fixed-effects: the factors in the experiment have a predetermined set of levels and the only inferences are for the levels of the factors actually used in the experiment.
- ▶ Random effects: the levels of factors used in the experiment are randomly selected from a population of possible levels. The inferences from the data in the experiment are for all levels of the factors in the population from which the levels were selected and not only the levels used in the experiment.
- Mixed-effects: the levels of some of the factors used in the experiment are randomly selected from a population of possible levels, whereas the levels of the other factors in the experiment are predetermined.

Introduction into Mixed Linear Models

Why use mixed models?

- To avoid mistakes.
- To get more appropriate fixed effect estimates.
- To broaden inference over wider population.
- To deal with missing data.
- To be able to handle correlation structure in data.
- To be able to handle heterscedasticity between treatment groups.
- More flexible for repeated measure and longitudinal data (allows missing data and?does not require measurements at the same time points).

Why is the terminology so various ...

Different names arose partially because these methods were re-invented in a variety of fields (psychology, education, agronomy, economics, ...), each with different slants and emphasis.

You can find ...

- Mixed-effects models
- Hierarchical linear models
- Multilevel models
- Variance component models
- Random-effects models
- Random-coefficients regression models
- **.**..

Fixed vs. random factors difference

Comparison of Fixed and Mixed model

	Fixed Model	Mixed Model	
Levels	Given number of possibilities	Selected at random	
Levels		from a population	
New experiment	Use same levels	Use different levels	
ivew experiment		from same population	
Goal	Estimates means of fixed levels	Estimate variance	
Goal		of population of means	
Inference	$H_0: \ \mu_1 = \mu_2 = \ldots = \mu_k$	$H_0: \ \sigma_{\mu}^2 = 0$ $H_1: \sigma_{\mu}^2 > 0$	
	$H_1: \exists i, j \in \{1, \ldots, k\} \ \mu_i \neq \mu_j$	$H_1: \sigma_{\mu}^2 > 0$	

Packages for Linear Mixed Effects models in R

There are several packages in R that deal with linear mixed models, most widely used are:

- nIme Non-Linear Mixed Effects, Part of R, library (nlme)
 Fit only Gaussian outcomes, it is possible to specify the variance-covariance matrix for the random effects.
- Ime4 Linear Mixed Effects, library (lme4)
 Can be used to fit generalized mixed-effects regression models, it is not possible to specify the variance-covariance matrix for the random effects, but can handle with diagonal covariance structures or unstructured covariance matrices.
- ▶ Other useful packages: merTools, glmmTMB, brms, modelr

All of them has some advantages and some disadvantages. Notations, specifications, special functions and classes are different.

For power analysis for random effects in mixed models is possible to use package called **pamm**.

Simple example:

```
library(nlme)
library(lme4)
library (MASS)
data (oats)
names(oats) = c('block', 'variety', 'nitrogen', 'yield')
oats$mainplot = oats$variety
oats$subplot = oats$nitrogen
attach (oats)
# SIMPLE MIXED EFFECTS MODEL
model1a=lme( yield~variety*nitrogen, random = ~ 1|block)
model2a=lmer(yield~variety*nitrogen + (1|block))
# NESTED MIXED EFFECTS MODEL
model1b=lme( yield~variety*nitrogen, random=~1|block/mainplot)
model2b=lmer(yield~variety*nitrogen + (1|block/mainplot))
```

Description of data set:

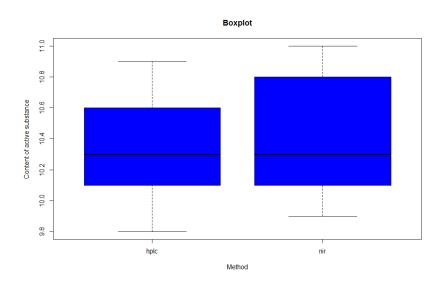
In a pharmaceutical company the use of NIR (Near Infrared Reflectance) spectroscopy was investigated as an alternative to the more cumbersome (and expensive) HPLC method to determine the content of active substance in tablets.

Source of data set:

Brockhoff and Thierry-Carstensen 2003, Test set validation using simple statistical methods.

	hplc	nir	difference
Tablet_1	10.4	10.1	0.3
Tablet_2	10.6	10.8	-0.2
Tablet_3	10.2	10.2	0.0
Tablet_4	10.1	9.9	0.2
Tablet_5	10.3	11.0	-0.7
Tablet_6	10.7	10.5	0.2
Tablet_7	10.3	10.2	0.1
Tablet_8	10.9	10.9	0.0
Tablet_9	10.1	10.4	-0.3
Tablet_10	9.8	9.9	-0.1

The aim is to study the method differences.



Simple analysis of the pharmaceutical data by the paired t-test

```
> mean(d) -0.05
> var(d) 0.08722222
> sd(d) 0.2953341
t.test(d)
# alternatively: t.test(hplc, nir, paired = TRUE)
Paired t-test
data: hplc and nir
t = -0.5354, df = 9, p-value = 0.6054
alternative hypothesis: true difference in means is not equal to 0
95 percent confidence interval: -0.2612693 0.1612693
sample estimates: mean of the differences -0.05
The standard error of the mean \bar{d} (ie. the uncertainty of the estimated difference) :
SE_{\bar{d}} = \frac{s_d}{\sqrt{n}} = \frac{0.295}{\sqrt{10}} = 0.0934
t-statistics: t = \frac{\bar{d}}{SE_z} = \frac{-0.05}{0.0934} = -0.5354
Final regression model: d_i = \mu + \varepsilon_i \quad \varepsilon_i \sim N(o, \sigma^2)
Estimated model parameters: \hat{\mu} = \bar{d}, \hat{\sigma} = s_d
```

Simple analysis of the pharmaceutical data by the ANOVA

We have randomized balanced design with two treatments - methods and 10 blocks - tablets.

Used regression model: $y_{ij} = \mu + \alpha_i + \beta_j + \varepsilon_{ij}$ $\varepsilon_{ij} \sim N(0, \sigma^2)$

The uncertainty of the average method difference:

$$SE_{\bar{y}_2 - \bar{y}_1} = \sqrt{MS_E\left(\frac{1}{n_1} + \frac{1}{n_2}\right)} = \sqrt{0.0436(\frac{1}{10} + \frac{1}{10})} = 0.0934$$

We have the same p-value as in the t-test approach and the uncertainty since the t statistic for testing $H_0: \mu_1 = \mu_2$ is

$$t = \frac{\bar{y}_2 - \bar{y}_1}{\sqrt{MS_E\left(\frac{1}{n_1} + \frac{1}{n_j}\right)}}.$$

Misleading information from the classical ANOVA approach

The ANOVA approach compare treatments means and is based on knowledge obtained only from certain measurements:

In pharmaceutical data example, the **ANOVA approach** is valid only for statements about the 10 specific tablets in the experiment, not for tablets in general.

The uncertainty of the average NIR value is given by using only the 10 NIR measurements: $SE_{\bar{y_1}} = \frac{s_1}{\sqrt{10}} = 0.127$, $s_1 = sd(NIR) = 0.4012$.

On the other hand, the **Mixed Models approach** is valid for tablets (blocks) in general, since we consider the 10 tablets as a random sample.

Let us consider Mixed Model with tablet as a random effects:

$$y_{ij} = \mu + a_i + \beta_j + \varepsilon_{ij}$$
 $\varepsilon_{ij} \sim N(0, \sigma^2), \ a_i \sim N(0, \sigma_T^2).$

Comparison of Fixed and Mixed model

The observations in Mixed Model are no longer independent and the tablet differences become a part of the variance structure.

- 1. The expected value of the observation y_{ij} .
- 2. The variance of the observation y_{ii} .
- 3. The relation between two observations.

Comparison table	OI IIXEU		IIIIXEU	Marian
Comparison table	of fivod	and	mivad	modal

		Fixed Model	Mixed Model
1	E [y _{ij}]	$\mu + \alpha_i + \beta_j$	$\mu + \beta_j$
2	var [y _{ii}]	σ^2	$\sigma_T^2 + \sigma^2$
3	$cov(y_{i_1j_1}, y_{i_2j_2}) j_1 \neq j_2$	0	σ_T^2 (if $i_1 = i_2$), 0 (if $i_1 \neq i_2$)

Analysis of the pharmaceutical data by the Mixed Models

Used mixed model:

$$y_{ij} = \mu + a_i + \beta_j + \varepsilon_{ij}$$
 $\varepsilon_{ij} \sim N(0, \sigma^2), \ a_i \sim N(0, \sigma_T^2).$

```
# ways how to estimate mixed models
# function lme (need library nlme)
# function lmer (need library lme4)
> model<-lme( y~method, random = ~1|tablet, data=temp)
> model<-lme( y~method, random=list(tablet=~1), data=temp)
> model<-lmer(y~method+(1|tablet), data=temp)
> summary(model)
```

Analysis of the pharmaceutical data by the Mixed Models

```
Used mixed model: y_{ij} = \mu + a_i + \beta_j + \varepsilon_{ij} \varepsilon_{ij} \sim N(0, \sigma^2), \ a_i \sim N(0, \sigma_T^2).
> summary(model)
Linear mixed model fit by REML
Formula: v ~ method + (1 | tablet)
Random effects:
Groups Name Variance Std.Dev.
 tablet (Intercept) 0.089333 0.29889
 Residual 0.043611 0.20883
Number of obs: 20, groups: tablet, 10
Fixed effects:
            Estimate Std. Error t value
(Intercept) 10.39000 0.11530 90.11
methodhplc -0.05000 0.09339 -0.54
Correlation of Fixed Effects:
            (Intr)
methodhplc -0.405
anova (model) numDF denDF F-value p-value
(Intercept) 1 9 9666.573 <.0001
                   1 9 0.287 0.6054
method
```

Analysis of the pharmaceutical data by the Mixed Models

Used mixed model:

$$y_{ij} = \mu + a_i + \beta_j + \varepsilon_{ij}$$
 $\varepsilon_{ij} \sim N(0, \sigma^2), \ a_i \sim N(0, \sigma_T^2),$

with estimated Random effects components:

$$\hat{\sigma}^2 = 0.043611, \quad \hat{\sigma}_T^2 = 0.089333,$$

and Fixed effects components:

$$\hat{\mu} = 10.39000, \quad \hat{\beta}_1 = 0, \quad \hat{\beta}_2 = -0.05000$$

We have the same pharmaceutical data set, but we add 5 non-paired measuremnt from each method.

```
tablet method
                              tablet method
                                                         tablet method
        1
            hplc 10.4
                          14
                                   7
                                         nir 10.2
                                                             14
                                                                   hplc
                                                                          NA
            nir 10.1
                          15
                                       hplc 10.9
                                                                    nir 10.3
                                                             14
            hplc 10.6
                          16
                                       nir 10.9
                                                      29
                                                             15
                                                                   hplc
                                                                          NA
            nir 10.8
                           17
                                       hplc 10.1
                                                      30
                                                             15
                                                                    nir
                                                                         9.7
            hplc 10.2
                           18
                                       nir 10.4
                                                      31
                                                             16
                                                                   hplc 10.3
            nir 10.2
                           19
                                  10
                                        hplc
                                             9.8
                                                      32
                                                             16
                                                                    nir
                                                                          NA
                           2.0
                                                      33
            hplc 10.1
                                  10
                                         nir
                                              9.9
                                                             17
                                                                   hplc
                                                                         9.6
            nir 9.9
                           21
                                  11
                                       hplc
                                                      34
                                                             17
                                                                    nir
                                               NA
                                                                          NA
9
            hplc 10.3
                           22
                                  11
                                        nir 10.8
                                                      35
                                                             18
                                                                   hplc 10.0
                           23
            nir 11.0
                                  12
                                       hplc
                                               NA
                                                      36
                                                             18
                                                                    nir
                                                                          NA
            hplc 10.7
                           24
                                  12
                                                             19
                                                                   hplc 10.2
                                        nir
                                              9.8
                                                      37
12
            nir 10.5
                           25
                                  13
                                       hplc
                                                             19
                                                                    nir
                                               NA
                                                      38
                                                                          NA
            hplc 10.3
                           2.6
                                  13
                                         nir 10.5
                                                      39
                                                             2.0
                                                                   hplc
                                                                         9.9
                                                      40
                                                                    nir
                                                                          NA
```

The mixed model in a direct way may give information about the key issues in a data set, that a straightforward fixed ANOVA does not.

Analysis by fixed effects ANOVA

The fixed effect analysis only uses the information in the first 10 tablets.

Note that only the Tablets row of the table has changed compared to the previous analysis.

Analysis by mixed model

```
> model2<-lme(v~method,random=~1| tablet,data=temp2t)</pre>
> anova(model2)
           numDF denDF F-value p-value
(Intercept) 1 19 15715.981 <.0001
method 1 9 0.687 0.4285
> summary(model2)
Linear mixed-effects model fit by REML
Random effects:
Formula: ~1 | tablet
       (Intercept) Residual
StdDev: 0.3192429 0.2085067
Fixed effects: y ~ method
               Value Std.Error DF t-value p-value
(Intercept) 10.283857 0.09259174 19 111.06668 0.0000
methodhplc -0.072111 0.08697180 9 -0.82913 0.4285
Correlation:
           (Intr)
methodhplc -0.47
```

Compariosn of Fixed and Mixed model approach

We use the R function estimable.

FIXED:

```
Estimate Std. Error t value DF Pr(>|t|) Lower.CI Upper.CI LS HPLC 10.2125 0.06177356 165.321550 9 0.0000000 10.1227585 10.3522415 LS NIR 10.2625 0.06177356 166.130957 9 0.0000000 10.1227585 10.4022415 LS DIF -0.0500 0.09339284 -0.535373 9 0.6053664 -0.2612693 0.1612693
```

MIXED:

```
Estimate Std. Error t value DF Pr(>|t|) Lower.CI Upper.CI LS HPLC 10.21174590 0.09259174 110.2878719 9 0.0000000 10.0022888 10.4212030 LS NIR 10.28385689 0.09259174 111.0666778 19 0.0000000 10.0900602 10.4776536 LS DIF -0.07211099 0.08697180 -0.8291307 9 0.4284719 -0.2688549 0.1246329
```

Note that apart from giving a slightly different value, the Mixed model estimation is also more precise, than the one only based on tablets 1-10. This is the kind of analysis that the mixed model for this situation leads to.

Recall simple linear regression model of **fixed effects** approach:

$$y_{ij} = \mu + \alpha_i + \beta_j + \varepsilon_{ij} \quad \varepsilon_{ij} \sim N(0, \sigma^2), \ i \in \{1, 2\}, \ j \in \{1, 2, 3\},$$

$$\begin{pmatrix} y_{11} \\ y_{21} \end{pmatrix} \quad \begin{pmatrix} 1 & 1 & 0 & 1 & 0 & 0 \\ 1 & 0 & 1 & 1 & 0 & 0 \\ 1 & 1 & 0 & 0 & 1 & 0 \end{pmatrix} \quad \begin{pmatrix} \mu \\ \alpha_1 \\ \theta_{21} \\ \theta_{21} \end{pmatrix}$$

$$\underbrace{\begin{pmatrix} y_{11} \\ y_{21} \\ y_{12} \\ y_{22} \\ y_{13} \\ y_{23} \end{pmatrix}}_{\mathbf{y}} = \underbrace{\begin{pmatrix} 1 & 1 & 0 & 1 & 0 & 0 \\ 1 & 0 & 1 & 1 & 0 & 0 \\ 1 & 1 & 0 & 0 & 1 & 0 \\ 1 & 0 & 1 & 0 & 1 & 0 \\ 1 & 1 & 0 & 0 & 0 & 1 \\ 1 & 0 & 1 & 0 & 0 & 1 \end{pmatrix}}_{\mathbf{X}} \cdot \underbrace{\begin{pmatrix} \mu \\ \alpha_1 \\ \alpha_2 \\ \beta_1 \\ \beta_2 \\ \beta_3 \end{pmatrix}}_{\beta} + \underbrace{\begin{pmatrix} e_{11} \\ e_{21} \\ e_{12} \\ e_{22} \\ e_{13} \\ e_{23} \end{pmatrix}}_{\mathbf{e}}.$$

equivalently,

$$\mathbf{Y} = \mathbf{X}\beta + \mathbf{e}$$

where \mathbf{y} is an vector of all observations - response variables, \mathbf{X} is an known matrix of predictors (usually called design matrix), β is a vector of unknown coefficients - fixed effects parameters and \mathbf{e} is an vector of unknown independent measurement errors $\mathbf{e} \sim N(\mathbf{0}, \sigma^2 \mathbf{I})$.

Regression model of **Mixed linear model** approach:

$$y_{ij} = \mu + \alpha_i + b_j + \varepsilon_{ij}$$
 $b_j \sim N(0, \sigma_B^2), \ \varepsilon_{ij} \sim N(0, \sigma^2)$ $i \in \{1, 2\}, \ j \in \{1, 2, 3\},$

$$\underbrace{\begin{pmatrix} y_{11} \\ y_{21} \\ y_{12} \\ y_{22} \\ y_{13} \\ y_{23} \end{pmatrix}}_{\mathbf{y}} = \underbrace{\begin{pmatrix} 1 & 1 & 0 \\ 1 & 0 & 1 \\ 1 & 1 & 0 \\ 1 & 0 & 1 \\ 1 & 1 & 0 \\ 1 & 0 & 1 \end{pmatrix}}_{\mathbf{X}} \cdot \underbrace{\begin{pmatrix} \mu \\ \alpha_{1} \\ \alpha_{2} \\ \beta \end{pmatrix}}_{\beta} + \underbrace{\begin{pmatrix} 1 & 0 & 0 \\ 1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & 0 & 1 \\ 0 & 0 & 1 \end{pmatrix}}_{\mathbf{z}} \cdot \underbrace{\begin{pmatrix} b_{1} \\ b_{2} \\ b_{3} \\ b_{2} \\ b_{3} \end{pmatrix}}_{u} + \underbrace{\begin{pmatrix} e_{11} \\ e_{21} \\ e_{12} \\ e_{22} \\ e_{13} \\ e_{23} \end{pmatrix}}_{\mathbf{e}},$$

equivalently,

$$\mathbf{Y} = \mathbf{X}\beta + \mathbf{Z}u + \mathbf{e}$$

where **y** is an vector of all observations - response variables, **X** is an known matrix of predictors (usually called design matrix), β^0 is a vector of unknown coefficients - fixed effects parameters, **Z** is the design matrix for random effects, **u** is the vector of random effects $\mathbf{u} \sim N(\mathbf{0}, \mathbf{G})$, $cov(u_i, u_j) = G_{ij}$ and **e** is an vector of unknown independent measurement errors $\mathbf{e} \sim N(\mathbf{0}, \mathbf{R})$, $cov(e_i, e_i) = R_{ii}$, typically **R** is diagonal.

Distribution of response variable **y** in Mixed Models:

Let us consider linear mixed model:

$$Y = X\beta + Zu + e$$
, where $u \sim N(0, G)$, $e \sim N(0, R)$.

Distribution of y is multivariate normal

$$\mathbf{y} \sim \mathit{N}(\mu, \mathbf{V})$$

with

$$\mu = E[X\beta + Zu + e] = X\beta$$

and

$$V = var[X\beta + Zu + e] = var[Zu] + var[e] = ZGZ^T + R.$$

$$\mathbf{y} \sim N(\mathbf{X}\beta, \mathbf{ZGZ}^T + \mathbf{R})$$

Notice that if ${\bf R}$ is diagonal and we have random block effect model then ${\bf V}$ is block diagonal matrix.

Example of a y distribution for simple linear Mixed Effects Models:

Let us consider linear mixed effect model:

$$y_{ij} = \mu + \alpha_i + b_j + \varepsilon_{ij}$$
 $b_j \sim N(0, \sigma_B^2), \ \varepsilon_{ij} \sim N(0, \sigma^2)$ $i \in \{1, 2\}, \ j \in \{1, 2, 3\},$

$$\mu = \left(\begin{array}{c} \mu + \alpha_1 \\ \mu + \alpha_2 \\ \mu + \alpha_1 \\ \mu + \alpha_2 \\ \mu + \alpha_1 \\ \mu + \alpha_2 \end{array} \right) \quad \mathbf{V} = \left(\begin{array}{ccccc} \sigma^2 + \sigma_B^2 & \sigma_B^2 & 0 & 0 & 0 & 0 & 0 \\ \sigma_B^2 & \sigma^2 + \sigma_B^2 & 0 & 0 & 0 & 0 & 0 \\ 0 & 0 & \sigma^2 + \sigma_B^2 & \sigma_B^2 & 0 & 0 \\ 0 & 0 & \sigma_B^2 & \sigma^2 + \sigma_B^2 & 0 & 0 \\ 0 & 0 & 0 & 0 & \sigma^2 + \sigma_B^2 & \sigma_B^2 \\ 0 & 0 & 0 & 0 & \sigma_B^2 & \sigma^2 + \sigma_B^2 \end{array} \right)$$

Notice that two observations from the same block are correlated !!

Theory of Mixed Models

The likelihood function L for mixed effect models:

For given parameter values, the likelihood function L returns a measure of the probability of observing response variables \mathbf{y} .

The negative log likelihood function for a mixed model is given by:

$$I(\mathbf{y}, \beta, \gamma) = \frac{1}{2} \left(n \ln(2\pi) + \ln|\mathbf{V}(\gamma)| + (\mathbf{y} - \mathbf{X}\beta)^{\mathsf{T}} \mathbf{V}(\gamma)^{-1} (\mathbf{y} - \mathbf{X}\beta) \right)$$

In the simple one way ANOVA with random block effect model is

$$\gamma = (\sigma^2, \sigma_b^2)^T$$
 and $\beta = (\alpha_1, \alpha_2)^T$

The maximum likelihood estimation is given by

$$(\hat{\beta}^{(ML)}, \hat{\gamma}^{(ML)}) = \operatorname{argmin}_{(\beta, \gamma)} I(\mathbf{y}, \beta, \gamma)$$

Assume γ is known then ML estimation of $\hat{\beta}(\gamma)$ is given by weighted least squares estimation

$$\hat{\beta}^{ML}(\gamma) = \operatorname{argmin}_{(\beta)} (\mathbf{y} - \mathbf{X}\beta)^{\mathsf{T}} \mathbf{V}(\gamma)^{-1} (\mathbf{y} - \mathbf{X}\beta),$$

by differentiate and equal to zero we obtain:

$$\hat{\beta}^{ML}(\gamma) = (\mathbf{X}^{\mathsf{T}} \mathbf{V}(\gamma)^{-1} \mathbf{X})^{-1} \mathbf{X}^{\mathsf{T}} \mathbf{V}(\gamma)^{-1} \mathbf{y}$$

Theory of Mixed Models

The restricted likelihood method for mixed effect models:

Since the maximum likelihood estimation is biased, we want to modify it to obtain unbiased estimator.

Idea of Restricted (residual) maximum likelihood (REML) is in linear transform of data which eliminates mean.

REML method is given by modification of classical ML function by

$$I_{\textit{REML}}(\mathbf{y},\beta,\gamma) = \frac{1}{2} \left(n \ln(2\pi) + \ln|\mathbf{V}(\gamma)| + (\mathbf{y} - \mathbf{X}\beta)^\mathsf{T} \mathbf{V}^{-1}(\gamma) (\mathbf{y} - \mathbf{X}\beta) + \ln\left|\mathbf{X}^\mathsf{T} \mathbf{V}^{-1}(\gamma) \mathbf{X}\right| \right).$$

This REML method gives (at least in balanced case) the unbiased estimates and is generally preferred in mixed models.

Theory of Mixed Models

Restricted maximum likelihood (REML)

- ► The default parameter estimation criterion for linear mixed models in lme and lmer functions is REML.
- Maximum likelihood (ML) estimates (sometimes called full maximum likelihood) can be requested by specifying REML=FALSE.
- Generally REML estimates of variance components are preferred. ML estimates are known to be biased. Although REML estimates are not guaranteed to be unbiased, they are usually less biased than ML estimates.

Example of Mixed effect model analysis

Drying of beech wood planks

To investigate the effect of drying of beech wood on the humidity percentage, the following experiment was conducted. Each of 20 planks was dried in a certain period of time. Then the humidity percentage was measured in 5 depths (1,3,5,7,9) and 3 widths (1,2,3) for each plank.

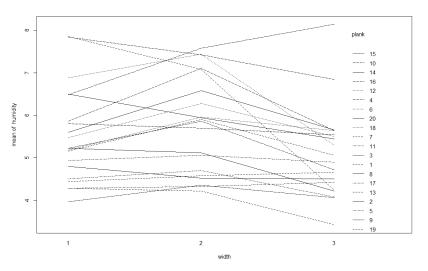
Variables:

- plank Numbered 1-20
- width Numbered 1,2,3
- depth Numbered 1,3,5,7,9
- humidity Humidity percentage

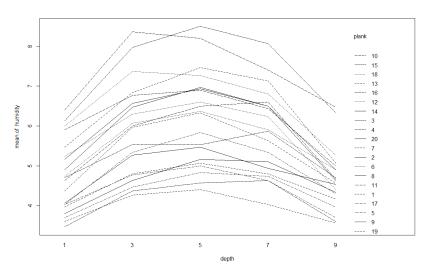
Source: The Royal Veterinary and Agricultural University, Denmark. **Number of observations:** 300 (20 planks)

```
depth 1: close to the top
depth 5: in the center
depth 9: close to the bottom
depth 3: between 1 and 5
depth 7: between 5 and 9
width 1: close to the side
width 3: in the center
width 2: between 1 and 3
```

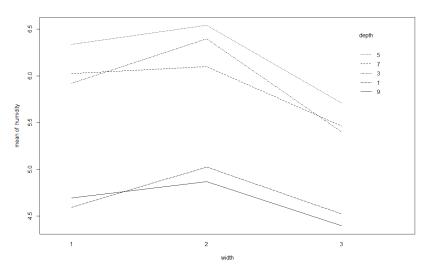
with(planks, interaction.plot(width,plank,humidity,legend=T))



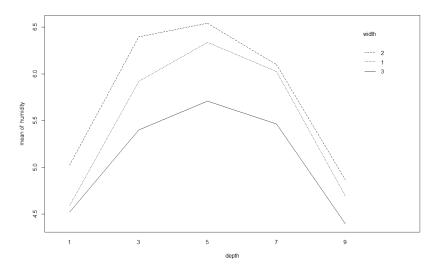
with (planks, interaction.plot(depth,plank,humidity,legend=T))



with(planks, interaction.plot(width,depth,humidity,legend=T))



with(planks, interaction.plot(depth,width,humidity,legend=T))



Exercise

Analyze data from the **Drying of beech wood planks** experiment.

- Plot four average humidity profiles:2 interaction plots for width and 2 for depth (done).
- Carrying out the fixed effects model analysis.
- Carry out the mixed model analysis.
- Run the post hoc analysis
- Compare the fixed parameters and use the p-value correction (TukeyHSD).
 - Hint: Use function lsmeans from the package lsmeans with adjust="tukey".
- Summarize results.