# CARDIOVASCULAR DISEASE PREDICTION

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# **AGENDA**

- Study, Data, & Primary outcome
- Hypothesis
- Data Cleaning & Preprocessing: Predictors
- Table I
- Exploratory Data Analysis: Visualization
- Unsupervised Learning
- Supervised Learning
- Ensemble Learning
- Extended Thinking

# STUDY, DATA, & PRIMARY OUTCOME

- World Health Organization estimated 12 million deaths worldwide per year due to Heart diseases.
- Half of the deaths in the US and other developed countries are due to Cardiovascular Diseases.
- The early prognosis of cardiovascular diseases can aid in **making decisions** on lifestyle changes in high risk patients and in turn reduce the complications. This research intends to pinpoint the **most relevant/risk factors** of heart disease as well as predict the overall risk using Machine Learning models.
- Source The dataset is publicly available on the **Kaggle** website, and it is from an **ongoing cardiovascular study** on residents of the town of **Framingham**, **Massachusetts**. The classification goal is to predict whether the patient has 10-year risk of future coronary heart disease (CHD)
- The dataset provides the patients' information. It includes over **4,000 records and 15 attributes**. Variables Each attribute is a potential risk factor. There are both demographic, behavioral and medical risk factors.
- TenYearCHD: 10-year coronary heart disease prediction 0/1

df0.shape
(4238, 16)

### **HYPOTHESIS**

- TenYearCHD is highly correlated with diabetes, glucose, and smoking.
- Education, age, and gender will not affect much to the TenYearCHD and the glucose level.
- Most of my prediction will be 0s, which means not risky for coronary disease in the next 10 years, because the glucose level for people is very low(mostly 55 - 100).
- (Normal blood sugar levels are less than 100 mg/dL after not eating for at least 8 hours; <140 mg/dL two hours after eating)
- Machine learning: Ensemble ML will give better prediction accuracy than Base ML models. Accuracy comparison

#### Gender and risk

Although men tend to develop coronary artery disease earlier in life, after age 65 the risk of heart disease in women is almost the same as in men. Women have many of the same risk factors for heart disease as men, such as smoking, high blood pressure, and high cholesterol.

How Age and Gender Affect Your Heart | Kaiser Permanente ...

https://www.niddk.nih.gov > diabetes > overview > preventing-problems > h...

Over time, high blood glucose from diabetes can damage your blood vessels and the nerves that control your heart and blood vessels. The longer you have diabetes, the higher the chances that you will develop heart disease. ... In adults with diabetes, the most common causes of death are heart disease and stroke.

#### Diabetes, Heart Disease, and Stroke | NIDDK

Smoking raises your risk of getting CAD and dying early from CAD. Carbon monoxide, nicotine, and other substances in tobacco smoke can promote atherosclerosis and trigger symptoms of coronary artery disease. ... Clumping platelets can then block your coronary arteries and cause a heart attack. 2012年

Smoking and Coronary Artery Disease - CardioSmart https://www.cardiosmart.org > Healthwise

# **CLEANING & PREPROCESSING**

#### number of predictors

#### df0.isnull().sum()

male	0
age	0
education	105
currentSmoker	0
cigsPerDay	29
BPMeds	53
prevalentStroke	0
prevalentHyp	0
diabetes	0
totChol	50
sysBP	0
diaBP	0
BMI	19
heartRate	1
glucose	388
TenYearCHD	0
dtype: int64	

#### • Introduction:

- Systolic pressure; Diastolic blood pressure
- Prevalent Stroke: whether or not the patient had previously had a stroke (Nominal)
- Prevalent Hyp: whether or not the patient was hypertensive

#### Missing Values Imputation:

- Mean and medium to fill the null values in the cigsPerDay, BPMeds, totChol, BMI, and heartRate column.
- KNN for education after scaling
- Glucose: (compared RMSE without using the Education, training set = 92.5%)
  - Multiple Regression: 15.1074
  - Gradient Boosting Regressor: 17.7771
  - XGBoost Regressor: 18.3610
- Error: 15/<u>100-150</u> =15/150 -> 15/100 = <u>10%-15%</u>.

# TABLE I

•	Outcome -	- Ten Yeaı	$^{\sim}$ CHD
	<b>-</b> 4 4 4 5 1 1 1 6		<b>—</b> : : —

relationship

relationships!

H<sub>a</sub>: some relationships

P<0.05 -> Rejecting null

hypothesis; there are

•	Age			Grouped	by	TenYearCHD isnull	0	1	pval
•	Education: 1, 2, 3, 4	variable	level						
		n					3594	644	
•	Male: 0, I	age				0	48.8 (8.4)	54.1 (8.0)	<0.001
i laic.	1 1410. 0, 1	education	1			0	1437 (40.0)	330 (51.2)	<0.001
•	Blood Pressure Meds: 0, I		2				1147 (31.9)	155 (24.1)	
	2.002 1.0004.01.020.0, 1		3				607 (16.9)	88 (13.7)	
•	Diabetes: 0, I		4				403 (11.2)	71 (11.0)	
	,	male	0			0	2118 (58.9)	301 (46.7)	<0.001
•	$H_0$ : $\mu_1 = \mu_2$ : no significance		1				1476 (41.1)	343 (53.3)	
	♥ 1 ± 1 <del>2</del> , <b>0</b>						155		

**BPMeds** 

diabetes

3511 (97.7) 603 (93.6)

0 3525 (98.1) 604 (93.8)

41 (6.4)

40 (6.2)

83 (2.3)

69 (1.9)

<0.001

<0.001

ptest

Two Sample T-test

Chi-squared

Chi-squared

Chi-squared

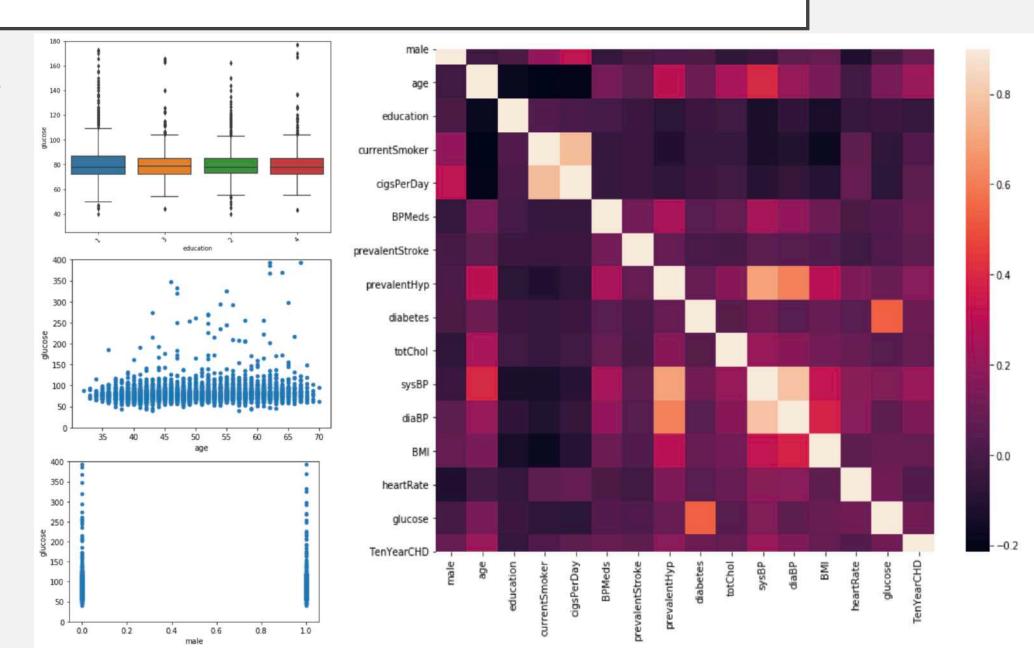
Chi-squared

<sup>[1]</sup> Warning, Hartigan's Dip Test reports possible multimodal distributions for: age.

<sup>[2]</sup> Warning, test for normality reports non-normal distributions for: age.

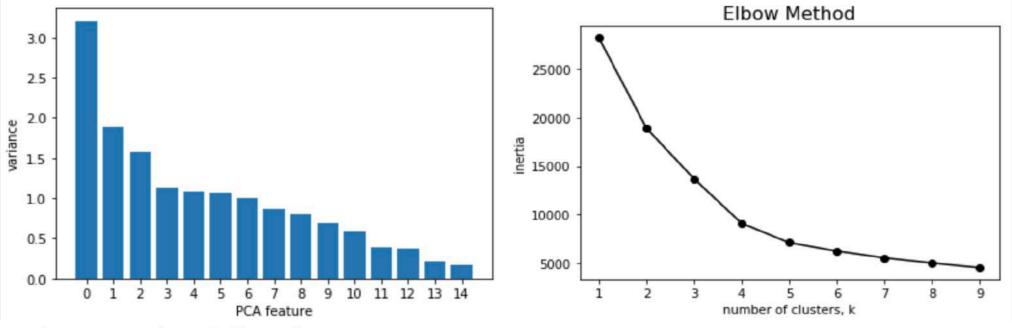
# **EXPLORATORY DATA ANALYSIS**

- Back to my hypothesis
- Age and education could indirectly affect TenYearCHD through affecting glucose.
- Highly correlated with <u>outcome</u>:
- Age
- prevalentHyp
- prevalentStroke
- Diabetes
- totChol
- sysBP
- diaBP



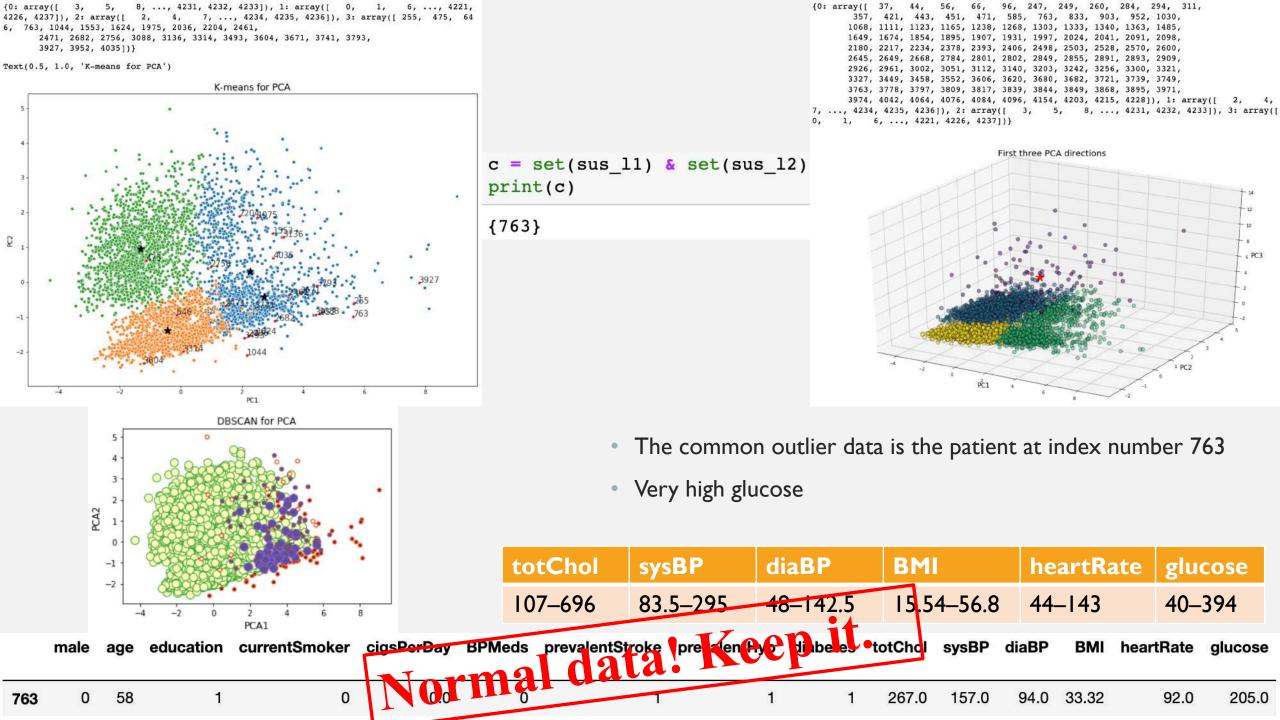
# UNSUPERVISED

- Scaling dataset
- PCA
- 15 PCs in total
- PCI: 21.3%
- PC2: 12.5%
- PC3: 10.5%
- Total: 44.3%
- Clusters to detect any anomalies or outliers



```
Variance: Projected dimension
```

```
21.3%:
         -0.05 * f1 + 0.30 * f2 + -0.11 * f3 + -0.20 * f4 + -0.17 * f5 + 0.20 * f6 + 0.07
* f7 +
       0.43 * f8 + 0.14 * f9 + 0.19 * f10
12.5%:
          0.35 * f1 + -0.11 * f2 + -0.02 * f3 + 0.59 * f4 + 0.63 * f5 + 0.04 * f6 + -0.02
* f7 +
       0.16 * f8 + -0.02 * f9 + 0.02 * f10
          0.06 * f1 + 0.02 * f2 + -0.03 * f3 + 0.06 * f4 + 0.05 * f5 + -0.05 * f6 + -0.02
10.5%:
* f7 + -0.12 * f8 + 0.69 * f9 + -0.02 * f10
          0.56 * f1 + 0.11 * f2 + -0.08 * f3 + -0.11 * f4 + -0.03 * f5 + 0.08 * f6 + 0.28
7.5%:
* f7 + 0.01 * f8 + 0.03 * f9 + -0.27 * f10
7.1%:
         -0.20 * f1 + -0.21 * f2 + 0.48 * f3 + 0.06 * f4 + 0.00 * f5 + 0.55 * f6 + 0.53
* f7 + 0.09 * f8 + 0.05 * f9 + -0.16 * f10
```



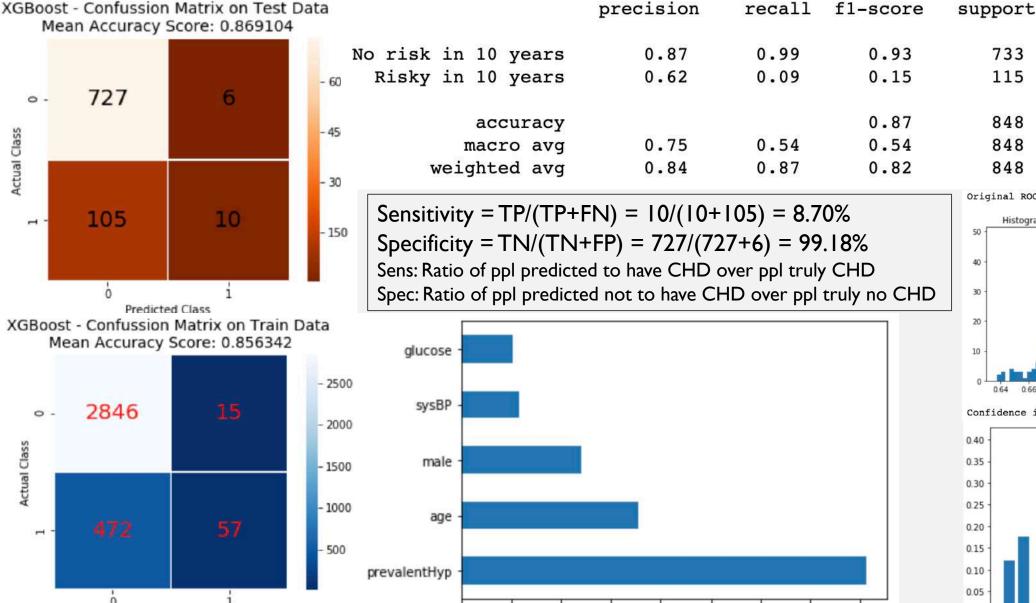
# SUPERVISED LEARNING MODELS

Training set = 80%; CV = 5

	<b>Model Name</b>	Accuracy on Testing	Accuracy on Training	AUC_ROC	DeLong 95% CI	I,000 Boostrapping 95% CI	Fitting condition
I	Logistic Regression	0.8656	0.8484	0.6629	(0.6074, 0.7184)	(0.6148, 0.7075)	Good
	Decision Tree						
2	Classifier	0.7807	0.8484	0.5405	(0.4957, 0.5852)	(0.5016, 0.5790)	Overfitting
	Random Forest						A little
3	Classifier	0.8620	0.9035	0.7125	(0.6599, 0.7651)	(0.6667, 0.7566)	overfitting
4	KNN with scaling	0.8514	0.8546	0.5952	(0.5387, 0.6518)	(0.5482, 0.6422)	Good
5	SVM with scaling	0.8585	0.8602	0.6050	(0.5426, 0.6674)	(0.5508, 0.6572)	Good
6	AdaBoost	0.8632	0.8584	0.7042	(0.6514, 0.7570)	(0.6594, 0.7474)	Good
7	GradientBoosting	0.8644	0.8584	0.7127	(0.6607, 0.7647)	(0.6689, 0.7554)	Good
8	XGBoost	0.8691	0.8563	0.7155	(0.6644, 0.7667)	(0.6732, 0.7561)	Good
	ANN with SMOTE						
9	and scaling	0.7783	0.9834	0.5232	(0.4622, 0.5842)	(0.4737, 0.5754)	Overfitting
10	Bagging Classifier	0.8644	0.8440	0.5893	(0.5323, 0.6463)	(0.5410, 0.6354)	Good

Predicted Class

# **XGBOOST**



0.05

0.10

0.15

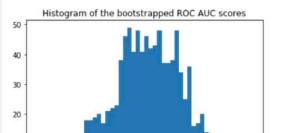
0.20

0.25

0.30

0.35

0.40



733

115

848

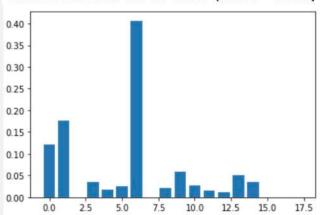
848

848

10

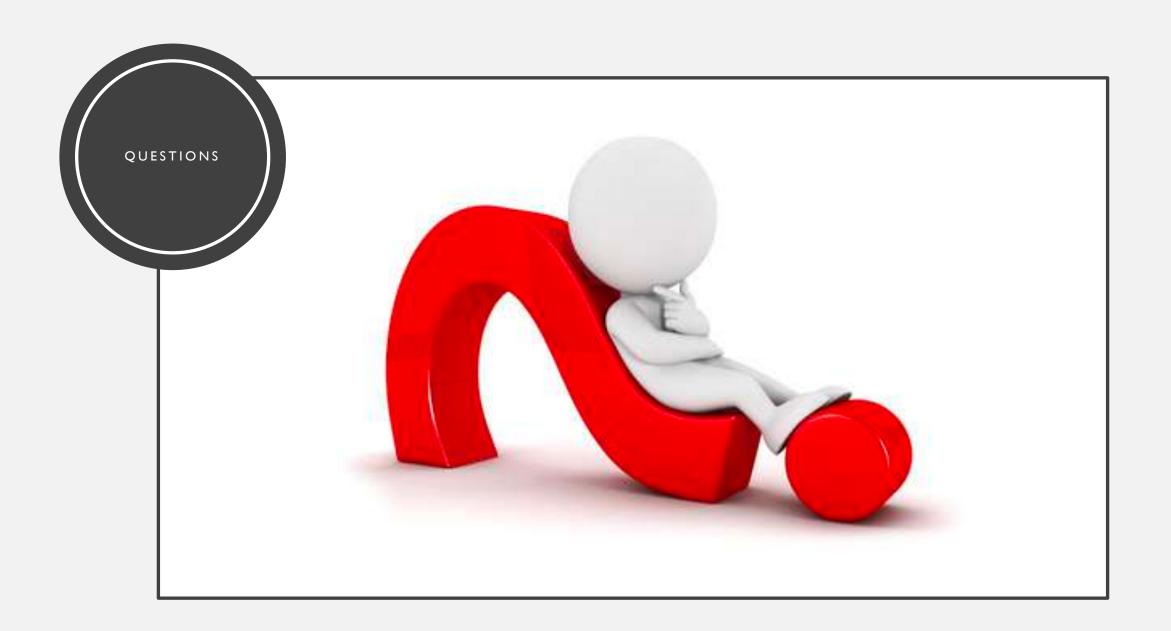
Original ROC area: 0.7155

0.68 0.70 0.72 0.74 0.76 0.78 Confidence interval for the score: [0.6732 - 0.7561]



## EXTENDED THINKING

- Feature Engineering
- My dataset is a mixture of continuous and discrete. I couldn't apply Naïve Bayes Model directly on my dataset. GaussianNB continuous; MultinomialNB discrete. 2 ways to do:
  - I can transfer all the continuous variables into categorical variable, such as "Very Low -0, Low -1, Medium -2, High -3, Very High -4" (20% each, (max min)/5). Remove the previous continuous columns, and then I can fit a MultinomialNB.
  - I can break my dataset into 2 parts: I with only categorical, and I with only continuous. GaussianNB continuous; MultinomialNB discrete. Then transform all the dataset by taking the class assignment probabilities (with predict\_proba method) as new features: np.hstack((multinomial\_probas, gaussian\_probas)).
    - Finally, refit a new model (e.g. a new gaussian NB) on the new features.
  - Found a package called "mixed\_naive\_bayes".
- Stacking & Soft/Hard voting classifiers
- Transform my dataset into a spectrum to apply **CNN** according to the color distribution to predict the 10-year coronary heart disease.



# Thank you!