

Detecting and Quantifying Variation In Effects of Program Assignment (ITT)

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This Session

Goal: To *illustrate* and *integrate* key concepts

Topics

- Defining variation in program effects
- Detecting and quantifying this variation

Empirical Examples

- A secondary analysis of three MDRC work/welfare studies
(59 sites with 1,176 individuals randomized per site, on average)
- A secondary analysis of the National Head Start Impact Study
(198 sites with 19 individuals randomized per site, on average)

Reference

- Bloom, H.S., S.W. Raudenbush, M.J. Weiss and K. Porter (conditional acceptance) *Journal of Research on Educational Effectiveness*.

Part I

Defining Individual Variation in Program Effects

Distribution of Individual Program Effects

Individual potential outcomes

$Y_i(1)$ = outcome with treatment

$Y_i(0)$ = outcome without treatment

Individual program effect

$$B_i = Y_i(1) - Y_i(0)$$

Population mean program effect

$$E(B) = E(Y(1) - Y(0))$$

Population program effect variance

$$\text{Var}(B) = \text{Var}(Y(1) - Y(0))$$

Population program effect distribution = ????

Distribution of Individual Program Effects

(continued)

The fundamental barrier to observing a program effect distribution for individuals

- One can only observe an outcome with a program or without the program for a given individual at a given time.
- Hence it is not possible to observe individual program effects
- Therefore one can only infer a distribution of individual program effects based on assumptions.

The fundamental barrier to estimating a variance of program effects for individuals

- The effect of a program on an outcome variance is not necessarily the same as the variance of the program effects.
- To see this, note that:

$$Y_i(1) = Y_i(0) + B_i$$

$$\text{Var}(Y(1)) = \text{Var}(Y(0)) + \text{Var}(B) + 2\text{Cov}(B, Y(0))$$

and

$$\text{Var}(Y(1)) - \text{Var}(Y(0)) = \mathbf{Var}(b) + 2\text{COV}(B, Y(0))$$

Some Implications of Individual Impact Variation For the National Head Start Impact Study

Estimated Parameter	Cognitive Outcome Measure	
	Receptive Vocabulary (PPVT)	Early Reading (WJ/LW)
<u>Mean Effect Size</u>		
For full sample	0.15***	0.16***
For lowest pretest quartile	0.16***	0.17***
For other sample members	0.08*	0.13**
<u>Individual Residual outcome variance</u> (in original units)		
Treatment group	545***	433***
Control group	667***	440***

NOTES: The full sample size varies by outcome from about 3500 to 3700 children and includes both three and four year olds. The statistical significance of individual estimates is indicated as * \leq 10 percent, ** \leq 5 percent and *** \leq 1 percent. Estimates that differ statistically significantly across subgroups at the 0.10 level are indicated in bold.

Part II

Defining, Identifying, Estimating and Reporting Cross-site Variation in Program Effects

A Cross-Site Distribution of Mean Program Effects

Theoretical Model

Level One: Individuals

$$Y_{ij} = A_j + B_j T_{ij} + e_{ij}$$

Level Two: Sites

$$A_j = \alpha + a_j$$

$$B_j = \beta + b_j$$

where:

Y_{ij} = the outcome for individual i from site j ,

T_{ij} = one if individual i from site j was assigned to the program and zero otherwise,

A_j = the site j population mean control group outcome,

B_j = the site j population mean program effect,

e_{ij} = a random error that varies across individuals with a zero mean and a variance that can differ between treatment and control group members

β = the cross-site grand mean program effect,

b_j = a random error that varies across sites with zero mean and variance $\tau_b^2 = \tau_B^2$

α and a_j = the cross-site grand mean control group outcome and a random error that varies across sites with zero mean and variance τ_a^2 , respectively

Some Important Goals of a Cross-Site Analysis

Goal #1

Estimate the cross-site grand mean program effect

Goal #2

Estimate the cross-site standard deviation of program effects

Goal #3

Estimate the cross-site distribution of program effects

Goal #4

Estimate the difference in mean program effects between two categories of sites (the simplest possible moderator analysis).

Goal #5

Estimate the mean program effect for each site

Estimating Impact Variation across Randomized Blocks¹

Identification strategy

- Randomizing individuals within a “block” to treatment or control status provides unbiased estimates of the mean program effect for each block.
- This makes it possible to estimate program effect variation across blocks.
- Blocks can be studies, sites, cohorts or portions of the preceding.

Important distinctions

- Effects of program assignment vs. effects of program participation
- Variation in effects vs. variation in effect estimates

¹ By definition, randomized blocks have subjects randomized within them. When entire blocks are randomized they typically are called *clusters*.

Cross-site Variation in Impacts
vs.
Cross-site Variation in Impact Estimates

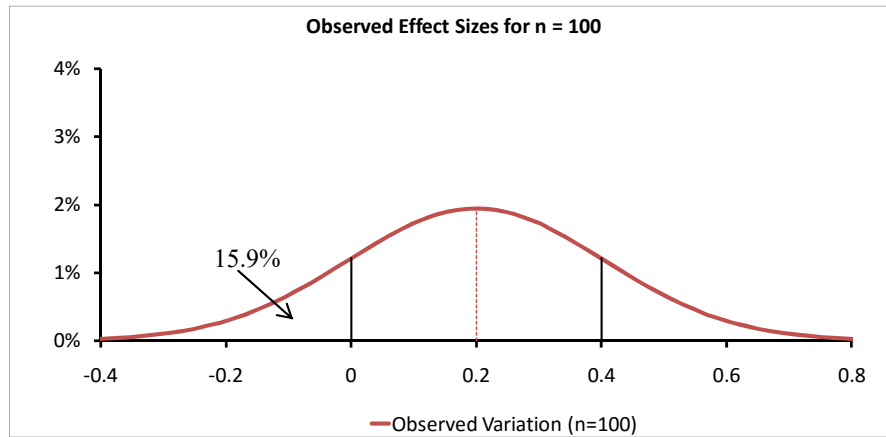
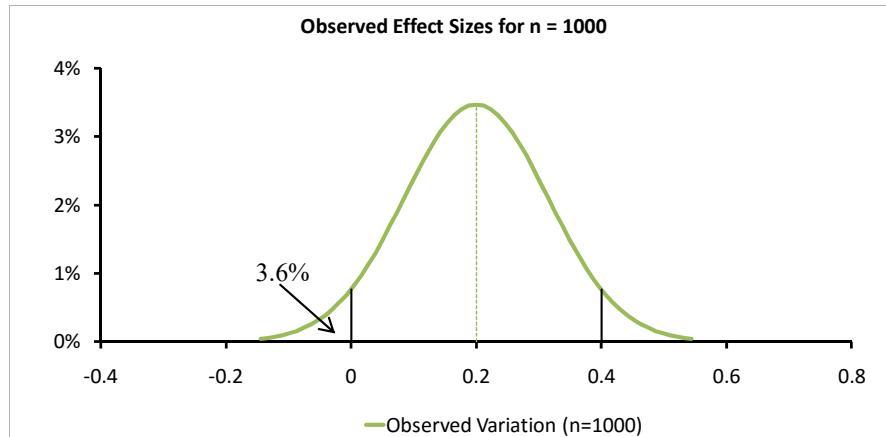
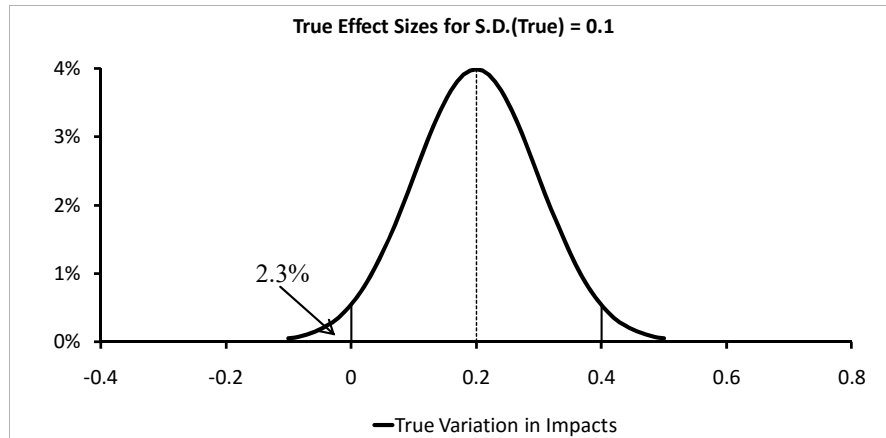
For Impact Estimation

$$\begin{aligned}\mathbf{Var}(\text{impact estimates}) &= \mathbf{Var}(\text{impacts}) + \mathbf{Var}(\text{impact estimation error}) \\ &= \tau_B^2 + V_j\end{aligned}$$

$$\mathbf{Reliability}(\text{impact estimates}) = \mathbf{Var}(\text{impacts}) / \mathbf{Var}(\text{impact estimates})$$

$$= \frac{\tau_B^2}{\tau_B^2 + V_j}$$

Figure 1



Estimation Model: FIRC

Fixed Site-Specific Intercepts, Random Site-Specific Program Effects and Separate Level-One Residual Variances for Ts and Cs (When necessary)

Level One: Individuals

$$Y_{ij} = \alpha_j + B_j \cdot T_{ij} + r_{ij}$$

Level Two: Sites

$$\alpha_j = \alpha_j$$

$$B_j = \beta + b_j$$

Why fixed site-specific intercepts?

- To account for cross-site variation in \bar{T}_j and hence the potential for bias in estimates of β and τ_B^2 due to a possible correlation between \bar{T}_j and α_j

An Alternative Expression of the Impact Estimation Model

Site-Center All Variables

- This is equivalent to specifying fixed site-specific intercepts after one accounts for the degrees of freedom lost when site-centering the dependent variable

Level One: Individuals

$$Y_{ij} - \bar{Y}_j = B_j(T_{ij} - \bar{T}_j) + r_{ij} - \bar{r}_j$$

Level Two: Sites

$$B_j = \beta + b_j$$

Specify a separate level-one residual variance for Ts and Cs

- Removes potential bias in cross-site variance estimates

How Many Level-One Residual Variances to Estimate?

A Cautionary Tale: Using Data from the *Head Start Impact Study*

- With a separate level-one residual variance for each site there appeared to be a huge amount of cross-site variation in program effects (which was highly statistically significant).
- With a single level-one residual variance for all sites and assignment groups there appeared to be much less cross-site variation in program effects (which was somewhat statistically significant).
- With a separate level-one residual variance for Ts and Cs the results were similar to those for a single variance.

Bottom Line

- Estimating too many variances reduces the sample size for each estimate and thereby increases the uncertainty about those estimates.
- This uncertainty (perhaps counter-intuitively) causes one to understate impact estimation error variance for each site (V_j) and thereby over-state true cross-site impact variation (τ_B^2).

Head Start Impact Study Example Of How Method Matters for Estimating Cross-Site Variation In Effects of Program Assignment

- **Sample size:** 119 centers, 1,056 children from the 3 year old cohort
- **Outcome:** Woodcock Johnson Letter Word Identification test score at the end of the first year after random assignment
- **Issue:** Massive difference in results from two different methods for estimating variation in effects of program assignment
 - Method #1:** Site centering the treatment indicator for a random Head Start impact model with data pooled across blocks (a single level-one residual variance)
 - Method #2:** A “split sample” model of Head Start impacts by site combined with a V-Known random-effects meta analysis (a separate level-one residual variance for each site)

**Head Start Impact Study Results
for Two Estimation Methods
(Three-year-old Cohort)**

Estimation Approach	Estimated Impact	True Impact Variation (τ)	Chi-sqr stat for τ	P-value
Single centering RE approach	6.071	35.737	125.705	0.296
Split sample + V-known approach	7.746	261.390	421.391	0.000

Key Results to Report From A Cross-Site Analysis Of Program Effects

Results to report

- Estimated grand mean program effect ($\hat{\beta}$)
- Estimated cross-site standard deviation of program effects (τ_B)
- Estimated cross-site distribution of program effects (Adjusted Empirical Bayes Estimates)
- Estimated mean program effect for each site (Empirical Bayes Estimates)
- Estimated difference in mean program effects for two categories of sites ($\hat{\beta}_2 - \hat{\beta}_1$)

Empirical Example: MDRC's Welfare-to-Work Studies¹

Research Design

- Secondary analysis of individual data from three MDRC multi-site randomized trials (GAIN, NEWWS and PI)

Study Sample

- 59 local welfare offices with an average of 1,176 randomized sample members per office (site)

Outcome Measure

- Total earnings (in dollars) during the first two years after random assignment

¹ Bloom, H S., C. J. Hill and J. A. Riccio (2003) "Linking Program Implementation and Effectiveness: Lessons from a Pooled Sample of Welfare-to-Work Experiments," *Journal of Policy Analysis and Management*, 22(4): 551 – 575.

Summary of Welfare-to-Work Parameter Estimates¹

Estimated Cross-site Grand Mean Program Effect ($\hat{\beta}$)

- Point estimate = \$875
- Estimated standard error = \$137
- P-value ≤ 0.001
- 95 percent confidence interval = \$606 to \$1,144

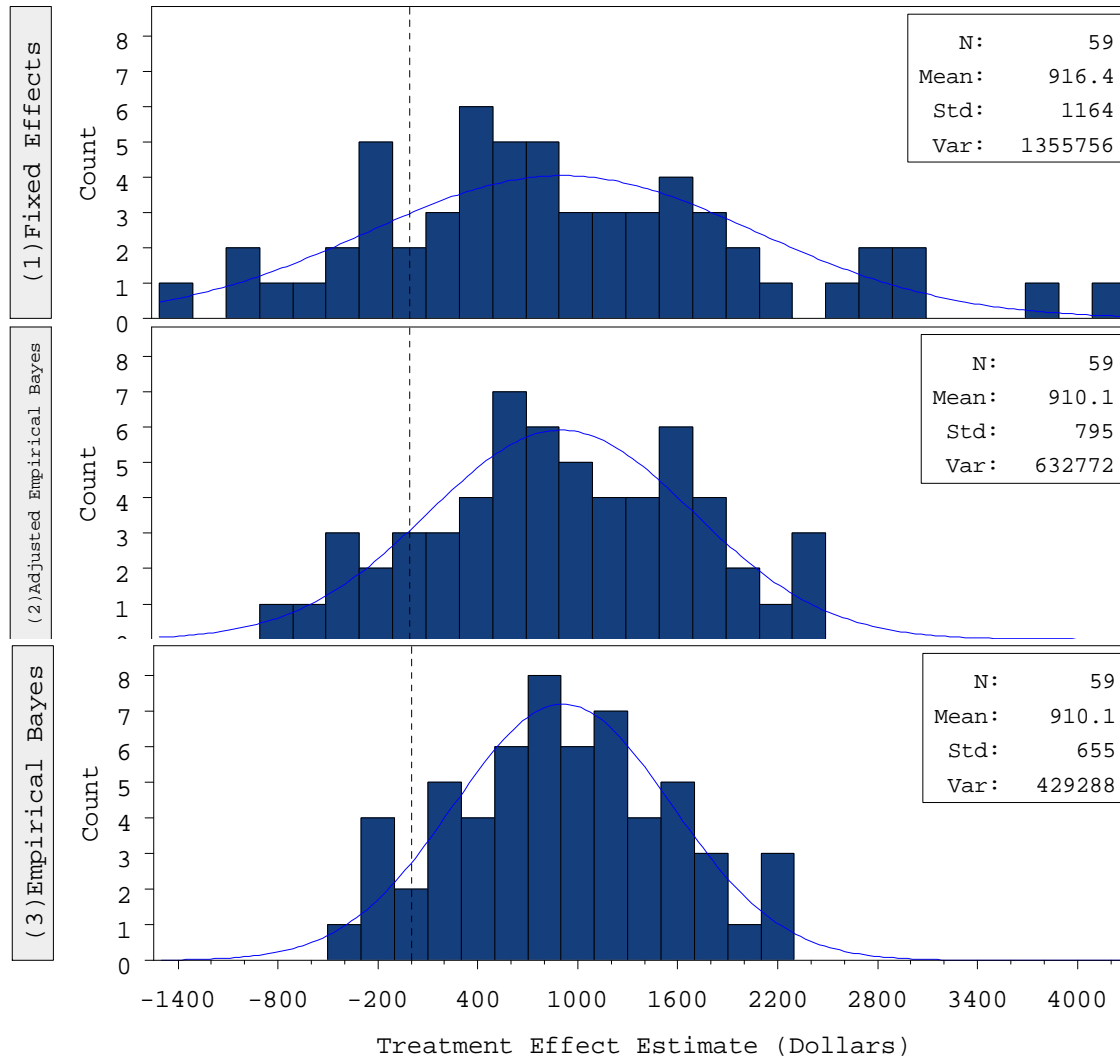
Estimated Cross-Site Standard Deviation of Program Effects (\hat{t}_B)

- Point estimate = \$742
- P-value ≤ 0.001
- Asymmetric 95 percent confidence interval = \$525 to \$1,048

NOTE: Cross-site reliability = 0.497 and $\sigma_T^2/\sigma_C^2 = 1.09$

¹ From Bloom, Raudenbush, Weiss and Porter (under review).

Cross-Site Distribution of Welfare-to-Work Program Effects on Total Two-Year Earnings



Some Important Diagnostics

Assessing the Implications of Uncertainty

- It is important to assess the implications of uncertainty for interpreting one's findings about cross-site variation
- This uncertainty is a function of the study design that produced the findings

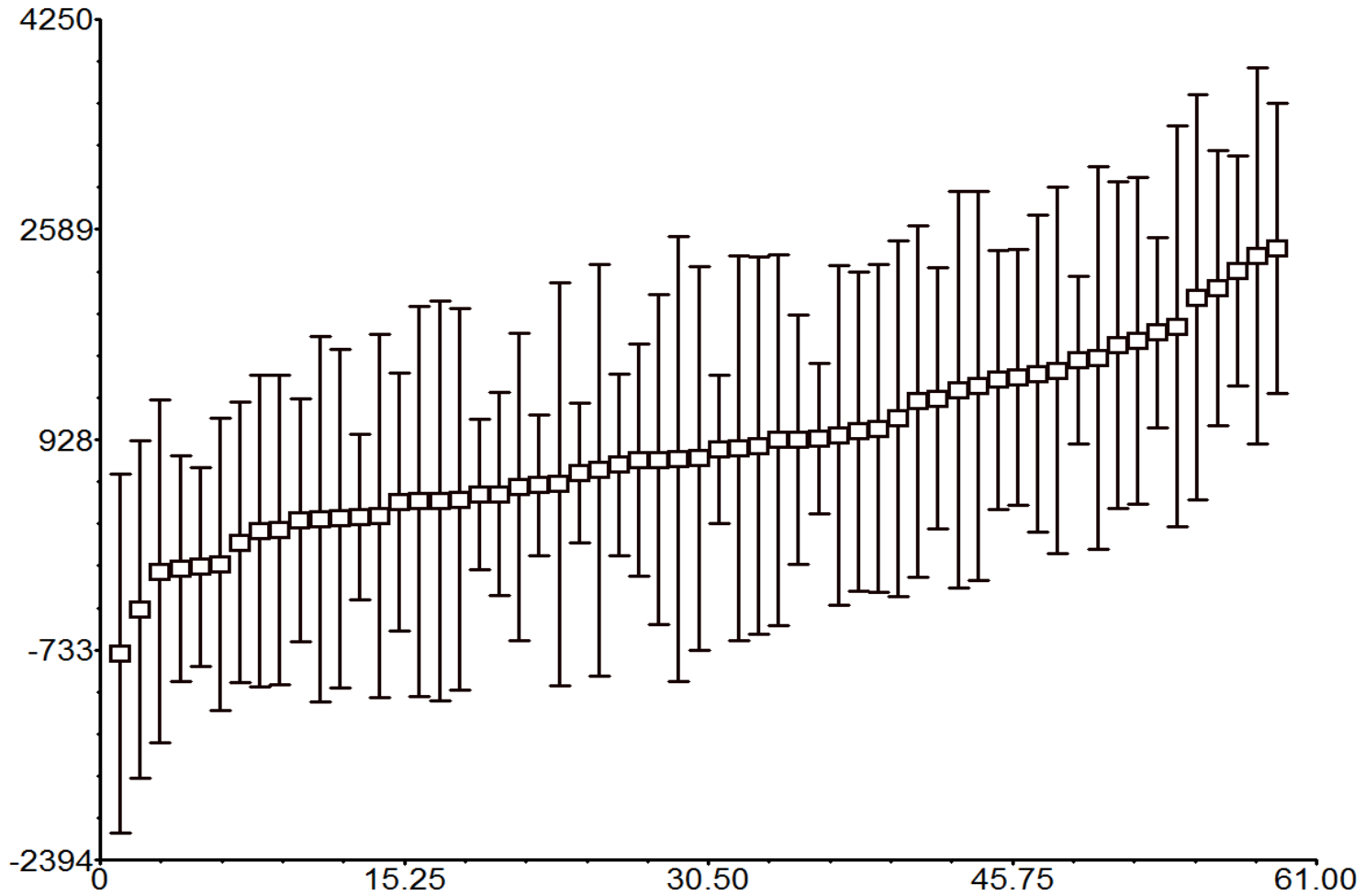
Caterpillar Plots

- graphically report confidence intervals of the OLS or Empirical Bayes estimates of the program effect for each site

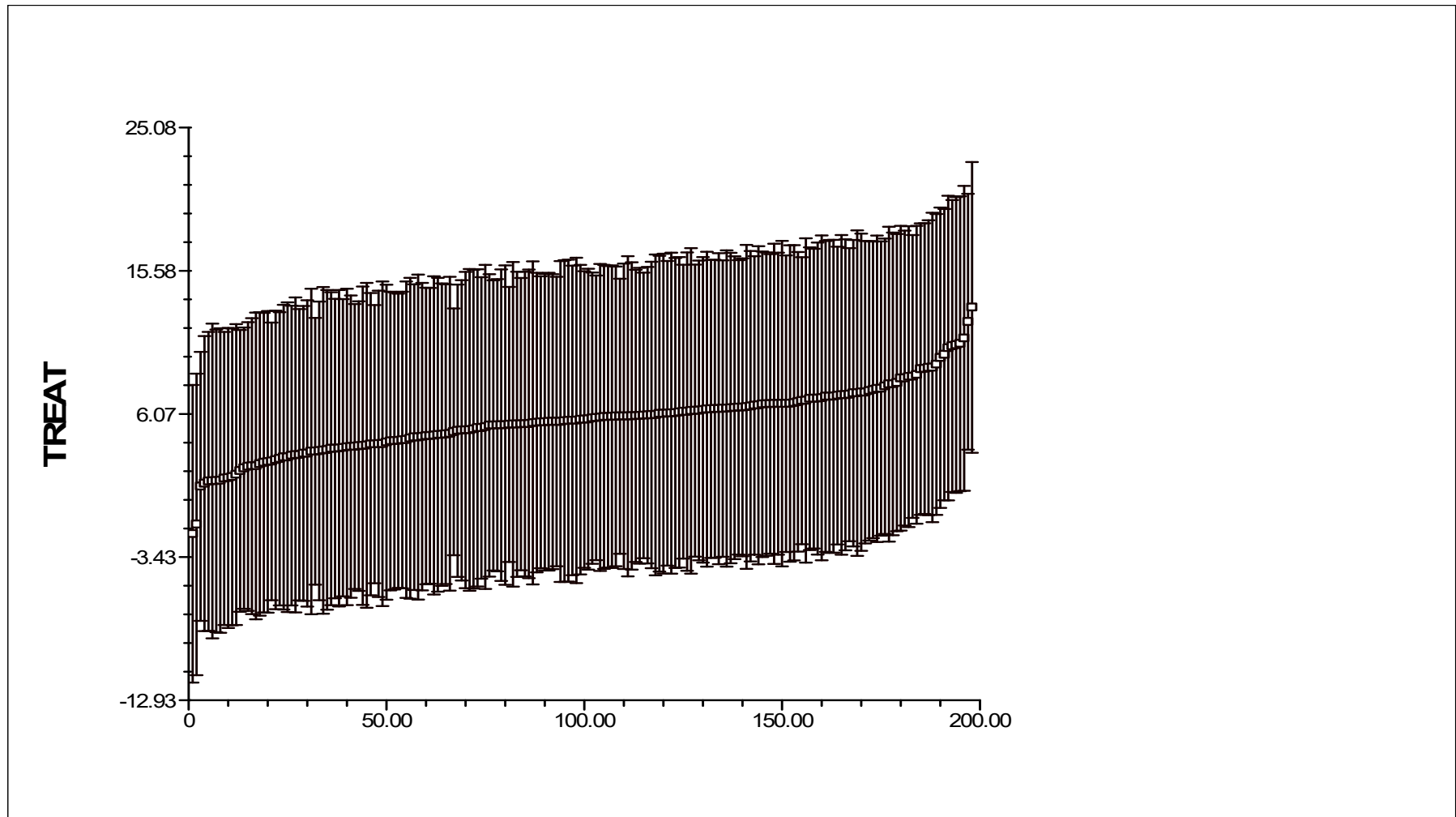
Likelihood Profile Graphs

- Superimpose a graph of the likelihood function for τ^2
- On a graph of the corresponding Empirical Bayes impact estimates for sites

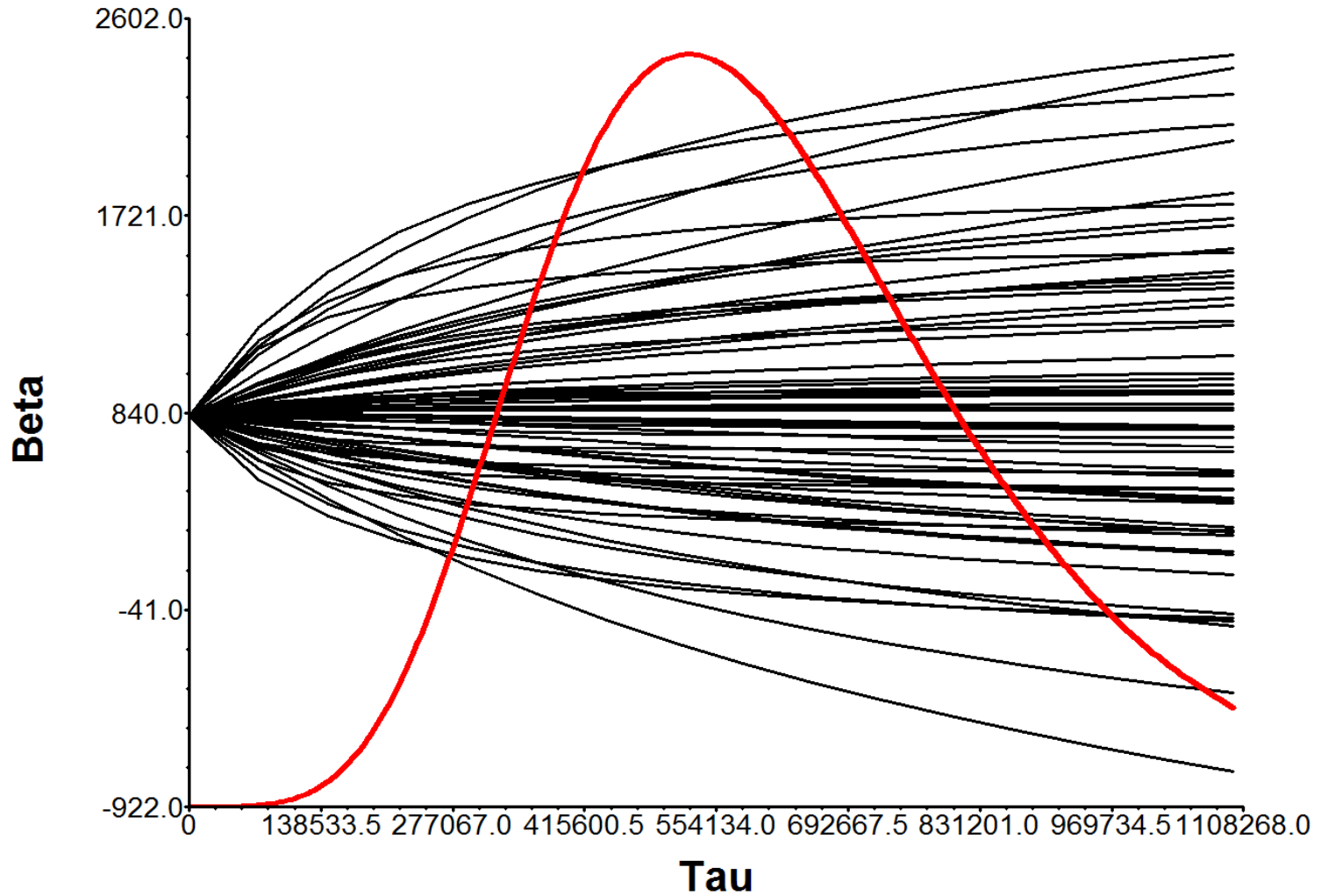
Caterpillar Plot of Empirical Bayes Estimates of Site-Specific Welfare-to-Work Program Effects



Caterpillar Plot For Empirical Bayes Estimates of Head Start Effects on Woodcock Johnson Letter Word Identification Scores



Likelihood Profile Graph for Empirical Bayes Estimates of Site-Specific Welfare-to-Work Program Effects



Profile Likelihood Graph For Empirical Bayes Estimates of Head Start Effects on Woodcock Johnson Letter Word Identification Scores

