Lecture 9: Matching

Introduction to Econometrics,Spring 2025

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Review the Last Two Lectures

Internal v.s. External Validity

- There are five primary threats to the internal validity of a multiple regression study:

  1. Omitted variables
  2. Functional form misspecification
  3. Errors in variables (measurement error in the regressors)
  4. Censored, Trancated and Selection Samples
  5. Simultaneous causality

- The data structure may violate the 2th OLS regression assumption, thus **random sampling**.

  ◦ adjusted the s.e. by clustering or other methods.

- Last but not least, the economic **magnitude** of $\hat{\beta}$ matters.

  ◦ economic significance is as important as statistical significance.

OLS and Controls

- The main identification strategy of OLS regression is **Control**, ie. putting covariates into the regression as control variables.

- The main identifying assumption of an OLS regression is

- Essentially, the strategy compares treatment and control subjects who have **the same observable characteristics**, which is often called **Selection on Observables**.

    - Besides the regression, we can also use **matching** to achieve this goal.

Matching: Introduction

Introduction

- In **observational** studies, as opposed to RCTs, we cannot directly determine the causal effect because the **counterfactual** outcome of the treated group is unknown.

  - In other words, we **cannot find a suitable control group** to compare with the treated group.

- The idea of **matching** method is quite simple:

- For simplicity, we focus on the former question, ie. **constructing a control group**, which is more common in practice.

Introduction

- Suppose Y_{i1} and Y_{i0} are the outcomes of the treated and untreated group, respectively.

- And we can use some or all samples from untreated group to construct the **counterfactual outcomes** of the treated group Y_{i1}^c

- Then the average treatment effect(ATE or ATT) easily by making the difference

- **Question**: How can we use samples from the untreated group to get the counterfactual outcomes of the treated group, Y_{1i}^c?

- **Answer**: select the untreated samples that are **similar to the treated ones** in terms of **the covariates** X_i

- **Assumption**: If CIA holds, thus $(Y_1, Y_0) \perp\!\!\!\perp D \,|\, X$, then the treatment status can be seen as randomized given the covariates X_i.

Example: Training Program Evaluation

- **Question**: What is the causal effect of a training program on the wage of workers?

- A simple OLS regression model can be written as

$$Y_i = \beta_0 + \beta_1 D_i + u_i$$

- The treated group is the workers who have received the training program $D = 1$

- The untreated group is the workers who have not received the training program $D = 0$

- The outcome is the log-wage of workers Y_i, and the covariates X_i form a vector including variables such as age, education, experience, etc.

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- The average wage gap between the treated group and the untreated group is

OLS Regression for the Training Program

- The main identifying assumption of the OLS regression here is

    - Conditional Independence Assumption(CIA): which means that if we can "balance" covariates X then we can take the treatment D as randomized.

- However, we may still suffer the **misspecification** of the model under the CIA, which can also make estimates β_1 biased.

A Trainning Example: matching samples

- Assume that the covariates X is the **age of the workers**, and to see **how the matching method works**.

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16	43	10700	16	24	9700			
17	28	11500	17	29	6200			
18	27	10700	18	35	30200			
19	28	16300	19	32	17800			
Average:	28.5	16426	20	23	9500	Average:		
			21	32	25900			
			Average:	33	20724			

Trainees			Non-Trainees			Matched Sample		
unit	age	earnings	unit	age	earnings	unit	age	earnings
1	28	17700	1	43	20900	8	28	8800
2	34	10200	2	50	31000	14	34	24200
3	29	14400	3	30	21000	17	29	6200
4	25	20800	4	27	9300	15	25	23300
5	29	6100	5	54	41100	17	29	6200
6	23	28600	6	48	29800	20	23	9500
7	33	21900	7	39	42000	10	33	15500
8	27	28800	8	28	8800	4	27	9300
9	31	20300	9	24	25500	12	31	26600
10	26	28100	10	33	15500	11,13	26	8450
11	25	9400	11	26	400	15	25	23300
12	27	14300	12	31	26600	4	27	9300
13	29	12500	13	26	16500	17	29	6200
14	24	19700	14	34	24200	9,16	24	17700
15	25	10100	15	25	23300	15	25	23300
16	43	10700	16	24	9700	1	43	20900
17	28	11500	17	29	6200			
18	27	10700	18	35	30200			
19	28	16300	19	32	17800			
Average:	28.5	16426	20	23	9500	Average:		
			21	32	25900			
			Average:	33	20724			

A Trainning Example: matching samples

	Trainees			Non-Trainees			Matched Sample	
unit	age	earnings	unit	age	earnings	unit	age	earnings
1	28	17700	1	43	20900	8	28	8800
2	34	10200	2	50	31000	14	34	24200
3	29	14400	3	30	21000	17	29	6200
4	25	20800	4	27	9300	15	25	23300
5	29	6100	5	54	41100	17	29	6200
6	23	28600	6	48	29800	20	23	9500
7	33	21900	7	39	42000	10	33	15500
8	27	28800	8	28	8800	4	27	9300
9	31	20300	9	24	25500	12	31	26600
10	26	28100	10	33	15500	11,13	26	8450
11	25	9400	11	26	400	15	25	23300
12	27	14300	12	31	26600	4	27	9300
13	29	12500	13	26	16500	17	29	6200
14	24	19700	14	34	24200	9,16	24	17700
15	25	10100	15	25	23300	15	25	23300
16	43	10700	16	24	9700	1	43	20900
17	28	11500	17	29	6200	8	28	8800
18	27	10700	18	35	30200			
19	28	16300	19	32	17800			
Average:	28.5	16426	20	23	9500	Average:		
			21	32	25900			
			Average:	33	20724			

A Trainning Example: matching samples

	Trainees			Non-Trainees			Matched Sample	
unit	age	earnings	unit	age	earnings	unit	age	earnings
1	28	17700	1	43	20900	8	28	8800
2	34	10200	2	50	31000	14	34	24200
3	29	14400	3	30	21000	17	29	6200
4	25	20800	4	27	9300	15	25	23300
5	29	6100	5	54	41100	17	29	6200
6	23	28600	6	48	29800	20	23	9500
7	33	21900	7	39	42000	10	33	15500
8	27	28800	8	28	8800	4	27	9300
9	31	20300	9	24	25500	12	31	26600
10	26	28100	10	33	15500	11,13	26	8450
11	25	9400	11	26	400	15	25	23300
12	27	14300	12	31	26600	4	27	9300
13	29	12500	13	26	16500	17	29	6200
14	24	19700	14	34	24200	9,16	24	17700
15	25	10100	15	25	23300	15	25	23300
16	43	10700	16	24	9700	1	43	20900
17	28	11500	17	29	6200	8	28	8800
18	27	10700	18	35	30200	4	27	9300
19	28	16300	19	32	17800	8	28	8800
Average:	28.5	16426	20	23	9500	Average:		
			21	32	25900			
			Average:	33	20724			

An Illustrated Example: matched samples

Trainees				Non-Trainees				Matched Sample		
unit	age	earnings		unit	age	earnings		unit	age	earnings
1	28	17700		1	43	20900		8	28	8800
2	34	10200		2	50	31000		14	34	24200
3	29	14400		3	30	21000		17	29	6200
4	25	20800		4	27	9300		15	25	23300
5	29	6100		5	54	41100		17	29	6200
6	23	28600		6	48	29800		20	23	9500
7	33	21900		7	39	42000		10	33	15500
8	27	28800		8	28	8800		4	27	9300
9	31	20300		9	24	25500		12	31	26600
10	26	28100		10	33	15500		11,13	26	8450
11	25	9400		11	26	400		15	25	23300
12	27	14300		12	31	26600		4	27	9300
13	29	12500		13	26	16500		17	29	6200
14	24	19700		14	34	24200		9,16	24	17700
15	25	10100		15	25	23300		15	25	23300
16	43	10700		16	24	9700		1	43	20900
17	28	11500		17	29	6200		8	28	8800
18	27	10700		18	35	30200		4	27	9300
19	28	16300		19	32	17800		8	28	8800
Average:	28.5	16426		20	23	9500		Average:	28.5	13982
				21	32	25900				
				Average:	33	20724				

- The average wage gap between the treated group and the **matched untreated group** is

Two Assumptions: One Old and One New

- We still rely on the **Conditional independence Assumption(CIA)**, which is akin to running an OLS regression.

- More specifically, we assume that the potential incomes for the workers are **independent of th e training status** given the age of the workers.

- It means that if CIA are not satisfied, then both the OLS and the matching estimator will be biased.

  - Matching is not a *silver bullet* for OVB in OLS.

- Besides, do you notice that **there are some untreated samples that are not matched with any treated samples**?

- This is due to the **Overlap Assumption**, a new assumption in the matching method that was not

Two Assumptions: One Old and One New

- The **Overlap Assumption** is to ensure that we can find a matched untreated sample for each treated sample.Mathematically, it is expressed as:

- This implies that the likelihood of receiving treatment is neither 0 nor 1 for any given covariates.

- Including either case in our comparison would bias the average treatment effect estimation.

- It suggests that we **change the samples explicitly** based on the covariates to ensure that the **overlap assumption** is satisfied.

A Trainning Example: before matching
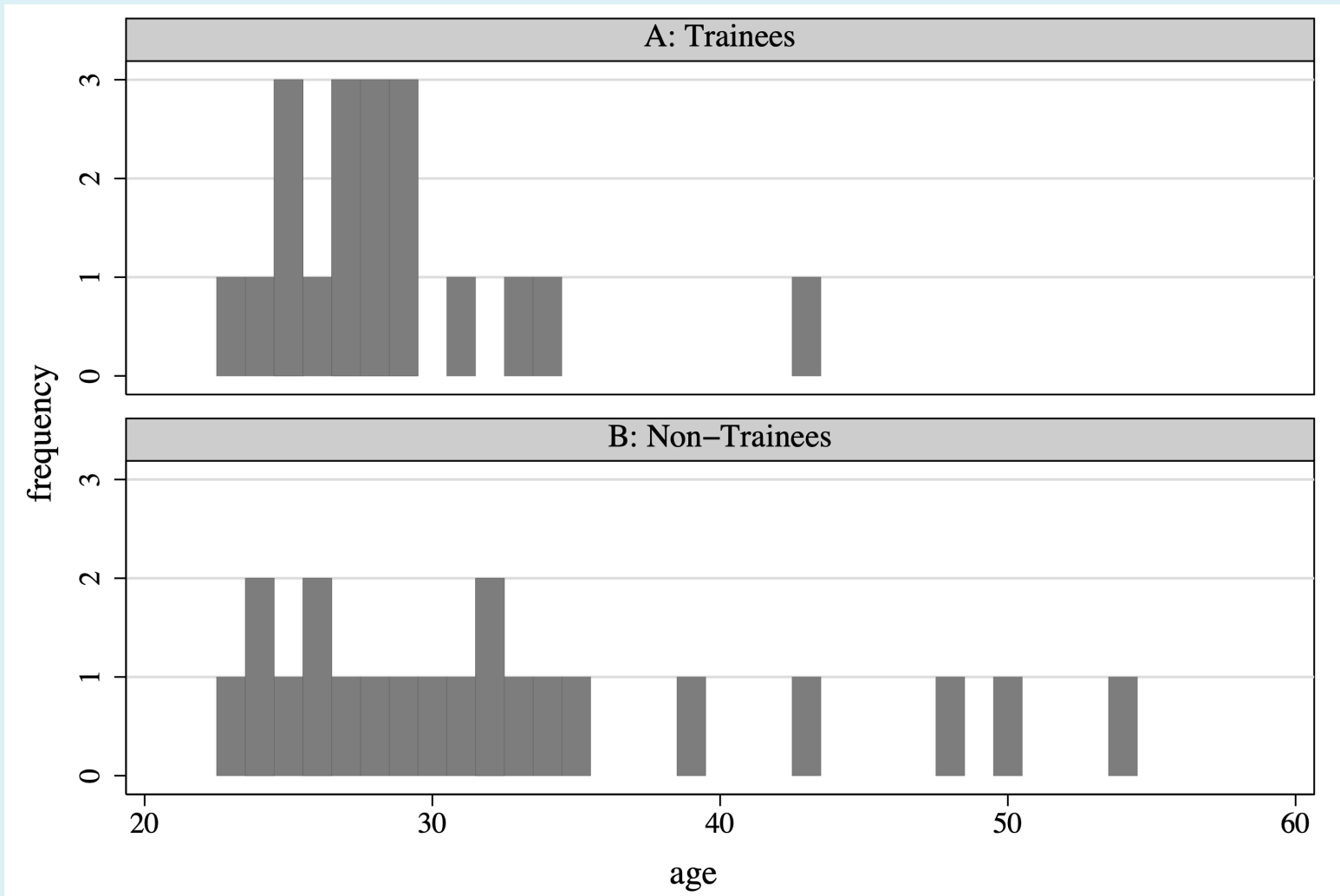
A Trainning Example: after matching
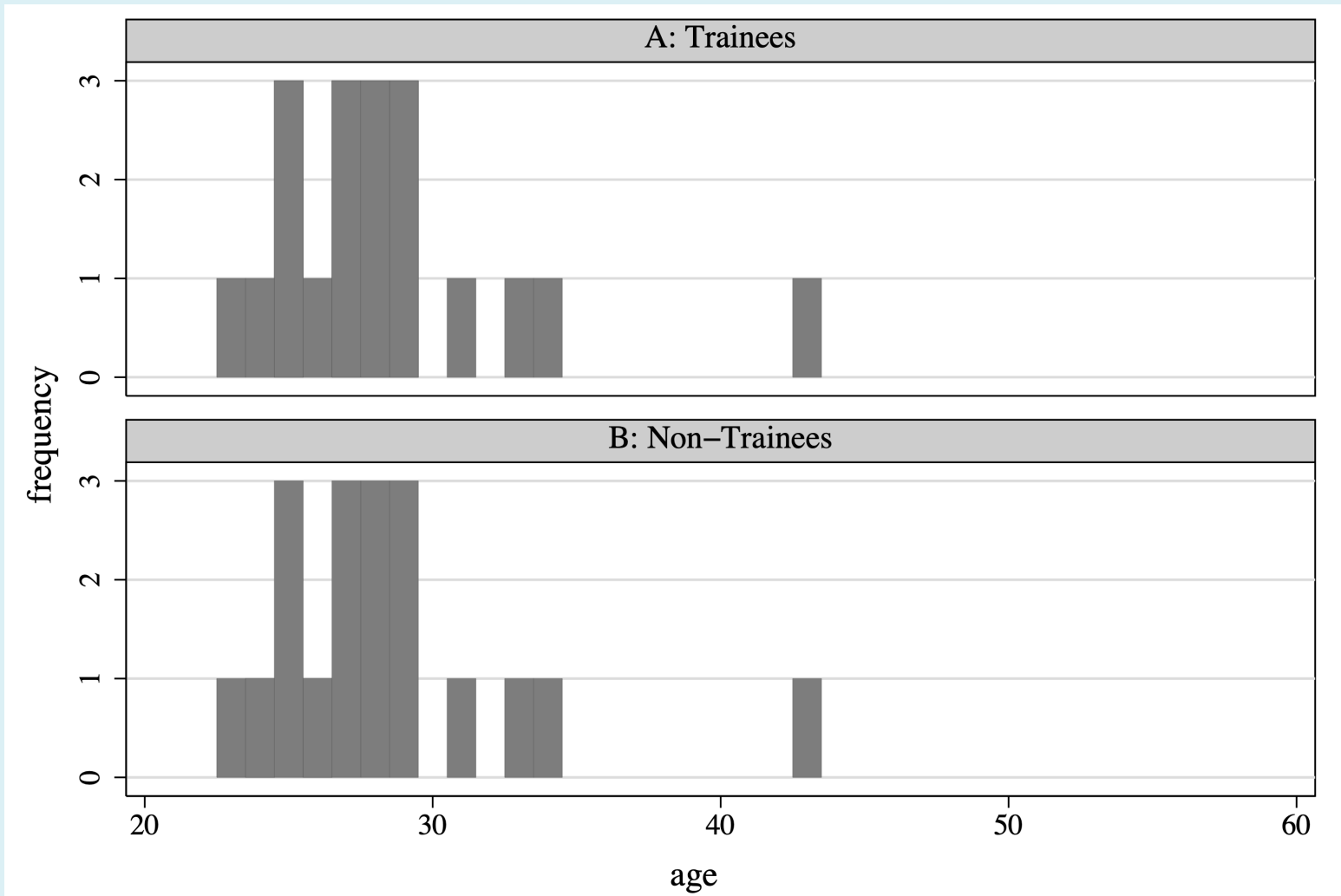
The Overlap Assumption in OLS
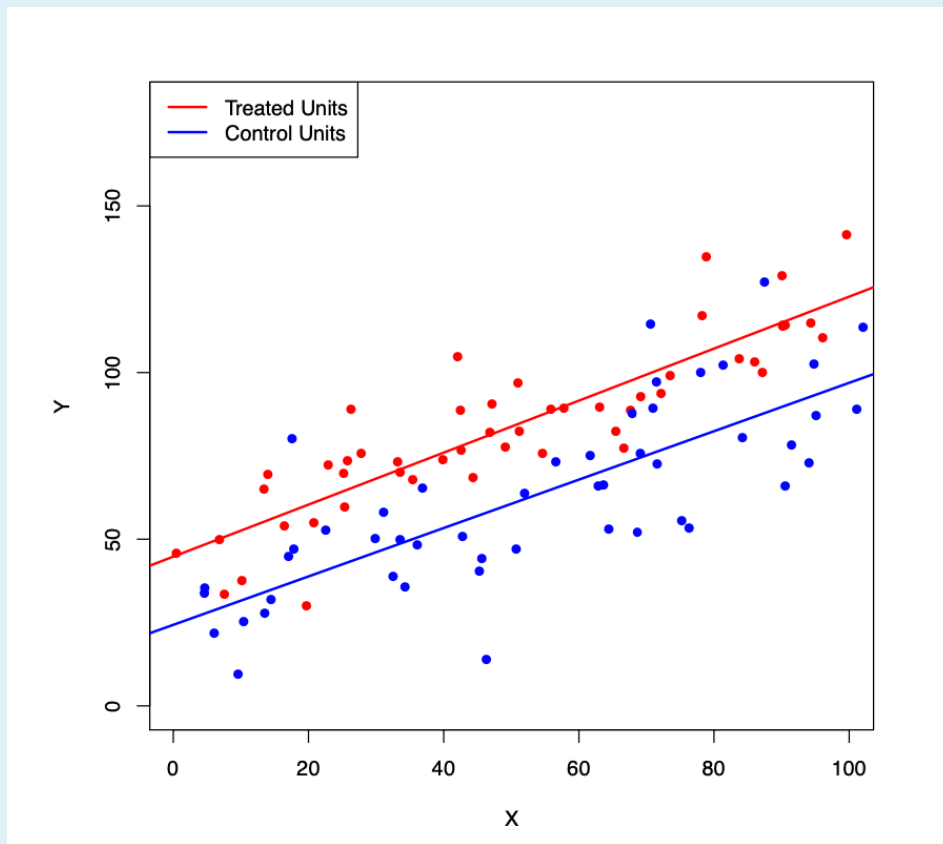
The overlap assumption is satisfied
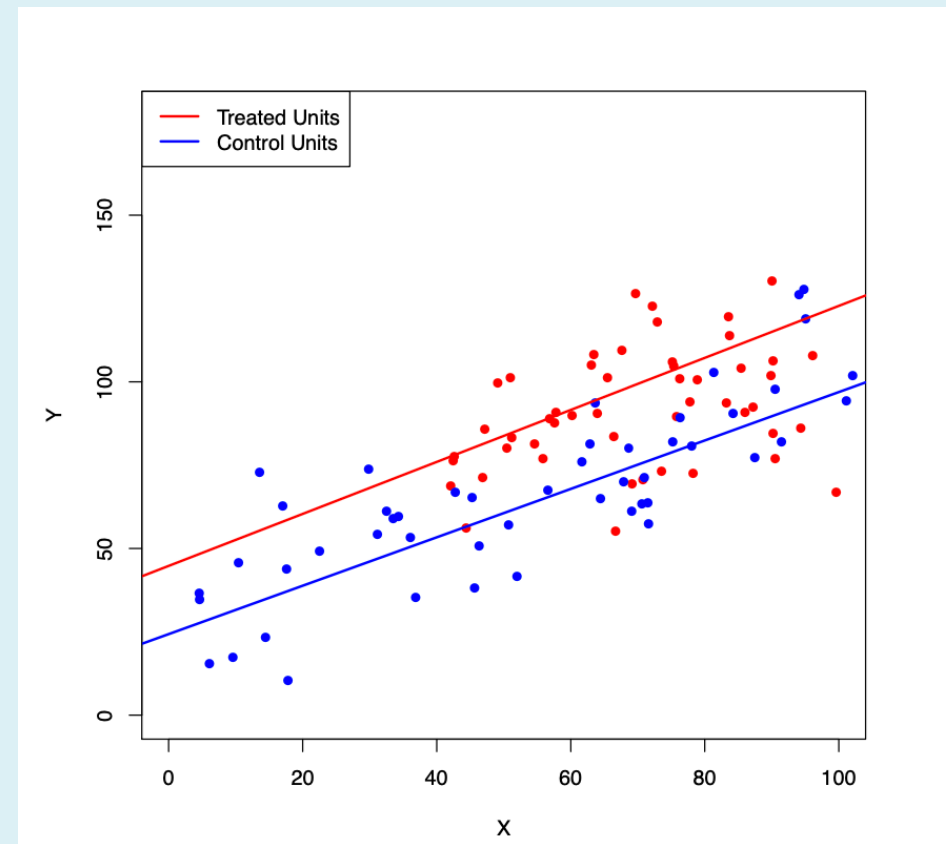
The overlap assumption is violated

- In the OLS regression, the overlap assumption is not **explicitly required**, which may lead to biased estimates.

Matching Estimators: Exact matching is hard

- The training case is an example of **Exact matching** which means that only units with identical covariate values are used to construct the control group.

- But what if we have multiple covariates using to match, thus $X = (X_1, X_2, \ldots X_k)'$?

  - In this case, it is **impossible** to find proper units with identical values in all covariates $X_1, X_2, \ldots X_k$.

- Two complementary solutions are running in parallel, representing the directions in which the matching method is developing.

1.
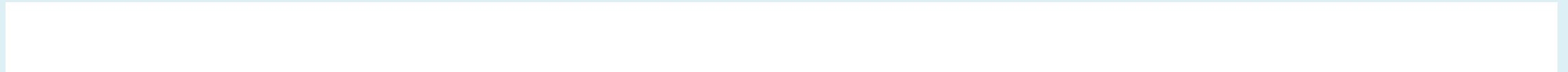
2.

Matching Estimator

Introduction

- The matching estimator can be divided into three steps: **Matching**, **Estimation** and **Inference**.

- Matching: Find a control group for each treated individual based on the covariates.

- Estimation: Estimate the average treatment effect(making a difference) using the matched samples.

- Inference: Test the statistical significance of the treatment effect(ATT or ATE) using the matched samples.

Reweight as Counterfactuals

- Basic settings: all notations are the same as before, like Y_{1i}, Y_{0i}, D_i, and X_i.

  - the sample size here is the only one need to noted : N_T treated individuals and N_C control individuals.

- The **counterfactual for treated individual** i that what we want is Y_{1i}^C, then **how to construct it by matching**?

- Because we construct the *counterfactuals* by using the **untreated samples**, therefore in a more general sense, the counterfactual for treated individual i is

- where $w_i(j)$ is a **weight** of untreated individual j for treated individual i, and normally $\sum_j w_i(j) = 1$

Matching Estimator

- Then individual treatment effect, δ_i, is

- A matching estimator for the average treatment effect on the treated(ATT) is

- Where C is the common support region of the treated and untreated individuals.

- And $j = 1, 2, \ldots, N^C$ and $i = 1, 2, \ldots, N^T$.

- $i \in (D = 1 \cap C)$ means that i is a treated individual and i is in the common support region.

Weight to Matching

- Question: **How to obtain these weights**, thus $w_i(j)$?

- **Answer**: *It is easy and hard at the same time.*

- E.g. if $w_i(j) = \begin{cases} 1 & \text{if } j = i \\ 0 & \text{otherwise} \end{cases}$

- In this case, the weights are **equal** for all the untreated samples.

- Then we're back to a **difference in means**, except now it's based on the N_T matched samples.

Weight to Matching

- **More Reasonable Weights**: The weights $w_i(j)$ should be **related with covariates** X_i in treated group and X_j in untreated group.

Proximity: When X is Discrete

- If X is **discrete**, then we can use the equality of X to construct the weights. Thus

- Where I(·) is an indicator function,

- This is the **Exact Matching** what we did in the trainning case.

Proximity: When X is Continuous

- If X is **continuous**, then we may not find a unit with the same covariate values. Then we may need **proximity** rather than **equality**.

- Then the weight $w_i(j)$ can be *a measure of how close* X_j of untreated group is to X_i of the treated group.

- If the gap(distance) is small, then the weight is large, and vice versa.

- Question: What do "**small**" and "**large**" mean in the previous sentence?

  - It depends on.

- If we just pick the smallest one as we did in the trainning case, then we have the **Nearest Neighbor Matching**.

Math Review: Distance between two vectors

- If X_i and X_j are both single-dimensional variables, then the distance between them is the difference between them,

$$|X_i - X_j|$$

- What if X_i and X_j are both multi-dimensional variables, thus k-dimensional vectors as follows

- **Question**: how to measure the distance between two vectors?

- **Answer**: The **Euclidean distance** can be as the measure of the distance between X_i and X_j,

Proximity: When X is a Vector

- The Euclidean distance is not invariant to changes in the **scale** of X. A more commonly used distance is the **normalized Euclidean distance**

- where V_X^{-1} is the symmetric and positive semidefinite variance matrix of X of X, thus

- $\hat{\sigma}_k^2$ is the variance of the k-th variable.

- No scale problem but still *no correlations* between Xs.

Proximity: When X is a Vector

- **Mahalanobis distance** between X_i and X_j is defined as

- where Σ_X^{-1} is the variance-covariance matrix of X.

- $\hat{\sigma}_{jk}$ is the covariance between the j-th and k-th variables.

- No scale problem and **taking correlations between Xs into account**.

Many Matching Methods

- There many methods to choose the matchers and weights. Here are some of them:

- **Exact Matching**:
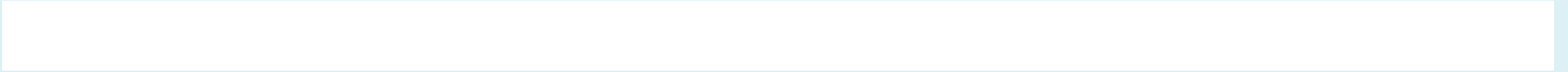
- **Nearest Neighbor Matching(NNM)**:
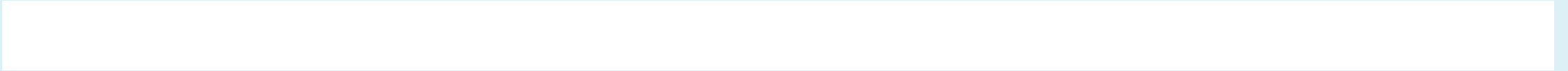
- **Radius Matching**: all the samples within a certain range are matched.
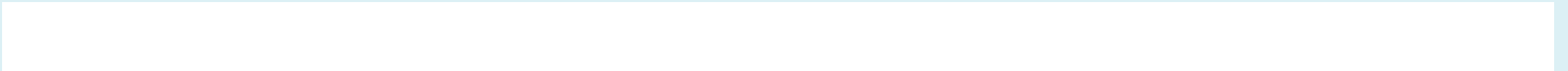
Many Matching Methods
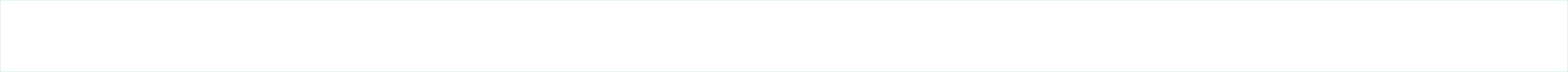
- **Radius Matching**: all the samples within a certain range are matched.

- **Subclassification** : Divide the treated and untreated group into subclasses based on the covariates and then match within each subclass.

- **Kernel Matching**: The weight is based on the kernel function, which is an estimated density function of the covariates.

The curse of dimensionality

- As the dimension of X expands (i.e., matching on more variables), whatever matching method we use, it becomes increasingly difficult to find a suitable or closely matched control for each treated sample, even if we have a large sample size.

- Need alternative ways to shrink the dimensions of X.

- It turns out that if CIA is satisfied,then we actually only need to match/conditional on the **propensity score** $p(x)$, instead of the entire X_i.

Propensity-Score Methods

The Magic of Propensity Scores

- Recall the CIA assumption:

$$\left(Y_{0i}, Y_{1i}\right) \perp\!\!\!\perp D_i \,|\, X_i$$

- The **propensity score** is defined as *the probability of treatment given X_i*, thus

- Formally the **Propensity Score Theorem** is

- If we control/adjust/balance the propensity score instead of the raw covariates, then the treatment is as good as random.

- This theorem extends CIA assumption from **multiple dimensions** to **a one-dimensional score**, avoiding the curse of dimensionality.

Propensity-Score Theorem

Theorem If $\left(Y_{0i},\ Y_{1i}\right) \perp\!\!\!\perp D_i \,|\, X_i,$ then $\left(Y_{0i},\ Y_{1i}\right) \perp\!\!\!\perp D_i \,|\, p(X_i).$

Proof

To prove this theorem, we will show

Propensity-Score Theorem

Theorem If $\left(Y_{0i},\ Y_{1i}\right) \perp\!\!\!\perp D_i \,|\, X_i,\ $ then $\left(Y_{0i},\ Y_{1i}\right) \perp\!\!\!\perp D_i \,|\, p(X_i)$.

Proof

Propensity-Score Theorem

Theorem If $\left(Y_{0i},\ Y_{1i}\right) \perp\!\!\!\perp D_i \,|\, X_i,$ then $\left(Y_{0i},\ Y_{1i}\right) \perp\!\!\!\perp D_i \,|\, p(X_i).$

Proof

Propensity-Score Matching

Intuition

- **Question**: X_i carries way more information than $p(X_i)$, so how can we still get conditional independence of treatment by only conditioning on $p(X_i)$?

- **Answer** Conditional independence of treatment is not about *extracting all of the information* possible from X_i. We actually *only care about creating a situation* in which $D_i \mid$ a function of X is independent of $\left(Y_{0i}, Y_{1i}\right)$.

Propensity-Score Matching

Estimation: Binary Dependent Regression

- **Question**:How to obtain the propensity scores $p(X_i)$?

- Recall the definition of propensity score, **does it sound familiar?**

- Yes,it is the **binary dependent regression** model that the independent variables are the covariates X_i.

- As we have learned in the previous lecture, we can estimate the propensity scores using three models:

- Of course there are another ways to estimate it like machine learning methods, but the most common way is to use **logit** regression.

Propensity-Score Matching

Estimation: Logit Regression

- The logit model of the propensity score is given by

- Where $\mathbf{X}_i$ is the vector of covariates and β is the vector of coefficients.

- Then we could get the estimated propensity scores $\hat{p}(\mathbf{X}_i)$ by plugging in the estimated coefficients $\hat{\beta}$.

Propensity-Score Matching

Estimation: Logit Regression

- However, for the nonlinearity of the model, the marginal effect of coviarites on the propensity score is not constant.

    - It means that *the same change of the covariates will not have the same effect* on the propensity score for all the values of the covariates.

- Therefore, a more common way to estimate the propensity score is to use the **log odds ratio**,

- Recall: We claimed that matching is over regression as it is **non-parametric**, don't need to specify the functional form of the model.

- However, in the propensity score method, we still need to **specify the functional form of the model and estimate the coefficients**.

Propensity-Score Matching

Estimation: Predicted instead of Explained

- Note: The focus in the model here is a little bit different from the one we learned in the binary dependent variable regression.

  - Here we focus on the predicted probability of being treated, which is the **propensity score**, and the covariates are the explanatory variables.

  - While in the binary dependent variable regression, we focus on the explanatory coefficient of the covariates(only one or two in most cases) on the treated variable(which actually is the dependent variable).

- Therefore, when we estimate the propensity score by the logit model, the function form should be **as flexible as possible** to capture the relationship between the covariates and the treatment variable.

  - Polynomial terms and interaction terms are often included in the model.

  - Even ML methods can be used to estimate the propensity score as well.

Overlap Assumption in Propensity Score Methods

- Recall: The **Overlap Assumption**

- Which is to ensure that we can find a matched untreated sample for each treated sample, or the distribution of X for the treated and control groups should overlap.

- In P-score methods, the overlap assumption is about the distribution of the propensity score rather than the covariates.

- The easiest way to check the overlap assumption is to **plot the distribution of covariates before and after matching**.

  ○ As we did it in the training example, in which we plotted the distribution of only one covariate.

  ○ Apparently when X is a vector which can be tough as the dimensions of X expand.
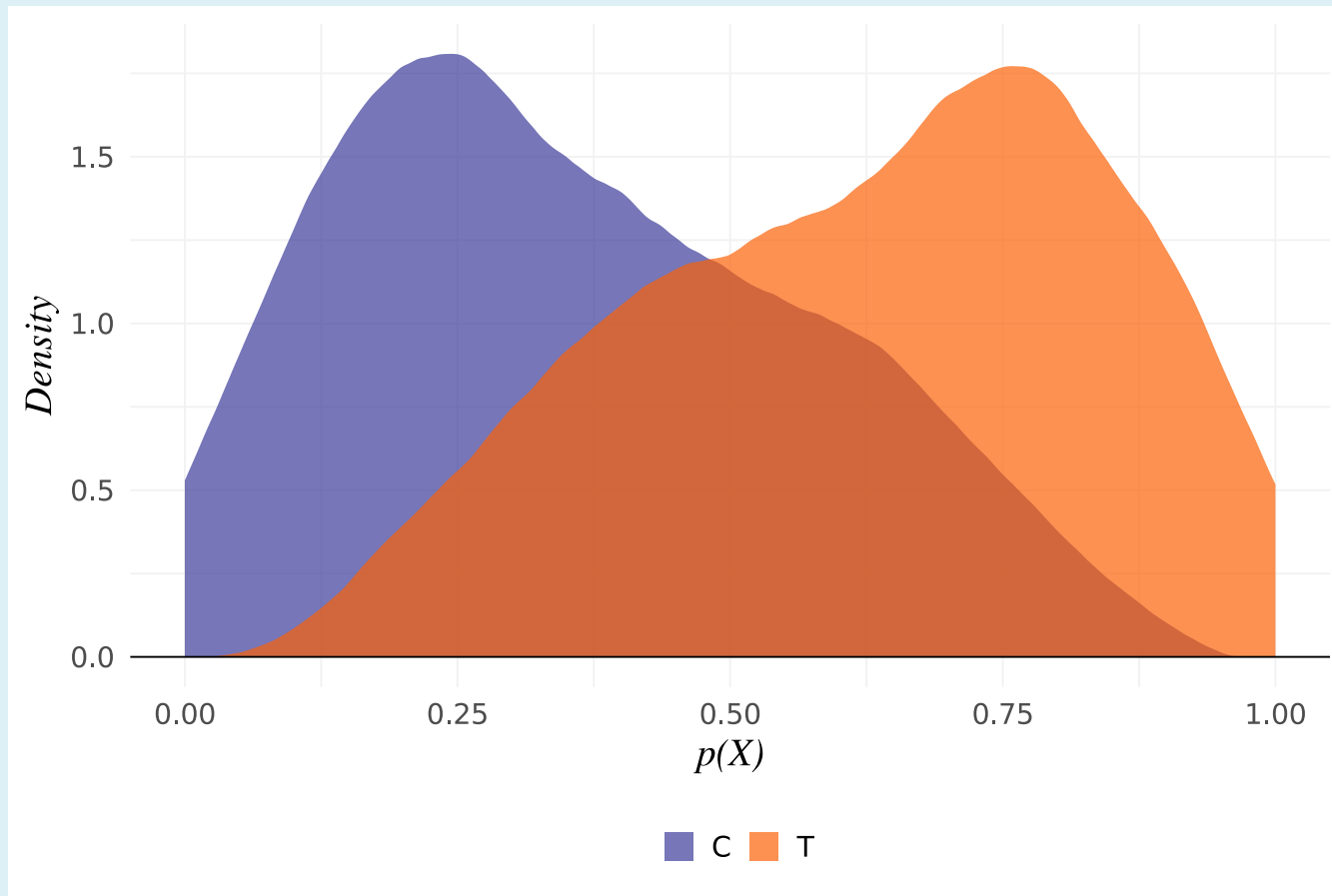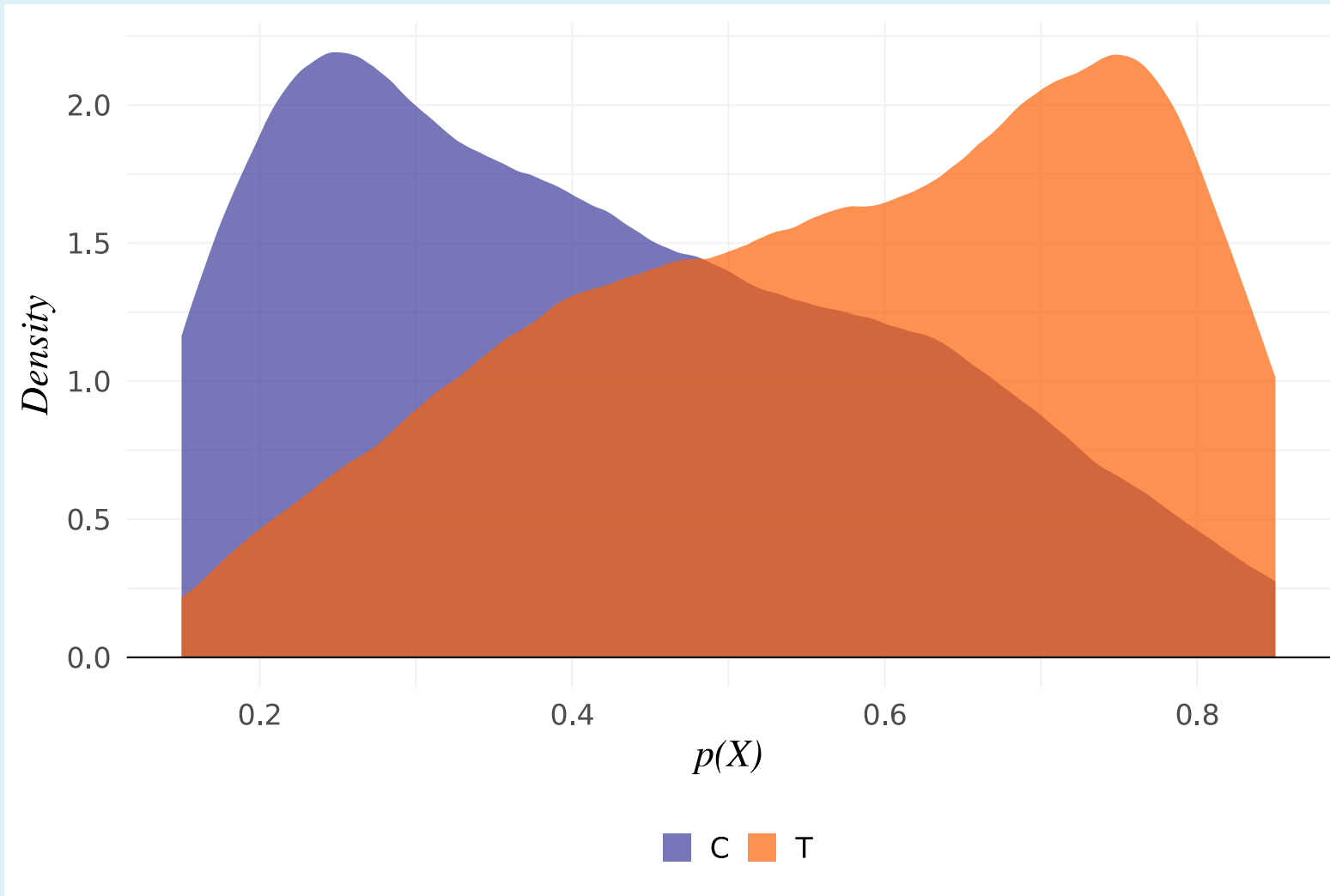
- Trimming samples to overlap in $p(X_i)$, thus we only keep the samples if $0.15 \leq p(X) \leq 0.85$

Regression and Propensity Scores Reweight

Regression with P-Scores

- Based on the **Propensity Score Theorem**, conditional on the propensity score, the treatment is as good as random.

- Then, the simple idea is to use propensity scores as a control variable instead of the raw covariates in the regression model

- Assumption: the relationship between the outcome and the propensity score is linear.

- To consider the non-linearity, we can add the polynomial terms or interaction terms between the propensity score and the treatment to make a more flexible model.

- Normally, the cubic term is enough for the flexibility.

Inverse Probability Weighting

- The **inverse probability weighting (IPW)** is an alternative way to use the propensity score to control the bias due to the selection on observables.

    ○ The idea is to weight the treated and control units by the inverse of the propensity score.

- The Average Treatment Effect (ATE) can be derived by the following formula:

- Under the CIA and Overlap Assumption, we could show that

Inverse Probability Weighting

- Thus, we have the following result:

$$E[Y_{1i}] = E\left[\frac{D_i}{p(X_i)}Y_i\right]$$

IPW Estimator for ATE

- Similarly, we could show that

- Then, we could get the ATE by the following

- This is the *Horvitz-Thompson* IPW estimator for the ATE.

IPW Estimator for ATE

- Then the IPW estimator for ATE is given by

- The IPW weights here are the **inverse of the propensity score**.

- HW weights do not necessarily average to 1,which can be a problem.

A more general IPW Estimator for ATE

- The standardization means dividing each group's weights by the sum of all weights within that group.

- A more general IPW estimand is given by

- Where $E(\frac{D_i}{p(X_i)})$ can be seen as the average weight for the treated group and $E(\frac{1-D_i}{1-p(X_i)})$ can be seen as the average weight for the control group.

- Then corresponding IPW estimator for ATE is given by

Practical Implications

- IPW provides a way to estimate causal effects without explicitly modeling the outcome process like matching.

- Practical challenges:

  - Extreme weights when $p(X)$ is close to 0 or 1
  - Need for careful diagnostics (covariate balance, weight distribution)

- Some Extensions:

- Double Robustness: Combine IPW with outcome regression

  - Consistent if either the propensity score is correctly specified or the outcome regression is correctly specified.

Matching in practice

Matching in Practice

Introduction

- Although matching is a simple concept, it can be more difficult to implement in practice.

- There are many decisions to make when matching units. The questions are as follows:

  1. How to choose variables as the matching covariates?

  2. Which matching methods should be used? distances and weights: Matching/Propensity Score Matching

  3. How many control units should be matched to each treatment unit?: one-to-one or many-to-one?

  4. The sample is matched with or without replacement?

  5. The order of matching: greedy or optimal?

Matching in Practice

Choosing Variables

- **Question**: Which variables should be used for matching treatment and control units?

- **Answer**: Include all variables that are likely to be confounders. (*Recall the "good and bad controls" framework*)

  - Irrelevant variables
  - Relevant variables
  - Omitted variables
  - Colliders
  - Confounders

- Selecting matching covariates follows similar principles as in regression analysis.

- As with OLS regression, comparing results across different sets of variables serves as a **sensitivity analysis**.

Matching in practice

With or Without Replacement

- Matching with replacement means that control units can be used as a match for **more than once**.

  - each control unit is "placed back" into the controls after being used once.

- Two advantages:

  - treatment and control units after matching will be better balanced.
  - the order in which we match the units does not matter, in turn the matching algorithm is reduced in complexity.

- Nonetheless, it is very common to match with replacement.

Matching in practice

Greedy v.s Optimal Matching

- The greedy matching is a simple and fast algorithm that matches each treated unit to the control unit with the closest distance.

- However, the closest control units for every single sample may not be the best match for the treated unit as a whole.

  - Thus the local optimal solution may not be the global optimal solution.

- The optimal matching is a more complex algorithm that finds the best possible match for each treated unit simultaneously.

  - It is often computationally expensive because it have to consider all possible matches for all treated units.

Matching in practice

1:1 v.s 1:m Matching

- **1:1 matching**: each treated unit can be matched to only one control.

- **1:m matching**: each one can be matched to more than one control.

- Benefit: This can be useful in large samples where there are more control units than treated units, because the inclusion of more units will increase the precision of our estimates.

- Cost: often the second, third and fourth matches may be poorer than the first match, meaning that we may end up including control units that are not very similar to the treatment

Matching in practice

Assessing Balance

- As in RCTs, after carrying out matching we should first carry out balance tests to compare the treatment and control units.

- If matching was successful, then by definition they should be very similar to each other in terms of their covariates.

- Balance tests are particularly useful in matching because they might be able to help us choose between different distance metrics or matching with vs. without replacement.

- Normally, matching procedures need a relatively large number of samples to be able to find a good match.

Matching in practice

In a Summary

- If matching was successful, then by definition they should be very similar to each other in terms of their covariates.

- Balance tests are particularly useful in matching because they might be able to help us choose between different distance metrics, matching with vs. without replacement.

- Choosing the **"best"** matching method highly depends on the unique characteristics of the dataset as well as the goals of the analysis.

    - Similar to the logic of Machine learning

- Therefore, sensitivity analysis is very crucial to Matching.

Wrap up

- Both matching and regression rely on CIA (selection on observables). Most biases we could suffer in regression, such as OVB, measurement error, and simultaneous causality, will not be avoided even if we use matching.

- Most importantly, matching is essentially as the same as regression, only different in the weight of estimating the CEF function.

- Question: Why we still need matching?

- Answer:

- In practice, using matching alone as main identification strategy is less common in economics, more frequently combined with other methods like DID and SCM, which we will discuss later on.

Appendix: Matching vs Regression

Matching in essential is Regression

- Although matching is a non-parametric or semi-parametric method, it is essentially as the same as regression.

- Suppose the ATT, thus the average treatment effect on the treated, is the parameter of interest.

$$ATT = \delta_{ATT} = E[Y_{i1} - Y_{i0} | D = 1]$$

- Under the CIA, if we can control/balance some covariates X_i, then we have the selection bias equals to zero, thus

$$E[Y_{0i} | X_i, D = 1] = E[Y_{i0} | X_i, D = 0]$$

Matching in essential is Regression

- Using the LIE and CIA,

$$\delta_{ATT} = E\big[(E[Y_{i1}|X_i, D = 1] - E[Y_{i0}|X_i, D = 1])|D_i = 1\big]$$

$$= E\big[(E[Y_{i1}|X_i, D = 1] - E[Y_{i0}|X_i, D = 0])|D_i = 1\big]$$

$$= E\big[(E[Y_i|X_i, D = 1] - E[Y_i|X_i, D = 0])|D_i = 1\big]$$

$$= E\big[\delta_X|D_i = 1\big]$$

- Where δ_X is the average outcomes gaps between two groups within observed covariates X_i.

$$\delta_X = E[Y_i|X_i, D = 1] - E[Y_i|X_i, D = 0]$$

Matching in essential is a regression

- Recall: the Bayes' rule

$$P\left(X_i = x \mid D_i = 1\right) = \frac{P\left(D_i = 1 \mid X_i = x\right) \cdot P\left(X_i = x\right)}{P\left(D_i = 1\right)}$$

- Then if X_i is discrete, then the matching estimator can be written as

$$\delta_M = E\left[\delta_X \mid D_i = 1\right] = \Sigma_x \delta_X P(X_i = x \mid D_i = 1)$$

$$= \sum_x \delta_x \frac{P\left(D_i = 1 \mid X_i = x\right) P\left(X_i = x\right)}{P\left(D_i = 1\right)}$$

$$= \sum_x \delta_x \left[\frac{P\left(D_i = 1 \mid X_i = x\right) P\left(X_i = x\right)}{\Sigma_x P\left(D_i = 1 \mid X_i = x\right) P\left(X_i = x\right)}\right]$$

Regression in essential is a matching

- Suppose we have a saturated regression model

$$Y_i = \sum_x I(X_i = x)\beta_x + \delta_R D_i + u_i$$

- The $I(X_i = x)$ is the indicator function for $X_i = x$, which means that

$$I(X_i = x) = \begin{cases} 1 & \text{if } X_i = x \\ 0 & \text{otherwise} \end{cases}$$

- D is the treatment status, u_i is the error term.

- The β_x is the coefficient of $X_i = x$ in the regression model.

- The δ_R is the regression estimator of the treatment effect.

- **Note**: Saturating X means allowing a separate intercept for each unique value of X_i, which makes the conditional expectation function $E[Y|X]$ to be linear in X.

Regression in essential is a matching

- We can prove that the regression estimator δ_R can be expressed as follows†

$$\delta_R = E\left[\frac{Var(D_i|X_i)}{E\left[Var(D_i|X_i)\right]}\delta_X\right]$$

- Where $Var(D_i|X_i)$ is the conditional variance of D_i given X_i, thus

$$Var(D_i|X_i) = E\left[(D_i - E[D_i|X_i])^2 \mid X_i\right]$$

- Let $p(x) = P(D_i = 1 | X_i = x)$, then

$$Var(D_i|X_i) = p(x)(1 - p(x)) = P\left(D_i = 1 \mid X_i = x\right)(1 - P\left(D_i = 1 \mid X_i = x\right))$$

†The detailed proof is somewhat complex, so it's placed in the appendix for those interested. You can also refer to the Mostly Harmless Econometrics (MHE) textbook for the proof (pp54-55).

Regression in essential is a matching

- Then the regression estimator δ_R

$$\delta_R = E\left[\frac{Var(D_i|X_i)}{E[Var(D_i|X_i)]}\delta_X\right]$$

$$= \frac{E[Var(D_i|X_i)\delta_X]}{E[Var(D_i|X_i)]}$$

$$= \frac{\sum_x \delta_X \cdot Var(D_i|X_i) \cdot P(X_i = x)}{\sum_x \cdot Var(D_i|X_i) \cdot P(X_i = x)}$$

$$= \sum_x \delta_X\left[\frac{P\big(D_i = 1 \mid X_i = x\big)\big(1 - P\big(D_i = 1 \mid X_i = x\big)\big)P\big(X_i = x\big)}{\sum_x P\big(D_i = 1 \mid X_i = x\big)\big(1 - P\big(D_i = 1 \mid X_i = x\big)\big)P\big(X_i = x\big)}\right]$$

Matching vs. Regression in essential

- The **matching estimator** δ_M

$$\delta_M = \sum_x \delta_X \left[\frac{P\left(D_i = 1 \mid X_i = x\right) P\left(X_i = x\right)}{\sum_x P\left(D_i = 1 \mid X_i = x\right) P\left(X_i = x\right)} \right]$$

- The **regression estimator** δ_R

$$\delta_R = \sum_x \delta_X \left[\frac{P\left(D_i = 1 \mid X_i = x\right)\left(1 - P\left(D_i = 1 \mid X_i = x\right)\right) P\left(X_i = x\right)}{\sum_x P\left(D_i = 1 \mid X_i = x\right)\left(1 - P\left(D_i = 1 \mid X_i = x\right)\right) P\left(X_i = x\right)} \right]$$

- The difference between the two estimators is **the weight** of the treatment effect δ_X for each unique value of X_i.

  - The matching estimator δ_M uses the weight $P(D_i = 1 \mid X_i = x)$, which is larger for more treated samples.
  - The regression estimator δ_R uses the weight $P(D_i = 1 \mid X_i = x)(1 - P(D_i = 1 \mid X_i = x))$, which is the largest when $P(D_i = 1 \mid X_i = x) = 0.5$, thus half treated and half untreated observations.

Appendix

- Then the regression model of D on X is

$$D_i = \sum_x I(X_i = x)\gamma_x + v_i$$

  - The $I(X_i = x)$ is still the indicator function for $X_i = x$

  - The γ_x is the coefficient of $X_i = x$ and v_i is the error term.

- Then the population regression function (PRF) in terms of conditional expectation function (CEF) as

$$E[D|X] = \sum_x I(X_i = x)\gamma_x = \gamma$$

- Then the residuals of the regression model of D on X,$\tilde{\mathrm{D}}_i$, is

$$\tilde{D}_i = D - E[D|X_i = x]$$

- Because our regression model is saturated as follows

$$Y_i = \sum_x I(X_i = x)\beta_x + \delta_R D_i + u_i$$

- Then the key coefficient of interest is δ_R. Based on FWL theorem, we have

$$\delta_R = \frac{\text{Cov}\left(\tilde{D}_i, Y_i\right)}{V\left(\tilde{D}_i\right)}$$

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$$= \frac{E[\tilde{D}_i, Y_i]}{\tilde{D}_i^2} \quad \because \text{Cov}\left(\tilde{D}_i, Y_i\right) = E[\tilde{D}_i, Y_i] - E[\tilde{D}_i]E[Y_i]$$

$$= \frac{E\left[\left(D_i - E\left[D_i \mid X_i\right]\right)Y_i\right]}{E\left[\left(D_i - E\left[D_i \mid X_i\right]\right)^2\right]} \quad \because \tilde{D}_i = D - E[D \mid X_i = x]$$

$$= \frac{E\left\{\left(D_i - E\left[D_i \mid X_i\right]\right)E\left[Y_i \mid D_i, X_i\right]\right\}}{E\left[\left(D_i - E\left[D_i \mid X_i\right]\right)^2\right]} \quad \because ILE$$

Proof of the regression estimator δ_R

- Because

$$E\left[Y_i \mid D_i, X_i\right] = E\left[Y_i \mid D_i = 0, X_i\right] + \delta_X D_i$$

- Then the numerator of δ_R is

$$E\left\{\left(D_i - E\left[D_i \mid X_i\right]\right)E\left[Y_i \mid D_i, X_i\right]\right\}$$

$$= E\left\{\left(D_i - E\left[D_i \mid X_i\right]\right)E\left[Y_i \mid D_i = 0, X_i\right]\right\} + E\left\{\left(D_i - E\left[D_i \mid X_i\right]\right)D_i\delta_X\right\}$$

$$= 0 + E\left\{\left(D_i - E\left[D_i \mid X_i\right]\right)D_i\delta_X\right\} \because Cov(\tilde{D}_i, X) = 0$$

$$= E\left\{\left(D_i - E\left[D_i \mid X_i\right]\right)^2\delta_X\right\} \because Cov(\tilde{D}_i, D) = \tilde{D}_i^2$$

Proof of the regression estimator δ_R

$$\delta_R = \frac{E\left\{\left(D_i - E[D_i \mid X_i]\right)E\left[Y_i \mid D_i, X_i\right]\right\}}{E\left[\left(D_i - E[D_i \mid X_i]\right)^2\right]}$$

$$= \frac{E\left[\left(D_i - E[D_i \mid X_i]\right)^2 \delta_X\right]}{E\left[[\left(D_i - E[D_i \mid X_i]\right)^2 \mid X_i]\right]}$$

$$= \frac{E\left[[\left(D_i - E[D_i \mid X_i]\right)^2 \mid X_i]\delta_X\right]}{E\left[\left(D_i - E[D_i \mid X_i]\right)^2\right]}$$

$$= \frac{E\left[Var^2\left(X_i \mid D_i\right)\delta_X\right]}{E\left[Var^2\left(X_i \mid D_i\right)\right]}$$