

# BIOSTAT620 HW2

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## PROBLEM 1

ALL THE CODE IS IN THE github, [https://github.com/ZihaoHanGitHub/biostat620\\_hw2](https://github.com/ZihaoHanGitHub/biostat620_hw2)

1a:B

1b:A

1c:AD

1d:D

1e:AB

## PROBLEM 2

(a)

## Equation 1: SUR for Total Screen Time:

```
##
## SUR estimates for 'eq1' (equation 1)
## Model Formula: Total.ST.min ~ Y1_lag1 + X + Z
##
##           Estimate Std. Error t value Pr(>|t|)
## (Intercept) 352.0822019  93.7943895  3.75377 0.00077732 ***
## Y1_lag1      0.0759624   0.1474990  0.51500 0.61045444
## X           113.7532152   64.1235419  1.77397 0.08656981 .
## Z           11.6993908   55.0486027  0.21253 0.83318166
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 158.330817 on 29 degrees of freedom
## Number of observations: 33 Degrees of Freedom: 29
## SSR: 726990.782164 MSE: 25068.647661 Root MSE: 158.330817
## Multiple R-Squared: 0.088851 Adjusted R-Squared: -0.005406

##
## Equation 2: SUR for Social Screen Time:
```

```
##
## SUR estimates for 'eq2' (equation 2)
## Model Formula: Social.ST.min ~ Y2_lag1 + X + Z
##
##           Estimate Std. Error  t value Pr(>|t|)
## (Intercept)  98.266204   36.251233   2.71070 0.011161 *
## Y2_lag1       0.196162    0.151791   1.29232 0.206450
## X             34.623299   27.181768   1.27377 0.212860
## Z            -25.896557   23.394815  -1.10694 0.277419
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 66.588376 on 29 degrees of freedom
## Number of observations: 33 Degrees of Freedom: 29
## SSR: 128586.344076 MSE: 4434.011865 Root MSE: 66.588376
## Multiple R-Squared: 0.11593 Adjusted R-Squared: 0.024475
```

(b)

**Identify covariates in each model that are statistically significant at level  $\alpha = 0.05$ . Explain.**

From the table, in the model of Total Screen Time, we get The intercept is statistically significant at the 0.05 level (with  $pvalue = 0.00077732 < 0.05$ ), suggesting that the average value of the response variable is significantly different from zero. For other variables, Y1\_lag1 with the  $pvalue = 0.61045444 > 0.05$ , X with the  $pvalue = 0.08656981 > 0.05$ , Z with the  $pvalue = 0.83318166 > 0.05$ , indicating these three variables does not have a significant effect on Total Screen Time.

From the table, in the model of Social Screen Time, we get The intercept is statistically significant at the 0.05 level (with  $pvalue = 0.011161 < 0.05$ ), suggesting that the average value of the response variable is significantly different from zero. For other variables, Y2\_lag1 with the  $pvalue = 0.206450 > 0.05$ , X with the  $pvalue = 0.212860 > 0.05$ , Z with the  $pvalue = 0.277419 > 0.05$ , indicating these three variables does not have a significant effect on Total Screen Time.

(c)

**Test the null hypothesis  $\beta_3 = \gamma_3 = 0$ , that is,  $Z(t)$  is not an important predictor in BOTH screen time outcomes. Draw conclusion at  $\alpha = 0.05$  level.**

```
## Do not have sufficient evidence to reject the null hypothesis,
##           with the pvalue 0.8438972
```

```
## This test use the Wald test, with the wald test statistics 0.03877393
```

## PROBLEM 3

(a)

**Explain why  $X_i$  and  $\epsilon_i$  are independent.**

$X_i$  denote the  $i^{th}$  patients using A or B drugs, and  $\epsilon_i$  denoted the error term in this model. Since in this model, the dataset is collected from a randomized clinical trail, it means that the drug selection in the clinical trial is random during the treatment, each individual has the equal chance to use A or B drugs, therefore,

$X_i$  would be independent on each other factors. More than that, in SLR assumption, the independence assumption is one of the basic assumption in SLR, therefore, the error term  $\epsilon_i$  is always independent on other variables. Hence,  $X_i$  and  $\epsilon_i$  are independent.

(b)

**In model (1), explain which parameter represents the treatment effect of drug A, and explain which parameter represents the treatment effect of drug B**

Since the individual who receive drug A (coded by  $X_1 = 1$ ), and who are randomized to receive drug B (coded by  $X_i = -1$ ). The effect parameter of drug A is  $\beta_1$ , and the effect parameter of drug B is  $-\beta_1$ .

(c)

**Show that the treatment effects identified in part (b) are invariant for the inclusion of any confounding covariate Z into the model (1)**

Since we plug in the confounding covariate Z into the model, the model equation is become  $Y_i = \beta_0 + \beta_1 X_i + \beta_2 Z_i + \epsilon_i, i = 1, \dots, 2n$ . However, the effect of drug A and drug B do not change, The effect parameter of drug A is still  $\beta_1$ , and the effect parameter of drug B is still  $-\beta_1$ . Plug in a confounding covariate Z does not change the explanation of effect parameter, meaning that the plug in the Z does not change the interpretation of effects of drug A or drug B. Hence, the treatment effects are invariant for the inclusion of any confounding covariates.

(d)

**Give the estimate of the causal effect (i.e. ATE) when drug B is a placebo.**

Since drug B is a placebo, ATE is defined as the average causal effect of the treatment across all subjects. Given that

$ATE = E[Y|X_i = 1] - E[Y|X_i = -1]$ , defined as the difference in the expected values of Y between those who receive drug A and those who receive drug B (placebo). Therefore,  $ATE = \beta_1 - (-\beta_1) = 2\beta_1$ .

Hence the estimate of causal effect ATE is  $2\beta_1$  when B is a placebo.

## PROBLEM 4

(a)

**What is the variance of the error  $\tilde{\epsilon} = \beta_1 \epsilon + e$  under the assumption that the two errors  $e$  and  $\epsilon$  are independent?**

Since two errors  $e$  and  $\epsilon$  are independent,

$$Var[\tilde{\epsilon}] = Var[\beta_1 \epsilon + e] = Var[\beta_1 \epsilon] + Var[e]$$

$$\text{Therefore, } Var[\tilde{\epsilon}] = \beta_1^2 Var[\epsilon] + Var[e] = \beta_1^2 \sigma_\epsilon^2 + \sigma_e^2$$

(b)

**What is the variance of the unbiased estimator of  $\beta_1 = \alpha_1\beta_1$  denoted by  $\hat{\beta}_1$ , when a random sample of  $n$  observations  $(Z_i, X_i, Y_i), i = 1, \dots, n$ , are collected from a biomedical study?**

Since  $Y = \beta_0 + \beta_1 Z + \varepsilon$ , where  $\beta_1 = \alpha_1\beta_1$ .

The unbiased estimator of  $\beta_1$  is  $\tilde{\beta}_1 = \frac{\sum_{i=1}^n (Z_i - \bar{Z})(Y_i - \bar{Y})}{\sum_{i=1}^n (Z_i - \bar{Z})^2}$

With the Variance  $Var(\tilde{\beta}_1) = \frac{Var[\varepsilon]}{SSZ}$ , where  $SSZ = \sum_{i=1}^n (Z_i - \bar{Z})^2$

Hence  $Var(\tilde{\beta}_1) = \frac{\beta_1^2 \sigma_\varepsilon^2 + \sigma_\varepsilon^2}{SSZ}$

(c)

**What is the variance of the unbiased estimator  $\alpha_1$ , denoted by  $\hat{\alpha}_1$ , with the random sample of  $n$  observations  $(Z_i, X_i, Y_i), i = 1, \dots, n$ ,**

Since the unbiased estimator of  $\alpha_1$  is  $\alpha_1 = SSXZ/SSZ$ , therefore it given the variance

$Var(\hat{\alpha}_1) = \frac{Var[\varepsilon]}{SSZ} = \frac{\sigma_\varepsilon^2}{SSZ}$ , where  $SSZ = \sum_{i=1}^n (Z_i - \bar{Z})^2$ .

(d)

**Deriving the variance of the IV estimator  $\hat{\beta}_1 = \frac{\hat{\beta}_1}{\hat{\alpha}_1}$  seems to be analytically challenging. One may invoke the method of bootstrap to numerically evaluate this variance with the random sample of  $n$  observations  $(Z_i, X_i, Y_i), i = 1, \dots, n$ . Describe the major steps and pseudo code that you design to implement the bootstrap method in the calculation of the variance of the IV estimator.**

Since the  $\hat{\beta}_1$  is the effect of  $Z$  on  $Y$ , and  $\hat{\alpha}_1$  is the effect of  $Z$  on  $X$ , therefore, they are independent of each other.

Therefore  $Var(\hat{\beta}_1) = Var(\frac{\hat{\beta}_1}{\hat{\alpha}_1}) = \frac{Var(\hat{\beta}_1)}{Var(\hat{\alpha}_1)}$

Hence,  $Var(\hat{\beta}_1) = \frac{\beta_1^2 \sigma_\varepsilon^2 + \sigma_\varepsilon^2}{SSZ} * \frac{SSZ}{\sigma_\varepsilon^2} = \frac{\beta_1^2 \sigma_\varepsilon^2 + \sigma_\varepsilon^2}{\sigma_\varepsilon^2}$

Pseudo Code for implement the Bootstrap method for calculating the variance of the IV estimator:

*Loading the Data*

```
$ data = read.csv("."), colname = c("Y", "X", "Z")$
```

*Define the Bootstrap*

```
iteration = n for (i in 1:iteration) { # sampling from the original dataset # calculate the variance of VI estimator }
```

*Calculate the Variance of VI estimator by bootstrap*

## PROBLEM 5

We denote that  $x_{i2}$  as an indicator of gender,  $x_{i2} = 0$  for male, and  $x_{i2} = 1$  for female.

Therefore,

$$y_i = \beta_0^F + \beta_1^F z_i + \beta_2^F x_{i1} + \varepsilon_i^F$$

$$y_i = \beta_0^M + \beta_1^M z_i + \beta_2^M x_{i1} + \varepsilon_i^M$$

$y_i = \beta_0 + \beta_1 z_i + \beta_2 x_{i1} + \beta_3 x_{i2} + \beta_4(z_i \times x_{i2}) + \beta_5(x_{i1} \times x_{i2}) + \varepsilon_i$ , when  $x_{i2} = 0$  is becomes to

$$y_i = \beta_0 + \beta_1 z_i + \beta_2 x_{i1} + \varepsilon_i, \text{ therefore, } \beta_0^M = \beta_0, \beta_1^M = \beta_1, \beta_2^M = \beta_2$$

When  $x_{i2} = 1$ , the linear model is given by:

$$y_i = \beta_0 + \beta_1 z_i + \beta_2 x_{i1} + \beta_3 + \beta_4 z_i + \beta_5 x_{i1}, \text{ therefore, } \beta_0^F = \beta_0 + \beta_3, \beta_1^F = \beta_1 + \beta_4, \beta_2^F = \beta_2 + \beta_5$$