



اونيورسيتي مليسىيا فهاغ السلطان عبد الله
UNIVERSITI MALAYSIA PAHANG
AL-SULTAN ABDULLAH

PROJECT STATISTICAL MODELLING & SIMULATION (BSO3443)

NAME		STUDENT ID
1	TAN CHEK CHENG	SD21031
2	WONG ZI MING	SD21037
3	DAVID LAU KING LUEN	SD21044
4	TEAN JIN HE	SD21063
LECTURER	DR. NORIANTI MUHAMMAD	
DATE RELEASE	17 OCTOBER 2023 (TUESDAY)	
DURATION	17 OCTOBER 2023 – 22 JANUARY 2024	
SESSION/SEMESTER	SESSION 2023/2024 SEMESTER I	
METHOD OF SUBMISSION	Upload R Code (rfile) and Output as report (pdf) in KALAM	

Marks Distribution			
Course Outcomes	Program Outcomes	Domain	Marks
CLO2 Formulate statistical models for various problems in science, engineering, and industry. (C5, PLO2)	Critical Thinking & Scientific Approach	CTPS	/ 10 (2%)
CLO3 Manipulate statistical modelling theory and methodology in solving various applications using appropriate statistical software. (P4, PLO3)	Technical Skills	Psychomotor	/ 15 (3%)
CLO4: Demonstrate good interest and initiative for exploring issues in statistical modelling analysis for a given task (A3, PLO7)	Lifelong Learning	Affective	/15 (2%)
CLO5 Plan a business strategy by generating new ideas and innovation in the application of statistical modelling and simulation. (A3, PLO8)	Entrepreneur Skills	Affective	/ 10 (3%)
Total			/ 50

BACKGROUND

Consider you are a Data Analyst at a certain department of organisation or company which objective to find and create a new or best solution to your organisation. Your organisation might consist of government sector, private company, industry, agency etc. Based on the data given, there are three main objectives you should focus on, which are:

1. To understand the data and variables by using any knowledge, method, and tools that you have learned.
2. To analyse the data and develop the best model which give the best solution to your organisation. Item you should do for the analysis;
 - a. Investigate the basic model obtained from the data set.
 - b. Develop the best model as your solution based on this course.

- c. Briefly explain your final model which consider as the best solution.
3. To plan the simple business of this task or assignment by considering developing an appropriate mobile apps/interactive dashboard/software or other appropriate device for the solution you obtained to be considered as your product.

Important Notes:

1. Please use the rubrics attached to guide your final report in order to get maximum marks.

APPENDIX: RUBRIC GUIDELINE FOR CLO2 (PLO2)

CRITERIA	LEVEL OF ACHIEVEMENT					WEIGHTAGE	SCORE (%)
	Inadequate 1	Emerging 2	Developing 3	Good 4	Excellent 5		
Identifying main problem and stating the subsidiary aspect of the problem	Unable to identifies the main problem and fail to state the subsidiary aspect of the problem	identifies little the main problem and fail to state the subsidiary aspect of the problem	Identifies the main problem without stating the subsidiary aspect of the problem	Identifies the main problem and considering little stating the subsidiary aspect of the problem	Identifies and clearly states both the main and subsidiary aspect of the problem	0.1	
Collecting and analysing information	Collects inadequate viable information unable to perform analysis	Collects little viable information unable to perform analysis	Collects adequate information and performs analysis based on two sources	Collects adequate information and performs analysis based on more sources	Collects a bundance of information and performs analysis based on multiple resources and justify.	0.1	
Identifying potential solutions	Identifies a single solution yet fail to present reasoning	Identifies two solutions yet fail to present reasoning	Identifies a few simple solutions and simple reasoning for the suggested solutions	Identifies more simple solutions and simple reasoning for the suggested solutions	Identifies and explains - accurately and thoroughly- multiple solutions and perspectives	0.1	
Selecting the most appropriate solution based on appropriate data or theory and provide alternative approach	Do not select any solutions that does not meet the required specifications.	Selects a solution that does not meet the required specifications.	Selects a reasonable solution but does not justify the solution. No discussion of alternate approaches included	Selects a reasonable solution and justify the solution. No discussion of alternate approaches included	Selects and articulates a solution based on appropriate data and discuss alternative approaches.	0.1	
						TOTAL	/2

APPENDIX: RUBRIC GUIDELINE FOR CLO3 (PLO3)

CRITERIA	LEVEL OF ACHIEVEMENT					WEIGHTAGE	SCORE (%)
	Inadequate 1	Emerging 2	Developing 3	Good 4	Excellent 5		
Theory/ Knowledge	Very little knowledge provided, or information is incorrect	Some knowledge or information provided but missing all major points	Some knowledge or information provided but still missing some major points	Good knowledge observed, missing some minor points	Excellent knowledge observed; provides all necessary background principles	0.2	
Measurement/ Techniques/ Data Validation	Inappropriate measurement techniques are demonstrated	Partly correct measurement techniques are demonstrated, with partly valid data	Correct measurement techniques are demonstrated, with partly valid data	Good measurement techniques are demonstrated, with valid but not accurate data	Competent measurement techniques are demonstrated, with valid and accurate data	0.2	
Results	Lack of results / zero readability of the result. Poor originality, taking credits of others work	Partly complete result	Result presented but at low readability / some result presented. Reader has to guess some of the missing information. Less originality, copy paste here and then	Clear, neat presentation. All required results are presented. Readability. Complete with labels, title, axes, etc.	Very Clear, neat presentation. All required results are presented. High readability. Complete with labels, title, axes, etc.	0.2	
						TOTAL	/3

APPENDIX: RUBRIC GUIDELINE FOR CLO4 (PLO7)

Criteria	LEVEL OF ACHIEVEMENT					WEIGHTAGE	SCORE (%)
	Very Weak 1	Weak 2	Fair 3	Good 4	Very Good 5		
Interest	No interest in exploring issues for a given task	Demonstrate limited interest in exploring issues for a given task	Demonstrate sufficient interest in exploring issues for a given task	Demonstrate good interest for exploring issues for a given task	Demonstrate excellent interest in exploring issues for a given task	0.2	
Optimisation	Not able to retrieve information	Able to retrieve Information from minimal references	Able to retrieve information from sufficient reference	Able to retrieve information from many references	Able to retrieve Information from Maximum references	0.2	
						TOTAL	/2

APPENDIX: RUBRIC GUIDELINE FOR CLO5 (PLO8)

Criteria	LEVEL OF ACHIEVEMENT					WEIGHTAGE	SCORE (%)
	Very Weak 1	Weak 2	Fair 3	Good 4	Very Good 5		
Vision	No vision to solve problem.	Minimal vision to solve problem.	Satisfactory vision to solve problem.	Good vision to solve problem.	Excellent vision to solve problem.	0.2	
Passionate	Dislike to organise an entrepreneurial activity.	Minimal liking to organise an entrepreneurial activity.	Satisfactory liking and enjoys organising an entrepreneurial activity.	Likes and enjoys organising an entrepreneurial activity.	Passionate to organise an entrepreneurial activity.	0.2	
Entrepreneurial Opportunity	No entrepreneurial idea for value adding/solving customer needs.	Has unclear entrepreneurial idea for value adding/ solving customer needs and is not relevant to customer needs.	Business idea is clear but does not fulfil the realistic customer needs.	Business idea is clear and fulfils the customer needs.	Able to mobilise the idea to become opportunity according to the business strategy and fulfil the customer needs.	0.2	
						TOTAL	/3

TABLE OF CONTENT

	TABLE OF CONTENT	i
1.0	INTRODUCTION	1
1.1	Case Study	1
1.2	Objectives	1
2.0	DATA DESCRIPTION	2
3.0	DATA ANALYSIS	4
3.1	Attributes Distributions	4
3.2	Exploratory Analysis and Visualization	10
4.0	DATA MODELLING	13
5.0	END PRODUCT	24
5.1	Introduction	24
5.2	Smoker Status Prediction Interface	24
6.0	CONCLUSION	28
7.0	REFERENCE	29
8.0	APPENDIX	30

1.0 INTRODUCTION

1.1 Case Study

Smoking is the process of burning a material—typically tobacco, cannabis, or opium—and then inhaling the smoke to become high. Depending on the material and the amount smoked, smoking can have either energizing or calming effects on the body. In addition to being extremely addictive, smoking is bad for your health since it increases your risk of developing heart disease, stroke, lung cancer, and chronic obstructive pulmonary disease (COPD). Many negative effects of smoking on health have been demonstrated. It has been discovered that smoking shortens the life expectancy of smokers overall and damages almost all of the body's organs. As of 2018, smoking is still a major contributor to avoidable morbidity and mortality worldwide, posing a persistent threat to global health. The World Health Organization analysis estimates that by 2030, smoking-related fatalities will account for 10 million deaths.

Evidence-based treatments to assist smokers in quitting have been proposed and promoted. Less than one-third of the participants were able to maintain sobriety, though. Many doctors felt that smoking cessation therapy was time-consuming and useless, thus they did not routinely offer it in their daily practices. The degree of nicotine dependence, the amount of carbon monoxide (CO) exhaled, the number of cigarettes smoked daily, the age at which smoking began, the history of failed quit attempts, marital status, emotional distress, temperament and impulsivity scores, and the desire to give up the habit are some of the variables that have been proposed to address this issue and determine which smokers would be more likely to quit successfully. However, when these characteristics are used individually for prediction, the results can be inconsistent and challenging for physicians and patients to comprehend and apply. Offering a prediction model could be a useful approach to comprehending each smoker's likelihood of stopping. In recent years, machine learning techniques have been used to construct health outcome prediction models.

1.2 Objectives

1. To develop a full model and do the comparison with a reduced model in order to choose the best prediction model used.
2. To predict the smoking status by considering various health-related variables
3. To visualize the prediction result.

2.0 DATA DESCRIPTION

Dataset: <https://www.kaggle.com/datasets/gauravduttakiit/smoker-status-prediction/data>

The data has been provided with the smoker status prediction and contains the following fields:

Attribute	Data Type	Description
age	Integer	Age of the individual in years
height.cm.	Integer	Height of the individual in centimeters
weight.kg.	Integer	Weight of the individual in kilograms
waist.cm.	Numeric	Waist circumference length in centimeters
eyesight.left.	Numeric	Eyesight in the left eye
eyesight.right.	Numeric	Eyesight in the right eye
hearing.left.	Integer	Hearing status in the left ear (1: normal, 2: impaired)
hearing.right.	Integer	Hearing status in the right ear (1: normal, 2: impaired)
systolic	Integer	Systolic blood pressure
relaxation	Integer	Diastolic blood pressure
fasting.blood.sugar	Integer	Fasting blood sugar level
Cholesterol	Integer	Total cholesterol level
triglyceride	Integer	Triglyceride level
HDL	Integer	High-density lipoprotein (HDL) cholesterol level
LDL	Integer	Low-density lipoprotein (LDL) cholesterol level
hemoglobin	Numeric	Hemoglobin level
Urine.protein	Integer	Urine protein status (1: present, 2: absent)
serum.creatinine	Numeric	Serum creatinine level
AST	Integer	Aspartate transaminase (AST) level

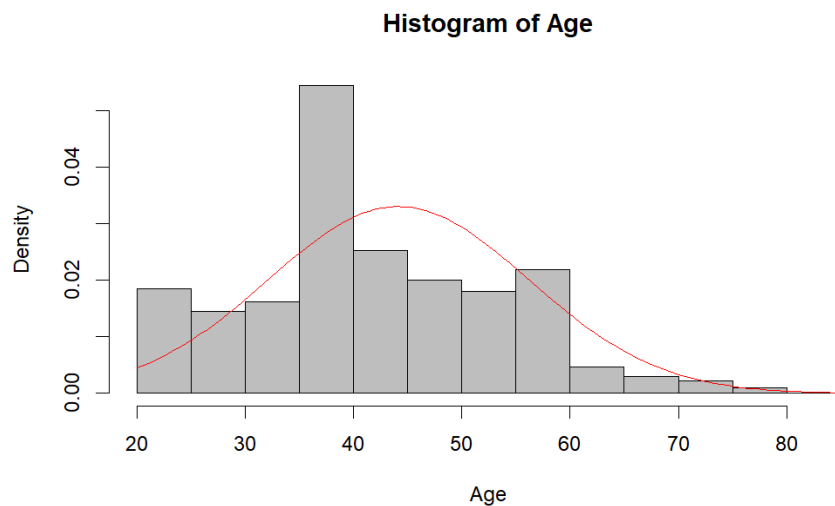
ALT	Integer	Alanine transaminase (ALT) level
Gtp	Integer	Gamma-glutamyl transferase (GTP) level
dental.caries	Integer	Dental caries status (1: present, 0: absent)
smoking	Integer	Smoking status (1: smoker, 0: non-smoker)

3.0 DATA ANALYSIS

3.1 Attributes Distributions

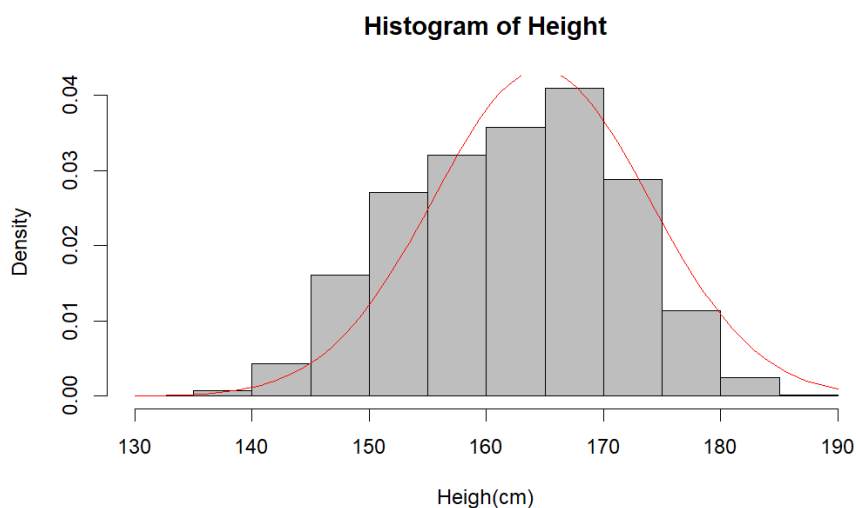
The attributes distributions shown below are followed by the significant attributes.

3.1.1 Age



Based on the histogram above, the 'Age' distribution is approximately normally distributed since it is symmetric bell-shaped with the highest point at the mean. The most common age is around 40, as indicated by the highest point on the histogram and density curve.

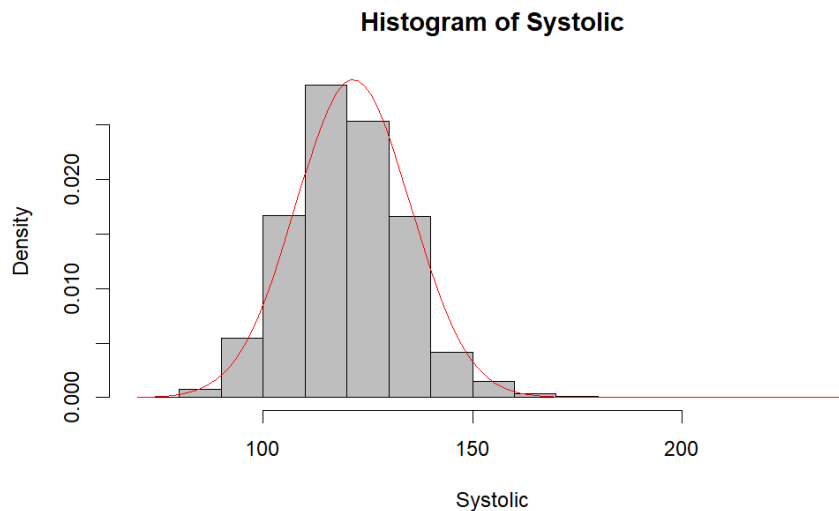
3.1.2 Height



Based on the histogram above, the 'Height' distribution is approximately normally distributed since it is symmetric bell-shaped with the highest point at the mean. The most

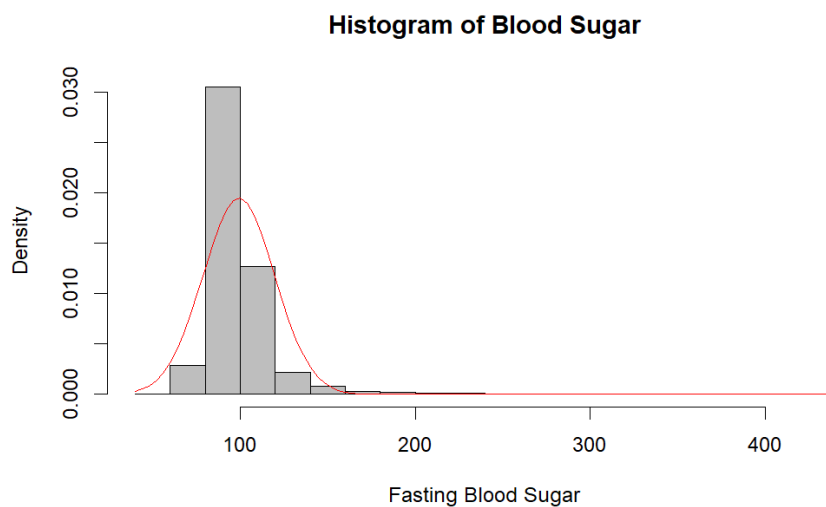
common height range is around 160-170 cm, as indicated by the highest point on the histogram and density curve. This means that most individuals in this dataset have a height within this range.

3.1.3 Systolic



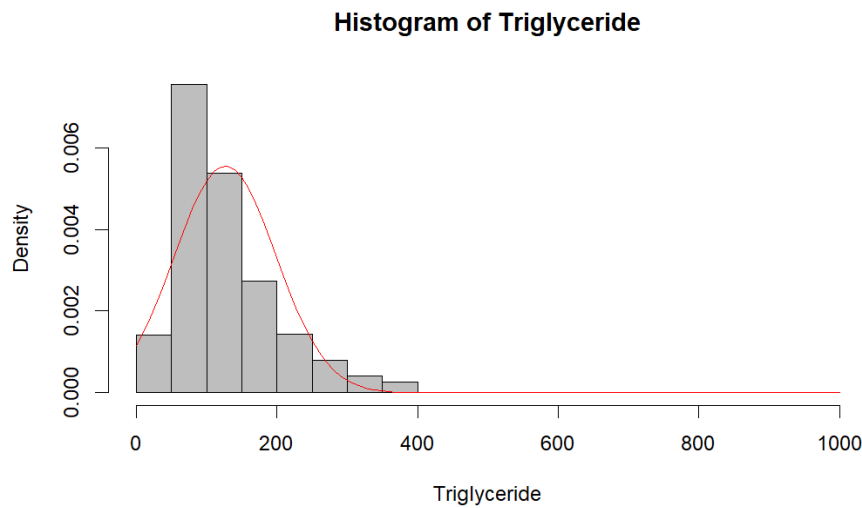
Based on the histogram above, the 'Systolic' distribution is approximately normally distributed since it is symmetric bell-shaped with the highest point at the mean. The most common systolic blood pressure is around 120, as indicated by the highest point on the histogram and density curve.

3.1.4 Blood Sugar



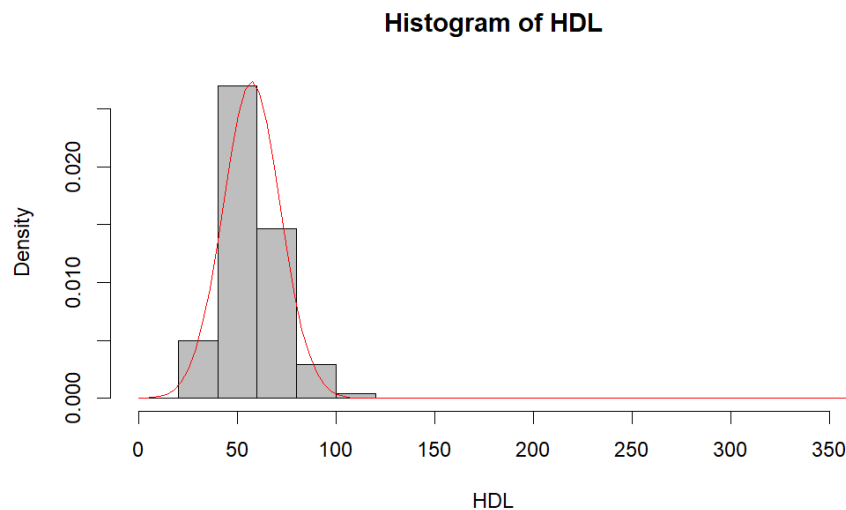
Based on the histogram above, the 'Blood Sugar' distribution is approximately skewed to the right with a peak at 100. This means that most individuals in this dataset have a fasting blood sugar level of around 100.

3.1.5 Triglyceride



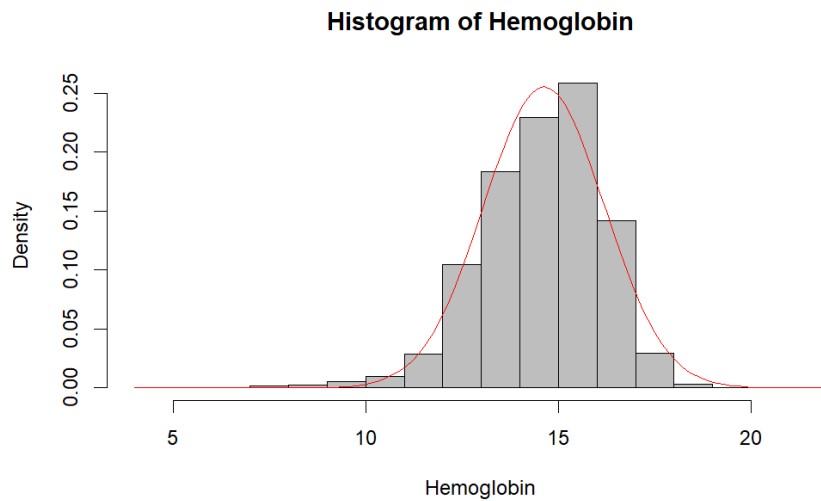
From the histogram and density curve, we can see that the distribution of triglyceride levels is skewed to the right, with a peak around 0-200. This means that most individuals in this dataset have low triglyceride levels, but there are also a significant number of individuals with higher levels.

3.1.6 HDL



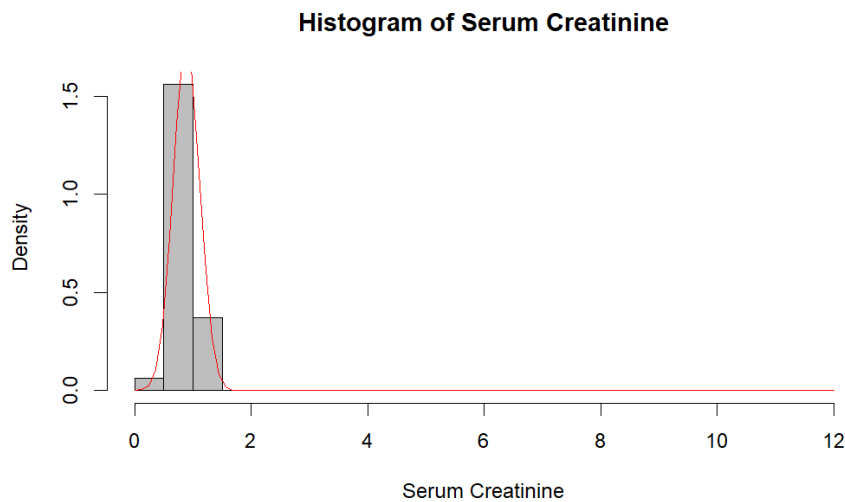
From the histogram and density curve, we can see that the distribution of HDL levels is skewed to the right, with a peak around 50-100. This means that most individuals in this dataset have their HDL levels within this range, but there are also a significant number of individuals with higher levels.

3.1.7 Hemoglobin



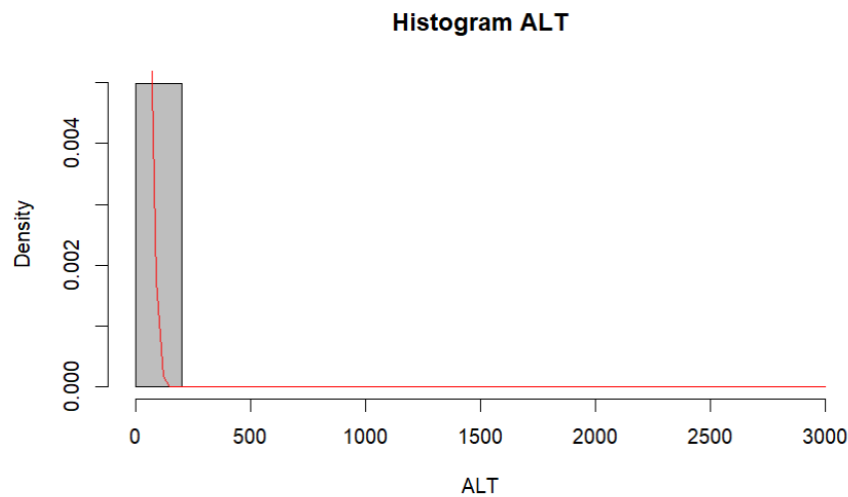
From the histogram and density curve, we can see that the distribution of hemoglobin levels is skewed to the left, with a peak around 15. This means that most individuals in this dataset have their hemoglobin levels within this range.

3.1.8 Serum Creatinine



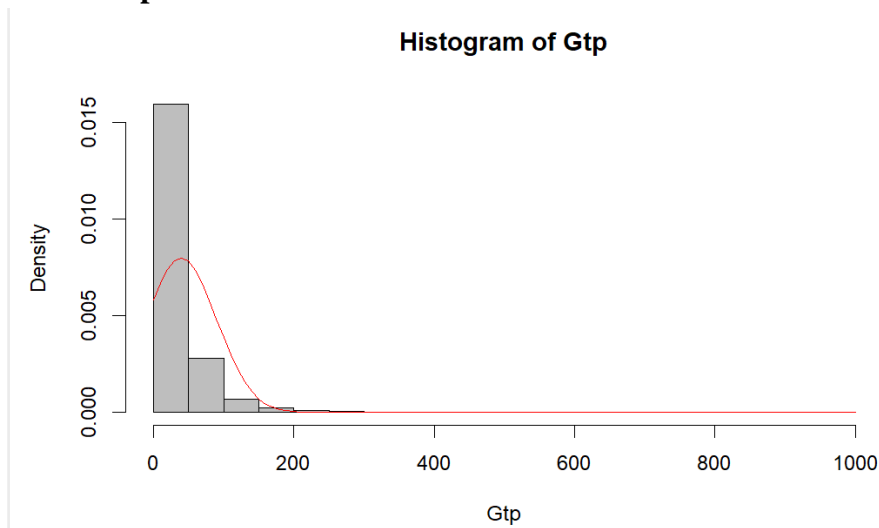
From the histogram and density curve, we can see that the distribution of serum creatinine levels is skewed to the right, with a peak around 1-2. This means that most individuals in this dataset have their serum creatinine levels within this range, but there are also a significant number of individuals with higher levels.

3.1.9 ALT



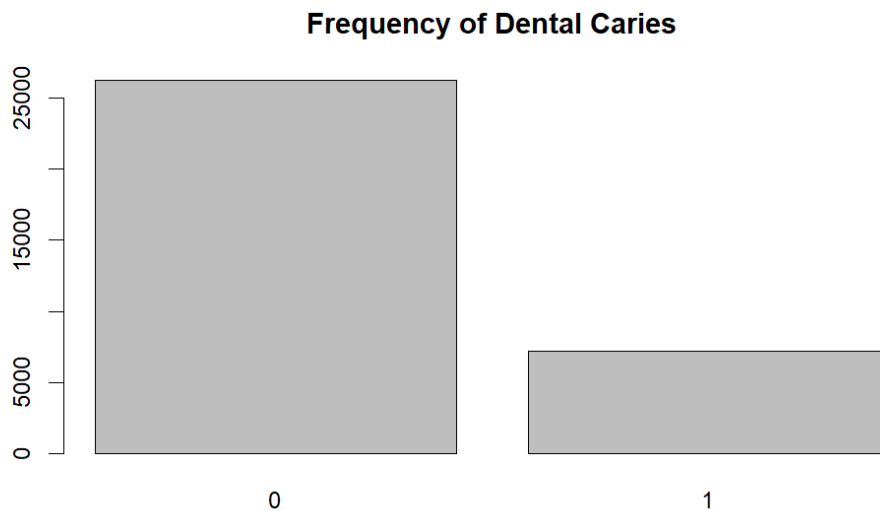
From the histogram and density curve, we can see that the distribution of ALT levels is skewed to the right, with a peak around 0-500. This means that most individuals in this dataset have their ALT levels within this range, but there are also a significant number of individuals with higher levels.

3.1.10 Gtp



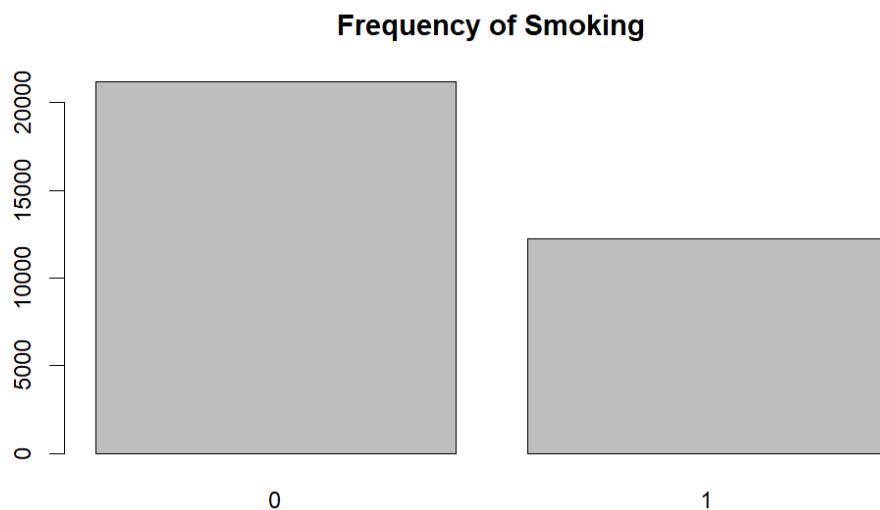
From the histogram and density curve, we can see that the distribution of Gtp levels is skewed to the right, with a peak around 0-200. This means that most individuals in this dataset have their Gtp levels within this range, but there are also a significant number of individuals with higher levels.

3.1.11 Dental Caries



Category “0” represents no dental caries while category “1” represents dental caries. From the bar graph, we can see that the category “0” has a significantly higher frequency of dental caries than the category “1” with around 25000 counts while category “1” has a frequency of around 7500 counts. This means that most of the people do not have dental caries.

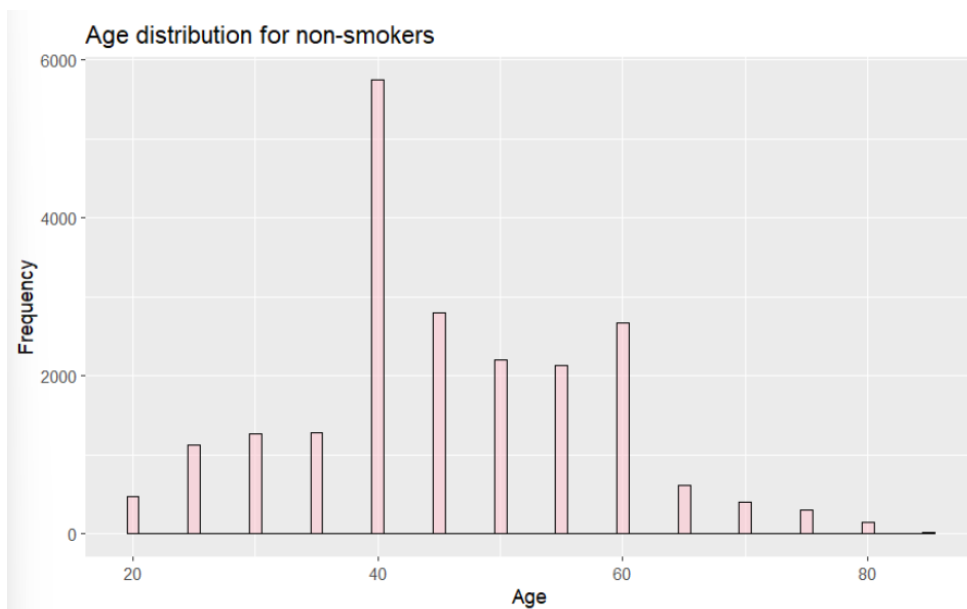
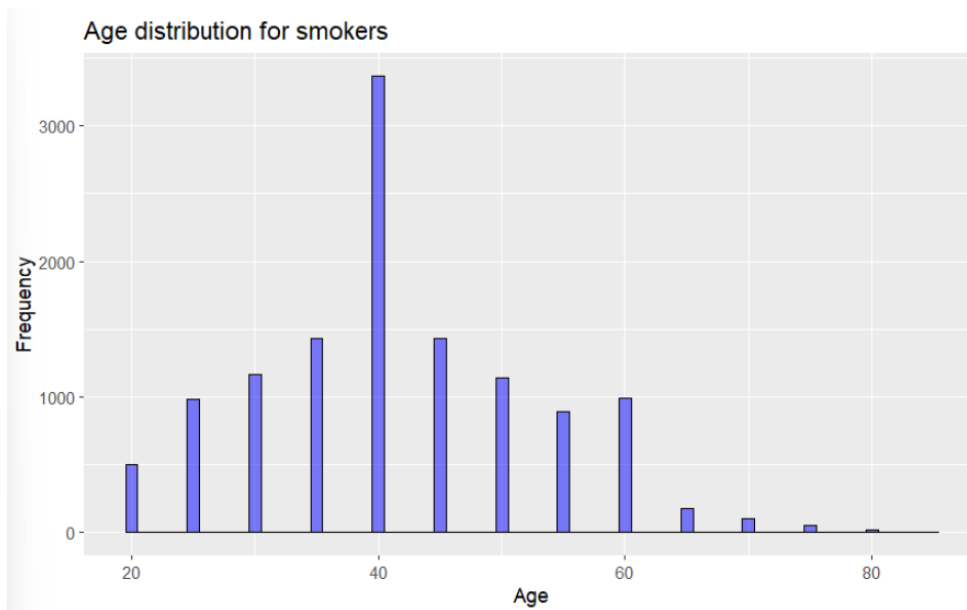
3.1.12 Smoking



Category “0” represents no smoking while category “1” represents smoking. From the bar graph, we can see that the category “0” has a significantly higher frequency of smoking than the category “1” with around 20000 counts while category “1” has a frequency of around 12500 counts. This means that most of the people are non-smokers.

3.2 Exploratory Analysis and Visualization

3.2.1 Age Distribution for Smokers and Non-smokers



Based on both the histograms, the frequency of smokers and non-smokers is the highest at age 40 while the lowest is at age 80 respectively.

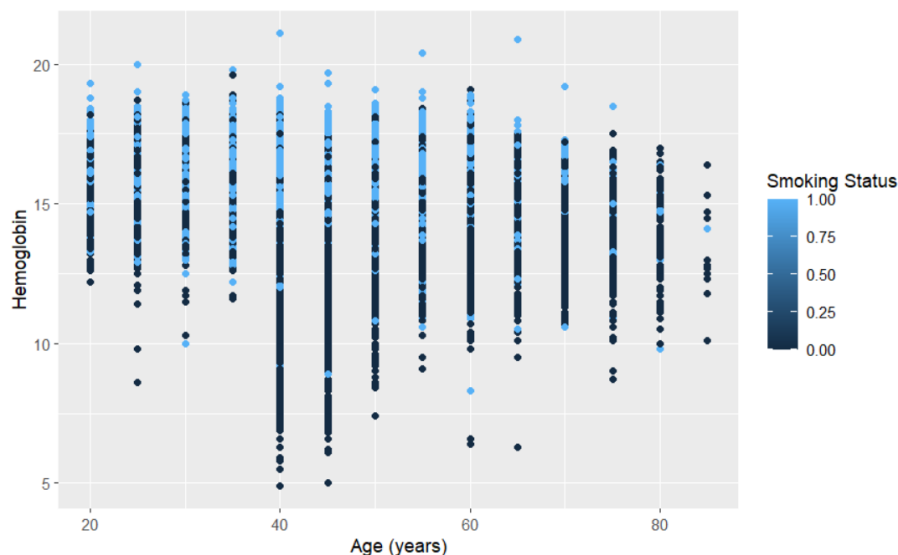
3.2.2 Percentage of Smoking Status

Percentage of Smoking Status



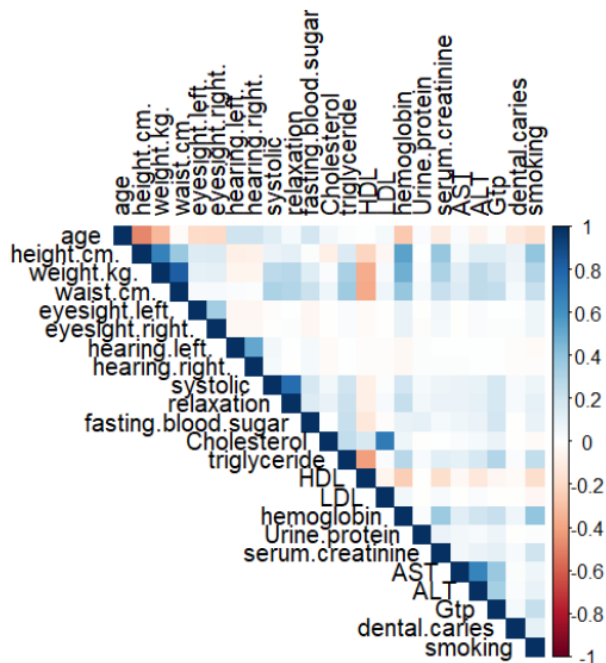
Based on the pie chart, non-smokers have a higher percentage of smoking status of 63.4% while another 36.6% are smokers.

3.2.3 Hemoglobin vs Age



Based on this scatter plot, we can see that the light blue colour represents the person who is a smoker and the darker blue colour represents the person who is not a smoker. Based on the scatter plot, people who are older are likely not a smoker. Moreover, people who smoke have an overly high level of hemoglobin. This is because the carbon monoxide in cigarette smoke blocks oxygen attachment to the red cells' empty hemoglobin slots, causing the body to increase red blood cell production.

3.2.4 Correlation Matrix among the Attributes



From the heatmap, the darker blue colour represents the strong correlation between the variables. There appears to be a number that shows a negative number, which means that when the number is negative, the data has a relationship that is rather than reversed. If the number is approaching zero, the variables do not correlate at all.

Based on the heatmap, when investigating the correlation between the response variable smoking and the explanatory variables, the explanatory variables that have a weak positive correlation with the highest to the response variable smoking among the attributes in this dataset are height.cm., weight.kg., waist.cm., triglyceride, hemoglobin, serum.creatinine and Gtp.

4.0 DATA MODELLING

Extract and load the dataset into R studio

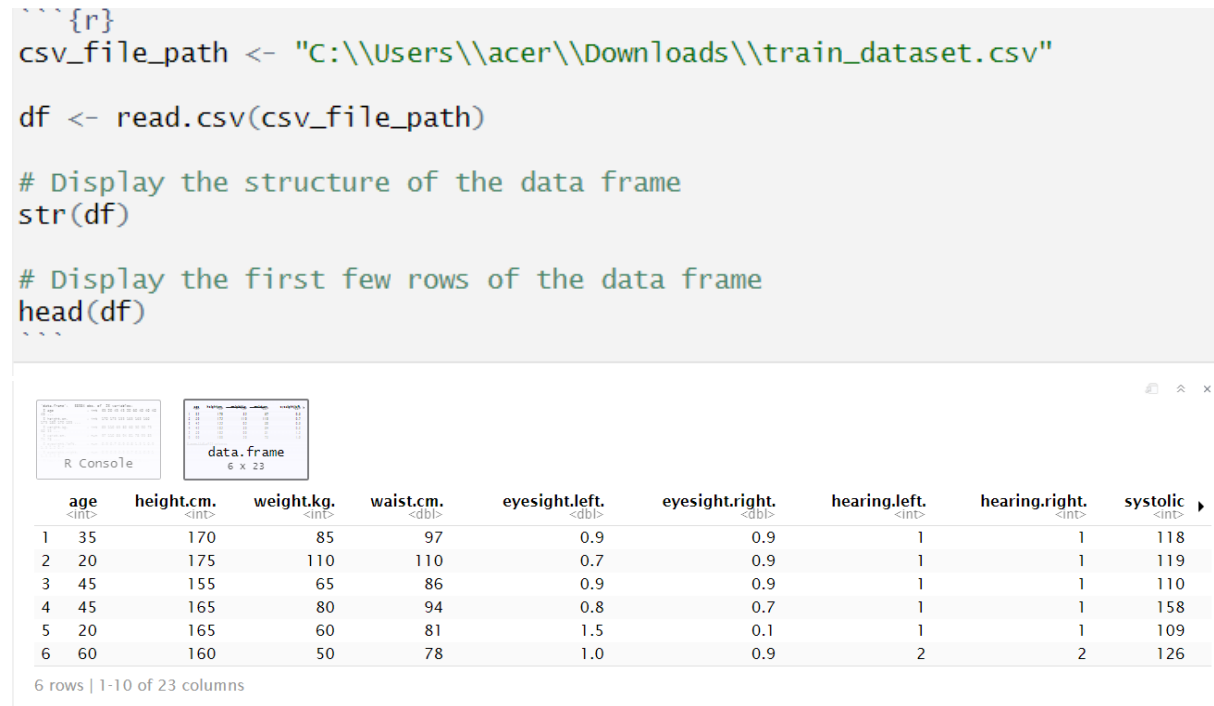


Figure 4.1 Loading Data

The dataset is loaded into R studio. The figure above shows the head of the data frame imported

```

'data.frame':  33467 obs. of  23 variables:
 $ age           : int  35 20 45 45 20 60 40 40 40 45 ...
 $ height.cm.    : int  170 175 155 165 165 160 175 180 170 155 ...
 $ weight.kg.    : int  85 110 65 80 60 50 90 75 60 55 ...
 $ waist.cm.     : num  97 110 86 94 81 78 95 85 74 78 ...
 $ eyesight.left.: num  0.9 0.7 0.9 0.8 1.5 1 0.9 1.5 1.2 0.7 ...
 $ eyesight.right.: num  0.9 0.9 0.9 0.7 0.1 0.9 1 1.5 1.5 1 ...
 $ hearing.left. : int  1 1 1 1 1 2 1 1 1 1 ...
 $ hearing.right.: int  1 1 1 1 1 2 1 1 1 1 ...
 $ systolic      : int  118 119 110 158 109 126 130 110 89 114 ...
 $ relaxation    : int  78 79 80 88 64 75 88 60 57 81 ...
 $ fasting.blood.sugar: int  97 88 80 249 100 114 90 100 83 96 ...
 $ cholesterol   : int  239 211 193 210 179 177 207 170 178 184 ...
 $ triglyceride  : int  153 128 120 366 200 74 331 62 69 177 ...
 $ HDL           : int  70 71 57 46 47 98 39 58 60 41 ...
 $ LDL           : int  142 114 112 91 92 64 102 99 104 107 ...
 $ hemoglobin    : num  19.8 15.9 13.7 16.9 14.9 13.9 16.5 14 12.9 13.1 ...
 $ Urine.protein : int  1 1 3 1 1 1 1 2 2 1 ...
 $ serum.creatinine : num  1 1.1 0.6 0.9 1.2 1 1 1.4 0.7 0.6 ...
 $ AST           : int  61 19 1090 32 26 47 19 29 17 22 ...
 $ ALT           : int  115 25 1400 36 28 23 22 20 17 15 ...
 $ Gtp           : int  125 30 276 36 15 70 19 32 14 56 ...
 $ dental.caries  : int  1 1 0 0 0 0 0 1 0 0 ...
 $ smoking       : int  1 0 0 0 0 1 0 1 0 0 ...

```

Figure 4.2 Data structure

The figure shows all the attribute names within the dataframe and their data type respectively.

age	height.cm.	weight.kg.	waist.cm.	eyesight.left.	eyesight.right.	hearing.left.	hearing.right.	systolic	relaxation
Min. :20.00	Min. :130.0	Min. : 30.00	Min. : 51.00	Min. :0.100	Min. :0.10	Min. :1.000	Min. :1.000	Min. : 71.0	Min. : 40.00
1st Qu.:40.00	1st Qu.:160.0	1st Qu. : 55.00	1st Qu. : 76.00	1st Qu.:0.800	1st Qu.:0.80	1st Qu.:1.000	1st Qu.:1.000	1st Qu.:112.0	1st Qu. : 70.00
Median :40.00	Median :165.0	Median : 65.00	Median : 82.00	Median :1.000	Median :1.00	Median :1.000	Median :1.000	Median :120.0	Median : 76.00
Mean :44.15	Mean :164.7	Mean : 65.93	Mean : 82.08	Mean :1.014	Mean :1.01	Mean :1.025	Mean :1.026	Mean :121.5	Mean : 76.02
3rd Qu.:55.00	3rd Qu.:170.0	3rd Qu. : 75.00	3rd Qu. : 88.00	3rd Qu.:1.200	3rd Qu.:1.20	3rd Qu.:1.000	3rd Qu.:1.000	3rd Qu.:130.0	3rd Qu. : 82.00
Max. :85.00	Max. :190.0	Max. :135.00	Max. :129.00	Max. :9.900	Max. :9.90	Max. :2.000	Max. :2.000	Max. :233.0	Max. :146.00
fasting.blood.sugar	cholesterol	triglyceride	HDL	LDL	hemoglobin	urine.protein	serum.creatinine	AST	ALT
Min. : 46.00	Min. : 55	Min. : 8.0	Min. : 4.00	Min. : 1.0	Min. : 4.90	Min. :1.000	Min. : 0.1000	Min. : 6.0	Min. : 1.00
1st Qu. : 89.00	1st Qu.:172	1st Qu. : 75.0	1st Qu. : 47.00	1st Qu. : 92.0	1st Qu.:13.60	1st Qu.:1.000	1st Qu. : 0.8000	1st Qu. : 19.0	1st Qu. : 15.00
Median : 96.00	Median :195	Median :108.0	Median : 55.00	Median :113.0	Median :14.80	Median :1.000	Median : 0.9000	Median : 23.0	Median : 21.00
Mean : 99.26	Mean :197	Mean :126.8	Mean : 57.26	Mean :115.2	Mean :14.62	Mean :1.087	Mean : 0.8865	Mean : 26.2	Mean : 27.14
3rd Qu.:104.00	3rd Qu.:220	3rd Qu.:160.0	3rd Qu. : 66.00	3rd Qu.:136.0	3rd Qu.:15.70	3rd Qu.:1.000	3rd Qu. : 1.0000	3rd Qu. : 29.0	3rd Qu. : 31.00
Max. :423.00	Max. :445	Max. :999.0	Max. :359.00	Max. :1860.0	Max. :21.10	Max. :6.000	Max. :11.6000	Max. :1090.0	Max. :2914.00
Gtp	dental.caries	smoking							
Min. : 2.00	Min. :0.0000	Min. :0.0000							
1st Qu. :17.00	1st Qu.:0.0000	1st Qu.:0.0000							
Median :26.00	Median :0.0000	Median :0.0000							
Mean :39.95	Mean :0.2147	Mean :0.3663							
3rd Qu. :44.00	3rd Qu.:0.0000	3rd Qu.:1.0000							
Max. :999.00	Max. :1.0000	Max. :1.0000							

Figure 4.3 Summary of dataframe

The provided summary depicts various statistical measures for each attribute in the dataset. It includes information such as the minimum and maximum values, quartiles, and the mean for all the attributes involved.

```

***{r}

logreg <- glm(formula = smoking ~ ., family = binomial(link="logit"), data = df)
summary(logreg)
***

Call:
glm(formula = smoking ~ ., family = binomial(link = "logit"),
    data = df)

Coefficients:
              Estimate Std. Error z value Pr(>|z|)
(Intercept) -1.993e+01  4.679e-01 -42.589  < 2e-16 ***
age          8.450e-04  1.399e-03   0.604  0.54593
height.cm.   8.282e-02  2.511e-03  32.974  < 2e-16 ***
weight.kg.   -2.441e-02  2.685e-03  -9.093  < 2e-16 ***
waist.cm.    1.453e-02  3.140e-03   4.627  3.72e-06 ***
eyesight.left. 2.359e-02  2.800e-02   0.842  0.39954
eyesight.right. 1.786e-03  2.797e-02   0.064  0.94910
hearing.left. -9.824e-02  1.030e-01  -0.954  0.34018
hearing.right. 1.169e-01  1.001e-01   1.168  0.24273
systolic     -9.436e-03  1.564e-03  -6.032  1.62e-09 ***
relaxation    4.526e-03  2.154e-03   2.102  0.03559 *
fasting.blood.sugar 4.174e-03  6.930e-04   6.024  1.71e-09 ***
cholesterol  -5.055e-03  6.157e-04  -8.210  < 2e-16 ***
triglyceride  4.566e-03  2.476e-04  18.442  < 2e-16 ***
HDL          -3.344e-03  1.244e-03  -2.689  0.00717 **
LDL          2.828e-04  4.641e-04   0.609  0.54223
hemoglobin    4.400e-01  1.268e-02  34.715  < 2e-16 ***
urine.protein -6.612e-02  3.372e-02  -1.961  0.04988 *
serum.creatinine 1.755e-01  6.809e-02   2.578  0.00994 **
AST          -3.490e-04  1.282e-03  -0.272  0.78549
ALT          -6.837e-03  9.982e-04  -6.849  7.45e-12 ***
Gtp          9.747e-03  4.346e-04  22.428  < 2e-16 ***
dental.caries  4.104e-01  3.157e-02  13.000  < 2e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for binomial family taken to be 1)

    Null deviance: 43972  on 33466  degrees of freedom
Residual deviance: 34100  on 33444  degrees of freedom
AIC: 34146

```

Figure 4.4 Full Logistic Model

All the attributes are used to fit into the basic logistic model with binomial distribution. The target variable is “smoking”. The figure above shows the null deviance and residual deviance. The AIC value is 34146

```

##{r}
null_model <- glm(formula = smoking ~ 1, family = binomial(link = "logit"), data = df)
summary(null_model)
##{r}

Call:
glm(formula = smoking ~ 1, family = binomial(link = "logit"),
    data = df)

Coefficients:
              Estimate Std. Error z value Pr(>|z|)
(Intercept) -0.54825     0.01135  -48.32  <2e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for binomial family taken to be 1)

    Null deviance: 43972  on 33466  degrees of freedom
Residual deviance: 43972  on 33466  degrees of freedom
AIC: 43974

Number of Fisher Scoring iterations: 4

```

Figure 4.5 Null Model

The figure above shows the null model with AIC value of 43974. The AIC (Akaike Information Criterion) is a measure that balances the goodness of fit of a model with its complexity, penalizing models that have more parameters. The goal is to find a model that fits the data well but is not overly complex. In the context of model selection criteria like AIC, lower values are preferred.

By comparing the two models, we can conclude that the null model does not offer a better fit as it has a higher AIC value than the full model.

```

##{r}
# Install and load the 'car' package (if not already installed)
# install.packages("car")
library(car)

# Assuming 'lm_model' is your linear regression model
# Make sure your model is fitted before calculating VIF

# Calculate VIF
vif_values <- car::vif(logreg)

# Print the VIF values
print(vif_values)
##{r}

```

age	height.cm	weight.kg	waist.cm	eyesight.left	eyesight.right	hearing.left	hearing.right	systolic	relaxation
1.687685	2.273006	6.248744	4.453792	1.137716	1.139278	1.358622	1.363709	2.447715	2.395523
fasting.blood.sugar	cholesterol	triglyceride	HDL	LDL	hemoglobin	urine.protein	serum.creatinine	AST	ALT
1.139308	2.808775	1.766252	1.694096	2.359566	1.389577	1.022667	1.161300	2.275334	2.791200
gtp	dental.caries								
1.455968	1.012046								

Figure 4.6 Multicollinearity Checking

```

***{r}
anova(logreg, test="Chisq")
***

Analysis of Deviance Table

Model: binomial, link: logit

Response: smoking

Terms added sequentially (first to last)


```

	Df	Deviance	Resid. Df	Resid. Dev	Pr(>Chi)
NULL			33466	43972	
age	1	956.1	33465	43016	< 2.2e-16 ***
height.cm.	1	4698.2	33464	38317	< 2.2e-16 ***
weight.kg.	1	55.0	33463	38262	1.218e-13 ***
waist.cm.	1	233.9	33462	38029	< 2.2e-16 ***
eyesight.left.	1	2.2	33461	38026	0.1421187
eyesight.right.	1	0.5	33460	38026	0.4785978
hearing.left.	1	0.0	33459	38026	0.8934461
hearing.right.	1	1.0	33458	38025	0.3084187
systolic	1	11.9	33457	38013	0.0005614 ***
relaxation	1	84.9	33456	37928	< 2.2e-16 ***
fasting.blood.sugar	1	220.6	33455	37707	< 2.2e-16 ***
Cholesterol	1	2.8	33454	37705	0.0919096 .
triglyceride	1	1099.1	33453	36606	< 2.2e-16 ***
HDL	1	5.6	33452	36600	0.0178052 *
LDL	1	1.0	33451	36599	0.3084939
hemoglobin	1	1617.7	33450	34981	< 2.2e-16 ***
Urine.protein	1	0.6	33449	34981	0.4505911
serum.creatinine	1	11.7	33448	34969	0.0006299 ***
AST	1	2.5	33447	34966	0.1104928
ALT	1	6.6	33446	34960	0.0099481 **
Gtp	1	691.0	33445	34269	< 2.2e-16 ***
dental.caries	1	168.7	33444	34100	< 2.2e-16 ***

```

---
signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

Figure 4.7 ANOVA test

Results & Interpretation#####

Significance Codes:

'***': Very highly significant (p-value < 0.001).

'**': Highly significant (p-value < 0.01).

'*': Significant at a 5% level (p-value < 0.05).

' ': Not significant (p-value > 0.1).

Results from the full logistic regression model show that there are some insignificant variables such as "age ", "eyesight.left.", "eyesight.right.", "hearing.left. ", "hearing.right. ", "Cholesterol ", "LDL ", "AST " " and "relaxation" based on its respective p-values.

Variance Inflation Factors (VIF).

The Variance Inflation Factor (VIF) measures the inflation in the coefficient of the independent variable due to the collinearities among the other independent variables.

A VIF of 1 means that the regression coefficient is not inflated by the presence of the other predictors, and hence multicollinearity does not exist.

Ideally, the Variance Inflation Factors are below 5.

Results from multicollinearity with VIF test shows that weight.kg. (VIF: 6.23): This variable has a relatively high VIF, indicating that its variance is inflated due to its correlation with other predictors and waist.cm. (VIF: 4.43): While this VIF is above 2, it is not extremely high. However, it suggests some correlation with other predictors.

Results from the ANOVA test show that "eyesight.left.", "eyesight.right.", "hearing.left. ", "hearing.right. ", "Cholesterol ", "LDL", "Urine.protein", and "AST " are insignificant to the fitted model.

Hence, fit the updated glm() model (logistic regression) as a reduced model without the insignificant variables and non-collinear variables into logreg2 in order to develop the best model in our case study.


```

```{r}
Fit the updated glm() model (logistic regression) into logreg2.
logreg2 <- glm(formula = smoking ~ age + height.cm. + systolic +
 fasting.blood.sugar + triglyceride +
 HDL + hemoglobin + serum.creatinine + ALT + Gtp + dental.caries,
 family = binomial(link="logit"),
 data = df)

summary(logreg2)

```

Call:
glm(formula = smoking ~ age + height.cm. + systolic + fasting.blood.sugar +
    triglyceride + HDL + hemoglobin + serum.creatinine + ALT +
    Gtp + dental.caries, family = binomial(link = "logit"), data = df)

Coefficients:
                Estimate Std. Error z value Pr(>|z|)
(Intercept)    -1.849e+01  3.781e-01 -48.888  < 2e-16 ***
age              3.241e-03  1.281e-03   2.529   0.0114 *
height.cm.       7.032e-02  2.030e-03  34.636  < 2e-16 ***
systolic        -9.216e-03  1.054e-03  -8.745  < 2e-16 ***
fasting.blood.sugar 4.213e-03  6.854e-04   6.148  7.87e-10 ***
triglyceride     3.474e-03  2.146e-04  16.185  < 2e-16 ***
HDL             -5.853e-03  1.075e-03  -5.447  5.13e-08 ***
hemoglobin       4.278e-01  1.248e-02  34.268  < 2e-16 ***
serum.creatinine  9.990e-02  6.859e-02   1.456   0.1453
ALT             -8.969e-03  7.179e-04 -12.493  < 2e-16 ***
Gtp              9.903e-03  4.288e-04  23.097  < 2e-16 ***
dental.caries    4.089e-01  3.144e-02  13.008  < 2e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for binomial family taken to be 1)

    Null deviance: 43972  on 33466  degrees of freedom
Residual deviance: 34369  on 33455  degrees of freedom
AIC: 34393

Number of Fisher Scoring iterations: 5

```

Figure 4.8 Reduced Model

```

```{r}
Use the anova() function to analyze the updated table of deviance.
anova(logreg2, test="chisq")
```

```

Analysis of Deviance Table

Model: binomial, link: logit

Response: smoking

Terms added sequentially (first to last)

| | Df | Deviance | Resid. Df | Resid. Dev | Pr(>Chi) |
|---------------------|----|----------|-----------|------------|---------------|
| NULL | | | 33466 | 43972 | |
| age | 1 | 956.1 | 33465 | 43016 | < 2.2e-16 *** |
| height.cm. | 1 | 4698.2 | 33464 | 38317 | < 2.2e-16 *** |
| systolic | 1 | 40.7 | 33463 | 38277 | 1.788e-10 *** |
| fasting.blood.sugar | 1 | 281.3 | 33462 | 37995 | < 2.2e-16 *** |
| triglyceride | 1 | 1116.1 | 33461 | 36879 | < 2.2e-16 *** |
| HDL | 1 | 21.0 | 33460 | 36858 | 4.532e-06 *** |
| hemoglobin | 1 | 1568.3 | 33459 | 35290 | < 2.2e-16 *** |
| serum.creatinine | 1 | 5.7 | 33458 | 35284 | 0.01699 * |
| ALT | 1 | 5.8 | 33457 | 35279 | 0.01649 * |
| Gtp | 1 | 740.9 | 33456 | 34538 | < 2.2e-16 *** |
| dental.caries | 1 | 168.8 | 33455 | 34369 | < 2.2e-16 *** |

 signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Figure 4.9 Reduced Model ANOVA test

Results & Interpretation

Results from the updated fitted logistic regression model show that only "serum.creatinine" variable is insignificant based on its p-value.

Additionally, results from the ANOVA test show all variables, including "serum.creatinine", are significant to the fitted model.

This is judged by the low deviance residuals as well as the Pr(>Chi) of > .05, respectively.

Hence, the "serum.creatinine" variable is kept in the fitted model.

```
```{r}
anova(logreg,logreg2, test="chisq")
```
```

Analysis of Deviance Table

Model 1: smoking ~ age + height.cm. + weight.kg. + waist.cm. + eyesight.left. + eyesight.right. + hearing.left. + hearing.right. + systolic + relaxation + fasting.blood.sugar + cholesterol + triglyceride + HDL + LDL + hemoglobin + Urine.protein + serum.creatinine + AST + ALT + Gtp + dental.caries

Model 2: smoking ~ age + height.cm. + systolic + fasting.blood.sugar + triglyceride + HDL + hemoglobin + serum.creatinine + ALT + Gtp + dental.caries

| | Resid. Df | Resid. Dev | Df | Deviance | Pr(>Chi) |
|---|-----------|------------|-----|----------|---------------|
| 1 | 33444 | 34100 | | | |
| 2 | 33455 | 34369 | -11 | -268.97 | < 2.2e-16 *** |

 signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Figure 4.10 Comparison of full model and reduced model

H_0 : The full model offers better fit

H_1 : The full model does not offer better fit

The p-value = 0.0000 and is less than $\alpha=0.05$, H_0 is rejected.

Therefore, we can conclude that the reduced model offers better fit than the full model

Test for the Absence of Strongly Influential Outliers

Notes

Test using standardized residuals and Cook's Distance.

Standardized residual values > 3 = influential outlier.

Cook's D value $>$ Cook's D Threshold $(4/N)$ = influential outlier.

```

```{r}
library(dplyr)
library(broom)
Place all the calculated values from the logistic regression model into a new data frame.
logreg.data <- augment(logreg2) %>%
 mutate(index = 1:n())

Show the top 6 highest standardized residuals (if > 3 = influential outlier).
head(logreg.data$.std.resid[order(-logreg.data$.std.resid)])

Plot of standardized residuals
plot(fitted(logreg2),
 rstandard(logreg2))
```

```

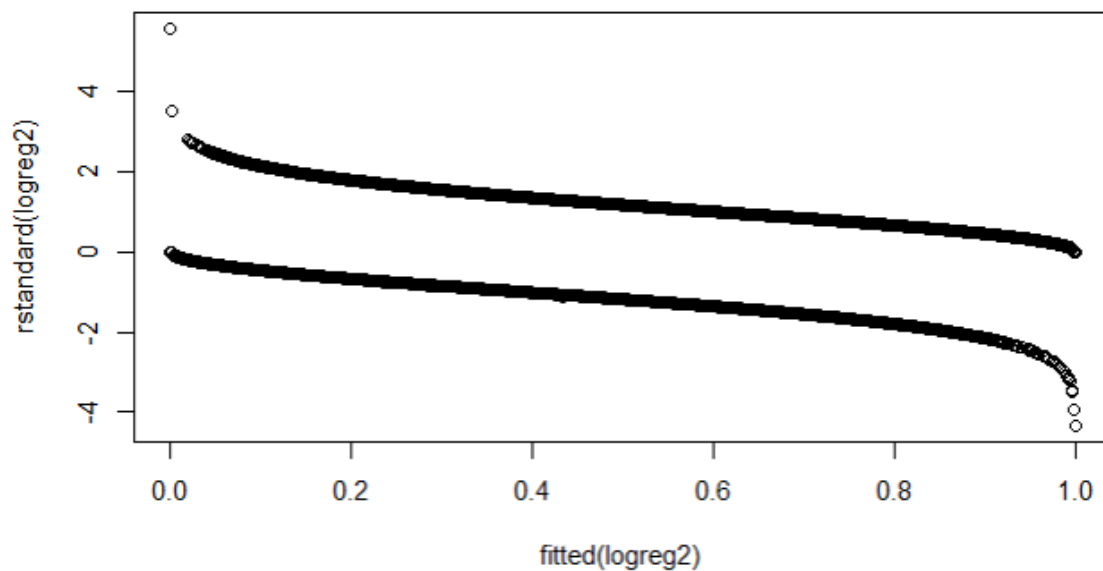


Figure 4.11 Standardized Residual Plot

```

```{r}
Set Cook's D Threshold.
cook_threshold <- 4 / nrow(df)

Cook's D Plot.
plot(logreg2, which = 4, id.n = 12)
abline(h = cook_threshold, col = "red")

Put outlier data into a new data frame where > Cook's D Threshold = influential outliers.
influ_out <- logreg.data %>%
 filter(.cooks_d > cook_threshold)

Get the percentage of influential outliers.
outliers <- round(100*(nrow(influ_out) / nrow(logreg.data)),1)
```

```

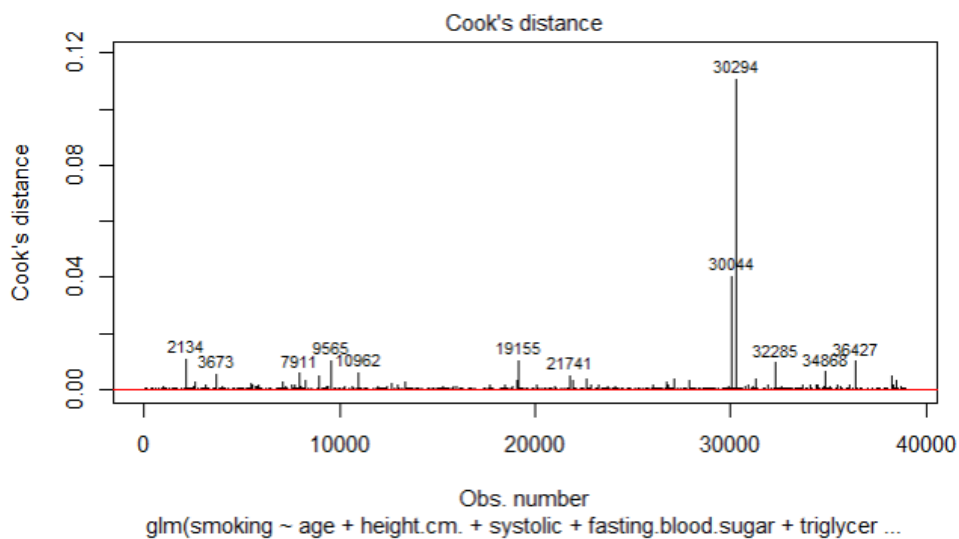


Figure 4.12 Cook Distance Plot

```
##{r}
### Get the percentage of influential outliers.
outliers <- round(100*(nrow(influ_out) / nrow(logreg.data)),1)

### Store values in a data variable.
print_outliers <- format(round(outliers, 2), nsmall = 2)

### Print the number of percentage of observations that exceed Cook's distance threshold.
sprintf('Proportion of data points that are highly influential = %s Percent', print_outliers)
##}
```

[1] "Proportion of data points that are highly influential = 4.10 Percent"

Figure 4.13 Percentage of Influential Outliers

Results & Interpretation

Standardized Residuals.

Results show that none of the data points of the fitted model consists of any outliers.

Cook's Distance.

In addition, based on the pre-defined threshold ($4/N$), only 4.1% of the data points are in the outlier zone, which is small as well.

This highlighted section outlines the justification for choosing the final model, which is regarded as the most favourable solution.

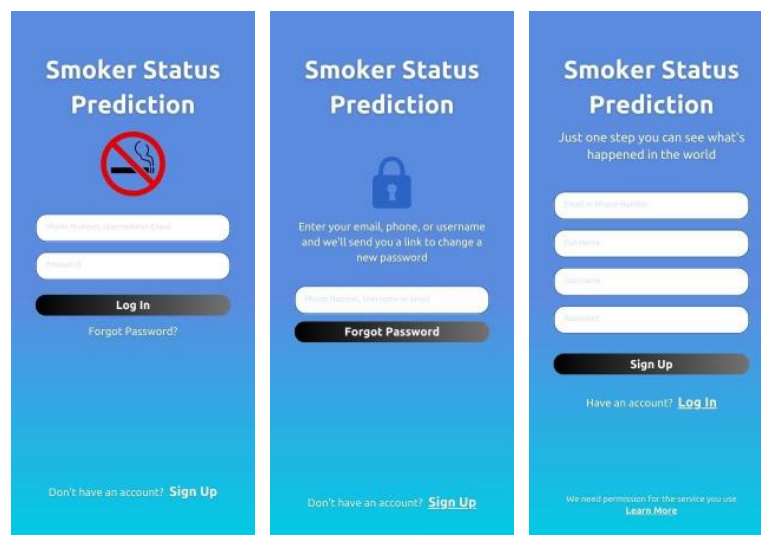
The final logistic regression model was constructed to predict smoking status by considering various health-related variables. Through a stepwise refinement process, insignificant predictors and potential multicollinearity issues were addressed. The resulting model includes significant predictors such as age, height, systolic blood pressure, fasting blood sugar, triglyceride levels, HDL cholesterol, hemoglobin levels, ALT (Alanine Aminotransferase), GTP (Gamma-Glutamyl Transferase), and the presence of dental caries. Notably, serum creatinine levels were found to be statistically insignificant in the full model. Then the reduced model is developed with better fit performance after comparison with full model. Diagnostic tests, including VIF for multicollinearity, standardized residuals and Cook's Distance for influential outliers, and deviance tests, were conducted to ensure the model's robustness. The findings indicate that the model provides a significant fit without strong evidence of multicollinearity or influential outliers. Interpretation of the coefficients suggests how each predictor contributes to the likelihood of smoking based on individual health characteristics, providing a valuable tool for understanding and predicting smoking behavior.

5.0 END PRODUCT

5.1 Introduction

A mobile application for smoker status prediction is a tool that allows people to quickly and efficiently identify whether he or she is a smoker or not. The software would utilize a person's health information such as age, height, weight, blood pressure, hemoglobin, urine protein, and so on to predict a person's smoker status. Furthermore. This tool was developed as a mobile application with a user-friendly interface that allows users to input their preferred data for predictions and review the expected results at the end of pages.

5.2 Smoker Status Prediction Interface



Step 1:

The user can log in by filling up their phone number, username or email and password. If a user forgets their password, they can click the 'Forgot Password?' button and it will redirect the user to a page where they can reset their password via a link after entering their phone number, username or email. If a user is new to this application and has not yet signed up, they can do so by clicking the 'Sign Up' button below and creating a new account.



Step 2:

Once logged in to an account, it will jump to the next page which is the main menu of the application. It shows that there is the title of the application 'Smoker Status Prediction' on the top and a greeting to the user below the title. After that, three buttons are given to the user to select, the user can choose the 'New Test' button to start to predict the smoker status. If the user has tested the prediction, they can click the 'History' button to review the preview results. Also, the user can choose the 'Quit' button to exit the interface.

PLEASE FILL YOUR INFORMATION

* You may refer to your medical report if you want to get a more accurate result.

Age :

Height (cm) :

Weight (kg) :

Waist circumference length (cm) :

Eyesight (left) :

Eyesight (right) :

Hearing (left) :

Hearing (right) :

Continue

PLEASE FILL YOUR INFORMATION

Systolic (Blood Pressure) :

Relaxation (Blood Pressure) :

Fasting Blood Sugar :

Cholesterol :

Triglyceride :

HDL (Cholesterol Type) :

LDL (Cholesterol Type) :

Hemoglobin :

Continue

PLEASE FILL YOUR INFORMATION

Urine Protein :

Serum Creatinine :

AST (glutamic oxaloacetic transaminase type) :

ALT (glutamic oxaloacetic transaminase type) :

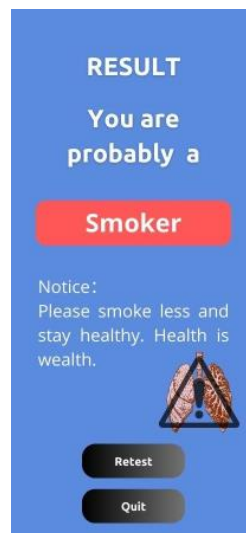
γ-GTP :

Dental caries :

SUBMIT

Step 3:

On these pages, the user can fill up their medical information to predict the result. If the user forgets or is not sure about their information, they may refer to their medical check-up report to enter the data in order to get more accurate results at the end. Moreover, the figures show the sample information of a person to test whether he/she is a smoker or non-smoker. After filling up the data, the user can click the 'SUBMIT' button to start the prediction.



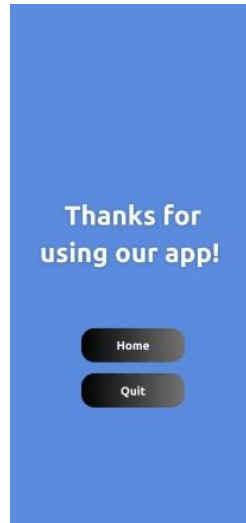
Step 4 a:

After submitting the data, the result will be shown in the figure above. For the sample data, the result shows the user is probably a smoker based on his / her medical information. In addition, there is also a notice to motivate the user to smoke less and stay healthy. Then, there are two buttons shown below the page which are the 'Retest' and 'Quit' buttons. The user can click the 'Retest' button to re-do the prediction again if they are not satisfied with the result or they can click the 'Quit' button to exit the interface.



Step 4 b:

This page shows a good result which the user is probably not a smoker based on the information given. Moreover, there is also a notice to the user and two buttons are shown below the page which are the 'Retest' and 'Quit' buttons. The user can click the 'Retest' button to redo the prediction again or they can click the 'Quit' button to exit the interface.



Step 5:

Lastly, if a user wants to log back into the home page, they can click the 'Home' button, which will take them to the login page. Otherwise, they can click the 'Quit' button to exit the interface.

6.0 CONCLUSION

In a nutshell, the conclusion of the project on statistical modelling and simulation for predicting smoking status using health-related variables is that a logistic regression model was developed to predict smoking status based on various health characteristics. The model was refined through a stepwise process, addressing predictors and potential multicollinearity issues. The resulting model includes significant predictors such as age, height, blood pressure, cholesterol levels, and other health-related variables. Diagnostic tests were conducted to ensure the model's robustness, indicating a significant fit without strong evidence of multicollinearity or influential outliers. The model provides a valuable tool for understanding and predicting smoking behaviour. Additionally, a mobile application was developed as the end product, providing a user-friendly interface for inputting medical information and predicting smoking status. This project demonstrates the potential of machine learning techniques in constructing health outcome prediction models and offers a practical tool for individuals and healthcare professionals to comprehend and apply smoking cessation strategies.

Moreover, this project is an excellent representation of the critical role that entrepreneurship skills play in a variety of fields: it requires proficiency in statistical analysis, application development, data analysis, predictive modelling, and business planning. This project highlights the entrepreneurial skills needed to interpret data, create user-friendly applications, and innovate in business strategy. It involves everything from identifying predictors for smoking status to developing a mobile app for health prediction and using machine learning techniques for outcome modelling. It emphasises how important it is to use data-driven insights to make strategic decisions and promote industry growth, and how important entrepreneurial savvy is to generating value and developing creative solutions.

7.0 REFERENCE

Centers for Disease Control and Prevention. “CDC - Fact Sheet - Health Effects of Cigarette Smoking - Smoking & Tobacco Use.” *Smoking and Tobacco Use*, U.S. Department of Health & Human Services, 29 Oct. 2021, www.cdc.gov/tobacco/data_statistics/fact_sheets/health_effects/effects_cig_smoking/index.htm.

How Does Smoking Affects Red Blood Cell Count? [Fact Checked!]. 8 Sept. 2022, scienceoxygen.com/how-does-smoking-affects-red-blood-cell-count/. Accessed 7 Jan. 2024.

“Tobacco: Health Benefits of Smoking Cessation.” *Www.who.int*, 25 Feb. 2020, www.who.int/news-room/questions-and-answers/item/tobacco-health-benefits-of-smoking-cessation.

“Why Is Hemoglobin Higher in Smokers? – Heimduo.” *Heimduo.org*, heimduo.org/why-is-hemoglobin-higher-in-smokers/. Accessed 7 Jan. 2024.

World Health Organization. “Tobacco.” *World Health Organization*, 31 July 2023, www.who.int/news-room/fact-sheets/detail/tobacco.

SMS Project

Group 3

Code

Hide

```
csv_file_path <- "C:/UMP SEM 5/BSD3443 STATISTICAL MODELLING AND SIMULATION/new title Group P
roject/train_dataset.csv"

df <- read.csv(csv_file_path)

# Display the first few rows of the data frame
head(df)
```

| a.. | height.cm. | weight.kg. | waist.cm. | eyesight.left. | eyesight.right. | hearing.left. |
|-------|------------|------------|-----------|----------------|-----------------|---------------|
| <int> | <int> | <int> | <dbl> | <dbl> | <dbl> | <int> |
| 1 35 | 170 | 85 | 97 | 0.9 | 0.9 | 1 |
| 2 20 | 175 | 110 | 110 | 0.7 | 0.9 | 1 |
| 3 45 | 155 | 65 | 86 | 0.9 | 0.9 | 1 |
| 4 45 | 165 | 80 | 94 | 0.8 | 0.7 | 1 |
| 5 20 | 165 | 60 | 81 | 1.5 | 0.1 | 1 |
| 6 60 | 160 | 50 | 78 | 1.0 | 0.9 | 2 |

6 rows | 1-9 of 23 columns

Hide

```
#get column names
names(df)
```

```
[1] "age"           "height.cm."    "weight.kg."    "waist.cm."
"eyesight.left." "eyesight.right."
[7] "hearing.left." "hearing.right." "systolic"      "relaxation"
"fasting.blood.sugar" "Cholesterol"
[13] "triglyceride"  "HDL"           "LDL"           "hemoglobin"
"Urine.protein"    "serum.creatinine"
[19] "AST"           "ALT"           "Gtp"           "dental.caries"
"smoking"
```

Hide

```
#check if any null value
sum(is.na(df))
```

```
[1] 0
```

Hide

```
#Check if any duplicated data  
sum(duplicated(df))
```

[1] 5517

Hide

```
duplicated_rows <- df[duplicated(df),]  
duplicated_rows
```

| | a..
<int> | height.cm.
<int> | weight.kg.
<int> | waist.cm.
<dbl> | eyesight.left.
<dbl> | eyesight.right.
<dbl> | hearing.le
<int> |
|------|--------------|---------------------|---------------------|--------------------|-------------------------|--------------------------|---------------------|
| 702 | 20 | 165 | 60 | 78.0 | 0.9 | 0.2 | |
| 1132 | 50 | 175 | 70 | 78.5 | 1.5 | 1.0 | |
| 1144 | 50 | 160 | 65 | 82.0 | 1.0 | 1.0 | |
| 1233 | 45 | 155 | 45 | 68.0 | 0.2 | 0.2 | |
| 1346 | 25 | 165 | 85 | 92.3 | 1.2 | 1.2 | |
| 1463 | 40 | 165 | 65 | 80.0 | 1.0 | 1.2 | |
| 1485 | 45 | 155 | 60 | 75.0 | 1.0 | 1.0 | |
| 1709 | 50 | 170 | 70 | 86.0 | 0.5 | 0.5 | |
| 1750 | 75 | 150 | 50 | 75.0 | 0.4 | 0.7 | |
| 1785 | 40 | 165 | 60 | 75.0 | 1.2 | 0.9 | |

1-10 of 5,517 rows | 1-8 of 23 columns

Previous 1 2 3 4 5 6 ... 100 Next

Hide

```
df<-unique(df)  
df
```

| | a..
<int> | height.cm.
<int> | weight.kg.
<int> | waist.cm.
<dbl> | eyesight.left.
<dbl> | eyesight.right.
<dbl> | hearing.lef
<int> |
|---|--------------|---------------------|---------------------|--------------------|-------------------------|--------------------------|----------------------|
| 1 | 35 | 170 | 85 | 97.0 | 0.9 | 0.9 | |
| 2 | 20 | 175 | 110 | 110.0 | 0.7 | 0.9 | |
| 3 | 45 | 155 | 65 | 86.0 | 0.9 | 0.9 | |
| 4 | 45 | 165 | 80 | 94.0 | 0.8 | 0.7 | |
| 5 | 20 | 165 | 60 | 81.0 | 1.5 | 0.1 | |
| 6 | 60 | 160 | 50 | 78.0 | 1.0 | 0.9 | |
| 7 | 40 | 175 | 90 | 95.0 | 0.9 | 1.0 | |

| | a.. | height.cm. | weight.kg. | waist.cm. | eyesight.left. | eyesight.right. | hearing.lef |
|----|-------|------------|------------|-----------|----------------|-----------------|-------------|
| | <int> | <int> | <int> | <dbl> | <dbl> | <dbl> | <int> |
| 8 | 40 | 180 | 75 | 85.0 | 1.5 | 1.5 | |
| 9 | 40 | 170 | 60 | 74.0 | 1.2 | 1.5 | |
| 10 | 45 | 155 | 55 | 78.0 | 0.7 | 1.0 | |

Hide

```
#Display the list structure
str(df)
```

```
'data.frame':  33467 obs. of  23 variables:
 $ age          : int  35 20 45 45 20 60 40 40 40 45 ...
 $ height.cm.   : int  170 175 155 165 165 160 175 180 170 155 ...
 $ weight.kg.    : int  85 110 65 80 60 50 90 75 60 55 ...
 $ waist.cm.     : num  97 110 86 94 81 78 95 85 74 78 ...
 $ eyesight.left. : num  0.9 0.7 0.9 0.8 1.5 1 0.9 1.5 1.2 0.7 ...
 $ eyesight.right. : num  0.9 0.9 0.9 0.7 0.1 0.9 1 1.5 1.5 1 ...
 $ hearing.left.  : int  1 1 1 1 1 2 1 1 1 1 ...
 $ hearing.right. : int  1 1 1 1 1 2 1 1 1 1 ...
 $ systolic      : int  118 119 110 158 109 126 130 110 89 114 ...
 $ relaxation    : int  78 79 80 88 64 75 88 60 57 81 ...
 $ fasting.blood.sugar: int  97 88 80 249 100 114 90 100 83 96 ...
 $ Cholesterol   : int  239 211 193 210 179 177 207 170 178 184 ...
 $ triglyceride  : int  153 128 120 366 200 74 331 62 69 177 ...
 $ HDL           : int  70 71 57 46 47 98 39 58 60 41 ...
 $ LDL           : int  142 114 112 91 92 64 102 99 104 107 ...
 $ hemoglobin    : num  19.8 15.9 13.7 16.9 14.9 13.9 16.5 14 12.9 13.1 ...
 $ Urine.protein  : int  1 1 3 1 1 1 1 2 2 1 ...
 $ serum.creatinine : num  1 1.1 0.6 0.9 1.2 1 1 1.4 0.7 0.6 ...
 $ AST           : int  61 19 1090 32 26 47 19 29 17 22 ...
 $ ALT           : int  115 25 1400 36 28 23 22 20 17 15 ...
 $ Gtp           : int  125 30 276 36 15 70 19 32 14 56 ...
 $ dental.caries  : int  1 1 0 0 0 0 0 1 0 0 ...
 $ smoking       : int  1 0 0 0 0 1 0 1 0 0 ...
```

Hide

```
# get number of rows and columns of data frame
dim(df)
```

```
[1] 33467    23
```

Hide

```
# check null values in dataset
colSums(is.na(df))
```

| | | | | | |
|---------|------------------|----------------|------------|---------------|--------------|
| | age | height.cm. | weight.kg. | waist.cm. | eyesigh |
| t.left. | eyesight.right. | | | | |
| 0 | 0 | 0 | 0 | 0 | |
| | 0 | | | | |
| | hearing.left. | hearing.right. | systolic | relaxation | fasting.bloo |
| d.sugar | Cholesterol | | | | |
| 0 | 0 | 0 | 0 | 0 | |
| | 0 | | | | |
| | triglyceride | HDL | LDL | hemoglobin | Urine.p |
| rotein | serum.creatinine | | | | |
| 0 | 0 | 0 | 0 | 0 | |
| | 0 | | | | |
| | AST | ALT | Gtp | dental.caries | s |
| moking | | | | | |
| 0 | 0 | 0 | 0 | 0 | |

Hide

```
summary(df)
```

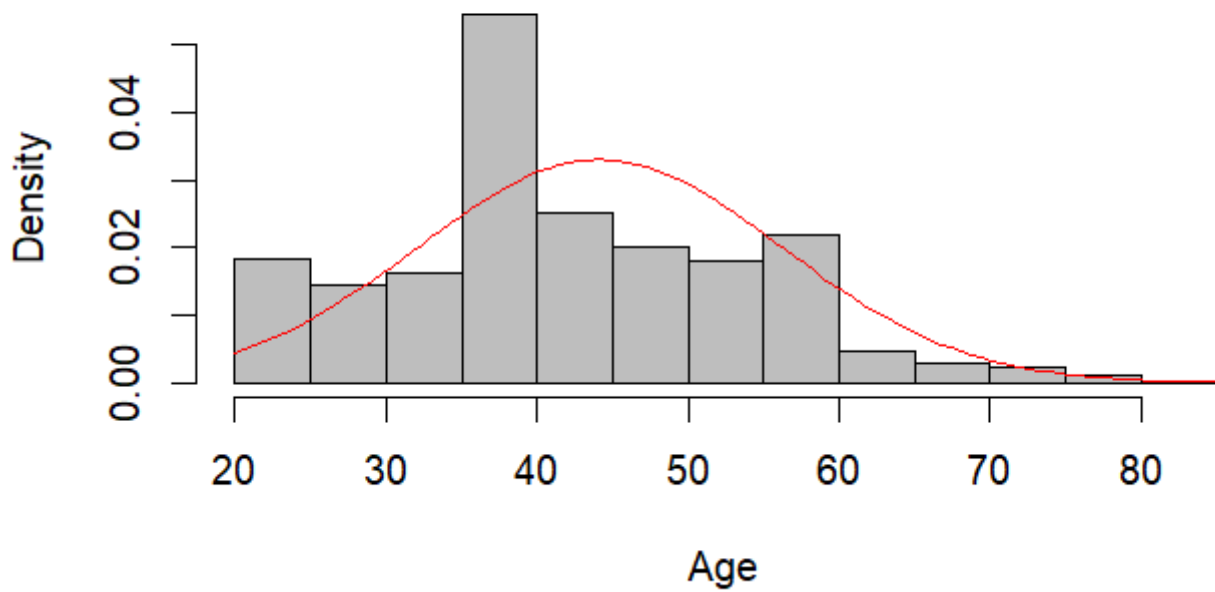

| age | height.cm. | weight.kg. | waist.cm. | eyesight.left. | eyesight.r |
|--|-----------------|----------------|----------------|----------------|------------|
| ight. hearing.left. hearing.right. | | | | | |
| Min. :20.00 | Min. :130.0 | Min. : 30.00 | Min. : 51.00 | Min. :0.100 | Min. :0. |
| 10 Min. :1.000 | Min. :1.000 | | | | |
| 1st Qu.:40.00 | 1st Qu.:160.0 | 1st Qu.: 55.00 | 1st Qu.: 76.00 | 1st Qu.:0.800 | 1st Qu.:0. |
| 80 1st Qu.:1.000 | 1st Qu.:1.000 | | | | |
| Median :40.00 | Median :165.0 | Median : 65.00 | Median : 82.00 | Median :1.000 | Median :1. |
| 00 Median :1.000 | Median :1.000 | | | | |
| Mean :44.15 | Mean :164.7 | Mean : 65.93 | Mean : 82.08 | Mean :1.014 | Mean :1. |
| 01 Mean :1.025 | Mean :1.026 | | | | |
| 3rd Qu.:55.00 | 3rd Qu.:170.0 | 3rd Qu.: 75.00 | 3rd Qu.: 88.00 | 3rd Qu.:1.200 | 3rd Qu.:1. |
| 20 3rd Qu.:1.000 | 3rd Qu.:1.000 | | | | |
| Max. :85.00 | Max. :190.0 | Max. :135.00 | Max. :129.00 | Max. :9.900 | Max. :9. |
| 90 Max. :2.000 | Max. :2.000 | | | | |
| systolic relaxation fasting.blood.sugar Cholesterol triglyceride HDL | | | | | |
| LDL hemoglobin | | | | | |
| Min. : 71.0 | Min. : 40.00 | Min. : 46.00 | Min. : 55 | Min. : 8.0 | Min. : |
| 4.00 Min. : 1.0 | Min. : 4.90 | | | | |
| 1st Qu.:112.0 | 1st Qu.: 70.00 | 1st Qu.: 89.00 | 1st Qu.:172 | 1st Qu.: 75.0 | 1st Qu.: |
| 47.00 1st Qu.: 92.0 | 1st Qu.:13.60 | | | | |
| Median :120.0 | Median : 76.00 | Median : 96.00 | Median :195 | Median :108.0 | Median : |
| 55.00 Median : 113.0 | Median :14.80 | | | | |
| Mean :121.5 | Mean : 76.02 | Mean : 99.26 | Mean :197 | Mean :126.8 | Mean : |
| 57.26 Mean : 115.2 | Mean :14.62 | | | | |
| 3rd Qu.:130.0 | 3rd Qu.: 82.00 | 3rd Qu.:104.00 | 3rd Qu.:220 | 3rd Qu.:160.0 | 3rd Qu.: |
| 66.00 3rd Qu.: 136.0 | 3rd Qu.:15.70 | | | | |
| Max. :233.0 | Max. :146.00 | Max. :423.00 | Max. :445 | Max. :999.0 | Max. :3 |
| 59.00 Max. :1860.0 | Max. :21.10 | | | | |
| Urine.protein serum.creatinine AST ALT Gtp denta | | | | | |
| l.caries smoking | | | | | |
| Min. :1.000 | Min. : 0.1000 | Min. : 6.0 | Min. : 1.00 | Min. : 2.00 | Min. |
| :0.0000 Min. :0.0000 | | | | | |
| 1st Qu.:1.000 | 1st Qu.: 0.8000 | 1st Qu.: 19.0 | 1st Qu.: 15.00 | 1st Qu.: 17.00 | 1st Q |
| u.:0.0000 1st Qu.:0.0000 | | | | | |
| Median :1.000 | Median : 0.9000 | Median : 23.0 | Median : 21.00 | Median : 26.00 | Median |
| :0.0000 Median :0.0000 | | | | | |
| Mean :1.087 | Mean : 0.8865 | Mean : 26.2 | Mean : 27.14 | Mean : 39.95 | Mean |
| :0.2147 Mean :0.3663 | | | | | |
| 3rd Qu.:1.000 | 3rd Qu.: 1.0000 | 3rd Qu.: 29.0 | 3rd Qu.: 31.00 | 3rd Qu.: 44.00 | 3rd Q |
| u.:0.0000 3rd Qu.:1.0000 | | | | | |
| Max. :6.000 | Max. :11.6000 | Max. :1090.0 | Max. :2914.00 | Max. :999.00 | Max. |
| :1.0000 Max. :1.0000 | | | | | |

#Exploratory Data Analysis (EDA)

Hide

```
hist(df$age, freq=FALSE, col="gray", xlab="Age", main = "Histogram of Age")
curve(dnorm(x, mean=mean(df$age), sd=sd(df$age)), add=TRUE, col="red") #line
```

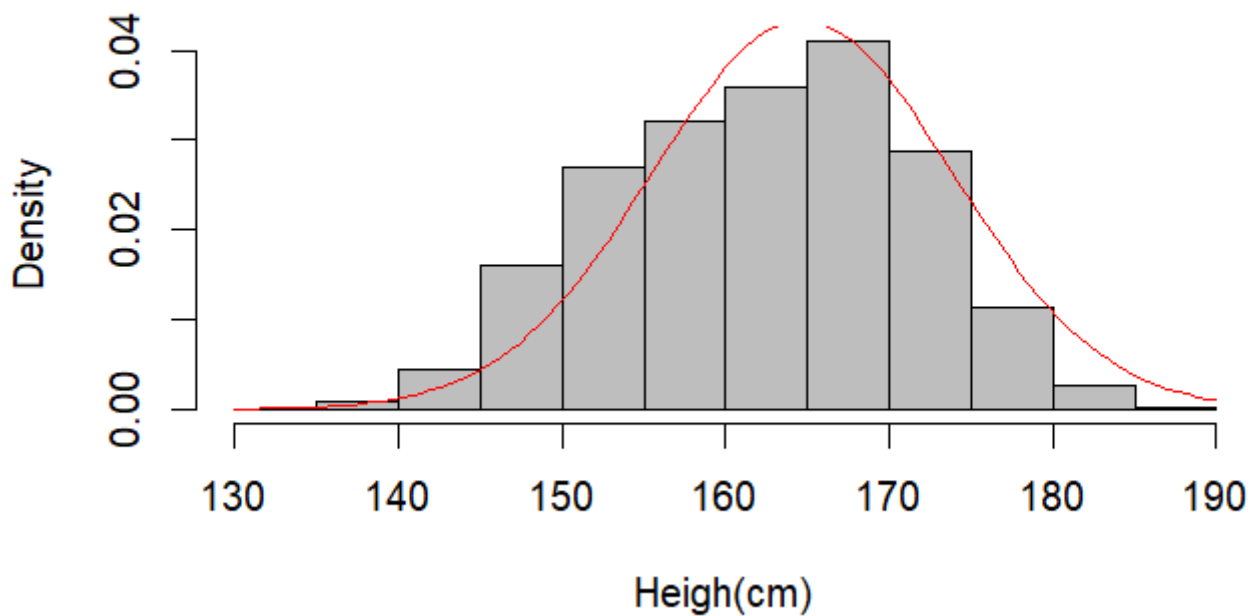
Histogram of Age



Hide

```
hist(df$height.cm., freq=FALSE, col="gray", xlab="Heigh(cm)", main = "Histogram of Height")  
curve(dnorm(x, mean=mean(df$height.cm.), sd=sd(df$height.cm.)), add=TRUE, col="red") #line
```

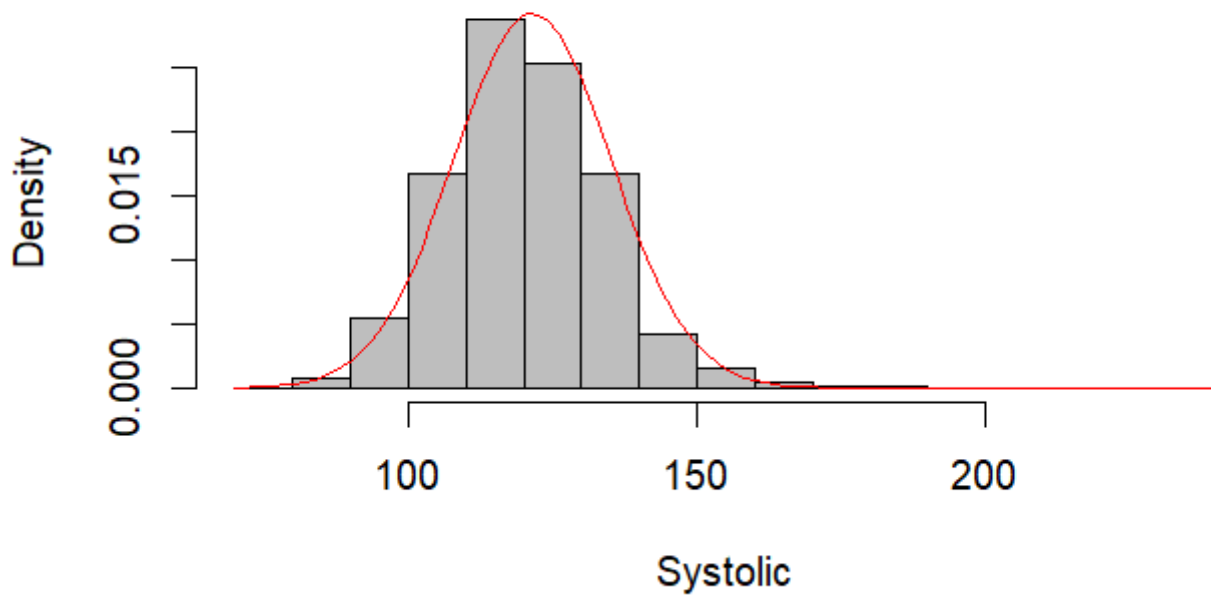
Histogram of Height



Hide

```
hist(df$systolic, freq=FALSE, col="gray", xlab="Systolic", main = "Histogram of Systolic")  
curve(dnorm(x, mean=mean(df$systolic), sd=sd(df$systolic)), add=TRUE, col="red") #line
```

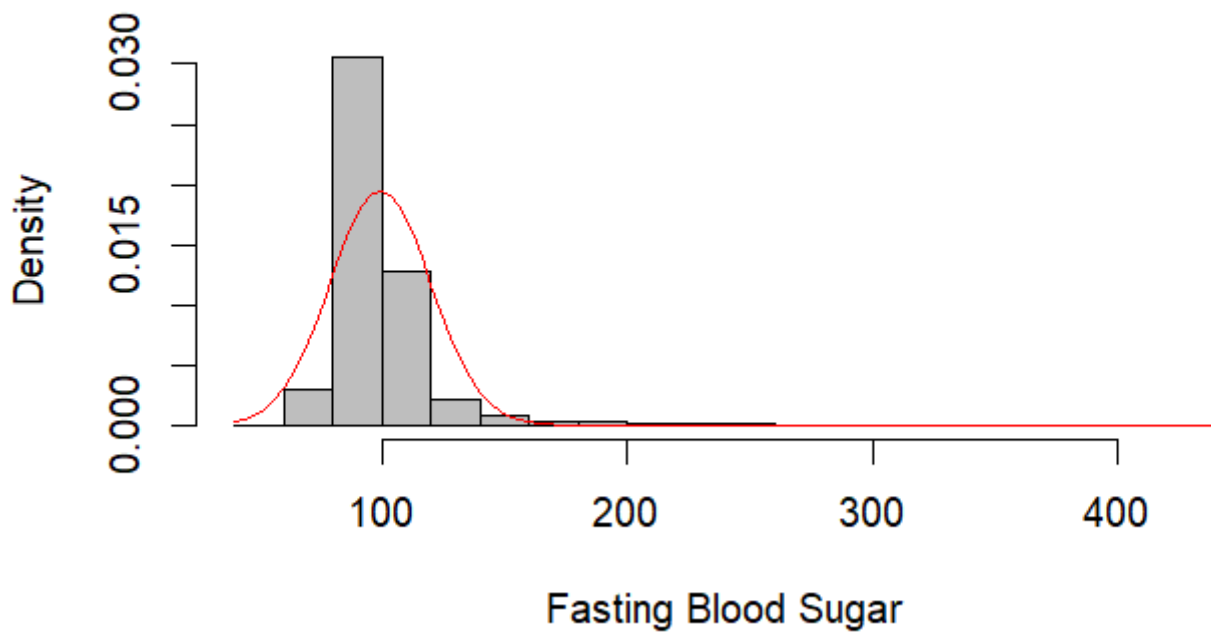
Histogram of Systolic



Hide

```
hist(df$fasting.blood.sugar, freq=FALSE, col="gray", xlab="Fasting Blood Sugar", main = "Histogram of Blood Sugar")  
curve(dnorm(x, mean=mean(df$fasting.blood.sugar), sd=sd(df$fasting.blood.sugar)), add=TRUE, col="red") #line
```

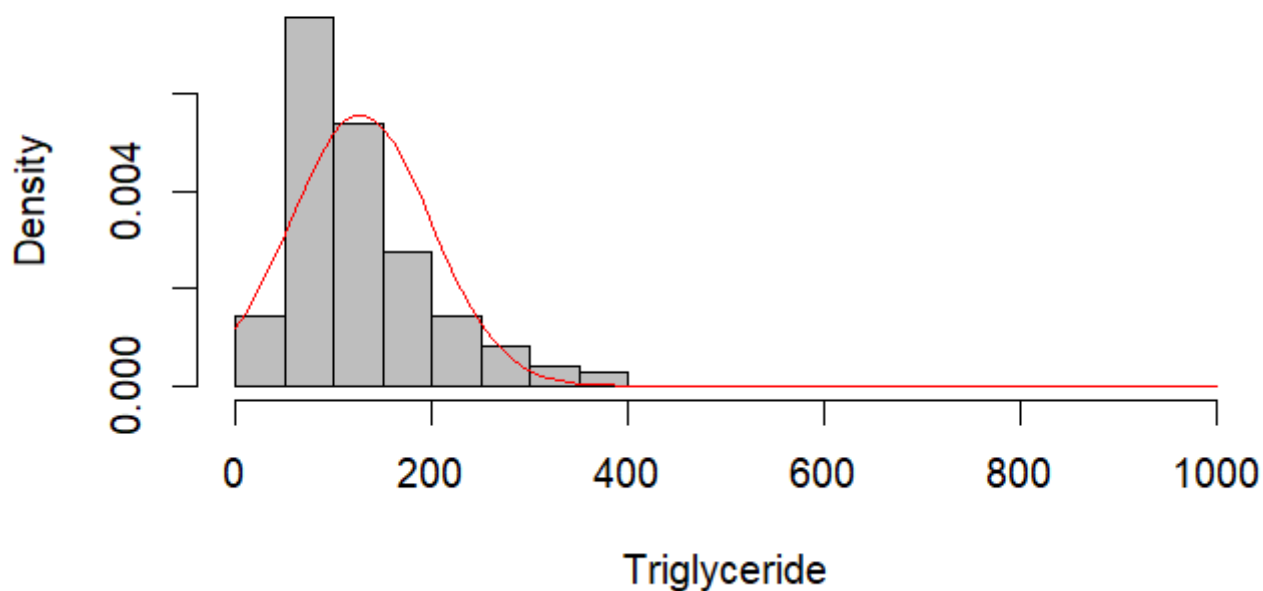
Histogram of Blood Sugar



Hide

```
hist(df$triglyceride, freq=FALSE, col="gray", xlab="Triglyceride", main = "Histogram of Triglyceride")
curve(dnorm(x, mean=mean(df$triglyceride), sd=sd(df$triglyceride)), add=TRUE, col="red") #line
```

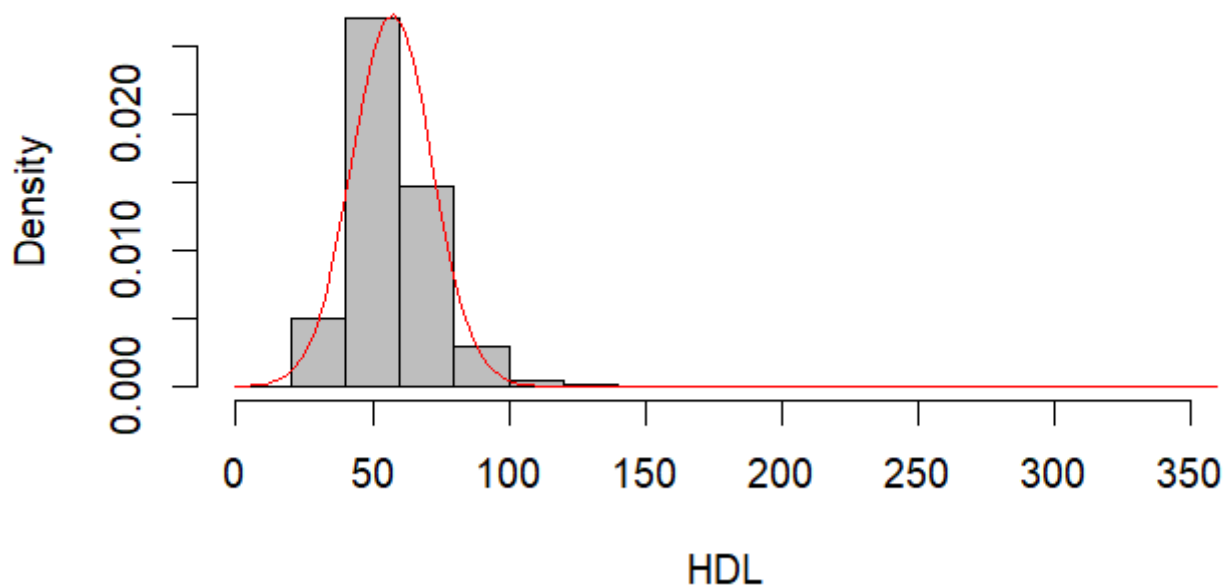
Histogram of Triglyceride



Hide

```
hist(df$HDL, freq=FALSE, col="gray", xlab="HDL", main = "Histogram of HDL")
curve(dnorm(x, mean=mean(df$HDL), sd=sd(df$HDL)), add=TRUE, col="red") #line
```

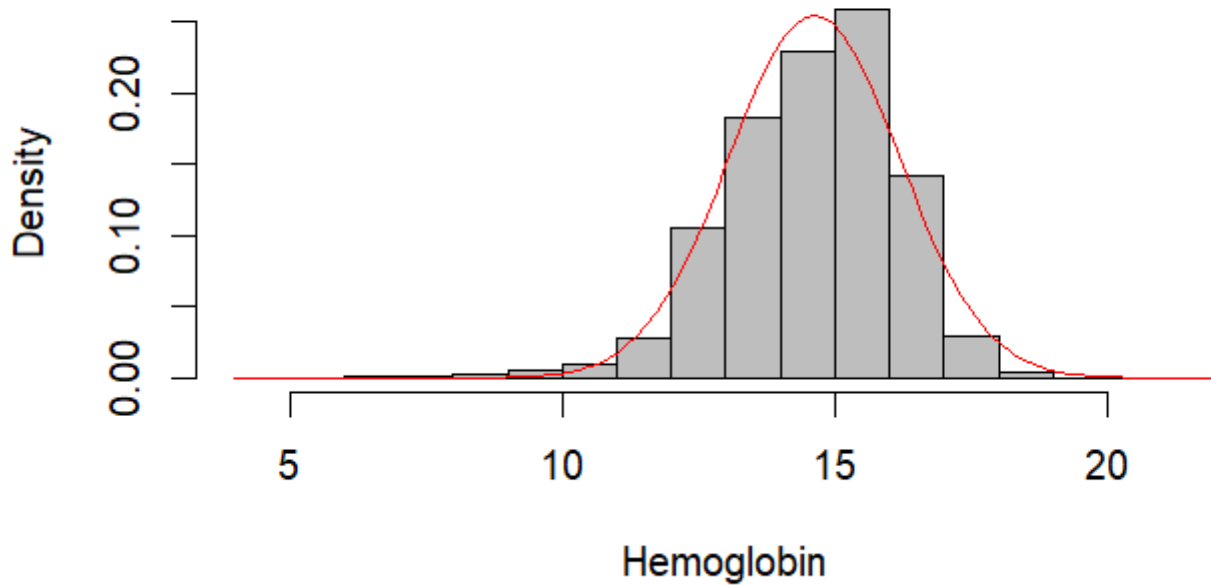
Histogram of HDL



[Hide](#)

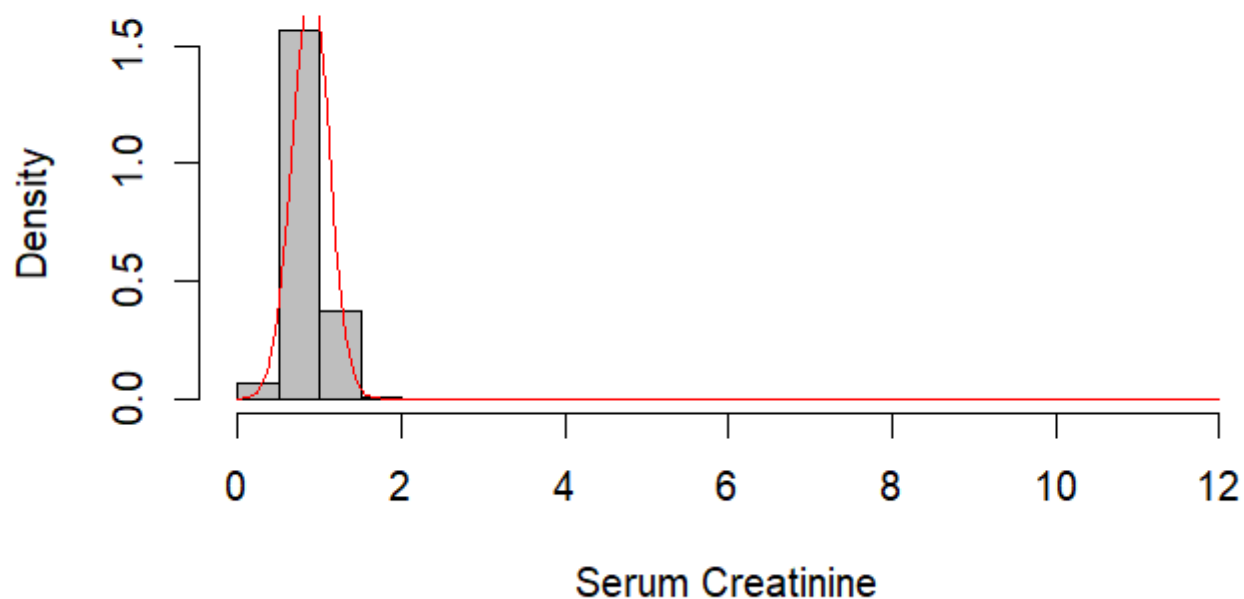
```
hist(df$hemoglobin, freq=FALSE, col="gray", xlab="Hemoglobin", main = "Histogram of Hemoglobin")  
curve(dnorm(x, mean=mean(df$hemoglobin), sd=sd(df$hemoglobin)), add=TRUE, col="red") #line
```

Histogram of Hemoglobin

[Hide](#)

```
hist(df$serum.creatinine, freq=FALSE, col="gray", xlab="Serum Creatinine", main = "Histogram of Serum Creatinine")  
curve(dnorm(x, mean=mean(df$serum.creatinine), sd=sd(df$serum.creatinine)), add=TRUE, col="red") #line
```

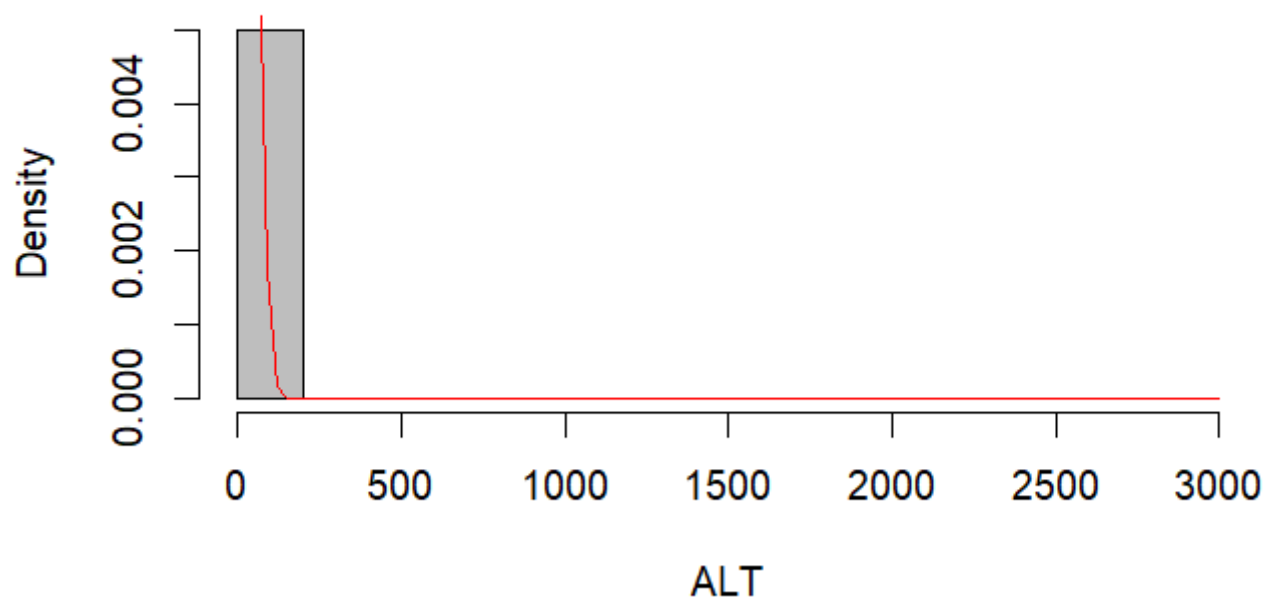
Histogram of Serum Creatinine



Hide

```
hist(df$ALT, freq=FALSE, col="gray", xlab="ALT", main = "Histogram of ALT")  
curve(dnorm(x, mean=mean(df$ALT), sd=sd(df$ALT)), add=TRUE, col="red") #line
```

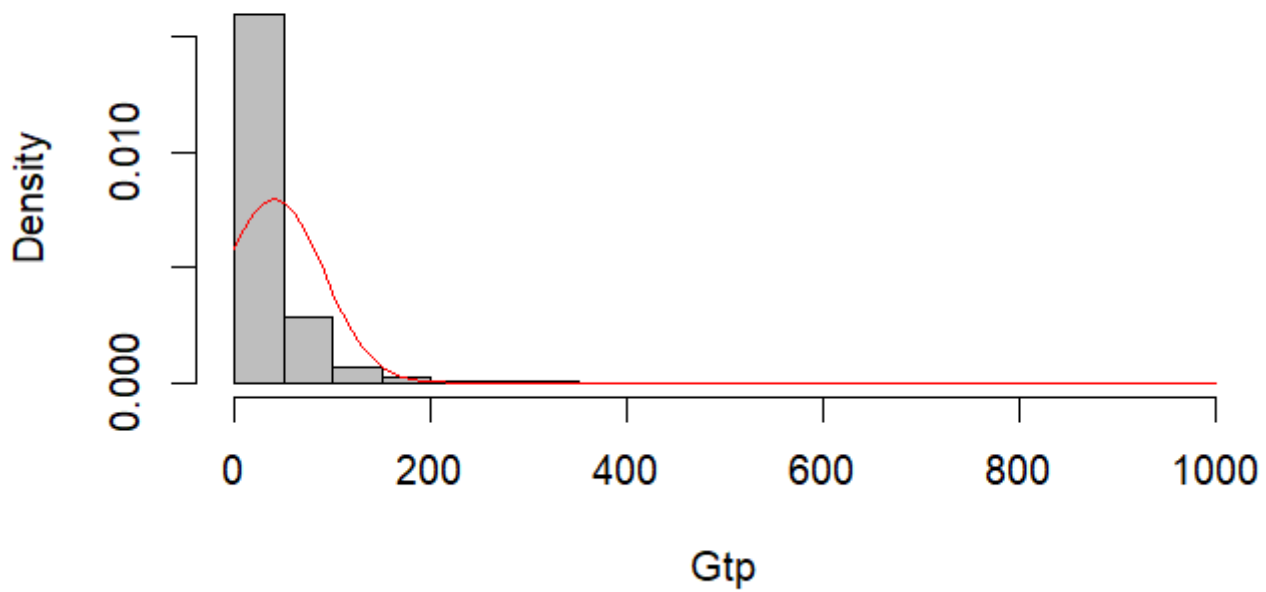
Histogram of ALT



Hide

```
hist(df$Gtp, freq=FALSE, col="gray", xlab="Gtp", main = "Histogram of Gtp")  
curve(dnorm(x, mean=mean(df$Gtp), sd=sd(df$Gtp)), add=TRUE, col="red") #line
```

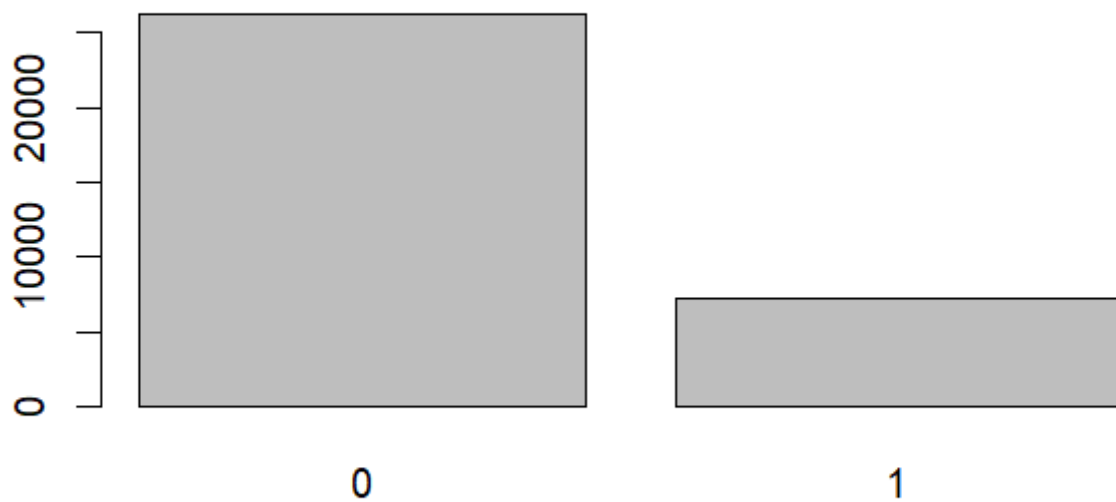
Histogram of Gtp



Hide

```
barplot(table(df$dental.caries), main = "Frequency of Dental Caries")
```

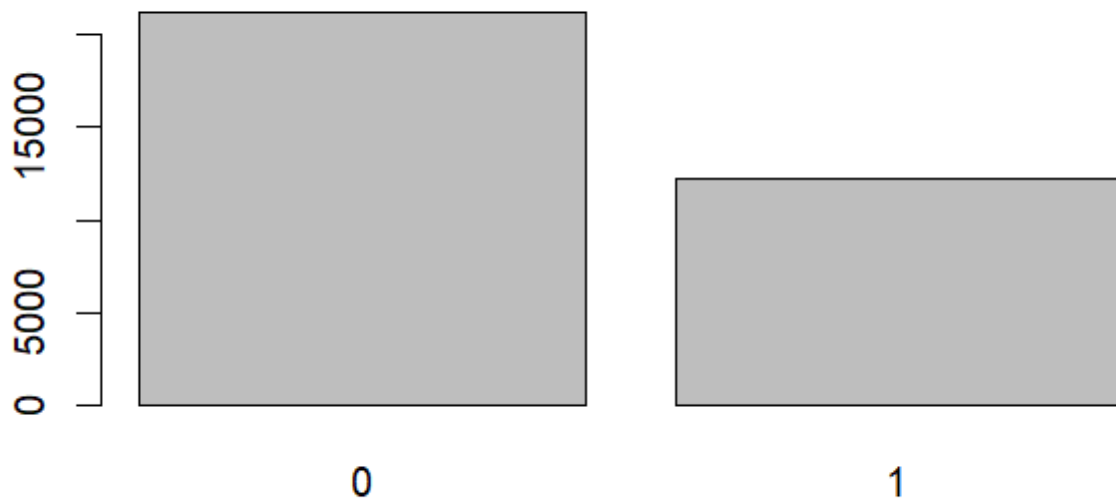
Frequency of Dental Caries



Hide

```
barplot(table(df$smoking), main = "Frequency of Smoking")
```

Frequency of Smoking

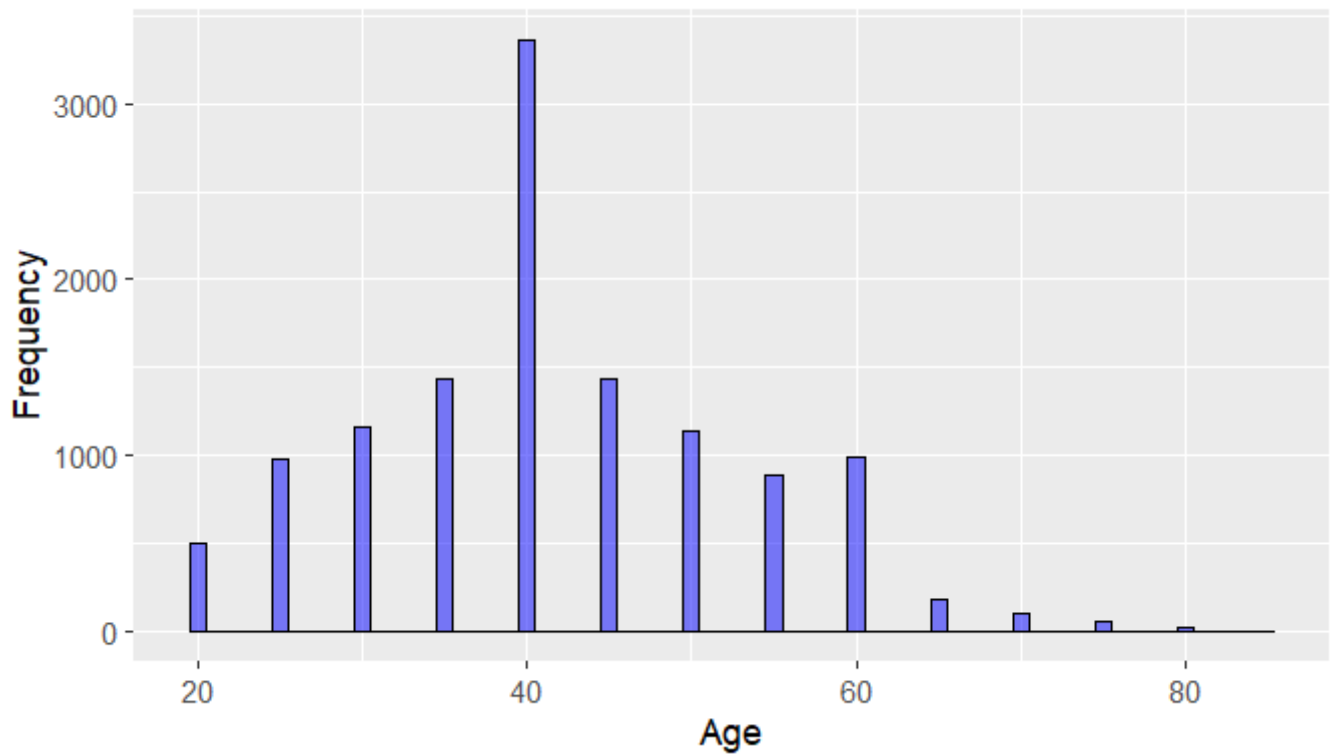


Hide

```
# Assuming you have the 'ggplot2' library installed. If not, install it using install.packages('ggplot2')
library(ggplot2)

# Histogram for smokers
ggplot(subset(df, smoking == 1), aes(x = age)) +
  geom_histogram(binwidth = 1, color = "black", fill = "blue", alpha = 0.5) +
  labs(title = "Age distribution for smokers") +
  xlab("Age") +
  ylab("Frequency")
```

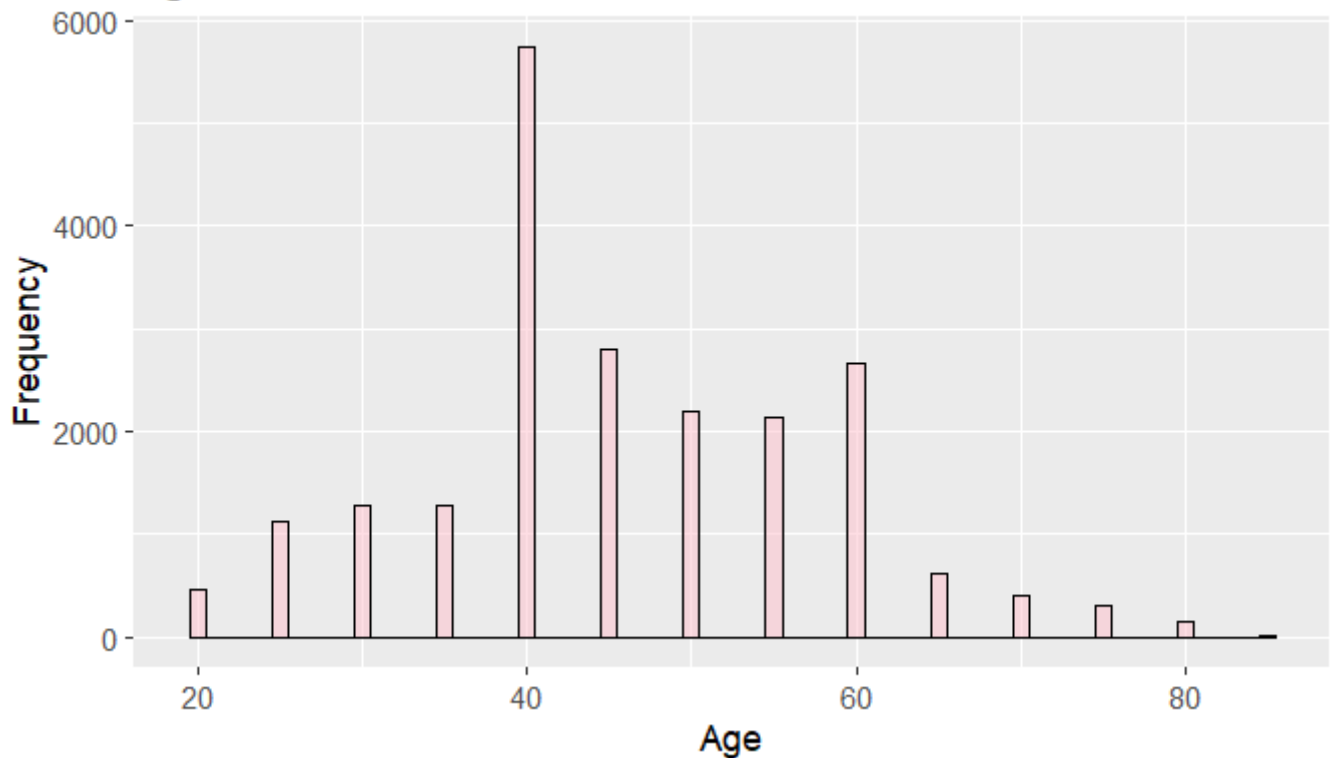

Age distribution for smokers



Hide

```
# Histogram for non-smokers
ggplot(subset(df, smoking == 0), aes(x = age)) +
  geom_histogram(binwidth = 1, color = "black", fill = "pink", alpha = 0.5) +
  labs(title = "Age distribution for non-smokers") +
  xlab("Age") +
  ylab("Frequency")
```

Age distribution for non-smokers



Based on both the histograms, the frequency of smokers and non-smokers is the highest at age 40 while the lowest is at age 80 respectively.

Hide

```
library(ggplot2)
library(plotrix)
library(scales)
# Pie chart for 'smoking'
smoking_counts <- table(df$smoking)

# Calculate percentages and format labels
percentages <- smoking_counts / sum(smoking_counts) * 100
labels_with_percentages <- paste(names(percentages), "\n", sprintf("%.1f%%", percentages))

# Plotting side by side with percentages
par(mfrow = c(1, 2), mar = c(5, 5, 2, 2))

pie(smoking_counts, labels = labels_with_percentages, col = c("red", "green"),
    main = "Percentage of Smoking Status", clockwise = TRUE,
    shadow = TRUE, explode = c(0, 0.1), percent = TRUE)
```

Percentage of Smoking Sta



Hide

```
NA
NA
```

Based on the pie chart, non-smokers have a higher percentage of smoking status of 63.4% while another 36.6% are smokers.

Hide

```
round(cor(df),4)
```

| | age | height.cm. | weight.kg. | waist.cm. | eyesight.left. | eyesight.right. | hearing.left. | hearing.right. | systolic | relaxation |
|---------------------|---------------------|-------------|--------------|-----------|----------------|-----------------|---------------|----------------|----------|------------|
| age | 1.0000 | -0.4809 | -0.3255 | -0.0271 | -0.1882 | -0.1918 | | | | |
| 0.2018 | 0.2066 | 0.1327 | 0.0484 | | | | | | | |
| height.cm. | -0.4809 | 1.0000 | 0.6744 | 0.3781 | 0.1489 | 0.1556 | | | | |
| -0.0808 | -0.0777 | 0.0820 | 0.1149 | | | | | | | |
| weight.kg. | -0.3255 | 0.6744 | 1.0000 | 0.8247 | 0.1068 | 0.1136 | | | | |
| -0.0519 | -0.0526 | 0.2683 | 0.2736 | | | | | | | |
| waist.cm. | -0.0271 | 0.3781 | 0.8247 | 1.0000 | 0.0307 | 0.0396 | | | | |
| 0.0215 | 0.0197 | 0.3193 | 0.2930 | | | | | | | |
| eyesight.left. | -0.1882 | 0.1489 | 0.1068 | 0.0307 | 1.0000 | 0.3444 | | | | |
| -0.0465 | -0.0484 | -0.0139 | 0.0072 | | | | | | | |
| eyesight.right. | -0.1918 | 0.1556 | 0.1136 | 0.0396 | 0.3444 | 1.0000 | | | | |
| -0.0374 | -0.0352 | -0.0083 | 0.0146 | | | | | | | |
| hearing.left. | 0.2018 | -0.0808 | -0.0519 | 0.0215 | -0.0465 | -0.0374 | | | | |
| 1.0000 | 0.5152 | 0.0570 | 0.0093 | | | | | | | |
| hearing.right. | 0.2066 | -0.0777 | -0.0526 | 0.0197 | -0.0484 | -0.0352 | | | | |
| 0.5152 | 1.0000 | 0.0482 | -0.0012 | | | | | | | |
| systolic | 0.1327 | 0.0820 | 0.2683 | 0.3193 | -0.0139 | -0.0083 | | | | |
| 0.0570 | 0.0482 | 1.0000 | 0.7606 | | | | | | | |
| relaxation | 0.0484 | 0.1149 | 0.2736 | 0.2930 | 0.0072 | 0.0146 | | | | |
| 0.0093 | -0.0012 | 0.7606 | 1.0000 | | | | | | | |
| fasting.blood.sugar | 0.1833 | 0.0155 | 0.1362 | 0.2138 | -0.0417 | -0.0424 | | | | |
| 0.0421 | 0.0450 | 0.1724 | 0.1496 | | | | | | | |
| Cholesterol | 0.0580 | -0.0809 | 0.0254 | 0.0635 | -0.0077 | -0.0089 | | | | |
| -0.0270 | -0.0242 | 0.0596 | 0.0947 | | | | | | | |
| triglyceride | 0.0147 | 0.1576 | 0.3228 | 0.3595 | 0.0225 | 0.0257 | | | | |
| 0.0047 | 0.0015 | 0.1971 | 0.2153 | | | | | | | |
| HDL | 0.0085 | -0.2143 | -0.3607 | -0.3762 | -0.0209 | -0.0239 | | | | |
| -0.0196 | -0.0144 | -0.0842 | -0.0884 | | | | | | | |
| LDL | 0.0422 | -0.0465 | 0.0368 | 0.0672 | -0.0093 | -0.0091 | | | | |
| -0.0193 | -0.0179 | 0.0136 | 0.0358 | | | | | | | |
| hemoglobin | -0.2652 | 0.5367 | 0.4888 | 0.3804 | 0.0956 | 0.0963 | | | | |
| -0.0319 | -0.0318 | 0.1851 | 0.2347 | | | | | | | |
| Urine.protein | 0.0273 | 0.0060 | 0.0284 | 0.0423 | -0.0088 | -0.0118 | | | | |
| 0.0168 | 0.0179 | 0.0480 | 0.0521 | | | | | | | |
| serum.creatinine | -0.1035 | 0.3801 | 0.3175 | 0.2272 | 0.0578 | 0.0464 | | | | |
| 0.0022 | 0.0105 | 0.0732 | 0.0879 | | | | | | | |
| AST | 0.0248 | 0.0498 | 0.1289 | 0.1463 | -0.0057 | -0.0021 | | | | |
| 0.0079 | 0.0100 | 0.0863 | 0.0865 | | | | | | | |
| ALT | -0.0681 | 0.1315 | 0.2513 | 0.2504 | 0.0216 | 0.0276 | | | | |
| 0.0004 | 0.0014 | 0.0935 | 0.1039 | | | | | | | |
| Gtp | 0.0113 | 0.1383 | 0.2087 | 0.2403 | 0.0029 | 0.0171 | | | | |
| 0.0084 | 0.0050 | 0.1714 | 0.1786 | | | | | | | |
| dental.caries | -0.1145 | 0.0829 | 0.0767 | 0.0476 | 0.0077 | 0.0169 | | | | |
| -0.0186 | -0.0165 | 0.0344 | 0.0363 | | | | | | | |
| smoking | -0.1671 | 0.3934 | 0.2981 | 0.2228 | 0.0601 | 0.0620 | | | | |
| -0.0233 | -0.0181 | 0.0704 | 0.1045 | | | | | | | |
| | fasting.blood.sugar | Cholesterol | triglyceride | HDL | LDL | hemoglobin | U | | | |
| rine.protein | serum.creatinine | AST | ALT | | | | | | | |
| age | | 0.1833 | 0.0580 | 0.0147 | 0.0085 | 0.0422 | -0.2652 | | | |
| 0.0273 | -0.1035 | 0.0248 | -0.0681 | | | | | | | |
| height.cm. | | 0.0155 | -0.0809 | 0.1576 | -0.2143 | -0.0465 | 0.5367 | | | |
| 0.0060 | 0.3801 | 0.0498 | 0.1315 | | | | | | | |
| weight.kg. | | 0.1362 | 0.0254 | 0.3228 | -0.3607 | 0.0368 | 0.4888 | | | |

| | | | | | | | | |
|---------------------|---------|---------|---------------|---------|---------|---------|---------|---------|
| 0.0284 | 0.3175 | 0.1289 | 0.2513 | | | | | |
| waist.cm. | | | 0.2138 | 0.0635 | 0.3595 | -0.3762 | 0.0672 | 0.3804 |
| 0.0423 | 0.2272 | 0.1463 | 0.2504 | | | | | |
| eyesight.left. | | | -0.0417 | -0.0077 | 0.0225 | -0.0209 | -0.0093 | 0.0956 |
| -0.0088 | 0.0578 | -0.0057 | 0.0216 | | | | | |
| eyesight.right. | | | -0.0424 | -0.0089 | 0.0257 | -0.0239 | -0.0091 | 0.0963 |
| -0.0118 | 0.0464 | -0.0021 | 0.0276 | | | | | |
| hearing.left. | | | 0.0421 | -0.0270 | 0.0047 | -0.0196 | -0.0193 | -0.0319 |
| 0.0168 | 0.0022 | 0.0079 | 0.0004 | | | | | |
| hearing.right. | | | 0.0450 | -0.0242 | 0.0015 | -0.0144 | -0.0179 | -0.0318 |
| 0.0179 | 0.0105 | 0.0100 | 0.0014 | | | | | |
| systolic | | | 0.1724 | 0.0596 | 0.1971 | -0.0842 | 0.0136 | 0.1851 |
| 0.0480 | 0.0732 | 0.0863 | 0.0935 | | | | | |
| relaxation | | | 0.1496 | 0.0947 | 0.2153 | -0.0884 | 0.0358 | 0.2347 |
| 0.0521 | 0.0879 | 0.0865 | 0.1039 | | | | | |
| fasting.blood.sugar | | | 1.0000 | 0.0091 | 0.2264 | -0.1222 | -0.0185 | 0.0985 |
| 0.1055 | 0.0433 | 0.0693 | 0.0888 | | | | | |
| Cholesterol | | | 0.0091 | 1.0000 | 0.2436 | 0.1684 | 0.7066 | 0.0670 |
| -0.0012 | 0.0063 | 0.0254 | 0.0473 | | | | | |
| triglyceride | | | 0.2264 | 0.2436 | 1.0000 | -0.4153 | 0.0221 | 0.2724 |
| 0.0330 | 0.1234 | 0.1037 | 0.1744 | | | | | |
| HDL | | | -0.1222 | 0.1684 | -0.4153 | 1.0000 | -0.0560 | -0.2409 |
| -0.0085 | -0.1719 | -0.0362 | -0.1298 | | | | | |
| LDL | | | -0.0185 | 0.7066 | 0.0221 | -0.0560 | 1.0000 | 0.0548 |
| -0.0086 | 0.0319 | -0.0017 | 0.0277 | | | | | |
| hemoglobin | | | 0.0985 | 0.0670 | 0.2724 | -0.2409 | 0.0548 | 1.0000 |
| 0.0238 | 0.3661 | 0.1239 | 0.1992 | | | | | |
| Urine.protein | | | 0.1055 | -0.0012 | 0.0330 | -0.0085 | -0.0086 | 0.0238 |
| 1.0000 | 0.0862 | 0.0539 | 0.0387 | | | | | |
| serum.creatinine | | | 0.0433 | 0.0063 | 0.1234 | -0.1719 | 0.0319 | 0.3661 |
| 0.0862 | 1.0000 | 0.0516 | 0.0861 | | | | | |
| AST | | | 0.0693 | 0.0254 | 0.1037 | -0.0362 | -0.0017 | 0.1239 |
| 0.0539 | 0.0516 | 1.0000 | 0.6671 | | | | | |
| ALT | | | 0.0888 | 0.0473 | 0.1744 | -0.1298 | 0.0277 | 0.1992 |
| 0.0387 | 0.0861 | 0.6671 | 1.0000 | | | | | |
| Gtp | | | 0.1801 | 0.0835 | 0.2988 | -0.0480 | -0.0088 | 0.2228 |
| 0.0706 | 0.1112 | 0.3790 | 0.3307 | | | | | |
| dental.caries | | | -0.0022 | -0.0004 | 0.0296 | -0.0273 | -0.0002 | 0.0697 |
| -0.0010 | 0.0316 | 0.0131 | 0.0259 | | | | | |
| smoking | | | 0.0978 | -0.0296 | 0.2495 | -0.1774 | -0.0420 | 0.3983 |
| 0.0134 | 0.2090 | 0.0635 | 0.0973 | | | | | |
| | | Gtp | dental.caries | smoking | | | | |
| age | 0.0113 | | -0.1145 | -0.1671 | | | | |
| height.cm. | 0.1383 | | 0.0829 | 0.3934 | | | | |
| weight.kg. | 0.2087 | | 0.0767 | 0.2981 | | | | |
| waist.cm. | 0.2403 | | 0.0476 | 0.2228 | | | | |
| eyesight.left. | 0.0029 | | 0.0077 | 0.0601 | | | | |
| eyesight.right. | 0.0171 | | 0.0169 | 0.0620 | | | | |
| hearing.left. | 0.0084 | | -0.0186 | -0.0233 | | | | |
| hearing.right. | 0.0050 | | -0.0165 | -0.0181 | | | | |
| systolic | 0.1714 | | 0.0344 | 0.0704 | | | | |
| relaxation | 0.1786 | | 0.0363 | 0.1045 | | | | |
| fasting.blood.sugar | 0.1801 | | -0.0022 | 0.0978 | | | | |
| Cholesterol | 0.0835 | | -0.0004 | -0.0296 | | | | |
| triglyceride | 0.2988 | | 0.0296 | 0.2495 | | | | |
| HDL | -0.0480 | | -0.0273 | -0.1774 | | | | |

| | | | |
|------------------|---------|---------|---------|
| LDL | -0.0088 | -0.0002 | -0.0420 |
| hemoglobin | 0.2228 | 0.0697 | 0.3983 |
| Urine.protein | 0.0706 | -0.0010 | 0.0134 |
| serum.creatinine | 0.1112 | 0.0316 | 0.2090 |
| AST | 0.3790 | 0.0131 | 0.0635 |
| ALT | 0.3307 | 0.0259 | 0.0973 |
| Gtp | 1.0000 | 0.0454 | 0.2380 |
| dental.caries | 0.0454 | 1.0000 | 0.1064 |
| smoking | 0.2380 | 0.1064 | 1.0000 |

Based on the correlation shown above, the variables that have positive correlation with response variables smoking are height.cm., weight.kg., waist.cm., triglyceride, hemoglobin, serum.creatinine and Gtp.

Hide

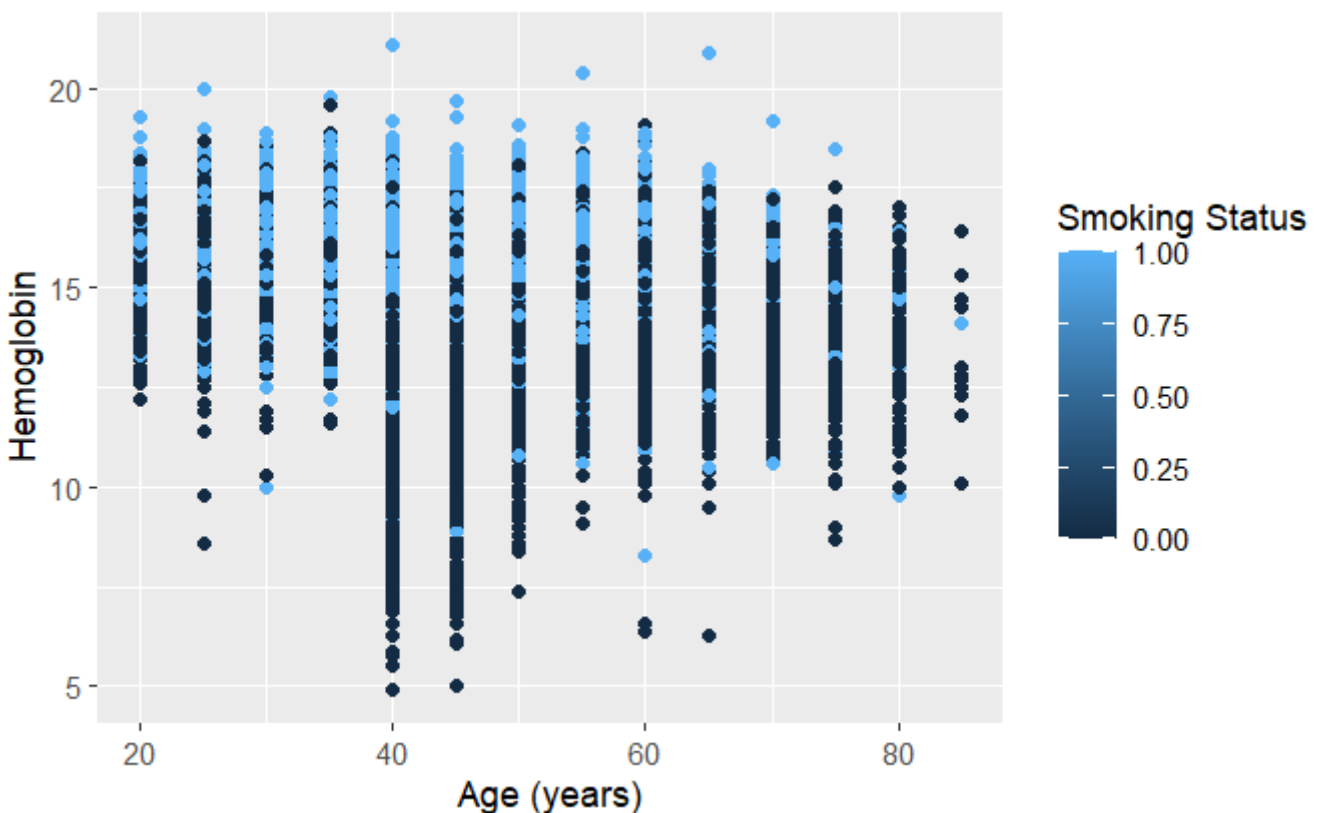
```
#scatter plot
# Load packages
library(tidyverse)

# Load data
data(df)
```

Warning: data set 'df' not found

Hide

```
# Create a scatter plot of mpg, disp, and cyl
# Create a scatter plot of age, height, and smoking
ggplot(df, aes(x = age, y = hemoglobin, color = smoking)) +
  geom_point() +
  labs(x = "Age (years)", y = "Hemoglobin", color = "Smoking Status")
```



Based on this scatter plot, we can see that the light blue colour represents the person who is a smoker and the darker blue colour represents the person who is not a smoker. Based on the scatter plot, people who are older are likely not a smoker. Moreover, people who smoke have an overly high level of hemoglobin. This is because the carbon monoxide in cigarette smoke blocks oxygen attachment to the red cells' empty hemoglobin slots, causing the body to increase red blood cell production.

Hide

```
# Load packages
library(tidyverse)
library(corrplot)

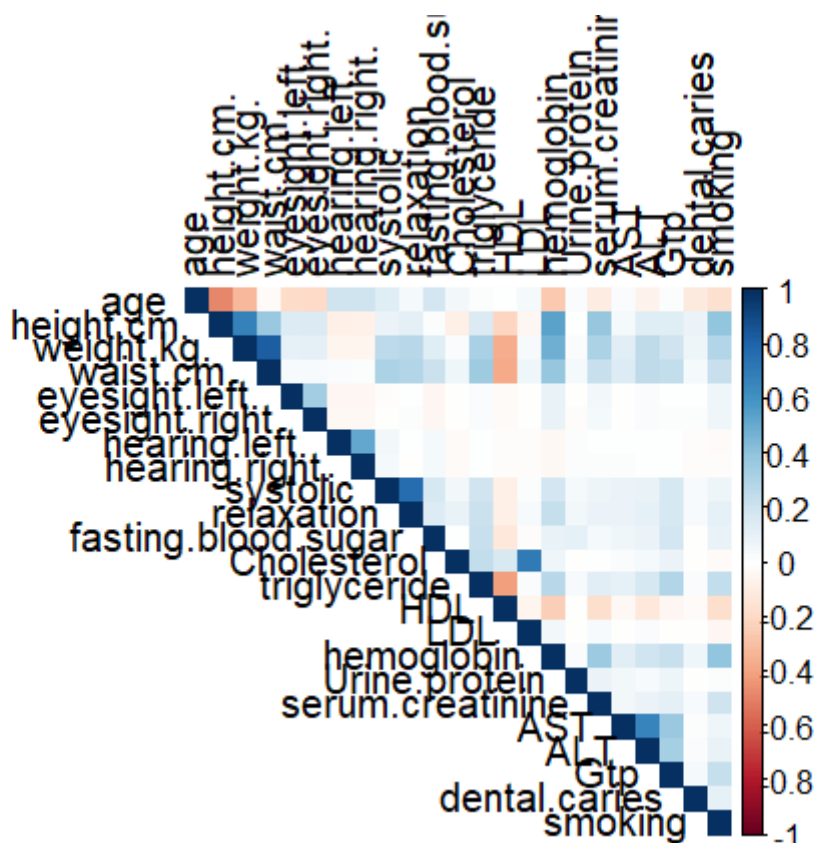
# Load data (Replace 'df' with your actual dataframe)
data(df)
```

Warning: data set 'df' not found

Hide

```
# Calculate correlation matrix
cor_mat <- cor(df)

# Plot correlation matrix with larger size
corrplot(cor_mat, method = "color", type = "upper", tl.col = "black", tl.srt = 90)
```



Hide

NA

NA

From the heatmap, the darker blue colour represents the strong correlation between the variables. There appears to be a number that shows a negative number, which means that when the number is negative, the data has a relationship that is rather than reversed.

If the number is approaching zero, the variables do not correlate at all.

Based on the heatmap, when investigating the correlation between the response variable smoking and the explanatory variables, the explanatory variables that have a weak positive correlation with the highest to the response variable smoking among the attributes in this dataset are height.cm., weight.kg., waist.cm., triglyceride, hemoglobin, serum.creatinine and Gtp.

a. Investigate the basic model obtained from the data set.

Hide

```
logreg <- glm(formula = smoking ~ ., family = binomial(link="logit"), data = df)
summary(logreg)
```

Call:

```
glm(formula = smoking ~ ., family = binomial(link = "logit"),  
     data = df)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|---------|---------|---------|--------|--------|
| -4.3034 | -0.8208 | -0.3947 | 0.9515 | 4.7542 |

Coefficients:

| | Estimate | Std. Error | z value | Pr(> z) | |
|---------------------|------------|------------|---------|----------|-----|
| (Intercept) | -1.993e+01 | 4.679e-01 | -42.589 | < 2e-16 | *** |
| age | 8.450e-04 | 1.399e-03 | 0.604 | 0.54593 | |
| height.cm. | 8.282e-02 | 2.511e-03 | 32.974 | < 2e-16 | *** |
| weight.kg. | -2.441e-02 | 2.685e-03 | -9.093 | < 2e-16 | *** |
| waist.cm. | 1.453e-02 | 3.140e-03 | 4.627 | 3.72e-06 | *** |
| eyesight.left. | 2.359e-02 | 2.800e-02 | 0.842 | 0.39954 | |
| eyesight.right. | 1.786e-03 | 2.797e-02 | 0.064 | 0.94910 | |
| hearing.left. | -9.824e-02 | 1.030e-01 | -0.954 | 0.34018 | |
| hearing.right. | 1.169e-01 | 1.001e-01 | 1.168 | 0.24273 | |
| systolic | -9.436e-03 | 1.564e-03 | -6.032 | 1.62e-09 | *** |
| relaxation | 4.526e-03 | 2.154e-03 | 2.102 | 0.03559 | * |
| fasting.blood.sugar | 4.174e-03 | 6.930e-04 | 6.024 | 1.71e-09 | *** |
| Cholesterol | -5.055e-03 | 6.157e-04 | -8.210 | < 2e-16 | *** |
| triglyceride | 4.566e-03 | 2.476e-04 | 18.442 | < 2e-16 | *** |
| HDL | -3.344e-03 | 1.244e-03 | -2.689 | 0.00717 | ** |
| LDL | 2.828e-04 | 4.641e-04 | 0.609 | 0.54223 | |
| hemoglobin | 4.400e-01 | 1.268e-02 | 34.715 | < 2e-16 | *** |
| Urine.protein | -6.612e-02 | 3.372e-02 | -1.961 | 0.04988 | * |
| serum.creatinine | 1.755e-01 | 6.809e-02 | 2.578 | 0.00994 | ** |
| AST | -3.490e-04 | 1.282e-03 | -0.272 | 0.78549 | |
| ALT | -6.837e-03 | 9.982e-04 | -6.849 | 7.45e-12 | *** |
| Gtp | 9.747e-03 | 4.346e-04 | 22.428 | < 2e-16 | *** |
| dental.caries | 4.104e-01 | 3.157e-02 | 13.000 | < 2e-16 | *** |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for binomial family taken to be 1)

Null deviance: 43972 on 33466 degrees of freedom

Residual deviance: 34100 on 33444 degrees of freedom

AIC: 34146

Number of Fisher Scoring iterations: 5

Hide

```
null_model <- glm(formula = smoking ~ 1, family = binomial(link="logit"), data = df)  
summary(null_model)
```



```
Call:
glm(formula = smoking ~ 1, family = binomial(link = "logit"),
     data = df)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|---------|---------|---------|--------|--------|
| -0.9551 | -0.9551 | -0.9551 | 1.4173 | 1.4173 |

Coefficients:

| | Estimate | Std. Error | z value | Pr(> z) |
|-------------|----------|------------|---------|------------|
| (Intercept) | -0.54825 | 0.01135 | -48.32 | <2e-16 *** |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for binomial family taken to be 1)

Null deviance: 43972 on 33466 degrees of freedom
Residual deviance: 43972 on 33466 degrees of freedom
AIC: 43974

Number of Fisher Scoring iterations: 4

Hide

```
#Install and load the 'car' package (if not already installed)
#install.packages("car")
library(car)

# Assuming 'lm_model' is your linear regression model
# Make sure your model is fitted before calculating VIF

# Calculate VIF
vif_values <- car::vif(logreg)

# Print the VIF values
print(vif_values)
```

| | | | | | |
|-----------------|---------------|---------------|---------------------|-----------|---------|
| t.left. | age | height.cm. | weight.kg. | waist.cm. | eyesigh |
| eyesight.right. | hearing.left. | | | | |
| 1.687685 | 2.273006 | 6.248744 | 4.453792 | 1. | |
| 137716 | 1.139278 | 1.358622 | | | |
| hearing.right. | systolic | relaxation | fasting.blood.sugar | Chole | |
| sterol | triglyceride | HDL | | | |
| 1.363709 | 2.447715 | 2.395523 | 1.139308 | 2. | |
| 808775 | 1.766252 | 1.694096 | | | |
| LDL | hemoglobin | Urine.protein | serum.creatinine | | |
| AST | ALT | Gtp | | | |
| 2.359566 | 1.389577 | 1.022667 | 1.161300 | 2. | |
| 275334 | 2.791200 | 1.455968 | | | |
| dental.caries | | | | | |
| 1.012046 | | | | | |

```
anova(logreg, test="Chisq")
```

Analysis of Deviance Table

Model: binomial, link: logit

Response: smoking

Terms added sequentially (first to last)

| | Df | Deviance | Resid. Df | Resid. Dev | Pr(>Chi) |
|---|----|----------|-----------|------------|---------------|
| NULL | | | 33466 | 43972 | |
| age | 1 | 956.1 | 33465 | 43016 | < 2.2e-16 *** |
| height.cm. | 1 | 4698.2 | 33464 | 38317 | < 2.2e-16 *** |
| weight.kg. | 1 | 55.0 | 33463 | 38262 | 1.218e-13 *** |
| waist.cm. | 1 | 233.9 | 33462 | 38029 | < 2.2e-16 *** |
| eyesight.left. | 1 | 2.2 | 33461 | 38026 | 0.1421187 |
| eyesight.right. | 1 | 0.5 | 33460 | 38026 | 0.4785978 |
| hearing.left. | 1 | 0.0 | 33459 | 38026 | 0.8934461 |
| hearing.right. | 1 | 1.0 | 33458 | 38025 | 0.3084187 |
| systolic | 1 | 11.9 | 33457 | 38013 | 0.0005614 *** |
| relaxation | 1 | 84.9 | 33456 | 37928 | < 2.2e-16 *** |
| fasting.blood.sugar | 1 | 220.6 | 33455 | 37707 | < 2.2e-16 *** |
| Cholesterol | 1 | 2.8 | 33454 | 37705 | 0.0919096 . |
| triglyceride | 1 | 1099.1 | 33453 | 36606 | < 2.2e-16 *** |
| HDL | 1 | 5.6 | 33452 | 36600 | 0.0178052 * |
| LDL | 1 | 1.0 | 33451 | 36599 | 0.3084939 |
| hemoglobin | 1 | 1617.7 | 33450 | 34981 | < 2.2e-16 *** |
| Urine.protein | 1 | 0.6 | 33449 | 34981 | 0.4505911 |
| serum.creatinine | 1 | 11.7 | 33448 | 34969 | 0.0006299 *** |
| AST | 1 | 2.5 | 33447 | 34966 | 0.1104928 |
| ALT | 1 | 6.6 | 33446 | 34960 | 0.0099481 ** |
| Gtp | 1 | 691.0 | 33445 | 34269 | < 2.2e-16 *** |
| dental.caries | 1 | 168.7 | 33444 | 34100 | < 2.2e-16 *** |
| --- | | | | | |
| Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1 | | | | | |

Results & Interpretation

Significance Codes: ‘*’: **Very highly significant (p-value < 0.001)**. ‘’: **Highly significant (p-value < 0.01)**. ‘.’: Significant at a 5% level (p-value < 0.05). ‘ ’: Not significant (p-value > 0.1).

Results from the full logistic regression model show that there are some insignificant variables such as “age”, “eyesight.left.”, “eyesight.right.”, “hearing.left.”, “hearing.right.”, “Cholesterol”, “LDL”, “AST” and “relaxation” based on its respective p-values.

Variance Inflation Factors (VIF).

The Variance Inflation Factor (VIF) measures the inflation in the coefficient of the independent variable due to the collinearities among the other independent variables.

A VIF of 1 means that the regression coefficient is not inflated by the presence of the other predictors, and hence multicollinearity does not exist.

Ideally, the Variance Inflation Factors are below 5.

Results from multicollinearity with VIF test shows that weight.kg. (VIF: 6.23): This variable has a relatively high VIF, indicating that its variance is inflated due to its correlation with other predictors and waist.cm. (VIF: 4.43): While this VIF is above 2, it is not extremely high. However, it suggests some correlation with other predictors.

Results from the ANOVA test show that “eyesight.left.”, “eyesight.right.”, “hearing.left.”, “hearing.right.”, “Cholesterol”, “LDL”, “Urine.protein”, and “AST” are insignificant to the fitted model.

Hence, fit the updated glm() model (logistic regression) as a reduced model without the insignificant variables and non-collinear variables into logreg2 in order to develop the best model in our case study.

b. Develop the best model as your solution based on this course.

Hide

```
### Fit the updated glm() model (logistic regression) into logreg2.
logreg2 <- glm(formula = smoking ~ age + height.cm. + systolic +
               fasting.blood.sugar + triglyceride +
               HDL + hemoglobin + serum.creatinine + ALT + Gtp + dental.caries,
               family = binomial(link="logit"),
               data = df)

summary(logreg2)
```

Call:

```
glm(formula = smoking ~ age + height.cm. + systolic + fasting.blood.sugar +  
    triglyceride + HDL + hemoglobin + serum.creatinine + ALT +  
    Gtp + dental.caries, family = binomial(link = "logit"), data = df)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|---------|---------|---------|--------|--------|
| -4.3039 | -0.8216 | -0.4068 | 0.9589 | 5.4841 |

Coefficients:

| | Estimate | Std. Error | z value | Pr(> z) | |
|---------------------|------------|------------|---------|----------|-----|
| (Intercept) | -1.849e+01 | 3.781e-01 | -48.888 | < 2e-16 | *** |
| age | 3.241e-03 | 1.281e-03 | 2.529 | 0.0114 | * |
| height.cm. | 7.032e-02 | 2.030e-03 | 34.636 | < 2e-16 | *** |
| systolic | -9.216e-03 | 1.054e-03 | -8.745 | < 2e-16 | *** |
| fasting.blood.sugar | 4.213e-03 | 6.854e-04 | 6.148 | 7.87e-10 | *** |
| triglyceride | 3.474e-03 | 2.146e-04 | 16.185 | < 2e-16 | *** |
| HDL | -5.853e-03 | 1.075e-03 | -5.447 | 5.13e-08 | *** |
| hemoglobin | 4.278e-01 | 1.248e-02 | 34.268 | < 2e-16 | *** |
| serum.creatinine | 9.990e-02 | 6.859e-02 | 1.456 | 0.1453 | |
| ALT | -8.969e-03 | 7.179e-04 | -12.493 | < 2e-16 | *** |
| Gtp | 9.903e-03 | 4.288e-04 | 23.097 | < 2e-16 | *** |
| dental.caries | 4.089e-01 | 3.144e-02 | 13.008 | < 2e-16 | *** |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for binomial family taken to be 1)

Null deviance: 43972 on 33466 degrees of freedom
Residual deviance: 34369 on 33455 degrees of freedom
AIC: 34393

Number of Fisher Scoring iterations: 5

Hide

```
### Use the anova() function to analyze the updated table of deviance.  
anova(logreg2, test="Chisq")
```

Analysis of Deviance Table

Model: binomial, link: logit

Response: smoking

Terms added sequentially (first to last)

| | Df | Deviance | Resid. Df | Resid. Dev | Pr(>Chi) |
|---------------------|----|----------|-----------|------------|-----------|
| NULL | | | 33466 | 43972 | |
| age | 1 | 956.1 | 33465 | 43016 | < 2.2e-16 |
| height.cm. | 1 | 4698.2 | 33464 | 38317 | < 2.2e-16 |
| systolic | 1 | 40.7 | 33463 | 38277 | 1.788e-10 |
| fasting.blood.sugar | 1 | 281.3 | 33462 | 37995 | < 2.2e-16 |
| triglyceride | 1 | 1116.1 | 33461 | 36879 | < 2.2e-16 |
| HDL | 1 | 21.0 | 33460 | 36858 | 4.532e-06 |
| hemoglobin | 1 | 1568.3 | 33459 | 35290 | < 2.2e-16 |
| serum.creatinine | 1 | 5.7 | 33458 | 35284 | 0.01699 |
| ALT | 1 | 5.8 | 33457 | 35279 | 0.01649 |
| Gtp | 1 | 740.9 | 33456 | 34538 | < 2.2e-16 |
| dental.caries | 1 | 168.8 | 33455 | 34369 | < 2.2e-16 |

| | |
|---------------------|-----|
| NULL | |
| age | *** |
| height.cm. | *** |
| systolic | *** |
| fasting.blood.sugar | *** |
| triglyceride | *** |
| HDL | *** |
| hemoglobin | *** |
| serum.creatinine | * |
| ALT | * |
| Gtp | *** |
| dental.caries | *** |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

Results & Interpretation

Results from the updated fitted logistic regression model shows that only “serum.creatinine” variable is insignificant based on its p-value.

Additionally, results from the ANOVA test shows all variables, including “serum.creatinine”, are significant to the fitted model.

This is judged by the low deviance residuals as well as the Pr(>Chi) of > 0.05 , respectively.

Hence, the “serum.creatinine” variable is kept in the fitted model.

Hide

```
car::vif(logreg2)
```

| | | |
|---------------------|------------------|----------|
| age | height.cm. | systolic |
| 1.421810 | 1.509450 | 1.118942 |
| fasting.blood.sugar | triglyceride | HDL |
| 1.116290 | 1.345118 | 1.269627 |
| hemoglobin | serum.creatinine | ALT |
| 1.362957 | 1.157226 | 1.413102 |
| Gtp | dental.caries | |
| 1.443219 | 1.011522 | |

Results show that all the variables have a VIF value of < 5 .

Test for the Absence of Strongly Influential Outliers

Notes

Test using standardized residuals and Cook's Distance.

Standardized residuals values > 3 = influential outlier.

Cook's D value $>$ Cook's D Threshold $(4/N)$ = influential outlier.

Hide

```
### Place all the calculated values from the logistic regression model into a new data frame.
library(dplyr)
library(magrittr)
```

Attaching package: 'magrittr'

The following object is masked from 'package:purrr':

set_names

The following object is masked from 'package:tidyr':

extract

Hide

```
library(tidyverse)
library(broom)
```

Warning: package 'broom' was built under R version 4.2.3

Hide

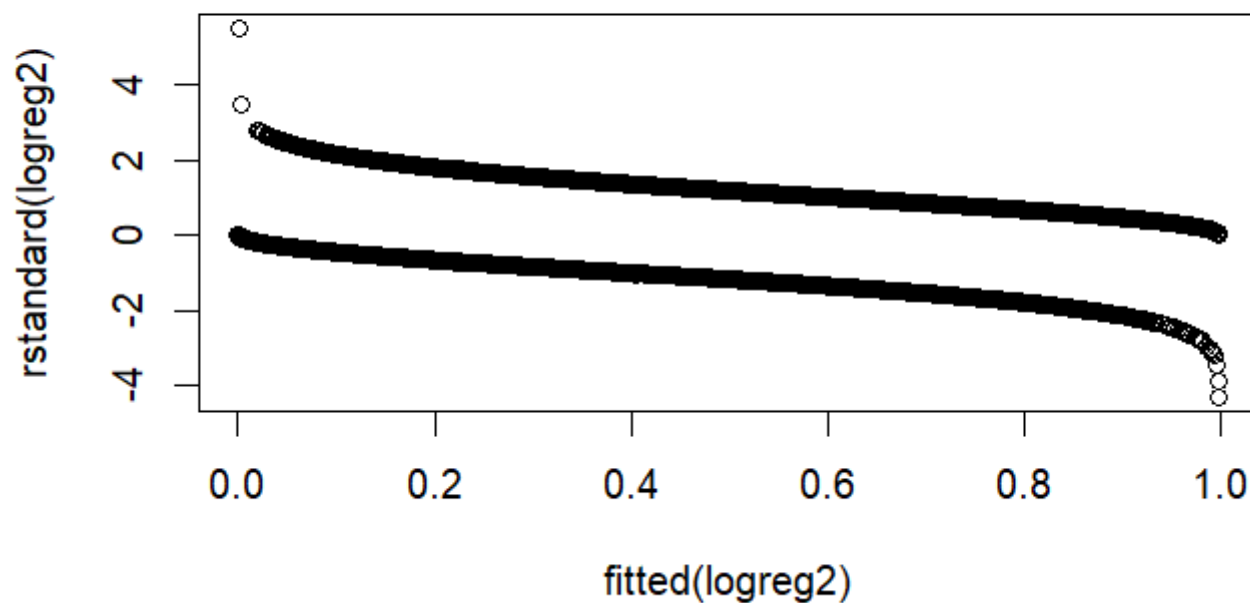
```
logreg.data <- augment(logreg2) %>%
  mutate(index = 1:n())
```

```
### Show the top 6 highest standardized residuals (if  $> 3$  = influential outlier).
head(logreg.data$.std.resid[order(-logreg.data$.std.resid)])
```

```
[1] 5.484120 3.451080 2.800086 2.791547 2.771910 2.737196
```

Hide

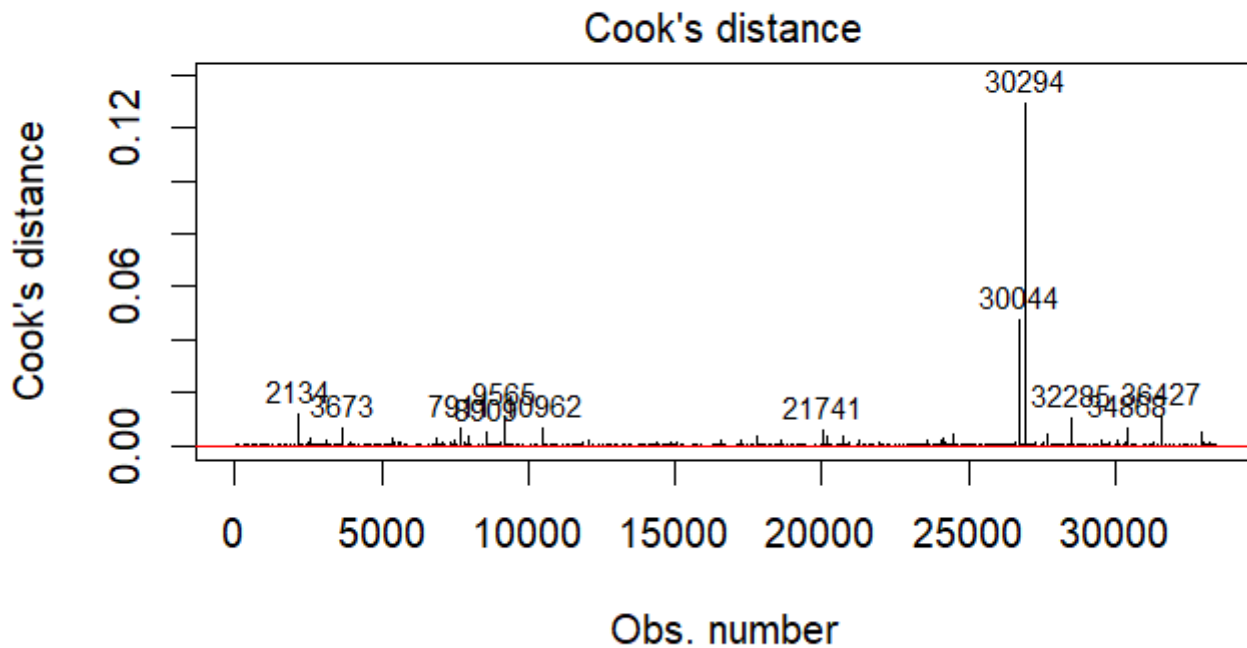
```
### Plot of standardized residuals
plot(fitted(logreg2),
     rstandard(logreg2))
```



Hide

```
### Set Cook's D Threshold.
cook_threshold <- 4 / nrow(df)

### Cook's D Plot.
plot(logreg2, which = 4, id.n = 12)
abline(h = cook_threshold, col = "red")
```



`glm(smoking ~ age + height.cm. + systolic + fasting.blood.sugar + triglycer`

Hide

```
### Put outlier data into a new data frame where > Cook's D Threshold = influential outliers.
influ_out <- logreg.data %>%
  filter(.cooks_d > cook_threshold)

### Get the percentage of influential outliers.
outliers <- round(100*(nrow(influ_out) / nrow(logreg.data)),1)
```

Hide

```
### Get the percentage of influential outliers.
outliers <- round(100*(nrow(influ_out) / nrow(logreg.data)),1)

### Store values in a data variable.
print_outliers <- format(round(outliers, 2), nsmall = 2)

### Print the number of percentage of observations that exceed Cook's distance threshold.
sprintf('Proportion of data points that are highly influential = %s Percent', print_outliers)
```

```
[1] "Proportion of data points that are highly influential = 4.10 Percent"
```

Results & Interpretation

Standardized Residuals.

Results show that none of the data points of the fitted model consist of any outliers.

Cook's Distance.

In addition, based on the pre-defined threshold ($4/N$), only 4.1% of the data points are in the outlier zone, which is small as well.

c. Briefly explain your final model which consider as the best solution.

The final logistic regression model was constructed to predict smoking status by considering various health-related variables. Through a stepwise refinement process, insignificant predictors and potential multicollinearity issues were addressed. The resulting model includes significant predictors such as age, height, systolic blood pressure, fasting blood sugar, triglyceride levels, HDL cholesterol, hemoglobin levels, ALT (Alanine Aminotransferase), GTP (Gamma-Glutamyl Transferase), and the presence of dental caries. Notably, serum creatinine levels were found to be statistically insignificant in the updated model. Diagnostic tests, including VIF for multicollinearity, standardized residuals and Cook's Distance for influential outliers, and deviance tests, were conducted to ensure the model's robustness. The findings indicate that the model provides a significant fit without strong evidence of multicollinearity or influential outliers. Interpretation of the coefficients suggests how each predictor contributes to the likelihood of smoking based on individual health characteristics, providing a valuable tool for understanding and predicting smoking behavior.