Quant II

Lab 12: Machine Learning and Generalization

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- Estimating heterogeneous treatment effects and generate the optimal assignment

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- What if it is too expensive? Active learning (Miller et al., 2019)

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- Some relationships in causal inference can be non-causal.
- We just need to fit/predict it with a high accuracy.
 - Example I: Propensity score
 - Example II: First stage of IV
 - Example III: Response surface (what covariates to control for)
- These are "nuisance parameters" that have no causal interpretation.

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- One model for the outcome, and the other for the treatment.
- But why?

• Let's consider the following DGP:

$$Y_i = \theta D_i + g_0(\mathbf{X}_i) + U_i$$

$$D_i = m_0(\mathbf{X}_i) + V_i$$

• We have $D_i \perp \{Y_i(1), Y_i(0)\} | \mathbf{X}_i$.

- The classic model-based approach will find an estimate \hat{g} for g_0 .
- Then,

$$\hat{\theta} = \frac{\sum_{i=1}^{N} D_i(Y_i - \hat{g}(\mathbf{X}_i))}{N_1} - \frac{\sum_{i=1}^{N} (1 - D_i)(Y_i - \hat{g}(\mathbf{X}_i))}{N_0}$$

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• This is "single selection."

• Nevertheless, we can show that:

$$\begin{split} & \sqrt{N}(\hat{\theta} - \theta) \\ = & \sqrt{N} \left[\frac{\sum_{i=1}^{N} D_i U_i}{N_1} - \frac{\sum_{i=1}^{N} (1 - D_i) U_i}{N_0} \right] \\ + & \sqrt{N} \left[\frac{\sum_{i=1}^{N} D_i (g_0(\mathbf{X}_i) - \hat{g}(\mathbf{X}_i))}{N_1} - \frac{\sum_{i=1}^{N} (1 - D_i) (g_0(\mathbf{X}_i) - \hat{g}(\mathbf{X}_i))}{N_0} \right] \end{split}$$

- The first part is just the Hajek estimator, which converges to N(0, I).
- But the second part may diverge to infinity as the convergence of \hat{g} to g_0 is often slow.

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- Denote $E[Y_i|\mathbf{X}_i] = m_0(\mathbf{X}_i)\theta + g_0(\mathbf{X}_i)$ as $I_0(\mathbf{X}_i)$.
- We use machine learning to estimate $m_0(\mathbf{X}_i)$ and $l_0(\mathbf{X}_i)$.
- Then, we take the residual: $\hat{V}_i = D_i \hat{m}(\mathbf{X}_i)$ and $\hat{W}_i = Y_i \hat{l}(\mathbf{X}_i)$.
- Finally, $\hat{\theta}$ is estimated by regressing \hat{W}_i on \hat{V}_i .

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- Finally, $\hat{\theta}$ is estimated by regressing \hat{W}_i on \hat{V}_i .
- Intuitively, the second part of the bias is now decided by $(\hat{m}(\mathbf{X}_i) m_0(\mathbf{X}_i))(\hat{l}(\mathbf{X}_i) l_0(\mathbf{X}_i))$ plus $V_i(\hat{g}(\mathbf{X}_i) g_0(\mathbf{X}_i))$.
- Even when each estimator converges to the true value slowly, their product may have a satisfying convergence rate.

- This is called "Robinson's Transformation" (Robinson, 1988).
- The transformation allows us to achieve "Neyman orthogonality," meaning the bias from estimating nuisance parameters have negligible influence on the estimation of causal parameters.

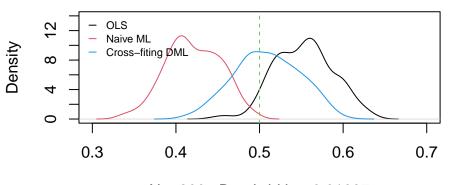
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- The transformation allows us to achieve "Neyman orthogonality," meaning the bias from estimating nuisance parameters have negligible influence on the estimation of causal parameters.
- Double machine learning is built upon the same idea as the doubly robust estimator.
- You need to get either the response surface or the propensity score correct, and you have higher efficiency by getting both correct.

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- We still have a remainder: $V_i(\hat{g}(\mathbf{X}_i) g_0(\mathbf{X}_i))$.
- This term only relies on the property of \hat{g} .
- If you use LASSO, the remainder converges to zero at a fast rate.
- For more general algorithms, we use sample splitting to eliminate it.
- As \hat{g} is generated on an independent sample, it should be orthogonal to V_i .
- We can split the sample multiple times and take the average over the estimates.
- There is no efficiency loss.



density.default(x = thetahat[, 1])



N = 200 Bandwidth = 0.01087

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- Belloni et al. (2012): use LASSO/Post-LASSO to select instruments.
- Belloni et al. (2013): use LASSO/Post-LASSO to select covariates.
- Chernozhukov et al. (2016): use LASSO/Post-LASSO to select covariates in panel data.
- Belloni et al. (2016): use double machine learning to to estimate any functional.
- Chernozhukov et al. (2018): use double machine learning to estimate nuisance parameters.

Generalization

- We have data from some samples/population
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Generalization

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- We care about the effect in other population
- Example: trials in one hospital. One cannot force patients to take the trials.
- Therefore: volunteer population and general population are different.

Generalization framework

- Use the setup from Imai, et al 2008
- Sample *n* from a finite population of N >> n.
- Insample indicator $I_i \in \{0, 1\}$. In sample if $I_i = 1$.
- (I_i, T_i, Y_i) are random variables.

Generalization framework

- Use the setup from Imai, et al 2008
- Sample *n* from a finite population of N >> n.
- Insample indicator $I_i \in \{0, 1\}$. In sample if $I_i = 1$.
- (I_i, T_i, Y_i) are random variables.
- $TE_i = Y_i(1) Y_i(0)$.
- $SATE = \frac{1}{n} \sum_{i \in \{I_i = 1\}} TE_i$
- $PATE = \frac{1}{N} \sum_{1}^{N} TE_{i}$

Estimator

- Assuming balance between treated group and control group
- Baseline estimator:

$$D = \frac{1}{n/2} \sum_{i \in \{l_i = 1, T_i = 1\}} Y_i - \frac{1}{n/2} \sum_{i \in \{l_i = 1, T_i = 0\}} Y_i$$

- Define estimation error: $\Delta = PATE D$
- Consider observed covariates X_i , unobserved covariates U_i
- $\bullet Y_i(t) = g_t(X_i) + h_t(U_i)$

Estimator

$$\Delta = \Delta_{\mathcal{S}} + \Delta_{\mathcal{T}} = \Delta_{\mathcal{S}_{\mathcal{X}}} + \Delta_{\mathcal{S}_{\mathcal{U}}} + \Delta_{\mathcal{T}_{\mathcal{X}}} + \Delta_{\mathcal{T}_{\mathcal{U}}}$$

where

$$\Delta_S = PATE - SATE$$
Sample selection

and

$$\underbrace{\Delta_T = SATE - D}_{\text{Treatment imbalance}}$$

Sample selection

- Δ_S vanish if we have census, or SATE = PATE = NATE.
- where $NATE = \frac{1}{N-n} \sum_{i \in \{I_i = 0\}} TE_i$

Sample selection

- Δ_S vanish if we have census, or SATE = PATE = NATE.
- where $NATE = \frac{1}{N-n} \sum_{i \in \{I_i=0\}} TE_i$
- Otherwise:

$$\Delta_{S_X} = \frac{N-n}{N} \left[\frac{1}{N-n} \sum_{i \in \{I_i=0\}} (g_1(X_i) - g_0(X_i)) - \frac{1}{n} \sum_{i \in \{I_i=1\}} (g_1(X_i) - g_0(X_i)) \right]$$

$$\Delta_{S_U} = \frac{N-n}{N} \left[\frac{1}{N-n} \sum_{i \in \{I_i=0\}} (h_1(U_i) - h_0(U_i)) - \frac{1}{n} \sum_{i \in \{I_i=1\}} (h_1(U_i) - h_0(U_i)) \right]$$

Sample selection (cont.)

Alternatively

$$\begin{split} &\Delta_{S_X} = \frac{N-n}{N} \int \{g_1(X) - g_0(X)\} d\{\hat{F}(X|I=0) - \hat{F}(X|I=1)\} \\ &\Delta_{S_X} = \frac{N-n}{N} \int \{h_1(U) - h_0(U)\} d\{\hat{F}(U|I=0) - \hat{F}(U|I=1)\} \end{split}$$

Treatment Imbalance

Similarly,

$$\Delta_{T_X} = \int \frac{g_1(X) + g_0(X)}{2} d\{\hat{F}(X|T=0, I=1) - \hat{F}(X|T=1, I=1)\}$$

$$\Delta_{T_U} = \int \frac{h_1(U) + h_0(U)}{2} d\{\hat{F}(U|T=0, I=1) - \hat{F}(U|T=1, I=1)\}$$

Generalization

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- ullet Usually, non-representative sample, or target population eq reference population
- Low-dimension X and RCT (without worrying U term)
 - Post-stratification:
 - Empirically get \hat{F} and $g_t(X)$ for each cell of X for target population and reference population
 - Same case if sample is non-representative

Generalization via Propensity Score (Stuart et al 2011)

- Key assumption:
 - $0 < P(I_i = 1|X_i) < 1 \text{ for all } X_i$
 - ② $I \perp \{Y(1), Y(0)\}|X \text{ or } E(\Delta_{S_{II}}) = 0$
 - **3** $T \perp \{I, Y(1), Y(0)\}|X$

Generalization via Propensity Score (Stuart et al 2011)

- Key assumption:
 - $0 < P(I_i = 1|X_i) < 1 \text{ for all } X_i$
 - **2** $I \perp \{Y(1), Y(0)\}|X \text{ or } E(\Delta_{S_{II}}) = 0$
 - **3** $T \perp \{I, Y(1), Y(0)\}|X$
- Propensity $p_i = Pr(I_i = 1|X_i)$
- Propensity difference $\Delta_p = \frac{1}{n} \sum_{i \in \{I_i = 1\}} \hat{p}_i \frac{1}{N-n} \sum_{i \in \{I_i = 0\}} \hat{p}_i$
- ullet If representative sample (random sample), $E(\Delta_p)=0$

- Otherwise:
 - IPTW
 - Full matching
 - sub-classification

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 - IPTW
 - Full matching
 - sub-classification
- Stuart et al 2011 finds they have similar results, and IPTW is slightly better.

• For IPTW, we can see:

$$\begin{split} &E\big[\frac{I(1-T)Y}{w(X)(1-e(X))}\big]\\ &=E(E\big[\frac{1(I=1)(1-1(T=1))Y}{w(X)(1-e(X))}\big|Y,X\big])\\ &=E(E\big[\frac{1(I=1)(1-1(T=1))Y(0)}{w(X)(1-e(X))}\big|Y(0),X\big])\\ &=E(\frac{Y(0)}{w(X)(1-e(X))}E\big[1(I=1)(1-1(T=1))\big|Y(0),X\big])\\ &=E(\frac{Y(0)}{w(X)(1-e(X))}P(I=1|X=x)(1-P(T=1|X=x)))\\ &=E(Y(0)) \end{split}$$

where w(X) = P(I = 1 | X = x), and e(x) = P(T = 1 | X = x)

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where
$$w(X) = P(I = 1 | X = x)$$
, and $e(x) = P(T = 1 | X = x)$

- It's possible to model them at different level, individual and context level
- It's accessible if population is well defined in dataset.

Generalization on PATT (Hartman et al. 2015)

• Hartman et al. 2015 further defines the sufficient conditions.

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- Hartman et al. 2015 further defines the sufficient conditions.
- Setup:
 - Y_{ist} represent potential outcome for a unit i assigned to study sample s
 and treatment t
 - s = 1 means reference context, and s = 0 means target population
 - t = 1 means setting the individual to treatment
 - S_i , sample indicator; T_i , treatment indicator
 - W_i^T , observed covariates related to the sample selection machenism for membership in reference group v.s. target population
 - $W_i^{C_T}$, observed covariates related to the sample assignment for inclusion of controls in reference group v.s. target population

Estimands:

•
$$\tau_{SATE} = E(Y_{11} - Y_{10}|S=1)$$

•
$$\tau_{SAT*} = E(Y_{11}|S=1, T=t) - E(Y_{10}|S=1, T=t)$$

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•
$$\tau_{PATE} = E(Y_{01} - Y_{00}|S = 0)$$

•
$$\tau_{PATC} = E(Y_{01} - Y_{00}|S = 0, T = 0)$$

•
$$\tau_{PATT} = E(Y_{01} - Y_{00}|S = 0, T = 1)$$

Assumption 1:

Consistency under parallel studies

$$Y_{i01}=Y_{i11}$$

$$Y_{i00} = Y_{i10}$$

Intuitively, potential outcome does not change by assigned contexts.

Assumption 2:

Strong Ignorability of Sample Assignment of Treated

$$\begin{split} (Y_{01},Y_{11}) \perp S | (W^T,T=1) \\ 0 < Pr(S=1|W^T,T=1) < 1 \\ \rightarrow E(Y_{s1}|S=0,T=1) = E_{01}(E(Y_{s1}|W^T,S=1,T=1)) \\ \text{for } s=0,1 \\ E_{01}(E(\cdot|W^T,...)) = E_{W^T|S=0,T=1}(E(\cdot|W^T,...)) \end{split}$$

i.e. the characteristics of **treated** units in reference population (W^T for s=1, t=1) are adjusted to match the characteristics of **treated** units in target population (s=0, t=1)

Assumption 3:

Strong Ignorability of Sample Assignment for Control

$$\begin{split} (Y_{00},Y_{10}) \perp S | (W^{C_T},T=1) \\ 0 < Pr(S=1|W^{C_T},T=1) < 1 \\ \rightarrow E(Y_{s0}|S=0,T=1) = E_{01}(E(Y_{s0}|W^{C_T},S=1,T=0)) \\ \text{for } s=0,1 \\ E_{01}(E(\cdot|W^{C_T},...)) = E_{W^{C_T}|S=0,T=1}(E(\cdot|W^{C_T},...)) \end{split}$$

i.e. the characteristics of **control** units in reference population (W^{C_T} for $s=1,\,t=1$) are adjusted to match the characteristics of **treated** units in target population ($s=0,\,t=1$)

Assumption 4:

Stable Unit Treatment Value Assumption

$$Y_{ist}^{L_i} = Y_{ist}^{L_j}, \forall$$
 treatment vectors L_i, L_j

Theorem 1 (Hartman et al. 2015)

Theorem 1. Assuming consistency and SUTVA hold, if

$$\begin{split} &E_{01}(E(Y_{s1}|W^T,S=0,T=1)) - E_{01}(E(Y_{s0}|W^{C_T},S=0,T=1)) \\ &= E_{01}(E(Y_{s1}|W^T,S=1,T=1)) - E_{01}(E(Y_{s0}|W^{C_T},S=1,T=1)) \end{split}$$

or sample assignment for treated units is strongly ignoreable given W^T , and sample assignment for controls is strongly ignoreable given W^{C_T} , then

$$au_{PATT} = E_{01}\{E(Y|W^T, S=1, T=1)\} - E_{01}\{E(Y|W^{C_T}, S=1, T=0)\}$$

Placebo Tests for Checking Assumptions

- Idea: Check whether adjusted outcome is the same as the true observed outcome in target population.
- Suppose one potential outcome is observed in target population. For example, Y_{00} is observed (S = 0, T = 0).

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- Idea: Check whether adjusted outcome is the same as the true observed outcome in target population.
- Suppose one potential outcome is observed in target population. For example, Y_{00} is observed (S = 0, T = 0).
- If assumption 1,3,4 hold, we have $E(Y|S=0, T=0) = E_{01}\{E(Y|W^{C_T}, S=1, T=0)\}$
- Similar if one can observe Y_{01} .
- Implementation: equavlent test (Hartman and Hidalgo 2011)

Estimation

- Create matched pairs or strata within reference population. (Genetic Mtaching, GenMatch)
- Reweight the matched pairs according to the characteristics of the target population. (Maximum Entropy Match, MaxEnt)

Some Extensions

- Some consider controlling context level heterogeneity
 - General meta-analysis approach: hierachical model
 - Meager 2018: bayesian hierarchical model on ATE of microcredit expansion in 7 settings
- Other post-stratification
 - MRP (Multilevel regression with poststratification, Starting from Gelman and Little 1997)
 - Even in experiment setting (Miratrix et al. 2019)
- Change estimation scheme of propensity
 - Kern et al, 2016, BART (Bayesian Additive Regression Tree)
 - Doubly robust

Reference

- Imai K, King G, Stuart E A. Misunderstandings between experimentalists and observationalists about causal inference[J].
 Journal of the royal statistical society: series A (statistics in society), 2008, 171(2): 481-502.
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The End

Good luck with your final!