



Lecture 12 - RDD.pdf

Regression Discontinuity

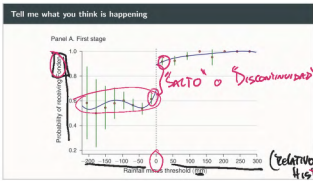
Mauricio Romero

Introduction

What is regression discontinuity design (RDD)?

- Donald Campbell, educational psychologist, invented regression discontinuity design but then it went dormant for decades
- Angrist and Pischke (1999) and Black (1999) independently rediscovered it
- It has become incredibly popular in economics

DICT
 RDD
 DID
 OLS ~ "hardwired"



Running example from Mexico 1

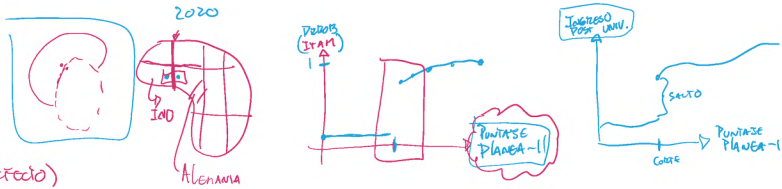
Rules for Recovery Impact of Indexed Disaster Funds on Shock Coping in Mexico?

By Agustín Rivera-Velazco, Juan Carlos Escobar-Castillo

Government provision of disaster relief is typically hampered by quality constraints such as need and administrative capacity. In addition, discretionary decisions by those that provide relief may create selection bias. Mexico's indexed disaster fund (IDF) controls both of these concerns: receipt of disaster relief is automatic based on receipt of a natural disaster, and the amount of relief is indexed to the amount of damage. In this paper, we use the IDF to estimate the causal effect of disaster relief on shock coping in Mexico. We use a regression discontinuity design to estimate the causal effect of a self-selected treatment on shock coping. We find that the IDF has a positive and significant effect on shock coping. The magnitude of the effect is similar to that of the control group. Our findings suggest that disaster relief programs should be designed to be automatic and indexed to the amount of damage.

What is a regression discontinuity design?

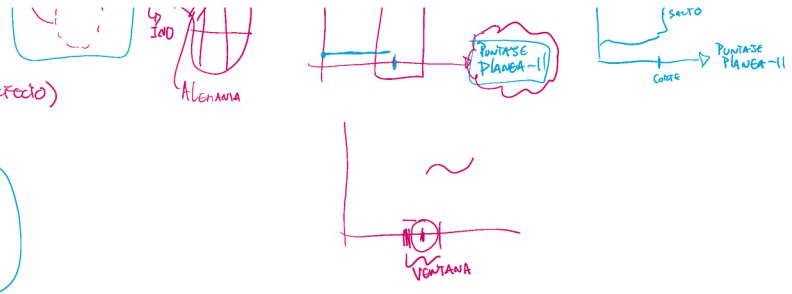
- Goal: estimate some causal effect of a treatment on some outcome
- Problem: Selection bias (i.e., $E[Y^1|D=1] \neq E[Y^0|D=0]$)
- RDD basic idea: if treatment assignment occurs abruptly when some underlying variable X (the "running variable") passes a cutoff c , then we can use that arbitrary rule to estimate the causal effect even of a self-selected treatment



• Problem: Selection bias (i.e., $E[Y^0|D=1] \neq E[Y^0|D=0]$)

• RDD basic idea: If treatment assignment occurs abruptly when some underlying variable X (the "running variable") passes a cutoff c_0 , then we can use that arbitrary rule to estimate the causal effect even of a self-selected treatment

(No sea Perfeccio)



Arbitrary rules

- Firms, schools and gov agencies assign "things" based on arbitrary thresholds of continuous variables
- Consequently, probabilities of treatment will "jump" when that running variable exceeds a known threshold
 - Academic test scores, scholarships or prizes, higher education admission, certification of merit
 - Poverty scores: (poor) income-based anti-poverty programs (generally: any program targeting that extreme poverty of households)
 - Land area: fertilizer program or debt relief initiative for owners of plots below a certain size
 - Child age cutoffs for pensions; date of birth for starting school with different cohorts; date of loan to determine eligibility for debt relief
 - Election: fraction that voted for a candidate of a particular party

Selection examples and solutions from the literature

- Think of these in light of a treatment where $E[Y^0|D=1] \neq E[Y^0|D=0]$
- Yelp rounded a continuous score of ratings to generate stars
 - US targeted air strikes in Vietnam using rounded risk scores
 - Universal healthcare after age 65
 - When a newborn's birthweight is below 1500 grams it gets intensive medical care

CONTINUOS → DISCRETOS

*... → 7, 8, 9, 10, 11, 12, 13, 14, 15

"EFFECTO 4 o 5 ESTRELLAS"

(3.5, 4.4) → 4
(4.5, 5) → 5

(3.49) vs (3.5)

Sharp vs. Fuzzy RDD

- There's traditionally thought to be two kinds of RD designs:
 - Sharp RDD: Treatment is a deterministic function of running variable, X
 - Fuzzy RDD: Discontinuous "jump" in the probability of treatment when $X > c_0$. Cutoff is used as an instrumental variable for treatment
- Fuzzy is a type of IV strategy and requires explicit IV estimators like 2SLS

Sharp Design

Treatment assignment in the sharp RDD

Deterministic treatment assignment ("Sharp RDD")

In Sharp RDD, treatment status is a deterministic and discontinuous function of a covariate, X_i :

$$T_i = \begin{cases} 1 & \text{if } X_i \geq c_0 \\ 0 & \text{if } X_i < c_0 \end{cases}$$

where c_0 is a known threshold or cutoff. In other words, if you know the value of X_i for a unit i , you know treatment assignment for unit i with certainty.

Treatment effect definition and estimation

Definition of treatment effect

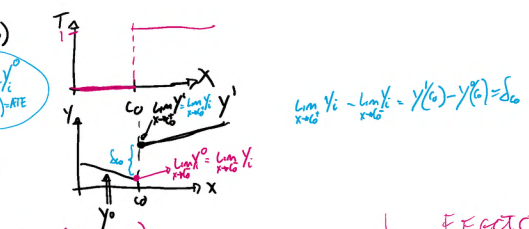
The treatment effect δ is the discontinuity in the conditional expectation function:

$$\delta = \lim_{x \rightarrow c_0^+} E[Y|X=x] - \lim_{x \rightarrow c_0^-} E[Y|X=x] = E[Y^1|X=c_0] - E[Y^0|X=c_0] = \delta(\kappa=c_0)$$

Average causal effect of the treatment at the discontinuity

$$\tau = E[Y^1] - E[Y^0] = \delta(\kappa=c_0)$$

T is correlated with X and deterministic function of X , overlap only occurs in the limit and thus the treatment effect is in the limit as X approaches c_0 .



```
## Basic RD Model
N=1000 # Number of observations
T=runif(N, 0, 1) # Treatment
X=runif(N, 0, 1) # Running variable
Y0=0.5 + 0.5*X # Control potential outcome
Y1=0.5 + 0.5*X + 0.2 # Treatment potential outcome
# Data matrix
dat = data.frame(X=X, T=T, Y0=Y0, Y1=Y1)
# Fit the model
fit = lm(Y ~ X + T, data=dat)
# Coefficients
beta0 = fit$coefficients["(Intercept)"]
beta1 = fit$coefficients["X"]
delta0 = fit$coefficients["T"]
# ATE = 2
ATE = beta0 + beta1 + delta0
```

$Y_0 = N(X, sd=1)$
 $Y_1 = N(X+Z, sd=1)$
 $\delta_0 = c_0 = 0$
 $\tau = N(Y_1 - Y_0 | X=c_0) = 2 - 1 = 1$

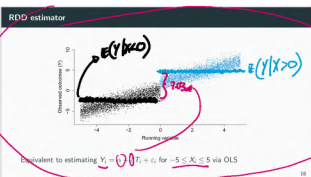
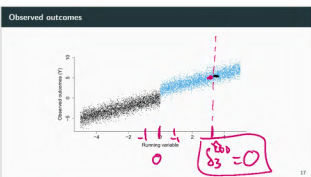
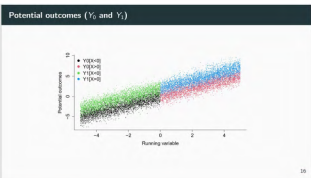
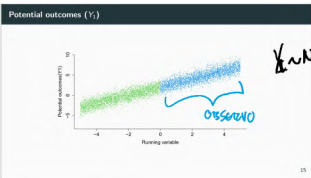
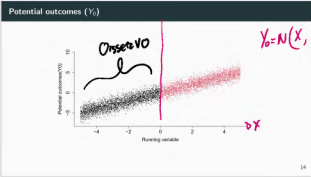
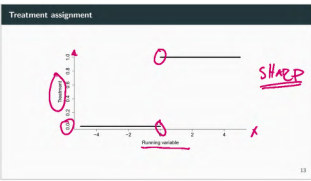
EFFECTO No POTERIZ ECO IV

$\tau = Y(c_0) - Y_0(c_0)$

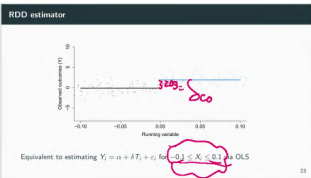
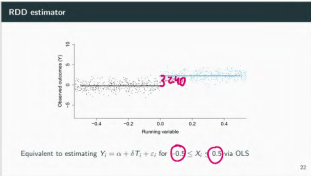
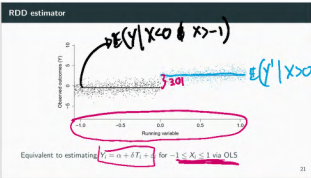
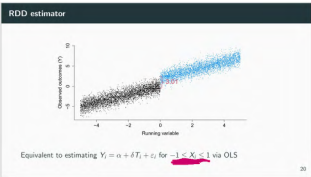
$\tau = Y(c_0) - Y_0(c_0) = 2 - 1 = 1$

Treatment assignment
 $Y_i = N(X_i, Z_i, sd=1)$
 $Y_{0i} = N(X_i, 0, sd=1)$
 $Y_{1i} = N(X_i, 1, sd=1)$

$Y_i \sim N(X_i, Z_i, sd=1)$
 $\sigma_0 = \sigma_1 = 0$
 $\sigma_1 = 1 = E(Y_i | Z_i=1) - Y_0(X_i, 1) = Z_i - 1$
 $\sigma_2 = Z_i$



Equivalent to estimating $Y_i = \beta_0 + \beta_1 X_i + \beta_2 D_i + \epsilon_i$ for $-5 \leq X_i \leq 5$ via OLS



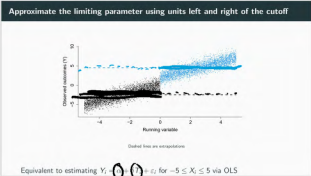
T_i si $Y_i > Y_0$

Como DETERMINAR VELOCIDAD \Rightarrow PROBLEMA "BIAS-VARIANCE TRADEOFF" \Rightarrow DENSITY

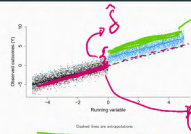
Extrapolation

- Extrapolation
- In RDD, the counterfactuals are conditional on X
 - We use extrapolation in estimating treatment effects with the sharp RDD because we do not have overlap
 - Left of cutoff, only non-treated observations, $T_i = 0$ for $X < c_0$
 - Right of cutoff, only treated observations, $T_i = 1$ for $X \geq c_0$
 - The extrapolation is to a counterfactual

No HAY Como SABER Como se COMPARA



Approximate the limiting parameter using units left and right of the cutoff



$$\delta(\alpha + \delta) + (\beta + \lambda)X_i$$

CUANDO T=c

$$\alpha + \beta X_i \quad \text{CUANDO T=0}$$

$$\frac{1}{\lambda} \times 200$$

Equivalent to estimating $Y_i = \alpha + \beta X_i + \lambda X_i + \gamma_i + \delta T_i$ for $-5 \leq X_i \leq 5$ via OLS

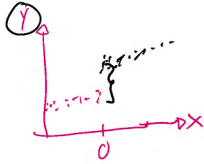
$$\begin{aligned} \gamma_i &= \alpha + \beta X_i + \lambda X_i + \delta \cdot 1 + \epsilon_i \\ &= (\alpha + \delta) + (\beta + \lambda) X_i \end{aligned}$$

Smoothness assumption

Key identifying assumption

Smoothness for continuity of conditional expectation functions (Hahn, Todd and Van der Klauw 2001)

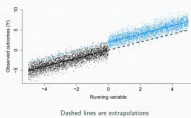
- Potential outcomes are not actual outcomes
- If population average potential outcomes, Y^1 and Y^0 , are smooth functions of X through the cutoff, c_0 , then potential average outcomes won't jump at c_0 .
- Implies the cutoff is exogenous - i.e., nothing else changes related to potential outcomes at c_0 .
- Unobservables are evolving smoothly, too, through the cutoff



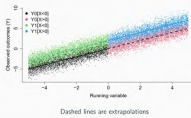
Smoothness is the identifying assumption and untestable

- The smoothness assumption allows us to use average outcome of units right below the cutoff as a valid counterfactual for units right above the cutoff
- Extrapolation is allowed if smoothness is credible, and extrapolation is non-sensical if smoothing isn't credible
- Why not directly testable? Because potential outcomes are not observable

Approximate the limiting parameter using units left and right of the cutoff



Approximate the limiting parameter using units left and right of the cutoff



Estimation

Re-centering the data

- It is common for authors to transform X by "centering" at \bar{x} . $\rightarrow \delta_{\bar{x}}$

$$10 + \beta(X - \bar{x}) = \lambda(X - \bar{x}) + \tau + \epsilon$$
- This doesn't change the interpretation of the treatment effect - only the interpretation of the intercept.

Nonlinearities

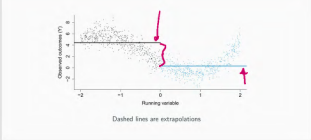
Nonlinearity bias

- Smoothness and linearity are different things.
- What if the trend relation $E[Y|X]$ does not jump at c_0 but rather is simply nonlinear?
- Then your linear model will identify a treatment effect when there isn't because the functional form had poor predictive properties beyond the cutoff.
- Let's look at a simulation

Simulations!

```
## Run linear OLS
N=1000 #number of observations
Treatment=(0,1)
X=(0,1)
#Outcome
#We only get treatment if X>0
Treatment=(X>0)
#DGP (notice there is NO treatment effect)
Y=X^2+epsilon
#Constant Model
Const=lm(Y~Treatment)
Const2=lm(Y~Treatment,subset=abs(X)<1)
Const3=lm(Y~Treatment,subset=abs(X)>0.5)
Const4=lm(Y~Treatment,subset=abs(X)>1)
```

Non-Linear



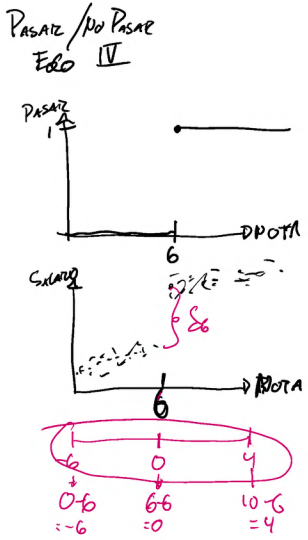
	(1)	(2)	(3)	(4)
Treatment	4.13***	-3.64***	-2.03***	-0.14*
Constant	4.45***	3.18***	2.02***	3.24***
Sample	1000	208	243	68
Observations	1000	208	243	68

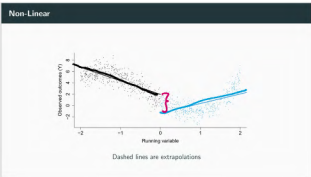
Note: *p<0.1; **p<0.05; ***p<0.01

Simulations!

$Y \sim \beta_0 + \beta_1 X + \lambda X T + \delta T + \epsilon$

```
##Linear Model
Linear=lm(Y~Treatment)
Linear2=lm(Y~Treatment+X*Treatment,subset=abs(X)<1)
Linear3=lm(Y~Treatment+X*Treatment,subset=abs(X)>0.5)
Linear4=lm(Y~Treatment+X*Treatment,subset=abs(X)>1)
```





	(1)	(2)	(3)	(4)
Treatment	3.30*** (0.17)	-0.45** (0.18)	0.13 (0.25)	-1.35*** (0.48)
X	-2.39*** (0.11)	-4.21*** (0.21)	5.82*** (0.64)	8.73 (5.57)
X*Treatment	4.08*** (0.15)	2.33*** (0.30)	3.27*** (0.85)	-4.94 (8.86)
Constant	1.97*** (0.12)	0.95*** (0.13)	0.56*** (0.19)	1.03*** (0.35)
Sample Observations	Full 1,000	$ X < 1$ 508	$ X < 0.5$ 243	$ X < 0.1$ 48

Note: ***p<0.01, **p<0.05, *p<0.10

Sharp RDD: Nonlinear Case

- Suppose the nonlinear relationship is $E[Y^0|X] = f(X)$ for some reasonably smooth function $f(X)$.
- In that case we'd fit the regression model $Y_i = f(X_i) + \delta T_i + \epsilon_i$
- There are 2 common ways of approximating $f(X)$.

Nonlinearities

"higher order polynomials" but problematic due to overfitting. Gelman and Imbens 2019 recommend at best a quadratic.

- Use global and local regressions with $f(X)$ equaling a 2nd order polynomial. $Y_i = \alpha_0 + \alpha_1 X_i + \alpha_2 X_i^2 + \epsilon_i$
- Or use some nonparametric kernel method (we won't cover that)

General case

Different polynomials on the 2 sides of the discontinuity

- We can generalize the function $f(x)$ by allowing it to differ on both sides of the cutoff by including them both individually and interacting them with T_i .
- In that case we have:

$$E[Y^0|X] = \alpha_0 + \alpha_1 X + \alpha_2 X^2 + \dots + \alpha_p X^p$$

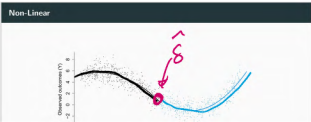
$$E[Y^1|X] = \alpha_0 + \delta + \alpha_1 X + \alpha_2 X^2 + \dots + \alpha_p X^p$$
 where X_i is the centered running variable (i.e. $X_i - \alpha_0$).
- Re-centering at α_0 ensures that the treatment effect at $X_i = \alpha_0$ is the coefficient on T_i in a regression model with interaction terms.

Different polynomials on the 2 sides of the discontinuity

- To derive a regression model, first note that the observed values must be used in place of the potential outcomes:

$$E[Y^1|X] = E[Y^0|X] + (E[Y^1|X] - E[Y^0|X]) T$$
- Regression model you estimate is:

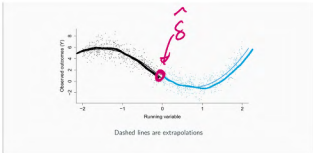
$$Y_i = \alpha_0 + \alpha_1 X_i + \alpha_2 X_i^2 + \dots + \alpha_p X_i^p + \delta T_i + \epsilon_i$$
 where $\alpha_0 = \beta_0 - \beta_1$, $\alpha_1 = \beta_1 - \beta_2$ and $\alpha_2 = \beta_2 - \beta_3$.
- The treatment effect at α_0 is δ .



SP #
Merge (X, Y, BY = "llaves", ALL=1)

Y llaves 1 2 3 4 5
X llaves 4 5 6 7 8
Merge llaves (4, 5)





	(1)	(2)	(3)	(4)
Treatment	0.21***	0.06	0.07	0.09
X	0.000	0.000	0.000	0.000
X2	(0.31)	(0.07)	(0.06)	(0.04)
X2	-2.20***	-1.89***	-1.87***	-2.00***
X2*Treatment	0.31	0.02	0.03	0.03
X2*Treatment	1.24***	1.07***	1.30***	1.02
X2*Treatment	0.04	0.02	0.03	0.03
X2*Treatment	5.84***	3.28***	3.79***	4.93***
Constant	0.91	0.12	0.12	0.11
Constant	0.44***	0.00***	0.18***	0.14*
Constant	0.14	0.19	0.20	0.19
Sample	F,df	X<1	X<1.5	X<1.1
Observations	1,000	500	500	500
Note			*p<0.1, **p<0.05, ***p<0.01	

Testing for violations

Robustness against what?

- Are you done now that you have your main results? No
- Your main results are only causal insofar as smoothness is a credible belief, so you need to convince the reader this is true
- You must now scrutinize alternative hypotheses that are consistent with your main results through sensitivity checks, placebo and alternative approaches

Main Challenges

- Classify your concern regarding smoothness violations into two categories:
 - Manipulation of the running variable
 - Endogeneity of the cutoff (¿MEJOR OTROS CASOS EN EL PUNTO CORTA?)
- Most robustness is aimed at building credibility around these

Manipulation of your running variable score

- Treatment is not as good as randomly assigned around the cutoff. Q, when agents can "perfectly" manipulate their running variable. This happens when:
 - The assignment rule is known in advance
 - Agents are interested in adjusting
 - Agents have the time/ability to adjust
- Since necessarily treatment assignment is no longer independent of potential outcomes, it's likely this implies smoothness has been violated

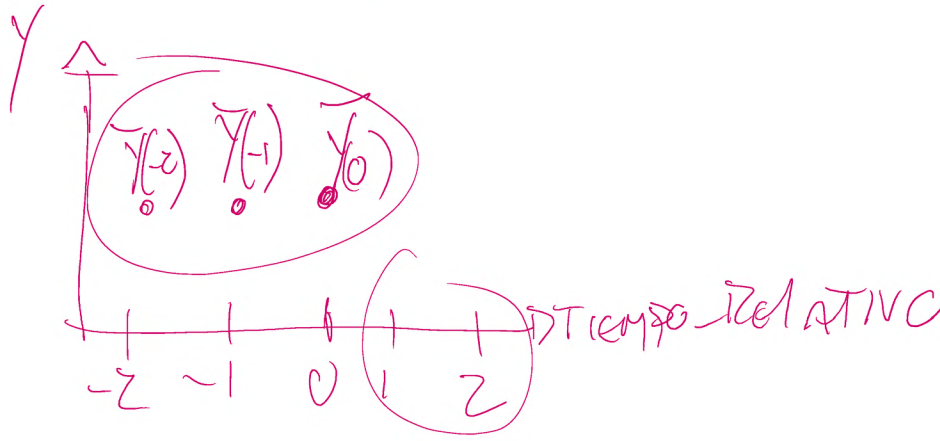
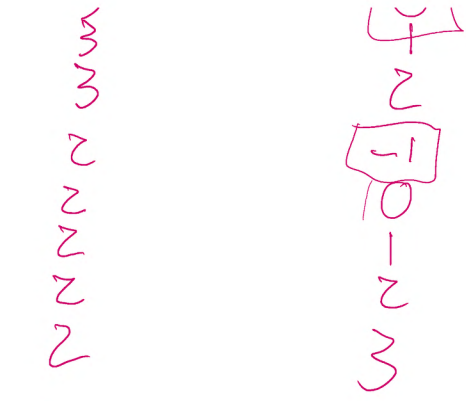
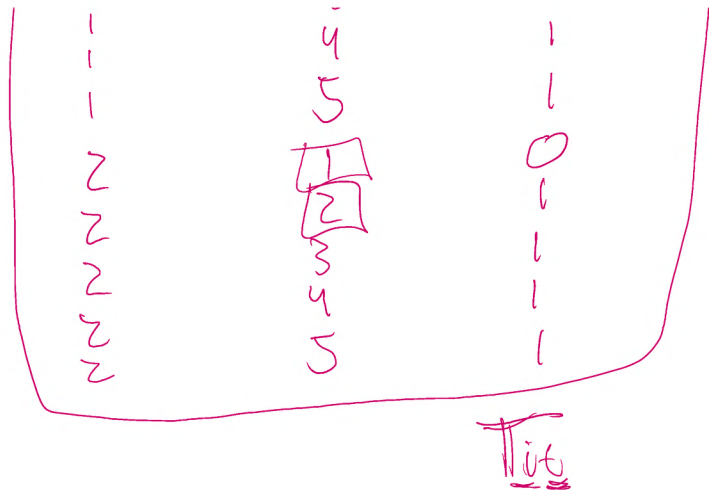
A badly designed RCT

- Suppose a doctor randomly assigns heart patients to statin and placebo to study the effect of the statin on heart attacks within 10 years
- Patients are placed in two different waiting rooms, A and B, and plans to give those in A the statin and those in B the placebo
- The doors are unlocked and movement between the two can happen

McCrary Density Test

We would expect waiting room A to become crowded. In the RDD context, sorting on the running variable implies heading on the "good side" of c_0

- McCrary (2008) test: under the null the density should be continuous at the cutoff of c_0
- Under the alternative hypothesis, the density should increase at the "good side" of c_0
 - Partition the running variable into bins and calculate frequencies in each bin
 - Treat those frequencies as dependent variable in an RDD regression



Indice INEGUALDAD
 0 | 30
 100 |
 F(y₀)

10 OBS	ID	ESTADO	AGE
1	1	0	1802
2	2	0	1802
3	3	0	1802
4	4	0	1802
5	5	0	1802
6	6	0	1802
7	7	0	1802
8	8	0	1802
9	9	0	1802
10	10	0	1802
11	11	1	1802
12	12	1	1802
13	13	1	1802
14	14	1	1802
15	15	1	1802
16	16	1	1802
17	17	1	1802
18	18	1	1802
19	19	1	1802
20	20	1	1802
21	21	1	1802
22	22	1	1802
23	23	1	1802
24	24	1	1802
25	25	1	1802
26	26	1	1802
27	27	1	1802
28	28	1	1802
29	29	1	1802
30	30	1	1802

$N = 32 \cdot 1000 = 30$
 $N_{treatment} = 1000 \cdot 30$
 $N_{control} = 32 \cdot 1000$



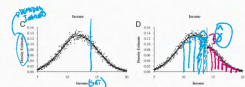
the running variable implies reaping on the "good side" of c_0

- McCrary (2008) test: under the null the density should be continuous at the cutoff
- Under the alternative hypothesis, the density should increase at the "good side" of c_0
 1. Partition the running variable into bins and calculate frequencies in each bin
 2. Treat those frequency counts as dependent variable in an RDD regression
- You need no jump to "pass" this test

McCrary density test

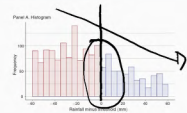
- The McCrary Density Test has become mandatory for every analysis using RDD.
- You can install `rddensity` for Stata/R, and it will implement the test

McCrary density test

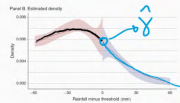


Panel C: density of income when there is no pre-announcement and no manipulation. Panel D: the density of income when there is pre-announcement and manipulation. From McCrary (2008).

McCrary density test – FONDEN running example



McCrary density test – FONDEN running example



Caveats about McCrary Density Test

- For RDD to be useful, you need to know something about the mechanism generating the running variable and how susceptible it could be to manipulation
- A discontinuity in the density is "suspicious" – it suggests manipulation of X around the cutoff is probably going on. In principle one doesn't need continuity
- This is a data-hungry test. You need a lot of observations at c_0 to distinguish a discontinuity from noise

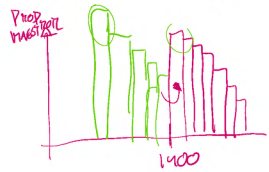
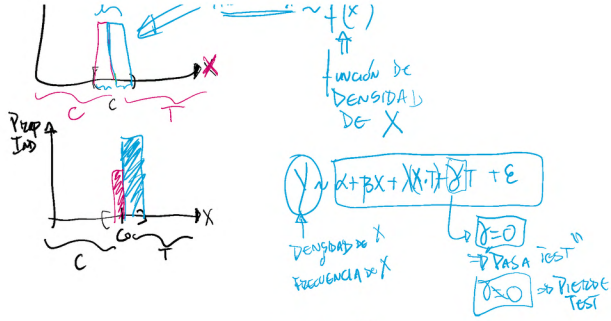
Visualizing manipulation — Proxy means test in Colombia

American Economic Journal, Economic Policy 1 (Fall 2012), 41-61
<http://www.aeaweb.org/economicpolicy/papers/1212/12121212.pdf>

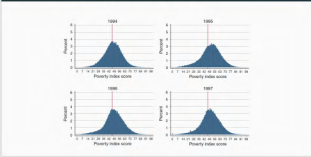
Manipulation of Social Program Eligibility*

By ANTONIO CARRASCO AND ESTER CRIVELLO**

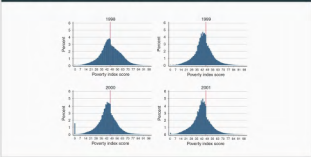
We document how manipulation of a targeting criterion for social welfare programs creates over time. First, there was unusual behavior of some local politicians in the timing of the household censuses around local elections. Then, there was a sharp increase in the sudden emergence of a sharp discontinuity in the same density around the eligibility threshold, which coincided with the change of the data reporter in that region. The discontinuity at the threshold is larger where regional elections are more competitive. These related facts are closely related to corruption, and results also highlight the importance of information and incentives. *LEZ*, 0712-12, 12A, 00A, 011



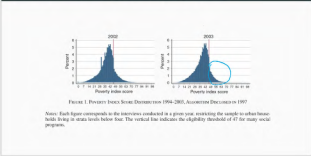
Visualizing manipulation — Proxy means test in Colombia



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Visualizing manipulation — Proxy means test in Colombia



Endogenous cutoffs: Evaluating smoothness through balance

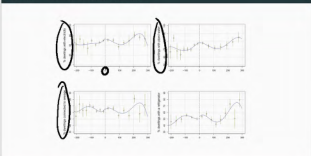
- Balance tests and placebo tests are related but distinct
- We can't directly test smoothness because we don't observe potential outcomes
- RD is like a "local RCT": Average values of exogenous covariates shouldn't jump around the cutoff
- Balance tests are indirect searching for evidence supporting smoothness

Balance implementation

- Don't make it hard — do what you did in Z only Y → CHARACTERISTICS & NO CHANGE
- Choose other nonconfounders associated with potential outcomes, Z
- Create similar graphical plots as you did for Y
- Could also conduct the parametric and nonparametric estimation on Z
- You do not want to see a jump around the cutoff, z

REG $\alpha + \beta X + \lambda(X-T) + \epsilon$
 DEBERIA SER CERO

Balance — FONDEN running example

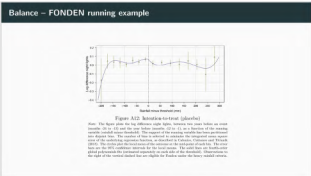


Balance — FONDEN running example



Placebos at non-discontinuous points

- Placebos in time are common with panel; placebo in running variables are their equivalent in RDD
- Imbens and Lemieux (2010) suggest we look at one side of the discontinuity (e.g., $X < c_0$), take the median value of the running variable in that section, and pretend it was a discontinuity, c_0^*
- Then test whether in reality there is a discontinuity at c_0^* . You do not want to find anything.
- Remember: smoothness at placebo points is neither necessary nor sufficient for smoothness in the potential outcomes at the cutoff



Fuzzy design

Fuzzy RDD, IV and ITT

- Fuzzy RDD is an IV estimator, and requires those assumptions
- You may be more comfortable with presenting the intent-to-treat (ITT) parameter which is just the reduced form regression of Y on Z , therefore
- Many papers will not present an IV-style parameter, but rather a lizzard of ITT parameters, out of a "fear" that the exclusion restrictions may not hold
- But let's review the IV approach anyway for completeness (more IV to come!)

Probability of treatment jumps at discontinuity

Probabilistic treatment assignment (i.e., "fuzzy RDD")
 The probability of receiving treatment changes discontinuously at the cutoff, c_0 , but need not go from 0 to 1

$$\lim_{x \rightarrow c_0^-} P(T = 1 | X = c_0) \neq \lim_{x \rightarrow c_0^+} P(T = 1 | X = c_0)$$

Examples: Incentives to participate in some program may change discontinuously at the cutoff but are not powerful enough to move everyone from non participation to participation.

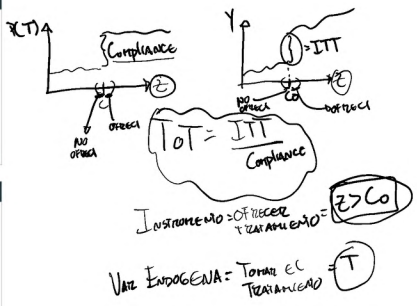
Deterministic (sharp) vs. probabilistic (fuzzy)

- In the sharp RDD, T is determined by $X > c_0$
- In the fuzzy RDD, the conditional probability of treatment jumps at c_0
- The relationship between the conditional probability of treatment and X can be written as:

$$P(T = 1 | X) = \beta_0(X) + \beta_1(X) \cdot \mathbb{1}(X > c_0)$$
 where $\beta_0 = 1$ if $X > c_0$ and 0 otherwise.

Instrumental variables

- As said, fuzzy designs are numerically equivalent and conceptually similar to IV (Instrument Z with X and $X > c_0$)
- "Reduced form": Numerator: "jump" in the regression of the outcome on the running variable X
- "First stage": Denominator: "jump" in the regression of the treatment indicator on the running variable X
- Same IV assumptions, covariates about compliers vs. defiers and statistical tests that we discussed with instrumental variables apply here



$$\frac{ITT \text{ (Salto en Y cuando } X > c_0)}{Compliance \text{ (Salto en T cuando } X > c_0)} = TOT$$

Wald estimator

Wald estimator of treatment effect under Fuzzy RDD
 Average causal effect of the treatment is the Wald IV parameter

$$\hat{\beta}_{\text{Wald RDD}} = \frac{E[Y|X = c_1] - E[Y|X = c_2]}{E[X|X = c_1] - E[X|X = c_2]}$$

ITT Compliance

Limitations of the LATE

- Fuzzy RDD has assumptions of all standard IV framework (exclusion, independence, nonzero first stage, and monotonicity)
- As with other binary IVs, the fuzzy RDD is estimating LATE: the local average treatment effect for the group of compliers
- In RDD, the compliers are those whose treatment status changed as we moved the value of x_i from just to the left of c_0 to just to the right of c_0

Balance - FONDEN running example

Then, we use local polynomial methods to estimate the first stage and the LATE. The typical causal parameter is the LATE.

$$F_{it} = \alpha_0 + \alpha_1 \text{Above} + \epsilon_{it}$$

$$Y_{it} = \gamma_0 + \gamma_1 F_{it} + \epsilon_{it}$$

where F_{it} is a binary variable that takes the value of one if the individual is eligible for funds. The variable for instrumenting the treatment is the difference in the probability of receiving the treatment between the compliers and the compliers who are not compliers. The variable F_{it} is an indicator variable for observed receipt exceeding the binary receipt threshold. Finally, ϵ_{it} and ϵ_{it} are error terms. The parameters of interest are the first stage estimate γ_1 , the LATE estimate α_1 , and the ratio γ_1/α_1 , which can be interpreted as the LATE under some additional assumptions.

Balance - FONDEN running example

	Below	Above
Panel A: Receipt (Y)	0.00	0.00
Fuzzy RDD	0.00	0.00
Panel B: Receipt (Y)	0.00	0.00
Fuzzy RDD	0.00	0.00
Panel C: LATE (Y)	0.00	0.00
Fuzzy RDD	0.00	0.00
Panel D: LATE (Y)	0.00	0.00
Fuzzy RDD	0.00	0.00
Panel E: LATE (Y)	0.00	0.00
Fuzzy RDD	0.00	0.00
Panel F: LATE (Y)	0.00	0.00
Fuzzy RDD	0.00	0.00

Visualization

Pictures, pictures and more pictures

- RDD is visually intense
- Eyeball tests are rampant (and disordered) in RDD studies
- Let's review some of the graphs you have to include

Outcomes

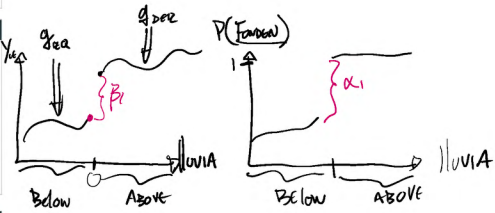
1. Outcome by running variable (X):

- Construct bins and average the outcome within bins on both sides of the cutoff
- Look at different bin sizes when constructing these graphs
- Plot the running variables, X_i , on the horizontal axis and the average of Y_i for each bin on the vertical axis
- Consider plotting a relatively flexible regression line on top of the bin means, but some readers prefer an eyeball test without the regression line to avoid "triming"

Probability of treatment

2. Probability of treatment by running variable if fuzzy RDD

- In a fuzzy RDD, you also want to see that the treatment variable jumps at c_0



$\beta_1 = \text{ITT}$

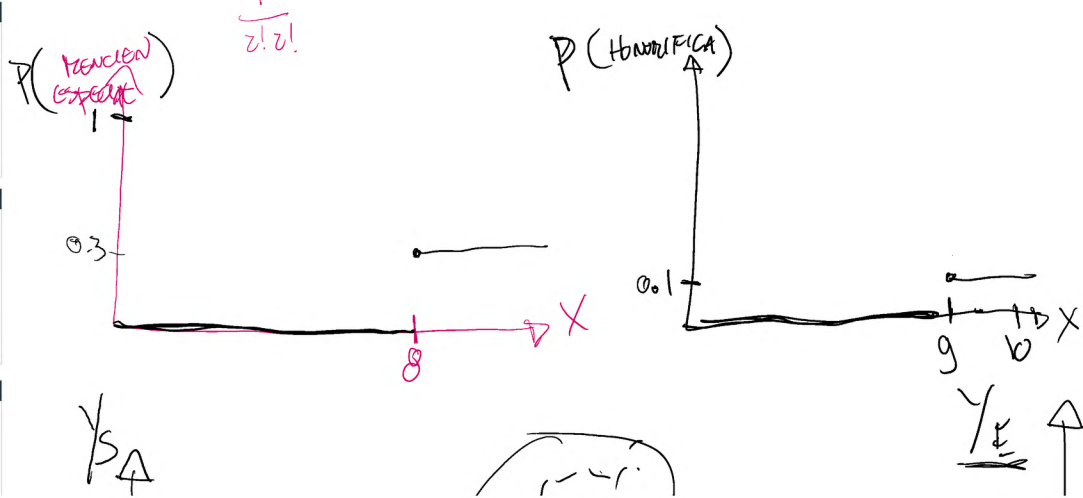
$\frac{\beta_1}{\alpha_1} = \text{LATE}$

$Y_{it} = \gamma_0 + \gamma_1 \text{Funds}_{it} + \epsilon_{it}$

$\text{Funds}_{it} = \alpha_0 + \alpha_1 \text{Above}_{it} + \nu_{it}$

$\nu_{it} \rightarrow \text{Instrument}$

$\frac{\gamma_1}{\alpha_1} = \text{LATE}$



Handwritten scribbles and notes.

2. Probability of treatment by running variable if fuzzy RDD

- In a fuzzy RDD, you also want to see that the treatment variable jumps at c_0 .
- This tells you whether you have a first stage ("fit")
- Let's look at that again from earlier Hoxby (2008) and enrollment at the flagship

McCrary Density

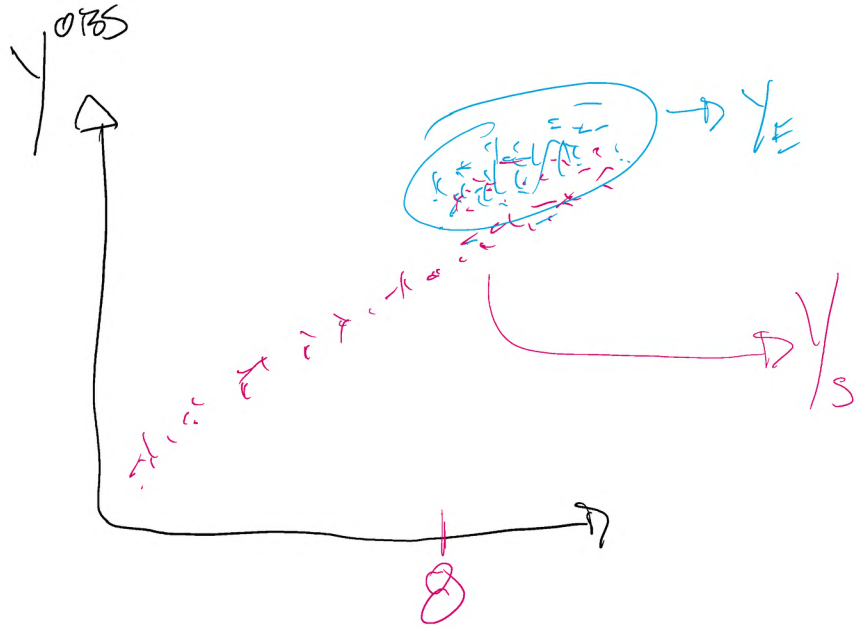
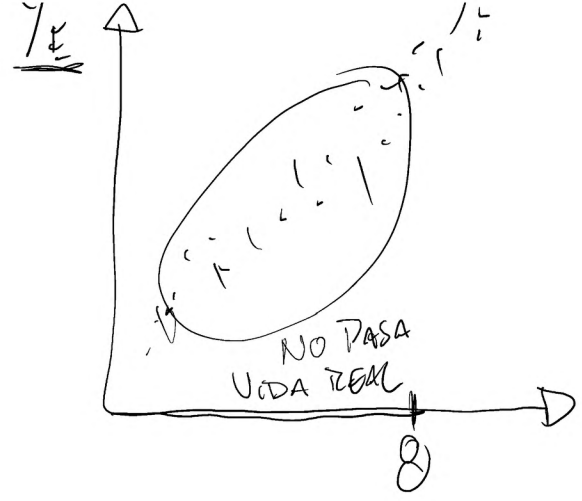
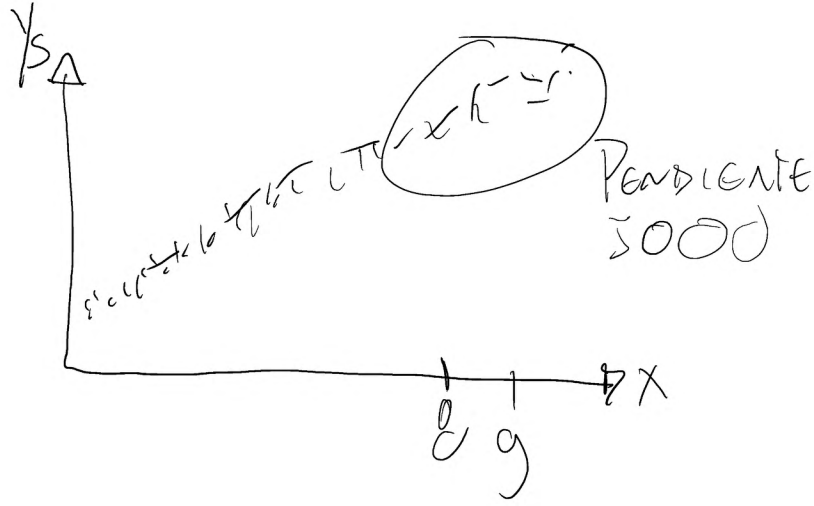
3. Density of the running variable

- One should plot the number of observations in each bin.
- This plot allows to investigate whether there is a discontinuity or heaping in the distribution of the running variable at the threshold.
- Heaping or discontinuities in the density suggest that people can manipulate their running variable score.
- This is an indirect test of the identifying assumption that each individual has imprecise control over the assignment variable, which may violate smoothness

Balance pictures

4. Covariates by a running variable

- Construct a similar graph to the outcomes graph but use a noncollider covariate as the "outcome"
- Balance implies smoothness through the cutoff, c_0 .
- If noncollider covariates jump at the cutoff, one is probably justified to reject that potential outcomes aren't also probably jumping there.



REG

$$Y \sim \alpha_0 + \alpha_1 \cdot X + \alpha_2 (\text{MENCION}) + \alpha_3 (\text{MENCION} \cdot X) + \epsilon.$$

Instrumento ($X > 8$ en la ventana $7.9 < X < 8.1$)

α_2^{IV} CAUSAL

Dummy