

# Sensitivity analysis for principal ignorability violation in estimating complier and noncomplier average causal effects

Trang Nguyen

joint work with Liz Stuart, Dan Scharfstein, Betsy Ogburn

arXiv:2303.05032 | [trang.nguyen@jhu.edu](mailto:trang.nguyen@jhu.edu)

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# Outline

## Background

- Principal causal effects

- Principal ignorability (PI) – one identification strategy

- Sensitivity analysis for PI violation

## Mean-based sensitivity analysis: expanding options

- Allow different sensitivity parameters

- Accommodate different estimation methods

## Distribution-based sensitivity analysis: a new thing

- Method limitations and information use

- A method using full information (ongoing work)

## The noncompliance problem

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$Y(z)$	potential outcomes
$X$	baseline covariates
$S$	treatment received

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Might be interested in  
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but  
those who received  $\neq$  those who did not

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## Principal stratification

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Principal stratification (Frangakis & Rubin 2002)  
avoids this problem  
by creating a new pre-treatment variable  
based on potential treatment received  
and stratifying on it

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C	principal stratum, defined based on potential values $S(1), S(0)$ of S

Principal causal effects:

$$E[Y(1) - Y(0) \mid C = c]$$

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Two-sided noncompliance: 4 principal strata

$S(1)$	$S(0)$	$C$
1	1	always taker
1	0	complier
0	1	defier
0	0	never taker

$Z$  treatment assigned  
 $Y$  outcome  
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One-sided noncompliance: 2 principal strata

$S(1)$	$S(0)$	$C$
1	0	complier
0	0	noncomplier

i.e.,  $C = S(1)$

$Z$	treatment assigned
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## Examples

Our focus is one-sided noncompliance

target: (non)complier average causal effects (CACE and NACE)

JOBS II for unemployed workers

- ▶ Z: week-long training on job search and mental health
- ▶ S: attending training

Experience Corps for the elderly

- ▶ Z: facilitated program for volunteering to help kids in school
- ▶ S: volunteering above a certain number of hours

## Identification challenge: $C$ is not observed under control

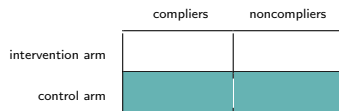
	compliers	noncompliers
intervention arm		
control arm		

## Two major identification strategies

		compliers	noncompliers
exclusion restriction/IV	intervention arm		
	control arm		

		compliers	noncompliers
principal ignorability	intervention arm		
	control arm		

## Principal ignorability (PI)



$$Y \perp\!\!\!\perp C \mid X, Z = 0$$

ie within covariate levels, compliers and noncompliers share the same outcome distribution under control

## Principal ignorability (PI)

	compliers	noncompliers
intervention arm	$\mu_{11}(X)$	$\mu_{10}(X)$
control arm	$\mu_{01}(X)$	$\mu_{00}(X)$

$$Y \perp\!\!\!\perp C \mid X, Z = 0$$

or

$$\underbrace{E[Y \mid X, Z = 0, C = 1]}_{\mu_{01}(X)} = \underbrace{E[Y \mid X, Z = 0, C = 0]}_{\mu_{00}(X)}$$

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Combined with treatment assignment ignorability, PI identifies CACE, NACE:

$$E[Y(1) - Y(0) \mid C = c] = \frac{E\{[\mu_{1c}(X) - \kappa_0(X)] \pi_c(X)\}}{E[\pi_c(X)]}$$

where

$$\begin{aligned} \kappa_0(X) &:= E[Y \mid X, Z = 0] && \text{(mixture outcome mean)} \\ \pi_c(X) &:= P(C = c \mid X, Z = 1) && \text{(principal score)} \end{aligned}$$

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PI is untestable

Need sensitivity analyses

## Prior sens analysis method that inspired this work

Ding and Lu (2017) use a mean ratio sensitivity parameter

$$\frac{\mu_{01}(X)}{\mu_{00}(X)} = \rho$$

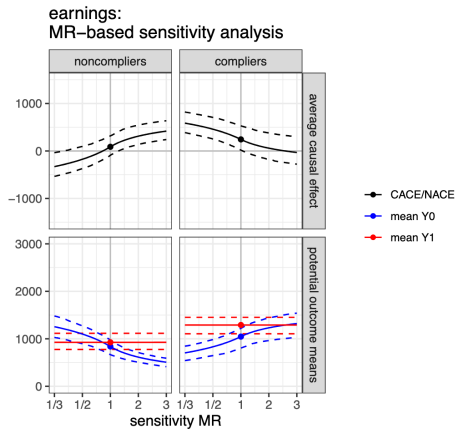
and modify a PI-based weighting estimator to incorporate  $\rho$

See also Jiang, Yang and Ding (2022)



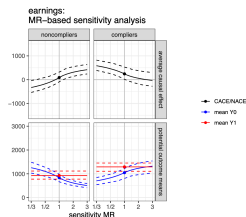
# Example of MR-based sens analysis

earnings in JOBS II



# Example of MR-based sens analysis

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other outcomes for which MR param not ideal

- ▶ JOBS II: having a job (binary), depressive symptoms (bounded)
- ▶ Experience Corps: generativity (bounded)

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## A range of sens assumptions with different sens params

PI:  $\mu_{01}(X) = \mu_{00}(X).$

sens-MR:  $\frac{\mu_{01}(X)}{\mu_{00}(X)} = \rho,$

sens-OR:  $\frac{\mu_{01}(X)/[1 - \mu_{01}(X)]}{\mu_{00}(X)/[1 - \mu_{00}(X)]} = \psi,$

sens-GOR:  $\frac{[\mu_{01}(X) - l]/[h - \mu_{01}(X)]}{[\mu_{00}(X) - l]/[h - \mu_{00}(X)]} = \psi$

where  $l$  and  $h$  are the outcome lower and upper bounds,

sens-SMD:  $\frac{\mu_{01}(X) - \mu_{00}(X)}{\sqrt{[\sigma_{01}^2(X) + \sigma_{00}^2(X)]/2}} = \eta$

where  $\sigma_{0c}^2(X) := \text{var}(Y | X, Z = 0, C = c),$

for some range of  $\rho$ ,  $\psi$  or  $\eta$  that is considered plausible.

## Identification

sens-MR and sens-GOR result in point identification of CACE, NACE

because they help solve the mixture equation

$$\pi_1(X)\mu_{01}(X) + \pi_0(X)\mu_{00}(X) = \kappa_0(X).$$

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sens-SMD obtains bounds for CACE, NACE

bounds are narrowed if also assume  $1/k \leq \frac{\sigma_{01}^2(X)}{\sigma_{00}^2(X)} \leq k$  for some  $k > 1$

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and reduce to point identification if assume  $\sigma_{01}^2(X) = \sigma_{00}^2(X)$  (aka sens-SMD<sub>e</sub>)

in all cases, effect identification is via identification of  $\mu_{0c}(X)$

by a function of sens param,  $\pi_c(X)$ ,  $\kappa_0(X)$  (and  $\text{var}(Y | X, Z = 0)$  w/ sens-SMD)

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Sens analysis = a modification of main analysis



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Sens analysis = a modification of main analysis

▶ Type A ( $\approx$  outcome regression estimators)

- ▶ estimates  $\kappa_0(X)$  to first estimate effects conditional on covariates and then aggregates them to estimate CACE/NACE, eg

$$\frac{\sum_{i=1}^n \hat{\pi}_c(X_i) [\hat{\mu}_{1c}(X_i) - \hat{\kappa}_0(X_i)]}{\sum_{i=1}^n \hat{\pi}_c(X_i)}, \quad \frac{\sum_{i=1}^n \frac{Z_i}{\hat{e}(X_i, Z_i)} I(C_i = c) [Y_i - \hat{\kappa}_0(X_i)]}{\sum_{i=1}^n \frac{Z_i}{\hat{e}(X_i, Z_i)} I(C_i = c)}$$

- ▶ sens analysis technique: replace  $\kappa_0(X)$  by the identification result of  $\mu_{0c}(X)$  under the sens assumption

## Sens analysis techniques to go with 3 types of PI-based estimators

Sens analysis = a modification of main analysis

▶ Type B ( $\approx$  influence function based estimators)

- ▶ write the CACE/NACE as

$$\frac{\nu_{1c} - \nu_{0c}^{\text{PI}}}{\pi_c}$$

where  $\nu_{zc} := E[\pi_c(X)\mu_{zc}(X)]$ ,  $\nu_{0c}^{\text{PI}} := E[\pi_c(X)\kappa_0(X)]$ ,  $\pi_c := E[\pi_c(X)]$

- ▶ a type B estimator can be expressed as combination of IF-based estimators of  $\pi_c$ ,  $\nu_{1c}$  and  $\nu_{0c}^{\text{PI}}$

$$\frac{\hat{\nu}_{1c,\text{IF}} - \hat{\nu}_{0c,\text{IF}}^{\text{PI}}}{\hat{\delta}_{c,\text{IF}}}$$

- ▶ sens analysis technique: replace  $\hat{\nu}_{0c,\text{IF}}^{\text{PI}}$  with an IF-based estimator of  $\nu_{0c}$  under the sens assumption

## Sens analysis techniques to go with 3 types of PI-based estimators

Sens analysis = a modification of main analysis

- ▶ Type C ( $\approx$  other/weighting estimators)
  - ▶ an example is the pure weighting estimator

$$\frac{\sum_{i=1}^n \frac{Z_i}{\hat{e}(X_i, Z_i)} I(C_i = c) Y_i}{\sum_{i=1}^n \frac{Z_i}{\hat{e}(X_i, Z_i)} I(C_i = c)} - \frac{\sum_{i=1}^n \frac{1-Z_i}{\hat{e}(X_i, Z_i)} \hat{\pi}_c(X_i) Y_i}{\sum_{i=1}^n \frac{1-Z_i}{\hat{e}(X_i, Z_i)} \hat{\pi}_c(X_i)}$$

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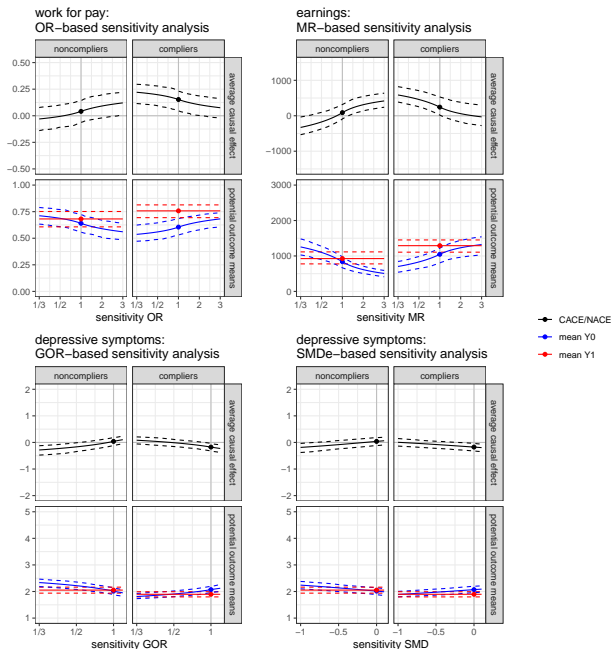
- ▶ no general sens analysis technique
    - ▶ sens-MR: scale  $Y$  in control units by a simple function of  $\rho$  and  $\pi_c(X)$

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# JOBS II results



## Some other things we noticed/figured out

- ▶ Partial loss of multiple robustness

because  $\mu_{0c}(X)$  is a function of  $\pi_c(X)$  and  $\kappa_0(X)$

- ▶ A pattern of finite-sample bias for the sens analysis where effect estimates are less extreme than should be

because  $E[Y_0 | C = c] = \frac{E[\pi_c(X)\mu_{0c}(X)]}{E[\pi_c(X)]}$  is a weighted average where the function being averaged depends on the weight

- ▶ If IF-based nonparametric estimation: Rate conditions on nuisance estimation for the estimator to be root-n consistent

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## A general limitation of above methods

is the risk of contradicting the observed data distribution

Assumption	Risk level	Info used from the observed mixture outcome distribution
sens-MR	greatest risk	mean $\kappa_0(X) = E[Y   X, Z = 0]$ lower bound (0)
sens-GOR (for nonbinary $Y$ )	less risk	mean both bounds
sens-SMD	less risk	mean variance $\text{var}(Y   X, Z = 0)$
sens-OR (binary $Y$ )	no risk	full distribution (as mean = probability)

For nonbinary  $Y$ , to avoid contradicting the observed data distribution,

sens analysis needs to be fully informed by  $P(Y | X, Z = 0)$

## Constructing that sens analysis: foundation

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(Everything here conditions on  $X, Z = 0$ ,  
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Principal scores

Recall/recast  $\pi_c(X) = P(C = c \mid X, Z = 0)$  (outcome-agnostic)

Now define  $\tilde{\pi}_c(X, Y) = P(C = c \mid X, Z = 0, Y)$  (outcome-specific)

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- ▶  $E[\tilde{\pi}_c(X, Y) \mid X, Z = 0] = \pi_c(X)$
- ▶ If  $Y$  and  $C$  are dependent,  $\tilde{\pi}_c(X, Y)$  is a function of  $Y$  in addition to  $X$

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(Side note: Very different from the unobserved confounding problem b/c of the observed mixture.  
Hence exponential tilting doesn't work, except for a binary outcome, in which case = sens-OR.)

## Constructing that sens analysis: 2 steps

Use shorthand  $\tilde{\pi}_1$  for  $\tilde{\pi}_1(X, Y)$

Step 1: Assume a distribution for  $\tilde{\pi}_1$  with mean  $\pi_1(X)$  that allows  $\tilde{\pi}_1$  to vary, indexed by a dispersion param

Step 2: Connect  $Y$  to  $\tilde{\pi}_1$  to induce  $Y$ - $C$  dependence

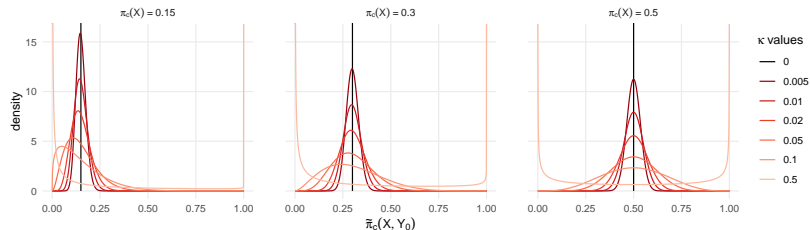
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- ▶ use the beta distribution (idea borrowed from Victor Veitch, 2020)

$$\tilde{\pi}_1 \mid X, Z = 0 \sim \text{Beta} \left( \pi_1(X) \frac{1 - \kappa}{\kappa}, \pi_0(X) \frac{1 - \kappa}{\kappa} \right)$$





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- ▶ use quantile-to-quantile mapping between distributions of  $Y$  and of  $\tilde{\pi}_1$  (given  $X, Z = 0$ )
  - ▶ same-quantiles: positive  $Y$ - $C$  association
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The sens param: the “signed”  $\kappa$

## Identification and estimation

Identification:

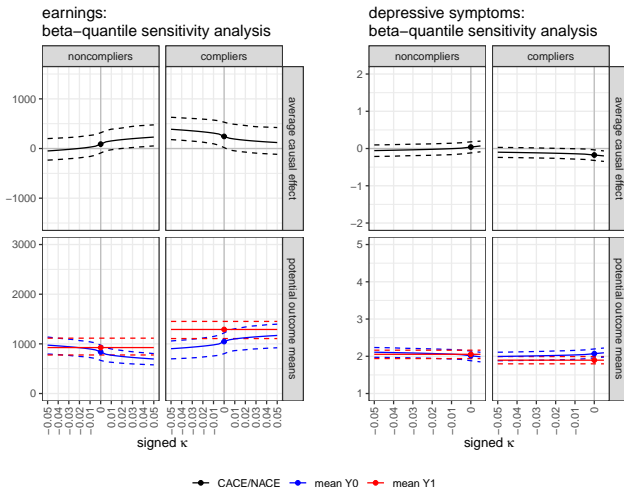
$$\mu_{0c}(X) = \frac{E[\tilde{\pi}_c Y \mid X, Z = 0]}{E[\tilde{\pi}_c \mid X, Z = 0]},$$

where  $\tilde{\pi}_c$  and  $Y$  are quantile-to-quantile connected

Estimation:

- ▶ estimate  $P(Y \mid X, Z = 0)$  so can compute quantiles
- ▶ then estimate  $\mu_{0c}(X)$  using numerical integration

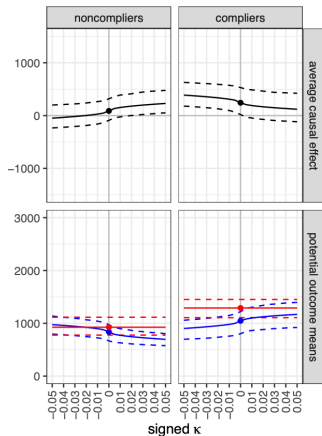
# Example: JOBS II preliminary results



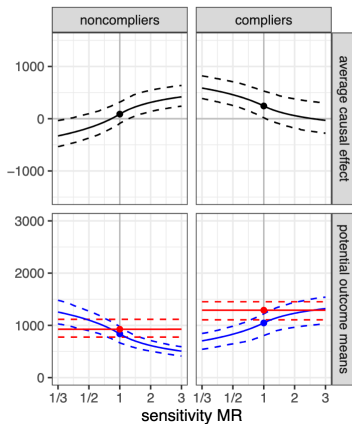
Results are less extreme than mean-based sens analysis

# earnings: distribution-based (left), mean-based (right)

earnings:  
beta-quantile sensitivity analysis

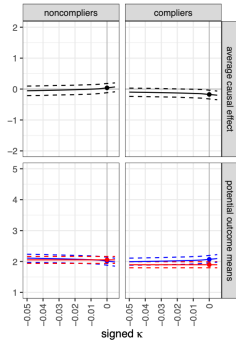


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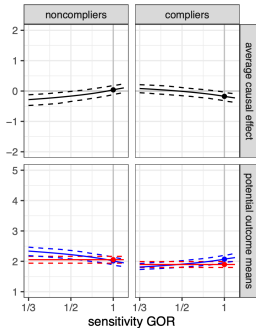


# depressive symptoms: distribution-based (left), mean-based (middle and right)

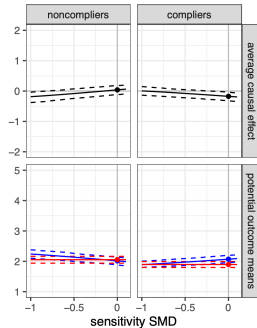
depressive symptoms:  
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depressive symptoms:  
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