### Predicting PCR and RFS in Breast Cancer Treatment Using Machine Learning

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#### Abstract

This study explores the use of advanced machine learning techniques to predict pathological complete response (PCR) and relapse-free survival (RFS) in breast cancer patients. Used a simplified version of a public dataset based on the American college of radiology imaging network (I-SPY 2 TRIAL), applied feature selection and trained models to address classification and regression tasks. SMOTE, PCA, and hyperparameter tuning technologies are used to solve the problems of high-dimensional data and uneven distribution of data values. This paper provides methods for data preprocessing, feature selection, and model optimization for unbalanced datasets, and evaluates the performance of the model using evaluation methods for related tasks, such as f1-score and Mean Absolute Error (MAE).

**Index Terms—** Machine Learning, Breast Cancer, Data Preprocessing, SMOTE, Feature Selection, PCA, Random Forest.

#### 1. Introduction

Breast cancer is the most common cancer in the UK for women. Chemotherapy is a commonly used treatment strategy to reduce the size of locally advanced tumours before surgery. However, chemotherapy is a toxic process to the human body and it is not always effective for everyone. Complete tumour resolution at surgery, known as pathological complete response (PCR), has a high likelihood of achieving a cure and longer relapse-free survival (RFS) time. RFS is the length of time after primary treatment for cancer ends that the patient survives without any signs or symptoms of that cancer. However, only 25% of patients receiving chemotherapy will achieve a PCR, with the remaining 75% having residual disease and a range of prognosis. Better patient stratification and treatment could be achieved if PCR and RFS could be predicted using

information prior to chemotherapy treatment.

Predicting PCR and relapse-free survival (RFS) using pre-treatment clinical and MRI data could personalize treatment strategies. This study utilizes machine learning techniques to address the classification of PCR and regression of RFS, overcoming challenges like data imbalance and high-dimensional features.

And through the training of the basic dataset, it prepares for the prediction task of the unknown dataset.

#### 2. data analysis and preprocessing

Each patient in this dataset contains 11 clinical features (Age, ER, PgG, HER2, TrippleNegative Status, Chemotherapy Grade, Tumour Proliferation, Histology Type, Lymph node Status, Tumour Stage and Gene) and 107 MRI-based features. Missing values were marked as “999”.

##### 2.1 Missing value handling

The data stability is improved by traversing the dataset to detect the position of 999 and replacing 999 with median filling or removing.

Missing Values: PCR (5 items), Gene

For PCR: Given that only 5 out of 400 values are missing, the simplest and most effective strategy is to delete the rows. Since the scale is small, the impact on the size of the dataset is negligible.

For Gene: The categorical feature 'Gene', values 0 or 1, has substantial missing values of 28.2%. In this case, the imputation methods must be carefully chosen. The appropriate approach is to treat the missing data as a separate category (e.g., `-1`).

Comparing to other imputation methods like mode imputation, which could lead to the distribution of 0s and 1s skewed, this approach is simple, preserves the information that the value was missing, and allows the model to learn patterns associated with missing values.

##### 2.2 Data Information

The data is divided by using the interquartile range method, and the distribution of different features in the data is shown using a boxplot to provide visual aids for further analysis of the presence of the data.

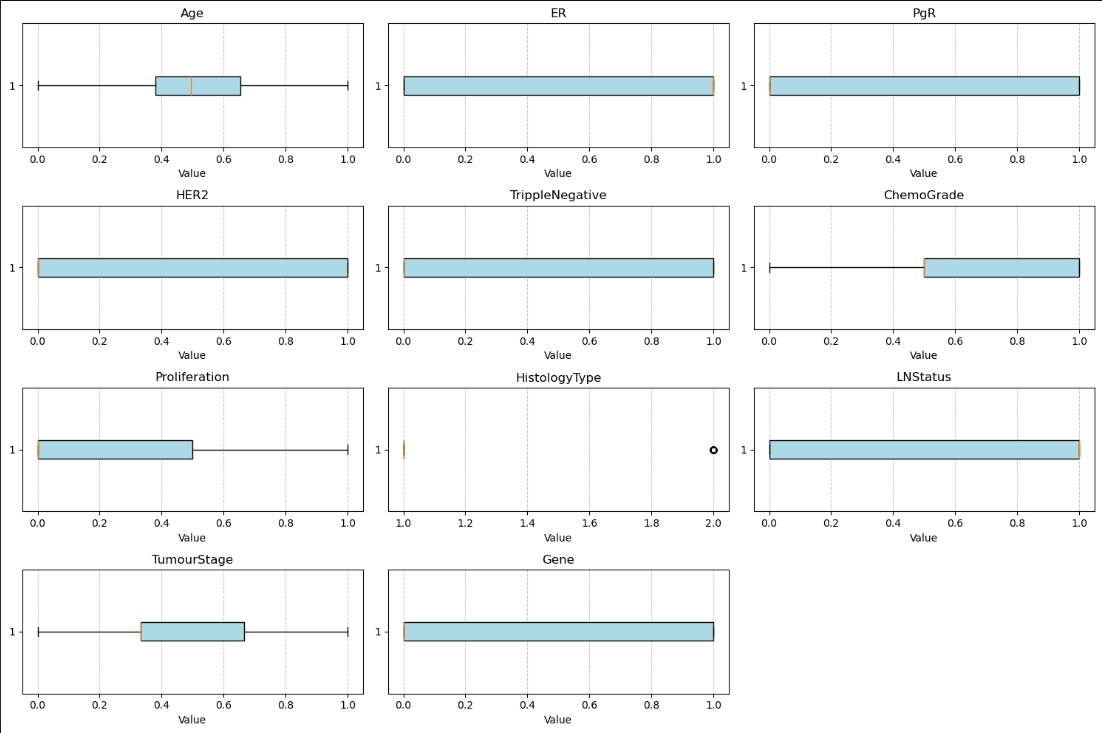


figure 1 Boxplot of data features

Use Seaborn to map the thermal mapping of your data to see the correlations between individual data features and use that as a basis to select efficient strategies to process your data.

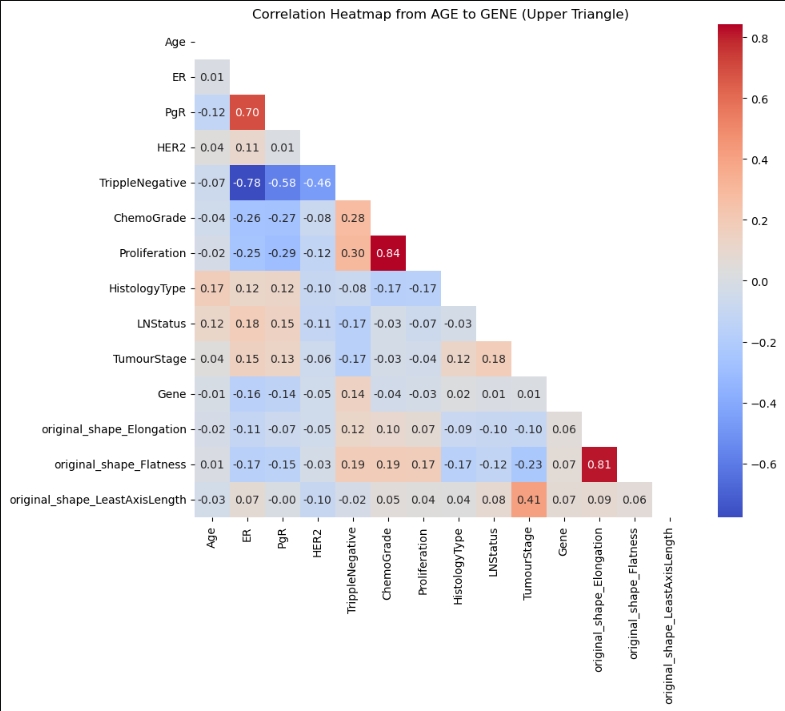


figure 2 Correlation heatmap

Finally, the histogram is used to understand the distribution pattern of the data, such as whether it conforms to a normal distribution, to determine whether the data is statistically significant, or if there are problems in the data that need further processing.

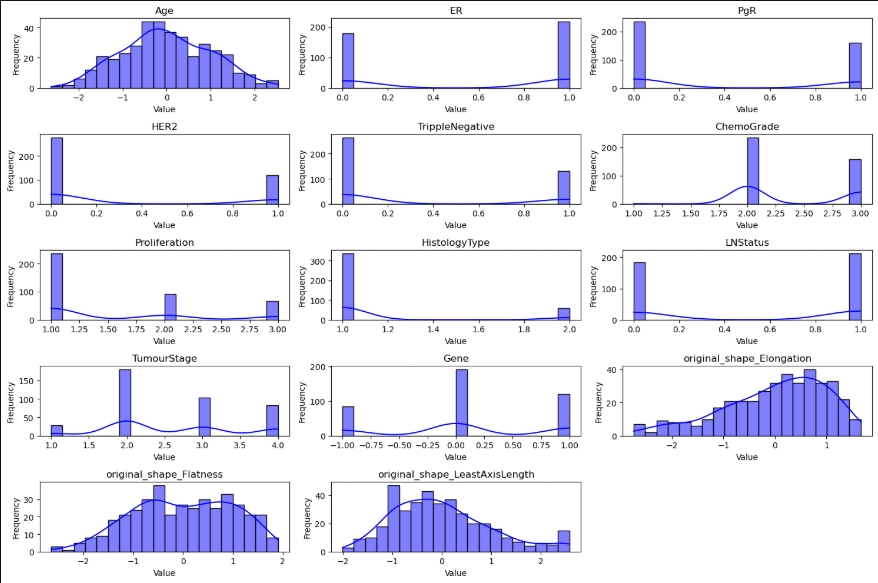


figure 3 Data histogram

Then, through data processing operations, the ID column is removed and the target column (PCR column and RFS column) is separated.

##### 2.3 Data imbalance issue

According to the analysis of histograms and boxplots, there may be a class imbalance in PCR (classification tasks). Workarounds include oversampling (such as SMOTE), undersampling, or adjusting how the model is trained through category weights. However, after comparing the two models and artificial parameters, it was found that SMOTENC performed the best.

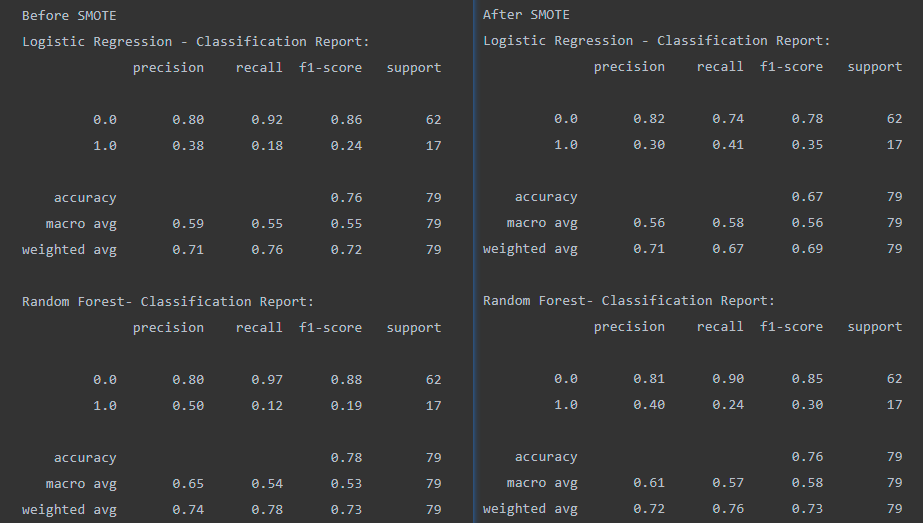


figure 4 SMOTE-Compare

#### 3. MODEL SELECTION AND TRAINING

##### 3.1 PCR (classification task)

Logistic Regression: Based on its efficiency and ease of interpretation for binary classification tasks.

Random Forest: Processes high-dimensional data and provides feature importance assessment.

##### 3.2 RFS (regression task)

Linear Regression: Used as a baseline model for comparison.

Random Forest: Processes high-dimensional data.

#### 4. MODEL EVALUATION

The first paragraph in each section should not be indented, but all following paragraphs within the section should be indented as these paragraphs demonstrate.

#### 5. CONCLUSION

Major headings, for example, “1. Introduction”, should appear in all capital letters, bold face if possible, centered in the column, with one blank line before, and one blank line after. Use a period (“.”) after the heading number, not a colon.

**11. References**

List and number all bibliographical references at the end of the paper. The references can be numbered in alphabetic order or in order of appearance in the document. When referring to them in the text, type the corresponding reference number in square brackets as shown at the end of this sentence [2]. An additional final page (the fifth page, in most cases) is allowed, but must contain only references to the prior literature.

[1] A.B. Smith, C.D. Jones, and E.F. Roberts, “Article Title,” *Journal*, Publisher, Location, pp. 1-10, Date.

[2] Jones, C.D., A.B. Smith, and E.F. Roberts, *Book Title*, Publisher, Location, Date.

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| **Task and Weighting** | **Data pre-processing (10%)** | **Feature**  **Selection**  **(25%)** | **ML method**  **development**  **(25%)** | **Method**  **Evaluation**  **(10%)** | **Report**  **Writing**  **(30%)** |
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