



## Bio-Inspired Spider Wasp Optimization-Based Convolutional Neural Network for Lung Cancer Classification

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### ABSTRACT

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*lung cancer classification, CNN, spider wasp optimization, hyperparameter tuning*

Lung cancer is one of the deadliest diseases in the world affecting nearly five million individuals per year. The accurate detection and classification of lung cancer are crucial for proper treatment planning and increasing survival rates. In recent years, the convolutional neural network (CNN) emerged as an effective approach in lung tumor identification. However, the CNN face challenges such as computational complexity, increased training time, and limited generalization. This study proposed an optimized version of CNN named Spider Wasp optimization CNN (SWoCNN) for precise classification of lung cancer from computed tomography (CT) images. The presented SWoCNN incorporates the fast exploration capacity of spider wasp optimization (SWO) into the CNN algorithm for enhanced lung cancer classification. The developed strategy begins with the collection of lung CT images from both normal and cancer patients and the images are preprocessed to increase their quality. Consequently, an attention neural network (ANN) is developed for capturing and extracting the relevant features from the preprocessed images. Further, feature selection was done using the waterwheel plant algorithm (WPA), which selects the most informative features crucial for distinguishing the patterns of lung cancer. Finally, the proposed SWoCNN was trained using the selected features to classify lung cancer types. The SWO leverages its searching capacity to optimize and fine-tune the CNN hyperparameters to its optimal value, leading to improved classification, reduced computational time and fast CNN training. The proposed strategy was modeled and implemented in the Python platform and experimental results highlighted that the designed methodology achieved an average performance such as accuracy of 0.96407, precision of 0.96396, specificity of 0.96503, and false negative rate of 0.02963. Furthermore, a comparative assessment with existing models depicted that metrics such as accuracy, precision, recall, and specificity are enhanced by 1.23%, 1.24%, 1.19%, and 1.18%, respectively, depicting its reliability in classifying lung cancer.

## 1. INTRODUCTION

Lung cancer is one of the deadliest diseases that occur due to the abnormal growth of cells in the lungs, causing damage to healthy lung tissues [1]. The common cause of this cancer is tobacco usage and it accounts for nearly 80% of lung cancer deaths [2]. The study by the World Health Organization (WHO) on cancer reported that lung cancer has led to approximately 1.61 million deaths and it is more prominent among males than females. Early and precise identification of this disease is crucial for improving the patient's health and survival rate [3]. Currently, various imaging techniques including magnetic resonance imaging (MRI), Computed Tomography (CT), and X-ray are employed for lung cancer prediction [4]. Among these techniques, CT was widely used for lung cancer detection because of its effectiveness in capturing minute variations in the lung region [5]. Conventionally, the medical images are manually examined

by healthcare professionals or radiotherapists to detect the presence of cancer [6]. However, this manual examination of cancer is time-consuming and it depends on the knowledge of the healthcare experts. Thus, researchers focused on developing an automatic framework leveraging advanced technologies such as artificial intelligence (AI), computer-aided processing, big data analysis, etc. [7]. In this automatic mechanism, the system was trained using the data containing both healthy and cancer features to understand the variations between them [8].

In recent times, AI approaches such as machine learning (ML) and deep learning (DL) have earned greater attention in image processing tasks. The AI-based models analyze the medical images and capture the features and hierarchical relations to detect the presence of lung cancer [9, 10]. Initially, the researchers utilized ML algorithms such as decision trees (DT), random forest classifier (RF), logistic regression, and support vector machine (SVM), for lung cancer detection [11,

12]. They process the medical images and train using either supervised or unsupervised learning strategies for classification tasks. The studies on ML models concluded that they offered automatic classification with increased accuracy than the manual process. Despite its increased performances, it faces challenges such as large computational demands, algorithmic bias, lack of adaptability, reduced accuracy in test cases, data dependency, etc. [13-15]. On the other hand, the DL models including deep neural network (DNN), CNN, and feed-forward neural network (FFNN) have shown promising results in image classification [16].

Among these neural networks, CNN is more suitable for medical image analysis because of its potential to capture features and relations within the images more effectively. Hence many researches are conducted on lung cancer classification using CNN models [17, 18]. The existing works developed techniques such as Fuzzy Particle Swarm Optimization with CNN [19], Deep CNN [20], 2-dimensional CNN [21], 3-dimensional CNN [22], autoencoder-based CNN [23], etc., for lung cancer detection. These strategies mainly focused on differentiating malignant and benign cells in the lung and achieved accuracy of around 86% to 95%. However, these frameworks face challenges such as computational complexity, minimum accuracy, instability, less generalization, and limited to binary classification. To address these issues, a novel optimized CNN was developed in this work for detecting and classifying lung cancer from CT images.

The major contributions of the presented framework are described below:

- This study develops a hybrid classifier by integrating a convolutional neural network with spider wasp optimization for lung cancer identification and classification.
- The proposed framework employs an attention neural network for capturing the informative features from the pre-processed images.
- A meta-heuristic-based feature selector was designed using the Waterwheel Plant Algorithm to optimally select the most relevant attribute from the extracted feature sequences.
- The CNN in the developed algorithm learns the hierarchical and spatial feature representations and recognizes the lung cancer pattern, while the SWO optimizes CNN's training by refining its parameters to optimal value.
- The presented technique was executed in Python software and its outcomes are validated with conventional classifier models using metrics like accuracy, precision, false positive rate (FPR), recall, f-measure, specificity, FNR, and execution time.

## 2. RELATED WORKS

Identifying the cancerous nodules in lung CT images is crucial for diagnosing and classifying lung cancer. However, it is a complex and time-consuming process because of the varying size, texture, and shape of nodules. Asuntha and Srinivasan [19] presented an innovative solution for this problem using the DL model. This study designed a unique feature selector model by combining fuzzy function with particle swarm optimization (FPSO) and the selected features using FPSO to train CNN. This strategy was validated using the real-time CT image dataset gathered from Arthi Scan

Hospital and the experimental outcomes demonstrated that the usage of FPSO reduces the computational complexity of the CNN. However, it faces complexity in validating the correctness and reliability of fuzzy variables, sets, and membership functions.

Faruqui et al. [20] designed a hybrid classifier model named LungNet having 22 layers of CNN. This study aims to identify and categorize lung cancer from CT scans using a deep CNN algorithm. This study used the lung cancer dataset containing 525,000 images. The results of the study illustrated that it achieved 94.81% accuracy and 3.35% false positive rate. However, the model's accuracy was reduced to 91.6% when classifying the different stages of lung cancer.

Biradar et al. [21] proposed an automatic lung cancer classifier using the DL algorithm. This study intends to detect lung cancer nodules with improved speed and accuracy. A 2-dimensional CNN was created to classify malignant and benign cells in the lungs and it was validated using the Kaggle CT scan dataset. The implementation results manifested that it achieved only 88.76% in detecting cancer nodules, which is not sufficient for real-world clinical scenarios.

Bangare et al. [22] developed a computer-aided approach to categorize lung cancer from CT images. Its primary concern is to increase the probability of survival rate of lung cancer-affected individuals through accurate and timely cancer detection. This work developed a 3-dimensional CNN with image pre-processor and segmentation to detect malignant and non-malignant cells in the lungs. The experimental results highlighted that this strategy achieved an accuracy of 86.42%, specificity of 86.72%, and recall of 86.11%. However, these performances are not sufficient for lung cancer classification in real healthcare institutions.

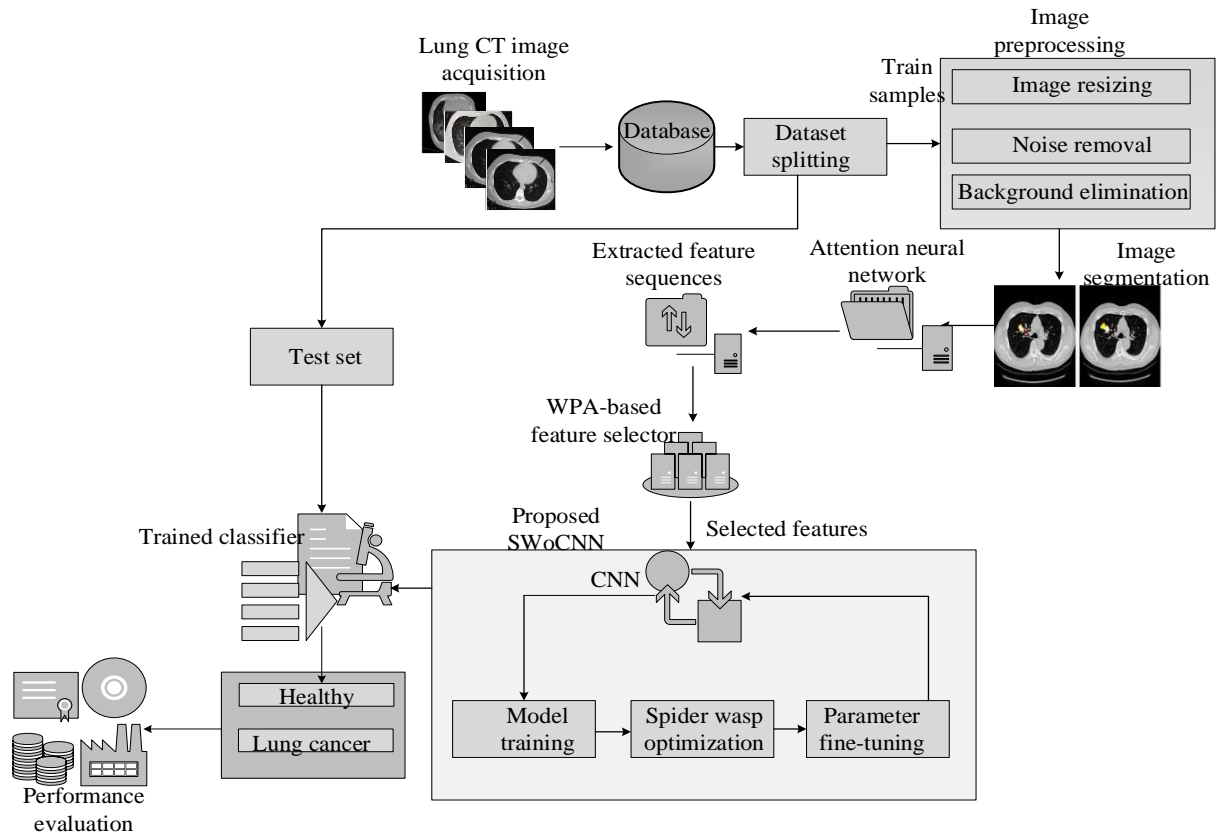
The previous studies [23-27] developed an integrated framework for classifying lung cancer. This strategy combined the efficiency of the autoencoder and CNN for efficient feature learning. In addition, an Adam optimizer was deployed for optimizing the entire classifier model, and a multispace image reconstruction algorithm was used to minimize the error. The results of the work suggested that it earned 95.79% accuracy in lung cancer nodule detection. Also, it obtained less processing time of 15 seconds (s). However, this methodology is data-dependent and cancer detection from low pixel images lowers its accuracy.

## 3. PROPOSED SWOCNN FOR LUNG CANCER CLASSIFICATION

An optimized CNN was developed to classify lung cancer using lung CT images. The presented framework comprises five modules: image acquisition, image preprocessing, feature analysis, feature selection, and optimized model training and classification. In the data collection module, the CT images of healthy and lung cancer-affected individuals are gathered and imported into the system. The accumulated images are preprocessed following steps like noise filtering, image resizing, and background removal. Also, the U-Net algorithm was used for segmenting the region of interest (ROI) from the preprocessed images. In the feature extraction module, ANN was designed to capture the relevant attributes from the segmented images, which reduces the dimensionality of the data and improves training speed. Consequently, the WPA-based feature selector was modeled to select the most informative and highly correlative features from the extracted

feature sets, enabling the classifier to concentrate on major features for tumor classification. Finally, an optimized CNN was developed by integrating the efficiency of SWO into it. The SWO continuously refines and optimizes the

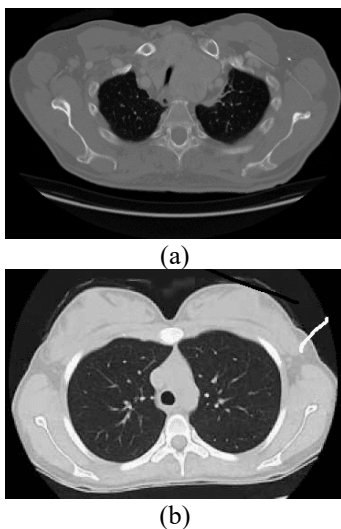
hyperparameters of CNN, enabling it /to adapt to evolving changes and enhancing its learning capacity. The architecture of the developed lung cancer classifier model is displayed in Figure 1.



**Figure 1.** Architecture of SWoCNN for lung cancer classification

### 3.1 Image acquisition

The first stage of the developed framework is the lung CT image acquisition from both healthy and cancer-affected persons as shown in Figure 2. The gathered images are labeled to train the classifier to recognize the patterns of lung cancer. The presented study used the publicly available Lung CT scan database from Kaggle site [27].



**Figure 2.** Dataset images of different classes: (a) Lung cancer; (b) Normal

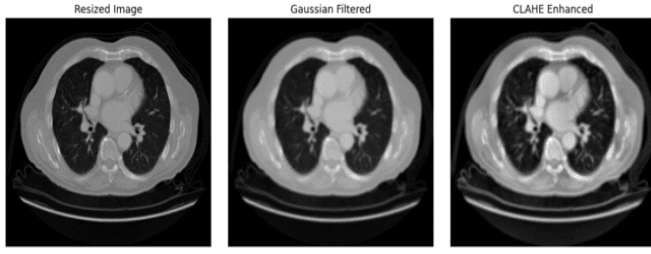
### 3.2 Image preprocessing

Image preprocessing defines the data preparation phase, where the system converts the raw CT images into an appropriate format for further processing. This stage follows a sequence of processes such as noise filtering, image resizing, and contrast enhancement or background elimination. In the noise filtering step, a Gaussian filter was applied to the images to eliminate the noise or unwanted random variations within each image. The Gaussian filter removes the noise attributes and replaces them with the average value of nearby or surrounding attributes, which is estimated by Gaussian distribution. The mathematical formulation of the Gaussian filter is expressed in Eq. (1):

$$Gs(m,n) = C_{im}(m,n) \times \frac{1}{2\pi\sigma^2} e^{-\frac{m^2+n^2}{2\sigma^2}} \quad (1)$$

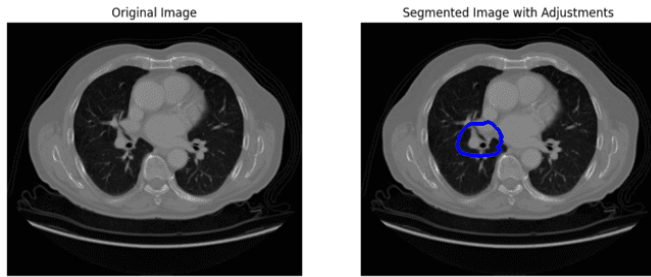
where,  $Gs(m, n)$  represents the Gaussian filter,  $C_{im}(m, n)$  denotes the input CT image,  $(m, n)$  indicates the kernel coordinates, and  $\sigma$  refers to the standard deviation. Consequently, the filtered images are resized into a common size to boost the dataset consistency. Finally, background removal was done to discard the unwanted regions from the CT images. This process is done using image thresholding. This step helps the system to focus on useful regions in the images. In addition, data augmentation was performed to reduce the risk of overfitting. This step creates new training

samples by performing steps like rotation, random translation, scaling, image mixing, etc. Figure 3 displays the preprocessed images.



**Figure 3.** Preprocessed images

Further, image segmentation was done using the U-Net algorithm to isolate ROI from the preprocessed images [28]. This technique has the potential to segment the regions with training using less data. It consists of three main elements namely: encoder, bottleneck, and decoder. The encoder component down-samples the preprocessed images through convolution and pooling operations and maps the regions within each image. The bottleneck is placed between encoder and decoder and it maps the most informative regions while retaining spatial information. The encoder performs upsampling and reconstructs the image from the feature representation. The reconstructed image is the segmentation output. Figure 4 displays the segmentation outcomes.



**Figure 4.** Segmentation images

### 3.3 Feature extraction

An attention model is a kind of DL approach, which tends to concentrate on significant parts of the input data. In the proposed work, it was used to capture the relevant features from the segmented results. Feature extraction is one of the crucial steps in cancer classification, which aims to minimize the data dimension and assists the classifier in quickly learning the feature representations [29]. The architecture of ANN is similar to the structure of the encoder-decoder and finds the relevant features through the calculation of attention weights. It includes three main components namely: encoder, attention, and decoder. The encoder accepts the segmented images as input and generates hidden states (features present in the segmented images). The attention module determines the relevance between the encoder's hidden state and the target hidden state and produces similarity scores as outcomes, which are mathematically expressed in Eq. (2):

$$Sw_{xt}^y = \omega \left( \tanh \left( w_e h_{xt}^y + b_{we} \right) \right) + \beta \quad (2)$$

where,  $Sw_{xt}^y$  represents the similarity score,  $\omega$  denotes associated with ANN,  $\tanh$  defines the activation function,  $h_{xt}^y$  indicates the hidden state generated by the encoder,  $b_{we}$  denotes the bias vector associated with the attention component,  $w_e$  represent the weight vector of the attention module, and  $\beta$  indicates the bias vector of ANN. The resultant of the attention component is converted into attention weights using a Softmax function, as defined in Eq. (3):

$$Aw_t = \text{soft max} \left( Sw_{xt}^y \right) \quad (3)$$

where,  $Aw_t$  denotes the attention weights, which represents the relevance of the extracted feature for lung cancer categorization. These weights are applied to the encoder's hidden state and all the relevant information is extracted based on their importance through a weighted sum function, as represented in Eq. (4):

$$W_{st} = \sum_{t=1}^{T_m} Aw_t h_{xt}^y \quad (4)$$

where,  $W_{st}$  denotes the weighted sum and  $T_s$  indicates the maximum iteration. The decoder component receives the weighted sum and transforms it into a feature vector. This extracted feature vector contains all important information related to lung cancer classification.

### 3.4 Feature selection

Feature selection represents the process of selecting the most relevant attributes while eliminating the irrelevant ones. This step aims to enhance interpretability, reduce complexity, and increase accuracy. In the developed work, a novel feature selector was modeled using the meta-heuristic optimization algorithm named "Waterwheel plant algorithm [30]." The WPA is a bio-inspired approach modeled based on the natural characteristics of waterwheel plants to solve complex optimization problems. This algorithm is mathematically designed following the hunting expedition behavior of waterwheel plants. They use search agents to find the prey. Here, the concept of this algorithm is applied to select the most relevant and highly informative features for classifier training. The feature selection process begins with the initialization of extracted feature sets in the search space. In WPA, the extracted feature sets by ANN are expressed as a matrix using Eq. (5) and each feature is initialized in the search space using Eq. (6):

$$F_e = \begin{bmatrix} F_{e1} \\ \vdots \\ F_{ei} \\ \vdots \\ F_{eQ} \end{bmatrix}_{Q \times R} = \begin{bmatrix} f_{e(1,1)} \cdots f_{e(1,j)} \cdots f_{e(1,R)} \\ \vdots & \ddots & \vdots & \ddots \\ f_{e(i,1)} \cdots f_{e(i,j)} \cdots f_{e(i,R)} \\ \vdots & \ddots & \vdots & \ddots \\ f_{e(Q,1)} \cdots f_{e(Q,j)} \cdots f_{e(Q,R)} \end{bmatrix} \quad (5)$$

$$f_{e(i,j)} = L_{bj} + \kappa_{i,j} \cdot (U_{bj} - L_{bj}) \quad (6)$$

$i = 1, 2, 3, \dots, Q, j = 1, 2, 3, \dots, R$

where,  $F_e$  indicates the extracted feature sets,  $F_{ei}$  denotes the  $i^{th}$  feature (candidate solution),  $f_{e(i,j)}$  represents the  $j^{th}$  problem variable,  $\kappa_{ij}$  represents a random number (0, 1),  $U_{bj}$  and  $L_{bj}$  refers to the upper bound and lower bound. Further, the fitness value was determined for each feature based on their relevance to the tumor classification task, which is defined in Eq. (7):

$$\gamma_v = \gamma_v \begin{bmatrix} F_{e1} \\ \vdots \\ F_{ei} \\ \vdots \\ F_{eQ} \end{bmatrix}_{Q \times R} \quad (7)$$

The candidate solution with a high fitness value indicates the best or optimal solution and the solution with the lowest value denotes the worst. After fitness evaluation, the next step is exploration, where the system explores the search space and updates its initial values. This exploration helps to find the optimal solution and mitigates the possibility of local optima. The exploration phase of WPA is modeled using Eqs. (8) and (9):

$$\vec{A} = \vec{d} \cdot (f_e(t) + 2\rho) \quad (8)$$

$$f_e(t+1) = f_e(t) + \vec{A} \cdot \left( 2\rho + \vec{d}_1 \right) \quad (9)$$

where,  $\vec{A}$  denotes the direction of moving in the search space,  $f_e(t+1)$  indicates the updated solution,  $f_e(t)$  defines the current solution,  $\vec{d}_1 \vec{d}$  refers to the random vector, and  $\rho$  represents the optimization factor. Consequently, the system exploits the current solutions to estimate the optimal features. This exploitation phase follows the hunting and feeding characteristics of WPA and it is formulated in Eqs. (10) and (11):

$$\vec{A}' = \vec{d}_2 \cdot \left( \rho \cdot f'_e(t) + \vec{d}_3 f_e(t) \right) \quad (10)$$

$$f_e(t+1) = f_e(t) + \rho \vec{A}'. \quad (11)$$

where,  $\vec{A}'$  denotes the direction of exploitation,  $\vec{d}_2$  and  $\vec{d}_3$  indicates the random vector. Further, fitness value was determined for the updated solutions. Then, the feature with a fitness value greater than 0.5 is selected and this process continuous until reaching maximum iteration or maximum convergence rate. The selected features are fed into the classifier for training.

### 3.5 Optimized CNN for lung cancer classification

A meta-heuristic optimizer-based CNN was developed to accurately identify and categorize lung cancer by processing the CT images [31]. The developed algorithm incorporates the optimization and exploration characteristics of SWO into

CNN, enabling the system to adapt to the evolving variations in the image features and to train the model with optimal values. The CNN is a popular DL model used for performing tasks such as object detection, speech recognition, video labeling, image classification, facial recognition, etc. It consists of five important layers namely: input, convolutional, pooling, and fully-connected (FC) layers. The input layer accepts the selected feature sets from the WPA module as classifier input and transforms them into a suitable format for feature learning. The convolutional layer is the building block of CNN, which is responsible for understanding the patterns within each image. It applies a sequence of kernel filters on the input data and creates a feature map, which represents the patterns influencing healthy and lung cancer-affected images. The feature map is mathematically expressed in Eq. (12):

$$F_{map}(p, q) = \sum_{i=0}^{x-1} \sum_{j=0}^{y-1} S_f(x, y) * K(p-x, q-y) \quad (12)$$

where,  $F_{map}(p, q)$  indicates the feature map at the position  $(p, q)$ , and  $K(p-x, q-y)$  represents the kernel filter applied at the location  $(p-x, q-y)$ . Each convolutional layer is followed by a Rectified Linear Unit (ReLU) activation function to introduce non-linearity in the system. After each convolution+ReLU combination, a pooling layer was placed to reduce the spatial dimension, while retaining the significant information. This layer aims to minimize the memory and helps prevent overfitting by performing pooling operations. The pooling operation defines the process of sliding a 2-D filter over each channel of the feature map. In the designed CNN, a max pooling layer was used, which chooses the maximum element from the feature map covered by the 2-D filter. After multiple convolutional and pooling layers, the resultant feature map was flattened into a one-dimensional vector. This learned feature vector contains the patterns and relations between them, enabling them to categorize the lung cancer classes. The FC layer accepts this feature and computes the lung cancer classification through a Softmax function. The Softmax function returns the possibility of the input image belonging to the class, as represented in Eqs. (13) and (14):

$$P_{vl}(C_{im}) = \text{Soft max}(W_{fc} * f_{le}) + b' \quad (13)$$

$$C_{cl} = \begin{cases} \text{if } (0 > P_{vl} \leq 0.2), & \text{normal} \\ \text{if } (P_{vl} \geq 0.3), & \text{SCC} \end{cases} \quad (14)$$

where,  $P_{vl}$  indicates the probability value,  $C_{cl}$  denotes the classification function,  $W_{fc}$  defines the weight matrix of the FC layer,  $f_{le}$  represents the flattened feature vector, and  $b'$  refers to the bias vector of the FC layer. Consequently, the loss or error was determined to estimate the efficiency of the classifier and it is formulated in Eq. (15):

$$Loss = \frac{1}{S} \sum_{i=1}^S \sum_{j=1}^C Y_i^j \cdot \log(Y_i'^j) \quad (15)$$

where,  $Y_i^j$  denotes the ground truth outcome,  $S$  refers to the number of training samples,  $C$  indicates the number of classes, and  $Y_i'^j$  represents the predicted class. Reducing this loss is

significant for improving classification accuracy. Also, the CNN's performance depends on the value of its hyperparameters such as several filters, kernel size, hidden layer, learning rate, batch size, weights, and bias vectors. To optimally select the values of these parameters, the SWO was integrated into CNN. The SWO is a meta-heuristic optimization algorithm developed based on the searching, hunting, mating, and nesting characteristics of female wasps [32]. The female wasps usually do these processes by depositing solitary egg within each spider's abdomen. In the beginning stage of optimization, the female wasps explore their surroundings for finding suitable spiders. Then, they immobilize and pull them to make nests. After identifying appropriate nests and finding prey, they intend to drag the prey into nests. Unlike conventional optimization models such as genetic algorithm (GA), particle swarm optimization (PSO), grey wolf optimization (GWO), etc., the SWO dynamically switches between exploration and exploitation phases, leading to faster convergence and prevents premature convergence. In addition, the SWO model is more robust and efficient in optimizing the complex and high-dimensional spaces. Moreover, it offers diversified search options, which avoids the local optima. Since the SWO was used for selecting the optimal values in the search range, only the hunting phase is used in the designed model. The mating and nesting phases are not considered as they don't directly relate with parameter update task, they are intended to reproduction and settlement of wasps in the biological systems. The proposed optimization involves four phases: (i) initialization, (ii) fitness evaluation, (iii) searching phase, and (iv) parameter selection and termination.

#### i. Initialization

The SWO algorithm commences with the initialization of the female wasps. Each female wasp represents the CNN parameter set and it is initialized with a random value in the search space. In the context of CNN parameter tuning, female wasp defines the hyperparameter population, and prey denotes their optimal value. The wasp population and the random initialization process are defined in Eqs. (16) and (17):

$$Pr = \begin{bmatrix} Pr_{(1,1)} & Pr_{(1,2)} & \dots & Pr_{(1,d)} \\ Pr_{(2,1)} & Pr_{(2,2)} & \dots & Pr_{(2,d)} \\ \vdots & \vdots & \vdots & \vdots \\ Pr_{(c,1)} & Pr_{(c,2)} & \dots & Pr_{(c,d)} \end{bmatrix} \quad (16)$$

$$\vec{Pr}_t = \vec{L}_u + \vec{r} \left( \vec{K} - \vec{L}_u \right) \quad (17)$$

where,  $Pr$  indicates the parameter population,  $\vec{L}_u$   $\vec{K}$  denotes the lower and upper bound of the search space and  $\vec{r}$  represents the random initialization vector.

#### ii. Fitness evaluation

For each parameter set, fitness was calculated based on the predefined objective function. The objective function of the SWO is to reduce the loss incurred by CNN, and it is formulated in Eq. (18):

$$Obj_{fun} = \min \left( \frac{1}{S} \sum_{i=1}^S \sum_{j=1}^C Y_i^j \cdot \log(Y_i'^j) \right) \quad (18)$$

The parameter set with high fitness indicates the optimal solution and vice versa. Then, the best candidate solution was estimated by sorting them as per their fitness value.

#### iii. Searching phase

This phase follows how female wasps search for the most suitable spiders for feeding their offspring. They use a constant step for searching for prey by randomly looking into the search space. Similarly, in the developed work, they look for the optimal value for the parameter set in the search range. In this searching phase, the values of the parameter set will be updated in two ways. The random exploration of search space and parameter update is formulated in Eq. (19):

$$Pr(t+1) = Pr(t) + f \cdot (Pr_a(t) - Pr_b(t)) \quad (19)$$

where,  $Pr(t)$  denotes the current parameter set,  $Pr(t+1)$  denotes the updated parameter set,  $\phi$  represents the constant motion,  $a$  and  $b$  indicates two randomly selected indices to choose the exploration direction. In some instances, the optimization process drops and selects the local optima. To prevent this, the SWO algorithm explores the surrounding search space with a small step size, as expressed in Eq. (20):

$$Pr(t+1) = Pr(t) + \phi \cdot \vec{L}_u + \vec{r} \left( \vec{K} - \vec{L}_u \right) \quad (20)$$

where,  $\vec{r}$  defines the random vector [0, 1]. In searching phase, the updated hyperparameters in the search space are determined using Eq. (21):

$$Pr(t+1) = \begin{cases} Pr(t) + f \cdot (Pr_a(t) - Pr_b(t)); & \text{if } c < g \\ Pr(t) + \phi \cdot \vec{L}_u + \vec{r} \left( \vec{K} - \vec{L}_u \right); & \text{otherwise} \end{cases} \quad (21)$$

where,  $\chi$  and  $\gamma$  defines the random number ranging [0, 1], representing the probability of obtaining local optima in the searching phase.

$$Pr(t+1) = Pr(t) + G \times |2 \times r_e \times Pr_a(t) - Pr_b(t)| \quad (22)$$

where,  $G$  denotes the deviation between the current solution and the best solution, and  $r_e$  indicates the randomness in escaping in range [0, 1]. The formula for calculating  $G$  is expressed in Eq. (23):

$$G = r_e \times \left( 2 - 2 \times \left( \frac{t}{T_m} \right) \right) \quad (23)$$

where,  $t$  indicates the current iteration, and  $T_m$  denotes the maximum iteration. This stage is the beginning of the exploitation stage in SWO approach.



#### v. Hunting phase

This phase follows how the female wasps move towards the region where the best spider (prey) can be found. The hyperparameter enters this stage only when its fitness is less than 0.3 (near optimal) and they are updated by exploiting locally. Instead of searching randomly, the algorithm adjusts current solutions by shifting them closer to the high-quality solutions, as formulated in Eq. (24):

$$Pr(t+1) = Pr(t)' + \cos(2\pi l) \times (Pr(t)' - Pr(t)) \quad (24)$$

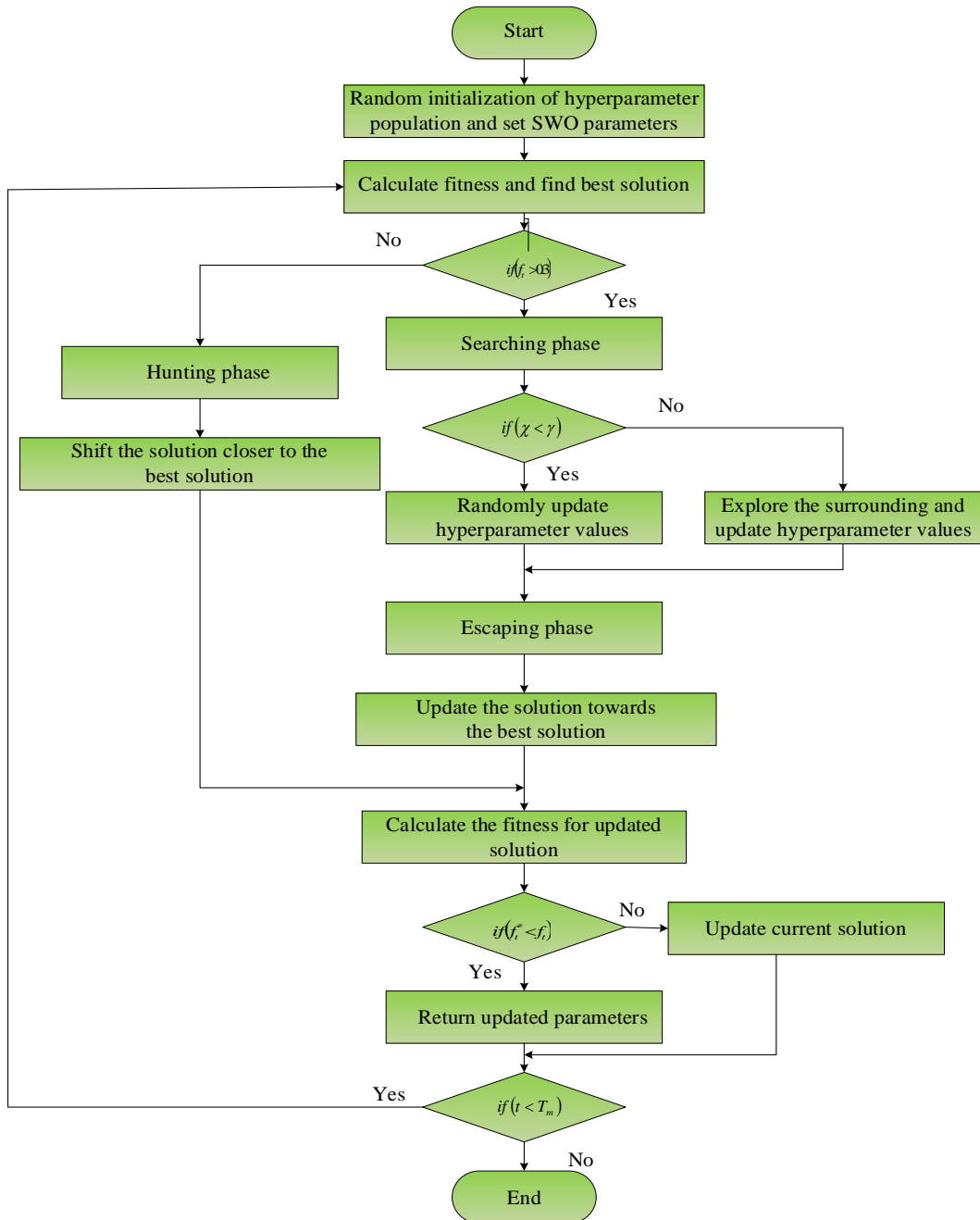
where,  $Pr(t)'$  defines the best available solution. After updating hyperparameters, the fitness value was determined for them for selection purpose.

#### vi. Parameter selection and termination

If the fitness of the updated sequence is greater than the old fitness, the updated sequence was selected for modeling CNN. If the new fitness is less than the old fitness, no changes in CNN design the parameter selection is formulated in Eq. (25):

$$Pr = \begin{cases} \text{if } (f_t^* < f_t), Pr(t+1) \\ \text{else}, Pr(t) \end{cases} \quad (25)$$

where,  $f_t^*$  denotes the fitness of the updated hyperparameter, and  $f_t$  indicates the fitness of the current solution. This process continues until it reaches maximum iteration. Algorithm 1 presents the working of the developed classifier in pseudocode format. Figure 5 presents the flowchart of the SWO algorithm for CNN parameter update.



**Figure 5.** The flowchart of the SWO algorithm for CNN parameter update

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**Algorithm: SWO-CNN**

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**Input:** Maximum iteration  $T_m$ , learning rate, kernel size, hidden layer, batch size, weights and bias vector, CT images;

**Output:** Lung cancer classification;

**Start {**

Design CNN with initial parameters (input, convolutional, pooling, and FC layers);

Train the model using selected features using WPA;

Estimate class probability;

Perform classification using Eq. (14);

Parameter optimization:

Define and initialize SWO parameters like population size  $c$  and decision variable  $d$ ;

Initialize the initial value of the parameter set  $Pr$ ;

Randomly initialize parameter population  $pr$  using Eq. (17)

**Fitness evaluation:**

$Obj_{fun} \rightarrow \min(loss)$

for  $t=1: T_m$

**Searching Phase:**

if ( $x < y$ )

Update parameter set using Eq. (19);

Else

Update parameter set  $Pr$  using Eq. (20);

End if;

**Escaping stage:**

Update the parameters towards the best solution using Eq. (21);

**Hunting Phase:**

if ( $f_i < 0.7$ )

Update the parameter using Eq. (22);

else

Exist; // the parameter set will not be updated in this phase

End if;

Determine fitness  $f_t^*$  for the updated parameter set  $Pr(t+1)$ ;

if ( $f_t^* < f_t$ )

Return  $Pr(t+1)$ ;

else

Return  $Pr(t)$ ;

$t++$ ;

end for;

Return optimal parameter value;

} End

---

$$Accuracy = \frac{t_p + t_n}{t_p + t_n + f_p + f_n} \quad (26)$$

where,  $t_p$ ,  $t_n$ ,  $f_p$  and  $f_n$  represent TP, TN, FP, and FN, respectively.

**Precision:** Precision measures the classifier's efficiency in identifying true positive cases to the total true instances, and it is formulated in Eq. (27). This metric quantifies the effectiveness of the model in classifying lung cancer among the true samples (real normal and lung cancer samples).

$$Precision = \frac{t_p}{t_p + t_n} \quad (27)$$

**Sensitivity/recall:** Recall determines the system's proficiency to detect all relevant instances and it defines the fraction of real positive cases to the total positive cases, as defined in Eq. (28):

$$Recall = \frac{t_p}{t_p + f_p} \quad (28)$$

**F-measure:** F1-score denotes the harmonic mean of sensitivity and precision. It determines the balanced performance of the classifier in categorizing lung cancer, as formulated in Eq. (29). This parameter enables to determine the efficiency of the model in classifying normal and lung cancer images considering the false positive and negatives.

$$F1-score = 2 \times \left( \frac{sensitivity * precision}{sensitivity + precision} \right) \quad (29)$$

**Specificity:** Specificity measures the model's effectiveness in detecting and classifying normal (true negative) instances. It is represented mathematically in Eq. (30):

$$Specificity = \frac{t_n}{t_n + f_p} \quad (30)$$

**False positive rate:** FPR represents the ratio of actual negative cases that the system incorrectly categorizes as positive, as formulated in Eq. (31). It determines the incorrect lung cancer classification made by the system.

$$FPR = \frac{f_p}{t_n + f_p} \quad (31)$$

**False negative rate:** FNR denotes the ratio of real positive cases incorrectly categorized as negative by the system. It is expressed in Eq. (32):

$$FNR = \frac{f_n}{t_p + f_n} \quad (32)$$

**Execution time:**

The execution time determines the overall time consumed

## 4. RESULTS AND DISCUSSION

The developed classifier was modeled and executed in Python software version 3.12.6 running on Jupyter Notebook 7.12, and supported by a Dell 11th Generation processor with 8GB RAM. The presented framework was trained and tested using the publicly available "Lung cancer CT scan dataset" from the Kaggle site.

### 4.1 Performance metrics

The definition and formula of parameters used for estimating the performance of the developed classifier in lung cancer categorization are defined below.

**Accuracy:** Accuracy defines the classifier's effectiveness in categorizing both healthy and lung cancer and it quantifies the proportion of truly classified cases to the total instances. It measures the correctness of the system in classifying both normal and lung cancer images present in the dataset. It is represented in Eq. (26):



by the designed model for performing lung cancer identification and categorization. This measures the overall computational efficiency of the model in performing lung cancer classification.

The investigation of these parameters allows us to estimate how the classifier performs lung cancer classification.

**Table 1.** The hyperparameter and its optimized value

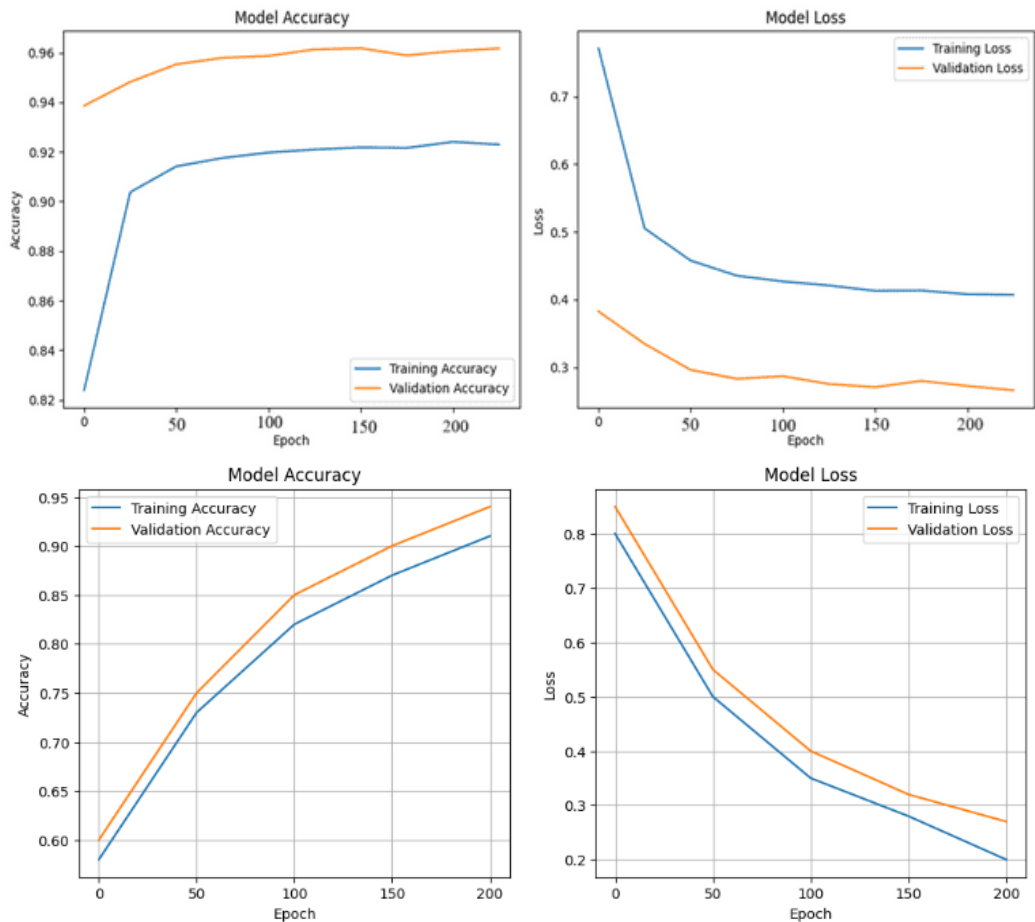
Hyperparameter	Range	Optimized Value
Number of filters for convolutional layer 1	[16, 32, 64, 96]	64
Kernel size for convolutional layer 1	[3, 4, 5]	3
Number of filters for convolutional layer 2	[48, 64, 96, 128]	96
Kernel size for convolutional layer 2	[3, 4, 5]	3
Number of filters for convolutional layer 3	[64, 96, 128]	128
Kernel size for convolutional layer 3	[3, 4, 5]	3
Batch size	[5, 10, 20]	20
Epoch	[50, 100, 200]	200
Hidden Layer (HL) 1	[75, 100, 125]	100
HL 2	[75, 100, 125]	125
Learning rate	[0.001, 0.01, 0.1, 0.2]	0.01
Activation	-	ReLU

4.2 Train and test phases

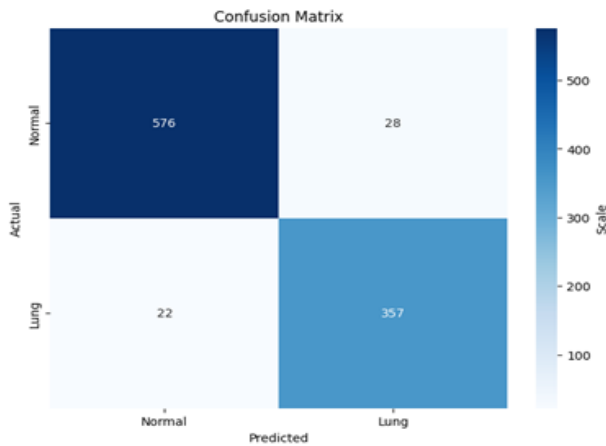
The presented framework was intensively trained with the Lung cancer CT scan dataset. Initially, the input dataset was split into the ratios of 80:20 for the train and test phases. The training phase begins with the initialization of CNN hyperparameters with initial value. The activation function of the CNN was ReLU, and the metrics like number of filters, kernel size, hidden layer range, learning rat, etc., are optimized by the SWO algorithm over the epoch (Table 1).

The outcomes during training and testing are determined using accuracy and loss metrics. Accuracy quantifies the classifier's efficiency in learning the hierarchical feature representations and its generalization capacity against real-world data. On the other hand, the loss measures the error made by the model during the training and test. Higher accuracy suggests that the model correctly classifies the lung cancer instances, while high loss indicates the model's inefficiency in categorizing the lung tumor cases.

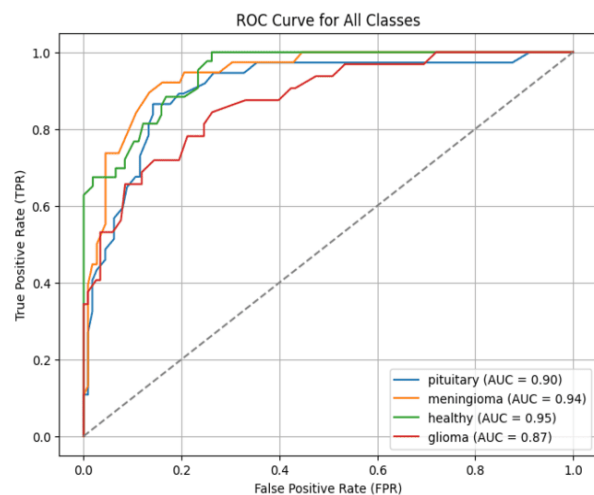
The analysis of accuracy and loss in train and test phases is presented in Figure 6. The improvement of accuracy over the epoch indicates that the designed SWoCNN has the greater potential of learning the features differentiating healthy and lung cancer images. Also, the proposed model obtained lower loss, which demonstrates that it correctly categorizes the tumor classes and generalizes well on unseen samples. Furthermore, it is noted that the presented model obtained higher accuracy irrespective of the datasplits, and the smooth training and testing curve highlights the model’s stability and its capacity to prevent overfitting issue.



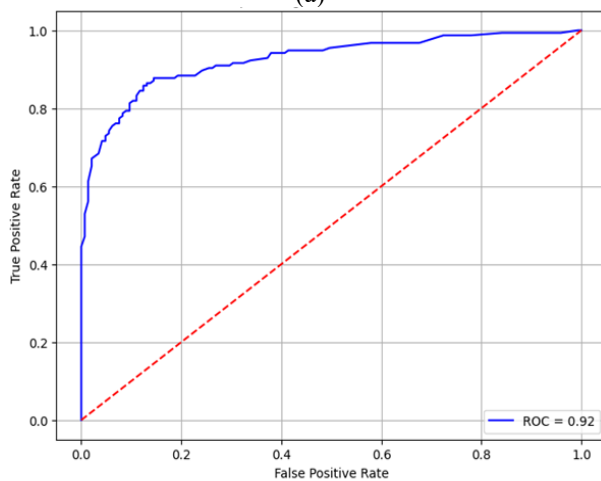
**Figure 6.** Accuracy and loss in train and test stages: 80/20 split and 70/30 split



**Figure 7.** Confusion matrix



(a)

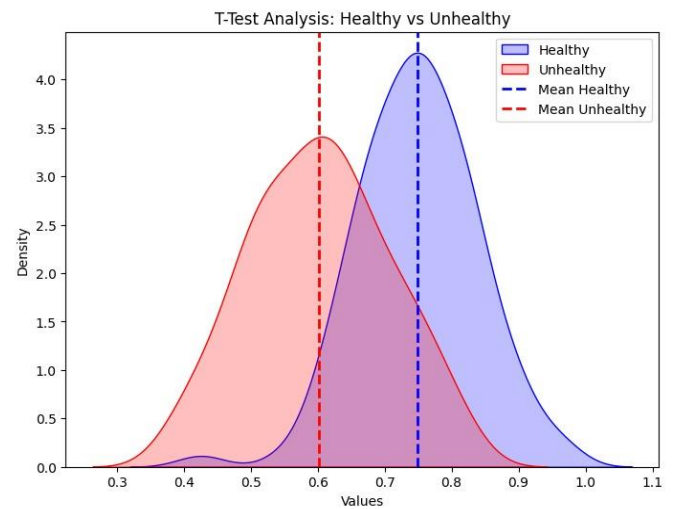


(b)

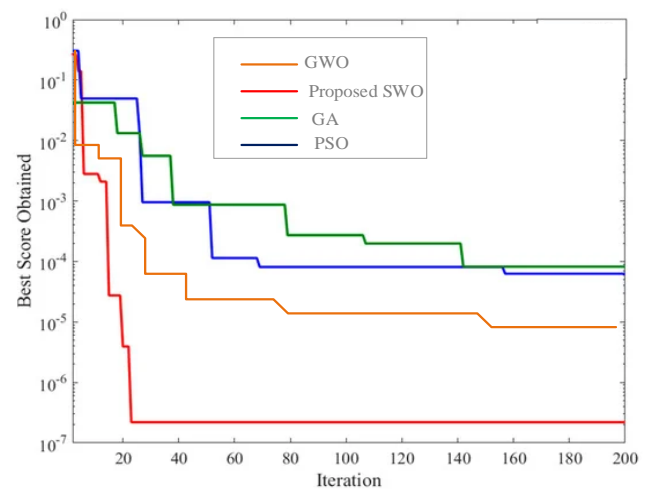
**Figure 8.** ROC curve (a) 80/20 datasplit (b) 70/30 datasplit

#### 4.2.1 Assessment of classification performances

The designed model's categorization efficiency was determined by estimating the confusion matrix. Figure 7 displays the confusion matrix of the presented technique. It is a tabular representation of the classifier's efficiency in categorizing lung cancer by comparing the classified outcomes with the real class. It also allows us to estimate the model's classification results such as accuracy, precision, recall, etc., through four cells namely: true positive (TP), true negative (TN), false positive (FP), and false negative (FN).



**Figure 9.** Statistical significance analysis: (a) 80/20 data split, and (b) 70/30 data split



**Figure 10.** Comparison of convergence curve of different optimization models

These cells evaluate the system's efficiency in correctly categorizing lung cancer and healthy instances from the images. TP indicates the case where it exactly classifies the lung cancer types, while TN defines the case where the system accurately detects and categorizes healthy images. In contrast, FP represents the case where the system incorrectly classifies the negative case as positive and FN denotes the case where it incorrectly categorizes the actual positive as negative.

On the other hand, receiver operating characteristic (ROC) is the graphical illustration of the system's classification results. It demonstrates how effectively the developed SWoCNN model categorizes lung cancer and healthy images across all possible thresholds. It is a plot of the TP rate against the FP rate, and it emphasizes the model's outcome by estimating the area under the curve (AUC). If AUC is close to 1, it indicates higher accuracy and vice versa.

Figure 8 displays the ROC curve of the presented model for 80/20 datasplit and 70/30 datasplit. It is observed that for 80/20 data split, the designed approach achieved AUC of 0.95, while for 70/30 data split it obtained AUC of 0.92. This higher AUC depicts that the designed framework precisely categorizes normal and lung cancer instances from the CT images.

Furthermore, to validate the SWO optimization model, its

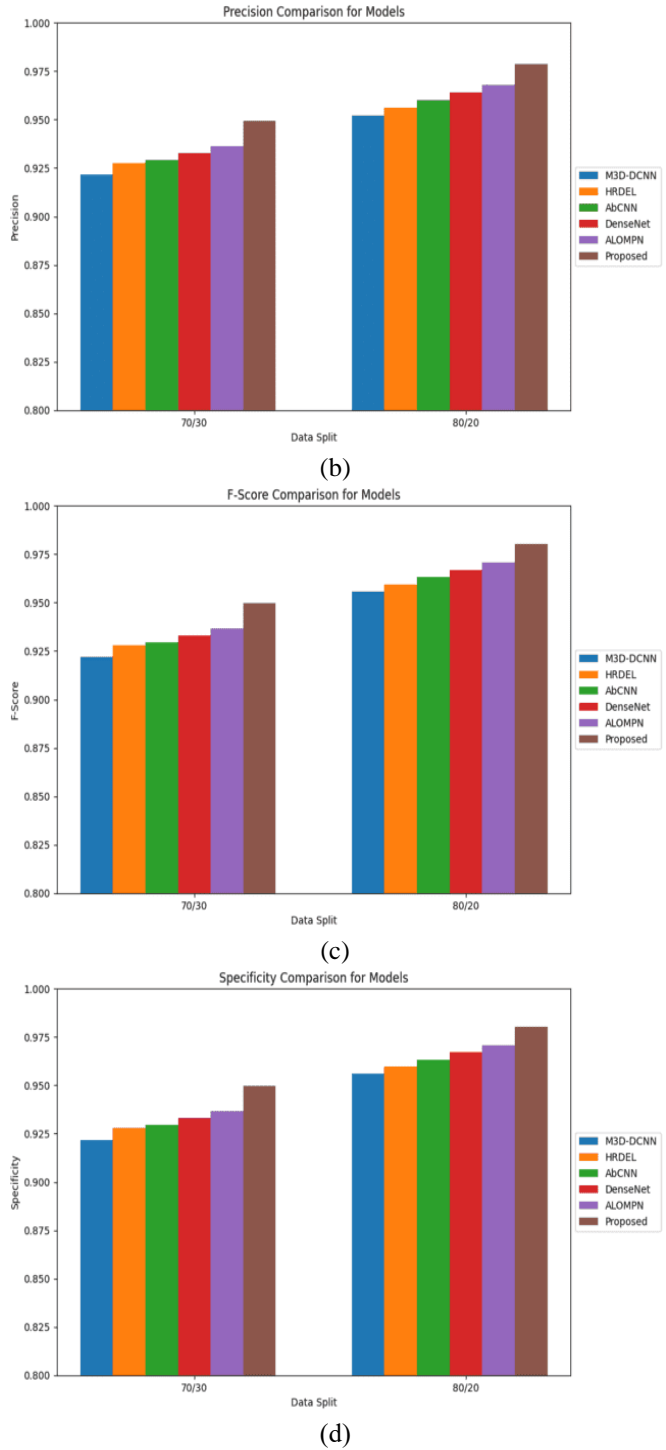
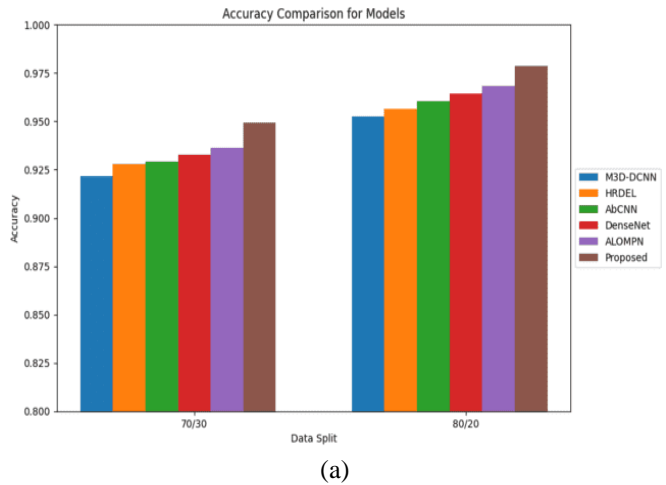
convergence rate was compared with other optimization algorithms such as GA, PSO, GWO, and firefly optimization algorithm (FOA). The convergence rate quantifies how quickly the optimization algorithm converges to the optimal solution set per generation in the decision space (Figure 9).

Figure 10 presents the comparative assessment of convergence curve of different optimization algorithms. Compared to other meta-heuristic optimization models, the proposed SWO approach achieved faster and highly stable convergence by reaching the best fitness score (lower training loss). This illustrates that utilizing SWO for CNN parameter fine-tuning results offer effective exploration and exploitation of search space and finds best value for each parameter.

### 4.3 Comparative assessment

The performances achieved by the developed SWoCNN model were compared and evaluated with the conventional lung cancer classification algorithms such as Ant lion Optimized Multilayer Perception network (ALOMPN) [33], DenseNet [34], Attention-based CNN (AbCNN) [35], High ranking deep ensemble learning (HRDEL) [36] and Multilevel 3D Deep CNN (M3D-DCNN) [37].

The metrics used in comparative analysis include accuracy, precision, FPR, f1-score, FNR, recall, computational time, and specificity, and these metrics are determined under two cases: 70/30 data split and 80/20 data split. Figures 11(a-d) present the comparative assessment of accuracy, precision, f1-score, and specificity. For 80/20 data split, the developed model and the existing techniques such as M3D-DCNN, HRDEL, AbCNN, DenseNet and ALOMPN obtained accuracy of 0.97872, 0.95238, 0.95634, 0.96436, and 0.96436, respectively, while for 70/30 dataset, these techniques earned accuracy of 0.94942, 0.92157, 0.92781, 0.93285, and 0.93629. This comparative analysis manifests that the developed methodology achieved improved accuracy than the conventional models, highlighting its proficiency in categorizing lung cancer from CT images. Similarly, the precision metric was compared with the existing models. The conventional classifiers and the presented SWoCNN achieved a precision of 0.95198, 0.95597, 0.96, 0.96406, 0.96815, and 0.97854, respectively for an 80/20 data split. On the other hand, they attained precision of 0.92153, 0.92773, 0.92930, 0.93282, 0.93624, and 0.94937, respectively for a 70/30 data split. The improvement of precision in the designed framework highlights its proficiency in recognizing the pattern of different lung cancer categories.



**Figure 11.** Performance comparison: (a) accuracy, (b) precision, (c) f1-score, and (d) specificity

Consequently, the designed strategy and the traditional models including M3D-DCNN, HRDEL, AbCNN, DenseNet and ALOMPN achieved f1-score of 0.98024, 0.95568, 0.95938, 0.96311, 0.96686, and 0.97065, respectively for 80/20 data split, while they earned f1-score of 0.94973, 0.92182, 0.92799, 0.92971, 0.93311, and 0.93682, respectively for 70/30 data split. The increased f1-score illustrates that the developed algorithm provides a balanced classification of healthy and lung cancer classes. The specificity of the model was also estimated and compared to the existing models to validate the efficiency of the designed SWoCNN model in recognizing the healthy image from tumor classes. The above-mentioned algorithm conventional models

and the proposed framework achieved specificity of 0.95594, 0.95962, 0.96332, 0.96705, 0.97082, and 0.98035 for 80/20 data split, while they earned specificity of 0.92175, 0.92795, 0.92966, 0.93308, 0.93673, and 0.94971, respectively for 70/30 data split as shown in Table 2 and Table 3. The

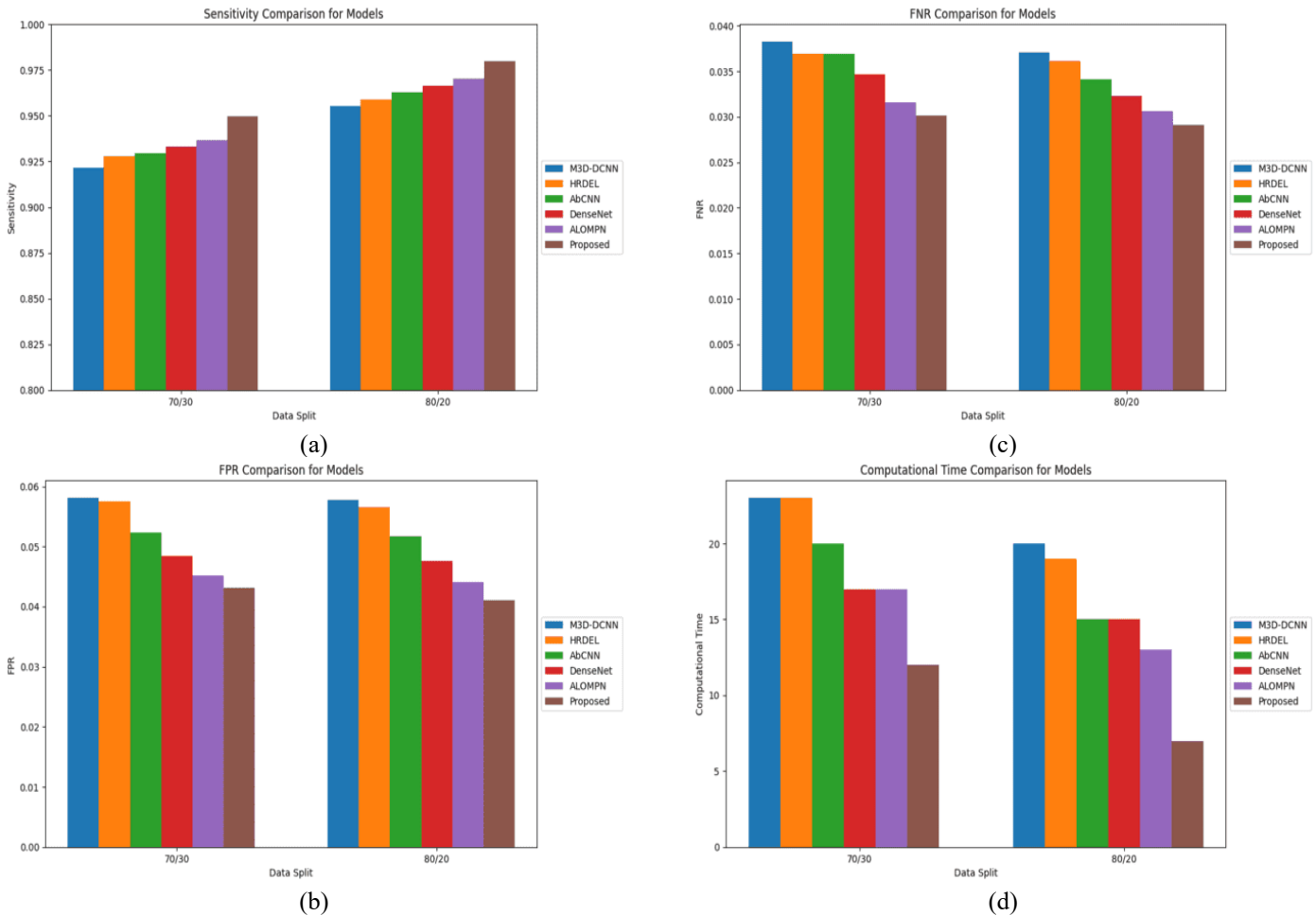
specificity comparison demonstrates that the developed model obtained greater specificity than the existing algorithms, which highlights its robustness in identifying and classifying normal images from the lung cancer-affected images.

**Table 2.** Comparative assessment of the developed model's performance with other techniques for an 80/20 ratio

Parameters	M3D-DCNN	HRDEL	AbCNN	DenseNet	ALOMP	Proposed
Accuracy	0.95238	0.95634	0.96033	0.96436	0.96842	0.97872
Precision	0.95198	0.95597	0.96	0.96406	0.96815	0.97854
F1-Score	0.95568	0.95938	0.96311	0.96686	0.97065	0.98024
Specificity	0.95594	0.95962	0.96332	0.96705	0.97082	0.98035
Sensitivity	0.95527	0.95900	0.96276	0.96655	0.97037	0.98005
FPR	0.05772	0.05660	0.05172	0.04762	0.04412	0.04110
FNR	0.03709	0.03614	0.03409	0.03226	0.03061	0.02913
Computational Time (Sec)	20	19	15	15	13	7

**Table 3.** Comparative performance measures of the proposed model with other techniques

Parameters	M3D-DCNN	HRDEL	AbCNN	DenseNet	ALOMP	Proposed
Accuracy	0.92157	0.92781	0.92934	0.93285	0.93629	0.94942
Precision	0.92153	0.92773	0.92930	0.93282	0.93624	0.94937
F1-Score	0.92182	0.92799	0.92971	0.93311	0.93682	0.94973
Specificity	0.92175	0.92795	0.92966	0.93308	0.93673	0.94971
Sensitivity	0.92172	0.92792	0.92963	0.93307	0.93669	0.94965
FPR	0.05809	0.05749	0.05227	0.04846	0.04524	0.04312
FNR	0.03824	0.03692	0.03690	0.03472	0.03160	0.03013
Computational Time (Sec)	23	23	20	17	17	12



**Figure 12.** Performance comparison: (a) sensitivity, (b) FPR, (c) FNR, and (d) computational time

The comparative assessment of metrics such as sensitivity, FPR, FNR, and computational time are graphically presented in Figure 12. The sensitivity is a crucial parameter that determines the classifier's effectiveness in recognizing all correct (lung cancer instances) from the lung cancer cases that occurred. The presented methodology and the above-stated conventional models obtained a sensitivity of 0.94965, 0.92172, 0.92792, 0.92963, 0.3308, and 0.93673, respectively for 70/30 data split, while they achieved 0.98005, 0.95527, 0.95900, 0.96276, 0.96655, and 0.97037, respectively for 80/20 data split. The increased sensitivity in the designed model validates its effectiveness in identifying all lung cancer categories from both true and false positives. On the other hand, the FPR and FNR metrics are determined to estimate the error rate of the classifier. The designed strategy achieved lower FPR and FNR of 0.04312 and 0.03013 for the 70/30 data split, while it earned an FPR of 0.04110 and FNR of 0.02913 for the 80/20 data split. Subsequently, the above-mentioned conventional models achieved FPR of 0.5809, 0.05749, 0.05227, 0.04846 and 0.04524, and FNR of 0.03824, 0.03692, 0.03690, 0.03472, and 0.03160, respectively, for 70/30 data split. Also, they earned FPR of 0.05772, 0.05660, 0.05172, 0.04762, and 0.04412, and FNR of 0.03709, 0.03614, 0.03409, 0.03226, and 0.03061, respectively for 80/20 data split. The lower FPR and FNR in the developed algorithm highlight that it correctly classifies the healthy and lung cancer classes compared to existing models. Finally, the execution time is also determined to manifest the computational efficiency of the model. The presented algorithm consumed a minimum computational time of 7s and 12s for 80/20 and 70/30 data splits, while the existing models have taken longer computational times of the 20s, 19s, 15s, 15s, and 13s, respectively for 80/20 data split and 23s, 23s, 20s, 17s, and 17s, respectively for 70/30 data split. The reduced time consumption illustrates that the presented model quickly learns the feature representations and performs classification tasks. This also highlights that the parameter optimization by SWO minimizes the computational demands and training time. Tables 2 and 3 list the performance validation of the existing models with the presented approach for 80/20 and 70/30 data split.

#### 4.4 Discussion

A hybrid classification strategy was developed to detect and classify the types of lung cancer from the CT images. The proposed model incorporates the optimization efficiency of SWO into the CNN to boost its adaptability and training speed through optimal parameter value selection. The presented work used the "Lung cancer CT scan database" from the Kaggle site as input and the images are preprocessed through steps like noise removal, image resizing, and background elimination. Also, the U-Net algorithm was developed to segment the ROI from the preprocessed images and the relevant features are captured using ANN. Consequently, a WPA-based feature selector was proposed to select the most informative and highly relevant attributes, assisting the classifier to focus on major features influencing the patterns of lung cancers. Finally, an optimized CNN was designed to perform the classification task. The CNN performs feature learning and estimates the lung cancer patterns by extracting spatial and local features. The SWO integrated into CNN refines its parameters continuously in an iterative manner, enabling the classifier to adapt to the evolving variations in

input images and boosting the training speed through optimal parameter selection.

The robustness of the designed methodology was validated by determining the results in two cases: 80/20 and 70/30 data splits and the experimental results highlight that the designed methodology earned improved outcomes. Furthermore, a comparative assessment was performed with conventional classification models such as M3D-DCNN, HRDEL, AbCNN, DenseNet, and ALOMPN and it demonstrated that on average the performances such as accuracy, precision, recall, f1-score, and specificity are increased by 0.01172, 0.01176, 0.01132, 0.01132, and 0.01126 in the designed model. On the other hand, the metrics such as FPR, FNR, and computational time are reduced by 0.00257, 0.00123, and 5.5s in the presented methodology. The comparative analysis highlighted that the designed strategy performs better lung cancer classification than the currently available classifier model, making it a reliable solution to the real-time medical industry for early and precise lung cancer classification.

## 5. CONCLUSIONS

A lung cancer classifier SWoCNN was designed by combining the advantages of SWO into the CNN. The major objective of the research is to categorize the various types of lung cancer from the CT images using optimized CNN. The CT images are gathered from healthy and lung cancer-affected persons and preprocessed to make them suitable for further processes. Consequently, image segmentation and feature extraction are done using U-Net and ANN to capture ROI and relevant features from the preprocessed CT scans. A feature selector was designed using WPA to optimally select the most relevant and informative attributes from the extracted sequences. Finally, the proposed SWoCNN classifier was trained using the selected features to classify healthy and lung cancer categories such as adenocarcinoma, LCC, and SCC. The implementation outcomes of the designed strategy demonstrate that it achieved an accuracy of 0.94942 and 0.97872 for lung cancer classification on 70/30 and 80/20 data split. Furthermore, the developed approach earned a minimum execution time of 7s and 12s, demonstrating its computational efficiency. Finally, the comparative assessment with the existing techniques such as M3D-DCNN, HRDEL, AbCNN, DenseNet, and ALOMPN demonstrated that the designed framework achieved better outcomes than others.

Although the designed model achieved improved results, it has several limitations. Firstly, the presented strategy is limited to binary classification; it doesn't classify the sub classes or different grades of lung cancer. Secondly, the developed framework is not validated across diverse databases, which limits its scalability across different clinical scenarios. Thirdly, the study has not focused on predicting the survival rate of cancer-affected individuals, which is crucial for preparing treatment plans. Hence future work focus on testing the scalability of the model and extending it to classify different lung cancer grades and predicting the survival rate of the patients.

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