Lecture 6: When Complete Randomization Fails Attrition after random assignment

Xi Chen

Rotterdam School of Management Erasmus University Rotterdam

June 3, 2023

Outline

1 Attrition and the conventional approaches

2 Bounding approach

3 Bounding approach in attrition



Outline

1 Attrition and the conventional approaches

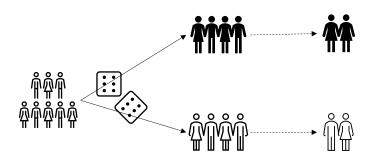
2 Bounding approach

3 Bounding approach in attrition



Failure to measure outcomes

The general idea of attrition: the failure to measure outcomes of some units.





Various reasons of attrition



Refuse to cooperate (two-stage studies)



(long-term studies)



Blocked by authorities (field experiments)



Unmeasurable (e.g., decease)



Attrition: an example

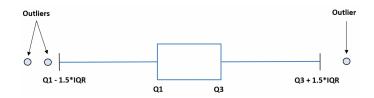
The introduction of e-commerce to countryside in China¹.



Attrition: local governments denied access to some villages.

RSM

Attrition: the deletion of outliers in experimental data



Should we delete outliers in experimental data?

- Deletion leads to missing outcomes for some units.
- Leading to the failure of randomization, eventually non-identification of ATE.



Attrition: the deletion of outliers in experimental data

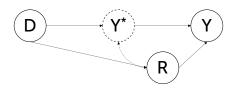


Should we delete outliers in experimental data?

- Deletion leads to missing outcomes for some units.
- Leading to the failure of randomization, eventually non-identification of ATE.



A DAG representation of attrition



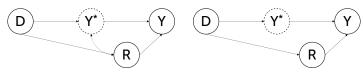
- *D* is the treatment, and **exogenous due to randomization**.
- lacksquare Y^* is a latent outcome, partially observed by researchers.
- Arr $R = \{0,1\}$ is the binary state of attrition, caused by or causing Y^* .
- Y is the observed outcome with $Y = (1 R) \times Y^*$

Fact (Intention-to-treat)

Without any assumption, we can identify the intention-to-treat effects with $P(Y \mid D)$.

Traditional approaches: Random attrition

Assuming the attrition is random...



- (a) DAG of Non-random Attrition
- (b) DAG of Random Attrition

Fact (Random Attrition)

Under random attrition, we can identify the average treatment effect with $P(Y^* \mid D) = P(Y^* \mid D, R)$.



Traditional approaches: Random attrition

Technical proof:

$$P(Y^* \mid do(D)) = P(Y^* \mid D, R = 0) P(R = 0) + P(Y^* \mid D, R = 1) P(R = 1)$$

$$\stackrel{Y^* \perp R \mid D}{=} P(Y^* \mid D) P(R = 0) + P(Y^* \mid D) P(R = 1)$$

$$= P(Y^* \mid D)$$

Because the attrition is at random, we can use observed outcome Y to calculate the conditional distribution:

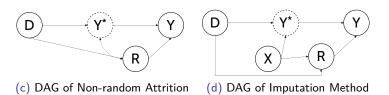
$$P(Y^* | D) = P(Y^* | D, R = 0) = P(Y | D)$$



Traditional approaches: imputation

Assumption for Imputation

We observe **non-descendant** variables X that **separate** Y^* **and** R.



Fact (Imputation Method`

Under the assumption of the imputation method, we can identify the average treatment effect with

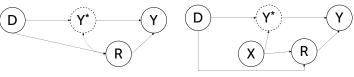
$$P(Y^* | D, X) = P(Y^* | D, X, R).$$



Traditional approaches: imputation

Assumption for Imputation

We observe **non-descendant** variables X that **separate** Y^* **and** R.



(e) DAG of Non-random Attrition (f) DAG of Imputation Method

Fact (Imputation Method)

Under the assumption of the imputation method, we can identify the average treatment effect with $P(Y^* | D, X) = P(Y^* | D, X, R).$



Traditional approaches: imputation

Technical proof:

$$P(Y^* \mid do(D), X) = P(Y^* \mid D, X, R = 0) P(R = 0) + P(Y^* \mid D, X, R = 1) P(R = 0)$$

$$\stackrel{Y^* \perp R \mid X}{=} P(Y^* \mid D, X) P(R = 0) + P(Y^* \mid D, X) P(R = 1)$$

$$= P(Y^* \mid D, X)$$

Because the attrition is at random conditional on X, we can use observed outcome Y to calculate the conditional distribution:

$$P(Y^* | D, X) = P(Y^* | D, X, R = 0) = P(Y | D, X)$$

We can then integrate over X to have the ATE:

$$P(Y \mid D, X) = \sum_{x \in X} P(Y \mid D, X = x)$$



Issues of the traditional approaches

The law of credibility

The credibility of inference decreases with the strength of the assumptions maintained.

Assumptions are strong and cannot be justified. Credibility is sacrificed for the "power" of conclusions.

- The random attrition is hardly a credible assumption.
- Difficult to find a set of X that d-separate Y^* and R.

Could we adopt a better strategy?



Outline

1 Attrition and the conventional approaches

2 Bounding approach

3 Bounding approach in attrition



Relaxing unrealistic assumptions

Many assumptions are non-refutable and unrealistic.

- Unconfoundedness
- Random attrition
- **...**

Forego the point identification, make weaker assumptions and increase credibility of analyses.

Strong assumptions identify a point: \bullet Weak assumptions identify a set: [---]



Some types of bounds: no-data bounds

Assume the potential outcomes Y^0 and Y^1 are bounded [0,1].

$$-1 \leq Y_i^1 - Y_i^0 \leq 1 \Rightarrow E\left(Y_i^1 - Y_i^0\right) \in [-1, 1]$$

In more general terms, if Y^0 and $Y^1 \in [a,b]$

$$E\left(Y_i^1 - Y_i^0\right) \in [a - b, b - a]$$

The length of the bound is 2(b-a).



Some types of bounds: no-assumption bounds

Assume we collect some data with $\{Y_i, D_i\}$, but random assignment is not ensured².

Can we do better with data?

$$E\left(Y_{i}^{1}-Y_{i}^{0}\right) = \underbrace{\pi E\left(Y_{i}^{1}\mid D_{i}=1\right)}_{\text{Observed}} + \underbrace{\left(1-\pi\right)}_{\text{Unobserved}} \underbrace{E\left(Y_{i}^{1}\mid D_{i}=0\right)}_{\text{Unobserved}} - \underbrace{\pi E\left(Y_{i}^{0}\mid D_{i}=1\right)}_{\text{Unobserved}} - \underbrace{\left(1-\pi\right)}_{\text{Observed}} \underbrace{E\left(Y_{i}^{0}\mid D_{i}=0\right)}_{\text{Observed}}$$

Under the bounds on potential outcomes [a, b]:

$$au^{obs} + (1-\pi) a - \pi b \leq E\left(Y_i^1 - Y_i^0\right) \leq au^{obs} + (1-\pi) b - \pi a$$

The length of the bound is (b-a), half of the no-data bounds.

²Let $\pi = P(D_i = 1)$ in the data collected.



Some types of bounds: monotone treatment response

Assume treatment always benefit people, or $Y_i^1 \geq Y_i^0, \forall i$ This means ITE is nonnegative, so the lower bound of ITE is 0. Under this assumption, the lower bound of ATE is also 0:

$$E\left(Y_i^1-Y_i^0\right)\geq 0$$

Using this assumption, we may further narrow the no-data and no-assumptions bounds.

Fact

Under monotone treatment response assumption, the no-data bound becomes $E(Y_i^1 - Y_i^0) \in [0, b - a]$.



Some types of bounds: monotone treatment selection

Assume the treatment group potential outcomes are better than control groups, with

$$E\left(Y_{i}^{1}\mid D_{i}=1\right) \geq E\left(Y_{i}^{1}\mid D_{i}=0\right)$$

$$E\left(Y_{i}^{0}\mid D_{i}=1\right) \geq E\left(Y_{i}^{0}\mid D_{i}=0\right)$$

Under this assumption, the ATE is bounded from above,

$$E\left(Y_{i}^{1}-Y_{i}^{0}\right)\leq E\left(Y_{i}^{1}\mid D_{i}=1\right)-E\left(Y_{i}^{0}\mid D_{i}=0\right)$$

We may use this assumption to further narrow the no-assumption bounds.



Outline

1 Attrition and the conventional approaches

2 Bounding approach

3 Bounding approach in attrition



A running example

- We have a data from a randomized experiment.
- The outcome Y_i measured with a scale from 1 to 10 (i.e., $Y_i \in [1, 10]$).
- The experiment has an attrition problem, with NA denoting missing values.

Obs	D_i	Yi	
1	1	10	
2	1	8	
3	1	NA	
4	0	3	
5	0	NA	
6	0	6	
$Y_i \in [0, 10]$			



A running example

Obs	D_i	Y_i	
1	1	10	
2	1	8	
3	1	NA	
4	0	3	
5	0	NA	
6	0	6	
$Y_i \in [0, 10]$			

- Without additional assumptions, the intention-to-treat is: $ITT_Y = \frac{10+8}{3} \frac{3+6}{3} = 3.$
- Under random attrition, the ATE is: ATE_{random} = $\frac{10+8}{2} \frac{3+6}{2} = 4.5$.



Bounding approach: Manski bounds

Manski bounds have no additional assumptions. See Horowitz and Manski (2000).

- Upper bound: substituting NA's in treatment with max values and control with min values.
- Lower bound: substituting NA's in treatment with min values and treatment with max values.

Obs	Di	Yi
1	1	10
2	1	8
3	1	NA
4	0	3
5	0	NA
6	0	6
$Y_i \in [0, 10]$		

RSM

Bounding approach: Manski bound

Obs	D_i	Y_i
1	1	10
2	1	8
3	1	NA
4	0	3
5	0	NA
6	0	6

■ Manski bound:
$$\left[\frac{10+8+1}{3} - \frac{3+10+6}{3}, \frac{10+8+10}{3} - \frac{3+1+6}{3}\right] = [0,6]$$

■ $ITT_Y = 3$ and $ATE_{random} = 4.5$

In practice, use max and min values of observed outcomes.

Uninformative but wide, as a benchmark for further analysissm

Can we narrow the bound?



Lee bounds Lee (2009)

Assumptions: monotonicity of treatment selection.

- Treatment status either increases / decreases attrition for all people.
- The effect of treatment on attrition has the same sign (+ or −).

How to use the condition?

- Trimming samples such that share of observed individuals is equal in the treatment and control.
- Trimming from above or below to obtain lower or upper bound.



- Suppose we have a treatment group with 40% of attrition and a control group with 50%.
- The treatment decreases the attrition likelihood for 10%.
- To get the same level of "missing," we need to trim x% of people from the treatment.
- To calculate x%, we do:

$$\frac{40\%}{1-x\%} = 50\% \Rightarrow x\% = \frac{50\% - 40\%}{50\%} = 20\%$$



- Suppose we have a treatment group with 40% of attrition and a control group with 50%.
- The treatment decreases the attrition likelihood for 10%.
- To get the same level of "missing," we need to trim x% of people from the treatment.
- To calculate x%, we do:

$$\frac{40\%}{1-x\%} = 50\% \Rightarrow x\% = \frac{50\% - 40\%}{50\%} = 20\%$$



- Suppose we have a treatment group with 40% of attrition and a control group with 50%.
- The treatment decreases the attrition likelihood for 10%.
- To get the same level of "missing," we need to trim x% of people from the treatment.
- To calculate x%, we do:

$$\frac{40\%}{1-x\%} = 50\% \Rightarrow x\% = \frac{50\% - 40\%}{50\%} = 20\%$$



- Suppose we have a treatment group with 40% of attrition and a control group with 50%.
- The treatment decreases the attrition likelihood for 10%.
- To get the same level of "missing," we need to trim x% of people from the treatment.
- To calculate x%, we do:

$$\frac{40\%}{1 - x\%} = 50\% \Rightarrow x\% = \frac{50\% - 40\%}{50\%} = 20\%$$



Question: which 20% of the treated units to trim?

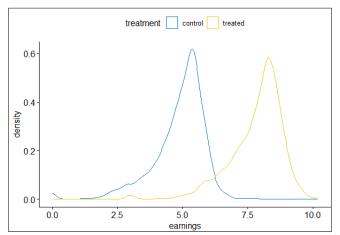
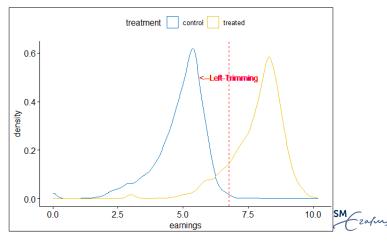
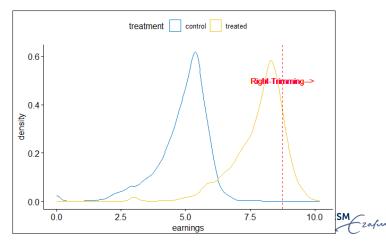


Figure: Density Plots of Outcomes for the Treated and Control Group of Con

To trim the people in the 20% lower quantile to get the upper bound:



To trim the people in the 20% upper quantile to get the lower bound:



Lee bounds: procedure

Y the outcomes, D the treatment, and W the non-attrition.

- Step 1: Get the share of individuals with observed Y in the treatment and control, as q_1 and q_0 .
- Step 2: Obtain the attrition difference:

$$q = egin{cases} rac{q_1 - q_0}{q_1} & ext{if } q_1 > q_0 \ rac{q_0 - q_1}{q_0} & ext{if } q_0 > q_1 \end{cases}$$

■ Step 3: Find the qth and (1-q)th quantile of observed outcomes in the treatment group as Y_q^1 and Y_{1-q}^1 .



Lee bounds: procedure

Y the outcomes, D the treatment, and W the non-attrition.

■ Step 4: Upper and lower bound:

$$ar{ au} = \underbrace{\frac{\sum \mathbb{1}\left(Y_i \geq Y_q^1\right) D_i W_i Y_i}{\sum \mathbb{1}\left(Y_i \geq Y_q^1\right) D_i W_i}}_{\text{Non-attriters in control}} - \underbrace{\frac{\sum (1 - D_i) W_i Y_i}{\sum (1 - D_i) W_i}}_{\text{Non-attriters in control}}$$

Non-attritors in treatment with higher outcome

$$\underline{\tau} = \underbrace{\frac{\sum 1 \left(Y_i \le Y_{1-q}^1 \right) D_i W_i Y_i}{\sum 1 \left(Y_i \le Y_{1-q}^1 \right) D_i W_i}}_{\text{Non-attriters in control}} - \underbrace{\frac{\sum (1 - D_i) W_i Y_i}{\sum (1 - D_i) W_i}}_{\text{Non-attriters in control}}$$

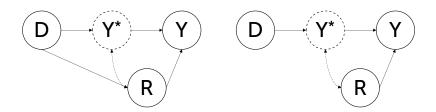
Non-attritors in treatment with lower outcome

RSM

Lee bounds: summary

Treatment influences attrition in one direction:

- By comparing the treatment and control, we may "partial out" the effect of treatment on attrition.
- ATE on the non-attritors!



References I

- Couture, Victor, Benjamin Faber, Yizhen Gu, and Lizhi Liu. 2021. "Connecting the Countryside via E-Commerce: Evidence from China." American Economic Review: Insights, 3 (1): 35-50.
- ► Horowitz, J. L., & Manski, C. F. (2000). Nonparametric analysis of randomized experiments with missing covariate and outcome data. Journal of the American statistical Association, 95(449), 77-84.
- Lee, D. S. (2009). Training, wages, and sample selection: Estimating sharp bounds on treatment effects. The Review of Economic Studies, 76(3), 1071-1102.

