Generalizability in Causal Inference

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Outline

- 1. What is causal inference?
- 2. Observational causal inference (internal validity)
- 3. Transportability of causal effects
- 4. Recovering from selection bias
- 5. Data fusion

What is causal inference?

Causal assumptions → Causal conclusions

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To make the leap from ______ to _____ we need **a model.** The model allows us to go from assumptions to conclusions, and the assumptions of your model must be in the same level of the leap you want to make.

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Formal language to represent the problem (nonparametrically), reduce it to an exercise of symbolic calculus, and derive complete solutions.

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Elias Bareinboim and Judea Pearl

PNAS July 5, 2016 113 (27) 7345-7352; published ahead of print July 5, 2016 https://doi.org/1

Edited by Richard M. Shiffrin, Indiana University, Bloomington, IN, and approved March 15, 20 June 29, 2015)

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Outputs: We will go over each of these for TR and SB problems. But first a quick review.

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Structural models: combine the power of potential outcomes, structural equations, and graphs.

The structural model

The **structural model is our oracle.** With a fully specified structural model we can answer **any** causal or counterfactual question.

Functional assignmentsDistribution unobserved factorsM: $Z = f_z(U_z)$ P: $P(U_z, U_x, U_y)$ $X = f_x(Z, U_x)$ $Y = f_y(X, Z, U_y)$

Causal (and counterfactual) quantities are defined in terms of our model.

M_x :	$Z = f_z(U_z)$	$E[Y_x] = E[Y do(x)] = E[f_y(x, Z, U_y)]$
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In most cases we don't have a fully specified model, but only a partial understanding of what is going on. How can we encode that knowledge?

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The question of whether our partial understanding + the data we have is sufficient for answering our query is known as the **identification problem**.

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Task: to express $P(Y|do(x)) = P(Y_x)$ in terms of P(Y,X,Z). **Symbolically,** this amounts to removing do() operators or counterfactual subscripts; **Graphically,** licensing assumptions checked via d-sep. in modified graphs.
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We also have **complete algorithms:** completeness assures us that, if we can't find a solution, it is impossible to identify the effect **without extra assumptions.** That is, no other method can do better.

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Our goal: extend our modeling tools to formally characterize when and how.

Transportability

(exp/obs) dist pop A, B, ... \rightarrow (exp/obs) dist target pop

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How can we operationalize this?

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description of which mechanisms are suspected to be different

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 $P^*(y \mid do(x)) = P(y \mid do(x), s)$
Encoding disparities: selection nodes

We will extend our causal diagram with "selection nodes" (S) which indicates *structural discrepancies* between populations.

Switching between the two populations is represented by conditioning on different values of S (or simply conditioning or not conditioning on S).

For instance, if $P(y \mid do(x))$ represents the experimental distribution of Y in the source domain Π and $P^*(y \mid do(x))$ the experimental distribution of Y in the target domain Π^* , the selection node act as a "switcher", and accounts for any discrepancy between the two populations. That is, by definition,

 $P^*(y | do(x)) = P(y | do(x), s)$

Thus, *symbolically*, our task is to *remove conditioning on S on any do() expression* (or counterfactual expression), since we do not have experimental data on the target domain.









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Thus, *graphically*, we will check for *separation of the source of discrepancy* (S) from key variables in the terms that describe out target quantity .

For clarity, selection nodes (S) are represented by square nodes (■).

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- Effect estimable directly from obs. distribution in target (vanilla identification)

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Lesson 1: differences in propensity to receive treatment do not matter for transportability of causal effects. What matters are potential effect-modifiers.

Is a randomized control trial really a gold standard?





Is a randomized control trial really a gold standard?





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Lesson 2: unless one wants to confine experimental results to the strict conditions of the studied subpopulation, even with a perfect RCT one still needs to go through a transportability exercise (ie, causal modeling).

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- **Strategy:** break relations that are not directly TR to find invariant pieces.

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A more elaborate example:

$$E^{*}[Y|do(x)] = E[Y|do(x), s]$$

= $\sum_{z} E[Y|do(x), z, s]P(z|do(x), s)$
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Now let us extend to **multiple populations**, each with **different experimental conditions:** for instance, in one domain only X was randomized while in another domain only Z was randomized... and so on.
Π^A :



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Not transportable from A.



X

 Π^B :

Not transportable from A.



Not transportable from A.

 Π^B :



Not transportable from B.





Not transportable from A.

Not transportable from B.

Not transportable at all?

 Π^B :





Not transportable from A.

Not transportable from B.

Not transportable at all?

What if we combine the experimental results of A and B?





Not transportable from A.

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 Π^B :

What if we combine the experimental results of A and B?

$$P^*(y | do(x)) = \sum_{z} P^*(y | do(x), z) P^*(z | do(x))$$

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$$= \sum_{z} P^B(y | do(z)) P^A(z | do(x))$$

RCT Z in B RCT X in A





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We have a *complete algorithms* that can decide how to combine results of several experimental and observational studies, each conducted on a different population and under a different set of conditions, so as to construct a valid estimate of the effect size for the target population.

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- It means that if the algorithm can't find a solution, then it is **impossible** to transport the causal effect of interest **without strengthening assumptions.**

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FUSION DEMO 1

Selection Bias

Selected Subpopulation
→ General Population

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- Here: **nonparametric, qualitative description** of the determinants of inclusion of units in the study sample.












Encoding the selection mechanism

Again we extend our causal diagram with "selection nodes" (S) which now indicate *selection to the study sample* (S = 1), or not (S = 0). Our target of inference is a quantity on the population as a whole, not conditioning on S.



Symbolically, our task is to express the query in terms of the available data, that is, the **distribution under selection bias** P(V | S = 1) - or more concisely $P(V | S) - \text{ and the$ **census data**we have available (if any).

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Graphically, we will check for separation of the selection mechanism S from key variables of interest that compose our query.

Very simple necessary and sufficient condition for conditional distributions.

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The conditional distribution P(y | x) is recoverable (without external data) if and only if: $(Y \perp S | X)$



 $P(y \mid x)$ recoverable

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Note this is different from recovering the causal effect P(y | do(x)).

For instance, in the third model, P(y|x) is not recoverable, while P(y|do(x)) is, as we show next.



Do we need external data?



E[Y|do(x)] = E[Y|do(x), s]= $\sum_{z} E[Y|do(x), z, s]P(z|do(x), s)$ = $\sum_{z} E[Y|x, z, s]P(z|s)$



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Don't need external data

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$$External data on Z$$

$$= \sum_{z} E[Y | do(x), z, s]P(z)$$

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FUSION DEMO 2

Data Fusion $(d_1, d_2, d_3, d_4) \rightarrow (d'_1, d'_2, d'_3, d'_4)$

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		Dataset 1	Dataset 2	Dataset 3
d_1	Population	Los Angeles	New York	Texas
d ₂	Obs. / Exp.	Experimental	Observational	Experimental
	Treat.Assign.	Randomized Z ₁	-	Randomized Z ₂
d ₃	Sampling	Selection on Age	Selection on SES	-
d4	Measured	X ₁ , Z ₁ , W, M, Y ₁	X ₁ , X ₂ , Z ₁ , N, Y ₂	X ₂ , Z ₁ , W, L, M, Y ₁

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-Observational Causal Inference: $(d_1, see(x), d_3, d_4) \rightarrow (d_1, do(x), d_3, d_4)$

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d ₃	Sampling	Selection on Age	Selection on SES	-
d4	Measured	X ₁ , Z ₁ , W, M, Y ₁	X ₁ , X ₂ , Z ₁ , N, Y ₂	X ₂ , Z ₁ , W, L, M, Y ₁

-Observational Causal Inference: $(d_1, see(x), d_3, d_4) \rightarrow (d_1, do(x), d_3, d_4)$ - Sampling Selection Bias: $(d_1, d_2, select(age), d_4) \rightarrow (d_1, d_2, \{\}, d_4)$

We can describe each data collection as the tuple:

 $(d_1, d_2, d_3, d_4) = (population, obs/exp., sampling selection, observed data)$

		Dataset 1	Dataset 2	Dataset 3
d ₁	Population	Los Angeles	New York	Texas
d ₂	Obs. / Exp.	Experimental	Observational	Experimental
	Treat.Assign.	Randomized Z ₁	-	Randomized Z ₂
d ₃	Sampling	Selection on Age	Selection on SES	_
d4	Measured	X ₁ , Z ₁ , W, M, Y ₁	X ₁ , X ₂ , Z ₁ , N, Y ₂	X ₂ , Z ₁ , W, L, M, Y ₁

-Observational Causal Inference: $(d_1, see(x), d_3, d_4) \rightarrow (d_1, do(x), d_3, d_4)$

- Sampling Selection Bias:
- Transportability:

$$(d_1, d_2, select(age), d_4) \rightarrow (d_1, d_2, \{\}, d_4)$$

 $(LA, d_2, d_3, d_4) \rightarrow (NY, d_2, d_3, d_4)$

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 -Observation
 $do(x), d_3, d_4$)

 - Sampling Se In general: $(d_1, d_2, d_3, d_4) \rightarrow (d_1', d_2', d_3', d_4')$ $(d_1, d_2, \{\}, d_4)$

 - Transportab
 $, d_3, d_4$)

Conclusions

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Software under development: Causal Fusion.

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Thank you!

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