# **Malaria Bounding Boxes**

#### **Final Notebook**

```
In [0]:
```

```
#Mouning Google Drive
from google.colab import drive
drive.mount('/content/drive', force_remount=True)
```

Go to this URL in a browser: https://accounts.google.com/o/oauth2/auth?client\_id=947318989803-6bn6 qk8qdgf4n4g3pfee6491hc0brc4i.apps.googleusercontent.com&redirect\_uri=urn%3aietf%3awg%3aoauth%3a2.0%b&response\_type=code&scope=email%20https%3a%2f%2fwww.googleapis.com%2fauth%2fdocs.test%20https%3a%2f%2fwww.googleapis.com%2fauth%2fdrive.photos.readonlyttps%3a%2f%2fwww.googleapis.com%2fauth%2fdrive.photos.readonlyttps%3a%2f%2fwww.googleapis.com%2fauth%2fdrive.photos.readonly

Enter your authorization code:
.....
Mounted at /content/drive

..... Þ

In [0]:

```
import cv2
import numpy as np
import pandas as pd
from sklearn.metrics import average_precision_score
from tqdm import tqdm
import pandas as pd
import matplotlib.pyplot as plt
from matplotlib import patches
import numpy as np
from os import path
import pandas as pd
import matplotlib.pyplot as plt
from matplotlib import patches
import numpy as np
from os import path
from tqdm import tqdm
import tensorflow.python.keras
from tensorflow.python.keras import models, layers
from tensorflow.python.keras.layers import SeparableConv2D, DepthwiseConv2D
from tensorflow.python.keras.models import Model, load_model
from tensorflow.python.keras.layers import BatchNormalization, Activation, Flatten
from tensorflow.python.keras.optimizers import Adam
from keras.preprocessing.image import ImageDataGenerator
```

The default version of TensorFlow in Colab will soon switch to TensorFlow 2.x.

We recommend you upgrade now or ensure your notebook will continue to use TensorFlow 1.x via the %tensorflow version

1.x magic: more info.

Using TensorFlow backend.

# **Mean Average Precision**

```
In [0]:
```

```
def map_value(annotate_path, result_df_path):
    """
    This Function takes annotations file path and path of result dataframe which genereates after ec
    aluating test images
    """

def get_data(input_path):
    """
    This Function takes annotations file path and gives a dictionory of ground truth labels and bou
    nding boxes coordinates
```

```
.....
 img data = []
 with open (input path, 'r') as f:
  for line in f:
   line split = line.strip().split(',')
   (filename, x1, y1, x2, y2, class_name) = line_split
   img data.append({'class': class name, 'x1': int(x1), 'x2': int(x2), 'y1': int(y1), 'y2': int(y2)
) })
 return img data
def union(au, bu, area intersection):
  This function gives the area of union for intersection over union calculation
 area a = (au[2] - au[0]) * (au[3] - au[1])
 area_b = (bu[2] - bu[0]) * (bu[3] - bu[1])
 area_union = area_a + area_b - area_intersection
 return area union
def intersection(ai, bi):
  This function gives the area of intersection for intersection over union calculation
 11 11 11
 x = max(ai[0], bi[0])
 y = max(ai[1], bi[1])
 w = min(ai[2], bi[2]) - x
 h = min(ai[3], bi[3]) - y
 if w < 0 or h < 0:
  return 0
 return w*h
def iou(a, b):
 \# a and b should be (x1,y1,x2,y2)
  This function takes two boxes cordinates (ground truth and predicted) and gives intersection ove
r union of 2 boxes
 if a[0] >= a[2] or a[1] >= a[3] or b[0] >= b[2] or b[1] >= b[3]:
 area i = intersection(a, b)
 area u = union(a, b, area i)
return float(area i) / float(area u + 1e-6)
gt = get_data(annotate path)
pred = pd.read csv(result df path)
pred.columns = ["name", "class", "x1","y1","x2","y2", "prob"]
Truth = {}
Predicted = {}
for bbox in gt:
 bbox['bbox_matched'] = False
pred probs = np.array(pred["prob"])
box_idx_sorted_by_prob = np.argsort(pred_probs)[::-1]
 # we compare each predicted bounding box with ground truth bounding box
for box idx in tqdm(box idx sorted by prob):
 pred box = pred.loc[box idx]
 pred_class = pred_box['class']
 pred_x1 = pred_box['x1']
 pred_x2 = pred_box['x2']
 pred_y1 = pred_box['y1']
 pred_y2 = pred_box['y2']
 pred prob = pred box['prob']
 if pred_class not in Predicted:
  Predicted[pred class] = []
  Truth[pred class] = []
```

```
Predicted[pred class].append(pred prob)
 found_match = False
 for gt box in gt:
  gt class = gt box['class']
  gt_x1 = gt_box['x1']
  gt x2 = gt box['x2']
  gt y1 = gt box['y1']
  gt_y2 = gt_box['y2']
  gt seen = gt box['bbox matched']
  if gt_class != pred_class:
   continue
  if gt seen:
   continue
  iou1 = iou((pred x1, pred y1, pred x2, pred y2), (gt x1, gt y1, gt x2, gt y2))
  # taking 0.5 as threshold i.e. if iou is greater than 0.5 then only it will consider as correct
matching
  if iou1 >= 0.5:
   found match = True
   gt box['bbox matched'] = True
   break
  else:
   continue
 Truth[pred class].append(int(found match))
for gt_box in gt:
 if not gt box['bbox matched']:
  if gt box['class'] not in Predicted:
   Predicted[gt box['class']] = []
   Truth[gt box['class']] = []
  Truth[gt box['class']].append(1)
  Predicted[gt box['class']].append(0)
t = \{ \}
g = g
for key in Truth.keys():
  if key not in t:
   t[key] = []
   p[key] = []
  t[key].extend(Truth[key])
  p[key].extend(Predicted[key])
all_aps = []
for key in t.keys():
 ap = average_precision_score(t[key], p[key])
 print('{} AP: {}'.format(key, ap))
 all_aps.append(ap)
print('mAP = {}'.format(np.mean(np.array(all_aps))))
```

# loading second level classifier best weights

```
In [0]:
```

```
def denseblock(input, num_filter = 12, dropout_rate = 0.5):
    global compression
    temp = input
    for in range(1):
        BatchNorm = layers.BatchNormalization()(temp)
        relu = layers.Activation('relu')(BatchNorm)
        Conv2D 3 3 = layers.SeparableConv2D(int(num filter*compression), (5,5), use bias=False ,pad
ding='same') (relu)
       if dropout rate>0:
            Conv2D 3 3 = layers.Dropout(dropout rate)(Conv2D 3 3)
        concat = layers.Concatenate(axis=-1)([temp,Conv2D 3 3])
        temp = concat
    return temp
## transition Blosck
def transition(input, num filter = 12, dropout rate = 0.5):
    global compression
    BatchNorm = lavers.BatchNormalization()(input)
```

```
relu = layers.Activation('relu')(BatchNorm)
    Conv2D BottleNeck = layers.SeparableConv2D(int(num filter*compression), (7,7), use bias=False,p
adding='same') (relu)
   if dropout rate>0:
        Conv2D BottleNeck = layers.Dropout(dropout rate)(Conv2D BottleNeck)
    avg = layers.AveragePooling2D(pool size=(2,2))(Conv2D BottleNeck)
    return ava
#output layer
def output layer(input):
    global compression
    BatchNorm = layers.BatchNormalization()(input)
    relu = layers.Activation('relu')(BatchNorm)
    AvgPooling = layers.AveragePooling2D(pool_size=(2,2))(relu)
    flat = layers.Flatten()(AvgPooling)
    output = layers.Dense(num classes, activation='softmax')(flat)
    return output
num filter = 64
batch size = 32
num classes = 6
epochs = 100
1 = 10
compression = 0.45
dropout_rate = 0
input = layers.Input(shape=(32, 32, 3,))
First Conv2D = layers.SeparableConv2D(num filter, (5,5), use bias=False, padding='same')(input)
First Block = denseblock(First Conv2D, num filter, dropout rate)
First Transition = transition(First Block, num filter, dropout rate)
Second_Block = denseblock(First_Transition, num_filter, dropout_rate)
Second Transition = transition(Second Block, num filter, dropout rate)
Third Block = denseblock(Second Transition, num_filter, dropout_rate)
Third Transition = transition(Third Block, num filter, dropout rate)
Last Block = denseblock(Third Transition, num filter, dropout rate)
output = output layer(Last Block)
model = Model(inputs=[input], outputs=[output])
model.load weights ("/content/drive/My Drive/rcnn/second classifier checkpoint/epochs:030-
val acc:0.715.hdf5")
WARNING:tensorflow:From /usr/local/lib/python3.6/dist-
packages/tensorflow core/python/ops/resource variable ops.py:1630: calling
```

packages/tensorflow\_core/python/ops/resource\_variable\_ops.py:1630: calling
BaseResourceVariable.\_\_init\_\_ (from tensorflow.python.ops.resource\_variable\_ops) with constraint i
s deprecated and will be removed in a future version.
Instructions for updating:
If using Keras pass \* constraint arguments to layers.

# **Function to print images**

In [0]:

```
def predicted(image_name):
    """
    This function takes imag name and prints original image, predicted images of first lavel classifier and second level classifier

    """
    #print("Original Image with ground truth bounding boxes and labels")

    fig = plt.figure(figsize=(12,6))
    ax = fig.add_axes([0,0,1,1])
    ax.set_title("Original Image with ground truth bounding boxes and labels")

# read and plot the image
    image = plt.imread('/content/drive/My Drive/rcnn/test_images/'+image_name)
    plt.imshow(image)
    edgecolor = 'r'
    test_df = pd.read_csv("/content/drive/My Drive/rcnn/test_df.csv")
# iterating over the image for different objects
    for _,row in test_df[test_df["image"]=="/images/"+image_name].iterrows():
        vmin = row v min
```

```
VIIITII - TOM *V IIITII
       #print(xmin)
       xmax = row.x max
       ymin = row.y_min
       ymax = row.y_max
       width = xmax - xmin
      height = ymax - ymin
      # assign different color to different classes of objects
       if row.class label == 'red blood cell':
           edgecolor = 'r'
           ax.annotate('RBC', xy=(xmax-40,ymin+20))
       elif row.class label == 'trophozoite':
           edgecolor = 'b'
           ax.annotate('trophozoite', xy=(xmax-40,ymin+20))
       elif row.class label == 'difficult':
           edgecolor = 'b'
           ax.annotate('difficult', xy=(xmax-40,ymin+20))
       elif row.class label == 'ring':
           edgecolor = 'b'
           ax.annotate('ring', xy=(xmax-40,ymin+20))
       elif row.class label == 'schizont':
           edgecolor = 'b'
           ax.annotate('schizont', xy=(xmax-40,ymin+20))
       elif row.class label == 'gametocyte':
           edgecolor = 'b'
           ax.annotate('gametocyte', xy=(xmax-40,ymin+20))
       elif row.class label == 'leukocyte':
           edgecolor = 'b'
           ax.annotate('leukocyte', xy=(xmax-40,ymin+20))
      # add bounding boxes to the image
       rect = patches.Rectangle((xmin,ymin), width, height, edgecolor = edgecolor, facecolor = 'non
e')
       ax.add patch (rect)
   #print("Prediction after first classifier(which classifies RBC and OTHER)")
   fig = plt.figure(figsize=(12,6))
   ax = fig.add axes([0,0,1,1])
   ax.set title("Prediction after first classifier(which classifies RBC and OTHER)")
   # read and plot the image
  image = plt.imread('/content/drive/My Drive/rcnn/test images/' + image name)
   plt.imshow(image)
  result df 2class = pd.read csv("/content/drive/My Drive/rcnn/Data model 2/result df 2class.csv"
  # iterating over the image for different objects
   for ,row in result df 2class.loc[result df 2class["name"]==image name].iterrows():
       xmin = row.x1
       #print(xmin)
       xmax = row.x2
       ymin = row.y1
       ymax = row.y2
       width = xmax - xmin
      height = ymax - ymin
       # assign different color to different classes of objects
       if row.label == 'other':
           edgecolor = 'b'
           ax.annotate('other', xy=(xmax-40,ymin+20))
       elif row.label == 'RBC':
           edgecolor = 'r'
           ax.annotate('RBC', xy=(xmax-40,ymin+20))
       # add bounding boxes to the image
       rect = patches.Rectangle((xmin,ymin), width, height, edgecolor = edgecolor, facecolor = 'non
e')
       ax.add patch (rect)
  testing image = result df 2class[result df 2class["label"] == "other"][result df 2class["name"] == i
mage_name]
   testing_image.reset_index(inplace=True)
   if (tacting image chang[]] |= ().
```

```
import cv2
    from tqdm import tqdm
    testing_img_arrays = []
    V = []
    WIDTH = 32
    HEIGHT = 32
    for i in range(testing image.shape[0]):
      image = plt.imread('/content/drive/My Drive/rcnn/test images/'+image name)
       image_crop = image[testing_image["y1"][i]:testing_image["y2"][i],testing_image["x1"][i]:test
ing image["x2"][i]]
      testing img arrays.append(cv2.resize(image crop, (WIDTH, HEIGHT), interpolation=cv2.INTER CUB
IC))
    testing img arrays final = np.array(testing img arrays)
    model.load_weights("/content/drive/My Drive/rcnn/second_classifier_checkpoint/epochs:030-
val acc:0.715.hdf5")
    model_prediction = model.predict(testing_img_arrays_final)
    result label = np.argmax(model_prediction, axis=1)
    feature names = ['difficult', 'gametocyte', 'leukocyte', 'ring', 'schizont', 'trophozoite']
    fig = plt.figure(figsize=(12,6))
    ax = fig.add axes([0,0,1,1])
    # read and plot the image
    image = plt.imread('/content/drive/My Drive/rcnn/test images/'+image name)
    plt.imshow(image)
    ax.set title("prediction after second classifier(final prediction)")
    # iterating over the image for different objects
    for _,row in result_df_2class[result_df_2class["name"] == image_name].iterrows():
        xmin = row.x1
         #print(xmin)
         xmax = row.x2
        ymin = row.y1
        ymax = row.y2
        width = xmax - xmin
        height = ymax - ymin
         # assign different color to different classes of objects
        if row.label == 'other':
            edgecolor = 'b'
            ax.annotate(feature names[result label[i]], xy=(xmax-40,ymin+20))
            i = i+1
         elif row.label == 'RBC':
            edgecolor = 'r'
            ax.annotate('RBC', xy=(xmax-40,ymin+20))
        # add bounding boxes to the image
        rect = patches.Rectangle((xmin,ymin), width, height, edgecolor = edgecolor, facecolor = 'n
one')
         ax.add patch (rect)
```

#### **Final Results**

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#### map of Test Data

```
In [0]:
```

```
print("MAP of test DATA:")
map_value("/content/drive/My Drive/rcnn/annotate_test_2class.txt","/content/drive/My
Drive/rcnn/Data_model_2/result_df_2class.csv")
```

MAP of test DATA:

```
100%| 9195/9195 [01:22<00:00, 111.75it/s]

other AP: 0.960051090010949

RBC AP: 0.998090609353597

mAP = 0.979070849682273
```

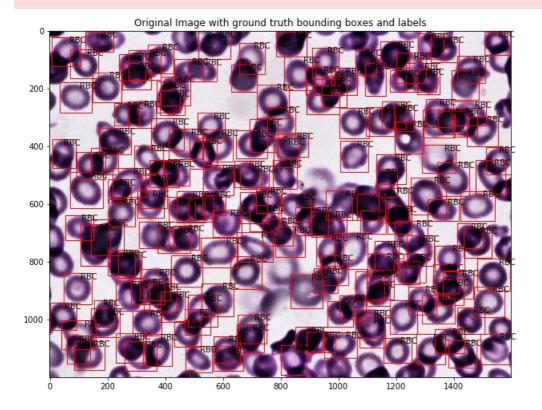
# **Image Predictions**

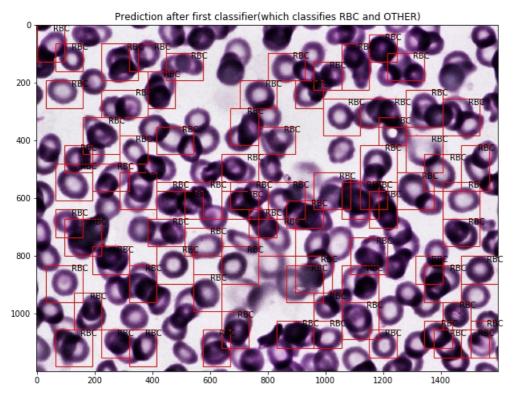
• sample\_image:1 Image with more nubmer of cells

#### In [0]:

predicted("46a2d369-4bea-4405-ac78-91bbe130b9a7.png")

/usr/local/lib/python3.6/dist-packages/ipykernel\_launcher.py:85: UserWarning: Boolean Series key w ill be reindexed to match DataFrame index.



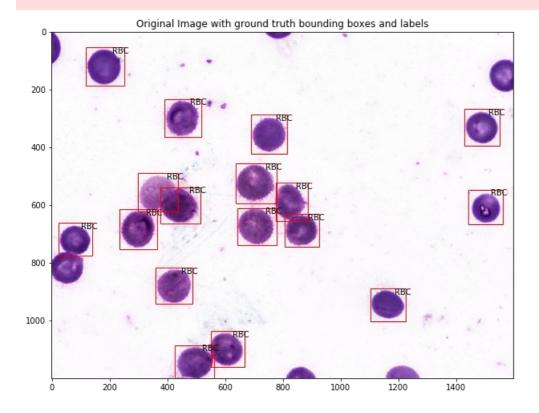


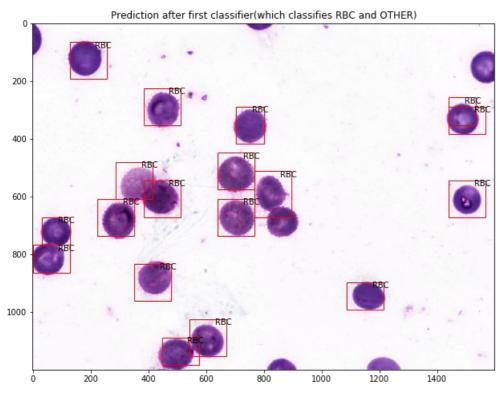
# • sample\_image:2 Image with very few cells

# In [0]:

predicted("c3209548-7f7c-4afd-b47c-b42a72b1ebf7.png")

/usr/local/lib/python3.6/dist-packages/ipykernel\_launcher.py:85: UserWarning: Boolean Series key w ill be reindexed to match DataFrame index.

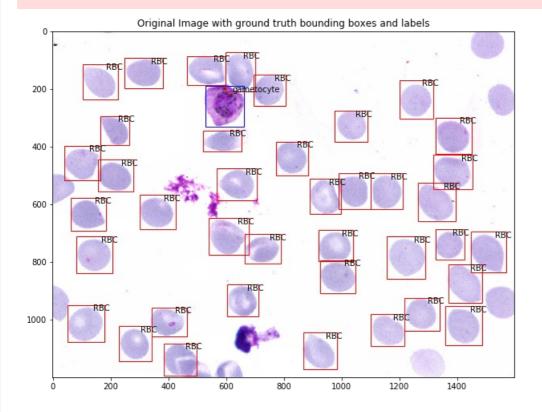


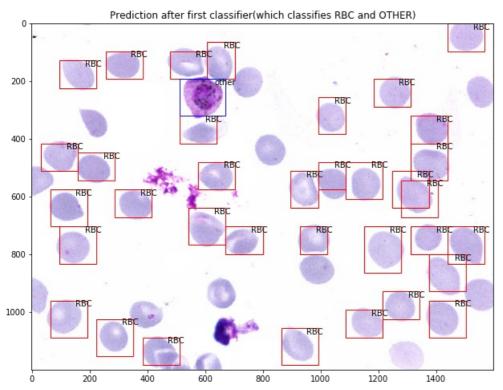


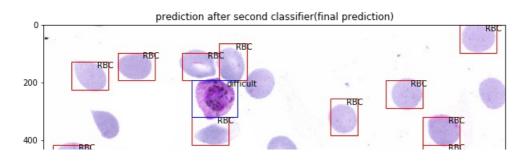
• sample\_image:3

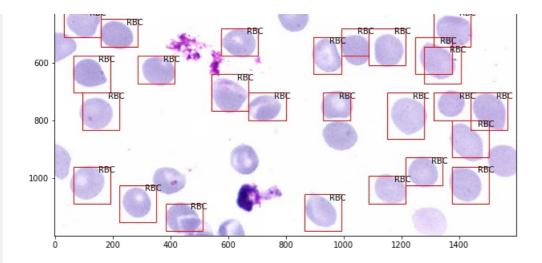
predicted("04de27ff-0d36-4da3-a829-e93851d76981.png")

/usr/local/lib/python3.6/dist-packages/ipykernel\_launcher.py:85: UserWarning: Boolean Series key w ill be reindexed to match DataFrame index.







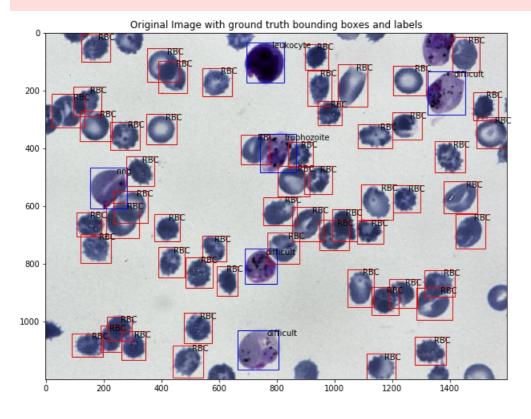


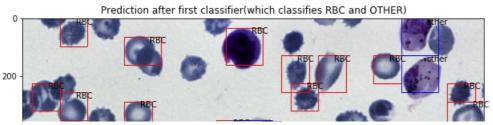
- First level classifier predicting correctly and in this image we have "gametocyte" but our model is predicting as difficult, but difficult class consists of the cells which are difficult to classify. So if our model predict class as difficult we have to manually classify the cell.
- sample\_image:4

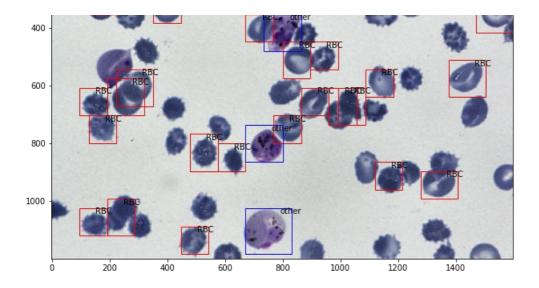
### In [0]:

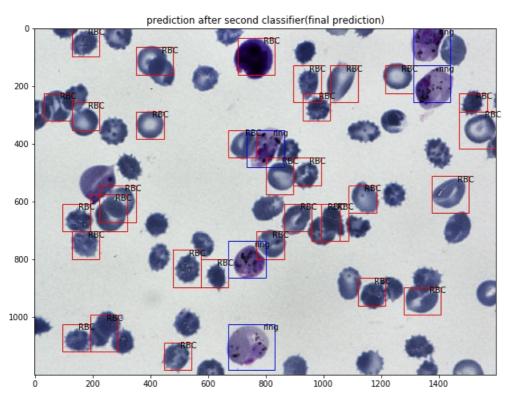
predicted("be6fe358-dee1-4dcf-9b0d-5d8612aba577.png")

/usr/local/lib/python3.6/dist-packages/ipykernel\_launcher.py:85: UserWarning: Boolean Series key w ill be reindexed to match DataFrame index.







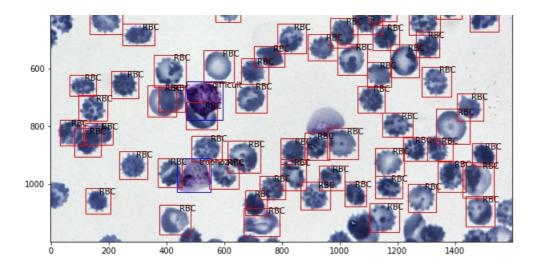


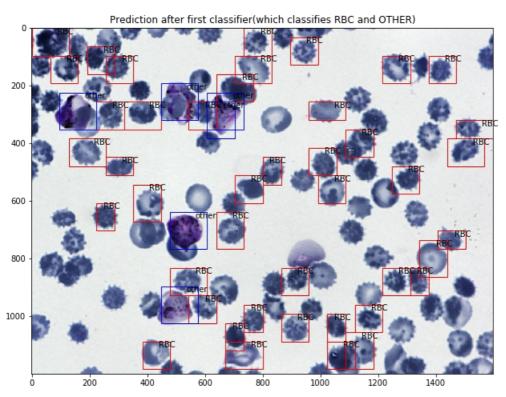
- in the Actual image we have class difficult but our model predict them as ring. This may be correct prediction.
- Model could not able to predict the uninfected cell "Leuckocyte". This is the failure of first level classifier.
- sample\_image:5

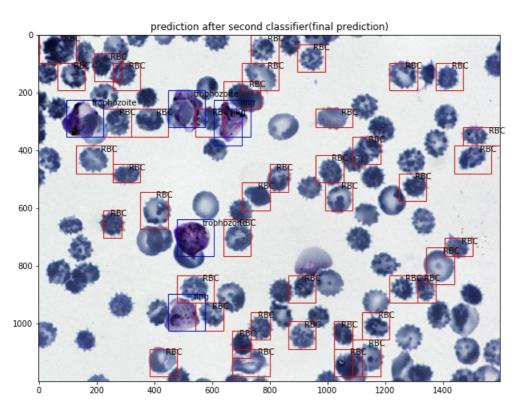
#### In [17]:

predicted("11b72c7b-5c52-44b7-adb7-21eacc25e441.png")

/usr/local/lib/python3.6/dist-packages/ipykernel\_launcher.py:91: UserWarning: Boolean Series key w ill be reindexed to match DataFrame index.







# Conclusion

- Our First level Classifier is doing decent job in classifying the infected and uninfected cells but not so well in classifying uninfected cells(RBC and Lueckocyte)
- Our second level classifier also working nice but i have taken very less resolution of cropped images due to memory issues. If we take good resolution we can get still more good results.
- our model is slightly confusing with "difficult" class, since actually it is not an infected cell type, it is just given because we could not able to classify few cells into any of 6 categories. So our model will predict correct class instead of difficult.

# **Future Work**

• Instead of taking two different classifiers, we could balance the dataset by creating duplicate images of minor class for example we can use GAN(Generative Adversarial Network) as a data augmentation technique.