## **Dev Joshi**

## **Final Report : Benign or Malignant Cancer Prediction**

## **Problem Statement**

The dataset consisting of digitized imaging of fine needle aspirate (FNA) of a breast tumor cell mass can be used to distinguish between a benign and malignant tumor to aid in clinical diagnosis. Each cell nucleus has ten real-valued features which describe characteristics of the cell nuclei present in the image. The project goal is to deploy machine learning algorithms to accurately distinguish between a benign and malignant tumor to aid in clinical diagnosis. The original dataset is available at the <a href="https://archive.ics.uci.edu/ml/datasets/Breast+Cancer+Wisconsin+Diagnostic">https://archive.ics.uci.edu/ml/datasets/Breast+Cancer+Wisconsin+Diagnostic</a>.

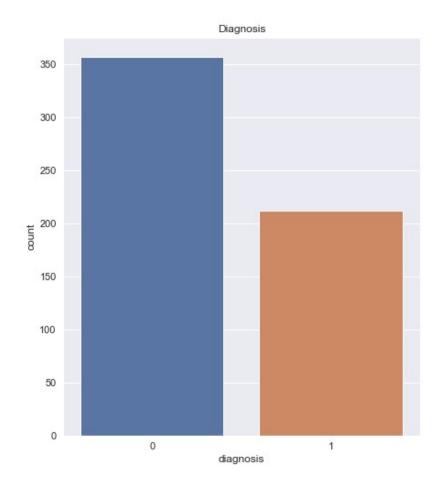
**<u>Data Wrangling:</u>** We began by downloading the dataset from the above link. The dataset consisted of 569 rows and 33 columns. We removed two columns – 'id' and 'Unnamed: 32' as they weren't essential for the analysis. The various columns of the dataset are:

- 0 diagnosis
- 1 radius mean
- 2 texture mean
- 3 perimeter mean
- 4 area mean
- 5 smoothness mean
- 6 compactness mean
- 7 concavity mean
- 8 concave points mean
- 9 symmetry mean
- 10 fractal dimension mean
- 11 radius se
- 12 texture se
- 13 perimeter se
- 14 area se
- 15 smoothness se
- 16 compactness se
- 17 concavity se
- 18 concave points se
- 19 symmetry se
- 20 fractal dimension se
- 21 radius worst

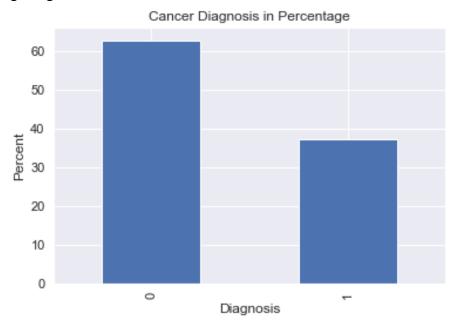
- 22 texture\_worst
- 23 perimeter\_worst
- 24 area worst
- 25 smoothness worst
- 26 compactness worst
- 27 concavity worst
- 28 concave points\_worst
- 29 symmetry\_worst
- 30 fractal\_dimension\_worst

We explored the data and looked at various columns of the dataframe. We made the target variable (column) 'diagnosis' as numeric by assigning '1' as Malignant and '0' as benign.

We plotted this column to observe that the number of the benign cases are 357 and the number of malignant cases are 212.

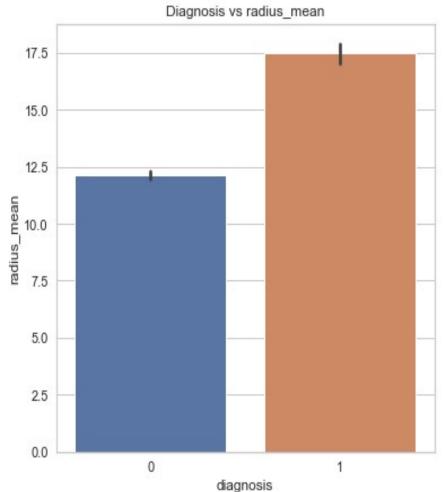


The same plot in terms of percentage yielded 62.7 % and 37.3 % for 0 (benign) and 1(malignant) respectively as seen in the following image.

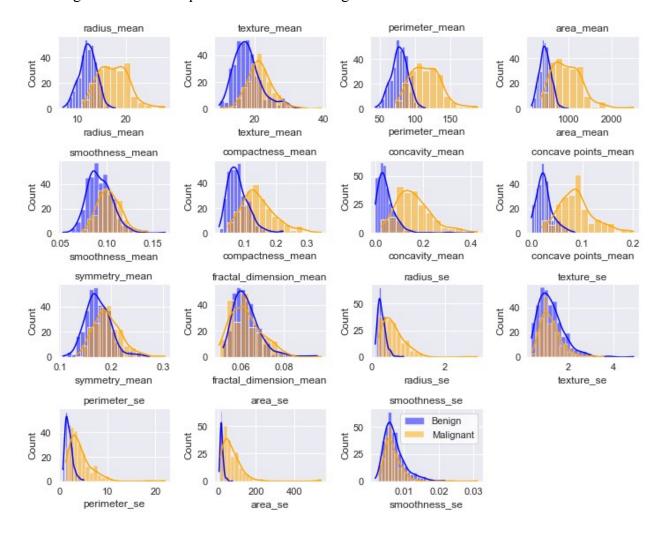


**Exploratory Data Analysis:** After initial data wrangling, we sought to explore the various features of the data. We looked at the various tumor features in relation to the diagnosis. For

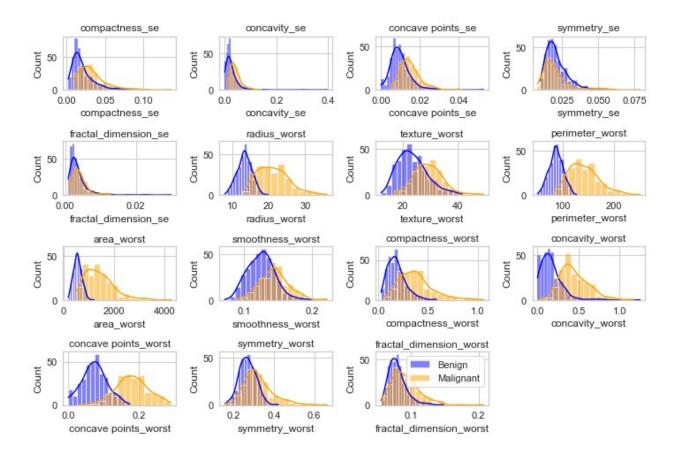
example, we see that the malign cells have greater mean radius than benign cell as seen in the adjacent image.



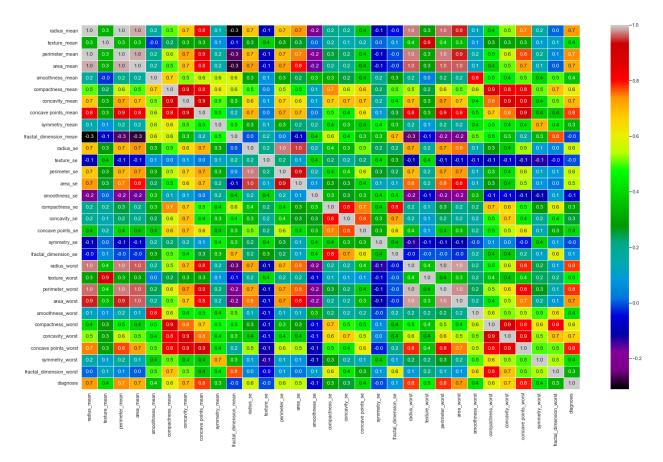
Similarly, the other features in the data can also be compared as seen in the image below. From this image also, we see that the mean radius is greater for the malign cells as compared to the benign cells (1<sup>st</sup> subplot). For most of the features (except few – namely – symmetry\_mean, fractal\_dimension\_mean, texture\_se, smoothness\_se), the feature corresponding to the malign cells has higher value as compared to that of the benign cells.



We plotted similar image for the remaining features as well. Here also, we see the malign cells have higher value, although lesser count, than the benign cells for most of the features.



We also checked the correlation map with respect to the diagnosis feature. As we can see in the image below, it has a low correlation - but positive - with most of the features. It has negative correlation with few of the features. Lastly, we saved the data for further analysis in the subsequent units.



## **Preprocessing, Training and Modeling:**

We downloaded the saved data from the previous section. We then divided the data into those that will be used to train the model and those that will be used to predict the approval: 70 % for training and 30 % for testing. We also applied standard scaling for X\_train and X\_test data so that no feature (with larger values) dominates over others (with smaller values).

