1. Preliminary analysis:

#

- # Perform preliminary data inspection and report the findings as to the structure of the data, missing values, duplicates, etc.
- # Based on the findings from the previous question remove duplicates (if any), treat missing values using an appropriate strategy.
- # 2. Prepare an informative report about the data explaining the distribution of the disease and the related factors. You could use the below approach to achieve the objective

#

- # Get a preliminary statistical summary of the data. Explore the measures of central tendencies and the spread of the data overall.
- # Identify the data variables which might be categorical in nature. Describe and explore these variables using appropriate tools e.g. count plot
- # Study the occurrence of CVD across Age.
- # Study the composition of overall patients w.r.t. Gender.
- # Can we detect a heart attack based on anomalies in the Resting Blood Pressure of the patient?
- # Describe the relationship between Cholesterol levels and our target variable.
- # What can be concluded about the relationship between peak exercising and the occurrence of a heart attack.
- # Is thalassemia a major cause of CVD?
- # How are the other factors determining the occurrence of CVD?
- # Use a pair plot to understand the relationship between all the given variables.
- # 3. Build a baseline model to predict using a Logistic Regression and explore the results.

```
library("readxl")
library("writexl")
library("dplyr")
library("ggplot2")
library("Hmisc")
library("treemapify")
```

```
library("scales")
library("ggthemes")
library("corrr")
library("GGally")
library("caret")
setwd("C:/Users/anbha/OneDrive/Desktop/Purdue University/Course6-Capstone Project/Project Data
Set/1582800613_project3datadictionary")
getwd()
data <- data.frame(read_excel("data.xlsx"))</pre>
data
View(data)
head(data)
tail(data)
dim(data)
class(data)
str(data)##Notice all the data types are in Numeric and some needs to be converted to char/factors
wherever applicable
##1.Perform preliminary data inspection and report the findings as the structure of the data, missing
values, duplicates, etc.
##2.Based on the findings from the previous question remove duplicates (if any) and treat missing
values using an appropriate strategy.
names(data)
names(data)<- gsub('','_',names(data)) ##Removing WS in col names if any and replacing with '_'
names(data)
colSums(is.na(data))###no missing values
sum(duplicated(data))##1 duplicated data
```

```
data <-unique(data)
dim(data)##duplicate removed
# age
        age in years
# sex
       (1 = male; 0 = female)
# ср
        chest pain type
# trestbps
                resting blood pressure (in mm Hg on admission to the hospital)
# chol serum cholestoral in mg/dl
# fbs
        (fasting blood sugar > 120 mg/dl) (1 = true; 0 = false)
# restecg
                resting electrocardiographic results
# thalach
                maximum heart rate achieved
                exercise induced angina (1 = yes; 0 = no)
# exang
# oldpeak
                ST depression induced by exercise relative to rest
# slope the slope of the peak exercise ST segment
        number of major vessels (0-3) colored by flourosopy
# ca
# thal 3 = normal; 6 = fixed defect; 7 = reversable defect
# target
                1 or 0
##changing column names into meaningful names
data <- rename(data,
        chest_pain_type=cp,
        resting_blood_pressure = trestbps,
        cholestrol = chol,
        fasting_blood_sugar = fbs,
        resting_ecg = restecg,
        max_heart_rate = thalach,
        exercise_induced_angina = exang,
        st_depression = oldpeak,
        st_slope = slope,
```

```
major_vessels = ca,
        thalessimia = thal
)
names(data)
# 3.Get a preliminary statistical summary of the data. Explore the measures of central tendencies and
the spread of the data overall.
summary(data)
# Identify the data variables which might be categorical in nature. Describe and explore these variables
using appropriate tools e.g., count plot.
##It is clear that columns such as sex,chest_pain_type,fasting_blood_sugar,
##resting_ecg,exercise_induced_angina,st_slope,major_vessels,thalessimia,target
##shouldnt be of numeric type and can be converted to factors for better analysis
##converting numeric to factors
rapply(data,function(x)length(unique(x)))
data %>% summarise_all(funs(n_distinct(.)))##summarizing cat vars
catcols <- c("sex", "chest_pain_type", "fasting_blood_sugar", "resting_ecg",
       "exercise_induced_angina", "st_slope", "thalessimia")
data[catcols] <- lapply(data[catcols], factor)
##changing gender to male/female
##frequency table
data$sex
table(data$sex)
data$sex <- recode_factor(data$sex, '0' = "female", '1' = "male")
```

```
table(data$sex)
##chart
# Describe and explore these variables using appropriate tools e.g., count plot.
# Male percentage is very high as comapred to female in this dataset
countsgender<-table(data$sex)</pre>
countsgender
pct<- round(countsgender/sum(countsgender)*100)</pre>
labels <- paste(" ",c("Female","Male"),"-",pct,"%" )
pie(countsgender, labels = labels, main = "Gender Wise Distribution of Data",
  col = c("red","green"))
##changing chest_pain_type to typical angina, atypical angina, non-anginal pain, asymptomatic
#frequency table
data$chest_pain_type
table(data$chest_pain_type)
data$chest_pain_type <- recode_factor(data$chest_pain_type, '0' = "typical angina", '1' = "angina",
              '2' = 'non-anginal pain',
              '3' = 'asymptomatic')
table(data$chest_pain_type)
#chart
##typical angina occurs most frequently
##and asymptomatic is the least occurring of the chest pain types
countChestPainType <- table(data$chest_pain_type)</pre>
countChestPainType
pct<- round(countChestPainType/sum(countChestPainType)*100)</pre>
labels <- paste(pct," %")
```

```
p <- barplot(countChestPainType,</pre>
    main = "Chest Pain Type Distribution",
    xlab = "Types of Chest Pain",
    ylab = "Number of Patients",
    ylim = c(0, max(countChestPainType) + 100),
    legend = rownames(countChestPainType),
    args.legend = list(x = "topright", inset = c(-0.1, -0.25), cex=0.5),
    col = c("red","blue","green","yellow"))
text(x = p,y = countChestPainType + 25,labels = labels)
##changing fasting blood sugar
#frequency table
data$fasting_blood_sugar
table(data$fasting_blood_sugar)
data$fasting_blood_sugar <- recode_factor(data$fasting_blood_sugar, '0' = "non-diabetic",
                     '1' = "diabetic")
table(data$fasting_blood_sugar)
#chart
##to see spread of diabetic people
##high percentage of people(85%) in this data set are non diabetic
countsdiabetic<-table(data$fasting_blood_sugar)</pre>
countsdiabetic
pct<- round(countsdiabetic/sum(countsdiabetic)*100)</pre>
pct
labels <- paste(pct,"%" )</pre>
xx<-barplot(countsdiabetic,
      width = 1,
```

```
main = "Diabetic and Non diabetic patients",
      xlab = "Diabetic/Non-Diabetic",
      ylab = "Number of Patients",
      ylim = c(0, max(countsdiabetic) + 100),
      legend = rownames(countsdiabetic),
      args.legend = list(x = "topright", inset = c(-0.1, -0.25), cex=0.5),
      col = rainbow(2)
text(x = xx,labels = labels,y=countsdiabetic+20,col = "black")
##changing resting_ecg
##frequency table
data$resting_ecg
table(data$resting_ecg)
data$resting_ecg <- recode_factor(data$resting_ecg, '0' = "normal",
                       '1' = "abnormal",
                       '2' = 'hyper' )
table(data$resting_ecg)
##chart
##TO SEE resting_ecg spread of its categories
## we can see hyper is very negligible quantity
##and both normal and abnormal are in equal quantity
plotdata <- data %>%
 count(resting_ecg)
```

```
ggplot(plotdata,
   aes(fill = resting_ecg,
     area = n,
     label = resting_ecg)) +
geom_treemap() +
 geom_treemap_text(colour = "white",
          place = "centre") +
labs(title = "Resting ECG Spread") +
theme(legend.position = "none")
##changing exercise_induced_angina
##frequency table
data$exercise_induced_angina
table(data$exercise_induced_angina)
data$exercise_induced_angina <- recode_factor(data$exercise_induced_angina, '0' = "no",
                  '1' = "yes")
table(data$exercise_induced_angina)
##chart
##TO SEE exercise_induced_angina spread of its categories
## Almost 67% of data doesnt have exercise_induced_angina
plotdata <- data %>%
count(exercise_induced_angina) %>%
 arrange(desc(exercise_induced_angina)) %>%
```

```
mutate(prop = round(n*100/sum(n), 1),
     lab.ypos = cumsum(prop) - 0.5*prop)
plotdata$label <- pasteO(plotdata$exercise_induced_angina, "\n",</pre>
             round(plotdata$prop), "%")
ggplot(plotdata,
   aes(x = "",
      y = prop,
      fill = exercise_induced_angina)) +
 geom_bar(width = 1,
      stat = "identity",
      color = "black") +
 geom_text(aes(y = lab.ypos, label = label),
      color = "black") +
 coord_polar("y",
       start = 0,
       direction = -1) +
 theme_void() +
 theme(legend.position = "FALSE") +
 labs(title = "Excercise Induced Angina Spread")
##changing st_slope
##frequency table
data$st_slope
table(data$st_slope)
data$st_slope <- recode_factor(data$st_slope, '0' = "unsloping",
```

```
'1' = "flat",
                         '2' = "downsloping")
##chart
##st_slope
##downsloping and flat are of equal proportion but unsloping is very less
plotdata <- data %>%
count(st_slope) %>%
arrange(desc(st_slope)) %>%
 mutate(prop = round(n*100/sum(n), 1),
     lab.ypos = cumsum(prop) - 0.5*prop)
plotdata
plotdata$label <- paste0(plotdata$st_slope, "-",
             round(plotdata$prop), "%")
ggplot(plotdata,
   aes(x = "",
     y = prop,
     fill = st_slope)) +
 geom_bar(width = 1,
     stat = "identity",
     color = "black") +
 geom_text(aes(y = lab.ypos, label = label),
      color = "black") +
theme_void() +
theme(legend.position = "FALSE") +
labs(title = "ST Slope Spread")
##changing thalessimia
##only three categories of thalessimia given
##1 = normal; 2 = fixed defect; 3 = reversable defect
```

```
##so converting 4th category i.e 0 to 2 (since 2 has the max no of values)
##frequency table
data$thalessimia
table(data$thalessimia)
str(data$thalessimia)
##chart
plotdata <- data %>%
count(thalessimia)
ggplot(plotdata,
   aes(fill = thalessimia,
     area = n,
     label = thalessimia)) +
 geom_treemap() +
 geom_treemap_text(colour = "white",
          place = "centre") +
labs(title = "Thalessimia Spread") +
theme(legend.position = "none")
##since we dont have any factor for 0 in thalessimia,
##we will convert the 2 rows where thalessimia = 0 as thalessimia = 2
data$thalessimia <- recode_factor(data$thalessimia, '0' = "2")
data$thalessimia <- recode_factor(data$thalessimia, '1' = "normal",
                  '2'= "fixed defect",
                  '3'= "reversable defect")
##plotting again after converting thalessimia = 2
##frequency table
table(data$thalessimia)
##chart
plotdata <- data %>%
count(thalessimia)
```

```
ggplot(plotdata,
   aes(fill = thalessimia,
      area = n,
      label = thalessimia)) +
 geom_treemap() +
 geom_treemap_text(colour = "white",
           place = "centre") +
 labs(title = "Thalessimia ECG Spread") +
 theme(legend.position = "none")
##5.Study the occurrence of CVD across different ages
## To analyze the CVD, let's explore the target variable first
##frequency table
str(data$target)
table(data$target)
## 0 - Disease- & 1 - disease+
data$target2 <- recode_factor(data$target, '0' = "Disease-",
                         '1' = "Disease+")
str(data$target2)
table(data$target2)
##chart for disease- and disease+
plotdata <- data %>%
 count(target2)
ggplot(plotdata,
   aes(x = target2,
      y = n)) +
 geom_bar(stat = "identity") +
 geom_text(aes(label = n),
```

```
vjust=-0.5) +
labs(x = "Target",
   y = "Number Of Patients",
   title = "Target Variable Distribution")
##This shows there are more people with CVD in this Data Set
##Now let us compare Age vs CVD
##Dividing the data set based on Target Variable
## as dataHealthy and dataDiseased for better understanding
dataHealthy <- data %>% filter(target2 == 'Disease-')
View(dataHealthy)
dataDiseased <- data %>% filter(target2 == 'Disease+')
View(dataDiseased)
##Study the occurrence of CVD across different ages
##bar plot (group) for health and Diseased
ggplot(data,
   aes(x = age,
     fill = target2 )) +
geom_bar(position = position_dodge(preserve = "single" ))+
labs(title = "Age Distribution of Diseased and Healthy", y="Number Of Patients",x="Age")
##kernel density plot for diseased
##the graph
ggplot(dataDiseased, aes(x=age))+
```

```
geom_density(color="darkblue", fill="lightblue")+
geom_vline(aes(xintercept=mean(age)),
       color="blue", linetype="dashed", size=1)+
labs(title = "Age Density Plot For Diseased",y="Density",x="Age")
ggplot(dataDiseased,
   aes(x = as.factor(age))) +
geom_bar(fill = "indianred3",
     color = "black")+
labs(title = "Frequency by age for diseased",x="Age",y="Number Of Patients")
##from the graph it is evident that CVD increases from 47 and peaks at 54
##so 47-54 is the riskiest age band for getting CVD
##6.Can we detect heart attack based on anomalies in resting blood pressure of the patient?
##box plot for resting_blood_pressure for diseased and healthy
ggplot(data, aes(x=target2, y=resting_blood_pressure, fill=target2)) +
geom_boxplot(alpha=0.3) +
labs(title = "Distribution of Resting Blood Pressure with Target Variable",y="Resting Blood
Pressure",x="Target Variable")+
theme(legend.position="none")
ggplot(data, aes(x=sex, y=resting_blood_pressure, fill=target2)) +
   geom_boxplot(alpha=0.3) +
labs(title = "Gender Based Distribution of Resting Blood Pressure with Target Variable",y="Resting
Blood Pressure",x="Gender")+
   theme(legend.title = element_blank())
##male patients have the same range but female patients range vary w.r.t Blood pressure
#Male range: 120-140(both diseased and Healthy)
```

```
#Female range: 120-140(diseased), 130-155(healthy)
#Density plot to identify relationship between resting_blood_pressure and CVD
ggplot(data, aes(x=resting_blood_pressure,fill=target2))+
geom_density(color="red")+
geom_vline(aes(xintercept=mean(resting_blood_pressure)),
       color="blue", linetype="dashed", size=1)+
labs(title = "Density Distribution of Blood Pressure with Target Variable",x="Resting Blood
Pressure",y="Density")
##We cannot say Resting Blood Pressure has connection to CDV in this case,
##as both the box plot and density plot suggest they have the same range
##7. Study the composition of overall patients w.r.t . gender.
##stacked bar for overall percentage of gender in data
plotdata <- data %>%
count(sex) %>%
arrange(desc(sex)) %>%
 mutate(prop = round(n*100/sum(n), 1),
    lab.ypos = cumsum(prop) - 0.5*prop)
plotdata
plotdata$label <- paste0(plotdata$sex, "-",
             round(plotdata$prop), "%")
ggplot(plotdata,
   aes(x = "",
     y = prop,
     fill = sex)) +
geom_bar(width = 1,
     stat = "identity",
```

```
color = "black") +
geom_text(aes(y = lab.ypos, label = label),
      color = "black") +
theme_void() +
theme(legend.position = "FALSE") +
labs(title = "Gender Spread")
##donut chart for percentage of male and female in diseased
plotdata <- dataDiseased %>%
count(sex) %>%
arrange(desc(sex)) %>%
 mutate(prop = round(n*100/sum(n), 1),
    lab.ypos = cumsum(prop) - 0.5*prop)
plotdata$label <- pasteO(plotdata$sex, "\n",
             round(plotdata$prop), "%")
plotdata
ggplot(plotdata,
   aes(x = 1,
     y = prop,
     fill = sex)) +
geom_col() +
geom_text(aes(y = lab.ypos, label = label),
      color = "black") +
coord_polar("y") +
xlim(c(0.2, 1 + 0.5))+
theme_void() +
```

```
theme(legend.position = "FALSE") +
labs(title = "Gender spread of Patients in Diseased Data")
##pie chart for percentage of male and female in healthy
plotdata <- dataHealthy %>%
count(sex) %>%
arrange(desc(sex)) %>%
 mutate(prop = round(n*100/sum(n), 1),
     lab.ypos = cumsum(prop) - 0.5*prop)
plotdata$label <- paste0(plotdata$sex, "\n",
             round(plotdata$prop), "%")
ggplot(plotdata,
   aes(x = "",
     y = prop,
     fill = sex)) +
geom_bar(width = 1,
     stat = "identity",
     color = "black") +
geom_text(aes(y = lab.ypos, label = label),
      color = "black") +
coord_polar("y",
       start = 0,
       direction = -1) +
theme_void() +
theme(legend.position = "FALSE") +
 labs(title = "Gender spread of Patients in Healthy Data")
```

```
##donut chart to calcualte diseased in female population
plotdata <- data %>% filter(sex == 'female') %>%
count(target2) %>%
arrange(desc(target2)) %>%
 mutate(prop = round(n*100/sum(n), 1),
     lab.ypos = cumsum(prop) - 0.5*prop)
plotdata$label <- paste0(plotdata$target2, "\n",
             round(plotdata$prop), "%")
plotdata
ggplot(plotdata,
   aes(x = 1,
     y = prop,
     fill = target2)) +
geom_col() +
geom_text(aes(y = lab.ypos, label = label),
      color = "black") +
coord_polar("y") +
xlim(c(0.2, 1 + 0.5))+
theme_void() +
theme(legend.position = "FALSE") +
 labs(title = "Diseased and Healthy in Female Patients")
##pie chart to calcualte diseased in male population
plotdata <- data %>% filter(sex == 'male') %>%
count(target2) %>%
```

```
arrange(desc(target2)) %>%
mutate(prop = round(n*100/sum(n), 1),
    lab.ypos = cumsum(prop) - 0.5*prop)
plotdata$label <- pasteO(plotdata$target2, "\n",
             round(plotdata$prop), "%")
ggplot(plotdata,
   aes(x = "",
     y = prop,
     fill = target2)) +
geom_bar(width = 1,
     stat = "identity",
     color = "black") +
geom_text(aes(y = lab.ypos, label = label),
      color = "black") +
coord_polar("y",
       start = 0,
       direction = -1) +
theme_void() +
theme(legend.position = "FALSE") +
labs(title = "Dieseased and Healthy in Male patients")
##from these graphs we are able to infer that
# 1.In overall data, male population is more than women
# but large percentage of women seem to be diseased (75%)
# as compared to men (45%). So 3/4th of female seeems to have CVD. So women in this data seems to
be at high risk
```

```
# 8.Describe the relationship between cholesterol levels and our target variable.
my_data <- data %>%
group_by(target2) %>%
summarise(mean = mean(cholestrol),
      std = sd(cholestrol),
      min = min(cholestrol),
      max = max(cholestrol),
      med = median(cholestrol))
my_data
e <- ggplot(data, aes(x = target2, y = cholestrol))
e + geom_violin(aes(fill = target2), trim = FALSE) +
geom_boxplot(width = 0.2)+
scale_fill_manual(values = c("#00AFBB", "#E7B800", "#FC4E07"))+
theme(legend.position = "none")+labs(title="Cholestrol Distibution with Target Variable",x="Target
Variable",y="Cholestrol")
##the graph and stat functions shows disease+ data has quite a lot of outliers
# and also the violin plot suggest data for diseased and healthy
# are distributed in the same range to an extent and hence it is inconclusive with this data set
# 9. What can be concluded about the relationship between peak exercising and occurrence of heart
attack?
plotdata <- data %>%
group_by(target2) %>%
summarise(n = n(),
      mean = mean(max_heart_rate),
```

```
sd = sd(max_heart_rate),
      se = sd / sqrt(n)
plotdata
ggplot(plotdata,
   aes(x = target2,
     y = mean,
     group = 1)) +
geom_point(size = 3) +
geom_line() +
geom_errorbar(aes(ymin = mean - se,
          ymax = mean + se),
        width = .1)+
labs(title="Mean Value Comparison of Max Heart Rate with Target Variable",x="Target
Variable",y="Mean")
ggplot(data, aes(x=max_heart_rate,fill = target2))+
geom_density(color="darkblue")+
geom_vline(aes(xintercept=mean(max_heart_rate)),
       color="blue", linetype="dashed", size=1)+
labs(title = "Density plot for Max Heart Rate with Target Variable",x="Max Heart Rate",y="Density")
##lets explore more
ggplot(data,
   aes(x = max_heart_rate ,
     fill = target2 )) +
```

```
geom_bar(position = position_dodge(preserve = "single" ))+
labs(title = "Max Heart Rate comparison of with Target",x="Max Heart Rate",y="Number Of Patients")
##dieseased seems to have max heart rate greater than mean value i.e 150
##max density concentration at 162 for diseased.
##max density concentration at 148 for healthy
#10.Is thalassemia a major cause of CVD? How are the other factors determining the occurrence of CVD?
ggplot(data,
   aes(x = target2,
     fill = thalessimia )) +
geom_bar(position = position_dodge(preserve = "single" ))+
labs(title = "Thalessimia comparison with Target Variable",x="Target",y="Number Of Patients")+
geom_text(aes(label =..count..),stat="count",position = position_dodge(width =1 ),vjust=0.01)
#thalessimia appears to be a major cause, accounting to max percentage of diseased patients
# out of 164 diseased, more than 125 seems to have irreversable thalessimia
##reversable defect and normal doesnt seem to have great impact on CVD
## lets see how diabetes relate to CVD
plotdata <- data %>% filter(fasting_blood_sugar == 'diabetic') %>%
count(target2) %>%
arrange(desc(target2)) %>%
 mutate(prop = round(n*100/sum(n), 1),
    lab.ypos = cumsum(prop) - 0.5*prop)
plotdata$label <- paste0(plotdata$target2, "\n",
             round(plotdata$prop), "%")
plotdata
```

```
ggplot(plotdata,
   aes(x = 1,
     y = prop,
     fill = target2)) +
geom_col() +
 geom_text(aes(y = lab.ypos, label = label),
      color = "black") +
coord_polar("y") +
xlim(c(0.2, 1 + 0.5))+
theme_void() +
theme(legend.position = "FALSE") +
 labs(title = "Diabetic patients in Healthy and Dieseased")
## lets see how diabetes relate to CVD
plotdata <- dataDiseased %>%
count(fasting_blood_sugar) %>%
arrange(desc(fasting_blood_sugar)) %>%
 mutate(prop = round(n*100/sum(n), 1),
     lab.ypos = cumsum(prop) - 0.5*prop)
plotdata$label <- paste0(plotdata$fasting_blood_sugar, "\n",
             round(plotdata$prop), "%")
plotdata
ggplot(plotdata,
   aes(x = 1,
     y = prop,
     fill = fasting_blood_sugar)) +
```

```
geom_col() +
 geom_text(aes(y = lab.ypos, label = label),
      color = "black") +
 coord_polar("y") +
 xlim(c(0.2, 1 + 0.5))+
 theme_void() +
 theme(legend.position = "FALSE") +
 labs(title = "Spread of Diabetic/Non Diabetic patients in diseased")
plotdata <- dataHealthy %>%
 count(fasting_blood_sugar) %>%
 arrange(desc(fasting_blood_sugar)) %>%
 mutate(prop = round(n*100/sum(n), 1),
     lab.ypos = cumsum(prop) - 0.5*prop)
plotdata$label <- pasteO(plotdata$fasting_blood_sugar, "\n",</pre>
             round(plotdata$prop), "%")
plotdata
ggplot(plotdata,
   aes(x = 1,
      y = prop,
      fill = fasting_blood_sugar)) +
 geom_col() +
 geom_text(aes(y = lab.ypos, label = label),
      color = "black") +
 coord_polar("y") +
 xlim(c(0.2, 1 + 0.5))+
```

```
theme_void() +
theme(legend.position = "FALSE") +
labs(title = "Spread of Diabetic/Non Diabetic patients in Healthy")
##diabetic doesnt play an important role in this Dataset
##relationship between chest pain type and target var
ggplot(data, aes(fill=target2, x=chest_pain_type)) +
geom_bar(position="stack", stat="count") +
ggtitle("Chest Pain type distribution between Diseased and Healthy patients") +
ylab("Number Of Patients")+coord_flip()
# '1' = "angina",
#'2' = 'non-anginal pain',
#'3' = 'asymptomatic' all seem to contribute more to diseased than typical angina
# i.e a patient would more likely to be healthy if cp type is typical angina
#than compared to other CP types
##in other words angina contributes to cvd the most among cp types
##resting_ecg relation with target
plotdata <- data %>% filter(resting_ecg == 'normal') %>%
count(target2) %>%
arrange(desc(target2)) %>%
 mutate(prop = round(n*100/sum(n), 1),
    lab.ypos = cumsum(prop) - 0.5*prop)
plotdata$label <- paste0(plotdata$target2, "\n",
             round(plotdata$prop), "%")
```

```
ggplot(plotdata,
   aes(x = "",
     y = prop,
     fill = target2)) +
geom_bar(width = 1,
     stat = "identity",
     color = "black") +
geom_text(aes(y = lab.ypos, label = label),
      color = "black") +
coord_polar("y",
       start = 0,
       direction = -1) +
theme_void() +
theme(legend.position = "FALSE") +
 labs(title = "Normal ECG Spread across Target Variable")
plotdata <- data %>% filter(resting_ecg == 'abnormal') %>%
count(target2) %>%
arrange(desc(target2)) %>%
 mutate(prop = round(n*100/sum(n), 1),
     lab.ypos = cumsum(prop) - 0.5*prop)
plotdata$label <- pasteO(plotdata$target2, "\n",
             round(plotdata$prop), "%")
ggplot(plotdata,
```

```
aes(x = "",
     y = prop,
     fill = target2)) +
geom_bar(width = 1,
     stat = "identity",
     color = "black") +
 geom_text(aes(y = lab.ypos, label = label),
      color = "black") +
coord_polar("y",
       start = 0,
       direction = -1) +
theme_void() +
theme(legend.position = "FALSE") +
labs(title = "Abnormal ECG Spread across Target Variable")
##Abnormal ECG seems to be associated more with Disease+ patients
plotdata <- dataDiseased %>%
count(resting_ecg) %>%
arrange(desc(resting_ecg)) %>%
 mutate(prop = round(n*100/sum(n), 1),
    lab.ypos = cumsum(prop) - 0.5*prop)
plotdata$label <- paste0(plotdata$resting_ecg, "\n",
             round(plotdata$prop), "%")
ggplot(plotdata,
   aes(x = "",
```

```
y = prop,
      fill = resting_ecg)) +
 geom_bar(width = 1,
      stat = "identity",
      color = "black") +
 geom_text(aes(y = lab.ypos, label = label),
      color = "black") +
 coord_polar("y",
        start = 0,
        direction = -1) +
 theme_void() +
 theme(legend.position = "FALSE") +
 labs(title = "ECG Spread Across Diseased")
##abnormal accounts to almost 60% of positive cases
#11.Use a pair plot to understand the relationship between all the given variables.
str(data)
d <- data[,sapply(data,class)=="numeric"]</pre>
d$target2 <- data$target2
ggpairs(d,
    columns = 1:(ncol(d)-1),
    aes(color = target2, alpha = 0.5))
##ST_Depression and Major_Vessels seems to have a tight co relation
##with the Target Var
res <- cor.test(data$st_depression, data$target,</pre>
         method = "pearson")
res
```

```
##box plot for St depression
ggplot(data, aes(x=target2, y=st_depression, fill=target2)) +
geom_boxplot(alpha=0.3) +
labs(title = "Distribution of ST Depression for Target Variable",y="ST Depression",x="Target")+
theme(legend.position="none")
##violin plot for Major Vessels
res <- cor.test(data$major_vessels, data$target,
        method = "pearson")
res
e <- ggplot(data, aes(x = target2, y = major_vessels))
e + geom_violin(aes(fill = target2), trim = FALSE) +
scale_fill_manual(values = c("#00AFBB", "#E7B800", "#FC4E07"))+
labs(title = "Distribution of Major Vessels Across Target Variable",x="Target",y="Major Vessels")+
theme(legend.position = "none")
##It is evident that
##ST_Deperesssion and Major Vessels have a strong impact on the predictor var target
##st_slope relation
plotdata <- dataDiseased %>%
count(st_slope) %>%
arrange(desc(st_slope)) %>%
mutate(prop = round(n*100/sum(n), 1),
    lab.ypos = cumsum(prop) - 0.5*prop)
plotdata
```

```
plotdata$label <- paste0(plotdata$st_slope, "-",
             round(plotdata$prop), "%")
ggplot(plotdata,
   aes(x = "",
      y = prop,
      fill = st_slope)) +
 geom_bar(width = 1,
      stat = "identity",
      color = "black") +
 geom_text(aes(y = lab.ypos, label = label),
      color = "black") +
 theme_void() +
 theme(legend.position = "FALSE") +
 labs(title = "ST Slope Spread across diseased")
plotdata <- dataHealthy %>%
 count(st_slope) %>%
 arrange(desc(st_slope)) %>%
 mutate(prop = round(n*100/sum(n), 1),
     lab.ypos = cumsum(prop) - 0.5*prop)
plotdata
plotdata$label <- paste0(plotdata$st_slope, "-",
             round(plotdata$prop), "%")
ggplot(plotdata,
   aes(x = "",
      y = prop,
      fill = st_slope)) +
```

```
geom_bar(width = 1,
     stat = "identity",
     color = "black") +
geom_text(aes(y = lab.ypos, label = label),
      color = "black") +
theme_void() +
theme(legend.position = "FALSE") +
labs(title = "ST Slope Spread Across Healthy")
##downsloping seems to contribute to CVD
##Relationship with exercise_induced_Angina
##No % is really high (67%), lets see how this plays with target var
plotdata <- dataDiseased %>%
count(exercise_induced_angina) %>%
arrange(desc(exercise_induced_angina)) %>%
mutate(prop = round(n*100/sum(n), 1),
    lab.ypos = cumsum(prop) - 0.5*prop)
plotdata
plotdata$label <- paste0(plotdata$exercise_induced_angina, "-",
             round(plotdata$prop), "%")
ggplot(plotdata,
   aes(x = "",
     y = prop,
     fill = exercise_induced_angina)) +
geom_bar(width = 1,
     stat = "identity",
```

```
color = "black") +
geom_text(aes(y = lab.ypos, label = label),
      color = "black") +
theme_void() +
theme(legend.position = "FALSE") +
labs(title = "Exercise Induced Angina Spread Across Diseased")
plotdata <- dataHealthy %>%
count(exercise_induced_angina) %>%
arrange(desc(exercise_induced_angina)) %>%
 mutate(prop = round(n*100/sum(n), 1),
    lab.ypos = cumsum(prop) - 0.5*prop)
plotdata
plotdata$label <- paste0(plotdata$exercise_induced_angina, "-",
             round(plotdata$prop), "%")
ggplot(plotdata,
   aes(x = "",
     y = prop,
     fill = exercise_induced_angina)) +
geom_bar(width = 1,
     stat = "identity",
     color = "black") +
geom_text(aes(y = lab.ypos, label = label),
      color = "black") +
theme_void() +
theme(legend.position = "FALSE") +
labs(title = "Exercise Induced Angina Spread Across Healthy")
```

```
# The percentage of 'no' in Disease+ data is very high whereas healthy patients
# dataset is domainated by 'yes'
# and hence it plays a role in deciding the CVD outcome.
##Let us write this excel to analyze with Tableau
write_xlsx(data,"cvd_data_latest.xlsx")
# 13. Visualize the variables using Tableau to create an understanding for attributes of a Diseased vs a
Healthy person.
#
# 14. Demonstrate the variables associated with each other and factors to build a dashboard
##Model building and Testing
# 12. Perform logistic regression, predict the outcome for test data, and validate the results by using the
confusion matrix.
data1 <- data %>% select(-target2)
data1
data1$target <- as.factor(data1$target)</pre>
train_indices <- sample(1:nrow(data1),0.7*nrow(data1))
train indices
train <- data1[train_indices,]
test <- data1[-train_indices,]</pre>
##building model by including all columns from the data (basemodel)
basemodel <- glm(target~.,data = train,family = 'binomial')
summary(basemodel)
pred_prob <- predict(basemodel,test)</pre>
pred <- as.factor(ifelse(pred_prob >= 0.5,1,0))
```

```
##drilling down with columns having high co-relation suggested by R
model1 <- glm(target~
sex+chest_pain_type+resting_blood_pressure+resting_ecg+max_heart_rate+thalessimia+major_vessels,
data = train, family = 'binomial')
summary(model1)
pred prob <- predict(model1,test)</pre>
pred <- as.factor(ifelse(pred prob >= 0.5,1,0))
caret::confusionMatrix(pred,test$target) ##accuracy reduces to 78%
##building model by including columns selected from my analysis (myModel)
myModel <- glm(target~
sex+age+chest pain type+resting ecg+max heart rate+exercise induced angina+st depression+st sl
ope+thalessimia+major_vessels,data = train,family = 'binomial')
summary(myModel)
pred_prob <- predict(myModel,test)</pre>
pred <- as.factor(ifelse(pred_prob >= 0.5,1,0))
caret::confusionMatrix(pred,test$target)
```