INTRODUCTION

Overview

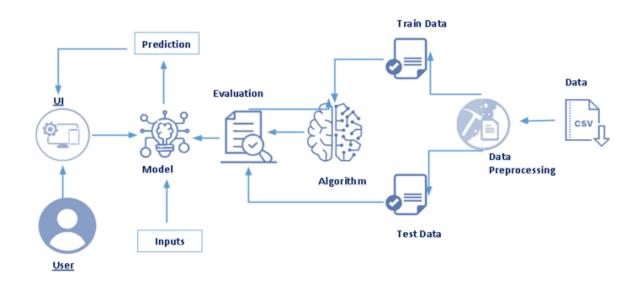
Chronic Kidney Disease (CKD) is a major medical problem and can be cured if treated in the early stages. Usually, people are not aware that medical tests, we take for different purposes could contain valuable information concerning kidney diseases. Consequently, attributes of various medical tests are investigated to distinguish which attributes may contain helpful information about the disease. The information says that it helps us to measure the severity of the problem and we make use of such information to build a machine learning model that predicts Chronic Kidney Disease

Proposed System

In this section, a detailed description of the data set creation, model preparation, and disease prediction has been given. The first action is data collection. Our proposed system collects structured and unstructured data obtained from various sources. After data collection, they are subjected to preprocessing and are split into cleaning and test data sets. Then the training data set is trained with the machine learning algorithms such as CNN and KNN to a number of epochs for improving the accuracy of the prediction results. After multiple epochs, once the desired target is achieved, the developed model is ready for testing.

3.THEORETICAL ANALYSIS

3.1 Block Diagram



3.2 Hardware/Software Designing

- 2 GB ram or above
- Dual core processor or above
- Internet connection

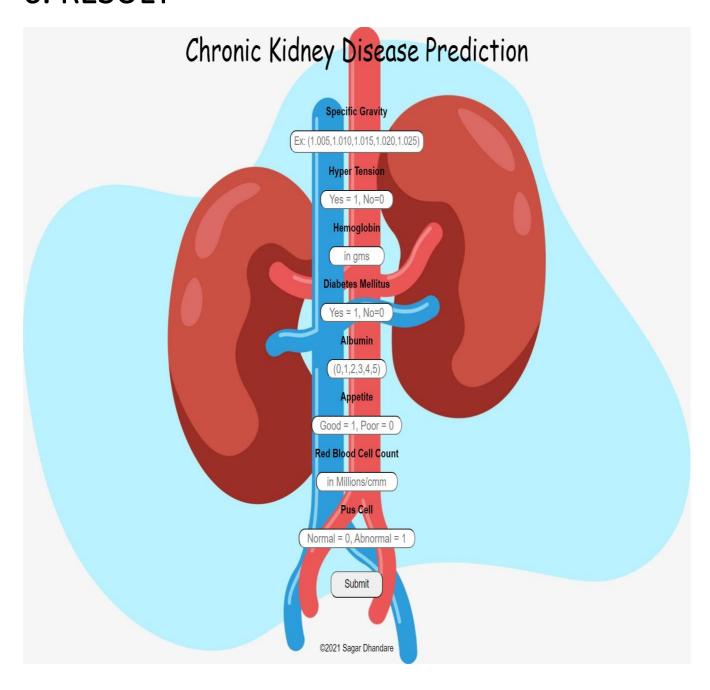
Software requirements

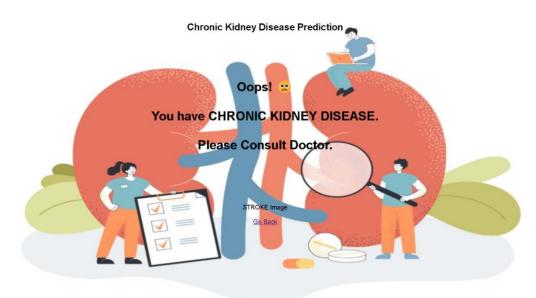
- Anaconda Navigator
- Python packages
- VS Studio

4. FLOWCHART

- User interacts with the UI (User Interface) to fill the information asked.
- Given input is analysed by the model which is integrated.
- Once model analyses the given information , the prediction is showcased on the UI
- 1. Data Collection
 - a. Collect the dataset or create the dataset
- 2. Understanding the data
 - a. Importing the required libraries
 - b. Reading the Dataset
 - c. EDA on Dataset
 - d. Take care of missing data
 - e. Data Visualization
 - f. Cleaning The Text
 - g. Building count vectors with scikit-learn Count-Vectorizer for text classification
 - h. Splitting Data into Train and Test
- 3. Model Building
 - a. Training and testing the model
 - b. Evaluation of Model
 - c. Saving the model
- 4. Application Building
 - a. Create an HTML file
 - b. Build Python Code
- 5. Final UI
 - a. Dashboard Of the flask app

6. RESULT





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6. ADVANTAGES AND DISADVANTAGES

Advantages

- Help physicians to identify effective treatments and best practices.
- Patients exploit better and greater affordable healthcare services.
- Increases in the speed of working with large datasets and rapid report
- generation, faster analysis, improved operational efficiency and
- reduced operating cost.
- Data Mining can extract predictive knowledge from large databases.

Disadvantages

- Data Ownership issues.
- Privacy and Security related to Human Data Administration.
- It Involves privacy issues and security issues and
- Misuse or incorrect information.

8. Applications

- App
- Websites

9. CONCLUSION

Using this project, we can predict if we have chronic kidney disease or not. This program will check the information of our health report and predicts whether we have chronic kidney disease or not. When our health report is bad then it say that we have chronic kidney disease.

10. FUTURE SCOPE

This program allows users to predict if that whether we have chronic kidney disease or not . By the help of this prediction we can take following measures to cure CKD . It helps to take measure for CKD before our kidney completely fails.

Appendix

```
import pandas as pd
import numpy as np
import matplotlib.pyplot as plt
import seaborn as sns
df = pd.read_csv("kidney_disease.csv")
df.info()
df.head(10)
```

```
df.describe()
df["rc"].unique()
df["rc"] = df["rc"].replace("\t?", '0')
df["rc"].unique()
df["rc"] = df["rc"].astype(float)
df["wc"].unique()
df["pcv"] = df["pcv"].replace(("\t?", "\t43"), (0, 43))
df["pcv"] = df["pcv"].astype(float)
df.info()
df.drop(["id"], axis = 1, inplace = True)
df.describe()
obj = (df.select dtypes(include = object)).columns
numeric = (df.select dtypes(include = np.number)).columns
print(numeric, obj)
for i in obj:
  print(i)
  print(df[i].unique())
df["dm"] = df["dm"].replace({"\tno":"no", "\tyes":"yes", " yes":"yes"})
df["cad"] = df["cad"].replace("\tno", "no")
df["classification"] = df["classification"].replace("ckd\t", "ckd")
for i in obj:
  print(i)
  print(df[i].unique())
df.isnull().sum()
```

```
df["rc"].fillna(df["rc"].mean(), inplace=True)
for i in numeric:
  df[i].fillna(df[i].mean(),inplace=True)
for i in obj:
  df[i].fillna(df[i].mode()[0], inplace=True)
df.head(10)
from sklearn.preprocessing import LabelEncoder
for i in obj:
  le = LabelEncoder()
  df[i] = le.fit transform(df[i])
df.info()
df.shape
df.corr()
import seaborn as sns
plt.figure(figsize = (15,15))
sns.heatmap(df.corr(), annot = True, fmt=".2f",linewidth=0.5)
df.corrwith(df.classification).plot(kind="bar", grid=True,figsize=(12,8),
title = "corr with target")
x = df.drop(["classification", "pot", "ba", "wc", "age", 'su', "ane"], axis =
1)
x.columns
x.columns=['Blood_pressure', "Specafic_gravity", "albumin",
"red_blood_cells", 'pus_cell', 'pus_cell_clumps',
"blood_glucose_random", 'blood_urea', 'serum_creatinine', 'sodium',
"hemoglobin", 'packed cell volume', 'red blood cell count',
```

```
'hypertension', 'diabetesmellitus', 'coronary artery disease',
'appetite', 'pedal edema']
x.columns
x.head(10)
y = df["classification"]
from sklearn.model selection import train test split
x_train,x_test,y_train, y_test = train_test_split(x,y, test_size = 0.25)
x train.shape, x test.shape, y train.shape
from sklearn.ensemble import RandomForestClassifier
from sklearn.naive bayes import GaussianNB
from sklearn.metrics import classification report,
confusion matrix, accuracy score
ran fore = RandomForestClassifier(n estimators=100)
ran fore.fit(x train, y train)
rf pred = ran fore.predict(x test)
rf score = round(ran fore.score(x train, y train)*100, 2)
rf test score = round(ran fore.score(x test, y test)*100, 2)
print("Random forest train score = \n", rf_score)
print("Random forest test score = \n", rf test score)
print("accuracy = \n", accuracy score(y test,rf pred))
print("confusion Matrix : \n", confusion_matrix(y_test, rf_pred))
print(classification report(y test, rf pred))
import pickle
pickle.dump(ran_fore, open("CKD.pkl",'wb'))
```