

Potential Outcomes: From Matching to Propensity Scores

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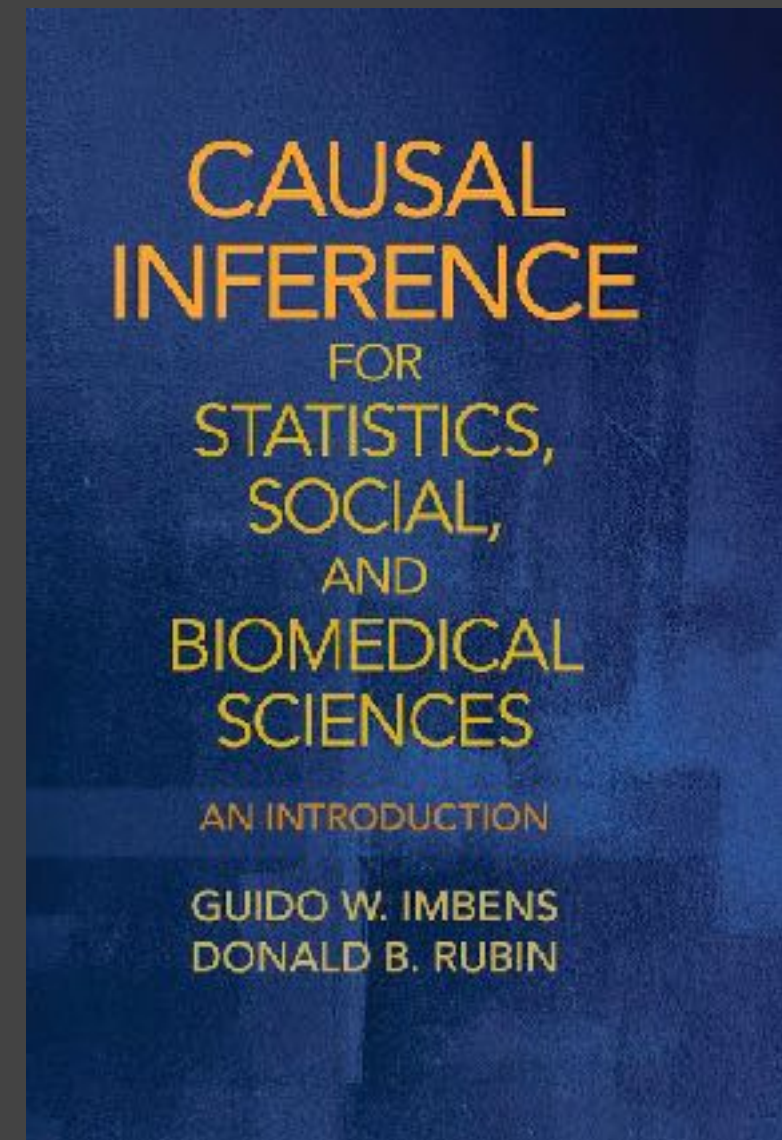
Intended Learning Outcomes

- Potential Outcomes framework
- The fundamental problem of CI
- Dealing with the CIA and Exchangeability
 - Matching
 - Propensity scores



Potential Outcomes framework a.k.a. (Neyman-)Rubin causal model

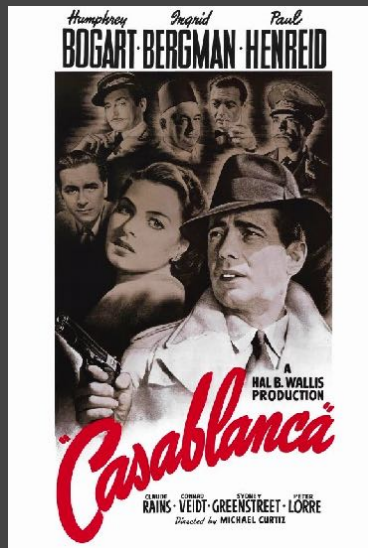
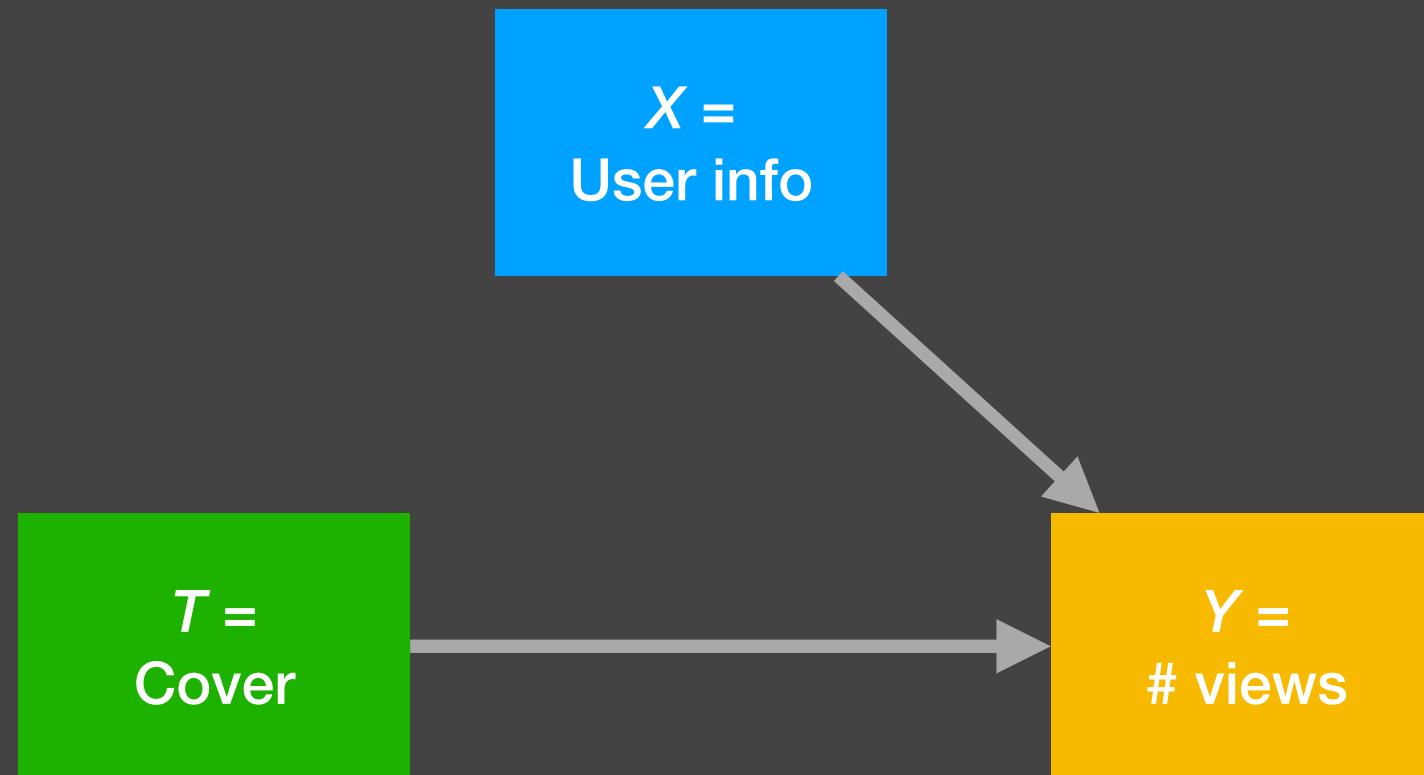
- 1923 - Jerzy **Neyman** - first idea, limited to RCTs
- 1974 - Donald **Rubin** - extension to observational studies
- 1994 - **Imbens & Angrist** - application to economics (instrumental variables)
- Today - Applied throughout medicine, economics, social sciences






Which cover would *make* you watch this movie?



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Which cover would *make you* watch this movie?

User features X				Treatment	Outcome
Age	Gender	Movies viewed	...	Cover	Watched?
55	F		Y
27	M		N
33		...

Looks like a prediction or
classification problem: $Y = f(X, T)$

The fundamental problem

User features X				Treatment	Outcome	Potential Outcomes		Causal effect
Age	Gender	Movies viewed	...	Cover	Watched?	$Y(T=0)$	$Y(T=1)$	$Y(1)-Y(0)$
55	F		Y	Y	?	?
27	M		N	?	N	?
33		...	?	Y	?
					$E[Y] =$	4.1%	5.9%	(+1.8%) «observational»



Looks like a prediction or classification problem: $Y = f(X, T)$

But actually we have two populations — are they «equivalent» («exchangeable»)?

Are the underlying populations similar across X ? Are there confounding features?

When can we mix data?

- **Intuition:** check if the two populations (*control* and *treatment*) are **exchangeable** (\sim i.i.d.)
- **Formally:**
Conditional Independence Assumption (CIA):
«Assignment to *Treatment* or *Control* group has been at random [w.r.t. observed features]»

User features X				Treatment
Age	Gender	Movies viewed	...	Cover
55	F	
...	
...	
27	M	
33	

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Conditional Independence Assumption (CIA):
«Assignment to *Treatment* or *Control* group has been at random [w.r.t. observed features]»
- In Randomized Controlled Trials (RCTs), validity of the CIA is assessed by checking e.g. averages of relevant features (age, sex...) → «Table 1»
- Unlikely to be satisfied in observational data.




Table 1. Baseline Characteristics of the 500 Patients.^A

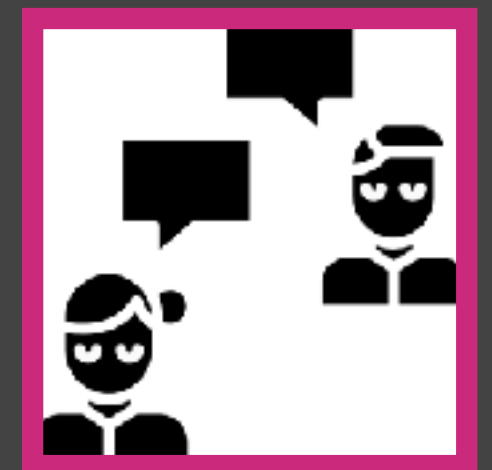
Characteristic	Intervention (N = 233)	Control (N = 267)
Age — yr		
Median	65.8	65.7
Interquartile range	54.5–76.0	55.5–76.4
Male sex — no. (%)	135 (57.9)	157 (58.8)
NIHSS score [†]		
Median (interquartile range)	17 (14–21)	18 (14–22)
Range	3–30	4–38
Location of stroke in left hemisphere — no. (%)	116 (49.8)	153 (57.3)
History of ischemic stroke — no. (%)	29 (12.4)	25 (9.4)
Atrial fibrillation — no. (%)	66 (28.3)	60 (22.8)
Diabetes mellitus — no. (%)	34 (14.6)	34 (12.7)
Prestroke modified Rankin scale score — no. (%) [‡]		
0	190 (81.5)	214 (80.1)
1	21 (9.0)	29 (10.9)
2	12 (5.2)	13 (4.9)
≥2	10 (4.3)	11 (4.1)
Systolic blood pressure — mm Hg [§]	146±25.0	145±24.4
Treatment with IV alteplase — no. (%)	203 (87.1)	242 (90.6)
Time from stroke onset to start of IV alteplase — min		
Median	85	87
Interquartile range	67–110	65–116
ASPECTS — median (interquartile range) [¶]	9 (7–10)	9 (8–10)
Intracranial arterial occlusion — no./total no. (%)		
Intracranial ICA	1/233 (0.4)	3/266 (1.1)
ICA with involvement of the M1 middle cerebral artery segment	59/233 (25.3)	75/266 (28.2)
M1 middle cerebral artery segment	154/233 (66.1)	165/266 (62.0)
M2 middle cerebral artery segment	18/233 (7.7)	21/266 (7.9)
A1 or A2 anterior cerebral artery segment	1/233 (0.4)	2/266 (0.8)
Extracranial ICA occlusion — no./total no. (%)	75/233 (32.2)	70/266 (26.3)
Time from stroke onset to randomization — min ^{‡‡}		
Median	204	196
Interquartile range	152–251	149–266
Time from stroke onset to groin puncture — min		
Median	260	NA
Interquartile range	210–313	

The recommender's view





A new user logs in...

What cover do we show them?

User features X				Treatment	Outcome
Age	Gender	Movies viewed	...	Cover	Watched?
55	F		Y
27	M		N
33		...
44	M	?	...







Matching

User features X				Treatment	Outcome
Age	Gender	Movies viewed	...	Cover	Watched?
55	F		Y
27	M		N
33		...
44	M		N
44	M	?	?

Intuition:

- See if you already met a similar case and apply what you learned

Matching

User features X				Treatment	Outcome
Age	Gender	Movies viewed	...	Cover	Watched?
55	F		Y
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33		...
44	M		N
44	M	?	?





Intuition:

- See if you already met a similar case and apply what you learned

Drawbacks:

- Need to search whole dataset — potentially slow
- No guarantee to find a match!
- Curse of dimensionality — things get worse the more you know about your users [larger $\dim(X)$]

Propensity Scores

User features X				Treatment	Outcome
Age	Gender	Movies viewed	...	Cover	Watched?
55	F		Y
27	M		N
33		...
44	M		N
44	M	?	?

Goals:

- Improve robustness by relying on more than $N=1$ observations.
- Exploit what we know about outcomes in C and T groups.
- Avoid curse of dimensionality
- Find objective way to define «distance» between units.





Idea:

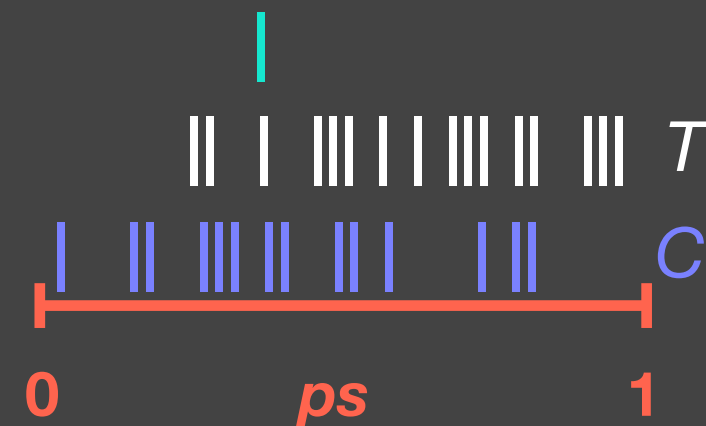
- Reduce information in X to a single number:
What is the probability that a user with features $X=x$ was in the treatment group?

$$ps = P(T|X)$$

- $P(T|X)$ model tries to capture how biases cropped up in the assignment to Treatment group in the real world.





Propensity Score Stratification

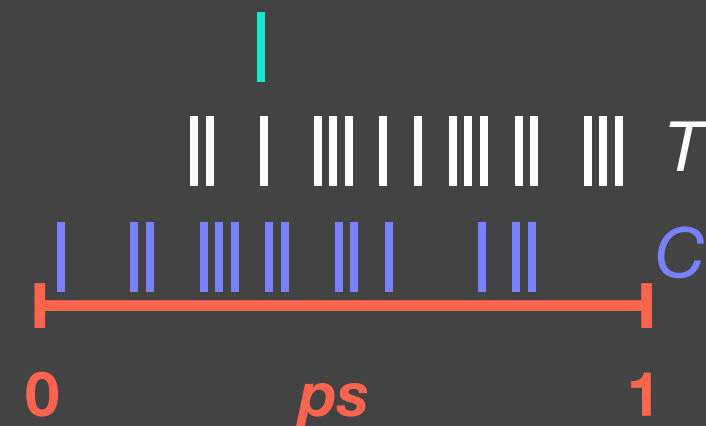
User features X				Treatment	PS	Outcome
Age	Gender	Movies viewed	...	Cover	$P(T X)$	Watched?
55	F		0.8	Y
27	M		0.95	N
33		0.65	...
44	M		0.45	N
44	M	?	0.45	?



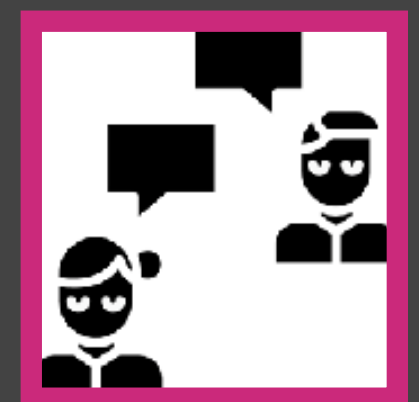
Recall: $P(T|X)$ model tries to capture how biases cropped up in the assignment to Treatment group in the real world. Contains no info on outcomes.

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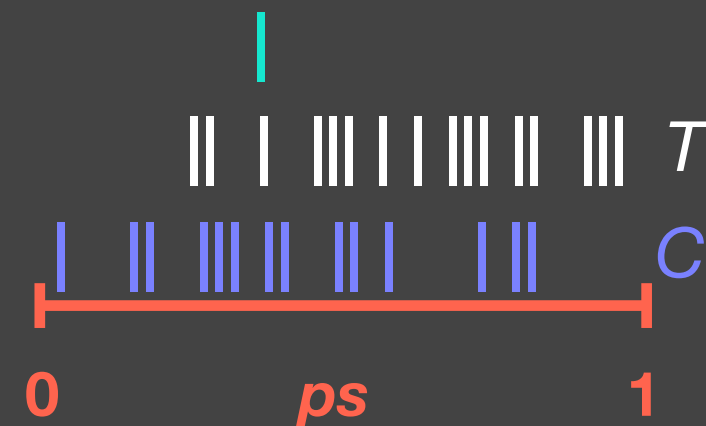
How would this look if assignment had been purely at random (CIA)?



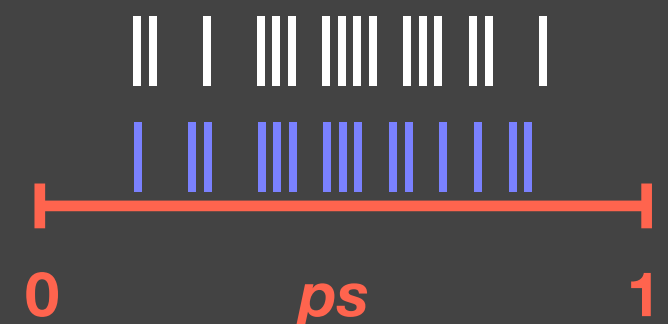
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Propensity Score Stratification




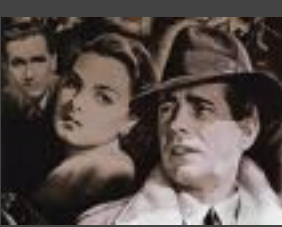
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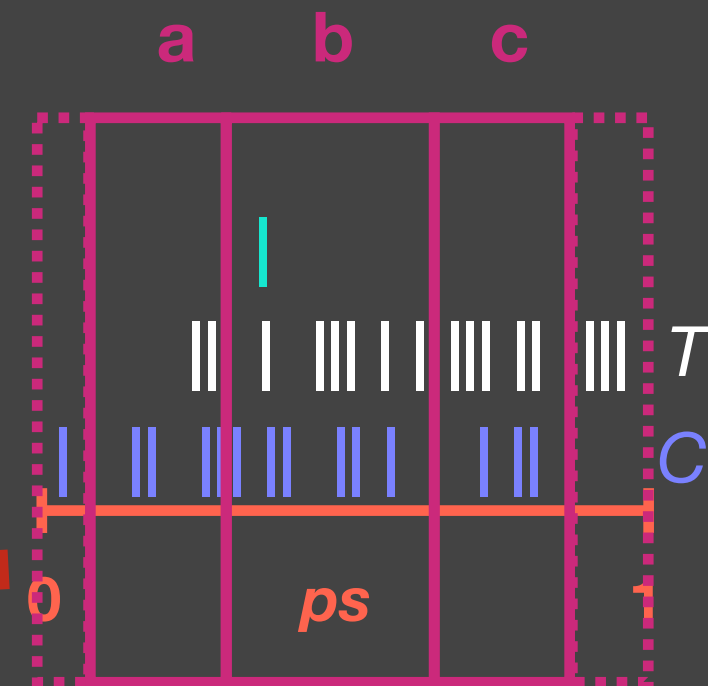


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Propensity Score Stratification

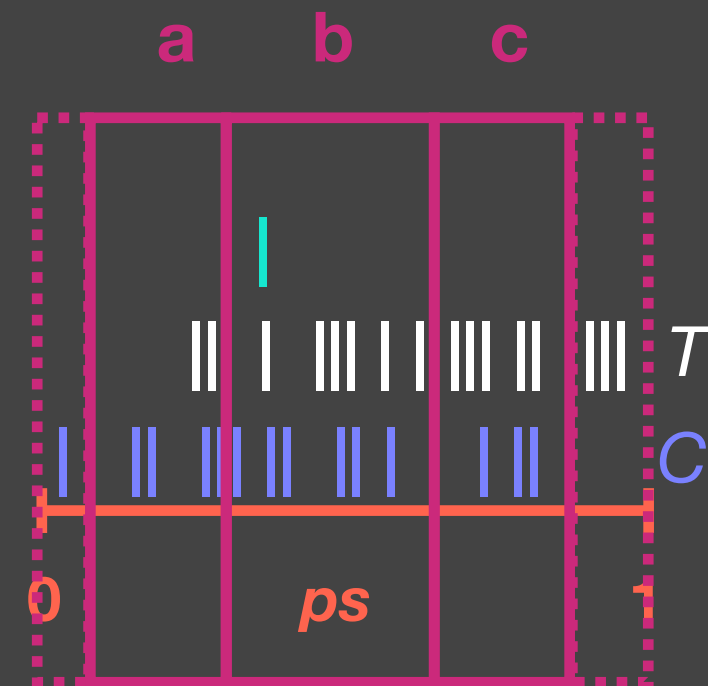
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Age	Gender	Movies viewed	...	Cover	$P(T X)$	Watched?
55	F		0.8	Y
27	M		0.95	N
33		0.65	...
44	M		0.45	N
44	M	?	0.45	?



- Drop outliers in ps-space, not in X-space.
- Adjust bin width to satisfy «exchangeability» within the bin.

Propensity Score Stratification

	a	b	c
ps	0.1-0.3	0.3-0.6	0.6-0.85
# obs.	100	200	150
Age	36(4)	48(3)	55(5)
Gender	M(52%)	F(50.1%)	F(55%)
ATE	-2%	-0.5%	5.4%



- Adjust bin width to satisfy «exchangeability» within the bin.
- Then extract causal effect by bin.
- Allows us to design group-targeted actions → customer segmentation.

Propensity Score Stratification

	a	b	c
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Age	36(4)	48(3)	55(5)
Gender	M(52%)	F(50.1%)	F(55%)
ATE	-2%	-0.5%	5.4%



Overall causal ATE:



$$\frac{100}{450}(-0.02) + \frac{200}{450}(-0.005) + \frac{150}{450}(0.054) = +1.1\%$$



Recall «Observational»
ATE: +1.8%

- Adjust bin width to satisfy «exchangeability» within the bin.
- Then extract causal effect by bin.
- Allows us to design group-targeted actions → customer segmentation.
- Causal estimate of population-wide ATE.

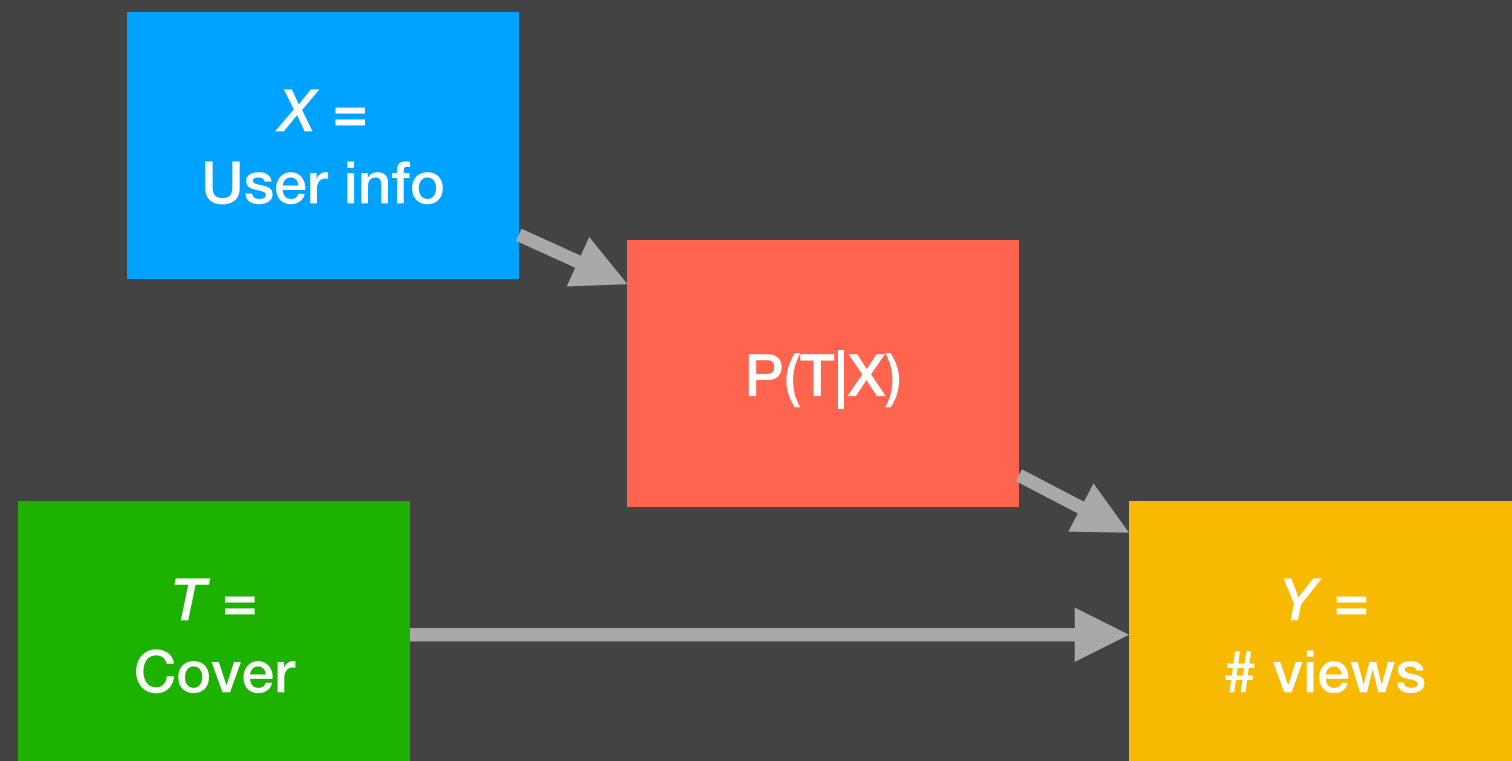
Connection with DAGs?

User features X				Treatment	PS	Outcome
Age	Gender	Movies viewed	...	Cover	$P(T X)$	Watched?
55	F		0.8	Y
27	M		0.2	N

a b c

A good DAG underlying your model to estimate $P(T|X)$ will allow you to:

- Not use irrelevant features
- Avoid biases
- Reduce uncertainty



Causal inference: Best practices

- **Always follow the four steps: Model, Identify, Estimate, Refute.**

Refute is the most important step.

«Try to prove yourself wrong»
—W.D. Phillips, Nobel prize in Physics 1997

- **Aim for simplicity.**

If your analysis is too complicated, it is most likely wrong.

- **Try at least two methods with different assumptions.**

Higher confidence in estimate if both methods agree.

- **Remember the order for validity of estimates obtained:
Randomization, Natural experiments, Conditioning.**

Consider observational methods as strong hints (but they can be misleading)

Adapted from Amit Sharma
(Microsoft Research, *DoWhy*'s lead developer)

Now turn to the Notebook pscore_oil_wells_analyse.ipynb



Charles Addams, «Skier», The New Yorker, 13 Jan 1940