

DATA SCIENCE CAPSTONE PROJECT

Predictive Medical Assistant

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Introduction - Problem Statement : The project introduces the development of the Predictive Medical Assistant, a predictive model focused on anticipating the probability of coronary heart disease (CHD) or myocardial infarction (MI) by leveraging various health-related attributes. By employing machine learning techniques, the project aims to analyze health-related data and create a predictive model to enhance early detection and timely, targeted intervention for individuals susceptible to heart diseases, thereby contributing significantly to preventive care within the healthcare domain. To kickstart the project, a comprehensive health-related dataset will be acquired and preprocessed from Kaggle (<https://www.kaggle.com/datasets/kamilpytlak/personal-key-indicators-of-heart-disease>).

Data description

The project utilizes the following data attributes:

- **Heart disease:** Indicates whether respondents have ever reported having coronary heart disease (CHD) or myocardial infarction (MI)
- **BMI:** Body Mass Index (BMI)
- **Smoking:** Determines if respondents have smoked at least 100 cigarettes in their lifetime. (Note: 5 packs = 100 cigarettes)
- **Alcohol drinking:** Indicates whether respondents are considered heavy drinkers. (Note: For adult men, heavy drinking refers to consuming more than 14 drinks per week, while for adult women, it means consuming more than 7 drinks per week)
- **Stroke:** Indicates whether respondents have ever suffered a stroke
- **Physical health:** Reflects the number of days during the past 30 days when respondents experienced poor physical health, including illness and injury
- **Mental health:** Indicates the number of days during the past 30 days when respondents experienced poor mental health
- **Walking difficulty:** Determines if respondents have serious difficulty walking or climbing stairs
- **Sex:** Identifies respondents' gender as male or female
- **Age category:** Represents respondents' age categorized into fourteen levels
- **Ethnicity:** Indicates the imputed race/ethnicity value
- **Diabetes:** Determines if respondents have ever had diabetes
- **Physical activity:** Indicates whether respondents have engaged in physical activity or exercise during the past 30 days, excluding regular job-related activities
- **General health:** Reflects respondents' perception of their overall general health
- **Sleep time:** Represents the average number of hours of sleep obtained within a 24-hour period
- **Asthma:** Determines if respondents have ever been told they had asthma.
- **Kidney disease:** Indicates if respondents have ever been diagnosed with kidney disease, excluding kidney stones, bladder infections, or incontinence
- **Skin cancer:** Determines if respondents have ever been told they had skin cancer

1. Initial setup, dataset acquisition and data preprocessing

This section encompasses initial setup code, data loading, and crucial preprocessing steps essential for effective data analysis and modeling. These steps include handling missing values, binary encoding, scaling numerical features, and one-hot encoding categorical data to prepare the dataset adequately.

In [3]:

```
# Import necessary libraries for data manipulation, preprocessing, model training, evaluation, and visualization
import pandas as pd
import numpy as np
from sklearn.preprocessing import MinMaxScaler
from sklearn.neighbors import KNeighborsClassifier
from sklearn.linear_model import LogisticRegression
from sklearn.ensemble import RandomForestClassifier
from sklearn.model_selection import train_test_split, GridSearchCV
from sklearn.decomposition import PCA
from sklearn.cluster import KMeans
from sklearn.metrics import accuracy_score
from joblib import dump, load
from tableone import TableOne
import seaborn as sns
import matplotlib.pyplot as plt
```

In [5]:

```
# Function for data preprocessing
def data_preprocessing(input_data=None):
    # Load data
    data = pd.read_csv("heart_2020_cleaned.csv")

    # Handle missing values
    data.dropna(inplace=True)

    # Binary encoding
    binary_columns = ['Smoking', 'AlcoholDrinking', 'Stroke', 'DiffWalking', 'Sex', 'PhysicalActivity', 'Asthma', 'KidneyDisease', 'SkinCancer']
    data[binary_columns] = data[binary_columns].replace({'Yes': 1, 'No': 0, 'Male': 1, 'Female': 0})

    # Scale numerical features
    scaler = MinMaxScaler()
    numerical_cols = ['BMI', 'PhysicalHealth', 'MentalHealth', 'SleepTime']
    data[numerical_cols] = scaler.fit_transform(data[numerical_cols])

    # One-hot encode categorical features with pd.get_dummies
    categorical_columns = ['AgeCategory', 'Race', 'Diabetic', 'GenHealth']
    data = pd.get_dummies(data, columns=categorical_columns)

    return data
```

In [6]:

```
# Load data and preprocess
```

```
data = data_preprocessing()
```

2. Exploratory data analysis (EDA)

EDA is an essential preliminary step in understanding and preparing data which helps uncover the insights into the dataset's characteristics, the underlying patterns and relationships in the dataset, ultimately leading to more informed and effective model building. Visualizations and statistical tests provide insights into the dataset's characteristics, including the correlation matrix, distribution of the target variable, and univariate statistical tests.

In [7]:

```
# Display the dimensions of the dataset
print("Number of rows and columns:")
print(data.shape)
```

Number of rows and columns:
(313014, 41)

In [8]:

```
# Display the column labels (names) of the dataset
print("Column names:")
print(data.columns)
```

Column names:
Index(['HeartDisease', 'BMI', 'Smoking', 'AlcoholDrinking', 'Stroke',
 'PhysicalHealth', 'MentalHealth', 'DiffWalking', 'Sex',
 'PhysicalActivity', 'SleepTime', 'Asthma', 'KidneyDisease',
 'SkinCancer', 'AgeCategory_18-24', 'AgeCategory_25-29',
 'AgeCategory_30-34', 'AgeCategory_35-39', 'AgeCategory_40-44',
 'AgeCategory_45-49', 'AgeCategory_50-54', 'AgeCategory_55-59',
 'AgeCategory_60-64', 'AgeCategory_65-69', 'AgeCategory_70-74',
 'AgeCategory_75-79', 'AgeCategory_80 or older',
 'Race_American Indian/Alaskan Native', 'Race_Asian', 'Race_Black',
 'Race_Hispanic', 'Race_Other', 'Race_White', 'Diabetic_No',
 'Diabetic_Yes', 'Diabetic_Yes (during pregnancy)',
 'GenHealth_Excellent', 'GenHealth_Fair', 'GenHealth_Good',
 'GenHealth_Poor', 'GenHealth_Very good'],
 dtype='object')

In [9]:

```
# Display overview of the dataset
print("Data overview:")
print(data.info())
```

Data overview:
<class 'pandas.core.frame.DataFrame'>
Index: 313014 entries, 0 to 319794
Data columns (total 41 columns):
Column Non-Null Count Dtype
--- --- -
0 HeartDisease 313014 non-null object
1 BMI 313014 non-null float64
2 Smoking 313014 non-null int64
3 AlcoholDrinking 313014 non-null int64
4 Stroke 313014 non-null int64
5 PhysicalHealth 313014 non-null float64
6 MentalHealth 313014 non-null float64
7 DiffWalking 313014 non-null int64
8 Sex 313014 non-null int64
9 PhysicalActivity 313014 non-null int64
10 SleepTime 313014 non-null float64
11 Asthma 313014 non-null int64
12 KidneyDisease 313014 non-null int64
13 SkinCancer 313014 non-null int64
14 AgeCategory_18-24 313014 non-null bool
15 AgeCategory_25-29 313014 non-null bool
16 AgeCategory_30-34 313014 non-null bool
17 AgeCategory_35-39 313014 non-null bool
18 AgeCategory_40-44 313014 non-null bool
19 AgeCategory_45-49 313014 non-null bool
20 AgeCategory_50-54 313014 non-null bool
21 AgeCategory_55-59 313014 non-null bool
22 AgeCategory_60-64 313014 non-null bool
23 AgeCategory_65-69 313014 non-null bool
24 AgeCategory_70-74 313014 non-null bool
25 AgeCategory_75-79 313014 non-null bool
26 AgeCategory_80 or older 313014 non-null bool
27 Race_American Indian/Alaskan Native 313014 non-null bool
28 Race_Asian 313014 non-null bool
29 Race_Black 313014 non-null bool
30 Race_Hispanic 313014 non-null bool
31 Race_Other 313014 non-null bool
32 Race_White 313014 non-null bool
33 Diabetic_No 313014 non-null bool
34 Diabetic_Yes 313014 non-null bool
35 Diabetic_Yes (during pregnancy) 313014 non-null bool
36 GenHealth_Excellent 313014 non-null bool
37 GenHealth_Fair 313014 non-null bool
38 GenHealth_Good 313014 non-null bool
39 GenHealth_Poor 313014 non-null bool
40 GenHealth_Very good 313014 non-null bool
dtypes: bool(27), float64(4), int64(9), object(1)
memory usage: 43.9+ MB
None

In [10]:

```
# Display descriptive statistics for the numerical columns of the dataset
print("Data description:")
```

```
print(data.describe())
```

Data description:

	BMI	Smoking	AlcoholDrinking	Stroke
count	313014.000000	313014.000000	313014.000000	313014.000000
mean	0.196298	0.411959	0.068399	0.037426
std	0.076472	0.492189	0.252431	0.189805
min	0.000000	0.000000	0.000000	0.000000
25%	0.144513	0.000000	0.000000	0.000000
50%	0.184233	0.000000	0.000000	0.000000
75%	0.233007	1.000000	0.000000	0.000000
max	1.000000	1.000000	1.000000	1.000000

	PhysicalHealth	MentalHealth	DiffWalking	Sex
count	313014.000000	313014.000000	313014.000000	313014.000000
mean	0.111554	0.129514	0.137249	0.475589
std	0.264016	0.264482	0.344111	0.499405
min	0.000000	0.000000	0.000000	0.000000
25%	0.000000	0.000000	0.000000	0.000000
50%	0.000000	0.000000	0.000000	0.000000
75%	0.066667	0.100000	0.000000	1.000000
max	1.000000	1.000000	1.000000	1.000000

	PhysicalActivity	SleepTime	Asthma	KidneyDisease
count	313014.000000	313014.000000	313014.000000	313014.000000
mean	0.776422	0.265154	0.133221	0.036695
std	0.416643	0.062348	0.339814	0.188012
min	0.000000	0.000000	0.000000	0.000000
25%	1.000000	0.217391	0.000000	0.000000
50%	1.000000	0.260870	0.000000	0.000000
75%	1.000000	0.304348	0.000000	0.000000
max	1.000000	1.000000	1.000000	1.000000

	SkinCancer
count	313014.000000
mean	0.092986
std	0.290414
min	0.000000
25%	0.000000
50%	0.000000
75%	0.000000
max	1.000000

In [11]:

```
# Calculate the number of missing values for each column
data.isnull().sum()
```

Out[11]:

HeartDisease	0
BMI	0
Smoking	0
AlcoholDrinking	0
Stroke	0
PhysicalHealth	0
MentalHealth	0
DiffWalking	0
Sex	0
PhysicalActivity	0
SleepTime	0
Asthma	0
KidneyDisease	0
SkinCancer	0
AgeCategory_18-24	0
AgeCategory_25-29	0
AgeCategory_30-34	0
AgeCategory_35-39	0
AgeCategory_40-44	0
AgeCategory_45-49	0
AgeCategory_50-54	0
AgeCategory_55-59	0
AgeCategory_60-64	0
AgeCategory_65-69	0
AgeCategory_70-74	0
AgeCategory_75-79	0
AgeCategory_80 or older	0
Race_American Indian/Alaskan Native	0
Race_Asian	0
Race_Black	0
Race_Hispanic	0
Race_Other	0
Race_White	0
Diabetic_No	0
Diabetic_Yes	0
Diabetic_Yes (during pregnancy)	0
GenHealth_Excellent	0
GenHealth_Fair	0
GenHealth_Good	0
GenHealth_Poor	0
GenHealth_Very good	0

dtype: int64

In [12]:

```
# Display the first few rows of the dataset
print("\nSample data:")
print(data.head())
```

Sample data:

	HeartDisease	BMI	Smoking	AlcoholDrinking	Stroke	PhysicalHealth
0	No	0.055294	1	0	0	0.100000
1	No	0.100447	0	0	1	0.000000

```

2      No  0.175782      1      0      0      0.666667
3      No  0.147169      0      0      0      0.000000
4      No  0.141132      0      0      0      0.933333

MentalHealth  DiffWalking  Sex  PhysicalActivity  ...  Race_Other  \
0      1.0      0      0      1      ...      False
1      0.0      0      0      1      ...      False
2      1.0      0      1      1      ...      False
3      0.0      0      0      0      ...      False
4      0.0      1      0      1      ...      False

Race_White  Diabetic_No  Diabetic_Yes  Diabetic_Yes (during pregnancy)  \
0      True      False      True      False
1      True      True      False      False
2      True      False      True      False
3      True      True      False      False
4      True      True      False      False

GenHealth_Excellent  GenHealth_Fair  GenHealth_Good  GenHealth_Poor  \
0      False      False      False      False
1      False      False      False      False
2      False      True      False      False
3      False      False      True      False
4      False      False      False      False

GenHealth_Very good
0      True
1      True
2      False
3      False
4      True

[5 rows x 41 columns]

```

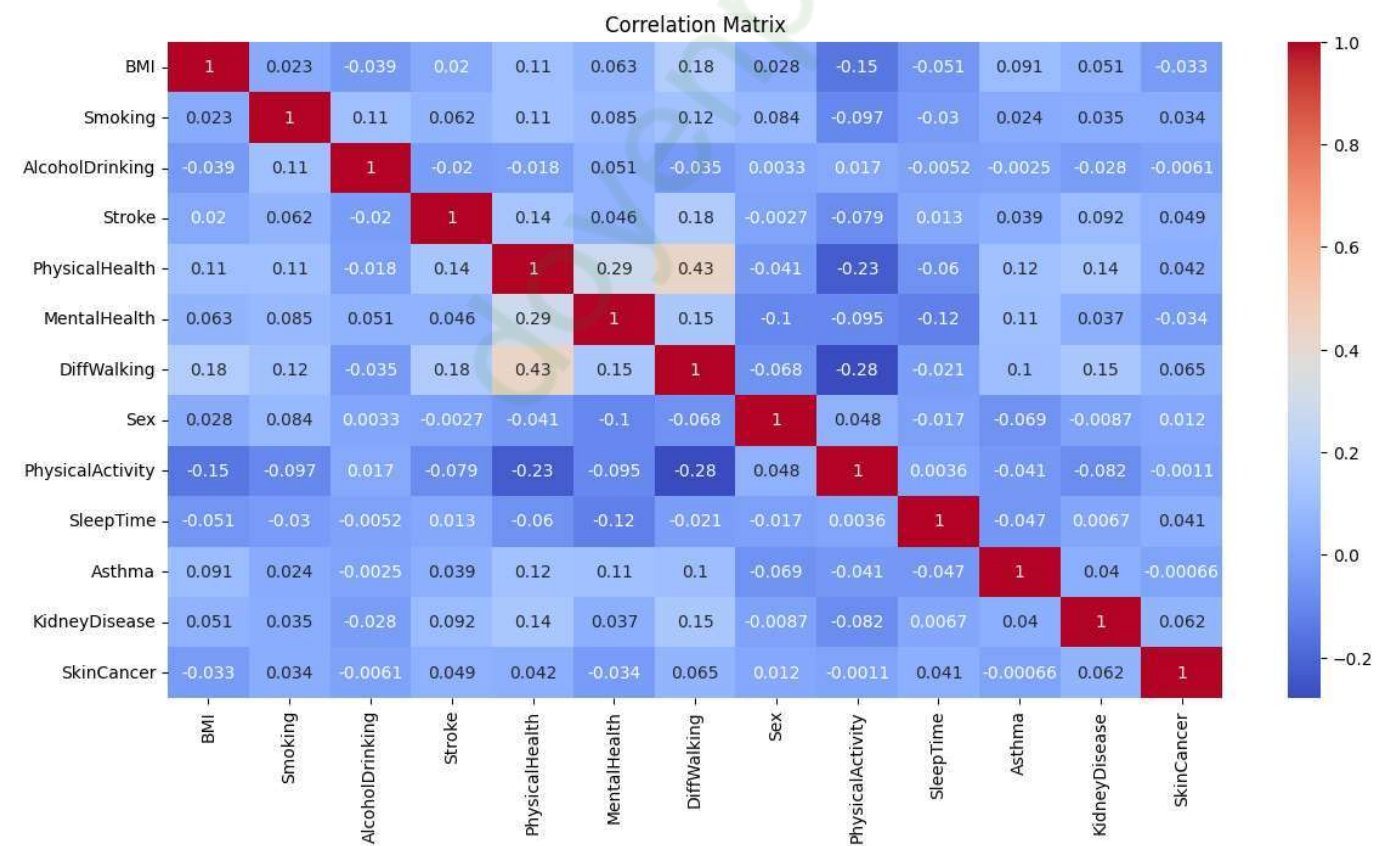
In [13]:

```

# Exclude non-numeric columns from correlation calculation
numeric_data = data.select_dtypes(include=['number'])

# Visualize correlation matrix
plt.figure(figsize=(14, 7))
sns.heatmap(numeric_data.corr(), annot=True, cmap='coolwarm')
plt.title('Correlation Matrix')
plt.show()

```

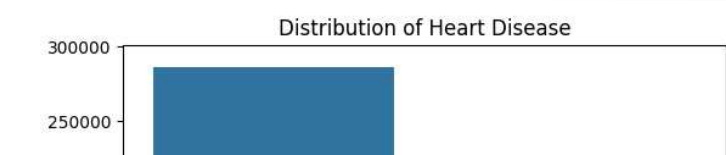


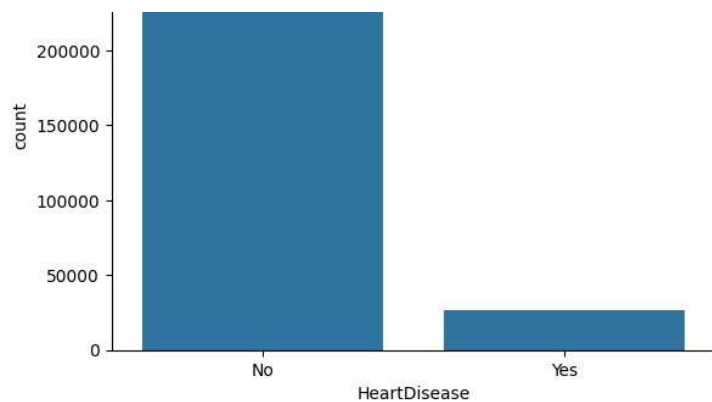
In [14]:

```

# Distribution of Target Variable
sns.countplot(x='HeartDisease', data=data)
plt.title('Distribution of Heart Disease')
plt.show()

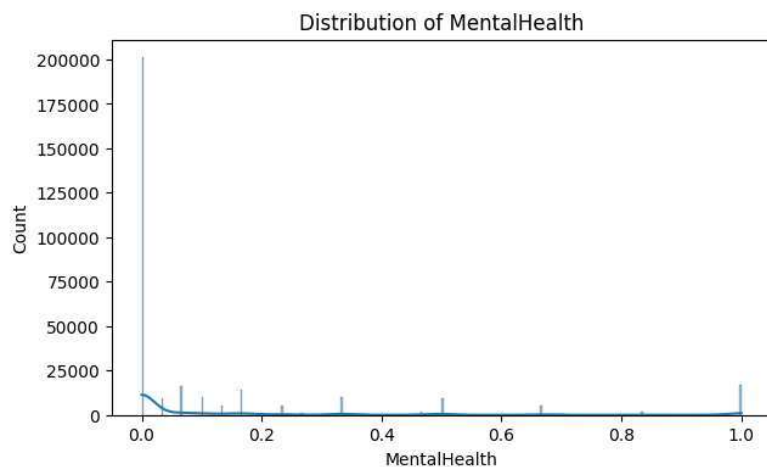
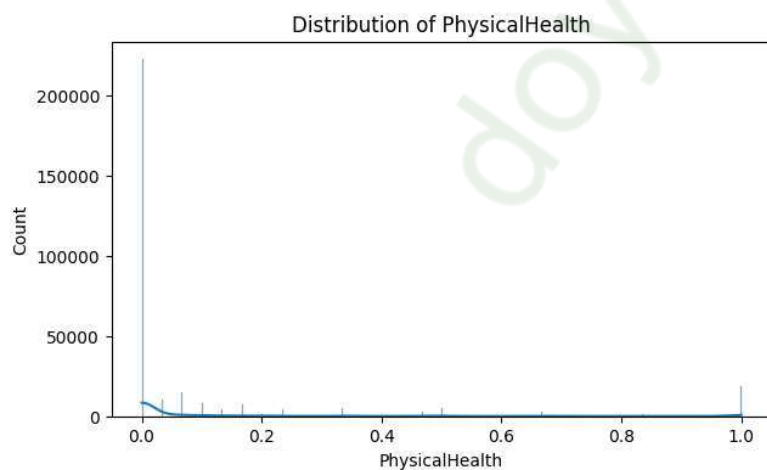
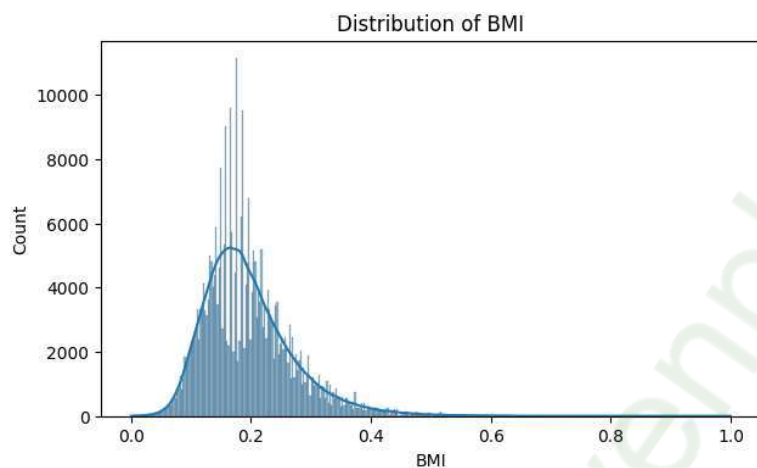
```

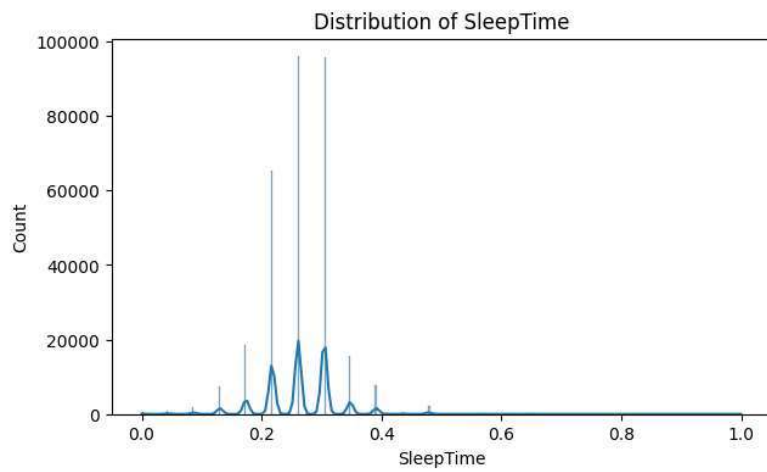




In [15]:

```
# Visualize distributions of numerical features
numerical_features = ['BMI', 'PhysicalHealth', 'MentalHealth', 'SleepTime']
for feature in numerical_features:
    plt.figure(figsize=(7, 4))
    sns.histplot(data=data, x=feature, kde=True)
    plt.title(f"Distribution of {feature}")
    plt.show()
```





In [46]:

```
# Assuming `data` is DataFrame
# Convert boolean columns to int
bool_columns = data.select_dtypes(include=['bool']).columns
data[bool_columns] = data[bool_columns].astype(int)

# Adjusting the categorical columns definition for TableOne
categorical_columns = list(data.select_dtypes(include=['object', 'category']).columns) # Convert to list

# Add bool_columns to categorical if not already present
categorical_columns += [col for col in bool_columns if col not in categorical_columns]

# Preliminary univariate statistical tests using TableOne
print("\nUnivariate statistical tests:")
try:
    # Attempt to create TableOne object
    mytable = TableOne(data,
                       columns=data.columns.tolist(),
                       categorical=categorical_columns,
                       groupby='HeartDisease',
                       pval=True)

    # Print the table
    print(mytable.tabulate(tablefmt="fancy_grid"))
except Exception as e:
    # If an error occurs, print the error message
    print("Error:", e)
```

Univariate statistical tests:

		Missing	Overall	No	Yes	P-Value
n			313014	286430	26584	
HeartDisease, n (%)	No	0	286430 (91.5)	286430 (100.0)		<0.001
	Yes		26584 (8.5)		26584 (100.0)	
BMI, mean (SD)		0	0.2 (0.1)	0.2 (0.1)	0.2 (0.1)	<0.001
Smoking, mean (SD)		0	0.4 (0.5)	0.4 (0.5)	0.6 (0.5)	<0.001
AlcoholDrinking, mean (SD)		0	0.1 (0.3)	0.1 (0.3)	0.0 (0.2)	<0.001
Stroke, mean (SD)		0	0.0 (0.2)	0.0 (0.2)	0.2 (0.4)	<0.001
PhysicalHealth, mean (SD)		0	0.1 (0.3)	0.1 (0.2)	0.3 (0.4)	<0.001
MentalHealth, mean (SD)		0	0.1 (0.3)	0.1 (0.3)	0.2 (0.3)	<0.001
DiffWalking, mean (SD)		0	0.1 (0.3)	0.1 (0.3)	0.4 (0.5)	<0.001
Sex, mean (SD)		0	0.5 (0.5)	0.5 (0.5)	0.6 (0.5)	<0.001
PhysicalActivity, mean (SD)		0	0.8 (0.4)	0.8 (0.4)	0.6 (0.5)	<0.001
SleepTime, mean (SD)		0	0.3 (0.1)	0.3 (0.1)	0.3 (0.1)	<0.001
Asthma, mean (SD)		0	0.1 (0.3)	0.1 (0.3)	0.2 (0.4)	<0.001
KidneyDisease, mean (SD)		0	0.0 (0.2)	0.0 (0.2)	0.1 (0.3)	<0.001
SkinCancer, mean (SD)		0	0.1 (0.3)	0.1 (0.3)	0.2 (0.4)	<0.001
AgeCategory_18-24, n (%)	0	0	292123 (93.3)	265664 (92.8)	26459 (99.5)	<0.001
	1		20891 (6.7)	20766 (7.2)	125 (0.5)	
AgeCategory_25-29, n (%)	0	0	296234 (94.6)	269781 (94.2)	26453 (99.5)	<0.001
	1		16780 (5.4)	16649 (5.8)	131 (0.5)	
AgeCategory_30-34, n (%)	0	0	294467 (94.1)	268108 (93.6)	26359 (99.2)	<0.001
	1		18547 (5.9)	18322 (6.4)	225 (0.8)	

AgeCategory_35-39, n (%)	0	0	292737 (93.5)	266441 (93.0)	26296 (98.9)	<0.001
	1		20277 (6.5)	19989 (7.0)	288 (1.1)	
AgeCategory_40-44, n (%)	0	0	292308 (93.4)	266198 (92.9)	26110 (98.2)	<0.001
	1		20706 (6.6)	20232 (7.1)	474 (1.8)	
AgeCategory_45-49, n (%)	0	0	291621 (93.2)	265761 (92.8)	25860 (97.3)	<0.001
	1		21393 (6.8)	20669 (7.2)	724 (2.7)	
AgeCategory_50-54, n (%)	0	0	288158 (92.1)	262914 (91.8)	25244 (95.0)	<0.001
	1		24856 (7.9)	23516 (8.2)	1340 (5.0)	
AgeCategory_55-59, n (%)	0	0	283989 (90.7)	259531 (90.6)	24458 (92.0)	<0.001
	1		29025 (9.3)	26899 (9.4)	2126 (8.0)	
AgeCategory_60-64, n (%)	0	0	280211 (89.5)	256854 (89.7)	23357 (87.9)	<0.001
	1		32803 (10.5)	29576 (10.3)	3227 (12.1)	
AgeCategory_65-69, n (%)	0	0	279765 (89.4)	257175 (89.8)	22590 (85.0)	<0.001
	1		33249 (10.6)	29255 (10.2)	3994 (15.0)	
AgeCategory_70-74, n (%)	0	0	282874 (90.4)	260978 (91.1)	21896 (82.4)	<0.001
	1		30140 (9.6)	25452 (8.9)	4688 (17.6)	
AgeCategory_75-79, n (%)	0	0	292129 (93.3)	269483 (94.1)	22646 (85.2)	<0.001
	1		20885 (6.7)	16947 (5.9)	3938 (14.8)	
AgeCategory_80 or older, n (%)	0	0	289552 (92.5)	268272 (93.7)	21280 (80.0)	<0.001
	1		23462 (7.5)	18158 (6.3)	5304 (20.0)	
Race_American Indian/Alaskan Native, n (%)	0	0	307971 (98.4)	281914 (98.4)	26057 (98.0)	<0.001
	1		5043 (1.6)	4516 (1.6)	527 (2.0)	
Race_Asian, n (%)	0	0	305285 (97.5)	278958 (97.4)	26327 (99.0)	<0.001
	1		7729 (2.5)	7472 (2.6)	257 (1.0)	
Race_Black, n (%)	0	0	290702 (92.9)	265805 (92.8)	24897 (93.7)	<0.001
	1		22312 (7.1)	20625 (7.2)	1687 (6.3)	
Race_Hispanic, n (%)	0	0	286330 (91.5)	261128 (91.2)	25202 (94.8)	<0.001
	1		26684 (8.5)	25302 (8.8)	1382 (5.2)	
Race_Other, n (%)	0	0	302472 (96.6)	276727 (96.6)	25745 (96.8)	0.047
	1		10542 (3.4)	9703 (3.4)	839 (3.2)	
Race_White, n (%)	0	0	72310 (23.1)	67618 (23.6)	4692 (17.6)	<0.001
	1		240704 (76.9)	218812 (76.4)	21892 (82.4)	
Diabetic_No, n (%)	0	0	43361 (13.9)	34296 (12.0)	9065 (34.1)	<0.001
	1		269653 (86.1)	252134 (88.0)	17519 (65.9)	
Diabetic_Yes, n (%)	0	0	272212 (87.0)	254585 (88.9)	17627 (66.3)	<0.001
	1		40802 (13.0)	31845 (11.1)	8957 (33.7)	
Diabetic_Yes (during pregnancy), n (%)	0	0	310455 (99.2)	283979 (99.1)	26476 (99.6)	<0.001
	1		2559 (0.8)	2451 (0.9)	108 (0.4)	
GenHealth_Excellent, n (%)	0	0	246919 (78.9)	221800 (77.4)	25119 (94.5)	<0.001
	1		66095 (21.1)	64630 (22.6)	1465 (5.5)	
GenHealth_Fair, n (%)	0	0	279464 (89.3)	259757 (90.7)	19707 (74.1)	<0.001
	1		33550 (10.7)	26673 (9.3)	6877 (25.9)	
GenHealth_Good, n (%)	0	0	222379 (71.0)	205060 (71.6)	17319 (65.1)	<0.001
	1		90635 (29.0)	81370 (28.4)	9265 (34.9)	
GenHealth_Poor, n (%)	0	0	302072 (96.5)	279226 (97.5)	22846 (85.9)	<0.001
	1		10942 (3.5)	7204 (2.5)	3738 (14.1)	
GenHealth_Very good, n (%)	0	0	201222 (64.3)	179877 (62.8)	21345 (80.3)	<0.001
	1		111792 (35.7)	106553 (37.2)	5239 (19.7)	
Cluster, mean (SD)		0	0.4 (0.5)	0.4 (0.5)	0.2 (0.4)	<0.001

3. Unsupervised learning - Clustering

In this section, unsupervised learning techniques, particularly clustering methods, are explored. PCA (Principal Component Analysis) is employed for dimensionality reduction, and KMeans clustering is utilized to uncover patterns and groupings within the data. The primary focus is on visualizing patient clusters based on heart disease risk factors, aiming to reveal potential insights and relationships. Prioritizing this exploration phase allows for valuable understanding of the data's inherent structure before proceeding to subsequent analyses.

In [18]:

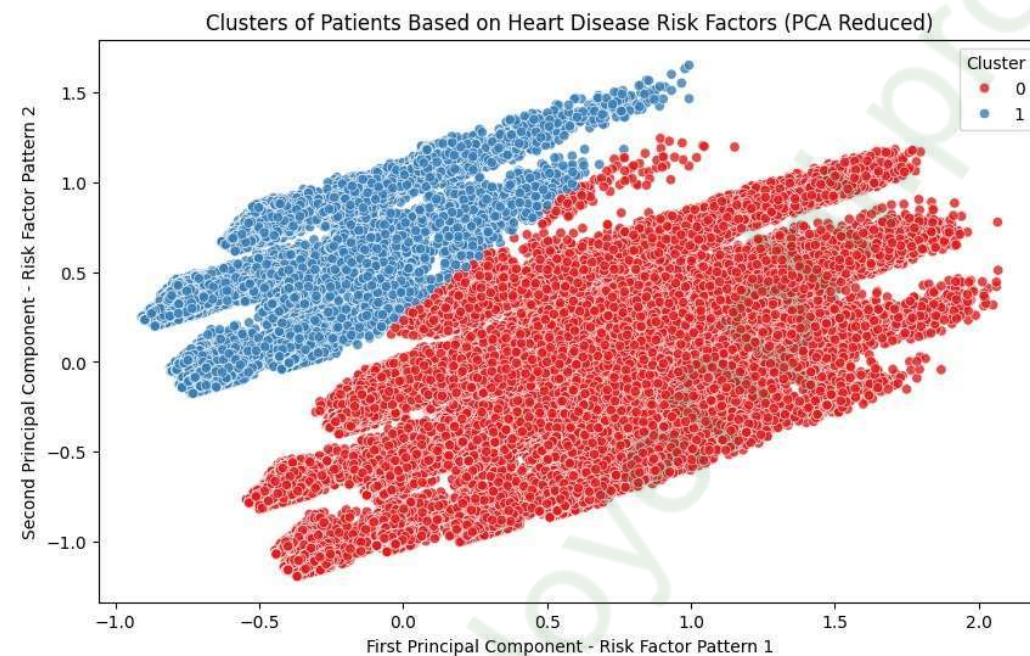
```
# Dimensionality Reduction using PCA
pca = PCA(n_components=2)
data_pca = pca.fit_transform(data.drop(columns=['HeartDisease']))
```

In [19]:

```
# KMeans clustering
kmeans = KMeans(n_clusters=2, random_state=1, n_init=10)
data['Cluster'] = kmeans.fit_predict(data_pca)
```

In [20]:

```
# Visualizing Clusters
plt.figure(figsize=(10, 6))
sns.scatterplot(x=data_pca[:,0], y=data_pca[:,1], hue=data['Cluster'], palette='Set1', alpha=0.8)
plt.title("Clusters of Patients Based on Heart Disease Risk Factors (PCA Reduced)")
plt.xlabel("First Principal Component - Risk Factor Pattern 1")
plt.ylabel("Second Principal Component - Risk Factor Pattern 2")
plt.legend(title='Cluster')
plt.show()
```



4. Model building and fitting

This section elaborates on the selection, training, and optimization of machine learning models aimed at predicting heart disease risk. It covers details regarding model training, parameter tuning using GridSearchCV, and model evaluation based on accuracy. Specifically, various machine learning algorithms including KNeighborsClassifier, LogisticRegression, and RandomForestClassifier, will be utilized for model building. Each model will undergo rigorous testing and evaluation, with the most accurate model being selected based on predefined criteria.

In [21]:

```
# Prepare data for modeling - Split data
X = data.drop(columns=['HeartDisease'])
y = data['HeartDisease']
X_train, X_test, y_train, y_test = train_test_split(X, y, test_size=0.25, random_state=1, stratify=y)
```

K-Nearest Neighbors (KNN): KNN is chosen for this project due to its simplicity, non-parametric nature, and ability to predict heart disease risk based on the similarity between data points. Its adaptability to various classification tasks and localized decision-making process make it suitable for capturing complex patterns in the data without strict assumptions. Despite its simplicity, KNN remains effective for exploring heart disease prediction, although its performance should be compared with other algorithms for validation.

In [22]:

```
# Initialize the K-Nearest Neighbors (KNN) classifier with default parameters. KNN is a simple, instance-based learning algorithm
# where the class of a sample is determined by the majority class among its k nearest neighbors.
knn = KNeighborsClassifier()

# Define the parameter grid for GridSearchCV to tune the KNN classifier. The grid specifies a range for 'n_neighbors'
# (i.e., the value of k in KNN) from 1 to 49. The optimal number of neighbors will be determined through cross-validation.
param_grid_knn = {'n_neighbors': np.arange(1, 50)}

# Perform grid search with 4-fold cross-validation to find the best number of neighbors ('n_neighbors') for the KNN classifier.
# GridSearchCV evaluates different 'n_neighbors' values to find the best parameter setting that maximizes model performance.
knn_gscv = GridSearchCV(knn, param_grid_knn, cv=4).fit(X_train, y_train)

# Retrieve the best KNN model (with the optimal 'n_neighbors' value) found by the grid search.
knn_opti = knn_gscv.best_estimator_
```

```
# Convert X_test to a NumPy array and ensure it is C-contiguous. This step may be necessary for performance reasons,
# as KNN can be sensitive to the layout of the input data in memory.
X_test_c_contiguous = np.asarray(X_test).copy(order='C')

# Calculate the accuracy of the optimized KNN model on the test dataset. This measures how well the model performs on
# unseen data, using the test set.
knn_accuracy = accuracy_score(y_test, knn_opti.predict(X_test_c_contiguous))

# Save the optimized KNN model to a file named 'knn_opti.joblib'. This allows the model to be used later without retraining,
# making it convenient for deployment or further evaluation.
dump(knn_opti, "knn_opti.joblib")
```

```
c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\model_selection\_validation.py:824: UserWarning: Scoring failed.
The score on this train-test partition for these parameters will be set to nan. Details:
Traceback (most recent call last):
```

```
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\model_selection\_validation.py", line 813, in _score
    scores = scorer(estimator, X_test, y_test)
    ~~~~~
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\metrics\_scorer.py", line 527, in __call__
    return estimator.score(*args, **kwargs)
    ~~~~~
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\base.py", line 705, in score
    return accuracy_score(y, self.predict(X), sample_weight=sample_weight)
    ~~~~~
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\neighbors\_classification.py", line 246, in predict
    if self._fit_method == "brute" and ArgKminClassMode.is_usable_for(
    ~~~~~
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\metrics\_pairwise_distances_reduction\_dispatcher.py", line 471, in is_usable_for
    ArgKmin.is_usable_for(X, Y, metric)
    ~~~~~
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\metrics\_pairwise_distances_reduction\_dispatcher.py", line 115, in is_usable_for
    and (is_numpy_c_ordered(X) or is_valid_sparse_matrix(X))
    ~~~~~
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\metrics\_pairwise_distances_reduction\_dispatcher.py", line 99, in is_numpy_c_ordered
    return hasattr(X, "flags") and X.flags.c_contiguous
    ~~~~~
AttributeError: 'Flags' object has no attribute 'c_contiguous'
```

```
warnings.warn(
c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\model_selection\_validation.py:824: UserWarning: Scoring failed.
The score on this train-test partition for these parameters will be set to nan. Details:
Traceback (most recent call last):
```

```
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\model_selection\_validation.py", line 813, in _score
    scores = scorer(estimator, X_test, y_test)
    ~~~~~
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\metrics\_scorer.py", line 527, in __call__
    return estimator.score(*args, **kwargs)
    ~~~~~
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    return accuracy_score(y, self.predict(X), sample_weight=sample_weight)
    ~~~~~
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\neighbors\_classification.py", line 246, in predict
    if self._fit_method == "brute" and ArgKminClassMode.is_usable_for(
    ~~~~~
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    ArgKmin.is_usable_for(X, Y, metric)
    ~~~~~
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\metrics\_pairwise_distances_reduction\_dispatcher.py", line 115, in is_usable_for
    and (is_numpy_c_ordered(X) or is_valid_sparse_matrix(X))
    ~~~~~
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\metrics\_pairwise_distances_reduction\_dispatcher.py", line 99, in is_numpy_c_ordered
    return hasattr(X, "flags") and X.flags.c_contiguous
    ~~~~~
AttributeError: 'Flags' object has no attribute 'c_contiguous'
```

```
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c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\model_selection\_validation.py:824: UserWarning: Scoring failed.
The score on this train-test partition for these parameters will be set to nan. Details:
Traceback (most recent call last):
```

```
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\model_selection\_validation.py", line 813, in _score
    scores = scorer(estimator, X_test, y_test)
    ~~~~~
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\metrics\_scorer.py", line 527, in __call__
    return estimator.score(*args, **kwargs)
    ~~~~~
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\base.py", line 705, in score
    return accuracy_score(y, self.predict(X), sample_weight=sample_weight)
    ~~~~~
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\neighbors\_classification.py", line 246, in predict
    if self._fit_method == "brute" and ArgKminClassMode.is_usable_for(
    ~~~~~
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\metrics\_pairwise_distances_reduction\_dispatcher.py", line 471, in is_usable_for
    ArgKmin.is_usable_for(X, Y, metric)
    ~~~~~
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\metrics\_pairwise_distances_reduction\_dispatcher.py", line 115, in is_usable_for
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    ~~~~~
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\metrics\_pairwise_distances_reduction\_dispatcher.py", line 99, in is_numpy_c_ordered
    return hasattr(X, "flags") and X.flags.c_contiguous
    ~~~~~
AttributeError: 'Flags' object has no attribute 'c_contiguous'
```

```
warnings.warn(
c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\model_selection\_validation.py:824: UserWarning: Scoring failed.
The score on this train-test partition for these parameters will be set to nan. Details:
```



```
# cross-validation results over the parameter grid.
logReg_opti = logReg_gscv.best_estimator_

# Evaluate the accuracy of the optimized Logistic Regression model on the test set to determine its performance.
logreg_accuracy = accuracy_score(y_test, logReg_opti.predict(X_test))

# Save the optimized Logistic Regression model to a file named 'logReg_opti.joblib' for later use or deployment,
# enabling the model to be loaded and used without retraining.
dump(logReg_opti, "logReg_opti.joblib")
```

```

solver = _check_solver(self.solver, self.penalty, self.dual)
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\linear_model\_logistic.py", line 56, in _check_solver
raise ValueError(
ValueError: Solver lbfgs supports only 'l2' or 'none' penalties, got l1 penalty.

warnings.warn(some_fits_failed_message, FitFailedWarning)
c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\model_selection\_search.py:976: UserWarning: One or more of the t
est scores are non-finite: [
nan 0.91678736 0.91679588      nan 0.91678736 0.91679588      nan 0.91677458 0.91679588
nan 0.91678736 0.91679588      nan 0.91678736 0.91679588      nan 0.91678736 0.91679588
nan 0.9167831 0.91679588      nan 0.9167831 0.91679588      nan 0.9167831 0.91679588
nan 0.91679162 0.91679588      nan 0.9167831 0.91679588      nan 0.9167831 0.91679588
nan 0.9167831 0.91679588      nan 0.91677884 0.91679588      nan 0.91677884 0.91679588
nan 0.9167831 0.91679588      nan 0.91677884 0.91679588      nan 0.91677884 0.91679588
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nan 0.9167831 0.91679588      nan 0.91678736 0.91679588      nan 0.91678736 0.91679588
nan 0.91679162 0.91679588      nan 0.9167831 0.91679588      nan 0.9167831 0.91679588
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nan 0.91678736 0.91679588      nan 0.9167831 0.91679588      nan 0.9167831 0.91679588
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nan 0.91679162 0.91679588      nan 0.9167831 0.91679588      nan 0.9167831 0.91679588
nan 0.91678736 0.91679588      nan 0.91679162 0.91679588      nan 0.91679162 0.91679588
nan 0.91679588 0.91679588      nan 0.9167831 0.91679588      nan 0.9167831 0.91679588
nan 0.91678736 0.91679588      nan 0.9167831 0.91679588      nan 0.9167831 0.91679588
nan 0.91678736 0.91679588      nan 0.9167831 0.91679588      nan 0.9167831 0.91679588]
warnings.warn(

```

Out[23]:

```
['logReg_opti.joblib']
```

Random Forest Classifier: The Random Forest Classifier is chosen for its ability to handle complex datasets like medical data, accurately predicting heart disease. Its capability to capture nonlinear relationships and manage missing data ensures robust performance. Additionally, its ensemble approach reduces overfitting, enhancing model reliability and interpretability for heart disease risk assessment.

In [24]:

```

# Initialize the Random Forest classifier without specifying any hyperparameters.
rf = RandomForestClassifier()

# Define the parameter grid for the Random Forest classifier. It specifies a range of values for 'n_estimators'
# (the number of trees in the forest) and 'max_features' (the number of features to consider when looking for the best split)
# to find the optimal combination that improves model performance.
param_grid_rf = {'n_estimators': [600, 730, 800], 'max_features': list(range(1, 3))}

# Perform grid search with 5-fold cross-validation on the Random Forest classifier using the defined parameter grid.
# The grid search explores different combinations of 'n_estimators' and 'max_features' to find the best settings based on the training data.
# 'n_jobs=-1' allows the process to use all available CPU cores for faster completion.
rf_gscv = GridSearchCV(rf, param_grid_rf, cv=5, n_jobs=-1).fit(X_train, y_train)

# Retrieve the best estimator (Random Forest model) found by the grid search. This model is considered optimized
# based on the cross-validation results over the specified parameter grid.
rf_opti = rf_gscv.best_estimator_

# Calculate and print the accuracy of the optimized Random Forest model on the test dataset to evaluate its performance.
rf_accuracy = accuracy_score(y_test, rf_opti.predict(X_test))

# Save the optimized Random Forest model to a file named 'randomForest_opti.joblib' for future use,
# allowing the model to be readily available without needing to retrain.
dump(rf_opti, "randomForest_opti.joblib")

```

```

c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\model_selection\_validation.py:425: FitFailedWarning:
24 fits failed out of a total of 30.
The score on these train-test partitions for these parameters will be set to nan.
If these failures are not expected, you can try to debug them by setting error_score='raise'.

```

Below are more details about the failures:

8 fits failed with the following error:

Traceback (most recent call last):

```

File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\model_selection\_validation.py", line 732, in _fit_and_score
estimator.fit(X_train, y_train, **fit_params)
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\base.py", line 1151, in wrapper
return fit_method(estimator, *args, **kwargs)
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\ensemble\_forest.py", line 456, in fit
trees = Parallel(
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\utils\parallel.py", line 65, in __call__
return super().__call__(iterable_with_config)
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\joblib\parallel.py", line 1863, in __call__
return output if self.return_generator else list(output)
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\joblib\parallel.py", line 1792, in _get_sequential_output
res = func(*args, **kwargs)
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\utils\parallel.py", line 127, in __call__
return self.function(*args, **kwargs)
File "c:\Users\Asus\scoop\apps\python\current\Lib\site-packages\sklearn\ensemble\_forest.py", line 188, in _parallel_build_trees
tree.fit(X, y, sample_weight=curr_sample_weight, check_input=False)

```

Out [24]:

```
['randomForest_opti.joblib']
```

In [25]:

```
# Results
print(f"K-Nearest Neighbors Accuracy: {knn_accuracy}")
print(f"Logistic Regression Accuracy: {logreg_accuracy}")
print(f"Random Forest Accuracy: {rf_accuracy}")
```

```
K-Nearest Neighbors Accuracy: 0.8713164822245508
Logistic Regression Accuracy: 0.915812610218008
Random Forest Accuracy: 0.9035065300176349
```

5. Results and Model Deployment:

Based on the results obtained from the three models, it is evident that Logistic Regression outperforms both K-Nearest Neighbors (KNN) and Random Forest Classifier in terms of accuracy. With an accuracy of 91.58%, Logistic Regression demonstrates superior predictive power in identifying heart disease. While both KNN and Random Forest also achieve respectable accuracies of 87.13% and 90.40%, respectively, Logistic Regression proves to be the most effective model for this task. Therefore, Logistic Regression is the recommended choice for accurately predicting heart disease in this dataset.

The following Python code is intended for deploying a trained machine learning model using Streamlit. The model is loaded from a file named "logReg_opti.joblib". The main function of the code creates a Streamlit web application titled "Heart Disease Prediction". It includes input fields for various features such as age, sex, chest pain type, blood pressure, cholesterol, blood sugar, resting ECG, max heart rate, exercise-induced angina, and oldpeak. The user can input values for these features, and upon clicking the "Predict" button, the code makes predictions using the loaded model and displays the result. This code is saved in a separate Python file named "heart_disease_prediction_app.py" within the same folder for deployment purposes.

In [1]:

```
import streamlit as st
import pandas as pd
from joblib import load

# Load the trained model
model = load('logReg_opti.joblib') # Adjust the filename as per your model

# Define the function to make predictions
def predict(input_features):
    try:
        prediction = model.predict(input_features)
        return prediction
    except Exception as e:
        return str(e)

# Create the Streamlit web application
def main():
    st.title('Heart Disease Prediction')

    # Add input fields for features
    heart_disease = st.radio('Heart Disease', ['Yes', 'No'])
    bmi = st.slider('BMI', 10.0, 50.0, 25.0)
    smoking = st.selectbox('Smoking', ['Yes', 'No'])
    alcohol_drinking = st.selectbox('Alcohol Drinking', ['Yes', 'No'])
    stroke = st.selectbox('Stroke', ['Yes', 'No'])
    physical_health = st.slider('Physical Health', 0, 100, 50)
    mental_health = st.slider('Mental Health', 0, 100, 50)
    diff_walking = st.selectbox('Difficulty Walking', ['Yes', 'No'])
    sex = st.selectbox('Sex', ['Male', 'Female'])
    age_category = st.selectbox('Age Category', ['18-24', '25-29', '30-34', '35-39', '40-44', '45-49', '50-54', '55-59', '60-64', '65-69', '70-74', '75-79', '80 or older'])
    race = st.selectbox('Race', ['White', 'Black', 'Asian', 'Hispanic'])
    diabetic = st.selectbox('Diabetic', ['Yes', 'No'])
    physical_activity = st.selectbox('Physical Activity', ['Yes', 'No'])
    gen_health = st.selectbox('General Health', ['Excellent', 'Very Good', 'Good', 'Fair', 'Poor'])
    sleep_time = st.slider('Sleep Time', 2, 12, 8)
    asthma = st.selectbox('Asthma', ['Yes', 'No'])
    kidney_disease = st.selectbox('Kidney Disease', ['Yes', 'No'])
    skin_cancer = st.selectbox('Skin Cancer', ['Yes', 'No'])

    # Convert categorical features to numerical format
    categorical_mapping = {'Yes': 1, 'No': 0, 'Male': 1, 'Female': 0,
                           '18-24': 0, '25-29': 1, '30-34': 2, '35-39': 3, '40-44': 4, '45-49': 5,
                           '50-54': 6, '55-59': 7, '60-64': 8, '65-69': 9, '70-74': 10, '75-79': 11, '80 or older': 12,
                           'White': 0, 'Black': 1, 'Asian': 2, 'Hispanic': 3,
                           'Excellent': 0, 'Very Good': 1, 'Good': 2, 'Fair': 3, 'Poor': 4}

    heart_disease_numeric = categorical_mapping[heart_disease]
    smoking_numeric = categorical_mapping[smoking]
    alcohol_drinking_numeric = categorical_mapping[alcohol_drinking]
    stroke_numeric = categorical_mapping[stroke]
    diff_walking_numeric = categorical_mapping[diff_walking]
    diabetic_numeric = categorical_mapping[diabetic]
    physical_activity_numeric = categorical_mapping[physical_activity]
    asthma_numeric = categorical_mapping[asthma]
    kidney_disease_numeric = categorical_mapping[kidney_disease]
    skin_cancer_numeric = categorical_mapping[skin_cancer]

    # Convert input into a DataFrame
    input_data = pd.DataFrame({
        'HeartDisease': [heart_disease_numeric],
        'BMI': [bmi],
        'Smoking': [smoking_numeric],
        'AlcoholDrinking': [alcohol_drinking_numeric],
        'Stroke': [stroke_numeric],
        'PhysicalHealth': [physical_health],
        'MentalHealth': [mental_health],
        'DiffWalking': [diff_walking_numeric],
        'Sex': [sex],
```

```

        'AgeCategory': [age_category],
        'Race': [race],
        'Diabetic': [diabetic_numeric],
        'PhysicalActivity': [physical_activity_numeric],
        'GenHealth': [gen_health],
        'SleepTime': [sleep_time],
        'Asthma': [asthma_numeric],
        'KidneyDisease': [kidney_disease_numeric],
        'SkinCancer': [skin_cancer_numeric]
    })

    # Make prediction
    if st.button('Predict'):
        prediction = predict(input_data)
        st.write(f'The prediction is: {prediction}')

if __name__ == '__main__':
    main()

```

2024-03-01 11:15:12.746

Warning: to view this Streamlit app on a browser, run it with the following command:

```
streamlit run C:\Users\Asus\AppData\Roaming\Python\Python311\site-packages\ipykernel_launcher.py [ARGUMENTS]
```

After creating the **heart_disease_prediction_app.py** file and running it, I executed the command **streamlit run heart_disease_prediction_app.py** in my terminal to initiate the Streamlit application. This command launched a local server and opened the application in a web browser. Upon opening the application, I was presented with a user interface where I could input various features related to heart disease prediction, such as age, sex, chest pain type, blood pressure, cholesterol, blood sugar, resting ECG, max heart rate, exercise-induced angina, and oldpeak. After inputting these features and clicking the "Predict" button, the application utilized the trained model to make predictions and displayed the result on the browser interface, similar to the image below.

6. Representing results - Dashboard insights

The dashboard illustrates the comparison of three machine learning models for heart disease prediction: K-Nearest Neighbors, Logistic Regression, and Random Forest. Each bar in the plot represents the accuracy achieved by the respective model on the test dataset. The accuracy scores, ranging from approximately 0.87 to 0.92, indicate the proportion of correctly predicted outcomes. Based on these results, it is evident that Logistic Regression achieved the highest accuracy of approximately 91.58%, making it the most effective model for heart disease prediction among the three considered.

In [35]:

```

import seaborn as sns
import matplotlib.pyplot as plt

# Set Seaborn theme and style
sns.set_theme(style='whitegrid')

# Sample data
models = ['K-Nearest Neighbors', 'Logistic Regression', 'Random Forest']
accuracies = [0.871252587727145, 0.9157742735195645, 0.9039793492984384]

# Create a horizontal bar plot using Seaborn
plt.figure(figsize=(8, 6))
ax = sns.barplot(x=accuracies, y=models, palette='mako')

# Set title and axis labels
plt.title('Model Comparison for Heart Disease Prediction', fontsize=16)
plt.xlabel('Accuracy', fontsize=14)
plt.ylabel('Models', fontsize=14)

# Set font size for tick labels
plt.xticks(fontsize=12)
plt.yticks(fontsize=12)

# Show grid lines
plt.grid(True, axis='x', linestyle='--', alpha=0.7)

# Add data labels to each bar
for i, v in enumerate(accuracies):
    ax.text(v + 0.01, i, f'{v:.3f}', color='black', va='center', fontsize=12)

# Show plot
plt.tight_layout()
plt.show()

```

C:\Users\Asus\AppData\Local\Temp\ipykernel_15760\2812506543.py:13: FutureWarning:

Passing `palette` without assigning `hue` is deprecated and will be removed in v0.14.0. Assign the `y` variable to `hue` and set `legend=False` for the same effect.

```
ax = sns.barplot(x=accuracies, y=models, palette='mako')
```

Model Comparison for Heart Disease Prediction





7. Conclusion:

The success of the Predictive Medical Assistant project relies on the availability of resources and technical feasibility, ensuring efficient model training through high-performance computing clusters and expert guidance. Compliance with healthcare regulations, such as HIPAA, is prioritized, with stringent measures like data encryption and access controls ensuring data privacy and security. Integration with existing healthcare systems is facilitated through APIs and standard data exchange formats, enhancing usability in real-world settings. A comprehensive cost-benefit analysis explores economic feasibility and potential health impacts, while scalability and maintenance protocols ensure adaptability and relevance over time. Success metrics tailored to medical diagnostics provide a comprehensive evaluation of model performance, with expected outcomes including high model accuracy, an interactive web application, and comprehensive documentation. This project lays a foundation for future advancements in medical diagnostics and decision support systems, addressing key areas of technical feasibility, compliance, integration, cost-benefit analysis, scalability, maintenance, and success metrics.

STREAMLIT BROWSER INTERFACE

Heart Disease Prediction

Heart Disease

- ☒ Yes
- ☐ No



Smoking

Yes

▼

Alcohol Drinking

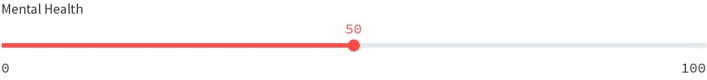
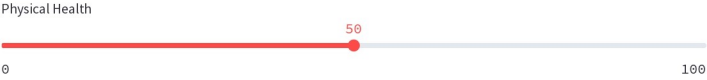
Yes

▼

Stroke

Yes

▼



Difficulty Walking

Yes

▼

Sex

Male

▼

Age Category

18-24

▼

Race

White

▼

Diabetic

Yes

▼

Physical Activity

Yes

▼

General Health

Excellent

▼



Asthma

Yes

▼

Kidney Disease

Yes

▼

Skin Cancer

Yes

▼

Predict