A Local Projections Approach to Difference-in-Differences Event Studies

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Research question

How to estimate Difference-in-Differences (DiD) with multiple treated groups & treatment periods?

- Recent literature shows that conventional TWFE implementations can be severely biased.
- A new regression-based framework: LP-DiD.
 - o Local projections (Jordà 2005) + clean controls (Cengiz et al 2019).
- Simulation evidence to assess its performance.
- Empirical applications:
 - 1. The effect of banking deregulation on the wage share.
 - 2. Democracy & growth.

Research question

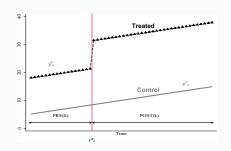
Why do we need yet another DiD estimator?

Advantages of LP-DiD:

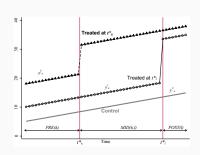
- Simpler, more transparent, easier to code, and faster to compute than other recent DiD estimators.
- Flexible: offers a general framework that can easily accommodate different settings.
- Allows matching on pre-treatment outcomes and other time-varying covariates.

Difference-in-Differences (DiD)

2x2 Setting



Staggered Setting



(Visual examples from Goodman-Bacon, 2021)

The conventional (until recently) DiD estimator: TWFE

• Static TWFE

$$y_{it} = \alpha_i + \delta_t + \beta^{TWFE} D_{it} + \epsilon_{it}$$

• Event-study (distributed lags) TWFE

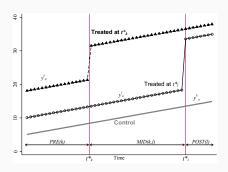
$$y_{it} = \alpha_i + \delta_t + \sum_{h=-Q}^{H} \beta_h^{TWFE} D_{it-h} + \epsilon_{it}$$

- OK in the 2x2 setting.
- Biased even under parallel trends with staggered treatment, if treatment effects are dynamic and heterogeneous.

Background

The problems with TWFE in the staggered setting

- TWFE as weighted-average of 2x2 comparisons (Goodman-Bacon 2021)
 - 1. Newly treated vs Never treated;
 - 2. Newly treated vs Not-yet treated;
 - 3. Newly treated vs Earlier treated.



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- Bias formula for TWFE (Goodman-Bacon 2021)

$$p \lim_{N \to \infty} \hat{\beta}^{TWFE} = VWATT + VWCT - \Delta ATT$$

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 TWFE as a weighted-average of cell-specific ATTs (de Chaisemartin & D'Haultfoeuille 2020)

$$E\left[\hat{\beta}^{TWFE}\right] = E\left[\sum_{(g,t):D_{gt}=1} \frac{N_{g,t}}{N_1} w_{g,t} \Delta_{g,t}\right]$$

o Weights can be negative!

LP-DiD: baseline version

A Local Projections Diff-in-Diff Estimator (LP-DiD) Baseline version

Setting & Assumptions:

- Binary absorbing treatment.
- Staggered adoption.
- Treatment effects can be dynamic & heterogeneous.
- No anticipation.
- Parallel trends.

LP-DiD: baseline versions

A Local Projections Diff-in-Diff Estimator (LP-DiD) Baseline version

Estimating equation:

$$y_{i,t+h} - y_{i,t-1} = eta_h^{LP-DiD} \Delta D_{it}$$
 } treatment indicator $+ \delta_t^h$ } time effects $+ e_{it}^h$; for $h = 0, \dots, H$.

restricting the sample to observations that are either:

$$\left\{ \begin{array}{ll} \text{newly treated} & \Delta D_{it} = 1 \,, \\ \\ \text{or clean control} & D_{i,t+h} = 0 \end{array} \right.$$

What does LP-DiD identify?

• A variance-weighted average effect:

$$E(\hat{\beta}_{h}^{LP-DiD}) = \sum_{g \neq 0} \omega_{g,h}^{LP-DiD} \tau_{g}(h)$$

o $\tau_g(h) = h$ -periods forward ATT for treatment-cohort g.

No negative weights.

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- o $\tau_g(h) = h$ -periods forward ATT for treatment-cohort g.
- No negative weights.
- Weights depend on subsample size & treatment variance:

$$\omega_{g,h}^{LP-DiD} = \frac{N_{CCS_{g,h}}[n_{gh}(n_{c,g,h})]}{\sum_{g\neq 0} N_{CCS_{g,h}}[n_{g,h}(n_{c,g,h})]},$$

- o $N_{CCS_{g,h}} =$ size of subsample including group g &its clean controls.
- o $[n_{gh}(n_{c,g,h})]$ = treatment variance in that subsample.

LP-DiD as a 'swiss knife'



Flexibility in choosing a weighting scheme

- Can apply any desired weights through weighted regression.
- Equally-weighted ATT: reweight by $1/(\omega_{g,h}^{LP-DiD}/N_g)$.
- $\omega_{g,h}^{LP-DiD}$ easy to compute from 'residualized' treatment indicator.
- Can also use regression adjustment.

LP-DiD as a 'swiss knife'



Easy to adapt to different settings

- Covariates & outcome lags
- Non-absorbing treatment
- Continuous treatment variable

LP-DiD with covariates

LP-DiD with covariates and outcome lags

Estimating equation:

```
\begin{array}{ll} y_{i,t+h} - y_{i,t-1} = & \beta_h^{LP-DiD} \Delta D_{it} & \} \  \, \text{treatment indicator} \\ & + \sum_{p=1}^P \gamma_p^h \Delta y_{i,t-p} & \} \  \, \text{outcome lags} \\ & + \sum_{m=1}^M \sum_{p=0}^P \gamma_{m,p}^h \Delta x_{m,i,t-p} & \} \  \, \text{covariates} \\ & + \delta_t^h & \} \  \, \text{time effects} \\ & + e_{it}^h \, ; & \text{for } h = 0, \dots, H \, , \end{array}
```

restricting the sample to observations that are either

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ight.$$

- Covariates will generally alter the weights.
- Can use p-score methods to make sure weights remain non-negative, or regression adjustment to get equally-weighted ATT.

LP-DiD with non-absorbing treatment

LP-DiD with non-absorbing or continuous treatment

• In general: Adapt the clean control condition to the specific setting.

LP-DiD with non-absorbing treatment

LP-DiD with non-absorbing or continuous treatment

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- Example for non-absorbing treatment:

$$\left\{ \begin{array}{ll} \text{treatment} & \left(\Delta D_{it}=1\right) & \& & \left(\Delta D_{i,t-j}=0 \text{ for } -h \leq j \leq L; j \neq 0\right) \\ \text{clean control} & \Delta D_{i,t-j}=0 \text{ for } -h \leq j \leq L \end{array} \right.$$

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Example for continuous treatment X_{it}:

$$\begin{cases} \text{movers} & \left(|\Delta X_{it}|>c\right) & \& \left(|\Delta X_{i,t-j}|\leq c \text{ for } -h\leq j\leq L; j\neq 0\right) \\ \text{quasi-stayers} & |\Delta X_{i,t-j}|\leq c \text{ for } -h\leq j\leq L \end{cases}$$

• Underlying assumption: treatment effects *stabilize* after *L* periods.

Simulation

- N=500; T=50.
- DGP:

$$Y_{0it} = \rho Y_{0,i,t-1} + \lambda_i + \gamma_t + \epsilon_{it}; \quad -1 < \rho < 1; \quad \lambda_i, \gamma_t, \epsilon_{it} \sim N(0,25)$$

- Binary staggered treatment.
- TE grows in time for 20 periods, and is stronger for early adopters.

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1 Exogenous treatment

- o Units randomly assigned to 10 groups of size N/10
- o One group never treated; others treated at $t = 11, 13, 15 \dots, 27$.

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1 Exogenous treatment

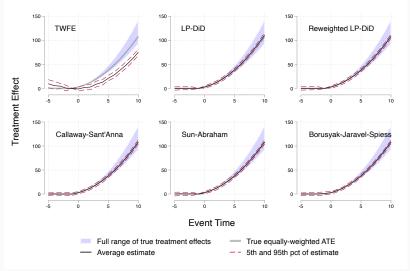
- o Units randomly assigned to 10 groups of size N/10
- o One group never treated; others treated at $t = 11, 13, 15 \dots, 27$.

2 Endogenous treatment

- o Probability of treatment depends on past outcome dynamics.
- o Negative shocks increase probability of treatment.
- o Parallel trends holds only conditional on outcome lag.

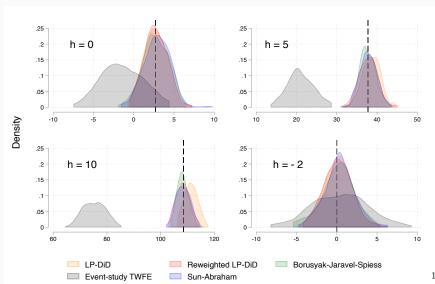
Simulation 1 – exogenous treatment scenario

True effect path and estimates from 200 replications



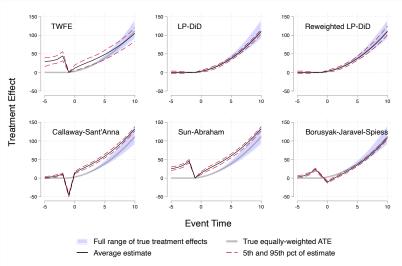
Simulation 1 – exogenous treatment scenario

Distribution of estimates from 200 replications.



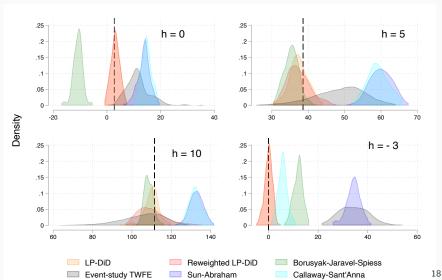
Simulation 2 – endogenous treatment scenario

True effect path and estimates from 200 replications



Simulation 2 – endogenous treatment scenario

Distribution of estimates from 200 replications.



Computational speed

Estimating the treatment effect path in a single simulation of the synthetic dataset with exogenous treatment timing:

• TWFE: 1.04 seconds

• LP-DiD: 1.2 seconds

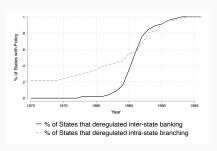
• Callaway-Sant'Anna (2020): 144.6 seconds

• Sun-Abraham (2020): 198.5 seconds

(using a laptop with 2.80 GHz Quad-core Intel i7 Processor and 16 GB of Ram)

Application: Banking Deregulation and the Labor Share

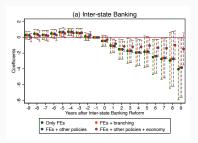
1970-1996: staggered introduction of (inter-state & intra-state) banking deregulation in US states.

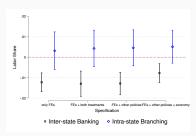


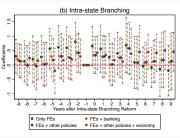
Leblebicioglu & Weinberger (EJ, 2020) use static & event-study
 TWFE to estimate effects on the labor share.

TWFE estimates

- Negative effect of *inter-state* bank deregulation (≈ -1 pp).
- No effect of *intra-state* branching deregulation.





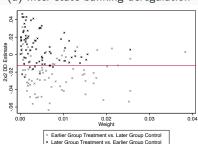


Forbidden comparisons in the TWFE specification

- TWFE uses 'forbidden' comparisons: earlier liberalizers are controls for later liberalizers.
- Goodman-Bacon (2021) decomposition to quantify their influence.
- Contribution of unclean comparisons to TWFE estimates:
 - o 36% for inter-state banking deregulation;
 - o 70% for intra-state branching deregulation.

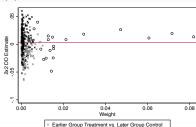
Goodman-Bacon (2021) decomposition diagnostic for the static TWFE estimate

(a) Inter-state banking deregulation



Later Group Treatment vs. Earlier Group Control

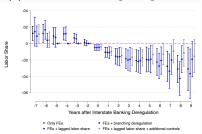
(b) Intra-state branching deregulation



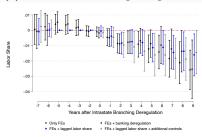
 Later Group Treatment vs. Earlier Group Control Treatment vs. Already Treated

Effect of banking deregulation on the labor share: LP-DiD estimates

(a) Inter-state banking deregulation



(b) Intra-state branching deregulation



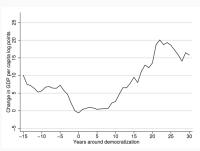
Application: Democracy and economic growth

- Acemoglu, Naidu, Restrepo and Robinson (2019).
- 1960-2010 panel on 175 countries & binary measure of democracy.
- Potential for negative weights.
- Non-absorbing treatment.
- Selection based on pre-treatment GDP dynamics.

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GDP per capita around democratization



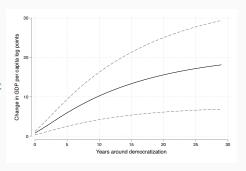
Effect of democracy on growth: dynamic panel estimates

• Dynamic fixed effects specification:

$$y_{ct} = \beta D_{ct} + \sum_{j=1}^{p} \gamma_j y_{c,t-j} + \alpha_c + \delta_t + \epsilon_{ct},$$

• Long-run effect: $\frac{\hat{\beta}}{1-\sum_{j=1}^{p}\hat{\gamma}_{j}}=21pp$ (s.e. 7pp)

IRF from the dynamic panel estimates



Effect of democracy on growth: LP-DiD specification

$$y_{c,t+h} - y_{c,t-1} = \beta_h^{LP\ DiD} \Delta D_{ct} + \delta_t^h + \sum_{j=1}^p \gamma_j^h y_{c,t-j} + \epsilon_{ct}^h$$
. (1)

restricting the estimation sample to:

$$\begin{cases} \text{ democratizations} & D_{it} = 1; D_{i,t-j} = 0 \text{ for } 1 \leq j \leq L \\ \text{ clean controls} & D_{i,t-j} = 0 \text{ for } 0 \leq j \leq L. \end{cases}$$
 (2)

Effect of democracy on growth: LP-DiD specification

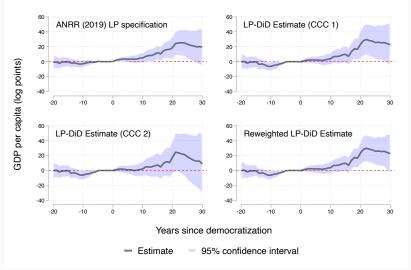
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 (2)

- We set L=20 years.
- Acemoglu et al. LP analysis: a version of this, but controls defined by $D_{it} = D_{i,t-1} = 0$, and treated defined as $\Delta D_{it} = 1$
- They still include countries that transition at or before t-1.

Effect of democracy on growth: LP-DiD estimates



Takeaways

Conclusions

- LP-DiD offers a flexible overarching framework for DiD settings.
- Simpler and less computationally intensive than estimators that aggregate many group-specific averages.
- Flexibility in defining the treatment and ('clean') control units based on the setting.
- Allows matching on pre-treatment outcomes and other time-varying covariates.

Additional Slides

Identification Assumptions (baseline specification)

No anticipation

$$E[y_{it}(p) - y_{it}(0)] = 0$$
, for all p and t such that $t < p$.

Units do not respond in anticipation of a future treatment.

Parallel trends

$$E[y_{it}(0) - y_{i1}(0)|p_i = p] = E[y_{it}(0) - y_{i1}(0)],$$

for all $t \in \{2, ..., T\}$ and for all $p \in \{1, ..., T, \infty\}.$

Average trends in untreated potential outcomes do not depend on treatment status.

A1 - Other new DiD estimators

Alternative estimators: de Chaisemartin & D'Haultfoeiulle

de Chaisemartin & D'Haultfoeiulle estimator

- For a given time-horizon ℓ , it estimates the average effect of having switched in or out of treatment ℓ periods ago.
- A weighted average, across time periods t and possible values of treatment d, of 2x2 DiD estimators.
- The constituent 2x2 DiDs compare the $t-\ell-1$ to t outcome change, in groups with a treatment equal to d at the start of the panel and whose treatment changed for the first time in $t-\ell$ (the first-time switchers) and in control groups with a treatment equal to d from period 1 to t (not-yet switchers).

Alternative estimators: Callaway-Sant'Anna

Callaway-Sant'Anna estimator

- Estimates each group specific effect at the selected time horizon.
- Take long-differences in the outcome variable, and compare each treatment group *g* with its control group.
- To control for covariates, re-weight observations based on outcome regression (OR), inverse-probability weighting (IPW) or doubly-robust (DR) estimation.
- Aggregate group-time effects into a single overall ATT using some weights.

Alternative estimators: Sun-Abraham

Sun-Abraham interaction-weighted estimator

- Event-study DiD specification, with leads and lags of the treatment variable.
- Includes a full set of interaction terms between relative time indicators D_{it}^k (ie, leads and lags of the treatment variable) and treatment cohort indicators $1\{G_g=g\}$ (dummies for when a unit switches into treatment).
- Then calculates a weighted average over cohorts g for each time horizon, in order to obtain a standard event-study plot.

Alternative estimators: Borusyak-Jaravel-Spiess

Borusyak-Jaravel-Spiess imputation estimator

- Estimate unit and time FEs only using untreated sample.
- Take them out from Y to form counterfactual Y'.
- Then for any treatment group, just compare Y and Y' for treated units around event time.
- Average these across events to get an average effect.