Causality, potential outcomes and randomization

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Principles of Empirical Analysis Lecture 4

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- marketing campaing on sales
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- These are **causal** questions
 - requires evaluating counterfactual states of the world
 - "how would Y change if we changed X?"
- Compare to **descriptive** questions (lectures 1-3)
 - requires measuring the actual state of the world
 - "what is joint distribution of X and Y?"

- The next four lectures will focus on answering causal questions using research designs based on **randomization**
 - the simplest context for learning relevant statistical concepts
- The prime example is randomized controlled trials (RCT)
 - RCTs have become an important part of economits' toolkit
 - you might end up running them for living
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 - you will definitely end up interpretting results from other people's RCTs
- Even when we can't run an experiment, it is often helpful to ask: what would be the **ideal experiment** for answering this question?
 - helpful benchmark for "naturally occurring" or "quasi" experiments
 - we'll discuss an example of a "natural experiment" involving actual randomization already on Wednesday's class
 - you'll see other types of quasi-experimental approaches in lectures 8–11

- Good understanding of **why randomization eliminates selection bias** and the content and importance of the following concepts:
 - 1 causality
 - 2 counterfactual
 - 3 potential outcomes
 - 4 treatment effect
 - 5 selection bias
- Basic understanding of the ethics and limitations of randomized controlled trials (RCTs) in the context of social experiments

- Imagine that you have been asked to assist the government to evaluate the following proposal by a private investor:
 - the investor has designed a new type of integration program for newly arrived immigrants (which seems reasonably good)
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- My take: helpful to break this into two parts
 - what is the question one needs to answer?
 - how to answer it?

1 Treatment

• impact of [...]

2 Counterfactual

- impact in comparison to [...]
- **3 Outcome** and **population**
 - impact on [...]

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- What is a well-defined question for our case study?
 - my take: "what is the impact of the **new program** in comparison to **business-as-usual programs** on **participants' cumulative unemployment benefits during their first three years in Finland**?
- Next: formal definitions using the potential outcomes framework

• We focus on binary (0/1) treatments and denote treament status of individual *i* as

$$D_i = \begin{cases} 1 & \text{if she receives the treament} \\ 0 & \text{if she doesn't} \end{cases}$$

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in words: y_{1i} is the outcome of individual *i* in the state of the world where she is treated and y_{0i} is her outcome in the state of the world where she was *not* treated (note: only one state of the world occurs)

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 - $y_{1i} y_{0i}$

in words: difference in the potential outcomes with and without the treatment

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 The fundamental challenge of causal inference is that we cannot observe both y_{1i} and y_{0i} for the same individual. Instead, we observe

$$y_i = \begin{cases} y_{1i} & \text{if } D_i = 1\\ y_{0i} & \text{if } D_i = 0 \end{cases}$$

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- Why ATE and ATT?
 - treatment effect may be different for those getting the treatment than it would be for those not getting it (e.g. specific integration policy)
 - internal validity: do we learn the true effect for the treated population?
 - external validity: can we extrapolate to other populations?

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- Invalid control group leads to selection bias
 - whether the control group provides a good counterfactual or not is the key question of all design-based causal inference

• As the amount of data increases, the sample averages approach the population average (expectations)

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where the first step is from the previous slide and the second step is taken by simply adding and substracting $\mathbb{E}[y_{0i}|D=1]$

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- in words: differences in the average outcomes between treatment and control groups include the treatment effect *and* the selection bias (the difference between the two groups if neither had been treated)

- Let's return to the case of new integration program and speculate about the likely selection bias in two alternative control groups:
 - 1 all immigrants not participating in the program
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 - 1 all immigrants not participating in the program
 - 2 unemployed immigrants entering the employment services at the same time, but participate in other types of programs
- Let's assume that the new program consists of
 - 60 days intensive language training
 - followed by 6 months of guaranteed real low-skilled job
 - while the business-as-usual model includes
 - 1yr standard language training
 - "graduation" into standard unemployment

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- Thus $\mathbb{E}[y_{0i}|D=1] \mathbb{E}[y_{0i}|D=0] = 0$, i.e. no selection bias
 - in words: the control group tells us what would have happened to the treatment group in the absence of the treatment

- The key ethical concern of RCTs is the unequal treatment of the treatment and control group
 - sometimes a question of life and death (e.g. early AIDS medication)
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 - sometimes a question of life and death (e.g. early AIDS medication)
- Nevertheless, drug approval requires extensive clinical trials. Why?
 - The 1960's thalidomide tragedy led to stricker requirements that drugs have to be proved to be safe and effective before they are marketed
 - the proof comes from clinical trials (RCTs)

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 - inevitable because not all policies can be tested with RCTs
- But why not study the impact of policies suitable for experimental research designs using the most reliable methods?
 - my interpretation: policy makers often have a gut feeling that RCTs are somehow immoral (without having really thought this through)
- Aalto Economic Institute is part of this debate
 - see e.g our recent reports on social experiments and ex-post evaluations (if you are interested; i.e. this is not a requirement for this course)

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 - benefits those potentially getting the treatment later
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- Typically we do not know whether the treatment is beneficial or not
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- Features of ethically sound experiments
 - always: never cause harm knowingly, privacy protection, pre-evaluation of risks and benefits, reliable measurement, approriate test population
 - usually: informed consent (e.g. possibility to opt-out)

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 - but: many questions cannot be be answered with RCTs
 - it would be crazy to focus only on question suitable for RCTs
- RCTs are not helpful and/or possible when
 - treatment affects everyone (e.g. monetary policy)
 - the experiment would be unethical or too impractical/expensive
 - the study population differs (too much) from the relevant population
 - relevant follow-up period is impractically long
- Even when RCTs are feasible, they only guarantee internal validity

- Causality: how one thing affects another thing
 - requires comparing counterfactual states of the world to each other ("how would Y change if we changed X?")
 - at most, one of them is observed
- Control group in an experimental research design
 - the outcomes of the control group are used to infer what would have happened to the treatment group in the absence of the treatment
- Selection bias occurs when the control group is not comparable to the treatment group, i.e. $\mathbb{E}[y_{0i}|D=0] \neq \mathbb{E}[y_{0i}|D=1]$
 - = potential outcomes differ between the treatment and control groups
- Randomization eliminates selection bias
 - on expectation, the only difference between the groups is that the treatment group gets the treatment and the control group does not
 - $\rightarrow\,$ differences in average outcomes must be due to the treatment