EDA - Haberman Dataset

February 6, 2022

Let's import useful packages to do EDA on the dataset.

```
[1]: import warnings
   warnings.filterwarnings('ignore')

[2]: from IPython.display import display
   from matplotlib import pyplot as plt
   from matplotlib import style

[3]: import numpy as np
   import pandas as pd
   import seaborn as sns

[4]: style.use(style='seaborn-whitegrid')

[5]: pd.set_option('display.precision', 4)
```

Dataset

The dataset contains cases from a study that was conducted between 1958 and 1970 at the University of Chicago's Billings Hospital on the survival of patients who had undergone surgery for breast cancer.

Download the Haberman Dataset from here.

Kaggle API: kaggle datasets download -d gilsousa/habermans-survival-data-set Now let's read the dataset.

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0	30	64	1	1
1	30	62	3	1
2	30	65	0	1
3	31	59	2	1
4	31	65	4	1
5	33	58	10	1

6	33	60	0	1
7	34	59	0	2
8	34	66	9	2
9	34	58	30	1

If we notice, **age** and **op_year** values are bit confusing. And we know from the description of the dataset at here, that **op_year** happens to be in 1900's.

Let's preprocess the op_year feature by adding 1900 to each data-point.

```
[8]: data_df['op_year'] = data_df['op_year'].apply(func=lambda x: 1900 + x)
[9]: display(data_df.head(n=10))
```

	1 ,	, · –		
	age	op_year	axil_nodes	surv_status
0	30	1964	1	1

0	30	1964	1	1
1	30	1962	3	1
2	30	1965	0	1
3	31	1959	2	1
4	31	1965	4	1
5	33	1958	10	1
6	33	1960	0	1
7	34	1959	0	2
8	34	1966	9	2
9	34	1958	30	1

Now the dataset looks understandable and cleaner.

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Some background understanding.

What are Axillary Nodes?

The axillary lymph nodes or armpit lymph nodes are lymph nodes in the human armpit. Between 20 and 49 in number, they drain lymph vessels from the lateral quadrants of the breast, the superficial lymph vessels from thin walls of the chest and the abdomen above the level of the navel, and the vessels from the upper limb. They are divided in several groups according to their location in the armpit. These lymph nodes are clinically significant in breast cancer, and metastases from the breast to the axillary lymph nodes are considered in the staging of the disease.

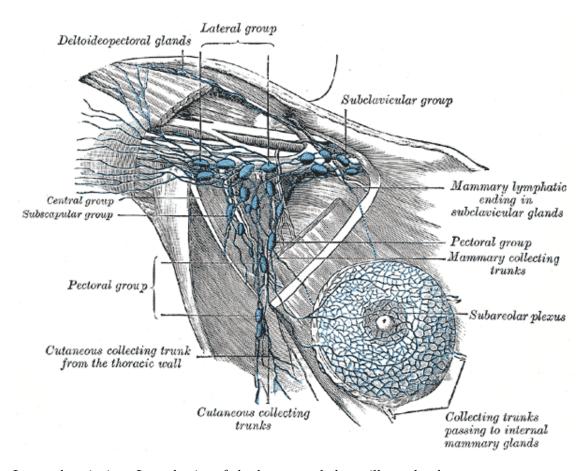


Image description: Lymphatics of the breast and the axillary glands

Image source: here

1.1 Analyze high level statistics of the dataset: number of points, number of features, number of classes, data-points per class

[10]: data_df.info()

<class 'pandas.core.frame.DataFrame'>
RangeIndex: 306 entries, 0 to 305
Data columns (total 4 columns):

#	Column	Non-Null Count	Dtype
0	age	306 non-null	int64
1	op_year	306 non-null	int64
2	axil_nodes	306 non-null	int64
3	surv_status	306 non-null	int64

dtypes: int64(4) memory usage: 9.7 KB

From the above output, we learn that, dataset has 4 attributes/columns which includes class attribute as well.

- age: Age of patient at time of operation.
- op_year: Patient's year of operation.
- axil_nodes: Number of positive axillary nodes detected.
- surv_status: Survival status. This is a class attribute, encoded as below,
 - -1 = the patient survived 5 years or longer
 - -2 = the patient died within 5 year

There are 306 rows. All the columns are of numerical type. We also learn that the dataset doesn't have any NULL values, which is good.

Features and Target(s)

We will now assign features and class-label (target) for ease of understanding and we shall use these features and the target in our further analysis.

```
[11]: features = list(data_df.columns)[:-1]
  target = list(data_df.columns)[-1]

print("All the features in the dataset: {}".format(features))
print("Target class in the dataset: '{}'".format(target))
```

```
All the features in the dataset: ['age', 'op_year', 'axil_nodes'] Target class in the dataset: 'surv_status'
```

Now let's understand how many data-points we have in surv_status target.

[12]: display(data_df[[target]].value_counts())

This dataset is an **imbalanced** dataset, because number of data-points of the class 1 is not equal to number of data-points of the class 2.

Let's calculate basic descriptive statistics for our dataset features.

[13]: display(data_df[features].describe())

	age	op_year	axil_nodes
count	306.0000	306.0000	306.0000
mean	52.4575	1962.8529	4.0261
std	10.8035	3.2494	7.1897
min	30.0000	1958.0000	0.0000
25%	44.0000	1960.0000	0.0000
50%	52.0000	1963.0000	1.0000
75%	60.7500	1965.7500	4.0000
max	83.0000	1969.0000	52.0000

age feature

• There are 306 data-points.

- The average age is 52.457 years.
- The standard deviation is 10.803.
- The minimum age in the dataset is 30 years.
- The 25^{th} percentile is 44 years, which implies 25% of the patients' age are ≤ 44 years.
- The 50^{th} percentile is 52 years, which implies 50% of the patients' age are ≤ 52 years.
- The 75th percentile is 60.75 years, which implies 75% of the patients' age are ≤ 60.75 years.
- The maximum age in the dataset is 83 years.

op_year feature

- There are 306 data-points.
- The average year date is 1962.853.
- The standard deviation is 3.249.
- The minimum operation year date in the dataset is 1958.
- The 25th percentile is 1960 year date, which implies 25% of the patients' operation year dates are < 1960.
- The 50^{th} percentile is 1963 year date, which implies 50% of the patients' operation year dates are ≤ 1963 .
- The 75^{th} percentile is 1965.75 year date, which implies 75% of the patients' operation year dates are < 1965.75.
- The maximum operation year date in the dataset is 1969.

axil_nodes feature

- There are 306 data-points.
- The average number of axillary nodes are 4.02.
- The standard deviation is 7.189.
- The minimum number of axillary nodes are 0.
- The 25^{th} percentile is 0, which implies 25% of the patients' axillary nodes in the body are ≤ 0 .
- The 50^{th} percentile is 1, which implies 50% of the patients' axillary nodes in the body are < 1.
- The 75^{th} percentile is 4, which implies 75% of the patients' axillary nodes in the body are < 4.
- The maximum number of axillary nodes are 52.

We will see this visually in the report.

Now, let's find the most repeated values for each feature in the dataset.

[14]: display(data_df[features].mode())

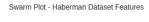
```
age op_year axil_nodes
0 52 1958 0
```

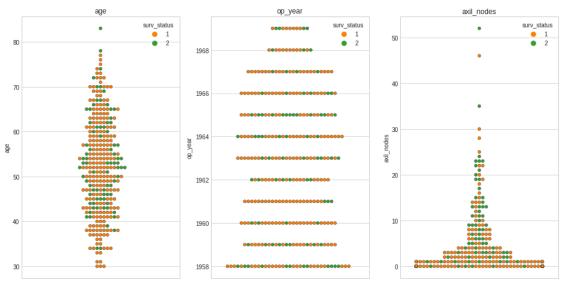
Let's understand a bit more, using seaborn.swarmplot().

```
[15]: def swarm_plotter(df, features):
```

This is funtion helps in plotting the dataframes.

[16]: swarm_plotter(df=data_df, features=features)





- From age feature, we observe, 52 is repeated quite often and is nearly equal to the average which we found earlier. There is one outlier in this feature, where the patient's age is 83 years.
- From op_year feature, we observe, more operations took place in the year date 1958. There is no sign of outliers in this feature.
- From axil_nodes feature, we observe, 0 and 1 axillary nodes are most common. We also observe couple of outliers present in this feature.

1.2 Explain the objective of the problem

Objective

- To understand the dataset and convert the raw data into useful reports/information.
- Reports of useful information can be obtained from the raw data by performing descriptive statistics, univariate and multivariate analysis.
- To articulate the reports with the rest of the team in an understandable way.

1.3 Perform univariate analysis - Plot PDF, CDF, Box plots

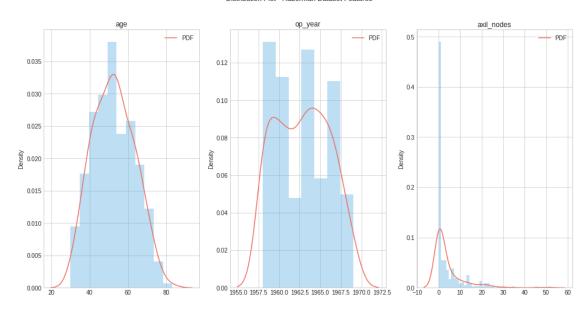
- Plot the required charts to understand which feature are important for classification.
- Make sure that you add titles, legends and labels for each and every plots.
- Suppress the warnings you get in Python, in that way it makes your notebook more presentable.
- Do write observations/inference for each plot.

Distribution Plot

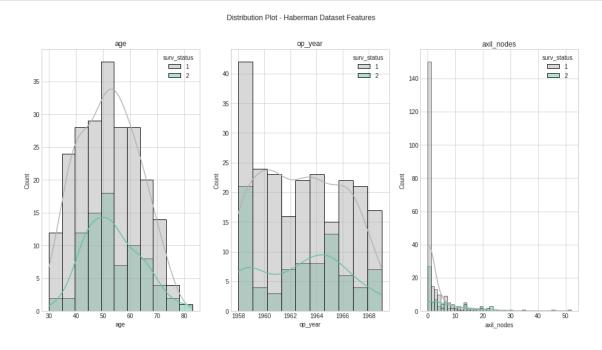
```
[17]: def dist_plotter(df, features, hue=False):
          This is funtion helps in plotting the dataframes.
          plt.figure(figsize=(16, 8))
          for (i, feature) in zip(range(len(features)), features):
              plt.subplot(int('13' + '{}'.format(i+1)))
              if hue:
                  sns.histplot(x=df[feature],
                                hue=df[target],
                                kde=True,
                                palette='Set2 r')
              else:
                  sns.distplot(x=df[feature],
                                kde=True,
                                hist_kws={'color': '#5DADE2'},
                                kde_kws={'color': '#EC7063', 'label': 'PDF'})
                  plt.legend()
              plt.title(label=feature)
              i += 1
          plt.suptitle(t='Distribution Plot - Haberman Dataset Features')
          plt.show()
          plt.close(fig='all')
          return None
```

```
[18]: dist_plotter(df=data_df, features=features, hue=False)
```

Distribution Plot - Haberman Dataset Features



[19]: dist_plotter(df=data_df, features=features, hue=True)



The histogram plot can also be combined with KDE to plot the PDF curve for each feature.

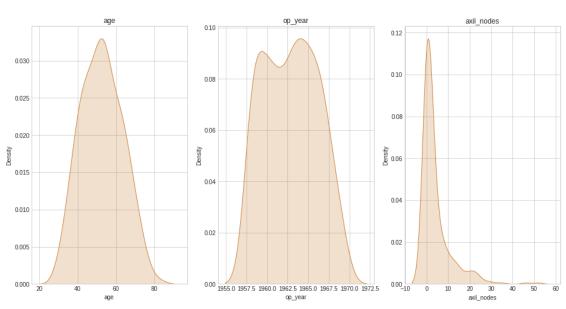
• From the above Distribution plot, we observe that, no feature is well separating the surv_status class attribute. We also see a lot of overlap between the surv_status classes.

PDF Plot

```
[20]: def pdf_plotter(df, features, hue=False):
          This is funtion helps in plotting the dataframes.
          plt.figure(figsize=(16, 8))
          for (i, feature) in zip(range(len(features)), features):
              plt.subplot(int('13' + '{}'.format(i+1)))
              if hue:
                  sns.kdeplot(x=df[feature],
                              hue=df[target],
                              shade=True,
                              palette='rainbow')
              else:
                  sns.kdeplot(x=df[feature],
                              shade=True,
                              color='peru')
              plt.title(label=feature)
              i += 1
          plt.suptitle(t='PDF Plot - Haberman Dataset Features')
          plt.show()
          plt.close(fig='all')
          return None
```

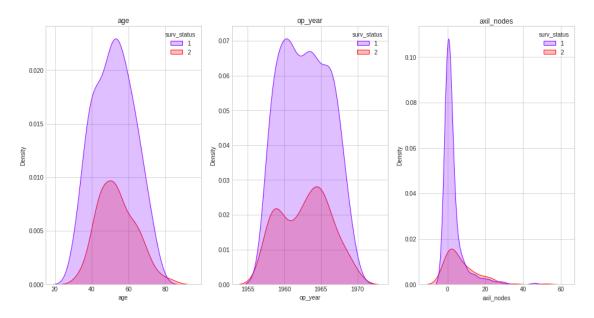
[21]: pdf_plotter(df=data_df, features=features, hue=False)





[22]: pdf_plotter(df=data_df, features=features, hue=True)





The above two plots are a PDF (Probability Density Function) plot with hue turned off and on.

• From the 2^{nd} PDF plot, we see that, none of the features separate the class attributes. Both the class attributes are overlapping. Hence it will be hard which determine which feature is good and which is not.

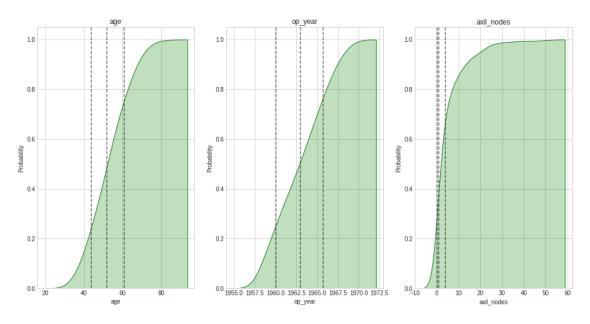
CDF Plot

```
[23]: def cdf_plotter(df, features):
          11 11 11
          This is funtion helps in plotting the dataframes.
          plt.figure(figsize=(16, 8))
          for (i, feature) in zip(range(len(features)), features):
              plt.subplot(int('13' + '{}'.format(i+1)))
              sns.kdeplot(x=df[feature],
                           shade=True,
                           cumulative=True,
                           color='g')
              plt.ylabel(ylabel='Probability')
              plt.title(label=feature)
              percentiles = np.percentile(a=df[feature],
                                           q=np.array([25, 50, 75]))
              for quantile in percentiles:
                  plt.axvline(x=quantile, ls='--', c='k', alpha=0.6)
              i += 1
```

```
plt.suptitle(t='CDF Plot - Haberman Dataset Features')
plt.show()
plt.close(fig='all')
return None
```

[24]: cdf_plotter(df=data_df, features=features)

CDF Plot - Haberman Dataset Features



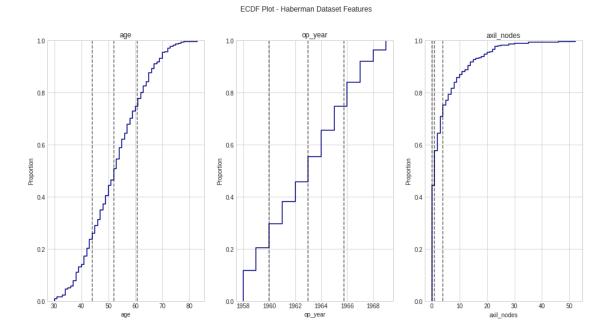
This is a cumulative distribution function plot of our features. On the Y-axis, CDF plot shows probability values upto to certain point on the X-axis. We also see three vertical lines that represent the quantiles.

- From age CDF plot, we observe that 50% of the patient's age lies between 30-52 years.
- From op_year CDF plot, we observe that 50% of patient's operations took place between 1958-1963 year dates.
- From axil nodes CDF plot, we oberve that 50% of patients' axillary nodes are upto 1.

ECDF Plot

```
[25]: def ecdf_plotter(df, features):
    """
    This is funtion helps in plotting the dataframes.
    """
    plt.figure(figsize=(16, 8))
    for (i, feature) in zip(range(len(features)), features):
        plt.subplot(int('13' + '{}'.format(i+1)))
        sns.ecdfplot(x=df[feature], color='navy')
        plt.title(label=feature)
```

```
[26]: ecdf_plotter(df=data_df, features=features)
```



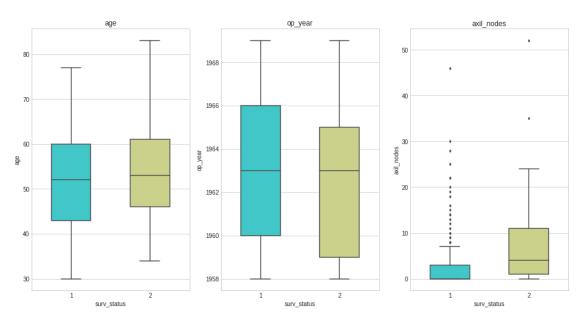
ECDF stands of Empirical Cumulative Distribution Function. Unlike Swarm plot which represents only count, ECDF plot represents the proportion or count of observations falling below each unique value in a dataset. We observe, 52, 1958 and 0 are most common (or modal) values in age, op_year, axil_nodes features.

Box Plot

```
[27]: def box_plotter(df, features):
    """
    This is funtion helps in plotting the dataframes.
    """
    plt.figure(figsize=(16, 8))
    for (i, feature) in zip(range(len(features)), features):
        plt.subplot(int('13' + '{}'.format(i+1)))
        sns.boxplot(x=target,
```

[28]: box_plotter(df=data_df, features=features)





The Box plot gives detailed visualization of quantiles, minimum and maximum values.

- From age, we observe that 50% of the patient's age lies between 30-52 years.
- From op_year CDF plot, we observe that 50% of patient's operations took place between 1958-1963 year dates.
- From axil_nodes CDF plot, we oberve that 50% of patients' axillary nodes are upto 1.

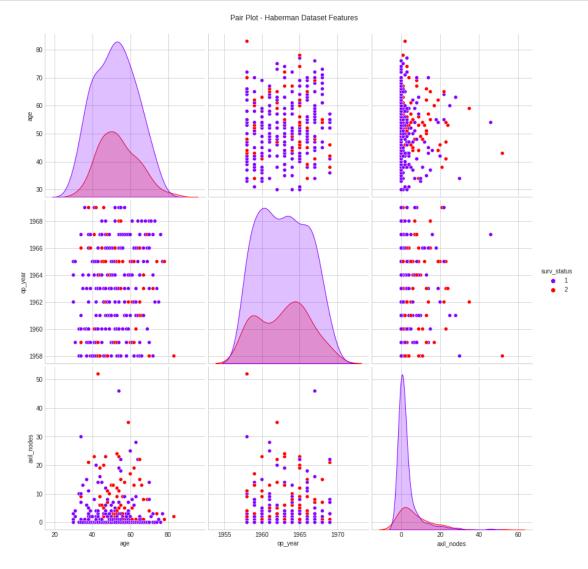
1.4 Perform bivariate analysis - Plot 2D Scatter plots and Pair plots

- Plot the required scatter plots and pair plots of different features to see which combination of features are useful for clasification task.
- Make sure that you add titles, legends and labels for each and every plots.

- Suppress the warnings you get in Python, in that way it makes your notebook more presentable.
- Do write observations/inference for each plot.

Pair Plot

```
[29]: g = sns.pairplot(data=data_df, hue=target, height=4, palette='rainbow')
    g.fig.suptitle(t='Pair Plot - Haberman Dataset Features', y=1.02)
    plt.show()
    plt.close(fig='all')
```



Pair plot represents pairwise relationships in a dataset. On the diagnols, univariate distribution plot is drawn to show the marginal distribution of the data in each column.

• We already observed diagnol PDF plots earlier. Every class in the surv_status target variable, overlaps with other class, hence it becomes harder to point-out which feature is fairly

separating each class and which is not.

• Also, if we notice the scatter plots, there is no strong correlation across the features.

1.5 Summarize your final conclusions of the exploration

- You can describe the key features that are important for the classification task.
- Try to quantify your results i.e., while writing observations include numbers, percentages, fractions etc.
- Write a brief of your exploratory analysis in 3-5 points.
- Write your observations in english as crisply and unambigously as possible.

Overall Conclusion

- Our dataset is an imbalanced dataset as the number of data-points of the classes for our target variable surv_status are not equal.
- Dataset has couple of outliers especially for axil_nodes feature. For some patients the count of axillary nodes cross 30.
- None of the features are fairly separating the classes of our target variable surv_status.

End of the file.	