

Lecture 17

Sensitivity Analysis

Outline

- Sensitivity analysis
 - Bound under no assumptions
 - Bound for the smoking example
 - A model-based approach
 - Rosenbaum sensitivity analysis

Sensitivity analysis

- Most often, validity of unconfoundedness can not be easily checked. Alternatively, one should check sensitivity of a causal analysis to unconfoundedness
- **Sensitivity analysis** aims at assessing the bias of causal effect estimates when the unconfoundedness assumption is assumed to fail in some specific and meaningful ways
- Sensitivity is different from testing – unconfoundedness is intrinsically non-testable, more of a “insurance” check
- Sensitivity analysis in causal inference dates back to the Hill-Fisher debate on causation between smoking and lung cancer, and first formalized in Cornfield (1959, JNCI)

Bounds under no assumptions

- Consider a simple case where: 1. no covariates; 2. binary outcome
- We are interested in the ATE

$$\tau_{\text{sp}} = \mu_t - \mu_c,$$

where

$$\mu_t = \mathbb{E}[Y_i(1)] = p \cdot \mu_{t,1} + (1 - p) \cdot \mu_{t,0},$$

and

$$\mu_c = \mathbb{E}[Y_i(0)] = p \cdot \mu_{c,1} + (1 - p) \cdot \mu_{c,0}.$$

$$\mu_{t,1} = \mathbb{E}[Y_i(1)|W_i = 1]$$

$$\mu_{t,0} = \mathbb{E}[Y_i(1)|W_i = 0]$$

$$\mu_{c,1} = \mathbb{E}[Y_i(0)|W_i = 1]$$

$$\mu_{c,0} = \mathbb{E}[Y_i(0)|W_i = 0]$$

$$p = P(W_i = 1)$$

Identifiable
from
observed
data

Bound the unknown $\mu_{t,0}$ and $\mu_{c,1}$ by $[0, 1]$ as the outcome is binary

Bounds under no assumptions

- So we get the bounds

$$\begin{aligned}\mu_t &\in [p \cdot \mu_{t,1}, p \cdot \mu_{t,1} + (1 - p)] \\ \mu_c &\in [(1 - p) \cdot \mu_{c,0}, (1 - p) \cdot \mu_{c,0} + p]\end{aligned}$$

- The the bound of ATE $\tau = \tau_{sp} = \mu_t - \mu_c$ is

$$\tau \in [p \cdot \mu_{t,1} - (1 - p) \cdot \mu_{c,0} - p, p \cdot \mu_{t,1} + (1 - p) - (1 - p) \cdot \mu_{c,0}]$$

- Unfortunately, because we don't have any assumptions at all, this bound is not very informative

- $\tau^{upper} - \tau^{lower}$

$$= p\mu_{t,1} + (1 - p) - (1 - p)\mu_{c,0} - p\mu_{t,1} + (1 - p)\mu_{c,0} + p \equiv 1$$

- By definition, $\tau^{upper} \leq 1$ and $\tau^{lower} \geq -1$, bound always covers 0
 - Better than the naive bound $[-1, 1]$

The Imbens-Rubin-Sacerdote lottery data

[Estimating the effect of unearned income on labor earnings, savings, and consumption: Evidence from a survey of lottery players. *American economic review*, 2001]

- Goal: Estimate magnitude of lottery prizes (unearned income) on economic behavior, including labor supply, consumption and savings
- Data collection:
 - “Winners”: individuals who had played and won large sums of money in the Massachusetts lottery
 - “Losers”: individuals who played the lottery and had won only small prizes
- We analyze a subset of $N_t = 259$ and $N_c = 237$ individuals with complete answers

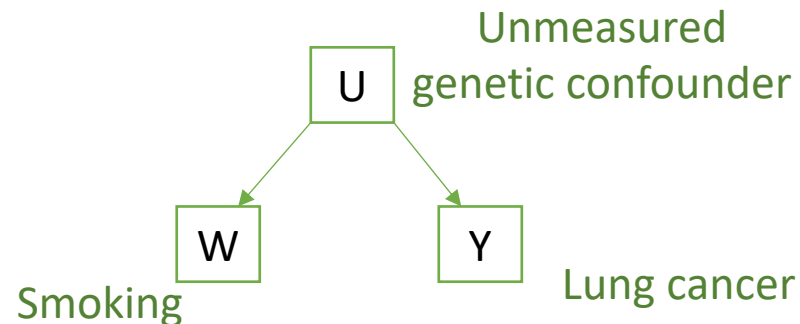
Result on the lottery data

- Binary outcome: whether the earning after treatment is positive or not
- Estimated quantities: $\hat{p} = \frac{N_t}{N} = 0.4675$, $\hat{\mu}_{t,1} = \bar{Y}_t^{\text{obs}} = 0.4106$ and $\hat{\mu}_{c,0} = \bar{Y}_c^{\text{obs}} = 0.5349$
- Plug in these quantities into our bound:
$$\tau \in [-0.56, 0.44]$$
- The two-sample difference estimate: $\bar{Y}_t^{\text{obs}} - \bar{Y}_c^{\text{obs}} = -0.124$

Sensitivity analysis bound: a more useful example

The smoking on lung cancer effect example (Cornfield et al. 1959 JNCI)

- Fisher argued the association between smoking and lung cancer may be due to a common gene that causes both



- Observed association between smoking and lung cancer
 - Risk ratio

$$RR_{WY} = \frac{P[Y_i^{\text{obs}} = 1 | W_i = 1]}{P[Y_i^{\text{obs}} = 1 | W_i = 0]}$$

- Observed risk ratio $RR_{WY} \approx 9$
- Can this be fully explained by U ?

Sensitivity analysis bound: a more useful example

- Assume that U_i are binary variables
- Define

$$p_0 = P[U_i = 1|W_i = 0], \quad p_1 = P[U_i = 1|W_i = 1]$$

- $RR_{WU} = \frac{p_1}{p_0}$
- If there is no causal effect of smoking on lung cancer, then $Y_i(0) = Y_i(1) = Y_i$
 $P[Y_i^{\text{obs}} = 1|W_i = 0, U_i = 0] = P[Y_i^{\text{obs}} = 1|W_i = 1, U_i = 0] = P[Y_i = 1|U_i = 0] = r_0,$
 $P[Y_i^{\text{obs}} = 1|W_i = 0, U_i = 1] = P[Y_i^{\text{obs}} = 1|W_i = 1, U_i = 1] = P[Y_i = 1|U_i = 1] = r_1$
- Then we have

$$RR_{WY} = \frac{P[Y_i^{\text{obs}} = 1|W_i = 0, U_i = 0]}{P[Y_i^{\text{obs}} = 1|W_i = 0, U_i = 1]} = \frac{r_0(1 - p_1) + r_1p_1}{r_0(1 - p_0) + r_1p_0}$$

Sensitivity analysis bound: a more useful example

$$RR_{WY} = \frac{r_0(1 - p_1) + r_1p_1}{r_0(1 - p_0) + r_1p_0}, \quad RR_{WU} = \frac{P[U_i = 1|W_i = 1]}{P[U_i = 1|W_i = 0]} = \frac{p_1}{p_0}$$

- As $p_1 \geq p_0$ because we observe $RR_{WY} > 1$, then (from some math)

$$RR_{WY} = \frac{r_0(1 - p_1) + r_1p_1}{r_0(1 - p_0) + r_1p_0} \leq \frac{p_1}{p_0} = RR_{WU}$$

- Cornfield showed that if Fisher is right, we have $RR_{WU} \geq RR_{WY} \approx 9$
- Such a genetic confounder might be too strong to be realistic
- If we believe that such genetic confounder does not exist, then smoking should have a causal effect on lung cancer

Another sensitivity analysis idea: base on a model

Idea:

$$W_i \perp (Y_i(0), Y_i(1)) \mid \mathbf{X}_i, U_i$$

observed
↓
unobserved ↙

- How sensitive is our estimate of causal effect to the presence of U_i ?
- A model-based approach (Rosenbaum and Rubin, 1983 JRSS-B)
 - Consider the scenario that $Y_i(w)$ is binary
 - Assume that the unmeasured confounding is binary
 - Build the following model

$$U_i \sim \text{Bernoulli}(q)$$
$$\text{logit}(P[W_i = 1 \mid \mathbf{X}_i, U_i]) = \gamma_0 + \mathbf{X}_i^T \boldsymbol{\kappa} + \gamma_1 U_i$$
$$\text{logit}(P[Y_i(0) = 1 \mid \mathbf{X}_i, U_i]) = \beta_0 + \mathbf{X}_i^T \mathbf{b}_0 + \beta_0 U_i$$
$$\text{logit}(P[Y_i(1) = 1 \mid \mathbf{X}_i, U_i]) = \alpha_0 + \mathbf{X}_i^T \mathbf{b}_1 + \alpha_1 U_i$$

Propensity score model

Outcome regression model

Another sensitivity analysis idea: base on a model

$$U_i \sim \text{Bernoulli}(q)$$

$$\text{logit}(P[W_i = 1 | \mathbf{X}_i, U_i]) = \gamma_0 + \mathbf{X}_i^T \boldsymbol{\kappa} + \gamma_1 U_i$$

Propensity score model

$$\text{logit}(P[Y_i(0) = 1 | \mathbf{X}_i, U_i]) = \beta_0 + \mathbf{X}_i^T \mathbf{b}_0 + \beta_1 U_i$$

Outcome regression model

$$\text{logit}(P[Y_i(1) = 1 | \mathbf{X}_i, U_i]) = \alpha_0 + \mathbf{X}_i^T \mathbf{b}_1 + \alpha_1 U_i$$

- Sensitivity parameters: $(q, \gamma_1, \beta_1, \alpha_1)$
- Sensitivity parameters can not be estimated as unmeasured confounder U_i is unobserved
- Sensitivity analysis: Set the sensitivity parameters to different values and see how estimates of causal effects change

An example of calculation

- Consider the simpler case where there is no \mathbf{X}_i

- Our observed data provides estimates of $p = \mathbb{E}(W_i) = \mathbb{P}(W_i = 1)$, $\mu_{t,1} = \mathbb{E}[Y_i^{obs} | W_i = 1]$ and

$$\mu_{c,0} = \mathbb{E}[Y_i^{obs} | W_i = 0]$$

$$p = q \cdot \frac{\exp(\gamma_0 + \gamma_1)}{1 + \exp(\gamma_0 + \gamma_1)} + (1 - q) \cdot \frac{\exp(\gamma_0)}{1 + \exp(\gamma_0)}$$

$$\begin{aligned} \mu_{t,1} &= \Pr(U_i = 1 | W_i = 1) \cdot \mathbb{E}[Y_i(1) | W_i = 1, U_i = 1] \\ &\quad + (1 - \Pr(U_i = 1 | W_i = 1)) \cdot \mathbb{E}[Y_i(1) | W_i = 1, U_i = 0] \\ &= \frac{q \cdot \frac{\exp(\gamma_0 + \gamma_1)}{1 + \exp(\gamma_0 + \gamma_1)}}{q \cdot \frac{\exp(\gamma_0 + \gamma_1)}{1 + \exp(\gamma_0 + \gamma_1)} + (1 - q) \cdot \frac{\exp(\gamma_0)}{1 + \exp(\gamma_0)}} \cdot \frac{\exp(\alpha_0 + \alpha_1)}{1 + \exp(\alpha_0 + \alpha_1)} \\ &\quad + \frac{(1 - q) \cdot \frac{\exp(\gamma_0)}{1 + \exp(\gamma_0)}}{q \cdot \frac{\exp(\gamma_0 + \gamma_1)}{1 + \exp(\gamma_0 + \gamma_1)} + (1 - q) \cdot \frac{\exp(\gamma_0)}{1 + \exp(\gamma_0)}} \cdot \frac{\exp(\alpha_0)}{1 + \exp(\alpha_0)}, \end{aligned}$$

$$\begin{aligned} \mu_{c,0} &= \frac{q \cdot \frac{1}{1 + \exp(\gamma_0 + \gamma_1)}}{q \cdot \frac{1}{1 + \exp(\gamma_0 + \gamma_1)} + (1 - q) \cdot \frac{1}{1 + \exp(\gamma_0)}} \cdot \frac{\exp(\beta_0 + \beta_1)}{1 + \exp(\beta_0 + \beta_1)} \\ &\quad + \frac{(1 - q) \cdot \frac{1}{1 + \exp(\gamma_0)}}{q \cdot \frac{1}{1 + \exp(\gamma_0 + \gamma_1)} + (1 - q) \cdot \frac{1}{1 + \exp(\gamma_0)}} \cdot \frac{\exp(\beta_0)}{1 + \exp(\beta_0)}. \end{aligned}$$

Given any value of $(q, \gamma_1, \beta_1, \alpha_1)$, we can solve the three equations to estimate $(\gamma_0, \beta_0, \alpha_0)$

$$\begin{aligned} U_i &\sim \text{Bernoulli}(q) \\ \text{logit}(P[W_i = 1 | \mathbf{X}_i, U_i]) &= \gamma_0 + \gamma_1 U_i \\ \text{logit}(P[Y_i(0) = 1 | \mathbf{X}_i, U_i]) &= \beta_0 + \beta_1 U_i \\ \text{logit}(P[Y_i(1) = 1 | \mathbf{X}_i, U_i]) &= \alpha_0 + \alpha_1 U_i \end{aligned}$$

An example of calculation

$$U_i \sim \text{Bernoulli}(q)$$

$$\text{logit}(P[W_i = 1|U_i]) = \gamma_0 + \gamma_1 U_i$$

$$\text{logit}(P[Y_i(0) = 1|U_i]) = \beta_0 + \beta_1 U_i$$

$$\text{logit}(P[Y_i(1) = 1|U_i]) = \alpha_0 + \alpha_1 U_i$$

- Consider the simpler case where there is no \mathbf{X}_i
- Our observed data provides estimates of $p = \mathbb{E}(W_i) = \mathbb{P}(W_i = 1)$, $\mu_{t,1} = \mathbb{E}[Y_i^{obs}|W_i = 1]$ and $\mu_{c,0} = \mathbb{E}[Y_i^{obs}|W_i = 0]$
- Given any value of $(q, \gamma_1, \beta_1, \alpha_1)$, we can solve the three equations to estimate $(\gamma_0, \beta_0, \alpha_0)$
- Then given the value of both $(q, \gamma_1, \beta_1, \alpha_1)$ and $(\hat{\gamma}_0, \hat{\beta}_0, \hat{\alpha}_0)$, we can estimate $\mu_{t,0} = \mathbb{E}[Y_i(1)|W_i = 0]$ and $\mu_{c,1} = \mathbb{E}[Y_i(0)|W_i = 1]$
- The average treatment effect will be

$$\tau_{sp} = \mu_t - \mu_c = p \cdot (\mu_{t,1} - \mu_{c,1}) + (1 - p) \cdot (\mu_{t,0} - \mu_{c,0}).$$

Sensitivity Analysis

A more general approach (Rosenbaum book 2002)

Define $\pi_j = e(X_j, U_j)$ for a unit j . For a given Γ , assume

$$\frac{1}{\Gamma} \leq \frac{\pi_j(1 - \pi_k)}{\pi_k(1 - \pi_j)} \leq \Gamma \text{ all pairs of units } (j, k) \text{ with } X_j = X_k$$

Then we assess how the inference on causal effect change within the set for different Γ

- Tutorial (R package [sensitivitymult](#)):

<https://rosenbap.shinyapps.io/learnsenShiny/>