Adjustment Sets and Approaches - and limitations / critiques

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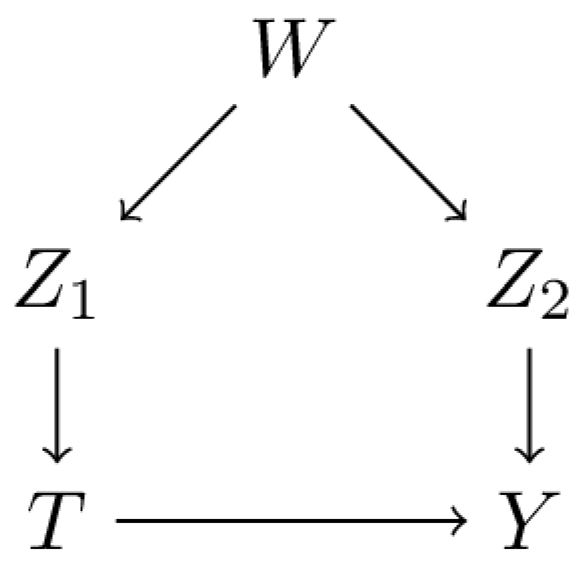
Adjustment sets and approaches

How to find adjustment sets?

- adjustment sets:
  - the back-door criterion states that any set Z that blocks all backdoor paths from X to Y is a sufficient adjustment set for causal effect estimation of $P(Y|\text{do}(X))$ using the backdoor formula.
  - how do we find these sufficient sets?
  - what if there are multiple?
- adjustment: how to do this?
  - stratification
  - what is regression adjustment?
  - T-learner vs S-learner

Valid adjustment sets

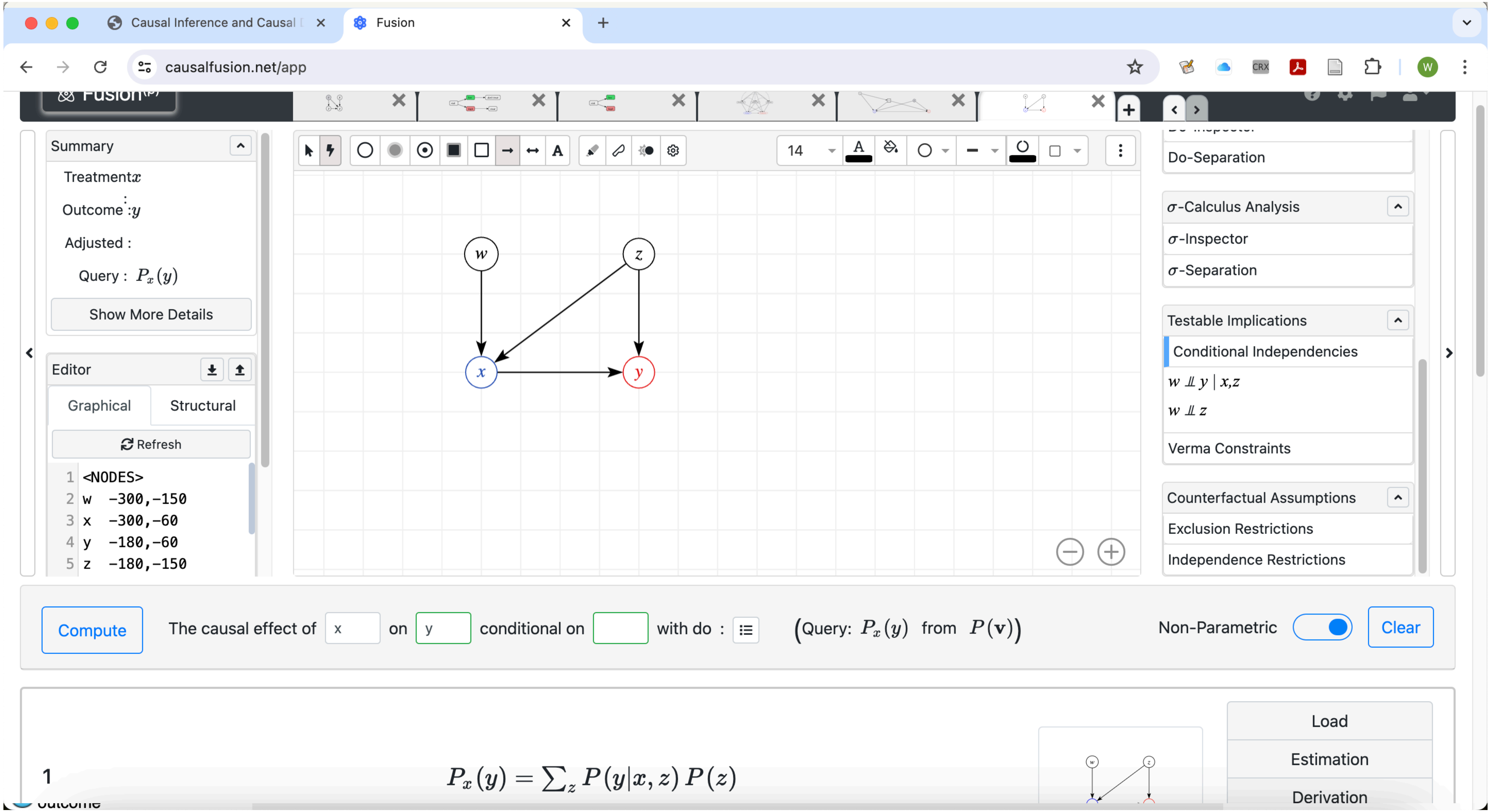


dag

- in general:
  - PA_T (the direct parents of treatment T: Z_1) are a valid adjustment set
  - PA_Y (the direct parents of outcome Y: Z_2) are a valid adjustment set
- in this case:
  - W is also a valid adjustment set

Valid adjustment sets: picking one

- websites like dagitty.net and causalfusion.net provide user-friendly interfaces for creating and exporting DAGs, in addition:
  - valid adjustment sets (if they exist)
  - testable conditional independencies



causalfusion.net

dagitty.net

How to do adjustment

What not to do

1. do univariable pre-screening against outcome (and / or treatment)

- this should maybe never be done
- especially not in the context of causal inference

Adjustment formula

$$P(y|\text{do}(x)) = \sum_z P(y|x,z)P(z)$$

- entails summing over all possible values of Z
- say Z is 5 categorical variables with each 3 categories, this means $4^5 = 1024$ estimates of:
  - $P(y|x,z)$ for each value of x
- what if Z is continuous?
- in practice, researchers rely on smoothness assumptions (e.g. regression) to estimate $P(Y|x,z)$ with a parametric model
- this assumption *can* be based on substantive causal knowledge, but often seems inspired rather pragmatism or necessity
- misspecification of this estimator leads to biased results (even if you know all the confounders)

Target queries

- up to now we've worked exclusively with $P(y|\mathrm{do}(t))$: the probability of observing outcome y when setting treatment T to t
- this is not typically what is of most interest, say there are two treatment options $T \in \{0,1\}$ (control and 'treatment')

1. *average treatment effect*

$$\mathrm{ATE} = E[y|\mathrm{do}(t=1)] - E[y|\mathrm{do}(t=0)]$$

2. *conditional average treatment effect*

$$\mathrm{CATE} = E[y|\mathrm{do}(t=1), w] - E[y|\mathrm{do}(t=0), w]$$

3. *prediction-under-intervention* $P(y|\mathrm{do}(t), w)$ (more on this on day 4)

- these can be computed from $P(y|\mathrm{do}(t), w)$

The simplest case: linear regression

- assume the following structural causal model (z is confounder, u is exogenous noise):

$$f_y(t, z, u) = \beta_t t + \beta_z z + \beta_u u$$

- then:

$$\text{ATE} = E[Y|\text{do}(t = 1)] - E[Y|\text{do}(t = 0)]$$

- i.e. the ATE collapses to the the regression parameter β_t in a linear regression model of y on t, z

General estimators for the ATE and the CATE (meta-learners)

- denote $\tau(w) = E[y|\text{do}(t=1), w] - E[y|\text{do}(t=0), w]$
  - (assuming W is a sufficient set)
- T-learner: model $T=0$ and $T=1$ separately (e.g. regression separetely for treated and untreated):

$$\mu_0(w) = E[Y|\text{do}(T=0), W=w]$$
$$\mu_1(w) = E[Y|\text{do}(T=1), W=w]$$
$$\tau(w) = \mu_1(w) - \mu_0(w)$$

- S-learner: use T as just another feature

$$\mu(t, w) = E[Y|T=t, W=w]$$
$$\tau(w) = \mu(1, w) - \mu(0, w)$$

- (many other variants combinations: this is a whole literature)

Intuitive way-pointers:

- where does the complexity come from?

a. variance in outcome under control: $E[y|\text{do}(T=0), w]$

b. variance CATE: $\tau(w)$ (in statistics: *interaction* between treatment and covariate)

Where does the variance come from?

$$\begin{tikzcd} & X \arrow[d] \\ T \arrow[r] & Y \end{tikzcd}$$

DAG

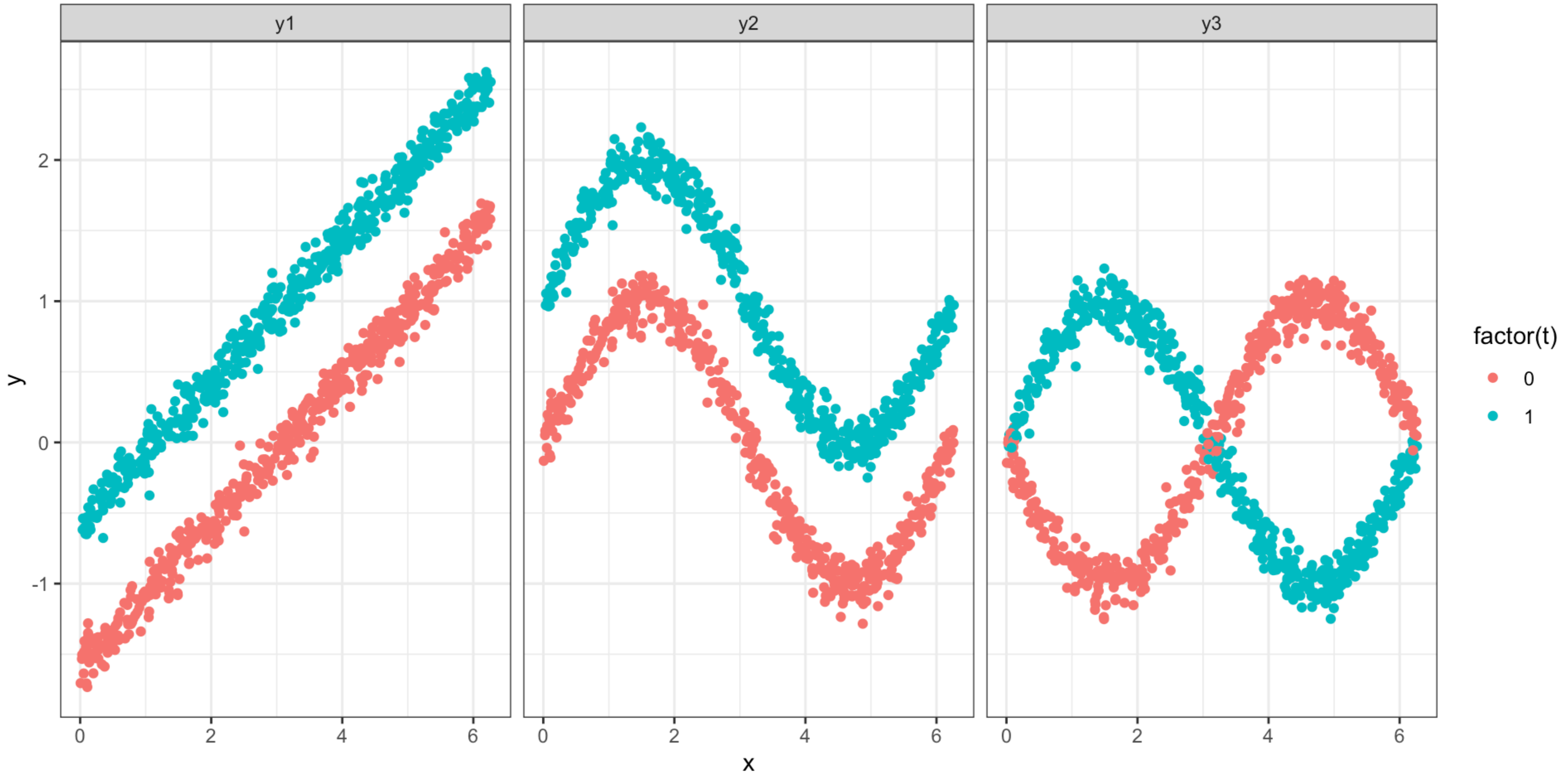


Figure 1: Three datasets with the same DAG

1. $Y = T + 0.5(X - \pi) + \epsilon$ (linear)
2. $Y = T + \sin(X) + \epsilon$ (non-linear additive)
3. $Y = T * \sin(X) - (1 - T)\sin(x) + \epsilon$ (non-linear + interaction)

Limitations of DAGs and SCMs

Making DAGs

- how do you get a DAG? up to now we assumed we had one
- based on prior evidence, expert knowledge
- “no causes in, no causes out”

A003024: The death of DAGs?

The number of possible DAGs grows super-exponentially in the number of nodes

n_nodes	n_dags	time at 1 sec / DAG
1	1	
2	3	
3	25	
4	543	
5	29281	> an hour
6	3781503	> a day
7	1138779265	> a year
8	783702329343	
9	1213442454842881	> human species
10	4175098976430598143	> age of universe

Do we need to consider all DAGs?

- a single sufficient set suffices
- adjusting for all direct causes of the treatment or all direct causes of the outcome are always sufficent sets
- can we judge these without specifying all covariate-covariate relationships?
- potential approach:
  - put all potential confounders in a cluster (e.g Anand et al. 2023)
  - ignore covariate-covariate relationships in that cluster
  - what happens when (partial) missing data?

SCM vs potential outcomes

- definition of causal effect
  - PO: averages of individual potential outcomes
  - SCM: submodel or mutilated DAG
- both require positivity
- d-separation implies conditional independence (exchangeability)

References

Anand, Tara V., Adele H. Ribeiro, Jin Tian, and Elias Bareinboim. 2023. "Causal Effect Identification in Cluster DAGs." *Proceedings of the AAAI Conference on Artificial Intelligence* 37 (10): 12172–79. https://doi.org/10.1609/aaai.v37i10.26435.