Calculating Effect Sizes for Single-Case Research

An Introduction to the SingleCaseES and scdhlm Web Applications and R Packages

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Outline

1. Organizing and curating data from single-case designs.

- 2. Within-study effect sizes.
 - Background
 - The SingleCaseES app
- 3. Between-case standardized mean differences
 - Background
 - \circ The scdhlm app

Organizing and curating data from single-case designs



Why organize and curate your data?

- 1. So that you can do statistical analysis and effect size calculations.
- 2. So that you can share your data.
 - Make it easily accessible for inclusion in systematic reviews!
- 3. Because graphing data usually involves loss of information.
- 4. To fully document your research study.

Tidy SCD data

- One column per variable
- One row per observation session
- Descriptive labels for
 - Case (participant)
 - Phase of design or treatment condition

Case	Phase	Session	Outcome
Deborah's Group	Baseline	1	62.63
Deborah's Group	Baseline	2	40.22
Deborah's Group	Baseline	3	54.26
Deborah's Group	Baseline	4	40.26
Deborah's Group	Baseline	5	46.82
Deborah's Group	Baseline	6	52.45
Deborah's Group	Intervention	7	25.37
Deborah's Group	Intervention	8	26.32
Deborah's Group	Intervention	9	7.65
Deborah's Group	Intervention	10	11.41
Deborah's Group	Intervention	11	13.30
Deborah's Group	Intervention	12	22.66
Deborah's Group	Intervention	13	13.34
Amy's Group	Baseline	1	16.67
Amy's Group	Baseline	2	28.43
Amy's Group	Baseline	3	29.41
Amy's Group	Baseline	4	30.39
Amy's Group	Baseline	5	45.10
Amy's Group	Baseline	6	37.25

6/49

Multiple dependent variables

• Wide format: Use separate columns for multiple outcome variables

Case	Phase	Session	Problem_Behavior	On_Task_Behavior
Deborah's Group	Baseline	1	16.7	56.7
Deborah's Group	Baseline	2	20.0	70.0
Deborah's Group	Baseline	3	26.7	66.7
Deborah's Group	Baseline	4	20.0	86.7
Deborah's Group	Baseline	5	16.7	56.7
Deborah's Group	Baseline	6	13.3	70.0
Deborah's Group	Intervention	7	16.7	46.7
Deborah's Group	Intervention	8	20.0	73.3
Deborah's Group	Intervention	9	20.0	56.7
Deborah's Group	Intervention	10	30.0	50.0
Deborah's Group	Intervention	11	30.0	63.3
Deborah's Group	Intervention	12	13.3	63.3

Multiple dependent variables

• Long format: One row per outcome measure per session

Case	Phase	Session	DV	Outcome
Deborah's Group	Baseline	1	On Task Behavior	56.7
Deborah's Group	Baseline	1	Problem Behavior	16.7
Deborah's Group	Baseline	2	On Task Behavior	70.0
Deborah's Group	Baseline	2	Problem Behavior	20.0
Deborah's Group	Baseline	3	On Task Behavior	66.7
Deborah's Group	Baseline	3	Problem Behavior	26.7
Deborah's Group	Baseline	4	On Task Behavior	86.7
Deborah's Group	Baseline	4	Problem Behavior	20.0
Deborah's Group	Baseline	5	On Task Behavior	56.7
Deborah's Group	Baseline	5	Problem Behavior	16.7
Deborah's Group	Baseline	6	On Task Behavior	70.0
Deborah's Group	Baseline	6	Problem Behavior	13.3

Adding more detail

- Add furthers details about what happened in the study.
- Some ideas:
 - Actual session date + times (YYYY-MM-DD-HH:MM)
 - Observation session lengths
 - Clinician/therapist IDs
 - Notes about events

Case	Phase	Session	Problem Behavior	On-Task Behavior	Date	Session length	Notes
Deborah's Group	Baseline	1	16.7	56.7			
Deborah's Group	Baseline	2	20.0	70.0			
Deborah's Group	Baseline	3	26.7	66.7			

Share Your Data!



Within-case effect size indices

IRL

LRR

Within-case effect size indices

- Single-number summary of the **direction** and **magnitude** of intervention effect (functional relation) *for each case* within a study.
- Use these if you want to:
 - Describe results separately for each participant
 - Examine heterogeneity of effects or associations with individual-level characteristics
 - Compare results across participants and SCED studies that use various outcome measures
- Lots of proposed effect size indices. Today we'll focus on
 - Non-overlap of all pairs
 - Within-case standardized mean difference
 - Log-response ratio

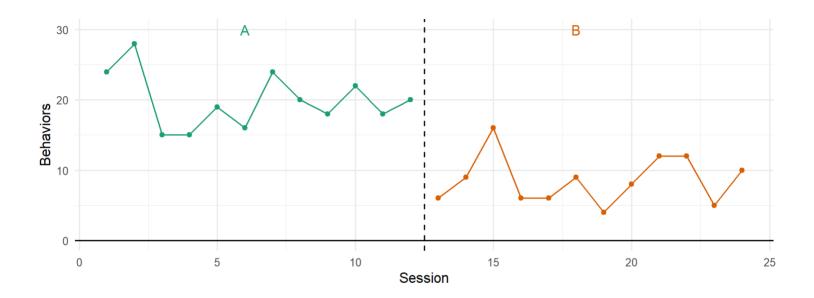
SingleCaseES: Single-series calculator

- Access SingleCaseES on the web at https://jepusto.shinyapps.io/SCD-effect-sizes/
- Or by opening RStudio and typing

```
library(SingleCaseES)
SCD_effect_sizes()
```

- Two parts to the app:
 - Single-series calculator (direct data entry)
 - Multiple-series calculator (using a data file)

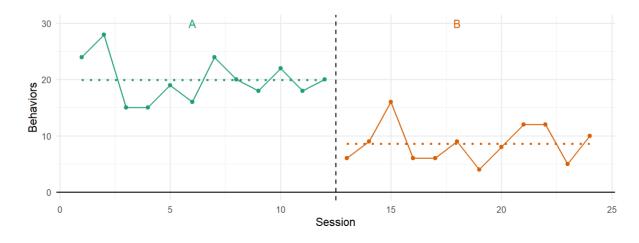
Simplest possible model



- Stable baseline and treatment phases (no time trends)
 - Immediate shift in level due to intervention
- Independence of outcome measurements

Notation

- n_A observations in phase A: $y_1^A, \ldots, y_{n_A}^A$
- n_B observations in phase B: $y_1^B, \ldots, y_{n_B}^B$
- Mean level of the outcome in each phase: μ_A, μ_B
 - $\circ~$ Estimated by sample means ${ar y}_A, {ar y}_B$
- Standard deviation of the outcome in each phase: σ_A, σ_B
 - $\circ~$ Estimated by sample standard deviations S_A, S_B



Non-overlap of all pairs

- Non-overlap measures are defined in terms of *ordinal comparisons* of outcomes
- Non-overlap of all pairs (Parker and Vannest, 2009) is defined in terms of all pairs of one observation from phase A and one observation from phase B.
- For every pair $i=1,\ldots,n_A$ and $j=1,\ldots,n_B$, take

$$q_{ij} = egin{cases} 1 & ext{if} & y^B_j ext{ better than } y^A_i \ rac{1}{2} & ext{if} & y^B_j = y^A_i \ 0 & ext{if} & y^B_j ext{ worse than } y^A_i \end{cases}$$

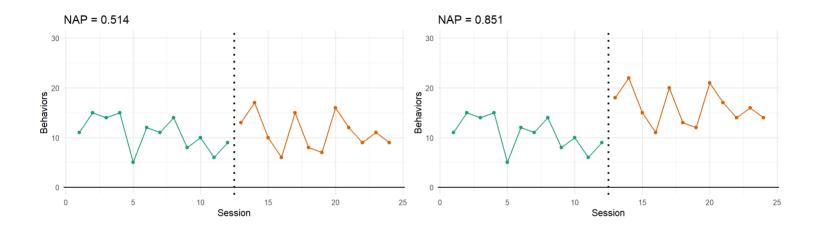
• NAP estimator:

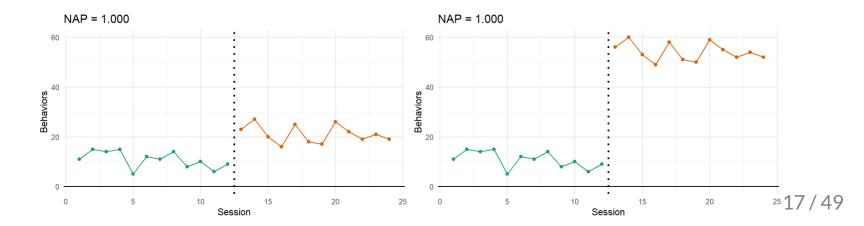
$$NAP = rac{1}{mn}\sum_{i=1}^{n_A}\sum_{j=1}^{n_B}q_{ij}$$

- Standard error based on unbiased estimator (Sen, 1967; Mee, 1990)
 - Methods assume that observations are independent and identically distributed within each phase.

Limited range of sensitivity

• Limited range where NAP (and other non-overlap measures) sensitive to change.





Within-case standardized mean difference

- Proposed by Gingerich (1984) and Busk and Serlin (1992)
- Parameter definition:

$$\delta = rac{\mu_B - \mu_A}{\sigma_A}$$

- Difference in means, "standardized" by baseline variation
- NOT equivalent to between-case SMD because σ_A includes only *within-case variation*.
- Appropriate for interval-scale outcomes
 - Is variability of outcomes approximately constant for different mean levels?
 - Standardizing by within-case variation means this measure will be strongly affected by reliability of measurements
 - Problematic for outcomes with restricted range in baseline

Within-case standardized mean difference: estimation

• Originally proposed estimator:

$$d=rac{{ar y}_B-{ar y}_A}{S_A}$$

• Estimator with small-sample bias correction:

$$g=\left(1-rac{3}{4n_A-5}
ight) imesrac{ar{y}_B-ar{y}_A}{S_A}$$

• Approximate standard error, *assuming independent observations*:

$$SE_g = \left(1 - rac{3}{4n_A - 5}
ight) \sqrt{rac{1}{n_A} + rac{S_B^2}{n_B S_A^2}} + rac{d^2}{2(n_A - 1)}$$

Proportional change in levels

- Percentage (proportional) change from baseline to intervention is an easily interpretable "informal" effect size (Campbell and Herzinger, 2010).
- The log response ratio is a formal measure of effect size that describes change in proportional terms (Pustejovsky, 2015; Pustejovsky, 2018).
- Parameter definition:

$$\psi = \log igg(rac{\mu_B}{\mu_A}igg)$$

- Appropriate for **ratio-scale** outcomes (frequency counts, percentage duration)
- Natural logarithm is used to make the range unrestricted.
- Transformation to percentage change:

$$\% ext{ change} = 100\% imes \left(e^{\psi} - 1
ight)$$

Log response ratio: estimation

• Basic estimator (biased if *m* or *n* is small):

$$R_1 = \logiggl(rac{{ ilde y}_B}{{ ilde y}_A}iggr)$$

• Bias-corrected estimator:

$$R_2 = \logiggl(rac{{ ilde y}_B}{{ ilde y}_A}iggr) + rac{{ ilde S}_B^2}{2n_B{ ilde y}_B^2} - rac{{ ilde S}_A^2}{2n_A{ ilde y}_A^2}$$

• Approximate standard error for R_2 , *assuming independent observations*:

$$SE_R = \sqrt{rac{{ ilde S}_A^2}{{n_A}{ ilde y}_A^2}} + rac{{ ilde S}_B^2}{{n_B}{ ilde y}_B^2}$$

Direction of improvement

- Two versions of LRR:
 - LRRi: Positive numbers represent increases in desirable outcomes
 - LRRd: Negative numbers represent decreases in undesirable outcomes
- Use the version that corresponds to *predominant valence* of outcomes in your data.
- For count outcomes, LRRi = -LRRd
- For proportion / percentage outcomes, the outcome valence is harmonized before calculation.
 - For proportion / percentage outcomes, LRRi \neq -LRRd

Truncation constants

- If ${ar y}_A=0$ or ${ar y}_B=0$ then LRR is undefined.

$$\circ \;$$
 If $S_A^2=0$ or $S_B^2=0$ then SE_R is undefined

• To handle such situations, the app uses **truncated mean** and **truncated SD** estimators:

$${ ilde y}_A = \max\left\{ {{ar y}_A},rac{1}{{2n_A}D}
ight\}, \qquad {{ ilde y}_B} = \max\left\{ {{ar y}_B},rac{1}{{2n_B}D}
ight\}$$

and

$${ ilde S}_A^2 = \max\left\{S_A^2, rac{1}{n_A^3 D^2}
ight\}, \qquad { ilde S}_B^2 = \max\left\{S_B^2, rac{1}{n_B^3 D^2}
ight\}$$

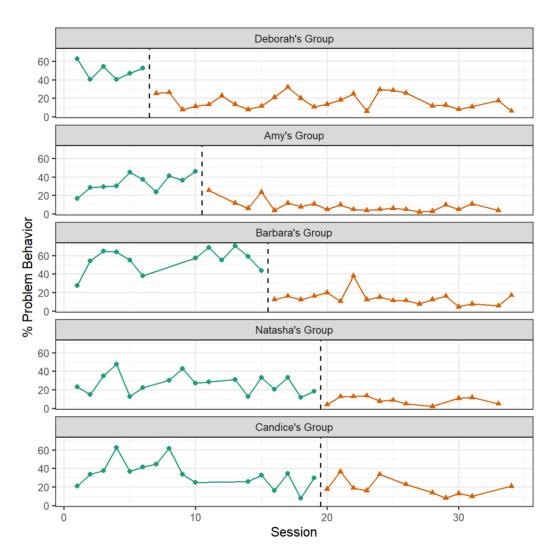
- *D* is a constant that depends on the outcome scale and measurement procedures
 - Number of intervals / items
 - Session length for direct observation
 - \circ Can also define your own D

SingleCaseES: Multiple-series calculator

- Basic walk-through with data from Rodriguez and Anderson (2014)
- Calculating phase-pairs in ABAB designs
- Aggregating effect sizes

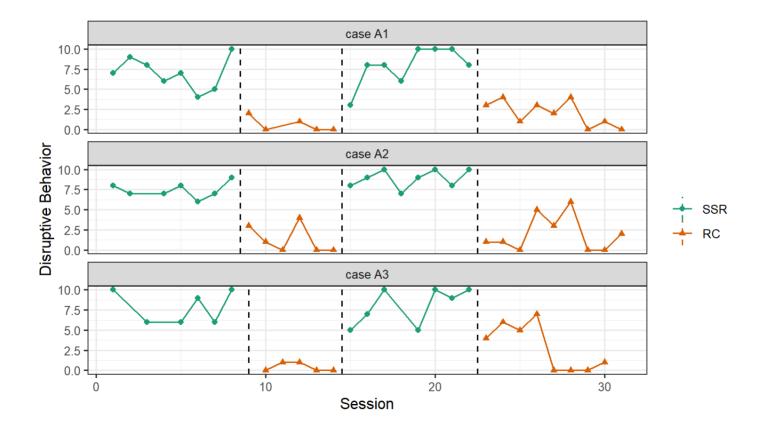
Rodriguez and Anderson (2014)

Integrating a social behavior intervention during small group academic instruction using a total group criterion intervention



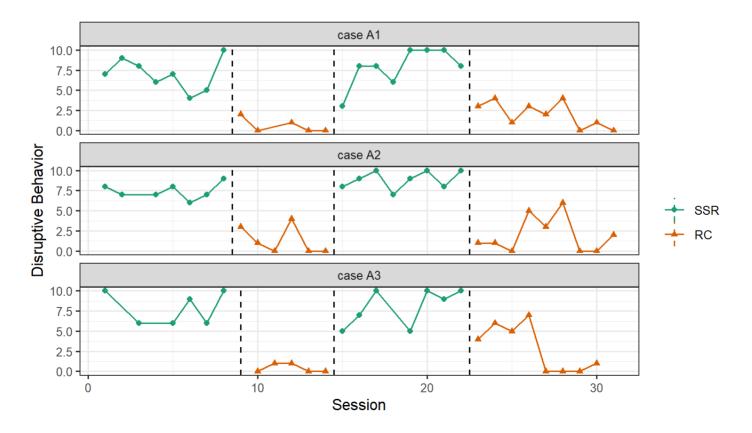
Lambert, Cartledge, Heward et al. (2006)

Effects of response cards on disruptive behavior and academic responding during math lessons by fourth-grade urban students



Calculating phase pairs

- Might want to calculate effect sizes for adjacent pairs of baseline and intervention phases.
- SingleCaseES provides an option to determine phase pairs automatically.



Aggregating effect sizes

- After calculating effect sizes for adjacent pairs of phases, we might want to **average them together** to simplify reporting or further analysis.
 - Average across phase pairs in an ABAB design
 - Average across cases to generate an overall summary effect size estimate
- Several options for taking weighted averages
 - Equal weighting
 - Inverse-variance weighting: $\frac{1}{V}$ (use for LRR)
 - $\circ n_A$
 - $\circ~n_B$
 - $\circ n_A n_B$ (use for NAP)
 - $\circ \ rac{1}{n_A} + rac{1}{n_B}$ (use for SMD)

Replication code

```
# Load packages
library(SingleCaseES)
# Load data
library(readx1)
library(janitor)
dat <-
 read_excel(path = "Small-is-Beautiful-effect-size-workshop.xlsx", sheet = "Lambert") %>%
 clean_names(case = "parsed")
# clean data
library(dplyr)
dat <-
 dat %>%
 group_by(case) %>%
 mutate(phase_pair_calculated = calc_phase_pairs(treatment, session = day)) %>%
 ungroup()
# Batch calculation
res <- batch_calc_ES(dat = dat,</pre>
                     grouping = c(case),
                     condition = treatment,
                     outcome = outcome,
                     aggregate = c(phase_pair_calculated),
                     weighting = "1/nA + 1/nB",
                     session_number = day,
                     baseline_phase = "SSR",
```

Between-case standardized mean differences



Premises

- **Goal**: Estimate an effect size using data from a single-case design that is *in the same metric* as the standardized mean difference effect size from a between-group experimental design.
- Why? (Shadish, Hedges, Horner et al., 2015)
 - **Translation** of single-case research for researchers who work primarily with between-groups designs
 - **Comparison** of results from single-case studies and betweengroups studies, for purposes of understanding the utility and limitations of each type of design
 - **Synthesis** involving both single-case and between-groups designs

SMD in between-group experiment

• What is the SMD from a between-group experiment?

$$\delta_{BC} = \frac{\begin{pmatrix} \text{Average outcome if} \\ \text{everybody gets intervention} \end{pmatrix} - \begin{pmatrix} \text{Average outcome if} \\ \text{nobody gets intervention} \end{pmatrix}}{\begin{pmatrix} \text{SD of outcome if} \\ \text{nobody gets intervention} \end{pmatrix}}$$
$$\delta_{BC} = \frac{\begin{pmatrix} \text{Average outcome if} \\ \text{everybody gets intervention} \end{pmatrix} - \begin{pmatrix} \text{Average outcome if} \\ \text{nobody gets intervention} \end{pmatrix}}{\sqrt{\begin{pmatrix} \text{Between-participant} \\ \text{variance} \end{pmatrix} + \begin{pmatrix} \text{Within-participant} \\ \text{variance} \end{pmatrix}}}$$

• We aim to estimate these component quantities using data from a single-case experimental design.

The broad strategy

(Pustejovsky, Hedges, and Shadish, 2014) described a general strategy for estimating BC-SMD:

- 1. Develop a hierarchical linear model that describes:
 - The form of time trends and intervention effects
 - How the trends and intervention effects vary across participants
- 2. Imagine a **hypothetical between-group experiment** with the same population of participants, same intervention, same dependent variable.
 - When is treatment initiated?
 - When are outcomes assessed?
- 3. Use the hierarchal model to estimate the components of δ_{BC} for the hypothetical experiment.
- 4. Make a small-sample correction (similar to Hedges' g)

Design translation

A multiple baseline across participants:

T1	Т2	Т3	T4	T5	Т6	Т7	Т8	Т9	T10	T11	T12	T13	T14	T15	T16	T17	T18
Х	Х	Х	Х	XT	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х
Х	Х	Х	Х	Х	Х	Х	XT	Х	Х	Х	Х	Х	Х	Х	Х	Х	Х
Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	XT	Х	Х	Х	Х	Х	Х	Х
Х	Х	Х	Х	Х	Х	Х	Х	Х	Х	х	Х	Х	ХТ	Х	Х	Х	Х

A hypothetical between-group design (with pre-test):

Т1	Т2	Т3	T4	Т5	Т6	Т7	Т8	Т9	T10	T11	T12	T13	T14	T15	T16	T17	T18	
				ХТ											Х			
				ХТ										Х				
				Х											Х			
				Х											Х			

Overview of methods literature

- BC-SMD estimators for a basic hierarchical linear model with no time trends:
 - Hedges, Pustejovsky, and Shadish (2012): Treatment reversal (ABAB) design replicated across 3+ participants
 - Hedges, Pustejovsky, and Shadish (2013): Multiple baseline / multiple probe design with 3+ participants
 - Shadish, Hedges, and Pustejovsky (2014): More worked examples
- Pustejovsky, Hedges, and Shadish (2014) described a more general strategy for multiple baseline / multiple probe designs across participants
 - Valentine, Tanner-Smith, Pustejovsky et al. (2016): Tutorial and practical guidance
- Swaminathan, Rogers, and Horner (2014) proposed Bayesian estimation methods
- Chen, Pustejovsky, Klingbeil et al. (2023) proposed BC-SMD methods for more complex designs:
 - Multiple baseline across behaviors, replicated across 3+ participants
 - Clustered multiple baseline design across participants (3+ clusters)
 - Multivariate multiple baseline design across 3+ participants

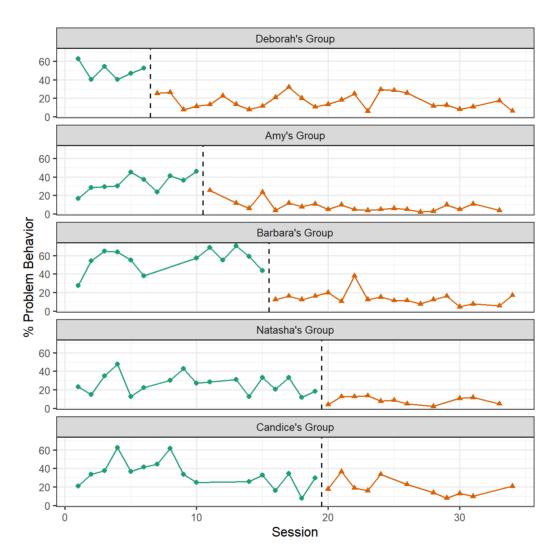
scdhlm web app

- Access scdhlm on the web at https://jepusto.shinyapps.io/scdhlm/
- Or by opening RStudio and typing

library(scdhlm)
shine_scd()

Rodriguez and Anderson (2014)

Integrating a social behavior intervention during small group academic instruction using a total group criterion intervention



The most basic HLM

• Level-1 model for each participant:

$$Y_{ij}=eta_{0j}+eta_{1j}(Tx)_{ij}+e_{ij}$$

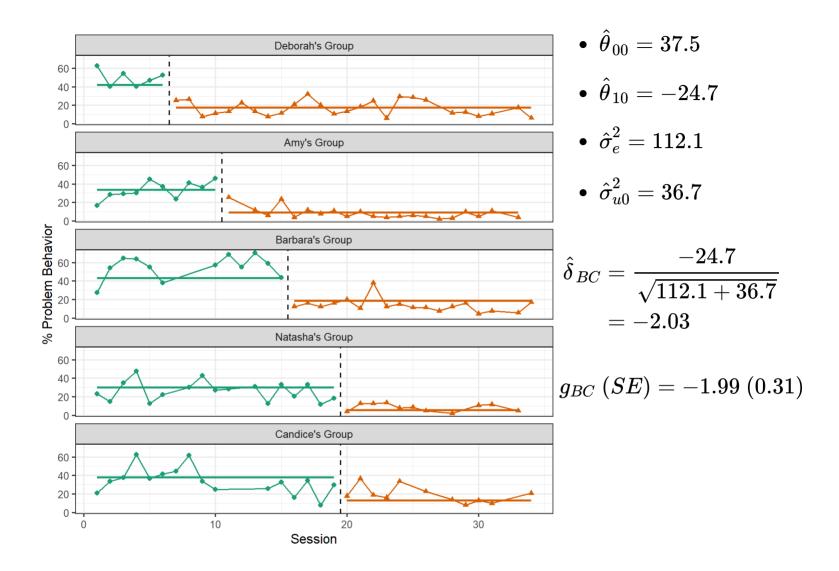
where $\operatorname{Var}(e_{ij}) = \sigma_e^2$ and $e_{1j}, \ldots, e_{Tj} \sim AR_1(\phi)$

• Level-2 model:

$$egin{aligned} eta_{0j} &= heta_{00} + u_{0j}, \qquad u_{0j} \sim N(0, \sigma_{u0}^2) \ eta_{1j} &= heta_{10} \end{aligned}$$

- Under this model:
 - \circ Average outcome if nobody gets intervention: $heta_{00}$
 - $\circ~$ Average outcome if everybody gets intervention: $heta_{00}+ heta_{10}$
 - $\circ~$ SD of outcome if nobody gets intervention: $\sqrt{\sigma_{u0}^2+\sigma_e^2}$

• BC-SMD effect size:
$$\delta_{BC} = \frac{\theta_{10}}{\sqrt{\sigma_{u0}^2 + \sigma_e^2}}$$
 38 / 49



A more flexible HLM

• Level-1 model for each participant:

$$Y_{ij}=eta_{0j}+eta_{1j}(\mathrm{Time})_{ij}+eta_{2j}(Tx)_{ij}+eta_{3j}(Tx)_{ij} imesig((\mathrm{Time})_{ij}-k_jig)+e_{ij}$$
 where $\mathrm{Var}(e_{ij})=\sigma_e^2$ and $e_{1j},\ldots,e_{Tj}\sim AR_1(\phi)$ and k_j is last baseline session.

• Level-2 model:

$$egin{aligned} eta_{0j} &= heta_{00} + u_{0j}, & u_{0j} \sim N(0, \sigma_{u0}^2) \ eta_{1j} &= heta_{10} + u_{1j}? \ eta_{2j} &= heta_{20} + u_{2j}? \ eta_{3j} &= heta_{30} + u_{3j}? \end{aligned}$$

- $\circ~$ Adding a random effect \rightarrow allowing slope / Tx effect to vary across cases
- $\circ~$ Omitting a random effect \rightarrow assuming slope / Tx effect is *constant*
- Models with more random effects require more cases

A more flexible HLM

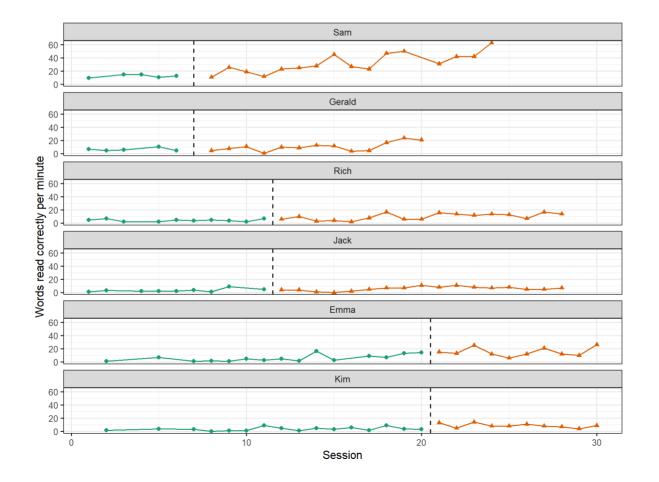
- Can also modify assumptions about level-1 errors
- Different variance by treatment phase:

$$ext{Var}(e_{ij}) = egin{cases} \sigma_{eC}^2 & ext{if} & (Tx)_{ij} = 0 \ \sigma_{eT}^2 & ext{if} & (Tx)_{ij} = 1 \end{cases}$$

- Correlation structure of level-1 errors:
 - $\circ\;$ First order auto-regression $(AR_1(\phi))$
 - $\circ\;$ First order moving average $(MA_1(\phi))$
 - \circ Independent

Model should be informed by theoretical expectations and visual inspection

Barton-Arwood, Wehby, and Falk (2005) Reading instruction for elementary-age students with emotional and behavioral disorders: Academic and behavioral outcomes



42/49

Models with time trends

- For models with time trends, we need to specify *timing* of pre-test and post-test for the hypothetical between-group design.
- **Initial treatment time**: Last session of baseline phase before being assigned to intervention or comparison condition.
 - Default: Length of shortest baseline phase
- Focal follow-up time: Session during which outcomes would be assessed in hypothetical experiment.
 - Default: Last measurement occasion for first case to enter intervention
 - This is not a particularly good default
 - Ideally, pick a focal follow-up time based on a meaningful or typical treatment duration

Barton-Arwood, Wehby, and Falk (2005) effect size calculations

- Model specification
 - Baseline level (random)
 - Baseline time trends (constant)
 - Treatment level change (constant)
 - Treatment trend change (random)
 - Level-1 variance differs by phase
- Initial treatment time: After 6 sessions
- Focal follow-up time of session 16 (10 sessions of treatment).
- BC-SMD estimate:

$$g_{BC} \; (SE) = 0.82 \; (0.75)$$

Illustrative application of BC-SMDs

- Calder and colleagues (2020, 2021) studied an explicit grammar instruction intervention for children with developmental language disorder.
- Calder, Claessen, Ebbels et al. (2020): multiple baseline across nine participants
 - Data available in the Excel workbook
 - Try calculating a BC-SMD estimate after 10 weeks of intervention
- Calder, Claessen, Ebbels et al. (2021): crossover randomized trial with ${\cal N}=21$ participants
 - 10 weekly intervention sessions
 - $\circ ~g=1.97, SE=0.11$ for expressive morphosyntax
 - $\circ g = 0.06, SE = 0.06$ for grammaticality judgements

Limitations of between-case SMD

- Tool for translating from single-case logic to group-design logic.
 - Premised on the idea that a hypothetical group design is theoretically plausible
- Describes a summary, average effect across a set of cases
 - Potentially concealing individual-level heterogeneity
- For some models, magnitude depends on the features (timing) of hypothetical between-group design
- Technical limitations
 - Only available for some designs
 - Requires at least 3 participants (preferably more!)
 - Models assume normal (Gaussian) errors
 - Care needed for model selection

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