



Prediction And Analysis Of Liver Patient Data Using Machine Learning

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1.INTRODUCTION

1.1 Project overview

Liver is the largest organ in the abdomen. This is the primary organ for maintaining the chemicals like glucose, balancing so many nutrients, fat, vitamins, cholesterol and hormones. Liver disease prevents normal liver function. Mainly due to the large amount of alcohol consumption liver disease develops. Early detection of liver disease using classification algorithms is an effective activity that can help doctors diagnose the disease in a short period of time. Early detection of liver disease is a daunting task for doctors. The main purpose of our project is to analyse the parameters of the various classification algorithms and compare their predictive accuracy to determine the best stage for determining liver disease.

1.2 Objectives

The primary objective is to develop a machine learning model that accurately predicts the likelihood of a patient developing a liver disease based on their medical history, symptoms, and other relevant data, enabling early intervention and improved health outcomes

2. Project Initialization and Planning Phase

2.1 Define Problem Statement

The early prediction of liver disease can significantly improve patient outcomes, reduce healthcare costs, and enhance the quality of life. With the advent of big data, machine learning, and advanced analytics, there is a growing potential to predict diseases before they manifest clinically. This predictive capability can aid in proactive treatment, preventive measures, and personalized healthcare. In the realm of healthcare, accurately predicting the onset and progression of disease is crucial for early intervention and improved patient outcomes. Despite advancements in medical technology and data analytics, current methods often lack precision and timeliness, leading to delayed diagnoses and suboptimal treatment plans. This project aims to develop a robust and scalable liver disease prediction model that leverages patient data, including medical history, genetic information, lifestyle factors, and real-time health indicators, to forecast the likelihood of disease occurrence with high accuracy. The goal is to enhance predictive capabilities, reduce healthcare costs, and ultimately improve patient care through timely and personalized interventions.

2.2 Project Proposal (Proposed solution)

This project proposal outlines a solution to address the problem of early liver disease detection through machine learning. With a clear objective to develop a predictive model for assessing disease risk based on symptoms, lifestyle factors, and health data, the proposal defines the scope of the project, including data collection, model development, and deployment. The proposed solution details the approach to be used, key features of the model, and specifies the resource requirements including hardware, software, and personnel. By creating an accurate and user friendly tool, the project aims to enable proactive health management and improve early liver disease detection.

2.3 Initial Project Planning

Develop a machine learning model to predict the likelihood of a liver disease based on patient data. The scope includes data collection, basic preprocessing, model development, and initial evaluation for accuracy. The model will be deployed in a simple application for healthcare use. Ongoing maintenance and updates are not included.

3. Data Collection and Preprocessing Phase

3.1 Data Collection Plan and Raw Data Sources Identified

- **Data Availability**: Determine which sources provide the required data and whether they are accessible.
- **Data Quality**: Evaluate the reliability, completeness, and accuracy of the data from each source.
- Data Consistency: Ensure consistency across different sources and data formats. Time: time in format hh:mm:ss
- **Data Extraction**: Design protocols for extracting data from electronic health records (EHRs), databases, or files.

- **Data Integration**: Merge data from multiple sources into a unified dataset suitable for analysis.
- **Data Cleaning**: Preprocess data to handle missing values, outliers, and inconsistencies.
- **Hospital EHRs**: Patient demographics, medical history, medications, and laboratory results.
- **Public Datasets**: Liver disease datasets from repositories like UCI Machine Learning Repository.
- Research Databases: Specific datasets from liver disease research studies.
- **Health Insurance Claims**: Historical claims data with diagnostic codes related to liver diseases.

• 3.2 Data Quality Report

• Data quality is ensured through thorough verification, addressing missing values, and maintaining adherence to ethical guidelines, establishing a reliable foundation for predictive modeling.

3.3 Data Exploration and preprocessing

- Data Exploration involves analyzing the liver disease patient dataset to understand patterns, distributions, and outliers.
- Preprocessing includes handling missing values, scaling, and encoding categorical variables.
- These crucial steps enhance data quality, ensuring the reliability and effectiveness of subsequent analysis.

4. Model Development Phase

4.1 Feature Selection Report

- The Feature Selection Report outlines the rationale behind choosing specific features (e.g., age, food plan, gender) for the liver disease prediction model.
- It evaluates relevance, importance, and impact on predictive accuracy, ensuring the inclusion of key factors influencing the model's ability.

4.2 Model Selection Report

 The Model Selection Report details the rationale behind choosing Linear Regression, Random Forest, Decision Tree, and XGB models for liver disease prediction. • It considers each model's strengths in handling complex relationships, interpretability, adaptability, and overall predictive performance, ensuring an informed choice aligned with project objectives.

4.3 Initial Model Training Code, Model Validation and Evaluation Report

- The Initial Model Training Code employs selected algorithms on the liver disease dataset, setting the foundation for predictive modeling.
- The subsequent Model Validation and Evaluation Report rigorously
 assesses model performance, employing metrics like Mean Absolute Error
 (MAE), Mean Squared Error (MSE), Root Mean Squared Error (RMSE),
 R-squared error to ensure reliability and effectiveness in predicting liver
 disease.

5.Model Optimization and Tuning Phase

Final Model Selection Justification

- The XGBoost Regressor is the final model chosen because of its best overall performance compared to the other models.
- It captures the variance in the data very well with minimal prediction error. XGBoost can capture complex non-linear relationships.

6. RESULTS

6.1 Output Screenshots

PCA.HTML

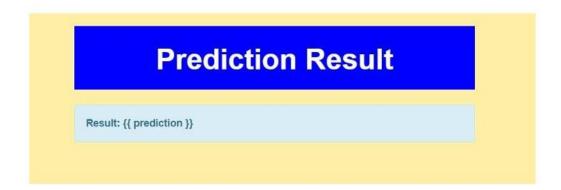
HOME PAGE



Age:	Gender:
651	1
Total_Bilirubin:	Direct_Bilirubin:
0.7	0.1
Alkaline_Phosphotase:	Alamine_Aminotransferase:
187	16
Aspartate_Aminotransferase:	Total_Protiens:
18	6.8
Albumin:	Albumin_and_Globulin_Ratio:
3.3	0.90

OUTPUT PAGE

RESULT.HTML



7. Advantages and disadvantages

7.ADVANTAGES AND DISADVANTAGES

Advantages:

- A. It can be widely used in Medical Field to predict liver disease.
- B. It would reduce the burden on doctors and medical staff leading to a more accurate treatment of the patients.

Disadvantages:

- A. The cost of misclassification of a single patient can be very high.
- B. Even the best performing Machine Learning Algorithm 20

8. Conclusion

The main objective of this study was to provide a summary of the classification algorithms in the field of data driven predictive data for liver disease. In this study, various classification algorithms were analysed to help doctors predict liver disease early. The purpose of this study was achieved by conducting comparative research in various papers. Based on this study, Random Forest has a much higher accuracy than other algorithms and can be used continuously in predicting user-recommended liver disease.

9. FUTURE SCOPE

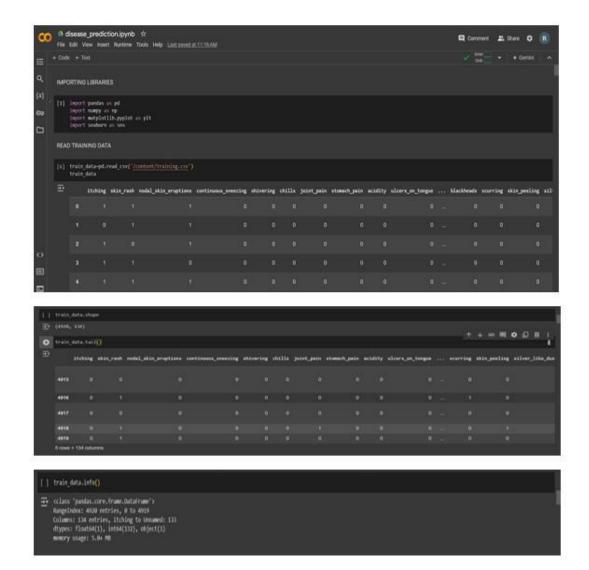
- Improved Accuracy: As more data becomes available and algorithms become more sophisticated, the accuracy of predicting liver disease will continue to improve. This includes both diagnosing the presence of liver disease and predicting its progression.
- Early Detection: Machine learning models can potentially detect liver disease at earlier stages than traditional methods, allowing for timely interventions and better outcomes for patients.
- Personalized Medicine: ML algorithms can analyze large datasets to identify patterns and correlations that can lead to personalized treatment plans. This could involve tailoring treatment strategies based on individual patient data, genetics, lifestyle factors, etc.
- Integration with Healthcare Systems: There is a growing trend towards integrating machine learning models into electronic health records (EHRs) and clinical decision support systems. This integration can streamline diagnosis and treatment planning processes.
- Feature Importance and Interpretability: Future research can focus on improving the interpretability of machine learning models used for liver disease prediction. Understanding which features (e.g., biomarkers, genetic markers) contribute most to predictions can enhance medical understanding and guide further research.
- Handling Imbalanced Data: Imbalanced datasets (where one class is more prevalent than another) are common in medical data. Future research can explore techniques to handle

- imbalanced data effectively, ensuring that models are robust and reliable.
- Telemedicine and Remote Monitoring: ML models can support telemedicine initiatives by enabling remote monitoring of liver disease patients. This can include predicting exacerbations, monitoring treatment effectiveness, and providing real-time alerts for healthcare providers.
- Drug Discovery and Development: Machine learning can aid in identifying potential drug targets and predicting the efficacy of treatments for liver disease. This can accelerate the drug discovery process and improve outcomes for patients with liver diseases.
- Ethical Considerations: As with any AI application in healthcare, there will be ongoing discussions about ethical considerations such as patient privacy, consent, and the responsible use of AI in medical decision-making.
- Global Health Impact: Machine learning can potentially have a significant impact on global health by improving access to diagnostic tools and personalized treatment options, particularly in underserved areas where access to healthcare is limited.

10.Appendix

10.1. Source Code

Code Snippets

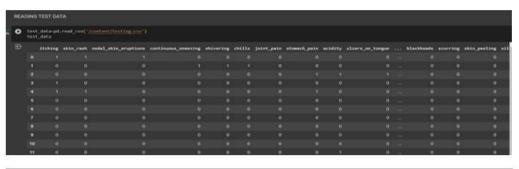




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4910		- 1			10	1 3	0.			. 0	6			



tre	in data describ										
	Stehing	skin_rash	model_skir_eruptions	continuous_sneesing	shivering	shille	300st_pain	storach_pain	ecidity	silders_un_tongue	pos_filled_pisples
.00	wet 4000.000000	4020.000000	4000,000000	4920.000000	4800.000000	4920.000000	4920.000000	4020.000000	4820.000000	4323.000000	4920.000000
	ean 0.137905	0.199756	0.021961	0.045122	0.021961	0.162196	0.139024	0.045122	0.046122	0.021961	0.021964
- 23	MF 0.344730	0.366417	0.146639		0.146539	0.366667	0.546007	0.297500	0.207593	0.146539	0.146539
	en 0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000
	9% 0.000000	0.000000	0.000000	0.000000	0.000000			0.000000	0.000000	0.000000	0.000000
	PK 0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000
	9% 0.000000	8.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	8.000000	8.000000
	unt 1,000000	1.000000	1.000000	1.000000	1.000000	1.000000	1.0000000	1.000000	1.000000	1.000000	1.000000



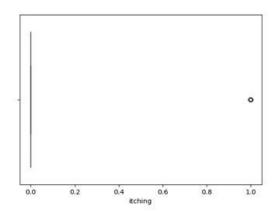


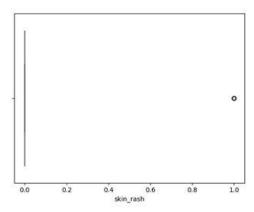
HAN	OLING MISSING VALUES	IN TEST DATA					
	test_data.ismull().sum						
	itching skin rush nodal_skin_eruptions continuous_sneezing shivering						
	Inflammatory nails blister red sore around tense yellow crust coce prognosis Length: 131, dtype: in	 9 0 0 0 0					
[11]	test_data.drop("fluid_	overload',axis -	1,implace-true)				

HSCRIPTIN	E ANALISIS														
D test	text_data_describe()														
	itching	ekin_resh	nodal_skin_eruptions	continuous_sneezing	shivering	chills	joint_pain	stomach_pain	ecidity	ulcers_on_tongue		pus_filled_pisples	blackheads		
coun		42.000000	42 000000				42.000000					42,000000			
mear	0.166667	0.190476	0.023810	0.047619	0.023610	0.166667	0.142857	0.047619	0.047619	0.023810		0.023810	0.023810		
stat		0.397437	0.154303	0.215540	0.154303		0.354189	0.215540	0.215540	0.154303			0.154303		
min	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000		0.000000	0.000000		
25%	0.000000	0.000000	0.000000	8.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000		0.000000	0.000000		
50%	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000		0.000000	0.000000		
75%	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000	0.000000		0.000000	0.000000		
max	1.000000	1.000000	1,000000	1.000000	1,000000	1.000000	1.000000	1.000000	1.000000	1.000000		1.000000	1.000000		
& rows	* 131 column														

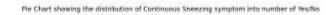
```
CHECKING OUTLAYERS BY VISUALIZING DATA

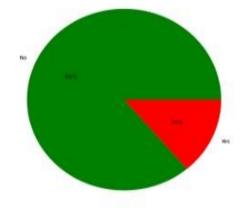
def func(col):
    sns.boxplot(xcol,data-train_data)
    plt.show()
    for i in train_data.columns:
        func(i)
```

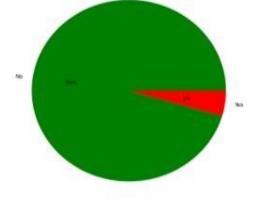


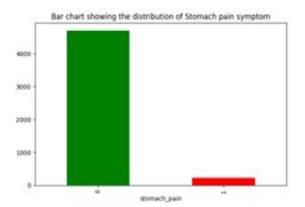


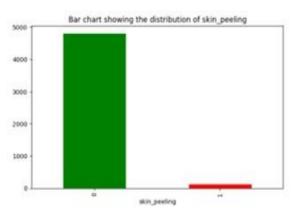
Pie chart showing the distribution of Itching symptom into number of Yes/No







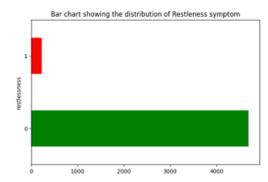


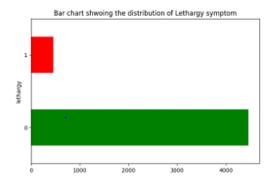


```
[ ] plt.subplot(1,2,1)
train_data['restlessess'].value_counts().plot(kind = 'barh', color = ['g','r'])
plt.title("War chart showing the distribution of Westleness symptom")

plt.subplot(1,2,2)
train_data['lethurgy'].value_counts().plot(kind = 'burh',color = ['g','r'])
plt.title("War chart showing the distribution of tethurgy symptom")

plt.subplots_adjust(left = 0.5,right = 2.5)
```





```
train_data['inflamatory_nails'].value_counts().sort_index().plot.line()
```

```
Bivariate Analysis

• a - len(train_data|train_data|'prognosis'] -- 'Fungal infection'])

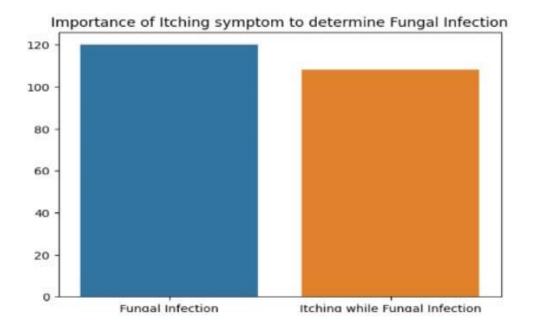
• b - len(train_data|(train_data|'itching'] -- 1) & (train_data|'prognosis'] -- 'Fungal infection')])

fi - pd.OstaFrame(data - [a,b], columes-['values'],index - ['Fungal Infection', 'Itching while Fungal Infection'])

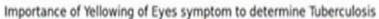
sns.barplot(data=fi, x=fi.index, y=fi['values'], color='skyblue')

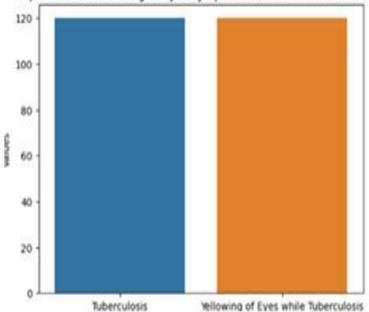
sns.harplot(data = fi , x = fi.index, y= fi['values'])

plt.title('importance of Itching symptom to determine Fungal Infection')
```



```
    a = len(train_data[train_data['prognosis'] == 'Tuberculesis'])
    b = len(train_data[(train_data['yelinwing of_eyes'] == 1) & (train_data['prognosis'] == 'Tuberculesis')))
    fi = pd.Dutaframe(data = [a,b], columns['Values'],index = ['Tuberculesis','yellowing of eyes shile scurring'])
    sns.barplot(data-fi, x-fi.index, y-fi['Values'], color-'green') = (thengof inlor in 'green'
    sns.barplot(data = fi , x = fi.index, y-fi['Values'])
    plt.title('Importance of yellowing of eyes symptom to determine Suberculesis')
```





multivariate analysis

corr=train_data.corr()
corr.style.background_gradient('coolwarm')

	itching	skin_rash	nodal_skin_eruptions	continuous_sneezing	shivering	chills	joint_pain	stomach_pain	acidity	ulcers_on_tongue	muscle_wasting
itching	1.000000	0.318158	0.326439	-0.086906	-0.059893	-0.175905	-0.160650	0.202850	-0.086906	-0.059893	-0.059893
skin_rash	0.318158	1.000000	0.298143	-0.094786	-0.065324		0.171134	0.161784	-0.094786	-0.065324	
nodal_skin_eruptions	0.326439	0.298143	1.000000		-0.022444	-0.065917	-0.060200	-0.032566	-0.032566	-0.022444	
continuous_sneezing	-0.086906	-0.094786	-0.032566	1.000000	0.608981	0.446238	-0.087351	-0.047254	-0.047254	-0.032566	
shivering	-0.059893		-0.022444	0.608981	1,000000	0.295332	-0.060200	-0.032566	-0.032566	-0.022444	
chills	-0.175905		-0.065917	0.446238	0.295332	1.000000	-0.004688	-0.095646	-0.095646	-0.065917	-0.065917
joint_pain	-0.160650	0.171134	-0.060200	-0.087351	-0.060200	-0.004688	1.000000	-0.087351	-0.087351	-0.060200	
stomach_pain	0.202850	0.161784	-0.032566	-0.047254		-0.095646	-0.087351	1,000000	0.433917	0.649078	
acidity	-0.086906	-0.094785	-0.032566		-0.032566	-0.095646	-0.087351	0,433917	1.000000	0.608981	
ulcers_on_tongue	-0.059893		-0.022444				-0.060200	0.649078	0.608981	1.000000	-0.022444
muscle_wasting	-0.059893	-0.065324	-0.022444		-0.022444	-0.065917	-0.060200	-0.032566	-0.032566	-0.022444	1.000000
vomiting	-0.057763	-0.225046	-0.119543	-0.173459	-0.119543	0.144263	0.199921			0.153603	-0.119543
burning_micturition	0.207896	0.166507	-0.032103			-0.094285	-0.086108	0.412239	-0.046581	-0.032103	
spotting_urination	0.350585	0.298143	-0.022444		-0.022444	-0.065917	-0.060200	0.608981	-0.032566	-0.022444	
fatigue	0.069744	-0.105248	-0.120465		-0.120465	0.269437		-0.174797	-0.174797	-0.120465	-0.120465
weight_gain	-0.061573	-0.067156	-0.023073			-0.067765	-0.061889	-0.033480	-0.033480	-0.023073	-0.02307
anxiety	-0.061573	-0.067156	-0.023073	-0.033480	-0.023073	-0.067765	-0.061889	-0.033480	-0.033480	-0.023073	-0.023073

DATA PREPROCESSING

• train_data.drop(['weight_gain', 'cold_hands_and_feets', 'anxiety', 'irregular_sugar_level',

'yellow_urine', 'acute_liver_failure', 'swelling_of_stomach', 'drying_and_tingling_lips', 'continuous_feel_of_urine',

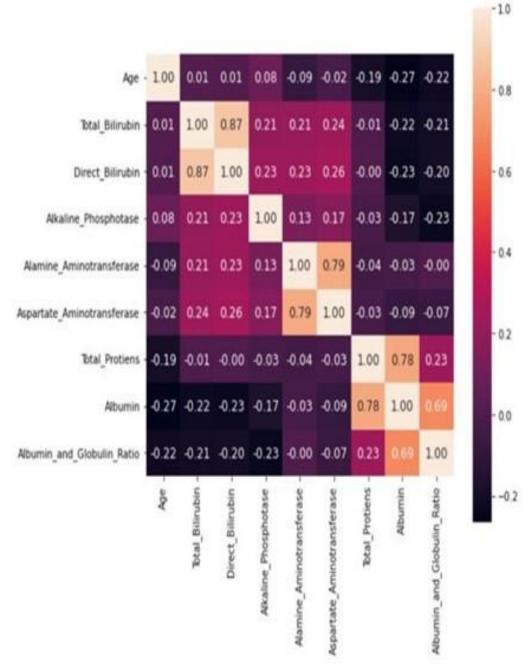
'internal_itching', 'polyuria', 'mood swings', 'receiving unsterile injections', 'stomach bleeding', 'prominent veins on calf', 'loss of smell', 'throat irritation',

'redness of eyes', 'sinus_pressure', 'runny_nose', 'pain_during_bowel_movements', 'pain_in_anal_region', 'cramps', 'bruising', 'enlarged_thyroid', 'brittle_nails',

'swollen_extremeties', 'slurred_speech', 'distention_of_sbdomen', 'fluid_overload.1', 'skin_peeling', 'silver_like_dusting', 'small_dents_in_sails', 'blister',

'red_sore_around_nose', 'bloody_stool', 'swollen_blood_vessels', 'hip_joint_pain',

'painful walking', 'spinning movements', 'altered sensorium', 'toxic look (typhos)'], axis -1, inplace - True



```
    * Irvin a Random torest Classifler and calculate accuracy
rfc - HandburgestClassifler(random_state=42)
rfc.filtOm_train_yl_train)
y_pred_rfc > rfc.predict(X1_val)

     } y_peed = rfc.predict(xt_val)
yt_pend = rfc.predict(xt_val)
y_pend = rfc.
    [94] from sklearn.metrics import confusion_matrix
                             * Calculate and print the confusion matrix cm - confusion_matrix(y1_val, y_pred_rfc)
     [89] from sklearn.svm import SVC
                                svm1.fit(X1_train,y1_train)
y_pred_svc - svm1.predict(X1_val)
y_pred - svm1.predict(X1_val)
                               yr_pred = swmi.predict(xi_ctain)
y_predi = swmi.predict(xi_ctain)
print('the Training Accuracy of the algorithm is',accuracy_score(yi_train,yt_pred))
print('the Validation Accuracy of the algorithm is',accuracy_score(yi_val,y_pred))
print('the Testing Accuracy of the algorithm is',accuracy_score(y_test,y_predi))

    The Training Accuracy of the algorithm is 0.9930313588850174
    the Validation Accuracy of the algorithm is 0.9959349593495935
    the Testing Accuracy of the algorithm is 1.0

         [95] from sklearn.metrics import confusion_matrix
                                           # Calculate and print the confusion matrix
cm = confusion_matrix(y1_val, y_pred_svc)
            ① [[32 0 0 ... 0 0 0]
[0 39 0 ... 0 0 0]
[0 0 41 ... 0 0 0]
                                                     ...
[ 0 0 0 ... 36 0 0]
[ 0 0 0 ... 0 37 0]
[ 0 0 0 ... 0 0 39]]
```

```
| Non-midphenclasifier() | Non-frick(non-production) | Non-production | No
```





PCA.HTML:

<!DOCTYPE html>

```
<html lang="en">
<head>
 <title>Liver Patient Analysis</title>
 <meta charset="utf-8">
 <meta name="viewport" content="width=device-width, initial-</pre>
scale=1">
 link rel="stylesheet"
href = "https://maxcdn.bootstrapcdn.com/bootstrap/3.3.7/css/bootstrap.\\
min.css">
 <style type="text/css">
  body {
   background-color: #ffcf0059;
  }
  nav {
   background-color: #ad38c2;
   height: 60px;
  .navbar-brand {
   color: white;
   font-size: 30px;
  nav ul li a {
   color: white;
   font-size: 20px;
```

```
</style>
</head>
<body>
<nav class="navbar">
 <div class="container-fluid">
  <div class="navbar-header">
  <a class="navbar-brand">Liver Patient Analysis</a>
 </div>
  <a href="#">Home</a>
  <a href="/predict">Goto Predict</a>
 </u1>
 </div>
</nav>
<div class="container">
 <h3>Introduction</h3>
```

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 paper is to analyze the parameters of various classification algorithms and compare their predictive accuracies to find the best classifier for determining liver disease. This paper focuses on the related works of various authors on liver disease such that algorithms were implemented using Weka tool that is a machine

learning software written in Java. Various attributes that are essential in the prediction of liver disease were examined and the dataset of liver patients was also evaluated. This paper compares various classification algorithms such as Random Forest, Logistic Regression

```
</div>
</body>
</html>
```

Result.html:

```
<!DOCTYPE html>
<html>
<head>
<title>Prediction Result</title>
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>Prediction Result</h1>
    </div>
   </div>
  </div>
 </div>
 <div class="container">
  <div class="row">
   <div class="col-md-3"></div>
   <div class="col-md-6">
```

App.py:

```
# import numpy as np
# from flask import Flask, request, jsonify, render_template
# import pickle

# app = Flask(__name__)
# model = pickle.load(open('indian_liver_patient.pkl', 'rb'))

# @app.route('/')
# def home():
# return render_template('index.html')

# @app.route('/predict', methods=['POST'])
```

```
# def predict():
#
    For rendering results on HTML GUI
#
#
#
    try:
       int features = [int(x) \text{ for } x \text{ in request.form.values()}]
#
       final features = [np.array(int features)]
#
       prediction = model.predict(final features)
#
#
       output = round(prediction[0], 2)
       return render template('index.html', prediction text='Predicted
Value: $ {}'.format(output))
#
    except Exception as e:
       return str(e)
#
# @app.route('/predict api', methods=['POST'])
# def predict api():
#
#
    For direct API calls through request
#
#
    try:
       data = request.get json(force=True)
#
#
       prediction = model.predict([np.array(list(data.values()))])
       output = prediction[0]
#
#
       return jsonify(output)
#
    except Exception as e:
#
       return jsonify({'error': str(e)})
```

```
# if name == " main ":
    app.run(debug=True)
"""from flask import Flask, request, render template
import pandas as pd
import pickle
app = Flask( name )
# Load the trained model
try:
  with open('indian liver patient.pkl', 'rb') as model file:
    model = pickle.load(model file)
except (EOFError, FileNotFoundError) as e:
  model = None
  print(f"Error loading model: {e}")
@app.route('/')
def home():
  return render template('index.html')
@app.route('/predict', methods=['GET', 'POST'])
def predict():
  if request.method == 'POST':
```

```
if model is None:
          raise ValueError("Model not loaded properly")
       # Get input data from form
       data = request.form.to dict()
       data = {k: float(v) for k, v in data.items()}
       # Convert to DataFrame
       df = pd.DataFrame([data])
       # Make prediction
       prediction = model.predict(df)
       # Return result
       result = 'Liver disease' if prediction[0] == 1 else 'No liver
disease'
       return render template('result.html', result=result)
     except Exception as e:
       return str(e)
  return render template('predict.html')
if name == " main ":
  app.run(debug=True)"""
```

try:

```
from flask import Flask, request, render template
import pandas as pd
import pickle
app = Flask( name )
# Load the model once during the app initialization
model = pickle.load(open('liver analysis.pkl', 'rb'))
@app.route('/')
def home():
  return render template('index.html')
@app.route('/predict')
def index():
  return render template('predict.html')
@app.route('/data predict', methods=['POST'])
def predict():
  if request.method == 'POST':
     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(tb), float(db), float(ap),
float(aa1), float(aa2), float(tp), float(a), float(agr)]]
     prediction = model.predict(data)[0]
     if prediction == 1:
       result = "You have a liver disease"
     else:
       result = "You don't have a liver disease"
     return render template('result.html', prediction=result)
if __name__ == '__main__ ':
  app.run(debug=True)
```

10.2 GitHub and project Demo link:

Github link: Click Here

Project Demo link: Click Here