PROJECT REPORT DOCUMENT

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TEAM ID:PNT2022TMID45417

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1.INTRODUCTION

1.1 PROJECT OVERVIEW

The chronic kidney disease popularly called as (CKD). The main causes of CKD are blood pressure, blood sugar and glucose level, strain on small blood vessels, excess sedimentation of salts on the kidney and also due to age factors. The end stage pf CKD leads to kidney failure that makes kidney unable to remove wastes from the body. And also fails to control the fluid levels in the body. It shows no symptoms at the earlier stages and results in high risks. Using, some machine learning models we can predict the CKD as early as possible. Data science using Machine learning is the most efficient to predict the CKD and prevent the patients from the risks.

1.2 PURPOSE

The experimental procedure concludes that advances in machine learning, with assist of predictive analytics, represent a promising setting by which to recognize intelligent solutions, which in turn prove the ability of predication in the kidney disease domain and beyond.

2.LITERATURE SURVEY

2.1 EXISTING PROBLEM

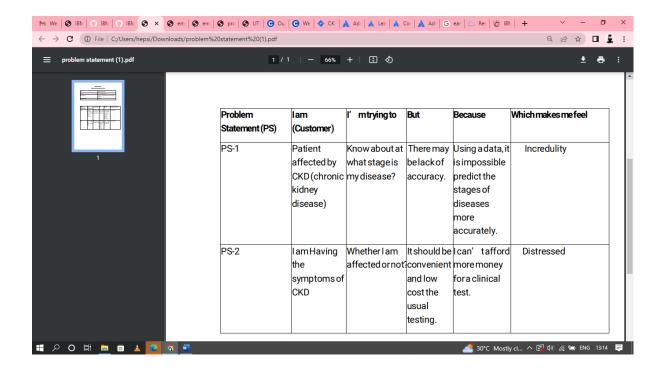
Shanila Yunus Yashfi, Md Ashikul Islam, Pritilata, Nazmus Sakib, Tanzila Islam, Mohammad Shahbaaz, Sadaf Salman Pantho [1] - "They have used UCI dataset and real time dataset and processed them. handled missing data, trained it and made a Random Forest and ANN model. Also implemented these two algorithms in python language. The gain using Random Forest algorithm is 97.12% and ANN is 94.5% respectively which is relatively very good" Siddheshwar Tekale, Pranjal Shingavi, Sukanya Wandhekar, Ankit Chatorikar[2] - " have analyzed 14 different attributes related to CKD patients and predicted accuracy for different machine learning algorithms like Decision tree and Support Vector Machine. From the results analysis, it is observed that the decision tree algorithms give the accuracy of 91.75% and SVM gives accuracy of 96.75%." PANKAJ CHITTORA 1, SANDEEP CHAURASIA1, (Senior Member, IEEE), PRASUN CHAKRABARTI2,3, (Senior Member, IEEE), GAURAV KUMAWAT1, TULIKA CHAKRABARTI4, ZBIGNIEW LEONOWICZ 5, (Senior Member, IEEE), MICHAŁ JASIŃSKI 5, (Member, IEEE), ŁUKASZ JASIŃSKI 5, RADOMIR GONO 6, (Senior Member, IEEE), ELŻBIETA JASIŃSKA 7, AND VADIM BOLSHEV 8 [3] - "three different techniques have been applied: correlation-based feature selection, Wrapper method and LASSO regression. In this perception, seven classifiers algorithm were applied viz. artificial neural

network, C5.0, logistic regression, CHAID, linear support vector machine (LSVM), K-Nearest neighbors and random tree. For each classifier, the results were computed based on full features, selected features by CFS, selected features by Wrapper, selected features by LASSO regression, SMOTE with selected features by LASSO, SMOTE with full features. It was observed that LSVM achieved the highest accuracy of 98.86% in SMOTE with full features." Saurabh Pal[4]-" Have used chronic kidney disease dataset collected from UCI machine leaning repository developed a chronic kidney disease prediction model using three machine learning classifiers Logistic Regression, Decision Tree and Support Vector Machine to measure the performance of the prediction model. After applying the base classifiers, we find decision tree classifier obtained better results in terms of Accuracy, Precision, Recall, F-score as 95.92%, 0.99, 0.98, and 0.98, respectively. The highest accuracy The application of MLAs in kidney diseases may enhance the ability of clinicians to predict CKD and RF, thus improving diagnostic assistance and providing suitable therapeutic decisions. However, it is necessary to improve the development process of MLA tools.

2.2 REFERENCES

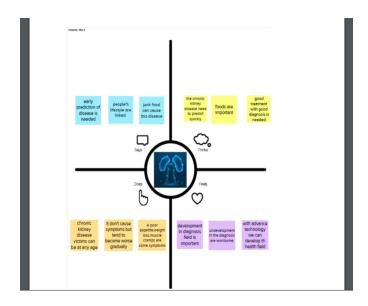
[1] S. Y. Yashfi et al., "Risk Prediction of Chronic Kidney Disease Using Machine Learning Algorithms," 2020 11th International Conference on Computing, Communication and Networking Technologies (ICCCNT), 2020, pp. 1-5, Doi: 10.1109/ICCCNT49239.2020.9225548. [2] International Journal of Advanced Research in Computer and Communication Engineering Vol. 7, Issue 10, October 2018 Copyright to IJARCCE DOI 10.17148/IJARCCE.2018.71021 92 "Prediction of Chronic Kidney Disease Using Machine Learning Algorithm" Siddheshwar Tekale1, Pranjal Shingavi2, Sukanya Wandhekar3, Ankit Chatorikar4 Vidya Pratishthan's Kamalnayan Bajaj Institute of Engineering and Technology, Baramati. [3] P. Chittora et al., "Prediction of Chronic Kidney Disease - A Machine Learning Perspective," in IEEE Access, vol. 9, pp. 17312-17334, 2021, Doi: 10.1109/ACCESS.2021.3053763. [4] Pal, S. Chronic Kidney Disease Prediction Using Machine Learning Techniques. Biomedical Materials & Devices (2022). https://doi.org/10.1007/s44174-022-00027-y [5] Schena, F.P., Anelli, V.W., Abbrescia, D.I. et al. Prediction of chronic kidney disease and its progression by artificial intelligence algorithms. J Nephrol (2022). https://doi.org/10.1007/s40620-022-01302-3

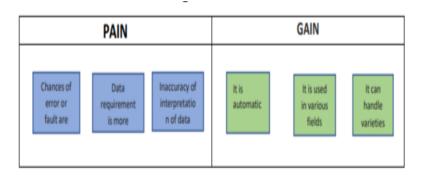
2.3 PROBLEM STATEMENT DEFINITION



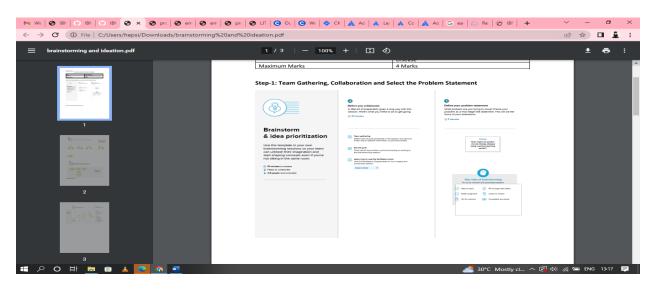
3. IDEATION & PROPOSED SOLUTION

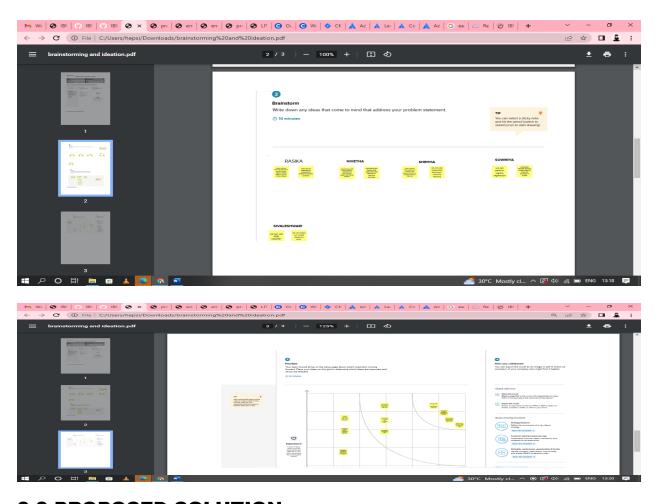
3.1 EMPATHY MAP CANVAS



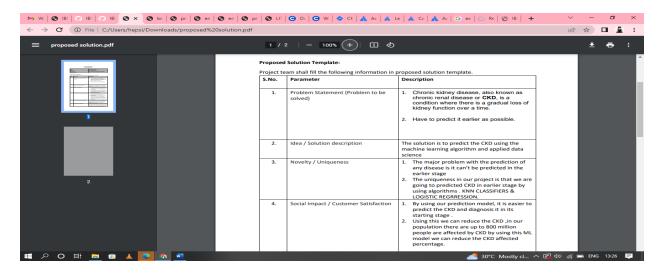


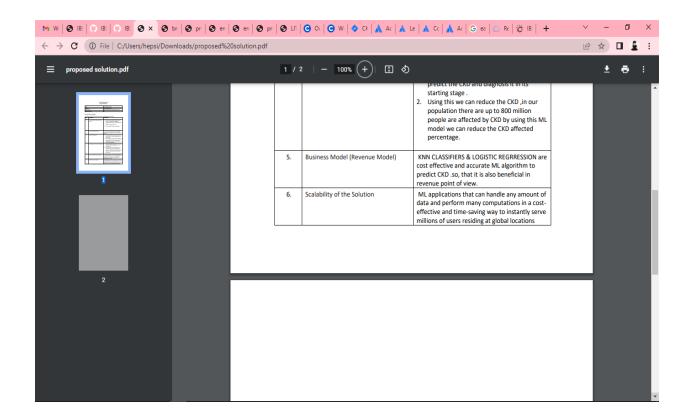
3.2 IDEATION & BRAINSTORMING



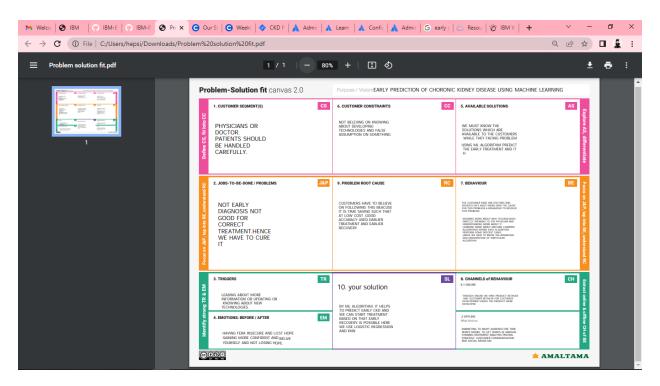


3.3 PROPOSED SOLUTION



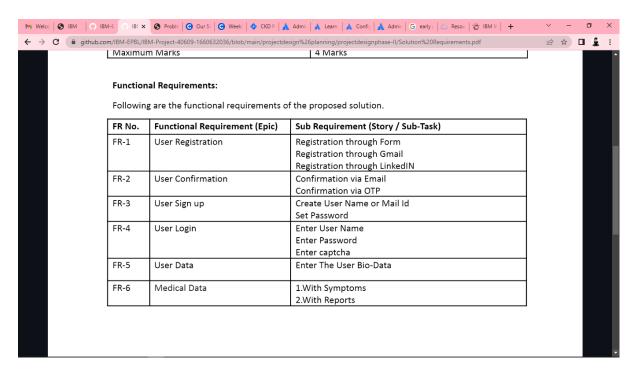


3.4 PROBLEM SOLUTION FIT

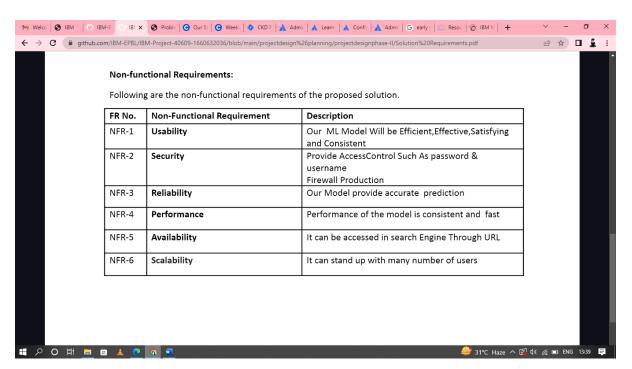


4.REQUIREMENT ANALYSIS

4.1 FUNCTIONAL REQUIREMENTS

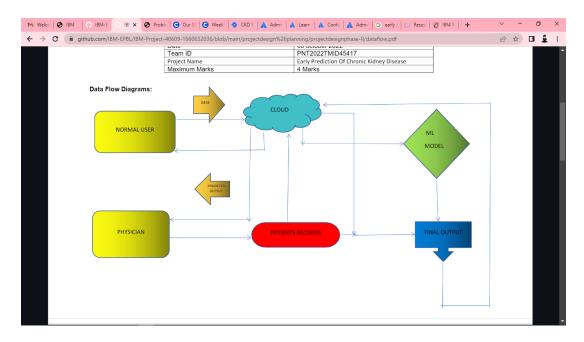


4.2 NON FUNCTIONAL REQUIREMENTS

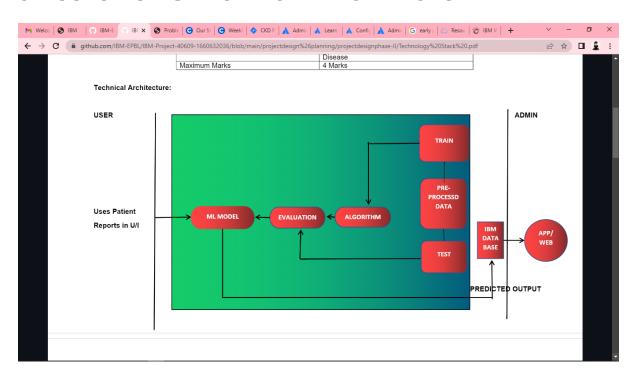


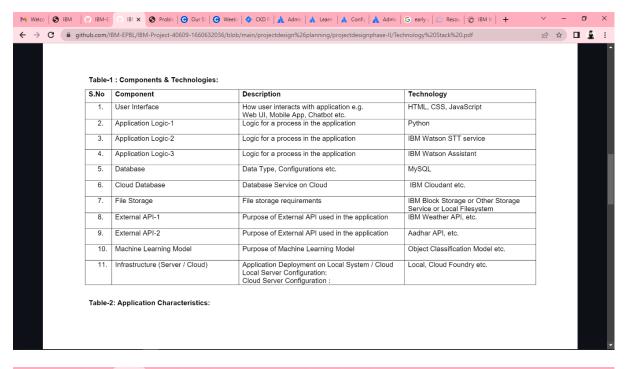
5. PROJECT DESIGN

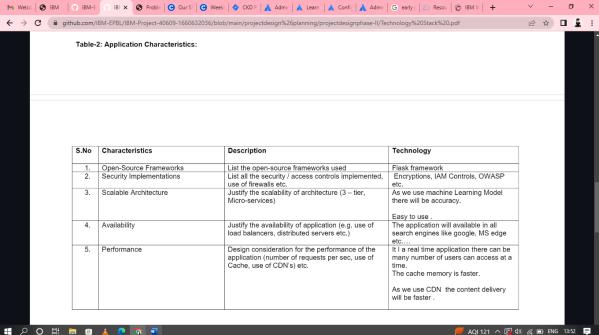
5.1 DATA FLOW DIAGRAM



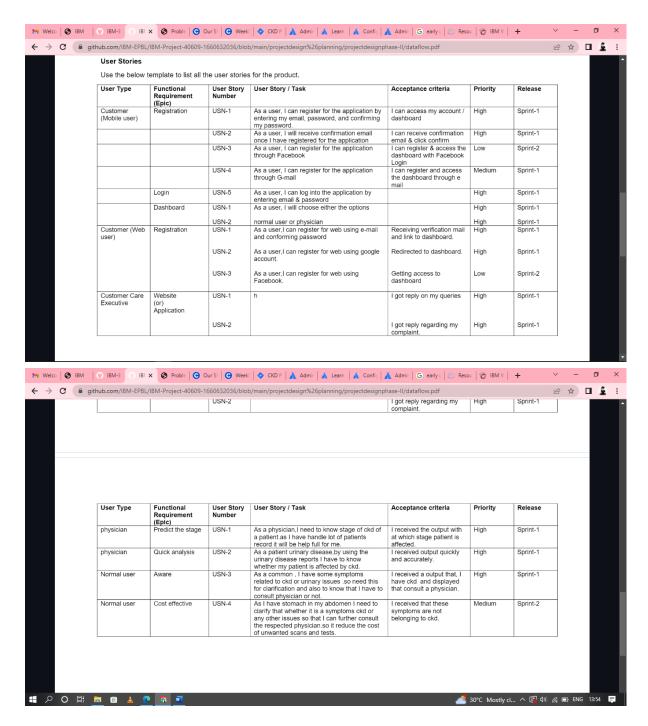
5.2 SOLUTION & TECHNICAL ARCHITECTURE





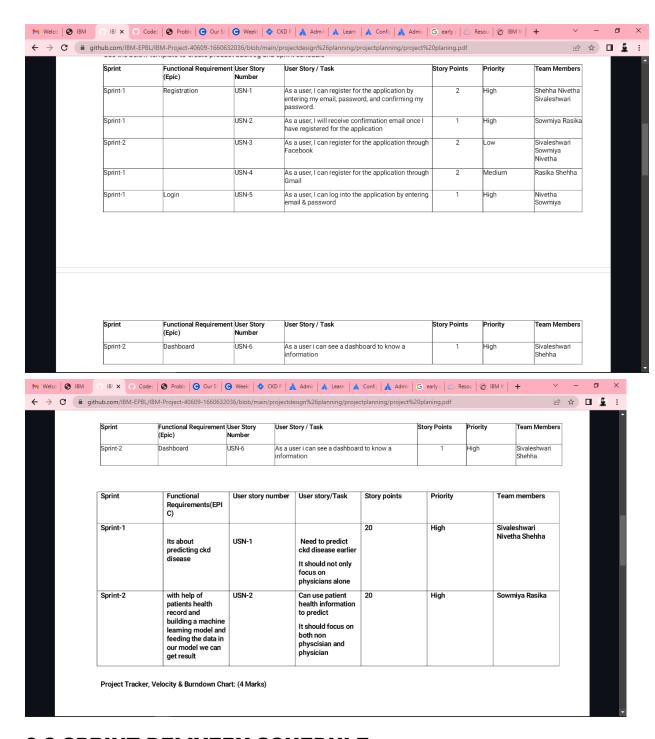


5.3 USER STORIES

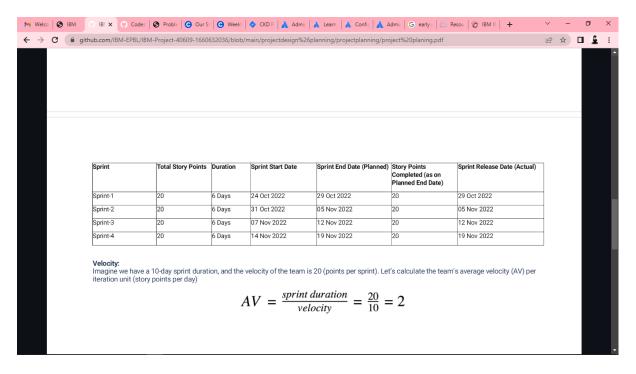


6. PROJECT PLANNING & SCHEDULING

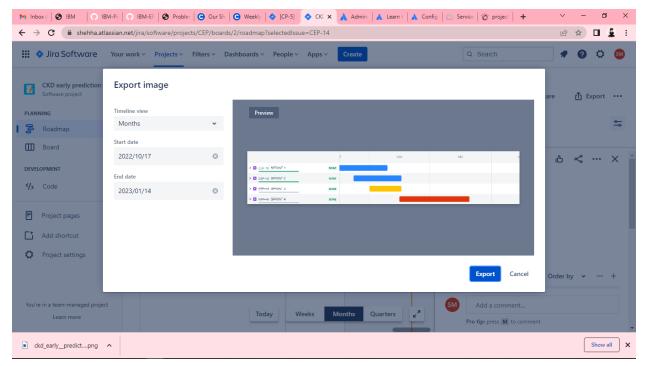
6.1 SPRINT PLANNING & ESTIMATION



6.2 SPRINT DELIVERY SCHEDULE



6.3 REPORTS FROM JIRA



7. CODING & SOLUTIONS

7.1 FEATURE 1

#importing libraries

```
import pandas as pd
import numpy as np
import matplotlib.pyplot as plt
import seaborn as sns
import os
os.getcwd()
path='C:\\Users\\ELCOT\\Downloads\\'
data=pd.read_csv(path+'chronickidneydisease.csv')
data.head(10)
data.tail(10)
data.shape
data.columns=['id','age','blood_pressure','specific_gravity','albumin','sugar','red_blood_cells',
       'pus_cell','pus_cell_clumps','bacteria','blood glucose random',
       'blood_urea','serum_creatinine','sodium','potassium','hemoglobin','packed_cell_volume',
       'white_blood_cell_count', 'red_blood_cell_count'
,'hypertension','diabetesmellitus','coronary_artery_disease',
       'apettite','pedal_edema','anemia','class']
data.columns
data.info()
data.drop(['id'],axis=1,inplace=True)
data
#target column
data['class'].unique()
#rectify target column
data['class']=data['class'].replace('ckd\t','ckd')
data['class'].unique()
#fetching categorical column
cat=data.select_dtypes(include=['object']).columns.tolist()
#removing column which are not categorical
cat.remove('red_blood_cell_count')
cat.remove('packed_cell_volume')
cat.remove( 'white_blood_cell_count')
cat
num=data.select_dtypes(include=['float64']).columns.tolist()#fetch numerical column
num.remove('specific_gravity')#remove which are not numerical
num.remove('albumin')
num.remove('sugar')
num
#adding column which is numerical
num.append('red_blood_cell_count')
```

```
num.append('packed_cell_volume')
num.append('white_blood_cell_count')
num
sns.pairplot(data,hue='class')
fig=plt.figure(figsize=(10,5))
fig
sns.barplot(x='blood glucose random',y='class',data=data)
cat.append('specific_gravity')#adding column which is categorical
cat.append('albumin')
cat.append('sugar')
cat
a=data['coronary_artery_disease'].unique()
#b=data['sugar'].unique()
#c=data['albumin'].unique()
#d=data['specific_gravity'].unique()
#e=data['anemia'].unique()
#f=data['pedal_edema'].unique()#
#g=data['apettite'].unique()
h=data['diabetesmellitus'].unique()
#i=data['bacteria'].unique()
#j=data['hypertension'].unique()#
#k=data['red_blood_cell_count'].unique()
#l=data['pus_cell'].unique()
#m=data['pus_cell_clumps'].unique()
a,h
#rectifying the categorical column classes
data['coronary_artery_disease']=data['coronary_artery_disease'].replace('\tno','no')
data['coronary_artery_disease'].unique()
data['diabetesmellitus']=data.diabetesmellitus.replace(to_replace={'yes':'yes','\tyes':'yes','\tno':'no'
})
data['diabetesmellitus'].unique()
#handling missing value
data.isna().sum()
#before handling the numeric variable which is considered as string should be convert to
numerical
data.red_blood_cell_count=pd.to_numeric(data.red_blood_cell_count,errors='coerce')
data.packed_cell_volume=pd.to_numeric(data.packed_cell_volume,errors='coerce')
data.white_blood_cell_count=pd.to_numeric(data.white_blood_cell_count,errors='coerce')
#handle numerical column null values
data['blood_pressure'].fillna(data['blood_pressure'].mean(),inplace=True)
data['blood_urea'].fillna(data['blood_urea'].mean(),inplace=True)
```

```
data['blood glucose random'].fillna(data['blood glucose random'].mean(),inplace=True)
data['serum_creatinine'].fillna(data['serum_creatinine'].mean(),inplace=True)
data['sodium'].fillna(data['sodium'].mean(),inplace=True)
data['potassium'].fillna(data['potassium'].mean(),inplace=True)
data['hemoglobin'].fillna(data['hemoglobin'].mean(),inplace=True)
data['pus_cell'].fillna(data['pus_cell'].mode()[0],inplace=True)
data['age'].fillna(data['age'].mode()[0],inplace=True)
data['pus_cell_clumps'].fillna(data['pus_cell_clumps'].mode()[0],inplace=True)
data['bacteria'].fillna(data['bacteria'].mode()[0],inplace=True)
data['red_blood_cell_count'].fillna(data['red_blood_cell_count'].mode()[0],inplace=True)
data['red_blood_cells'].fillna(data['red_blood_cells'].mode()[0],inplace=True)
data['white_blood_cell_count'].fillna(data['white_blood_cell_count'].mode()[0],inplace=True)
data['packed_cell_volume'].fillna(data['packed_cell_volume'].mode()[0],inplace=True)
data['hypertension'].fillna(data['hypertension'].mode()[0],inplace=True)
data['diabetesmellitus'].fillna(data['diabetesmellitus'].mode()[0],inplace=True)
data['coronary_artery_disease'].fillna(data['coronary_artery_disease'].mode()[0],inplace=True)
data['apettite'].fillna(data['apettite'].mode()[0],inplace=True)
data['pedal_edema'].fillna(data['pedal_edema'].mode()[0],inplace=True)
data['anemia'].fillna(data['anemia'].mode()[0],inplace=True)
data['specific_gravity'].fillna(data['specific_gravity'].mode()[0],inplace=True)
data['albumin'].fillna(data['albumin'].mode()[0],inplace=True)
data['sugar'].fillna(data['sugar'].mode()[0],inplace=True)
data.isnull().sum()
from sklearn.preprocessing import LabelEncoder
for i in cat:
  print('label of encoder= ',i)
  lei=LabelEncoder()
  print(data[i])
  data[i]=lei.fit_transform(data[i])
  print(data[i])
  print('*'*100)
data.corr().T
selcols=['red_blood_cells','pus_cell',
'diabetesmellitus', 'coronary_artery_disease','blood_urea','pedal_edema','anemia',
'blood glucose random']
x=pd.DataFrame(data,columns=selcols)
y=pd.DataFrame(data,columns=['class'])
print(x.shape)
print(y.shape)
from sklearn.model_selection import train_test_split
x_train,x_test,y_train,y_test=train_test_split(x,y,test_size=0.2,random_state=2)
```

```
print(x_train.shape)
print(y_train.shape)
print(x_test.shape)
print(y_test.shape)
from sklearn.linear_model import LogisticRegression
lgr=LogisticRegression()
lgr.fit(x_train,y_train)
y_pred=lgr.predict(x_test)
y_pred1=lgr.predict([[140,45,0,0,0,0,0,0]])
print(y_pred1)
from sklearn.metrics import accuracy_score
acc=accuracy_score(y_test,y_pred)
acc
from sklearn.metrics import confusion_matrix
conf_mat=confusion_matrix(y_test,y_pred)
conf_mat
import pickle
pickle.dump(lgr,open('CKD.pkl','wb'))
```

7.2 FEATURE 2

We import pandas for the analytics of the data set. Numpy library is used for the computation of array and seaborn for the visualization. We split the data in to train and test, and we logistic regression model to train the model and predict the ckd, by integrate the model in the web application developed in the flask.

8.TESTING

8.1 TEST CASES

We split the data in to train and test and train the model

from sklearn.model_selection import train_test_split

#SPLITING

```
x_train,x_test,y_train,y_test=train_test_split(x,y,test_size=0.2,random_state=2)
print(x_train.shape)
print(y_train.shape)
```

```
print(x_test.shape)
print(y_test.shape)
#IMPORT MODEL FOR TRAIN
from sklearn.linear_model import LogisticRegression
lgr=LogisticRegression()
lgr.fit(x_train,y_train)
#TEST
y_pred=lgr.predict(x_test)
y_pred1=lgr.predict([[140,45,0,0,0,0,0,0]])
print(y_pred1)
```

8.2 USER ACCEPTANCE TESTING

User Acceptance Testing (UAT), which is performed on most UIT projects, sometimes called beta testing or end-user testing, is a phase of software development in which the software is tested in the "real world" by the intended audience or business representative. This type of testing is not intended to be menu-driven, but rather to be performed by business users to verify that the application will meet the needs of the end-user, with scenarios and data representative of actual usage in the field. We make a web application to provide the better user interface.

9.RESULTS

9.1 PERFORMANCE METRICES

We get the performance accuracy of 91% for the prediction of our data set using logistic regression.

10. ADVANTAGES& DISADVANTAGES

ADVANTAGES:

- The website provide better user interface.
- It provide output with simple steps

- It is used to predict the disease quickly.
- It will be available for everyone.

DISADVANTAGES:

- Some times the prediction may be unreliable.
- When data is not accurate, the prediction will be wrong.

11. CONCLUSION:

Chronic Kidney Disease (CKD) is a serious condition that affects over 14% of the world's population. Because it can be predicted with a 91% overall accuracy, individuals may detect it early and receive treatment with the least amount of expense and risk. In practice, effective feature engineering minimizes the amount of characteristics required for the prediction algorithm and thus, the number of required medical tests. In comparison to similar work performed with the same dataset, filling in missing values based on their distribution and the cooccurrence of other characteristics using a Logistic Regression leads to improved accuracy in prediction models. Furthermore, compared to other models, the extratrees classifier and the random forest classifier are the superior algorithms for making predictions for CKD since they have 100% overall accuracy and less bias towards certain features. This project includes data preparation, treatment of missing information, and feature selection to predict the presence or absence of CKD. The necessity of adding domain expertise into feature selection while analyzing clinical data linked to CKD is further highlighted by this work. In light of this, it may be desirable to investigate the usage of a Logistic Regression based technique in the future to manage missing values in datasets relevant to various illnesses. Additionally, by including information about food kinds, water consumption habits, and genetics into the study, additional understanding of CKD may be acquired.

12. FUTURE SCOPE

The future scope of the project is very high. It will prevent the CKD in earlier and reduce the risk factors. As it can be used by normal people other than physician it give guidence to the people whether they to consult the respective physician or no need.

13.APPENDIX

SOURCE CODE:IBM

GITHUB: https://github.com/IBM-EPBL/IBM-Project-40609-1660632036

DEMO LINK

https://drive.google.com/file/d/1q5XwOlydh1vbQ3Joz0jGFkn1Wxpi5dKp/view?usp=sharing