# Review of CLDP 944: Multilevel Models for Longitudinal Data

- Topics:
  - Review of general MLM concepts and terminology
  - Model comparisons and significance testing
  - Fixed and random effects of time
  - > Significance testing and effect size in MLM

# What is a Multilevel Model (MLM)?

- Same as other terms you have heard of:
  - General(*ized*) Linear Mixed Model (if you are from statistics)
    - *Mixed* = Fixed and Random effects
  - Random Coefficients Model (also if you are from statistics)
    - Random coefficients = Random effects = latent variables/factors
  - > Hierarchical Linear Model (if you are from education)
- MLM is for modeling **dependency**. Special cases include:
  - > Random Effects ANOVA or Repeated Measures ANOVA
  - > (Latent) Growth Curve Model (where "Latent" implies use of SEM)
  - > Within-Person Fluctuation Model (e.g., for daily diary data)
  - Clustered/Nested Observations Model (e.g., for kids in schools)
  - > Cross-Classified Models (e.g., "value-added" models)
  - > Psychometric Models (e.g., factor analysis, item response theory)

## The Two Sides of \*Any\* Model

## Model for the Means:

- > Aka Fixed Effects, Structural Part of Model
- > What you are used to **caring about for testing hypotheses**
- How the expected outcome for a given observation varies as a function of values on predictor variables

## Model for the Variance:

- > Aka Random Effects and Residuals, Stochastic Part of Model
- What you \*were\* used to making assumptions about instead
- ➤ How residuals are distributed and related across observations (persons, groups, time, etc.) → these relationships are called "dependency" and this is the primary way that multilevel models differ from general linear models (e.g., regression)

## For Example: A Single-Level (BP) Model

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

• <u>Model for the Means (→Predicted Values)</u>:



- 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,  $\beta_0$ ,  $\beta_1$ ,  $\beta_2$ , and  $\beta_3$ )
- Model for the Variance (→ "Piles" of Variance):
  - e<sub>i</sub> ~ N(0, σ<sub>e</sub><sup>2</sup>) → ONE residual (unexplained) deviation, so
     estimated parameter is residual variance in single-level (BP) model
  - $e_i$  residuals have a mean of 0 with some estimated **constant variance**  $\sigma_e^2$ , are **normally distributed**, are unrelated to X and Z, and are **independent** across all observations
  - We should change models when any of these assumptions do not hold...

## Models We Will Learn in CLDP 945

- <u>Outcome type</u>: General (normal) vs. General*ized* (not normal)
- <u>Dimensions of sampling</u>: One (so one variance term per outcome) vs.
   Multiple (so multiple variance terms per outcome) → OUR WORLD
- <u>General Linear Models</u>: conditionally normal outcome distribution, fixed effects (identity link; only one dimension of sampling)

Note: Least Squares is only for GLM

- <u>Generalized Linear Models</u>: any conditional outcome distribution, fixed effects through link functions, no random effects (one dimension)
- <u>General Linear Mixed Models</u>: conditionally normal outcome distribution, fixed and random effects (identity link, but multiple sampling dimensions)
- <u>Generalized Linear Mixed Models</u>: any conditional outcome distribution, fixed and random effects through link functions (multiple dimensions)
  - > Many of the same concepts, but with more complexity in estimation
- "Linear" means fixed effects predict the *link-transformed* <u>conditional mean</u> of DV in a linear combination of (effect\*predictor) + (effect\*predictor)...

## What kinds of designs will we analyze?

- "Longitudinal" data (still, but with more complexity)
  - Same individual units of analysis measured at different occasions (which could range from milliseconds to days to years)
  - > Accelerated longitudinal designs; multiple levels of "time"
  - > Multivariate models (e.g., for families, dyads, and mediation)
- "Repeated measures" (RM) data (not involving "time")
  - Same individual units of analysis measured via different *items*, using different *stimuli*, or under different *conditions*
- "Clustered" and "cross-classified" data
  - Same individual units of analysis (one or more kinds of groups) measured via different *people* (cross-sectionally or longitudinally)

# **Options for Modeling Dependency**

- Many sampling designs have one or more sources/types of dependency, or correlation of observations from same unit
- Three main ways of building dependency into a model:
  - Fixed effects in the model for the means: add ID variable as a categorical predictor to represent differences across upper-level units
    - Main effects of ID represent intercept dependency; interactions of ID with lower-level predictors represent predictor-specific dependency
    - Does not allow prediction of why those differences occurred
  - Multivariate variance–covariance structures: for balanced longitudinal or repeated measures data; those using lags also require equal intervals
    - e.g., VC(H), CS(H), AR1(H), TOEP(H), or Unstructured (UN) as "answer key"
    - Can create a pattern of non-constant variance and covariance over time/RM
  - > Add one level (or more): add random intercept (and slope) variances
    - Can create multiple patterns of non-constant variance and covariance even with unbalanced data (longitudinal or clustered) → LET'S REVIEW THIS...

## **Two-Level Longitudinal Data**

- <u>Between-Person (BP) Variation:</u>
  - Level 2 "INTER-individual Differences" Time-Invariant
  - > All longitudinal studies begin as cross-sectional studies
- <u>Within-Person (WP) Variation</u> over Time:
  - Level 1 "INTRA-individual Differences" Time-Varying
  - > Only longitudinal studies can provide this extra information
- Longitudinal studies allow examination of both types of relationships simultaneously (and their interactions)
  - > Any variable measured over time usually has both BP and WP variation
  - > BP = more/less than other people; WP = more/less than one's average
- I use "person" here, but level 2 can be any entity that is measured repeatedly (like animals, schools, houses, countries...)

## Characterizing Longitudinal Data

- What should "time" be?
  - > e.g., time in study, age, grade, time from event/diagnosis
- Does time vary both **within-** AND **between-persons**?
  - > Model will need to differentiate each level of time effect
  - > Often known as "accelerated" longitudinal designs
- Is time balanced or unbalanced?
  - Balanced = everyone has a shared measurement schedule
    - Some people may miss occasions, making their data "incomplete"
  - > Unbalanced = people have different possible "time" values
    - By definition, observations are "incomplete" across persons
    - This is a consequence of any time metric varying between persons

# Characterizing Longitudinal Data



#### **Role of "Time" in the Model for the Means:**

- WP Change  $\rightarrow$  describe pattern of *average* change (e.g., growth curves)
- WP Fluctuation → describe average time-specific trends that may not have been expected (e.g., reactivity, day of the week, circadian/schedule effects)

#### **Role of "Time" in the Model for the Variance:**

- WP Change → describe *individual differences* in change (random effects)
   → this allows variances and covariances to differ over time
- WP Fluctuation  $\rightarrow$  describe pattern of variance and covariance over time

## ANOVA for two-level 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 when using 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 for *Time*:  $\beta_0 + \beta_1(T_1) + \beta_2(T_2) + \beta_3(T_3)$
  - > 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 variance...
  - i.e., how they "handle dependency" due to persons, or what they says the variance and covariance of the y<sub>ti</sub> residuals should look like...

# Summary: ANOVA models for longitudinal data are like "one size fits most"

- Saturated Model for the Means (balanced time required)
  - > All possible mean differences across time
  - > Unparsimonious, but best-fitting (is a description of the complete data)
- 3 kinds of Models for the Variance (need complete data in least squares)
  - > BP ANOVA ( $\sigma_e^2$  only; **VC**) → independence and constant variance over time
  - > Univ. RM ANOVA  $(\tau_{U_0}^2 + \sigma_e^2; \mathbf{CS}) \rightarrow$  constant variance and covariance over time
  - > Multiv. RM ANOVA (**unstructured**)  $\rightarrow$  is a description of the (complete) data

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 over time (polynomials, pieces)
  - Random intercepts and slopes and/or alternative covariance structures that predict intermediate patterns of variance and covariance over time

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



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



# Adding Within-Person Information... (i.e., to become a Two-Level Model)

Full Sample Distribution 3 People, 5 Occasions each



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## **Empty +Within-Person Model**



## $y_{ti}$ variance (V) $\rightarrow$ 2 sources:

### <u>Level-2 Random Intercept</u> <u>// Variance</u> (of U<sub>0i</sub>, as $\tau_{U_0}^2$ ):

**Between**-Person Variance in **G** Differences from **GRAND** mean

**INTER**-Individual Differences

### <u>Level-1 Residual Variance</u> (of $e_{ti}$ , as $\sigma_e^2$ ):

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

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



## BP vs. +WP Empty Models

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

 $y_i = \beta_0 + e_i$ 

- >  $\beta_0$  = fixed intercept = grand mean
- e<sub>i</sub> = residual deviation from GRAND mean
- Empty +Within-Person Model (for >1 occasions):

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

- >  $\beta_0$  = fixed intercept = grand mean
- > **U**<sub>0i</sub> = random intercept = individual deviation from GRAND mean
- e<sub>ti</sub> = time-specific residual deviation from OWN mean

## Same Model Using Multilevel Notation: Empty Means, Random Intercept Model

## **GLM Empty Model:**

- y<sub>i</sub> = β<sub>0</sub> + e<sub>i</sub>
- MLM Empty Model:
- Level 1:

 $y_{ti} = \beta_{0i} + e_{ti}$ 

• Level 2:

 $\beta_{0i} = \gamma_{00} + \mathbf{U}_{0i}$ 

Fixed Intercept = mean of means (=mean because no predictors yet) Random Intercept = individual-specific deviation from predicted intercept

3 Parameters: Model for the Means (1):

Fixed Intercept γ<sub>00</sub>

**Model for the Variance (2):** 

• Level-1 Variance of 
$$e_{ti} 
ightarrow \sigma_e^2$$

• Level-2 Variance of 
$$U_{0i} \rightarrow \tau_{U_0}^2$$

<u>Residual</u> = time-specific deviation from individual's predicted outcome

> Composite equation:  $y_{ti} = (\gamma_{00} + U_{0i}) + e_{ti}$

## Intraclass Correlation (ICC)

### **Intraclass Correlation (ICC):**

$$ICC = \frac{BP}{BP + WP} = \frac{Intercept Var.}{Intercept Var. + Residual Var.} = \frac{\tau_{U_0}^2}{\tau_{U_0}^2 + \sigma_e^2}$$
$$Corr(y_1, y_2) = \frac{Cov(y_1, y_2)}{\sqrt{Var(y_1)} * \sqrt{Var(y_2)}} \begin{bmatrix} V \text{ matrix } VCORR \text{ Matrix} \\ \begin{bmatrix} \sigma_e^2 + \tau_{u_0}^2 & \tau_{u_0}^2 & \tau_{u_0}^2 \\ \tau_{u_0}^2 & \sigma_e^2 + \tau_{u_0}^2 & \tau_{u_0}^2 \end{bmatrix} \begin{bmatrix} 1 & ICC & ICC \\ ICC & 1 & ICC \\ ICC & ICC \end{bmatrix}$$

- ICC = Proportion of total variance that is between persons
- ICC = Correlation of occasions from same person (in VCORR)
- 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 <u>constant</u> person dependency)

# Augmenting the empty means, random intercept model with *time*

• 2 questions about the possible effects of *time*:

## 1. Is there an effect of time on average?

- > If the line describing the sample means not flat?
- > Significant FIXED effect of time

# 2. Does the average effect of time vary across individuals?

- > Does each individual need his or her own line?
- > Significant RANDOM effect of time

Fixed and Random Effects of Time

(Note: The intercept is random in every figure)



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## B. Fixed Linear Time, Random Intercept Model (4 parameters: effect of time is FIXED only)



## C or D: Random Linear Time Model (6 parms)

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## Random Linear Time Model



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## Summary: Sequential Models for Effects of Time

Level 1:  $\mathbf{y}_{ti} = \mathbf{\beta}_{0i} + \mathbf{e}_{ti}$ Level 2:  $\mathbf{\beta}_{0i} = \mathbf{\gamma}_{00} + \mathbf{U}_{0i}$ Composite:  $\mathbf{y}_{ti} = \mathbf{\gamma}_{00} + \mathbf{U}_{0i} + \mathbf{e}_{ti}$  Empty Means, Random Intercept Model: 3 parms =  $\gamma_{00}$ ,  $\sigma_e^2$ ,  $\tau_{U_0}^2$ 

Level 1: $y_{ti} = \beta_{0i} + \beta_{1i} (Time_{ti}) + e_{ti}$	Fixed Linear Time,
Level 2: $\beta_{0i} = \gamma_{00} + U_{0i}$	Random Intercept Model:
$\beta_{1i} = \gamma_{10}$	4 parms = $\gamma_{00}$ , $\gamma_{10}$ , $\sigma_e^2$ , $\tau_{U_0}^2$

Composite:  $y_{ti} = (\gamma_{00} + U_{0i}) + \gamma_{10}(Time_{ti}) + e_{ti}$ 

Level 1:  $\mathbf{y}_{ti} = \mathbf{\beta}_{0i} + \mathbf{\beta}_{1i}(\mathsf{Time}_{ti}) + \mathbf{e}_{ti}$ Level 2:  $\mathbf{\beta}_{0i} = \mathbf{\gamma}_{00} + \mathbf{U}_{0i}$   $\mathbf{\beta}_{1i} = \mathbf{\gamma}_{10} + \mathbf{U}_{1i}$ Random Linear Time Model: 6 parms =  $\mathbf{\gamma}_{00}, \mathbf{\gamma}_{10}, \sigma_{e}^{2}, \tau_{U_{0}}^{2}, \tau_{U$ 

Composite:  $y_{ti} = (\gamma_{00} + U_{0i}) + (\gamma_{10} + U_{0i})(Time_{ti}) + e_{ti}$ 

## Random Linear Time Models Imply:

- People differ from each other systematically in TWO ways—in intercept (U<sub>0i</sub>) and slope (U<sub>1i</sub>), which implies TWO kinds of BP variance, which translates to TWO sources of person dependency (covariance or correlation in the outcomes from the same person)
- If so, after controlling for both BP intercept and slope differences (by estimating the  $\tau_{U_0}^2$  and  $\tau_{U_1}^2$  variances in the **G** matrix), the **e**<sub>ti</sub> **residuals** (whose variance and covariance are estimated in the R matrix) should be **uncorrelated with homogeneous variance across time**, as shown (or else a different **R** matrix is needed):

Level	-2	Level-1 <b>R</b> matrix:				
<b>G</b> mat	rix:	REPEATED TYPE=V			E=VC	
RAND	MC	[	$\sigma^2$	0	0	0 ]
TYPE=	UN		° e	2	ů O	
$\int \tau^2$	<b>-</b> ]		0	$\sigma_{e}^{-}$	0	0
	U <sub>10</sub>		0	0	$\sigma_e^2$	0
$\tau_{U_{01}}$	$\tau^2_{U_1}$		0	0	0	$\sigma_e^2$

G and R combine to create a total
V matrix whose per-person
structure depends on the specific
time occasions for each person in
Z (flexible for unbalanced time)

## What Does Each Side of the Model Need?

- Nested models (i.e., in which one is a subset of the other) can now differ from each other in two important ways
- Model for the Means → which predictors and which fixed effects of them are included in the model
  - <u>Does not</u> require assessment of relative model fit using LL or -2LL (can use univariate or multivariate Wald tests for this)
- Model for the Variance → what the pattern of variance and covariance of residuals from the same unit should be
  - ▹ DOES require assessment of relative model fit using LL or −2LL
  - Cannot use the Wald test *p*-values that show up on the output for testing significance of variances because those *p*-values are use a two-sided sampling distribution for what the variance could be (but variances cannot be negative, so those *p*-values are not valid)

# Testing Significance of Fixed Effects (of Predictors) in the Model for the Means

- Any single-df **fixed effect** has 4-5 relevant pieces of output:
  - Estimate = best guess for the fixed effect from our data
  - Standard Error = precision of fixed effect estimate (quality of most likely estimate)
  - *t*-value or *z*-value = Estimate / Standard Error
  - > *p***-value** = probability that fixed effect estimate is  $\neq 0$
  - > 95% Confidence Interval = Estimate ± 1.96\*SE = range in which true (population) value of estimate is expected to fall 95% of the time
- Compare test statistic (t or z) to critical value at chosen level of significance (known as alpha): this is a "univariate Wald test"
- Whether the *p*-value is based on *t* or *z* varies by program...

## Evaluating Significance of Fixed Effects

Fixed effects can be tested via **Wald** tests: the ratio of its estimate/SE forms a statistic we compare to a distribution

	Denominator DF is infinite (Proper Wald test)	Denominator DF is estimated instead ("Modified" Wald test)
Numerator DF = 1 (test one fixed effect) is <b>Univariate Wald Test</b>	use <b>z</b> distribution (Mplus, STATA)	use <b>t</b> distribution (SAS, SPSS)
Numerator DF > 1 ( <i>test 2</i> + <i>fixed effects</i> ) is <b>Multivariate Wald Test</b>	use <b>χ²</b> distribution (Mplus, STATA)	use <b>F</b> distribution (SAS, SPSS)
Denominator DF options	not applicable, so DDF is not given	SAS, STATA 14: BW, KR SAS, STATA 14, SPSS: Satterthwaite

## Evaluating Effect Size of Fixed Effects

- Most common measure of effect size in MLM is Pseudo-R<sup>2</sup>
  - > Is supposed to be variance accounted for by predictors
  - Multiple piles of variance mean multiple possible values of pseudo R<sup>2</sup> (can be calculated per variance component or per model level)
  - > A fixed linear effect of time will reduce level-1 residual variance  $\sigma_e^2$  in  ${\bm R}$
  - > By how much is the residual variance  $\sigma_e^2$  reduced?

Pseudo  $R_e^2 = \frac{\text{residual variance}_{\text{fewer}} - \text{residual variance}_{\text{more}}}{\text{residual variance}_{\text{fewer}}}$ 

> If time varies between persons, then level-2 random intercept variance  $\tau_{U_0}^2$  in **G** may also be reduced:

Pseudo  $R_{U0}^2 = \frac{\text{random intercept variance}_{\text{fewer}} - \text{random intercept variance}_{\text{more}}}{\text{random intercept variance}_{\text{fewer}}}$ 

> But you are likely to see a (net) INCREASE in  $\tau_{U_0}^2$  instead.... Here's why:

## Increases in Random Intercept Variance

- Level-2 random intercept variance  $\tau_{U_0}^2$  will often increase as a consequence of reducing level-1 residual variance  $\sigma_e^2$
- Observed level-2  $\tau_{U_0}^2$  is NOT just between-person variance
  - > Also has a small part of within-person variance (level-1  $\sigma_e^2$ ), or: **Observed**  $\tau_{U_0}^2$  = **True**  $\tau_{U_0}^2$  + ( $\sigma_e^2/n$ )
    - As *n* occasions increases, bias of level-1  $\sigma_e^2$  is minimized
  - > Likelihood-based estimates of "true"  $\tau_{U_0}^2$  use  $(\sigma_e^2/n)$  as correction factor: **True**  $\tau_{U_0}^2$  = **Observed**  $\tau_{U_0}^2$  –  $(\sigma_e^2/n)$
- For example: observed level-2  $\tau_{U_0}^2$ =4.65, level-1  $\sigma_e^2$ =7.06, n=4
  - > True  $\tau_{U_0}^2$  = 4.65 (**7.60**/4) = **2.88** in empty means model
  - > Add fixed linear time slope  $\rightarrow$  reduce  $\sigma_e^2$  from 7.06 to 2.17 (R<sup>2</sup> = .69)
  - > But now True  $\tau_{U_0}^2 = 4.65 (2.17/4) = 4.10$  in fixed linear time model

# Significance Tests for Choosing Models for the Variance

- Requires assessment of **relative model fit**: how well does the model fit relative to other possible models?
  - > Assessment of *absolute* model fit is only possible for balanced data
- Relative fit is indexed by overall model log-likelihood (LL):
  - > Log of likelihood for each person's outcomes given model parameters
  - Sum log-likelihoods across all independent persons = model LL
  - > Two flavors: Maximum Likelihood (ML) or Restricted ML (REML)
- What you get for this on your output varies by software...
- Given as -2\*log likelihood (-2LL) in SAS or SPSS MIXED:
   -2LL gives BADNESS of fit, so smaller value = better model
- Given as just log-likelihood (LL) in STATA MIXED and Mplus:
   LL gives GOODNESS of fit, so bigger value = better model

## Comparing Models for the Variance

#### • Two strategies for choosing a model for the variance:

- > Does the more complex model fit better (than a simpler model)?
- Does the simpler model fit worse (than a more complex model)?
- Nested models are compared using a "likelihood ratio test":
   -2ΔLL test (aka, "χ<sup>2</sup> test" in SEM; "deviance difference test" in MLM)

"fewer" = from model with fewer parameters "more" = from model with more parameters Results of 1. & 2. must be positive values!

- 1. Calculate -2 $\Delta$ LL: if given -2LL, do -2 $\Delta$ LL = (-2LL<sub>fewer</sub>) (-2LL<sub>more</sub>) if given LL, do -2 $\Delta$ LL = -2 \*(LL<sub>fewer</sub> - LL<sub>more</sub>)
- 2. Calculate  $\Delta df = (\# Parms_{more}) (\# Parms_{fewer})$
- 3. Compare  $-2\Delta LL$  to  $\chi^2$  distribution with df =  $\Delta df$
- 4. Get *p*-value from CHIDIST in excel or LRTEST option in STATA

## Comparing Models for the Variance

- What your *p*-value for the  $-2\Delta LL$  test means:
  - > If you ADD parameters, then your model can get better (if −2ΔLL test is significant ) or not better (not significant)
  - > If you **REMOVE** parameters, then your model can get **worse** (if −2ΔLL test is significant ) or **not worse** (not significant)
- Nested or non-nested models can also be compared by Information Criteria that also reflect model parsimony
  - > No significance tests or critical values, just "smaller is better"
  - > **AIC** = Akaike IC = -2LL + 2\*(# parameters)
  - > **BIC** = Bayesian IC =  $-2LL + \log(N)^*(\# parameters)$
  - > What "parameters" means depends on flavor (except in stata):
    - ML = ALL parameters; REML = variance model parameters only

## Flavors of Maximum Likelihood

- Remember that Maximum likelihood comes in two flavors:
- "Restricted (or residual) maximum likelihood"
  - Only available for general linear models or general linear mixed models (that assume normally distributed residuals)
  - > Is same as LS given complete outcomes, but it doesn't require them
  - > Estimates variances the same way as in LS (accurate)  $\rightarrow \frac{\sum(y_i y_{pred})^2}{N k}$
- "Maximum likelihood" (ML; also called FIML\*)
  - Is more general, is available for the above plus for non-normal outcomes and latent variable models (CFA/SEM/IRT)
  - > Is NOT the same as LS: it under-estimates variances by  $\frac{\sum(y_i y_{pred})^2}{N}$ not accounting for the # of estimated fixed effects  $\rightarrow \frac{N}{N}$
- \*FI = Full information  $\rightarrow$  it uses all original data (they both do)

# Flavors of Full-Information Maximum Likelihood

- <u>Restricted maximum likelihood</u> (**REML**; used in MIXED)
  - Provides unbiased variances
  - > Especially important for small N (< 100 units)</p>
  - -2ΔLL test cannot be used to compare models differing in fixed effects (no biggee; we can do this using univariate or multivariate Wald tests)
  - **-2ΔLL test** MUST be used to compare different models for the variance
- Maximum likelihood (ML; also used in MIXED)
  - Variances (and SEs) are too small in small samples
  - Is only option in most software for path models and SEM
  - –2ΔLL test can be used to compare any nested model; must be used to compare different models for the variance

$$\frac{\sum(y_i - y_{pred})^2}{N}$$

 $\frac{\sum (y_i - y_{pred})^2}{\sum (y_i - y_{pred})^2}$ N - k

## ML vs. REML in a nutshell

Remember "population" vs. "sample" formulas for calculating variance?		<b>'Population''</b> $\frac{\Sigma(y_i - y_{pred})^2}{N}$	"Sample" $\frac{\Sigma(y_i - y_{pred})^2}{N - k}$		
All comparisons must have same N!!!	ML		REML		
To select, type	METHOD=ML (-2 log likelihood)		METHOD=REML <i>default</i> (-2 res log likelihood)		
In estimating variances, it treats fixed effects as	<b>Known</b> (df for having to also estimate fixed effects is not factored in)		<b>Unknown</b> (df for having to estimate fixed effects is factored in)		
So, in small samples, L2 variances will be	<b>Too small</b> (less difference after N=30-50 or so)		<b>Too small</b> (less difference after N=30-50 or so)		Unbiased (correct)
But because it indexes the fit of the	<b>Entire model</b> (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 Models

#### All observations must be the same across models!

## **Compare Models Differing In:**

Type of Comparison:	Means Model (Fixed) Only	Variance Model (Random) Only	Both Means and Variances Model (Fixed and Random)
<u>Nested?</u> 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 (NO REML AIC, BIC)	REML AIC, BIC (ML ok if big N)	ML AIC, BIC only (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

## 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**?
  - > Means? Can only use  $-2\Delta LL$  tests if ML (or *p*-value of each fixed effect)
  - > Variances? Can use ML (or preferably REML)  $-2\Delta$ LL tests, no *p*-values
  - > Both sides? Can only use  $-2\Delta LL$  tests if ML
- 3. Models estimated using **ML** or **REML**?
  - > ML: All model comparisons are ok
  - > REML: Model comparisons are ok for the variance parameters only

## Effect Size for Random Effects Variances

- We can test if a random effect variance is significant, but the variance estimates are not likely to have inherent meaning
  - > e.g., "I have a significant fixed linear time effect of  $\gamma_{10} = 1.72$ , so people increase by 1.72/time on average. I also have a significant random linear time slope variance of  $\tau_{U_1}^2 = 0.91$ , so people need their own slopes (people change differently). But how much is a variance of 0.91, really?"

#### • 95% Random Effects Confidence Intervals can tell you

- > Can be calculated for each effect that is random in your model
- Provide range around the fixed effect within which 95% of your sample is predicted to fall, based on your random effect variance:

Random Effect 95% CI = fixed effect  $\pm (1.96*\sqrt{\text{Random Variance}})$ 

Linear Time Slope 95% CI =  $\gamma_{10} \pm \left(1.96^* \sqrt{\tau_{U_1}^2}\right) \rightarrow 1.72 \pm \left(1.96^* \sqrt{0.91}\right) = -0.15 \text{ to } 3.59$ 

 So although people improve on average, individual slopes are predicted to range from -0.15 to 3.59 (so some people may actually decline)

# Another Variance Model Effect Size: Intercept/Slope Reliability

• Another measure of effect size for random effects variances is Intercept Reliability (IR) or Slope Reliability (SR)

$$\begin{aligned} & \tau_{U_1}^2 = \text{random slope variance} \\ & \sigma_e^2 = \text{residual variance} \\ & L1n = L1 \text{ sample size per L2 unit} \\ & \sigma_{L1}^2 = \text{variance of L1 predictor} \end{aligned} \qquad \mathbf{SR} = \frac{\tau_{U_1}^2}{\tau_{U_1}^2} + \frac{\sigma_e^2}{L1n * \sigma_{L1}^2} \end{aligned}$$

- IR formula is the same, just replacing  $\sigma_{L1}^2$  with 1
- SR is known as growth rate reliability in context of time (Willett, 1989)

## The Big Picture of Longitudinal Data: Models for the Means for Time

- What kind of change occurs on average over "time"? There are two baseline models to consider:
  - *F* "Empty" → only a fixed intercept (predicts no change)
  - > "Saturated" → all occasion mean differences from time 0 (ANOVA model that uses # fixed effects = n) \*\*\* may not be possible in unbalanced data



<u>In-between options:</u> polynomial slopes, piecewise slopes, nonlinear slopes...

#### **Saturated Means:**

Reproduces mean at each occasion

**# Fixed Effects** 

= # Occasions

# **Good fit**

#### Name... that... Trajectory!

## The Big Picture of Longitudinal Data: Models for the Variance for Time



#### What is the pattern of variance and covariance over time?

CS and UN are just two of the many, many options available within MLM, including *random effects models* (for change) and *alternative covariance structure models* (for fluctuation).

## Summary: Unconditional Longitudinal Models

#### **Model for the Means for Time:**

- What kind of **fixed effects of time** are needed to create a function with which to parsimoniously **represent the pattern of saturated means** across time?
  - > Continuous or discontinuous? This choice likely comes from the design!
  - > Polynomials? Pieces? Log time? Truly nonlinear? This comes from the means plot!
  - > Use obtained *p*-values to test significance of fixed effects (Wald test)
  - > Use pseudo-R<sup>2</sup> values to describe effect size (just for residual if time is WP only)

#### **Model for the Variance for Time (building V):**

- What kind of **random effects of time** in G are needed :
  - > To account for **individual differences in each aspect of change**?
  - > To describe any non-constant variance and covariance across occasions?
  - > Do the residuals in R show any covariance after accounting for random effects?
  - > Use REML  $-2\Delta$ LL tests to test significance of new effects (or ML if big upper-level *N*)
  - > Use random effects CIs and intercept/slope reliability to describe effect size

## Summary of Unconditional Time Models

- Each source of correlation or dependency goes into a new variance component (or pile of variance) until each source meets the usual assumptions of GLM: normality, independence, constant variance
- Example two-level longitudinal model:



Next we will add predictors to account for each pile!