

**EARLY DETECTION OF CHRONIC KIDNEY DISEASE USING  
MACHINE LEARNING**

**TEAM ID: PNT2022TMID25581**

**IBM-PROJECT-53532-1661414687**

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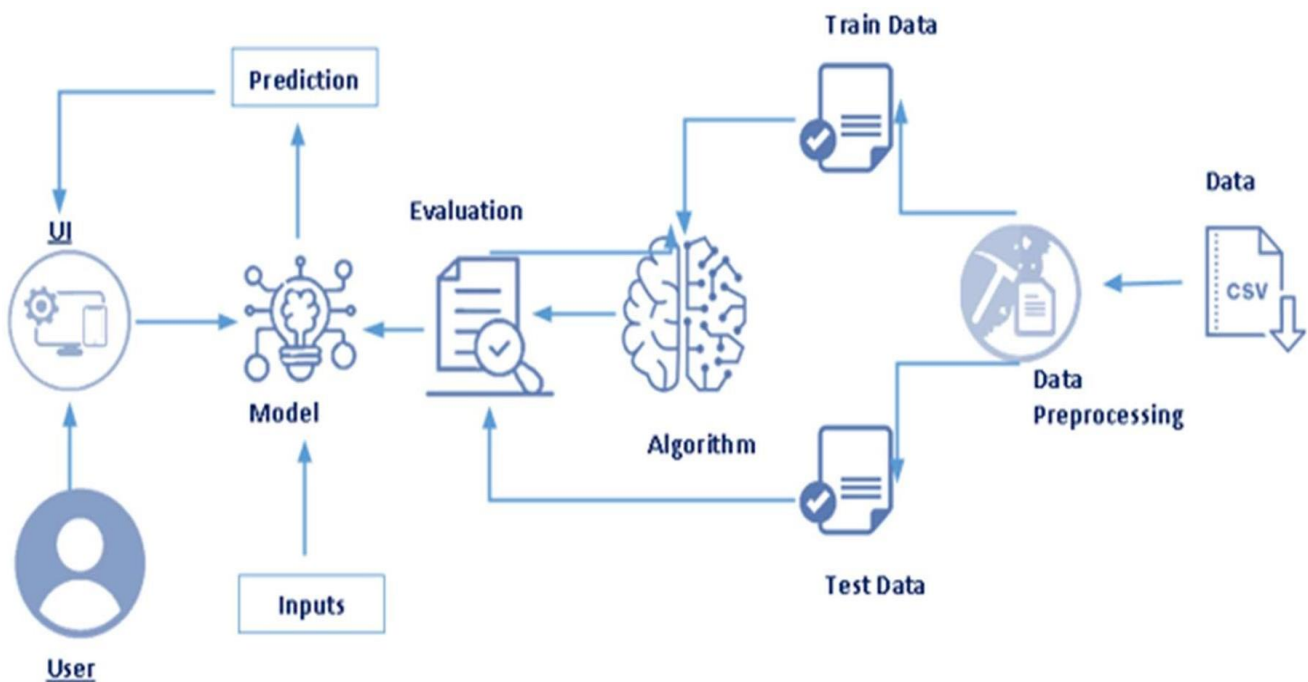
## 1.0 Introduction:

Chronic kidney disease prediction is one of the most important issues in health care-analytics. The most interesting and challenging tasks in day-to-day lives as one third of adult population is affected by chronic kidney disease (CKD), and millions die each year because they do not have access to affordable treatment. Chronic Kidney Disease can be cured, if treated in the early stages. The main aim of the project is to predict whether the patient have chronic kidney disease or not in a painless, accurate and faster way based on certain diagnostic measurement like Blood Pressure(BP), Albumin(AI) etc., and then appropriate treatment can be given based on the details provided by the model.

## 2.0 Skills Required:

Python, Python Web Frameworks, Python For data Visualization, Data Preprocessing Techniques, Machine Learning, IBM Cloud, IBM Watson Studio, Python-Flask

## 3.0 Architecture:



In this project we have followed the entire software development life cycle process using agile methodology.

## 4.0 LITERATURE SURVEY:

### INTRODUCTION:-

There are a large number of people in the world who gets affected to kidney related diseases. Our country India ranks first in terms of people getting affected by kidney related disease. Though population plays a major factor in it, the fact that China which has higher population than us is actually ranking in a much lower place than us in the term of people getting affected by kidney related disease. Generally, the kidney diseases are caused by the abnormal physiological functionalities of human kidney. Therefore, the characteristic symptoms are generated based on the differentiation between normal physiological functionalities and abnormal physiological functionalities of the kidney. The difference in the functionality of kidney is mainly due to the lifestyle and the food preference people have in their life. So the reason we were doing the project is to find the symptoms of kidney related diseases for the users and alert them before the disease becomes fatal for the patient. The main motive of this project is to find the symptom and its cure as soon as possible.

### Literature Review:-

[1]

DATE OF PUBLICATION	PAPER NAME	AUTHOR NAME	LINK
February 12, 2019	Detailed Review of Chronic Kidney Disease	Yesubabu Kakitapallia, Janakiram Ampolua, Satya Dinesh Madasa, M.L.S. Sai Kumar	<a href="https://www.karger.com/Article/Pdf/504622">https://www.karger.com/Article/Pdf/504622</a>

The proposed method is used to explain about the effects of kidney diseases. In this method Kidney damage can be assessed by albumin creatinine rate (ACR); albuminuria is one of the identifiers of kidney function in a timed urine collection. It was stated that one of the reasons for CKD, i.e., excretion of proteinuria, which is due to the intake of cooked meat or increased intake of protein or any kidney infection.

[2]

DATE OF PUBLICATION	PAPER NAME	AUTHOR NAME	LINK
October 1, 2019	Chronic Kidney Disease Diagnosis and Management	Teresa K. Chen, MD, MHS, Daphne H. Knicely, MD, and Morgan E Grams, MD, PhD	<a href="https://www.ncbi.nlm.nih.gov/pmc/articles/PMC7015670/">https://www.ncbi.nlm.nih.gov/pmc/articles/PMC7015670/</a>

The proposed system uses CKD method for diagnosis. In this method once adiagnosis of CKD has been made, the next step is to determine staging, whichis based on GFR, albuminuria, and cause of CKD . Staging of GFR is classifiedas G1 (GFR ≥90 ml/min/1.73 m²), G2 (GFR 60–89 ml/min/1.73 m²), G3a (45–59 ml/min/1.73 m²), G3b (30–44 ml/min/1.73 m²), G4 (15–29 ml/min/1.73 m²), and G5 (<15 ml/min/1.73 m²).

[3]

DATE OF PUBLICATION	PAPER NAME	AUTHOR NAME	LINK
27 November 2013	Risk factors for chronic kidney disease: an update	Rumeyza Kazancioğlu	<a href="https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4089662/">https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4089662/</a>

This system explains about inheritance in the kidney disease such as Uromodulin. It is another identified mutation is related to APOL1. An autosomal recessive pattern of inheritance is demonstrated and associated witha substantially higher risk of ESRD (10- fold higher risk of ESRD due to focal glomerulosclerosis and 7-fold higher risk of ESRD due to (hypertension)).

[4]

DATE OF PUBLICATION	PAPER NAME	AUTHOR NAME	LINK
31 December2014	Systems biology towards novel chronic kidney disease diagnosis and treatment.	Dr. Bernd Mayer	<a href="https://cordis.europa.eu/docs/results/241/241544/final1-syskid-final-report-2015-03-13.pdf">https://cordis.europa.eu/docs/results/241/241544/final1-syskid-final-report-2015-03-13.pdf</a>

This system explains about the hemodynamic in our glomerulor. Alterations in glomerular hemodynamic were considered of upmost importance. Both afferent arteriolar glomerular vasodilatation and efferent vasoconstriction increase intra-glomerular filtration pressure thus leading to hyper filtration, which on the short term stabilizes GFR but on the long term leads to progressing glomerular sclerosis thereby initiating a vicious cycle

[5]

DATE OF PUBLICATION	PAPER NAME	AUTHOR NAME	LINK
June 2020	Chronic kidney disease: prevalence and risk factors.	A.M. Aringazina, O.Zh.Narmanova, G.O. Nuskabaeva, Zh.A. Tagaeva	<a href="http://www.researchgate.net/publication/342930212_Chronic_kidney_disease_prevalence_and_risk_factors_literature_review">www.researchgate.net/publication/342930212_Chronic_kidney_disease_prevalence_and_risk_factors_literature_review</a>

This system explains about the disease risks and affects that has on us and our surrounding. It also explains about the ill factor it might have and chances of spreading to anyone in our surrounding.

[6]

DATE OF PUBLICATION	PAPER NAME	AUTHOR NAME	LINK
July 2011	The Burden of Chronic Kidney Disease on Developing Nations: A 21st Century Challenge in Global Health	Nugent R.A, Fathima S.F., Feigl A.B, Chyung D	<a href="https://www.karger.com/Article/Fulltext/321382">https://www.karger.com/Article/Fulltext/321382</a>

The proposed system explains about the ill effects these kidney diseases have on developing countries. In developing nations, the growing prevalence of chronic diseases such as chronic kidney disease has severe implications on health and economic output. The rapid rise of common risk factors such as diabetes, hypertension, and obesity, especially among the poor, will result in even greater and more profound burdens that developing nations are not equipped to handle.

[7]

DATE OF PUBLICATION	PAPER NAME	AUTHOR NAME	LINK
FEBRAURY ,2020	A Comparative Analysis of Machine Learning Techniques	Alvaro Sobrinho, Andessa C.M. Da Silveira, Leandro Dias da Silva, Evandro de B. Costa	<a href="https://www.researchgate.net/publication/339014686_ComputerAided_Diagnosis_of_Chronic_Kidney_Disease_in_Developing_Countries_A_Comparative_Analysis_of_Machine_Learning_Techniques">https://www.researchgate.net/publication/339014686_ComputerAided_Diagnosis_of_Chronic_Kidney_Disease_in_Developing_Countries_A_Comparative_Analysis_of_Machine_Learning_Techniques</a>

This system explains about the use of diagnosing with the help of computer. Software systems have been developed to assist physicians during CKD monitoring and diagnosis. For example, CKD-Go is a web application (app) to help users verify their kidney function by inputting their ACR and GF

[8]

DATE OF PUBLICATION	PAPERNAME	AUTHORNAME	LINK
January 2018	Diagnosis of Kidney Conditions Using Low-Cost Paper Diagnostics	Md. Nazibul Islam	<a href="https://www.researchgate.net/publication/325499924_Diagnosis_of_Kidney_Conditions_Using_Low-Cost_Paper_Diagnostics">https://www.researchgate.net/publication/325499924_Diagnosis_of_Kidney_Conditions_Using_Low-Cost_Paper_Diagnostics</a>

This proposed system uses Low cost paper diagnosis method. The method Paper diagnostic devices (PADs) can play a vital role in low-cost and rapid diagnosis of kidney condition, resulting in early detection of kidney complications. Paper diagnostics are paper and cellulose based analytical devices capable of qualitative or quantitative detection of biomarker.

[9]

Date OF PUBLICATION	PAPERNAME	AUTHOR NAME	LINK
December 2019	Machine Learning Applied to Kidney Disease Prediction: Comparison Study	Akm Shahariar Azad Rabby, Rezwana Mamata, Monira Akter Laboni, Ohidujjaman	<a href="https://www.researchgate.net/publication/338356158_Machine_Learning_Applied_to_Kidney_Disease_Prediction_Comparison_Study">https://www.researchgate.net/publication/338356158_Machine_Learning_Applied_to_Kidney_Disease_Prediction_Comparison_Study</a>

The proposed system uses applied machine language for early detection of kidney disease. The main aim is to find an optimized and efficient machine learning (ML) technique that can effectively recognize and predict the condition of chronic kidney disease. the data has been divided into two sections. In one section traindataset got trained and another section got evaluated by test dataset. The analysis results show that the Decision Tree Classifier and Gaussian Naive Bayes achieved the highest performance than the other classifiers, obtaining the accuracy score of 100% and 1 recall(Sensitivity) score.



<b>Date OF PUBLICATION</b>	<b>PAPER NAME</b>	<b>AUTHORNAME</b>	<b>LINK</b>
January 2022	Development of a Graphical User Interface Software for The Prediction of Chronic Kidney Disease	S.C. Nwaneri and H.C. Ugo	<a href="https://www.ajol.info/index.php/njt/article/view/225270/212516">https://www.ajol.info/index.php/njt/article/view/225270/212516</a>

The proposed system explains about the training and development of model for kidney diagnosing. This process involves building the neural network architecture and determining the activation functions to be used at the output of each layer of the network. The ANN model is built using the Keras libraries in python. Dense is used to assign the number of layers for the network. The ANN model consists of 25 neurons in the input layer and one neuron at the output layer.

As per agile methodology there are various phases in our project.

1. Project Design and Planning Phase
2. Project Development Phase
3. Train the Machine learning model on IBM.

## **5.0 Project Design and Planning Phase:**

In the Project Design and planning phase we have

5.0 Ideation phase

5.1 Design Phase I

5.2 Design phase II

5.3 Project planning phase

### **5.1 IDEATION PHASE**

During the ideation phase we have done problem definition, Empathy map canvas and Brainstorming and ideation process. Let us see in detail each activity.

### **5.1.1 PROBLEM DEFINITION**

**Team ID** : PNT2022TMID25581

**PROJECT NAME** : EARLY DETECTION OF CHRONIC KIDNEY  
DISEASE

# PROBLEM STATEMENT

## EARLY DETECTION OF CHRONIC KIDNEY DISEASE USING MACHINE LEARNING

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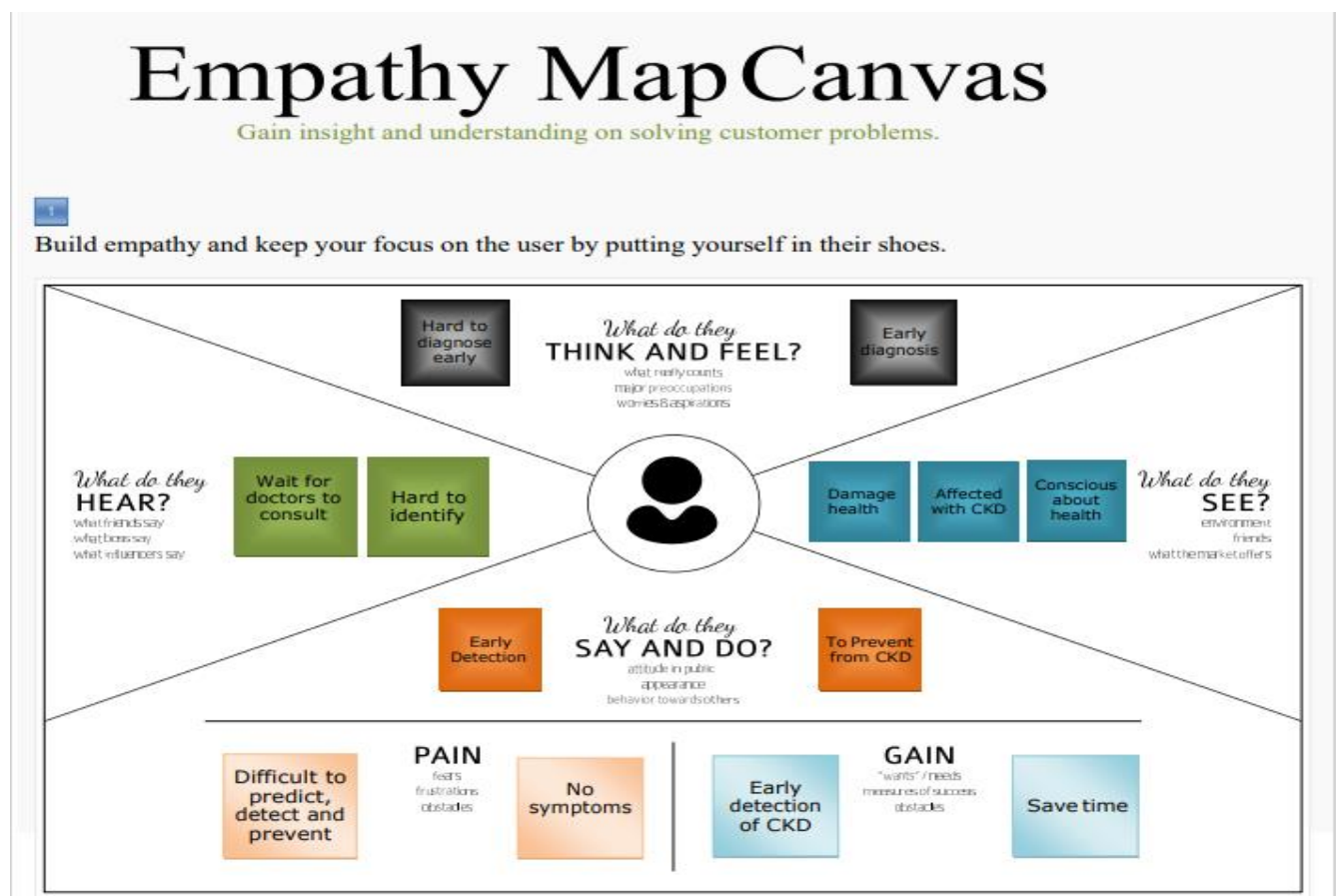
- Irregular lifestyle of current days leads to many disease chronic diseases. One of them being Chronic Kidney Disease (CKD), it has been of a growing concern, kidney is one of the most important organs in the body required for filtering blood, once a person has lost their kidneys, they could survive only for 18 days without their kidneys, it would take a fortune to just keep the person alive, with treatments like dialysis, transplant etc.
- 10% of the population worldwide is affected by chronic kidney disease (CKD), and millions die each year because they do not have access to affordable treatment.
- People usually don't realize that the medical tests we perform for various purposes can contain valuable information related to kidney disease. Subsequently, the attributes of various medical tests are examined to distinguish which attributes may contain useful information about the disease. The information, they say, helps us gauge the severity of the problem, and we use that information to build a machine learning model that predicts chronic kidney disease.
- Early detection of kidney disease can help in treatment which could save lives. Analyzing various medical tests, would give us an idea about which attributes help us distinguish the disease.
- The main aim of this project is to predict whether the patient have chronic kidney disease or not, in more accurate and faster way based on certain diagnostic measurements like Blood Pressure (BP), Albumin (AI).

## 5.1.2 EMPATHY MAP CANVAS

An empathy map is a collaborative visualization used to articulate what we know about a particular type of user. It externalizes knowledge about users in order to

- 1) Create a shared understanding of user needs, and
- 2) Aid in decision making.

An empathy map helps to map what a design team knows about the potential audience. This tool helps to understand the reason behind some actions a user takes deeply. This tool helps build Empathy towards users and helps design teams shift focus from the product to the users who are going to use the product.



### **5.1.3 SOLUTION ARCHITECTURE**

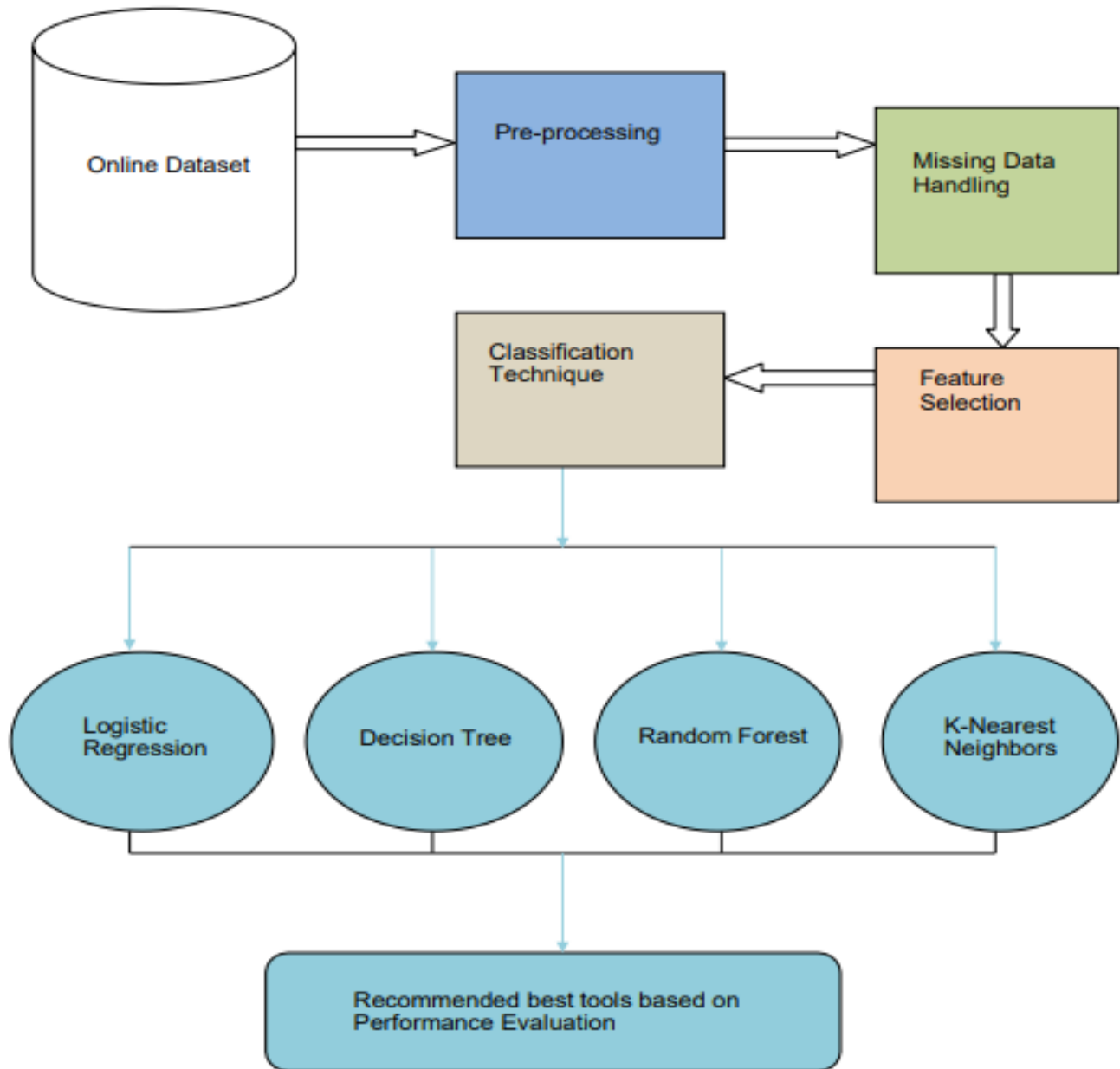
Solution Architecture:

Solution architecture is a complex process with many sub processes that bridges the gap between business problems and technology solutions.

Its goals are to:

- Find the best tech solution to solve existing business problems.
- Describe the structure, characteristics, behavior, and other aspects of the software to project stakeholders.
- Define features, development phases, and solution requirements.
- Provide specifications according to which the solution is defined, managed, and delivered.

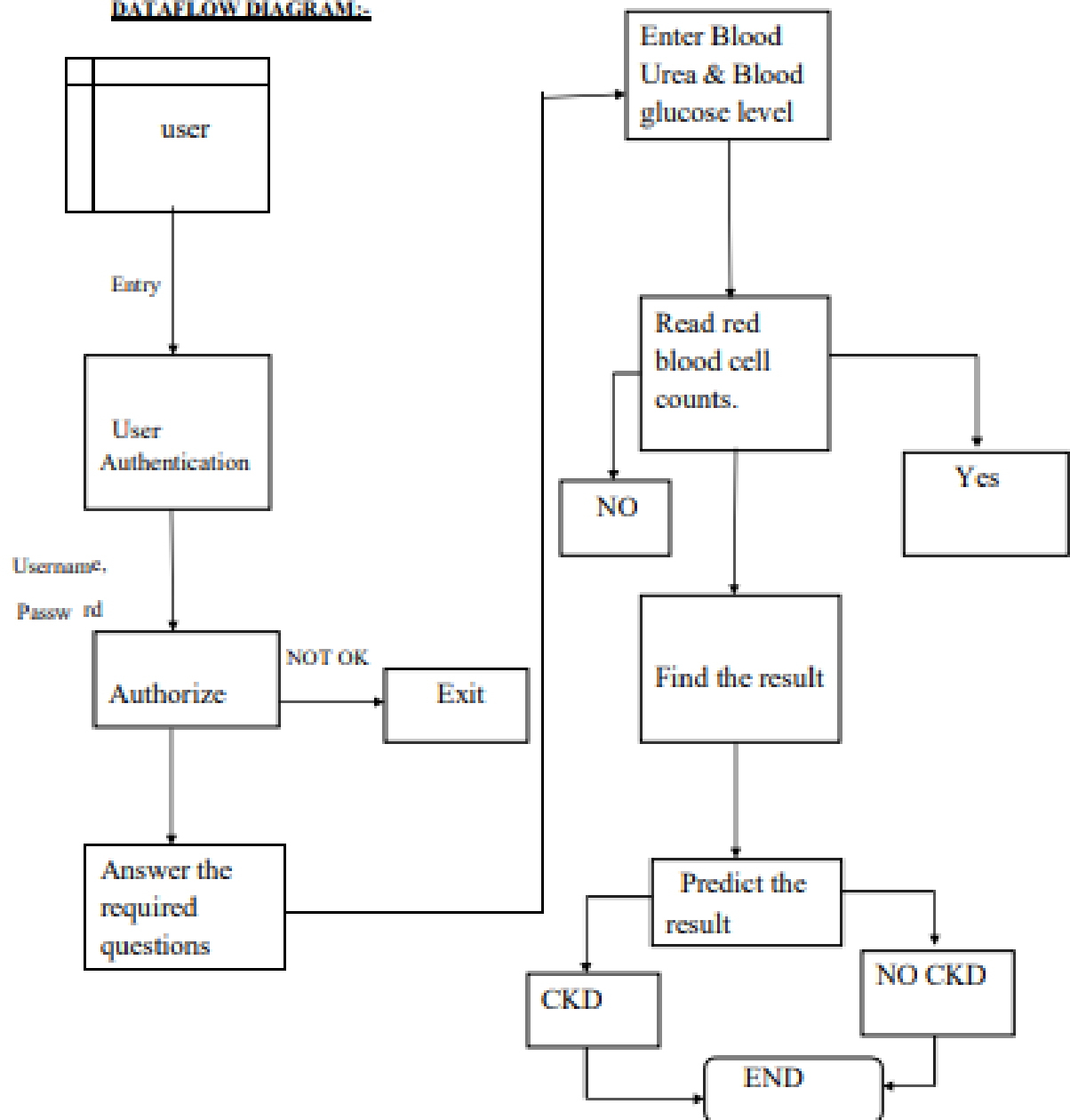
## ARCHITECTURE DIAGRAM:



## 5.2 DATA FLOW DIAGRAM AND USER STORIES

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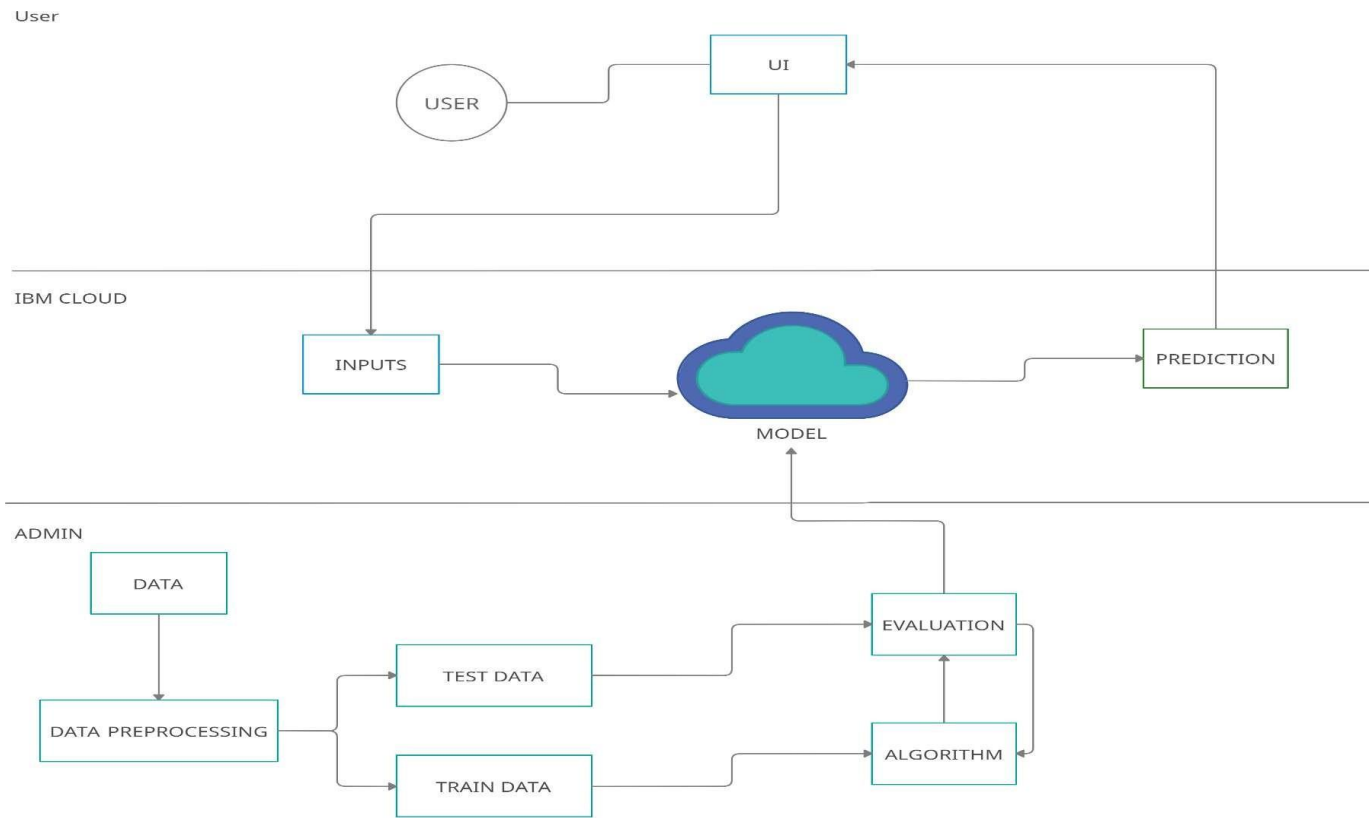
DATAFLOW DIAGRAM:-



### 5.3 TECHNICAL ARCHICTECTURE

Technology Architecture describe the logical software and hardware capabilities that are required to support the deployment of business, data, and application services. This includes IP infrastructure, middleware, networks, communications, processing, standards, etc.

Technology architecture deals with the deployment of application components on technology components. A standard set of predefined technology components is provided in order to represent servers, network, workstations.





## 6.0 CODING

```
In [1]: import pandas as pd

In [2]: import numpy as np

In [3]: from collections import Counter as c

In [4]: import matplotlib.pyplot as plt

In [5]: import seaborn as sns

In [6]: import missingno as msno

In [7]: from sklearn.metrics import accuracy_score, confusion_matrix

In [8]: from sklearn.model_selection import train_test_split

In [9]: from sklearn.preprocessing import LabelEncoder

In [10]: from sklearn.linear_model import LogisticRegression

In [11]: import pickle

In [12]: data=pd.read_csv("F:\SEM 7\IBM Project\kidney_disease.csv")

In [13]: data.head()
```

Out[13]:

	id	age	bp	sg	al	su	rbc	pc	pcc	ba	...	pcv	wc	rc	htn	dm	cad	appet	pe	ane	classification
0	0	48.0	80.0	1.020	1.0	0.0	NaN	normal	notpresent	notpresent	...	44	7800	5.2	yes	yes	no	good	no	no	ckd
1	1	7.0	50.0	1.020	4.0	0.0	NaN	normal	notpresent	notpresent	...	38	6000	NaN	no	no	no	good	no	no	ckd
2	2	62.0	80.0	1.010	2.0	3.0	normal	normal	notpresent	notpresent	...	31	7500	NaN	no	yes	no	poor	no	yes	ckd
3	3	48.0	70.0	1.005	4.0	0.0	normal	abnormal	present	notpresent	...	32	6700	3.9	yes	no	no	poor	yes	yes	ckd
4	4	51.0	80.0	1.010	2.0	0.0	normal	normal	notpresent	notpresent	...	35	7300	4.6	no	no	no	good	no	no	ckd

5 rows × 26 columns

```
In [14]: data.tail()
```

Out[14]:

	id	age	bp	sg	al	su	rbc	pc	pcc	ba	...	pcv	wc	rc	htn	dm	cad	appet	pe	ane	classification
395	395	55.0	80.0	1.020	0.0	0.0	normal	normal	notpresent	notpresent	...	47	6700	4.9	no	no	no	good	no	no	notckd
396	396	42.0	70.0	1.025	0.0	0.0	normal	normal	notpresent	notpresent	...	54	7800	6.2	no	no	no	good	no	no	notckd
397	397	12.0	80.0	1.020	0.0	0.0	normal	normal	notpresent	notpresent	...	49	6600	5.4	no	no	no	good	no	no	notckd
398	398	17.0	60.0	1.025	0.0	0.0	normal	normal	notpresent	notpresent	...	51	7200	5.9	no	no	no	good	no	no	notckd
399	399	58.0	80.0	1.025	0.0	0.0	normal	normal	notpresent	notpresent	...	53	6800	6.1	no	no	no	good	no	no	notckd

5 rows × 26 columns

```
In [15]: data.head(10)
```

Out[15]:

	id	age	bp	sg	al	su	rbc	pc	pcc	ba	...	pcv	wc	rc	htn	dm	cad	appet	pe	ane	classification
0	0	48.0	80.0	1.020	1.0	0.0	NaN	normal	notpresent	notpresent	...	44	7800	5.2	yes	yes	no	good	no	no	ckd
1	1	7.0	50.0	1.020	4.0	0.0	NaN	normal	notpresent	notpresent	...	38	6000	NaN	no	no	no	good	no	no	ckd
2	2	62.0	80.0	1.010	2.0	3.0	normal	normal	notpresent	notpresent	...	31	7500	NaN	no	yes	no	poor	no	yes	ckd
3	3	48.0	70.0	1.005	4.0	0.0	normal	abnormal	present	notpresent	...	32	6700	3.9	yes	no	no	poor	yes	yes	ckd
4	4	51.0	80.0	1.010	2.0	0.0	normal	normal	notpresent	notpresent	...	35	7300	4.6	no	no	no	good	no	no	ckd
5	5	60.0	90.0	1.015	3.0	0.0	NaN	NaN	notpresent	notpresent	...	39	7800	4.4	yes	yes	no	good	yes	no	ckd
6	6	68.0	70.0	1.010	0.0	0.0	NaN	normal	notpresent	notpresent	...	36	NaN	NaN	no	no	no	good	no	no	ckd
7	7	24.0	NaN	1.015	2.0	4.0	normal	abnormal	notpresent	notpresent	...	44	6900	5	no	yes	no	good	yes	no	ckd
8	8	52.0	100.0	1.015	3.0	0.0	normal	abnormal	present	notpresent	...	33	9600	4	yes	yes	no	good	no	yes	ckd
9	9	53.0	90.0	1.020	2.0	0.0	abnormal	abnormal	present	notpresent	...	29	12100	3.7	yes	yes	no	poor	no	yes	ckd

10 rows × 26 columns

```
In [17]: data.head(10)
```

```
Out[17]:
```

	age	bp	sg	al	su	rbc	pc	pcc	ba	bgr	...	pcv	wc	rc	htn	dm	cad	appet	pe	ane	classification
0	48.0	80.0	1.020	1.0	0.0	NaN	normal	notpresent	notpresent	121.0	...	44	7800	5.2	yes	yes	no	good	no	no	ckd
1	7.0	50.0	1.020	4.0	0.0	NaN	normal	notpresent	notpresent	NaN	...	38	6000	NaN	no	no	no	good	no	no	ckd
2	62.0	80.0	1.010	2.0	3.0	normal	normal	notpresent	notpresent	423.0	...	31	7500	NaN	no	yes	no	poor	no	yes	ckd
3	48.0	70.0	1.005	4.0	0.0	normal	abnormal	present	notpresent	117.0	...	32	6700	3.9	yes	no	no	poor	yes	yes	ckd
4	51.0	80.0	1.010	2.0	0.0	normal	normal	notpresent	notpresent	106.0	...	35	7300	4.6	no	no	no	good	no	no	ckd
5	60.0	90.0	1.015	3.0	0.0	NaN	NaN	notpresent	notpresent	74.0	...	39	7800	4.4	yes	yes	no	good	yes	no	ckd
6	68.0	70.0	1.010	0.0	0.0	NaN	normal	notpresent	notpresent	100.0	...	36	NaN	NaN	no	no	no	good	no	no	ckd
7	24.0	NaN	1.015	2.0	4.0	normal	abnormal	notpresent	notpresent	410.0	...	44	6900	5	no	yes	no	good	yes	no	ckd
8	52.0	100.0	1.015	3.0	0.0	normal	abnormal	present	notpresent	138.0	...	33	9600	4	yes	yes	no	good	no	yes	ckd
9	53.0	90.0	1.020	2.0	0.0	abnormal	abnormal	present	notpresent	70.0	...	29	12100	3.7	yes	yes	no	poor	no	yes	ckd

10 rows x 25 columns

```
In [18]: data.columns
```

```
Out[18]: Index(['age', 'bp', 'sg', 'al', 'su', 'rbc', 'pc', 'pcc', 'ba', 'bgr', 'bu',  
              'sc', 'sod', 'pot', 'hemo', 'pcv', 'wc', 'rc', 'htn', 'dm', 'cad',  
              'appet', 'pe', 'ane', 'classification'],  
              dtype='object')
```

```
In [19]: data.columns=['age','blood_pressure','specific_gravity','albumin','sugar','red_blood_cells','pus_cell','pus_cell_clumps','bacteria','bloo
```

```
In [20]: data.columns
```

```
Out[20]: Index(['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'],  
              dtype='object')
```

```
In [21]: data.columns
```

```
Out[21]: Index(['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'],  
              dtype='object')
```

```
In [22]: data.info()
```

RangeIndex: 400 entries, 0 to 399

Data columns (total 25 columns):

#	Column	Non-Null Count	Dtype
0	age	391 non-null	float64
1	blood_pressure	388 non-null	float64
2	specific_gravity	353 non-null	float64
3	albumin	354 non-null	float64
4	sugar	351 non-null	float64
5	red_blood_cells	248 non-null	object
6	pus_cell	335 non-null	object
7	pus_cell_clumps	396 non-null	object
8	bacteria	396 non-null	object
9	blood glucose random	356 non-null	float64
10	blood_urea	381 non-null	float64
11	serum_creatinine	383 non-null	float64
12	sodium	313 non-null	float64
13	potassium	312 non-null	float64
14	hemoglobin	348 non-null	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
18	hypertension	398 non-null	object
19	diabetesmellitus	398 non-null	object
20	coronary_artery_disease	398 non-null	object
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

```
In [23]: data['class'].unique()
```

```
Out[23]: array(['ckd', 'ckd\t', 'notckd'], dtype=object)
```

```
In [24]: data['class']=data['class'].replace("ckd\t","ckd")  
data['class'].unique()
```

```
Out[24]: array(['ckd', 'notckd'], dtype=object)
```





```
In [25]: catcols=set(data.dtypes[data.dtypes!='0'].index.values)
print(catcols)
```

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

```
In [26]: for i in catcols:
print("Columns:",i)
print(c(data[i]))
print('***120+\\n')
```

Columns: anemia

Counter({'no': 339, 'yes': 60, nan: 1})

\*\*\*\*\*

Columns: appetite

Counter({'good': 317, 'poor': 82, nan: 1})

\*\*\*\*\*

Columns: pus\_cell\_clumps

Counter({'notpresent': 354, 'present': 42, nan: 4})

\*\*\*\*\*

Columns: packed\_cell\_volume

Counter({nan: 70, '52': 21, '41': 21, '44': 19, '48': 19, '40': 16, '43': 14, '45': 13, '42': 13, '32': 12, '36': 12, '33': 12, '28': 12, '50': 12, '37': 11, '34': 11, '35': 9, '29': 9, '30': 9, '46': 9, '31': 8, '39': 7, '24': 7, '26': 6, '38': 5, '47': 4, '49': 4, '53': 4, '51': 4, '54': 4, '27': 3, '22': 3, '25': 3, '23': 2, '19': 2, '16': 1, '\t?': 1, '14': 1, '18': 1, '17': 1, '15': 1, '21': 1, '20': 1, '\t43': 1, '9': 1})

\*\*\*\*\*

Columns: red\_blood\_cell\_count

Counter({nan: 130, '5.2': 18, '4.5': 16, '4.9': 14, '4.7': 11, '3.9': 10, '5': 10, '4.8': 10, '4.6': 9, '3.4': 9, '3.7': 8, '6.1': 8, '5.5': 8, '5.9': 8, '3.8': 7, '5.4': 7, '5.8': 7, '5.3': 7, '4': 6, '4.3': 6, '4.2': 6, '5.6': 6, '4.4': 5, '3.2': 5, '4.1': 5, '6.2': 5, '5.1': 5, '6.4': 5, '5.7': 5, '6.5': 5, '3.6': 4, '6': 4, '6.3': 4, '3.5': 3, '3.3': 3, '3': 3, '2.6': 2, '2.8': 2, '2.5': 2, '3.1': 2, '2.1': 2, '2.9': 2, '2.7': 2, '2.3': 1, '8': 1, '2.4': 1, '\t?': 1})

\*\*\*\*\*

Columns: class

Counter({'ckd': 250, 'notckd': 150})

\*\*\*\*\*

Columns: bacteria

Counter({'notpresent': 374, 'present': 22, nan: 4})

\*\*\*\*\*

Columns: hypertension

Counter({'no': 251, 'yes': 147, nan: 2})

\*\*\*\*\*

Columns: pus\_cell

Counter({'normal': 259, 'abnormal': 76, nan: 65})

\*\*\*\*\*

Columns: pedal\_edema

Counter({'no': 323, 'yes': 76, nan: 1})

\*\*\*\*\*

Columns: red\_blood\_cells

Counter({'normal': 201, nan: 152, 'abnormal': 47})

\*\*\*\*\*

Columns: coronary\_artery\_disease

Counter({'no': 362, 'yes': 34, '\tno': 2, nan: 2})

\*\*\*\*\*

Columns: white\_blood\_cell\_count

Counter({nan: 105, '9800': 11, '6700': 10, '9600': 9, '9200': 9, '7200': 9, '6900': 8, '11000': 8, '5800': 8, '7800': 7, '9100': 7, '9400': 7, '7000': 7, '4300': 6, '6300': 6, '10700': 6, '10500': 6, '7500': 5, '8300': 5, '7900': 5, '8600': 5, '5600': 5, '10200': 5, '5000': 5, '8100': 5, '9500': 5, '6000': 4, '6200': 4, '10300': 4, '7700': 4, '5500': 4, '10400': 4, '6800': 4, '6500': 4, '4700': 4, '7300': 3, '4500': 3, '8400': 3, '6400': 3, '4200': 3, '7400': 3, '8000': 3, '5400': 3, '3800': 2, '11400': 2, '5300': 2, '8500': 2, '14600': 2, '7100': 2, '13200': 2, '9000': 2, '8200': 2, '15200': 2, '12400': 2, '12800': 2, '8800': 2, '5700': 2, '9300': 2, '6600': 2, '12100': 1, '12200': 1, '18900': 1, '21600': 1, '11300': 1, '\t6200': 1, '11800': 1, '12500': 1, '11900': 1, '12700': 1, '13600': 1, '14900': 1, '16300': 1, '\t8400': 1, '10900': 1, '2200': 1, '11200': 1, '19100': 1, '\t?': 1, '12300': 1, '16700': 1, '2600': 1, '26400': 1, '4900': 1, '12000': 1, '15700': 1, '4100': 1, '11500': 1, '10800': 1, '9900': 1, '5200': 1, '5900': 1, '9700': 1, '5100': 1})

\*\*\*\*\*

Columns: diabetesmellitus

Counter({'no': 258, 'yes': 134, '\tno': 3, '\tyes': 2, nan: 2, 'yes': 1})

\*\*\*\*\*

```

In [30]: contcols.remove('specific_gravity')
         contcols.remove('albumin')
         contcols.remove('sugar')
         print(contcols)

{'serum_creatinine', 'blood glucose random', 'potassium', 'sodium', 'blood_pressure', 'blood_urea', 'hemoglobin', 'age'}

In [31]: contcols.add('red_blood_cell_count')
         contcols.add('packed_cell_volume')
         contcols.add('white_blood_cell_count')
         print(contcols)

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

In [32]: catcols.add('specific_gravity')
         catcols.add('albumin')
         catcols.add('sugar')
         print(catcols)

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

In [33]: data['coronary_artery_disease']=data.coronary_artery_disease.replace('\tno','no')
         c(data['coronary_artery_disease'])

Out[33]: Counter({'no': 364, 'yes': 34, nan: 2})

In [34]: data['diabetesmellitus']=data.diabetesmellitus.replace(to_replace=('\tno':'no','\tyes':'yes',' yes':'yes',' no':'no'))
         c(data['diabetesmellitus'])

Out[34]: Counter({'yes': 137, 'no': 261, nan: 2})

```

```

In [35]: data.isnull().any()

Out[35]: age                True
         blood_pressure      True
         specific_gravity    True
         albumin             True
         sugar               True
         red_blood_cells     True
         pus_cell            True
         pus_cell_clumps     True
         bacteria            True
         blood glucose random True
         blood_urea          True
         serum_creatinine    True
         sodium              True
         potassium           True
         hemoglobin          True
         packed_cell_volume  True
         white_blood_cell_count True
         red_blood_cell_count True
         hypertension        True
         diabetesmellitus    True
         coronary_artery_disease True
         appetite            True
         pedal_edema         True
         anemia              True
         class               False
         dtype: bool

```

```

In [36]: data.isnull().sum()

Out[36]: age                9
         blood_pressure      12
         specific_gravity    47
         albumin             46
         sugar               49
         red_blood_cells     152
         pus_cell            65
         pus_cell_clumps     4
         bacteria            4
         blood glucose random 44
         blood_urea          19
         serum_creatinine    17
         sodium              87
         potassium           88
         hemoglobin          52
         packed_cell_volume  70
         white_blood_cell_count 105
         red_blood_cell_count 130
         hypertension        2
         diabetesmellitus    2
         coronary_artery_disease 2
         appetite            1
         pedal_edema         1
         anemia              1
         class               0
         dtype: int64

```

```

In [37]: 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')

In [38]: 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)

In [39]: 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)

```

```

In [40]: from sklearn.preprocessing import LabelEncoder

```

```

In [41]: 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)

```

```

LABEL ENCODING OF: albumin
Counter({0: 245, 1: 44, 2: 43, 3: 43, 4: 24, 5: 1})
Counter({0: 245, 1: 44, 2: 43, 3: 43, 4: 24, 5: 1})

```

```

*****
LABEL ENCODING OF: sugar
Counter({0: 339, 2: 18, 3: 14, 4: 13, 1: 13, 5: 3})
Counter({0: 339, 2: 18, 3: 14, 4: 13, 1: 13, 5: 3})

```

```

*****
LABEL ENCODING OF: anemia
Counter({'no': 340, 'yes': 60})
Counter({0: 340, 1: 60})

```

```

*****
LABEL ENCODING OF: appetite
Counter({'good': 318, 'poor': 82})
Counter({0: 318, 1: 82})

```

```

*****
LABEL ENCODING OF: pus_cell_clumps
Counter({'notpresent': 358, 'present': 42})
Counter({0: 358, 1: 42})

```

```

*****
LABEL ENCODING OF: class
Counter({'ckd': 250, 'notckd': 150})
Counter({0: 250, 1: 150})

```

```

*****
LABEL ENCODING OF: bacteria
Counter({'notpresent': 378, 'present': 22})
Counter({0: 378, 1: 22})

```

```

*****
LABEL ENCODING OF: hypertension
Counter({'no': 253, 'yes': 147})
Counter({0: 253, 1: 147})

```

```

*****
LABEL ENCODING OF: pus_cell
Counter({'normal': 324, 'abnormal': 76})
Counter({1: 324, 0: 76})

```

```

*****
LABEL ENCODING OF: pedal_edema
Counter({'no': 324, 'yes': 76})
Counter({0: 324, 1: 76})

```

```

*****
LABEL ENCODING OF: red_blood_cells
Counter({'normal': 353, 'abnormal': 47})
Counter({1: 353, 0: 47})

```

```

*****
LABEL ENCODING OF: coronary_artery_disease
Counter({'no': 366, 'yes': 34})
Counter({0: 366, 1: 34})

```

```

*****
LABEL ENCODING OF: specific_gravity
Counter({1.02: 153, 1.01: 84, 1.025: 81, 1.015: 75, 1.005: 7})
Counter({3: 153, 1: 84, 4: 81, 2: 75, 0: 7})

```

```

*****
LABEL ENCODING OF: diabetesmellitus
Counter({'no': 263, 'yes': 137})
Counter({0: 263, 1: 137})

```

```

In [42]: selcols=['red_blood_cells','pus_cell','blood glucose random','blood_urea','pedal_edema','anemia','diabetesmellitus','coronary_artery_dis

In [43]: x=pd.DataFrame(data,columns=selcols)
y=pd.DataFrame(data,columns=['class'])
print(x.shape)
print(y.shape)

(400, 8)
(400, 1)

In [44]: 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)

In [45]: from sklearn.ensemble import RandomForestClassifier
lgr = RandomForestClassifier()

In [46]: #from sklearn.linear_model import LogisticRegression

In [47]: #lgr=LogisticRegression(solver='lbfgs', max_iter=1000)
lgr.fit(x_train.values, y_train.values.ravel())

Out[47]: RandomForestClassifier()

In [48]: y_pred=lgr.predict(x_test.values)

In [49]: y_pred1=lgr.predict([[90,157,0,1,1,0,1,1]])
print(y_pred1)
c(y_pred1)

[0]
Out[49]: Counter({0: 1})

In [50]: accuracy_score(y_test,y_pred)

Out[50]: 0.95

In [51]: conf_mat=confusion_matrix(y_test,y_pred)
conf_mat

Out[51]: array([[52,  2],
               [ 2, 24]], dtype=int64)

In [52]: pickle.dump(lgr,open('CKD.pkl','wb'))

```



```
CKD FINAL
> .idea
> .ipynb_checkpoints
> templates
> images
  > home.html
  > indexnew.html
  > result.html
> venv
  > app.py
  > CKD Prediction Notebook.ipynb
  > CKD.pkl
  > kidney_disease.csv

app.py > predict
1  import numpy as np
2  import pandas as pd
3  from flask import Flask, request, render_template
4  import pickle
5
6  app = Flask(__name__)
7  model = pickle.load(open('CKD.pkl', 'rb'))
8
9
10 @app.route('/')
11 def home():
12     return render_template('home.html')
13
14
15 @app.route('/Prediction', methods=['POST', 'GET'])
16 def prediction():
17     return render_template('indexnew.html')
18
19
20 @app.route('/Home', methods=['POST', 'GET'])
21 def my_home():
22     return render_template('home.html')
23
24
25 @app.route('/predict', methods=['POST'])
26 def predict():
27     #input_features = ([int(x) for x in request.form.values()])
28     blood_urea = request.form["blood_urea"]
29     blood_glucose_random = request.form["blood_glucose_random"]
30     anemia = request.form["Anemia"]
31     if (anemia == "no"):
32         anemia = 0
33     if (anemia == "yes"):
```

Building a flask file :

```
CKD FINAL
> .idea
> .ipynb_checkpoints
> templates
> images
  > home.html
  > indexnew.html
  > result.html
> venv
  > app.py
  > CKD Prediction Notebook.ipynb
  > CKD.pkl
  > kidney_disease.csv

app.py > predict
51     red_blood_cell = 1
52
53     diabetics_mellitus = request.form["diabetics_mellitus"]
54     if (diabetics_mellitus == "no"):
55         diabetics_mellitus = 0
56     if (diabetics_mellitus == "yes"):
57         diabetics_mellitus = 1
58
59     pedal_edema = request.form["pedal_edema"]
60     if (pedal_edema == "no"):
61         pedal_edema = 0
62     if (pedal_edema == "yes"):
63         pedal_edema = 1
64
65     input_features = [int(blood_urea),int(blood_glucose_random),int(anemia),int(coronary_artery_disease),int(pus_cell),int(red_blood_cell)]
66     #input_features = [int(red_blood_cell),int(pus_cell),int(blood_glucose_random),int(blood_urea),int(pedal_edema),int(anemia),int(diabetics_mellitus)]
67     print(input_features)
68     features_value = [np.array(input_features)]
69
70
71     #features_name = ['red_blood_cells','pus_cell','blood glucose random','blood_urea','pedal_edema','anemia','diabetesmellitus','coronary_artery_disease']
72     features_name = ['blood_urea','blood glucose random','anemia','coronary_artery_disease','pus_cell','red_blood_cells','diabetesmellitus']
73     df = pd.DataFrame(features_value, columns=features_name)
74     output = model.predict(df)
75     return render_template('result.html', prediction_text=output)
76
77
78 # Press the green button in the gutter to run the script.
79 if __name__ == '__main__':
80     app.run(host='localhost', debug=True)
81
82 # See PyCharm help at https://www.jetbrains.com/help/pycharm/
```

Home.html screen shots:

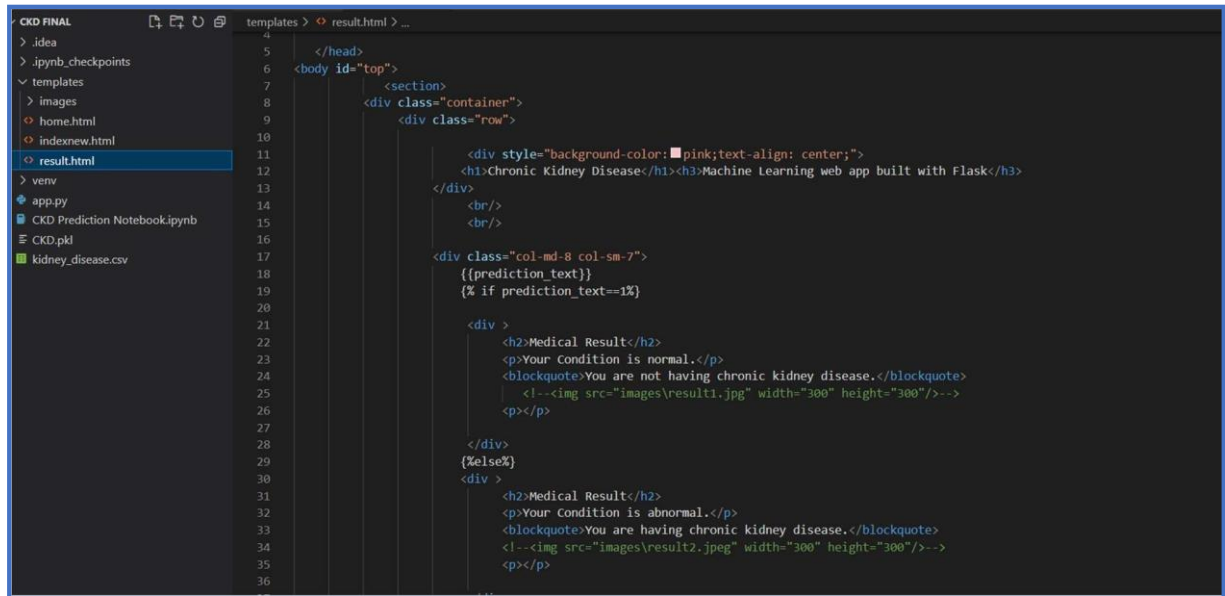
```
CKD FINAL
> .idea
> .ipynb_checkpoints
> templates
> images
> home.html
> indexnew.html
> result.html
> venv
app.py
CKD Prediction Notebook.ipynb
CKD.pkl
kidney_disease.csv

templates > home.html > html > body > form > section#home > div.container > div.row > div.owl-carousel.owl-theme
1 <html lang="en">
2 <head>
3
4 <title>Chronic Kidney Disease prediction</title>
5
6 <!-- <link rel="stylesheet" href="css/tooplate-style.css"-->
7
8 </head>
9 <body>
10 <form action="{{url_for('prediction')}}" method="POST">
11 <!--<img src='C:\Users\kgaru\Desktop\ibm_chronic_kidney_disease\app_ibm\templates\images\slider5.jpeg' width="400" height="200" alt=
12
13 <section id="home">
14
15 <div class="container">
16 <div class="row">
17 <div class="owl-carousel owl-theme">
18 <div class="item item-first">
19 <div class="caption">
20 <div style="background-color: #pink;text-align: center;">
21 <h1>Chronic Kidney Disease</h1><h3>Machine Learning web app built with Flask</h3>
22 </div>
23 <div class="col-md-12 col-sm-12">
24 <button type="submit" class="form-control" id="cf-submit" name="submit">prediction</button>
25 <!--
26
27 </div>
28 </div>
29
30 </div>
31
32 </div>
33 </div>
```

```
CKD FINAL
> .idea
> .ipynb_checkpoints
> templates
> images
> home.html
> indexnew.html
> result.html
> venv
app.py
CKD Prediction Notebook.ipynb
CKD.pkl
kidney_disease.csv

templates > indexnew.html > ...
61
62
63
64
65 <div class="col-md-6 col-sm-6">
66 <label for="select">Select Diabetics Mellitus</label>
67 <select name = "diabetics_mellitus">
68 <option value = "yes"> yes</option>
69 <option value = "no">no </option>
70 </select>
71 </div>
72
73 <div class="col-md-6 col-sm-6">
74 <label for="select">Select Pedal Edema</label>
75 <select name = "pedal_edema">
76 <option value = "yes"> yes</option>
77 <option value = "no">no </option>
78 </select>
79 </div>
80 <div class="col-md-12 col-sm-12">
81 <button type="submit" class="form-control" id="cf-submit" name="submit">predict</button>
82 </div>
83 </form>
84
85 </div>
86
87 </div>
88 </section>
89
90 </body>
91 </html>
92
```

Result to display the screen shot



```
4
5 </head>
6 <body id="top">
7   <section>
8     <div class="container">
9       <div class="row">
10
11         <div style="background-color: #pink;text-align: center;">
12           <h1>Chronic Kidney Disease</h1><h3>Machine Learning web app built with Flask</h3>
13         </div>
14         <br/>
15         <br/>
16
17         <div class="col-md-8 col-sm-7">
18           {{prediction_text}}
19           {% if prediction_text==1%}
20
21             <div>
22               <h2>Medical Result</h2>
23               <p>Your Condition is normal.</p>
24               <blockquote>You are not having chronic kidney disease.</blockquote>
25               <!---->
26               <p></p>
27             </div>
28
29           {%else%}
30           <div>
31             <h2>Medical Result</h2>
32             <p>Your Condition is abnormal.</p>
33             <blockquote>You are having chronic kidney disease.</blockquote>
34             <!---->
35             <p></p>
36           </div>
```

## Running the app

```
(c) Microsoft Corporation. All rights reserved.

C:\Users\kgaru\Desktop\ckd final>c:\users\kgaru\anaconda3\scripts\activate

(base) C:\Users\kgaru\Desktop\ckd final>conda activate deployment

(deployment) C:\Users\kgaru\Desktop\ckd final>python app.py
c:\Users\kgaru\anaconda3\envs\deployment\lib\site-packages\sklearn\base.py:329: UserWarning: Trying to unpickle estimator DecisionTreeClassifier from version 1.0.2 when using version 1.1.2. This might lead to breaking code or invalid results. Use at your own risk. For more info please refer to: https://scikit-learn.org/stable/model_persistence.html#security-maintainability-limitations
  warnings.warn(
c:\Users\kgaru\anaconda3\envs\deployment\lib\site-packages\sklearn\base.py:329: UserWarning: Trying to unpickle estimator RandomForestClassifier from version 1.0.2 when using version 1.1.2. This might lead to breaking code or invalid results. Use at your own risk. For more info please refer to: https://scikit-learn.org/stable/model_persistence.html#security-maintainability-limitations
  warnings.warn(
```

## TEST CASE 1: CKD

Chronic Kidney Disease

Machine Learning web app built with Flask

prediction

Chronic Kidney Disease

Machine Learning web app built with Flask

Blood Urea

Blood Glucose Random

Select Anemia

Select Coronary Artery Disease

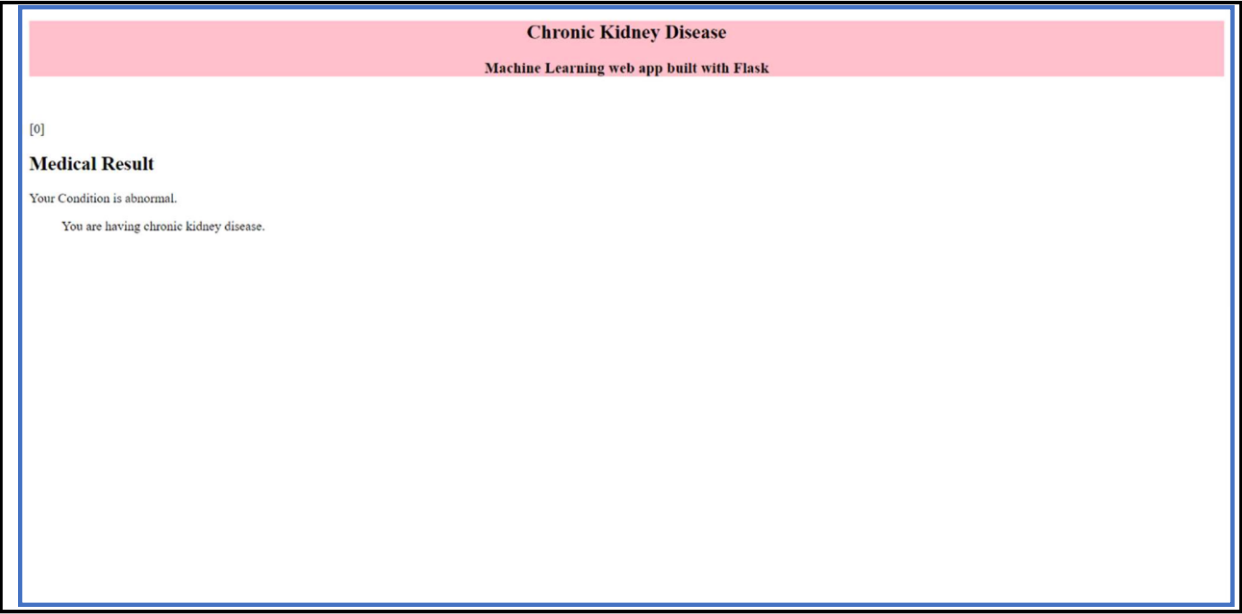
Select Pus Cell

Select Red Blood Cell

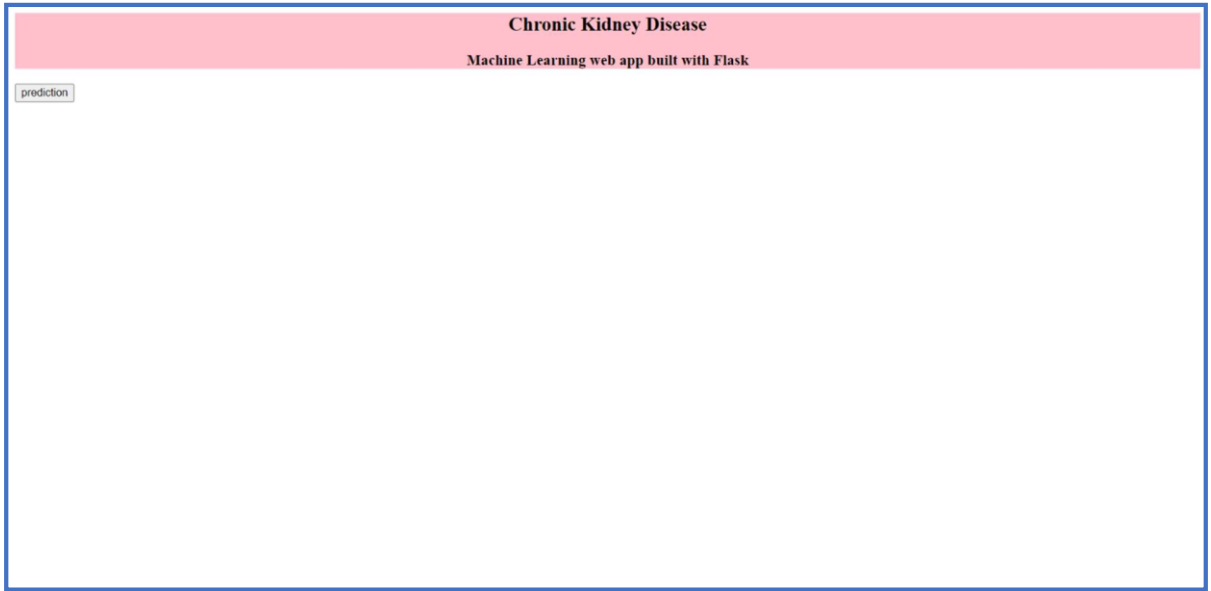
Select Diabetes Mellitus

Select Pedal Edema

predict



**TEST CASE 2: NO CKD**



Chronic Kidney Disease

Machine Learning web app built with Flask

Blood Urea

46

Blood Glucose Random

117

Select Anemia

no

Select Coronary Artery Disease

no

Select Pus Cell

no

Select Red Blood Cell

no

Select Diabetes Mellitus

no

Select Pedal Edema

no

predict

Chronic Kidney Disease

Machine Learning web app built with Flask

[1]

Medical Result

Your Condition is normal.

You are not having chronic kidney disease.

## 7.0 Conclusion:

This Project has helped team members to understand various concepts of Machine learning, Flask file, IBM cloud and Python notebook.

This project can be scaled for usage in prediction of other chronic diseases which will help doctors in diagnosis of disease at an early stage thereby helping in early detection of various disease.

### Github link:

<https://github.com/IBM-EPBL/IBM-Project-53532-1661414687>

