

The Aria International

Abstract

The project aims to develop predictive and classification algorithms to forecast or classify diseases and mortality rates based on historical data. This data contains detailed information on mortality rates caused by various diseases over several years, categorized by countries, organizations, and regions. Additionally, the study will analyze trends and patterns in death rate changes over time for specific areas.

The primary objective is to identify critical diseases that require immediate attention and understand which diseases are contributing most significantly to child mortality. This analysis will provide actionable insights for organizations like the WHO and other health agencies, empowering them with valuable decision-making tools to look over those decease which are causing high death rate and decrease it.

The study employs a variety of machine learning algorithms, including:

For Regression Analysis: Linear Regression, K-Nearest Neighbors (KNN) Regressor, and Decision Tree Regressor.

For Classification Analysis: Decision Tree Classifier, K-Nearest Neighbors (KNN) Classifier, and Gradient Boosting Classifier.

By leveraging these models, the project aims to enhance predictive capabilities and support strategic interventions to mitigate child mortality on a global scale.

Table Of Content

| Abstract | I |
|--|----|
| Keywords | V |
| 1. Introduction | 1 |
| 1.1. Background | 1 |
| 1.1.1 The Importance of Health on a Global Scale. | 1 |
| 1.1.2 Human Rights and Equity: | 1 |
| 1.1.3 Impact on Families and Communities: | 1 |
| 1.1.4 Global Development Goals: | 1 |
| 1.1.5 Challenges and Opportunities in Addressing Under-5 Mortality | 1 |
| 1.2 The Relationship between AI and Health | 2 |
| 1.2.1 How Machine Learning Is Used in Healthcare | 3 |
| 1.3 What is the ML Process? | 3 |
| 1.4 Problem-Statement: | 3 |
| 1.5 Aims: | 4 |
| 1.6 Objectives: | 4 |
| 1.7 Scope and Structure of the Project and Report: | 5 |
| 2. Literature Review | 6 |
| 2.1. Related Work: | 6 |
| 2.2. Gaps in Existing Research Addressed by the Case Study | 7 |
| 3. Dataset Description | 8 |
| 3.1 Key Features: | 8 |
| 4. Methodology | 9 |
| 4.1 Data Collection: | 9 |
| 4.1.1. Exploratory Data Analysis (EDA): | 9 |
| 4.1.2. Data Preprocessing: | 9 |
| 4.1.3. Data Visualization: | 9 |
| 4.1.4. Machine Learning Models: | 10 |
| 4.2 Model Implementation: | 10 |
| 4.3 Model Evaluation: | 10 |
| 4.4 Tools and Libraries Used: | 10 |
| 5. Result and Analysis | 11 |
| 5.1 Four Case Studies: | 11 |
| 5.1.1 Regression Analysis: | 11 |

| 5.1.2 Classification Analysis: | 11 |
|---|----|
| 6. Discussion | 14 |
| 6.1. Significance of the Findings | 14 |
| 6.2. Limitations of the Study | |
| 6.3. Potential Improvements and Future Research Directions | |
| | |
| 7. Conclusion: | 16 |
| 7.1. The broader impact of the study in medical applications and research | 16 |
| 8. References: | 17 |
| 9. Appendix | 18 |
| 9.1. Codes | 18 |
| 9.2. Graphs | |
| Table Of Tables | |
| Table 2-1 : Literature Review | 7 |
| Table 9-1 : Models Accuracy Or RMSE For Case Studies | |
| Table Of Figures | |
| Figure 1.2-1 AI Transforming On Health Care | 2 |
| Figure 5.1.1-1: Taking User Input, Printing RMSE OF Each Model And Predicting Death Co. | |
| Figure 5.1.2-1: Taking User Input, Printing Accuracy Score And Classifying Top Disease | |
| Figure 5.1.2-2: Taking User Input, Printing User Input And Classifying Top Cause Of Death | |
| Disease | |
| Figure 5.1.2-3 Continue Output | |
| Figure 5.1.2-4 : Continue Output | |
| Figure 5.1.2-5 : Continue Output | |
| Figure 5.1.2-6: Taking User Input, Printing Accuracy And Classifying Top 5 Countries | |
| Figure 9-1 : Loading Dataset | |
| Figure 9-2: Pre-processing | 18 |
| Figure 9-3: Handling Missing Values | |
| Figure 9-4: Handling Categorical Data By Applying Label Encoder | |
| Figure 9-5 : Splitting and Scaling Features For Prediction Task | 19 |
| Figure 9-6: Evaluation Function- Training Models, Calculating RMSE Comparing RMSE An | |
| Selecting Best Model | |
| Figure 9-7: Predicting Death Count By Taking User Input | |
| Figure 9-8: Preprocess Function- Filtering Data According To Country And Year | |
| Figure 9-9: Evaluate Classifier Model Function By Accuracy Score | |
| Figure 9-10: Predicting Disease Name By Country And Year | |
| Figure 9-11 : Taking User Input For Prediction | |
| Figure 9-12 : Filtering Data By Year | |
| Figure 9-13: Predicting Top Cause Of Death By Disease In Year | |
| Figure 9-14: TAking Year As User Input | |
| Figure 9-15: Filtering Data By Year And Disease And Providing Top 5 Countries | |
| Figure 9-16: Printing The Table | |
| Figure 9.2-1 : Co-relation Matrix | |
| Figure 9.2-2: Malaria By YearFigure 9.2-3: Hiv/aids By Year | |
| Figure 9.2-4: Meningitis By YearFigure 9.2-5: Nutritional By Year | 27 |

| Figure 9.2-6: Other Neontal Disorders ByYearFigure 9.2-7: Whooping Cough By Year | 27 |
|---|-----------|
| Figure 9.2-8: LRI By YearFigure 9.2-9: Congenital Birth Defects By Year | 27 |
| Figure 9.2-10: Measies By YearFigure 9.2-11: Neonatal Sepsis And Infection By Year. | 28 |
| Figure 9.2-12: Neonatal Due To Birth And Trauma By YearFigure 9.2-13: Drowning I | 3y Year28 |
| Figure 9.2-14: Tuberculosis By YearFigure 9.2-15: Neonatal Pertem Birth By Year | 28 |
| Figure 9.2-16: Diarrheal By YearFigure 9.2-17: Neoplasms By Year | 29 |
| Figure 9.2-18 : Syphilis By Year | 29 |
| Figure 9.2-19: Graphical Representation-Taking User Input To Visualize Top 10 Entities | 29 |
| Figure 9.2-20: Output Of Figure 9.2-19 | 30 |
| Figure 9.2-21: Graphical Representation-Taking User Input To Visualize Top 5 Diseases | 30 |
| Figure 9.2-22: Pie Chart To Visualize The Output Of 9.2-21 | 31 |
| Figure 9.2-23: Visualizing Top 10 Most And Least Occurring Diseases In Specific Entity | 31 |
| Figure 9.2-24 Output Of Figure: 9.2-23 | 32 |
| Figure 9.2-25: Visualization By Using Plotly-Sort The Diseases By Total Death Count For | Specific |
| Entity | 32 |
| Figure 9.2-26 : Output Of 9.2-25 | 33 |
| Figure 9.2-27: By Using Plotly Graph - Taking Year And Diseases As User Input For Visua | |
| Figure 9.2-28: Output Of 9.2-27 Visualizing Top 10 Entities | 34 |

Keywords

- 1. Machine Learning
- 2. Random Forest
- 3. Regression
- 4. Classification
- 5. KNN
- 6. Prediction
- 7. Data Visualization
- 8. Diseases
- 9. Findings
- 10. Gradient Boosting Classifier.

1. Introduction

1.1. Background

Child mortality, especially among children under the age of 5, remains one of the most pressing global health challenges. Despite notable progress over the past decades, approximately 5 million children under 5 years of age died in 2020 alone. Many of these deaths were caused by conditions such as pneumonia, diarrhea, malaria, complications during childbirth, and neonatal disorders—conditions that could largely have been prevented or effectively treated with timely and accessible healthcare interventions.

1.1.1 The Importance of Health on a Global Scale.

The health and survival of children under the age of 5 are critical indicators of a nation's overall development and well-being. Despite significant advancements, under-5 mortality remains a pressing global issue, with millions of preventable deaths occurring each year. Ensuring good health for populations worldwide is not just a moral obligation but also a strategic priority that underpins the sustainability and growth of nations.

1.1.2 Human Rights and Equity:

Every child has the right to survive and thrive. Reducing under-5 mortality ensures equitable access to healthcare and opportunities for a better future.

1.1.3 Impact on Families and Communities:

The death of a child has devastating emotional and economic impacts on families and communities.

1.1.4 Global Development Goals:

Addressing under-5 mortality is essential for achieving Sustainable Development Goal 3 (Good Health and Well-being), which aims to end preventable deaths of newborns and children under 5 by 2030.

1.1.5 Challenges and Opportunities in Addressing Under-5 Mortality

Reducing under-5 mortality faces significant challenges, including inadequate healthcare access, socioeconomic inequalities, persistent infectious diseases, malnutrition, and geographic disparities, particularly in low- and middle-income countries. Climate change and limited data also exacerbate the issue. However, opportunities exist to address these challenges through technological advancements like predictive analytics, global initiatives such as UNICEF's immunization campaigns, and community-based healthcare models. Focused interventions in maternal care, nutrition, and sanitation, alongside policy reforms and increased funding, can

significantly reduce child mortality. By leveraging innovation, education, and collaboration, we can save millions of lives and improve global child health outcomes.

1.2 The Relationship between AI and Health

Artificial Intelligence (AI) is transforming healthcare by improving efficiency, accuracy, and accessibility in diagnosis, treatment, and prevention. AI-driven technologies, such as machine learning algorithms and natural language processing, enable healthcare professionals to analyze vast amounts of data, identify patterns, and make predictions that were previously unattainable. For example, AI models can predict death rate by diseases over years, disease outbreaks, personalize treatment plans based on patient data, and assist in early detection of illnesses like cancer..

Despite its benefits, challenges such as data privacy, algorithm bias, and the need for robust regulatory frameworks remain. Nonetheless, the synergy between AI and healthcare offers immense potential to address global health challenges, reduce disparities, and improve outcomes for patients worldwide.



Figure 1.2-1 AI Transforming On Health Care

1.2.1 How Machine Learning Is Used in Healthcare

Machine Learning (ML) is a powerful tool for addressing child mortality by identifying risk factors, improving early diagnosis, and enabling timely interventions. Here's how ML is applied to reduce under-5 deaths caused by diseases:

• Reducing Neonatal Mortality

Birth Risk Prediction: ML models identify pregnancies at risk of complications, such as preterm births or birth asphyxia, enabling early interventions.

Neonatal Monitoring: ML-powered devices monitor vital signs in newborns, detecting conditions like sepsis or respiratory distress early for timely treatment.

• Early Detection of Diseases

Predictive Analytics: ML models analyze health records, environmental factors, and genetic data to predict diseases such as pneumonia, malaria, and diarrhea, which are leading causes of under-5 deaths. Early detection allows for quicker treatment and better survival outcomes.

• Nutritional Support

ML algorithms identify patterns in malnutrition data and predict areas with food insecurity, enabling targeted nutritional interventions for at-risk children.

• Public Health Monitoring and Disease Forecasting

By analyzing environmental and epidemiological data, ML models predict disease outbreaks like malaria or cholera, allowing health systems to prepare and respond effectively.

1.3 What is the ML Process?

ML is a data driven approach to discovering patterns in the data that can be exploited for making predictions. Data collection, cleansing and preparation for analysis are the initial steps in this process. In addition, the data is treated with various machine learning algorithms in order to determine patterns. In order to make predictions, the results of that analysis shall be used.

1.4 Problem-Statement:

Child mortality under age 5 remains a critical global health issue, with millions of preventable deaths annually due to diseases like pneumonia, malaria, and diarrhea etc. Despite abundant historical data, there is a lack of advanced tools to forecast mortality trends and prioritize high-risk diseases effectively.

Organizations like WHO struggle to derive actionable insights from complex datasets to guide interventions. This project aims to address these challenges by developing machine learning models for predicting and classifying mortality rates.

1.5 Aims:

- Develop predictive and classification models for forecasting mortality rates in children under age 5.
- Identify and prioritize high-risk diseases contributing to child mortality, such as pneumonia, malaria, and diarrhea.
- Provide actionable insights to organizations like the WHO for more informed decisionmaking.
- Optimize resource allocation by identifying high-risk regions and critical diseases for targeted interventions.
- Analyze trends and patterns in mortality rates over time, categorized by countries, regions, and organizations.

1.6 Objectives:

- Collect and preprocess historical data on mortality rates, diseases, and socio-economic factors.
- Implement regression analysis models (e.g., Linear Regression, KNN Regressor, Decision Tree Regressor) to predict mortality rates.
- Implement classification models (e.g., Decision Tree Classifier, KNN Classifier, Gradient Boosting Classifier) to classify regions or countries based on mortality risk levels.
- Evaluate the models using appropriate metrics (e.g., MSE for regression, Accuracy for classification).
- Conduct trend analysis to identify significant changes in mortality rates over time.
- Visualize results through heatmaps and other interactive tools to provide clear insights into high-risk areas.

1.7 Scope and Structure of the Project and Report:

This report covers the implementation of machine learning algorithms in child mortality analysis, focusing on datasets that combine numerical and categorical variables. It includes a detailed description of the ML process, from data collection to model deployment. The structure of the report is as follows:

- **Introduction:** Overview of child mortality and the role of machine learning in healthcare.
- **Machine Learning Process**: Explanation of data collection, preprocessing, feature engineering, model selection, and evaluation.
- Case Study Application: Implementation of machine learning techniques to child mortality detection and prediction.
- **Results and Evaluation:** Evaluation of model performance and insights drawn from the data.
- Conclusion: Despite the challenges, the project aims to provide a reliable tool for healthcare professionals to predict and personalize child mortality different factors like cause of death, most important factor influencing death rate, yearly or country wise death rate which can improve early detection and treatment.

2. Literature Review

2.1. Related Work:

[1]. Author: Susan Idicula-Thomas & Ulka Gawde Year: 2021. Machine learning (ML) algorithms have been successfully employed for prediction of outcomes in clinical research.

[2] .Author: Lily Tapak, MSc1, Hossein Mahjub, PhD2, Omid Hamidi, MSc3, Jalal Poorolajal, PhD2 Year: 2013. This study compared two traditional classification methods (logistic regression and Fisher linear discriminant analysis) and four machine-learning classifiers (neural networks, support vector machines, fuzzy c-mean, and random forests) to classify persons with and without diabetes.

[3]. Author: Harimoorthy, K (Harimoorthy, Karthikeyan); Thangavelu, M (Thangavelu, Menakadevi)Year: 2021. Data in the healthcare industry consists of patient information and disease – related information. This medical data and machine learning techniques will help us to analyse a large amount of data to find out the hidden patterns in the disease.

| Research Paper | Health Analysis | Variables/ Features | Methodology | Accuracy |
|---|---|---|-------------------------------------|---|
| Comparison of machine learning algorithms applied to symptoms to determine infectious causes of death in children | SVM, ANN,CART, gradient boosting modeling, and KNN, c5 | Cause of death, disease code, ICD- 10 codes, Number of cases, %cases | Machine Learning | Based on disease SVM 80%-98%. ANN, KNN, CART, GBM 70%-90% C5 80%-90% |
| Real-Data Comparison of Data Mining Methods in Prediction of Diabetes in Iran | Fuzzy-C-mean, SVM, Random Forest, Logistic Regression, Linear Discriminant Analysis, | Sex,Smoking, Job, Hypertension, Residential, Physical Activity | Data Mining And Machine Learning | Lr = 0.93 Lda= 0.92 Fcm = 0.85 SVM = 0.98 NN = 0.93% RF = 0.93 |

| | Neural Network. | | | |
|--|---|--|--------------------|--|
| Multi-disease prediction model using improved SVM-radial bias | SVM-Radial bias kernel, SVM-Linear, SVM- Polynomial, Random forest and Decision tree | Datasets: Chronic kidney disease, Diabetes | Big Data Analytics | 98.3%, 98.7%, and 89.9% in Chronic Kidney Disease, Diabetes |

Table 2-1: Literature Review

2.2. Gaps in Existing Research Addressed by the Case Study

• Limited Adoption of Advanced Machine Learning Techniques

The studies reviewed, including [1], [2], and [3], highlight the use of traditional statistical and predictive models in survival prediction. However, they do not extensively explore the integration of modern machine learning techniques such as Random Forest or XGBoost for improving predictive accuracy. The case study fills this gap by leveraging advanced machine learning models and comparing their performance to identify the most effective approach for survival status prediction.

Challenges in Handling Class Imbalance

A recurring challenge in the studies [1] and [3] is the imbalance in survival datasets, particularly with fewer instances of advanced-stage cases. The case study addresses this gap by applying preprocessing techniques that mitigate the impact of imbalanced classes, ensuring that the predictive models are robust and unbiased.

Absence of Real-World Applicability

Existing research, such as in [2], focuses on theoretical or experimental validations without clear pathways for real-world implementation. This case study emphasizes actionable insights, demonstrating how the model's outputs can inform healthcare organization, such as early identification of high-risk disease which can cause high risk death and which specific area/country to be first prioritize for health aids.

• Limited Use of Comprehensive Evaluation Metrics

While [1] and [2] primarily assess model performance using accuracy or basic statistical measures, they do not evaluate using comprehensive metrics like R² and Accuracy score. The case study addresses this gap by adopting a multifaceted evaluation strategy, enabling a more thorough understanding of model performance.

3. Dataset Description

The dataset appears to track child mortality rates for various diseases under the age of 5, including causes such as malaria, HIV/AIDS, meningitis, pneumonia, and more. Each record includes the following columns:

- Entity: The country or region or organizations (e.g., Afghanistan).
- · Code: A unique identifier for the country or region (e.g., AFG for Afghanistan).
- · Year: The year the data was recorded (e.g., 1990, 2019).
- · These 17 columns representing the number of deaths due to various diseases
 - Malaria
 - HIV/AIDS.
 - Meningitis.
 - Nutritional Deficiencies.
 - Other Neonatal Disorders
 - Other diseases
 - Whooping Cough
 - Lower Respiratory Infections
 - Congenital birth defects
 - Measles
 - Neonatal sepsis and other neonatal infections
 - Neonatal encephalopathy due to birth asphyxia and trauma
 - Drowning
 - Tuberculosis
 - Neonatal preterm birth
 - Diarrheal Diseases.
 - Neoplasms.
 - Syphilis.

3.1 Key Features:

- 1. **Deaths per Disease**: The number of deaths for each disease under 5 years of age is recorded separately.
- 2. **Time-Series Data:** The data is available over multiple years, allowing for trend analysis and the identification of changes over time.
- 3. **Geographical Coverage**: The dataset includes information for different countries, which will enable regional comparisons

4. Methodology

Following is a step-by-step process for building a Predicting and Classifying Child Mortality Rates model.

4.1 Data Collection:

Data Sources: The dataset "cause-of-death-in-children is downloaded from kaggle.

Data Description: Entity (Countires or Organizations), Year (1990-2019), Code (Country code),

Disease Names 17 columns: Malaria, HIV/AIDS, Meningitis, Nutritional deficiencies, Other neonatal disorders, Whooping cough, Lower respiratory infections, Congenital birth defects, Measles, Neonatal sepsis and other neonatal infections, Neonatal encephalopathy due to birth asphyxia and trauma, Drowning, Tuberculosis, Neonatal preterm birth, Diarrheal diseases, Neoplasms, Syphilis

Data Availability: The "Code" column contains missing values because some regions or organizations may not have a country abbreviation

4.1.1. Exploratory Data Analysis (EDA):

Regular Expression: Applied regex to remove specific part from column.

Descriptive Statistics: Calculate mean, median, standard deviation, and correlation among the features.

4.1.2. Data Preprocessing:

Data Cleaning: Handling missing data by imputation (fillna= 'Unknown')

Data Transformation: Encoding categorical columns Entity and Code by applying LabelEncoder.

4.1.3. Data Visualization:

- Plotting trends of mortality rates by year, disease, or region.
- Visualizing correlations between diseases, years, country/area and mortality rates.
- Using plotly, bar charts, and scatter plots to understand relationships.

4.1.4. Machine Learning Models:

Model Selection:

Regression Models (for predicting mortality rates):

- 1. Linear Regression
- 2. K-Nearest Neighbors (KNN) Regressor.
- 3. Decision Tree Regressor.

Classification Models (for classifying regions or countries into high-risk categories based on mortality rates, Year, Diseases, Country/area):

- 1. Decision Tree Classifier.
- 2. K-Nearest Neighbors (KNN) Classifier.
- 3. Gradient Boosting Classifier.

4.2 Model Implementation:

- Scaling numerical data for machine learning models that require scaling (e.g., normalization or standardization.
- Training the models using the training dataset.

4.3 Model Evaluation:

- Regression Model Evaluation: Using performance metrics such as RMSE.
- Classification Model Evaluation: Using metrics such as Accuracy.
- Cross-validation to ensure model robustness.
- **Comparing Models:** Comparing the performance of different models to select the best one based on the evaluation metrics.

4.4 Tools and Libraries Used:

- Programming Languages: Python.
- Data Analysis Libraries: Pandas, NumPy.
- Data Visualization: Matplotlib, Seaborn, Plotly.
- Table: Tabulate
- Machine Learning Libraries: Scikit-learn

5. Result and Analysis

5.1 Four Case Studies:

5.1.1 Regression Analysis:

Case Study 1: Predicting Death Count: In this predictive analysis, the model takes the region, disease, and year as inputs and provides the predicted death count for specific disease in specific region/country as output.

Model Evaluation: By using RMSE evaluating model performance of Linear Regression, KNN Regressor and Decision Tree Regressor.

```
Enter the region (e.g., Africa, Asia, Europe): Afghanistan
Enter the disease (e.g., Malaria, HIV/AIDS, Tuberculosis): Malaria
Enter the year you want to predict: 2030
Model RMSE Scores:
KNN: 94.61, Linear Regression: 46.25, Decision Tree: 53.86
The best model is Linear Regression.
Predicted deaths for Malaria in Afghanistan in 2030: 111
```

Figure 5.1.1-1: Taking User Input, Printing RMSE OF Each Model And Predicting Death Count

5.1.2 Classification Analysis:

Case Study 2: Analysis By Entity And Year: In this classification analysis, the model takes country and year as input and provide top disease causing high death rate in specific region/country and in specific year.

Model Evaluation: By using Accuracy Score evaluating model performance of Random Forest Classifier, Gradient Boosting Classifier and Decision Tree Classifier.

Figure 5.1.2-1: Taking User Input, Printing Accuracy Score And Classifying Top Disease

Case Study 3: Analysis By Year: In this classification analysis, the model takes year as input and provide top cause of disease in specific year for every region/country.

Model Evaluation: By using Accuracy Score evaluating model performance of Random Forest Classifier, Gradient Boosting Classifier and Decision Tree Classifier.

| Enter the year: 2018 Random Forest Accuracy: 0.70 Gradient Boosting Accuracy: 0. Decision Tree Accuracy: 0.94 Best Model: Decision Tree with Top cause of death by disease | Accuracy: 0.94 | |
|---|--------------------------|-------------|
| Country | Disease | Death Count |
| Afghanistan | Congenital birth defects | 16521 |
| African Region (WHO) | Diarrheal diseases | 368837 |
| Albania | Congenital birth defects | 102 |
| Algeria | Neonatal preterm birth | 5914 |
| American Samoa | Neonatal preterm birth | 2 |

Figure 5.1.2-2: Taking User Input, Printing User Input And Classifying Top Cause Of Death By Disease

| Angola | Diarrheal diseases | 7700 |
|--|--|-------|
| Antigua and Barbuda Neonatal preterm birth | | 3 |
| Argentina | Congenital birth defects | 2101 |
| Armenia | Congenital birth defects | 135 |
| Australia | Congenital birth defects | 290 |
| Austria Congenital birth defects | | 91 |
| Azerbaijan | Lower respiratory infections | 1426 |
| Bahamas | Congenital birth defects | 7 |
| Bahrain | Congenital birth defects | 31 |
| Bangladesh | Neonatal encephalopathy due to birth asphyxia and trauma | 15700 |

Figure 5.1.2-3 Continue Output

| Barbados | Congenital birth defects | 7 |
|------------------------|--------------------------|------|
| Belarus | Congenital birth defects | 179 |
| Belgium | Congenital birth defects | 113 |
| Belize | Neonatal preterm birth | 26 |
| Benin | Malaria | 8224 |
| Bermuda | Malaria | 9 |
| Bhutan | Neonatal preterm birth | 90 |
| Bolivia | Neonatal preterm birth | 1701 |
| Bosnia and Herzegovina | Neonatal preterm birth | 51 |
| Botswana | Neonatal preterm birth | 296 |

Figure 5.1.2-4: Continue Output

| Western Pacific Region (WHO) | Congenital birth defects | 49549 |
|--------------------------------|------------------------------|--------|
| World | Lower respiratory infections | 709340 |
| World Bank High Income | Congenital birth defects | 15683 |
| World Bank Low Income | Lower respiratory infections | 219630 |
| World Bank Lower Middle Income | Lower respiratory infections | 444305 |
| World Bank Upper Middle Income | Congenital birth defects | 95665 |
| Yemen | Neonatal preterm birth | 10526 |
| Zambia | Diarrheal diseases | 4102 |
| Zimbabwe | Lower respiratory infections | 4675 |

Figure 5.1.2-5: Continue Output

Case Study 4: Analysis By Disease And Year: In this classification analysis, the model will take year as input and provide top 5 countries with highest occurrence of specific disease and specific year.

Model Evaluation: By using Accuracy Score evaluating model performance of Random Forest Classifier, Gradient Boosting Classifier and Decision Tree Classifier.

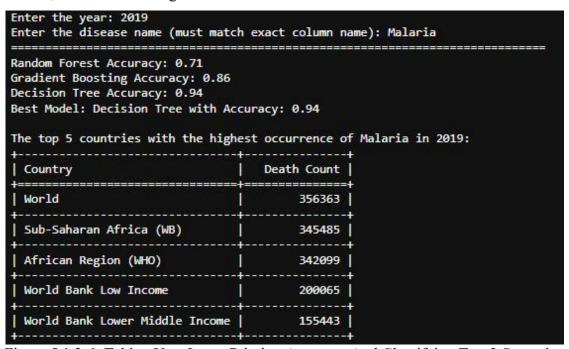


Figure 5.1.2-6: Taking User Input, Printing Accuracy And Classifying Top 5 Countries

6. Discussion

6.1. Significance of the Findings

The case study on child mortality rates due to various diseases provides significant insights into the global health challenges faced by under-5 children. The key findings of the study can be summarized as follows:

- Identifying Key Diseases: The study highlights the leading causes of child mortality, such as malaria, HIV/AIDS, pneumonia, and neonatal disorders, providing valuable information for public health efforts targeting high-mortality diseases.
- Trends Over Time: By analyzing mortality trends over multiple years, the study helps track changes in child mortality rates, showcasing the effectiveness of health interventions or identifying emerging challenges that need more focus.
- **Geographical Insights:** The dataset enables regional comparisons, showing how mortality rates vary across countries or regions. This information can guide policy makers and health organizations in targeting interventions more effectively in regions with higher mortality rates.
- **Predictive Modeling**: Through machine learning models, the study provides predictive insights, enabling the forecasting of mortality rates, which can be instrumental for resource allocation, health infrastructure planning, and early intervention.
- Classifying High-Risk Areas: The classification models help categorize regions or countries into high-risk categories based on mortality rates, helping prioritize regions that need urgent health interventions.

6.2. Limitations of the Study

While the study provides valuable insights, there are several limitations that must be considered:

- **Data Completeness**: The presence of missing values, especially in the "Code" column, can impact the reliability of the data, limiting the analysis to regions with available country codes. Some regions or organizations may not be fully represented, leading to biased conclusions.
- External Factors Not Included: The study focuses on mortality due to specific diseases but does not account for external factors like healthcare quality, socio-economic conditions, or vaccination rates, which could influence mortality rates.
- **Predictive Model Accuracy**: The accuracy of the predictive models, particularly for regression tasks such as predicting death counts, may not be perfect. Regression models like Linear Regression, Decision Tree Regressor, or KNN may be limited by their assumptions and the complexity of the data.
- **Limited to Available Data**: The study only covers data from 1990 to 2019, limiting its ability to capture very recent trends or the impacts of new health interventions.

6.3. Potential Improvements and Future Research Directions

- **Data Augmentation and Cleaning:** Further data collection and refinement could reduce missing values and improve data completeness. Imputation techniques like KNN imputation or using external datasets for filling missing data could be explored.
- Incorporating External Variables: Including socio-economic factors, healthcare infrastructure, vaccination rates, and other demographic details could lead to a more holistic analysis of child mortality rates.
- Advanced Machine Learning Models: While the current study uses basic models such as Decision Trees and KNN, exploring more advanced models like XGBoost, Random Forests, or deep learning techniques could improve the predictive power and accuracy of mortality predictions.
- Cross-National Comparisons: Future studies could compare mortality rates in countries with similar economic conditions, healthcare systems, and educational levels to identify effective interventions that could be adopted in other regions.
- **Longitudinal Analysis:** The study could extend beyond 2019 to understand the evolving nature of child mortality rates, particularly in light of ongoing global health crises like the COVID-19 pandemic or changes in healthcare practices.
- Classifying High-Risk Areas: Further refinement of the classification models using more granular data could improve the identification of high-risk regions, allowing for better-targeted interventions.
- **Real-Time Data Integration:** Incorporating real-time or near real-time data from health organizations could allow for more timely interventions and improvements in child mortality reduction strategies.

By addressing these limitations and exploring the outlined future directions, the study could provide more comprehensive and actionable insights for reducing child mortality rates globally.

7. Conclusion:

In conclusion, our project represents a significant step forward in leveraging technology to address the critical global issue of under-5 child mortality. By developing machine learning models for classification and prediction, we aim to provide actionable insights into the leading causes of death among children under the age of five. These insights will help identify high-risk regions, prioritize diseases requiring urgent attention, and guide effective resource allocation. We remain committed to refining our methodologies, improving data quality, and enhancing analytical techniques to ensure a meaningful impact in reducing child mortality rates and strengthening global health initiatives.

7.1. The broader impact of the study in medical applications and research

The broader impact of this study in medical applications and research is significant across various areas:

- **Improved Health Policy:** The study aids in identifying high-risk regions and diseases, helping policymakers allocate resources effectively and design targeted interventions.
- Enhanced Public Health Decision-Making: It supports data-driven decisions on disease prevention and management, optimizing public health strategies.
- Advancement in Epidemiology: The study provides insights into disease interactions, helping researchers understand factors contributing to child mortality and guiding policy development.
- **Global Health Interventions:** By identifying high mortality causes, it informs global strategies and supports efforts toward achieving SDG 3 (reducing child mortality).
- **Disease Prediction and Prevention:** The predictive models could lead to early warning systems for outbreaks and improved preventive healthcare strategies.
- Addressing Health Inequities: The study highlights disparities in child mortality, enabling further research into reducing healthcare access gaps and social determinants of health.
- **Pediatric Research:** It provides data for focused medical research on diseases like malaria, pneumonia, and neonatal infections, leading to better treatments and interventions.
- **Machine Learning in Healthcare:** The study demonstrates the potential of machine learning in predicting disease trends, which can be applied across various healthcare domains.

8. References:

- [1]. https://rdcu.be/d6DFS
- [2]. https://doi.org/10.4258/hir.2013.19.3.177
- [3]. https://www.webofscience.com/wos/woscc/full-record/WOS:000505384300008

9. Appendix

9.1. Codes

| lf | e_path = = pd.read head() | | | | sktop\cau | ises-of-de | ath-in-chi | ldren.csv | | | | | | |
|----|---------------------------------|------|------|--|---|---|---|---|---|---|--|--|--|---|
| | Entity | Code | Year | Deaths - Malaria - Sex: Both - Age: Under 5 (Number) | Deaths - HIV/AIDS - Sex: Both - Age: Under 5 (Number) | Deaths - Meningitis - Sex: Both - Age: Under 5 (Number) | Deaths - Nutritional deficiencies - Sex: Both - Age: Under 5 (Number) | Deaths - Other neonatal disorders - Sex: Both - Age: Under 5 (Number) | Deaths - Whooping cough - Sex: Both - Age: Under 5 (Number) | Deaths - Lower respiratory infections - Sex: Both - Age: Under 5 (Number) | Deaths - Congenital birth defects - Sex: Both - Age: Under 5 (Number) | Deaths - Measles - Sex: Both - Age: Under 5 (Number) | Deaths - Neonatal sepsis and other neonatal infections - Sex: Both - Age: Under 5 (Number) | Deaths - Neonatal encephalopathy due to birth asphyxia and trauma - Sex: Both - Age: Under 5 (Number) |
| 0 | Afghanistan | AFG | 1990 | 21 | 10 | 1709 | 1779 | 7112 | 2455 | 20224 | 12850 | 8649 | 420 | 1599 |
| 1 | Afghanistan | AFG | 1991 | 41 | 12 | 1743 | 1822 | 7574 | 2385 | 20879 | 13701 | 8669 | 520 | 1804 |
| 2 | Afghanistan | AFG | 1992 | 51 | 13 | 1954 | 2069 | 8614 | 2370 | 23585 | 15812 | 8539 | 662 | 2160 |
| 3 | Afghanistan | AFG | 1993 | 24 | 16 | 2252 | 2427 | 9458 | 2659 | 27116 | 17855 | 8949 | 723 | 241 |
| 4 | Afghanistan | AFG | 1994 | 52 | 19 | 2446 | 2649 | 9823 | 3187 | 29271 | 18835 | 10642 | 736 | 251 |

Figure 9-1: Loading Dataset

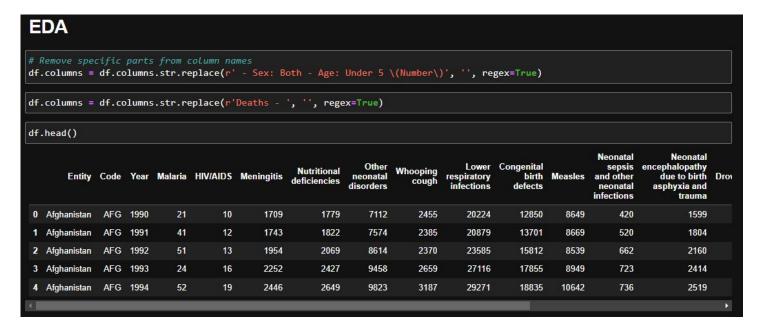


Figure 9-2: Pre-processing

```
print("\nHandling Missing Values...")
missing_values = df.isnull().sum()
print(missing_values[missing_values > 0])

Handling Missing Values...
Code 690
dtype: int64

df['Code'] = df['Code'].fillna('Unknown')

print(df.isnull().sum().sum())
```

Figure 9-3: Handling Missing Values

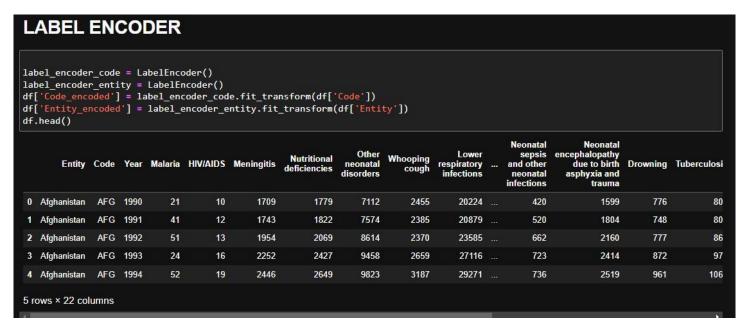


Figure 9-4: Handling Categorical Data By Applying Label Encoder

Case Study 1: Predicting Death Count:

```
def preprocess_data(df, region, disease):
    df_region = df[df['Entity'] == region]
    df_region = df_region[['Year', disease]].dropna() # Remove rows with missing values

X = df_region[['Year']] # Independent variable: Year
    y = df_region[disease] # Dependent variable: Disease

X_train, X_test, y_train, y_test = train_test_split(X, y, test_size=0.2, random_state=42)

# Scaling (Standardization)
    scaler = StandardScaler()
    X_train_scaled = scaler.fit_transform(X_train)
    X_test_scaled = scaler.transform(X_test)

    return X_train_scaled, X_test_scaled, y_train, y_test, scaler
```

Figure 9-5: Splitting and Scaling Features For Prediction Task

```
def evaluate_models(X_train_scaled, X_test_scaled, y_train, y_test):
   # Train KNN model
   knn_reg_model = KNeighborsRegressor(n_neighbors=5)
   knn_reg_model.fit(X_train_scaled, y_train)
   y_pred_knn = knn_reg_model.predict(X_test_scaled)
   rmse_knn = np.sqrt(mean_squared_error(y_test, y_pred_knn))
   # Train Linear Regression model
   lr model = LinearRegression()
   lr_model.fit(X_train_scaled, y_train)
   y_pred_lr = lr_model.predict(X_test_scaled)
   rmse_lr = np.sqrt(mean_squared_error(y_test, y_pred_lr))
   dt_reg_model = DecisionTreeRegressor(random_state=42)
   dt_reg_model.fit(X_train_scaled, y_train)
   y_pred_dt = dt_reg_model.predict(X_test_scaled)
   rmse_dt = np.sqrt(mean_squared_error(y_test, y_pred_dt))
   print(f"Model RMSE Scores:\nKNN: {rmse_knn:.2f}, Linear Regression: {rmse_lr:.2f}, Decision Tree: {rmse_dt:.2f}")
   if rmse_knn <= rmse_lr and rmse_knn <= rmse_dt:
       return knn_reg_model, "KNN Regres
   elif rmse_lr <= rmse_knn and rmse_lr <= rmse_dt:
       return lr_model, "Linear Regression
       return dt_reg_model, "Decision Tree Regressor"
```

Figure 9-6: Evaluation Function- Training Models, Calculating RMSE Comparing RMSE And Selecting Best Model

```
def predict_deaths(region, disease, year_to_predict, best_model, scaler):
   year_scaled = scaler.transform([[year_to_predict]]) # Scale the input year
   predicted deaths = best model.predict(year scaled)
   print(f"Predicted deaths for {disease} in {region} in {year_to_predict}: {predicted_deaths[0]:.0f}")
try:
   region = input("Enter the region (e.g., Africa, Asia, Europe): ")
   disease = input("Enter the disease (e.g., Malaria, HIV/AIDS, Tuberculosis): ")
   year_to_predict = int(input("Enter the year you want to predict: "))
   # Step 2: Preprocess the data
   X_train_scaled, X_test_scaled, y_train, y_test, scaler = preprocess_data(df, region, disease)
   # Step 3: Train models and evaluate RMSE
   best model, best model name = evaluate models(X train scaled, X test scaled, y train, y test)
   print(f"The best model is {best_model_name}.")
   # Step 4: Predict deaths using the best model
   predict deaths(region, disease, year to predict, best model, scaler)
except ValueError as e:
   print(f"Error: {e}")
```

Figure 9-7: Predicting Death Count By Taking User Input

Case Study 2: Classification Analysis By Country And Year:

```
def preprocess_top_diseases_data(df, country, year):
    # Filter data for the selected country and
   df_filtered = df[(df['Entity'] == country) & (df['Year'] == year)]
    if df_filtered.empty:
        raise ValueError(f"No data available for the selected country ({country}) and year ({year}).")
   disease_columns = [col for col in df.columns if col not in ['Entity', 'Code', 'Year']]
    if df_filtered[disease_columns].isnull().all().all():
        raise ValueError(f"No death count data for any diseases in {country} in {year}.")
   disease_counts = df_filtered[disease_columns].iloc[0]
   top_diseases = disease_counts.sort_values(ascending=False).head(5).index.tolist()
   df['High_Death_Disease'] = df[disease_columns].idxmax(axis=1) # Add top disease as Label
   label_encoder_disease = LabelEncoder()
df['High_Death_Disease'] = label_encoder_disease.fit_transform(df['High_Death_Disease'])
   label_encoder_country = LabelEncoder()
df['Entity_encoded'] = label_encoder_country.fit_transform(df['Entity'])
   # Select features and target
X = df[['Year', 'Entity_encoded']]
y = df['High_Death_Disease']
   # Split into train and test sets
   X_train, X_test, y_train, y_test = train_test_split(X, y, test_size=0.2, random_state=42)
   scaler = StandardScaler()
   X_train_scaled = scaler.fit_transform(X_train)
   X_test_scaled = scaler.transform(X_test)
   return X_train_scaled, X_test_scaled, y_train, y_test, top_diseases, scaler, label_encoder_disease, label_encoder_country
```

Figure 9-8: Preprocess Function-Filtering Data According To Country And Year

```
def evaluate_classifier_model(X_train_scaled, X_test_scaled, y_train, y_test):
    models = {
        "Random Forest": RandomForestClassifier(random_state=42),
        "Gradient Boosting": GradientBoostingClassifier(random_state=42),
        "Decision Tree": DecisionTreeClassifier(random_state=42)
}

accuracies = {}
for model_name, model in models.items():
    model.fit(X_train_scaled, y_train)
    y_pred = model.predict(X_test_scaled)
    accuracy = accuracy_score(y_test, y_pred)
    accuracies[model_name] = accuracy
    print(f"{model_name} Accuracy: {accuracy:.2f}")

# Select the best model
best_model_name = max(accuracies, key=accuracies.get)
best_model_s[best_model_name]
    print(f"Best Model: {best_model_name} with Accuracy: {accuracies[best_model_name]:.2f}")
    return best_model, best_model_name
```

Figure 9-9: Evaluate Classifier Model Function By Accuracy Score

```
def predict_top_diseases(country, year, classifier, scaler, label_encoder_disease, label_encoder_country, df):
   # Prepare the data for the specific country and year
   df_filtered = df[(df['Entity'] == country) & (df['Year'] == year)]
   if df_filtered.empty:
       print(f"No data available for {country} in {year}.")
       return
   # Extract the features
   X_predict = df_filtered[['Year', 'Entity_encoded']]
   # Ensure 'Entity_encoded' is available
   if 'Entity_encoded' not in X_predict.columns:
       X_predict['Entity_encoded'] = label_encoder_country.transform(X_predict['Entity']) # Transform if missing
   # Scale the features
   X_predict_scaled = scaler.transform(X_predict[['Year', 'Entity_encoded']]) # Only use relevant columns for s
   # Make predictions
   predicted diseases = classifier.predict(X predict scaled)
   predicted_disease_names = label_encoder_disease.inverse_transform(predicted_diseases)
   print("=====
   print(f"The top diseases causing high death rates in {country} in {year} are:")
    for disease in predicted_disease_names:
       print(disease)
```

Figure 9-10: Predicting Disease Name By Country And Year

```
try:
    # User Input
    country = input("Enter the country name: ")
    year = int(input("Enter the year: "))
    print("==========")

# Preprocess data
    X_train_scaled, X_test_scaled, y_train, y_test, top_diseases, scaler, label_encoder_disease, label_encoder_country = preproce
# Train models and select the best one
    best_model, best_model_name = evaluate_classifier_model(X_train_scaled, X_test_scaled, y_train, y_test)

# Predict top diseases
    predict_top_diseases(country, year, best_model, scaler, label_encoder_disease, label_encoder_country, df)

except ValueError as e:
    print(f"Error: {e}")
```

Figure 9-11: Taking User Input For Prediction

Case Study 3: Classification Analysis By Year:

```
Analysis By Year ¶

def preprocess top_diseases_data_by_year(df, year):
    # Filter data for the selected year
    df_filtered = df[df['Year'] == year]

if df_filtered.empty:
    raise ValueError(f"No data available for the year {year}.")

non_disease_columns = ['Entity', 'Code', 'Year', 'Entity_encoded', 'Code_encoded']
    disease_columns = [col for col in df.columns if col not in non_disease_columns]

# Check if the dataset contains any death counts for diseases
if df_filtered[disease_columns].isnull().all().all():
    raise ValueError(f"No death count data for any diseases in {year}.")

return df_filtered, disease_columns
```

Figure 9-12: Filtering Data By Year

```
def predict_top_diseases(year, df):
     # Preprocess the data for the selected year
df_filtered, disease_columns = preprocess_top_diseases_data_by_year(df, year)
     X = df[['Year', 'Entity_encoded', 'Code
y = df[disease_columns].idxmax(axis=1)
     # Encode categorical features
label_encoder_disease = LabelEncoder()
df['High_Death_Disease'] = label_encoder_disease.fit_transform(y)
     label_encoder_country = LabelEncoder()
     df['Entity_encoded'] = label_encoder_country.fit_transform(df['Entity'])
     label_encoder_code = LabelEncoder()
df['Code_encoded'] = label_encoder_code.fit_transform(df['Code'])
     X_train, X_test, y_train, y_test = train_test_split(X, y, test_size=0.2, random_state=42)
     scaler = StandardScaler()
X_train_scaled = scaler.fit_transform(X_train)
X_test_scaled = scaler.transform(X_test)
     best_model, best_model_name = evaluate_classifier_model(X_train_scaled, X_test_scaled, y_train, y_test)
     results = []
     for _, row in df_filtered.iterrows():
          disease_counts = pd.to_numeric(row[disease_columns], errors='coerce')
          high_cause_of_death = disease_counts.idxmax()
          death_count = disease_counts.max()
          if high_cause_of_death != 'High_Death_Disease':
                results.append([row['Entity'], high_cause_of_death, death_count])
     # Output the results as a table, ensuring that High_Death_Disease doesn't appear
print(f"Top cause of death by disease in {year}:")
print(tabulate(results, headers=["Country", "Disease", "Death Count"], tablefmt="grid"))
```

Figure 9-13: Predicting Top Cause Of Death By Disease In Year

```
# Example Usage
try:
    # User Input
    year = int(input("Enter the year: "))
    # Predict top diseases for the selected year
    predict_top_diseases(year, df)
except ValueError as e:
    print(f"Error: {e}")
```

Figure 9-14: TAking Year As User Input

Case Study 4: Classification Analysis By Disease:

```
Analysis By Disease
def preprocess_top_diseases_data1(df, year):
    df_filtered = df[df['Year'] == year]
    if df_filtered.empty:
       raise ValueError(f"No data available for the year {year}.")
    non_disease_columns = ['Entity', 'Code', 'Year', 'Entity_encoded', 'Code_encoded']
disease_columns = [col for col in df.columns if col not in non_disease_columns]
    if df_filtered[disease_columns].isnull().all().all():
       raise ValueError(f"No death count data for any diseases in {year}.")
    return df_filtered, disease_columns
def get_top_disease_countries(year, df, disease_name):
    # Preprocess the data for the selected year
    df_filtered, disease_columns = preprocess_top_diseases_data1(df, year)
    if disease_name not in disease_columns:
       raise ValueError(f"{disease_name} data is missing from the dataset.")
    df_sorted = df_filtered[['Entity', disease_name]].sort_values(by=disease_name, ascending=False)
    # Get the top 5 countries
    top_5_countries = df_sorted.head(5)
    # Output the results as a table using tabulate
    print(f"\nThe top 5 countries with the highest occurrence of {disease_name} in {year}:")
```

Figure 9-15: Filtering Data By Year And Disease And Providing Top 5 Countries

```
year = int(input("Enter the year: "))
disease_name = input("Enter the disease name (must match exact column name): ").strip()
    print("=====
    # Prepare features and target for training
    df_filtered, disease_columns = preprocess_top_diseases_data1(df, year)
    x = df[['Year', 'Entity_encoded', 'Code_encoded']] # Features: Year, Entity_encoded, Code_encoded
y = df[disease_columns].idxmax(axis=1) # Target: Disease with highest death count
    # Encode categorical features
label_encoder_disease = LabelEncoder()
df['High_Death_Disease'] = label_encoder_disease.fit_transform(y)
    label_encoder_country = LabelEncoder()
df['Entity_encoded'] = label_encoder_country.fit_transform(df['Entity'])
    label_encoder_code = LabelEncoder()
df['Code_encoded'] = label_encoder_code.fit_transform(df['Code'])
    X_train, X_test, y_train, y_test = train_test_split(X, y, test_size=0.2, random_state=42)
    scaler = StandardScaler()
    X_train_scaled = scaler.fit_transform(X_train)
X_test_scaled = scaler.transform(X_test)
    # Evaluate classifier models and determine the best one
    best_model, best_model_name = evaluate_classifier_model(X_train_scaled, X_test_scaled, y_train, y_test)
    get_top_disease_countries(year, df, disease_name)
except ValueError as e:
    print(f"Error: {e}")
```

Figure 9-16: Printing The Table

| Models | Case Study 1 (Regression Task) (RMSE) | Case Study 2 (Classification Task) (Accuracy Score) | Case Study 3 (Classification Task) (Accuracy Score) | Case Study 4 (Classification Task) (Accuracy Score) |
|----------------------|--|--|--|--|
| Random Forest | - | 0.27 | 0,72 | 0.68 |
| Classifier | | | | |
| Gradient | - | 0.84 | 0.86 | 0.87 |
| Boosting | | | | |
| Classifier | | | | |
| Decision Tree | - | 0.95 | 0.94 | 0.95 |
| Classifier | | | | |
| KNN Regressor | 94.61 | - | - | - |
| Linear | 46.25 | - | - | - |
| Regresson | | | | |
| Decision Tree | 53.86 | - | - | - |
| Regression | | | | |

Table 9-1: Models Accuracy Or RMSE For Case Studies

9.2. Graphs

HeatMap

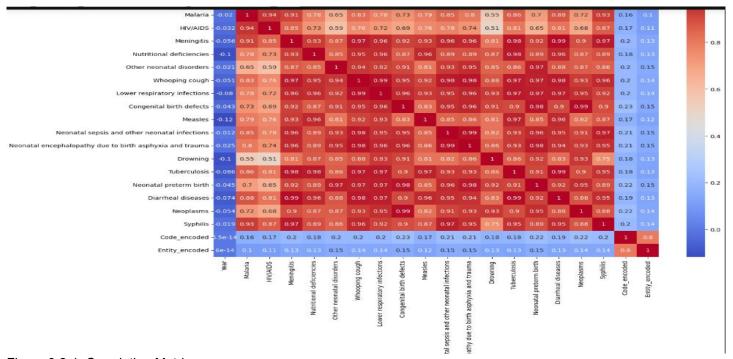


Figure 9.2-1: Co-relation Matrix

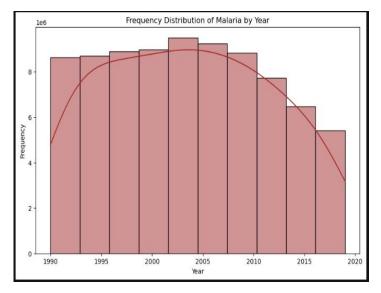


Figure 9.2-2: Malaria By Year

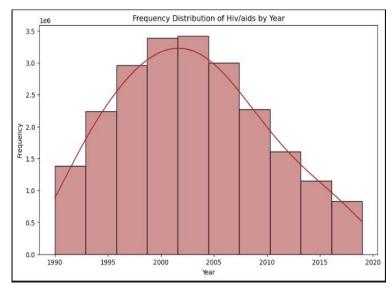


Figure 9.2-3: Hiv/aids By Year

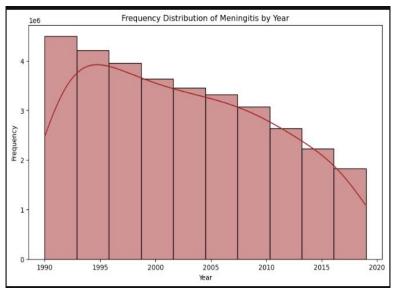


Figure 9.2-4: Meningitis By Year

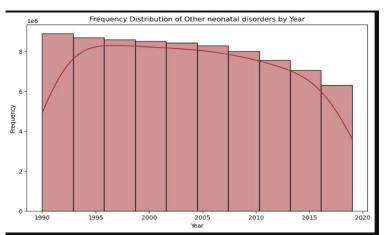


Figure 9.2-6: Other Neontal Disorders ByYear

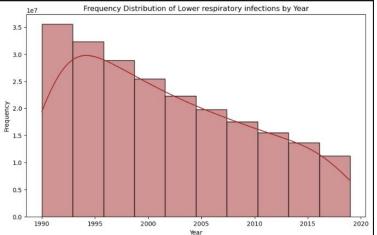


Figure 9.2-8: LRI By Year

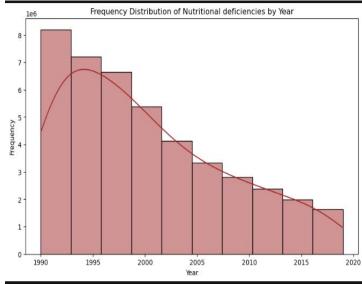


Figure 9.2-5: Nutritional By Year

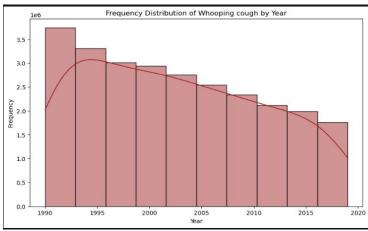


Figure 9.2-7: Whooping Cough By Year

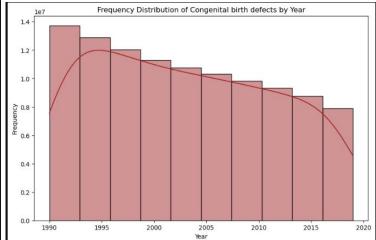


Figure 9.2-9: Congenital Birth Defects By Year

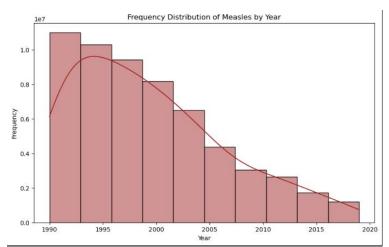


Figure 9.2-10: Measies By Year

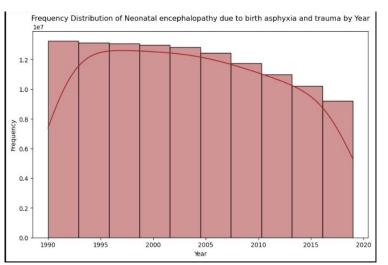


Figure 9.2-12: Neonatal Due To Birth And Trauma By Year

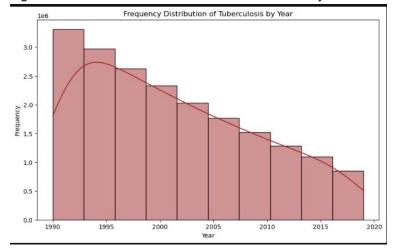


Figure 9.2-14: Tuberculosis By Year

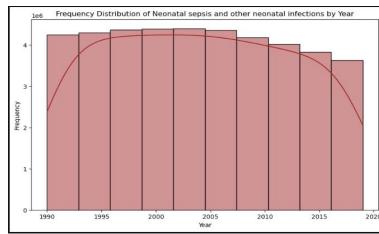


Figure 9.2-11: Neonatal Sepsis And Infection By Year

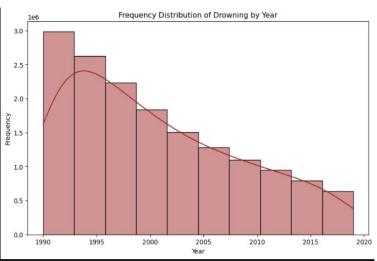


Figure 9.2-13: Drowning By Year

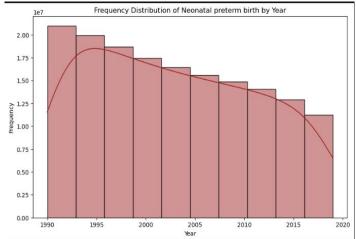
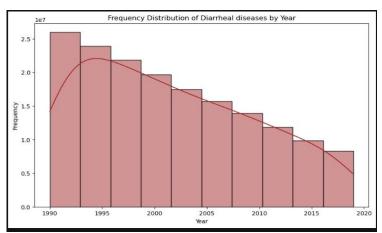


Figure 9.2-15: Neonatal Pertem Birth By Year



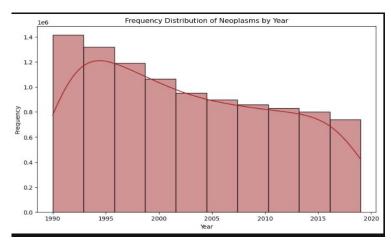


Figure 9.2-16: Diarrheal By Year

Figure 9.2-17: Neoplasms By Year

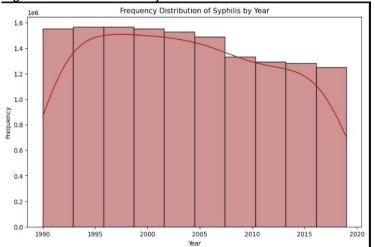


Figure 9.2-18: Syphilis By Year

```
disease_name = input("Enter the disease name (e.g., Malaria, HIV/AIDS, Tuberculosis): ")

if disease_name in df.columns:

    disease_counts = df[['Entity', disease_name]].groupby('Entity')[disease_name].sum().reset_index()
    top_10_disease = disease_counts.sort_values(by=disease_name, ascending=False).head(10)
    plt.figure(figsize=(12, 8))
    plt.bar(top_10_disease['Entity'], top_10_disease[disease_name], color='green')
    plt.vitle(f'Top_10_Entities with Highest Occurrence of {disease_name}')
    plt.ylabel('Entity (Country/Area/Organization)')
    plt.ylabel(f'{disease_name} Death Count')
    plt.xticks(rotation=45, ha='right')
    plt.tight_layout()
    plt.show()

else:
    print(f"Error: The disease '{disease_name}' does not exist in the dataset. Please check your input.")

Enter the disease name (e.g., Malaria, HIV/AIDS, Tuberculosis): Malaria
```

Figure 9.2-19: Graphical Representation-Taking User Input To Visualize Top 10 Entities

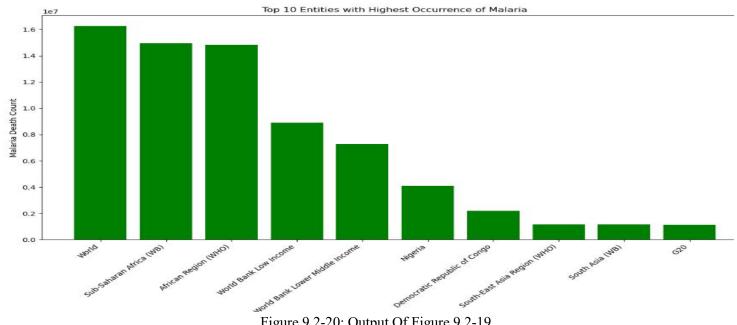


Figure 9.2-20: Output Of Figure 9.2-19

```
entity_name = input("Enter the country or region (e.g., India, Pakistan, Afghanistan): ")
entity_data = df[df['Entity'] == entity_name]
if not entity_data.empty:
    disease_columns = [col for col in df.columns if col not in ['Entity', 'Code', 'Year', 'Code_encoded', 'Entity_encoded']]
    disease_death_counts_entity = entity_data[disease_columns].sum()
    top_5_diseases_entity = disease_death_counts_entity.nlargest(5)
plt.figure(figsize=(8, 8))
    plt.pie(top_5_diseases_entity, labels=top_5_diseases_entity.index, autopct='%1.1f%%', startangle=140)
    plt.title(f'Proportion of Top 5 Diseases with Highest Death Count in {entity_name}')
plt.axis('equal')
    plt.show()
    print(f"Error: No data found for the entity '{entity_name}'. Please check your input.")
Enter the country or region (e.g., India, Pakistan, Afghanistan): Pakistan
```

Figure 9.2-21: Graphical Representation-Taking User Input To Visualize Top 5 Diseases

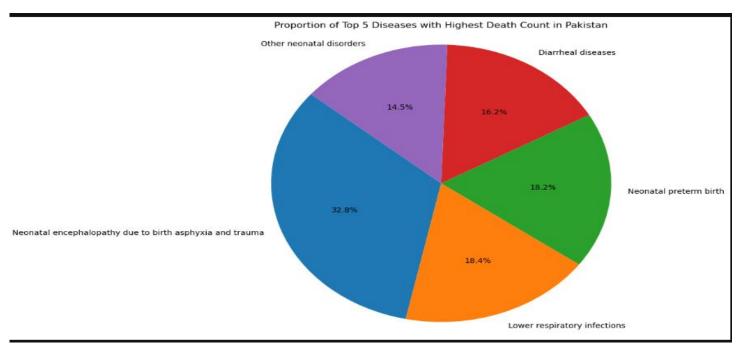


Figure 9.2-22: Pie Chart To Visualize The Output Of 9.2-21

```
entity_name = input("Enter the country or region (e.g., Algeria, India, Pakistan): ")
entity_data = df[df['Entity'] == entity_name]
if not entity_data.empty:
    disease_columns = [col for col in df.columns if col not in ['Entity', 'Code', 'Year', 'Code_encoded', 'Entity_encoded']]
    disease_death_counts_entity = entity_data[disease_columns].sum()
    top_10_max_diseases = disease_death_counts_entity.nlargest(10) top_10_min_diseases = disease_death_counts_entity.nsmallest(10)
    plt.figure(figsize=(14, 8))
    plt.subplot(1, 2, 1)
    sns.lineplot(x=top_10_max_diseases.index, y=top_10_max_diseases.values, marker='o', color='green')
    plt.title(f'Top 10 Most Occurring Diseases in {entity_name}')
plt.xlabel('Disease')
    plt.ylabel('Death Count')
    plt.xticks(rotation=90)
    plt.grid(True)
    plt.subplot(1, 2, 2)
    plt.title(f'Top 10 Least Occurring Diseases in {entity_name}')
plt.xlabel('Disease')
plt.ylabel('Death Count')
    sns.lineplot(x=top_10_min_diseases.index, y=top_10_min_diseases.values, marker='o', color='red')
    plt.xticks(rotation=90)
    plt.grid(True)
    # Adjust the layout and show the plot
plt.tight_layout()
    plt.show()
    print(f"Error: No data found for the entity '{entity_name}'. Please check your input.")
Enter the country or region (e.g., Algeria, India, Pakistan): Afghanistan
```

Figure 9.2-23: Visualizing Top 10 Most And Least Occurring Diseases In Specific Entity

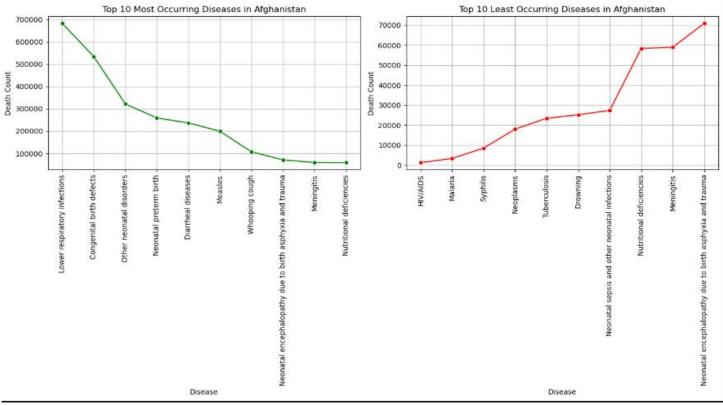


Figure 9.2-24 Output Of Figure: 9.2-23

Figure 9.2-25: Visualization By Using Plotly-Sort The Diseases By Total Death Count For Specific Entity

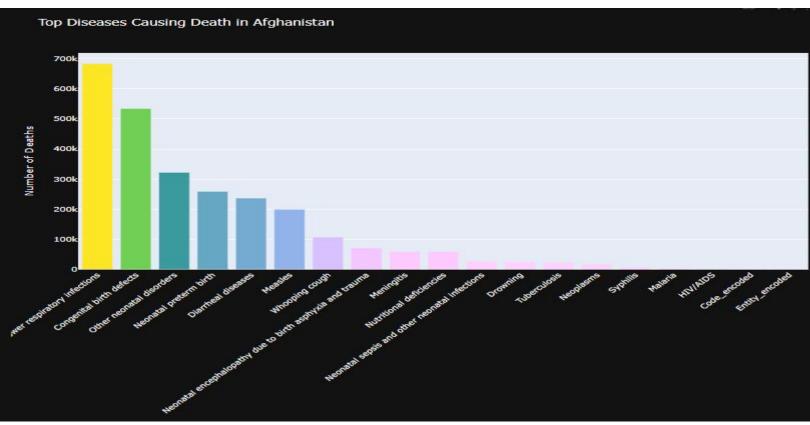


Figure 9.2-26: Output Of 9.2-25

Figure 9.2-27: By Using Plotly Graph - Taking Year And Diseases As User Input For Visualization

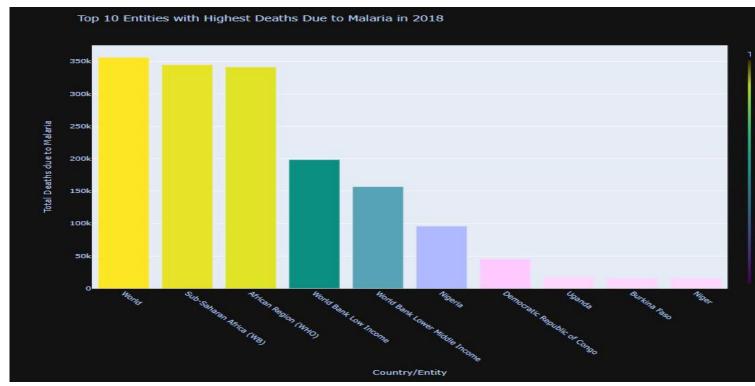


Figure 9.2-28: Output Of 9.2-27 Visualizing Top 10 Entities