

Title in Progress

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- ▶ Propensity score matching (PSM) is a tool that can help us mitigate the effects of confounders. . .
- ▶ . . . but there is no consensus on the best way to estimate standard errors when using the PSM algorithm.
- ▶ How can we assess which procedures reliably estimate standard errors?

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A simulation study!

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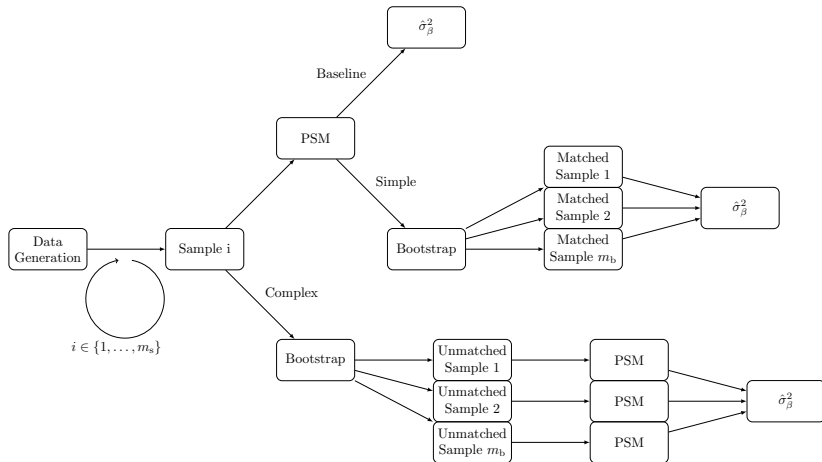
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- ▶ The PSM algorithm intakes an unmatched dataset and outputs a matched one.
- ▶ When do we execute the bootstrap - before the match or after it?
- ▶ Let's try both!

Roadmap of the Simulation Study



Data Generation - Continuous Outcome

For each individual $i \in \{1, \dots, n\}$, we consider covariates $L_{1i}, L_{2i}, L_{3i} \sim N(0, 1)$. Treatments are distributed according to law $A_i \sim B(\pi_i)$, where π_i - the true propensity to be treated - is subject to the data-generating process

$$\log \left(\frac{\pi_i}{1 - \pi_i} \right) = \alpha_0 + \alpha_1 L_{1i} + \alpha_2 L_{2i}.$$

Given this, we further define the data-generating process of our continuous outcome via

$$Y_i = \beta_1 A_i + \beta_2 L_{2i} + \beta_3 L_{3i} + \varepsilon_i,$$

where ε_i denotes random error. Because L_{2i} effects both A_i and Y_i , it acts as a confounder in estimating the treatment effect.

Data Generation - Binary Outcome

For each individual $i \in \{1, \dots, n\}$, we consider covariates $L_{1i}, L_{2i}, L_{3i} \sim N(0, 1)$. Treatments are distributed according to law $A_i \sim B(\pi_i)$, where π_i - the true propensity to be treated - is subject to the data-generating process

$$\log \left(\frac{\pi_i}{1 - \pi_i} \right) = \alpha_0 + \alpha_1 L_{1i} + \alpha_2 L_{2i}.$$

Given this, we further define the data-generating process of our binary outcome via $Y_i \sim B(\tau_i)$ where

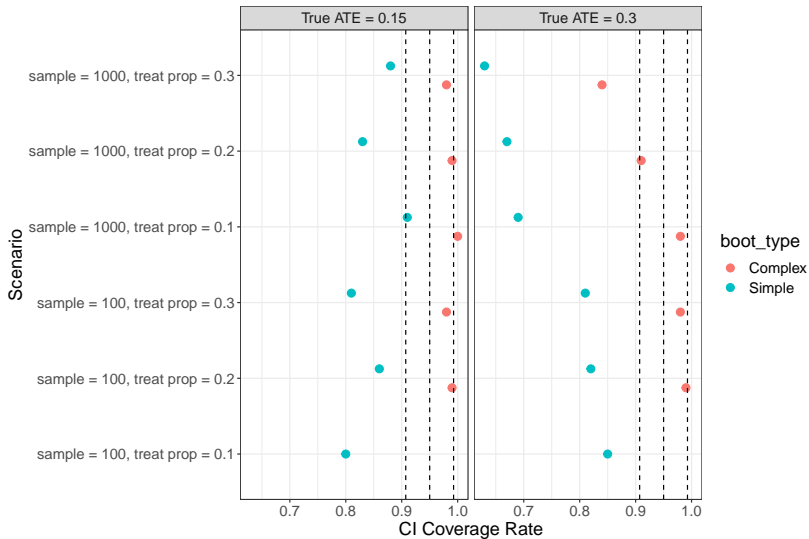
$$\log \left(\frac{\tau_i}{1 - \tau_i} \right) = \beta_1 A_i + \beta_2 L_{2i} + \beta_3 L_{3i}.$$

Observe that we have omitted a random error term, as realizations of Y_i are innately subject to noise.

Measures

Results

Binary Coverage Rates by Parameters of Interest

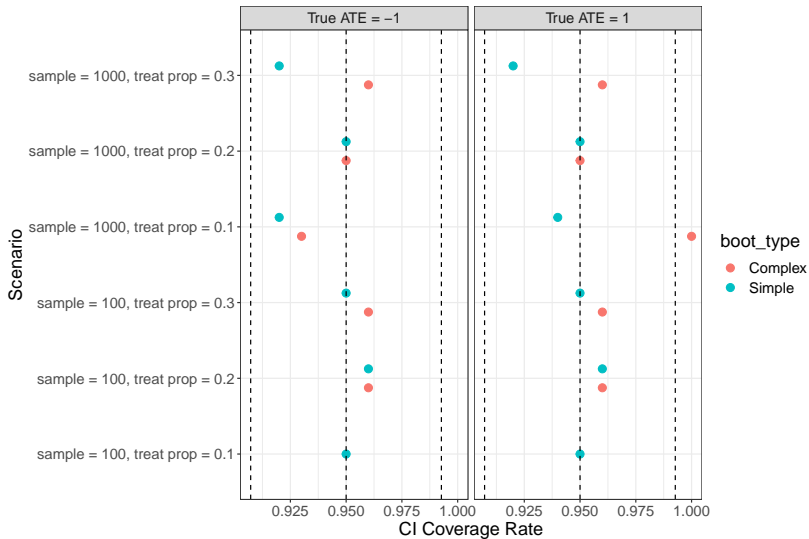


Results

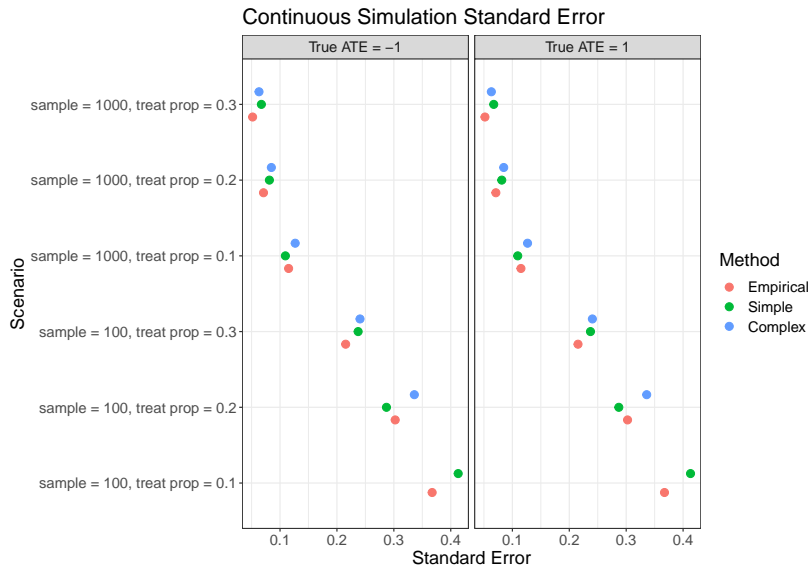


Results

Continuous Coverage Rates by Parameters of Interest



Results



Limitations

- ▶ Sample size / treatment (or exposure) prevalence

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- ▶ Small number of initial samples, limited in detecting differences in coverage rate

Future Work

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- ▶ Increased sample size, changes in bootstrap performance?
- ▶ Changes in treatment propensity model
- ▶ Non-normal distributions of covariates