# Glaucoma Classification

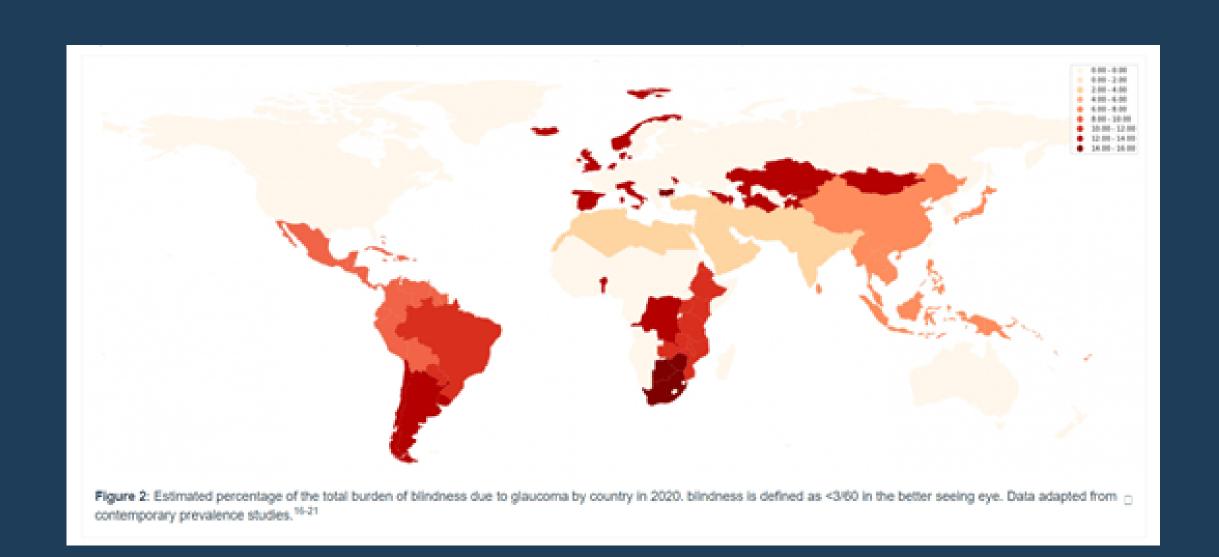
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# Member

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## Problem Statement



ต้อหินเป็นโรคที่พบได้บ่อยในผู้สูงอายุและถือเป็นโรคที่ทำให้เกิดภาวะสูญเสียการมอง เห็นชนิดถาวรที่พบได้บ่อยที่สุดทั่วโลกเป็นอันดับ 2 รองจากโรคต้อกระจก โดยในปี ค.ศ. 2020 มีผู้เป็นโรคต้อหินถึง 76 ล้านคน

## **OBJECTIVES**



เพื่อทำการสร้าง AI ที่ใช้ในการตรวจคัดกรองโรค glaucoma, ตาปกติ และ โรคอื่นๆด้วย Machine learning และ deep learning

# Machine Learning

## Data Collection

ประเภท	จำนวนข้อมูล
Glaucoma	1,363
Normal	2,009
Others	896
Total	4,268

## EXPERIMENTAL SETUP

- 1) **คัดรูป และ ทำการสกัด** Features
- 2) **ทำการเก็บ** Dataset **จาก** Features **ที่สกัดได้**
- 3) **นำ** Dataset **ไปทำการ** Train Model **ด้วย** Quadratic Discriminant Analysis Algorithm และ RBF SVM Algorithm
- 4) **ทดสอบ** Model **ที่ได้ ด้วย** Test Data

## EXPERIMENTAL SETUP

5-Fold Cross Validation

	Data Train					Test Data
Model 1	Fold 1	Fold 2	Fold 3	Fold 4	Fold 5	Test Data
Model 2	Fold 1	Fold 2	Fold 3	Fold 4	Fold 5	Test Data
Model 3	Fold 1	Fold 2	Fold 3	Fold 4	Fold 5	Test Data
Model 4	Fold 1	Fold 2	Fold 3	Fold 4	Fold 5	Test Data
Model 5	Fold 1	Fold 2	Fold 3	Fold 4	Fold 5	Test Data

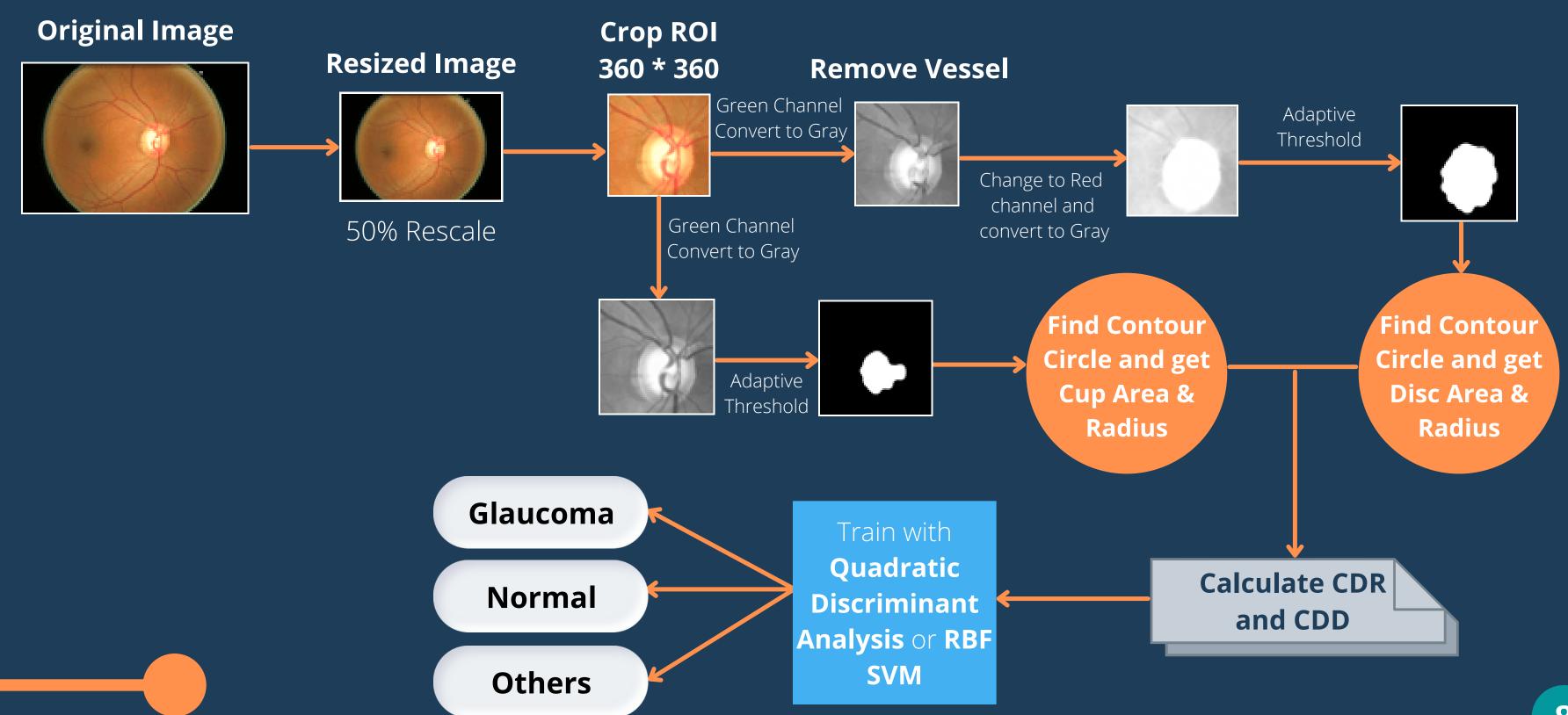
Data Validation





Data Test

## **Machine Learning Methods**



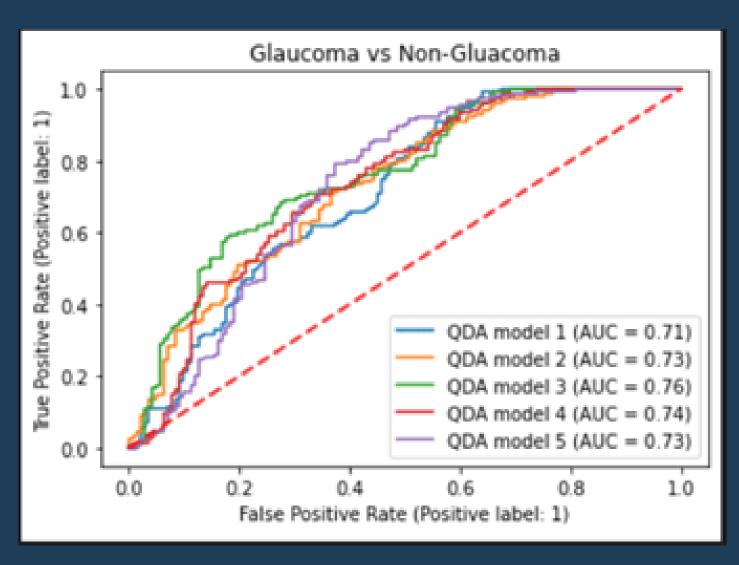
## ARCHITECTURE

Quadratic
Discriminant
Analysis



#### Glaucoma vs Non- Glaucoma

Quadratic Discriminant Analysis (Model 3)

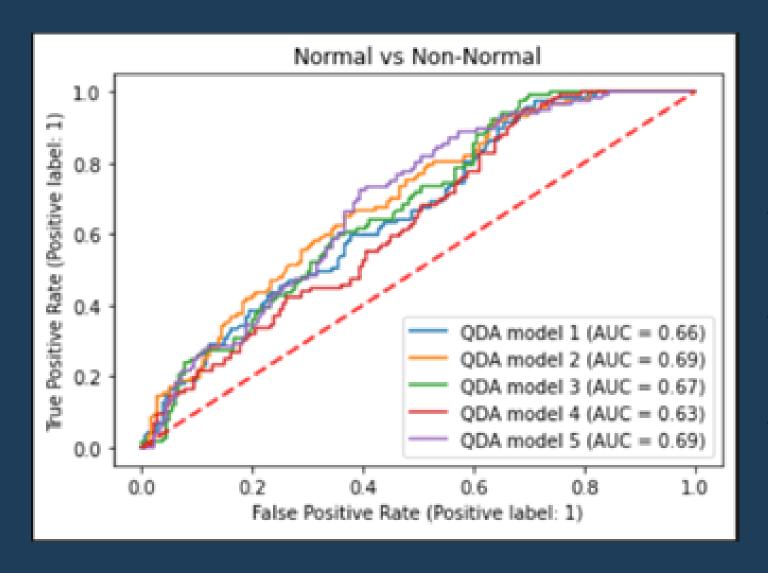


ประเภทจริง	ประเภ	SDU	
(True class)	บวก(Positive)	ลบ(Negative)	(Total)
บวก(Positive)	149(80.11%)	37(19.89%)	186
aບ(Negative)	78(42.16%)	107(57.84%)	185
sɔu(Total)	227	144	371

Accuracy	Specificity	Sensitivity	Precision	F1 Score
0.690027	0.578378	0.801075	0.656388	0.690027

Quadratic Discriminant Analysis (Model 2)

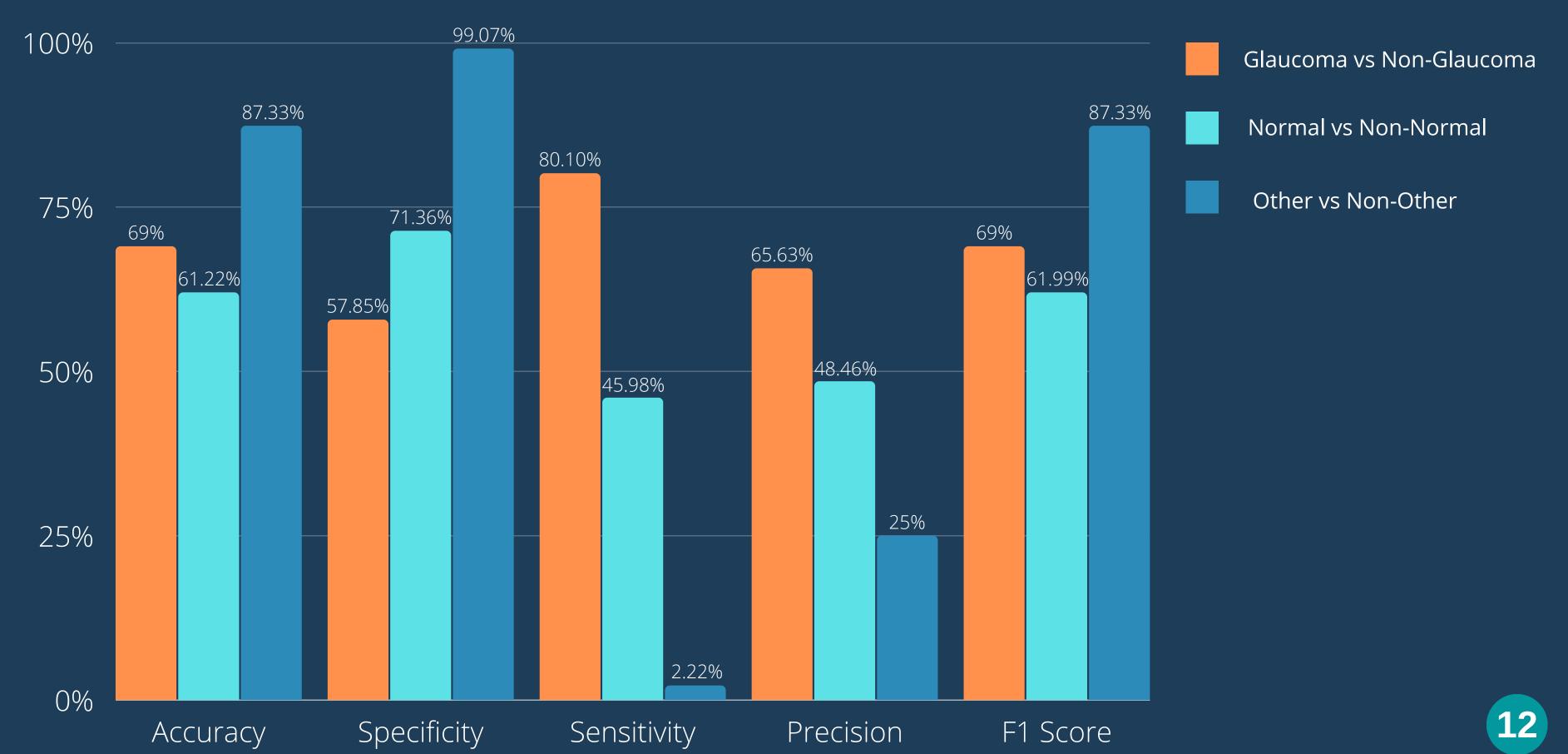
#### Normal vs Non- Normal



ประเภทจริง	ประเภ	SOU	
(True class)	บวก(Positive)	au(Negative)	(Total)
บวก(Positive)	63(45.98%)	74(54.02%)	137
<b>au</b> (Negative)	67(28.63%)	167(71.37%)	234
<b>sວມ</b> (Total)	130	241	371

Accuracy	Specificity	Sensitivity	Precision	F1 Score
0.619946	0.713675	0.459854	0.484615	0.619946

## Quadratic Discriminant Analysis



RBF SVM (Model 3)

# Glaucoma vs Non-Gluacoma 1.0 0.8 0.6 RBF SVM model 1 (AUC = 0.75) RBF SVM model 2 (AUC = 0.74) RBF SVM model 3 (AUC = 0.79) RBF SVM model 4 (AUC = 0.78) RBF SVM model 5 (AUC = 0.76) RBF SVM model 5 (AUC = 0.76) RBF SVM model 5 (AUC = 0.76) RBF SVM model 5 (AUC = 0.76)

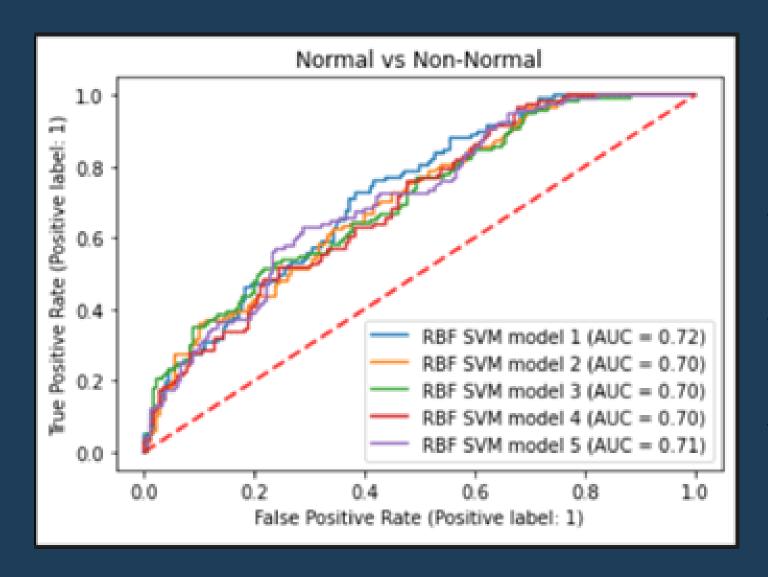
#### Glaucoma vs Non- Glaucoma

ประเภทจริง	ประเภ	SDU	
(True class)	บวก(Positive)	<b>aບ</b> (Negative)	(Total)
<b>บวก</b> (Positive)	161(86.56%)	25(13.44%)	186
<b>aບ</b> (Negative)	80(43.24%)	105(56.76%)	185
<b>sɔu</b> (Total)	241	130	371

Accuracy	Specificity	Sensitivity	Precision	F1 Score
0.716981	0.567567	0.865591	0.668050	0.716981

RBF SVM (Model 1)

#### Normal vs Non- Normal

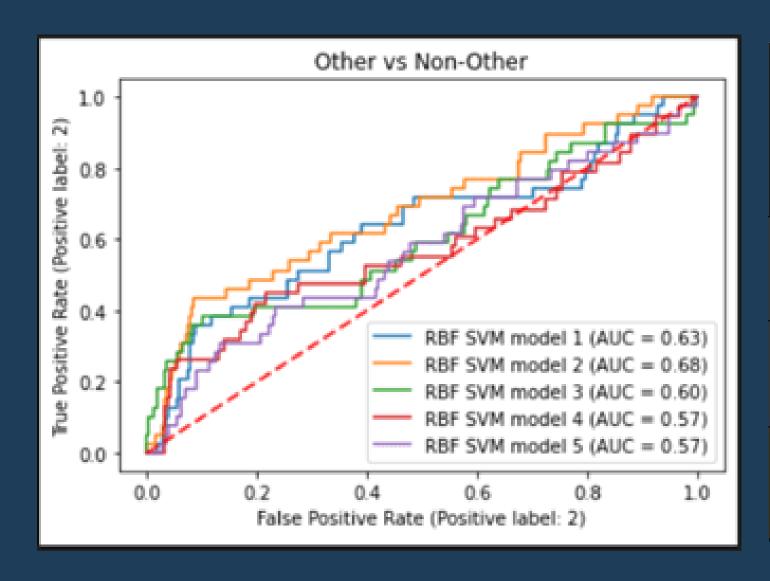


ประเภทจริง	ประเภ	SOU	
(True class)	บวก(Positive)	au(Negative)	(Total)
บวก(Positive)	50(36.50%)	87(63.50%)	137
<b>au</b> (Negative)	41(17.52%)	193(82.48%)	234
<b>sວມ</b> (Total)	91	280	371

Accuracy	Specificity	Sensitivity	Precision	F1 Score
0.649596	0.829060	0.343066	0.540230	0.617251

RBF SVM (Model 5)

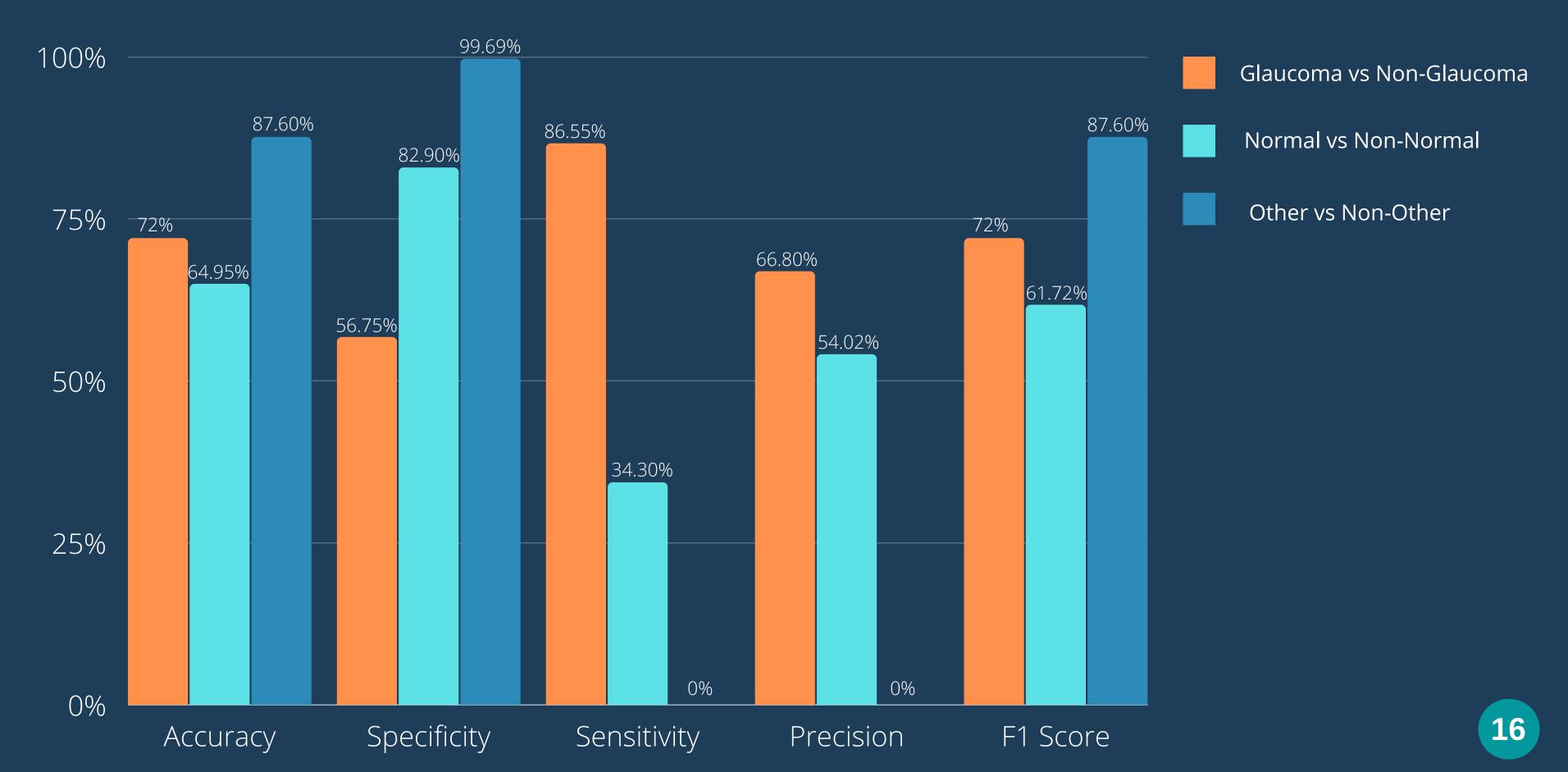
#### Other vs Non- Other



ประเภทจริง	ประเภ	SOU	
(True class)	บวก(Positive)	ลบ(Negative)	(Total)
บวก(Positive)	0(0%)	45(100%)	45
<b>aບ</b> (Negative)	1(0.30%)	325(99.70%)	326
sɔu(Total)	1	370	371

Accuracy	Specificity	Sensitivity	Precision	F1 Score
0.876010	0.996932	0	0	0.876010

## RBF SVM



```
def delVessel(image):
    blue,green,red = cv.split(image)
    kernel = cv.getStructuringElement(cv.MORPH_ELLIPSE,(26,26))
    ves = cv.morphologyEx(green, cv.MORPH_BLACKHAT, kernel)
    vessel2 = cv.bitwise_or(ves,green)
    vessel = cv.bitwise_or(ves,red)
    vessel = cv.medianBlur(red,7)
    plt.imshow(vessel2)
    return vessel
```

```
def GetDisc(image):
   M = 60 #filter size
   filter = signal.gaussian(M, std=7) #Gaussian Window
   filter=filter/sum(filter)
   STDf = filter.std() #It'standard deviation
   image_pre = image-image.mean()-image.std()
   thr = (0.5 * M) - (2*STDf) - image_pre.std()
   r,c = image.shape
   Dd = np.zeros(shape=(r,c))
   for i in range(1,r):
       for j in range(1,c):
           if image_pre[i,j]>thr:
               Dd[i,j]=255
           else:
               Dd[i,j]=0
   Dd = cv.morphologyEx(Dd, cv.MORPH_CLOSE, cv.getStructuringElement(cv.MORPH_ELLIPSE,(2,2)), iterations = 1)
   Dd = cv.morphologyEx(Dd, cv.MORPH_OPEN, cv.getStructuringElement(cv.MORPH_ELLIPSE,(7,7)), iterations = 1)
   Dd = cv.morphologyEx(Dd, cv.MORPH_CLOSE, cv.getStructuringElement(cv.MORPH_ELLIPSE,(1,21)), iterations = 1)
   Dd = cv.morphologyEx(Dd, cv.MORPH_OPEN, cv.getStructuringElement(cv.MORPH_ELLIPSE,(21,1)), iterations = 1)
   Dd = cv.morphologyEx(Dd, cv.MORPH_CLOSE, cv.getStructuringElement(cv.MORPH_ELLIPSE,(23,23)), iterations = 1)
   Dd = cv.morphologyEx(Dd, cv.MORPH_OPEN, cv.getStructuringElement(cv.MORPH_ELLIPSE,(43,43)), iterations = 1)
   Dd = np.uint8(Dd)
   return Dd
```

```
def GetCup(image):
    blue, green, red = cv.split(image)
    green = cv.medianBlur(green,7)
             #filter size
    M = 60
    filter = signal.gaussian(M, std=7) #Gaussian Window
    filter = filter/sum(filter)
    STDf = filter.std() #It'standard deviation
    green_pre = green-green.mean()-green.std()
    thr = (0.5 * M) + (2 * STDf) + (green_pre.std()) + (green_pre.mean())
    r,c = green.shape
    Dc = np.zeros(green.shape[:2])
    for i in range(1,r):
        for j in range(1,c):
            if green_pre[i,j]>thr:
               Dc[i,j]=255
            else:
               Dc[i,j]=0
    Dc = cv.morphologyEx(Dc, cv.MORPH_CLOSE, cv.getStructuringElement(cv.MORPH_ELLIPSE,(2,2)), iterations = 1)
    Dc = cv.morphologyEx(Dc, cv.MORPH_OPEN, cv.getStructuringElement(cv.MORPH_ELLIPSE,(7,7)), iterations = 1)
    Dc = cv.morphologyEx(Dc, cv.MORPH_CLOSE, cv.getStructuringElement(cv.MORPH_ELLIPSE,(1,21)), iterations = 1)
    Dc = cv.morphologyEx(Dc, cv.MORPH_OPEN, cv.getStructuringElement(cv.MORPH_ELLIPSE,(21,1)), iterations = 1)
    Dc = cv.morphologyEx(Dc, cv.MORPH_CLOSE, cv.getStructuringElement(cv.MORPH_ELLIPSE,(33,33)), iterations = 1)
    Dc = cv.morphologyEx(Dc, cv.MORPH_OPEN, cv.getStructuringElement(cv.MORPH_ELLIPSE,(33,33)), iterations = 1)
    Dc = np.uint8(Dc)
    return Dc
```

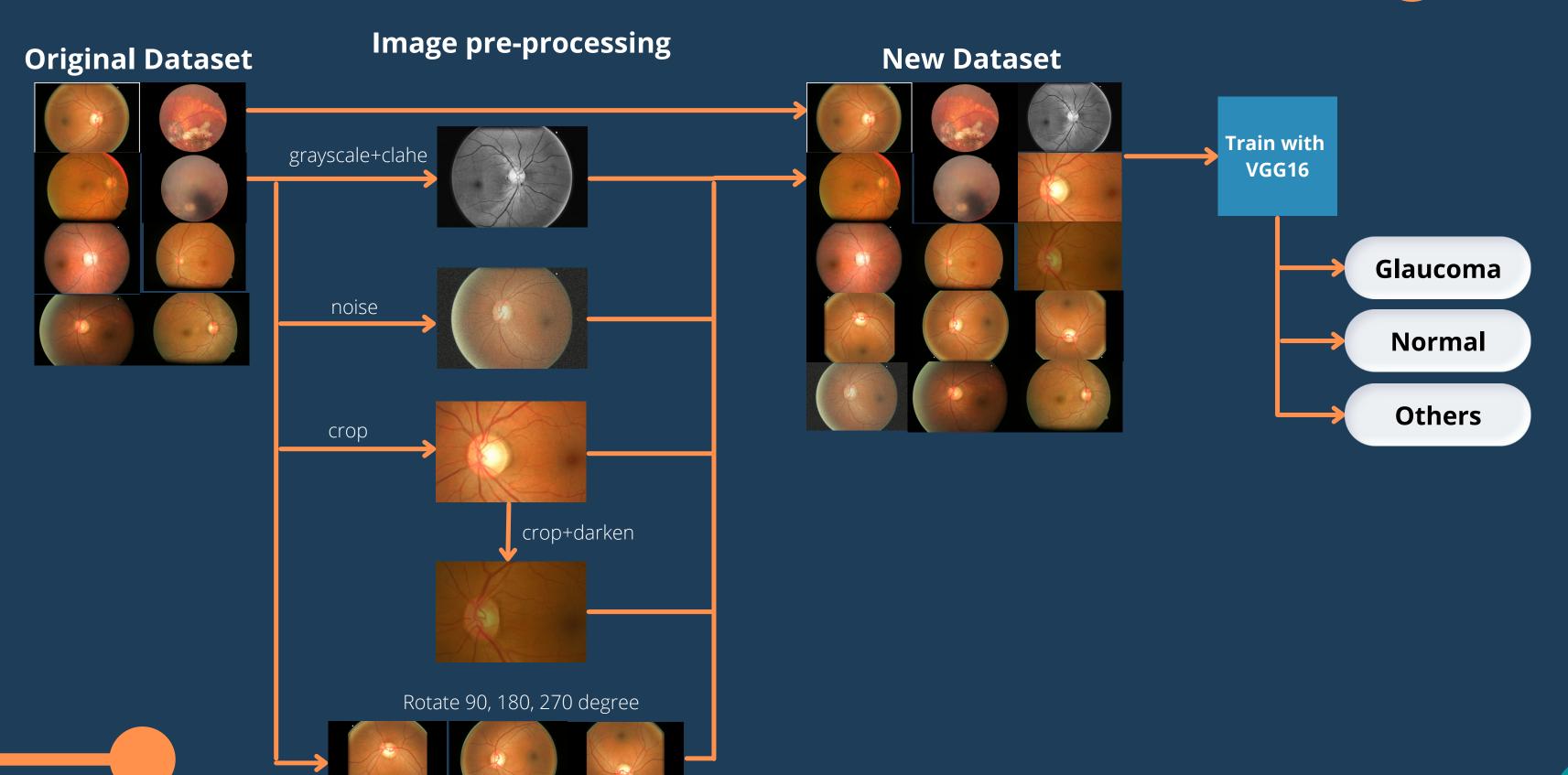
```
trainData01,testData01,trainType01,testType01 = train_test_split(data01, eye01, test_size=0.2, random_state=1)
skf = StratifiedKFold(n_splits=5)
plt.scatter(data01[:,0],data01[:,1], c=eye01)
plt.xlabel('CDR')
plt.ylabel('y')
# QDA model for Glucoma vs Non-Glaucoma
clf = QuadraticDiscriminantAnalysis()
tprs = []
aucs = []
mean_fpr = np.linspace(0, 1, 100)
fig, ax = plt.subplots()
i = 1
for train_index, test_index in skf.split(trainData01, trainType01):
   x_train, x_test = trainData01[train_index], trainData01[test_index]
   y_train, y_test = trainType01[train_index], trainType01[test_index]
   clf.fit(x_train,y_train)
   filename = 'savemodel/GOQDA_model'+str(i)+'.sav'
   joblib.dump(clf,filename)
   viz = plot_roc_curve(clf,x_test, y_test,name='QDA model {}'.format(i), ax=ax)
   interp_tpr = np.interp(mean_fpr, viz.fpr, viz.tpr)
   interp_tpr[0] = 0.0
   tprs.append(interp_tpr)
   aucs.append(viz.roc_auc)
   i+=1
```

# Deep Learning

## Data Collection

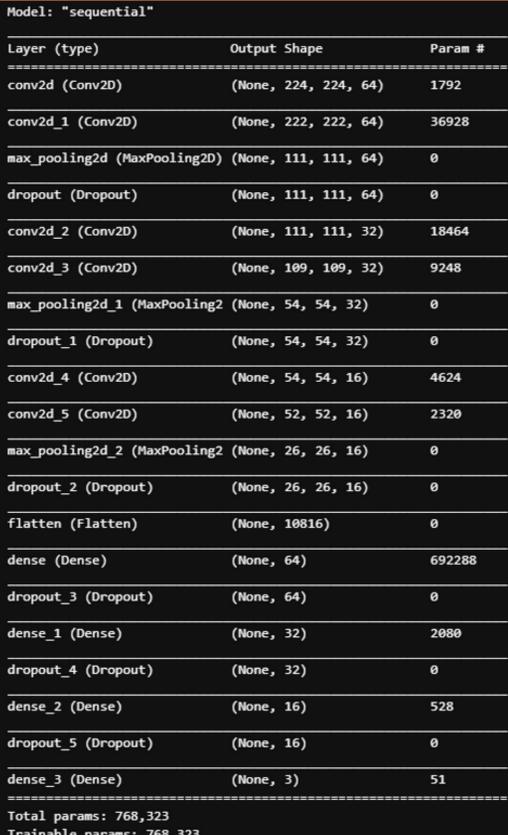
ประเภท	จำนวนข้อมูล	
Glaucoma	12,516	
Normal	9,050	
Others	5,049	
Total	26,615	

## **Deep Learning Methods**



## ARCHITECTURE

#### Model 1



Trainable params: 768,323 Non-trainable params: 0



Trainable params: 1,539

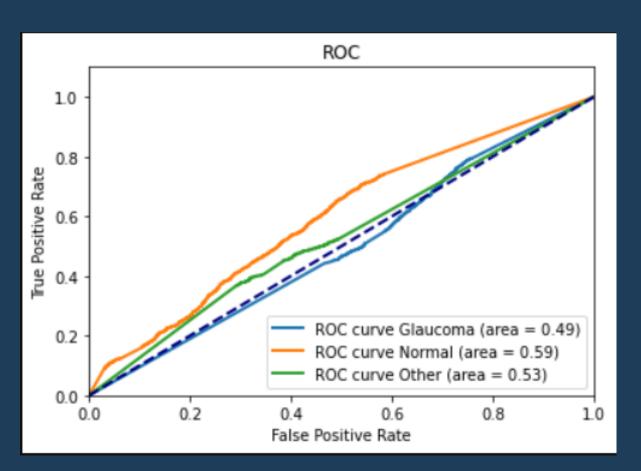
Non-trainable params: 14,714,688



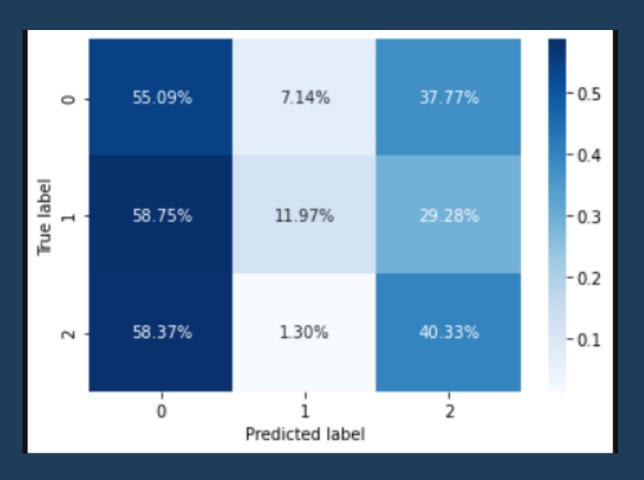
Model: "sequential"			
Layer (type)	Output	Shape	Param #
vgg16 (Functional)	(None,	7, 7, 512)	14714688
global_average_pooling2d (Gl	(None,	512)	0
dense (Dense)	(None,	3)	1539
Total params: 14,716,227			

## Model 1

#### ROC



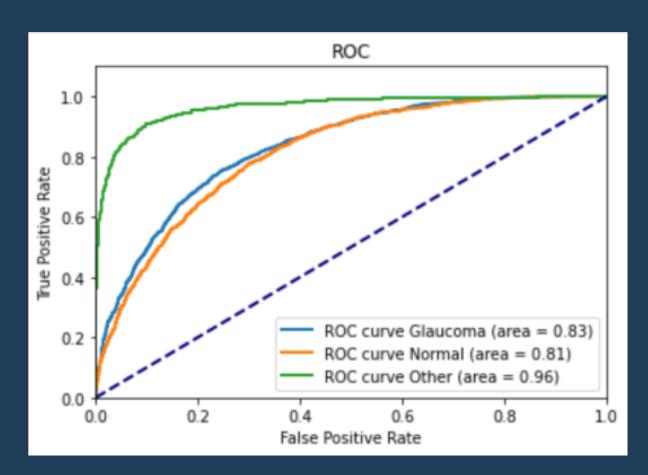
#### confusion table



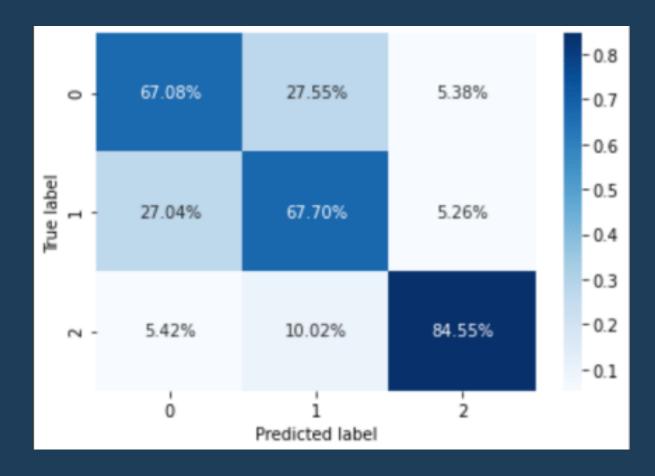
Type	Precision	Recall	F1 Score
Glaucoma	0.45483	0.55090	0.49828
Normal	0.53061	0.11974	0.19538
Other	0.21632	0.40330	0.28160

#### Model 2

#### ROC



#### confusion table



Type	Precision	Recall	F1 Score
Glaucoma	0.75522	0.67079	0.71051
Normal	0.60780	0.67697	0.64052
Other	0.78791	0.84552	0.81570

```
import tensorflow as tf
import numpy as np
from tensorflow.keras.preprocessing import image
from sklearn.metrics import accuracy_score, f1_score, precision_score, confusion_matrix, roc_curve, auc, plot_roc_curve
path_train = "dataset/train"
img_train = tf.keras.preprocessing.image_dataset_from_directory(
   path_train,
    validation_split=0.2,
    subset = "training",
    seed = 125,
   image_size = (224,224),
    batch_size = 32
img_validation = tf.keras.preprocessing.image_dataset_from_directory(
    path_train,
    validation_split=0.2,
   subset = "validation",
    seed = 125,
    image_size = (224,224),
    batch_size = 32
```

```
VGG16_MODEL=tf.keras.applications.VGG16(input_shape=(224,224,3),
                                               include top=False,
                                               weights='imagenet')
VGG16_MODEL.trainable=False
global average layer = tf.keras.layers.GlobalAveragePooling2D()
prediction_layer = tf.keras.layers.Dense(3,activation='softmax')
model = tf.keras.Sequential([
  VGG16 MODEL,
  global_average_layer,
  prediction_layer
model.compile(optimizer="adam",
              loss=tf.keras.losses.sparse_categorical_crossentropy,
              metrics=["accuracy"])
model.summary()
```

```
history = model.fit(img_train,
                    epochs=18,
                    steps_per_epoch=30,
                    validation_steps=2,
                    validation_data=img_validation)
model.save("deeplearningVGG16.h5")
#import Test img
path_test = "dataset/test"
img_test = tf.keras.preprocessing.image_dataset_from_directory(
    path_test,
   image_size = (224,224)
label = ["glaucoma", "normal", "other"]
loadmodel = tf.keras.models.load_model("deeplearningVGG16.h5")
predict = loadmodel.predict(img_test)
prediction = np.argmax(predict)
```

```
i = 0
glau = 0
norm = 0
oth = 0
for i in predict:
    index = np.argmax(i)
    if index == 0:
        glau+=1
    elif index == 1:
        norm+=1
    elif index==2:
        oth+=1

print(glau)
print(norm)
print(oth)
```

```
# change file type to jpeg

import imghdr
import cv2
import os
import glob

for file in glob.glob('dataset/test/other/*.jpg'):
    image = cv2.imread(file)
    file_type = imghdr.what(file)
    if file_type != 'jpeg':
        print(file + " - invalid - " + str(file_type))
        cv2.imwrite(file, image)
    print("finish jpg",image)
```

## **PROBLEM**

GPU Error.



Some images can't crop.



The feature can't extract other images.





## Reference

Machine Learning

[1] A. Issac, M. Parthasarthi and M. K. Dutta, "An adaptive threshold based algorithm for optic disc and cup segmentation in fundus images," 2015 2nd International Conference on Signal Processing and Integrated Networks (SPIN), Noida, India, 2015, pp. 143-147, doi: 10.1109/SPIN.2015.7095384. (access Mar. 23th 2021)

[2] Ahmed Almazroa, Ritambhar Burman, Kaamran Raahemifar, Vasudevan Lakshminarayanan, "Optic Disc and Optic Cup Segmentation Methodologies for Glaucoma Image Detection: A Survey", Journal of Ophthalmology, vol. 2015, Article ID 180972, 28 pages, 2015. https://doi.org/10.1155/2015/180972 (access Mar. 23th 2021)

[3] Atheesan S. and Yashothara S., "Automatic glaucoma detection by using funduscopic images," 2016 International Conference on Wireless Communications, Signal Processing and Networking (WiSPNET), Chennai, India, 2016, pp. 813-817, doi: 10.1109/WiSPNET.2016.7566246.

(access Mar. 24th 2021)

[4] Ashish Issac, M. Parthasarthi, Malay Kishore Dutta, Flowchart for the segmentation of optic disc and cup, [Online]. Available: https://ieeexplore.ieee.org/document/7095384. (access Mar. 24th 2021)



## Reference

Deep Learning

[1] Lorenzo Baraldi, "VGG-16 pre-trained model for Keras", 2015. [Online]. Available: https://gist.github.com/baraldilorenzo/07d7802847aaad0a35d3 (access May. 7th 2021)

[2] Siladittya Manna, "K-Fold Cross Validation for Deep Learning Models using Keras", Mar. 2020. [Online]. Available: https://medium.com/the-owl/k-fold-cross-validation-in-keras-3ec4a3a00538 (access May. 7th 2021)

[3] Machinecurve, "How to predict new samples with your TensorFlow / Keras model?", Feb. 2020. [Online]. Available: https://www.machinecurve.com/index.php/2020/02/21/how-to-predict-new-samples-with-your-keras-model (access May. 7th 2021)

[4] Patrick Kalkman, "Increase the Accuracy of Your CNN by Following These 5 Tips I Learned From the Kaggle Community", Feb. 2021. [Online]. Available: https://towardsdatascience.com/increase-the-accuracy-of-your-cnn-by-following-these-5-tips-i-learned-from-the-kaggle-community-27227ad39554 (access May. 8th 2021)



## Reference

Deep Learning

[5] Brijesh Thumar, "How to use VGG model in TensorFlow Keras", May. 2019. [Online]. Available: https://androidkt.com/how-to-use-vgg-model-in-tensorflow-keras/ (access May. 9th 2021)

[6] JIACHENG DANG, "Detecting Open-angle Glaucoma Using a Two-parts Deep Learning Architecture", Mar. 2020. [Online]. Available: https://ysjournal.com/detecting-open-angle-glaucoma-using-a-two-parts-deep-learning-architecture/ (access May. 9th 2021)

[7]Ahn JM, Kim S, Ahn KS, Cho SH, Lee KB, et al. (2019) Correction: A deep learning model for the detection of both advanced and early glaucoma using fundus photography. PLOS ONE 14(1): e0211579. https://doi.org/10.1371/journal.pone.0211579 (access May. 9th 2021)