Week 6: Instrumental Variables

PLSC 30600 - Causal Inference

Last two weeks

- Identification under conditional ignorability
 - \circ Treatment assignment is independent of the potential outcomes given observed confounders ${f X}$
 - "Selection-on-observables"
- "Selection-on-observables" isn't a testable assumption
 - \circ Relies on theory to decide which **X** to include.
 - DAGs can help here.
- Lots of estimation strategies
 - \circ Stratify with low-dimensional ${f X}$
 - IPTW to eliminate treatment-covariate relationship, regression to model the outcomecovariate relationship.
 - Matching to reduce model dependence.
 - Or consider more modern flexible modelling techniques for $E[Y_i(d)|X_i]$

This week

- Can we estimate a treatment effect when neither ignorability nor conditional ignorability hold for treatment?
 - Can we get rid of *unobserved* confounding?
- "Instrumental variables" designs are one way of dealing with this
- We can identify some average of treatment effects if...
 - There does exist an ignorable or conditionally ignorable instrument which...
 - ...has a monotonic effect on the treatment...
 - ...and has no effect on the outcome except through its effect on the treatment.
- What's the average? The "Local Average Treatment Effect"
 - Average effect among those who are moved to take treatment by the instrument

Instrumental Variables

Treatment non-compliance

- Often experiments suffer from treatment non-compliance
 - Participants randomized to receive a phone call don't pick up.
 - Participants randomized to wear surgical masks choose not to.
- New notation!
 - \circ Let Z_i denote whether i is assigned to receive a treatment.
 - \circ Let D_i denote the treatment actually *taken* by an individual.
- Can we just take the simple difference-in-means between $D_i=1$ and $D_i=0$?
 - No! Non-compliance affected by other factors which might also affect the outcome.
 - We're stuck with an observational design.

• Unless...

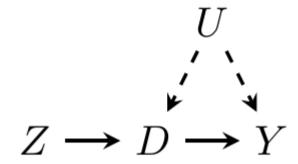
Intent-to-treat effect

- We can first just change the question instead of the effect of **treatment**, we can make our estimand the effect of being **assigned to treatment**.
- ullet Our estimator for the ITT is just the difference in means between the $Z_i=1$ and $Z_i=0$ arms

$$\hat{ au}_{ ext{ITT}} = \hat{E}[Y_i|Z_i=1] - \hat{E}[Y_i|Z_i=0]$$

- Identified under randomization of Z_i even if D_i is not randomized.
 - \circ But combines two effects: the actual effect of D_i and the effect of Z_i on D_i .

Instrumental variables



• Suppose though that we're interested in the *actual* effect of receiving treatment (the effect of D_i). What can we do?

Instrumental variables

• Start by writing down potential outcomes for D_i along with joint potential outcomes of Y_i in terms of Z_i and D_i

$$D_i(z) = D_i ext{ if } Z_i = z$$
 $Y_i(d,z) = Y_i ext{ if } D_i = d, Z_i = z$

• Observed treatment
$$D_i$$
 is a function of treatment assignment (Z_i) - it's a post-treatment quantity (and so has potential outcomes).

Assumptions

- 1. Randomization of instrument
- 2. Exclusion restriction
- 3. Non-zero first-stage relationship
- 4. Monotonicity

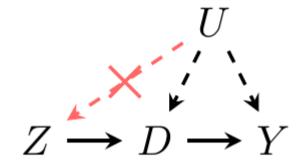
Assumption 1: Randomization

• Z_i is is independent of both sets of potential outcomes (potential outcomes for the treatment and potential outcomes for the outcome).

$$\{D_i(1),D_i(0)\}\!\perp\!\!\!\perp\!\!\! Z_i$$
 $\{Y_i(d,z)orall d,z\}\!\perp\!\!\!\perp\!\!\! Z_i$

- We can weaken this to conditional ignorability (where Z_i is randomized conditional on X_i), which is common in observational settings.
 - But if we don't believe conditional ignorability for the treatment, why would we believe it for the instrument?
- Sufficient to identify the intent-to-treat (ITT) effect

Assumption 1: Randomization



ullet The randomization assumption eliminates any arrows from U to Z.

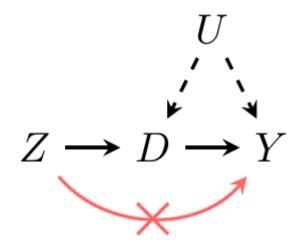
Assumption 2: Exclusion restriction

- Z_i only affects Y_i by way of its effect on D_i .
- In other words, if D_i were set at some level d, the potential outcome for $Y_i(d, z)$ does not depend on z.

$$Y_i(d,z) = Y_i(d,z') ext{ for any } z
eq z'$$

- Not a testable assumption! -- we have to justify this with substantive knowledge.
 - Easiest in the treatment non-compliance case
 - But consider what might happen in a non-blinded situation where respondents knew their treatment assignments.
- "Surprise" factor -- If I told you Z was associated with Y, would you think "that's odd"?

Assumption 2: Exclusion restriction



• The exclusion restriction eliminates any causal paths from Z to Y except for $Z \to D \to Y$.

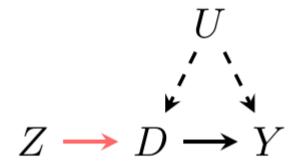
Assumption 3: Non-zero first stage

• Z_i has an effect on D_i

$$E[D_i(1)-D_i(0)]
eq 0$$

- Seems trivial, but we need this to make the estimator work.
- Magnitude matters for estimator performance a "weak" first-stage → heavily biased IV estimator
 - IV estimators are *consistent* but not *unbiased*.

Assumption 3: Non-zero first stage



ullet The non-zero first stage assumption requires a path from Z to D.

Assumption 4: Monotonicity

• Z_i 's effect on D_i only goes in one direction at the individual level

$$D_i(1)-D_i(0)\geq 0$$

- If it goes the other way, we can always flip the direction of the treatment to make this hold
 - \circ The key is that the instrument does not have a positive effect on D_i for some units and a negative effect for others.
- Not a testable assumption

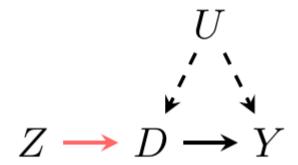
Assumption 4: Monotonicity

• In binary instrument/binary treatment world, this is sometimes called a "no defiers" assumption.

Stratum	$D_i(1)$	$D_i(0)$
"Always-takers"	1	1
"Never-takers"	0	0
"Compliers"	1	0
"Defiers"	0	1

• Under no defiers, every unit with $D_i=1$ and $Z_i=0$ is an always-taker, every unit with $D_i=0$ and $Z_i=1$ is a never-taker.

Assumption 4: Monotonicity



• Can't represent the monotonicity assumption in a DAG - it's an assumption about the form of the relationship between Z and D.

• The classic IV estimand with one instrument is a ratio of sample covariances.

$$au_{ ext{IV}} = rac{Cov(Y,Z)}{Cov(D,Z)}$$

• With a binary instrument, this is sometimes called the "Wald" estimand - a ratio of differences in means

$$au_{ ext{IV}} = rac{E[Y_i|Z_i=1] - E[Y_i|Z_i=0]}{E[D_i|Z_i=1] - E[D_i|Z_i=0]}$$

What does the Wald estimand correspond to in terms of causal effects?

$$au_{ ext{IV}} = rac{E[Y_i|Z_i=1] - E[Y_i|Z_i=0]}{E[D_i|Z_i=1] - E[D_i|Z_i=0]}$$

- Under our identification assumptions:
 - The numerator is the ITT
 - The denominator is the first-stage effect

Let's decompose the denominator first - under randomization:

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

 With binary treatment/binary instrument, we can use law of total expectation to decompose by principal stratum

$$egin{aligned} E[D_i(1) - D_i(0)] &= E[D_i(1) - D_i(0)|D_i(1) = D_i(0)] imes P(D_i(1) = D_i(0)) + \ E[D_i(1) - D_i(0)|D_i(1) > D_i(0)] imes P(D_i(1) > D_i(0)) + \ E[D_i(1) - D_i(0)|D_i(1) < D_i(0)] imes P(D_i(1) < D_i(0)) \end{aligned}$$

- The first term is 0
- And by no defiers, the last term is 0 since $P(D_i(1) < D_i(0)) = 0$

$$E[D_i(1) - D_i(0)] = Pr(D_i(1) > D_i(0))$$

• Next, the numerator (the ITT). Under the exclusion restriction and randomization:

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

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

• The difference (with some algebra) is

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

Conditioning on the principal strata:

$$egin{aligned} &= Eigg[(Y_i(1)-Y_i(0)) imes(0)|(D_i(1)=D_i(0))igg] imes P(D_i(1)=D_i(0))+\ &= igg[(Y_i(1)-Y_i(0)) imes(1)|(D_i(1)>D_i(0))igg] imes P(D_i(1)>D_i(0))+\ &= igg[(Y_i(1)-Y_i(0)) imes(-1)|(D_i(1)< D_i(0))igg] imes P(D_i(1)< D_i(0)) \end{aligned}$$

ullet Again, first term is zero because $D_i(1)-D_i(0)=0$, third is zero by "no defiers" and we have

$$E[Y_i|Z_i=1] - E[Y_i|Z_i=0] = Eigg[Y_i(1) - Y_i(0)|D_i(1) > D_i(0)igg] imes P(D_i(1) > D_i(0))$$

• The ITT is the product of a conditional average treatment effect and the proportion of compliers.

The LATE Theorem

• The IV estimand, under our identification assumptions, is a Local Average Treatment Effect (LATE):

$$rac{E[Y_i|Z_i=1]-E[Y_i|Z_i=0]}{E[D_i|Z_i=1]-E[D_i|Z_i=0]}=E[Y_i(1)-Y_i(0)|D_i(1)>D_i(0)]$$

- The LATE is a conditional average treatment effect within the *subpopulation* of **compliers**
- If treatment effects are constant, we can generalize this to the whole sample.
 - But if effects are heterogeneous, we are not necessarily getting a "representative" treatment effect.

Better LATE than never?

- How should we interpret the LATE?
 - It's not necessarily the quantity we care about we care about the effect of the treatment in the entire sample.
- Compliers are those compelled to take treatment by our encouragement. Would estimates generalize to those who are less encourageable?
 - The LATE is design-specific. If we came up with a different instrument, that changes the population on which we're estimating an effect!
 - What can we do?
 - We could describe the distribution of covariates among compliers vs. the population as a whole (Abadie's kappa-weighting).

- Gerber, Karlan and Bergan (2009, AEJ:AE) estimate the effect of reading the Washington Post (or Washington Times) on political attitudes and voting behavior.
 - $\circ Z_i$: Random assignment to receive a free subscription to the Washington Post
 - \circ D_i : Actually subscribing to the Washington Post (as measured by a post-encouragement survey)
 - \circ Y_i : 2005 Turnout (measured in the survey)
- Assumptions:
 - Assignment to get the free subscription offer is ignorable/exogenous
 - Getting the free subscription offer affects actual subscriptions (non-zero first stage)
 - No one would subscribe to the Post if they didn't receive the offer but not subscribe if they did. (monotonicity/no defiers)
 - Assignment to get the free subscription offer doesn't affect voting except through actually subscribing to the Post (exclusion restriction)

• First, subset the data to WaPo or control observations that completed the follow-up survey

```
green <- read_dta("assets/publicdata.dta")
wapost <- green %>% filter(treatment != "TIMES"&!is.na(getpost)&!is.na(voted))
```

• Is there a first-stage effect?

• About 34 percent of the sample is a "complier" - quite substantial!

• Is there an ITT?

• ITT is essentially zero.

Compare with the naive OLS estimate

- Post subscribers are 6pp more likely to vote in the 2005 VA gubernatorial election.
 - But is this causal? No!

Let's estimate the LATE using the Wald estimator

```
(mean(wapost$voted[wapost$post == 1]) - mean(wapost$voted[wapost$post == 0]))/(mean(wapost$getp
## [1] -0.00396
```

• Equivalent to a ratio of regression coefficients

```
coef(lm_robust(voted ~ post, data=wapost))[2]/coef(lm_robust(getpost ~ post, data=wapost))[2]
## post
## -0.00396
```

We'll talk about inference later, but take note: the SE for the LATE can be much larger than the SE for the ITT

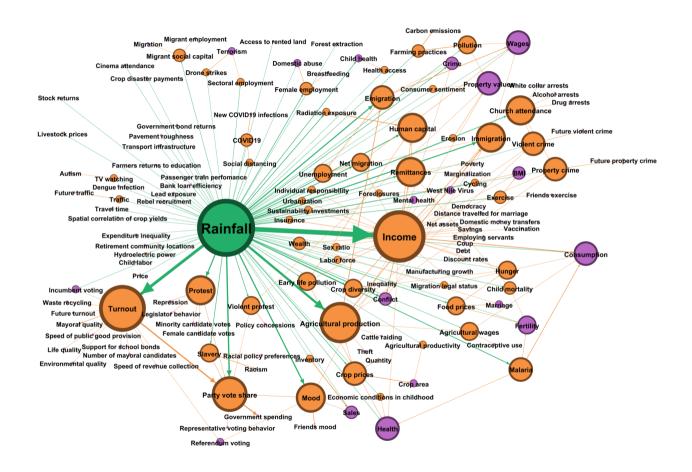
IV in observational studies

- Most applications of IV are not treatment non-compliance.
- But all follow the same underlying logic.
 - Treatment of interest is not randomized...but there exists a real or "natural" experiment that is.
 - And this natural experiment affects the outcome only through its affect on the treatment of interest.
- Examples:
 - Angrist (1990) Vietnam draft lottery number as an instrument for the effect of military service on income.
 - Angrist and Krueger (1991) Birth quarter as an instrument for education's effect on income.
 - Acemoglu et. al. (2001) European settler mortality as an instrument for the effect of institutional quality on GDP per capita.
 - Kern & Hainmueller (2009) West German TV signal strength as an instrument for the effect of watching West German TV on support for the East German regime
- Challenges
 - Exogeneity/ignorability isn't guaranteed
 - If an instrument has a large effect on your treatment of interest, it probably has an effect on other stuff that could affect the outcome as well (violating the exclusion restriction)

Discussion: The rainfall instrument

- Miguel, Satyanth, and Sergenti (2004, JPE) look at the effect of economic growth on civil conflict in 41 African countries.
 - Growth and conflict are confounded (e.g. by political institutions).
 - Instrument for GDP growth using the annual change in rainfall -- for heavily agrarian countries, rainfall fluctuations determine crop yields which are a large component of GDP.
 - Observe that changes in rainfall are associated with changes in GDP and negative GDP shocks (instrumented by rainfall) increase civil conflict.
- Does this satisfy the IV identification assumptions?
 - Exogeneity? Is rainfall as-good-as randomly assigned?
 - Monotonicity? Do positive rainfall shocks strictly boost GDP per capita?
 - Exclusion restriction? Is rainfall's effect transmitted only through the mechanism the authors define?

Discussion: The rainfall instrument



Mellon (2023) "Rain, Rain, Go Away: 195 Potential Exclusion-Restriction Violations for Studies Using Weather as an Instrumental Variable"

Estimation and inference for IV

IV with constant effects

• Let's consider a linear model for the potential outcomes

$$Y_i(d) = \alpha + au d + \gamma U_i + \eta_i$$

• If we could control for U_i , we could estimate the regression to get an estimate of τ

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

• But we can't - and regressing Y_i on D_i alone will not give a consistent estimator of τ since $Cov(\gamma U_i + \eta_i, D_i) \neq 0$

IV with constant effects

ullet Suppose Z_i is an instrument that is exogeneous and satisfies the exclusion restriction

$$Cov(\gamma U_i + \eta_i, Z_i) = 0$$

• Then, we can identify τ

$$egin{aligned} Cov(Y_i,Z_i) &= Cov(lpha+ au D_i+\gamma U_i+\eta_i,Z_i) \ &= Cov(lpha,Z_i) + Cov(au D_i,Z_i) + Cov(\gamma U_i+\eta_i,Z_i) \ &= au Cov(D_i,Z_i) \end{aligned}$$

Which gives us our IV estimand

$$au = rac{Cov(Y_i, Z_i)}{Cov(D_i, Z_i)}$$

IV estimator

• We can estimate τ by plugging in the sample quantities.

$$au_{ ext{IV}}^{\hat{}} = rac{\widehat{Cov(Y_i,Z_i)}}{\widehat{Cov(D_i,Z_i)}}$$

• This also can be written as a ratio of two regression coefficients

$$au_{ ext{IV}} = rac{\widehat{Cov(Y_i,Z_i)}/\widehat{Var(Z_i)}}{\widehat{Cov(D_i,Z_i)}/\widehat{Var(Z_i)}}$$

- Denominator: "First stage": Regression of D_i on Z_i
- ullet Numerator: "Reduced form": Regression of Y_i on Z_i

2SLS - Including covariates

- What if ignorability of Z_i only holds conditional on X_i -- or we want to include X_i as predictors to improve precision.
- We'll assume a particular structure for the outcome and treatment models

$$Y_i = X_i' \beta + \tau D_i + \epsilon_i$$

$$D_i = X_i' \alpha + \gamma Z_i + \nu_i$$

- Assume the X_i are exogenous but not excluded (appear in both equations). D_i is still endogenous so we can't get the treatment effect by just regressing outcome on treatment and covariates.
- Can we get an expression for Y_i in the form of Z_i alone?

2SLS - Including covariates

• Substitute in for D_i

$$egin{aligned} Y_i &= X_i'eta + au\left[X_i'lpha + \gamma Z_i +
u_i
ight] + \epsilon_i \ &= X_i'eta + au\left[X_i'lpha + \gamma Z_i
ight] + \left[au
u_i + \epsilon_i
ight] \ &= X_i'eta + au E[D_i|X_i,Z_i] + \epsilon_i^* \end{aligned}$$

- We can identify τ by regressing Y_i on X_i and the **fitted values** from a regression of D_i on X_i and the instrument Z_i .
- Intuition -- we want to only use the variation in D_i that is driven by the exogenous factor Z_i .

2SLS - Including covariates

You can also still get the ratio form of the IV estimator

$$Y_i = X_i'eta + au \left[X_i'lpha + \gamma Z_i +
u_i
ight] + \epsilon_i \ = X_i'(eta + aulpha) + au \gamma Z_i + \left[au
u_i + \epsilon_i
ight]$$

- The coefficient on Z_i in the reduced form regression is $\tau\gamma$
- The coefficient on Z_i in the first-stage is γ .
- So the ratio of the reduced form to the first-stage regression is τ .

Two-stage least squares

ullet First stage - Regress D_i on X_i and Z_i . Get the fitted values $\hat{D_i}$

$$\hat{D}_i = X_i'\hat{lpha} + \hat{\gamma}Z_i$$

ullet Second stage - Regress Y_i on X_i and fitted values \hat{D}_i

$$\hat{Y}_i = X_i'\hat{eta} + \hat{ au}\hat{D}_i$$

- The coefficient on the fitted values is the IV estimate
 - But, the standard errors will be wrong Why?

- Recall our Gerber, Karlan and Bergan (2009, AEJ:AE) experiment
 - $\circ Z_i$: Random assignment to receive a free subscription to the Washington Post
 - \circ D_i : Actually subscribing to the Washington Post (as measured by a post-encouragement survey)
 - \circ Y_i : 2005 Turnout (measured in the survey)
 - $\circ X_i$: Gender, Age
- Let's load and subset

```
green <- read_dta("assets/publicdata.dta")
wapost <- green %>% filter(treatment != "TIMES"&!is.na(getpost)&!is.na(voted)&!is.na(Bfemale)&!
```

• Our first stage regresses subscription on assignment + covariates

```
first stage <- lm robust(getpost ~ post + Bfemale + reportedage , data= wapost)
summary(first stage)
##
## Call:
## lm robust(formula = getpost ~ post + Bfemale + reportedage, data = wapost)
##
## Standard error type: HC2
##
## Coefficients:
##
              Estimate Std. Error t value Pr(>|t|) CI Lower CI Upper DF
## (Intercept) 0.12097
                         0.06406 1.889 5.94e-02 -0.004786 0.24673 729
                      0.03482 10.118 1.31e-22 0.283965 0.42069 729
       0.35233
## post
## Bfemale -0.00435 0.03505 -0.124 9.01e-01 -0.073156 0.06445 729
## reportedage 0.00170 0.00125 1.360 1.74e-01 -0.000756 0.00416 729
##
## Multiple R-squared: 0.134 , Adjusted R-squared: 0.13
## F-statistic: 35.4 on 3 and 729 DF, p-value: <2e-16
```

Let's actually run 2SLS - I like two routines: iv_robust in estimatr (does 2SLS with robust SEs)
 and ivmodel in ivmodel (does robust 2SLS and weak-instrument robust tests + other diagnostics)

```
wapo 2sls <- iv robust(voted ~ getpost + Bfemale + reportedage | post + Bfemale + reportedage,
summary(wapo 2sls)
##
## Call:
## iv robust(formula = voted ~ getpost + Bfemale + reportedage
      post + Bfemale + reportedage, data = wapost)
##
##
## Standard error type: HC2
##
## Coefficients:
##
              Estimate Std. Error t value Pr(>|t|) CI Lower CI Upper
              0.22562
                         0.07334 3.0766 2.17e-03 0.08165
                                                            0.3696 729
## (Intercept)
               0.00428
## getpost
                         0.09086 0.0471 9.62e-01 -0.17410
                                                            0.1827 729
## Bfemale -0.03495 0.03354 -1.0423 2.98e-01 -0.10079
                                                            0.0309 729
## reportedage 0.01040 0.00127 8.1879 1.19e-15 0.00791
                                                            0.0129 729
##
## Multiple R-squared: 0.093 , Adjusted R-squared: 0.0893
## F-statistic: 23.3 on 3 and 729 DF, p-value: 2.15e-14
```

LIML

##

1.00000

0.00428

```
wapo 2sls2 <- ivmodelFormula(voted ~ getpost + Bfemale + reportedage | post + Bfemale + report
summary(wapo 2sls2)
##
## Call:
## ivmodel(Y = Y, D = D, Z = Z, X = X, intercept = intercept, beta0 = beta0,
      alpha = alpha, k = k, manyweakSE = manyweakSE, heteroSE = heteroSE,
##
      clusterID = clusterID, deltarange = deltarange, na.action = na.action)
##
## sample size: 733
##
##
## First Stage Regression Result:
##
## F=111, df1=1, df2=729, p-value is <2e-16
## R-squared=0.132, Adjusted R-squared=0.131
## Residual standard error: 0.444 on 730 degrees of freedom
##
##
## Coefficients of k-Class Estimators:
##
##
               k Estimate Std. Error t value Pr(>|t|)
                  0.04708
## OLS
         0.00000
                           0.03247
                                        1.45
                                                 0.15
## Fuller 0.99863
                  0.00472
                          0.08979
                                      0.05 0.96
## TSLS
         1.00000
                  0.00428
                          0.09060
                                      0.05
                                               0.96
```

0.05

0.09060

0.96

• Our ratio estimator is consistent

$$\hat{ au}_{ ext{IV}} = rac{\widehat{Cov(Y_i, Z_i)}}{\widehat{Cov(D_i, Z_i)}} \stackrel{p}{
ightarrow} au + rac{Cov(U_i, Z_i)}{Cov(D_i, Z_i)}$$

- Under exogeneity $Cov(Z_i, U_i)$ is zero.
- However, when there are small violations of exogeneity, a weak instrument will amplify them.
- More generally, with a weak instrument, our t-ratio hypothesis tests assuming asymptotic normality will have **incorrect** type-1 error rates.
 - Why? Distributions of ratios are poorly behaved.

- Let's use a simulation to see how bad the bias can be in IV versus just a simple OLS regression of outcome on treatment under unobserved confounding.
- Let $U_i \sim \mathcal{N}(0,1)$ be an unobserved confounder. $Z_i \sim \mathrm{Bern}(.5)$ is an **exogenous** instrument.
- The probability of treatment is modeled via a logit

$$\logigg(rac{P(D_i=1|Z_i,U_i)}{1-P(D_i=1|Z_i,U_i)}igg) = \gamma Z_i + U_i$$

- γ here captures the relationship between the exogenous instrument Z_i and the treatment
- The outcome is a function of U and a mean zero error term ϵ_i only, so the true treatment effect is 0

$$Y_i = U_i + \epsilon_i$$

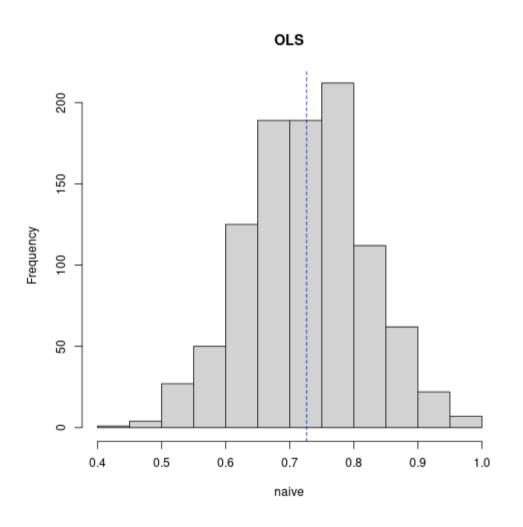
[1] -0.0196

ullet Let's see how the Wald estimator performs when we have a pretty large effect of Z_i on D_i : $\gamma=2$ and N=1000

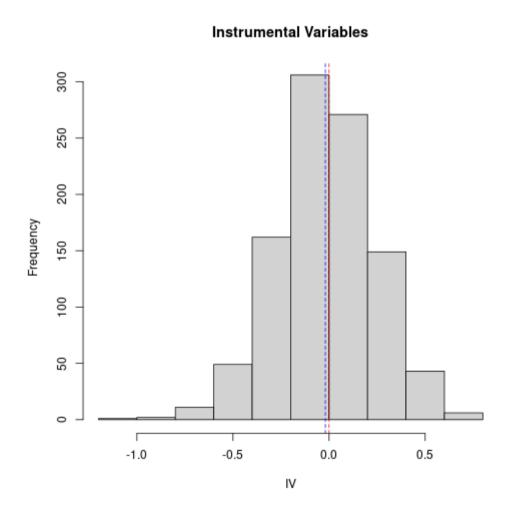
```
## First stage effect
 mean(firststage)
## [1] 0.345
## F-statistic from the first stage
 mean(firststageF)
## [1] 157
 ## Bias of the naive OLS Y ~ X
 mean(naive)
## [1] 0.726
 ## Bias of IV
 mean(IV)
```

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• Sampling distribution of the naive OLS estimator



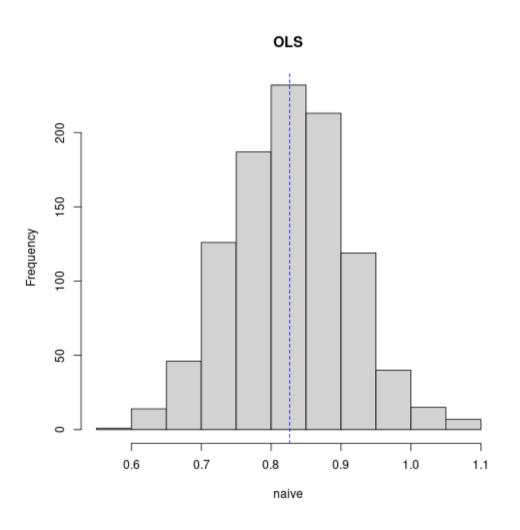
• Sampling distribution of the IV estimator



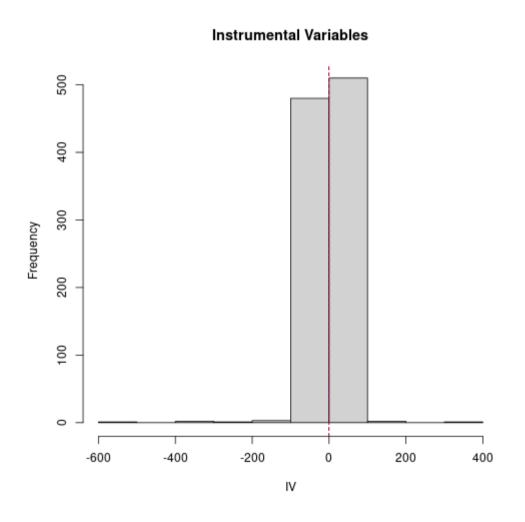
• Now, what happens when our instrument is weak: $\gamma = .2$ and N = 1000

```
## First stage effect
 mean(firststage)
## [1] 0.041
## F-statistic from the first stage
 mean(firststageF)
## [1] 2.6
 ## Bias of the naive OLS Y ~ X
 mean(naive)
## [1] 0.826
## Bias of IV
 mean(IV)
## [1] -1.04
```

• Sampling distribution of the naive OLS estimator



• Sampling distribution of the IV estimator



- When is an instrument too weak?
- Classic result: Staiger and Stock (1997), Stock and Yogo (2005) use first stage F-statistic thresholds

 ∼→ heuristic of first-stage F-statistic below 10.
- Recently: Lee, Moreira, McCrary, Porter (2020) -- If we want to use the F-statistic as a screen then we actually need F>104.7
- Suggestions:
 - Permutation tests using a test statistic that does not depend on the first stage.
 - Anderson-Rubin (1949) approach
 - Angrist and Pischke (2009) with "just-identified" IV (number of instruments = number of endogenous variables) bias is usually overwhelmed by the large standard errors.

Permutation test

- When the assignment process of Z_i is known, we can construct hypothesis tests using permutation inference assuming a constant treatment effect τ (Imbens and Rosenbaum, 2005).
 - \circ With a single, de-meaned instrument $\tilde{Z}_i = Z_i \bar{Z}$, we can construct a test statistic based on the sample covariance between Z_i and Y_i with the effect removed:

$$T(au) = rac{1}{N} \sum_{i=1}^N ilde{Z_i} imes (Y_i - au D_i)$$

- If the instrument is valid, under the null hypothesis that $\tau = \tau_0$, we can get the **randomization distribution** of the test statistic by simply re-randomizing treatment according to the known assignment process.
 - \circ Construct confidence intervals by "inverting the test" what values of au_0 does the test fail to reject?
- Alternative test statistics based on ranks of $Y_i \tau D_i$) (possibly within strata) can also be used.

Anderson-Rubin Test

- Even when the assignment process is not known, the IV assumptions allow us to construct a test statistic that does not depend on the first stage.
 - This is the **Anderson-Rubin (1949)** approach **Andrews, Stock and Sun (2019)** provide a good explanation especially for the "just-identified" case
- Let $\hat{\delta}_{\text{ITT}}$ be the **reduced form** or intent-to-treat estimate.
- The instrumental variables assumptions that the reduced form is related to the first stage π and the treatment effect τ

$$\delta_{ ext{ITT}} = \pi imes au$$

• Assuming a particular null $H_0: au = au_0$ implies that

$$\delta_{\mathrm{ITT}} - \pi imes au_0 = 0$$

Anderson-Rubin Test

 And so we can construct a test statistic based on the difference between the estimated ITT and the estimated first stage adjusted by the null which we know is normal in large samples.

$$g(au_0) = \hat{\delta}_{ ext{ITT}} - \hat{\pi} au_0 \sim \mathcal{N}(0,\Omega(au_0))$$

• The Anderson-Rubin (1949) test statistic is:

$$AR(au) = g(au)'\Omega(au_0)^{-1}g(au)$$

Under the null $H_0: \tau = \tau_0$, this has a chi-squared distribution which does not depend on the value of the first stage.

- Intuitively: Statistical properties of differences in two normal random variables are well-known and easy. Statistical properties of ratios are much more complicated!
 - Again, invert the test to get a confidence interval
 - \circ Can get **infinite** confidence bounds with a weak instrument the test **never rejects** for any value of au_0

Example: Strong instrument

```
wapo_iv <- ivmodelFormula(voted ~ getpost | post , data= wapost, heteroSE=T)
print(AR.test(wapo_iv))</pre>
```

```
## $Fstat
## [1] 0.0171
##
## $df
## [1] 1 731
##
## $p.value
## [1] 0.896
##
## $ci.info
## [1] "[-0.2058034696935, 0.175035583124674]"
##
## $ci
##
        lower upper
## [1,] -0.206 0.175
```

Example: Weak instrument

```
weak iv data <- data.frame(Y = Y, D = D, Z = Z)
 weak iv <- ivmodelFormula(Y ~ D | Z , data= weak iv data, heteroSE=T)</pre>
 print(AR.test(weak iv))
## $Fstat
## [1] 2.26
##
## $df
## [1] 1 998
##
## $p.value
## [1] 0.133
##
## $ci.info
## [1] "Whole Real Line"
##
## $ci
##
        lower upper
## [1,] -Inf Inf
```

Conclusion

- Instrumental variables lets us leverage *alternative* sources of randomness to learn about an otherwise confounded causal relationship.
- An instrument:
 - Affects treatment
 - Doesn't affect the outcome except through treatment
 - Is ignorable w.r.t the outcome.
- LATE theorem: The IV estimand is the ATE among those who would take treatment due to the instrument.
 - With continuous treatment/instrument a weighted average of LATEs (Angrist and Imbens, 1995)
 - With covariates a weighted average of covariate-specific LATEs
 - But be careful with this interpretation when the model is not fully saturated (Słoczyński, 2022)
- Statistical inference is tricky
 - Beware weak instruments typical large-sample asymptotics do poorly when instruments are irrelevant.
 - Consider weak-instrument robust tests (Anderson-Rubin)
 - If it's not in the reduced form, it's not real.