

CANCERS DIAGNOSIS PROJECT WITH CNN [CANCERAPY APPLICATION]

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Technical Report

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Abstract: This project is going to consider more than **three cancers** and tumors like breast tumor, skin cancer, and colorectal tumor for now and each model of them will aim to detect or to be more specific to classify the given data like an image (Radiology) which will be given as an input to the model which will be deployed as a mobile application and classify the cancer, or the tumor based on their different types. The problem of tumor classification for its malignancy level and the cancer for its type has been studied for the last few years and has a reasonable number of applications here and there for everyone who's trying to improve a model which has a reasonable level of predictability. There are enormous number of algorithms to be used in machine learning for such a classification problem and this would a great challenge to choose between, not only this we have also to figure out which hyperparameters for every model we are trying to deploy (Tuning Process). There are different number of algorithms and architectures to be used when dealing with neural networks models, starting with **conventional** neural network model with specific number of layers and nodes or units and apply different techniques to get the best test accuracy but as we are dealing with images (Computer Vision Model), it's better to use **CNN** which is a class of ANN as well, and it's predictable to be more accurate and less computation time as well. All of these models are modeled by the aid of TensorFlow framework for training and evaluating process, also it will be very effective for Converting the best model for each cancer or tumor to a TensorFlow Lite model to be deployed as mobile application using Flutter framework.

Keywords: Machine Learning, Computer Vision, Image Classification, Binary Classification, Multi-Class Classification Tuning, CNN, Neural Network, ANN, TensorFlow, Keras, Flutter, dart.



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1. MOTIVATION

Cancer is a **leading** cause of death worldwide, accounting for nearly 10 million deaths in 2020, or nearly one in six deaths [1], but the patient can handle that cancer or at least limit its symptoms with early detection which is the main key here.

The number of cases for every cancer or tumor changes significantly but its not all about the number of each case yearly, it's about the death rate for each one as well. According to [1], we are going to introduce the most dangerous cancers and tumors.

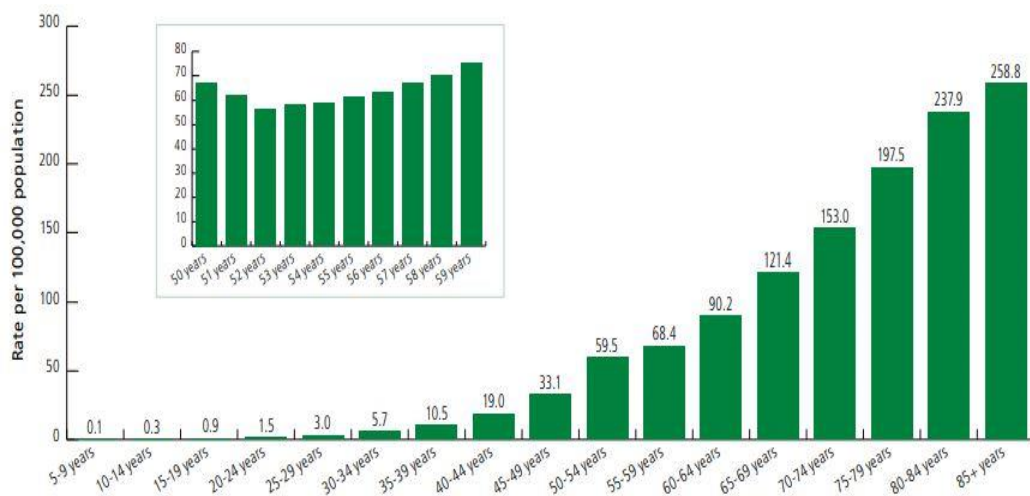
Table 1. Cancer New Cases and Deaths

New Cases		Cancer Deaths	
Breast	2.26 million	Lung	1.80 million
Lung	2.21 million	Colon and Rectum	916,000 deaths
Colon and Rectum	1.93 million	Liver & Stomach	900,000 deaths
Skin	1.2 million	Breast	685,000 deaths

First version of that project is including three types of cancers which has been bolded above in Table 1, which are Breast, Colorectal (Colon and Rectum Combined), and Skin Cancer and as time moves on, the project will hold most of known cancers with a reasonable dataset to be trained.

Breast Cancer is considered as the most common type of cancer between females and the first cause of cancer death among them In Egypt, it constitutes 33% of female cancer cases and more than 22,000 new cases diagnosed each year. This is expected to rise **exponentially** over the next years given the enlarging population [2]. **Early Detection and Diagnosis**, it's not just a combination of two words it's the key for treatment and the only way to be far away from the danger zone [3].

Approximately 4.4% of men (1 in 23) and 4.1% of women (1 in 25) will be diagnosed with **CRC** (Colon and Rectal Cancer) in their lifetime [4]. Lifetime risk is similar in men and women despite higher incidence rates in men because women have longer life expectancy. Age feature had a huge influence on CRC rates between population. The following figure is showing a left skewed distribution for rates of CRC with age.



Source: Main figure: NAACCR, 2019. Inset: Surveillance, Epidemiology, and End Results (SEER) Program, 2019.

Figure 1. CRC Rates with Age

Skin Cancer as well is one of the most dangerous cancers as most people at first stages ignores any symptoms on skin considering it as a normal lesion. Skin cancer is challenging case to be studied as it has large ratio of **misdiagnosed** cases as it contains multi classes to classify between and they are very close to each other, also the one type of them has dozens of shapes depends on diameter of lesions, the color as well, ... etc.

Therefore, it's important to develop an effective classifier for each cancer to detect or classify its type with the aid of machine learning. It could be of great help to automate the detection and locate tumor tissue cells and to speed up the process. This way one would be able to overcome the dependence on the pathologist which would be especially useful in regions where no experts are available but that's not the main aim of the project which will be discussed latter.

2. INOTRODUCTION

Machine learning algorithms with just few years after putting the eyes of interest on it, exponentially increased in terms of applications involved in and the accuracy of detecting as well. Computer vision is a great topic in machine learning which is simply machine with an eye, so if we are talking about images with tracing something on it or detecting which class the image belongs to, all of this is under the term computer vision.

Therefore, our project is considered as a computer vision application by training a large set of images of specific cancer as input, then deploy the model which will predict the type of that cancer immediately with reasonable percent of accuracy.

Neural Network is the main algorithm to be working on, images trained by taking each pixel as a single unit of input considering we have only one channel for simplicity. So, if we are talking about (200, 200) resolution by RGB mode three channels involved then we have about 120,000 features for the input layer only, also nowadays we have a variety and large number of datasets as every institution is trying hard to help with improving the diagnosis with computers.

All the above arguments makes classical machine learning algorithms fades in some applications, and the neural networks are leading and improving day by day. Figure 2 shows by a simple figure the concept of what we talked about.

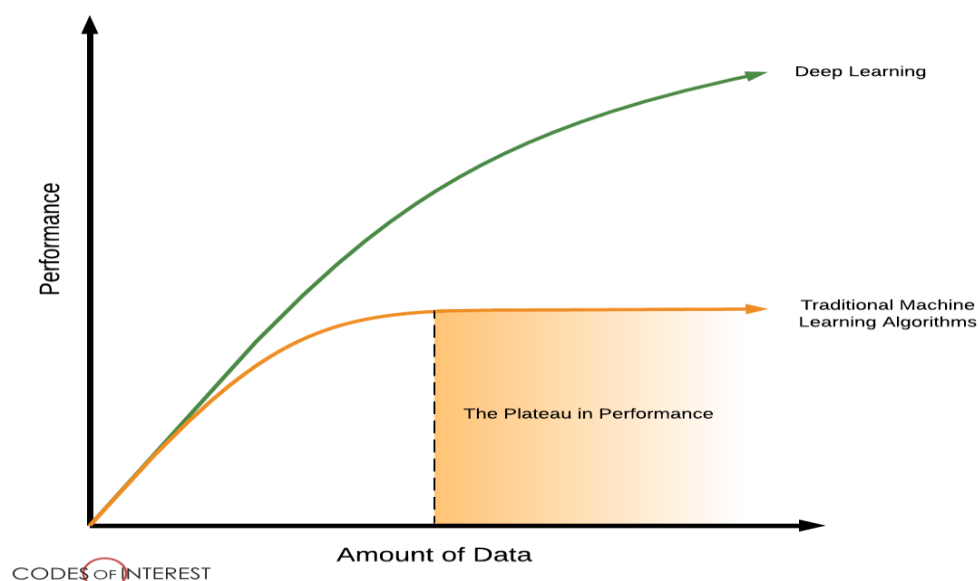


Figure 2. Performance of Classical ML and Deep Learning with Data

The Project (Cancers Classifier) Main goal is not just to help people at wild and abandoned areas, but hopefully deploying a versatile and large system with most of cancers on it to be deployed in every medical institution by uploading the radiology as usual but the system will automatically predict the cancer type or it's malignancy level and report all of that on the patient medical record and label it as a machine guess.

But for the first version of the project, it will not be embedded at any institution, it will just be deployed in a mobile application with quite good UI with flutter framework.

3. METHODS

3.1 Datasets

3.1.1 Breast Cancer Dataset

Our model trained on a dataset of **Mammography** images of INbreast database was originally collected from Centro Hospitalar de S. Joao [CHSJ], Breast center, Porto. INbreast database collects data from Aug. 2008 to July 2010, which contains 115 cases with a total of 410 images and becomes **7632 images** with augmentation process [5].

This dataset not just consist of 2 classes benign and malignant, but also the masses of dense in this mammography image of the breast, and the breast density is divided into four categories according to BI-RADS standards, which are Entirely fat (Density 1), Scattered fibro glandular densities (Density 2), Heterogeneously dense (Density 3), and extremely dense (Density 4), therefore we have 8 classes to classify between as shown in Table 2.

Table 2. Breast Cancer Classes

Benign	Malignant
Density 1	Density 1
Density 2	Density 2
Density 3	Density 3
Density 4	Density 4

3.1.2 Skin Cancer Dataset

Harvard Data verse introduces one of the biggest datasets for a certain cancer or tumor which is HAM10000 ("Human Against Machine with 10000 training images"), data is introduced as **Dermatoscopic** images from different populations, acquired and stored by different modalities [6].

Our model will train on three different classes of skin cancer which are melanoma (**Mel**), melanocytic nevi (**Nv**), and benign keratosis-like lesions (**bkl**), but unfortunately the three types of cancer are not balanced which will affect the prediction process as it is going to be biased, so we have mission here to do with data before processing.

3.1.3 Colorectal Cancer Dataset

This data set represents a collection of textures in histological images of human colorectal cancer. All images are RGB, 0.495 μm per pixel, digitized with an Aperio ScanScope (Aperio/Leica biosystems), magnification 20x. Histological samples are fully anonymized images of formalin-fixed paraffin-embedded human colorectal adenocarcinomas (primary tumors) from our pathology archive (Institute of Pathology, University Medical Center Mannheim, Heidelberg University, Mannheim, Germany) [7].

3.2 Models

The key difference between plain NN or Connected layers NN and CNN, is that for plain NN as shown in Figure 3, the input pass through each unit in the next layer by two steps one is linear step and the second is adding some non-linearity to the system (activation function) and the number of input in the previous layer is equal to the number of parameters of that node which are going to be trained, therefore it has a huge number of parameters to train.

The diagram illustrates a neural network architecture. It consists of four layers of nodes: an input layer with 3 orange nodes, two hidden layers each with 4 green nodes, and an output layer with 1 red node. Arrows indicate the flow of information from left to right, with every node in one layer connected to every node in the subsequent layer. Labels at the bottom identify the layers: 'INPUT LAYER' (orange), 'HIDDEN LAYERS' (green), and 'OUTPUT LAYER' (red).

Figure 3. Plain Neural Network

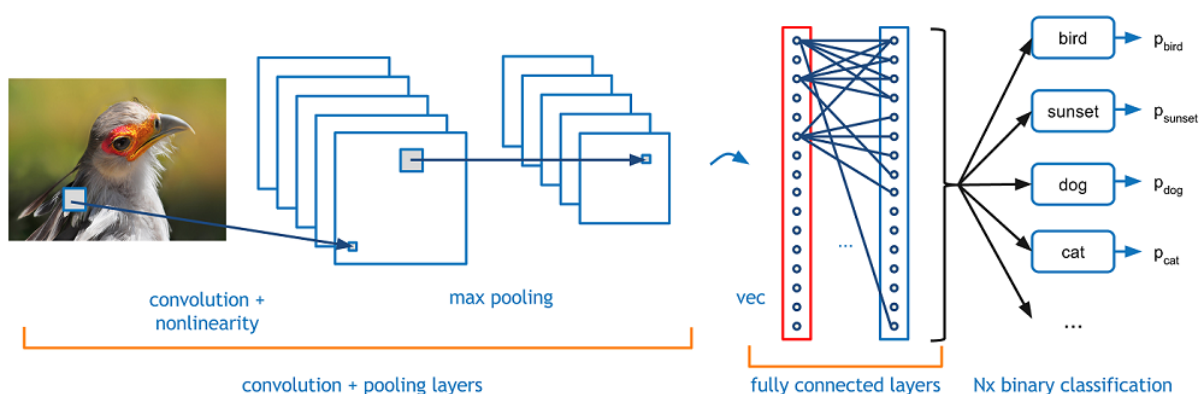


Figure 4. Convolutional Neural Network

Then we will be fining the model by make some of last layers of MobileNetv2 to be trainable and the last layers having the most parameters though. Then after 5 more epochs. We got **94%** training accuracy and **83%** test accuracy which is much better than before, and all of this takes small amount of time.



Using the main dataset which mentioned before in Dataset Section, which contains about 7800 images, 6000 of them to be trained. But this time converting to ResNet101 architecture, and applying the same procedure exactly as MobileNetv2, this time we got eight classes benign and malignant plus **masses of densities** as shown in Figure 8.

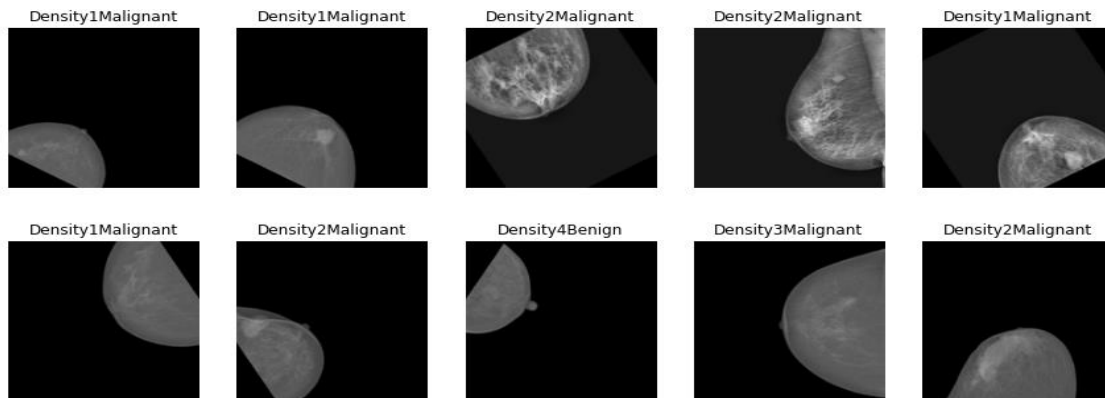


Figure 8. Breast Cancer Classes with masses (Authentic)

Dense breast tissue is detected on a mammogram [8], dense breasts indicates increasing in the chance that breast cancer may go **undetected by a mammogram**, since dense breast tissue can mask a potential cancer and increasing the risk of getting breast cancer. Therefore, even if the model indicated a benign for the patient but with high dense, so extra tests for the patient is mandatory to ensure that the benign class is benign and high dense of the breast not giving a false positive. We got an **overall accuracy** for the eight classes which is much harder as a model than above to classify:

$$\text{Training Acc} = 92\%, \quad \text{Test Acc} = 85\%$$

How can we evaluate our model, or what's the above test accuracy gives us actually? , what we can really do is to compare it with real life (compare apples to apples) to figure out the limitation of the model. Overall, the **sensitivity** of mammography is about **87 percent** [9]. This means mammography correctly identifies about 87 percent of women who truly have breast cancer in real life. So according to our 85 percent for 8 classes classification, we are doing great.

Figure 9, shows a simple code for developing random take of the validation set every time the code is running and evaluate the image described (It's Working well).

```
for imgs, labels in validation_dataset.take(1):
    plt.imshow(imgs[11].numpy().astype("uint8"))
    detect = list(labels[11].numpy())
    print('The truth ground class is ' + class_names[detect.index(max(detect))])
    img = tf.keras.preprocessing.image.img_to_array(imgs[11])
    img = np.expand_dims(img, axis = 0)
    predict = reconstructed_model.predict(img)
    predict = list(predict[0])
    print('The predicted class is ' + class_names[predict.index(max(predict))])
```

The truth ground class is Density3Malignant
The predicted class is Density3Malignant

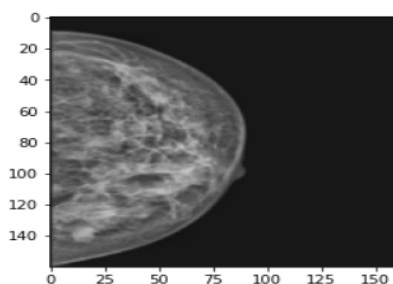


Figure 9. Evaluating Breast Cancer Multiclass Model (Authentic)

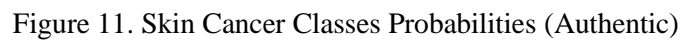
Training Acc = 98.6%, Test Acc = 96.7



Adam combines the best properties of the AdaGrad and RMSProp algorithms to provide an optimization algorithm that can doing well with noisy problems, after missing around here and there by different algorithms Adam is the choice for every model we trained already, and this might be the only property or hyperparameter won't change with models.

We are working on three classes for skin cancer to be classified by our model, one is malignant (**mel**) and the other two are benign (**nv** and **bkl**). But these data are not equal in terms of number of images or not even close as shown in Table 3 and Figure 11.

Number of Images for each Class	
Melanocytic Nevi (nv)	6705 Images
Melanoma (mel)	1113 Images
Benign lesions of the keratosis (bkl)	1099 Images



Using a little trick here to determine the const, is to get the accuracy of number of samples from validation test for each class. Without class weighting we got high accuracy for nv class and low accuracy for other two classes, then adding class weights to the TensorFlow model to fit and tune the const in the equation 1 to get **accuracy for each class** close as possible.



Image width to height ratio was 1.3333 (600,450) px originally, at first, we ignored that ratio and move with (160,160) px instead, but it might affect model learning somehow as skin cancer diagnosis is very sensitive to lesion diameter and dimensions. Therefore, we kept that ratio in other iteration (200,150) px which leads to higher rates of learning and slightly better accuracy with less epochs.

3.2.3 Colorectal Cancer Model

Figure 13 indicates eight random images with their label taking from the training dataset which is going to be trained for our model.



Using ResNet101 Model with categorical cross entropy for loss, and apply augmentation with flipping and rotations as well, we got an overall training accuracy about **98 percent**, and precision **98.5** and validation accuracy **89 percent** and precision **88.6** percent at first sight it might be indicated as overfitting with high bias. Nevertheless, we iterates different times not any epoch the validation accuracy exceeds **89 percent** even if training accuracy is low. So, it might be an indicating of the limitation of the model to be classified as better than this time.

4. DEPLOYMENT

The three final models created one for breast cancer, skin cancer, and colorectal cancer will be converted into TensorFlow lite model [13] to be deployed for low computation devices like mobile phones, so we can use function in model to predict the cancer type with just an image uploaded by camera or from gallery using Flutter Framework.

5. FURTHER WORK

There are still much more to do for this project to be mature:

- We must try other architectures other than ResNet101 and MobilNetv2 and comparing between them by computation time and accuracy.
- There's other trick for solving imbalanced data may work will, it's like oversampling without really duplicating images exactly, we can augment images until they catch the required number of images to solve imbalanced data.
- Increase the number of classifiers (Cancers) in the application and improves the UI of the mobile application.
- For later versions, we can take the uploaded image and classify it and added to the database with other trained images to improves the number of images to be trained. This might lead to mislabeled images but overall, it's going better.
-

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