

### Outline

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- Methodology
- Results
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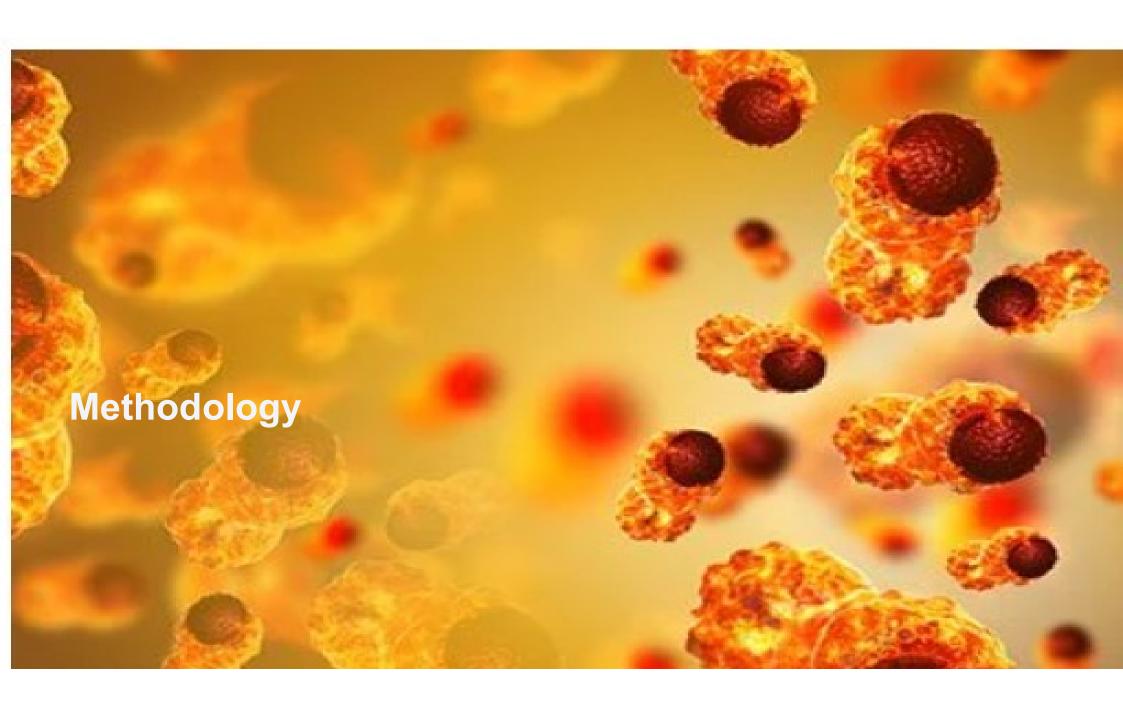
### Introduction

#### Background of the project:

During the IBM course on Coursera we have done some exercises. One of them was a cancer prediction with SVM (Support Vector Machine) method. After completing the course, I used the same data-set and have decided to improve the prediction based on what I have learned to have a better predicting model.

#### Problems we want to find answers:

Classification of samples of tumor to find out whether it is benign or malignant based on the attributes of the tumor.



## Methodology

#### **Executive Summary**

- Perform data wrangling
- Perform interactive visual analytics using Plotly Dash
- Perform predictive analysis using classification models

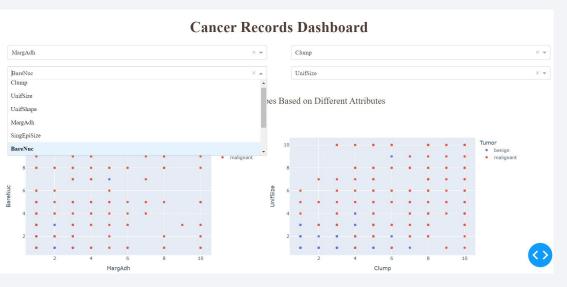
## **Data Wrangling**

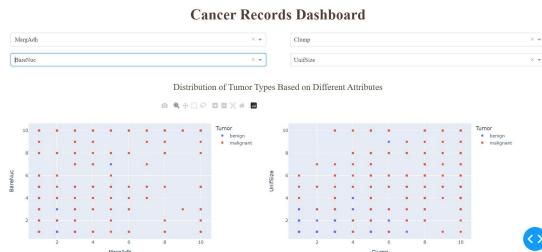
- Data wrangling in this project is as follows:
  - Creating a data frame from csv file
  - Get the shape of the data frame
  - Create a new column called Tumor to be used for plot legend
  - Get the data types of each column
  - Convert the non numeric values (numbers that are defined as an object) to numeric ones
  - Replace all the non digits (here "?") with NaN
  - Remove the NaNs
  - Convert the values to integer type

### **EDA** with Data Visualization

Interactive visual analytics using Plotly Dash

The attributes are selected from drop-down list and scatter plot is plotted





## Predictive Analysis (Classification)

- For predictive analysis
  - The target values list is converted into a numpy array

```
Y=df['Class'].to_numpy()
Y
```

X=transform.fit transform(X)

transform=preprocessing.StandardScaler()

Selected features are converted to have 0 mean and unit variance

```
#Standardize features by removing the mean and scaling to unit variance
#It should be a standard normally distributed data (e.g. Gaussian with 0 mean and unit variance)
```

The data set is splitted into train and test sets

## Predictive Analysis (Classification)

 Defining the parameters for the classification model (e.g. Support Vector Machine) and using the grid search method to find the best parameters

```
[ ] parameters = {'kernel':('linear', 'rbf', 'poly', 'rbf', 'sigmoid'),
                   'C': np.logspace(-3, 3, 5),
                   'gamma':np.logspace(-3, 3, 5)}
     svm = SVC()
    svmcv=GridSearchCV(svm,parameters,scoring='accuracy',cv=10)
     svm cv=svmcv.fit(X train,Y_train)
[ ] print("tuned hpyerparameters :(best parameters) ",svm cv.best params)
     print("accuracy :",svm cv.best score )
    tuned hpyerparameters :(best parameters) {'C': 0.03162277660168379, 'gamma': 1.0, 'kernel': 'si
    accuracy: 0.9817508417508417
```

## Predictive Analysis (Classification)

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```
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Y
```

transform=preprocessing.StandardScaler()

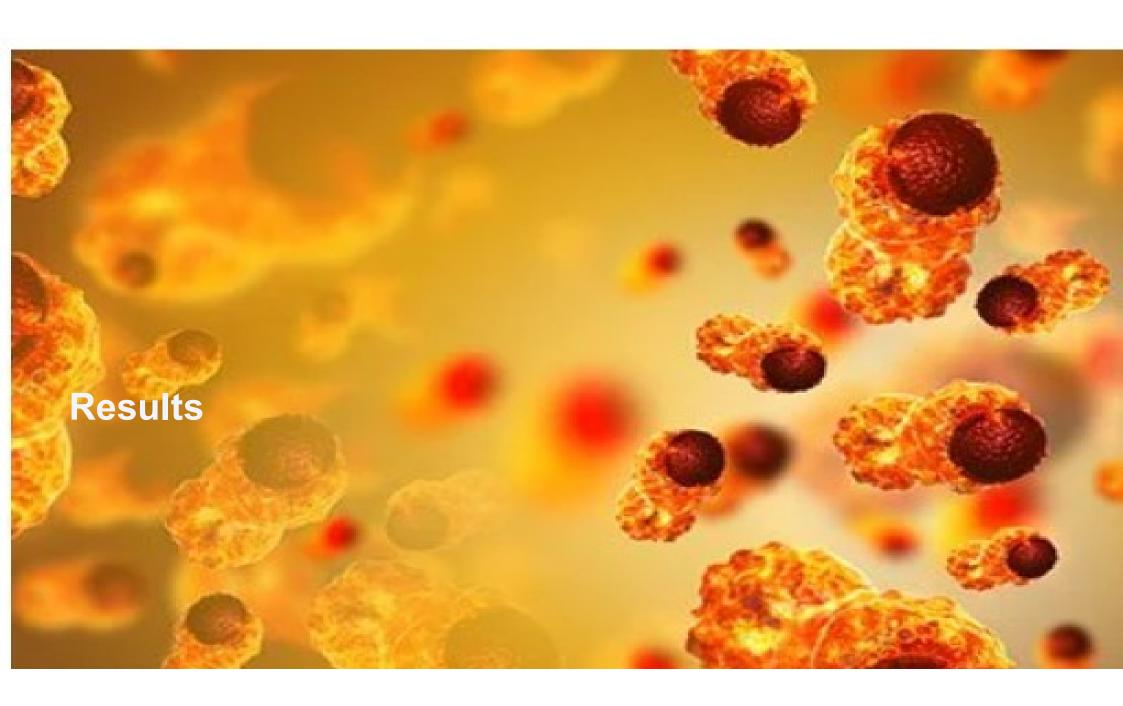
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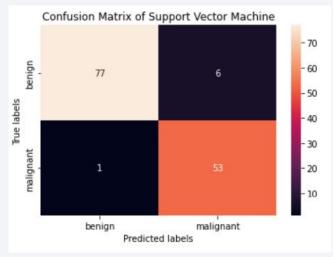
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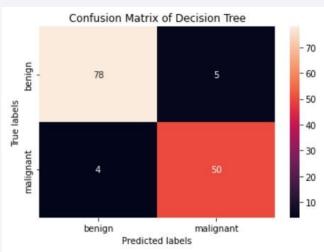
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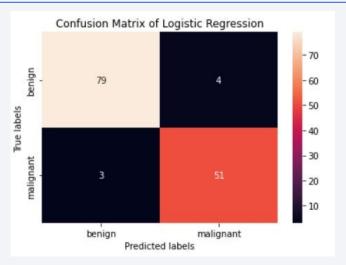
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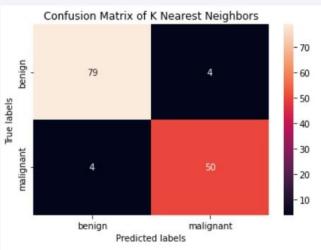


### **Results- Confusion Matrix**









### Results and Conclusion

- According to our predictive analyses, the best scores belong to Logistic Regression (LR) and Support Vector Machine(SVM).
   The best parameters for these two models are:
  - SVM
     C=0.031, gamma=1.0, Kernel=Sigmoid
  - LR
     C= 1, penalty= I2, solver= lbfgs

Accuracy	Model	
0.948905	logistic regression	0
0.948905	support vector machine	1
0.934307	decision tree	2
0.941606	k nearest neighbors	3

# **Appendix**

• Github link of the notebook file:

https://github.com/Amorovati/Cancer\_Prediction