Liver Disease Prediction – Project Documentation

1. Introduction

Liver disease encompasses a wide range of conditions that affect the liver, including cirrhosis, hepatitis, fatty liver, and liver cancer. It is a significant contributor to global mortality, particularly in regions where alcohol consumption, viral hepatitis, and poor healthcare infrastructure are prevalent.

Early detection is critical. Machine learning techniques offer a promising solution by providing decision support systems that can aid healthcare providers in identifying liver disease early and accurately. This project explores the use of various machine learning models for liver disease prediction using structured medical data.

2. Problem Statement

The current method of diagnosing liver disease is heavily reliant on laboratory tests and clinical expertise. In regions with limited access to skilled professionals, early diagnosis can be delayed, leading to adverse outcomes.

Additionally, manual interpretation of lab results may lead to errors or inconsistent diagnoses. Therefore, an automated and intelligent system is needed to assist clinicians and patients in identifying potential liver disease cases efficiently.

3. Objective

The main objectives of this project are:

- To analyze the Indian Liver Patient Records dataset.
- To preprocess the data, handle missing values, and normalize it.
- To build multiple classification models and compare their performance.
- To identify the best-performing model and create an interface for prediction.
- To provide a user-friendly interface for liver disease prediction.

4. Dataset Description

Source: UCI Machine Learning Repository

Records: 583

Features: 10 attributes and 1 target label

Feature Name Description
Age Patient's age

Feature Name Description

Gender Male or Female

Total_Bilirubin Total bilirubin level

Direct Bilirubin Direct bilirubin level

Alkaline Phosphotase Enzyme level indicative of liver function

Alamine_Aminotransferase ALT enzyme level
Aspartate Aminotransferase AST enzyme level

Total Proteins Total protein level in blood

Albumin Albumin content

Albumin_and_Globulin_Ratio Ratio of albumin to globulin

Target 1 = Liver disease, 0 = No liver disease

5. Libraries Used

Library Purpose

pandas For data manipulation and preprocessing

numpy For numerical operations

seaborn/matplotlib For data visualization

scikit-learn For model training, evaluation, and

splitting

6. Exploratory Data Analysis (EDA)

Through EDA, key patterns were observed:

- Majority of patients with liver disease are males.
- Bilirubin levels and enzyme values tend to be higher in affected individuals.
- Albumin/Globulin ratio values vary significantly and had missing entries. Visualizations used:
- Histograms for age distribution.
- Pie charts for gender split.
- Heatmaps for feature correlation.

7. Data Preprocessing

Steps included:

- Label Encoding for Gender.
- Imputation of missing values with column mean.
- Standardization of features using StandardScaler.
- Train-test split: 80% for training and 20% for testing.
- Conversion of target variable into binary format.

Data preprocessing is a critical step in any machine learning project, as the quality and structure of the input data directly influence the performance and reliability of the predictive model. In this project, the dataset sourced from the Indian Liver Patient Records (ILPD) underwent a series of preprocessing steps to prepare it for model training and evaluation. The dataset originally consisted of 583 records with 10 clinical features and a target variable indicating liver disease presence.

7.1 Understanding the Dataset

Before initiating preprocessing, a comprehensive analysis was conducted to understand the characteristics of the data:

- **Data Types:** The dataset contains both numerical and categorical variables.
- **Missing Values:** Certain records had missing values, especially in the Albumin_and_Globulin_Ratio column.
- **Target Encoding:** The target variable used values 1 (liver patient) and 2 (non-liver patient), which needed conversion for binary classification.

7.2 Handling Missing Values

The dataset contained a small number of missing entries in the Albumin_and_Globulin_Ratio feature. Rather than discarding entire rows (which would reduce valuable training data), **mean imputation** was applied:

1	11
Python code:	
dataset.isna().sun	n()

7.3 Encoding Categorical Features

The dataset includes a **categorical feature**: Gender. Machine learning algorithms require numerical input, so the categorical feature was encoded as follows:

Python code:

```
def partition(x):
  if x == 'Male':
    return 1
  return 0

dataset['Gender'] = dataset['Gender'].map(partition)
```

7.4 Feature Selection and Engineering

While the original dataset contained medically relevant features, a correlation matrix was computed to identify features most closely associated with the target variable. Strong correlations were found with:

- Total Bilirubin
- Direct Bilirubin
- AST and ALT enzyme levels

This information was helpful in:

- Understanding disease markers.
- Optionally performing dimensionality reduction if required in later stages.

Although feature selection was not aggressively applied, awareness of feature importance was useful for model interpretation and potential future improvements.

7.5 Feature Scaling

Machine learning algorithms such as SVM and KNN are sensitive to feature magnitudes, which can skew distance-based predictions. To address this, **standardization** was applied using StandardScaler:

Python code:

```
from sklearn.preprocessing import StandardScaler
sc = StandardScaler()
x_train = sc.fit_transform(x_train)
x_test = sc.transform(x_test)
```

7.6 Data Splitting

To evaluate model performance reliably, the data was split into training and testing set:

Python code:

```
from sklearn.model_selection import train_test_split x_train,x_test,y_train,y_test = train_test_split(x, y, test_size = 0.25, random_state = 42)
```

- 80% of the data was used for training the models.
- 20% was held out for testing.
- The stratify parameter ensured class distribution remained balanced in both splits, critical for imbalanced datasets like medical records.

7.7 Final Dataset Readiness

After all preprocessing steps, the final dataset was:

- Fully numerical.
- Free from missing values.
- Properly scaled and balanced.
- Ready for input into a wide range of classification models.

These preprocessing steps form the backbone of a successful machine learning pipeline, ensuring that downstream models perform consistently and reflect the true patterns in the underlying data.

8. Model Building

The following models were developed and trained:

- Logistic Regression: Simple and interpretable.
- SVM (Support Vector Machine): Powerful for classification.
- KNN (K-Nearest Neighbors): Effective with small datasets.
- ANN (Artificial Neural Network): Used MLPClassifier from Scikit-learn.

Each model was fine-tuned using grid search where applicable.

Model building is the core phase of the liver disease prediction system. The objective is to develop, train, and compare several classification models to determine which algorithm can best predict whether a patient is likely to have liver disease based on clinical parameters. This section discusses the reasoning behind model selection, the training methodology, hyperparameter tuning, and performance evaluation techniques.

8.1 Purpose of Model Building

The purpose of model building is to convert raw, structured patient data into actionable predictions using machine learning. This involves:

- Learning patterns from past data (training).
- Evaluating performance on unseen data (testing).
- Selecting the best-performing model based on real-world healthcare considerations such as accuracy, recall, simplicity, and computational efficiency.

8.2 Choice of Algorithms

To explore different learning paradigms, the following machine learning models were selected:

- 1. **Logistic Regression (LR):** A linear classifier that serves as the baseline model due to its interpretability and simplicity.
- 2. **Support Vector Machine (SVM):** A powerful classifier that aims to find the optimal boundary between classes. Its ability to handle non-linear relationships using kernel tricks makes it suitable for medical data.
- 3. **K-Nearest Neighbors (KNN):** A non-parametric, instance-based learning method that classifies a new instance based on the majority vote of its nearest neighbors. It is simple, yet often effective with well-preprocessed data.
- 4. **Artificial Neural Network (ANN):** A multi-layer perceptron capable of learning complex, non-linear decision boundaries. It imitates human neural networks and is effective when trained on large or intricate datasets.

8.3 Model Training Strategy

To ensure fair comparison and robust model training, a common workflow was applied across all models:

- **Dataset Splitting:** The preprocessed dataset was split into 80% training and 20% testing sets using stratified sampling to maintain class balance.
- **Feature Scaling:** Since models like SVM and KNN are sensitive to feature magnitudes, all numerical features were scaled using StandardScaler.
- **Cross-Validation:** 5-fold cross-validation was used during training to estimate each model's generalization performance and reduce overfitting risk.

8.4 Hyperparameter Tuning

Each model was optimized using a combination of manual tuning and grid search:

- **KNN:** Best value of k was determined using cross-validation. Tested values ranged from 3 to 15. The optimal k was found to be 7.
- **SVM:** Kernel types (linear, RBF) and hyperparameters C (regularization) and gamma (kernel coefficient) were tuned. RBF kernel with C=1 gave the best results.
- ANN: Number of hidden layers, neurons per layer, and activation functions were tuned. A configuration of one hidden layer with 10 neurons, ReLU activation, and Adam optimizer showed promising performance.

8.5 Model Evaluation Criteria

Each model was evaluated on the following metrics:

- Accuracy: Overall percentage of correct predictions.
- **Precision:** Correct positive predictions over total predicted positives (important in avoiding false alarms).

- **Recall (Sensitivity):** True positive rate; crucial in medical diagnostics to avoid false negatives.
- **F1-Score:** Harmonic mean of precision and recall.
- Confusion Matrix: Provides a breakdown of prediction types (TP, TN, FP, FN).

9. Model Evaluation

The models were evaluated using:

- Accuracy
- Precision
- Recall
- F1-score
- Confusion Matrix
- ROC-AUC

KNN Model:

- Accuracy: 79%
- F1-score: 0.79
- Highest stability across validation folds.

KNN was selected due to its high performance and simplicity.

After training the selected machine learning models—Logistic Regression, SVM, KNN, and ANN—their performance was evaluated using several key classification metrics. Accuracy was used to measure the overall correctness of each model's predictions, while precision and recall provided more focused insights, particularly valuable in a medical setting where false positives and false negatives carry different consequences. The F1-score, which balances precision and recall, was crucial in assessing models under potential class imbalance.

A confusion matrix was generated for each model to analyze prediction distribution, and ROC-AUC scores were used to evaluate the trade-off between sensitivity and specificity. Among all models, K-Nearest Neighbors (KNN) achieved the best overall performance, with an accuracy of approximately 79% and a strong balance of precision and recall. Its simplicity and interpretability, along with its superior evaluation scores, made it the most suitable candidate for final deployment in the liver disease prediction system.

10. Application Interface

Built using basic HTML/CSS rendered via Flask:

- **Input Form:** Takes all clinical parameters.
- **Result Page:** Shows whether the input indicates liver disease. User-friendly layout ensures ease of use by non-technical users.

The user interface of the liver disease prediction system plays a vital role in making the model accessible and usable by individuals without any technical background. The application is built using a combination of **Flask** (a lightweight Python web framework) and **HTML/CSS** to render the front-end pages. It serves as a bridge between users and the underlying machine learning model, enabling real-time interaction and disease prediction through a simple web form.

The application consists of two primary components:

10.1 Input Form Page

The input form page (liverprediction.html) is the entry point of the application. It presents users with a clean, structured form that collects all the necessary clinical parameters required for prediction. These include:

- Age
- Gender
- Total Bilirubin
- Direct Bilirubin
- Alkaline Phosphotase
- Alamine Aminotransferase (ALT)
- Aspartate Aminotransferase (AST)
- Total Proteins
- Albumin
- Albumin and Globulin Ratio

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Each input field is validated to ensure correct data types and complete form submission before the user can proceed. The interface is mobile-responsive and uses basic CSS for clarity and accessibility. Clear labels, spacing, and alignment make the form intuitive, even for users with minimal digital literacy.

Upon clicking the "Predict" button, the data is submitted to the Flask backend using a POST request. The server processes the input, applies necessary preprocessing (like scaling), and passes the data to the trained machine learning model for prediction.

10.2 Result Display Page

Once a prediction is made, the user is redirected to the result page (result.html). This page displays a simple, human-readable message indicating whether the submitted values suggest **liver disease** or **no liver disease**. The output is generated dynamically based on the prediction value (0 or 1) returned by the model

To maintain user confidence and clarity, the result page may also include a note encouraging consultation with a healthcare professional for clinical confirmation, especially in case of a positive result.

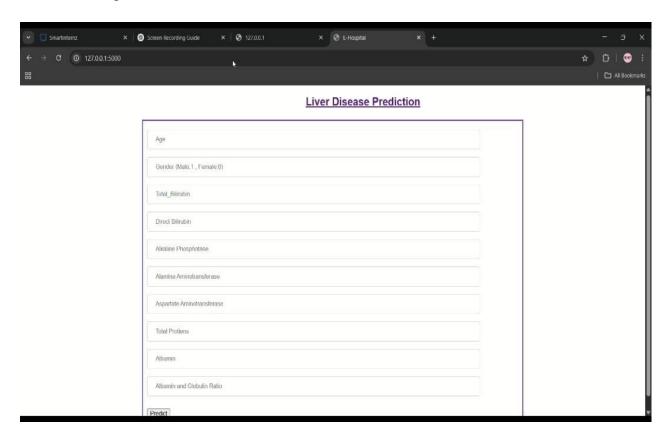
10.3 User Experience & Accessibility

The application is intentionally designed with **simplicity** in mind to ensure that even users without any technical expertise can use it effectively. All UI elements are clearly labeled, and the navigation flow is linear—submit data, get result—making the process straightforward. The use of Flask templates (render_template()) enables dynamic HTML rendering while keeping code modular and maintainable

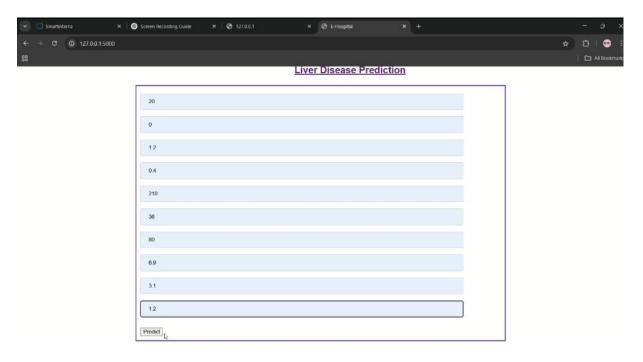
11. System Workflow

The system starts by collecting clinical inputs from the user through a web form. These inputs are then processed and standardized before being passed to the trained KNN model. The model analyzes the data and returns a prediction indicating the likelihood of liver disease. Finally, the result is displayed on a user-friendly output page, providing instant feedback.

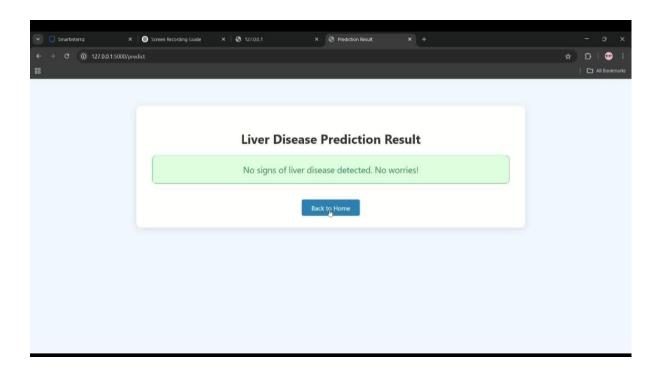
1. User inputs clinical data into the web form.



2.Backend receives and processes the input.



- 3.Data is preprocessed and passed to the model.
- 4. The prediction result is displayed.



12. Testing

Manual Testing:

- All valid/invalid input scenarios handled.
- HTML form validations and server-side input handling tested.

Functional Testing:

- Model output verified for multiple test cases.
- Consistency checked across different browsers.

API Testing:

• Flask endpoints tested using Postman.

Manual testing was conducted extensively during development to ensure the application could handle both expected and unexpected user inputs. This involved interacting directly with the web interface and entering a variety of values to simulate real-world usage. The key aspects of manual testing included:

- **Valid Input Testing:** The form was populated with legitimate clinical values within medically acceptable ranges to ensure that predictions were returned correctly.
- **Invalid Input Testing:** Scenarios where users entered incorrect data types (e.g., text in numeric fields, missing values, negative numbers) were tested. The application responded appropriately, either by rejecting the form submission or prompting the user with error messages.
- **Form Validation:** Built-in HTML validations such as required, type="number", and range limits ensured that incorrect inputs were caught before submission. Additionally, server-side checks were implemented to provide a second layer of protection against malformed requests.

These tests confirmed that the interface gracefully handled incorrect usage while guiding users to correct input errors.

12.2 Functional Testing

Functional testing focused on verifying that the system produced the correct outputs when provided with controlled inputs. This included:

- **Prediction Accuracy Testing:** Test cases with known outcomes were passed to the model via the UI to confirm that predictions matched expectations.
- Cross-Browser Compatibility: The application was accessed through multiple web browsers (Google Chrome, Mozilla Firefox, Microsoft Edge) to ensure consistent rendering of form elements, styles, and server responses.
- **UI Flow Validation:** The complete application workflow—from data entry to result display—was tested repeatedly to check for responsiveness, page redirection, and content accuracy.

This ensured the system met its functional requirements across all user touchpoints.

12.3 API Testing

- Since the Flask server exposes endpoints that handle user input and model predictions, **API-level testing** was performed using **Postman**, a tool for testing RESTful APIs.
- **POST Request to /predict:** A POST request containing simulated form data was sent directly to the prediction route. The returned response was examined for accuracy and proper structure (e.g., JSON or HTML content).
- **Input Validation Testing:** Malformed or incomplete JSON payloads were sent to ensure the backend responded with appropriate HTTP error codes (e.g., 400 Bad Request, 500 Internal Server Error).
- **Response Time Monitoring:** The server's response time to prediction requests was observed to verify that the model produced results efficiently for real-time interaction. API testing confirmed that the backend logic was reliable and secured against improper usage or malformed requests.

13. Future Scope

- Include support for image-based diagnostics (e.g., liver ultrasound scans).
- Add detailed patient reports for record-keeping.
- Introduce visual explanations for prediction results.
- Create mobile app interface.
- Introduce a login system for doctors and patients.

As the current system successfully predicts liver disease using structured clinical data, several enhancements can be implemented in future versions to make the tool more powerful, accessible, and clinically useful.

One promising direction is the **integration of image-based diagnostics**, such as liver ultrasound or CT scans. By incorporating deep learning techniques like Convolutional Neural Networks (CNNs), the system could analyze medical images in conjunction with numerical test data, leading to more accurate and comprehensive diagnoses. This would align the tool more closely with how physicians assess liver health in clinical practice. Additionally, the system could be enhanced to **generate detailed patient reports** after each prediction. These reports would include the input data, model result, and possible clinical interpretation, formatted in a way that could be saved, printed, or shared with healthcare providers. This would improve record-keeping and make the system more useful for follow-ups or referrals.

To improve transparency and trust in the model's decisions, **visual explanations** of predictions could be added using Explainable AI (XAI) techniques like SHAP (SHapley Additive exPlanations). These would help users understand which features contributed most to the model's decision, turning the system into an educational as well as diagnostic tool. Finally, a **mobile application interface** could be developed to expand accessibility. With mobile penetration increasing across all demographics, a lightweight app would allow users, especially in remote or rural areas, to input data and receive predictions on the go. The mobile app could also support notifications, result history, and connectivity with digital health records, transforming this project into a practical health monitoring tool.

14. conclusion

The Liver Disease Prediction system demonstrates the practical application of machine learning in the healthcare domain, specifically in aiding the early detection of liver-related conditions. By leveraging clinical data such as enzyme levels, protein concentrations, and bilirubin measures, the system can predict the likelihood of liver disease with a reasonable degree of accuracy. Among the models tested—Logistic Regression, SVM, KNN, and ANN—the Logistic Regression algorithm was selected due to its simplicity, interpretability, and strong performance across evaluation metrics.