

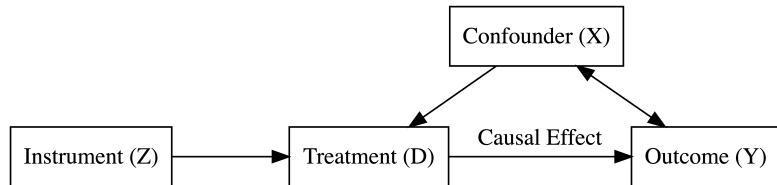
# Instrumental Variables and 2SLS

## Lecture 7 - Introduction to Causal Inference

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## Setup

**Issue:** We want to find the effect of treatment  $D$  on outcome  $Y$ , but there is a confounder  $X$ . We have an extra variable  $Z$ :

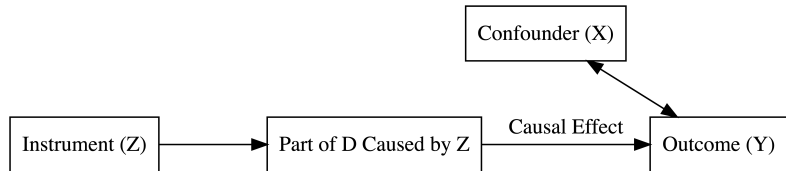


This **instrumental variable**  $Z$  has a few assumptions:

1.  $Z$  is correlated with the treatment  $D$  (**Relevance**).
2.  $Z$  is uncorrelated with any confounder  $X$  (**Exogeneity**).
3.  $Z$  has no direct effect on the outcome  $Y$ , only indirectly through  $D$  (**Exclusions Restriction**).
4.  $Z$  is exogenous to  $Y$  and  $D$  (also **Exogeneity**).

## Inducing Exogeneity in Treatment

Instead of using our original treatment variable  $D$ , let us instead only use the **part of  $D$  caused by  $Z$** . Let us call this  $\hat{D}$ .



The confounder  $X$  does not cause  $\hat{D}$ . Because  $\hat{D}$  is the part of  $D$  caused by  $Z$ .

- ▶ Thus,  $X$  is no longer a confounder - since it doesn't affect selection in  $\hat{D}$ .

Since  $\hat{D}$  is exogenous and there are no more confounders between  $\hat{D}$  and  $Y$ , we can calculate the causal effect of  $\hat{D}$  on  $Y$ .

## Two-Stage Least Squares

The way to estimate the causal effect in IV is with 2-stage least squares (2SLS) estimator.

**1st stage:** Regress  $D$  on  $Z$  to estimate  $\hat{D}$ .

$$D_i = \delta + Z_i\beta + \varepsilon_i$$

**2nd stage:** Regression  $Y$  on  $\hat{D}$  to estimate the causal effect:

$$Y_i = \alpha + \hat{D}_i\tau_{LATE} + u_i$$

The OLS estimate for  $\tau_{LATE}$  is our causal estimate.

- Note: Standard errors will be incorrect if you manually run the two stages. Use software like **R** with **fixest** package.

## Local Average Treatment Effect

The calculated treatment effect of  $\hat{D}$  on  $Y$  is called the local average treatment effect (LATE).

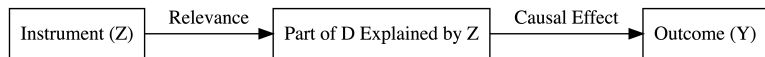
- ▶ Substantively, it is the causal effect of  $D$  on  $Y$  for the part of  $D$  explained by  $Z$
- ▶ This is also called the causal effect for **compliers**. Compliers are the units whose treatment  $D$  that “comply” (are influenced/caused) by the exogenous  $Z$ .

As noted before, this might not be equal to the total average treatment effect (ATE) between  $D$  and  $Y$ .

(LATE is sometimes called the Average Causal Response if  $D$  is continuous).

# Relevance and Weak Instruments

The relevance assumption is that  $Z$  must be correlated with  $D$ .



We can test relevance by running a regression of  $D$  on  $Z$  (which is the 1st stage of 2SLS), and see if  $\beta$  is significant:

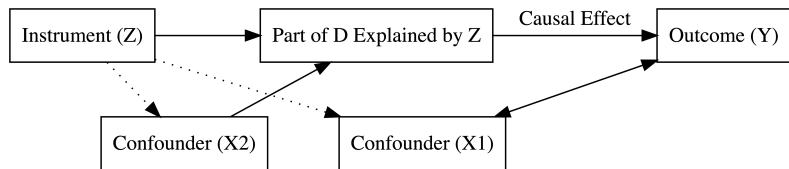
$$D_i = \delta + Z_i\beta + \varepsilon_i$$

If the correlation between  $Z$  and  $D$  is significant but weak, we have a weak instrument.

- ▶ If  $F$  test statistic in 1st stage is lower than 10,  $Z$  is weak.
- ▶ Weak instruments can have very biased  $\tau_{\text{LATE}}$  estimates, especially in small samples.

## Exogeneity Assumption

$Z$  must be exogenous/randomly assigned in respect to both  $D$  and  $Y$ . The dotted lines below shows violations to exogeneity:



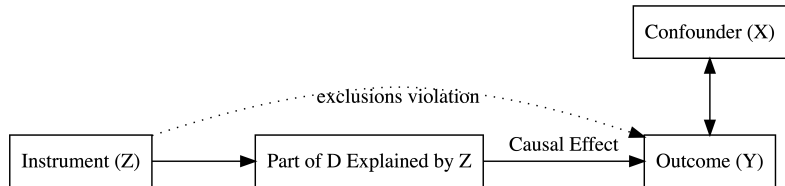
We can solve exogeneity violations by **controlling/accounting** for the confounders. We include confounders in both stages of 2SLS:

- ▶ 1st stage:  $D_i = \delta_0 + Z_i\delta_1 + X_i\delta_2 + \varepsilon_i$
- ▶ 2nd stage:  $Y_i = \alpha + \hat{D}_i\tau_{\text{LATE}} + X_i\beta + u_i$

Note: panel data - include fixed effects in both stages.

## Exclusions Restriction

The exclusions restriction states  $Z$  must not have a direct effect on  $Y$ . It can only have an indirect effect through  $D$ .



Why? Well if  $Z$  has an independent effect on  $Y$  outside of  $D$ , then  $Z$  is a confounder between  $\hat{D}$  and  $Y$ , and  $\hat{D}$  will no longer be exogenous.

There is no way to test the exclusions restriction. You can only justify it through your own understanding of the research topic in question.



## Reduced Form of 2SLS

The reduced form regression is the regression of  $Y$  on  $Z$ :

$$Y_i = \gamma_0 + Z_i\gamma_1 + \epsilon_i$$

Assuming  $Z$  is exogenous (add controls if necessary), that means the estimate  $\gamma_1$  is the causal effect of  $Z$  on  $Y$ .

► This effect is also called the **intent-to-treat effect**  $\tau_{ITT}$ .

If  $Z$  and  $D$  are both binary (quite common in causal inference), our LATE can also be estimated as:

$$\tau_{LATE} = \frac{\tau_{ITT}}{Pr(\text{compliers})} = \frac{\tau_{ITT}}{\widehat{Cov}(D_i, Z_i)}$$

► This is where the interpretation of LATE as the causal effect of compliers comes from.

# Finding Valid Instruments

It is difficult finding an instrument that plausibly satisfies relevance, exogeneity, and exclusions.

- ▶ In the econometrics literature, a lot of attention is put on trying to find an instrument that doesn't violate exclusions.
- ▶ However, **Exogeneity** is actually probably the more difficult assumption to meet - it is hard to find a  $Z$  that is truly randomly assigned in terms of both  $D$  and  $Y$ .
- ▶ The most reliable way to find instruments is with a non-compliance or fuzzy regression discontinuity, which we cover in lectures 9 and 11.
- ▶ Other common instruments are often random by nature: Lotteries, rainfall, natural disasters, random selection of beneficiaries for policy pilots, etc.

## Extension: Multiple Instruments

You do not need to stick with just one valid instrument. Including more instruments has a few advantages:

1. Having multiple valid instruments provides more variation in  $\hat{D}$ , allowing for more precise (and less variance) estimates.
2. Multiple instruments can also be a solution for weak-instruments.

**First Stage** ( $p$  number of instruments):

$$D_i = \delta_0 + \delta_1 Z_{1i} + \delta_2 Z_{2i} + \cdots + \delta_p Z_{pi} + \varepsilon_i$$

**Second Stage:** remains the same.

$$Y_i = \alpha + \hat{D}_i \tau_{\text{LATE}} + u_i$$