PSTAT 126 Project Summary

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Introduction

In this project, we used multivariate linear regression to determine potential predictors of maximum heart rate. We used an OLS method to fit a model, and ultimately chose seven predictors: slope, age, cp, exang, target, trestbps and restecg. Specifically, an OLS method aims to minimize the sum of squared residuals. Finally, we used logistic regression to examine whether patients have heart disease based on certain predictors selected by LASSO.

Data Description

This data set dates from 1988 and consists of four databases: Cleveland, Hungary, Switzerland, and Long Beach V. It contains information about patients and risks for heart disease as well as whether or not they eventually end up developing heart disease. The data set originally contains 1025 observations. For the sake of convenience of the scatter plot, we randomly chose 500 from the original dataset.

The age variable is a numeric variable that lists the age of the patient in years. The sex variable is a binary variable that states the sex of the patient, with 0 representing female and 1 representing male. The cp variable is a categorical variable that classifies the type of chest pain the patient is experiencing; 0 represents typical angina, 1 represents atypical angina, 2 represents non-anginal pain, and 3 represents asymptomatic. The trestbps variable is a numeric variable that states the resting blood pressure in mmHg.

The chol variable is a numeric data which give an overview of person's cholesterol levels. The fbs variable is a binary data with 1 means the fast blood sugar is greater than 120mg/dl, and 0 means the fast blood sugar is not. The restecg variable is a nominal data with scale of 0 (normal), 1 (abnormality in ST-T waves), and 2 (show left ventricular hypertrophy), which record the resting electrocardiogram results. The thalach variable is a numeric data which records the maximum heart rate achieved.

The variable exang indicates that if the patient have exercise induced angina (chest pain), which is shown with 0 indicating "no" and 1 indicating "yes". The variable oldpeak is a float type data records the ST depression, which is a measure of abnormality of an electrocardiogram, and the measurement is in unit depression. The variable slope is the slope of the peak exercise ST segment, which is an electrocardiography read out indicating quality of blood flow to the heart, and the data are in nominal type including 0 (upsloping), 1 (flat) and 2 (downsloping). Finally, the variable target is a binary data showing whether the patient is having heart disease (1 = having heart disease and 0 = normal).

Summary Statistics and Graphs

Table 1: Data summary

Name	heart_disease_data
Number of rows	500
Number of columns	14
Column type frequency:	
factor	7
numeric	7
Group variables	None

Variable type: factor

skim_variable	n_missing	complete_rate	ordered	n_unique	top_counts
sex	0	1	FALSE	2	1: 355, 0: 145
ср	0	1	FALSE	4	0: 223, 2: 159, 1: 86, 3: 32
fbs	0	1	FALSE	2	0: 418, 1: 82
restecg	0	1	FALSE	3	0: 263, 1: 233, 2: 4
exang	0	1	FALSE	2	0: 330, 1: 170
slope	0	1	FALSE	3	2: 238, 1: 227, 0: 35
target	0	1	FALSE	2	1: 266, 0: 234

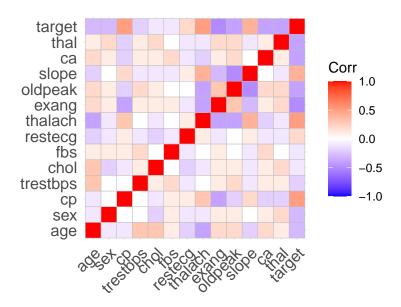
Variable type: numeric

skim_variable	n_missing	$complete_rate$	mean	sd	p0	p25	p50	p75	p100	hist
age	0	1	54.02	9.22	29	47	55.0	60.25	77.0	
trestbps	0	1	132.46	18.06	94	120	130.0	140.00	200.0	
chol	0	1	248.13	55.44	126	211	243.0	282.00	564.0	
thalach	0	1	150.19	23.31	71	137	152.0	168.00	202.0	
oldpeak	0	1	1.03	1.17	0	0	0.8	1.60	6.2	
ca	0	1	0.76	1.07	0	0	0.0	1.00	4.0	
thal	0	1	2.29	0.63	0	2	2.0	3.00	3.0	

Correlation

The following correlation matrix represent the relationship between different variables and the interaction between each other.

Each cell represents a correlation coefficient, the red color, which has a coefficient of 1, represents the strong positive correlation between two variables; whereas, the white color with coefficient of 0 indicates little (no) correlation between two variables, and the purple color with coefficient of -1 indicates a strong negative correlation.



The correlation matrix of numeric data shows that there isn't a strong positive correlation between any two variables, and between maximum heart rate achieved and target; a relatively strong negative correlation between ST slope and oldpeak; and little correlation between sex and chest pain type, cholesterol and fasting blood sugar.

Selecting Model

We decided to use an OLS model because our predictors were not highly correlated and we have a relatively large sample size compared to the number of predictors, so variable selection is not as big of an issue.

We used a forward step-wise model, meaning we assumed no relationship between the variables, and then added predictors one by one until no other significant predictors were left.

Our criteria for choosing the variables is based on the AIC, meaning that starting with an intercept-only model, we add the variable with the lowest AIC value at a time, then re-examine the new model to find the next variable until no variables significant.

By going through forward step-wise selection, we chose 7 predictors for the model, which are slope, age, cp, exang, target, trestbps and restecg.

```
##
##
  lm(formula = thalach ~ slope + age + cp + exang + target + trestbps +
##
       restecg, data = train_data)
##
## Residuals:
##
       Min
                1Q
                    Median
                                 3Q
                                        Max
##
  -55.480
           -9.282
                     1.679
                            12.213
                                     49.131
##
```

```
## Coefficients:
##
                Estimate Std. Error t value Pr(>|t|)
## (Intercept) 169.35644
                            8.77777
                                     19.294 < 2e-16 ***
                            3.57973
                                     -0.325 0.745187
## slope1
                -1.16422
## slope2
                11.27038
                            3.70469
                                      3.042 0.002508 **
                -0.78128
                            0.10136
                                     -7.708 1.08e-13 ***
## age
## cp1
                10.55720
                            2.84049
                                      3.717 0.000232 ***
## cp2
                 5.17212
                            2.45548
                                      2.106 0.035815 *
## cp3
                15.10773
                            3.79034
                                      3.986 8.03e-05 ***
## exang1
                -8.35457
                            2.22378
                                     -3.757 0.000198 ***
## target1
                 6.43474
                            2.32472
                                      2.768 0.005911 **
## trestbps
                 0.11511
                            0.05034
                                      2.286 0.022769
                -3.17946
                            1.79352
                                     -1.773 0.077055
## restecg1
## restecg2
               -12.76272
                           10.06948
                                    -1.267 0.205749
## ---
                   0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
## Residual standard error: 17.15 on 388 degrees of freedom
## Multiple R-squared: 0.4909, Adjusted R-squared: 0.4765
## F-statistic: 34.02 on 11 and 388 DF, p-value: < 2.2e-16
```

Finding R^2 of Predicted Values

About 49% of the variance in maximum heart rate in the test data is accounted for by the slope of the peak exercise ST segment, age, chest pain type, exercise-induced angina, heart disease, resting blood pressure, and resting ecg.

```
## [1] "R-square value on test data: 0.493739305422597"
```

Checking Influential Points

To check outliers, leverage points, and influence points, we created graphs of the residuals, the diagonal values of the hat matrix, and Cook's distance. In each graph, we selected the largest value in red to visualize it on each plot.

The residuals appear randomly scattered with equal variance and no obvious pattern, and based on the Q-Q plot, are approximately normal, indicating that the model is a good fit.

We then refit the model without the influential point:

```
##
## Call:
##
  lm(formula = thalach ~ target + age + slope + exang + cp + trestbps +
##
       restecg, data = train_data[-unusual_idx, ])
##
## Residuals:
##
                                3Q
       Min
                1Q
                   Median
                                        Max
  -52.044 -9.493
                     1.371
                           11.650
##
## Coefficients:
##
                Estimate Std. Error t value Pr(>|t|)
## (Intercept) 170.20376
                            8.54789 19.912 < 2e-16 ***
```

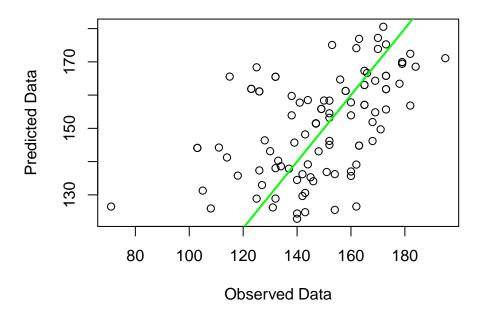


Figure 1: Plotting Predictions for OLS Model

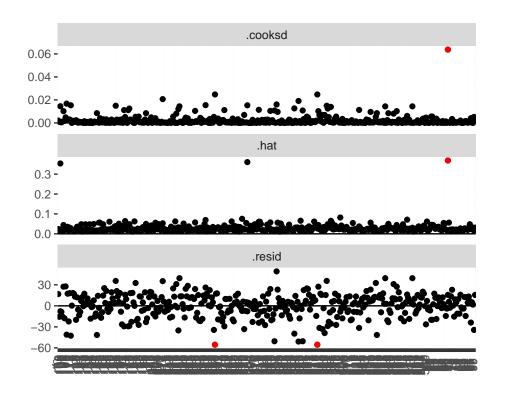


Figure 2: Influential Points Analysis

Normal Q-Q Plot

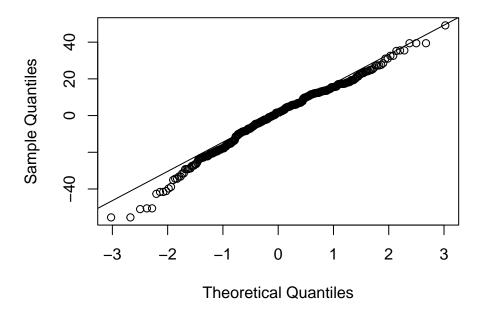


Figure 3: Q-Q Plot of Residuals

```
## target1
                 5.82986
                             2.26682
                                       2.572 0.010490 *
                                      -7.425 7.28e-13 ***
                -0.73603
                             0.09913
## age
## slope1
                -0.80053
                             3.48606
                                      -0.230 0.818496
## slope2
                11.07447
                             3.60713
                                       3.070 0.002291 **
## exang1
                -9.70972
                             2.18320
                                      -4.447 1.14e-05 ***
## cp1
                 9.76435
                             2.77038
                                       3.525 0.000475 ***
## cp2
                 4.29509
                             2.39756
                                       1.791 0.074005 .
## cp3
                14.44117
                             3.69288
                                       3.911 0.000109 ***
## trestbps
                 0.09797
                             0.04914
                                       1.993 0.046916 *
                                      -1.260 0.208537
## restecg1
                -2.21404
                             1.75759
                                      -1.354 0.176448
## restecg2
               -13.27747
                             9.80428
##
## Signif. codes:
                     '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## Residual standard error: 16.7 on 386 degrees of freedom
## Multiple R-squared: 0.4911, Adjusted R-squared: 0.4766
## F-statistic: 33.86 on 11 and 386 DF, p-value: < 2.2e-16
```

From the plot and the best fit lines (Fig. 4), the lines only change a little for the model with or without the influential points, so we shouldn't be concerned about those point affecting the results.

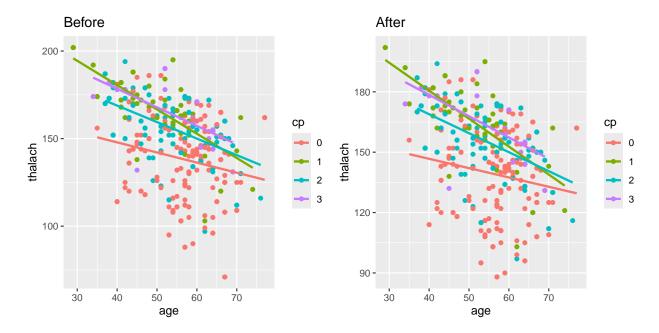


Figure 4: With/Without Influential Points

Innovation

Logistic regression is used to predict the outcome of a binary variable, for example 0 or 1. Suppose the probability of 1 occurs is p and the probability of 1 does not occur is 1-p, then the ratio of 1 occurring vs. 1 not-occurring is simply $\frac{1}{1-p}$.

Since we then need a linear regression to estimate the coefficient, we then apply log map the probability to real number for linear regression, that is

$$Logit(p) = log(\frac{p}{1-p})$$

Apply logit(p) as the new response and fit a linear model with the predictor variables.

$$Logit(p) = \beta_0 + \beta_1 x_1 + \beta_2 x_2 + \dots + \beta_n x_n$$

Then we inverse the logit to get p, namely,

$$p = \frac{1}{1+e^{-(\beta_0+\beta_1x_1+\beta_2x_2+\cdots+\beta_nx_n)}}$$

Technical Conditions

- 1. The response variable is binary. This fits our model that the response is the target variable, which indicates whether the patient have heart disease.
- 2. The model should have minimal multicollinearity. The LASSO regression is removing predictors that are having high correlations.
- 3. Sufficient sample size and independent observation. Our dataset is large and the patients conditions are independent.

- 4. There shouldn't be prefect separation. We don't have some predictors that can precisely predict the outcome.
- 5. There shouldn't be highly influential outliners. For this one we didn't particularly remove the outliners.

Method Justification

To explore further models, we decided to use logistic regression since we were interested in seeing how the variables can be used to predict heart disease, or the target variable. Since heart disease is a binary variable, OLS, ridge, and LASSO cannot be used since the response variable in these cases must be continuous.

Fitting Logistic Model

First, we used LASSO to determine which variables to include in the model. Based on the results from the LASSO regression, we fit a logistic model using the non-zero coefficients.

```
##
## Call:
##
  glm(formula = target ~ age + sex + cp + trestbps + fbs + restecg +
       thalach + exang + slope + ca + thal, family = binomial, data = train data)
##
## Coefficients:
##
                Estimate Std. Error z value Pr(>|z|)
## (Intercept)
               3.215911
                           2.691304
                                      1.195
                                             0.23212
               -0.013983
                           0.023098
                                     -0.605 0.54493
## age
               -2.011873
                           0.445663
                                     -4.514 6.35e-06
## sex1
## cp1
                0.838787
                           0.482932
                                      1.737
                                             0.08241
## cp2
                2.817552
                           0.506933
                                      5.558 2.73e-08 ***
                3.350009
                           0.717658
                                      4.668 3.04e-06 ***
## cp3
## trestbps
               -0.017784
                           0.009636
                                     -1.846 0.06496 .
                                     -1.098
## fbs1
               -0.539926
                           0.491566
                                             0.27204
                0.540507
                           0.351903
                                      1.536
                                             0.12455
## restecg1
               -0.380370
## restecg2
                           2.182778
                                     -0.174
                                             0.86166
## thalach
                0.019662
                           0.011372
                                      1.729
                                             0.08381
## exang1
               -0.853594
                           0.419316
                                     -2.036
                                             0.04178
                0.173169
                           0.746474
                                      0.232
## slope1
                                            0.81655
## slope2
                2.248040
                           0.765454
                                      2.937 0.00332 **
                                     -5.769 7.95e-09 ***
## ca
               -1.116049
                           0.193442
## thal
               -1.307612
                           0.284819
                                     -4.591 4.41e-06 ***
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
## (Dispersion parameter for binomial family taken to be 1)
##
       Null deviance: 552.83 on 399
                                      degrees of freedom
## Residual deviance: 235.42 on 384
                                      degrees of freedom
## AIC: 267.42
##
## Number of Fisher Scoring iterations: 6
```

ROC Curve

A tool to visualize logistic regression is a ROC (Receiver Operating Characteristic) curve, in which the true positive rate (sensitivity) is plotted against the false positive rate (1 - specificity) at different threshold values. The closer the ROC curve is to the top-left corner, the better the model is at distinguishing between positive and negative classes. A model with no discriminative power will have an ROC curve along the diagonal line, essentially equivalent to random guessing. As can be seen in Fig. 5, the ROC curve is relatively close to the top-left corner, indicating that the model has high sensitivity and high specificity.

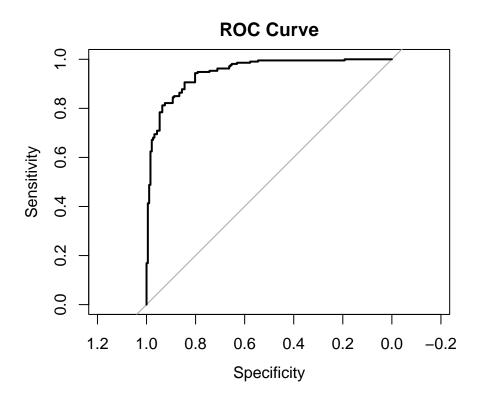


Figure 5: ROC Curve for logisitic model

AUC

The AUC, or area under the curve, is a measure of how well the model is able to discriminate between positive and negative cases. The AUC ranges from 0 to 1, with the following interpretations:

- < 0.5: No discriminative power (equivalent to random guessing).
- 0.5 0.7: Poor discrimination.
- 0.7 0.8: Acceptable discrimination.
- 0.8 0.9: Excellent discrimination.
- > 0.9: Outstanding discrimination.

[1] "AUC: 0.947176822073259"

Since the AUC is above 0.9, we can conclude that our model is highly accurate at distinguishing between cases.

Optimal Threshold Value

Based on the ROC curve, we can determine the optimal threshold value, which is the value that maximizes the distance from the diagonal line (representing random chance) or maximizes the area under the ROC curve (AUC). If the predicted probability is less than the threshold, then the response can be classified as 0, or no heart disease. If the predicted probability is greater than threshold, then the response can be classified as 1, or having heart disease.

[1] "Optimal threshold value: 0.524617014252765"

Conclusion

Overall, our analysis revealed some interesting results. We determined that heart disease, age, quality of blood flow to the heart, exercise induced angina, chest pain type, resting blood pressure, and resting electrocardiogram results are major factors when examining maximum heart rate. However, our data set does not account for other important factors such as family history, diet, environment, and mental health, which all play an important role that should be evaluated in further studies.

Studying maximum heart rate is crucial as it is a key component in cardiovascular health. Maximum heart rate can be used to determine a recommended level of exercise intensity and prevent individuals from overexertion and reduces the risk of heart-related incidents during exercise. Additionally, doctors often use maximum heart rate to monitor cardiovascular health and adjust medication dosages.

We were particularly fascinated by our logistic model, which was fairly accurate in determining whether or not patients had heart disease. However, logistic models are limited by their sensitivity to outliers and their tendency to overfit the data. Going forward, we would also like to examine how other factors can be used to predict heart disease in patients.

In conclusion, while our results yielded valuable insights into studying maximum heart rate, further research is needed to better examine its role in heart function. In the future, we are excited to examine new methods of regression to study maximum heart rate.