# Lecture 10: Instrumental variables III

#### **PPHA 34600**

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# From last time: applications of IV

## $Z_i$ is a valid instrument when the following are satisfied:

- **1** First stage:  $Cov(Z_i, D_i) \neq 0$
- **2** Exclusion restriction:  $Cov(Z_i, \varepsilon_i) = 0$

#### When we have these two conditions, we can...:

- Handle OVB
- Handle measurement error

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- All we need is a first stage...
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- But!  $Z_i$  is just generating variation in part of  $C_i$
- If this part affects  $Y_i$  differently than the non-moved bit,  $\hat{\tau} \neq \tau^{ATE}$

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- $Y_i(D_i, Z_i)$  is the outcome as a function of both treatment and the instrument
- $Y_i(D_i = 1, Z_i) Y_i(D_i = 0, Z_i)$ : Causal effect of treatment given your instrument
- $Y_i(D_i, Z_i = 1) Y_i(D_i, Z_i = 0)$ : Causal effect of your instrument given your treatment status

## In our intended causal chain, $Z_i \rightarrow D_i \rightarrow Y_i$ :

- We want notation to think about  $Z_i$  having a causal effect on  $D_i$ . Define:
  - $D_i(Z_i = 1)$  or just  $D_i(1)$  is treatment status when  $Z_i = 1$
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- Observed treatment status is just:

$$D_i = D_i(0) + (D_i(1) - D_i(0))Z_i = \alpha + \gamma_i Z_i + \nu_i$$

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(This should look familiar!)

- As before,  $\alpha = E[D_i(0)]$
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- $\rightarrow$  We can't observe both  $D_i(1)$  and  $D_i(0)$  (why?)
- $\rightarrow$  We can hope for the *average* causal effect of  $Z_i$  on  $D_i = E[\gamma_i]$

# With this framework, we need some (new) assumptions

We'll make four assumptions:

- **1 First stage:**  $E[D_i|Z_i=1] \neq E[D_i|Z_i=0]$  for some i
  - This is the same as before:  $Cov(D_i, Z_i) \neq 0$
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- **2** Independence:  $Y_i(D_i, Z_i), D_i(1), D_i(0) \perp Z_i$
- **3** Exclusion restriction:  $Y_i(Z_i = 1, D_i) = Y_i(Z_i = 0, D_i)$
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What used to just be the exclusion restriction,  $Cov(Z_i, \varepsilon_i) = 0$  is now: **(A) Independence:**  $Y_i(D_i, Z_i), D_i(1), D_i(0) \perp Z_i$ 

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$$E[Y_i|Z_i = 1] - E[Y_i|Z_i = 0]$$

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We can combine these two to express:

$$Y_i = Y_i(D_i = 0, Z_i) + (Y_i(D_i = 1, Z_i) - Y_i(D_i = 0, Z_i))D_i$$
  
=  $Y_i(0) + (Y_i(1) - Y_i(0))D_i$ 

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- **4** Monotonicity:  $D_i(Z_i = 1) D_i(Z_i = 0) \ge 0$  for all i

# Monotonicity

This new assumption says:

$$D_i(Z_i = 1) - D_i(Z_i = 0) \ge 0$$
 for all i

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- While Z<sub>i</sub> need not move everybody's treatment status...
- ... all affected units move in the same way
- Either  $D_i(Z_i = 1) \ge D_i(Z_i = 0)$  for all i
- Or  $D_i(Z_i = 1) \le D_i(Z_i = 0)$  for all i
- Moving from  $Z_i = 0$  to  $Z_i = 0$  doesn't move some units from  $D_i = 0$  to  $D_i = 1$  and others from  $D_i = 1$  to  $D_i = 0$

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- **4 Monotonicity:**  $D_i(Z_i = 1) D_i(Z_i = 0) \ge 0$  for all i

As always, we'd (ideally) estimate the following regression:

$$Y_i = \alpha + \tau D_i + \varepsilon_i$$

Since  $D_i$  is not randomly assigned, we also need an instrument,  $Z_i$  Recall that we can estimate  $\hat{\tau}^{IV}$  using two regressions:

$$\underbrace{D_i = \alpha + \gamma Z_i + \eta_i}_{\text{first stage}}$$

and

$$Y_i = \alpha + \theta Z_i + \nu_i$$
reduced form

Then

$$\hat{\tau}^{IV} = \frac{\hat{\theta}}{\hat{\gamma}} = \frac{E[Y_i|Z_i = 1] - E[Y_i|Z_i = 0]}{E[D_i|Z_i = 1] - E[D_i|Z_i = 0]}$$

Let's decompose 
$$\hat{\tau}^{IV} = \frac{E[Y_i|Z_i=1] - E[Y_i|Z_i=0]}{E[D_i|Z_i=1] - E[D_i|Z_i=0]}$$
:
$$E[Y_i|Z_i=1] = \underbrace{E[Y_i(0) + (Y_i(1) - Y_i(0))D_i|Z_i=1]}_{\text{exclusion restriction}}$$

$$= \underbrace{E[Y_i(0) + (Y_i(1) - Y_i(0))D_i(Z_i=1)]}_{\text{independence}}$$

and

$$E[Y_i|Z_i = 0] = \underbrace{E[Y_i(0) + (Y_i(1) - Y_i(0))D_i|Z_i = 0]}_{\text{exclusion restriction}}$$

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Taken together, these two yield

$$E[Y_i|Z_i = 1] - E[Y_i|Z_i = 0] = E[(Y_i(1) - Y_i(0))(D_i(1) - D_i(0))]$$

$$= \underbrace{E[Y_i(1) - Y_i(0)|D_i(1) > D_i(0)]Pr(D_i(1) > D_i(0))}_{\text{monotonicity}}$$

where  $E[Y_i(1) - Y_i(0)]$  is some kind of treatment effect

 $|D_i(1) > D_i(0)|$ : for compliers only

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where  $E[Y_i(1) - Y_i(0)]$  is some kind of treatment effect  $|D_i(1) > D_i(0)|$ : for compliers only  $Pr(D_i(1) > D_i(0))$ : share of compliers in the population. By the same logic:

$$E[D_i|Z_i=1] - E[D_i|Z_i=0] = \underbrace{E[D_i(1) - D_i(0)]}_{ ext{independence}}$$

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By the same logic:

$$\begin{split} E[D_{i}|Z_{i} = 1] - E[D_{i}|Z_{i} = 0] &= \underbrace{E[D_{i}(1) - D_{i}(0)]}_{\text{independence}} \\ &= \underbrace{Pr(D_{i}(1) > D_{i}(0))}_{\text{monotonicity}} \\ \hat{\tau}^{IV} = \frac{E[Y_{i}|Z_{i} = 1] - E[Y_{i}|Z_{i} = 0]}{E[D_{i}|Z_{i} = 1] - E[D_{i}|Z_{i} = 0]} = E[Y_{i}(1) - Y_{i}(0)|D_{i}(1) > D_{i}(0)] \end{split}$$

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# What happens without monotonicity?

Monotonicity,  $D_i(Z_i = 1) - D_i(Z_i = 0) \ge 0$  for all i, is a new assumption

- Without it, we have  $D_i(Z_i = 1) D_i(Z_i = 0) < 0$  for some i
- This breaks our ability to estimate  $\tau^{LATE}$  using  $\hat{\tau}^{IV}$

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- We had:

$$E[Y_i|Z_i=1]-E[Y_i|Z_i=0]=E[(Y_i(1)-Y_i(0))(D_i(1)-D_i(0))]$$

But without monotonicity:

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- → We can't deal with this
  - $\tau^i$  could be > 0 for all i, but we could mistakenly estimate 0 effect
- $\rightarrow$  We would have **defiers** ( $\nearrow$ 2)

$$\hat{\tau}^{IV} = E[Y_i(1) - Y_i(0)|D_i(1) > D_i(0)]$$

What is this "conditional on  $D_i(1) > D_i(0)$ " beast?

- $\hat{\tau}^{IV}$  estimates the (L)ATE, conditional on  $D_i(1) > D_i(0)$
- $D_i(1) > D_i(0)$  means  $Z_i$  moves  $D_i$

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- We can divide the world into three groups:
  - $\mathbf{0}$   $D_i(1) > D_i(0)$ : Compliers
  - **2**  $D_i(1) = D_i(0) = 1$ : Always-takers
  - **3**  $D_i(1) = D_i(0) = 0$ : Never-takers

$$\hat{\tau}^{IV} = E[Y_i(1) - Y_i(0)|D_i(1) > D_i(0)]$$

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- $\rightarrow$  Note that  $Z_i$  doesn't affect  $D_i$  for never-takers or always-takers
- → The instrument is useless for them
- → We can't learn about their treatment effects!
- → (They essentially have no first stage)
- → We can estimate LATEs for compliers only

#### Non-compliance throwback

#### We looked at several scenarios of non-compliance:

• If only T can non-comply, we can show:

$$\hat{\tau}^{IV} = \frac{E[Y_i|Z_i=1] - E[Y_i|Z_i=0]}{E[D_i|Z_i=1]} = E[Y_i(1) - Y_i(0)|D_i=1] = \tau^{LATE}$$

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#### Non-compliance throwback

If both T and C can non-comply:

$$\hat{\tau}^{IV} = \frac{E[Y_i|Z_i = 1] - E[Y_i|Z_i = 0]}{E[D_i|Z_i = 1] - E[D_i|Z_i = 0]}$$
$$= E[Y_i(1) - Y_i(0)|D_i(1) > D_i(0)] = \tau^{LATE}$$

#### Why does this work?

- We have an as-good-as-random estimate,  $E[Y_i|Z_i=1]-E[Y_i|Z_i=0]$
- We need to scale this by the complier proportion

### Counting compliers

The fraction of compliers is just:

$$\pi^{C} = Pr(D_{i}(1) > D_{i}(0)) = E[D_{i}(1) - D_{i}(0)]$$

$$= E[D_{i}(1)] - E[D_{i}(0)]$$

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We can also count the fraction of the treatment group which complies:

$$\begin{aligned} Pr(D_i(1) > D_i(0)|D_i = 1) &= \frac{Pr(D_i = 1|D_i(1)) > D_i(0))Pr(D_i(1) > D_i(0)}{Pr(D_i = 1)} \\ &= \frac{Pr(Z_i = 1)(E[D_i|Z_i = 1] - E[D_i|Z_i = 0])}{Pr(D_i = 1)} \end{aligned}$$

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- We can just count them
- But we can actually learn something more about them!

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$$\frac{Pr(\textit{Male}_i = 1 | D_i(1) > D_i(0)}{Pr(\textit{Male}_i = 1)}$$

$$= \frac{Pr(D_i(1) > D_i(0)|Male_i = 1)}{Pr(D_i(1) > D_i(0))}$$

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Let's ask: are compliers more likely to be men?

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ightarrow This is just the first stage for men divided by the overall first stage!

#### Heterogeneous $\tau_i$ makes things interesting:

- With homogenous  $\tau_i$ , all instruments should yield the same  $\tau^{LATE}$
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With multiple instruments, we get multiple estimates of

$$E[Y_i(1) - Y_i(0)|D_i(1) > D_i(0)]$$

Each instrument  $Z_i^1,...Z_i^K$  will have its own compliers where  $D_i(1) > D_i(0)$ 

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And the 2SLS estimator will be:

$$\hat{\tau}^{2SLS} = \frac{Cov(Y_i, \hat{D}_i)}{Cov(D_i, \hat{D}_i)} = \frac{\pi_1 Cov(Y_i, Z_i^1)}{Cov(D_i, \hat{D}_i)} + \frac{\pi_2 Cov(Y_i, Z_i^2)}{Cov(D_i, \hat{D}_i)}$$

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 $\rightarrow$  This is just a weighted average of each instrument's  $\hat{\tau}^{IV}$ 

### Non-binary treatments

#### What happens with non-binary treatment?

• Binary treatment: there is only  $Y_i(1)$  and  $Y_i(0)$ 

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- Binary treatment: there is only  $Y_i(1)$  and  $Y_i(0)$
- Non-binary treatment: define  $S_i \in \{0, 1, ..., \bar{S}\}$
- This has many potential outcomes  $Y_i(0), Y_i(1), ..., Y_i(\bar{S})$
- And many causal effects:  $Y_i(1) Y_i(0), Y_i(2) Y_i(1)...$

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- This has many potential outcomes  $Y_i(0), Y_i(1), ..., Y_i(\bar{S})$
- And many causal effects:  $Y_i(1) Y_i(0), Y_i(2) Y_i(1)...$
- In a linear model, these are all the same
- But that's unrealistic
- → 2SLS to the rescue!

### The average causal response

#### We can get a weighted average response with some assumptions:

- Independence + exclusion:  $\{Y_i(0), Y_i(1), ... Y_i(\bar{S})\} \perp Z_i$
- First stage:  $E[S_i(1) S_i(0)] \neq 0$
- Monotonicity:  $S_i(1) S_i(0) \ge 0$  for all i (or vice versa)

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Then:

$$\hat{\tau}^{IV} = \frac{E[Y_i|Z_i=1] - E[Y_i|Z_i=0]}{E[S_i|Z_i=1] - E[S_i|Z_i=0]}$$

$$= \sum_{s=1}^{S} \omega_{s} E[Y_{i}(s) - Y_{i}(s-1) | S_{i}(1) \geq s \geq S_{i}(0)]$$

where

$$\omega_s = \frac{Pr(S_i(1) \ge s > S_i(0)}{\sum_{j=1}^{\bar{S}} Pr(S_i(1) \ge j > S_i(0))}$$

### The average causal response

$$\hat{\tau}^{IV} = \sum_{s=1}^{\bar{S}} \omega_s E[Y_i(s) - Y_i(s-1)|S_i(1) \ge s \ge S_i(0)]$$

- $\rightarrow \hat{\tau}^{IV}$  gives a weighted average of the unit causal response
- $\rightarrow$  The unit causal response,  $E[Y_i(s) Y_i(s-1)|S_i(1) \ge s \ge S_i(0)]$  is the average difference in potential outcomes for compliers at  $S_i = s$
- $\rightarrow$  The size of the compliance group is  $Pr(S_i(1) \ge s > S_i(0))$

## What do we get from the IV?

#### We've talked through several cases

- Constant  $\tau$ :
  - $\hat{\tau}^{IV} = \tau^{ATE}$
- Perfect compliance:
  - $\hat{\tau}^{IV} = \tau^{ATE}$
- Heterogeneous treatment effects, one IV:
  - $\hat{\tau}^{IV} = \tau^{LATE}$
- Heterogeneous treatment effects, multiple IVs:
  - $\hat{\tau}^{IV} = \frac{1}{K} \sum_{k} \omega_k \tau_k^{LATE}$
- Multiple values of treatment:
  - $\hat{\tau}^{IV} = \sum_{s=1}^{\bar{S}} \omega_s E[Y_i(s) Y_i(s-1)|S_i(1) \ge s \ge S_i(0)]$

### Taking stock of IV

#### We've come a long way from RCTs:

- Took a brief detour through the thicket of SOO
- Started our discussion of SOU

### Taking stock of IV

#### We've come a long way from RCTs:

- Took a brief detour through the thicket of SOO
- Started our discussion of SOU
- → **IV** is our first SOU design
  - IV helps us do causal inference with non-random treatment
  - We just need some random leverage over treatment

### Taking stock of IV

#### Under the right assumptions, we can use IV for...

- Eliminating bias due to measurement error
- Eliminating bias due to omitted variables
- Eliminating bias due to simultaneity
- Translating from ITT to LATE
- Estimating (L)ATEs

The trick is satisfying the exclusion restriction!

#### Recap

#### TL;DR:

- 1 Instrumental variables are very powerful
- We can use them to handle non-compliance
- 3 More generally, the IV estimates LATE (not ATE) with heterogeneity

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