Splitting datasets and metrics examples

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Import libraries

Import libraries

```
In[14]:
```

```
%pylab inline
import time # used for timing the training
import os # operating system commands
import shutil # High level operations for files
import random # Random generators
import pandas as pd # Pandas data read, maybe not needed in this case??
import tensorflow as tf # Tensorflow
from tensorflow.keras import layers, models, optimizers # Neural network stuff
from tensorflow.keras.preprocessing.image import ImageDataGenerator # For image processing with keras
from tensorflow.keras.metrics import Accuracy, FalseNegatives, FalsePositives # Specific metrics
from sklearn.metrics import classification_report, confusion_matrix, roc_curve # For final results analysis
print('Tensorflow version = ', tf.__version__) # Check the version
```

Populating the interactive namespace from numpy and matplotlib Tensorflow version = 2.1.0

Read the original file names

```
In[7]:
    # Original training files can be found here.
    orig_dir = '/kaggle/input/chest-xray-pneumonia/chest_xray/chest_xray/train'

# List only all jpeg-files (images)
    normal_images = [x for x in os.listdir(os.path.join(orig_dir,'NORMAL')) if x.endswith(".jpeg")]
    pneumonia_images = [x for x in os.listdir(os.path.join(orig_dir,'PNEUMONIA')) if x.endswith(".jpeg")]

# Show the statistics
    N_NORMAL = len(normal_images)
    N_PNEUMONIA = len(pneumonia_images)
    TOTAL = N_NORMAL + N_PNEUMONIA

print('Original training images:')
    print(f'(N_NORMAL:5d) normal cases')
    print(f'(N_PNEUMONIA:5d) pneumonia cases')
    print(f'(TOTAL:5d) totally.')
```

```
Original training images:
1341 normal cases
3875 pneumonia cases
5216 totally.
```

Create own directories

```
In[8]:
    # Own libraries, will be deleted when restarted
    train_dir = './train'
    valid_dir = './validation'
    test_dir = './test'
    all_dirs = [train_dir, valid_dir, test_dir]
    try:
        for d in all_dirs:
            os.mkdir(d)
            os.mkdir(os.path.join(d,'NORMAL'))
            os.mkdir(os.path.join(d,'PNEUMONIA'))
    except:
        pass
    print('Training directory = ', train_dir)
    print(os.listdir(train_dir))
```

```
Training directory = ./train
['NORMAL', 'PNEUMONIA']
```

Suffle the data

```
# Shuffle the lists in random order
random.shuffle(normal_images)
random.shuffle(pneumonia_images)
```

Copy original images to dataset folders

```
In[10]:
        # Copy the original images into train, validation, and test directories
        # Normal test images
        for fname in normal_images[:500]:
            src = os.path.join(orig_dir, 'NORMAL', fname)
            dst = os.path.join(test_dir, 'NORMAL', fname)
            shutil.copyfile(src, dst)
        # Normal validation images
        for fname in normal_images[501:1001]:
            src = os.path.join(orig_dir, 'NORMAL', fname)
            dst = os.path.join(valid_dir, 'NORMAL', fname)
            shutil.copyfile(src, dst)
        # Normal training images
        for fname in normal_images[1002:]:
            src = os.path.join(orig_dir, 'NORMAL', fname)
            dst = os.path.join(train_dir, 'NORMAL', fname)
            shutil.copyfile(src, dst)
```

```
# Pneumonia test images
for fname in pneumonia_images[:500]:
    src = os.path.join(orig_dir, 'PNEUMONIA', fname)
    dst = os.path.join(test_dir, 'PNEUMONIA', fname)
    shutil.copyfile(src, dst)

# Pneumonia validation images
for fname in pneumonia_images[501:1001]:
    src = os.path.join(orig_dir, 'PNEUMONIA', fname)
    dst = os.path.join(valid_dir, 'PNEUMONIA', fname)
    shutil.copyfile(src, dst)

# Pneumonia training images
for fname in pneumonia_images[1002:]:
    src = os.path.join(orig_dir, 'PNEUMONIA', fname)
    dst = os.path.join(train_dir, 'PNEUMONIA', fname)
    shutil.copyfile(src, dst)
```

Data generators

```
In[11]:
        TS = (150, 150) \# Image size
        BS = 128 # Batch size
        print('Training:')
        tg = ImageDataGenerator(rescale=1./255)
        train_generator = tg.flow_from_directory(
            train_dir,
            target_size = TS,
            batch_size = BS,
            class_mode = 'binary')
        print('Validation:')
        devg = ImageDataGenerator(rescale=1./255)
        dev_generator = devg.flow_from_directory(
            valid_dir,
            target_size = TS,
            batch_size = BS,
            shuffle = False,
            class_mode = 'binary')
        print('Testing:')
        testg = ImageDataGenerator(rescale=1./255)
        test_generator = devg.flow_from_directory(
            test_dir,
            target_size = TS,
            batch_size = BS,
            class_mode = 'binary')
```

```
Training:
Found 3212 images belonging to 2 classes.
Validation:
Found 1000 images belonging to 2 classes.
Testing:
Found 1000 images belonging to 2 classes.
```

```
In[12]:
    # Check the indices
    train_generator.class_indices

Out[12]:
    {'NORMAL': 0, 'PNEUMONIA': 1}
```

Model

Your own CNN model comes here

Training and watching the elapsed time

```
E = 3 #Number of epochs
# Start the clock
t_start = time.time()
print('Training ...', end='')
h = model.fit_generator(
      train_generator,
      steps_per_epoch = None,
      verbose = 0.
      epochs = E,
      validation_data = dev_generator,
      validation_steps = None
# Check the time and calculate the elapsed time and time per epoch
t_end = time.time()
t_{elapsed} = t_{end} - t_{start}
t_{per_epoch} = t_{elapsed/E}
print('Done.')
print(f'Time elapsed = {t_elapsed:.0f} seconds.')
print(f'Time per epoch = {t_per_epoch:.2f} seconds.')
# Save the model
model.save('case_2_run_007.h5')
print('Model saved.')
```

```
Training ...Done.
Time elapsed = 131 seconds.
Time per epoch = 43.57 seconds.
Model saved.
```

Check the keys and use them

and false negatives is increasing whenever

you rerun the model code (!?)

```
# Extract the metrics and loss
hh = h.history
acc = hh['accuracy']
acc_v = hh['val_accuracy']
fn = hh['false_negatives_2']
fn_v = hh['val_false_negatives_2']
fp = hh['false_positives_2']
fp_v = hh['val_false_positives_2']
loss = hh['loss']
loss_v = hh['val_loss']
epochs = arange(len(loss)) + 1
```

How to avoid increasing indexing in metrics

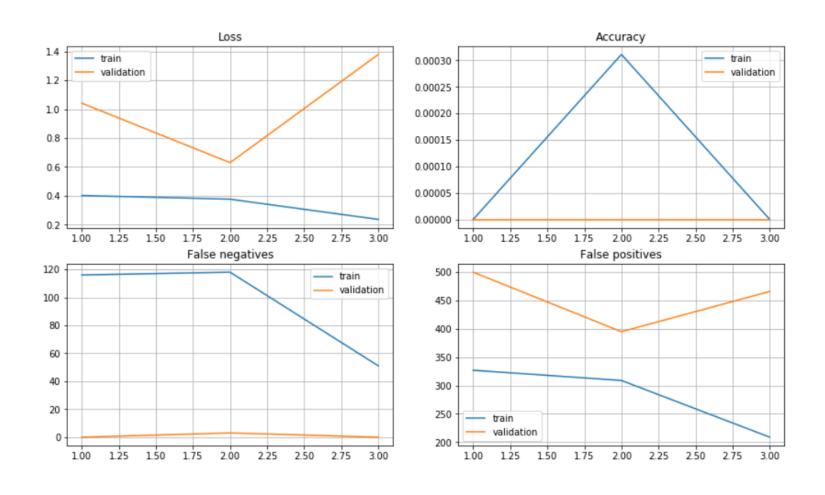
Create a list of metrics beforehand

```
ถ้าจะให้ดีใช้ "acc"
```

my_metrics = [Accuracy(), FalseNegatives(), FalsePositives()]

Then use the list in model compilation step:

Training results (bad)

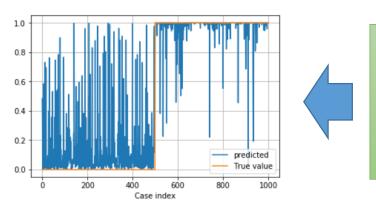


Finding the labels and predictions

```
devg = ImageDataGenerator(rescale=1./255)
dev_generator = devg.flow_from_directory(
    valid_dir,
    target_size = TS,
    batch_size = BS,
    shuffle = False,
    class_mode = 'binary')
```

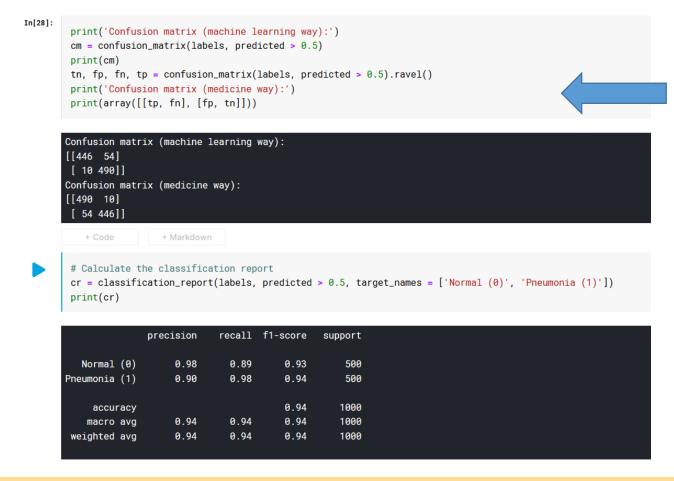
Note: You need to have shuffle = False for the validation generator!

Otherwise, the images are shuffled and you don't know which is the right label.



Aim is the predicted values are as close as possible for true values (=labels).

Confusion matrix and classification report



This is how you can convert the confusion matrix into format usually shown in medicine literature.

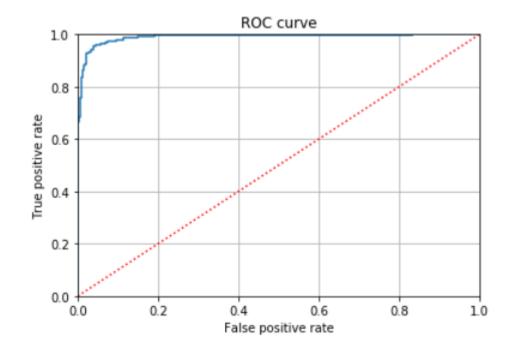
Remember: recall = *sensitivity,* recall for Pneumonia (1) = Sensitivity.

Q: How can you get the *specificity* from the classification report?

ROC curve analysis

```
# Calculate the ROC curve analysis
fpr, tpr, thresholds = roc_curve(labels, predicted, pos_label = 1)
```

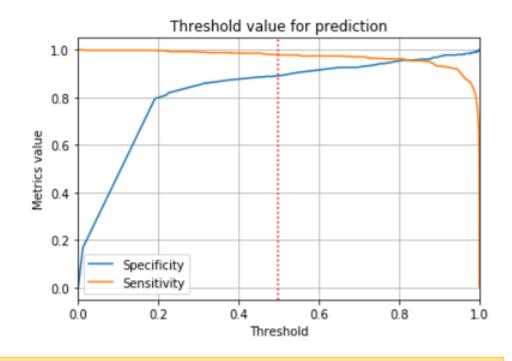
```
# Show the ROC curve
plot(fpr, tpr)
plot([0, 1], [0, 1], 'r:')
xlabel('False positive rate')
ylabel('True positive rate')
title('ROC curve')
xlim([0, 1])
ylim([0, 1])
grid()
```



Q: What does the ROC curve tell about the classifier?

Threshold analysis

```
# Find the best threshold
plot(thresholds, 1 - fpr, label = 'Specificity')
plot(thresholds, tpr, label = 'Sensitivity')
axvline(0.5, color = 'red', linestyle = ':')
xlim([0, 1])
title('Threshold value for prediction')
xlabel('Threshold')
ylabel('Metrics value')
legend()
grid()
```



Q: How would you select the theshold for the prediction?

Q: Is 0.5 (see classification report and confusion matrix code, predicted > 0.5) the optimal value for making decisions?

Q: How sensitivity and specificity are usually tuned for diagnostic screening purposes?