

A Review of Liver Patient Analysis Methods Using Machine Learning

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1.1 Overview

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.

About the Project:

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.

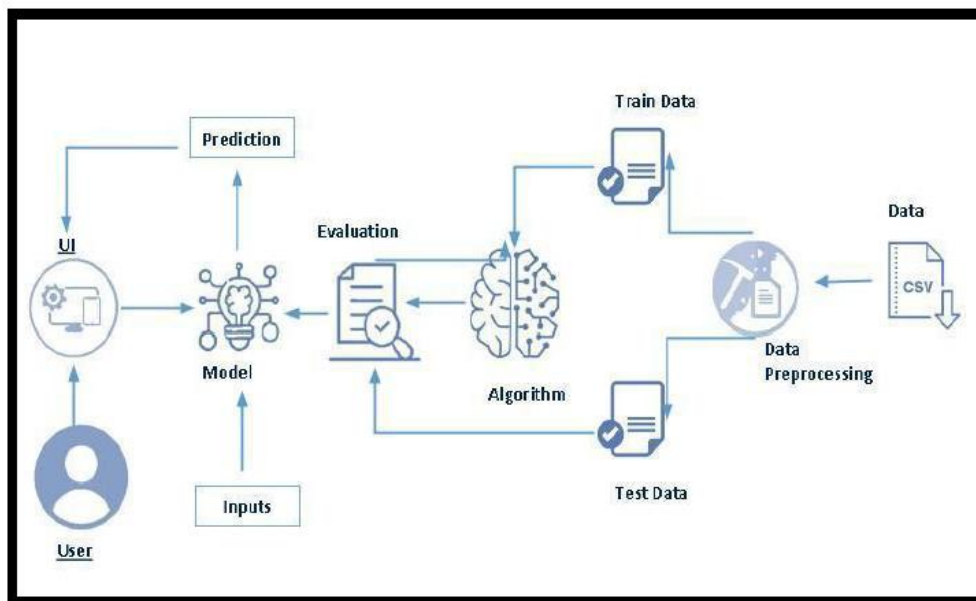
A Project Description:

In India, delays in diagnosing diseases are a major problem due to a lack of medical professionals. The typical scenario, which is mainly in rural and slightly urban areas:

1. A patient who sees a doctor with certain symptoms.
2. The doctor will perform some tests, such as blood and urine tests, depending on the symptoms.
3. The patient undergoes the above tests in the analytical laboratory.
4. The patient takes the reports back to the hospital, where they are examined and diagnosed.

The goal of this project is to reduce some of the delays caused by unnecessary detours between the hospital and the pathology laboratory. Historically, work has been done to detect the onset of heart disease, such as Parkinson's, and machine learning algorithms have been developed to predict liver disease.

Technical Architecture:

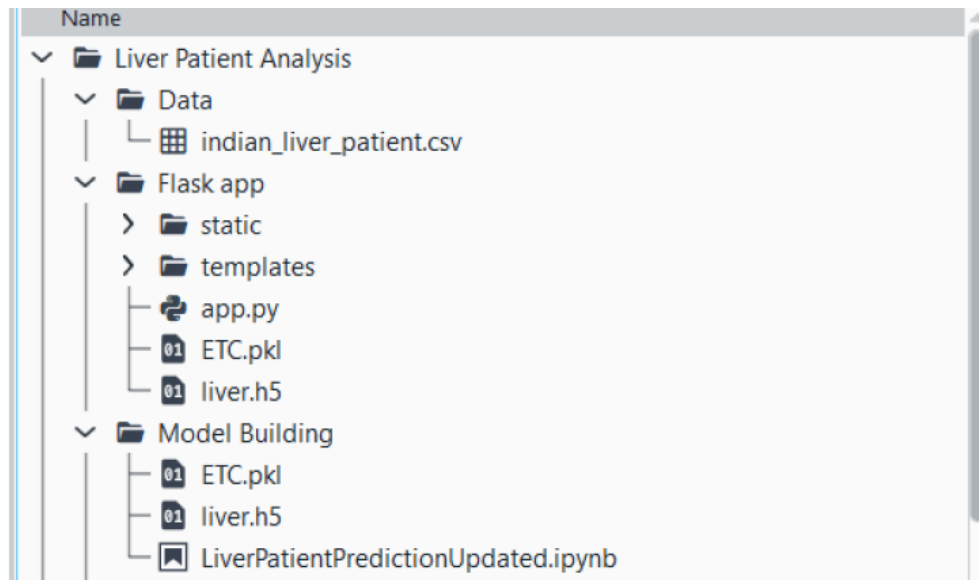


Project Flow:

- 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

Project Structure:

Create the Project folder which contains files as shown below



We are building a flask application which needs HTML pages stored in the templates folder and python script app.py for scripting.

- ETC.pkl is our saved model. Further we will use this model for flask integration.
- Training folder contains a model training file.

1.2 Purpose

The goal of this project is to reduce some of the delays caused by unnecessary detours between the hospital and the pathology laboratory.

Problem Definition & Design Thinking

2.1 Empathy Map



2.2 Ideation & Brainstorming Map



RESULT

```
import pandas as pd
import numpy as np
import plotly.express as px
import matplotlib.pyplot as plt
from google.colab import files
import io
import seaborn as sns
from sklearn.ensemble import VotingClassifier
from sklearn.linear_model import LogisticRegression
from sklearn.linear_model import LinearRegression
from sklearn.svm import SVC
from sklearn.naive_bayes import MultinomialNB
from sklearn.tree import DecisionTreeClassifier
from sklearn.preprocessing import StandardScaler
from sklearn.neighbors import KNeighborsClassifier
from sklearn.metrics import accuracy_score, mean_absolute_error, r2_score
from sklearn.model_selection import train_test_split
from sklearn.metrics import confusion_matrix, classification_report, accuracy_score
from sklearn.ensemble import RandomForestClassifier, ExtraTreesClassifier, AdaBoostClassifier, BaggingClassifier, GradientBoostingClassifier
import xgboost as xgb
%matplotlib inline
```

[5] data=files.upload()

Choose Files No file chosen

Upload widget is only available when the cell has been executed in the current browser session. Please rerun this cell to enable.

Liver_Patient.ipynb - Collaborator x +

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Liver_Patient.ipynb ☆

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data=pd.read_csv('/content/indian_liver_patient.csv.csv')

[7] data.head()

	Age	Gender	Total_Bilirubin	Direct_Bilirubin	Alkaline_Phosphotase	Alamine_Aminotransferase	Aspartate_Aminotransferase	Total_Protiens	Albumin	Albumin_and_Globulin_Ratio
0	65	Female	0.7	0.1	187	16	18	6.8	3.3	
1	62	Male	10.9	5.5	699	64	100	7.5	3.2	
2	62	Male	7.3	4.1	490	60	68	7.0	3.3	
3	58	Male	1.0	0.4	182	14	20	6.8	3.4	
4	72	Male	3.9	2.0	195	27	59	7.3	2.4	

[8] 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
```

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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
dtypes: float64(5), int64(5), object(1)
memory usage: 50.2+ KB
```

data.isnull().any()

	Age	Gender	Total_Bilirubin	Direct_Bilirubin	Alkaline_Phosphotase
	False	False	False	False	False

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data.isnull().any()

Age	False
Gender	False
Total_Bilirubin	False
Direct_Bilirubin	False
Alkaline_Phosphotase	False
Alamine_Aminotransferase	False
Aspartate_Aminotransferase	False
Total_Protiens	False
Albumin	False
Albumin_and_Globulin_Ratio	True
Dataset	False
dtype: bool	

[10] 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

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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.fillna(data['Albumin_and_Globulin_Ratio'].mean(), inplace=True)

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

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```
data.Albumin_and_Globulin_Ratio.fillna(data['Albumin_and_Globulin_Ratio'].mean(),inplace=True)
data.isnull().sum()
```

```
Age      0
Gender    0
Total_Bilirubin    0
Direct_Bilirubin    0
Alkaline_Phosphatase    0
Alamine_Aminotransferase    0
Aspartate_Aminotransferase    0
Total_Protiens    0
Albumin    0
Albumin_and_Globulin_Ratio    0
Dataset    0
dtype: int64
```

```
[12] data.head()
```

	Age	Gender	Total_Bilirubin	Direct_Bilirubin	Alkaline_Phosphatase	Alamine_Aminotransferase	Aspartate_Aminotransferase	Total_Protiens	Albumin	Albumin_and_Globulin_Ratio
0	65	Female	0.7	0.1	187	16	18	6.8	3.3	
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```
Albumin_and_Globulin_Ratio
Dataset
dtype: int64
```

```
[12] data.head()
```

	Age	Gender	Total_Bilirubin	Direct_Bilirubin	Alkaline_Phosphatase	Alamine_Aminotransferase	Aspartate_Aminotransferase	Total_Protiens	Albumin	Albumin_and_Globulin_Ratio
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4	72	Male	3.9	2.0	195	27	59	7.3	2.4	

```
[13] 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
```

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Liver_Patient.ipynb

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RAM Disk

Code Text

[12] 2 62 Male 7.3 4.1 490 60 68 7.0 3.3

3 58 Male 1.0 0.4 182 14 20 6.8 3.4

4 72 Male 3.9 2.0 195 27 59 7.3 2.4

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 583 non-null float64
10 Dataset 583 non-null int64
dtypes: float64(5), int64(5), object(1)
memory usage: 50.2+ KB
```

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Liver_Patient.ipynb

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RAM Disk

Code Text

[12] 2 62 Male 7.3 4.1 490 60 68 7.0 3.3

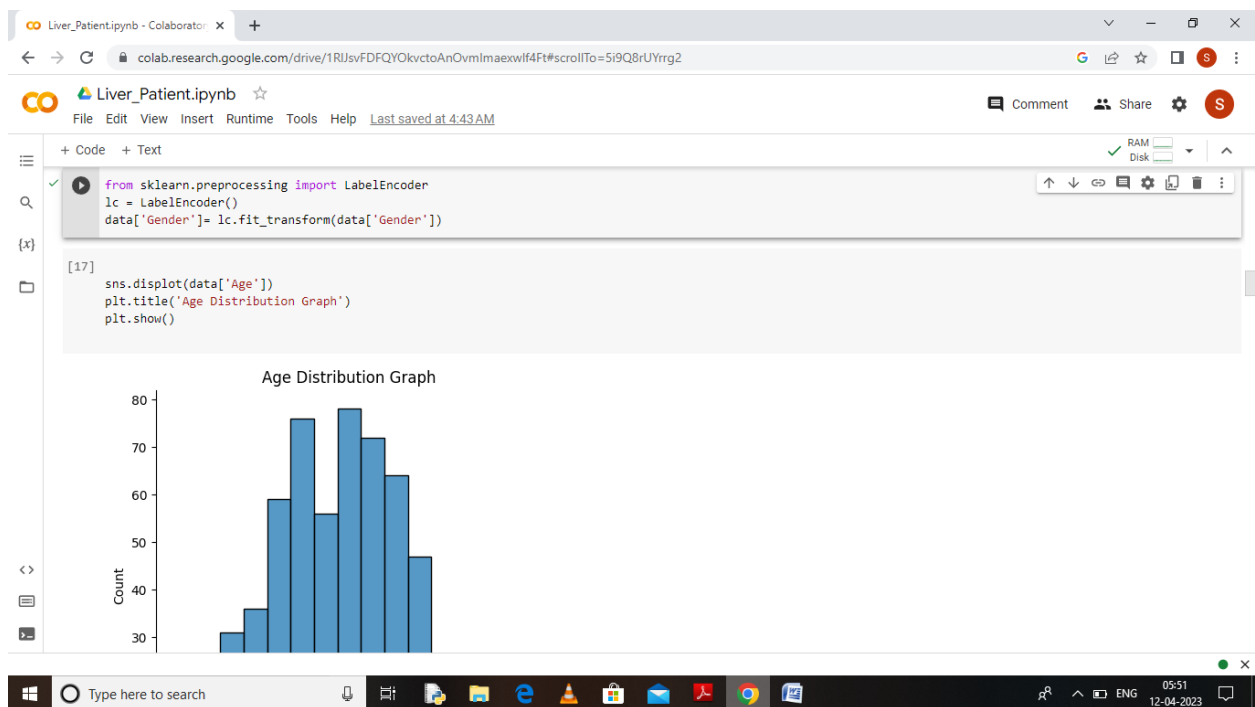
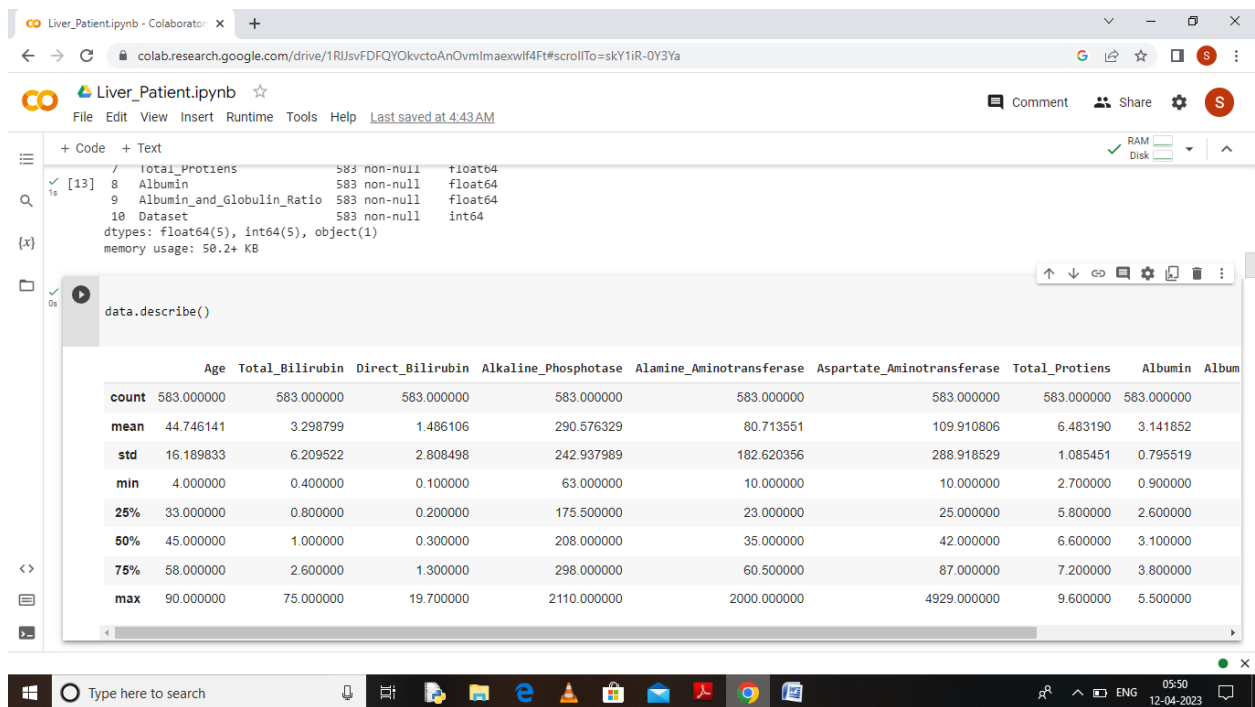
3 58 Male 1.0 0.4 182 14 20 6.8 3.4

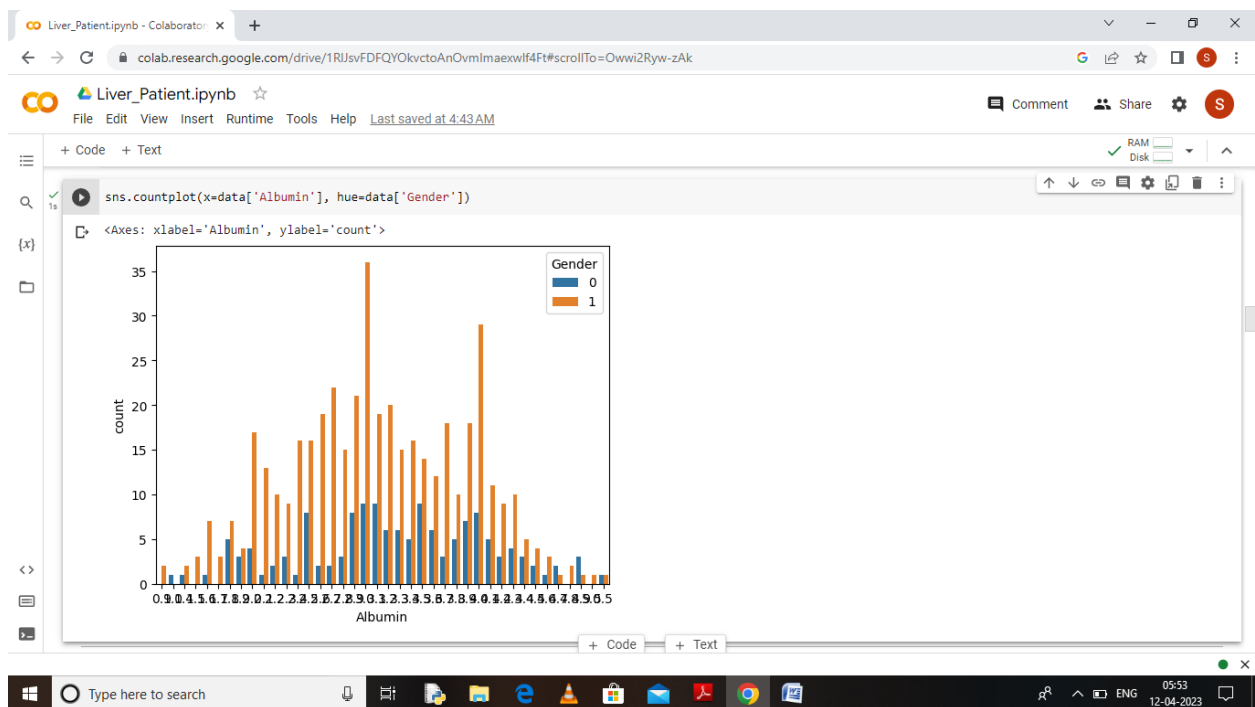
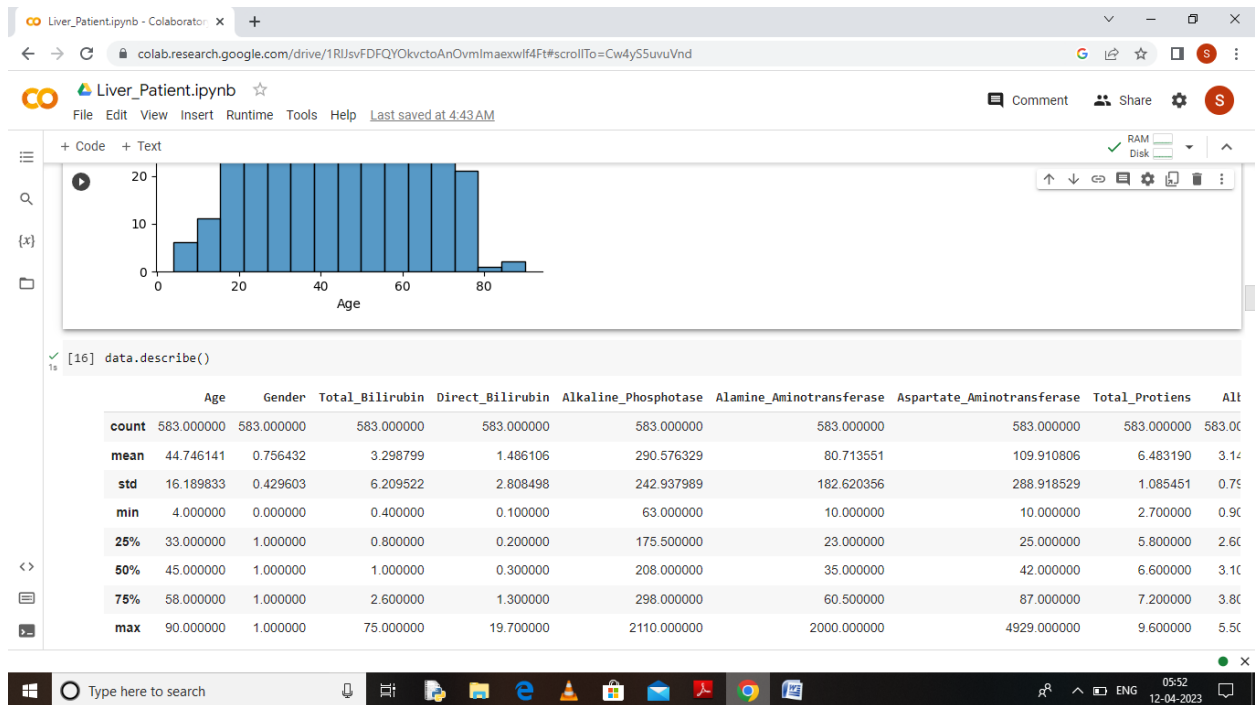
4 72 Male 3.9 2.0 195 27 59 7.3 2.4

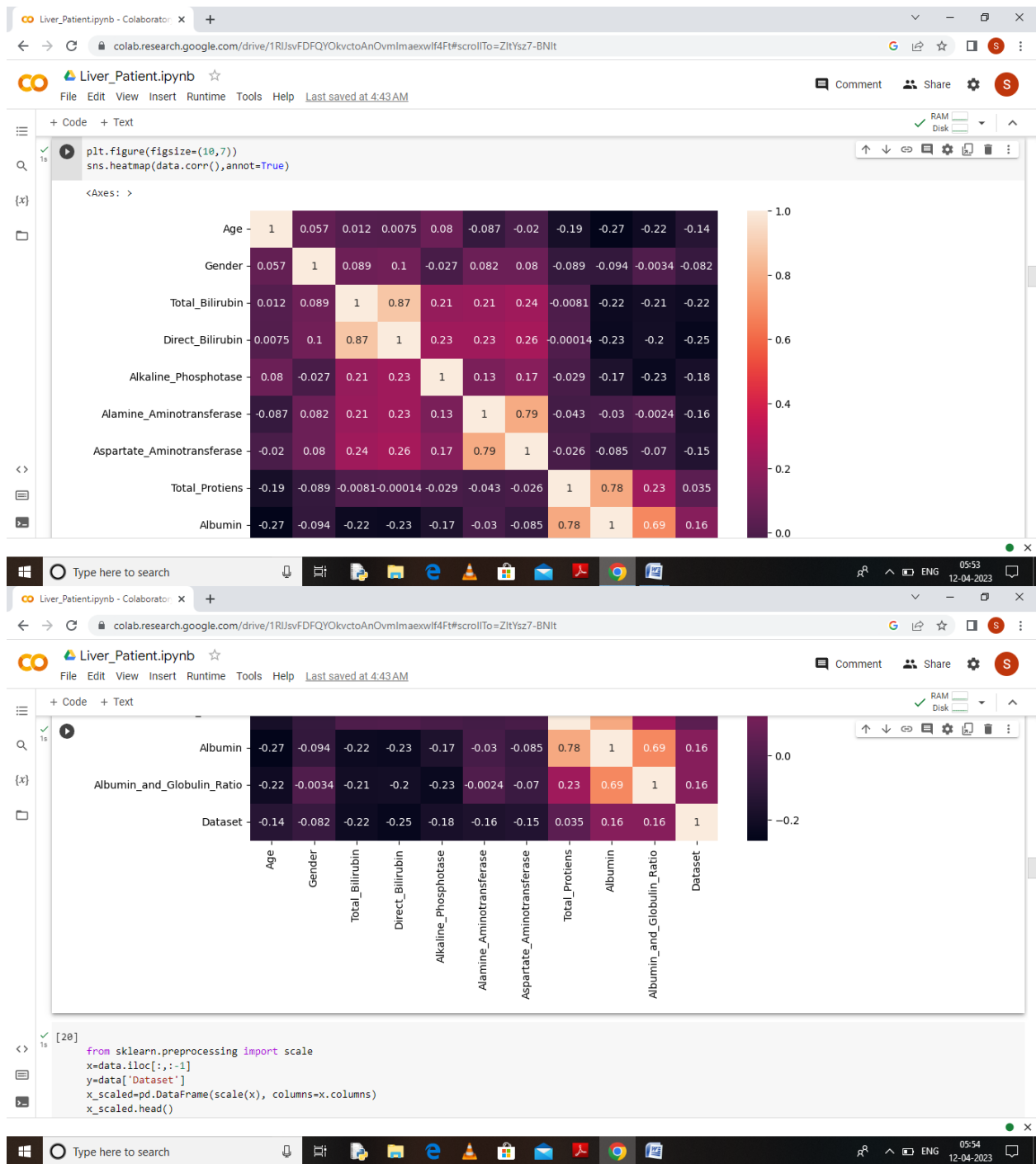
data.info()

```
<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
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3 Direct_Bilirubin 583 non-null float64
4 Alkaline_Phosphotase 583 non-null int64
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7 Total_Protiens 583 non-null float64
8 Albumin 583 non-null float64
9 Albumin_and_Globulin_Ratio 583 non-null float64
10 Dataset 583 non-null int64
dtypes: float64(5), int64(5), object(1)
memory usage: 50.2+ KB
```

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Liver_Patient.ipynb ☆

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✓ [19]

✓ [20]

```
from sklearn.preprocessing import scale
x=data.iloc[:, :-1]
y=data['Dataset']
x_scaled=pd.DataFrame(scale(x), columns=x.columns)
x_scaled.head()
```

	Age	Gender	Total_Bilirubin	Direct_Bilirubin	Alkaline_Phosphatase	Alamine_Aminotransferase	Aspartate_Aminotransferase	Total_Protiens	Albumin	All
0	1.252098	-1.762281	-0.418878	-0.493964	-0.426715	-0.354665	-0.318393	0.292120	0.198969	
1	1.066637	0.567446	1.225171	1.430423	1.682629	-0.091599	-0.034333	0.937566	0.073157	
2	1.066637	0.567446	0.644919	0.931508	0.821588	-0.113522	-0.145186	0.476533	0.198969	
3	0.819356	0.567446	-0.370523	-0.387054	-0.447314	-0.365626	-0.311465	0.292120	0.324781	
4	1.684839	0.567446	0.096902	0.183135	-0.393756	-0.294379	-0.176363	0.753153	-0.933340	

✓ [21] 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)

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Liver_Patient.ipynb - Collaborator x +

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Liver_Patient.ipynb ☆

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✓ [21] 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)

✓ [22] pip install imblearn

```
Looking in indexes: https://pypi.org/simple, https://us-python.pkg.dev/colab-wheels/public/simple/
Collecting imblearn
  Downloading imblearn-0.0-py2.py3-none-any.whl (1.9 kB)
Requirement already satisfied: imbalanced-learn in /usr/local/lib/python3.9/dist-packages (from imblearn) (0.10.1)
Requirement already satisfied: numpy>=1.17.3 in /usr/local/lib/python3.9/dist-packages (from imbalanced-learn->imblearn) (1.22.4)
Requirement already satisfied: joblib>=1.1.1 in /usr/local/lib/python3.9/dist-packages (from imbalanced-learn->imblearn) (1.2.0)
Requirement already satisfied: scipy>=1.3.2 in /usr/local/lib/python3.9/dist-packages (from imbalanced-learn->imblearn) (1.10.1)
Requirement already satisfied: threadpoolctl>=2.0.0 in /usr/local/lib/python3.9/dist-packages (from imbalanced-learn->imblearn) (3.1.0)
Requirement already satisfied: scikit-learn>=1.0.2 in /usr/local/lib/python3.9/dist-packages (from imbalanced-learn->imblearn) (1.2.2)
Installing collected packages: imblearn
Successfully installed imblearn-0.0
```

✓ [23] from imblearn.over_sampling import SMOTE
smote = SMOTE()

✓ from sklearn.preprocessing import StandardScaler

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Liver_Patient.ipynb

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```
from sklearn.preprocessing import StandardScaler
```

```
[25] sc = StandardScaler()
x_train_std = sc.fit_transform(x_train)
x_test_std = sc.transform(x_test)
```

```
[26] from imblearn.over_sampling import SMOTE
smote = SMOTE()
y_train.value_counts()

1    329
2    137
Name: Dataset, dtype: int64
```

```
[27] from sklearn.neighbors import KNeighborsClassifier
```

```
[28] 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
```

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Liver_Patient.ipynb

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```
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
```

```
[29] from sklearn.ensemble import RandomForestClassifier
model1=RandomForestClassifier()
model1.fit(x_train_smote, y_train_smote)
y_predict=model1.predict(x_test)
rfc1=accuracy_score(y_test,y_predict)
rfc1
pd.crosstab(y_test, y_predict)
print(classification_report(y_test, y_predict))
```

	precision	recall	f1-score	support
1	0.82	0.79	0.81	87
2	0.45	0.50	0.48	30
accuracy			0.72	117
macro avg	0.64	0.65	0.64	117
weighted avg	0.73	0.72	0.72	117

```
from sklearn.tree import DecisionTreeClassifier
```

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Liver_Patient.ipynb ☆

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```
from sklearn.linear_model import LogisticRegression
model5=LogisticRegression()
model5.fit(x_train_smote,y_train_smote)
y_predict=model5.predict(x_test)
logit=accuracy_score(y_test,y_predict)
logit
pd.crosstab(y_test,y_predict)
print(classification_report(y_test,y_predict))
```

	precision	recall	f1-score	support
1	0.96	0.63	0.76	87
2	0.47	0.93	0.62	30
accuracy			0.71	117
macro avg	0.72	0.78	0.69	117
weighted avg	0.84	0.71	0.73	117

```
/usr/local/lib/python3.9/dist-packages/sklearn/linear_model/_logistic.py:458: ConvergenceWarning: lbfgs failed to converge (status=1):
STOP: TOTAL NO. of ITERATIONS REACHED LIMIT.

Increase the number of iterations (max_iter) or scale the data as shown in:
https://scikit-learn.org/stable/modules/preprocessing.html
Please also refer to the documentation for alternative solver options:
```

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Liver_Patient.ipynb ☆

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```
import tensorflow.keras
from tensorflow.keras.models import Sequential
from tensorflow.keras.layers import Dense
```

```
[34] # Initialising the ANN
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'])
```

```
[35] # Fitting the ANN to the Training set
model_history = classifier.fit(x_train, y_train, batch_size=100, validation_split=0.2, epochs=100)
```

```
Epoch 1/100
4/4 [=====] - 2s 128ms/step - loss: 20.6170 - accuracy: 0.1586 - val_loss: -2.4078 - val_accuracy: 0.7234
Epoch 2/100
4/4 [=====] - 0s 25ms/step - loss: -5.4694 - accuracy: 0.7016 - val_loss: -9.4452 - val_accuracy: 0.7234
Epoch 3/100
4/4 [=====] - 0s 21ms/step - loss: -12.8357 - accuracy: 0.7016 - val_loss: -15.3244 - val_accuracy: 0.7234
Epoch 4/100
4/4 [=====] - 0s 27ms/step - loss: -19.3103 - accuracy: 0.7016 - val_loss: -20.7903 - val_accuracy: 0.7234
Epoch 5/100
4/4 [=====] - 0s 19ms/step - loss: -25.3693 - accuracy: 0.7016 - val_loss: -26.4392 - val accuracy: 0.7234
```

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Colaboratory interface showing the execution of a neural network training script. The script is titled "Liver_Patient.ipynb" and is running on a Google Colab environment. The code defines a function to fit an ANN to the training set, using a batch size of 100, a validation split of 0.2, and 100 epochs.

The output shows the progress of the training, including the loss and accuracy for each epoch. The training is divided into four groups of 25 epochs each, with a 4/4 split indicating the end of each group. The loss decreases over time, while the accuracy remains relatively stable around 0.7234.

The first screenshot shows the first 12 epochs of training. The second screenshot shows the remaining 12 epochs, from epoch 13 to epoch 24. The training process is complete, and the final loss and accuracy values are displayed.

Epoch 1/100
4/4 [=====] - 2s 128ms/step - loss: 20.6170 - accuracy: 0.1586 - val_loss: -2.4078 - val_accuracy: 0.7234
Epoch 2/100
4/4 [=====] - 0s 25ms/step - loss: -5.4694 - accuracy: 0.7016 - val_loss: -9.4452 - val_accuracy: 0.7234
Epoch 3/100
4/4 [=====] - 0s 21ms/step - loss: -12.8357 - accuracy: 0.7016 - val_loss: -15.3244 - val_accuracy: 0.7234
Epoch 4/100
4/4 [=====] - 0s 27ms/step - loss: -19.3103 - accuracy: 0.7016 - val_loss: -20.7903 - val_accuracy: 0.7234
Epoch 5/100
4/4 [=====] - 0s 19ms/step - loss: -25.3693 - accuracy: 0.7016 - val_loss: -26.4392 - val_accuracy: 0.7234
Epoch 6/100
4/4 [=====] - 0s 19ms/step - loss: -31.6509 - accuracy: 0.7016 - val_loss: -32.4281 - val_accuracy: 0.7234
Epoch 7/100
4/4 [=====] - 0s 26ms/step - loss: -38.8414 - accuracy: 0.7016 - val_loss: -38.7394 - val_accuracy: 0.7234
Epoch 8/100
4/4 [=====] - 0s 20ms/step - loss: -46.4067 - accuracy: 0.7016 - val_loss: -45.7490 - val_accuracy: 0.7234
Epoch 9/100
4/4 [=====] - 0s 27ms/step - loss: -54.4914 - accuracy: 0.7016 - val_loss: -53.7416 - val_accuracy: 0.7234
Epoch 10/100
4/4 [=====] - 0s 27ms/step - loss: -63.7798 - accuracy: 0.7016 - val_loss: -62.7697 - val_accuracy: 0.7234
Epoch 11/100
4/4 [=====] - 0s 24ms/step - loss: -74.4876 - accuracy: 0.7016 - val_loss: -72.9588 - val_accuracy: 0.7234
Epoch 12/100
4/4 [=====] - 0s 25ms/step - loss: -86.1563 - accuracy: 0.7016 - val_loss: -84.6048 - val_accuracy: 0.7234
Epoch 13/100
4/4 [=====] - 0s 22ms/step - loss: -100.1355 - accuracy: 0.7016 - val_loss: -98.0134 - val_accuracy: 0.7234
Epoch 14/100
4/4 [=====] - 0s 27ms/step - loss: -116.7105 - accuracy: 0.7016 - val_loss: -113.3326 - val_accuracy: 0.7234
Epoch 15/100
4/4 [=====] - 0s 23ms/step - loss: -134.2424 - accuracy: 0.7016 - val_loss: -131.3577 - val_accuracy: 0.7234
Epoch 16/100
4/4 [=====] - 0s 25ms/step - loss: -156.2685 - accuracy: 0.7016 - val_loss: -151.8920 - val_accuracy: 0.7234
Epoch 17/100
4/4 [=====] - 0s 19ms/step - loss: -180.5942 - accuracy: 0.7016 - val_loss: -175.4873 - val_accuracy: 0.7234
Epoch 18/100
4/4 [=====] - 0s 22ms/step - loss: -207.7542 - accuracy: 0.7016 - val_loss: -202.6962 - val_accuracy: 0.7234
Epoch 19/100
4/4 [=====] - 0s 16ms/step - loss: -240.5221 - accuracy: 0.7016 - val_loss: -233.4873 - val_accuracy: 0.7234
Epoch 20/100
4/4 [=====] - 0s 16ms/step - loss: -279.2705 - accuracy: 0.7016 - val_loss: -267.8985 - val_accuracy: 0.7234
Epoch 21/100
4/4 [=====] - 0s 14ms/step - loss: -317.2207 - accuracy: 0.7016 - val_loss: -307.9347 - val_accuracy: 0.7234
Epoch 22/100
4/4 [=====] - 0s 14ms/step - loss: -362.6669 - accuracy: 0.7016 - val_loss: -352.7855 - val_accuracy: 0.7234
Epoch 23/100
4/4 [=====] - 0s 21ms/step - loss: -416.4498 - accuracy: 0.7016 - val_loss: -402.2024 - val_accuracy: 0.7234
Epoch 24/100
4/4 [=====] - 0s 19ms/step - loss: -473.1902 - accuracy: 0.7016 - val_loss: -457.9147 - val_accuracy: 0.7234

Liver_Patient.ipynb - Collaborator x

colab.research.google.com/drive/1RJsVDFQYOKvctoAnOvmlmaexwf4Ft#scrollTo=NbQ6_EsH3_v_

Liver_Patient.ipynb

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[35] 4/4 [=====] - 0s 19ms/step - loss: -778.6101 - accuracy: 0.7016 - val_loss: -747.9950 - val_accuracy: 0.7234
Epoch 29/100
4/4 [=====] - 0s 21ms/step - loss: -873.7567 - accuracy: 0.7016 - val_loss: -840.2831 - val_accuracy: 0.7234

[36] model14.predict([[50,1,1.2,0.8,150,70,80,7.2,3.4,0.8]])

/usr/local/lib/python3.9/dist-packages/sklearn/base.py:439: UserWarning: X does not have valid feature names, but DecisionTreeClassifier was fitted
warnings.warn(array([1]))

[37] model11.predict([[50,1,1.2,0.8,150,70,80,7.2,3.4,0.8]])

/usr/local/lib/python3.9/dist-packages/sklearn/base.py:439: UserWarning: X does not have valid feature names, but RandomForestClassifier was fitted
warnings.warn(array([1]))

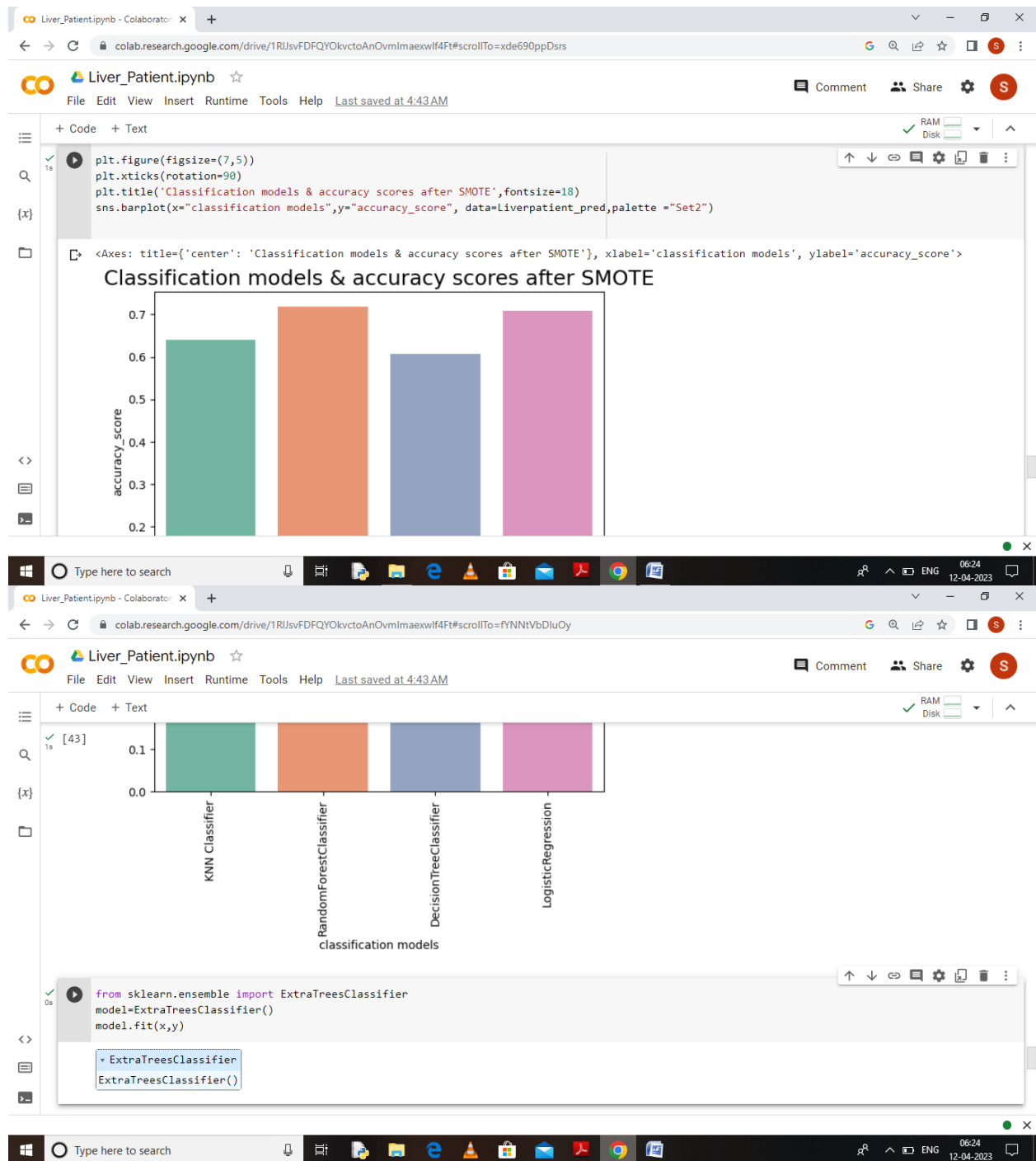
classifier.save("liver.h5")
y_pred = classifier.predict(x_test)

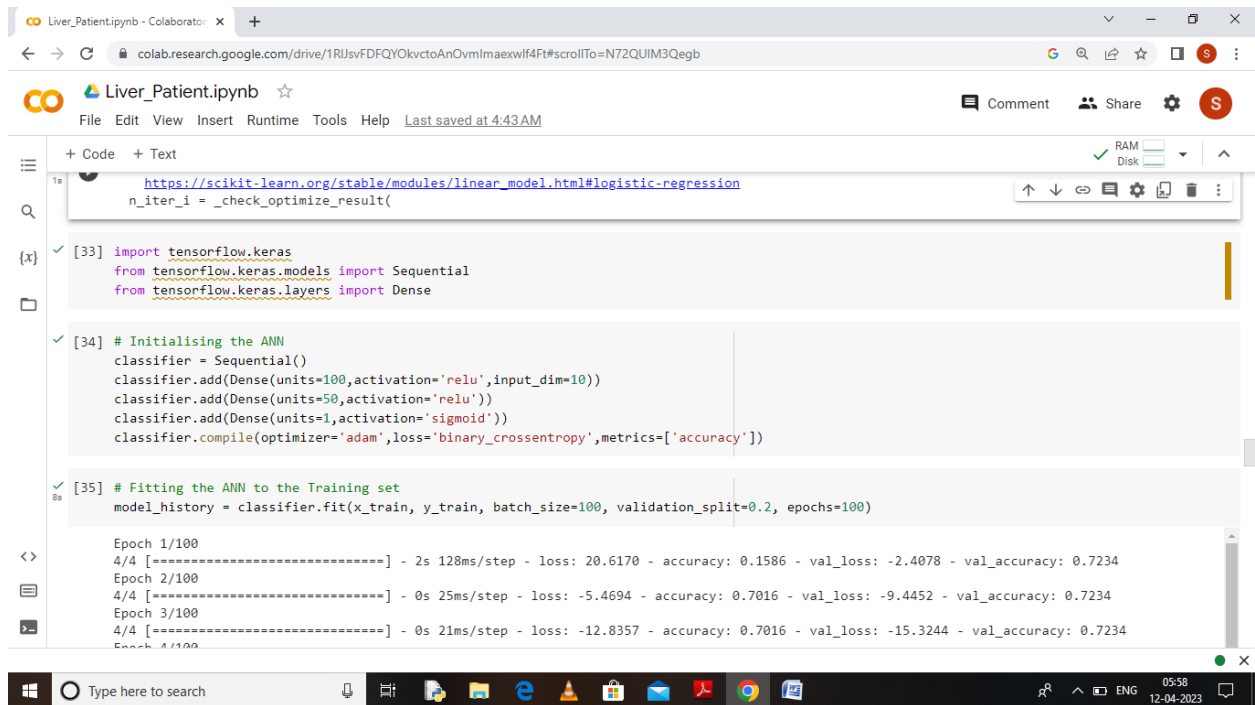
4/4 [=====] - 0s 3ms/step

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```
https://scikit-learn.org/stable/modules/linear_model.html#logistic-regression
n_iter_1 = _check_optimize_result(

[33] import tensorflow.keras
      from tensorflow.keras.models import Sequential
      from tensorflow.keras.layers import Dense

[34] # Initialising the ANN
      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'])

[35] # Fitting the ANN to the Training set
      model_history = classifier.fit(x_train, y_train, batch_size=100, validation_split=0.2, epochs=100)

Epoch 1/100
4/4 [=====] - 2s 128ms/step - loss: 20.6170 - accuracy: 0.1586 - val_loss: -2.4078 - val_accuracy: 0.7234
Epoch 2/100
4/4 [=====] - 0s 25ms/step - loss: -5.4694 - accuracy: 0.7016 - val_loss: -9.4452 - val_accuracy: 0.7234
Epoch 3/100
4/4 [=====] - 0s 21ms/step - loss: -12.8357 - accuracy: 0.7016 - val_loss: -15.3244 - val_accuracy: 0.7234
Epoch 4/100
```

4 ADVANTAGES & DISADVANTAGES

List of advantages and disadvantage

Advantage

A liver biopsy may be done to: **Diagnose a liver problem that can't be otherwise identified with a health care provider's exam, blood tests or imaging studies.** Obtain a sample of tissue from an irregularity found by an imaging study. Determine the severity of liver disease, a process called staging.

Disadvantage

It's possible, and dangerous, to get too much vitamin A. Eating large amounts of liver can lead to symptoms of vitamin A toxicity, which happens when your own liver can't process the excess vitamin A quickly enough. Most doctors recommend that people without vitamin deficiencies eat just one serving of liver per week.

5 APPLICATIONS

Testing the model Define Problem / Problem Understanding

Specify the business problem

Business requirements

Literature Survey

Social or Business Impact.

Data Collection & Preparation

Collect the dataset

Data Preparation

Exploratory Data Analysis

Descriptive statistical

Visual Analysis

Model Building

Training the model in multiple algorithms

6 CONCLUSION

Explore and understand your data

Visualize the data at hand to gain a better intuition — Word cloud, N-gram Bar Chart

Text Cleaning — Word Stemmer and Word Lemmatization

Feature Extraction — Count Vectorizer, Tfidf Vectorizer, Word Embedding

Algorithm — Naive Bayes

Scoring & Metrics — Accuracy, Precision, Recall

7 FUTURE SCOPE

Generally, a **90% spam catch rate** (90 out of 100 spam messages are correctly identified as spam) and a false positive rate of less than 1% (less than 1 legitimate message out of a hundred incorrectly identified as spam) is considered good.

Help prevent similar attacks and respond to changing behaviour.

Time Saving

Cyber Security analyse Pattern

Fraud Detection.

8 APPENDIX

Source Code

```
import pandas as pd
import numpy as np
import plotly.express as px
import matplotlib.pyplot as plt
from google.colab import files
import io
import seaborn as sns
from sklearn.ensemble import VotingClassifier
from sklearn.linear_model import LogisticRegression
from sklearn.linear_model import LinearRegression
from sklearn.svm import SVC
from sklearn.naive_bayes import MultinomialNB
from sklearn.tree import DecisionTreeClassifier
from sklearn.preprocessing import StandardScaler
from sklearn.neighbors import KNeighborsClassifier
from sklearn.metrics import accuracy_score, mean_absolute_error, r2_score
from sklearn.model_selection import train_test_split
from sklearn.metrics import confusion_matrix, classification_report, accuracy_score
from sklearn.ensemble import
RandomForestClassifier, ExtraTreesClassifier, AdaBoostClassifier, BaggingClassifier, GradientBoostingC
lassifier
import xgboost as xgb
# %matplotlib inline

data=files.upload()

data=pd.read_csv('/content/indian_liver_patient.csv.csv')

data.head()

data.info()

data.isnull().any()

data.isnull().sum()

data.Albumin_and_Globulin_Ratio.fillna(data['Albumin_and_Globulin_Ratio'].mean(),inplace=True)
data.isnull().sum()

data.head()
```

```

data.info()

data.describe()

from sklearn.preprocessing import LabelEncoder
lc = LabelEncoder()
data['Gender']= lc.fit_transform(data['Gender'])

data.describe()

sns.displot(data['Age'])
plt.title('Age Distribution Graph')
plt.show()

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

plt.figure(figsize=(10,7))
sns.heatmap(data.corr(),annot=True)

from sklearn.preprocessing import scale
x=data.iloc[:, :-1]
y=data['Dataset']
x_scaled=pd.DataFrame(scale(x), columns=x.columns)
x_scaled.head()

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)

pip install imblearn

from imblearn.over_sampling import SMOTE
smote = SMOTE()

from sklearn.preprocessing import StandardScaler

sc = StandardScaler()
x_train_std = sc.fit_transform(x_train)
x_test_std = sc.transform(x_test)

from imblearn.over_sampling import SMOTE
smote = SMOTE()
y_train.value_counts()

from sklearn.neighbors import KNeighborsClassifier

x_train_smote, y_train_smote=smote.fit_resample(x_train,y_train)

```

```
y_train_smote.value_counts()
```

```
from sklearn.ensemble import RandomForestClassifier
model1=RandomForestClassifier()
model1.fit(x_train_smote, y_train_smote)
y_predict=model1.predict(x_test)
rfc1=accuracy_score(y_test,y_predict)
rfc1
pd.crosstab(y_test, y_predict)
print(classification_report(y_test, y_predict))
```

```
from sklearn.tree import DecisionTreeClassifier
model4=DecisionTreeClassifier()
model4.fit(x_train_smote,y_train_smote)
y_predict=model4.predict(x_test)
dtc1=accuracy_score(y_test,y_predict)
dtc1
pd.crosstab(y_test,y_predict)
print(classification_report(y_test,y_predict))
```

```
from sklearn.neighbors import KNeighborsClassifier
model2=KNeighborsClassifier()
model2.fit(x_train_smote,y_train_smote)
y_predict=model2.predict(x_test)
knn1=(accuracy_score(y_test,y_predict))
knn1
pd.crosstab(y_test,y_predict)
print(classification_report(y_test,y_predict))
```

```
from sklearn.linear_model import LogisticRegression
model5=LogisticRegression()
model5.fit(x_train_smote,y_train_smote)
y_predict=model5.predict(x_test)
logi1=accuracy_score(y_test,y_predict)
logi1
pd.crosstab(y_test,y_test)
print(classification_report(y_test,y_predict))
```

```
import tensorflow.keras
from tensorflow.keras.models import Sequential
from tensorflow.keras.layers import Dense
```

```
# Initialising the ANN
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'])

# Fitting the ANN to the Training set
model_history = classifier.fit(x_train, y_train, batch_size=100, validation_split=0.2, epochs=100)

model4.predict([[50,1,1.2,0.8,150,70,80,7.2,3.4,0.8]])

model1.predict([[50,1,1.2,0.8,150,70,80,7.2,3.4,0.8]])

classifier.save("liver.h5")
y_pred = classifier.predict(x_test)

y_pred
y_pred = (y_pred > 0.5)
y_pred

def predict_exit(sample_value):
    # Convert list to numpy array
    sample_value = np.array(sample_value)
    #Reshape because sample_value contains only 1 record
    sample_value = sample_value.reshape(1, -1)
    #Feature Scaling
    sample_value = scale(sample_value)
    return classifier.predict(sample_value)

#Age-->| Gender-->| Total_Bilirubin-->| Direct_Bilirubin->| Alkaline_phosphatase-->|
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 ')

acc_smote= [['KNN Classifier', knn1], ['RandomForestClassifier', rfc1],
            ['DecisionTreeClassifier',dtc1],['LogisticRegression',logi1]]
Liverpatient_pred=pd.DataFrame(acc_smote, columns= ['classification models','accuracy_score'])
Liverpatient_pred

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 ExtraTreesClassifier
model=ExtraTreesClassifier()
model.fit(x,y)

```

```
model.feature_importances_  
  
dd=pd.DataFrame(model.feature_importances_,index=x.columns).sort_values(0,ascending=False)  
dd  
  
dd.plot(kind='barh',figsize=(7,6))  
plt.title("FEATURE IMPORTANCE",fontsize=14)  
  
import joblib  
joblib.dump(model1, 'ETC.pkl')
```