ADVANCED DATA ANALYSIS FOR PSYCHOLOGICAL SCIENCE

Part 1. Introduction to multilevel modeling

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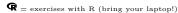
Master degree in Developmental and Educational Psychology
University of Padova
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Outline of Part 1

- LM recap: Short recap of linear regression modeling 🖢 🗨
- LMER: Introduction to multilevel modeling (linear mixed-effects regression)
- Data processing: How to approach a multilevel data structure?
 How to manipulate and pre-process multilevel data?
- **Descriptives**: Which descriptive stats should be reported from a multilevel dataset? How to compute and interpret them?
- Model fit: How to fit a multilevel model in R? How to inspect, report, visualize, and interpret the results of a multilevel model? \P
- Model evaluation: Which are the assumptions of multilevel models? How to evaluate them? How to compare multiple models and select the best model? •
- Related: Summaries & in-depth topics related to multilevel modeling (e.g., generalized and Bayesian LMER, power analysis)

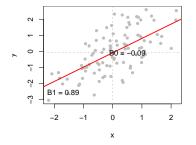
[■] not for the exam



Linear regression models

Linear models (LM) allow to determinate the link between two variables as expressed by a linear function: $y_i = \beta_0 + \beta_1 x_i + \epsilon_i$ Such a function can be graphically represented as a **straight line**, where:

- β_0 is the **intercept** (value assumed by y when x = 0)
- β_1 is the **slope** (predicted change in y when x increases by 1 unit)
- ϵ_i are the **errors** (distance between observation i and the regression line)



 x_i and y_i are the values of observation i for the casual variables x and y

 β_0 , β_1 , and ϵ_i are called "**parameters**", or "**coefficients**". They are *estimated* from the sampled data and *generalized* to the whole population.

Fitting linear models in R

```
data("children", package = "npregfast") # loading children dataset from npregfast pkg
```

R uses the lm() function to fit linear models with the arguments formula $(y \sim x1 + x2 + ...)$ and data (identifying the dataframe with the model variables).

Null model

[1] 243.9085

Children' height is only predicted by the model intercept $\beta_0=$ expected (i.e., mean) value of height in the sample. σ^2 is the variance of the residuals ϵ_i (deviations from the intercept).

```
m0 <- lm(formula = height ~ 1,
data = children)

coefficients(m0) # model parameters

(Intercept)
153.4013

summary(m0)$sigma^2 # residual variance
```

Simple regression model

```
height is now predicted by the intercept \beta_0 (mean value when age is 0), the slope \beta_1 (expected change for 1-unit increase in age), and the residual variance \sigma^2.
```

```
m1 <- lm(formula = height - age,
data = children)

coefficients(m1) # model parameters

(Intercept) age
94.904099 4.388803

summary(m1)$sigma^2 # residual variance

[1] 56.19656
```

Multiple regression & interactions

LM also allow to include **multiple predictors** and the **interactions**¹ among them. This is done by estimating a separate slope (thus, a separate line) for each predictor by *holding constant* the value of the other predictors, which are fixed to zero.

Multiple regression model

```
eta_0 = 	ext{expected value in girls with age} = 0
eta_1 = 	ext{age effect}^2 	ext{ within the same sex}
eta_2 = 	ext{sex difference when age} = 0
	ext{m2} < - 	ext{lm(formula = height - age + sex,}
	ext{data = children)}
	ext{coefficients(m2)}
(Intercept) age sextale 95.0075706 4.3887983 -0.2001025
```

Interactive model $\beta_1 = \text{age effect in girls}$

104.25

3.70

-19.04

1.41

¹The interaction between x_1 and x_2 is computed as the product of x_1 and x_2 .

 $^{^2}$ In this context, "effect" is used as a synonym of "relationship" (not a causal effect).

Model comparison & model selection

Likelihood ratio test

Compares the fit of two nested models (i.e., predicting the same y variable, with the more complex model including all predictors included in the simpler model).

```
library(lmtest)
```

```
lrtest(m0.m1.m2.m3) # returns Chisa statistic
  #Df
        LogLik Df
                    Chisa
                            Pr(>Chisa)
   2 -10417.84 NA
                       NA
      -8582.42 1 3670.84 0.000000e+00
```

```
4 -8582 19 1
                 0.45 5.046155e-01
  -8468.86 1
               226.67 3.176229e-51
```

Information criteria

The Akaike (AIC) and the Bayesian Information Criterion (BIC) compare multiple models in terms of fit & parsimony (the lower number of parameters the better)

```
AIC(m0,m1,m2,m3) # AIC: the lower the better
[1] 20839.68 17170.83 17172.39 16947.72
# Akaike weights: from 0 (-) to 1 (+)
MuMIn::Weights(AIC(m0,m1,m2,m3))
model weights
[1] 0 0 0 1
```

Here, model fit to the data is expressed by its likelihood = probability of observing the sampled data given the parameters estimated by the model, sometimes referred as the evidence of a model, or its ability to predict/forecast new data that are similar to the sampled data (see interactive visualization by Kristoffer Magnusson).

 β_0 , β_1 , and ϵ must be **estimated** based on data sampled from a population:

$$\hat{\beta}_0 = b_0; \, \hat{\beta}_1 = b_1; \, \hat{\epsilon} = e$$
).

b There are several methods to estimate unknown parameters, such as:

- Ordinary least squares (OLS): finds the parameter values that minimize the sum of the squared residuals (default LM estimator)
- Maximum likelihood estimator (MLE): finds the parameter values that maximize the model likelihood, making the observed data the most probable under that model
- Bayesian estimator: finds the parameter posterior distributions based on prior knowledge/beliefs (prior) and observed data (likelihood)

Regardless of the used method, parameters values (or distributions) are always accompanied with a measure of the uncertainty/precision associated with their estimate:

Standard errors (SE) = predicted variability in the parameter estimate if the data were collected from different random samples from the same population.

SE are used for computing test statistics (Est/SE) & confidence intervals (Est \pm 1.96 \times SE)

head(data.frame(observed = children\$height,

var(residuals(m3)) # residual variance SIGMA2

What are residuals?

Residuals are the model-based estimates of the population errors.

```
Linear model:
                                                                  predicted = fitted(m3),
u_i = \beta_0 + \beta_1 x_i + \epsilon_i
                                                                  residuals = residuals(m3)
                                                                  squared = residuals(m3)^2 ))
Predicted values:
                                                  observed predicted residuals squared
\hat{y}_i = \beta_0 + \beta_1 x_i
                                                    150.77
                                                               152.90
                                                                          -2.13
                                                                                    4.55
                                                    170.59
                                                              156.61
                                                                          13.98 195.33
                                                    167.31
                                                              160.31
                                                                           7.00
                                                                                 49.01
                                                              165.52
                                                    165.72
                                                                           0.20
                                                                                    0.04
Observed values:
                                                    171.67
                                                              160.31
                                                                          11.36 129.06
                                                    143.74
                                                              151.07
                                                                          -7.33
                                                                                   53.74
y_i = \hat{y}_i + \hat{\epsilon}_i
                                               sum(residuals(m3)^2) # sum of squared (SS) residuals
Residuals = observed - predicted
                                               ## [1] 128188.3
```

[1] 51.29585

In LM, model parameters include:

 $\hat{\epsilon}_i = y_i - \hat{y}_i$

- (1) intercept, (2) slope(s), and (3) **residual variance** σ^2
- \rightarrow How many parameters in the previous models? (= No. predictors + 2)

Statistical inference on regression coefficients

In the NHST approach, we can **test the statistical** significance of regression coefficients (two-tail t-test).

This is automatically done by R in the model summary.

summary(m3) # model results

	Estimate	Std.	Error	t value	Pr(> t)
(Intercept)	104.25		0.88	118.22	0.000000e+00
age	3.70		0.06	57.45	0.000000e+00
sexmale	-19.04		1.26	-15.14	1.237494e-49
age:sexmale	1.41		0.09	15.39	3.897810e-51

- Estimate = estimated parameter
- Std. Error = parameter standard error
- ${\tt t}$ value = test statistic computed as
- t = Estimate/Std.Error
- p-value = p corresponding to the t-value with No. Obs. No. Coeff. 1
 degrees of freedom

Effect size:

Coefficient of determination

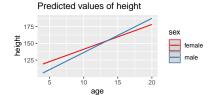
$$R^2 = 1 - SS_{residuals} / SS_{total}$$

[1] 0.79

The model explains 79% of the variance in height.

Plotting effects:

```
sjPlot::plot_model(m3,type="pred",terms=c("age","sex"))
```



Hands on **Q**

1. Download & read the dataset from the "Pregnancy during pandemics" study

```
depr = postnatal depression, age = mother's age, NICU = intensive care, threat = fear of COVID
library(osfr) # package to interact with the Open Science Framework platform
proj <- "https://osf.io/ha5dp/" # link to the OSF project
osf_download(osf_ls_files(osf_retrieve_node(proj))[2, ],conflicts="overwrite") # download
preg <- na.omit(read.csv("OSFData_Upload_2023_Mar30.csv",stringsAsFactors=TRUE)) # read data
colnames(preg)[c(2,5,12,14)] <- c("age","depr","NICU","threat") # set variable names</pre>
```

- Explore the the variables depr, threat, NICU, and age (descr., corr., & plots)
- 3. Fit a null model m0 of depr
- Fit a simple regression model m1 with depr being predicted by threat
- Fit a multiple regression model m2 also controlling for NICU and age
- Fit an interactive model m3 to check whether age moderates the relationship between threat and depr.

- 7. Compare the models with AIC and likelihood ratio test: which is the best model?
- Print & interpret the coefficients estimated by the selected model
- Print & interpret the statistical significance of the estimated coefficients
- 10. Plot the effects of the selected model
- 11. Compute the determination coefficient of the selected model

One step back: Linear model assumptions

Core assumptions:

- 1. Linearity: x_i and y_i are linearly associated \rightarrow the expected (mean) value of ϵ_i is zero
- 2. Normality: residuals ϵ_i are normally distributed with $\epsilon_i \sim \mathcal{N}(0, \sigma^2)$
- 3. Homoscedasticity: ϵ_i variance is constant over the levels of x_i (homogeneity of variance)
- 4. Independence of predictors & errors: predictors x_i are unrelated to residuals ϵ_i
- 5. Independence of observations: for any two observations i and j with $i \neq j$, the residual terms ϵ_i and ϵ_j are independent (no common disturbance factors)

Additional assumptions:

- 6. Absence of influential observations (multivariate outliers)
- 7. Absence of multicollinearity (for multiple regression):

lack of linear relationship between x_1 and x_2

Model diagnostics: Assessing LM assumptions

Normality & linearity ©

hist(residuals(m3))

qqnorm(residuals(m3)); qqline(residuals(m3))





Homoscedasticity & independence $x, \epsilon \odot$

plot(residuals(m3) ~ children\$sex)
plot(residuals(m3) ~ children\$age)

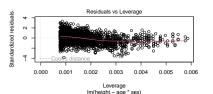




Independence of observations ?

Absence of influential cases ©

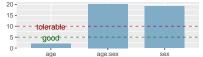
plot(m3,which=5)



Absence of multicollinearity

siPlot::plot model(m3."diag")[[1]]

Variance Inflation Factors (multicollinearity)



Are the unmeasured factors influencing y unrelated from one individual to another?

Cluster variables & nested data

In many cases, the sampling method creates clusters of individual observations

- students → schools
- children \rightarrow families \rightarrow neighborhoods \rightarrow cities \rightarrow regions \rightarrow states \rightarrow planets \P

Nested data structure (= multilevel or hierarchical data structure)

- = when data points at the **individual level** appear *in only one group* of the **cluster level** variable
- \rightarrow individual observations are **nested** within clusters

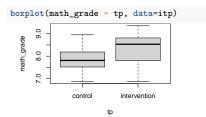
How do you imagine such a nested dataset?

Individual observation = statistical unit = individual entity within a sample or population that is the subject of data collection & analysis (not necessarily a person)

We're hired by a school principal to assess whether an innovative teaching program can improve math achievement in first-year high-school students.

```
# reading data
itp <- read.csv("data/studentData.csv")</pre>
# frequency table class by intervention
table(itp[,c("classID","tp")])
```

	tp		
intervention	control	classID	
0	30	A	
0	22	В	
27	0	C	
- 11	0	D	



The teaching program tp was delivered over the first semester to 2 out of 4 classes and we got the students' end-of-semester math_grade (1-10).

Nested dataset: students are nested within classes, with each student only belonging to one class.

head(itp[,1:4],12)

	-			
	studID	classID	tp	math_grade
1	s1	A	control	7.74
2	s2	A	control	8.31
3	s3	A	${\tt control}$	7.09
4	s4	A	control	7.80
5	s5	A	control	7.21
6	s6	A	${\tt control}$	8.95
7	s7	A	${\tt control}$	7.48
8	s8	A	${\tt control}$	7.86
9	s9	A	${\tt control}$	7.85
10	s10	A	${\tt control}$	7.13
11	s11	A	${\tt control}$	7.87
12	s12	A	control	6.88

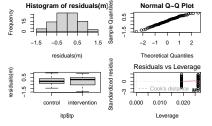
Non-independence of observations with nested data

Let's try with a linear regression model:

```
m <- lm(math_grade ~ tp, data=itp)
summary(m)$coefficients[,1:3]
## Estimate Std. Error t value
## (Intercept) 7.85 0.08 97.60
## tpintervention 0.48 0.12 3.87</pre>
```

Model diagnostics (see slide #11):

hist(residuals(m)); qqnorm(residuals(m))
boxplot(residuals(m)~itp\$tp); plot(m,5)



- Coefficient meaning?
- Linear model assumptions?
- Independent observations?

Are ϵ_i and ϵ_j independent for any $i \neq j$? Are the unmeasured factors influencing y unrelated from one individual to another?

NO: students are nested within classes and such cluster variable is likely to explain differences in the y variable (as well as in the relationship between x and y)

Thus, we cannot rely on linear models to analyze these data.

LMER

Local dependencies

Local dependencies = correlations that exist among observations within a specific cluster (but the software doesn't know that!)

e.g., grades from the same class will be more correlated than they are between different classes

Why is this a problem?

- 1) Can result in biased estimates of the standard errors \rightarrow underestimated p-values (+false positive)
- Potentially important variables at the cluster level are neglected e.g., teachers' characteristics, teaching CV, class social climate

When is this a problem?

clustering factors for both y and x

Virtually, any time that a cluster variable is potentially related to y Pragmatically, we cannot account for all potential clusters e.g., children \to families \to neighborhoods \to cities \to regions \to states \to planets \P Based on theory & logic, we should focus on what we consider the most influential

Mixed-effects models

Multilevel models are part of the largest linear mixed-effects regression (LMER) family that include additional variance terms for handling local dependencies.

Why 'mixed-effects'?

Because such additional terms come from the distinction between:

- Fixed effects: effects that remain constant across clusters, whose levels are
 exhaustively considered (e.g., gender, levels of a Likert scale) and generally
 controlled by the researcher (e.g., experimental conditions)
- Random effects: effects that vary from cluster to cluster, whose levels are randomly sampled from a population (e.g., schools)

b When individual observations can change cluster over time, it is still a mixed-effects model but not a multilevel model.

b Here, "levels" refers to the possible categories/classes of a categorical variable, but from now on we will use this term with a different meaning...

From LM to LMER.

LM formula: $y_i = \beta_0 + \beta_1 x_i + \epsilon_i$ Intercept and slope are **constant across** all individual observations i within the population; x, y, and the error term ϵ only variate across individual observations i LMER formula: $y_{ij} = \beta_{0j} + \beta_{1j}x_{ij} + \epsilon_{ij}$ Intercept and slope have both a fixed (0/1) and a random component (j); y, x, and ϵ variate across individual observations i as well as across clusters j

$$y_{ij} = \beta_{0j} + \beta_{1j}x_{ij} + \epsilon_{ij} = (\beta_{00} + \lambda_{0j}) + (\beta_{10} + \lambda_{1j})x + \epsilon_{ij}$$

LMER are an extension of LM where the intercept and the slope are decomposed into the fixed components β_{00} and β_{10} referred to the whole sample, and the random components λ_{0j} and λ_{1j} randomly varying across clusters.

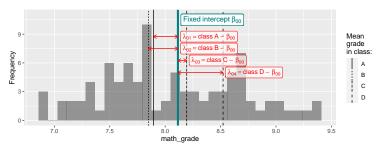
In LMER, x variables (predictors) always variate across clusters j, but not necessarily across individual observations i (e.g., school principals' age only variate across schools, whereas students' age variate across students within schools)

LMER

Random intercept

Let's start with an **intercept-only model** (i.e., *unconditional* or *null model*), where math grades (y_{ij}) are only predicted by the intercept β_{00} and the residuals ϵ_{ij}

- Linear model: $y_i = \beta_0 + \epsilon_i$ The intercept value β_0 is common to all individuals within the population
- Linear mixed-effects model: $y_{ij} = \beta_{0j} + \epsilon_{ij} = (\beta_{00} + \lambda_{0j}) + \epsilon_{ij}$
 - β_{00} is the fixed intercept (also called 'average' or 'general intercept') that applies to the whole population
 - λ_{0j} is the random intercept = cluster-specific deviation from the fixed intercept (i.e., mean class grade fixed intercept)



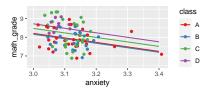
Random slope

Let's now add a predictor: students' anxiety levels x_{ij} .

Random intercept model

$$y_{ij} = \beta_{0j} + \beta_1 x_{ij} + \epsilon_{ij}$$
$$= (\beta_{00} + \lambda_{0j}) + \beta_1 x_{ij} + \epsilon_{ij}$$

Math grades y_{ij} are predicted by the overall mean grade β_{00} , their average relationship with anxiety β_{10} , the random variation among clusters λ_{0j} (random intercept), and the random variation among individuals within clusters ϵ_{ij} (residuals).

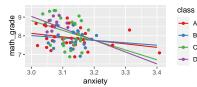


Random intercept & random slope model

$$y_{ij} = \beta_{0j} + \beta_{1j} x_{ij} + \epsilon_{ij}$$

= $(\beta_{00} + \lambda_{0j}) + (\beta_{10} + \lambda_{1j}) x_{ij} + \epsilon_{ij}$

Since the effect of anxiety might not be the same across all classes, we partition β_1 into the overall $average\ relationship$ between anxiety and grades β_{10} ($fixed\ slope$) and the cluster-specific variation in the relationship λ_{1j} ($random\ slope$) - basically, an interaction between anxiety and class.



LMER

From LMER to multilevel modeling

LMER is often called 'multilevel modeling' due to the underlying variance decomposition of the y_{ij} variable into the within-cluster and the between-cluster levels.

That is, the LMER formula $y_{ij} = (\beta_{00} + \lambda_{0j}) + (\beta_{10} + \lambda_{1j}) + \epsilon_{ij}$ can be expressed in two separate levels:

Level 1 (within):
$$y_{ij} = \beta_{0j} + \beta_{1j}x_{ij} + \epsilon_{ij}$$

Level 2 (between): $\beta_{0j} = \beta_{00} + \lambda_{0j}$
 $\beta_{1j} = \beta_{10} + \lambda_{1j}$

b In some papers and textbooks, the coefficients β_{00} and β_{01} are indicated with γ_{00} and γ_{01} , while λ_{0i} and λ_{1i} are sometimes indicated with U_{0i} and U_{1j} , respectively.

That's all for now!

Questions?

LMER

Homework (optional):

- read the slides presented today and write in the Moodle forum if you have any doubts
- refresh your familiarity with **Q**: R-intro.pdf
- exe cises 1-3 from exeRcises.pdf

For each exercise, the solution (or one of the possible solutions) can be found in dedicated chunk of commented code within the exeRcises. Rmd file

The problem

Sometimes the sampling method creates *clusters* of individual observations: **nested data structure** where individuals observations are *nested within* clusters.

$\rightarrow \ Local \ dependencies$

- = correlations among observations within a cluster, violating the LM assumption of independence.
- \rightarrow We cannot use ordinary LM

The solution

Linear mixed-effects regression (LMER) includes additional variance terms¹ to handle local dependencies.

$$y_{ij} = \beta_{0j} + \beta_{1j} x_{ij} + \epsilon_{ij}$$

= $(\beta_{00} + \lambda_{0j}) + (\beta_{10} + \lambda_{1j}) x_{ij} + \epsilon_{ij}$

These can be expressed in two separate levels:

Level 1 (within):
$$y_{ij} = \beta_{0j} + \beta_{1j}x_{ij} + \epsilon_{ij}$$

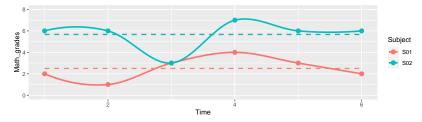
Level 2 (between): $\beta_{0j} = \beta_{00} + \lambda_{0j}$
 $\beta_{1j} = \beta_{10} + \lambda_{1j}$

¹The additional variance terms are the variance τ_{00}^2 of the random intercept λ_{0j} and the variance τ_{10}^2 of the random slope λ_{1j} . We will see this later...

Multilevel modeling in longitudinal designs

Longitudinal assessments (or repeated-measure designs) involve the collection of multiple data from the same subjects at multiple time points.

- \rightarrow Observations from the same subject are not independent (local dependencies).
 - Individual observations = time points (level 1: within-subject)
 - $\bullet \ \ \text{Clusters} = \text{subjects} \ (\textit{level 2: between-subjects})$



b If individuals are further nested within higher-level clusters, we can specify a 3-level model (time points \rightarrow students \rightarrow classes)

Case study: Adolescent insomnia



Contents lists available at ScienceDirect

Sleep Health

Journal of the National Sleep Foundation journal homepage: sleephealthjournal.org

Wearable and mobile technology to characterize daily patterns of sleep, stress, presleep worry, and mood in adolescent insomnia

Luca Menghini, PhD^a, Dilara Yuksel, PhD^b, Devin Prouty, PhD^b, Fiona C. Baker, PhD^{b,c}, Christopher King, PhD^d, Massimiliano de Zambotti, PhD^{bs}





Motion and heart rate continuous passive recording



Bedtime electronic diary ratings of stress, worry, and mood

A sample of 93 US adolescents undertook a semi-structured clinical interview for DSM-5 insomnia symptomatology (insomnia vs. healthy sleepers).

Then, they were provided with a Fitbit wristband (recording **sleep** data) for 2 months. Over the same period, every evening they responded short questionnaires on their **stress** levels at bedtime.

We want to understand whether daily stress predicts lower sleep time (HP1); whether the stress impact on sleep is moderated by insomnia symptomatology (HP2).

Hands on **R**

1. Download & read the datasets from https://github.com/SRI-human-sleep/INSA-home

```
\label{eq:dayNr} \begin{split} \text{ID} &= \text{subject ID, dayNr} = \text{day, stress} = \text{daily stress rating (1-5), TST} = \text{total sleep time (min),} \\ \text{insomnia} &= \text{subject's group (insomnia vs. healthy)} \end{split}
```

```
repo <- "https://github.com/SRI-human-sleep/INSA-home" # loading datasets from GitHub
load(url(paste0(repo,"/raw/main/Appendix%20D%20-%20Data/emaFINAL.RData")))
load(url(paste0(repo,"/raw/main/Appendix%20D%20-%20Data/demosFINAL.RData")))
# selecting columns
ema <- ema[,c("ID","dayNr","stress","TST")] # ema = time-varying variables
demos <- demos[,c("ID","insomnia")] # demos = time-invariant variables</pre>
```

- 2. Print the first rows of the datasets: How many rows per subject?
- 3. Which variable includes individual observations, which is the cluster variable, which is the predictor?
- Which variable(s) at the within-cluster level (Level 1)? Which variable(s) at the between-cluster level (Level 2)

- 5. Explore (descript., correlations, plots)
- Compute the cluster mean for each level-1 variable using aggregate()
- Join the cluster means to the demos dataset using cbind()
- 8. Join the cluster means to the ema dataset using plyr::join()
- 9. Subtract individual obs. from cluster means

Wide & Long data structure

Wide-form dataset

```
one row per cluster
```

```
clustMeans <- # computing cluster means
  aggregate(x = ema[,c("TST","stress")],
  by = list(ema$ID), FUN = mean, na.rm = T)
# join cluster means to the wide-form dataset
demos <- cbind(demos, clustMeans[,2:3])
colnames(demos)[3:4] <- c("TST.m","stress.m")
head(demos)</pre>
```

```
TD insomnia TST.m stress.m

1 s001 0 466.1786 1.707317

2 s002 0 431.0745 2.175000

4 s005 0 415.2059 1.872727

5 s006 1 413.1111 3.393443

6 s007 0 445.7642 1.983333

7 s008 0 422.8468 3.045455
```

Level-2 (between) variables:

ID, insomnia, TST.m, stress.m

Long-form dataset

one row per individual observation

```
ID davNr stress
                      TST insomnia TST.m stress.m
1 s001
           1
                  3 507.0
                                  0.466.2
                                               1.7
2 s001
                  1 502.5
                                  0 466.2
                                               1.7
                  3 469.5
3 s001
                                  0 466.2
                                               1.7
4 s001
                                  0 466.2
                       NΑ
                                               1.7
5 s001
                       NA
                                  0 466.2
                 NΑ
                                               1.7
6 s001
                       NA
                                  0 466.2
                                               1.7
```

Level-1 (within) variables:

dayNr, stress, TST

Between & within cluster

Long-form dataset

one row per individual observation

hea	ad(em	a[,-6]	, 20)			
	ID	dayNr	stress	TST	insomnia	stress.m
1	s001	1	3	507.0	0	1.7
2	s001	2	1	502.5	0	1.7
3	s001	3	3	469.5	0	1.7
4	s001	4	2	NA	0	1.7
5	s001	5	NA	NA	0	1.7
6	s001	6	3	NA	0	1.7
7	s001	7	1	NA	0	1.7
8	s001	8	2	NA	0	1.7
9	s001	9	1	NA	0	1.7
10	s001	10	2	NA	0	1.7
11	s001	11	2	NA	0	1.7
12	s001	12	1	NA	0	1.7
13	s001	13	2	NA	0	1.7
14	s001	14	1	NA	0	1.7
15	s001	15	1	NA	0	1.7
16	s001	16	NA	NA	0	1.7
17	s001	17	NA	NA	0	1.7
18	s001	18	NA	NA	0	1.7
19	s001	19	NA	510.5	0	1.7
20	s001	20	NA	515.5	0	1.7

Long-form data structures are needed to fit multilevel models.

Here, level-1 variables x_{ij} (stress) and y_{ij} (TST) change both between and within cluster.

In contrast, level-2 variables x_j (insomnia, stress.m) only change between clusters, whereas they keep identical values across all the rows associated with the same cluster.

Data centering

Data centering = subtracting the mean of a variable from each variable value.

- The mean of a centered variables is always 0.
- Its variance and covariances are equivalent to those of the original variable.
- Centered scores represent deviations from the mean.

In both LM and LMER, centering the predictors is useful to reduce collinearity (linear relationship between predictors) and for better interpreting a model intercept (= value of y when x is at its mean); but it does not affect the slopes.

```
demos$stress.gmc <- # grand-mean centering
demos$stress.m - mean(demos$stress.m)

Non-centered

Centered

Output
Description
Ou
```

Grand mean vs. Cluster mean centering

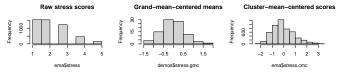
With LMER, we can distinguish two main ways to center the data:

1) Grand mean centering = subtracting the mean of the whole sample (grand-mean or grand-average) from each cluster's mean.

```
# gmc stress = mean cluster's stress - grand mean
demos$stress.gmc <- demos$stress.m - mean(demos$stress.m)</pre>
```

2) Cluster mean centering (or 'group mean centering') = subtracting the mean of the cluster (group mean) from each individual observation nested within that cluster.

```
# cmc stress = individual obs. - mean of the corresponding cluster
ema$stress.cmc <- ema$stress.m</pre>
```



Hands on **Q**: Compute the grand-mean-centered & the cluster-mean-centered values of stress and TST. Then, compute their Pearson's correlation with the cor() function

That's all for now!

Questions?

Homework (optional):

- read the slides presented today and write in the Moodle forum if you have any doubts
- exe cises 4-5 from exeRcises.pdf

For each exercise, the solution (or one of the possible solutions) can be found in dedicated chunk of commented code within the exercises.Rmd file

In the last episodes...

Problem & solution

The sampling method can create clusters of individual observations = $nested\ data$ leading to $local\ dependencies$

→ Multilevel modeling (or LMER) includes additional variance terms to handle local dependencies.

Level 1 (within):
$$y_{ij} = \beta_{0j} + \beta_{1j}x_{ij} + \epsilon_{ij}$$

Level 2 (between): $\beta_{0j} = \beta_{00} + \lambda_{0j}$
 $\beta_{1j} = \beta_{10} + \lambda_{1j}$

Wide and long datasets

LMER require **long-form datasets**, with one row per each individual observation (level 1) and multiple rows for each cluster (level 2)

Between and within

In such datasets, within-cluster (level-1) variables variate both between and within clusters, while between-cluster (level-2) variables only variate across clusters, keeping identical values across the rows belonging to the same cluster.

Data centering

& Variance decomposition

Data centering (= subtracting the mean from each variable value) can be used to decompose the variance into:

- the between-cluster component = grand-mean-centered means
- the within-cluster component
 = cluster-mean-centered values

The adolescent insomnia case study

A sample of 93 US adolescents undertook a semi-structured clinical interview for DSM-5 insomnia symptomatology (insomnia vs. healthy sleepers).

Then, they were provided with a Fitbit wristband (recording sleep data) for 2 months. Over the same period, every evening they rated their stress (1-5) at bedtime.

We want to test whether day-to-day fluctuations in stress predict lower total sleep time TST (HP1), and whether the stress impact on TST is moderated by insomnia symptomatology (HP2).

```
load("insa.RData") # read processed data
insa[,c("ID","TST","TST.m","TST.gmc","TST.cmc")]
         TST TST.m TST.gmc TST.cmc
1 s001 507.0 466.18
                      53.73
                             40.82
2 s001 502.5 466.18
                      53.73
                             36.32
3 s001 469.5 466.18
                      53.73
                             3.32
21 s001 496.0 466.18
                      53.73
                             29.82
22 s001 447.5 466.18
                      53.73 -18.68
23 s001 450.5 466.18
                      53.73
                            -15.68
24 s001 423.0 466.18
                      53.73 -43.18
29 s001 483.5 466.18
                      53.73 17.32
30 s001 450.0 466.18
                      53.73
                            -16.18
31 s001 529.0 466.18
                      53.73
                             62.82
TST = raw total sleep time (minutes)
```

TST.gmc = grand-mean-centered cluster means
of TST (level-2 component)

TST.cmc = cluster-mean-centered TST (level-1 component)

Descriptive statistics of multilevel data

The first section of the results section in any quantitative report (including published papers) includes the **descriptive statistics** of the considered variables in the examined sample. Descriptive statistics are also the main output of any quantitative report you might draft or read in your **professional practice**.

With mutlilevel datasets, the descriptive statistics to be reported are the following:

- 1. **Mean and SD** of any considered quantitative variable
- 2. Frequency (%) of any considered categorical variable
- 3. Level-specific correlations among quantitative variables
- 4. Intraclass correlation coefficient (ICC) of any quantitative variable measured at the *within-cluster* level
- © Compute descriptive statistics 1-3, considering the variables TST, stress, and insomnia (*Note*: correlations can be computed with the cor() function; level-2 correlations should be computed on the cluster means in the demos dataset)
- **b** Response rate (or missing data) is a further important descriptive to report. Here, for simplicity, we omitted missing data points from the insa dataset.

Level-specific correlations

Between-cluster (level 2)

Cluster means

Level-2 correlation

= linear relationship across clusters

Do stressed subjects sleep worse than unstressed subjects?

```
wide <- insa[!duplicated(insa$ID),]
cor(wide[,c("stress.m", "TST.m")])</pre>
```

stress.m TST.m stress.m 1.000 -0.067 TST.m -0.067 1.000

Within-cluster (level 1)

Individual deviations from cluster mean = cluster-mean-centered values

Level-1 correlation

= linear relationship within cluster

Do subjects sleep worse than usual in those days where they are more stressed than usual?

```
cor(insa[,c("stress.cmc", "TST.cmc")])
```

```
stress.cmc TST.cmc
stress.cmc 1.00 -0.06
TST.cmc -0.06 1.00
```

Additional variance (& covariance) terms

LMER includes additional variance and covariance terms to handle local dependencies. \rightarrow Variance and covariance what?!

Rembember the LMER formula:

$$y_{ij} = (\beta_{00} + \lambda_{0j}) + (\beta_{10} + \lambda_{1j})x_{ij} + \epsilon_{ij}$$

 λ_{0j} are the random deviations of cluster $intercepts \ {\it from the} \ fixed \ intercept \ \beta_{00}$

 λ_{1j} are the random deviations of cluster slopes from the fixed slope β_{10}

 ϵ_{ij} is the **residual term** indicating the random deviations of *observed values* from *predicted values* (see slide #8) In both LM and LMER, we don't report each single residual value ϵ_{ij} , but we use $\sigma^2 = \text{variance of the residuals } \epsilon$

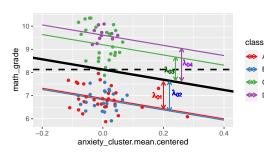
Similarly, in LMER we summarize the random effects by reporting their variances:

 $au_{00}^2 = ext{variance of random intercept } \lambda_{0j}$ $au_{11}^2 = ext{variance of random slope } \lambda_{1j}$

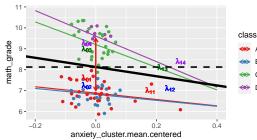
Moreover, when both λ_{0j} and λ_{1j} are included, we need to also consider the covariance term: $\rho_{01} = \text{covariance between } \lambda_{0j} \text{ and } \lambda_{1j}$

 $o au_{00}^2, au_{11}^2, hinspace
ho_{01}$ are the additional variance & covariance terms included in LMER

Random intercept and random slope (1/2)



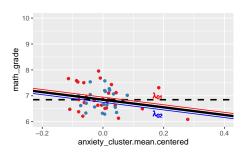
Random intercept (RI) $y_{ij} = (\beta_{00} + \lambda_{0j}) + \beta_1 x_{ij} + \epsilon_{ij}$ RI = distances between each cluster's intercept and the fixed intercept Parallel lines: there is no random slope $\tau_{00}^2 = \text{variance of the RI (how much the RI differ among each other)}$

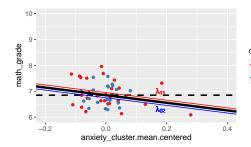


• RI and random slope (RS) $y_{ij} = (\beta_{00} + \lambda_{0j}) + (\beta_{10} + \lambda_{1j})x_{ij} + \epsilon_{ij}$ RS = distances between each cluster's slope and the fixed slope $\tau_{00}^2 = \text{variance of the RI} = 2.22$ $\tau_{10}^2 = \text{variance of the RS}$ $= \text{var}(\lambda_{11}, \lambda_{12}, \lambda_{13}, \lambda_{14}) = 6.27$ $\rho_{01} = \text{covariance between } \lambda_{0j} \& \lambda_{1j}$

 $= var(\lambda_{01}, \lambda_{02}, \lambda_{03}, \lambda_{04}) = 2.22$

Random intercept & random slope (2/2)





What happens if we remove class C and D? \rightarrow Both random effects become smaller class lower variance τ_{00} and τ_{10}

• A

Random intercept (RI) $y_{ij} = (\beta_{00} + \lambda_{0j}) + \beta_1 x_{ij} + \epsilon_{ij}$

Class A and class B's intercepts are very close, their distances from the **fixed intercept** are very small

 $u_{ii} = (\beta_{00} + \lambda_{0i}) + (\beta_{10} + \lambda_{1i})x_{ii} + \epsilon_{ii}$

$$\lambda_{01} \sim \lambda_{02} \rightarrow \tau_{00}^2 \sim 0$$

• RI and random slope (RS)

class Class A and class B's slopes are very close \rightarrow their distances from the **fixed**

slope are very small
$$\lambda_{11} \sim \lambda_{12} \rightarrow \tau_{11}^2 \sim 0$$

Conclusions: It makes no sense to use LMER (better using LM!)

Null model & variance decomposition (1/2)

A null model only includes the intercept and residual terms (see slide #20).

In LM null models $(y_i = \beta_0 + \epsilon_i)$

the intercept β_0 is simply the mean of y_i , and the variance of ϵ_i (σ^2) is simply the variance of y_i .

In LMER null models $(y_{ij} = \beta_{00} + \lambda_{0j} + \epsilon ij)$

the y variance is decomposed into:

- the variance σ^2 of the residuals ϵ_{ij} across both levels
- the between-cluster (level-2) variance $au_{00}^2 = \text{variance of the random intercept } \lambda_{0j}$

Null model & variance decomposition (2/2)

Spoiler alert: How to fit LMER in R

```
# fitting a null LMER model
library(lme4)
m0 <- lmer(TST ~ (1 | ID), data = insa)
summary(m0)
Linear mixed model fit by REML ['lmerMod']
Formula: TST ~ (1 | ID)</pre>
```

Data: insa

REML criterion at convergence: 49553.2

Scaled residuals:

Min 1Q Median 3Q Max -3.4233 -0.6134 -0.0285 0.5760 5.6047

Random effects:

 Groups
 Name
 Variance
 Std.Dev.

 ID
 (Intercept)
 1183
 34.39

 Residual
 5158
 71.82

 Number of
 obs: 4333, groups:
 ID, 93

Fixed effects:

Estimate Std. Error t value (Intercept) 410.838 3.769 109

If we inspect the summary of a null LMER model, starting from the bottom, we can see that:

- Fixed effects only include the fixed intercept β_{00} (= 410.838 minutes).
- Random effects include variance & SD of the random intercept λ_{0j} ($\tau_{00}^2 = 1183$) and that of the residuals ϵ_{ij} ($\sigma^2 = 5158$).

The sum $\sigma^2 + \tau_{00}^2$ of the residual (level-1) and the random intercept variance (level-2) is the model estimate of the population-level total variance in y_{ij}

▶ Variance decomposition & Data centering

The variance decomposition implemented by LMER is basically equivalent to the data centering procedures shown in the last lecture (see slide #32).

```
# random intercept LAMBDA Oj
                                                       # arand-mean-centered TST cluster means
round(head( ranef(m0)$ID[[1]] ).1)
                                                       round(head( wide$TST.gmc ).1)
[1] 50.0 6.2 4.7 4.1 31.1 7.9
                                                       [1] 53.7 18.6 2.8 0.7 33.3 10.4
# random intercept variance TAU^2
                                                       # variance of TST cluster means
(tau2 <- round(summarv(m0)$varcor$ID[[1]]))</pre>
                                                       var(wide$TST.m)
[1] 1183
                                                       [1] 1241.19
# residual variance SIGMA^2
                                                       # variance of cluster-mean-centered TST
(sigma2 <- summarv(m0)$sigma^2)
                                                       var(insa$TST.cmc, na.rm=TRUE)
[1] 5157.676
                                                       [1] 5072.426
                                                       # observed total variance in TST
# estimated total variance in TST
tau2 + sigma2
                                                       var(insa$TST, na.rm=TRUE)
[1] 6340.676
                                                       [1] 6291.752
```

■ The small differences between model-based (on the left) and observed values (on the right) are due to slight adjustments (e.g., accounting for the number of clusters) used by LMER models (for details, see Finch & Bolin, 2014, chapter 2)

Intraclass correlation coefficient (ICC)

The last 'descriptive' statistics to be reported is the ICC

- = Proportion of between-cluster variance over the total variance The ICC is estimated from the null model as $ICC = \tau_{00}^2/(\tau_{00}^2 + \sigma^2)$ and can range between 0 and 1.
 - ICC = 1: the variable only varies across clusters ('cluster-only variable')
 - 0.50 < ICC < 1: the variable mainly varies across clusters
 - ICC = 0.50: the variable equally varies across & within clusters
 - 0 < ICC < 0.50: the variable mainly varies within clusters*
 - ICC = 0: the variable only varies within cluster ('individual-only variable')

The ICC is important in multilevel modeling, because it indicates the degree to which the nested data structure may impact a level-1 variable \rightarrow it indexes of the local dependencies implied by the nested data structure.

Descriptive statistics of multilevel data

Now we have all the core descriptive statistics! ©

Variable	Mean~(SD)/Freq.~(Prop.)	ICC	1.	2.
1. TST (minutes)	413.69 (79.32)	0.19	1.00	-0.06
2. Stress (1 - 5)	2.21 (1.06)	0.26	-0.07	1.00
3. Insomnia group	47 (50.54%)	NA	NA	NA

Note: lv-1 and lv-2 correlations are shown below and above the main diagonal, respectively. In this case, the two variable are not so correlated at any level Θ

- Download and read the file studentData.csv
- DESC: Compute the mean and SD of anxiety and math_grade; compute the number of students per classID
- Compute the cluster mean for anxiety using aggregate() → wide-form
- 4. Join the cluster means to the long-form: plyr::join(long,wide,by="cluster")
- Compute the cluster-mean-centered values of anxiety
- 6. Repeat points 4-5 for math_grade

- DESC: Compute the between-cluster (lv2) correlation from the wide-form dataset (1 row per cluster)
- DESC: Compute the within-cluster (lv1) correlation from the long-form dataset (1 row per individual obs.)
- Fit a null multilevel model with the lme4 package:

```
m0 <- lmer(y ~ (1|cluster), data) and get \sigma^2: summary(m0)$sigma^2 and \tau^2_{00}: summary(m0)$varcor$ID[[1]]
```

10. DESC: Compute and interpret the ICC $= \tau_{00}^2/(\tau_{00}^2 + \sigma^2)$

That's all for now!

Questions?

Homework (optional):

- read the slides presented today and write in the Moodle forum if you have any doubts
- exe cises 6-7 from exeRcises.pdf

For each exercise, the solution (or one of the possible solutions) can be found in dedicated chunk of commented code within the exercises.Rmd file

In the last episodes...

Problem & solution

The sampling method can create clusters of individual observations = $nested\ data$ leading to $local\ dependencies$

→ Multilevel modeling (or LMER)

 $includes\ additional\ variance\ (and$ $covarariance)\ terms\ for\ local\ dependencies.$

Level 1 (within):
$$y_{ij} = \beta_{0j} + \beta_{1j}x_{ij} + \epsilon_{ij}$$

Level 2 (between): $\beta_{0j} = \beta_{00} + \lambda_{0j}$
 $\beta_{1j} = \beta_{10} + \lambda_{1j}$

Wide and long datasets

LMER require **long-form datasets**, with one row per each individual observation (level 1) and multiple rows for each cluster (level 2)

Variance decomposition

LMER automatically decompose the Y variance into its within-cluster (lv1) and between-cluster (lv2) components.

Similarly, we can use data centering to better express predictors (X variables) at level 1 (cluster mean centering) or at level 2 (cluster means).

Descriptive statistics

- Mean (SD) / Freq. of any variable
- Level-specific correlations
- $ICC = \tau_{00}^2/(\tau_{00}^2 + \sigma^2)$

indexing the proportion of level-2 variance, where τ_{00}^2 is the variance of the random intercept β_{00} (lv2) and σ^2 is the variance of the residuals ϵ_{ij} (lv1) from a null model

Fitting multilevel models (in R): Null model

We will use the lme4 package (Bates et al 2014), which uses the lmer() function to fit linear models the exact same way of lm() (i.e., formula & data arguments).

```
library(lme4) # loading package
```

Ordinary linear model (LM)

TST is predicted by the **intercept** β_0 (expected value of TST in the sample = grand average) & the **residual variance** σ^2 , without accounting for local dependencies and the multilevel data structure.

Multilevel model (LMER)

TST is predicted by the fixed intercept β_{00} (lv2), the variance of the random intercept τ_{00}^2 (lv2), & the residual variance σ^2 (lv1).

(Intercept) 410.8383

summary(lmer0)\$varcor\$ID[[1]] # RI variance
[1] 1182.746

summary(lmer0)\$sigma^2 # residual variance

[1] 5157.676

[1] 1186.171

[1] 5137.951

Random intercept (RI) model

A RI model can include 1+ predictors, but their effect does not variate across clusters.

Ordinary linear model (LM)

TST is predicted by the **intercept** β_0 (expected value when stress.cmc = 0),

the slope β_1 (indexing the predicted change in TST for a 1-unit increase in stress.cmc), and the residual variance σ^2 .

coefficients(lm1) # intercept & slope

(Intercept) stress.cmc 413.701214 -4.762748

```
summary(lm0)$sigma^2 # residual variance
```

[1] 6291.752

Multilevel model (LMER)

TST is predicted by the fixed intercept β_{00} (lv2), the variance of the RI τ_{00}^2 (lv2), the slope β_1 (same meaning than in LM), & the residual variance σ^2 (lv1).

summary(lmer1)\$sigma^2 # residual variance

Random slope (RS) model

In a **RS model** the effect of 1+ level-1 predictors randomly varies across clusters.

Random intercept (RI) model

The within-individual effect of stress on TST is fixed across clusters. The model only includes a fixed slope β_1 indexing the overall relationship between the two variables.

```
lmer1 <-
  lmer(TST ~ stress.cmc + (1 ID),
       data = insa)
fixef(lmer1) # fixed effects
(Intercept)
            stress.cmc
 410.848597
             -4.920536
summary(lmer1)$varcor$ID[[1]] # RI var
[1] 1186.171
summary(lmer1)$sigma^2 # residual var
```

[1] 5137.951

Random slope (RS) model

The effect of stress varies across clusters. The model also includes the RS variance τ_{10}^2

and the **covariance** ρ_{01} between RI and RS.

```
1mer2 <-
  lmer(TST ~ stress.cmc + (stress.cmc ID).
       data = insa)
fixef(lmer2) # fixed effects
(Intercept)
            stress.cmc
 410 909025
             -5.685554
# RI variance, RS variance, RI-RS covariance
matrix(summary(lmer2)$varcor$ID)[c(1,4,2),]
[1] 1183.70745
                87.26116
                           21.22170
```

summary(lmer2)\$sigma^2 # residual variance

[1] 5071.189

```
From the previous examples, we saw that lmer() includes an additional term using the syntax (1 | cluster_variable), standing for the random intercept:

lmer(formula = TST ~ stress.cmc + (1 | ID), data = insa)
```

If we replace the value 1 in the first term between brackets with the name of a level-1 predictor included in the model, we get (predictor | cluster_variable), standing for the random intercept and the random slope:

```
lmer(formula = TST ~ stress.cmc + (stress.cmc | ID), data = insa)
```

```
It is also possible to add further level-1 and level-2 predictors (multiple\ regression) lmer(TST ~ stress.cmc + x2 + x3 + x4 + ... + (stress.cmc | ID), data = insa) ...and their interactions:
```

```
lmer(TST ~ stress.cmc + x2 + x2:stress.cmc + (stress.cmc | ID), data = insa)
```



Download & read the pre-processed dataset insa. RData (omitting missing data)

```
TST = total sleep time (min), stress.cmc = cluster-mean-centered stress (1-5), insomnia = insomnia group, ID = participant identifier getwd() # get where your working directory is, and save the data file in it load("insa.RData") # read data
```

- 2. Mean, SD, correlations & plots
- Fit a null LMER model m0 of TST and compute the ICC
- Fit a model m1 with TST being predicted by stress.cmc
- Fit a model m2 with a random slope for stress.cmc
- Inspect the summary() of each model:
 Is there a substantial within-individual relationship between TST and stress
 (hupothesis 1)

- 7. Fit a model m3 that also includes insomnia group differences: Any group differences? Does it change the effect of stress?
- Fit a model m4 that also includes the interaction between insomnia and stress.cmc
- Inspect the summary() of of model m4:
 Does insomnia moderate the
 within-individual relationship between
 stress and TST? (hypothesis 2)

lmer() model summary

Here we print and comment the summary of the interactive model m4.

```
m4 <- lmer(TST ~ stress.cmc * insomnia + (stress.cmc | ID), data = insa)
```

summary(m4)

Linear mixed model fit by REML ['lmerMod']

Formula: TST ~ stress.cmc * insomnia + (stress.cmc | ID)
Data: insa

REML criterion at convergence: 49511.7

Scaled residuals:

Min 1Q Median 3Q Max -3.4787 -0.6086 -0.0211 0.5756 5.5474

Random effects:

Groups Name Variance Std.Dev. Corr
ID (Intercept) 1196.32 34.588
stress.cmc 86.44 9.297 0.06
Residual 5071.75 71.216
Number of obs: 4333, groups: ID, 93

Fixed effects:

TIACU CIICCOD.				
	Estimate	Std.	Error	t value
(Intercept)	409.505		5.395	75.900
stress.cmc	-7.187		2.290	-3.138
insomnia1	2.759		7.572	0.364
stress.cmc:insomnia1	2.923		3.188	0.917

- First lines: model formula, data, and parameter estimation method (here, REML), info on estimation convergence
- Scaled residuals: descriptives of the model residuals
- Random effects: estimated variance
 (τ₀₀², τ₁₀²), SD (τ₀₀, τ₁₀), and correlation
 (ρ₁₀) of random intercept and random
 slope, residual variance (σ²) and SD (σ)
- Number of individual observations (lv1) and clusters (lv2) used by the model
- Fixed effects: fixed intercept and fixed slope for stress, insomnia, and their interaction (i.e., product)

LMER coefficient interpretation

Here, we interpret the fixed coefficients estimated by model m4.

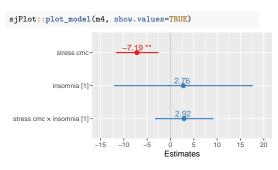
round(summary(m4)\$coefficients, 1) # fixed effects part of the summary

	Estimate	Std.	Error	t	value
(Intercept)	409.5		5.4		75.9
stress.cmc	-7.2		2.3		-3.1
insomnia1	2.8		7.6		0.4
stress.cmc:insomnia1	2.9		3.2		0.9

- Fixed intercept: the predicted value of TST when stress.cmc = 0 (average stress level) and insomnia = 0 (controls = reference group) is 409.5 minutes.
- Fixed stress slope: when insomnia = 0 (controls), TST is predicted to decrease
 by -7.2 minutes for each 1-point increase in stress.cmc (more stressed than usual).
- Fixed insomnia slope: when stress.cmc = 0 (average stress), the insomnia is expected to show an average TST of 2.8 minutes higher than the control group.
- Interaction: when insomnia = 1, the stress-related decrease in TST is predicted to be reduced by 2.9 minutes (i.e., -7.2 + 2.9 = -4.3 minutes per 1-unit increase in stress).
- t values (= Estimate/Std.Error) suggest that stress.cmc (higher stress than usual) predicts lower TST (|t| > 1.96), but their relationship does not change across the insomnia and the control group (|t| < 1.96) → HP1 supported, HP2 not supported

Visualizing fixed estimates & standard errors

Forest plot: The plot_model() function of the sjPlot package allows visualizing fixed estimates (dots) with their 95% confidence intervals (CI) = Estimate ± 1.96 Std.Err. indexing the precision of the estimate value (line limits).



Interpretation:

- Consistently with the previous slide, the only 95% CI excluding zero are those of stress.cmc (in line with HP1 but not HP2).
- The insomnia estimate (lv2) varies more than that of stress (lv1) - also due to the *lower sample size at the* between-cluster level

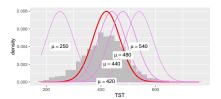
Both 95% CI and the t-value are derived from the standard error (SE) = predicted variability in the estimate if the data were collected from different random samples.



▶ Parameter estimation in LMER

LMER coefficients and SE can be estimated with various methods (or algorithms), including the Bayesian estimator (see slide #7), but the most used are MLE and REML.

Maximum Likelihood Estimation (MLE) Finds the combination of parameter values that maximize the likelihood function (= probability of observing our data given the model) using an iterative approach (the model is repeatedly fitted with different parameter values until the maximum is identified).



Restricted Maximum Likelihood (REML) Similar to MLE, but estimates the variance components in a different way:

- MLE firstly estimates the mean μ and then the variance (as the distance from μ), but this was found to underestimate the variance
- REML applies a correction based on the number of fixed coefficients to get

less biased variance estimates

Since variance components are critical in LMER (random effects), REML is generally preferred (default in R), but with large sample they are basically the same.

Visualizing fixed and random effects

stress.cmc

The plot_model() function also allows to visualize fixed and random effects.

Random effects Fixed effects Regression line & 95% CI 📤 Estimate & 95% CL plot_model(m4, type = "pred") # main effects plot_model(m4, type = "re") Predicted values of TST Predicted values of TST Random effects 450 -420 -ID (Intercept) 425 415 s041 -E 400 E 410 s040 s039 -405 s038 -375 s037 -400 s035 s034 s033 stress.cmc insomnia s031 s030 s029 plot_model(m4, type = "int") # interaction s028 s027 s026 -Predicted values of TST s025 s024 -450 s023 s022 -425 insomnia s021 s019 -LS 400 s018 s017 s016 -375 s013 s012 s011 s008 stress.cmc s007 -

LMER results in a scientific paper/report

While the output of summary() is quite exhaustive, it slightly differs from what typically reported in scientific papers/reports. The tab_model() from sjPlot provides such a format.

You should now be able to understand the meaning of any reported value.

ightharpoonup sjPlot calls random effect variances au rather than au^2 .

tab_model(m4, show.se=TRUE, collapse.se=TRUE, string.est="b (SE)")

Predictors	b (SE)	CI	P
(Intercept	409.50 (5.40)	398.93 - 420.08	< 0.001
Stress	-7.19 (2.29)	-11.682.70	0.002
Group [Insomnia]	2.76 (7.57)	-12.09 - 17.60	0.716
$Stress \times Group \; [Insomnia]$	2.92(3.19)	-3.33 - 9.17	0.359
Random Effects			
σ^2	5071.7	75	
$ au_{00}$ ID	1196.3	32	
$ au_{11}$ ID.stress.cmc	86.44		
$ ho_{01}$ ID	0.06		
N _{ID}	93		
Observations	4333		

That's all for now!

Questions?

Homework (optional):

- read the slides presented today and write in the Moodle forum if you have any doubts
- exe cises 8-9 from exeRcises.pdf

For each exercise, the solution (or one of the possible solutions) can be found in dedicated chunk of commented code within the exercises.Rmd file

Problem & solution

The sampling method can create *nested* data structures (obs. within clusters).

 $\label{local_local} \textbf{LMER} \ \text{includes} \ additional \ (co) variance \\ terms \ \text{to handle local dependencies}.$

Level 1 (within):
$$y_{ij} = \beta_{0j} + \beta_{1j}x_{ij} + \epsilon_{ij}$$

Level 2 (between): $\beta_{0j} = \beta_{00} + \lambda_{0j}$
 $\beta_{1j} = \beta_{10} + \lambda_{1j}$

Variance decomposition

Based on long-form datasets, LMER decompose the Y variance into within-cluster & between-cluster. The same can be done by cluster-mean-centering the predictors X.

LMER descriptives

Mean (SD) / Freq. of any variable; Level-specific correlations; $ICC = \tau_{00}^2/(\tau_{00}^2 + \sigma^2)$

LMER model fit & output

```
lmer(Yij ~ (1|cluster), data) # null RI model
lmer(Yij ~ Xij + (1|cluster), data) # RI
lmer(Yij ~ Xij + (Xij|cluster), data) # RS
```

summary(fit)\$coefficients # fixed effects

	Estimate	Std.	Error	t value
(Intercept)	409.50		5.40	75.90
stress.cmc	-7.19		2.29	-3.14
insomnia1	2.76		7.57	0.36
stress.cmc:insomnia1	2.92		3.19	0.92

Random effect variances $(\tau_{00}^2, \tau_{10}^2, \sigma^2)$:

```
summary(fit)$varcor[[1]][c(1,4,2)]
```

[1] 1196.32 86.44 18.29

Reading the Results section of a paper (pt1)

Based on what we saw in the previous lectures, you should now be able to understand the results of scientific papers/reports reporting on multilevel analyses.

Try answering the following questions by looking at the results of the linked papers.

Note: Similar questions will be included in the final exam.

- 1. Which variable identifies individual observations and which is the cluster variable?
- Which predictors are at level 1 (within-cluster)? Which at level 2 (between-cluster)?
- 3. Do the authors report the random effects? Which ones?
- 4. Does the model include 1+ random slopes? For which predictor(s)?
- Do the authors report estimate SE, t-value, 95% CI?

- Graham et al (2020): Neighborhood disadvantage & children's sleep health (Table 3)
 - DOI: 10.1016/j.sleh.2020.05.002
- Ersan & Rodriguez (2020): Socioeconomic status & math achievement (**Table 5**)
 - DOI: 10.1186/s40536-020-00093-y
- Juvrud et al (2021): Infants' attention, maternal affect, & emotional context (Supplementary Table 2)
 - DOI: 10.3389/fpsyg.2021.700272

Multilevel model evaluation

With 'model evaluation' we refer to two main procedures:

- Model diagnostics: Evaluating whether the model fits the data consistently
 with the underlying model assumptions (e.g., see LM assumptions in slide #11)
- Model comparison: Evaluating whether the model fits substantially better or
 worse than alternative models (e.g., see LM model comparison in slide #6)
 → model selection (choosing the best model)
- Data analysis pipeline
 - 1. Data exploration & descriptives
 - 2. Model fit
 - 3. Model diagnostics
 - 4. Model comparison
 - 5. Model selection & coefficient interpretation
 - 6. Result visualization

LMER assumptions

Similar to LM, LMER models require that some assumptions about the data hold true. Otherwise, we cannot trust the estimated parameters or any other result.

Assumptions common to LM:

- 1. Linearity: x_i and y_i are linearly associated \rightarrow expected (mean) value of ϵ_{ij} is zero
- 2. Normality: residuals ϵ_{ij} are normally distributed $\rightarrow \epsilon_{ij} \sim \mathcal{N}(0, \sigma^2)$
- 3. Homoscedasticity: ϵ_{ij} variance is constant over the levels of x_i (homogeneity of variance)
- 4. Independence: predictors x_{ij} and x_j are unrelated to residuals ϵ_{ij}
- ${\bf 5. \ Absence \ of \ influential \ observations} \ ({\rm multivariate \ outliers})$
- $\textbf{6. Absence of multicollinearity}: \ no \ linear \ relationship \ between \ different \ predictors$

Additional LMER assumptions:

- 7. Linearity, Normality, Homoscedasticity, & Independence of random effects:
- In LMER, assumptions 1-4 also apply to 'cluster-level residuals' (i.e., random effects).

Random intercept λ_{0j} and random slope λ_{1j} should be normally distributed with

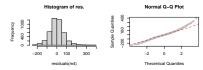
 $\lambda_{0j} \sim \mathcal{N}(0, \tau_{00}^2)$ and $\lambda_{1j} \sim \mathcal{N}(0, \tau_{11}^2)$, their variance should be homogeneous across the levels of x variables, and they should be independent from predictors

LMER diagnostics: Residuals (lv1)

Let's evaluate whether model m4 (adolescent insomnia) meets LMER assumptions.

Normality & linearity: symmetric histogram centered on 0, straight normal QQ plot ©

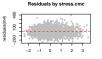
hist(residuals(m4))
qqnorm(residuals(m4)); qqline(residuals(m4))



${\bf Homoscedasticity} \ \& \ independence:$

 $\textit{\textbf{no}}$ trends in ϵ_{ij} or their variance over x Θ

plot(residuals(m4) ~ insa\$stress.cmc)
plot(residuals(m4) ~ insa\$insomnia)

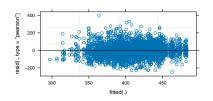




A faster way to evaluate assumptions 1-4 is to plot residuals vs. predicted values ('summary' of predictor information): the points (residuals) should be evenly divided above & below (normality) their mean value of zero (linearity), with no strong trends (independence & homoscedasticity)

(independence & homoscedasticity)





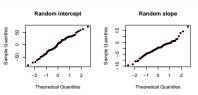
LMER diagnostics: Random effects (lv2)

Random effects can be extracted using the function ranef (model_name), which returns a dataset with 2 columns (RI & RS) and a number of rows = number of clusters (lv2).

```
# from long to wide: 1 row per subject
wide <- insa[!duplicated(insa$ID),]
# extract random effects
RI <- ranef(m4)[[1]][,1] # r. intercept
RS <- ranef(m4)[[1]][,2] # r. slope</pre>
```

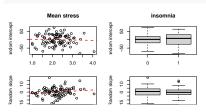
Normality & linearity: Straight QQ ©

qqnorm(RI); qqline(RI)
qqnorm(RS); qqline(RS)



Homoscedasticity & independence: No marked trends in random effects or their variance ©, but slightly higher RI var. and lower RS var. in insomnia than in controls ©

```
plot(RI ~ wide$stress.m) # RI
plot(RI ~ wide$insomnia)
plot(RS ~ wide$stress.m) # RS
plot(RS ~ wide$insomnia)
```



LMER diagnostics: Multicollinearity & influential cases

With both LM & LMER, we need to avoid using too correlated predictors (multicollinearity), otherwise they will 'steal' each other's explained variance.

ightarrow Variance inflation factors (VIF) tell us how much the standard errors are increased due to multicollinearity $lap{1}{2}$ $VIF=1/(1-R_{x_i}^2)$

Influential cases are data points that substantially change (influence) one or more parameter estimates (multivariate outliers). With LMER, influential cases can be at lv1 or at lv2 (clusters).

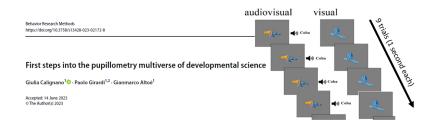
→ The Cook's distance (CD) tells us how much the parameter estimates change after the exclusion of each case. If too extreme, we remove that case and check again.

```
VIF > 5 = highly correlated x; here ok \odot
car::vif(m4)
stress 2.07 : insomnia 1 : inter. 2.07 :
barplot(car::vif(m4)); abline(h=5)
                   Variance inflation factors
   ^{\circ}
          stress cmc
Extreme CD for 1 obs. & 2 clusters 

boxplot(cooks.distance(m4)) # lv1
library(influence.ME) # lv2
plot(influence(m4, group="ID"), which="cook")
 0.2 -
 0.0 -
-0.2 -
-0.4 -
                                  Lv-2 Cook's distance
```

I v-1 Cook's distance

Case study: Infants' pupil dilation •



A sample of 16 12-month-olds undertook 2 blocks of familiarization task with 9 one-sec trials each. In each block, they were familiarized with a novel visual object presented on a screen either with (audiovisual) or without (visual) an auditory label (e.g., "coba").

Eye tracking was used to record their **pupil dilation** (in millimeters) over the experiment, as a measure of infant online processing & attention deployment.

We want to test whether **pupil dilation is lower in the audiovisual trials** (HP1), since the auditory label is expected to improve familiarization and require less processing efforts.

1. Download & read the dataset from the Pupillometry multiverse study

id = infant's ID, fam = familiar. type (labeled vs. unlabeled), pupil = pupil dilation (mm)

library(osfr) # package to interact with the OSF platform

proj <- "https://osf.io/p8nfh/" # link to the OSF project (see protocol paper & data dictionary)

osf_download(osf_ls_files(osf_retrieve_node(proj))[5,],conflicts="overwrite") # download

infants <- read.csv2("data/multiverse.csv",stringsAsFactors=TRUE) # read dataset

colnames(infants)[c(18,17,19)] <- c("id","fam","pupil") # shortening variable names

Explore the the variables id, fam, and pupil (descriptives & correlations)

infants\$pupil <- as.numeric(infants\$pupil) # pupil as numeric

1. Download & read the dataset from the $Pupillometry\ multiverse$ study

```
id = infant's ID, fam = familiar. type (labeled vs. unlabeled), pupil = pupil dilation (mm)
library(osfr) # package to interact with the OSF platform
proj <- "https://osf.io/p8nfh/" # link to the OSF project (see protocol paper & data dictionary)
osf_download(osf_ls_files(osf_retrieve_node(proj))[5,],conflicts="overwrite") # download
infants <- read.csv2("data/multiverse.csv",stringsAsFactors=TRUE) # read dataset
colnames(infants)[c(18,17,19)] <- c("id","fam","pupil") # shortening variable names
infants$pupil <- as.numeric(infants$pupil) # pupil as numeric</pre>
```

- Explore the the variables id, fam, and pupil (descriptives & correlations)
- 3. Which variable identifies individual observations and which is the cluster variable? How many clusters?

1. Download & read the dataset from the *Pupillometry multiverse* study

```
id = infant's ID, fam = familiar. type (labeled vs. unlabeled), pupil = pupil dilation (mm)
library(osfr) # package to interact with the OSF platform
proj <- "https://osf.io/p8nfh/" # link to the OSF project (see protocol paper & data dictionary)
osf_download(osf_ls_files(osf_retrieve_node(proj))[5,],conflicts="overwrite") # download
infants <- read.csv2("data/multiverse.csv",stringsAsFactors=TRUE) # read dataset
colnames(infants)[c(18,17,19)] <- c("id","fam","pupil") # shortening variable names
infants$pupil <- as.numeric(infants$pupil) # pupil as numeric</pre>
```

- Explore the the variables id, fam, and pupil (descriptives & correlations)
- 3. Which variable identifies individual observations and which is the cluster variable? How many clusters?
- 4. Which predictor(s) at lv1, which at lv2?

```
1. Download & read the dataset from the Pupillometry multiverse study
```

```
id = infant's ID, fam = familiar. type (labeled vs. unlabeled), pupil = pupil dilation (mm)
library(osfr) # package to interact with the OSF platform
proj <- "https://osf.io/p8nfh/" # link to the OSF project (see protocol paper & data dictionary)
osf_download(osf_ls_files(osf_retrieve_node(proj))[5,],conflicts="overwrite") # download
infants <- read.csv2("data/multiverse.csv",stringsAsFactors=TRUE) # read dataset
colnames(infants)[c(18,17,19)] <- c("id","fam","pupil") # shortening variable names
infants$pupil <- as.numeric(infants$pupil) # pupil as numeric</pre>
```

- Explore the the variables id, fam, and pupil (descriptives & correlations)
- 3. Which variable identifies individual observations and which is the cluster variable? How many clusters?
- 4. Which predictor(s) at lv1, which at lv2?
- Fit a null LMER model m0 and compute the ICC for the variable pupil

6. Fit a random-intercept model m1 that

includes the fixed effect fam

Hands on **Q**

1. Download & read the dataset from the Pupillometry multiverse study

```
id = infant's ID, fam = familiar. type (labeled vs. unlabeled), pupi1 = pupil dilation (mm)
library(osfr) # package to interact with the OSF platform
proj <- "https://osf.io/p8nfh/" # link to the OSF project (see protocol paper & data dictionary)
osf_download(osf_ls_files(osf_retrieve_node(proj))[5,],conflicts="overwrite") # download
infants <- read.csv2("data/multiverse.csv",stringsAsFactors=TRUE) # read dataset
colnames(infants)[c(18,17,19)] <- c("id","fam","pupi1") # shortening variable names
infants$pupi1 <- as.numeric(infants$pupil) # pupil as numeric</pre>
```

- Explore the the variables id, fam, and pupil (descriptives & correlations)
- Which variable identifies individual observations and which is the cluster variable? How many clusters?
- 4. Which predictor(s) at lv1, which at lv2?
- Fit a null LMER model m0 and compute the ICC for the variable pupil



1. Download & read the dataset from the Pupillometry multiverse study

```
id = infant's ID, fam = familiar. type (labeled vs. unlabeled), pupil = pupil dilation (mm)
library(osfr) # package to interact with the OSF platform
proj <- "https://osf.io/p8nfh/" # link to the OSF project (see protocol paper & data dictionary)
osf_download(osf_ls_files(osf_retrieve_node(proj))[5,],conflicts="overwrite") # download
infants <- read.csv2("data/multiverse.csv",stringsAsFactors=TRUE) # read dataset
colnames(infants)[c(18,17,19)] <- c("id","fam","pupil") # shortening variable names
infants$pupil <- as.numeric(infants$pupil) # pupil as numeric</pre>
```

- Explore the the variables id, fam, and pupil (descriptives & correlations)
- 3. Which variable identifies individual observations and which is the cluster variable? How many clusters?
- 4. Which predictor(s) at lv1, which at lv2?
- Fit a null LMER model m0 and compute the ICC for the variable pupil

- Fit a random-intercept model m1 that includes the fixed effect fam
- Fit a random-slope model m2 (i.e., 'free' the random slope for fam)

```
1. Download & read the dataset from the Pupillometry multiverse study
```

```
id = infant's ID, fam = familiar. type (labeled vs. unlabeled), pupil = pupil dilation (mm)
library(osfr) # package to interact with the OSF platform
proj <- "https://osf.io/p8nfh/" # link to the OSF project (see protocol paper & data dictionary)
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infants <- read.csv2("data/multiverse.csv",stringsAsFactors=TRUE) # read dataset
colnames(infants)[c(18,17,19)] <- c("id","fam","pupil") # shortening variable names
infants$pupil <- as.numeric(infants$pupil) # vupil as numeric</pre>
```

- Explore the the variables id, fam, and pupil (descriptives & correlations)
- 3. Which variable identifies individual observations and which is the cluster variable? How many clusters?
- 4. Which predictor(s) at lv1, which at lv2?
- Fit a null LMER model m0 and compute the ICC for the variable pupil

- Fit a random-intercept model m1 that includes the fixed effect fam
- Fit a random-slope model m2 (i.e., 'free' the random slope for fam)
- Assess model m2 diagnostics

Hands on **R**

1. Download & read the dataset from the Pupillometry multiverse study

```
id = infant's ID, fam = familiar. type (labeled vs. unlabeled), pupil = pupil dilation (mm)
library(osfr) # package to interact with the OSF platform
proj <- "https://osf.io/p8nfh/" # link to the OSF project (see protocol paper & data dictionary)
osf_download(osf_ls_files(osf_retrieve_node(proj))[5,],conflicts="overwrite") # download
infants <- read.csv2("data/multiverse.csv",stringsAsFactors=TRUE) # read dataset
colnames(infants)[c(18,17,19)] <- c("id","fam","pupil") # shortening variable names
infants$pupil <- as.numeric(infants$pupil) # pupil as numeric</pre>
```

- Explore the the variables id, fam, and pupil (descriptives & correlations)
- Which variable identifies individual observations and which is the cluster variable? How many clusters?
- 4. Which predictor(s) at lv1, which at lv2?
- Fit a null LMER model m0 and compute the ICC for the variable pupil

- Fit a random-intercept model m1 that includes the fixed effect fam
- Fit a random-slope model m2 (i.e., 'free' the random slope for fam)
- 8. Assess model m2 diagnostics
- Print, visualize, & interpret the fixed effects estimated by model m2:
 Is hypothesis HP1 confirmed?

Statistical inference on LMER coefficients

We saw that a coefficient estimate and its standard error (SE) are used to compute t-values and 95% CI (see slides #52-54). library(lme4) m2 <- lmer(pupil ~ fam + (1|id), data=infants) (s <- summarv(m2)\$coefficients)</pre> Estimate Std. Error t value (Intercept) 5819.1904 718.67898 8.097065 famunlabeled -349 4742 55 83258 -6 259324 s[2,1] / s[2,2] # t-value for fam (Est/SE)[1] -6.259324 s[2,1]-1.96*s[2,2] : s[2,1] + 1.96*s[2,2] # CI[1] -458.9061 [1] -240.0424 A fixed effect can be considered 'substantial' if t > 1.96 & CI exclude zero. Why is that? 'Rule of thumb' based on the standardized

normal distribution, where 1.96 corresponds to a probability of 0.05 (sounds familiar?)

We saw that a coefficient estimate and its standard error (SE) are used to compute *t*-values and 95% CI (see slides #52-54).

```
library(lme4)
m2 <- lmer(pupil ~ fam + (1|id), data=infants)
(s <- summary(m2)$coefficients)</pre>
```

Estimate Std. Error t value (Intercept) 5819.1904 718.67898 8.097065 famunlabeled -349.4742 55.83258 -6.259324

```
s[2,1] / s[2,2] # t-value for fam (Est/SE)
```

[1] -6.259324

```
s[2,1]-1.96*s[2,2]; s[2,1] + 1.96*s[2,2] # CI
[1] -458.9061
```

[1] -240.0424

A fixed effect can be considered 'substantial' if t>1.96 & CI exclude zero. Why is that? 'Rule of thumb' based on the standardized normal distribution, where 1.96 corresponds to a probability of 0.05 (sounds familiar?)

However, rules of thumb are insufficient to draw statistical inference on population parameters \rightarrow we need an *inference criterion*.



Within the NHST approach, p-values are used to determinate whether an effect is significant or not. Yet, in LMER p-values cannot be computed with the standard approach.

→ corrections have been proposed (e.g.,

Satterthwaite method used by lmerTest pkg).

library(lmerTest)

```
m2 <- lmer(pupil ~ fam + (1|id), data=infants)
summary(m2)$coefficients</pre>
```

Estimate Std. Error t value Pr(>|t|) (Intercept) 5819.19 718.68 8.10 7.225612e-07 famunlabeled -349.47 55.83 -6.26 3.928467e-10

LMER model comparison

An alternative way to quantify the 'importance' of a predictor is by comparing two models that only differ by the presence vs. absence of that predictor:

```
m0 <- lmer(pupil ~ (1|id), data = infants, REML = FALSE) # null model (intercept-only)
m1 <- lmer(pupil ~ fam + (1|id), data = infants, REML = FALSE) # model including fam
```

Statistical models aim at identifying the underlying process that generated the data, but many models can explain the same data, and none of that might be 'the true one'

Model comparison = identifying the model that best approximates the true model

LMER model comparison

An alternative way to quantify the 'importance' of a predictor is by *comparing* two models that only differ by the presence vs. absence of that predictor:

```
m0 <- lmer(pupil ~ (1|id), data = infants, REML = FALSE) # null model (intercept-only)
m1 <- lmer(pupil ~ fam + (1|id), data = infants, REML = FALSE) # model including fam
```

Statistical models aim at identifying the underlying process that generated the data, but many models can explain the same data, and none of that might be 'the true one'

Model comparison = identifying the model that best approximates the true model

Likelihood ratio test

likelihood) of the two models is equivalent.

If significant, it means that the more complex model improves the fit beyond what would be expected with the additional predictor added.

Tests the hypothesis H_0 that the fit (i.e.,

```
lmtest::lrtest(m0,m1)

#Df LogLik Df Chisq Pr(>Chisq)
1 3 -247926.1 NA NA NA
2 4 -247906.6 1 39.15 3.922796e-10
```

LMER model comparison

An alternative way to quantify the 'importance' of a predictor is by *comparing* two models that only differ by the presence vs. absence of that predictor:

```
m0 <- lmer(pupil ~ (1|id), data = infants, REML = FALSE) # null model (intercept-only)
m1 <- lmer(pupil ~ fam + (1|id), data = infants, REML = FALSE) # model including fam
```

Statistical models aim at identifying the underlying process that generated the data, but many models can explain the same data, and none of that might be 'the true one'

Model comparison = identifying the model that best approximates the true model

Likelihood ratio test

Tests the hypothesis H_0 that the fit (i.e., likelihood) of the two models is equivalent. If significant, it means that the more complex model improves the fit beyond what would be expected with the additional predictor added.

```
lmtest::lrtest(m0,m1)
```

```
#Df LogLik Df Chisq Pr(>Chisq)
1 3 -247926.1 NA NA NA
2 4 -247906.6 1 39.15 3.922796e-10
```

Information criteria

Measure model efficiency in terms of data forecasting, accounting for likelihood (better

fit) & parsimony (less predictors). Akaike

(AIC) & Bayesian Information Criterion

(BIC): the lower the better

```
AIC(m0,m1); BIC(m0,m1)

df AIC BIC
```

```
df AIC BIC
m0 3 495858.3 495882.7
m1 4 495821.1 495853.7
```

Effect sizes in multilevel modeling

Coefficient of determination (R^2)

Reflecting the **proportion of variance** in the dependent variable y that is **explained** by the independent variables x (see slide #9)

With LMER, we can compute to types of \mathbb{R}^2 :

- $Marginal R^2$: variance explained by fixed effects only / total variance
- \bullet $Conditional \ R^2\colon$ variance explained by fixed & random effects / total variance

In our case, the variance explained by the fixed slope of fam (R2m) is quite low (0.11%). It slightly increases with the random slope (0.30%), but it's still very low \odot In contrast, substantial variance is explained by the random effects (about 31-36%)

```
r.squaredGLMM(lmer(pupil~fam+(fam|id),data=infants)) # random slope model
```

Hands on **Q**, eyes on papers

Infants' pupil dilation

1. Fit models m0, m1, and m2 as in slide #66

Reading the Results section (pt2)

Infants' pupil dilation

- 1. Fit models m0, m1, and m2 as in slide #66
- We want to account the habituation effect
 on pupil dilation: fit a third model m3 that
 also includes time (time in ms over the trial),
 and a fourth model m4 including session
 (reflecting time on task)

Reading the Results section (pt2)

Infants' pupil dilation

- 1. Fit models m0, m1, and m2 as in slide #66
- We want to account the habituation effect
 on pupil dilation: fit a third model m3 that
 also includes time (time in ms over the trial),
 and a fourth model m4 including session
 (reflecting time on task)
- Evaluate model m4 diagnostics

Reading the Results section (pt2)

Infants' pupil dilation

- 1. Fit models m0, m1, and m2 as in slide #66
- We want to account the habituation effect
 on pupil dilation: fit a third model m3 that
 also includes time (time in ms over the trial),
 and a fourth model m4 including session
 (reflecting time on task)
- Evaluate model m4 diagnostics
- 4. Compare all models with the likelihood ratio test and the AIC: which is the best model?

Reading the Results section (pt2)

Infants' pupil dilation

- 1. Fit models m0, m1, and m2 as in slide #66
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 also includes time (time in ms over the trial),
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- 4. Compare all models with the likelihood ratio test and the AIC: which is the best model?
- 5. Print, visualize, and interpret the coefficients estimated by the selected model: which fixed effects are significant?

Reading the Results section (pt2)

Infants' pupil dilation

- 1. Fit models m0, m1, and m2 as in slide #66
- We want to account the habituation effect
 on pupil dilation: fit a third model m3 that
 also includes time (time in ms over the trial),
 and a fourth model m4 including session
 (reflecting time on task)
- Evaluate model m4 diagnostics
- 4. Compare all models with the likelihood ratio test and the AIC: which is the best model?
- 5. Print, visualize, and interpret the coefficients estimated by the selected model: which fixed effects are significant?
- 6. Print and interpret the coefficient of determination \mathbb{R}^2 of the selected model

Reading the Results section (pt2)

Infants' pupil dilation

- 1. Fit models m0, m1, and m2 as in slide #66
- We want to account the habituation effect
 on pupil dilation: fit a third model m3 that
 also includes time (time in ms over the trial),
 and a fourth model m4 including session
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- Evaluate model m4 diagnostics
- 4. Compare all models with the likelihood ratio test and the AIC: which is the best model?
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- 6. Print and interpret the coefficient of determination \mathbb{R}^2 of the selected model

Reading the Results section (pt2)

For each of the papers linked in slide #59:

Did the authors compare multiple models?
 Based on which criteria?

Infants' pupil dilation

- 1. Fit models m0, m1, and m2 as in slide #66
- We want to account the habituation effect
 on pupil dilation: fit a third model m3 that
 also includes time (time in ms over the trial),
 and a fourth model m4 including session
 (reflecting time on task)
- Evaluate model m4 diagnostics
- 4. Compare all models with the likelihood ratio test and the AIC: which is the best model?
- 5. Print, visualize, and interpret the coefficients estimated by the selected model: which fixed effects are significant?
- 6. Print and interpret the coefficient of determination \mathbb{R}^2 of the selected model

Reading the Results section (pt2)

- 1. Did the authors compare multiple models? Based on which criteria?
- 2. Do the authors report the likelihood ratio test of their models? Which is the best model?

Infants' pupil dilation

- 1. Fit models m0, m1, and m2 as in slide #66
- We want to account the habituation effect
 on pupil dilation: fit a third model m3 that
 also includes time (time in ms over the trial),
 and a fourth model m4 including session
 (reflecting time on task)
- Evaluate model m4 diagnostics
- 4. Compare all models with the likelihood ratio test and the AIC: which is the best model?
- 5. Print, visualize, and interpret the coefficients estimated by the selected model: which fixed effects are significant?
- 6. Print and interpret the coefficient of determination \mathbb{R}^2 of the selected model

Reading the Results section (pt2)

- Did the authors compare multiple models?
 Based on which criteria?
- 2. Do the authors report the likelihood ratio test of their models? Which is the best model?
- 3. Do the authors report the AIC and BIC indicators? Which is the best model?

Hands on **Q**, eyes on papers

Infants' pupil dilation

- 1. Fit models m0, m1, and m2 as in slide #66
- We want to account the habituation effect
 on pupil dilation: fit a third model m3 that
 also includes time (time in ms over the trial),
 and a fourth model m4 including session
 (reflecting time on task)
- 3. Evaluate model m4 diagnostics
- 4. Compare all models with the likelihood ratio test and the AIC: which is the best model?
- 5. Print, visualize, and interpret the coefficients estimated by the selected model: which fixed effects are significant?
- Print and interpret the coefficient of determination R² of the selected model

Reading the Results section (pt2)

- 1. Did the authors compare multiple models? Based on which criteria?
- 2. Do the authors report the likelihood ratio test of their models? Which is the best model?
- 3. Do the authors report the AIC and BIC indicators? Which is the best model?
- 4. Do the authors report the statistical significance of the estimated parameters? Which fixed effect is significant?

Infants' pupil dilation

- 1. Fit models m0, m1, and m2 as in slide #66
- We want to account the habituation effect
 on pupil dilation: fit a third model m3 that
 also includes time (time in ms over the trial),
 and a fourth model m4 including session
 (reflecting time on task)
- 3. Evaluate model m4 diagnostics
- 4. Compare all models with the likelihood ratio test and the AIC: which is the best model?
- 5. Print, visualize, and interpret the coefficients estimated by the selected model: which fixed effects are significant?
- 6. Print and interpret the coefficient of determination \mathbb{R}^2 of the selected model

Reading the Results section (pt2)

- 1. Did the authors compare multiple models? Based on which criteria?
- 2. Do the authors report the likelihood ratio test of their models? Which is the best model?
- 3. Do the authors report the AIC and BIC indicators? Which is the best model?
- 4. Do the authors report the statistical significance of the estimated parameters? Which fixed effect is significant?
- 5. Do the authors report the coefficient of determination? If yes, what proportion the y variance is explained by the models?

Questions?

Homework (optional):

- read the slides presented today and write in the Moodle forum if you have any doubts
- exe cises 10-11 from exeRcises.pdf

Mid-course survey on Moodle (please!): Open untill November 22nd

For each exercise, the solution (or one of the possible solutions) can be found in dedicated chunk of commented code within the exercises.Rmd file

Credits

The present slides are partially based on:

- Altoè, G. (2023) Corso Modelli lineari generalizzati ad effetti misti 2023. https://osf.io/b7tkp/
- Finch, W. H., Bolin, J. E., Kelley, K. (2014). Multilevel Modeling Using R (2nd edition). Boca Raton: CRC Press
- Pastore, M. (2015). Analisi dei dati in psicologie (e applicazioni in R). Il Mulino.

Useful resources on multilevel modeling

- Bates, D. (2022). lme4: Mixed-effects modeling with R. https://stat.ethz.ch/~maechler/MEMo-pages/lMMwR.pdf
- Baayen, R. H., Davidson, D. J., & Bates, D. M. (2008). Mixed-effects modeling with crossed random effects for subjects and items. *Journal of memory and language*, 59(4), 390-412.
- Bliese, P. (2022). Multilevel modeling in R (2.7).
 https://cran.r-project.org/doc/contrib/Bliese_Multilevel.pdf
- McElreath, R. (2020). Statistical rethinking: A Bayesian course with examples in R and Stan. Chapman and Hall/CRC.
- Pinheiro, J., & Bates, D. (2006). Mixed-effects models in S and S-PLUS. Springer science & business media.

Suggested papers on specific topics (see Moodle)

Data centering

 Enders, C. K., & Tofighi, D. (2007). Centering predictor variables in cross-sectional multilevel models: A new look at an old issue. *Psychological Methods*, 12(2), 121–138. https://doi.org/10.1037/1082-989X.12.2.121

Model selection & Information criteria

- Akaike, H. (1974). A new look at the statistical model identification. IEEE transactions on automatic control, 19(6), 716-723. https://doi.org/10.1109/TAC.1974.1100705
- Vrieze, S. I. (2012). Model selection and psychological theory: a discussion of the differences between the Akaike information criterion (AIC) and the Bayesian information criterion (BIC). Psychological methods, 17(2), 228. https://psycnet.apa.org/doi/10.1037/a0027127

Suggested papers on related topics (see Moodle)

Missing data in multilevel modeling

- Little, R. J. (1988). A test of missing completely at random for multivariate data with missing values. *Journal of the American statistical Association*, 83(404), 1198-1202. https://doi.org/10.1080/01621459.1988.10478722
- Newman, D. A. (2014). Missing data: Five practical guidelines. Organizational Research Methods, 17(4), 372-411.

Psychometrics of multilevel measures

- Cranford, J. A., Shrout, P. E., Iida, M., Rafaeli, E., Yip, T., & Bolger, N. (2006). A
 procedure for evaluating sensitivity to within-person change: Can mood measures in
 diary studies detect change reliably?. Personality and Social Psychology Bulletin, 32(7),
 917-929. https://doi.org/10.1177/0146167206287721
- Geldhof, G. J., Preacher, K. J., & Zyphur, M. J. (2014). Reliability estimation in a multilevel confirmatory factor analysis framework. Psychological methods, 19(1), 72. https://psycnet.apa.org/doi/10.1037/a0032138
- Stapleton, L. M., Yang, J. S., & Hancock, G. R. (2016). Construct meaning in multilevel settings. *Journal of Educational and Behavioral Statistics*, 41(5), 481-520. https://doi.org/10.3102/1076998616646200

Suggested papers on related topics (see Moodle)

Statistical power in multilevel models

- Kumle, L., Vō, M. L. H., & Draschkow, D. (2021). Estimating power in (generalized) linear mixed models: An open introduction and tutorial in R. Behavior research methods, 53(6), 2528-2543. https://doi.org/10.3758/s13428-021-01546-0
- Lafit, G., Adolf, J. K., Dejonckheere, E., Myin-Germeys, I., Viechtbauer, W., & Ceulemans, E. (2021). Selection of the number of participants in intensive longitudinal studies: A user-friendly shiny app and tutorial for performing power analysis in multilevel regression models that account for temporal dependencies. Advances in methods and practices in psychological science, 4(1), 2515245920978738. https://doi.org/10.1177/2515245920978738

Bayesian LMER

- Sorensen, T., & Vasishth, S. (2015). Bayesian linear mixed models using Stan: A tutorial for psychologists, linguists, and cognitive scientists. arXiv preprint arXiv:1506.06201. https://doi.org/10.20982/tqmp.12.3.p175
- Van de Schoot, R., Kaplan, D., Denissen, J., Asendorpf, J. B., Neyer, F. J., & Van Aken,
 M. A. (2014). A gentle introduction to Bayesian analysis: Applications to developmental
 research. Child development, 85(3), 842-860. https://doi.org/10.1111/cdev.12169

Suggested online resources

- Kristoffer Magnusson's website about R, statistics, psychotherapy, open science, and data visualization: https://rpsychologist.com/viz
- Quant Psych: very nice and funny YouTube channel on statistics applied to
 psychology data, including the topics of our course (e.g., LM, LMER, GLMER).
 https://www.youtube.com/@QuantPsych

Suggested online resources on specific topics

Coefficient of determination R^2

 Jason Fernando (2023) R-Squared: Definition, Calculation Formula, Uses, and Limitations. Available at this link

Introduction to LMER

 Quant Psych YouTube channel (2021). Mixed Models, Hierarchical Linear Models, and Multilevel Models: A simple explanation. Available at this link

Generalized LMER (GLMER)

- Quant Psych YouTube Channel. Understanding Generalized Linear Models (Logistic, Poisson, etc.). Available at this link
- Quant Psych YouTube Channel. Generalized Mixed Models in R. Available at this link

Bayesian LMER

Qixiang Fang and Rens van de Schoot (2019). Intro to Frequentist (Multilevel)
 Generalised Linear Models (GLM) in R with glm and lme4. Available at this link

- AIC: Akaike Information Criterion
- BIC: Bayesian Information Criterion
- ICC: intraclass correlation coefficient
- LM: linear models
- CI: confidence intervals
- MLE: maximum likelihood estimator
- OLS: ordinary least squares
- NHST: null hypothesis significance testing
- SD: standard deviation
- SE: standard error
- SS: sum of squares

- β = beta, indexing population-level intercept (β₀)
 and slope (β₁, β₂, etc.) parameters
- $\epsilon = epsilon$, indexing population-level errors to be estimated based on model residuals
- $\lambda = lambda$, indexing random effects (cluster-specific deviation from fixed coefficients)
- $\sigma = sigma$, indexing the variance σ^2 of population-level errors (or model residual)
- N = capital nu, indexing that a variable is normally distributed
- ρ = rho, indexing the correlation between random effects
- $\tau = tau$, indexing the variance of the random effects

Achronyms & Greek letters

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- ICC: intraclass correlation coefficient
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 - ciao