Heart Failure Analysis and Prediction



About this dataset

Cardiovascular diseases (CVDs) are the number 1 cause of death globally, taking an estimated 17.9 million lives each year, which accounts for 31% of all deaths worlwide. Heart failure is a common event caused by CVDs and this dataset contains 12 features that can be used to predict mortality by heart failure. Most cardiovascular diseases can be prevented by addressing behavioural risk factors such as tobacco use, unhealthy diet and obesity, physical inactivity and harmful use of alcohol using population-wide strategies. People with cardiovascular disease or who are at high cardiovascular risk (due to the presence of one or more risk factors such as hypertension, diabetes, hyperlipidaemia or already established disease) need early detection and management wherein a machine learning model can be of great help.

Importing libraries

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```
library(ggplot2)
library(dplyr);
library(rpart)
library(rpart.plot)
library(caret);
library(reshape2)
library(DataExplorer)
library(gridExtra)
```

Impontring the data

We will import heart_failure_clinical_records_dataset.csv

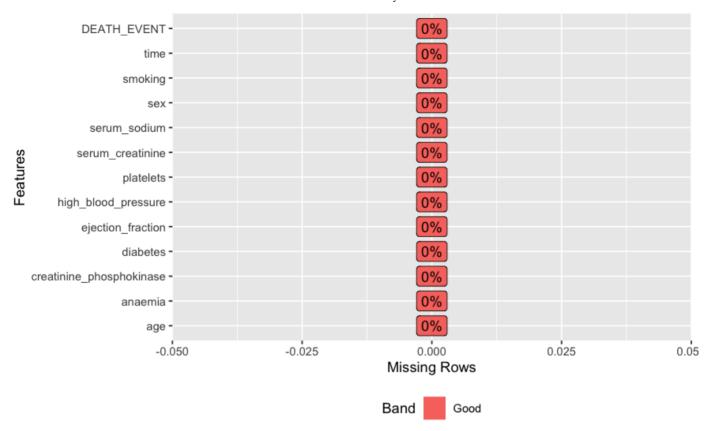
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```
data <- read.csv("heart failure clinical records dataset.csv")
```

First, let's check nulls in the dataset

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plot_missing(data)



Get a look at the dataset

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head(data, 10)

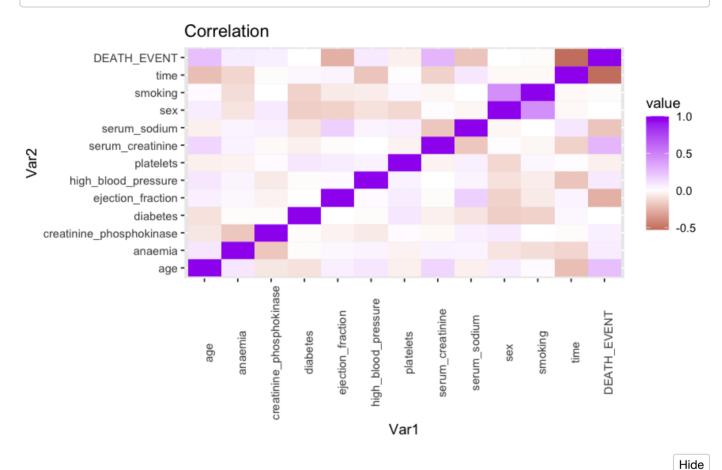
	 <dbl:< th=""><th>anae > <int></int></th><th>creatinine_phosphokinase <int></int></th><th>diabetes <int></int></th><th>ejection_fraction <int></int></th><th>high_blood_pre</th></dbl:<>	anae > <int></int>	creatinine_phosphokinase <int></int>	diabetes <int></int>	ejection_fraction <int></int>	high_blood_pre
1	75	0	582	0	20	
2	55	0	7861	0	38	
3	65	0	146	0	20	
4	50	1	111	0	20	
5	65	1	160	1	20	
6	90	1	47	0	40	
7	75	1	246	0	15	
8	60	1	315	1	60	
9	65	0	157	0	65	
10	80	1	123	0	35	
1-1	0 of 1	0 rows 1-	8 of 13 columns			

Fining correlations between features

```
# calculating correlations and rounding to nearest 2 decimal points
cormap <- round( cor(data), 2)

#convert the matrix to a dataframe
cormap_melted<-melt(cormap)

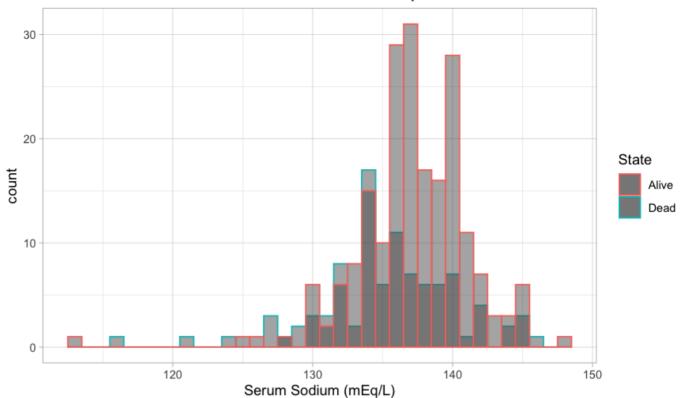
#creating heatmap
ggplot(data = cormap_melted, aes(x=Var1, y=Var2, fill=value) ) +
   geom_tile() + scale_fill_gradient2(low="darkred",high="purple",mid="white") + theme
(axis.text.x = element_text(angle = 90)) + labs(title = 'Correlation')</pre>
```



dead_sodium <- filter(data, DEATH_EVENT==1) %>% select(serum_sodium)
alive_sodium <- filter(data, DEATH_EVENT==0) %>% select(serum_sodium)

ggplot() +
 geom_histogram(data = dead_sodium, aes(serum_sodium, color='Dead'), alpha=0.5, binw
idth = 1) +
 geom_histogram(data = alive_sodium, aes(serum_sodium, color='Alive'), alpha = 0.5, b
inwidth = 1) +
 theme_light() + labs(color='State', title='Distribution of serum sodium for dead an
d alive patients', x='Serum Sodium (mEq/L)')

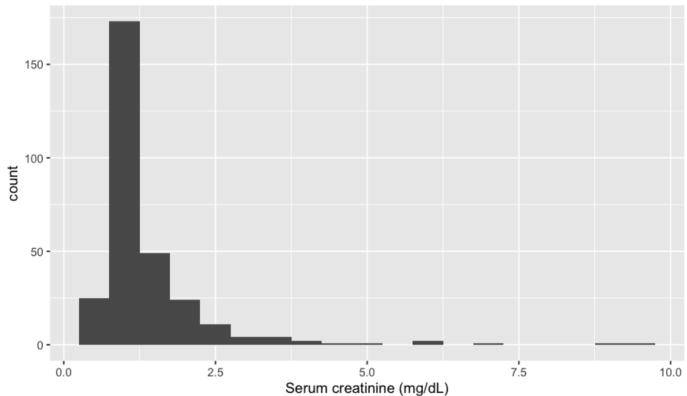
Distribution of serum sodium for dead and alive patients



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 $ggplot(data, aes(x = serum_creatinine)) + geom_histogram(binwidth = 0.5) + labs(title = 'Histogram of serum creatinine distribution', x='Serum creatinine (mg/dL)')$

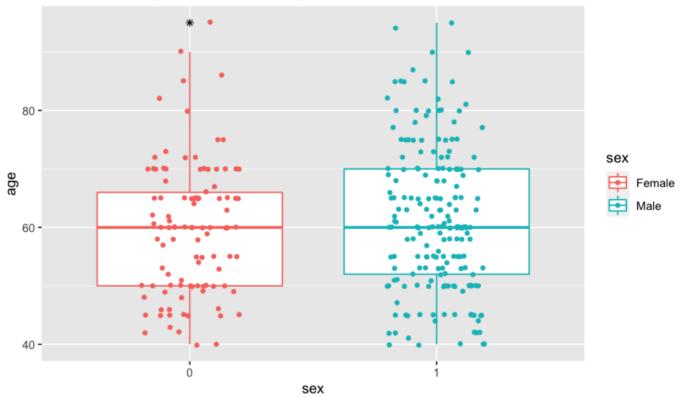
Histogram of serum creatinine distribution



The distribution of serum_sodium is left skewed, while serum_creatinine is right skewed.

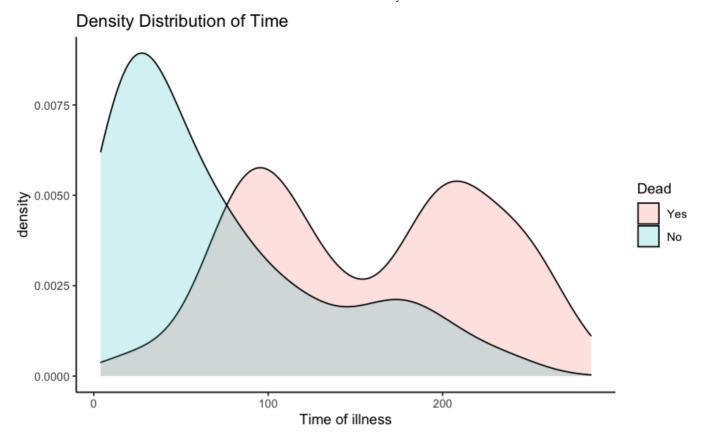
ggplot(data, aes(x=factor(sex), y=age, color=factor(sex))) + geom_boxplot(outlier.col
or = 'black', outlier.shape = 8) + geom_jitter(shape=16, position=position_jitter(0.2
)) + labs(x='sex', title='Distribution of gender and their ages', color='sex') + scal
e_color_discrete(labels=c('Female', 'Male'))

Distribution of gender and their ages



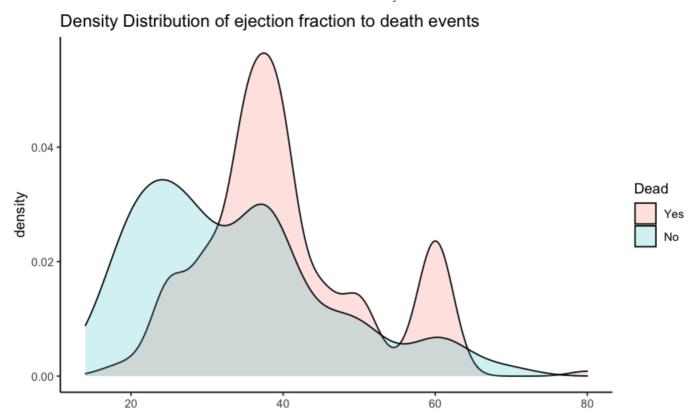
Next, lets investigate some relations

The duration of illness of dead patient and recovered patient



At first, patient have higher probablility of recovery, while passing 100+ days incease the probablility of death.

Let's see ejection fraction affects the occurance of heart failure.



Ejection Fraction

Dead patients have higher probablility of having ejection fraction.

Predict using Logistic Regression

```
set.seed(123) #setting the seed to make sure that the split function give us the same
results

taining.samples <- data$DEATH_EVENT %>%
    createDataPartition(p=0.5, list=FALSE) # splitting the data 50-50, to avoid over fi
tting

# store the test data and train data
train.data <- data[taining.samples, ] #first partition of sample
test.data <- data[- taining.samples, ] #last partition of sample

# run the logistic regression model using all of the features we have using binomial
classification
model <- glm(DEATH_EVENT~., data = train.data, family=binomial())

# inspect the coefficients resulted from the logistic regression
summary(model)$coef</pre>
```

```
Std. Error
                                                       z value
                              Estimate
                                                                   Pr(>|z|)
                          4.997730e+00 9.422564e+00 0.5304002 5.958345e-01
(Intercept)
                          4.585887e-02 2.462071e-02 1.8626135 6.251667e-02
age
                          3.921844e-01 5.633341e-01 0.6961843 4.863134e-01
anaemia
creatinine_phosphokinase -4.944054e-05 3.384572e-04 -0.1460762 8.838612e-01
                          7.300236e-01 5.704696e-01 1.2796889 2.006546e-01
diabetes
ejection_fraction
                         -9.656250e-02 2.568859e-02 -3.7589649 1.706178e-04
                         3.043644e-01 5.365284e-01 0.5672849 5.705207e-01
high blood pressure
                         -2.082016e-06 2.489475e-06 -0.8363275 4.029707e-01
platelets
                          9.266276e-01 2.658751e-01 3.4851988 4.917715e-04
serum creatinine
                         -2.615251e-02 6.628936e-02 -0.3945205 6.931968e-01
serum sodium
                          1.069458e-01 6.458398e-01 0.1655919 8.684781e-01
sex
smoking
                         -1.866607e-01 6.101210e-01 -0.3059405 7.596499e-01
                         -2.710215e-02 5.310282e-03 -5.1037120 3.330550e-07
time
```

```
# testing the model
probabilities <- model %>% predict(test.data, type = "response")

# thresholding results; results from the logistic function >= 0.5 lies in the 1 clas
s, while < 0.5 lies in the opposite one
predicted.classes <- ifelse(probabilities >= 0.5, "1", "0")

# Model accuracy
mean(predicted.classes == test.data$DEATH_EVENT) * 100
```

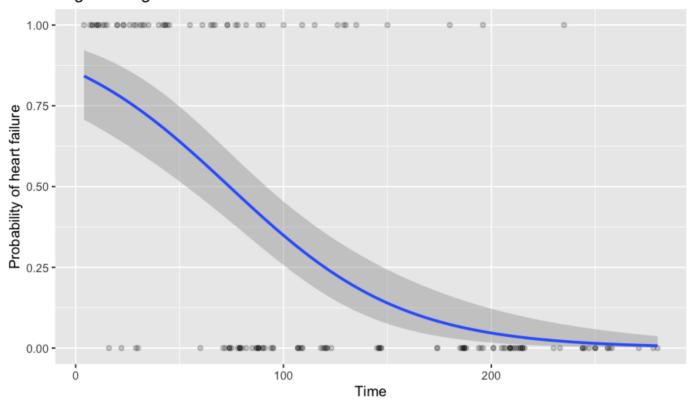
```
[1] 81.20805
```

plotting logistic function with time parameter

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```
train.data %>%
  mutate(prob = ifelse(DEATH_EVENT == "1", 1, 0)) %>%
  ggplot(aes(time, prob)) +
  geom_point(alpha = 0.2) +
  geom_smooth(method = "glm", method.args = list(family = "binomial")) +
  labs(
    title = "Logistic Regression Model",
    x = "Time",
    y = "Probability of heart failure")
```

Logistic Regression Model



Creating the descision tree

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```
set.seed(123)
data$Dead<-ifelse(data$DEATH_EVENT!=1, "No", "Yes")

#This variable is no needed for constructing a classification tree
data$DEATH_EVENT<-NULL

#creating tree, I will use a small critical point
heartTree<-rpart(Dead~.,data=data,control=rpart.control(cp=0.00001))

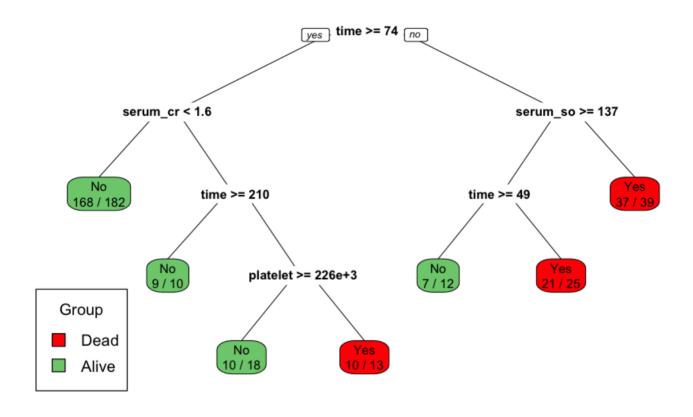
#Creating a matrix to check the accuracy of decision tree
conf.matrix <- table(data$Dead, predict(heartTree, type="class"))

rownames(conf.matrix) <- paste("Actual", rownames(conf.matrix), sep = ":")

colnames(conf.matrix) <- paste("Pred", colnames(conf.matrix), sep = ":")

print(conf.matrix)</pre>
```

```
Pred:No Pred:Yes
Actual:No 194 9
Actual:Yes 28 68
```



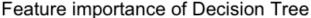
```
Accuracy<-(conf.matrix[1,1] + conf.matrix[2,2])/sum(conf.matrix)*100

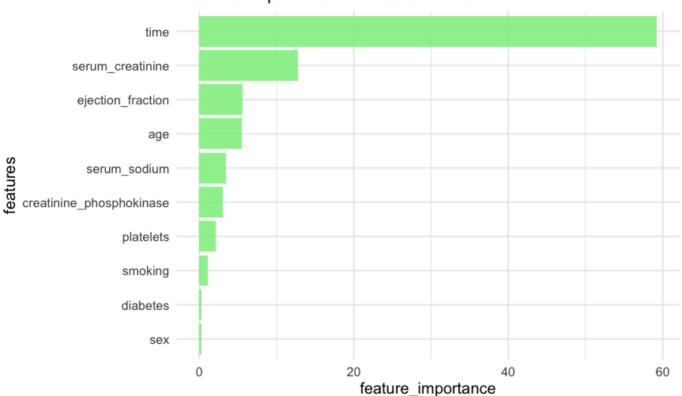
Accuracy
```

[1] 87.62542

Finding the real impact of features

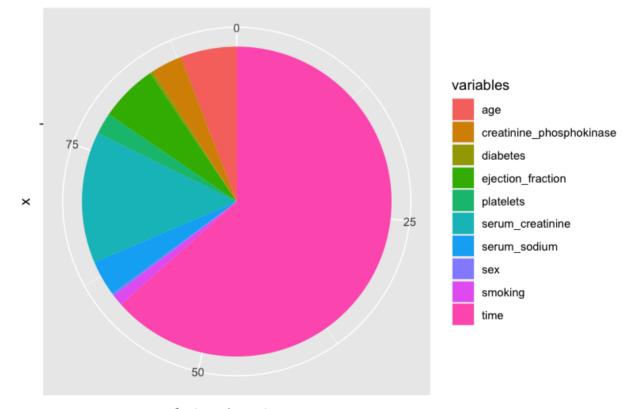
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#Visualize the feature importance using pie chart
importance <- data.frame(variables=names(heartTree\$variable.importance), feature_impo
rtance=heartTree\$variable.importance)
ggplot(data=importance, aes(x="", y=feature_importance, fill=variables)) + geom_bar(s
tat="identity", width=1) +
 coord_polar("y", start=0)</pre>



feature_importance

- * time is the most effective of all, while diabetes is the least effective
- * sex is not as effective as time