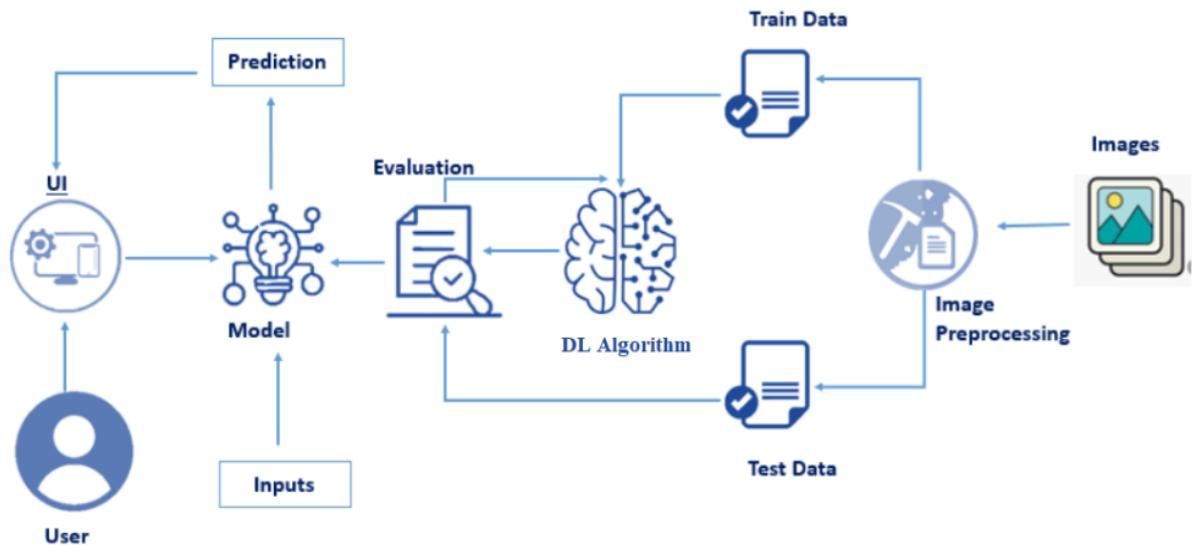


# Microbe Mapper: Visual Recognition of Micro-organisms

## Introduction:

Microorganisms such as protozoa and bacteria play very important roles in many practical domains, like agriculture, industry and medicine. To explore functions of different categories of microorganisms is a fundamental work in biological studies, which can assist biologists and related scientists to get to know more properties, habits and characteristics of these tiny but obligatos living beings. However, taxonomy of microorganisms (microorganism classification) is traditionally investigated through morphological, chemical or physical analysis, which is time and money consuming. In order to overcome this, since the 1970s CBMIA methods are used to classify microorganisms into different categories using multiple artificial intelligence approaches, such as machine vision, pattern recognition and machine learning algorithms. With the advancement of technology, many new techniques of Deep learning have contributed towards classification of image in a more efficient way such as ResNet, VGG16, Inception V3 etc. Here in the given project, we are using the Inception V3 to classify the microorganisms into their original classes.

## Technical Architecture:



## **Pre-requisites:**

To complete this project, you must require the following software's, concepts, and packages

Anaconda Navigator is a free and open-source distribution of the Python and R programming languages for data science and machine learning related applications. It can be installed on Windows, Linux, and macOS. Conda is an open-source, cross-platform, package management system. Anaconda comes with so very nice tools like JupyterLab, Jupyter Notebook, Spyder, Visual Studio Code. For this project, we will be using Jupyter notebook and Spyder

To install Anaconda navigator and to know how to use Jupyter Notebook & Spyder using Anaconda watch the video

Link: [Click here to watch the video](#)

### 1. To build Machine learning models you must require the following packages

- Numpy:** o It is an open-source numerical Python library. It contains a multidimensional array and matrix data structures and can be used to perform mathematical operations
- Scikit-learn:** o It is a free machine learning library for Python. It features various algorithms like support vector machine, random forests, and k-neighbours, and it also supports Python numerical and scientific libraries like NumPy and SciPy
- Flask:** Web framework used for building Web applications
- Python packages:**
  - ✓ open anaconda prompt as administrator
  - ✓ Type “pip install numpy” and click enter.
  - ✓ Type “pip install pandas” and click enter.
  - ✓ Type “pip install scikit-learn” and click enter.
  - ✓ Type “pip install tensorflow==2.3.2” and click enter.
  - ✓ Type “pip install keras==2.3.1” and click enter.
  - ✓ Type “pip install Flask” and click enter.

## **Deep Learning Concepts**

- CNN:** a convolutional neural network is a class of deep neural networks, most commonly applied to analysing visual imagery.

## **CNN Basic**

- Flask:** Flask is a popular Python web framework, meaning it is a third-party Python library used for developing web applications.

## **Flask Basics**

If you are using Pycharm IDE, you can install the packages through the command prompt and follow the same syntax as above.

### **Project Objectives:**

By the end of this project, you will:

- Know fundamental concepts and techniques of Convolutional Neural Network.
- Gain a broad understanding of image data.
- Know how to pre-process/clean the data using different data pre-processing techniques.
- know how to build a web application using the Flask framework.

### **Project Flow:**

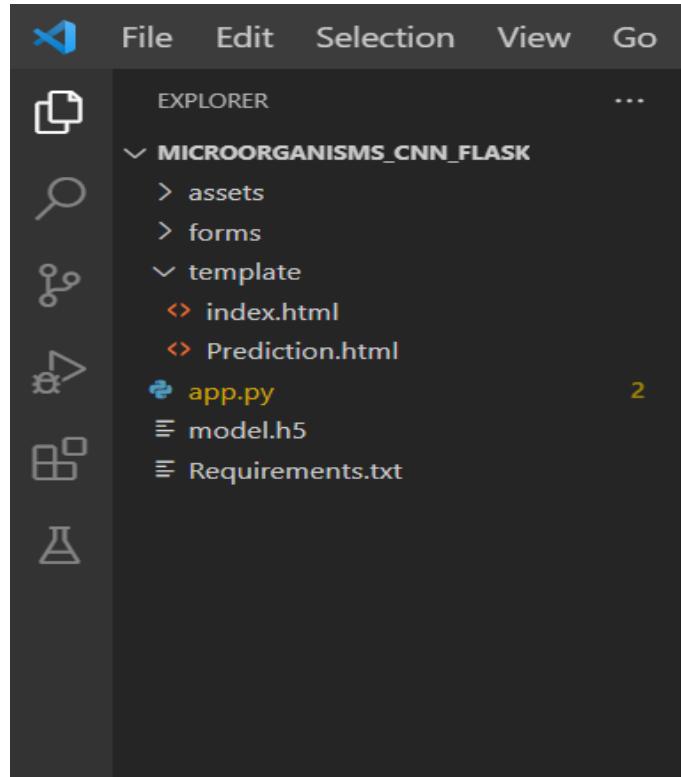
- The user interacts with the UI (User Interface) to choose the image.
- The chosen image analysed by the model which is integrated with flask application.
- Inceptionv3 Model analyse the image, then prediction is showcased on the Flask UI.

To accomplish this, we have to complete all the activities and tasks listed below

- Data Collection.
  - ✓ Create Train and Test Folders.
- Data Pre-processing.
  - ✓ Import the ImageDataGenerator library
  - ✓ Configure ImageDataGenerator class
  - ✓ Apply ImageDataGenerator functionality to Trainset and Test set
- Model Building
  - ✓ Import the model building Libraries
  - ✓ Initializing the model
  - ✓ Adding Input Layer
  - ✓ Adding Hidden Layer
  - ✓ Adding Output Layer
  - ✓ Configure the Learning Process
- Training and testing the model
  - ✓ Save the Model
  - ✓ Application Building
  - ✓ Create an HTML file
  - ✓ Build Python Code

## Project Structure:

Create a Project folder which contains files as shown below



- The Dataset folder contains the training and testing images for training our model.
- We are building a Flask Application that needs HTML pages stored in the templates folder and a python script app.py for server-side scripting
- we need the model which is saved and the saved model in this content is a model.h5 templates folder contains base.html, index.html pages.

## Milestone 1: Collection of Data

Data Collection Collect images of micro-organisms then organized into subdirectories based on their respective names as shown in the project structure. Create folders of types of microbes that need to be recognized.

The given dataset has 8 different types of folders given for Micro Organism they are following:

- Amoeba
- Euglena
- Hydra
- Paramecium
- Rod bacteria
- Spherical bacteria

- Spiral bacteria
- Yeast

## Download the Dataset-

<https://www.kaggle.com/datasets/mdwaquarazam/microorganism-image-classification>

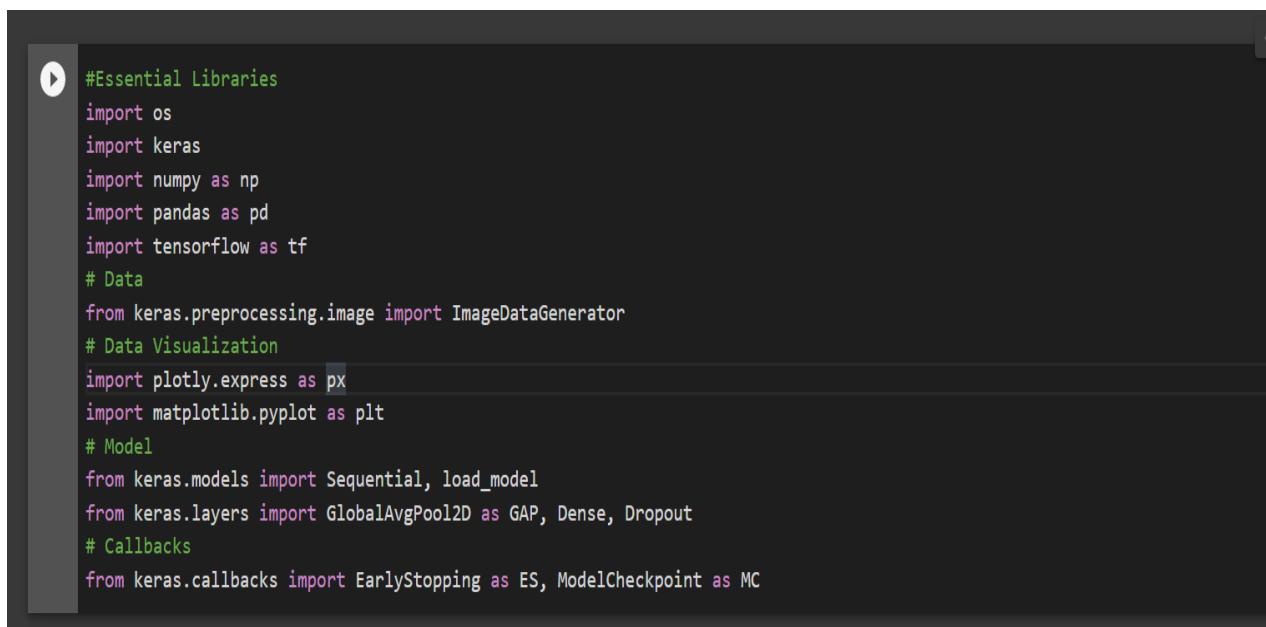
## Milestone 2: Image Pre-processing

In this milestone we will be improving the image data that suppresses unwilling distortions or enhances some image features important for further processing, although perform some geometric transformations of images like rotation, scaling, translation, etc.

### **Activity 1: Import the ImageDataGenerator library**

Image data augmentation is a technique that can be used to artificially expand the size of a training dataset by creating modified versions of images in the dataset.

The Keras deep learning neural network library provides the capability to fit models using image data augmentation via the ImageDataGenerator class. Let us import the ImageDataGenerator class from TensorFlow Keras



```
#Essential Libraries
import os
import keras
import numpy as np
import pandas as pd
import tensorflow as tf
# Data
from keras.preprocessing.image import ImageDataGenerator
# Data Visualization
import plotly.express as px
import matplotlib.pyplot as plt
# Model
from keras.models import Sequential, load_model
from keras.layers import GlobalAvgPool2D as GAP, Dense, Dropout
# Callbacks
from keras.callbacks import EarlyStopping as ES, ModelCheckpoint as MC
```

### **Activity 2: Configure ImageDataGenerator class**

ImageDataGenerator class is instantiated and the configuration for the types of data augmentation

```
[ ] # Initialize generator
gen = ImageDataGenerator(
    rescale=1./255,
    rotation_range=10,
    horizontal_flip=True,
    brightness_range=[0.3,0.8],
    validation_split=0.1
)
```

There are five main types of data augmentation techniques for image data; specifically:

- Image shifts via the width\_shift\_range and height\_shift\_range arguments.
- The image flips via the horizontal flip and vertical flip arguments.
- Image rotations via the rotation range argument
- Image brightness via the brightness range argument.
- Image zoom via the zoom range argument. An instance of the ImageDataGenerator class can be constructed for train and test.

### Activity 3: Apply ImageDataGenerator functionality to Trainset and Test set

Let us apply ImageDataGenerator functionality to Trainset and Test set by using the following code. For Training set using flow\_from\_directory function.

This function will return batches of images from the different classes as 'Amoeba': 0, 'Euglena': 1, 'Hydra': 2, 'Paramecium': 3, 'Rod\_bacteria': 4, 'Spherical\_bacteria': 5, 'Spiral\_bacteria': 6, 'Yeast': 7

Arguments:

- directory: Directory where the data is located. If labels are "inferred", it should contain subdirectories, each containing images for a class. Otherwise, the directory structure is ignored.
- batch\_size: Size of the batches of data which is 64.
- target\_size: Size to resize images after they are read from disk.
- class\_mode:
  - 'int': means that the labels are encoded as integers (e.g., for sparse\_categorical\_crossentropy loss).
  - 'categorical' means that the labels are encoded as a categorical vector (e.g., for categorical\_crossentropy loss).
  - 'binary' means that the labels (there can be only 2) are encoded as float32 scalars with values 0 or 1 (e.g., for binary\_crossentropy).
  - None (no labels).

```
▶ # Load Data
train_ds = gen.flow_from_directory(
    root_path,
    batch_size=128, # For better utilization of GPU, I have kept the batch size a little bit
    shuffle=True,
    class_mode='binary',
    target_size=(256,256), # This image size is generally sufficient for better image classifications.
    subset='training'
)
valid_ds = gen.flow_from_directory(
    root_path,
    batch_size=64, # For faster inference, the batch size here is small.
    shuffle=True,
    class_mode='binary',
    target_size=(256,256), # This image size is generally sufficient for better image classifications.
    subset='validation'
)

Found 714 images belonging to 8 classes.
Found 75 images belonging to 8 classes.
```

## Milestone 3: Model Building

Now it's time to build our Convolutional Neural Networking which contains an input layer along with the convolution, max-pooling, and finally an output layer.

### **Activity 1: Importing the Model Building Libraries**

Importing the necessary libraries

```
# Model
from keras.models import Sequential, load_model
from keras.layers import GlobalAvgPool2D as GAP, Dense, Dropout
# Callbacks
from keras.callbacks import EarlyStopping as ES, ModelCheckpoint as MC

# Pre-Trained Model
from tensorflow.keras.applications import InceptionV3
```

## Activity 2: Exploratory Data Analysis

```
# Get class names
class_names = sorted(os.listdir(root_path))
n_classes = len(class_names)
class_names

['Amoeba',
 'Euglena',
 'Hydra',
 'Paramecium',
 'Rod_bacteria',
 'Spherical_bacteria',
 'Spiral_bacteria',
 'Yeast']

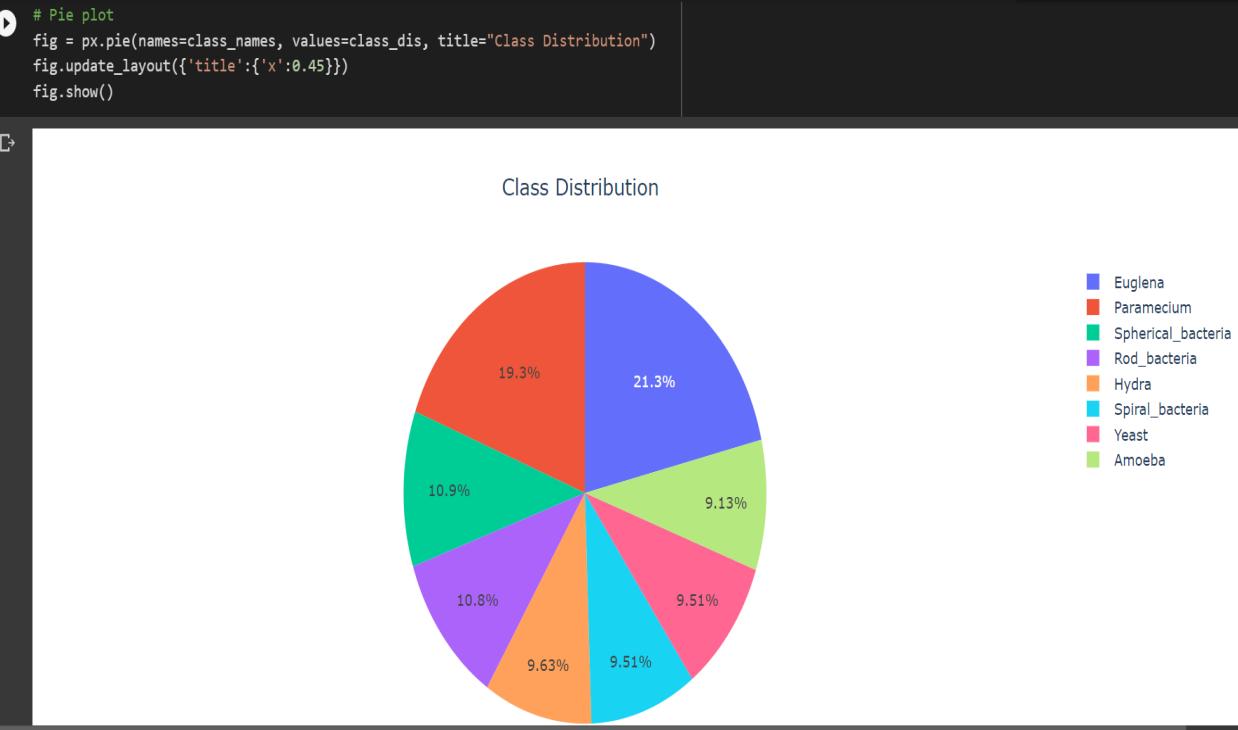
[] n_classes

8

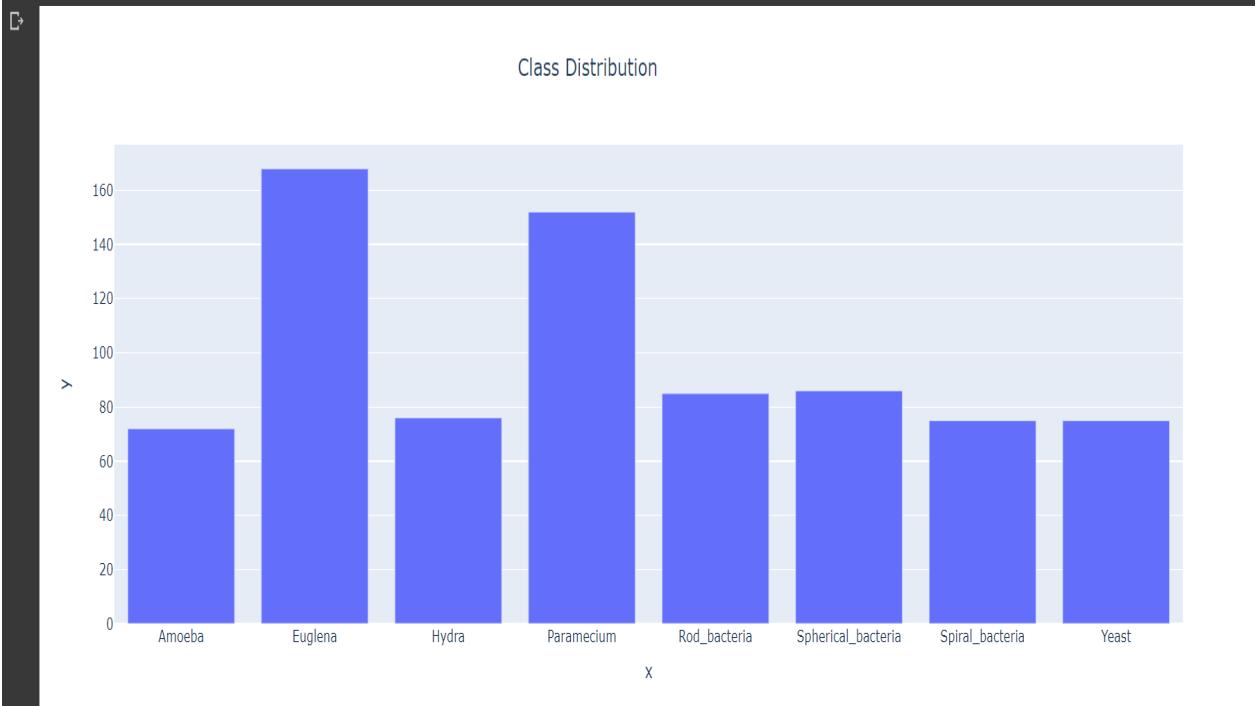
[] # Calculate class distribution
class_dis = [len(os.listdir(root_path + '/' + name)) for name in class_names]
class_dis

[72, 168, 76, 152, 85, 86, 75, 75]
```

## Activity 3: Visualisation of the data



```
# Bar Plot  
fig = px.bar(x=class_names, y=class_dis, title="Class Distribution")  
fig.update_layout({'title': {'x': 0.45}})  
fig.show()
```



```

def show_images(data, GRID=[2,6], model=None, size=(25,10)):

    # The plotting configurations
    n_rows, n_cols = GRID
    n_images = n_rows * n_cols
    plt.figure(figsize=size)

    # Data for visualization
    images, labels = next(iter(data)) # This process can take a little time because of the large batch size

    # Iterate through the subplots.
    for i in range(1, n_images+1):
        # Select a random data
        id = np.random.randint(len(images)) # This is a dynamic function because for validation data and training data, the length of total images is different.
        image, label = images[id], class_names[int(labels[id])]

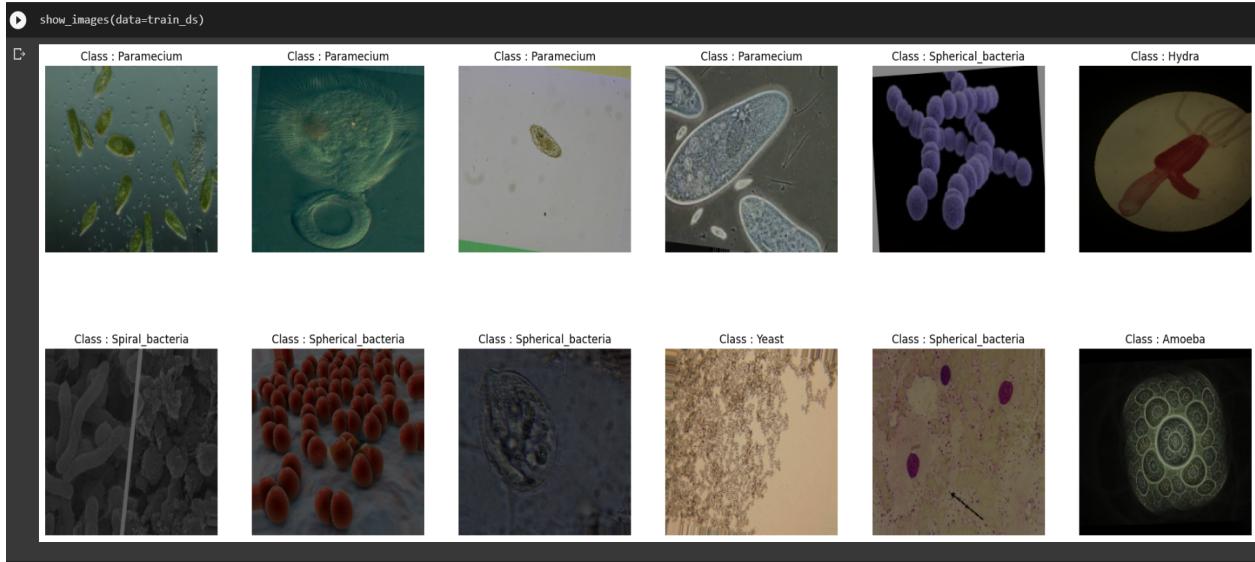
        # Plot the sub plot
        plt.subplot(n_rows, n_cols, i)
        plt.imshow(image)
        plt.axis('off')

        # If model is available make predictions.
        if model is not None:
            pred = class_names[np.argmax(model.predict(image[np.newaxis,...]))]
            title = f"Class : {label}\nPred : {pred}"
        else:
            title = f"Class : {label}"

        plt.title(title)
    plt.show()

[ ] show_images(data=train_ds)

```



## Activity 4: Initializing the model

Keras has 2 ways to define a neural network:

- Sequential
- Function API

The Sequential class is used to define linear initializations of network layers which then, collectively, constitute a model. In our example below, we will use the Sequential constructor to create a model, which will then have layers added to it using the add() method.

### Activity 5: Configure the Learning Process

- The compilation is the final step in creating a model. Once the compilation is done, we can move on to the training phase. The loss function is used to find errors or deviations in the learning process. Keras requires a loss function during the model compilation process.
- Optimization is an important process that optimizes the input weights by comparing the prediction and the loss function. Here we are using adam optimizer
- Metrics are used to evaluate the performance of your model. It is similar to the loss function, but not used in the training process

### Activity 6: Train The model

Now, let us train our model with our image dataset. The model is trained for 30 epochs and after every epoch, the current model state is saved if the model has the least loss encountered till that time. We can see that the training loss decreases in almost every epoch till 30 epochs and probably there is further scope to improve the model.

`fit_generator` functions used to train a deep learning neural network

Arguments:

- `steps_per_epoch`: it specifies the total number of steps taken from the generator as soon as one epoch is finished and the next epoch has started. We can calculate the value of `steps_per_epoch` as the total number of samples in your dataset divided by the batch size.
- `Epochs`: an integer and number of epochs we want to train our model for.
- `validation_data` can be either:
  - an inputs and targets list
  - a generator
  - an inputs, targets, and sample\_weights list which can be used to evaluate the loss and metrics for any model after any epoch has ended.
  - `validation_steps`: only if the `validation_data` is a generator then only this argument can be used. It specifies the total number of steps taken from the generator before it is stopped at every epoch and its value is calculated as the total number of validation data points in your dataset divided by the validation batch size

```

▶ # Give you a model, a name
name = "inception-v3"

# Base model
base = InceptionV3(input_shape=(256,256,3), include_top=False)
base.trainable = False

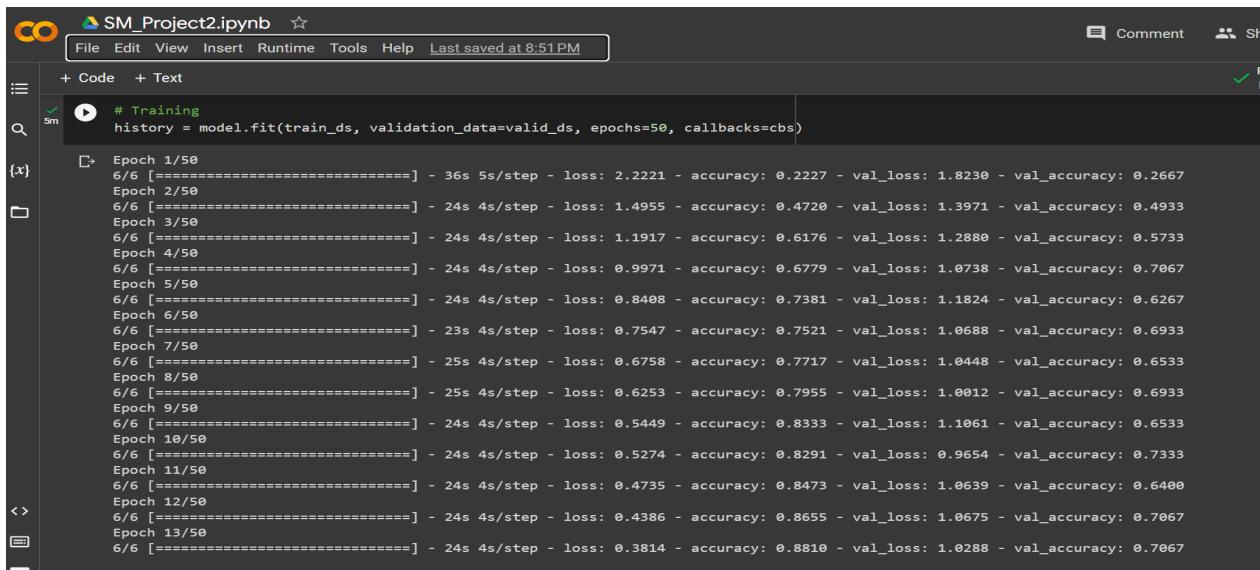
# Model Architecture
model = Sequential([
    base, 
    Dense(256, kernel_initializer='he_normal', activation='relu'),
    Dropout(0.2),
    Dense(n_classes, activation='softmax')
])
# Callbacks
cbs = [ES(patience=3, restore_best_weights=True), MC(name + ".h5", save_best_only=True)]

# Compile Model
opt = tf.keras.optimizers.Adam(learning_rate=1e-3) # Higher than the default learning rate
model.compile(loss='sparse_categorical_crossentropy', optimizer=opt, metrics=['accuracy'])

# Training
history = model.fit(train_ds, validation_data=valid_ds, epochs=15, callbacks=cbs)

```

Analysing the trained model by looking at the accuracy through the epochs:



The screenshot shows a Jupyter Notebook cell titled "SM\_Project2.ipynb". The cell contains Python code for training a model and printing the training history. The output of the code is displayed below, showing the progress of 15 epochs. Each epoch shows the training loss, validation loss, and validation accuracy. The validation accuracy remains relatively stable around 0.6 to 0.7.

```

# Training
history = model.fit(train_ds, validation_data=valid_ds, epochs=15, callbacks=cbs)

Epoch 1/50
6/6 [=====] - 36s 5s/step - loss: 2.2221 - accuracy: 0.2227 - val_loss: 1.8230 - val_accuracy: 0.2667
Epoch 2/50
6/6 [=====] - 24s 4s/step - loss: 1.4955 - accuracy: 0.4720 - val_loss: 1.3971 - val_accuracy: 0.4933
Epoch 3/50
6/6 [=====] - 24s 4s/step - loss: 1.1917 - accuracy: 0.6176 - val_loss: 1.2880 - val_accuracy: 0.5733
Epoch 4/50
6/6 [=====] - 24s 4s/step - loss: 0.9971 - accuracy: 0.6779 - val_loss: 1.0738 - val_accuracy: 0.7067
Epoch 5/50
6/6 [=====] - 24s 4s/step - loss: 0.8408 - accuracy: 0.7381 - val_loss: 1.1824 - val_accuracy: 0.6267
Epoch 6/50
6/6 [=====] - 23s 4s/step - loss: 0.7547 - accuracy: 0.7521 - val_loss: 1.0688 - val_accuracy: 0.6933
Epoch 7/50
6/6 [=====] - 25s 4s/step - loss: 0.6758 - accuracy: 0.7717 - val_loss: 1.0448 - val_accuracy: 0.6533
Epoch 8/50
6/6 [=====] - 25s 4s/step - loss: 0.6253 - accuracy: 0.7955 - val_loss: 1.0012 - val_accuracy: 0.6933
Epoch 9/50
6/6 [=====] - 24s 4s/step - loss: 0.5449 - accuracy: 0.8333 - val_loss: 1.1061 - val_accuracy: 0.6533
Epoch 10/50
6/6 [=====] - 24s 4s/step - loss: 0.5274 - accuracy: 0.8291 - val_loss: 0.9654 - val_accuracy: 0.7333
Epoch 11/50
6/6 [=====] - 24s 4s/step - loss: 0.4735 - accuracy: 0.8473 - val_loss: 1.0639 - val_accuracy: 0.6400
Epoch 12/50
6/6 [=====] - 24s 4s/step - loss: 0.4386 - accuracy: 0.8655 - val_loss: 1.0675 - val_accuracy: 0.7067
Epoch 13/50
6/6 [=====] - 24s 4s/step - loss: 0.3814 - accuracy: 0.8810 - val_loss: 1.0288 - val_accuracy: 0.7067

```

Visualising Model Summary:

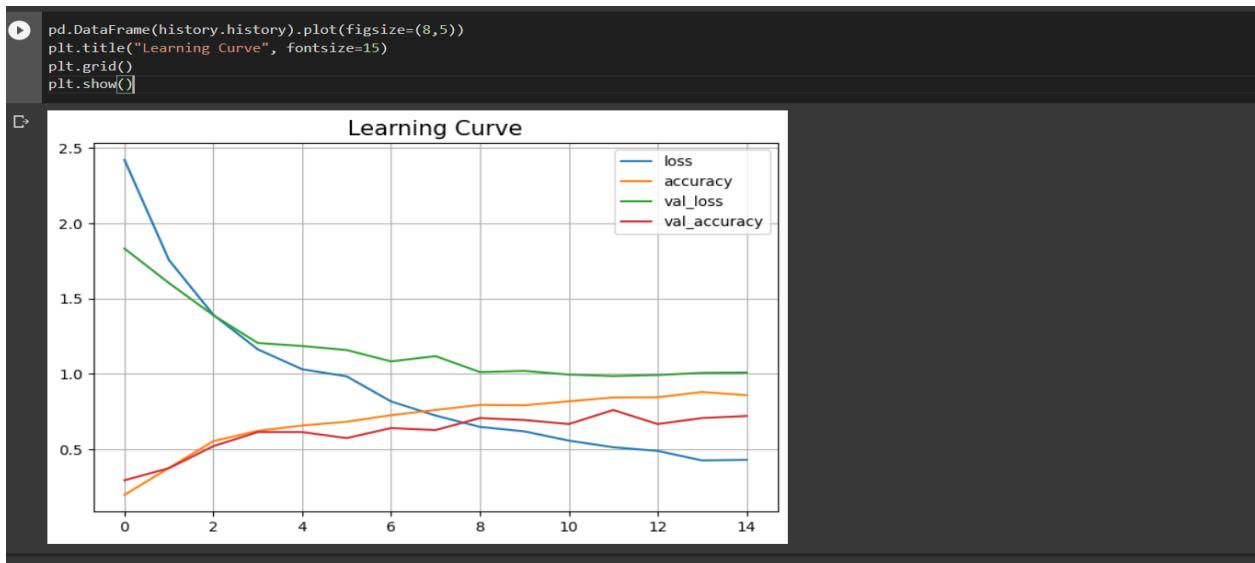
```
▶ model.summary()

Model: "sequential_2"
-----
```

Layer (type)	Output Shape	Param #
inception_v3 (Functional)	(None, 6, 6, 2048)	21802784
global_average_pooling2d_2 (GlobalAveragePooling2D)	(None, 2048)	0
dense_4 (Dense)	(None, 256)	524544
dropout_2 (Dropout)	(None, 256)	0
dense_5 (Dense)	(None, 8)	2056

```
=====
Total params: 22,329,384
Trainable params: 526,600
Non-trainable params: 21,802,784
```

Visualising the loss and accuracy of the trained model.



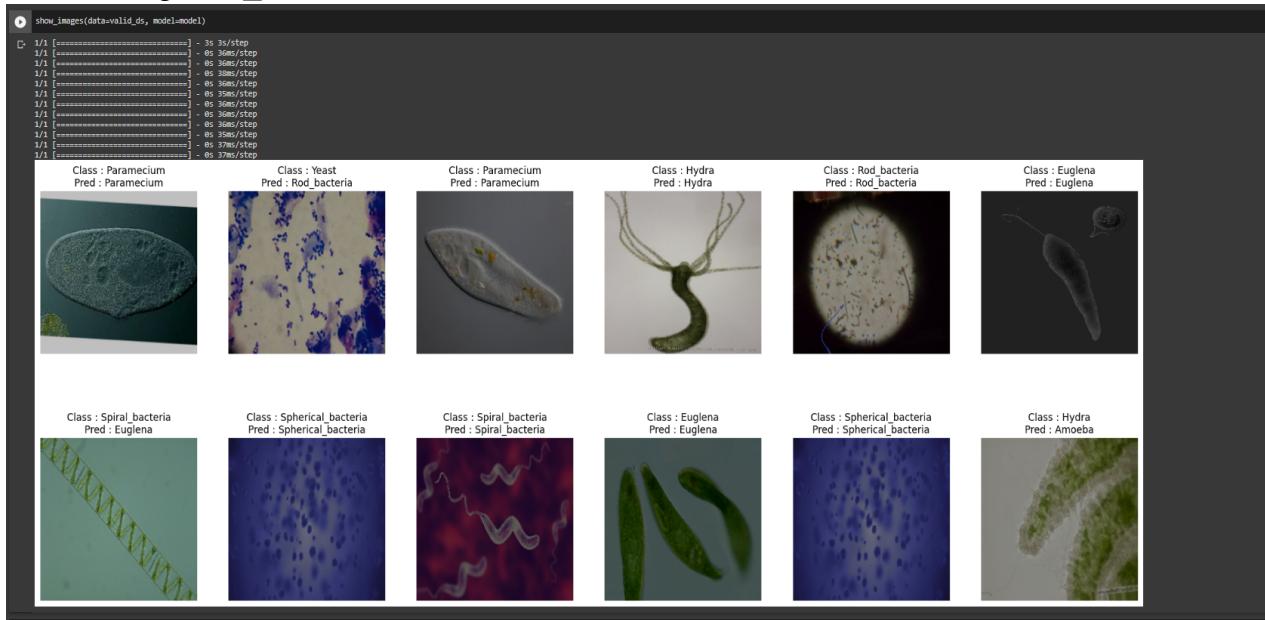
## Activity 7: Save the Model

The model is saved with .h5 extension as follows An H5 file is a data file saved in the Hierarchical Data Format (HDF). It contains multidimensional arrays of scientific data.

```
[ ] model.save("model1.h5")
```

## Activity 8: Test The model

Evaluation is a process during the development of the model to check whether the model is the best fit for the given problem and corresponding data. Load the saved model using `load_model`



Taking an image as input and checking the results. **Displaying 4 Test Results.**

```

[ ] from tensorflow.keras.preprocessing import image
import numpy as np

[ ] ## Testing 1

img= image.load_img('/content/drive/MyDrive/Micro_Organism/Spiral_bacteria/Image_11.jpg',target_size=(192,192))    #read image
X= image.img_to_array(img) # converting the image into array

X= np.expand_dims(X,axis=0)
pred = np.argmax(model.predict(X)) # predicting the higher probability index
op=['Amoeba', 'Euglena', 'Hydra', 'Paramecium', 'Rod_bacteria', 'Spherical_bacteria', 'Spiral_bacteria', 'Yeast'] # creating a list
op[pred] # List indexing with output

1/1 [=====] - 0s 29ms/step
'Spiral_bacteria'

```

```

[ ] ## Testing 2

img= image.load_img('/content/drive/MyDrive/Micro_Organism/Rod_bacteria/Image_1.jpg',target_size=(120,120))    #read image
X= image.img_to_array(img) # converting the image into array

X= np.expand_dims(X,axis=0)
pred = np.argmax(model.predict(X)) # predicting the higher probability index
op=['Amoeba', 'Euglena', 'Hydra', 'Paramecium', 'Rod_bacteria', 'Spherical_bacteria', 'Spiral_bacteria', 'Yeast'] # creating a list
op[pred] # List indexing with output

1/1 [=====] - 0s 42ms/step
'Rod_bacteria'

```

```
## Testing 3

img= image.load_img('/content/drive/MyDrive/Micro_Organism/Spiral_bacteria/Image_10.jpg',target_size=(256,256)) #read image
X= image.img_to_array(img) # converting the image into array
|
X= np.expand_dims(X,axis=0)
pred = np.argmax(model.predict(X)) # predicting the higher probability index
op=['Amoeba', 'Euglena', 'Hydra', 'Paramecium', 'Rod_bacteria', 'Spherical_bacteria', 'Spiral_bacteria', 'Yeast'] # creating a list
op[pred] # List indexing with output

1/1 [=====] - 0s 29ms/step
'Spiral_bacteria'
```

```
## Testing 4

img= image.load_img('/content/drive/MyDrive/Micro_Organism/Amoeba/Image_1.jpg',target_size=(64,64)) #read image
X= image.img_to_array(img) # converting the image into array

X= np.expand_dims(X,axis=0)
pred = np.argmax(model.predict(X)) # predicting the higher probability index
op=['Amoeba', 'Euglena', 'Hydra', 'Paramecium', 'Rod_bacteria', 'Spherical_bacteria', 'Spiral_bacteria', 'Yeast'] # creating a list
op[pred] # List indexing with output

1/1 [=====] - 1s 683ms/step
'Amoeba'
```

## Milestone 4: Application Building

Now that we have trained our model, let us build our flask application which will be running in our local browser with a user interface.

In the flask application, the input parameters are taken from the HTML page. These factors are then given to the model to know to predict the type of Garbage and showcased on the HTML page to notify the user. Whenever the user interacts with the UI and selects the “Image” button, the next page is opened where the user chooses the image and predicts the output.

### Activity 1: Create HTML Pages

- We use HTML to create the front end part of the web page.
- Here, we have created 3 HTML pages- home.html, intro.html, and upload.html
- home.html displays the home page.
- Intro.html displays an introduction about the project
- upload.html gives the emergency alert. For more information regarding HTML <https://www.w3schools.com/html/>
- We also use JavaScript-main.js and CSS-main.css to enhance our functionality and view of HTML pages.
- Link :CSS , JS

```

PROBLEMS OUTPUT DEBUG CONSOLE TERMINAL
PS C:\Users\nitis\Downloads\SM_Project2\MicroOrganisms_CNN_Flask> C:/Users/nitis/anaconda3/scripts/activate
PS C:\Users\nitis\Downloads\SM_Project2\MicroOrganisms_CNN_Flask> conda activate MiniProject
PS C:\Users\nitis\Downloads\SM_Project2\MicroOrganisms_CNN_Flask> & C:/Users/nitis/anaconda3/envs/MiniProject/python.exe c:/Users/nitis/Downloads/SM_Project2/MicroOrganisms_CNN_Flask/app.py
* Serving Flask app 'app'
* Debug mode: on
WARNING: This is a development server. Do not use it in a production deployment. Use a production WSGI server instead.
Press CTRL+C to quit
* Restarting with watchdog (windowsapi)
* Debugger is active!
* Debugger PIN: 261-368-191
127.0.0.1 - - [03/May/2023 14:45:30] "GET / HTTP/1.1" 200 -
127.0.0.1 - - [03/May/2023 14:45:31] "GET /static/vendor/remixicon/remixicon.css HTTP/1.1" 304 -
127.0.0.1 - - [03/May/2023 14:45:31] "GET /static/vendor/boxicons/css/boxicons.min.css HTTP/1.1" 304 -
127.0.0.1 - - [03/May/2023 14:45:31] "GET /static/vendor/aos/aos.css HTTP/1.1" 304 -
127.0.0.1 - - [03/May/2023 14:45:31] "GET /static/vendor/lightbox/css/lightbox.min.css HTTP/1.1" 304 -

```

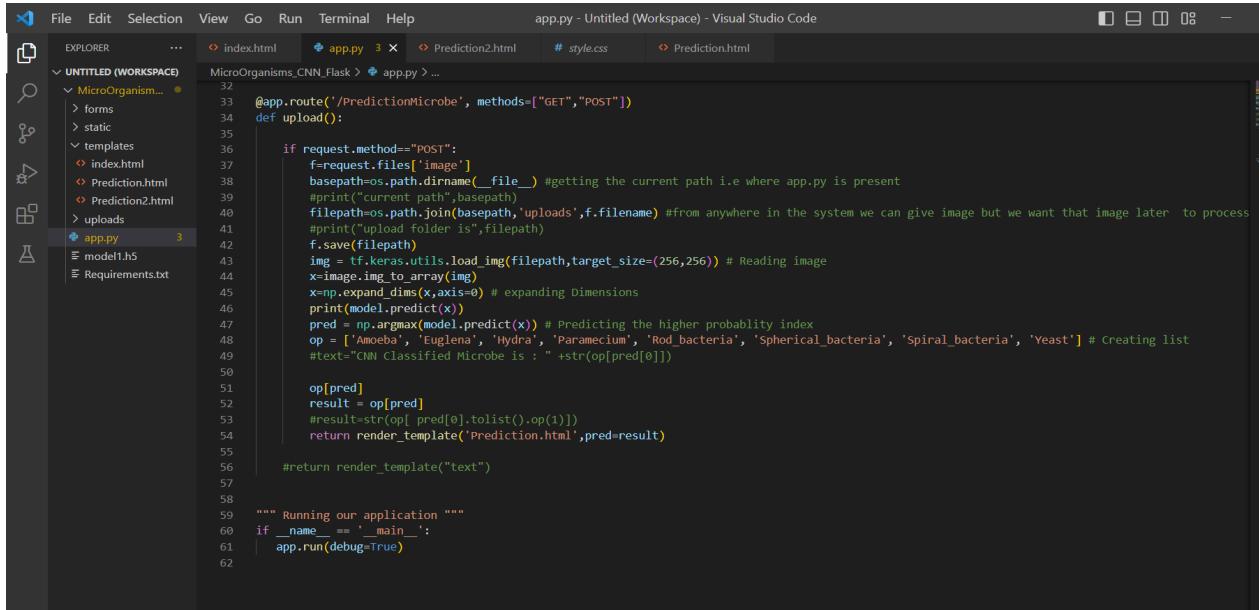
## Create app.py (Python Flask) file: -

Write below code in Flask app.py python file script to run Micro-Organism Classification Project.

```

File Edit Selection View Go Run Terminal Help app.py - Untitled (Workspace) - Visual Studio Code
EXPLORER UNTITLED (WORKSPACE) MicroOrganisms_CNN_Flask > app.py ...
index.html Prediction2.html # style.css Prediction.html
MicroOrganisms_CNN_Flask > app.py ...
1 import numpy as np
2 import pandas as pd
3 import os
4 import tensorflow as tf
5 from flask import Flask, render_template, url_for, request
6 from tensorflow.keras.models import Model
7 from tensorflow.keras.preprocessing import image
8 from tensorflow.python.ops.gen_array_ops import Concat
9 from tensorflow.keras.models import load_model
10
11 #Loading the model
12 model=load_model("model1.h5",compile=False)
13 app=Flask(__name__)
14
15 #default home page or route
16 @app.route('/')
17 def home():
18     return render_template('index.html')
19
20 @app.route('/Prediction')
21 def Prediction():
22     return render_template('Prediction.html')
23
24

```



```

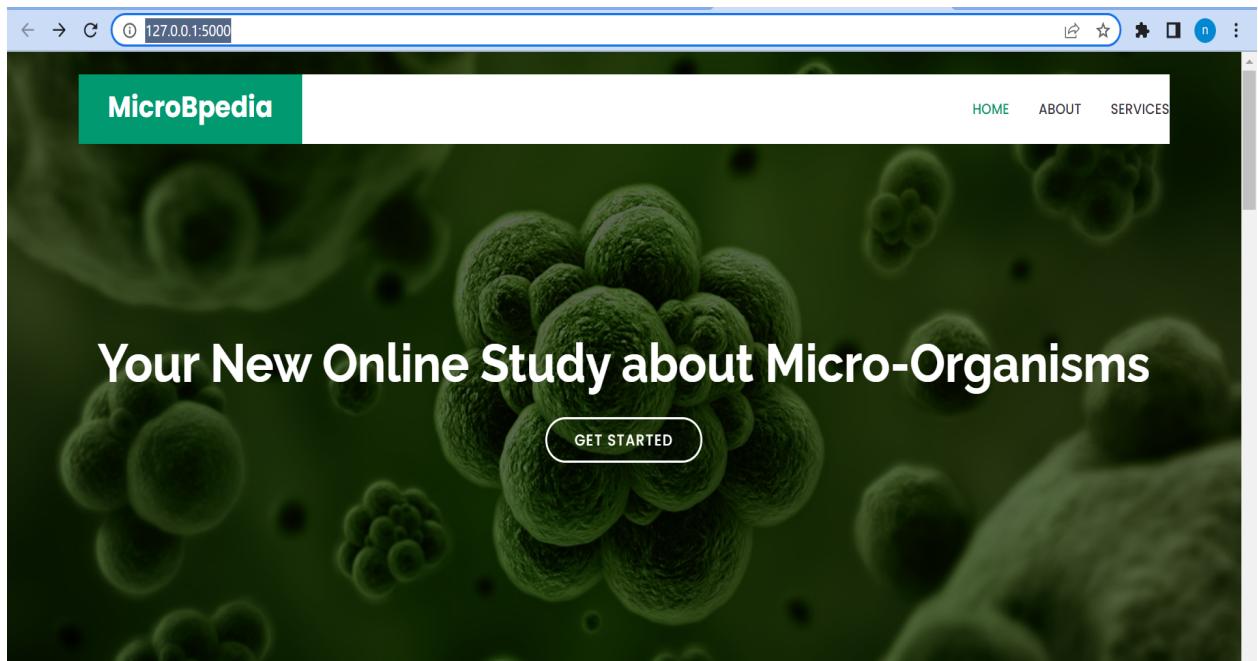
File Edit Selection View Go Run Terminal Help
EXPLORER index.html app.py Prediction2.html # style.css Prediction.html
UNTITLED (WORKSPACE)
MicroOrganisms_CNN_Flask
> forms
> static
templates
index.html
Prediction.html
Prediction2.html
uploads
app.py 3
model1.h5
Requirements.txt

app.py - Untitled (Workspace) - Visual Studio Code

32 @app.route('/PredictionMicrobe', methods=["GET", "POST"])
33 def upload():
34     if request.method=="POST":
35         f=request.files['image']
36         basepath=os.path.dirname( __file__ ) #getting the current path i.e where app.py is present
37         #print("current path",basepath)
38         filepath=os.path.join(basepath,'uploads',f.filename) #from anywhere in the system we can give image but we want that image later to process
39         #print("upload folder is",filepath)
40         f.save(filepath)
41         img = tf.keras.utils.load_img(filepath,target_size=(256,256)) # Reading image
42         x=image.img_to_array(img)
43         x=np.expand_dims(x,axis=0) # expanding Dimensions
44         print(model.predict(x))
45         pred = np.argmax(model.predict(x)) # Predicting the higher probability index
46         op = ['Amoeba', 'Euglena', 'Hydra', 'Paramecium', 'Rod_bacteria', 'Spherical_bacteria', 'Spiral_bacteria', 'Yeast'] # Creating list
47         #text="CNN Classified Microbe is : "+str(op[pred[0]])
48
49
50
51
52
53
54
55
56
57
58 """
59     Running our application """
60 if __name__ == "__main__":
61     app.run(debug=True)
62

```

**Index.html is displayed below (Webpage named “MicroBpedia”):**



## INTRODUCTION

The 17th-century discovery of living forms existing invisible to the naked eye was a significant milestone in the history of science, for from the 13th century onward it had been postulated that “invisible” entities were

**About Section is displayed below:**

The screenshot shows a browser window with the URL 127.0.0.1:5000. The page title is "MicroBpedia". The navigation bar includes links for "HOME", "ABOUT", and "SERVICES".

# INTRODUCTION

Microbiology, study of microorganisms, or microbes, a diverse group of generally minute simple life-forms that include bacteria, archaea, algae, fungi, protozoa, and viruses. The field is concerned with the structure, function, and classification of such organisms and with ways of both exploiting and controlling their activities.

The 17th-century discovery of living forms existing invisible to the naked eye was a significant milestone in the history of science, for from the 13th century onward it had been postulated that "invisible" entities were responsible for decay and disease. The word *microbe* was coined in the last quarter of the 19th century to describe these organisms, all of which were thought to be related. As microbiology eventually developed into a specialized science, it was found that microbes are a very large group of extremely diverse organisms.

- ✓ Microbiology essentially began with the development of the microscope. Although others may have seen microbes before him, it was Antonie van Leeuwenhoek, a Dutch draper whose hobby was lens grinding and making microscopes, who was the first to provide proper documentation of his observations. Leeuwenhoek conveyed his findings in a series of letters to the British Royal Society during the mid-1670s.
- ✓ Girolamo Fracastoro, an Italian scholar, advanced the notion as early as the mid-1500s that contagion is an infection that passes from one thing to another. A description of precisely what is passed along eluded discovery until the late 1800s, when the work of many scientists, Pasteur foremost among them, determined the role of bacteria in fermentation and disease. Robert Koch, a German physician, defined the procedure (Koch's postulates) for proving that

The screenshot shows a browser window with the URL 127.0.0.1:5000. The page title is "MicroBpedia". The navigation bar includes links for "HOME", "ABOUT", and "SERVICES".

**Types of different Microrganisms**

The major groups of microorganisms—namely bacteria, archaea, fungi (yeasts and molds), algae, protozoa, and viruses—are summarized below. Links to the more detailed articles on each of the major groups are provided.

[Learn More >](#)

**Spiral\_Bacteria**  
Spiral bacteria, bacteria of spiral (helical) shape, form the third major morphological category of prokaryotes along with the rod-shaped bacilli and round cocci.

**Rod\_Bacteria**  
Bacillus (Latin "stick") is a genus of Gram-positive, rod-shaped bacteria, a member of the phylum Bacillota, with 266 named species.

**Yeast**  
Yeasts are eukaryotic, single-celled microorganisms classified as members of the fungus kingdom.

**Sperical\_Bacteria**  
A colony of *Escherichia coli* [11] Unlike in multicellular organisms, increases in cell size (cell growth) and reproduction by cell division are tightly linked in unicellular organisms

**Paramecium**  
It's a genus of eukaryotic, unicellular ciliates, commonly studied as a model organism of the ciliate group.

**Euglena**  
Euglena, genus of more than 1,000 species of single-celled flagellated (i.e., having a whiplike appendage) microorganisms that feature both plant and animal characteristics

**Services section is displayed below with a click button.**

The screenshot shows a web browser window with the URL 127.0.0.1:5000. The page has a green header bar with the logo "MicroBpedia". The main content area is titled "Our Services" and contains a sub-section for "Prediction". It includes a placeholder text "Upload a picture to know the type of Microorganism it is.", a circular icon with a magnifying glass, and a button labeled "Predict the type of Micro-Organism". A green arrow points from the text to the button. The top right corner of the page has navigation links for "HOME", "ABOUT", and "SERVICES".

**Finally, the Footer of the Webpage along with the choosing of Image for prediction.**

The screenshot shows a web browser window with the URL 127.0.0.1:5000/Prediction. The page title is "Prediction Page" and the breadcrumb navigation shows "Home / Prediction Page". The main content area is titled "Upload Image Here To Identify the MicroOrganism" and features a file input field with the placeholder "Choose... Choose File Image\_5.jpg" and a blue "Predict!" button. The top right corner of the page has navigation links for "HOME", "ABOUT", and "SERVICES".

**Final Predicted Output (after I click on the Predict Button) is displayed as follows:**

← → ⌂ 127.0.0.1:5000/PredictionMicrobe

**MicroBpedia**

Prediction Page

HOME ABOUT SERVICES

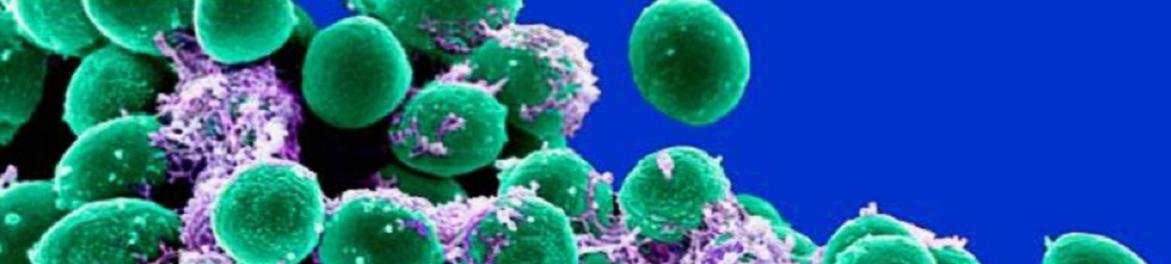
Home / Prediction Page

Upload Image Here To Identify the MicroOrganism

Choose... Choose File No file chosen

Predict!

**Prediction: Spherical\_bacteria**



← → ⌂ 127.0.0.1:5000/PredictionMicrobe

**MicroBpedia**

Prediction Page

HOME ABOUT SERVICES

Upload Image Here To Identify the MicroOrganism

Choose... Choose File No file chosen

Predict!

**Prediction: Spiral\_bacteria**



**MicroBpedia**

Blue Ridge, Rajiv Gandhi Infotech Park,  
 Hinjewadi

**Useful Links**

- > Home
- > Prediction
- > About us

**Our Services**

- > Home
- > Prediction
- > About us

**Join Our Newsletter**

Please subscribe and sign-in to get notification from our weekly newsletter and avail its benefits.

↑

← → C ① 127.0.0.1:5000/PredictionMicrobe

MicroBpedia

HOME ABOUT SERVICES

Prediction Page

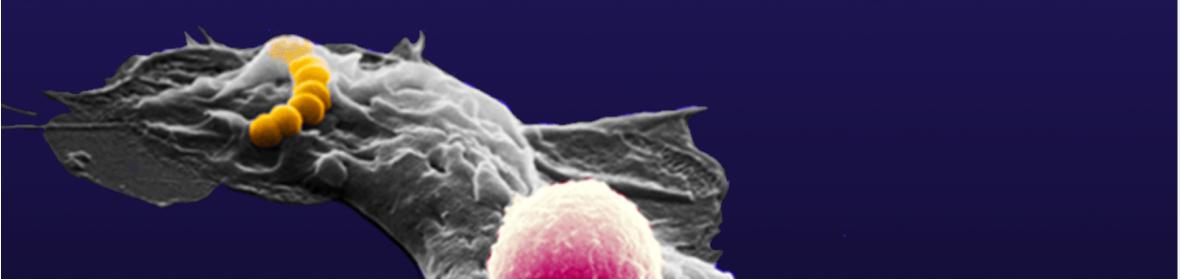
Home / Prediction Page

Upload Image Here To Identify the MicroOrganism

Choose... Choose File No file chosen

Predict!

**Prediction: Amoeba**



\*\*\*\*Thank You\*\*\*\*