

HematoVision: Advanced Blood Cell Classification Using Transfer Learning

HematoVision aims to develop an accurate and efficient model for classifying blood cells by employing transfer learning techniques. Utilizing a dataset of 12,000 annotated blood cell images, categorized into distinct classes such as eosinophils, lymphocytes, monocytes, and neutrophils, the project leverages pre-trained convolutional neural networks (CNNs) to expedite training and improve classification accuracy. Transfer learning allows the model to benefit from pre-existing knowledge of image features, significantly enhancing its performance and reducing computational costs. This approach provides a reliable and scalable tool for pathologists and healthcare professionals, ensuring precise and efficient blood cell classification.

Scenario 1: Automated Diagnostic Systems for Healthcare

Integrating HematoVision into automated diagnostic systems in clinical settings can revolutionize blood analysis. By using transfer learning, the system quickly adapts to the specifics of blood cell classification, capturing images of blood samples, classifying the cells in real-time, and generating detailed reports. This automation reduces the manual workload on pathologists, speeds up diagnostic processes, and ensures high accuracy in results, ultimately improving patient care and treatment efficiency.

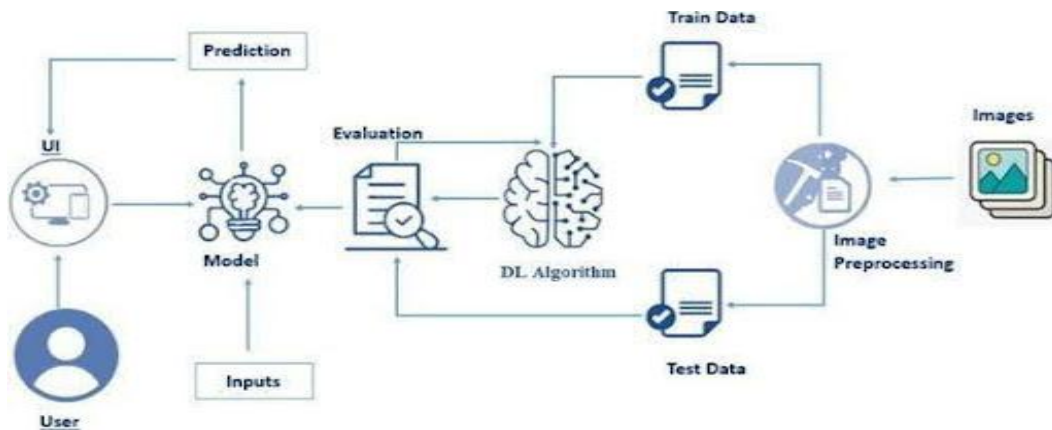
Scenario 2: Remote Medical Consultations

HematoVision can be employed in telemedicine platforms to enhance remote consultations and diagnostics. With transfer learning, the model's ability to accurately classify blood cells from diverse sources is improved, allowing healthcare providers to upload blood cell images for automated analysis. This enables timely and accurate assessments without the need for in-person visits, facilitating better access to specialized medical expertise and improving healthcare delivery in remote or underserved areas.

Scenario 3: Educational Tools for Medical Training

HematoVision's transfer learning-based classification model can be integrated into educational tools for medical training. By incorporating this advanced technology into interactive learning platforms, students and laboratory technicians can upload and analyze blood cell images to receive instant feedback. This hands-on learning experience enhances their understanding of blood cell morphology and classification, providing practical skills and knowledge that are crucial for accurate diagnostic practice and medical training.

Architecture:



Prerequisites

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

Anaconda Navigator:

Refer to the link below to download Anaconda Navigator

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 matplotlib" and click enter.

Type "pip install scipy" and click enter.

Type "pip install seaborn" and click enter.

Type "pip install tensorflow" and click enter.

Type "pip install Flask" and click enter.

Prior Knowledge

You must have prior knowledge of the following topics to complete this project.

DL Concepts

Neural Networks:: <https://www.analyticsvidhya.com/blog/2020/02/cnn-vs-rnn-vs-mlp-analyzing-3-types-of-neural-networks-in-deep-learning/>

Deep Learning Frameworks:: <https://www.knowledgehut.com/blog/data-science/pytorch-vs-tensorflow>

Transfer Learning: <https://towardsdatascience.com/a-demonstration-of-transfer-learning-of-vgg-convolutional-neural-network-pre-trained-model-with-c9f5b8b1ab0a>

VGG16: <https://www.geeksforgeeks.org/vgg-16-cnn-model/>

Convolutional Neural Networks

(CNNs): <https://www.analyticsvidhya.com/blog/2021/05/convolutional-neural-networks-cnn/> [s://www.javatpoint.com/k-nearest-neighbor-algorithm-for-machine-learning](https://www.javatpoint.com/k-nearest-neighbor-algorithm-for-machine-learning)

Overfitting and

Regularization: <https://www.analyticsvidhya.com/blog/2021/07/prevent-overfitting-using-regularization-techniques/>

Optimizers: <https://www.analyticsvidhya.com/blog/2021/10/a-comprehensive-guide-on-deep-learning-optimizers/>

Flask Basics: https://www.youtube.com/watch?v=lj4l_CvBnt0

Project Objectives

- By the end of this project, you will:
- Know fundamental concepts and techniques used for Deep Learning.
- Gain a broad understanding of data.
- Have knowledge of pre-processing the data/transformation techniques on outliers and some visualization concepts.

Project Flow

- The user interacts with the UI (User Interface) to choose the image.
- The chosen image is analyzed by the model which is integrated with the flask application.
- Once the model analyses the input the prediction is showcased on the UI

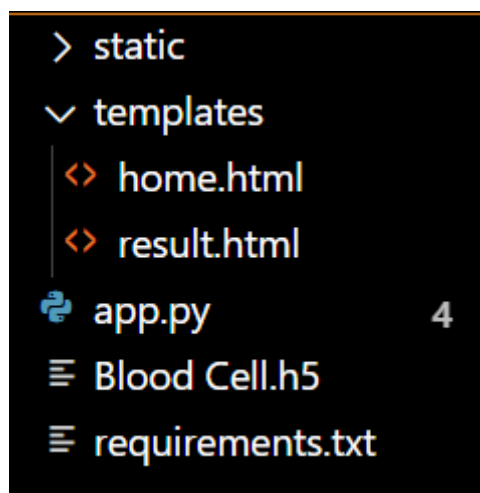
To accomplish this, we have to complete all the activities listed below,

- Data Collection: Collect or download the dataset that you want to train.
- Data pre-processing

- Data Augmentation
- Splitting data into train and test
- Model building
- Import the model-building libraries
- Initializing the model
- Training and testing the model
- Evaluating the performance of the model
- Save the model
- Application Building
- Create an HTML file
- Build python code

Project Structure

Create the Project folder which contains files as shown below



- We are building a Flask application with HTML pages stored in the templates folder and a Python script app.py for scripting.
- Blood Cell.h5 is our saved model. Further, we will use this model for flask integration.

• Data Collection and Preparation

- ML depends heavily on data. It is the most crucial aspect that makes algorithm training possible. So, this section allows you to download the required dataset.

Split Data and Model Building

Train-Test-Split:

In this project, we have already separated data for training and testing.

```
train_images, test_images = train_test_split(bloodCell_df, test_size=0.3, random_state=42)
train_set, val_set = train_test_split(bloodCell_df, test_size=0.2, random_state=42)
```

```
print(train_set.shape)
print(test_images.shape)
print(val_set.shape)
print(train_images.shape)
```

```
(7965, 2)
(2988, 2)
(1992, 2)
(6969, 2)
```

```
image_gen = ImageDataGenerator(preprocessing_function= tf.keras.applications.mobilenet_v2.preprocess_input)
train = image_gen.flow_from_dataframe(dataframe= train_set,x_col="filepaths",y_col="labels",
                                     target_size=(244,244),
                                     color_mode='rgb',
                                     class_mode="categorical",
                                     batch_size=8,
                                     shuffle=False
                                   )
test = image_gen.flow_from_dataframe(dataframe= test_images,x_col="filepaths", y_col="labels",
                                    target_size=(244,244),
                                    color_mode='rgb',
                                    class_mode="categorical",
                                    batch_size=8,
                                    shuffle= False
                                   )
val = image_gen.flow_from_dataframe(dataframe= val_set,x_col="filepaths", y_col="labels",
                                   target_size=(244,244),
                                   color_mode= 'rgb',
                                   class_mode="categorical",
                                   batch_size=8,
                                   shuffle=False
                                   )
```

```
Found 7965 validated image filenames belonging to 4 classes.
Found 2988 validated image filenames belonging to 4 classes.
Found 1992 validated image filenames belonging to 4 classes.
```

Model Building:

Mobilenet V2 Transfer-Learning Model:

The MobileNetV2-based neural network is created using a pre-trained MobileNetV2 architecture with frozen weights. The model is built sequentially, incorporating the MobileNetV2 base, a flattening layer, dropout for regularization, and a dense layer with SoftMax activation for classification into four categories of blood cells. The model is compiled using the Adam optimizer and categorical cross-entropy loss. During training, which spans 5 epochs, a generator is employed for the training data, and validation is conducted with callbacks such as Model Checkpoint and Early Stopping. The best-performing model is saved as "blood_cell.h5" for future use. The model summary provides an overview of the architecture, showcasing the layers and parameters involved.

```

model = keras.models.Sequential([
    keras.layers.Conv2D(filters=128, kernel_size=(8, 8), strides=(3, 3), activation='relu', input_shape=(224, 224, 3)),
    keras.layers.BatchNormalization(),

    keras.layers.Conv2D(filters=256, kernel_size=(5, 5), strides=(1, 1), activation='relu', padding="same"),
    keras.layers.BatchNormalization(),
    keras.layers.MaxPool2D(pool_size=(3, 3)),

    keras.layers.Conv2D(filters=256, kernel_size=(3, 3), strides=(1, 1), activation='relu', padding="same"),
    keras.layers.BatchNormalization(),
    keras.layers.Conv2D(filters=256, kernel_size=(1, 1), strides=(1, 1), activation='relu', padding="same"),
    keras.layers.BatchNormalization(),
    keras.layers.Conv2D(filters=256, kernel_size=(1, 1), strides=(1, 1), activation='relu', padding="same"),
    keras.layers.BatchNormalization(),

    keras.layers.Conv2D(filters=512, kernel_size=(3, 3), activation='relu', padding="same"),
    keras.layers.BatchNormalization(),
    keras.layers.MaxPool2D(pool_size=(2, 2)),

    keras.layers.Conv2D(filters=512, kernel_size=(3, 3), activation='relu', padding="same"),
    keras.layers.BatchNormalization(),

    keras.layers.Conv2D(filters=512, kernel_size=(3, 3), activation='relu', padding="same"),
    keras.layers.BatchNormalization(),

    keras.layers.MaxPool2D(pool_size=(2, 2)),

    keras.layers.Conv2D(filters=512, kernel_size=(3, 3), activation='relu', padding="same"),
    keras.layers.BatchNormalization(),

    keras.layers.MaxPool2D(pool_size=(2, 2)),

    keras.layers.Flatten(),
    keras.layers.Dense(1024, activation='relu'),
    keras.layers.Dropout(0.5),
    keras.layers.Dense(1024, activation='relu'),
    keras.layers.Dropout(0.5),
    keras.layers.Dense(4, activation='softmax')
])

model.compile(
    loss='categorical_crossentropy',
    optimizer=tf.optimizers.SGD(learning_rate=0.001),
    metrics=['accuracy']
)

model.summary()

```

Model: "sequential"

Layer (type)	Output Shape	Param #
conv2d (Conv2D)	(None, 73, 73, 128)	24704
batch_normalization (Batch Normalization)	(None, 73, 73, 128)	512
conv2d_1 (Conv2D)	(None, 73, 73, 256)	819456
batch_normalization_1 (Batch Normalization)	(None, 73, 73, 256)	1024
max_pooling2d (MaxPooling2D)	(None, 24, 24, 256)	0
conv2d_2 (Conv2D)	(None, 24, 24, 256)	590080
batch_normalization_2 (Batch Normalization)	(None, 24, 24, 256)	1024
conv2d_3 (Conv2D)	(None, 24, 24, 256)	65792
batch_normalization_3 (Batch Normalization)	(None, 24, 24, 256)	1024
conv2d_4 (Conv2D)	(None, 24, 24, 256)	65792
batch_normalization_4 (Batch Normalization)	(None, 24, 24, 256)	1024
conv2d_5 (Conv2D)	(None, 24, 24, 512)	1180160
batch_normalization_5 (Batch Normalization)	(None, 24, 24, 512)	2048
max_pooling2d_1 (MaxPooling2D)	(None, 12, 12, 512)	0
conv2d_6 (Conv2D)	(None, 12, 12, 512)	2359808
batch_normalization_6 (Batch Normalization)	(None, 12, 12, 512)	2048
conv2d_7 (Conv2D)	(None, 12, 12, 512)	2359808
batch_normalization_7 (Batch Normalization)	(None, 12, 12, 512)	2048

```
history = model.fit(train, epochs=5, validation_data=val, verbose=1)
```

Epoch 1/5

WARNING:tensorflow:From C:\Users\Dell\OneDrive\Documents\ANACONDA\Lib\site-packages\keras\src\utils\tf_utils.py:492: The name tf.ragged.RaggedTensorValue is deprecated. Please use tf.compat.v1.ragged.RaggedTensorValue instead.

WARNING:tensorflow:From C:\Users\Dell\OneDrive\Documents\ANACONDA\Lib\site-packages\keras\src\engine\base_layer_utils.py:384: The name tf.executing_eagerly_outside_functions is deprecated. Please use tf.compat.v1.executing_eagerly_outside_functions instead.

```
996/996 [=====] - 4064s 4s/step - loss: 1.6087 - accuracy: 0.3605 - val_loss: 1.0419 - val_accuracy: 0.5341
Epoch 2/5
996/996 [=====] - 3582s 4s/step - loss: 1.1049 - accuracy: 0.5171 - val_loss: 0.8389 - val_accuracy: 0.6401
Epoch 3/5
996/996 [=====] - 3307s 3s/step - loss: 0.8307 - accuracy: 0.6457 - val_loss: 0.5835 - val_accuracy: 0.7440
Epoch 4/5
996/996 [=====] - 6200s 6s/step - loss: 0.5703 - accuracy: 0.7602 - val_loss: 0.3648 - val_accuracy: 0.8529
Epoch 5/5
996/996 [=====] - 9178s 9s/step - loss: 0.3828 - accuracy: 0.8507 - val_loss: 0.2819 - val_accuracy: 0.8901
```

```
history1 = model.fit(train, epochs=1, validation_data=val, verbose=1)
```

```
996/996 [=====] - 3392s 3s/step - loss: 0.2661 - accuracy: 0.8925 - val_loss: 0.2942 - val_accuracy: 0.8800
```

Testing Model & Data Prediction

Evaluating the model

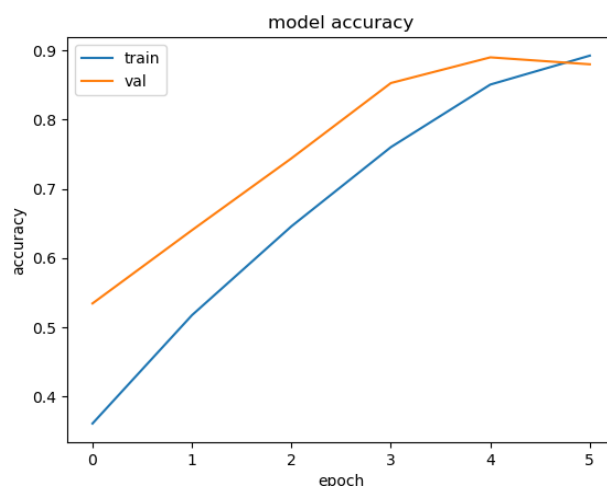
Here we have tested with the Mobilenet V2 Model With the help of the predict () function.

```
pred = model.predict(test)
pred = np.argmax(pred, axis=1) #pick class with highest probability

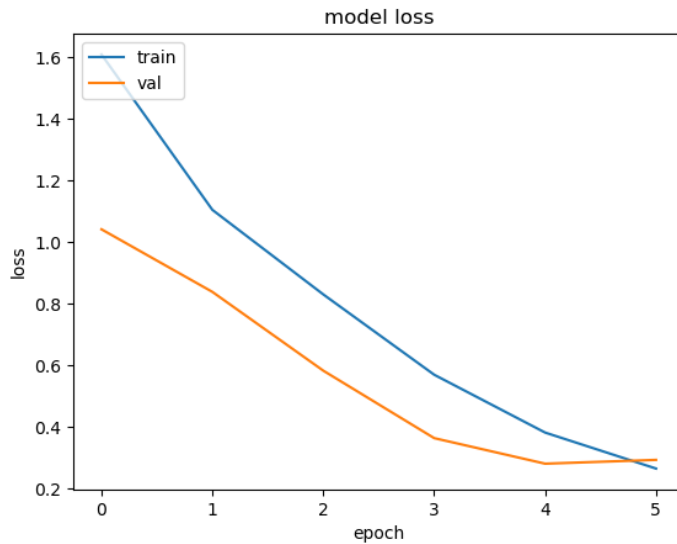
labels = (train.class_indices)
labels = dict((v,k) for k,v in labels.items())
pred2 = [labels[k] for k in pred]
```

```
374/374 [=====] - 332s 886ms/step
```

```
plt.plot(history.history['accuracy'] + history1.history['accuracy'])
plt.plot(history.history['val_accuracy'] + history1.history['val_accuracy'])
plt.title('model accuracy')
plt.ylabel('accuracy')
plt.xlabel('epoch')
plt.legend(['train', 'val'], loc='upper left')
plt.show()
```



```
plt.plot(history.history['loss'] + history1.history['loss'])
plt.plot(history.history['val_loss'] + history1.history['val_loss'])
plt.title('model loss')
plt.ylabel('loss')
plt.xlabel('epoch')
plt.legend(['train', 'val'], loc='upper left')
plt.show()
```



```
from sklearn.metrics import confusion_matrix, accuracy_score
from sklearn.metrics import classification_report

y_test = test_images.labels # set y_test to the expected output
print(classification_report(y_test, pred2))
print("Accuracy of the Model:", "{:.1f}%".format(accuracy_score(y_test, pred2)*100))
```

	precision	recall	f1-score	support
eosinophil	0.82	0.81	0.82	725
lymphocyte	0.90	0.99	0.94	762
monocyte	0.98	0.96	0.97	759
neutrophil	0.87	0.80	0.83	742
accuracy			0.89	2988
macro avg	0.89	0.89	0.89	2988
weighted avg	0.89	0.89	0.89	2988

Accuracy of the Model: 89.3%


```

import matplotlib.pyplot as plt
import seaborn as sns
from sklearn.metrics import confusion_matrix

class_labels = ['EOSINOPHIL', 'LYMPHOCYTE', 'MONOCYTE', 'NEUTROPHIL']

cm = confusion_matrix(y_test, pred2)

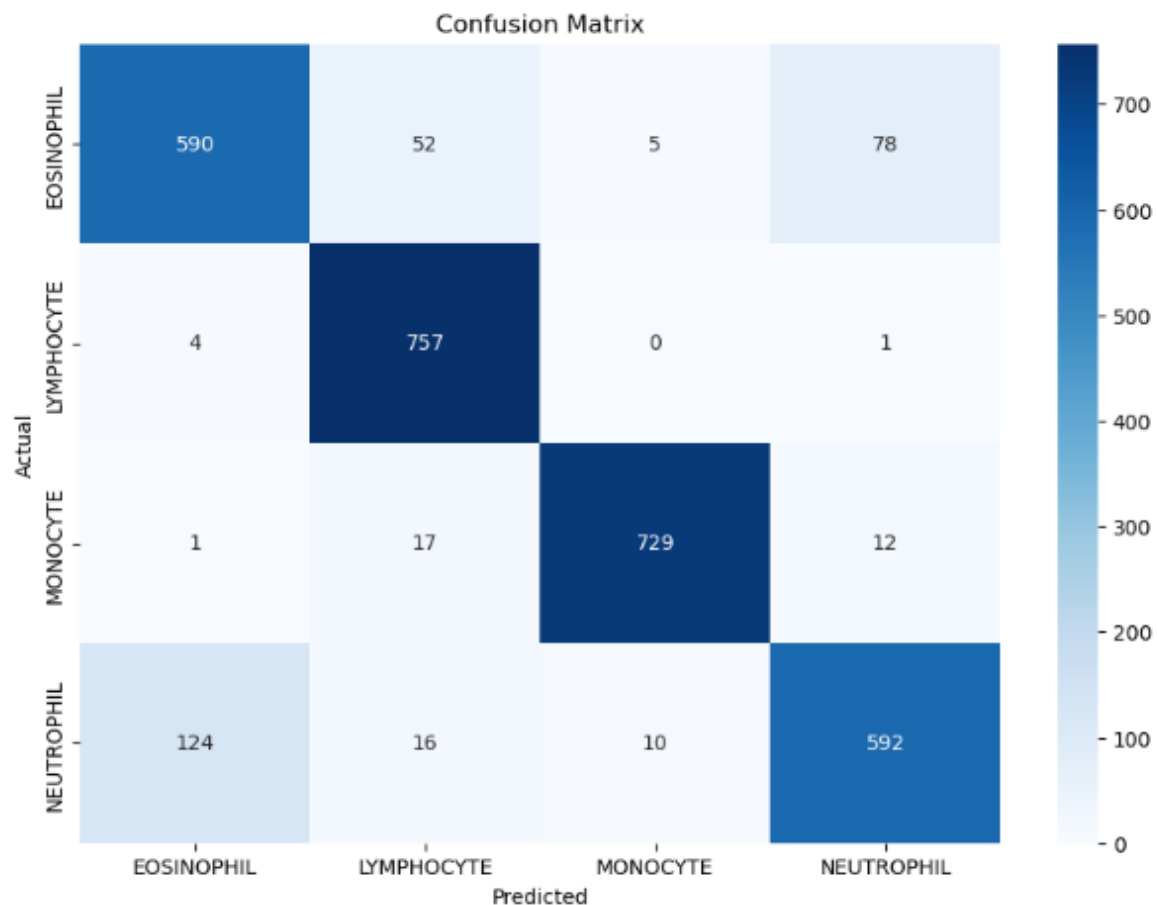
plt.figure(figsize=(10, 7))
sns.heatmap(cm, annot=True, fmt='g', vmin=0, cmap='Blues')

plt.xticks(ticks=[0.5, 1.5, 2.5, 3.5], labels=class_labels)
plt.yticks(ticks=[0.5, 1.5, 2.5, 3.5], labels=class_labels)
plt.xlabel("Predicted")
plt.ylabel("Actual")

plt.title("Confusion Matrix")

plt.show()

```



Saving the model

Saving the model

Finally, we have chosen the best model now saving that model

```
model.save("Blood Cell.h5")
```

Application Building

In this section, we will be building a web application that is integrated into the model we built. A UI is provided for the user where he has to enter the values for predictions. The entered values are given to the saved model and prediction is showcased on the UI.

This section has the following tasks

- Building HTML Pages
- Building server-side script

Building HTML Pages:

For this project create three HTML files namely

- home.html
- result.html

• Build Python code:

- Import the libraries

```
import os
import numpy as np
import cv2
from flask import Flask, request, render_template, redirect, url_for
from tensorflow.keras.models import load_model
from tensorflow.keras.applications.mobilenet_v2 import preprocess_input
import matplotlib.pyplot as plt
import io
import base64

app = Flask(__name__)
model = load_model("Blood_Cell.h5")
class_labels = ['eosinophil', 'lymphocyte', 'monocyte', 'neutrophil']
```

- Load the saved model. Importing the Flask module in the project is mandatory. An object of the Flask class is our WSGI application. The Flask constructor takes the name of the current module (__name__) as argument.
- Here we will be using the declared constructor to route to the HTML page which we have created earlier.
- In the above example, the '/' URL is bound with the index.html function. Hence, when the index page of the web server is opened in the browser, the html page will be rendered. Whenever you enter the values from the html page the values can be retrieved using POST Method.
- Retrieves the value from UI:

```

app = Flask(__name__)
model = load_model("Blood_Cell.h5")
class_labels = ['eosinophil', 'lymphocyte', 'monocyte', 'neutrophil']

def predict_image_class(image_path, model):
    img = cv2.imread(image_path)
    img_rgb = cv2.cvtColor(img, cv2.COLOR_BGR2RGB)
    img_resized = cv2.resize(img_rgb, (224, 224))
    img_preprocessed = preprocess_input(img_resized.reshape((1, 224, 224, 3)))
    predictions = model.predict(img_preprocessed)
    predicted_class_idx = np.argmax(predictions, axis=1)[0]
    predicted_class_label = class_labels[predicted_class_idx]
    return predicted_class_label, img_rgb

@app.route("/", methods=["GET", "POST"])
def upload_file():
    if request.method == "POST":
        if "file" not in request.files:
            return redirect(request.url)
        file = request.files["file"]
        if file.filename == "":
            return redirect(request.url)
        if file:
            file_path = os.path.join("static", file.filename)
            file.save(file_path)
            predicted_class_label, img_rgb = predict_image_class(file_path, model)

            # Convert image to string for displaying in HTML
            _, img_encoded = cv2.imencode('.png', cv2.cvtColor(img_rgb, cv2.COLOR_RGB2BGR))
            img_str = base64.b64encode(img_encoded).decode('utf-8')

            return render_template("result.html", class_label=predicted_class_label, img_data=img_str)
    return render_template("home.html")

```

- Here we are routing our app to the output() function. This function retrieves all the values from the HTML page using a Post request. That is stored in an array. This array is passed to the model. Predict () function. This function returns the prediction. This prediction value will rendered to the text that we have mentioned in the output.html page earlier.
- Main Function:

```

if __name__ == "__main__":
    app.run(debug=True)

```

Run the web application

- : Run the application
 - Open Anaconda prompt from the start menu
 - Navigate to the folder where your Python script is.
 - Now type the “app.py” command
 - Navigate to the local host where you can view your web page.

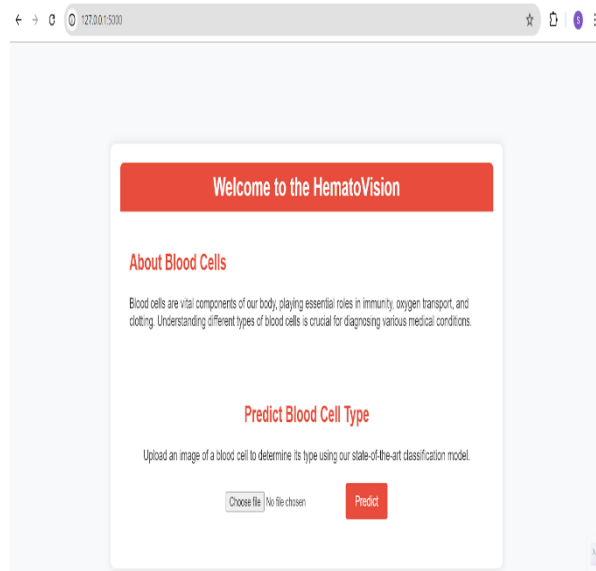
- Click on the inspect button from the top right corner, enter the inputs, click on the predict button, and see the result/prediction on the web.

```
* 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.
* Running on http://127.0.0.1:5000
Press CTRL+C to quit
* Restarting with watchdog (windowsapi)
```

Now, Go the web browser and write the localhost url (http://127.0.0.1:5000) to get the below results

- UI Image preview:

Let's see what our index.html page looks like:



By clicking on choose file it will ask us to upload the image , then by clicking on the predict button , it will take us to the result.html

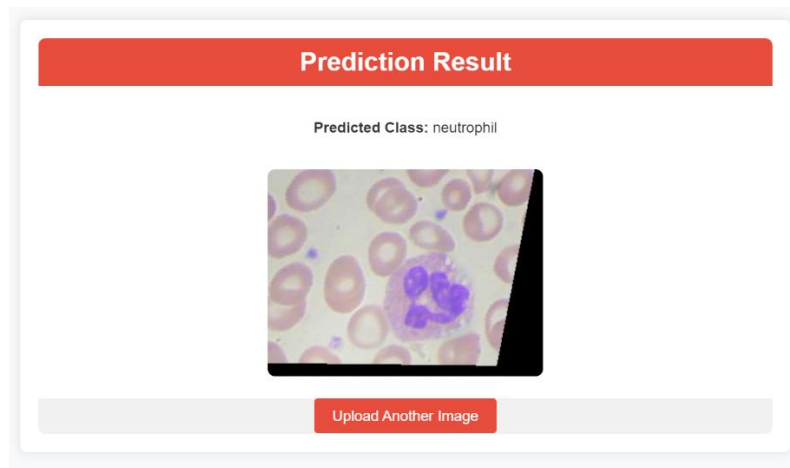
Test For Class-1 : Neutrophil

Predict Blood Cell Type

Upload an image of a blood cell to determine its type using our state-of-the-art classification model.

Choose file _8_9488.jpeg

Predict



Test For Class-2 : Monocyt

Predict Blood Cell Type

Upload an image of a blood cell to determine its type using our state-of-the-art classification model.

_3_9423.jpeg

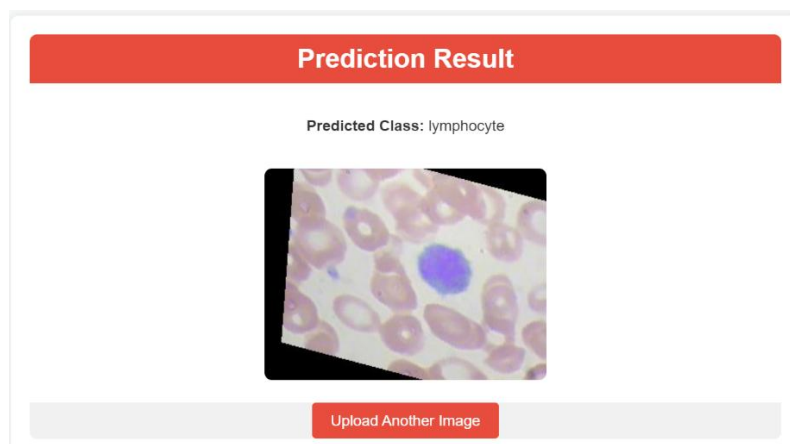


Test For Class-3 : Lymphocyte

Predict Blood Cell Type

Upload an image of a blood cell to determine its type using our state-of-the-art classification model.

_5_9201.jpeg



Test For Class-4 : Eosinophil

Predict Blood Cell Type

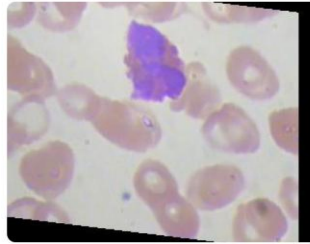
Upload an image of a blood cell to determine its type using our state-of-the-art classification model.

Choose file _3_9885.jpeg

Predict

Prediction Result

Predicted Class: eosinophil



Upload Another Image