

Configuration Manual of Classification models for Improving Identification of heart diseases

MSc Research Project
MSc. in Data Analytics

Omkar Terdal Ramesh
Student ID: x16104439

School of Computing
National College of Ireland

Supervisor: Dr.Catherine Mulwa

National College of Ireland
Project Submission Sheet
School of Computing



Student Name:	Omkar Terdal Ramesh
Student ID:	x16104439
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Configuration Manual of Classification models for Improving Identification of heart diseases

Omkar Terdal Ramesh
x16104439

1 Introduction

Our Project "Classification Models for Improving Identification of Heart Diseases in Healthcare Industry : Eastern Europe" is about creating model for improving identification of heart diseases using data collected from a repository and showcase its severity level using Machine learning algorithms and Artificial Neural network over a train and test dataset. This configuration manual accompanies the Project report and helps in understanding the configuration process for the duration of the project.

2 Environment configuration

We have used python through jupyter notebook to execute our project. As the data was collected in a .csv file, we stored the data onto the system as the jupyter notebook can access files directly on the system and can the software on the system itself. We are using Anaconda v2.2019.10.

2.1 Anaconda

Here are the steps to download Anaconda v2.2019.10. We can access the installer from the following link <https://docs.anaconda.com/anaconda/install/hashes/win-2-64/> and we can select the version we want to download.

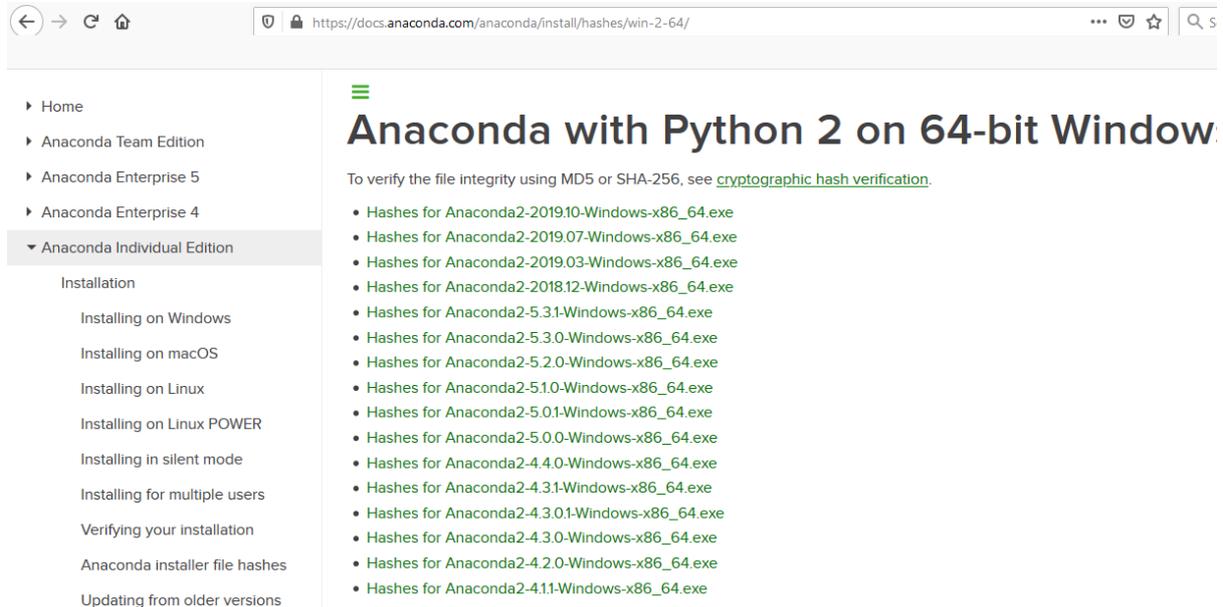


Figure 1: Anaconda Download site

2.2 Local system specifications

We used the local system to make a remote connection to the jupyter notebook. The laptop used is a Asus GL702VM and its specifications are shown below in Figure 2

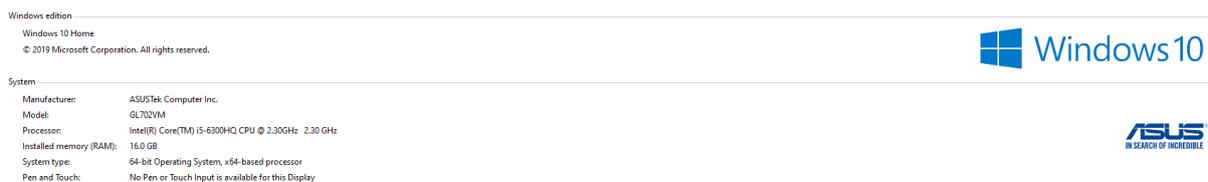


Figure 2: Local system specifications

2.3 Python Libraries

We have installed the enlisted python libraries with corresponding versions as shown below in the Table.

Libraries	Versions
numpy (numpy, 2020)	1.18.2
Pandas (Pands, 2020)	1.0.3
Matplotlib (matplotlib, 2020)	3.2.1
cv2	4.1.2
sklearn 1.8.0	(scikit, 2020)
tensorflow (tensorflow, 2020)	2.2.0-rc3

3 Data Collection

Shown Below is a part of our dataset which is stored locally on the system. The dataset has 16 attributes and over 4000 rows. They are sufficient enough to run this project. In the the below shown Figure 3 you can see the dataset.

A	B	C	D	E	F	G	H	I	J	K	L	M	N	O	P
gender	age	education	currentSmoker	cigsPerDay	BPMeds	prevalentStroke	prevalentHyp	diabetes	totChol	sysBP	diaBP	BMI	heartRate	glucose	Label
1	39	4	0	0	0	0	0	0	195	106	70	26.97	80	77	0
0	46	2	0	0	0	0	0	0	250	121	81	28.73	95	76	0
1	48	1	1	20	0	0	0	0	245	127.5	80	25.34	75	70	0
0	61	3	1	30	0	0	0	1	225	150	95	28.58	65	103	1
0	46	3	1	23	0	0	0	0	285	130	84	23.1	85	85	0
0	43	2	0	0	0	0	0	1	228	180	110	30.3	77	99	0
0	63	1	0	0	0	0	0	0	205	138	71	33.11	60	85	1
0	45	2	1	20	0	0	0	0	313	100	71	21.68	79	78	0
1	52	1	0	0	0	0	0	1	260	141.5	89	26.36	76	79	0
1	43	1	1	30	0	0	0	1	225	162	107	23.61	93	88	0
0	50	1	0	0	0	0	0	0	254	133	76	22.91	75	76	0
0	43	2	0	0	0	0	0	0	247	131	88	27.64	72	61	0
1	46	1	1	15	0	0	0	1	294	142	94	26.31	98	64	0
0	41	3	0	0	1	0	0	1	332	124	88	31.31	65	84	0
0	39	2	1	9	0	0	0	0	226	114	64	22.35	85	NA	0
0	38	2	1	20	0	0	0	1	221	140	90	21.35	95	70	1
1	48	3	1	10	0	0	0	1	232	138	90	22.37	64	72	0
0	46	2	1	20	0	0	0	0	291	112	78	23.38	80	89	1
0	38	2	1	5	0	0	0	0	195	122	84.5	23.24	75	78	0
1	41	2	0	0	0	0	0	0	195	139	88	26.88	85	65	0
0	42	2	1	30	0	0	0	0	190	108	70.5	21.59	72	85	0
0	43	1	0	0	0	0	0	0	185	123.5	77.5	29.89	70	NA	0
0	52	1	0	0	0	0	0	0	234	148	78	34.17	70	113	0
0	52	3	1	20	0	0	0	0	215	132	82	25.11	71	75	0
1	44	2	1	30	0	0	0	1	270	137.5	90	21.96	75	83	0
1	47	4	1	20	0	0	0	0	294	102	68	24.18	62	66	1
0	60	1	0	0	0	0	0	0	260	110	72.5	26.59	65	NA	0
1	35	2	1	20	0	0	0	1	225	132	91	26.09	73	83	0
0	61	3	0	0	0	0	0	1	272	182	121	32.8	85	65	1
0	60	1	0	0	0	0	0	0	247	130	88	30.36	72	74	0
1	36	4	1	35	0	0	0	0	295	102	68	28.15	60	63	0
1	43	4	1	43	0	0	0	0	226	115	85.5	27.57	75	75	0
0	59	1	0	0	0	0	0	1	209	150	85	20.77	90	88	1
1	61	NA	1	5	0	0	0	0	175	134	82.5	18.59	72	75	1
1	54	1	1	20	0	0	0	1	214	147	74	24.71	96	87	0
1	37	2	0	0	0	0	0	1	225	124.5	92.5	38.53	95	83	0
1	56	NA	0	0	0	0	0	0	257	153.5	102	28.09	72	75	0

Figure 3: Dataset and Attributes

3.1 Data Storage

Shown in Figure 4 below is the path where the collected data is stored for the project. The data is stored in the datasets folder to access the data through the Jupyter notebook. This is the installation folder for Anaconda.

Name	Date modified	Type	Size
.ipynb_checkpoints	15-08-2020 15:13	File folder	
Datasets	12-06-2020 11:14	File folder	
model	12-06-2020 11:14	File folder	
test_sample	12-06-2020 11:14	File folder	
Traning_Testing	12-06-2020 11:14	File folder	
Anaconda.bat	05-08-2020 11:41	Windows Batch File	1 KB
Anaconda.txt	05-08-2020 11:40	Text Document	1 KB
Heart_Disease_Using_ML.ipynb	14-08-2020 20:36	IPYNB File	175 KB
Heart_Disease_Using_ML-Copy1.ipynb	16-08-2020 11:43	IPYNB File	184 KB
Heart_Disease_Using_ML-Copy2.ipynb	15-08-2020 15:13	IPYNB File	183 KB
Test.ipynb	31-07-2020 11:28	IPYNB File	14 KB

Figure 4: Dataset and Attributes

4 Jupyter notebook

In this section we will run jupyter notebook with the help of anaconda. The below Figure shows that in order to run jupyter notebook, we need to first launch the command prompt from the same directory. And the run the anaconda batch file to load the required libraries into jupyter notebook.

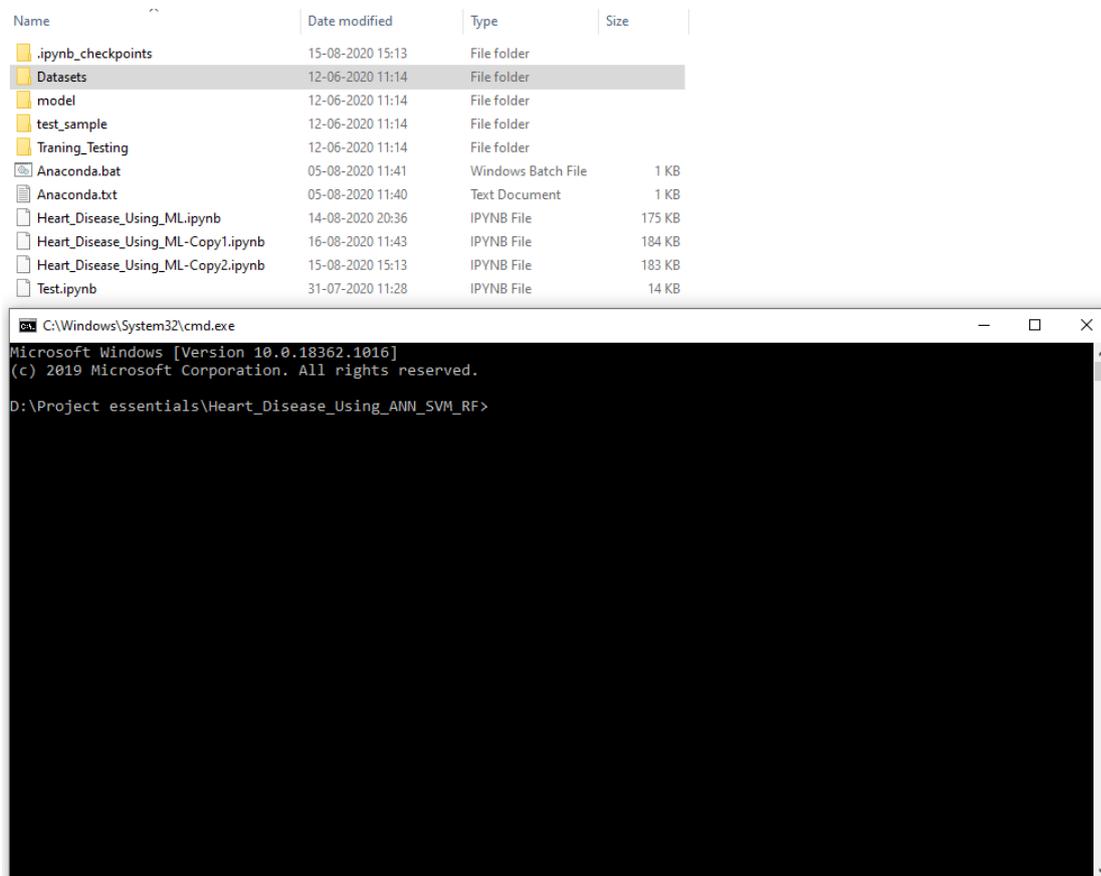


Figure 5: Command prompt

We then launch jupyter notebook after the libraries have been loaded.

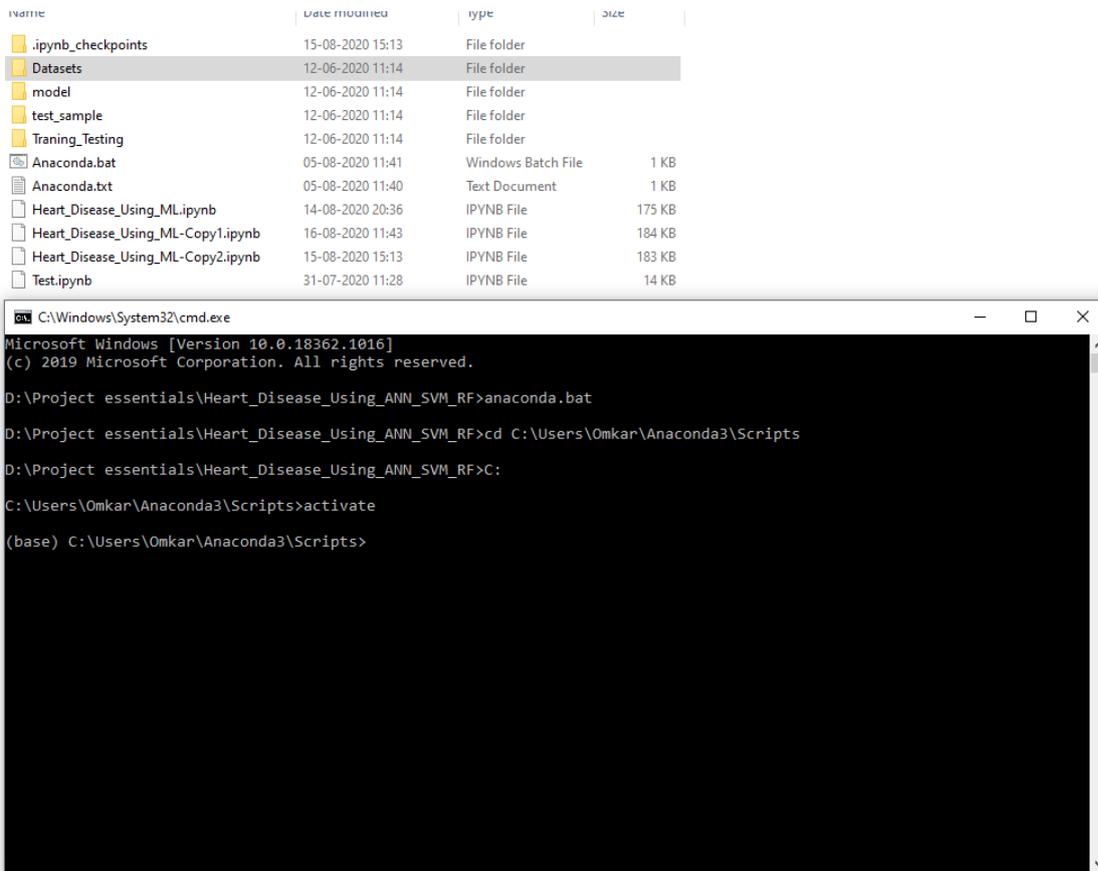


Figure 6: Launching Jupyter notebook

After the Jupyter notebook is launched it establishes a connection to the directory and opens in the default web browser with the files from the directory.

Name	Date modified	Type	Size
.ipynb_checkpoints	15-08-2020 15:13	File folder	
Datasets	12-06-2020 11:14	File folder	
model	12-06-2020 11:14	File folder	
test_sample	12-06-2020 11:14	File folder	
Traning_Testing	12-06-2020 11:14	File folder	
Anaconda.bat	05-08-2020 11:41	Windows Batch File	1 KB
Anaconda.txt	05-08-2020 11:40	Text Document	1 KB
Heart_Disease_Using_ML.ipynb	14-08-2020 20:36	IPYNB File	175 KB
Heart_Disease_Using_ML-Copy1.ipynb	16-08-2020 11:43	IPYNB File	184 KB
Heart_Disease_Using_ML-Copy2.ipynb	15-08-2020 15:13	IPYNB File	183 KB
Test.ipynb	31-07-2020 11:28	IPYNB File	14 KB

```

C:\Windows\System32\cmd.exe
Microsoft Windows [Version 10.0.18362.1016]
(c) 2019 Microsoft Corporation. All rights reserved.

D:\Project essentials\Heart_Disease_Using_ANN_SVM_RF>anaconda.bat

D:\Project essentials\Heart_Disease_Using_ANN_SVM_RF>cd C:\Users\Omkar\Anaconda3\Scripts

D:\Project essentials\Heart_Disease_Using_ANN_SVM_RF>C:

C:\Users\Omkar\Anaconda3\Scripts>activate

(base) C:\Users\Omkar\Anaconda3\Scripts>D:

(base) D:\Project essentials\Heart_Disease_Using_ANN_SVM_RF>jupyter notebook

```

Figure 7: Loading Jupyter notebook

5 Implementation

We go ahead and import all the necessary libraries required for our project. The Figure 9 below shows the libraries imported.

We go ahead and load the data into the Jupyter notebook by using the read command. The Figure 10 shows how.

```

C:\Windows\System32\cmd.exe - jupyter notebook
(c) 2019 Microsoft Corporation. All rights reserved.
D:\Project essentials\Heart_Disease_Using_ANN_SVM_RF>anaconda.bat
D:\Project essentials\Heart_Disease_Using_ANN_SVM_RF>cd C:\Users\Omkar\Anaconda3\Scripts
D:\Project essentials\Heart_Disease_Using_ANN_SVM_RF>C:
C:\Users\Omkar\Anaconda3\Scripts>activate
(base) C:\Users\Omkar\Anaconda3\Scripts>D:
(base) D:\Project essentials\Heart_Disease_Using_ANN_SVM_RF>jupyter notebook
[I 02:30:40.234 NotebookApp] The port 8888 is already in use, trying another port.
[I 02:30:40.365 NotebookApp] JupyterLab extension loaded from C:\Users\Omkar\Anaconda3\lib\site-packages\jupyterlab
[I 02:30:40.365 NotebookApp] JupyterLab application directory is C:\Users\Omkar\Anaconda3\share\jupyter\lab
[I 02:30:40.369 NotebookApp] Serving notebooks from local directory: D:\Project essentials\Heart_Disease_Using_ANN_SVM_R
F
[I 02:30:40.369 NotebookApp] The Jupyter Notebook is running at:
[I 02:30:40.369 NotebookApp] http://localhost:8889/?token=5c1d4505739736c40a9d9a06a14a3043ee250c0475188436
[I 02:30:40.369 NotebookApp] or http://127.0.0.1:8889/?token=5c1d4505739736c40a9d9a06a14a3043ee250c0475188436
[I 02:30:40.370 NotebookApp] Use Control-C to stop this server and shut down all kernels (twice to skip confirmation).
[C 02:30:40.469 NotebookApp]

To access the notebook, open this file in a browser:
file:///C:/Users/Omkar/AppData/Roaming/jupyter/runtime/nbserver-12964-open.html
Or copy and paste one of these URLs:
http://localhost:8889/?token=5c1d4505739736c40a9d9a06a14a3043ee250c0475188436
or http://127.0.0.1:8889/?token=5c1d4505739736c40a9d9a06a14a3043ee250c0475188436

```

Figure 8: Jupyter Notebook

The screenshot shows the Jupyter Notebook interface. The title bar reads "jupyter Heart_Disease_Using_ML-Copy1 Last Checkpoint: 08/05/2020 (autosaved)". The menu bar includes File, Edit, View, Insert, Cell, Kernel, Widgets, and Help. The toolbar contains icons for file operations, zooming, and running code. The main area shows a code cell with the following content:

```

import lib

In [1]: import pandas as pd
import numpy as np
import itertools
from sklearn import preprocessing
from sklearn.model_selection import train_test_split
import matplotlib.pyplot as plt
import seaborn as sns
from sklearn.metrics import classification_report, confusion_matrix, accuracy_score
import pickle

```

Figure 9: Jupyter Libraries

5.1 Implementation of ANN

We will first divide the data into training data and test data. This is ensure that the models can perform and hope to achieve the maximum efficiency. Figure 11

5.1.1 Data preprocessing

We preprocess the data as required by the ANN model by using keras and converting the data into arrays of data. The following figure 12 demonstrates it.

5.1.2 Building and training ANN

We will build a sequential model, compile the model and Fit the model to run through epochs. This model runs successfully through a 100 epochs and yields a result of 85.61 accuracy.

load data

```
In [2]: df = pd.read_csv('Datasets/Heart_Data.csv')
In [3]: df.head()
Out[3]:
```

	gender	age	education	currentSmoker	cigsPerDay	BPMeds	prevalentStroke	prevalentHyp	diabetes	totChol	sysBP	diaBP	BMI	heartRate	glucose	La
0	1	39	4.0	0	0.0	0.0	0	0	0	195.0	106.0	70.0	26.97	80.0	77.0	
1	0	46	2.0	0	0.0	0.0	0	0	0	250.0	121.0	81.0	28.73	95.0	76.0	
2	1	48	1.0	1	20.0	0.0	0	0	0	245.0	127.5	80.0	25.34	75.0	70.0	
3	0	61	3.0	1	30.0	0.0	0	1	0	225.0	150.0	95.0	28.58	65.0	103.0	
4	0	46	3.0	1	23.0	0.0	0	0	0	285.0	130.0	84.0	23.10	85.0	85.0	

```
< >
In [4]: df.columns
Out[4]: Index(['gender', 'age', 'education', 'currentSmoker', 'cigsPerDay', 'BPMeds',
              'prevalentStroke', 'prevalentHyp', 'diabetes', 'totChol', 'sysBP',
              'diaBP', 'BMI', 'heartRate', 'glucose', 'Label'],
              dtype='object')
```

Figure 10: Loading Data into Jupyter notebook

ANN

```
In [5]: df.dropna(axis=0,inplace=True)
In [6]: dataset = df.values
In [7]: dataset
Out[7]: array([[ 1., 39., 4., ..., 80., 77., 0.],
               [ 0., 46., 2., ..., 95., 76., 0.],
               [ 1., 48., 1., ..., 75., 70., 0.],
               ...,
               [ 0., 52., 2., ..., 80., 107., 0.],
               [ 1., 40., 3., ..., 67., 72., 0.],
               [ 0., 39., 3., ..., 85., 80., 0.]])
In [8]: X = dataset[:,0:15]
         print(X)
         Y = dataset[:,15]
[[ 1.  39.  4.  ... 26.97 80.  77. ]
 [ 0.  46.  2.  ... 28.73 95.  76. ]
 [ 1.  48.  1.  ... 25.34 75.  70. ]
 ...
 [ 0.  52.  2.  ... 21.47 80. 107. ]
 [ 1.  40.  3.  ... 25.6  67.  72. ]
 [ 0.  39.  3.  ... 20.91 85.  80. ]]
```

Figure 11: Data Distribution to training and test data

5.1.3 Visualization

The Following plot graph shows the loss during epochs (Figure 15). And followed by that we can see the Accuracy graph of the model displayed with the help of the plot command (Figure 16)

5.1.4 Conclusion

The following figure 17 shows the accuracy, precision, recall, f1-score and support of the ANN model.

preprocessing

```
In [9]: from keras.models import Sequential
        from keras.layers import Dense, InputLayer, Flatten, BatchNormalization

        Using TensorFlow backend.

In [10]: min_max_scaler = preprocessing.MinMaxScaler()
         X_scale = min_max_scaler.fit_transform(X)

In [11]: X_scale
Out[11]: array([[1.          , 0.18421053, 1.          , ..., 0.27702375, 0.36363636,
                0.10451977],
                [0.          , 0.36842105, 0.33333333, ..., 0.31968008, 0.51515152,
                0.10169492],
                [1.          , 0.42105263, 0.          , ..., 0.23751818, 0.31313131,
                0.08474576],
                ...,
                [0.          , 0.52631579, 0.33333333, ..., 0.14372273, 0.36363636,
                0.18926554],
                [1.          , 0.21052632, 0.66666667, ..., 0.24381968, 0.23232323,
                0.09039548],
                [0.          , 0.18421053, 0.66666667, ..., 0.13015027, 0.41414141,
                0.11299435]])

In [12]: X_train, X_val_and_test, Y_train, Y_val_and_test = train_test_split(X_scale, Y, test_size=0.3)
         X_val, X_test, Y_val, Y_test = train_test_split(X_val_and_test, Y_val_and_test, test_size=0.5)
         print(X_train.shape, X_val.shape, X_test.shape, Y_train.shape, Y_val.shape, Y_test.shape)
(2560, 15) (549, 15) (549, 15) (2560,) (549,) (549,)
```

Figure 12: Data preprocessing for ANN

5.2 Implementation of Random Forest and SVM Models

5.2.1 Data preprocessing

We need to preprocess the data again in a different way to feed it to both the random forest and SVM models. These are machine learning algorithms and hence need to be preprocessed differently compared to Artificial Neural networks. Figure 18

We go ahead and plot the attributes of the dataset. This provides an idea on how to approach the Models for increased performance and accuracy. Figure 19

We consider the gender attribute to be labelled and plotted to see the number of entries they have and how we can make use of it. Figure 20

We then plot the gender compared to age to see calculate the age group for the most affected are. This figure 21 shows us that the most case received are between the ages of 39-48.

Keeping age groups and gender as our two main attributes for the models we will further train the data into training set and test set. As shown in the Figure 22.

5.2.2 Random Forest Model

We train the random forest model and input the training and test data. As shown in Figure 23. The Random Forest model yielded an accuracy of 85.06. With the precision being 0.85, recall being 1.00 and f1-score being 0.92. as shown in figure 24.

Building and Training Our Artificial Neural Network

```
In [13]: model = Sequential([
          Dense(32, activation='relu', input_shape=(15,)),
          Dense(32, activation='relu'),
          Dense(1, activation='sigmoid'),
        ])

WARNING:tensorflow:From C:\Users\Omkar\Anaconda3\lib\site-packages\keras\backend\tensorflow_backend.py:74: The name tf.get_default_graph is deprecated. Please use tf.compat.v1.get_default_graph instead.

WARNING:tensorflow:From C:\Users\Omkar\Anaconda3\lib\site-packages\keras\backend\tensorflow_backend.py:517: The name tf.placeholder is deprecated. Please use tf.compat.v1.placeholder instead.

WARNING:tensorflow:From C:\Users\Omkar\Anaconda3\lib\site-packages\keras\backend\tensorflow_backend.py:4138: The name tf.random_uniform is deprecated. Please use tf.random.uniform instead.

In [14]: model.compile(optimizer='sgd',
                      loss='binary_crossentropy',
                      metrics=['accuracy'])

WARNING:tensorflow:From C:\Users\Omkar\Anaconda3\lib\site-packages\keras\optimizers.py:790: The name tf.train.Optimizer is deprecated. Please use tf.compat.v1.train.Optimizer instead.

WARNING:tensorflow:From C:\Users\Omkar\Anaconda3\lib\site-packages\keras\backend\tensorflow_backend.py:3376: The name tf.log is deprecated. Please use tf.math.log instead.

WARNING:tensorflow:From C:\Users\Omkar\Anaconda3\lib\site-packages\tensorflow\python\ops\nn_impl.py:180: add_dispatch_support.<locals>.wrapper (from tensorflow.python.ops.array_ops) is deprecated and will be removed in a future version.
Instructions for updating:
Use tf.where in 2.0, which has the same broadcast rule as np.where
```

Figure 13: Training ANN

5.2.3 SVM Model

We train and fit the SVM model with the training and test data. The following Figure 25 shows how. The SVM model yielded an accuracy of 84.79. The precision 0.85, recall is 1.0, and f1-score is 0.92. As shown in Figure 26.

6 Comparison

Once we acquired all the accuracy from the models we ran, we felt the need to compare the results just to see how close or how far apart the artificial neural network techniques is from the machine learning techniques. The highest accuracy was yielded by ANN. Though the ANN technique's accuracy was a bit higher than other techniques, we can conclude that in the case of disease identification every small amount is considered and can come to great help. The following plot will show the difference in the accuracies. Figure 27

7 Section 6

Your sixth section. Change the header and label to something appropriate.

References

```

In [15]: hist = model.fit(X_train, Y_train,
                        batch_size=32, epochs=100,
                        validation_data=(X_val, Y_val))

WARNING:tensorflow:From C:\Users\Omkar\Anaconda3\lib\site-packages\keras\backend\tensorflow_backend.py:986: The name tf.assign_add is deprecated. Please use tf.compat.v1.assign_add instead.

Train on 2560 samples, validate on 549 samples
Epoch 1/100
2560/2560 [=====] - 0s 190us/step - loss: 0.6465 - acc: 0.6555 - val_loss: 0.5266 - val_acc: 0.8689
Epoch 2/100
2560/2560 [=====] - 0s 35us/step - loss: 0.5035 - acc: 0.8422 - val_loss: 0.4504 - val_acc: 0.8689
Epoch 3/100
2560/2560 [=====] - 0s 34us/step - loss: 0.4656 - acc: 0.8422 - val_loss: 0.4248 - val_acc: 0.8689
Epoch 4/100
2560/2560 [=====] - 0s 35us/step - loss: 0.4531 - acc: 0.8422 - val_loss: 0.4145 - val_acc: 0.8689
Epoch 5/100
2560/2560 [=====] - 0s 34us/step - loss: 0.4475 - acc: 0.8422 - val_loss: 0.4095 - val_acc: 0.8689
Epoch 6/100
2560/2560 [=====] - 0s 34us/step - loss: 0.4475 - acc: 0.8422 - val_loss: 0.4095 - val_acc: 0.8689

In [16]: model.evaluate(X_test, Y_test)[1]

549/549 [=====] - 0s 60us/step

Out[16]: 0.8561020037515568

```

Figure 14: Building ANN

Visualizing Loss and Accuracy

```

In [17]: plt.plot(hist.history['loss'])
plt.plot(hist.history['val_loss'])
plt.title('Model loss')
plt.ylabel('Loss')
plt.xlabel('Epoch')
plt.legend(['Train', 'Val'], loc='upper right')
plt.show()

```

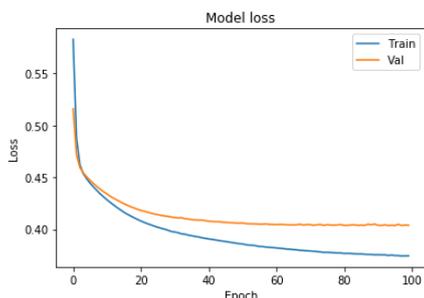


Figure 15: Loss in ANN

```

In [18]: plt.plot(hist.history['acc'])
plt.plot(hist.history['val_acc'])
plt.title('Model accuracy')
plt.ylabel('Accuracy')
plt.xlabel('Epoch')
plt.legend(['Train', 'Val'], loc='lower right')
plt.show()

```

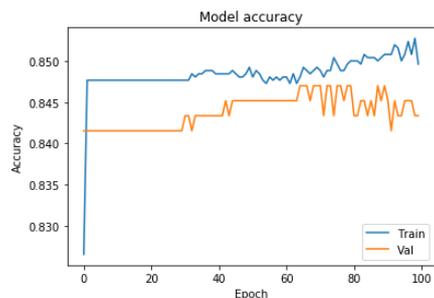


Figure 16: Accuracy for ANN

```
In [22]: Y_pred = model.predict(X_test)
Y_pred = np.round(Y_pred.flatten())

In [23]: accuracy = accuracy_score(Y_test, Y_pred)
print("Accuracy: %.2f%%" % (accuracy*100))
Accuracy: 85.25%

In [24]: print(classification_report(Y_test, Y_pred))
```

	precision	recall	f1-score	support
0.0	0.85	1.00	0.92	468
1.0	0.50	0.01	0.02	81
accuracy			0.85	549
macro avg	0.68	0.51	0.47	549
weighted avg	0.80	0.85	0.79	549

Figure 17: Conclusion for ANN

Data preprocessing and Data Analysis for ML

```
In [22]: df = pd.read_csv('Datasets/Heart_Data.csv')

In [23]: df=df.dropna(axis=0)

In [24]: from numpy import set_printoptions
from sklearn.feature_selection import SelectKBest
from sklearn.feature_selection import f_classif

X = df.drop(['Label'], axis=1)
y = df['Label']
# feature extraction
test = SelectKBest(score_func=f_classif, k=4)
fit = test.fit(X, y)
# summarize scores
set_printoptions(precision=3)
print(fit.scores_)
features = fit.transform(X)
# summarize selected features
print(features[0:5,:])
```

```
[ 30.996 211.751 14.698 1.343 9.918 29.291 8.572 124.379 32.195
 30.757 191.001 84.296 24.783 1.539 55.229]
[[ 39.  0. 106.  70. ]
 [ 46.  0. 121.  81. ]
 [ 48.  0. 127.5 80. ]
 [ 61.  1. 150.  95. ]
 [ 46.  0. 130.  84. ]]
```

Figure 18: Data preprocessing for ML Methods

```
In [28]: df.columns
Out[28]: Index(['gender', 'age', 'education', 'currentSmoker', 'cigsPerDay', 'BPMeds',
              'prevalentStroke', 'prevalentHyp', 'diabetes', 'totChol', 'sysBP',
              'diaBP', 'BMI', 'heartRate', 'glucose', 'Label'],
              dtype='object')

In [29]: name = ['gender', 'age', 'education', 'currentSmoker', 'cigsPerDay', 'BPMeds',
                'prevalentStroke', 'prevalentHyp', 'diabetes', 'totChol', 'sysBP',
                'diaBP', 'BMI', 'heartRate', 'glucose']
          importance = fit.scores_

In [30]: plt.figure(figsize=(40,15), dpi=100)
          sns.barplot(x=name, y=importance)

Out[30]: <matplotlib.axes._subplots.AxesSubplot at 0x23490238508>
```

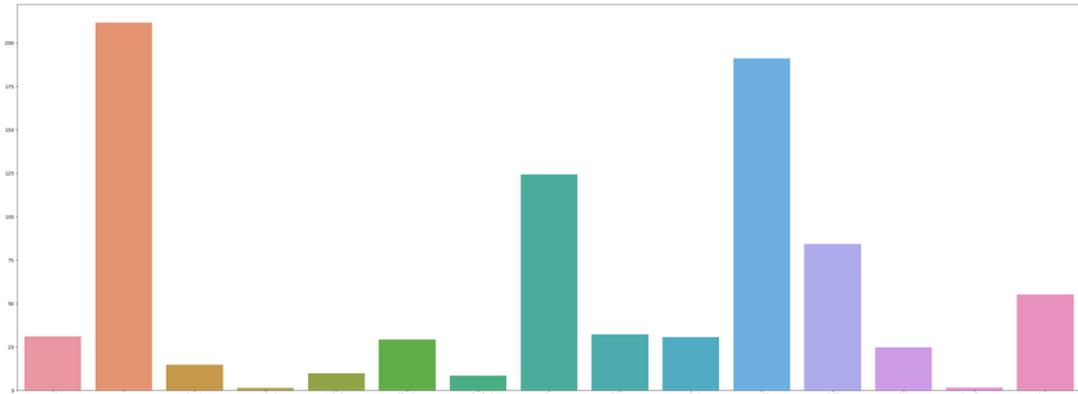


Figure 19: Plotting Attributes

```
In [28]: plt.figure(figsize=(15,7), dpi=100)
          sns.countplot(x='gender', hue='Label', data=df, palette='Set1')

Out[28]: <matplotlib.axes._subplots.AxesSubplot at 0x14b70d191c8>
```

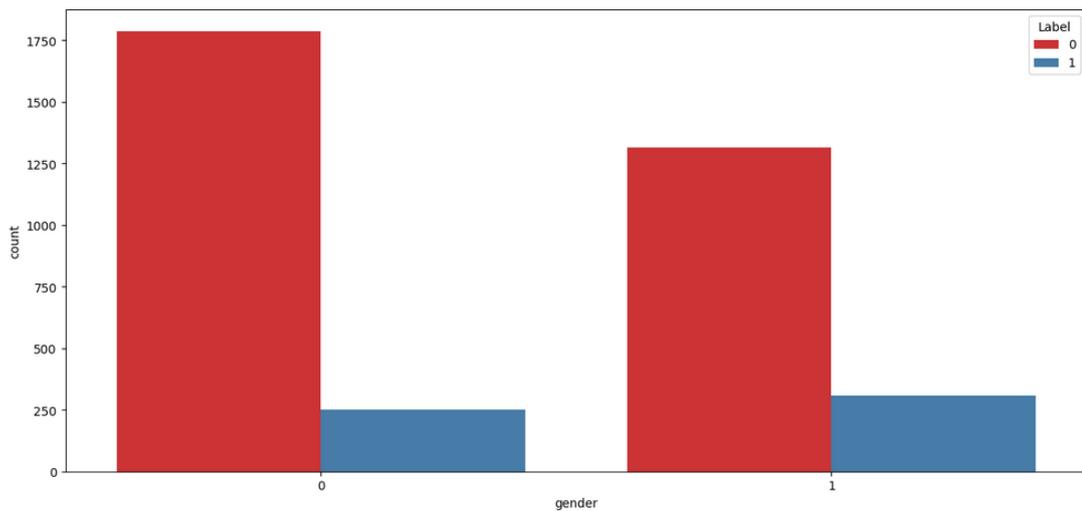


Figure 20: Gender plot

```
In [29]: plt.figure(figsize=(15,7), dpi=100)
sns.countplot(x='age',hue='Label',data=df, palette='Set1')
```

```
Out[29]: <matplotlib.axes._subplots.AxesSubplot at 0x14b70e12248>
```

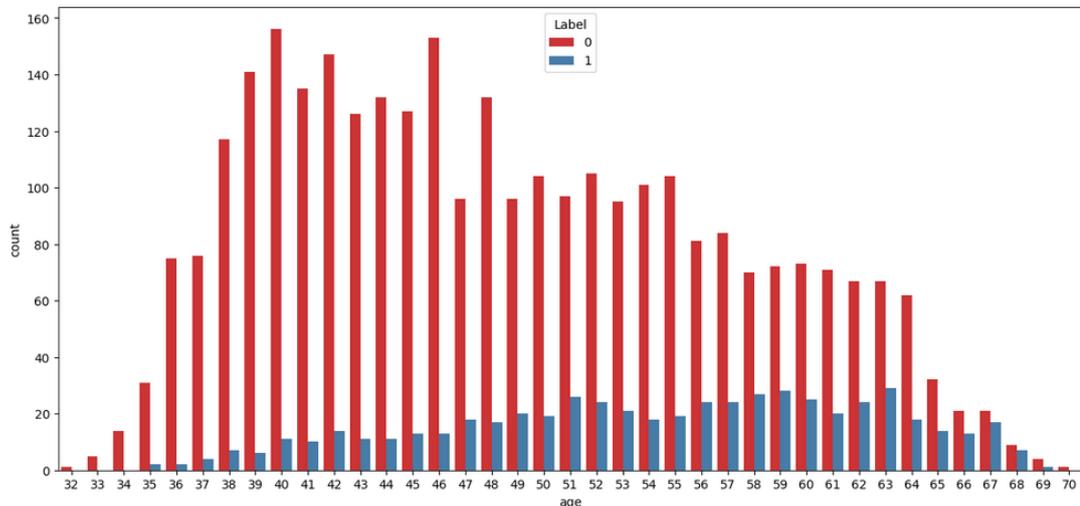


Figure 21: Age group plot

```
In [36]: X = df.drop('Label', axis=1)
y = df['Label']
x_train, x_test, y_train, y_test = train_test_split(X, y, test_size=0.30, random_state=1031)
x_train.head(10)
```

```
Out[36]:
```

	gender	age	education	currentSmoker	cigsPerDay	BPMeds	prevalentStroke	prevalentHyp	diabetes	totChol	sysBP	diaBP	BMI	heartRate	glucose
1111	0	52	2.0	0	0.0	0.0	0	1	1	600.0	159.5	94.0	28.27	78.0	140.0
3938	0	49	3.0	1	4.0	0.0	0	1	0	227.0	150.0	91.0	24.30	88.0	83.0
2469	1	48	4.0	1	20.0	0.0	0	1	0	259.0	135.0	90.0	20.72	102.0	81.0
221	0	38	2.0	1	9.0	0.0	0	0	0	180.0	124.0	66.0	29.29	85.0	68.0
2916	0	45	1.0	0	0.0	0.0	0	0	0	210.0	120.0	72.0	22.01	75.0	93.0
2416	1	56	2.0	1	20.0	0.0	0	1	0	205.0	210.0	130.0	25.49	95.0	127.0
2225	1	38	4.0	0	0.0	0.0	0	0	0	240.0	122.5	80.0	23.97	60.0	43.0
875	1	39	2.0	1	30.0	0.0	0	0	0	225.0	128.0	86.5	25.13	74.0	100.0
3042	0	38	4.0	1	3.0	0.0	0	0	0	177.0	126.0	80.0	23.84	90.0	79.0
1653	1	39	3.0	1	20.0	0.0	0	0	0	148.0	101.0	62.0	24.47	70.0	81.0

```
In [37]: print(x_train.shape, x_test.shape, y_train.shape, y_test.shape)
```

```
(2560, 15) (1098, 15) (2560,) (1098,)
```

```
In [38]: x_train.to_excel('Traning_Testing/X_train.xlsx', index=False)
x_test.to_excel('Traning_Testing/X_test.xlsx', index=False)
y_train.to_excel('Traning_Testing/y_train.xlsx', index=False)
y_test.to_excel('Traning_Testing/y_test.xlsx', index=False)
```

Figure 22: Training the Data

Training the Random Forest model

```
In [34]: from sklearn.ensemble import RandomForestClassifier
```

```
In [35]: rfc = RandomForestClassifier(n_estimators=600)
```

```
In [36]: rfc = rfc.fit(x_train,y_train)
```

Figure 23: Training Random Forest

Predictions and Evaluation of RF Model

```
In [42]: predictions_rf = rfc.predict(x_test)

In [43]: acc_rf = accuracy_score(y_true=y_test, y_pred=predictions_rf)
print("Overall accuracy of ADA model using test-set is : %f" %(acc_rf*100))

Overall accuracy of ADA model using test-set is : 85.063752

In [44]: print(classification_report(y_test,predictions_rf))

              precision    recall  f1-score   support

     0       0.85         1.00         0.92         931
     1       0.64         0.04         0.08         167

 accuracy          0.85         0.85         0.85         1098
 macro avg         0.74         0.52         0.50         1098
 weighted avg      0.82         0.85         0.79         1098

In [45]: print(confusion_matrix(y_test,predictions_rf))

[[927  4]
 [160  7]]

In [46]: filename='model/rf_model.sav'
pickle.dump(rfc,open(filename, 'wb'))
```

Figure 24: Evaluation of RF

Training the SVM model

```
In [47]: from sklearn.svm import SVC

In [48]: model_svm = SVC()

In [49]: model_svm.fit(x_train, y_train)

C:\Users\Omkar\Anaconda3\lib\site-packages\sklearn\svm\base.py:193: FutureWarning: The default value of gamma will change from 'auto' to 'scale' in version 0.22 to account better for unscaled features. Set gamma explicitly to 'auto' or 'scale' to avoid this warning.
  "avoid this warning.", FutureWarning)

Out[49]: SVC(C=1.0, cache_size=200, class_weight=None, coef0=0.0,
decision_function_shape='ovr', degree=3, gamma='auto_deprecated',
kernel='rbf', max_iter=-1, probability=False, random_state=None,
shrinking=True, tol=0.001, verbose=False)
```

Figure 25: Training SVM Model

Predictions and Evaluation

```
In [45]: predictions_svm = model_svm.predict(x_test)

In [46]: acc_svm = accuracy_score(y_true=y_test, y_pred=predictions_svm)
print("Overall accuracy of SVM model using test-set is : %f" %(acc_svm*100))

Overall accuracy of SVM model using test-set is : 84.790528

In [48]: print(classification_report(y_test,predictions_svm))

              precision    recall  f1-score   support

     0       0.85         1.00         0.92         931
     1       0.00         0.00         0.00         167

 accuracy          0.85         0.85         0.85         1098
 macro avg         0.42         0.50         0.46         1098
 weighted avg      0.72         0.85         0.78         1098

In [48]: print(confusion_matrix(y_test,predictions_svm))

[[931  0]
 [167  0]]

In [49]: filename='model/svm_model.sav'
pickle.dump(model_svm,open(filename, 'wb'))
```

Figure 26: Evaluation of SVM

Comparison

```
In [50]: li_x = ['ANN', 'RF', 'SVM']  
li_y = [accuracy, acc_rf, acc_svm]
```

```
In [51]: li = []  
for i in li_y:  
    print(i)  
    li.append(i*100)  
  
0.8597449908925319  
0.8479052823315118  
0.8479052823315119
```

```
In [52]: import seaborn as sns  
print(li)  
sns.barplot(x=li_x, y=li)  
  
[85.97449908925319, 84.79052823315118, 84.79052823315118]
```

```
Out[52]: <matplotlib.axes._subplots.AxesSubplot at 0x15502c0d408>
```

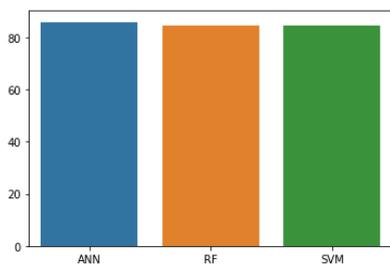


Figure 27: Comparison of ANN, RF and SVM