

# EE19MTECH01008 - Assignment 3

Wednesday, 5 May 2021 10:07 AM

3) Exclusion Restriction  $\rightarrow$  Z causal effect on Y is fully mediated by T. This assumption excludes Z from structural equation of Y and from any other structural equations that would make causal association flow from Z to Y without going through T.

Relevance  $\rightarrow$  Z has causal effect on T. It corresponds to the existence of an active edge from Z to T in the causal graph.

Exogeneity  $\rightarrow$  Given a simple regression model

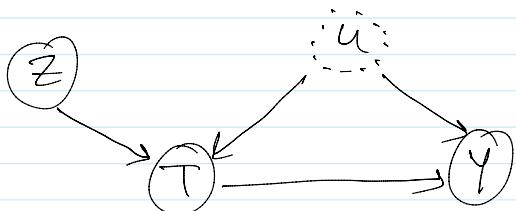
$$Y = \beta_0 + \beta_1 X + u$$

Z is used as an instrumental variable for x

$$\text{Cov}(Z, u) = 0 \quad \& \quad \text{Cov}(Z, x) \neq 0$$

x, u are uncorrelated then  $\beta_0$  &  $\beta_1$  are consistent

Instrument exogeneity means Z should have no partial effect on Y (after x and omitted variables have been controlled for) & Z should be uncorrelated with omitted variables



Graph used for reference of the assumptions.

4)  $P(X=1 | Z=g)$

$$P(X=1 | Z=1) = \frac{0.116 + 0.009}{0.116 + 0.009 + 0.334 + 0.051} = 0.245$$

0.116 + 0.009

+ 0.334 + 0.051

= 0.245

$$0.116 + 0.009 + 0.334 + 0.051$$

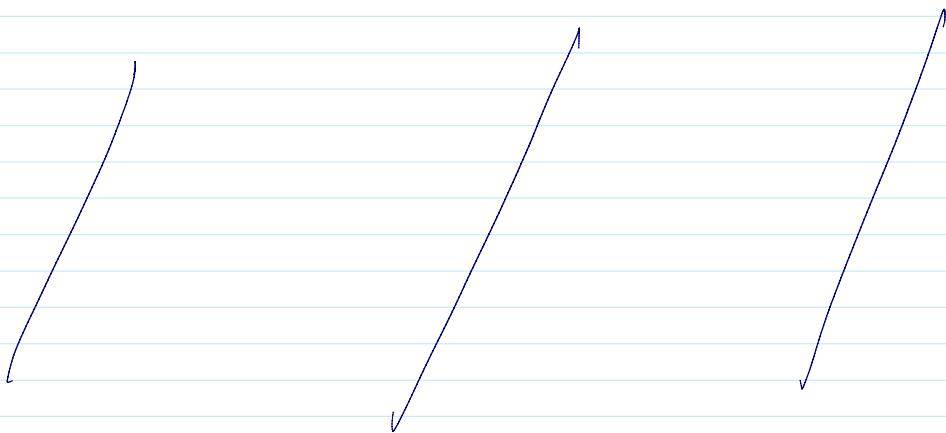
$$P(X=1 | Z=0) = \frac{0.274 + 0.101}{0.274 + 0.101 + 0.079 + 0.036} = 0.765$$

We multiply the corresponding values by  $\frac{1}{0.245}$  &  $\frac{1}{0.765}$

$$P(X=0 | Z=1) = 0.755 \quad (\because 1 - 0.245)$$

$$P(X=0 | Z=0) = 0.235 \quad (\because 1 - 0.765)$$

$$\begin{aligned} \therefore T &= E[Y(1) - Y(0)] \\ &= E\left[\frac{\mathbb{I}(X=1)Y}{e(w)}\right] - E\left[\frac{\mathbb{I}(X=0)Y}{1-e(w)}\right] \\ &= \left[ \frac{0.116 \times 1}{0.245} + \frac{0.274 \times 1}{0.765} + \frac{0.009 \times 0}{0.245} + \frac{0.101 \times 0}{0.765} \right] \\ &\quad - \left[ \frac{0.334 \times 1}{0.755} + \frac{0.079 \times 1}{0.235} + \frac{0.051 \times 0}{0.765} + \frac{0.031 \times 0}{0.235} \right] \\ &= 0.053 \end{aligned}$$



5) The propensity score is given by

$$f(x) = P(T=1 | X=x)$$

For treated individuals,  $w(x) = \frac{1}{\cdot}$ .

For treated individuals,  $\omega(u) = \frac{1}{p(u)}$

For control individuals,  $\omega(u) = \frac{1}{1-p(u)}$

In the presence of measured confounders, the causal effect is estimated by IPW  $\rightarrow$

$$\tau = \frac{1}{n_T} \sum_{i:T_i=1} \frac{Y_i}{p(x_i)} - \frac{1}{n_C} \sum_{i:T_i=0} \frac{Y_i}{1-p(x_i)}$$

$$= \frac{1}{n} \sum_i \frac{T_i Y_i}{p(x_i)} - \frac{1}{n} \sum_i \frac{(1-T_i) Y_i}{1-p(x_i)}$$

$p(x)$  is logistic regression based estimator of propensity score

The original quantity  $\tau = E[Y(1)] - E[Y(0)]$   
is unbiased, we have to show

$$E\left[\frac{YT}{p(x)}\right] = E[Y(1)] \quad \& \quad E\left[\frac{Y(1-T)}{1-p(x)}\right] = E[Y(0)]$$

$$\begin{aligned} \therefore E\left[\frac{YT}{p(x)}\right] &= E\left[E\left[\frac{YT}{p(x)} \mid X\right]\right] \\ &= E\left[E\left[\frac{Y(1)T}{p(x)} \mid X\right]\right] \\ &= E\left[\frac{E[Y(1)|X]E[T|X]}{p(x)}\right] \\ &= E\left[E[Y(1)|X]\right] = E[Y(1)] \end{aligned}$$

Similarly,

$$E\left[\frac{Y(1-T)}{1-p(x)}\right] = E[Y(0)]$$

$$(1/n_1) \sum Y | T=1 = b \left( 1 + \frac{p(u_1=0)p(u_1=1)}{m_1 - n_1 m_1 + 1 + m_1 - 1} \right)$$

$$6) a) E[Y_{x=1} | Z=1] = b \left( 1 + \frac{P(Y_1=0)P(Y_2=1)}{P(Y_1=0)P(Y_2=1) + P(Y_1=1)P(Y_2=0)} \right)$$

b)	$U_1$	$U_2$	$X_{U_1 U_2}$	$Z_{U_1 U_2}$	$Y_{U_1 U_2}$	$Z_{X=0}_{U_1 U_2}$	$Z_{X=1}_{U_1 U_2}$	$Y_{X=0}_{U_1 U_2}$	$Y_{X=1}_{U_1 U_2}$
	0	0	0	0	0	0	a	0	ab
	0	1	0	1	b	1	a+1	b	b(a+1)
	1	0	1	a	ba	0	a	0	ab
	1	1	1	a+1	b(a+1)	1	a+1	b	b(a+1)

7)  $\tau$  be the slope of total effect of  $X$  on  $Y$

$$\tau = E[Y|d_0(u+1)] - E[Y|d_0(u)]$$

for any evidence  $E=e$ ,

$$E[Y_{X=n}|E=e] = E[Y|E=e] + \tau(n - E[X|E=e])$$

The quantity can be calculated by best estimate of  $Y$  conditioned on evidence "e",  $E[Y|e]$  and adding to the change  $Y$  when  $X$  is shifted from current best estimate

$E[X|E=e]$  to hypothetical value  $n$ .

This is used in computing effect of treatment on treated

$$ETT = E[Y_1 - Y_0 | X=1]$$

evidence  $e = \{X=1\}$

$$\begin{aligned} ETT &= E[Y_1 | X=1] - E[Y_0 | X=1] \\ &= E[Y | X=1] - E[Y | X=1] + \tau(1 - E[X | X=1]) \\ &\quad - \tau(0 - E[X | X=1]) \\ &= \tau \end{aligned}$$

$\therefore$  The effect of treatment on treated is basically equal to the effect on treatment on entire population.

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2) a) For  $w = 0 \quad X = 0$

$$\frac{30+20}{200} = \frac{50}{200} = \frac{1}{4} = 25\%.$$

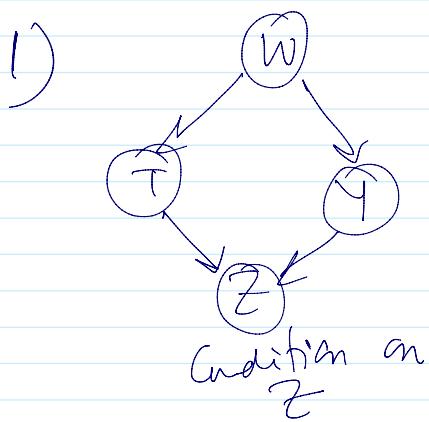
$w = 1 \quad X = 0$

$$\frac{30+20}{200} = \frac{1}{4} = 25\%.$$

So we can assume the treatment assignment is unconfounded.

b)  $P(w=1 | X=1) = \frac{75}{100} = 0.75$

$$P(w=1 | X=0) = \frac{30+20}{30+30+20+20} = 0.5$$



If we condition on a descendant of  $T$  that isn't a mediator, it could unblock a path from  $T$  to  $Y$  that was blocked by a collider. In this figure conditioning on  $Z$  induces non-causal association between  $T$  and  $Y$ , which biases the estimate of the causal effect.

So conditioning on  $Z$  or any descendant of  $Z$  in a path like this, will induce collider bias.

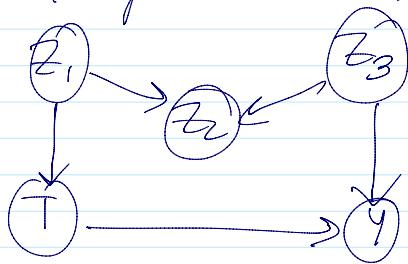
Selection bias  $\Rightarrow$  it is a distortion in measure of association due to sample selection that does not accurately reflect

selection bias  $\rightarrow$  SR is a distortion in measure of associations due to sample selection that does not accurately reflect the target population. SR occurs when investigators use improper procedures for selecting a sample population.

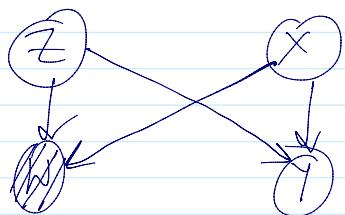
M-bias  $\rightarrow$  If we condition on pre treatment covariates

we can induce collider bias. In the following figure

if we condition on  $Z_2$  open a back door path for non causal association to flow. This is known as M-bias due to M-shape that this non causal association flows.



In the given graph, if we condition on  $W$ , there's



Collider bias

$$P(Y=y | do(X=x), W=w)$$

$$= \sum_z P(Y=y | X=x, W=w, Z=z)$$

$$\begin{aligned}
 2C) \quad & \tau = E[Y(1) - Y(0)] \\
 & = E\left[\frac{\mathbb{I}(W=1)Y}{e(w)}\right] - E\left[\frac{\mathbb{I}(W=0)Y}{e(w)}\right] \\
 & = 7.80 - 6.07 \\
 & = 1.7299
 \end{aligned}$$