

Propensity score matching (PSM)

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January 2024

Estimation of the treatment effect

- Examples:
 - evaluation of the efficiency of the activation programmes in reducing unemployment
 - evaluation of the effects of the innovation grants

Ideal social experiment

- ① members of the experimental group and control group has the same distribution of unobserved characteristics
- ② members of the experimental group and control group has the same distribution of observed characteristics
- ③ the same questionnaire was used in experimental and control group - characteristics and outcomes measured in the same way
- ④ both groups are in the same economic environment

- Ideal evolution study is based on the random experiment: for such a study the treatment effect can be measured simply as the difference between the mean outcome variable in the treatment and control groups
- in such a case assumptions (1)-(4) are satisfied by definition
- Traditional econometrics analysis of problems related to evolution studies based on nonexperimental data was concentrated on assumption (1) - e.g. Heckman model
- in practice it seems that more important are the problems related to assumptions (2), (3), (4).
- PSM make possible to reduce bias of the estimated treatment effect related to invalidity of the assumption (2).

- $i \in I_1$ experimental group
- $j \in I_0$ control group
- $D \in \{0, 1\}$
 - $D = 0$ unit is untreated
 - $D = 1$ unit is treated
- X - vector of characteristics of the unit
- $\Pr(X) = \Pr(D = 1|X)$ conditional probability to be a member of experimental group (propensity score)
- Treatment effect is measured with the outcome variable Y
 - Y_0 value of the outcome variable if untreated
 - Y_1 value of the outcome variable if treated

- Treatment Effect for a unit with characteristics given by X

$$E(Y_1 - Y_0 | D = 1, X)$$

- Average Treatment Effect - ATE

$$ATE = E(Y_1 - Y_0)$$

- Average Treatment Effect on the treated - ATE_1

$$ATE_1 = E(Y_1 - Y_0 | D = 1)$$

- We observe Y

$$Y = DY_1 + (1 - D) Y_0$$

- Every unit has the observed effect and counter-factual effect

	I_0	I_1
$D = 0$	Y_0	Y_1
$D = 1$	Y_0	Y_1

- It is not possible to observe the outcome of the treatment for the control group $E(Y_0 | D = 1, X)$ (counterfactual effect)

Intuition of PM

- Treatment effect is based on comparison of the outcomes for treated units and untreated units.
- The more similar is the untreated unit to the treated unit the larger is the weight of this observation in this comparison
- Estimate of the treatment effect for unit i

$$Y_{1i} - \sum_{j \in I_0} W_{N_0, N_1}(i, j) Y_{0j}$$

- Estimated of the average treatment effect (ATE)

$$\sum_{i \in I_1} w_{N_0, N_1}(i) \left[Y_{1i} - \sum_{j \in I_0} W_{N_0, N_1}(i, j) Y_{0j} \right]$$

- Various PM variants differ in the way the weight functions $w_{N_0, N_1}(i)$ and $W_{N_0, N_1}(i, j)$ are defined.

Statistical independence

- Definition

$$\Pr(D = 1 | Y_0, Y_1, X) = \Pr(D = 1 | X) \quad (1)$$

- Selection based only on observable variables
- We assume that there is no self selection based on unobserved variables
- Excluded is the selection based on unobserved variables correlated with (influencing) the outcome variable as in Heckman model
- Equivalent notation

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

- Stable Unit Treatment Assumption SUTVA - outcome of one unit is independent from the outcomes of other units

Strong statistical independence

- Common support assumption

$$0 < \Pr(X) < 1 \quad (2)$$

- Strong statistical independence: (1) (2) satisfied
- Condition(2) is necessary condition for application of PSM.
- If $\Pr(X) = 0$ or $\Pr(X) = 1$, the such at unit is never treated or always treated and then forming the pair of the treated and untreated observations is impossible
- If condition (2) is invalid for some X we have to restrict our population to subpopulation for which this condition is satisfied
- Define

$$S_0 = \text{Supp}(X|D = 0)$$

$$S_1 = \text{Supp}(X|D = 1)$$

- Common support

$$S = S_0 \cap S_1$$

Bias decomposition in evaluation studies

- Difference in outcomes in treatment and control groups:
 $E(Y_0|D=1, X \in S_1) - E(Y_0|D=0, X \in S_0) = B_1 + B_2 + B_3$
- Bias related to different in supports

$$B_1 = E[E(Y_0|D=1, X \in S_1 \setminus (S_0 \cap S_1))|D=1] \\ - E[E(Y_0|D=0, X \in S_0 \setminus (S_0 \cap S_1))|D=0]$$

- Bias related to differences in distributions of observed characteristics in experimental and control groups:

$$B_2 = E[E(Y_0|D=0, X \in S_0 \cap S_1)|D=1] \\ - E[E(Y_0|D=0, X \in S_0 \cap S_1)|D=0]$$

- Bias related to differences in distributions of unobserved characteristics in experimental and control groups:

$$B_3 = E[E(Y_0|D=1) - E(Y_0|D=0)|D=1, X \in S_0 \cap S_1]$$

- Application of PSM makes possible the correction of the bias B_2 and the determination of the common support for which bias B_1 is eliminated.

Implications of the statistical independence

- Strong statistical independence implies that, the average counterfactual effect can be calculated as follows

$$\begin{aligned} E(Y_0 | D = 1) &= E[E(Y_0 | D = 1, X) | D = 1] \\ &= E[E(Y_0 | D = 0, X) | D = 1] \end{aligned}$$

- Assume that we succeeded to estimate (parametrically or non parametrically) the regression $\hat{r}_0(X_i) = E(Y_0 | D = 0, X_i)$ and $\hat{r}_1(X_i) = E(Y_1 | D = 1, X_i)$
- Estimate of ATE_1 can be obtained as the following mean

$$\frac{1}{N_1} \sum_{i \in I_1} (\hat{r}_1(X_i) - \hat{r}_0(X_i))$$

- The assumption needed for this estimator is that $Y_0 \perp D | X$
- In fact to estimate ATE_1 sufficient condition is that of independence of the conditional expectations from X

$$E(Y_0 | D = 1, X) = E(Y_0 | D = 0, X)$$

- Strong statistical independence implies that (Resenbaum, Rubin 1983)

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

- Proof

$$\begin{aligned} E[D | Y_0, Y_1, \Pr(X)] &= E \{ E[D | Y_0, Y_1, X] | \Pr(X) \} \\ &= E [\Pr(D = 1 | Y_0, Y_1, X) | \Pr(X)] \\ &= E \{ \Pr(D = 1 | X) | \Pr(X) \} \\ &= E \{ \Pr(X) | \Pr(X) \} = \Pr(X) \\ &= E[D | \Pr(X)] \end{aligned}$$

Implications of the statistical independence

- Resenbaum and Rubina result implies, that the independence of outcome from the membership in the group (experimental/control) is satisfied not only conditional on all the characteristics but also conditional on propensity score $\Pr(X)$ only
- Notice that if we have observations from experimental and control groups with identical $\Pr(X)$, then ATE_1 can be calculated as follows:

$$E[Y_1 | D = 1, \Pr(X)] - E[Y_0 | D = 0, \Pr(X)] = E[Y_1 - Y_0 | \Pr(X)]$$

- Choosing the observations to the control group we can base the choice on the similarity of the propensity score $\Pr(X)$ only

In PSM control group is chosen from the sample of untreated units on the basis of the similarity of the estimated values of the propensity scores $\Pr(X)$

- Parametric methods (probit, logit)
- Nonparametric methods
- Crucial element in estimation of $\Pr(X)$ is to make sure that the common support assumption is satisfied
 - range of the predicted values should be similar
 - predictors must not perfectly predict outcomes
- In some sense the model for $\Pr(X)$ should not be "too well fitted"

PSM - choosing the observations to the control group

- Selecting the observations with replacement or without replacement
- Number of the untreated units chosen for one treated: 1 do 1, 1 do n
- Selection algorithm:
 - nearest neighbor
 - nearest neighbor with caliper
 - radius matching
 - kernel method

$$w_{ij} = \frac{K\left(\frac{\text{Pr}(X_i) - \text{Pr}(X_j)}{h}\right)}{\sum_{j \in I_0} \left(\frac{\text{Pr}(X_i) - \text{Pr}(X_j)}{h}\right)}$$

where h is the bandwidth

- The choice of the algorithm depends mainly on the sample size

PSM - checking the matching quality

- In the best case the control group selected on the basis of PSM should be large and has the distribution of observable characteristics identical to experimental group
- When choosing the selection algorithm we should take into account two factors:
 - the bias of evaluated treatment effect: the smaller the more similar are the units in control group to the units in treatment group
 - the variance of the estimate of the treatment effect: the smaller the larger is the control group
- The quality of matching is tested by comparing mean characteristics in treatment and control groups
- For this effect we can use the formal tests for equal means

PSM - advantages and disadvantages

- Advantages

- In the case of PSM we make no assumptions about the functional form of the relationship between the expected outcomes and the values of characteristics - relationship $E(Y_1|D=1,X) - E(Y_0|D=1,X)$ is left unspecified and can have quite general form
- eliminates two most important sources of the estimation bias

- Disadvantages

- the strong statistical independence assumptions must be satisfied
 - absence of self selection
 - absence of selection based on unobserved characteristics
- needs large number of observations - especially the sample of untreated observations should be large to make possible selection of the untreated units sufficiently similar to treated observations