

Irregular Assignment Mechanism
(Continued from lecture 3/3)

Example from last lecture: Trial – effect of vitamin A on infant mortality

Assignment Z	Trt received W	Outcome Y	# children	
0	0	0	74	
0	0	1	11514	.9936
1	0	0	34	
1	0	1	2385	.9859
1	1	0	12	
1	1	1	9663	.9988

$$- \text{ITT}_{z \text{ on } Y} = \frac{12048}{12094} - \frac{11514}{11588} = 0.0026 \text{ (compare } Z = 1 \text{ to } Z = 0, \text{ on } Y)$$

$$- \text{ITT}_{z \text{ on } w} = \frac{9675}{12094} - \frac{0}{11588} = 0.80 \text{ (compare } Z = 1 \text{ to } Z = 0, \text{ on } W)$$

$$- \text{Per protocol} = .9988 - .9936 = 0.0052 \text{ (compare } w = z = 1 \text{ to } w = z = 0, \text{ on } Y)$$

$$- \text{As treated} = .9988 - (13899/14007) = 0.0065 \text{ (compare } w = 1 \text{ to } w = 0)$$

If we are interested in the effect of W on Y, then there is a problem in that assignment of W is not randomized or even necessarily un-confounded.

The specific concern is that we don't know about the Z = 1, W = 0, and Z = 0, W = 0 groups, we don't know whether these two groups are the same or not.

Since we have no covariate to condition on, the best thing we could do is to control for compliance.

Approach: 2 steps

1. Define “compliance” in a way that we can condition on it
2. We estimate treatment effect of W on a subgroup with some assumptions.

Step 1, define compliance

Four subclasses	Z = 0	Z = 1	Notes
Complier	W = 0	W = 1	
Always taker	W = 1	W = 1	<i>Empty set in our example</i>
Never taker	W = 0	W = 0	<i>Also called non-compliers in our example</i>
Defier	W = 1	W = 0	<i>Generally assumed empty</i>

Compliance

- Property of the unit
- Defined for both Z's
- Not observed for all units

Step 2, now we can talk about causal inference for W on Y in compliance strata.

W is un-confounded given compliance strata, as shown:

$$P(W_i^{obs} = 1 | y(0,0), y(1,0), y(0,1), y(1,1), C_i) = P(W_i^{obs} = 1 | C_i)$$

If $C_i = \text{non-complier/never taker}$, LHS = RHS = 0

If $C_i = \text{complier}$, $W_i = Z_i$ and LHS = RHS by randomization of Z

If we could condition on C, then earlier work (matching/stratification) solves the problem.

Don't observe C for everyone. But can learn about causal effect on the subgroup of compliers (with some assumptions)

From last time,

$$ITT_{z \text{ on } y} = ITT_{z \text{ on } y, c} \times \pi_c + ITT_{z \text{ on } y, n} \times \pi_n$$

Add exclusion restrictions:

$$y_i(0,0) = y_i(1,0), \text{ for } c_i = n, \text{ which means that } ITT_{z \text{ on } y, n} = 0$$

Then we obtain:

$$ITT_{z \text{ on } y, c} = \text{treatment effect of W on Y for compliers (Z=W, for } C_i=c)$$

$$= \frac{ITT_{z \text{ on } y}}{\pi_c} = \frac{ITT_{z \text{ on } y}}{ITT_{z \text{ on } w}} = \frac{0.0026}{0.8} = 0.0032$$

= Compiler Average Causal Effect (CACE) or

Local Average Treatment Effect(LATE)

Variance for estimate: could be estimated using delta method and the fact that the estimate is derived as a function of sample proportions.

Instrumental Variable Approach

Previous example is an instance of the instrumental variable approach. We now describe the approach in more general terms.

Instrument: variable that randomly induces variation in the treatment of interest (e.g., Z = randomization to treatment group is an instrument in the Vitamin A study)

Key assumptions for IV approach (Gelman & Hill, Chap 10.5, 10.6)

- unconfoundedness of instrument
 - o trivially true in example because Z is randomized;
 - o it may require use of covariates
- non-zero association between instrument and treatment
 - o this can be checked using correlation of Z and W .

For
$$ITT_{y,c} = \frac{ITT_y}{ITT_w} \rightarrow \text{the effect of randomization on treatment}$$

If $ITT_w = 0$, it doesn't help

- monotonicity
 - o no defiers
- exclusion restriction
 - o no direct effect of instrument on the outcome

For future reference we now note that the CACE can be expressed as the ratio of “regression estimates”

$$CACE = ITT_{y,c} = \frac{ITT_y}{ITT_w}$$

$$\rightarrow \hat{ITT}_{y,c} = \frac{\hat{ITT}_y}{\hat{ITT}_w} = \frac{\bar{y}_{z=1} - \bar{y}_{z=0}}{\bar{w}_{z=1} - \bar{w}_{z=0}} = \frac{\hat{\beta}_{reg\ y\ on\ z}}{\hat{\beta}_{reg\ w\ on\ z}}$$

Regression view of the IV approach

Outcome Y }
 Instrument Z } all can be continuous
 Treatment T } (T is W we used in lecture before)

- We build two regression models: Y on T and T on Z

$$\left\{ \begin{array}{l} Y = \beta_0 + \beta_1 T + \varepsilon \\ \text{T may be correlated with error (which is why we can't estimate effect of T} \\ \text{using this regression)} \\ T = \gamma_0 + \gamma_1 Z + v \end{array} \right.$$

- Four assumptions of IV in this regression setup:

Non-zero association: $\gamma_1 \neq 0$

Monotonicity : linearity of relationship of T on Z

Unconfoundedness : Z is independent of v

(assuming no covariates X, but we can also accommodate X if present)

Exclusion restriction: Z is independent of Y_i and ε , given T_i

IV estimation approach

Regress T on Z $\Rightarrow \hat{\gamma}_0, \hat{\gamma}_1 \Rightarrow \hat{T}$

Regress Y on $\hat{T} \Rightarrow \hat{\beta}_1$

Standard error requires some adjustments

- Where does exclusion restriction come from? Start with more general regression

(no ER) $\left\{ \begin{array}{l} Y = \beta_0 + \beta_1 T + \beta_2 Z + \varepsilon \\ T = \gamma_0 + \gamma_1 Z + v \end{array} \right.$

$\longrightarrow Y = (\beta_0 + \beta_1 \gamma_0) + (\beta_2 + \beta_1 \gamma_1) Z + \varepsilon$, let $\beta_2 + \beta_1 \gamma_1 = \delta_1$

We want to obtain $\beta_1 = \frac{\delta_1 - \beta_2}{\gamma_1}$

Can estimate δ_1 and γ_1 , because Z is randomized

But no unbiased estimate for β_2 , since T maybe correlated with ε , so we need exclusion restriction to let $\beta_2 = 0$.