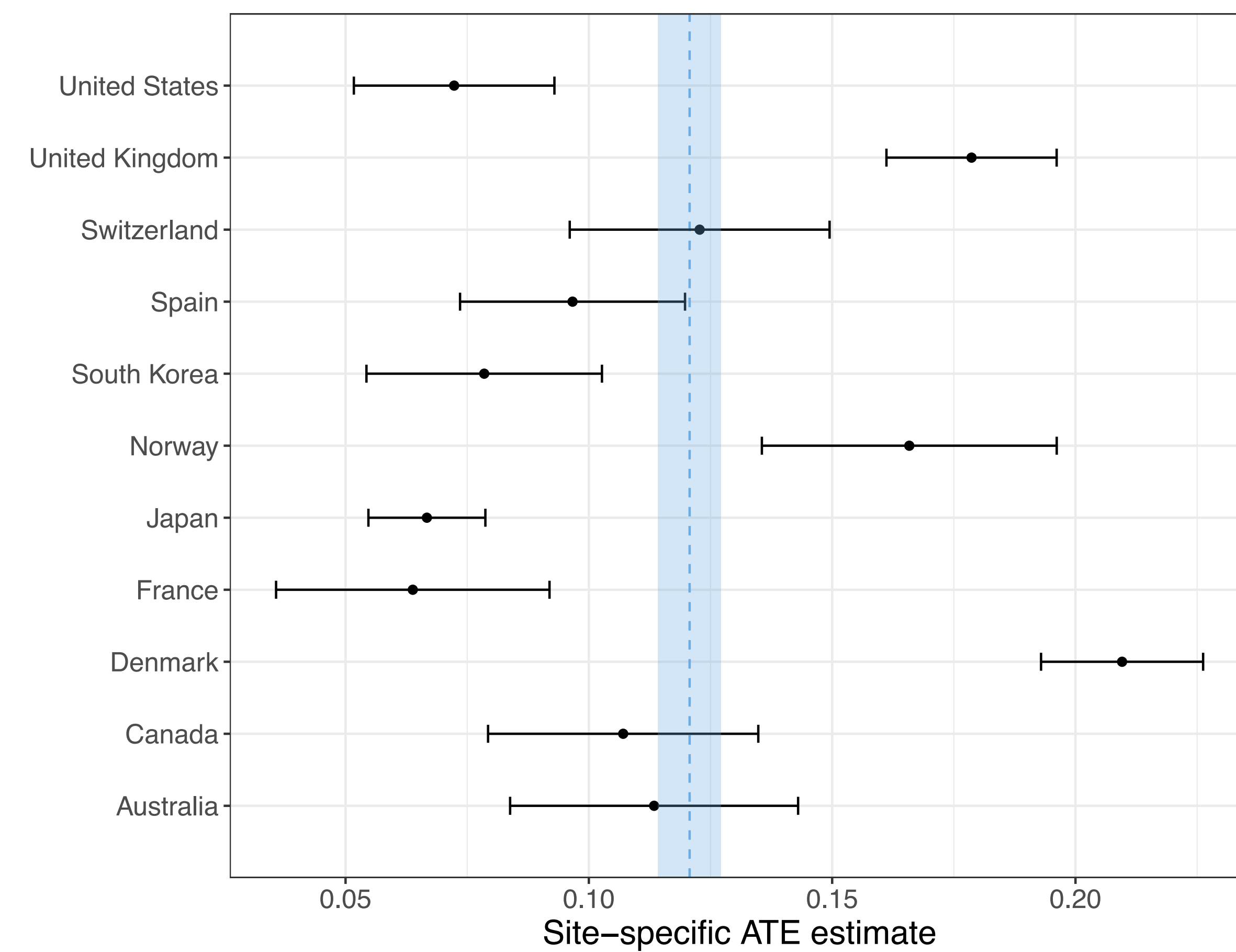


Decomposing Treatment Effect Heterogeneity in Multisite Experiments

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Problem

What explains the variation across sites in a multisite experiment?



Example from: Valentino, N. A., Soroka, S. N., Iyengar, S., Aalberg, T., Duch, R., Fraile, M., Hahn, K. S., Hansen, K. M., Harell, A., Helbling, M., Jackman, S. D., & Kobayashi, T. (2019). Economic and Cultural Drivers of Immigrant Support Worldwide. *British Journal of Political Science*, 49(4), 1201–1226.

Contribution

Existing methods:

- Meta-analysis: is there between-study variance, net of sampling variation?
- Meta-regressions: does variation in specific site-level covariates correlate with cross-site variation?
- Reweighting: Lu et al. (2023) - how much variation would remain if all sites had the same distribution of observed unit-level covariates? Lu, B., Ben-Michael, E., Feller, A., & Miratrix, L. (2023). Is It Who You Are or Where You Are? Accounting for Compositional Differences in Cross-Site Treatment Effect Variation. *Journal of Educational and Behavioral Statistics*, 48(4), 420–453.

New question: based on observed covariates, do **site-level** or **unit-level** features explain more of the heterogeneity?

Quantity of interest: how much variation would remain if covariates at one level were held identical in expectation across sites?

Helps answer:

1. Are population or context differences driving the heterogeneity?
2. How do the modeled unit- and site-explained variations compare to the total systematic heterogeneity?

Formal Setup

- Outcome Y , treatment T , sites $\{1, \dots, K\}$
- Observed unit-level covariates X , observed site-level covariates M
- Unobserved unit-level covariates U_x , site-level covariates U_m
- CATE on observed covariates:

$$\tau(X, M) = \mathbb{E}_{U_x, U_m}[\tau(X, M, U_x, U_m) \mid X, M]$$

Estimands:

$$\tau_{site}^2 = \text{Var}(\mathbb{E}[\tau(X, M) \mid X = \tilde{x}, M])$$

$$\tau_{unit}^2 = \text{Var}(\mathbb{E}[\tau(X, M) \mid X, M = \tilde{m}])$$

Key assumption: conditional cross-level independence

$$U_x \perp M \mid X, M \text{ and } U_m \perp X \mid X, M$$

Under this assumption, while τ_{site}^2 and τ_{unit}^2 might also capture variation explained by unobserved covariates, they do not inadvertently capture variation explained by the other level.

Estimation

Algorithm - estimation of τ_{site}^2 and τ_{unit}^2

Input: pooled.data (experimental data pooled across sites)

Output: $\tau_{site}^2, \tau_{unit}^2$

1. $M, X, Y, T \leftarrow$ pooled.data[M], pooled.data[X], pooled.data[Y], pooled.data[T];
2. $\tau(\cdot) \leftarrow$ outcome model estimated using M, X, Y, T ;

Shuffle covariates at each level

3. data.site \leftarrow sample(X), M, Y ;
4. data.unit $\leftarrow X$, sample(M), Y ;

Predict potential outcomes

5. data.site[\tilde{Y}_1, \tilde{Y}_0] $\leftarrow \tau(\text{data.site})$;
6. data.unit[\tilde{Y}_1, \tilde{Y}_0] $\leftarrow \tau(\text{data.unit})$;

Estimate site average treatment effects

7. **for** site j **do**
- 8 | $\widehat{ATE}_{site,j} \leftarrow \text{mean}(\text{data.site}[\tilde{Y}_1] - \text{data.site}[\tilde{Y}_0])$;
- 9 | $\widehat{ATE}_{unit,j} \leftarrow \text{mean}(\text{data.unit}[\tilde{Y}_1] - \text{data.unit}[\tilde{Y}_0])$;
10. **end**

Estimate cross-site variances

11. $\hat{\tau}_{site}^2 \leftarrow$ estimated between-site variance of \widehat{ATE}_{site} ;
12. $\hat{\tau}_{unit}^2 \leftarrow$ estimated between-site variance of \widehat{ATE}_{unit} ;

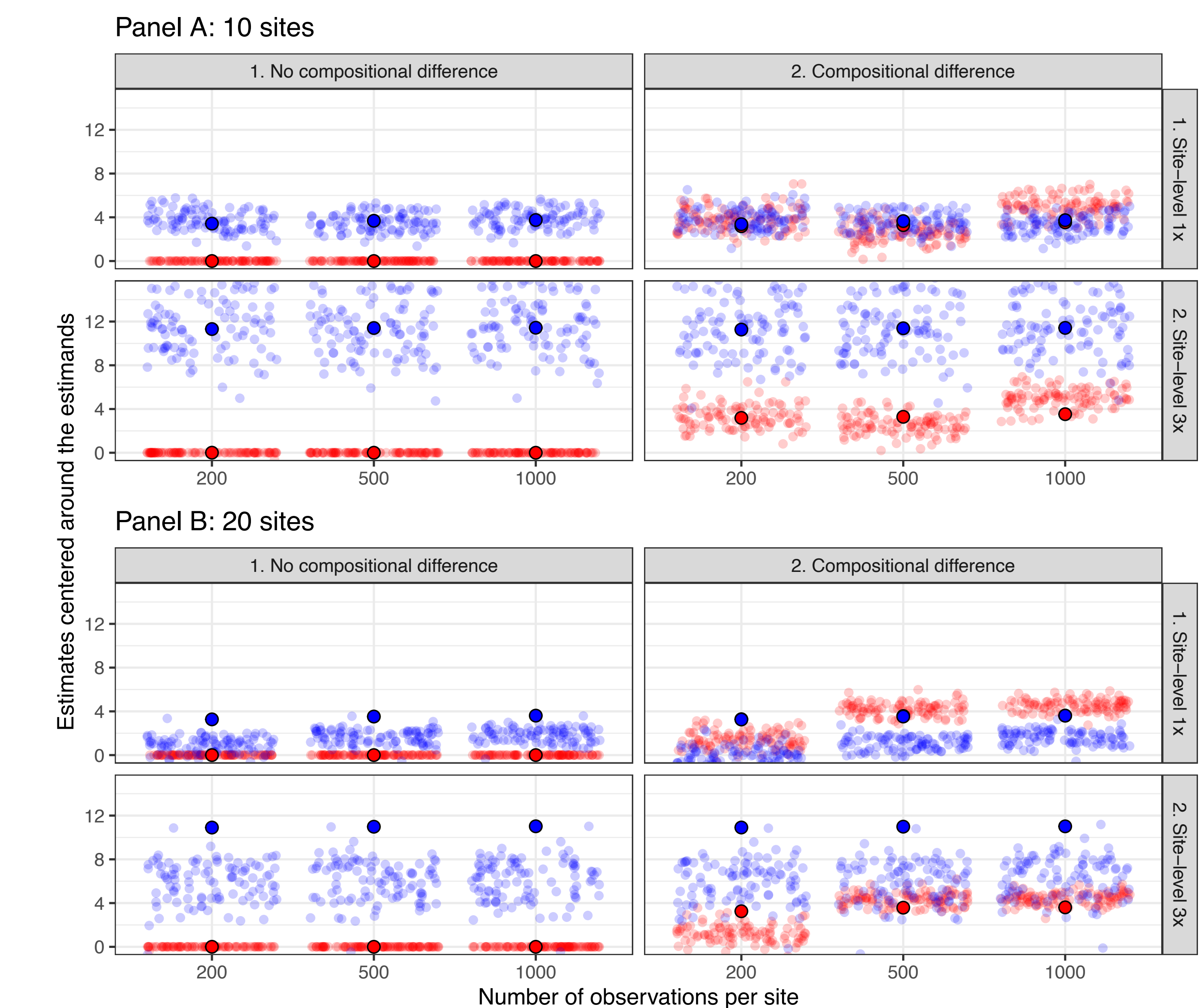
Permutation test - difference between $\hat{\tau}_{site}^2$ and $\hat{\tau}_{unit}^2$:

The site-level datasets \widehat{ATE}_{site} and \widehat{ATE}_{unit} are combined. The “site” and “unit” labels are randomly permuted. Each time, $\hat{\tau}_{site}^2 - \hat{\tau}_{unit}^2$ is computed, providing a distribution under the null of no difference.

Simulation

With different coefficients, compositional differences, number of observations, and number of sites. Outcome model selected: BART

1. How do the estimands (τ_{site}^2 and τ_{unit}^2) behave?
2. How well does the estimator recover them?



With 20 sites, $\hat{\tau}_{site}^2$ is consistently underestimated.

Tentative reason: data suggests that with more sites, BART gives less importance to the unit-level covariates and might not be adapted to the nested structure of the data.

Tentative solution: use a multilevel outcome model.

Application

STAR experiment (Tennessee, 1985-1989): Does reducing class-size lead to better educational outcomes?

- Reports focus on how site-level (e.g. inner-city versus rural) moderate the treatment effect.
- The decomposition suggests that $\hat{\tau}_{unit}^2$ is actually three times as large as $\hat{\tau}_{site}^2$ and significantly so.
- **Limitation:** can composition really be distinguished from context?

Word, E., Johnston, J., Pate Bain, H., DeWayne Fulton, B., Boyd Zaharias, J., Achilles, C., Nannette Lintz, M., Folger, J., & Breda, C. (1990). *The State Of Tennessee's Student/Teacher Achievement Ratio (Star) Project, Technical Report*. Tennessee State Department of Education.