White Rose Social Sciences Doctoral Training Partnership

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- 8 April 2025, University of Leeds

Potential outcomes framework

Goal in causal inference is to assess the causal effect of a treatment/exposure on some outcome

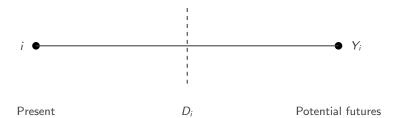
- → Does raising the minimum wage reduce employment?
- → Does housing assistance reduce homelessness?
- → Do body-worn cameras reduce police use of force?
- → Does voting by mail increase voter turnout?
- → Does exposure to misinformation reduce political trust??

~→ ...

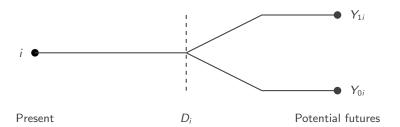


Potential futures Present

Potential outcomes framework



Potential outcomes framework



Y_i : Observed outcome variable of interest for unit i

Potential outcomes

 Y_{0i} and Y_{1i} : Potential outcomes for unit

$$Y_{\cdot i} = \left\{ egin{array}{ll} Y_{1i} & ext{Potential outcome for unit } i ext{ with treatment} \\ Y_{0i} & ext{Potential outcome for unit } i ext{ without treatment} \end{array} \right.$$

 D_i : Indicator of treatment intake for *unit*

$$D_i = \begin{cases} 1 & \text{if unit } i \text{ received the treatment} \\ 0 & \text{otherwise.} \end{cases}$$

Definition of causal effect

$$\delta_i = Y_{1i} - Y_{0i}$$

Fundamental problem of causal inference

 \rightarrow We cannot observe both potential outcomes for the same unit i!

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Randomisation solves the problem!

Logic of randomised control trials

- → Randomly divide a sample in two groups
- → Because this was random, both groups are on average the same
- Then apply the treatment/exposure to one group (the treatment group), but not the other (control group)
- → Because the exposure happened after the treatment assignment, the only difference between the two groups is the treatment/exposure
- Therefore, any subsequently observed differences are attributable to the treatment/exposure
- we we randomisation, we can thus find the average treatment effect

What if we cannot conduct an experiment?

- → Randomised Experiments
- ∴ Observational Studies
 - Selection on observables
 - Regression
 - Matching
 - Weighting
 - Selection on unobservables
 - Difference-in-Differences and synthetic control
 - Instrumental Variables
 - Regression Discontinuity Designs

- Causality is defined by potential outcomes, not by realised (observed) outcomes
- → Observed association is neither necessary nor sufficient for causality
- → Estimation of causal effects of a treatment (usually) starts with studying the assignment mechanism
- The goal is to mimic the features of a randomised experiment even if we don't have one
- when we don't have an RCT, our ability to make causal inferences often relies on making untestable assumptions about the assignment mechanism
- ⇒ Now let's see how we can leverage panel data to make causal inferences!

Potential outcomes framework

Difference-in-differences

⇒ What if we use **time** in our favour?

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- \rightsquigarrow Collect data on Y at two points in time: before and after the treatment/exposure/policy intervention

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- \rightsquigarrow Collect data on Y at two points in time: before and after the treatment/exposure/policy intervention
- Analyse the extent to which Y changes in units that received the treatment
- Analyse the extent to which Y changes in units that did NOT receive the treatment
- → Compare the two changes

Some conceptual clarification to make our lives easier

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- → Variation between units: difference
- → Variation within units (over time): changes

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- ⇒ We want to estimate the difference in changes or (difference-in-differences)

Some conceptual clarification to make our lives easier

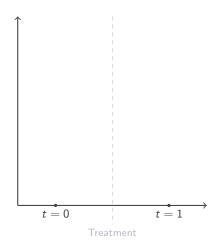
→ Variation between units: difference

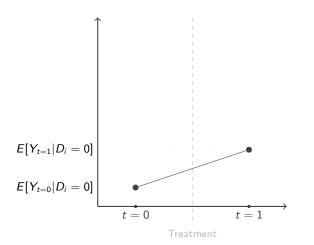
or (difference-in-differences)

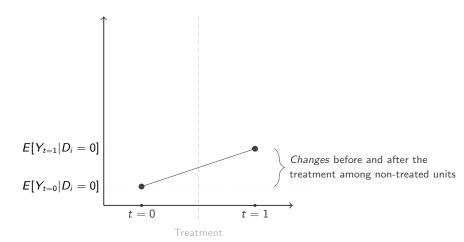
- → Variation within units (over time): changes
- ⇒ We want to estimate the difference in changes
 - \rightarrow The difference between (a) changes in Y before and after the intervention among treated units and (b) changes in Y before and after

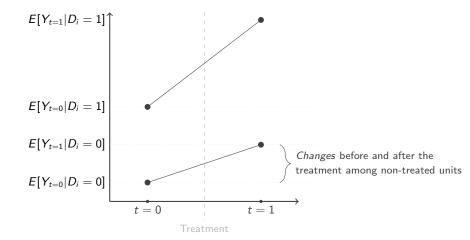
the intervention among non-treated units is the causal effect!

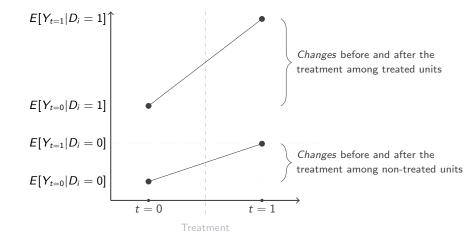
(under some assumptions regarding those changes... Let's dive into it)

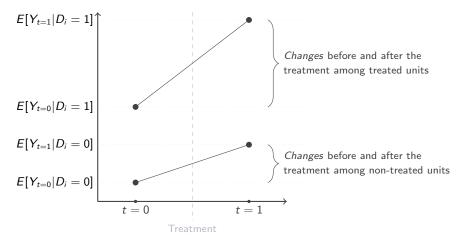






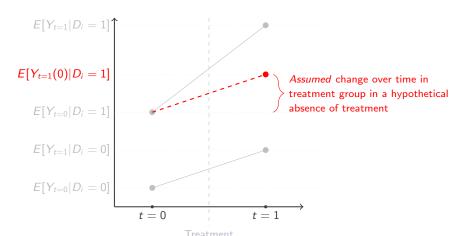






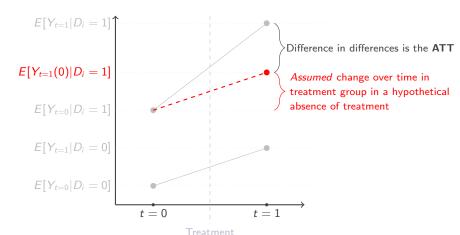
 \rightarrow Problem: Missing potential outcomes: $E[Y_{i,t=1}(0)|D_i=1]$ and $E[Y_{i,t=1}(1)|D_i=0]$

Strategy: Use the change in the control group to assume $E[Y_{t=1}(0)|D_i=1]$



Assumption: Trend over time would be the same for treatment and control

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Assumption: Trend over time would be the same for treatment and control

Identification assumption

Parallel trends

→ Had the treated units not received the treatment, they would have followed the same trend as the control units

Difference-in-differences estimator

Difference in changes:

$$\delta_{ATT} = \Big\{ \text{Changes in treatment group before and after treatment} \Big\} \\ - \Big\{ \text{Changes in control group before and after treatment} \Big\}$$

Threats to validity

Non-parallel trends

- Very critical assumption: treatment units have similar trends to control units in the absence of treatment
- ~ Fundamental problem of causal inference: we cannot observe potential outcome under the control condition for treated units in the post-treatment period
 - ⇒ What can we do? (more on that later...)
 - · Careful assessment: is assuming parallel trends plausible?
 - · Estimate treatment effects at different time points (placebo tests)

Using regression to estimate the difference-in-differences

We can obtain the difference in differences using regression techniques.

$$Y_i = \alpha + \beta_1 \cdot D_i + \beta_2 \cdot T_i + \delta \cdot (D_i \cdot T_i) + \varepsilon.$$

We can see that

$$E[Y_i|D_i,T_i]$$
 $T_i=0$ $T_i=1$ Changes after - before $D_i=0$ $lpha$ $lpha+eta_1$ $lpha+eta_1+eta_2+\delta$ $eta_2+\delta$ Treated - control eta_1 $eta_1+\delta$ δ



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$E[Y_i D_i,T_i]$	$T_i = 0$	$T_i = 1$	Changes after - before
$D_i = 0$	α	$\alpha + \beta_2$	eta_2
$D_i = 1$	$\alpha + \beta_1$	$\alpha + \beta_1 + \beta_2 + \delta$	$\beta_2 + \delta$
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Estimator (Regression 2)

With panel data we can use regression with first differences:

$$\Delta Y_i = \alpha + \delta \cdot D_i + X'\beta + u.$$

where
$$\Delta Y_i = Y_i(1) - Y_i(0)$$
.

With two periods this gives the same result as other regressions

1. We can include covariates

- Controlling for some covariates may increase precision
- Time-varying covariates may strengthen the parallel assumptions
- (add covariates cautiously! e.g., beware of post-treatment bias)
- - (not just binary)

Advantages of the regression estimator

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- 2. Easy to calculate standard errors
 - (though be careful about clustering)
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Advantages of the regression estimator

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 - (add covariates cautiously! e.g., beware of post-treatment bias)
- 2. Easy to calculate standard errors
 - (though be careful about clustering)
- 3. Easy to extend to other types of treatment
 - (not just binary)

- >> This setup only works for the simplest scenario with two time periods
 - → It doesn't make use more periods
 - Useful to make careful assessments of time trends
 - → Sometimes different units are treated at different time points

Difference-in-differences with multiple periods

>> Assume a pool of structured data

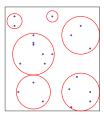


Multiple periods

>> Assume a pool of structured data



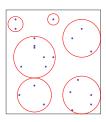
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 \rightsquigarrow Each dot represents a unit *i*

 \leadsto Each circle represents a group j

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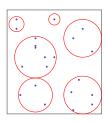


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· Pooled approach

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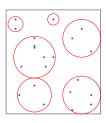


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- · Pooled approach
- · Between approach

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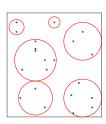


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- · Pooled approach
- · Between approach
- · Random Effects

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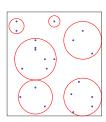
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- · Pooled approach
- · Between approach
- · Random Effects
- · Fixed Effects

- >> Focus on within-group variation
- >> Implementation: dummy variables for each group j (γ_i)

$$Y_{ij} = \gamma_j + \beta \cdot X_{ij} + \varepsilon$$

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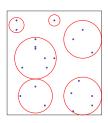
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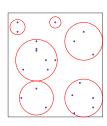
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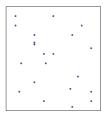
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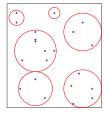
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 \leadsto Each dot represents a measure t

Fixed-effect regression with panel data

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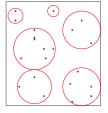
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$$Y_{it} = \gamma_i + \beta \cdot X_{it} + \varepsilon$$

Multiple periods

Fixed-effect regression with panel data

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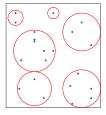
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Fixed-effect regression with panel data

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>> What about time fixed-effect?

DiD: Two-way fixed-effect regression

Estimator (Regression with Multiple Time Periods)

We can generalise to multiple groups/time periods using unit and period fixed-effects ('two-way' fixed-effect model):

$$Y_{it} = \gamma_i + \alpha_t + \delta \cdot D_{it} + \varepsilon$$

- γ_i is a fixed-effect for units (dummy for each unit)
- α_t is a fixed-effect for time periods (dummy for each period)
- δ is the DiD estimate based on D_{it}

- we can replace D_{it} with almost any type of treatment (not only binary)
- we can extend easily to multiple periods
- we can have units treated at different times
- we can estimate unit-specific time trends by including a unit-period interaction

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Very flexible approach

- we can replace D_{it} with almost any type of treatment (not only binary)
- we can extend easily to multiple periods
- we can have units treated at different times
- we can estimate unit-specific time trends by including a unit-period interaction
 - → useful when treatment occurs at different times for different units and there are slight deviations from parallel trends

Multiple periods

DiD: Two-way fixed-effect regression

- - i.e., changes over time!

$$ightarrow \hat{\delta}
ightarrow \hat{\delta}_{ATT}$$
 (it might not be that simple...)

- Fine in panel data, as we have same units at several points in time
- units in each time period Longitudinal Data Analysis

- \rightarrow Unit FEs means that we are only using within unit variation in Y to calculate the effect of D
 - i.e., changes over time!
 - This removes all time-constant confounders

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- Time FEs means that we remove the effect of any changes to the response variable that affect all units at the same time

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Slides: ThiagoROliveira.com/2-LDA-2025.pdf

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- It is hard to provide a visual inspection of the parallel trends assumption here as treatment may switch on at different time for different units
- Nevertheless, we are still assuming that treated/control units would have evolved identically over time in absence of treatment
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- It is hard to provide a visual inspection of the parallel trends assumption here as treatment may switch on at different time for different units
- Nevertheless, we are still assuming that treated/control units would have evolved identically over time in absence of treatment
- >> Why not always use unit dummies?
 - Fine in panel data, as we have same units at several points in time
 - Not possible with repeated cross-section when we do not have the same units in each time period Longitudinal Data Analysis

Some caution with two-way fixed-effect models



A rare photo of an applied economist keeping up with the difference-indifferences literature





When Should We Use Unit Fixed Effects Regression Models for Causal Inference with Longitudinal Data? 🛍 😝

Kosuke Imai

Harvard University

In Song Kim Massach

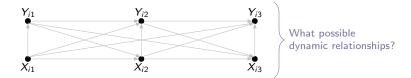
Massachusetts Institute of Technology

- → Imai & Kim (2019) show that unit FEs might not be that effective in adjusting for unobserved time-constant confounders
- → The issue is related to possible dynamic causal relationships



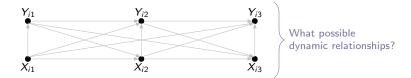
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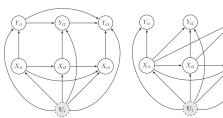
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→ Some dynamic causal relationships compromise unit FEs

FIGURE 2 Identification Assumptions of Regression Models with Unit Fixed Effects



(a) past outcome affects current outcome

(b) past treatments affect current outcome



 X_{i3}

(d) past outcomes affect both current outcome and treatment. Longitudinal Data Analysis

(1) Past outcome affects current outcome

(2) Past treatments affect current outcome

→ (3) Past outcomes affect current treatment

(4) Past outcomes affect current outcome and treatment

(c) past outcomes affect current treatment Thiago Oliveira

Slides: ThiagoROliveira.com/2-LDA-2025.pdf

Key assumptions of unit fixed effects models

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- 1. Past treatments do not directly influence current outcome

Key assumptions of unit fixed effects models

- 1. Past treatments do not directly influence current outcome
- 2. Past outcomes do not affect current treatment

- → Causal inference with observational data is really hard!
- → Longitudinal data can help, but it's not a silver bullet
 - · Have a look at all assumptions involved
 - · Parallel trends is an untestable assumption
- → This is a fast-changing topic. Keep up with the literature!
 - Callaway and Sant'Anna (2020); Callaway et al. (2021); Imai et al. (2021); Goodman-Bacon (2018); Imai and Kim (2019)
- Now let's see how to estimate those models using R!
 - Find the lab notes here: thiagoroliveira/2-LDA-lab.html

Thank you!

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