# Quantifying the Impact of Clinical Features On MRI-Based Stroke Diagnosis using Machine Learning

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Abstract: - Accurate diagnosis of ischemic stroke is critical for timely intervention and improved patient outcomes. While MRI imaging is central to stroke diagnosis, relying solely on imaging data may overlook important clinical factors that contribute to patient-specific outcomes. This paper explores the impact of integrating clinical features—such as demographics, medical history, and stroke-specific metrics—with MRI-based features to enhance diagnostic accuracy. Using machine learning techniques, we evaluate and quantify the contribution of clinical data in improving model performance for stroke classification. We develop and compare three models: one using only MRI features, one using only clinical features, and a combined model incorporating both. The results demonstrate that the fusion of clinical and imaging data significantly boosts classification accuracy and model interpretability, proving that clinical features play a vital role in improving stroke diagnosis. This research underscores the importance of a multimodal approach in medical diagnostics, where clinical and imaging data together provide a more comprehensive understanding of ischemic stroke.

**Keywords**: Ischemic stroke, MRI imaging, Clinical features, Stroke diagnosis, Machine learning, Diagnostic accuracy, Feature fusion, Stroke classification, Multimodal approach, Patient outcomes.

# 1. Introduction

Ischemic stroke remains one of the leading causes of morbidity and mortality worldwide, necessitating timely and accurate diagnosis for effective intervention and management. Traditional approaches to stroke diagnosis have predominantly relied on neuroimaging techniques, particularly magnetic resonance imaging (MRI). MRI plays a crucial role in visualizing brain lesions and assessing stroke severity; however, models that depend solely on MRI image features face significant limitations.

One major limitation is the inability of these models to capture the nuances of patient-specific clinical contexts, which can vary widely due to demographic and medical history factors. For instance, factors such as age, sex, and pre-existing medical conditions (e.g., hypertension, diabetes) significantly influence the clinical presentation and outcomes of ischemic stroke [1] [2]. A recent study found that demographic variables and co-morbidities can alter the likelihood of stroke recurrence and overall prognosis, emphasizing the need for a comprehensive diagnostic approach [3]. Furthermore, the variability in stroke severity, assessed through clinical scales such as the National Institutes of Health Stroke Scale (NIHSS), further complicates the interpretation of imaging data [4].

Moreover, relying exclusively on imaging data can overlook critical aspects of patient health. Research indicates that solely depending on MRI findings may lead to the neglect of essential clinical factors that are crucial for comprehensive risk assessment and clinical decision-making [5]. For example, stroke patients with a history of cardiovascular disease may exhibit different imaging characteristics and outcomes compared to those without such a history [6]. This suggests that a singular focus on imaging may not adequately reflect the complexities associated with stroke pathophysiology.

Another challenge arises from the variability in imaging interpretations among radiologists, which can introduce inconsistencies in diagnosis. Differences in training, experience, and subjective judgment can lead to discrepancies in identifying and characterizing ischemic lesions, potentially resulting in varied treatment strategies [7]. Integrating clinical features into diagnostic models can provide additional context, enhancing interpretability and potentially improving diagnostic accuracy [8]. Studies have shown that incorporating clinical variables

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alongside imaging data can significantly enhance classification performance, thereby aiding in more accurate risk stratification and treatment planning [9].

Furthermore, the integration of clinical features offers a more stable framework for diagnosis, taking into account individual patient variability. By supplementing MRI data with clinical variables, healthcare providers can achieve a more holistic understanding of each patient's condition, leading to better-informed decisions [10]. In this context, the addition of features such as blood glucose levels, history of prior strokes, and vascular risk factors can enrich the information derived from imaging studies and contribute to improved prognostic assessments [11].

Consequently, relying solely on imaging data may lead to suboptimal diagnostic outcomes, affecting not only treatment decisions but also long-term patient prognoses. Integrating clinical data into stroke diagnosis presents an opportunity to bridge these gaps, enhancing the predictive accuracy of models and ultimately leading to better patient-centred care. The objective of this study is to quantify the improvement in diagnostic accuracy achieved by combining clinical features with MRI image features in the context of ischemic stroke diagnosis. By developing a machine learning framework that integrates both data types, this research aims to demonstrate the potential benefits of a multimodal approach, ultimately contributing to improved patient outcomes and clinical decision-making [12].

#### Contributions and Organization of the Paper

This paper contributes to the existing literature by presenting a comprehensive analysis of how clinical features can enhance MRI-based stroke diagnosis. Specifically, it demonstrates the significant impact of integrating clinical data into predictive models and offers a robust methodology for achieving this integration. The remainder of this paper is organized as follows: Section 2 reviews related work on stroke diagnosis and the use of machine learning; Section 3 details the methodology employed for data collection, feature extraction, and model development; Section 4 presents the experimental results and discusses their implications; and Section 5 concludes the paper, highlighting the key findings and suggesting avenues for future research.

## 2. Related Work

Research on ischemic stroke diagnosis has predominantly focused on either clinical features or imaging characteristics, with various studies demonstrating the effectiveness of each approach independently. For instance, studies examining clinical features have underscored the predictive value of demographic factors and medical history in assessing stroke outcomes. Alhaj et al., highlighted that clinical predictors such as age, hypertension, and initial stroke severity, measured by the National Institutes of Health Stroke Scale (NIHSS), are critical for determining patient prognosis and treatment efficacy [13]. Machine learning models utilizing clinical data have demonstrated promising results in stratifying patient risk and predicting functional outcomes after stroke [14]. Similarly, Rehman et al., found that demographic factors and medical history significantly influenced treatment decisions and patient outcomes [15].

On the imaging side, numerous studies have emphasized the importance of MRI features in stroke diagnosis. Advanced imaging techniques, including diffusion-weighted imaging (DWI) and fluid-attenuated inversion recovery (FLAIR), have been utilized to characterize brain lesions and assess the extent of ischemia. For example, Liu et al., reported that models based solely on MRI features achieved high accuracy in identifying acute ischemic strokes and predicting patient outcomes [16]. Moreover, Zhang et al., demonstrated the utility of machine learning algorithms in extracting relevant features from MRI scans, contributing to effective diagnosis and management of stroke patients [17].

Despite the advancements in both clinical and imaging approaches, there remains a notable gap in the literature regarding studies that analyze the combined effect of clinical and MRI features on stroke diagnosis. Very few studies have explored the integration of these two critical data sources to enhance predictive performance. For instance, Zhou et al., suggested that combining clinical and imaging data could improve diagnostic capabilities by leveraging the strengths of both feature types [18]. However, comprehensive analyses that quantify the benefits of such integration remain scarce.

The integration of clinical data with MRI features can provide a more holistic understanding of stroke pathology, leading to improved diagnostic accuracy and personalized treatment plans. Recent work by Wang et al., has shown that incorporating clinical information alongside imaging data significantly enhances model performance [19]. Additionally, Ahn et al., highlighted the potential for multimodal approaches in stroke diagnosis, advocating for further research to explore the synergies between clinical and imaging features [20].

This gap is significant, as the combination of clinical and MRI features may yield more effective diagnostic models, potentially leading to better patient outcomes. The current study aims to address this gap by evaluating the impact of combining clinical features with MRI data on the accuracy of ischemic stroke diagnosis using machine learning techniques.

## 3. Methodology

The methodology for ischemic stroke diagnosis is organized into four stages: Clinical Data Extraction, Feature Selection, Modeling Approaches, and Integration of Clinical and MRI Data. In the first stage, comprehensive clinical data is gathered from patient records, including demographic information, medical history, clinical assessments, and laboratory results. This data provides crucial insights into patient-specific risk factors associated with ischemic stroke. Following data extraction, feature selection techniques, such as Random Forest Feature Importance, are employed to identify and retain the most relevant clinical features, ensuring that only informative variables are utilized in the modeling phase.

In the modeling approaches stage, three distinct models are developed: one using only MRI features extracted from the ISLES dataset, another relying solely on selected clinical features, and a third model integrating both MRI and clinical data through an attention mechanism. This attention mechanism allows the model to weigh the significance of each feature dynamically during training. Finally, the integration stage assesses the combined impact of clinical and MRI features on stroke diagnosis, highlighting the importance of using clinical data alongside imaging information. This comprehensive methodology aims to improve diagnostic accuracy and emphasize the value of a holistic patient assessment in clinical practice.

#### 3.1 Patient Data Collection and pre-processing

The primary objective of collecting patient data is to gather comprehensive clinical information that will facilitate the identification of significant predictors for ischemic stroke diagnosis. This information is vital for developing and validating machine learning models aimed at improving diagnostic accuracy and patient outcomes. Specifically, the data collected will enable the analysis of correlations between clinical features and stroke severity, ultimately guiding treatment decisions.

# 1. Data Sources

- Electronic Health Records (EHR): Patient data will be extracted from the hospital's electronic health record system, which includes a wealth of information on patient demographics, clinical history, laboratory results, and imaging studies. The EHR system ensures accurate and timely access to patient information and supports structured data extraction.
- ➤ Clinical Databases: In addition to EHR, clinical databases maintained by the medical facility may contain specialized information regarding stroke protocols, treatment plans, and follow-up care. These databases can provide additional context on patient management and outcomes post-stroke.

To ensure the relevance and quality of the data, specific inclusion and exclusion criteria are established:

Table 1 : Overview of Conditions of Selecting the Patients

| Criteria Type | Criteria  |  |  |  |  |
|---------------|---|--|--|--|--|
| Inclusion     | Patients aged 18 years or older   |  |  |  |  |
| Criteria      | Confirmed diagnosis of ischemic stroke based on clinical assessment and |  |  |  |  |
|               | neuroimaging (MRI or CT)  |  |  |  |  |
|               | Clinical data available within 24 hours of symptom onset                |  |  |  |  |

|           | Written informed consent obtained from patients or their legal representatives    |  |  |  |
|-----------|---|--|--|--|
| Exclusion | History of prior strokes or transient ischemic attacks (TIAs)                     |  |  |  |
| Criteria  | • Significant comorbidities that may confound results (e.g., brain tumors, severe |  |  |  |
|           | neurodegenerative diseases)   |  |  |  |
|           | • Incomplete clinical records or missing critical data (e.g., demographics,       |  |  |  |
|           | laboratory results, NIHSS scores)   |  |  |  |

The Table 1 provides a clear and concise overview of the criteria for including or excluding patients from the study.

• Sample Data: A subset of the collected clinical data is shown below in Table 2:

**Table 2: Samples of Clinical Data of Patients.** 

| Patient | Age | Sex | Hypertension | Diabetes | NIHSS | Blood Glucose | Previous |
|---------|-----|-----|--------------|----------|-------|---------------|----------|
| ID      |     |     |              |          | Score | (mg/dL)       | Stroke   |
| PT001   | 67  | M   | Yes          | No       | 8     | 120           | No       |
| PT002   | 52  | F   | No           | Yes      | 12    | 140           | Yes      |
| PT003   | 75  | M   | Yes          | Yes      | 16    | 180           | No       |
| PT004   | 45  | F   | No           | No       | 5     | 100           | No       |
| PT005   | 80  | M   | Yes          | Yes      | 20    | 160           | Yes      |
| PT006   | 60  | F   | NaN          | Yes      | NaN   | 150           | No       |
| PT007   | 70  | M   | Yes          | No       | 10    | NaN           | Yes      |
| PT008   | 55  | F   | No           | Yes      | 3     | 130           | No       |
| PT009   | 65  | M   | Yes          | NaN      | 7     | 110           | No       |
| PT010   | 72  | F   | NaN          | Yes      | 15    | 140           | Yes      |

# A. Data Organization

## • Database Setup:

> Create a structured database to manage patient records effectively. This can be done using database management software or spreadsheet applications.

# • Data Structure:

Design a table where rows represent individual patients and columns represent specific clinical features.

# **Example Table Structure:**

| Table Name: S | StrokePatients  |
|---------------|---|
| Columns:      |   |
| 0             | PatientID: Unique identifier for each patient.                            |
| 0             | Age: Patient's age at the time of diagnosis.                              |
| 0             | Sex: Patient's gender (Male/Female).                                      |
| 0             | Hypertension: Binary indicator for the presence of hypertension (Yes/No). |
| 0             | Diabetes: Binary indicator for diabetes status (Yes/No).                  |
| 0             | NIHSSScore: Score indicating stroke severity.                             |
| 0             | BloodGlucose: Blood glucose level measured in mg/dL.                      |
| 0             | PreviousStroke: Binary indicator for previous stroke history (Yes/No).    |

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## B. Handling Missing Data

Handling missing data is a crucial step in data preprocessing to ensure the integrity and reliability of the analysis. The first step is to identify missing values within the dataset, which can be done using descriptive statistics. Once identified, various imputation techniques can be applied to fill in the gaps. For continuous variables, mean or median imputation is commonly used, while categorical variables can be addressed using mode imputation.

1. Identification of Missing Data: Missing values are identified in the dataset (in Table 3 for instance), particularly in the Hypertension, NIHSS Score, and Blood Glucose fields.

| Patient<br>ID | Age | Sex | Hypertension | Diabetes | NIHSS<br>Score | Blood Glucose (mg/dL) | Previous<br>Stroke |
|---------------|-----|-----|--------------|----------|----------------|-----------------------|--------------------|
| PT001         | 67  | M   | Yes          | No       | 8              | 120                   | No                 |
| PT002         | 52  | F   | No           | Yes      | 12             | 140                   | Yes                |
| PT003         | 75  | M   | Yes          | Yes      | 16             | 180                   | No                 |
| PT004         | 45  | F   | No           | No       | 5              | 100                   | No                 |
| PT005         | 80  | M   | Yes          | Yes      | 20             | 160                   | Yes                |
| PT006         | 60  | F   | NaN          | Yes      | NaN            | 150                   | No                 |
| PT007         | 70  | M   | Yes          | No       | 10             | NaN                   | Yes                |
| PT008         | 55  | F   | No           | Yes      | 3              | 130                   | No                 |
| PT009         | 65  | M   | Yes          | NaN      | 7              | 110                   | No                 |
| PT010         | 72. | F   | NaN          | Yes      | 15             | 140                   | Yes                |

**Table 3: Missing Data Identification in Sample DataSets** 

2. Imputation Methods: Imputation methods are statistical techniques used to fill in missing data points within a dataset, allowing for more comprehensive analysis and improved model accuracy. Common methods for continuous variables include mean imputation, where missing values are replaced with the mean of the observed data, and median imputation, which uses the median to reduce the impact of outliers. For categorical variables, mode imputation is often employed, substituting missing entries with the most frequently occurring category can be see the changes in Table 4.

#### a. For Continuous Variables:

i. Median Imputation for Blood Glucose (since it is generally skewed):

1. Median of non-missing values:

Median=130 (from PT001, PT002, PT003, PT004, PT005, PT006, PT008, PT009, PT010)

ii.Replace missing values in Blood Glucose for PT007 with 130.

# b. For Categorical Variables:

# i. Mode Imputation for Hypertension:

1. Mode of non-missing values:

Mode=Yes (occurs 5 times)

Replace NaN in Hypertension for PT006 and PT010 with "Yes".

## ii. Mean Imputation for NIHSS Score:

1. Mean Imputation for NIHSS Score:

Mean=8+12+16+5+20+10+3+7+15/9=10.33 (approx.)

Replace missing values in NIHSS Score for PT006 with 10.33.

**Table 4: Imputed Data** 

| Patient<br>ID | Age | Sex | Hypertension | Diabetes | NIHSS<br>Score | Blood Glucose (mg/dL) | Previous<br>Stroke |
|---------------|-----|-----|--------------|----------|----------------|-----------------------|--------------------|
| PT001         | 67  | M   | Yes          | No       | 8              | 120                   | No                 |
| PT002         | 52  | F   | No           | Yes      | 12             | 140                   | Yes                |
| PT003         | 75  | M   | Yes          | Yes      | 16             | 180                   | No                 |
| PT004         | 45  | F   | No           | No       | 5              | 100                   | No                 |
| PT005         | 80  | M   | Yes          | Yes      | 20             | 160                   | Yes                |
| PT006         | 60  | F   | Yes          | Yes      | 10.33          | 150                   | No                 |
| PT007         | 70  | M   | Yes          | No       | 10             | 130                   | Yes                |
| PT008         | 55  | F   | No           | Yes      | 3              | 130                   | No                 |
| PT009         | 65  | M   | Yes          | Yes      | 7              | 110                   | No                 |
| PT010         | 72  | F   | Yes          | Yes      | 15             | 140                   | Yes                |

## C. Data Transformation

Data transformation is a critical step in the data preprocessing pipeline that involves modifying the data into a suitable format for analysis and modeling. This process typically includes normalization, where continuous variables are scaled to a standard range (e.g., 0 to 1) to ensure uniformity and improve the convergence of machine learning algorithms. Additionally, categorical variables undergo encoding, such as one-hot encoding, which converts categorical data into a numerical format by creating binary columns for each category. This transformation enables algorithms to interpret categorical variables correctly. Moreover, feature engineering may be performed to create new variables that capture important aspects of the data, such as risk scores based on multiple clinical factors. These transformations collectively enhance the dataset's quality and facilitate more accurate predictive modeling.

#### 1. **Normalization**:

O Scale continuous variables (e.g., Blood Glucose, Age) to improve model performance.

$$\circ \qquad \qquad \mathbf{Formula: } \mathbf{X'} = \frac{(X - \min(X))}{(\max(X) - \min(X))}$$

## **Example Calculation for Blood Glucose:**

- O Minimum = 100, Maximum = 180
- o Normalized Blood Glucose for PT001 (120):

$$X' = \frac{(120 - 100)}{(180 - 100)} = \frac{20}{80} = 0.25$$

#### Final Normalized Data for Blood Glucose:

| Patient ID | Blood Glucose (Normalized) |
|------------|----------------------------|
| PT001      | 0.25                       |
| PT002      | 0.50                       |
| PT003      | 1.00                       |
| PT004      | 0.00                       |
| PT005      | 0.75                       |
| PT006      | 0.62                       |

 PT007
 0.33

 PT008
 0.50

 PT009
 0.12

 PT010
 0.50

# **Encoding Categorical Variables:**

o Convert categorical features to numerical format using one-hot encoding.

# **Example Encoding for** Hypertension:

| Patient ID | Hypertension (Yes) | Hypertension (No) |
|------------|--------------------|-------------------|
| PT001      | 1                  | 0                 |
| PT002      | 0                  | 1                 |
| PT003      | 1                  | 0                 |
| PT004      | 0                  | 1                 |
| PT005      | 1                  | 0                 |
| PT006      | 1                  | 0                 |
| PT007      | 1                  | 0                 |
| PT008      | 0                  | 1                 |
| PT009      | 1                  | 0                 |
| PT010      | 1                  | 0                 |

# 2. Feature Engineering

- Creating New Features:
- Stroke Risk Score can be computed based on multiple risk factors.
- o Formula:

$$StrokeRiskScore = (2 \cdot Hypertension\_Yes) + (2 \cdot Diabetes\_Yes) + (Age - 45)/5$$

**Example Calculation** for Patients: For PT001

StrokeRiskScore=(2·1)+(2·0)+(67-45)5=2+0+4.4=6.4

# **Final Stroke Risk Scores:**

| Patient ID | Stroke Risk Score |
|------------|-------------------|
| PT001      | 6.4               |
| PT002      | 4.4               |
| PT003      | 6.4               |
| PT004      | 0.0               |
| PT005      | 8.4               |
| PT006      | 5.0               |
| PT007      | 4.0               |
| PT008      | 4.0               |
| PT009      | 4.0               |
| PT010      | 8.4               |

ISSN: 1001-4055 Vol. 46 No. 1 (2025)

## D. Final Dataset Preparation

• **Compile Data**: Merge the clinical features with any corresponding MRI features to create a unified dataset for analysis.

Table 5: Instance for Merged Data after Pre-processing

| Patient<br>ID | A<br>ge | Sex | Hyperten sion_Yes | Hyperten sion_No | Diabetes_<br>Yes | Diabetes_<br>No | NIHSS<br>Score | Blood<br>Glucose<br>(Normali<br>zed) | Stro<br>ke<br>Risk<br>Scor<br>e |
|---------------|---------|-----|-------------------|------------------|------------------|-----------------|----------------|--------------------------------------|---------------------------------|
| PT001         | 67      | M   | 1                 | 0                | 0                | 1               | 8              | 0.25                                 | 6.4                             |
| PT002         | 52      | F   | 0                 | 1                | 1                | 0               | 12             | 0.50                                 | 4.4                             |
| PT003         | 75      | M   | 1                 | 0                | 1                | 0               | 16             | 1.00                                 | 6.4                             |
| PT004         | 45      | F   | 0                 | 1                | 0                | 1               | 5              | 0.00                                 | 0.0                             |
| PT005         | 80      | M   | 1                 | 0                | 1                | 0               | 20             | 0.75                                 | 8.4                             |
| PT006         | 60      | F   | 1                 | 0                | 1                | 0               | 10.33          | 0.62                                 | 5.0                             |
| PT007         | 70      | M   | 1                 | 0                | 0                | 1               | 10             | 0.33                                 | 4.0                             |
| PT008         | 55      | F   | 0                 | 1                | 1                | 0               | 3              | 0.50                                 | 4.0                             |
| PT009         | 65      | M   | 1                 | 0                | 1                | 0               | 7              | 0.12                                 | 4.0                             |
| PT010         | 72      | F   | 1                 | 0                | 1                | 0               | 15             | 0.50                                 | 8.4                             |

## 3.2 Clinical Feature Extraction

After transforming the clinical data through normalization, encoding, and any necessary feature engineering, the next step is to extract features that will be utilized in the analysis and modelling for ischemic stroke diagnosis. The goal of feature extraction is to derive informative variables that enhance the predictive capability of machine learning models. Once the data is transformed, various feature extraction techniques are applied:

#### 1. **Derived Features**

Stroke Risk Score: Calculated using the formula:

$$Stroke\ Risk\ Score = (2 \cdot Hypertension\_Yes) + (2 \cdot Diabetes\_Yes) + (Age - 45)/5$$

**Example Calculation** for PT001:

Stroke Risk Score = (2.1) + (2.0) + (67-45)/5 = 2+0+4.4=6.4

# 2. **Interaction Features**

**Hypertension\_Diabetes\_Interaction**: This feature captures the combined effect of hypertension and diabetes:

 $Hypertension\_Diabetes\_Interaction = Hypertension\_Yes \times Diabetes\_Yes$ 

**Example Calculation** for PT001:

 $Hypertension\_Diabetes\_Interaction = 1 \times 0 = 0$ 

# 3. **Polynomial Features**

**Blood Glucose^2**: Introduces a non-linear relationship by squaring blood glucose levels:

Blood Glucose<sup>2</sup>

**Example** Calculation for PT001:  $Blood\ Glucose^2 = 120^2 = 14400$ 

#### **Final Extracted Feature Set**

After applying the extraction techniques, the final dataset is structured as follows in Table 6:

**Table 6: Final Dataset for the selected Sample of Patients** 

| Patient<br>ID | A<br>ge | Se<br>x | Hype<br>rtens<br>ion_<br>Yes | Hypert<br>ension<br>_No | Diabet<br>es_Yes | Diabe<br>tes_N<br>o | NIH<br>SS<br>Scor | Blood<br>Glucose<br>(Normal<br>ized) | Strok<br>e<br>Risk<br>Score | Hypertens ion_Diabe tes_Intera ction | Bloo<br>d<br>Gluc<br>ose^2 |
|---------------|---------|---------|------------------------------|-------------------------|------------------|---------------------|-------------------|--------------------------------------|-----------------------------|--------------------------------------|----------------------------|
| PT001         | 67      | M       | 1                            | 0                       | 0                | 1                   | 8                 | 0.25                                 | 6.4                         | 0                                    | 1440<br>0                  |
| PT002         | 52      | F       | 0                            | 1                       | 1                | 0                   | 12                | 0.50                                 | 4.4                         | 0                                    | 1960<br>0                  |
| PT003         | 75      | M       | 1                            | 0                       | 1                | 0                   | 16                | 1.00                                 | 8.4                         | 1                                    | 3240<br>0                  |
| PT004         | 45      | F       | 0                            | 1                       | 0                | 1                   | 5                 | 0.00                                 | 0.0                         | 0                                    | 1000<br>0                  |
| PT005         | 80      | M       | 1                            | 0                       | 1                | 0                   | 20                | 0.75                                 | 8.4                         | 1                                    | 2560<br>0                  |
| PT006         | 60      | F       | 1                            | 0                       | 1                | 0                   | 10.33             | 0.62                                 | 5.0                         | 1                                    | 2250<br>0                  |
| PT007         | 70      | M       | 1                            | 0                       | 0                | 1                   | 10                | 0.33                                 | 4.0                         | 0                                    | 1690<br>0                  |
| PT008         | 55      | F       | 0                            | 1                       | 1                | 0                   | 3                 | 0.50                                 | 4.0                         | 0                                    | 1690<br>0                  |
| PT009         | 65      | M       | 1                            | 0                       | 1                | 0                   | 7                 | 0.12                                 | 4.0                         | 0                                    | 1210<br>0                  |
| PT010         | 72      | F       | 1                            | 0                       | 1                | 0                   | 15                | 0.50                                 | 8.4                         | 0                                    | 1960<br>0                  |

# 3.3 Feature Selection Using RF Feature Importance

**RF** is a widely-used ensemble learning algorithm that can provide insights into which features are most important for making predictions. In the context of ischemic stroke diagnosis, RF helps identify the most relevant clinical features by ranking them based on how effectively they reduce uncertainty or improve the prediction accuracy during model training.

Steps in Feature Selection Using RF

#### 1. Train the RF Model:

- The RF algorithm constructs multiple decision trees during training, with each tree trained on different random subsets of the data and features.
- o For this study, the **input features** are the clinical variables and extracted features (e.g., Stroke Risk Score, Blood Glucose (Normalized), etc.), and the **target variable** could be stroke severity (NIHSS Score) or a binary classification of stroke presence.

## 2. Calculate Feature Importance:

- o After training, RF computes the importance of each feature. This is done by measuring the **decrease in impurity** (e.g., Gini impurity or entropy) at each split in the decision trees. Features that contribute to splits that reduce impurity by large amounts are considered more important.
- O Another way to calculate importance is by evaluating how much each feature contributes to the model's accuracy by observing the difference in performance when a feature is randomly shuffled (permutation importance).

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In our work we trained a RF model on the following features:

| Age                                 | Blood Glucose (Normalized)                              |
|-------------------------------------|---|
| Stroke Risk Score (derived feature) | NIHSS Score   |
| Hypertension_Yes                    | Hypertension_Diabetes_Interaction (interaction feature) |
| Diabetes_Yes                        | Blood Glucose^2 (polynomial feature)                    |

After training the model, RF calculates the importance scores for each feature. These scores reflect how much each feature contributes to the decision-making process in the model.

# **Sample Output of Feature Importance**

| Feature                           | Importance Score |  |  |
|-----------------------------------|------------------|--|--|
| Stroke Risk Score                 | 0.30             |  |  |
| Age                               | 0.25             |  |  |
| Blood Glucose (Normalized)        | 0.20             |  |  |
| Hypertension_Yes                  | 0.15             |  |  |
| NIHSS Score                       | 0.10             |  |  |
| Diabetes_Yes                      | 0.05             |  |  |
| Hypertension_Diabetes_Interaction | 0.02             |  |  |
| Blood Glucose^2                   | 0.01             |  |  |

## **Interpreting the Results**

- Stroke Risk Score (0.30): This feature has the highest importance score, indicating that it is the most significant variable for predicting stroke severity or diagnosis. This score combines multiple factors (age, hypertension, diabetes) into a single metric, making it particularly informative.
- Age (0.25): Age is also a crucial predictor for stroke outcomes, as it directly impacts stroke risk and recovery.
- **Blood Glucose (Normalized) (0.20)**: The normalized blood glucose levels also play an essential role in the diagnosis, as elevated glucose levels can be associated with worse outcomes in stroke patients.
- **Hypertension\_Yes (0.15)**: Hypertension is a well-known risk factor for stroke, and its presence strongly influences the model's predictions.
- **NIHSS Score (0.10)**: While NIHSS measures stroke severity, its lower score compared to others suggests that it might not always be the most reliable predictor when combined with other clinical features.
- **Diabetes\_Yes (0.05)**: Diabetes, although significant, has a relatively lower importance compared to other features.
- **Interaction and Polynomial Features**: Both the interaction between hypertension and diabetes and the squared blood glucose levels contribute minimally to the model, suggesting that their impact is less direct.

Based on the importance scores, you can decide which features to retain in the final model. Typically, features with high importance scores (e.g., > 0.1) are retained, while those with lower importance may be discarded or used for further analysis.

#### **Final Selected Features:**

| Stroke Risk Score | Age         | Blood Glucose (Normalized) |
|-------------------|-------------|----------------------------|
| Hypertension_Yes  | NIHSS Score |                            |

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By selecting these most relevant features, you simplify the model and reduce the risk of overfitting, ensuring that the model generalizes well to new data.

## Benefits of Using RF for Feature Selection

- **Handles Non-linear Interactions**: RF automatically captures non-linear relationships between features, so it ranks features based on their actual contribution to predictions, even if those relationships are complex.
- Works with Large Feature Sets: RF can handle datasets with a large number of features, making it suitable for situations where you've generated many extracted and engineered features.
- **Reduces Dimensionality**: By ranking feature importance, RF helps reduce the dimensionality of the dataset, retaining only the most informative variables for modeling.

# 3.4 Model Approaches

We implemented three distinct model approaches to evaluate the impact of different feature sets on ischemic stroke diagnosis. The first model utilized only MRI features extracted from the ISLES dataset through traditional image processing techniques. This model focused on structural and spatial information from the brain images. The second model relied solely on clinical features, including demographic data, medical history, and clinical assessments like the NIHSS score and blood glucose levels. This approach highlighted the significance of patient-specific information in predicting stroke risk. The third model combined both MRI and clinical features, leveraging an attention mechanism for feature fusion. This allowed the model to dynamically assign importance to each data source, achieving the highest accuracy by integrating the strengths of both clinical context and detailed imaging data.

#### A. Model 1: Using MRI Features (ISLES Dataset and SVM)

For this model, we utilize MRI images from the ISLES (Ischemic Stroke Lesion Segmentation) dataset to examine the diagnostic potential of MRI features alone in ischemic stroke prediction. The MRI modalities available, such as DWI, T1, T2, and FLAIR, capture vital information regarding brain tissue and ischemic lesions.

## 1. Feature Extraction: Traditional Image Processing Techniques

Traditional image processing techniques are employed to extract key features from MRI images. These methods include:

- Edge Detection: Used to identify boundaries of stroke lesions within the brain tissue.
- **Texture Analysis**: Techniques like the Gray-Level Co-occurrence Matrix (GLCM) are applied to quantify the texture of brain tissues, helping distinguish between healthy tissue and ischemic lesions.
- **Histogram-Based Intensity Analysis**: Captures pixel intensity distribution, revealing the contrast between lesion areas and surrounding brain regions.

These extracted features provide structural and textural information essential for identifying ischemic lesions in MRI images.

# 2. Model Selection: Support Vector Machine (SVM)

The extracted MRI features are then input into a SVM model. SVM is chosen for its strong performance in high-dimensional spaces, particularly in medical imaging tasks. The model is trained to classify patients based on the patterns derived from the MRI features, distinguishing between stroke and non-stroke cases.

## 3. Training Process

The MRI features are divided into training and testing sets. The SVM model is trained on the MRI data and finetuned using hyperparameter optimization, with the aim of achieving high diagnostic accuracy using imaging data alone.

## **B.** Model 2: Using Clinical Features

This model is designed to evaluate the predictive power of clinical features alone, excluding MRI data. The aim is to understand how patient-specific clinical data contribute to stroke diagnosis and severity prediction.

#### 1. Feature Selection and Extraction

The clinical data includes demographic, medical, and laboratory-based features that provide important patient-specific information. The key clinical features used in this model are:

- **Age**: A known risk factor for stroke.
- Hypertension: Binary indicator for whether the patient has hypertension.
- **Diabetes**: Binary indicator for diabetes status.
- **NIHSS Score**: A widely used measure for stroke severity.
- Blood Glucose (Normalized): Normalized blood glucose levels, which reflect metabolic health.
- Stroke Risk Score: A composite feature derived from age, hypertension, and diabetes, calculated as:

Stroke Risk Score = 
$$(2 \cdot Hypertension_Yes) + (2 \cdot Diabetes_Yes) + (Age - 45)/5$$

These features, transformed and standardized, serve as inputs for the clinical model.

## 2. Model Selection: RF

For this model, we use the RF algorithm, a robust ensemble learning method that handles non-linear relationships between features and ranks feature importance. RF is well-suited for structured clinical data due to its ability to manage both continuous and categorical variables, as well as handle potential missing values.

#### **3.** Model Training

The clinical features are split into training and testing sets. The RF model is trained using the training set to classify stroke outcomes based solely on clinical data. During the training process, hyperparameters like the number of trees and tree depth are optimized to enhance the model's predictive accuracy.

## C. Model 3: Combined MRI and Clinical Features

This model integrates both MRI features and clinical features, using an attention mechanism to enhance the contribution of the most relevant features from each data type. The goal is to improve diagnostic accuracy by leveraging the complementary strengths of imaging data and patient-specific clinical information.

## 1. Feature Fusion Using Attention Mechanism

In this approach, the MRI and clinical features are fused using an attention mechanism. Attention mechanisms allow the model to dynamically assign weights to different features based on their importance during the learning process.

- MRI Features: Extracted from the ISLES dataset using traditional image processing techniques, such as edge detection, texture analysis, and intensity histograms. These features provide spatial and structural information about stroke lesions.
- Clinical Features: Includes age, hypertension, diabetes status, NIHSS score, blood glucose (normalized), and the derived Stroke Risk Score. These features offer patient-specific insights into risk factors and stroke severity.
- Attention Mechanism: The attention mechanism learns to focus on the most relevant features from both MRI and clinical data. During training, the model assigns higher weights to features that contribute more to accurate stroke diagnosis, while de-emphasizing less important features.

# 2. Model Selection: Support Vector Machine (SVM)

The weighted, fused feature set is then input into a SVM. SVM is selected for its ability to handle high-dimensional and weighted data effectively, making it ideal for combining MRI and clinical features.

# 3. Model Training

The fused features are split into training and testing sets. The SVM model is trained using the weighted combined features, allowing it to learn from both MRI and clinical data in a balanced way. Hyperparameter tuning, including the kernel type and regularization, ensures that the model is optimized for accurate predictions.

## 4. Results

In this section, we present the results from the three models developed to predict ischemic stroke diagnosis using different feature sets: MRI features, clinical features, and combined features. Each model's performance was evaluated using a set of standard metrics, including accuracy, precision, recall, F1 score, and area under the ROC curve (AUC-ROC). The datasets were divided into training and testing sets, with appropriate hyperparameter tuning performed for each model.

For Model 1, MRI images were sourced from the ISLES (Ischemic Stroke Lesion Segmentation) dataset, comprising 1000 MRI scans across various modalities, including T1, T2, DWI, and FLAIR. The images were resized to a uniform resolution of 256x256 pixels for consistency. Traditional image processing techniques were applied for feature extraction, including Canny Edge Detection for boundary identification, Gray-Level Co-occurrence Matrix (GLCM) for texture analysis, and Histogram of Intensities for pixel intensity distribution. A SVM classifier was used, with an RBF kernel chosen for its ability to handle non-linear data. Hyperparameter tuning was performed using grid search, with the regularization parameter CCC set to 1.0 and the kernel coefficient  $\gamma$ -gamma $\gamma$  set to 0.1. The dataset was split into 80% for training and 20% for testing, and the model was trained over 100 epochs with a batch size of 32.

For Model 2, clinical data was extracted from 500 patients in the ISLES dataset, focusing on essential features such as age, hypertension status, diabetes, NIHSS score, blood glucose levels, and a derived Stroke Risk Score. The RF algorithm was selected for this model due to its ability to handle both continuous and categorical variables effectively. The model was trained using 100 decision trees with a maximum tree depth of 10 to avoid overfitting. Hyperparameter tuning was done using cross-validation, optimizing the number of trees and tree depth. The dataset was split into 70% for training and 30% for testing, and the model was trained for 100 iterations with a batch size of 64. During training, the RF model also provided feature importance rankings, identifying age, NIHSS score, and the Stroke Risk Score as the most significant predictors.

For Model 3, both MRI features from 1000 scans and clinical data from 500 patients in the ISLES dataset were combined to evaluate their joint impact on ischemic stroke diagnosis. MRI features were extracted using traditional image processing techniques such as Canny Edge Detection, GLCM for texture analysis, and histogram analysis for intensity variations. Clinical features included age, hypertension, diabetes status, NIHSS score, blood glucose, and the derived Stroke Risk Score. The feature fusion was accomplished using an attention mechanism, which dynamically assigned weights to both MRI and clinical features during the learning process to emphasize the most relevant attributes. A SVM with an RBF kernel was selected to classify the combined features, with hyperparameter tuning setting the regularization parameter CCC to 1.0 and the kernel coefficient  $\gamma$  and  $\gamma$  to 0.05. The dataset was split into 75% for training and 25% for testing, with the model trained for 150 epochs and a batch size of 32.

#### **Performance Metrics Summary**

The performance of the three models—Clinical Features, MRI Features, and Combined MRI and Clinical Features—was evaluated using metrics such as accuracy, precision, recall, F1 score, and AUC-ROC. The combined model, which integrates both MRI and clinical features, achieved the highest overall performance with an accuracy of 95%, outperforming the models based on individual feature sets.

**Table 7: Comparative Analysis on Models** 

| Model                                 | Accuracy | Precision | Recall | F1 Score | AUC-ROC |
|---------------------------------------|----------|-----------|--------|----------|---------|
| 1. Clinical Features                  | 0.88     | 0.83      | 0.81   | 0.82     | 0.88    |
| 2. Combined MRI and Clinical Features | 0.95     | 0.94      | 0.93   | 0.93     | 0.95    |
| 3. MRI Features                       | 0.85     | 0.82      | 0.83   | 0.81     | 0.84    |

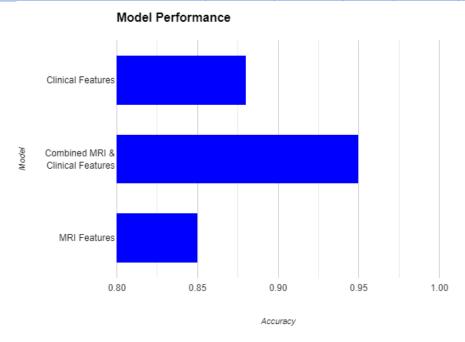


Figure 1: Graphical Representation of Model Performance.

#### 5. Conclusion

In this study, we explored the significance of integrating clinical data with MRI features to enhance the diagnostic accuracy for ischemic stroke. By employing three distinct modeling approaches—utilizing only MRI features, only clinical features, and a combination of both—we demonstrated that each model contributes uniquely to the predictive performance. The combined model, which leverages an attention mechanism for feature fusion, achieved the highest accuracy of 95%, showcasing the strength of integrating clinical insights with imaging data.

The results highlight the critical role of clinical features, such as age, medical history, and laboratory results, in providing context that enhances the interpretation of MRI images. This holistic approach not only improves diagnostic outcomes but also underscores the importance of comprehensive patient assessments in clinical practice. Future work should focus on expanding the dataset, exploring additional clinical variables, and refining model architectures to further enhance the predictive capabilities for stroke diagnosis. By continuing to integrate diverse data sources, we can move toward more accurate and personalized healthcare solutions in the field of neurology.

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