Difference-in-Differences Models

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Agenda for today

- What are difference-in-differences models?
- 2. Key «ingredients»
- 3. A textbook example (also using Stata)
- 4. Testing assumptions
- Potential extensions on evaluation models

Key readings:

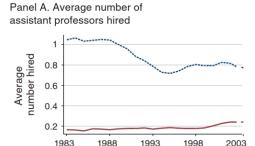
- Angrist and Pischke, «Mostly Harmless Econometrics: An Empiricist's Companion»,
 Chapter 5
- Or: Scott Cunningham, «The Mixtape», <u>Chapter 9, Difference-in-Differences</u>



Matching methods vs. Difference-in-Differences

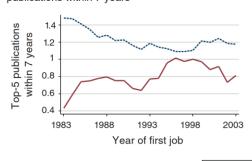
	Matching Models (PSM)	Difference-in-Differences (DiD)
When	We need to assess the effect of a "treatment" (e.g., choice, policy)	We need to assess the effect of a "treatment" (e.g., choice, policy)
Problem	T & C groups are very different ("selection on observables"); cross-section, no panel data	Requires data "before" and "after" the treatment + "treated" & "control" units. Is the "treatment" exogenous (e.g. natural experiment?)
Stata commands	teffects psmatch, tebalance, teffects overlap	Regular OLS regression or panel regression (though there are automatic Stata commands you can explore)
Key tests	Balancing and overlapping conditions (quality checks after PSM)	Parallel trends assumption (before the "intervention"), i.e. T and C's outcomes trajectories are "parallel"
Attention!	T & C only matched on observable characteristics. If unobservables matter, PSM does not provide causal effects → IV or panel DiD	If parallel trends assumption is violated, we may need to construct matched samples first and then run a DiD regression
First stage	Probit predicting assignment to "treatment" (X)	Obtain matched samples of treated and control units (via PSM/CEM etc)

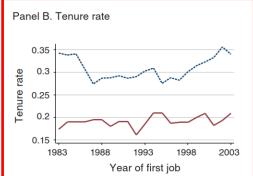
Example 1: Equal but Inequitable



Year of first job

Panel C. Average number of top-5 publications within 7 years





Panel D. Average number of non-top-5 publications within 7 years

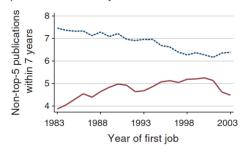


FIGURE 2. TRENDS IN TOP-50 ECONOMICS DEPARTMENTS

Women

Tenure clock stopping policies:

Assistant professors are allowed to stop their tenure clock for 1 year after childbirth or adoption. No research is expected during this time.

Do **gender-neutral** clock stopping (GNCS) policies level the playing field in terms of tenure rates (i.e. do the effects differ for M and W and affect the gender gap)?

Top-50 Econ departments in the US; some universities adopted them (at different points in time), others didn't.

Example 1: Equal but Inequitable

	Total effects (1)	Male-female (2)
Panel A. Policy effects years 0–3		
Men FOCS	-0.008 (0.067)	-0.181 (0.140)
Women FOCS	0.172 (0.140)	
Men GNCS	0.051 (0.079)	0.068 (0.145)
Women GNCS	-0.017 (0.107)	
Panel B. Policy effects years 4+		
Men FOCS	0.002 (0.075)	-0.047 (0.128)
Women FOCS	0.049 (<u>0.101</u>)	0.00
Men GNCS	$0.176 \\ (0.083)$	0.370 (0.146)
Women GNCS	$ \begin{array}{c} -0.194 \\ (0.106) \end{array} $	
Sample size	1,392	

DV = 1 if the individual makes tenure, 0 if not. GNCS = Gender Neutral tenure clock stopping policy/FOCS = Female only. Second column shows difference in the male and female coefficients for each policy type. Std. errors clustered at policy-university level. Examples of controls: time-varying university characteristics, PhD rank, having done a postdoc... Gender-specific university FE included.

- a) Men whose first job was at a top-50 university with a gender-neutral tenure clock stopping policy in place for 4+ years have a 17.6 percentage point tenure rate advantage over men at the same university prior to the implementation of any policy (diff: male T & C)
- b) Women, in turn, are 19.4 percentage points less likely to get tenure relative to other women hired by the same university prior to the clock stopping policy (diff: female T & C)
- c) This increased the gender gap between men and women by 37 percentage points: women are even less likely to get tenure compared to men after the introduction of GNCS policies. (diff-in-diff)



Example 2: Inventor death and innovation

If the collaboration between two patent inventors were to exogenously end, would this have a significant and long-lasting impact on the career, compensation, and patents of co-inventors? Or are co-inventors easily substituted for, beyond shortterm disruption of ongoing work?

Data

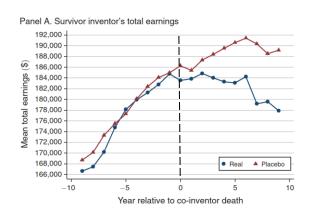
- USPTO patents data and Treasury administrative tax data
- Some inventors die suddenly before or at the age of 60 (4,714 inventors):
 exogenous shock in collaborative networks
- Compare inventors whose co-inventors did not pass away but who are
 otherwise similar to inventors who experienced the premature death of a coinventor (i.e. matched sample of inventors with and without loss of a
 team member treated vs control groups)

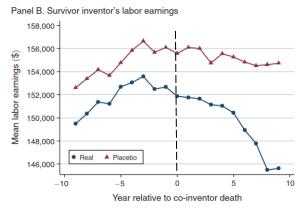


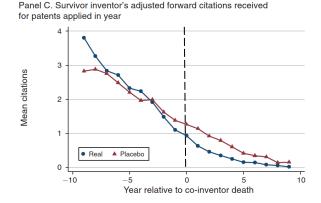
Example 2: Inventor death and innovation

Ending a collaboration causes a large and long-lasting decline in an inventor's:

- labor earnings (-3.8 percent after 8 years)
- total earnings (-4 percent after 8 years)
- citation-weighted patents (-15 percent after 8 years)







Note: "control inventors" are exactly matched to "real inventors" in age, year, and total nr of patent applications at the time of (real/control) death to secure parallel trends prior to the "shock"

Jaravel, X., Petkova, N., & Bell, A. (2018). Team-specific capital and innovation. *American Economic Review*, 108(4-5), 1034-73.

Example 3: Why Marathons Can Be Deadly

Large marathons frequently involve widespread road closures and infrastructure disruptions, which may create delays in emergency care for individuals with acute medical conditions who live in proximity to marathon routes ("treated" by this exogenous shock).

Data

- Hospitalizations for acute myocardial infarction or cardiac arrest (age ≥ 65)
- 11 U.S. cities that hosted marathons (2002-2012)

Summary here:

ns-running-heart-attack/

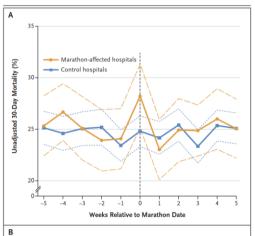
- Mortality of those hospitalized the day of the marathon vs. those hospitalized
 - in the same week day but 5 weeks bef/after the marathon
 - In the same day but in surrounding ZIP codes unaffected by the marathon

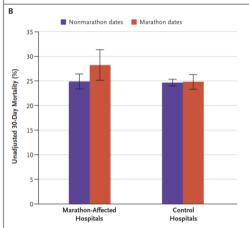




https://time.com/4736467/maratho

Example 3: Why Marathons Can Be Deadly

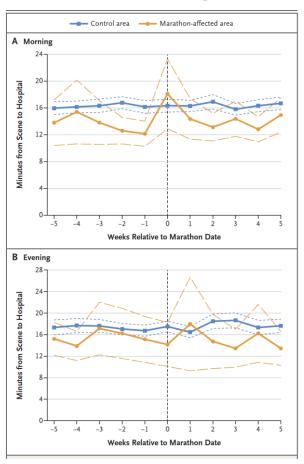




People who were admitted to marathon-affected hospitals on marathon dates had:

- longer ambulance transport times before noon (4.4 minutes longer)
- higher 30-day mortality than beneficiaries who were hospitalized on non-marathon dates
- higher 30-day mortality than those who were hospitalized on the same day as the marathon but in unaffected surrounding ZIP code areas

Jena, A. B., Mann, N. C., Wedlund, L. N., & Olenski, A. (2017). Delays in emergency care and mortality during major US marathons. *New England Journal of Medicine*, *376*(15), 1441-1450.



«Parallel worlds»: key ingredients

Time

The "**treatment**" – i.e., policy change, intervention, event – takes place at a certain point in time (pooled cross-section or panel data)

Policy change or treatment

We identify (at least one) **before** and **after period** with respect to the treatment

Comparison groups

Treated group receives the intervention or is subject to the policy change only in the post-period

Control group is not affected by the treatment

Fixed factors

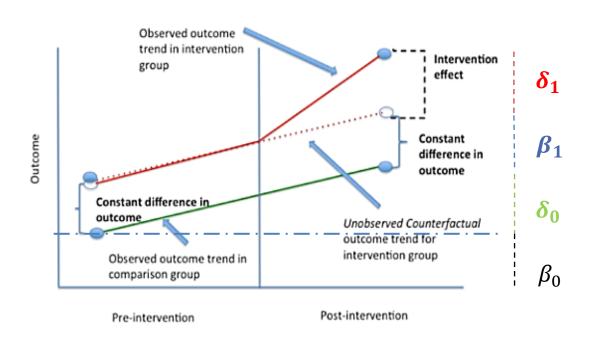
Assume that important factors associated with the outcome Y are fixed during the pre- and post-periods

Time invariant factors

If observed, we can control for those factors that could affect trends and vary over time (parallel trends or constant bias)

More formally, with a textbook example:

 $y = \beta_0 + \beta_1 dTreated + \delta_0 dPost + \delta_1 dTreated \cdot dPost + controls + u$



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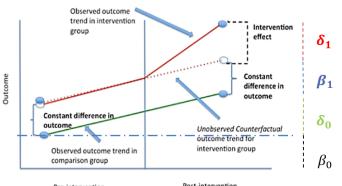
More formally, with a textbook example:

$$y = \beta_0 + \beta_1 dTreated + \delta_0 dPost + \delta_1 dTreated \cdot dPost + controls + u$$

	Before	After	After-Before
Control	eta_0	$\beta_0 + \delta_0$	δ_0
Treated	$\beta_0 + \beta_1$	$\beta_0 + \delta_0 + \beta_1 + \frac{\delta_1}{\delta_1}$	$\delta_0 + \delta_1$
Treated-Control	eta_1	$\beta_1 + \delta_1$	δ_1

 δ_1 is the "average treatment effect" (ATE)

$$\widehat{\delta_1} = (\bar{y}_{2,T} - \bar{y}_{2,C}) - (\bar{y}_{1,T} - \bar{y}_{1,C})$$



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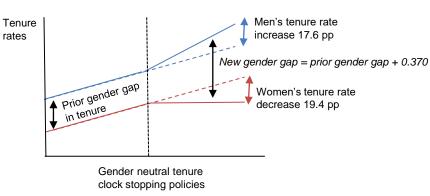
Recalling the tenure-gender example:

Recall slide 5

	After policy change in tenure rates
Men	0.176
Women	-0.194
Gender gap	0.370

 δ_1 is the "average treatment effect" (ATE)

$$\widehat{\delta_1} = 0.176 - (-0.194) = 0.370$$



DiD Assumptions

COMMON TRENDS OR CONSTANT BIAS	Both treatment and control groups would have the same trends over time (possibly conditioning for other factors) if the intervention had not happened. Treated and control groups are not equivalent, but their difference remains constant over time.			
RANDOM ASSIGNMENT INTO TREATMENT	The policy/intervention must "naturally" affect certain group of subjects (firms, persons, families), but not all. Alternatively, the treatment may affect all but its effects may differ across groups (e.g. slide 5). Manipulation is the key: "No causation without (explicit, natural, or whatever) manipulation"			
EXOGENEITY	The covariates X are not influenced by the treatment.			
SUTVA	No interference (spillovers/externalities) and variation in treatment among the groups.			



Example

- Introduction of a minimum wage policy (Card and Krueger, 1994)
- In a nutshell: in theory, a firm makes hiring decisions based on wages and the contribution of employees to revenue
- In a perfectly competitive market, a higher minimum wage implies that firms will demand fewer workers (or hours worked)
- Thus, a policy that helps those who can get jobs at the higher wage may harm some other workers, who won't find employment because of the higher minimum wage

Authors use a dramatic change in the New Jersey state minimum wage to test whether this is true.



Example

- In 1992, New Jersey raised the state minimum wage by about 19% from \$4.25 to \$5.05
- Card and Krueger (1994) obtained data from February 1992 ("before" or pre-period) and November 1992 ("after" or post-period) from fastfood chains*, which usually pay minimum wages (TREATED)
- They collected data from similar fast-food restaurants in eastern Pennsylvania – just across the Delaware river –, which did not change the minimum wage (\$4.25) (<u>CONTROL</u>)

We have the key elements of a basic two-period DiD:

- observations over time (both pre- and post-treatment)
- exogeneous and random treatment
- policy change applies only to treated group in the post-period
- the control group is not exposed to the experiment during the time





Data structure

For each store (id), we have two observations (balanced two-year panel). Some are treated, some are in the control group; both have 2 periods

. list id fte treated post in 1/10, sep(2) nolabel

	id	fte	treated	post
1.	1	16	1	0
2.	1	20	1	1
3.	2	10	1	0
4.	2	7.5	1	1
5.	3	6	1	0
6.	3	4	1	1
7.	4	10	1	0
8.	4	5	1	1
9.	5	5	1	0
10.	5	10	1	1

. d id treated	. d id treated post fte chain						
variable name	storage type	display format	value label	variable label			
id treated post fte chain	float float float float float	%9.0g %9.0g %9.0g %9.0g %9.0g	treated post	Restaurant ID NJ = 1; PA = 0 Feb.92 = 0; Nov. 1992 = 1 Output: Full Time Employees Burger King = 1; KFC = 2; Roys = 3; Wendy's = 4			

We want to estimate the **causal effect of** *x* (*treated*) **on** *y* (*fte*), keeping other things equal.



Interpretation of the results

- Restaurants in PA employed on average 10 FTEs before the policy change (β_0)
- Restaurants in NJ employed on average 2.6 fewer FTEs than PA before the policy change (β₁) – significant at 5%
- There is no average significant effect (at the 5% significance level) for restaurants in PA on number of FTEs after the introduction of the policy (δ_0)
- The average number of FTEs in NJ increased by 3,44 FTEs units after the policy change with respect to PA – in other words, due to the increase in minimum wages (δ₁)



 $y = \beta_0 + \beta_1 dTreated + \delta_0 dTime + \delta_1 dTreated \cdot dTime + controls + u$

. reg fte i.treated##i.post, robust

Sinear regression	Number of obs	=	784
	F(3, 780)	=	1.56
	Prob > F	=	0.1970
	R-squared	=	0.0084
	Root MSE	=	8.3213

fte	Coef.	Robust Std. Err.	t	P> t	[95% Conf.	. Interval]
treated NJ 1.post	-2.6006 -2.743421	1.319187 1.578217	-1.97 -1.74	0.049 0.083	-5.190177 -5.841476	0110226 .354634
treated#post NJ#1	3.442788	1.700103	2.03	0.043	.1054694	6.780107
_cons	10.31579	1.239793	8.32	0.000	7.882063	12.74952

	Before	After	After-Before
Control	eta_0	$\beta_0 + \delta_0$	δ_0
Treated	$\beta_0 + \beta_1$	$\beta_0 + \delta_0 + \beta_1 + \delta_1$	$\delta_0 + \delta_1$
Treated-Control	eta_1	$\beta_1 + \delta_1$	δ_1

Estimate and compare means

. * Treated
. sum fte if treated ==1 & post==1

fte	316	8.414557	7.870619	0	40
Variable	Obs	Mean	Std. Dev.	Min	Max

. scalar y_tpost = r(mean)

. sum fte if treated ==1 & post==0

fte	316	7.71519	8.004734	0	60
Variable	Obs	Mean	Std. Dev.	Min	Max

. scalar y_tpre = r(mean)

. * Control

. sum fte if treated ==0 & post==1

fte	76	7.572368	8.548179	0	35
Variable	Obs	Mean	Std. Dev.	Min	Max

. scalar y cpost = r(mean)

. sum fte if treated ==0 & post==0

				Max
fte	76	10.31579	10.85229	50

. scalar y_cpre = r(mean)



 δ_1 is the "average treatment effect" (ATE)

•
$$\widehat{\delta}_1 = (\bar{y}_{2,T} - \bar{y}_{2,C}) - (\bar{y}_{1,T} - \bar{y}_{1,C})$$



- . * DiD estimator
- . di y_tpost y_tpre (y_cpost y_cpre)
- 3.4427881

Adding controls

- Adding control variables reduces the residual variance, which in turn lowers the standard error of the regression estimates.
- If the treatment is really random, the point estimate should not change by adding more controls. Often this is seen as a robustness check on the claim of random assignment.
- Here, we are comparing the same restaurants before and after, so chain type couldn't affect the difference-indifferences because it is not changing over time.



Linear regress	ion			Number o	f obs	=	784
ninear regress	51011			F(3, 780		=	1.56
				Prob > F		=	0.1970
				R-square		-	0.0084
				Root MSE		=	8.3213
		Robust					
fte	Coef.	Std. Err.	t	P> t	[95%	Conf.	Interval]
treated							
NJ	-2.6006	1.319187	-1.97	0.049	-5.190	0177	0110226
1.post	-2.743421	1.578217	-1.74	0.083	-5.84	1476	.354634
reated#post							
NJ#1	3.442788	1.700103	2.03	0.043	.105	4694	6.780107
cons	10.31579	1.239793	8.32	0.000	7.882	2063	12.74952

. reg fte i.t	reaced##1.pos	c I.chain, I	obust				
Linear regres:	sion			Number o	f obs	=	784
_				F(6, 777)	-	12.19
				Prob > F		=	0.0000
				R-square	d	=	0.0607
				Root MSE		-	8.1144
		Robust					
fte	Coef.	Std. Err.	t	P> t	[95%	Conf.	Interval]
treated							
NJ	-2.30713	1.258303	-1.83	0.067	-4.77	7207	.1629459
1.post	-2.743421	1.509528	-1.82	0.070	-5.70	6658	.2198161
treated#post							
NJ#1	3.442788	1.632124	2.11	0.035	.238	8929	6.646683
chain							
KFC	-5.123389	.6552841	-7.82	0.000	-6.40		-3.837052
Roys	-1.690295	.7306062	-2.31	0.021	-3.12	4491	2560989
Wendy's	-1.00196	1.05185	-0.95	0.341	-3.06	6764	1.062845
_cons	11.67423	1.289992	9.05	0.000	9.14	1944	14.20651

Adding unit and time FEs

So far, we have seen the standard 2x2 DiD regression model:

$$y = \beta_0 + \beta_1 dTreated + \delta_0 dPost + \delta_1 dTreated \cdot dPost + u$$

We can extend the model adding unit (λ_i) and time (μ_t) fixed effects

$$y = \beta_0 + \beta_1 dTreated + \delta_0 dPost + \delta_1 dTreated \cdot dPost + \lambda_i + \mu_t + u$$

 λ_i captures time-invariant characteristics, including those that could affect self-selection (or assignment) into the treatment/program

 μ_t identifies time-variant characteristics, regardless of the group the individual belongs to

If the treatment is really random, the point estimate should not change by adding FEs.



Adding control variables

We can also control for variables that we think could affect the evolution of the trends between treatment and control groups (remember, variables that affect the difference between trends):

$$y = \beta_0 + \beta_1 dTreated + \delta_0 dPost + \delta_1 dTreated \cdot dPost + \theta X_{it} + \lambda_i + \mu_t + u$$

Control variables can significantly affect the "treatment effect"!

If some variables in X do not change over time, they won't affect the DiD estimator



Checking assumptions I



COMMON TRENDS OR CONSTANT BIAS

RANDOM ASSIGNMENT INTO TREATMENT

EXOGENEITY

SUTVA

There are some "exclusion restrictions" that you need to discuss:

For example, exogeneity or the SUTVA assumption

When is the SUTVA assumption violated?

Whenever it is not possible to argue that the effect of the treatment affects only treated individuals (i.e., ∄ externalities or spillovers). For example, there are interactions between individuals in the treatment and in the control group.

What about exogeneity?

The classical problem of OLS biased estimates due to various sources of endogeneity potentially occurring together so that sometimes it is not easy to distinguish between them.



Checking assumptions II

Other assumptions can be tested with data, or at least you can see if the data seems consistent with an assumption, even though it may not guarantee that the assumption is valid

- Graphically showing that trends are parallel is a good first step, but we can test the assumption as well
- We will see adjusted and unadjusted plots sometimes we may need to adjust for factors that affect the difference in trends



DiD vs. Experiments

In order to mimic a randomized controlled trial, we need a "natural" or "quasi-experiment" (exogeneous and unforeseen!) to reveal the causal effect of the "policy"

But ... What if the selection into treatment is not random?

We can relax this assumption with Conditional Independence Assumption

Or in other words:

once you control for X, being treated is "as good as random"



Recap so far

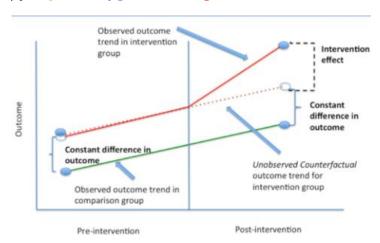
- To address your research question, consider the randomized case (i.e. the "ideal experiment") as an hypothetical benchmark.
- This is the experiment you would like to run, if you could!
- Think about your problem in terms of the potential outcome framework and then check if the assignment to the treatment is any close to the ideal experiment (it must be independent of the potential outcomes).
- Analyze the source of variation of the assignment to treatment. This is crucial
 and it must be exogenous (even conditional on observables, CIA).



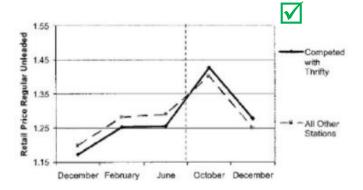


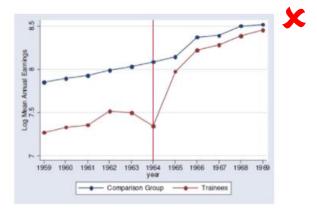
Parallel trends assumption in DiD

 $y = \beta_0 + \delta_0 dTime + \beta_1 dTreated + \delta_1 dTreated \cdot dTime + controls + u$



In the absence of treatment, the difference between T and C is constant over time. Visual inspection can be useful.





Stata exercise: Prison rates

We use the dataset "prison" (from J. Wooldridge).

Assume a policy change lowering prison sentences for some states (state = 17-51), but not for other states.

Assume that this policy started in 1986.

. d pris treated post did state year

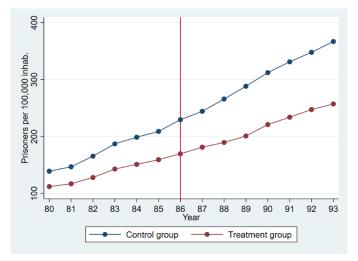
variable name	storage type	display format	value label	variable label
pris	float	%9.0g		prison pop. per 100,000
treated	float	%9.0g		<pre>0 = Control states; 1 = Treated States</pre>
post	float	%9.0g		<pre>0 = Before treatment; 1 = After Treatment</pre>
did	float	%9.0g		DiD estimator
state	byte	%9.0g		alphabetical; DC = 9
year	byte	%9.0g		80 to 93



Parallel trends: plot average values

```
bysort year: egen pris0=mean(pris) if treated==0
lab var pris0 "Control group"
bysort year: egen pris1=mean(pris) if treated==1
lab var pris1 "Treatment group"

twoway (connected pris0 pris1 year), xline(86) ytitle("Prisoners per 100,000 inhab.") ///
xtitle("Year") xlab(80(1)93)
graph export "trends1.png", as(png) replace
```





Parallel trends: same plot with *margins*

We can also plot the same graph using the *margins* command.

First, we would run a saturated model with dummy variables for each year, another dummy for "treated", and each of their interactions.

reg pris i.year##i.treated

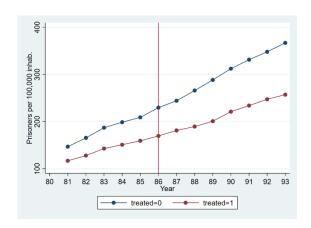
This approach is very helpful when we relax the assumption and we want to test for parallel trends conditional on observables!

71	er of obs =		MS	df	SS	Source
7.5	686) =					
0.000			106894.021	27	2886138.58	Model
0.229	ared =		14098.8798	686	9671831.57	Residual
0.199	R-squared =					
118.7	MSE =	Root	17612.8614	713	12557970.1	Total
Interval	[95% Conf.	P> t	t	Std. Err.	Coef.	pris
						year
90.327	-74.52338	0.851	0.19	41.98047	7.902258	81
108.969	-55.88218	0.527		41.98047	26.54346	82
130.567	-34.28386	0.252	1.15	41.98047	48.14177	83
142.180	-22.67054	0.155		41.98047	59.75509	84
152.397	-12.4541	0.096	1.67	41.98047	69.97154	85
172.988	8.137071	0.031		41.98047	90.56271	86
187.625	22.77434	0.012		41.98047	105.2	87
209.342	44.49122	0.003		41.98047	126.9169	88
231.732	66.88103	0.000		41.98047	149.3067	89
255.621	90.7699	0.000		41.98047	173.1955	90
274.64	109.7907	0.000		41.98047	192.2164	91
291.38	126.5368	0.000		41.98047	208.9624	92
310.35	145.5017	0.000	5.43	41.98047	227.9274	93
43.3978	-97.3133	0.452	-0.75	35.83302	-26.95774	1.treated
						year#treated
96.5684	-102.4271	0.954		50.67554	-2.929324	81 1
88.9385	-110.057	0.835		50.67554	-10.55925	82 1
82.2497	-116.7458	0.734		50.67554	-17.24801	83 1
78.7743	-120.2212	0.683		50.67554	-20.72339	84 1
76.8307	-122.1648	0.655		50.67554	-22.66707	85 1
66.5365	-132.459	0.516		50.67554	-32.96124	86 1
63.5216	-135.4739	0.478		50.67554	-35.97616	87 1
50.0570	-148.9385	0.330		50.67554	-49.44072	88 1
39.0391	-159.9564	0.233		50.67554	-60.45866	89 1
35.3380	-163.6575	0.206		50.67554	-64.1597	90 1
29.2014	-169.7941	0.166		50.67554	-70.29631	91 1
25.9020	-173.0935	0.147		50.67554	-73.59571	92 1
16.6725	-182.323	0.103	-1.63	50.67554	-82.8252	93 1
197.237	80.67003	0.000	4.68	29.68468	138.9538	cons



Parallel trends: same plot with *margins*

margins treated, at(year=(81(1)93)) vsquish
marginsplot, noci xline(86) xlab(80(1)93) ytitle("Prisoners per 100,000 inhab.")
xtitle("Year") title("")



It looks the same as the previous one!



Adjusted pre				Number	of obs =	71
Model VCE	: OLS					
Expression	: Linear pred	iction, pred	ict()			
1at	: year	-	81			
2at	: year	-	82			
	: year	=	83			
	: year	-	84			
5at	: year	=	85			
6at	: year	-	86			
7at	: year	-	87			
8at	: year	-	88			
9at	: year	=	89			
	: year	=	90			
11at	: year	-	91			
12at	: year	=	92			
13at	: year	-	93			
		Delta-method				
	Margin	Std. Err.	t	P> t	[95% Conf.	Interva
at#treated						
1 0	146.856	29.68468	4.95	0.000	88.57228	205.13
1 1	116.9689	20.0705	5.83	0.000	77.56195	156.37
2 0	165.4972	29.68468	5.58	0.000	107.2135	223.780
2 1	127.9802	20.0705	6.38	0.000	88.57324	167.38
3 0	187.0955	29.68468	6.30	0.000	128.8118	245.379
3 1	142.8898	20.0705	7.12	0.000	103.4828	182.29
4 0	198.7088	29.68468	6.69	0.000	140.4251	256.99
4 1	151.0277	20.0705	7.52	0.000	111.6207	190.43
5 0	208.9253	29.68468	7.04	0.000	150.6416	267.2
5 1	159.3005	20.0705	7.94	0.000	119.8935	198.70
6 0	229.5165	29.68468	7.73	0.000	171.2327	287.80
6 1	169.5975	20.0705	8.45	0.000	130.1905	209.00
7 0	244.1537	29.68468	8.22	0.000	185.87	302.43
7 1	181.2198	20.0705	9.03	0.000	141.8128	220.62
8 0	265.8706	29.68468	8.96	0.000	207.5869	324.15
8 1	189.4721	20.0705	9.44	0.000	150.0652	228.87
9 0	288.2604	29.68468	9.71	0.000	229.9767	346.54
9 1	200.844	20.0705	10.01	0.000	161.437	240.2
10 0	312.1493	29.68468	10.52	0.000	253.8656	370.43
10 1	221.0319	20.0705	11.01	0.000	181.6249	260.438
11 0	331.1701	29.68468	11.16	0.000	272.8864	389.453
11 0						
11 1	233.9161	20.0705	11.65	0.000	194.5091	273.323
	347.9162	29.68468	11.72 12.32	0.000	289.6324	406.199
				0.000	207.9557	286.769
12 1 13 0	247.3627 366.8811	29.68468	12.36	0.000	308.5974	425.164

A fully saturated regression model

- In control states, prison rates increase over time (i.year)
- In treated states, the passage of time has a positive, but lower effect on prison rates (sum i.year & i.treated#i.year)
- In other words, prison rates increase in all states (magnitude of coefficients) but they are smaller in treated than in control states (sign of coefficients)
- Note that here 1980 is the reference year, so comparing preand post-policy is quite difficult



* A) Fully saturated model: reg pris i.treated##i.year, cluster (state

Linear regression

Number of obs = 714 F(27, 50) = 150.08 Prob > F = 0.0000 R-squared = 0.2298 Root MSE = 118.74

(Std. Err. adjusted for 51 clusters in state)

pris	Coef.	Robust Std. Err.	t	P> t	[95% Conf.	Interval]
1.treated	-26.95774	23.03849	-1.17	0.248	-73.2319	19.31642
year						
81	7.902258	2.572847	3.07	0.003	2.734542	13.06997
82	26.54346	4.055156	6.55	0.000	18.39844	34.68848
83	48.14177	7.273863	6.62	0.000	33.53179	62.75176
84	59.75509	8.602639	6.95	0.000	42.47618	77.034
85	69.97154	11.12025	6.29	0.000	47.63585	92.30722
86	90.56271	18.27964	4.95	0.000	53.84698	127.2784
87	105.2	19.39936	5.42	0.000	66.23521	144.1648
88	126.9169	26.52769	4.78	0.000	73.63443	180.1993
89	149.3067	35.73909	4.18	0.000	77.5226	221.0907
90	173.1955	34.7398	4.99	0.000	103.4186	242.9725
91	192.2164	37.49753	5.13	0.000	116.9004	267.5324
92	208.9624	41.13044	5.08	0.000	126.3495	291.5753
93	227.9274	45.63096	5.00	0.000	136.2749	319.5798
treated#year						
1 81	-2.929324	3.026349	-0.97	0.338	-9.007925	3.149277
1 82	-10.55925	4.564066	-2.31	0.025	-19.72644	-1.392048
1 83	-17.24801	8.315161	-2.07	0.043	-33.9495	5465155
1 84	-20.72339	10.23496	-2.02	0.048	-41.28092	1658684
1 85	-22.66707	12.78809	-1.77	0.082	-48.3527	3.01856
1 86	-32.96124	19.38873	-1.70	0.095	-71.90466	5.98218
1 87	-35.97616	21.1049	-1.70	0.094	-78.36659	6.414272
1 88	-49.44072	27.81833	-1.78	0.082	-105.3155	6.434047
1 89	-60.45866	36.97279	-1.64	0.108	-134.7207	13.80338
1 90	-64.1597	36.27455	-1.77	0.083	-137.0193	8.699881
1 91	-70.29631	39.17234	-1.79	0.079	-148.9763	8.383664
1 92	-73.59571	43.01121	-1.71	0.093	-159.9863	12.79485
1 93	-82.8252	47.50028	-1.74	0.087	-178.2323	12.58192
_cons	138.9538	20.97328	6.63	0.000	96.82768	181.0798

Testing pre-trends

. reg pris i.treated##i.years pre if year>= 83 & year <= 89 , cluster(state) Linear regression F(7.50)13.52 Prob > F 0.0000 R-squared 0.1007 Root MSE 108.57 (Std. Err. adjusted for 51 clusters in state) Robust pris Coef. Std. Err. P>|t| [95% Conf. Interval] 1.treated -71.66694 45.43946 -1.58 0.121 -162.9348 19.6009 years pre -3.21 0.002 -78.09804 -17.95199 17.17109 -3.39 0.001 -92.73061 -23.75231 -69.85478 18.29084 -3.82 0.000 -106.593-33.11654treated#years pre 22.04213 -8.803008 52.88726 1 1 15.35685 1.44 0.157 1 2 23.9858 17.65608 0.180 -11.47747 59.44907

0.155

0.000

1.44

5.98

-10.77429

170.7132

65.69667

343.1874

19.03627

42.93482

- Compare trends 3 years pre-policy for treated and control states
- Using 86 (year of the policy introduction) as a baseline, control states had lower prison rates in the years just before the treatment (i.year_pre)
- But this trend was not significantly different for treated states (*i.treated#i.years_pre*)

There is no significantly different pre-trend for the treated group



27.46119

256.9503

1 3

cons

Testing pre-trends, adding FEs

. xtset state year

panel variable: state (strongly balanced)
time variable: year, 80 to 93

delta: 1 unit

. xtreg pris i.treated##i.years_pre if year>= 83 & year <= 89 , fe cluster(state)
note: 1.treated omitted because of collinearity</pre>

Fixed-effects (within) regression Number of obs 357 Number of groups = 51 Group variable: state R-sa: Obs per group: within = 0.3785min = 7 between = 0.0699avg = 7.0 overall = 0.0185max = F(6,50)15.51 corr(u i, Xb) = -0.0660Prob > F 0.0000

(Std. Err. adjusted for 51 clusters in state)

pris	Coef.	Robust Std. Err.	t	P> t	[95% Conf	. Interval]
1.treated	0	(omitted)				
years pre						
1	-48.02502	14.95103	-3.21	0.002	-78.05505	-17.99498
2	-58.24146	17.14654	-3.40	0.001	-92.68131	-23.80162
2 3	-69.85478	18.26469	-3.82	0.000	-106.5405	-33.1690
reated#years pre						
1 1	22.04213	15.33489	1.44	0.157	-8.758912	52.8431
1 2	23.9858	17.63083	1.36	0.180	-11.42677	59.39838
1 3	27.46119	19.00906	1.44	0.155	-10.71963	65.642
_cons	207.7671	2.548821	81.51	0.000	202.6477	212.8866
sigma u	109.85421					
sigma e	30.799141					
rho	.92712446	(fraction	of varia	nce due t	oui)	

Same result as in the previous example:

There is no significantly different pre-trend for the treated group

Simple DiD estimation

reg pris i.treated##i.post, cluster (state)

Linear regression	Number of obs	=	714
-	F(3, 50)	=	44.58
	Prob > F	=	0.0000
	R-squared	=	0.1775
	Root MSE	=	120.61

(Std. Err. adjusted for 51 clusters in state)

pris	Coef.	Robust Std. Err.	t	P> t	[95% Conf.	Interval]
1.treated 1.post	-39.31225 123.9003	26.77102 26.98897	-1.47 4.59	0.148 0.000	-93.08343 69.69136	14.45893 178.1092
treated#post 1 1	-46.35971	27.96324	-1.66	0.104	-102.5255	9.806117
_cons	174.3394	24.38248	7.15	0.000	125.3658	223.3131

How do you interpret the results?

diff pris, treated(treated) period(post) cluster(state)

. diff pris, treated(treated) period(post) cluster(state)

DIFFERENCE-IN-DIFFERENCES ESTIMATION RESULTS

Number of ol	bservations	in the	DIFF-IN-DIFF:	714
	Before	Aft	ter	
Control:	96	128	224	
Treated:	210	280	490	
	306	408	3	

Outcome var.	pris	S. Err.	t	P> t
Before				
Control	174.339			
Treated	135.027			
Diff (T-C)	-39.312	26.771	-1.47	0.148
After				
Control	298.240			
Treated	212.568			
Diff (T-C)	-85.672	52.324	1.64	0.108
Diff-in-Diff	-46.360	27.963	1.66	0.104

R-square: 0.18

* Means and Standard Errors are estimated by linear regression

**Clustered Std. Errors

Inference: * p<0.01; ** p<0.05; * p<0.1



^{*} diff is a user-written command to be installed 35

Testing before/after trends

. reg pris i.treated##i.years_post i.treated##i.years_pre if year>= 83 & year <= 89 , cluster (state)

(Std. Err. adjusted for 51 clusters in state)

		(Std.	Err. ad	justed for	r 51 clusters	in state)
pris	Coef.	Robust Std. Err.	t	P> t	[95% Conf.	Interval]
1.treated	-59.91898	39.08622	-1.53	0.132	-138.426	18.588
years_post 1 2 3	14.63727 36.35415 58.74396	2.218082 8.914252 18.72665	6.60 4.08 3.14	0.000 0.000 0.003	10.18212 18.44935 21.13037	19.09242 54.25895 96.35755
treated#years_post	-3.01492 -16.47948 -27.49742	3.389226 9.481491 19.31209	-0.89 -1.74 -1.42	0.378 0.088 0.161	-9.82238 -35.52362 -66.2869	3.79254 2.564654 11.29206
years_pre 1 2 3	-20.59117 -30.80762 -42.42093	8.149227 10.58 11.9499	-2.53 -2.91 -3.55	0.015 0.005 0.001	-36.95938 -52.05818 -66.42301	-4.222969 -9.557056 -18.41886
treated#years_pre	10.29417 12.23785 15.71323	8.334959 10.86806 12.44591	1.24 1.13 1.26	0.223 0.266 0.213	-6.447086 -9.591288 -9.285112	27.03543 34.06698 40.71158
_cons	229.5165	36.6686	6.26	0.000	155.8654	303.1675

- Now compare 3 years pre- and postpolicy
- Compared to 86, control states had lower prison rates in the years just before the treatment (i.year_pre)
- But this trend was not significantly different for treated groups (i.treated#i.years_pre)
- After the treatment, prison rates increase in control states (i.years_post)
- This increase was smaller (but not significantly different) for treated states (i.treated#i.years_post)



Testing before/after trends with FEs

. xtset state year

panel variable: state (strongly balanced)

time variable: year, 80 to 93

delta: 1 unit

. xtreg pris i.treated##i.years_post i.treated##i.years_pre if year>= 83 & year <= 89 , fe cluster(state)
note: l.treated omitted because of collinearity</pre>

Fixed-effects (within) regression Number of obs Group variable: state Number of groups = R-sq: Obs per group: within = 0.4874min = between = 0.0699avg = 7.0 overall = 0.0514 max = F(12,50) 14.06 corr(u i, Xb) = 0.0060Prob > F 0.0000

(Std. Err. adjusted for 51 clusters in state)

pris	Coef.	Robust Std. Err.	t	P> t	[95% Conf.	Interval
1.treated	0	(omitted)				
years post						
	14.63727	2.214856	6.61	0.000	10.1886	19.0859
2 3	36.35415	8.901286	4.08	0.000	18.47539	54.2329
3	58.74396	18.69941	3.14	0.003	21.18508	96.3028
treated#years post						
1 1	-3.01492	3.384296	-0.89	0.377	-9.812478	3.78263
1 2	-16.47948	9.467699	-1.74	0.088	-35.49591	2.53695
1 3	-27.49742	19.284	-1.43	0.160	-66.23047	11.2356
years pre						
1	-20.59117	8.137373	-2.53	0.015	-36.93557	-4.24677
1 2 3	-30.80762	10.56461	-2.92	0.005	-52.02727	-9.58796
3	-42.42093	11.93251	-3.56	0.001	-66.38809	-18.4537
treated#years pre						
1 1	10.29417	8.322835	1.24	0.222	-6.422735	27.0110
1 2	12.23785	10.85225	1.13	0.265	-9.559537	34.0352
1 3	15.71323	12.42781	1.26	0.212	-9.248751	40.6752
_cons	188.3956	1.030672	182.79	0.000	186.3254	190.465
sigma_u	108.29803					
sigma_e	28.252839			mana garas succession		
rho	.93627821	(fraction	of varia	ace due t	0 11 1)	

Same result as in the previous example:

There is no significant pretrend, but there is no significant effect of the policy either

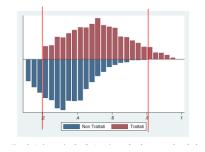
Recap

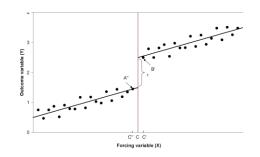
- Trends could be non-linear. Maybe the best fitting model is a quadratic trends model or other functional form
- Remember that the difference between the groups may not be parallel in the raw, unadjusted data, but they could become parallel after "holding" other variables constant or after "taking into account" the effect of other variables (in other words, the trends could become parallel conditional on other covariates)
- This is a common situation. The parallel trends test may fail with raw data (unadjusted) but it could pass when we control for covariates
- "Passing" here means that we do not reject the null hypothesis that trends are parallel

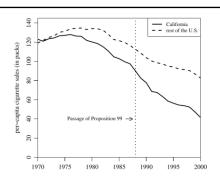


Extensions on DiD – related approaches

	MATCHING AND DID	REGRESSION DISCONTINUITY DESIGN	SYNTHETIC CONTROL GROUP
Assumption violated	Parallel Trends Random Selection into Treatment	Random Selection into Treatment	Parallel Trends Random Selection into Treatment
Main idea	Identification of untreated units (control group) similar in several respects to treated units, before the treatment, as counterfactual. Based CIA: conditional on observables, the difference between the two groups is only the exposure to the treatment	Assignment to treatment depends on (a set of) variables (i.e., forcing variables) satisfying a set of known conditions. The effect of the treatment is estimated by the discontinuity of the outcome variable at the cut-off (also, geographic).	Since only a few units are treated, the counterfactual group is built as a weighted average of other untreated units.

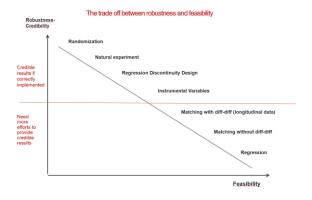


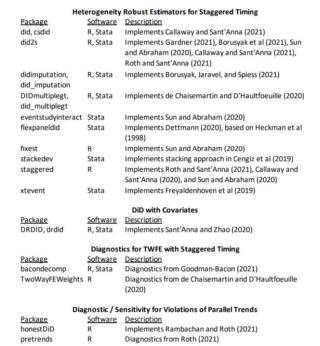




A Checklist for DiD Practitioners (to start)

- Is everyone treated at the same time?
- Are you sure about the validity of the parallel trends assumption?
- Do you have a large number of treated and untreated clusters sampled from a super-population?





More on: https://asjadnaqvi.github.io/DiD/





Extensions on DiD – complex settings

	MULTIPLE TIME PERIODS	MULTIPLE GROUPS	DYNAMIC (STAGGERED) TREATMENT
Main takeaways	Adding μ_t would control for "time trends" and $dPost$ could be more general because it could accommodate different timing of treatment for some units Adding λ_i captures time-invariation group/individual characteristics including those that could affect selection (or assignment) into the treatment/program. Interacting $\gamma_s * \mu_t$ (state-specific time trends) is often described as a robustness check: the DiD estimator shouldn't change		Not everyone is treated at the same time Standard "static" TWFE models may not represent a straightforward weighted average of unit-level treatment effects when treatment effects are allowed to be heterogeneous across time or units.
Some references	assumptions. <i>Empirical Economics</i> , 39(1) Callaway, B., & Sant'Anna, P. H. (2021). D 230. Goodman-Bacon, A. (2021). Difference-in-Bertrand, M., Duflo, E., & Mullainathan, S. journal of economics, 119(1), 249-275.	tion of the effects of dynamic treatments by se), 111-137. ifference-in-differences with multiple time periodifferences with variation in treatment timing. J (2004). How much should we trust differences collusion enforcement: Justice for consumers an	lournal of Econometrics, 225(2), 200-lournal of Econometrics, 225(2), 254-277in-differences estimates?. The Quarterly

Studies, 32(7), 2587-2624

Roth, J., Sant'Anna, P. H., Bilinski, A., & Poe, J. (2022). What's Trending in Difference-in-Differences? A Synthesis of the Recent Econometrics Literature. arXiv preprint arXiv:2201.01194. 41

It's over! Good luck with the exam!



And remember the **motivate** command in Stata, if needed!

. motivate

'If you are going through hell, keep going.'

Winston Churchill

Otherwise, there is always demotivate ©

. demotivate

Best not to dwell on what your R-using colleagues really think of you.

