Introduction to Within-Person Analysis and RM ANOVA

- Today's Class:
 - From between-person to within-person
 - > ANOVAs for longitudinal data
 - Model comparisons under ML (and now REML!)

The Two Sides of a (BP) Model

$$y_i = \beta_0 + \beta_1 X_i + \beta_2 Z_i + \beta_3 X_i Z_i + e_i$$

Model for the Means (Predicted Values):

Our focus today

- Each person's expected (predicted) outcome is a weighted linear function of his/her values on X and Z (and here, their interaction), each measured once per person (i.e., this is a between-person model)
- Estimated parameters are called fixed effects (here, β_0 , β_1 , β_2 , and β_3)
- The number of fixed effects will show up in formulas as k (so k = 4 here)

Model for the Variance ("Piles" of Variance):

- $e_i \sim N(0, \sigma_e^2) \rightarrow ONE$ residual (unexplained) deviation
- e_i has a mean of 0 with some estimated constant variance σ_e^2 , is normally distributed, is unrelated to X and Z, and is unrelated across people (across all observations, just people here)
- Contains residual variance only in above BP model

Review: Variances and Covariances

Variance:

Dispersion of y

Variance
$$(y_t) = \frac{\sum_{i=1}^{N} (y_{ti} - \hat{y}_{ti})^2}{N - k}$$

Covariance:

How y's go together, unstandardized

Covariance
$$(y_1, y_2) = \frac{\sum_{i=1}^{N} (y_{1i} - \hat{y}_{1i})(y_{2i} - \hat{y}_{2i})}{N - k}$$

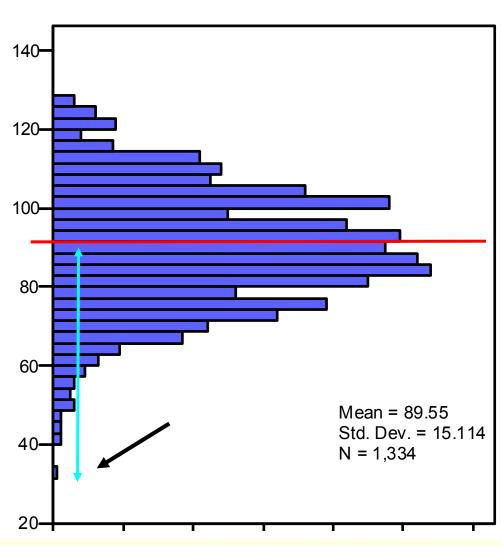
Correlation:

How y's go together, standardized (-1 to 1)

Correlation
$$(y_1, y_2) = \frac{\text{Covariance}(y_1, y_2)}{\sqrt{\text{Variance}(y_1)} * \sqrt{\text{Variance}(y_2)}}$$

N = # people, t = time, i = personk = # fixed effects, $\hat{y}_{ti} = y$ predicted from fixed effects

An Empty Between-Person Model (i.e., Single-Level)



$$y_i = \beta_0 + e_i$$

Filling in values:

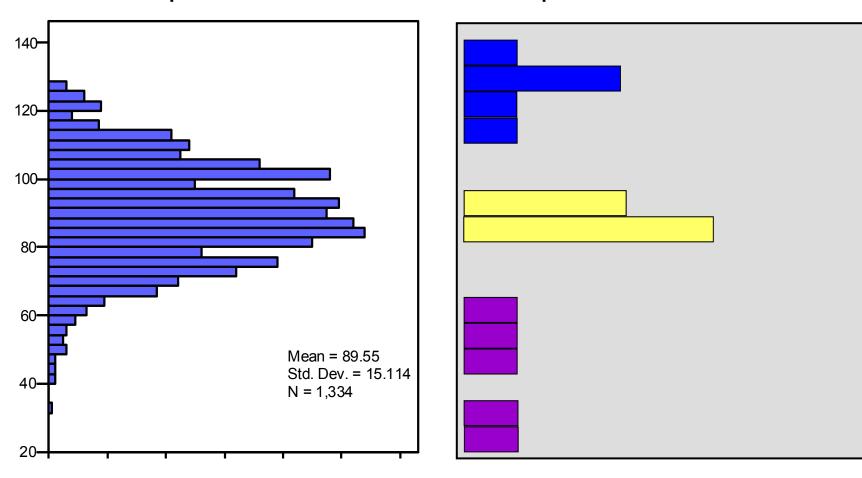
Y Error

Variance:

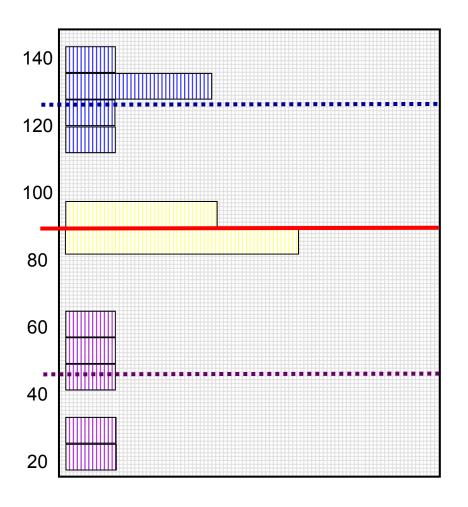
$$\frac{\Sigma (y - y_{\text{pred}})^2}{N - 1}$$

Adding Within-Person Information... (i.e., to become a Multilevel Model)

Full Sample Distribution 3 People, 5 Occasions each



Empty +Within-Person Model



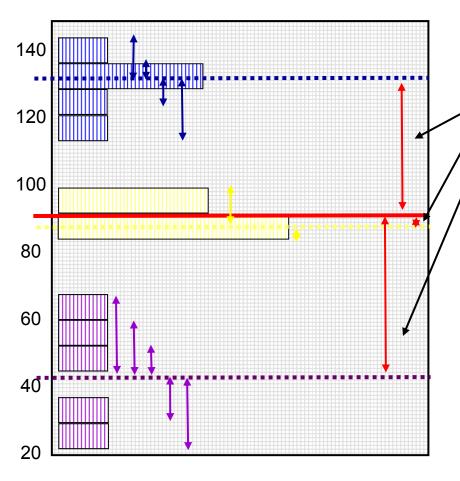
Start off with Mean of Y as "best guess" for any value:

- = Grand Mean
- = Fixed Intercept

Can make better guess by taking advantage of repeated observations:

- = Person Mean
- → Random Intercept

Empty +Within-Person Model



Variance of Y \rightarrow 2 sources:

Between-Person (BP) Variance:

Differences from **GRAND** mean

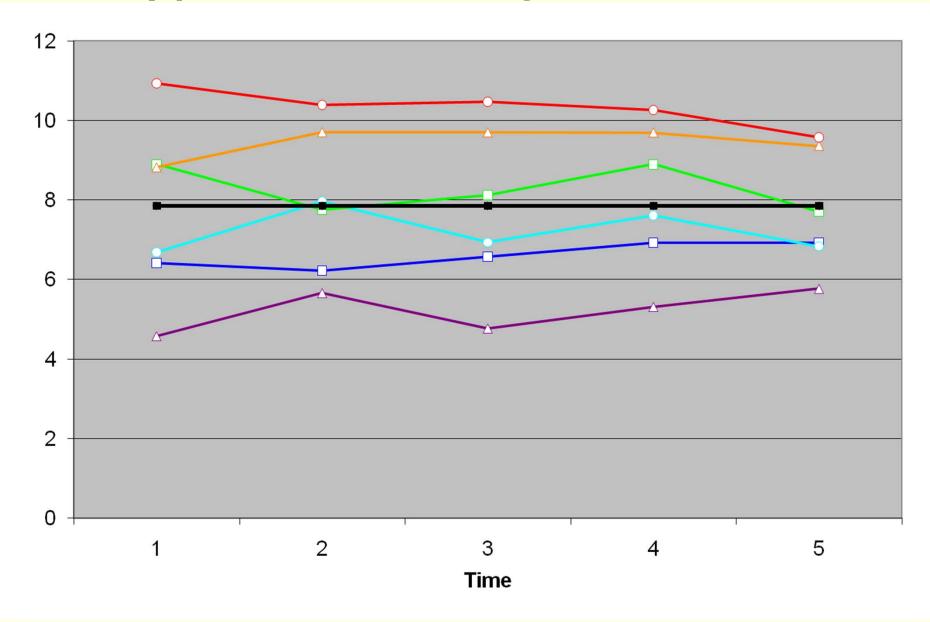
INTER-Individual Differences

Within-Person (WP) Variance:

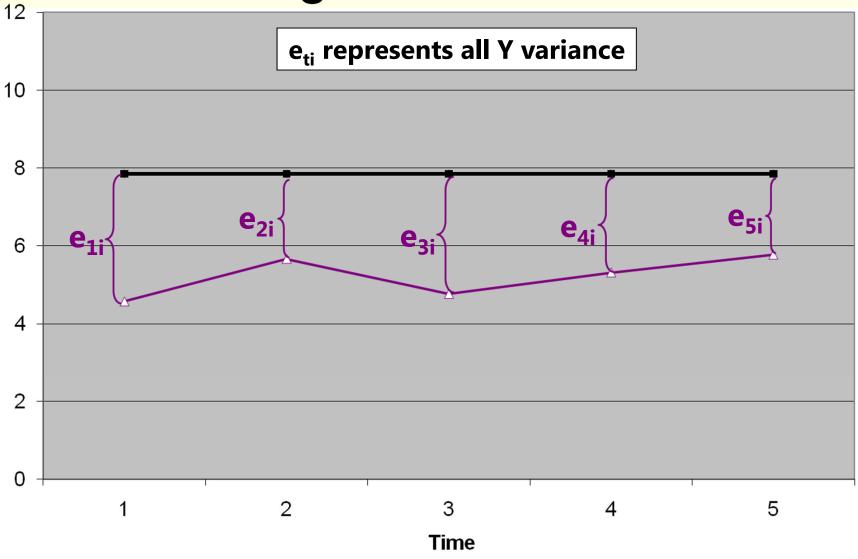
- → Differences from **OWN** mean
- → **INTRA**-Individual Differences
- → This part is only observable through longitudinal data.

Now we have 2 piles of variance in Y to predict.

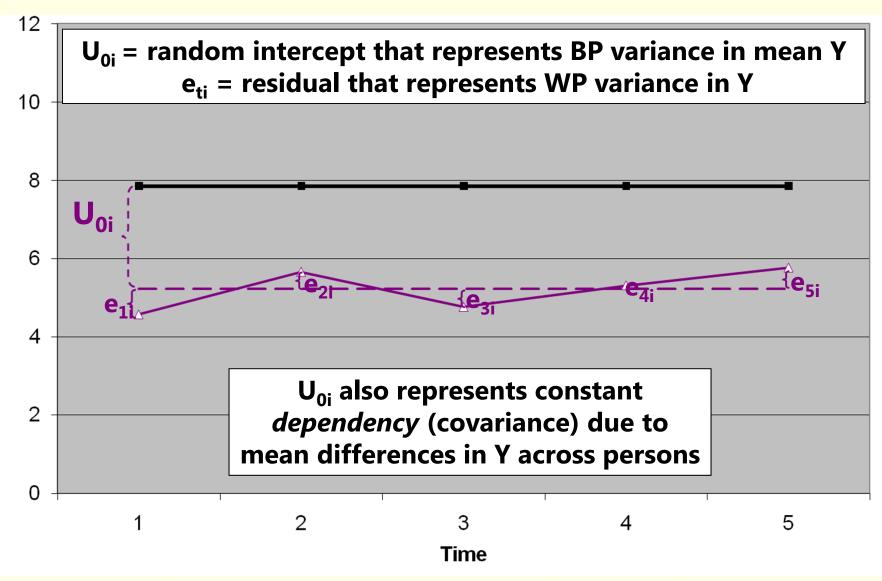
Hypothetical Longitudinal Data



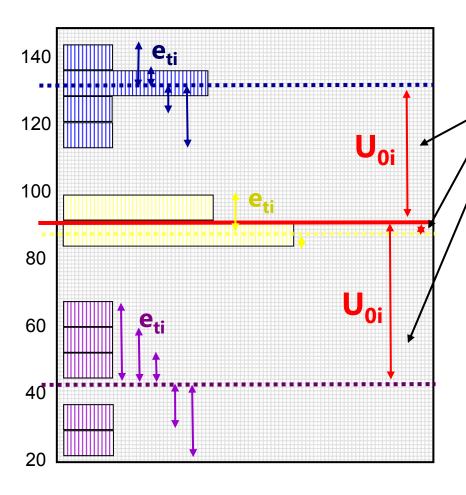
"Error" in a BP Model for the Variance: Single-Level Model



"Error" in a +WP Model for the Variance: Multilevel Model



Empty +Within-Person Model



Variance of Y \rightarrow 2 sources:

Level 2 Random Intercept

<u>Variance</u> (of U_{0i} , as $\tau_{U_0}^2$):

Between-Person Variance

Differences from **GRAND** mean

→ INTER-Individual Differences

Level 1 Residual Variance (of e_{ti} , as σ_e^2):

- → Within-Person Variance
- → Differences from **OWN** mean
- → **INTRA**-Individual Differences

BP vs. +WP Empty Models

• Empty Between-Person Model (used for 1 occasion):

$$y_i = \beta_0 + e_i$$

- β_0 = fixed intercept = grand mean
- e_i = residual deviation from GRAND mean
- Empty +Within-Person Model (>1 occasions):

$$y_{ti} = \beta_0 + U_{0i} + e_{ti}$$

- β_0 = fixed intercept = grand mean
- U_{0i} = random intercept = individual deviation from GRAND mean
- e_{ti} = time-specific residual deviation from OWN mean

Intraclass Correlation (ICC)

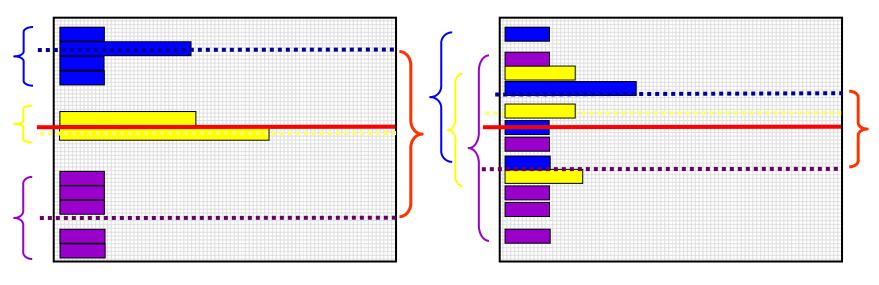
Intraclass Correlation (ICC):

$$\begin{split} ICC &= \frac{BP}{BP + WP} = \frac{Intercept\ Variance}{Intercept\ Variance + \ Residual\ Variance} \\ &= \frac{\tau_{U_0}^2}{\tau_{U_0}^2 + \sigma_e^2} \quad \begin{array}{c} R\ matrix \\ \begin{bmatrix} \sigma_e^2 + \tau_{u_0}^2 & \tau_{u_0}^2 \\ \tau_{u_0}^2 & \sigma_e^2 + \tau_{u_0}^2 \end{bmatrix} \begin{bmatrix} 1\ ICC\ ICC \\ ICC\ 1\ ICC \end{bmatrix} \\ \begin{bmatrix} \tau_{u_0}^2 & \tau_{u_0}^2 & \sigma_e^2 + \tau_{u_0}^2 \end{bmatrix} \\ \begin{bmatrix} \tau_{u_0}^2 & \tau_{u_0}^2 & \tau_{u_0}^2 \\ \tau_{u_0}^2 & \tau_{u_0}^2 & \sigma_e^2 + \tau_{u_0}^2 \end{bmatrix} \begin{bmatrix} 1\ ICC\ ICC \end{bmatrix} \\ ICC\ ICC\ 1 \end{bmatrix} \end{split}$$

- ICC = Proportion of total variance that is between persons
- ICC = Average correlation among occasions (in RCORR)
- ICC is a standardized way of expressing how much we need to worry about dependency due to person mean differences
 (i.e., ICC is an effect size for constant person dependency)

$$ICC = \frac{Between - Person}{Between - Person + Within - Person}$$

<u>Counter-Intuitive:</u> Between-Person Variance is in the numerator, but the ICC is the correlation over time!



ICC = BTW / BTW + within

- → Large ICC
- → Large correlation over time

ICC = btw / btw + WITHIN

- → Small ICC
- → Small correlation over time

BP and +WP Conditional Models

Multiple Regression, Between-Person ANOVA: 1 PILE

$$y_i = (\beta_0 + \beta_1 X_i + \beta_2 Z_i...) + e_i$$

- e_i → ONE residual, assumed uncorrelated with equal variance across observations (here, just persons) → "BP (all) variation"
- Repeated Measures, Within-Person ANOVA: 2 PILES
 - $y_{ti} = (\beta_0 + \beta_1 X_i + \beta_2 Z_i...) + U_{0i} + e_{ti}$
 - → U_{0i} → A random intercept for differences in person means, assumed uncorrelated with equal variance across persons → "BP (mean) variation" = $\tau_{U_0}^2$ is now "leftover" after predictors
 - $ightharpoonup \mathbf{e_{ti}}
 ightharpoonup A$ residual that represents remaining time-to-time variation, usually assumed uncorrelated with equal variance across observations (now, persons and time) ightharpoonup "**WP variation**" = σ_e^2 is also now "leftover" after predictors

Example Data for BP and WP Models

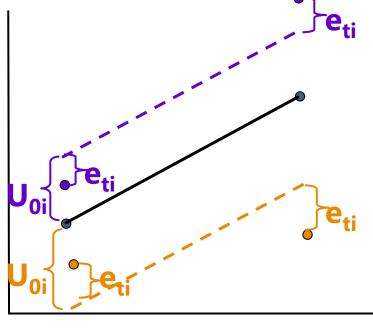
- 50 kids ages 10 and 11 assigned to control or treatment group
- Hypothesis: Outcome should be higher with age, with a greater age difference in the treatment group

Means (SE)	Age 10	Age 11	Marginal
Control	49.08 (1.14) 5.8 1.68 2.0		51.99 <i>(0.89)</i>
Treatment	50.76 (0.91) 7.8		54.70 (0.87)
Marginal	49.92 <i>(0.73)</i> 6.8	4 56.76 <i>(0.79)</i>	53.34 (0.64)

PSYC 944: Lecture 3

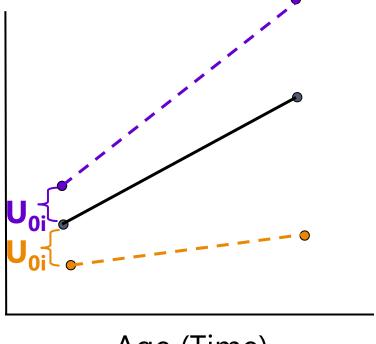
Why error and person*age are the same thing in two-occasion data

Same age slope, so error is leftover



Age (Time)

Different age slope, so no error is leftover



Age (Time)

ANOVA for longitudinal data?

- There are 3 possible "kinds" of ANOVAs we could use:
 - Between-Persons/Groups, Univariate RM, and Multivariate RM

NONE OF THEM ALLOW:

- > **Missing occasions** (do listwise deletion due to least squares)
- > **Time-varying predictors** (covariates are BP predictors only)
- Each includes the same model for the means for time: all possible mean differences (so 4 parameters to get to 4 means)
 - > "Saturated means model": $\beta_0 + \beta_1(T_1) + \beta_2(T_2) + \beta_3(T_3)$
 - > The *Time* variable must be balanced and discrete in ANOVA!
- These ANOVAs differ by what they predict for the correlation across outcomes from the same person in the model for the variances...
 - \triangleright i.e., **how they "handle dependency"** due to persons, or what they says the variance and covariance of the y_{ti} residuals should look like...

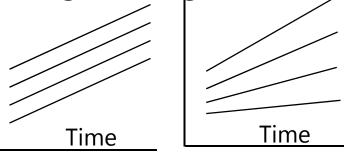
I. Between-Groups ANOVA

- Uses e_{ti} only (total variance = a single variance term of σ_e^2)
- Assumes no covariance at all among observations from the same person: Dependency? What dependency?
- Will usually be very, very wrong for longitudinal data
 - WP effects tested against wrong residual variance (significance tests will often be way too conservative)
 - Will also tend to be wrong for clustered data, but less so (because the correlation among persons from the same group is not as strong as the correlation among occasions from the same person)
- Predicts a variance-covariance matrix over time (here, 4 occasions) like this, called "Variance Components" (R matrix is TYPE=VC on REPEATED):

$$egin{bmatrix} \sigma_{e}^{2} & 0 & 0 & 0 \ 0 & \sigma_{e}^{2} & 0 & 0 \ 0 & 0 & \sigma_{e}^{2} & 0 \ 0 & 0 & \sigma_{e}^{2} & 0 \ \end{pmatrix}$$

2a. Univariate Repeated Measures

- Separates total variance into two sources:
 - **Between-Person** (mean differences due to U_{0i} , or $\tau_{U_0}^2$)
 - ightarrow Within-Person (remaining variance due to e_{ti} , or σ_e^2)
- Predicts a variance-covariance matrix over time (here, 4 occasions) like this, called "Compound Symmetry" (R matrix is TYPE=CS on REPEATED):
- $\begin{bmatrix} \sigma_{e}^{2} + \tau_{u_{0}}^{2} & \tau_{u_{0}}^{2} & \tau_{u_{0}}^{2} & \tau_{u_{0}}^{2} \\ \tau_{u_{0}}^{2} & \sigma_{e}^{2} + \tau_{u_{0}}^{2} & \tau_{u_{0}}^{2} & \tau_{u_{0}}^{2} \\ \tau_{u_{0}}^{2} & \tau_{u_{0}}^{2} & \sigma_{e}^{2} + \tau_{u_{0}}^{2} & \tau_{u_{0}}^{2} \\ \tau_{u_{0}}^{2} & \tau_{u_{0}}^{2} & \tau_{u_{0}}^{2} & \sigma_{e}^{2} + \tau_{u_{0}}^{2} \end{bmatrix}$
- Mean differences from U_{0i} are the only reason why occasions are correlated
- Will usually be at least somewhat wrong for longitudinal data
 - If people change at different rates, the variances and covariances over time have to change, too



The Problem with Univariate RM ANOVA

- Univ. RM ANOVA $(\tau_{U_0}^2 + \sigma_e^2)$ predicts **compound symmetry:**
 - > All variances and all covariances are equal across occasions
 - > In other words, the amount of error observed should be the same at any occasion, so a single, pooled error variance term makes sense
 - > If not, tests of fixed effects may be biased (i.e., sometimes tested against too much or too little error, if error is not really constant over time)
 - COMPOUND SYMMETRY RARELY FITS FOR LONGITUDINAL DATA
- But to get the correct tests of the fixed effects, the data must only meet a less restrictive assumption of sphericity:
 - ➤ In English → pairwise differences between adjacent occasions have equal variance and covariance (satisfied by default with only 2 occasions)
 - If compound symmetry is satisfied, so is sphericity (but see above)
 - > Significance test provided in ANOVA for where data meet sphericity assumption
 - > Other RM ANOVA approaches are used when sphericity fails...

The Other Repeated Measures ANOVAs...

2b. Univariate RM ANOVA with sphericity corrections

- \rightarrow Based on ε \rightarrow how far off sphericity (from 0-1, 1=spherical)
- Applies an overall correction for model df based on estimated ε, but it doesn't really address the problem that data ≠ model

• 3. Multivariate Repeated Measures ANOVA

All variances and covariances are estimated separately over time (here, 4 occasions), called "Unstructured" (R matrix is TYPE=UN on REPEATED)—it's not a model, it IS the data:

$$\begin{bmatrix} \sigma_{11}^2 & \sigma_{12} & \sigma_{13} & \sigma_{14} \\ \sigma_{21} & \sigma_{22}^2 & \sigma_{23} & \sigma_{24} \\ \sigma_{31} & \sigma_{32} & \sigma_{33}^2 & \sigma_{43} \\ \sigma_{41} & \sigma_{42} & \sigma_{43} & \sigma_{44}^2 \end{bmatrix}$$

Because it can never be wrong, UN can be useful for complete and balanced longitudinal data with few occasions (e.g., 2-4)

> Parameters =
$$\frac{\#occasions * (\#occasions + 1)}{2}$$
 so can be hard to estimate

- > Unstructured can also be specified to include random intercept variance $\tau_{U_0}^2$
- Every other model for the variances is nested within Unstructured (we can do model comparisons to see if all other models are NOT WORSE)

PSYC 944: Lecture 3

Summary: ANOVA approaches for longitudinal data are "one size fits most"

- Saturated Model for the Means (balanced time required)
 - > All possible mean differences
 - > Unparsimonious, but best-fitting (is a description, not a model)
- 3 kinds of Models for the Variances (complete data required)
 - \rightarrow BP ANOVA (σ_e^2 only) \rightarrow assumes independence and constant variance over time
 - ▶ Univ. RM ANOVA $(\tau_{U_0}^2 + \sigma_e^2)$ → assumes constant variance and covariance
 - ➤ Multiv. RM ANOVA (whatever) → no assumptions; is a description, not a model

there is no structure that shows up in a scalar equation (i.e., the way $U_{0i} + e_{ti}$ does)

MLM will give us more flexibility in both parts of the model:

- > Fixed effects that *predict* the pattern of means (polynomials, pieces)
- Random intercepts and slopes and/or alternative covariance structures that predict intermediate patterns of variance and covariance over time

3 Decision Points for Model Comparisons

1. Are the models **nested** or **non-nested**?

- > Nested: have to add OR subtract effects to go from one to other
 - Can conduct significance tests for improvement in fit
- Non-nested: have to add AND subtract effects
 - No significance tests available for these comparisons

2. Differ in model for the **means**, **variances**, or **both**?

- \rightarrow Means? Can only use ML -2Δ LL tests (or p-value of each fixed effect)
- \triangleright Variances? Can use ML (or preferably REML) $-2\Delta LL$ tests, no p-values
- \triangleright Both sides? Can only use ML -2Δ LL tests

3. Models estimated using **ML** or **REML**?

- > ML: All model comparisons are ok
- > REML: Model comparisons are ok for the variance parameters only

Likelihood-Based Model Comparisons

- Relative model fit is indexed by a "deviance" statistic → -2LL
 - > Log of likelihood (LL = total data height!) of observing the data given model parameters, -2*LL so that the differences between model LL values follow $\sim \chi^2$
 - > -2LL is a measure of BADNESS of fit, so smaller values = better models
 - Models are compared using their deviance values (significance tests)
 - ➤ Comes in two estimation flavors (labeled as -2 log likelihood on output): Maximum Likelihood (ML) or Restricted (Residual) ML (REML)
- Fit is also indexed by Information Criteria that reflect -2LL deviance
 AND # parameters used and/or sample size
 - \rightarrow **AIC** = Akaike IC = -2LL + 2 *(#parameters)
 - > **BIC** = Bayesian IC = -2LL + log(N)*(#parameters) → penalty for complexity
 - ➤ In ML → #parameters = all parameters (means and variances models)
 - ➤ In REML → #parameters = variance model parameters only (except in STATA!)
 - No significance tests or critical values, just "smaller is better"

$-2\Delta LL$ (i.e., LRT, Deviance) Tests:

(models must use the same estimator & N)

- 1. Calculate $-2\Delta LL$: $(-2LL_{fewer}) (-2LL_{more})$
- 2. Calculate Δdf : (# Parms_{more}) (# Parms_{fewer})

- 1. & 2. must be positive values!
- 3. Compare $-2\Delta LL$ to χ^2 distribution with df = Δ df CHIDIST function in excel will give exact p-values for the difference test
- Fixed effects p < .05: $-2\Delta LL(1) > 3.84$, $-2\Delta LL(2) > 5.99$, $-2\Delta LL(3) > 7.82$
- Some controversy about $-2\Delta LL$ tests when testing random effects variances that cannot be negative (i.e., the "boundary problem")
 - > χ^2 is not distributed as usual (mean=df) \rightarrow is actually a mixture χ^2 with df and df-1, so using the critical χ^2 for actual df results in conservative model comparison test
 - \rightarrow e.g., $-2\Delta LL(df=2)>5.99$, whereas $-2\Delta LL(df=mixture of 1,2)>5.14$
- Two proposed solutions when testing random effects variances:
 - For random intercepts, can use a 1-tailed test (χ^2 for p < .10): $-2\Delta LL(1) > 2.71$
 - ▶ Use mixture *p*-value = 0.5*prob($\chi^2_{df-1} > -2\Delta LL$) + 0.5*prob($\chi^2_{df} > -2\Delta LL$)
 - ➤ In practice these assume no relationship among how well variance parameters are estimated, which is suspect → I tend to just use the conservative test and call it good

Critical Values for 50:50 χ^2 Mixtures

Significance Level

df (q)	0.10	0.05	0.025	0.01	0.005
0 vs. 1	1.64	2.71	3.84	5.41	6.63
1 vs. 2	3.81	5.14	6.48	8.27	9.63
2 vs. 3	5.53	7.05	8.54	10.50	11.97
3 vs. 4	7.09	8.76	10.38	12.48	14.04
4 vs. 5	8.57	10.37	12.10	14.32	15.97
5 vs. 6	10.00	11.91	13.74	16.07	17.79
6 vs. 7	11.38	13.40	15.32	17.76	19.54
7 vs. 8	12.74	14.85	16.86	19.38	21.23
8 vs. 9	14.07	16.27	18.35	20.97	22.88
9 vs. 10	15.38	17.67	19.82	22.52	24.49
10 vs. 11	16.67	19.04	21.27	24.05	26.07

This may work ok if only one new parameter is bounded ... for example:

+ Random Intercept df=1: 2.71 vs. 3.84

+ Random Linear df=2: 5.14 vs. 5.99

+ Random Quad df=3: 7.05 vs. 7.82

Critical values such that the right-hand tail probability = $0.5 \times Pr (\chi^2_q > c) + 0.5 \times Pr (\chi^2_{q+1} > c)$

Source: Appendix C (p. 484) from Fitzmaurice, Laird, & Ware (2004). Applied Longitudinal Analysis. Hoboken, NJ: Wiley

ML vs. REML (more details to follow)

All comparisons must have same N!!!	ML	REML	
To select, type	METHOD=ML (-2 log likelihood)	METHOD=REML default (-2 res log likelihood)	
In estimating variances, it treats fixed effects as	Known (df for having to also estimate fixed effects is not factored in)	Unknown (df for having to estimate fixed effects is factored in)	
So, in small samples, L2 variances will be	Too small (less difference after N=30-50 or so)	Unbiased (correct)	
But because it indexes the fit of the	Entire model (means + variances)	Variances model only	
You can compare models differing in	Fixed and/or random effects (either/both)	Random effects only (same fixed effects)	

Rules for Comparing Multilevel Models

All observations must be the same across models!

Compare Models Differing In:

Type of Comparison:	Means Model	Variance Model	Both Means and
	(Fixed)	(Random)	Variances Model
	Only	Only	(Fixed and Random)
Nested? YES, can do significance tests via	Fixed effect p-values from ML or REML OR ML –2ΔLL only (NO REML –2ΔLL)	NO <i>p</i> -values REML –2ΔLL (ML –2ΔLL is ok if big N)	ML –2ΔLL only (NO REML –2ΔLL)
Non-Nested? NO signif. tests, instead see	ML AIC, BIC	REML AIC, BIC	ML AIC, BIC only
	(NO REML AIC, BIC)	(ML ok if big N)	(NO REML AIC, BIC)

<u>Nested</u> = one model is a <u>direct subset</u> of the other

Non-Nested = one model is not a direct subset of the other

Summary: Model Comparisons

- Significance of **fixed effects** can be tested with EITHER their
 p-values OR **ML** –2ΔLL (LRT, deviance difference) tests
 - \rightarrow p-value \rightarrow Is EACH of these effects significant? (fine under ML or REML)
 - \rightarrow ML –2 Δ LL test \rightarrow Does this SET of predictors make my model better?
 - \rightarrow REML -2 Δ LL tests are WRONG for comparing models differing in fixed effects
- Significance of random effects can only be tested with –2ΔLL tests
 (preferably using REML; here ML is not wrong, but results in too small
 variance components and fixed effect SEs in smaller samples)
 - > Can get *p*-values as part of output but *shouldn't* use them
 - > #parms added (df) should always include the random effect covariances
- My recommended approach to building models:
 - \triangleright Stay in REML (for best estimates), test new fixed effects with their p-values
 - \triangleright THEN add new random effects, testing $-2\Delta LL$ against previous model