FLS 6441 - Methods III: Explanation and Causation

Week 3 - Field Experiments

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April 2019

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 - Design-Based Solutions to the Fundamental Problem of Causal Inference: Which treatment assignment mechanisms avoid biases and provide plausible counterfactuals

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 - Design-Based Solutions to the Fundamental Problem of Causal Inference: Which treatment assignment mechanisms avoid biases and provide plausible counterfactuals
 - How much can we learn with better research design?
 - Model-Based Solutions: Not so much.

Rest of the Course

Independence

		Independence of Treatment Assignment	Researcher Controls Treatment Assignment?
Controlled Experi- ments	Field Experiments	✓	√
	Survey and Lab Experiments	√	√
Natural Ex- periments	Randomized Natural Experiments	✓	
	Instrumental Variables	✓	
	Discontinuities	✓	
Observational Studies	Difference-in-Differences		
	Controlling for Confounding		
	Matching		
	Comparative Cases and Process Tracing		

Section 1

Independence

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- The Treatment Assignment Mechanism depends on Potential Outcomes
- ► So estimates of the ATF are biased
- ➤ The solution?
- Treatment Assignment Mechanisms that ARE independent of potential outcomes

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 - ▶ We want to estimate:

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(5)

▶ Potential outcomes in the treatment and control groups are now **unbiased** and representative of all the units

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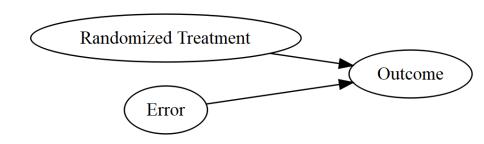
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 - It has nothing to do with potential outcomes!
 - So we get a representative sample of Y₀ and Y₁
 - Every unit has exactly the same probability of treatment
 - Potential outcomes are 'Completely Missing at Random'
 - ► No omitted variable bias is possible
 - ► No self-selection is possible
 - ► No reverse causation is possible

► This is the **entire** causal diagram:

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- Why does randomization remove selection bias?
- ► Assume: $Y_{1i} = Y_{0i} + \alpha$, where α is the real constant treatment effect

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$$A\hat{T}E = \underbrace{\alpha}_{\text{Real ATE}} + \underbrace{E(Y_0|D=1) - E(Y_0|D=0)}_{\text{Bias}}$$

► Now, use the Independence of Treatment Assignment:

$$E(Y_0|D=1) = E(Y_0|D=0)$$

$$A\hat{T}E = \underbrace{\alpha}_{\text{Real ATE}}$$

► This works for observable *and* unobservable variables

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 - We have no way of verifying if potential outcomes are biased

Balance in Repeateed Experiments

Section 2

Analysis

Analyzing Field Experiments

▶ If treatment is random we know that:

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 - NO modelling assumptions ("non-parametric")

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Analyzing Field Experiments

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$$Y_i = Y_{0i} + (Y_{1i} - Y_{0i})D_i + \epsilon_i$$

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Regression Results $(Y_i = \alpha + \beta D_i = \epsilon_i)$:

	term	estimate	std.error	statistic	p.value
1	(Intercept)	0.03459	0.07110	0.48647	0.62664
2	treatment	0.27065	0.10044	2.69472	0.00706

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- Treatment effects are still probabilistic (random variables) so we may get the wrong answer by chance
- ▶ In repeated experiments, 95% of confidence intervals will cross the true treatment effect

```
## Error in eval(predvars, data, env): object
not found
```

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Independence

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- ▶ In general, inference is more efficient with more higher-level units (more villages, less people per village)
 - But there is usually a cost trade-off

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Critiquing Field Experiments

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- ► Three reasons to include controls:
 - Small sample, but note causal inference is now model-dependent
 - Chance/residual imbalance on a specific variable which we want to adjust for
 - ▶ To improve precision, i.e. reduce the standard errors on β
 - ► The more variation in Y we can explain with covariates, the more certain we can be on the effect of D

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- ► Average treatment effects are easiest (difference-in-means equals mean-difference)
- ▶ But we can also estimate Quantile treatment effects, eg. the effect of treatment on the bottom 10% of the distribution

Section 3

Assumptions

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- 1. Compliance with Randomization procedure
- 2. Randomization produced balance on potential outcomes
- 3. SUTVA
- 4. Excludability

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- **Design:** Double-blind assignment
- ► Checks: Qualitative fieldwork

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- ► Check: Or a Kolmogorov-Smirnov Test of identical distributions

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Implementing Field Experiments

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- ▶ But spillovers are common! If you get an award, I might feel more motivated or less motivated
- Why are spillovers a problem?
 - Design: Limit risk of spillovers, eq. leave 20 miles between each unit
 - Check: Qualitative fieldwork
 - Analysis: Try to measure spillovers

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- ▶ Our treatment effect is no longer *only* the effect of our information
- Or do we want to measure these additional effects?

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- Others we don't want to capture
 - ► Eq. Measurement bias: Researchers treat treated units differently and record higher outcomes for them
 - Or Hawthorne Effects arising from being studied, not treatment (more next week)
- Design: Careful specification of treatment and control

► What if we find zero effect of government investment of \$1000 in healthcare on health outcomes, because households responded by reducing their spending by exactly \$1000? Independence

- What if we find zero effect of government investment of \$1000 in healthcare on health outcomes, because households responded by reducing their spending by exactly \$1000?
- ► Experimental treatment effects capture all net downstream effects

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 - We don't want to be guinea pigs!

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- ▶ How do we randomize?
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 - We can't directly test potential outcomes
 - 1. Qualitative research: to reconstruct the treatment process
 - Balance tests: We can directly test other variables between treatment and control
 - Randomization balances all variables, not just potential outcomes

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- ▶ What's the difference between these three options?
- ▶ What % treated? 50:50 is usually most efficient

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 - So why leave this to chance??
 - ► We can measure these variables and *enforce* balance (50%) female in both treatment and control)
 - Blocking means randomizing within fixed groups

- Randomization is *inefficient* and risky
- ▶ We know we need balance on key covariates, eg. gender, democracy
 - So why leave this to chance??
 - ► We can measure these variables and *enforce* balance (50%) female in both treatment and control)
 - Blocking means randomizing within fixed groups
 - ► Eq. We have 10 states and a sample size of 5000 so we fix 250 treated and 250 control in each state
- "Block what you can; randomize what you cannot"

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- Both work in the same way randomization avoids selection (into the data/treatment)

Section 5

Critiquing Field Experiments

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- ▶ What theory is this testing? Does it reject any theory?

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2. Generalizability of Context

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 - Selection bias has come back!

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 - General Equilibrium Effects: Average test scores went from 75% to 95%, so the exam board readjusted the test and made it harder.

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Independence

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- But sites were not representative (close to main roads and cities so they're easy to visit)
- ➤ Treatment could not be scaled (Every village cannot get visits from Columbia professors twice a year)
- And politics was ignored (No implementation unless you give them responsibility, but lose control)

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