Capstone 3: Project Report

Del Wester

4/06/2021

Breast Cancer

Most types of breast cancer are easy to diagnose by microscopic analysis of a sample - or biopsy - of the affected area of the breast. The two most commonly used screening methods, physical examination of the breasts by a healthcare provider and mammography, can offer an approximate likelihood that a lump is cancer, and may also detect some other lesions, such as a simple cyst. When these examinations are inconclusive, a healthcare provider can remove a sample of the fluid in the lump for microscopic analysis (a procedure known as fine needle aspiration, or fine needle aspiration, FNA) to help establish the diagnosis. A needle aspiration can be performed in a healthcare provider's office or clinic. Together, physical examination of the breasts, mammography, and FNA can be used to diagnose breast cancer with a good degree of accuracy.

The features for this dataset are computed from a digitized image of a fine needle aspirate (FNA) of a breast mass. They describe characteristics of the cell nuclei present in the image. This dataset was obtained from the University of Wisconsin Hospitals, Madison from https://archive.ics.uci.edu/ml/datasets/breast+cancer+wisconsin+(diagnostic).

Data Wrangling

I loaded the dataset into Excel and replaced the numbered columns with the column names. I then saved as a csv file and loaded into a pandas dataframe. There are 699 records in this dataset.

Number of Attributes: 10 + output attribute

Attribute information: except for ID and Class, all columns had values ranging from 1 – 10.

Input variables:

1 - ID

2 – Clump_Thickness

3 - Uniformity_of_Cell_Size

4 - Uniformity_of_Cell_Shape

5 - Marginal_Adhesion

6 - Single_Epithelial_Cell_Size

7 - Bare_Nuclei

8 - Bland_Chromatin

9 - Normal_Nucleoli

10 - Mitoses

Output variable:

11 - Class (2 for benign, 4 for malignant)

Missing Attribute Values: Bare_Nuclei was missing 16 values which I replaced with the mean.

Exploratory Data Analysis

The Breast Cancer data includes mostly continuous data with a single categorical column. . Exploratory data analysis was used to derive relationships between the class and the various features available from the data profile. The Class feature of this set was determined by the image of a fine needle aspirate (FNA) of a breast mass and takes into consideration the characteristics of the cell nuclei present in the image.
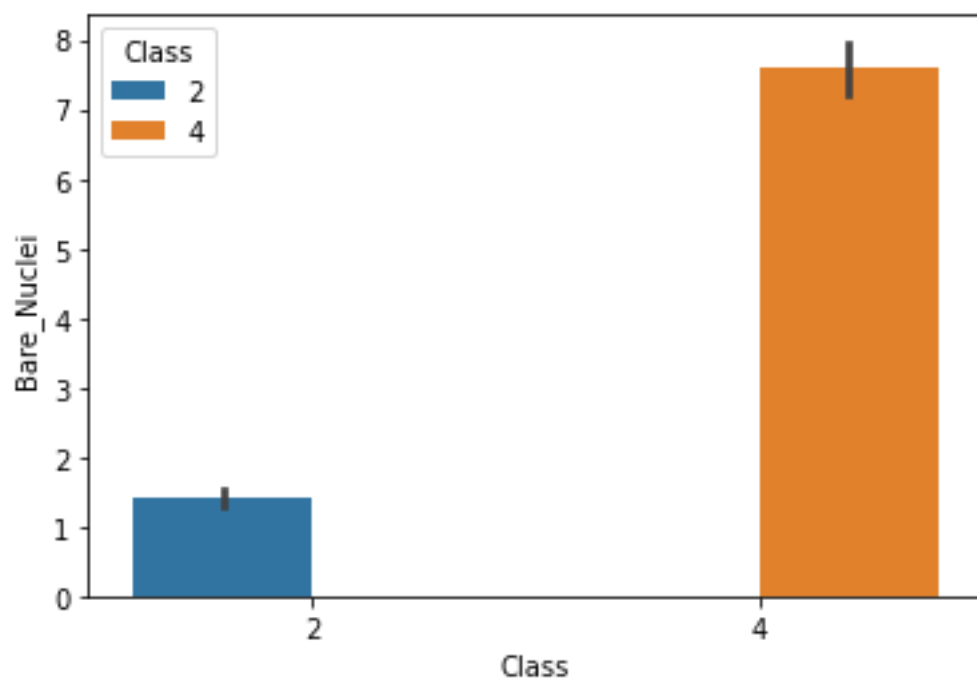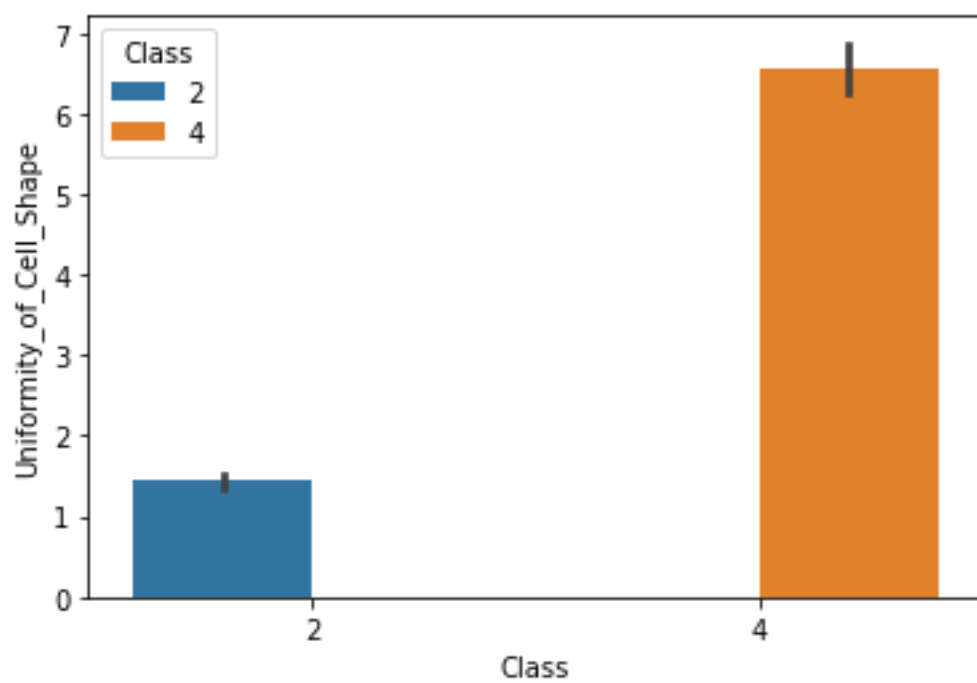
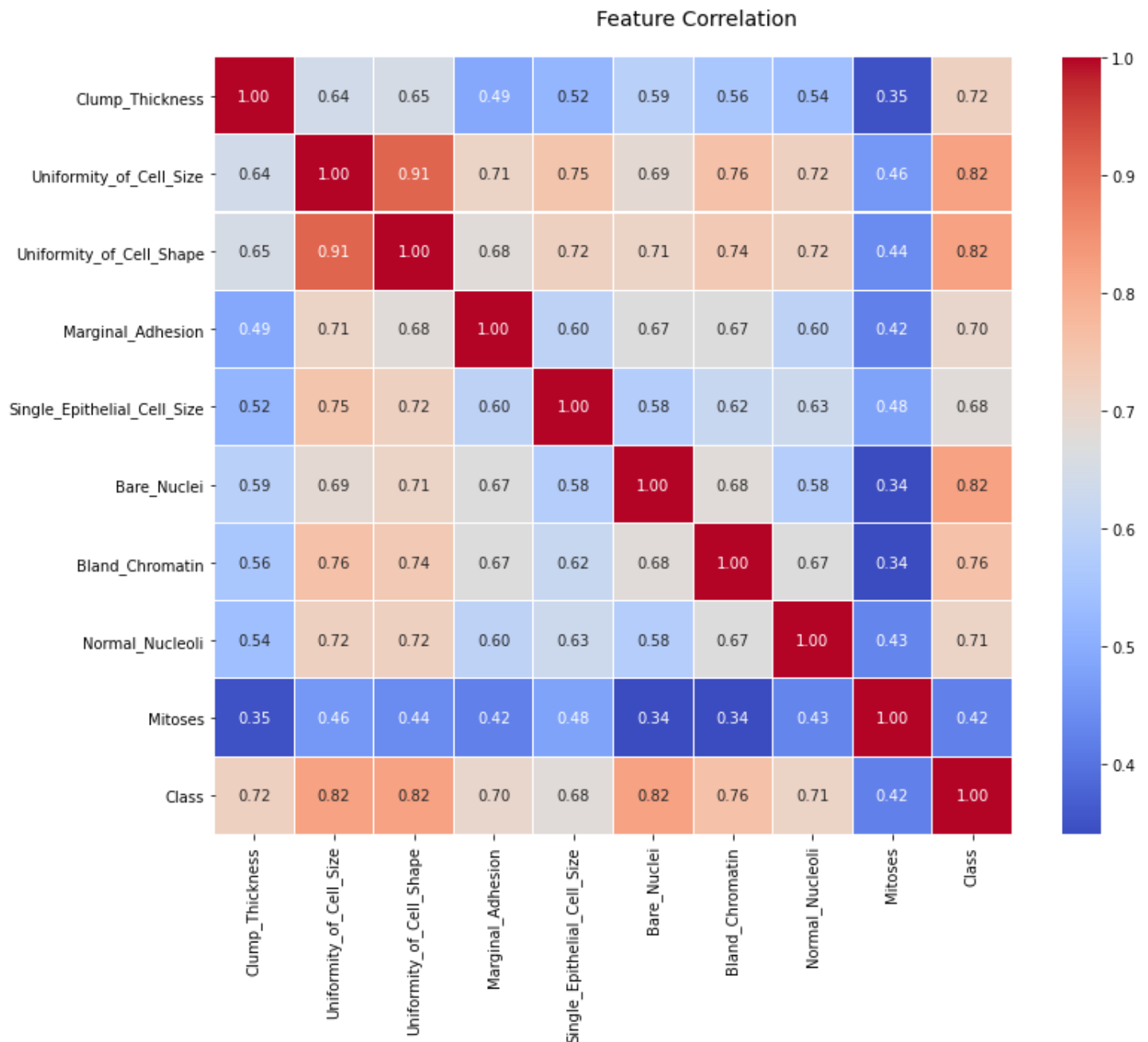"Class" Distribution:

Benign: 458

Malignant: 241
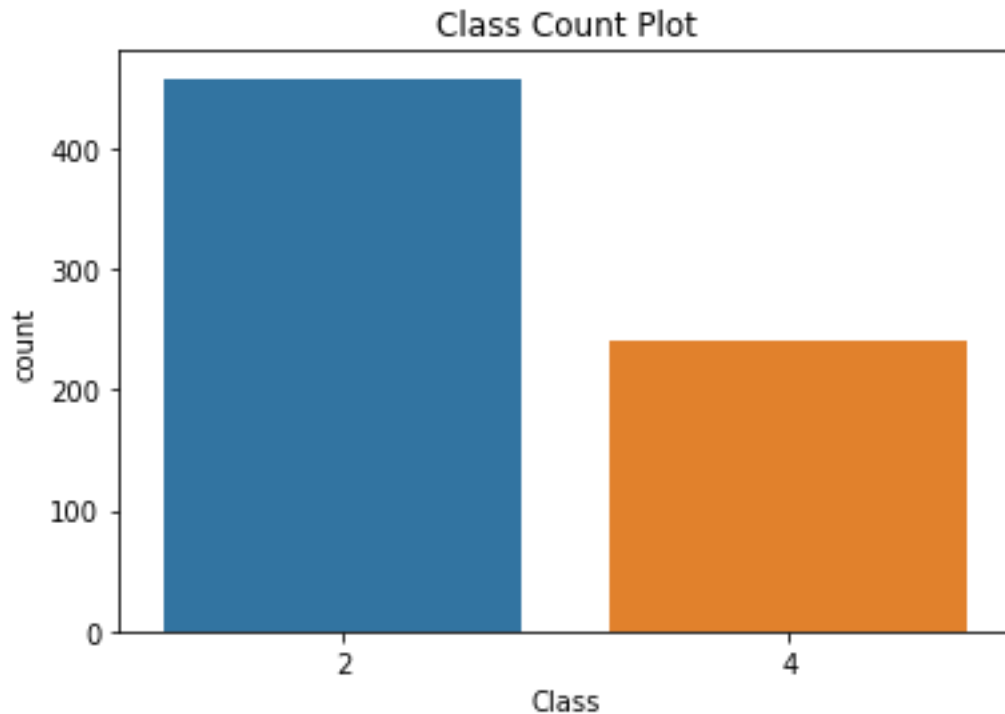
The plots for the output variable vs each of the top features are below.

Correlation Matrix

Feature Correlation

Almost perfect linear patterns between the Uniformity_of_Cell_Size and Uniformity_of_Cell_Shape features are highly linearly related (multicollinearity). Multicollinearity is a problem as it undermines the significance of independent variables. It can be fixed by removing the highly correlated features from the model using regression methods that cut the number of predictors to a smaller set of uncorrelated components, such as Partial Least Squares or Principal Components Analysis. I simply removed one of the colinear features before modeling.

There's one other feature that has colinearity with other features, and that's Mitosis, which is colinear with Clump_thickness, Bare_Nuclei, and Bland_Chromatin. So I drop both Uniformity_of_Cell_Size and Mitosis and re-run the heatmap.
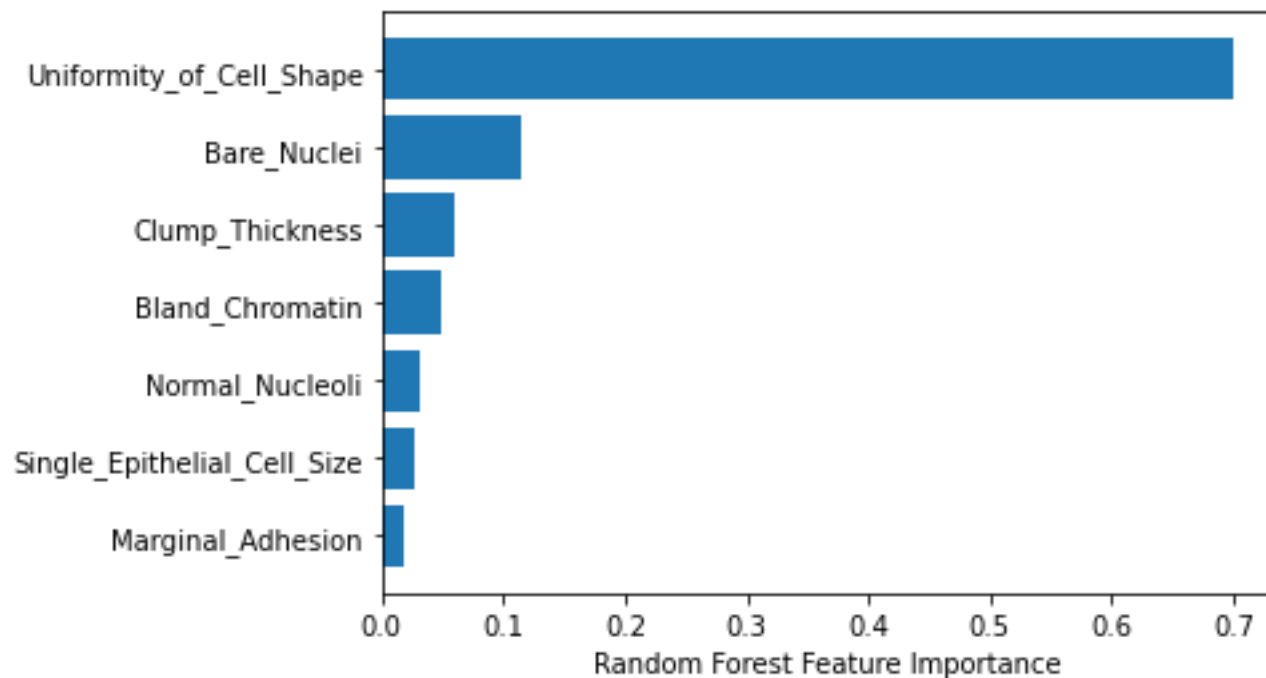
Feature Correlation
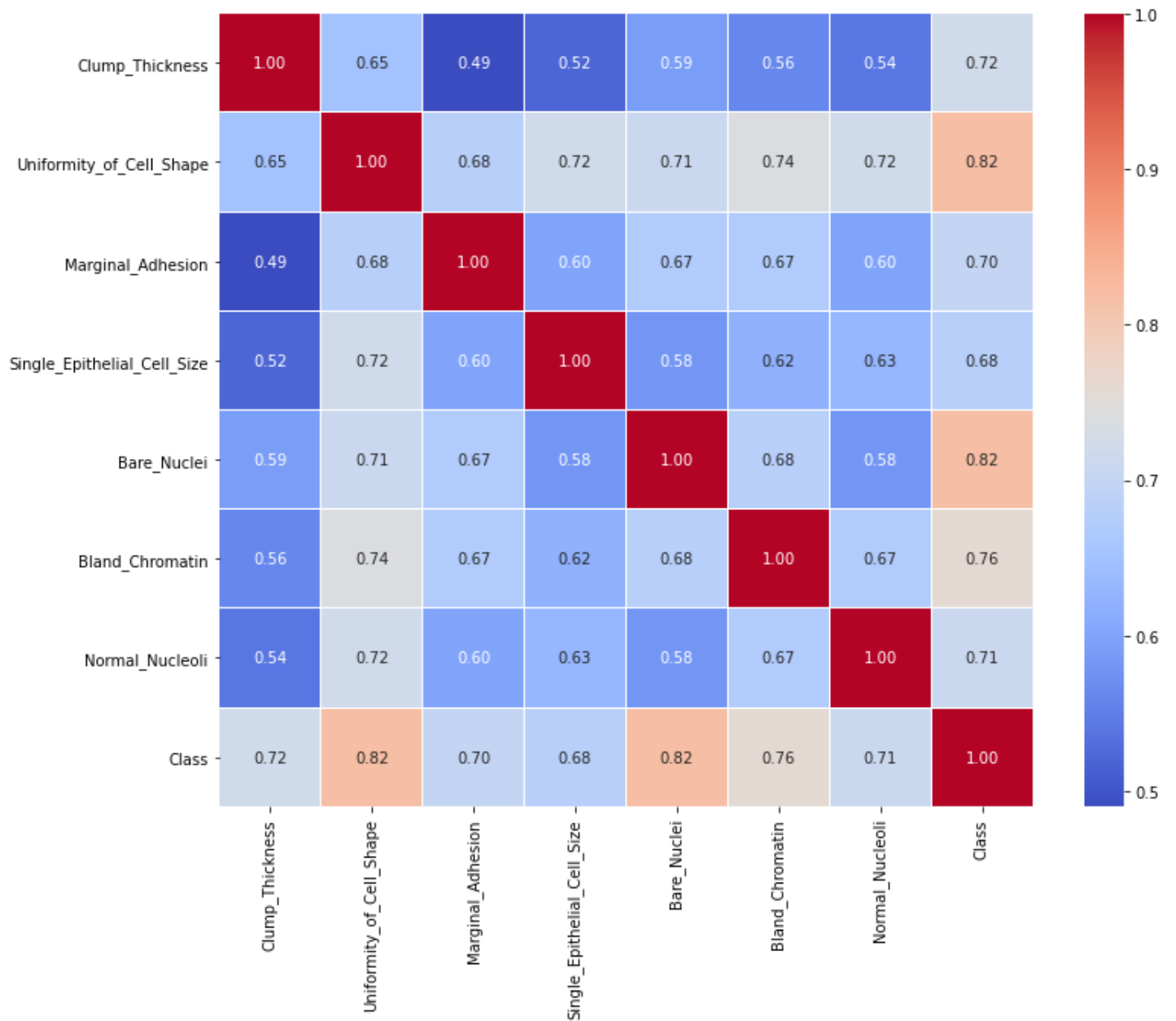
Now we're sitting pretty for the next step.

Preprocessing and Training

In this section, the data was scaled using a Robust Scaler and I ran a Random Forest Regressor to find the Feature Importance which is displayed below.

Top Features - Uniformity_of_Cell_Shape and Bare_Nuclei.

Modeling

I tested and scored the following models: RandomForestClassifier, KNeighborsClassifier, and Logistic Regression. I used GridsearchCV hyper parameter tuning for the models as well. I then ran the ROC_AUC scores/curve and Classification Reports. Logistic Regression scored higher than the others and was what I recommend for the final model.

ROC-AUC Scores/Curve

Logistic Regression   99.6
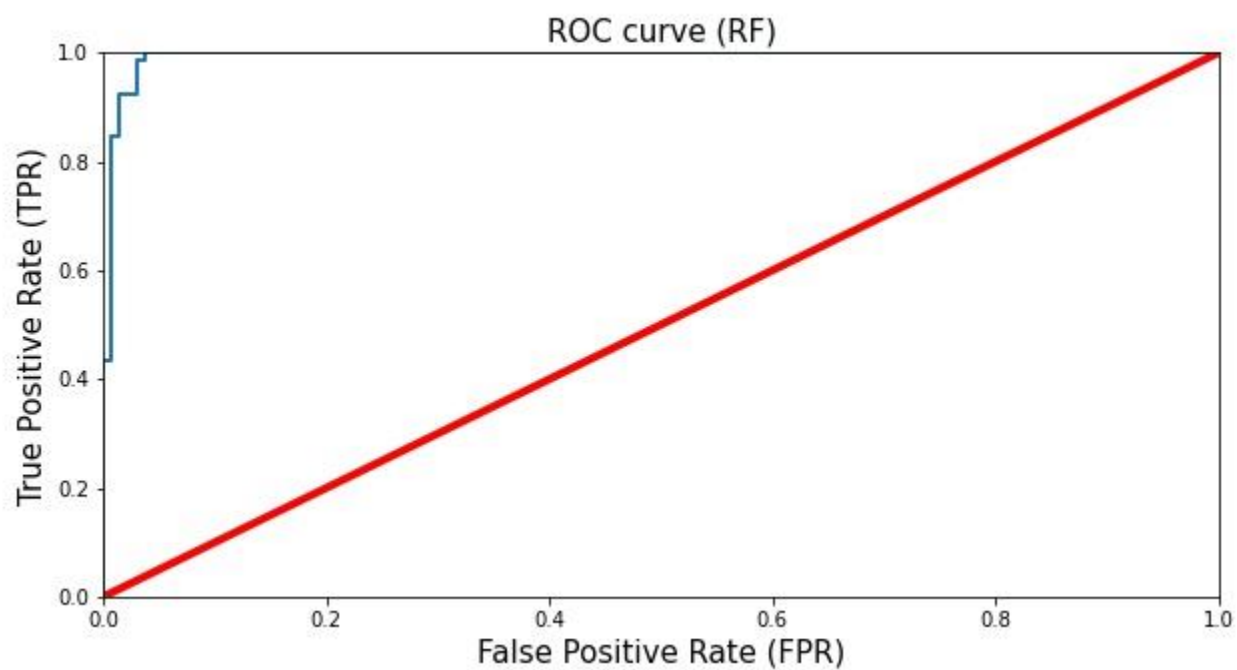

ROC curve (RF)
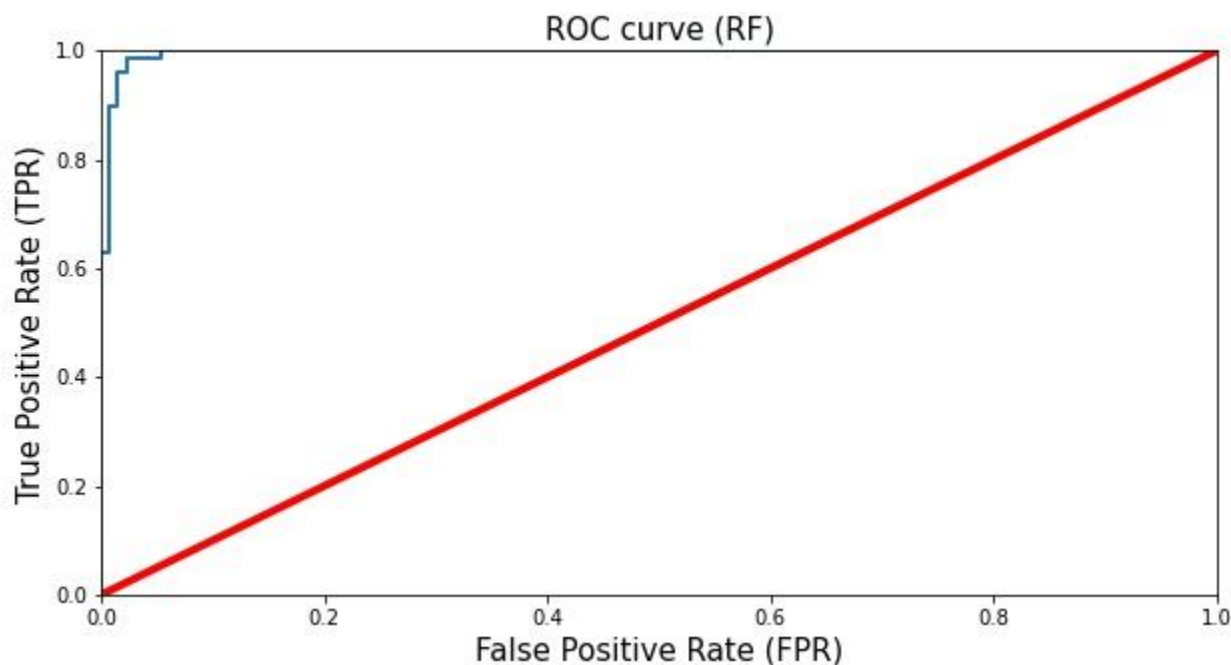
Random Forest Classifier   99.4

ROC curve (RF)
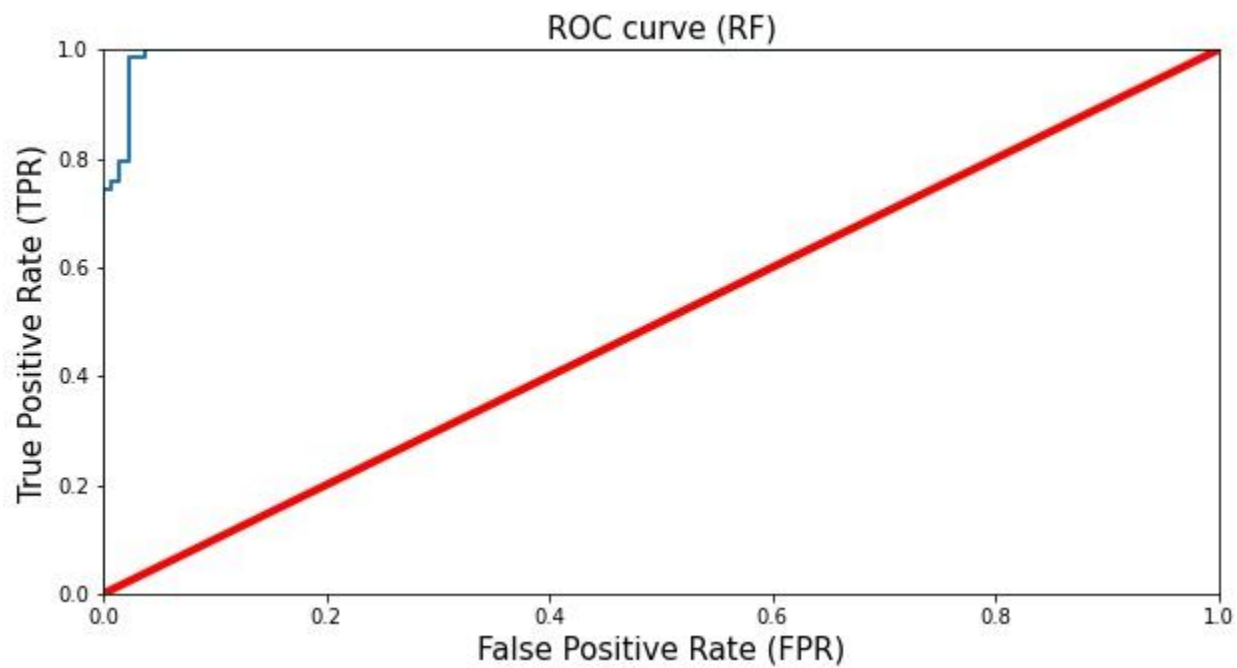
KNeighbors Classifier   99.3

ROC curve (RF)

A very high TPR is good considering you don't want to tell someone they don't have cancer when they do.

Classification Reports

Logistic Regression

Class	precision	recall	f1-score	support
2	0.99	0.98	0.98	132
4	0.96	0.99	0.97	78
accuracy			0.98	210
macro avg	0.98	0.98	0.98	210
weighted avg	0.98	0.98	0.98	210

Random Forest Classifier

Class	precision	recall	f1-score	support
2	0.99	0.96	0.98	132
4	0.94	0.99	0.96	78
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KNeighbors Classifier

Class	precision	recall	f1-score	support
2	0.97	0.97	0.97	132
4	0.95	0.95	0.95	78
accuracy			0.96	210
macro avg	0.96	0.96	0.96	210
weighted avg	0.96	0.96	0.96	210

Precision - percent of predictions correct.

Recall – percent of positive cases identified.

F1 Score – percent of positive predictions correct.

Support – number of actual cases for that Class type.

Conclusion

There are different techniques that can be used for the prediction of breast cancer. In this paper, I've analyzed breast cancer data using three classification techniques to predict the type of cancer and compared the results. They indicate that Logistic Regression is the best classifier with this dataset, followed by Random Forest and KNeighbors. Further studies could be conducted to improve performance of these classification techniques by identifying more features that could be used in the analysis.

Sources & Citations:

University of Wisconsin Hospitals, Madison from
https://archive.ics.uci.edu/ml/datasets/breast+cancer+wisconsin+(diagnostic)

https://en.wikipedia.org/wiki/Breast_cancer#Diagnosis