BIOSTAT 200B HW3

- 1. Questions for Lab3
- (1) The intercept from such a regression of residuals on residuals will always be zero (disregarding rounding error). Why? (Hint: what is the sample mean of the residuals from a linear regression model? And what is the formula for the intercept in a simple linear regression model?)

The intercept in a simple regression model is $\widehat{\beta_0} = \overline{Y} - \widehat{\beta_1} \overline{X}$. Since we know the sample mean of the residuals from a linear regression model will always be 0, we can get $\overline{X} = \overline{Y} = 0$ in this regression model, which regresses residuals on residuals. Therefore, $\widehat{\beta_0}$ will always be zero in this case.

(2) How do we interpret the parameter estimates for the coefficients for regnc, regs, regw? How do we interpret the intercept in this model?

The parameter estimates for the coefficients for **regnc** is -0.46696, which means when the hospital is located in North Central, then the infection risk would decrease by 0.46696. Similarly, the parameter estimates for the coefficients for **regs** and **regw** are -0.93369 and -0.47946, meaning that the infection risk would decrease by 0.93369 and 0.47946 when the hospital is located in South and West, respectively.

| Analysis of Variance | | | | | | | | | | | |
|------------------------|----------------|-------------------|-----|----------------|--------|-------|--------|--|--|--|--|
| Source | DF | Sum of Squares | | Mean Square | F | Value | Pr > F | | | | |
| Model | 3 | 13.99694 | . 4 | 4.66565 | 2.71 0 | | 0.0484 | | | | |
| Error | 109 | 187.38288 | | 1.71911 | | | | | | | |
| Corrected Total | 112 | 201.37982 | | | | | | | | | |
| | | | | | | | | | | | |
| Root MSE | Root MSE | | | R-Square | | 0.069 | 5 | | | | |
| Depender | Dependent Mean | | | Adj R-S | q | 0.043 | 9 | | | | |
| Coeff Var | | 30.1076 | 5 | | | | | | | | |

| Parameter Estimates | | | | | | | | | | | |
|---------------------|-----------------------|--|--|--|--|--|--|--|--|--|--|
| DF | Parameter Estimate | Standard Error | t Value | Pr > t | | | | | | | |
| 1 | 4.86071 | 0.24778 | 19.62 | <.0001 | | | | | | | |
| 1 | -0.46696 | 0.33929 | -1.38 | 0.1716 | | | | | | | |
| 1 | -0.93369 | 0.32842 | -2.84 | 0.0053 | | | | | | | |
| 1 | -0.47946 | 0.41090 | -1.17 | 0.2458 | | | | | | | |
| | 1 1 | Parameter Estimate 1 4.86071 1 -0.46696 1 -0.93369 | DF Parameter Estimate Standard Error 1 4.86071 0.24778 1 -0.46696 0.33929 1 -0.93369 0.32842 | DF Parameter Estimate Standard Error t Value 1 4.86071 0.24778 19.62 1 -0.46696 0.33929 -1.38 1 -0.93369 0.32842 -2.84 | | | | | | | |

The intercept in this model is 4.86071, and we can interpret it as the estimated infection risk when the hospital is located in North East since it is our reference category for these dummy variables.

(3) What region did we make the reference region by using the above code? Write code to fit a model using a different region as the reference group and run a regression model. Compare the output, especially the parameter estimates. Did R^2 change? Did the ANOVA table change?

As we stated in question (2), North East is the region where we made the reference region by using the code. We write another code, which lets North Central be the reference

region.

```
data senic1; set senic;

if region=1 then regne=1; else regne=0;

if region=3 then regs=1; else regs=0;

if region=4 then regw=1; else regw=0;

run;

proc reg data=senic1;

model risk = regne regs regw;

run; quit;
```

The results of ANOVA table and the parameter estimates are shown beside.

We can find that the parameter estimates for the coefficients for **regs**, **regw**, and intercept all

| Analysis of Variance | | | | | | | | | | | | | | |
|----------------------|------------------|---------|------|---------------|---------|----------------|------|----------------|----|---------|------|------|--------|----|
| Sc | Source | | | DF | | Sum Squa | | Mean Square | | F Value | | ıe | Pr> | F |
| M | Model | | | 3 | | 13.996 | 394 | 4.6656 | 5 | | 2.7 | 71 | 0.04 | 84 |
| E | Error | | | 109 | 1 | 87.382 | 288 | 1.7191 | 1 | | | | | |
| C | orre | cted To | tal | 112 | 201.379 | | 982 | | | | | | | |
| | | Root N | /ISE | | | 1.3 | 1115 | R-Sq | ua | re | 0.0 |)695 | 5 | |
| | | Depen | den | t Mea | n | 4.38 | 5487 | Adj R-Sq | | q | 0.0 |)439 | 9 | |
| | Coeff Var | | | | 30.10 | | | | | | | | | |
| Parameter Estimates | | | | | | | | | | | | | | |
| | Variable DF Esti | | • | neter mate | Sta | ndard Error | ť | Val | ue | Pr | > t | | | |
| | Int | ercept | 1 | 4 | 4.39375 | | 0. | 0.23178 | | 18.96 | | <.(| <.0001 | |
| | re | gne | 1 | (| 0.4 | 6696 | 0. | 33929 | | 1. | 38 | 0.1 | 1716 | |
| | re | gs | 1 | -(| 0.4 | 6672 | 0. | 31652 | | -1. | 47 | 0.1 | 1432 | |
| | re | gw | 1 | -(| 0.0 | 1250 | 0. | 40146 | | -0. | 03 | 0.9 | 9752 | |

change. This is because we have changed our reference region form North East to North Central. Actually, the interpretation of these parameter estimates for the coefficients remain the same. In question (2), we knew that the infection risk would decrease by 0.46696 when the hospital is located in North Central, which means the estimated infection rate is 4.86071-0.46696=4.39375. In this question, we can also get the same coefficient for the intercept, as we can interpret it as the estimated infection risk when the hospital is located in North Central since it is our reference category in this case. While we comparing the R² and the ANOVA table, we can see that they are the same, because changing the reference category for dummy variables does not change the meaning of conducting these regression models.

(4) How do we interpret the p-values in the Parameter Estimates table, in terms of testing for differences in means by region? What means are being compared?

For the p-values in the Parameter Estimates table, if they are greater than the significance level α (let α be 0.05 in this question), than we cannot reject the null hypothesis, meaning that there is no statistically significant difference between the parameter estimates for the coefficients and 0. That is to say, we are comparing the mean infection risk of each

region and the reference region. For example, for the p-value of variable **regne** shown in question 3, p-value = 0.1716 > 0.05, and we can conclude that there is no statistically significant difference between the mean infection risk of North East and our reference category North Central.

(5) Write out the null and alternative hypotheses for the test conducted by "test_region". Give the distribution of the test statistic under the null, the value of the test statistic and the p-value. What do you conclude?

The null hypothesis H_0 is the parameter estimates for the coefficient of variable **regnc**, **regs**, and **regw** are all equal to 0. The alternative hypothesis H_1 is as least one of the parameter estimates for the coefficient of variable **regnc**, **regs**, and **regw** is not equal to 0.

| Test test_region Results for Dependent Variable risk | | | | | | | | | | |
|--|-----|----------------|---------|--------|--|--|--|--|--|--|
| Source | DF | Mean Square | F Value | Pr > F | | | | | | |
| Numerator | 3 | 3.04987 | 2.50 | 0.0636 | | | | | | |
| Denominator | 107 | 1.22083 | | | | | | | | |

The distribution of the test statistics is $F^*\sim F_{3,107}$, and the value of the test statistic F^* is 2.50, with the p-value = 0.0636. Since the p-value = 0.0636 > 0.5, we cannot reject H_0 and can conclude that there is no significant difference between each of the parameter estimates for the coefficient of variable **regnc**, **regs**, **regw** and 0.

(6) Conduct the overall (omnibus) F test for the model risk = length census regnc regs regw. Write out the null and alternative hypotheses. Give the distribution of the test statistic under the null, the value of the test statistic and the p-value. What do you conclude?

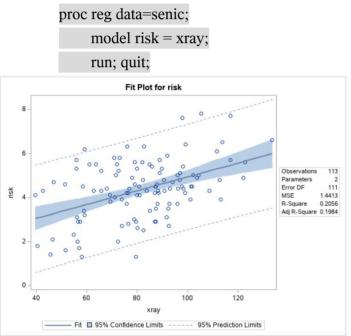
The null hypothesis H_0 is the parameter estimates for the coefficient of variable **length**, **census**, **regnc**, **regs**, and **regw** are all equal to 0. The alternative hypothesis H_1 is as least one of the parameter estimates for the coefficient of variable **length**, **census**, **regnc**, **regs**, and **regw** is not equal to 0.

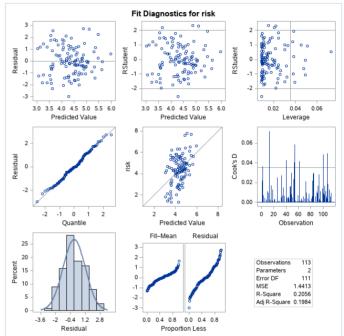
| Analysis of Variance | | | | | | | | | | |
|----------------------|-----|-------------------|----------------|---------|--------|--|--|--|--|--|
| Source | DF | Sum of Squares | Mean Square | F Value | Pr > F | | | | | |
| Model | 5 | 70.75096 | 14.15019 | 11.59 | <.0001 | | | | | |
| Error | 107 | 130.62886 | 1.22083 | | | | | | | |
| Corrected Total | 112 | 201.37982 | | | | | | | | |

The distribution of the test statistics is $F^* \sim F_{5.107}$, and the value of the test statistic F^* is

11.59, with the p-value < 0.0001. Since the p-value < 0.5, we can reject H_0 and can conclude that there is significant difference between at least one of the parameter estimates for the coefficient of variable **length**, **census**, **regnc**, **regs**, **regw** and 0.

(7) A regression of risk on xray shows a highly significant relationship. Fit the model and conduct model diagnostics (residuals analysis). Report this relationship. Provide an interpretation of the regression coefficients, using appropriate units.





The model fitting plot and residuals analysis are shown above. It looks like **xray** has a positive correlation with **risk**. Form residual analysis, we can find the regression function linear, and the error variances are constant; it also seems to meet the assumption of normality of residual according to Q-Q plot. We can interpret the regression coefficient of **xray** as the infection risk would increase by 0.0314 when the ratio of number of x-rays performed to number of patients without signs or symptoms of pneumonia X 100 increase 1 unit. The regression

| | Analysis of Variance | | | | | | | | | | | | |
|---------------------|----------------------|--------|-----|------------------|---------------------|-------|--------|-------------|--------------|---------|--------|--------|------|
| So | Source | | DF | Sum o Squares | | | | Mea Squa | | F Value | | Pr > F | |
| Mo | Model 1 | | 4 | 1.396 | 42 | 4 | 1.3964 | 12 | 28. | 72 | <.0001 | | |
| En | rror 111 1 | | 15 | 9.983 | 40 | 1 | 1.4412 | 29 | | | | | |
| Со | Corrected Total 112 | | | | 20 | 1.379 | 82 | | | | | | |
| | | | | | | | | | | | | | |
| | | Root N | ISE | | | 1.20 | 054 | 4 R-Square | | 0.2 | 0.2056 | | |
| | Depender | | | Mea | an | 4.35 | 5487 | 7 Adj R-9 | | ≀-Sq | q 0.19 | | |
| | Coeff Var | | | | | 27.56 | 3773 | 3 | | | | | |
| | | | | | | | | | | | | | |
| Parameter Estimates | | | | | | | | | | | | | |
| | Variable D | | DF | | rameter Estimate | | St | | dard rror | t Value | | Pr | > t |
| | Int | ercept | 1 | | 1.7 | 9202 | 0 | .49 | 9136 | | 3.65 | 0.0 | 004 |
| | xra | ay | 1 | | 0.0 | 3140 | 0 | 0.0 | 0586 | | 5.36 | <.0 | 001 |

coefficient of intercept is 1.79202, which means that the estimated infection risk is 1.79202 when the ratio of number of x-rays is 0.

Investigators hypothesize that xray will be significantly related to risk after controlling for beds, nurses and svcs. Run the appropriate model and report the results. Provide an interpretation of each of the regression coefficients, using appropriate units. Also report the partial correlation of risk and xrays controlling for beds, nurses and svcs. Do the

proc reg data=senic; model risk = xray beds nurses svcs; run; quit;

findings support their hypothesis or not?

The parameter estimates for the coefficients are

| Parameter Estimates | | | | | | | | | |
|---------------------|----|-----------------------|-------------------|---------|---------|--|--|--|--|
| Variable | DF | Parameter Estimate | Standard Error | t Value | Pr > t | | | | |
| Intercept | 1 | 0.86917 | 0.53682 | 1.62 | 0.1083 | | | | |
| xray | 1 | 0.02858 | 0.00542 | 5.27 | <.0001 | | | | |
| beds | 1 | -0.00033930 | 0.00141 | -0.24 | 0.8106 | | | | |
| nurses | 1 | 0.00223 | 0.00191 | 1.17 | 0.2457 | | | | |
| svcs | 1 | 0.01976 | 0.01162 | 1.70 | 0.0918 | | | | |

shown beside in the table. With all other variables holding constant, we can interpret the regression coefficient of **xray** as the infection risk would increase by 0.02858 when the ratio of number of x-rays performed to number of patients without signs or symptoms of pneumonia X 100 increase 1 unit. Similarly, the interpretation of regression coefficient of **beds** is the infection risk would decrease by 0.00003393 when the average number of beds in hospital increase 1 unit; that of **nurses** is the infection risk would increase by 0.00223 when the average number of full-time equivalent nurses increase 1 unit; that of **svcs** is the infection risk would increase by 0.001976 when the percent of 35 potential facilities and services that are provided by the hospital increase 1 unit. The regression coefficient of intercept is 0.86917, which means that the estimated infection risk is 0.86917 when all the other variables are set as 0.

proc corr data = senic; var risk; with xray; partial beds nurses svcs; run;

The result of the partial correlation of **risk** and **xrays** controlling for **beds**, **nurses** and **svcs** is shown beside. We can find that Pearson Partial Correlation Coefficients of **risk** and **xrays** is 0.4526, and the p-

| Variable N Mean Std Dev Sum Minimum Maximum Variance Std Dev beds 113 252.16814 192.84269 28495 29.00000 835.00000 nurses 113 173.24779 139.26539 19577 14.00000 656.00000 svcs 113 43.15929 15.20086 4877 5.70000 80.00000 | 1 With Variables: xray 1 Variables: risk 1 Variables: xray 1 Variables: risk 1 Variables xray 1 Variables: xray | | |
|--|---|--------------|----------|
| 1 Variables: risk | 1 Variables: risk | | |
| Simple Statistics Simple Statistics | Variable N Mean Std Dev Sum Minimum Maximu beds 113 252.16814 192.84269 28495 29.00000 835.000 nurses 113 173.24779 139.26539 19577 14.00000 656.000 svcs 113 43.15929 15.20086 4877 5.70000 80.000 | | |
| Variable N Mean Std Dev Sum Minimum Maximum Partial Variance Partial Std Dev beds 113 252.16814 192.84269 28495 29.00000 835.00000 nurses 113 173.24779 139.26539 19577 14.00000 656.00000 svcs 113 43.15929 15.20086 4877 5.70000 80.00000 | Variable N Mean Std Dev Sum Minimum Maximum beds 113 252.16814 192.84269 28495 29.00000 835.000 nurses 113 173.24779 139.26539 19577 14.00000 656.000 svcs 113 43.15929 15.20086 4877 5.70000 80.000 | | |
| Variable N Mean Std Dev Sum Minimum Maximum Variance Std Dev beds 113 252.16814 192.84269 28495 29.00000 835.00000 835.00000 835.00000 805.000000 805.000000 | beds 113 252.16814 192.84269 28495 29.00000 835.000 nurses 113 173.24779 139.26539 19577 14.00000 656.000 svcs 113 43.15929 15.20086 4877 5.70000 80.000 | | |
| nurses 113 173.24779 139.26539 19577 14.00000 656.00000 svcs 113 43.15929 15.20086 4877 5.70000 80.00000 | nurses 113 173.24779 139.26539 19577 14.00000 656.000 svcs 113 43.15929 15.20086 4877 5.70000 80.000 | | Std Dev |
| svcs 113 43.15929 15.20086 4877 5.70000 80.00000 | svcs 113 43.15929 15.20086 4877 5.70000 80.000 | 00 | |
| | | 00 | |
| Yray 113 81 62832 19 36383 9224 39 60000 133 50000 377 06899 19 4183 | vrav 113 81 62832 10 36383 0224 30 60000 133 500 | 00 | |
| 110 01.02002 10.00000 100.00000 011.00000 10.4102 | 110 01.02002 10.00000 9224 09.00000 100.000 | 00 377.06899 | 19.41826 |
| risk 113 4.35487 1.34091 492.10000 1.30000 7.80000 1.50313 1.2260 | risk 113 4.35487 1.34091 492.10000 1.30000 7.800 | 00 1.50313 | 1.22602 |
| | Prob > r under H0: Partial Rho=0 | | |
| Pearson Partial Correlation Coefficients, N = 113 Prob > r under H0: Partial Rho=0 | ris | k | |
| | xray 0.4526 < .000 | | |

value < 0.001, meaning that these two variables are significantly related after controlling for **beds**, **nurses** and **svcs**. Therefore, this finding do support investigators' hypothesis.

Conduct a joint test of whether beds and nurses contribute to explaining variation in risk after controlling for svcs and xray.

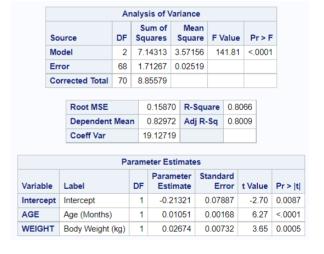
proc reg data=senic;
 model risk = xray beds nurses svcs;
 join_test: test beds, nurses;
run; quit;

| Test join_test Results for Dependent Variable risk | | | | | | | | | | |
|--|-----|----------------|---------|--------|--|--|--|--|--|--|
| Source | DF | Mean Square | F Value | Pr > F | | | | | | |
| Numerator | 2 | 1.50050 | 1.24 | 0.2924 | | | | | | |
| Denominator | 108 | 1.20629 | | | | | | | | |

The result of the joint test is shown above. In this test, our null hypothesis is the parameter estimates for the coefficient of variable **beds** and **nurses** are both equal to 0. From the table, we can find that the p-value is 0.2924, which is greater than our significant level 0.05, meaning that we cannot reject the null hypothesis and can conclude that there is no significant difference between each of the parameter estimates for the coefficient of variable **beds**, **nurses**, and 0.

2.(a) proc reg data=d.spirometry; model fev1 = age weight; run; quit;

The ANOVA table and the parameter estimates of the fitting model is shown beside. Form the table, we can obtain the value of R^2 is 0.8066.



(b)
proc reg data=d.spirometry;
model fev1 = age weight;

output out=out_2 predicted=pre_value;

quit;
proc corr data = out_2;
var fev1;
with pre_value;
run;

run;



As the table we shown below, we obtain the predicted values for the model in question 2(a) and the Pearson correlation between the observed FEV1 values and the predicted values. The correlation between observed FEV1 values and the predicted one is 0.89811, and the square of it is $0.89811^2 = 0.8066$, which equals to the value of R^2 we obtained in previous question.