

*Interpreting Did and IV Estimates:
ATE, ATET, LATE, and All That*

Bernard Black

Univ of Texas (through 2010)

Northwestern (from 09.01.2010)

Law School & Kellogg School of Management

CELS, 2009

Main goal

- Assume: You have a (reasonably) **valid** instrumental variable (IV) or difference-in-differences (DiD) study design
 - How to achieve that: different talk(s)
- This provides evidence of causation for a **subpopulation**
- Goal here: What can you learn from the study
 - Hint: Less than you might think

Proceed by Example

- See what we learn from:
 - One of my wife's studies
 - Litvak, *The Impact of the Sarbanes-Oxley Act on Non-US Companies Cross-Listed in the US* (J Corp Fin 2007)
 - One of my own studies
 - Black, Jang & Kim, *Does Corporate Governance Affect Firms' Market Values? Evidence from Korea* (JLEO 2006)
 - Two classic IV studies
 - Angrist & Kreuger, *Does Compulsory School Attendance Affect Schooling and Earnings?* (QJE 1991)
 - McClellan, McNeil & Newhouse, *Does More Intensive Treatment of Acute Myocardial Infarction in the Elderly Reduce Mortality?* (JAMA 1994)

Simple setup (0-1 “treatment”)

- Treatment is “on” ($D_i = 1$) or “off” ($D_i = 0$)
 - *i indexes the subjects of the treatment: people or firms or states*
- *We are interested in how treatment affects outcome y_i for subject i*
 - $[y_i | D_i = 1] - [y_i | D_i = 0]$
 - Or, more compactly: $y_{1i} - y_{0i}$**
- *Any subject is treated or not*
- *So we observe one of these, but not the other*
 - *Regression analysis lets us estimate the unobserved outcome*

Regression design

- OLS: $y_i = \alpha + \beta^* D_i + \boldsymbol{\gamma}^* \mathbf{X}_i + \varepsilon$
 - \mathbf{X} = set of control variables
 - $\underline{\beta}$ is estimate of $E[y_{1i} - y_{0i}]$
- This is association, not causation. Could have:
 - **reverse causation**: y causes D
 - **omitted variable**: both y and D are associated with an unobserved variable w
 - if so, regression estimate $\underline{\beta}$ is biased
- IV and DiD provide ways to get unbiased estimate

Litvak (2007) research design

- “natural” experiment: SOX applies to all US firms (so no within-US control group), and to **some** foreign firms
 - SOX applies, if firm is cross-listed in US on level 2 or 3
 - Does not apply if firm is cross-listed on level 1 and 4
 - (Obviously) does not apply if firm is not cross-listed
- *DiDiD* design:
 - stock price return during SOX adoption events in 2002 (first difference)
 - “pair return”: return to level-23 firm relative to propensity matched non-cross-listed firm from same country (second “pair” difference)
 - $n = 431$ (this will become important later)
 - to level-23 pairs relative to level-14 pairs (third difference)

Litvak main result

- Negative return to level-23 pairs versus level-14 pairs: $\sum(\text{SOX events}) = -5.8\%$ ($t = 5.48$)
- Assume investors are right in predicting effect of SOX on firm value
 - apparently valid through 2005 (see Litvak 2008).
- What can we learn (and not learn)?

Logic of research design

- Narrow event study windows → stock price return more likely due to SOX
- **Pair** return controls for unobserved developments in home country
- Difference in level-23 and level-14 pair returns controls for unobserved US developments that affect **all** cross-listed firms
- But:
 - limited control variables
 - limited data availability, especially for non-cross-listed firms
 - must assume: no relevant unobserved difference between level-23 and level-14 pairs
 - usual event study assumptions (smart investors, efficient markets)

What inferences can we draw?

- All level-23 firms were “treated” by SOX
- Mean -5.8% difference in pair returns is estimate of **Average Treatment Effect (ATE)**
- In symbols: $ATE = E[y_{1i} - y_{0i}]$
 - E = expectation operator
 - y_{1i} = value of firm i if treated by SOX ($D_i = 1$)
 - y_{0i} = value of firm i if **not** treated by SOX ($D_i = 0$)
not observed; estimated using the DiDiD procedure
- US firms were treated too . . .
 - Can we estimate a -5.8% reaction to SOX for these firms?

ATE for whom?

- On the **treated population**
 - Litvak studies only level-23 foreign firms
 - no assurance that **observed** ATE on level-23 firms = **unobserved** treatment effect on US firms
 - Compare Zhang (2008) (event study of US firm reactions, using non-US firms as control)
 - advantage: measures ATE for US firms
 - concern: weaker control group → unobserved differences between treatment and control groups

ATET (Average Treatment Effect on the Treated)

- **ATET = ATE on the treated**
- Formally: $ATET = E[(y_{1i} - y_{0i}) | D_i = 1]$
- If whole population is treated, then $ATE = ATET$
 - For Litvak's study, $ATE = ATET$
 - But **ATET** terminology is still useful: suggests need to focus on who is treated

How about other foreign firms?

- Level-14 firms were **not** treated
 - Technical answer: Strictly speaking, we don't know ATE
 - Practical question: Are they similar enough to level-23 firms to infer SOX impact, ***if they had been treated?***
 - No clear answer, must study how they differ.
- Non-cross-listed matched firms?
 - Matching procedure ensures they are similar to level-23 firms in propensity to cross-list
 - Does that imply similar reaction to SOX?
- Other non-cross listed firms?
 - Less similar to level-23 firms → less likely that one can infer hypothetical effect if treated

Black, Jang and Kim

- IV study (2006); DiD study (2009, with Park)
- Large Korean firms (assets > 2 trillion won) are subject to legal shock in 1999. Must have:
 - 50% outside directors
 - audit committee, majority of outside directors
 - outside director nomination committee
- Small firms are not treated, form control group
- Core result: Large firm Tobin's q rises by 0.16 ($t = 3.86$) relative to small firms

Korea Study: ATE and ATET

- All large firms are treated:
 - ATE = ATET = 0.16 increase in Tobin's q
- *What can we infer about value of these governance reforms for **small** firms?*
 - Technically, nothing (they are not treated)
 - *But if similar to large firms, could see similar effect*
 - *mid-sized (say 0.5-2T won): likely similar effects*
 - *smaller firms: less clear*
 - *BJK report similar coefficients for large firms and for small firms who are voluntary adopters*

Angrist & Krueger on school dropouts

- Core idea: dropouts allowed at age 16 (say)
- Some students are closer to graduation when they turn 16 (depends on birth date)
 - Assume start K if age 5 by Jan. 1 of next year
 - Each K class includes ages 4.08-5.07 (year.months)
 - 4.08 kids will be almost halfway through grade 11 when they turn 16
 - 5.07 kids will turn 16 about halfway through grade 10
 - Predict (and observe): more schooling/fewer dropouts if younger when start K

Birthdate as instrument for schooling

- Classic problem in labor economics: impact of schooling on future income
- Want to estimate:
- $\text{income} = \alpha + \beta * (\text{years of school}) + \boldsymbol{\nu} * \mathbf{X} + \delta * \text{ability} + \varepsilon$
- But can't observe ability, so actually estimate:
- $\text{income} = \alpha + \beta * (\text{years of school}) + \boldsymbol{\nu} * \mathbf{X} + \varepsilon$
 - omitted variable problem
 - estimate of β biased **upward** if ability \rightarrow schooling
- need instrument for schooling, uncorrelated with ability
 - birthdate, relative to K cutoff date, is plausible instrument

More complex setup (still 0-1 variables)

- A treatment is “on” ($D_i = 1$) or “off” ($D_i = 0$)
 - Kids enter K younger ($D_i = 1$) or older ($D_i = 0$)
- *Treatment affects outcomes: Younger kids stay in school longer, on average*
 - $E[\text{school years}_{1i}] > E[\text{school years}_{0i}]$
- Treatment never causes reverse response
 - $\text{school years}_{1i} \geq \text{school years}_{0i}$ for all i
 - *No one gets less school because entered K younger*

Instrument for whom?

- Put aside “weak instrument” problem with their study (not known at the time):
 - Estimated 4-6% higher earnings per year of school
 - Not far from OLS estimates with good controls
- Who is affected by the “treatment”?
 - not your kids or mine, who will graduate HS anyway
 - Let $G_i = 1$ if graduate; 0 otherwise
 - Observe: $[\text{school years}_i | G_i = 1]$ is **independent** of D_i
 - instead marginal kids, who would have dropped out if they were older at entry $(G_i = 0 | D_i = 0) \approx G_{0i} = 0$
 - some graduate if $D_i = 1$; some still don't

Don't know ATE or ATET

- No reason to believe value of schooling is same for both groups
- → don't know ATE = average effect on all kids of being “treated” (being young when enter K)
 - don't know ATET either (average treatment effect on kids who were young when enter K)
- What do we know?

LATE

- We know the **local** average treatment effect (LATE) – ATE for the marginal kids
- Formally: We observe, through the 2SLS procedure:
- $LATE = E[(y_{1i} - y_{0i}) | G_{0i} = 0]$
 - $G_{Di} = 1$ if graduate; 0 if drop out
 - y_{1i} = earnings if treated (if young when enter K)
 - y_{0i} = earnings if not treated (older when enter K)
- Can think of this as *ATE on the effectively treated (subpopulation of treated for whom treatment matters)*

Is This Useful to Know?

- Not if your policy concern is the average return to extra education for all kids.
- But suppose you are a state legislator, considering changing the minimum school leaving age, or the age at school entry?
 - You might care about effect on the marginal kids

McClellan et al. on AMI treatment

- Observe:
 - **huge** difference in AMI (heart attack) survival for who receive cardiac catheterization ($D_i = 1$), and those who don't
 - 37% higher 4-year survival (67% vs. 30%)
- But also observe:
 - patients who receive catheterization are younger and healthier than those who don't → better outcomes in any case
- Survival difference shrinks if control for observable health markers, still large
 - could reflect unobserved health differences
 - could reflect unobserved treatment differences
 - cath-equipped hospitals might be better at non-cath treatment

Instrument: Distance to cath hospital

- Observe: only major hospitals perform cath
- AMI victims usually taken to nearest hospital
 - some later transfer to cath hospital, some don't
 - let $b_i = 1$ if **extra distance** from home to cath hospital, versus nearest hospital < 2.5 miles
 - $E[D_i | b_i = 1] > E[D_i | b_i = 0]$
 - 26.2% vs. 19.5% (within 90 days)
- Observable health markers for AMI victims are independent of b_i
- Use b_i as instrument for D_i

McClellan et al. results

- 4-year survival:
- $E[y_i | b_i=1] - E[y_i | b_i=0] = 58.5\% - 58.1\% = 0.4\%$
- Implies that $D_i = 1$ improves survival by:

$$\text{effect of cath on survival} = \frac{\Delta(\text{survival})}{\Delta(\text{cath rate})} = \frac{0.4\%}{.067} = 6.0\%$$

$$E[y_i | D_i=1] - E[y_i | D_i=0] = \frac{E[y_i | b_i=1] - E[y_i | b_i=0]}{E[D_i | b_i=1] - E[D_i | b_i=0]}$$

Sometimes called a “Wald” estimate

This is LATE -- on whom?

- Who is affected by background condition ($b_i = 0$ or 1)?
- Some patients will get cath anyway
 - Those who benefit the most, one hopes
- Some won't get cath anyway
 - Hopefully those who won't benefit (too sick already, otherwise not indicated)
- Distance to cath hospital affects **marginal** patient
 - gets cath only if close to cath hospital

Terminology: always takers; never takers, compliers

- **Always takers:** get treatment ($D_i = 1$) no matter what
- **Never takers:** no treatment ($D_i = 0$) no matter what
 - For both: D_i indep. of b_i
- **Compliers: Treatment depends on background**
 - $D_i = 1$ if and only if $b_i = 1$
- LATE (6.0% higher survival) is average effect on **compliers** (on the marginal patients)
- Terms developed for partly voluntary action
 - e.g., enroll in training program if eligible; complete diet or exercise program; serve in Army if drafted
 - but apply here too

More terminology

- Clinical medical trial (low-fat diet, say):
 - Sign up 2,000 people; half get “treated”; half are controls
 - Only some complete the program
 - Dropouts are **not** random
 - So ATE for those who complete treatment is endogenous and biased.
- Use **intention to treat** ($b_i = 1$) as instrument for actual treatment

LATE with intent to treat

- Wald estimate of LATE

$$LATE = \frac{\Delta(\text{expected outcome based on intent-to-treat})}{\Delta(\text{likelihood of treatment based on intent-to-treat})}$$

- Compare McClellan AMI study:
 - “intent”: treat those close to cath hospital with cath
 - But some who lived close ($b_i=1$) got no cath
 - And some who lived farther away ($b_i = 0$) got cath
 - Denominator is difference in cath rate between these two groups.

LATE versus ATET

- ΔTET is average effect on those who are
- $ATET =$ weighted average of LATE and (effect on always takers)

Instrument Validity

- Standard, and large, problem with instrument validity: There is no test!
- Valid instrument must be (i) exogenous; (ii) correlated, preferably strongly, with instrumented variable, and (iii) predict dependent variable only **indirectly**, through the instrumented variable
 - (i) is sometimes easy
 - (ii) can be estimated
 - (iii) [called the “**exclusion restriction**”] is the hardest to satisfy, and there is no test, only logic.

Difference in Hansen test

- There is, however, a Hausman test for endogeneity, **assuming** instrument is valid
 - Is IV coefficient estimate significantly different than OLS estimate
 - Caveat: weak instrument → weak test
- And a Hansen “test of overidentifying restrictions”: validity of two or more instruments **assuming** one is valid
 - does IV coefficient estimate change significantly when add second instrument
 - and “difference in Hansen” test when add third instrument

What can these tests really tell you?

- Hausman: Is there evidence of endogeneity for the **compliers**?
- Hansen: Can fail this test if:
 - (i) second instrument is not valid, or
 - (ii) different instruments operate on different subpopulations with different LATEs
 - Test can't distinguish between these
 - two effects could offset, leading to false comfort with multiple instruments

Example of different subpopulations

- Effect of additional child on mother's earnings
 - focus on 3+ kids versus 2 (Angrist & Evans, 1998)
 - instrument 1: first two kids of same sex
 - weakly predicts greater probability of third child
 - instrument 2: second birth is twins
 - strongly predicts greater probability of third child
- What's your prediction:
 - Will treatment effects be similar?
 - If different, which should be larger?

Same sex vs. twins estimates

- Married women, age 21-35, lots of controls
- Same sex estimate on probability of working:
–13.5% [95% CI = 7.7-19.3%]
- Twins estimate:
–8.3% [95% CI = 4.9-11.7%]
- (Probably) different, with expected sign
- If twins are random event,
 - Twins estimate is close to ATET = ATE
 - But on whom?: Effect on whom?
 - women with one child, and desire or willingness to have 2+
- Same sex estimate is LATE

Wrapup

- Limitations of LATE are frustrating
 - Example: Benmelech (2009): railroad gauge in 1800s as instrument for asset specificity of railroad care
 - test effect of asset specificity on capital structure
 - generalization to other assets?
 - generalization to today's financial markets??
 - Maybe still “Better LATE than Nothing” (Imbens, 2009)
- Lots of complexity left untouched:
 - variable intensity treatments (not just 0-1)
 - adding control variables