

Robust inference on the causal effects of stochastic interventions in two-phased vaccine efficacy trials Nima Hejazi, David Benkeser, and Mark van der Laan

EMORY

Graduate Group in Biostatistics & Department of Statistics, University of California, Berkeley Department of Biostatistics and Bioinformatics, Emory University

OVERVIEW & MOTIVATIONS

- We consider estimating the effect of a stochastic intervention in vaccine trials with two-phase sampling of the treatment.
- We present efficient estimators of the mean counterfactual outcome under a stochastic intervention with
 - consistency and efficiency guarantees,
 - a *multiple* double robustness property.
- The proposed estimators are asymptotically normal with a variance estimator based on a well-studied efficient influence function.
- Our txshift R package [3] implements these estimators and leverages state-of-theart machine learning in the procedure.
- Our haldensify R package [2] estimates the propensity score, a conditional density.

HIV VACCINE EFFICACY TRIALS

- Question: How does HIV infection risk vary with the shifting of an immunogenic response profile in a trial's vaccine arm.
- Analysis of data from the HIV Vaccine Trials Network 505 efficacy trial, as in [4]:
 - RCT with n=2504; only n=189 from vaccine arm in second-stage sample.
 - Background covariates (W): sex, age, BMI, behavioral HIV risk score.
 - Treatment (*A*): immunogenic response profiles, post-vaccination.
 - Outcome (Y): HIV-1 infection status at week 28 of trial.
 - All individuals with HIV-1 infections by week 28 in second-stage sample.
- Goal: Develop a ranking of immunogenic responses as study endpoints for future HIV-1 vaccine efficacy trials.

THE EFFECT OF A STOCHASTIC TREATMENT REGIME

- 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 $d(A, W) = A + \delta$ for a given shift δ . To protect against violations of the positivity assumption, make the shifting mechanism a function of the observed data, where u(w) is the maximum shift supported in the observed data:

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

• The causal parameter has been shown to be identified by a functional of the observed data [1]:

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

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

ullet The efficient influence function (EIF) of Ψ relative to the nonparametric model ${\cal M}$ is

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

where the auxiliary covariate 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)

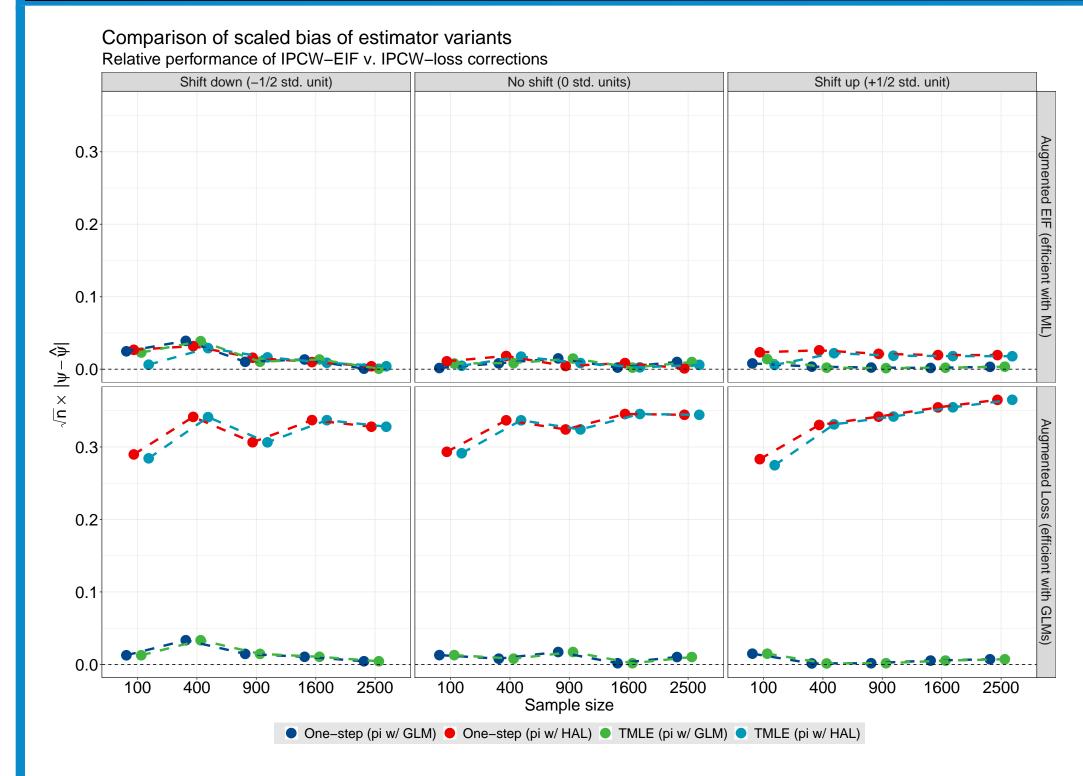
EFFICIENT CORRECTIONS FOR TWO-PHASE SAMPLING

- In the HVTN 505 HIV-1 trial, immunogenic responses A are sequenced for infected individuals and a matched subset of controls, making $O=(W,\Delta,\Delta A,Y)$ the observed data structure.
 - $\Delta \in \{0,1\}$ is the sampling mechanism introduced by two-phase sampling, under which the observed immunogenic response (ΔA) is arbitrarily set to 0 when unobserved.
 - Given V := (W, Y), we assume that $\Delta \sim \text{Bern}(\pi_0(V))$.
- The IPCW-TMLE [5] estimates the target parameter by adding *inverse weights* to the loss function.
- Efficiency improvements are attainable by constructing estimators based on the augmented EIF:

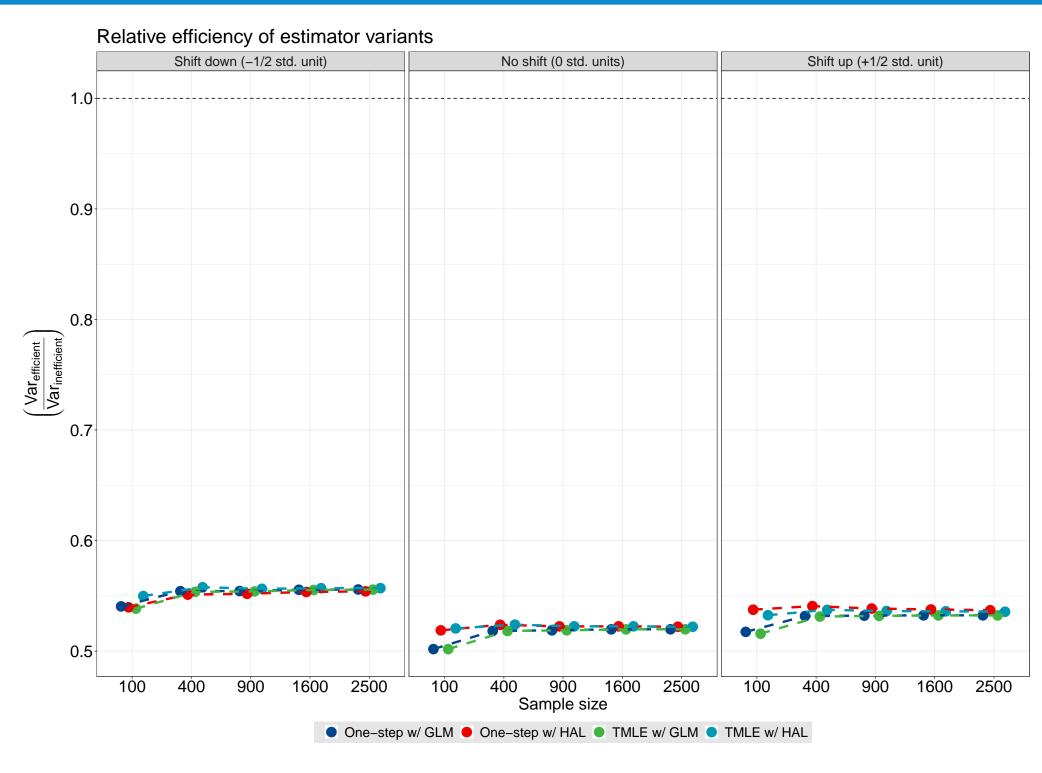
$$D(P_0)(o) = \frac{\Delta}{\pi(v)} D^F(P_0^X)(x) - \left(1 - \frac{\Delta}{\pi(v)}\right) \mathbb{E}(D^F(P_0^X)(x) \mid \Delta = 1, V = v)$$

- This augmentation allows the construction of estimators that
 - achieve the efficiency bound for the class of regular asymptotically linear estimators;
 - are multiply robust, with consistency of parameter estimates when one of $\{g,Q\}$ and one of $\{\pi_0(V), \mathbb{E}_0(D^F(P_0^X)(X) \mid \Delta=1,V)\}$ are consistently estimated;
 - allow valid statistical inference to be derived even when π_0 is estimated nonparametrically.

NUMERICAL STUDY & RESULTS



(a) Most estimator variants unbiased in large samples; inefficient variants with HAL fail to converge.



(b) Fitting π with HAL or GLM, efficient estimators show relative efficiency similar to simpler variants.

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CONTACT INFORMATION

- N. Hejazi, Graduate Group in Biostatistics, UC Berkeley, nhejazi@berkeley.edu
- D. Benkeser: Asst. Prof. of Biostatistics, Emory University, benkeser@emory.edu
- M. van der Laan, Prof. of Biostatistics, UC Berkeley, laan@berkeley.edu