Treatment Effects Practical Session #1: Testing the LATE Model

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Introduction

This practical session is based on Huber & Mellace (2015). You may find it helpful to consult the paper and or my lecture notes. See Hands-On Programming with R for a review of basic R that you will need below. My notes on this book are available here.

Exercises

 Write a function to simulate n iid draws from the model given below, with arguments n, alpha and beta. Your function should return a data frame (or tibble) with named columns D, Z, and Y.

$$\begin{split} Y &= D + \beta Z + U \\ D &= 1\{\alpha Z + \epsilon > 0\} \\ \begin{bmatrix} U \\ \epsilon \end{bmatrix} &\sim \text{Normal}(0, \Sigma), \quad \Sigma = \begin{bmatrix} 1 & 0.5 \\ 0.5 & 1 \end{bmatrix} \\ Z &\sim \text{Bernoulli}(0.5), \text{ indep. of } (U, \epsilon) \end{split}$$

- 2. Answer the following questions about the model from the preceding part.
 - (a) Is the treatment D endogenous? How can you tell?
 - (b) What is the distribution of treatment effects? What is the LATE in this model?
 - (c) What is the role of β ?
 - (d) What is the role of α ?
 - (e) Which of the LATE assumptions does the model satisfy?
- 3. Write a function called get_theta() to compute the sample analogues of θ_1 , θ_2 , θ_3 , θ_4 defined in Equation (7) of Huber & Mellace (2015). Your function should take a single input argument: a data frame (or tibble) with columns named D, Z, and Y corresponding to the model from above. It should return a vector with four named elements: theta1, theta2, theta3, and theta4.

- 4. Check your function from the preceding part by generating 100,000 observations from the model in part 1 with parameter values $\alpha = 0.6$ and $\beta = 1$. You should detect a violation of the LATE assumptions. Calculate the Wald estimand. Does it equal the LATE? Repeat for $\beta = 0$. How do you results change?
- 5. Repeat the preceding part for a variety of values of β until you find one for which the LATE assumptions are violated but you *cannot* detect a violation of the inequalities from the paper. Why is this possible?
- 6. Load the wooldridge dataset and read the documentation for the card dataset. Once you understand the contents of the dataset, carry out the following steps to construct a data frame (or tibble) called card_dat:
 - (a) Define the instrument Z as a dummy variable for living near a 4-year college in 1966. (The idea here is that living near a college reduces your costs of attending in a way that doesn't affect wages.)
 - (b) Define the outcome Y as the log of weekly earnings in 1976.
 - (c) Construct the treatment D as a dummy variable that equals one if a person has completed 16 years of education or more by 1976. This is effectively a proxy for "has a four-year degree."
- 7. Apply your function get_theta() to card_dat. Do you detect any violations of the LATE model? Re-read the documentation for card to see if you can find any potential explanation for your results. Interpret the IV estimate for card_dat in light of this.
- 8. **Bonus Question:** If you found the preceding parts too easy, here's a challenge for you! We did not consider statistical significance when looking for a violation of the LATE model in the preceding part. Use the function boot() from the R package boot, along with your function get_theta() from above to implement the "simple bootstrap with Bonferroni adjustment" described on page 402 of Huber & Mellace (2015) and apply it to card_dat. Briefly discuss your findings.

Solutions

```
Y <- D + beta * Z + U
tibble(Z, D, Y)
}
get_IV <- function(dat) {
  cov(dat$Y, dat$Z) / cov(dat$D, dat$Z)
}</pre>
```

Problem 2

- (a) The treatment D is indeed endogenous since it is correlated with the error term U, as we see from the variance-covariance matrix Σ .
- (b) The parameter β controls the strength of the *direct effect* of the instrument Z on the outcome Y. Unless $\beta = 0$, mean exclusion is violated.
- (c) The treatment effects in this model are in fact homogenous so their distribution is degenerate: $Y_1 Y_0 = 1$ for everyone, so this is the LATE as well as the ATE! This isn't a problem: the LATE model allows but does not require homogeneous treatment effects.
- (d) The parameter α determines the share of compliers, i.e. the share of people who will have D=1 if and only if Z=1. Since $D=1\{\alpha Z+\epsilon>0\}$, compliers are individuals with anyone with $-\alpha<\epsilon\leq 0$. Since $\epsilon \operatorname{Normal}(0,1)$, the share of compliers is $\operatorname{pnorm}(0)-\operatorname{pnorm}(-\alpha)=0.5-\operatorname{pnorm}(-\alpha)$.
- (e) It depends on the parameter values. The only two assumptions that may be violated are mean exclusion and the existence of compliers. As long as $\alpha \neq 0$, there are some compliers in the population. Mean exclusion is only satisfied if $\beta = 0$. Unconfounded type holds regardless of the values of α and β since ϵ is independent of Z. Monotonicity also holds regardless of the values of α and β because the first-stage take-up model shifts everyone in the same way: it is a threshold crossing model.

```
get_theta <- function(dat) {
    D <- dat$D
    Z <- dat$Z
    Y <- dat$Y

q <- mean(D[Z == 0]) / mean(D[Z == 1])
    if(q > 1) q <- 1
    if(q < 0) q <- 0

r <- mean(1 - D[Z == 1]) / mean(1 - D[Z == 0])
    if(r > 1) r <- 1
    if(r < 0) r <- 0

Y11 <- Y[(D == 1) & (Z == 1)]</pre>
```

```
set.seed(12345)
get IV <- function(dat) {</pre>
  D <- dat$D
  7 < - dat$7
 Y <- dat$Y
  cov(Z, Y) / cov(Z, D)
}
sims0 \leftarrow draw_sims(alpha = 0.6, beta = 0, n = 1e5)
get IV(sims0) # Should equal one and it does
## [1] 1.00287
get_theta(sims0) # Everything should be negative and everything is
##
       theta1
                  theta2
                              theta3
                                          theta4
## -0.6467863 -0.3120119 -0.4502310 -0.8680653
sims1 \leftarrow draw sims(alpha = 0.6, beta = 1, n = 1e5)
get_IV(sims1) # Way too high!
## [1] 5.413024
get theta(sims1) # A violation for both theta1 and theta4
##
       theta1
                   theta2
                              theta3
                                          theta4
```

```
## 0.3417739 -1.3105387 -1.4459594 0.1207921
```

Problem 5

```
sims2 <- draw_sims(alpha = 0.6, beta = 0.5, n = 1e5)
get_IV(sims2) # Way too high!

## [1] 3.230891
get_theta(sims2) # No violations

## theta1 theta2 theta3 theta4
## -0.1460979 -0.8244632 -0.9774652 -0.3703117</pre>
```

Problem 6

```
library(dplyr)
library(wooldridge)
data(card) #?card for documentation
card_dat <- card %>%
  mutate(Y = lwage, Z = nearc4, D = 1 * (educ >= 16)) %>%
  select(Y, Z, D) %>%
  tibble()

# Check some values against Huber & Mellace
# Share of compliers matches the paper
card_dat %>%
  group_by(Z) %>%
  summarize(takeup = mean(D)) %>%
  pull(takeup) %>%
  diff()
```

```
## [1] 0.06856902
get_IV(card_dat)
```

```
## [1] 2.273731
```

The LATE estimate suggests an *insanely* high return to college: remember that the outcome here is on the log scale!

```
# Check for violations of the LATE model
get_theta(card_dat) # There seems to be a violation in theta4
```

```
## theta1 theta2 theta3 theta4
## -0.09029760 -0.25126153 -0.23175056 0.09925123
```

In the documentation, we see that IQ is a crucial omitted variable and that it is correlated with the instrument. This may be partially responsible for the outrageously high LATE estimate.

```
library(boot)
get_boot_p <- function(dat) {</pre>
  boot results <- boot(dat,</pre>
                         function(data, boot_rows) get_theta(data[boot_rows,]),
                         R = 499
  theta_hat <- boot_results$t0</pre>
  theta hat boot <- boot results$t
  colnames(theta hat boot) <- names(theta hat)</pre>
  theta_tilde_boot <- sweep(theta_hat_boot, 2, theta_hat)</pre>
  colMeans(sweep(theta_tilde_boot, 2, theta_hat, FUN = ">"))
}
pvalues <- get_boot_p(card_dat)</pre>
pvalues
##
      theta1
                 theta2
                            theta3
                                       theta4
## 0.9759519 1.0000000 1.0000000 0.0000000
4 * min(pvalues) # Bonferroni correction
## [1] 0
```