## TESTING THE VALIDITY OF THE COUNTERFACTUAL

Jesse Lecy

## CORE CONCEPTS



The Selection Problem



**Tests for Group Equivalence** 

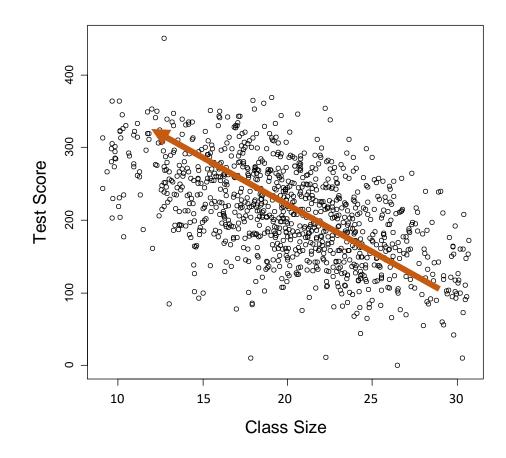
Omnibus Test
Bonferroni Correction



**Tests for Nonrandom Attrition** 

## NATURE GIVES US CORRELATIONS: THE SELECTION PROBLEM IN EVALUATION RESEARCH

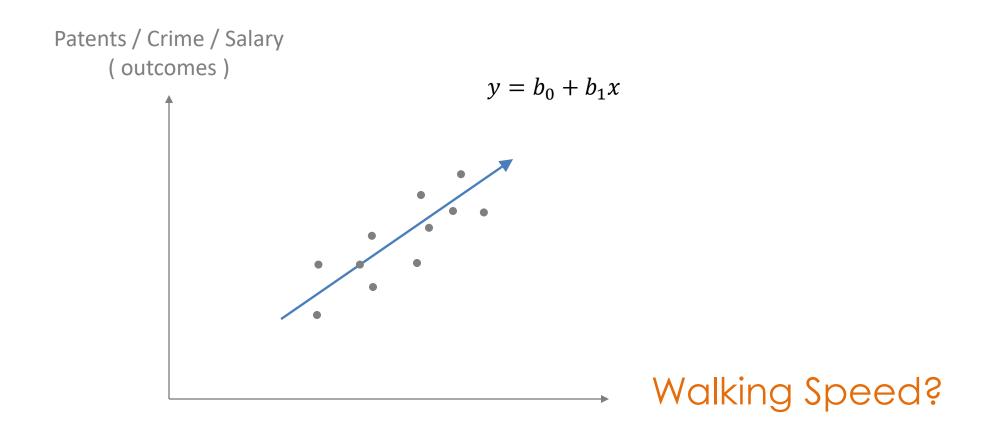
If we reduce classroom size we should see student performance improve. Right?



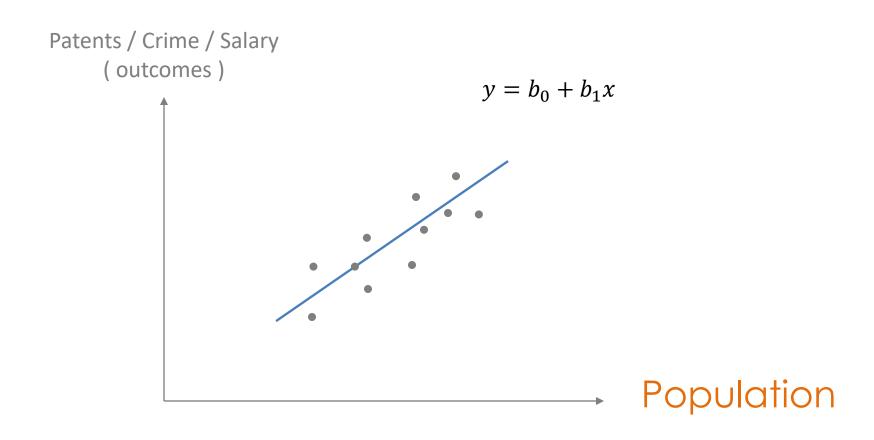
Listen: 4:30 – 15:30

http://www.radiolab.org/2010/oct/08/its-alive/



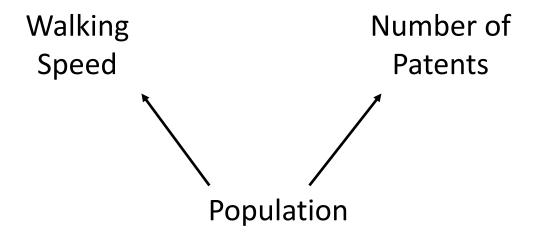


If you increase the WALKING SPEED of a city, would you increase the number of patents produced?



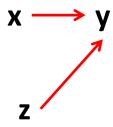
If you increase the POPULATION of a city, would you increase the number of patents produced?

## The problem with correlations:

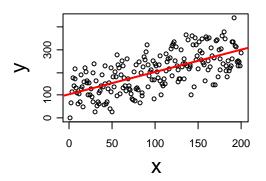


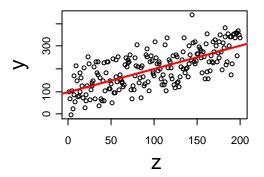
## NATURE GIVES US CORRELATIONS

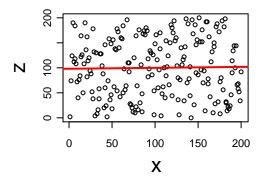
### Example #1



X and Z both impact Y

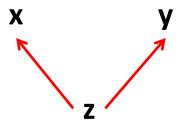




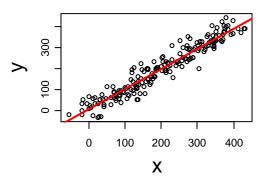


no correlation

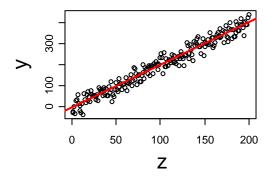
#### Example #2

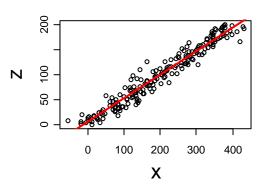


Z impacts both X and Y

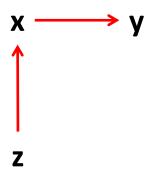


No causal relationship but high correlation

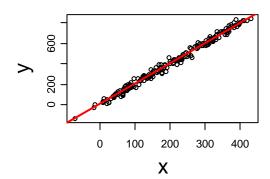


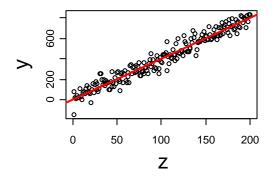


### Example #3

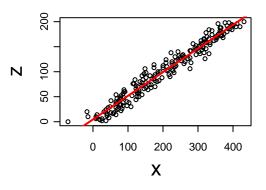


### Causal chain

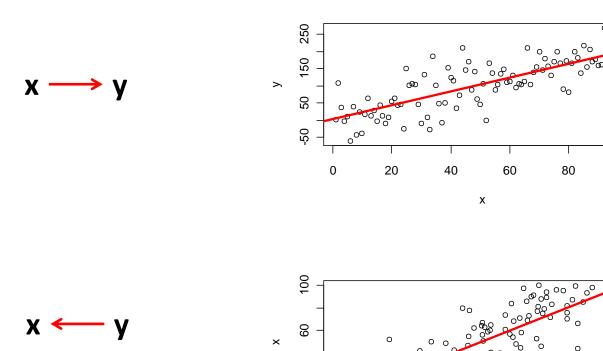




no direct causal relationship but highly correlated



## Reverse causality



$$Y = b_0 + b_1 X + e$$

Both models are highly significant, how do we know which one is the causal relationship?

100

100

у

200

250

$$X = b_0 + b_1 Y + e$$

#### THE PROGRAM EVALUATION FRAMEWORK

$$Heartrate = b_0 + b_1 \cdot Caffeine + \varepsilon$$

## Treatment Effect = typical dosage x slope b<sub>1</sub>

Slopes represent change in Y as a result of a one-unit change in X

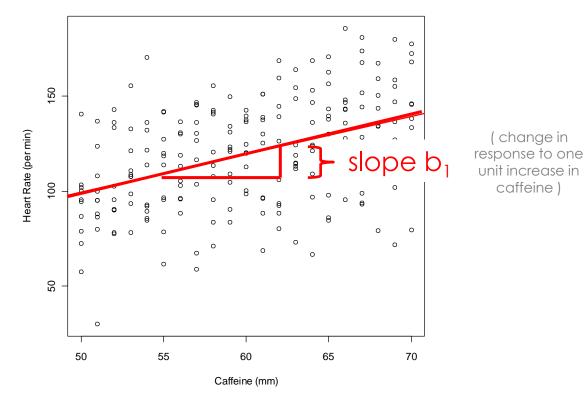
Rarely is one-unit of X meaningful, so be sure to translate the slope to an appropriate dosage to generate a meaningful impact measure

If the treatment is one espresso:

Espresso contains ~ 50mg caffeine Heart rate increase = (50)(b1)

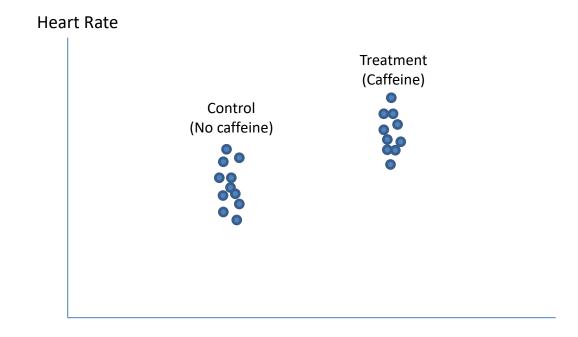
Effect Size = (dosage)(b1)

#### **Dosage and Response**

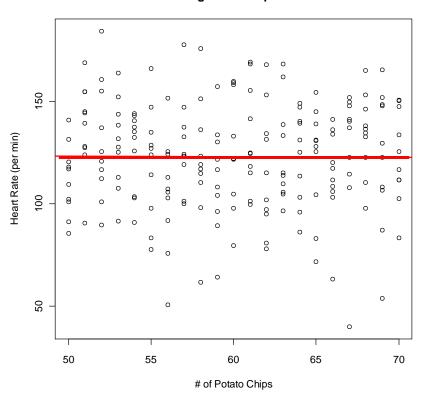


## DISCRETE TREATMENT CASE: MODEL IS THE SAME EXCEPT CAFFEINE IS A DUMMY NOT A LEVEL

$$Heartrate = b_0 + b_1 \cdot Caffeine + \varepsilon$$



#### **Dosage and Response**



## No effect of treatment on outcome

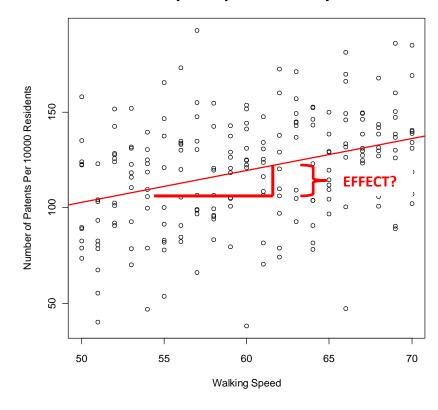
## HOW DO WE KNOW WHEN THE INTERPRETATION IS CAUSAL?

When is b<sub>1</sub> an impact, and when is it just a relationship in data?

#### **Dosage and Response**

## 150 Heart Rate (per min) 0 50 55 60 65 70 Caffeine (mm)

#### **City Density and Productivity**



## THE SELECTION PROBLEM IN EVALUATION RESEARCH

## Microfinance example of bias from selection INTO a study group

#### Number of each "type" of person in the study

(latent traits can be hard to observe thus hard to control for in a study)

	NOT Entrepreneurial	Entrepreneurial
No Loan	30	15
Takes a Loan	20	35

#### Average weekly income AFTER loan period

	NOT Entrepreneurial	Entrepreneur
No Loan	\$10	\$20
Takes a Loan	\$10	\$20
Income not impact by the loan	cted	

You are more likely to take a loan if you know you are good at business

Takes Loan?

$$NO: \quad \frac{30 \cdot \$10 + 15 \cdot \$20}{45} = \$13.33$$

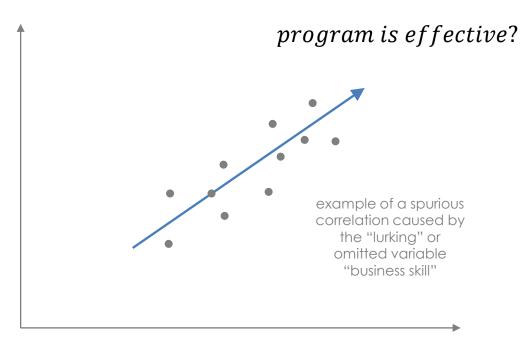
$$YES: \quad \frac{20 \cdot \$10 + 35 \cdot \$20}{55} = \$16.37$$

The loan appears to have an impact!

Even though we know it didn't.

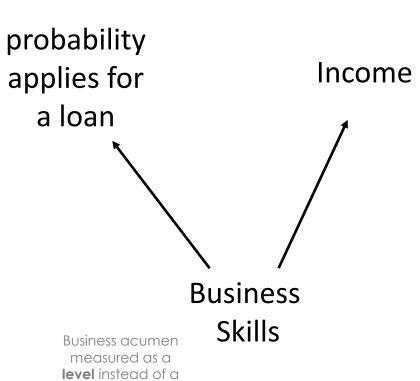
## CORRELATION VERSION OF LAST EXAMPLE:

#### Income



Probability of Taking a Loan

### Selection mechanism



type here

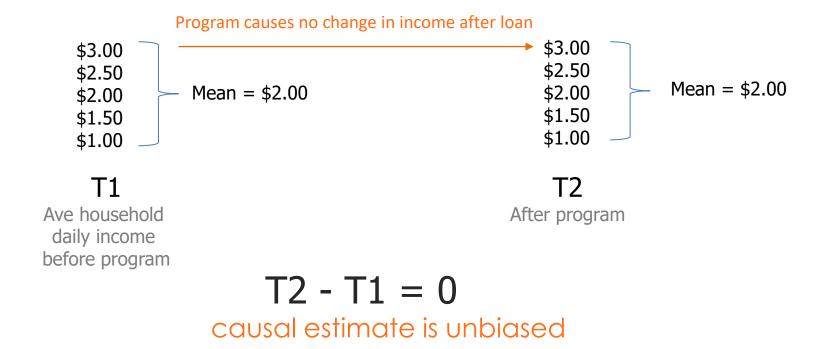
## "Selection" Problem

Those that participate in the program are different from those that do not participate.

This is the biggest problem in impact evaluation!

## Microfinance example of bias from selection OUT OF a study group

#### Reflexive design



## Random Attrition Example



$$T2 - T1 = 0$$

Impact study accurately represents program effects

Program is not determined to be effective (no change)

(impact estimate is unbiased)

## Non-Random Attrition Example

T2 - T1 > 0

We over-estimate program effects
Program appears to be effective

(impact estimate BIASED)

## Non-Random Attrition Example

T2 - T1 < 0

We under-estimate program effects
Program appears to harm families

(impact estimate is BIASED)

## What if the program has impact?

#### Program is effective: increases income by \$1



True impact = T2-T1 = \$1.00

#### Random Attrition

#### 

## T2-T1 = \$1.00 (UNBIASED)

#### Non-Random Attrition A

$$T2-T1 = $1.50$$

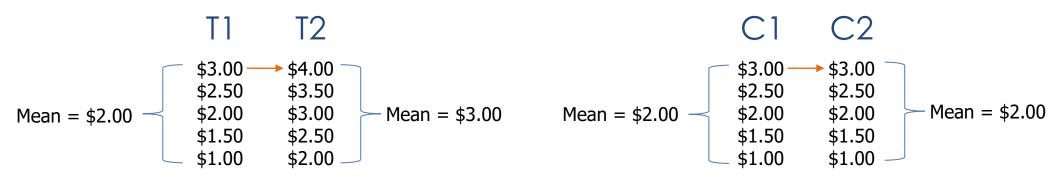
(too big: BIASED)

#### Non-Random Attrition B

$$T2-T1 = $0.50$$

(too small: BIASED)

### Attrition with two study groups



Total gains: T2-T1 = \$1.00

No secular trends: C2-C1 = 0

True impact: (T2-T1) - (C2-C1) = \$1.00

#### Random Attrition

T1	T2	C1	C2
\$3.00	\$4.00	\$3.00	\$3.00
\$2.50	<del>\$3.50</del>	\$2.50	<del>\$2.50</del>
\$2.00	\$3.00	\$2.00	\$2.00
\$1.50	<del>\$2.50</del>	\$1.50	<del>\$1.50</del>
\$1.00	\$2.00	\$1.00	\$1.00

#### (3.00-2.00)-(2.00-2.00)=\$1.00 UNBIASED

#### Non-Random Attrition A

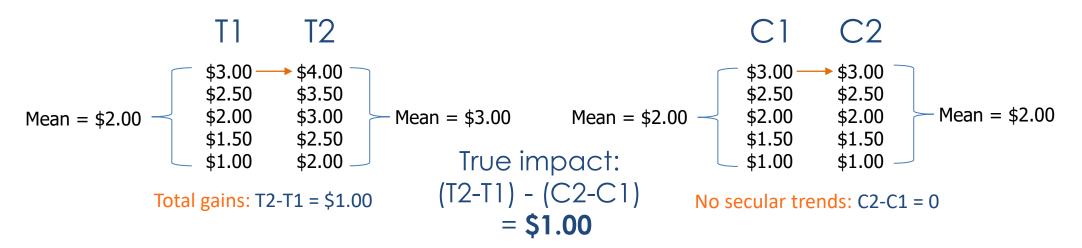
T1	T2	C1	C2
\$3.00	\$4.00	\$3.00	\$3.00
\$2.50	\$3.50	\$2.50	\$2.50
\$2.00	\$3.00	\$2.00	\$2.00
\$1.50	<del>\$2.50</del>	\$1.50	<del>\$1.50</del>
\$1.00	<del>\$2.00</del>	\$1.00	<del>\$1.00</del>

#### Non-Random Attrition B

T1	T2	C1	C2
\$3.00	<del>\$4.00</del>	\$3.00	<del>\$3.00</del>
\$2.50	<del>\$3.50</del>	\$2.50	<del>\$2.50</del>
\$2.00	\$3.00	\$2.00	\$2.00
\$1.50	\$2.50	\$1.50	\$1.50
\$1.00	\$2.00	\$1.00	\$1.00

$$(3.50-2.00)-(2.50-2.00)=$$
 \$1.00  $(2.50-2.00)-(2.00-1.50)=$  \$1.00 UNBIASED

### Attrition with two study groups



#### Random Attrition

T1	T2	C1	C2
\$3.00	\$4.00	\$3.00	\$3.00
\$2.50	<del>\$3.50</del>	\$2.50	<del>\$2.50</del>
\$2.00	\$3.00	\$2.00	\$2.00
\$1.50	<del>\$2.50</del>	\$1.50	<del>\$1.50</del>
\$1.00	\$2.00	\$1.00	\$1.00

$$T2-T1 = (3.00-2.00) = $1.00$$
  
UNBIASED

#### Non-Random Attrition A

$$T2-T1 = (3.50-2.00) = $1.50$$
BIASED

#### Non-Random Attrition B

T1	T2	C1	C2
\$3.00	<del>\$4.00</del>	\$3.00	<del>\$3.00</del>
\$2.50	<del>\$3.50</del>	\$2.50	<del>\$2.50</del>
\$2.00	\$3.00	\$2.00	\$2.00
\$1.50	\$2.50	\$1.50	\$1.50
\$1.00	\$2.00	\$1.00	\$1.00
T2-T1 = (2.50-2.00) = \$0.50			

12-11 = (2.50-2.00) = \$0.50BIASED

Note that we have no trend in the data so we can usually substitute the reflexive and diff-in-diff estimators (T2-T1 = [T2-T1]-[C2-C1]), but when non-random attrition is present we cannot

### Attrition with two study groups

#### Random Attrition

## T1 T2 C1 C2 \$3.00 \$4.00 \$3.00 \$3.00 \$2.50 \$3.50 \$2.50 \$2.50 \$2.00 \$3.00 \$2.00 \$2.00 \$1.50 \$2.50 \$1.50 \$1.50 \$1.00 \$2.00 \$1.00 \$1.00

(3.00-2.00)-(2.00-2.00)=\$1.00 UNBIASED

#### Non-Random Attrition A

(3.50-2.00)-(2.50-2.00)=\$1.00

#### Non-Random Attrition B

(2.50-2.00)-(2.00-1.50)=\$1.00 UNBIASED

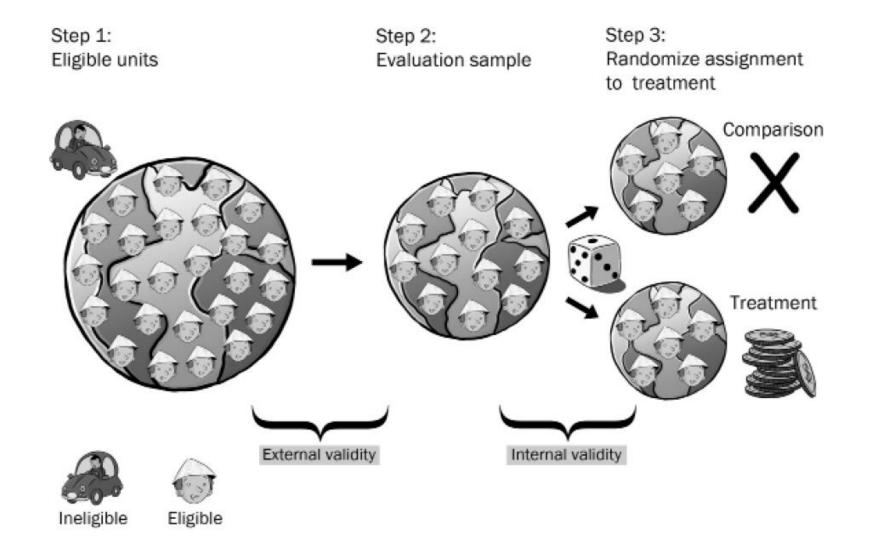
Note that this only works when the attrition process is the same in both groups (random cases, mostly high performers, or mostly low performers exiting from the treatment and study groups at the same rate)

# RANDOMIZED CONTROL TRIALS (RCT'S): THE "GOLD STANDARD" FOR INTERNAL VALIDITY

Gertler, P. J., Martinez, S., Premand, P., Rawlings, L. B., & Vermeersch, C. M. (2016). *Impact evaluation in practice*. The World Bank.

>> Chapter 4. Randomized Selection Methods

Figure 4.3 Steps in Randomized Assignment to Treatment



## Our counterfactual framework is valid / robust when the groups only DIFFER BY THE TREATMENT but are OTHERWISE "IDENTICAL"

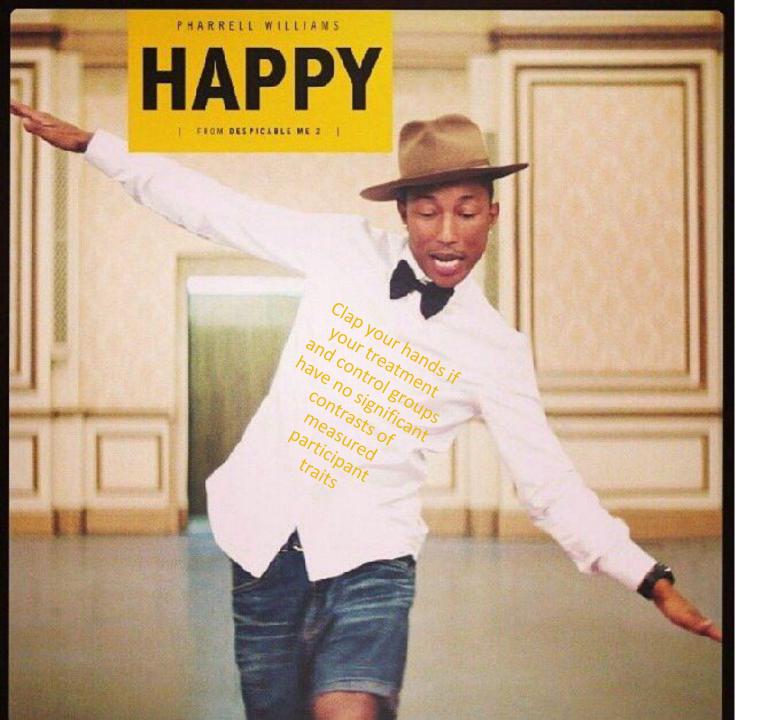


When true, we can interpret the differences in group outcomes after the treatment period to be caused by the treatment

#### How do we test the criteria:

#### groups only OTHERWISE "IDENTICAL" ???





## "HAPPY" RANDOMIZATION

### MATHEMATICALLY: ALWAYS !!!

weights will never be exactly identical

so what do we mean by "different"?

### STATISTICALLY:

### TEST OF GROUP MEANS

Use a t-test and select a level of confidence that we are comfortable with

### STATISTICALLY:

### TEST OF GROUP MEANS

What does alpha=0.05 mean?

[--- 95% confidence interval ---]

5 times out of 100 the two samples are drawn from the same population (or weight distribution), but we will still consider them to be different.

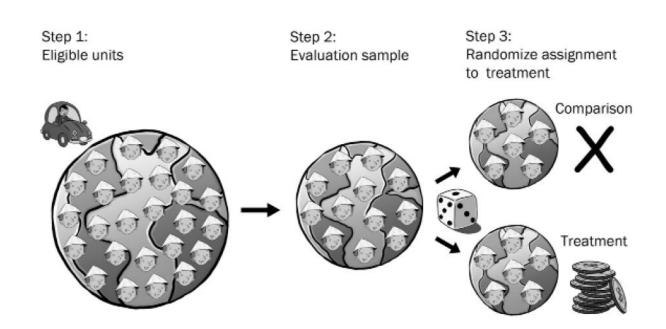
### STATISTICALLY:

### TEST OF GROUP MEANS

What does alpha=0.05 mean?

[--- 95% confidence interval --- ]

### How often will randomization "fail"?



Unhappy randomization is not failed randomization (process applied improperly), rather just bad luck of the draw

By definition of a test at a 95% confidence level, each measured characteristic like weight will differ

5 out of 100 times



"Unhappy Randomization"



Table 4.1 Case 3—Balance between Treatment and Comparison Villages at Baseline

Household	Treatment villages	Comparison villages		_
characteristics	(N = 2964)	(N = 2664)	Difference	<i>t</i> -stat
Health expenditures (\$ yearly per capita)	14.48	14.57	-0.09	-0.39
Head of household's age (years)	"cont 41.6 <b>←</b>	rasts" → 42.3	-0.7	-1.2
Spouse's age (years)	36.8	36.8	0.0	0.38
Head of household's education (years)	2.9	2.8	0.1	2.16*
Spouse's education (years)	2.7	2.6	0.1	0.006
Head of household is female = 1	0.07	0.07	-0.0	-0.66
Indigenous = 1	0.42	0.42	0.0	0.21
Number of household members	5.7	5.7	0.0	1.21
Has bathroom = 1	0.57	0.56	0.01	1.04
Hectares of land	1.67	1.71	-0.04	-1.35
Distance to hospital (km)	109	106	3	1.02

Source: Authors' calculation.

The most important table in every study: comparisons of treatment and control group characteristics

For the counterfactual to be <u>valid</u>, the groups can ONLY differ by the treatment, not by any measured traits.

— Is this problematic?

What is the appropriate test for "identical" or equivalent groups?

We should observe no differences in measured traits.

Assume a 95% confidence interval.

<sup>\*</sup> Significant at the 5 percent level.

### Bonferroni Correction:

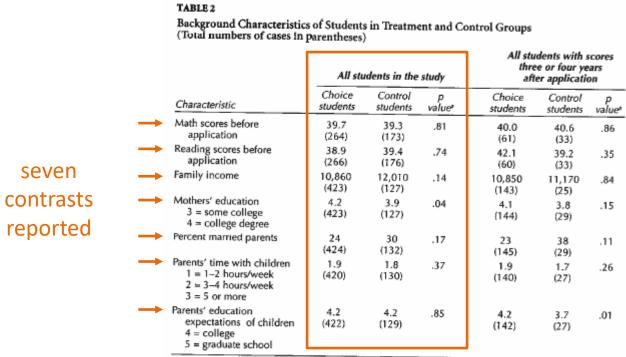
When we want to be 95% confident that two groups are the same, and we can measure those groups using a set of contrasts, then our decision rule is no longer to reject the null (that the groups are the same) if the p-value < 0.05. A "contrast" is a comparison of means of any measured characteristic between two groups.

If we have a 5% chance of observing a p-value of less than 0.05 for each contrast, then the probability of observing at least one contrast with a p-value that small is greater than 5%! It is actually n\*0.05 (minus prob of observing multiple < 0.05 at same time) where n is the number of contrasts.

So if we want to be 95% confident that the groups are different (not just the contrasts), we have to adjust our decision rule to  $\alpha/n$ .

For example, if we have 10 contrasts, then our decision rule is now 0.05/10, or 0.005. The p-value of at least one contrast must be below 0.005 for us to conclude that the groups are different.

### Test for Group Equivalence



a. The tests of significance are suggestive of the equivalence of the two groups. Technically, tests of significance should be done at each point of random assignment, but the number of cases at each point is too few for such tests to be meaningful.

Smallest p-value in table

0.04 > 0.0071

New alpha = (0.05 / 7) = 0.0071

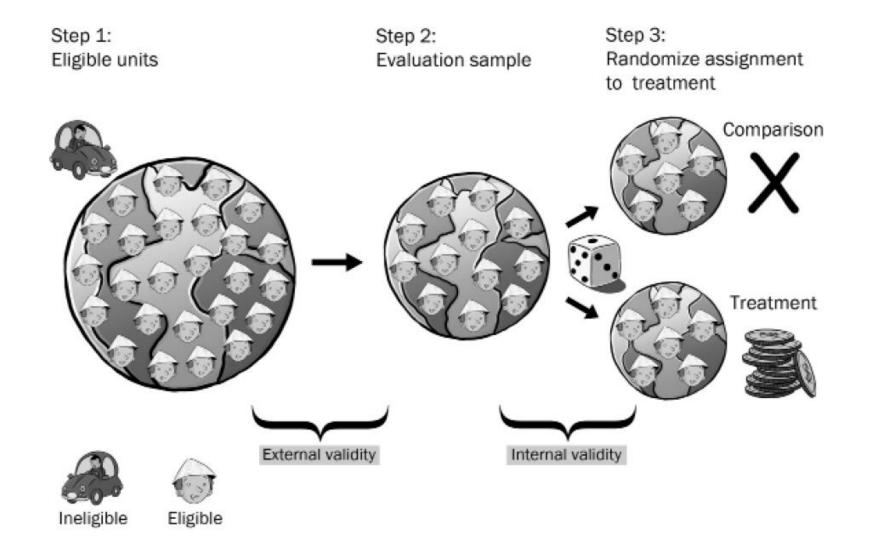
Do not reject :: Groups are equivalent

### RCT versus Natural Experiments:

- 1. RCT assumes complete control over the assignment process
- 2. Natural Experiments often utilize randomization:
  - → Charter School lotteries
  - → Vietnam draft
- 3. Quasi-Experimental techniques can use other methods to create group equivalence (for example, matching)

## **ATTRITION**

Figure 4.3 Steps in Randomized Assignment to Treatment



### Tests for Selection-Into Study Group

TABLE 2

Background Characteristics of Students in Treatment and Control Groups (Total numbers of cases in parentheses)

	All students in the study			All students with scores three or four years after application		
Characteristic	Choice students	Control students	p value*	Choice students	Control students	p value
Math scores before application	39.7 (264)	39.3 (173)	.81	40.0 (61)	40.6 (33)	.86
Reading scores before application	38.9 (266)	39.4 (176)	.74	42.1 (60)	39.2	.35
Family income	10,860 (423)	12,010 (127)	.14	10,850 (143)	11,170 (25)	.84
Mothers' education 3 = some college 4 = college degree	4.2 (423)	3.9 (127)	.04	4.1 (144)	3.8 (29)	.15
Percent married parents	24 (424)	30 (132)	.17	23 (145)	38 (29)	.11
Parents' time with children 1 = 1-2 hours/week 2 = 3-4 hours/week 3 = 5 or more	1.9 (420)	1.8 (130)	.37	1.9 (140)	1.7 (27)	.26
Parents' education expectations of children 4 = college 5 = graduate school	4.2 (422)	4.2 (129)	.85	4.2 (142)	3.7 (27)	.01

a. The tests of significance are suggestive of the equivalence of the two groups. Technically, tests of significance should be done at each point of random assignment, but the number of cases at each point is too few for such tests to be meaningful.

#### **Non-Random Attrition**

If the people that leave a program or study are different than those that stay, the calculation of effects will be biased.

#### The Fix:

Examine characteristics of those that stay versus those that leave.

# Non-Random attrition tests for selection OUT of the study group

TABLE 2

Background Characteristics of Students in Treatment and Control Groups
(Total numbers of cases in parentheses)

	All students in the study			All students with scores three or four years after application		
Characteristic	Choice students	Control students	p value*	Choice students	Control students	p value*
Math scores before application	39.7 (264)	39.3 (173)	.81	40.0 (61)	40.6 (33)	.86
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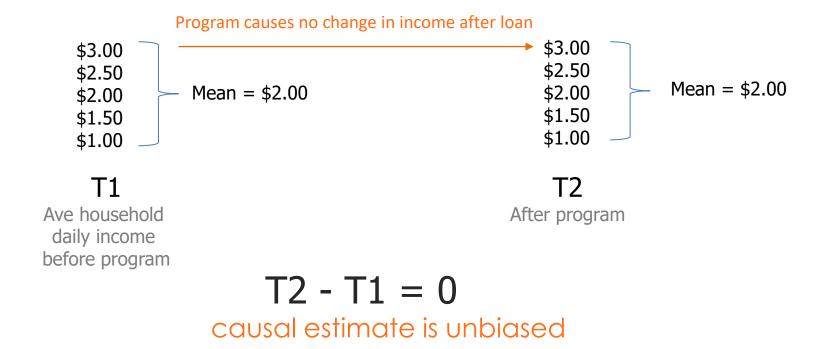
Do group traits differ after attrition occurs?

Attrition is natural, the question is whether it is random (will not change the groups) or non-random (will change the groups)

Can also be tested by comparing traits of those that stay to those that leave.

# Microfinance example of bias from selection OUT OF a study group

### Reflexive design



### Random Attrition Example



$$T2 - T1 = 0$$

Impact study accurately represents program effects

Program is not determined to be effective (no change)

### Non-Random Attrition Example

 $T2 - T1 \neq 0$ 

We over-estimate program effects

Program appears to be effective

### Non-Random Attrition Example

 $T2 - T1 \neq 0$ 

We under-estimate program effects
Program appears to harm families

### WHAT DO WE MEAN BY "TREATED"?

HOW NON-COMPLIANCE
CHANGES OUR
MEASURE OF
EFFECTS

## One of The Physicists Behind The Higgs Boson Has Made an Algorithm to Replace The Pill It's up to 99.5% effective at stopping pregnancy.

Those more effective methods are ones that don't require people to remember to take a pill, put on a condom, or record their temperature daily, such as intrauterine contraception or implants.

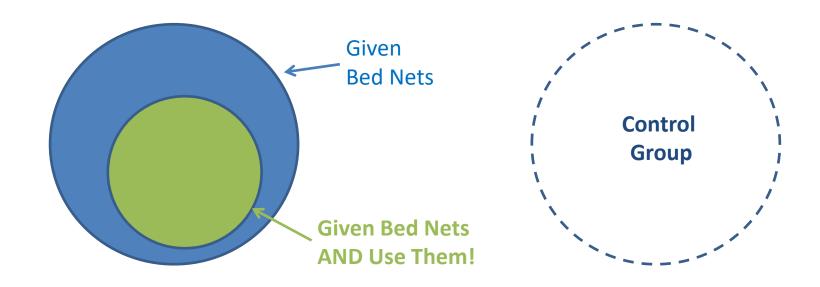
That's because human error can mess with things quite a lot. In fact, the UK's National Health Service (NHS) explained that when the app was used perfectly all the time, only five out of every 1,000 women would fall pregnant every year - a rate slightly better than the pill (99.5 percent).

But for "typical use" - where the app isn't used entirely correctly every day - it's more likely that seven out of every 100 women would experience accidental pregnancies, which is around 93 percent efficiency.

What is the TRUE effectiveness of the app?

99.5 percent, or 93 percent?

### Estimation of the counter-factual:

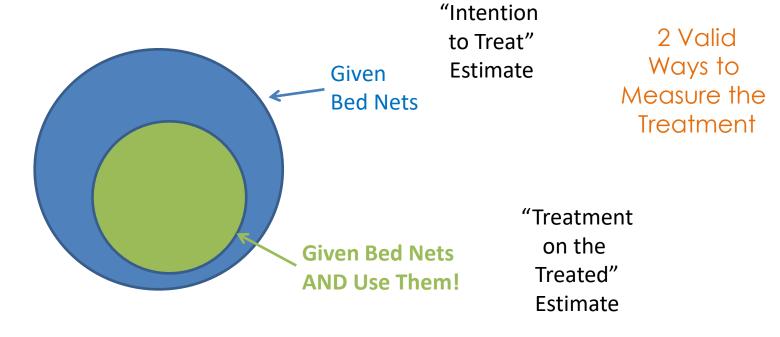


Program Effect = T2 - C2

Is Group T (for treatment) those that are GIVEN bed nets, or those that USE them?

### **TERMINOLOGY:**

- "Average" treatment effects
  - Treatment on the Treated (TOT) Effects
  - Intention to Treat (ITT) Effects

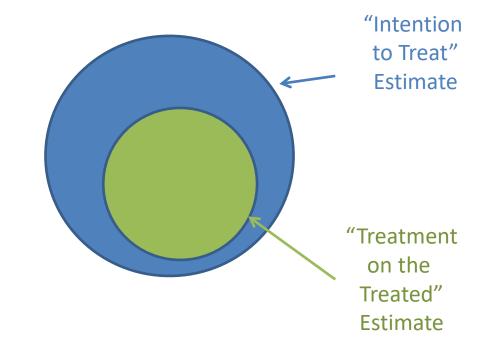


### WHY DO WE NEED TWO MEASURES?

THE TOT MEASURE is the optimistic or best-case scenario. It tells us how effective the program or intervention is when followed with HIGH FIDELITY.

THE ITT MEASURE is the more cynical or realistic version. We never expect that programs work exactly as designed. The ITT is closer to a measure of how the laboratory equipment works once it's in the field. It is also a better estimate of how much change we can expect at the population level.

The different between the TOT and the ITT tells us how many gains can be made by improving program implementation! So both are useful and important!



## One of The Physicists Behind The Higgs Boson Has Made an Algorithm to Replace The Pill It's up to 99.5% effective at stopping pregnancy.

Those more effective methods are ones that don't require people to remember to take a pill, put on a condom, or record their temperature daily, such as intrauterine contraception or implants.

That's because human error can mess with things quite a lot. In fact, the UK's National Health Service (NHS) explained that when the app was used perfectly all the time, only five out of every 1,000 women would fall pregnant every year - a rate slightly better than the pill (99.5 percent).

But for "typical use" - where the app isn't used entirely correctly every day - it's more likely that seven out of every 100 women would experience accidental pregnancies, which is around 93 percent efficiency.

Which is treatment on the treated effect?

Which is the intention-to-treat effect?

### NOTE !!!

These two measures are **NOT** about attrition.

They are about **refusing the treatment** or failing to follow the properly-prescribed treatment regiment (or one school burning down during the study).

Non-compliance means participants refuse to receive or comply with the treatment. But they do not refuse to participate in the study – we can still measure their performance in the second time period.

Attrition means people left the study. You can accept the treatment and still attrit. Or you can refuse treatment and refuse participation. Either way, attrition means we can't measure the outcome in the post-treatment study period.

