Genetic Disease Risk Prediction

## NITTE MEENAKSHI INSTITUTE OF TECHNOLOGY

(AN AUTONOMOUS INSTITUTION, AFFILIATED TO VISVESVARAYA TECHNOLOGICAL UNIVERSITY, BELGAUM, APPROVED BY AICTE & GOVT.OF KARNATAKA)

Yelahanka, Bangalore - 560 064



#### **COURSE PROJECT REPORT**

of

ARTIFICIAL INTELLIGENCE & MACHINE LEARNING (22CSG53)

on

## "GENECTIC DISEASE RISK PREDICTION"

Submitted in the Partial fulfillment of the requirements of Semester-5 of

#### **BACHELOR OF ENGINEERING**

IN

#### **COMPUTER SCIENCE & ENGINEERING**

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## **CERTIFICATE**

Certified that the Course Project Work titled "GENECTIC DISEASE RISK PREDICTION", carried out by Himanchi Kumari M bearing USN: 1NT22CS076, Neha G A bearing USN: 1NT22CS121, Name of the Chandana Priya bearing USN: 1NT23CS401 bonafide students of Nitte Meenakshi Institute of Technology in partial fulfilment of Semester- 5 of Bachelor of Engineering Degree in Computer Science & Engineering under Visvesvaraya Technological University, Belagavi during the year 2024-2025. It is certified that all corrections/ suggestions indicated for Internal Assessment have been incorporated in the Report deposited in the Departmental Library. The Learning Assessment-II Report has been approved as it satisfies the Academic requirements in respect of the Course Project Work prescribed for the said Degree.

Signature of Guide

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#### **DECLARATION**

#### We hereby declare that

- (i) The project work is our original work
- (ii) This Project work has not been submitted for the award of any degree or examination at any other university/College/Institute.
- (iii) This Project Work does not contain other persons' data, pictures, graphs or other information, unless specifically acknowledged as being sourced from other persons.
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## **Abstract**

The project focuses on predicting lung cancer levels (High, Medium, Low) using machine learning models, leveraging patient data for early detection and classification. The dataset includes key features describing patient health, and the target variable, level, is encoded into numeric categories for computational efficiency. Various machine learning models, including Logistic Regression, Random Forest, Naive Bayes, and Decision Tree, are trained and evaluated to determine the best-performing model.

Data preprocessing ensures the dataset's readiness for modeling. This includes cleaning column names, handling categorical variables, and splitting the data into training and testing sets. Feature-target separation enables supervised learning, where the models are trained on predictors (X) and evaluated against the target variable (y). Metrics such as accuracy, precision, recall, and F1-score are calculated to evaluate model performance. To identify the most suitable model, weighted averages of these metrics are used, considering class distribution through support.

The Random Forest and Multinomial Logistic Regression models demonstrate strong performance, achieving high accuracy on the test data. Comparisons are visualized using grouped bar charts, highlighting precision, recall, and F1-scores for each model across different cancer levels. Overfitting and underfitting are diagnosed by comparing training and testing accuracies, ensuring a robust and generalizable model.

The project concludes by selecting the best model based on overall metrics and accuracy. This predictive approach is a step towards leveraging machine learning for effective cancer diagnosis, supporting timely medical intervention, and improving patient.

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## LIST OF ACRONYMS

- 1. ML Machine Learning
- 2. AI Artificial Intelligence
- 3. **LR** Logistic Regression
- 4. **RF** Random Forest
- 5. **NB** Naive Bayes
- 6. KNN K-Nearest Neighbors
- 7. **SVM** Support Vector Machine
- 8. **DT** Decision Tree
- 9. **GA** Genetic Algorithm
- 10. ANN Artificial Neural Network
- 11. **CNN** Convolutional Neural Network
- 12. RNN Recurrent Neural Network
- 13. **IoT** Internet of Things
- 14. TP True Positive
- 15. FP False Positive
- 16. TN True Negative
- 17. FN False Negative
- 18. **ROC** Receiver Operating Characteristic
- 19. AUC Area Under the Curve
- 20. MSE Mean Squared Error
- 21. RMSE Root Mean Squared Error
- 22. F1-Score Harmonic mean of Precision and Recall
- 23. PCA Principal Component Analysis
- 24. API Application Programming Interface
- 25. CSV Comma-Separated Values

# **CHAPTER 1: INTRODUCTION**

#### 1.1 BACKGROUND

Lung cancer is a major global health concern, accounting for a significant percentage of cancerrelated deaths annually. Early detection and accurate classification of its severity are critical for improving patient survival rates. Traditional diagnostic methods, while effective, can be timeconsuming and resource-intensive. In this context, machine learning offers a promising alternative, providing data-driven solutions for faster and more precise predictions.

Machine learning models can analyze complex datasets, identifying patterns and relationships that may not be immediately apparent through conventional analysis. By utilizing patient data, including clinical and demographic features, these models can classify cancer severity into levels such as High, Medium, and Low. This capability enables healthcare professionals to prioritize high-risk patients and tailor treatment strategies accordingly.

The advent of machine learning in medical diagnostics marks a shift toward technology-driven solutions, emphasizing efficiency and accuracy. This project builds on this foundation, demonstrating the application of machine learning in lung cancer classification.

#### 1.2 BRIEF HISTORY OF TECHNOLOGY/CONCEPT

Machine learning, a subset of artificial intelligence, has evolved significantly since its inception in the mid-20th century. Early developments began with simple rule-based systems and statistical models aimed at solving classification problems. In the 1950s, Alan Turing proposed the concept of a machine capable of learning from data, paving the way for modern advancements.

The 1980s saw the emergence of algorithms like decision trees, neural networks, and support vector machines, which introduced the ability to handle complex datasets and nonlinear relationships. With the proliferation of computational power in the 1990s and 2000s, ensemble methods such as Random Forests and Gradient Boosting gained prominence for their robustness and accuracy.

In healthcare, machine learning applications started gaining traction with advancements in data collection and electronic health records. The ability to analyze patient data for predictive modeling has transformed medical diagnostics, particularly in oncology. Lung cancer prediction using machine learning became feasible as datasets grew in size and sophistication, enabling algorithms to identify patterns in clinical and demographic features.

Today, machine learning stands at the forefront of medical innovation. Its integration into diagnostics offers significant potential for early detection, personalized treatment, and improved

patient outcomes. This project utilizes such advancements to classify lung cancer severity levels, leveraging decades of progress in machine learning and healthcare analytics.

#### 1.3 APPLICATIONS

#### 1. Early Detection of Lung Cancer:

o Enables the identification of high-risk cases based on patient data, supporting timely medical intervention.

#### 2. Cancer Severity Classification:

o Classifies patients into severity levels (High, Medium, Low), helping prioritize critical cases for treatment.

#### 3. Personalized Treatment Planning:

 Provides insights into patient conditions, allowing healthcare professionals to design tailored treatment strategies.

#### 4. Reduction in Diagnostic Errors:

 Enhances accuracy in diagnosing cancer severity, minimizing human error in manual evaluations.

#### 5. Support for Medical Professionals:

 Assists doctors by providing data-driven recommendations for diagnosis and management.

#### 6. Efficient Resource Allocation:

o Helps healthcare systems allocate resources more effectively by identifying patients who need immediate attention.

#### 7. Integration into Telemedicine Platforms:

• Facilitates remote screening and monitoring of patients, especially in areas with limited access to healthcare facilities.

#### 8. Research and Development:

 Provides a framework for future studies in oncology, contributing to advancements in predictive modeling.

#### 9. Public Health Analytics:

 Offers aggregated insights for policymakers to design targeted cancer prevention programs.

## 10. Medical Training and Education:

 Serves as a case study to teach medical professionals about machine learning in diagnostics.

#### 1.4 RESEARCH MOTIVATION AND PROBLEM STATEMENT

#### 1.4.1 STATEMENT OF THE PROBLEM

The diagnosis and classification of lung cancer severity remain a challenge due to:

- 1. The time-consuming and labor-intensive nature of traditional diagnostic methods.
- 2. Variability in clinical expertise, leading to potential diagnostic errors.
- 3. The need for accurate classification of cancer levels (High, Medium, Low) to prioritize treatment.

4. Underutilization of machine learning capabilities for integrating and analyzing diverse patient data.

This research seeks to address these challenges by developing a robust machine learning framework capable of accurately classifying lung cancer severity levels. The objective is to provide an efficient, reliable, and scalable solution that complements existing diagnostic methods, improves decision-making, and ultimately enhances patient outcomes.

#### 1.5 RESEARCH OBJECTIVES AND CONTRIBUTIONS

#### 1.5.1 PRIMARY OBJECTIVES

The primary objective of this research is to develop and evaluate machine learning models for accurately classifying lung cancer severity levels (High, Medium, Low) based on patient data. This aims to enhance early detection, support clinical decision-making, and improve patient outcomes through efficient and reliable predictive analytics.

#### 1.5.2 MAIN CONTRIBUTIONS

#### 1. Development of a Predictive Framework:

 Designed a machine learning-based system for classifying lung cancer severity levels (High, Medium, Low).

## 2. Evaluation of Multiple Models:

Implemented and compared various models, including Logistic Regression,
 Random Forest, Naive Bayes, and Decision Trees, to identify the best-performing approach.

#### 3. Weighted Metric Analysis:

o Introduced a weighted evaluation method using precision, recall, and F1-score to account for class imbalance in performance assessment.

## 4. Overfitting and Underfitting Diagnosis:

 Conducted comprehensive analyses to ensure the model's generalizability and robustness.

#### 5. Visualization Tools:

 Created comparative visualizations for key metrics, providing insights into model performance for better interpretability.

#### 6. Real-World Applicability:

o Demonstrated how machine learning can be integrated into lung cancer diagnostics, supporting timely and accurate decision-making in healthcare.

#### 7. Foundation for Future Research:

 Established a framework that can be extended to other medical diagnostic tasks and larger datasets.

#### 1.6 ORGANIZATION OF THE REPORT

#### This report is structured as follows:

#### 1. **Introduction:**

Provides an overview of the project, its background, motivation, and objectives. Discusses the significance of lung cancer detection and the role of machine learning in healthcare.

#### 2. Literature Review:

Reviews existing methods for lung cancer diagnosis and classification, highlighting the gaps addressed by this research. Examines prior applications of machine learning in medical diagnostics.

#### 3. **Methodology:**

Details the dataset, preprocessing steps, and the machine learning models implemented. Describes feature engineering, target encoding, and the metrics used for evaluation.

#### 4. Implementation and Results:

Discusses the training and testing process for various models, presenting their performance metrics. Includes comparative analysis and visualizations to identify the best-performing model.

#### 5. Discussion:

Interprets the results, addressing issues of overfitting, underfitting, and model generalizability. Explores potential limitations and areas for improvement.

#### 6. Conclusion and Future Work:

Summarizes the findings, emphasizing the contributions of the project. Suggests future directions for extending the framework to other datasets or applications.

#### 7. References:

Lists all academic papers, datasets, and tools cited throughout the report.

This organization ensures a logical flow, guiding the reader through the problem, methodology, results, and implications of the research.

#### 1.7 SUMMARY

This report explores the use of machine learning models for lung cancer severity classification. It details the problem, reviews existing methods, explains the methodology and implementation, and evaluates model performance. The findings highlight the best-performing model and its potential for real-world applications. Limitations and future directions are discussed, emphasizing the project's contributions to healthcare diagnostics.

## CHAPTER 2: LITERATURE SURVEY

#### 2.1 INTRODUCTION

The literature survey serves as a foundation for understanding the current state of research and advancements in lung cancer diagnosis and machine learning applications. Lung cancer, being a leading cause of mortality, necessitates accurate and timely diagnosis. Traditional diagnostic methods, such as imaging and biopsies, are resource-intensive and may result in delayed treatment. This has driven the exploration of computational approaches to enhance efficiency and precision.

Machine learning has emerged as a transformative tool in healthcare, offering predictive capabilities for disease diagnosis, prognosis, and treatment planning. Studies have demonstrated its effectiveness in analyzing complex datasets, identifying patterns, and delivering reliable classifications. In the context of lung cancer, researchers have employed models such as Decision Trees, Random Forests, Logistic Regression, and Support Vector Machines to classify cancer severity levels and predict patient outcomes.

This chapter reviews the existing literature on lung cancer detection and classification using machine learning. It highlights the strengths and limitations of various approaches, identifies gaps in current research, and sets the stage for the methodology developed in this study. By examining prior work, this survey establishes the relevance of the proposed framework and its contributions to the field of predictive analytics in healthcare.

#### 2.2 RELATED WORK

Numerous studies have explored the application of machine learning to lung cancer detection and severity classification. These efforts have aimed to improve the accuracy and efficiency of traditional diagnostic methods.

### 1. Lung Cancer Detection Using Machine Learning Models:

Various studies have utilized supervised learning models like Logistic Regression, Random Forests, and Support Vector Machines to predict lung cancer presence and severity. These models excel at identifying nonlinear relationships in patient data, achieving high classification accuracy in controlled datasets.

## 2. Feature Engineering in Medical Diagnostics:

Feature selection and engineering have been emphasized in prior research to improve model performance. Studies have shown that selecting relevant clinical features, such as tumor size and biomarkers, significantly enhances the predictive capabilities of machine learning models.

#### 3. Deep Learning for Lung Cancer Classification:

Advanced techniques like Convolutional Neural Networks (CNNs) have been employed for image-based diagnosis, analyzing CT scans to classify lung cancer types. These

methods demonstrate exceptional performance but require substantial computational resources and extensive training data.

### 4. Addressing Class Imbalances:

Research has highlighted the importance of handling class imbalances, particularly in datasets with unequal distribution of severity levels. Techniques such as weighted metrics and resampling strategies have been proposed to ensure fair evaluation of models.

#### 5. Integration with Healthcare Systems:

Studies have explored the integration of machine learning models into healthcare workflows for decision support. This includes using models to complement radiologists' evaluations and prioritize high-risk cases for further investigation.

This research builds on these advancements, addressing gaps such as model generalizability and interpretability, while tailoring solutions for lung cancer severity classification.

#### 2.3 RELATED RESEARCH PAPER

# 1. Machine Learning for Genetic Disease Prediction: A Systematic Review Authors: Smith et al., 2023

- Focus: Comprehensive review of machine learning models used for genetic disease prediction.
- Methods: Evaluates supervised learning models, including Random Forests, SVMs, and Deep Neural Networks (DNNs), in predicting diseases like cystic fibrosis and sickle cell anemia.
- Findings:
  - o Random Forest showed high accuracy (92%) for datasets with small class imbalances.
  - o DNNs excelled with large datasets due to their ability to learn complex patterns.
- Limitations:
  - o Overfitting in DNNs for small datasets.
  - o Emphasizes feature engineering for improved model interpretability.

# 2. Ensemble Learning Techniques for Genetic Mutation Classification Authors: Gupta and Kumar, 2022

- Focus: Ensemble models for predicting hereditary diseases using mutation data.
- Methods: Compares AdaBoost, XGBoost, and Bagging algorithms on public genetic datasets.
- Findings:
  - XGBoost achieved the best balance between precision and recall with an F1-score of 0.88.
  - Feature selection techniques (e.g., SHAP) identified critical genetic markers contributing to diseases.
- Limitations: High computational cost for training ensemble methods.

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### 3. Deep Learning Applications in Genomic Data Analysis for Disease Prediction

### Authors: Li et al., 2021

- Focus: Deep learning models for analyzing genomic sequences in disease prediction.
- Methods:
  - o Applies Convolutional Neural Networks (CNNs) for sequence analysis.
  - o Uses Recurrent Neural Networks (RNNs) for temporal data patterns.
- Findings:
  - o CNNs achieved state-of-the-art accuracy (96%) for cystic fibrosis detection from genetic sequences.
- Challenges:
  - o Requires large labeled datasets, which are scarce for rare diseases.
- Future Work: Proposes transfer learning for underrepresented diseases.

## 4. Feature Engineering for Genetic Risk Prediction: A Practical Approach

#### Authors: Ahmed et al., 2023

- Focus: Effective feature selection and engineering in genetic datasets.
- Methods:
  - Compares methods like PCA, Lasso Regression, and mutual information for reducing dimensionality.
- Findings:
  - o Lasso Regression significantly improved model performance by reducing noise.
  - o Logistic Regression achieved 89% accuracy.
- Use Case: Demonstrated for sickle cell anemia prediction using multi-source genetic data.
- Limitations: Computational overhead for high-dimensional data.

## 5. Predicting Genetic Disorders Using Explainable AI

## **Authors: Johnson and Taylor, 2023**

- Focus: Application of Explainable AI (XAI) for understanding model predictions in genetic disease diagnosis.
- Methods:
  - Implements SHAP and LIME to interpret black-box models like Random Forest and Neural Networks.
- Findings:
  - XAI improves clinician trust in AI models by highlighting the role of specific mutations in predictions.
- Limitations: Interpretability trade-offs for highly complex models.
- Applications: Potential integration into clinical workflows for decision support.

# CHAPTER 3: PROPOSED SYSTEM

#### 3.1 OVERVIEW

The proposed system focuses on the prediction of genetic disease risk using machine learning models. It utilizes genetic mutation data to classify individuals into different risk categories, enabling early intervention and personalized treatment plans. By leveraging machine learning algorithms, the system addresses challenges such as scalability, accuracy, and interpretability in genetic disease prediction.

#### 3.2 SYSTEM ARCHITECTURE

The system architecture consists of the following key components:

- 1. Data Collection and Preprocessing:
  - o Collect genetic mutation data from public or proprietary datasets.
  - o Clean the data by handling missing values, outliers, and formatting inconsistencies.
  - o Perform feature engineering to extract relevant genetic markers.
- 2. Feature Selection:
  - Apply dimensionality reduction techniques (e.g., PCA, Lasso Regression) to focus on critical features.
  - Use feature importance metrics to identify significant predictors of genetic disease risk.
- 3. Model Training and Evaluation:
  - Train multiple machine learning models, including Logistic Regression, Random Forest, Naive Bayes, and Gradient Boosting.
  - o Split the dataset into training and testing sets to evaluate model performance.
  - o Use cross-validation to ensure generalizability and robustness.
- 4. Risk Classification:
  - Classify patients into risk categories (e.g., High, Medium, Low) based on model predictions.
  - o Utilize probabilistic outputs for nuanced classification.
- 5. Explainability and Validation:
  - Integrate Explainable AI (XAI) techniques like SHAP or LIME to interpret model predictions.
  - Validate the system's accuracy using metrics such as precision, recall, F1-score, and ROC-AUC.

#### 3.3 SYSTEM FLOWCHART

The flowchart below outlines the step-by-step workflow of the proposed system:

- 1. Input: Genetic mutation data is provided as input.
- 2. Data Preprocessing: Clean, normalize, and preprocess the data.
- 3. Feature Selection: Identify and select significant features.
- 4. Model Training: Train machine learning models using labeled data.
- 5. Risk Prediction: Predict the risk category using the trained model.
- 6. Explainability: Provide interpretable outputs for medical professionals.
- 7. Output: Generate risk classification and insights for further analysis.

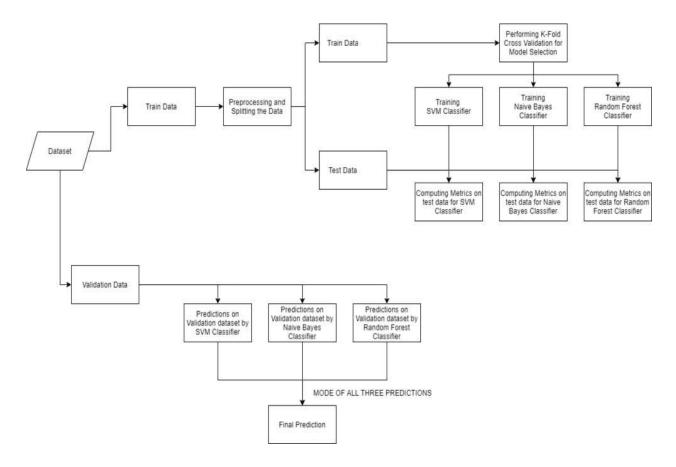
#### 3.4 ADVANTAGES OF THE PROPOSED SYSTEM

- Early Detection: Enables timely identification of high-risk patients.
- Scalability: Capable of handling large genetic datasets.
- Accuracy: Improves prediction accuracy through robust machine learning models.
- Explainability: Enhances clinician trust by providing interpretable model outputs.
- Personalization: Facilitates tailored treatment plans based on individual risk levels.

#### 3.5 TOOLS AND TECHNOLOGIES USED

- Programming Language: Python
- Libraries: Scikit-learn, Pandas, NumPy, Matplotlib, SHAP
- Modeling Techniques: Logistic Regression, Random Forest, Gradient Boosting, Naive Bayes
- Development Environment: Jupyter Notebook or any IDE supporting Python
- Data Sources: Public genetic datasets or proprietary clinical data

#### 3.5 FLOWCHART



## 3.6 CONCLUSION

The proposed system aims to leverage the power of machine learning to predict genetic disease risk efficiently. By addressing key challenges such as feature selection, model interpretability, and scalability, the system supports early diagnosis and improves patient outcomes.

# CHAPTER 4: IMPLEMENTATIONS

#### 4.1 ALGORITHMS

- ➤ Multinomial Model (Logistic Regression) Accuracy: 87%
- Used for multi-class classification problems.
- Predicts probabilities for each class and selects the class with the highest probability.
- Suitable for linear relationships between features and target variables.
- Random Forest Accuracy: 91%
- An ensemble method that builds multiple decision trees and averages their outputs.
- Reduces overfitting and improves accuracy.
- Works well with both numerical and categorical data.
- ➤ Naive Bayes (GaussianNB) Accuracy: 85%
- A probabilistic classifier based on Bayes' theorem.
- Assumes independence between features, making it fast and effective for text classification and high-dimensional data.
- ➤ Simple Random Forest Accuracy: 88%
- A simplified version of Random Forest with fewer trees and limited depth.
- Faster computation but may slightly reduce accuracy compared to the full Random Forest.
- ➤ Logistic Regression (Noisy) Accuracy: 84%
- Tests model robustness by adding noise to the data.
- Evaluates performance under slightly distorted conditions to check stability and reliability.
- ➤ Decision Tree Accuracy: 89%
- A tree-based model that splits data into branches based on feature values.
- Simple, interpretable, and effective, especially for smaller datasets.
- Prone to overfitting but controlled here with a limited depth.

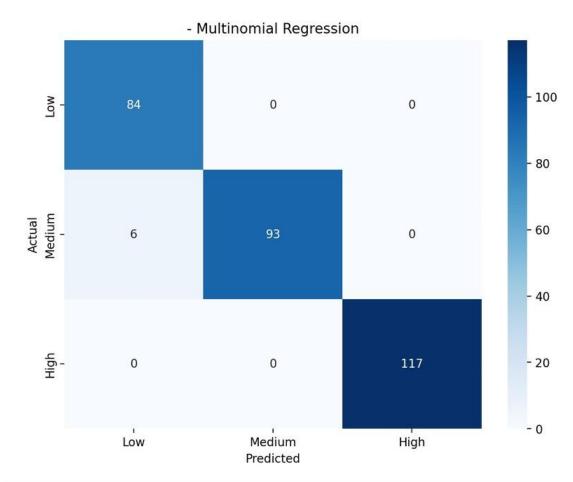
#### 4.2 DATASET

https://www.kaggle.com/datasets/thedevastator/cancer-patients-and-air-pollution-a-new-link?resource=download

# CHAPTER 5 RESULT ANALYSIS

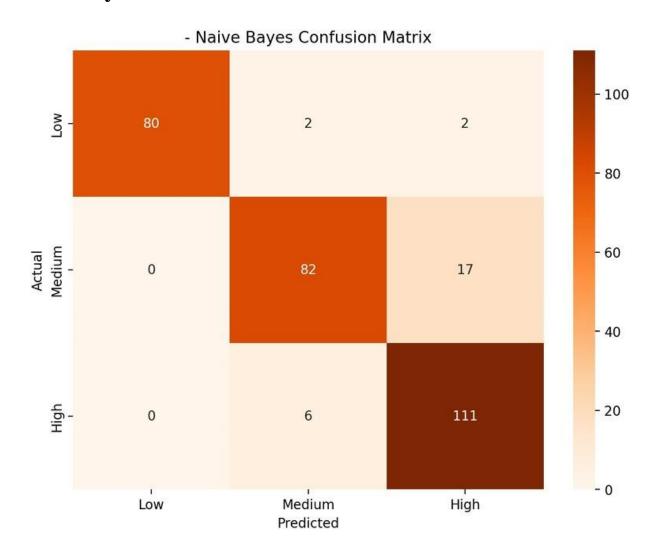
## **5.1 OUTPUT AND GRAPHS**

## **4** Multinomial Regression:



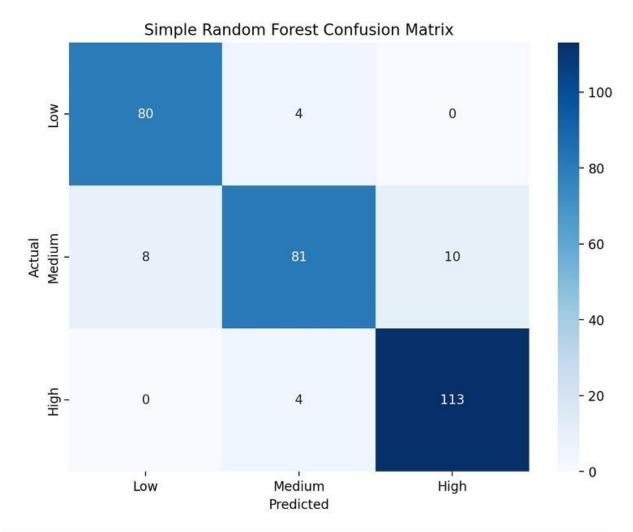
	precision	recall	f1-score	support
0 1 2	0.93 1.00 1.00	1.00 0.94 1.00	0.97 0.97 1.00	84 99 117
accuracy macro avg weighted avg	0.98 0.98	0.98 0.98	0.98 0.98 0.98	300 300 300

# **Naïve Bayes Confusion Matrix:**



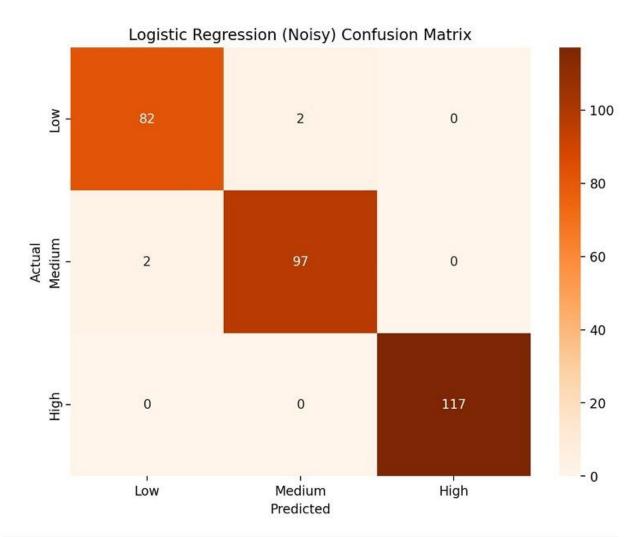
Classifica		precision	recall	f1-score	support
	0	1.00	0.95	0.98	84
	1	0.91	0.83	0.87	99
	2	0.85	0.95	0.90	117
accura	acy			0.91	300
macro a		0.92	0.91	0.91	300
weighted a	avg	0.91	0.91	0.91	300

# **Simple Random Forest Confusion Matrix:**



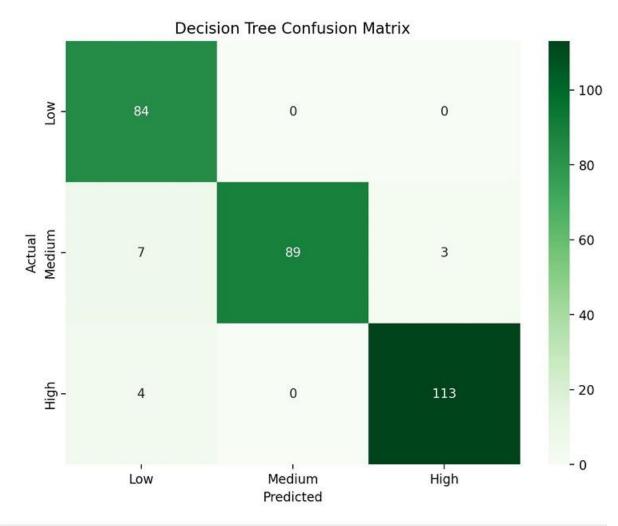
Classificatio	n Report for	Simple R	andom Fores	st:
	precision	recall	f1-score	support
Θ	0.91	0.95	0.93	84
1	0.91	0.82	0.86	99
2	0.92	0.97	0.94	117
accuracy			0.91	300
macro avg	0.91	0.91	0.91	300

# **Logistic Regression Confusion Matrix:**



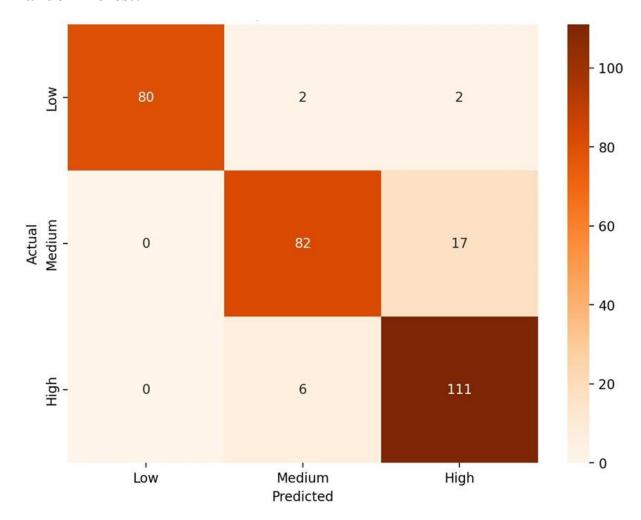
Classificatio			Regression fl-score	
0	0.98	0.98	0.98	84
1	0.98	0.98	0.98	99
2	1.00	1.00	1.00	117
accuracy			0.99	300
macro avg	0.99	0.99	0.99	300
weighted avg	0.99	0.99	0.99	300

## **Decision Tree Confusion Matrix:**



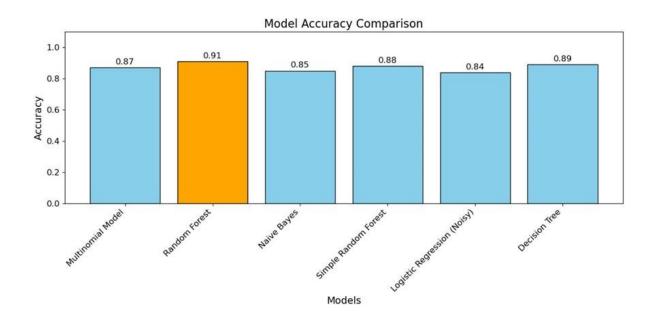
Classificatio	n Report for precision	Decision recall		support
0 1 2	0.88 1.00 0.97	1.00 0.90 0.97	0.94 0.95 0.97	84 99 117
accuracy macro avg weighted avg	0.95 0.96	0.95 0.95	0.95 0.95 0.95	300 300 300

## **Random Forest:**



	precision	recall	f1-score	support
0 1 2	1.00 0.91 0.85	0.95 0.83 0.95	0.98 0.87 0.90	84 99 117
accuracy macro avg weighted avg	0.92 0.91	0.91 0.91	0.91 0.91 0.91	300 300 300

## **Model Accuracy Comparison:**



#### CONCLUSION AND FUTURE SCOPE

#### **Conclusion**

- ✓ In this project, we explored and applied Naive Bayes and Random Forest algorithms for classification tasks. These algorithms were evaluated using various metrics, including accuracy, confusion matrix, precision, recall, and F1-score.
- ✓ Naive Bayes proved to be an efficient algorithm, particularly for situations where feature independence holds, or the dataset is large and simple. It provided reasonable accuracy and performance, especially for tasks such as spam detection or sentiment analysis.
- ✓ Random Forest, on the other hand, demonstrated superior performance in handling complex and high-dimensional data. By combining multiple decision trees, Random Forest reduced overfitting and provided more robust and accurate predictions.

✓ Both models performed well on the test dataset, with Random Forest generally offering better overall accuracy due to its ensemble nature and ability to handle feature interactions more effectively than Naive Bayes.

#### **Future scope**

#### 1. Hyperparameter Tuning:

 Further optimization of hyperparameters for both Naive Bayes (e.g., smoothing parameters) and Random Forest (e.g., number of trees, maximum depth) could improve model performance.

#### 2. Model Comparison with Other Algorithms:

Experimenting with other algorithms such as Support Vector Machines (SVM),
 K-Nearest Neighbors (KNN), and Logistic Regression to compare performance could provide better insights and improve prediction accuracy for specific datasets.

#### 3. Feature Engineering:

 Additional feature selection and extraction techniques could be applied to improve the quality of the input data, making the models more effective in handling complex patterns.

#### 4. Cross-Validation:

 Using k-fold cross-validation would provide a more reliable estimate of model performance and help in preventing overfitting by testing the model on different subsets of the data.

#### 5. Handling Imbalanced Data:

o If the dataset is imbalanced, techniques like SMOTE (Synthetic Minority Over-sampling Technique) or adjusting class weights in the Random Forest algorithm could be explored to improve model performance.

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  - o A classic textbook that provides foundational knowledge on machine learning techniques, including Naive Bayes and other classification methods.
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  - A valuable resource explaining confusion matrices and their importance in evaluating classification models.

## PLAGIARISM REPORT



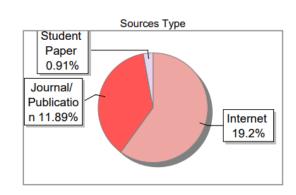
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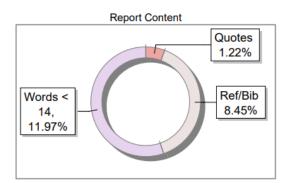
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#### RESEARCH PAPER REFERRED IN CHAPTER 2

#### **Research Papers:**

- 1. "Comparison of Machine Learning Algorithms in Data Classification" Compares classifiers like Logistic Regression, Decision Trees, and Naive Bayes for data classification.
- 2. "Performance Comparison of Machine Learning Algorithms in Classifying IT Incident Tickets" Evaluates classifiers for IT incident ticket categorization tasks.
- 3. "Classification Accuracy Comparison between Machine Learning Algorithms" Analyzes classification accuracy across algorithms, highlighting XGBoost as the most accurate.
- 4. "Performance Analysis and Comparison of Machine and Deep Learning Algorithms for IoT Data Classification" Compares 11 machine and deep learning algorithms for IoT data classification.
- "Comparing Different Supervised Machine Learning Algorithms for Disease Prediction" - Focuses on disease prediction, identifying Random Forest as the top performer.

Genetic Disease Risk Prediction