```
import tensorflow as tf
import tensorflow hub as hub
import matplotlib.pyplot as plt
import numpy as np
import pandas as pd
import seaborn as sns
from tensorflow.keras.utils import get file
from sklearn.metrics import roc curve, auc, confusion matrix
from imblearn.metrics import sensitivity_score, specificity_score
import os
import glob
import zipfile
import random
# to get consistent results after multiple runs
tf.random.set_seed(7)
np.random.seed(7)
random.seed(7)
# 0 for benign, 1 for malignant
class_names = ["benign", "malignant"]
```

Preparing the Dataset

```
def download_and_extract_dataset():
    # dataset from https://github.com/udacity/dermatologist-ai
    # 5.3GB
    train_url = "https://s3-us-west-1.amazonaws.com/udacity-dlnfd/datasets/skin-cancer/train.zip"
    # 824.5MB
    valid_url = "https://s3-us-west-1.amazonaws.com/udacity-dlnfd/datasets/skin-cancer/valid.zip"
```

```
# 5.1GB

test_url = "https://s3-us-west-1.amazonaws.com/udacity-
dlnfd/datasets/skin-cancer/test.zip"

for i, download_link in enumerate([valid_url, train_url, test_url]):
    temp_file = f"temp{i}.zip"
    data_dir = get_file(origin=download_link,
fname=os.path.join(os.getcwd(), temp_file))
    print("Extracting", download_link)
    with zipfile.ZipFile(data_dir, "r") as z:
        z.extractall("data")
    # remove the temp file
    os.remove(temp_file)

# comment the below line if you already downloaded the dataset
download_and_extract_dataset()
```

```
# preparing data
# generate CSV metadata file to read img paths and labels from it
def generate csv(folder, label2int):
    folder_name = os.path.basename(folder)
    labels = list(label2int)
    # generate CSV file
    df = pd.DataFrame(columns=["filepath", "label"])
    i = 0
    for label in labels:
        print("Reading", os.path.join(folder, label, "*"))
        for filepath in glob.glob(os.path.join(folder, label, "*")):
            df.loc[i] = [filepath, label2int[label]]
            i += 1
    output_file = f"{folder_name}.csv"
    print("Saving", output_file)
    df.to_csv(output_file)
# generate CSV files for all data portions, labeling nevus and
seborrheic keratosis
```

```
# as 0 (benign), and melanoma as 1 (malignant)
# you should replace "data" path to your extracted dataset path
# don't replace if you used download_and_extract_dataset() function
generate_csv("data/train", {"nevus": 0, "seborrheic_keratosis": 0,
"melanoma": 1})
generate_csv("data/valid", {"nevus": 0, "seborrheic_keratosis": 0,
"melanoma": 1})
generate_csv("data/test", {"nevus": 0, "seborrheic_keratosis": 0,
"melanoma": 1})
```

```
# loading data
train_metadata_filename = "train.csv"
valid_metadata_filename = "valid.csv"

# load CSV files as DataFrames

df_train = pd.read_csv(train_metadata_filename)

df_valid = pd.read_csv(valid_metadata_filename)

n_training_samples = len(df_train)

n_validation_samples = len(df_valid)

print("Number of training samples:", n_training_samples)

print("Number of validation samples:", n_validation_samples)

train_ds = tf.data.Dataset.from_tensor_slices((df_train["filepath"],
    df_train["label"]))

valid_ds = tf.data.Dataset.from_tensor_slices((df_valid["filepath"],
    df_valid["label"]))
```

```
Number of training samples: 2000
Number of validation samples: 150
```

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Let's load the images:

```
# preprocess data
def decode_img(img):
    # convert the compressed string to a 3D uint8 tensor
    img = tf.image.decode_jpeg(img, channels=3)
    # Use `convert_image_dtype` to convert to floats in the [0,1] range.
    img = tf.image.convert_image_dtype(img, tf.float32)
    # resize the image to the desired size.
    return tf.image.resize(img, [299, 299])
```

```
def process_path(filepath, label):
    # load the raw data from the file as a string
    img = tf.io.read_file(filepath)
    img = decode_img(img)
    return img, label

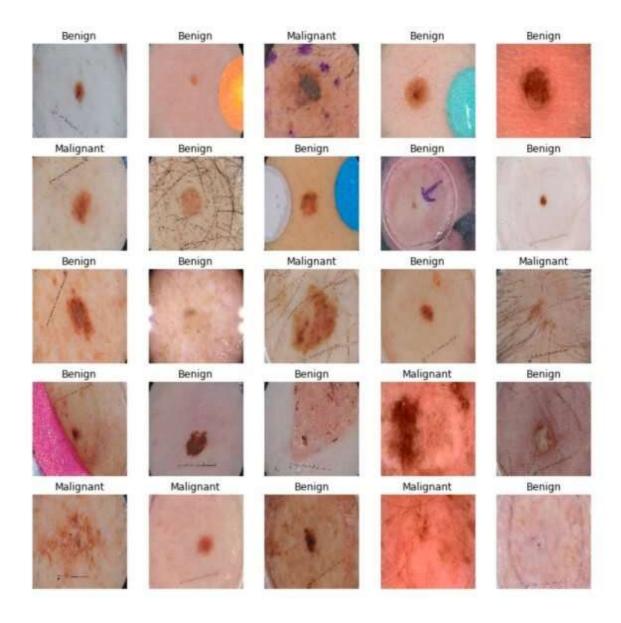
valid_ds = valid_ds.map(process_path)
train_ds = train_ds.map(process_path)
# test_ds = test_ds
for image, label in train_ds.take(1):
    print("Image shape:", image.shape)
    print("Label:", label.numpy())
Image shape: (299, 299, 3)
Label: 0
```

```
# training parameters
batch_size = 64
optimizer = "rmsprop"

def prepare_for_training(ds, cache=True, batch_size=64, shuffle_buffer_size=1000):
    if cache:
        if isinstance(cache, str):
            ds = ds.cache(cache)
        else:
            ds = ds.cache()
        # shuffle the dataset
        ds = ds.shuffle(buffer_size=shuffle_buffer_size)
        # Repeat forever
        ds = ds.repeat()
```

```
# split to batches
  ds = ds.batch(batch_size)
  # `prefetch` lets the dataset fetch batches in the background while
the model
 # is training.
 ds = ds.prefetch(buffer_size=tf.data.experimental.AUTOTUNE)
  return ds
valid ds = prepare for training(valid ds, batch size=batch size,
cache="valid-cached-data")
train_ds = prepare_for_training(train_ds, batch_size=batch_size,
cache="train-cached-data")
batch = next(iter(valid_ds))
def show_batch(batch):
  plt.figure(figsize=(12,12))
 for n in range(25):
      ax = plt.subplot(5,5,n+1)
      plt.imshow(batch[0][n])
      plt.title(class_names[batch[1][n].numpy()].title())
      plt.axis('off')
show_batch(batch)
```

Output:



```
# building the model
# InceptionV3 model & pre-trained weights
module_url = "https://tfhub.dev/google/tf2-
preview/inception_v3/feature_vector/4"
m = tf.keras.Sequential([
    hub.KerasLayer(module_url, output_shape=[2048], trainable=False),
    tf.keras.layers.Dense(1, activation="sigmoid")
])
m.build([None, 299, 299, 3])
m.compile(loss="binary_crossentropy", optimizer=optimizer,
metrics=["accuracy"])
m.summary()
```

```
Model: "sequential"

Layer (type) Output Shape Param #

keras_layer (KerasLayer) multiple 21802784

dense (Dense) multiple 2049

Total params: 21,804,833

Trainable params: 2,049

Non-trainable params: 21,802,784
```

Training the Model

We now have our dataset and the model, let's get them together:

Here is a part of the output during training:

```
Epoch 00001: val_loss improved from inf to 0.49703, saving model to
benign-vs-malignant 64 rmsprop 0.497.h5
accuracy: 0.7722 - val loss: 0.4970 - val accuracy: 0.8125
<...SNIPED...>
Epoch 27/100
accuracy: 0.8708
Epoch 00027: val loss improved from 0.40253 to 0.38991, saving model to
benign-vs-malignant 64 rmsprop 0.390.h5
31/31 [============ ] - 21s 691ms/step - loss: 0.3025
- accuracy: 0.8684 - val_loss: 0.3899 - val_accuracy: 0.8359
<...SNIPED...>
Epoch 41/100
accuracy: 0.8802
Epoch 00041: val_loss did not improve from 0.38991
31/31 [============= ] - 21s 690ms/step - loss: 0.2829
- accuracy: 0.8790 - val_loss: 0.3948 - val_accuracy: 0.8281
Epoch 42/100
accuracy: 0.8859
Epoch 00042: val_loss did not improve from 0.38991
- accuracy: 0.8831 - val loss: 0.4572 - val accuracy: 0.8047
```

Model Evaluation

First, let's load our test set, just like previously:

```
# evaluation
# load testing set

test_metadata_filename = "test.csv"

df_test = pd.read_csv(test_metadata_filename)
n_testing_samples = len(df_test)

print("Number of testing samples:", n_testing_samples)

test_ds = tf.data.Dataset.from_tensor_slices((df_test["filepath"], df_test["label"]))

def prepare_for_testing(ds, cache=True, shuffle_buffer_size=1000):
```

```
if cache:
    if isinstance(cache, str):
        ds = ds.cache(cache)
    else:
        ds = ds.cache()

    ds = ds.shuffle(buffer_size=shuffle_buffer_size)
    return ds

test_ds = test_ds.map(process_path)

test_ds = prepare_for_testing(test_ds, cache="test-cached-data")
```

The above code loads our test data and prepares it for testing:

```
Number of testing samples: 600
```

images of the shape (299, 299, 3) can fit our memory, let's convert our test set from tf.data into a NumPy array:

```
# convert testing set to numpy array to fit in memory (don't do that
when testing
# set is too large)
y_test = np.zeros((n_testing_samples,))
X_test = np.zeros((n_testing_samples, 299, 299, 3))
for i, (img, label) in enumerate(test_ds.take(n_testing_samples)):
    # print(img.shape, label.shape)
    X_test[i] = img
    y_test[i] = label.numpy()

print("y_test.shape:", y_test.shape)
# load the weights with the least loss
m.load_weights("benign-vs-malignant_64_rmsprop_0.390.h5")
print("Evaluating the model...")
loss, accuracy = m.evaluate(X_test, y_test, verbose=0)
print("Loss:", loss, " Accuracy:", accuracy)
```

Copy

Output:

```
Evaluating the model...

Loss: 0.4476394319534302 Accuracy: 0.8
```

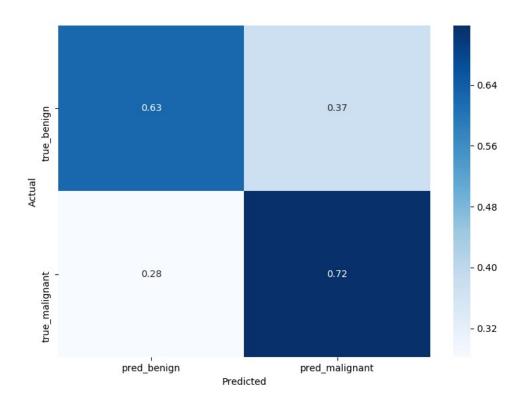
The below function does that:

```
def get predictions(threshold=None):
  Returns predictions for binary classification given `threshold`
  For instance, if threshold is 0.3, then it'll output 1 (malignant)
for that sample if
  the probability of 1 is 30% or more (instead of 50%)
 y_pred = m.predict(X_test)
  if not threshold:
    threshold = 0.5
  result = np.zeros((n_testing_samples,))
 for i in range(n_testing_samples):
   # test melanoma probability
    if y pred[i][0] >= threshold:
      result[i] = 1
   # else, it's 0 (benign)
  return result
threshold = 0.23
# get predictions with 23% threshold
# which means if the model is 23% sure or more that is malignant,
# it's assigned as malignant, otherwise it's benign
y pred = get predictions(threshold)
```

Now let's draw our confusion matrix and interpret it:

```
def plot_confusion_matrix(y_test, y_pred):
    cmn = confusion_matrix(y_test, y_pred)
    # Normalise
    cmn = cmn.astype('float') / cmn.sum(axis=1)[:, np.newaxis]
```

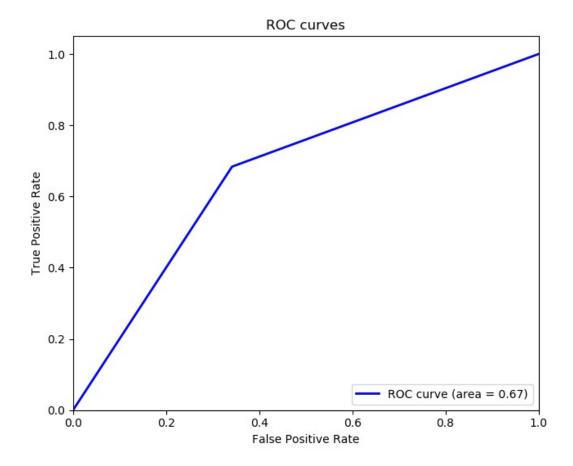
Output:



```
def plot_roc_auc(y_true, y_pred):
    This function plots the ROC curves and provides the scores.
    # prepare for figure
    plt.figure()
    fpr, tpr, _ = roc_curve(y_true, y_pred)
    # obtain ROC AUC
    roc_auc = auc(fpr, tpr)
    # print score
    print(f"ROC AUC: {roc_auc:.3f}")
    # plot ROC curve
    plt.plot(fpr, tpr, color="blue", lw=2,
                label='ROC curve (area = {f:.2f})'.format(d=1,
f=roc_auc))
    plt.xlim([0.0, 1.0])
    plt.ylim([0.0, 1.05])
    plt.xlabel('False Positive Rate')
    plt.ylabel('True Positive Rate')
    plt.title('ROC curves')
    plt.legend(loc="lower right")
    plt.show()
plot_roc_auc(y_test, y_pred)
```

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Output:



ROC AUC: 0.671