RR_Project_Report

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1 REPRODUCIBLE RESEARCH PROJECT

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1.1 INTRODUCTION

Our project is centered on replicating and expanding upon the groundbreaking machine learning research detailed in the Asian Journal of Computer and Information Systems (AJCIS), specifically the article titled 'Utilisation of Machine Learning Techniques in Testing and Training of Different Medical Datasets'. This initiative seeks to deepen our understanding of how machine learning can revolutionize the analysis of medical data, thereby enhancing disease detection and improving diagnostic accuracy across various health conditions.

By engaging with this research, we aim to assess the reproducibility of the results and explore potential enhancements in the methodology. Our study will utilize a similar approach to analyze medical datasets, focusing on maximizing the potential of machine learning in healthcare. Through rigorous testing and training, our objective is to contribute meaningful advancements to the medical field, offering insights that could be vital in developing more effective diagnostic tools and treatments.

1.2 DATASETS

The study we replicated utilized machine learning algorithms to analyze six different healthcare datasets, all sourced from the **UC Irvine Machine Learning Repository**. Each of these datasets varies in the number of attributes and instances, as detailed below.

1.3 METHODOLOGY

1.3.1 DATA PREPROCESSING

For each dataset involved in our study, detailed data preprocessing steps are applied to ensure compatibility with machine learning models and to maintain the integrity of the data analysis. Below are the dataset-specific preprocessing methods implemented:

COVID-19 Dataset * **Binary Remapping:** Each categorical variable representing binary states (e.g., positive '+' or negative '-') is replaced with numerical values where '+' is mapped to 1 and '-' is mapped to 0. * **Numerical Conversion:** All columns are converted to numeric types to facilitate mathematical operations and model fitting. * **Column Renaming:** To clearly indicate

the binary nature of the data, columns are renamed by adding a 'is_' prefix, enhancing clarity and manageability in the dataset.

Hepatitis Dataset * Binary Remapping: Specific columns that contain categorical data related to the patient's conditions (e.g., 'Steroid', 'Antivirals', 'Fatigue') are converted to binary format, where applicable values are adjusted to 0 and 1 to reflect absence and presence of the condition. * Mean and Median Imputation: Missing values are addressed by calculating and substituting the mean and median for each column, allowing for robustness against outliers and skewed data. * Dropping Missing Data: Rows containing missing data are removed to avoid the introduction of bias or inaccuracies in the model's predictions. Corresponding entries in the target variable are also excluded to maintain dataset alignment.

Chronic Kidney Disease Dataset * Binary Mapping: Categorical descriptions (e.g., 'normal', 'abnormal'; 'present', 'notpresent') in symptoms and diagnostic results are mapped to binary (1, 0) to simplify the input for algorithm processing. * Mean and Median Imputation: Implements both mean and median imputation for handling missing values, providing two sets of data for comparative analysis and model robustness. * Dropping Missing Data: Ensures that only complete data entries are processed, by excluding rows with any missing values from the analysis, thus maintaining the reliability of the dataset.

Breast Cancer Dataset * Binary Mapping: The target variable 'Diagnosis' is converted from its original categorical labels ('M' for malignant, 'B' for benign) into a binary format (1 for malignant, 0 for benign), aligning it with binary classification tasks in machine learning. * Data Checks and Cleaning: Verifies data types, checks for outliers, and ensures that all entries are consistent and appropriate for analysis.

Immunotherapy Dataset * Binary Mapping and Renaming: Converts 'sex' from a categorical to a binary format and updates the column name to 'is_female' to reflect this binary distinction. * One-Hot Encoding: Applies one-hot encoding to the 'Type' column, which involves transforming it into multiple binary columns, each representing a different type of immunotherapy, thereby avoiding ordinal implications in the categorical data. * Concatenation: The one-hot encoded columns are then concatenated back to the main dataset, preserving the original data structure while adding new binary indicators.

Cryotherapy Dataset * Binary Mapping: Converts 'sex' to a binary format and adjusts column naming similarly to the Immunotherapy dataset to maintain consistency. * ne-Hot Encoding: The 'Type' column is treated with one-hot encoding to convert categorical treatment types into a series of binary columns, which are more suitable for machine learning models. * Concatenation of One-Hot Encoded Data: These binary columns are merged back into the primary dataset, ensuring that all data remains integrated and accessible for subsequent analysis.

1.3.2 MODELING AND HYERPARAMETER OPTIMIZATION

In aligning with the methods of the replicated study, we employ the same machine learning models, each optimized through a systematic hyperparameter tuning process. The models and their respective hyperparameters are as follows:

• Random Forest (RF): Optimized for n_estimators (50, 100, 200), criterion ('gini', 'entropy'), max_depth (None, 10, 20), min_samples_split (2, 5, 10), min_samples_leaf (1, 2, 4), max_features ('sqrt', 'log2').

- K-Nearest Neighbors (KNN): Optimized for n_neighbors (3, 5, 7), weights ('uniform', 'distance'), algorithm ('auto', 'ball_tree', 'kd_tree', 'brute'), p (1, 2).
- Support Vector Machine (SVM): Optimized for C (0.1, 1, 10), kernel ('poly', 'rbf'), degree (2, 3).
- Decision Tree (DT): Optimized for criterion ('gini', 'entropy'), splitter ('best', 'random'), max_depth (None, 10, 20), min_samples_split (2, 5, 10), min_samples_leaf (1, 2, 4), max_features ('sqrt', 'log2').

Hyperparameter tuning is conducted using **GridSearchCV**, which incorporates a **5-fold cross-validation** to ensure robustness and reliability in model performance. This meticulous optimization process ensures that each model is finely tuned to provide the best possible accuracy and generalizability on diverse medical datasets.

1.3.3 PERFORMANCE METRICS

In alignment with the replicated study from the Asian Journal of Computer and Information Systems, we utilize the same performance metrics to evaluate the effectiveness of the machine learning algorithms implemented in our research. Each model was applied to the designated healthcare datasets, and their performance was quantitatively assessed using the following metrics:

- Training Accuracy: This metric indicates the proportion of correct predictions made by the model on the training dataset. It provides insight into how well the model learns from the data, mirroring the approach taken in the replicated study.
- **Testing Accuracy:** Reflects the proportion of correct predictions on the testing dataset, crucial for understanding how well the model generalizes to new, unseen data.
- **Training Time:** Measures the duration of time the model takes to learn from the training data, highlighting the computational efficiency of the model during the training phase.
- **Testing Time:** The time required by the model to make predictions on the testing dataset, indicative of the model's operational efficiency.

Additionally, to further enhance our evaluation framework, we have incorporated the measurement of the **Training and Testing F1 Score**, metrics not used in the replicated study. The F1 Score is a harmonic mean of precision and recall, providing a more nuanced view of model performance, particularly in the context of imbalanced datasets:

- Training F1 Score: This metric helps assess the balance between precision and recall during the model's training phase, offering deeper insights into the model's predictive performance and accuracy.
- Testing F1 Score: Evaluates the model's precision and recall on the testing dataset, which is crucial for applications where the cost of false positives and false negatives is significant.

By utilizing these metrics, which were also employed in the original study, we ensure a consistent and comparative approach to evaluating model performance.

1.4 RESULTS

•	dataset		train_time	eval_time	\
0	breast_cancer	rf	0.183281	0.027984	
1	breast_cancer	knn	0.002406	0.018223	
2	breast_cancer	svm	0.029788	0.012569	
3	breast_cancer	dt	0.017596	0.005135	
4	chronic_kidney_disease_mean_imputted	rf	0.269601	0.044630	
5	chronic_kidney_disease_mean_imputted	knn	0.005406	0.024386	
6	chronic_kidney_disease_mean_imputted	svm	0.010864	0.006508	
7	chronic_kidney_disease_mean_imputted	dt	0.004596	0.001432	
8	<pre>chronic_kidney_disease_median_imputted</pre>	rf	0.191878	0.040303	
9	<pre>chronic_kidney_disease_median_imputted</pre>	knn	0.005036	0.017453	
10	<pre>chronic_kidney_disease_median_imputted</pre>	svm	0.010226	0.003573	
11	<pre>chronic_kidney_disease_median_imputted</pre>	dt	0.003194	0.001441	
12	<pre>chronic_kidney_disease_missing_dropped</pre>	rf	0.273432	0.035553	
13	<pre>chronic_kidney_disease_missing_dropped</pre>	knn	0.004863	0.015425	
14	<pre>chronic_kidney_disease_missing_dropped</pre>	svm	0.009367	0.004073	
15	<pre>chronic_kidney_disease_missing_dropped</pre>	dt	0.008460	0.003340	
16	covid_19	rf	0.194533	0.030823	
17	covid_19	knn	0.003052	0.031629	
18	covid_19	svm	0.007811	0.003878	
19	covid_19	dt	0.007249	0.006057	
20	cryotherapy	rf	0.219233	0.040224	
21	cryotherapy	knn	0.007742	0.034397	
22	cryotherapy	svm	0.007215	0.003353	
23	cryotherapy	dt	0.003139	0.002361	
24	hepatitis_mean_imputted	rf	0.237288	0.038766	
25	hepatitis_mean_imputted	knn	0.006000	0.018121	
26	hepatitis_mean_imputted	svm	0.012891	0.005145	
27	hepatitis_mean_imputted	dt	0.009227	0.006996	
28	hepatitis_median_imputted	rf	0.376774	0.041732	
29	hepatitis_median_imputted	knn	0.005754	0.022186	
30	hepatitis_median_imputted	svm	0.010769	0.006719	
31	hepatitis_median_imputted	dt	0.007189	0.003993	
32	hepatitis_missing_dropped	rf	0.208498	0.039103	
33	hepatitis_missing_dropped	knn	0.005799	0.018653	
34	hepatitis_missing_dropped	svm	0.013068	0.006910	
35	hepatitis_missing_dropped	dt	0.004419	0.002771	
36	immunotherapy	rf	0.318120	0.035794	
37	immunotherapy	knn	0.006032	0.025356	
38	immunotherapy	svm	0.004922	0.003697	
39	immunotherapy	dt	0.005246	0.002195	
	17				
	train_acc eval_acc train_f1 eval_f1				
0	1.000000 0.964912 1.000000 0.952381				

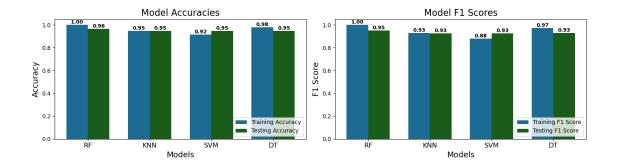
1 0.949451 0.947368 0.929231 0.926829

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                          0.880503
                                     0.925000
3
     0.980220
                0.947368
                          0.973451
                                     0.930233
4
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                1.000000
                          1.000000
                                     1.000000
5
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                0.700000
                          0.865922
                                     0.727273
6
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                0.662500
                          0.743100
                                     0.776860
7
     0.971875
                0.987500
                          0.976864
                                     0.990291
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                0.987500
                          1.000000
                                     0.990291
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                          0.865169
                                     0.741573
     0.618750
                0.637500
                          0.752033
                                     0.771654
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11
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                0.987500
                          0.984615
                                     0.990291
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                1.000000
                          1.000000
                                     1.000000
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                          0.612245
                                     0.500000
     0.849206
14
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                          0.341463
                                     0.500000
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                          1.000000
                                     1.000000
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                          1.000000
                                     0.166667
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                0.333333
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18
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     0.902778
                          0.909091
                                     0.666667
22
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                          0.888889
                                     0.666667
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23
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                          1.000000
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26
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                          0.887892
                                     0.872727
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                0.741935
                          0.959596
                                     0.840000
                0.774194
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                          1.000000
                                     0.857143
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                          0.924528
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39
     0.902778
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                                     0.857143
```

1.4.1 Wisconsin Breast Cancer

Training And Testing Time

Accuracy and F1 Score

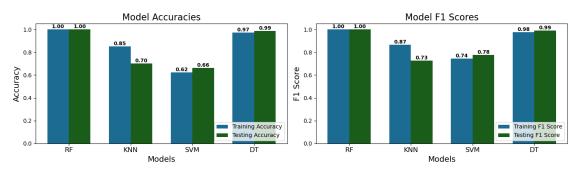


1.4.2 Chronic Kidney Disease

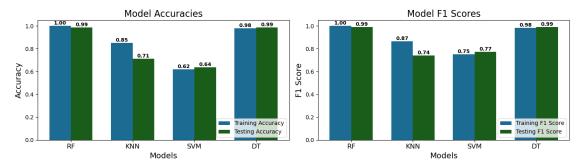
Training And Testing Time

Accuracy and F1 Score

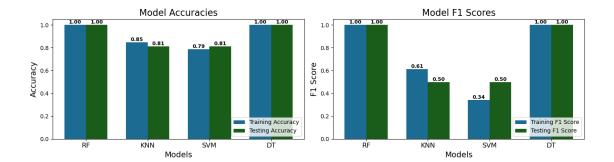
Mean Imputted



Median Imputted



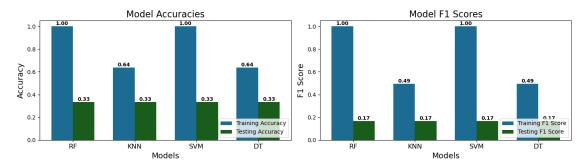
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1.4.3 Covid 19

Training And Testing Time

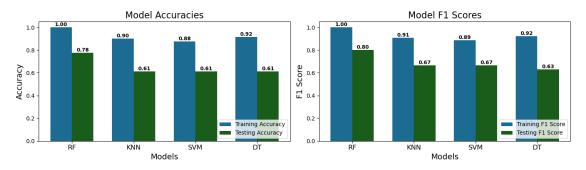
Accuracy and F1 Score



1.4.4 Cryotherapy

Training And Testing Time

Accuracy and F1 Score

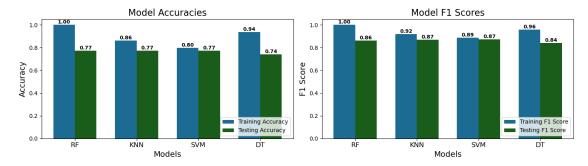


1.4.5 Hepatisis

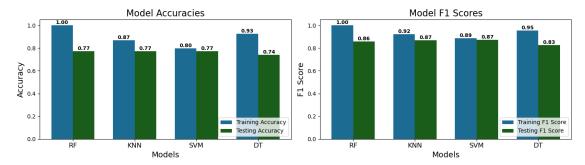
Training And Testing Time

Accuracy and F1 Score

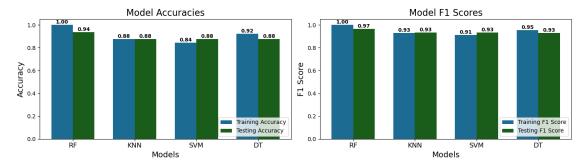
Mean Imputted



Median Imputted



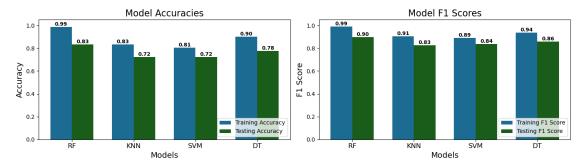
Missing Dropped



1.4.6 Immunotherapy

Training And Testing Time

Accuracy and F1 Score



1.5 COMPARISON

In our replication of the study, we follow the original approach of categorizing model performance into four levels: **excellent execution, good execution, fair execution, and inadequate execution.** However, the original study does not specify the metrics or methodology used to classify these performance categories. In the absence of this information, we have structured our comparison based on a combination of quantitative measures: **Testing Accuracy, Testing F1 Score, and Testing Time.**

In our study, we primarily use **Testing Accuracy** to compare the performance of models across various datasets. If models show identical testing accuracy, we then assess the **Testing F1 Score** for a deeper evaluation of model precision and recall. In cases where the F1 scores are also equivalent, **Testing Time** serves as the final criterion to determine the model's efficiency.

Our Results Comparison

Replicated Study Comparison

Our results in the replicated study exhibit significant differences across almost all datasets when compared to the original study. This variation can primarily be attributed to the lack of detailed methodology disclosed in the original study regarding the specific metrics, hyperparameters, and comparison criteria used. Without this information, our replication had to rely on standard performance metrics (Testing Accuracy, F1 Score, and Testing Time) and commonly used hyperparameters for each model, which likely led to the observed discrepancies in model performance classification across the datasets. This highlights the importance of transparency and detailed documentation in research to ensure reproducibility and accurate comparison of results.