

Robust Nonparametric Inference for Stochastic Interventions under Multi-Stage Sampling Nima S. Hejazi, Mark J. van der Laan, and David C. Benkeser

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OVERVIEW & MOTIVATIONS

- We consider the problem of efficiently estimating the effect of a stochastic shift interventions in studies with two-phase sampling of the treatment.
- We present an estimator of the average counterfactual outcome under a stochastic shift intervention with
 - consistency and efficiency guarantees,
 - a multiple double robustness property.
- The proposed estimator is asymptotically normal with estimable variance, thereby allowing for the construction of confidence intervals and hypothesis tests.
- The *txshift* R package [2] implements these estimators and leverages state-of-the-art machine learning in the procedure.

DATA: HIV VACCINE TRIALS

- Our approach is motivated by application to investigations of the effects of immune responses on HIV vaccine efficacy.
- Question: How does risk of HIV infection differ under shifts of an immune response in the vaccine arm of an efficacy trial?
- We simulate a data structure based on the HVTN 505 HIV-1 efficacy trial, as in [3]:
 - 2504 participants, with all observed cases matched to controls.
 - Background (W): sex, age, BMI, etc.
 - Intervention (*A*): immunobiomarkers (i.e., T-Cell profiles from ICS assays on preserved HIV-1-stimulated PBMCs).
 - Outcome (Y): HIV-1 infection status.
- Takeaway: Variable importance measure for ranking immune responses by utility as immunogenicity study endpoints in all future HIV-1 vaccine efficacy trials.

METHODOLOGY I: THE EFFECT OF A STOCHASTIC INTERVENTION

- Consider $X = (W, A, Y) \sim P_0^X \in \mathcal{M}$, where W is a set of baseline covariates, A a treatment, and Y an outcome of interest, with no assumptions placed on the statistical model \mathcal{M} .
- Consider a shift of the treatment (i.e., d(A, W)) so that $A = A + \delta$ for a user-specified shift δ .
- To protect against violations of the assumption of positivity, the shifting mechanism may be made a function of the observed data, where u(w) is the maximum shift with support in the data:

$$d(a, w) = \begin{cases} a + \delta, & a + \delta < u(w) \\ a, & \text{otherwise} \end{cases}$$

• The causal parameter is identified by the observed data parameter [4]:

$$\Psi(P_0^X) = \mathbb{E}_{P_0} \overline{Q}(d(A, W), W), \tag{1}$$

where $\overline{Q}(d(A, W), W)$ is the conditional mean of the outcome given A = d(A, W) and W.

• The efficient influence function (EIF) of Ψ relative to a nonparametric model is

$$D^{F}(P_{0}^{X})(x) = H(a, w)(y - \overline{Q}(a, w)) + \overline{Q}(d(a, w), w) - \Psi(P_{0}^{X})(x),$$
(2)

where the auxiliary term, H(a, w), may be expressed as

$$H(a, w) = \mathbb{I}(a < u(w)) \frac{g(a - \delta \mid w)}{g(a \mid w)} + \mathbb{I}(a \ge u(w) - \delta).$$
(3)

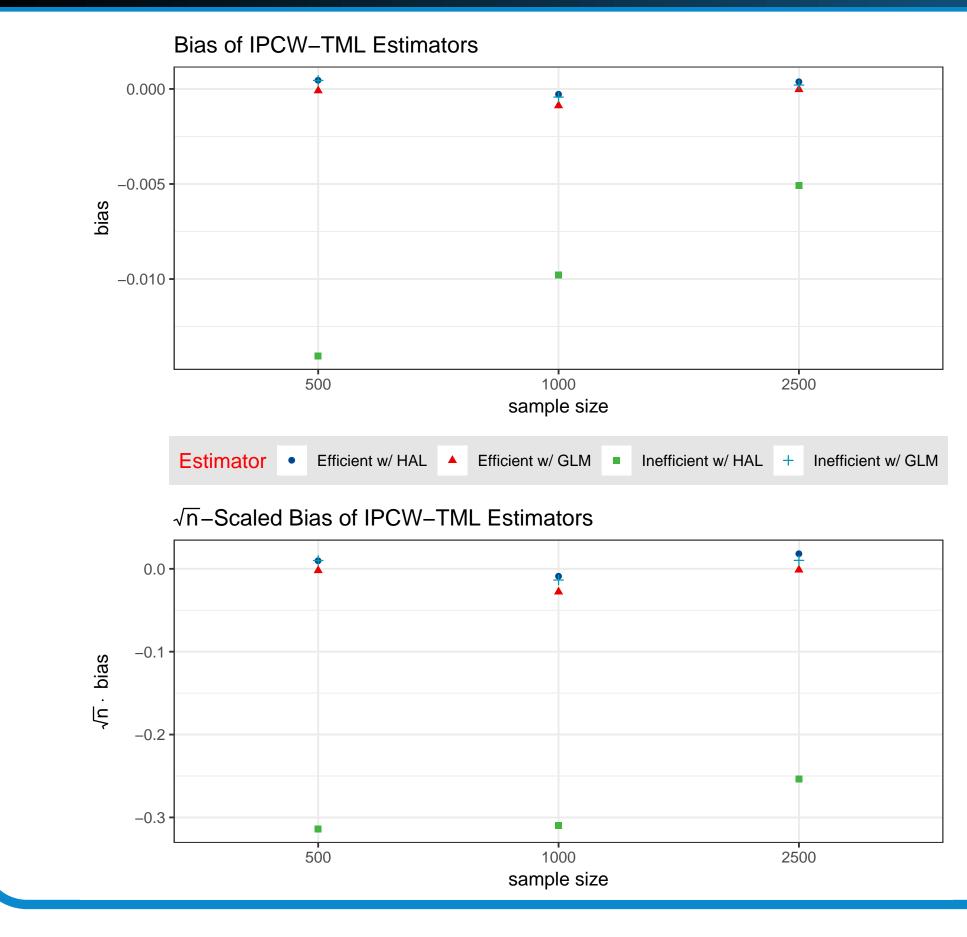
METHODOLOGY II: CORRECTIONS FOR TWO-PHASE SAMPLING

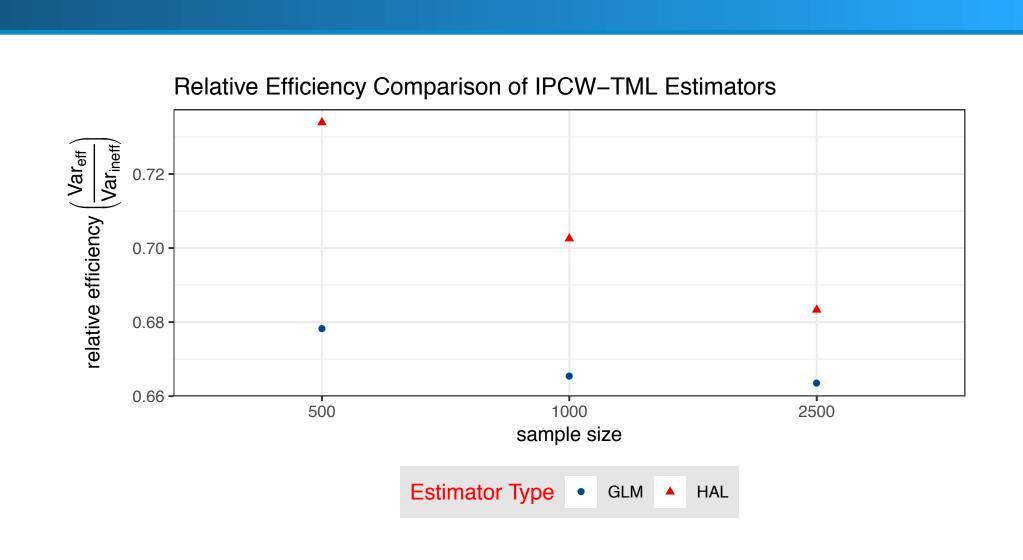
- In the HVTN 505 HIV-1 trial, all infected participants and the matched subset of controls have A (immunobiomarkers) measured, thus making the observed data structure $O = (W, \Delta A, Y) \sim P_0$.
 - $\Delta \in \{0,1\}$ is the missingness mechanism introduced by sampling, under which the observed immune response (ΔA) is arbitrarily set to 0 when unobserved.
 - We assume that, given V := (W, Y), Δ is Bernoulli distributed with probability $\Pi_0(V)$.
- The IPCW-TMLE [5] estimates the target parameter by adding *inverse weights* to the loss function.
- Improvements in the efficiency are attainable using a TMLE based on the EIF:

$$D(P_0)(o) = \frac{\Delta}{\Pi(v)} D^F(P_0^X)(x) - \left(1 - \frac{\Delta}{\Pi(v)}\right) \mathbb{E}(D^F(P_0^X)(x) \mid \Delta = 1, V = v)$$

- This augmented estimator exhibits several desirable properties
 - efficiency, achieving the CR bound for the class of regular asymptotically linear estimators;
 - multiple robustness, consistency of the parameter estimate when one of (g,Q) and one of $(\Pi, \mathbb{E}_0(D^F(P_0^X)(X) \mid \Delta = 1, V))$ is consistently estimated;
 - valid statistical inference even when Π is estimated nonparametrically.

RESULTS & DISCUSSION





- All estimators approx. unbiased in large samples; however, inefficient TMLE with HAL has bias not converging at $n^{-\frac{1}{2}}$.
- Fitting II with HAL or GLM, efficient TMLE has lower variance than the inefficient.

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