

# Capstone Project - 3 Supervised - ML - Cardiovascular Risk Prediction

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## Addressing the problem

The purpose of this project is to try a machine learning approach for **Cardiovascular Risk Prediction** by given the id, sex, and information about the health condition of the person. This project contains: Exploratory data analysis, feature engineering, choosing appropriate features, cross algorithms, cross validation, tuning the algorithms, analysis of feature importance, analysis of model performance. The predictions of future could help for saving a person's life and could help them to get rid off unhealthy habits. Another point of view is to test the machine learning algorithms how good are at solving this problem.



## Addressing the problem

#### KNOW YOUR HEART

#### **How Cardiovascular Heart Disease Risk Prediction helps?**

Cardiovascular disease is the leading cause of death worldwide and a major public health concern. Therefore, its risk assessment is crucial to many existing treatment guidelines.

Risk estimates are also being used to predict the magnitude of future cardiovascular disease mortality and morbidity at the population level and in specific subgroups to inform policymakers and health authorities about these risks. Additionally, risk prediction inspires individuals to change their lifestyle and behavior and to adhere to medications.



## Why CHD Risk prediction is important?

The disease is caused when the heart's blood vessels, the coronary arteries, become narrowed or blocked and can't supply enough blood to the heart. This can cause a heart attack or angina. The good news is that if we tackle these risk factors, we can reduce our numbers of people developing heart disease.

By CHD risk prediction, we can analyze the risk factors which are contributing more to develop a heart disease.

The ability to predict risk for and from cardiovascular disease is increasingly important for several reasons. Perhaps foremost among these is the need to determine individual risk to better plan investigation and management with the greatest accuracy and safety for patients and lowest costs for the health care system. This will be even more important as personalized medicine becomes more widespread.



#### **Features Summary**

#### **Independent variables:**

- Id Unique identification number for each patient.
- Age Age of the patient(Continuous).
- Education Educational qualification of the patients(ordinal).
- Sex Male or Female('M' or 'F').
- Is\_smoking Whether or not the patient is a current smoker(Binary).
- **CigsPerDay** The number of cigarettes that the patient smoked on average in one day(Continuous).
- **BPMeds** Whether or not the patient was on blood pressure medication(Nominal).
- prevalentStroke Whether or not the patient previously had a stroke(Nominal).



#### Features Summary (continued)

#### **Independent variables:**

- prevalentHyp Whether or not the patient was hypertensive(Nominal).
- Diabetes Whether or not the patient had diabetes(Nominal).
- Tot\_Chol Total Cholesterol level(Continuous).
- Sys\_BP Systolic blood pressure(Continuous).
- Dia\_BP Diastolic blood pressure(Continuous).
- BMI Body Mass Index(Continuous).
- Heart rate The speed at which the heart beats(Continuous).
- Glucose Glucose level in blood(Continuous).

#### **Dependent variable:**

 TenYearCHD – Whether the patient will develop a risk of Cardiovascular Disease in a 10 years(Binary).



#### **Outliers**

- An outlier is an **extremely high or extremely low data point** relative to the nearest data point and the rest of the neighboring co-existing values in a data graph or dataset you're working with.
- Ways to detect outliers:
- Interquartile range
- Box plot
- Scatter plot
- Z score
- In the given dataset we have used Box plot to detect outliers.



#### Outliers (continued)

**Z-Score** - To handle outliers.

```
# Z Score based technique to remove outliers
lst = ['cigsPerDay','totChol','sysBP','diaBP','BMI','heartRate','glucose']
for i in lst:
    lower_limit = df[i].mean() - 3*df[i].std()
    print(i+'Lower_limit:',round(lower_limit,2))
    upper_limit = df[i].mean() + 3*df[i].std()
    print(i+'Upper_limit:',round(upper_limit,2))
    df[i] = np.where(df[i]>upper_limit,upper_limit,np.where(df[i]<lower_limit,lower_limit,df[i]))</pre>
```



#### **NULL** value treatment

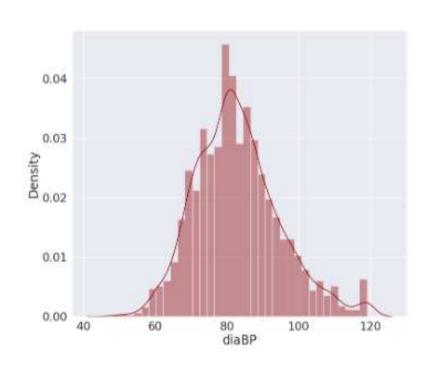
Null values are imputed with most closest value as possible by grouping features

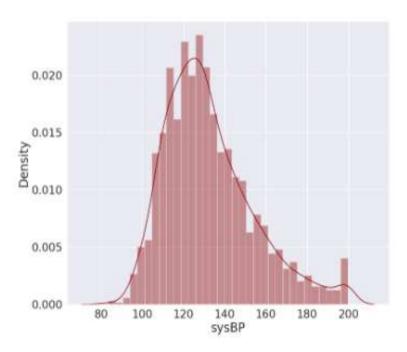
```
# Treating NULL values
# 'education'
df['education'].fillna(0,inplace = True)
df.groupby('education').agg({'age':'mean'})
# print(df.groupby('education').agg({'age':'mean'}))
# From this we can replace education '0' with '1' because based on the average the education varies.
df['education'] = df['education'].apply(lambda x : 1 if x == 0 else x)
# 'ciasPerDay'
df[df['is smoking'] == 1].groupby('sex').agg({'cigsPerDay':'mean'})
# print(df[df['is smoking'] == 1].groupby('sex').agg({'cigsPerDay':'mean'}))
# As the average number of cigarettes consumed by Females is 14 and Males is 22
df.loc[ (df.sex == 0) & (df['cigsPerDay'].isna()), 'cigsPerDay'] = 14
df.loc[ (df.sex == 1) & (df['cigsPerDay'].isna()), 'cigsPerDay'] = 22
# 'alucose'
df.groupby('diabetes').agg({'glucose':'mean'})
# print(df.groupby('diabetes').agg({'glucose':'mean'}))
# Average Glucose level of Patients without Diabetes is 79 and with Diabetes is 172
df.loc[ (df.diabetes == 0) & (df['glucose'].isna()), 'glucose'] = 79
df.loc[ (df.diabetes == 1) & (df['glucose'].isna()), 'glucose'] = 172
# 'BPMeds'
df.loc[(df.prevalentHyp == 1) & (df['BPMeds'].isna()), 'BPMeds'] = 1
```



## **Exploratory Data Analysis**

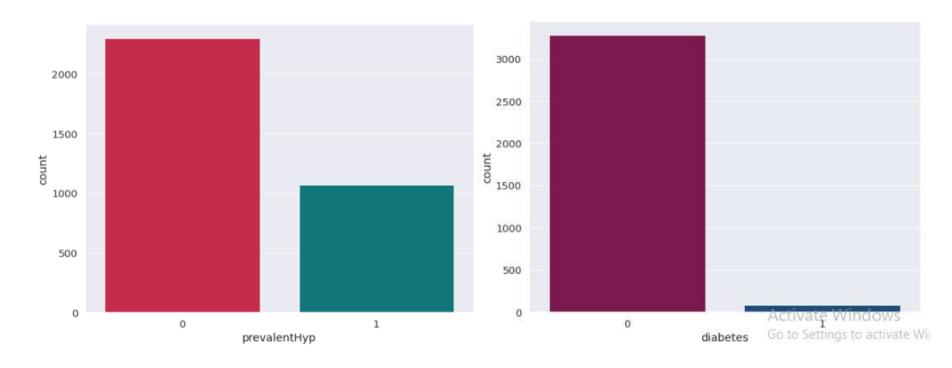
Distribution plot for Numeric independent variable.







Count plot for Categorical independent variables.

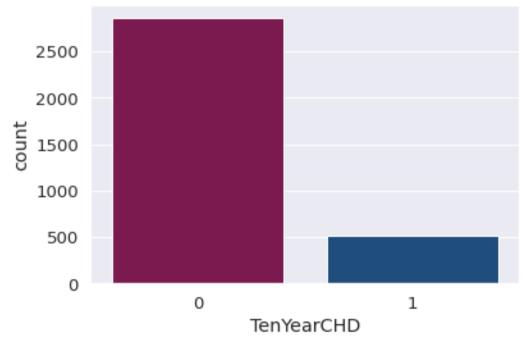




#### Count plot for dependent variable.

Data Imbalance:

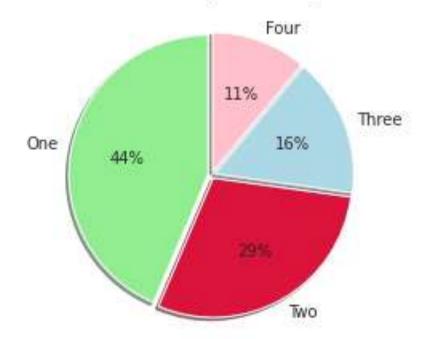
From this we could clearly see the data imbalance.





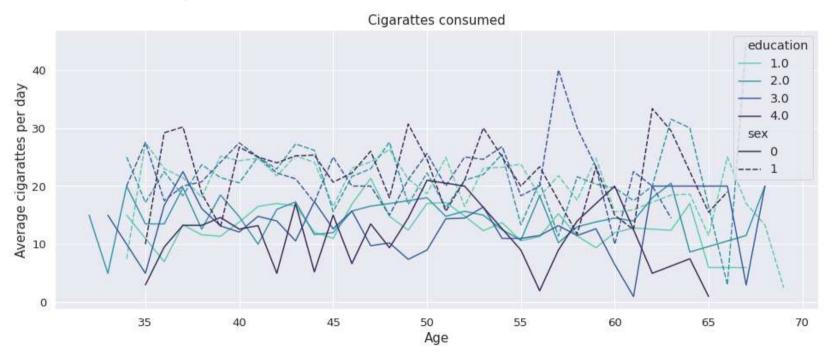
Pie chart for Categorical independent variable

Education percenatge



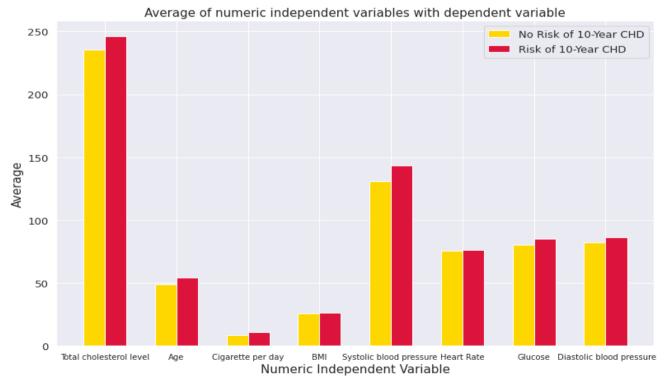


#### **Line plot for Cigarettes consumption**





Bar plot for comparison between patients with and without risk of CHD.





#### Multicollinearity

id	1	0.016	-0.027	-0.0013	-0.0049	-0.017	0.011	-0.039	0.013	0.0036	-0.017	0.021	0.0015	0.04	0.025	0.017	0.0089
age	0.016	1	-0.17	-0.042	-0.21	-0.19	0.14	0.059	0.31	0.11	0.28	0.41	0.22	0.14	-0.0041	0.12	0.23
education	-0.027	-0.17	1	0.025	0.03	0.015	-0.017	-0.032	-0.083	-0.052	-0.021	-0.13	-0.061	-0.13	-0.038	-0.038	-0.054
sex	-0.0013	-0.042	0.025	1	0.22	0.34	-0.051	-0.012	0.0011	0.0084	-0.075	-0.037	0.059	0.098	-0.12	-0.0059	0.082
is_smoking	-0.0049	-0.21	0.03	0.22	1	0.78	-0.052	-0.044	-0.12	-0.053	-0.049	-0.14	-0.12	-0.17	0.065	-0.063	0.034
cigsPerDay	-0.017	-0.19	0.015	0.34	0.78	1	-0.049	-0.043	-0.082	-0.05	-0.025	-0.1	-0.07	-0.1	0.068	-0.073	0.065
BPMeds	0.011	0.14	-0.017	-0.051	-0.052	-0.049	1	0.12	0.28	0.07	0.089	0.28	0.21	0.079	0.016	0.038	0.09
revalentStroke	-0.039	0.059	-0.032	-0.012	-0.044	-0.043	0.12	1	0.071	0.01	-0.011	0.06	0.049	0.0055	-0.019	0.028	0.069
prevalentHyp	0.013	0.31	-0.083	0.0011	-0.12	-0.082	0.28	0.071	1	0.082	0.16	0.71	0.62	0.3	0.15	0.088	0.17
diabetes	0.0036	0.11	-0.052	0.0084	-0.053	-0.05	0.07	0.01	0.082	1.	0.052	0.12	0.06	0.086	0.041	0.51	0.1
totChol	-0.017	0.28	-0.021	-0.075	-0.049	-0.025	0.089	-0.011	0.16	0.052	1	0.2	0.16	0.13	0.086	0.05	0.09
sysBP	0.021	0.41	-0.13	-0.037	-0.14	-0.1	0.28	0.06	0.71	0.12	0.2	1	0.78	0.33	0.18	0.14	0.21
diaBP	0.0015	0.22	0.061	0.059	-0.12	-0.07	0.21	0.049	0.62	0.06	0.16	0.78	1	0.38	0.18	0.069	0.13
BMI	0.04	0.14	-0.13	0.098	-0.17	-0.1	0.079	0.0055	0.3	0.086	0.13	0.33	0.38	1	0.068	0.094	0.065
heartRate	0.025	-0.0041	-0.038	-0.12	0.065	0.068	0.016	-0.019	0.15	0.041	0.086	0.18	0.18	0.068	1	0.1	0.021
glucose	0.017	0.12	-0.038	-0.0059	-0.063	-0.073	0.038	0.028	0.088	0.51	0.05	0.14	0.069	0.094	0.1	1	0.11
TenYearCHD	0.0089	0.23	-0.054	0.082	0.034	0.065	0.09	0.069	0.17	0.1	0.09	0.21	0.13	0.065	0.021	0.11	.1



- 0.0



## **Feature Engineering**

#### The Metric Trap

One of the major issues when dealing with unbalanced datasets relates to the metrics used to evaluate their model. Using simpler metrics like accuracy score can be misleading. In a dataset with highly unbalanced classes, the classifier will always "predict" the most common class without performing any analysis of the features and it will have a high accuracy rate, obviously not the correct one.

#### Imbalance in our dataset

```
No Risk of 10-Year CHD: 84.87 %
Risk of 10-Year CHD: 15.13 %
```



## Feature Engineering (continued)

## To encounter imbalance dataset following sampling techniques are used: Random Over Sampling

Randomly duplicate examples in the minority class.

#### **Random Under Sampling**

Randomly delete examples in the majority class.

#### **Synthetic Minority Oversampling Technique (SMOTE)**

New examples are synthesized from the existing examples. This is a type of data augmentation for the minority class.

#### **SMOTETomek**

A hybrid method which is a mixture of under-sampling method (Tomek) with an oversampling method (SMOTE).



## Feature Engineering (continued)

For our dataset Random Over Sampling is found to be effective.

```
# Random Over Sampling
from imblearn.over_sampling import RandomOverSampler

ros = RandomOverSampler(sampling_strategy= 1.0,random_state = 0)
X_ros,y_ros = ros.fit_resample(data.iloc[:,0:-1], data['TenYearCHD'])
print('X ros :',X_ros.shape)
print('y ros :',y_ros.shape)
```

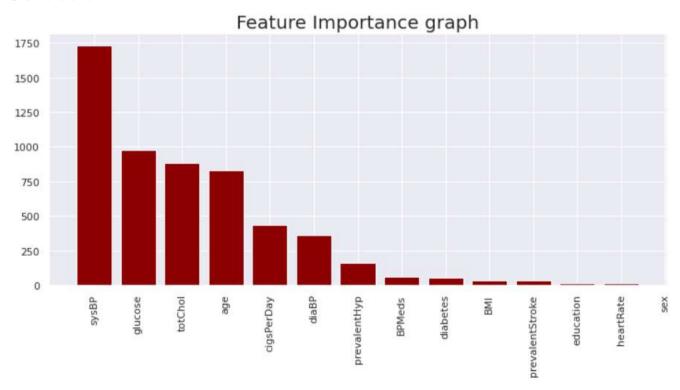
```
# Classes after sampling
ros_xy['TenYearCHD'].value_counts()

1    2856
0    2856
Name: TenYearCHD, dtype: int64
```



## Feature Engineering (continued)

#### **Feature Selection**





## Preparing dataset for modelling

#### **Train Test Split**

Train test split is a model validation procedure that allows you to simulate how a model would perform on new/unseen data.

```
X = data[independent variables].values
y = data[dependent variable].values
# Splitting the dataset into the Training set and Test set
X train, X test, y train, y test = train test split(X, y, test size = 0.25, random state = 0)
```

- X\_train: (2523, 15) X\_test : (842, 15) y\_train: (2523, 1)
- y\_test : (842, 1)



## Preparing dataset for modelling

#### **Standardization**

Standardizing a dataset involves rescaling the distribution of values so that the mean of observed values is 0 and the standard deviation is 1.

```
# Transforming data

# Standardization
scaler = MinMaxScaler()
X_train = scaler.fit_transform(X_train)
X_test= scaler.transform(X_test)
```



## **Applying Model**

#### **Logistic Regression**

```
# Logistic Regression
 # Creating an instance of Logistic Regression
 logistic = LogisticRegression(fit intercept = True)
 # Hyperparameters
 grid = {'C':10.0**np.arange(-2.3), 'penalty':['l1','l2']}
 # KFol.d
 cv = KFold(n splits = 5, random state = None, shuffle = False)
 # Grid Search
 clf = GridSearchCV(logistic,grid,cv = cv,n jobs = -1,scoring = 'f1 macro')
 # Model fit
 clf.fit(X train, y train)
GridSearchCV(cv=KFold(n splits=5, random state=None, shuffle=False),
             estimator=LogisticRegression(), n jobs=-1,
             param grid={'C': array([1.e-02, 1.e-01, 1.e+00, 1.e+01, 1.e+02]),
                         'penalty': ['l1', 'l2']},
             scoring='f1 macro')
```

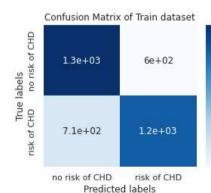


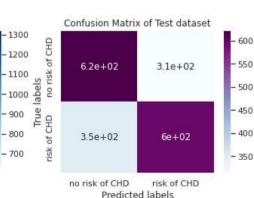
## **Applying Model**

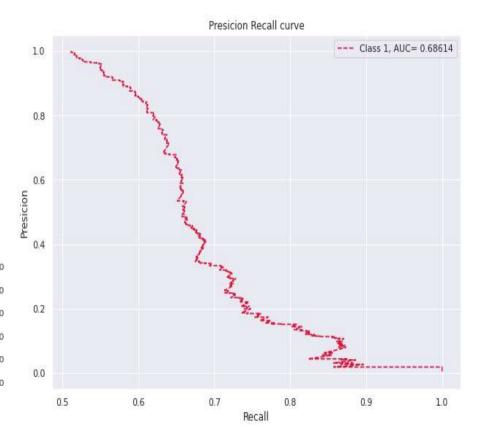
#### **Model Validation(Logistic regression)**

The ROC AUC score on train data is : 0.71434 The ROC AUC score on test data is : 0.69948 The Precision Recall score on train data is: 0.70048 The Precision Recall score on test data is : 0.68614 recall f1-score precision support 0.64 0.66 0.65 935 0.66 0.63 0.64 950

accuracy 0.65 1885 macro avg 0.65 0.65 0.65 1885 weighted avg 0.65 0.65 0.65 1885







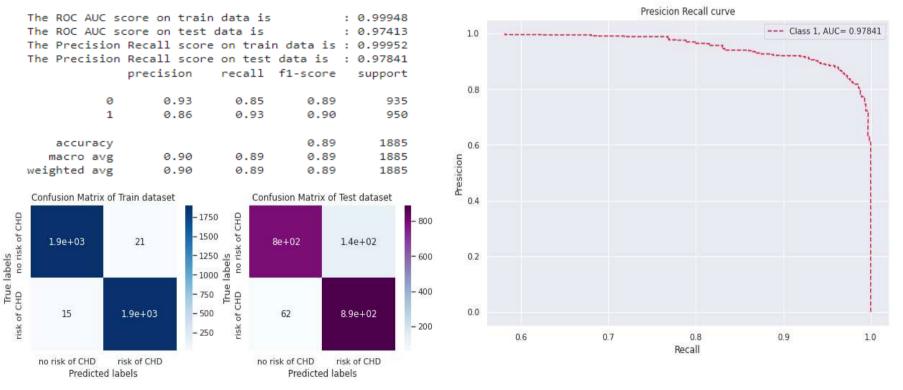


#### **Random Forest Classifier**

```
# Create an instance of the RandomForestClassifier
 rfc model = RandomForestClassifier(criterion = 'gini',class weight = {0:1,1:1.5},random state = 0)
# By adding weights to Class 1, False Negative can be reduced
# Grid search
 rfc grid = GridSearchCV(estimator=rfc model,param grid = param dict,cv = 5, verbose=2, scoring='f1 macro')
# Model Fit
rfc grid.fit(X train, v train)
Fitting 5 folds for each of 1 candidates, totalling 5 fits
[CV] END max depth=22, min samples leaf=3, min samples split=4, n estimators=100; total time=
                                                                                                 0.5s
[CV] END max depth=22, min samples leaf=3, min samples split=4, n estimators=100; total time=
                                                                                                 0.5s
[CV] END max depth=22, min samples leaf=3, min samples split=4, n estimators=100; total time=
                                                                                                 0.55
[CV] END max depth=22, min samples leaf=3, min samples split=4, n estimators=100; total time=
                                                                                                 0.55
[CV] END max depth=22, min samples leaf=3, min samples split=4, n estimators=100; total time=
                                                                                                 0.55
GridSearchCV(cv=5.
             estimator=RandomForestClassifier(class weight={0: 1, 1: 1.5},
                                              random state=0),
             param grid={'max depth': [22], 'min samples leaf': [3],
                         'min samples split': [4], 'n estimators': [100]},
             scoring='f1 macro', verbose=2)
```



#### Model Validation(Random forest classifier)



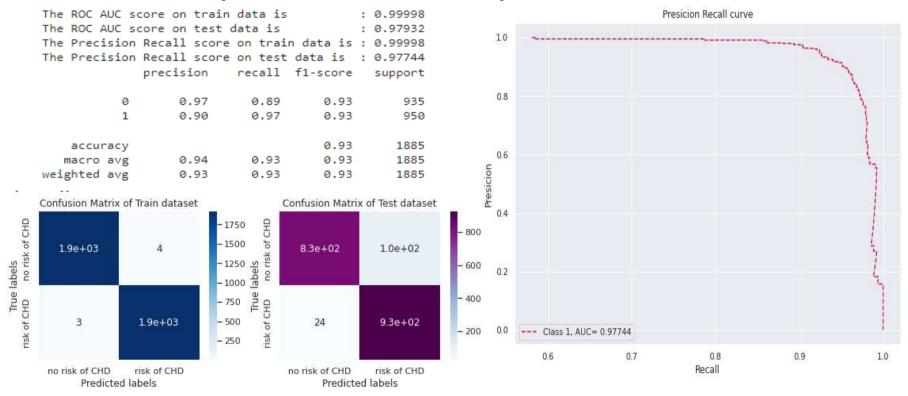


#### **Gradient Boost Classifier**

```
# Create an instance of the RandomForestClassifier
gbc model = GradientBoostingClassifier(random state = 0)
# Grid search
gbc grid = GridSearchCV(estimator=gbc model,param grid = param dict,cv = 5, verbose=2, scoring='roc auc')
# Model Fit
gbc grid.fit(X train,y train)
Fitting 5 folds for each of 1 candidates, totalling 5 fits
[CV] END max depth=50, min samples leaf=70, min samples split=60, n estimators=200; total time=
                                                                                                  3.1s
[CV] END max depth=50, min samples leaf=70, min samples split=60, n estimators=200; total time=
                                                                                                  3.05
[CV] END max depth=50, min samples leaf=70, min samples split=60, n estimators=200; total time=
                                                                                                  3.0s
[CV] END max depth=50, min samples leaf=70, min samples split=60, n estimators=200; total time=
                                                                                                  3.05
[CV] END max depth=50, min samples leaf=70, min samples split=60, n estimators=200; total time=
                                                                                                  3.1s
GridSearchCV(cv=5, estimator=GradientBoostingClassifier(random state=0),
             param grid={'max depth': [50], 'min samples leaf': [70],
                         'min samples split': [60], 'n estimators': [200]},
             scoring='roc auc', verbose=2)
```



#### Model Validation(Gradient Boost classifier)





#### **XGB Classifier**

```
# Create an instance of the RandomForestClassifier
xgb model = XGBClassifier(random state = 0,seed = 1)
# Grid search
xgb grid = GridSearchCV(estimator=xgb model,param grid = param dict,cv = 5, verbose=2, scoring='roc auc')
# Model Fit
xgb grid.fit(X train,y train)
Fitting 5 folds for each of 1 candidates, totalling 5 fits
[CV] END gamma=1, learning rate=0.2, max depth=10, min child weight=1, n estimators=100, reg lambda=1, subsample =0.5; total time=
                                                                                                                                     0.65
[CV] END gamma=1, learning rate=0.2, max depth=10, min child weight=1, n estimators=100, reg lambda=1, subsample =0.5; total time=
                                                                                                                                    0.65
[CV] END gamma=1, learning rate=0.2, max depth=10, min child weight=1, n estimators=100, reg lambda=1, subsample =0.5; total time= 0.6s
[CV] END gamma=1, learning rate=0.2, max depth=10, min child weight=1, n estimators=100, reg lambda=1, subsample =0.5; total time= 0.6s
[CV] END gamma=1, learning rate=0.2, max depth=10, min child weight=1, n estimators=100, reg lambda=1, subsample =0.5; total time= 0.6s
GridSearchCV(cv=5, estimator=XGBClassifier(seed=1),
             param grid={'gamma': [1], 'learning rate': [0.2],
                         'max depth': [10], 'min child weight': [1],
                         'n estimators': [100], 'reg lambda': [1],
                         'subsample ': [0.5]},
             scoring='roc auc', verbose=2)
```



## Model Validation(XGB Classifier) The ROC AUC score on train data is : 0.9

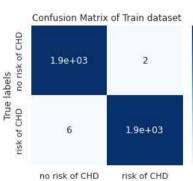
0.89

The ROC AUC score on train data is : 0.99995
The ROC AUC score on test data is : 0.97255
The Precision Recall score on train data is : 0.99995
The Precision Recall score on test data is : 0.97338
precision recall f1-score support

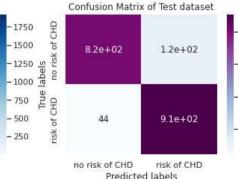
0 0.95 0.88 0.91 935

accuracy 0.92 1885 macro avg 0.92 0.92 0.92 1885 weighted avg 0.92 0.92 0.92 1885

0.95

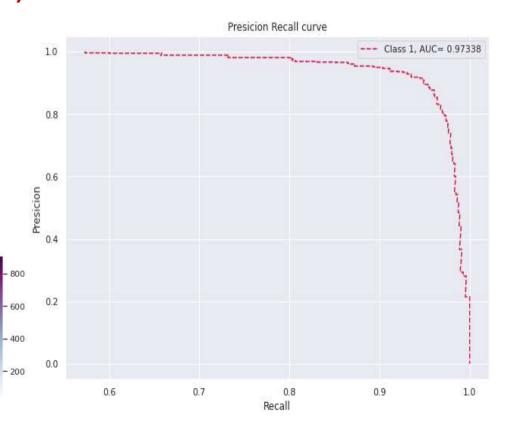


Predicted labels



0.92

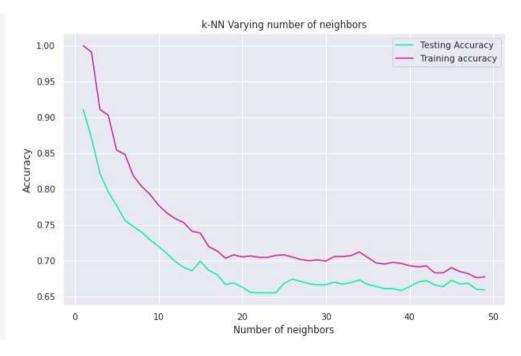
950





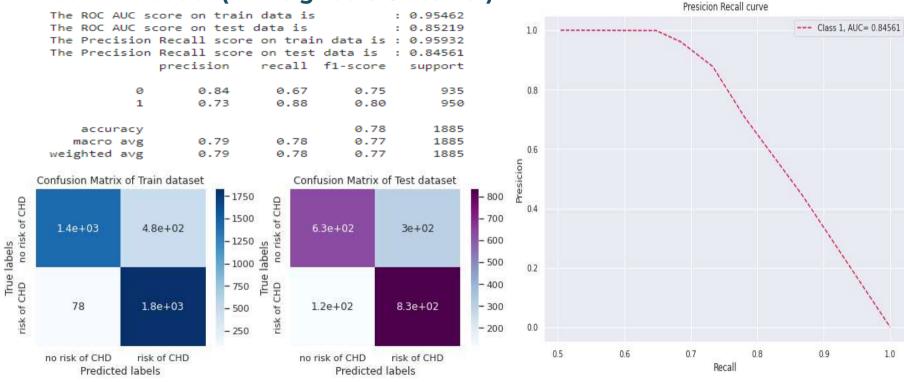
#### **KNNeighbors Classifier**

```
#Setup arrays to store training and test accuracies
neighbors = np.arange(1,50)
train accuracy =np.empty(len(neighbors))
test accuracy = np.empty(len(neighbors))
for i,k in enumerate(neighbors):
   # Setup a knn classifier with k neighbors
    knn = KNeighborsClassifier(n_neighbors=k)
   # Fit the model
    knn.fit(X train, y train)
    # Compute accuracy on the training set
   train accuracy[i] = knn.score(X train, y train)
    # Compute accuracy on the test set
   test accuracy[i] = knn.score(X_test, y_test)
```





#### Model Validation(KNNeighbors Classifier)





#### **Model Performance**

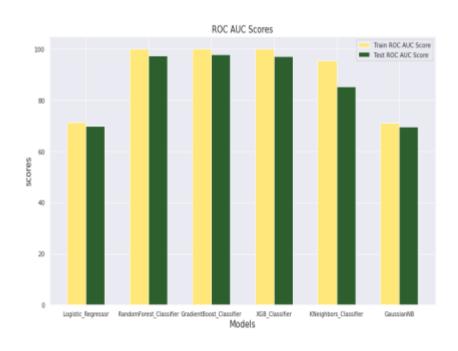
#### Used model's scores

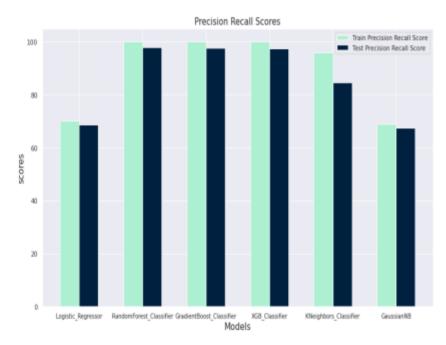
	index	Models	Train ROC AUC score	Test ROC AUC score	Train Precision recall score	Test Precision recall score
0	0	Logistic_Regressor	71.4341	69.9479	70.0481	68.6141
1	1	RandomForest_Classifier	99.9481	97.4129	99.9523	97.8414
2	2	${\sf GradientBoost\_Classifier}$	99.9982	97.9320	99.9982	97.7439
3	3	XGB_Classifier	99.9950	97.2553	99.9952	97.3383
4	4	KNeighbors_Classifier	95.4622	85.2187	95.9316	84.5614
5	5	GaussianNB	70.9767	69.6526	68.9231	67.4698



## Model Performance (continued)

#### Visualization of model scores







#### Conclusion

- It's quite obvious that smokers have a high risk of 10 year CHD.
- Patients with no history of hypertensive, stroke, diabetes have less risk of 10 year CHD.
- By using simple Logistic Regressor algorithm we were able to get the Precision Recall AUC score of 68 %
- Very little improvement in Precision Recall AUC score after using Random Forest classifier because of data imbalance.
- Data Imbalance is addressed by Random Over sampling technique...
- We got the best Precision Recall AUC score of 97.8 percent using <u>Random</u>
   <u>Forest Classifier</u> with cross validation and hyper parameter tunning.
- Top five most important features are SysBP,Glucose
   ,Totchol,age,Cigsperday



#### **Conclusion**

 For the given dataset, <u>Random Forest Classifier</u> has proven to be the best fit model with,

Train Precision Recall AUC score: 99.9 %

**Test Precision Recall AUC score** : 97.8%

Train ROC AUC score :99.9 %

**Test** ROC AUC score : 97.4%



# Thank you