**Assignment Part Two** Chun-Ting Wu r0915592

**Statistical Methods for Bioinformatics**  Yu-Jie Qiu r0823712

1. **Study and describe the predictor variables. Do you see any issues that are relevant for making predictions?**

|  |  |  |
| --- | --- | --- |
| **Attributes** | **Description** | **Mean±SD** |
| Cscore | progression of the cancer score | 36.152±52.72 |
| lcavol | log of cancer volume (𝑐𝑐) | 1.350±1.179 |
| lweight | log of prostate weight (𝑔) | 3.629±0.428 |
| age | age of a patient (years) | 68.866±7.445 |
| lbph | log of the amount of benign prostatic hyperplasia (BPH) (𝑐𝑚2). A noncancerous enlargement of the prostate gland, as an area in a digitized image | 0.100±1.451 |
| svi | Seminal vesicle invasion; 1=Yes, 0=No.  Indicator of whether prostate cancer cells have invaded the seminal vesicle. | {'0'=76, '1'=21} |
| lcp | log of capsular penetration (𝑐𝑚).  Represents the level of extension of cancer into the capsule (the fibrous tissue which acts as an outer lining of the prostate gland) | -0.179±1.398 |
| lpsa | log Prostate specific antigen (PSA) (𝑛𝑔/𝑚𝐿) | 2.478±1.154 |

The dataset comprises 97 clinical measurements and 8 variables. Table 1 provides a summary of the variables, including a brief statistical description. Notably, *svi* (seminal vesicle invasion) is represented as a dummy variable (no = 0 / yes = 1).

Table 1. Description and statistics summary of all eight attributes for prostate cancer dataset.

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自動產生的描述The distribution of the response variable, Cscore, is shown in Figure 1, ranging from -19.473 to 373. The Shapiro test was used to assess the normality of the distribution, and the results showed that the Cscore was not normally distributed (p-value = 4.139e-13).

Figure 1. The distribution of Cscore

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
|  | Cscore | | | |
| group | **N** | **mean** | **std. error** | **Variance** |
| 0 | 76 | 20.5 | 24.1 | 579 |
| 1 | 21 | 92.8 | 82.9 | 6870 |

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自動產生的描述Figure 2 and Table 2 compare the Cscore between group 0 and group 1. As the Cscore distribution is non-normal, we performed the Mann-Whitney U test to analyze whether the scores differed between the two groups. The results showed that the Cscore in group 0 was significantly lower than in group 1 (p < 0.001). However, the number of patients in group 1 (n = 21) was much smaller than in group 0 (n = 76), leading to unequal sample sizes that could cause heterogeneity of variance (F-test to compare two variances: p < 0.05). This heterogeneity could dramatically affect the type I error rate (p-value) and result in a loss of statistical power (type II error), which would decrease the accuracy of prediction. Therefore, caution is needed when interpreting the results.

Table 2. The summary of Cscore in two svi groups.

Figure 2. Comparison of Cscore in 2 svi group

In terms of issues relevant for making predictions, it's important to note that the dataset has a relatively small sample size, with only 97 observations, which may limit the generalizability of any predictive model and may lead to overfitting and hinder accurate predictions.

Additionally, there may be issues with multicollinearity, as some of the predictor variables may be correlated with each other. Multicollinearity refers to high correlation among predictor variables in multiple regression, which can increase the standard errors of the regression coefficients and result in noisy estimates. To validate the effect of correlations between variables, we utilized a scatterplot, which is shown in Figure 3. The results indicate that the correlation between the variables *lcavol* and *lpsa* is higher than 0.7, which can be considered a strong correlation.

To quantify the effect of collinearity on the variance of our regression estimates, we calculated the variance inflation factor (VIF) using the formula VIF = 1/(1 - R-squared). A VIF greater than 5 is typically considered indicative of multicollinearity. Our results indicate that all VIF values are lower than 3, suggesting that there is no multicollinearity issue that need to be addressed in our dataset.

Finally, it's important to carefully consider the interpretation of the natural logarithm transformation applied to some of the variables when interpreting the results of any predictive models.

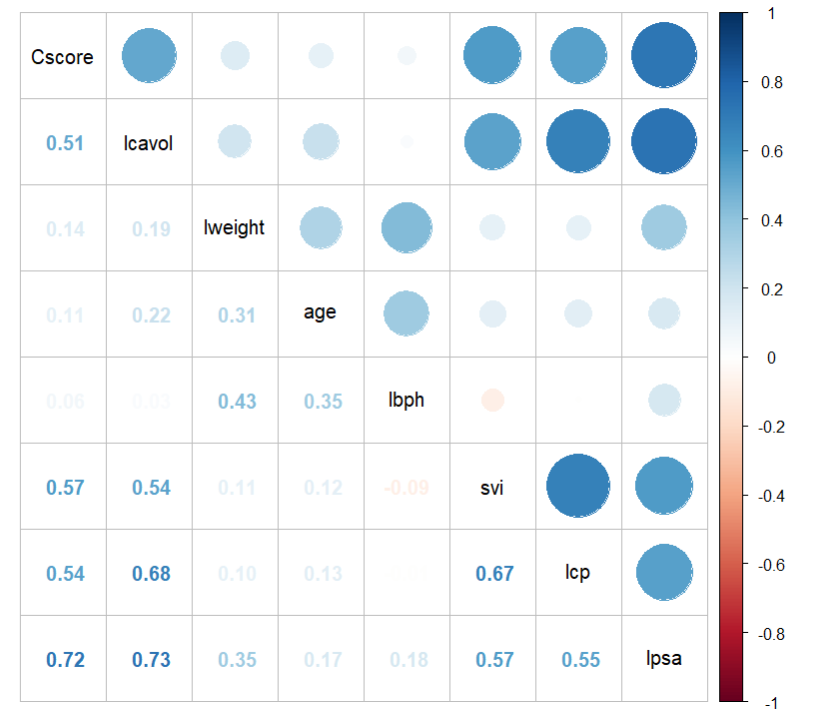
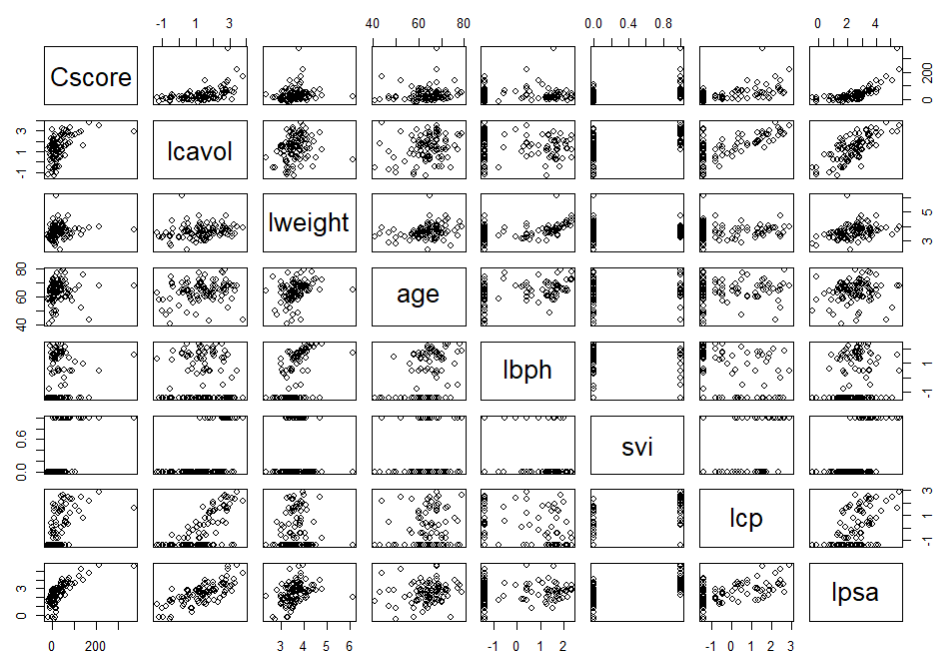


Figure 3. Result of correlation between variables.

1. **Generate your best linear regression model using only linear effects. Are there any indications that assumptions underlying inferences with the model are violated? Evaluate the effect of any influential point, or outlier.**

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Description automatically generated Outliers can have a significant impact on prediction, especially in models that rely on minimizing errors or distances between the predicted values and the actual values. In regression models, residuals should fit a dataset well and fall randomly around zero, following a normal distribution for an unbiased fit. However, residual plots and a quantile-quantile (Q-Q) plot (Figure 5) revealed that the patient at index 96 was an outlier. The influence plot (Figure 4) also confirmed that the patient at index 96 was an influential point with a Cook's Distance larger than 0.5. The presence of an outliers can pull the regression line or decision boundary away from most of the data, which will increase the variance of estimated coefficients and lead to less accurate predictions. Therefore, the decision was made to remove the entire row at index 96 from the dataset. After removing the outlier, both the residual standard error and variance decreased (residual error: 35 -> 24, variance: 2779 -> 1601), indicating a more accurate fit. It is important to note that high leverage points with high residuals can have a significant impact on the slope of the regression line. On the other hand, the point at index 32 is a high leverage point with a low residual. Therefore, it is unlikely to have a significant impact on the coefficient or accuracy of the regression model, and it was decided to keep this point in the analysis.

Figure 4 .Influence Plot

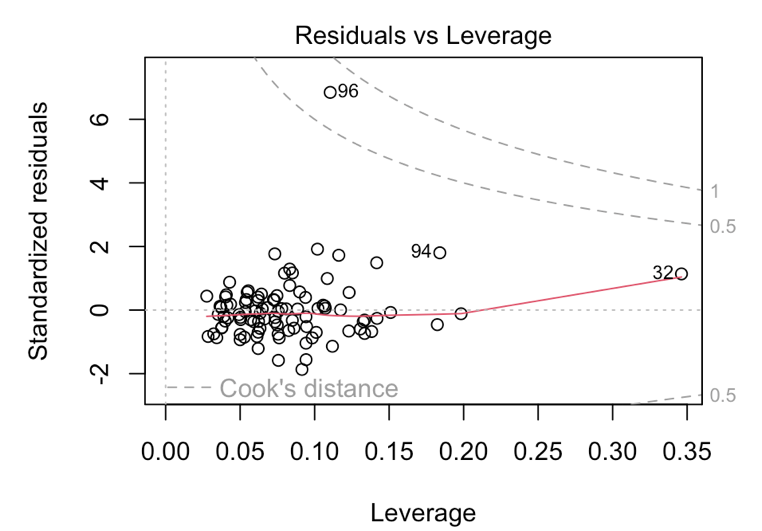
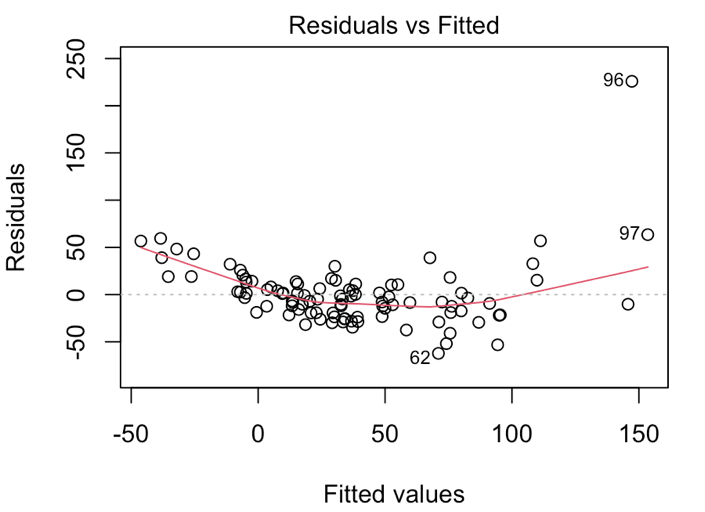
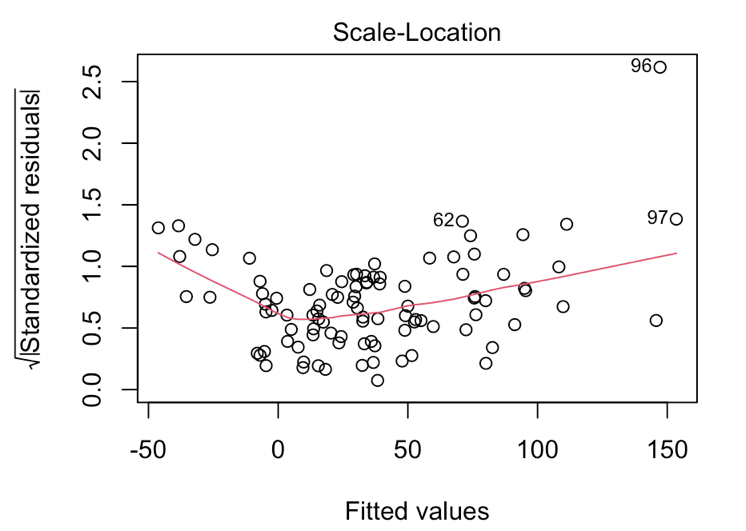
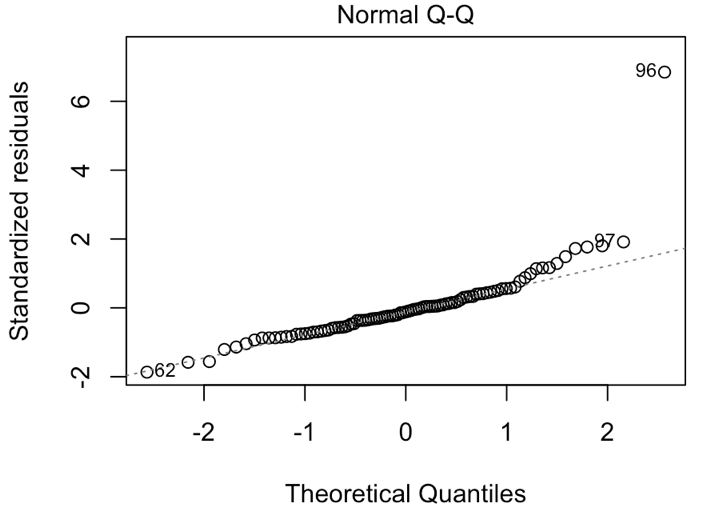
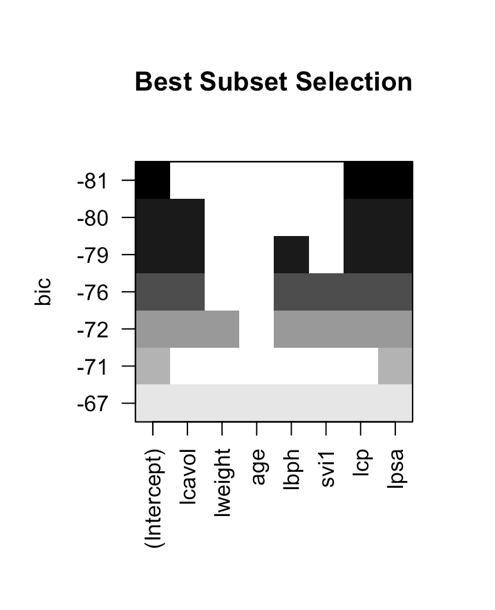
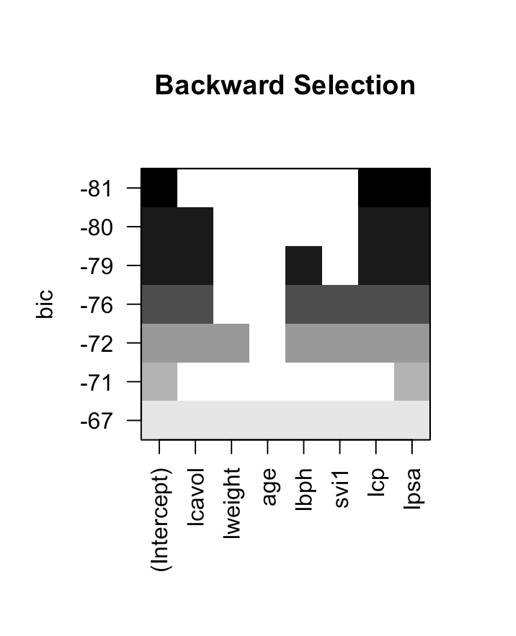
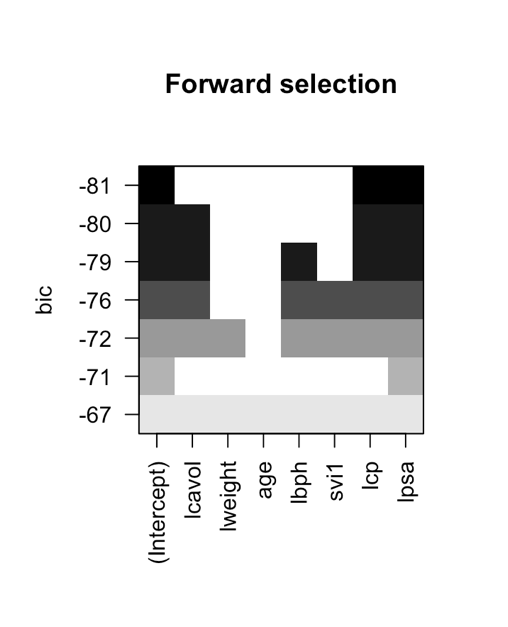
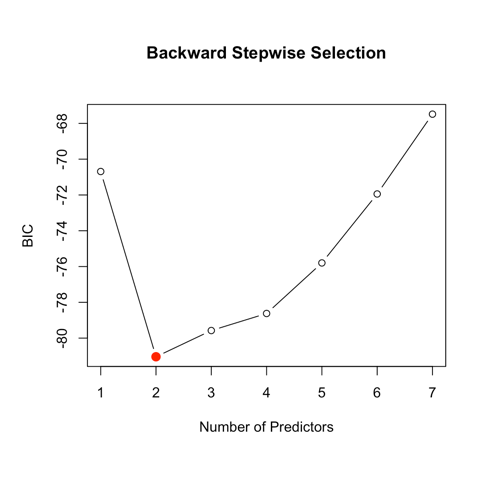
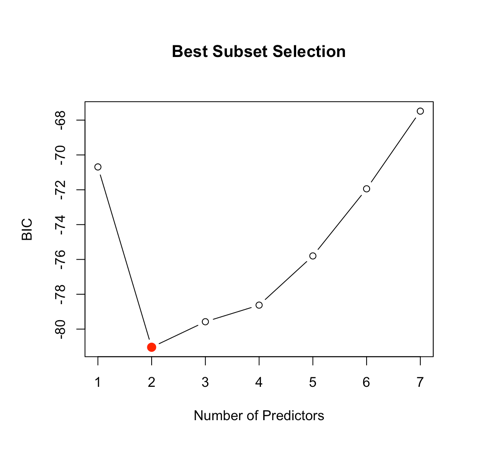
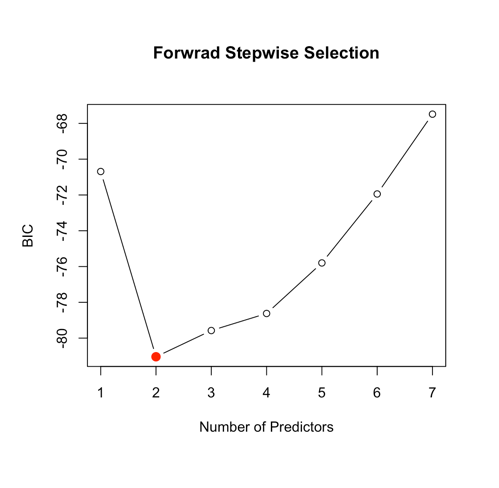


Figure 5. Diagnostic Plot for evaluating the assumptions of linear regression model

The Best Subset Selection, Forward Selection, and Backward Selection techniques are employed to identify the optimal model based on the Bayesian information criterion (BIC) score. The BIC score penalizes the model for its complexity, aligning with the principle of Occam's razor, which states that the simplest explanation is often the best one. Additionally, within the Bayesian probability framework, the BIC provides a higher probability of selecting the true model based on the prior probability of the candidate models that include the true model. As shown in Figure 6, the results of all three techniques indicate that the two variables *lcp* and *lpsa* are the best predictors of Cscore.

Best Subset Selection, Forward Selection, and Backward Selection are all examples of deterministic search algorithms that are relatively fast and straightforward to implement. These methods can provide a good balance between model complexity and predictive accuracy and can help identify a small set of predictors that are highly related to the outcome variable. While cross-validation is a more data-driven approach that involves splitting the data into training and testing sets to evaluate model performance. Cross-validation is a more robust method to estimate the predictive performance of a model because it evaluates the model's performance on a hold-out set of data that was not used to fit the model. This provides a more realistic estimate of the model's performance on new, unseen data, which help estimate the generalizability of a model and identify overfitting.

*Figure 6. Best Subset Selection, Forward Selection and Backward Selection*



As shown in Figure 7, the validation set approach and cross-validation have resulted in different models. The validation set approach selected the model with only one variable, lpsa (Figure 7. Left)., while cross-validation selected the model with two variables, lcp and lpsa (Figure 7. Right). The two functions perform model selection using different approaches. Validation set approach split the data into training and validation sets. The training set is used to fit models of increasing complexity and the validation set is used to evaluate the performance of each model and choose the best one based on the lowest validation error. This approach is prone to overfitting since the model selection is based on a single validation set, and the results may not be reliable. K-fold cross-validation, on the other hand, uses all the data for training and validation and therefore provides a more robust estimate of the test error. In summary, the validation set approach is simpler and faster to implement but can lead to overfitting, while k-fold cross-validation is more reliable but requires more computation time.

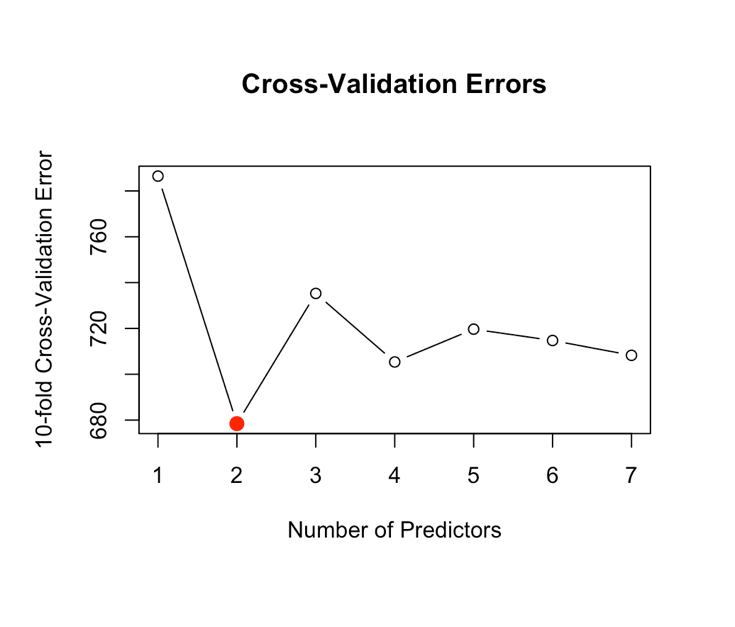
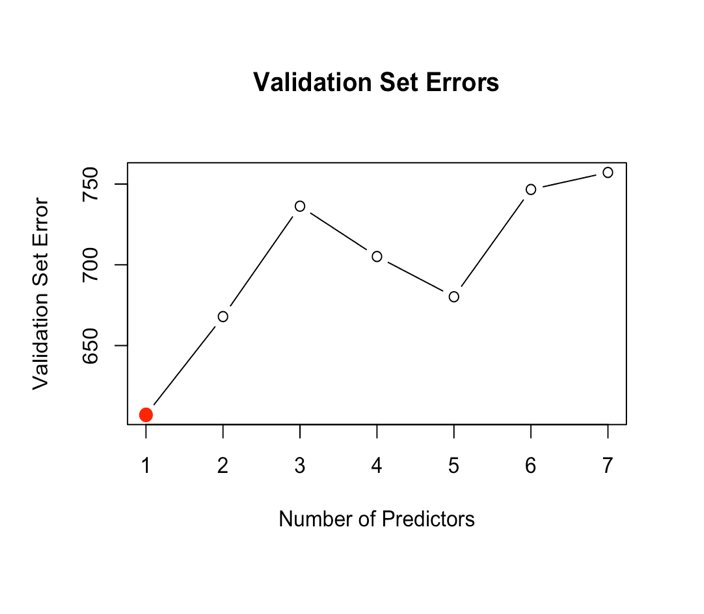


Figure 7. Validation Set vs 10-fold Cross Validation

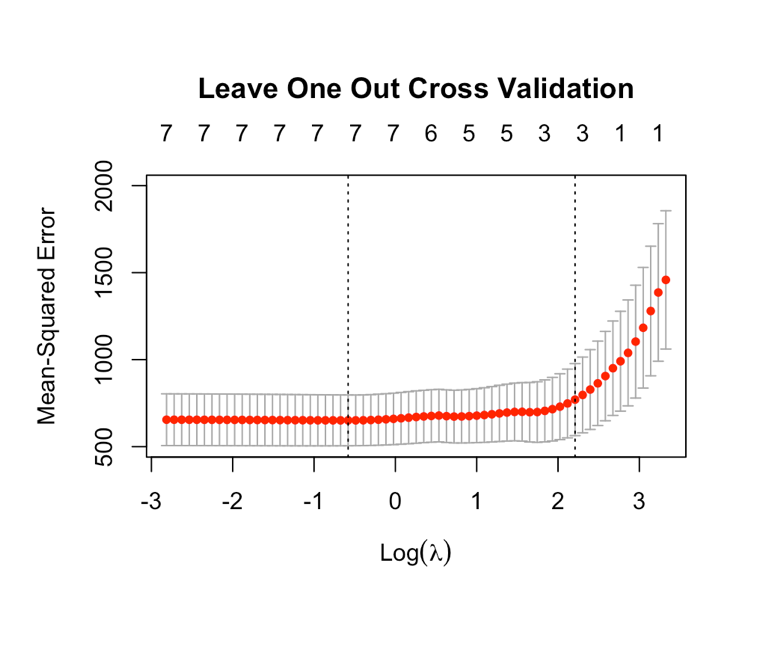
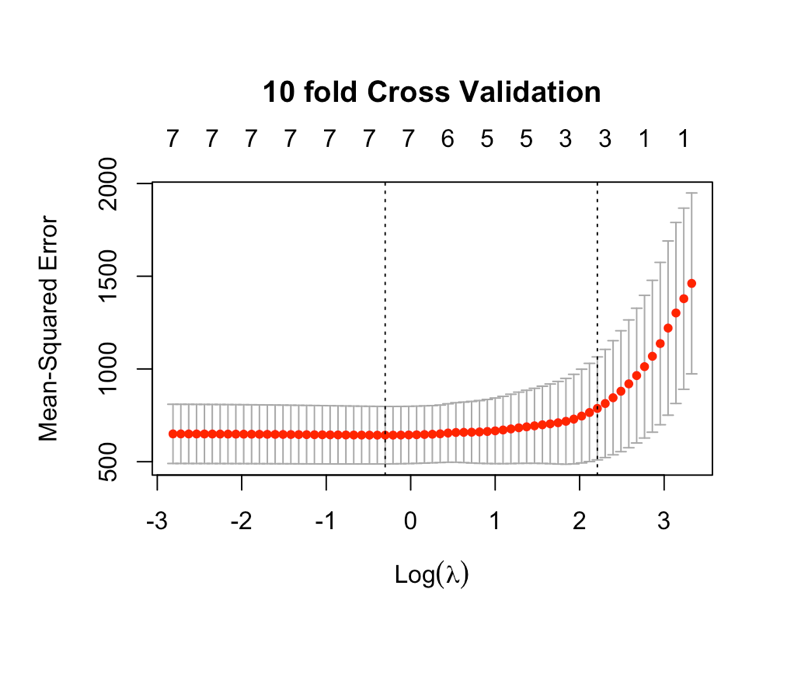
Base I the consistency we are able to obtain among different approach, the final linear model for the selection of variables includes the predictors and , and the formula for this model is . This outcome aligns with our expectations since the severity of cancer is likely to increase with a higher level of invasion or migration of cancer cells into the capsule, which is represented by the lcp variable. The PSA protein, which is produced by both normal and malignant prostate cells, is a useful indicator for prostate cancer detection. Higher PSA levels are often associated with prostate cancer, and this is reflected in the positive coefficient of 21.2 for the lpsa variable in the final model. The coefficient values indicate that a one-unit increase in lcp is associated with an increase of 8.5 in the mean value of the response variable, holding other variables constant, and a one-unit increase in lpsa is associated with an increase of 21.2 in the mean value of the response variable, holding other variables constant.

1. **Make an appropriate LASSO model, with the appropriate link and error function, and evaluate the prediction performance. Do you see evidence that over-learning is an issue?**

Lasso regression is a linear regression technique that incorporates a penalty term to the cost function, which is controlled by the tuning parameter 𝜆. The penalty term imposes a constraint on the sum of the absolute values of the regression coefficients, resulting in the shrinking of coefficients towards zero. This leads to variable selection, whereby some coefficients are set exactly to zero, making lasso particularly useful in high-dimensional datasets where not all features may be important in predicting the outcome variable. By selecting the most important features, lasso regression can reduce overfitting and improve model interpretability. Unlike ridge regression, which shrinks all coefficients towards zero, lasso can eliminate some coefficients, resulting in a sparse model with only a subset of the original predictors.

Overfitting occurs when a model is too complex and captures noise instead of the underlying signal in the data, leading to poor generalization performance on new data. One way to assess overfitting in Lasso is to compare the model performance on the training and testing sets. If the model has learned too much from the training set and does not generalize well to the test set, this may indicate overfitting. The prediction performance of the Lasso model can be evaluated using the test set, and the mean squared error (MSE) can be calculated. If the MSE on the test set is significantly higher than the MSE on the training set, this may indicate overfitting.

Figure 8. (Left) 10-fold cross-validation error (Right) Leave-One-Out-Cross-Validation error as a function of log λ



In our study, we fitted a Lasso model using the glmnet package to predict *Cscore* based on all predictor variables. The model was trained on a randomly selected two-thirds of the data and tested on the remaining one-third. The optimal lambda parameter was chosen via cross-validation, as shown in the Figure 8(Left). We observed that a shrinkage penalty λ of 0.74 resulted in the smallest cross-validation error of 643.20. We calculated the MSE for both the training and test sets using the selected lambda. We found that the MSE on the test set (750.17) was higher than the MSE on the training set (468.57), suggesting that the model may be overfitting to the training data. This indicates that the model may have learned the noise in the training data and may not generalize well to new data. However, the small size of the test set (only 32 observations) makes it difficult to draw definitive conclusions about the model's performance. By varying the size of the training and test groups, we observed that the test and training MSEs differed significantly. Therefore, additional evaluation may be necessary. We decided to perform LOOCV (leave-one-out-cross-validation) to further assess the Lasso model's performance.

LOOCV is a resampling technique that is used to estimate the performance of a statistical model on new data. It works by fitting the model on n-1 data points and testing it on the remaining one. This process is repeated n times, each time leaving out a different data point. By averaging the results over all the n iterations, we obtain an estimate of the model's prediction performance on new data. LOOCV is often used when the sample size is small, as it uses all the data for training and testing and can give a good estimate of the model's performance on new data. In our case, LOOCV was performed with nfolds=96, which means that the model is trained and tested 96 times, leaving out one observation at a time. As shown in Figure 8(Right), a shrinkage penalty of λ=0.55 was selected to minimize the cross-validation error, resulting in a value of 651.01. However, it is worth noting that LOOCV did not result in better performance than using 10-fold cross-validation. The performance on the test set was not as good as expected, with a MSE of 757.24, suggesting that the model may still be overfitting.

1. **Look at the coefficient for “lcavol” in your LASSO model. Does this coefficient correspond to how well it can predict Cscore? Explain your observation.**

The coefficient of *lcavol* in a LASSO model represents its correlation with the target variable, *Cscore*, while accounting for the impact of other predictor variables in the model. A negative coefficient indicates an inverse relationship between *lcavol* and *Cscore*, implying that as *lcavol* increases, the predicted value of *Cscore* decreases. However, the coefficient alone does not provide an accurate measure of *lcavol*'s predictive ability and must be considered in the context of other variables in the model, as well as the model's overall performance in predicting the target variable.

As shown in Figure 9, there are two different LASSO models with distinct coefficient values for *lcavol.* The 10-fold Cross-validation LASSO model's coefficient for *lcavol* is -3.097977, indicating that a one-unit increase in the natural log of *lcavol* corresponds to a -3.097977 unit decrease in *Cscore* while keeping other predictors constant. On the other hand, the LOOCV LASSO model's coefficient for *lcavol* is -3.969890. Therefore, it is not appropriate to draw direct comparisons between these two coefficients or to determine *lcavol*'s predictive capability based solely on the coefficient value.

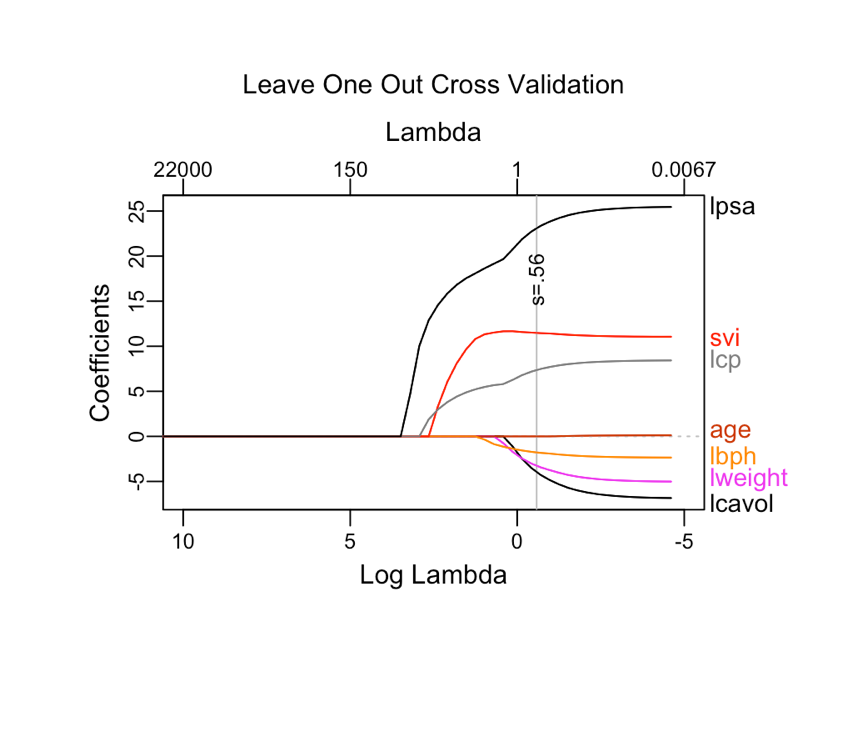
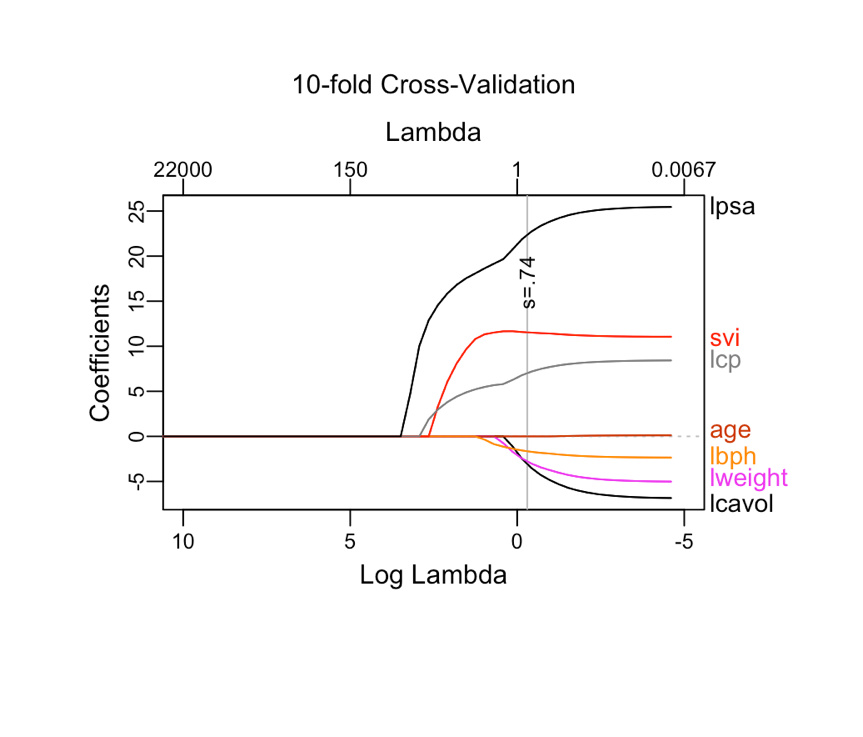


Figure 9. (Left) 10-fold Cross Validation (Right) LOOCV standardized Lasso coefficients as functions of log λ

We conducted a linear regression analysis with "lcavol" as the only predictor variable and "Cscore" as the response variable. The Adjusted R-squared value was 0.3138, indicating that only 31.38% of the variability in "Cscore" can be explained by the variability in "lcavol", while controlling for other predictor variables in the model. As shown in Figure 10, we also generated a scatter plot with "lcavol" on the x-axis and "Cscore" on the y-axis, which included a linear regression line to visualize the overall trend. This plot allowed us to visually evaluate how well "lcavol" predicts "Cscore", and we observed a moderate positive correlation of 0.5665924 between the two variables.

It is important to keep in mind that correlation does not necessarily imply causation, and that there may be other variables not included in the model that can influence the relationship between "lcavol" and "Cscore". In the context of the LASSO regression model, it is possible for the coefficient for "lcavol" to be negative, even if there is a positive correlation between "lcavol" and "Cscore". This can occur because other predictor variables in the model may have a stronger association with "Cscore" or because the relationship between "lcavol" and "Cscore" may be non-linear or conditional on the values of other variables. Therefore, it is essential to evaluate the overall performance of the model and consider the interpretation of coefficients in the context of the other predictor variables.

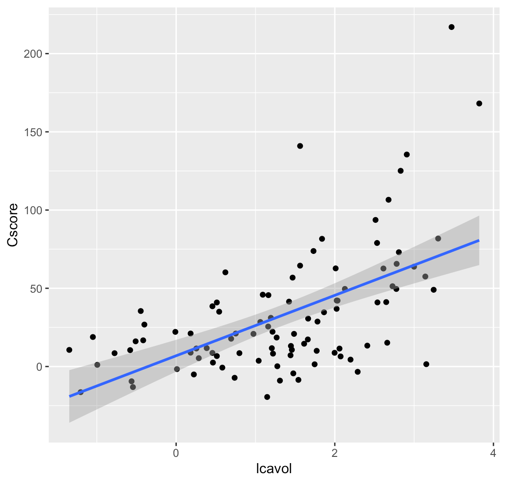


Figure 10. Relationship between lcavol and Cscore

As illustrated in Figure 3, "lcavol" and "lpsa" exhibit high correlation, which suggests that they can be used interchangeably to predict the response variable. However, when dealing with a group of variables that have high correlations, the LASSO may select one variable from the group while reducing the influence of the others. This means that the coefficient of "lcavol" in the LASSO model may not be representative of its true relationship with "Cscore", and it may be necessary to consider other models or techniques to better understand the relationship between the variables.

1. **Fit your best model with *appropriate* non-linear effects. Report a comparison of performance to LASSO and your model reported under question 2. Explain what you find and indicate relevant issues or limitations of your analysis.**

To fit the best model with appropriate non-linear effects, we utilized the gam function from the gam package to fit generalized additive models that incorporate non-linear effects of variables using smooth functions. To evaluate the performance of this model, we compared it to the LASSO and the best linear model using the mean squared error (MSE) on the test set. It is important to note that adding more terms or interactions can increase the complexity of the model and risk overfitting, so careful evaluation and consideration of the trade-off between model complexity and performance is necessary. In other words, simpler models are preferred within the marginal worse performance.

To determine the best subset of predictors, we split the data into a training and test set using a 2/3 split ratio and performed forward stepwise selection. We found that the best model includes the predictors *lcavol*, *lbph*, *lcp*, and *lpsa*. We then fit a generalized additive model using smoothing spline functions with 4 degrees of freedom to identify the non-linear effects of each predictor. The performance of the GAM is evaluated by computing the MSE (379.97) on the test data. We found that only *lpsa* had a significant non-linear effect and simplified the model by removing the predictor *lbph* and the smoothed term of predictor *lcp* and *lcavol*, respectively. For further simplification, we reduce the degrees of freedom and then the test MSE (332.66) was computed for the simplified models. An ANOVA test was performed to compare the performance of the models, simplification justified as expected.

The reason why we choose to use smoothing splines as a starting point of fitting GAM model is that they do not rely on any pre-specified functional form of the non-linear relationship between variables. This means that they can capture complex and intricate non-linear relationships, without the researcher having to make any assumptions about the shape of the relationship. Additionally, smoothing splines are a type of flexible regression model that can be adjusted to fit different degrees of non-linearity. This can be useful in situations where the degree of non-linearity is unknown or varies across the range of the predictor variable. Finally, smoothing splines have been shown to have good performance characteristics in terms of minimizing mean squared error and avoiding overfitting, especially when compared to other non-linear regression methods such as stepwise regression.

A natural spline function was then applied to *lpsa*, and we found that the natural spline function had a lower MSE (310.34) on the test set than the GAM model, indicating better performance. We also identified that removing *lcavol* resulted in a model with a lower MSE (267.09). Further simplification of the model by reducing the degrees of freedom resulted in marginally worse test MSE, which was justified by an ANOVA test. However, further simplification by removing *lcp* led to a significant increase in the MSE (324.33, p-value 0.0009). Finally, a polynomial regression model was fitted with linear predictors *lcp* and a quadratic term of *lpsa*. The test MSE for this model was significantly lower than all other models(257.29), and it maintained high simplicity. Cubic term of *lpsa* was also fitted and display even lower MSE(246.70). However, with p-value of 0.1838 on ANOVA, simpler model was favoured.

Comparing the performance of the best non-linear model to LASSO (MSE:750.17) and the best linear model reported in question 2(MSE: 684.49), we found that the non-linear model had a significantly lower MSE. Therefore, a polynomial regression model with linear predictors *lcp* and a quadratic term of *lpsa* was identified as the best model for predicting the prostate cancer Cscore, which explains 80.56% of the variability in Cscore. The formula for this model is .

One limitation of this analysis is that the sample size is relatively small, which may limit the generalizability of the findings. Additionally, the models are limited by the available predictors and potential unmeasured confounding variables that may affect the relationship between the predictors and the response variable. It is important to note that we only used the mean squared error as a measure of predictive performance, and non-linear models may have potential interpretability issues since the effects of the variables are not easily summarized by a single coefficient. Further research and data collection may be needed to improve the accuracy of the models.