

Sample Size Planning for MLM

PSYC 575

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Updated: 2022-10-02

Week Learning Objectives

- Describe the importance of having sufficient sample size for scientific research
- Describe conceptually the steps for sample size planning: precision analysis and power analysis
- Perform power analysis for MLM using the PowerUpR application and the `simr` package
- Understand the effect of uncertainty in parameter values and explore alternative approaches for sample size planning

Why Sample Size?

Small Sample Size is a Problem Because . . .

Low power

Misleading and noisy results¹

- When coupled with publication bias (statistical significance filter)^{2 3}

Nonreproducible findings

[1] See [Maxwell \(2004\)](#)

[2] See the graph on [this blog post](#)

[3] See also [Vasishth et al. \(2018\)](#)

Review: Sampling distributions

What is the null distribution?

- Suppose we examine the effect of a therapy on eating disorder
- We test against the null hypothesis $H_0 : \gamma_{01} = 0$, where γ_{01} is the fixed effect of the therapy on eating disorder

What is the alternative distribution?

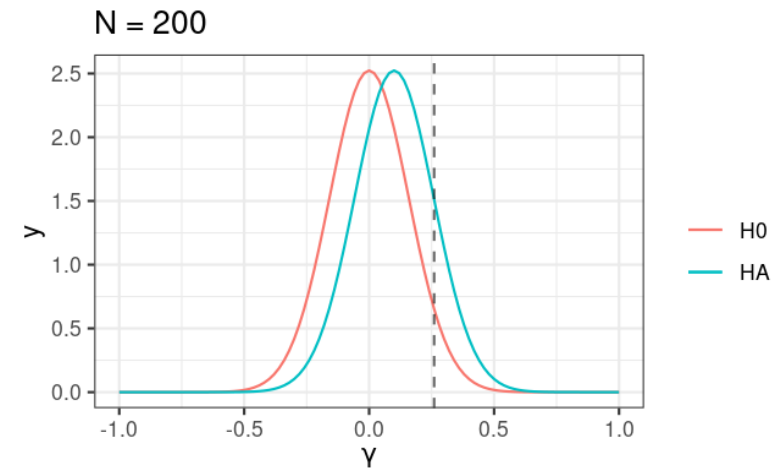
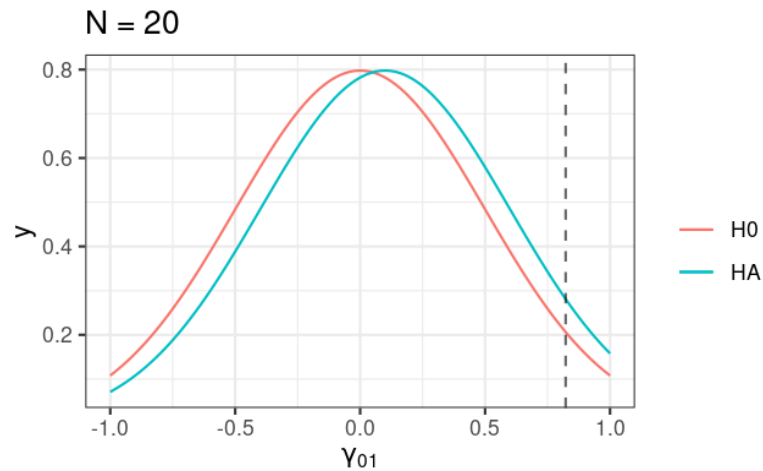
- Assume that the true effect of this therapy is $\gamma_{01} = .1$

Sampling Distribution as a Function of Sample Size

Assume true effect is $\gamma_{01} = 0.10$

Let's say

- when $N = 20$, $p < .05$ when $\hat{\gamma} \geq 0.82$
- when $N = 200$, $p < .05$ when $\hat{\gamma} \geq 0.26$



Steps for Sample Size Planning

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1. Write down your model equations
2. List out all parameters in the model
3. Determine if you want to achieve a desired level of
 - a. Power, or
 - b. Precision

Step 1: Write down model equations

Group-based therapy for eating disorder (cluster-randomized trial)

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Level-1

$$Y_{ij} = \beta_{0j} + \beta_{1j}X_{cmc_{ij}} + e_{ij}$$

$$e_{ij} \sim N(0, \sigma)$$

Level-2

$$\beta_{0j} = \gamma_{00} + \gamma_{01}W_j + u_{0j}$$

$$\beta_{1j} = \gamma_{10} + \gamma_{11}W_j + u_{1j}$$

$$\begin{bmatrix} u_{0j} \\ u_{1j} \end{bmatrix} \sim N \left(\begin{bmatrix} 0 \\ 0 \end{bmatrix}, \begin{bmatrix} \tau_0^2 & \\ \tau_{01} & \tau_1^2 \end{bmatrix} \right)$$

- γ_{10} : X (purely level-1 with ICC = 0)
- γ_{01} : W (level-2)
- γ_{11} : $W \times X$ (cross-level interaction)

Step 2: List out all parameters

1. Fixed effects: $\gamma_{00}, \gamma_{01}, \gamma_{10}, \gamma_{11}$

2. Random effects: $\tau_0^2, \tau_1^2, \tau_{01}$

3. Number of clusters: J

4. Cluster size: n

Level-1

$$Y_{ij} = \beta_{0j} + \beta_{1j}X_{cmc_{ij}} + e_{ij}$$

$$e_{ij} \sim N(0, \sigma)$$

Level-2

$$\beta_{0j} = \gamma_{00} + \gamma_{01}W_j + u_{0j}$$

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$$\begin{bmatrix} u_{0j} \\ u_{1j} \end{bmatrix} \sim N \left(\begin{bmatrix} 0 \\ 0 \end{bmatrix}, \begin{bmatrix} \tau_0^2 & \\ & \tau_1^2 \end{bmatrix} \right)$$

Standard Error and Precision Analysis

Sample Size and $SE/Post. SD$

In the previous graph, when $N = 20$, the sample estimate is likely to be anywhere between -0.4 and 0.6

$$SE \propto \frac{1}{\sqrt{N}}$$

One goal of sample size planning is to

- Have sufficient sample size to get precise (low SE) sample estimates of an effect

Analytic Formulas of SE

J = Number of clusters; n = Cluster size

- E.g., $J = 100$ schools; $n = 10$ students per school

Assuming $\tau_{01} = 0$

$$SE(\gamma_{01}) = \sqrt{\frac{1}{S_W^2} \left(\frac{\tau_0^2}{J} + \frac{\sigma^2}{Jn} \right)}$$

$$SE(\gamma_{10}) = \sqrt{\frac{\tau_1^2}{J} + \frac{\sigma^2}{JnS_X^2}}$$

$$SE(\gamma_{11}) = \sqrt{\frac{1}{S_W^2} \left(\frac{\tau_1^2}{J} + \frac{\sigma^2}{JnS_X^2} \right)}$$

Precision Analysis

Group-based therapy for eating disorder (cluster-randomized trial)

- Intervention at group level
- 10 participants per group
- Outcome standardized (i.e., $SD = \sqrt{\tau_0^2 + \sigma^2} = 1$)
 - $\gamma = \text{Cohen's } d$
- ICC = .3 (i.e., $\tau_0^2 = .3$)
- Goal: estimate J such that $SE(\gamma_{10}) \leq .1$
 - E.g., if we estimated the sample effect size to be $d = .25$, the 95% CI would be approximately [.05, .45].

Calculating J

When the predictor is binary (e.g., treatment-control), if half of the groups is in one condition, $S_W^2 = 0.25$

- Otherwise, if 30% in one condition, $S_W^2 = 0.3 \times 0.7$
- $\tau_0^2 = 0.3, \sigma^2 = 0.7, n = 10$

E.g., if $J = 30$

$$SE(\gamma_{01}) = \sqrt{\frac{1}{S_W^2} \left(\frac{\tau_0^2}{J} + \frac{\sigma^2}{Jn} \right)} = \sqrt{\frac{1}{0.25} \left(\frac{0.3}{30} + \frac{0.7}{(30)(10)} \right)} = 0.222$$

Keep trying, and you'll find ...

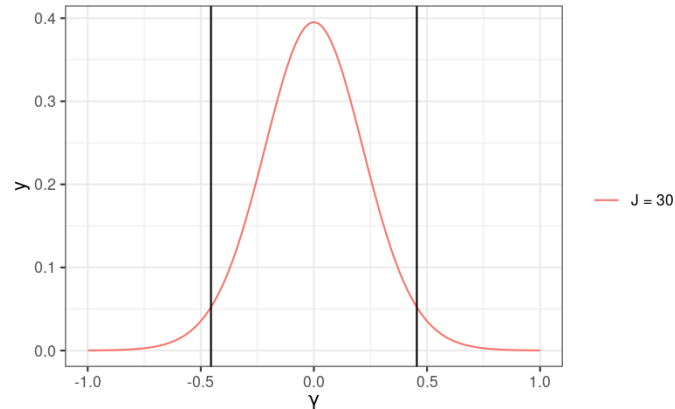
When $J = 148$, $SE(\gamma_{01}) = 0.1$

So you'll need 148 groups (74 treatment, 74 control)

Power Analysis

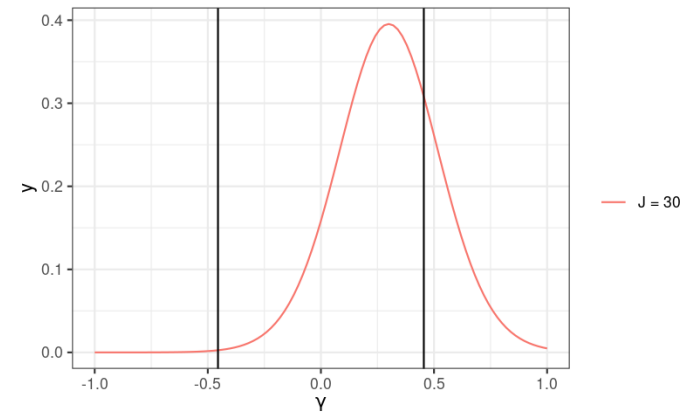
Two-tailed test, $\alpha = .05$

$$H_0 : \gamma_{01} = 0$$



Critical region: $\hat{\gamma}_{01} \leq -0.45$ or $\hat{\gamma}_{01} \geq 0.45$

$$H_1 : \gamma_{01} = 0.3$$



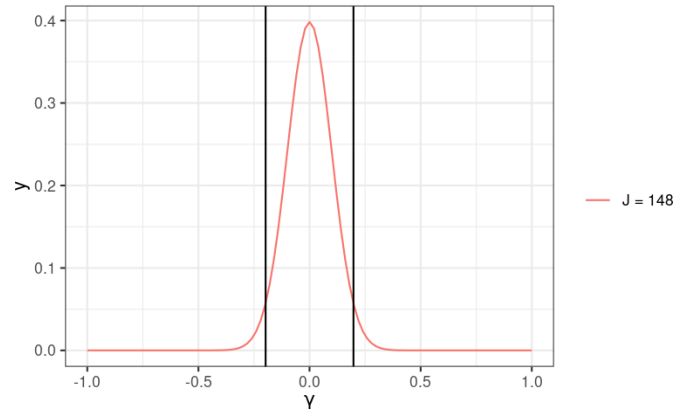
Power¹

$$\approx P(\hat{\gamma}_{01} \leq -0.45) + P(\hat{\gamma}_{01} \geq 0.45) = 0.247$$

[1] In practice, we need to incorporate the sampling variability of the standard error as well, so this power calculation is only a rough approximation.

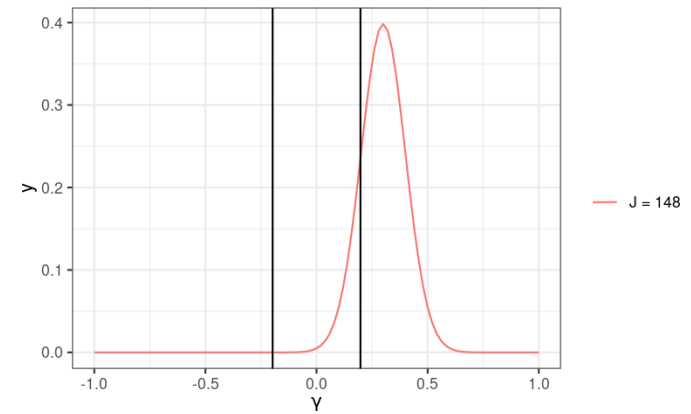
Two-tailed test, $\alpha = .05$

$$H_0 : \gamma_{01} = 0$$



Critical region: $\hat{\gamma}_{01} \leq -0.2$ or $\hat{\gamma}_{01} \geq 0.2$

$$H_1 : \gamma_{01} = 0.3$$



Power

$$\approx P(\hat{\gamma}_{01} \leq -0.2) + P(\hat{\gamma}_{01} \geq 0.2) = 0.846$$

Tools for Power Analysis

1. Stand-alone programs

- [Optimal Design](#)
- [PinT](#)

2. R packages

- `simr`

3. Spreadsheet/Webapp

- [PowerUp!](#)

See more discussion in [Arend & Schäfer \(2019\)](#)

PowerUpR Shiny App

<https://powerupr.shinyapps.io/index/>

Monte Carlo Simulation for Power Analysis

- Simulate a large number (e.g., $R = 1,000$) of data sets based on given effect size, ICC, etc
- Fit an MLM to each simulated data
- Power \approx Proportion of times $p < \alpha$

See sample R code for using `simr`

Uncertainty in Parameter Values

Uncertainty in Parameter Values

In the PowerUpR demo, to calculate the number of clusters J need to achieve 80% power, we determined

1. Type I error rate = .05
2. Two tailed test = TRUE
3. $g_2, r_{21}, r_{22} = 0$, as we did not include any covariates
4. $p = .5$, for a balanced design (half treatment, half control)

However, we need to guess the values of

1. Effect size = .3?
2. ICC = .3?

The Effect of Uncertainty in Power

Ignoring uncertainty

- The more uncertainty we have but ignore about a parameter value, the more power loss we will have in our study (red curve)
- Uncertainty in both effect size and ICC can further reduce our power
- The more uncertainty we have, the more samples we need to achieve 80% power



Hybrid Classical-Bayesian approach

- Incorporates uncertainty for sample size planning
- Instead of plugging in a point value of a guess, we can specify how much uncertainty we have (e.g., standard error of γ_{01} from a previous study)

$$\delta \sim N(.3, .1) \quad \rho \sim \text{Beta}(a, b)$$

- where a, b can be calculated by $\hat{\rho} = .3$ and $\sigma_\rho = .1$ (estimate and uncertainty about ρ)

hcbR Shiny App

http://winnie-wy-tse.shinyapps.io/hcb_shiny

Additional Notes on Power

- Increasing J usually leads to higher power than increasing n
- Balanced designs generally have higher power than unbalanced designs
- Larger sample size required for testing level-2 predictors
- Testing an interaction requires a much larger sample size
 - E.g., 16 times larger than for a main effect