

# Beyond Estimation: Next Steps for Precision Medicine

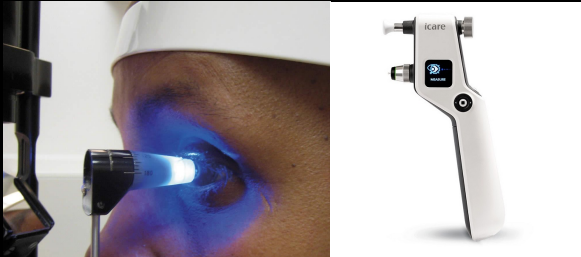
Michael Wallace, University of Waterloo

Slides available at: `mpwallace.github.io`

# Glaucoma: One Disease, Many Treatments

Glaucoma: group of eye diseases associated with elevated intraocular pressure (IOP).

IOP can be measured in various ways.



# Glaucoma: One Disease, Many Treatments

Treatment options attempt to lower IOP, they include:

- Lifestyle changes.
- Eye drops (numerous options).
- Surgery.

# Glaucoma: One Disease, Many Treatments

Treatment decisions are made based on various factors:

- Current and past IOP.
- Current and past treatments.
- Concerns over side effects.
- Broader risk factors.
- Other characteristics (such as age).

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Precision Medicine: tailoring treatment decisions to patient-level characteristics.

- Dynamic treatment regimes (DTRs) 'formalize' personalized treatment:

*"Patient presents with historic IOP of 13 and is taking Azarga. If current IOP is 15 or higher, add Alphagan, otherwise continue with only Azarga."*



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- How do we choose the best DTR?  
Should our IOP cut-off be 13, 15, 20?
- What makes this difficult?

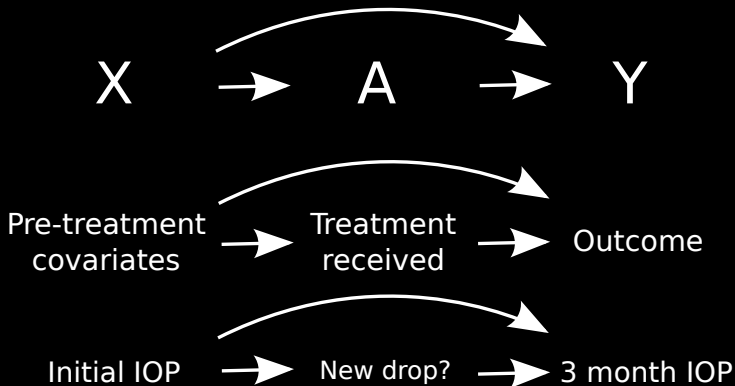
We typically work with data from observational studies.

Patient	Observed IOP	Drop added?	IOP at 3 months
1	16	No	15
2	20	Yes	16
3	21	Yes	17
4	16	Yes	16
5	15	No	18
...	...	...	...



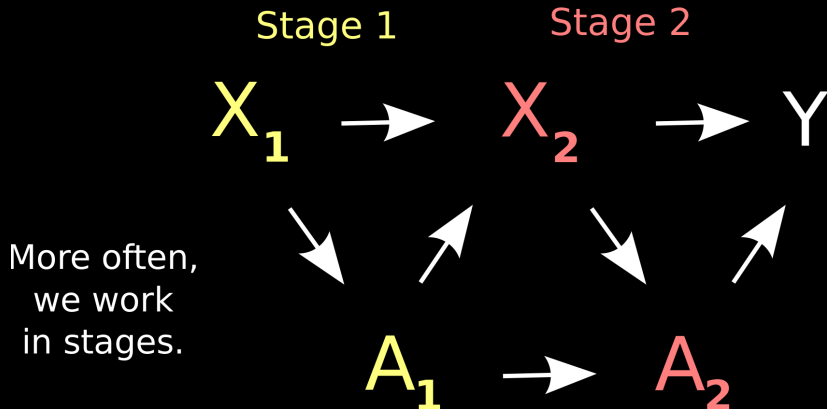
We typically work with data from observational studies.

Patient	Observed IOP $X$	Drop added? $A$	IOP at 3 months $Y$
1	16	No	15
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DTR: treatment  $A^{opt}$  that optimizes  $E[Y|X, A^{opt}]$

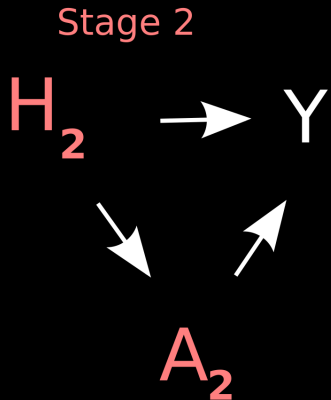
# Identifying the best treatment regime: multi-stage



DTR: treatment sequence  $A_1^{opt}, A_2^{opt}$

# Identifying the best treatment regime: multi-stage

$$H_2 = (X_1, A_1, X_2)$$





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Lots of methods available:

**Q-learning**

**MSMs**

**G-estimation**

**IPTW**

**dWOLS**

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**etc...**

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Expected outcome  
(to be maximized)



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- We might propose the following model

$$E[Y|X, A; \beta, \psi] = \beta_0 + \beta_1 \text{IOP} + A(\psi_0 + \psi_1 \text{IOP})$$

“Treat ( $A = 1$ ) if  $\psi_0 + \psi_1 \text{IOP} > 0$ ”

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- More generally, split outcome into two components:

$$\underbrace{E[Y|X, A; \beta, \psi]}_{\text{Expected outcome (to be maximized)}} = \underbrace{\text{Impact of patient history in the absence of treatment}}_{G(X; \beta)} + \underbrace{\gamma(X, A; \psi)}_{\text{Impact of treatment on outcome}}$$

- Simplifies focus: find  $A^{opt}$  that maximizes  $\gamma(X, A; \psi)$ .

# Identifying the best treatment regime

- Suppose the true outcome model is:

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$$E[Y|X, A; \beta, \psi] = G(X; \beta) + \gamma(X, A; \psi)$$

■ Three models to specify:

1. Blip model:  $\gamma(X, A; \psi)$ .
2. Treatment-free model:  $G(X; \beta)$ .
3. Treatment model:  $P(A = 1|X; \alpha)$ .

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- Estimate  $\psi$  via WOLS of  $Y$  on covariates in blip and treatment-free models, with weights  $w = |A - P(A = 1|X; \hat{\alpha})|$ .

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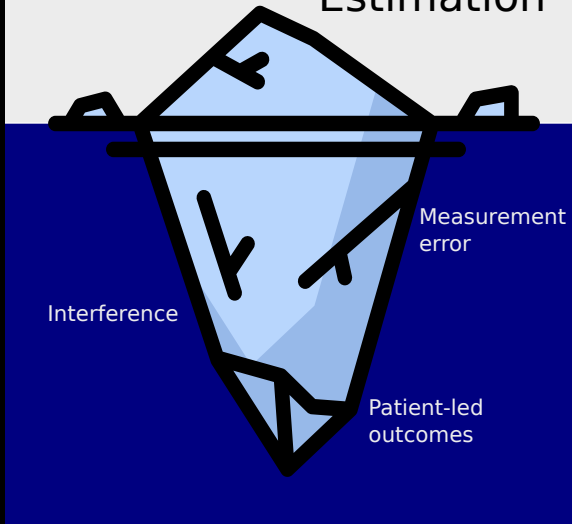
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- A weighted regression with weights  $w = |A - P(A = 1|X; \hat{\alpha})|$  will still yield consistent estimators of  $\psi_0, \psi_1$ .
- The estimators are “doubly robust”: consistent if at least one of the **treatment-free** or **treatment** components is correctly specified.
- The **blip** must always be correct.



So all we have to do is specify some models, estimate the model parameters, then choose the treatment that maximizes the expected outcome - easy!

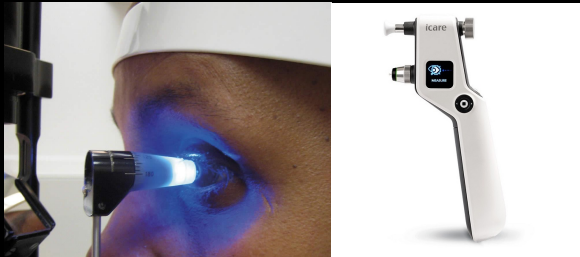
# Estimation

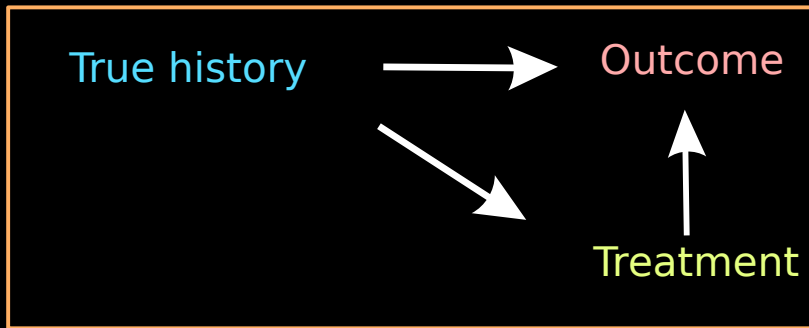


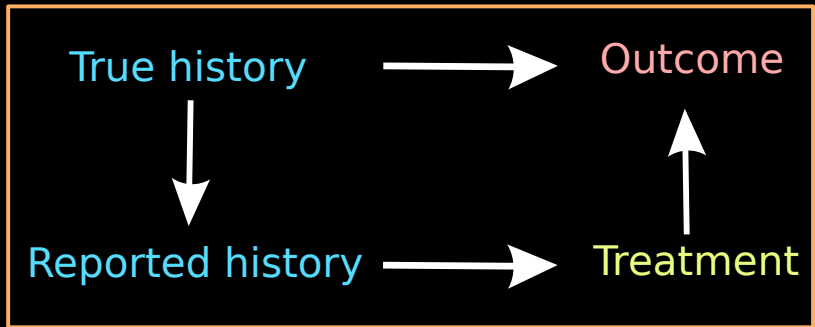
Target measurement: 'average' IOP.

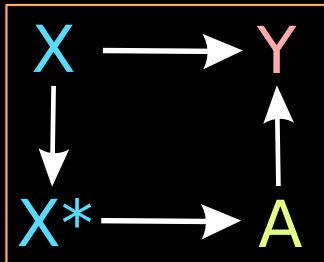
Observed measurement: 1-3 in-clinic readings within  $< 5$  minutes.

Some patients have access to more regular at-home tonometry.







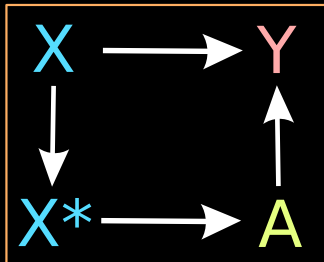


Estimation: suppose the true outcome model is:

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If we only observe  $X^*$ . What happens? What can we do about it?





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If we only observe  $X^*$ . What happens? What can we do about it?

Solution: 'correct' for the measurement error using additional data.

Assume: classical additive measurement error:

$$\text{Observed} = \text{True} + \text{Error}$$

$$X^* = X + U$$

$$U \sim N(0, \sigma_u^2); Y \perp X^* | X$$

Assume: replicate measurements available on at least some patients.

Patient	First IOP measurement	Second IOP measurement
1	16	15
2	20	16
3	21	17
4	16	16
5	15	18
...	...	...

Simple correction method: Regression Calibration.

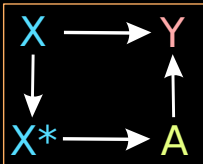
Principle:

1. Use additional data to estimate  $E[X|X^*, A] = X_{rc}$ .
2. Replace  $X$  with  $X_{rc}$  and carry out a standard analysis.
3. Adjust the resulting standard errors to account for the estimation in step 1.

# Identifying the best treatment regime

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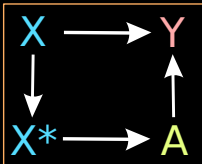
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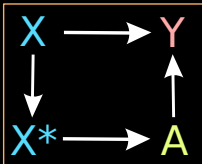
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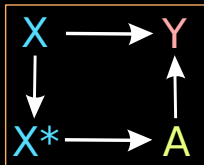
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where  $A$  depends on  $X^*$ .

- Establish (approximate) covariate balance in  $X_{rc}$  by regressing  $A$  on  $X_{rc}$ .



Suppose we conclude that our treatment rule should be:

“If 3-month average IOP  $> 15$  add secondary drop, otherwise,  
maintain current treatment regime.”



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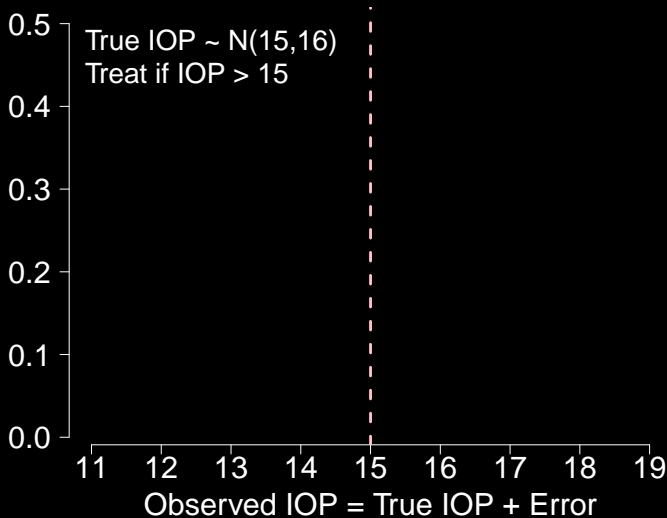
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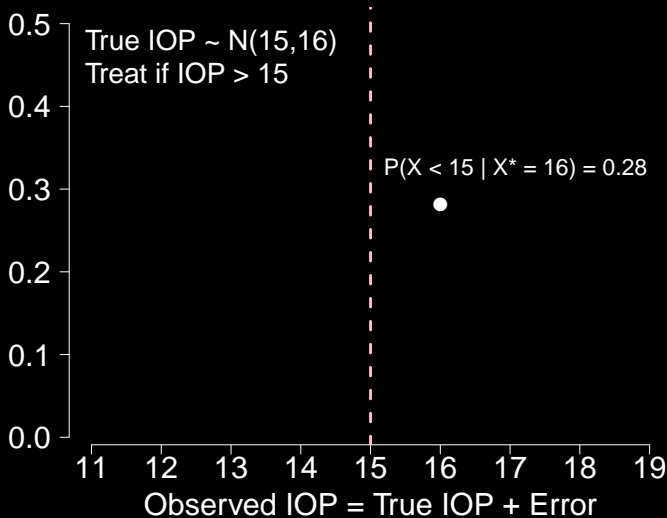
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What is  $P(X \leq 15 | X^* = 16)$ ?

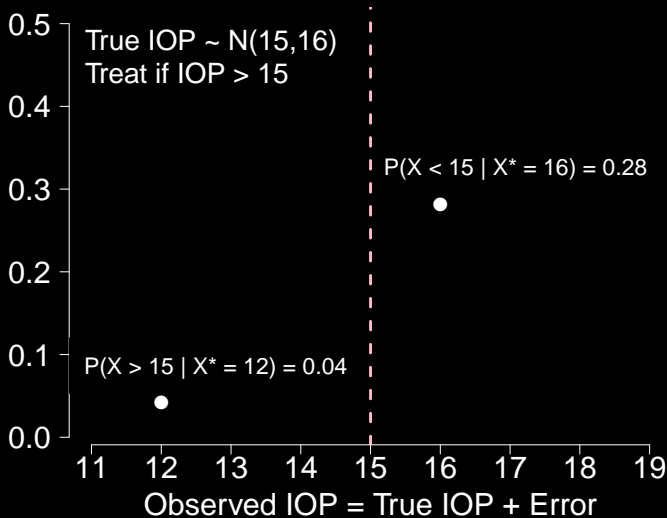
In some settings, results fairly intuitive:



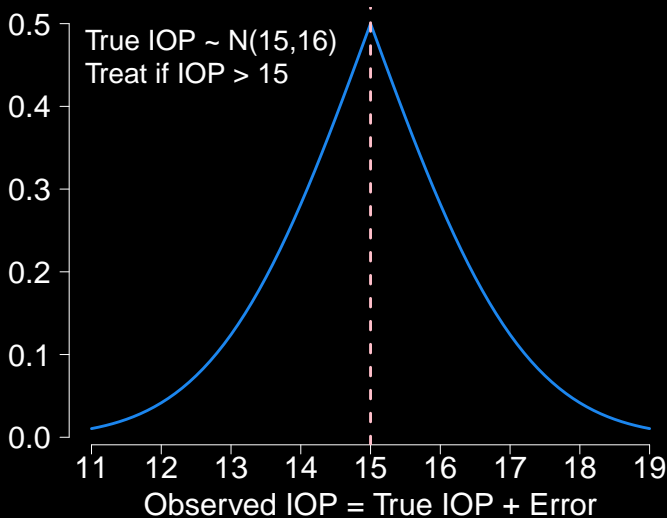
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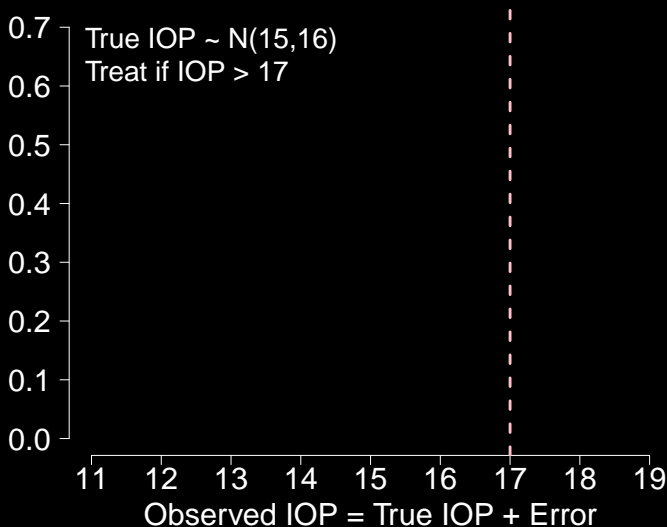
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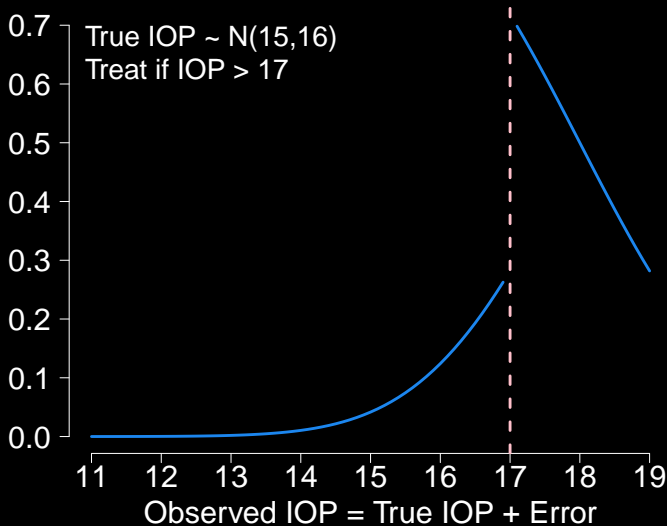
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In others, perhaps more of a surprise (to some):



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# “Isn't this just a prediction problem?”

Data availability will vary by study and by variable:

Scenario	Analysis	Application
1: “We never observe the truth.”	$X^*$	$X^*$

# “Isn't this just a prediction problem?”

What if error-free data are possible, but expensive?

Scenario	Analysis	Application
1: “We never observe the truth.”	$X^*$	$X^*$
2: “Past data are error-prone, but future data may not be.”	$X^*$	$X$
3: “Past data are not error-prone, but future data may be.”	$X$	$X^*$

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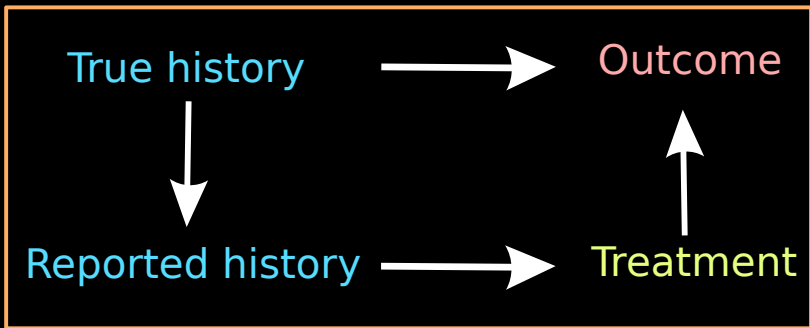
Only Scenario 4 is well-studied.

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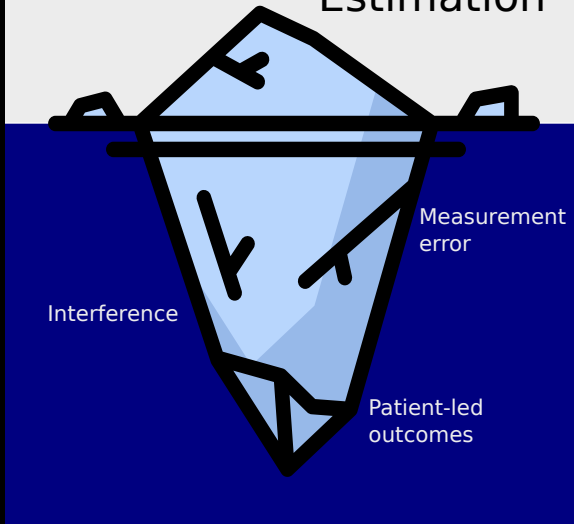
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Correcting for measurement error is at worst competitive with an analysis that ignores it completely.

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# Estimation



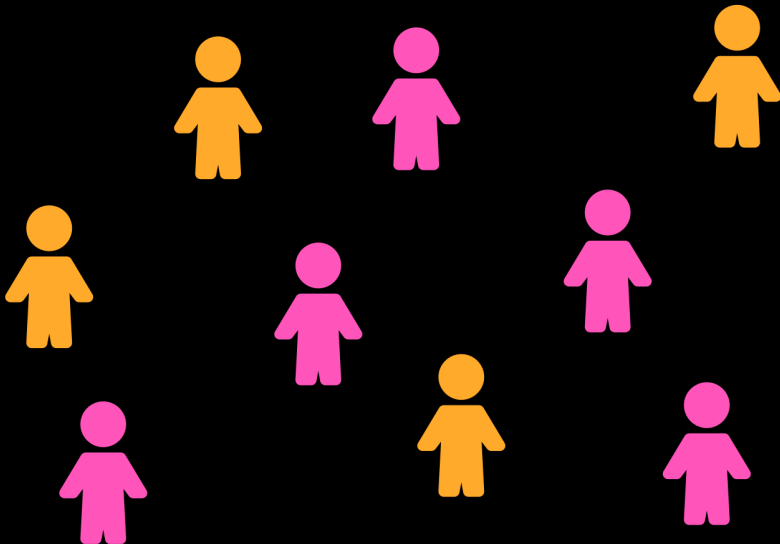


No man is an Island, entire of itself; every man is a piece of the Continent, a part of the main; if a clod be washed away by the sea, Europe is the less, as well as if a promontory were, as well as if a manor of thy friends or of thine own were; any man's death diminishes me, because I am involved in Mankind; And therefore never send to know for whom the bell tolls; It tolls for thee.

(John Donne)

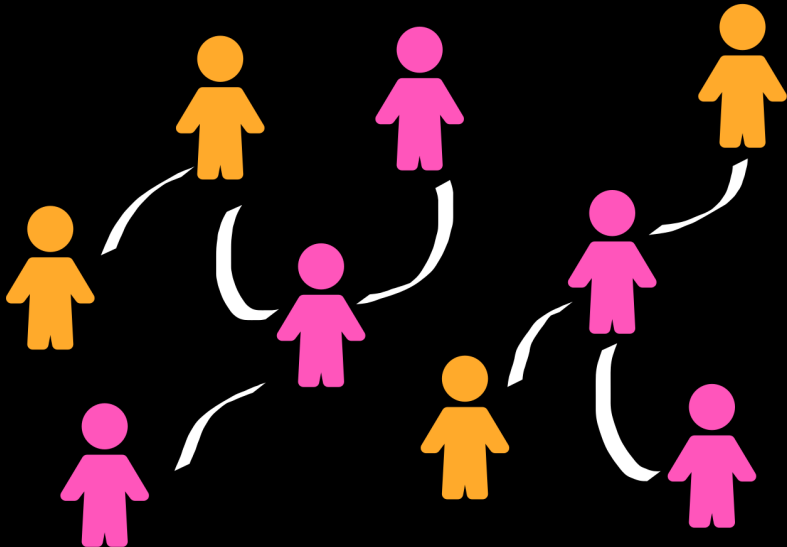
Interference: another patient's treatment doesn't affect my outcome.

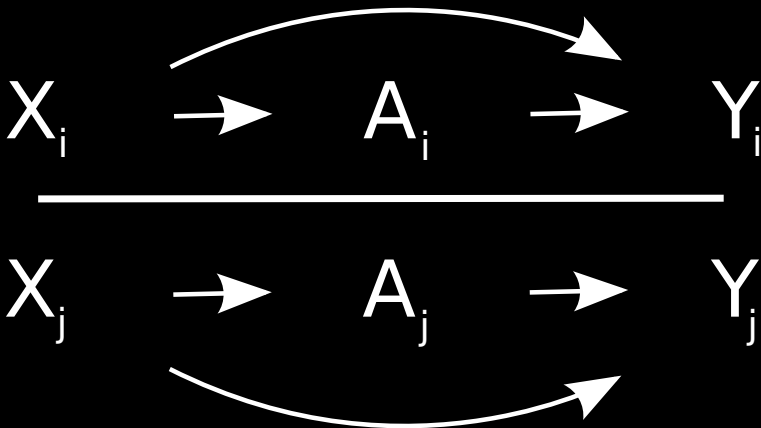
Common assumption: no interference (or 'spillover'):

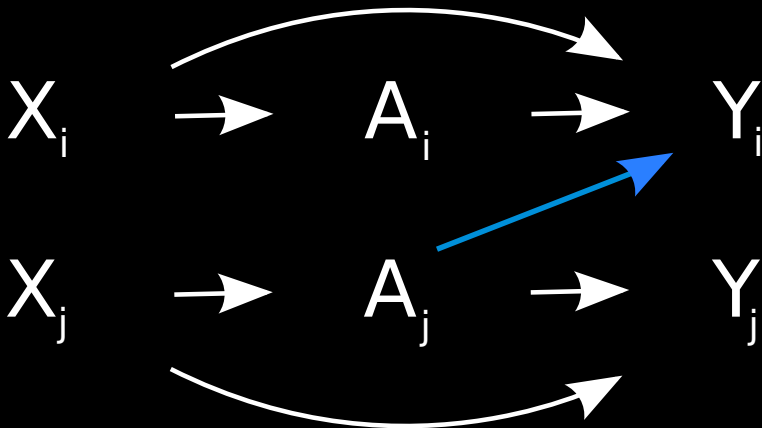




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# Identifying the best treatment regime

- No interference:

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“Treat ( $A_i = 1$ ) if  $\psi_0 + \psi_1 X_i > 0$ ”

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With interference: two ideas:

- (1) Add interaction terms:

$$E[Y|X, A; \beta, \psi] = \beta_0 + \beta_1 X_i + \beta_2 X_j + A_i(\psi_0 + \psi_1 X_i + \psi_2 A_j)$$

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- (2) Use ‘network propensity weights’: for each individual  $i$ , apply weights based on the probability their neighbour is treated.

Estimation is only half the battle.

Example: Alex and Blake share a household. The following table summarizes the ranking of the four possible combinations of a binary treatment:

Rank	Alex	Blake
1	o	-
2	-	o
3	o	o
4	-	-

Treatment: o

No treatment: -

- In what order do we prescribe treatments to patients?
- What if resources are limited?
- How do we balance the needs of the individual against the needs of the population?

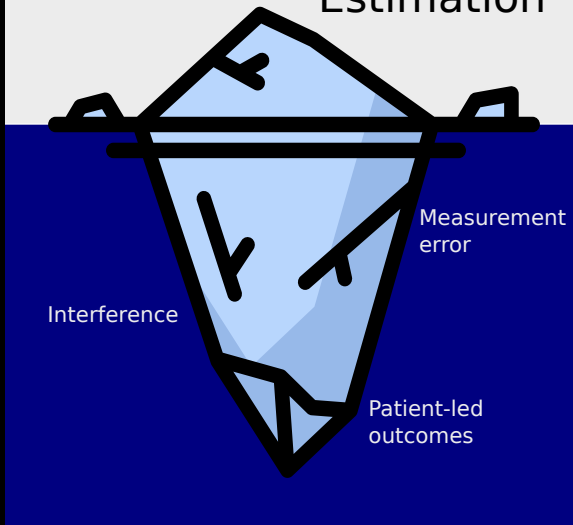
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# Estimation



Recall the treatment options for glaucoma:

- Lifestyle changes.
- Eye drops (numerous options).
- Surgery.

Treatments have various side effects, including:

- 'Minor': eyelash growth, iris discoloration.
- 'Major': additional vision loss.

Prior research tells us the probability of side effects, and the 'average' IOP decrease following each treatment.

Suppose you had the swallowing predicted effects:

Treatment	Iris discoloration	Eyelash growth	Vision loss	$E[\text{IOP change}]$
Drop A	80%	40%	1%	2
Drop B	90%	30%	1%	2

Which treatment do you choose? What influences your decision?

How do we elicit this information from patients?

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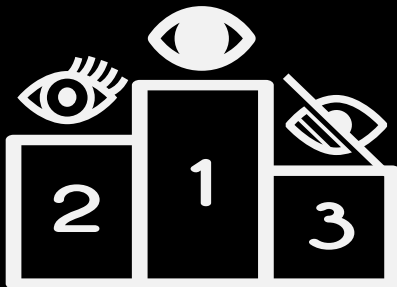
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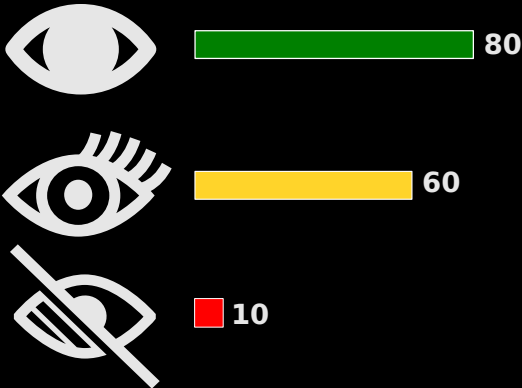
Patients can provide information to varying degrees:

- Ranking: “I would prefer iris discoloration to eyelash growth, and prefer both to additional vision loss.”



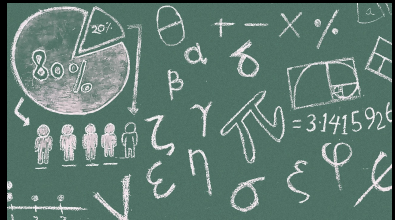
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- Utility functions: “I want to maximize this function of the possible outcomes.”



Methods from *multi-attribute decision making* can be applied to recommend treatment options.

We can also consider non-probabilistic properties of treatments, such as their cost.

Treatment	Cost	Iris discoloration	Eyelash growth	Vision loss	IOP change
Drop A	\$	80%	40%	1%	2
Drop B	\$\$	90%	30%	1%	2
Surgery	\$\$\$	1%	0%	5%	3

Such methods can be implemented via web-based applications:

<https://shiny.math.uwaterloo.ca/sas/mwallace/mapp/madm/>

**Assign Preference Scores to Attributes:**

Assign preference scores to the attributes such that a greater preference score represents a more valued attribute and a lower preference score represents a less valued attribute.

Attribute #1

0 10 20 30 40 50 60 70 80 90 100

Attribute #2

0 10 20 30 40 50 60 70 80 90 100

**Select Types:**

Benefit attributes are attributes such that a greater value is more preferable. Cost attributes are attributes such that a smaller value is more preferable.

Attribute #1

Benefit

Attribute #2

Benefit

**Enter Attribute Values:**

Enter the values for all treatment/attribute combinations.

	Attribute #1	Attribute #2
Treatment #1	0.80	0.40
Treatment #2	0.90	0.30

**Calculate**

Such methods can be implemented via web-based applications:

<https://shiny.math.uwaterloo.ca/sas/mwallace/mapp/madm/>

## Enter Attribute Values:

Enter the values for all treatment/attribute combinations.

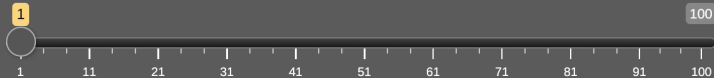
	Attribute #1	Attribute #2
Treatment #1	0.80	0.40
Treatment #2	0.90	0.30

Such methods can be implemented via web-based applications:

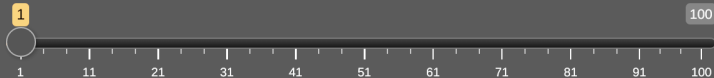
<https://shiny.math.uwaterloo.ca/sas/mwallace/mapp/madm/>

Assign preference scores to the attributes such that a greater preference score represents a more valued attribute and a lower preference score represents a less valued attribute.

**Attribute #1**

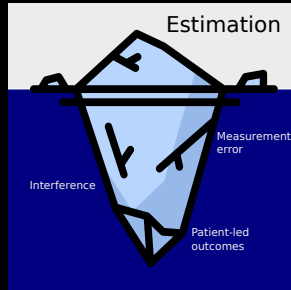


**Attribute #2**



# So where are we now?

- DTRs an important tool in precision medicine.
- Lots of estimation methods available, ease of implementation, and avoiding 'black boxes' a big challenge.
- Beyond estimation: lots of interesting, open, practical problems to work on.
  - Measurement error.
  - Interference.
  - Patient-led outcomes.
  - Model assessment.
  - Treatment quality.
  - etc. etc. etc.



# Acknowledgments



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Error



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Interference and  
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Patient-Led  
Outcomes



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Interference and  
Networks



- **dWOLS**: M. P. Wallace and E. E. M. Moodie (2015). Doubly-robust dynamic treatment regimen estimation via weighted least squares. *Biometrics* **71(3)** 636-644.
- **Precision Medicine and Measurement Error**: D. Spicker and M. P. Wallace (2020). Measurement error and precision medicine: error-prone tailoring covariates in dynamic treatment regimes. *Statistics in Medicine* **39(26)**  
<https://doi.org/10.1002/sim.8690>
- **Precision Medicine and Interference**: C. Jiang, M. E. Thompson, M. P. Wallace (2022). Dynamic treatment regimes with interference. *Canadian Journal of Statistics*. *In press*



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[mpwallace.github.io](https://github.com/mpwallace)