

# 7. Difference-in-Differences

Empirical Evaluation of Economic Policy

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# Overview

- 1 Panel data methods
  - Error-components model
  - Estimators
- 2 Difference-in-Differences designs
  - Binary designs
  - Complex designs
- 3 Application

# Overview

## 1 Panel data methods

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# Error-components model

- Traditionally, the most commonly used model to study relationships in panel data settings
- **Two-way error-components model:**

$$y_{it} = \mathbf{x}_{it}'\boldsymbol{\beta} + \varepsilon_{it} \quad i \in \{0, \dots, N\}, \quad t \in \{0, \dots, T\} \quad (1)$$

with

$$\varepsilon_{it} = \alpha_i + \gamma_t + \eta_{it}. \quad (2)$$

Here,

- $\alpha_i$ : **unobserved unit-specific effect** (e.g., innate ability)
- $\gamma_t$ : **unobserved period-specific effect** (e.g., national policies)
- $\eta_{it}$ : remaining disturbance

## Two-way fixed effects model

Plugging (2) into equation (1) gives us the **Two-way fixed effects model**:

$$y_{it} = \alpha_i + \gamma_t + \mathbf{x}'_{it}\boldsymbol{\beta} + \eta_{it} \quad (3)$$

where

- $\alpha_i$ : unit-specific intercept
- $\gamma_t$ : period-specific intercept

**Note:** Effects of *unit-specific time-invariant* characteristics **and** *period-specific unit-invariant* characteristics will be absorbed by  $\alpha_i$  and  $\gamma_t$ , resp.

# Estimating fixed effects models

Recall,  $\alpha_i$  and  $\gamma_t$  are **unobserved**, but primary interest is  $\beta$

**Three** possible ways to go about estimation:

- Within-estimator
- Least Squares Dummy Variables (LSDV) estimator
- First-difference estimator (will not be discussed)

## Within-estimator – OWFE estimator

If  $\gamma_t = 0$  for all  $t$  (no period-specific effects), (3) reduces to a **One-way fixed effects model**:

$$y_{it} = \alpha_i + \mathbf{x}'_{it}\boldsymbol{\beta} + \eta_{it} \quad (4)$$

We will first consider the within-estimator in this simplified setting

## Within-estimator – OWFE estimator

Consider transforming all variables – **including** individual fixed effects  $\alpha_i$  – as follows (taking the dependent variable as example):

$$\tilde{y}_{it} = y_{it} - \frac{1}{T} \sum_{t=1}^T y_{it}$$

Clearly, given this transformation, the individual-specific effects  $\alpha_i$  will be **eliminated** from the regression:

$$\begin{aligned}\tilde{\alpha}_i &= \alpha_i - \frac{1}{T} \sum_{t=1}^T \alpha_i \\ &= 0.\end{aligned}$$



## Within-estimator – OWFE estimator

The **within-estimator** is then simply defined as the *pooled* OLS estimator of the **transformed** model:

$$\tilde{y}_{it} = \tilde{\mathbf{x}}'_{it}\beta + \tilde{\eta}_{it}$$

In the one-way fixed effects model, the within estimator is often referred to as the **One-way fixed effects (OWFE) estimator**

## Within-estimator – TWFE estimator

Consider transforming all variables – **including** individual and time fixed effects  $\alpha_i$  and  $\gamma_t$  – as follows (taking the dependent variable as example):

$$\check{y}_{it} = y_{it} - \frac{1}{T} \sum_{t=1}^T y_{it} - \frac{1}{N} \sum_{i=1}^N y_{it} + \frac{1}{NT} \sum_{t=1}^T \sum_{i=1}^N y_{it}$$

Clearly, given this transformation, **both** the individual-specific effects  $\alpha_i$  and period-specific effects  $\gamma_t$  will be **eliminated** from the regression:

$$\check{\alpha}_i = \check{\gamma}_t = 0.$$

## Within-estimator – TWFE estimator

Again, the **within-estimator** in this setting is defined as the *pooled* OLS estimator of the **transformed** model:

$$\check{y}_{it} = \check{\mathbf{x}}'_{it}\beta + \check{\eta}_{it}$$

In the one-way fixed effects model, the within estimator is often referred to as the **Two-way fixed effects (TWFE) estimator**

**Note:** The **TWFE estimator** is historically the most commonly used estimator in Difference-in-Differences settings

# LSDV estimator

It can be shown that the TWFE estimator is *equivalent* to a particular **Least Squares Dummy Variables (LSDV) estimator**:

$$y_{it} = \sum_{j=1}^N \alpha_j 1\{i = j\} + \sum_{t'=1}^T \gamma_{t'} 1\{t = t'\} + \mathbf{x}'_{it} \boldsymbol{\beta} + \eta_{it} \quad (5)$$

This can easily be implemented in Stata using `regress`:

```
regress y i.unit_id i.period_id x
```

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## Simple DiD

Often in economics, having randomized treatments is **not possible**

- RCTs may be unfeasible due to practical or ethical reasons
- RCTs may lack *external validity*

**Hence**, researchers rely on **natural experiments** to estimate treatment effects

- Induced by policy changes (e.g., changes in U.S. state's minimum wage laws)
- Assignment to treatment is generally **not randomized**
- Simply comparing control and treated units leads to biased estimates for treatment effects

# Simple DiD

Consider two locations  $g \in \{s, c\}$  and two time periods  $t \in \{0, 1\}$ :

- $Y_{g,t}(0)$ : Potential outcome at location  $g$  in  $t$  *without* treatment  
→ employment level with low minimum wages
- $Y_{g,t}(1)$ : Potential outcome at location  $g$  in  $t$  *with* treatment  
→ employment level with high minimum wages

In  $t = 0$ , both locations  $g$  are untreated so that  $Y_{g,0} = Y_{g,0}(0)$ .

**However**, in  $t = 1$ :

- For the treated group  $s$ :  $Y_{s,1} = Y_{s,1}(1)$
- For the control group  $c$ :  $Y_{c,1} = Y_{c,1}(0)$

## Simple DiD

We would like to estimate  $E[Y_{s,1}(1) - Y_{s,1}(0)]$

→ average effect of increasing minimum wages in location  $s$  at period  $t = 1$  (Card & Krueger, 1994)

To overcome that treatment assignment to  $g$  may not be random, we could use the following **simple DiD estimator**:

$$\text{DiD} = (Y_{s,1} - Y_{s,0}) - (Y_{c,1} - Y_{c,0})$$

Under the assumption that in the absence of treatment, both locations  $g$  would have experienced the **same** average outcome **evolution**, the **simple DiD estimator** is unbiased:

$$E[\text{DiD}] = E[Y_{s,1}(1) - Y_{s,1}(0)].$$



## Dynamic potential outcomes

To make the exposition slightly more general, we extend the standard potential outcomes framework

- Assume a panel of  $G$  groups observed for  $T$  periods
- Assume treatment is assigned at the  $(g, t)$  level and is binary  $D_{g,t} \in \{0, 1\}$
- Assume **SUTVA** holds: Potential outcomes of group  $g$  only depend on treatments received by group  $g$

Let  $(d_1, \dots, d_T) \in \{0, 1\}^T$  be a particular sequence of treatments for group  $g$  in all periods  $t$ , then

$$Y_{g,t}(d_1, \dots, d_T)$$

will denote the associated potential outcomes

# Assumptions

To identify treatment effects, we will rely on **two** key assumptions:

- **No anticipation:** For all groups  $g$  and all possible treatment values  $(d_1, \dots, d_T) \in \{0, 1\}^T$

$$Y_{g,t}(d_1, \dots, d_T) = Y_{g,t}(d_1, \dots, d_t) \quad (\text{NA})$$

- **Parallel trends:** For all time periods  $t \geq 2$

$$E[Y_{g,t}(\mathbf{0}_t) - Y_{g,t-1}(\mathbf{0}_{t-1})] \quad (\text{PT})$$

does not vary across groups  $g$ .

# Assumptions

Sometimes, an additional assumption is made:

- **No dynamic effects:** For all groups  $g$  and all possible treatment values  $(d_1, \dots, d_T) \in \{0, 1\}^T$

$$Y_{g,t}(d_1, \dots, d_t) = Y_{g,t}(d_t) \quad (\text{ND})$$

In this case, **parallel trends** reduces to: For all time periods  $t \geq 2$

$$E[Y_{g,t}(0) - Y_{g,t-1}(0)] \quad (\text{PT})$$

does not vary across groups  $g$ .

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## Definition

Assume treatment is binary and that all groups  $g$  are treated at the same time – i.e., **no variation in treatment timing**

- Let  $T_g \in \{0, 1\}$  be an indicator for treatment groups
- Assume there is at least one treated group and one control group
- Let  $F \geq 2$  denote the time period at which all treatment groups become treated

Formally, this design is summarized as follows: For all groups  $g$

$$D_{g,t} = 1\{t \geq F\} T_g. \quad (\text{D1})$$

## TWFE is simple DiD

In design (D1), the TWFE estimator is a simple DiD estimator

- Let  $G_0$  and  $G_1$  denote the number of control and treatment groups, resp.
- Let  $T_0$  and  $T_1$  denote the number of control and treatment periods, resp.

It can be shown that

$$\beta_{TWFE} = \left( \frac{1}{G_1 T_1} \sum_{g: T_g=1, t \geq F}^T Y_{g,t} - \frac{1}{G_1 T_0} \sum_{g: T_g=1, t < F}^T Y_{g,t} \right) \quad (6)$$

$$- \left( \frac{1}{G_0 T_1} \sum_{g: T_g=0, t \geq F}^T Y_{g,t} - \frac{1}{G_0 T_0} \sum_{g: T_g=0, t < F}^T Y_{g,t} \right) \quad (7)$$

## TWFE is unbiased for ATT

A natural target parameter in design (D1) is the **Average Treatment Effect on the Treated (ATT)**:

$$\mathbf{ATT} = \frac{1}{G_1 T_1} \sum_{(g,t): D_{g,t}=1} E[Y_{g,t}(\mathbf{0}_{F-1}, \mathbf{1}_{t-F+1}) - Y_{g,t}(\mathbf{0}_t)].$$

If assumptions (NA) and (PT) hold, then the TWFE estimator is **unbiased** for the **ATT**:

$$E[\beta_{TWFE}] = \mathbf{ATT}.$$

## Dynamic effects and event-study designs

To estimate dynamic treatment effects in design (D1), researchers have often relied on two-way fixed effects **event-study** regressions:

$$\begin{aligned}
 Y_{g,t} = & \alpha_0 + \alpha_1 T_g + \sum_{t'=1, t' \neq F-1}^T \gamma_{t'} 1\{t = t'\} \\
 & + \sum_{\ell=-F+2, \ell \neq 0}^{T-F+1} \beta^{\ell} 1\{t = F-1 + \ell\} T_g + \varepsilon_{g,t}. \quad (\text{ES1})
 \end{aligned}$$

where  $1\{t = F-1 + \ell\} T_g$  are *relative-time* indicators equal to one **if** at period  $t$  group  $g$  has been treated for  $\ell$  periods.



## TWFE ES is simple DiD

The TWFE estimator in event-study setup (ES1) is again a simple DiD estimator:

$$\beta_{TWFE}^{\ell} = \frac{1}{G_1} \sum_{g: T_g=1}^T (Y_{g,F-1+\ell} - Y_{g,F-1}) - \frac{1}{G_0} \sum_{g: T_g=0}^T (Y_{g,F-1+\ell} - Y_{g,F-1}).$$

For  $\ell \leq -1$ ,  $\beta_{TWFE}^{\ell}$  is often referred to as a **pre-trend** or **placebo estimator**: Can be used to formally test (NA) and (PT)

## TWFE ES is unbiased for $ATT_\ell$

In design (D1), two-way fixed effects event-studies allow us to consider dynamic treatment effects –  $ATT_\ell$ :

$$ATT_\ell = \frac{1}{G_1} \sum_{g: T_g=1} E[Y_{g,F-1+\ell}(\mathbf{0}_{F-1}, \mathbf{1}_\ell) - Y_{g,F-1+\ell}(\mathbf{0}_{F-1})].$$

Again, if assumptions (NA) and (PT) hold, then for all periods  $\ell > 0$  the TWFE ES estimator is **unbiased** for the  $ATT_\ell$ :

$$E[\beta_{TWFE}^\ell] = ATT_\ell.$$

## TWFE ES allows to test NA and PT

Furthermore, if assumptions (NA) and (PT) hold, then for all periods  $\ell < 0$ :

$$E[\beta_{TWFE}^{\ell}] = 0. \quad (8)$$

Hence, the **placebo estimators** can be used to formally test the null of (NA) and (PT) **jointly** being satisfied

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## Staggered adoption

Assume treatment is binary **but** groups  $g$  may get treated at different time periods – i.e., **staggered treatment adoption**

- Let  $F_g$  denote the time period at which group  $g$  becomes treated
- Assume that for untreated groups  $F_g > T$
- Assume that all groups not yet treated at  $t = 1$ , do not all receive treatment in the same time period

Formally, in a staggered adoption design: For all groups  $g$

$$D_{g,t} = 1\{t \geq F_g\}. \quad (\text{D2})$$

## Failure of TWFE in staggered designs

If there is variation in treatment timing across groups  $g$  and if treatment effect may vary across time  $t$ , then

- $\beta_{TWFE}$  may be biased for the **ATT**
- $\beta_{TWFE}^\ell$  may be biased for the **ATT** <sub>$\ell$</sub>

Why? TWFE effectively estimates a **weighted sum** of treatment effects across all treated  $(g, t)$  cells with weights that may become **negative** if treatment effects vary across  $t$

These weights can be calculated in Stata with the `twfeweights` package

## Robust estimators

Most of recent lit. using TWFE (ES) regressions are embedded in more complex designs than the simple binary treatment, no variation in treatment timing design (D1) ...

In a staggered adoption design like (D2), estimators **robust** to heterogeneous effects across time **exist** – see, e.g., [Callaway and Sant'Anna \(2021\)](#) and [Sun and Abraham \(2021\)](#)

For an overview of (robust) estimators in more complex and general designs, see [de Chaisemartin and D'Haultfoeuille \(2023\)](#)

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## Bailey and Goodman-Bacon (2015)

What is the impact of providing access to primary care on longer-term health?

- Use the rollout of Community Health Centers (CHCs)
- CHCs can help lower **mortality among elderly** by providing accessible preventive care

Exploit the staggered adoption of CHCs across U.S. counties:

- *Our empirical strategy uses variation in when and where CHC programs were established to quantify their effects on mortality rates*
- Since CHCs are started in different counties in different time periods, effects are estimated in event-time – i.e., relative to initial rollout

# Bailey and Goodman-Bacon (2015)

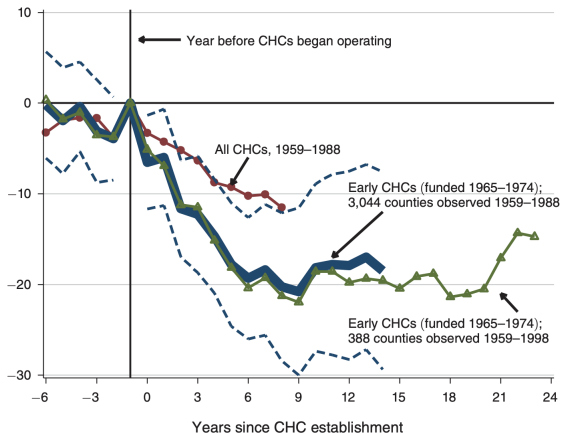
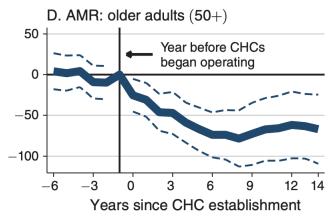
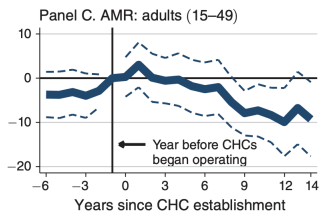
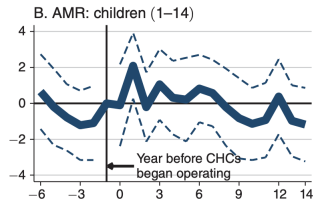
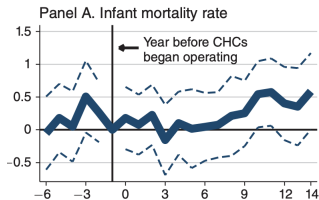


FIGURE 5. THE RELATIONSHIP BETWEEN COMMUNITY HEALTH CENTERS AND MORTALITY RATES

# Bailey and Goodman-Bacon (2015)



## References I

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## References II

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