Causal Inference Methods and Case Studies

STAT24630

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Lecture 5

Topic: Classical randomized experiments

- Neyman's repeated sampling approach

  - Motivation

  - Variance calculation

  - CI and hypothesis testing

  - Fisher VS Neyman

Motivation

- Limitations of the Fisher's randomization inference
  - Do not allow heterogeneity of causal effects across individuals
  - Do not have inference for the population

- Neyman's approach
  - Allow heterogeneity of causal effects across individuals
  - Focus on estimation and inference for the average treatment effect: either just for the N samples or for the whole population (PATE)
  - Repeated sampling: sampling generated b drawing from both the population units, and from the randomization distribution of assignment vector W
  - Only provide asymptotic approximation for large N instead of the exact inference

Example: Duflo-Hanna-Ryan teacher-incentive experiment

- Conducted in rural India, designed to study the effect of financial incentives on teacher performance
- In total N = 107 single-teacher schools, 53 schools are randomly chosen and are given a salary that's tied to their attendance
- One outcome: open (proportion of times the school is open during a random visit)

Table 6.1. *Summary Statistics for Duflo-Hanna-Ryan Teacher-Incentive Observed Data*

	Variable	Control ($N_c = 54$)		Treated ($N_t = 53$)			
		Average	(S.D.)	Average	(S.D.)	Min	Max
Pre-treatment	pctprewritten	0.19	(0.19)	0.16	(0.17)	0.00	0.67
Post-treatment	open	0.58	(0.19)	0.80	(0.13)	0.00	1.00
	pctpostwritten	0.47	(0.19)	0.52	(0.23)	0.05	0.92
	written	0.92	(0.45)	1.09	(0.42)	0.07	2.22
	written_all	0.46	(0.32)	0.60	(0.39)	0.04	1.43

Example: Duflo-Hanna-Ryan teacher-incentive experiment

Standard two-sample test:

$$\hat{\tau}^{\mathrm{dif}} = 0.80 - 0.58 = 0.22$$

$$s.e. = \sqrt{\frac{0.19^2}{54} + \frac{0.13^2}{53}} \approx 0.032$$

$$95\% \ CI: [0.22 - 1.96 * 0.032, 0.22 + 1.96 * 0.032]$$

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- This calculation ignores the randomization procedure of the treatment assignment
- Can we justify this standard difference-in-means analysis from the randomization perspective?

Estimation of the sample average treatment effect

- Causal estimand: $\mathrm{SATE} = \tau_{\mathrm{fs}} = \frac{1}{N}\Sigma_{i=1}^{N}\{Y_i(1) - Y_i(0)\}$ for the sampled N units

- Difference-in-means estimator:

$$\hat{\tau}^{\mathrm{dif}} = \overline{Y}_{\mathrm{t}}^{\mathrm{obs}} - \overline{Y}_{\mathrm{c}}^{\mathrm{obs}}$$

where $\quad \overline{Y}_{\mathrm{c}}^{\mathrm{obs}} = \dfrac{1}{N_{\mathrm{c}}}\displaystyle\sum_{i:W_i=0} Y_i^{\mathrm{obs}} \quad$ and $\quad \overline{Y}_{\mathrm{t}}^{\mathrm{obs}} = \dfrac{1}{N_{\mathrm{t}}}\displaystyle\sum_{i:W_i=1} Y_i^{\mathrm{obs}}$

- Under complete randomization (random W) and treat the potential outcomes as fixed (fixed $\boldsymbol{Y}(0) = \{Y_i(0), i = 1, \cdots, N\}$ and $\boldsymbol{Y}(1) = \{Y_i(1), i = 1, \cdots, N\}$), this estimator is unbiased

$$\mathbb{E}_W\left[\hat{\tau}^{\mathrm{dif}}\,\middle|\,\boldsymbol{Y}(0), \boldsymbol{Y}(1)\right] = \frac{1}{N}\sum_{i=1}^{N}\left(\frac{\mathbb{E}_W[W_i]\cdot Y_i(1)}{N_{\mathrm{t}}/N} - \frac{\mathbb{E}_W[1 - W_i])\cdot Y_i(0)}{N_{\mathrm{c}}/N}\right)$$

$$= \frac{1}{N}\sum_{i=1}^{N}(Y_i(1) - Y_i(0)) = \tau_{\mathrm{fs}}.$$

First, we can re-write $\hat{\tau}^{dif}$:

$$
\begin{aligned}
\hat{\tau}^{dif} &= \overline{Y_t^{obs}} - \overline{Y_c^{obs}} \\
&= \frac{1}{N_t} \sum_{i:W_i=1} Y_i^{obs} - \frac{1}{N_c} \sum_{i:W_i=0} Y_i^{obs} \\
&= \frac{1}{N_t} \sum_{i:W_i=1} W_i Y_i(1) - \frac{1}{N_c} \sum_{i:W_i=0} (1-W_i) Y_i(0) \\
&= \frac{1}{N} \sum_{i=1}^{N} \left(\frac{N}{N_t} W_i Y_i(1) - \frac{N}{N_c} (1-W_i) Y_i(0) \right)
\end{aligned}
$$

And now we can take an expectation:

$$
\begin{aligned}
E(\hat{\tau}^{dif}) &= \frac{1}{N} \sum_{i=1}^{N} \left(\frac{N}{N_t} E(W_i) Y_i(1) - \frac{N}{N_c} E(1-W_i) Y_i(0) \right) \\
&= \frac{1}{N} \sum_{i=1}^{N} \left(\frac{N}{N_t} \frac{N_t}{N} Y_i(1) - \frac{N}{N_c} \frac{N_c}{N} Y_i(0) \right) \\
&= \frac{1}{N} \sum_{i=1}^{N} (Y_i(1) - Y_i(0)) \\
&= \tau
\end{aligned}
$$

Calculate the variance of the estimator

- Causal estimand: $\mathrm{SATE} = \tau_{\mathrm{fs}} = \frac{1}{N}\sum_{i=1}^{N}\{Y_i(1) - Y_i(0)\}$ for the sampled N units

- Difference-in-means estimator: $\hat{\tau}^{\mathrm{dif}} = \overline{Y}_t^{\mathrm{obs}} - \overline{Y}_c^{\mathrm{obs}}$

- Under complete randomization and fixed potential outcomes, we can also calculate the variance of $\hat{\tau}^{\mathrm{dif}}$ (if you are interested in the proof see Appendix A of Chapter 6)

$$V_W[\hat{\tau}^{\mathrm{dif}}|\boldsymbol{Y}(0), \boldsymbol{Y}(1)] = \frac{S_c^2}{N_c} + \frac{S_t^2}{N_t} - \frac{S_{ct}^2}{N}$$

where

$$S_c^2 = \frac{1}{N-1}\sum_{i=1}^{N}(Y_i(0) - \overline{Y}(0))^2, \quad \text{and} \quad S_t^2 = \frac{1}{N-1}\sum_{i=1}^{N}(Y_i(1) - \overline{Y}(1))^2$$

Sample variance of $Y_i(0)$ and $Y_i(1)$

$$S_{ct}^2 = \frac{1}{N-1}\sum_{i=1}^{N}(Y_i(1) - Y_i(0) - (\overline{Y}(1) - \overline{Y}(0)))^2$$

$$= \frac{1}{N-1}\sum_{i=1}^{N}(Y_i(1) - Y_i(0) - \tau_{\mathrm{fs}})^2.$$

Sample variance of the unit-level treatment effects

Conservative approximation of the variance of the estimator

- $\mathbb{V}_W[\hat{\tau}^{\text{dif}}|Y(0), Y(1)] = \dfrac{S_c^2}{N_c} + \dfrac{S_t^2}{N_t} - \dfrac{S_{ct}^2}{N}$

  where

  $$S_c^2 = \frac{1}{N-1}\sum_{i=1}^{N}(Y_i(0) - \bar{Y}(0))^2, \quad \text{and} \quad S_t^2 = \frac{1}{N-1}\sum_{i=1}^{N}(Y_i(1) - \bar{Y}(1))^2$$

  Sample variance of $Y_i(0)$ and $Y_i(1)$

  $$S_{ct}^2 = \frac{1}{N-1}\sum_{i=1}^{N}(Y_i(1) - Y_i(0) - (\bar{Y}(1) - \bar{Y}(0)))^2$$

  $$= \frac{1}{N-1}\sum_{i=1}^{N}(Y_i(1) - Y_i(0) - \tau_{\text{fs}})^2.$$

  Sample variance of the unit-level treatment effects

- Estimate S_c^2 and S_t^2 by sample variance of observed outcomes

  $$s_c^2 = \frac{\sum_{i:W_i=0}(Y_i^{\text{obs}} - \bar{Y}_c^{\text{obs}})^2}{N_c - 1}, s_c^2 = \frac{\sum_{i:W_i=1}(Y_i^{\text{obs}} - \bar{Y}_t^{\text{obs}})^2}{N_t - 1}$$

- S_{ct}^2 is not identifiable
  - No heterogeneity of treatment effects across individuals $S_{ct}^2 = 0$
  - In general, $S_{ct}^2 \geq 0$ though the exact value is unknown

Conservative approximation of the variance of the estimator

- $\mathbb{V}_W[\hat{\tau}^{\text{dif}}|\boldsymbol{Y}(0), \boldsymbol{Y}(1)] = \dfrac{S_c^2}{N_c} + \dfrac{S_t^2}{N_t} - \dfrac{S_{ct}^2}{N}$

  where

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  Sample variance of $Y_i(0)$ and $Y_i(1)$

  $$S_{ct}^2 = \frac{1}{N-1}\sum_{i=1}^{N}(Y_i(1) - Y_i(0) - (\overline{Y}(1) - \overline{Y}(0)))^2$$

  $$= \frac{1}{N-1}\sum_{i=1}^{N}(Y_i(1) - Y_i(0) - \tau_{\text{fs}})^2.$$

  Sample variance of the unit-level treatment effects

- A conservative estimator of $\text{Var}_W[\hat{\tau}^{\text{dif}}|\boldsymbol{Y}(0), \boldsymbol{Y}(1)]$

  $$\mathbb{V}_W[\hat{\tau}^{\text{dif}}|\boldsymbol{Y}(0), \boldsymbol{Y}(1)] \leq \frac{S_c^2}{N_c} + \frac{S_t^2}{N_t} = \mathbb{E}_W\left[\frac{s_c^2}{N_c} + \frac{s_t^2}{N_t}\Big|\boldsymbol{Y}(0), \boldsymbol{Y}(1)\right]$$

  Neyman's estimator of the variance, same as *s.e.* on slide 5

Estimation of the population average treatment effect

- Causal estimand: $\mathrm{PATE} = \tau_{\mathrm{sp}} = \mathbb{E}\big(Y_i(1) - Y_i(0)\big) = \mathbb{E}(\mathrm{SATE}) = \mathbb{E}(\tau_{\mathrm{fs}})$
- We assume that $(Y_i(0), Y_i(1))$ are jointly i.i.d samples from a super population with variance σ_c^2 and σ_s^2
- We still use difference-in-means estimator:

$$\hat{\tau}^{\mathrm{dif}} = \overline{Y}_t^{\mathrm{obs}} - \overline{Y}_c^{\mathrm{obs}}$$

- $\hat{\tau}^{\mathrm{dif}}$ is still unbiased for τ_{sp}: $\mathbb{E}\big(\hat{\tau}^{\mathrm{dif}}\big) = \mathbb{E}(\mathbb{E}_W[\hat{\tau}^{\mathrm{dif}}|\boldsymbol{Y}(0), \boldsymbol{Y}(1)]) = \mathbb{E}(\tau_{\mathrm{fs}}) = \tau_{\mathrm{sp}}$

- The variance of $\hat{\tau}^{\mathrm{dif}}$ (variance decomposition formula):
  - Check Wikipedia if you do not know the variance decomposition formula
    https://en.wikipedia.org/wiki/Law_of_total_variance

$$\mathbb{V}\big(\hat{\tau}^{\mathrm{dif}}\big) = \mathbb{E}\big(\mathbb{V}_W\big[\hat{\tau}^{\mathrm{dif}}|\boldsymbol{Y}(0), \boldsymbol{Y}(1)\big]\big) + \mathbb{V}(\mathbb{E}_W[\hat{\tau}^{\mathrm{dif}}|\boldsymbol{Y}(0), \boldsymbol{Y}(1)])$$

Variance calculation for the population

$$\mathbb{V}(\hat{\tau}^{\text{dif}}) = \mathbb{E}\big(\mathbb{V}_W\big[\hat{\tau}^{\text{dif}}|\boldsymbol{Y}(0), \boldsymbol{Y}(1)\big]\big) + \mathbb{V}(\mathbb{E}_W[\hat{\tau}^{\text{dif}}|\boldsymbol{Y}(0), \boldsymbol{Y}(1)])$$

- $\mathbb{V}_W\big[\hat{\tau}^{\text{dif}}|\boldsymbol{Y}(0), \boldsymbol{Y}(1)\big] = \dfrac{S_c^2}{N_c} + \dfrac{S_t^2}{N_t} - \dfrac{S_{ct}^2}{N}$, with
$$\mathbb{E}(S_c^2) = \sigma_c^2, \mathbb{E}(S_t^2) = \sigma_t^2, \mathbb{E}(S_{ct}^2) = \mathbb{V}(Y_i(1) - Y_i(0))$$

- $\mathbb{V}\big(\mathbb{E}_W[\hat{\tau}^{\text{dif}}|\boldsymbol{Y}(0), \boldsymbol{Y}(1)]\big) = \mathbb{V}(\tau_{\text{fs}}) = \mathbb{V}\left(\dfrac{1}{N}\sum_{i=1}^{N}\{Y_i(1) - Y_i(0)\}\right) = \dfrac{1}{N}\mathbb{V}(Y_i(1) - Y_i(0))$

- So $\mathbb{V}_W\big(\hat{\tau}^{\text{dif}}\big) = \dfrac{\sigma_c^2}{N_c} + \dfrac{\sigma_t^2}{N_t}$ exactly the same as in two-sample testing
  - In two-sample testing, we assume that observed outcome Y_i are i.i.d. in the treatment group and Y_i are i.i.d. in the control group
  - Under complete randomization, $Y_i = Y_i(W_i)$ are not i.i.d. even with the treatment/control group because W_i are negatively correlated across i

Construct confidence intervals for τ_{fs} or τ_{sp}

- We have the same estimator $\hat{\tau}^{\mathrm{dif}}$ and the same variance approximation of $\hat{\tau}^{\mathrm{dif}}$

$$\widehat{\mathbb{V}}(\hat{\tau}^{\mathrm{dif}}) = \frac{s_c^2}{N_c} + \frac{s_t^2}{N_t}$$

no matter we are interested about SATE τ_{fs} or PATE τ_{sp}

- When N is large enough, we can approximate the distribution of $\hat{\tau}^{\mathrm{dif}}$ by a normal distribution

- Then the 95% CI for either τ_{fs} or τ_{sp} is
$[\hat{\tau}^{\mathrm{dif}} - 1.96 \times \sqrt{\widehat{\mathbb{V}}(\hat{\tau}^{\mathrm{dif}})}, \hat{\tau}^{\mathrm{dif}} - 1.96 \times \sqrt{\widehat{\mathbb{V}}(\hat{\tau}^{\mathrm{dif}})}]$
same as we we had earlier

Hypothesis testing for τ_{fs} or τ_{sp}

- We have the same estimator $\hat{\tau}^{\text{dif}}$ and the same variance approximation of $\hat{\tau}^{\text{dif}}$

$$\widehat{\mathbb{V}}(\hat{\tau}^{\text{dif}}) = \frac{s_c^2}{N_c} + \frac{s_t^2}{N_t}$$

  no matter we are interested about SATE τ_{fs} or PATE τ_{sp}

- When N is large enough, we can approximate the distribution of $\hat{\tau}^{\text{dif}}$ by a normal distribution

- When can test for the null hypothesis $\boldsymbol{H_0: \tau_{fs} = 0}$ or $\boldsymbol{H_0: \tau_{sp} = 0}$

- The t-statistics: $t = \dfrac{\hat{\tau}^{\text{dif}}}{\sqrt{\widehat{\mathbb{V}}(\hat{\tau}^{\text{dif}})}}$

- Under either H_0 and when N is large, we have t approximately follows a $N(0, 1)$ distribution
- Two-sided p-value: $2(1 - \phi(|t|))$

Application to the Duflo-Hanna-Ryan data

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	written	0.92	(0.45)	1.09	(0.42)	0.07	2.22
	written_all	0.46	(0.32)	0.60	(0.39)	0.04	1.43

Confidence interval for each of the four outcomes:

$\hat{\tau}$	$\widehat{(\text{s.e.})}$	95% C.I.
0.22	(0.03)	(0.15,0.28)
0.05	(0.04)	($-$0.03,0.13)
0.17	(0.08)	(0.00,0.34)
0.14	(0.07)	(0.00,0.28)

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Analysis on two different subgroups: within each subgroup we still have complete randomization of assignments

Variable	pctpre $= 0$ ($N = 40$)			pctprewritten > 0 ($N = 67$)			Difference		
	$\hat{\tau}$	(s.e.)	95% C.I.	$\hat{\tau}$	(s.e.)	95% C.I.	EST	(s.e.)	95% C.I.
open	0.23	(0.05)	(0.14,0.32)	0.21	(0.04)	(0.13,0.29)	0.02	(0.06)	(−0.10,0.14)
pctpost written	−0.004	(0.06)	(−0.16,0.07)	0.11	(0.05)	(0.01,0.21)	−0.15	(0.08)	(−0.31,0.00)
written	0.20	(0.10)	(0.00,0.40)	0.18	(0.10)	(−0.03,0.38)	0.03	(0.15)	(−0.26,0.31)
written _all	0.04	(0.07)	(−0.10,0.19)	0.22	(0.09)	(0.04,0.40)	−0.18	(0.12)	(−0.41,0.05)

Fisher v.s. Neyman

- Like Fisher, Neyman proposed randomization-based inference

- Unlike Fisher,
    - estimands are average treatment effects
    - heterogenous treatment effects are allowed
    - population as well as sample inference is possible
    - asymptotic approximation is required for inference

- Fisher's approach can easily be applied to deal with any randomization mechanism in an experiment, but it can be much harder for Neyman's approach