

Research Design

03: elements of causal inference; experiments

goals of data science research

goals of data science research

descriptive

associative

predictive

explicative

goals of data science research, explicative

What is causation?

goals of data science research, what is causation?

CAUSE, N. | That which produces an effect; that which gives rise to any action, phenomenon, or condition.
Cause and *effect* are correlative terms.

How can we learn or test
if thing A causes thing B?

causal inference and experiments

causal inference, the potential outcomes approach

Causal effects involve the comparison of the outcome actually observed with other potential outcomes that could have been observed had the treatment taken on a different level, but that are not, in fact, observed. Causal inference is therefore fundamentally a missing data problem.

— Imbens & Rubin

causal inference, which concerns what *would happen* to an outcome y as a result of a treatment, intervention, or exposure z , given pre-treatment information x .

— Gelman, Hill, Ventari

What's a *treatment*? Why can't we observe these *potential* outcomes, these *missing* data?

the potential outcomes approach, a metaphor for missing outcomes

The Road Not Taken

Two roads diverged in a yellow wood,
And *sorry I could not travel both*
And be one traveler, long I stood
And looked down one as far as I could
To where it bent in the undergrowth;

Then took the other, as just as fair,
And having perhaps the better claim,
Because it was grassy and wanted wear;
Though as for that the passing there
Had worn them really about the same,

And both that morning equally lay
In leaves no step had trodden black.
Oh, I kept the first for another day!
Yet knowing how way leads on to way,
I doubted if I should ever come back.

I shall be telling this with a sigh
Somewhere ages and ages hence:
Two roads diverged in a wood, and I—
I took the one less traveled by,
And that has made all the difference.

— Robert Frost

the potential outcomes approach, common notation for causal inference in experiments

i , an experimental unit

$z = 0$, the control group

$z = 1$, the treatment group

y_i^0 , the potential outcome of unit i if no treatment

y_i^1 , the potential outcome of unit i if treatment

$y_i = y_i^0 \cdot (1 - z_i) + y_i^1 \cdot z_i$, the observed outcome of unit i

$\tau_i = y_i^1 - y_i^0$, causal effect for unit i

$\hat{\tau} = \frac{1}{n} \sum_{i=1}^n (y_i^1 - y_i^0)$, sample average treatment effect

The fundamental problem of causal inference: we can never observe both y_i^0 and y_i^1 . And we can only attribute an **average treatment effect $\hat{\tau}$** to a unit if we assume that effects are constant across units.

$\bar{\tau} = \frac{1}{N} \sum_{i=1}^N (y_i^1 - y_i^0)$, population average treatment effect

the potential outcomes approach, hypothetical data — *balanced* treatment and control groups?

Unit i	Female, x_{1i}	Age, x_{2i}	Treatment, z_i	Potential outcomes		Observed outcome, y_i
				if $z_i = 0$, y_i^0	if $z_i = 1$, y_i^1	
Audrey	1	40	0	140	135	140
Anna	1	40	0	140	135	140
Bob	0	50	0	150	140	150
Bill	0	50	0	150	140	150
Caitlin	1	60	1	160	155	155
Cara	1	60	1	160	155	155
Dave	0	70	1	170	160	160
Doug	0	70	1	170	160	160

Of note, with just 8 units, split equally between treatment and control groups, there are

$$\binom{n+k-1}{k} = 330$$

unique possible experiments!

Do you think this treatment assignment *balances* the treatment and control groups, or is it *biased*?

What's the **sample average treatment effect** $\hat{\tau}$ for this particular treatment assignment?

How does $\hat{\tau}$ compare with the *unknown true average treatment effect*?

Now re-assign the units to treatment and control groups *randomly* where $z \perp y^0, y^1$ and repeat. What do you get?

```
set.seed(3)
z <- sample(c(0,0,0,0,1,1,1,1), 8)
```

the potential outcomes approach, properties of randomization

```
d <-
  read.table(text = '
Unit      Female Age z yi0 yi1
Audrey    1      40  0 140 135
Anna      1      40  0 140 135
Bob       0      50  0 150 140
Bill      0      50  0 150 140
Caitlin   1      60  1 160 155
Cara      1      60  1 160 155
Dave      0      70  1 170 160
Doug      0      70  1 170 160
', header = TRUE)

tau_tru <- with(d, mean(yi1 - yi0) )

d$yi      <- with(d, yi0 * (1 - z) + yi1 * z)
y1        <- with(d, mean(yi[z == 1]) )
y0        <- with(d, mean(yi[z == 0]) )
tau_hat   <- y1 - y0

set.seed(123)

d$z       <- sample(c(0, 0, 0, 0, 1, 1, 1, 1), 8)
d$yi      <- with(d, yi0 * (1 - z) + yi1 * z)
y1        <- with(d, mean(yi[z == 1]) )
y0        <- with(d, mean(yi[z == 0]) )
tau_hat   <- y1 - y0
```

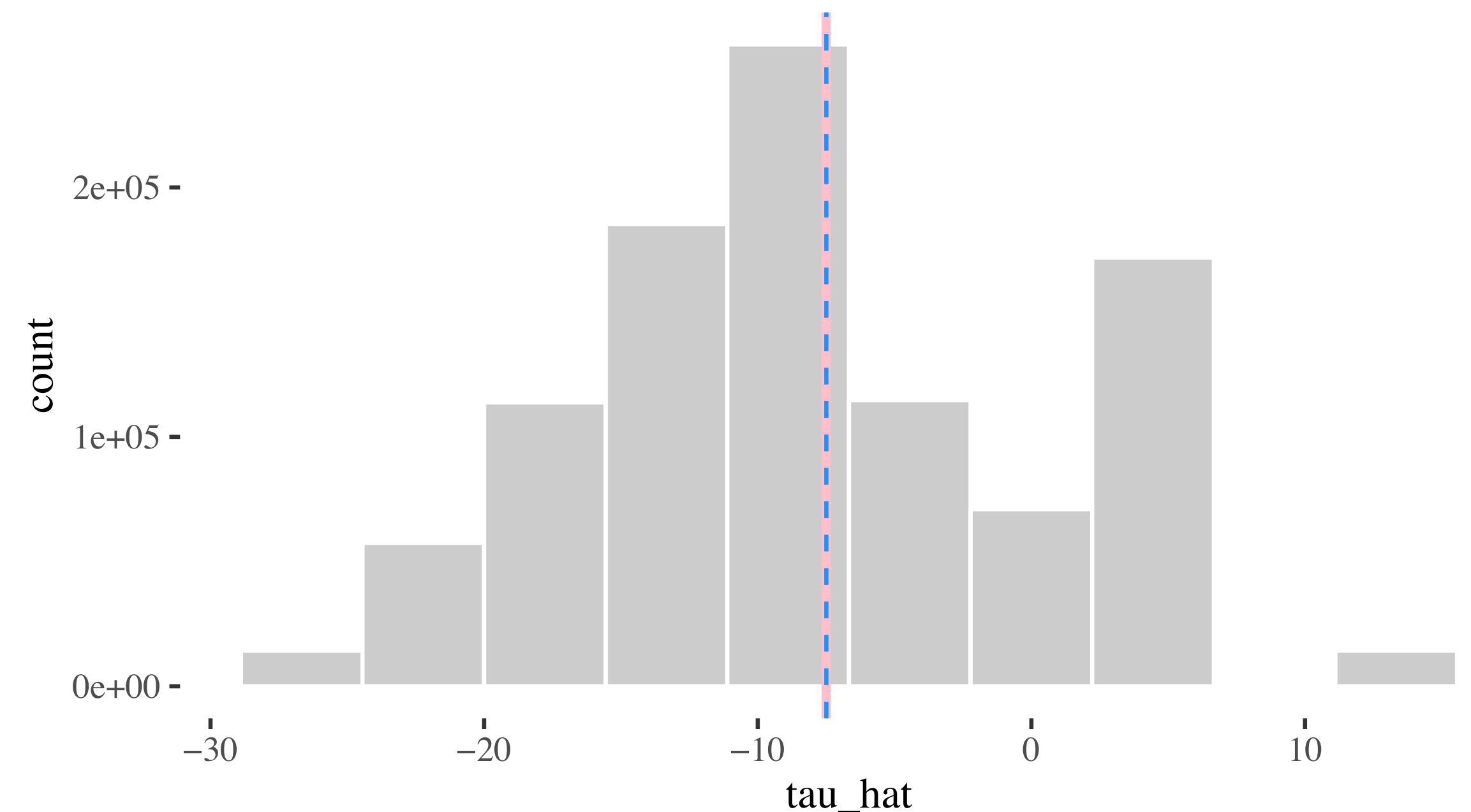
No *single* randomized experiment guarantees that $\hat{\tau}$ will be close to the *unknown true average treatment effect*.

Try experimenting with different seeds in this code, and re-run to see how individual $\hat{\tau}$ is affected by the sample.

the potential outcomes approach, properties of randomization

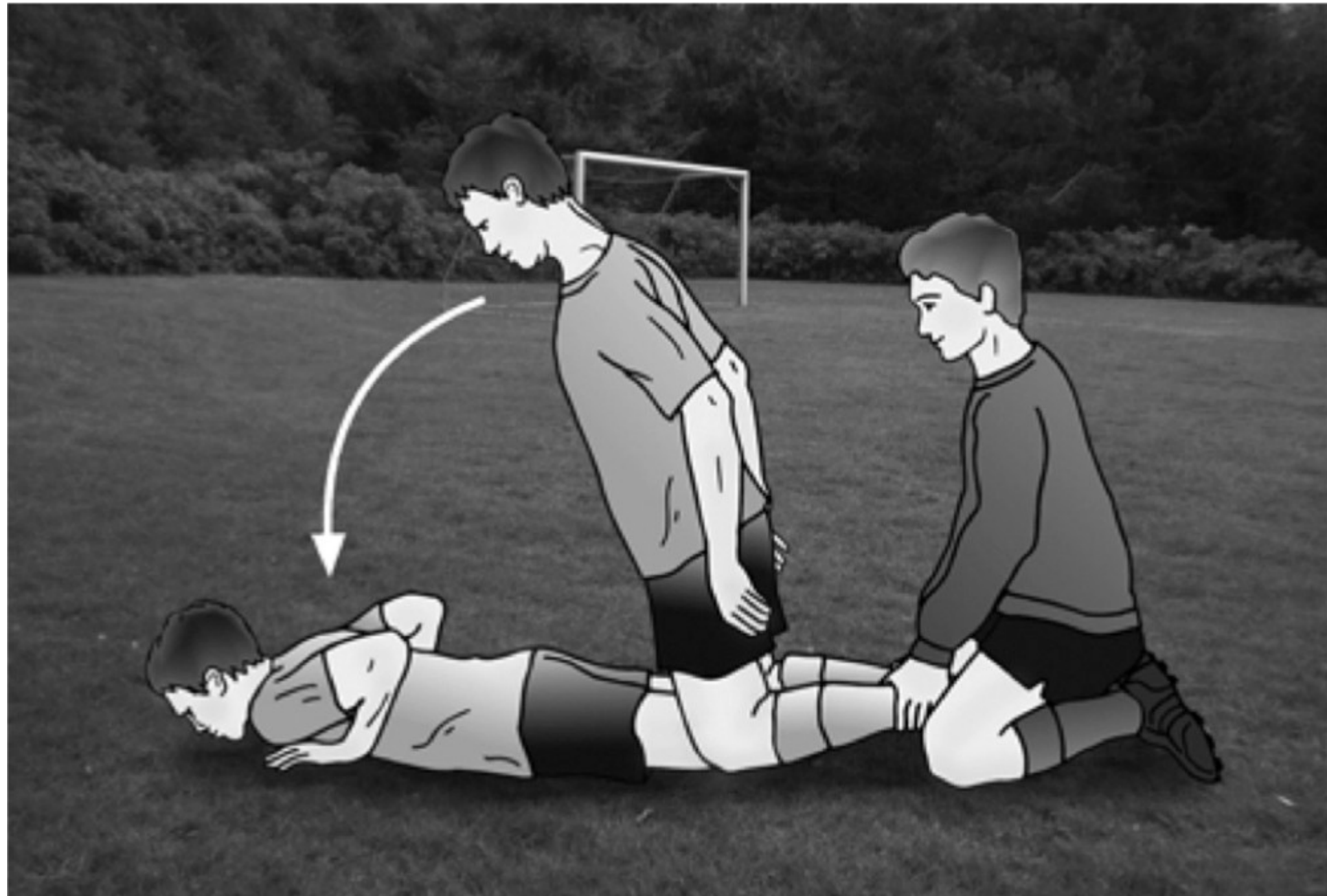
```
sim_experiment <- function(d) {  
  d$z <- sample(c(0, 0, 0, 0, 1, 1, 1, 1), 8)  
  y1 <- with(d, mean(yi1[z == 1]) )  
  y0 <- with(d, mean(yi0[z == 0]) )  
  
  return(y1 - y0)  
}  
  
tau_hat <- replicate( 1e6, sim_experiment(d) )  
  
library(ggplot2)  
library(ggthemes)  
  
ggplot() +  
  theme_tufte() +  
  geom_histogram(aes(tau_hat),  
                 bins = 10,  
                 fill = "lightgray",  
                 color = "white") +  
  geom_vline(aes(xintercept = tau_tru),  
            color = "pink",  
            lwd = 1.1) +  
  geom_vline(aes(xintercept = mean(tau_hat)),  
            color = "dodgerblue",  
            linetype = "dashed")  
  
E_tau_hat <- mean(tau_hat)
```

But randomly assigning units to treatment and control groups ensures that there are *no differences in expectation in the distribution* of potential outcomes between groups receiving different treatments — it's an *unbiased* estimator. In these simulations, $\mathbb{E}(\hat{\tau}) = -7.497 \simeq -7.5$



By collecting *more units*, we can improve balance in single experiments, and by collecting *pre-treatment* information, we can *adjust for imbalances* — techniques we cover later.

review of a published, randomized controlled experiment



Purpose?

Null hypothesis?

Alternative hypothesis?

Experimental design?

Results?

introducing your group projects

References

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