Counterfactuals and directed acyclic graphs in epidemiology

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Erasmus MC, Rotterdam

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Causal inference

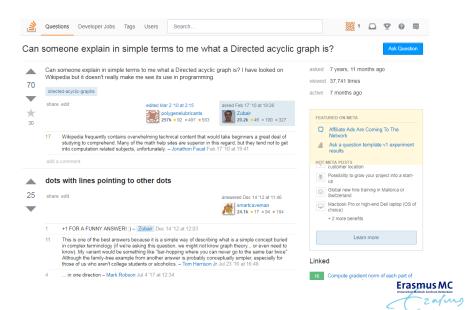
- ▶ The goal of causal inference: to know how outcome Y would change if we changed exposure A from a = 0 to a = 1
- ▶ Y^a: the value outcome Y would take if we set A to a
- So $Y^{a=1}$ is the value Y would take if we set a=1 (N.B.: this is not Y|A=1)
- ▶ Ideally, we would like to observe $Y^{a=1}$ and $Y^{a=0}$ for everyone
- ▶ But we observe at most half of the vector Y^a because we can only observe each person as $Y^{a=1}$ or $Y^{a=0}$
- ▶ We write the causal effect we're interested in as $E[Y^{a=1} Y^{a=0}]$ or $Pr[Y^{a=1} = 1] Pr[Y^{a=0} = 1]$ or $\frac{Pr[Y^{a=1} = 1]}{Pr[Y^{a=0} = 1]}$



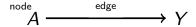
Counterfactuals

- Counterfactual notation has been used to achieve insight into estimation of longitudinal causal effects, instrumental variables and a number of other topics
- Some of the insight provided by counterfactuals can also be depicted in graphical form



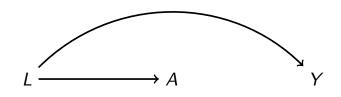


- ▶ DAGs are diagrams with simple rules that allow us to clearly think about our causal questions to determine what sources of bias we might have and how we can avoid them
- ▶ DAGs are composed of two elements: nodes and edges
- Edges are directed (i.e. one causes the other)
- ▶ DAGs must be acyclic, no feedback loops are allowed
- ▶ The edges are not deterministic



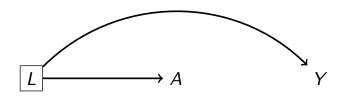


 $ightharpoonup Y \not\perp\!\!\!\perp A$ and $Y^a \perp\!\!\!\perp A$



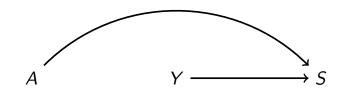


 $ightharpoonup Y \perp \!\!\!\perp A|L$ and $Y^a \perp \!\!\!\perp A|L$



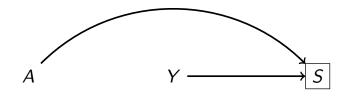


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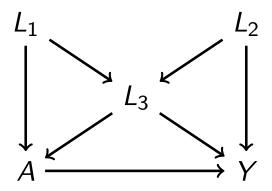


 $ightharpoonup Y \not\!\perp\!\!\!\perp A|S$ and $Y^a \!\!\perp\!\!\!\perp A|S$



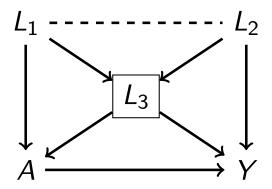


- With a DAG you can identify whether it's possible to achieve conditional exchangeability
- ▶ Here adjusting for L_1 , L_3 or L_2 , L_3 or L_1 , L_2 , L_3 are all sufficient adjustment sets





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Epidemiologists rejoice! "'the task of selecting an appropriate set of covariates to control for confounding has been reduced to a simple "roadblocks" puzzle manageable by a simple algorithm" arxiv.org/abs/1801.04016 and

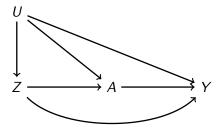


Comments on: The tale wagged by the DAG | International ...
I am grateful to the editors for the opportunity to comment on Nancy Krieger and George Davey Smith's article, 'The tale wagged by the DAG', which appeared in t

This is true! But the DAG has to be correctly specified which is the hard (impossible) part.

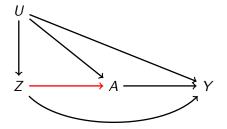
Erasmus MG

► This DAG makes the least amount of assumptions about the observed data. Every node is connected to every other



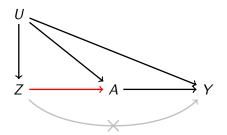


- ► The relevance assumption is represented by the red arrow, showing that there is a relationship between Z and A
- ► Z ⊥ A



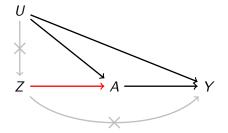


- ► The exclusion restriction (ER) assumption assumes that the direct edge between Z and A is not present
- $\blacktriangleright \mathsf{E}[Y^{z,a}] = \mathsf{E}[Y^{z',a}]$



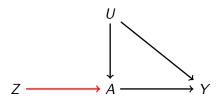


- ▶ The exchangeability assumption assumes that Z is not confounded with Y
- $ightharpoonup Y^{a,z} \perp \!\!\! \perp Z$





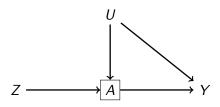
- ▶ With our three assumptions, we end up with our classic IV DAG which implies that $Z \perp \!\!\! \perp Y^a$
- ▶ Therefore if $Z \not\perp\!\!\!\perp Y$ then it must be that $E[Y^a Y^{a'}] \neq 0$
- ► IV estimation requires monontonicity and homogeneity cannot be drawn on a DAG because they are parametric concepts





Conditioning on exposure in IV analyses

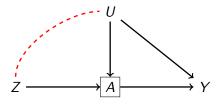
- Sometimes people like to stratify or restrict on exposure
- ▶ But if we do, now $Z \not\perp\!\!\!\perp U$ and therefore $Z \not\perp\!\!\!\!\perp Y^{z,a}$, AKA, the exchangeability assumption is violated
- Swanson et al, Am J Epi, 2015





Conditioning on exposure in IV analyses

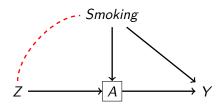
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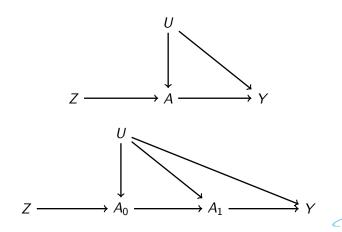
An example where a simple DAG could have helped

- "Strong evidence of collider bias was observed for smoking behaviour" (Cho et al, Sci Rep, 2015)
- "To minimize the effect of risk factors susceptible to collider bias, associations between the SNP and cardiovascular outcomes were then assessed with adjustments for smoking"
- By saying smoking is what drives collider bias, they are implying smoking is the only confounder alcohol and cardiovascular outcomes. Better off simply adjusting then?

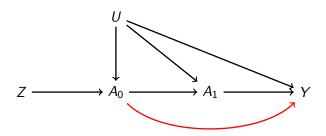




- ► We can depict a time-varying exposure by splitting the *A* node into separate nodes for each time point
- ▶ We are still assuming the IVs assumptions but splitting *A* into two nodes adds assumptions to the DAG

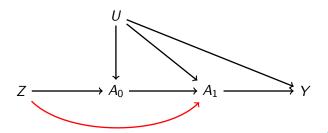


- \blacktriangleright We have assumed that A_0 does not have a direct effect on Y
- If A₀ does affect Y then we can no longer estimate the effect of A₁ ony Y
- ► This is because, although the ER holds for A as a whole, it does not hold for A₁
- ▶ So if we measure A_0 we're alright?



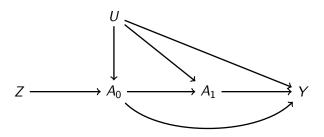


- We have also assumed that Z does not have a direct effect on A₁
- ▶ If Z does affect A_1 then we can no longer estimate the effect of A_0 ony Y
- ▶ This is because, although the ER still holds for A as a whole, it does not hold for A_0
- All this to illustrate that, in contexts such as Mendelian randomization, important thought must be given to how exposures change longitudinally



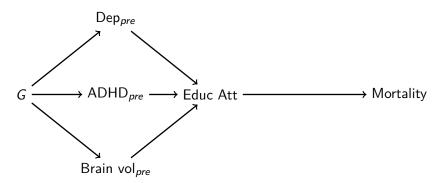


- What if we think about joint effects?
- ► First, though "lifetime effects" is often used in Mendelian randomization, it has no precise defintion
- We propose $E[Y_k^{\bar{A}+\bar{1}}-Y_k^{\bar{A}}]$
- Only identifiable with IV if the effect of the Z on A does not change with time (N.B.: A can change as long as the effect of G doesn't)
- This also has implications for which null hypotheses are testable (Swanson et al, Eur J Epi, 2018)



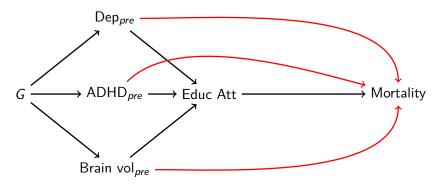


▶ DAG for a MR study of the effect of educational attainment on mortality



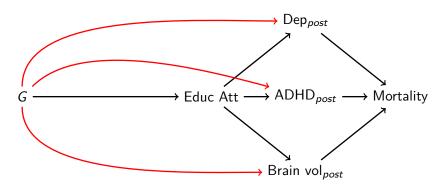


▶ But what if the intermediates between *G* and education attainment have a direct effect on mortality?



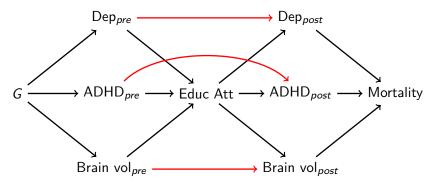


 Or what if G has a direct affect on the intermediates after educational attainment





▶ We have also assumed that Z does not have a direct effect on A₁





Thank you

- We are Erasmus MC Causal Inference Group (we don't have a name yet)
 - ► Sonja Swanson (assistant professor): Instrumental variable analysis, target trial emulation, causal inference
 - Jeremy Labrecque (postdoc): Instrumental variable analysis, target trial emulation, causal inference
 - Elizabeth Diemer (PhD student): Methodological issues in perinatal MR
 - Paloma Rojas Saunero (PhD student): Target trial emulation using data from the Rotterdam Study
- Email us at: s.swanson@erasmusmc.nl or j.labrecque@erasmusmc.nl

