

Machine Learning for Health Care

**PROJECT 1**

**Interpretable and Explainable Classification**

**for Medical Data**

**Students**

Teodora Petrovic, 23-745-797

Rohit Koonireddy, 20-622-924

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# Heart Disease Prediction Dataset

## Dataset Description

The dataset comprises the following attributes for each individual:

* Age: The age of the individual.
* Sex: The gender of the individual (Male or Female).
* Chest Pain Type: The type of chest pain experienced by the individual.
* Resting Blood Pressure: The individual's blood pressure at rest.
* Cholesterol: The cholesterol level of the individual.
* Fasting Blood Sugar: The fasting blood sugar level of the individual.
* Resting Electrocardiographic Results: Results of the resting electrocardiogram.
* Maximum Heart Rate: The maximum heart rate achieved during exercise.
* Exercise-Induced Angina: Whether the individual experienced angina during exercise.
* ST Depression: ST depression induced by exercise relative to rest.
* ST Slope: The slope of the peak exercise ST segment.
* Heart Disease: The presence or absence of heart disease (1 indicates presence, 0 indicates absence).

## Exploratory Data Analysis

Exploratory Data Analysis (EDA) was conducted to understand the characteristics of the heart disease prediction dataset. Various visualizations, including violin plots and histograms, were used to explore the distribution of features and their relationships with the target variable.

Sample view of the dataset:

A screenshot of a computer

Description automatically generated

Simple describe view of the data:

A screenshot of a computer screen

Description automatically generated

Sample Violon chart:

A diagram of a kayak

Description automatically generated with medium confidence

## Logistic Lasso Regression

We trained a logistic regression model to predict the presence of heart disease using the provided dataset. First, we split the data into features and target variables, encoding categorical features and dividing the dataset into training and testing sets. Next, the logistic regression model was fitted to the training data. Subsequently, we used the trained model to predict heart disease presence on the test set. Finally, we evaluated the model's performance using accuracy and F1-score metrics, assessing the proportion of correct predictions and the balance between precision and recall, respectively.

A graph of blue rectangular shapes

Description automatically generated with medium confidence

## Multi-Layer Perceptrons

Multi-Layer Perceptrons (MLPs), a class of feedforward neural networks, were employed for heart disease prediction. MLPs are capable of learning complex relationships between features and the target variable. The model's hyperparameters, such as the number of hidden layers and neurons, activation functions, and optimization algorithms, were fine-tuned to achieve optimal performance. Evaluation metrics such as accuracy and F1-score were used to assess the model's predictive capability.

## Neural Additive Models

Neural Additive Models (NAMs) offer an interpretable framework for understanding the predictions of complex neural network models. By decomposing the model predictions into additive components, NAMs provide insights into the contribution of each feature to the overall prediction. Shapely values, a popular technique for interpreting machine learning models, were employed to quantify the impact of individual features on the model predictions. Summary plots generated from Shapely values help visualize feature importance and aid in understanding the model's behavior.

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# Pneumonia Prediction Dataset

## Exploratory Data Analysis

### Label distribution and qualitative description of the data

The dataset is structured into three main folders: train, val and test. Within each of these folders, there are subfolders representing the two categories of X-Ray images: Pneumonia and Normal. In total, there are 5,856 X-Ray images in JPEG format. Their distribution is shown in table 1.

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| Class | Train | Validation | Test | Total |
| Normal | 1341 | 8 | 234 | 1583 |
| Pneumonia | 3875 | 8 | 390 | 4273 |

*Table 1. Label distribution*

A close-up of x-ray images

Description automatically generatedThe normal chest X-ray typically shows clear lungs without any areas of abnormal opacification. However, in cases of bacterial pneumonia, distinct white areas may appear on the image, indicating focal lobar consolidation, as illustrated in Figure 1. These differences can be discovered even by an untrained observer.

*Figure 1. Distinct white areas are marked*

A close-up of an x-ray of a chest

Description automatically generatedIt is also easy to notice differences if pneumonia has progressed significantly, as depicted in Figure 2.

*Figure 2. Disease progressed significantly*

A x-ray of a person's chest

Description automatically generatedOn the other hand, distinguishing between healthy and diseased samples is often more challenging in cases of pneumonia in early stages. This is because viral pneumonia is often manifested with a more diffuse ‘‘interstitial’’ pattern in both lungs, which can closely resemble the appearance of a healthy chest X-ray. I was not able to see these differences. Less pronounced visual differences between healthy and disease samples are shown in Figure 3.

*Figure 3. Samples that are dificult to classify by untrained eye*

However, in most cases it is very difficult for someone who is not an expert in the field of medicine to see visual differences between healthy and disease samples as well as differences between virus and bacterial pneumonia.

### Source of bias that could influence model performance

There are significantly more pneumonia samples than healthy samples, as shown in Table 1. The potential source of bias that could influence model performance would be bias towards the majority class (pneumonia class). In a classification task with imbalanced classes, a model might tend to predict the majority class more frequently due to its prevalence in the dataset. Consequently, the model may have lower sensitivity or recall for the minority class (normal images) and higher false negative rates for detecting normal cases.

### Data preprocessing

The data has undergone cropping, resizing, and normalization. This was executed with the aid of torchvision library.

* torchvision.transforms.Resize
* torchvision.transforms.CenterCrop
* torchvision.transforms.Normalize

To address bias explained earlier, we utilized data augmentation techniques. Data augmentation is creating additional training examples by applying transformations to the existing data. This can help us balance the dataset and improve model generalization. We implemented intensity transformations (sharpness adjustments, random noise) and positional transformations. We applied transformations by using functionalities from torchvision library:

* torchvision.transforms.RandomApply
* torchvision.transforms.RandomAffine
* torchvision.transforms.GaussianBlur
* torchvision.transforms.RandomHorizontalFlip
* torchvision.transforms.RandomVerticalFlip
* torchvision. transforms.RandomAdjustSharpness

A computer screen shot of code

Description automatically generatedImplementation of these transformations can be seen in Figure 4.

*Figure 4. Transforming images*

A computer code with numbers and symbols

Description automatically generatedPart of the code in which we add noise to the image is shown in Figure 5.

*Figure 5. Adding noise to the image*

A close-up of a radiograph

Description automatically generatedFigure 6 shows one image after cropping, then after resizing to (3, 222, 224), and after all other transformations.

*Figure 6. Image preprocessing and augmentations*

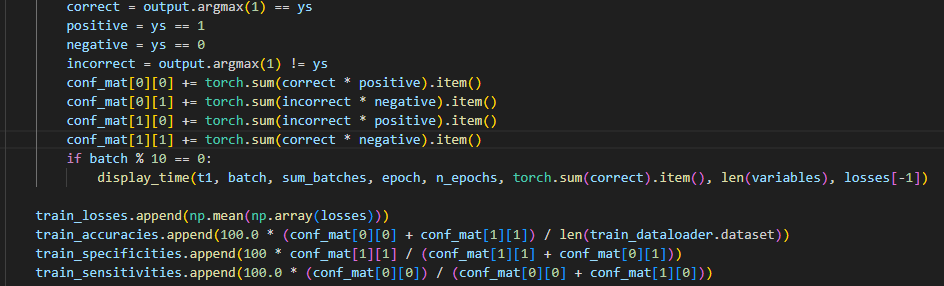
## CNN Classifier

### Network architecture and implementation

A computer screen with text

Description automatically generatedWe are using ResNet18, which was pretrained on ImageNet dataset. In python script dataset.py, in def \_\_getitem\_\_(self, index) we are processing images. We load one image, do the preprocessing and transformations and return it together with its label (0 for healty sample, 1 for pneumonia sample). File 2d\_test.py is the heart of task 2 and used to split data into train and validation data and call main function. Test data is unchanged. Training is done in epochs, and data is divided in batches. We are making predictions if the person has disease or not, calculate loss, back propagate it and update the weights. This is shown in Figure 7.

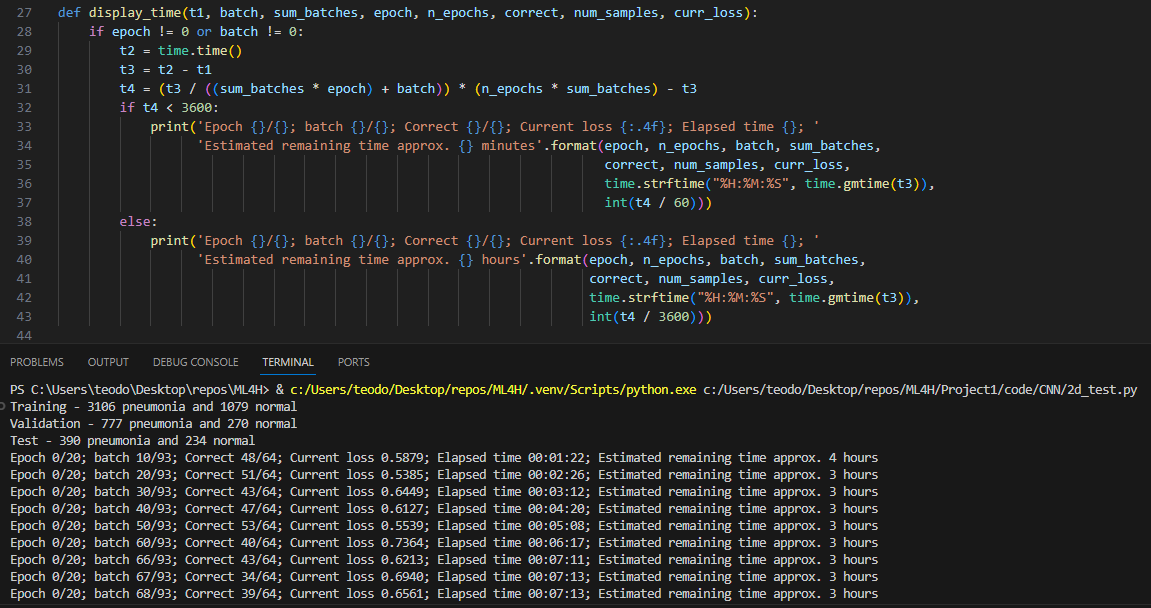
*Figure 7. Epoch and time tracking*

After every epoch we are appending loss, accuracy, specificity and sensitivity using the coeficiency matrix. We do this for train, validation and test. This is shown in figure 8.

*Figure 8. Calculating accuracy, specificity and sensitivity*

Then, we plot the results using code from Figure 9.

*Figure 9. Plotting the results*

To keep track of time and progress, we implemented function shown in Figure 10.

*Figure 10. Printing progress and time*

### Results

CNN performed best with parameters shown in table 2 and achieved results presented in table 3.

|  |  |  |
| --- | --- | --- |
|  | Unbalanced dataset | Balanced dataset |
| number of epochs | 20 | 20 |
| batch size | 64 | 64 |
| learning rate | 0.0001 | 0.0001 |
| scheduler step size | 10 | 10 |
| scheduler gama | 0.33 | 0.33 |
| optimization function | sgd | sgd |
| loss function | cross entropy | cross entropy |
| weight decay | 0.0005 | 0.0005 |

*Table 2. Best parameters for training the CNN*

|  |  |  |
| --- | --- | --- |
|  | Unbalanced dataset | Balanced dataset |
| Accuracy |  |  |
| Specificity |  |  |
| Sensitivity |  |  |

*Table 3. Best results*

We have calculated the performance using formulas (1), (2) and (3).

|  |  |
| --- | --- |
|  | (1) |
|  | (2) |
|  | (3) |

A graph showing the growth of the number of countries/regions

Description automatically generated with medium confidenceLoss function on train and validation data is given in Figure 7.

*Figure 7. Loss function*

A graph of different types of graphs

Description automatically generated with medium confidencePerformance parameters, such as accuracy, specificity and sensitivity during epochs is shown in Figure 8.

*Figure 8. accuracy, specificity and sensitivity during epochs*

## Integrated Gradients

## Grad-CAM

## Data Randomization Test

A close-up of x-ray images

Description automatically generated

*Figure 9. View for Integrated Gradients approach with 70% target randomization*

# References