# MachineLearningCode

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# 1 CSC8635 Machine Learning Extended Technical Report

# 1.1 What is the need for the project?

'Training of neural networks for automated diagnosis of pigmented skin lesions is hampered by the small size and lack of diversity of available datasets of dermatoscopic images.' (Tschandl, Rosendahl and Kittler, 2018). In this project, I will be exploring two machine learning models and running a comparison between them in order to choose the more suitable model for the project domain. As there is such 'a small size and lack of diversity' of datasets surrounding this area, the project will provide good groundwork for future projects to base off of.

'There are few comprehensive empirical studies comparing learning algorithms' (Caruana and Niculescu-Mizil, 2006). Machine learning comparisons are few and far between. In comparison with the small work put towards datsets in this domain, I believe that a comparison of machine learning technologies in this area would prove useful in automating skin cancer diagnosis with dermatoscopic images. 'I am using the paper 'An empirical comparison of supervised learning algorithms' as a basis for the comparison, and while the paper provides a much more detailed comparison than I am likely to need it still provides a solid background and good practises to follow for the comparison.

### 1.2 What did I do?

In order to help achieve success in this project, a methodology was needed to follow a clear structure. KDD was decided as a suitable methodology for this project, as a large portion of the methodology is dedicated to data miniming i.e. machine learning in this case.

Cookie Cutter was used to help structure the project in a reproducable format as well as providing ease of use for navigating around the directory. A cookie cutter build for data science was found that was excellent for the project at hand, and therefore used (Medium, 2019).

GitHub was used for version control and also allowed for backups in case of unforeseen errors. Due to the limited time available for the project and often massive runtimes of each model, only a two models were able to be compared. In this particular instance, I compared a custom built Convolutional Neural Network and a Virtual Geometry Group (VGG) pre-defined neural network (Simonyan and Zisserman, 2014).

# 1.2.1 Learning the Application Domain

In order to understand the domain and the datasets of the project, I found it useful to read the paper from the developers of the dataset itself, 'The HAM10000dataset, a large collection of multisource dermatoscopic images of common pigmented skin lesions' (Tschandl, Rosendahl and Kittler, 2018).

While an artificial neural network was already built to differentiate between melanomas and melanocytic nevi, there was a lack of data for both of these cell types and others.

The original datasets used to generate the datasets for this project were discovered to have bias towards melanocytic lesions, which is explored in my own project when plotting the dataset in the 'Plotting the Data' section.

While there do exists multi-class predictions for skin diseases, at the time of writing the paper there were no models for dermatoscopic images, and at the time of writing the report there are very few working models of which the validity of the results is questionable.

# 1.2.2 Sources used to help produce code

https://www.kaggle.com/kmader/dermatology-mnist-loading-and-processing https://www.kaggle.com/sid321axn/step-wise-approach-cnn-model-77-0344-accuracy?fbclid=IwAR2JWtc6\_nC3PZwfg5THA1qjTx1qaLc6NllbXIAftRkZQu\_AnQ6VMgaDsIs https://engmrk.com/vgg16-implementation-using-keras/

```
In [1]: # import libraries
        import os
        import numpy as np
        import pandas as pd
        from glob import glob
        import seaborn as sns
        import matplotlib.pyplot as plt
        from PIL import Image
        from sklearn.model_selection import train_test_split
        from keras.utils.np_utils import to_categorical
        from keras.models import Sequential
        from keras.layers import Dense, Dropout, Flatten, Conv2D, MaxPool2D, BatchNormalization
        from keras.optimizers import Adam
        from keras.callbacks import ReduceLROnPlateau
        from keras.preprocessing.image import ImageDataGenerator
        from sklearn.metrics import confusion_matrix
        from keras import applications
        from sklearn.linear_model import LogisticRegression
        from keras.layers import MaxPooling2D
```

Using TensorFlow backend.

Due to the nature of the document, it is not possible to associate every KDD step with the associated code, so some steps may be out of place but all relevant information is covered at some point in the report. I have aimed to include each step at the point where it was first used.

Creating a Target Dataset

As the dataset is quite large and the machine learning models can be very demanding in terms of resources, it is important that only relevant data was used. The metadata was used for the exploratory data analysis process, but ultimately only the images were used for the machine learning process itself. The source location

Data Cleaning and Preprocessing

In [5]: # dictionary for image paths

While it was initially thought that there would be a large amount of data cleaning and preprocessing, after accelerating past this step it was noticed that data cleaning was mostly unneccesary for the project, as explained later in the report. While it was useful to understand the dataset further, problems like this highlight the issues of KDD.

Relevant data cleaning is discussed later in the discussion for the models.

Image paths were assigned to the metadata dataframe in order to provide a 'merge' and connect the data.

A dictionary was produced for readability during the exploratory data analysis phase and general ease of use throughout the process.

```
In [7]: # dictionary to rename cell types for readability
    lesion_type_dict = {
        'nv': 'Melanocytic Nevi',
        'mel': 'Melanoma',
        'bkl': 'Benign Keratosis-like Lesions ',
        'bcc': 'Basal Cell Carcinoma',
        'akiec': 'Bowens Disease',
        'vasc': 'Vascular Lesions',
        'df': 'Dermatofibroma'
}
```

Cells were assigned an ID to be used as a response variable later in the project.

```
In [8]: # add column for path of image
    metadata_df['path'] = metadata_df['image_id'].map(imageid_path_dict.get)
    # add column for cell types, for readability
    metadata_df['cell_type'] = metadata_df['dx'].map(lesion_type_dict.get)
    # assign each cell type an ID and add it to a column
    metadata_df['cell_type_id'] = pd.Categorical(metadata_df['cell_type']).codes
```

Some introductory exploratory data analysis was performed in order to get a feel and general understanding for the data. This included use of the head, tail, and columns functions.

```
In [9]: # print first 5 values of dataframe
    print(metadata_df.head(5))
```

```
lesion_id
                   image_id
                              dx dx_type
                                                 sex localization \
                                           age
O HAM_0000118 ISIC_0027419
                             bkl
                                   histo 80.0 male
                                                           scalp
1 HAM_0000118 ISIC_0025030
                                   histo 80.0 male
                             bkl
                                                           scalp
2 HAM_0002730
               ISIC_0026769
                             bkl
                                   histo 80.0 male
                                                           scalp
3 HAM_0002730
               ISIC_0025661
                             bkl
                                   histo 80.0 male
                                                           scalp
4 HAM_0001466
               ISIC_0031633
                             bkl
                                   histo 75.0 male
                                                             ear
                                     path
                                                                cell_type
O HAM10000_images_part_1\ISIC_0027419.jpg
                                           Benign Keratosis-like Lesions
1 HAM10000_images_part_1\ISIC_0025030.jpg
                                           Benign Keratosis-like Lesions
2 HAM10000_images_part_1\ISIC_0026769.jpg
                                           Benign Keratosis-like Lesions
  HAM10000_images_part_1\ISIC_0025661.jpg
3
                                           Benign Keratosis-like Lesions
  HAM10000_images_part_2\ISIC_0031633.jpg
                                           Benign Keratosis-like Lesions
   cell_type_id
0
             1
1
2
             1
3
4
             1
```

# 

```
lesion_id
                       image_id
                                    dx dx_type
                                                 age
                                                         sex localization
10010 HAM_0002867
                   ISIC_0033084 akiec
                                         histo 40.0
                                                                  abdomen
                                                        male
                   ISIC_0033550
                                         histo 40.0
10011 HAM_0002867
                                 akiec
                                                        male
                                                                  abdomen
10012 HAM_0002867
                   ISIC_0033536
                                 akiec
                                         histo 40.0
                                                        male
                                                                  abdomen
10013 HAM_0000239
                   ISIC_0032854
                                 akiec
                                         histo 80.0
                                                        male
                                                                     face
10014 HAM_0003521
                   ISIC_0032258
                                         histo 70.0 female
                                   mel
                                                                     back
                                         path
                                                    cell_type
                                                              cell_type_id
10010 HAM10000_images_part_2\ISIC_0033084.jpg Bowens Disease
10011 HAM10000_images_part_2\ISIC_0033550.jpg Bowens Disease
                                                                          2
10012 HAM10000_images_part_2\ISIC_0033536.jpg Bowens Disease
                                                                          2
```

```
10013 HAM10000_images_part_2\ISIC_0032854.jpg Bowens Disease 2
10014 HAM10000_images_part_2\ISIC_0032258.jpg Melanoma 5
```

This allows us to see the first and last 5 values of the metadata dataset, and get a general understanding of how the data is formatted.

```
In [11]: # print column names
         print(metadata_df.columns)
Index(['lesion_id', 'image_id', 'dx', 'dx_type', 'age', 'sex', 'localization',
       'path', 'cell_type', 'cell_type_id'],
      dtype='object')
In []: # print columns and their data type
        print(metadata_df.dtypes)
In [12]: # summary of numeric values
         print(metadata_df.describe())
                    cell_type_id
               age
       9958.000000
                    10015.000000
count
         51.863828
                        3.528208
mean
         16.968614
                         1.377071
std
min
          0.000000
                        0.000000
25%
         40.000000
                         4.000000
50%
         50.000000
                        4.000000
         65.000000
75%
                        4.000000
         85.000000
                         6.000000
max
```

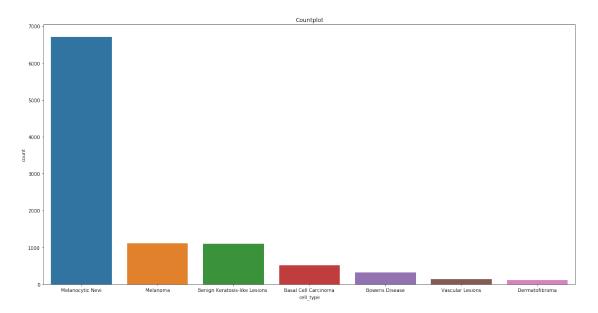
While the mean age is calculated as 52, as the ages in the dataset are given in intervals of 5, this may not actually be a true representation of the ages. However, for our purposes with the dataset as long as the age is kept consistent throughout this is not a large issue, although in order to get a more precise model in a future project it may help to give the exact ages.

Despite images only being used for the final model, data cleaning was still used due to the methodologys structure and general good practise. While it was good practise, for my particular models these steps were irrelevant for training the models and optimising accuracy. The documentation for this data cleaning and commented code has been left in the report for reproducability. This also provides a basis for future work on the project if someone chose to include the meta data in their model, which may be beneficial for comparing image classification and extracting features from the image and classifying using the data.

For data cleaning, it is often important to deal with null/NA values from data to remove redundant rows from the dataset. In this situation, as age was the only column with null values I opted to replace them with the mean value of the age column. In order to improve accuracy further, it may be worthwhile in a future project to look at other factors that my affect age and calculate the age based of that e.g. are certain ages more likely to have a certain cell type? While not null values, I removed any rows that had unknown values (prevalent in the sex and localization columns) to maximise information gain.

```
In [13]: # gives us data types and how many values of each column are non-null
         print(metadata_df.info())
<class 'pandas.core.frame.DataFrame'>
RangeIndex: 10015 entries, 0 to 10014
Data columns (total 10 columns):
lesion_id
                10015 non-null object
image id
                10015 non-null object
                10015 non-null object
dx
                10015 non-null object
dx_type
                9958 non-null float64
age
                10015 non-null object
sex
localization
               10015 non-null object
                10015 non-null object
path
cell_type
                10015 non-null object
               10015 non-null int8
cell_type_id
dtypes: float64(1), int8(1), object(8)
memory usage: 714.0+ KB
None
In [14]: # fill null values of age with the mean value of the age column
         \#metadata\_df["age"] = metadata\_df["age"].fillna(int(metadata\_df["age"].mean()))
In [15]: # display unique values in cell type column
         print(metadata_df.dx.unique())
['bkl' 'nv' 'df' 'mel' 'vasc' 'bcc' 'akiec']
In [16]: # display unique values in diagnosis column
         print(metadata_df.dx_type.unique())
['histo' 'consensus' 'confocal' 'follow_up']
In [17]: # display unique values in sex column
         print(metadata_df.sex.unique())
['male' 'female' 'unknown']
In [18]: # drop rows where sex = unknown
         metadata_df = metadata_df [metadata_df.sex!='unknown']
In [19]: # display unique values in sex column
         print(metadata_df.localization.unique())
['scalp' 'ear' 'face' 'back' 'trunk' 'chest' 'upper extremity' 'abdomen'
 'unknown' 'lower extremity' 'genital' 'neck' 'hand' 'foot' 'acral']
In [20]: # drop rows where localization = unknown
         metadata_df = metadata_df[metadata_df.localization!='unknown']
```

**Plotting the Data** Data was plotted to allow further exploration in the dataset as well as giving the oppurtunity to spot any patterns that the dataset may contain.



Out[22]: lesion\_id image\_id dx dx\_type age \ cell\_type Basal Cell Carcinoma Benign Keratosis-like Lesions Bowens Disease Dermatofibroma Melanocytic Nevi Melanoma Vascular Lesions sex localization path cell\_type\_id cell\_type Basal Cell Carcinoma 

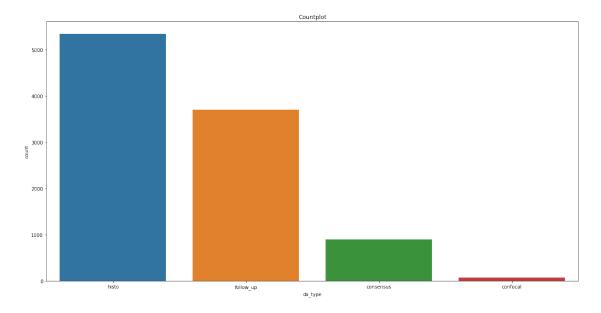
In [22]: metadata\_df.groupby('cell\_type').count()

Benign Keratosis-like Lesions

1099 1099

Bowens Disease	327	327	327	327
Dermatofibroma	115	115	115	115
Melanocytic Nevi	6705	6705	6705	6705
Melanoma	1113	1113	1113	1113
Vascular Lesions	142	142	142	142

Here we can see that there are a much larger amount of Melanocytic Nevi cells than any other type, with approximately 6 times more values then the next most frequent cell, melanoma. The least frequent cell type is Dermatofibroma. This means that there is a huge range of 6384. This is important to consider for the machine learning models, as it may lead to overfitting. For example, if the model was incorrect and predicted every image as Melanocytic Nevi, it would result in a high accuracy for the training data but low accuracy for the test data. This also stresses the importance of using a confusion matrix, as it allows me to verify the validity of my results.

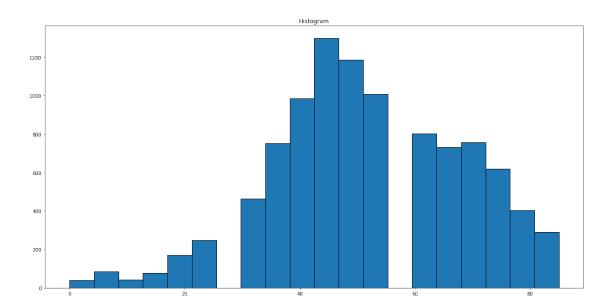


A countplot of the diagnosis type was produced, which allows us to see how the cells in the images were diagnosed. Although this will not necessarily affect our results, it is worth noting that this gives us some insights into how the cancer is diagnosed and potential false diagnoses.

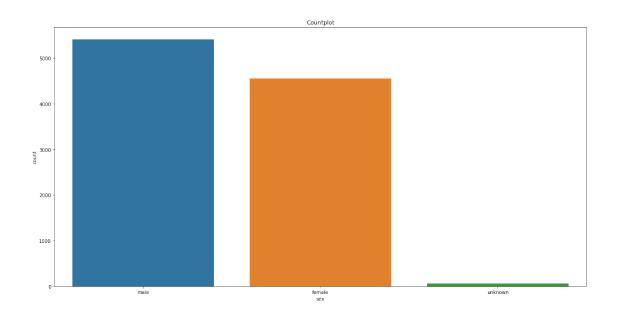
```
In [24]: # histogram of age
     plt.figure(figsize=(20, 10))
```

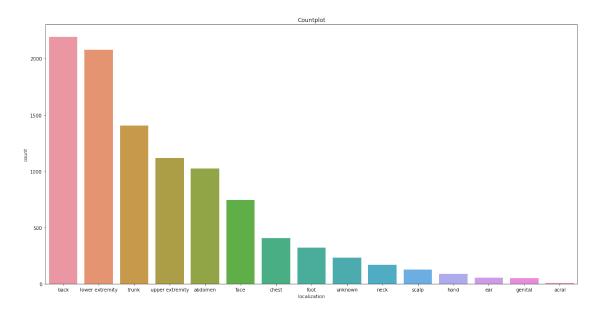
```
plt.hist(metadata_df['age'], bins = 20, histtype='bar', ec='black')
plt.title("Histogram")
plt.show()
```

- C:\Users\b5034806\AppData\Local\Continuum\anaconda3\envs\MLEnv3\lib\site-packages\numpy\lib\hi
  keep = (tmp\_a >= first\_edge)
- C:\Users\b5034806\AppData\Local\Continuum\anaconda3\envs\MLEnv3\lib\site-packages\numpy\lib\hi
  keep &= (tmp\_a <= last\_edge)</pre>



To see the age distribution of the dataset, a histogram was produced. As we can see, the majority of people are in the 40-60 range, with very few people under the age of 20 included in the dataset.





# 1.2.3 Data Reduction and Projection

As neural networks are a natural choice for images and models excel without extracting the features, the metadata was not included for the final model and only the images were used.

# 1.2.4 Choosing Function of Data Mining

Using the background reading, it was clear that this is a classification problem and that the models should predict the cell type, similarly to the old artificial neural network discussed in the 'Learning the application domain' section. The models will essentially be an expansion on that, with five more cell types.

## 1.2.5 Choosing the data mining algorithm

I have decided that CNN and VGG will be a natural fit for this project, as they are both convolutional neural networks which are a natural solution for image classification, and it allows me to compare a custom built model with a pre-defined model.

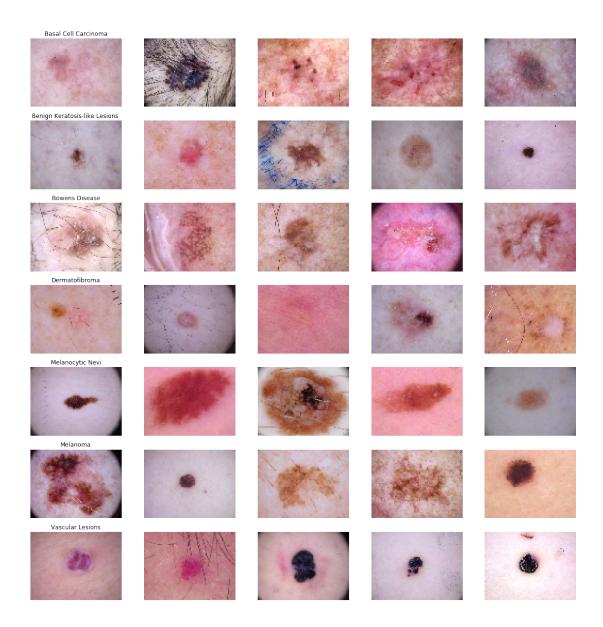
# 1.2.6 Data mining

The data mining process is discussed in depth throughout the discussion of the models:

#### 1.2.7 Convolutional Neural Network

CNNs are based upon neural networks in that they are based upon neurons. However, they differ from neural networks by taking in a multi chaneled image, which is suitable for our dataset as the images are RGB values which is read in as a 600x600x3 array, which equates to 3 chanels. CNNs are composed of convolutional, nonlinear, pooling and fully connected layers. The convolution layer works by convolving a filter around the image and computing element wise multiplications based upon the receptive field, which is summed up and repeated for each location on the image, which results in the feature map. This is how the model identified the features. The fully connected layer works by taking the high level features and associates these features with a class. In order to train a CNN, back propogation is used.

While all of the images are the same size and image resizing is not necessary, due to the timescale of the project it was essential for me to make the image sizes smaller to allow for faster run times, although this ultimately could affect the accuracy of the models due to a loss of data in the images. I have therefore tried to balance run time and accuracy, and resized the images to 1/3 of the original size. In the future, given a larger timescale/ better resources I would like to rexplore the dataset using the original image sizes and compare the accuracies and see if there is any significant differences between the original and rescaled images. While not directly applicable to our dataset, it is also important to note that padding is inefficient for classification problems, as it may cost some epochs for the neural network to calculate that there is no correlation in the black pixels using gradient descent (Howard, 2019).



The predictor variables and response variable were split, so that the model would attempt to learn what cell type was in the image i.e. the image ID.

It was important to find a good balance between the training and test split, as a split too large will result in greater performance statistic variance, whereas a split too small will result in greater parameter estimate variance.

```
In [90]: # create training and test data for x and y variables at a 75:25 ratio
x_train, x_test, y_train, y_test = train_test_split(predictor, response, test_size=0.5
```

Data was normalised to get the values on a common scale, while also not distorting the differences in the range.

```
In [91]: # normalise data
         x_train = np.asarray(x_train["image"].tolist())
         x_test = np.asarray(x_test['image'].tolist())
         x_train = (x_train - np.mean(x_train)) / (np.std(x_train))
         x_test = (x_test - np.mean(x_test) / np.std(x_test))
In [92]: # split data into training and validation data set at a 70:30 ratio
         \#x\_train, x\_validate, y\_train, y\_validate = train\_test\_split(x\_train, y\_train, test\_s)
```

One hot encoding was performed on the labels in order to allow the models to bypass the issue of categorical values.

```
In [93]: # one hot encoding
         y_train = to_categorical(y_train, num_classes = 7)
         y_test = to_categorical(y_test, num_classes = 7)
```

A canal value of 3 was used for the convolutional layer, as previously explained in the 'Convolutional Neural Network' section.

```
In [94]: # canal value of 3 as images are RGB
         x_train = x_train.reshape(x_train.shape[0], *(75, 100, 3))
         x_{test} = x_{test.reshape}(x_{test.shape}[0], *(75, 100, 3))
In [95]: # thanks to https://github.com/yuguan1/example-ML-code/blob/master/DL1/CNN.ipynb
         # set the CNN model
         input_shape = (75, 100, 3)
         nClasses = 7
         model = Sequential()
         model.add(Conv2D(32, kernel_size=(3, 3),
                          activation='relu',
                          input_shape=(input_shape)))
         model.add(MaxPooling2D(pool_size=(2, 2)))
         model.add(Conv2D(64, (3, 3), activation='relu'))
         model.add(MaxPooling2D(pool_size=(2, 2)))
         model.add(Dropout(0.25))
         model.add(Flatten())
         model.add(Dense(128, activation='relu'))
         model.add(Dropout(0.5))
         model.add(Dense(nClasses, activation='softmax'))
         model.summary()
                                            Param #
```

Output Shape

Layer (type)

conv2d_9 (Conv2D)	(None,	73, 98,	32)	896
max_pooling2d_9 (MaxPooling2	(None.	 36, 49,	 32)	0
conv2d_10 (Conv2D)	(None,	34, 47,	64)	18496
max_pooling2d_10 (MaxPooling	(None,	17, 23,	64)	0
dropout_9 (Dropout)	(None,	17, 23,	64)	0
flatten_5 (Flatten)	(None,	25024)		0
dense_9 (Dense)	(None,	128)		3203200
dropout_10 (Dropout)	(None,	128)		0
dense_10 (Dense)	(None,	7)		903
Total params: 3,223,495				
Trainable params: 3,223,495 Non-trainable params: 0				

The adam optimiser is a generalised optimizer that works for many models, and is particularly efficient for deep learning.

A learning rate annealer was used to stablise the learning rate. This was used as initially, the learning rate is too high which causes a high variance in accuracy and therefore divergant results. Stabilising the learning rate over time reduces the impact of this. It is also worth noting that having the learning rate too slow would require many epochs before reaching the minimum point on the gradient descent.

Data augmentation is used to prevent overfitting. As the dataset is not particularly large, data augmentation helps to prevent overfitting by generating relevant images with slight variations, to provide more data for the dataset and the variations help to encourage generalisation and therefore reduces the likelihood of overfitting.

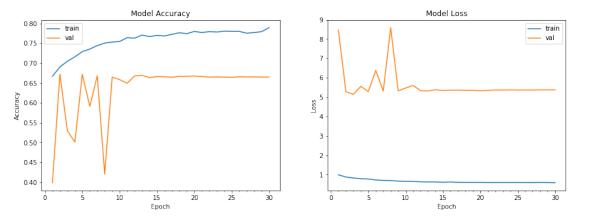
'It has been observed in practice that when using a larger batch there is a significant degradation in the quality of the model...' (Keskar, Mudigere, Nocedal, Smelyanskiy and Tang, 2016). After reading this paper, it was clear that increasing the batch size too much would not be good, due to generalization. Too low of a batch size would result in a low accuracy and a slow computational process. It is also important to consider that there may be interactions with other hyperparameters. Therefore, epochs and batch sizes were experimented with to find the best balance between accuracy and speed.

```
In [99]: # train\ the\ model\ CNN\ RESULTS
   epochs = 30
   batch_size = 16
   history = model.fit_generator(datagen.flow(x_train,y_train, batch_size=batch_size),
               epochs = epochs, validation_data = (x_test,y_test),
               verbose = 1, steps_per_epoch=x_train.shape[0] // batch_
                , callbacks=[learning_rate_reduction])
Epoch 1/30
Epoch 2/30
Epoch 3/30
Epoch 4/30
Epoch 5/30
Epoch 00005: ReduceLROnPlateau reducing learning rate to 0.00050000000237487257.
Epoch 6/30
Epoch 7/30
Epoch 8/30
Epoch 00008: ReduceLROnPlateau reducing learning rate to 0.0002500000118743628.
Epoch 9/30
Epoch 10/30
```

```
Epoch 11/30
Epoch 00011: ReduceLROnPlateau reducing learning rate to 0.0001250000059371814.
Epoch 12/30
Epoch 13/30
Epoch 14/30
Epoch 00014: ReduceLROnPlateau reducing learning rate to 6.25000029685907e-05.
Epoch 15/30
Epoch 16/30
Epoch 17/30
Epoch 00017: ReduceLROnPlateau reducing learning rate to 3.125000148429535e-05.
Epoch 18/30
Epoch 19/30
Epoch 20/30
Epoch 00020: ReduceLROnPlateau reducing learning rate to 1.5625000742147677e-05.
Epoch 21/30
Epoch 22/30
Epoch 23/30
Epoch 00023: ReduceLROnPlateau reducing learning rate to 1e-05.
Epoch 24/30
Epoch 25/30
Epoch 26/30
Epoch 27/30
Epoch 28/30
Epoch 29/30
```

```
Epoch 30/30
- 10s 21ms/step - loss: 0.5778 - acc: 0.7894 - val_los
In [100]: # thanks to https://www.kaggle.com/sid321axn/step-wise-approach-cnn-model-77-0344-ac
                          # function to plot model's validation loss and validation accuracy
                         def plot_model_history(model_history):
                                    fig, axs = plt.subplots(1,2,figsize=(15,5))
                                    # summarize history for accuracy
                                    axs[0].plot(range(1,len(model_history.history['acc'])+1),model_history.history['acc'])
                                    axs[0].plot(range(1,len(model_history.history['val_acc'])+1),model_history.history
                                    axs[0].set_title('Model Accuracy')
                                    axs[0].set_ylabel('Accuracy')
                                    axs[0].set_xlabel('Epoch')
                                    axs[0].set_xticks(np.arange(1,len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history['acc'])+1),len(model_history[
                                    axs[0].legend(['train', 'val'], loc='best')
                                    # summarize history for loss
                                    axs[1].plot(range(1,len(model_history.history['loss'])+1),model_history.history[
                                    axs[1].plot(range(1,len(model_history.history['val_loss'])+1),model_history.history.history.history.history.
                                    axs[1].set_title('Model Loss')
                                    axs[1].set_ylabel('Loss')
                                    axs[1].set_xlabel('Epoch')
                                    axs[1].set_xticks(np.arange(1,len(model_history.history['loss'])+1),len(model_history['loss'])+1)
                                    axs[1].legend(['train', 'val'], loc='best')
                                   plt.show()
```

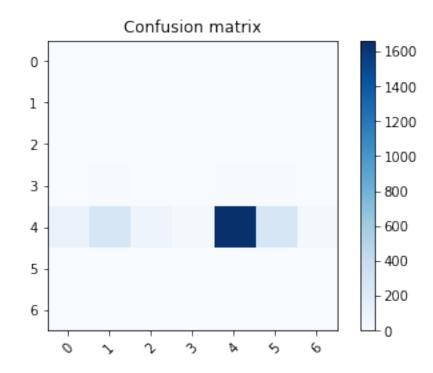
In [101]: plot\_model\_history(history)



From this plot we can immediately see how the learning rate annealer affects the model. Initially there are huge spikes in the accuracy for the test data, although after some epochs this stablises roughly at its peak (Approx 66%) due to the learning rate being reduced. This also applies to the test data's loss, which we can see stablises at the same Epoch as the accuracy. While the accuracy score is okay, the loss value is incredibly high and suggests that a lot of errors are being made every epoch, which may be due to the overfitting which will be explored with the next plot.

```
In [104]: # thanks to https://www.kaggle.com/sid321axn/step-wise-approach-cnn-model-77-0344-ac
          y_pred = model.predict(x_test)
          y_pred = np.argmax(y_pred, axis = 1)
          y_true = np.argmax(y_test,axis = 1)
          confusion = confusion_matrix(y_pred, y_true)
          # Function to plot confusion matrix
          def plot_confusion_matrix(cm, classes,
                                    normalize=False,
                                    title='Confusion matrix',
                                    cmap=plt.cm.Blues):
              HHHH
              This function prints and plots the confusion matrix.
              Normalization can be applied by setting `normalize=True`.
              plt.imshow(cm, interpolation='nearest', cmap=cmap)
              plt.title(title)
              plt.colorbar()
              tick_marks = np.arange(len(classes))
              plt.xticks(tick_marks, classes, rotation=45)
              plt.yticks(tick_marks, classes)
              if normalize:
                  cm = cm.astype('float') / cm.sum(axis=1)[:, np.newaxis]
              thresh = cm.max() / 2.
              for i, j in itertools.product(range(cm.shape[0]), range(cm.shape[1])):
                  plt.text(j, i, cm[i, j],
                           horizontalalignment="center",
                           color="white" if cm[i, j] > thresh else "black")
              plt.tight_layout()
              plt.ylabel('True label')
              plt.xlabel('Predicted label')
          plot_confusion_matrix(confusion, classes = range(7))
        NameError
                                                  Traceback (most recent call last)
        <ipython-input-104-e8c4038ddc62> in <module>
         34
                plt.xlabel('Predicted label')
         35
    ---> 36 plot_confusion_matrix(confusion, classes = range(7))
```

NameError: name 'itertools' is not defined



Unfortunately, the confusion matrix confirms that the model has been overfitted, as the model is only predicted for cell id 4, Melanocytic Nevi, which was previously discussed to potentially cause overfitting. As there is a significantly higher amount of Melanocytic Nevi than any other cell, the machine gets into the habit of only predicting these values, which likely explains the high loss experienced in the model.

#### 1.2.8 VGG Neural Network

The VGG Neural Network is a pre-defined neural network which was developed by 2 Oxford Students for a neural network competition, and received first place for image identification and 2nd place for image classification. It is popular for being a very efficient CNN and therefore in theory should perform better than the custom built CNN.

```
# images rescaled to 224x224 for VGG
         metadata_df['image'] = metadata_df['path'].map(lambda x: np.asarray(Image.open(x).res
         # change working directory back
         os.chdir(r"C:\Users\b5034806\Documents\Machine-Learning-Extended-Technical-Project\mle
In [28]: # get predictor variables and add them to a dataframe
         predictor = metadata_df.drop(columns=['cell_type_id'],axis=1)
         # get response variables and add them to a series
         response = metadata_df['cell_type_id']
In [29]: # create training and test data for x and y variables at a 75:25 ratio
         x_train, x_test, y_train, y_test = train_test_split(predictor, response, test_size=0.3
In [30]: # normalise data
         x_train = np.asarray(x_train["image"].tolist())
         x_test = np.asarray(x_test['image'].tolist())
         x_train = (x_train - np.mean(x_train)) / (np.std(x_train))
         x_test = (x_test - np.mean(x_test) / np.std(x_test))
In [31]: # Perform one-hot encoding on the labels
         y_train = to_categorical(y_train, num_classes = 7)
         y_test = to_categorical(y_test, num_classes = 7)
In [32]: # reshape images for VGG
         x_train = x_train.reshape(x_train.shape[0], *(224, 224, 3))
         x_{test} = x_{test.reshape}(x_{test.shape}[0], *(224, 224, 3))
In [33]: # thanks to https://engmrk.com/vgg16-implementation-using-keras/
         # set the VGG model
         input_shape = (224, 224, 3)
         model = Sequential([
             Conv2D(64, (3, 3), input_shape=input_shape, padding='same',
                    activation='relu'),
             Conv2D(64, (3, 3), activation='relu', padding='same'),
             MaxPooling2D(pool_size=(2, 2), strides=(2, 2)),
             Conv2D(128, (3, 3), activation='relu', padding='same'),
             Conv2D(128, (3, 3), activation='relu', padding='same',),
             MaxPooling2D(pool_size=(2, 2), strides=(2, 2)),
             Conv2D(256, (3, 3), activation='relu', padding='same',),
             Conv2D(256, (3, 3), activation='relu', padding='same',),
             Conv2D(256, (3, 3), activation='relu', padding='same',),
             MaxPooling2D(pool_size=(2, 2), strides=(2, 2)),
             Conv2D(512, (3, 3), activation='relu', padding='same',),
             Conv2D(512, (3, 3), activation='relu', padding='same',),
             Conv2D(512, (3, 3), activation='relu', padding='same',),
```

```
MaxPooling2D(pool_size=(2, 2), strides=(2, 2)),
Conv2D(512, (3, 3), activation='relu', padding='same',),
Conv2D(512, (3, 3), activation='relu', padding='same',),
Conv2D(512, (3, 3), activation='relu', padding='same',),
MaxPooling2D(pool_size=(2, 2), strides=(2, 2)),
Flatten(),
Dense(4096, activation='relu'),
Dense(4096, activation='relu'),
Dense(7, activation='softmax')
```

# model.summary()

228
  56
7584
168
080
080
80160
9808
9808
9808
9808

```
flatten 1 (Flatten)
                 (None, 25088)
    ______
dense 1 (Dense)
                     (None, 4096)
                                          102764544
-----
dense_2 (Dense)
                     (None, 4096)
                                          16781312
              (None, 7) 28679
dense_3 (Dense)
______
Total params: 134,289,223
Trainable params: 134,289,223
Non-trainable params: 0
In [34]: # compile the model
       model.compile(optimizer = 'adam', loss = "categorical_crossentropy", metrics=["accura
In [35]: # Set a learning rate annealer
       learning_rate_reduction = ReduceLROnPlateau(monitor='val_acc',
                                         patience=3,
                                         verbose=1,
                                         factor=0.5,
                                         min_lr=0.00001)
In [36]: # data augmentation
       datagen = ImageDataGenerator(
             rotation_range=30, # randomly rotate images in the range (degrees, 0 to 180)
             horizontal_flip=True, # randomly flip images
             vertical_flip=True) # randomly flip images
       datagen.fit(x_train)
In [37]: # train the model VGG RESULTS
       # epochs reduced and batch size increased due to slow runtime
       epochs = 20
       batch_size = 32
       history2 = model.fit_generator(datagen.flow(x_train,y_train, batch_size=batch_size),
                              epochs = epochs, validation_data = (x_test,y_test),
                              verbose = 1, steps_per_epoch=x_train.shape[0] // batch_
                               , callbacks=[learning_rate_reduction])
Epoch 1/20
```

0

conv2d\_13 (Conv2D) (None, 14, 14, 512) 2359808

\_\_\_\_\_

max\_pooling2d\_5 (MaxPooling2 (None, 7, 7, 512)

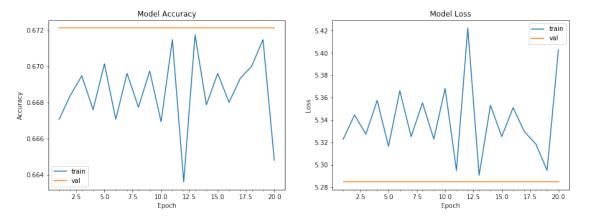
```
Epoch 3/20
Epoch 4/20
Epoch 00004: ReduceLROnPlateau reducing learning rate to 0.0005000000237487257.
Epoch 5/20
Epoch 6/20
Epoch 7/20
Epoch 00007: ReduceLROnPlateau reducing learning rate to 0.0002500000118743628.
Epoch 8/20
Epoch 9/20
Epoch 00010: ReduceLROnPlateau reducing learning rate to 0.0001250000059371814.
Epoch 11/20
Epoch 12/20
Epoch 13/20
Epoch 00013: ReduceLROnPlateau reducing learning rate to 6.25000029685907e-05.
Epoch 14/20
Epoch 15/20
Epoch 16/20
Epoch 00016: ReduceLROnPlateau reducing learning rate to 3.125000148429535e-05.
Epoch 17/20
Epoch 18/20
Epoch 19/20
```

Epoch 2/20

Epoch 00019: ReduceLROnPlateau reducing learning rate to 1.5625000742147677e-05.

```
In [39]: # thanks to https://www.kaggle.com/sid321axn/step-wise-approach-cnn-model-77-0344-acc
         # function to plot model's validation loss and validation accuracy
         def plot_model_history(model_history):
             fig, axs = plt.subplots(1,2,figsize=(15,5))
              # summarize history for accuracy
             axs[0].plot(range(1,len(model_history.history['acc'])+1),model_history.history['a
             axs[0].plot(range(1,len(model_history.history['val_acc'])+1),model_history.history
             axs[0].set_title('Model Accuracy')
             axs[0].set_ylabel('Accuracy')
             axs[0].set_xlabel('Epoch')
             axs[0].set_xticks(np.arange(1,len(model_history.history['acc'])+1),len(model_history.history['acc'])+1),
             axs[0].legend(['train', 'val'], loc='best')
              # summarize history for loss
             axs[1].plot(range(1,len(model_history.history['loss'])+1),model_history.history['.
             axs[1].plot(range(1,len(model_history.history['val_loss'])+1),model_history.history
             axs[1].set_title('Model Loss')
             axs[1].set_ylabel('Loss')
             axs[1].set_xlabel('Epoch')
             axs[1].set_xticks(np.arange(1,len(model_history.history['loss'])+1),len(model_history.history['loss'])+1)
             axs[1].legend(['train', 'val'], loc='best')
             plt.show()
```

In [41]: plot\_model\_history(history2)



Even though VGG was not successful in working with the test data, we are still able to perform an analysis on the training data. Despite providing data annealing, the variation is high throughout every epoch suggesting that the learning rate was still too high, and data annealing needed to be applied more aggressively. On the other hand, you can actually see that the variance is only for a very small range and in reality the learning rate may not have been high enough to begin with. Even though the accuracy lies at a respectable 67.21%, as the validation accuracy and loss

are flat it is likely that the model does not work as intended and the model is not actually learning anything about the images over time. This could potentially be explained to a very low training rate, and is something to keep in consideration for future implementation of VGG for this dataset. Due to VGG being quite demanding in terms of resources and taking large amounts of time each time the model is trained it was not possible to tune the parameters in many ways and try to 'fix' the model. This is something that can be explored in future projects.

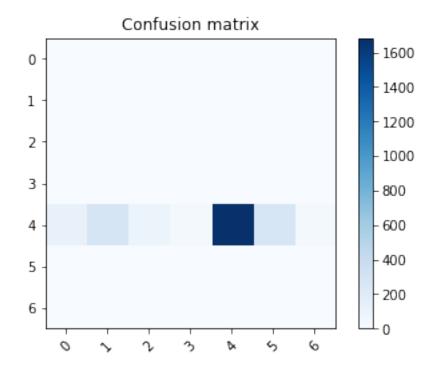
In [42]: # thanks to https://www.kaggle.com/sid321axn/step-wise-approach-cnn-model-77-0344-acc y\_pred = model.predict(x\_test) y\_pred = np.argmax(y\_pred, axis = 1) y\_true = np.argmax(y\_test,axis = 1) confusion = confusion\_matrix(y\_pred, y\_true) # Function to plot confusion matrix def plot\_confusion\_matrix(cm, classes, normalize=False, title='Confusion matrix', cmap=plt.cm.Blues): 11 11 11 This function prints and plots the confusion matrix. Normalization can be applied by setting `normalize=True`. plt.imshow(cm, interpolation='nearest', cmap=cmap) plt.title(title) plt.colorbar() tick\_marks = np.arange(len(classes)) plt.xticks(tick\_marks, classes, rotation=45) plt.yticks(tick\_marks, classes) if normalize: cm = cm.astype('float') / cm.sum(axis=1)[:, np.newaxis] thresh = cm.max() / 2.for i, j in itertools.product(range(cm.shape[0]), range(cm.shape[1])): plt.text(j, i, cm[i, j], horizontalalignment="center", color="white" if cm[i, j] > thresh else "black") plt.tight\_layout() plt.ylabel('True label') plt.xlabel('Predicted label') plot\_confusion\_matrix(confusion, classes = range(7))

-----

```
Traceback (most recent call last)
```

NameError: name 'itertools' is not defined

NameError



The confusion matrix for VGG is very similar to that of the custom built CNN, and also suggests that the model is overfitted, potentially giving an explanation for the results that were gathered.

# 1.2.9 Interpretation

While it is likely that the results are not accurate, a comparison will still be made with the assumption that the results are 'correct'. Only the test data is compared, as training data results are not very relevant for comparisons between models. While the custom built CNN ends with an accuracy of 66.49%, VGG ends with the slightly higher accuracy of 67.21%. Similarly, the custom built CNN ends with a loss of 5.3748, whereas VGG ends with a loss of 5.2847. Taking both of these into account, VGG has the highest accuracy and lowest loss by a small margin, making it slightly better than the custom built model. It is also worth noting that it was significantly slower to train VGG than the custom built CNN, and so if the dataset is large enough it may be worth the tradeoff for a small amount of optimisation in loss and accuracy to get much faster run times. However, to reiterate, it is likely that the models are not working as intended and these results should be treated cautiously.

# 1.3 How Successful was my Project?

While a conclusion was met that satisfied the initial goal, the project may be deemed as unsuccesful as the models are likely not functioning as intended. However, I would argue that the project has seen success in many ways. For example, the groundwork has been layed out for anyone to pick up the project, add in a model of their choosing that they would like to compare, fine-tune the hyperparameters and use the tools provided such as the graphs for the models and confusion matrices for a comparison between any model. Another example would be that I have not yet discovered an attempt for an implementation of VGG on this dataset, and therefore the groundwork has also been layed out for the projects code to be tuned in such a way that VGG may receive more desireable results.

If using 'The Logical Framework Method for Defining Project Success' (Baccarini, 1999), the project can be deemed a failure in terms of 'project management success' as the original 'deliverable' was not achieved as desired. However, the project success can be deemed as a success as groundwork has been provided for future work in this area and although it was not the original deliverable, a successful output has been produce. The paper also makes a very good point about success not being 'black or white' and it may be more suitable to deem this project as a partial success.

In the future, I believe that a large scale comparison is needed, as if the 'most efficient' model is found, it could potentially be used in place of diagnosing by eye providing automated and fast diagnoses for dermatoscopic images.

# 1.4 Reflection

This project also introduced me to many new technologies, which has been a pleasure. As this was my first large project with python, it was nice to learn a new programming language and the strengths and weaknesses that it comes with. Python was a very successful language for machine learning as it is widely supported and relatively simple to pick up, key for the short time scale of the project. As a Data Scientist, I believe that both Python and R will prove to be useful tools to use in future projects as they excel in many tasks that are required of data science projects. Anaconda was a great way to produce environments for the project and easily download and manage the packages required for the code. Jupyter Notebooks was a great tool for producing markdown documents such as this one, and provides numerous advantages and disadvantages over R, such as code being much easier to manage in my opinion, but also a lot more tedious to use with the

natural implementation of cells for programming. Cookie cutter lacks the automation that Project Template provides, although it adds a lot more customisation in that there are a huge amount of premade directories online, which allowed me to find the data science cookie cutter build which worked well for this project.

KDD was much more successful for this project than the cloud computing project, as it utilised all of the steps (cloud computing did not make use of the three data mining steps). However, that is not to say that it did not come without issues. As previously mentioned, the linearity of KDD can result in wasted time, or if followed precisely with no going back between steps can actually force a project to stop before starting another 'cycle' after the interpretation phase.

Ironically, the most straightforward part was likely implementing the models themselves (Although not to much success), as there are many existing frameworks that you can base yours off of. The most difficult part of the project for me was editing the models and data in a way such that accuracy and loss are optimised. When given a large dataset such as this one computation times can be quite slow which does not give many attempts at fine tuning the models.

For future projects, it is always worth following a methodology as it provides a natural work flow and clear objectives to achieve your goal, although I would like to explore a new methodology in my next project such as AGILE or SCRUM. It may also be worth considering my strengths/weaknesses for future projects, as while I attempted to tackle a quite difficult dataset in domains I have never worked with before (Medical and images for machine learning) which definitely improved my skills and abilities, it also led to the original goal not being met. Therefore, rather than taking on a whole new area at once, it may be wise to approach a project with more caution.

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