

TWFE Metrics Discussion Group: Implementing Callaway & Sant'Anna (2020) DID Estimator

Apr 29, 2021

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NOTE: current as of 4/29/2021; information on resources and package availability likely to change



Current state of the literature

(incomplete, but a start)

Why is
staggered DiD
a problem?

How do we
do things
differently?

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Two reasons, according to Goodman-Bacon (2021)

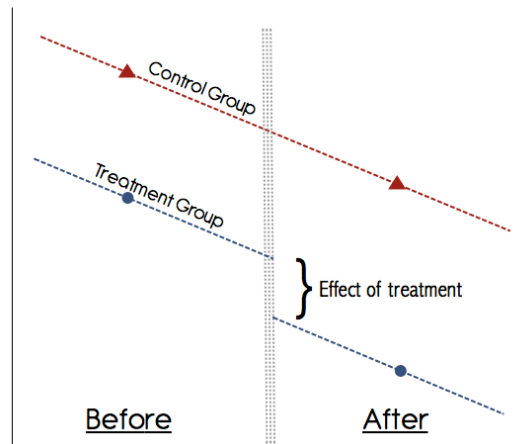
1. Variance weighting (implicit in OLS)
2. Use of past treated units as controls for later-treated units

Why is staggered DiD a problem?

Two reasons, according to Goodman-Bacon (2021)

1. Variance weighting (implicit in OLS)
2. Use of past treated units as controls for later-treated units

Goodman-Bacon (2021) *<-- DiD with constant post-period effects*



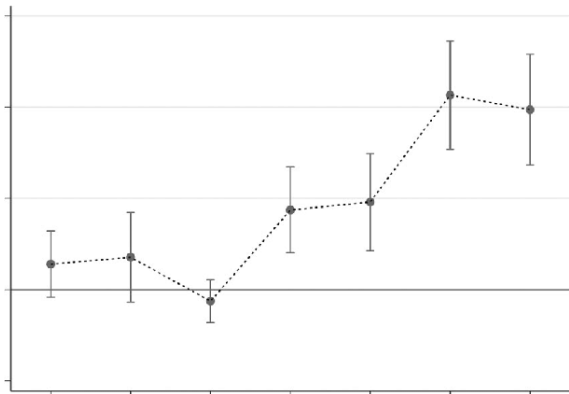
Why is staggered DiD a problem?

Two reasons, according to Goodman-Bacon (2021)

- ~~1. Variance weighting (implicit in OLS)~~
2. Use of past treated units as controls for later-treated units

Sun & Abraham (2020) *<-- Focus on event-study DiD (dynamic post period effects)*

Show that event-study fixes problem #1 but not #2



Current state of the literature:

1. Why is staggered DiD a problem?

→ *Goodman-Bacon (2020) provides tools to diagnose how bad of a problem staggered DiD is in your setting.*

→ *That said, we know there's some degree of a problem, so let's move onto the fix.*

Current state of the literature:

1. Why is staggered DiD a problem?

→ *Goodman-Bacon (2020) provides tools to diagnose how bad of a problem staggered DiD is in your setting.*

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2. How do we do things differently?

How do we do things differently?

Current options for doing things differently (Baker, 2020)

1. Stacked DiD Estimator
2. Sun & Abraham (2020) Estimator
3. Callaway & Sant'Anna (2020) Estimator

How do we do things differently?

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The are the **same** in that they all:

- Rely on event-study models not pooled DiD
- Modify the set of units that act as effective controls in the estimation process

How do we do things differently?

Current options for doing things differently (Baker, 2020)

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They **vary** ONLY in terms of:

- Which units are used as effective controls (for treatment cases)
- How covariates are incorporated into the analysis

How do we do things differently?

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3. **Callaway & Sant'Anna (2020) Estimator**

They **vary** ONLY in terms of:

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- How covariates are incorporated into the analysis

*Baker (2020) suggests you implement one of these three methods to test robustness of inferences.
My take is that Callaway & Sant'Anna allows for the most flexibility + customization*

Callaway & Sant'Anna Estimator:

1. Estimate group-time effects: unique estimates per cohort of units treated at the same point in time
2. Identify appropriate comparison units for each group-time treated group
 - Pairings of units in the treatment and comparison groups per group-time treatment

Implement the Callaway & Sant'Anna Estimator in R

Stata wrappers by: Jonathan Roth & Nick Huntington-Klein

- <https://github.com/jonathandroth/staggered#stata-implementation>
- <https://github.com/NickCH-K/did>

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April 2021

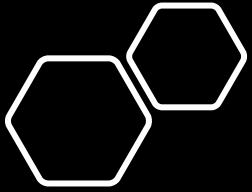
Replying to @peternka

For now, the Stata implementations don't have the richness and options of the R version.

That richness is important. For example, we report multiple aggregations of the ES parameters.

The R package is easy to use and we are happy to share our code or discuss!





Or wait for a
user-written
Stata package to
emerge



Asjad Naqvi @AsjadNaqvi · 15h

Half way done with hard-coding Sant'Anna, Pedro H. C., and Zhao, Jun (2020) DRDID R-package in Stata. The main hurdle is Mata coding for propensity score matching. Took a while to get this exact match to the R output.

```

: tempname
: experimental
re
: yname
tname
: idname
dname
w
post
y1
y0
D
deltaV
intercept
control
: end

      grad
      1      2      3      4      5      6      7      8
1  -.0168279125  .006237643  .0006962939  -.0002628433  .0006649905  .0000160856  -1.273542535  .0007238796

      hess
[symmetric]
      1      2      3      4      5      6      7      8
1  16.56968726
2  6.187131496  2.87741089
3  .5009888221  .2165617164  .0214065333
4  .1281830305  .0384025548  .0029220035  .0038182921
5  .4892017775  .2052684591  .017519195  .0027493258  .0217407047
6  .0653057833  .028780705  0  .0005467719  .0025486715  .0029398841
7  2041.000555  957.6316493  75.61879034  33.92870332  66.44260266  10.38161259  1149471.474
8  .6160526382  .2699886328  .0214065333  .0038182921  .0217407047  .0029398841  93.79981573  .0266116788
    
```

4 13 171 [share icon]

*Note:

The DRDID R-package is different from the the Callaway & Sant'Anna estimator for multi-period fixed effects ("DID" package in R). In fact, DID builds on top of DRDID.

DRDID focuses on scenarios with only two time periods (pre-treatment and post-treatment). It essentially reweights by propensity score and controls for vars included in propensity score calculation ... a.k.a. "doubly" robust ("dr")

Crash course in using R

*From one novice to
another*

<https://uvastatlab.github.io/phdplus/installR.html>

- A. **Download/install** R and RStudio if you've never used either before
 - You need both “R-base” and “RStudio”

- B. Or, **check your RStudio version + update** to the newest if needed
 - Check for most current version on <https://www.r-project.org/>
→ “R version 4.0.5 (Shake and Throw) has been released on 2021-03-31.”

Install packages

In Stata, **ssc install**

In R, there are two necessary steps:

(1) Install

```
> install.packages("<package name>")
```

(2) “activate” the package during each analysis session you plan to use it

```
> library("<package name>")
```

<https://www.datacamp.com/community/tutorials/top-ten-most-important-packages-in-r-for-data-science>

Package Info: <https://bcallaway11.github.io/did/>

Steps

1. Install the Callaway & Sant'Anna package



You can install `did` from CRAN with:

```
install.packages("did")
```

or get the latest version from github with:

```
# install.packages("devtools")
devtools::install_github("bcallaway11/did")
```

2. Activate the package



```
library(did)
```

3. Bring in data



```
maindata <- read_dta("D:/TFDW/D_R_mat_201006.dta")
```

4. Implement/run function...

Estimate Group- Time Average Treatment Effects

Start with the most basic approach,
relying on package defaults

Estimate Group-Time Average Treatment Effects

- Function to use in R = “`att_gt`”

```
example_attgt <- att_gt(yname = "Y",  
                        tname = "period",  
                        idname = "id",  
                        gname = "G",  
                        xformula = ~X,  
                        data = dta  
                        )
```

Estimate Group-Time Average Treatment Effects

- Function to use in R = `att_gt`

```
example_attgt <- att_gt(yname = "Y",  
                       tname = "period",  
                       idname = "id",  
                       gname = "G",  
                       xformula = ~X,  
                       data = dta  
                       )
```

Calendar time or ordered units

1st period treated (constant w/in id)

Estimate Group-Time Average Treatment Effects

- Function to use in R = “att_gt”

```
example_attgt <- att_gt(yname = "Y",  
                        tname = "period",  
                        idname = "id",  
                        gname = "G",  
                        xformla = ~X,  
                        data = dta  
                        )
```

Specify pre-treatment covariates that predict treatment or “NULL” for no covariates

Estimate Group-Time Average Treatment Effects

- What use are covariates here?

This feature builds in a DRDID (doubly robust DID) estimator where you:

- (1) reweight control units per group-time cohort using propensity score to ensure control units are similar on observables
- (2) Partials out variation in outcomes due to covariates, as estimated among control units

Specify pre-treatment covariates that predict treatment or “NULL” for no covariates

```
xformula = ~X,
```

In other words, this options controls for covariates + reweights with propensity score

Estimate Group-Time Average Treatment Effects

Example produces estimates ATE for groups 2, 3, and 4

Recall group (gname) = first period treated

```
summary(example_attgt)
#> Group-Time Average Treatment Effects:
#> Group Time ATT(g,t) Std. Error [95% Simult. Conf. Band]
#> 2 2 0.9615 0.0593 0.8035 1.1196 *
#> 2 3 2.0424 0.0587 1.8859 2.1989 *
#> 2 4 3.0104 0.0601 2.8503 3.1705 *
#> 3 2 -0.0392 0.0551 -0.1862 0.1077
#> 3 3 1.0877 0.0617 0.9233 1.2522 *
#> 3 4 1.9938 0.0589 1.8369 2.1506 *
#> 4 2 -0.0537 0.0585 -0.2096 0.1022
#> 4 3 0.0382 0.0563 -0.1119 0.1883
#> 4 4 0.9410 0.0578 0.7869 1.0951 *
#> ---
#> Signif. codes:  `*' confidence band does not cover 0
#>
#> P-value for pre-test of parallel trends assumption: 0.79243
#> Control Group: Never Treated, Anticipation Periods: 0
#> Estimation Method: Doubly Robust
```


Estimate Group-Time Average Treatment Effects

Example produces estimates ATE for groups 2, 3, and 4

Recall group (gname) = first period treated

```
summary(example_attgt)
#> Group-Time Average Treatment Effects:
#>   Group Time ATT(g,t) Std. Error [95% Simult. Conf. Band]
#>   2     2    0.9615    0.0593    0.8035    1.1196 *
#>   2     3    2.0424    0.0587    1.8859    2.1989 *
#>   2     4    3.0104    0.0601    2.8503    3.1705 *
#>   3     2   -0.0392    0.0551   -0.1862    0.1077
#>   3     3    1.0877    0.0617    0.9233    1.2522 *
#>   3     4    1.9938    0.0589    1.8369    2.1506 *
#>   4     2   -0.0537    0.0585   -0.2096    0.1022
#>   4     3    0.0382    0.0563   -0.1119    0.1883
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```

Effect in 1st year treated for group 2

Estimate Group-Time Average Treatment Effects

Example produces estimates ATE for groups 2, 3, and 4

Recall group (gname) = first period treated

```
summary(example_attgt)
#> Group-Time Average Treatment Effects:
#> Group Time ATT(g,t) Std. Error [95% Simult. Conf. Band]
#>      2      2  0.9615    0.0593    0.8035    1.1196 *
#>      2      3  2.0424    0.0587    1.8859    2.1989 *
#>      2      4  3.0104    0.0601    2.8503    3.1705 *
#>      3      2 -0.0392    0.0551   -0.1862    0.1077
#> 3      3  1.0877    0.0617    0.9233    1.2522 *
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#> P-value for pre-test of parallel trends assumption: 0.79243
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```

Effect in 1st year treated for group 3

Estimate Group-Time Average Treatment Effects

Example produces estimates ATE for groups 2, 3, and 4

Recall group (gname) = first period treated

```
summary(example_attgt)
#> Group-Time Average Treatment Effects:
#> Group Time ATT(g,t) Std. Error [95% Simult. Conf. Band]
#>      2      2  0.9615    0.0593    0.8035    1.1196 *
#>      2      3  2.0424    0.0587    1.8859    2.1989 *
#>      2      4  3.0104    0.0601    2.8503    3.1705 *
#>      3      2 -0.0392    0.0551   -0.1862    0.1077
#>      3      3  1.0877    0.0617    0.9233    1.2522 *
#>      3      4  1.9938    0.0589    1.8369    2.1506 *
#>      4      2 -0.0537    0.0585   -0.2096    0.1022
#>      4      3  0.0382    0.0563   -0.1119    0.1883
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#> ---
#> Signif. codes:  `*' confidence band does not cover 0
#>
#> P-value for pre-test of parallel trends assumption: 0.79243
#> Control Group: Never Treated, Anticipation Periods: 0
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```

Effect in 1st year treated for group 4

Estimate Group-Time Average Treatment Effects

Example produces estimates ATE for groups 2, 3, and 4

Recall group (gname) = first period treated

```
summary(example_attgt)
#> Group-Time Average Treatment Effects:
#> Group Time ATT(g,t) Std. Error [95% Simult. Conf. Band]
#>      2      2  0.9615    0.0593    0.8035    1.1196 *
#>      2      3  2.0424    0.0587    1.8859    2.1989 *
#>      2      4  3.0104    0.0601    2.8503    3.1705 *
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Effect in 1st year treated for group 4

Estimate Group-Time Average Treatment Effects

What are the group-time point estimates relative to?

```
summary(example_attgt)
#> Group-Time Average Treatment Effects:
#> Group Time ATT(g,t) Std. Error [95% Simult. Conf. Band]
#>      2      2      0.9615      0.0593      0.8035      1.1196 *
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```

Pre-treatment estimates:

- Reference is always the period before ($t = t - 1$)*
*this is why period=1 is not estimated

Post-treatment estimates:

- Reference point is the period before treatment ($t = -1$)

Estimate Group-Time Average Treatment Effects

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Estimate Group-Time Average Treatment Effects


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```

Built in parallel trends test of all pre-treatment estimates

Estimate Group-Time Average Treatment Effects

Default settings shown below results

```
summary(example_attgt)
#> Group-Time Average Treatment Effects:
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```

Estimate Group- Time Average Treatment Effects

Move beyond package defaults

Select Alternative Control Groups

DEFAULT = never treated

- If no never treated, last treated group serves as control group for all prior treated groups (a warning will pop up)

```
example_attgt_altcontrol <- att_gt(yname = "Y",  
                                   tname = "period",  
                                   idname = "id",  
                                   gname = "G",  
                                   xformula = ~X,  
                                   data = dta,  
                                   control_group = "notyettreated"  
                                   )
```

Includes:

- (a) never treated (if any) +*
- (b) not yet treated*

Quick Note on Coding var for “gname”

Make sure to code your “never treated” group as “0”

```
#Replace missing values of first treatment year to 0 for never treated  
dta$G[is.na(dta$G)] <- 0
```

```
example_attgt <- att_gt(yname = "Y",  
                       tname = "period",  
                       idname = "id",  
                       gname = "G", 1st period treated  
                       xformula = ~X,  
                       data = dta Name of dataset  
                       )
```

Estimate Group-Time Average Treatment Effects

Move from individual estimates of group-time ATEs by period pre/post treatment to something more useful

Aggregating group-time average treatment effects

Recall, example produces estimates ATE for groups 2, 3, and 4, where group (gname) = first period treated

Function in R = “**aggte**”

Options for aggregation

1. Simple → Weighted average of all group time ATEs, weights proportional to group size
2. Dynamic/event-study → Average group time ATEs at each time point post treatment
3. Group-specific → Average all post-treatment time points by treatment time group
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Aggregating group-time average treatment effects

Recall, example produces estimates ATE for groups 2, 3, and 4, where group (gname) = first period treated

Function in R = “**aggte**”

Options for aggregation

1. Simple

```
agg.simple <- aggte(example_attgt, type = "simple")
```

2. Dynamic/event-study

```
agg.es <- aggte(example_attgt, type = "dynamic")
```

3. Group-specific

```
agg.gs <- aggte(example_attgt, type = "group")
```

4. Calendar time

```
agg.ct <- aggte(example_attgt, type = "calendar")
```

Implementing the Callaway & Sant'Anna Estimator in R

Can specify/customize many other options!

Helpful resources include:

- <https://causalinf.substack.com/p/callaway-and-santanna-dd-estimator>
- <https://bcallaway11.github.io/did/articles/did-basics.html>

Recent paper using this method -- Gilpin, Karger, Nencka (2021):

- <https://www.chicagofed.org/publications/working-papers/2021/2021-06>

Thanks!