Day 10: Noncompliance

Erin Rossiter

March, 2022

- Draft of final paper due tomorrow at 3pm

- » Github and/or email
- » Saving time today to talk about them
- Next week
 - » Online markets & related issues (demand effects, manipulation checks, attention checks)
 - » Attrition (moved!)
 - » Let me know about any other topics
- Methods workshop on Friday
 - » please tell other first years to come!
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- Difference-in-means is an **unbiased** estimator of the ATE assuming:
 - 1. Randomization of treatment
 - » $E[Y_i(1)|D_i=1] = E[Y_i(1)]$
 - » $E[Y_i(0)|D_i=0] = E[Y_i(0)]$
 - » We can estimate left-hand terms using our observed data!
 - 2. Excludability
 - » what does this mean?
 - 3. Noninterference
 - » what does this mean?
- But, things don't always go as planned.

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 - » one-sided, or failure-to-treat
 - some subject assigned to treatment aren't actually treated
 - » two-sided
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What's the difference between noncompliance and attrition?

Key difference:

- noncompliance
 - » means you measured the outcomes for everyone, but people received treatments they weren't assigned to
- attrition
 - » is more a missing data problem; everyone complied, you just didn't get Y measured for everyone

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(Board 1)
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Canvassing experiment

- » People assigned to treatment don't answer the door, or canvassers mess up and don't knock!
- » Should be treated (canvassed), but instead doesn't receive treatment
- Social interaction experiment
 - » I ask people to talk about gun control, but they talk about meaning of life (control topic)
 - Should be treated (political convo), but instead doesn't receive treatment
- Drug trial
 - » People assigned to take a pill don't do it
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One-sided noncompliance

- Treatment: face-to-face canvassing
- Outcome: voter turnout
- Suppose 1,000 assigned to treatment, 1,000 assigned to control
- BUT only 250 assigned to treatment are actually reached by canvassers
 - » Gives us 3 groups of subjects:
 - 1. 250 assigned to treatment who actually were treated
 - 2. 750 assigned to treatment who remain untreated
 - 3. 1,000 (untreated) in the control group

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Sequence of an experiment $Z \rightarrow D \rightarrow Y$

- Z (treatment assignment)
- D (treatment actually received)
- Y (outcome)

In the voter mobilization experiment example, what is the difference when thinking about the:

- causal effect of Z on Y vs
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Compliers' "compliance" potential outcomes

- $-d_i(z=1)=1$
 - » receive treatment if assigned to treatment
 - $-d_i(z=0)=0$
 - » do not receive treatment if assigned to control

Never-takers' "compliance" potential outcomes:

- $-d_i(z=1)=0$
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- $-d_i(z=1)=0$
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z is the assigned treatment and $d_i(z)$ is a function of the actual treatment that results from an assignment

Compliers' "compliance" potential outcomes:

- $d_i(z=1) = 1$
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- $d_i(z=0) = 0$
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Visual

D(0)	D(1)	Type
0	1	Complier
0	0	Never-taker

DeclareDesign for potential outcomes

Defining causal effects

Notation note: \mathbf{z}' is a vector of all treatment assignments, possibly altered except for i

-z=z' i keeps same treatment assignment even if others' change

Part A

$$d_i(\mathbf{z}) = d_i(\mathbf{z}')$$
 if $\mathbf{z} = \mathbf{z}'$

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Part B:

$$Y_i(\mathbf{z}, \mathbf{d}) = Y_i(\mathbf{z}', \mathbf{d}') \ \mathbf{z_i} = \mathbf{z_i}' \ \text{and} \ \mathbf{d_i} = \mathbf{d_i}$$

assignment and what I do given that assignment

» my potential outcomes don't depend on other peoples' z and a

Can you think of examples that violate this assumption?

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- basically, my potential outcomes depend only on my assignment and what I do given that assignment
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Intent-to-treat effect of z_i on d_i for each subject:

$$ITT_{i,D} = d_i(1) - d_i(0)$$

Average Intent-to-treat effect (assuming one-sided noncompliance):

$$ITT_D = E[ITT_D] = E[d_i(1)] - E[d_i(0)] = E[d_i(1)] - 0 = E[d_i(1)]$$

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If an experiment has 100% compliance, what quantity is the ITT_Y equivalent to?

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We usually *want* to estimate ATE, but we don't have enough info to when there's noncompliance:

$$ATE = \frac{1}{N} \sum_{i=1}^{N} (Y_i(1) - Y_i(0)) = E[Y_i(d=1) - Y_i(d=0)]$$

Complier average causal effect (CACE) is more realistic to estimate

$$CACE = \frac{\sum_{i=1}^{N} (Y_i(1) - Y_i(0)) d_i(1)}{\sum_{i=1}^{N} d_i(1)}$$
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Calculating estimands

(Board 3) let's calculate each estimand

DeclareDesign for estimands

- We are limited in what we can learn from experiments with non-compliance
- And of course, we don't have full potential outcomes!
- What estimators work?
 - » no estimator for ATE :(
 - » unbiased estimator for ITT OLS or DIM
 - » unbiased estimator for ITT_D OLS or DIM
 - » consistent estimator for $CACE = ITT/ITT_D$ (See Theorem 5.1) 2SLS

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Table 5.2, pg 150, New Haven voter mobilization experiment

- group means
- N size for group in parentheses

	Treatment Grp	Control Grp
Turnout rate among those contacted by canvassers	53.43 (395)	
Turnout rate among those not contacted by canvassers Overall turnout rate	36.48 (1050) 41.38 (1445)	37.54 (5645) 37.52 (5645)

(Board 4)

- What is the cell we're not used to seeing?
- What is the $I\hat{T}T$?
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- What is the $I\hat{T}T$?
- What is the ITT_D ?

- What is the CACE?

Table 5.2, pg 150, New Haven voter mobilization experiment

- group means
- N size for group in parentheses

	Treatment Grp	Control Grp
Turnout rate among those contacted by canvassers	53.43 (395)	
Turnout rate among those not contacted by canvassers	36.48 (1050)	37.54 (5645)
Overall turnout rate	41.38 (1445)	37.52 (5645)

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2SLS (or IV estimation)

2SLS is a consistent estimator for $CACE = ITT/ITT_D$ (See Theorem 5.1)

```
Model1: TREATED_i = \alpha_0 + \alpha_1 ASSIGNED_i + \epsilon_i
Model2: VOTED_i = \beta_0 + \beta_1 TREATED_i + u_i
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- What parameter is the CACE?
- Exclusion restriction: ASSIGNED affects VOTED only through TREATED
- Instrument: ASSIGNED
 - » predicts TREATED
 - » independent of u

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DeclareDesign for estimation + simulation

Two-sided noncompliance

Two-sided noncompliance

D(Z=0)	D(Z=1)	Туре
0	1	Complier
0	0	Never-taker
1	1	Always-taker
1	0	Defier

- still able to estimate CACE, ITT, and ITT_D
- comes up when subjects have access to treatments and discretion for whether to take them
 - » not a problem in survey experiments

You must be able to distinguish between assigned and actual treatment when estimating causal effects (i.e., these need to be columns in your data)

Be sure to estimate ITT or CACE

- these utilize random assignment and we can get unbiased/consistent estimates of our estimands
- yes, it changes interpretation of estimates
 - » ITT effect of assignment on outcomes
 - » CACE ATE among compliers
 - who are these people? who can I generalize to?
- avoid estimating ATE based on what units received not random! prone to bias

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