

Quantitative Social Science in the Age of Big Data and AI

Lecture 7: Matching

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Outline

1. Review the last three lectures
2. Matching: Introduction
3. Matching: Propensity Score Matching
4. Matching in Practice
5. Wrap up

Review the Last Three Lectures

Causal Inference and OLS Regression

- Causal inference in nature is a comparison about two potential outcomes Y_1 and Y_0 for the same unit.
- The main challenge of causal inference is that we cannot observe the counterfactual outcome Y_0 for the treated group.
- Using RCTs, we can compare the treated group and the untreated group directly.
- However, in many cases, we do not have a RCT, thus we need to use some methods to make causal inference.
- The first method using observational data that we learned to make causal inference is **OLS regression**.

Causal Inference and OLS Regression

- The main identification strategy of OLS regression is **Control**, ie. putting covariates into the regression as control variables.
- The identifying assumption of OLS regression is
 - Conditional Independence Assumption(CIA): which means that if we can balance / adjust / control for the covariates X to make the treatment D as randomized, thus

$$(Y_1, Y_0) \perp\!\!\!\perp D | X$$

- Then the estimated coefficient $\hat{\beta}_1$ can be interpreted as the ATE / ATT of the treatment D on the outcome Y .
- And hypothesis testing and confidence interval can help us to see if the treatment has a significant effect on the outcome in the *population*.

OLS Regression: Nonlinearity

- Recall the OLS regression model is a linear model in terms of the parameters β_i s.

$$Y_i = \beta_0 + \beta_1 X_{1i} + \beta_2 X_{2i} + \dots + \beta_k X_{ki} + u_i$$

- Nonlinearity in Xs
 - Polynomial terms: X_{1i}^2 , $X_{1i}X_{2i}$, etc.
 - Logarithmic terms: $\ln(X_{1i})$ and $\ln(Y_i)$, etc.
 - Interaction terms: $X_{1i} \times X_{2i}$, etc.
- Nonlinearity in Ys or β_i s
 - Discrete Choice models:
 - Binary/Dichotomous Ys: $Y_i \in \{0, 1\}$
 - Count Ys: $Y_i \in \{0, 1, 2, \dots\}$
 - Multinomial Ys: $Y_i \in \{1, 2, \dots, J\}$
 - Ordinal Ys: $Y_i \in \{1, 2, \dots, J\}$
- However, suppose any function form of Y_i or β_i s could be biased. Then we need to use *non-parametric* methods to estimate the relationship between Y_i and X_i s.

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:
 - What if we can **construct a reasonable control group** by **selecting some(or all) samples in untreated group** in some way?
 - A **reasonable control group** should be **similar** to the treated group in terms of the covariates X before the treatment.
 - A similar but not equivalent question is find a suitable treat group by selecting some samples in treated group.
- 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

$$\delta_{ATE} = E[Y_{i1} - Y_{i1}^c]$$

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

- **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.
- Then the key coefficient $\hat{\beta}_1$ is the difference in log-wage between the treated group and untreated group.
- If OLS assumptions hold, then the estimated coefficient $\hat{\beta}_1$ can be interpreted as the causal effect of the training program in terms of the wage of workers.

Unmatched Samples by training status

Trainees			Non-Trainees		
unit	age	earnings	unit	age	earnings
1	28	17700	1	43	20900
2	34	10200	2	50	31000
3	29	14400	3	30	21000
4	25	20800	4	27	9300
5	29	6100	5	54	41100
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- The average wage gap between the treated group and the untreated group is

$$\delta = E[Y_{1i} - Y_{0i}] = 16426 - 20724 = -4298$$

- It appears that joining the training program will **reduce** the wages of workers by 4298.
- Question:** Can you find the bias of this estimation?

OLS Regression for the Training Program

- **Answer:** Yes, we may suffer the **OV**B. The common way to solve this problem is to add some covariates into the regression model.

$$Y_i = \beta_0 + \beta_1 D_i + \beta_2 X_i + u_i$$

- where X_i is the covariate of the workers, like age, education, experience, etc.
- 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.
- Therefore, we may use a method which can balance the covariates X like OLS but **without the assumption of function forms**:
 - **Matching**

A Training Example: matching samples

- Assume that the covariates X is the **age of the workers**, and to see **how the matching method works**.
- We pick **the untreated samples** that are **similar to the treated samples** in terms of the age of the workers.

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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			
16	43	10700	16	24	9700			
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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			

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13	29	12500	13	26	16500	17	29	6200
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15	25	10100	15	25	23300	15	25	23300
16	43	10700	16	24	9700	1	43	20900
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13	29	12500	13	26	16500	17	29	6200
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An Illustrated Example: matched samples

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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

$$\delta = E[Y_{1i} - Y_{0i}] = 16426 - 13982 = 2444$$

- Now, joining the training program will **increase** the wage of workers by 2444 .

Two Assumptions: One Old and One New

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

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

- More specifically, we assume that the potential incomes for the workers are **independent of the 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**?
 - It means that these samples are *not used* in the estimation of the average treatment effect at all.
 - This is due to the **Overlap Assumption**, a new assumption in the matching method that was not discussed in the OLS regression.

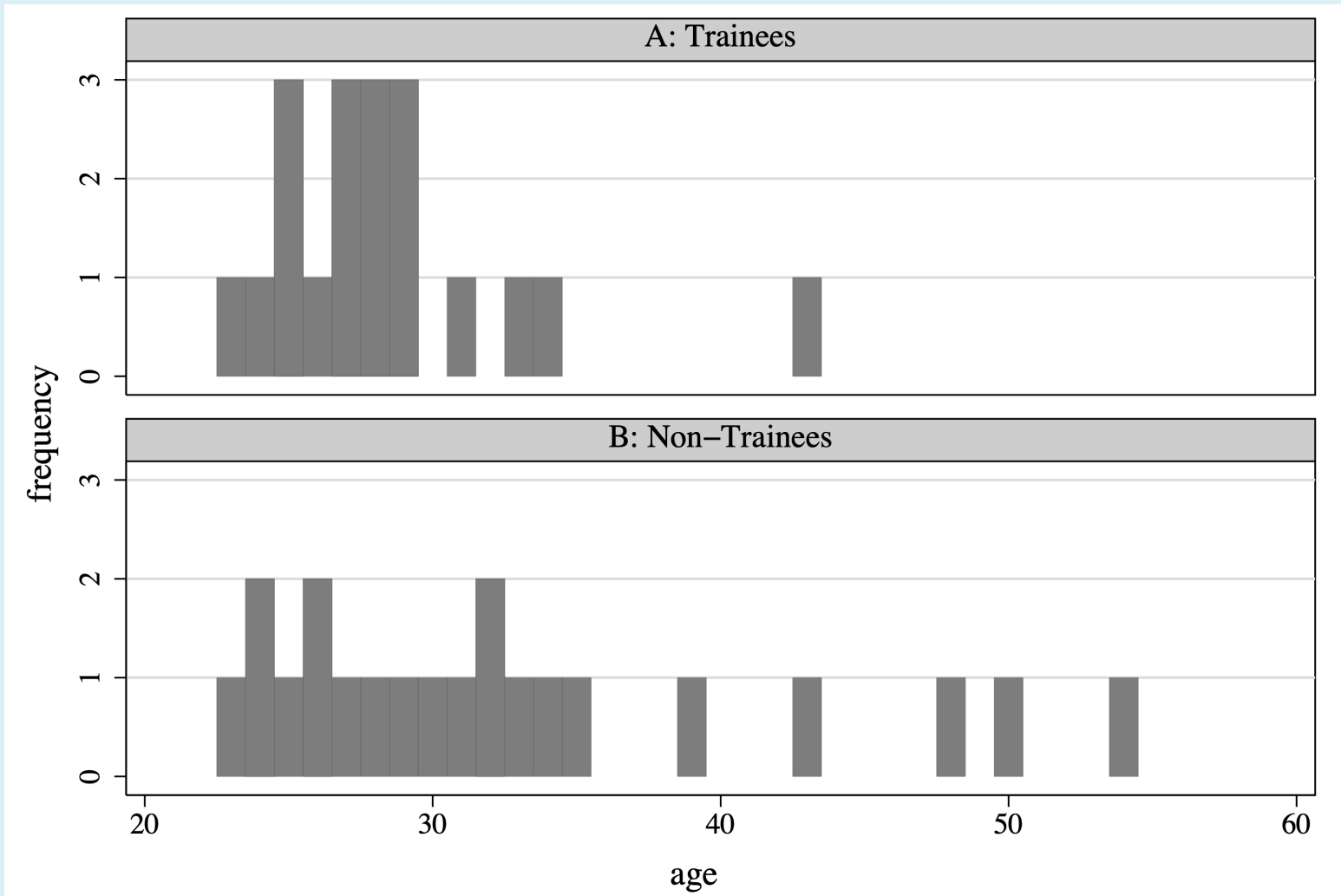
Two Assumptions: One Old and One New

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

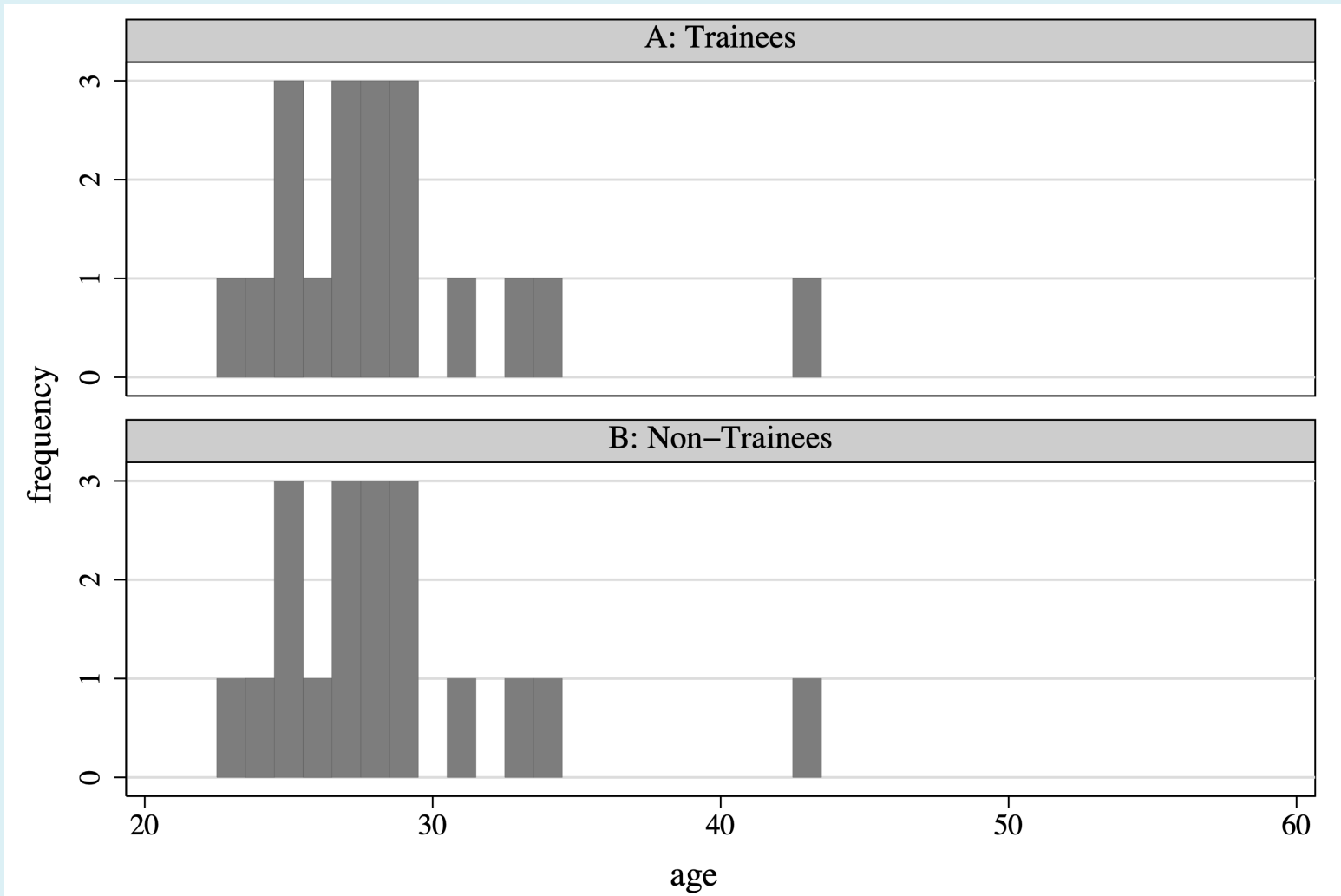
$$0 < \Pr(D_i = 1 \mid X_i) < 1$$

- This implies that the likelihood of receiving treatment is neither 0 nor 1 for any given covariates.
 - If the probability of receiving treatment is 0 for some X_i , then no samples with these characteristics receive treatment, making them unavailable for matching.
 - If the probability of receiving treatment is 1 for some X_i , then all samples with these characteristics receive treatment, making it impossible to find untreated matches.
 - 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.

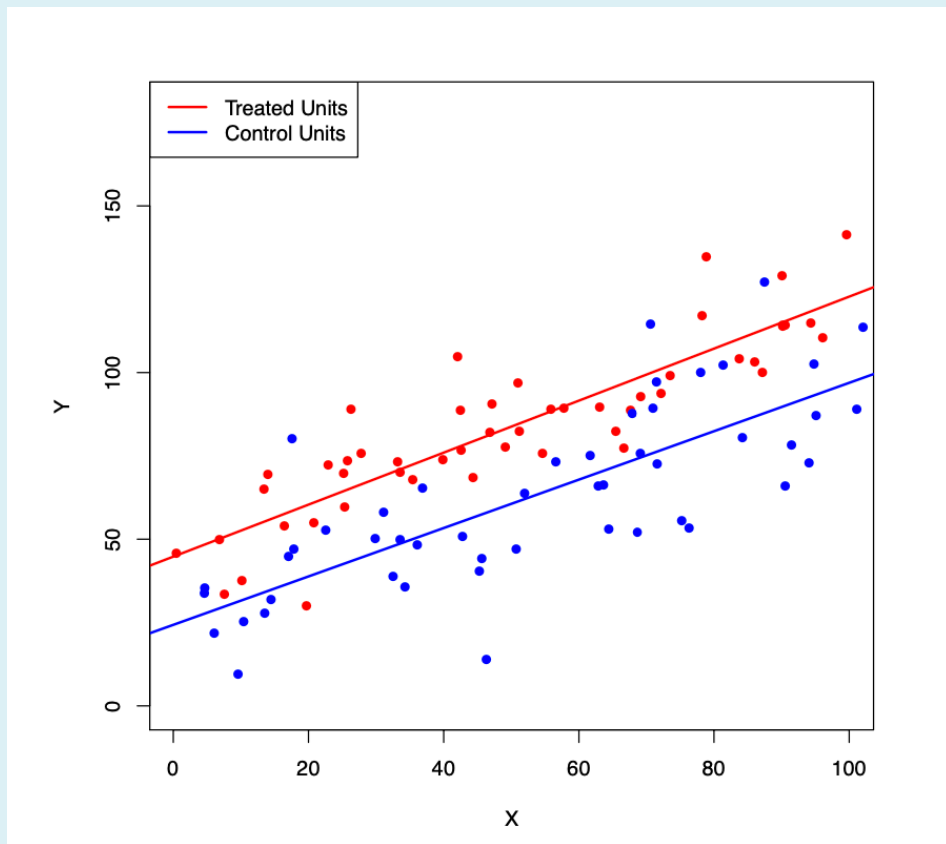
A Training Example: before matching



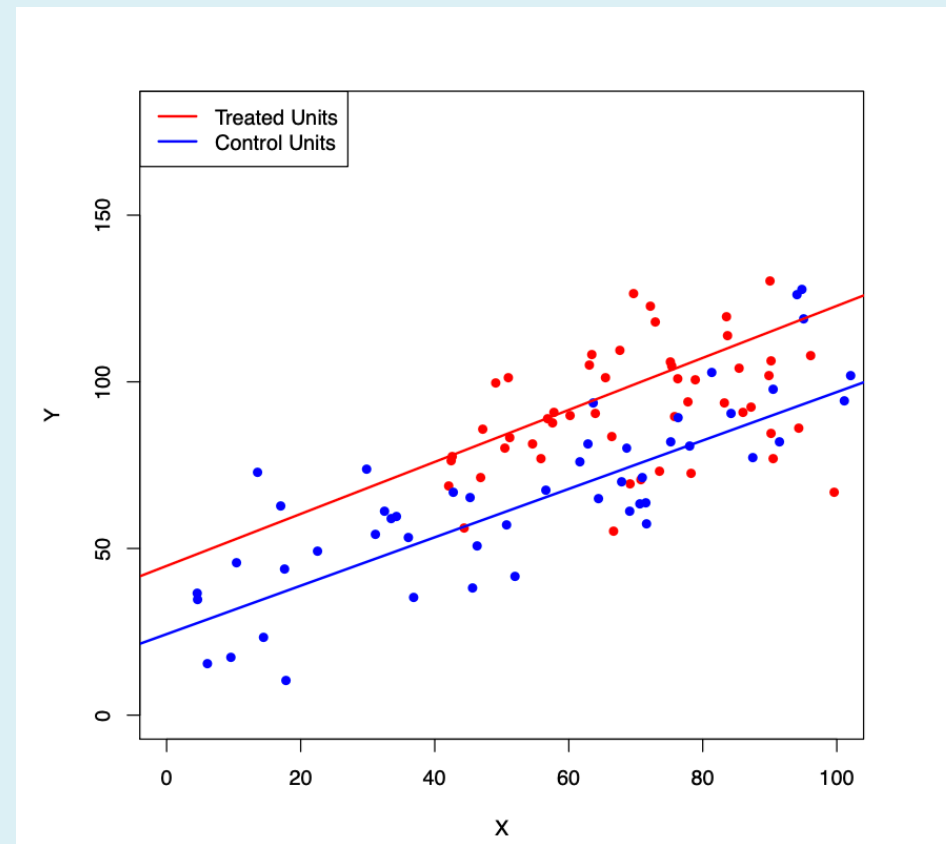
A Training Example: after matching



The Overlap Assumption in OLS



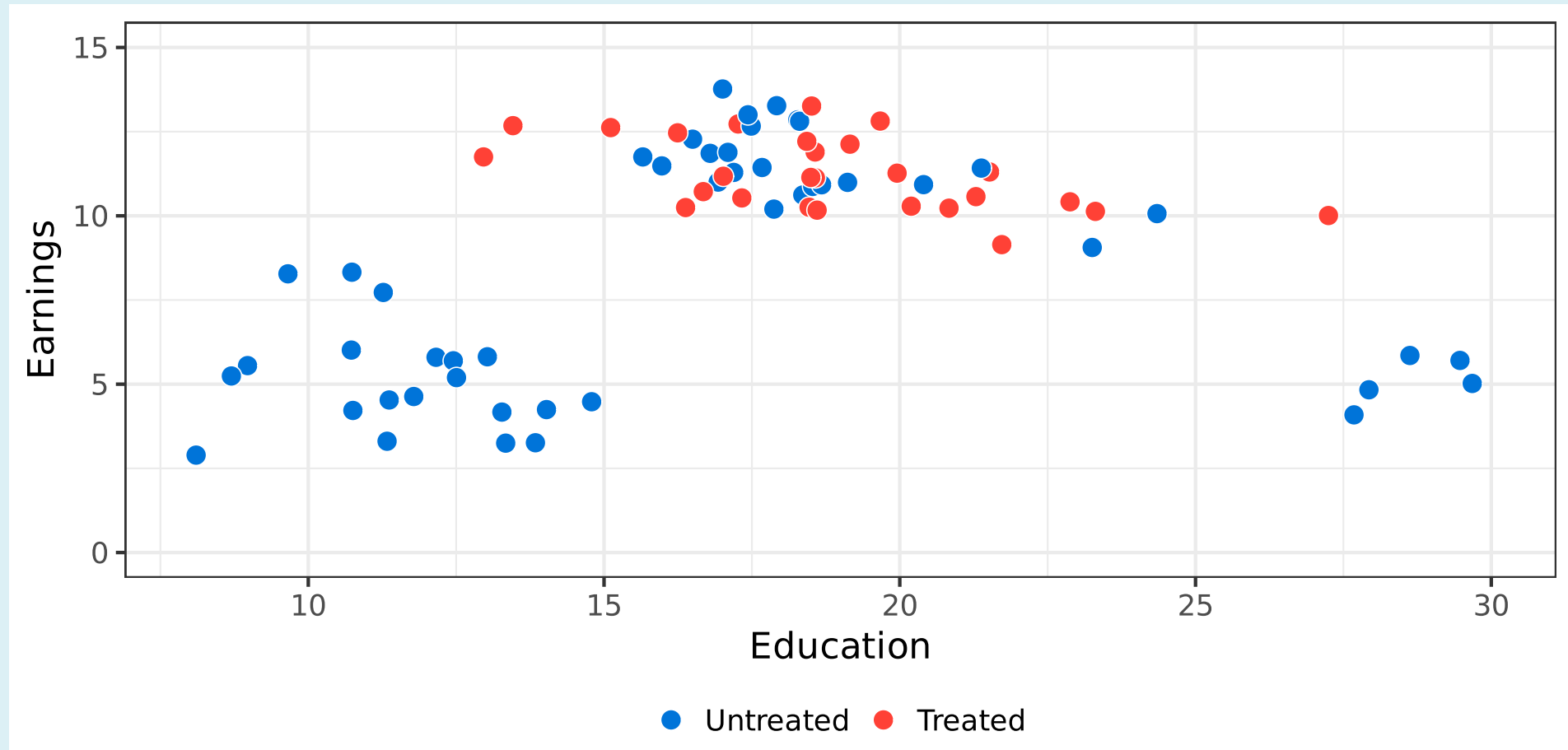
The overlap assumption is satisfied



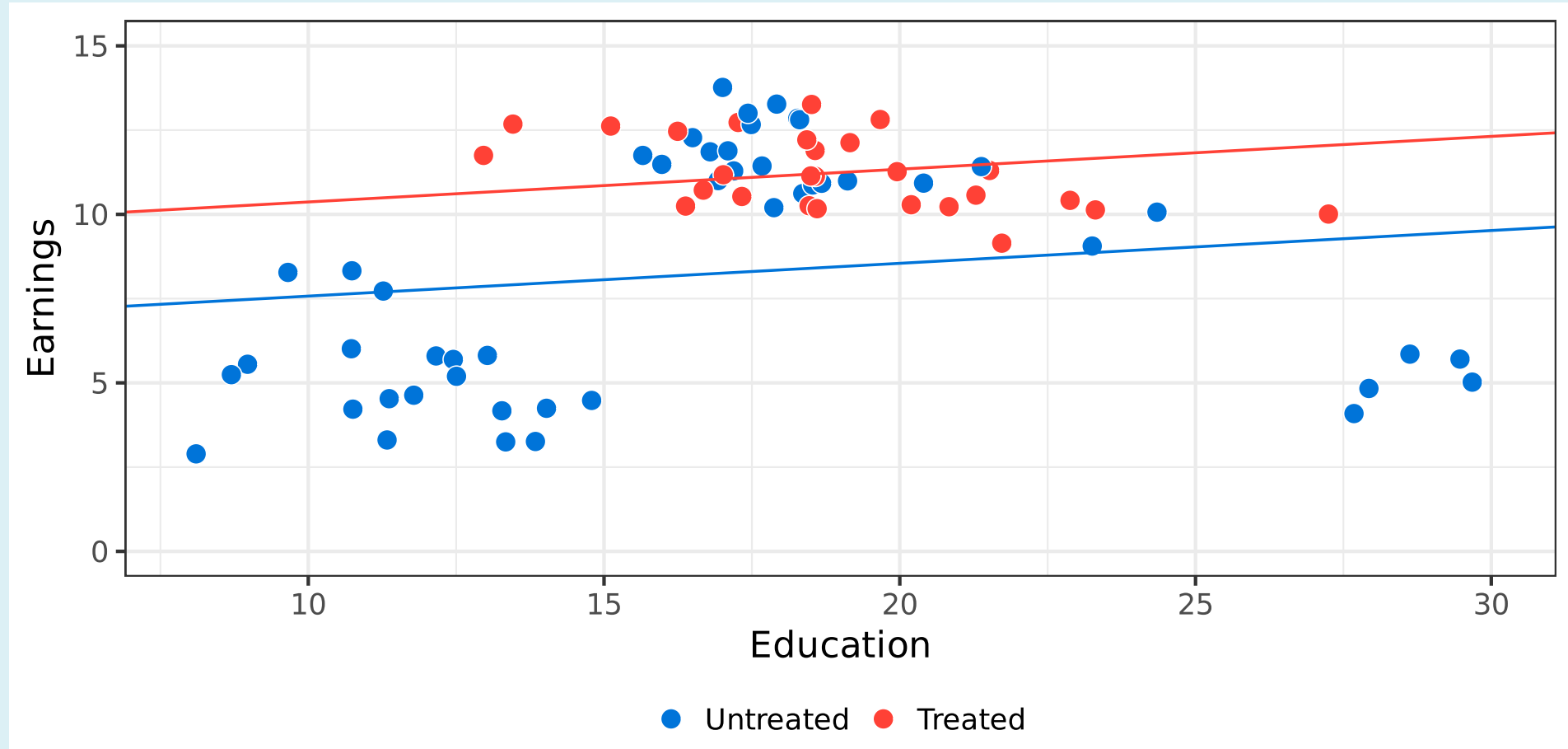
The overlap assumption is violated

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

OLS vs Matching: Overlap Assumption



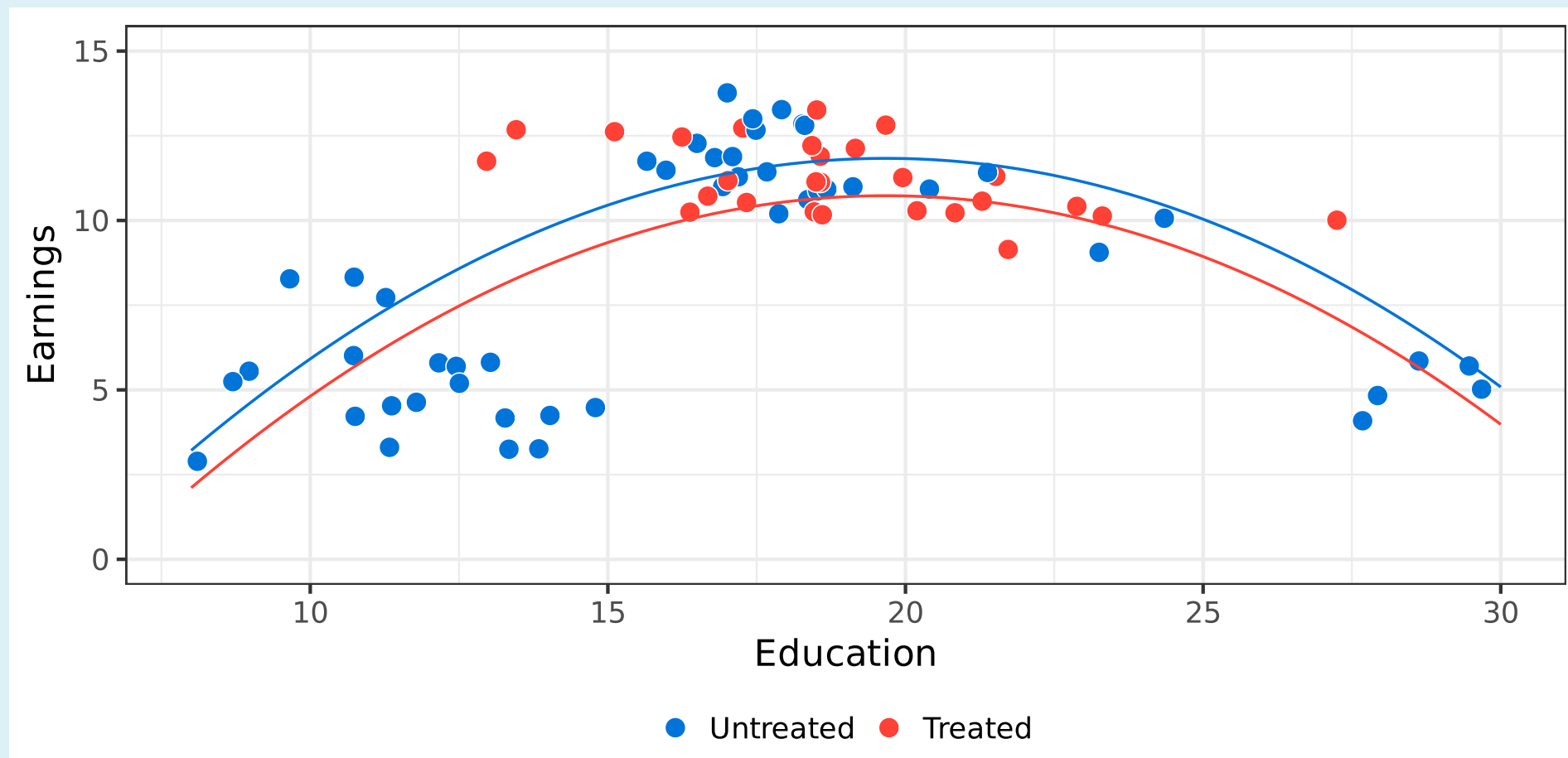
OLS vs Matching: Overlap Assumption



$$\text{Earnings} = \beta_0 + \beta_1 \text{Education} + \beta_2 \text{Treatment} + u$$

- Basic take-away: *treated* group is **higher** than the *untreated* group in the OLS regression.

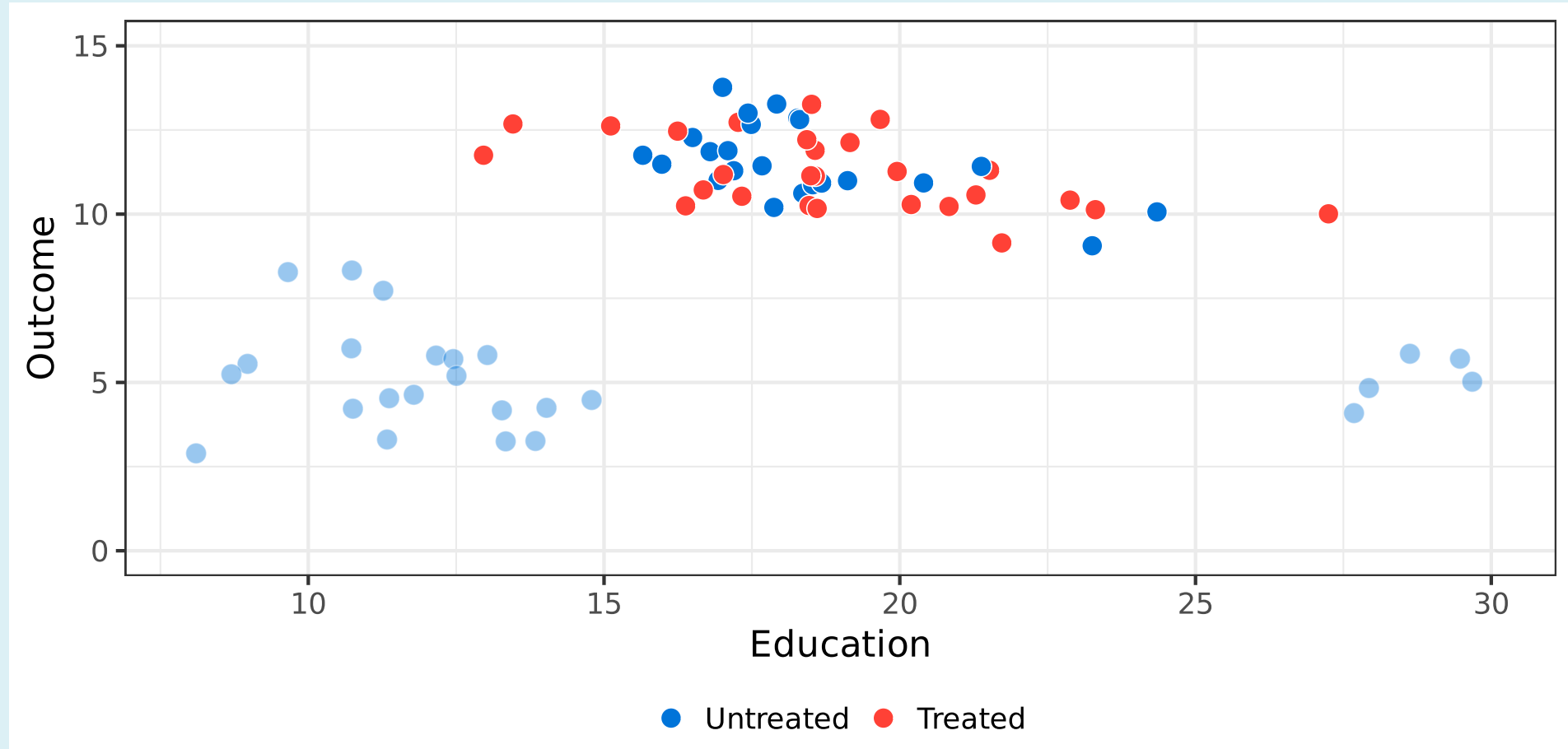
Overlap Assumption in Nonlinearity



$$\text{Earnings} = \beta_0 + \beta_1 \text{Education} + \beta_2 \text{Education}^2 + \beta_3 \text{Treatment} + u$$

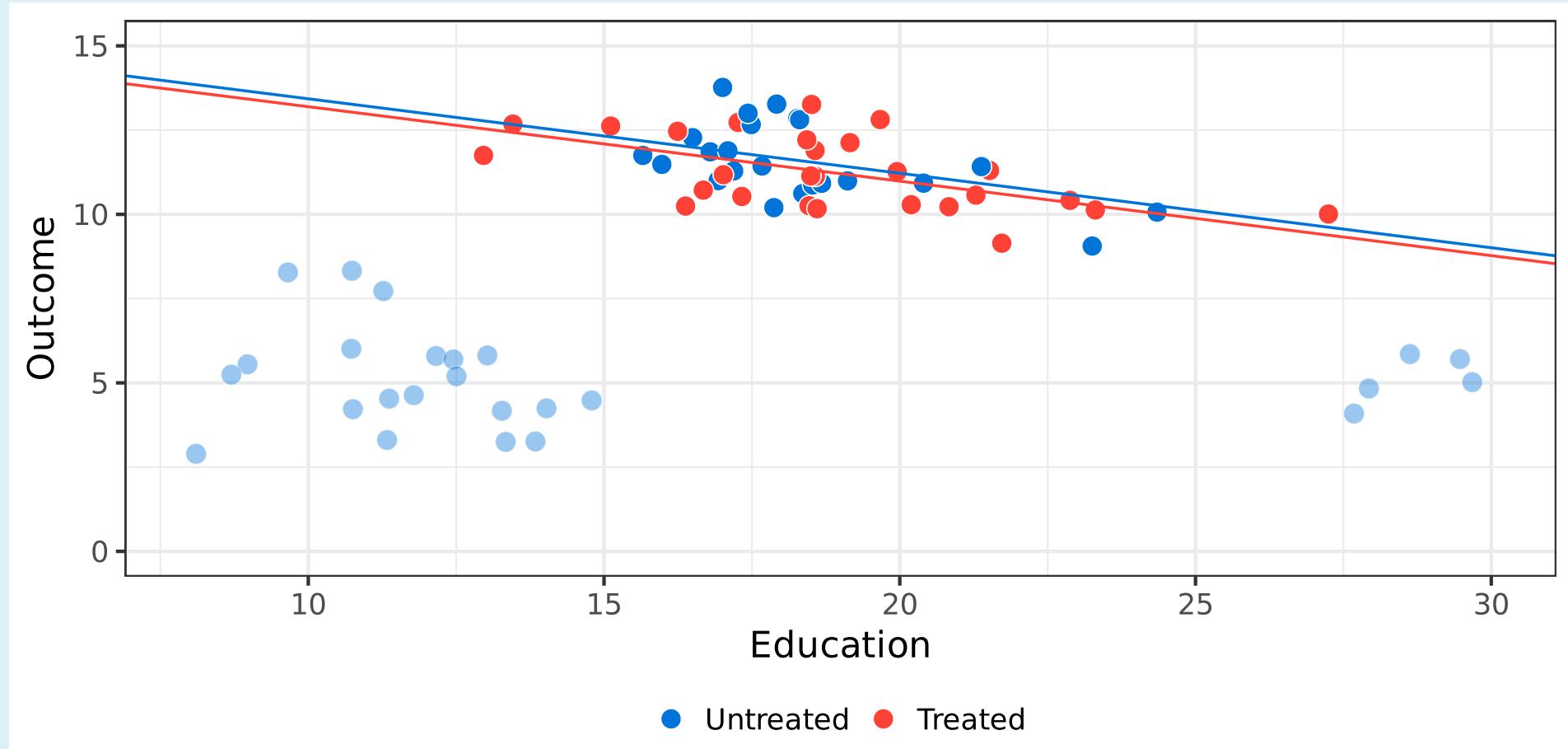
- Basic take-away: *untreated* group is **higher** than the *treated* group in the OLS regression.

Overlap Assumption: Common Support



- The **common support region** is the region where *both the treated and untreated groups have data*.

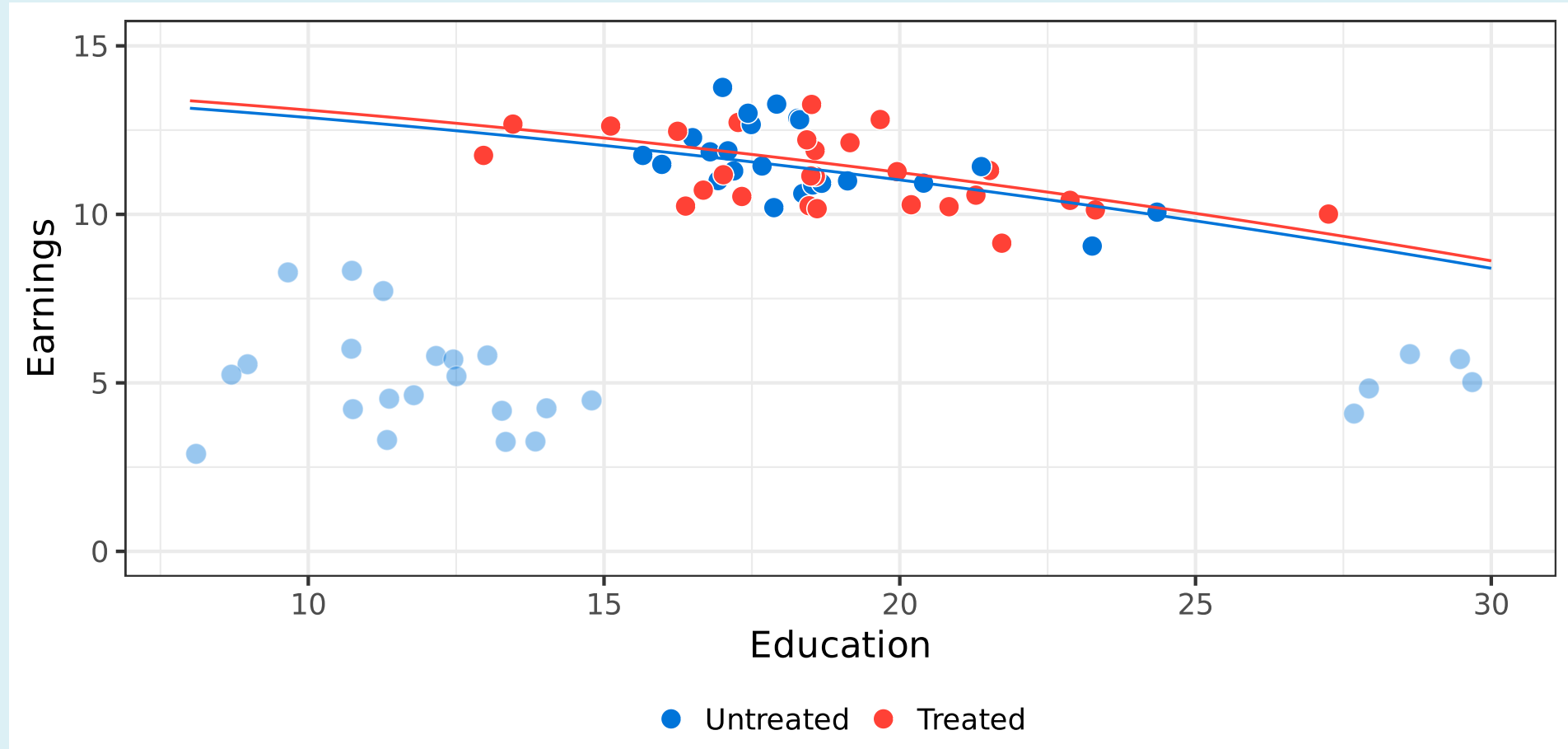
Common Support: After Trimming Data



$$\text{Earnings} = \beta_0 + \beta_1 \text{Education} + \beta_2 \text{Treatment} + u$$

- Basic take-away: *treated* group is **very close** to the *untreated* group in the OLS regression.

Common Support: After Trimming Data



$$\text{Earnings} = \beta_0 + \beta_1 \text{Education} + \beta_2 \text{Education}^2 + \beta_3 \text{Treatment}$$

- Basic take-away: *treated* group is still **very close** to the *untreated* group in the OLS regression.

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 $X = (X_1, X_2, \dots, X_k)'$?
 - In this case, it is **impossible** to find proper units with identical values in all covariates X_1, X_2, \dots, X_k .
- Two complementary solutions are running in parallel, representing the directions in which the matching method is developing.

1. Lower the accuracy of the comparison.

- From *find a unit in the untreated group with the same covariate values* to *find a unit in the untreated group with similar covariate values*.

2. Directly reduce dimensions.

- Converting multiple variables into *a single numerical value*, then use the numerical value to match the samples.

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.
 1. define how to measure the similarity between the treated and untreated samples.
 2. choose the criteria to match the samples.
 3. evaluate the quality of the matching.
- 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.
- We will focus on the **Matching** and **Estimation** in this lecture.

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

$$Y_{1i}^C = \sum_j w_i(j) Y_{0j}$$

- 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

$$\delta_i = Y_{1i} - Y_{1i}^C = Y_{1i} - \sum_j w_i(j) Y_{0j}$$

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

$$\hat{\delta}_M = \frac{1}{N_T} \sum_{i \in (D=1 \cap C)} \delta_i = \frac{1}{N_T} \sum_{i \in (D=1 \cap C)} (Y_{1i} - Y_{1i}^c) = \frac{1}{N_T} \sum_{i \in (D=1)} \left(Y_{1i} - \sum_{j \in (D=0 \cap C)} w_i(j) Y_{0j} \right)$$

- Where C is the common support region of the treated and untreated individuals.
- And $j = 1, 2, \dots, N^C$ and $i = 1, 2, \dots, N^T$.
- $i \in (D = 1 \cap C)$ 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.

$$\begin{aligned}\hat{\delta}_M &= \frac{1}{N_T} \sum_{i \in (D=1 \cap C)} \left(Y_{1i} - \sum_{j \in (D=0 \cap C)} w_i(j) Y_{0j} \right) \\ &= \frac{1}{N_T} \sum_{i \in (D=1 \cap C) \& j=i} (Y_{1i} - Y_{0j})\end{aligned}$$

- 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.
 - The idea: **more similar the covariates are, more weight the untreated sample should have.**

Proximity: When X is Discrete

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

$$w_i(j) = \mathbb{I}(X_i = X_j)$$

- Where $\mathbb{I}(\cdot)$ is an indicator function,

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

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

$$|X_i - X_j|$$

- 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 the **distance metric**.
- If we just pick **the smallest one** as we did in the training 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

$$X_i = (X_{i1}, X_{i2}, \dots, X_{ik}) \quad \text{and} \quad X_j = (X_{j1}, X_{j2}, \dots, X_{jk})$$

- **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 ,

$$\|X_i - X_j\| = \sqrt{(X_i - X_j)'(X_i - X_j)}$$

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**

$$\|(X_i - X_j)\| = \sqrt{(X_i - X_j)' V_X^{-1} (X_i - X_j)}$$

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

$$V_X^{-1} = \begin{bmatrix} \hat{\sigma}_1^2 & 0 & 0 & \cdots & 0 \\ 0 & \hat{\sigma}_2^2 & 0 & \cdots & 0 \\ 0 & 0 & \hat{\sigma}_3^2 & \cdots & 0 \\ \vdots & \vdots & \vdots & \ddots & \vdots \\ 0 & 0 & 0 & \cdots & \hat{\sigma}_k^2 \end{bmatrix}$$

- $\hat{\sigma}_k^2$ is the variance of the k -th variable.
- No scale problem but still *no correlations* between X s.

Proximity: When X is a Vector

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

$$\|(X_i - X_j)\| = \sqrt{(X_i - X_j)' \Sigma_X^{-1} (X_i - X_j)}$$

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

$$\Sigma_X^{-1} = \begin{bmatrix} \hat{\sigma}_{11} & \hat{\sigma}_{12} & \hat{\sigma}_{13} & \cdots & \hat{\sigma}_{1k} \\ \hat{\sigma}_{11} & \hat{\sigma}_{22} & \hat{\sigma}_{23} & \cdots & \hat{\sigma}_{2k} \\ \hat{\sigma}_{11} & \hat{\sigma}_{32} & \hat{\sigma}_{33} & \cdots & \hat{\sigma}_{3k} \\ \vdots & \vdots & \vdots & \ddots & \vdots \\ \hat{\sigma}_{k1} & \hat{\sigma}_{k2} & \hat{\sigma}_{k3} & \cdots & \hat{\sigma}_{kk} \end{bmatrix}$$

- $\hat{\sigma}_{jk}$ 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:**
 - the weight $w_i(j)$ is 1 if j is the exact match of i and 0 otherwise.
 - **problem:** The exact match may not be found.
- **Nearest Neighbor Matching(NNM):**
 - The smallest distance between the treated and untreated group.
 - the weight $w_i(j)$ is 1 if j is the nearest neighbor of i and 0 otherwise.
 - **problem:** The nearest neighbor may not be a good match.
- **Caliper Matching:** Samples within a certain range are matched.
 - the weight $w_i(j)$ is 1 if j is in the range and 0 otherwise.
 - **problem:** How to choose the range?

Many Matching Methods

- **Radius Matching**: all the samples within a certain range are matched.
 - the weight $w_i(j)$ is 1 if j is in the range and 0 otherwise.
 - **problem**: How to choose the **radius**?
- **Subclassification** : Divide the treated and untreated group into subclasses based on the covariates and then match within each subclass.
 - the weight $w_i(j)$ is 1 if j is in the same subclass as i and 0 otherwise.
 - **problem**: How to choose the subclasses?
- **Kernel Matching**: The weight is based on the kernel function, which is an estimated density function of the covariates.
 - all the samples in the untreated group are used to estimate the counterfactual outcome.
 - the weight is based on the specific kernel function

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 .
- **Propensity scores**
- 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:

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

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

$$p(X_i) = E[D_i | X_i]$$

- Formally the **Propensity Score Theorem** is

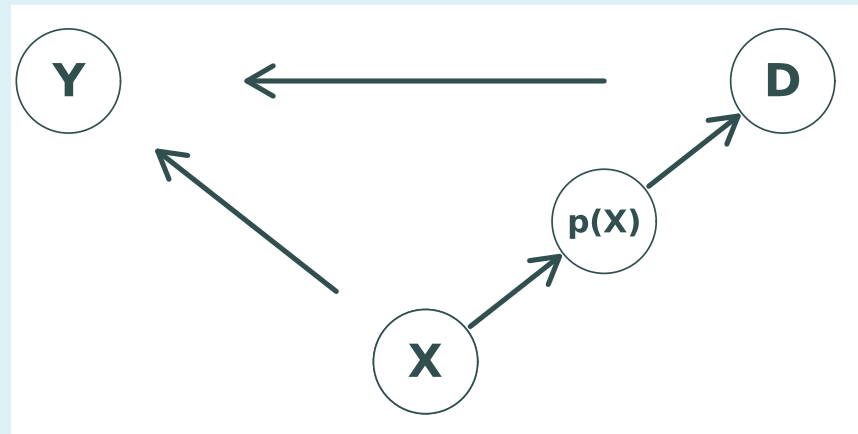
- If $(Y_{0i}, Y_{1i}) \perp\!\!\!\perp D_i | X_i$, then $(Y_{0i}, Y_{1i}) \perp\!\!\!\perp D_i | p(X_i)$.

- 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 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 $D_i | \text{a function of } X \text{ is independent of } (Y_{0i}, Y_{1i})$.



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?**

$$p(X_i) = \Pr(D_i = 1 \mid X_i)$$

- Yes, it is the **binary dependent regression** model that the independent variables are the covariates X_i .
- We can estimate the propensity scores using three models we covered previously:
 1. LPM
 2. Logit
 3. Probit
- 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

$$p(\mathbf{X}_i) = E(D = 1 | \mathbf{X}_i) = \frac{1}{1 + e^{(-\mathbf{X}_i\beta)}}$$

- Where \mathbf{X}_i is the vector of covariates and β is the vector of coefficients.
- The **estimated propensity scores** $\hat{p}(\mathbf{X}_i)$ can be obtained by plugging in the estimated coefficients $\hat{\beta}$.

$$\hat{p}(\mathbf{X}_i) = \frac{1}{1 + e^{(-\mathbf{X}_i\hat{\beta})}}$$

Propensity-Score Matching

Estimation: Logit Regression

- However, for the *nonlinearity* of the model, the marginal effect of covariates 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**,

$$\ln\left(\frac{p(\mathbf{X}_i)}{1 - p(\mathbf{X}_i)}\right) = \mathbf{X}_i\beta$$

- 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

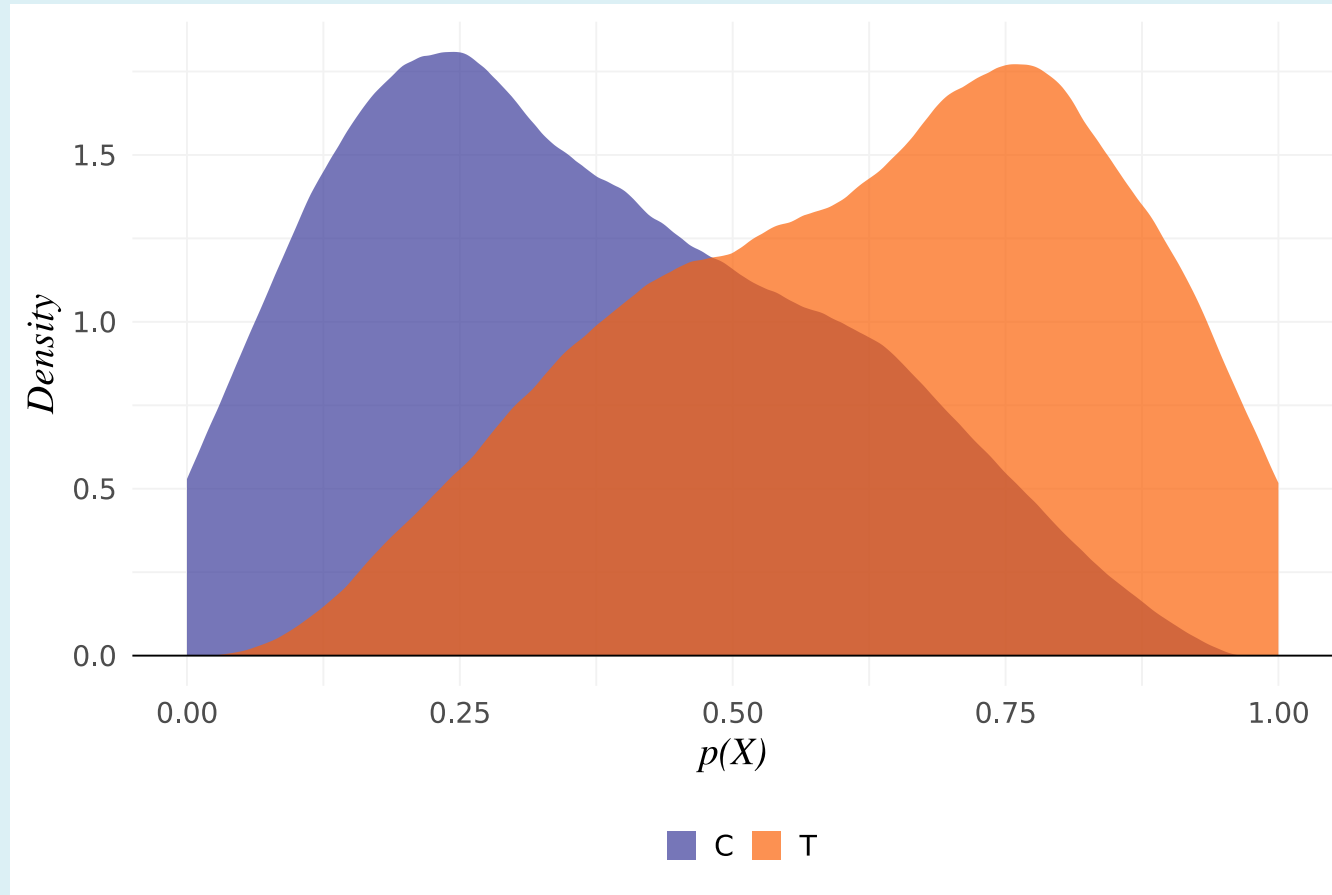
Estimation: Predicted instead of Explained

- 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.

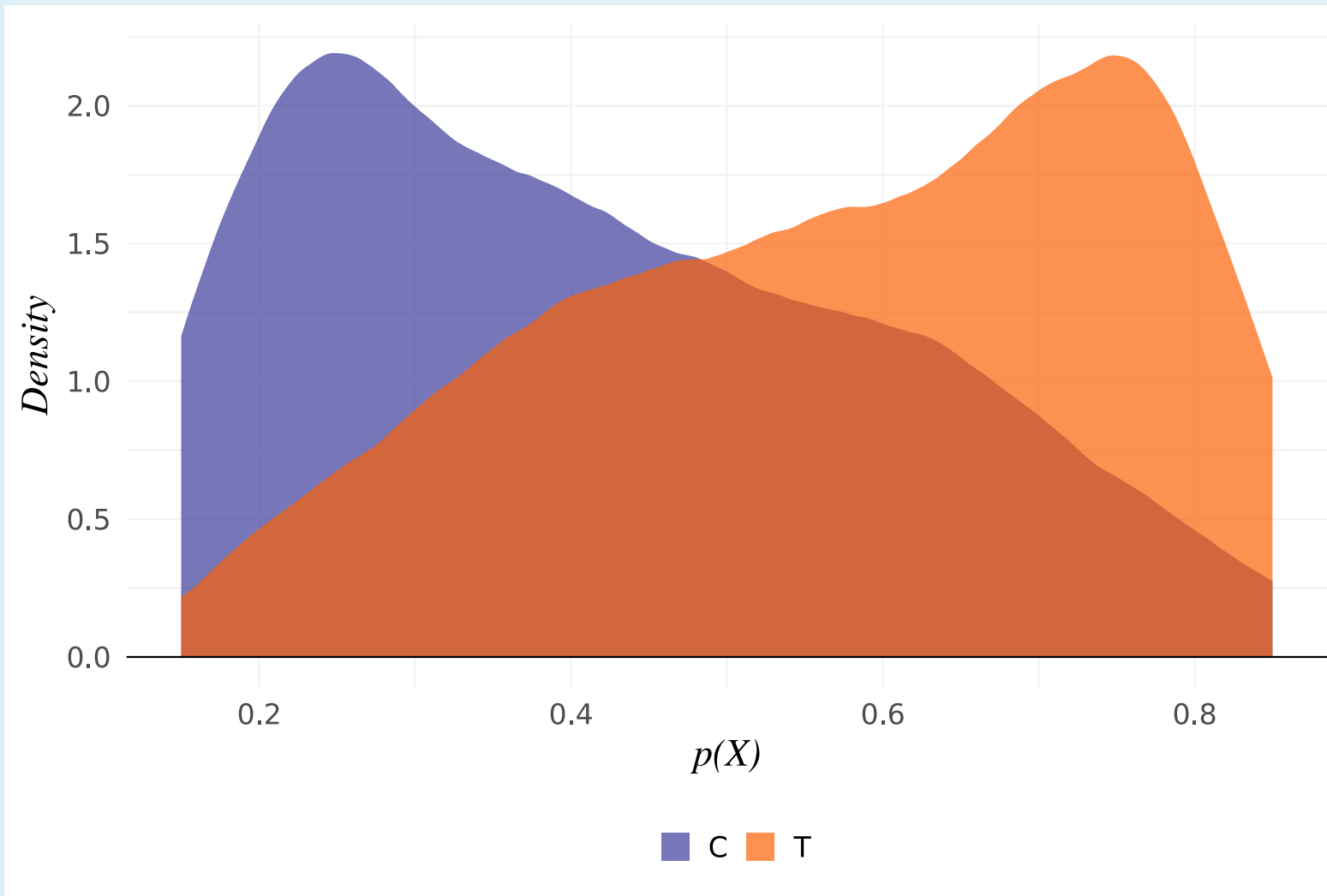
Propensity-Score Matching

Overlap Assumption in Propensity Score Methods

- Recall: The **Overlap Assumption** $0 < \Pr(D_i = 1 \mid X_i) < 1$
- 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.
- It is much more easier when using p-scores to check the overlap assumption.



- Example: The plot below shows the distribution of the estimated propensity score $p(X_i)$ for the treated and control groups.
 - The pscore in the treated group is in $[0.15, 1]$
 - The pscore in the control group is in $[0, 0.85]$



- 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 Propensity 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

$$Y_i = \beta_0 + \beta_1 D_i + \beta_2 p(X_i) + u_i$$

- 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.

$$Y_i = \beta_0 + \beta_1 D_i + \beta_2 p(X_i) + \beta_3 D_i \cdot p(X_i) + \beta_4 p(X_i)^2 + \beta_5 D_i \cdot p(X_i)^2 + \beta_6 p(X_i)^3 + \beta_7 D_i \cdot p(X_i)^3 + \dots + u_i$$

- 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:

$$\delta_{ATE} = E(Y_{1i} - Y_{0i}) = E \left[\frac{D_i}{p(X_i)} Y_i \right] - E \left[\frac{1 - D_i}{1 - p(X_i)} Y_i \right]$$

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

$$E[Y_{1i}] = E \left[\frac{D_i}{p(X_i)} Y_i \right]$$
$$E[Y_{0i}] = E \left[\frac{1 - D_i}{1 - p(X_i)} Y_i \right]$$

IPW Estimator for ATE

- Then, we could get the ATE by the following formula:

$$\delta_{ATE} = E \left[\frac{D_i}{p(X_i)} Y_i \right] - E \left[\frac{1-D_i}{1-p(X_i)} Y_i \right]$$

$$= E \left[\frac{(D_i - D_i \cdot p(\mathbf{X}_i) - p(\mathbf{X}_i) + D_i \cdot p(\mathbf{X}_i))}{p(\mathbf{X}_i)(1-p(\mathbf{X}_i))} Y_i \right]$$

$$= E \left[\frac{(D_i - p(\mathbf{X}_i))}{p(\mathbf{X}_i)(1-p(\mathbf{X}_i))} Y_i \right]$$

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

IPW Estimator for ATE

- Then the IPW estimator for ATE is given by

$$\hat{\delta}_{ATE}^{HW} = \frac{1}{N} \sum_{i=1}^N \frac{D_i - p(X_i)}{p(X_i)(1 - p(X_i))} Y_i$$

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

$$\frac{D_i - \hat{p}(\mathbf{X}_i)}{\hat{p}(\mathbf{X}_i)(1 - \hat{p}(\mathbf{X}_i))} = \begin{cases} \frac{1}{\hat{p}(\mathbf{X}_i)} & \text{if } D_i = 1 \\ -\frac{1}{1 - \hat{p}(\mathbf{X}_i)} & \text{if } D_i = 0 \end{cases}$$

- 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

$$\delta_{ATE} = E(Y_{1i} - Y_{0i}) = \frac{E \left[\frac{D_i}{p(X_i)} Y_i \right]}{E \left[\frac{D_i}{p(X_i)} \right]} - \frac{E \left[\frac{1-D_i}{1-p(X_i)} Y_i \right]}{E \left[\frac{1-D_i}{1-p(X_i)} \right]}$$

- Where $E\left(\frac{D_i}{p(X_i)}\right)$ can be seen as the **average weight** for the treated group and $E\left(\frac{1-D_i}{1-p(X_i)}\right)$ can be seen as the **average weight** for the control group.
- Then corresponding IPW estimator for ATE is given by

$$\hat{\delta}_{ATE}^{IPW} = \left[\sum_{i=1}^N \frac{Y_i D_i}{p(X_i)} \right] / \left[\sum_{i=1}^N \frac{D_i}{p(X_i)} \right] - \left[\sum_{i=1}^N \frac{Y_i (1 - D_i)}{(1 - p(X_i))} \right] / \left[\sum_{i=1}^N \frac{(1 - D_i)}{(1 - p(X_i))} \right]$$

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: Combine IPW with outcome regression
 - **Double Robustness method**: 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.
- Shortcomings of matching with replacement:
 - reduces the effective sample size when the same control units are used multiple times.
 - may lead to estimates being overly influenced by a small number of frequently used control units.

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:** Matching is over regression in the following aspects:
 1. Due to its non-parametric characteristics, matching does not impose any restrictions on empirical specification or estimate specific parameters of the CEF function.
 2. Regression does not account for the common support issue explicitly, while matching does.
- In practice, using matching alone as main identification strategy is less common in economics, more frequently combined with other methods like DID and SCM.