#### Panel Data Analysis

Lecture 2: From randomized controlled trial to two-way fixed effects 5 May, 2015 Prof. Andrew Eggers

#### Last week

#### Parallel trends assumption

**Diff-in-diff:** Binary treatment applied at a point in time to a subset of the units in the dataset



#### This week

Generalizing and expanding

- other kinds of grouped data (not necessarily time)
- general pattern of treatment application (not necessarily
- at **same point** in time)
- generalized treatment (not necessarily binary)

#### Back to RCT



#### More simulations!

Last time: from randomized controlled trial (RCT), we added a "baseline" measure: i.e. pre-treatment outcome for all units, with possible time trend.

This time: we start with an RCT in grouped data (e.g. pairs of twins participating in a drug trial; municipalities in districts participating in a field experiment) and add a second group (e.g. time periods).

Simulation 1: random assignment in grouped data (1) Generate data according to  $x_j \sim N(0,1)$ Voi ~  $N(x_i,i)$ 

$$y_{1i} \sim N(x_{j(i)} + \tau, 1)$$
$$\tau = 1$$

where i indexes units, j indexes groups, and j(i) indicates the group of unit i
(2) Assign treatment (d) randomly
(3) Estimate ATT (effect of d on y)
by

(3a) **Difference-in-means**: average difference in observed y between treated and control units

i.e. 
$$\mathrm{E}[y_i|d_i=1]$$
 -  $\mathrm{E}[y_i|d_i=0]$ 



Is the unconfoundedness assumption met in this case?

(3b) **Controlling for**  $x_j$ : Regression of observed  $y_i$  on  $d_i$  and  $x_{j(i)}$ i.e.  $y_i = \alpha + \beta_1 d_i + \beta_2 x_{j(i)} + \varepsilon_i$ (3c) **LSDV**: Regression of observed  $y_i$  on  $d_i$  and indicator for each j i.e.  $y_i = \alpha_{j(i)} + \beta_1 d_i + \varepsilon_i$ 

#### Simulation 1 (random assignment in grouped data): distribution of estimates across replications



(The more units per group, the more similar the densities for the two control strategies.)

Estimated ATT

Simulation 2: non-random assignment in grouped data

(1) Generate data according to  $\mathbf{x}_{i} \sim \mathbf{N}(0,1)$  $y_{0i} \sim N(x_{j(i)}, 1)$  $y_{1i} \sim N(x_{j(i)} + \tau, 1)$  $\tau = 1$ where i indexes units, j indexes groups, and j(i) indicates the group of unit i (2) Assign treatment (d) as function of  $x_i$ :  $Pr(d_i=1) = 1/(1 + exp(-x_i))$ (3) Estimate ATT (effect of d on y) by (3a) **Difference-in-means**: average difference in observed y between treated and control units i.e.  $E[y_i|d_i = 1] - E[y_i|d_i = 0]$ 



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**Important fact:** They yield the same estimate for  $\beta_1$ . Why?

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**Connection:** If the z's are the group dummies,  $\tilde{y} = \tilde{\beta}_0 + \tilde{\beta}_1 \tilde{d} + \tilde{\epsilon}$  is the deviations-from-means regression.

No matter how complex your analysis (fixed effects, clustering, etc), when your treatment is continuous you should show the conditional bivariate relationship (y on x, controlling for  $z_1$ ,  $z_2$ ,  $z_3$ , etc) in a **partial regression plot:** 

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## Partial regression plot

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# More intuition of fixed effects

Fixed effects regression addresses group-specific unobservable confounders. How? Think about what would happen if you **observed** and **included** these confounders:

• Consider LSDV version: 
$$y_i = \sum_{j=1}^J \alpha_j + \beta_1 d_i + \epsilon_i$$

If you included group-specific unobservable confounders, they would drop out due to multicollinearity with group dummies.

• Consider deviation from means version:  $\tilde{y_i} = \beta_1 \tilde{d_i} + \tilde{\epsilon_i}$ 

If you included group-specific unobservable confounders (before calculating deviations from means), they would drop out because they are constant within groups.

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#### Key features:

• Data are organized into groups (e.g. individuals within households, households within municipalities, municipalities within districts, countries within regions, etc).

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Does this apply to your case?

## Adding a second group dimension

In simulations I and 2, each unit belonged to I group



We can easily think about situations where each belongs to 2 or more groups



**Most common:** geographic unit and time period.

But also: class and ethnicity, education level and gender, etc.

# Treatment assignment on the basis of two group dimensions

Suppose there is an unobserved confounder (i.e. something related to treatment that affects potential outcomes) that varies by district and year.

Suppose the relationship between the confounder and the district and year is additive (**not** interactive), i.e.

- A. the confounder is higher for some **districts** than others
- B. the confounder is higher for some years than others
- but NOT true that confounder is higher for some district-year combinations than others, once we account for A and B

Then we get the right answer through LSDV with dummies for district and year, or regression with district & year fixed effects.

#### Simulation 3: random assignment in twice-grouped data

(1) Generate data according to  $x_j \sim N(0,1)$   $\lambda_t \sim N(0,1)$   $y_{0i} \sim N(x_{j(i)}+\lambda_{t(i)},1)$   $y_{1i} \sim N(x_{j(i)}+\lambda_{t(i)}+\tau,1)$  $\tau = 1$ 

where i indexes units, j indexes group 1, t indexes group t, and j(i) and t(i) indicate the group and time of unit i (2) Assign treatment (d) randomly (3) Estimate ATT (effect of d on y) by

(3a) **Difference-in-means**: average difference in observed y between treated and control units i.e.  $E[y_i|d_i = 1] - E[y_i|d_i = 0]$ 

(3b) Controlling for  $x_i$  and  $\lambda_t$ : Regression of observed  $y_i$  on  $d_i$ ,  $x_{i(i)}$ , and  $\lambda_{t(i)}$ i.e.  $y_i = \alpha + \beta_1 d_i + \beta_2 \times_{j(i)} + \beta_3$  $\lambda_{t(i)} + \epsilon_i$ (3c) Controlling for group 1: Regression of observed  $y_i$  on  $d_i$ and indicator for each j i.e.  $y_i = \alpha_{i}(i) + \beta_1 d_i + \varepsilon_i$ (3d) Controlling for groups 1 and 2: Regression of observed y<sub>i</sub> on d<sub>i</sub> and indicator for each j and t i.e.  $y_i = \alpha_{j(i)} + \alpha_{t(i)} + \beta_1 d_i + \varepsilon_i$ 

# Simulation 3 (random assignment in twice-grouped data): distribution of estimates across replications



Estimated ATT

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# Simulation 4 (non-random assignment in twice grouped data): distribution of estimates across replications



Recall: for binary treatment, the difference in means  $\mathrm{E}[y_{1i}|d_i{=}1] \text{ - } \mathrm{E}[y_{0i}|d_i{=}0]$ 

can be rewritten as

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$$\begin{split} E[y_{0i}|X_i,d_i=1] &= E[y_{0i}|X_i,d_i=0],\\ \text{then we have}\\ E[y_{1i}|X_i,d_i=1] - E[y_{0i}|X_i,d_i=0] &= E[y_{1i}|X_i,d_i=1] - E[y_{0i}|X_i,d_i=1] \end{split}$$

Diff-in-means controlling for  $\boldsymbol{X}$ 

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## CIA and panel data

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In diff-in-diff, CIA amounts to: parallel trends assumption. Generally in panel data (with unit and time fixed effects), CIA amounts to: no time-varying confounders within units.

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$$\begin{split} E[y_{0i}|X_{i},d_{i}{=}1] &= E[y_{0i}|X_{i},d_{i}{=}0], \\ & \text{then we have} \\ E[y_{1i}|X_{i},d_{i}{=}1] - E[y_{0i}|X_{i},d_{i}{=}0] &= E[y_{1i}|X_{i},d_{i}{=}1] - E[y_{0i}|X_{i},d_{i}{=}1] \\ & \text{Diff-in-means controlling for X} \end{split}$$
## **CIA** beyond binary treatment

Recall: for binary treatment, the difference in means  $\mathrm{E}[y_{1i}|d_i{=}1] \text{ - } \mathrm{E}[y_{0i}|d_i{=}0]$ 

can be rewritten as

$$\underbrace{E[y_{1i}|d_i=1] - E[y_{0i}|d_i=1]}_{ATT} + \underbrace{E[y_{0i}|d_i=1] - E[y_{0i}|d_i=0]}_{Selection bias}$$

Sometimes the independence assumption does not hold:

$$E[y_{0i}|d_i=1] \neq E[y_{0i}|d_i=0]$$

But if the conditional independence assumption (CIA) does hold



## Going beyond binary treatment

CIA extends beyond binary treatment:

Conditional on X, the units that get a particular value of treatment have the same potential outcomes for ALL values of treatment as the units that get another particular value of treatment. (See MHE 54-59).

Panel fixed effects regression can be seen as a particular set of controls that might make the CIA believable.

Whenever we causally interpret a regression, we make a conditional independence assumption (CIA); what varies is what is in the conditioning set and how convincing it is.

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You should always state your CIA and provide conditions in which it might be violated.

#### $limit_{it} = a_t + b_t population_{it} + c_t county_{it} + d_t county_{it} \times population_{it}$



In UK from 1885, constituency spending limits are a deterministic function of population and borough/county classification.

 $limit_{it} = a_t + b_t population_{it} + c_t county_{it} + d_t county_{it} \times population_{it}$ 



Win  $Margin_{it} = \alpha_i + \delta_t + \beta_1 Limit_{it} + f(population_{it}) + \varepsilon_{it}$ 

**Question**: how does spending limit affect avg margin between winner and loser?

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So, can think of this as:

- LSDV with dummies for constituency and year
- Regression of "demeaned" (by constituency and year) win margin on "demeaned" (by constituency and year) spending limits
- Regression of residuals from [regression of win margin on constituency and year dummies] on residuals from [regression of spending limits on constituency and year dummies]

Relationship between spending limits and WinMargin:





y-axis: residuals from regressing WinMargin on constituency and year dummies



y-axis: residuals from regressing WinMargin on constituency and year dummies x-axis: residuals from regressing spending limits on constituency and year dummies



What is the CIA here?

What is the CIA here?

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"In absence of treatment, affected constituencies would have evolved as unaffected [constituencies]."

i.e. no time-varying confounders.

i.e. changes in spending limits not systematically related to changes in competition, controlling for changes in other places

## Example: Levitt (1994) on effects of campaign spending

Levitt (1994), "Using Repeat Challengers to Estimate the Effect of Campaign Spending on Election Outcomes in the U.S. House".

**Question:** What is the effect of campaign spending on election outcomes?

Consider running this cross-sectional regression:

 $\label{eq:def_DemCongVoteShare_i} DemCongVoteShare_i = \beta_0 + \beta_1 \; (DemSpend_i - RepSpend_i) + \beta_2 \; DemPresVoteShare_i + \epsilon_i$  where

*DemCongVoteShare*<sub>i</sub>: Vote share for Democratic congressional candidate in district i *DemSpend*<sub>i</sub>, *RepSpend*<sub>i</sub>: Spending by Democratic and Republican congressional candidates in district i *DemPresVoteShare*<sub>i</sub>: Vote share for Democratic presidential candidate in district i

- Would you expect  $\beta_1$  to be positive or negative?
- What CIA is necessary to interpret that coefficient causally?
- Why might this CIA be violated?

## Example: Levitt (1994) on effects of campaign spending (2)

Question: What is the effect of campaign spending on election outcomes?

**Unmeasured confounder**: Candidate quality (i.e. attractiveness)

**Research design:** Two-way fixed effects, where groups are (i) *pairs* of candidates and (ii) years. (Also includes scandal and incumbency dummies.)

#### Ways of stating the CIA:

- Levitt (782-783): "An individual candidate's quality must be constant over time."
- Election-relevant features of pairs of candidates are fixed over time.
- The same pair of candidates in a different year is comparable, after controlling for nationwide year-to-year swings in electoral outcomes.
- Variation in spending over time within a given pair of candidates is unrelated to potential outcomes conditional on the year (and other controls).

To discuss:

- How might this CIA be violated?
- Do pairs of candidates who only appear once in the dataset contribute anything to the estimation of the effect?

# Fixed effects and first-differences

Consider situation where (as in simulation)  $x_{j(i)}$  is a time-invariant confounder:  $y_{it} = \beta_0 + \beta_1 d_{it} + \beta_2 x_{j(i)} + \epsilon_{it}$ 

Above we addressed this with LSDV/group fixed effects, which we showed was equivalent to a deviation-from-means regression:

$$\widetilde{y}_{it} = \beta_{0} + \beta_{1} \ \widetilde{d}_{it} + \epsilon_{it}$$

Another way to address time-invariant confounders in panel data: first-differences. Calculate  $\Delta y_{it} = y_{it} - y_{i,t-1}$ , etc and estimate

$$\Delta y_{it} = \beta_0 + \beta_1 \Delta d_{it} + \Delta \epsilon_{it}$$

#### As explained in MHE (224):

- Algebraically equivalent when just two periods
- Not otherwise, but consistent (i.e. as sample size increases, both converge to truth)

# Example: Ansell (2014) on effect of house prices on political preferences

Ansell (2014), "The Political Economy of Ownership: Housing Markets and the Welfare State"

**Question:** How does variation in house prices affect homeowners' preferences regarding redistribution?

Consider running this cross-sectional regression:

 $SupportForRedistribution_{i} = \beta_{0} + \beta_{1} PriceOfHouse_{i} + \beta_{2} Income_{i} + \beta_{3} Age_{i} \varepsilon_{i}$ 

- Would you expect  $\beta_1$  to be positive or negative?
- What CIA is necessary to interpret that coefficient causally?
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# Example: Ansell (2014) on effect of house prices on political preferences

Ansell (2014), "The Political Economy of Ownership: Housing Markets and the Welfare State"

**Question:** How does variation in house prices affect homeowners' preferences regarding redistribution?

**Research design:** First-difference regression in seven-wave British Household Panel Survey

**The CIA:** Confounding variables are constant within individuals between two waves of panel.

How might this be violated?

## Not all panel FE analysis is causal

Example: Fowler (2015), "Do elections select better representatives?"

**Question:** Why do incumbents win more electoral support than non-incumbents? **Research design:** Descriptive decomposition.

• Estimate

 $DemVoteShare_{i} = \beta_{0} + \beta_{1} (DemIncumbent_{i} - RepIncumbent_{i}) + \varepsilon_{i}$ 

• Estimate same thing with state-decade and year fixed effects:

 $DemVoteShare_{i} = \alpha_{j} + \gamma_{t} + \beta_{1} (DemIncumbent_{i} - RepIncumbent_{i}) + \varepsilon_{i}$ 

- Interpret difference in  $\beta_1$  as a measure of "party match" component of incumbent success (tendency of incumbents to belong to locally-popular party)
- (Estimate "officeholder benefits" via RDD)

The CIA: Not relevant, because no causal claims.
#### How do you test the CIA?

Because of the fundamental problem of causal inference, CIA is **always** an assumption; it cannot be directly tested.

Diff-in-diff is a special case of panel fixedeffects regression that allows for a very nice indirect test.

Why is it possible?

Parallel trends assumption looks good



How do you test the CIA? (2)

**Transparent indirect tests of the CIA**: a characteristic of my favorite research designs.

e.g. regression discontinuity designs (RDD) (at right: Lee (2008), "Randomized experiments from nonrandom selection in U.S. House elections")



# How do you test the CIA? (3)

What do these cases have in common? The fact that we slightly **extend** the CIA and test the additional implications.

- Diff-in-diff:
  - CIA: Parallel trends assumption applies between pre- and post-treatment (i.e. potential outcomes are same for treatment group and control group post-treatment, after adjusting for time-invariant difference)
  - Extension to CIA: Parallel trends assumption should apply between pre-pretreatment and pre-treatment period too
- RDD:
  - CIA: At 0% threshold of vote share margin, potential outcomes are independent of treatment status (i.e. difference in outcomes between winners and losers of dead heat is due to treatment)
  - Extension to CIA: At 0% threshold of vote share margin, pre-treatment covariates should also be independent of treatment status

## How do you test the CIA? (4)

• Generalization of diff-in-diff (panel fixed effects design):

- CIA: No time-variant unit specific confounders
- Extension to CIA: When including dummies for "2 years before treatment", "1 year before treatment", etc in regression, lagged treatment has no effect (see Kuziemko & Werker 2006 for example; also MHE 237-238)

#### General pattern:

- does something that should have no effect have an effect ("placebo")
- does treatment affect something it shouldn't? ("placebo outcome", falsification test)

As we add complexity from simple diff-in-diff (esp when we go beyond binary treatment), testing the CIA becomes harder and harder.

A problem with transparent designs: You get asked for placebo tests!

## Wrapping up

If there are unobserved confounders, but you think they may be fixed within groups (or within units over time), fixed effects (or first differences) can help.

All causal inference requires a conditional independence assumption.

If panel methods are especially credible, it is because

- CIA may be particularly credible within units over time
- some cases (e.g. simple diff-in-diff) allow for transparent indirect tests of CIA

**Next time:** lagged dependent variables, synthetic control method, random effects, standard errors (or a subset)