IBM NALAIYA THIRAN – PROJECT REPORT EARLY DETECTION OF CHRONIC KIDNEY DISEASE USING MACHINE LEARNING

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1: INTRODUCTION

1.1 Project Overview

Early detection of chronic kidney disease using machine learning is to test the presence of kidney disease quickly and diagnose with appropriate medications. Since late detection of such diseases might lead to fatal reactions and medications might be expensive. Because to reduce such impact of kidney disease with detecting the disease in early stages. So that by utilizing ML algorithms, detection of kidney diseases can be done earlier and the risks due to the kidney diseases can also be reduced if it is detected earlier.

1.2 Purpose

Chronic kidney disease is A severe medical problem that should be treated earlier. Several medical tests taken for some purposes may also contain valuable information related to kidney disease. As a result the attributes of tests are investigated to distinguish which attribute may contain helpful information about the disease. Further on this information helps to measure the severity of the problem earlier and helps to make use of those information to build a machine learning model to predict the data. This research also considers the practical aspects of data collection and highlights the importance of incorporating domain knowledge when using machine learning for CKD status prediction.

2: LITERATURE SURVEY

2.1: Existing Problem

Chronic kidney disease, also called chronic kidney failure, involves a gradual loss of kidney function. Kidneys filter wastes and excess fluids from your blood, which are then removed in urine. Advanced chronic kidney disease can cause dangerous levels of fluid, electrolytes and wastes to build up in your body. Initially there are generally no symptoms; later, symptoms may include leg swelling, feeling tired, vomiting, loss of appetite, and confusion. In the early stages of chronic kidney disease, few signs or symptoms are noticeable which cannot make to realize that we have kidney disease until the condition is advanced.

2.2: References

S.No	Paper Title	Author	Journal	Publication	Description
			Name	Year	
1	Prediction of kidney disease-A machine learning perspective.	Pankaj Chittora	IEEE Access	2021	The paper explains the use of Machine Learning to act as a tool to detect diseases on time. Seven classifier algorithms have been applied in this research such as Artificial Neural Network, C5.0, Chisquare Automatic interaction detector, logistic regression, linear support vector machine with penalty L1 & with penalty L2 and random tree. The dataset used here consists of 400 instances and 24 attributes. The attributes are labelled in two classes as CKD (chronic kidney disease) and non-CKD. Out of all classifiers used, SVM was found to

					generate consistent
					results with higher
					accuracy.
2	XGBoost Model for chronic kidney disease	Adeola Ogunleye, Qing-Guo Wang	IEEE	2019	The paper explains the use of XGBoost Algorithm for detecting Chronic Kidney Disease. XGBoost is an extendible and cutting-edge application of gradient boosting machines and it has proven to push the limits of computing power for boosted trees algorithms. In this algorithm, decision trees are created in sequential form. It was developed for the sole purpose of model performance and computational speed. The model was found to starkly identify CKD apart from other disease with the attributes in the data set. The proposed model was applied and found to have achieved an accuracy, sensitivity and specificity of 1.000, 1.000
3	Early detection of kidney disease using advanced machine learning models.	A.Vaishnov i Anuhya, Ayyala Ganesh, Nallabathini Poojitha, Amandeep Singh, Amitha S K, Dr. Dhananjay a. V	International Research Journal of Engineering and Technology (IRJET)	2022	and 1.000, respectively. Every year, an increasing number of patients are diagnosed with late stages of renal disease. Chronic Kidney Disease, also known as Chronic Renal Disease, is characterized by abnormal kidney function or a breakdown of renal function that progresses over months or years. Chronic kidney disease is often found during screening of persons who are known to be at risk for

models are compared to select the optimal model for prediction. 4 Machine Zvi Segal, BMC 2017 End stage renal disease learning Dan Kalifa, algorithm for kira most severe stage of chronic kidney disease	Vector Machine (SVR), Random Forest (LR), Artificial Neural Network (ANN), and Decision Tree are four master teaching methodologies investigated (DT). The components are built using chronic kidney disease datasets, and the outcomes of these	The primary goals of this research are to design and suggest a machine learning method for predicting CKD. Support	4	learning algorithm for	Dan Kalifa, Kira	BMC	2017	The primary goals of this research are to design and suggest a machine learning method for predicting CKD. Support Vector Machine (SVR), Random Forest (LR), Artificial Neural Network (ANN), and Decision Tree are four master teaching methodologies investigated (DT). The components are built using chronic kidney disease datasets, and the outcomes of these models are compared to select the optimal model for prediction. End stage renal disease (ESRD) describes the most severe stage of
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		_		(0)(D)
	detection of	Bar		(CKD), when patients
1 .	end-stage	Ehrenberg,		need dialysis or renal
	renal disease	Guy Elad,		transplant. There is often
		Gal Maor,		a delay in recognizing,
		Maor Maor,		
				diagnosing, and treating
		Lewis,		the various etiologies of
		Muhamma		CKD. The objective of the
		d Tibi, Liat		present study was to
		Korn &		employ machine learning
		Gideon		algorithms to develop a
		Koren		prediction model for
		Roleii		•
				progression to ESRD
				based on a large-scale
				multidimensional
				database.
5	Chronic	G.	2021	Early diagnosis and
	kidney	Nandhini, J	· -	characterization are the
	•	•		
	disease	Aravinth		important components in
	prediction			determining the treatment
	using			of chronic kidney disease
	machine			(CKD). CKD is an ailment
	learning			which tends to damage
	techniques			the kidney and affect their
	tooriinquoo			effective functioning of
				•
				excreting waste and
				balancing body fluids.
				Some of the
				complications included
				are hypertension, anemia
				(low blood count), mineral
				bone disorder, poor
				•
				nutritional health, acid
				base abnormalities, and
				neurological
				complications. Early and
				error-free detection of
				CKD can be helpful in
				averting further
				9
				deterioration of patient's
				health. These chronic
				diseases are
				prognosticated using
				various types of data
				mining classification
				· ·
				approaches and machine
				learning (ML) algorithms.
1				This Prediction is

					performed using Random Forest (RF) Classifier, Logistic Regression (LR) and K-Nearest Neighbor (K-NN) algorithm and Support Vector Machine (SVM). The data used is collected from the UCI Repository with 400 data sets with 25 attributes. This data has been fed into Classification algorithms. The experimental results show that K-NN, LR, SVM hands out. Repository with 400 data sets with 25 attributes. This data has been fed into Classification algorithms. The experimental results show that K-NN, LR, The experimental results show that K-NN, LR,
6	Detection of Chronic Kidney Disease using Machine Learning Algorithms with Least Number of Predictors	Marwa Almasoud, Tomas E Ward	International Journal of Advanced Computer Science and Applications	2018	show that K-NN, LR, SVM hands out an accuracy of 94%, 98% and 93.75% respectively. The RF classifier gives out a maximum accuracy of 100%. Chronic kidney disease (CKD) is one of the most critical health problems due to its increasing prevalence. In this paper, we aim to test the ability of machine learning algorithms for the prediction of chronic kidney disease using the smallest subset of features. Several
					statistical tests have been done to remove redundant features such as the ANOVA test, the Pearson's correlation, and the Cramer's V test.

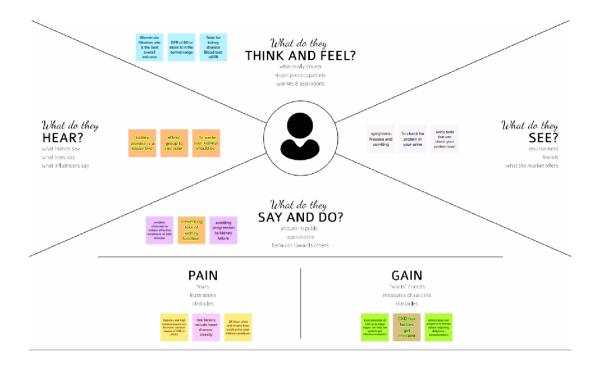
Logistic regression,
support vector machines,
random forest, and
gradient boosting.
earning algorithms for the
prediction of chronic
kidney disease using the
smallest subset of
features. Several
statistical tests have been
done to remove
redundant features such
as the ANOVA test, the
Pearson's correlation,
and the Cramer's V test.
Logistic regression,
support vector machines,
random forest, and
gradient boosting
algorithms have been
trained and tested using
10-fold cross-validation.
We achieve an accuracy
of 99.1 according to F1-
measure from Gradient
Boosting classifier. Also,
we found that hemoglobin
has higher importance for
both random forest and
Gradient boosting in
detecting CKD. Finally,
our results are among the
highest compared to
previous studies but with
a smaller number of
features reached so far.

2.3: Problem Statement Definition

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.

3: IDEATION AND PROPOSED SOLUTION

3.1: Empathy Map Canvas



3.2: Ideation and Brainstorming



Define your problem statement

What problem are you trying to solve? Frame your problem as a How Might We statement. This will be the focus of your brainstorm.

5 minutes

PROBLE

How might we predict the presence of Kidney Disease in a person



Brainstorm

Write down any ideas that come to mind that address your problem statement.

10 minutes

You can select a sticky note and hit the pencil [switch to sketch] icon to start drawing!

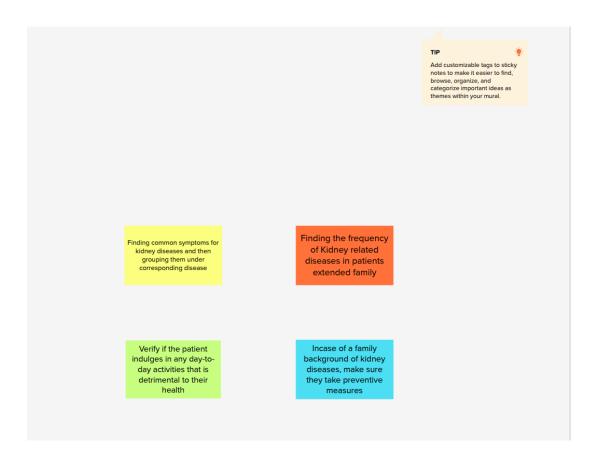
A ARCHANA G BHUVANESH S ABARNA SRI RAM R Tracking history of Kidney Finding if Finding the Tracing close contact with Analyzing **Analysing** Age of disease is extent of Workplace Diseases **Patient** other infected contagious Kidney Environment **Symptoms** patient amongst ancestors **Habits** people or not infection Finding Checking if Checking History of **Travel** others with personal and disease if relevant patient same age group having similar disease surrounding History fatal medication history hygiene symptoms



Group ideas

Take turns sharing your ideas while clustering similar or related notes as you go. Once all sticky notes have been grouped, give each cluster a sentence-like label. If a cluster is bigger than six sticky notes, try and see if you and break it up into smaller sub-groups.

0 20 minutes

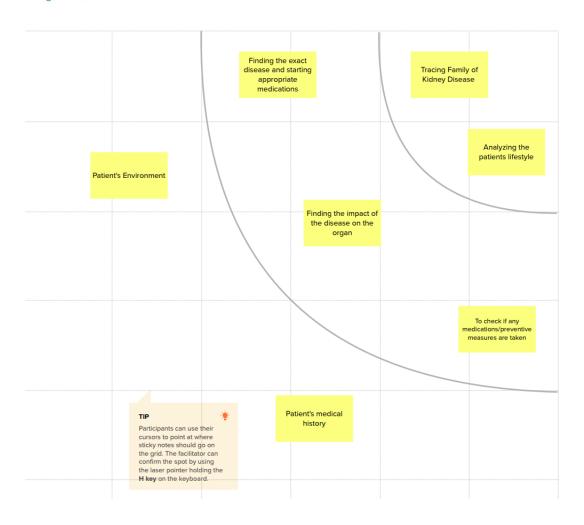




Prioritize

Your team should all be on the same page about what's important moving forward. Place your ideas on this grid to determine which ideas are important and which are feasible.

0 20 minutes



3.3: Proposed Solution

S.No.	Parameter	Description
1.	Problem Statement (Problem to be solved)	To develop a machine learning model and further, an application to detect chronic kidney disease in early stages
2.	Idea / Solution description	Utilizing the patient's health record data we propose a solution where a machine learning model is used to predict if patient is suffering from CKD or not.
3.	Novelty / Uniqueness	Early detection can be done using this solution. Easy to use for people who are not so familiar with applications.
4.	Social Impact / Customer Satisfaction	CKD can be detected at very early stages and the patient can get the necessary medical care at the earliest to get cured.
5.	Business Model (Revenue Model)	Apart from patients, this solution can be directly used at hospitals as well. So, the application can be purchased by hospitals or medical labs.
6.	Scalability of the Solution	The ML model can be scaled such that it can detect other diseases which are a side effect or can occur along with CKD.

3.4: Problem Solution Fit



4: REQUIREMENT ANALYSIS

4.1: Functional Requirements

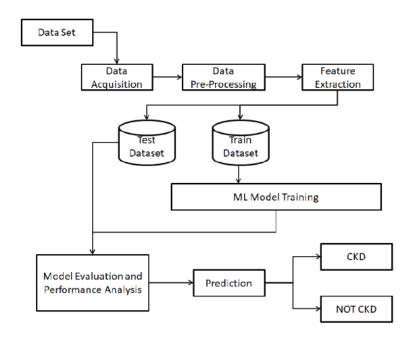
FR	Functional	Sub Requirement (Story / Sub-Task)
No.	Requirement (Epic)	
FR-1	User Registration	Registration through Google Form
		Registration through Hospitals
		Registration through Medical Camps
FR-2	User Confirmation	Confirmation via Message
		Confirmation via Mail
FR-3	User Requirements	Knowledge in application to submit required
		data
		Inferring the output and acting accordingly
FR-4	User Infrastructure	A system to support ML data modelling
		A Suitable GPU and CPU
FR-5	User Network	Network infrastructure to connect the User
		Interface and the Data Storage
FR-6	User Cost	No Expenditure will be necessary

4.2: Non – Functional Requirements

FR	Non-Functional	Description
No.	Requirement	
NFR-1	Usability	It can be used for various Datasets and
		trained for specific models and prediction
		methods.
NFR-2	Security	Secure Cloud Storage can be used to store
		data
NFR-3	Reliability	The Reliability of the predicted outcomes
		should be sufficient
NFR-4	Performance	The Computer's Hardware will play a major
		role in determining the performance of the
		System
NFR-5	Availability	It is available as a software package.
NED C	O I - I - iliu	It can be easiled on and interconnected with
NFR-6	Scalability	It can be scaled up and interconnected with
		other applications to provide various
		medical services.

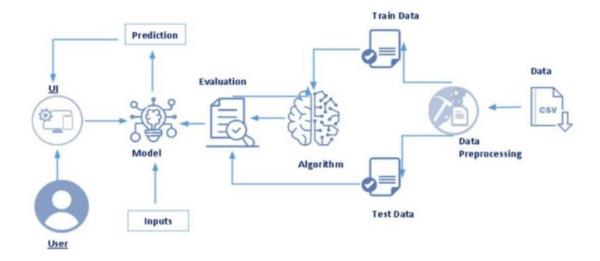
5: PROJECT DESIGN

5.1: Data Flow Diagrams



- 1. The Collected Data is pre-processes and is fed as input to the ML model.
- 2. Individual Feature Extraction is carried out before it is fed as input to the ML model.
- 3. The Dataset is divided into Test and Train Datasets.
- 4. The Model is trained using the Train Dataset.
- 5. The Model is then analysed using the Test Dataset and the predictions are carried out.
- 6. The error in prediction is calculated and used as feedback to train the model.
- 7. The model accuracy increases over the course.

5.2: Solution and Technical Architecture



5.3: User Stories

User Type	Functional Requirement (Epic)	User Story Number	User Story / Task	Acceptance criteria
Administrator	Data Collection	USN-1	As an Admin, I collect the appropriate dataset for predicting the chronic kidney disease.	I can submit images to the dataset under the guidelines and requirements mentioned
		USN-2	As an Admin, I split the dataset as train and test datasets.	From the Dataset images are split into Train and Test Data by an 80-20 ratio
	Model Building	USN-3	As an Admin, I split the Model into Training and Testing from the overall dataset.	I can feed the Separate Datasets to the ML Model for Testing and Training
Customer		USN-4	As a User, I submit my blood pressure and sugar level and other required information	I can view the required details that are to be used for prediction for specific individuals
Administrator	Training and Testing	USN-5	As an Admin, I Train the Model using	I can train the model for multiple epochs,

User Type	Functional Requirement (Epic)	User Story Number	User Story / Task	Acceptance criteria
			Regression algorithm and Test the Performance of the model.	improving efficiency of the prediction
	Implementation of the Application	USN-6	As an Admin, I work on predicting the spread of chronic kidney diseaseand predict the possibility of kidney failure	The Final System is deployed and the prediction is made
Customer		USN-7	As a User, I collect information and inferences from the system and analyse my lifestyle	I can view the out-come of the prediction and the analysis of the predicted outcome
Customer Care Executive		USN -8	As a Customer Care Executive, I shall attend the calls and guide the user.	I must clearly know the details of the model and the UI
	Maintenance	USN-9	I must be sure that the ML Model is up to date and is working in proper condition	I can check the outcome and update the model when required

6: PROJECT PLANNING AND SCHEDULING

6.1: Sprint Planning and Estimation

Sprint	Milestone
Sprint 1	User Registers into the application
	User inputs data of past illness, habits and other information relating to chronic kidney diseases
	3. User submits the details regarding Blood Sugar Levels, Blood
	Pressure and other required information relating to his current health conditions
Sprint 2	User can access the prediction model
	 After collecting the required information from the User, the ML model can now be accessed and used to predicted the presence of kidney diseases
Sprint 3	Application stores the predictions, that can be used for future analysis.
	2. The data stored has to be maintained securely.
Sprint 4	 Administrator should properly maintain the data and the prediction algorithm should be updated whenever required.

6.2 Sprint Delivery Schedule

Sprint	Total Story Points	Duration	Sprint Start Date	Sprint End Date (Planned)	Story Points Completed (as on Planned End Date)	Sprint Release Date (Actual)
Sprint-1	20	6 Days	24 Oct	29 Oct	20	29 Oct
			2022	2022		2022
Sprint-2	20	6 Days	31 Oct	05 Nov	20	04 Nov
			2022	2022		2022
Sprint-3	20	6 Days	07 Nov	12 Nov	20	09 Nov
			2022	2022		2022
Sprint-4	20	6 Days	14 Nov	19 Nov	20	16 Nov
			2022	2022		2022

6.3: Reports from JIRA



7. CODING AND SOLUTION

7.1 FEATURE 1 - DATA PREPROCESSING

Importing the necessary libraries for processing the dataset and building the model

```
import pandas as pd
import numpy as mp
from collections import Counter as c
import matplotlib.pyplot as plt
import seaborn as sns
import missingno as msno
from sklearn.metrics import accuracy_score, confusion_matrix
from sklearn.model_selection import train_test_split
from sklearn.preprocessing import LabelEncoder
from sklearn.linear_model import LogisticRegression
import pickle
```

The dataset is loaded into the jupyter notebook file.

```
data=pd.read_csv(r"C:\Users\archa\Desktop\IBM Datasets\chronickidneydisease.csv")
```

Information of the dataset is observed.

```
data.info()
<class 'pandas.core.frame.DataFrame'>
RangeIndex: 400 entries, 0 to 399
Data columns (total 25 columns):
# Column
                            Non-Null Count Dtype
                           391 non-null float64
0 age
1
    blood pressure
                            388 non-null
                                           float64
                           353 non-null
    specific_gravity
                                           float64
    albumin
                           354 non-null
                                          float64
                            351 non-null
                                           float64
    sugar
    red_blood_cells
                           248 non-null
                                           object
   pus cell
                           335 non-null
                                           object
    pus_cell_clumps
                           396 non-null
7
                                           object
8
    bacteria
                            396 non-null
                                           object
    blood glucose random 356 non-null
                                           float64
9
381 non-null serum_creatinine 383 non-null sodium
                                           float64
                                           float64
                                           float64
                           312 non-null
13 potassium
                                           float64
                           348 non-null
14 hemoglobin
                                           float64
15 packed_cell_volume
                            330 non-null
                                           object
16 white_blood_cell_count 295 non-null
                                           object
17 red_blood_cell_count 270 non-null
                                           object
19 diabetesmellitus
18 hypertension
                            398 non-null
                                           object
                            398 non-null
                                           object
                                           object
20 coronary_artery_disease 398 non-null
21 appetite
                  399 non-null
                                           object
22 pedal edema
                            399 non-null
                                           object
23 anemia
                           399 non-null
                                           object
24 class
                            400 non-null
                                           object
dtypes: float64(11), object(14)
memory usage: 78.2+ KB
```

The id column in the dataset is dropped and the columns are renamed for easy understanding.

The class, which is the output column is observed. The classes are changed so that there can be only two output – 'ckd' or 'notckd'.

Target Column

```
In [98]: data['class'].unique()
Out[98]: array(['ckd', 'ckd\t', 'notckd'], dtype=object)

Rectifying the target column

In [99]: data['class']=data['class'].replace("ckd\t","ckd")
data['class'].unique()
Out[99]: array(['ckd', 'notckd'], dtype=object)
```

The categorical columns are analysed. The fields which are not categorical are removed from the set 'catcols' and the fields(columns) which are categorical are added to the set.

```
In [100]: catcols=set(data.dtypes[data.dtypes=='0'].index.values)
    print(catcols)

    {'class', 'packed_cell_volume', 'red_blood_cells', 'pus_cell', 'appetite', 'bacteria', 'pedal_edema', 'hypertension', 'red_blood_cell_count', 'white_blood_cell_count', 'anemia', 'diabetesmellitus', 'coronary_artery_disease', 'pus_cell_clumps'}

In [101]: for i in catcols:
    print("Columns:",i)
    print((data[i]))
    print('**120+'\n')

catcols.add('specific_gravity')
    catcols.add('sloumin')
    catcols.add('sugar')
    print(catcols)

{'class', 'red_blood_cells', 'pus_cell', 'appetite', 'bacteria', 'albumin', 'pedal_edema', 'sugar', 'hypertension', 'anemia', 'specific_gravity', 'diabetesmellītus', 'coronary_artery_disease', 'pus_cell_clumps'}
```

Similarly, this is done for continuous columns. The continuous columns are analysed. The fields which are not continuous are removed from the set 'contcols' and the fields(columns) which are continuous are added to the set.

```
contcols=set(data.dtypes[data.dtypes!='0'].index.values)
print(contcols)

{'blood_urea', 'age', 'potassium', 'hemoglobin', 'sodium', 'sugar', 'albumin', 'blood glucose random', 'specific_gravity', 'ser
um_creatinine', 'blood_pressure'}

for i in contcols:
    print("Continuous Columns :",i)
    print(c(data[i]))
    print('*'*120+'\n')
```

Removing the Columns which are not Numerical

```
contcols.remove('specific_gravity')
contcols.remove('albumin')
contcols.remove('sugar')
print(contcols)
{'blood_urea', 'age', 'potassium', 'hemoglobin', 'sodium', 'blood glucose random', 'serum_creatinine', 'blood_pressure'}
```

Adding the Columns which are continuous

```
contcols.add('red_blood_cell_count')
contcols.add('packed_cell_volume')
contcols.add('white_blood_cell_count')
print(contcols)

{'blood_urea', 'packed_cell_volume', 'age', 'potassium', 'hemoglobin', 'sodium', 'blood glucose random', 'red_blood_cell_count', 'white_blood_cell_count', 'serum_creatinine', 'blood_pressure'}
```

The columns coronary artery disease and diabetes mellitus have ambiguous values. So this is rectified by either assigning the values to yes or no.

```
data['coronary_artery_disease']=data.coronary_artery_disease.replace('\tno','no')
c(data['coronary_artery_disease'])

Counter({'no': 364, 'yes': 34, nan: 2})

data['diabetesmellitus']=data.diabetesmellitus.replace(to_replace={'\tno':'no','\tyes':'yes','yes':'yes'})
c(data['diabetesmellitus'])

Counter({'yes': 136, 'no': 261, ' yes': 1, nan: 2})
```

The nest step is to verify if there are any null values present in the dataset.

<pre>data.isnull().any()</pre>		sodium	True
		potassium	True
age	True	hemoglobin	True
blood pressure	True	packed_cell_volume	True
specific gravity	True	white_blood_cell_count	True
albumin	True	red_blood_cell_count	True
sugar	True	hypertension	True
red blood cells	True	diabetesmellitus	True
pus cell	True	coronary_artery_disease	True
pus cell clumps	True	appetite	True
bacteria	True	pedal_edema	True
blood glucose random	True	anemia	True
blood urea	True	class	False
serum creatinine	True	dtype: bool	

The null values must be removed. But before that some continuous data are represented as a string. These are converted to numeric using to_numeric() function.

```
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')
data.red_blood_cell_count = pd.to_numeric(data.red_blood_cell_count, errors='coerce')
```

Now, the null values must be removed. This can be done by replacing the null values using either mean, median or mode using fillna() function.

```
data['blood glucose random'].fillna(data['blood glucose random'].mean(),inplace=True)
data['blood_pressure'].fillna(data['blood_pressure'].mean(),inplace=True)
data['blood_urea'].fillna(data['blood_urea'].mean(),inplace=True)
data['packed_cell_volume'].fillna(data['pecked_cell_volume'].mean(),inplace=True)
data['packed_cell_volume'].fillna(data['potassium'].mean(),inplace=True)
data['potassium'].fillna(data['potassium'].mean(),inplace=True)
data['red_blood_cell_count'].fillna(data['red_blood_cell_count'].mean(),inplace=True)
data['sodium'].fillna(data['sodium'].mean(),inplace=True)
data['sodium'].fillna(data['sodium'].mean(),inplace=True)
data['white_blood_cell_count'].fillna(data['white_blood_cell_count'].mean(),inplace=True)

data['age'].fillna(data['age'].mode()[0],inplace=True)
data['hypertension'].fillna(data['pus_cell_clumps'].mode()[0],inplace=True)
data['appetite'].fillna(data['appetite'].mode()[0],inplace=True)
data['appetite'].fillna(data['albumin'].mode()[0],inplace=True)
data['pus_cell'].fillna(data['pus_cell'].mode()[0],inplace=True)
data['red_blood_cells'].fillna(data['red_blood_cells'].mode()[0],inplace=True)
data['coronary_artery_disease'].fillna(data['coronary_artery_disease'].mode()[0],inplace=True)
data['anemia'].fillna(data['bacteria'].mode()[0],inplace=True)
data['anemia'].fillna(data['sugar'].mode()[0],inplace=True)
data['anemia'].fillna(data['sugar'].mode()[0],inplace=True)
data['diabetesmellitus'].fillna(data['pedal_edema'].mode()[0],inplace=True)
data['apdal_edema'].fillna(data['pedal_edema'].mode()[0],inplace=True)
data['specific_gravity'].fillna(data['specific_gravity'].mode()[0],inplace=True)
```

Both numerical and categorical values are present in the dataset. Only numerical can be processed by the computer. So, the categorical values must be encoded. This is done Label encoding in the sklearn library. Here, each of the integer is assigned a unique integer based on alphabetical ordering.

```
for i in catcols:
    print("Label Encoding of: ",i)
    LEi = LabelEncoder()
    print(c(data[i]))
    data[i]=LEi.fit_transform(data[i])
    print(c(data[i]))
    print("*"*100)
```

7.2 - MODEL BUILDING

In machine learning, the concept of the dependent variable (y) and independent variables(x) is important to understand. Here, the dependent variable is nothing but output in the dataset and the independent variable is all inputs in the dataset. We can denote any symbol (alphabets). In our dataset, we can say that class is the dependent variable and all other columns are independent. But in order to select the independent columns, we will be selecting only those columns which are highly correlated and some value to our dependent column.

```
selcols=['red_blood_cells','pus_cell','blood glucose random','blood_urea','pedal_edema','anemia','diabetesmellitus','coronary_art
x=pd.DataFrame(data,columns=selcols)
y=pd.DataFrame(data,columns=['class'])
print(x.shape)
print(y.shape)

4
(400, 8)
(400, 1)
```

The next step is to split the data set into training data and testing data. This is done using train_test_split. The train-test split is a technique for evaluating the performance of a machine learning algorithm.

Train Dataset: Used to fit the machine learning model.

Test Dataset: Used to evaluate the fit machine learning model.

About 80% of the data is used for training the machine learning model and 20% is used for testing.

```
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)

(320, 8)
(320, 1)
(80, 8)
(80, 1)
```

The dataset is trained using the Logistic Regression Model as it was found to have more accuracy over other models such as Decision Tree Classifier.

```
from sklearn.linear_model import LogisticRegression
lgr=LogisticRegression()
lgr.fit(x_train, y_train)

G:\Anaconda\lib\site-packages\sklearn\utils\validation.py:63: DataConversionWarning: A column-vector y was passed when a 1d arr
ay was expected. Please change the shape of y to (n_samples, ), for example using ravel().
    return f(*args, **kwargs)

LogisticRegression()
```

To the trained model, both the test input and an external input is given and the output is verified.

For the trained ML model. The accuracy and confusion matrix is plotted.

Finally, the model is saved as a pickle file.

```
pickle.dump(lgr,open('CKD.pkl','wb'))
```

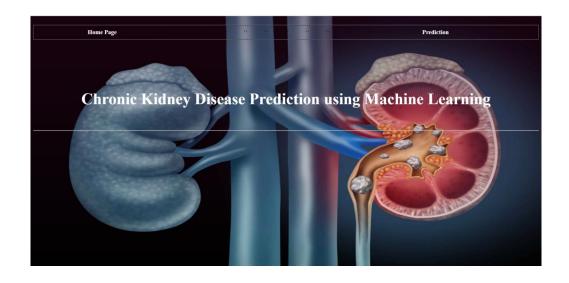
7.3 - Local Deployment

The HTML code for the home page, index page and the two prediction pages are written.

The app.py flask file which is a web framework written in python for server-side scripting is coded and run on jupyter notebook.

```
import numpy as np
import pandas as pd
from flask import Flask,request,render_template
import pickle as pk
app=Flask(__name__)
model=pk.load(open('CKD.pkl','rb'))
@app.route('/')
def home():
   return render_template('homepage.html')
p.route('/Prediction',methods=['POST','GET'])
def prediction():
"return render_template('indexpage.html')
@app.route('/Home',methods=['POST','GET'])
def my_home():
input_features=[float(x) for x in request.form.values()]
     features_value=[np.array(input_features)]
    features_name=['blood_urea','blood_glucose_random','coronary_artery_disease','anemia','pus_cell','red_blood_cells','diabetesmdf=pd.DataFrame(features_value,columns=features_name)
     output=model.predict(df)
    if(output==1):
         return render_template('predictionNo.html')
         return render_template('predictionYes.html')
                    main
if name
    app.run(debug=False)
```

The app.py runs on the local host: 5000 and the web page is viewed.



Enter your Blood year
Enter your Blood urea:
Enter your Blood Glucose Random:
Select Anemia or not : Yes ✓
Select Coronary Artery Disease or not : 🚾
Select Pus Cell Normal or Abnormal : Normal
Select Red Blood Cell Level Normal or Abnormal : Normal
Select Diabetes Mellitus or not : Y95 v
Select Pedal Enema or not : Yes v
Submit

Chronic Kidney Disease Prediction using Machine Learning

Prediction: You are not at risk of Chronic Kidney Disease! Keep up your good health!

FEATURE 7.4 - IBM CLOUD DEPLOYMENT

Once the local deployment is completed, the next step is to deploy the ML model from the cloud. Here, we utilise the IBM cloud platform. In the IBM cloud platform, Watson studio, Machine learning and Object Storage are used.

The same code that was used for training the model is loaded in the Watson studio. Few minor corrections are made in the code.

The pd.read_csv is replaced using a code which utilises API key to load the dataset stored in the IBM cloud space.

The ibm Watson machine learning library is installed.

```
In [50]: ! pip install -U ibm-watson-machine-learning
```

Using the unique API key generated in IBM Cloud and mentioning our server location. Using the API credentials a new space is created in IBM Watson. The space has its unique Space id. This space id is set as default.

The required ML model is downloaded. Then, we look for the version that is being supported by IBM and downloading the correct version. This is followed by creating a new deployment space for the model. The next step is to set up the model requirements and link it to the deployment space followed by saving the model to the space by mentioning the attributes of the model.

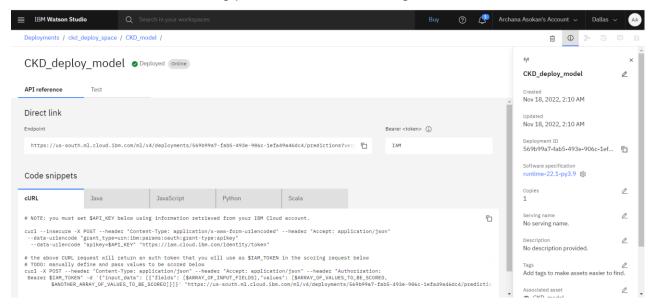
```
In [61]: model_name="CKD_model"
    deployment = "CKD_deploy_model"
model_deploy = lgr

In [62]: software_spec_uid = wml_client.software_specifications.get_uid_by_name("runtime-22.1-py3.9")
In [63]: software_spec_uid
Out[63]: '12b83a17-24d8-5082-900f-0ab31fbfd3cb'
In [64]: model_props = {
        wml_client.repository.ModelMetaNames.NAME:model_name,
        wml_client.repository.ModelMetaNames.TYPE:"scikit-learn_1.0",
        wml_client.repository.ModelMetaNames.SOFTWARE_SPEC_UID:software_spec_uid
    }
In [65]: model_details = wml_client.repository.store_model(
        model = model_deploy,
        meta_props = model_props,
        training_data = x_train,
        training_target = y_train)
```

The details of the saved model are generated. This includes the API key required for saving the model.

```
In [66]: model details
   Out[66]: {'entity': {'hybrid pipeline software specs': [],
                  'label_column': 'class',
'schemas': {'input': [{'fields': [{'name': 'red_blood_cells',
                        'type': 'int64'},
                       {'name': 'pus_cell', 'type': 'int64'},
                      {'name': 'blood glucose random', 'type': 'float64'}, 
{'name': 'blood_urea', 'type': 'float64'}, 
{'name': 'pedal_edema', 'type': 'int64'},
                       {'name': 'anemia', 'type': 'int64'},
                       {'name': 'diabetesmellitus', 'type': 'int64'},
                       {'name': 'coronary_artery_disease', 'type': 'int64'}],
                     'id': '1',
                     'type': 'struct'}],
                  'output': []},
'software_spec': {'id': '12b83a17-24d8-5082-900f-0ab31fbfd3cb',
                 'name': 'runtime-22.1-py3.9'},
'type': 'scikit-learn_1.0'},
'metadata': {'created_at': '2022-11-17T20:31:04.627Z',
                  'id': 'bc2b6428-307b-49b1-b087-cb8f5093c963',
                  'modified at': '2022-11-17T20:31:07.423Z',
                  'name': 'CKD model',
                  'owner': 'IBMid-6640045IHE',
                  'resource key': '2cf40232-410c-42e5-b99a-6b3b4acf99c5',
                  'space_id': '97486295-15b7-4879-81d2-eceeac8158b5'},
                 'system': {'warnings': []}}
```

Once the model has been created, saved, and deployed in the IBM cloud, we can see the below window. This consists of the Key required for deploying the model and also the header file and end scoring point which is to be integrated with the flask file.



INTEGRATING FLASK WITH SCORING END POINT:

The below code is used to link the code with the model in IBM cloud. This consists of the user specific API key.

```
import requests
import json
# NOTE: you must manually set API_KEY below using information retrieved from your IBM Cloud account.
API_KEY = "alcVu7bGrEpjyRSovDbNDF8tBZ89tU9aaQ3UjK-18Nbg"
token_response = requests.post('https://iam.cloud.ibm.com/identity/token', data={"apikey":
API_KEY, "grant_type": 'urn:ibm:params:oauth:grant-type:apikey'})
mltoken = token_response.json()["access_token"]
header = {'Content-Type': 'application/json', 'Authorization': 'Bearer ' + mltoken}
```

The necessary libraries are imported.

```
import numpy as np
import pandas as pd
from flask import Flask, request, render_template
import pickle as pk
```

The program given below serves as the backend for our Web page API and linking our Machine Learning model with it. We have a home page. From that you will be directed to the index page where you can give the inputs. This input received from that page is then sent to out ML model to do the prediction and the output will be displayed at the next web page, which can be either a prediction yes page (CKD positive) or prediction no page (CKD negative).

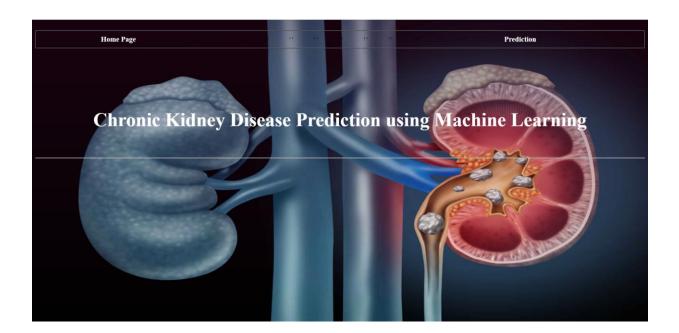
HTML PAGES:

Given below are the HTML codes for the four webpages and its corresponding result.

Home page:

```
k!DOCTYPE html>
<html lang="en">
 <head>
   <meta charset="utf-8" />
   <meta name="viewport" content="width=device-width, initial-scale=1" />
   <title>Chronic Kidney Disease Prediction using Maachine Learning</title>
  <body>
   <style>
     body {
       min-height: 100%;
       background: linear-gradient(
           ødeg,
           rgba(0, 0, 0, 0),
           rgba(78, 5, 48, 0.3)
         url(https://d3b6u46udi9ohd.cloudfront.net/wp-content/uploads/2022/05/24051731/Kidney-1.jpg);
       background-size: cover;
       background-repeat: no-repeat;
       background-attachment: fixed;
       background-size: 100% 100%;
   </style>
   <br />
   <table
           border="0"
           cellpadding="10"
           cellspacing="0"
          width="80%"
align="center"
           <a href="/Home" style="text-decoration: none"</pre>
                ><font
                  face="'Monaco'"
                  style="font-weight: bold"
                  size="4"
```

```
</font
              ></a>
             <a href="/Prediction" style="text-decoration: none"</pre>
                  face="'Monaco'"
                  style="font-weight: bold"
                  size="4"
                  color="white"
                  >Prediction</font
                ></a
             <br />
   <h1 align="center">
       <font face="Monaco" color="white" size="35"</pre>
        >Chronic Kidney Disease Prediction using Machine Learning</font
   </h1>
   <br /><br />
   <hr />
 </body>
</html>
```



Index page:

```
<!DOCTYPE html>
<html lang="en">
  <head>
   <meta charset="utf-8" />
<meta name="viewport" content="width=device-width, initial-scale=1" />
    <title>Chronic Kidney Disease Prediction using Machine Learning</title>
  <body>
   <style>
     body {
       min-height: 100%;
       background: linear-gradient(
            rgba(0, 0, 0, 0),
            rgba(173, 216, 230, 0.3)
          url();
       background-size: cover;
       background-repeat: no-repeat;
       background-attachment: fixed;
       background-size: 100% 100%;
    </style>
    <br />
    <h4 align="center">
       <font face="Monaco" color="darkblue" size="20"</pre>
          >Chronic Kidney Disease Prediction using Machine Learning</font>
       </b >
    </h4>
    </div>
   <br /><br />
    <form action="{{ url for('predict') }}" class="predict" method="POST"><br>
   <label for="bglucose">Enter your Blood Glucose Random:</label>
        <input type="number" id="bglucose" name="bglucose"><br><br>
        <label for="anemia">Select Anemia or not :</label>
        <select name="anemia" id="anemia">
        <option value="1">Yes</option>
<option value="0">No</option><br><br>
        <label for="Coronary">Select Coronary Artery Disease or not :</label>
<select name="Coronary" id="Coronary">
        <option value="1">Yes</option>
        <option value="0">No</option><br><br>
        <label for="cell">Select Pus Cell Normal or Abnormal :</label>
        <select name="cell" id="cell">
        <option value="1">Normal</option>
        <option value="0">Abnormal</option><br><br>
        </select><br/>//select><br/>//select Red Blood Cell Level Normal or Abnormal :</label>
        <select name="rbc" id="rbc">
        <option value="1">Normal</option>
        <option value="0">Abnormal</option><br><br>
        <label for="dia">Select Diabetes Mellitus or not :</label>
        <select name="dia" id="dia">
        <option value="1">Yes</option>
<option value="0">No</option><br><br>
        </select><br><br><
        <label for="enema">Select Pedal Enema or not :</label>
<select name="enema" id="enema">
        <option value="1">Yes</option>
        <option value="0">No</option><br><br>
        <button>Submit</button>
    </div>
    </form>
   khr />
  </body>
</html>
```

Enter your Blood urea:
Enter your Blood Glucose Random:
Select Anemia or not : ⋈₀₅ ✓
Select Coronary Artery Disease or not : [Yes v
Select Pus Cell Normal or Abnormal : Normal
Select Red Blood Cell Level Normal or Abnormal : Normal
Select Diabetes Mellitus or not : [Yes v]
Select Pedal Enema or not : Yes v
Submit

Chronic Kidney Disease Positive Page:

```
k!DOCTYPE html>
<html lang="en">
  <head>
    <meta charset="utf-8" />
<meta name="viewport" content="width=device-width, initial-scale=1" />
    <title>Chronic Kidney Disease Prediction using Machine Learning</title>
  </head>
  <body>
    <style>
      body {
        min-height: 100%;
        background: linear-gradient(
             0deg,
            rgba(0, 0, 0, 0),
rgba(255, 204, 203, 0.3)
          ),
          url();
        background-size: cover;
        background-repeat: no-repeat;
        background-attachment: fixed;
        background-size: 100% 100%;
    </style>
    <br />
    <div
      id="demobox"
      style="
        background-color: darkred;
        padding: 10px;
        border: 1px solid green;">
      <h4 style="color: white; text-align: center;">
      </h4>
    </div>
    <h4 align="center">
        <font face="Monaco" color="darkred" size="20"</pre>
          >Chronic Kidney Disease Prediction using Machine Learning</font
        ></b >
    </h4>
    </div>
    <br /><br />
    <div style="font-size: xx-large; text-align: center;">
        <br/> <b>Prediction: You are at risk of Chronic Kidney Disease! Please consult a doctor immediately!!</b>
    </div>
```

Chronic Kidney Disease Prediction using Machine Learning

Prediction: You are at risk of Chronic Kidney Disease! Please consult a doctor immediately!!

Chronic Kidney Disease Negative Page:

```
k!DOCTYPE html>
<html lang="en">
  <head>
    <meta charset="utf-8" />
    <meta name="viewport" content="width=device-width, initial-scale=1" />
    <title>Chronic Kidney Disease Prediction using Machine Learning</title>
  </head>
  <body>
    <style>
      body {
        min-height: 100%;
        background: linear-gradient(
            rgba(0, 0, 0, 0),
            rgba(144, 238, 144, 0.3)
          ),
          url();
        background-size: cover;
        background-repeat: no-repeat;
        background-attachment: fixed;
        background-size: 100% 100%;
    </style>
    <br />
    <div
     id="demobox"
      stvle="
        background-color: darkgreen;
        padding: 10px;
        border: 1px solid green;">
      <h4 style="color: white; text-align: center;">
      </h4>
    </div>
    <h4 align="center">
      <b>
        <font face="Monaco" color="darkgreen" size="20"</pre>
          >Chronic Kidney Disease Prediction using Machine Learning</font
        ></b >
    </h4>
    </div>
    <br /><br />
    <div style="font-size: xx-large; text-align: center;">
        <b>Prediction: You are not at risk of Chronic Kidney Disease! Keep up your good health!</b>
    </div>
```

Chronic Kidney Disease Prediction using Machine Learning

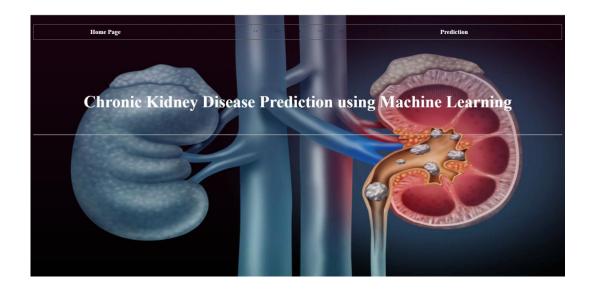
Prediction: You are not at risk of Chronic Kidney Disease! Keep up your good health!

When the flask file is run, the ML model that is saved in the IBM cloud is deployed. This results in the generation of the home page in the browser used. From the home page, we can navigate to and provide inputs in the index page. The ML model will then compute the result and display either one of the two prediction pages.

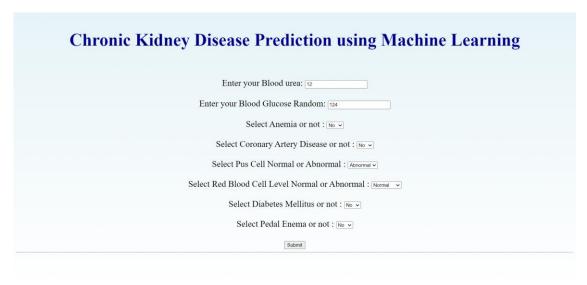
8. TESTING

8.1 TEST CASES

Test	Blood	Blood	Anemia	Coronary	Pus	RBC	Diabetes	Pedal	CKD
Case	Urea	Glucose		artery	Cell	Level	Mellitus	Enema	or
Number		Random		disease					Not
1	53	423	Yes	No	Abnormal	Abnormal	Yes	No	CKD
2	109	26	No	Yes	Normal	Normal	Yes	No	Not
									CKD
3	75	38	No	No	Normal	Normal	No	Yes	Not
									CKD
4	54	100	No	Yes	Normal	Abnormal	No	Yes	CKD
5	140	16	Yes	No	Normal	Abnormal	No	No	Not
									CKD

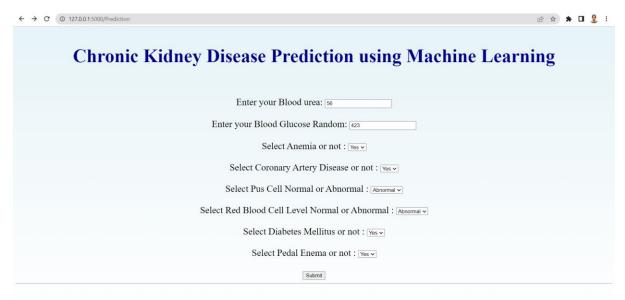


TEST CASE 1:



Chronic Kidney Disease Prediction using Machine Learning Prediction: You are not at risk of Chronic Kidney Disease! Keep up your good health!

TEST CASE 2:



Chronic Kidney Disease Prediction using Machine Learning Prediction: You are at risk of Chronic Kidney Disease! Please consult a doctor immediately!!

As observed, different inputs are given and the result is seen. Similarly, different values of inputs such as urea, blood sugar, anaemia, coronary artery disease etc are given and the result is observed and noted.

8.2 USER ACCEPTANCE TESTING

DEFECT ANALYSIS:

The number of resolved or closed bugs at each severity level, and how they were resolved

Resolution	Severity	Severity	Severity	Severity	Subtotal	
	1	2	3	4		
By Design	10	3	1	1	18	
Fixed	14	3	2	2	12	
Skipped	0	0	0	0	0	
Won't Fix	0	0	0	0	0	
Totals	24	6	3	3	30	

TEST CASE ANALYSIS:

This report shows the number of test cases that have passed, failed, and untested

Section	Total Cases	Not	Fail	Pass
		Tested		
Print Engine	1	0	0	13
	3			
Client Application	2	0	0	21
	1			
Security	2	0	0	2
Exception Reporting	1	0	0	1
Final Report Output	8	0	0	8
Version Control	1	0	0	1

9. RESULTS

9.1 PERFORMANCE METRICS

The Logistic Regression ML model that we have used here has better performance in accuracy compared to other models. We have compared the performance metrics of 2 models and selected this as the best for the application. The model performed well for all the test cases. The API developed also performed good with no glitches or lag found during the deployment and testing phase.

The accuracy score, confusion matrix and error values are attached below to support the above statement.

Accuracy = 91.25%

Mean absolute error = 0.0875 = 8.75%

```
from sklearn.metrics import mean_absolute_error
mean_absolute_error(y_test,y_pred)
0.0875
```

Classification Report:

```
from sklearn.metrics import classification report
print(classification_report(y_test, y_pred))
            precision recall f1-score support
                 1.00
                        0.87
                                   0.93
                0.79
                         1.00
                                  0.88
                                             26
   accuracy
                                   0.91
                                             80
  macro avg
                 0.89
                         0.94
                                   0.91
weighted avg
                0.93
                          0.91
                                   0.91
```

The ML model trained is found to have good accuracy with very minimal error. So, the developed ML model for Early Detection of Chronic Kidney Disease is highly accurate.

10. ADVANTAGES AND DIS-ADVANTAGES

ADVANTAGES:

Chronic kidney disease involves a gradual loss of kidney function. Advanced chronic kidney disease can cause dangerous levels of fluid, electrolytes and wastes to build up in your body. In the early stages of chronic kidney disease, you might have few signs or symptoms.

Detecting the Chronic kidney disease in the early stage has more advantage to take medications in the first stage

DIS-ADVANTAGE:

The limitations of early detection include the following: issues related to the sensitivity of currently available tests; the pre-test probability of disease in the population of interest; a lack of ability to accurately predict progression in individuals; and in some parts of the world the inability to offer therapy irrespective of diagnosis.

11. CONCLUSION

The early identification of CKD should increase the amount of time that patients are exposed to therapeutic strategies of proven benefit. Such increased exposure is expected to have several beneficial effects: preventing or delaying progression to ESRD; improving patient safety through avoidance of nephrotoxic effects of drugs and/or procedures (for example, contrast-based imaging); and, possibly, preventing AKI episodes in patients identified as having CKD.

In turn, these improvements should lead to a reduction in the need for RRT and improved health for these individuals. Given the already large burden of CKD on health-care resources, which is projected to increase still further in both developed and developing countries, improvements in CKD detection, an improved ability to predict individual patients' prognosis and better treatment strategies are necessary. As the incidence of diabetes increases in the developing world, and exposure to infections and drug therapies remain a constant threat to all, the impetus to ensure that identification of CKD occurs as early as possible must continue.

12. FUTURE WORKS

This project may help the future to detect the Chronic Kidney Disease early and to proceed with the treatments during the initial stage itself. Also, In Future, is model can also be improvised to give even more accuracy which helps to provide finite information.

13. APPENDIX

13.1 SOURCE CODE

DATA PRE-PROCESSING AND MODEL BUILDING CODE:

```
import pandas as pd
import numpy as mp
from collections import Counter as c
import matplotlib.pyplot as plt
import seaborn as sns
import missingno as msno
from sklearn.metrics import accuracy_score, confusion_matrix
from sklearn.model_selection import train_test_split
from sklearn.preprocessing import LabelEncoder
from sklearn.linear_model import LogisticRegression
import pickle
data=pd.read_csv(r"C:\Users\archa\Desktop\IBM Datasets\chronickidneydisease.csv")
data.head(5)
data.tail(5)
data.drop(["id"],axis=1,inplace=True)
data.columns
data.columns=['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',
```

'appetite', 'pedal_edema', 'anemia', 'class']

data.columns

```
data.info()
data['class'].unique()
data['class']=data['class'].replace("ckd\t","ckd")
data['class'].unique()
### Categorical Columns
catcols = set(data.dtypes[data.dtypes == 'O'].index.values) \\
print(catcols)
  print("Columns:",i)
  print(c(data[i]))
  print('*'*120+'\n')
catcols.remove('red_blood_cell_count')
catcols.remove('packed_cell_volume')
catcols.remove('white_blood_cell_count')
print(catcols)
# ## Numerical Columns
contcols=set(data.dtypes[data.dtypes!='O'].index.values)
print(contcols)
for i in contcols:
  print("Continuous Columns :",i)
  print(c(data[i]))
  print('*'*120+'\n')
contcols.remove('specific_gravity')
contcols.remove('albumin')
```

```
contcols.remove('sugar')
print(contcols)
contcols.add('red\_blood\_cell\_count')
contcols.add('packed_cell_volume')
contcols.add('white_blood_cell_count')
print(contcols)
catcols.add('specific_gravity')
catcols.add('albumin')
catcols.add('sugar')
print(catcols)
data['coronary_artery_disease']=data.coronary_artery_disease.replace('\tno','no')
c(data['coronary_artery_disease'])
data['diabetesmellitus']=data.diabetesmellitus.replace(to_replace={'\tno':'no',\tyes':'yes','yes':'
yes'})
c(data['diabetesmellitus'])
data.isnull().any()
data.isnull().sum()
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')
data.red_blood_cell_count = pd.to_numeric(data.red_blood_cell_count, errors='coerce')
data['blood glucose random'].fillna(data['blood glucose random'].mean(),inplace=True)
data['blood_pressure'].fillna(data['blood_pressure'].mean(),inplace=True)
data['blood_urea'].fillna(data['blood_urea'].mean(),inplace=True)
data['hemoglobin'].fillna(data['hemoglobin'].mean(),inplace=True)
```

```
data['packed_cell_volume'].fillna(data['packed_cell_volume'].mean(),inplace=True)
data['potassium'].fillna(data['potassium'].mean(),inplace=True)
data['red_blood_cell_count'].fillna(data['red_blood_cell_count'].mean(),inplace=True)
data['serum creatinine'].fillna(data['serum creatinine'].mean(),inplace=True)
data['sodium'].fillna(data['sodium'].mean(),inplace=True)
data['white_blood_cell_count'].fillna(data['white_blood_cell_count'].mean(),inplace=True)
data['age'].fillna(data['age'].mode()[0],inplace=True)
data['hypertension'].fillna(data['hypertension'].mode()[0],inplace=True)
data['pus_cell_clumps'].fillna(data['pus_cell_clumps'].mode()[0],inplace=True)
data['appetite'].fillna(data['appetite'].mode()[0],inplace=True)
data['albumin'].fillna(data['albumin'].mode()[0],inplace=True)
data['pus_cell'].fillna(data['pus_cell'].mode()[0],inplace=True)
data['red blood cells'].fillna(data['red blood cells'].mode()[0],inplace=True)
data['coronary_artery_disease'].fillna(data['coronary_artery_disease'].mode()[0],inplace=True
data['bacteria'].fillna(data['bacteria'].mode()[0],inplace=True)
data['anemia'].fillna(data['anemia'].mode()[0],inplace=True)
data['sugar'].fillna(data['sugar'].mode()[0],inplace=True)
data['diabetesmellitus'].fillna(data['diabetesmellitus'].mode()[0],inplace=True)
data['pedal_edema'].fillna(data['pedal_edema'].mode()[0],inplace=True)
data['specific_gravity'].fillna(data['specific_gravity'].mode()[0],inplace=True)
data.isnull().any()
for i in catcols:
  print("Label Encoding of: ",i)
```

```
LEi = LabelEncoder()
  print(c(data[i]))
  data[i]=LEi.fit_transform(data[i])
  print(c(data[i]))
  print("*"*100)
selcols=['red_blood_cells','pus_cell','blood glucose
random', 'blood_urea', 'pedal_edema', 'anemia', 'diabetesmellitus', 'coronary_artery_disease']
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([[129,99,1,0,0,1,0,1]])
print(y_pred)
print(c(y_pred))
print(y_pred1)
```

```
accuracy_score(y_test,y_pred)
confusion_mat = confusion_matrix(y_test,y_pred)
confusion_mat
pickle.dump(lgr,open('CKD.pkl','wb'))
```

TRAINING THE MODEL ON IBM:

import pandas as pd

import numpy as mp

from collections import Counter as c

import matplotlib.pyplot as plt

import seaborn as sns

from sklearn.metrics import accuracy_score, confusion_matrix

from sklearn.model_selection import train_test_split

from sklearn.preprocessing import LabelEncoder

from sklearn.linear_model import LogisticRegression

import pickle

import os, types

import pandas as pd

from botocore.client import Config

import ibm_boto3

def __iter__(self): return 0

@hidden_cell

```
# The following code accesses a file in your IBM Cloud Object Storage. It includes your
credentials.
# You might want to remove those credentials before you share the notebook.
cos_client = ibm_boto3.client(service_name='s3',
  ibm_api_key_id='8spio8rnxJxnzQCZqCY91nFWkd5pcolpt-ewKwuL3Ztj',
  ibm_auth_endpoint="https://iam.cloud.ibm.com/oidc/token",
  config=Config(signature_version='oauth'),
  endpoint_url='https://s3.private.us.cloud-object-storage.appdomain.cloud')
bucket = 'ckddetection-donotdelete-pr-yhimgpvl3j9jee'
object_key = 'chronickidneydisease.csv'
body = cos_client.get_object(Bucket=bucket,Key=object_key)['Body']
# add missing __iter__ method, so pandas accepts body as file-like object
if not hasattr(body, "__iter__"): body.__iter__ = types.MethodType( __iter__, body )
data = pd.read\_csv(body)
data.head()
data.head(5)
data.tail(5)
data.drop(["id"],axis=1,inplace=True)
data.columns
```

```
data.columns=['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',
    'appetite', 'pedal_edema', 'anemia', 'class']
data.columns
data.info()
data['class'].unique()
data['class']=data['class'].replace("ckd\t","ckd")
data['class'].unique()
### Categorical Columns
catcols=set(data.dtypes[data.dtypes=='O'].index.values)
print(catcols)
  print("Columns:",i)
  print(c(data[i]))
  print('*'*120+'\n')
catcols.remove('red_blood_cell_count')
catcols.remove('packed_cell_volume')
catcols.remove('white_blood_cell_count')
print(catcols)
```

```
### Numerical Columns
contcols=set(data.dtypes[data.dtypes!='O'].index.values)
print(contcols)
for i in contcols:
  print("Continuous Columns :",i)
  print(c(data[i]))
  print('*'*120+'\n')
contcols.remove('specific_gravity')
contcols.remove('albumin')
contcols.remove('sugar')
print(contcols)
contcols.add('red_blood_cell_count')
contcols.add('packed_cell_volume')
contcols.add('white_blood_cell_count')
print(contcols)
catcols.add('specific_gravity')
catcols.add('albumin')
catcols.add('sugar')
print(catcols)
data['coronary_artery_disease']=data.coronary_artery_disease.replace('\tno','no')
c(data['coronary_artery_disease'])
```

```
data['diabetesmellitus']=data.diabetesmellitus.replace(to_replace={'\tno':'no', \tyes':'yes', 'yes':'
yes'})
c(data['diabetesmellitus'])
data.isnull().any()
data.isnull().sum()
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')
data.red_blood_cell_count = pd.to_numeric(data.red_blood_cell_count, errors='coerce')
data['blood glucose random'].fillna(data['blood glucose random'].mean(),inplace=True)
data['blood_pressure'].fillna(data['blood_pressure'].mean(),inplace=True)
data['blood_urea'].fillna(data['blood_urea'].mean(),inplace=True)
data['hemoglobin'].fillna(data['hemoglobin'].mean(),inplace=True)
data['packed_cell_volume'].fillna(data['packed_cell_volume'].mean(),inplace=True)
data['potassium'].fillna(data['potassium'].mean(),inplace=True)
data['red blood cell count'].fillna(data['red blood cell count'].mean(),inplace=True)
data['serum_creatinine'].fillna(data['serum_creatinine'].mean(),inplace=True)
data['sodium'].fillna(data['sodium'].mean(),inplace=True)
data['white_blood_cell_count'].fillna(data['white_blood_cell_count'].mean(),inplace=True)
data['age'].fillna(data['age'].mode()[0],inplace=True)
data['hypertension'].fillna(data['hypertension'].mode()[0],inplace=True)
```

```
data['pus_cell_clumps'].fillna(data['pus_cell_clumps'].mode()[0],inplace=True)
data['appetite'].fillna(data['appetite'].mode()[0],inplace=True)
data['albumin'].fillna(data['albumin'].mode()[0],inplace=True)
data['pus_cell'].fillna(data['pus_cell'].mode()[0],inplace=True)
data['red_blood_cells'].fillna(data['red_blood_cells'].mode()[0],inplace=True)
data['coronary_artery_disease'].fillna(data['coronary_artery_disease'].mode()[0],inplace=True
)
data['bacteria'].fillna(data['bacteria'].mode()[0],inplace=True)
data['anemia'].fillna(data['anemia'].mode()[0],inplace=True)
data['sugar'].fillna(data['sugar'].mode()[0],inplace=True)
data['diabetesmellitus'].fillna(data['diabetesmellitus'].mode()[0],inplace=True)
data['pedal_edema'].fillna(data['pedal_edema'].mode()[0],inplace=True)
data['specific_gravity'].fillna(data['specific_gravity'].mode()[0],inplace=True)
data.isnull().any()
for i in catcols:
  print("Label Encoding of: ",i)
  LEi = LabelEncoder()
  print(c(data[i]))
  data[i]=LEi.fit_transform(data[i])
  print(c(data[i]))
  print("*"*100)
```

```
selcols=['red_blood_cells','pus_cell','blood glucose
random', 'blood_urea', 'pedal_edema', 'anemia', 'diabetesmellitus', 'coronary_artery_disease']
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([[129,99,1,0,0,1,0,1]])
print(y_pred)
print(c(y_pred))
print(y_pred1)
accuracy_score(y_test,y_pred)
confusion_mat = confusion_matrix(y_test,y_pred)
confusion_mat
get_ipython().system(' pip install -U ibm-watson-machine-learning')
from ibm_watson_machine_learning import APIClient
```

```
wml_credentials = {
  'apikey': "a1cVu7bGrEpjyRSOvDbNDF8tBZ89tU9aaQ3UjK-18Nbg",
  "url": "https://us-south.ml.cloud.ibm.com"
}
wml_client = APIClient(wml_credentials)
wml_client.spaces.list()
space_id = "97486295-15b7-4879-81d2-eceeac8158b5"
wml_client.set.default_space(space_id)
wml_client.software_specifications.list()
model_name="CKD_model"
deployment = "CKD_deploy_model"
model_deploy = lgr
software_spec_uid = wml_client.software_specifications.get_uid_by_name("runtime-22.1-
py3.9")
=software_spec_uid
=model_props = {
  wml_client.repository.ModelMetaNames.NAME:model_name,
wml_client.repository.ModelMetaNames.TYPE:"scikit-learn_1.0",
wml_client.repository.ModelMetaNames.SOFTWARE_SPEC_UID:software_spec_uid
}
model_details = wml_client.repository.store_model(
  model = model_deploy,
```

```
meta_props = model_props,
training_data = x_train,
training_target = y_train)
model_details
```

HOMEPAGE HTML CODE:

```
<!DOCTYPE html>
<html lang="en">
 <head>
  <meta charset="utf-8"/>
  <meta name="viewport" content="width=device-width, initial-scale=1" />
  <title>Chronic Kidney Disease Prediction using Maachine Learning</title>
 </head>
 <body>
  <style>
   body {
    min-height: 100%;
    background: linear-gradient(
       0deg,
       rgba(0, 0, 0, 0),
       rgba(78, 5, 48, 0.3)
      ),
      url(https://d3b6u46udi9ohd.cloudfront.net/wp-
content/uploads/2022/05/24051731/Kidney-1.jpg);
    background-size: cover;
    background-repeat: no-repeat;
```

```
background-attachment: fixed;
 background-size: 100% 100%;
 }
</style>
<br/>br />
>
  <table
   border="0"
   cellpadding="10"
   cellspacing="0"
   width="80%"
   align="center"
   >
     <a href="/Home" style="text-decoration: none"
      ><font
       face="'Monaco"
       style="font-weight: bold"
       size="4"
       color="white"
       >Home Page</font
```

```
></a
>
' '
' '
' '
' '
' '
' '
' '
' '
' '
' '
' '
' '
<a style="text-decoration: none"
 ><font
  face="'Monaco""
  style="font-weight: bold"
  size="4"
  color="white"
 >
 </font
></a>
```

```
<a href="/Prediction" style="text-decoration: none"
       ><font
        face="'Monaco""
        style="font-weight: bold"
        size="4"
        color="white"
        >Prediction</font
       ></a
     <br/>br />
<br/>br />
<br/>br />
<br />
<br/>br />
<br />
<br />
<h1 align="center">
```

INDEX PAGE HTML CODE

```
rgba(173, 216, 230, 0.3)
   ),
   url();
  background-size: cover;
  background-repeat: no-repeat;
  background-attachment: fixed;
  background-size: 100% 100%;
 }
</style>
<br/>br />
<h4 align="center">
 \langle b \rangle
  <font face="Monaco" color="darkblue" size="20"
   >Chronic Kidney Disease Prediction using Machine Learning</font>
  </b>
</h4>
</div>
<br /><br />
<form action="{{ url_for('predict') }}" class="predict" method="POST"><br>
<div style="font-size: x-large; text-align: center;">
  <label for="burea">Enter your Blood urea:</label>
  <input type="number" id="fname" name="fname"><br><br>
  <label for="bglucose">Enter your Blood Glucose Random:</label>
  <input type="number" id="bglucose" name="bglucose"><br><br>
  <label for="anemia">Select Anemia or not :</label>
```

```
<select name="anemia" id="anemia">
<option value="1">Yes</option>
<option value="0">No</option><br><br>
</select><br><br>
<label for="Coronary">Select Coronary Artery Disease or not :</label>
<select name="Coronary" id="Coronary">
<option value="1">Yes</option>
<option value="0">No</option><br><br>
</select><br><br>
<label for="cell">Select Pus Cell Normal or Abnormal :</label>
<select name="cell" id="cell">
<option value="1">Normal</option>
<option value="0">Abnormal</option><br><br>
</select><br><br>
<label for="rbc">Select Red Blood Cell Level Normal or Abnormal :</label>
<select name="rbc" id="rbc">
<option value="1">Normal</option>
<option value="0">Abnormal</option><br><br>>
</select><br><br>
<label for="dia">Select Diabetes Mellitus or not :</label>
<select name="dia" id="dia">
<option value="1">Yes</option>
<option value="0">No</option><br><br>
</select><br><br>
<label for="enema">Select Pedal Enema or not :</label>
```

PREDICTION YES PAGE HTML CODE

```
rgba(255, 204, 203, 0.3)
   ),
   url();
  background-size: cover;
  background-repeat: no-repeat;
  background-attachment: fixed;
  background-size: 100% 100%;
 }
</style>
<br/>br />
<div
 id="demobox"
 style="
  background-color: darkred;
  padding: 10px;
  border: 1px solid green;">
 <h4 style="color: white; text-align: center;">
 </h4>
</div>
<h4 align="center">
 <b>
  <font face="Monaco" color="darkred" size="20"</pre>
   >Chronic Kidney Disease Prediction using Machine Learning</font
  ></b>
</h4>
```

```
</div>
<br/>
<br/>
<br/>
<div style="font-size: xx-large; text-align: center;">
<br/>
<br/>
<br/>
<br/>
<br/>
/div>
</div>
</html>
```

PREDICTION NO PAGE HTML CODE

```
),
   url();
  background-size: cover;
  background-repeat: no-repeat;
  background-attachment: fixed;
  background-size: 100% 100%;
 }
</style>
<br/>br />
<div
 id="demobox"
 style="
  background-color: darkgreen;
  padding: 10px;
  border: 1px solid green;">
 <h4 style="color: white; text-align: center;">
 </h4>
</div>
<h4 align="center">
 <b>
  <font face="Monaco" color="darkgreen" size="20"
   >Chronic Kidney Disease Prediction using Machine Learning</font
  ></b>
</h4>
</div>
```

```
<br /><br />
  <div style="font-size: xx-large; text-align: center;">
    <br/> <b > Prediction: You are not at risk of Chronic Kidney Disease! Keep up your good
health!</b>
  </div>
  <hr />
 </body>
</html>
FLASK (app.py) CODE:
import requests
import json
# NOTE: you must manually set API_KEY below using information retrieved from your
IBM Cloud account.
API_KEY = "a1cVu7bGrEpjyRSOvDbNDF8tBZ89tU9aaQ3UjK-18Nbg"
token_response = requests.post('https://iam.cloud.ibm.com/identity/token', data={ "apikey":
API_KEY, "grant_type": 'urn:ibm:params:oauth:grant-type:apikey'})
mltoken = token_response.json()["access_token"]
header = {'Content-Type': 'application/json', 'Authorization': 'Bearer ' + mltoken}
import numpy as np
import pandas as pd
from flask import Flask,request,render_template
import pickle as pk
app=Flask(__name__)
model=pk.load(open('CKD.pkl','rb'))
@app.route('/')
```

```
def home():
  return render_template('homepage.html')
@app.route('/Prediction',methods=['POST','GET'])
def prediction():
              return render_template('indexpage.html')
@app.route('/Home',methods=['POST','GET'])
def my_home():
              return render_template('homepage.html')
@app.route('/predict',methods=['POST'])
def predict():
  payload_scoring = {"input_data": [{"field":
[['blood_urea','blood_glucose_random','coronary_artery_disease','anemia','pus_cell','red_bloo
d_cells','diabetesmellitus','pedal_edema']], "values": [input_features]}]}
  response_scoring = requests.post('https://us-
south.ml.cloud.ibm.com/ml/v4/deployments/569b99a7-fab5-493e-906c-
1efa49a46dc4/predictions?version=2022-11-17', json=payload_scoring,
  headers={'Authorization': 'Bearer ' + mltoken})
  print("Scoring response")
  predictions=response_scoring.json()
  pred=predictions['predictions'][0]['values'][0][0]
  if(pred==1):
    return render_template('predictionNo.html')
```

```
else:
    return render_template('predictionYes.html')

if __name__ == '__main__':
    app.run(debug=False)
```

13.2 GIT REPO LINK AND PROJECT DEMO LINK

Git Repo: https://github.com/IBM-EPBL/IBM-Project-42119-1660650072

Project demo video link: https://youtu.be/nDAlhi5cPxM