

# **Political Methodology II**

## **Section: Difference in Differences**

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February 12, 2022

Stanford University

SOO Loose-ends

Difference-in-differences

General Exam Advice

Generalising to multiple time periods: Fixed Effects

Synthetic Control Methods

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## Regression as a type of propensity weighting (Angrist and Pischke, p.83)

Let's our estimand for the ATE comes from a regression of the form

$$\mathbb{E}[Y_i|D_i, X_i] = \alpha + X_i^T \beta + \delta_R D_i$$

where  $X_i$  is a vector of covariates and  $D_i$  is the treatment indicator.

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Now let's substitute in the definition of the propensity score function,  $p(X_i) = \mathbb{E}[D_i|X_i]$ .

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We can see that this estimand is equal to the weighted propensity score estimand:

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where  $\frac{p(X_i)(1-p(X_i))}{\mathbb{E}[p(X_i)(1-p(X_i))]}$  is the weight for observations with covariates  $X_i$ .

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$$\delta_{ATE} = \mathbb{E}\left[\frac{Y_i D_i}{p(X_i)} - \frac{Y_i(1 - D_i)}{1 - p(X_i)}\right]$$

When will these two coincide?

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When will these two coincide? Constant treatment effects across strata of  $X_i$ .

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When will these two coincide? Constant treatment effects across strata of  $X_i$ . Otherwise, OLS does not estimate ATE/ATT.

## OLS with heterogeneous treatment effects



In this case your estimator **or** is picking your estimand.





SOO Loose-ends

**Difference-in-differences**

General Exam Advice

Generalising to multiple time periods: Fixed Effects

Synthetic Control Methods

## DiD: Two Groups and Two Periods

Denote potential outcomes  $Y_{(d)}(t)$  for  $d \in \{0, 1\}$ ,  $t \in \{0, 1\}$

**Estimand (ATT in the 2nd period)**

$$\tau_{ATT} = E[Y_1(1) - Y_0(1)|D = 1]$$

	<b>Post-Period (T=1)</b>	<b>Pre-Period (T=0)</b>
Treated D=1	$E[Y_1(1) D = 1]$	$E[Y_0(0) D = 1]$
Control D=0	$E[Y_0(1) D = 0]$	$E[Y_0(0) D = 0]$

### **Problem**

*Missing potential outcome:  $E[Y_0(1)|D = 1]$ , ie. what is the average post-period outcome for the treated in the absence of the treatment?*

# Identification with Difference-in-Differences

## Identification Assumption (parallel trends)

$$E[Y_0(1) - Y_0(0)|D = 1] = E[Y_0(1) - Y_0(0)|D = 0]$$

## Identification Result

Given parallel trends the ATT is identified as:

$$\begin{aligned} E[Y_1(1) - Y_0(1)|D = 1] &= \left\{ E[Y(1)|D = 1] - E[Y(1)|D = 0] \right\} \\ &\quad - \left\{ E[Y(0)|D = 1] - E[Y(0)|D = 0] \right\} \end{aligned}$$

Implicit functional form assumption: Parallel trends in levels  $\neq$  Parallel trends in logs (growth rates). (cf Jensen's Inequality)

# Non-parametric Identification with Difference-in-Differences

Start with the estimand we want:

$$\tau_{ATT} = E[Y_1(1)|D = 1] - E[Y_0(1)|D = 1]$$

What is the missing data here?

# Non-parametric Identification with Difference-in-Differences

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What is the missing data here? The missing data is  $E[Y_0(1)|D = 1]$  (the control counterfactual in the post-period for the treated group). To obtain an estimate of this, we assume parallel trends:

$$\begin{aligned} E[Y_0(1) - Y_0(0)|D = 1] &= E[Y_0(1) - Y_0(0)|D = 0] \\ \implies E[Y_0(1)|D = 1] &= \underbrace{E[Y_0(0)|D = 1]}_{\text{Level at } t = 0} + \underbrace{E[Y_0(1) - Y_0(0)|D = 0]}_{\text{Trend for control group}} \end{aligned}$$

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Substitute the assumption in for the missing data and rearrange:

$$\begin{aligned} \tau_{ATT} &= E[Y_1(1)|D = 1] - E[Y_0(1)|D = 1] \\ &= E[Y_1(1)|D = 1] - E[Y_0(0)|D = 1] + E[Y_0(1)|D = 0] - E[Y_0(0)|D = 0] \\ &= \underbrace{(E[Y(1)|D = 1] - E[Y(0)|D = 1])}_{\text{Before after for treated}} - \underbrace{(E[Y(1)|D = 0] - E[Y(0)|D = 0])}_{\text{Before after for control}} \end{aligned}$$

This is just the difference of pre-post differences between the treated and control groups.

### Estimand (Sample Means: Panel)

$$\left\{ \frac{1}{N_1} \sum_{D_i=1} Y_i(1) - \frac{1}{N_0} \sum_{D_i=0} Y_i(1) \right\} - \left\{ \frac{1}{N_1} \sum_{D_i=1} Y_i(0) - \frac{1}{N_0} \sum_{D_i=0} Y_i(0) \right\}$$
$$= \left\{ \frac{1}{N_1} \sum_{D_i=1} \{Y_i(1) - Y_i(0)\} - \frac{1}{N_0} \sum_{D_i=0} \{Y_i(1) - Y_i(0)\} \right\},$$

where  $N_1$  and  $N_0$  are the number of treated and control units respectively.

This implies a fully saturated regression model with a two-way interaction:

$$E[Y_{igt}|g, t] = \beta_0 + \beta_1 \text{Treated}_g + \beta_2 \text{Post}_t + \beta_3 (\text{Treated}_g \times \text{Post}_t)$$

The subscripts are conventions to indicate the level of variation ( $i$  = Individual,  $g$  = Group,  $t$  = Time).



## Difference-in-Differences estimator

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The subscripts are conventions to indicate the level of variation ( $g = \text{Group}$ ,  $t = \text{Time}$ ). This translates to our two-by-two table as follows:

	Post-Period (T=1)	Pre-Period (T=0)	Pre/Post Diff.
Treated D=1	$\beta_0 + \beta_1 + \beta_2 + \beta_3$	$\beta_0 + \beta_1$	$\beta_2 + \beta_3$
Control D=0	$\beta_0 + \beta_2$	$\beta_0$	$\beta_2$
Treated/Control Diff.	$\beta_1 + \beta_3$	$\beta_1$	$\beta_3$

$\beta_3$  is the diff-in-diff estimate.

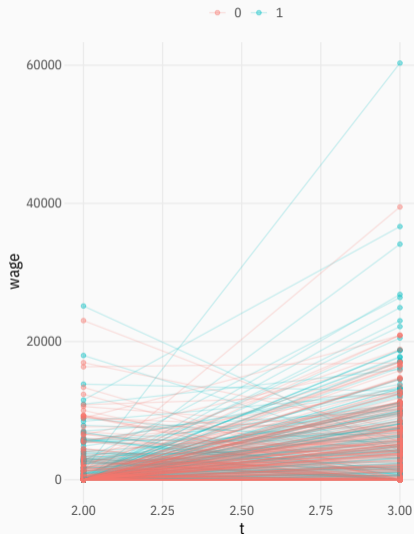
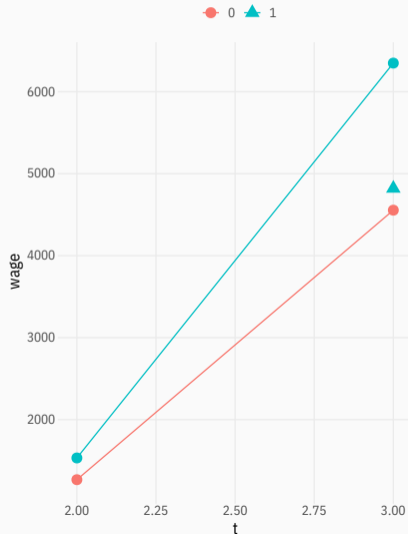
# HOW MUCH SHOULD WE TRUST DIFFERENCES-IN-DIFFERENCES ESTIMATES?\*

MARIANNE BERTRAND  
ESTHER DUFLO  
SENDHIL MULLAINATHAN

Standard advice: Cluster at least the unit level. More on this later.

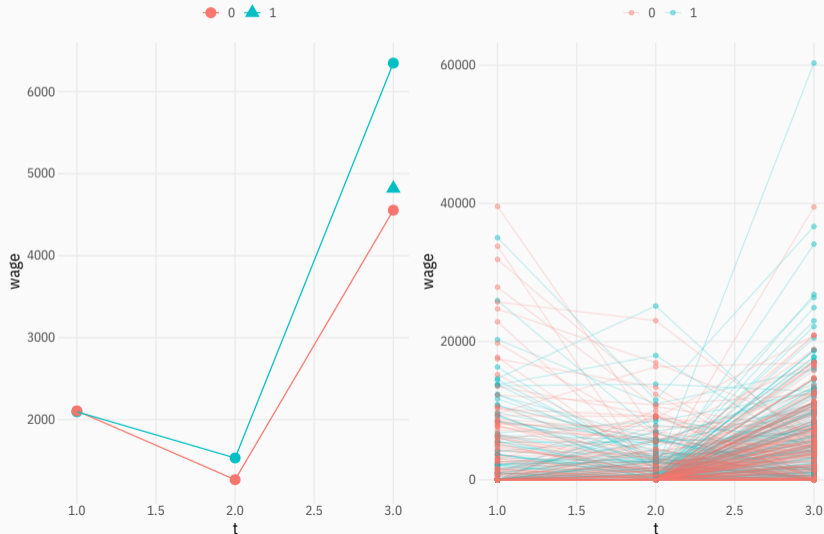
# Lalonde Experimental Sample: 2 time periods

## Lalonde: Experimental



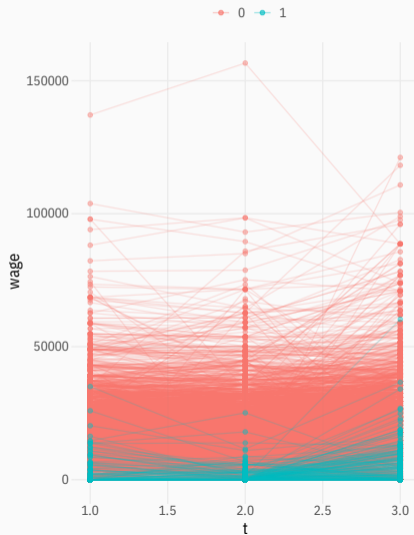
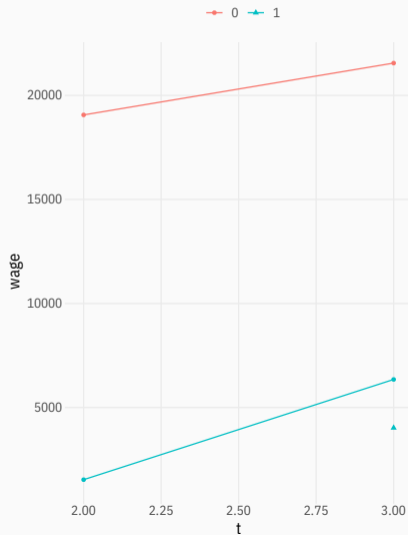
# Lalonde Experimental Sample: 3 time periods - Ashenfelter Dip

## Lalonde: Experimental



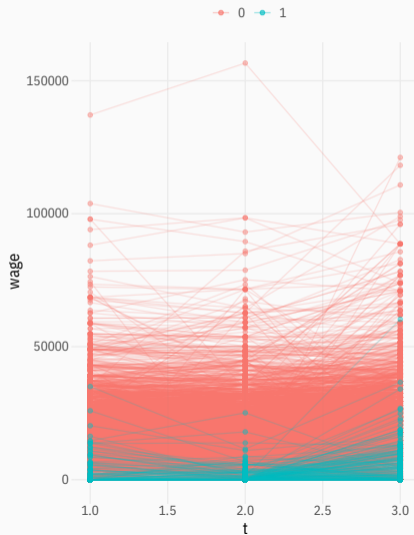
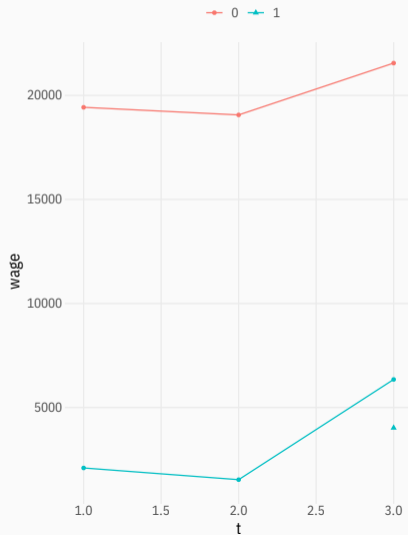
# Lalonde PSID: 2 time periods

## Lalonde: Observational



# Lalonde PSID: 3 time periods

## Lalonde: Observational



SOO Loose-ends

Difference-in-differences

**General Exam Advice**

Generalising to multiple time periods: Fixed Effects

Synthetic Control Methods

- Identify the relevant counterfactual
- What assumptions can we use to impute that counterfactual? Are those assumptions plausible?
- When writing identification results:
  - Start with observables (i.e.,  $E[Y_i | D_i = 1]$ , not  $E[Y_{1i} | D_i = 1]$ )
  - Manipulate the expressions to get to causal estimands
  - Explicitly note whenever you invoke an assumption. E.g., don't simply write  $E[Y_{1i} | X_i, D_i = 1] = E[Y_{1i} | X_i]$  without explaining what assumption justifies it.



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- When showing properties of estimators:
  - First write down an expression for the estimator
  - Then apply expectation, variance, etc.
  - Get to some population quantity, then your identification results can kick in

# Review Topics

- Power: intuition and calculation
- SOO assumptions
- OLS estimator
- Sensitivity analysis

SOO Loose-ends

Difference-in-differences

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Synthetic Control Methods

# Fixed Effects Regressions

- We often have access to panel data, wherein each individual  $i \in \{1, \dots, N\}$  is observed for  $T \geq 2$  time periods
- Stipulate following potential outcomes  $Y_{it}^{(d)}$ 
  - $\mathbb{E} [Y_{it}^0 | \alpha_i, t, D_{it}] = \alpha_i + \lambda_t$ 
    - $\alpha_i$  is a unit fixed-effect: each individual has an intercept  $\alpha_i$  - absorbs time-invariant unit-specific confounders
    - $\lambda_t$  is a time fixed-effect: each time period has an intercept  $\lambda_t$  - absorbs unit-invariant time-specific confounders
  - Suppose  $D_{it}$  is as-good as randomly assigned conditional on  $\alpha_i$
  - Stipulate constant, additive effect of treatment. Then,  $\mathbb{E} [Y_{it}^1] = \mathbb{E} [Y_{it}^0] + \tau$
- This motivates the popular **two-way fixed-effects** regression

$$Y_{it} = \tau D_{it} + \alpha_i + \gamma_t + \varepsilon_{it}$$

## Within Estimator

- With large datasets, estimating individual  $\alpha_i$ s can involve inverting a very large matrix
  - With short panels, the estimates of  $\alpha_i$ s are inconsistent anyway := incidental parameters problem (Neyman-Scott)
- Instead, we can use Frisch-Waugh-Lovell (again!) and partial out FEs
- Calculate individual averages of the 2wFE equation

$$\bar{Y}_i = \alpha_i + \bar{\lambda} + \tau \bar{D}_i + \bar{\varepsilon}$$

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- Calculate individual averages of the 2wFE equation

$$\bar{Y}_i = \alpha_i + \bar{\lambda} + \tau \bar{D}_i + \bar{\varepsilon}$$

- Subtract this from the FE equation

$$Y_{it} - \bar{Y}_i = \lambda_t - \bar{\lambda} + \tau(D_{it} - \bar{D}_i) + (\varepsilon_{it} - \bar{\varepsilon})$$

# Staggered Adoption, Treatment Reversals, and other complications

- Recall that we stipulated a **constant, additive treatment effect** and **treatment timing as-good-as-random (conditional on FEs)**
- Last 5 years of methods literature on panel data studies what happens when we relax these parametric assumptions
- Weird weights redux: 2WFE no longer consistent for ATT

What's Trending in Difference-in-Differences?

A Synthesis of the Recent Econometrics Literature

Jonathan Roth\* Pedro H. C. Sant'Anna<sup>†</sup> Alyssa Bilinski<sup>‡</sup> John Poe<sup>§</sup>

January 3, 2022

Two-Way Fixed Effects and Differences-in-Differences with  
Heterogeneous Treatment Effects: A Survey\*

Clément de Chaisemartin<sup>†</sup> Xavier D'Haultfoeuille<sup>‡</sup>

- review paper 1
- review paper 2

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**Synthetic Control Methods**



## Synthetic Control: Motivation

- We're sometimes interested in estimating a treatment effect where only a single unit is treated.
- In these cases, it's really important to estimate a good counterfactual for that particular unit – not just on average as in the matching or traditional diff-in-diff cases.
- The synthetic control method introduced by Abadie, Diamond, and Hainmueller (2010) is useful when there are many pre-treatment outcome observations, and perhaps relatively few untreated units.
- The intuition is to create a “synthetic control” that is a weighted average of control units. We pick the weights so that the pre-treatment outcome of the synthetic control looks similar to the pre-treatment outcomes of the treated unit.

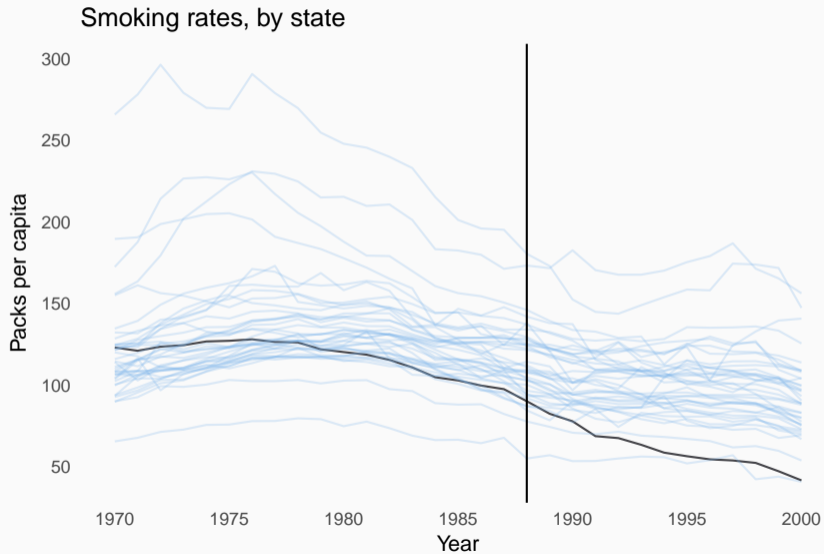
# Smoking Data

ADH (2010) study the effect on smoking rates of an increase in the tobacco tax in California in 1988.

```
load("synth.rdata")
head(synth.long[c(1:3, 1206:1209), c("statename", "year", "smoking")])
```

```
##      statename year smoking
## 1      Alabama 1970   89.8
## 2      Arkansas 1970  100.3
## 3      Colorado 1970  124.8
## 1206 West Virginia 2000  107.9
## 1207 Wisconsin 2000   80.1
## 1208 Wyoming   2000   90.5
```

# Smoking Data



The problem is to impute what California's level of smoking would have been in the post-treatment period if Proposition 99 hadn't been passed.

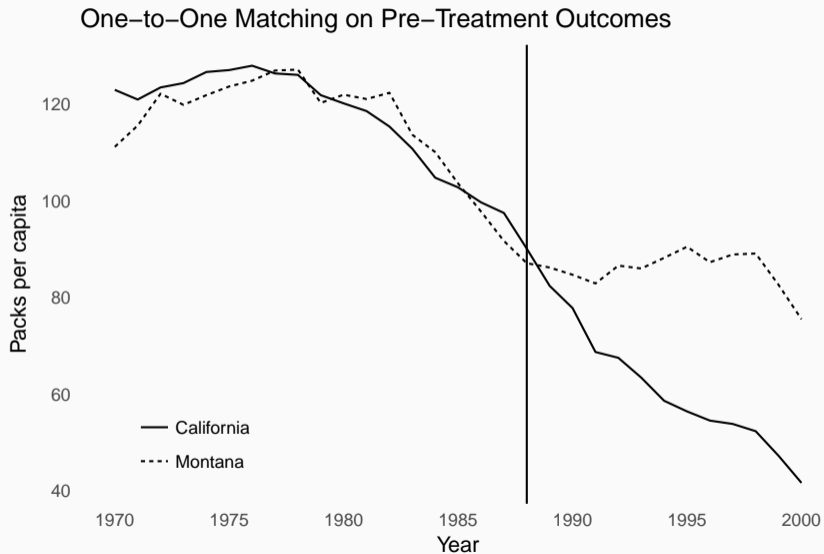
Potential strategies:

- Matching on pre-treatment covariates (possibly including lagged outcomes).
- Difference-in-differences.
- Regression.
- Synthetic control.

## Matching on Pre-Treatment Outcomes

```
genmatch = GenMatch(synth$statename = "California",  
                    synth[, paste0("smoking-", 1970:1988)])  
matched.unit = genmatch$matches[,2]  
match.statename = synth[matched.unit, statename]  
out = Match(Y = synth$smoking_1996, Tr = synth$statename == "California",  
            X = synth[, paste0("smoking-", 1970:1988), with=F],  
            Weight.matrix = genmatch, estimand = "ATT")
```

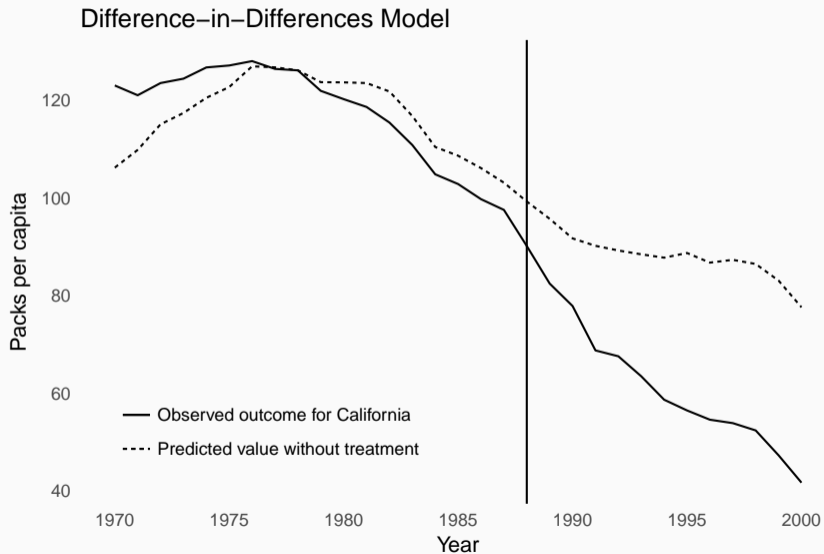
# Matching on Pre-Treatment Outcomes



# Difference-in-Differences

```
synth.long = synth.long %>%  
  mutate(treatment = statename == 'California' & year ≥ 1989)  
did = lm(smoking ~ treatment + statename + factor(year), synth.long)  
calif = synth.long[synth.long$statename == 'California', ]  
calif$treatment = F  
counterfactuals = predict(did, newdata = calif)
```

# Difference-in-Differences





## Synthetic Control Method

Synthetic control setup: suppose unit  $i = 0$  is treated after time  $T_0$ , and units  $i = 1, \dots, n$  are never treated. We observe outcomes for time periods  $t = 1 < T_0 \leq T$ . In potential outcomes notation, we observe:

$$Y_{0,t}(0) \quad \text{for } t = 1, \dots, T_0 - 1$$

$$Y_{0,t}(1) \quad \text{for } t = T_0, \dots, T$$

$$Y_{i,t}(0) \quad \text{for } t = 1, \dots, T \text{ and } i = 1, \dots, n$$

We want to estimate  $Y_{0,t}(1) - Y_{0,t}(0)$  for time periods  $t \geq T_0$ , but we can't observe  $Y_{0,t}(0)$  after  $T_0$ .

# What We Observe

The matrix we observe is:

$$\mathbf{Y}^{obs} = \begin{pmatrix} \mathbf{Y}_{treat,pre}^{obs} & \mathbf{Y}_{control,pre}^{obs} \\ \mathbf{Y}_{treat,post}^{obs} & \mathbf{Y}_{control,post}^{obs} \end{pmatrix} = \begin{pmatrix} \mathbf{Y}_{treat,pre}(0) & \mathbf{Y}_{control,pre}(0) \\ \mathbf{Y}_{treat,post}(1) & \mathbf{Y}_{control,post}(0) \end{pmatrix}$$

# What We Observe

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To estimate the ATT we need  $\mathbf{Y}_{treat,post}(0)$ . We only observe:

$$\mathbf{Y}(0) = \begin{pmatrix} \mathbf{Y}_{treat,pre}(0) & \mathbf{Y}_{control,pre}(0) \\ ? & \mathbf{Y}_{control,post}(0) \end{pmatrix}$$

## Synthetic Control Method

The synthetic control estimator imputes  $Y_{0,t}(0)$  as a weighted average of the observed outcomes for the control units (plus possibly an intercept shift):

$$\hat{Y}_{0,t}(0) = \mu + \sum_{i=1}^n \omega_i Y_{i,t}$$

for some weight vector  $\omega = (\omega_1, \dots, \omega_n)$  and intercept  $\mu$ . The parameters  $(\mu, \omega)$  define the “synthetic control” for the treated unit.

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The parameters are typically picked by minimizing the squared distance between the synthetic control’s pre-treatment outcomes and the treatment unit’s pre-treatment outcomes:

$$\omega^* = \arg \min_{(\mu, \omega_1, \dots, \omega_n)} \sum_{t=1}^{T_0-1} \left( Y_{0,t} - \mu - \sum_{i=1}^n \omega_i Y_{i,t} \right)^2$$

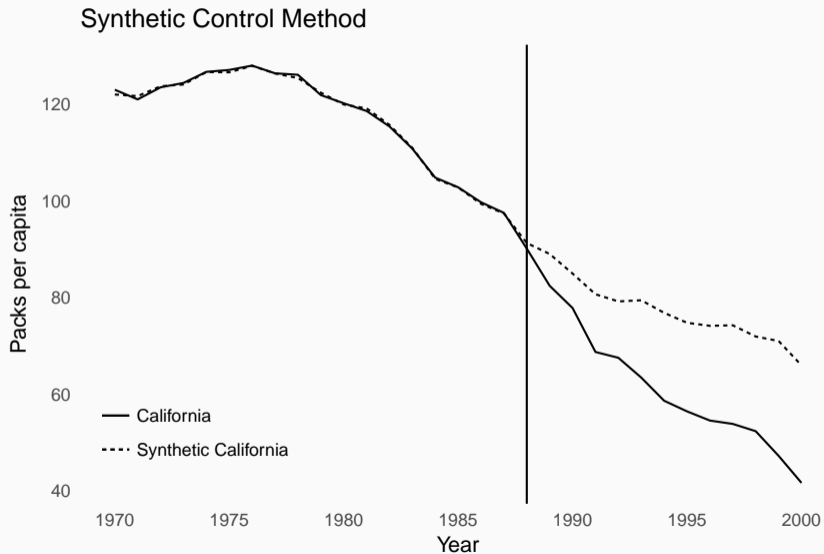
The natural method would be to estimate  $(\mu, \omega)$  using OLS, but if  $n > T_0 - 1$  it requires additional constraints (e.g., weights summing to 1 or regularization).

# Synthetic Control Method: Estimation Using glmnet

```
library(glmnet)
control.pre = synth[synth$state≠3, paste0("smoking_", 1970:1988)] %>%
  as.matrix %>% t
control.full = synth[synth$state≠3, paste0("smoking_", 1970:2000)] %>%
  as.matrix %>% t
treat.pre = synth[synth$state==3, paste0("smoking_", 1970:1988)] %>%
  as.matrix %>% t

# estimate weights and intercept using LASSO
weightsout = cv.glmnet(x = control.pre, y = treat.pre)
predictions = predict(weightsout, newx = control.full, s = "lambda.min")
calif$scm.pred = as.numeric(predictions)
```

# Synthetic Control Method: Results



## Synthetic Control Method: Uncertainty

How to assess uncertainty with synthetic control methods? There's only one treated unit so asymptotics are not helpful. Instead, ADH suggest a procedure like Bertrand, Duflo, and Mullainathan's placebo laws to assess what the null distribution of the SCM estimator looks like.