### P8130hw4

Ze Li

2023-11-25

#### Problem 1

(a)

H0: The median blood sugar readings was equal to 120 in the population from which the 25 patients were selected

Ha:The median blood sugar readings was less than 120 in the population from which the 25 patients were selected.

```
data1 = c(125,123, 117, 123, 115, 112, 128, 118, 124, 111, 116, 109, 125, 120, 113, 123, 112, 118, 121,
data1
  [1] 125 123 117 123 115 112 128 118 124 111 116 109 125 120 113 123 112 118 121
## [20] 118 122 115 105 118 131
medianvalue = 120
diff = data1-medianvalue
num_neg = sum(diff<0)</pre>
result1.1 <- binom.test(num_neg, length(diff), p = 0.5, alternative = "less")
result1.1
##
## Exact binomial test
##
## data: num_neg and length(diff)
## number of successes = 14, number of trials = 25, p-value = 0.7878
## alternative hypothesis: true probability of success is less than 0.5
## 95 percent confidence interval:
## 0.0000000 0.7301469
## sample estimates:
## probability of success
                     0.56
```

The test statistics 0.56. Since the p-value is 0.7878, which is greater than 0.05, so we fail to reject the null hypothesis, indicating we have no evidence that the median blood sugar levels are less than 120.

(b)

```
result1.2 <- wilcox.test(data1, mu = medianvalue, alternative = "less")

## Warning in wilcox.test.default(data1, mu = medianvalue, alternative = "less"):
## cannot compute exact p-value with ties

## Warning in wilcox.test.default(data1, mu = medianvalue, alternative = "less"):
## cannot compute exact p-value with zeroes

result1.2

##
## Wilcoxon signed rank test with continuity correction
##
## data: data1
## V = 112.5, p-value = 0.1447
## alternative hypothesis: true location is less than 120</pre>
```

The test statistics is 112.5. Since the p-value is 0.1447, which is greater than 0.05, so we fail to reject the null hypothesis, indicating we have no evidence that the median blood sugar levels are less than 120.

#### Problem 2

(a)

```
data2 = read_excel("Brain.xlsx") |>
  janitor::clean_names()
data2_nohomo = data2 |>
  filter(species != "Homo sapiens")
model=lm(glia_neuron_ratio ~ ln_brain_mass,data2_nohomo)
summary(model)
```

```
##
## lm(formula = glia_neuron_ratio ~ ln_brain_mass, data = data2_nohomo)
##
## Residuals:
       Min
                  1Q
                      Median
                                    3Q
                                            Max
## -0.24150 -0.12030 -0.01787 0.15940 0.25563
##
## Coefficients:
                 Estimate Std. Error t value Pr(>|t|)
##
                  0.16370
                             0.15987
                                       1.024 0.322093
## (Intercept)
                            0.03604
                                       5.026 0.000151 ***
## ln_brain_mass 0.18113
## ---
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' ' 1
## Residual standard error: 0.1699 on 15 degrees of freedom
## Multiple R-squared: 0.6274, Adjusted R-squared: 0.6025
## F-statistic: 25.26 on 1 and 15 DF, p-value: 0.0001507
```

(b)

```
predicted = model$coefficients[1]+7.22*model$coefficients[2]
predicted

## (Intercept)
## 1.471458
```

(c)

The interval for the prediction of a single new observation is more relevant for your prediction of human glia-neuron ratio than an interval for the predicted mean glia-neuron ratio at the given brain mass.

(d)

```
# Method1
prediction_interval = predict(model, newdata = data.frame(ln_brain_mass= 7.22), interval = "prediction"
prediction_interval
##
          fit
                   lwr
                             upr
## 1 1.471458 1.036047 1.906869
# Method2
tcrit = qt(df=15, 0.975)
se=sqrt(0.1699)
lowerbound = predicted - tcrit*se
upperbound = predicted + tcrit*se
c(lowerbound, upperbound)
## (Intercept) (Intercept)
     0.5928973
                 2.3500186
```

The 95% prediction interval for human glia-neuron ratio is (1.04, 1.91). So we can conclude that human brain doesn't have an excessive glia-neuron ratio for its mass compared with other primates.

(e)

Considering the position of the human data point relative to those data used to generate the regression line, we can see that the point falls beyond the range of the variable used to fit the line, so we are not certain that the regression line could be used to predict the glia\_neuron ratio of humans.

#### Problem 3

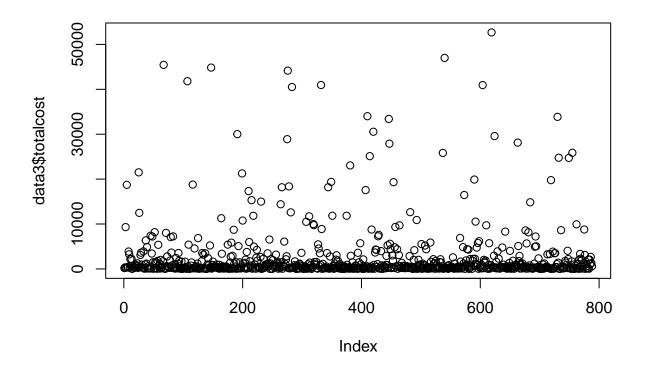
(a)

The main outcome is total cost (in dollars) of patients diagnosed with heart disease. The main predictor is number of emergency room (ER) visits. And other important covariates are age, gender, number of complications that arose during treatment, and duration of treatment condition.

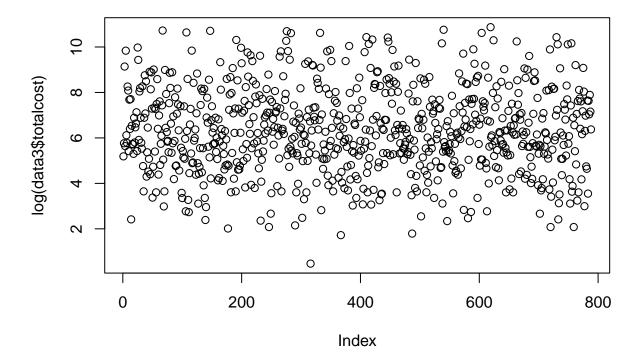
```
##
         id
                   totalcost
                                                    gender
                                       age
         : 1.0
                       : 0.0
                                 Min.
                                        :24.00
                                                 female:608
   Min.
                 Min.
  1st Qu.:197.8
                  1st Qu.: 161.1
                                                 male :180
                                  1st Qu.:55.00
## Median :394.5
                  Median : 507.2
                                  Median :60.00
         :394.5
                  Mean : 2800.0
## Mean
                                  Mean :58.72
   3rd Qu.:591.2
                  3rd Qu.: 1905.5
                                  3rd Qu.:64.00
##
## Max.
         :788.0
                  Max. :52664.9
                                  Max. :70.00
                                   e_rvisits
                                                  complications
## interventions
                      drugs
## Min. : 0.000
                   Min.
                         :0.0000
                                  Min. : 0.000
                                                  Min. :0.00000
## 1st Qu.: 1.000
                   1st Qu.:0.0000
                                  1st Qu.: 2.000
                                                  1st Qu.:0.00000
## Median : 3.000
                   Median :0.0000
                                  Median : 3.000
                                                  Median :0.00000
## Mean : 4.707
                   Mean :0.4467
                                  Mean : 3.425
                                                  Mean :0.05711
## 3rd Qu.: 6.000
                   3rd Qu.:0.0000
                                  3rd Qu.: 5.000
                                                  3rd Qu.:0.00000
## Max.
        :47.000
                   Max.
                         :9.0000
                                  Max. :20.000
                                                  Max. :3.00000
## comorbidities
                     duration
## Min. : 0.000
                  Min. : 0.00
                   1st Qu.: 41.75
## 1st Qu.: 0.000
## Median : 1.000
                   Median :165.50
## Mean : 3.767
                   Mean :164.03
## 3rd Qu.: 5.000
                   3rd Qu.:281.00
## Max. :60.000
                   Max. :372.00
```

(b)

plot(data3\$totalcost)



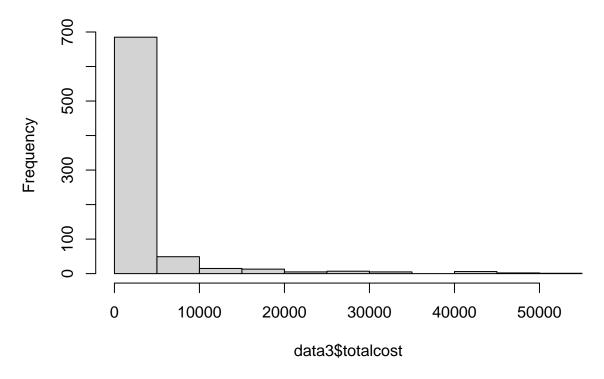
plot(log(data3\$totalcost))



It seems that after log transformation, the plot is approximately to normality since the points are randomly distributed.

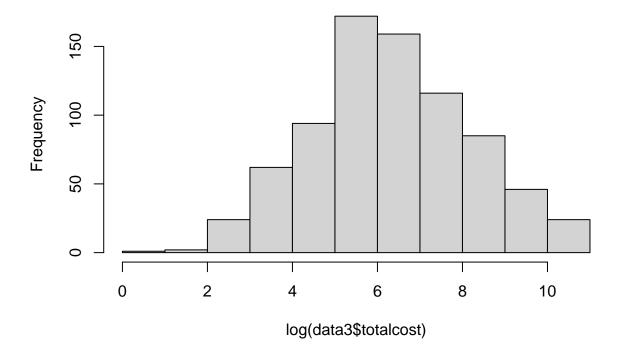
hist(data3\$totalcost)

# Histogram of data3\$totalcost



hist(log(data3\$totalcost))

## Histogram of log(data3\$totalcost)

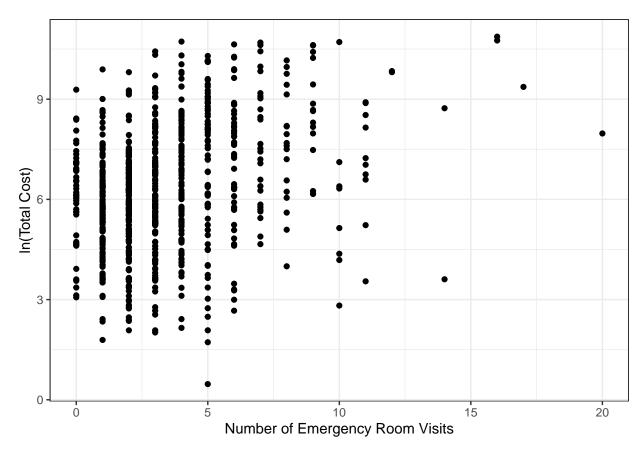


We can also see from the histogram plot that log transformation helps improve normality.

(c)

```
heart_data =
  data3 |>
  mutate(
    comp_bin =
        case_when(
        complications == 0 ~ "0",
        TRUE ~ "1"
        )) |>
  filter(totalcost > 0) |>
  mutate(ln_cost = log(totalcost))
```

(d)



```
model3 = lm(ln_cost~e_rvisits,heart_data)
summary(model3)
```

```
##
## Call:
## lm(formula = ln_cost ~ e_rvisits, data = heart_data)
##
## Residuals:
##
       Min
                1Q Median
                               3Q
                                      Max
## -6.2013 -1.1265 0.0191 1.2668 4.2797
##
## Coefficients:
##
              Estimate Std. Error t value Pr(>|t|)
## (Intercept) 5.53771
                          0.10362
                                     53.44
                                            <2e-16 ***
## e_rvisits
                0.22672
                           0.02397
                                     9.46
                                            <2e-16 ***
## ---
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' 1
## Residual standard error: 1.772 on 783 degrees of freedom
## Multiple R-squared: 0.1026, Adjusted R-squared: 0.1014
## F-statistic: 89.5 on 1 and 783 DF, p-value: < 2.2e-16
t_crit = qt(p=.05/2, df=783, lower.tail=FALSE)
t_crit
```

## [1] 1.962998

The slope is 0.22672, at a 5% significance level,  $t > t_783,0.975$ , we reject the null and conclude that there is a significant linear association between the number of Emergency room visits and  $\ln(\text{Total cost})$ .

It means that holding all other variable constant, as the risk of ERvisits goes up by 1 percent point, the predicted ln(Total cost) will increase by approximately 0.22672 dollars.

(e)

(i) Test if comp\_bin is an effect modifier of the relationship between totalcost and ERvisits. Comment.

```
fit_inter = lm(totalcost ~ e_rvisits*comp_bin, data = heart_data)
summary(fit_inter)
##
## Call:
## lm(formula = totalcost ~ e_rvisits * comp_bin, data = heart_data)
## Residuals:
             1Q Median
##
     Min
                           3Q
                                 Max
## -14973 -2187 -973
                          247
                               42326
##
## Coefficients:
##
                      Estimate Std. Error t value Pr(>|t|)
## (Intercept)
                       -566.69
                                   367.27 -1.543 0.12325
## e_rvisits
                        922.13
                                    87.07 10.590 < 2e-16 ***
## comp bin1
                       5423.48
                                  1937.91
                                            2.799 0.00526 **
## e_rvisits:comp_bin1 -277.03
                                   336.56 -0.823 0.41069
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 6148 on 781 degrees of freedom
## Multiple R-squared: 0.1614, Adjusted R-squared: 0.1582
## F-statistic: 50.1 on 3 and 781 DF, p-value: < 2.2e-16
```

The comp\_bin is not an effect modifier of the relationship between totalcost and ERvisit, since the p-value for the coefficient of e\_rvisits\*comp\_bin is not significant.

(ii) Test if comp\_bin is a confounder of the relationship between totalcost and ERvisits. Comment.

```
fit1 = lm(ln_cost ~ e_rvisits, data = heart_data)
fit2 = lm(ln_cost ~ e_rvisits + comp_bin, data = heart_data)
fit1$coefficients

## (Intercept) e_rvisits
## 5.5377096 0.2267218

fit2$coefficients

## (Intercept) e_rvisits comp_bin1
## 5.5210974 0.2046044 1.6858626
```

The coefficients of e\_rvisits in the regression model with or without comp\_bin did not show much difference, showing that comp\_bin might not be considered a confounder of the relationship between totalcost and ERvisits.

### (iii) Decide if comp\_bin should be included along with ERvisits. Why or why not?

```
fit2 |>
anova()
```

From the ANOVA test, the comp\_bin should be included with ERvisits as the p-value for the coefficient of comp\_bin is less than 0.05 in this model.

(f)

(i) Fit a MLR, show the regression results and comment.

```
fit_mlr =
  lm(ln_cost ~ e_rvisits + comp_bin + age + gender + duration, data = heart_data)
fit_mlr |>
  summary()
```

```
##
## Call:
## lm(formula = ln_cost ~ e_rvisits + comp_bin + age + gender +
       duration, data = heart_data)
##
##
## Residuals:
      Min
               1Q Median
                               30
                                      Max
## -5.0823 -1.0555 -0.1352 0.9533
                                  4.3462
##
## Coefficients:
##
                Estimate Std. Error t value Pr(>|t|)
## (Intercept) 6.0449619 0.5063454 11.938 < 2e-16 ***
## e_rvisits
                                      7.874 1.15e-14 ***
               0.1757486 0.0223189
## comp_bin1
               1.4921110
                          0.2554883
                                      5.840 7.65e-09 ***
               -0.0221376
## age
                          0.0086023
                                     -2.573
                                              0.0103 *
## gendermale -0.1176181
                          0.1379809
                                     -0.852
                                              0.3942
               0.0055406 0.0004848 11.428 < 2e-16 ***
## duration
## ---
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' ' 1
## Residual standard error: 1.605 on 779 degrees of freedom
## Multiple R-squared: 0.268, Adjusted R-squared: 0.2633
## F-statistic: 57.03 on 5 and 779 DF, p-value: < 2.2e-16
```

```
fit_mlr |>
anova()
```

```
## Analysis of Variance Table
##
## Response: ln_cost
##
             Df
                 Sum Sq Mean Sq F value
                                            Pr(>F)
## e_rvisits
              1
                 281.16 281.16 109.1541 < 2.2e-16 ***
## comp_bin
               1
                 112.84 112.84 43.8083 6.738e-11 ***
## age
               1
                    3.06
                            3.06
                                   1.1896
                                             0.2757
                    0.99
                                   0.3832
                                             0.5361
## gender
               1
                            0.99
              1 336.40
                         336.40 130.6016 < 2.2e-16 ***
## duration
## Residuals 779 2006.55
                            2.58
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' 1
```

The fitted model is  $\ln(\text{totalcost}) = 6.0449619 + 0.1757486$  ERvisits  $+ 1.4921110 \text{comp\_bin} + 0.0055406 \text{duration}$ . As the covariates age and gender didn't make any significant difference to the model under a 5% confidence level, they should not be included along with other variables.

(ii) Compare the SLR and MLR models. Which model would you use to address the investigator's objective and why?

```
anova(fit2,fit_mlr)
```

```
## Analysis of Variance Table
##
## Model 1: ln_cost ~ e_rvisits + comp_bin
## Model 2: ln_cost ~ e_rvisits + comp_bin + age + gender + duration
##
    Res.Df
              RSS Df Sum of Sq
                                    F
                                         Pr(>F)
## 1
       782 2347.0
## 2
        779 2006.5 3
                        340.46 44.058 < 2.2e-16 ***
## ---
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' 1
```

Since the p-value of anova test is less than 0.05, we reject the null hypotheses and conclude that the larger model is superior. As a result, we will choose MLR models.