

# Causal Data Science

Lecture 7:2 Estimating causal effects

Lecturer: Sara Magliacane



### Estimands for binary treatments

- We generally cannot estimate unit-level causal effect:  $Y_i(t=1)-Y_i(t=0)$
- We can estimate the average causal effect/average treatment effect

ATE = 
$$\mathbb{E}[Y(t=1) - Y(t=0)] = \mathbb{E}[Y|do(T=1)] - \mathbb{E}[Y|do(T=0)]$$



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We can also estimate the average causal effect of treatment on the treated:

$$ATT = \mathbb{E}[Y(t=1) - Y(t=0) | T=1]$$

We can also estimate the average causal effect of treatment on the control:

ATC = 
$$\mathbb{E}[Y(t=1) - Y(t=0) | T=0]$$



### Estimands for binary treatments

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- We can estimate the average causal effect/average treatment effect  $ATE = \mathbb{E}[Y(t=1) Y(t=0)] = \mathbb{E}[Y|\operatorname{do}(T=1)] \mathbb{E}[Y|\operatorname{do}(T=0)]$
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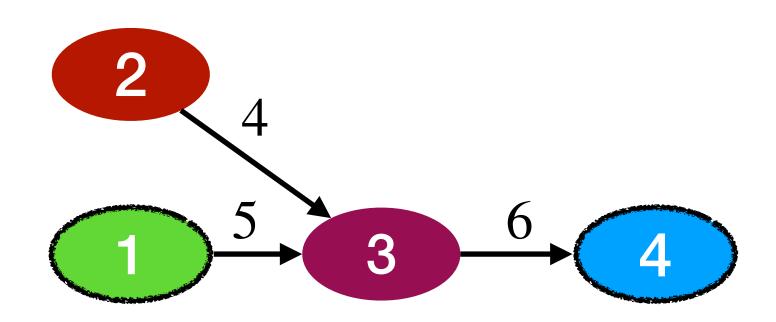
$$ATT = \mathbb{E}[Y(t=1) - Y(t=0) | T=1]$$

• For all, we assume that our covariates X form a valid adjustment set (e.g. we can check them/filter them with backdoor criterion)



#### Average causal effect/average treatment effect (ATE)

• ATE =  $\mathbb{E}[Y(t=1) - Y(t=0)] = \mathbb{E}[Y|do(T=1)] - \mathbb{E}[Y|do(T=0)]$ 



```
x2_1 = randn(n_samples)
x1_1 = 1
x3_1 = 5 * x1_1 + 4 * x2_1 + randn(n_samples)
x4_1 = 6 * x3_1 + randn(n_samples)

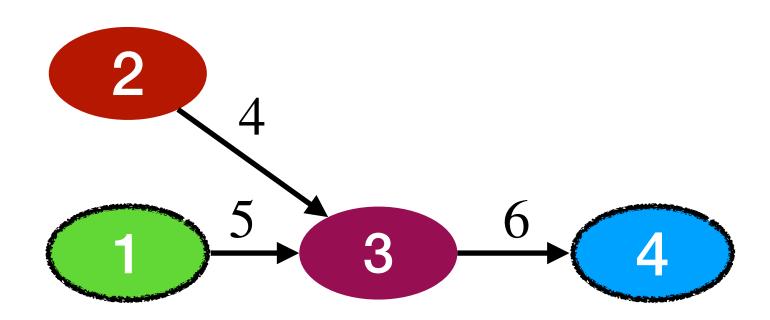
x2_0 = randn(n_samples)
x1_0 = 0
x3_0 = 5 * x1_0 + 4 * x2_0 + randn(n_samples)
x4_0 = 6 * x3_0 + randn(n_samples)
diff = np.mean(x4_1) - np.mean(x4_0)
print(diff)
```

30.514748479180785



#### Average causal effect/average treatment effect (ATE)

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How well does the treatment work on the patients who choose it?

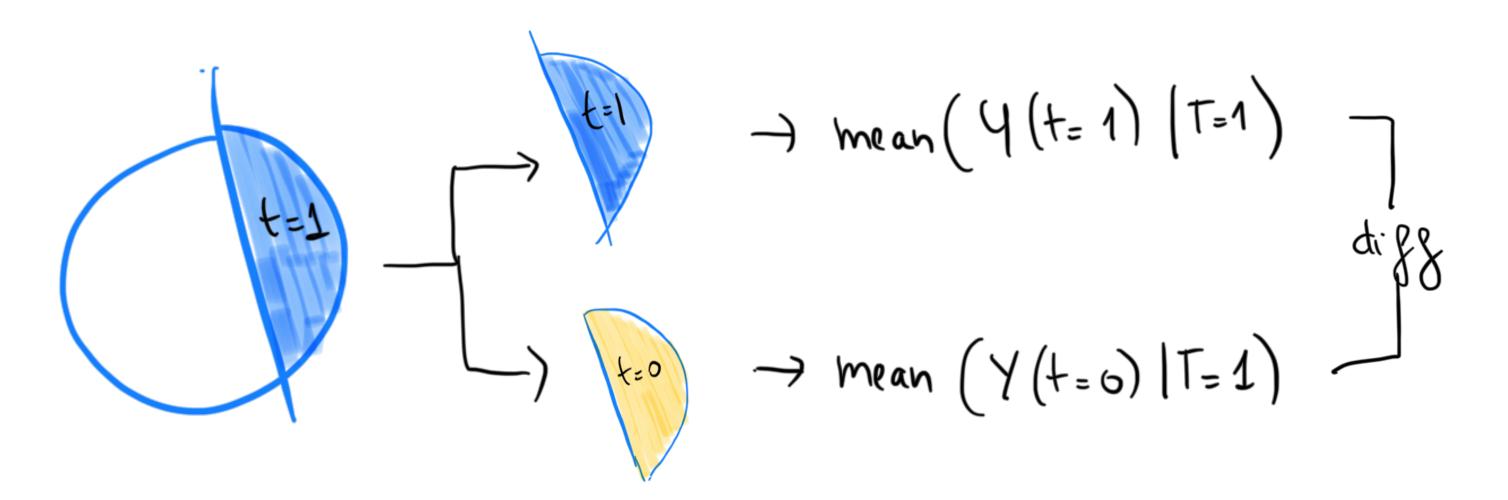
$$ATT = \mathbb{E}[Y(t=1) - Y(t=0) | T=1] \qquad \text{call of with in do () notation}$$



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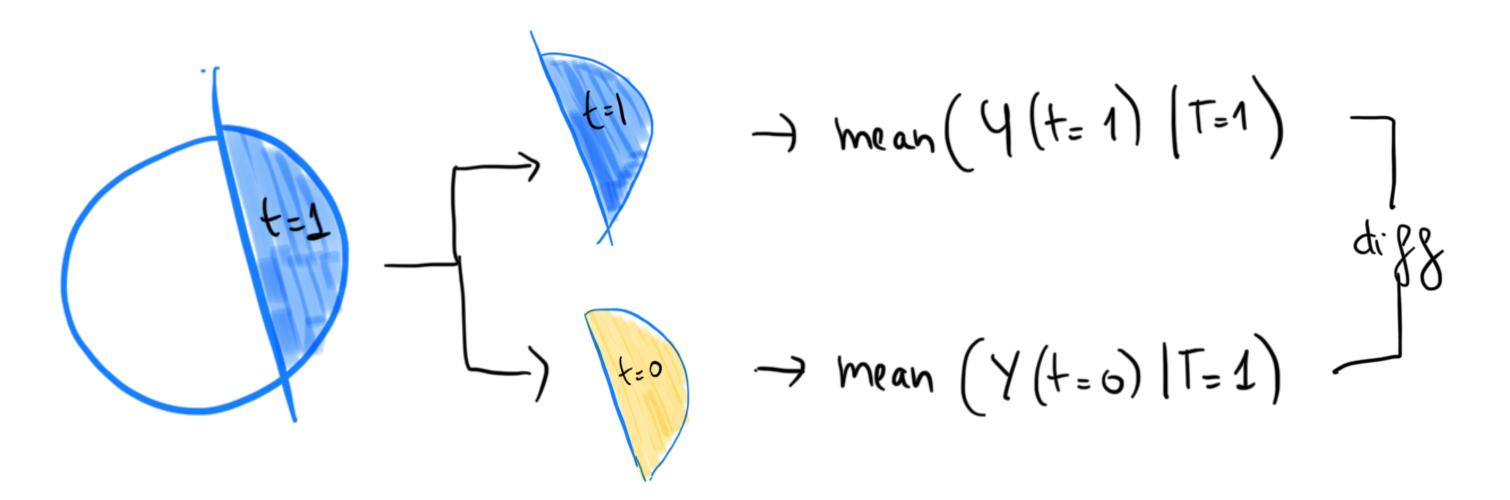




#### Average causal effect of treatment on the treated (ATT)

How well does the treatment work on the patients who choose it?

$$ATT = \mathbb{E}[Y(t=1) - Y(t=0) | T=1]$$



 Not the same as ATE: For example, people who choose a treatment could be more health-conscious, which means they get anyway better outcomes



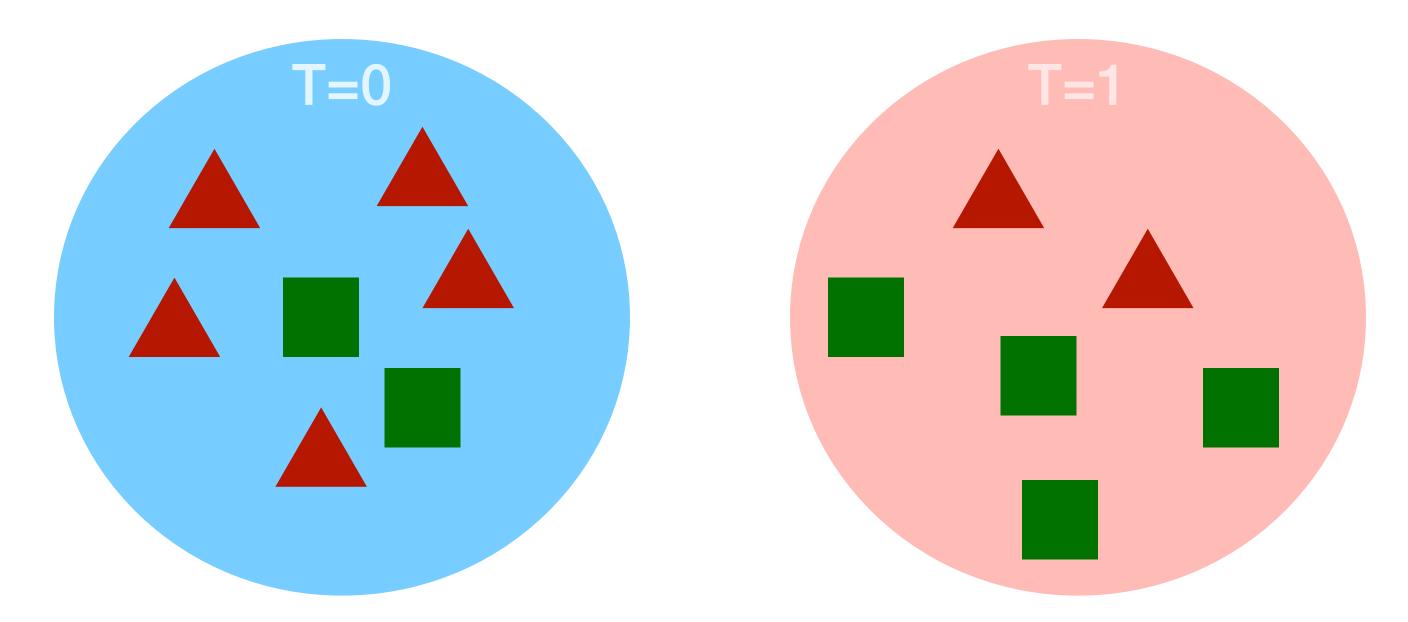
- Usually for ATT, sometimes for ATE
- Intuition: find the most similar couple of patients in terms of covariates  $\mathbf{X}$ , such that one is in the treatment and the other in the control group



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  - If successful, it's like an RCT
  - For example: I want to compare the outcomes of other people of my age

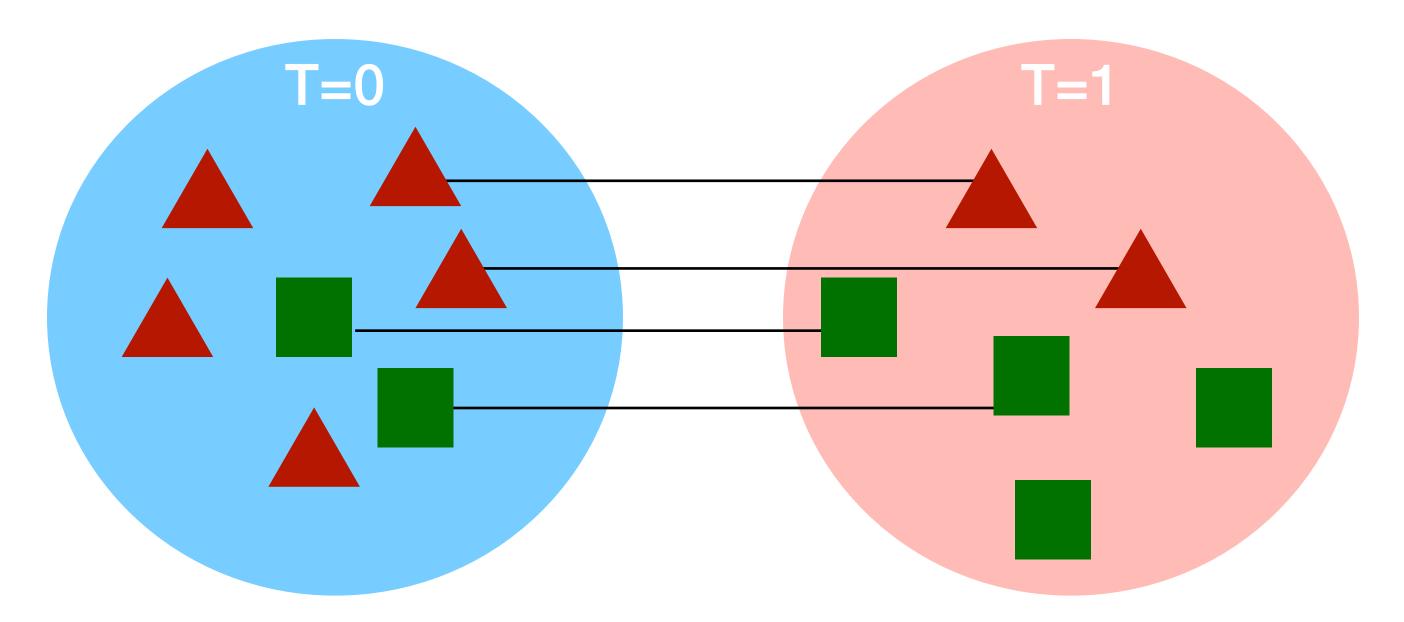


- Usually for ATT, sometimes for ATE
- Intution: find the most similar couple of units in terms of covariates X, such that one is in the treatment and the other in the control group



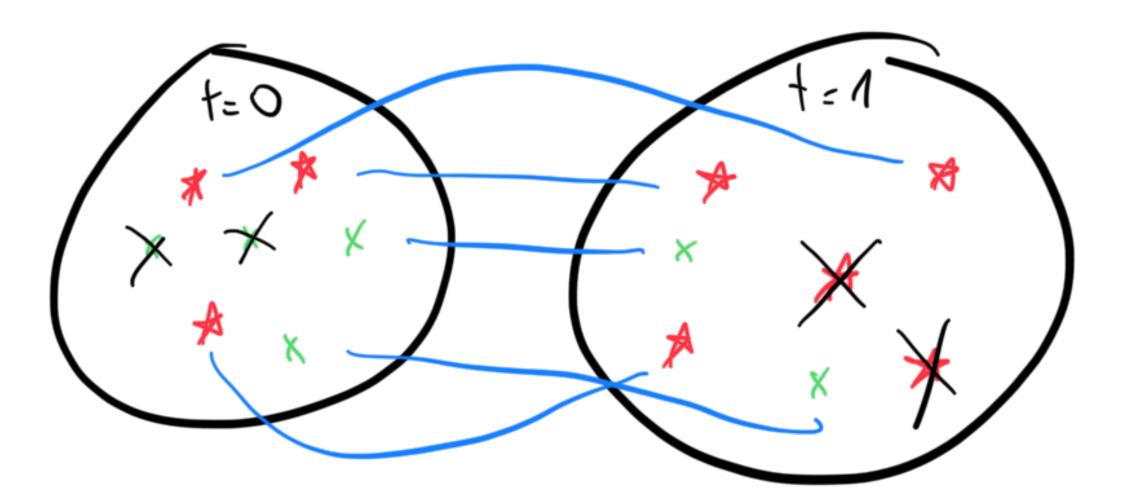


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- kNN
- Covariate balancing  $T \perp \mathbf{X} \equiv P(\mathbf{X} | T = 0) = P(\mathbf{X} | T = 1)$



### **Exact matching**

- Intuition: find the most similar couple of patients in terms of covariates  $\mathbf{X}$ , such that one is in the treatment and the other in the control group
  - For example: I want to compare the outcomes of other people of my age
- Note: we can only match units on variables we are adjusting:
  - Units with same values X = x in each group are indistinguishable
- Goal: discard unmatched units, so we have the same number of units with the same combination of values for X in treatment and control (balancing)



#### Matching - continuous covariates, greedy/optimal

• If exact matching on the value is not possible, e.g. because we have continuous covariates, we can use any distance, e.g Mahalanobis distance



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- Many variants exist, in general two types of algorithms:
  - Greedy matching: greedily and incrementally match treated with control based on distance
  - Optimal matching: optimize for the smallest total distance, can be slow



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- Many variants exist, in general two types of algorithms:
  - Greedy matching: greedily and incrementally match treated with control based on distance
  - Optimal matching: optimize for the smallest total distance, can be slow
- Need to check covariate balancing after matching (e.g. std mean difference)

$$T \perp \!\!\!\perp \mathbf{X} \equiv P(\mathbf{X} \mid T = 0) = P(\mathbf{X} \mid T = 1)$$



#### Estimation method: Propensity score matching (PSM)

- Assumptions: binary treatment T, X is valid adjustment set
- Propensity score: the probability of getting assigned the treatment

$$e(x)$$
  $\pi(x) := P(T = 1 | X = x)$ 



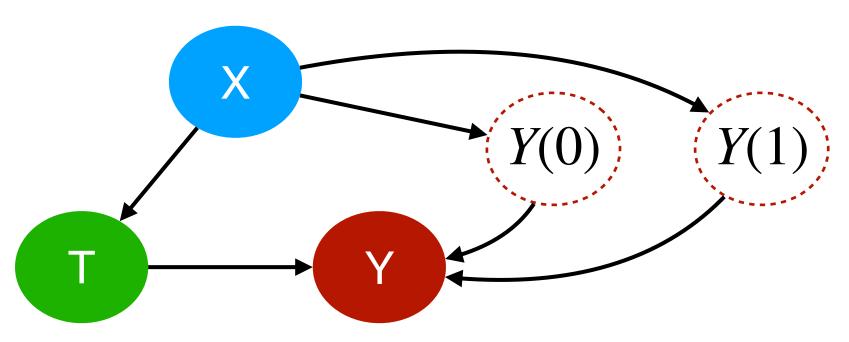
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Conditional ignorability/No unmeasured confounding

• We can show that  $T \perp \!\!\! \perp \mathbf{X} \mid \pi(\mathbf{X})$  and that if  $Y(0), Y(1) \perp \!\!\! \perp T \mid \mathbf{X}$  then





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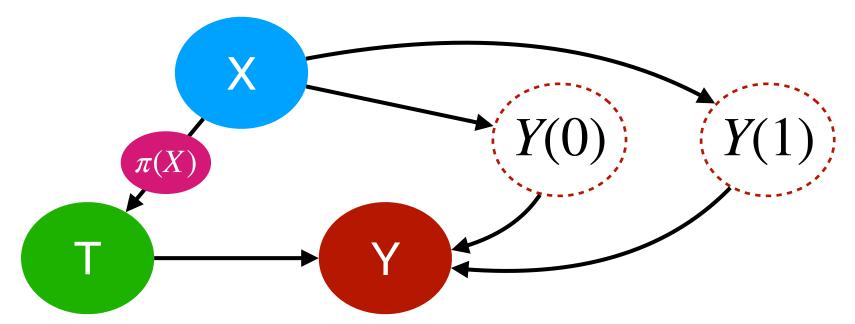
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$$Y(0), Y(1) \perp \!\!\! \perp T \mid \pi(X)$$

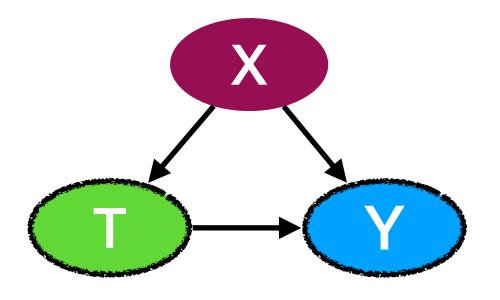
e.g. with logistic regression

- We can estimate  $\pi$  from data and use it to match
  - If X has a lot of covariates, it is easier to match since it's a single number





### Matching and IPW Jupyter notebook



$$P(\mathbf{X} = 1) = 0.3$$

$$P(T = 1 | \mathbf{X} = 1) = 0.1$$

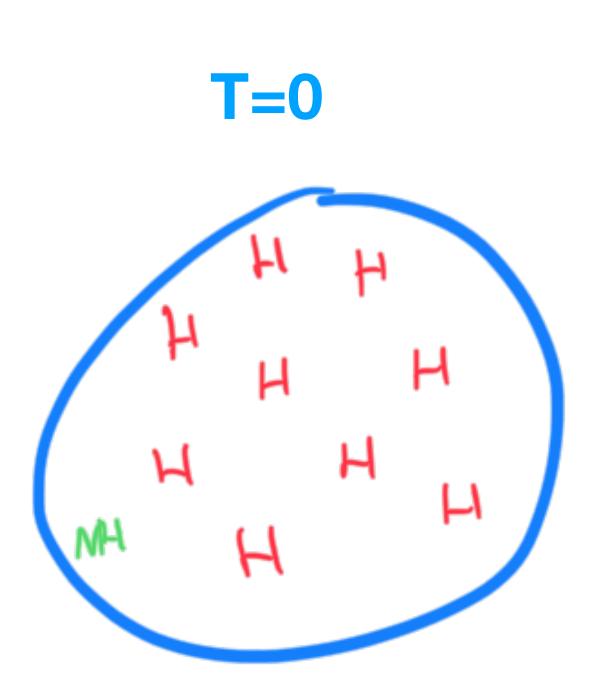
$$P(T = 1 | \mathbf{X} = 0) = 0.9$$

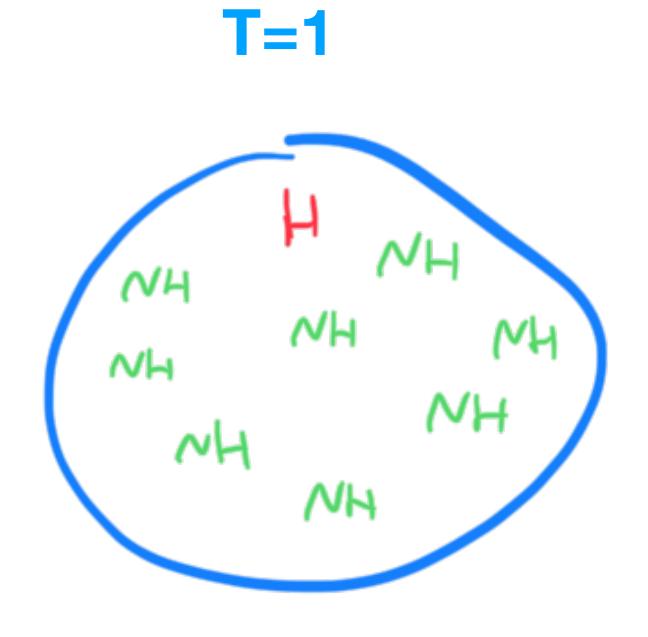
$$P(Y = 1 | T = 1, X = 1) = 0.75$$

$$P(Y = 1 | T = 0, X = 1) = 0.5$$

$$P(Y = 1 | T = 0, \mathbf{X} = 0) = 0.6$$

$$P(Y = 1 | T = 1, \mathbf{X} = 0) = 0.9$$







### Matching and IPW Jupyter notebook

```
treatment_group_x_0 = treatment_group[treatment_group["x"]==0]
treatment_group_x_1 = treatment_group[control_group["x"]==0]
control_group_x_0 = control_group[control_group["x"]==0]
control_group_x_1 = control_group[control_group["x"]==1]

print("Number of people with X=0 in treatment: ", len(treatment_group_x_0)," and in control: ", len(control_group_x_0))
print("Number of people with X=1 in treatment: ", len(treatment_group_x_1)," and in control: ", len(control_group_x_1))

Number of people with X=0 in treatment: 3159 and in control: 338
Number of people with X=1 in treatment: 157 and in control: 1346
```

$$P(X = 1) = 0.3$$
  $P(T = 1 | X = 1) = 0.1$   $P(T = 1 | X = 0) = 0.9$ 

```
min_number_x1 = min(len(treatment_group_x_1), len(control_group_x_1))
balanced_treatment_x_1 = treatment_group_x_1[0:min_number_x1]
balanced_control_x_1 = control_group_x_1[0:min_number_x1]
print("After balancing: number of people with X=1 in treatment: ", len(balanced_treatment_x_1)," and in control: ", len(balanced_control_x_1))
After balancing: number of people with X=0 in treatment: 338 and in control: 338
After balancing: number of people with X=1 in treatment: 157 and in control: 157
```

# Next class: Inverse probability weighting (IPW)

- Idea: rather than match, reweight (downweight or upweight) observations
- Inverse probability (of treatment) weighting: weight by inverse of probability of treatment received:
  - For treated T=1: weight by the inverse of  $\pi=P(T=1|\mathbf{X})$
  - For untreated T=0: weight by the inverse of  $1-\pi=P(T=0|\mathbf{X})$



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$$\hat{\mathbb{E}}(Y(t=1)) = \frac{1}{n} \sum_{i=1}^{n} Y_i \cdot 1\{T=1\} \cdot \frac{1}{P(T=1 \mid X_i)}$$

$$\hat{\mathbb{E}}(Y(t=0)) = \frac{1}{n} \sum_{i=1}^{n} Y_i \cdot 1\{T=0\} \cdot \frac{1}{P(T=0 \mid X_i)}$$



#### Questions? JELLY BEANS





WE FOUND NO

LINK BETWEEN

WE FOUND NO

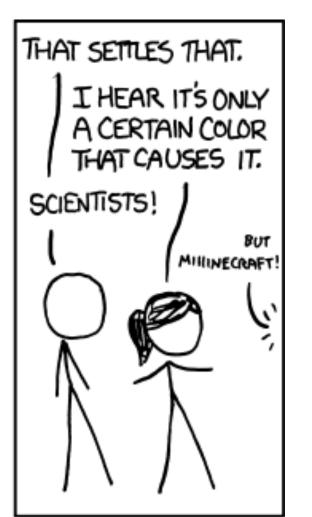
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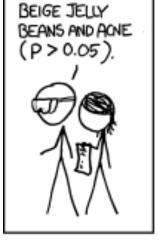
TURQUOISE JELLY

BEANS AND ACNE

(P > 0.05)

PINK JELLY





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GREY JELLY

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BEANS AND ACNE

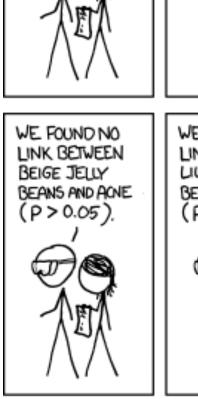
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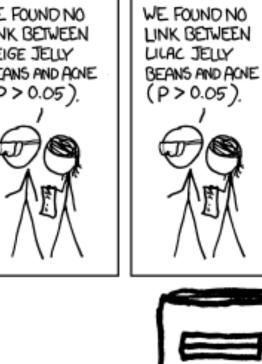
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(P>0.05).

TAN JELLY







BLACK JELLY

BEANS AND ACNE



WE FOUND A

LINK BETWEEN



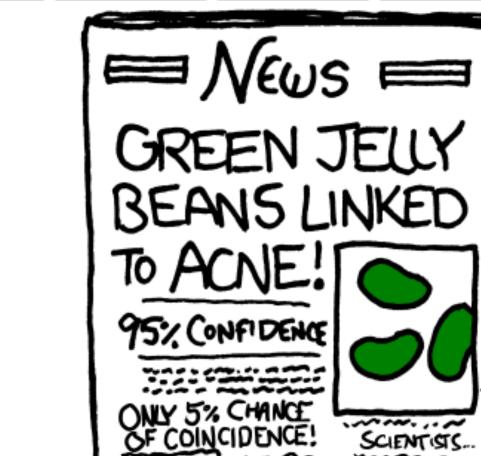






PEACH JELLY







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SALMON JELLY

(P>0.05)

BEANS AND ACNE



WE FOUND NO

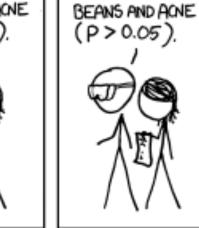
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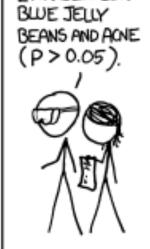
BEANS AND ACNE

(P > 0.05)

RED JELLY

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WE FOUND NO

(P>0.05).

WE FOUND NO

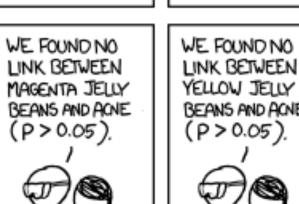
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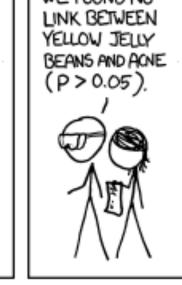


WE FOUND NO

LINK BETWEEN

TEAL JELLY





https://xkcd.com/882/