

HANDOUT 10 – RANDOMIZED EXPERIMENTS

AGENDA

- Introduction
- Policy question: Does deworming raise students’ performance?
- Trichuris trichiura, growth and students’ performance: Evidence from Jamaica
- Threats to internal validity
- Takeaways
- In preparation for Monday’s class with Lant Pritchett
- Vocabulary

BIBLIOGRAPHY FOR TODAY’S CLASS

- Simeon et al. (1995). Treatment of Trichuris trichiura infections improves growth, spelling scores and school attendance in some children. (*)
- Miguel, E., & Kremer, M. (2004). “Worms: identifying impacts on education and health in the presence of treatment externalities.” *Econometrica*, 72, 159-217. (**)
- Angrist and Pischke (2015), 1 (**)
- Stock and Watson (2007), 8.3 (**)

INTRODUCTION

Randomly assigning subject to control and treatment groups:

- Ensures that treatment is not correlated with individual characteristics (even unobservable characteristics)
- Ensures that any difference in outcomes between the treatment and control groups are attributable to the treatment, and not to other differences between the two groups.

We have two ways of estimate effects of randomized experiments:

	<u>Difference in Means</u>	<u>Regression Analysis</u>
Estimation	$Impact = \bar{Y}_T - \bar{Y}_C$	$Y = \beta_0 + \beta_1 TREAT + u$ TREAT is a dummy variable indicating person is in treatment group. Impact: $\beta_1 = 0$
Hypothesis Testing	Conduct a t-test for the difference in means: $H_0: u_r - u_c = 0$	Conduct a t-test to determine whether β_1 is significantly different from zero: $H_0: \beta_1 = 0$
Additional Benefits of Method		Can include additional controls to improve the precision of the estimate: $Y = \beta_0 + \beta_1 TREAT + \beta_2 X_2 + \dots + \beta_k X_k + u$ where $X_2 \dots X_k$ are several characteristics at baseline

POLICY QUESTION: DOES DEWORMING RAISE STUDENTS PERFORMANCE?

Each year, approximately 1.3 billion people suffer from hookworm and roundworm infections.

There is concern that poor health due to such infections may reduce students' performance in developing countries.

Policy Question:

What are the effects of implementing a deworming program on students' performance?

TRICHURIS TRICHIURA, GROWTH AND STUDENTS' PERFORMANCE EVIDENCE FROM JAMAICA

Simeon DT, Grantham-McGregor SM, Callender JE, Wong MS (1995): "*Treatment of Trichuris trichiura infections improves growth, spelling scores and school attendance in some children,*" Journal of Nutrition, 125, 1875-83.

Study design:

- 407 children in ages 6-12 in grades 2 to 5 from 14 public schools
- Schools chosen on the advice of local health professionals: children already infected with *Trichuris trichiura* infections were recruited for study
- Individual students were then randomly assigned to:
 - Treatment: Deworming drug given
 - Control: Placebo drug given
- Length of study: 6 months
- School attendance, performance, and anthropometric data were collected before and during the intervention (only available for 264 children due to loss of class registers in some schools/grades)

What do researchers mean by "residualized gain scores for each outcome variable"

Treatment effects were determined using multiple regression analyses. First, residualized gain scores were calculated for each outcome variable. To do this, linear regression analyses were conducted with the post-test measures as the dependent variable and the pretest measures as the independent variable. The residuals from these analyses were saved for use as the dependent variables in subsequent analyses. In these multiple regression analyses, treatment group was entered as a dummy variable and other potentially confounding variables were entered as covariates. The latter included gender, age, socioeconomic status, the intensity of *Ascaris* infections (log transformed) and school attended (as dummy variables). The analyses were repeated with treatment-by-stunting and treatment-by-intensity interactions as additional independent variables.

Trichuris trichiura, growth and school performance: Evidence from Jamaica: Team Work #2

TABLE 3

Results of the multiple regression analyses showing the significant treatment-by-stunting and treatment-by-intensity interactions for Jamaican school children, after controlling for the covariates¹

	b^2	P
Spelling		
Treatment	-0.1	0.62
Intensity	-0.3	0.57
Treatment × intensity	1.6	<0.05
School attendance		
Treatment	-3.2	0.08
Stunting ³	-6.8	0.01
Treatment × stunting	9.9	0.007
Body mass index		
Treatment	0.12	0.003
Intensity ⁴	0.13	0.15
Treatment × intensity	-0.35	0.009

¹Covariates included age, gender, socioeconomic status, intensity of *Ascaris* infection and school attended.

² b is the unstandardized regression coefficient.

³Defined as height-for-age Z-score <-1.

⁴Defined as *Trichuris* infection intensities above and below 7000 eggs per gram of stool.

When researchers switch from comparing means to regression analysis (Table 3), what is the sample regression function with interactions they are running?

What additional information do the interactions provide in this specification?

THREATS TO INTENAL VALIDITY

Threat #1: Unbalanced Groups

- If the treatment and controls group are different to begin with, the estimated treatment effect is likely to be biased.
- Are the treatment and control groups balanced here (Table 1)?

TABLE 1
Baseline characteristics of the treatment and placebo groups of Jamaican school children

	Group		Significance P
	Treatment (n = 206)	Placebo (n = 201)	
Age, ¹ y	9.2 ± 1.2	9.2 ± 1.3	0.63
Gender, % boys	52	47	0.35
Residence, % Kingston	35	33	0.77
<i>Ascaris</i> infection			
% Infected	42	50	0.13
Light	11	10	
Moderate	25	29	
Heavy	7	11	
Socioeconomic index ¹	5.9 ± 2.2	6.1 ± 2.2	0.28
<i>Trichuris</i> intensity, ^{2,3} epg	2421 (1200-25,458)	2667 (1200-41,733)	0.06
Anemia, ⁴ % hemoglobin <110 g/L	10	16	0.20
Low ferritin, ⁴ % <12 µg/L	6	9	0.39

¹Means ± SD.

²epg = eggs per gram of stool.

³Values are medians, with the range in parentheses.

⁴Blood samples were obtained from only 264 children.

Threat #2: Imperfect compliance

- In practice, the researcher may not be able to enforce that the treatment protocol is followed
- There are two forms of imperfect compliance:
 - **No-shows:** Not all the treatment group members get treated (i.e. take-up rate <100%).
 - **Crossovers:** Some control group members get treated.

Suppose some children assigned to the treatment group never took the deworming drug. Will the mean difference in table 2 be larger or smaller than the true effect of taking deworming drugs?

Suppose some children assigned to the control group ended up taking deworming drugs. Will the mean difference in table 2 be larger or smaller than the true effect of taking deworming drugs?

Summary: no-shows and crossovers lead to _____ estimation of the true treatment effects.

- Jamaica study: no direct evidence of non-shows and crossovers.
- We will learn statistical solution (instrumental variables) to estimate the treatment effect even in the presence of no-shows and crossovers.

Threat #3: Attrition

- Attrition – or non-response – means that some individuals disappear from the final sample. This may lead to an attrition bias.
 - If attrition is not correlated with treatment status, this does not create bias (but smaller sample size reduces precision).
 - If attrition is correlated with treatment status (i.e. those who left treatment group differ from those who left control group), this creates a bias i.e. attrition bias.

In the Jamaica study, 10 children from the treatment group and 5 children from the control group changed schools and thus were not in the final sample. Is this a threat to the internal validity of the experiment? Under what conditions is the answer yes?

Threat #4: Spillover Effects

- If the treatment affects outcomes of control group members indirectly, there are spillover effects.
- We will _____ estimate the true treatment effect if some control group members are affected indirectly by treatment.

Could there be spillover effects in the Jamaica deworming experiment?

WORMS: IDENTIFYING IMPACTS ON EDUCATION AND HEALTH IN THE PRESENCE OF TREATMENT EXTERNALITIES

BY EDWARD MIGUEL AND MICHAEL KREMER¹

Intestinal helminths—including hookworm, roundworm, whipworm, and schistosomiasis—infect more than one-quarter of the world’s population. Studies in which medical treatment is randomized at the individual level potentially doubly underestimate the benefits of treatment, missing externality benefits to the comparison group from reduced disease transmission, and therefore also underestimating benefits for the treatment group. We evaluate a Kenyan project in which school-based mass treatment with deworming drugs was randomly phased into schools, rather than to individuals, allowing estimation of overall program effects. The program reduced school absenteeism in treatment schools by one-quarter, and was far cheaper than alternative ways of boosting school participation. Deworming substantially improved health and school participation among untreated children in both treatment schools and neighboring schools, and these externalities are large enough to justify fully subsidizing treatment. Yet we do not find evidence that deworming improved academic test scores.

KEYWORDS: Health, education, Africa, externalities, randomized evaluation, worms.

TAKEAWAYS

- To assess the validity of any study, we should focus on both **internal and external validity**.
- Randomization is a particularly useful way of improving internal validity.
- We should still consider whether any of the problems discussed today (unbalanced groups, imperfect compliance, attrition, spillovers) might create bias in our estimates.
- In particular, we should think hard about whether the experimental design used correctly mimics the public policy measures that would be enacted in reality.

IN PREPARATION FOR MONDAY’S CLASS WITH LANT PRITCHETT

- Randomized experiments were once considered the “gold standard” for estimating causal effects and all World Bank programs were subject to a “specific program evaluation” (RCT)
- RCTs absorbed much of the funds available for development research and compromised funding for other disciplines addressing more comprehensive questions (with potentially larger impacts)
- Context matters: In trying to adapt every policy/program to the specifics of each context, we give studies lots of internal validity but compromise entirely external validity

VOCABULARY

- **Randomized experiment:** an experiment in which participants are randomly assigned to a control group, which receives no treatment, or to a treatment group, which receives a treatment.
- **Selection bias:** is the bias introduced by the selection of individuals, groups or data for analysis in such a way that proper randomization is not achieved, thereby ensuring that the sample obtained is not representative of the population.
- **Internal and external validity (review):**
 - Internal validity: when inferences about causal effects in a statistical study are valid for the population being studied.
 - External validity: when inferences and conclusions from a statistical study can be generalized from the population and the setting studied to other populations and settings.
- **Threats to internal validity:**
 - **Unbalanced groups:** whether treatment and controls are indeed similar at the beginning.
 - **Imperfect compliance:**
 - **No shows:** participants who are randomized to the treatment group, but do not received the treatment or did not follow the treatment protocol.
 - **Cross-overs:** participants who are randomized to the control group, but acquire the treatment by other means.
 - **Attrition:** loss of subjects from a study after assignment to the treatment or control group.
 - Attrition bias: If the reason for attrition is related to the treatment itself, then the attrition results in a biased OLS estimator of the causal effect.
 - **Spillover effects** (in the context of experiments): Occur when treatment affects outcomes of control group members indirectly, there are spillover effects.