Laboratory Based Datasets

Report

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# **Albert Einstein Hospital, Brazil Dataset**

This dataset contains anonymized data from patients visited at the Hospital Israelita Albert Einstein in So Paulo, Brazil, who had samples collected to perform the SARS-CoV-2 RT-PCR and other laboratory tests during their visit.

All data were anonymized in accordance with international best practices and standards.

## **Our Modeling**

The Albert Einstein Hospital dataset consisted of 5644 samples and, 111 variables. The variables are divided into 4 dependent variables and 107 independent variables, the dependent variables are ‘SARS-Cov-2 exam result’, ‘Patient admitted to regular ward’, ‘Patient admitted to semi-intensive unit’ and ‘Patient admitted to intensive care unit’. In our implementation we only used 1 dependent variable ‘SARS-Cov-2 exam result’ to predict COVID-19 infection.

Points analyzed from the dataset:

* Large number of missing values.
* Most features are Imbalanced
* Imbalanced positive and negative cases.

**Missing Values**

Other than ‘Patient Age quantile’ and ‘SARS-Cov-2 exam result’, all features have missing values. The least feature with missing values is 76.01% empty, that is 1354 out of 5644. See Figure 1 for a chart of how much values are there in each feature. Our approach into dealing with missing values, 1) we first shrunk the dataset approximately to the least feature with missing values ‘Parainfluenza 1’ which results into 1352 samples dataset. Figure 2 presents the new features value count. 2) then we dropped features with more than 75% missing values and deleted 1 feature with only 1 unique value (zero-variance feature), which leaves us a new dataset with 1352 samples and 31 features. 15 of the 31 features are categorical and have no missing values. Also, we transformed categorical features using dummy encoding, and standardized all features to have a mean of 0 and a variance of 1. 3) finally, we used KNN imputation with k=2 to fill the missing values.

**Imbalanced features**

As for imbalanced features, we in this implementation didn’t use any techniques regarding this issue. But as illustration of how the features are imbalanced, we created Table 1 for categorical features. As for numeric features they have a strange value distribution for instance the ‘Red blood cell distribution width (RDW)’ feature has 55 different values and, from the 1352 samples 986 of them are for one value, this issue is present in all of the numeric features except ‘Patient age quantile’ which has a fair value distribution.

**Imbalanced Target Variable**

The shape of the dataset after the above preprocessing is now 1352 samples and 47 features. The target variable has a class distribution of 1240 negative samples and 112 positive samples, which is very imbalanced, and training a model on a highly imbalanced set will only lead to a model that predicts the minority class poorly or simply a biased-model.

So, in this implementation we created 11 splits of the dataset every set has all the 112 positive samples and a unique 112 negative samples. We ran 7 classifiers and used 10-fold cross validation to test and train the models. Each model is trained and tested on each split of the dataset, and every split produces 10 results using 10-fold cross validation, from the 10 results we take the average which then gives us the average result for one split, this is done for every split which means 11 results will be there, we take the average of the 11 splits to produce the finale result for a specific classifier. See Table 2 for models results.

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Figure 1: Bar chart showcasing features value count.

Chart

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Figure 2: Bar chart of the features value count after shrinking the dataset.

Table 1: Summary of features and their value distribution.

|  |  |  |
| --- | --- | --- |
| Feature | Detected | Not detected |
| Respiratory Syncytial Virus | 52 (3.8%) | 1300 (96.2%) |
| Influenza A | 18 (1.3%) | 1334 (98.7%) |
| Influenza B | 76 (5.6%) | 1276 (94.4%) |
| Parainfluenza 1 | 3 (0.2%) | 1349 (99.8%) |
| CoronavirusNL63 | 45 (3.3%) | 1307 (96.7%) |
| Rhinovirus/Enterovirus | 379 (28%) | 973 (72%) |
| Coronavirus HKU1 | 20 (1.5%) | 1332 (98.5%) |
| Parainfluenza 3 | 10 (0.7%) | 1342 (99.3%) |
| Chlamydophila pneumoniae | 9 (0.7%) | 1343 (99.3%) |
| Adenovirus | 13 (1%) | 1339 (99%) |
| Parainfluenza 4 | 19 (1.4%) | 1333 (98.6%) |
| Coronavirus229E | 9 (0.7%) | 1343 (99.3%) |
| CoronavirusOC43 | 8 (0.6%) | 1344 (99.4%) |
| Inf A H1N1 2009 | 98 (7.2%) | 1254 (92.8%) |
| Bordetella pertussis | 2 (0.2%) | 1350 (99.8%) |
| Metapneumovirus | 14 (1%) | 1338 (99%) |

Table 2: Models performance

|  |  |  |  |
| --- | --- | --- | --- |
| Model | Average Precision | Average Recall | Average F1-Score |
| Logistic Regression (LR) | 79% | 76% | 75% |
| Support Vector Machine (SVM) | 81% | 77% | 77% |
| K-Nearest Neighbors (KNN) K=3 | 76% | 74% | 73% |
| Gaussian Naive Bayes (GNB) | 76% | 60% | 53% |
| Decision Tree (DT) | 74% | 72% | 72% |
| Random Forest (RF) | 78% | 75% | 75% |
| Extreme Gradient Boosting (XGBoost) | 78% | 76% | 75% |

Table 3 below presents some results of related work done using the same dataset, compared with ours.

Table 3: Comparison of our model with other existing models.

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| Reference | Features | Models used | Precision | Recall | F1-Score |
| [1] | 25 | Stacked Ensemble Machine Learning (See Table 4) | 100% | 100% | 100% |
| [2] | 24 | SVM, MPL, RF,  Naive Bayes,  Bayesian network | 93.77% | 96.76% | 93.80% |
| [3] | 14 | CR | 83.70% | 84.20% | 83.70% |
| Couldn’t find the paper, but was cited by [1] | 15 | SVM, RF, DT | - | - | 92.00% |
| Our model | 31 | LR, SVM, KNN, GNB, DT, RF, XGB | 78% | 74% | 73% |

Table 4: Performance Metrics of Stacked Ensemble Model. [1]

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
|  | Layer-1 | | | | Layer-2 |
| Performance Metrics | KNN | SVM | XGB | RF | AdB |
| Precision | 98.73% | 96.84% | 97.47% | 99.37% | 100% |
| Recall | 77.23% | 90.00% | 93.33% | 95.15% | 100% |
| F1-Score | 86.67% | 93.29% | 95.36% | 97.21% | 100% |

# **Italian Society of Medical Radiology Dataset**

This dataset is composed of 68 COVID-19 cases from the Italian society of medical and intervention radiology society (SIRM) database, and 62 Flu cases from the influenza research database (IRD).

## **Our Modeling**

The SIRM Dataset has 130 samples and 18 variables, 1 dependent variable ‘Decision label’ and 16 independent variables if you exclude sample id ‘Number’. Out of the 130 samples 68 of them COVID-19 positive and the remaining 62 are Flu positive, so we set the 68 COVID-19 cases as positive or ‘1’ and the 62 Flu cases as negative or ‘0’.

The people who constructed the dataset assigned missing values with a star symbol ‘\*’ instead of leaving it empty. To see how much missing values, we are dealing with we replaced every ‘\*’ with a NaN value. Figure 3 presents a bar chart of variables value count.

Features with more than 50% missing values were dropped, also 1 feature ‘High risk zone’ which has only 1 unique value and therefore dropped. The remaining are 8 features, see Figure 4 for the updated features value count.

We transformed categorical features into numerical features and standardized all features to have a mean of 0 and a variance of 1. Finally, we used Simple Imputer with a mean strategy to fill the missing values.

We ran 7 models and used 5-fold cross validation to test and train the models, Table 5 presents the results.

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Figure 3: Bar chart of features value count.

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Figure 4: Bar chart of the updated features value count.

Table 5: Model Performance.

|  |  |  |  |
| --- | --- | --- | --- |
| Model | Precision | Recall | F1-Score |
| Logistic Regression (LR) | 86% | 87% | 84% |
| Support Vector Machine (SVM) | 93% | 95% | 93% |
| K-Nearest Neighbors (KNN) K=6 | 87% | 89% | 86% |
| Gaussian Naive Bayes (GNB) | 90% | 92% | 89% |
| Decision Tree (DT) | 97% | 97% | 97% |
| Random Forest (RF) | 100% | 100% | 100% |
| Extreme Gradient Boosting (XGBoost) | 100% | 100% | 100% |

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