

Most of treatment effect methods extend to situations with imperfect compliance without imposing much stronger assumptions

- RCTs with perfect compliance identify the ATE without having to make any assumption.
- RCTs with imperfect compliance identify the LATE under exclusion restriction and monotonicity: often credible assumptions in context of RCTs

Most of treatment effect methods extend to situations with imperfect compliance without imposing much stronger assumptions

- RCTs with perfect compliance identify the ATE without having to make any assumption.
- RCTs with imperfect compliance identify the LATE under exclusion restriction and monotonicity: often credible assumptions in context of RCTs
- Sharp RDs identify the ATE at the threshold under the assumption that the conditional mean of the outcome is continuous
- Fuzzy RDs identify the LATE at the threshold under the same assumption plus monotonicity
- DID with imperfect compliance rely however on much stronger assumptions than standard DID

Many natural experiments do not lead to a sharp change in treatment rate for any group defined by a set of observable characteristics, but only to a larger increase of the treatment rate in some groups than in others.

- A good example is Duflo (AER, 2001), who uses a primary school construction program in Indonesia to measure returns to education
- Many primary schools were constructed in districts where there were few schools before the program: Treatment Group
- Few primary schools were constructed in districts which already had many schools: **Control Group**
- Primary school completion rate increased more in treatment group (but not from 0 to 100) compared to control group (didn't remain constant)

=

=

Table 1: Share of individuals completing primary school

	Older cohort	Younger cohort
High treatment regions	81.2%	90.0%
Low treatment regions	89.8%	94.3%

• In such fuzzy design, a commonly used estimand is the Wald-DID estimand

$$W_{DID} = \frac{E(Y_{1,1}) - E(Y_{1,0}) - \left(E(Y_{0,1} - E(Y_{0,0}))\right)}{E(D_{1,1}) - E(D_{1,0}) - \left(E(D_{0,1}) - E(D_{0,0})\right)}$$
(1)

- ullet ratio of DID of outcome Y variable and DID of treatment rates D
- What happens to W_{DID} in sharp design?
- In fuzzy designs, Wald-DID estimand relies on stronger assumptions than DID in sharp designs (De Chaisemartin & D'Haultfoeuille, RESTUD 2018)

- ullet Notation: drop i subscript from notation in Session 1
- For any random variable R and for every (d,g,t), let $R_{g,.}\sim R|G=g$, $R_{.,t}\sim R|T=t,\ R_{g,t}\sim R|G=g,T=t$, and $R_{d,g,t}\sim R|D=d,G=g,T=t$
- For any $(g,t)\in\{0,1\}^2$, let $\Delta_{g,t}^{TR}=Eig(Y(1)-Y(0)|D=1,G=g,T=tig)$: ATT in group g and period t
- DID in treatment rates: $DID_D = E(D_{1,1}) E(D_{1,0}) \left(E(D_{0,1} E(D_{0,0}))\right)$
- Recall DID assumption: Assumption 1

$$E(Y_{i1}(0)|G_i=1) - E(Y_{i0}(0)|G_i=1) = E(Y_{i1}(0)|G_i=0) - E(Y_{i0}(0)|G_i=0)$$

• Theorem 1: If Assumption 1 is satisfied

$$W_{DID} = \frac{E(D_{1,1})}{DID_D} \Delta_{1,1}^{TR} - \frac{E(D_{1,0})}{DID_D} \Delta_{1,0}^{TR} - \frac{E(D_{0,1})}{DID_D} \Delta_{0,1}^{TR} + \frac{E(D_{0,0})}{DID_D} \Delta_{0,0}^{TR}$$
 (2)



Proof Theorem 1

$$\begin{split} W_{DID} &= \frac{DID}{DID_D} \\ DID &= E(Y_{1,1}) - E(Y_{1,0}) - E(Y_{0,1}) + E(Y_{0,0}) \\ &= E(Y_{1,1}(1)|D = 1)E(D_{1,1}) + E(Y_{1,1}(0)|D = 0) \left(1 - E(D_{1,1})\right) \\ &- E(Y_{1,0}(1)|D = 1)E(D_{1,0}) - E(Y_{1,0}(0)|D = 0) \left(1 - E(D_{1,0})\right) \\ &- E(Y_{0,1}(1)|D = 1)E(D_{0,1}) - E(Y_{0,1}(0)|D = 0) \left(1 - E(D_{0,1})\right) \\ &+ E(Y_{0,0}(1)|D = 1)E(D_{0,0}) + E(Y_{0,0}(0)|D = 0) \left(1 - E(D_{0,0})\right) \\ &= E(Y_{1,1}(1) - Y_{1,1}(0)|D = 1)E(D_{1,1}) + E(Y_{1,1}(0)) \\ &- E(Y_{1,0}(1) - Y_{1,0}(0)|D = 1)E(D_{1,0}) - E(Y_{1,0}(0)) \\ &- E(Y_{0,1}(1) - Y_{0,1}(0)|D = 1)E(D_{0,1}) - E(Y_{0,1}(0)) \\ &+ E(Y_{0,0}(1) - Y_{0,0}(0)|D = 1)E(D_{0,0}) + E(Y_{0,0}(0)) \\ &= E(D_{1,1})\Delta_{1,1}^{TR} - E(D_{1,0})\Delta_{1,0}^{TR} - E(D_{0,1})\Delta_{0,1}^{TR} + E(D_{0,0})\Delta_{0,0}^{TR} \end{split}$$

- ullet Under common trends alone, W_{DID} identifies a weighted sum of the 4 ATTs in each group and time period
 - 2 of the ATTs enter with a negative weights
 - Weights can be estimated from data

Intuition:

- In a sharp DID, the only departure from the scenario where nobody is treated is that units in group 1 and period 1 receive the treatment
- any discrepancy between the trends of the mean outcome in the two groups must come from the effect of the treatment in group 1 and period 1: DID identifies ATT
- In a fuzzy design, there are potentially four departures from the scenario where nobody is treated
- the discrepancy between the trends of the mean outcome in the two groups can come from the treatment effect in any group and time period
- 2 ATTs enter with a negative sign because of first difference inherent to DID
- Why is this an issue?



Wald- DID

- Why is this an issue?
- ullet W_{DID} may not even have the same sign as any of the $\Delta_{g,t}^{TR}$ if they are heteregeneous
- ullet Under common trends assumption alone, W_{DID} does not identify anything meaningful: link to structural models
- \bullet We can impose more assumptions to make W_{DID} identify something meaningful

Wald - DID: More assumptions

Assumption 2 (*Treatment monotonicity*): \exists 2 random variables D(0), D(1) for period 0 and 1:

- D = D(T)
- **2** $(D(0), D(1)) \perp T|G$
- **3** $P(D(1) \ge D(0)|G) = 1$ or $P(D(1) \le D(0)|G) = 1$

Assumption 3 (Stable treatment effect): $\forall g \in \{0, 1\}$

$$E\big(Y(1) - Y(0)|G = g, T = 1, D(0) = 1\big) = E\big(Y(1) - Y(0)|G = g, T = 0, D(0) = 1\big)$$

- Assumption 2.2: Distribution of D stable across period in each group
- Assumption 2.3: between each pair of consecutive periods, in a given group there cannot be both units whose treatment increases and units whose treatment decreases

Wald - DID: More assumptions

- Assumption 3: in every group, the average treatment effect among units treated in period 0 does not change between 0 and 1.
- This restricts treatment effect heterogeneity over time, but not between groups
- $S = \{D(1) \neq D(0), T=1\}$: units whose treatment status switches between $\mathsf{T} = \mathsf{0}$ and $\mathsf{T} = \mathsf{1}$
- $\forall g \in \{0,1\}, \Delta_{g,1}^S = E\big(Y(1) Y(0)|S,G=g,T=1\big)$: LATE of switchers in group g and at period 1

Wald - DID: More assumptions

• **Theorem 2:** If Assumption 1 to 3 are satisfied,

$$W_{DID} = \frac{E(D_{1,1}) - E(D_{1,0})}{DID_D} \Delta_{1,1}^S - \frac{E(D_{0,1}) - E(D_{0,0})}{DID_D} \Delta_{0,1}^S$$
 (3)

- ullet W_{DID} identifies in this case a weighted sum of LATEs of switchers in both groups
- Weights can be estimated: we can check whether this estimator is identifying something meaningful or not
- Proof

$$\begin{split} E(D_{1,1}) - E(D_{1,0}) &= P(D(1) = 1 | G = 1, T = 1) - P(D(0) = 1 | G = 1, T = 0) \\ &= P(D(1) = 1 | G = 1, T = 1) - P(D(0) = 1 | G = 1, T = 1) \\ &= P(D(1) = 1, D(0) = 0 | G = 1, T = 1) \\ &= P(S|G = 1, T = 1). \end{split} \tag{1.22}$$

Moreover.

$$\begin{split} E(Y_{1,1}) - E(Y_{1,0}) \\ &= E(Y|G=1,T=1) - E(Y|G=1,T=0) \\ &= E(Y(1)|G=1,T=1,D(0)=1) \, P(D(0)=1|G=1,T=1) \\ &+ E(Y(1)|G=1,T=1,S) \, P(S|G=1,T=1) \\ &+ E(Y(0)|G=1,T=1,D(1)=0) P(D(1)=0|G=1,T=1) \\ &+ E(Y(0)|G=1,T=1,D(1)=0) P(D(1)=0|G=1,T=0) \\ &- E(Y(1)|G=1,T=0,D(0)=1) \, P(D(0)=1|G=1,T=0) \\ &- E(Y(0)|G=1,T=0,D(0)=0) P(D(0)=0|G=1,T=0) \\ &= E(Y(1) - Y(0)|G=1,T=1,D(0)=1) \, P(D(0)=1|G=1,T=1) \\ &+ E(Y(1) - Y(0)|G=1,T=1,S) \, P(S|G=1,T=1) \\ &+ E(Y(0)|G=1,T=1) \\ &- E(Y(1) - Y(0)|G=1,T=0,D(0)=1) \, P(D(0)=1|G=1,T=0) \\ &- E(Y(1) - Y(0)|G=1,T=1,S) \, P(S|G=1,T=1) \\ &+ E(Y(0)|G=1,T=1) \\ &- E(Y(1) - Y(0)|G=1,T=0,D(0)=1) \, P(D(0)=1|G=1,T=1) \\ &- E(Y(1) - Y(0)|G=1,T=1,S) \, P(S|G=1,T=1) + E(Y(0)|G=1,T=1) - E(Y(0)|G=1,T=0) \\ &= \Delta_{1}^{S}, P(S|G=1,T=1) + E(Y_{1,1}(0)) - E(Y_{1,0}(0)). \end{split}$$

(1.23)

Similarly, one can show that

$$E(D_{0,1}) - E(D_{0,0}) = P(S|G = 0, T = 1)$$
 (1.24)

$$E(Y_{0,1}) - E(Y_{0,0}) \ = \ \Delta_{0,1}^S P(S|G=0,T=1) + E(Y_{0,1}(0)) - E(Y_{0,0}(0)). \eqno(1.25)$$

Taking the difference between Equations (1.23) and (1.25), and using Assumption (1.25), and using Assumption (1.25), and (1.25), an

$$DID = \Delta_{1,1}^S P(S|G=1,T=1) - \Delta_{0,1}^S P(S|G=0,T=1).$$

Dividing each side by DID_D and using Equations (1.22) and (1.24) yields the result.

13/30

- \bullet In which cases is W_{DID} estimate meaningful or not under these assumptions?
- ullet When the share of treated units does not change over time in the control group, W_{DID} identifies the treatment effect among treatment group switchers
 - under: common trends + treatment monotonicity + stable treatment effect

- \bullet In which cases is W_{DID} estimate meaningful or not under these assumptions?
- ullet When the share of treated units does not change over time in the control group, W_{DID} identifies the treatment effect among treatment group switchers
 - under: common trends + treatment monotonicity + stable treatment effect
- ullet W_{DID} does not identify treatment effect when treatment rates increase in both treatment and control groups
 - ullet Need to assume treatment effect homogeneity across groups: $\Delta_{1,1}^S=\Delta_{0,1}^S$

To sum up

- Sharp design: DID estimand only relies on a common trends assumption
- Fuzzy design: Wald-DID estimand identifies some interpretable measure of the treatment effect only if the treatment effect is homogeneous, at least over time,
- Sometimes it is necessary to impose treatment effect homogeneity both over time and between groups
- Have positive weights in Wald-DID only in very specific cases
- De Chaisemartin & D'Haultfoeuille (RESTUD 2018) propose an alternative estimand that relies on weaker assumptions

Time-Corrected Wald Estimand (De Chaisemartin & D'Haultfoeuille, RESTUD 2018)

- Assumption 4: Conditional common trend $E\big(Y(0)|G,T=1,D(0)=0\big)-E\big(Y(0)|G,T=0,D(0)=0\big) \text{ and } E\big(Y(1)|G,T=1,D(0)=1\big)-E\big(Y(1)|G,T=0,D(0)=1\big) \text{ do not depend on } G$
- Mean of Y(0) (resp. Y (1)) follows the same evolution over time among treatment and control group units that were untreated (resp. treated) at T=0
- Define $\delta_d = E(Y_{d,0,1}) E(Y_{d,0,0})$: change in the mean outcome between period 0 and 1 for control group units with treatment status d

$$W_{TC} = \frac{E(Y_{1,1}) - E(Y_{1,0} + \delta_{D_{1,0}})}{E(D_{1,1}) - E(D_{1,0})} \tag{4}$$



Time-Corrected Wald Estimand

$$\begin{split} W_{TC} &= \frac{E(Y_{1,1}) - E(Y_{1,0} + \delta_{D_{1,0}})}{E(D_{1,1}) - E(D_{1,0})} \\ &= \frac{E(Y_{1,1}) - E(Y_{1,0} + (1 - D_{1,0})\delta_0 + D_{1,0}\delta_1)}{E(D_{1,1}) - E(D_{1,0})} \\ &= \frac{E(Y_{1,1}) - E(Y_{1,0}) - (1 - P(D_{1,0} = 1))\delta_0 - P(D_{1,0} = 1)\delta_1}{E(D_{1,1}) - E(D_{1,0})} \end{split}$$

- Theorem 3: If Assumption (2) and (4) are satisfied and if $E(D_{0,1})=E(D_{0,0})$, then $W_{TC}=\Delta_{1,1}^S$
- Proof:

Following the same steps as those used to derive Equation (1.23), we obtain

$$\begin{split} &E(Y_{1,1})-E(Y_{1,0})\\ &=& E(Y(1)-Y(0)|S,G=1,T=1)P(S|G=1)\\ &+& (E\left(Y(1)|D(0)=1,G=1,T=1\right)-E\left(Y(1)|D(0)=1,G=1,T=0\right))P(D(0)=1|G=1)\\ &+& (E\left(Y(0)|D(0)=0,G=1,T=1\right)-E\left(Y(0)|D(0)=0,G=1,T=0\right))P(D(0)=0|G=1). \ \ (1.27) \end{split}$$

Then,

$$\begin{split} \delta_1 &= E(Y_{1,0,1}) - E(Y_{1,0,0}) \\ &= E(Y(1)|D(1) = 1, G = 0, T = 1) - E(Y(1)|D(0) = 1, G = 0, T = 0) \\ &= E(Y(1)|D(0) = 1, G = 0, T = 1) - E(Y(1)|D(0) = 1, G = 0, T = 0). \end{split} \tag{1.28}$$

• Last equality follows from $E(D_{01})=E(D_{0,0})+\mathsf{A2}$ (treatment monotonicity) $\implies \{G=0,D(1)=1\}=\{G=0,D(0)=1\}$

Similarly

$$\delta_0 = E(Y(0)|D(0) = 0, G = 0, T = 1) - E(Y(0)|D(0) = 0, G = 0, T = 0).$$
 (1.29)

• the result follows combining Equations 1.27, 1.28, 1.29 and Assumption 4 (conditionnal common trend) once we note that:

$$P(D(0)=1|G=1)=P(D=1|G=1,T=0)$$
 and $P(S|G=1)=P(D=1|G=1,T=1)-P(D=1|G=1,T=0)$

Time-Corrected Wald Estimand: Intuition

$$W_{TC} = \frac{E(Y|G=1, T=1) - E(Y + (1-D)\delta_0 + D\delta_1|G=1, T=0)}{E(D|G=1, T=1) - E(D|G=1, T=0)}$$

- This is almost the Wald ratio in the treatment group with time as the instrument
- ullet except that we have $Y+(1-D)\delta_0+D\delta_1$ instead of Y in 2nd term
- This difference arises because time is not a standard instrument: it can directly affect the outcome
- When the treatment rate is stable in the control group, we can identify the trends on Y(0) and Y(1) by looking at how the mean outcome of untreated and treated units changes over time in this group
- Under Assumption 4, these trends are the same in the 2 groups



Time-Corrected Wald Estimand: Intuition

- We can add these changes to the outcome of untreated and treated units in the treatment group in period 0,
- and recover the mean outcome we would have observed in this group in period 1 if switchers had not changed their treatment between the 2 periods
- This is what $Y + (1 D)\delta_0 + D\delta_1$ does
- ullet the numerator of W_{TC} compares the mean outcome in the treatment group in period 1 to the counterfactual mean we would have observed if switchers had remained untreated
- This gives LATE after a normalization

Time-Corrected Wald Estimand: extentions

- Theorem 3 assumes that $E(D_{0,1}) = E(D_{0,0})$
- When the share of treated units is not stable over time in the control group, $\Delta_{1,1}^S$ is only partially identified if Y is bounded
- One can therefore estimate the bounds of $\Delta_{1,1}^S$ (See De Chaisemartin & D'Haultfoeuille (RESTUD, 2018))
- Partial identification is still very useful for applied economist, specially when the identified set is compact
- For a scalar parameter, it makes a lot of sense, when you think of how it relates to confidence bands in the fully identified setting

Change-in-Changes Estimands in Fuzzy DID

23 / 30

Fuzzy CIC

- This is an alternative estimator to the Wald estimators in Fuzzy DID setting
- Assumption 5: Monotonicity and time invariance of unobservables $Y(d) = h_d(U_d,T)$ with $U_d \in \mathbb{R}$ and $h_d(u,t)$ strictly increasing in u for all $(d,t) \in \mathcal{S}((D,T))$. Moreover, $U_d \perp T | G, D(0)$
- Assuption 2 (treatment monotonicity) and 5 generalize the CIC model in Athey & Imbens (2006)
- They both imply $U_d \perp T|G$: they require that at each period, both potential outcomes are strictly increasing functions of a scalar unobserved heterogeneity term
- whose distribution is stationary over time, as in Athey & Imbens (2006)
- Assumption 5 also imposes that $U_d \perp T|G,D(0)$: the distribution of U_d must be stationary within subgroup of units sharing the same treatment status at T=0

Fuzzy CIC

Assumption 6: Data restrictions (testable)

- $\mathfrak{S}(Y_{d,g,t}) = \mathcal{S}(Y) \text{ for } (d,g,t) \in \mathcal{S}((D,G,T)) \text{ and } \mathcal{S}(Y) \text{ is a closed interval of } \mathbb{R}$
- $\textbf{9} \ \ F_{Y_{d,g,t}} \ \text{is continuous on} \ \mathbb{R} \ \text{and strictly increasing on} \ \mathcal{S}(Y) \ \text{for} \\ (d,g,t) \in \mathcal{S}((D,G,T))$
 - 1st condition requires that the outcome have the same support in each of the eight treatment \times group \times period cells as in Athey & Imbens (2006)
 - The 2nd condition requires that the distribution of Y be continuous with positive density in each of the eight cells
- Consider $Q_d(y) = F_{Y_{d,0,1}}^{-1} \circ F_{Y_{d,0,0}}(y)$ the quantile-quantile transform of Y from period 0 to 1 in control group conditional on D=d
- \bullet This transform maps y at rank q in period 0 into the corresponding y' at rank q in period 1



Fuzzy CIC

Let's also define

$$F_{CIC,d}(y) = \frac{P(D_{1,1} = d)F_{Y_{d,1,1}}(y) - P(D_{1,0} = d)F_{Q_d(Y_{d,1,0})}(y)}{P(D_{1,1} = d) - P(D_{1,0} = d)}$$

$$W_{CIC} = \frac{E(Y_{1,1}) - E(Q_{D_{1,0}}(Y_{1,0}))}{E(D_{1,1}) - E(D_{1,0})}$$

- Theorem 4: If assumptions 5 and 6 are satisfied, and if $E(D_{0,1})=E(D_{0,0})$, then $W_{CIC}=\Delta_{1,1}^S$ and $F_{CIC,1}^{-1}(q)-F_{CIC,0}^{-1}(q)= au_q$
- ullet This result combines ideas from Imbens & Rubin (1997) and Athey & Imbens (2006)

Fuzzy CIC: Intuition

- \bullet We seek to recover the distribution of, say, Y(1) among switchers in the treatment group \times period 1 cell
- Let's start from the distribution of Y among all treated observations of this cell: both switchers and units already treated at T=0
- ullet We need to withdraw from this distribution, that of units treated at T=0, but it's not observed
- To construct it, we can apply the quantile-quantile transform from period 0
 to 1 among treated observations in the control group to the distribution of Y
 among treated units in the treatment group in period 0.
- In a way, this quantile-quantile transform uses a double-matching to reconstruct the unobserved distribution

Fuzzy CIC: Intuition

- ullet Consider a treated unit in the treatment group imes period 0 cell. She is 1rst matched to a treated unit in the control group imes period 0 cell with same y
- Those two units are observed at the same period of time and are both treated.
- ullet Under Assumption 5 they must have the same u_1
- ullet Second, the control imes period 0 unit is matched to her rank counterpart among treated units of the control group imes period 1 cell
- ullet Denote by y^* the outcome of this last observation
- $U_1 \perp T | G, D(0) = 1 \implies$ those two observations must also have the same u_1
- So $y^* = h_1(u_1, 1)$: outcome that treatment \times period 0 cell unit would have obtained in period 1



W_{CIC} can be re-written as

$$W_{CIC} = \frac{E(Y|G=1,T=1) - E((1-D)Q_0(Y) + DQ_1(Y)|G=1,T=0)}{E(D|G=1,T=1) - E(D|G=1,T=0)}$$

- Here again, W_{CIC} is almost the standard Wald ratio in the treatment group with T as the instrument
- ullet Except that we have $(1-D)Q_0(Y)+DQ_1(Y)$ instead of Y in second term
- it accounts for the fact that time directly affects the outcome, just as $(1-D)\delta_0 + D\delta_1$ does in the W_{TC} estimand
- Under Assumption 4, the trends affecting the outcome are identified by additive shifts
- Under Assumption 5 and 6, they are identified by possibly non-linear quantile-quantile transforms

Next session

- DID with Variation in Treatment Timing
- Read: Goodman-Bacon, Andrew. "Difference-in-differences with variation in treatment timing." Journal of Econometrics (2021)
- optional: De Chaisemartin, Clement and Xavier D'Haultfoeuille. "Two-Way Fixed Effects Estimators with Heterogeneous Treatment Effects". AER(2020)