Gov 2002 - Causal Inference II: Instrumental Variables

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- But if you can identify some exogenous sources of variation that drive the treatment, even if the treatment was not randomly assigned, you may be able to make headway.

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- But if you can identify some exogenous sources of variation that drive the treatment, even if the treatment was not randomly assigned, you may be able to make headway.
- The basic idea behind instrumental variables is that we have a treatment with unmeasured confounding, but that we have another variable, called the instrument, that affects the treatment, but not the outcome, and thus give us that exogenous variation.

U $Z \to A \to Y$



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 - no common causes of the instrument and the outcome
 - no direct or indirect effect of the instrument on the outcome not through the treatment.

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- Exclusion restriction
 - no common causes of the instrument and the outcome
 - no direct or indirect effect of the instrument on the outcome not through the treatment.
- First-stage relationship: Z affects A

An IV is only as good as its assumptions



 Finding a believable instrument is incredibly difficult and some people never believe any IV setups.

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An IV is only as good as its assumptions



- Finding a believable instrument is incredibly difficult and some people never believe any IV setups.
- We will see that even if all of the untestable assumptions are met, the IV approach estimates a "local" ATE. That is, local to this particular case/instrument.

 Angrist (1990): Draft lottery as an IV for military service (income as outcome)

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- Acharya, Blackwell, Sen (2014): cotton suitability as IV for proportion slave in 1860 (outcome is white attitudes today)

► Let's write down a causal model for *Y_i* with constant effects and an unmeasured confounder, *U_i*:

$$Y_i(a, u) = lpha + au a + \gamma u + \eta_i$$

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Let's write down a causal model for Y_i with constant effects and an unmeasured confounder, U_i:

$$Y_i(a, u) = \alpha + \tau a + \gamma u + \eta_i$$

If we connect this with a consistency assumption, we get the this regression form:

$$Y_i = \alpha + \tau A_i + \gamma U_i + \eta_i$$

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- Here we assume that $E[A_i\eta_i] = 0$, so if we measured U_i , then we would be able to estimate τ .
- But cov(γU_i + η_i, A_i) ≠ 0 because U is a common cause of A and Y.

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$$\operatorname{cov}(\gamma U_i + \eta_i, Z_i) = 0$$

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 $\operatorname{cov}(Y_i, Z_i) = \operatorname{cov}(\alpha + \tau A_i + \gamma U_i + \eta_i, Z_i)$

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$$cov(Y_i, Z_i) = cov(\alpha + \tau A_i + \gamma U_i + \eta_i, Z_i)$$

= cov(\alpha, Z_i) + cov(\alpha A_i, Z_i) + cov(\gamma U_i + \eta_i, Z_i)

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= cov(\alpha, Z_i) + cov(\alpha A_i, Z_i) + cov(\gamma U_i + \eta_i, Z_i)
= 0 + \tau cov(A_i, Z_i) + 0

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$$Y_i = \alpha + \tau A_i + \gamma U_i + \eta_i$$

With this in hand, we can formulate an expression for the average treatment effect here:

$$\tau = \frac{\operatorname{Cov}(Y_i, Z_i)}{\operatorname{Cov}(A_i, Z_i)} = \frac{\operatorname{Cov}(Y_i, Z_i)/V[Z_i]}{\operatorname{Cov}(A_i, Z_i)/V[Z_i]}$$

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• Reduced form coefficient: $Cov(Y_i, Z_i)/V[Z_i]$

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- Reduced form coefficient: $Cov(Y_i, Z_i)/V[Z_i]$
- First stage coefficient: $Cov(A_i, Z_i)/V[Z_i]$

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- Reduced form coefficient: $Cov(Y_i, Z_i)/V[Z_i]$
- ▶ First stage coefficient: Cov(A_i, Z_i)/V[Z_i]
- What happens with a weak first stage?

With a binary instrument, there is a simple estimator based on this formulation called the Wald estimator. It is easy to show that:

$$\tau = \frac{\text{Cov}(Y_i, Z_i)}{\text{Cov}(A_i, Z_i)} = \frac{E[Y_i | Z_i = 1] - E[Y_i | Z_i = 0]}{E[A_i | Z_i = 1] - E[A_i | Z_i = 0]}$$

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Intuitively, the effects of Z_i on Y_i divided by the effect of Z_i on A_i

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No covariates up until now. What if we have a set of covariates X_i that we are also conditioning on?

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- No covariates up until now. What if we have a set of covariates X_i that we are also conditioning on?
- Let's start with linear models for both the outcome and the treatment:

$$Y_i = X'_i\beta + \tau A_i + \varepsilon_i$$
$$A_i = X'_i\alpha + \gamma Z_i + \nu_i$$

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▶ Now, we assume that X_i are **exogenous** along with Z_i:

$$E[Z_i\nu_i] = 0 \quad E[Z_i\varepsilon_i] = 0$$
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• ... but A_i is endogenous: $E[A_i \varepsilon_i] \neq 0$

We can plug the treatment equation into the outcome equation:

$$Y_{i} = X'_{i}\beta + \tau [X'_{i}\alpha + \gamma Z_{i} + \nu_{i}] + \varepsilon_{i}$$

= $X'_{i}\beta + \tau [X'_{i}\alpha + \gamma Z_{i}] + [\tau\nu_{i} + \varepsilon_{i}]$
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- Red value in the brackets is the population fitted value of the treatment, E[A_i|X_i, Z_i]
- Because Z_i and X_i are uncorrelated with ν_i and ε_i, then this fitted value is also independent of ε^{*}_i.
- Thus, the population regression coefficient of a Y_i on $[X'_i\alpha + \gamma Z_i]$ is the average treatment effect, τ .

In practice, we estimate the first stage from a sample and calculate OLS fitted values:

$$\hat{A}_i = X_i'\hat{\alpha} + \hat{\gamma}Z_i.$$

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$$\hat{A}_i = X_i'\hat{\alpha} + \hat{\gamma}Z_i.$$

Here, â and γ are estimates from OLS. Then, we estimate a regression of Y_i on X_i and Â_i. We plug this into our equation for Y_i and note that the error for A_i is now a residual:

$$Y_i = X'_i eta + au \hat{A}_i + [arepsilon_i + au (A_i - \hat{A}_i)]$$

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• Key question: is \hat{A}_i uncorrelated with the error?

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- Key question: is \hat{A}_i uncorrelated with the error?
- \hat{A}_i is just a function of X_i and Z_i so it is uncorrelated with ε_i .
- We also know that \hat{A}_i is uncorrelated with $(A_i \hat{A}_i)$?

Heuristic procedure:



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 - 1. Run regression of treatment on covariates and instrument

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2. Construct fitted values of treatment

Heuristic procedure:

- 1. Run regression of treatment on covariates and instrument
- 2. Construct fitted values of treatment
- 3. Run regression of outcome on covariates and fitted values

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Heuristic procedure:

- 1. Run regression of treatment on covariates and instrument
- 2. Construct fitted values of treatment
- 3. Run regression of outcome on covariates and fitted values
- Note that this isn't how we actually estimate 2SLS because the standard errors are all wrong.

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Heuristic procedure:

- 1. Run regression of treatment on covariates and instrument
- 2. Construct fitted values of treatment
- 3. Run regression of outcome on covariates and fitted values
- Note that this isn't how we actually estimate 2SLS because the standard errors are all wrong.
- Computer wants to calculate the standard errors based on ε^{*}_i, but what we really want is the standard errors based on ε_i.

Nunn & Wantchekon IV example

	Trust of relatives (1)	Trust of neighbors (2)	Trust of local council (3)	Intragroup trust (4)	Intergroup trust (5)
Second stage: Dependent variable	is an individual's	trust			
ln (1+exports/area)	-0.190*** (0.067)	-0.245*** (0.070)	-0.221*** (0.060)	-0.251*** (0.088)	-0.174** (0.080)
Hausman test (<i>p</i> -value) R^2	0.88 0.13	0.53 0.16	0.09 0.20	0.44 0.15	0.41 0.12
First stage: Dependent variable is	ln (1+exports/a	rea)			
Historical distance of ethnic group from coast	-0.0014*** (0.0003)	-0.0014*** (0.0003)	-0.0014*** (0.0003)	-0.0014*** (0.0003)	-0.0014*** (0.0003)
Colonial population density Ethnicity-level colonial controls Individual controls District controls Country fixed effects	Yes Yes Yes Yes Yes	Yes Yes Yes Yes Yes	Yes Yes Yes Yes Yes	Yes Yes Yes Yes Yes	Yes Yes Yes Yes Yes
Number of observations Number of clusters F-stat of excl. instrument R^2	16,709 147 / 1,187 26.9 0.81	16,679 147 / 1,187 26.8 0.81	15,905 146 / 1,194 27.4 0.81	16,636 147 / 1,186 27.1 0.81	16,473 147 / 1,184 27.0 0.81

TABLE 5—IV ESTIMATES OF THE EFFECT OF THE SLAVE TRADE ON TRUST

Notes: The table reports IV estimates. The top panel reports the second-stage estimates, and the bottom panel reports first-stage estimates. Standard errors are adjusted for two-way clustering at the ethnicity and district levels. The individual controls, district controls, ethnicity-level colonial controls, and colonial population density measures are described in Table 3. The null hypothesis of the Hausman test is that the OLS estimates are consistent.

***Significant at the 1 percent level.

**Significant at the 5 percent level.

*Significant at the 10 percent level.

General 2SLS

To save on notation, we'll roll all the variables in the structural model in one vector, X_i, of size k, some of which may be endogenous.

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$$Y_i = X'_i\beta + \varepsilon_i$$

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 Z_i will be a vector of *l* exogenous variables that includes any exogenous variables in X_i plus any instruments. Key assumption:

$$E[Z_i\varepsilon_i]=0$$

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Useful quantities:

 $\Pi = (E[Z_i Z_i'])^{-1} E[Z_i X_i']$ (projection matrix) $V_i = \Pi' Z_i$ (fitted values)

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- Collect X_i into a $n \times k$ matrix $X = (X'_1, \ldots, X'_n)$
- Collect Z_i into a $n \times I$ matrix $Z = (Z'_1, \ldots, Z'_n)$

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- Take the population formula for the parameters:

$$\beta = (E[Z_i X_i'](E[Z_i Z_i'])^{-1}E[Z_i X_i'])^{-1}E[Z_i X_i'](E[Z_i Z_i'])^{-1}E[Z_i Y_i]$$

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And plug in the sample values (the n cancels out):

$$\hat{\beta} = [(X'Z)(Z'Z)^{-1}(Z'X)]^{-1}(Z'X)(Z'Z)^{-1}(Z'Y)$$

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How to estimate the parameters

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This is how R/Stata estimates the 2SLS parameters

Let V = Z(Z'Z)⁻¹Z'X be the matrix of fitted values for X, then we have

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$$\hat{\beta} = (V'V)^{-1}V'(X\beta + \varepsilon)$$

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• We can insert the true model for Y:

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• Using the matrix party trick and that V'X = V'V, we have

$$\hat{\beta} = (V'V)^{-1}V'X\beta + (V'V)^{-1}V'\varepsilon$$
$$= \beta + \left[n^{-1}\sum_{i}V_{i}V'_{i}\right]^{-1}n^{-1}\sum_{i}V_{i}\varepsilon_{i}$$

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• Consistent because $n^{-1}\sum_i V_i \varepsilon_i \xrightarrow{p} E[V_i \varepsilon_i] = 0$.

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$$\sqrt{n}(\hat{\beta}-\beta) = \left(n^{-1}\sum_{i}V_{i}V_{i}'\right)^{-1}\left(n^{-1/2}\sum_{i}V_{i}\varepsilon_{i}\right)$$

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By the CLT, n^{-1/2} ∑_i V_iε_i converges in distribution to N(0, B), where B = E[V'_iε'_iε_iV_i].

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$$n^{-1}\sum_i V_i V'_i \xrightarrow{p} E[V'_i V_i]$$
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- By the LLN, $n^{-1}\sum_i V_i V'_i \xrightarrow{p} E[V'_i V_i]$.
- Thus, we have that $\sqrt{n}(\hat{\beta} \beta)$ has asymptotic variance:

$$(E[V_i'V_i])^{-1}E[V_i'\varepsilon_i'\varepsilon_iV_i](E[V_i'V_i])^{-1}$$

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$$(E[V_i'V_i])^{-1}E[V_i'\varepsilon_i'\varepsilon_iV_i](E[V_i'V_i])^{-1}$$

Replace with the sample quantities to get estimates:

$$\widehat{\operatorname{var}}(\hat{\beta}) = (V'V)^{-1} \Big(\sum_{i} \hat{u}_i^2 V_i V_i'\Big) (V'V)^{-1}$$

where $\hat{u}_i = Y_i - X'_i \hat{\beta}$

What if we have more instruments than endogenous variables?

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What if we have more instruments than endogenous variables?

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Is it plausible to find more than one instrument?

Sargan test, Hansen test, J-test, etc.

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- Basic idea: under null that all instruments are good, running it with different subset of the instruments should only differ due to sampling noise.

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- If we reject the null hypothesis in these overidentification tests, then it means that the exclusion restrcitions for our instruments are probably incorrect. Note that it won't tell us which of them are incorrect, just that at least one is.

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- These overidentification tests depend heavily on the constant effects assumption
- Once we move away from constant effects, we no longer can generally pool multiple instruments together in this way.

Reading

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Reading









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The basic idea behind instrumental variable approaches is that we do not have ignorability for A_i, but we do have a variable, Z_i, that affects A_i, but only affects the outcome through A_i.

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1. Randomization



- 1. Randomization
- 2. Exclusion Restriction

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- 1. Randomization
- 2. Exclusion Restriction
- 3. First-stage relationship

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4. Monotonicity

Need the instrument to be randomized:

 $[\{Y_i(a,z), \forall a,z\}, A_i(1), A_i(0)] \perp \mathbb{Z}_i$

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- We can weaken this to conditional ignorability
- But why believe conditional ignorability for the instrument but not the treatment?
- Best instruments are truly randomized.
- Identifies the intent-to-treat (ITT) effect:

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

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Exclusion Restriction

The instrument has no direct effect on the outcome, once we fix the value of the treatment.

$$Y_i(a, 1) = Y_i(a, 0)$$
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NOT A TESTABLE ASSUMPTION

The linear model with heterogeneous effects

Rewriting the usual consistency assumption gives us a linear model with heterogeneous effects (we have seen this before in randomized experiments):

$$Y_i = Y_i(0) + (Y_i(1) - Y_i(0))A_i$$

= $\alpha_0 + \tau_i A_i + \eta_i$

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• Here, we have $\alpha_0 = E[Y_i(0)]$ and $\tau_i = Y_i(1) - Y_i(0)$.

First Stage

This next assumption is a little mundane, but turns out to be very important: the instrument must have an effect on the treatment.

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 Otherwise, what would we be doing? The instrument wouldn't affect anything.

Monotonicity

Lastly, we need to make another assumption about the relationship between the instrument and the treatment.

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- Monotonicity says that the presence of the instrument never dissuades someone from taking the treatment:

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Note if this holds in the opposite direction A_i(1) − A_i(0) ≤ 0, we can always rescale A_i to make the assumption hold.

This is sometimes called "no defiers". It turns out that with a binary treatment and a binary instrument, we can group units into four categories:

Name	$A_i(1)$	$A_{i}(0)$
Always Takers	1	1
Never Takers	0	0
Compliers	1	0
Defiers	0	1

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- These compliance groups are sometimes called "principal strata."
- The monotonicity assumption remove the possibility of there being defiers in the population.
- Anyone with A_i = 1 when Z_i = 0 must be an always-taker and anyone with A_i = 0 when Z_i = 1 must be a never-taker.

 Under these four assumptions, the Wald estimator is equal what we call Local average treatment effect (LATE) or the complier average treatment effect (CATE).

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$$\frac{E[Y_i|Z_i=1] - E[Y_i|Z_i=0]}{E[A_i|Z_i=1] - E[A_i|Z_i=0]} = E[Y_i(1) - Y_i(0)|A_i(1) > A_i(0)]$$

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 This fact was a massive intellectual jump in our understanding of IV.

Under the exclusion restriction and randomization,

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

= $E[Y_i(0) + (Y_i(1) - Y_i(0))A_i(1)]$ (randomization

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$$\begin{split} E[Y_i|Z_i = 1] &= E[Y_i(0) + (Y_i(1) - Y_i(0))A_i|Z_i = 1] \\ &= E[Y_i(0) + (Y_i(1) - Y_i(0))A_i(1)] \quad (\text{randomization}) \end{split}$$

• The same applies to when $Z_i = 0$, so we have

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 $E[(Y_i(1) - Y_i(0))(A_i(1) - A_i(0))]$ = $E[(Y_i(1) - Y_i(0))(1)|A_i(1) > A_i(0)] \Pr[A_i(1) > A_i(0)]$ + $E[(Y_i(1) - Y_i(0))(-1)|A_i(1) < A_i(0)] \Pr[A_i(1) < A_i(0)]$

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 $E[(Y_i(1) - Y_i(0))(A_i(1) - A_i(0))]$ =E[(Y_i(1) - Y_i(0))(1)|A_i(1) > A_i(0)] Pr[A_i(1) > A_i(0)] +E[(Y_i(1) - Y_i(0))(-1)|A_i(1) < A_i(0)] Pr[A_i(1) < A_i(0)] =E[Y_i(1) - Y_i(0)|A_i(1) > A_i(0)] Pr[A_i(1) > A_i(0)]

Under the exclusion restriction and randomization,

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

= $E[Y_i(0) + (Y_i(1) - Y_i(0))A_i(1)]$ (randomization

• The same applies to when $Z_i = 0$, so we have

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

▶ Thus, \$E[Y_i |Z_i = 1] - E[Y_i |Z_i = 0] = \$

 $E[(Y_i(1) - Y_i(0))(A_i(1) - A_i(0))]$ =E[(Y_i(1) - Y_i(0))(1)|A_i(1) > A_i(0)] Pr[A_i(1) > A_i(0)] +E[(Y_i(1) - Y_i(0))(-1)|A_i(1) < A_i(0)] Pr[A_i(1) < A_i(0)] =E[Y_i(1) - Y_i(0)|A_i(1) > A_i(0)] Pr[A_i(1) > A_i(0)]

► The third equality comes from monotonicity: with this assumption, A_i(1) < A_i(0) never occurs.

Proof (continued)

 $E[Y_i|Z_i = 1] - E[Y_i|Z_i = 0] = E[Y_i(1) - Y_i(0)|A_i(1) > A_i(0)] \Pr[A_i(1) > A_i(0)]$

- We can use the same argument for the denominator:

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

= Pr[A_i(1) > A_i(0)]

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- Dividing these two expressions through gives the LATE.

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Reading







 Once we allow for heterogeneous effects, all we can estimate with IV is the effect of treatment among compliers.

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- Once we allow for heterogeneous effects, all we can estimate with IV is the effect of treatment among compliers.
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- Without further assumptions, this estimand is not equal to overall treatment effect or the treatment effect on the treated.

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- ► This is a unknown subset of the data. Among treated units with Z_i = 1, we cannot distinguish them from the always-takers and similarly for the control units with Z_i = 0.
- Without further assumptions, this estimand is not equal to overall treatment effect or the treatment effect on the treated.
- Furthermore, since the complier group depends on the instrument, an IV estimate with one instrument will generally estimate a different quantity than an IV estimate of the same effect with a different instrument.
- 2SLS "cheats" by assuming that the effect is constant, so it is the same for compliers and non-compliers.
Will the LATE ever be equal to a usual causal quantity?

- ▶ Will the LATE ever be equal to a usual causal quantity?
- When non-compliance is **one-sided**, then the LATE is equal to the ATT.

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Think of a randomized experiment:

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- One-sided noncompliance: only those assigned to treatment (control) can actually take the treatment (control). Or

$$\Pr[A_i = 1 | Z_i = 0] = 0$$

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$$\Pr[A_i=1|Z_i=0]=0$$

Maybe this is because only those treated actually get pills or only they are invited to the job training location.

▶ With this assumption, we know that there are no "always-takers" and since there are no defiers, anyone treated (Z_i = 1) that takes the treatment (A_i = 1) is a complier.

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

 $= E[Y_i(1) - Y_i(0)|A_i = 1, Z_i = 1] \Pr[A_i = 1|Z_i = 1]$

(law of iterated expectations + binary treatment)

- ▶ With this assumption, we know that there are no "always-takers" and since there are no defiers, anyone treated (Z_i = 1) that takes the treatment (A_i = 1) is a complier.
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Noting that Pr[A_i = 1|Z_i = 0] = 0, then the Wald estimator is just the ATT:

$$\frac{E[Y_i|Z_i=1] - E[Y_i|Z_i=0]}{\Pr[A_i=1|Z_i=1]} = E[Y_i(1) - Y_i(0)|A_i=1]$$

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Thus, under the additional assumption of one-sided compliance, we can estimate the ATT using the usual IV approach. ▶ Noting that Pr[A_i = 1|Z_i = 0] = 0, then the Wald estimator is just the ATT:

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- Thus, under the additional assumption of one-sided compliance, we can estimate the ATT using the usual IV approach.
- The ATT is a combination of the LATE and the ATE for the always-takers. If we remove the possibility of the always takers, then anyone who actually takes the treatment is a complier.

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- Thus, under the additional assumption of one-sided compliance, we can estimate the ATT using the usual IV approach.
- The ATT is a combination of the LATE and the ATE for the always-takers. If we remove the possibility of the always takers, then anyone who actually takes the treatment is a complier.
- It's also easy to see that if we switch the direction of one-sided compliance, then we can esimate the average treatment effect for the controls.

The exclusion restriction cannot be tested directly, but it can be falsified.

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- The exclusion restriction cannot be tested directly, but it can be falsified.
- ► Under the exclusion restriction, Z_i only has an effect on Y_i because it has an effect on A_i.

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- ► Under the exclusion restriction, Z_i only has an effect on Y_i because it has an effect on A_i.
- ► Falsification test Test the reduced form effect of Z_i on Y_i in situations where it is impossible or extremely unlikely that Z_i could affect A_i.

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- Because Z_i can't affect A_i, then the exclusion restriction implies that this falsification test should have 0 effect. If we find an effect, instrument is suspicious.
- Nunn & Wantchekon (2011): use distance to coast as an instrument for Africans, use distance to the coast in an Asian sample as falsification test.

Nunn & Wantchekon falsification test

VOL. 101 NO. 7 NUNN AND WANTCHEKON: THE ORIGINS OF MISTRUST IN AFRICA 3243

	Trust of local government council			
	Afrobarometer sample		Asiabarometer sample	
	(1)	(2)	(3)	(4)
Distance from the coast	0.00039*** (0.00009)	0.00031*** (0.00008)	-0.00001 (0.00010)	0.00001 (0.00009)
Country fixed effects Individual controls	Yes No	Yes Yes	Yes No	Yes Yes
Number of observations Number of clusters R^2	19,913 185 0.16	19,913 185 0.18	5,409 62 0.19	5,409 62 0.22

TABLE 7—REDUCED FORM RELATIONSHIP BETWEEN THE DISTANCE FROM THE COAST AND TRUST WITHIN AFRICA AND ASIA

Notes: The table reports OLS estimates. The unit of observation is an individual. The dependent variable in the Asiabarometer sample is the respondent's answer to the question: "How much do you trust your local government?" The categories for the answers are the same in the Asiabarometer as in the Afrobarometer. Standard errors are clustered at the ethnicity level in the Afrobarometer regressions and at the location (city) level in the Asiabarometer and the WVS samples. The individual controls are for age, age squared, a gender indicator, education fixed effects, and religion fixed effects.

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- ***Significant at the 1 percent level.
- **Significant at the 5 percent level.
- *Significant at the 10 percent level.

Size, characteristics of the compliers

While we cannot identify who is a complier and who is not a complier in general, we can estimate the size of the complier group:

$$\Pr[A_i(1) > A_i(0)] = E[A_i(1) - A_i(0)] = E[A_i | Z_i = 1] - E[A_i | Z_i = 0]$$

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 Angrist and Pischke describe ways to calculate the difference between the compliers and overall population in terms of binary covariates.

Size, characteristics of the compliers

While we cannot identify who is a complier and who is not a complier in general, we can estimate the size of the complier group:

 $\Pr[A_i(1) > A_i(0)] = E[A_i(1) - A_i(0)] = E[A_i | Z_i = 1] - E[A_i | Z_i = 0]$

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- Angrist and Pischke describe ways to calculate the difference between the compliers and overall population in terms of binary covariates.
- Abadie (2003) shows how to calculate the mean of any covariate in the complier group.

Multiple instruments

 Since each instrument implies a different complier group, each instrument estimates a causal effect for a different subset of the population.

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Multiple instruments

- Since each instrument implies a different complier group, each instrument estimates a causal effect for a different subset of the population.
- Thus, if we had two instrument, then there would be two different LATEs, ρ₁ and ρ₂ for instruments Z_{1i} and Z_{2i}. We might try to use 2SLS to estimate an overall effect with these instruments with following first stage:

$$\hat{A}_i = \pi_1 Z_{1i} + \pi_2 Z_{2i}.$$

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2SLS as weighted average

In Angrist and Pischke, they show that the 2SLS estimator using these two instruments is a weighted sum of the two component LATEs:

$$\rho_{2SLS} = \psi \rho_1 + (1 - \psi) \rho_2,$$

where the weights are:

$$\psi = \frac{\pi_1 \operatorname{Cov}(A_i, Z_{1i})}{\pi_1 \operatorname{Cov}(A_i, Z_{1i}) + \pi_2 \operatorname{Cov}(A_i, Z_{2i})}$$

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Thus, the 2SLS estimate is a weighted average of causal effects for each instrument, where the weights are related to the strenght of prediction for each of the first stage effects of the instruments.

Covariates and heterogeneous effects

It might be the case that the above assumptions only hold conditional on some covariates, X_i. That is, instead of randomization, we might have conditional ignorability:

 $[\{Y_i(a,z), \forall a,z\}, A_i(1), A_i(0)] \perp \mathbb{Z}_i | X_i$

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We would also have exclusion conditional on the covariates:

$$\Pr[Y_i(a,0) = Y_i(a,1)|X_i] = 1$$
 for $a = 1,0$

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$$\Pr[Y_i(a,0) = Y_i(a,1)|X_i] = 1 \text{ for } a = 1,0$$

Under these assumptions, Angrist and Pischke show that if you fully saturate the first stage and the second stage in the covariates, then 2SLS estimates a weighted average of the covariates-specific LATEs (very similar to regression).
Covariates and heterogeneous effects

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- Under these assumptions, Angrist and Pischke show that if you fully saturate the first stage and the second stage in the covariates, then 2SLS estimates a weighted average of the covariates-specific LATEs (very similar to regression).
- Abadie (2003) shows how to estimate the overall LATE using a weighting approach based on a "propensity score" for the instrument.