# STAT/BIOST 572 Update Student Presentation

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April 26, 2012

# Recap

- Transparent Parametrizations of Models for Potential Outcomes
  - Model used to estimate causal effects
  - Imperfect compliance
  - Separate the identifiable parameters from the unidentified

## General Potential Outcomes model

- Observed
  - *Z* = Assignment to treatment (Instrument)
  - X = Receipt/Exposure to treatment
  - Y = Response
- Unobserved (describing *potential outcomes*)
  - $t_X =$  Underlying compliance "type"
  - t<sub>Y</sub> = Underlying response "type"



# Compliance Types $(t_X)$

- Z = Assigned treatment
- $X_z$  = Potential receipt/exposure to treatment given assignment Z = z
- $X_z$  = observed X under the consistency axiom

$X_{z=0}$	$X_{z=1}$	Compliance Type $t_X$		
0	0	NT	Never Taker	
1	0	DE	Defier	
0	1	CO	Complier	
1	1	AT	Always Taker	

Table: Compliance types  $(t_X)$  based on potential outcomes

### **Potential Outcomes**

- Y<sub>xz</sub> = Potential response under receipt/exposure to treatment x, and treatment assignment z
- In a general model, Z may have an effect on Y, and  $t_Y$  would take  $16 = 2^{2^2}$  possible states.
- Under the Instrumental Variable model [Angrist et al., 1996], the effect of Z on Y is only through X.

# Response Types $(t_Y)$

- This is represented by the Exclusion Restriction:  $Y_{xz} = Y_{xz'} = Y_{x}$  for  $z, z' \in \{0, 1\}$ .
- $Y_{x.}$  = Potential response under receipt/exposure to treatment x, regardless of assigned treatment

<i>Y</i> <sub>0</sub> .	$Y_{1.}$	Response Type $t_Y$			
0	0	NR	Never Recover		
1	0	ΗU	Hurt		
0	1	ΗE	Helped		
1	1	AR	Always Recover		

Table: Response types  $(t_Y)$  based on potential outcomes under Instrumental Variable model

#### Instrumental Variable Model

- Exclusion Restriction
  - Effect of Assignment (Z) on Response (Y) is only through Receipt/Exposure (X)
- Causal estimand of interest: Average Causal Effect (ACE)

• 
$$ACE(X \rightarrow Y) = p(Helped) - p(Hurt)$$



# What's wrong with this approach?

- Non-identifiability
  - Distribution over potential outcomes p(t<sub>X</sub>, t<sub>Y</sub>) may only be partially-identified
  - Causal estimands of interest (e.g. ACE) would hence depend on parameters that are not fully-identified.
- How to get meaningful information about causal effects?
  - Pearl [2000] proposed bounds on ACE
  - Find the set of distributions  $p(t_X, t_Y)$  compatible with observed data
  - Find the minimum and maximum ACE

## Motivating Example

- Data from a double-blind placebo-controlled randomized trial
- Compliance and response were dichotomized
- There are no DEfiers and no Always Takers since  $Z = 0 \Rightarrow X = 0$

Ζ	X	y	count	Z	X	y	count
0	0	0	158	1	0	0	52
0	0	1	14	1	0	1	12
0	1	0	0	1	1	0	23
0	1	1	0	1	1	1	78

Table: Lipid data; there are two structural zeros

# Finding the ACE

• Bounds

$$\begin{split} & \left[ p(y=1 \mid z=1) - p(y=1 \mid z=0) \right] \\ & - \left[ p(y=1, x=0 \mid z=1) + p(y=0, x=1 \mid z=0) \right] \\ & \leq ACE(X \to Y) \leq \\ & \left[ p(y=1 \mid z=1) - p(y=1 \mid z=0) \right] \\ & + \left[ p(y=0, x=0 \mid z=1) + p(y=1, x=1 \mid z=0) \right] \end{split}$$

• From the data: (0.39, 0.78)

## Bayesian Approach

• Prior distribution over potential outcomes  $p(t_X, t_Y)$ 

- Reflects proportion of individuals in population that possess characteristics corresponding to  $(t_X, t_Y)$
- Dirichlet prior
- Posterior
  - Gibbs sampling to sample  $(t_X, t_Y)$  from resulting posterior

• Find  $ACE(X \rightarrow Y) = p(Helped \mid data) - p(Hurt \mid data)$ 

#### Posterior: Sensitive to prior?

- Uniform
  - Dir  $(1,\ldots,1) \rightarrow$  Dir  $(1,\ldots,1.2,1,0.8)$
- Unit

• Dir 
$$(\frac{1}{8}, \dots, \frac{1}{8}) \to \text{Dir} (1, \dots, \frac{3}{16}, \frac{1}{8}, \frac{1}{16})$$

• Such perturbation should not have large effect on posterior ACE

#### Posterior: Sensitive to prior?

#### Prior and posterior on ACE(X->Y) for Lipid data Uniform and perturbed uniform priors on potential outcomes

#### Prior and posterior on ACE(X->Y) for Lipid data Unit and perturbed unit priors on potential outcomes



#### **Transparent Parametrizations**

- Re-parameterize  $p(t_X, t_Y)$  into  $f(\theta, \zeta)$ 
  - $\theta$  = identifiable parameter (estimable from observed (X, Y, Z))
  - $\zeta = \text{non-identifiable parameter}$



#### **Transparent Parametrizations**



**Figure** 4: A graph representing the functional dependencies in the analysis of the simple IV model with no Always Takers or Defiers. Rectangular nodes are observed; oval nodes are unknown parameters. p(x=1|z=0) = 0, so p(y|x=1, z=0) is undefined, hence these nodes are omitted.

## Next Steps

- Inference based on posterior distribution
  - Work out the distribution of the observed data  $p(y, x \mid z)$  implied by the transparent model
  - Compute the posterior distribution of the parameters
  - Apply inequality restrictions ("truncate") by Monte-Carlo rejection sampling
  - Find ACE  $(X \rightarrow Y)$  as a function of the unidentified parameter(s)

## Next Steps

- Extend to general setting
  - Various assumptions or restrictions: 8 different models
  - Derive bounds on conditional causal contrasts e.g. Average Controlled Direct Effect  $ACDE_{NT}(x_0) = E(Y_{x=0,z=1} - Y_{x=0,z=0} | NT)$
  - Evaluate bounds conditional Intention-To-Treat effect e.g.  $ITT_{CO} = E(Y_{x=1,z=1} - Y_{x=0,z=0} | CO)$

#### References

- J.D. Angrist, G.W. Imbens, and D.B. Rubin. Identification of causal effects using instrumental variables. *Journal of the American Statistical Association*, 91(434):444–455, 1996.
- J. Pearl. *Causality: models, reasoning, and inference,* volume 47. Cambridge Univ Press, 2000.