# Causal inference weeks 2 & 3: Selection on observables

Regression, matching, and sub-classification

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DIGM is also known as the naive estimator. Let's not be so naive!

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Ultimately use regression, but understand others.

## Introduction to covariate adjustment

Covariate adjustment based on categorical variables

Covariate adjustment using the propensity score

Covariate adjustment in sparse data without the propensity score

Two important facts about regression

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**To discuss:** Think of an example where  $D_i$  and  $X_i$  might be related in this way. What does Y represent in your example?

# Outcomes by X and D

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So, what is the effect of  $D_i$  on  $Y_i$ ?

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- treatment group: ATT (ATE on the treated)
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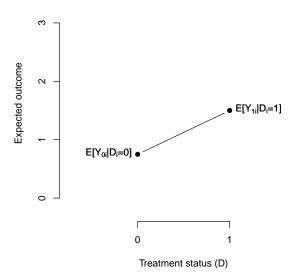
control group: ATC (ATE on the control)

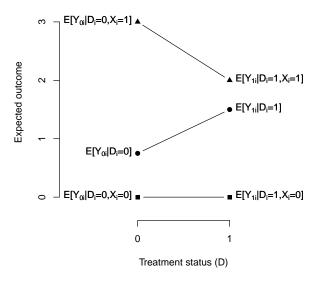
So what is ATE, ATT, ATC in this example (given CIA)?

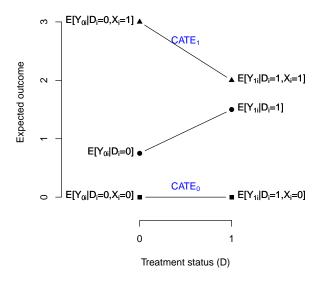


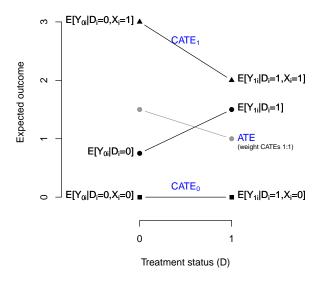


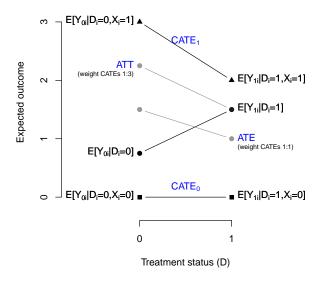
Treatment status (D)



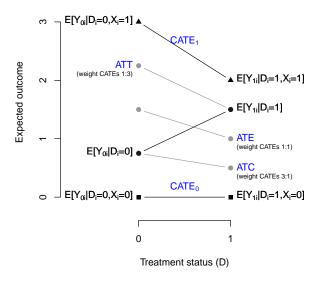




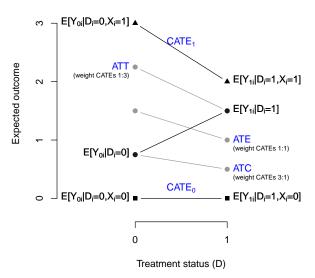




## Illustration for this example



# Illustration for this example (2)



#### Check understanding:

- Why are ATE, ATT, ATC all the same in a randomized experiment?
- How can the DIGM be positive when neither CATE is positive?

## Mathier explanation

The ATT is the weighted average of the CATEs, where the weights reflect the distribution of  $X_i$  in the treatment group.

ATT = 
$$\sum_{x=0,1} E[Y_i(1) - Y_i(0)|X_i = x] Pr(X_i = x|D_i = 1)$$
 (1)

$$= \sum_{x=0,1} CATE_x Pr(X_i = x | D_i = 1)$$
 (2)

In this case, this is

$$ATT = 0 \times 1/4 + -1 \times 3/4 = -3/4.$$

The weights of 1/4 and 3/4 come from the joint distribution: the probability of  $X_i = 1$  given that  $D_i = 1$  can be calculated as the ratio  $\frac{3/8}{3/8+1/8}$ , which is a simple application of Bayes' Theorem.

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- ► There might be another variable Z<sub>i</sub> related to D<sub>i</sub> that affects Y<sub>i</sub> (selection bias persists).
- X<sub>i</sub> might be an outcome, so controlling for it introduces post-treatment bias. e.g. suppose D<sub>i</sub> had been randomly assigned – how would we interpret this data?

### What do we condition on/control for?

For the CIA to hold, we must control for every variable that

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**To discuss**: In regressions, generally only one coefficient (at most) can be interpreted as a (causal) effect. Why is that?

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Link to CIA more obvious for sub-classification and matching, but regression more flexible and common.

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What about the DIGM as an estimator for ATE?

## Covariates

#### For each candidate, we have

- party
- electoral results
- year of birth
- secondary education
- university education
- profession

How can we use these to make better comparisons?

#### RETHNAL GREEN

Electorate: 42,172

*Holman, P. (Co-op. & Lab.)	)	20,519
Harris, Sir P. (L.)		9,715
Welfare, Mrs. D. (C.)		1,582
Mildwater, G. (Comm.)		610

Co-op. & Lab. majority 10,804

Mr. P. HOLMAN was elected for S.W. Bethnal Green in 1945. Born in 1891 and educated at Mill Hill and the London School of Economics, he has been a member of Middlesex County Council and Teddington U.D.C. He was sometime lecturer for the Workers' Educationa Association. He was a member of the Parlia-



member of the Parliamentary Labour Party groups on finance and industry.

Six Percy Haris, who is 73, first elected to Parliament for Harborough in 1916-18, represented S.W. Bethnal Green from 1922 to 1945. For 35 years he has served on the L.C.C. and after the 1949 elections was the only Libera on the council. Educated at Harrow and at Trinity Hall, Cambridge, he was called to the Bar in 1899. He was Chief Whip of the Parliamentary Liberal Party from 1935 to 1945, and deputy-leader in the war-time Parliament, and also chairman of the House of Commons All-Party Panel, and treasurer of the Inter-Parliamentary Union.

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#### Suppose we believe that CIA holds given candidate's

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#### Plausible?

## Number of candidates by party & education

	Par	ty: Con.		Par	ty: Lab.	
University:	Oxbridge	Other	None	Oxbridge	Other	None
School: Eton	16	4	5	2	0	0
Other public	40	25	31	21	10	3
Other	10	25	31	19	49	31
Not mentioned	7	12	17	5	27	37

## Number of MPs and unsuccessful candidates by cell

Note: (2,1) indicates 2 elected candidates and 1 unelected candidate

	Party: Con.			Party: Lab.		
University:	Oxbridge	Other	None	Oxbridge	Other	None
School: Eton	(14, 2)	(3, 1)	(4, 1)	(0, 2)	(0, 0)	(0, 0)
Other public	(18, 22)	(11, 14)	(11, 20)	(6, 15)	(4, 6)	(1, 2)
Other	(2, 8)	(8, 17)	(10, 21)	(2, 17)	(15, 34)	(13, 18)
Not mentioned	(4, 3)	(7, 5)	(12, 5)	(1, 4)	(8, 19)	(11, 26)

# Difference in group means (of log wealth at death) by cell

	Party: Con.			Par	ty: Lab.	
University:	Oxbridge	Other	None	Oxbridge	Other	None
School: Eton	2.61	2.66	-0.67	-	-	-
Other public	0.35	0.15	0.48	0.65	-0.27	0.58
Other	1.05	0.33	0.51	-0.02	-0.01	0.45
Not mentioned	0.06	1.48	0.25	-0.6	0.27	0.06

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Not mentioned	0.06	1.48	0.25	-0.6	0.27	0.06

### Average DIGM across cells

weighting by #candidates in each cell:	.40	(ATE
weighting by #MPs in each cell:	.54	(ATT)
weighting by #non-MPs in each cell:	.31	(ATC

## Matching

**Matching**: fill in missing potential outcomes using "nearest neighbor" with opposite treatment status.

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Party	School	University	Treated	In(Wealth)	$Y_{0i}$	$Y_{1i}$
Labour	Other Public	Not Mentioned	1	12.7	?	12.7
Labour	Other Public	Not Mentioned	0	11.8	11.8	
Labour	Other Public	Not Mentioned	0	12.5	12.5	

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To get ATE: take difference in mean of  $Y_{1i}$  and  $Y_{0i}$  (with imputations).

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To get ATE: take difference in mean of  $Y_{1i}$  and  $Y_{0i}$  (with imputations). To get ATT: same, but only using treated rows.

**Matching**: fill in missing potential outcomes using "nearest neighbor" with opposite treatment status. (e.g. for Tory **MP** born in 1928 who went to Oxford, use Tory **non-MP** born in 1927 who went to Cambridge)

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To get ATE: take difference in mean of  $Y_{1i}$  and  $Y_{0i}$  (with imputations).

To get ATT: same, but only using treated rows.

To get ATC: same, but only using control rows.

### Exact matching: implementation

#### Using Matching library:

```
> match.ate = Match(Y = d$Inrealgross, Tr = d$treated, X = d[,c("tory"
, "uni.cat", "sch.cat")], exact = T, estimand = "ATE")
> round(match.ate$est, 2)
     [,1]
[1,] 0.4
```

## Exact matching: implementation

```
Using Matching library:
> match.ate = Match(Y = d$lnrealgross, Tr = d$treated, X = d[,c("tory"
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     Γ,17
[1,] 0.4
> round(Match(Y = d$Inrealgross, Tr = d$treated, X = d[,c("tory", "uni
 .cat", "sch.cat")\, exact = T, estimand = "ATT")\est, 2)
     [,1]
Γ1, 7 0.54
> round(Match(Y = d$lnrealgross, Tr = d$treated, X = d[,c("tory", "uni
 .cat", "sch.cat")\uparrow, exact = T, estimand = "ATC")\$est, 2)
     Γ.17
[1,] 0.31
```

### Regression

Given CIA, **regression** implies regressing  $Y_i$  on an indicator for treatment and a dummy for every cell.

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```
> summary(lm(lnrealgross ~ treated + cell.cat, data = d))
Call:
lm(formula = lnrealgross ~ treated + cell.cat, data = d)
Residuals:
   Min
           10 Median
                          30
                                Max
-4.8977 -0.4868 -0.0022 0.4877 3.7358
Coefficients:
           Estimate Std. Error t value Pr(>|t|)
(Intercept) 13.0120
                      0.7079 18.382 < 2e-16 ***
treated
        0.3710
                      0.1057 3.510 0.000499 ***
cell.cat2 -0.7396
                      0.7274 -1.017 0.309864
cell.cat3 -0.9216
                      0.7343 -1.255 0.210151
cell.cat4 -0.1767
                      0.8378 -0.211 0.833110
cell.cat5 -0.8024 0.9145 -0.877 0.380777
cell ca+6 _0 5110
                      0 7766 _0 658 0 510017
```

### Regression (2)

Equivalent: regressing  $Y_i$  on each of the categorical variables and their interactions:

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```
> summary(lm(lnrealgross ~ treated + party*scat*ucat, data = d))
Call:
lm(formula = lnrealaross ~ treated + party * scat * ucat. data = d)
Residuals:
   Min
            10 Median
                            30
                                  Max
-4.8977 -0.4868 -0.0022 0.4877 3.7358
Coefficients: (2 not defined because of singularities)
                                         Estimate Std. Error t value Pr(>|t|)
(Intercept)
                                         13.04495
                                                     0.99091 13.165 < 2e-16 ***
treated
                                          0.37101
                                                    0.10570 3.510 0.000499 ***
partytory
                                          0.66122
                                                     0.88687 0.746 0.456362
scatnotMentioned
                                         -0.77257
                                                     1.00459 -0.769 0.442319
scatotherPublic
                                         -0.83536
                                                     1.14731 -0.728 0.466969
scatsecondary
                                         -1.10119
                                                     0.97491 -1.130 0.259342
ucatotherDearee
                                         -0.53300
                                                     0.75924 -0.702 0.483069
ucatoxbridae
                                         -0.03294
                                                     0.69342 -0.047 0.962139
partytory:scatnotMentioned
                                         -0.24588
                                                     0.93177 -0.264 0.791999
partytory:scatotherPublic
                                         -0.40490
                                                     1.07359 -0.377 0.706263
nartytory:scatsecondary
                                         -0.15465
                                                     0.85057
                                                              -0.182 0.855815
```

This is a **saturated model**, i.e. one with a dummy for each possible combination of the explanatory variables

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- all "main effects" (Labour, Eton, ...), and
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Given categorical variables, a saturated model is the most flexible possible functional form.

**To discuss:** Why would you prefer a saturated model to a model with only main effects, no interactions?

# Comparison of estimates

	ATE	ATT	ATC
Sub-classification	.40	.54	.31
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Regression	.37	-	-

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#### To note:

► ATE from sub-classification is average of cell DIGMs weighted by {#units in cell} (definition).

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#### To note:

- ATE from sub-classification is average of cell DIGMs weighted by {#units in cell} (definition).
- ATE from (exact) matching is exactly the same thing.
- ► ATE from saturated regression is the average of cell DIGMs weighted by {#units in cell × variance of treatment in cell} (Angrist and Pischke MHE p. 75).

Covariate adjustment using the propensity score

Introduction to covariate adjustment

Covariate adjustment based on categorical variables

Covariate adjustment using the propensity score

Covariate adjustment in sparse data without the propensity score

Two important facts about regression

Perhaps we don't quite believe the CIA based only on party, secondary education, and university education.

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But here we run into a problem of sparse data (curse of dimensionality):

- Sub-classification: many empty cells
- Matching: few exact matches
- (Saturated) regression: many empty groups, NA coefficients

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- Sub-classification: many empty cells
- Matching: few exact matches
- (Saturated) regression: many empty groups, NA coefficients

You may get an estimate, but it will be based on an unrepresentative subset (far from true ATE).

## How to proceed when many cells are empty

#### What can we do?

- Sub-classification: propensity score methods
- Matching: propensity score methods, nearest neighbor, coarsened exact matching
- ► Regression: propensity score methods, stronger CIA (i.e. less flexible functional form, e.g. drop interactions)

The propensity score is the probability of treatment, given covariates:

$$p(X_i) \equiv \Pr(D_i = 1|X_i) = E[D_i|X_i]$$

This can be estimated with OLS (linear probability model) or logistic regression (logit).

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The CIA becomes:  $Y_{0i}$ ,  $Y_{1i} \perp \!\!\!\perp D_i \mid p(X_i)$ 

Having estimated the propensity score, we can

- Sub-classification: calculate DIGM within bands of the propensity score
- Matching: match units based on nearby propensity scores
- Regression: regress outcome on treatment controlling flexibly for the propensity score

## Propensity score example

I regress the treatment indicator on party, secondary school category, university category, year of birth (yob), yob<sup>2</sup>, yob<sup>3</sup>, gender, 11 profession indicators.

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I regress the treatment indicator on party, secondary school category, university category, year of birth (yob), yob<sup>2</sup>, yob<sup>3</sup>, gender, 11 profession indicators.

```
ps.model = lm(treated ~ labour + scat + ucat + xxyob + I(xxyob^2) + I(xxyob^3) + xxfemale + xxoc_teacherall + xxoc_barrister + xxoc_solicitor + xxoc_dr + xxoc_civil_serv + xxoc_local_politics + xxoc_business + xxoc_white_collar + xxoc_union_org + xxoc_journalist + xxoc_miner, data = d, na.action = na.exclude) # na .exclude so that an NA is included in predictions for units with missing values
```

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```

#### Take a look at the results:

```
> summarv(ps.model)$coefficients
                                               t value
                         Estimate
                                    Std. Error
                                                              Pr(>|t|)
(Intercept)
                    2.640142e+04 9.015006e+04 0.2928608 0.7697793131
                   -1.535037e-01 5.526546e-02 -2.7775703 0.0057325165
labour
scatnotMentioned
                   -3.021275e-01 1.132277e-01 -2.6683185 0.0079308561
scatotherPublic
                    -3.154628e-01 1.036949e-01 -3.0422208 0.0025020298
scatsecondary
                    -3.864579e-01 1.072719e-01 -3.6026008 0.0003544281
ucatotherDearee
                    3.350661e-02 5.849440e-02 0.5728174 0.5670878012
ucatoxbridge
                    -4.119867e-02 6.544398e-02 -0.6295257 0.5293616454
xxvob
                    -4.187426e+01 1.410154e+02 -0.2969482 0.7666590682
I(xxvob^2)
                    2.213470e-02 7.352562e-02 0.3010475 0.7635335588
I(xxyob^3)
                    -3.899376e-06 1.277858e-05 -0.3051494 0.7604098834
xxfemale
                    -9.486131e-02 1.173128e-01 -0.8086184 0.4192117588
xxoc teacherall
                    -6.575377e-02 8.060895e-02 -0.8157131 0.4151461145
xxoc_barrister
                    -7.703770e-02 8.274819e-02 -0.9309896 0.3524163260
xxoc solicitor
                    -6.257366e-02 9.595262e-02 -0.6521308 0.5146885466
                    -3.314176e-02.1.551896e-01.-0.2135566.0.8310008086
xxoc_dr
```

## Propensity score example (2)

The propensity score is the prediction from this model:

pX = predict(ps.model)

### Propensity score example (2)

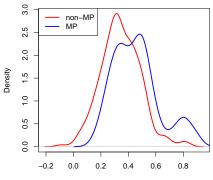
#### The propensity score is the prediction from this model:

```
pX = predict(ps.model)
```

#### Compare the distribution by treatment status:

```
plot(density(pX[d$treated == 0], na.rm = T), lwd = 2, col = "red", main = "Propensity score by treatment
status", xlab = "Probability of being an MP")
lines(density(pX[d$treated == 1], na.rm = T), lwd = 2, col = "blue")
legend("topleft", lwd = c(2,2), col = c("red", "blue"), legend = c("non-MP", "MP"))
```

#### Propensity score by treatment status



# Sub-classification on the propensity score

Let's start with 10 sub-classes of the propensity score:

```
library(dplyr)
d$pX.tile = ntile(pX, 10)
```

Counts and DIGMs in each sub-class:

Subclass	#units	#MPs	#non-MPs	DIGM
1	43	4	39	-0.24
2	43	13	30	0.01
3	43	13	30	0.9
4	42	12	30	0.33
5	43	18	25	0.24
6	43	15	28	-0.13
7	42	15	27	0.35
8	43	20	23	0.5
9	43	26	17	0.47
10	40	29	11	1.69

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7	42	15	27	0.35
8	43	20	23	0.5
9	43	26	17	0.47
10	40	29	11	1.69

ATE: 0.40 ATT: 0.57 ATC: 0.30

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7	42	15	27	0.35
8	43	20	23	0.5
9	43	26	17	0.47
10	40	29	11	1.69

ATE: 0.40 ATT: 0.57 ATC: 0.30

How many sub-classes? Bias-variance tradeoff.

# Nearest-neighbor matching on the propensity score

### Using defaults in Matching:

# Nearest-neighbor matching on the propensity score

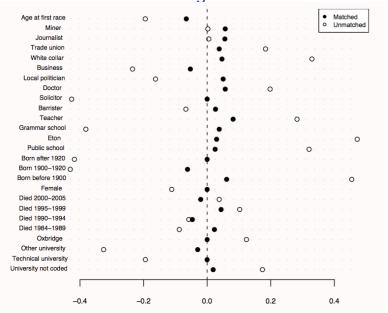
### Using defaults in Matching:

ATE: 0.42

ATT: 0.50

ATC: 0.38

## Balance statistics for matching



## Regression controlling for the propensity score

### Controlling via dummies for 10 sub-classes:

```
 d pX.tile = ntile(pX, 10) \\ summary(lm(lnrealgross \sim treated + as.factor(pX.tile), data = d)) \\ scoefficients \\
```

# Regression controlling for the propensity score

### Controlling via dummies for 10 sub-classes:

```
d$pX.tile = ntile(pX, 10)
summary(lm(lnrealgross ~ treated + as.factor(pX.tile), data = d))$coefficients
                         Estimate Std. Error
                                                 t value
                                                               Pr(>|t|)
(Intercept)
                     12.310737706
                                   0.1591787 77.33909546 3.258338e-248
treated
                      0.430298252
                                   0.1103101
                                              3.90080605
                                                          1.118504e-04
                      0.138439205
                                   0.2258278
                                              0.61302995
                                                          5.401932e-01
as.factor(pX.tile)2
as.factor(pX.tile)3
                      0.111779595
                                   0.2258278
                                              0.49497712
                                                          6.208789e-01
as.factor(pX.tile)4
                      0.065073825
                                   0.2269751
                                              0.28670023
                                                          7.744853e-01
as.factor(pX.tile)5
                      0.127734431
                                   0.2274973
                                              0.56147673
                                                          5.747764e-01
as.factor(pX.tile)6
                      0.158400433
                                   0.2264099
                                              0.69961800
                                                          4.845588e-01
as.factor(pX.tile)7
                      0.020971809
                                   0.2278481
                                              0.09204294
                                                          9.267084e-01
as.factor(pX.tile)8
                     -0.003937183
                                   0.2283635 -0.01724086
                                                          9.862528e-01
as.factor(pX.tile)9
                      0.227911766
                                   0.2316254
                                              0.98396686
                                                          3.257065e-01
as.factor(pX.tile)10
                      0.597870989
                                   0.2392019
                                              2.49944111
                                                          1.282519e-02
```

# Regression controlling for the propensity score (2)

### Controlling via polynomials of propensity score:

# Comparison of techniques (propensity score version)

### **Comparison of estimates**

AIE	AH	AIC
.40	.57	.30
.42	.50	.38
.41	-	-
.43	-	-
	.40 .42 .41	.42 .50 .41 -

# Comparison of techniques (propensity score version)

### Comparison of estimates

	ATE	ATT	ATC
Sub-classification	.40	.57	.30
Matching	.42	.50	.38
Regression: bins	.41	-	-
Regression: polynomials	.43	-	-

### To note:

 Propensity score is estimated, which should be considered in variance

# Comparison of techniques (propensity score version)

### Comparison of estimates

AIE	AH	AIC
.40	.57	.30
.42	.50	.38
.41	-	-
.43	-	-
	.40 .42 .41	.42 .50 .41 -

### To note:

- Propensity score is estimated, which should be considered in variance
- Our model for the propensity score has no interactions stronger assumptions than previous exercise

Covariate adjustment in spa	se data without the propensity score
-----------------------------	--------------------------------------

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Two important facts about regression

# Sub-classification without the propensity score

To avoid empty cells, you might try playing around with how cells are defined.

# Sub-classification without the propensity score

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This is basically how I think about **coarsened exact matching (CEM)** by King et al.

We can do nearest-neighbor matching with the covariates themselves, rather than propensity score.

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But which units are "near" each other?

We can do nearest-neighbor matching with the covariates themselves, rather than propensity score.

But which units are "near" each other?

- e.g. Should I match a Tory born in 1928 who went to Oxford to a
  - Tory born in 1927 with no university listed?
  - Labour candidate born in 1928 who went to the LSE?
  - ► Tory born in 1942 who went to Oxford?

We can do nearest-neighbor matching with the covariates themselves, rather than propensity score.

But which units are "near" each other?

- e.g. Should I match a Tory born in 1928 who went to Oxford to a
  - Tory born in 1927 with no university listed?
  - Labour candidate born in 1928 who went to the LSE?
  - Tory born in 1942 who went to Oxford?

#### Some of the options:

- scale distance on each variable by inverse of the variable's sample variance (default in Matching when not exact)
- scale distance by the inverse of the covariance matrix (Mahalanobis distance)
- genetic matching: search for a weight matrix that yields overall covariate balance (Diamond and Sekhon)

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1. Estimate propensity score  $p(X_i)$  based on model:

$$D_i = \alpha_0 + \alpha_1 X_{i1} + \alpha_2 X_{i2} + \ldots + \alpha_K X_{iK}$$

Regress  $Y_i$  on  $D_i$  and a flexible function of the propensity score.

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Regress  $Y_i$  on  $D_i$  and a flexible function of the propensity score.

2. Regress  $Y_i$  on  $D_i$  and covariates  $X_{i1}$  to  $X_{iK}$ .

## Regression: propensity score vs. covariates

### In the "MPs for Sale" example:

Approach	ATE
Regress $Y_i$ on $D_i$ and 10 bins of propensity score	.43
Regress $Y_i$ on $D_i$ and 4 polynomials of propensity score	.41
Regress $Y_i$ on $D_i$ and covariates from propensity score model	.41

#### My view:

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  - Statistical inference straightforward

## Covariate adjustment: bottom line

To make the "selection on observables" assumption credible, you need **good observables**, i.e. good measures of characteristics that affect the outcome and differ between treatment and control groups.

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Keep the statistics simple, focus on the data, and be opportunistic.

Two important facts about regression

Introduction to covariate adjustment

Covariate adjustment based on categorical variables

Covariate adjustment using the propensity score

Covariate adjustment in sparse data without the propensity score

Two important facts about regression

# Omitted variable bias formula Suppose two covariates, $X_{1i}$ and $X_{2i}$ .

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Then the omitted variables bias formula tells us

$$\beta^{s} = \beta^{l} + \pi_{21}\gamma.$$

Explain the last line in plain English.

#### OVB formula example: long regression

```
> long = lm(lnrealgross ~ treated + labour + scat + ucat, data = d)
> summary(long)
```

#### Call:

lm(formula = lnrealgross ~ treated + labour + scat + ucat, data = d)

#### Residuals:

Min 1Q Median 3Q Max -4.9695 -0.4504 -0.0260 0.3951 3.7635

#### Coefficients:

Signif. codes: 0 '\*\*\*' 0.001 '\*\*' 0.01 '\*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 0.9991 on 419 degrees of freedom Multiple R-squared: 0.1409, Adjusted R-squared: 0.1265 F-statistic: 9.816 on 7 and 419 DF, p-value: 2.386e-11

# OVB formula example: short regression

```
> short = lm(lnrealgross ~ treated + scat + ucat, data = d) # cut out labour
> summary(short)
Call:
lm(formula = lnrealgross ~ treated + scat + ucat, data = d)
Residuals:
   Min
            10 Median
                           30
                                 Max
-4.9081 -0.4111 -0.0555 0.4335 3.6594
Coefficients:
                 Estimate Std. Error t value Pr(>|t|)
(Intercept) 13.176890 0.233147 56.517 < 2e-16 ***
treated
            0.410704 0.103199 3.980 8.13e-05 ***
scatnotMentioned -0.817735   0.231155   -3.538   0.000449 ***
scatotherPublic -0.701988 0.218545 -3.212 0.001419 **
scatsecondary -0.955895 0.223573 -4.276 2.36e-05 ***
ucatotherDearee 0.005271 0.115758 0.046 0.963700
ucatoxbridge 0.233832
                           0.131855 1.773 0.076886 .
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
Residual standard error: 1.009 on 420 degrees of freedom
Multiple R-squared: 0.1219. Adjusted R-squared: 0.1093
F-statistic: 9.717 on 6 and 420 DF. p-value: 4.968e-10
```

#### OVB formula example: auxiliary regression

```
> auxiliary = lm(labour ~ treated + scat + ucat, data = d)
> summary(auxiliary)
Call:
lm(formula = labour \sim treated + scat + ucat, data = d)
Residuals:
    Min
             10 Median
                              30
                                      Max
-0.76547 -0.34044 -0.06668 0.34706 0.89441
Coefficients:
               Estimate Std. Error t value Pr(>|t|)
(Intercept)
                0.07963
                          0.10549
                                    0.755 0.45072
treated
               -0.12232 0.04669 -2.620 0.00912 **
scatnotMentioned 0.57331 0.10459 5.482 7.28e-08 ***
scatotherPublic 0.14828 0.09888 1.500 0.13447
scatsecondary 0.48775 0.10116 4.822 1.99e-06 ***
ucatotherDearee 0.11252 0.05237 2.148 0.03225 *
ucatoxbridae
               0.10936
                          0.05966 1.833 0.06749 .
Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' 1
Residual standard error: 0.4565 on 420 degrees of freedom
Multiple R-squared: 0.1786. Adjusted R-squared: 0.1669
```

F-statistic: 15.22 on 6 and 420 DF, p-value: 8.518e-16

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### OVB formula example: confirming equality

So is it true that

$$\beta^{s} = \beta^{l} + \pi_{21}\gamma.$$

i.e. "short equals long plus the effect of omitted times the regression of omitted on included"?

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$$\beta^{s} = \beta^{l} + \pi_{21}\gamma.$$

i.e. "short equals long plus the effect of omitted times the regression of omitted on included"?

```
> all.equal(coef(short)["treated"],
+ coef(long)["treated"] + coef(auxiliary)["treated"]*coef(long)["labour"])
[1] TRUE
```

Yes.

#### Lessons from the OVB formula

Omitting a variable causes bias in our estimate of ATE if and only if

- it is related to the treatment, conditional on other covariates, and
- it is related to the outcome, conditional on other covariates.

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Omitting a variable causes bias in our estimate of ATE if and only if

- ▶ it is related to the treatment, conditional on other covariates, and
- ▶ it is related to the outcome, conditional on other covariates.

#### This is why

- you don't have to control for anything in a randomized experiment
- you don't have to control for everything you can think of that affects Y<sub>i</sub>
   only variables related to D<sub>i</sub> (and Y<sub>i</sub>) conditional on other covariates
- you don't have to control for anything other than the running variable in an RDD

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$$X_{1i} = \alpha^a + \pi_{12} X_{2i} + e_i^a,$$

getting the residuals,

$$\tilde{X}_{1i} = X_{1i} - (\hat{\alpha}^a + \hat{\pi}_{12} X_{2i}),$$

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and regressing  $Y_i$  on those ("outcome-on-residuals" regression):

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and regressing  $Y_i$  on those ("outcome-on-residuals" regression):

$$Y_i = \alpha^* + \beta^* \tilde{X}_{1i} + e_i^*.$$

i.e., 
$$\beta^* = \beta^I$$
.

#### Regression anatomy: "reverse auxiliary" regression

```
> reverse.auxiliary = lm(treated ~ labour + scat + ucat, data = d)
> summary(reverse.auxiliary)
Call:
lm(formula = treated \sim labour + scat + ucat. data = d)
Residuals:
   Min
            10 Median
                           30
                                  Max
-0.8272 -0.3891 -0.2577 0.5499 0.7952
Coefficients:
                 Estimate Std. Error t value Pr(>|t|)
(Intercept) 0.827168 0.101705 8.133 4.77e-15 ***
labour
               -0.131437 0.050172 -2.620 0.009119 **
scatnotMentioned -0.322600 0.111115 -2.903 0.003887 **
scatotherPublic -0.371681 0.101160 -3.674 0.000269 ***
scatsecondary -0.432604 0.105632 -4.095 5.06e-05 ***
ucatotherDearee -0.005414 0.054588 -0.099 0.921037
ucatoxbridge -0.058278 0.062022 -0.940 0.347949
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
Residual standard error: 0.4732 on 420 degrees of freedom
Multiple R-squared: 0.07118, Adjusted R-squared: 0.05791
F-statistic: 5.364 on 6 and 420 DF, p-value: 2.364e-05
> resids.from.ra = resid(reverse.auxiliary)
```

### Regression anatomy: "outcome-on-residuals" regression

```
> star.reg = lm(d$lnrealgross ~ resids.from.ra)
> summary(star.rea)
Call:
lm(formula = d$Inrealgross ~ resids.from.ra)
Residuals:
   Min
            10 Median 30
                                  Max
-4.0169 -0.4474 -0.0796 0.4241 3.6068
Coefficients:
              Estimate Std. Error t value Pr(>|t|)
(Intercept) 12.6190 0.0511 246.939 < 2e-16 ***
resids.from.ra 0.3709 0.1089 3.406 0.000721 ***
Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
Residual standard error: 1.056 on 425 degrees of freedom
Multiple R-squared: 0.02658, Adjusted R-squared: 0.02429
F-statistic: 11.6 on 1 and 425 DF, p-value: 0.0007208
> all.equal(coef(long)["treated"], coef(star.reg)["resids.from.ra"], check.attributes = F)
Γ17 TRUE
```

### Regression anatomy: "outcome-on-residuals" regression

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> all.equal(coef(long)["treated"], coef(star.reg)["resids.from.ra"], check.attributes = F)
Γ17 TRUE
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#### It works!

#### Lessons from regression anatomy

The OLS coefficient on  $D_i$  measures the relationship between  $Y_i$  and the part of  $D_i$  not "explained" by  $X_i$ .

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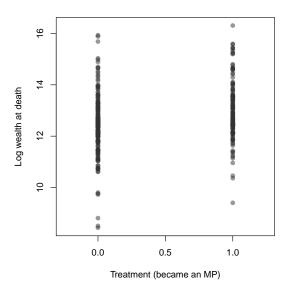
What is the CIA in any regression that claims to measure the effect of  $D_i$  on  $Y_i$ ?

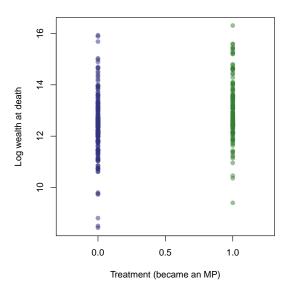
#### Lessons from regression anatomy

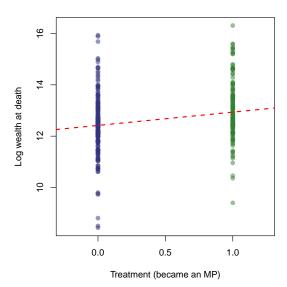
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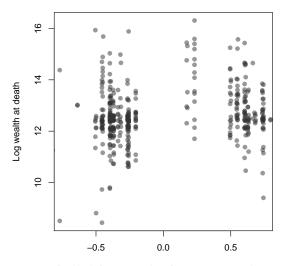
What is the CIA in any regression that claims to measure the effect of  $D_i$  on  $Y_i$ ?

The part of  $D_i$  not "explained" by  $X_i$  (the residual from the "reverse auxiliary regression") is not related to the potential outcomes, i.e. as-if random.

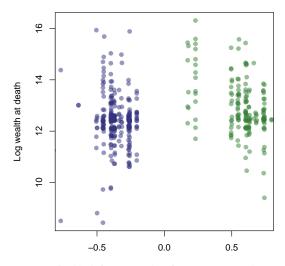




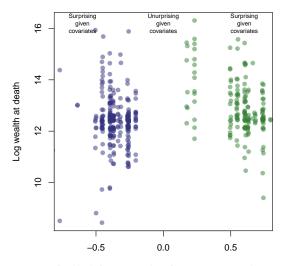




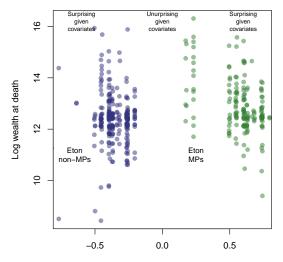
Residuals from regression of treatment on covariates



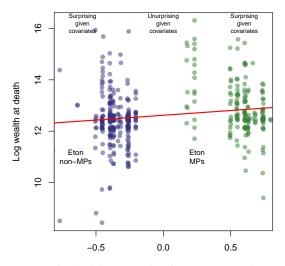
Residuals from regression of treatment on covariates



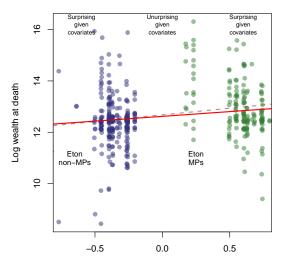
Residuals from regression of treatment on covariates



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