# Linear Mixed-Effects Models

Dale Barr

University of Glasgow

# Model specification

- specifying fixed effects
- specifying random effects

# Categorical predictors

- Recommendation: Make your own, don't rely on R defaults.
   Why?
  - model comparison doesn't work with variables of type factor
  - defaults don't support ANOVA-style interpretation

## LMEM versus t-test

You have run a study looking at the effects of alcohol consumption on simple reaction time. Data is stored in in the tables subjects and simple\_rt. Subjects (sub) were randomly assigned to one of two groups (cond). One group drank alcohol before performing the task, while the other had a placebo drink.

As a dependent variable, you measured how quickly each subject pressed a button in response to a flashing light (RT, in milliseconds). Each subject provided 8 measurements. Remove data from subjects S01 and S11 before analysis.

# t-test on subject means (1)

#### simple\_rt.zip

```
library("tidyverse")
subjects <- read csv("simple rt/subjects.csv",</pre>
                     col types = "icc",
                     progress = FALSE)
simple_rt <- read_csv("simple_rt/simple_rt.csv",</pre>
                      col types = "icci",
                      progress = FALSE)
combined <- subjects %>%
  filter(sub != "S01",
         sub != "S11") %>%
 inner join(simple rt, "sub") %>%
  select(sub, cond, RT)
subj means <- combined %>%
  group_by(sub, cond) %>%
  summarise(mean RT = mean(RT),
            .groups = "drop")
subj_means
```

```
# A tibble: 14 \times 3
   sub
         cond
                 mean_RT
   <chr> <chr>
                   <dbl>
         placebo
 1 S02
                    514.
 2 S03
         placebo
                    528.
 3 S04
         alcohol
                    507
         placebo
                    476.
 4 S05
 5 S06
         alcohol
                    450.
         placebo
 6 S07
                    488.
                    411.
 7 S08
         placebo
 8 S09
         alcohol
                    430.
                    458.
 9 S10
         alcohol
10 S12
         alcohol
                    537.
11 S13
         alcohol
                    500
         placebo
12 S14
                    434.
13 S15
         placebo
                    393.
14 S16
         alcohol
                    425
```

# t-test on subject means (2)

data: mean\_RT by cond
t = 0.35278, df = 12, p-value = 0.7304
alternative hypothesis: true difference in means
between group alcohol and group placebo is not equal
to 0
95 percent confidence interval:
 -46.21515 64.07230
sample estimates:
mean in group alcohol mean in group placebo

463.3214

Two Sample t-test

472.2500

# Random-intercepts LMEM

#### Level 1:

$$Y_{ij} = \beta_0 + \beta_1 X_{ij} + e_{ij}$$

## Level 2:

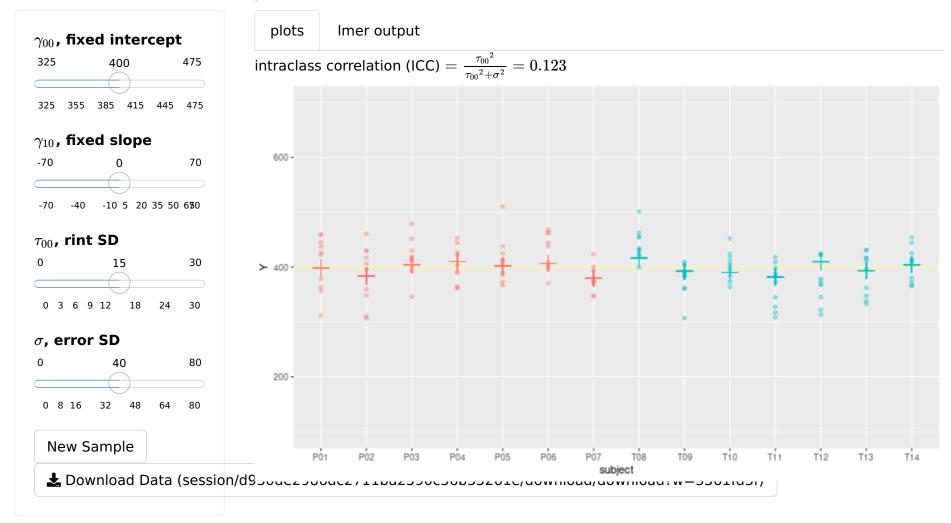
$$eta_0 = \gamma_{00} + S_{0i}$$
  $eta_1 = \gamma_{10}$ 

## **Variance Components**

$$S_{0i} \sim N\left(0,{ au_{00}}^2
ight)$$

$$e_{ij} \sim N\left(0,\sigma^2
ight)$$

## Random intercepts with ICC



For multi-level data, random-intercepts linear mixed-effects modeling can replace these analyses:

## between-subjects

- one-sample t-test
- independent samples t-test
- one-way ANOVA

## within/mixed designs

- paired samples t-test
- repeated-measures one-way ANOVA
- fully-within factorial ANOVA
- mixed-design ANOVA

NB: one obs per factor/cell

# Rules for random effects (1)

Always include random intercepts for any random factor (e.g., subjects) where you have multiple observations on the DV.

$$Y \sim (1 \mid subject)$$

Do I also need a random slope for factor A?

- 1. A is within-subjects
- 2. multiple observations per level of A

$$Y \sim A + (1 + A \mid subject)$$

# Rules for random effects (2)

What random slopes do I need for interaction ABC?

- identify highest-order combination of within factors
- if you have multiple observations per level of that factor / per cell of those factors, then you need a random slope for that factor / interaction of factors

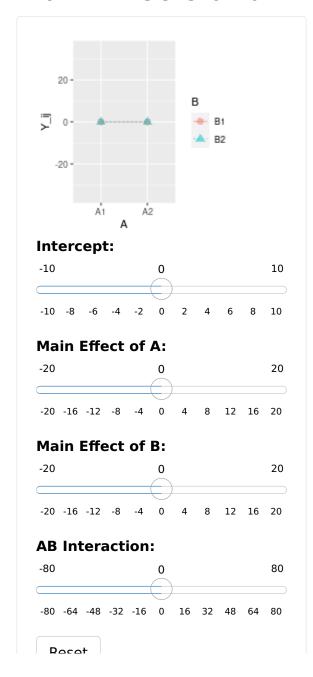
# Coding factorial predictors

Scheme	$A_1$	$A_2$
Treatment (dummy)	0	1
Sum	-1	1
Deviation	$-\frac{1}{2}$	$\frac{1}{2}$

Choice of a coding scheme impacts interpretation of:

- 1. the intercept term; and
- 2. the interpretation of the tests for all but the highest-order effects and interactions in a factorial design.

#### Main Effects and Interactions: 2x2 Factorial



#### **ANOVA**

Cell and Marginal Means

	B1	B2	
A1	0	0	0
A2	0	0	0
	0	0	NA

#### Simple Effects

eff		eff	
A@B1	0	B@A1	0
A@B2	0	B@A2	0

## Decomposition of Cell Means

cell	Y_ij	int	<b>A</b> _i	B_j	AB_ij
A1B1	0	0	0	0	0
A1B2	0	0	0	0	0
A2B1	0	0	0	0	0
A2B2	0	0	0	0	0

#### Regression

#### **Coding of A**

Dummy (A1=0,	
A2=1)	•

#### **Coding of B**

$$Y_{ijk}=eta_0+eta_1A_i+eta_2B_j+eta_3AB_{ij}+e_{ijk}$$
  $Y_{ijk}=0+0A_i+0B_j+0AB_{ij}+e_{ijk}$ 

cell	Y_ij	b0	<b>b1</b>	<b>A_i</b>	b2	B_j
A1B1	0	-0.00	0.00	0.00	-0.00	0.00
A1B2	0	-0.00	0.00	0.00	-0.00	1.00
A2B1	0	-0.00	0.00	1.00	-0.00	0.00
A2B2	0	-0.00	0.00	1.00	-0.00	1.00