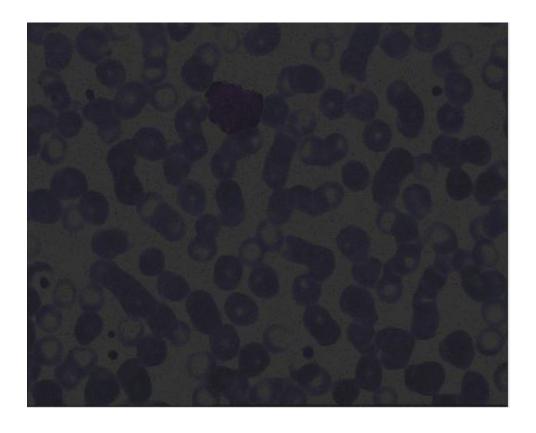
Part 1: Extraction of White Blood Cells

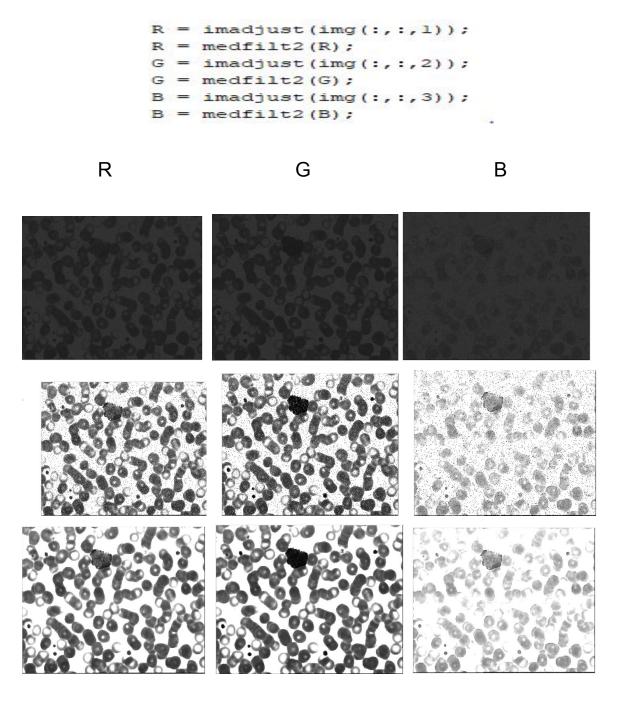
1) Iterate through the folders to process each image

```
Folder = dir('output_pre\');
Folder2 = dir('output\');
totalSimilarity=0;
total=0;
for j = 1 : 5
    File = dir(strcat('output_pre\',int2str(j),'\*.bmp'));
    File2 = dir(strcat('output\',int2str(j),'\*.bmp'));
    for i = 1 : length(File)
        filename2 = strcat('output\',int2str(j),'\',int2str(i),'.bmp');
        groundTruth = imread(filename2);
        filename = strcat('output_pre\',int2str(j),'\',int2str(i),'.bmp');
        img = imread(filename);
```

2) Example: Fist Basophil RGB image

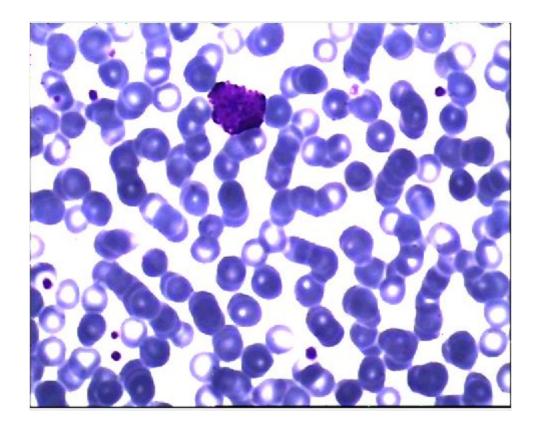


3) As we can see the image suffers from low contrast and very noisy so we will split the image into the 3 channels (R, G, B) and apply imadjust function and median filter to each channel



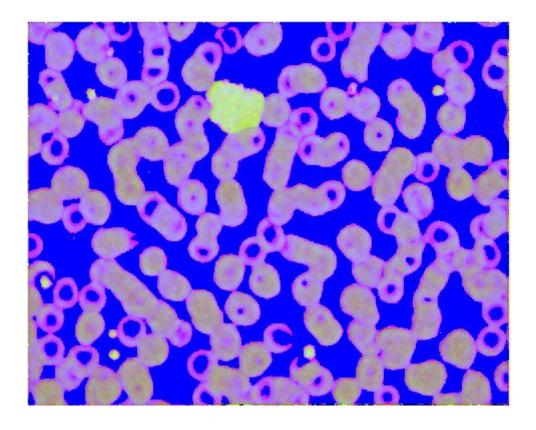
4) Concatenate again the 3 channels to display the processed RGB image

$$RGB = cat(3,R,G,B);$$

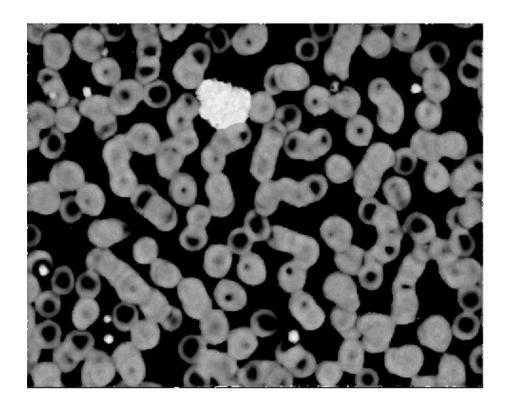


```
HSV = rgb2hsv(im2double(RGB));
sThresh = HSV(:,:,2) > 0.7 & HSV(:,:,2) < 1;</pre>
```

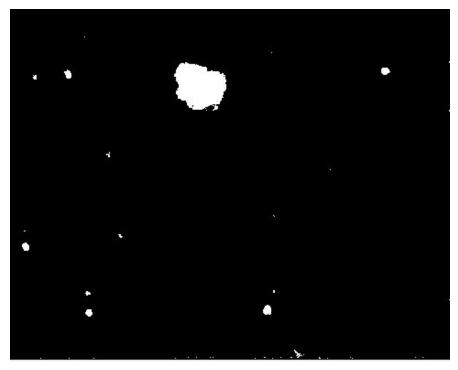
5) Convert the RGB image to HSV image



6) The saturation channel appears to be a good channel for segmenting the purple regions



7) Threshold the intensities less than 1 and greater than 0.7



8) As we can see there are a lot of noise that have been segmented with the required region so we will extract the larger regions only

final1 = bwareafilt(sThresh, [480, 100000000]);



9) Other images have some holes after the last step so we will apply dilation to fill those holes

```
se = strel('disk',1,0);
final2 = imdilate(final1,se);
```



10) We are going to calculate the similarity between the output images and the ground truth using Jaccard index

```
sim = getJaccard(final2,groundTruth);
totalSimilarity = totalSimilarity + sim;
total = total+1;
```

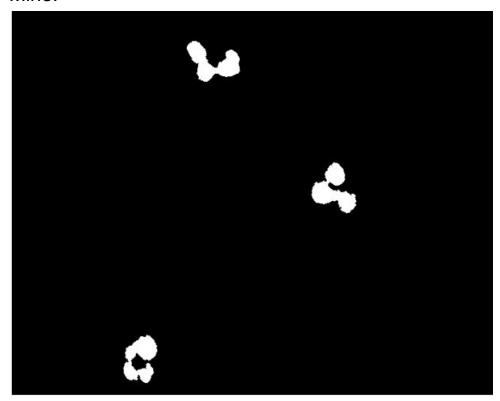
11) Jaccard Function

```
function J = getJaccard(A,B)

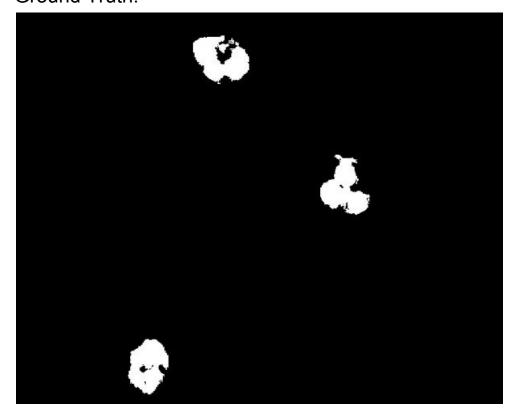
J = sum(A(:) & B(:))/sum(A(:) | B(:));
```

12) Sample from the dataset

Mine:



Ground Truth:



13) Output images will be saved in folders like the input

```
fileName = sprintf('%d.bmp',i);
folder = strcat('myOutput\',int2str(j));
fullFileName = fullfile(folder,fileName);
imwrite(final2,fullFileName);
```

Part 2: Classification of White Blood Cells

This is a multi-class classification problem, meaning that there are more than two classes to be predicted, in fact there are five types of White Blood Cells. I am going to classify then using VGG19 CNN pretrained model implemented with keras.

1) Batch size = 1 (Every image will update the weights)

```
train_data_dir = '/content/drive/MyDrive/WhiteBloodCellsData'
validation_data_dir = '/content/drive/MyDrive/WhiteBloodCellsData'
batch_size=1
img_width,img_height =500,500
```

2) Data Augmentation (to artificially expand the size of a training dataset by creating modified versions of images in the dataset.) Note: Split the dataset into 70% Training Data and 30% Validation Data

```
datagen = ImageDataGenerator(
    rescale = 1./255,
    shear_range=0.3,
    horizontal_flip=True,
    rotation_range = 2,
    width_shift_range = 0.3,
    height_shift_range = 0.3,
    validation_split = 0.3
)
```

3) The training dataset (171 images)

Found 171 images belonging to 5 classes.

4) The validation dataset (70 images)

Found 70 images belonging to 5 classes.

5) Import VGG19 pretrained model trained on imagenet dataset

```
[9] base_model = VGG19(include_top = False, weights = 'imagenet', input_shape = (500,500,3))
```

6) Create a new model and add VGG19 to it

```
[10] model= Sequential()
    model.add(base model)
    model.add(Flatten())
[11] model.summary()
    Model: "sequential"
    Layer (type)
                                  Output Shape
                                                             Param #
    vgg19 (Functional)
                                   (None, 15, 15, 512)
                                                             20024384
    flatten (Flatten)
                                   (None, 115200)
    Total params: 20,024,384
    Trainable params: 20,024,384
    Non-trainable params: 0
```

7) Add some fully connected layers after VGG19

```
model.add(Dense(1024,activation=('relu'),input dim=512))
   model.add(Dense(512,activation=('relu')))
   model.add(Dropout(.4))
   model.add(Dense(256,activation=('relu')))
   model.add(Dropout(.3))
   model.add(Dense(128,activation=('relu')))
   model.add(Dense(5,activation=('softmax')))
   model.summary()
Model: "sequential"
   Layer (type)
                                 Output Shape
                                                            Param #
   vgg19 (Functional)
                                 (None, 15, 15, 512)
                                                            20024384
   flatten (Flatten)
                                  (None, 115200)
   dense (Dense)
                                  (None, 1024)
                                                            117965824
   dense 1 (Dense)
                                  (None, 512)
                                                            524800
   dropout (Dropout)
                                  (None, 512)
                                                            Θ
   dense 2 (Dense)
                                  (None, 256)
                                                            131328
                                  (None, 256)
   dropout 1 (Dropout)
   dense 3 (Dense)
                                  (None, 128)
                                                            32896
   dense 4 (Dense)
                                  (None, 5)
                                                            645
   Total params: 138,679,877
   Trainable params: 138,679,877
   Non-trainable params: 0
```

8) Compiling the model and choosing the optimizer

```
batch_size= batch_size
epochs=500
learn_rate=.00001
sgd=SGD(lr=learn_rate,momentum=.9,nesterov=False)
adam=Adam(lr=learn_rate, beta_1=0.9, beta_2=0.999, epsilon=None, decay=0.0, amsgrad=False)
model.compile(optimizer=sgd,loss='categorical_crossentropy',metrics=['accuracy'])
```

9) Create function to stop the training process when a chosen validation accuracy is met

```
[14] import tensorflow as tf
    class MyThresholdCallback(tf.keras.callbacks.Callback):
        def __init__(self, threshold):
            super(MyThresholdCallback, self).__init__()
            self.threshold = threshold

        def on_epoch_end(self, epoch, logs=None):
        val_acc = logs["val_accuracy"]
        if val_acc >= self.threshold:
            self.model.stop_training = True
```

10) Start training the model and stop the training process when validation accuracy is equal 80%

11) The model stopped training when accuracy is met stopped in epoch 237(Last 4 epochs)

12) Training Accuracy Vs Validation Accuracy

```
#Accuracy
ax[1].plot(model.history.history['accuracy'],color='b',label='Training Accuracy')
ax[1].plot(model.history.history['val_accuracy'],color='r',label='Validation Accuracy')

[<matplotlib.lines.Line2D at 0x7fc2ce760828>]

09
08
07
06
05
04
03
02
05
01
00
150
200
```

13) And finally the model succeeded to recognise the type of this white blood cell (0 = Basophil)

/usr/local/lib/python3.6/dist-packages/tensorflow/ş
 warnings.warn('`model.predict_classes()` is depre
[0]

