Unsupervised Convolutional Filter Learning for COVID-19 Classification

ABSTRACT

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The outbreak of the SARS CoV-2, referred to as COVID-19, was initially reported in 2019 and has swiftly spread around the world. The identification of COVID-19 cases is one of the key factors to inhibit the spread of the virus. While there are multiple ways to diagnose COVID-19, these techniques are often expensive, time-consuming, or not readily available. Detection of COVID-19 using a radiological examination of Chest X-Rays provides a more viable, rapid, and efficient solution as it is easily available in most countries. The paper outlines a method that employs an unsupervised convolutional filter learning using Convolutional Autoencoder (CAE) followed by applying it to COVID-19 classification as a downstream task. This shows that the proposed technique provides state-of-the-art results with an average accuracy of 99.7%, AUC of 99.7%, specificity of 99.8%, sensitivity of 99.6%, and F1-score of 99.6%. We release the data and code for this work to aid further research.

1. INTRODUCTION

COVID-19 is one of the most highly infectious diseases in the last decade. As per the World Health Organization (WHO) report, as of July 22, 2021, over 192 million cases have been confirmed, with over 4.1 million deaths worldwide. This alarming rate of infection spread calls for critical and efficient diagnosis to curb the spread of the virus. Currently, the majority of the cases are being diagnosed using techniques such as RT-PCR, LFT, and Antibody Testing, which often produce highly accurate results. However, these techniques are generally time-consuming and expensive, posing a significant issue for developing and under-developed countries.

Chest X-Rays offer an alternate screening method for detecting COVID-19, where they are analyzed to look for visible evidence associated with SARS CoV-2 viral infection. The recent studies of the chest radiographic images show that viruses belonging to this family demonstrate substantial manifestation in radiographic images [1]. Thus, the categorization with the help of Chest X-rays could be a potentially cost-effective and accurate solution.

In our paper, we propose Convolutional Autoencoding as a pre-training technique for X-ray images. This model will learn to compress an image using an encoder, and again, decompress it to the original form using a decoder. Due to the unsupervised nature of this approach, the potential cost of labeling the images is alleviated. The resulting encoder will be capable of extracting complex features from an image, which will then be used as a pre-trained network, along with transfer learning to classify images. We also apply Gradual Unfreezing for the fine-tuning process as different layers in a network learn different features. For example, the lower layers of the CNN learn basic features such as edges and lines, while the higher layers learn more complex features such as textures and patterns. So, instead of refining all layers at once, which risks catastrophic interference, Gradual Unfreezing [2] suggests slowly un-freezing the model starting from the last layer as it contains the least general knowledge.

2. RELATED WORK

Recently, several AI systems based on deep learning have been implemented with promising results to detect COVID-19 using chest radiography images. Rahimzadeh and Attar [3] proposed a concatenated neural network of Xception and ResNet50V2 by using 180 COVID chest X-rays which obtained an accuracy of 99.56% and recall of 80.56%. A Generative Adversarial Network (GAN) approach was introduced by Loey et al. [4] using GoogleNet, AlexNet, and ResNet18 in low data settings with 69, 79, 79, 79 images for Covid, bacterial pneumonia, viral pneumonia, and normal cases respectively. This approach achieved test accuracy of 80.6% with GoogleNet, 85.2% with AlexNet, 99.9% with GoogleNet for 4-class, 3-class, and 2-class scenarios respectively. Ucar and Korkmaz [5] used SqueezeNet with Bayesian optimization with 76 COVID X-ray images and achieved 98.3% accuracy. Apostolopoulos and Mpesiana [6] implemented transfer learning with CNN using VGG19, Inception, MobileNet, Xception, and Inception-ResNetV2 and selected VGG19 as the final model with accuracy, sensitivity, specificity of 93.48%, 98.75%, and 92.85% respectively. Bandyopadhyay and Dutta [7] introduced an LSTM-GRU framework for confirming positive, negative, death and release cases of COVID with an accuracy of 87%, 67.8%, 62%, and 40.5% respectively.

Khan et al. [8] proposed CoroNet, a Deep CNN model, using Xception architecture with 284, 330, 327, 310 images for COVID, bacterial pneumonia, viral pneumonia, and normal cases respectively and achieved an accuracy of 85.9%,



a precision of 97% and a recall of 100%. Horry et al. [9] presented a COVID classification system using Xception, VGG, ResNet, and Inception and got an accuracy of 80%. Hemdan et al. [10] proposed COVIDX-Net, a framework of Deep Learning Classifiers, utilizing 7 Deep CNN architectures - VGG19, ResNetV2, InceptionV3, Inception-ResNetV2, MobileNetV2, DenseNet201, and Xception and achieved scores of 90% and 83% for accuracy and precision respectively. Singh et al. [11] performed COVID classification with 4 classifiers - Deep CNN, Extreme Learning Machine (ELM), Online sequential ELM, and Bagging Ensemble with Support Vector Machine (SVM). Bagging Ensemble with SVM is the best performing model with 95.7% accuracy and 95.8% precision. Ahuja et al. [12] proposed a 3 phasedetection model consisting of data augmentation phase, COVID detection phase, and abnormality localization phase to improve detection accuracy and achieved a test dataset accuracy of 99.4%. Fong et al. [13] conducted a case study that used composite Monte-Carlo (CMC) and fuzzy rule induction addressing the data limitation for forecasting. Islam et al. [14] proposed Deep CNN and CNN-LSTM models and achieved an accuracy of 99.4%, 99.9% AUC, 99.2% specificity, 99.3% sensitivity, and F1-score of 98.9% with the CNN-LSTM model.

3. RESEARCH METHOD

In this section, we present the proposed methodology for the identification of COVID-19 from chest X-Rays using an Unsupervised pre-training approach. Our proposed technique relies on a variant of Autoencoding called Convolutional Auto-encoder (CAE). An autoencoder has two main parts: an encoder that maps the input into the code and a decoder that maps the code to reconstruct the input. An autoencoder is a specific type of artificial neural network mainly employed to handle unsupervised machine learning tasks. In particular, an autoencoder is a feedforward neural network that is trained to predict the input itself. Thus, the system can minimize the reconstruction error by ensuring the hidden units capture the most appropriate features of the data.

Autoencoding is a data compression algorithm in which both compression and decompression are data-specific and automatically learned from samples instead of human engineering. This type of automatic or unsupervised learning is paramount to domains having less supervised data but huge unsupervised data. In our approach, we use Convolutional Neural Networks (CNNs or convnets) for compression and decompression as they have a proven record for images, and this variant is Convolutional Autoencoder (CAE). Figure 1 shows the block diagram of the Convolutional Auto-Encoder consisting of an Encoder and Decoder model.

The encoder tries to compress the input image by extracting important features such that the decoder can recreate the original image with minimal loss. This acts as a pre-training objective that allows the encoder to extract important features.

After training this encoder-decoder model on the training data, the encoder model is saved, and the decoder model is discarded. This encoder can then be used as a data preparation technique to perform feature extraction on the raw data, which in turn can be used to train a different machine learning model for downstream tasks.



Figure 1. Block diagram of convolutional auto-encoder model

The proposed technique achieves this in a two-step process: (1) Unsupervised Pre-training of Convolutional Auto-

Encoder.
(2) Transfer Learning with Encoder for downstream classification task – Detection of Chest X-Rays affected with COVID-19.

3.1 Unsupervised pre-training of convolutional autoencoder

Dataset: We collected 18617 chest X-ray images from various sources – covid19-radiography-database [15], covid19-chest-xray-image-dataset [16], chest-xray-pneumonia [17], covid-chestxray-dataset [18], Figure 1-COVID-chestxray-dataset [19] and Actualmed-COVID-chestxray- dataset [20].

Modeling:

The Encoder (E) takes the input image 'x', and compresses it into lower-dimensional features 's' known as 'bottleneck':

$$\mathbf{s} = \mathbf{E}(\mathbf{x}) \tag{1}$$

Layer (type)	Input Shape	Kernel #	Kernel Size	Output Shape	Param #		
Conv2D	224 x 224 x 3	64	3 x 3	224 x 224 x 64	1792		
Conv2D	224 x 224 x 64	64	3 x 3	224 x 224 x 64	36928		
MaxPooling2D	224 x 224 x 64	-	2 x 2	112 x 112 x 64	0		
Conv2D	112 x 112 x 64	128	3 x 3	112 x 112 x 128	73856		
MaxPooling2D	112 x 112 x 128	-	2 x 2	56 x 56 x 128	0		
Conv2D	56 x 56 x 128	256	3 x 3	56 x 56 x 256	295168		
MaxPooling2D	56 x 56 x 256	-	2 x 2	28 x 28 x 256	0		
Conv2D	28 x 28 x 256	512	3 x 3	28 x 28 x 512	1180160		
MaxPooling2D	28 x 28 x 512	-	2 x 2	14 x 14 x 512	0		
Conv2D	14 x 14 x 512	512	3 x 3	14 x 14 x 512	2359808		
MaxPooling2D	14 x 14 x 512	-	2 x 2	7 x 7 x 512	0		
Total params: 3,947,712							
Trainable params: 3,947,712							
Non-trainable params: 0							

Table 1. Architecture of encoder block

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Layer (type)	Input Shape	Kernel #	Kernel Size	Output Shape	Param #		
Conv2DTranspose	7 x 7 x 512	512	3 x 3	14 x 14 x 512	2359808		
Conv2DTranspose	14 x 14 x 512	256	3 x 3	28 x 28 x 256	1179904		
Conv2DTranspose	28 x 28 x 256	128	3 x 3	56 x 56 x 128	295040		
Conv2DTranspose	56 x 56 x 128	64	3 x 3	112 x 112 x 64	73792		
Conv2DTranspose	112 x 112 x 64	3	3 x 3	224 x 224 x 3	1731		
Total params: 3,910,275							
Trainable params: 3,910,275							
Non-trainable params: 0							

Even though most X-ray images are of single channel, it is possible for the images to have markings or coloration to highlight specific portions. Training the encoder layers with 3channel images instead of single-channel images allows us to apply the pretrained encoder model for various downstream Xray applications. Hence, we resize the input X-ray images to (224x224x3) and rescale all the pixel values in the range 0-1 for normalization. This ensures that each input has a similar data distribution and also improves computational efficiency. This preprocessed image is then fed to the Encoder (E). We design the Encoder as a 6-layer 2D Convolutional Network with Rectified Linear Unit (ReLU) as activation function, kernel size of (3x3), and Max-Pooling with a pool size of (2x2). This 6-layer Convolutional Encoder Network outputs (7x7x512) extracted features as shown in Table 1.

The Decoder (D) accepts the lower dimensional extracted features as inputs and reconstructs the original image with shape (224x224x3). Our Decoder as shown in Table 2 is a 5-layer Transposed 2D Convolutional Network with a stride of 2, ReLU activation function, and a kernel size of (3x3). This kernel size allows the model to learn complex features with less computation.

If we denote reconstructed image as 'y', then,

$$y = D(s) \tag{2}$$

From Eq. (1) and Eq. (2), the output image 'y' can be denoted as:

$$y = D(E(x)) \tag{3}$$

The reconstructed image 'y' is then compared with input 'x' and loss is calculated which is used to update the weights of all the layers. We used 'Mean Squared Error' (MSE) as the loss function. The MSE represents the cumulative squared error between the compressed and the original image, and lower the value of MSE, the lower the error. The Mean Squared Error can be calculated using the following equation:

$$MSE = \frac{\sum_{M,N} [I_1(m,n) - I_2(m,n)]^2}{M * N}$$
(4)

where, I_1 and I_2 are the input and output images with dimensions (m, n) respectively. 'Adam' is used as the optimizer to update weights with an initial learning rate of 0.01.

3.2 Transfer learning with encoder for downstream classification task – Detection of Chest X-Rays affected with COVID-19

Dataset: We collected 3709 COVID-19 chest X-rays from various sources - covid19-radiography-database [15], covid19-chest-xray-image-dataset [16], Figure1-COVID-chestxray-dataset [19] and Actualmed-COVID-chestxray-dataset [20]. Then we collected 3700 images for both normal and pneumonia cases from 'covid19-radiography-database [15] and chest-xray-pneumonia [17] respectively. Similar to the data preparation step during pre-training, we resized all images to (224,224,3) and rescaled each pixel to be in 0-1 range as a preprocessing step.

Layer (type)	Input Shape	Kernel #	Kernel Size	Output Shape	Param #	
Conv2D	224 x 224 x 3	64	3 x 3	224 x 224 x 64	1792	
Conv2D	224 x 224 x 64	64	3 x 3	224 x 224 x 64	36928	
MaxPooling2D	224 x 224 x 64	-	2 x 2	112 x 112 x 64	0	
Conv2D	112 x 112 x 64	128	3 x 3	112 x 112 x 128	73856	
MaxPooling2D	112 x 112 x 128	-	2 x 2	56 x 56 x 128	0	
Conv2D	56 x 56 x 128	256	3 x 3	56 x 56 x 256	295168	
MaxPooling2D	56 x 56 x 256	-	2 x 2	28 x 28 x 256	0	
Conv2D	28 x 28 x 256	512	3 x 3	28 x 28 x 512	1180160	
MaxPooling2D	28 x 28 x 512	-	2 x 2	14 x 14 x 512	0	
Conv2D	14 x 14 x 512	512	3 x 3	14 x 14 x 512	2359808	
MaxPooling2D	14 x 14 x 512	-	2 x 2	7 x 7 x 512	0	
Reshape	7 x 7 x 512	-	-	49 x 512	0	
LSTM	49 x 512	-	-	49 x 512	2099200	
Flatten	49 x 512	-	-	25088	0	
Dense	25088	-	-	64	1605696	
Dense (output)	64	-	-	3	195	
Total params: 7,652,803						
Trainable params: 3,705,091						
Non-trainable params: 3,947,712						

Table 3. Architecture of proposed classification network

Modelling: In this section, we explain the process of transfer learning with the pre-trained encoder for COVID-19 classification. The encoder model from the previous step is attached to a classification head for downstream classification. Here, we chose LSTM and Dense Layers that result in a network that is very efficient for classification tasks. Figure 2 shows the block diagram of the classification network with output shapes of each layer. In Figure 2, the input image is a test image fed to the pre-trained encoder obtained from the trained CAE model. This encoder extracts the important features from the image and outputs a compressed image of size (7x7x512). Table 3 shows the detailed architecture along with the total trainable parameters.



Figure 2. Block diagram of proposed convolutional network

This classification model is fine-tuned by applying the 'Gradual Unfreezing' technique. As the weights of the classification head (LSTM & Dense) are randomly initialized, we first freeze all the encoder layers, so that the pre-trained weights are not affected. Once the classification head is trained, we unfreeze the encoder layers and train the entire model. This technique helps the model to learn layer-level features effectively as the lower layers learn simple representations such as edges and curves while the higher layers learn complex representations.

4. RESULTS

In this experiment, the Chest X-Ray dataset was split into 73-13.5-13.5 sets for training, validation, and testing respectively. The proposed architecture is a combination of 6 Convolutional layers, an LSTM, and 2 Dense layers as shown in Table 3, trained using a single NVIDIA Tesla T4 with 16GB memory. We used 'Adam' optimizer with a learning rate of 0.001, categorical cross-entropy loss, early stopping with patience of 5, and batch size of 16. The total fine-tuning process took only 25 minutes and 18 epochs to converge because of the efficient pre-training method.

We experimented with various architectures for the classification head and Table 4 shows the performance comparison of these models. Although the proposed model is the best performing, it is worth acknowledging the competitiveness of these architectures because of the robustness of the pre-trained encoder.

Figure 3 shows the confusion matrix of the proposed method. Figure 4 shows the graphical representation of the performance metrics. Most of the existing systems rely on models that were pretrained on image datasets not specific to X-ray images. Hence, these models rely entirely on the fine-

tuning phase to learn about the features present in X-rays. Whereas the proposed CAE architecture is pretrained on a large corpus of X-ray images, allowing the model to extract intricate features useful for a wide range of X-ray related downstream tasks. Table 5 corroborates the effectiveness of the proposed system by comparing it with contemporary architectures.



Figure 3. Confusion matrix of proposed model



Figure 4. Representation of performance metrics

Table 4. Performance comparison of classification heads

Classification Head	Accuracy (%)	Precision (%)	Recall (%)	F1- score (%)	AUC (%)
3 LSTM + 2 Dense	97.5	96.3	96.3	96.3	97.2
1 LSTM + 3 Dense	97.7	96.6	96.6	96.6	97.4
5 Dense	98.4	97.5	97.6	97.55	98.2
1 LSTM + 2 Dense (Proposed)	99.7	99.6	99.6	99.6	99.7

Table 5. Comparison of proposed system with existing systems

Author	Architecture	Training Sample (COVID-19)	Testing Sample (COVID-19)	SAccuracy (%)	Accuracy COVID-19) (%)
Rahimzadeh et al.	ResNet50V2 + Xception	180	31	91.4	99.6
Loey et al.	GoogleNet	69	9	80.6	100.0
Ucar et al.	COVIDiagnosis-Net	76	10	98.3	100.0
Khan et al.	CoroNet	284	29	89.5	96.6
Hemdan et al	COVIDX-Net	25	5	90.0	-
Islam et al.	CNN-LSTM	1220	305	99.4	99.2
Proposed System	CAE	2703	503	99.73	99.6

5. CONCLUSION

In this paper, we suggest an unsupervised pre-training technique for diagnosing COVID-19 from chest X-rays. The proposed technique utilizes a combination of Convolutional Neural Network (CNN) which can be used as a pre-trained network to extract features for any chest X-ray tasks and an LSTM layer and Dense layer as classification head. According to the experiment results, the proposed method achieved an accuracy of 99.73%, a precision of 99.6%, a sensitivity of 99.6%, and an AUC of 99.7% with an F1-score of 99.6%. We hope that the proposed system would be able to help patients and reduce the workload of the medical diagnosis of COVID-19. Finally, the performance of our proposed system was not compared with radiologists and that would be part of a future study.

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