Introductory Econometrics Lecture 26: Treatment Effect Model

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Introduction

- \blacktriangleright In this class, we consider the problem of estimating the causal effect of a binary explanatory variable, which is referred as the treatment effect in the literature. The treatment effect model is different from the linear regression model.
- \blacktriangleright In econometrics, the treatment effect model is very often used for evaluating social program/experiment.
- \blacktriangleright Example 1: Suppose that a selected set of individuals receive training or education initiated by the government with a view to enhancing their employment prospects. Suppose that the government has collected the earnings data for the individuals who received the training and for the individuals who did not. The main purpose of methods of program evaluations is to quantify and estimate the effect of the training program.
- \blacktriangleright Example 2: Suppose that an education program required high schools to agree to assign teachers and students to small (13 to 17 students) or large (22 to 26 students) classes. The government is interested in the effect of class size on student achievement.
- \triangleright Such a question can arise in various other situations. A medical experiment studies on the effects of new treatment ask similar questions. One group of patients has received new treatment, and the other group has not.

Potential outcome variables

- ▶ Y_i : outcome variable; $D_i \in \{0, 1\}$: the binary explanatory variable; $X_{i1}, ..., X_{ik}$: other observed explanatory variables; ϵ_i : unobserved explanatory factors.
- \blacktriangleright The variable D_i is a binary variable taking 1 if the individual has gone through the treatment and 0 otherwise. The treatment here represents the actual treatment. The econometrician usually observes the treatment status for each individual D_i .
- \blacktriangleright $(X_{i1},..., X_{ik})$ represents a vector of various demographic characteristics for individual i . E.g., the variables can be annual income, age, gender, status of marriage, the number of children, education, etc. These represent all the observable characteristics of individual i .
- ► Suppose that Y_i is generated by $Y_i = g(D_i, X_{i1}, ..., X_{ik}, \epsilon_i)$.
- \blacktriangleright g is unknown and in the treatment effect model, we do not assume g is linear.
- \blacktriangleright The outcome variable $Y_i(1) = g(1, X_{i1}, \ldots, X_{ik}, \epsilon_i)$ represents a potential outcome of an individual i in the treatment state (e.g. training is received or studying in a reduced-size class). The variable $Y_i(0) = g(0, X_{i1},..., X_{ik}, \epsilon_i)$ represents a potential outcome of the same individual i in the control state (e.g. training is received or studying in a normal-size class).
- In Thus, each individual has a random vector $(Y_i(1), Y_i(0))$ that represents potential outcomes depending on the state (treatment or control). Certainly, $(Y_i(1), Y_i(0))$ are correlated.
- \blacktriangleright The econometrician cannot observe the random vector $(Y_i(1), Y_i(0))$ jointly, because for each individual, only one potential outcome $(Y_i(1)$ or $Y_i(0)$ is realized, depending on whether the individual i has gone through the treatment or not.

The relationship between D_i and $(Y_i(1), Y_i(0))$

- \blacktriangleright In a medical experiment, the individual is chosen to be in the treatment group through some randomization device or a lottery. In these cases, $D_i \perp (Y_i(1), Y_i(0))$ (i.e., D_i is independent of $(Y_i(1), Y_i(0))$.
- \triangleright For evaluating social experiment/program with observational data, it may not be convincing to assume $D_i \perp (Y_i(1), Y_i(0)).$

Treatment effects

 \blacktriangleright The individual treatment effect (ITE) for each individual *i* is defined as:

$$
Y_{i}\left(1\right) -Y_{i}\left(0\right) .
$$

- \triangleright The ITE is the difference between the potential outcomes in two different states for the same person.
- \triangleright The ITE is a counterfactual quantity, in the sense that in the actual world, we cannot observe the vector $(Y_i(1), Y_i(0))$.
- \blacktriangleright There are mainly two quantities of interest: ATE (average treatment effect)

 $ATE = E[Y_i(1) - Y_i(0)],$

and ATT (average treatment effect on the treated)

$$
ATT = E[Y_i(1) - Y_i(0) | D_i = 1].
$$

 \blacktriangleright The average treatment effect on the treated is the treatment effect of the people who have gone through the treatment.

- \triangleright Note that the expectation in the definition of ATE involves the joint distribution of $(Y_i(1), Y_i(0))$, and the expectation in the definition of ATT involves the joint distribution of $(Y_i(1), Y_i(0), D_i)$, which are both unobserved.
- \blacktriangleright ATE or ATT can not be estimated accurately merely by collecting a large size of samples.

The observed information

 \blacktriangleright The econometrician observes the treatment status D_i and covariates X_i . She also observes the outcome variable:

$$
Y_i = D_i Y_i (1) + (1 - D_i) Y_i (0).
$$

- \blacktriangleright The observed outcome variable Y_i is not the same as the potential outcomes $Y_i(1)$ or $Y_i(0)$. It is a realized outcome for an individual *i* depending on whether she has received treatment (Y_i) is realized to be $Y_i(1)$ or not $(Y_i$ is realized to be $Y_i(0)$).
- \blacktriangleright Identification of these parameters is concerned with the following question: can we uniquely determine the value of these parameters once we know the joint distribution of the observable random variables?

Randomized experiments

- \blacktriangleright In medical experiments, the treatment is performed using a randomization device. More specifically, for patient i , a lottery is run, and the patient is selected into the treated group with the design probability p , and stays in the control group with the design probability $1-p$.
- In these cases, we have $D_i \perp (Y_i(1), Y_i(0), X_{i1}, \ldots, X_{ik}).$ Randomized experiment assumption requires that knowing whether patient i is treated or not gives one no informational advantage in predicting the potential outcomes of i over another who does not know whether patient i is treated or not.
- \triangleright This assumption is still possibly violated in medical studies if only those patients who have higher potential treatment effect are selected into treatment among all the patients in the study on purpose.
- In this case, observing D_i will give information about the treatment effect $(Y_i(1) - Y_i(0))$ for individual *i*.

 \triangleright We use the following result from probability theory: if $V \perp W$, then for any function f ,

$$
E[f(V, W) | W = w] = E[f(V, w)].
$$
 (1)

 \blacktriangleright By [\(1\)](#page-10-0) and the randomized experiment assumption, $D_i \perp (Y_i(1), Y_i(0))$, we have

$$
\begin{aligned}\n\text{ATE} &= \mathbb{E}\left[Y_i\left(1\right) - Y_i\left(0\right)\right] \\
&= \mathbb{E}\left[Y_i\left(1\right)\right] - \mathbb{E}\left[Y_i\left(0\right)\right] \\
&= \mathbb{E}\left[D_i Y_i\left(1\right) + \left(1 - D_i\right) Y_i\left(0\right) \mid D_i = 1\right] \\
&- \mathbb{E}\left[D_i Y_i\left(1\right) + \left(1 - D_i\right) Y_i\left(0\right) \mid D_i = 0\right] \\
&= \mathbb{E}\left[Y_i \mid D_i = 1\right] - \mathbb{E}\left[Y_i \mid D_i = 0\right].\n\end{aligned}
$$

\blacktriangleright By LIE,

$$
E[Y_i D_i] = E[E[Y_i D_i | D_i]]
$$

= Pr [D_i = 1] E[Y_i D_i | D_i = 1]
+ Pr [D_i = 0] E[Y_i D_i | D_i = 0]
= E [D_i] E[Y_i | D_i = 1],

where

$$
E[Y_i D_i | D_i = 0] = E[(D_i Y_i(1) + (1 - D_i) Y_i(0)) D_i | D_i = 0]
$$

= 0

follows from [\(1\)](#page-10-0).

 \blacktriangleright Similarly, we have

$$
E[Y_i | D_i = 0] = \frac{E[Y_i (1 - D_i)]}{E[1 - D_i]}.
$$

\triangleright We can write

$$
ATE = \frac{E[Y_i D_i]}{E[D_i]} - \frac{E[Y_i (1 - D_i)]}{E[1 - D_i]},
$$

where the right hand side depends on the joint distribution of the observed random variables.

 \triangleright For estimation, we replace the population mean by the sample mean (this is sometimes called the analogue principle):

$$
\widehat{\text{ATE}} = \frac{\frac{1}{n} \sum_{i=1}^{n} Y_i D_i}{\frac{1}{n} \sum_{i=1}^{n} D_i} - \frac{\frac{1}{n} \sum_{i=1}^{n} Y_i (1 - D_i)}{\frac{1}{n} \sum_{i=1}^{n} (1 - D_i)}.
$$

- \triangleright We can check its consistency by using LLN and Slutsky's lemma.
- \triangleright This randomization assumption is not convincing when the individuals in the social experiments are people who may select into the treatment or not.

Comparison with the linear regression

- It seems that D_i is nothing but a dummy variable. Can we run a regression of Y_i on D_i and X_{i1}, \ldots, X_{ik} to estimate the ATE? Can the parameter of interest, the ATE, be formulated as a coefficient in a regression model.
- \triangleright One possible assumption is that

$$
Y_i = g(D_i, X_{i1}, ..., X_{ik}, \epsilon_i) = \gamma_0 + \gamma_1 D_i + \sum_{j=1}^k \beta_j X_{ij} + \epsilon_i.
$$

In this case, the ITE $Y_i(1) - Y_i(0) = \gamma_1$ is constant. This is very unrealistic. We investigate alternative model assumptions.

 \triangleright We first consider the following model assumption

$$
Y_i(0) = \mu_0 + U_i(0)
$$

$$
Y_i(1) = \mu_1 + U_i(1),
$$

where μ_0 and μ_1 are constants common across individuals and assumed to be nonstochastic and $(U_i(0), U_i(1))$ are stochastic components.

- \blacktriangleright We denote $X_i = (X_{i1}, ..., X_{ik})^\top$ for the vector of observed covariates.
- \blacktriangleright We assume $E[U_i(0) | X_i] = E[U_i(1) | X_i]$, which implies

$$
E[Y_i(1) - Y_i(0) | X_i] = \mu_1 - \mu_0,
$$

i.e., the ITE is mean independent of X_i but it can be random. And by LIE,

$$
ATE = E[Y_i(1) - Y_i(0)] = \mu_1 - \mu_0.
$$

- \blacktriangleright We assume $E[Y_i(1) | D_i, X_i] = E[Y_i(1) | X_i]$ and $E[Y_i(0) | D_i, X_i] = E[Y_i(0) | X_i]$, i.e., the conditional mean independence of potential outcomes with treatment status, conditional on demographic status X_i .
- \triangleright When we focus on a sub-population of indivdiuals with specific demographic status X_i , Y_i (1) and Y_i (0) are both mean independent of D_i .

 \blacktriangleright Let us write

$$
E[Y_i | D_i, X_i] = D_i E[Y_i(1) | D_i, X_i] + (1 - D_i) E[Y_i(0) | D_i, X_i]
$$

=
$$
D_i E[Y_i(1) - Y_i(0) | D_i, X_i] + E[Y_i(0) | D_i, X_i]
$$

=
$$
D_i E[Y_i(1) - Y_i(0) | X_i] + E[Y_i(0) | X_i],
$$

where the last equality follows from the conditional mean independence assumption.

 \blacktriangleright By the assumption $E[U_i(0) | X_i] = E[U_i(1) | X_i]$, we have

$$
D_i E [Y_i (1) - Y_i (0) | X_i] + E [Y_i (0) | X_i]
$$

= $D_i (\mu_1 - \mu_0) + E [Y_i (0) | X_i]$
= $\mu_0 + D_i (\mu_1 - \mu_0) + g (X_{i1}, ..., X_{ik}),$

where we denote $g(X_{i1},..., X_{ik}) = E[U_i(0) | X_i]$.

 \blacktriangleright Therefore, we have

$$
E[Y_i | D_i, X_i] = \mu_0 + (\mu_1 - \mu_0) D_i + g(X_{i1}, ..., X_{ik}).
$$

 \blacktriangleright Define

$$
V_i = Y_i - E[Y_i | D_i, X_i]
$$

and now we have the following regression model:

$$
Y_i = \mu_0 + (\mu_1 - \mu_0) D_i + g(X_{i1}, ..., X_{ik}) + V_i.
$$

- \blacktriangleright We have E $[V_i | D_i, X_i] = 0$ by definition.
- \blacktriangleright We assume g is linear in $X_{i1}, ..., X_{ik}$:

$$
g(X_{i1},...,X_{ik}) = \sum_{j=1}^{k} \beta_j X_{ij},
$$

and then

$$
Y_i = \mu_0 + (\mu_1 - \mu_0) D_i + \sum_{j=1}^k \beta_j X_{ij} + V_i.
$$

A multiple linear regression of Y_i on D_i and $X_{i1}, ..., X_{ik}$ consistently estimates ATE = $(\mu_1 - \mu_0)$.

 \blacktriangleright We assume $E[U_i(0) | X_i] = E[U_i(1) | X_i]$, which implies

 $E[Y_i(1) - Y_i(0) | X_i] = \mu_1 - \mu_0.$

- \blacktriangleright This assumption implies that the conditional average treatment effect given X_i does not depend on X_i , the characteristics of individual i .
- \blacktriangleright This assumption can be unrealistic. E.g., Average treatment of the class-size is the same between students from high-income family and students from low-income family.

Unconfoundedness assumption

- \triangleright Unconfoundedness is the key assumption of the basic treatment effect model.
- \blacktriangleright Unconfoundedness assumption: $(Y_i(1), Y_i(0)) \perp D_i | X_i$, i.e., $(Y_i(1), Y_i(0))$ and D_i are conditionally independent given X_i .
- \triangleright Unconfoundedness can be thought of as an assumption that the decision to take the treatment is purely random for individuals with similar values of the covariates.
- \blacktriangleright Suppose that we have three random vectors V, W and X, where (V, W) is a continuous random vector. Then we say V and W are conditionally independent given X , if for all possible values of ν , w and x ,

$$
f(v, w)|x(v, w | x) = f_{V|X}(v | x) f_{W|X}(w | x).
$$

Inconfoundedness is satisfied if (Y_i, D_i) are generated by the model

$$
Y_i = g(D_i, X_{i1}, ..., X_{ik}, \epsilon_i)
$$

$$
D_i = m(X_{i1}, ..., X_{ik}, \eta_i)
$$

and $\epsilon_i \perp \eta_i \mid X_{i1}, ..., X_{ik}$.

More on conditional independence

 \blacktriangleright When V and W are conditionally independent given X, one can easily see that for any function φ ,

 $E[\varphi(V) | W, X] = E[\varphi(V) | X].$

I.e., once we observe X , knowledge of W does not give us any further advantage in predicting the value of $\varphi(V)$.

 \triangleright We notice that

$$
f(v,w)|x(v,w|x) = \frac{f(v,w,x)(v,w,x)}{f_X(x)} = \frac{f(v,w,x)(v,w,x)}{f(w,x)(w,x)} \frac{f(w,x)(w,x)}{f_X(x)} = f_{V|(w,x)}(v|w,x) f_{W|X}(w,x).
$$

Finerefore, we have $f_{V|X}(v|x) = f_{V|(W,X)}(v|w,x)$, if (V,W) are conditionally independent given X . Hence,

$$
E[\varphi(V) | W = w, X = x] = \int \varphi(v) f_{V|(W,X)}(v | w, x) dv
$$

=
$$
\int \varphi(v) f_{V|X}(v | x) dv
$$

=
$$
E[\varphi(V) | X = x].
$$

 \blacktriangleright Therefore, the unconfoundedness assumption $(Y_i(1), Y_i(0)) \perp D_i | X_i$ implies the conditional mean independence assumption:

$$
E[Y_i(1) | D_i, X_i] = E[Y_i(1) | X_i]
$$

$$
E[Y_i(0) | D_i, X_i] = E[Y_i(0) | X_i].
$$

 \triangleright We can also show: if $V \perp W \mid X$,

$$
E[\eta(V, W) | X, W = w] = E[\eta(V, w) | X].
$$
 (2)

The unconfoundedness and randomization assumptions

- \blacktriangleright It can be shown that the randomization assumption $(Y_i(1), Y_i(0), X_i) \perp D_i$ implies the unconfoundedness assumption $(Y_i(1), Y_i(0)) \perp D_i | X_i$.
- \blacktriangleright The randomized experiment assumption does not allow $X_{i1}, ..., X_{ik}$ to be correlated with D_i ,
- \blacktriangleright The unconfounded condition allows D_i to be affected by X_{i1}, \ldots, X_{ik} , while the randomized experiment assumption does not.

Identification of ATE

 \blacktriangleright By LIE, we have

ATE = E[
$$
Y_i(1) - Y_i(0)
$$
]
= E[E[$Y_i(1) | X_i$]] - E[E[$Y_i(0) | X_i$]], (3)

and

$$
E[Y_i D_i | X_i] = E[E[Y_i D_i | X_i, D_i] | X_i]
$$

= Pr[D_i = 1 | X_i] E[Y_i D_i | X_i, D_i = 1]
+ Pr[D_i = 0 | X_i] E[Y_i D_i | X_i, D_i = 0].

► By the unconfoundedness assumption: $(Y_i(1), Y_i(0)) \perp D_i | X_i$, the result [\(2\)](#page-21-0) and the relation $Y_i = D_i Y_i (1) + (1 - D_i) Y_i (0)$, we have

$$
E[Y_i D_i | X_i, D_i = 1]
$$

= E[(D_i Y_i (1) + (1 - D_i) Y_i (0)) D_i | X_i, D_i = 1] = E[Y_i (1) | X_i] and

$$
E[Y_iD_i | X_i, D_i = 0] = 0.
$$

 \blacktriangleright Therefore, we have

$$
E[Y_i D_i | X_i] = Pr[D_i = 1 | X_i] E[Y_i(1) | X_i]
$$
 (4)

and similarly,

$$
E[Y_i (1 - D_i) | X_i] = Pr[D_i = 0 | X_i] E[Y_i (0) | X_i].
$$
 (5)

 \blacktriangleright Now [\(3\)](#page-23-0), [\(4\)](#page-24-0), [\(5\)](#page-24-1) and LIE imply

$$
\begin{aligned} \text{ATE} &= \mathbb{E} \left[\frac{\mathbb{E} \left[Y_i D_i \mid X_i \right]}{\Pr \left[D_i = 1 \mid X_i \right]} \right] - \mathbb{E} \left[\frac{\mathbb{E} \left[Y_i \left(1 - D_i \right) \mid X_i \right]}{\Pr \left[D_i = 0 \mid X_i \right]} \right] \\ &= \mathbb{E} \left[\frac{Y_i D_i}{\Pr \left[D_i = 1 \mid X_i \right]} - \frac{Y_i \left(1 - D_i \right)}{\Pr \left[D_i = 0 \mid X_i \right]} \right]. \end{aligned}
$$

Now the right hand side depends only on the joint distribution of observed random variables.

 \blacktriangleright Denote

$$
p(x) = Pr[D_i = 1 | X_i = x].
$$

This function is called propensity score. It is the probability of the event that the individual belongs to the treatment group, given that the observed characteristics are $\mathbf{x} \in \mathbb{R}^k$.

Estimation of ATE

 \blacktriangleright Let $\hat{p}(x)$ be an estimator of the propensity score, then we can estimate the ATE:

$$
\widehat{\text{ATE}} = \frac{1}{n} \sum_{i=1}^{n} \left\{ \frac{Y_i D_i}{\hat{p}(X_i)} - \frac{Y_i (1 - D_i)}{1 - \hat{p}(X_i)} \right\}.
$$

It is straightforward to construct $\hat{p}(x)$ if X_i is discrete:

$$
\hat{p}(x) = \frac{\sum_{i=1}^{n} 1 (D_i = 1, X_i = x)}{\sum_{i=1}^{n} 1 (X_i = x)}
$$

.

If X_i is continuous, we specify a parametric model for the propensity score:

$$
Pr[D_i = 1 | X_i] = \Phi(\beta_0 + \beta_1 X_{i1} + \dots + \beta_k X_{ik})
$$

as what we did for the Probit model. This gives a parametric model for the propensity score. $(\beta_0, ..., \beta_k)$ can be estimated by MLE (denoted by $(\hat{\beta}_0, ..., \hat{\beta}_k)$).

 \blacktriangleright The estimated propensity score is

$$
\hat{p}(X_i) = \Phi\left(\hat{\beta}_0 + \hat{\beta}_1 X_{i1} + \cdots + \hat{\beta}_k X_{ik}\right).
$$

- \triangleright This estimator is known to be consistent and asymptotically normally distributed, if our propensity score model is correct.
- \blacktriangleright This approach has the drawback that if our model for the propensity score is wrong, the ATE estimator is inconsistent.
- Actually, $p(x) = E[D_i | X_i = x]$ can be estimated without specifying a parametric model for it.

k-NN estimator

- \blacktriangleright The *k*-nearest neighbor (*k*-NN) estimator is the simplest nonparametric estimator of $p(x)$.
- Fix x_0 and suppose that we want to estimate $p(x_0)$ at this point. Assume that p is a smooth function, which means that its graph does not change too much.
- \blacktriangleright p(x) should be close to p(x₀) when x is close enough to x₀. $p(X_i)$ would be close to $p(x_0)$ for observations X_i close to x_0 .
- \blacktriangleright We simply average these $p(X_i)$ for observations X_i close to x_0 . We do not observe $p(X_i)$ but use D_i instead.

 \blacktriangleright Let

$$
d_i(\mathbf{x}_0) = ||\mathbf{X}_i - \mathbf{x}_0|| = \sqrt{(\mathbf{X}_i - \mathbf{x}_0)^\top (\mathbf{X}_i - \mathbf{x}_0)}
$$

denote the distance of X_i to x_0 .

 \blacktriangleright After computing the distance for all *n* observations in the sample, we sort them in the increasing order

$$
d_{(1)}(x_0) \leq d_{(2)}(x_0) \leq \cdots \leq d_{(n)}(x_0).
$$

 \blacktriangleright Let $N_k(x_0)$ denote the identities of the k-nearest neighbors of x_0 :

$$
N_k(x_0) = \{i : d_i(x_0) \leq d_{(k)}(x_0)\}.
$$

 \blacktriangleright The *k*-NN nonparametric estimator of $p(x_0)$ is

$$
\hat{p}_{kNN}\left(\boldsymbol{x}_0\right) = \frac{1}{k} \sum_{i \in N_k(\boldsymbol{x}_0)} D_i.
$$

- \triangleright The k-NN estimator is simply an average of the values of D_i across the k closest observations in terms of X_i .
- \blacktriangleright There is a data-driven procedure to select k in practical applications.
- \blacktriangleright The nonparametric ATE estimator using $\hat{p}_{kNN}(X_i)$ is consistent and asymptotically normal. It does not require a parametric model for the propensity score.