LIVER PATIENT ANALYSIS USING MACHINE LEARNING

A UG PROJECT PHASE-1 REPORT

Submitted to

JAWAHARLAL NEHRU TECHNOLOGICAL UNIVERSITY, HYDERABAD

In partial fulfillment of the requirements for the award of the degree of

BACHELOR OF TECHNOLOGY

IN

COMPUTER SCIENCE AND ENGINEERING

Submitted by

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(Affiliated to JNTUH, Hyderabad) Bollikunta, Warangal – 506005 **2018–2022**

DEPARTMENT OF COMPUTER SCIENCE AND ENGINEERING VAAGDEVI ENGINEERING COLLEGE

BOLLIKUNTA, WARANGAL – 506005 2018 – 2022



<u>UG PROJECT PHASE-1</u>

This is to certify that the UG Project Phase-1 entitled "LIVER PATIENT ANALYSIS USING MACHINE LEARNING" is being submitted by *B.ANUHYA*(H.NO:18UK1A0505), *M.SINDHU* (H.NO:18UK1A0552), *D.AJAY*(H.NO:18UK1A0570), *J.VAMSHI*(H.NO:18UK1A0577) in partial fulfillment of the requirements for the award of the degree of Bachelor of Technology in Computer Science and Engineering to Jawaharlal Nehru Technological University Hyderabad during the academic year 2021-22, is a record of work carried out by them under the guidance and supervision.

Project Guide
Mr. P. Niranjan Reddy
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Head of the Department Dr. R. Naveen Kumar (Professor)

External

ACKNOWLEDGEMENT

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ABSTRACT

Liver diseases avert the normal function of the liver. Mainly due to the large amount of alcohol consumption liver disease arises. 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. Discovering the existence of liver disease at an early stage is a complex task for the doctors. The main objective of this project is to 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 examines data from liver patients concentrating on relationships between a key list of liver enzymes, proteins, age and gender using them to try and predict the likeliness of liver disease. Here we are building a model by applying various machine learning algorithms find the best accurate model. And integrate to flask based web application. User can predict the disease by entering parameters in the web application.

Keywords – Liver diseases, enzymes, machine learning.

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1. INTRODUCTION

1.1. MOTIVATION:

Liver disease is any disturbance of liver function that causes illness. The liver is responsible for many critical functions within the body and should it become diseased or injured, the loss of those functions can cause significant damage to the body. Liver disease is also referred to as hepatic disease. Liver disease is a broad term that covers all the potential problems that cause the liver to fail to perform its designated functions. Usually, more than 75% or three quarters of liver tissue needs to be affected before a decrease in function occurs.

1.2. PROBLEM DEFINITION:

Liver diseases avert the normal function of the liver. Mainly due to the large amount of alcohol consumption liver disease arises. 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. Discovering the existence of liver disease at an early stage is a complex task for the doctors.

1.3. PROJECT OBJECTIVE:

The main objective of this project is to analyze 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 examines data from liver patients concentrating on relationships between a key list of liver enzymes, proteins, age and gender using them to try and predict the likeliness of liver disease. User can predict the disease by entering parameters in the web application.

1.4. LIMITATIONS OF PROJECT:

Many liver diseases are manageable if you catch them early. Without treatment, however, they can cause permanent damage. The complications of untreated or unmanaged liver disease can lead to cirrhosis, severe scarring that cannot be reversed. If cirrhosis has gone too far, a liver transplant may be your only option. Because some liver diseases can

develop without symptoms, making it a point to schedule annual physicals, along with the typical physical blood work, can help you and your doctor stay one step ahead. Focusing on a nutritious diet, physical exercise, and other healthy lifestyle choices such as limiting alcohol can also help with prevention or management.

1.5. ORGANIZATION OF DOCUMENTATION:

Actually there has been many theoretical projects and several experimental projects individually done based on liver patient analysis and many different algorithms have been developed for forecasting the liver diseases. But Machine Learning (ML) is a type of artificial intelligence (AI) that allows software applications to become more accurate at predicting outcomes without being explicitly programmed to do so. Our aim from the project is to make use of NumPy or pandas libraries in Machine Learning and predict whether a person has Liver disease or not. Secondly, to learn how to visualize the data by using graphs. In the end, we are predicting the liver disease based on the key list of liver enzymes, proteins, age and gender. The prediction is to be done using Machine Learning algorithms and withdrawing the conclusions.

2. PROBLEM STATEMENT

The liver is an organ about the size of a football. It sits just under your rib cage on the right side of your abdomen. The liver is essential for digesting food and ridding your body of toxic substances. Liver disease can be inherited (genetic). Liver problems can also be caused by a variety of factors that damage the liver, such as viruses, alcohol use and obesity. Over time, conditions that damage the liver can lead to scarring (cirrhosis), which can lead to liver failure, a life-threatening condition. But early treatment may give the liver time to heal. Liver failure occurs when your liver isn't working well enough to perform its functions (for example, manufacturing bile and ridding the body of harmful substances). Symptoms include nausea, loss of appetite, and blood in the stool. Treatments include avoiding alcohol and avoiding certain foods.

Other conditions that can lead to liver failure include:

- **Hepatitis A:** Contact with food or water contaminated with the hepatitis A virus, or with a person who's infected with virus, can cause liver inflammation. This type usually goes away on its own.
- **Autoimmune hepatitis:** In this type, your body's immune system, not a virus, attacks your liver and causes inflammation
- **Cirrhosis**: Things like drinking alcohol for many years or having hepatitis scar your liver can make it hard or impossible for your liver to work.
- **Primary sclerosing cholangitis**: This disease slowly damages your bile ducts. It mostly affects young men.
- Oxalosis: This is when your kidneys can't get rid of calcium oxalate crystals through your urine.
- Wilson's disease: People with this rare inherited disease store too much copper in their brain and liver.
- Alpha-1 antitrypsin deficiency: This genetic condition can lead to lung or liver disease.
- **Liver cancer:** People with long-term hepatitis B or hepatitis C often get this.
- **Liver adenoma:** This is when benign liver tumors are on an otherwise healthy liver. This often affects women between ages 20 and 44.

- **Fatty liver disease:** Extra fat cells can build up on your liver. Nonalcoholic fatty liver disease often affects people who are overweight, obese, or have high cholesterol. Alcohol-related fatty liver disease affects heavy drinkers.
- Alcoholic hepatitis: Liver inflammation that results from heavy or long-term drinking.
- Alagille syndrome: A genetic disorder that results in fewer bile ducts than normal in the liver.
- **Primary biliary cholangitis (PBC):** Over time, this disease destroys your small bile ducts. You might still hear it called by its former name, primary biliary cirrhosis.
- Galactosemia: People with this condition can't process galactose, a sugar found in many foods. It can cause liver damage.
- Lysosomal acid lipase deficiency (LAL-D): With this genetic condition, you can't produce an enzyme called lysosomal acid lipase (LAL), which helps your body break down fats and cholesterol in your cells. As a result, fats stay in your liver and cause damage.

Liver diseases avert the normal function of the liver. Mainly due to the large amount of alcohol consumption liver disease arises. 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. Discovering the existence of liver disease at an early stage is a complex task for the doctors. The main objective of this project is to 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.

3. LITERATURE SURVEY

3.1. INTRODUCTION:

The liver is an organ about the size of a football. It sits just under your rib cage on the right side of your abdomen. The liver is essential for digesting food and ridding your body of toxic substances. Liver disease can be inherited (genetic). Liver problems can also be caused by a variety of factors that damage the liver, such as viruses, alcohol use and obesity.

Over time, conditions that damage the liver can lead to scarring (cirrhosis), which can lead to liver failure, a life-threatening condition. But early treatment may give the liver time to heal. Liver disease doesn't always cause noticeable signs and symptoms. If signs and symptoms of liver disease do occur, the may include:

- Skin and eyes that appear yellowish (jaundice)
- Abdominal pain and swelling
- Swelling in the legs and ankles
- Itchy skin
- Dark urine colour
- Pale stool colour
- Chronic fatigue
- Nausea or vomiting
- Loss of appetite
- Tendency to bruise easily

3.2. EXISTING SYSTEM:

The purpose of the present model is to predict the whether a person has a liver disease or not. The model examines data from liver patients concentrating on relationships between a key list of liver enzymes, proteins, age and gender using them to try and predict the likeliness of liver disease. Here we are building a model by applying various machine learning algorithms find the best accurate model. And integrate to flask based web application. User can predict the disease by entering parameters in the web application.

3.3. DISADVANTAGES OF EXISTING SYSTEM:

- Collecting large amount of data set.
- Large number of training data and annotations are needed which may not be practical in some problems.

3.4. PROPOSED SYSTEM:

Here we are building a model by applying various machine learning algorithms find the best accurate model. Thus a person will get to know whether he/she is having a liver disease or not. Some of the machines learning algorithms are:

1. Linear Regression:

Linear Regression is a supervised machine learning algorithm where the predicted output is continuous and has a constant slope. It's used to predict values within a continuous range, (e.g. sales, price) rather than trying to classify them into categories (e.g. cat, dog).

Steps to implement Linear regression model:

- 1. Initialize the parameters.
- 2. Predict the value of a dependent variable by given an independent variable.
- 3. Calculate the error in prediction for all data points.
- 4. Calculate partial derivative w.r.t a0 and a1.
- 5. Calculate the cost for each number and add them.

2. Multiple Linear Regression:

Multiple Linear Regression is one of the important regression algorithms which models the linear relationship between a single dependent continuous variable and more than one independent variable. Multiple regression is a broader class of regressions that encompasses linear and nonlinear regressions with multiple explanatory variables.

3.Random Forest:

A random forest is a machine learning technique that's used to solve regression and classification problems. It utilizes ensemble learning, which is a technique that combines many classifiers to provide solutions to complex problems.

Working of Random Forest Algorithm

- Step 1 First, start with the selection of random samples from a given dataset.
- Step 2 Next, this algorithm will construct a decision tree for every sample. Then it will get the prediction result...
- Step 3 In this step, voting will be performed for every predicted result.
- Step 4 At last, select the most voted prediction result as the final prediction result.

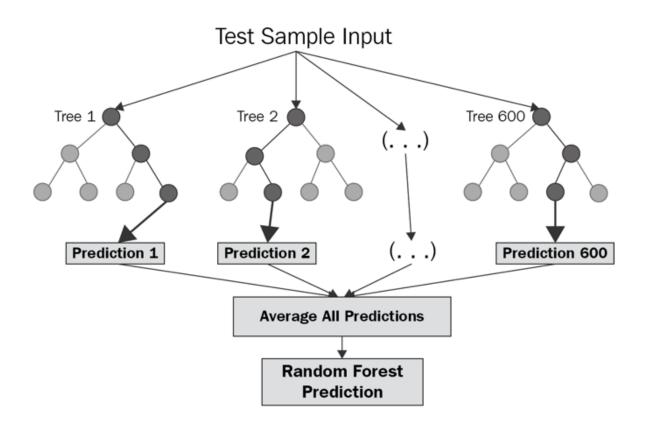


Figure 1 : Random Forest

4. <u>Logistic regression:</u>

- Logistic regression is a supervised learning classification algorithm used to predict the probability of a target variable. The nature of target or dependent variable is dichotomous, which means there would be only two possible classes.... Mathematically, a logistic regression model predicts P(Y=1) as a function of X.
- Logistic Regression is used when the dependent variable (target) is categorical. For example,
- To predict whether an email is a spam (1) or (0)
- Whether the tumor is malignant (1) or not (0)

You will need to train the datasets to run smoothly and see an incremental improvement in the prediction rate.

5. k-nearest neighbor algorithm:

- It is a supervised machine learning algorithm. The algorithm can be used to solve both classification and regression problem statements. The number of nearest neighbor's to a new unknown variable that has to be predicted or classified is denoted by the symbol 'K'.
- KNN algorithm at the training phase just stores the dataset and when it gets new data, then it classifies that data into a category that is much similar to the new data.
- K-Nearest Neighbors (KNN) is one of the simplest algorithms used in Machine
 Learning for regression and classification problem. KNN algorithms use data and
 classify new data points based on similarity measures (e.g. distance function). The
 data is assigned to the class which has the nearest neighbors.
- It's also worth noting that the KNN algorithm is also part of a family of "lazy learning" models, meaning that it only stores a training dataset versus undergoing a training stage. This also means that all the computation occurs when a classification or prediction is being made. Since it heavily relies on memory to store all its training data, it is also referred to as an instance-based or memory-based learning method.

• The K-NN working can be explained on the basis of the below algorithm:

Step-1: Select the number K of the neighbors

Step-2: Calculate the Euclidean distance of K number of neighbors

Step-3: Take the K nearest neighbors as per the calculated Euclidean distance.

Step-4: Among these k neighbors, count the number of the data points in each category.

Step-5: Assign the new data points to that category for which the number of the neighbor is maximum.

Step-6: Our model is ready.

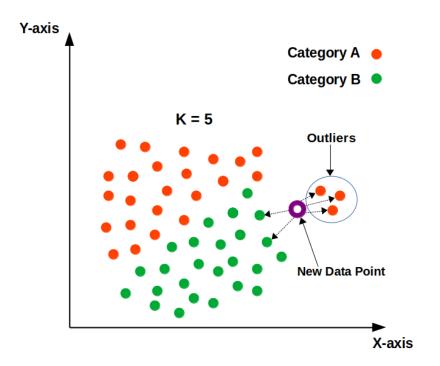


Figure 2: k-nearest neighbor

4. EXPERIMENTAL ANALYSIS

Milestone 1: Data Collection

ML depends heavily on data, without data, a machine can't learn. It is the most crucial aspect that makes algorithm training possible. In Machine Learning projects, we need a training data set. It is the actual data set used to train the model for performing various actions.

You can collect datasets from different open sources like kaggle.com, data.gov; UCI machine learning repository etc. The dataset used for this project was obtained from Kaggle.

Milestone 2: Data Pre-processing

Data Pre-processing includes the following main tasks

- Importing the libraries.
- Importing the dataset.
- Analyse the data.
- Taking care of Missing Data.
- Data Visualisation.
- Splitting Data into Train and Test.

Milestone 3: Model Building

The model building process involves setting up ways of collecting data, understanding and paying attention to what is important in the data to answer the questions you are asking, finding a statistical, mathematical or a simulation model to gain understanding and make predictions. Model Building Includes:

- Import the model building libraries.
- Initialising the model.
- Training the model.
- Model Evaluation.
- Save the Model.

Milestone 4: Application Building

- Create an HTML File.
- Build python code.
- Run the app in local browser.
- Show casting the prediction on UI.

4.1. PROJECT ARCHITECTURE:

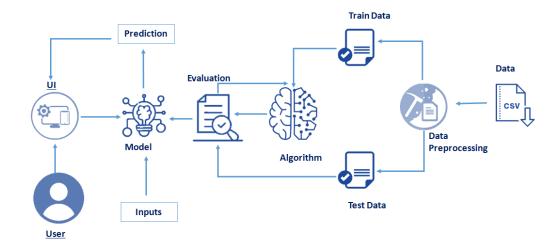


Figure 3: Project Architecture

4.2. SOFTWARE AND HARDWARE REQUIREMENTS:

Software Requirements:

- > Anaconda Environment
- > Flask
- > Python 3.9
- ➤ And other python libraries like NumPy, pandas.

Hardware Requirements:

- > Operating system
- Processing
- > RAM
- > Operating system specifications
- Disk space

4.3. BLOCK DIAGRAM:

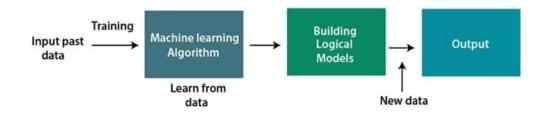


Figure 4: Block Diagram

4.4. PROJECTFLOW:

- User interacts with the UI (User Interface) to upload the input features.
- Uploaded features/input is analysed by the model which is integrated.
- Once a model analyses the uploaded inputs, the prediction is showcased on the UI.

5. DESIGN

5.1. USE CASEDIAGRAM:

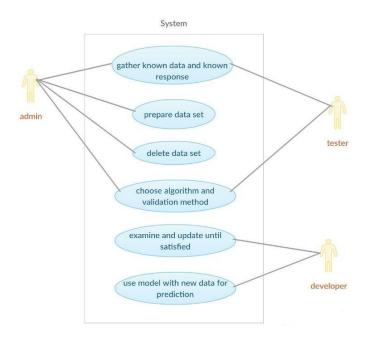


Figure 5: Use case Diagram

5.2. FLOWCHART:

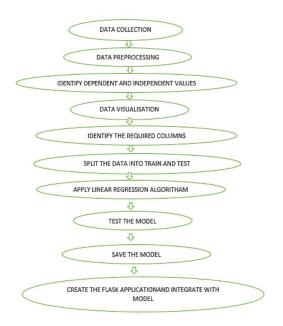


Figure 6: Flowchart

5.3. DECISION TREE:

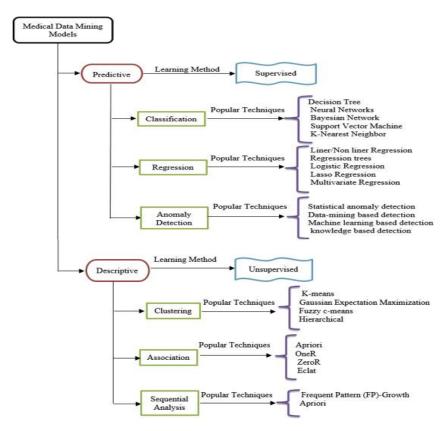


Figure 7: Decision Tree

5.3. SEQUENCE DIAGRAM:

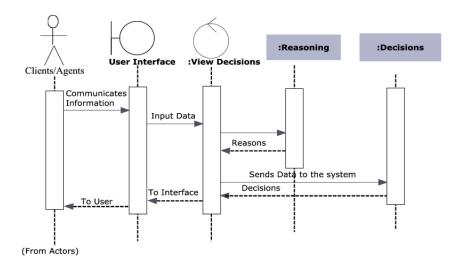


Figure 8: Sequence Diagram

5.4. COMMUNICATIO DIAGRAM:

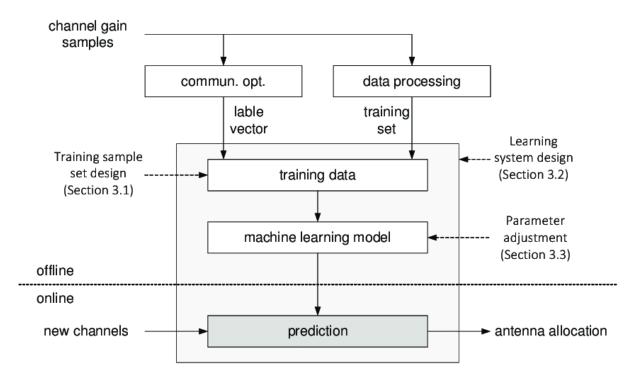


Figure 9: Communication Diagram

6. CONCLUSION

In UG Project Phase-1, we have worked on problem statement, literature survey and also done the experimental analyses which are required for the project to move forward. In experimental analysis we have discussed about the machine learning concepts and models and explained the algorithms to be used in the project. We also discussed about the flowcharts, use case diagrams, decision tree and sequence diagrams which are used in the project. Based on the experimental analysis we have designed the model for the project. Entire designing part is involved in UG Project Phase-1.

7. FUTURE SCOPE

UG Project Phase-2 is the extension of UG Project Phase-1. UG Project Phase-2 involves all the coding and implementation of the design which we have retrieved from UG Project Phase-1. All the implementation is done and conclusions will be retrieved in the phase. We will also work on the applications, advantages, and disadvantages of the project in this phase. Future scope of the project will be also discussed in the UG Project Phase-2.

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CERTIFICATE OF COMPLETION UG PROJECT PHASE-2

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1. INTRODUCTION

The liver is responsible for many critical functions within the body and should it become diseased or injured, the loss of those functions can cause significant damage to the body. Discovering the existence of liver disease at an early stage is a complex task for the doctors. The main objective of this project is to 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.

Machine learning (ML) is a type of artificial intelligence (AI) that allows software applications to become more accurate at predicting outcomes without being explicitly programmed to do so. Machine learning algorithms use historical data as input to predict new output values. Here we are building a model by applying various machine learning algorithms find the best accurate model. Thus a person will get to know whether he/she is having a liver disease or not. On our Dataset, we have applied Random Forest Regression and KNN algorithm. Random forest is a Supervised Machine Learning Algorithm that is used widely in Classification and Regression problems. It builds decision trees on different samples and takes their majority vote for classification and average in case of regression. The abbreviation KNN stands for "K-Nearest Neighbour". It is a supervised machine learning algorithm. The algorithm can be used to solve both classification and regression problem statements. The number of nearest neighbours to a new unknown variable that has to be predicted or classified is denoted by the symbol 'K'.

UG Project Phase-2 involves all the coding and implementation of the design which we have retrieved from UG Project Phase-1. All the implementation is done and conclusions are retrieved in this phase. We will also work on the applications, advantages, and disadvantages of the project in this phase. Future scope of the project will be also discussed in the UG Project Phase-2.

2. CODE SNIPPETS

2.1.MODELCODE:

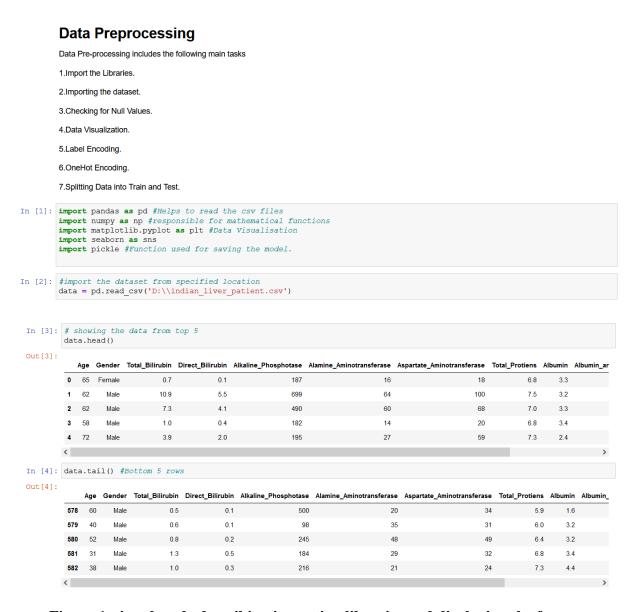


Figure 1: .ipynb code describing importing libraries and displaying the few rows from the Dataset.

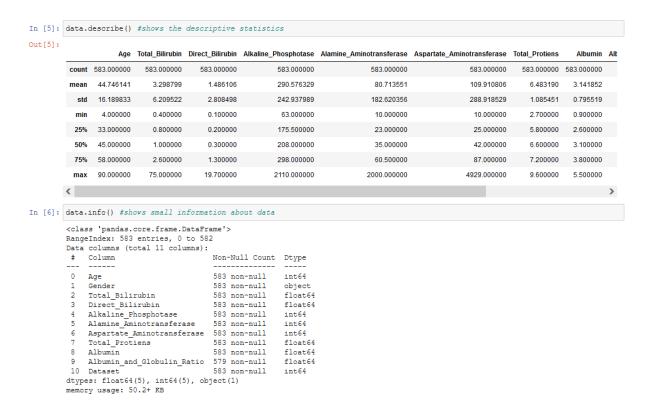


Figure 2: .ipynb code describing describe() and info() methods.

```
In [7]: data.isnull().any()
Out[7]: Age
                                             False
         Total_Bilirubin
                                              False
          Direct_Bilirubin
         Alkaline_Phosphotase
Alamine_Aminotransferase
                                              False
         Aspartate Aminotransferase
                                              False
          Total_Protiens
                                              False
         Albumin
                                             False
          Albumin_and_Globulin_Ratio
         Dataset
                                             False
         We can see that there are null values in the Albumin_and_Globulin_Ration Column.
In [8]: data.isnull().sum()
Out[8]: Age
         Gender
Total Bilirubin
         \overline{\text{Direct\_Bilirubin}}
         Alkaline_Phosphotase
Alamine_Aminotransferase
Aspartate_Aminotransferase
          Total_Protiens
         Albumin
          Albumin_and_Globulin_Ratio
         Dataset
         dtype: int64
```

Figure 3: .ipynb code describing whether they are any NULL values in the Dataset.

```
In [10]: data['Dataset'].unique()
Out[10]: array([1, 2], dtype=int64)
In [11]: data['Albumin_and_Globulin_Ratio'].fillna(data['Albumin_and_Globulin_Ratio'].mode()[0], inplace=True)
In [12]: #checking for the missing data after cleaning data
          data['Albumin_and_Globulin_Ratio'].fillna(data['Albumin_and_Globulin_Ratio'].mode()[0], inplace=True) data.isnull().sum()
Out[12]: Age
          Gender
          Total_Bilirubin
          Direct Bilirubin
                                           0
          Alkaline Phosphotase
          Alamine_Aminotransferase
Aspartate_Aminotransferase
          Total_Protiens
          Albumin
          Albumin_and_Globulin_Ratio
          Dataset
          dtype: int64
```

We can see that all the null values are filled.

Figure 4: .ipynb code describing filling of Null Values.

```
In [13]: plt.figure(figsize=(15,10))
          plt.subplot(3,3,1)
          plt.scatter(data['Age'], data['Dataset'])
         plt.ylabel('Dataset')
          plt.xlabel('Age')
          plt.subplot(3,3,2)
          plt.scatter(data['Gender'], data['Dataset'],)
         plt.ylabel('Dataset')
plt.xlabel('Gender')
          plt.subplot(3,3,3)
          plt.scatter(data['Total_Bilirubin'], data['Dataset'],)
          plt.ylabel('Dataset')
          plt.xlabel('Total_Bilirubin')
          plt.subplot(3,3,4)
          plt.scatter(data['Direct_Bilirubin'], data['Dataset'],)
         plt.ylabel('Dataset')
plt.xlabel('Direct_Bilirubin')
          plt.subplot(3,3,5)
          plt.scatter(data['Alkaline_Phosphotase'], data['Dataset'],)
          plt.ylabel('Dataset')
plt.xlabel('Alkaline_Phosphotase')
          plt.subplot(3,3,6)
          plt.scatter(data['Alamine Aminotransferase'], data['Dataset'],)
          plt.ylabel('Dataset')
          plt.xlabel('Alamine_Aminotransferase')
          plt.subplot(3,3,7)
          plt.scatter(data['Aspartate_Aminotransferase'], data['Dataset'],)
          plt.ylabel('Dataset')
          plt.xlabel('Aspartate Aminotransferase')
          plt.subplot(3,3,8)
          plt.scatter(data['Total_Protiens'], data['Dataset'],)
          plt.ylabel('Dataset')
plt.xlabel('Total_Protiens')
```

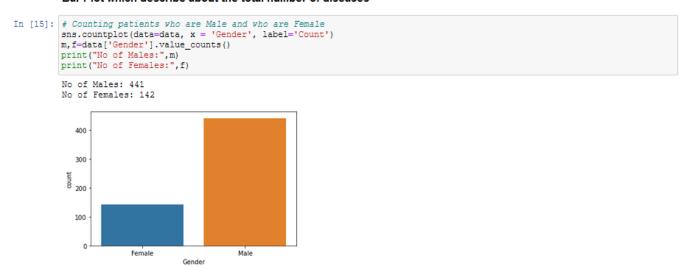
```
plt.subplot(3,3,9)
             plt.scatter(data['Albumin_and_Globulin_Ratio'], data['Dataset'])
             plt.ylabel('Dataset')
plt.xlabel('Albumin_and_Globulin_Ratio')
Out[13]: Text(0.5, 0, 'Albumin_and_Globulin_Ratio')
                 2.0
                                                                         2.0
                                                                                                                                2.0
                 1.8
                                                                         1.8
                                                                                                                                1.8
               Dataset
14
                                                                      Dataset
1.4
                                                                                                                             1.6
1.4
                 1.2
                                                                         1.2
                                                                                                                                1.2
                                                                                                                       Male
                                                                                                                                1.0
                 1.0
                                                                         1.0
                                         40
                                                  60
                                                            80
                                                                                                                                                 20
                                                                                                                                                            40
                                                                                                                                                                       60
                                                                                                                                                     Total Bilirubin
                  2.0
                                                                         2.0
                                                                                                                                2.0
                 1.8
                                                                         1.8
                                                                                                                                1.8
               Dataset
14
                                                                      Dataset
14
                                                                                                                             1.4
1.6
                 1.2
                                                                         1.2
                                                                                                                                1.2
                                                                         1.0
                                            10
                                                                                                1000
                                                                                                          1500
                                                                                                                     2000
                                                                                                                                                500
                                                                                                                                                         1000
                                                                                                                                                                    1500
                                                                                                                                                                              2000
                                      Direct Bilirubin
                                                                                          Alkaline Phosphotase
                                                                                                                                               Alamine Aminotransferase
                  2.0
                                                                         2.0
                                                                                                                                2.0
                 1.8
                                                                         1.8
                                                                                                                                1.8
               Dataset
14
                                                                      14 Dataset
14
                                                                                                                             Dataset
14
                 1.2
                                                                         1.2
                                                                                                                                1.2
                 1.0
                                                                         1.0
                                                                                             5 6 7
Total_Protiens
                                                                                                                                              1.0 1.5 2.0
Albumin_and_Globulin_Ratio
                                .000 2000 3000 400
Aspartate_Aminotransferase
                                                                                                                                         0.5
                              1000
                                                        4000
                                                                5000
```

Figure 5: .ipynb code describing Scatter Plot.

```
In [14]: 
# Counting patients who are diagnosed and not diagnoised with liver disease
sns.countplot(data=data, x = 'Dataset')
LD, NLD=data('Dataset').value_counts()
print("Non-liver disease patinets:", NLD)

liver disease patinets: 416
Non-liver disease patinets: 167
```

Bar Plot which describe about the total number of diseases



Bar plot between Gender and Count

Figure 6: .ipynb code describing Bar Plot.

```
In [16]: # Importing the LabelEncoder libraray from scikit-learn
           #encoding technique to convert categorical data to numeric data from sklearn.preprocessing import LabelEncoder
           le=LabelEncoder()
           # Converting Textual data into numeric data
data['Gender'] = le.fit_transform(data['Gender'])
           data.head()
Out[16]:
               Age Gender Total_Bilirubin Direct_Bilirubin Alkaline_Phosphotase Alamine_Aminotransferase Aspartate_Aminotransferase Total_Protiens Albumin Albu
                                            0.1
            0 65
                    0 0.7
                                                                         187
                                                                                                   16
                                                                                                                                          6.8
                                                                                                                                                  3.3
                                                                                                                                          7.5
                                                                                                                                                   3.2
            2
                62
                                    7.3
                                                    4.1
                                                                         490
                                                                                                   60
                                                                                                                            68
                                                                                                                                          7.0
                                                                                                                                                  3.3
                58
                                     1.0
                                                    0.4
                                                                         182
                                                                                                   14
                                                                                                                             20
                                                                                                                                          6.8
                                                                                                                                                  3.4
                      1 3.9
                                                    2.0
                                                                                                   27
                                                                                                                                          7.3
                                                                                                                                                  2.4
In [17]: # dividing the data into input and output
           x=data.iloc[:,0:-1]
           y=data.iloc[:,-1]
```

In the above code we are creating array or list of the independent variable x with our selected columns and for dependent variable y we are only taking the dependent or output or target column.

Figure 7: .ipynb code describing label encoding and splitting dataset into independent and dependent variables

Model Building

-Predictive modeling is a mathematical approach to create a statistical model to forecast future behavior based on input test data.

Model building includes the following main tasks

Training and testing the model, Evaluation of Model, Save the model, Predicting the output using the model.

Figure 8: .ipynb code describing model building.

```
In [27]: # Checking for accuracy score from actual data and predicted data
         RFaccuracy_score(RFpred, ytest)
         RFaccuracy
Out [27]: 0.7008547008547008
In [28]: # showing the confusion matrix
         RFcm=confusion_matrix(RFpred, ytest)
         RFcm
Out[28]: array([[72, 24],
                [11, 10]], dtype=int64)
In [29]: # K-Nearest Neighbors Model
         from sklearn.neighbors import KNeighborsClassifier
         KNN = KNeighborsClassifier()
In [30]: # train the data with K-Nearest Neighbors Model
         KNN.fit(xtrain, ytrain)
Out[30]: KNeighborsClassifier()
In [31]: KNNpred=KNN.predict(xtest)
In [32]: # Checking for accuracy score from actual data and predicted data
         KNNaccuracy=accuracy_score(KNNpred, ytest)
         KNNaccuracy
Out [32]: 0.6581196581196581
In [33]: # showing the confusion matrix
         KNNcm=confusion_matrix(KNNpred, ytest)
         KNNcm
Out[33]: array([[67, 24],
                [16, 10]], dtype=int64)
In [34]: print("Random Forest Algorithm accuracy score : {value:.2f} %".format(value=RFaccuracy*100))
         print("K-Nearest Neighbors Algorithm accuracy score : {value:.2f} %".format(value=KNNaccuracy*100))
         Random Forest Algorithm accuracy score : 70.09 %
         K-Nearest Neighbors Algorithm accuracy score : 65.81 %
In [35]: # saving the model import pickle
         pickle.dump(RFmodel, open('liver_analysis_RF.pkl','wb'))
```

Figure 9: .ipynb code describing finding Accuracy.

2.2.HTML CODE AND PYTHON CODE

1. app.py code:

```
File Edit Search Source Run Debug Connoise Projects Tools View Help

The Edit Search Source Run Debug Connoise Projects Tools View Help

The Edit Search Source Run Debug Connoise Projects Tools View Help

Those Flask Search Tasks, Prender, Seeplate, Prequest # Lask %s. a population

# under to run/serve on repplication

# under to run/serve on repplication

# under to run/serve on repplication

# upport pickle # pickle is used for seeplate; Prequest # Lask %s. a population

# under to run/serve on repplication

# upport pickle # pickle is used for seeplate; The search was predicted to the search of the used for seeplate; The search was predicted to the search of the used for seeplate; The search was predicted to the search of the unit seeplate of search;

# apport pickle # pickle is used for serializing and de-serializing Python object structures

## apport pickle # pickle is used for serializing and de-serializing python object structures

## apport pickle # pickle is used for serializing and de-serializing python object structures

## apport pickle # pickle is used for serializing and de-serializing python object structures

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## apport pickle # pickle is used for serializing python object structures

## apport pickle # pickle is used for serializing python object structures

## apport pickle # pickle is used for serializing python object structures

## apport pickle # pickle is used for serializing python object structures

## apport pickle # pickle is used for serializing python object structures

## apport pickle # pickle is used for serializing python object structures

## apport pickle # pickle is used for serializing python object s
```

Figure 10: .python code used for rendering all the HTML pages.

2. home.html:

```
File Edit Search Source Run Debug Connoies Projects Tools View Help

Cultivarily Decision Floats immobilish to produce the project Search Source Run Debug Connoies Project Souls View Help

Supply X | Doma Brill X | Production Floats | Doma Brill X | Doma Brill
```

Figure 11: home.html page is the code for home page of our Web Application.

3.index.html:

```
Spyder (Python 3.9)
  File Edit Search Source Run Debug Consoles Projects Tools View Help
    C:\Users\hp\Desktop\Flask\templates\index.html
  app.py X home.html X index.html X chance.html X nochance.html X
           1 klDCTYPE html>
4 chtml>
3 chead>
4 ctitle-liver Patient Analysis</title>
5 ct-- Latest compiled and minified CSS -->
6 clink rel="stylesheet" href="https://maxcdn.bootstrapcdn.com/bootstrap/3.3.7/css/bootstrap.min.css">
6 clink rel="stylesheet" href="https://maxcdn.bootstrapcdn.com/bootstrap/3.3.7/css/bootstrap.min.css">
7 clink rel="stylesheet" href="https://maxcdn.bootstrap.dn.com/bootstrap/3.3.7/css/bootstrap.min.css">
7 clink rel="stylesheet" href="https://maxcdn.bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/bootstrap.dn.com/
                             cstyle type="text/css">
body{
background-color:#FFFEF9;
background-simage:url('../static/img/index.jpg');
background-position: center;
background-attachment: fixed;
background-size:fill;
background-size: 110% 100%;
background-repeat: no-repeat;
background-position: center bottom;
}
                                    background;

} page-header(
background-color: coral;
width: 100%;
height: auto;
text-slign: center;
padding-top: 5px;
color: #fff;
}
                                     }
h1{
font-size: 40px;
font-weight: bold;
```

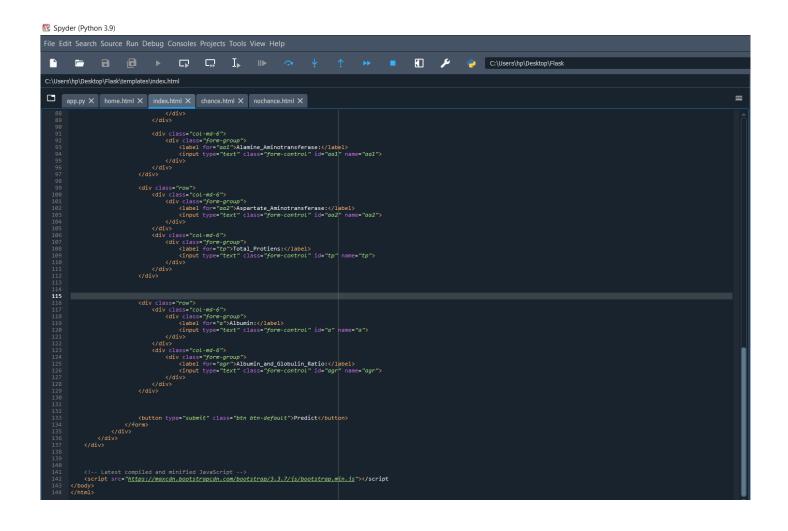


Figure 12: index.html is the page which displays all the inputs that are needed to be given by the user.

4.chance.html:

```
Spyder (Python 3.9)
 File Edit Search Source Run Debug Consoles Projects Tools View Help
  🖺 🗁 🗟 🖻 🕨 🗔 🎧 🔝 🔝 🔝 🕨 💀 🖢 🖸 C:\Users\hp\Desktop\Flask
 C:\Users\hp\Desktop\Flask\templates\chance.html
 app.py X chance.html X home.html X index.html X nochance.html X
     1 k!DOCTYPE html>
2 <html>
3 <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">
              cstyle type="text/css">
body{
   background-color:#FFFEF9;
   background-image:url('../static/img/chance.jpg');
   background-sposition: center;
   background-attachment: fixed;
   background-size:Fill;
   background-size: 70% 70%;
   background-repeat: no-repeat;
   background-position: center bottom;
}
                     }
.page-header{
                        page-meader(
background-color: coral;
width: 100%;
height: auto;
text-align: center;
padding-top: 5px;
color: #fff;
                    }
h1{
font-size: 40px;
font-weight: bold;
                           color: black;
font-size: 20px;
font-weight: bold;
font-position:center;
                </div>
                <!-- Latest compiled and minified JavaScript -->
<script src="https://maxcdn.bootstrapcdn.com/bootstrap/3.3.7/js/bootstrap.min.js"></script</pre>
```

Figure 13: chance.html is the page which displays that the user is having liver disease.

5.nochance.html:

```
Spyder (Python 3.9)
 File Edit Search Source Run Debug Consoles Projects Tools View Help
        C:\Users\hp\Desktop\Flask\templates\nochance.html
 app.py X home.html X index.html X chance.html X nochance.html X
    1 <IDCTYPE html>
2 <html>
3 <head>
4 <title>liver Patient Analysis</title>
5 <{!-- Latest compiled and minified CSS -->
4 <link rel="stylesheet" href="https://maxcdn.bootstrapcdn.com/bootstrap/3.1.Z/css/bootstrap.min.css">
7 
               <style type="text/css">
body{
   background-color:#FFFFF9;
   background-image:url('../static/img/Nochance.jpg');
   background-position: center;
   background-sitze:fill;
   background-size:Fill;
   background-size:60% 70%;
   background-repeat: no-repeat;
   background-position: center bottom;
}
                   backgrown
}
page-header{
background-color: coral;
width: 100%;
height: auto;
text-align: center;
padding-top: 5px;
color: #fff;
}
                     }
h1{
font-size: 40px;
font-weight: bold;
                   }
h3{
color: black;
font-size: 20px;
font-weight: bold;
font-position:center;
              <!-- Latest compiled and minified JavaScript -->
<script src="https://maxcdn.bootstrapcdn.com/bootstrap/3.3.7/js/bootstrap.min.js"></script</pre>
          </body>
```

Figure 14: nochance.html is the page which displays that the user is not having liver disease.

3.CONCLUSION

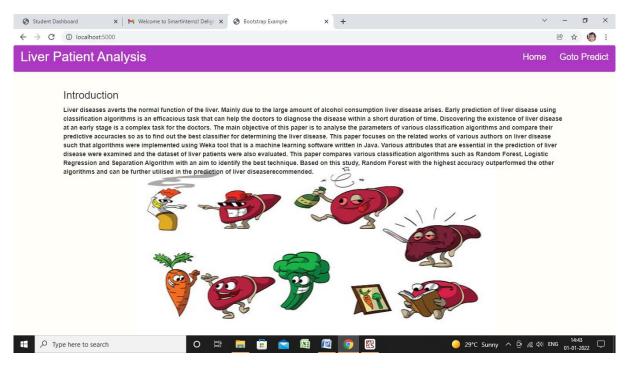


Figure:15: Home page (Which gives introduction to Liver Patient Analysis)

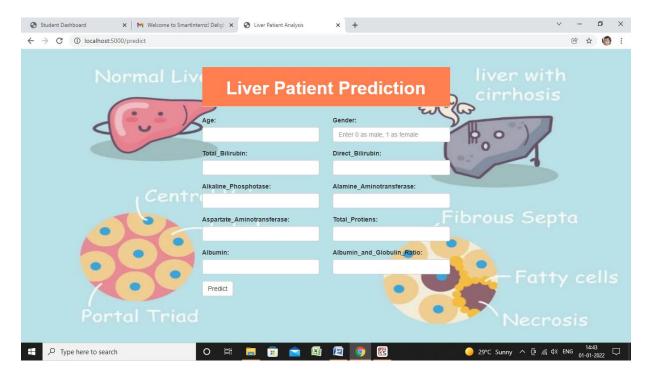


Figure: 16: Input pages (Which takes inputs from User)

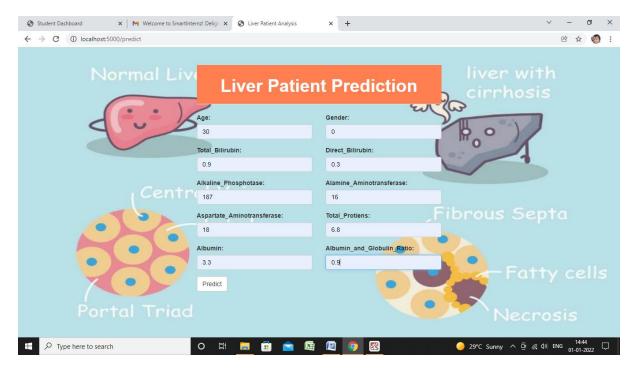


Figure: 17: Input pages (Which takes inputs from User)

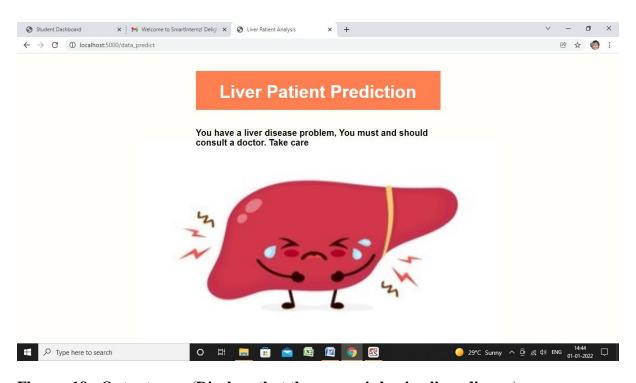


Figure: 18: Output page (Displays that the person is having liver disease)

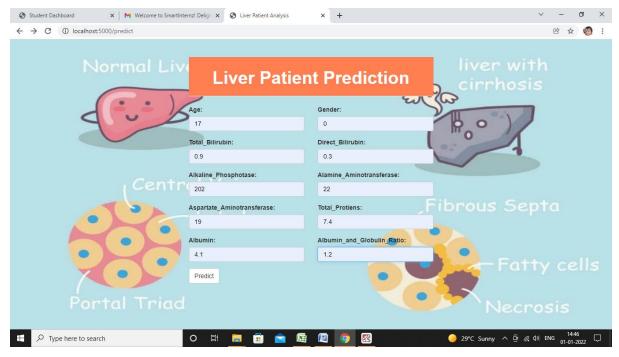


Figure: 19: Input pages (Which takes inputs from User)

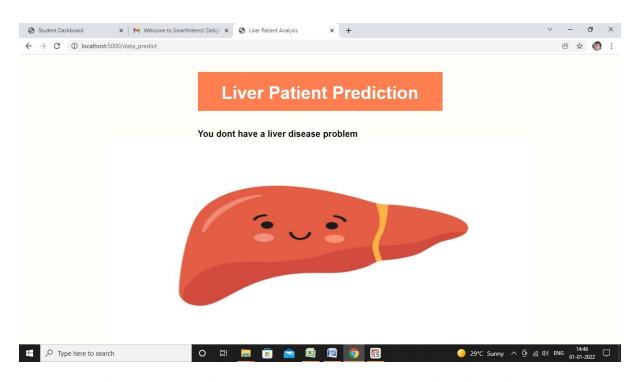


Figure: 20: Output page (Displays that the person is not having liver disease)

4. APPLICATION

The areas where this solution can be applied:

- Can be applied in each and every individual's Daily Life.
- Can be used in medical fields for faster prediction in determining the disease.

5. ADVANTAGES

Those with liver disease may qualify for **Social Security disability**. If you suffer from autoimmune hepatitis, cirrhosis, and other chronic liver conditions, you may qualify for Social Security disability benefits if the condition meets the Social Security blue book's listing.

6. DISADVANTAGES

Liver disease can progress to cirrhosis and liver failure. Associated complications may include **increased risk of bleeding and infection, malnutrition and weight loss**, and decreased cognitive function. Some liver diseases are associated with an increased risk of developing liver cancer.

Liver failure, or liver failure that occurs over many years, may cause:

- Fatigue
- Nausea
- Loss of appetite
- Diarrhoea
- Vomiting blood
- Blood in the stool

As liver failure advances, symptoms become more severe. In later stages, symptoms of liver failure may include:

- Jaundice(yellowing of the skin and eyes)
- Extreme tiredness
- Disorientation (confusion and uncertainty)
- Fluid build-up in the abdomen and extremities (arms and legs)

Sometimes, the liver fails suddenly, which is known as acute liver failure. People with acute liver failure may have the following symptoms:

- Bleeding
- Changes in mental status
- Musty or sweet breath odour
- Movement problems
- Loss of appetite
- General feeling of being unwell

7. FUTURE SCOPE

On our Dataset, we have applied Random Forest Regression and KNN algorithm. Random forest has got the highest accuracy of 70%.

Enhancements that can be made in the future:

This model can be further developed to suggest what are the preventions that a person diagnosed with liver disease should be follow and Can also enhance by adding the strict diet schedules which should be followed to be strong and fit. And we can further classify type of disease also based on the inputs given by the user.

8. BIBILOGRAPHY

- [1] Parminder Kaur1and Aditya Khamparia, —Classification Of Liver Based Diseases Using Random Tree", International Journal of Advances in Engineering & Technology, June, 2015, ISSN: 22311963, Vol. 8, Issue 3, pp. 306-313
- [2] Lichman, M. (2013). UCI Machine Learning Repository [http://archive.ics.uci.edu/ml]. Irvine, CA: University of California, School of Information and Computer Science.
- [3] Bendi Venkata Ramana, Prof. M.Surendra Prasad Babu,, Prof. N. B. Venkateswarlu, A Critical Study of Selected Classification Algorithms for Liver Disease Diagnosis, International Journal of Database Management Systems (IJDMS), Vol.3, No.2, May 2011.
- [4] Chuan Choong Yang, Chit Siang Soh and Vooi Voon Yap, —A nonintrusive appliance load monitoring for efficient energyconsumption based on Naive Bayes classifier, Sustainable Computing: Informatics and Systems 14 (2017) 34–42.
- [5] Cleary, J. and L. Trigg, —K*: An Instance-based Learner Using an Entropic Distance Measure, in 12th International Conference on Machine Learning. 1995. p. 108-114.
- [6] Ross J. Quinlan: —Learning with Continuous Classes In Proceedings AI'92 (Adams & Sterling, Eds), 343-348, Singapore: World Scientic, 1992.
- [7] Youvrajsinh Chauhan and Jignesh Vania, —J48 Classifier Approach to Detect Characteristic of Bt Cotton base on Soil Micro Nutrientl, International Journal of Computer Trends and Technology (IJCTT) volume 5 number 6 –Nov 2013.
- [8] S. Muthuselvan and Dr. K. Soma Sundaram, —An Analysis of Knowledge Discovery Process Over a Cloud Environment A Surveyl, International Journal of Applied Engineering Research ISSN 0973-4562 Volume 10, Number 17 (2015).
- [9] Inderjit Kaur, Deep Mann, Data Mining in Cloud Computing, International Journal of Advanced Research in Computer Science and Software Engineering.
- [10] Emmanuel Ahishakiye, Elisha Opiyo Omulo, Danison Taremwa and Ivan Niyonzima, —Crime Prediction Using Decision Tree (J48) Classification Algorithm International Journal of Computer and Information Technology (ISSN: 2279 0764) Volume 06 Issue 03, May 2017.

9.HELP FILE

PROJECT EXECUTION:

STEP-1: Go to Start, search and launch ANACONDA NAVIGATOR.

STEP-2: After launching of ANACONDA NAVIGATOR, launch JUPYTER NOTEBOOK.

STEP-3: Open "Major project code" IPYNB file.

STEP-4: Then run all the cells.

STEP-5: All the data preprocessing, training and testing, model building, accuracy of the model can be showcased.

STEP-6: And a pickle file will be generated.

STEP-7: Create a Folder named **FLASK** on the **DESKTOP.** Extract the pickle file into this Flask Folder.

STEP-8: Extract all the html files (home.html, index.html, chance.html, nochance.html) and python file(app.py) into the **FLASK Folder.**

STEP-9: Then go back to ANACONDA NAVIGATOR and the launch the SPYDER.

STEP-10: After launching Spyder, give the path of **FLASK FOLDER** which you have created on the DESKTOP.

STEP-11: Open all the app.py and html files present in the Flask Folder.

STEP-12: After running of the app.py, open **ANACONDA PROMPT** and follow the below steps:

cd File Path → click enter

python app.py -> click enter (We could see running of files).

STEP-13: Then open BROWSER, at the URL area type "localhost:5000".

STEP-14: Home page of the project will be displayed.

STEP-15: Click on "Go to Predict". Directly it will be navigated to index page.

STEP-16:A index page will be displayed where the user needs to give the inputs and then click on "**Predict**". Output will be generated whether a person is having liver disease or not.