Discussion Assignment #1

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Question 8: Give a detailed implementation of longitudinal IPTW to estimate parameters of an MSM without effect modifiers (Section 7-8).

Implementing the Horvitz-Thompson Estimator

We are interested in the expected Y if everyone got treatment regime $\bar{A}(t)=\bar{a}$, for t=0,1.

$$\Psi^F(P_{U,X}) = E_{U,X}[Y_{\bar{A}(t)=\bar{a}}]$$

The Horvitz-Thompson IPTW estimator for $\Psi^F(P_{U,X})$ is:

$$\hat{\Psi}(P_n) = \frac{1}{n} \sum_{i=1}^n \frac{\mathbb{I}[\bar{A}_i(t) = \bar{a}]}{g_n(A_i(0)|L_i(0)) \times g_n(A_i(1)|A_i(0), L_i(0), L_i(1))} Y_i$$

Step 1: Calculate appropriate stabilized weights using the modified Horvitz-Thomas estimator

Weights =
$$\frac{1}{g_n(A_i(0)|L_i(0)) \times g_n(A_i(1)|A_i(0),L_i(0),L_i(1))}$$

Part A:

Estimate the probability of receiving treatment using correctly specified parametric regression models (logistic regression)

$$g_0(A(0) = a(0)|L(0)) = expit[\beta_0 + \beta_1 L(0)]$$

$$g_0(A(1) = a(1)|\bar{L}(1), A(0)) = expit[\beta_0 + \beta_1 L(0) + \beta_2 L(1)]$$

In this example we are estimating the probability of being treated with AZT at each time point, given the covariate pattern and the prior history of AZT treatment.

Step 1: Calculate appropriate stabilized weights using the modified Horvitz-Thomas estimator

Part B:

Predict each subject's probability of the exposure at each time t, given his or her observed exposure and covariate history.

$$g_n(A_i(t)=a_i(t)|\bar{A}_i(t-1),\bar{L}_i(t))$$

In this example:

- for time points where AZT treatment is NOT occurring it is the predicted probability of NOT being treated, given the observed past.
- for time points where AZT treatment IS occurring it is the predicted probability of being treated, given the observed past.

Step 1: Calculate appropriate stabilized weights using the modified Horvitz-Thomas estimator

Part C:

Predict each subject's probability of the entire exposure history, which is the product of the time point specific probabilities.

$$\prod_{t=1}^k (A_i(t)|\bar{A}_i(t-1),\bar{L}_i(t))$$

In this example we are estimating the probability of their entire AZT treatment hitory pattern.

The weights, as given earlier, are thus the inverse of these products.

Step 2: Take the weighted average of observed outcomes across the population

The Horvitz-Thompson IPTW estimator for $\Psi^F(P_{U,X})$ is:

$$\hat{\Psi}(P_n) = \frac{1}{n} \sum_{i=1}^n \frac{\mathbb{I}[\bar{A}_i(t) = \bar{a}]}{g_n(A_i(0)|L_i(0)) \times g_n(A_i(1)|A_i(0), L_i(0), L_i(1))} Y_i$$

The Modified or Stabilized Horvitz-Thompson IPTW estimator for $Psi^F(P_{U,X})$ is:

$$\hat{\Psi}(P_n) = \frac{\frac{1}{n} \sum_{i=1}^{n} \frac{\mathbb{I}[\bar{A}_i(t) = \bar{a}]}{g_n(A_i(0)|L_i(0)) \times g_n(A_i(1)|A_i(0),L_i(0),L_i(1))} Y_i}{\frac{1}{n} \sum_{i=1}^{n} \frac{\mathbb{I}[\bar{A}_i(t) = \bar{a}]}{g_n(A_i(0)|L_i(0)) \times g_n(A_i(1)|A_i(0),L_i(0),L_i(1))}}$$

Question 9: How would you modify the above procedure when the target causal parameter is a MSM with effect modification (Section 9)?

Including Baseline Covariates in the Model

If effect modification by baseline covariates V (a subset of L(t)) is of interest to the target causal parameter, inclusion of those baseline characteristics in the MSM allows for their incorporation into the counterfactual pseudopopulations.

Therefore, we want to condition on baseline covariates \boldsymbol{V} in our model, where the model is now:

$$E[Y_{\bar{a}}|V] = m(\bar{a}, V|\beta) = \beta_0 + \beta_1 \sum_{t=0}^{1} a(t) + \beta_2 V + \beta_3 \sum_{t=0}^{1} a(t) \times V$$

The choice of numerator in the MSM changes the target parameter being measured, and the stabilized weights can be improved:

$$\hat{sw}_i = rac{g_n(ar{A}_i(1)|V_i)}{\prod_{t=0}^1 g_n(A_i(t)|ar{A}(t-1),ar{L}_i(t))}$$