

MA-4710 Regression Analysis Final Project



Michigan Tech

Final Project -Multilinear Regression Analysis

BY GROUP 18:

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ABSTRACT

This study delves into the comprehensive analysis of hospitals participating in the Study on the Efficacy of Nosocomial Infection Control (SENIC) project, leveraging a dataset comprising 113 samples and 11 variables. The dataset encapsulates crucial aspects of hospital characteristics, from the average length of stay and patient age to infection risk and the presence of medical schools. Employing statistical methodologies, this investigation aims to unravel patterns, relationships, and insights within the dataset, shedding light on the factors contributing to the efficacy of infection control measures in healthcare institutions.

INTRODUCTION

In the context of healthcare, effective infection control is paramount to ensuring the well-being of patients and the overall quality of medical services. This study delves into the intricacies of hospital operations by analyzing the Study on the Efficacy of Nosocomial Infection Control (SENIC) dataset, a compilation of 113 samples featuring 11 variables. These variables encompass crucial aspects of hospital characteristics, ranging from the average length of patient stays and age demographics to infection risk estimations and institutional practices such as routine culturing and X-ray ratios. Hospitals' structural components are also explored, including bed capacity, association with medical schools, and geographic regions. By employing rigorous statistical methodologies, this analysis aims to unravel patterns and relationships within the dataset, shedding light on the nuanced factors that contribute to the efficacy of infection control measures in healthcare institutions. The findings are poised to inform and enhance strategies for mitigating hospital-acquired infections and improving the overall resilience of healthcare systems.

THE GOAL OF THE PROJECT:

The goal of the analysis is to determine which predictor variables in this dataset can help to better understand and predict the length of stay of the patient in the US hospital by building the multiple linear regression model.

Description of the variables in the dataset (in ascending order):

- Length of Stay (Y): The average length of stay of all patients in the hospital (in days).
- Age (X1): The average age of patients (in years).
- Infection Risk (X2): The average estimated probability of acquiring infection in a hospital (in percent).
- Routine Culturing Ratio (X3): The ratio of the number of cultures performed to the number of patients without signs or symptoms of hospital-acquired infection, times 100.
- Routine Chest X-ray Ratio (X4): The ratio of the number of X-rays performed to the number of patients without signs or symptoms of pneumonia, times 100.
- Number of Beds (X5): The average number of beds in the hospital during the study period.
- Medical School (X6): Indicator of whether the hospital is associated with a medical school (1 = Yes, 2 = No).
- Region (X7): Indicator of the geographic region for the hospital (1 = NE, 2 = NC, 3 = S, 4 = W).
- Average daily Census (X8): The average number of patients per day in the hospital during the study period.
- Number of nurses (X9): The average number of full-time equivalent registered and licensed practical nurses during the study period (number of full-time plus one-half the number of part-time).
- Available facilities and services (X10): A percent of 35 potential facilities and services are provided by the hospital.

Numerical Variable: "X1" "X2" "X3" "X4" "X5" "X8" "X9" "X10"

Categorical Variable: "X6", "X7"

Response Variable: "Y"

EXPLORATORY DATA ANALYSIS:

Histograms of Y and Xs:

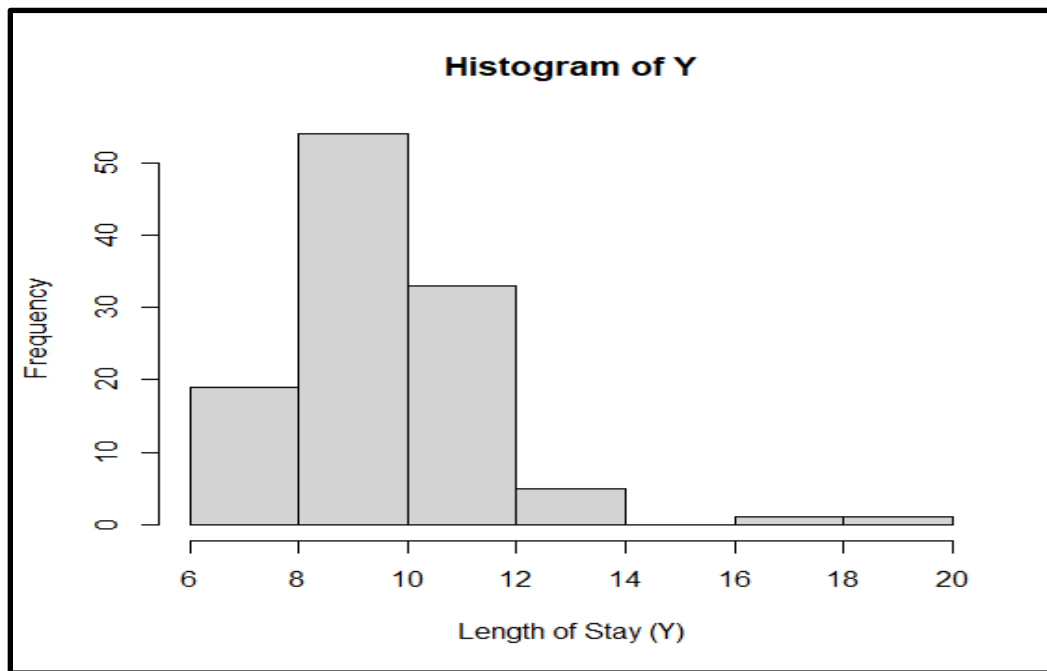
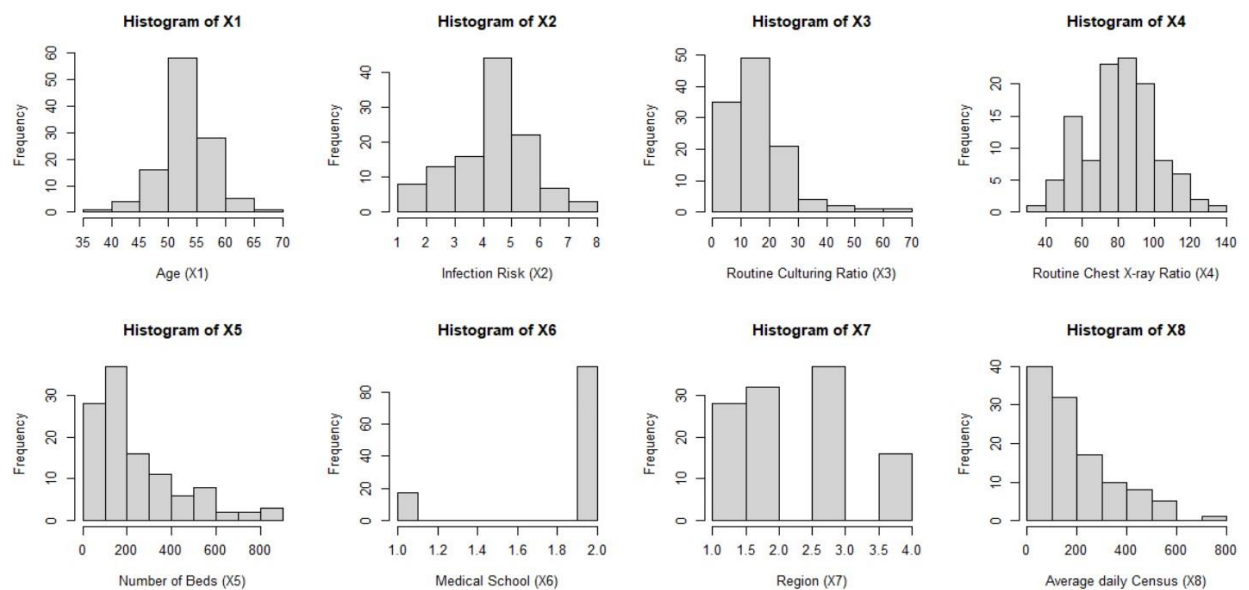


Figure 1. Histogram of the response variable; Length of Stay (Y).

The histogram for our response variable (i.e., Length of Stay), which is displayed in above figure 1 has the right-skewed distribution.



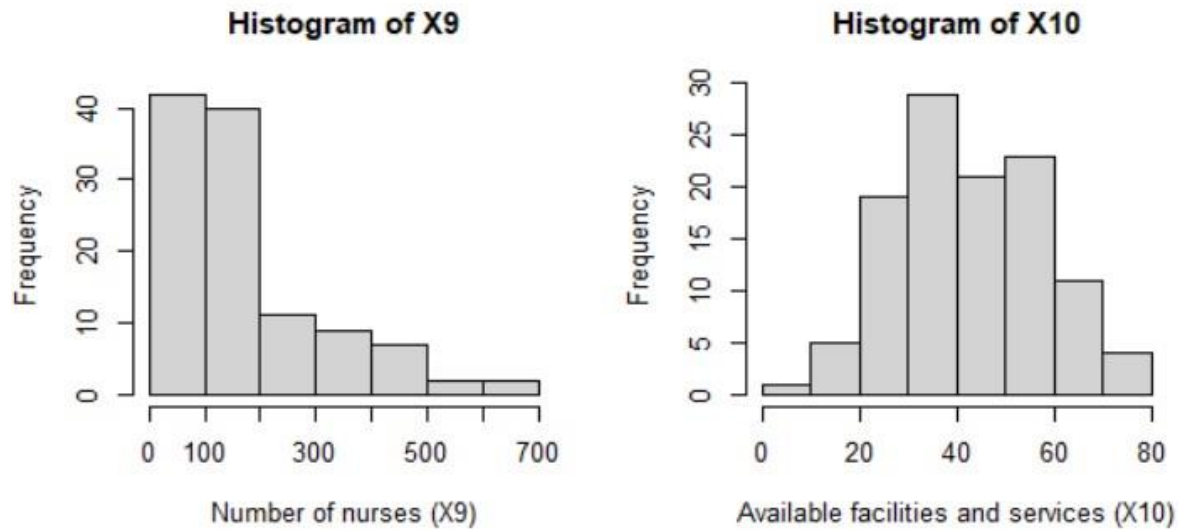


Figure 2. Histograms of the predictor variables (Xs).

```
> skew_summary
```

X1	X2	X3	X4	X5	X6
-0.101237983	-0.116597463	1.567681306	0.007669835	1.342231775	-1.929640399
X7	X8	X9	X10		
0.063487521	1.342984491	1.342382454	0.072223045		

Figure 3. Skewness of the predictor variables (Xs).

The histogram displayed in Figures 2 & 3 reveals that our predictor variable, Routine Culturing Ratio (X3) **with skewness of 1.567681306**, Number of Beds (X5) **with skewness of 1.342231775**, Average Daily Census (X8) **with skewness 1.342984491**, and Number of Nurses (X9) **with skewness 1.342382454** have a **right-skewed distribution**. On the other hand, Age (X1) **with skewness -0.101237983**, Infection Risk (X2) **with skewness -0.116597463**, Routine Chest X-ray Ratio (X4) **with skewness 0.007669835**, and Available Facilities and Services (X10) **with skewness 0.072223045**, have a **normal distribution**. In addition to these variables, we have two categorical variables, Medical School (X6) and Region (X7).

Boxplots of Y and Xs:

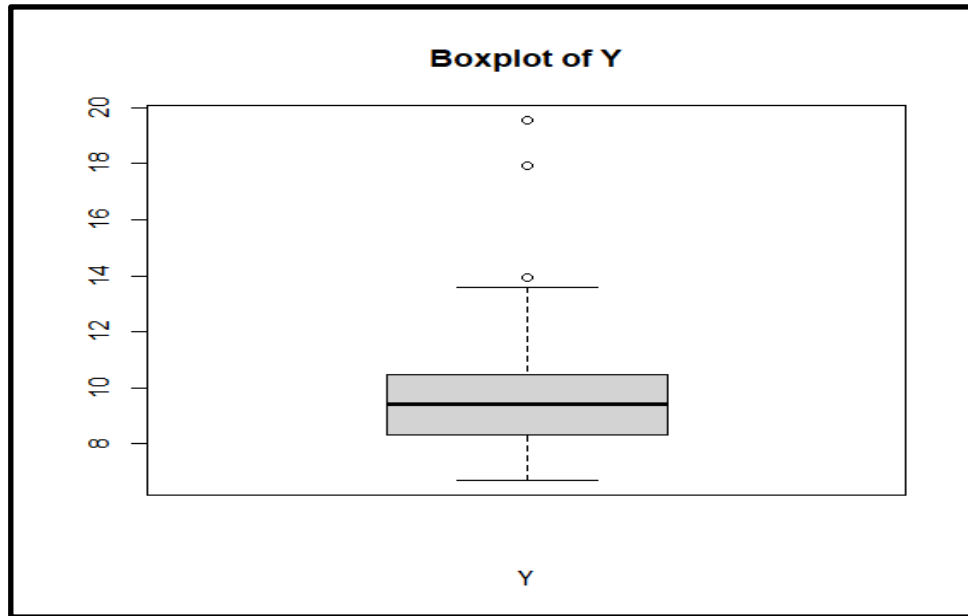


Figure 4. Boxplot of the response variable; Length of Stay (Y).

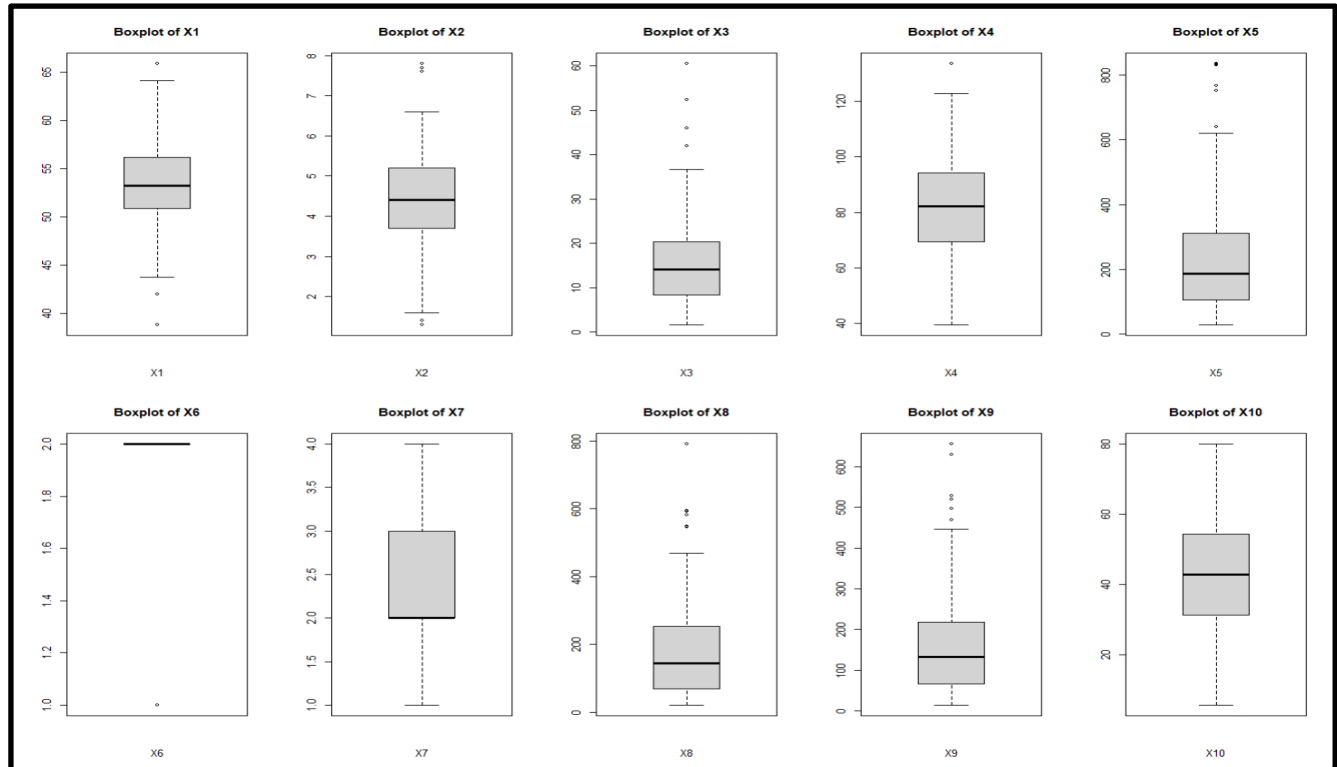


Figure 5. Boxplots of the predictor variables (Xs).

According to Figures 4 & 5, the boxplot of the response variable is not centered since one side of the median is larger than the other, and it contains a few outliers. This suggests that the response variable is not normally distributed.

Figure 4 shows that the boxplots of Routine Culturing Ratio (X3), Number of Beds (X5), Average daily Census (X8), and Number of Nurses (X9) are right skewed with multiple outliers. This indicates that these predictor variables are also not normally distributed. On the other hand, the boxplot for Age (X1), Infection Risk (X2), Routine Chest X-ray Ratio (X4), and Available facilities and services (X10) are centered around the median. Therefore, we can say that they are normally distributed.

We also have two categorical variables, Medical School (X6) and Region (X7). Since they cannot be plotted on a box plot, we cannot make any assumptions about their distribution.

Summary Statistics:-

Y	X1	X2	X3	X4	X5	X6
Min. : 6.700	Min. :38.80	Min. :1.300	Min. : 1.60	Min. : 39.60	Min. : 29.0	Min. :1.00
1st Qu.: 8.340	1st Qu.:50.90	1st Qu.:3.700	1st Qu.: 8.40	1st Qu.: 69.50	1st Qu.:106.0	1st Qu.:2.00
Median : 9.420	Median :53.20	Median :4.400	Median :14.10	Median : 82.30	Median :186.0	Median :2.00
Mean : 9.648	Mean :53.23	Mean :4.355	Mean :15.79	Mean : 81.63	Mean :252.2	Mean :1.85
3rd Qu.:10.470	3rd Qu.:56.20	3rd Qu.:5.200	3rd Qu.:20.30	3rd Qu.: 94.10	3rd Qu.:312.0	3rd Qu.:2.00
Max. :19.560	Max. :65.90	Max. :7.800	Max. :60.50	Max. :133.50	Max. :835.0	Max. :2.00
X7	X8	X9	X10			
Min. :1.000	Min. : 20.0	Min. : 14.0	Min. : 5.70			
1st Qu.:2.000	1st Qu.: 68.0	1st Qu.: 66.0	1st Qu.:31.40			
Median :2.000	Median :143.0	Median :132.0	Median :42.90			
Mean :2.363	Mean :191.4	Mean :173.2	Mean :43.16			
3rd Qu.:3.000	3rd Qu.:252.0	3rd Qu.:218.0	3rd Qu.:54.30			
Max. :4.000	Max. :791.0	Max. :656.0	Max. :80.00			

Figure 6. Summary Statistics of the data

It appears that the predictor variable Age (X1) does not provide much information about children or elderly people, as the minimum and maximum ages are between 38 and 66. Additionally, the Average Census (X8) has a wide range of values, from 20.0 to 791.0. The difference between the median and mean suggests that there may be outliers in X8. Furthermore, the Number of Beds (X5), Number of Nurses (X9), and Available Facilities and Services (X10) also have a wide range of values, indicating the potential presence of outliers.

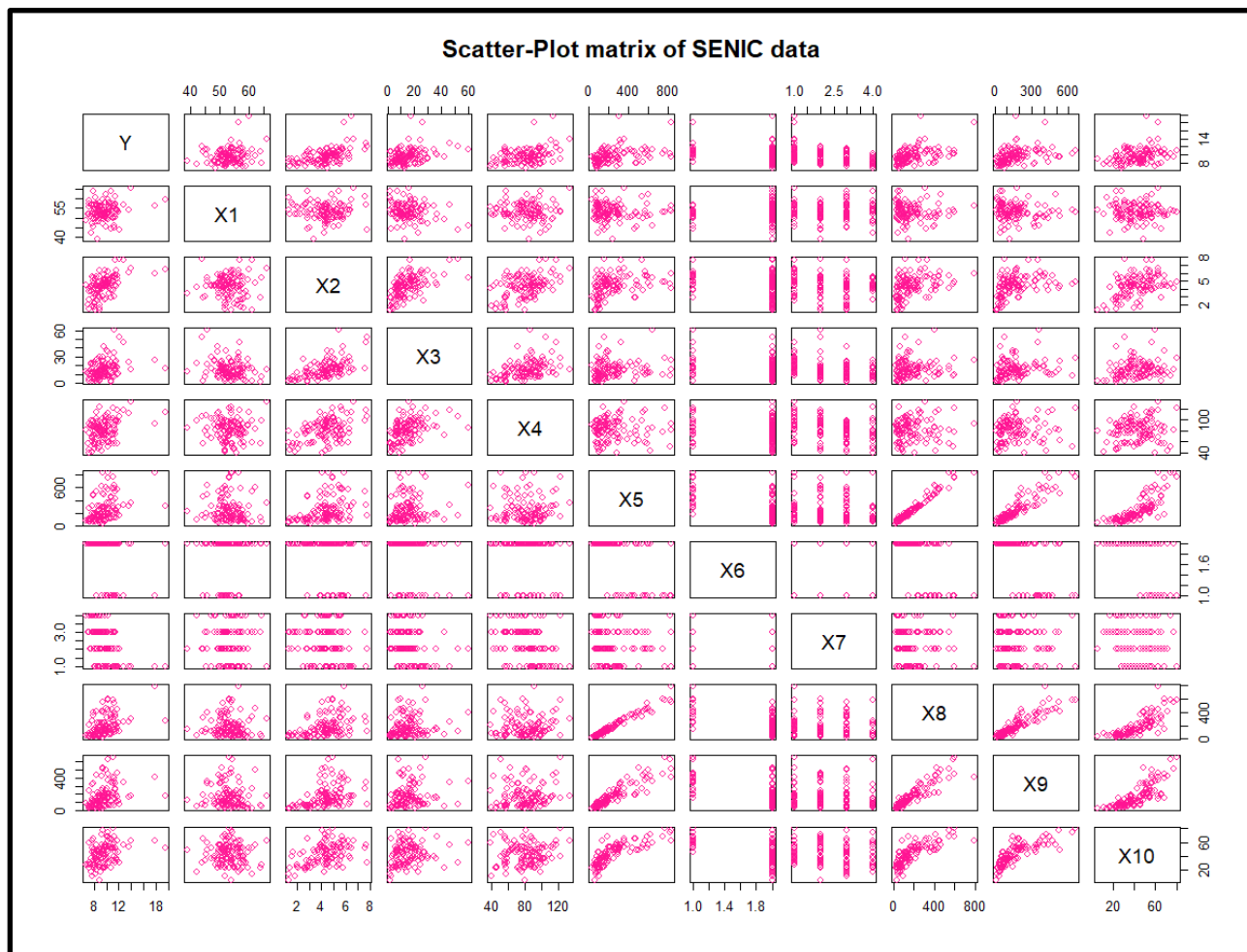


Figure 7. Scatter Plot Matrix

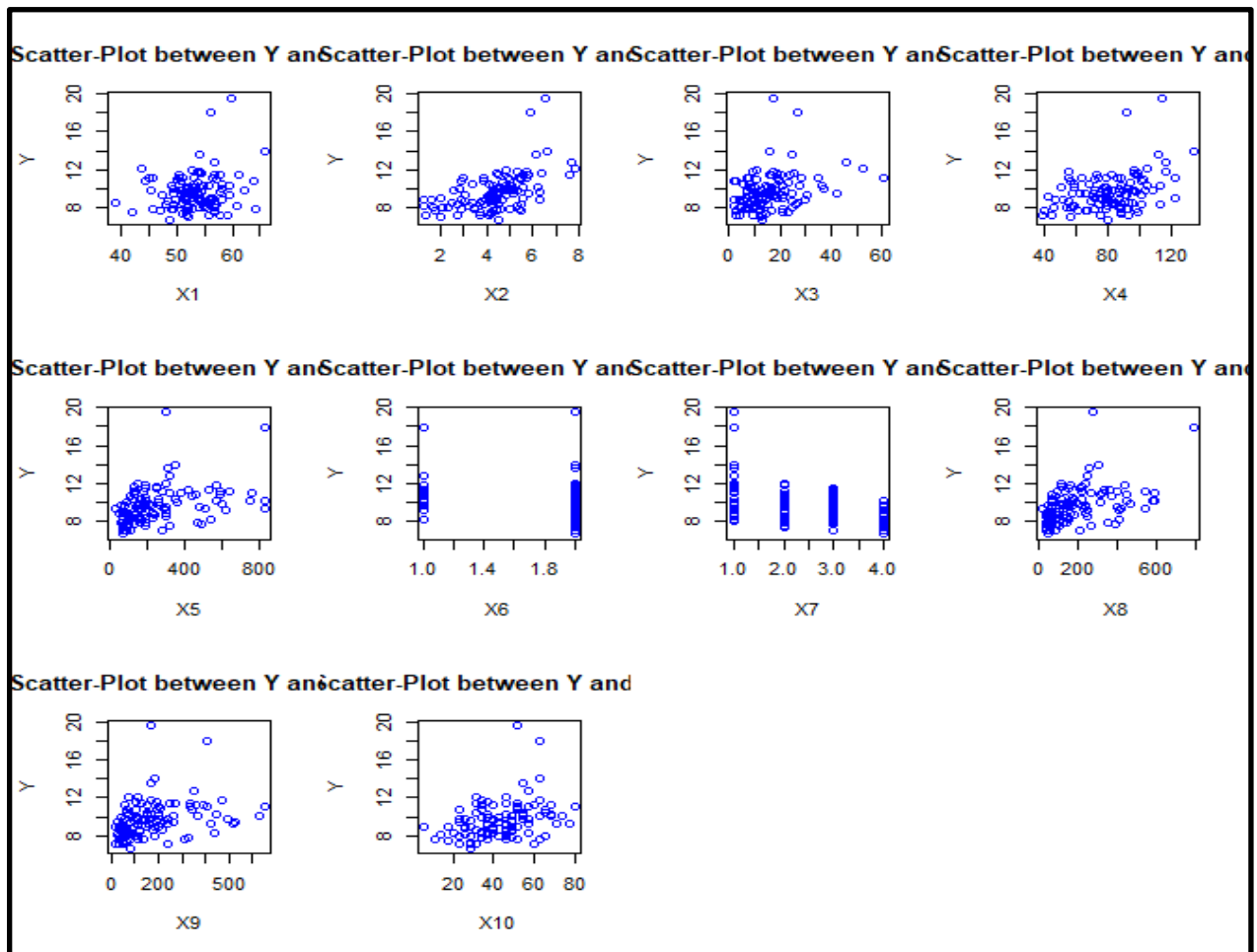


Figure 8. Scatter Plot Matrix of the response variable against the predictor variables.

We can conclude from the Scatter plot (Figures 7&8) that all predictor variables have a kind of linear relationship with the response variable, except for the two categorical variables, Medical School (X6) and Region (X7).

Added variable plots:

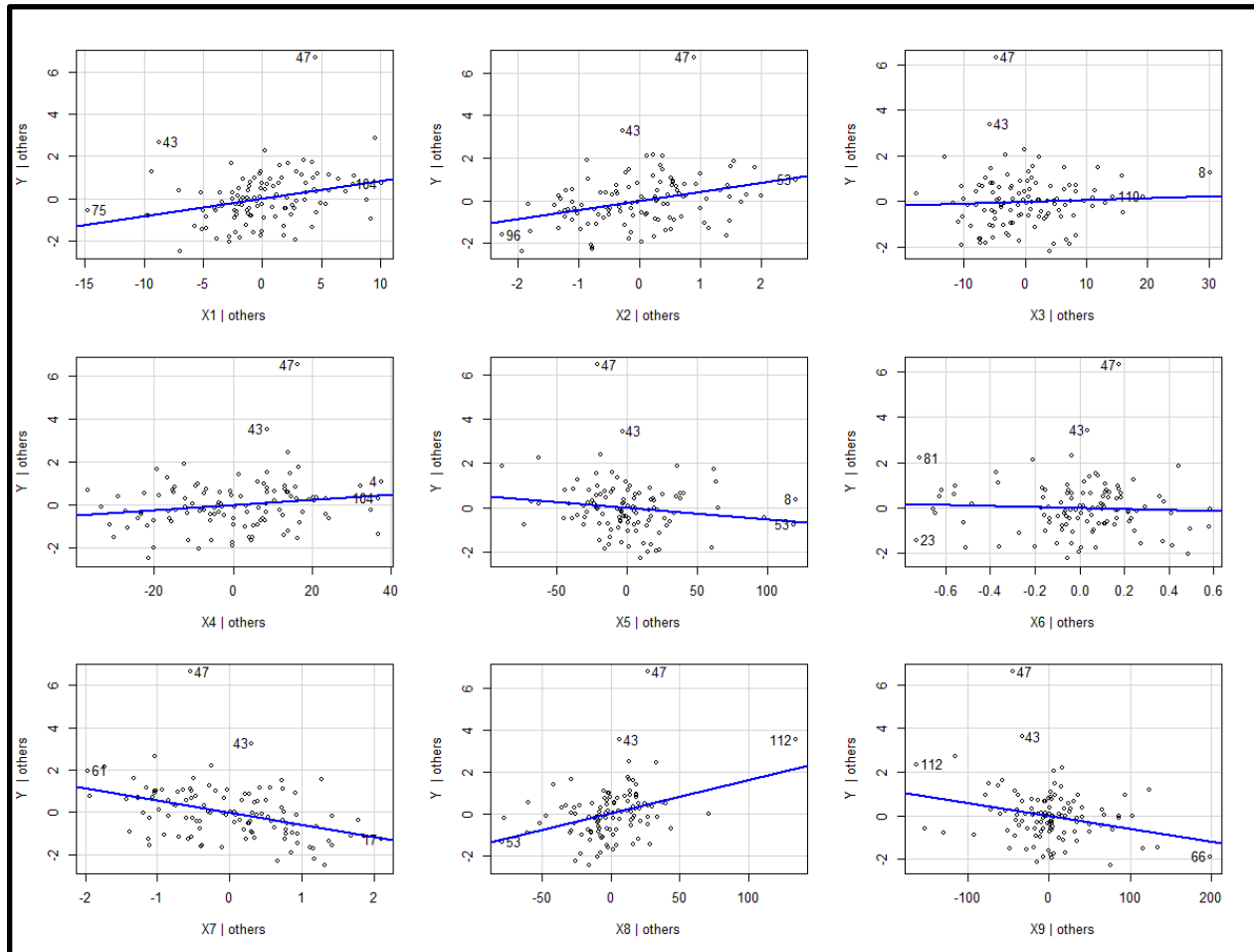


Figure 9. Added-Variable Plots of the response variable against the predictor variables.

The relationship regression lines between the predictor and response variables are provided by this, based on the Added-Variable Plots (Figure 9). Regression analysis reveals that the variables with the highest impact on the response variable are Infection Risk (X2), Average Daily Census (X8), and Routine Culturing Ratio (X3), Medical School (X6). The variables with the lowest impact are others.

Correlation Matrix:

```
> correlation_matrix
```

	Y	X1	X2	X3	X4	X5
Y	1.0000000	0.188913972	0.533443831	0.3266838	0.38248193	0.40926525
X1	0.1889140	1.000000000	0.001093166	-0.2258468	-0.01885490	-0.05882316
X2	0.5334438	0.001093166	1.000000000	0.5591589	0.45339156	0.35977000
X3	0.3266838	-0.225846789	0.559158869	1.0000000	0.42496204	0.13972495
X4	0.3824819	-0.018854897	0.453391557	0.4249620	1.00000000	0.04581997
X5	0.4092652	-0.058823160	0.359770000	0.1397249	0.04581997	1.00000000
X6	-0.2969510	0.145126369	-0.233029901	-0.2427441	-0.08669664	-0.59117997
X7	-0.4921304	-0.020431944	-0.192280702	-0.3082778	-0.29634411	-0.10562663
X8	0.4738855	-0.054774667	0.381411081	0.1429482	0.06291352	0.98099774
X9	0.3403671	-0.082944616	0.393981340	0.1988998	0.07738133	0.91550415
X10	0.3555379	-0.040451379	0.412600675	0.1851311	0.11192761	0.79452438

	X6	X7	X8	X9	X10
Y	-0.29695100	-0.49213043	0.47388550	0.34036706	0.35553792
X1	0.14512637	-0.02043194	-0.05477467	-0.08294462	-0.04045138
X2	-0.23302990	-0.19228070	0.38141108	0.39398134	0.41260068
X3	-0.24274409	-0.30827778	0.14294821	0.19889983	0.18513114
X4	-0.08669664	-0.29634411	0.06291352	0.07738133	0.11192761
X5	-0.59117997	-0.10562663	0.98099774	0.91550415	0.79452438
X6	1.00000000	0.10266758	-0.61475733	-0.58823974	-0.52439032
X7	0.10266758	1.00000000	-0.15274400	-0.11268137	-0.21153192
X8	-0.61475733	-0.15274400	1.00000000	0.90789698	0.77806330
X9	-0.58823974	-0.11268137	0.90789698	1.00000000	0.78350550
X10	-0.52439032	-0.21153192	0.77806330	0.78350550	1.00000000

Figure 10. Correlation matrix of the response variable against the predictor variables.0.9

Correlation plot:

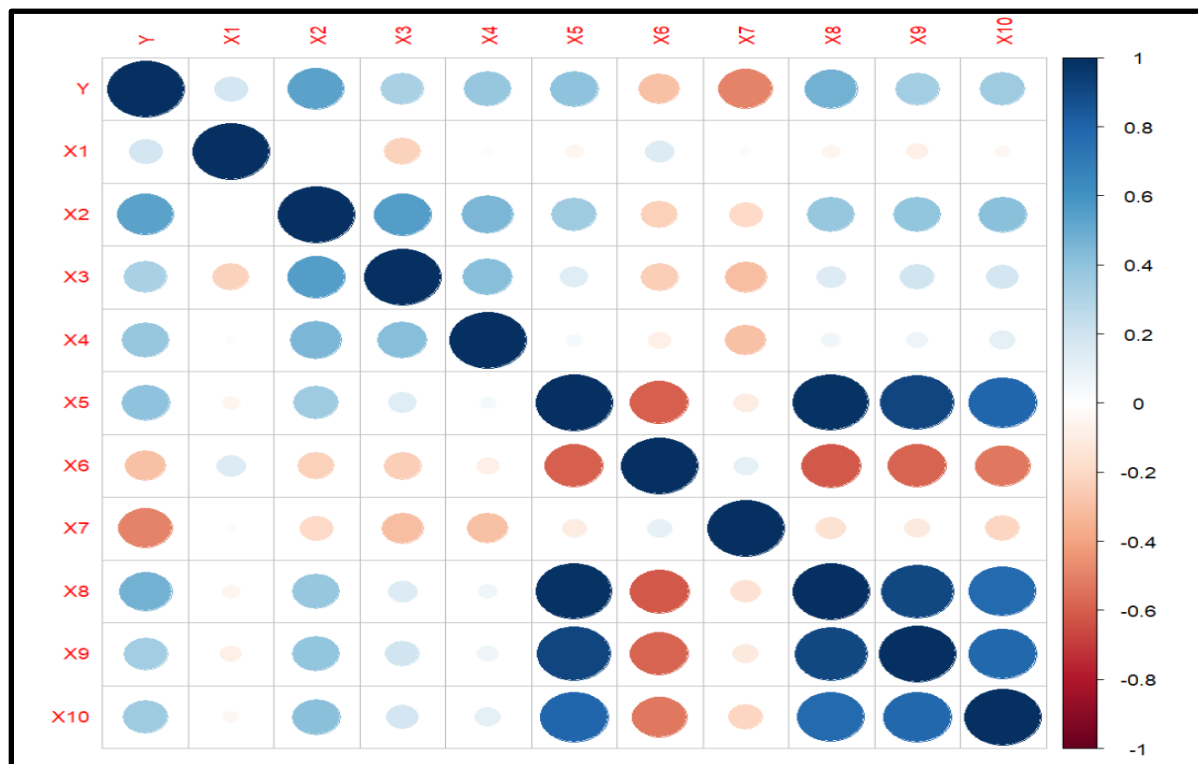


Figure 11. Correlation plot of the response variable against the predictor variables.

The number of beds (X5) has a strong correlation with X8(0.98099774) , X9(0.91550415), and X10(0.79452438), as shown by the correlation matrix and plot (Figures 10&11). A strong correlation has also been observed between X8, the average daily Census, and X9(0.90789698) and X10(0.77806330). In addition, X9—the number of nurses—correlates strongly with X10(0.78350550).

2. MODEL/METHODS

2.1 Model Fitting

Given our dataset with a response variable Y (Length of stay) and predictor variables X_1 to X_{10} , a potential multiple linear regression model could be formulated as follows:

$$Y = \beta_0 + \beta_1 X_1 + \beta_2 X_2 + \beta_3 X_3 + \beta_4 X_4 + \beta_5 X_5 + \beta_6 X_6 + \beta_7 X_7 + \beta_8 X_8 + \beta_9 X_9 + \beta_{10} X_{10} + \epsilon$$

Where:

β_0 is the y-intercept of the regression line.

$\beta_1, \beta_2, \dots, \beta_{10}$ are the coefficients for each predictor variable, representing the change in the response variable for a one-unit change in the predictor, all else being equal.

ϵ is the error term, representing the residual effect unexplained by the predictors.

We will now fit the full model including all predictor variables using the ‘lm’ function in R. This model will serve as a baseline for comparison.

```

> summary(full.lmfit)

Call:
lm(formula = Y ~ X1 + X2 + X3 + X4 + X5 + X6 + X7 + X8 + X9 +
    X10, data = senic)

Residuals:
    Min       1Q   Median       3Q      Max
-2.2346 -0.6592 -0.0699  0.6304  6.3389

Coefficients:
            Estimate Std. Error t value Pr(>|t|)
(Intercept)  3.720403   1.888078   1.970  0.051495 .
X1           0.085177   0.027282   3.122  0.002337 **
X2           0.426433   0.124402   3.428  0.000879 ***
X3           0.007916   0.015634   0.506  0.613704
X4           0.012513   0.007092   1.764  0.080670 .
X5          -0.005403   0.003513  -1.538  0.127110
X6          -0.204155   0.430168  -0.475  0.636091
X7          -0.580146   0.132088  -4.392  2.75e-05 ***
X8           0.015991   0.004282   3.734  0.000311 ***
X9          -0.005853   0.002180  -2.685  0.008463 **
X10         -0.012627   0.013594  -0.929  0.355161
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 1.223 on 102 degrees of freedom
Multiple R-squared:  0.6273,    Adjusted R-squared:  0.5907
F-statistic: 17.16 on 10 and 102 DF,  p-value: < 2.2e-16

```

Figure 12: Full Model

The linear regression model was applied to predict the response variable (Y) based on ten predictor variables (X1 to X10). The linear regression model was fitted with response variable Y and predictors X1 through X10. Among the predictors, X2, X7, and X8 showed statistically significant positive effects on Y, while X5 and X9 had significant negative effects. However, variables X3, X6, and X10 were not statistically significant ($p > 0.05$) and can be removed from the model. The adjusted R-squared was 0.5907, indicating that the model explained approximately 59.07% of the variability in Y. The p-value of X5 was very close to 0.1 so I had to investigate it further.

So now the regression equation from the above data is:

$$Y = 3.720 + 0.085X_1 + 0.426X_2 + 0.0079X_3 + 0.0125X_4 - 0.0054X_5 - 0.204X_6 - 0.580X_7 + 0.016X_8 - 0.0059X_9 - 0.0126X_{10}$$

Now we will test the significance of the model through the ANOVA test.

```
> anova(full.lmfit)
Analysis of Variance Table

Response: Y
      Df Sum Sq Mean Sq F value    Pr(>F)
X1      1  14.604   14.604   9.7660 0.0023154 **
X2      1 116.356  116.356  77.8089 3.284e-14 ***
X3      1   3.248    3.248   2.1720 0.1436244
X4      1   8.606    8.606   5.7549 0.0182590 *
X5      1  31.087   31.087  20.7886 1.430e-05 ***
X6      1   1.514    1.514   1.0124 0.3167176
X7      1  46.675   46.675  31.2122 1.931e-07 ***
X8      1  20.324   20.324  13.5910 0.0003663 ***
X9      1  12.975   12.975   8.6765 0.0039937 **
X10     1   1.290    1.290   0.8628 0.3551614
Residuals 102 152.531    1.495
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

Fig 13: ANNOVA MODEL

The ANOVA table indicates a highly significant F-statistic of 17.16 ($p < 2.2e-16$), confirming the model's overall explanatory power. Significant predictors include X1, X2, X4, X5, X7, X8, and X9, with positive impacts on Y. Notably, X2 has the highest impact ($F = 77.81$, $p < 3.284e-14$), while X5 has a negative effect. In contrast, X3, X6, and X10 do not significantly contribute to Y. The residual mean square is 1.495, representing unexplained variability. Model refinement is suggested, focusing on removing non-significant predictors and validating assumptions like normality and homoscedasticity.

We can see that the model is significant, but some of the individual predictors are not significant. Running the best subset and stepwise regression on this full model results in the following. First, we will look at the best subset for each number of predictor variables selected based on the highest adjusted R2.

3.2 Model Selection/ Step-wise Regression

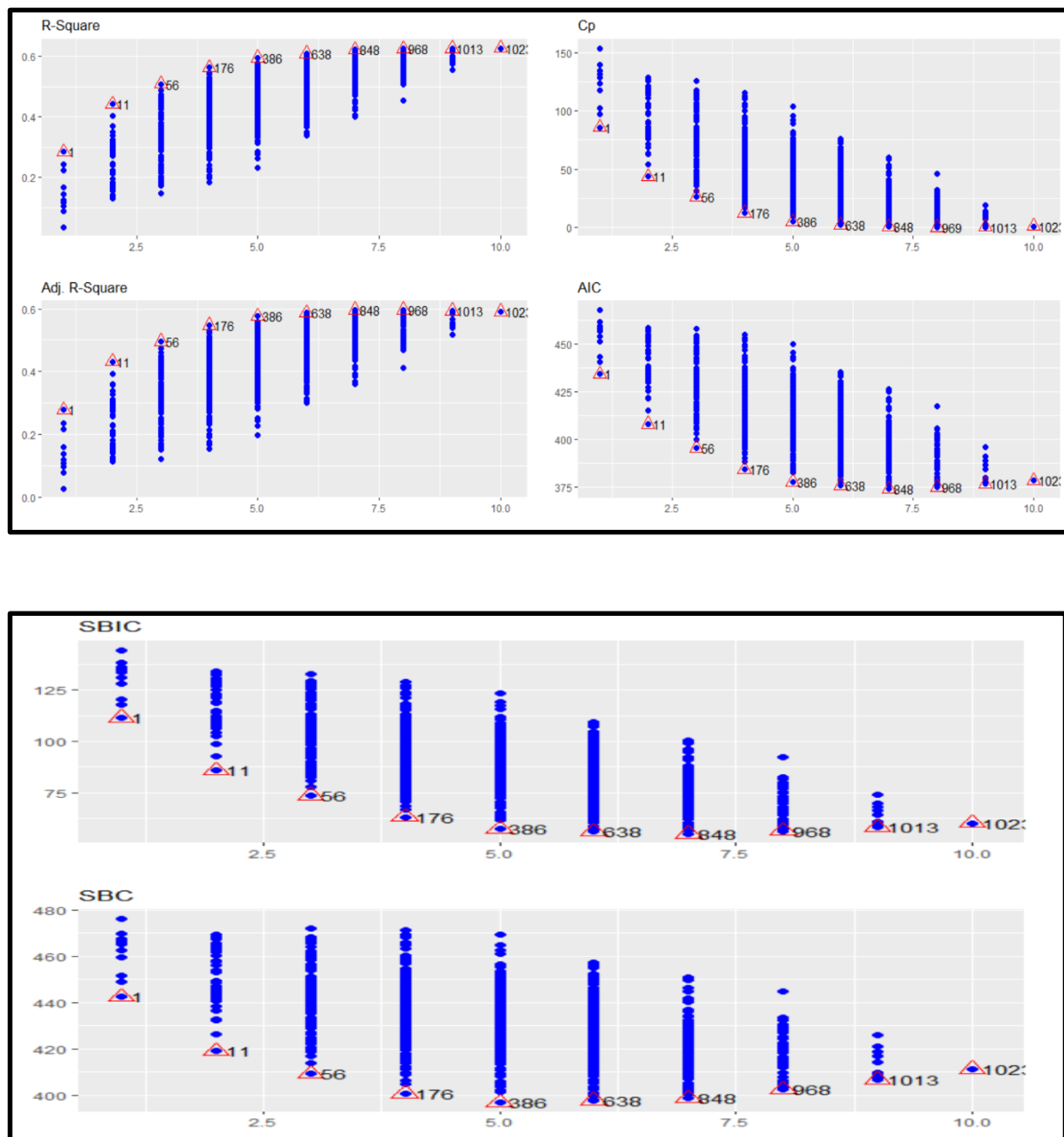


Figure 14: R square, Adjusted \square^{\square} , Mallows' Cp, AIC, BIC, SBIC, SBC

The goal of these criteria is to find a model that has the best trade-off between explaining the data and not becoming overly complex. Overly complex models may fit the current data well but can fail to generalize to new data. These criteria help to identify a model that is expected to have the best predictive performance on data. Comparing the above plot we can come up with the following subset of the full model. We can see that all our procedures agree on a model.

```
> print(b.cp[c(638, 848, 968, 1013, 1023),])
```

	n	predictors	cp
638	6	X1 X2 X4 X7 X8 X9	7.994230
848	7	X1 X2 X4 X5 X7 X8 X9	6.483045
968	8	X1 X2 X4 X5 X7 X8 X9 X10	7.617453
1013	9	X1 X2 X3 X4 X5 X7 X8 X9 X10	9.225240
1023	10	X1 X2 X3 X4 X5 X6 X7 X8 X9 X10	11.000000

```
> print(b.aic[c(638, 848, 968, 1013, 1023),])
```

	n	predictors	aic
638	6	X1 X2 X4 X7 X8 X9	375.9798
848	7	X1 X2 X4 X5 X7 X8 X9	374.2093
968	8	X1 X2 X4 X5 X7 X8 X9 X10	375.2601
1013	9	X1 X2 X3 X4 X5 X7 X8 X9 X10	376.8274
1023	10	X1 X2 X3 X4 X5 X6 X7 X8 X9 X10	378.5781

```
> print(b.press[c(638, 848, 968, 1013, 1023),])
```

	n	predictors	press
638	6	X1 X2 X4 X7 X8 X9	170.6394
848	7	X1 X2 X4 X5 X7 X8 X9	166.6266
968	8	X1 X2 X4 X5 X7 X8 X9 X10	166.8370
1013	9	X1 X2 X3 X4 X5 X7 X8 X9 X10	167.8287
1023	10	X1 X2 X3 X4 X5 X6 X7 X8 X9 X10	169.1170

Figure 15: The subset of the full model.

Stepwise Model Selection:

```
> k <- ols_step_both_p(full.lmfit,pent=0.10,prem=0.1,details=TRUE)
```

Stepwise Selection Method

Candidate Terms:

1. X1
2. X2
3. X3
4. X4
5. X5
6. X6
7. X7
8. X8
9. X9
10. X10

We are selecting variables based on p value...

Stepwise Selection: Step 1

+ X2

Model Summary

R	0.533	RMSE	1.624
R-Squared	0.285	Coef. Var	16.832
Adj. R-Squared	0.278	MSE	2.638
Pred R-Squared	0.254	MAE	1.104

RMSE: Root Mean Square Error

MSE: Mean Square Error

MAE: Mean Absolute Error

ANOVA

	Sum of Squares	DF	Mean Square	F	Sig.
Regression	116.446	1	116.446	44.15	0.0000
Residual	292.765	111	2.638		
Total	409.210	112			

Parameter Estimates

model	Beta	Std. Error	Std. Beta	t	Sig.	lower	upper
(Intercept)	6.337	0.521		12.156	0.000	5.304	7.370
X2	0.760	0.114	0.533	6.645	0.000	0.534	0.987

Stepwise Selection: Step 2

+ X7

Model Summary				
R	0.665	RMSE		1.441
R-Squared	0.442	Coef. Var		14.931
Adj. R-Squared	0.432	MSE		2.075
Pred R-Squared	0.405	MAE		0.955

RMSE: Root Mean Square Error
MSE: Mean Square Error
MAE: Mean Absolute Error

ANOVA					
	Sum of Squares	DF	Mean Square	F	Sig.
Regression	180.930	2	90.465	43.592	0.0000
Residual	228.280	110	2.075		
Total	409.210	112			

Parameter Estimates							
model	Beta	Std. Error	Std. Beta	t	Sig.	lower	upper
(Intercept)	8.630	0.619		13.944	0.000	7.403	9.856
X2	0.650	0.103	0.456	6.279	0.000	0.445	0.855
X7	-0.766	0.137	-0.405	-5.574	0.000	-1.038	-0.494

Model Summary				
R	0.665	RMSE		1.441
R-Squared	0.442	Coef. Var		14.931
Adj. R-Squared	0.432	MSE		2.075
Pred R-Squared	0.405	MAE		0.955

RMSE: Root Mean Square Error
MSE: Mean Square Error
MAE: Mean Absolute Error

ANOVA					
	Sum of Squares	DF	Mean Square	F	Sig.
Regression	180.930	2	90.465	43.592	0.0000
Residual	228.280	110	2.075		
Total	409.210	112			

Parameter Estimates							
model	Beta	Std. Error	Std. Beta	t	Sig.	lower	upper
(Intercept)	8.630	0.619		13.944	0.000	7.403	9.856
X2	0.650	0.103	0.456	6.279	0.000	0.445	0.855
X7	-0.766	0.137	-0.405	-5.574	0.000	-1.038	-0.494

Stepwise Selection: Step 3

+ X8

Model Summary				
R	0.714	RMSE		1.358
R-Squared	0.509	Coef. Var		14.070
Adj. R-Squared	0.496	MSE		1.843
Pred R-Squared	0.456	MAE		0.909

RMSE: Root Mean Square Error
MSE: Mean Square Error
MAE: Mean Absolute Error

ANOVA					
	Sum of Squares	DF	Mean Square	F	Sig.
Regression	208.335	3	69.445	37.683	0.0000
Residual	200.876	109	1.843		
Total	409.210	112			

Parameter Estimates							
model	Beta	Std. Error	Std. Beta	t	Sig.	lower	upper
(Intercept)	8.495	0.584		14.541	0.000	7.337	9.653
X2	0.503	0.105	0.353	4.809	0.000	0.296	0.710
X7	-0.722	0.130	-0.381	-5.555	0.000	-0.980	-0.464
X8	0.003	0.001	0.281	3.856	0.000	0.002	0.005

Model Summary				
R	0.714	RMSE		1.358
R-Squared	0.509	Coef. Var		14.070
Adj. R-Squared	0.496	MSE		1.843
Pred R-Squared	0.456	MAE		0.909

RMSE: Root Mean Square Error
MSE: Mean Square Error
MAE: Mean Absolute Error

ANOVA					
	Sum of Squares	DF	Mean Square	F	Sig.
Regression	208.335	3	69.445	37.683	0.0000
Residual	200.876	109	1.843		
Total	409.210	112			

Parameter Estimates							
model	Beta	Std. Error	Std. Beta	t	Sig.	lower	upper
(Intercept)	8.495	0.584		14.541	0.000	7.337	9.653
X2	0.503	0.105	0.353	4.809	0.000	0.296	0.710
X7	-0.722	0.130	-0.381	-5.555	0.000	-0.980	-0.464
X8	0.003	0.001	0.281	3.856	0.000	0.002	0.005

Stepwise Selection: Step 4

+ X9

Model Summary							
R-Squared	0.751	RMSE				1.286	
Adj. R-Squared	0.547	Coef. Var			13.327		
Pred R-Squared	0.503	MSE			1.653		
		MAE			0.903		
RMSE: Root Mean Square Error							
MSE: Mean Square Error							
MAE: Mean Absolute Error							
ANOVA							
	Sum of Squares	DF	Mean Square	F	Sig.		
Regression	230.640	4	57.660	34.873	0.0000		
Residual	178.571	108	1.653				
Total	409.210	112					
Parameter Estimates							
model	Beta	Std. Error	Std. Beta	t	Sig.	Lower	upper
(Intercept)	8.339	0.555		15.026	0.000	7.239	9.439
X2	0.552	0.100	0.387	5.523	0.000	0.354	0.751
X7	-0.685	0.124	-0.362	-5.542	0.000	-0.930	-0.440
X8	0.010	0.002	0.782	5.115	0.000	0.006	0.013
X9	-0.008	0.002	-0.563	-3.673	0.000	-0.012	-0.004

Model Summary							
R-Squared	0.751	RMSE				1.286	
Adj. R-Squared	0.547	Coef. Var			13.327		
Pred R-Squared	0.503	MSE			1.653		
		MAE			0.903		
RMSE: Root Mean Square Error							
MSE: Mean Square Error							
MAE: Mean Absolute Error							
ANOVA							
	Sum of Squares	DF	Mean Square	F	Sig.		
Regression	230.640	4	57.660	34.873	0.0000		
Residual	178.571	108	1.653				
Total	409.210	112					
Parameter Estimates							
model	Beta	Std. Error	Std. Beta	t	Sig.	Lower	upper
(Intercept)	8.339	0.555		15.026	0.000	7.239	9.439
X2	0.552	0.100	0.387	5.523	0.000	0.354	0.751
X7	-0.685	0.124	-0.362	-5.542	0.000	-0.930	-0.440
X8	0.010	0.002	0.782	5.115	0.000	0.006	0.013
X9	-0.008	0.002	-0.563	-3.673	0.000	-0.012	-0.004

Stepwise Selection: Step 5

+ X1

Model Summary							
R-Squared	0.772	RMSE				1.244	
Adj. R-Squared	0.595	Coef. Var			11.893		
Pred R-Squared	0.576	MSE			1.547		
	0.528	MAE			0.865		
RMSE: Root Mean Square Error							
MSE: Mean Square Error							
MAE: Mean Absolute Error							
ANOVA							
	Sum of Squares	DF	Mean Square	F	Sig.		
Regression	243.634	5	48.727	31.489	0.0000		
Residual	165.576	107	1.547				
Total	409.210	112					
Parameter Estimates							
model	Beta	Std. Error	Std. Beta	t	Sig.	Lower	upper
(Intercept)	4.237	1.514		2.799	0.006	1.236	7.238
X2	0.544	0.097	0.381	5.618	0.000	0.352	0.736
X7	-0.678	0.120	-0.358	-5.672	0.000	-0.915	-0.441
X8	0.009	0.002	0.763	5.153	0.000	0.006	0.013
X9	-0.007	0.002	-0.528	-3.549	0.001	-0.011	-0.003
X1	0.077	0.026	0.179	2.898	0.005	0.024	0.129

Model Summary							
R-Squared	0.772	RMSE				1.244	
Adj. R-Squared	0.595	Coef. Var			11.893		
Pred R-Squared	0.576	MSE			1.547		
	0.528	MAE			0.865		
RMSE: Root Mean Square Error							
MSE: Mean Square Error							
MAE: Mean Absolute Error							
ANOVA							
	Sum of Squares	DF	Mean Square	F	Sig.		
Regression	243.634	5	48.727	31.489	0.0000		
Residual	165.576	107	1.547				
Total	409.210	112					
Parameter Estimates							
model	Beta	Std. Error	Std. Beta	t	Sig.	Lower	upper
(Intercept)	4.237	1.514		2.799	0.006	1.236	7.238
X2	0.544	0.097	0.381	5.618	0.000	0.352	0.736
X7	-0.678	0.120	-0.358	-5.672	0.000	-0.915	-0.441
X8	0.009	0.002	0.763	5.153	0.000	0.006	0.013
X9	-0.007	0.002	-0.528	-3.549	0.001	-0.011	-0.003
X1	0.077	0.026	0.179	2.898	0.005	0.024	0.129

Stepwise Selection: Step 6

+ X4

Model Summary			
R-Squared	0.780	RMSE	1.229
Adj. R-Squared	0.609	Coef. Var	12.734
Pred R-Squared	0.587	MSE	1.509
	0.536	MAE	0.849

RMSE: Root Mean Square Error
MSE: Mean Square Error
MAE: Mean Absolute Error

ANOVA					
	Sum of Squares	DF	Mean Square	F	Sig.
Regression	249.211	6	41.535	27.517	0.0000
Residual	160.000	106	1.509		
Total	409.210	112			

Parameter Estimates							
model	Beta	Std. Error	Std. Beta	t	Sig.	Lower	Upper
(Intercept)	3.241	1.583		2.048	0.043	0.104	6.379
X2	0.453	0.107	0.318	4.247	0.000	0.241	0.664
X7	-0.618	0.122	-0.327	-5.064	0.000	-0.860	-0.376
X8	0.010	0.002	0.786	5.356	0.000	0.006	0.013
X9	-0.007	0.002	-0.530	-3.609	0.000	-0.011	-0.003
X1	0.079	0.026	0.184	3.003	0.003	0.027	0.131
X4	0.013	0.007	0.137	1.922	0.057	0.000	0.027

Model Summary			
R-Squared	0.780	RMSE	1.229
Adj. R-Squared	0.609	Coef. Var	12.734
Pred R-Squared	0.587	MSE	1.509
	0.536	MAE	0.849

RMSE: Root Mean Square Error
MSE: Mean Square Error
MAE: Mean Absolute Error

ANOVA					
	Sum of Squares	DF	Mean Square	F	Sig.
Regression	249.211	6	41.535	27.517	0.0000
Residual	160.000	106	1.509		
Total	409.210	112			

Parameter Estimates							
model	Beta	Std. Error	Std. Beta	t	Sig.	Lower	Upper
(Intercept)	3.241	1.583		2.048	0.043	0.104	6.379
X2	0.453	0.107	0.318	4.247	0.000	0.241	0.664
X7	-0.618	0.122	-0.327	-5.064	0.000	-0.860	-0.376
X8	0.010	0.002	0.786	5.356	0.000	0.006	0.013
X9	-0.007	0.002	-0.530	-3.609	0.000	-0.011	-0.003
X1	0.079	0.026	0.184	3.003	0.003	0.027	0.131
X4	0.013	0.007	0.137	1.922	0.057	0.000	0.027

Stepwise Selection: Step 7

+ X5

Model Summary			
R-Squared	0.789	RMSE	1.214
Adj. R-Squared	0.622	Coef. Var	12.583
Pred R-Squared	0.597	MSE	1.474
	0.545	MAE	0.841

RMSE: Root Mean Square Error
MSE: Mean Square Error
MAE: Mean Absolute Error

ANOVA					
	Sum of Squares	DF	Mean Square	F	Sig.
Regression	254.461	7	36.352	24.665	0.0000
Residual	154.749	105	1.474		
Total	409.210	112			

Parameter Estimates							
model	Beta	Std. Error	Std. Beta	t	Sig.	Lower	Upper
(Intercept)	3.251	1.564		2.079	0.040	0.150	6.351
X2	0.436	0.106	0.306	4.121	0.000	0.226	0.646
X7	-0.571	0.123	-0.302	-4.639	0.000	-0.816	-0.327
X8	0.017	0.004	1.333	4.113	0.000	0.009	0.025
X9	-0.006	0.002	-0.441	-2.891	0.005	-0.010	-0.002
X1	0.079	0.026	0.184	3.051	0.003	0.028	0.130
X4	0.014	0.007	0.137	1.951	0.054	0.000	0.027
X5	-0.006	0.003	-0.632	-1.887	0.062	-0.013	0.000

Model Summary			
R-Squared	0.789	RMSE	1.214
Adj. R-Squared	0.622	Coef. Var	12.583
Pred R-Squared	0.597	MSE	1.474
	0.545	MAE	0.841

RMSE: Root Mean Square Error
MSE: Mean Square Error
MAE: Mean Absolute Error

ANOVA					
	Sum of Squares	DF	Mean Square	F	Sig.
Regression	254.461	7	36.352	24.665	0.0000
Residual	154.749	105	1.474		
Total	409.210	112			

Parameter Estimates							
model	Beta	Std. Error	Std. Beta	t	Sig.	Lower	Upper
(Intercept)	3.251	1.564		2.079	0.040	0.150	6.351
X2	0.436	0.106	0.306	4.121	0.000	0.226	0.646
X7	-0.571	0.123	-0.302	-4.639	0.000	-0.816	-0.327
X8	0.017	0.004	1.333	4.113	0.000	0.009	0.025
X9	-0.006	0.002	-0.441	-2.891	0.005	-0.010	-0.002
X1	0.079	0.026	0.184	3.051	0.003	0.028	0.130

No more variables to be added/removed.

Final Model Output

Model Summary

R	0.789	RMSE	1.214
R-Squared	0.622	Coef. Var	12.583
Adj. R-Squared	0.597	MSE	1.474
Pred R-Squared	0.545	MAE	0.841

RMSE: Root Mean Square Error

MSE: Mean Square Error

MAE: Mean Absolute Error

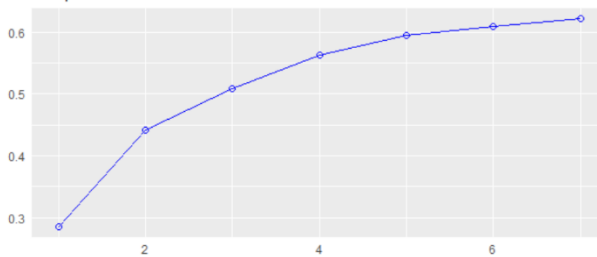
ANOVA

	Sum of Squares	DF	Mean Square	F	Sig.
Regression	254.461	7	36.352	24.665	0.0000
Residual	154.749	105	1.474		
Total	409.210	112			

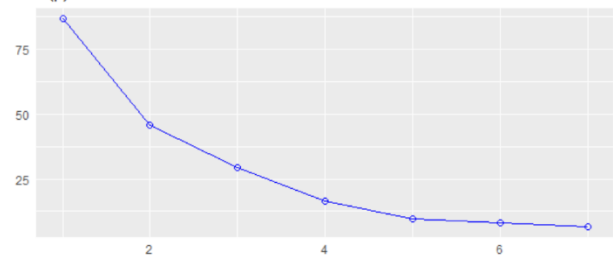
Parameter Estimates

model	Beta	Std. Error	Std. Beta	t	Sig	lower	upper
(Intercept)	3.251	1.564		2.079	0.040	0.150	6.351
X2	0.436	0.106	0.306	4.121	0.000	0.226	0.646
X7	-0.571	0.123	-0.302	-4.639	0.000	-0.816	-0.327
X8	0.017	0.004	1.333	4.113	0.000	0.009	0.025
X9	-0.006	0.002	-0.441	-2.891	0.005	-0.010	-0.002
X1	0.079	0.026	0.184	3.051	0.003	0.028	0.130
X4	0.014	0.007	0.137	1.951	0.054	0.000	0.027
X5	-0.006	0.003	-0.632	-1.887	0.062	-0.013	0.000

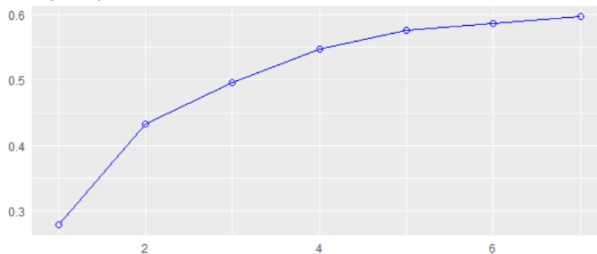
R-Square



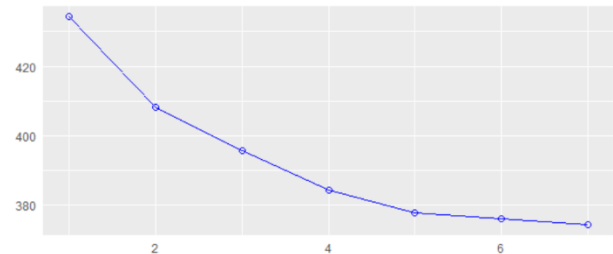
C(p)

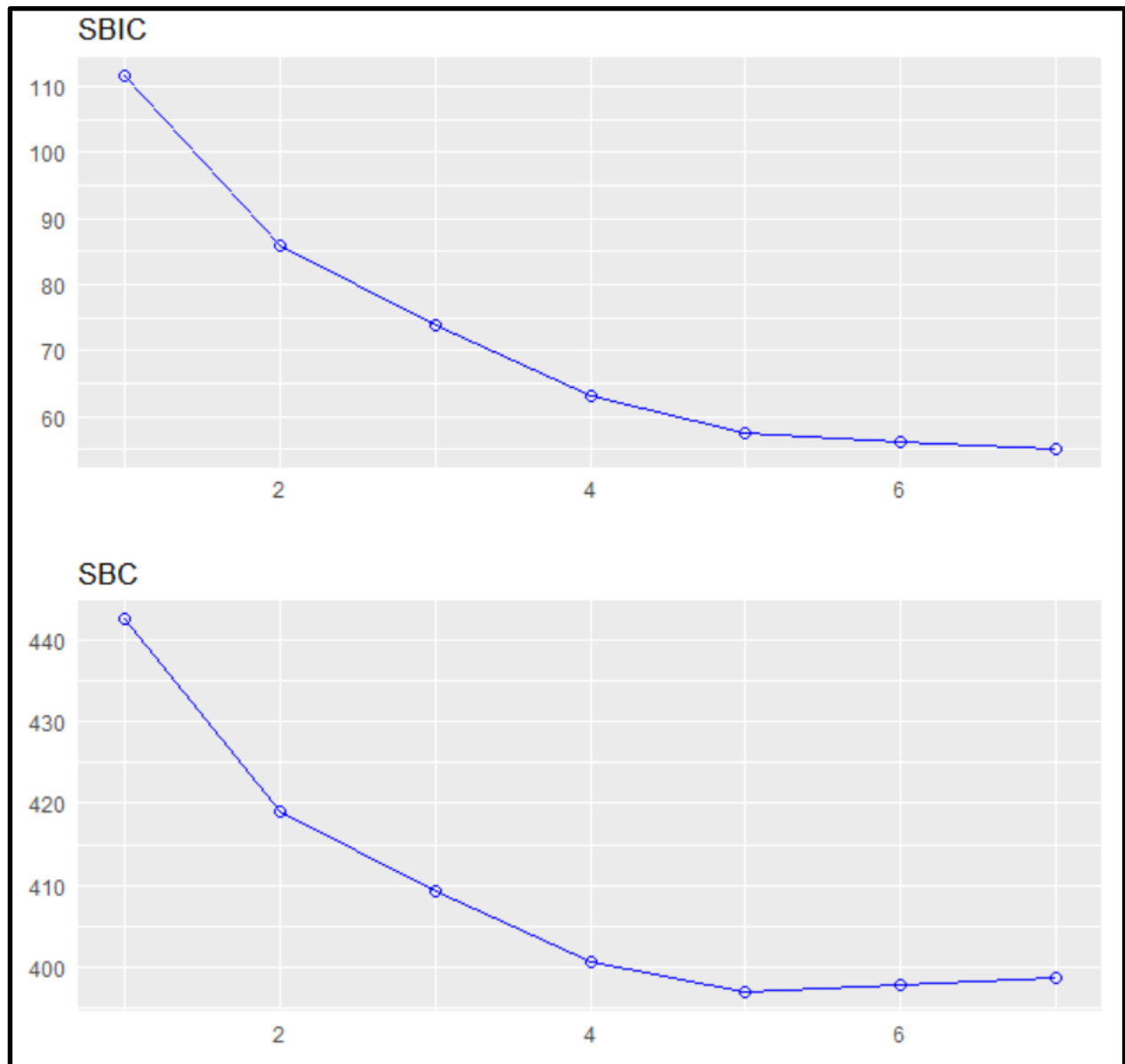


Adj. R-Square



AIC





MODEL EVALUATION

```
> reduced.lmfit <- lm(Y ~ x1+x2+x4+x5+x7+x8+x9, data=senic)
> summary(reduced.lmfit)

Call:
lm(formula = Y ~ x1 + x2 + x4 + x5 + x7 + x8 + x9, data = senic)

Residuals:
    Min       1Q   Median       3Q      Max
-2.1930 -0.6733 -0.0521  0.5819  6.2142

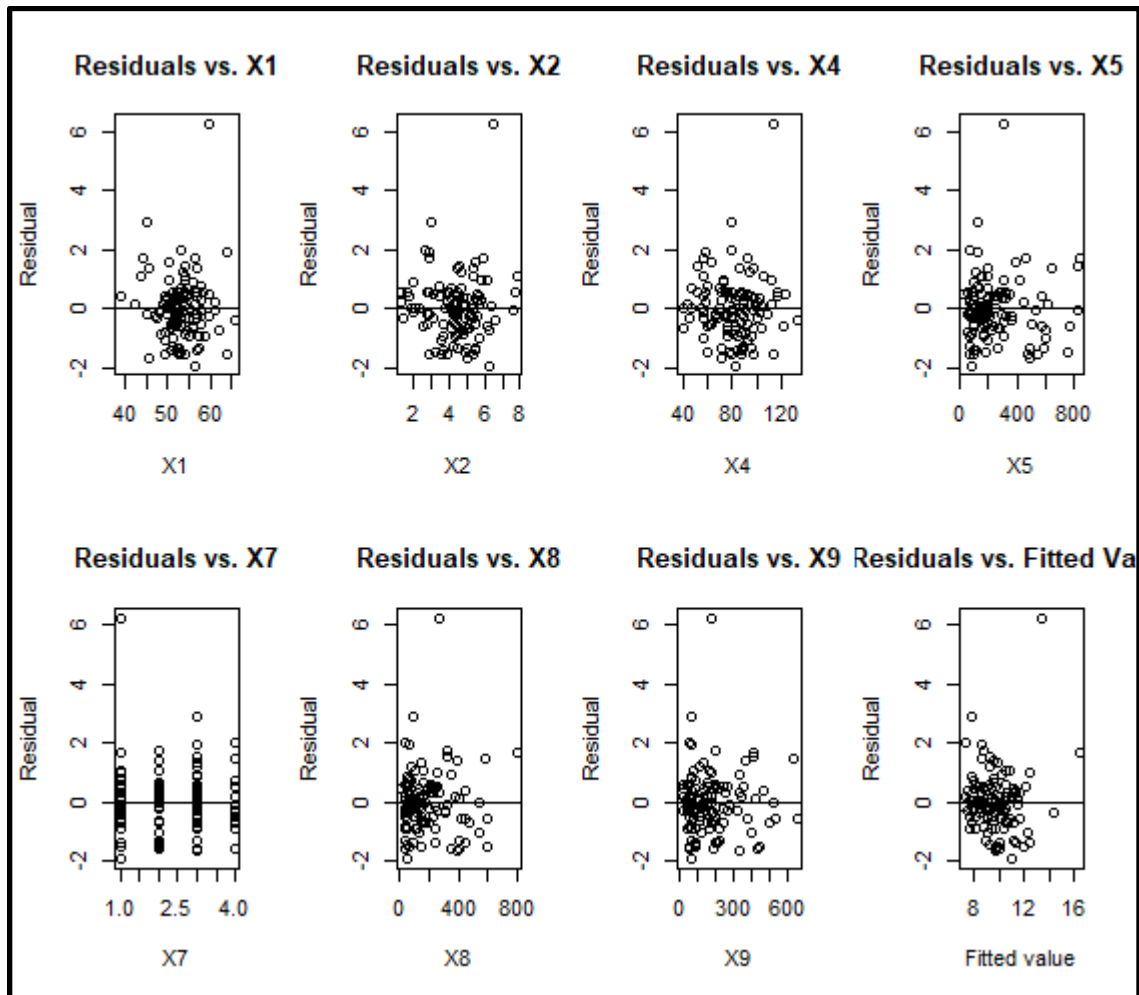
Coefficients:
              Estimate Std. Error t value Pr(>|t|)
(Intercept)   3.250675    1.563726   2.079  0.04007 *
x1             0.078936    0.025869   3.051  0.00289 **
x2             0.435866    0.105757   4.121 7.54e-05 ***
x4             0.013536    0.006939   1.951  0.05376 .
x5            -0.006262    0.003317  -1.887  0.06185 .
x7            -0.571442    0.123172  -4.639 1.01e-05 ***
x8             0.016573    0.004030   4.113 7.79e-05 ***
x9            -0.006059    0.002096  -2.891  0.00467 **
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 1.214 on 105 degrees of freedom
Multiple R-squared:  0.6218,    Adjusted R-squared:  0.5966
F-statistic: 24.67 on 7 and 105 DF,  p-value: < 2.2e-16
```

In this reduced linear regression model, we assess the impact of predictors on the response variable Y. The intercept of 3.2507 is statistically significant ($p = 0.04007$), suggesting that when all predictors are zero, the estimated mean response is 3.2507. Notably, X2 has a substantial positive effect (estimate = 0.4359, $p < 7.54e-05$), indicating that a one-unit increase in X2 is associated with an increase in Y. Conversely, X7 has a negative impact (estimate = -0.5714, $p < 1.01e-05$), suggesting that higher values of X7 correspond to lower values of Y. The model overall is significant (F-statistic = 24.67, $p < 2.2e-16$), explaining 62.18% of the variance in Y. However, attention should be given to predictors X4 and X5, which have marginal significance, and further model refinement is recommended. The residuals exhibit a standard error of 1.214, indicating the unexplained variability in the model, and the adjusted R-squared is 0.5966.

3.3 Model diagnostics/ Regression Diagnostics:

Linearity



Residual plot of Final Model

Looking at our residual vs. fitted value plots, we can see that our model meets the linearity assumption as the residuals are randomly distributed around the fitted values. Similarly, the jackknifed residual vs. predictor value plots also indicates that the linearity assumption is met as the residuals are randomly distributed around the predictor variables. That means our model is a good fit for our data.

The constancy of error Variance

```
> bptest(reduced.lmfit)

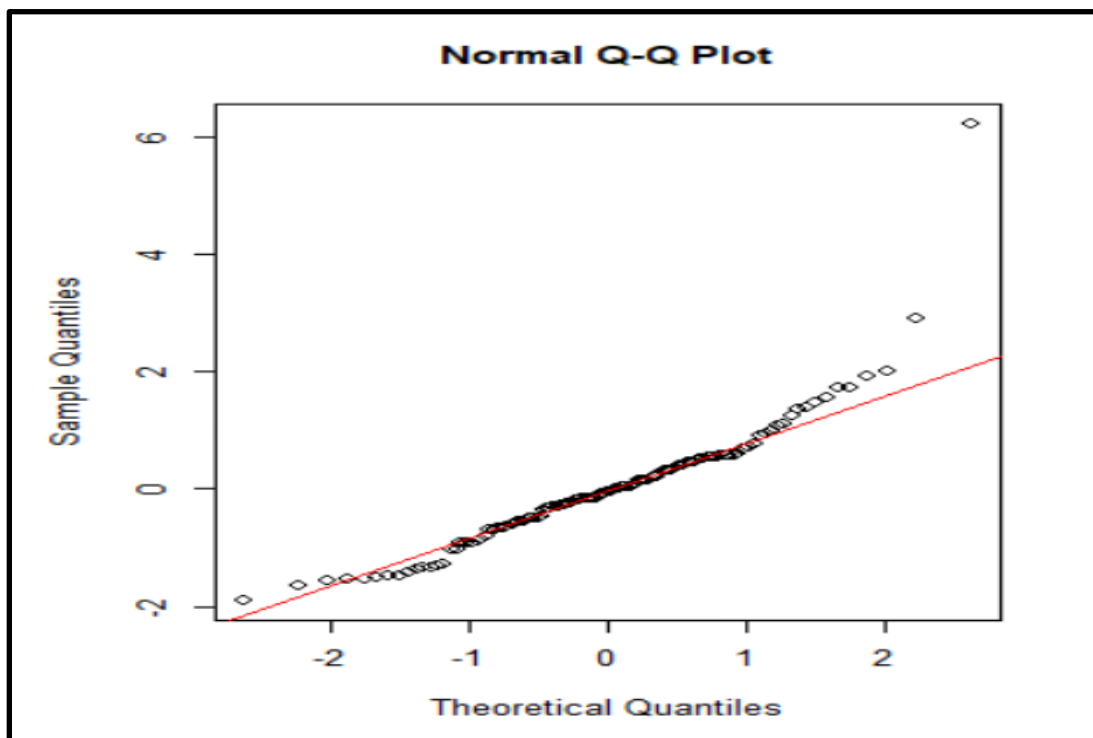
studentized Breusch-Pagan test

data: reduced.lmfit
BP = 7.9695, df = 7, p-value = 0.3353
```

In the Breusch-Pagan test, the null hypothesis states that there is constant error variance and the alternative hypothesis states that there is not constant variance. The decision rule is that if the p-value is less than the significance level of 0.05, we will reject the null hypothesis and conclude that the error variance is not constant. If the p-value is greater than the significance level of 0.05, we will fail to reject the null hypothesis and conclude that the error variance is constant.

We calculated a test statistic of 7.9695 and a p-value of 0.3353 so we failed to reject the null hypothesis and conclude that the error terms are constant.

Normality



```
> shapiro.test(res)

      Shapiro-Wilk normality test

data:  res
W = 0.87698, p-value = 3.23e-08
```

The normal probability plot indicates that the data nearly forms a normal distribution because the data points mostly align with only some slight deviation near the lowest and highest values. In the Shapiro-Wilk test, the null hypothesis states that the error terms are normally distributed while the alternative hypothesis states that the error terms are not normally distributed. Our decision rule states that if the test statistic is small and the p-value is less than the significance level ($\alpha = 0.05$), then we must reject the null hypothesis. If the test statistic is large and the p-value is greater than the significance level, we must fail to reject the null hypothesis.

We calculated a test statistic of 0.87698 and a p-value of 3.23e-08. Thus, we can reject the null hypothesis and conclude that the error terms are not normally distributed.

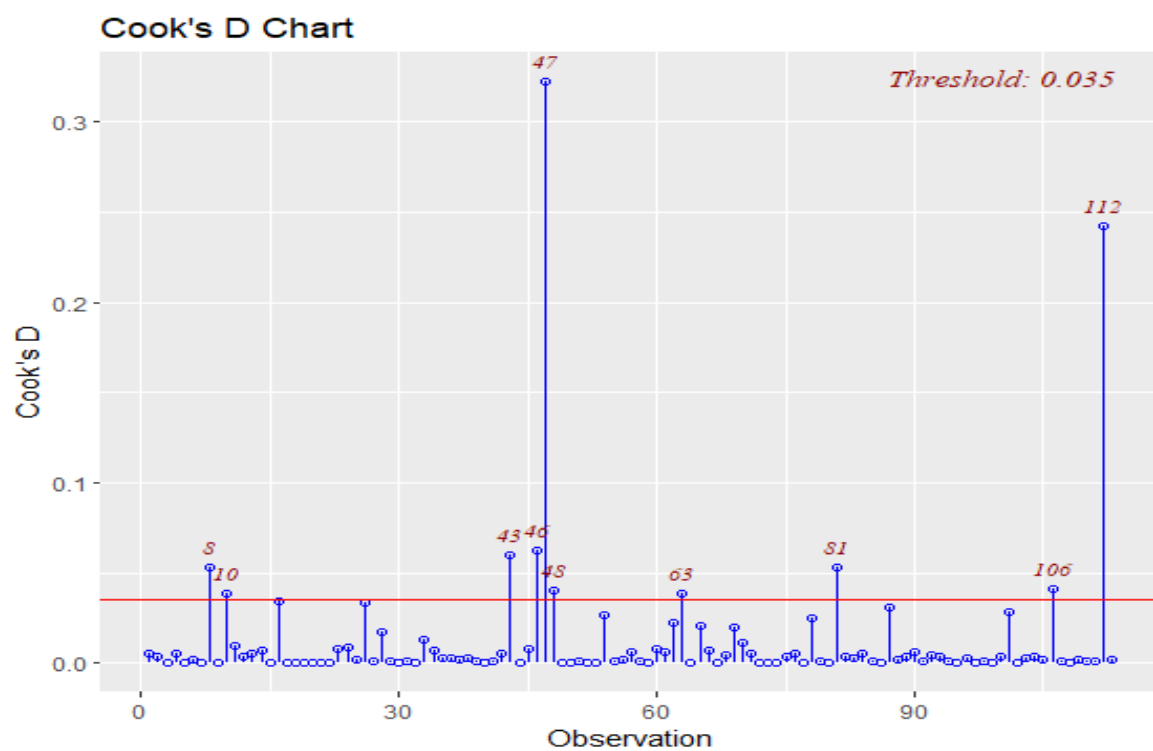
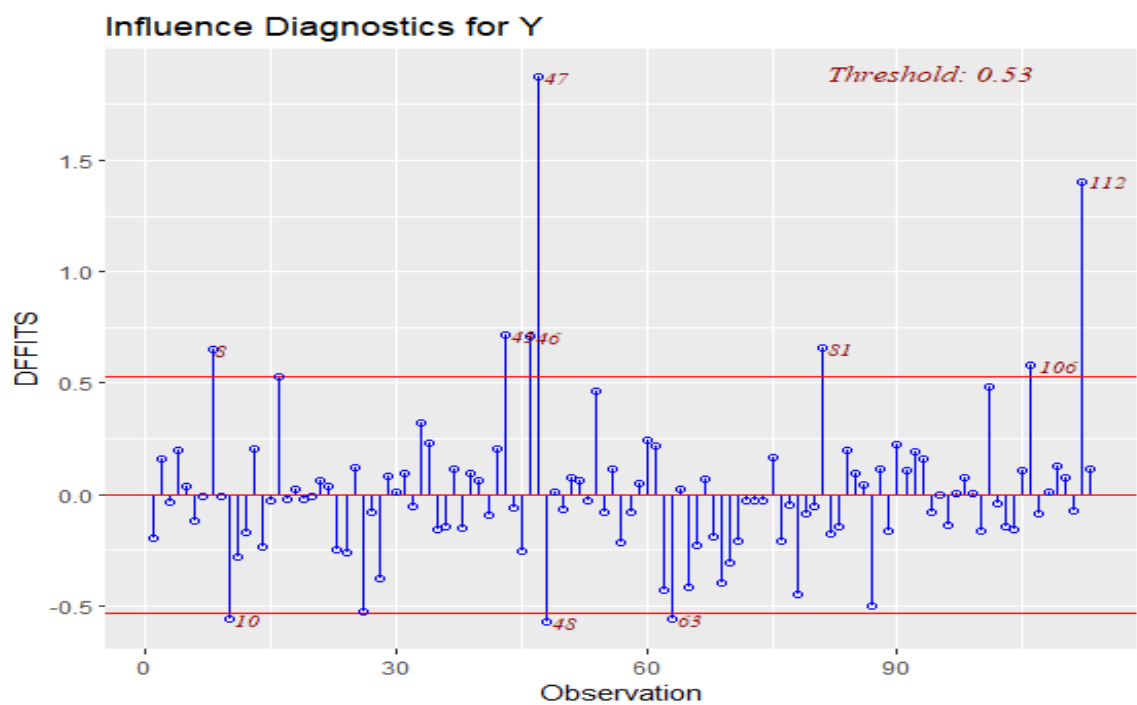
Multicollinearity

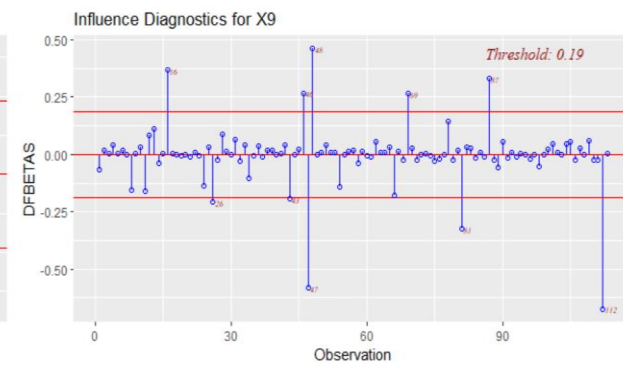
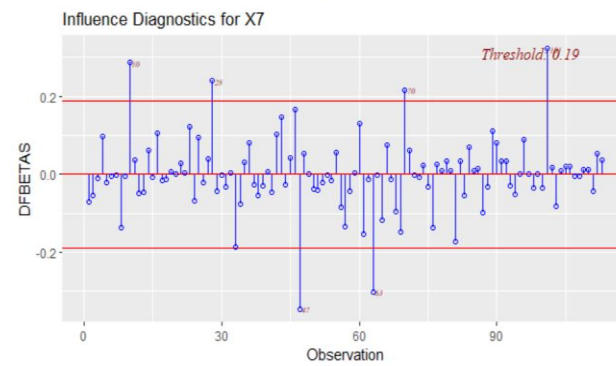
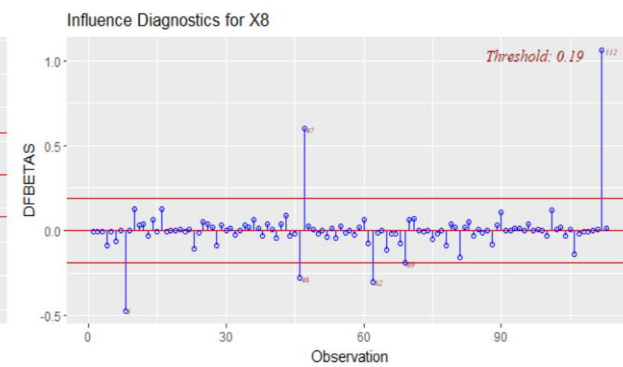
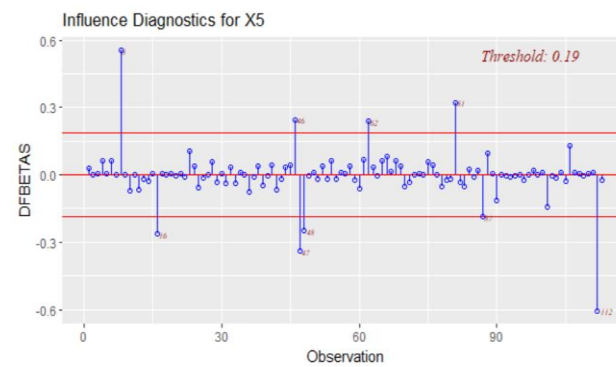
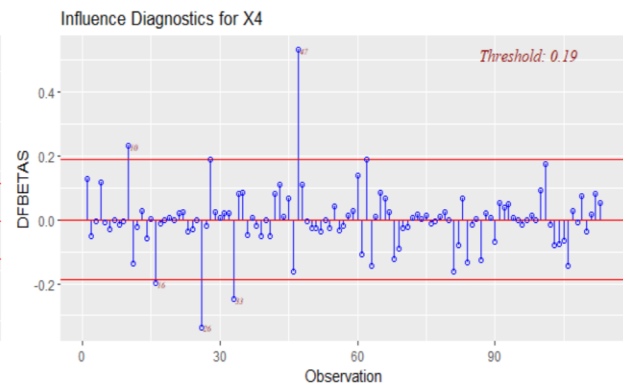
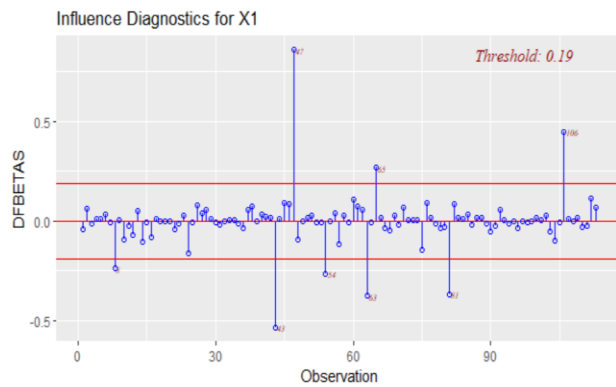
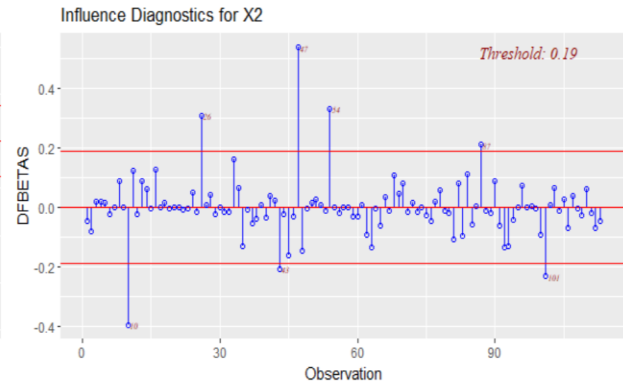
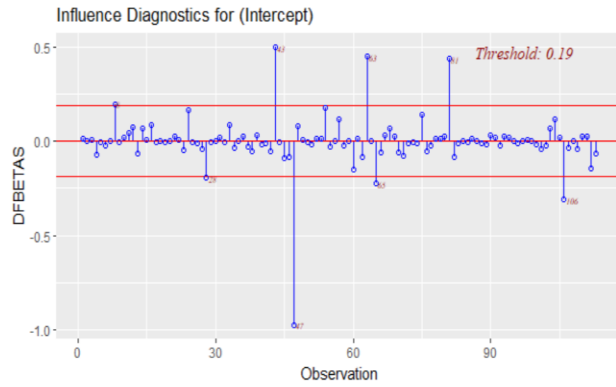
```
> vif(reduced.lmfit)

      x1      x2      x4      x5      x7      x8      x9
1.012338 1.528244 1.372107 31.102062 1.174799 29.173884 6.474638
```

The values obtained from the variance inflation factor analysis indicate that multicollinearity is a major problem in our final model. This is because the VIF values for X5 and X8 are more than 10.

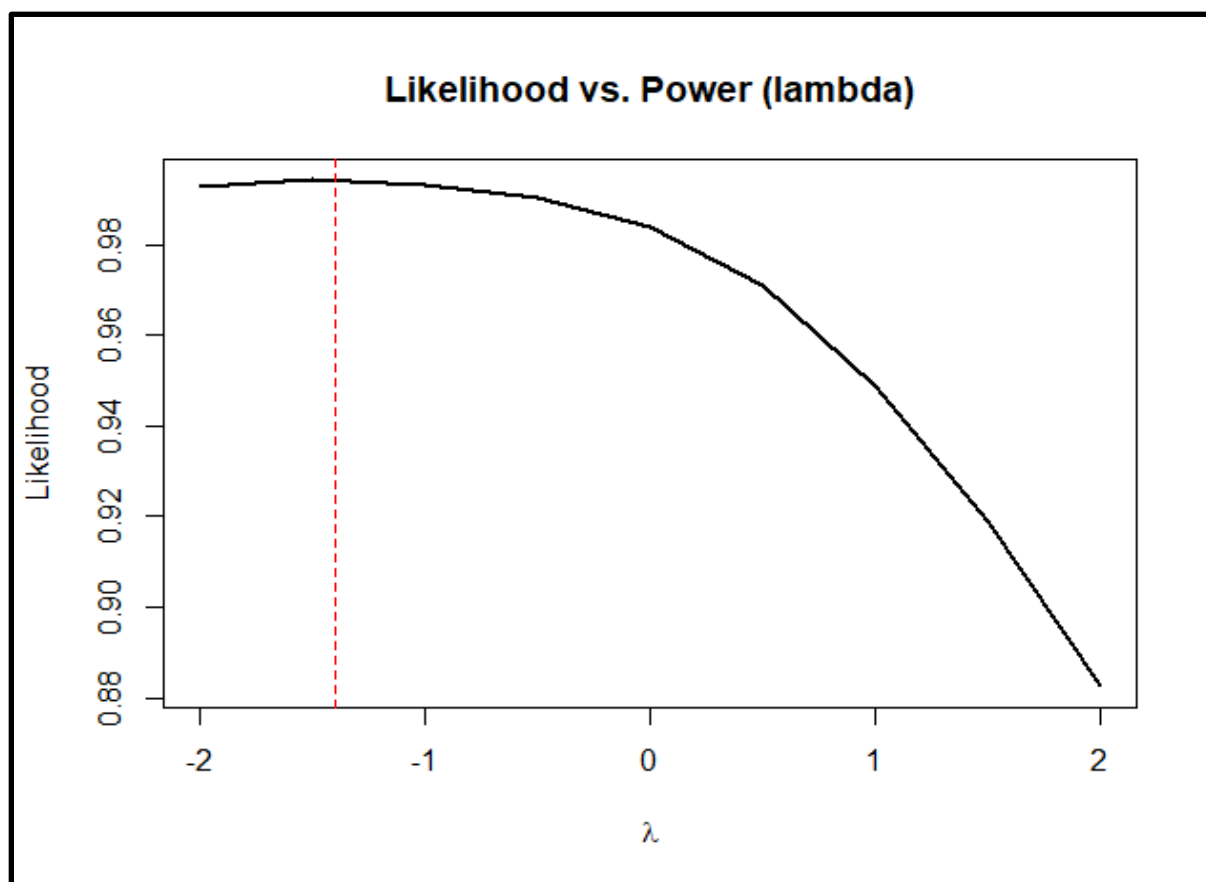
Influential Plots





From the graphs, we can see that there are many influential observations. The Cook's D plot shows that there are many outliers outside of the threshold of 0.035. Some of these major outliers include 8, 10, 43, 46, 47, 48, 63, 81, 112, etc. The DFFITS plot also indicates that there are many outliers both above and below the threshold of 0.53, including 8, 47, 81, 106, 112, etc.

Remedial Actions/Transformations:: Based on our assumption checking, we found that our model satisfied the linearity and the homoscedasticity assumptions. Therefore we don't have to use the method of Weighted Least Squares (WLS). However, a box-cox transformation is required to make the model normal. The box-cox transformation will not correct the multicollinearity discovered earlier. Using the box-cox function in R, the lambda value needed can be determined.



```
> lambda <- boxcox.summary$lambda
> lambda
[1] -1.396303
```

With a lambda value of -1.396303, the response variable can be transformed and a new model created.

```

> summary(boxcox.lmfit)

Call:
lm(formula = trans.Y ~ x1 + x2 + x4 + x5 + x7 + x8 + x9, data = senic)

Residuals:
    Min       1Q   Median       3Q      Max
-0.0199913 -0.0041916 -0.0003834  0.0039982  0.0153018

Coefficients:
            Estimate Std. Error t value Pr(>|t|)
(Intercept)  7.142e-02  8.756e-03   8.157 7.93e-13 ***
x1          -3.339e-04  1.448e-04  -2.305  0.0231 *
x2          -2.401e-03  5.921e-04  -4.054  9.67e-05 ***
x4          -5.376e-05  3.885e-05  -1.384  0.1694
x5           1.233e-05  1.857e-05   0.664  0.5082
x7           3.787e-03  6.897e-04   5.492  2.79e-07 ***
x8          -4.755e-05  2.256e-05  -2.107  0.0375 *
x9           1.558e-05  1.174e-05   1.327  0.1873
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 0.006797 on 105 degrees of freedom
Multiple R-squared:  0.5839,    Adjusted R-squared:  0.5562
F-statistic: 21.05 on 7 and 105 DF,  p-value: < 2.2e-16

```

Fig Summary of Boxcox transformed model

To be sure that the model was truly appropriate for the data, the assumptions for linearity, normality, homoscedasticity, outlier/influential points, and multicollinearity had to be checked again with the transformed values.

Multicollinearity

```

> vif(boxcox.lmfit)

```

	x1	x2	x4	x5	x7	x8	x9
	1.012338	1.528244	1.372107	31.102062	1.174799	29.173884	6.474638

Fig VIF

The values obtained from the variance inflation factor analysis indicate that multicollinearity is a major problem in our final model. This is because the VIF values for X5 and X8 are more than 10. The variance inflation factors did not change between the untransformed and transformed models.

Did the transformation method work?

Our transformed regression model was able to solve the problem of normality. However, our model still has multicollinearity. So we will remove one of both (X5 and X8) as they are highly correlated variables.

Final Model

We tried removing one of X5 And X8. Removing X5 gave best model.

```
> summary(boxcox.lmfit)

Call:
lm(formula = Y ~ X1 + X2 + X4 + X7 + X8 + X9, data = senic)

Residuals:
    Min       1Q   Median       3Q      Max
-2.2738 -0.6768 -0.0659  0.6496  6.3338

Coefficients:
              Estimate Std. Error t value Pr(>|t|)
(Intercept)  3.241386   1.582508   2.048 0.043006 *
X1           0.078625   0.026179   3.003 0.003332 **
X2           0.452894   0.106637   4.247 4.66e-05 ***
X4           0.013498   0.007023   1.922 0.057275 .
X7          -0.618258   0.122099  -5.064 1.75e-06 ***
X8           0.009770   0.001824   5.356 4.97e-07 ***
X9          -0.007281   0.002017  -3.609 0.000471 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

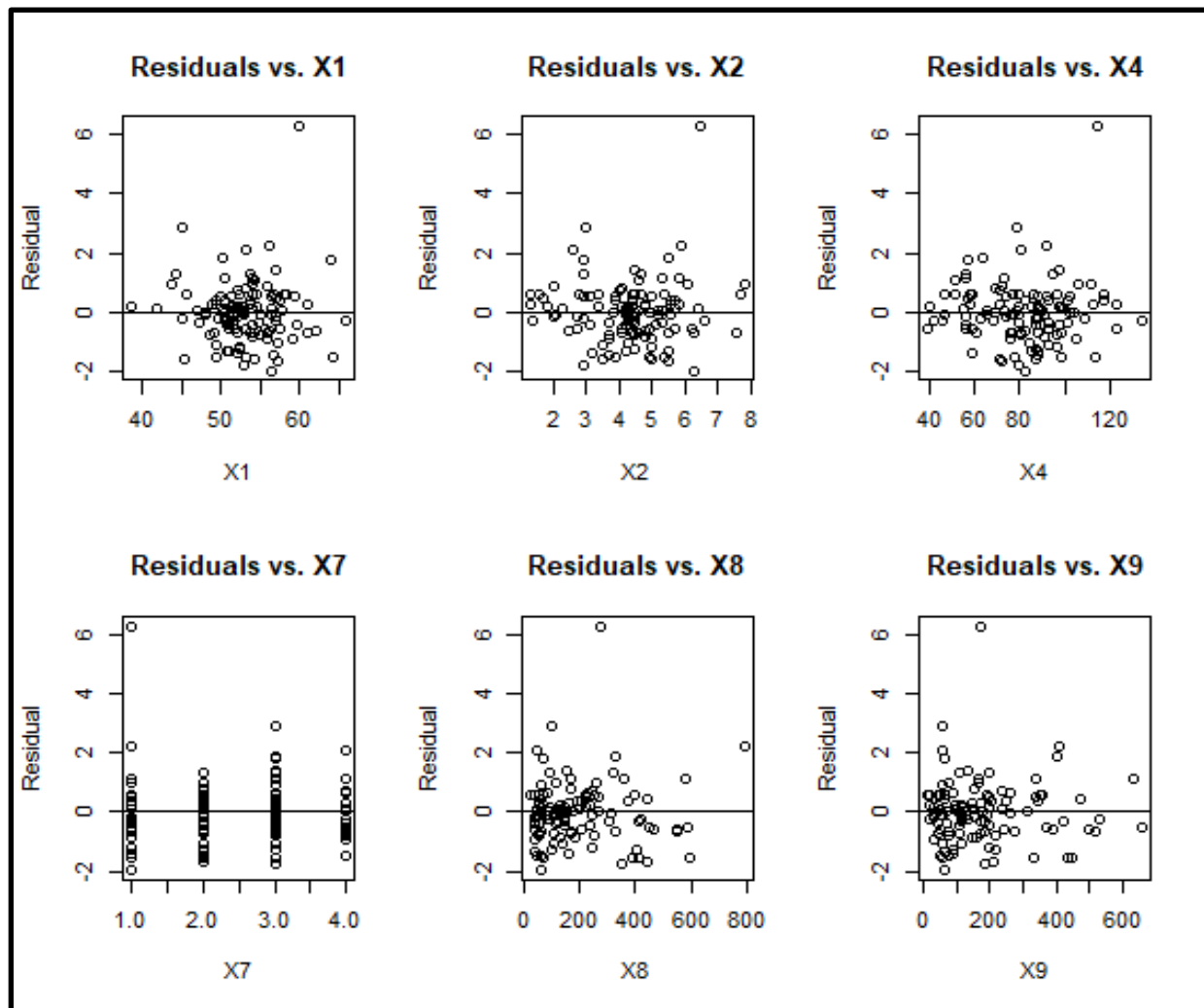
Residual standard error: 1.229 on 106 degrees of freedom
Multiple R-squared:  0.609,    Adjusted R-squared:  0.5869
F-statistic: 27.52 on 6 and 106 DF,  p-value: < 2.2e-16
```

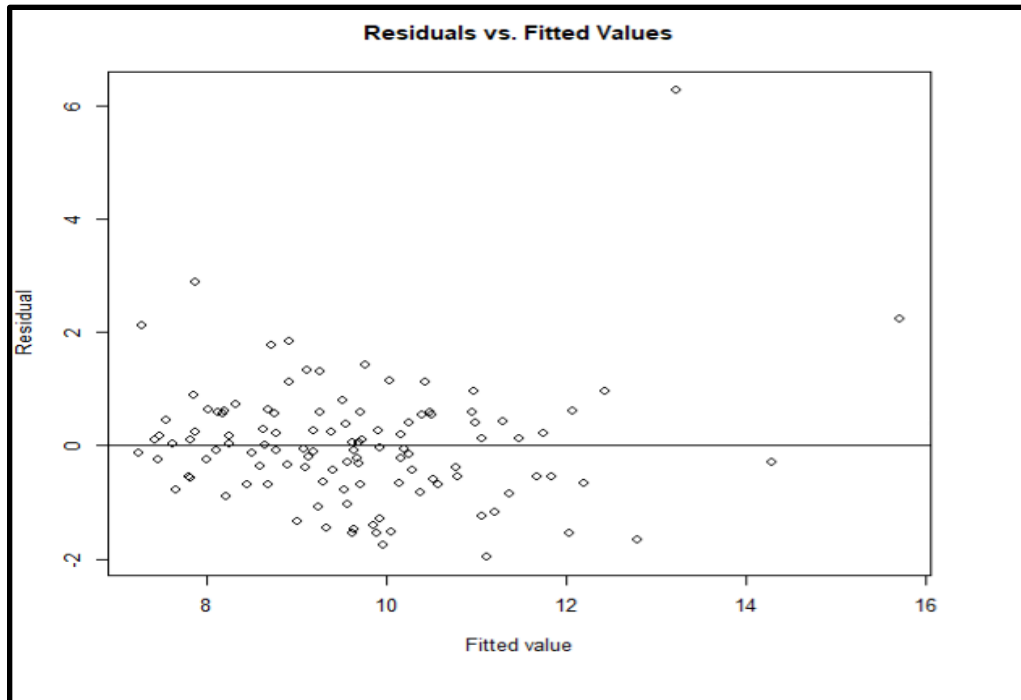
Multicollinearity

```
> vif(boxcox.lmfit)
          X1          X2          X4          X7          X8          X9
1.012297 1.517125 1.372096 1.127161 5.835974 5.856661
```

After removing the highly correlated variable X5, the values obtained from the variance inflation factor analysis indicate that multicollinearity is removed from our final model.

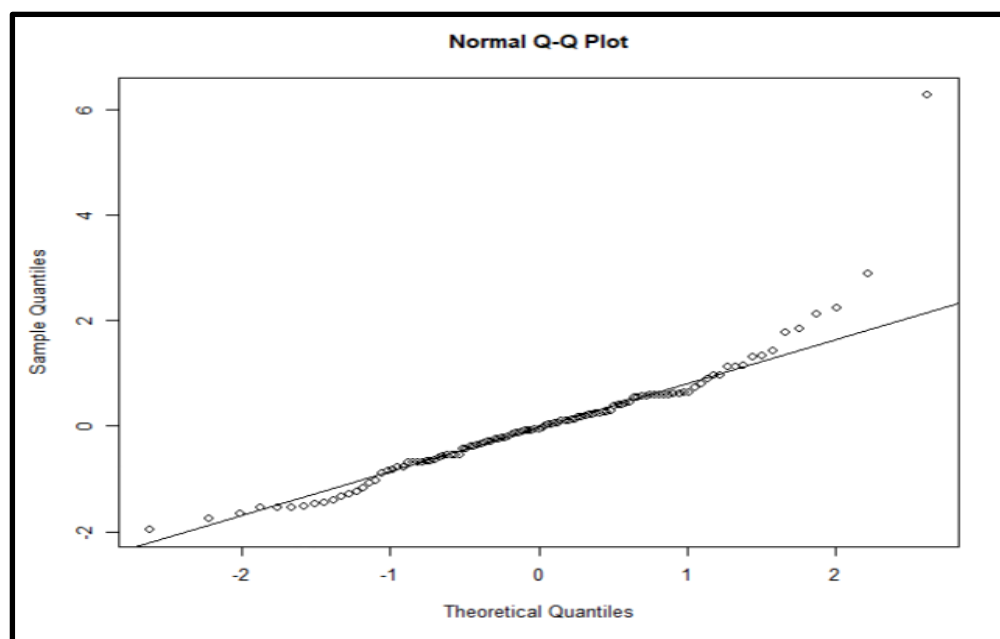
Linearity





To assess the linearity of our chosen transformed model we analyzed the scatter plots between the jackknifed residual and fitted values, we can see that the model meets the linearity assumption as the residuals are randomly distributed around the fitted values. Similarly, the jackknifed residual vs. predictor value plots for the transformed model also indicate that the linearity assumption is met as the residuals are randomly distributed around the predictor variables.

Normality



```
> shapiro.test(boxcox.res)

      Shapiro-Wilk normality test

data:  boxcox.res
W = 0.87439, p-value = 2.487e-08
```

The normal probability plot of the transformed model indicates that the data nearly forms a normal distribution because the data points mostly align with only some slight deviation near the lowest and highest values.

In the Shapiro-Wilk test, the null hypothesis states that the error terms are normally distributed while the alternative hypothesis states that the error terms are not normally distributed. Our decision rule states that if the test statistic is small and the p-value is less than the significance level ($\alpha = 0.05$), then we must reject the null hypothesis. If the test statistic is large and the p-value is greater than the significance level, we must fail to reject the null hypothesis.

For our transformed model:

We calculated a test statistic of 0.87439 and a p-value of 2.487e-08. Thus, we fail to reject the null hypothesis and conclude that the error terms are normally distributed.

Homoscedasticity Assumption

```
> bptest(boxcox.lmfit)

      studentized Breusch-Pagan test

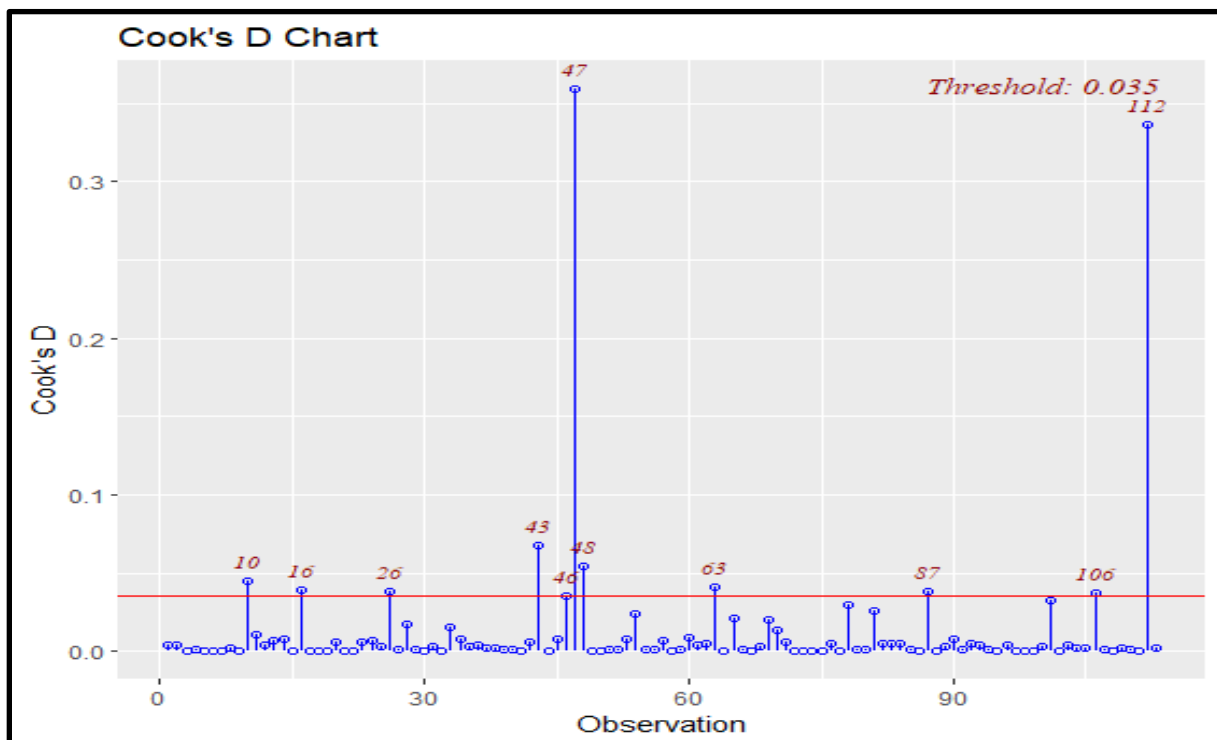
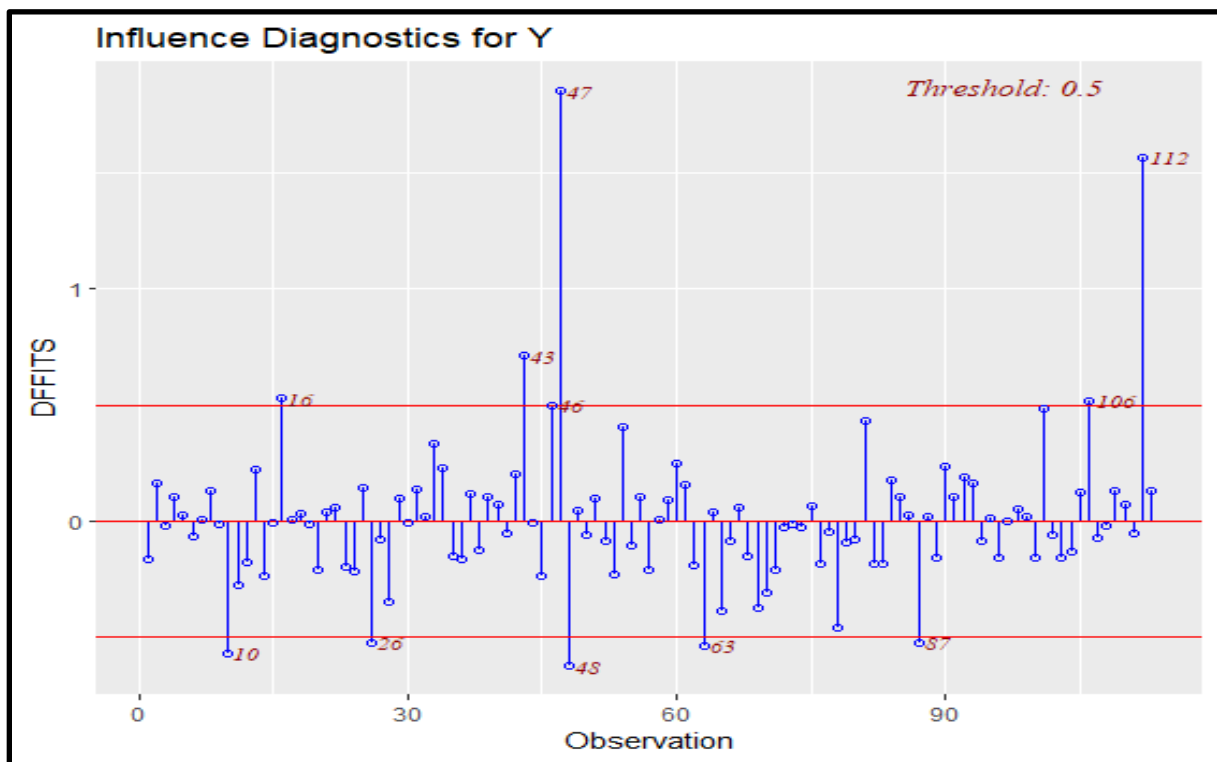
data:  boxcox.lmfit
BP = 9.9119, df = 6, p-value = 0.1284
```

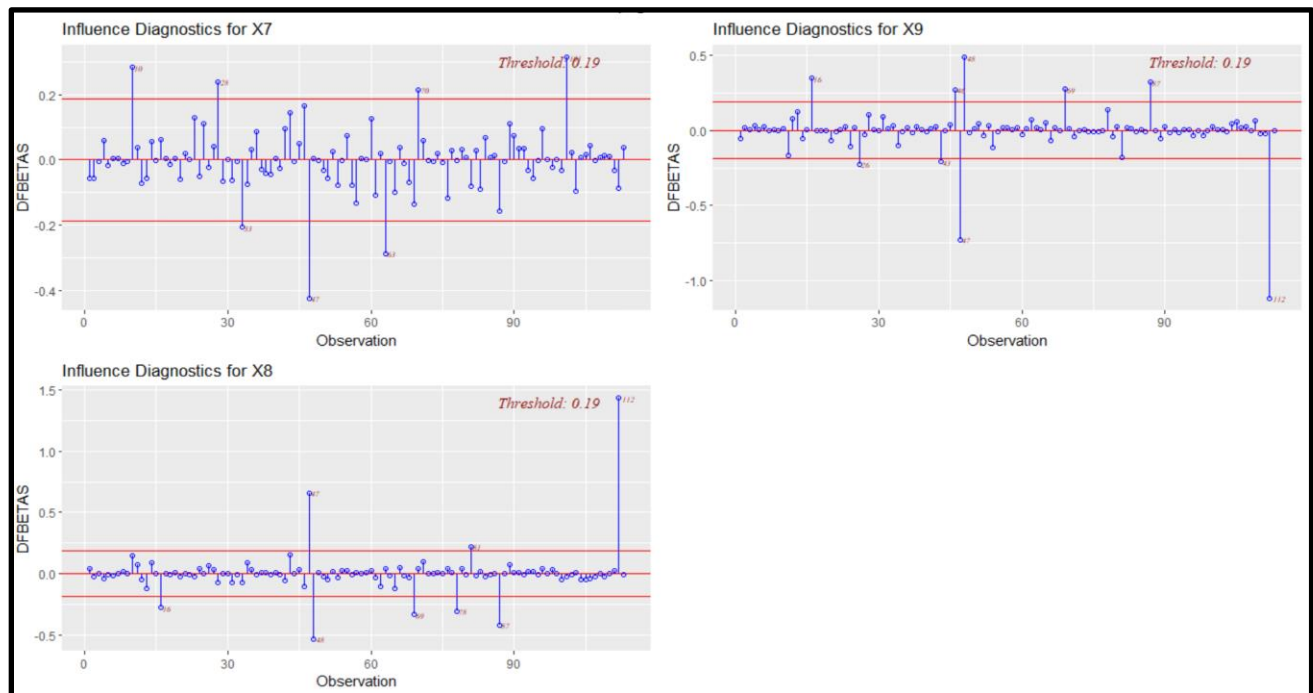
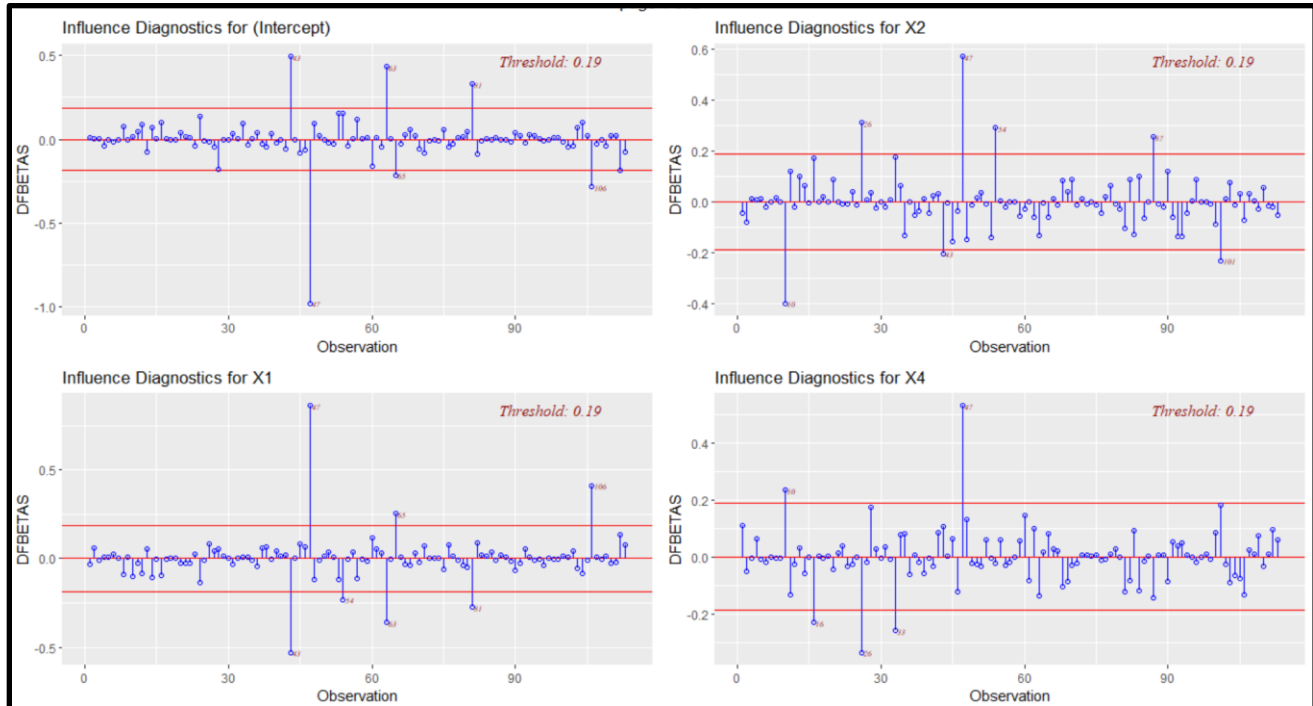
In the Breusch-Pagan test, the null hypothesis states that there is constant error variance and the alternative hypothesis states that there is no constant variance. The decision rule is that if the p-value is less than the significance level of 0.05, we will reject the null hypothesis and conclude that the error variance is not constant. If the p-value is greater than the significance level of 0.05, we will fail to reject the null hypothesis and conclude that the error variance is constant.

For our transformed model:

We calculated a test statistic of 9.9119 and a p-value of 0.1284 so we fail to reject the null hypothesis and conclude that the error terms are constant.

Influential Observations:





From the graphs, we can see that there are many influential observations. The Cook's D plot shows that there are many outliers outside of the threshold of 0.035. Some of these major outliers include 10, 26, 43, 76, 81, 101, 106. The DFFITS plot also indicates that there are many outliers both above and below the threshold of 0.53, including 26, 43, 46, 76, 81, 101, 106, etc.

Result

```
> summary(final.lmfit)

Call:
lm(formula = Y ~ x1 + x2 + x4 + x7 + x8 + x9, data = senic)

Residuals:
    Min       1Q   Median       3Q      Max
-2.2738 -0.6768 -0.0659  0.6496  6.3338

Coefficients:
            Estimate Std. Error t value Pr(>|t|)
(Intercept)  3.241386   1.582508   2.048  0.043006 *
x1           0.078625   0.026179   3.003  0.003332 **
x2           0.452894   0.106637   4.247  4.66e-05 ***
x4           0.013498   0.007023   1.922  0.057275 .
x7          -0.618258   0.122099  -5.064  1.75e-06 ***
x8           0.009770   0.001824   5.356  4.97e-07 ***
x9          -0.007281   0.002017  -3.609  0.000471 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Residual standard error: 1.229 on 106 degrees of freedom
Multiple R-squared:  0.609,    Adjusted R-squared:  0.5869
F-statistic: 27.52 on 6 and 106 DF,  p-value: < 2.2e-16

> anova(final.lmfit)
Analysis of Variance Table

Response: Y
      Df Sum Sq Mean Sq F value    Pr(>F)
x1      1  14.604   14.604   9.6752 0.0023996 **
x2      1 116.356  116.356  77.0859 3.099e-14 ***
x4      1  10.726   10.726   7.1058 0.0088860 **
x7      1  54.518   54.518  36.1185 2.663e-08 ***
x8      1  33.345   33.345  22.0914 7.853e-06 ***
x9      1  19.661   19.661  13.0255 0.0004708 ***
Residuals 106 160.000    1.509
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
> f_value<-qf(0.95, 7, 105, lower.tail=TRUE)
> f_value
[1] 2.098005
> #t-tests for individual regression coefficients
> t_value<-qt(0.975, 105)
> t_value
[1] 1.982815
```

Final model:

$Y = 3.241386 + 0.078625 X_1 + 0.452894 X_2 + 0.013498 X_4 - 0.618258 X_7 + 0.009770 X_8 - 0.007281 X_9.$

F-test: ($\alpha = 0.05$)

Hypothesis:

H_0 : All the regression coefficients = 0 vs H_a : not H_0

$F^* = 27.52$

$f\text{-value}(0.95, 7, 105) = 2.098005$

At a significance level of 0.05, $F^* > f\text{-value}$, so we reject the null hypothesis and conclude that a model with the set of the seven predictor variables is a better fit than an only-intercept model.

Also, $p\text{-value} = 2.2e-16$ which is less than 0.05, so we reject the null hypothesis and conclude that a model with the set of the seven predictor variables is a better fit than an only-intercept model.

Adjusted R2:

The adjusted R^2 value is 0.5869, which means that about 58% (more than half) of the variation in the length of the stay in hospitals can be described by our model.

Significance of individual predictors:

T-test: ($\alpha = 0.05$)

$t\text{-value}(0.975, 105) = 1.982815$

We can see that the absolute value of the t-statistic for variables X4, and X9 is less than the t-value at a significance level of 0.05. Hence, we cannot conclude that they are significant in the response variable. Also,

for X1, $p\text{-value} = 0.0231 < 0.05$ so we conclude that it is significant on the response variable.

For X2, $p\text{-value} = 9.67e-05 < 0.05$ so we conclude that it is significant on the response variable.

For X4, $p\text{-value} = 0.1694 > 0.05$ so we cannot conclude that it is significant on the response variable.

For X7, $p\text{-value} = 2.79e-07 < 0.05$ so we conclude that it is significant on the response variable.

For X8, $p\text{-value} = 0.0375 < 0.05$ so we conclude that it is significant on the response variable.

For X9, $p\text{-value} = 0.1873 > 0.05$ so we cannot conclude that it is significant on the response variable.

Interpretation of Coefficients:

For unit increase in X1, the mean of probability distribution of Y changes by $-3.339e-04$ when X2, X4, X7, X8, X9 are held constant.

For unit increase in X2, the mean of probability distribution of Y changes by $-2.401e-03$ when X1, X4, X7, X8, X9 are held constant.

For unit increase in X4, the mean of the probability distribution of Y changes by $-5.376e-05$ when X1, X2, X7, X8, and X9 are held constant.

For unit increase in X7, the mean of probability distribution of Y changes by $3.787e-03$ when X1, X2, X4, X8, X9 are held constant.

For unit increase in X8, the mean of probability distribution of Y changes by $-4.755e-05$ when X1, X2, X4, X7, X9 are held constant.

For unit increase in X9, the mean of the probability distribution of Y changes by $1.558e-05$ when X1, X2, X4, X7, and X8 are held constant.

Conclusion:

I initially studied our data using visualizations like histograms, boxplots, scatterplots, correlation plots, and added-variance plots before developing a linear regression model. These plots helped me comprehend the skewness of our data and the relationship between the predictors and the response.

Following that, I fitted numerous regression models with various predictor variables to determine the relevance of certain variables based on their p-values. Then I ran model selection to see which variables the stepwise regression function in R recommended we maintain for our linear model. I kept seven predictor variables for the final model based on the AIC, BIC, Adj. R2, and Mallow's CP values.

I next examined the assumptions of this chosen model and discovered that it was not normal. As a result, I conducted a Box-Cox transformation on the altered model and questioned the assumptions. Overall, I found that the modified model was normal. According to the hypothesis test, all seven predictors are significant for the model.

With an adjusted R2 of 0.5869, I picked this changed model as my final model. Based on their age, infection risk, area, the routine x-ray, average census, and the number of nurses and beds, this model may be used to forecast the duration of stay of patients in hospitals.

APPENDIX

Read a data into R

```
senic <- read.csv("C:/Users/ravih/downloads/SENIC.csv", header=TRUE)
head(senic)
```

Check for missing values in the entire dataset

```
missing_values <- sum(is.na(senic))
```

Display the number of missing values

```
cat("Number of missing values in the dataset:", missing_values, "\n")
```

No missing values

Analysing response variable

```
par(mfrow= c(1,1))
```

```
hist(senic$Y)
```

```
boxplot(senic$Y)
```

1. Introduction

1.1 Exploratory Data Analysis.

1. Histograms of Y and Xs

```
library(dplyr)
```

```
par(mfrow= c(3,4))
```

```
for (col in c(names(senic))){
```

```
senic %>% pull(col) %>% hist(main= col)
}
```

```
library(e1071)
skew_summary <- sapply(senic, function(x) skewness(x))
skew_summary
```

2. Boxplots of Y and Xs

```
par(mfrow= c(3,4))
for (col in c(names(senic))){
  senic %>% pull(col) %>% boxplot(main= col)
}
```

3. Summary Statistics

```
summary(senic)
```

4. Scatter Plot Matrix

```
pairs(senic, col= "#FF1493E2", main = "Scatter-Plot matrix of SENIC data")
```

```
par(mfrow=c(3,4))
plot(Y~X1, senic,col="blue", main="Scatter-Plot between Y and X1")
plot(Y~X2, senic,col="blue", main="Scatter-Plot between Y and X2")
```

```
plot(Y~X3, senic,col="blue", main="Scatter-Plot between Y and X3")
plot(Y~X4, senic,col="blue", main="Scatter-Plot between Y and X4")
plot(Y~X5, senic,col="blue", main="Scatter-Plot between Y and X5")
plot(Y~X6, senic,col="blue", main="Scatter-Plot between Y and X6")
plot(Y~X7, senic,col="blue", main="Scatter-Plot between Y and X7")
plot(Y~X8, senic,col="blue", main="Scatter-Plot between Y and X8")
plot(Y~X9, senic,col="blue", main="Scatter-Plot between Y and X9")
plot(Y~X10, senic,col="blue", main="Scatter-Plot between Y and X10")
```

5. Added-Variable Plots

```
library(car)
dev. off()
senic.lmfit <- lm(Y ~ X1+X2+X3+X4+X5+X6+X7+X8+X9+X10, data = senic)
avPlots(senic.lmfit)
```

6. correlation matrix

```
library(caret)
library(corrplot)

#dev.new()
correlation_matrix <- cor(senic)
correlation_matrix
# Create a correlation plot
corrplot(correlation_matrix, method = "circle", diag = TRUE, tl.cex = 0.8)
```

```
#####2. Model/Methods #####
```

```
## Fit a regression model with all of the predictors
```

```
full.lmfit <- lm(Y ~ X1+X2+X3+X4+X5+X6+X7+X8+X9+X10, data = senic)
```

```
summary(full.lmfit)
```

```
anova(full.lmfit)
```

```
## Model Selection (Stepwise Regression)
```

```
# Install packages for the model selection
```

```
# install.packages("leaps")
```

```
# install.packages("HH")
```

```
# install.packages("StepReg")
```

```
# Load HH, leaps, and StepReg packages
```

```
library(leaps)
```

```
library(HH)
```

```
library(StepReg)
```

```
#### Stepwise Regression
```

```
par(mfrow=c(3,2))
```

```
library(olsrr)
```

```
b<- ols_step_all_possible(full.lmfit )
```

```
plot(b)
```

```
b.adjr = data.frame(n=b$n,predictors=b$predictors,adjr=b$adjr)
```

```
#print(b.adjr)
```

```
print(b.adjr[c(638, 848, 968, 1013, 1023),])
```

```
b.cp = data.frame(n=b$n,predictors=b$predictors,cp=b$cp)
```

```
#print(b.cp)
```

```
print(b.cp[c(638, 848, 968, 1013, 1023),])
```

```
b.aic = data.frame(n=b$n,predictors=b$predictors,aic=b$aic)
```

```
#print(b.aic)
```

```
print(b.aic[c(638, 848, 968, 1013, 1023),])
```

```
b.press = data.frame(n=b$n,predictors=b$predictors,press=b$msep)
```

```
#print(b.press)
```

```
print(b.press[c(638, 848, 968, 1013, 1023),])
```

```
k <- ols_step_both_p(full.lmfit,pent=0.10,prem=0.1,details=TRUE)
```

```
plot(k)
```

```
# ##### Checking for correlated variables
```

```
# senic_data <- cbind(senic$Y, x1, x2, x4, x5, x7, x8, x9)
```

```
# senic_data <- as.data.frame(senic_data)
```

```
#
```

```
# # Create a correlation plot
```

```

# dev.new()

# corrplot(cor_matrix, method = "circle", diag = TRUE, tl.cex = 0.8)


library(dplyr)

Cols<- c("X3", "X6", "X10")

senic<-senic[, -which(names(senic) %in% Cols)]


cor_matrix <- cor(senic)

cor_matrix

corrplot(cor_matrix, method = "circle", diag = TRUE, tl.cex = 0.8)


# # As we can see, lots of variables are highly correlated.
# #Therefore Standardization needed for our variables.
#
# x1 <- (senic$X1 -mean(senic$X1))/sd(senic$X1)
# x2 <- (senic$X2 -mean(senic$X2))/sd(senic$X2)
# x3 <- (senic$X3 -mean(senic$X3))/sd(senic$X3)
# x4 <- (senic$X4 -mean(senic$X4))/sd(senic$X4)
# x5 <- (senic$X5 -mean(senic$X5))/sd(senic$X5)
# x6 <- (senic$X6 -mean(senic$X6))/sd(senic$X6)
# x7 <- (senic$X7 -mean(senic$X7))/sd(senic$X7)
# x8 <- (senic$X8 -mean(senic$X8))/sd(senic$X8)
# x9 <- (senic$X9 -mean(senic$X9))/sd(senic$X9)


# #Now very few of them are highly correlated to each other compare to without
# standardization.
# senic.itact.Std <- cbind(senic$Y,x1,x2,x4,x5,x7,x8,x9)
# senic.itact.Std <- as.data.frame(senic.itact.Std) # Converting to data Frame.

```

```

# head(senic.itact.Std)

# colnames(senic.itact.Std)[1] <- "Y"


##### Fit a reduced regression model
# reduced.lmfit <- lm(Y ~ x1+x2+x4+x5+x7+x8+x9, data=senic.itact.Std)
# summary(reduced.lmfit)


reduced.lmfit <- lm(Y ~ X1+X2+X4+X5+X7+X8+X9, data=senic)
summary(reduced.lmfit)


#####3 Regression Diagnostics
res <- rstudent(reduced.lmfit)
fitted.y <- fitted(reduced.lmfit)


##### Residual Plots #####
par(mfrow=c(2,4))
plot(res ~ senic$X1, xlab="X1", ylab="Residual", main="Residuals vs. X1")
abline(h=0)
plot(res ~ senic$X2, xlab="X2", ylab="Residual", main="Residuals vs. X2")
abline(h=0)
plot(res ~ senic$X4, xlab="X4", ylab="Residual", main="Residuals vs. X4")
abline(h=0)
plot(res ~ senic$X5, xlab="X5", ylab="Residual", main="Residuals vs. X5")
abline(h=0)
plot(res ~ senic$X7, xlab="X7", ylab="Residual", main="Residuals vs. X7")

```

```

abline(h=0)
plot(res ~ senic$X8, xlab="X8", ylab="Residual", main="Residuals vs. X8")
abline(h=0)
plot(res ~ senic$X9, xlab="X9", ylab="Residual", main="Residuals vs. X9")
abline(h=0)
plot(res ~ fitted.y, xlab="Fitted value", ylab="Residual", main="Residuals vs. Fitted
Values")
abline(h=0)

##### Normality #####
qqnorm(res);
qqline(res, col= "red")
shapiro.test(res)

##### Constancy of Error Variances #####
library(lmtest)
bptest(reduced.lmfit)

##### Multicollinearity #####
vif(reduced.lmfit)

##### performing transformations as we have high multicollinearity
install.packages("EnvStats")
library(EnvStats)

boxcox.summary <- boxcox(reduced.lmfit, optimize=TRUE)
lambda <- boxcox.summary$lambda
lambda

```



```
trans.Y <- senic$Y^lambda
```

```
senic <- cbind(senic,trans.Y)
```

```
senic
```

```
##### Re-fitting a model using the transformed response variable. #####
```

```
boxcox.lmfit <- lm(trans.Y ~ X1 + X2 + X4 + X5+ X7+ X8 +X9, data=senic)
```

```
summary(boxcox.lmfit)
```

```
boxcox.res <- rstudent(boxcox.lmfit)
```

```
boxcox.fitted.y <- fitted(boxcox.lmfit)
```

```
##### Checking if transformation decreased the multicollinearity problem
```

```
library(car)
```

```
vif(boxcox.lmfit)
```

```
##### Transformation didn't decrease the multi collinearity problem
```

```
## So removing highly correlated variable X5 and fitting the model again
```

```
boxcox.lmfit <- lm(Y ~ X1+X2+X4+X7+X8+X9, data = senic)
```

```
summary(boxcox.lmfit)
```

```
##### Now check the multicollinearity
```

```
library(car)
```

```
vif(boxcox.lmfit)
```

```
#### Stepwise Regression
```

```
par(mfrow=c(3,3))
```

```
library(olsrr)
```

```
b<- ols_step_all_possible(full.lmfit )
```

```
plot(b)
```

```
b.adjr = data.frame(n=b$n,predictors=b$predictors,adjr=b$adjr)
```

```
print(b.adjr)
```

```
print(b.adjr[c(256, 382, 466, 502, 511),])
```

```
b.cp = data.frame(n=b$n,predictors=b$predictors,cp=b$cp)
```

```
print(b.cp)
```

```
print(b.cp[c(256, 382, 466, 502, 511),])
```

```
b.aic = data.frame(n=b$n,predictors=b$predictors,aic=b$aic)
```

```
print(b.aic)
```

```
print(b.aic[c(256, 382, 466, 502, 511),])
```

```
b.press = data.frame(n=b$n,predictors=b$predictors,press=b$msep)
```

```
print(b.press)
```

```
print(b.press[c(256, 382, 466, 502, 511),])
```

```
k <- ols_step_both_p(full.lmfit,pent=0.10,prem=0.1,details=TRUE)
plot(k)
```

```
##### Fitting the reduced model
```

```
reduced.lmfit <- lm(Y ~ x1+x2+x4+x7+x8+x9, data=senic)
summary(reduced.lmfit)
```

```
##### Checking if removing x5 variable removed multi collinearity
```

```
vif(reduced.lmfit)
```

```
##### yes it did
```

```
##### MModel Diagnostics
```

```
##### Residual Plots #####
```

```
final.lmfit <- boxcox.lmfit
```

```
summary(final.lmfit)
```

```
anova(final.lmfit)
```

```
boxcox.res <- rstudent(boxcox.lmfit)
```

```
boxcox.fitted.y <- fitted(boxcox.lmfit)
```

```
par(mfrow=c(2,3))
```

```
plot(boxcox.res ~ senic$X1, xlab="X1", ylab="Residual", main="Residuals vs. X1")
```

```

abline(h=0)

plot(boxcox.res ~ senic$X2, xlab="X2", ylab="Residual", main="Residuals vs. X2")
abline(h=0)

plot(boxcox.res ~ senic$X4, xlab="X4", ylab="Residual", main="Residuals vs. X4")
abline(h=0)

plot(boxcox.res ~ senic$X7, xlab="X7", ylab="Residual", main="Residuals vs. X7")
abline(h=0)

plot(boxcox.res ~ senic$X8, xlab="X8", ylab="Residual", main="Residuals vs. X8")
abline(h=0)

plot(boxcox.res ~ senic$X9, xlab="X9", ylab="Residual", main="Residuals vs. X9")
abline(h=0)

plot(boxcox.res ~ boxcox.fitted.y, xlab="Fitted value", ylab="Residual", main="Residuals
vs. Fitted Values")
abline(h=0)


##### Normality #####

qqnorm(boxcox.res);
qqline(boxcox.res)
shapiro.test(boxcox.res)


##### Constancy of Error Variances #####

library(lmtest)
bptest(boxcox.lmfit)


# 1. DFFITS
ols_plot_dffits(boxcox.lmfit)

```

2. Cook's D

```
ols_plot_cooksd_chart(boxcox.lmfit)
```

3. DFBETAS

```
ols_plot_dfbetas(boxcox.lmfit)
```

```
##### Multicollinearity #####
```

```
library(car)
```

```
vif(boxcox.lmfit)
```