→ Breast Cancer Analysis

!pip install dataprep

```
Requirement already satisfied: entrypoints>=0.2.2 in /usr/local/lib/python3.7/dist -
Requirement already satisfied: defusedxml in /usr/local/lib/python3.7/dist-package
Requirement already satisfied: bleach in /usr/local/lib/python3.7/dist-packages (f
Requirement already satisfied: testpath in /usr/local/lib/python3.7/dist-packages
Requirement already satisfied: pandocfilters>=1.4.1 in /usr/local/lib/python3.7/di
Requirement already satisfied: mistune<2,>=0.8.1 in /usr/local/lib/python3.7/dist-
Requirement already satisfied: webencodings in /usr/local/lib/python3.7/dist-packag
Building wheels for collected packages: metaphone, pystache, python-Levenshtein
  Building wheel for metaphone (setup.py) ... done
  Created wheel for metaphone: filename=Metaphone-0.6-py3-none-any.whl size=13918
  Stored in directory: /root/.cache/pip/wheels/1d/a8/cb/6f8902aa5457bd71344e00665c
  Building wheel for pystache (PEP 517) ... done
  Created wheel for pystache: filename=pystache-0.6.0-py3-none-any.whl size=83635
  Stored in directory: /root/.cache/pip/wheels/78/87/45/383bd15701a08a94c735e9eaf3
  Building wheel for python-Levenshtein (setup.py) ... done
  Created wheel for python-Levenshtein: filename=python_Levenshtein-0.12.2-cp37-cp
  Stored in directory: /root/.cache/pip/wheels/05/5f/ca/7c4367734892581bb5ff896f150
Successfully built metaphone pystache python-Levenshtein
Installing collected packages: jinja2, locket, Werkzeug, partd, multidict, itsdange
  Attempting uninstall: jinja2
    Found existing installation: Jinja2 2.11.3
    Uninstalling Jinja2-2.11.3:
      Successfully uninstalled Jinja2-2.11.3
  Attempting uninstall: Werkzeug
    Found existing installation: Werkzeug 1.0.1
    Uninstalling Werkzeug-1.0.1:
      Successfully uninstalled Werkzeug-1.0.1
  Attempting uninstall: itsdangerous
    Found existing installation: itsdangerous 1.1.0
    Uninstalling itsdangerous-1.1.0:
      Successfully uninstalled itsdangerous-1.1.0
  Attempting uninstall: click
    Found existing installation: click 7.1.2
    Uninstalling click-7.1.2:
      Successfully uninstalled click-7.1.2
  Attempting uninstall: regex
    Found existing installation: regex 2019.12.20
    Uninstalling regex-2019.12.20:
      Successfully uninstalled regex-2019.12.20
  Attempting uninstall: flask
    Found existing installation: Flask 1.1.4
    Uninstalling Flask-1.1.4:
      Successfully uninstalled Flask-1.1.4
  Attempting uninstall: dask
    Found existing installation: dask 2.12.0
    Uninstalling dask-2.12.0:
      Successfully uninstalled dask-2.12.0
  Attempting uninstall: wordcloud
    Found existing installation: wordcloud 1.5.0
    Uninstalling wordcloud-1.5.0:
      Successfully uninstalled wordcloud-1.5.0
  Attempting uninstall: nltk
```

```
Found existing installation: nltk 3.2.5
         Uninstalling nltk-3.2.5:
           Successfully uninstalled nltk-3.2.5
     ERROR: pip's dependency resolver does not currently take into account all the pack
     datascience 0.10.6 requires folium==0.2.1, but you have folium 0.8.3 which is incor
import numpy as np
import pandas as pd
import matplotlib.pyplot as plt
import seaborn as sns
from sklearn.decomposition import PCA
from dataprep.eda import create_report
from dataprep.eda import plot_missing
from dataprep.eda import plot_correlation
from dataprep.eda import plot
from sklearn.model_selection import train_test_split
from sklearn.metrics import f1_score, accuracy_score, confusion_matrix, classification_rep
from sklearn.model_selection import learning_curve, cross_val_score, GridSearchCV
from sklearn.model_selection import train_test_split
from sklearn.ensemble import RandomForestClassifier, AdaBoostClassifier
from sklearn.svm import SVC
from sklearn.pipeline import make pipeline
from sklearn.neighbors import KNeighborsClassifier
from sklearn.linear_model import LogisticRegression
from sklearn.preprocessing import RobustScaler,StandardScaler,MinMaxScaler
import warnings
warnings.filterwarnings('ignore')
import numpy as np
import pandas as pd
import matplotlib.pyplot as plt
import seaborn as sns
from sklearn.decomposition import PCA
from dataprep.eda import create report
from dataprep.eda import plot_missing
from dataprep.eda import plot correlation
from dataprep.eda import plot
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from sklearn.metrics import f1_score, accuracy_score, confusion_matrix, classification_rep
from sklearn.model_selection import learning_curve, cross_val_score, GridSearchCV
from sklearn.model selection import train test split
```

from sklearn.ensemble import RandomForestClassifier, AdaBoostClassifier

```
from sklearn.svm import SVC
from sklearn.pipeline import make_pipeline
from sklearn.neighbors import KNeighborsClassifier
from sklearn.linear_model import LogisticRegression
from sklearn.preprocessing import RobustScaler,StandardScaler,MinMaxScaler
import warnings
warnings.filterwarnings('ignore')
```

Dataset Analysis

```
data = pd.read_csv('BRCA.csv')
df = data.copy()
pd.set_option('display.max_row',df.shape[0])
pd.set_option('display.max_column',df.shape[1])
df.head()
```

	Patient_ID	Age	Gender	Protein1	Protein2	Protein3	Protein4	Tumour_Stage
0	TCGA-D8- A1XD	36	FEMALE	0.080353	0.42638	0.54715	0.273680	III
1	TCGA-EW- A1OX	43	FEMALE	-0.420320	0.57807	0.61447	-0.031505	11
2	TCGA-A8- A079	69	FEMALE	0.213980	1.31140	-0.32747	-0.234260	111
3	TCGA-D8- A1XR	56	FEMALE	0.345090	-0.21147	-0.19304	0.124270	II
4	TCGA-BH- A0BF	56	FEMALE	0.221550	1.90680	0.52045	-0.311990	II



(df.isna().sum()/df.shape[0]*100).sort_values(ascending=False)

```
Date_of_Last_Visit
                      5.089820
Patient_Status
                      3.892216
Patient_ID
                      0.000000
Age
                      0.000000
Gender
                      0.000000
Protein1
                      0.000000
Protein2
                      0.000000
Protein3
                      0.000000
Protein4
                      0.000000
```

Tumour Stage	0.000000
Histology	0.000000
ER status	0.000000
PR status	0.000000
HER2 status	0.000000
Surgery_type	0.000000
Date_of_Surgery	0.000000

dtype: float64

plot_missing(df)

Stats	Bar Chart	Spectrum	Heat Map	Dendrogram
		Missing Stati	stics	
Missing Cell	S			30
Missing Cell	s (%)			0.6%
Missing Colu	ımns			2
Missing Row	/S			17
Avg Missing	Cells per Column			1.88
Avg Missing	Cells per Row			0.09

```
plt.figure(figsize=(10,8))
sns.heatmap(df.isna(),cbar=False)
plt.show()
```

```
print('There is' , df.shape[0] , 'rows')
print('There is' , df.shape[1] , 'columns')

There is 334 rows
There is 16 columns
```

Checking for duplicates

```
df.duplicated().sum()

0

df.loc[df.duplicated(keep=False),:]
```

Patient_ID Age Gender Protein1 Protein2 Protein3 Protein4 Tumour_Stage H



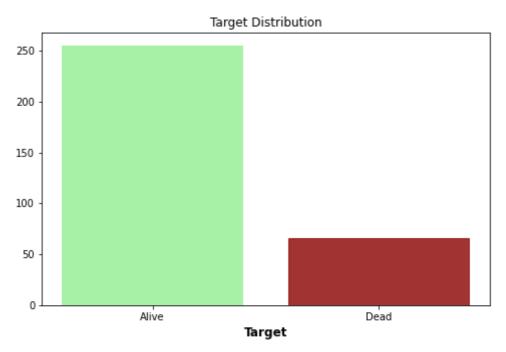
▼ Target Distribution

```
df = data.copy()
df = df.drop(['Patient_ID','Date_of_Surgery','Date_of_Last_Visit'],axis=1)
df['Patient_Status'].value_counts(normalize=True) #Classes déséquilibrées

Alive    0.794393
    Dead    0.205607
    Name: Patient_Status, dtype: float64

target_dist = df['Patient_Status'].value_counts()
fig, ax = plt.subplots(1, 1, figsize=(8,5))
barplot = plt.bar(target_dist.index, target_dist, color = 'lightgreen', alpha = 0.8)
barplot[1].set_color('darkred')

ax.set_title('Target Distribution')
percentage = df['Patient_Status'].value_counts(normalize=True)[0]*100
```



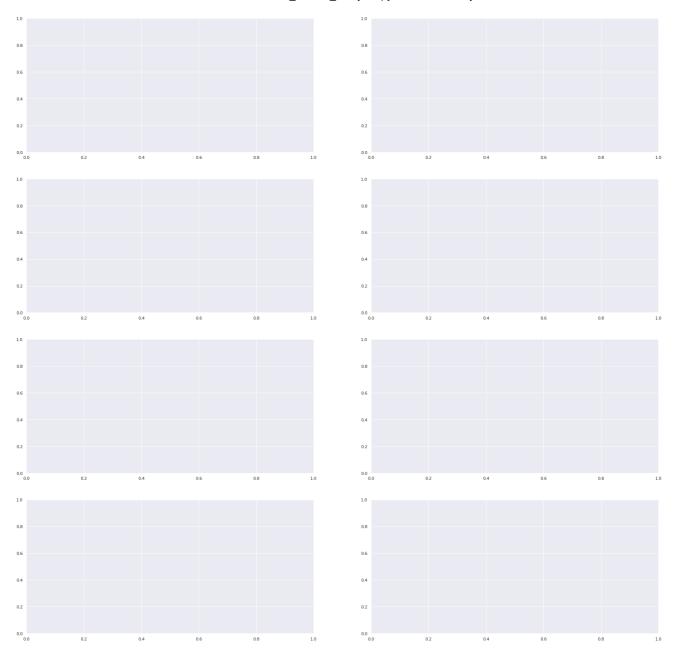
percentage of Alive Patients: 79.43925233644859%

```
Random under-sampling:
Alive 66
Dead 66
Name: Patient_Status, dtype: int64
Count (target)

60 -
50 -
40 -
10 -
```

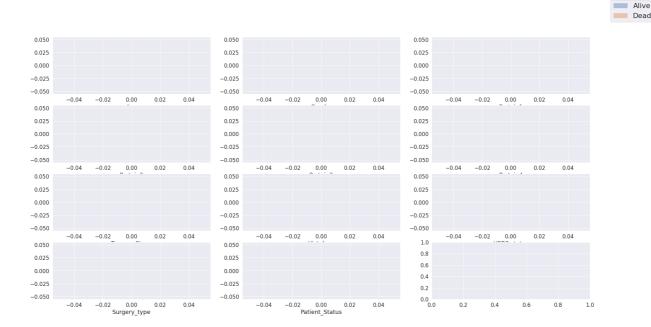
Categorical Features

```
for col in df.select_dtypes("object"):
  print(f'{col :-<50} {df[col].unique()}')</pre>
   Gender----- ['FEMALE' 'MALE']
   Tumour Stage-----['III' 'II' 'I']
   Histology------['Infiltrating Ductal Carcinoma'
    'Infiltrating Lobular Carcinoma']
   ER status----- ['Positive']
   PR status-----['Positive']
   HER2 status----- ['Negative' 'Positive']
   Surgery_type----- ['Modified Radical Mastectomy' 'Lu
                          ----- ['Alive' 'Dead' nan]
   Patient Status-----
fig, ax = plt.subplots(4,2, figsize=(30, 30))
i=0
sns.set(font_scale = 1.5)
for col in df.select dtypes('object'):
   sns.countplot(df_under[col], hue=df_under['Patient_Status'], ax=ax[i//2][i%2])
  i=i+1
plt.show()
import warnings
warnings.filterwarnings('ignore')
```



Continuous Features

```
Alive_df = df[df['Patient_Status']=="Alive"]
Dead_df = df[df['Patient_Status']=="Dead"]
sns.set(font_scale = 1.5)
fig, ax = plt.subplots(4,3, figsize=(30, 15))
i=0
for col in df.select_dtypes(include=['float64','int64']):
    sns.distplot(Alive_df[col],label='Alive',ax=ax[i//3][i%3])
    sns.distplot(Dead_df[col],label='Dead',ax=ax[i//3][i%3])
    i=i+1
fig.legend(labels=['Alive','Dead'],fontsize='22')
fig.show()
```



A bit od Data Engineering

```
def encoding(df):
                                                            #We can now analyze categorical
    code = {'FEMALE':0,
            'MALE':1,
            'III':3,
            'II':2,
            'I':1,
            'Infiltrating Ductal Carcinoma':0,
            'Mucinous Carcinoma':1,
            'Infiltrating Lobular Carcinoma':2,
            'Negative':0,
            'Positive':1,
            'Modified Radical Mastectomy':0,
            'Lumpectomy':1,
            'Simple Mastectomy':2,
            'Other':3,
            'Alive':1,
            'Dead':0
           }
    for col in df.select_dtypes('object'):
        df.loc[:,col]=df[col].map(code)
    return df
def imputation(df):
    df = df.fillna(df.median())
    df = df.dropna()
    return df
def feature_engineering(df):
    useless_columns = ['Patient_ID','Date_of_Surgery','Date_of_Last_Visit','ER status','PR
    df = df.drop(useless_columns,axis=1)
    return df
def preprocessing(df):
    df = encoding(df)
    df = feature engineering(df)
    df = imputation(df)
    X = df.drop('Patient Status',axis=1)
    y = df['Patient_Status']
    return df,X,y
df = data.copy()
df,X,y=preprocessing(df)
# Class count
count_class_0, count_class_1 = df['Patient_Status'].value_counts()
# Divide by class
df_class_0 = df[df['Patient_Status'] == 1]
df_class_1 = df[df['Patient_Status'] == 0]
```

```
df_class_0_under = df_class_0.sample(count_class_1,random_state=42)
df_under = pd.concat([df_class_0_under, df_class_1], axis=0)

print('Random under-sampling:')
print(df_under['Patient_Status'].value_counts())

# Resampling
```

df_under['Patient_Status'].value_counts().plot(kind='bar', title='Count (target)');

Random under-sampling:

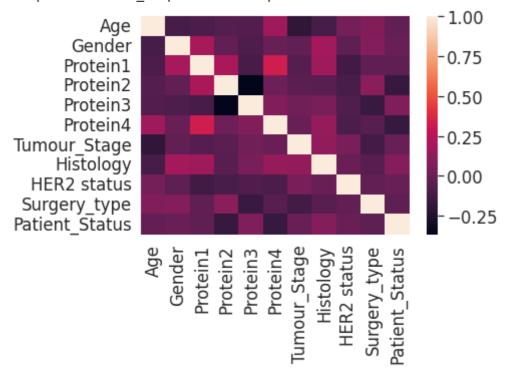
1.0 66 0.0 66

Name: Patient_Status, dtype: int64

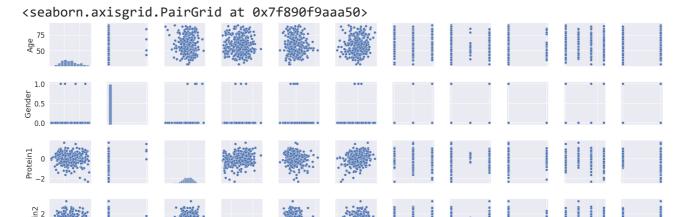


sns.heatmap(df_under.corr())





sns.pairplot(df, height=2)



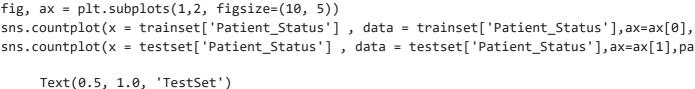
Modelling

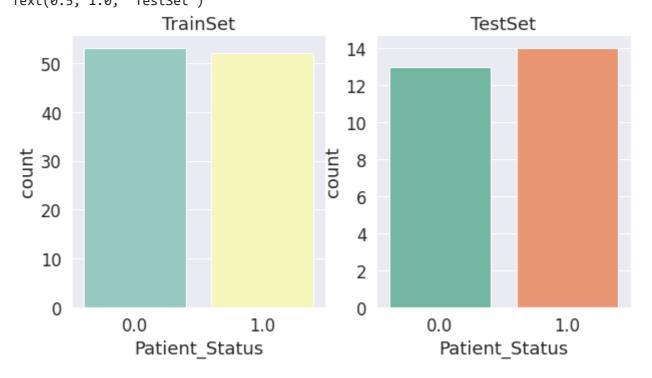


from sklearn.model_selection import train_test_split

from sklearn.metrics import f1_score, accuracy_score, confusion_matrix, classification_rep from sklearn.model_selection import learning_curve, cross_val_score, GridSearchCV from sklearn.model_selection import train_test_split

trainset, testset = train_test_split(df_under, test_size=0.2, random_state=0) fig, ax = plt.subplots(1,2, figsize=(10, 5)) sns.countplot(x = trainset['Patient_Status'] , data = trainset['Patient_Status'],ax=ax[0],





 $from \ sklearn. ensemble \ import \ Random Forest Classifier, \ AdaBoost Classifier$ from sklearn.svm import SVC from sklearn.pipeline import make_pipeline from sklearn.neighbors import KNeighborsClassifier

```
from sklearn.linear_model import LogisticRegression
from sklearn preprocessing import RobustScaler StandardScaler MinMayScaler

X_train = trainset.drop(['Patient_Status'],axis=1)
y_train = trainset['Patient_Status']

X_test = testset.drop(['Patient_Status'],axis=1)
y_test = testset['Patient_Status']

preprocessor = make_pipeline(RobustScaler())

PCAPipeline = make_pipeline(preprocessor, PCA(n_components=3,random_state=42))

RandomPipeline = make_pipeline(preprocessor,RandomForestClassifier(random_state=42))
AdaPipeline = make_pipeline(preprocessor,AdaBoostClassifier(random_state=42))

SVMPipeline = make_pipeline(preprocessor,SVC(random_state=42,probability=True))
KNNPipeline = make_pipeline(preprocessor,LogisticRegression(solver='sag',random_state=42))

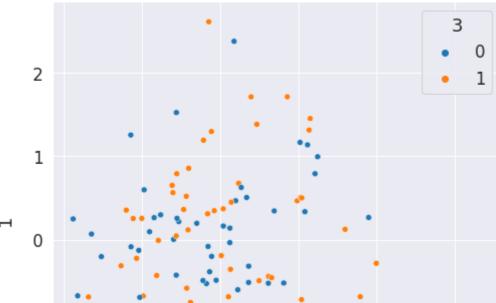
LRPipeline = make_pipeline(preprocessor,LogisticRegression(solver='sag',random_state=42))
```

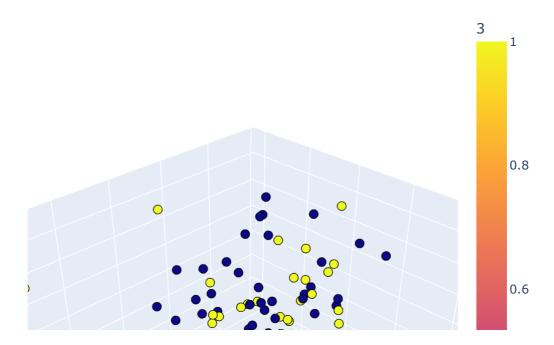
→ PCA Analysis

```
PCA_df = pd.DataFrame(PCAPipeline.fit_transform(X_train))
y_train = y_train.astype(int)
y_train.reset_index(drop=True, inplace=True)
PCA_df = pd.concat([PCA_df, y_train], axis=1, ignore_index=True )
PCA_df.head()
```

	0	1	2	3	7
0	-0.792496	-0.009582	-0.707232	1	
1	-0.561121	1.524745	-0.453318	0	
2	-0.072062	-1.668854	-0.343776	0	
3	0.236570	0.675897	-0.401078	1	
4	-0.275412	-0.856502	0.431383	0	

```
plt.figure(figsize=(8,8))
sns.scatterplot(PCA_df[0],PCA_df[1],hue=PCA_df[3],palette=sns.color_palette("tab10", 2))
plt.show()
```





Training models

```
dict_of_models = {'RandomForest': RandomPipeline,
'AdaBoost': AdaPipeline,
'SVM': SVMPipeline,
'KNN': KNNPipeline,
'LR': LRPipeline}
                                                                             0.2
def evaluation(model):
    # calculating the probabilities
    y_pred_proba = model.predict_proba(X_test)
    # finding the predicted valued
    y pred = np.argmax(y pred proba,axis=1)
    print('Accuracy = ', accuracy_score(y_test, y_pred))
    print('-')
    print(confusion_matrix(y_test,y_pred))
    print('-')
    print(classification_report(y_test,y_pred))
    print('-')
    N, train_score, test_score = learning_curve(model, X_train, y_train,
                                               cv=4, scoring='f1',
                                               train sizes=np.linspace(0.1,1,10))
    plt.figure(figsize=(5,5))
    plt.plot(N, train_score.mean(axis=1), label='train score')
    plt.plot(N, test_score.mean(axis=1), label='validation score')
    plt.legend()
    plt.show()
```

```
sns.set(font_scale = 1)
for name, model in dict_of_models.items():
    print('-----')
    print(name)
    model.fit(X_train,y_train)
    evaluation(model)
```

RandomForest

Accuracy = 0.48148148148148145

[[8 5] [9 5]]

_

	precision	recall	f1-score	support
0.0	0.47	0.62	0.53	13
1.0	0.50	0.36	0.42	14
accuracy			0.48	27
macro avg	0.49	0.49	0.47	27
weighted avg	0.49	0.48	0.47	27



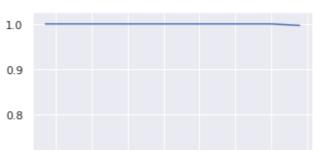
AdaBoost

Accuracy = 0.48148148148145

[[6 7] [7 7]]

_

-	precision	recall	f1-score	support
0.	0 0.46	0.46	0.46	13
1.	0.50	0.50	0.50	14
accurac	y		0.48	27
macro av	g 0.48	0.48	0.48	27
weighted av	g 0.48	0.48	0.48	27





SVM

Accuracy = 0.48148148148148145

[[10 3] [11 3]]

precision recall f1-score support 0.0 0.48 0.77 0.59 13 1.0 0.50 0.21 0.30 14 0.48 27 accuracy 0.49 0.44 27 macro avg 0.49 weighted avg 0.49 0.48 0.44 27

train score validation score 0.9 0.8 0.7 0.6 0.5 0.4 0.3 70 80 10 20 30 50 60

KNN

[[4 9] [9 5]]

-	precision	recall	f1-score	support
0.0	0.31	0.31	0.31	13
1.0	0.36	0.36	0.36	14
accuracy			0.33	27
macro avg	0.33	0.33	0.33	27
weighted avg	0.33	0.33	0.33	27

Tuning threshold

```
best_classifier = KNNPipeline
thresholds = [0.3, 0.4, 0.5, 0.6, 0.7, 0.8]
best_t = 0.3
best acc = 0
for t in thresholds:
   y_pred = (best_classifier.predict_proba(X_test)[:,1] >= t).astype(int)
   acc = accuracy_score(y_test, y_pred)
   if acc > best_acc:
       best_acc=acc
       best_t=t
    LR
print('Accuracy on test set :',round(best_acc*100),"%")
print('Best threshold :',best_t)
    Accuracy on test set: 44 %
    Best threshold: 0.7
```

Training Artificial Neural Network

```
macro avg
                        0.44
                                  0.44
                                                         27
# Importing the Keras libraries and packages
import tensorflow as tf
from keras.models import Sequential
from keras.layers import Dense, Dropout
X train.shape
     (105, 10)
# Initialising the ANN
classifier = Sequential()
# Adding the input layer and the first hidden layer
classifier.add(Dense(units = 16, kernel_initializer = 'uniform', activation = 'relu', inpu
classifier.add(Dropout(0.2))
# Adding the second hidden layer
classifier.add(Dense(units = 32, kernel_initializer = 'uniform', activation = 'relu'))
classifier.add(Dropout(0.2))
# Adding the third hidden layer
classifier.add(Dense(units = 8, kernel_initializer = 'uniform', activation = 'relu'))
classifier.add(Dropout(0.2))
# Adding the output layer
classifier.add(Dense(units = 1, kernel_initializer = 'uniform', activation = 'sigmoid'))
classifier.add(Dropout(0.2))
```

```
# Compiling the ANN
```

classifier.compile(optimizer = 'adam', loss = 'binary_crossentropy', metrics = ['accuracy'
callback = tf.keras.callbacks.EarlyStopping(monitor='accuracy', patience=80)
history =classifier.fit(X_train, y_train, batch_size = 10, epochs = 100, callbacks=callbac

```
Epoch 72/100
Epoch 73/100
Epoch 74/100
Epoch 75/100
Epoch 76/100
Epoch 77/100
11/11 [============= ] - 0s 2ms/step - loss: 2.0054 - accuracy: 0.
Epoch 78/100
Epoch 79/100
Epoch 80/100
Epoch 81/100
Epoch 82/100
11/11 [============= ] - 0s 2ms/step - loss: 2.7609 - accuracy: 0.4
Epoch 83/100
Epoch 84/100
Epoch 85/100
Epoch 86/100
Epoch 87/100
Epoch 88/100
Epoch 89/100
Epoch 90/100
Epoch 91/100
Epoch 92/100
Epoch 93/100
Epoch 94/100
Epoch 95/100
Epoch 96/100
Epoch 97/100
Epoch 98/100
```

Conclusion

According to the results shown above, these models (RF, AdaBoost, KNN, SVM, XGBoost, LR, ANN) can't make the classification between Dead and Alive patients. Best we can do is getting a 1/2 chance of guessing right...

Hypothesis

The features have no impact on the target There isn't enough rows in the dataset (need more people) The dataset isn't representative of the population As we undersampled the dataset, we only have 66*2 rows in the end. I could have tried to oversample instead

X