

KidniFy

Your Kidney Care Companion

23-032



Outline

❑ Our Team

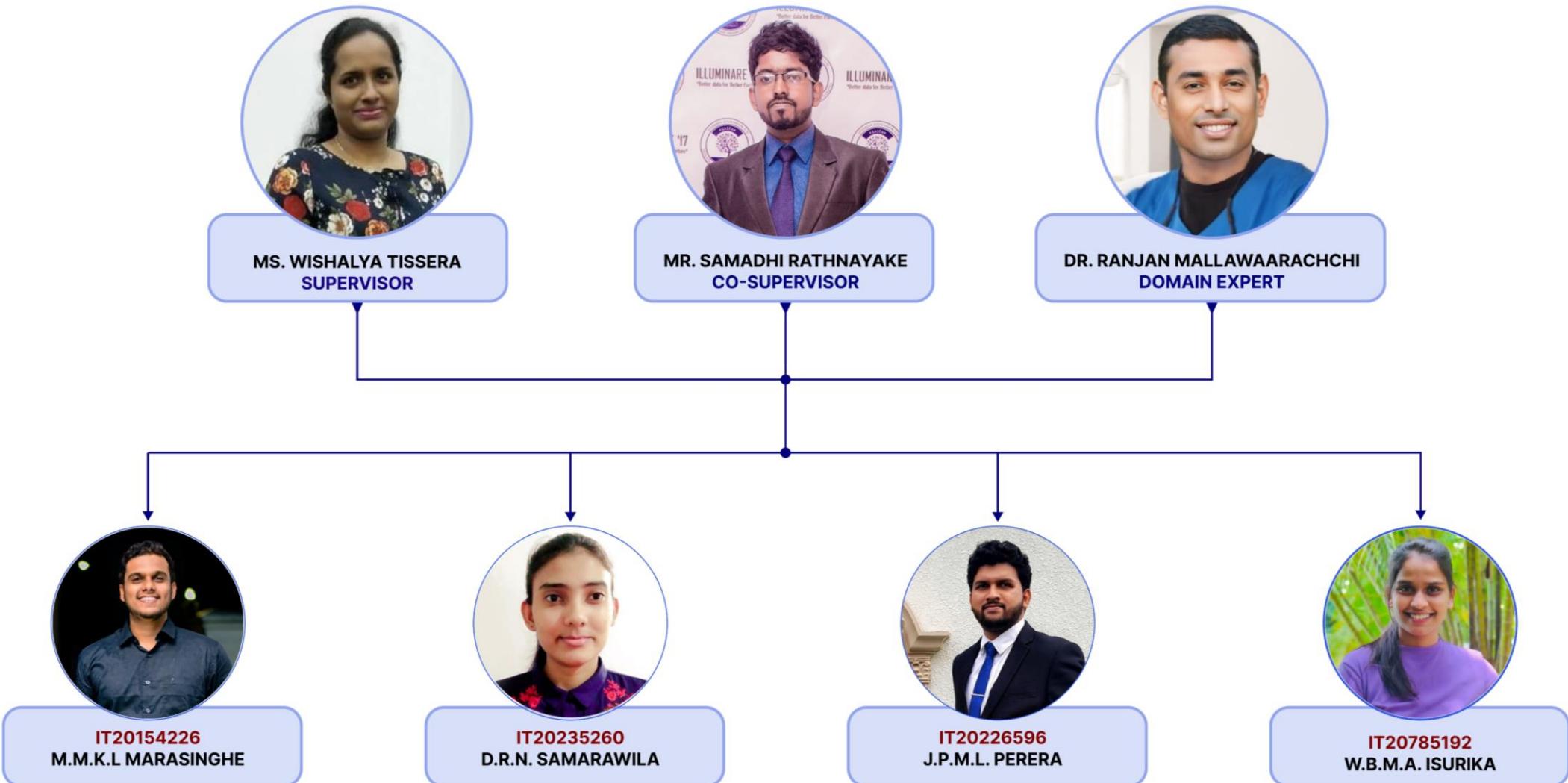
❑ Introduction

- Background
- Research Problem
- Overall Problem
- System Overview
- Gantt Chart
- Commercialization

❑ Individual Components

- Background
- Research Problem
- Research Gap
- Objectives
- Methodology

Our Team



Research Problem

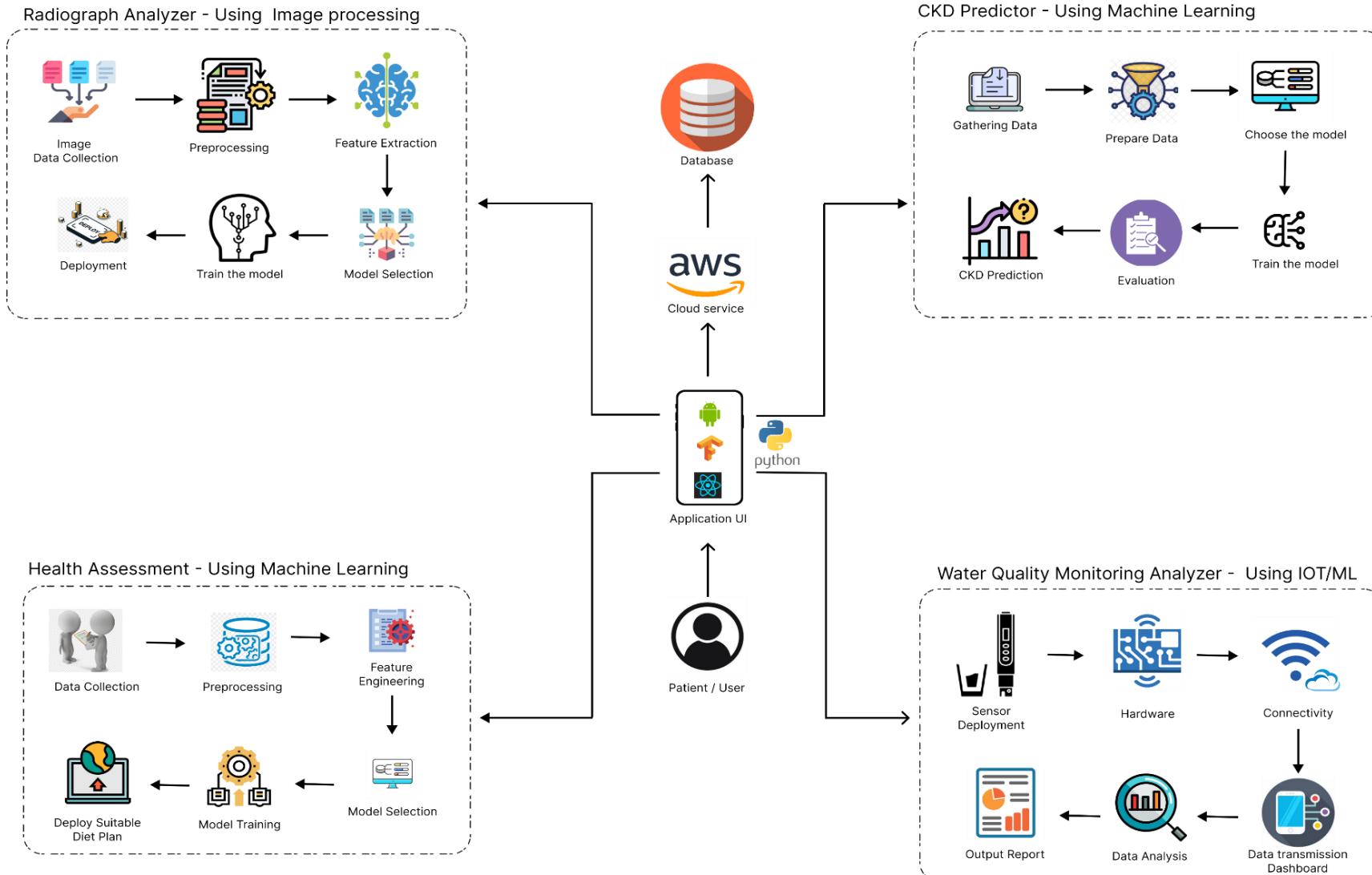
- How to develop an accurate ML model for kidney disease prediction?
- How to use image processing to analyze CKD patients' medical images?
- What are the efficient ways to test the water quality that affects kidney patients, combining IoT as well as ML?
- How to develop a personalized dietary prediction system for CKD patients?

Objectives

To develop a mobile app that is easy to use yet effective to

- Accurately predict about kidney health based on real time data.
- Get timely diagnosis of kidney disease using image processing
- Measure the quality of water with the use of an IOT device
- Analyse patient data and provide personalized diet plans

System Overview





Individual Components

IT20235260 | D.R.N. Samarawila

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Predictive analysis for risks of having a Kidney Disease

Background

- CKD is a global problem, identified as the 7th most common cause of death in the world.
- In Sri Lanka prevalence of CKD also has been increased alarmingly in recent decades.
- Kidney diseases are major health concern, impacting a considerable amount of the population, affecting approximately 18% of the population and it is considerable value from a population.
- In addition to these main types of CKD, there are also other less common causes of kidney disease, such as kidney stones, tumors, and structural abnormalities.
- Identify a CKD in early stages and the importance of that.
- Use the modern technology to aware and to prevent from the CKD.

Research Problem

Detecting and treating kidney disease early is essential for preventing complications and improving patient outcomes.

1. How accurate are existing ML-based models for kidney disease prediction in the Sri Lankan population?
2. What factors may influence the performance of these models, such as age, gender, socioeconomic status, and geographic location?
3. What is the level of awareness among the general public in Sri Lanka about the risks and symptoms of kidney disease, and how can this be improved to facilitate early detection and treatment?
4. What are the current barriers to accessing timely and accurate diagnostic tools for kidney disease in Sri Lanka, and how can these be addressed?

- Despite the importance of the early detection and treatment there are significant challenges in identifying individuals at risk of kidney disease in Sri Lanka.
- One major challenge is the lack of access to timely and accurate diagnostic tools, particularly in rural areas where healthcare resource are often limited.



Research Gap

- This research gap highlights the need for a comprehensive solution that is accurate, easily accessible, and localized.
- Will increase the availability of kidney disease prediction tools for patients in Sri Lanka.
- Aims to address this research gap by providing an accurate prediction of a user's kidney health status using multiple machine learning models.
- Based on the patient's location, the application will display a list of the nearest medical facilities where they can go through the necessary tests.

Objectives

Main Objective

- Develop an effective solution for predicting a user's kidney health condition with high accuracy.

Sub Objectives

- Gathering the data and preprocessed that data.
- Train highly accurate ML model with collected data.
- Giving the prediction according to the real time test data.
- Giving the instructions and the recommend treatments
- Based on patient's health condition and the current location, display a list of the nearest medical facilities where they can go through the necessary tests.

Completion of the Component

❑ Requirements Analysis

- Collect data manually and create data set

Attributes

- | | |
|-------------------|------------------------------|
| 1. ID | 9.Prolong use of medications |
| 2. Age | 10.Urinary Obstructions |
| 3. Gender | 11.Edema Symptoms |
| 4. Diabetic | 12. Urinary frequency stage |
| 5. Family History | 13.Urine color |
| 6. Obesity | 14.Location |
| 7. Smoking | |
| 8. Alcohol | |

Target variable : **Diagnose**

Gathered data

id	age	gender	diabetic	family_history	obesity	smoking	alcohol	prolong_use_of_medication	urinary_obstructions	edema_symptoms	urine_frequency_stage	urine_color	location	diagnose
1	26	Male	No	No	Yes	Yes	Yes	No	Yes	No	2	2	Kurunegala	Risky
2	57	Female	Yes	Yes	No	No	Yes	Yes	Yes	Yes	0	1	Kurunegala	Risky
3	59	Male	Yes	No	No	No	No	Yes	Yes	Yes	0	0	Kurunegala	Risky
4	39	Female	No	Yes	No	Yes	No	No	Yes	Yes	1	0	Kurunegala	Risky
5	57	Male	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	2	1	Kurunegala	Risky
6	18	Female	Yes	No	No	No	Yes	Yes	Yes	No	2	1	Kurunegala	Risky
7	41	Female	No	No	Yes	Yes	Yes	Yes	Yes	No	0	1	Kurunegala	Risky
8	19	Male	No	Yes	Yes	Yes	No	Yes	No	Yes	2	2	Kurunegala	Risky
9	27	Male	No	No	Yes	Yes	No	Yes	No	Yes	1	2	Kurunegala	Risky
10	55	Female	Yes	No	No	Yes	No	Yes	Yes	Yes	1	2	Kurunegala	Risky
11	19	Female	No	No	Yes	Yes	Yes	No	Yes	No	0	2	Kurunegala	Risky
594	40	Female	No	Yes	Yes	No	Yes	Yes	No	Yes	0	0	Kurunegala	Not Risky
595	61	Male	No	No	No	No	No	No	No	No	1	0	Kurunegala	Not Risky
596	70	Male	No	Yes	No	Yes	No	No	No	No	0	0	Kurunegala	Not Risky
597	42	Male	No	Yes	No	No	Yes	No	Yes	Yes	0	0	Kurunegala	Not Risky
598	43	Male	Yes	Yes	No	Yes	Yes	Yes	No	No	0	0	Kurunegala	Not Risky
599	55	Female	Yes	No	No	No	No	No	Yes	No	0	0	Kurunegala	Not Risky
600	26	Male	Yes	No	Yes	Yes	No	Yes	No	Yes	0	2	Kurunegala	Not Risky

Import libraries

The screenshot shows a Jupyter Notebook interface with the title "jupyter model" and a status bar indicating "Last Checkpoint: Last Tuesday at 18:58 (autosaved)". The toolbar includes File, Edit, View, Insert, Cell, Kernel, Widgets, Help, and a Python 3 (ipykernel) kernel selector. The notebook has one cell, In [111], containing Python code for importing various machine learning and visualization libraries, and loading a dataset from a CSV file. The output, Out[111], displays the first 10 rows of the dataset as a Pandas DataFrame.

```
In [111]: import pandas as pd
from sklearn.model_selection import train_test_split
from sklearn.linear_model import LinearRegression
from sklearn.metrics import accuracy_score
from sklearn.preprocessing import MinMaxScaler
from sklearn.preprocessing import LabelEncoder
from sklearn.metrics import mean_squared_error
from sklearn.linear_model import LogisticRegression

import seaborn as sns
import matplotlib.pyplot as plt

# Load the dataset
df = pd.read_csv('risk_of_ckd.csv')
df.head(10)
```

	id	age	gender	diabetic	family_history	obesity	smoking	alcohol	prolong_use_of_medication	urinary_obstructions	edema_symptoms	urine_frequency_sta
0	1	26	Male	No	No	Yes	Yes	Yes	No	Yes	Yes	No
1	2	57	Female	Yes	Yes	No	No	Yes	Yes	Yes	Yes	Yes
2	3	59	Male	Yes	No	No	No	No	Yes	Yes	Yes	Yes
3	4	39	Female	No	Yes	No	Yes	No	No	Yes	Yes	Yes
4	5	57	Male	Yes	No	Yes	Yes	Yes	Yes	Yes	Yes	Yes
5	6	18	Female	Yes	No	No	No	Yes	Yes	Yes	Yes	No
6	7	41	Female	No	No	Yes	Yes	Yes	Yes	Yes	Yes	No
7	8	19	Male	No	Yes	Yes	Yes	No	Yes	No	Yes	Yes
8	9	27	Male	No	No	Yes	Yes	No	Yes	No	Yes	Yes
9	10	55	Female	Yes	No	No	Yes	No	Yes	Yes	Yes	Yes

Data preprocessing

jupyter model Last Checkpoint: Last Tuesday at 18:58 (autosaved)

File Edit View Insert Cell Kernel Widgets Help

In [114]: `print(missing_values)`

```
id          0
age         0
gender      0
diabetic    0
family_history 0
obesity     0
smoking     0
alcohol      0
prolong_use_of_medication 0
urinary_obstructions 0
edema_symptoms 0
urine_frequency_stage 0
urine_color   0
location     0
diagnose     0
dtype: int64
```

In [115]: `print(df.dtypes)`

```
id           int64
age          int64
gender        object
diabetic      object
family_history object
obesity       object
smoking       object
alcohol        object
prolong_use_of_medication object
urinary_obstructions object
edema_symptoms object
urine_frequency_stage int64
urine_color    int64
location       object
diagnose       object
dtype: object
```

In [119]: `# Create a LabelEncoder object`
`label_encoder = LabelEncoder()`

jupyter model Last Checkpoint: Last Tuesday at 18:58 (autosaved)

File Edit View Insert Cell Kernel Widgets Help

In [119]: `# Create a LabelEncoder object`
`label_encoder = LabelEncoder()`

```
# Encode the "gender" column
df['gender_encoded'] = label_encoder.fit_transform(df['gender'])
df['diabetic_encoded'] = label_encoder.fit_transform(df['diabetic'])
df['alcohol_encoded'] = label_encoder.fit_transform(df['alcohol'])
df['urinary_obstructions_encoded'] = label_encoder.fit_transform(df['urinary_obstructions'])
df['smoking_encoded'] = label_encoder.fit_transform(df['smoking'])
df['obesity_encoded'] = label_encoder.fit_transform(df['obesity'])
df['edema_symptoms_encoded'] = label_encoder.fit_transform(df['edema_symptoms'])
df['prolong_use_of_medication_encoded'] = label_encoder.fit_transform(df['prolong_use_of_medication'])
df['family_history_encoded'] = label_encoder.fit_transform(df['family_history'])

print(df.dtypes)
```

id	int64
age	int64
gender	object
diabetic	object
family_history	object
obesity	object
smoking	object
alcohol	object
prolong_use_of_medication	object
urinary_obstructions	object
edema_symptoms	object
urine_frequency_stage	int64
urine_color	int64
location	object
diagnose	object
gender_encoded	int32
diabetic_encoded	int32
alcohol_encoded	int32
urinary_obstructions_encoded	int32
smoking_encoded	int32
obesity_encoded	int32

Jupyter model Last Checkpoint: Last Tuesday at 18:58 (autosaved)

File Edit View Insert Cell Kernel Widgets Help

Not Trusted Python 3 (ipykernel) Logout

In [131]: `columns_to_drop = ['location', 'id'] # Replace with the actual column names you want to drop
df = df.drop(columns_to_drop, axis=1)`

In [132]: `print(df.dtypes)`

age	int64
urine_frequency_stage	int64
urine_color	int64
diagnose	object
gender_encoded	int32
diabetic_encoded	int32
alcohol_encoded	int32
urinary_obstructions_encoded	int32
smoking_encoded	int32
obesity_encoded	int32
edema_symptoms_encoded	int32
prolong_use_of_medication_encoded	int32
family_history_encoded	int32
dtype: object	

In [74]:

In []: `# Select the features to scale (assuming numerical features)
numeric_features = ['blood_pressure', 'age', 'blood_sugar', 'albumin', 'hemoglobin', 'serum_creatinine', 'blood_urea_nitrogen']

Exclude non-numeric columns
numeric_df = df[numeric_features]

Create a scaler object
scaler = MinMaxScaler()

Scale the numeric features
scaled_features = scaler.fit_transform(numeric_df.values)
df[numeric_features] = scaled_features

Print the scaled dataset
print(df)`

Jupyter model Last Checkpoint: Last Tuesday at 18:58 (autosaved)

File Edit View Insert Cell Kernel Widgets Help

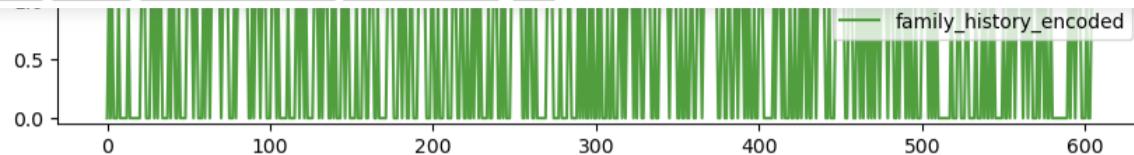
Not Trusted

Print the scaled dataset
print(df)

In [129]: `# Plot all the columns
df.plot(subplots=True, layout=(len(df.columns), 1), figsize=(10, 20))
plt.show()`

File Edit View Insert Cell Kernel Widgets Help

Not Trusted | Python 3 (ipykernel)



```
In [133]: X = df.drop('diagnose', axis=1) # Update 'target_variable' with the actual column name  
y = df['diagnose']
```

```
In [134]: print(df.dtypes)
```

```
age                int64  
urine_frequency_stage    int64  
urine_color            int64  
diagnose              object  
gender_encoded          int32  
diabetic_encoded        int32  
alcohol_encoded         int32  
urinary_obstructions_encoded    int32  
smoking_encoded         int32  
obesity_encoded         int32  
edema_symptoms_encoded    int32  
prolong_use_of_medication_encoded    int32  
family_history_encoded    int32  
dtype: object
```

```
In [135]: X_train, X_test, y_train, y_test = train_test_split(X, y, test_size=0.2, random_state=0)
```

```
In [136]: # Create a LabelEncoder object  
label_encoder = LabelEncoder()  
  
# Encode the target variable  
y_train_encoded = label_encoder.fit_transform(y_train)  
y_test_encoded = label_encoder.transform(y_test)  
  
# Build the Linear regression model  
model = LinearRegression()
```

Build the model and train

jupyter model Last Checkpoint: Last Tuesday at 18:58 (autosaved)

In [136]:

```
# Create a LabelEncoder object
label_encoder = LabelEncoder()

# Encode the target variable
y_train_encoded = label_encoder.fit_transform(y_train)
y_test_encoded = label_encoder.transform(y_test)

# Build the Linear regression model
model = LinearRegression()
model.fit(X_train, y_train_encoded)

# Predict the target variable for the test set
y_pred = model.predict(X_test)

# Calculate mean squared error (MSE)
mse = mean_squared_error(y_test_encoded, y_pred)

# Print the MSE
print("Mean Squared Error (MSE):", mse)

Mean Squared Error (MSE): 0.10486331057702959
```

In [137]:

```
from sklearn.metrics import r2_score

# Predict the target variable for the test set
y_pred = model.predict(X_test)

# Calculate R-squared
r2 = r2_score(y_test_encoded, y_pred)

# Print the R-squared
print("R-squared:", r2)

R-squared: 0.37468529061357825
```

In [138]:

```
X_train, X_test, y_train, y_test = train_test_split(X, y, test_size=0.2, random_state=42)

# Create a Logistic regression model
```

jupyter model Last Checkpoint: Last Tuesday at 18:58 (autosaved)

```
# Predict the target variable for the test set
y_pred = model.predict(X_test)

# Calculate R-squared
r2 = r2_score(y_test_encoded, y_pred)

# Print the R-squared
print("R-squared:", r2)

R-squared: 0.37468529061357825
```

In [138]:

```
X_train, X_test, y_train, y_test = train_test_split(X, y, test_size=0.2, random_state=42)

# Create a Logistic regression model
model = LogisticRegression()

# Fit the model on the training data
model.fit(X_train, y_train)

# Predict the target variable for the test set
y_pred = model.predict(X_test)

# Calculate the accuracy of the model
accuracy = accuracy_score(y_test, y_pred)

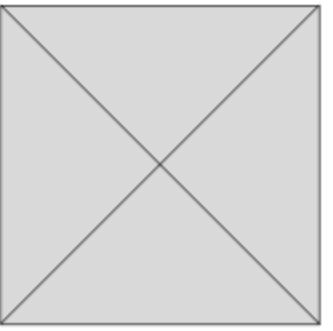
# Print the accuracy
print("Accuracy:", accuracy)

Accuracy: 0.8934426229508197
```

In []:

Wireframes

Predict your Kidney Status



Lorem ipsum dolor sit amet, consectetur adipiscing elit. Duis facilisis laoreet nisi, vel dignissim quam interdum a. Fusce nec ipsum.

Continue

Predict your Kidney Status

1 2 3

Select your gender

Female
 Male

Select your age group

Below 18
 18 - 30
 30 - 60

Next

Predict your Kidney Status

1 2 3

Do you smoke ?

Yes
 No

Do you consume Alcohol ?

Yes
 No

Family history of CKD ?

Yes
 No

Check your results

Your chance of having CKD



75%

Lorem ipsum dolor sit amet, consectetur adipiscing elit. Duis facilisis laoreet nisi, vel dignissim quam interdum a. Fusce nec ipsum.

Get more details

According to your Details

Tests

Medical Centers

Test again

What is planned to do next

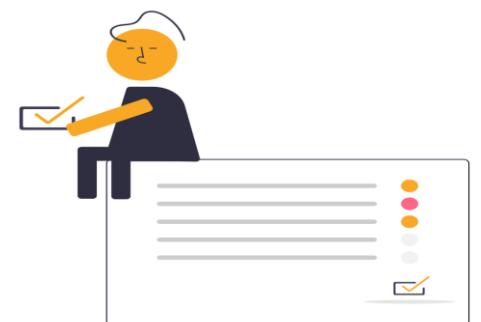
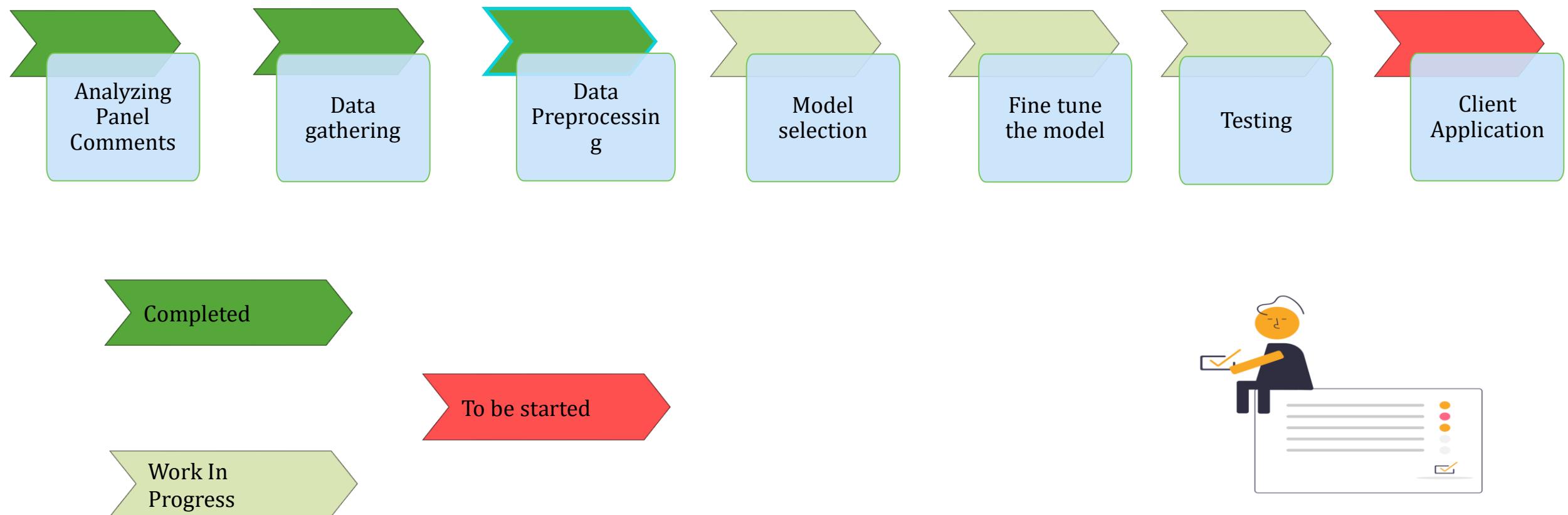
- Plan to get results accuracies for algorithms

Used Algorithm	Overall Accuracy
Linear regression	85.54%
Logistic Regression	89.17%
Random Forest	Plan to implement
Support Vector Machines (SVM)	Plan to implement
Naive Bayes	Plan to implement

- Recommend the nearest kidney treatment unit for a specific person where they can go through the necessary tests.
- Implementing the integration between the backend and frontend components of the mobile application.



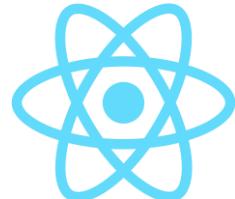
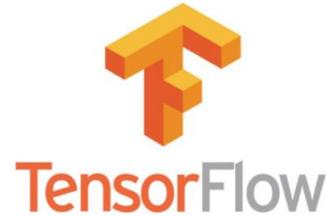
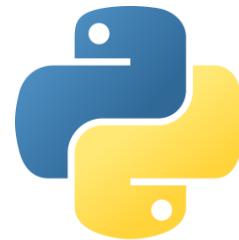
Overall Progress



Tools and Technologies

Software technologies

- Python
- TensorFlow
- AWS / Azure
- React Native
- MongoDB
- Node JS



React Native



Requirement Analysis

Functional requirements

- Gathering the data and preprocessed that data.
- Train highly accurate ML model with collected data.
- Giving the prediction according to the real time test data.
- Giving recommendations for the treatments.

Non-functional requirements

- Accuracy
- Availability
- Performance
- Speed

References

- [1] Gazi Mohammed Ifraz, "Comparative Analysis for Prediction of Kidney Disease Using Intelligent Machine Learning Methods," National Library of Medicine, 2021.
- [2] H. B. A. S. B. Abhijit V. Kshirsagar, "A Simple Algorithm to Predict Incident Kidney Disease," JAMA Network, 2008.
- [3] A. Vijayalakshmi, "Survey on Diagnosis of Chronic Kidney Disease Using Machine Learning Algorithms," IEEE, 2020.

IT20154226 | M.M.K.L. Marasinghe

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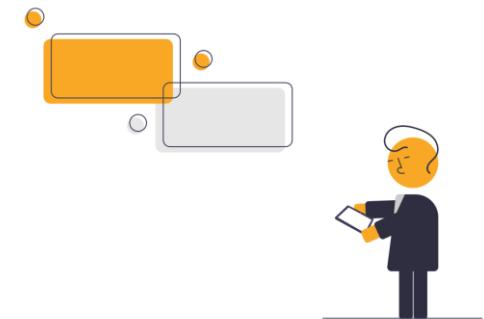
Kidney Disease Diagnosis using Image Processing

Background

- Types of image modalities
 - MRI, CT, Ultrasound, X-Ray
- Importance of medical imaging when diagnosing CKD patients
 - Accessibility, Point of Care analysis, Patient Education
- Available methods and architectures that can be used for analyze medical images
 - CNN
 - Deep Convolutional Autoencoders
 - Graph Convolutional Networks
 - Transfer Learning
 - Image Registration and Fusion

Research Problem

- How to use image processing to analyze CKD patients' medical images?
- What are the parameters to identify when analyzing medical images using image processing?
- How can image fusion techniques be utilized to integrate information from multiple imaging modalities?



Research Gap

- Developing a comprehensive, technology-driven approach to accurately diagnose kidney diseases using more advanced technologies.
- Automatic abnormality detection of kidney using CNN
- Multimodal Image Fusion to get more accurate predictions.
- Using standardization to keep track of the medical images uploaded by the user.

Objectives

- Identify parameters of the medical images
- Analyse medical images to diagnose the patient more accurately using image processing.
- Using standardization, create a visual representation of the progression of the kidneys over time.
- Using multimodal image fusion to get better accurate results.

Requirement Analysis

Functional requirements

- Analyse medical images to diagnose the patient more accurately
- Create a visual representation of the progression of the kidneys over time.

Non-functional requirements

- Accuracy
- Availability
- Performance
- Speed

Completion of the Component

- Gathering medical imaging dataset from internet resources
- Analyse the Data set
- Pre-process the Data Set
- Build the Model and Train the Model.

What's next ?

- Finalize the CNN model.
- Implement multimodal image fusion functionality to get more accurate results.
- Using standardization to track the progression of the kidney over time.
- Implementing the integration between the backend and frontend components of the mobile application.

Implementation

Loading the Image Dataset

```
import cv2
import glob

# Specifying the directory path containing the images
image_dir = 'D:\kidneyUS_images_14_june_2022\kidneyUS_images_14_june_2022/*.png' # Update with the appropriate file extension as per your dataset

# Use glob to retrieve the file paths of all images in the directory
image_paths = glob.glob(image_dir)

# Create an empty list to store the loaded images
images = []

# Loading each image using OpenCV and handle loading errors
for image_path in image_paths:
    image = cv2.imread(image_path)

    # Checking if the image is loaded successfully
    if image is not None:
        images.append(image)
    else:
        print(f"Error loading image: {image_path}")

# Checking the number of successfully loaded images
num_loaded_images = len(images)
print(f"Successfully loaded {num_loaded_images} images out of {len(image_paths)}")

Successfully loaded 534 images out of 534
```

Resizing and splitting the image dataset

```
In [2]: import cv2

# Specify the desired width and height for the resized images
desired_width = 1024
desired_height = 768

# Create an empty list to store the resized images
resized_images = []

# Resize each image using OpenCV
for image in images:
    resized_image = cv2.resize(image, (desired_width, desired_height))
    resized_images.append(resized_image)

# Check if the images are resized successfully
for i, image in enumerate(images):
    original_height, original_width = image.shape[:2]
    resized_height, resized_width = resized_images[i].shape[:2]

    if original_height != resized_height or original_width != resized_width:
        print(f"Image {i+1} was not resized correctly.")
    else:
        print(f"Image {i+1} was resized successfully.")
```

```
Image 1 was resized successfully.
Image 2 was resized successfully.
Image 3 was not resized correctly.
Image 4 was resized successfully.
Image 5 was resized successfully.
Image 6 was resized successfully.
Image 7 was resized successfully.
Image 8 was resized successfully.
Image 9 was resized successfully.
Image 10 was resized successfully.
```

```
# Check if any images were dropped during resizing
if len(images) == len(resized_images_success):
    print("All images were successfully resized.")
else:
    print(f"{len(images) - len(resized_images_success)} images were dropped during resizing.")
```

```
All images were successfully resized.
```

```
# Create an empty list to store the successfully resized images
resized_images_success = []

# Create a new list to store the indices of the successfully resized images
resized_indices_success = []

# Iterate over the original images and their corresponding resized images
for i, (image, resized_image) in enumerate(zip(images, resized_images)):
    if resized_image is not None:
        # If the resized image is not None, it indicates successful resizing
        resized_images_success.append(resized_image)
        resized_indices_success.append(i)
    else:
        print(f"Image {i+1} was not resized successfully and will be dropped.")

# Update the 'images' list with the successfully resized images
images = resized_images_success
```

```
from sklearn.model_selection import train_test_split
import random

# Define the labels
labels = ["1", "2", "3", "3", "4", "5"]

# Assign labels randomly to the resized images
random_labels = random.choices(labels, k=len(resized_images_success))

# Split the data into training, validation, and testing sets
train_images, test_images, train_labels, test_labels = train_test_split(resized_images_success, random_labels, test_size=0.2, random_state=42)
train_images, val_images, train_labels, val_labels = train_test_split(train_images, train_labels, test_size=0.2, random_state=42)

# Print the sizes of the resulting sets
print("Training set size:", len(train_images))
print("Validation set size:", len(val_images))
print("Testing set size:", len(test_images))
```

```
Training set size: 341
Validation set size: 86
Testing set size: 107
```

Pre-processing the input data for TensorFlow model training

```
In [6]: import tensorflow as tf
import numpy as np

# Convert images to NumPy arrays
train_images = np.array(train_images)
val_images = np.array(val_images)
test_images = np.array(test_images)

# Convert images to float32 and normalize
train_images_tensor = tf.convert_to_tensor(train_images.astype(np.float32) / 255.0)
val_images_tensor = tf.convert_to_tensor(val_images.astype(np.float32) / 255.0)
test_images_tensor = tf.convert_to_tensor(test_images.astype(np.float32) / 255.0)

# Convert labels to TensorFlow tensors
train_labels_tensor = tf.convert_to_tensor(train_labels)
val_labels_tensor = tf.convert_to_tensor(val_labels)
test_labels_tensor = tf.convert_to_tensor(test_labels)
```

Defining the CNN Model

```
In [7]: import tensorflow as tf
from tensorflow.keras import layers

image_height = 1024
image_width = 768
num_channels = 3

# Define the CNN model
model = tf.keras.Sequential([
    layers.Conv2D(32, (3, 3), activation='relu', input_shape=(768, 1024, 3)),
    layers.MaxPooling2D((2, 2)),
    layers.Conv2D(64, (3, 3), activation='relu'),
    layers.MaxPooling2D((2, 2)),
    layers.Flatten(),
    layers.Dense(64, activation='relu'),
    layers.Dense(1, activation='sigmoid')
])
```

CNN model compilation for image classification

```
In [8]: model.compile(optimizer='adam',
                     loss='sparse_categorical_crossentropy',
                     metrics=['accuracy'])
```

Label Encoding and One-Hot Encoding for Classification Labels

```
In [10]: import numpy as np
import tensorflow as tf

actual_labels = ['3', '4', '5', '2', '3', '3', '3', '3', '5', '3', '1', '5', '3', '1', '4', '3', '1', '2', '3', '5']

# Create a dictionary to map the label names to integer values
label_to_integer = {label: i for i, label in enumerate(set(actual_labels))}

# Convert the actual labels to numerical values using the mapping
train_labels = np.array([label_to_integer[label] for label in actual_labels])

# Convert the numerical labels to one-hot encoded format
num_classes = len(label_to_integer)
train_labels_encoded = tf.one_hot(train_labels, num_classes)

In [12]: train_labels_encoded = tf.reshape(train_labels_encoded, (-1, num_classes))

In [13]: import numpy as np

num_classes = len(np.unique(train_labels))from collections import Counter

label_counts = Counter(train_labels)
num_classes = len(label_counts)

In [14]: num_classes = len(set(train_labels))

In [15]: from collections import Counter

label_counts = Counter(train_labels)
num_classes = len(label_counts)

In [16]: model.add(layers.Dense(534, activation='softmax'))
```

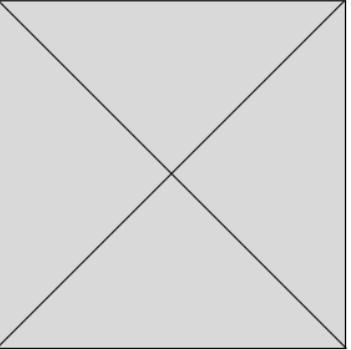
Training the model

```
In [*]: history = model.fit(train_images_tensor, train_labels_encoded, epochs=10, validation_data=(val_images_tensor, val_labels_tensor))  
Epoch 1/10
```



Wire Frames

Visual Insight for your Kidney Health



Lorem ipsum dolor sit amet, consectetur adipiscing elit. Duis facilisis laoreet nisi, vel dignissim quam interdum a. Fusce nec ipsum.

Continue

Instructions

1. Lorem ipsum dolor sit amet, consectetur adipiscing elit. Fusce tincidunt.

2. Lorem ipsum dolor sit amet, consectetur adipiscing elit. Fusce tincidunt.
3. Lorem ipsum dolor sit amet, consectetur adipiscing elit. Fusce tincidunt.
4. Lorem ipsum dolor sit amet, consectetur adipiscing elit. Fusce tincidunt.

Get Insights

Check progression

Get your insights

Select the image type

MRI

CT

Ultrasound

Choose one or more images



Continue

Get your insights

Select the image type

MRI

CT

Ultrasound

Choose one or more images



Continue

Your insights

Lorem ipsum dolor sit amet, consectetur adipiscing elit. Duis facilisis laoreet nisi, vel dignissim quam interdum a. Fusce nec ipsum.

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Lorem ipsum dolor sit amet, consectetur adipiscing elit. Fusce tincidunt.

Save as PDF

Let's Check your CKD progression

Lorem ipsum dolor sit amet, consectetur adipiscing elit. Fusce tincidunt.

Choose an image



Lorem ipsum dolor sit amet, consectetur adipiscing elit. Fusce tincidunt.

Check progression

Your CKD progression

Lorem ipsum dolor sit amet, consectetur adipiscing elit. Fusce tincidunt.



Lorem ipsum dolor sit amet, consectetur adipiscing elit. Fusce tincidunt.

References

- [1] Siddharth Rajput, “Automated Kidney Stone Detection Using Image Processing Techniques,” IEEE, 2021.
- [2] Ahmed Soliman, “Kidney segmentation from CT images using a 3D NMF-guided active contour model,” IEEE, 2021 .

IT20785192 | W.B.M.A. Isurika

Specializing in Data Science



Personalized Health Assessment and Management

Background

Important of Diet plan for CKD patients

- Slowing the Progression of CKD
- Managing Fluid Balance
- helps regulate fluid intake to prevent fluid overload, thereby reducing the strain on the kidneys and managing blood pressure.



Research Problem



- Despite the critical role of personalized diet management in slowing the progression of Chronic Kidney Disease (CKD) and improving patient outcomes.

- Currently they use one leaflet for all patients.

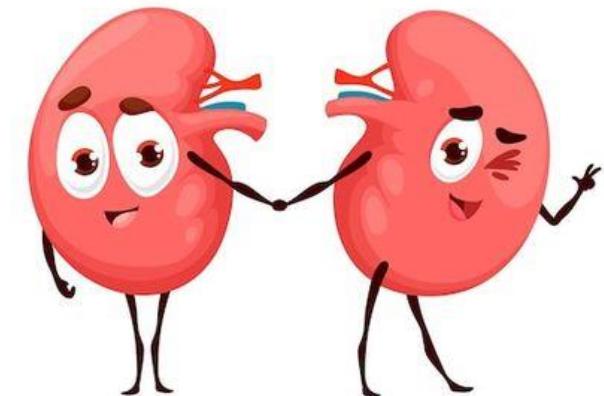
Research Gap

- Recent research has focused on Potassium level of blood to help manage the diet plans for CKD patients.
- Increase the accuracy of model and give proper diet plan
- Most studies have focused on short-term outcomes, such as changes in blood pressure or kidney function, rather than long-term outcomes such as financial side of patients.



What is Zone Attribute ?

- Existing dietary managements needs will be determined based on the ZONE Attribute
 - ✓ Safe - blood potassium level 3.5 - 5.0
 - ✓ Caution - blood potassium level 5.1 - 6.0
 - ✓ Danger - blood potassium level higher than 6.1



What tests they do ?

- "Serum Potassium Test" or "Potassium Blood Test"

Objectives

Main Objective : provide personalized recommendations to kidney patients regarding the most appropriate diet, exercise, and lifestyle changes based on their health condition.

Sub Objectives :

- Based on the Potassium level of the blood, categorize the patient into three zones.
- Categorize and prepare proper diet plan

Completion of the Component

□ Requirements Analysis

- Collect data manually and create data set

- Attributes

ID	Serum Albumin (g/dL)
Gender	Potassium
Age	Calcium
Height(m)	Phosphorus
Weight(kg)	Sodium
Stage of CKD	Hemoglobin
Current Medications	Cholesterol



Data Set

AutoSave (● Off) Isurika W. B. M. A. it20785192

Diet Prediction Data 01

File Home Insert Page Layout Formulas Data Review View Automate Help

Cut Copy Paste Format Painter

Font: Calibri 11pt

Number: General

Styles: Conditional Formatting

Cells: Insert Delete Format

Editing: AutoSum Fill Clear Sort & Filter Find & Select

Analysis: Analyze Data Sensitivity

Sensitivity

M20

13.4

	A	B	C	D	E	F	G	H	I	J	K	L	M	N	O
1	ID	Gender	Age	Height(m)	Weight(kg)	Current Medications	Stage of CKD	Serum Albumin (g/dL)	Potassium	Calcium	Phosphorus	Sodium	Hemoglobin	Cholesterol	Zone
2	1	Male	47	1.2	51.7	FALSE	4	4.9	5.4	7	3.9	138.2	11.3	14	3
3	2	Male	42	1.5	73.4	TRUE	3	5.4	3.8	11	5.9	139.4	12.3	14.3	2
4	3	Male	38	1.2	53.7	FALSE	3	2.5	4.8	8.8	6.3	146.1	15.6	16.3	2
5	4	Male	86	1.5	70.3	TRUE	5	2.5	6.9	8.3	6.7	123.7	12.3	13.8	4
6	5	Male	67	1.4	73.4	FALSE	3	2.7	2.9	10.5	4.6	133.3	11.8	14.7	1
7	6	Male	43	1.3	66	FALSE	3	3.6	3.6	11.3	6.4	154.4	16	17.6	2
8	7	Male	75	1.6	73.2	TRUE	2	5	2.9	11.5	7	155.9	12	10.6	1
9	8	Male	69	1.5	78.4	FALSE	3	3	5.7	10.2	2.4	129.6	13.7	17.9	3
10	9	Male	59	1.3	66.5	FALSE	5	5.9	2.8	11.6	6.2	148.3	13.6	17.3	1
11	10	Male	80	1.5	49.7	TRUE	5	2.7	3.5	11.6	6.3	120.3	13.6	14.8	2
12	11	Male	61	1.3	71.4	FALSE	3	3	7	8.9	6.3	155.1	13.5	16	4
13	12	Male	46	1.3	66	FALSE	1	2.5	4.4	11.8	5.9	129.8	10.6	13.1	2
14	13	Male	66	1.3	79.1	TRUE	3	5.9	3.3	6.5	3.2	157.8	17.3	13.3	1
15	14	Male	46	1.6	64	FALSE	3	4.5	3.9	6.6	6.3	153.9	10.9	16.2	2
16	15	Male	73	1.3	71.2	FALSE	3	3.2	6.2	8.1	6.9	158.8	18.2	12.4	4
511	510	Female	39	1.1	118.78	TRUE	3	3.1	3.7	7.4	6	137	10.7	13.1	2
512	511	Female	55	1	101.1	FALSE	1	4.2	3.4	11.7	5.7	149.4	13.6	16.7	1
513	512	Female	47	1.2	95.96	TRUE	1	3.5	6.6	7.4	6.3	128.5	10.6	12.8	4
514	513	Female	40	1.6	108.64	FALSE	2	5.1	5.7	10.9	6	157.1	13.5	17.8	3
515	514	Female	53	1.6	76.91	FALSE	4	5.2	2.4	10.2	2.2	152.5	12.6	11.7	1
516	515	Female	61	1.3	61.33	FALSE	3	4.5	4.2	9.2	3.5	133.1	13	15.6	2
517	516	Female	47	1.3	59.17	FALSE	2	5.4	5.6	8.7	5.4	130.8	10.7	11.4	3
518	517	Female	61	1.6	103.46	FALSE	2	3.1	4.6	11.1	5.8	124.1	11.3	13.8	2
519															

Import libraries

The screenshot shows a Jupyter Notebook interface with the following details:

- File Bar:** Untitled30.ipynb, Comment, Share, Settings, Help.
- Toolbar:** + Code, + Text, RAM Disk status (green checkmark).
- Sidebar:** Search (Q), Cell (x), File (□), Cell Counter (0s).
- Code Cells:**
 - #import libraries
 - import pandas as pd
 - import numpy as np
 - import matplotlib.pyplot as plt
 - import seaborn as sns
 - import numpy
 - from sklearn.preprocessing import MinMaxScaler
 - from sklearn import metrics
 - from sklearn import svm
 - from sklearn.ensemble import RandomForestClassifier
 - from sklearn.model_selection import train_test_split
 - from sklearn.metrics import accuracy_score, r2_score
 - import warnings
 - %matplotlib inline
 - warnings.filterwarnings('ignore')- # Assuming you have a dataset with 'X' as input features and 'y' as the target variable
- data = pd.read_csv('Diet Prediction Data 01.csv')
- data.head()

Data Preview:

ID	Gender	Age	Height(m)	Weight(kg)	Current Medications	Stage of CKD	Serum Albumin (g/dL)	Pottassium	Calcium	Phosphorus	Sodium	Himoglobin	Cholesterol	Zone	
0	1	Male	47	1.2	51.7	False	4.0	4.9	5.4	7.0	3.9	138.2	11.3	14.0	3
1	2	Male	42	1.5	73.4	True	3.0	5.4	3.8	11.0	5.9	139.4	12.3	14.3	2
2	3	Male	38	1.2	53.7	False	3.0	2.5	4.8	8.8	6.3	146.1	15.6	16.3	2
3	4	Male	86	1.5	70.3	True	5.0	2.5	6.9	8.3	6.7	123.7	12.3	13.8	4
4	5	Male	67	1.4	73.4	False	3.0	2.7	2.9	10.5	4.6	133.3	11.8	14.7	1

Data Preprocessing

```
[145] #To view the last few rows,  
data.tail()
```

ID	Gender	Age	Height(m)	Weight(kg)	Current Medications	Stage of CKD	Serum Albumin (g/dL)	Pottassium	Calcium	Phosphorus	Sodium	Himoglobin	Cholesterol	Zone	
721	722	Female	35	1.4	82.76	False	3.0	6.3	3.5	9.1	3.4	147.0	13.2	15.2	2
722	723	Female	29	1.0	67.30	False	1.0	4.6	5.0	9.2	6.9	125.1	13.5	10.9	2
723	724	Female	61	1.1	100.45	False	3.0	4.4	6.5	9.9	2.7	139.8	14.2	10.5	4
724	725	Female	31	1.2	110.13	False	3.0	3.4	2.5	10.0	5.7	147.5	13.5	16.8	1
725	726	Female	44	1.6	95.65	False	3.0	4.0	6.1	6.7	2.3	140.3	13.1	14.8	4

```
[146] #get the column headings of the data set  
data.columns.values
```

```
array(['ID', 'Gender', 'Age', 'Height(m)', 'Weight(kg)',  
       'Current Medications', 'Stage of CKD', 'Serum Albumin (g/dL)',  
       'Pottassium', 'Calcium', 'Phosphorus', 'Sodium', 'Himoglobin',  
       'Cholesterol', 'Zone'], dtype=object)
```

```
[146] #get the column headings of the data set  
data.columns.values
```

```
array(['ID', 'Gender', 'Age', 'Height(m)', 'Weight(kg)',  
       'Current Medications', 'Stage of CKD', 'Serum Albumin (g/dL)',  
       'Pottassium', 'Calcium', 'Phosphorus', 'Sodium', 'Himoglobin',  
       'Cholesterol', 'zone'], dtype=object)
```

```
[147] #Explore the data types of the columns  
data.dtypes
```

```
ID          int64  
Gender      object  
Age          int64  
Height(m)   float64  
Weight(kg)   float64  
Current Medications  object  
Stage of CKD    float64  
Serum Albumin (g/dL) float64  
Pottassium   float64  
Calcium      float64  
Phosphorus   float64  
Sodium       float64  
Himoglobin   float64  
Cholesterol  float64  
Zone          int64  
dtype: object
```

```

[148] #describe the data set
data.describe()

    ID      Age   Height(m)  Weight(kg) Stage of CKD  Serum Albumin (g/dL)  Potassium  Calcium  Phosphorus  Sodium  Hemoglobin  Cholesterol  Zone
count  726.000000  726.000000  726.000000  726.000000  716.000000  726.000000  726.000000  726.000000  726.000000  726.000000  720.000000  726.000000
mean   363.500000  57.071625  1.344766  75.338802  3.001397   4.442287  4.770386  9.230139  4.429063  140.465840  13.429477  14.487361  2.442149
std    209.722436  17.026562  0.157983  19.197597  1.097358   1.177497  1.306648  1.587852  1.487213  11.533203  2.135800  2.254686  1.035288
min    1.000000  28.000000  1.000000  48.000000  1.000000   2.400000  2.400000  6.500000  1.900000  120.100000  10.500000  10.500000  1.000000
25%   182.250000  42.000000  1.200000  60.100000  2.750000   3.500000  3.625000  7.900000  3.100000  130.425000  11.700000  12.600000  2.000000
50%   363.500000  57.000000  1.300000  72.200000  3.000000   4.400000  4.800000  9.300000  4.400000  141.400000  13.100000  14.500000  2.000000
75%   544.750000  72.000000  1.500000  88.152500  3.000000   5.400000  5.900000  10.600000  5.700000  150.100000  14.500000  16.325000  3.000000
max   726.000000  87.000000  1.600000  119.690000  5.000000   6.500000  7.000000  12.000000  7.000000  160.000000  18.500000  18.500000  4.000000

[149] #To get an idea about the null values
data.info()

<class 'pandas.core.frame.DataFrame'
RangeIndex: 726 entries, 0 to 725
Data columns (total 15 columns):
 #   Column          Non-Null Count  Dtype  
--- 
 0   ID              726 non-null    int64  
 1   Gender          726 non-null    object  
 2   Age             726 non-null    int64  
 3   Height(m)       726 non-null    float64 
 4   Weight(kg)      726 non-null    float64 
 5   Current Medications 784 non-null  object  
 6   Stage of CKD    716 non-null    float64 
 7   Serum Albumin (g/dL) 726 non-null  float64 
 8   Potassium       726 non-null    float64 
 9   Calcium          720 non-null    float64 
 10  Phosphorus       726 non-null    float64 
 11  Sodium           726 non-null    float64 
 12  Hemoglobin       726 non-null    float64 
 13  Cholesterol     720 non-null    float64 
 14  Zone             726 non-null    int64  
dtypes: float64(10), int64(3), object(2)

```

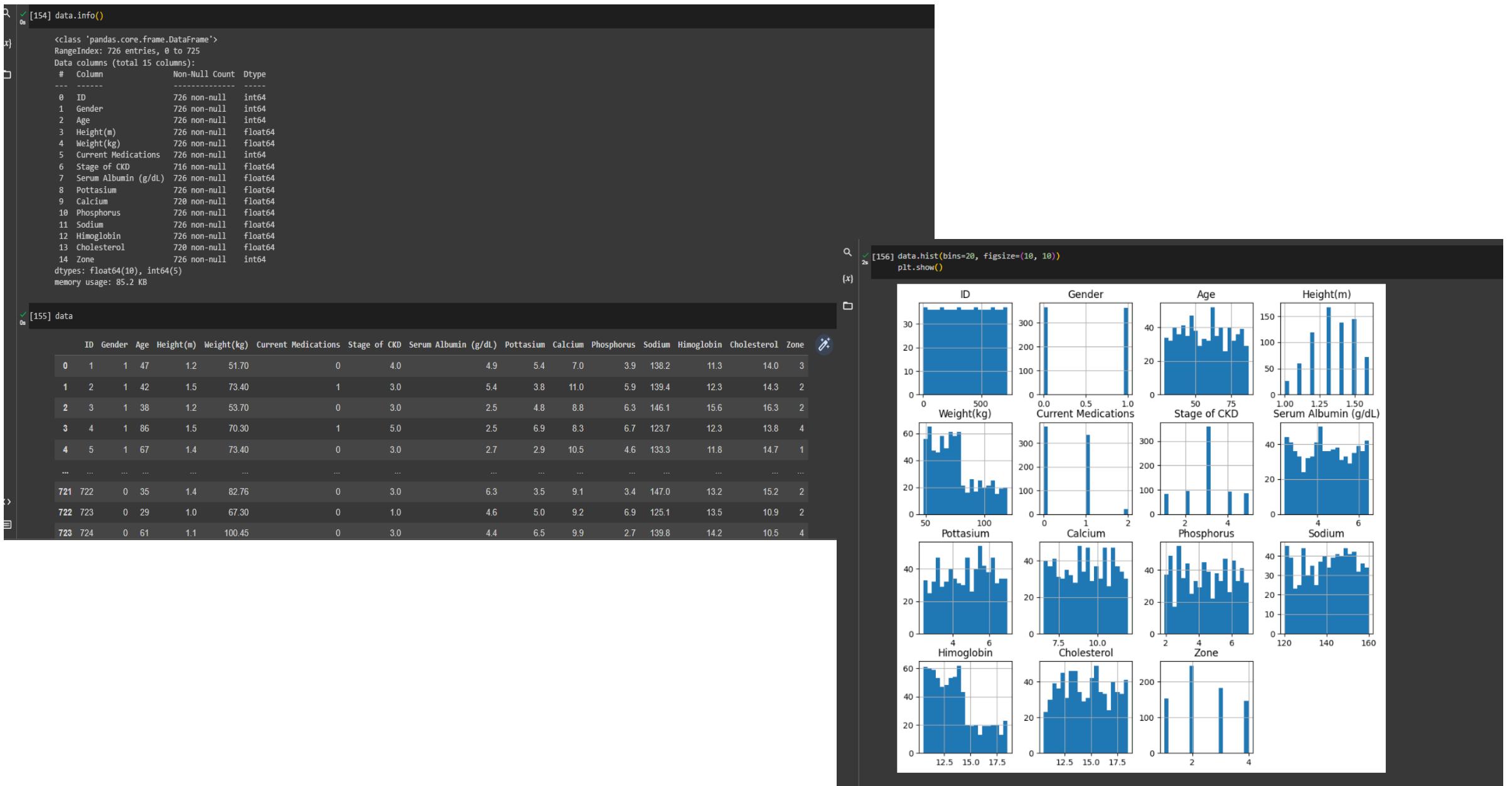
[150] #Getting the null value count
data.isnull().sum()

	ID	Gender	Age	Height(m)	Weight(kg)	Current Medications	Stage of CKD	Serum Albumin (g/dL)	Potassium	Calcium	Phosphorus	Sodium	Hemoglobin	Cholesterol	Zone
	0	0	0	0	0	22	10	0	0	6	0	0	0	6	0

[151] #replace null values using mean and value counts
Df_dataset = data.apply(lambda x: x.fillna(x.mean()))
if x.dtype == 'float'
else x.fillna(x.value_counts().index[0])

[152] from sklearn.preprocessing import LabelEncoder

Convert categorical variable using LabelEncoder
label_encoder = LabelEncoder()
data['Gender'] = label_encoder.fit_transform(data['Gender'])
data['Current Medications'] = label_encoder.fit_transform(data['Current Medications'])



```
[158] #split the dataset before balancing the class.
X = Df_dataset.drop(columns=[ 'Zone'], axis=1)
y = Df_dataset['Zone']

[159] X.columns.values
array(['ID', 'Gender', 'Age', 'Height(m)', 'Weight(kg)',
       'Current Medications', 'Stage of CKD', 'Serum Albumin (g/dL)',
       'Potassium', 'Calcium', 'Phosphorus', 'Sodium', 'Hemoglobin',
       'Cholesterol'], dtype=object)

[160] y.value_counts()
2    245
3    182
1    153
4    146
Name: Zone, dtype: int64

[161] #delete missing values rows
data = data.dropna(axis=0)

[163] # Split the data into train and test sets
X_train, X_test, y_train, y_test = train_test_split(data.drop("Zone", axis=1), data["Zone"], test_size=0.25)

# Create the random forest model
rf_model = RandomForestClassifier(n_estimators=100, max_depth=10)

# Fit the model to the training data
rf_model.fit(X_train, y_train)

# Predict the labels for the test data
y_pred = rf_model.predict(X_test)

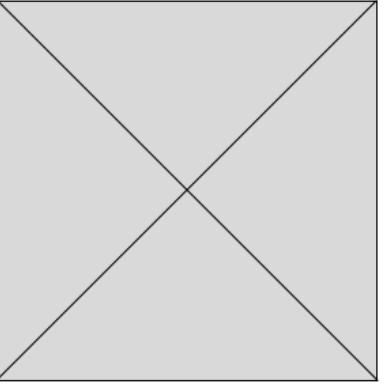
# Calculate the accuracy of the model
accuracy = accuracy_score(y_test, y_pred)
print("Accuracy:", accuracy)

# Calculate the R-squared value of the model
r2 = r2_score(y_test, y_pred)
print("R-squared:", r2)

Accuracy: 0.95
R-squared: 0.94269853783
```

Wireframes

Empowering knowledge nutrition for a healthier life



Lorem ipsum dolor sit amet, consectetur adipiscing elit. Duis facilisis laoreet nisi, vel dignissim quam interdum a. Fusce nec ipsum.

Continue

Embrace a nourishing journey towards kidney health and well-being

1 ————— 2 ————— 3

Select your gender

Female

Male

Select your age group

Below 18

18 - 30

30 - 60

Next

Do you have your Serum Potassium Test or Potassium Blood Test Reports with you ?

Yes

No

Height(m) _____

Weight (kg) _____

BMI

Any Current medications?

Yes

No

Next

Serium Albumin (g/DL) _____

Pottassium _____

Calcium _____

Phosphorus _____

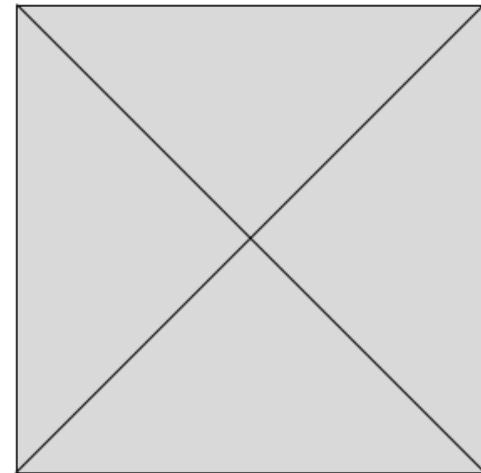
Sodium _____

Himoglobin _____

Check your results

Here's proper diet meal for you

*Lorem ipsum dolor sit amet,
consectetur adipiscing elit. Duis
facilisis laoreet nisi, vel dignissim
quam interdum a. Fusce nec
ipsum.*



What we plan to do next ?

- Plan to get results accuracies for algorithms

Used Algorithm	Overall Accuracy
Random Forest	95.54%
Logistic Regression	89.17%
Decision Trees	Plan to implement
Support Vector Machines (SVM)	Plan to implement
Naive Bayes	Plan to implement

- Implement ChatBot give proper Diet plan for patients

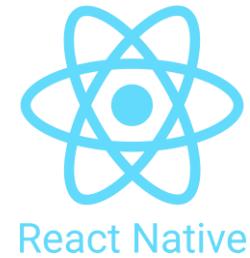


1. Natural Language Processing (NLP)

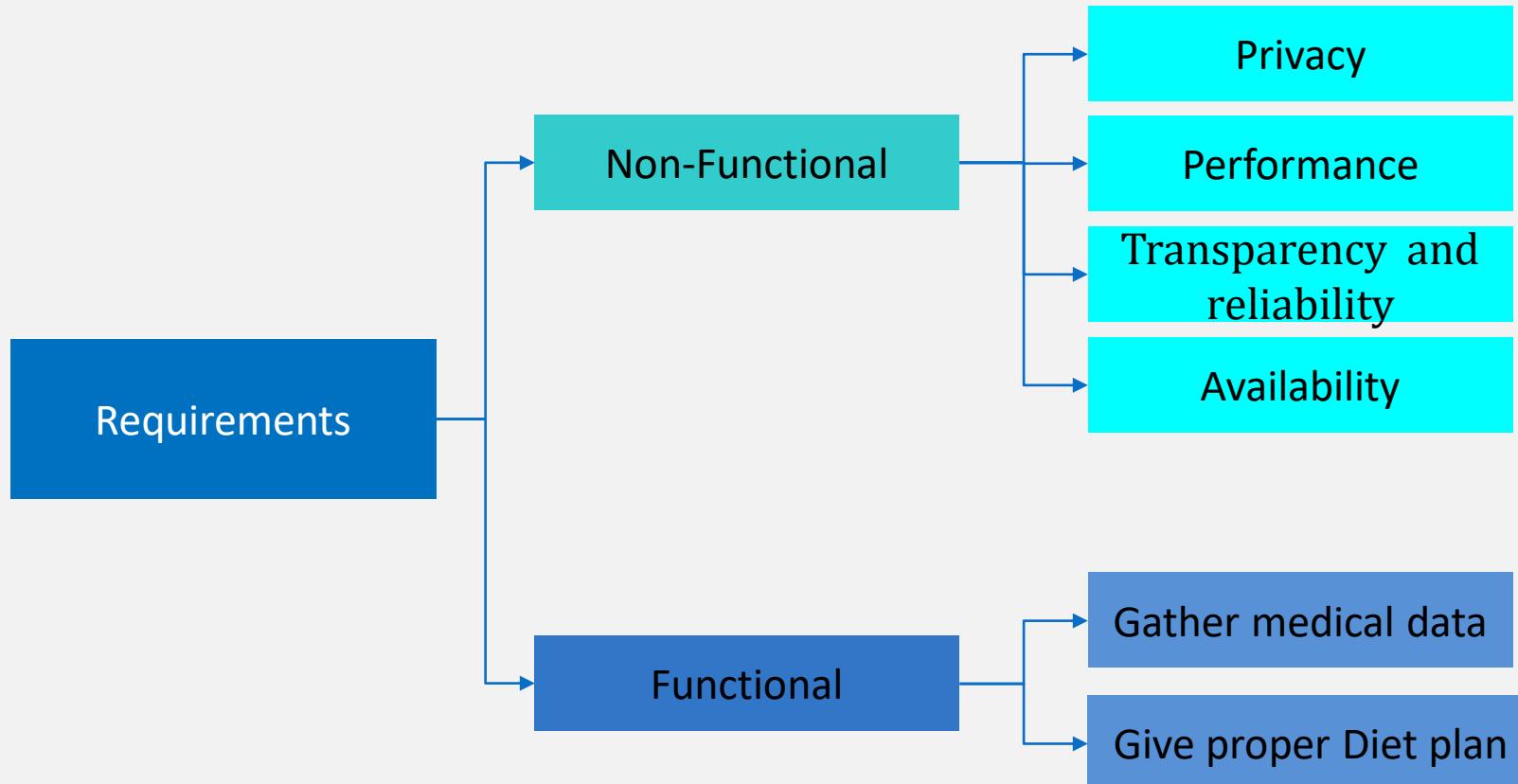
- Implementing the integration between the backend and frontend components of the mobile application.

Tools and Technologies

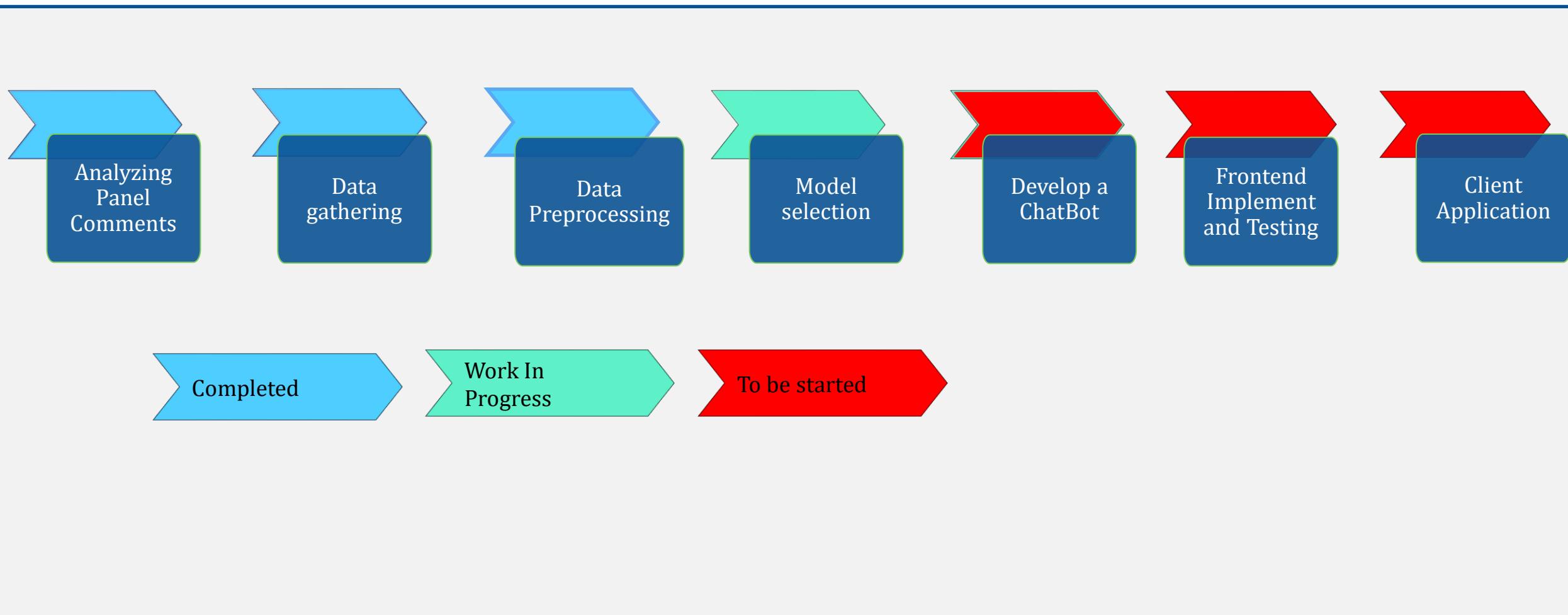
- Python
- Pandas , NumPy
- TensorFlow
- MongoDB
- AWS
- NodeJS
- React Native



Requirements



Overall Progress



References

- [10] N. Y. N., R. R. S. S. P. K. Annapoorna B. A, "Prediction of chronic kidney disease and diet recommendation," IJARIIT, 2021.
- [11] D. P. K. M.P.N.M. Wickramasinghe, "Dietary prediction for patients with Chronic Kidney Disease (CKD) by considering blood potassium level using machine learning algorithms," Research Gate, 2017.
- [12] National Kidney Foundation, " [Online]. Available: <https://www.kidney.org/kidneydisease/howkidneyswrk>.

IT20226596 | J.P.M.L. Perera

Specializing in Data Science



IoT & Machine Learning based Water quality Monitoring System for kidney patients

Background

- In Sri Lanka, prevalence of Kidney diseases (CKDu) have increased rapidly in recent decades.
- Most affected people are living in rural areas, and they have limited access to modern health care.
- Poor water quality has been identified as a possible contributor to kidney diseases.
- It is important to being aware of whether the daily used drinking water is harmful to kidneys or not.

Research Problem

- Lack of reliable and efficient real-time water quality monitoring system in Sri Lanka.
- Inadequate identification of sudden changes in water quality, impacting kidney patients' health.
- Absence of proper data management systems for recording, analyzing, and responding to water quality issues.
- Limited access to and understanding of water quality particularly for non-English speaking kidney patients.



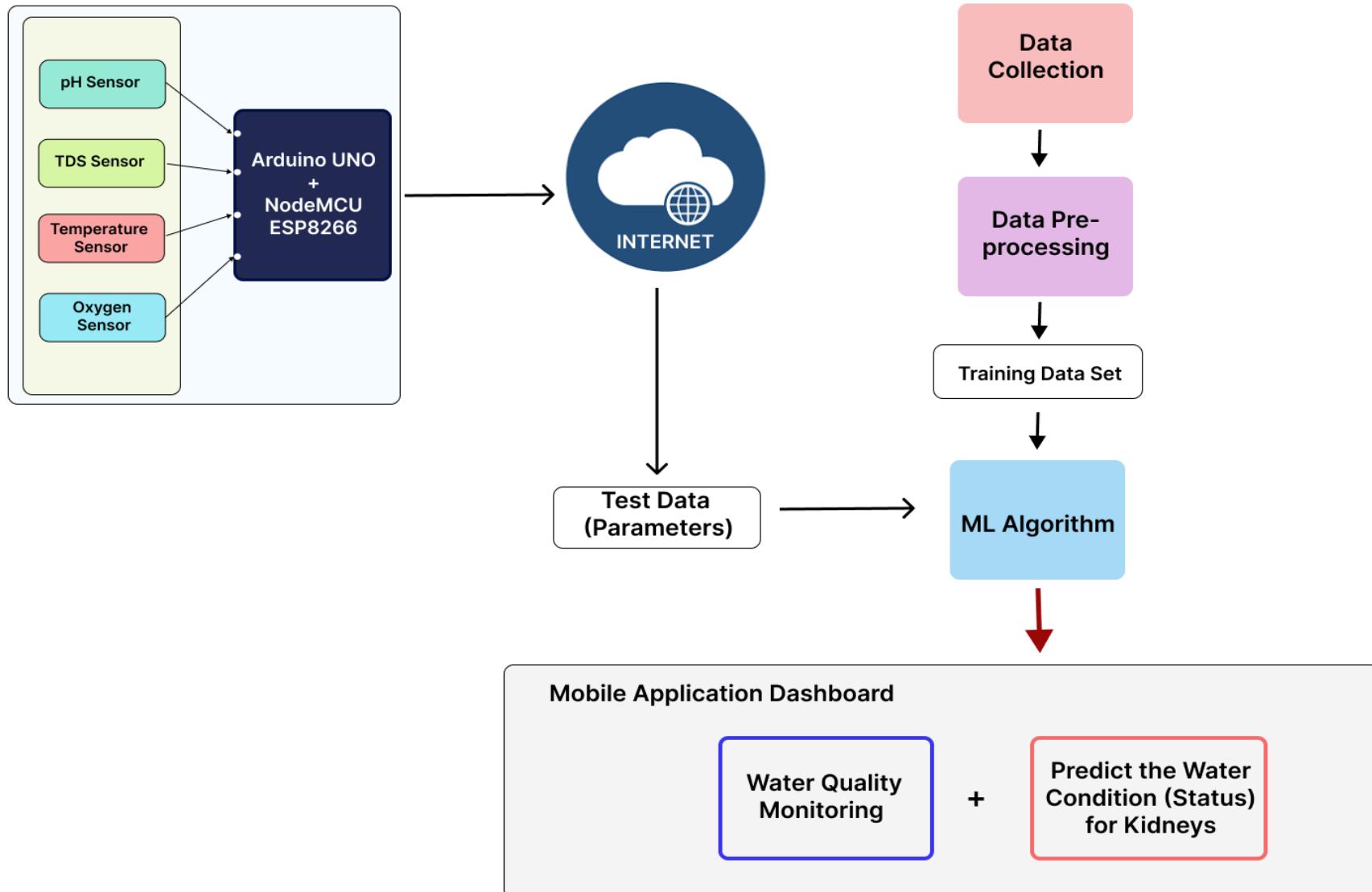
Research Gap

- The lack of a reliable and efficient real-time water quality monitoring system using IoT technology in Sri Lanka. Existing manual monitoring methods are unable to detect sudden changes in water quality.
- The absence of proper data management systems for recording, analyzing, and responding to water quality issues in Sri Lanka.
- Limited access to and understanding of water quality information, particularly for non-English speaking kidney patients in Sri Lanka.
- Insufficient integration of IoT and ML technologies for water quality monitoring and analysis in the context of Sri Lanka.

Novelty

- Implementing IoT technology specifically designed for monitoring water quality in Sri Lanka, which is currently absent or limited.
- Applying machine learning (ML) algorithms to analyze the collected water quality data and identify patterns, trends, and anomalies.
- Providing multilingual support to ensure that water quality information is available in local languages, improving accessibility and understanding for all users.
- Integrating IoT sensors and devices with ML algorithms to create a comprehensive solution for real-time monitoring, analysis, and management of water quality.

Component Overview



Completion of the Component

Hardware Solution – 50%

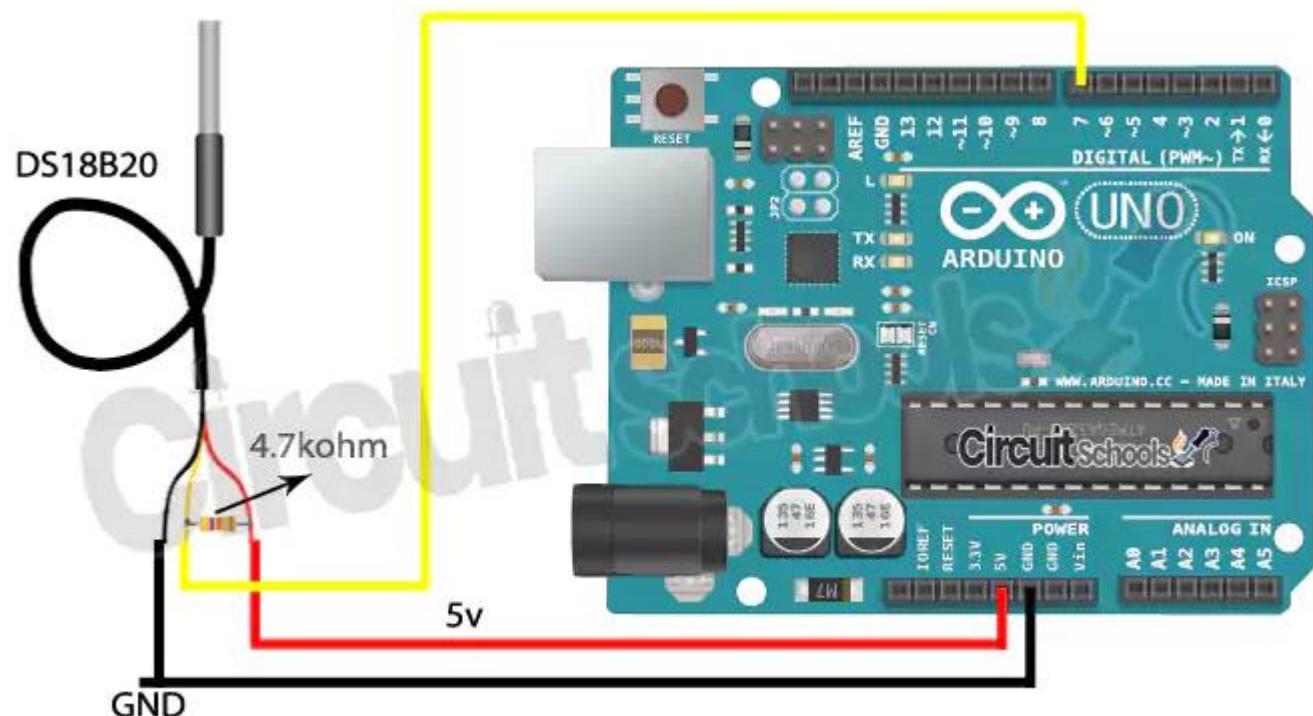
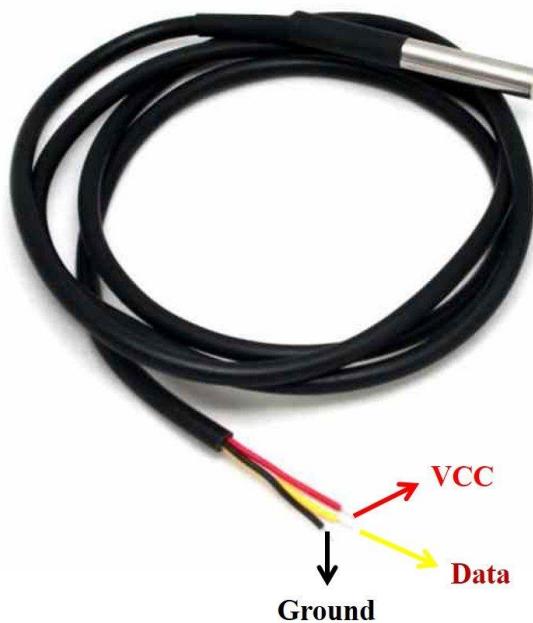
- **Gathering Hardware Components**
 - Arduino Uno Board
 - Sensors
 - Resistors
 - Jump wires
- **Design the Circuits**
- **Programming the Arduino Board for test the Sensors**
- **Get values Separately from Sensors**

Implementation

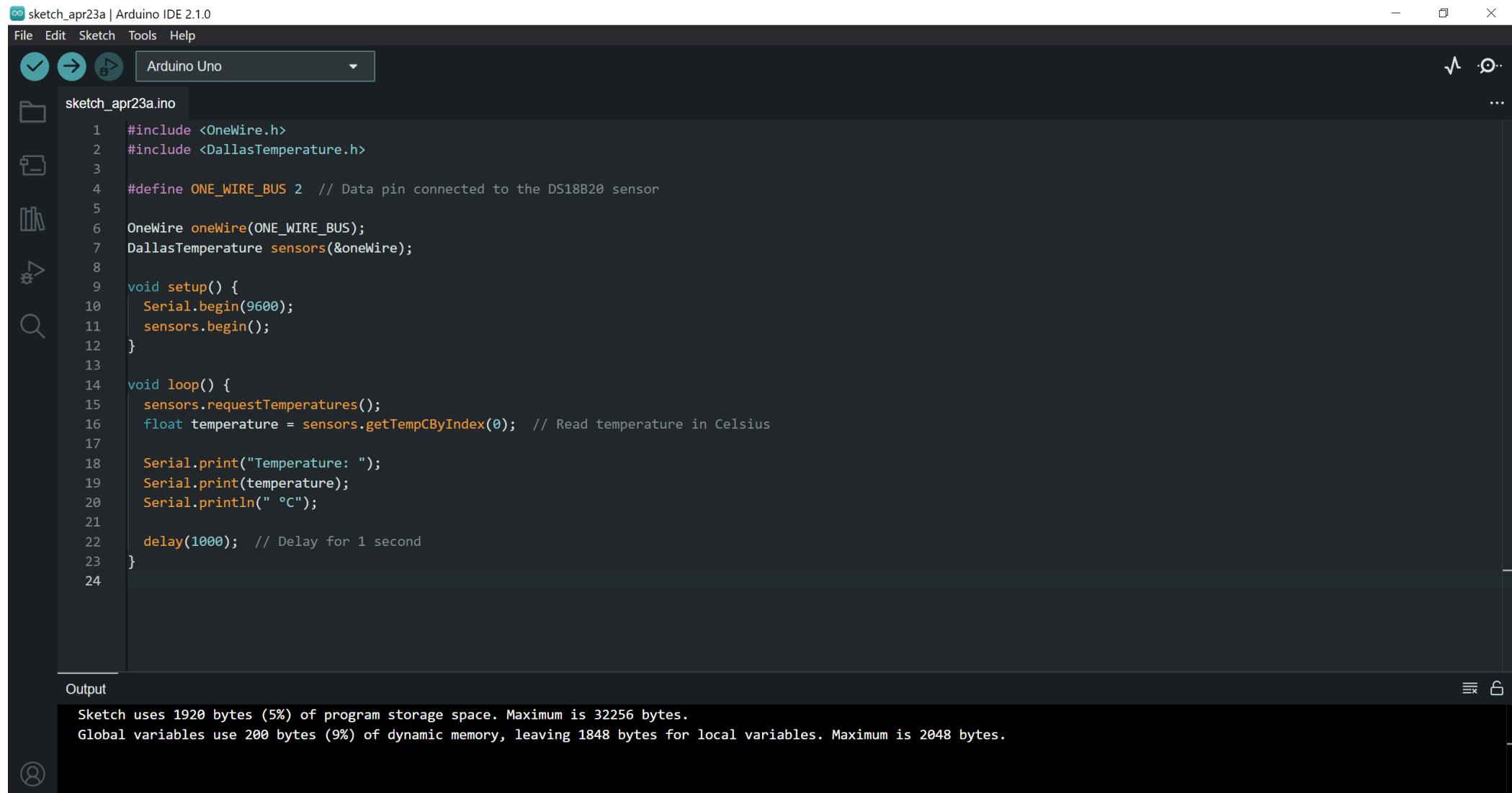
Hardware Solution

DS18B20 Waterproof Temperature Sensor

Circuit Diagram



Config Temperature sensor in Arduino Uno



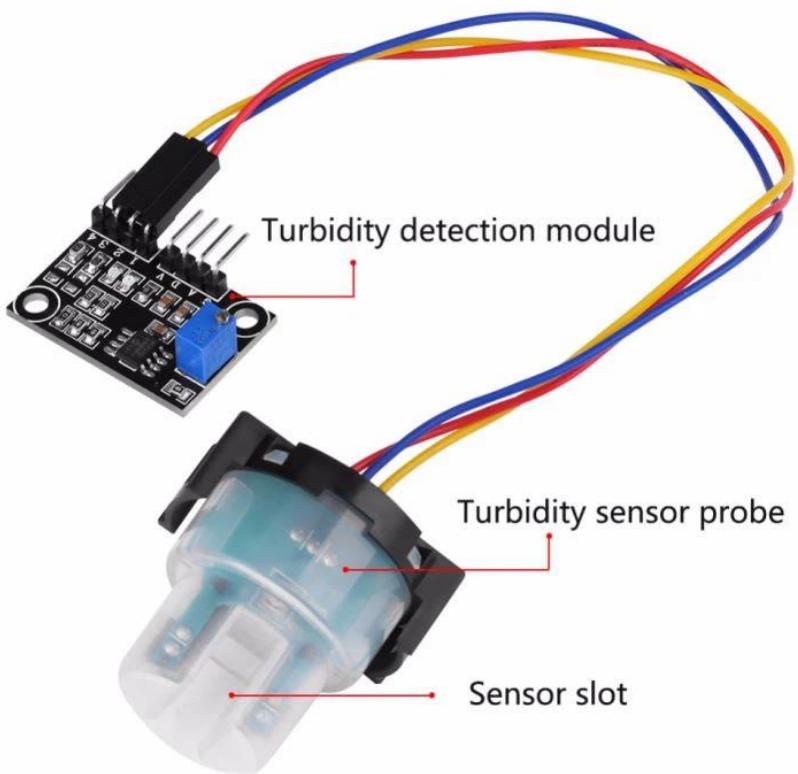
The screenshot shows the Arduino IDE 2.1.0 interface with a dark theme. The title bar reads "sketch_apr23a | Arduino IDE 2.1.0". The menu bar includes File, Edit, Sketch, Tools, and Help. The toolbar has icons for save, build, upload, and refresh. The board selector is set to "Arduino Uno". The left sidebar contains icons for file, folder, and search. The main code editor window displays "sketch_apr23a.ino" with the following code:

```
1 #include <OneWire.h>
2 #include <DallasTemperature.h>
3
4 #define ONE_WIRE_BUS 2 // Data pin connected to the DS18B20 sensor
5
6 OneWire oneWire(ONE_WIRE_BUS);
7 DallasTemperature sensors(&oneWire);
8
9 void setup() {
10   Serial.begin(9600);
11   sensors.begin();
12 }
13
14 void loop() {
15   sensors.requestTemperatures();
16   float temperature = sensors.getTempCByIndex(0); // Read temperature in Celsius
17
18   Serial.print("Temperature: ");
19   Serial.print(temperature);
20   Serial.println(" °C");
21
22   delay(1000); // Delay for 1 second
23 }
```

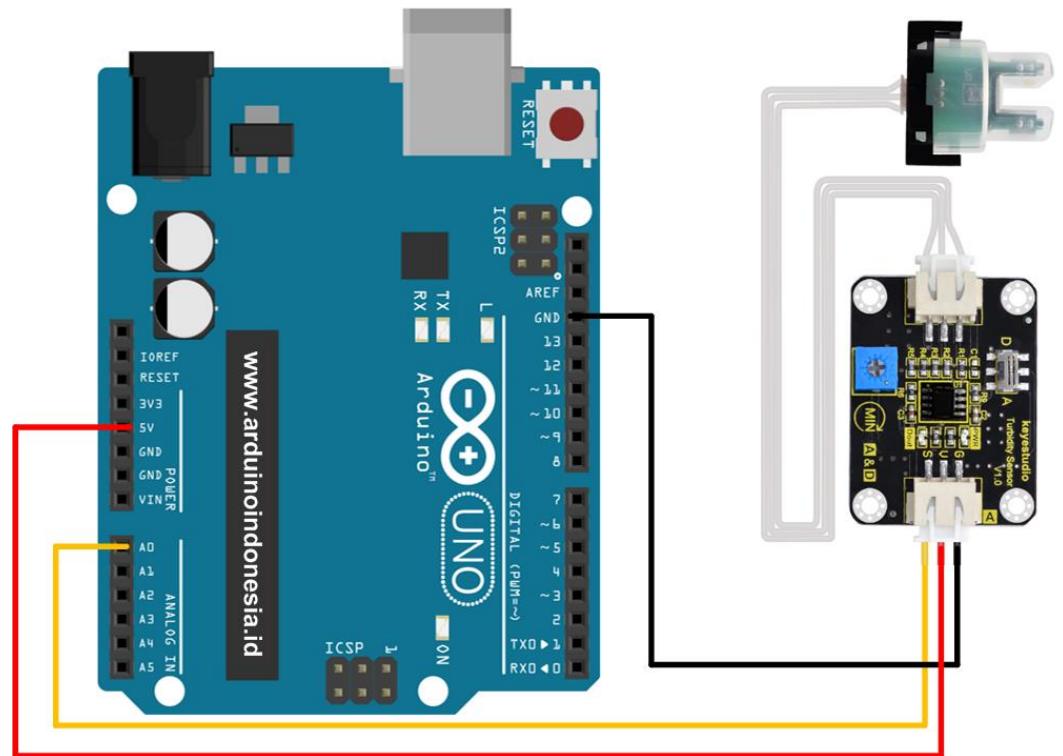
The bottom output window shows memory usage information:

```
Sketch uses 1920 bytes (5%) of program storage space. Maximum is 32256 bytes.
Global variables use 200 bytes (9%) of dynamic memory, leaving 1848 bytes for local variables. Maximum is 2048 bytes.
```

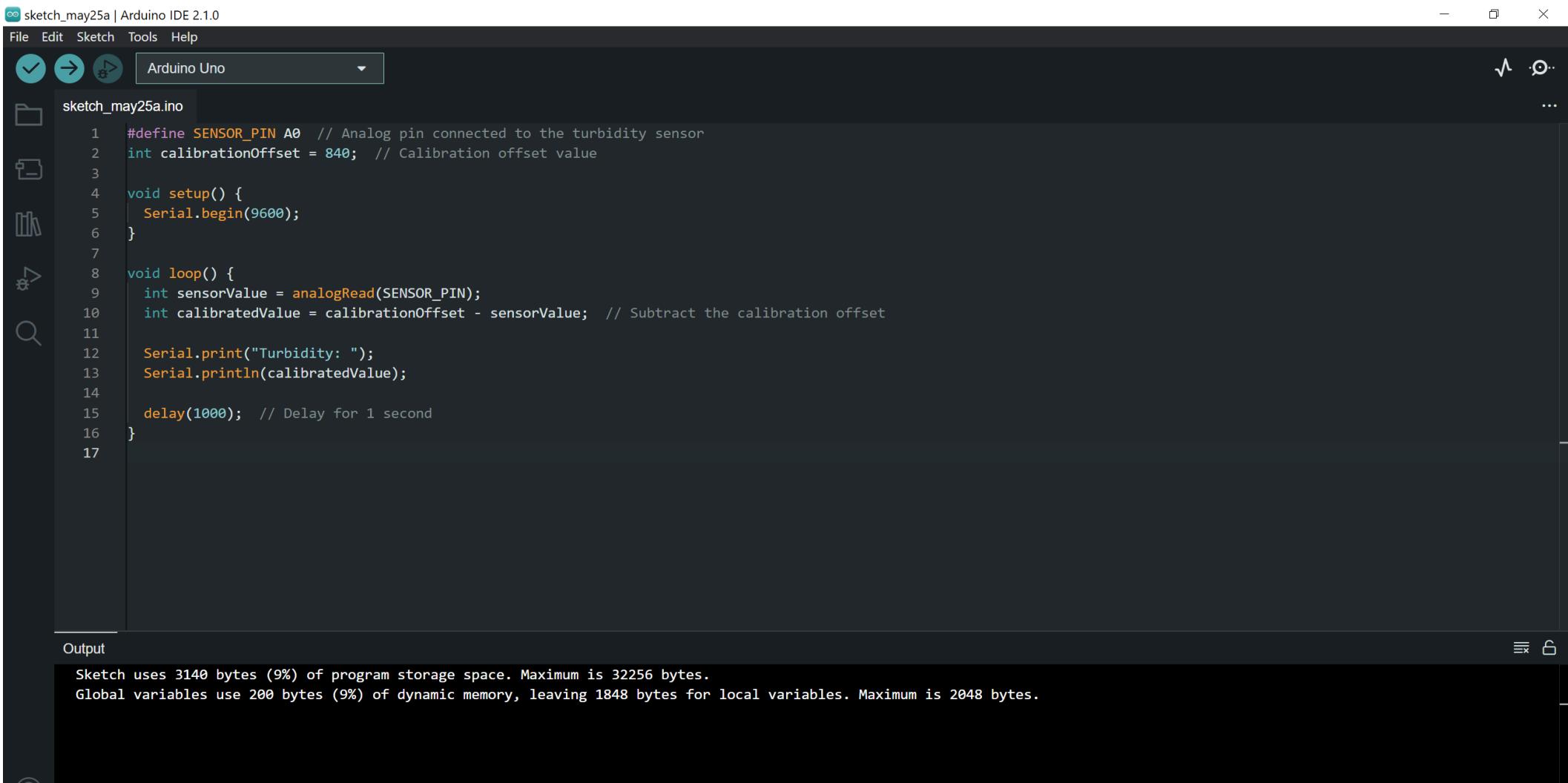
SENO189 Analog Turbidity Sensor



Circuit Diagram



Config Turbidity sensor in Arduino Uno



The screenshot shows the Arduino IDE interface with a dark theme. The title bar reads "sketch_may25a | Arduino IDE 2.1.0". The menu bar includes File, Edit, Sketch, Tools, and Help. The toolbar has icons for save, upload, and refresh. The board selector dropdown is set to "Arduino Uno". The left sidebar shows a file tree with "sketch_may25a.ino" selected. The code editor contains the following sketch:

```
1 #define SENSOR_PIN A0 // Analog pin connected to the turbidity sensor
2 int calibrationOffset = 840; // Calibration offset value
3
4 void setup() {
5     Serial.begin(9600);
6 }
7
8 void loop() {
9     int sensorValue = analogRead(SENSOR_PIN);
10    int calibratedValue = calibrationOffset - sensorValue; // Subtract the calibration offset
11
12    Serial.print("Turbidity: ");
13    Serial.println(calibratedValue);
14
15    delay(1000); // Delay for 1 second
16 }
17
```

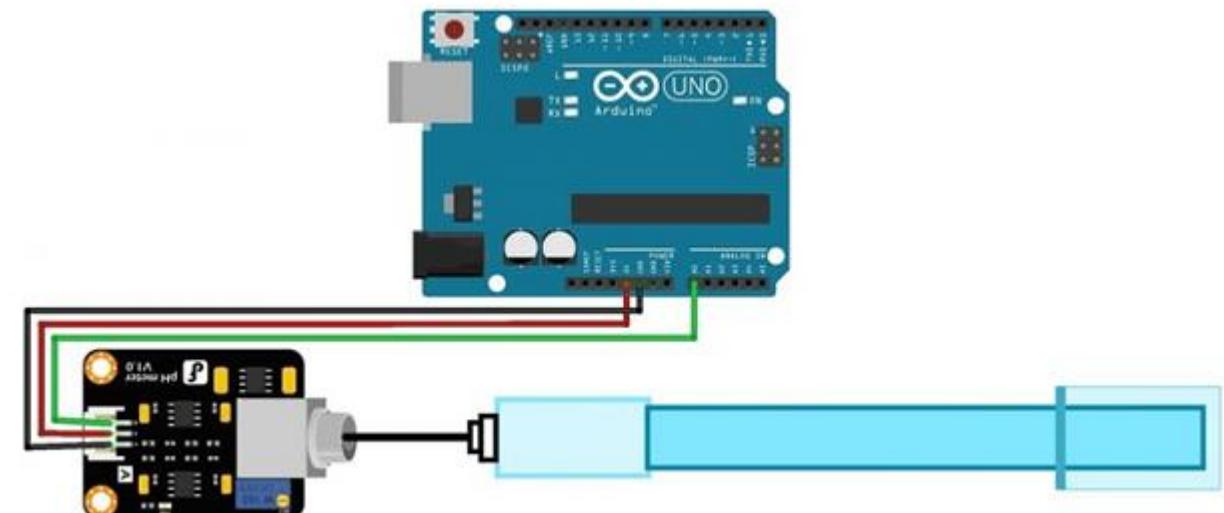
The bottom output window displays memory usage information:

```
Sketch uses 3140 bytes (9%) of program storage space. Maximum is 32256 bytes.
Global variables use 200 bytes (9%) of dynamic memory, leaving 1848 bytes for local variables. Maximum is 2048 bytes.
```

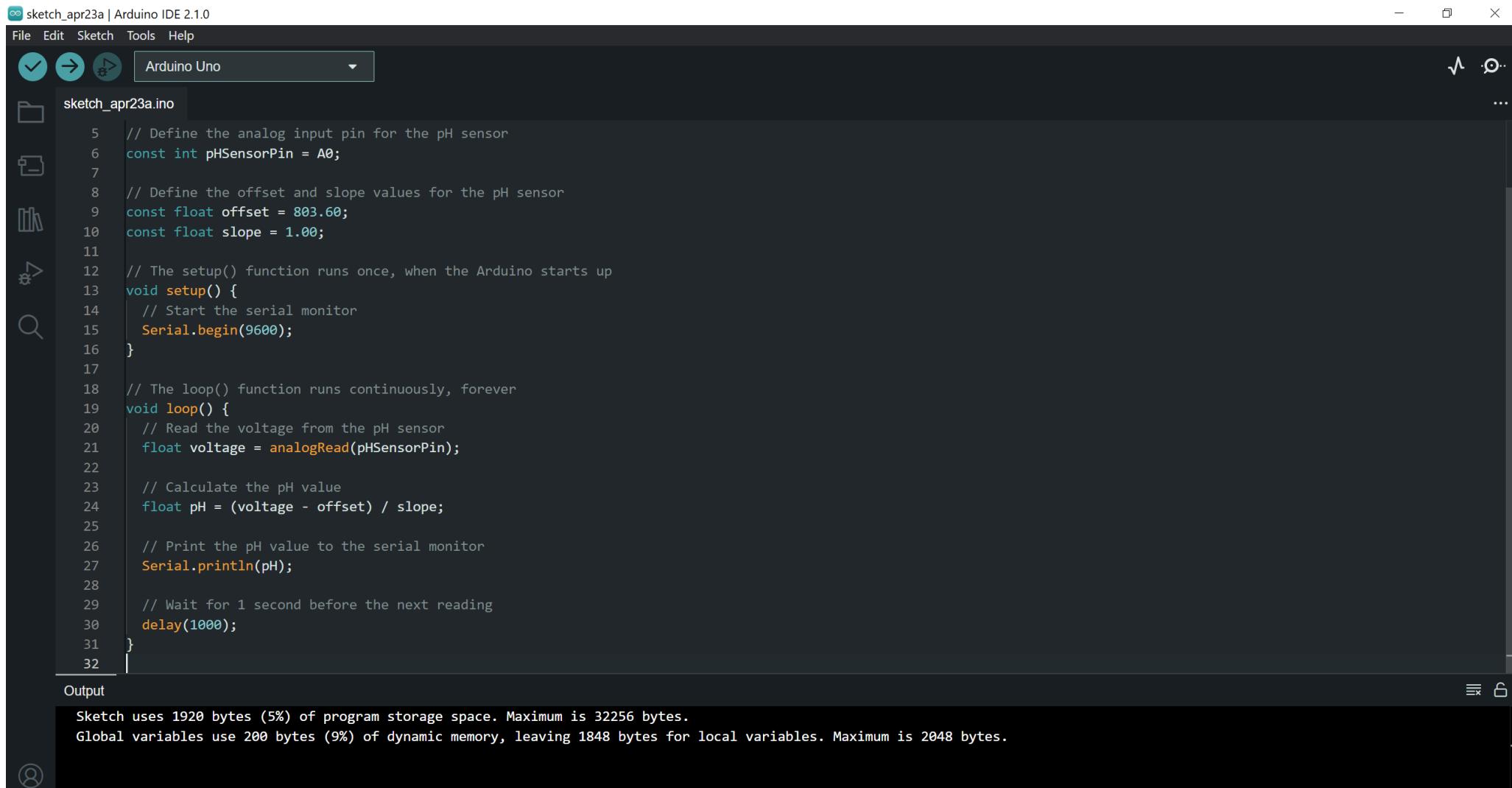
Analog pH Sensor with probe



Circuit Diagram



Config pH sensor in Arduino Uno



The screenshot shows the Arduino IDE interface with the following details:

- Title Bar:** sketch_apr23a | Arduino IDE 2.1.0
- Menu Bar:** File, Edit, Sketch, Tools, Help
- Toolbar:** Includes icons for save, upload, and refresh.
- Sketch Selection:** Arduino Uno
- Code Editor:** Displays the `sketch_apr23a.ino` file content. The code configures an analog input pin A0 for a pH sensor, defines offset and slope values, and implements setup() and loop() functions to read the voltage, calculate pH, and print it to the serial monitor with a 1-second delay between readings.

```
sketch_apr23a.ino
5 // Define the analog input pin for the pH sensor
6 const int pHsensorPin = A0;
7
8 // Define the offset and slope values for the pH sensor
9 const float offset = 803.60;
10 const float slope = 1.00;
11
12 // The setup() function runs once, when the Arduino starts up
13 void setup() {
14     // Start the serial monitor
15     Serial.begin(9600);
16 }
17
18 // The loop() function runs continuously, forever
19 void loop() {
20     // Read the voltage from the pH sensor
21     float voltage = analogRead(pHsensorPin);
22
23     // Calculate the pH value
24     float pH = (voltage - offset) / slope;
25
26     // Print the pH value to the serial monitor
27     Serial.println(pH);
28
29     // Wait for 1 second before the next reading
30     delay(1000);
31 }
32
```

- Output Panel:** Shows memory usage statistics:
 - Sketch uses 1920 bytes (5%) of program storage space. Maximum is 32256 bytes.
 - Global variables use 200 bytes (9%) of dynamic memory, leaving 1848 bytes for local variables. Maximum is 2048 bytes.
- Status Bar:** Shows a circular icon with the number 8.

Completion of the Component

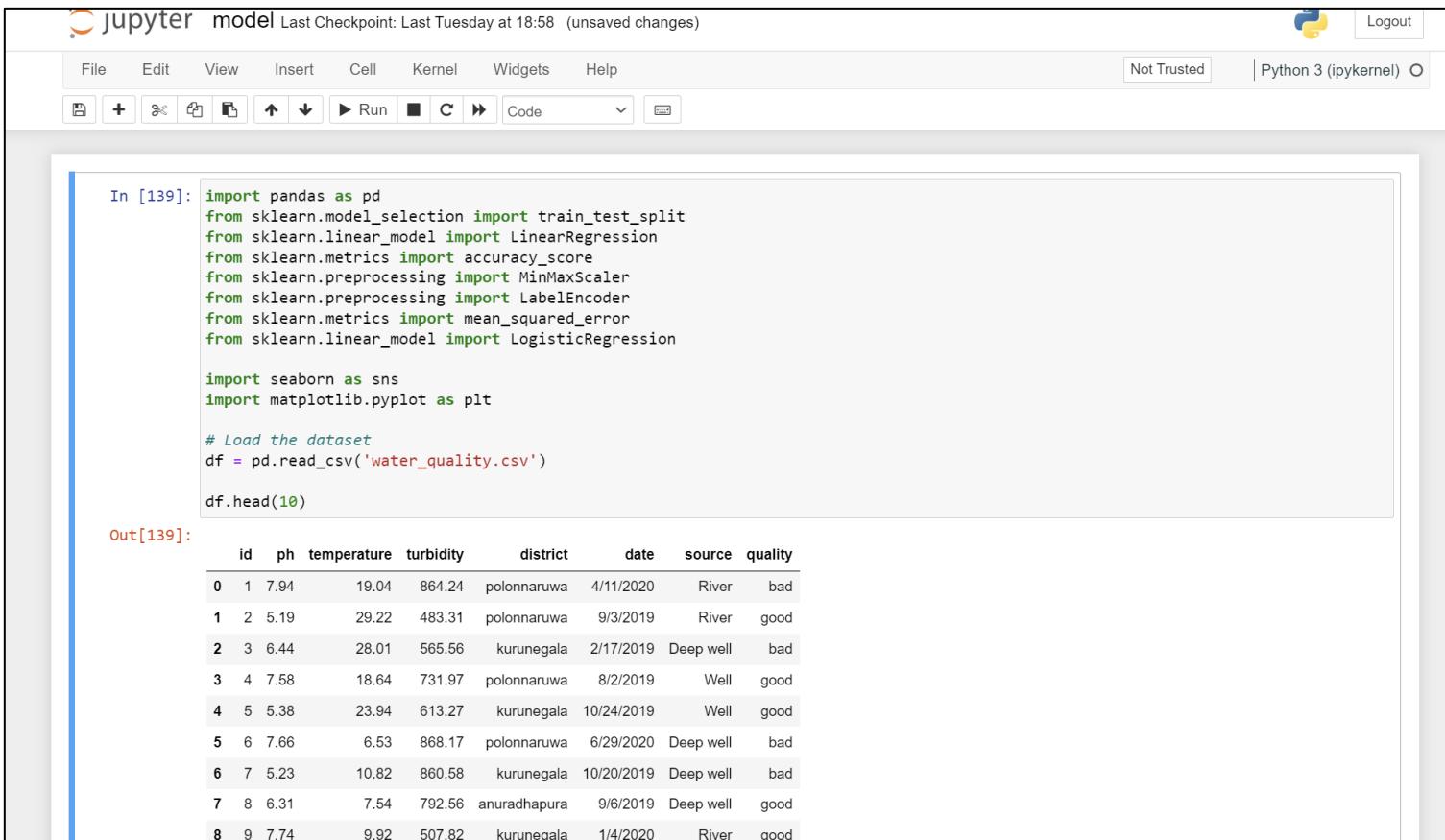
Software Solution – 50%

- **Gathering Dataset from NIFS Kandy**
- **Analyse the Data set**
- **Pre-process the Data Set**
- **Build the Models and Train the Model**
- **Get the Accuracy**

Implementation

Machine Learning (Software) Solution

- Import Libraries & Load the Data set into Jupyter Notebook



The screenshot shows a Jupyter Notebook interface with the following details:

- Title Bar:** jupyter model Last Checkpoint: Last Tuesday at 18:58 (unsaved changes)
- Toolbar:** File, Edit, View, Insert, Cell, Kernel, Widgets, Help, Run, Stop, Kernel, Code, Help.
- Status Bar:** Not Trusted | Python 3 (ipykernel)
- In [139]:** Contains Python code for importing various machine learning and visualization libraries, and loading a dataset from 'water_quality.csv'.
- Out[139]:** Displays the first 10 rows of the loaded dataset as a Pandas DataFrame.

	id	ph	temperature	turbidity	district	date	source	quality
0	1	7.94	19.04	864.24	polonnaruwa	4/11/2020	River	bad
1	2	5.19	29.22	483.31	polonnaruwa	9/3/2019	River	good
2	3	6.44	28.01	565.56	kurunegala	2/17/2019	Deep well	bad
3	4	7.58	18.64	731.97	polonnaruwa	8/2/2019	Well	good
4	5	5.38	23.94	613.27	kurunegala	10/24/2019	Well	good
5	6	7.66	6.53	868.17	polonnaruwa	6/29/2020	Deep well	bad
6	7	5.23	10.82	860.58	kurunegala	10/20/2019	Deep well	bad
7	8	6.31	7.54	792.56	anuradhapura	9/6/2019	Deep well	good
8	9	7.74	9.92	507.82	kurunegala	1/4/2020	River	good

- **Checking Missing values & Data types**

The screenshot shows a Jupyter Notebook interface with the following details:

- Header:** jupyter model Last Checkpoint: Last Tuesday at 18:58 (unsaved changes), Python 3 (ipykernel)
- Toolbar:** File, Edit, View, Insert, Cell, Kernel, Widgets, Help, Run, Code
- Data Preview:** A preview of a DataFrame with columns: id, ph, temperature, turbidity, district, date, source, quality. Rows 7, 8, and 9 are shown.
- In [140]:** `# Calculate the missing values in each column
missing_values = df.isnull().sum()`
- In [141]:** `print(missing_values)`
Output:
id 0
ph 0
temperature 0
turbidity 0
district 0
date 0
source 0
quality 0
dtype: int64
- In [142]:** `print(df.dtypes)`
Output:
id int64
ph float64
temperature float64
turbidity float64
district object
date object
source object
quality object
dtype: object

- **Transform the Data types using Label Encoding**

The screenshot shows a Jupyter Notebook interface with the following content:

```
In [143]: # Create a LabelEncoder object
label_encoder = LabelEncoder()

# Encode the "gender" column
df['district_encoded'] = label_encoder.fit_transform(df['district'])
df['source_encoded'] = label_encoder.fit_transform(df['source'])

print(df.dtypes)
```

Output of In [143]:

```
id          int64
ph         float64
temperature  float64
turbidity   float64
district    object
date        object
source      object
quality     object
district_encoded  int32
source_encoded  int32
dtype: object
```

```
In [144]: # Drop specific columns
columns_to_drop = ['date','district'] # Replace with the actual column names you want to drop
df = df.drop(columns_to_drop, axis=1)
df.head()
```

Output of In [144]:

	id	ph	temperature	turbidity	source	quality	district_encoded	source_encoded
0	1	7.94	19.04	864.24	River	bad	2	1
1	2	5.19	29.22	483.31	River	good	2	1
2	3	6.44	28.01	565.56	Deep well	bad	1	0

- **Update the target variable and Split the data set into Train & Test datasets**

The screenshot shows a Jupyter Notebook interface with the title "jupyter model" and a status bar indicating "Last Checkpoint: Last Tuesday at 18:58 (unsaved changes)". The toolbar includes File, Edit, View, Insert, Cell, Kernel, Widgets, Help, Not Trusted, and Python 3 (ipykernel). Below the toolbar is a toolbar with various icons for file operations and cell execution.

The main area contains three code cells:

```
In [149]: X = df.drop('quality', axis=1) # Update 'target_variable' with the actual column name  
y = df['quality']
```

```
In [150]: print(df.dtypes)
```

Column	Dtype
ph	float64
temperature	float64
turbidity	float64
quality	object
district_encoded	int32
source_encoded	int32
dtype:	object

```
In [151]: X_train, X_test, y_train, y_test = train_test_split(X, y, test_size=0.2, random_state=0)
```

- Build the model, Train the dataset & Get the accuracy

- Logistic Regression

```
In [154]: X_train, X_test, y_train, y_test = train_test_split(X, y, test_size=0.2, random_state=42)

# Create a logistic regression model
model = LogisticRegression()

# Fit the model on the training data
model.fit(X_train, y_train)

# Predict the target variable for the test set
y_pred = model.predict(X_test)

# Calculate the accuracy of the model
accuracy = accuracy_score(y_test, y_pred)

# Print the accuracy
print("Accuracy:", accuracy)

Accuracy: 0.5217391304347826
```

- Random Forest Classifier

```
In [155]: from sklearn.ensemble import RandomForestClassifier

# Split the data into training and test sets
X_train, X_test, y_train, y_test = train_test_split(X, y, test_size=0.2, random_state=42)

# Create a random forest classifier
model = RandomForestClassifier(n_estimators=100, random_state=42)

# Fit the model on the training data
model.fit(X_train, y_train)

# Predict the target variable for the test set
y_pred = model.predict(X_test)

# Calculate the accuracy of the model
accuracy = accuracy_score(y_test, y_pred)

# Print the accuracy
print("Accuracy:", accuracy)

Accuracy: 0.6521739130434783
```

What's Next ?

- Complete the IoT device
- Create the connectivity between device and mobile app
- Improve the ML model accuracy
- Give the Real time Analytical Quality Result of tested water sample through Our Mobile Application

References

- [1] D. P. K. K. M.P.N.M. Wickramasinghe, "Dietary prediction for patients with Chronic Kidney Disease (CKD) by considering blood potassium level using machine learning algorithms," Research Gate, 2017.
- [2] N. K. K. S. Moldobaeva Munara, "Recommending IoT based Real-time Water Quality Monitoring System in Malaysia," IEEE, 2022.
- [3] A. J. N. B. J. D. P. K. Shashika Lokuliyana, "A Survey: IoT Enable Framework for Water Quality Measurement and Distribution," IEEE, 2018.

Commercialization

- Partner with healthcare providers to offer our app as a resource for their patients.
- Collect data from app users to identify patterns and trends in kidney disease prevalence and risk factors. Use this data to improve the app's features and functionality and to develop targeted marketing campaigns.

2023-04-29 At Kurunegala Teaching Hospital | Consultant Nephrologist Dr. Premil Rajakrishnan


**SRI LANKA
INSTITUTE OF INFORMATION TECHNOLOGY**
16th Floor, BoC Merchant Tower, No. 2B, St. Michael's Road, Colombo 03

Date:28/04/2023 Your Ref : My Ref :2023-032

Dr. Premil Rajakrishna,
Consultant Nephrologist,
Teaching Hospital,
Kurunegala.

Dear Sir,

Certifying the project titled “KidniFY - A mobile based Chronic kidney Disease Patient care System Using ML and IoT” is conducting as a BSc in IT final year research project.

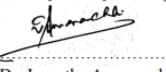
The Sri Lanka Institute of Information Technology (SLIIT) is the largest Degree Awarding Institute in the field of information Technology recognized by the University Grants Commission under the Universities Act. It was established in the year 1999 to educate and train Information Technology (IT) Professionals required by the fast-growing IT Industry in Sri Lanka.

This letter is to certify that the following students.
IT20154226 - M.M.K.L.Marasinghe
IT20226596 - J.P.M.L. Perera
IT20235260 - D.R.N. Samarawila
IT20785120 - W.B.M.A. Isurika

They are final year undergraduate students who conduct research entitled “KidniFY - A mobile based Chronic kidney Disease Patient care System Using ML and IoT” as partial fulfillment of the B.Sc. in Information Technology degree at Sri Lanka Institute of Information Technology (SLIIT). The students are conducting the research under the supervision of Ms. Wishalya Vanshanee Tissera

I kindly request your assistance in enabling these students to collect data from your organization to build their dataset for the research project. If you have any questions or require further clarification about the project, please do not hesitate to contact me.

Thank you for your cooperation



Dr. Jayantha Amararachchi
Assistant Professor/
Research Project Coordinator,
jayantha.a@sliit.lk
+94 11 754 4103

Tel: +94(0)11 2301904 - 5 Fax: +94(0)11 2301906 E-mail: info@sliit.lk URL: www.sliit.lk



2023-05-01 At Anuradhapura Teaching Hospital | Kidney Dialysis Unit



Thank You!

Q & A

