Matching and Re-weighting

When X differ...

Fernando Rios-Avila

Recap: Potential outcomes and Identification

To identify treatment effects one could **just** compare potential outcomes in two states:

- with treatment
- without treatment

Mathematically, average treatment effects would be:

$$ATE = E(Y_i(1) - Y_i(0))$$

the problem: with real data, we are only able to see one outcome. The counter factual is not observed:

$$Y_i=Y_i(1)\ast D+Y_i(0)\ast (1-D)$$

and simple differences may not capture ATE, because of selection bias and heterogeneity in effects.

Recap: Gold Standard - RCT

The easiest, but most expensive, way to deal with the problem is using **Randomized Control Trials**.

Effectively, you randomize Treatment, so that potential outcomes are independent of treatment:

$$Y(1), Y(0) \perp D$$

In other words, the distribution of potential outcomes is the same for those treated or untreated units.

$$\begin{split} E(Y,D=1) &= E(Y(1),D=1) = E(Y(1),D=0) \\ E(Y,D=0) &= E(Y(0),D=1) = E(Y(0),D=0) \\ ATT &= E(Y,D=1) - E(Y,D=0) \end{split}$$

But what if you can't Randomize

When unconditional fails

More often than not, specially if we didn't construct the data, it would be impossible to find that unconditional independence assumption holds.

For example, treatment (say having health insurance) may vary by age, gender, race, location, etc.

This is similar to the selection bias: Outcomes across treated and untreated groups will be different because:

- Composition: Characteristics of people among the treated could be different than those among the untreated For example, they could be older, more educated, mostly men, etc.
- Other factors: There could be factors we cannot control for, that also affect outcomes.

There is conditional

When unconditional independence assumption fails, we can call on Conditional independence assumption:

$Y(1), Y(0) \perp D|X$

In other words, If we can look into specific groups (given X), it may be possible to impose the Independence assumption.

This relaxes the independence condition, but assumes selection is due to observable characteristics only. (it still needs to be as good as randomized given X)

Implications:

$$\begin{split} E(Y|D=1,X) &= E(Y(1)|D=1,X) = E(Y(1)|D=0,X) \\ E(Y|D=0,X) &= E(Y(0)|D=1,X) = E(Y(0)|D=0,X) \end{split}$$

Intuition

Matching is a methodology that falls within quasi-experimental designs. You cannot or could not decide the assignment rules, so now are using data as given.

The idea is to construct an artificial control and use it as a counter-factual, so that both treated and control groups "look similar" in terms of observables.

Once a group of synthetic controls has been constructed, treatment effects can be calculated for the whole population:

$$\begin{split} ATE(X) &= E(Y|D=1,X) - E(Y|D=0,X) \\ ATE &= \int ATE(X) dFx \end{split}$$

How can we do this?

we just need to find observational twins!

Matching Twins



Figure 1: Matching on Observables

Subclassification or stratification

Consider the following dataset:

```
frause titanic, clear
expand freq
drop if freq==0
gen class1=class==1
tab survived class1 , nofreq col
<IPython.core.display.HTML object>
(Data downloaded from R base)
(8 zero counts ignored; observations not deleted)
(2,177 observations created)
(8 observations deleted)
          class1
 Survived | 0 1 | Total

        No
        72.92
        37.54
        67.70

        Yes
        27.08
        62.46
        32.30

Total | 100.00 100.00 | 100.00
```

If we assume full Independence assumption we would believe that being in first class increased chance of survival in 35.4%. but is that the case?

What if the composition of individuals differs across classes (women and children)

tab age class1, nofreq col tab sex class1, nofreq col

	class1		
Age	0	1	Total
Child Adult	5.49 94.51	1.85 98.15	4.95 95.05
Total	100.00	100.00	100.00
Sex	class1 0	1	Total

Male		82.68	55.38	I	78.65
Female	1	17.32	44.62	1	21.35
Total		100.00	100.00		100.00

There were fewer children, but more women in first class. Perhaps that explains the difference in survival rates

A better approach would be to look into the survival probabilities stratifying the data:

```
gen surv=survived==2
bysort age sex class1:egen sr_mean=mean(survived==2)
table (age sex) (class1), stat(mean surv) nototal
```

	cla O	ass1 1
Ι		
Ι	.4067797	1
Ι	.6136364	1
Ι		
	.1883378	.3257143
Ι	.6263345	.9722222
		cla 0 1 1 .4067797 .6136364 1 .1883378 .6263345

So even within each group, the survival probability is larger in first class. What about Average?

```
bysort age sex:egen sr_mean_class1=max(sr_mean*(class1==1))
bysort age sex:egen sr_mean_class0=max(sr_mean*(class1==0))
gen teff = sr_mean_class1-sr_mean_class0
sum teff if class1==1 // ATT
sum teff if class1==0 // ATU
sum teff // ATE
```

Max	Min	Std. dev.	Mean	Obs	Variable
.5932204	.1373765	.1125033	. 2375421	325	teff
Max	Min	Std. dev.	Mean	Obs	Variable
.5932204	.1373765	.1089261	.1887847	1,876	teff
Max	Min	Std. dev.	Mean	Obs	Variable
.5932204	.1373765	.1107948	. 1959842	2,201	teff

What did we do?

The procedure above is a simple stratification approach, aka matching, to analyze the true impact of the treatment (being a 1st class passenger).

- 1. Stratified the sample in groups by age and gender.
 - Identify the shares of each group by class1
- 2. Predict probability of survival per strata and class1
- 3. Obtain the Strata level Effects
- 4. Aggregate as needed.
 - Here, we could estimate ATE, ATT or ATU!

Where could things go wrong?

Overlapping

The procedure describe above works well whenever there is data overlapping.

+ For every combination of X, you see data on the control and treated group 0 < P(D|X) < 1

When this fails, you wont be able to estimate ATE's, although ATT's or ATU's might still be possible:

- for ATT: P(D|X) < 1
- for ATU: 0 < P(D|X)

For example:

frause hhprice, clear
keep price rooms type_h
tab rooms type_h

	Ι	=0 if house,	=1		
Number of		TownHouse			
rooms	Ι	0	1	I	Total
	-+-			-+-	
1		37	72		109
2		1,134	751		1,885
3		4,634	648	Ι	5,282
4	Ι	2,465	115	Ι	2,580
5	Ι	465	2	Ι	467
6	Ι	46	0	Ι	46
7	Ι	7	0	Ι	7
	-+-			-+-	
Total	Ι	8,788	1,588	I	10,376

Would not be able to estimate ATE nor ATU. Only ATT for townhouses.

Curse of dimensionality

There is a second problem in terms of stratification. How would we deal with Multiple dimensions? Would it be possible to find "twins" for every observation?

The answer is, probably no. Too many groups to track, to many micro cells to make use of:

```
frause oaxaca, clear
drop if lnwage==.
egen strata=group(educ isco)
bysort strata:egen flag=mean(female)
list educ isco female if (flag==0 | flag==1) & educ == 10, sep(0)
```

```
(Excerpt from the Swiss Labor Market Survey 1998)
(213 observations deleted)
```

```
+----+
| educ isco female |
```

	-			
158.	Ι	10	1	0
159.	Ι	10	1	0
197.	Ι	10	7	0
198.	Ι	10	7	0
199.	Ι	10	9	1
200.	Ι	10	9	1
	+-			+

Alternative: Matching as a weighted

The problem of curse of dimensional states that as the number of desired characteristics to match increase, fewer "twins" will be available in the data. At the end...no one will be like you!

The alternative, is to look into People that are sufficiently close so they can be used for matching.

$$\begin{split} ATT_i &= Y_i - \sum_{j \in C} w(x_j, x_i) Y_j \\ ATT &= \frac{1}{N_T} \sum (ATT_i) \\ ATT &= E(Y|D=1) - E_i \left(\sum_{j \in C} w(x_j, x_i) Y_j \Big| D = 0 \right) \end{split}$$

Depending how w(.) is defined, we would be facing different kinds of matching estimators.

Types of Matching

Matching on covariates

The first decision to take is whether one should find matches based on covariates, or based on scores (propensity scores).

Using covariates implies that will aim to find the closest "twin" possible, based on multiple dimensions:

$$\begin{split} Eclidean &= d(x_i, x_j) = \sqrt{(x_i - x_j)'(x_i - x_j)} \\ WEclidean &= d(x_i, x_j) = \sqrt{(x_i - x_j)'W(x_i - x_j)} \\ Maha &= d(x_i, x_j) = \sqrt{(x_i - x_j)'S^{-1}(x_i - x_j)} \end{split}$$

Distance measures are used to identify the closest matches to a given observation, and thus the weight assigned to that observation.

Has the advantage of looking at individuals who are indeed close to each other, but becomes more difficult as the dimensionality of X's increase. (you will not find close matches)

Matching on Scores

A second approach is to match individuals based on some summary index that condenses the information in X into a single scalar h(x), reducing the dimensionality problem from K to 1.

Few candidates:

- Propensity Score: P(D|X) based on a logit/probit/binomial model. Most common approach!
- Predicted Mean: $X\beta$ if there is information on outcome to be predicted
- PCA: Using Principal components to reduce dimensionality before Matching

Since there is only 1 dimension to consider, multiple distance measures are possible:

• nearest neighbors, kernel weight matching, radious matching.

But one has to be careful with the approach. King and Nielsen (2019) Argue about the risks of PSM

1 vs K matching; With and without replacement

Two additional questions remain regarding matching. How many "twins" to use, and if twins will be obtained with/without replacement.

- Fewer matches reduce bias (choosing only the closest observation), but increase variance.
- More matches increase bias, but reduce variance. (because of less optimal matches)
- with replacement: control units may be used more than once. This will improve matching quality reducing bias. But by using the same units multiple times, it will increase variance.
- without replacement: Control units are used once, potentially reducing matching quality, but reducing variance. It will be order dependent.

see Caliendo and Kopeing (2008)

What about SE? and Statistical inference?

Well....this is one of the few cases where Bootstrapping WON'T work!

Standard errors are more cumbersome. So we will just rely on software results

Other considerations

Once you have chosen your matching method, find your "statistical twins", and estimate your differences you are done! (or are you)

Not yet...common practice: Evaluate the balance of your data

Matching aims to reduce or eliminate differences in characteristics between treatment and control units. Thus, one should evaluate the differences (before and after match) of your characteristis

- 1. Check for overlapping condition.
- either variable by variable or with pscore
- 2. Assess Matching Quality: Have differences across groups vanished?
- Check Standardized differences $\frac{\mu_1 \mu_2}{\sqrt{0.5*(V_1 + V_2)}}$
- t-tests
- PR2 of regression with matched data

Implementation

In Stata, there are at least two approaches that can be used for matching:

- psmatch2 (from ssc)
- teffects (Official Stata command)

We will use this to answer a simple question:

What is the impact of Traing Jobs on Earnings?

Example

This file contains information on experimental and observed data for the analysis of training on earnings program:

```
use https://friosavila.github.io/playingwithstata/drdid/lalonde.dta, clear
keep if year==1978
drop if dwincl==0
label define sample 1 "exper" 2 "CPS" 3 "PSID"
label values sample sample
tab sample treated,m
```

```
(19,204 observations deleted)
(277 observations deleted)
```

	1	treated		
sample	0	1		Total
exper	260	185	0	445
CPS	0	0	15,992	15,992
PSID	0	0	2,490	2,490
Total	260	185	18,482	18,927

First Experimental design - RCT

```
reg re treated
tabstat age educ black married nodegree , by(treated)
logit treated age educ black hisp married nodegree
```

Source	SS SS	df	MS	Number of obs	; =	445
	+			F(1, 443)	=	8.04
Model	348013183	1	348013183	Prob > F	=	0.0048
Residual	1.9178e+10	443	43290369.3	R-squared	=	0.0178
+	+			Adj R-squared	l =	0.0156
Total	1.9526e+10	444	43976681.9	Root MSE	=	6579.5
re	Coefficient	Std. err.	t	P> t [95% c	onf.	interval]
	+					

treat _cc	ced ons	1794 4554	.342 .801	632. 408.	8534 0459	2.84 11.16	0.0 0.0	05 55 00 37	50.5745 752.855	3038.11 5356.747
Summary st Group vari	tatis	stics: Me	ean ed							
treated		age	ec	luc	blac	k maj	rried	nodegree	9	
0 1	 28 28	5.05385 5.81622	10.088 10.345	346 595	.826923	1 .153	38462 91892	.8346154	- L -	
Total	25	5.37079	10.19	551	.833707	9.168	35393 	.7820225	5	
Iteration Iteration Iteration Iteration	0: 1: 2: 3:	Log like Log like Log like Log like	elihood elihood elihood elihood	d = -d =	-302 294.729 294.714 294.714	. 1 08 64 64		Name	or of ob	- 44E
Log likeli	ihood	d = -294	.71464					LR o Prot Pseu	chi2(6) > > chi2 1do R2	= 14.77 = 0.0221 = 0.0244
treat	ced	Coeffi	cient	Std.	err.	z	P>	z [9	95% conf.	interval]
a ec bla hi marri nodegr cc	age duc ack isp ied ree ons	.0059 0639 2543 8299 .2342 8388 1.053	9171 9597 3689 1587 2415 5524 3028	.014 .07 .363 .504 .266 .309	2668 1354 9735 2305 1824 93833 7384	0.41 -0.90 -0.70 -1.64 0.88 -2.71 1.01	0.6 0.3 0.4 0.1 0.3 0.0 0.3	780 70 859 00 -1. 792 07 -1. 159	0220452 203811 0677438 817432 2874665 444933 0998064	.0338794 .0758916 .4590061 .159115 .7559495 2321722 3.105862

Then using PScore Matching CPS

```
keep if treated == 1 | sample ==2
replace treated=0 if treated==.
reg re treated
tabstat age educ black hisp married nodegree , by(treated)
```

```
(2,750 observations deleted)
```

(15,992 real changes made)

Source	SS SS	df	MS	Numb	per of obs	s =	16,177
	+			F(1,	16175)	=	142.43
Model	1.3206e+10	1	1.3206e+10	Prob) > F	=	0.0000
Residual	1.4997e+12	16,175	92717515.8	R-sc	luared	=	0.0087
	+			Adj	R-squared	i =	0.0087
Total	1.5129e+12	16,176	93528158.4	Root	MSE	=	9629
re	Coefficient	Std. err.	t	P> t	[95% c	conf.	interval]
treated _cons	-8497.516 14846.66	712.0207 76.14292	-11.93 194.98	0.000	-9893.1 14697.	156 .41	-7101.877 14995.91

Summary statistics: Mean Group variable: treated

nodegree	married	hisp	black	educ	age	treated
.2958354 .7081081	.7117309 .1891892	.072036 .0594595	.0735368 .8432432	12.02751 10.34595	33.22524 25.81622	0 1
. 3005502	.7057551	.0718922	.0823391	12.00828	33.14051	Total

We need to do trimming

bysort educ black hisp married:egen n11=sum(treated==1)
bysort age black hisp married:egen n22=sum(treated==1)
drop if n11==0 | n22 ==0

tabstat age educ black hisp married nodegree , by(treated) reg re treated

(13,536 observations deleted)

Summary statistics: Mean Group variable: treated

treated	age	educ	black	hisp	married	nodegree
0 1	24.24145 25.81622	11.69788 10.34595	. 252443 . 8432432	.0260586 .0594595	.3346906 .1891892	.2569218 .7081081
Total	24.35176	11.60318	.2938281	.0283983	.3244983	.2885271

Source	SS	df	MS	Number of obs	=	2,641
+	+			F(1, 2639)	=	73.89
Model	5.7607e+09	1	5.7607e+09	Prob > F	=	0.0000
Residual	2.0575e+11	2,639	77964783.1	R-squared	=	0.0272
+	+			Adj R-squared	=	0.0269
Total	2.1151e+11	2,640	80117339.3	Root MSE	=	8829.8

re	Coefficient	Std. err.	t	P> t	[95% conf.	interval]
treated	-5786.584	673.1834	-8.60	0.000	-7106.605	-4466.564
_cons	12135.73	178.1702	68.11	0.000	11786.36	12485.1

Lets do some matching

teffects nnmatch (re age educ black married nodegree) (treated)
tebalance summarize
teffects nnmatch (re age educ black married nodegree) (treated), nn(2)
tebalance summarize
teffects psmatch (re) (treated age educ black married nodegree)
tebalance summarize

teffects psmatch (re) (treated age educ black married nodegree) , nn(2)
tebalance summarize

Treatment-eff Estimator Outcome model Distance metr:	ects estimatio : nearest-ne : matching ic: Mahalanobi	Number of Matches	of obs = : requested = min = max =	2,641 1 1 138		
re	 Coefficient	AI robust std. err.	z	P> z	[95% conf.	interval]
ATE treated (1 vs 0)	 -3685.665	1188.666	-3.10	0.002	-6015.407	-1355.923

(refitting the model using the generate() option)

Covariate balance summary

		Raw	Matched
Numbor of obs		 2 6/1	۶ ۵۵۵
Treated obs	=	185	2,641
Control obs	=	2,456	2,641

		Standardized Raw	differences Matched	Var: Raw	iance ratio Matched
age		.2342346	015417	1.305844	.8410946
educ	I	7684118	0812288	1.881909	.8598207
black	Ι	1.473105	0	.7039609	1
married	I	3351313	0008087	.6923501	.999393
nodegree	Ι	1.010393	0	1.088086	1

Treatment-effects estimation			Number of obs		2,641
Estimator	:	nearest-neighbor matching	Matches: requested	l =	2
Outcome model	:	matching	mir	L =	2
Distance metric	:	Mahalanobis	max	: =	138

 re	Coef	ficient	AI r std.	obust err.	:	z	P> z	[95%	conf.	interval]
ATE treated (1 vs 0)	-51	66.888	1107	.653	-4.	 66	0.000	-7337		-2995.929
(refitting the	e mode	l using	the g	enerat	e() oj	ptio	n)			
Covariate bala	ince s	ummary								
		Ra	.w	Matc	hed					
Number of obs Treated obs Control obs	= = =	2,64 18 2,45	1 5 6 	5,2 2,0 2,0	 282 641 641					
	St: 	andardiz Ra	ed di w	fferen Match	ces ed		 Vari Raw	ance ra Mato	tio hed	
ag edu blac marrie nodegre	ge nc · k ed ·	.234234 768411 1.47310 335131 1.01039	6 – 8 – 5 3 3	.02090 .038523 .00746 00453 .00167	48 84 73 86 05	1 1 1	.305844 .881909 7039609 6923501 .088086	.7345 .8978 1.006 .9965 1.001	9997 301 716 432 557	
Treatment-effe Estimator Outcome model Treatment mode	ects e : pr : ma el: log	stimatio opensity tching git	n -scor	e matc]	hing		Number o Matches:	f obs reques	= sted = min = max =	2,641 1 1 138
 re	Coef	ficient	AI r std.	obust err.	:	 z	P> z	[95%	conf.	interval]
ATE treated (1 vs 0)	-42'	78.549	1135	.847	-3.	77	0.000	-6504	.768	-2052.331
(refitting the	mode	l using	the g	enerat	e() o	ptio	n)			

Covariate balance summary

	Raw	Matched
Number of obs =	2,641	5,282
Treated obs =	185	2,641
Control obs =	2,456	2,641

	Standardized	differences	Variar	nce ratio
	Raw	Matched	Raw	Matched
age	.2342346	.0014058	1.305844	.9313458
educ	7684118	1308249	1.881909	.9665937
black	1.473105	0926638	.7039609	.90999
married	3351313	0973289	.6923501	.9197524
nodegree	1.010393	.0821105	1.088086	1.07103

Treatment-effe	ects estimation	n		Number of	f obs	=	2,641
Estimator	: propensity	-score match	ing	Matches:	requested	=	2
Outcome model	: matching				min	=	2
Treatment mode	el: logit				max	=	138
		AI robust					
re	Coefficient	std. err.	Z	P> z	[95% co	nf.	interval]

ATE |

treated |

(1 vs 0) | -4380.078 1158.019 -3.78 0.000 -6649.754 -2110.403

(refitting the model using the generate() option)

Covariate balance summary

	Raw	Matched
Number of obs =	2,641	5,282
Treated obs =	185	2,641
Control obs =	2,456	2,641

Standardized differences Variance ratio Raw Matched Raw Matched age .2342346 06133 1.305844 .8834346 educ 7684118 1321518 1.881909 1.021302 black 1.473105 0698339 .7039609 .933348 married 3351313 0414439 .6923501 .9674741 nodegree 1.010393 .0939209 1.088086 1.080951						
age .234234606133 1.305844 .8834346 educ 76841181321518 1.881909 1.021302 black 1.4731050698339 .7039609 .933348 married 33513130414439 .6923501 .9674741 nodegree 1.010393 .0939209 1.088086 1.080951		S 	tandardized Raw	differences Matched	Vari Raw	ance ratio Matched
	age educ black married nodegree	 	.2342346 7684118 1.473105 3351313 1.010393	06133 1321518 0698339 0414439 .0939209	1.305844 1.881909 .7039609 .6923501 1.088086	.8834346 1.021302 .933348 .9674741 1.080951

A missing variable? Earnings in previous year. May capture information of Need to do treatment (selection)

```
tabstat age educ black hisp married nodegree re74, by(treated)
gen dre = re-re74
teffects nnmatch (dre age educ black married nodegree ) (treated)
teffects nnmatch (dre age educ black married nodegree ) (treated), nn(2)
teffects psmatch (dre) (treated age educ black married nodegree )
teffects psmatch (dre) (treated age educ black married nodegree ), nn(2)
```

Summary statistics: Mean Group variable: treated

reated	l age	educ	black	hisp	married	nodegree
0 1	24.24145 25.81622	11.69788 10.34595	.252443 .8432432	.0260586 .0594595	.3346906 .1891892	.2569218 .7081081
Total	24.35176	11.60318	.2938281	.0283983	.3244983	.2885271

treated | re74

0		9347.406
	। +−	2095.574
Total	I	8839.421

Treatment-effects estimation Number of obs = 2,641 Estimator : nearest-neighbor matching Matches: requested = 1 Outcome model : matching min = 1 Distance metric: Mahalanobis max = 138 _____ AI robust dre | Coefficient std. err. z P>|z| [95% conf. interval] ______ ATE treated | (1 vs 0) | 2616.653 1803.172 1.45 0.147 -917.4997 6150.806 _____ Treatment-effects estimation Number of obs = 2.641 Estimator : nearest-neighbor matching Matches: requested = 2 Outcome model : matching 2 min = Distance metric: Mahalanobis max = 138 _____ AI robust dre | Coefficient std. err. z P>|z| [95% conf. interval] _____+ ATE | treated (1 vs 0) | 730.2925 1674.91 0.44 0.663 -2552.47 4013.055 _____ Treatment-effects estimation Number of obs = 2,641 1 Estimator : propensity-score matching Matches: requested = Outcome model : matching min = 1 138 Treatment model: logit max = -----AI robust dre | Coefficient std. err. z P>|z| [95% conf. interval] ------_____ ATE treated | (1 vs 0) | 2162.311 1740.12 1.24 0.214 -1248.262 5572.884

_____ Number of obs = Treatment-effects estimation 2,641 Estimator : propensity-score matching Matches: requested = 2 2 Outcome model : matching min = Treatment model: logit max = 138 _____ AI robust dre | Coefficient std. err. z P>|z| [95% conf. interval] ATE | treated | (1 vs 0) | 1833.03 1739.496 1.05 0.292 -1576.318 5242.379 _____

In this case, Matching alone could not get the right answer. Who were the most likely to "go to the training?"

So instead we change the question: How much the change in earnings compare across groups.

Wait: What about Reweighting?

An alternative method to Matching is to do Re-weighting.

We have seen this!

Your control group has a distribution g(x) and your treatment f(x). We can use some weighting factors h(x) that reshapes $g(x) \to \hat{f}(x)$.

How? Using Propensity scores

Why does it work? Just as matching, your goal is to compare distributions of outcomes, forcing differences in observed characteristics to be the same.

IPW, does this by reweighting the distribution! (rather than matching)

Inverse Probability Weighting: IPW

s1: Estimate Pscore

$$p(D=1|X) = F(X\beta)$$

S2: Estimate IPW

For ATT: W(D = 1, x) = 1 & $W(D = 0, X) = \frac{\hat{p}(x)}{1 - \hat{p}(x)}$ For ATU: W(D = 0, x) = 1 & $W(D = 1, X) = \frac{1 - \hat{p}(x)}{\hat{p}(x)}$ For ATE: $W(D = 0, x) = \frac{1}{1 - \hat{p}(x)}$ & $W(D = 1, X) = \frac{1}{\hat{p}(x)}$ s3: Estimate Treatment effect:

$$TE = \sum_{i\in D=1} w_i^s(1)Y_i - \sum_{i\in D=0} w_i^s(0)Y_i$$

Even Better: Go DR!

An interesting advantage of IPW approach is that you can gain efficiency using Doubly Robust Methods. Namely, instead of comparing outcomes directly, you could compare predicted outcomes!

$$\begin{split} ATT &= \frac{1}{N_t} \sum (Y_1 - X' \hat{\beta}^0_w) \\ ATU &= \frac{1}{N_c} \sum (X' \hat{\beta}^1_w - Y_0) \\ ATE &= \frac{1}{N} \sum (X' \hat{\beta}^1_w - X' \hat{\beta}^0_w) \end{split}$$

where $\hat{\beta}_w^k$ can be modeled using weighted least squares

Comparing to Matching

teffects ipw (re) (treated age educ black married nodegree), iter(3) nolog teffects ipwra (re age educ black married nodegree) (treated age educ black married nodegre teffects ipw (dre) (treated age educ black married nodegree), iter(3) nolog teffects ipwra (dre age educ black married nodegree) (treated age educ black married nodegree)

Treatment-effects estimation				Number	of	obs	=	2,641
Estimator	: inverse-pr							
Outcome model	: weighted m	ean						
Treatment mode	l: logit							
		Robust						
re	Coefficient	std. err.	Z	P> z		[95%	conf.	interval]

_____+ ATE treated | (1 vs 0) | -4833.352 1088.667 -4.44 0.000 -6967.101 -2699.603 _____ POmean treated | 0 | 11979.19 179.1903 66.85 0.000 11627.99 12330.4 Treatment-effects estimation Number of obs = 2,641 Estimator : IPW regression adjustment Outcome model : linear Treatment model: logit _____ Robust re | Coefficient std. err. z P>|z| [95% conf. interval] _____ ATE treated | (1 vs 0) | -4835.38 1012.598 -4.78 0.000 -6820.036 -2850.724 _____ POmean treated | 0 | 11976.52 179.0958 66.87 0.000 11625.49 12327.54 _____ Warning: Convergence not achieved. Number of obs = 2,641Treatment-effects estimation Estimator : inverse-probability weights Outcome model : weighted mean Treatment model: logit _____ Robust dre | Coefficient std. err. z P>|z| [95% conf. interval] _____ ____+ _____ ATE treated (1 vs 0) | 1475.71 1792.427 0.82 0.410 -2037.382 4988.802 _____ POmean treated |

0		2746.475	161.2845	17.03	0.000	2430.363	3062.587
Warning: Cor	ive	rgence not ac	hieved.				
Treatment-ef Estimator Outcome mode Treatment mo	ffe el ode	cts estimatio : IPW regres : linear l: logit	n sion adjustr	nent	Number	of obs =	2,641
dre	 e	Coefficient	Robust std. err.	z	P> z	[95% conf.	interval]
ATE treated (1 vs 0)	 1 1	1286.605	1516.493	0.85	0.396	-1685.666	4258.875
POmean treated 0	+ 1 1 	2754.756	161.4406	17.06	0.000	2438.338	3071.173
Warning: Cor	ive	rgence not ac	hieved.				

Next: Regression Discontinuity