



EDUCATION

# **Authentic Power Calculations for RD Studies**

**J.R. Lockwood**

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# Power in a Randomized Experiment

$$d\sqrt{n} \approx 4$$

- $d$ : standardized effect size
- $n$ : # units in each arm
- Can approximately handle most issues by fiddling with  $n$ ; e.g.
  - Clustering: replace  $n$  with  $ESS = \frac{n}{DEFF}$
  - Covariates: replace  $n$  with  $ESS = \frac{n}{1-R^2}$
  - Imbalance: replace  $n$  with  $ESS = 4np(1-p)$

## Why is Power in RD Worse?

- $S$  = “forcing variable”
- $T$  = treatment =  $1\{S < 0\}$  (WLOG)
- Power degraded due to collinearity between  $S$  and  $T$ 
  - e.g. if  $S$  is uniform and  $T$  is split at the midpoint,  $R_{ST}^2 = 0.75$ 
    - Variance inflation is  $\frac{1}{1-R_{ST}^2} = 4$
    - → sample size required for power equivalent to randomized experiment is 4 times larger
    - Equivalently, minimum detectable effect for equivalent sample size is 2 times larger

# Why is Assessing RD Power More Challenging?

Primarily because power is affected by:

- Shape of distribution of  $S$  and where cutoff  $c$  determining  $T$  is in that distribution
  - Schochet (2008) provides clear description
- Estimators for  $E(Y|c^-)$  and  $E(Y|c^+)$  might be complex and may involve data-dependent data restrictions
  - E.g. Cross-validation choice of bandwidth (Ludwig and Miller, 2005; Imbens and Lemieux, 2008) or simultaneous choice of bandwidth and model complexity (Kirby, McCombs, and Mariano, 2009)
- Other complications like fuzziness and clustering exacerbate these issues

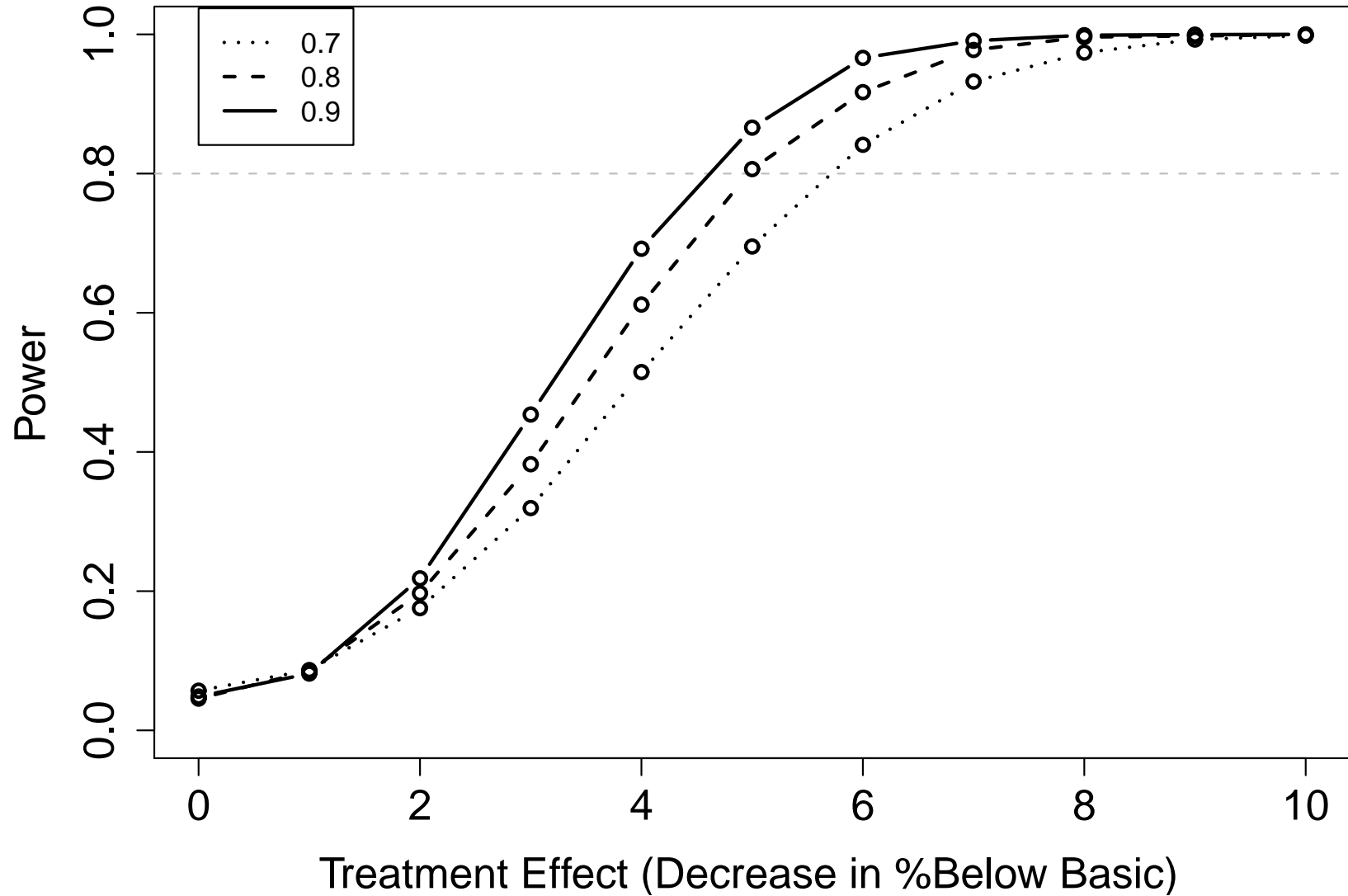
## Simulation as an Alternative Approach

- Often know a lot about data during design of RD studies
  - “Happenstance RD”: May have actual values of  $S$  and  $T$  and past values of  $Y$  (e.g. NCLB, RTTT)
  - “Designed RD”: Will know how you intend to construct  $S$  and  $T$  and again probably have good proxies for  $Y$
- Rather than trying to map knowledge about the data into power formulas, use knowledge about the data to simulate outcomes and analysis procedure

# Sketch of Approach

- $\beta$ : True treatment effect
- $D(\beta)$ : Simulated data, depends on  $\beta$
- $\hat{\beta}(D)$ : Estimated treatment effect, depends on  $D$ 
  - “Black Box” - make it as complicated as analysis will be
- Step 1: Estimate distribution of  $\hat{\beta}(D)$  given  $\beta = 0$ 
  - Use this to determine rejection region  $R$
- Step 2: Estimate  $Pr\{\hat{\beta}(D) \in R\}$  for selected sequence of alternatives  $\beta$ 
  - “Outer” loop: sequence of  $\beta$
  - “Inner” loop:  $M$  Monte Carlo iterations and count how often estimated effect is in rejection region

# Example Output



# Advantages of Simulation Approach

- Anything can be inserted in the analysis no matter how hard it would be to examine analytically; e.g.
  - Cluster corrections with imbalanced samples, including the use of random effects models to aid efficiency
  - Complex model selection criteria, such as bandwidth and functional form choice via cross-validation
- No need to agonize over what is meant by an “effect size” in RD - outcomes of simulation study get reported on the natural scale of the outcome measure
- Simulation approach naturally provides power curves rather than MDE at a single value of power (e.g. 0.80) which is more informative



# Conclusions

- RD is unlike a randomized experiment because careful statistical model selection and specification is inherent to obtaining valid impact estimates
  - i.e. in RD there is generally not a simple, analytically tractable procedure that will provide a compelling estimate.
- As standard practice for RD becomes more sophisticated (e.g. by WWC standards setting a high bar), simple formulas are less likely to provide authentic assessments of power
- Simulation is a defensible and relatively easy alternative
  - And can benefit from the fact that very specific data is often available during the design phase