5	Many of us are familiar with heart desiase and stroke. According World Health Organization World Health Organization heart disease, stroke is the number one cause of death and stroke is the second and there is a need for research. The heart failure prediction data serprovides valuable information about heart failure parameters. In my research, I will use logistic regression to predict whether a patient a heart failure or not using Heart Failure Prediction Dataset. 2- INTRODUCTION 2.1 Information About The Data Set
2	 2.1 Information About The Data Set Original Data Set and Source Info: Retrieved from fedesoriano. Heart Failure Prediction Dataset 12 Attributes 918 observations 2.2 Atribute(feature) Information Age: age of the patient [years] Sex: sex of the patient [M: Male, F: Female]
	 Sex: sex of the patient [M: Male, F: Female] ChestPainType: chest pain type [TA: Typical Angina, ATA: Atypical Angina, NAP: Non-Anginal Pain, ASY: Asymptomatic] RestingBP: resting blood pressure [mm Hg] Cholesterol: serum cholesterol [mm/dl] FastingBS: fasting blood sugar [1: if FastingBS > 120 mg/dl, 0: otherwise] RestingECG: resting electrocardiogram results [Normal: Normal, ST: having ST-T wave abnormality (T wave inversions and/or ST elevation or depression of > 0.05 mV), LVH: showing probable or definite left ventricular hypertrophy by Estes' criteria] MaxHR: maximum heart rate achieved [Numeric value between 60 and 202] ExerciseAngina: exercise-induced angina [Y: Yes, N: No] Oldpeak: oldpeak = ST [Numeric value measured in depression]
	• Oldpeak: oldpeak = SI [Numeric value measured in depression] • ST_Slope: the slope of the peak exercise ST segment [Up: upsloping, Flat: flat, Down: downsloping] • HeartDisease: output class [1: heart disease, 0: Normal] 3- Import Import Python Libraries Below are the list of python libraries that we need for this project # for data processing import pandas as pd import numpy as np # for vizulazations import matplotlib.pyplot as plt import seaborn as sns # scalling the data for the machine learning model from sklearn.preprocessing import StandardScaler # machine learning libraries for data modeling from sklearn.model_selection import train_test_split from sklearn.linear model import LogisticRegression
	<pre>from sklearn.pipeline import make_pipeline from sklearn import preprocessing from sklearn.metrics import confusion_matrix,ConfusionMatrixDisplay,accuracy_score,plot_roc_curve from sklearn.feature_selection import SelectKBest, mutual_info_classif from sklearn.metrics import log_loss, roc_auc_score, recall_score, precision_score, average_precision_sc from sklearn.preprocessing import LabelEncoder from sklearn import metrics</pre> 4. Exploring and Cleaning the Data 4.1 Loading the raw data
]: [data_heart = pd.read_csv("heart.csv") print("onservation size: ", len(data_heart)) data_heart.head() onservation size: 918 Age Sex ChestPainType RestingBP Cholesterol FastingBS RestingECG MaxHR ExerciseAngina Oldpeak ST_Slope HeartDisease 0 40 M ATA 140 289 0 Normal 172 N 0.0 Up 1 49 F NAP 160 180 0 Normal 156 N 1.0 Flat
	2 37 M ATA 130 283 0 ST 98 N 0.0 Up 3 48 F ASY 138 214 0 Normal 108 Y 1.5 Flat 4 54 M NAP 150 195 0 Normal 122 N 0.0 Up 4.2 Remove duplicate or irrelevant observations In this step I will be checking to see if there are any dublicates or irrelavant observations. In our data set we have no dublicates. # this checks to see if df contains any dublicates
]: [<pre>data_heart.duplicated().sum() 4.3 Checking for missing data Checking for null and NA values: We have no null values. We got zero for each features. print(data_heart.info()) <class 'pandas.core.frame.dataframe'=""> RangeIndex: 918 entries, 0 to 917</class></pre>
	RangeIndex: 918 entries, 0 to 917 Data columns (total 12 columns): # Column Non-Null Count Dtype
4	9 Oldpeak 918 non-null float64 10 ST_Slope 918 non-null object 11 HeartDisease 918 non-null int64 dtypes: float64(1), int64(6), object(5) memory usage: 86.2+ KB None 4.4 Investigating all the elements whithin each Feature In this section, we will check the data to see if there is any variable that loo different than what is listed on the data source website. I did not see any abnormality. feature_no =0
	<pre>for column in data_heart: unique_values = np.unique(data_heart[column]) nr_values = len(unique_values) # number of unique_values feature_no+=1 # countin feature to make sure we screen all of the features print("Feature: ",feature_no, column) if nr_values <= 10: # if the number of values are less than 10 print("The number of unquie values for feature {} is: {} {} ".format(column, nr_values, unique else:</pre>

	********************************** Feature: 10 Oldpeak The number of unquie values for feature Oldpeak is: 53 ***********************************
	<pre>4.5 Summarizing and Vizulation Data for this step we need split our data two as categorical and numerical categ_data =[] numeric_data = [] binary_data = [] for column in data_heart: unique_values = np.unique(data_heart[column]) nr_values = len (unique_values) if data_heart.dtypes[column] == "object":</pre>
	<pre>categ_data.append(column) else: if nr_values == 2: binary_data.append(column) else: numeric_data.append(column) print("categorical data") print(categ_data) print("numeric_data) print(numeric_data) print("binary_data (numeric 0 and 1)")</pre>
4	categorical data ['Sex', 'ChestPainType', 'RestingECG', 'ExerciseAngina', 'ST_Slope'] numeric data ['Age', 'RestingBP', 'Cholesterol', 'MaxHR', 'Oldpeak'] binary data (numeric 0 and 1) ['FastingBS', 'HeartDisease'] 4.5a numerical data summary Now we can vizulazi our numeric data and check to see if there is any skweness and any linear relationship in between our features. Between the state of the second state of the sec
]:	plot indicates that we do not have any high correlation in our data. However we see some skewnees in oldpeak and chelestrol p = sns.pairplot(data_heart.loc[:,numeric_data], corner=True) plt.gcf().set_size_inches(15, 8)
	200
	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0
]: [data_comparison = data_heart.loc[:,numeric_data] data_comparison["HeartDisease"]= data_heart["HeartDisease"] g = sns.PairGrid(data_comparison, hue="HeartDisease", corner=True, palette = "mako") g.map_diag(sns.histplot, multiple="stack", element="step", color=".3", palette = "mako") g.map_offdiag(sns.scatterplot) g.add_legend()
	He Wax Have 100 -
,	we do not see high correlation in between variables
]:	<pre># correlation matrix data_comparison = data_heart.loc[: ,numeric_data] data_comparison["HeartDisease"]= data_heart["HeartDisease"] matrix = data_heart.loc[: ,numeric_data].corr().round(3) mask = np.triu(np.ones_like(matrix, dtype=bool)) sns.heatmap(data_heart.loc[: ,numeric_data].corr(),annot=True, vmax=1, vmin=-1, center=0, cmap='PiYG', plt.show()</pre> Age - 100 -0.75
	RestingBP - 0.25 Cholesterol0.095
	4.5b Outliers Detection I noticed some outliers, these outliers needs further investigation to see if we need to remove them. sns.boxplot(x=data_heart.Cholesterol) <axessubplot:xlabel='cholesterol'></axessubplot:xlabel='cholesterol'>
]: [0 100 200 300 400 500 600 data_heart["Cholesterol"].hist() <axessubplot:></axessubplot:>
	250 200 150 50
]:	<pre>from scipy import stats z = np.abs(stats.zscore(data_heart["Cholesterol"])) threshold = 3 print("##########") print(np.where(z > 3))</pre>
	<pre>######### (array([76, 149, 616], dtype=int64),) sns.boxplot(x=data_heart.Oldpeak) <axessubplot:xlabel='oldpeak'></axessubplot:xlabel='oldpeak'></pre>
]:	Oldpeak from scipy import stats z_2 = np.abs(stats.zscore(data_heart["Oldpeak"])) threshold = 3 print("##########") print(np.where(z_2 > 3)) ##################################
]: [<pre>(array([166, 324, 702, 771, 791, 850, 900], dtype=int64),) q_low = data_heart["Oldpeak"].quantile(0.01) q_hi = data_heart["Oldpeak"].quantile(0.99) df_filtered = data_heart[(data_heart["Oldpeak"] < q_hi) & (data_heart["Oldpeak"] > q_low)] q_hi 4.0</pre>
	 4.5c Categorical Data Summary reviwing the categoerical data we see that our data has more patiant that does not have heart failure so when we are creating our model we need to split accord so that we do not have bias. We can identify some of the classifications by just reviewing the plots below, for example, there is not a clear difference in resting in between heart disease and not hear disease butin the ST_slope: we see a big number of patient who has heart desiase have a Stope flat because of the difference you can easily tell.
]:	<pre>ax = sns.countplot(x="HeartDisease", data=data_heart,palette = "mako") for container in ax.containers:</pre>
	400 - 300 - 200 - 100 -
]: [HeartDisease ax = sns.countplot(x="HeartDisease", data=data_heart,palette = "mako") for container in ax.containers: ax.bar_label(container) plt.show() 500
	400 - 410 300 - 200 - 100 - 0
]: [# we need to combine categorical data and binary data to review category_data_joined = categ_data + binary_data print(category_data_joined) for f in categ_data: ax = sns.countplot(x = f, data = data_heart.loc[:,category_data_joined], hue = 'HeartDisease', pal for container in ax.containers: ax.bar_label(container)
	plt.show() ['Sex', 'ChestPainType', 'RestingECG', 'ExerciseAngina', 'ST_Slope', 'FastingBS', 'HeartDisease'] 400 - 458
	200 - 143 143 143 140
	HeartDisease
	26 20 ATA NAP ASY TA ChestPainType HeartDisease 0 1 1
	150 - 100 - 100 - 106 - 82 - 106 - 107 - 106 - 107 - 1
	350 - HeartDisease
	ExerciseAngina HeartDisease 0 10 11 11 11 11 11 11 11 11 11 11 11
	200 - 150 - 100 - 78 79 49 14 Down ST_Slope
l r	5. Transforming the Categorical Variables: Creatinging Dummy Variables In this section we will transfer our categorical data to dummy variables. We already have some binary data we just need to transform rest of it. I did not want to use to pandas dummy variable builtin function because of the "dummy variable trap". It made my code ver messy therefore I decided to transfrom them one by using my own function. print("we need to transform these below three columns", categ_data)
	<pre>def transformCatData(colname, data): le = LabelEncoder() def transformCate(colname, df): new_col_name = colname+"_nu" df_new = pd.DataFrame(le.fit_transform(df[colname].values), columns =[new_col_name]) merged= pd.concat([df, df_new], axis="columns") del merged[colname] return merged for i in range(len(categ_data)): if i == 0:</pre>
	<pre>if i == 0:</pre>
	<pre>'Cholesterol', 'FastingBS', 'MaxHR', 'Oldpeak', 'Sex_nu', 'ChestPainType_nu', 'RestingECG_nu', 'ExerciseAngina_nu', 'ST_Slope_nu', 'HeartDisease',]]</pre> <pre>print(len(data_heart.columns.tolist()))</pre>
	<pre>print("new order, ", data_heart.columns.tolist()) we need to transform these below three columns ['Sex', 'ChestPainType', 'RestingECG', 'ExerciseAngina', 'pe'] original order Index(['Age', 'RestingBP', 'Cholesterol', 'FastingBS', 'MaxHR', 'Oldpeak',</pre>
-	<pre>formula</pre>
	<pre>print("**** y***") #print(y.head()) x_train,x_test,y_train, y_test = train_test_split(x, y,</pre>
	**** y*** Age RestingBP Cholesterol FastingBS MaxHR Oldpeak Sex_nu \ 506 75 136 225 0 112 3.0 1 576 62 139 170 0 120 3.0 1 339 64 95 0 1 145 1.1 0 383 38 110 0 0 156 0.0 0 580 51 131 152 1 130 1.0 1 ChestPainType_nu RestingECG_nu ExerciseAngina_nu ST_Slope_nu 506 0 1 1 1 1 576 0 2 1 1 1 339 0 1 0 0 0
	339 0 1 0 1 0 1 580 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
j: S t	
-	8- Feature Selection Before evaluating our model I want evaluate the some of the features that are not very important for my model. first we check coef valuating of each features and plot def dropFeatureAccr(x_train, x_test, y_train, y_test): accuracy_featuresDrop = []
	<pre>accuracy_featuresDrop = [] column_name= list(x_train.columns.values.tolist()) for column in range(len(column_name)): x_train_modified = x_train.drop(columns=[column_name[column]]) x_test_modified = x_test.drop(columns=[column_name[column]]) log_reg_classifier = LogisticRegression(solver='lbfgs', max_iter=1000) log_reg_classifier.fit(x_train_modified,y_train) y_pred_mod = log_reg_classifier.predict(x_test_modified) accuracy_featuresDrop.append(accuracy_score(y_test,y_pred_mod)) dictionary_feature_drop = dict(zip(column_name, accuracy_featuresDrop)) return dictionary_feature_drop logisticRegDrop =dropFeatureAccr(x_train= x_train,y_train= y_train, x_test= x_test, y_test = y_test)</pre>
	<pre>logisticRegDrop =dropFeatureAccr(x_train= x_train,y_train= y_train, x_test= x_test, y_test = y_test) print(logisticRegDrop) print("max effect ") print("feature", max(logisticRegDrop, key=logisticRegDrop.get)) print("accuracy score ", logisticRegDrop.get(max(logisticRegDrop, key=logisticRegDrop.get))) print("when we drop it the accuracy score went down effect") print("feature", min(logisticRegDrop, key=logisticRegDrop.get)) print("accuracy score ", logisticRegDrop.get(min(logisticRegDrop, key=logisticRegDrop.get))) lists = logisticRegDrop.items() # sorted by key, return a list of tuples</pre>
	lists = logisticRegDrop.items() # sorted by key, return a list of tuples x, y = zip(*lists) # unpack a list of pairs into two tuples plt.plot(x, y) plt.xticks(x,[0,1,2,3,4,5,6,7,8,9,10]) plt.show() {'Age': 0.8583877995642701, 'RestingBP': 0.8605664488017429, 'Cholesterol': 0.8518518518518519, 'FastingBesterol': 0.8562091503267973, 'MaxHR': 0.8758169934640523, 'Oldpeak': 0.8518518518519, 'Sex_nu': 0.8540305010893240523, 'St_Slope_nu': 0.8649237472766884, 'RestingECG_nu': 0.8583877995642701, 'ExerciseAngina_nu': 0.8605664429, 'ST_Slope_nu': 0.8366013071895425} max effect
	29, 'ST_Slope_nu': 0.8366013071895425} max effect feature MaxHR accuracy score 0.8758169934640523 when we drop it the accuracy score went down effect feature ST_Slope_nu accuracy score 0.8366013071895425 0.875 0.870 0.865
	0.865 0.860 0.855 0.845 0.840 0.835 0 1 2 3 4 5 6 7 8 9 10
f (0.8823529411764706 10- Evaluate the Model
f	<pre>fpr, tpr, _ = metrics.roc_curve(y_test, y_pred_mod) auc = metrics.roc_auc_score(y_test, y_pred_mod) #create ROC curve plt.plot(fpr,tpr,label="AUC="+str(auc))</pre>
: : :	<pre>auc = metrics.roc_auc_score(y_test, y_pred_mod) #create ROC curve plt.plot(fpr,tpr,label="AUC="+str(auc)) plt.ylabel('True Positive Rate') plt.xlabel('False Positive Rate') plt.legend(loc=4) plt.show()</pre>
	<pre>auc = metrics.roc_auc_score(y_test, y_pred_mod) #create ROC curve plt.plot(fpr,tpr,label="AUC="+str(auc)) plt.ylabel('True Positive Rate') plt.xlabel('False Positive Rate') plt.legend(loc=4) plt.show()</pre>

TN_log, FP_log, FN_log, TP_log = cm.ravel() print(TN_log, FP_log, FN_log, TP_log) group_names = ["TN", "FP", "TN", "TP"] group_counts = ["(0:0.0f)".format(value) for value in	<pre>accuracy</pre>				
The control of the co					
TIR Log = TID Lod / LTP Log + FN Log	<pre>cm.flatten()/np.sum(cm)] labels = [f"{v1}\n{v2}\n{v3}" for v1, v2, v3 in</pre>	<pre>group_percentages = ["{0:.2%}".format(value) for value in</pre>			
my_sccuracy_score_log= accuracy_score(y_test, y_pred_mcd) my_marrix_values_l = {""":"ST_log, ""p":"FT_log, """:"ST_log, ""2"":TT_log, ""accuracy":my_accuracy_score_log,	-150 -125 -100 -75 -50 -50 -25 TPR log = TP log/(TP log + FN log)				
that we can also add our prediction for each data points. Please see example below. [52]: test_prob = log_reg_classifier.predict_proba(x_test_modified) [;, 1] # probability test_prob = log_reg_classifier.predict(x_test_modified) # target = values print(test_prob[0]) olumination of the below results gives us the odd ratio for our each feature. For example sex, for male (sex=1) to the odds of having heart disease for females is exp(1.215323). You can derive it based on the logistic regression equation. [88]: coefficients = np.hstack((log_reg_classifier.intercept_, log_reg_classifier.coef_[0])) print(len(coefficients)) print(x_train.columns.values.tolist()) pd.DataFrame(dats=('variable': ['intercept'] + x_train_modified.columns.values.tolist(), 'coefficient': coe [88]: variable coefficient nu', 'ExerciseAngina_nu', 'ST_Slope_nu'] Age 0.021428 RestingBP 0.002181 Age 0.021428 RestingBP 0.002181 Age 0.0233080 Sex_nu 1.215323 FexciseAngina_nu 1.146983	<pre>my_accuracy_score_log= accuracy_score(y_test, y_pred_mod) my_matrix_values_1 = {"TN":TN_log, "FP":FP_log, "FN":FN_log</pre>	e. Over all our model did a pretty good job but we can still			
<pre>print(len(coefficients)) print(x_train.columns.values.tolist()) pd.DataFrame(data={'variable': ['intercept'] + x_train_modified.columns.values.tolist(), 'coefficient': coe 9 ['Age', 'RestingBP', 'Cholesterol', 'FastingBS', 'MaxHR', 'Oldpeak', 'Sex_nu', 'ChestPainType_nu', 'RestingEnu', 'ExerciseAngina_nu', 'ST_Slope_nu'] 1</pre>	that we can also add our prediction for each data points. Please see example [52]: test_prob = log_reg_classifier.predict_proba(x_test_modified) test_pred = log_reg_classifier.predict(x_test_modified) #tast_print(test_prob[0]) print(test_pred[0]) 0.8321832370727031 1 the below results gives us the odd ratio for our each feature. For example see	e below. ed) [:, 1] # probabilty rget = values ex, for male (sex=1) to the odds of having heart disease for			
 RestingBP 0.002181 Cholesterol -0.002316 FastingBS 0.795913 Oldpeak 0.233080 Sex_nu 1.215323 ExerciseAngina_nu 1.146983 	<pre>print(len(coefficients)) print(x_train.columns.values.tolist()) pd.DataFrame(data={'variable': ['intercept'] + x_train_mode 9 ['Age', 'RestingBP', 'Cholesterol', 'FastingBS', 'MaxHR', 'onu', 'ExerciseAngina_nu', 'ST_Slope_nu'] t[88]:</pre>	ified.columns.values.tolist(), 'coefficient': coe			
	 RestingBP 0.002181 Cholesterol -0.002316 FastingBS 0.795913 Oldpeak 0.233080 Sex_nu 1.215323 ExerciseAngina_nu 1.146983 				