ARIGNAR ANNA GOVERNMENT ARTS COLLEGE

VILLUPURAM - 605 602.



DEPARTMENT OF COMPUTER APPLICATIONS

MACHINE LEARNING WITH PYTHON

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

Learning

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A(9E9E4F14F07DEF59D674DB495A7853

86)

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).

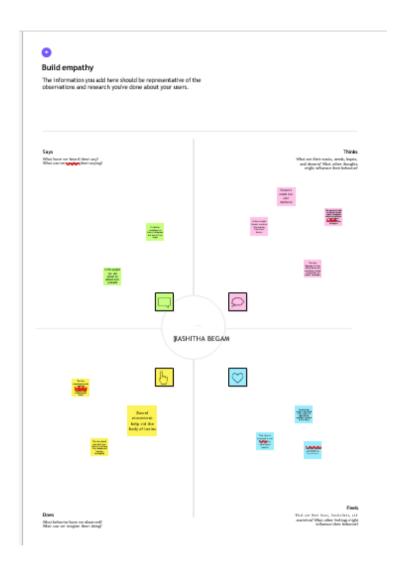
Diagnosis of liver diseases can be divided into three stages i.e., the first stage is liver inflammation, the second is liver scarring (cirrhosis), and the final stage is liver cancer or failure. Since these scenarios are present in liver disease, early prediction is significant to provide better health 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 (Shaheamlunget al., 2020), which is very expensive, particularly in New Zealand (Hepatitis C, 2021). Apparently, in New Zealand, 35 40% of thepopulation 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>

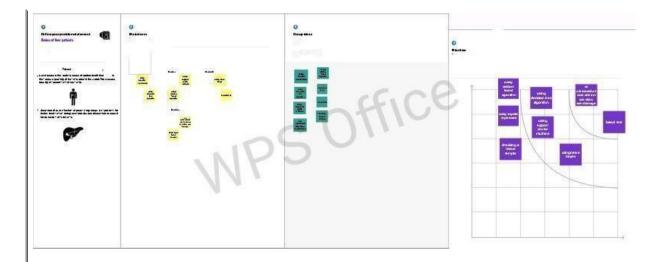
Liver function tests can be used to: Screen for liver infections, such as hepatitis. Monitor the progression of a disease, such as viral or alcoholic hepatitis, and determine how well a treatment is working. Measure the severity of a disease, particularly scarring of the liver (cirrhosis)

Problem Definetion & Design Thinking		

Empathy map



Ideation & brainstorming n	nap



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;
}
```

```
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 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 • Specify

Index.html Source code

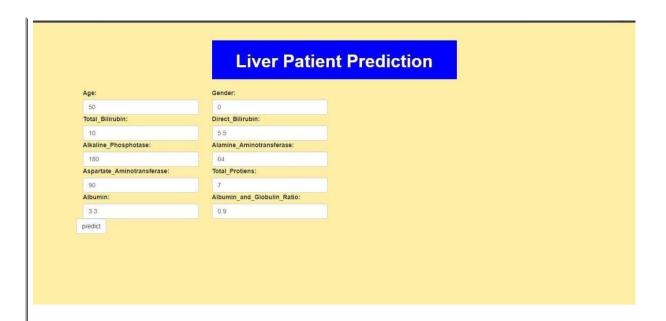
```
<!DOCTYPE html>
<html>
<head>
<title>Liver Patient Analysis</title>
<!-- Latest Compiled and minified CSS -->
<Link rel="stylesheet" href="https://maxcdn.bootstrapcdn.com/bootstrap/3.3.7/css/bootstrap.min.css">
<style type="text/css">
body{ background-color:
#ffcf0059;
}
.page-header{
background-color:
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:</label>
<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:</label>
<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 treatments for 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
2	Total_Bilirubin	583 non-null	float64
3	Direct_Bilirubin	583 non-null	float64
4	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
dty	pes: float64(5), int64(5), object(1)	
mer	mory usage: 50.2+ KB		
data	a.isnull().any()		
Age		False	
Gen	nder	False	
Tota	al_Bilirubin	False	
Dire	ect_Bilirubin	False	
Alka	aline_Phosphotase	False	
Alar	mine_Aminotransferase	False	
Asp	artate_Aminotransferase	False	
Tota	al_Protiens	False	
Albu	umin	False	
Albı	umin_and_Globulin_Ratio	True	

Dataset	False
dtype: bool	
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	4
Dataset	0
dtype: int64	
data['Albumin_and_Globulin_Ratio'] =	
data['Albumin_and_Globulin_Ratio'].fil	llna(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	
LabelEncoder lc = LabelEncoder()	

Milestone 3:

lc.fit_transform(data['Gender'])

data.describe()

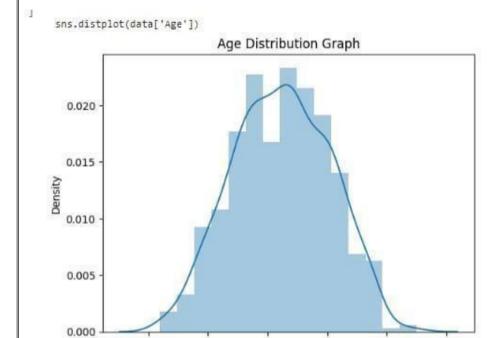
data['Gender'] =

	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	2000.000000	4929.000000	9.600000	5.500000	2.800000	2.000000

sns.distplot(data['Age'])

plt.title('Age Distribution Graph')

plt.show()



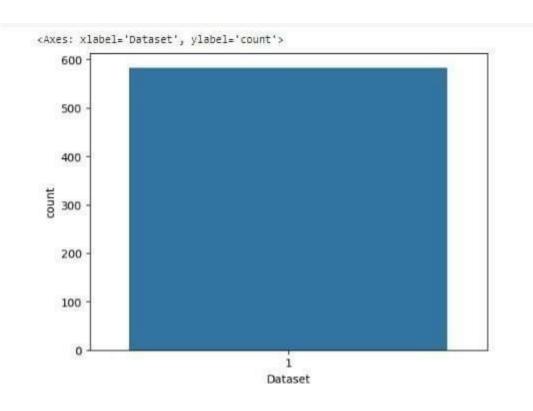
40

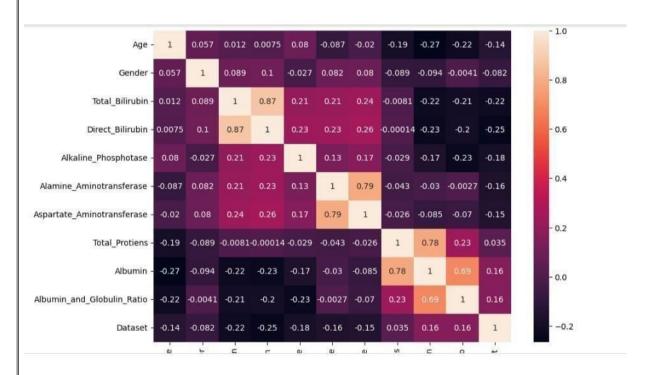
Age

60

80

100





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)

```
* RandomForestClassifier
RandomForestClassifier()
```

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
val loss:
-593204.0625 - val_accuracy: 0.7234
Epoch 2/100
val loss:
-601997.3125 - val accuracy: 0.7234
Epoch 3/100
val loss:
-610722.8125 - val accuracy: 0.7234
Epoch 4/100
val loss:
-619532.7500 - val accuracy: 0.7234
Epoch 5/100
val loss:
-628574.0000 - val accuracy: 0.7234
Epoch 6/100
val loss:
-637505.2500 - val accuracy: 0.7234
Epoch 7/100
```

4/4 [======================] - 0s 41ms/step - loss: -717873.1875 - accuracy: 0.7016 - val_loss: -646466.5625 - val_accuracy: 0.7234
Epoch 8/100
4/4 [======================] - 0s 52ms/step - loss: -727852.1250 - accuracy: 0.7016 - val_loss: -655572.5625 - val_accuracy: 0.7234
Epoch 9/100

```
4/4 [=======] - 0s
                             - loss:
                                        - accuracy: 0.7016 - val loss:
       - val accuracy: 0.7234
  [=======] - Os
                             - loss:
                                        - accuracy: 0.7016 -
                        29ms/step
                              -738061.7500
-664791.8125
Epoch 10/100
4/4
                        24ms/step -748137.7500
                                                 val loss:
-674201.8125 - val_accuracy: 0.7234
Epoch 11/100
val loss:
-683482.2500 - val accuracy: 0.7234
Epoch 12/100
val_loss:
-692927.8125 - val accuracy: 0.7234
Epoch 13/100
val loss:
-702565.5625 - val accuracy: 0.7234
Epoch 14/100
val loss:
-712313.8125 - val accuracy: 0.7234
Epoch 15/100
val_loss:
-721993.8125 - val accuracy: 0.7234
Epoch 16/100
                                        - accuracy: 0.7016 -
                     - 0s
                             - loss:
Epoch
```

```
4/4 [=======] - 0s
                              - loss:
                                         - accuracy: 0.7016 - val loss:
       - val accuracy: 0.7234
  [=======] - 0s
                              - loss:
                                         - accuracy: 0.7016 -
-731739.5625 - val accuracy: 0.7234
Epoch 17/100
-741573.6875 - val accuracy: 0.7234
Epoch 18/100
4/4 [===============] - 0s 43ms/step - loss: -834615.9375 - accuracy: 0.7016 - val loss:
-751650.8125 - val accuracy: 0.7234
Epoch 19/100
20ms/step
                                  -846041.3125
                                                    val loss:
-761723.0000 - val accuracy: 0.7234
   20/100
                        27ms/step
                                  -857278.9375
-771933.6875
Epoch 21/100
4/4
                        20ms/step
                                  -868970.2500
                                                    val_loss:
-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
                      - 0s
                              - loss:
                                         - accuracy: 0.7016 -
Epoch
```

```
4/4 [=======] - 0s
                        - loss:
                                - accuracy: 0.7016 - val loss:
     - val accuracy: 0.7234
 [=======] - 0s
                        - loss:
                                - accuracy: 0.7016 -
Epoch 25/100
val loss:
-824129.3750 - val accuracy: 0.7234
Epoch 26/100
val loss:
-834718.0625 - val_accuracy: 0.7234
Epoch 27/100
val loss:
-845578.3750 - val accuracy: 0.7234
Epoch 28/100
val loss:
-856461.6250 - val_accuracy: 0.7234
Epoch 29/100
val loss:
-867335.2500 - val_accuracy: 0.7234
Epoch 30/100
val_loss:
-878283.5625 - val_accuracy: 0.7234
  31/100
                   18ms/step
                          -988350.1875
val_loss:
Epoch
0.7234
Epoch
```

```
4/4 [=======] - 0s
                           - loss:
                                     - accuracy: 0.7016 - val loss:
      - val accuracy: 0.7234
 [=======] - Os
                           - loss:
                                    - accuracy: 0.7016 -
-889272.2500
Epoch 32/100
4/4
                     16ms/step
                             -1000706.9375
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
                    - 0s
                           - loss:
                                 accuracy: 0.7016 -
Epoch
```

```
val accuracy: 0.7234
Epoch
4/4 [======] - 0s
                       - loss: - accuracy: 0.7016 val loss: -
val accuracy: 0.7234
                     14ms/step
                             -1129659.7500
    -1016444.1875
   43/100
                     13ms/step -1143343.8750
    -1028518.8125
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
val_loss:
           - val accuracy: 0.7234
Epoch
4/4 [========== 0.7016 val loss: - accuracy: 0.7016 val loss: -
val accuracy: 0.7234
```

```
4/4 [=======] - 0s
                           - loss: - accuracy: 0.7016 val loss: -
val accuracy: 0.7234
Epoch
4/4 [=======] - 0s
                           - loss: - accuracy: 0.7016 val loss: -
val accuracy: 0.7234
Epoch 50/100
val_loss: -1115452.1250 - val_accuracy: 0.7234
Epoch 51/100
-1128213.5000
   52/100
                         19ms/step
                                   -1268405.0000
    -1141175.3750
   53/100
                         13ms/step
                                   -1283378.7500
    -1153975.8750
   54/100
                         13ms/step
                                  -1297842.6250
     -1166885.2500
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_accuracy: 0.7234
val_loss:
Epoch
4/4 [=======] - 0s
                           - loss: - accuracy: 0.7016 val loss: -
```

val_accuracy: 0.7234

```
val accuracy: 0.7234
Epoch
val accuracy: 0.7234
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
-1274690.7500
  63/100
                14ms/step
                      -1433544.1250
   -1288476.2500
  64/100
                19ms/step
                      -1449153.0000
val_loss:
    - val accuracy: 0.7234
Epoch
val accuracy: 0.7234
Epoch
```

```
val accuracy: 0.7234
Epoch
val accuracy: 0.7234
   -1302399.7500
  65/100
                 13ms/step
                        -1464224.3750
   -1316549.5000
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:
    - val accuracy: 0.7234
Epoch
val accuracy: 0.7234
Epoch
```

```
4/4 [=======] - 0s
                          - loss: - accuracy: 0.7016 val loss: -
val accuracy: 0.7234
Epoch
4/4 [=======] - 0s
                          - loss: - accuracy: 0.7016 val loss: -
val accuracy: 0.7234
val_loss: -1417268.3750 - val_accuracy: 0.7234
Epoch 73/100
-1432073.5000
   74/100
                        13ms/step
                                 -1610046.1250
    -1447196.8750
   75/100
                        20ms/step
                                 -1627160.6250
    -1462295.5000
   76/100
                        14ms/step
                                 -1644354.2500
    -1477444.3750
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
            - val_accuracy: 0.7234
val_loss:
Epoch
4/4 [=======] - 0s
                          - loss: - accuracy: 0.7016 val loss: -
```

val accuracy: 0.7234

```
val accuracy: 0.7234
Epoch
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
-1602683.8750
  85/100
                 14ms/step
                       -1801669.3750
   -1618731.8750
  86/100
                 20ms/step
                       -1819357.3750
   -1634839.8750
  87/100
val_loss:
        - val _accuracy: 0.7234
Epoch
val accuracy: 0.7234
```

```
4/4 [=======] - 0s
                      - loss: - accuracy: 0.7016 val loss: -
val accuracy: 0.7234
Epoch
4/4 [=======] - 0s
                      - loss: - accuracy: 0.7016 val loss: -
val accuracy: 0.7234
                    20ms/step
                            -1838195.5000
   -1650720.6250
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
val_loss:
     - val accuracy: 0.7234
Epoch
val accuracy: 0.7234
Epoch
```

```
val_loss: - val_accuracy: 0.7234

Epoch

4/4 [========] - 0s - loss: - accuracy: 0.7016 val_loss: - val_accuracy: 0.7234

Epoch
```

```
4/4 [=======] - Os
                               - loss: - accuracy: 0.7016 val loss: - val accuracy:
0.7234
Epoch
4/4 [=======] - 0s
                               - loss: - accuracy: 0.7016 val loss: - val accuracy:
0.7234
                            13ms/step
                                       -2024367.1250
     -1818068.7500
    98/100
                            14ms/step
                                       -2043511.3750
     -1835457.7500
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], ['LogisticRegression', 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

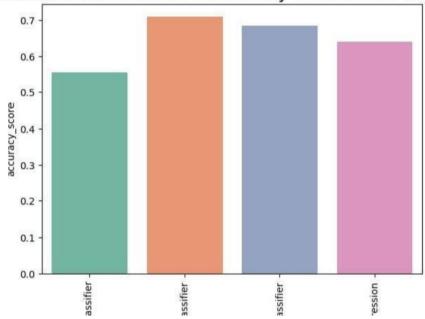
plt.figure(figsize=(7,5))

plt.xticks(rotation=90)

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

 $sns.barplot (x = "classification models", y = "accuracy_score", data = Liverpatient_pred, palette = "Set2")$





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

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

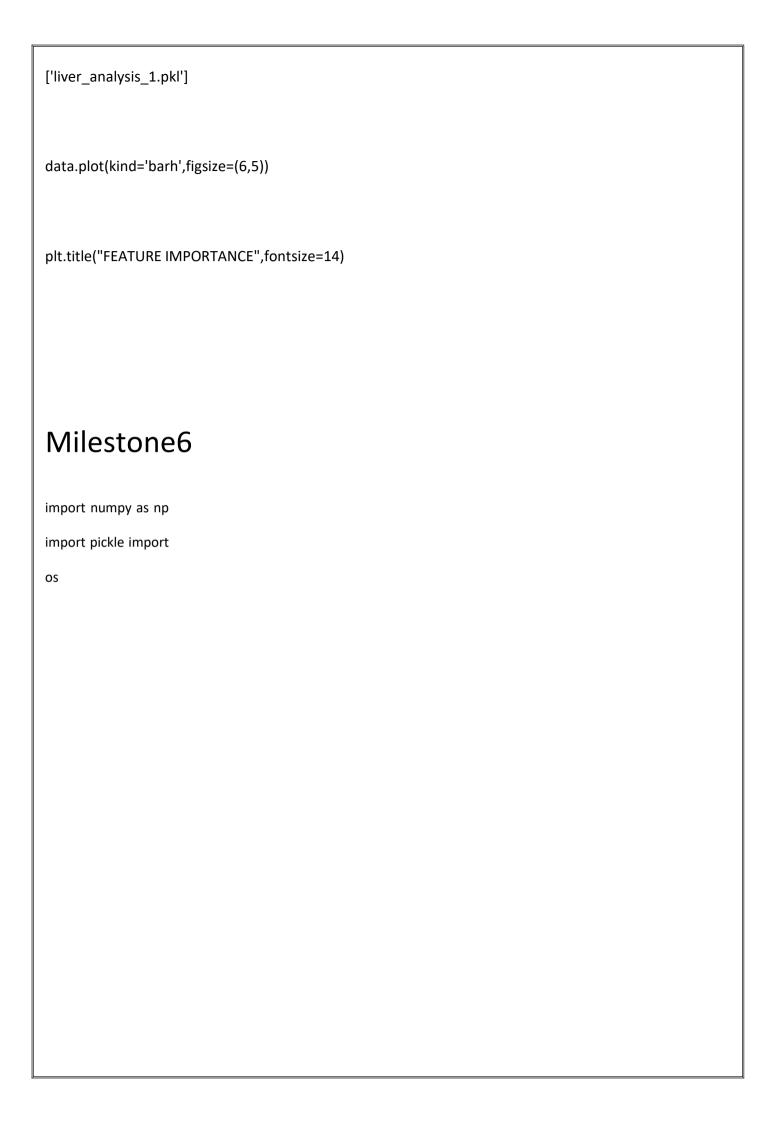
data.plot(kind='barh',figsize=(7,6))

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



import joblib

joblib.dump(RFmodel,'liver_analysis_1.pkl')



```
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(a),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)
```