

Lung cancer detection using novel residual unity AlexNet-based optimized mish dropout-deep convolutional neural network

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Abstract

Globally, one among the most prominent cause of death is lung cancer, which is the most malignant tumorin human health. Hence, automatically detecting or diagnosing lung disease as of computerized tomographic scan image is the necessary application. Numerous cancer classification systems have been engendered for this. However, it is not easy to determine the presence of tumors in small nodules. Hence, a novel Aquila-optimized mish dropout-deep convolutional neural network (AmiD-DCNN) cancer classification system was proposed in this paper. The noises are eliminated by utilizing an adaptive median filter initially and the Chi-square distribution adapted contrast limited adaptive histogram equalization is utilized to elevate the contrast. The residual unity AlexNet is utilized to segment the lung regions from the preprocessed image; also the Jaccard similarity and quadratic kernel-induced profuse clustering are employed to extract the cancerous region. The features are extracted after those steps, which are then fed to the Amid-DCNN classifier to classify cancer. The experiments are evaluated along with analogized with the benchmark models. The proposed model's efficient performance was demonstrated by the experimental outcomes.

Keywords Adaptive median filter (AMF) \cdot Chi-square distribution adapted contrast limited adaptive histogram equalization (Chi-CLAHE) algorithm \cdot Residual unity AlexNet (RU-AlexNet) \cdot Jaccard similarity and quadratic kernel-induced profuse clustering (JQPC) \cdot Aquila-optimized mish dropout-deep convolutional neural network (AmiD-DCNN)

1 Introduction

With 8.2 million deaths per year or so, cancer is the leading cause of mortality globally. However, the top of this list with 1.69 million fatalities annually is the LC (Perez and Arbelaez 2020). LC is a lung disease condition and can be either primary (originating as of the lungs) or metastasis (originating as of other organs). Small cell lung cancer (SCLC) and non-small cell lung cancer (NSCLC) are the extensions of LC. The most malignant cancer that occurs in 15% of cases is the SCLC, whereas NSCLC occurs in the remaining 85%. Adenocarcinoma, large-cell carcinoma,

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and squamous-cell carcinoma are the subcategories of NSCLC. Since the true cause of cancer and its whole therapy have not been discovered, lung cancer detection (LCD) continues to be a challenge for medical practitioners. However, cancer that is detected early enough can be treated (Akter et al. 2021) and survival rates can be elevated.

This is why accurate classification of lung illness is necessary, whereas LCDs can be created using medical imaging techniques as chest X-rays, MRI scans, computerized tomography (CT) scans, etc. Since CT scans create cross-sectional body images using CT and X-rays, which are crucial for the medical condition's diagnosis it is more appropriate to utilize CT scans for human body organs like the lungs (Asuntha and Srinivasan 2020). Manually segmenting the LC such as lesions, which calls for the skill of a radiologist drawing, is the conventional method for identifying LC in scan images. For a variety of reasons, it is difficult to engender strong and effective outcomes (Avanzo et al. 2020). Additionally, the sort, shape, size,

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and location of lung abnormalities may vary from what lung specialists predict. Thus, a computer-aided design module (CAD) was engendered to deal with morphological operators to execute the pre-processing over lung nodules (Firdaus Abdullah et al. 2020). For the early detection and classification of LC in the CT scan images, several machine learning (ML) and deep learning (DL) algorithms were developed.

Graph convolution networks (GCN), artificial neural networks (ANN), deep neural networks (DNN), support vector machines (SVM), convolutional neural networks (CNN) or deep CNN (CNN), and hybrid DNN with optimization algorithms are a few ML techniques utilized for the LC categorization (Firdaus et al. 2020). Since DL networks, also known as DCNN, accept images as input and can be trained end-to-end utilizing a supervised methodology, while learning highly discriminative image features, it is picked as the accurate lung cancer classifier among these models (Gao et al. 2021). Owing to learning numerous features of the input CT scan images, the DCNNs, however, have a temporal complexity. Hence, a novel AmiD-DCNN-based LC classification framework is proposed to tackle this issue.

1.1 Problem statement

Certain limitations are still needed to be solved for better classification even though several ML- and DL-based models are available for LC classification. A few limitations of the prevailing models are framed as follows,

- In the prevailing methodologies, grading the images grounded on the degree of the pulmonary nodule was not evaluated, which is of great significance for the LC diagnosis and treatment in clinical applications.
- Numerous irregular details along with low-quality pixels that diminished the LC prediction accuracy are contained in the captured CT images. The classification process' accuracy was significantly influenced by this.

By analyzing these downsides, the proposed scheme's objective is to engender an accurate classifier with a reliable segmentation and patch extraction technique.

This paper is structured as follows; Sect. 2 describes the recent related works of LC classification. Section 3 explains the proposed methodology. Section 4 analyzes the proposed model's experimental outcomes. Finally, Sect. 5 concludes this paper and gives a future suggestion.

2 Related works

Investigated LCD model grounded on morphological feature extraction (FE) and Kernel-based non-Gaussian-CNN (KNG-CNN) classification of CT images (Jena and George 2020). The false positives in the work were diagnosed by utilizing the KNG computation. Consequently, the KNG-CNN technique obtained an accuracy of 87.3%. But, the input image quality was not pondered, which would affect the classifier's accuracy.

Developed a DL-assisted prediction of LC on CT images grounded on the adaptive hierarchical heuristic mathematical model (AHHMM) (Karthiga and Rekha 2020). Image acquisition, pre-processing, binarization, thresholding and segmentation, FE, and detection by DNN were various stages comprised in this system. The test evaluation exhibited that the presence or absence of LC was detected by the AHHMM with 96.67% accuracy. However, the accuracy was analyzed only for ten images. Hence, when executed with enormous images, the accuracy could deviate.

K-nearest Neighbor (KNN) method for the LC prognosis. To elevate the KNN algorithm's accuracy, the features were chosen in the presented scheme by the genetic algorithm (GA) (Kavithaa et al. 2021). The presented approaches' implementation on an LC database exposed that better accuracy was attained by the suggested model. However, the GA could not achieve the repetitive fitness evaluation for the complex data. Hence, the k value was chosen accurately.

An approach grounded on the improved Naive Bayes classifier (I-NBC) algorithm for the early LC prediction on the CT images. An accelerated wrapper-based binary artificial bee colony (AWB-ABC) algorithm was utilized in this model for effective feature selection (Kumbhar et al. 2022). A superior trade-off betwixt, the prevailing and the engendered techniques were exhibited by the experiential outcomes. However, the linear data that limits the model's performance was not classified by the INBC model.

DL framework for the detection of lung nodules on CT images (Majidpourkhoei et al. 2021). The features as of the lung images were automatically extracted by engendering the CNN layers centered on the LeNet-5 model succeeded by the suspicious regions' classification as either nodule or non-nodule objects. The experiential outcomes demonstrated that a better accuracy than the prevailing works was attained by the developed CNN model. However, an overfitting issue was caused by the LeNet-5 for a few images that could affect the outcomes.

An autonomous decision support scheme for LC detection and classification. A multidimensional region-based fully convolutional network (mRFCN) algorithm was developed in this model for that (Maleki et al. 2021). When contrasted to the benchmark nodule detection/classification models, better detection performance was exhibited by the experiential outcomes. However, more false positives were provided by the model on the clinical dataset than the compared model.

A CNN-based framework for the LC diagnosis. An input image was classified by the developed CNN into one of three classes (benign, malignant, and non-cancerous) (Marentakis et al. 2021). A classification accuracy of 93.9% with diminished false positives was achieved by the suggested CNN. Yet, better sensitivity than the compared model was not achieved.

Improved DNN (IDNN) and ensemble classifier for LC identification from the CT image automatically. The noise was diminished with a multilevel brightness-preserving approach from the CT image. The affected region was segmented with IDNN (Masood et al. 2020). Regarding logarithmic loss, the discussed model elevated the LC prediction rate. Nevertheless, predicting the ensemble classifier's output was hard, which could cause computational complexity. Deep Gaussian mixture model in a region-centered CNN (DGMM-RBCNN) may have reduced computing cost, but it was difficult to anticipate the ensemble classifier's output (Nageswaran et al. 2022). For the elimination of over-parameterized solutions in the RBCNN model, DGMM-based dimensionality reduction was introduced in the given model. In the experiential analysis, better accuracy was achieved by the DGMM-RBCNN model than the prevailing system. In the initial stages, the model's precision was poor, which could diminish the average precision that could affect the model's performance.

Elastic deformation (ED)-based ML to augment the subtype classification of NSCLC (Nanglia et al. 2021). For the output prediction, '2' classifier models were trained on the original and augmented dataset. Regarding accuracy, sensitivity, specificity, and f1-score, the developed model's performance was achieved. The model was made by processing more features which takes more time to train the parameters that could create time complexity.

Methodology for LC detection with enhanced segmentation accuracy. Here, the segmented data was classified by a neuro-fuzzy classifier (Perez and Arbelaez 2020). The suggested model concluded with 90% accuracy with a diminished false alarm rate. As the model analyzed outcomes only for a limited number of images, the outcomes might deviate when processed with more images.

DL-based approach for LC detection and classification. The optimal features were selected by the fuzzy particle swarm optimization (FPSO) algorithm (Raoof et al. 2020). It was verified as of the experimental outcomes that better performance than the other techniques was attained by the developed technique. Yet, more accurate outcomes than the surveyed works were not produced by the work. A scheme of LC histology classification as of CT images grounded on radiomics and DL models. '4' diverse families of techniques were investigated for that (Rehman et al. 2021). The outcomes exhibited that Inception and LSTM + Inception was the best CNN that generated better performance than the other models. However, more time was taken by the model to evaluate all models that caused memory and time complexity.

A DL-centered methodology for LCD as of denoised CT scan images. "Denoising first" two-path CNN called denoising first detection-Nnetwork (DFD-Net) for LC detection was the introduced DL model (Sori et al. 2021). Regarding accuracy, sensitivity, and specificity, better LCD outcomes than the recently introduced approach were revealed by the DFD-Net. However, there was a loss in a few significant features that arouses the overfitting issue.

An early LCD model for breast image segmentation and analysis is discussed on (Surendar 2021). For lung lobe segmentation, global thresholding and guided 3D watershed transform were the models employed. It can be claimed as of the obtained outcomes that more accuracy was attained by the suggested model. But, a better segmentation cannot be executed for noisy images that degraded the model.

3 Proposed lung cancer classification system

For the efficient identification and classification of LC, a novel AmiD-DCNN classifier is future herein paper. The lung region is originally segmented herein scheme. The landscapes are extracted as of that segmented regions and fed into the AmiD-DCNN, which classifies cancer. Figure 1 articulated the future copy's block drawing.

3.1 Preprocessing

The input chest CT image (I) was used by the proposed work for categorizing the LCs that had been gathered from publically available datasets. The input image is induced into the preprocessing step owing to the input CT image's lower quality. Noise removal and contrast enhancement are the two steps in preprocessing.

3.1.1 Noise removal

Utilizing an adaptive median filter (AMF), the noise is eliminated from the input image. The drawbacks faced by the standard median were eliminated by designing AMF. The filter performs under two steps.

Step 1: After choosing the input images' window size, the median value (G_{med}) for the image is computed.

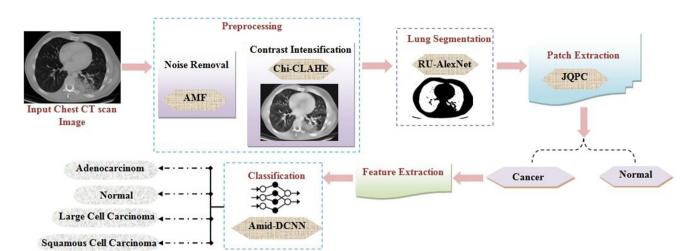


Fig. 1 Block drawing of the future practice

$$L_{\rm lev\,1}^1 = G_{\rm med} - G_{\rm min} \tag{1}$$

$$L_{\rm lev\,1}^2 = G_{\rm med} - G_{\rm max} \tag{2}$$

where $G_{\min} \in I$ is the minimum gray level value, $G_{\max} \in I$ is the maximum gray level value in the selected window W, the parameters of gray level value in step 1 is notated as $L_{\text{lev 1}}^1$ and $L_{\text{lev 1}}^2$.

$$\begin{cases} \text{if } L_{\text{lev }1}^{1} > 0 \text{ and } L_{\text{lev }1}^{2} < 0, \text{ go to step 2} \\ \text{else,} \text{ increase window size} \end{cases}$$
(3)

$$\begin{array}{l} \text{if } W \leq W_{\text{increased}}, \text{ then repeat step 1} \\ \text{otherwise, } \text{old grey level}\left(G_{\text{old}}\right) \end{array}$$

$$\tag{4}$$

Here, the elevated window size is signified as $W_{increased}$. Step 2: Here, it checks whether the noise corrupted the current pixel value in the input image or not. The pixel changes with the median if the pixel is corrupted or else retains the grayscale's pixel value.

$$l_{\text{lev 2}}^1 = G_{\text{old}} - G_{\min} \tag{5}$$

$$l_{lev2}^2 = G_{old} - G_{\max} \tag{6}$$

$$\begin{cases} \text{if } l_{\text{lev}2}^1 > 0 \text{ and } l_{\text{lev}2}^2 < 0, \text{ output}G_{\text{old}} \\ \text{otherwise, } G_{\text{med}} \end{cases}$$
(7)

where the parameters of the gray level value in step 2 are signified as $l_{\text{lev}2}^1$ and $l_{\text{lev}2}^2$. The output from these two processes can be obtained as I_{removed} .

3.1.2 Contrast intensification

The Chi-square distribution adapted contrast limited adaptive histogram equalization (Chi-CLAHE) algorithm was utilized to elevate the filtered CT lung image I_{removed} in this section. To enhance the image's contrast, a variant of

the adaptive histogram equalization technique applied over all neighborhood pixels is the CLAHE. The control parameters to adjust the contrast are chosen randomly in the conventional CLAHE algorithm. This may impact the contrast enhancement performance as well as elevate the number of iterations. Therefore to compute the control parameter, the Chi-square distribution function was utilized in this proposed work to overcome this drawback. The control parameters in a single test were computed by the distribution without elevating the error probability.

The filtered region is initially partitioned into contextual regions with an equal number of pixels. They are named tiles. For each tile, the pixels present in the image compute the histogram. Then, the average pixel in the gray level G_{avg} is computed as,

$$G_{\rm avg} \in \mathbf{I} = \frac{P_s - P_t}{N_g} \tag{8}$$

where the number of pixels in *s* direction of the contextual region is notated as P_s , the number of pixels in *t* direction of the contextual region is represented as P_t , and the number of gray levels notated as N_g . Using Chi-square distribution, the clip limit $cl_{chi sq}$ is set grounded on the average gray level and it is articulated as,

$$cl_{\text{chi sq}} = \frac{1}{(R/2)2^{R/2}} \left(G_{\text{avg}}\right)^{R/2-1} e^{G_{\text{avg}}/2}$$
(9)

Here, a random number is denoted by R. Then, the pixels over the control parameter are pondered as excess pixels and are redistributed for each gray level. To remove induced boundaries in the input image, it merges all the neighboring tiles using bilinear interpolation after executing the equalization. Hence, I_{pre} denotes the image after the contrast enhancement.

3.2 Lung segmentation

Here, the residual unity AlexNet (RU-AlexNet) is utilized to segment the lung image from $I_{\rm pre}$. A graphics processing unit (GPU) to accelerate image classification was utilized by the AlexNet, which is a DCNN model. However, more time was required by AlexNet to segment the lung region. This may lead to some sort of error. The proposed work adapted residual network along with unity normalization function to tackle these issues. The residual module utilizing a method of the fitting residual map was the most vital one in the residual network. The convolutional layer's outcome was not provided directly however chooses the residual mapping. Moreover, unity normalization is utilized to normalize the image pixels. The proposed segmentation algorithm is called RU-AlexNet owing to the addition of residual network (R) and unity normalization (U).

Here, the preprocessed lung image is provided as input to the RU-AlexNet. Also, the kernels are utilized to convolute the image and are articulated as,

$$\omega_n = \{\omega_1, \omega_2, \omega_3, \dots, \omega_N\} \tag{10}$$

where the number of kernels is denoted as ω_n . In the mathematical representation, the convolution for the given input data *C* is expressed as,

$$C = I_{\rm pre} \cdot \omega \tag{11}$$

Unity normalization (U) that provides nonlinearity in the neural network modifies the network structure. The gradient is also enhanced by the function. The normalization can be defined as,

$$\operatorname{Re} Lu(C) = \max(C, 0) \tag{12}$$

Further, the convoluted features are provided in the residual network containing a number of residual blocks. Every residual block is followed by a unit normalization layer, convolution, and a ReLU activation function. The input is added directly to the activation function (skip connection) in each residual block.

The first residual block is directly inputted by the convolution's output. Convolution and unit normalization is executed as stated earlier. The output is injected directly into the ReLU activation function. It may aid to study the complex patterns in the data and the residual block's output. It can be articulated as,

$$h(\Re) = f(C) \tag{13}$$

Now, the output with the introduction of skip connection is changed to,

$$h(C) = f(C) + C \tag{14}$$

Till the last residual block, this process is repeated. The residual network's output is transferred into the max pooling operation that diminished the number of features. The pooling operation \Im can be formulated as,

$$\Im = \frac{C - \omega}{\omega_{\text{stride}}} + 1 \tag{15}$$

Anywhere, the kernel's steps are written as ω_{stride} . Till the last layer, this process is continued.

All the features are flattened and provided to the completely linked coating (CLC) after the combining operation. A similar number of production lumps is limited in the last CLC identical to the amount of lessons. To board lesson likelihoods, the softmax activation purpose was utilized by the RU-AlexNet to normalize production actual standards in the series [0, 1] from the last CLC. The softmax equation *S* is signified as follows,

$$S = \frac{e^{\Im_k}}{\sum\limits_{k=1}^b \Im_k} \tag{16}$$

Anywhere, the CLC production at *k*th lump is signified for example \Im_k , *b* is the entire amount of production lumps. The segmented lung can be obtained from this process, and it is denoted as I_{seg} .

3.3 Patch extraction

Here, the Jaccard similarity and quadratic kernel-induced profuse clustering (JQPC) is utilized to extract the abnormal regions (cancerous region) from I_{seg} . As per the pixel similarity, the lung region was inspected by the traditional profuse clustering (PC). For predicting the affected region, the images are segmented into several subimages by the PC. Here, the Jaccard similarity index executed the grouping of similar superpixels into the same group. Also, the *k*-means have the issue of distance computation in PC during the clustering process. The quadratic kernel function is applied to overcome this. The extraction occurred in the following steps.

Step 1: Take the lung segmented region I_{seg} .

Step 2: Simple Linear Iterative Clustering (SLIC) is utilized to compute superpixels. To diminish the operational complexity, similar value pixels are grouped utilizing this algorithm.

Step 3: The K means algorithm is utilized to cluster all the pixels present in the image. The centroid is chosen randomly for this purpose. Then, the distance $\text{Dis}_{q \text{ker}}$ betwixt the centroids \cup_k and the selected pixel p_i is computed as, $\text{Dis}_{q \text{ker}} = \varphi(\cup_k), \varphi(p_i \in \mathbf{I}_{\text{seg}})$

$$\varepsilon_n = \{\varepsilon_1, \varepsilon_2, \varepsilon_3, \dots, \varepsilon_N\}$$
(18)

Here, the kernel function is represented as ϕ .

Step 4: A cluster is chosen for pixels where the distance betwixt the pixels, and the centroid is minimal.

Step 5: Fuse superpixels.

Step 6: Till the entire superpixels end, the process is repeated. The output can be obtained as normal β and abnormal regions α as of this process. For further processing, this abnormal region is utilized. The proposed JQPC's pseudocode is,

Anywhere, the numbers of removed structures are signified for example
$$\varepsilon_n$$
.

3.5 Classification

(17)

For labeling the lung stages, the AmiD-DCNN further examined the extracted features ε_n . To identify patterns in images and video, DCNN is the most commonly utilized type. However, owing to the elevated number of layers

Begin				
	Initialize parameters p_i, \bigcup_k , iteration <i>i</i> , maximum iteration i_{max}			
	Compute superpixels using SLIC			
	Perform clustering			
Select the number of centroid				
	Set $i = 1$			
	For $i \leq i_{\max}$			
	For each pixel p_i , do			
	Compute distance <i>Dis</i> _{<i>q</i>ker}			
	$Dis_{q \ker} = \varphi(\cup_k), \varphi(p_i \in \mathbf{I}_{seg})$			
	End for			
	Check all the pixels are presented under the cluster			
	If $(p_i == undercluster)$ {			
	Stop criteria			
	} Else {			
	Set $i = i + 1$			
	}End if			
	End for			
	Fuse superpixels			
	Return α			

3.4 Feature extraction

Here, the input acquired is the cancerous region α and passed on to the FE stage. Intensity histogram, histogram of oriented gradients, Gabor filter, entropy filter, grayscale contrast, grayscale correlation, grayscale energy, grayscale homogeneity, standard deviation, Haarwavelet, etc., are the several spectral features derived by the FE stage. The extracted features are articulated as,

betwixt input and output layers along with the random weights engendered at each layer due to active neurons, the traditional DCNN has a computational complexity issue. By replacing the ReLu activation with the mish activation function, a dropout connection is added to the DCNN network to execute effectively by overwhelming such limitations. The adaptive learning strategy-centered Aquila optimizer (ALSAQ) Algorithm was utilized to optimize the weight values in the DCNN. The proposed AmiD-DCNN's architecture is exhibited in Fig. 2,

Weight initialization: The adaptive learning strategybased Aquila optimizer (ALSAQ) algorithm was utilized to compute the proposed model's weight. Generally, a new population-centered optimizer is the Aquila optimizer (AQ), which is classified as a metaheuristic optimization technique. However, the AQ may fall readily into the local optimal solution. A very low convergence rate was found. For distributing Aquila's position in the search space, an adaptive learning model is introduced in the proposed model to elevate the convergence speed along with prevent the solution from local optima.

Initially, a set of random numbers (number of Aquila) are initialized for this. It can be articulated as,

$$\aleph_n = \{\aleph_1, \aleph_2, \aleph_3, \dots, \aleph_N\}$$
(19)

The novel algorithm was utilized to update the weights utilizing these initialized numbers. Utilizing adaptive learning strategy, each Aquila updated its position in betwixt an upper limit (UL) and a lower limit (LL). It is expressed as,

$$\aleph_{pq} = R * (UL_q - LL_q) + LL \quad \text{where } p = 1, 2,m \text{ and } q$$

= 1, 2, 3.....d (20)

where the total number of population is notated as m, and each Aquila's dimension size is signified as d, \aleph_{pq} represents pth Aquila at pth dimension, the random value is specified as R, UL_q and LL_q represents upper and lower bound at dimension q.

To determine the position of prey, the area of the search space from the sky was explored by the Aquila. The prey areas are identified by the Aquila, which chooses the best areas for hunting. The exploration is computed as,

$$\aleph_n^{\text{exploration}}(\tau+1) = \aleph_{\text{best}}(\tau) \left(1 - \frac{\tau}{t}\right) + \left(\aleph_{\text{avg}}(\tau) - \aleph_{\text{best}}(\tau) * R\right)$$
(21)

$$\aleph_{\text{avg}}(\tau) = \frac{1}{m} \sum_{p=1}^{m} \aleph_p(\tau)$$
(22)

where the solution of the next iteration of τ is represented as $\aleph_n^{\text{exploration}}(\tau + 1)$, the best-obtained solution until τ the iteration is signified as $\aleph_{\text{best}}(\tau)$, which describes the prey's exact position, the current solutions' average value at τ th iteration is notated as $\aleph_{\text{avg}}(\tau)$, the parameter to control the search process are signified as $(1 - \frac{\tau}{t})$, $\aleph_m(\tau)$ is the *p*th Aquila at τ th iteration, the total amount of repetitions is meant for example *t*.

Subsequently, the area of prey was chosen by the Aquila, which is found at a high level of altitude. The Aquila circle in the clouds, get into position along with prepares to attack the prey. The step $\aleph_n^{\text{prepare}}$ is expressed mathematically as,

$$\aleph_n^{\text{prepare}} = \aleph_{\text{best}}(\tau) * L(d) + \aleph_r(\tau) + (s_1 - s_2) * R$$
(23)

where the random Aquila between [1 - m] is symbolized as $\aleph_r(\tau)$, s_1 and s_2 are the spiral shape in search space, the distribution function of levy flights at dimensionis notated as L(d). The distribution function is articulated as,

$$L(d) = f \times \frac{R_1 \cdot \ell}{|R_2|^{\frac{1}{h}}}$$

$$\tag{24}$$

$$\ell = \frac{(1+\hbar) \cdot \sin e \frac{\pi \hbar}{2}}{\left(\frac{1+\hbar}{2}\right) \times \hbar \times 2\left(\frac{\hbar-1}{2}\right)}$$
(25)

$$s_1 = R * \cos(\theta) \tag{26}$$

$$s_2 = R * \sin e(\theta) \tag{27}$$

where the fixed constant values with the range up to 0.01 and 1.5 were represented by f and ℓ , respectively, the random value between 0 and 1 is denoted as R_1 and R_2 , and the constant value is notated as \hbar .

Consequently, the Aquila is in an exploitation position approaching the prey along with providing a pre-emptive attack. This behavior can be articulated as,

$$\aleph_n^{\text{behav}} = (\aleph_{\text{best}}(\tau) - \aleph_r(\tau)) * \lambda - R + (UL - LL) * R + LL \\ \times \zeta$$
(28)

where the adjustment parameteris denoted as λ and ζ .

Consequently, the Aquila takes the prey by walking on the ground. The last location at which the Aquila attacked the prey is computed as,

$$\aleph_n^{\text{exploitation}} = Q \times \aleph_{best}(\tau) - (\upsilon_1 \cdot \aleph_n(\tau) - R) - \upsilon_2 \cdot L(d) + R \cdot \upsilon_1$$
(29)

where a quality function was signified by Q, the motions employed during tracking the best solution is notated as v_1 , and it is articulated as,

$$\upsilon_1 = 2 \times R - 1 \tag{30}$$

 v_2 is a lowering worth as of 2 to 0, and it is computed as,

$$v_2 = 2 \times \left(1 - \frac{\tau}{t}\right) \tag{31}$$

The weights are updated in the developed network identical to the Aquila attacking prey among the number of prey. The updated weights φ_n are articulated as,

$$\phi_n \in \aleph_n = \{\phi_1, \phi_2, \phi_3, \dots, \phi_N\}$$
(32)

The input features are subjected to the developed neural network's convolution layer after updating weights. For the assumed contribution statistics Ψ , the difficulty is exactly spoken for example,

$$\Psi = \varepsilon_n \cdot \phi_n \tag{33}$$

After the convolution operation, the activation function is applied to convolutional layers. Learning trivial linear combinations of the inputs was avoided by the proposed work by employing activation functions. Hence, the dropout with a mish activation function was proposed now this paper. The initiation purpose is exactly represented for example,

$$f(\Psi) = \Psi \cdot \tanh(\Omega(\Psi)) \tag{34}$$

Here, the mish activation function is signified as $f(\Psi)$, and the softplus function is notated Ω . To prevent a model from overfitting, the dropout function after the activation function is articulated as,

 $D = \operatorname{prob}(f(\Psi))$

possibilities, the array remains provided to the softmax function that normalized the production actual standards in the variety [0, 1] as of the last CLC. The softmax purpose
$$\Gamma_{\text{soft}}$$
 is signified for example follows,

$$\Gamma_{\text{soft}} = \frac{e^{\Gamma_m}}{\sum\limits_{m=1}^h \Gamma_m}$$
(37)

Anywhere, the CLC production on *m*th lump remains denoted, for example, Γ_m , and the entire amount of production lumps remains illustrated, for example, *h*. Thus, LC can be classified with higher accuracy by this process. The proposed AmiD-DCNN's pseudocode is,

Input: extracted features ε_n **Output**: classification of cancer

(35)

Begin

Initialize parameters \aleph_n, ϕ_n , Compute weight value using ALSAQ $\phi_n \in \aleph_n = \{\phi_1, \phi_2, \phi_3, \dots, \phi_N\}$ For i = 1 to NCompute convolution operation CEvaluate activation function $f(C) = C \bullet \tanh(\Omega(C))$ Perform dropout function D = prob(f(C))Calculate combining process Δ Finish aimed at Flattening all the layers Evaluate softmax activation function Γ_{soft} Return classified cancer End begin

Here, the probability function is represented as *prob*, and the dropout purpose's production is meant for example D. Aggregating the material lengthways by weakening the information was the goal of the combining layer to which the obtained output is passed. The combining process's Δ outcome remains:

$$\Delta = \frac{D - \phi_n}{S} + 1 \tag{36}$$

Anywhere, the kernel pace remain mentioned for example S. The acquired features are flattened and transferred into the FCL where the entire features are converted into the one-dimensional array Γ . Toward board lesson

4 Results and discussions

In comparison with the benchmark techniques, the future LCD replica's presentation remains analyzed experimentally in this section. The experiments are executed on the working platform of Python. The data are gathered as of the publically available chest CT scan images dataset for experiential analysis. Figure 3 expresses a few sample image outcomes on the chest CT scan images dataset by the proposed model.

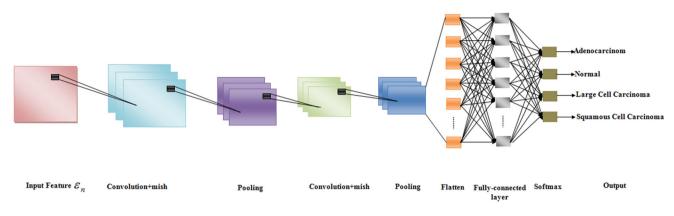
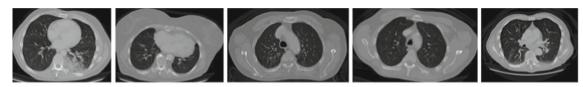


Fig. 2 Structure of proposed AmiD-DCNN



(i)



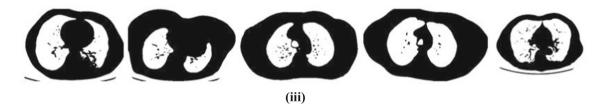


Fig. 3 Sample (i) input (ii) contrast intensified and (iii) segmented images of the proposed model on Chest CT scan images

 $\ensuremath{\text{Table 1}}$ MSE and RMSE analysis on the chest CT scan images dataset

Algorithms	MSE	RMSE
Proposed JQPC	9.981387	3.159333
PC	15.8978	3.987205
K-Medoid	25.82079	5.081416
FCM	28.20381	5.310726

4.1 Performance analysis

Here, patch extraction and classification are the two segments in which the proposed models' performance analysis is executed.

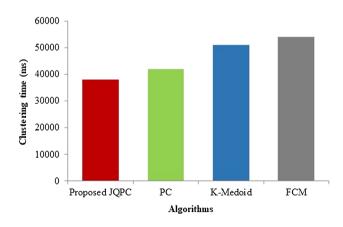
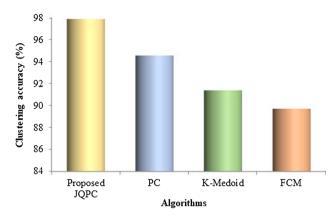


Fig. 4 Time taken for clustering



DNN BBN CNN Proposed AmiD-DCNN 80 85 90 95 100 Precision (%)

Fig. 5 Clustering accuracy analysis

4.1.1 Performance analysis of the patch extraction

Regarding mean square error (MSE), root MSE (RMSE), clustering time, and clustering accuracy, the proposed JQPC algorithm's performance is comparatively analyzed with the prevailing PC algorithms.

In comparison with the prevailing PC, K-Medoid, and FCM algorithms, the experimental outcomes of MSE and RMSE metrics for the proposed JQPC algorithm are demonstrated in Table 1. Here, 9.98 MSE was produced by the proposed model which is lower than the prevailing PC (15.89), *K*-Medoid (25.82), and FCM (28.20). The MSE's square root values are the RMSE values. Abnormal regions and normal regions are portioned with the least error values by utilizing the proposed JQPC algorithm, which is proved in Table 1.

To partition image regions into normal and abnormal regions, the time taken by the algorithm is the clustering time. It is verified as of Fig. 4 that 4051 ms, 12,972 ms, and 15,959 ms less than the existing PC, *K*-Medoid, and FCM algorithms were attained by the proposed JQPC. This exhibits that the regions were clustered faster than the prevailing algorithms by the proposed JQPC.

Figure 5 gives the comparative analysis of the proposed and the prevailing model's clustering accuracy. The clustered output's quality was determined by evaluating the clustering accuracy. Here, PC gives more accurate

Table 2 Training time analysis for the classifier algorithms

Algorithms	Training time (ms)	
Proposed Amid DCNN	40,007	
CNN	45,005	
RNN	50,010	
DBN	55,002	
DNN	60,011	

Fig. 6 Precision analysis of the proposed AmiD-DCNN algorithm

(94.57%) results than the existing *K*-Medoid (91.31%) and FCM (89.72%) among the prevailing algorithms. However, 3.53% more accurate results than the PC algorithm was attained by the proposed algorithm. Implementing Jaccard similarity and Quadratic kernel in the PC algorithm caused this improvement.

4.1.2 Performance analysis of classification

Regarding training time, precision, recall, f-measure, sensitivity, specificity, and accuracy, the proposed Amid DCNN's experiential analysis is analyzed with the prevailing CNN, recurrent neural network (RNN), deep belief network (DBN), and DNN.

The time taken by the classifier to train all the input parameters is termed training time. Table 2 depicts the experimental results obtained during the experimental analysis. Here, 40,007 ms is the proposed Amid DCNN's training time, which is the least time taken than the CNN (45,005 ms), RNN (50,010 ms), DBN (55,002 ms), and DNN (60,011 ms). This exhibits that fewer time computations were taken by the proposed classifier.

Regarding precision, the proposed classifier's experimental analysis and its comparison with existing

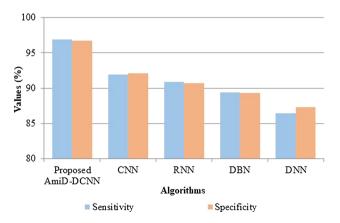


Fig. 7 Performance analysis in terms of sensitivity and specificity

 Table 3 Experimental outcomes of classification accuracy

Algorithms	Accuracy (%)	
Proposed AmiD-DCNN	96.7968	
CNN	91.99199	
RNN	90.79079	
DBN	89.4	
DNN	86.89718	

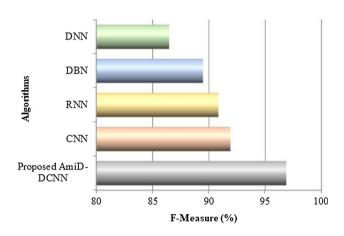


Fig. 8 F-measure analysis on the chest CT scan images dataset

algorithms are signified in Fig. 6. The precision value determines whether the LC stages are predicted correctly or incorrectly. Here, poor outcomes than the other classifiers were provided by the DNN (86.46%). Followed by CNN, the DBN and RNN perform slightly better than the DNN. By attaining a precision of 96.88%, the other baseline techniques were dominated by the proposed AmiD-DCNN. Owing to the utilization of Aquila-optimized mish dropout in the DCNN classifier, a better value is obtained.

The output result's quality was evaluated by the metrics: sensitivity and specificity. Figure 7 pictorially represented the experimental outcomes of sensitivity and specificity. It is evident from Fig. 7 that a better performance than the baseline classifiers was attained by the proposed AmiD-DCNN. 96.88% is the proposed classifier's sensitivity which is higher than the prevailing CNN (91.91%), DBN

Table 4 Comparative analysis of the recent works

Algorithms	Accuracy (%)	Specificity (%)
DFD-Net	87.8	89.1
CNN	90.1	91.7
IDNN	93.9	93
FPSOCNN	95.62	96.32
Proposed AmiD-DCNN	96.79	96.70

(89.4), and DNN (86.46%). The future replica's specificity is 96.70%, which remains better than the other algorithms. This analysis exhibits more quality output than the state-ofthe-art classifiers were given by the proposed AmiD-DCNN.

During the experimental analysis for the proposed and the existing classifiers, the accuracy outcomes are given in Table 3. As the accuracy determines how accurately the outcomes are predicted, the system's accuracy should be high as possible. Here, 96.79% is the proposed AmiD-DCNN's accuracy, which is higher than the CNN, RNN, and DNN by 5.22%, 6.61%, and 11.39%. This exhibits that the proposed classifier is more appropriate than the baseline classifiers for LC stages classification.

The f-measure values obtained by the proposed and the existing classifier algorithms are illustrated in Fig. 8. To give accurate outcomes, the f-measure value should be high for a better scheme. 96.88% is the AmiD-DCNN's F-measure value, which is followed by CNN (91.91%), then RNN (90.83%), and so on. This proves that the proposed classifier predicts the output more reliably than the prevailing baseline classifiers.

4.2 Comparative analysis with the literature papers

Regarding accuracy and specificity, the future replica's performance is analyzed now comparison with the prevailing works like CNN (Yu et al. 2020), IDNN, FPSOCNN, and DFD-Net.

The proposed AmiD-DCNN algorithm's accuracy and specificity analysis with the prevailing algorithms was unveiled in Table 4. It is seen from Table 4 that 1.22%, 3.29%, and 10.23% more accurate results than the existing FPSOCNN, IDNN, and DFD-Net models were attained by the proposed AmiD-DCNN. Thus, it can be concluded that the LC could be detected more effectively than the prevailing works by the proposed model.

5 Conclusion

In this paper, a novel AmiD-DCNN-founded LC organization framework is future. Here, the RU-AlexNet segmented the tumor regions, and the JQPC algorithm partitioned the patches. Finally, the data were classified by the AmiD-DCNN into the stages of LC. The proposed model's experiential outcomes are proved as a better model than the prevailing algorithms. For instance, 94.57% was achieved by the proposed JQPC, which was far better than the compared algorithms. By achieving the average accuracy of 96.79% with less training time of 40007 ms, the proposed AmiD-DCNN classifier performance was proved. In the comparative analysis with recent works, a improved presentation stood showed through the future perfect. Aimed at LC prediction, the analyses exhibited that dominance over other models was attained by the proposed models. The LC stages were successfully classified by the proposed model. However, the risk rate was not predicted. In the future, an advanced risk prediction approach will be developed along with the proposed model grounded on the cancer-affected region.

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Data availability Data included in article/supplementary material referenced in article.

Declarations

Conflict of interest The authors have no conflicts of interest to declare relevant to this article's content.

Human and animal rights This research does not involve any human participants and/or animals; hence, any informed consent or statement on the welfare of animals does not apply to this research.

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