# ARIGNAR ANNA GOVERNMENT ARTS COLLEGE VILLUPURAM – 605 602.



#### **DEPARTMENT OF COMPUTER SCIENCE**

#### **MACHINE LEARNING WITH PYTHON**

Project Title: A Review of Liver Patient Analysis Method using Machine

Learning

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# **Abstract**

Around a million deaths occur due to liver diseases globally.

There are several traditional methods to diagnose liver diseases, but they are expensive. Early prediction of liver disease would

benefit all individuals prone to liver diseases by providing early treatment. As technology is growing in health care, machine learning significantly affects health care for predicting conditions at early stages. This study finds how accurate machine learning is in predicting liver disease.

This present study introduces the liver disease prediction (LDP) method in predicting liver disease that can be utilised by health professionals, stakeholders, students and researchers. Five algorithms, namely Support Vector Machine (SVM), Naïve Bayes, K-Nearest Neighbors (K-NN), Linear Discriminant Analysis (LDA), and Classification and Regression Trees (CART), are selected. The accuracy is compared to uncover the best classification method for predicting liver disease using R and Python. From the results, K-NN obtains the best accuracy with 91.7%, and the autoencoder network achieved 92.1% accuracy, which is above the acceptable level of accuracy and can be considered for liver disease prediction.

## **Introduction**

In this project we will analyse the parameters of various classification algorithms and compare their predictive accuracies so as to find out the best classifier for determining the liver disease. This project compares various classification algorithms such as Random Forest, Logistic Regression, KNN and ANN Algorithm with an aim to identify the best technique. Based on this study, Random Forest with the highest accuracy outperformed the other algorithms and can be further utilised in the prediction of liver disease and can be recommended to the Liver. According to World Health Organization

(WHO) report in 2018, the number of deaths due to liver diseases is around one million and ranked 11th in the world with a critical number of fatalities (World Total Deaths, n.d.). Unnoticed at the initial stages, these symptoms are only visible when the disease turns chronic. However, even though the liver is partially infected, it can still function (Devikanniga et al., 2020).

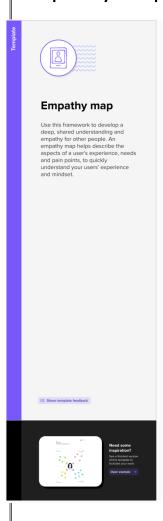
Diagnosis of liver diseases can be divided into three stages i.e., the first stage is liver inflammation, the second is liver scarring (cirrhosis), and the final stage is liver cancer or failure. Since these scenarios are present in liver disease, early prediction is significant to provide better health for New Zealanders. If liver disease is diagnosed early, there will be a chance of early treatment and control of deaths due to liver diseases (Arbain & Balakrishnan, 2019). But when the liver fails to function, few treatments are available except liver transplantation (Shaheamlung et al., 2020), which is very expensive, particularly in New Zealand (Hepatitis C, 2021). Apparently, in New Zealand, 35 -40% of the population are not diagnosed with Hepatitis C at the early stages because of the asymptomatic behaviour of liver disease. Unfortunately, most of these individuals do not know the risks linked to liver disease. Due to the asymptomatic behaviour and higher costs of liver disease treatment, it is essential to prevent or diagnose early for better treatment.

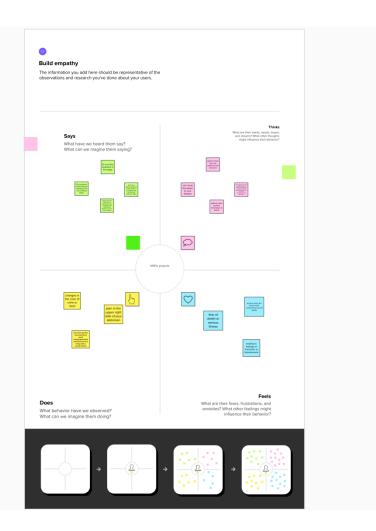
## <u>Purpose</u>

Liver function tests can be used to: Screen for liver infections, such as hepatitis. Monitor the progression of a disease, such as viral or alcoholic hepatitis, and determine how well a treatment is working. Measure the severity of a disease, particularly scarring of the liver (cirrhosis)

#### Problem Definetion & Design Thinking

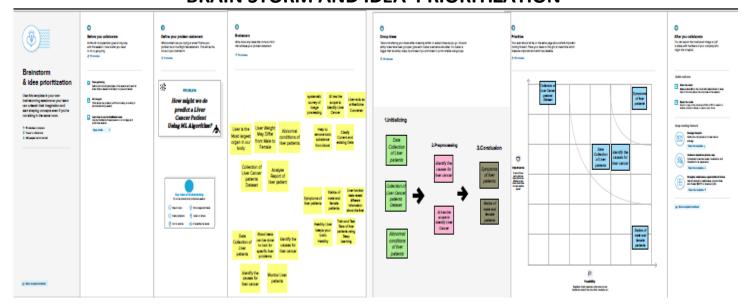
# **Empathy** map





# Ideation & brainstorming map

#### **BRAIN STORM AND IDEA PRIORITIZATION**



# Result:

Liver Patients analysis using machine learning can provide accurate and realiable results for the diagnosis, prognosis, and treatment of liver disease. Machine learning algorithm can analyse large amounts of patients data and identify patterns that may be difficult for human experts to detect.

## Home.html Source code

<!DOCTYPE html>

<html lang="en">

<head>

<title>Bootstrap Example</title>

```
<meta charset="utf-8">
<meta name="viewport" content="width=device-width,initial-scale=1">
link rel="stylesheet" href="https://maxcdn.bootstrapcdn.com/bootstrap/3.4.1/css/bootstrap.min.css">
<style type="text/css">
body{
background-color:#ffcf0059;
}
nav{
```

```
background-color:#ad38c2;
height:60px;
}
.navbar-brand{
color:white
}
</style>
</head>
<body>
<nav class="navbar">
<div class="containe-field">
<div class="navbar-header">
<a class="navbar-brand"><h1>Liver Patient Analysis</h1></a></nav>
</div>
 <h2>Introduction</h2>
 Liver diseases averts the normal function of the liver. This disease is caused by an assortment of
elements
that harm the liver. Diagnosis of liver infection at the preliminary stage is important for better treatment.
In
today's scenario devices like sensors are used for detection of infections. Accurate classification
techniques are required for automatic identification of disease samples. This disease diagnosis is very
 costly and complicated. Therefore, the goal of this work is to evaluate the performance of different
Machine
Learning algorithms in order to reduce the high cost of liver disease diagnosis. Early prediction of liver
 disease using classification algorithms is an efficacious task that can help the doctors to diagnose the
 disease within a short duration of time. In this project we will analyse the parameters of various
```

classification algorithms and compare their predictive accuracies so as to find out the best classifier for determining the liver disease. This project compares various classification algorithms such as Random Forest, Logistic Regression, KNN and ANN Algorithm with an aim to identify the best technique. Based on this study, Random Forest with the highest accuracy outperformed the other algorithms and can be further

utilised in the prediction of liver disease and can be recommended to the user.

Technical Architecture:

Project Flow:

- User interacts with the UI to enter the input.
- Entered input is analysed by the model which is integrated.
- Once model analyses the input the prediction is showcased on the UI

To accomplish this, we have to complete all the activities listed below,

- Define Problem / Problem Understanding
- O Specify

</div>

</html>

#### <u>Output</u> :

#### Liver Patient Analysis

#### Introduction

Liver diseases averts the normal function of the liver. This disease is caused by an assortment of elements that harm the liver. Diagnosis of liver infection at the preliminary stage is important for better treatment. In today's scenario devices like sensors are used for detection of infections. Accurate classification techniques are required for automatic identification of disease samples. This disease diagnosis is very costly and complicated. Therefore, the goal of this work is to evaluate the performance of different Machine Learning algorithms in order to reduce the high cost of liver disease diagnosis. Early prediction of liver disease using classification algorithms is an efficacious task that can help the doctors to diagnose the disease within a short duration of time. In this project we will analyse the parameters of various classification algorithms and compare their predictive accuracies so as to find out the best classifier for determining the liver disease. This project compares various classification algorithms such as Random Forest, Logistic Regression, KNN and ANN Algorithm with an aim to identify the best technique. Based on this study, Random Forest with the highest accuracy outperformed the other algorithms and can be further utilised in the prediction of liver disease and can be recommended to the user. Technical Architecture: Project Flov: • User interacts with the UI to enter the input. • Entered input is analysed by the model which is integrated. • Once model analyses the input the prediction is showcased on the UI To accomplish this, we have to complete all the activities listed below, • Define Problem / Problem Understanding • Specify

# Index.html Source code

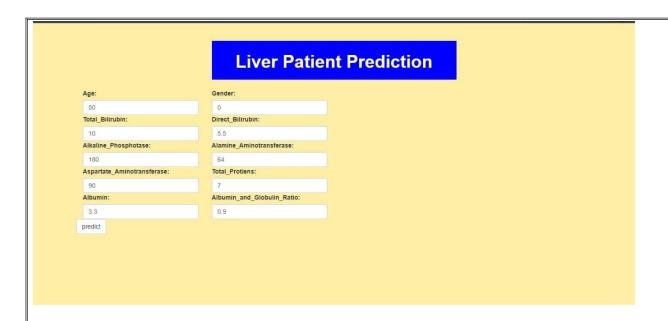
```
<!DOCTYPE html>
<html>
<head>
<title>Liver Patient Analysis</title>
<!-- Latest Compiled and minified CSS -->
<Link rel="stylesheet" href="https://maxcdn.bootstrapcdn.com/bootstrap/3.3.7/css/bootstrap.min.css">
<style type="text/css">
body{
background-color: #ffcf0059;
 }
.page-header{
background-color: blue;
width: 100%;
height: auto;
text-align: center;
padding-top: 5px;
color: #fff
}
h1{
 font-size: 40px;
 font-weight: bold;
 }
</style>
</head>
</body>
 <div class="container">
 <div class="row">
  <div class="col-md-3"></div>
  <div class="col-md-6">
```

```
<div class="page-header">
<h1>Liver Patient Prediction</h1>
</div>
</div>
</div>
</div>
 <div class="container">
 <div class="row">
 <div class="col-mid-3"></div>
 <div class="col-md-6">
 <form action="/data_predict" method="post">
 <div class="row">
 <div class="col-md-6">
 <div class="form-gorup">
 <label for="Age">Age:</label>
 <input type="text" class="form-control" id="age" name="age">
 </div>
 </div>
 <div class="col-md-6">
 <div class="form-gorup">
 <label for="gender">Gender:</label>
 <input type="text" class="form-control" id="gender" name="gender">
 </div>
 </div>
 <div class="col-md-6">
 <div class="form-gorup">
 <label for="tb">Total_Bilirubin:</label>
 <input type="text" class="form-control" id="tb" name="tb">
 </div>
 </div>
```

```
</div>
<div class="col-md-6">
<div class="form-gorup">
<label for="db">Direct_Bilirubin:</label>
<input type="text" class="form-control" id="db" name="db">
</div>
</div>
<div class="col-md-6">
<div class="form-gorup">
<label for="ap">Alkaline_Phosphotase:</label>
<input type="text" class="form-control" id="ap" name="ap">
</div>
</div>
<div class="col-md-6">
<div class="form-gorup">
<label for="aa1">Alamine_Aminotransferase:</label>
<input type="text" class="form-control" id="aa1" name="aa1">
</div>
</div>
<div class="col-md-6">
<div class="form-gorup">
<label for="aa2">Aspartate_Aminotransferase:
<input type="text" class="form-control" id="aa2" name="aa2">
</div>
</div>
<div class="col-md-6">
<div class="form-gorup">
<label for="tp">Total_Protiens:</label>
<input type="text" class="form-control" id="tp" name="tp">
</div>
```

```
</div>
<div class="col-md-6">
<div class="form-gorup">
<label for="a">Albumin:</label>
<input type="text" class="form-control" id="a" name="a">
</div>
</div>
<div class="col-md-6">
<div class="form-gorup">
<label for="agr">Albumin_and_Globulin_Ratio:
<input type="text" class="form-control" id="agr" name="agr">
</div>
<button type="submit" class="btn btn-default" > predict</button>
</form>
</div>
</div>
</div>
<!-- Latest Compiled and minified javascript -->
<script src="https://maxcdn.bootstrapcdn.com/bootstrap/3.3.7/js/bootstrap.min.js"></script>
</body>
</html>
```

#### **Output:**



# No chance.html source code

```
<!DOCTYPE html>
<html>
<head>
 <title>Liver Patient Analysis</title>
 <!-- Latest Compiled and minified CSS -->
  <Link rel="stylesheet" href="https://maxcdn.bootstrapcdn.com/bootstrap/3.3.7/css/bootstrap.min.css">
  <style type="text/css">
    body{
      background-color: #ffcf0059;
    }
    .page-header{
      background-color: blue;
      width: 100%;
      height: auto;
      text-align: center;
      padding-top: 5px;
```

```
color: #fff;
       }
       h1{
     font-size: 40px;
     font-weight: bold;
   }
 </style>
 </head>
 </body>
   <div class="container">
   <div class="row">
   <div class="col-md-3"></div>
   <div class="col-md-6">
   <div class="page-header">
   <h1>Liver Patient Prediction</h1></div>
   You have a liver desease problem, You must and should consult a doctor. Take care</h3>
   </style>
   </div>
   </div>
   </div>
   </div>
 <!-- Latest Compiled and minified javascript -->
 <script src="https://maxcdn.bootstrapcdn.com/bootstrap/3.3.7/js/bootstrap.min.js"></script>
</html>
```

#### Output:

#### **Liver Patient Prediction**

You have a liver desease problem You must and should consult a doctor. Take care

# **Advantages**

General

Diagnoses, grades and stages:

Hepatitis C

Hepatitis B

Steatohepatitis

Autoimmune hepatitis

Evalutes abnormal liver function tests

Identifies hepatotoxicity

Clarifies uncertain diagnoses

Confirms etiology of liver masses

Defines extent of necroinflammatory activity

Differentiates fiabrosis from cirrhosis

Liver transplant

Identifies acute cellular rejection

Defines recurrence of original disease

Identifies progressive fibrosis

Diagnoses other liver processes.

# Disadvantage

#### General

**Invasive** 

Accessibility to the procedure

Need for training

Cost

## Sample

Sampling error

Intraobserver and interobserver variations in

Interpretatooin

Specimen length and width

#### **Patient**

Site pain

Shoulder pain

Neuralgia

**Hypotension** 

Bleeding

Hemothorax

Hemoblilia

## **Application:**

- Diagnosis: Liver patients analysis can be used to diagnose liver diseases such as cirrhosis, hepatitis, and Livercancer, by analysing various biomarkers such as liver enyzmes, bilirubin, and albumin, doctors can determine the health of the liver and diagnose any underlying diseases.
- Treatment: Liver patients analysis can also to monitor the effectiveness of treatmentsfor liver diseases. By regulary analysing liver function tests and other biomarkers, doctors can achieve optimal results

## Conclusion

- pertinent Since the liver disease is not easy to diagnose, given the delicate nature of its signs, this research is in determining the algorithms that have better accuracy in predicting this dreadful disease.
- Once the dataset is selected, the preprocessing step is conducted by replacing the missing values and balancing the dataset.
- After that, using R, five different supervised learning methods are applied (i.e., SVM, Naïve Bayes, K-NN, LDA, and CART), and the accuracy with confusion matrix metrics are recorded.

- In this study, the autoencoder with 3-layers achieved an accuracy of 92.1%, slightly higher than K-NN due to its ability to ascertain overlapping features better than conventional K-NNs. Most of the algorithms are more than the acceptable level of accuracy, which is 75%.
- The results from this study would be able to assist health professionals and relevant stakeholders in the early detection of liver disease.

## Future scope

- In this paper, we proposed and built a machine learning based on a hybrid classifier to be used as a classification model for liver diseases diagnosis to improve performance and experrts to identify the chances of disease and conscious orescription of further treatment healthcare and examination.
- In future work, the use of fast datasets technique like apache hadoop or spark can be incorporated with this technique. In addition to this, we can use distributed refined algorithm like forest tree implement in apache hadoop to increase scalability and efficieny.

# **Appenix**

Source code

# Mliestone 2:

import pandas as pd

import numpy as np import seaborn as sns import matplotlib.pyplot as plt from matplotlib import rcParams from scipy import stats import warnings warnings.filterwarnings('ignore') from sklearn.tree import DecisionTreeClassifier from sklearn.ensemble import RandomForestClassifier from sklearn.model\_selection import train\_test\_split from sklearn.metrics import classification report, confusion matrix data=pd.read\_csv("indian\_liver\_patient.csv") data.head() data.info() <class 'pandas.core.frame.DataFrame'> RangeIndex: 583 entries, 0 to 582 Data columns (total 11 columns): # Column Non-Null Count Dtype 0 Age 583 non-null int64 1 Gender 583 non-null object float64 2 Total\_Bilirubin 583 non-null 3 Direct\_Bilirubin 583 non-null float64 Alkaline\_Phosphotase 583 non-null int64

5	Alamine_Aminotransferase	583 non-null	int64		
6	Aspartate_Aminotransferase	583 non-null	int64		
7	Total_Protiens	583 non-null	float64		
8	Albumin	583 non-null	float64		
9	Albumin_and_Globulin_Ratio	579 non-null	float64		
10	Dataset	583 non-null	int64		
dtyp	pes: float64(5), int64(5), object(1)				
mer	nory usage: 50.2+ KB				
data	ı.isnull().any()				
Age		False			
Gen	der	False			
Tota	ıl_Bilirubin	False			
Dire	ct_Bilirubin	False			
Alka	line_Phosphotase	False			
Alar	nine_Aminotransferase	False			
Aspa	artate_Aminotransferase	False			
Tota	l_Protiens	False			
Albı	ımin	False			
Albumin_and_Globulin_Ratio		True			
Dataset		False			
dtype: bool					
data.isnull().sum()					
Age		0			
Gen	der	0			
Tota	ıl_Bilirubin	0			
Dire	ct_Bilirubin	0			
Alka	line_Phosphotase	0			
Alar	nine_Aminotransferase	0			

Aspartate_Aminotransferase	0					
Total_Protiens	0					
Albumin	0					
Albumin_and_Globulin_Ratio	4					
Dataset	0					
dtype: int64						
data['Albumin_and_Globulin_Ratio'] data['Albumin_and_Globulin_Ratio'].	= fillna(data['Albumin_and_Globulin_Ratio'].mode()[0])					
data.isnull().sum()						
Age	0					
Gender	0					
Total_Bilirubin	0					
Direct_Bilirubin	0					
Alkaline_Phosphotase	0					
Alamine_Aminotransferase	0					
Aspartate_Aminotransferase	0					
Total_Protiens	0					
Albumin	0					
Albumin_and_Globulin_Ratio	0					
Dataset	0					
dtype: int64						
from sklearn.preprocessing import La	abelEncoder					
lc = LabelEncoder()						
data['Gender'] = lc.fit_transform(data	a['Gender'])					
Milestone 3:						

## Milestone 3:

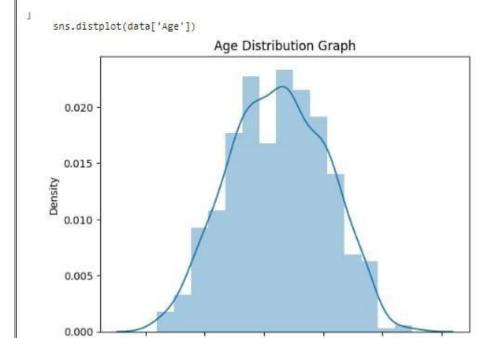
data.describe()

	Age	Gender	Total_Bilirubin	Direct_Bilirubin	Alkaline_Phosphotase	Alamine_Aminotransferase	Aspartate_Aminotransferase	Total_Protiens	Albumin	Albumin_and_Globulin_Ratio	Dataset
count	583.000000	583.000000	583.000000	583.000000	583.000000	583.000000	583.000000	583.000000	583.000000	583.000000	583.000000
mean	44,746141	0.756432	3.298799	1.486106	290.576329	80.713551	109.910806	6.483190	3.141852	0.947427	1.286449
std	16.189833	0.429603	6.209522	2.808498	242.937989	182.620356	288.918529	1.085451	0.795519	0.318522	0.452490
min	4.000000	0.000000	0.400000	0.100000	63.000000	10.000000	10.000000	2.700000	0.900000	0.300000	1.000000
25%	33.000000	1.000000	0.800000	0.200000	175.500000	23.000000	25.000000	5.800000	2.600000	0.700000	1.000000
50%	45.000000	1.000000	1.000000	0.300000	208.000000	35,000000	42.000000	6.600000	3.100000	0.950000	1.000000
75%	58.000000	1.000000	2.600000	1.300000	298.000000	60.500000	87.000000	7.200000	3.800000	1.100000	2.000000
max	90.000000	1.000000	75.000000	19 700000	2110.000000	2000 0000000	4929 000000	9.600000	5.500000	2 800000	2.000000

sns.distplot(data['Age'])

plt.title('Age Distribution Graph')

plt.show()



40

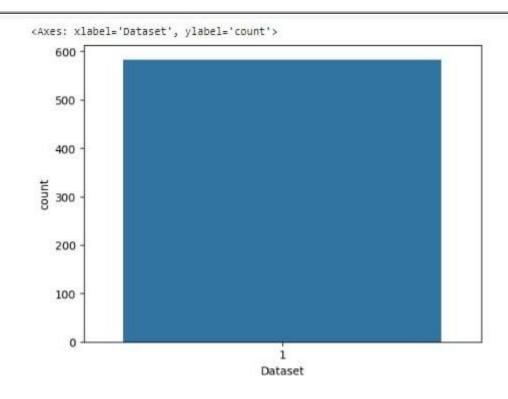
Age

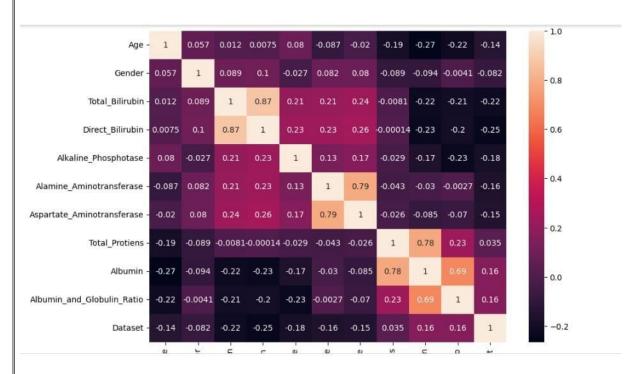
60

80

100

20





sns.countplot(data['Dataset'],x=data['Gender'])

plt.figure(figsize=(10,7))

sns.heatmap(data.corr(),annot=True)

from sklearn.preprocessing import scale

x=pd.DataFrame (scale(x),columns=x.columns)

x\_scaled.head()

	age	gender	Total_Bilirubin	Direct_Bilirubin	Alkaline_Phosphotase
0	1.252098	-1.762281	-0.418878	-0.493964	-0.426715
1	1.066637	0.567446	1.225171	1.430423	1.682629
2	1.066637	0.567446	0.644919	0.931508	0.821588
3	0.819356	0.567446	-0.370523	-0.387054	-0.447314
4	1.684839	0.567446	0.096902	0.183135	-0.393756

x=data.iloc[:,:-1]

y=data.Dataset

from sklearn.model\_selection import train\_test\_split

x\_train,x\_test,y\_train,y\_test=train\_test\_split(x,y,test\_size=0.2,random\_state=42)

from imblearn.over\_sampling import SMOTE

smote = SMOTE()

y\_train.value\_counts()

1 329

2 137

Name: Dataset, dtype: int64

x\_train\_smote, y\_train\_smote = smote.fit\_resample(x\_train,y\_train)

y\_train\_smote.value\_counts()

1 329

2 329

Name: Dataset, dtype: int64

## Milestone 4:

from sklearn.ensemble import RandomForestClassifier

RFmodel=RandomForestClassifier()

RFmodel.fit(x\_train,y\_train)

```
RandomForestClassifierRandomForestClassifier()
```

```
RFpred=RFmodel.predict(x_test)
RFaccuracy=accuracy_score(RFpred,y_test)
RFaccuracy
0.7521367521367521
RFcm=confusion_matrix(RFpred,y_test)
RFcm
array([[75, 17],
   [12, 13]])
from sklearn.neighbors import KNeighborsClassifier
KNN=KNeighborsClassifier()
KNN.fit(x_train,y_train)
KNNpred=KNN.predict(x_test)
KNNaccuracy=accuracy_score(KNNpred,y_test)
KNNaccuracy
0.6837606837606838
KNNcm=confusion_matrix(KNNpred,y_test)
KNNcm
array([[69, 19],
   [18, 11]])
```

from sklearn.tree import DecisionTreeClassifier

```
DTC=DecisionTreeClassifier()
DTC.fit(x_train,y_train)

    LogisticRegression

      LogisticRegression()
DTCpred=DTC.predict(x_test)
DTCaccuracy=accuracy_score(DTCpred,y_test)
DTCaccuracy
0.717948717948718
DTCcm=confusion matrix(DTCpred,y test)
DTCcm
array([[66, 12],
   [21, 18]])
from sklearn.linear_model import LogisticRegression
LR=LogisticRegression()
LRpred=LR.predict(x_test)
LRaccuracy=accuracy_score(LRpred,y_test)
LRaccuracy
0.7435897435897436
LRcm=confusion_matrix(LRpred,y_test)
LRcm
array([[82, 25],
   [5, 5]])
import tensorflow.keras
from tensorflow.keras.models import Sequential
from tensorflow.keras.layers import Dense
```

```
classifier = Sequential()
classifier.add(Dense(units=100,activation='relu',input dim=10))
classifier.add(Dense(units=50,activation='relu'))
classifier.add(Dense(units=1,activation='sigmoid'))
classifier.compile(optimizer='adam',loss='binary crossentropy',metrics=['accuracy'])
RFmodel history=classifier.fit(x train,y train,batch size=100,validation split=0.2,epochs=100)
Epoch 1/100
-593204.0625 - val accuracy: 0.7234
Epoch 2/100
-601997.3125 - val accuracy: 0.7234
Epoch 3/100
-610722.8125 - val accuracy: 0.7234
Epoch 4/100
-619532.7500 - val accuracy: 0.7234
Epoch 5/100
-628574.0000 - val_accuracy: 0.7234
Epoch 6/100
-637505.2500 - val accuracy: 0.7234
Epoch 7/100
-646466.5625 - val_accuracy: 0.7234
Epoch 8/100
-655572.5625 - val accuracy: 0.7234
Epoch 9/100
```

```
-664791.8125 - val_accuracy: 0.7234
Epoch 10/100
-674201.8125 - val accuracy: 0.7234
Epoch 11/100
-683482.2500 - val accuracy: 0.7234
Epoch 12/100
-692927.8125 - val accuracy: 0.7234
Epoch 13/100
-702565.5625 - val accuracy: 0.7234
Epoch 14/100
-712313.8125 - val accuracy: 0.7234
Epoch 15/100
-721993.8125 - val accuracy: 0.7234
Epoch 16/100
-731739.5625 - val accuracy: 0.7234
Epoch 17/100
-741573.6875 - val accuracy: 0.7234
Epoch 18/100
-751650.8125 - val accuracy: 0.7234
Epoch 19/100
-761723.0000 - val accuracy: 0.7234
Epoch 20/100
```

```
-771933.6875 - val_accuracy: 0.7234
Epoch 21/100
-782164.2500 - val accuracy: 0.7234
Epoch 22/100
-792485.2500 - val accuracy: 0.7234
Epoch 23/100
-803042.0625 - val accuracy: 0.7234
Epoch 24/100
-813523.7500 - val accuracy: 0.7234
Epoch 25/100
-824129.3750 - val accuracy: 0.7234
Epoch 26/100
-834718.0625 - val accuracy: 0.7234
Epoch 27/100
-845578.3750 - val accuracy: 0.7234
Epoch 28/100
-856461.6250 - val accuracy: 0.7234
Epoch 29/100
-867335.2500 - val accuracy: 0.7234
Epoch 30/100
-878283.5625 - val accuracy: 0.7234
Epoch 31/100
```

```
-889272.2500 - val_accuracy: 0.7234
Epoch 32/100
val loss: -900362.7500 - val accuracy: 0.7234
Epoch 33/100
val loss: -911565.0000 - val accuracy: 0.7234
Epoch 34/100
val loss: -922823.0000 - val accuracy: 0.7234
Epoch 35/100
val loss: -934481.6250 - val accuracy: 0.7234
Epoch 36/100
val loss: -945996.4375 - val accuracy: 0.7234
Epoch 37/100
val loss: -957543.7500 - val accuracy: 0.7234
Epoch 38/100
val loss: -969030.5625 - val accuracy: 0.7234
Epoch 39/100
val loss: -980694.3125 - val accuracy: 0.7234
Epoch 40/100
val loss: -992526.8125 - val accuracy: 0.7234
Epoch 41/100
val loss: -1004365.9375 - val accuracy: 0.7234
Epoch 42/100
```

```
val_loss: -1016444.1875 - val_accuracy: 0.7234
Epoch 43/100
val loss: -1028518.8125 - val accuracy: 0.7234
Epoch 44/100
val loss: -1040690.8750 - val accuracy: 0.7234
Epoch 45/100
val loss: -1052845.2500 - val accuracy: 0.7234
Epoch 46/100
val loss: -1065299.1250 - val accuracy: 0.7234
Epoch 47/100
val loss: -1077800.1250 - val accuracy: 0.7234
Epoch 48/100
val_loss: -1090211.3750 - val_accuracy: 0.7234
Epoch 49/100
val loss: -1102746.2500 - val accuracy: 0.7234
Epoch 50/100
val loss: -1115452.1250 - val accuracy: 0.7234
Epoch 51/100
val loss: -1128213.5000 - val accuracy: 0.7234
Epoch 52/100
val loss: -1141175.3750 - val accuracy: 0.7234
Epoch 53/100
```

```
val_loss: -1153975.8750 - val_accuracy: 0.7234
Epoch 54/100
val loss: -1166885.2500 - val accuracy: 0.7234
Epoch 55/100
val loss: -1179737.3750 - val accuracy: 0.7234
Epoch 56/100
val loss: -1193090.7500 - val accuracy: 0.7234
Epoch 57/100
val loss: -1206648.1250 - val accuracy: 0.7234
Epoch 58/100
val loss: -1220226.7500 - val accuracy: 0.7234
Epoch 59/100
val loss: -1233714.5000 - val accuracy: 0.7234
Epoch 60/100
val loss: -1247228.3750 - val accuracy: 0.7234
Epoch 61/100
val loss: -1260871.1250 - val accuracy: 0.7234
Epoch 62/100
val loss: -1274690.7500 - val accuracy: 0.7234
Epoch 63/100
val loss: -1288476.2500 - val accuracy: 0.7234
Epoch 64/100
```

```
val_loss: -1302399.7500 - val_accuracy: 0.7234
Epoch 65/100
val loss: -1316549.5000 - val accuracy: 0.7234
Epoch 66/100
val loss: -1330783.8750 - val accuracy: 0.7234
Epoch 67/100
val loss: -1344791.1250 - val accuracy: 0.7234
Epoch 68/100
val loss: -1359089.2500 - val accuracy: 0.7234
Epoch 69/100
val loss: -1373541.0000 - val accuracy: 0.7234
Epoch 70/100
val_loss: -1388008.6250 - val_accuracy: 0.7234
Epoch 71/100
val loss: -1402453.5000 - val accuracy: 0.7234
Epoch 72/100
val loss: -1417268.3750 - val accuracy: 0.7234
Epoch 73/100
val loss: -1432073.5000 - val accuracy: 0.7234
Epoch 74/100
val loss: -1447196.8750 - val accuracy: 0.7234
Epoch 75/100
```

```
val_loss: -1462295.5000 - val_accuracy: 0.7234
Epoch 76/100
val loss: -1477444.3750 - val accuracy: 0.7234
Epoch 77/100
val loss: -1492718.0000 - val accuracy: 0.7234
Epoch 78/100
val loss: -1508000.1250 - val accuracy: 0.7234
Epoch 79/100
val loss: -1523874.7500 - val accuracy: 0.7234
Epoch 80/100
val loss: -1539401.7500 - val accuracy: 0.7234
Epoch 81/100
val loss: -1555277.7500 - val accuracy: 0.7234
Epoch 82/100
val loss: -1571173.1250 - val accuracy: 0.7234
Epoch 83/100
val loss: -1586805.2500 - val accuracy: 0.7234
Epoch 84/100
val loss: -1602683.8750 - val accuracy: 0.7234
Epoch 85/100
val loss: -1618731.8750 - val accuracy: 0.7234
Epoch 86/100
```

```
val_loss: -1634839.8750 - val_accuracy: 0.7234
Epoch 87/100
val loss: -1650720.6250 - val accuracy: 0.7234
Epoch 88/100
val loss: -1667040.0000 - val accuracy: 0.7234
Epoch 89/100
val loss: -1683222.5000 - val accuracy: 0.7234
Epoch 90/100
val loss: -1699956.3750 - val accuracy: 0.7234
Epoch 91/100
val loss: -1716632.1250 - val accuracy: 0.7234
Epoch 92/100
val loss: -1733106.0000 - val accuracy: 0.7234
Epoch 93/100
val loss: -1749683.3750 - val accuracy: 0.7234
Epoch 94/100
val loss: -1766571.7500 - val accuracy: 0.7234
Epoch 95/100
val loss: -1783625.3750 - val accuracy: 0.7234
Epoch 96/100
val loss: -1800817.8750 - val accuracy: 0.7234
Epoch 97/100
```

```
val_loss: -1818068.7500 - val_accuracy: 0.7234
Epoch 98/100
val loss: -1835457.7500 - val accuracy: 0.7234
Epoch 99/100
val_loss: -1853025.7500 - val_accuracy: 0.7234
Epoch 100/100
val_loss: -1870389.7500 - val_accuracy: 0.7234
DTC.predict([[50,1,1.2,0.8,150,70,80,7.2,3.4,0.8]])
array([2])
RFmodel.predict([[50,1,1.2,0.8,150,70,80,7.2,3.4,0.8]])
array([1])
classifier.save("liver.h5")
y_test =(y_test>0.5)
y_test
355 True
407 True
90 True
402 True
268 True
  ...
516 True
305 True
167 True
312 True
329 True
Name: Dataset, Length: 117, dtype: bool
```

```
def predict exit(sample value):
sample_value =np.array(sample_value)
sample_value =sample_value.reshape(1,-1)
sample_value =scale(sample_value)
return classifier.predict(sample_value)
sample value=[[50,1,1.2,0.8,150,70,80,7.2,3.4,0.8]]
if predict_exit(sample_value)>0.5:
print('Prediction: Liver Patient')
else:
print('Prediction: Healthy')
1/1 [=======] - 0s 105ms/step
Prediction: Liver Patient
1/1 [=======] - 0s 24ms/step
Prediction: Liver Patient
Milestone 5:
acc_smote=[['KNN
Classifier',KNN],['RandomForestClassifier',RFaccuracy],['DecisionTreeClassifier',DTCaccuracy],['LogisticRegr
ession',LRaccuracy]]
Liverpatient pred=pd.DataFrame(acc smote,columns=['classification models','accuracy score'])
Liverpatient_pred
```

	classification models	accuracy_score
0	KNN Classifier	0.555556
1	RandomForestClassifier	0.709402
2	DecisionTreeClassifier	0.683761
3	LogisticRegression	0.641026

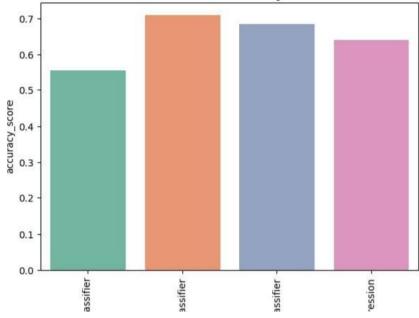
plt.figure(figsize=(7,5))

plt.xticks(rotation=90)

plt.title('Classification models & accuracy scores after SMOTE',fontsize=18)

sns.barplot(x="classification models",y="accuracy\_score",data=Liverpatient\_pred,palette="Set2")

## Classification models & accuracy scores after SMOTE



 $from \ sklearn. ensemble \ import \ ExtraTrees Classifier$ 

model1=ExtraTreesClassifier()

model1.fit(x,y)

```
model1.feature_importances

array([0.13287739, 0.01991026, 0.09463777, 0.08566328, 0.14577484,

0.13219089, 0.12830574, 0.09378994, 0.09210753, 0.07474237])

dd.pd.DataFrame(model1.feature_importances_,index=x.columns).sort_values(0,ascending=False)

dd

data.plot(kind='barh',figsize=(7,6))

plt.title("FEATURE IMPORTANCE",fontsize=14)
```



import joblib

joblib.dump(RFmodel,'liver\_analysis\_1.pkl')

```
['liver_analysis_1.pkl']

data.plot(kind='barh',figsize=(6,5))

plt.title("FEATURE IMPORTANCE",fontsize=14)
```

## Milestone6

```
import numpy as np
import pickle
import os
app=Flask(_name_)
@app.route('/')
def home():
return render_template('home.html')
@app.route('/predict')
def index():
return render_template("index.html")
@app.route('/data_predict',methods=['POST'])
def predict():
  age = request.form['age']
  gender=request.form['gender']
  tb = request.form['tb']
  db = request.form['db']
  ap = request.form['ap']
  aa1 = request.form['aa1']
```

```
aa2 = request.form['aa2']
  tp = request.form['tp']
  a = request.form['a']
  agr = request.form['agr']
  data = [[float(age), float(gender), float(db), float(ap), float(aa1), float(aa2), float(tp), float(agr)]
  model=pickle.load(open(os.path.join('c:Users/91630/Desktop/liver patient/Liver Patient Analysis/Flask
app,pkl_objects','liver_analysis_1.pkl'),'rb'))
  prediction= model.predict(data)[0]
  if (prediction == 1):
   return render_template('noChance.html',prediction='You have a liver desease problem,You must and should
consult a doctor. Take care')
  else:
   return render_template('chance.html', prediction='You dont have a liver desease problem')
if___name___=='_main_':
  app.run(debug=True)
```