

## Understanding Propensity scores: From naïve enthusiasm to intuitive understanding

New methodologies have been emerging to estimate the effect of a binary exposure on an outcome or result. Propensity scores has been proven to be useful to estimate the causal effect of exposure under certain assumptions even in the presence of confounding variables. This paper aims to define what is causal effect, detail how the propensity score can estimate causal effects as well as the assumptions and concerns about this method using an example of exposure to breast milk and the infant's consequent neurodevelopment (IQ) after 7.5 years. In this example, each individual has two potential outcomes, one in the individual was exposed to breastfeeding or not but only one outcome can be observed from the data and the other outcome is defined as counterfactual. Therefore, the causal effect can be defined as the mean of the individual causal effect and can be estimated by using results of other individuals to calculate counterfactual outcomes that were not observed.

Three different causal effects are of interest and are used depending on the questions that wants to be answered: the Average Causal Effect of the Exposure ( $ACE_{ALL}$ ) which is the causal effect in the entire population, the Average Causal Effect of the Exposure on the Exposed ( $ACE_{EXP}$ ) which considers the subgroup of the population that is exposed, and the Average Causal Effect of the Unexposed ( $ACE_{UNE}$ ) that considers the subgroup of the unexposed.

Given that is our intention to convert causal estimands to statistical estimands as part of the identification step, a few assumptions and properties are needed. For this specific example taken from [1] the intention is to be able to estimate the Average Treatment Effect represented by  $E[Y_1 - Y_0]$  using propensity scores. The assumptions made were that the treatment assignment precedes the effect  $Y$ , that every subject must have the potential to be exposed and unexposed, Stable Unit Treatment Value Assumption (SUTVA) given that data were sample independently from the population and that there are not unobserved confounders. This last assumption is also known as Strongly Ignorable Treatment Assignment (SITA) given that the observed covariates are conditionally independent.

Figure 1 shows the appropriate Single World Intervention Graph (SWIG) for this problem that is used to represent potential outcomes and their conditional independence relations where  $X$  are the covariates of the,  $A$  is the treatment and  $Y$  is the outcome.

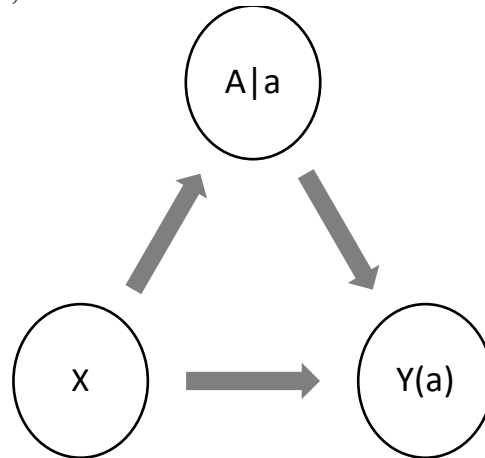


Figure 1: Single World Intervention Graph.

The following steps demonstrates how to estimate the Average Treatment Effect ( $ATE_{all}$ )

$$ATE_{all} = E[Y(1) - Y(0)] \quad (1)$$

Assumptions:

- $Y(a) \perp A|e(x)$
- Consistency

First, we are going to look at

$$E[Y(1) - Y(0)|e(X) = x] = E[Y(1)|e(x) = x] - E[Y(0)|e(x) = x] \quad (2)$$

which is possible because of expectation properties.

In second place, if  $Y(a) \perp A|e(x)$  and because of consistency:

$$\begin{aligned} E[Y(1)|A = 1, e(x) = x] - E[Y(0)|A = 0, e(x) = x] \\ E[Y|A = 1, e(x) = x] - E[Y|A = 0, e(x) = x] \end{aligned} \quad (3)$$

Then, Because of the Law of iterated Expectations:

$$E[E[Y(1) - Y(0)|e(x)]] = E[Y(1) - Y(0)] \quad (4)$$

Therefore,

$$E[E[Y(1) - Y(0)|e(x)]] = E[Y|A = 1, e(x)] - E[Y|A = 0, e(x)] \quad (5)$$

$$E[E[Y(1) - Y(0)|e(x)]] = E[E[Y|A = 1, e(x)] - [Y|A = 0, e(x)]] \quad (6)$$

$$E[Y(1) - Y(0)] = E[E[Y|A = 1, e(x)] - E[Y|A = 0, e(x)]] \quad (7)$$

In (7), we showed that it is possible to estimate the Average Treatment Effect from the data provided.

Causal Effects can be estimated using four different propensity scores methods: stratification, matching, inverse weighting, and covariate adjustment. The stratification method consists of making strata or groups of the individuals who have the same propensity score, estimating the exposure effect between exposed and unexposed group and using a weighted average. The matching method consist of pairing each exposed individual with another unexposed with the same propensity score and calculating the average pair estimate of the effect of the exposure after taking the difference in the two outcomes of within-pair effect. The inverse weighting creates two potential samples, each one for each outcome, assigning a replica to the original individual's outcome based on the propensity score and individual's characteristics. That is, if the propensity score is 1/2 for low-income individuals, then the replica should have the same number of individuals for high income entities and the same outcome of the original individual. Finally, the covariate adjustment can be estimated using a linear regression where the dependent variable would be the outcome and the independent variables to be the exposure and the propensity score. The regression coefficient for the propensity score variable should result in

the estimate of the exposure effect. Figure 2 provides a graphical representation of the four different estimation methods.

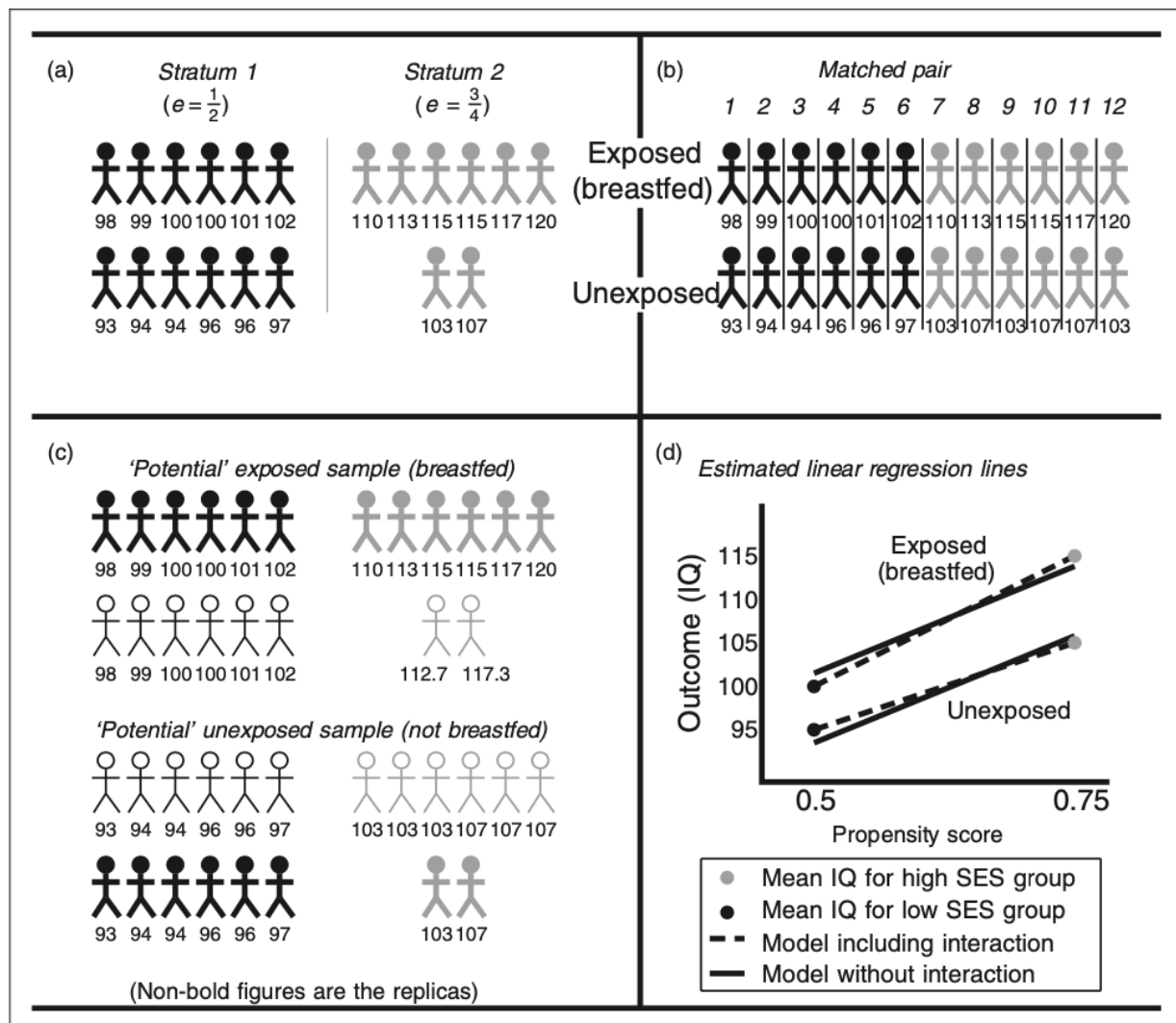


Figure 2: Estimation Methods [1]

To show that the procedure to estimate the causal effect using propensity scores, a logistic regression was performed to determine the propensity scores and the matching procedure was followed to estimate the average treatment effect using the NHEFS data from 1629 cigarette smokers aged 25-74 years who had a baseline visit and a follow-up visit about 10 years later.

The first step to implement this methodology is to estimate the propensity scores. In this case, a logistic regression was performed where the response variable is “quit smoking”. After that, propensity scores are obtained by extracting the response of the logistic regression model.

```
# Create Logistic Regression Model

propensity_model <- glm(qsmk ~ (seqn+death+yrdth+modth+dadth+sbp+dbp+sex+age+r
ace+income+marital+school+education+ht+wt71+wt82+birthplace+smokeintensity+sm
kintensity82_71+smokeyrs+asthma+bronch+tb+hf+hbp+pepticulcer+hepatitis+chroni
ccough+hayfever+diabetes+polio+tumor+nervousbreak+ alcoholpy+alcoholfreq+alco
holtype+alcoholhowmuch+pica+headache+otherpain+weakheart+allergies+nerves+lac
kpep+hbpmed+boweltrouble+wtloss+infection+active+exercise+birthcontrol+pregna
ncies+cholesterol+hightax82+price71+price82+tax71+tax82+price71_82+tax71_82),
family="binomial", data = df)

# calculate predicted propensity scores
df$ps_lgt <- predict(propensity_model, type = "link") #Logit
df$ps <- predict(propensity_model, type = "response")
```

After obtaining the propensity score, the second step is to perform matching to be able to estimate the causal effects. For this pick a treated subject and match them to an untreated subject with the closest propensity score one by one using 1-1 nearest neighbor matching without replacement, based on the logit of the propensity score.

```
NN_match <- matchit(qsmk ~ (seqn+death+yrdth+modth+dadth+sbp+dbp+sex+age+race+
income+marital+school+education+ht+wt71+wt82+birthplace+smokeintensity+smkint
ensity82_71+smokeyrs+asthma+bronch+tb+hf+hbp+pepticulcer+hepatitis+chroniccou
gh+hayfever+diabetes+polio+tumor+nervousbreak+ alcoholpy+alcoholfreq+alcoholt
ype+alcoholhowmuch+pica+headache+otherpain+weakheart+allergies+nerves+lackpep
+hbpmed+boweltrouble+wtloss+infection+active+exercise+birthcontrol+pregnancie
s+cholesterol+hightax82+price71+price82+tax71+tax82+price71_82+tax71_82),data
= df,distance = "logit", method = "nearest")
```

```
summary(NN_match)$nn
```

##	Control	Treated
## All (ESS)	31	13
## All	31	13
## Matched (ESS)	13	13
## Matched	13	13
## Unmatched	18	0
## Discarded	0	0

Finally, using the matched sample you can estimate the average treatment effect of the causal model proposed.

```
NNreplace_match_dat <- match.data(NNreplace_match)
NNreplace_match_dat <- data.frame(NNreplace_match_dat)

NNreplace_meandiff <- lm_robust(wt82_71 ~ qsmk, weights = weights, data = NNr
eplace_match_dat)
summary(NNreplace_meandiff)

##
## Call:
## lm_robust(formula = wt82_71 ~ qsmk, data = NNreplace_match_dat,
##           weights = weights)
##
## Weighted, Standard error type: HC2
##
## Coefficients:
##              Estimate Std. Error t value Pr(>|t|) CI Lower CI Upper DF
## (Intercept)    6.387      6.200   1.0301  0.3204  -6.912   19.686 14
## qsmk          -5.732      7.011  -0.8175  0.4273  -20.770    9.306 14
##
## Multiple R-squared:  0.0404 ,    Adjusted R-squared:  -0.02814
## F-statistic: 0.6683 on 1 and 14 DF,  p-value: 0.4273
```

Choosing a method and estimating the propensity score is not considered to be a trivial decision given that each one has advantages and disadvantages. Propensity scores are often estimated using logistic regression using all the confounders, but other non and semi-parametric approaches to this model have been proposed. In the breastfeeding example, a total of 926 babies were considered and were assessed at approximately 7.5 years of age by measuring their IQ. Additionally, other variables like social class, mother's education, family structure, marital status, infant age and gender, birthweight and birth order were considered. Propensity scores were estimated using a logistic regression and the three causal effects were calculated with the four methods but only 487 follow-up samples were collected. Results show that each methods performance was similar as shown in Figure 3, suggesting a positive relationship between breast milk consumption and IQ.

Causal inference is still a topic that deserves to be investigated more rigorously. Although relevance has been found in the studies carried out, there is still debate about the assumptions that are needed for these models to be used. Many of the areas of interest include estimating confidence intervals for propensity scores, using these propensity scores to train nonparametric models, and establishing a consensus for their proper use.

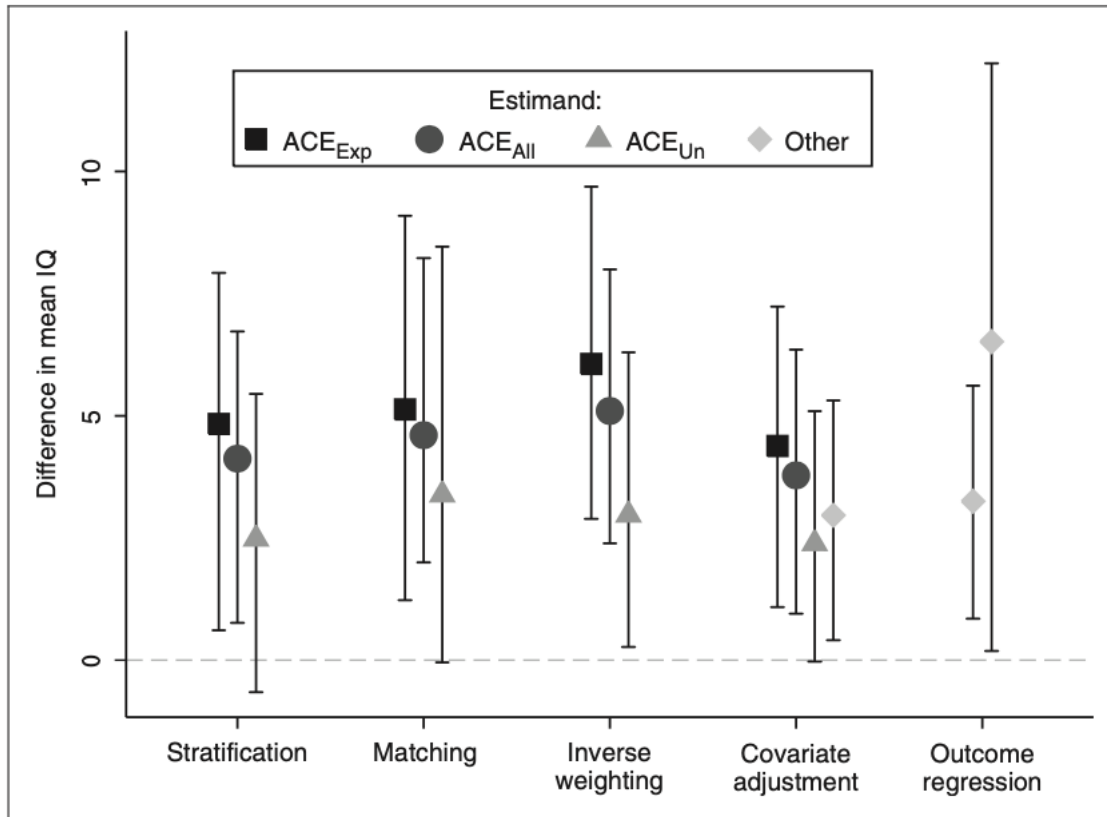


Figure 3: Results for each method in the BreastFeeding example [1]

- [1] Williamson, E., Morley, R., Lucas, A., & Carpenter, J. (2011). Propensity scores: From naïve enthusiasm to intuitive understanding. *Statistical Methods in Medical Research*, 21(3), 273–293. <https://doi.org/10.1177/0962280210394483>
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- [8] Weschler D. Weschler Intelligence Scale for Children, Anglicized revised edition. Sidcup, Kent: The Psychological Corporation Ltd.; 1974.
- [9] Paul R. Rosenbaum, Donald B. Rubin, The central role of the propensity score in observational studies for causal effects, *Biometrika*, Volume 70, Issue 1, April 1983, Pages 41–55,