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# Optimization-based convolutional neural model for the classification of white blood cells

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## Abstract

White blood cells (WBCs) are one of the most significant parts of the human immune system, and they play a crucial role in diagnosing the characteristics of pathologists and blood-related diseases. The characteristics of WBCs are well-defined based on the morphological behavior of their nuclei, and the number and types of WBCs can often determine the presence of diseases or illnesses. Generally, there are different types of WBCs, and the accurate classification of WBCs helps in proper diagnosis and treatment. Although various classification models were developed in the past, they face issues like less classification accuracy, high error rate, and large execution. Hence, a novel classification strategy named the African Buffalo-based Convolutional Neural Model (ABCNM) is proposed to classify the types of WBCs accurately. The proposed strategy commences with collecting WBC sample databases, which are preprocessed and trained into the system for classification. The preprocessing phase removes the noises and training flaws, which helps improve the dataset's quality and consistency. Further, feature extraction is performed to segment the WBCs, and African Buffalo fitness is updated in the classification layer for the correct classification of WBCs. The proposed framework is modeled in Python, and the experimental analysis depicts that it achieved 99.12% accuracy, 98.16% precision, 99% sensitivity, 99.04% specificity, and 99.02% f-measure. Furthermore, a comparative assessment with the existing techniques validated that the proposed strategy obtained better performances than the conventional models.

**Keywords:** Convolutional neural network, White blood cells, African Buffalo Optimization

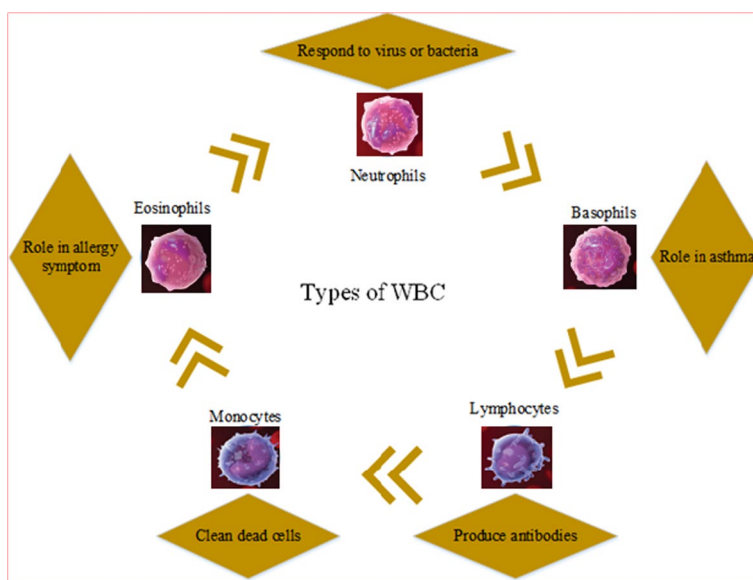
## Introduction

Generally, WBC is also called a white corpuscle or leukocyte; the functions of WBC contain a nucleus capable of mobility and defending the human body from disease and infection by feeding cellular debris and foreign materials by creating antibodies [1]. Besides, WBC produces multipotent cells in the bone marrow, also called hematopoietic stem cells [2]. Furthermore, the quantity of WBC in blood indicates the disease; WBC count is the most important subset for completing blood count [3]. Additionally, the average WBC count is approximately 1% in the blood, significantly changing health since immunity depends on the WBC [4, 5]. The different types of WBC play different jobs in our body, such as killing harmful bacteria, recognizing intruders, create antibodies to

protect the body against viruses and bacteria [6, 7]. The types of WBC are detailed in Fig. 1.

Generally, WBC begins in the bone marrow, and every blood cell descends after the Hematopoietic Stem Cell (HSC) [8]. Initially, HSC cells are isolated into a myeloid or lymphoid cell, which provides rise to the lymphoid cell line; the family of cells creates T and B cells [9]. As well, the infection affects elevated WBC in terms of WBC count, which is increased by overproduction. Some reasons to increase WBC counts are pregnancy, infections, cancers, asthma, inflammation, allergy, exercise, and Trauma [10, 11]. WBC plays an important role in the immune system. It contains different types for performing the various functions [12]. Also, WBC protects against parasites, viruses, and bacteria [13]. Likewise, a high WBC count causes infection and illness, and a low WBC count causes vulnerable serious infections such as chemotherapy [14, 15].

Hence, the accurate detection and classification of WBCs is significant for effective disease diagnosis and treatment planning. The traditional way of WBC classification relies on manual analysis by trained clinical experts, which is typically time-consuming and prone to errors. These issues in traditional approaches demand an automatic WBC classification strategy. Consequently, the researchers focused on developing a classification mechanism using advanced data analysis tools, including artificial intelligence approaches such as machine learning and deep learning [16]. The AI-based classification models have shown more potential in automating the detection and classification of WBCs compared to traditional approaches regarding accuracy and precision. These models learn by analyzing the complex patterns and relations within the input blood sample images, enabling them to identify the difference between various types of WBCs more effectively [17]. However, AI-based models face significant issues like overfitting, lack of generalizability, data complexity, and high computational time. To address these issues, several types of research are developed by using deep learning techniques such



**Fig. 1** Types and function of WBC

as Capsule network-based deep learning [20], CNN using image processing [21], computer-aided automated system [22], Canonical Correlation Analysis (CCA) [23], and WBC-Net based ResNet and Unet++ [24] but still having the problem of less successive rate, less prediction accuracy and high error rate [18]. Furthermore, current studies develop techniques like advanced convolutional neural networks [36], Deep neural networks [37], efficient lightweight CNN [38], etc., for detecting and classifying different kinds of WBCs. These models offer improved precision and accuracy than the conventional models. However, they cannot handle the dynamic variations in the input samples, leading to suboptimal performances and reduced efficiency in real-world scenarios. To resolve these issues, a hybrid classification strategy is proposed in the presented work using an optimized deep learning model [19].

The classification of WBC includes three main processes: data preprocessing, feature extraction, and classification [20]. Initially, microscopic input images are trained to the system, and the input images contain small particles such as noise and errors in the staining process [21]. Those errors are removed in the preprocessing layer using suitable filters. Furthermore, features are extracted based on morphological operation, a relationship of colors, and granularity features [22]. Also, extracted features are updated for CNN to classify WBC [23]. However, there are some issues during classification, such as less detection accuracy, high execution time, and errors [24]. The main motive of this current work is there is no proper system to detect and classify WBC because the classification of WBC is essential for identifying health conditions [25]. CNN is also used to classify WBC and attain accurate results, but it contains some issues like noisy errors, less accuracy, and high False Positive Rate (FPR) [26]. Moreover, many developed techniques only involve the process of feature extraction and classification [27]. So, accurate identification and detection of WBC is more risky. These reasons motivated this present research towards designing the optimized deep learning model.

This article is structured as follows: Sect. "Related works" details the related work based on WBC classification. Sect. "Proposed methodology" describes the process of the proposed methodology. Sect. "Results and discussion" mentions the achieved outcomes. Finally, Sect. "Discussion" concludes with a conclusion and future work about the developed model.

### **Related works**

Yusuf Yarg and Umit Atila [28] have developed a Capsule network-based deep learning framework for improving the accuracy rate of the data testing process. Thus, the developed capsule network attains better results, such as high accuracy during training data, and the gained accuracy rate is 96.86%. However, it takes more time to classify the WBC because of the data complexity.

Nowadays, image processing is used in biomedical systems to analyze various diseases. Burhanet et al. [29] have proposed a convolutional neural system using image processing methods to improve classification accuracy performance in WBC. Moreover, maximal information coefficient and ridge feature selection are used to extract the relevant features. Finally, the proposed technique obtains a 97.95% successive rate to classify WBC but has the problem of overfitting.

Huseyin et al. [30] developed a computer-aided automated system technique to locate and identify WBC types. It involves five processes: preprocessing, feature extraction, segmentation, feature selection, and classification. Moreover, the developed Convolutional Neural Network (CNN) has extracted the features based on the object by object recognition. It shows better performance in accuracy, yet it has taken a high execution time to classify the WBC.

WBC is used to detect human health conditions related to blood disease. It consists of characterization and identification of sample blood from the patient. Patil et al. [31] developed Connected Component Analysis (CCA) to overcome the problem of multiple cell overlap. Moreover, extracting multiple nuclei patches from the dataset using CCA has reduced classification time, but the false prediction rate is high compared to other techniques.

Yan Lu et al. [32] developed deep learning WBC-Net-based ResNet and Unet++ techniques. Initially, the designed WBC-Net extracts the multi-scale features based on residual blocks of encoder context-aware feature. Furthermore, the WBC will be segmented based on extracted features, and decoder deconvolution and convolution incorporation will be used to refine the segmentation of the WBC mask. However, the developed technique does not perform a classification process, and the classification of WBC is essential to identifying the types of WBC. Some recent work related to the proposed technique is detailed in the Table 1.

WBCs are a significant part of the immune system, which protects our human body against infectious diseases and foreign invaders. Ahmed et al. [33] developed a hybrid technique for improving the classification of Leukemia. First, the features from the WBC images will be extracted using the salp swarm algorithm. Thus, the developed algorithm identifies and selects the relevant features and also removes the noisy and

**Table 1** Literature survey of medical image analysis

Author	Technique	Advantage	Disadvantage
Yusuf Yarg and Umit Atila [28]	Capsule network-based deep learning framework	High rate of accuracy	High time consumption
Burhan et al. [29]	Convolutional neural system using image processing methods	Enhance medical image processing system	Overfitting
Huseyin et al. [30]	Computer-aided automated system technique	Attain a high rate for classifying WBC	Complex in Design
Patil et al. [31]	Canonical Correlation Analysis	Enhance the sensitivity	Because of the large size of the data, it takes more time to process
Yan Lu et al. [32]	Deep learning WBC-Net-based ResNet and Unet++ technique	The best score to validate the type of WBC	Not supported for noisy sample data
Sahlol, Ahmed T et al. [33]	Salp swarm algorithm model	Attain high accuracy and reduce computational complexity	Misclassification problems
Quan et al. [34]	Deep learning based on CNN	High rate of accuracy	High error rate
Shahin, Ahmed Ismail et al. [35]	End to End CNN system	Better classification accuracy	High computational complexity

highly correlated features. The developed technique attains high accuracy and reduces computational complexity but has missed classification problems.

Quan et al. [34] proposed a CNN-based classification network for improving the classification performance of Red Blood Cells (RBC). Consequently, an attentive, dense circular net is designed in CNN to enhance performance. Additionally, classification is processed using feature sets, and the experimental results are compared with other DL techniques. Finally, the network attains 94.61% accuracy, but the error rate is high.

Nowadays, transfer learning techniques are utilized to complete the identification of WBCs in blood smear images. Shahin et al. [35] developed an end-to-end CNN system for the identification of various types of WBCs. Moreover, the developed design improves the classification of WBCs by attaining better classification accuracy. However, computational complexity is high due to the vast amount of data.

### **Motivation**

WBCs play a significant role in the human immune system by diagnosing pathological conditions such as infections, immune system disorders, etc. Accurate identification and classification of WBCs is important for effective diagnosis of health conditions and proper treatment planning. The conventional way of WBCs classification involves a manual examination of blood samples by trained healthcare professionals. However, this traditional methodology is time-consuming, prone to human error, and demands significant expertise. These drawbacks of the traditional method create a demand for automatic classification mechanisms. The evolution of Artificial Intelligence (AI) techniques in the medical field has paved the way for automatically classifying WBCs. The AI techniques, including deep learning and machine learning approaches, have offered greater WBC classification accuracy than traditional methods.

The AI models are trained using the blood sample images using supervised or unsupervised learning mechanisms. These learning processes enable the AI models to learn the pattern difference between the different types of WBCs, allowing them to classify more accurately. Although various AI-based models have been developed, including deep neural networks, artificial neural networks, support vector machines, etc., they face certain challenges. Firstly, training the AI-based models is complex and time-consuming due to the vast data required for effective training. Secondly, the existing models cannot handle dynamic changes in the blood sample images, leading to inaccurate classifications. Moreover, the overlapping of cells, noise features, etc., in the images makes the existing techniques ineffective, leading to suboptimal classification performance. In addition, scalability and interpretability are the other challenging issues in the existing techniques. These challenges make the existing models ineffective for real-time WBC classification.

Hence, to address these issues, an innovative solution is proposed in this study by leveraging the benefits of the African Buffalo Optimization algorithm and Convolutional Neural Network. The novelty of the approach lies in the seamless integration of deep learning and meta-heuristic optimization algorithms for WBC classification. The adaptive nature of the ABO approach enables the system to understand the dynamic variations in cell appearance, providing optimal classification results. In addition, the integration of ABO into CNN minimizes the computational efforts, thereby lowering the

training time and resource demands. Moreover, iterative updation of ABO fitness in the classification layer enhances the scalability and reliability of the approach in WBC classification. The model upgrades its fitness and parameters at each iteration, enabling the system to handle more images and ensuring scalability and reliability in real-world scenarios. The major contributions of the proposed work are described below.

1. This study proposes a unique solution for accurately classifying different types of WBCs by combining the efficiency of African Buffalo Optimization and Convolutional Neural Network.
2. The convolutional layer of CNN performs image preprocessing and feature extraction, which improves the accuracy of the classification process by reducing noise and retaining the most significant features.
3. The fitness of the ABO is integrated into the pooling layer of the CNN approach, which enables the system to segment WBCs by detecting variations in pixel values, leading to better classification performance.
4. Finally, the fully connected layer of the CNN performs WBC classification by understanding the patterns and interconnections within the segmented images.
5. The proposed strategy is implemented in the Python tool, and the results are determined and validated with the existing techniques in terms of accuracy, precision, sensitivity, specificity and f-measure.

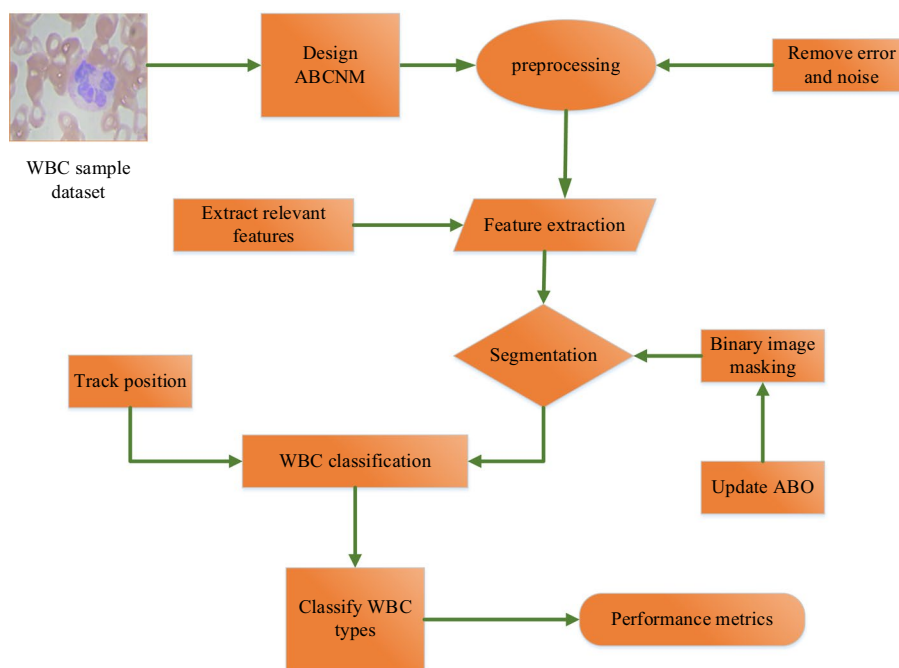
### **Proposed methodology**

Generally, White Blood Cells (WBC) provide more relevant information about the health status of human beings. Furthermore, the ratio of cells and deformation shows the diagnosis process of some diseases. Therefore, correct identification and classification of WBC type are essential. The WBC sample image dataset is collected from the net source and trained in the system in this work. Hereafter, a novel African Buffalo-based Convolutional Neural Model (ABCNM) is designed with suitable parameters. Immediately, the training errors are preprocessed in the preprocessing frame, and feature extraction is executed for the segment of the WBC cells. Followed by, extracted features are sent to the classification layers to classify the WBC cells. The proposed architecture is shown in Fig. 2.

Here, the best solution for the African buffalo is upgraded in the classification layer to identify the type of WBC cells. Consequently, the segmentation process is performed, and the WBC type is predicted successfully. Thus, the developed framework enhances the classification accuracy of WBC and also performs better in sensitivity and specificity.

### **Dataset description**

The presented study used the publicly available blood sample images dataset from Kaggle, accessible at <https://www.kaggle.com/datasets/paultimothymooney/blood-cells>. This dataset contains blood sample images from four types of leukocytes: Neutrophil (NE), Eosinophil (EO), Lymphocyte (LM), and Monocyte (MN). It contains a total of 12,444 images, with each leukocyte type indicated as follows: 3144 images of Eosinophils, 3139 images of Lymphocytes, 3132 images of Monocytes, and 3171 images of Neutrophils. This database was split into ratios of 80:20 for the training and testing



**Fig. 2** Proposed methodology

**Table 2** Dataset details

WBC types	Number of images in each class	Training images (80%)	Testing images (20%)
Neutrophil	3,171	2,537	634
Eosinophil	3,144	2,515	629
Lymphocyte	3,139	2,511	628
Monocyte	3,132	2,512	620
<b>Total</b>	<b>12,444</b>	<b>9,955</b>	<b>2,489</b>

process. 80% of the images were used for model training, while 20% of the data was used for model testing. The size of the image present in the dataset is  $256 \times 192$  pixels. Table 2 presents the details of the dataset.

**Process of ABCNM**

Generally, ABCNM is developed to classify the WBC in blood samples. Moreover, the fitness function of AB is updated to the developed CNN, and it contains five layers: input layer, convolutional layer, pooling layer, fully connected layer, and output layer. Furthermore, African Buffalo Optimization (ABO) is based on searching the location of each buffalo depending upon the two sounds, such as the 'maa' and 'waa' sound. Additionally, the maa sound denotes the buffalo's to a safe place, and the waa sound denotes the search place for food. Based on the sound location, buffalos can search for food. The purpose of using AB fitness in the classification layer is to search the types of WBC present in the blood sample using their fitness function and also classify the types of WBC. Thus, the fitness function is denoted as  $f(t_1)$  and  $f(t_2)$  where,  $f(t_1)$  it is used to segment the WBC cells based



on extracted features and  $f(t_2)$  is used to classify WBC type. The blood sample image is initially tested and trained to the system, and the collected dataset is imported to the developed ABCNM.

First, WBC sample images are trained to the system; trained datasets are updated to the input layer of developed ABCNM, as given in Eq. (1).

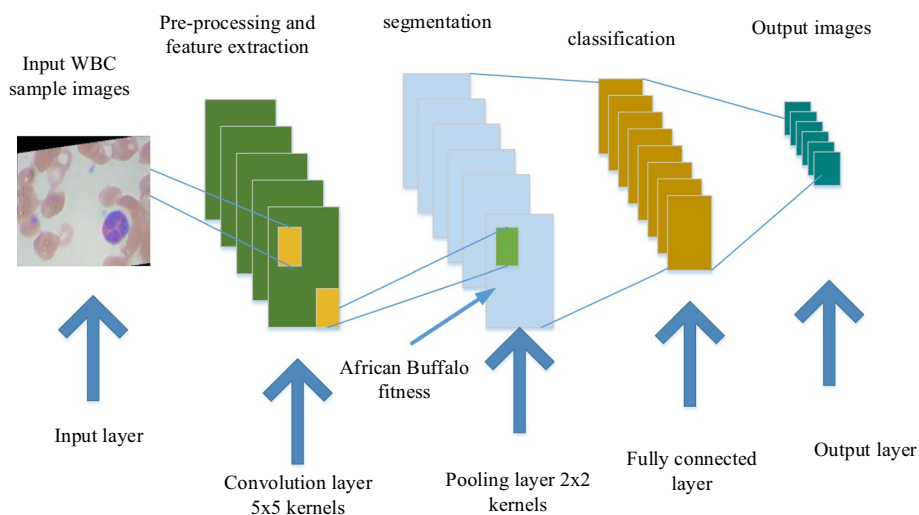
$$D(i) = k(n - 1), k(n - 2), k(n - 3), \dots, k(n - d) \tag{1}$$

where  $D(i)$  is denoted as the total quantity of updated datasets  $k(n - 1), k(n - 2), \dots$ , which are considered different types of WBC sample datasets. Figure 3 illustrates the ABCNM technique process.

- Preprocessing

Image preprocessing indicates converting the raw input data into an appropriate format for subsequent processing. In the proposed framework, the data preprocessing was performed in the convolutional layer of the CNN model. The input layer of the CNN forwards the input raw database into the convolutional layer for preprocessing. The preprocessing phase involves the removal of errors and noise from the raw blood sample images. The convolutional layer serves as a filter bank that applies various filters to the input image, which helps reduce the noises and other unwanted features from the input image. The errors, training flaws, and noise are filtered through convolutional operations. This is achieved by convolving the input image with specific kernels. In addition, the original input images were converted into grayscale images for effective and accurate classification of WBC. Moreover, a probability density function of preprocessing is obtained using Equation (2).

$$L(p) = \left[ L(o) \frac{k_o}{k_p} \right] \tag{2}$$



**Fig. 3** Process of ABCNM technique



where,  $L(p)$  is represented as probability density function of preprocessing and  $L(o)$  is denoted as an errors and noise presented in the dataset. Also  $k_o$  and  $k_p$  is considered as the input blood sample WBC image.

- Feature extraction

Feature extraction is an essential step in White blood cell classification. The objective of the step is to extract and capture the most relevant and significant attributes from the preprocessed blood sample images. In this phase, the meaningful features and patterns inherent in the data are identified and captured, which plays a crucial role in classifying different types of WBCs. This step reduces the dimensionality of the data and enhances the discriminative power of the classification model. In this process, the distinct features of WBC morphology, including color, shape, texture, size, length, and other spatial characteristics, are extracted through a series of convolution operations in the convolution layer. Additionally, feature extraction is obtained by Eq. (3).

$$K(w) = \frac{(K(w) - F_a) \times (F(e) - H_a)}{F_a - H_a} \tag{3}$$

where,  $K(w)$  is denoted as feature extraction of collected WBC blood samples and  $F_a$  is represented as redundant information of the WBC blood sample. Moreover,  $H_a$  it is represented as relevant information about WBC blood samples like size, shape, color, etc.

- Segmentation

In the pooling layer, the fitness function of ABO is updated for detecting WBC. Moreover, this layer segments and classifies WBC types using the fitness function of AB. In this technique, segmentation is done using mask images, and masking is used to identify the WBC types using color, size, shape, and length features. It helps classify variation among pixels in the segmented region. It can perform parallel prediction on the WBC blood samples. Moreover, the large image can be modified with the help of a binary image because binary images contain zero and non-zero values. Where the gained value is 1 in a binary mask, it is a real value, but if the value is 0 in a binary mask, it means null. Thus, the process of segmentation is obtained by Eq. (4).

$$N_s(h) = \frac{b(t) \times K(w)f(t_1) \times D(i)}{2} \tag{4}$$

where,  $N_s(h)$  is called the segmentation of WBC blood samples and  $b(t)$  is denoted as a mask image. Moreover,  $f(t_1)$  is denoted as fitness function segmentation based on extracted features. Moreover, WBC types are classified using the proposed ABCNM as detailed in Algorithm.1.

**Algorithm: 1**

**Input:** Blood sample images;  
**Output:** Classified WBC type;  
**Start {**

Initialize the input database; *//Import the dataset into the Python system*

Design CNN architecture;

Initialize the CNN parameters and define the CNN layers (convolution, pooling, and fully connected layer);

Preprocess the raw dataset; *// Remove errors and noise from the raw input images*

Feature extraction; *//Extract and capture the most significant features*

Segmentation:

Initialize the ABO parameters (maximum iteration, population);

Initialize the mask and binary image;

For each image in the dataset:

Update fitness function of ABO;

Perform segmentation using the updated fitness;

Detect variation among pixels in the segmented region;

If  $(N, (h) \rightarrow 1)$  Segment WBC;

Else No segmentation;

End for;

Classification: *//fully connected layer*

For each segmented image:

Track the position of the cell in the image;

Perform WBC classification; *//Classify the image as Monocytes, Lymphocytes, Neutrophils, Eosinophils*

End for;

Evaluate model performances; *//determine accuracy, precision, f-measure, specificity, and sensitivity*

**} End**

- Classification

Then, the segmented datasets are sent to the fully connected layer, which searches for the correct position of WBC types based on the labels such as 0 and 1 and classifies the WBC types such as Monocytes, Lymphocytes, Neutrophils, and Eosinophils. It can

classify the WBC types based on all object binary images assigned in unique numeric values. Then, the binary image values are computed using maximum and minimum pixels for each label. Finally, various types of WBC present in the collected blood sample dataset are classified using Eq. (5).

$$T_p = N_s(h) \times \frac{D(i)}{K(w)} \times f(t_2) \tag{5}$$

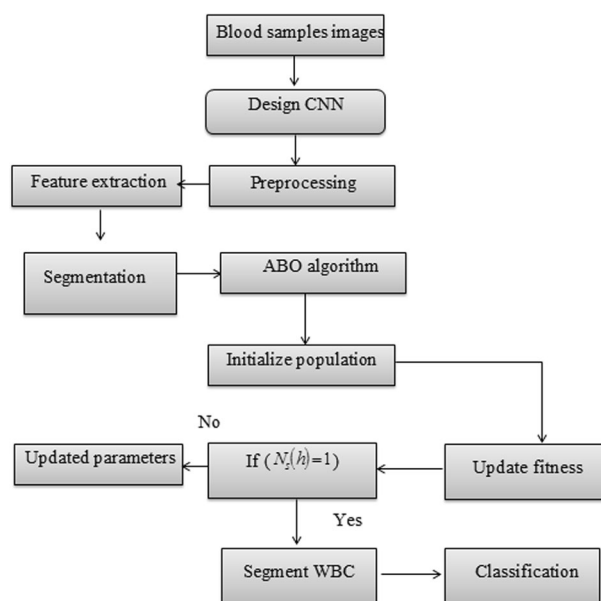
where,  $f(t_2)$  denotes the fitness function to classify WBC types from the collected blood samples. At last, the output layer provided the segmented output using a multilayer network and it classify WBC types based on the variation of pixels which is detailed in Fig. 4.

### Results and discussion

This study presented a unique solution for accurate classification of WBCs by combining the efficiency of ABO and CNN. The proposed strategy is modeled and executed in Python, version 3.7.8. Metrics like accuracy, precision, sensitivity, f-measure, specificity, and execution time determine the experimental results. Finally, a comparative assessment was made with the conventional models to validate the effectiveness of the proposed algorithm. Table 3 tabulates the implementation parameters and its description.

### Case study

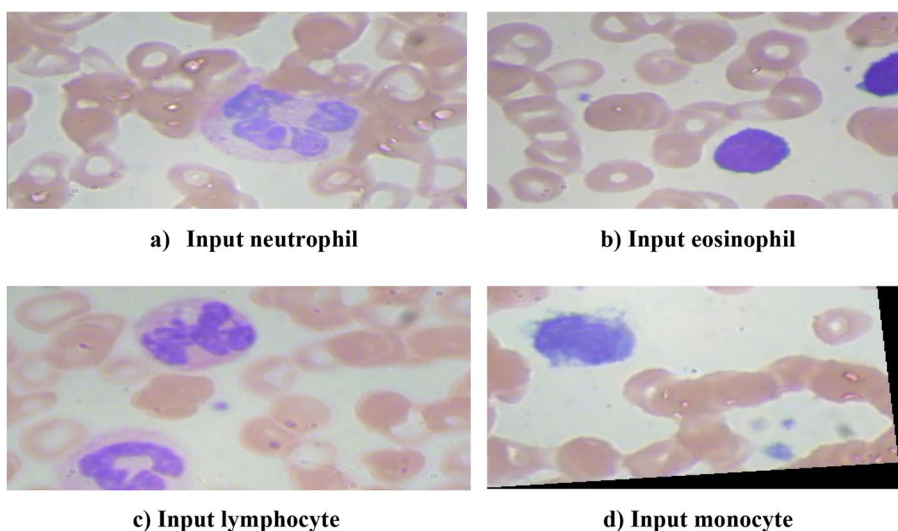
Classification of WBC is the most common and crucial issue in humans, and our bodies protect it against infections by eliminating bacteria, viruses, fungi, and parasites. The segmentation and classification of WBC is the most challenging task. Initially, blood sample datasets are trained for the system and updated to the proposed ABCNM to execute WBC-type preprocessing, feature extraction, segmentation, and classification.



**Fig. 4** Workflow of developed ABCNM

**Table 3** Implementation parameters and its description

Parameter	Description
Programming language	Python
Version	3.7.8
Operating system	64-bit Windows
CPU	CORE i7
Hard drive	2 Terabyte (TB)
Memory	8 Gigabyte (GB)
Library	TensorFlow, Keras, Scikit-learn



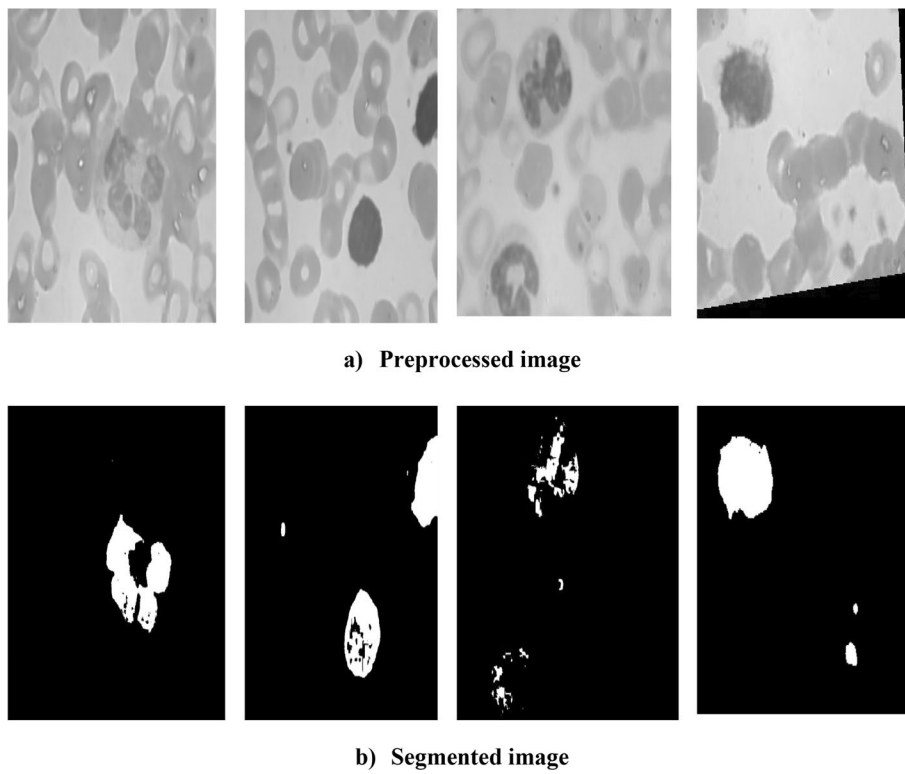
**Fig. 5** Input blood sample images

In this case study, several patient datasets with various WBC types are taken, and the input sample blood images of neutrophils, lymphocytes, monocytes, and eosinophils are mentioned in Fig. 5.

Consequently, preprocessing executed in the convolution layer removes noise and errors in the dataset. Moreover, the original input images are converted into greyscale images for segmentation. After that, feature extraction is processed to extract the relevant information in the WBC based on size, color, shape, etc.

- Segmentation

Next, in the Pooling layer, WBC in the blood samples are segmented with the help of fitness function AB of the proposed technique. Initially, mask the WBC using a binary mask. Then, segmentation happens based on the pixel values. Thus, the developed technique enhances the accuracy and performance of the classification of WBC types. Figure 6 (a, b) presents the preprocessed and corresponding segmentation images. The preprocessed images indicate the input image after applying steps like grayscale conversion, noise removal, and standardization, while the segmented image defines the outcome of the segmentation process. It indicates where the white



**Fig. 6** a Preprocessed image, b Segmented image

**Table 4** Classification of WBC type

SI.No	Classified WBC
a)	Neutrophil
b)	Eosinophil
c)	Lymphocyte
d)	Monocyte

blood cells are located in the blood sample images. The segmented images isolate the WBCs from the background and other cells.

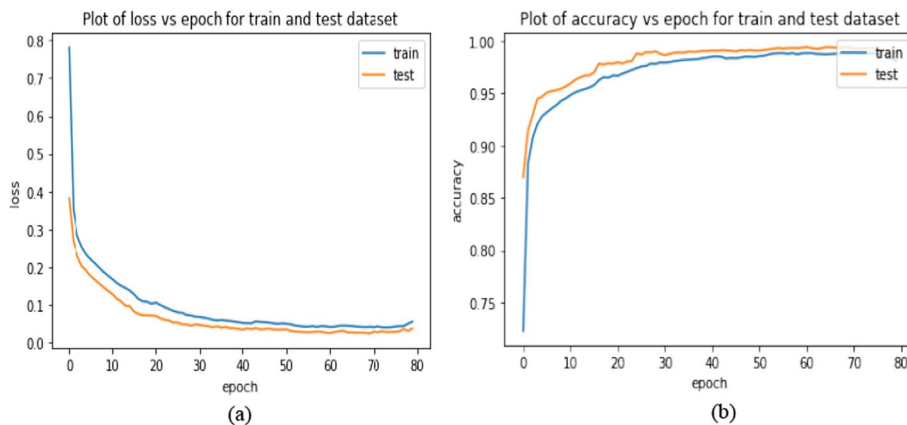
- Classification

With the help of a segmented image, developed ABCNM track the position of the WBC in blood sample based on labels 0 and 1. In the fully connected layer, detect the accurate position of WBC also classify the types of WBC based on the segmented results. Moreover, the classified WBC type is detailed in Table 4.

Finally, the type of WBC present in the sample blood dataset is identified and the calculation is made using segmented images from the developed ABCNM. Moreover, the developed technique performs outcomes for classifying WBC types.

**Table 5** ABCNM model parameters and its values

Hyperparameters	Search range	Optimal value
Dropout rate	[0.3, 0.4, 0.5, 0.6, 0.7, 0.8]	0.5
Learning rate	[0.1, 0.01, 0.005, 0.001, 0.0005]	0.001
Batch size	[8, 16, 32]	32
Kernel size	[3 × 3, 5 × 5, 7 × 7]	3 × 3
Number of filters	[16, 32, 64]	32



**Fig. 7** Training and testing performance: **a** accuracy, **b** loss

**Model training and testing**

This subsection assesses the proposed model’s training and testing performance by increasing the epoch. Firstly, the input database is split into ratios of 80:20 for the training and testing phases, and the performances were assessed in terms of accuracy and loss. Table 5 tabulates the implementation parameters and their values.

The accuracy metric measures the effectiveness of the developed model in identifying the pattern difference between the different types of WBCs. In contrast, the loss metric determines the misclassification made by the model in WBC classification. Figure 7 (a, b) presents the training and testing performance of the developed model. The training accuracy demonstrates how quickly the proposed model understands the patterns and relations influencing the types of WBCs, and the improvement of the model’s training accuracy over increasing epochs highlights that the designed strategy effectively learns different WBC patterns. On the other hand, testing accuracy indicates the model’s generalization efficiency over the unseen data samples (data not used for model training). The enhancement of the model’s testing accuracy over epochs highlights that the presented model identifies and classifies the different types of WBCs precisely on unseen data.

Consequently, the training loss quantifies the deviation between the actual and the predicted outcomes within the training sequence. It measures how effectively the proposed model fits into the training data. The testing loss defines the difference between the predicted and actual outcomes on real-world data (unseen data). It measures how effectively the developed algorithm applies the learned information on unknown data and predicts

the WBC types. Reducing training and testing accuracy highlights the model's learning efficiency and reliability across real-world scenarios.

The confusion matrix is a tabular representation of the performance of the classification model. It enables us to understand how effectively the classification model differentiates the different types of WBCs: NE, EO, LM, and MN. It evaluates the classification performance using four different metrics, namely: True Positive (TP), False Positive (FP), True Negative (TN), and False Negative (FN). TP indicates the scenarios where the model correctly identifies the specific class of WBC, while FP defines scenarios where the model incorrectly classifies the specific class of WBCs. TN represents the case where the model accurately predicts the absence of the specific WBC class. At the same time, FN indicates the scenario where the model inaccurately identifies the absence of a specific WBC class. Figure 8 presents the confusion matrix of the proposed strategy.

**Performance metrics**

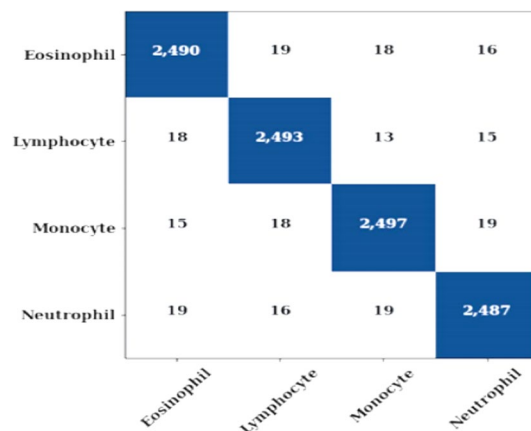
The implementation work of the developed ABCNM is done using the Python tool, and parameters like accuracy, sensitivity, specificity, execution time, F1-measure, and precision are calculated. Moreover, the developed approach is validated using existing methods like the Classification of WBC using Capsule Networks (CN) [28], the Classification of WBC using CNN-based Feature Selection Technique (FST) [29], WBC detection and classification-based Regional CNN (RCNN) [30], WBC image classification using Canonical Correlation Analysis (CCA) [31], and WBC classification based Optimized Swarm (OS) [33].

**Accuracy**

Accuracy is defined as the degree of calculation of the efficiency of the proposed model functioning. Also, it is the fraction of properly expected observance of all the observations that are expressed in Eq. (6),

$$A = \frac{T_p + T_n}{T_p + T_n + F_p + F_n} \tag{6}$$

where,  $T_p$  is denoted as accurate prediction and exact segmentation,  $T_n$  is denoted as accurate prediction and incorrect segmentation,  $F_p$  is considered as inaccurate



**Fig. 8** Confusion matrix



prediction and exact segmentation, and  $F_n$  is denoted as inaccurate prediction and incorrect segmentation of WBC. Moreover, accuracy comparison with the existing technique is detailed in Fig. 9. Generally, increasing the number of samples helps to reduce overfitting when training and testing also increase the generalization of our model. Increasing the amount of data automatically increases the accuracy of the testing and training. The benefits of increasing sample datasets are high speed, better prediction, classification results, and reduced cost.

The developed technique’s achieved accuracy rate is compared with other existing methods such as CN, FST, RCNN, OS, and CCA. Moreover, the RCNN replica attained 96%, and the FST technique gained 94.88%. Also, the CN and OS techniques’ gained accuracy rates are 96.86% and 96.11%, respectively. The developed ABCNM attained high accuracy while comparing other techniques to classify WBC as 99.12%.

**Precision**

This process has been evaluated to identify the correct positive estimates that are aligned with the overall positive estimates. Also, precision is the proportion of WBC classification, which is computed using Eq. (7),

$$P = \frac{T_p}{T_p + F_p} \tag{7}$$

The precision of the proposed ABCNM model is calculated and validated using prevailing methods like CN, RCNN, FST, CCA, and OS approaches. Initially, the RCNN technique gained a 96% precision rate, and the CN replica achieved 92.50% precision using a 2000 sample dataset. Moreover, the FST and CCA methods gained a 94.75% and 92.5% precision rate. Finally, the developed ABCNM technique gained 98.16% in precision rate. Thus, the comparison of precision rate is detailed in Fig. 10.

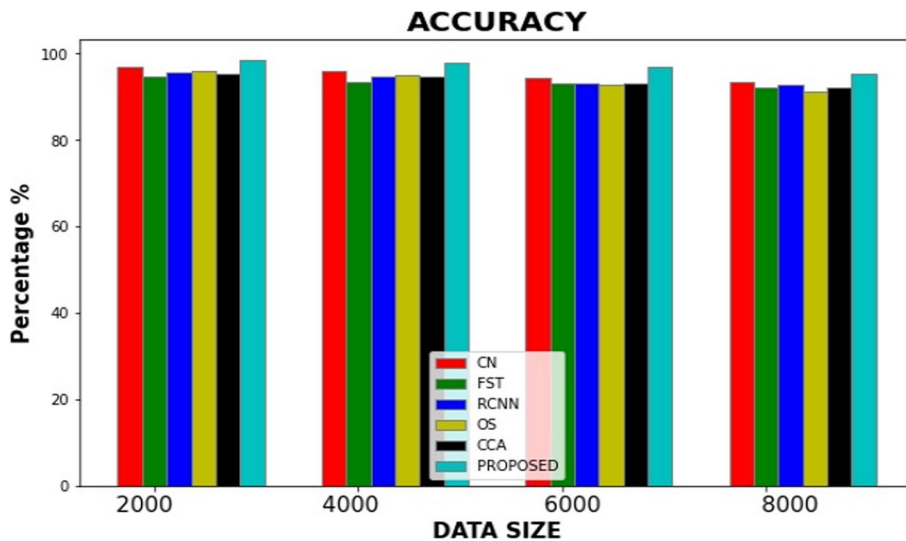
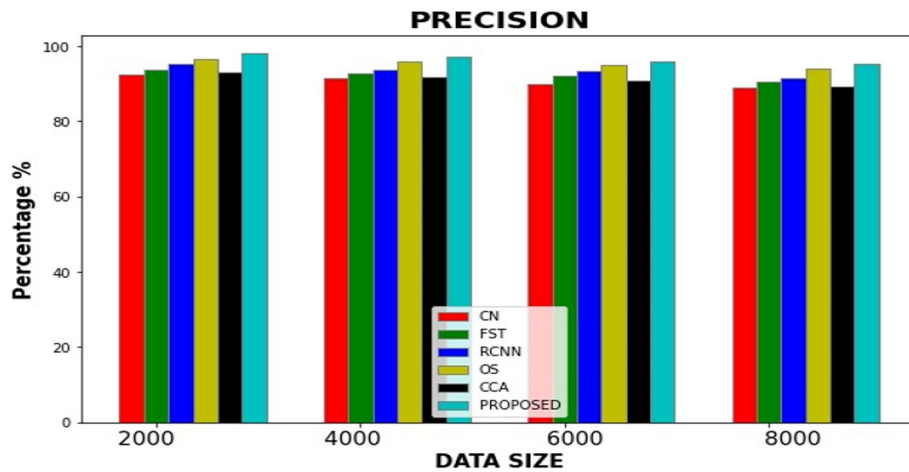


Fig. 9 Comparison of accuracy



**Fig. 10** Comparison of Precision

Here, the existing approaches have achieved a lower precision of only almost 97%. Additionally, the developed ABCNM approach has obtained a higher precision value of 98.16% than other methods.

**F-measure**

The calculation is based on precision and recall measurements to recognize the efficiency of tracing images, which is measured using Eq. (8),

$$F - measure = 2 \left( \frac{P * R}{P + R} \right) \tag{8}$$

Let P denote the precision value, and R represents the recall value. The F1-score of the proposed ABCNM model is calculated and validated using existing methods like OS, FST, and CCA approaches. Initially, the FST technique gained 95% in F-measure, and the OS replica achieved 98.5% in precision for using a 2000 sample dataset. Moreover, the CCA method gained a 92% precision rate. Finally, the developed ABCNM technique gained an F-measure rate of 99.02%. Thus, the comparison of the F-measure rate is detailed in Fig. 11. Here, the existing approaches achieved a lower F1-score, and the proposed model has attained a higher F1-score value of 99.02% than other methods.

**Sensitivity**

Sensitivity calculates the precise number of predicted true positives. Also, the probability of segmenting the WBC is measured using Eq. (9),

$$S_N = \frac{T_p}{T_p + T_n} \tag{9}$$

The sensitivity of the proposed ABCNM model is calculated and validated using existing methods like OS, CN, RCNN, FST, and CCA approaches.

The developed technique’s achieved sensitivity rate is compared with existing methods such as CN, FST, RCNN, OS, and CCA. Moreover, the RCNN replica attained 98%, and

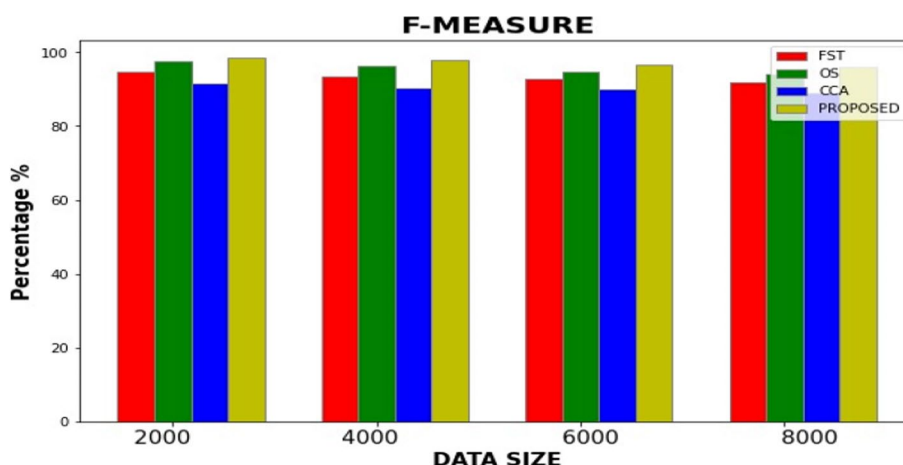


Fig. 11 Comparison of F-measure

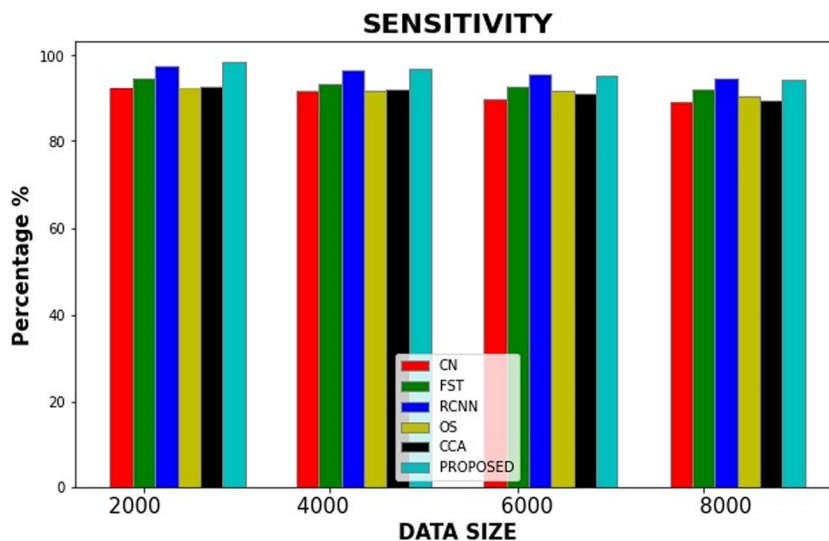


Fig. 12 Comparison of Sensitivity

the FST technique gained 95%. Also, the CN and OS techniques gained sensitivity rates of 92.50% and 93%, respectively.

The existing approaches have achieved a lower sensitivity of almost 98% for the 2000 data size. This ABCNM method has achieved a 99% higher sensitivity value than other methods, which proves the effectiveness of the developed model. The comparison of sensitivity is graphically represented in Fig. 12.

**Specificity**

Specificity is defined as the degree to which the number of true negatives recognized accurately is determined. Specificity is also employed to calculate the efficiency of detecting WBC using segmented images. The mathematical expression to calculate specificity is detailed in Eq. (10).

$$S = \frac{T_n}{T_p + T_n} \tag{10}$$

The achieved specificity rate of the developed technique is compared with other existing methods such as CN, OS, and CCA. Moreover, the CN replica attained 98.01 for using 500 sample WBC datasets. Also, OS and CCA techniques gained specificity rates are 95% and 97%, respectively. The developed ABCNM attained high specificity while comparing other techniques to classify WBC as 99.04%. The comparison of specificity is illustrated in Fig. 13.

**Execution time**

The execution time defines the time taken by the developed strategy to perform tasks such as data preprocessing, feature extraction, segmentation, classification, and optimization. It defines the computational efficiency of the model in classifying the different types of WBCs. The Fig. 14 presents the computational time comparison.

Here, the execution time of the developed algorithm was compared and validated with existing techniques such as CN, OS, FST, RCNN, and CCA. These models consumed 12 s, 13 s, 17 s, 9 s, and 11 s, respectively, to perform the above tasks, while the designed strategy incurred a minimum execution time of 4 s. The reduction of execution time in the proposed algorithm validates its computational efficiency.

**ROC curve comparison**

The Receiver Operating Characteristic (ROC) curve indicates the graphical representation classification performance of the developed model in WBC categorization. The ROC curve plots the true positive rate (sensitivity) against the false positive rate (1-specificity) across a range of threshold values. This enables us to determine how well the classification model differentiates between different classes of WBCs. The Area Under the ROC Curve (AUC) in the ROC curve determines the model’s discrimination efficiency.

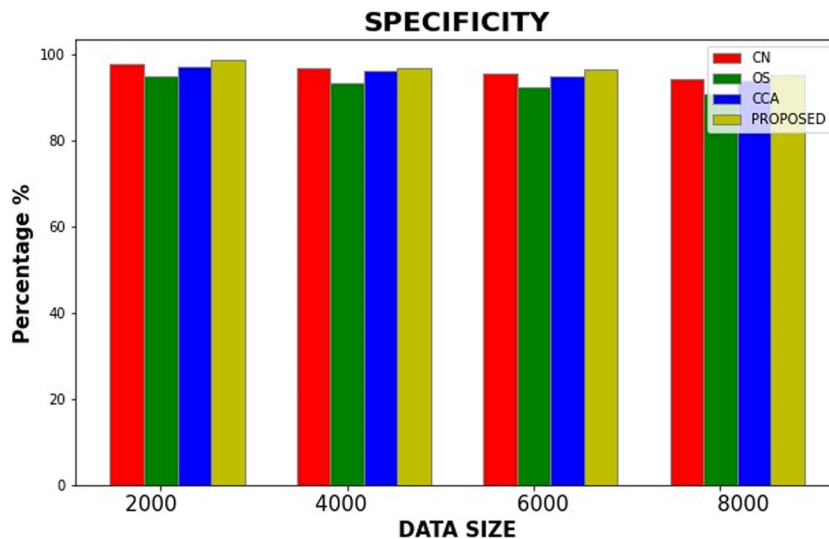
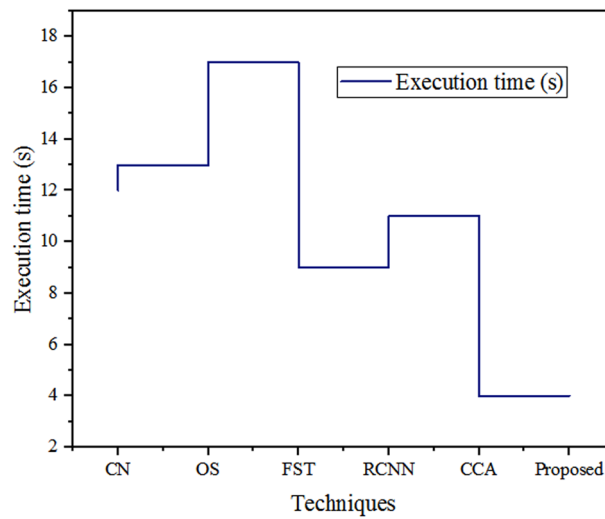
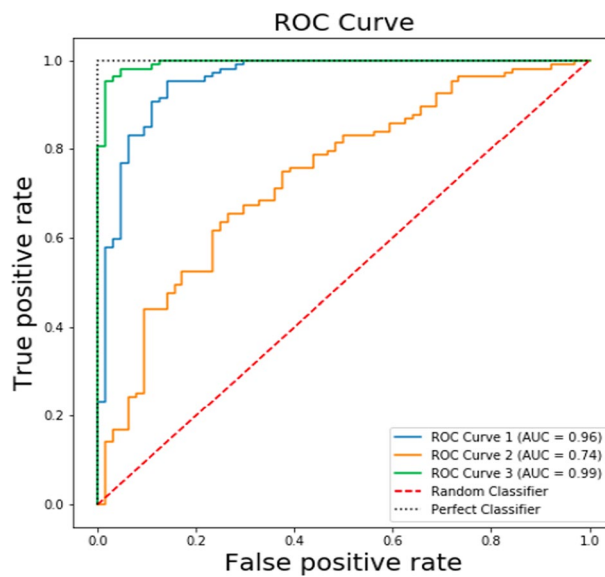


Fig. 13 Comparison of Specificity



**Fig. 14** Execution time comparison



**Fig. 15** AUC and ROC Curves Comparison

An AUC of 1 represents perfect classification and vice versa. High AUC values indicate that the algorithm effectively differentiates between various types of white blood cells, leading to more reliable diagnostic outcomes. Figure 15 compares the ROC curve of the proposed model with conventional models like CN and RCNN. The proposed strategy achieved an AUC of 0.99, perfectly classifying the WBC types. ROC curve 1 defines the AUC of RCNN, while ROC curve 2 defines the AUC of CN. The ROC curve 3 indicates the curve achieved by the proposed strategy.

**Comparison with hybrid methods**

In this module, the performance of the proposed strategy was compared with the hybrid classification models. The existing techniques included in the comparative evaluation include deep convolutional autoencoder (DCAE) [39], Grey Wolf Optimization with Discrete Charlier-Meixner hybrid polynomials (GWO-DCMP) [40], Hybrid Mayfly Algorithm with Particle Swarm Optimization (HMA-PSO) [41], and Support Vector Machine-Convolutional Neural Network (SVM-CNN) [42]. The parameters used for comparative assessment include accuracy, precision, sensitivity, f-measure, and specificity.

The accuracy represents the overall effectiveness and correctness of the proposed strategy in classifying WBCs. The existing models, including DCAE, GWO-DCMP, HMA-PSO, and SVM-CNN, obtained an accuracy of 94.62%, 96.17%, 92.65%, and 94.36%, respectively, while the proposed strategy obtained an accuracy of 99.12%. The significant enhancement of accuracy indicates the proposed strategy classifies the WBCs more accurately than the conventional models. This illustrates that the integration of ABO into the CNN approach improves the classification of WBCs, providing a promising solution for real-time WBC classification. On the other hand, precision measures the model’s effectiveness in accurately identifying the true positives from all positive instances. The above-stated conventional models achieved a precision of 94.18%, 95.68%, 91.37%, and 94.48%, respectively, while the proposed strategy obtained a higher precision of 98.16%. The improved precision of the developed algorithm highlights its effectiveness in classifying the specific class of WBCs. Consequently, the sensitivity quantifies the model’s ability to identify all relevant instances in WBC classification. The existing techniques, such as DCAE, GWO-DCMP, HMA-PSO, and SVM-CNN, earned sensitivity of 94.34%, 95.68%, 91.37%, and 94.48%, respectively, while the developed algorithm achieved 99% sensitivity. The higher sensitivity rate of the proposed methodology highlights the model’s effectiveness in precisely detecting all relevant instances. Finally, the f-measure performance was compared and validated with the existing hybrid models. The f-measure metric provides a balanced evaluation of system performances considering positive and negative cases. The above-stated conventional models obtained f-measures of 94.37%, 95.06%, 92.33%, and 94.05%, respectively. However, the proposed strategy earned a higher f-measure of 99.02%, indicating that it accurately classifies both positive and negative instances. Table 6 presents the comparative evaluation of the proposed model’s performance with the existing techniques.

**Table 6** Comparison with hybrid models

Methods	Accuracy	Precision	Sensitivity	F-measure
DCAE	94.62	94.18	94.34	94.37
GWO-DCMP	96.17	95.68	96.22	95.06
HMA-PSO	92.65	91.37	92.50	92.33
SVM-CNN	94.36	94.48	94.10	94.05
<b>Proposed</b>	<b>99.12</b>	<b>98.16</b>	<b>99</b>	<b>99.02</b>

To highlight the results of proposed model the Bold format is used that attracts the researchers for comparison with existing models

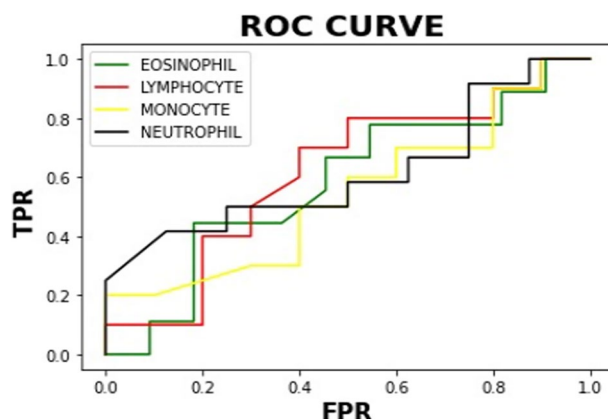


Fig. 16 ROC curve

Table 7 Overall performance metrics

Performance assessment with key metrics					
Methods	Accuracy	Precision	Sensitivity	Specificity	F-measure
CN	96.86	92.50	92.50	98.01	–
FST	94.88	94.75	95	–	95
RCNN	96	96	98	–	–
OS	96.11	97.1	93	95	98.5
CCA	95.89	92.5	93.7	97	92
<b>Proposed</b>	<b>99.12</b>	<b>98.16</b>	<b>99</b>	<b>99.04</b>	<b>99.02</b>

To highlight the results of proposed model the Bold format is used that attracts the researchers for comparison with existing models

The comprehensive comparative evaluation clearly shows that the proposed strategy achieved improved results compared to the existing hybrid models. These improved outcomes depict its effectiveness and robustness in identifying and classifying the different types of WBCs. This also highlights that integrating African Buffalo Optimization into the Convolutional Neural Network seamlessly enhances the classification of WBCs.

### Discussion

The proposed ABCNM model attained the best results in accuracy, sensitivity, specificity, F-measure, precision, and execution time. Thus, the developed scheme removed the training flaws in the initial stage. At next, extract features based on colours, morphological operation, and granularity features. The graphical representation of ROC curve is shown in Fig. 16.

Moreover, segmentation is executed in the pooling layer and WBC is classified in the fully connected layer. Thus, the developed ABCNM technique enhances the performance of WBC classification.

The outstanding metrics comparisons are tabulated in Table 7, in all parameter validation, the proposed ABCNM has gained the finest results. Moreover, they gained less execution time as 40 ms, high sensitivity as 99% and high classification accuracy as 99.12%. Hence, the robustness of the proposed ABCNM is verified and it can classify WBC.



## Conclusions and future work

A novel ABCNM is developed to classify WBC type in the blood sample, enhancing classification accuracy. Here, blood sample images are taken as input images and trained in the system. The developed ABCNM involves preprocessing, feature extraction, segmentation, and classification. Moreover, the proposed technique is processed on blood sample datasets, and the WBC is finally segmented. Furthermore, segmented images are employed in the fully connected layer to track the position of the WBC and classify the WBC type.

The developed technique has also achieved better accuracy, precision, F-measure, recall, sensitivity, execution time, and error rate. Thus, it achieved 99.12% accuracy in classifying WBC types. Comparing the past methods, it has shown a 2% accuracy improvement for classifying WBC types such as neutrophil, lymphocyte, eosinophil, and monocyte.

Although the proposed strategy achieved improved outcomes in WBC classification, it faces certain challenges. Firstly, the classification performance of the developed algorithm relies on the quality and quantity of the data used, which makes the system data-dependent and restricts its application in real-world scenarios. Secondly, the proposed algorithm may struggle to handle imbalanced datasets, reducing its classification. The other limitation of the study is that it is not tested or validated across diverse databases, which restricts its generalization efficiency. Hence, future work should concentrate on developing a multi-objective classification model, which produces accurate results irrespective of the quality and quantity of the data used in model training. Also, future work should focus on evaluating the developed strategy across imbalanced and diverse blood sample image databases to ensure its generalization across real-world scenarios.

### Author contributions

Author 1 has worked on the problem statement and drafted the entire manuscript with experimental results. Author 2 reviewed the manuscript and guided with valuable inputs.

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### Availability of data and materials

Data sharing is not applicable to this article as no new data were created or analyzed in this study.

### Code availability

Not applicable.

## Declarations

### Ethics approval and consent to participate

Not applicable.

### Consent for publication

Informed consent does not apply as this was a retrospective review with no identifying patient information.

### Human and animal rights

This article does not contain any studies with human or animal subjects performed by any of the authors.

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The authors declare no competing interests.

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## References

1. Saghiv MS, Michael SS. The immune system basic exercise physiology. Cham: Springer; 2020.

2. Chen X, et al. Bone marrow myeloid cells regulate myeloid-biased hematopoietic stem cells via a histamine-dependent feedback loop. *Cell Stem Cell*. 2017;21(6):747–60.
3. Ruberto Di, Cecilia AL, Putzu L. Detection of red and white blood cells from microscopic blood images using a region proposal approach. *Comput Biol Med*. 2020;116:103530.
4. Wirth MD, et al. The dietary inflammatory index is associated with elevated white blood cell counts in the national health and nutrition examination survey. *Brain, Behavior Immunity*. 2018;69:296–303.
5. Gupta D, et al. Optimized binary bat algorithm for classification of white blood cells. *Measurement*. 2019;143:180–90.
6. Maresso AW. Innate immunological defenses against bacterial attack. In: Maresso AW, editor. *Bacterial Virulence*. Cham: Springer; 2019. p. 31–46.
7. Schuppan D, Gisbert-Schuppan K. "Immunology of the intestine wheat syndromes. Cham: Springer; 2019. p. 11–23.
8. Wolfien M, et al. Hematopoietic stem-cell senescence and myocardial repair-Coronary artery disease genotype/phenotype analysis of post-MI myocardial regeneration response induced by CABG/CD133+ bone marrow hematopoietic stem cell treatment in RCT PERFECT Phase 3. *EBioMedicine*. 2020;57:102862.
9. Giger S, et al. A single metabolite which modulates lipid metabolism alters hematopoietic stem/progenitor cell behavior and promotes lymphoid reconstitution. *Stem cell reports*. 2020;15(3):566–76.
10. Ghosh S, et al. A review on the effect of COVID-19 in type 2 asthma and its management. *Int Immunopharmacol*. 2021;91:107309.
11. Khazdair MR, et al. A qualitative and quantitative comparison of *Crocus sativus* and *Nigella sativa* immunomodulatory effects. *Biomed Pharmacother*. 2021;140:111774.
12. Harshanand BA, Sangaiah AK. Comprehensive analysis of deep learning methodology in classification of leukocytes and enhancement using swish activation units. *Mobile Netw Appl*. 2020;25(6):2302–20.
13. Muriel J, et al. Ontogeny of leukocyte profiles in a wild altricial passerine. *J Comp Physiol B*. 2021;191(1):195–206.
14. Abdul-Jawad S, et al. Acute immune signatures and their legacies in severe acute respiratory syndrome coronavirus-2 infected cancer patients. *Cancer Cell*. 2021;39(2):257–75.
15. Zeng Z, et al. Longitudinal changes of inflammatory parameters and their correlation with disease severity and outcomes in patients with COVID-19 from Wuhan, China. *Crit Care*. 2020;24(1):1.
16. Baydilli YY, Atila U, Elen A. Learn from one data set to classify all—a multi-target domain adaptation approach for white blood cell classification. *Computer Methods Programs Biomed*. 2020;196:105645.
17. Ma Li, et al. Combining DC-GAN with ResNet for blood cell image classification. *Med Biol Eng Comput*. 2020;58(6):1251–64.
18. Teng, Chieh-Fang, et al. 2021 "Convolutional neural network-aided bit-flipping for belief propagation decoding of polar codes." ICASSP 2021–2021 IEEE International Conference on Acoustics, Speech and Signal Processing (ICASSP). IEEE.
19. Sevinc O, et al. White blood cell classification using genetic algorithm-enhanced deep convolutional neural networks. In: Arabia HR, Ferens K, de la Fuente D, Kozerenko EB, Varela JAO, Tinetti FG, editors., et al., *Advances in Artificial Intelligence and Applied Cognitive Computing*. Cham: Springer; 2021. p. 29–43.
20. Al-Dulaimi K, Khudhair, A, et al. "Classification of white blood cell types from microscope images: Techniques and challenges. In: Mendez-Vilas A, Torres-Hergueta E, editors., et al., *Microscopy science: last approaches on educational programs and applied research (Microscopy Book Series, 8)*. Spain: Formatex Research Center; 2018. p. 17–25.
21. Battiato, Sebastiano, et al. 2020 "Detection and classification of pollen grain microscope images." Proceedings of the IEEE/CVF Conference on Computer Vision and Pattern Recognition Workshops.
22. Li L, et al. Recent advances in artificial intelligence and machine learning for nonlinear relationship analysis and process control in drinking water treatment: a review. *Chem Eng J*. 2021;405:126673.
23. Tavakoli S, et al. New segmentation and feature extraction algorithm for classification of white blood cells in peripheral smear images. *Sci Rep*. 2021;11(1):1–13.
24. Saadatnejad S, Oveisi M, Hashemi M. LSTM-based ECG classification for continuous monitoring on personal wearable devices. *IEEE J Biomed Health Informat*. 2020;24(2):515–23.
25. Ferdosi, Bilkis Jamal, Sharmilee Nowshin, and Farzana Ahmed Sabera. 2018 "White blood cell detection and segmentation from fluorescent images with an improved algorithm using k-means clustering and morphological operators." 2018 4th International Conference on Electrical Engineering and Information & Communication Technology (ICEEICT). IEEE.
26. Özyurt F. A fused CNN model for WBC detection with MRMR feature selection and extreme learning machine. *Soft Comput*. 2020;24(11):8163–72.
27. Hong K-S, Jawad Khan M, Hong MJ. Feature extraction and classification methods for hybrid fNIRS-EEG brain-computer interfaces. *Front Human Neurosci*. 2018;12:246.
28. Baydilli YY, Atila Ü. Classification of white blood cells using capsule networks. *Comput Med Imaging and Graph*. 2020;80:101699.
29. Toğaçar M, Ergen B, Cömert Z. Classification of white blood cells using deep features obtained from convolutional neural network models based on the combination of feature selection methods. *Appl Soft Comput*. 2020;97:106810.
30. Kutlu H, Avci E, Özyurt F. White blood cells detection and classification based on regional convolutional neural networks. *Med Hypotheses*. 2020;135:109472.
31. Patil AM, Patil MD, Birajdar GK. White blood cells image classification using deep learning with canonical correlation analysis. *IRBM*. 2021;42(5):378–89.
32. Lu Y, et al. WBC-Net: a white blood cell segmentation network based on UNet++ and ResNet. *Appl Soft Comput*. 2021;101:107006.
33. Sahlol AT, Kollmannsberger P, Ewees AA. Efficient classification of white blood cell leukemia with improved swarm optimization of deep features. *Sci Rep*. 2020;10(1):1–11.
34. Quan Q, Wang J, Liu L. An effective convolutional neural network for classifying red blood cells in malaria diseases. *Int Sci Comput Life Sci*. 2020;12(2):217–25.

35. Shahin AI, et al. White blood cells identification system based on convolutional deep neural learning networks. *Comput Method Program Biomed.* 2019;168:69–80.
36. Ghaderzadeh M, Hosseini A, Asadi F, Abolghasemi H, Bashash D, Roshanpoor A. Automated detection model in classification of B-lymphoblast cells from normal B-lymphoid precursors in blood smear microscopic images based on the majority voting technique. *Sci Program.* 2022;2022:1–8.
37. Fasihfar Z, Rokhsati H, Sadeghsalehi H, Ghaderzadeh M, Gheisari M. AI-driven malaria diagnosis: developing a robust model for accurate detection and classification of malaria parasites. *Iranian J Blood Cancer.* 2023;15(3):112–24.
38. Hosseini A, Eshraghi MA, Taami T, Sadeghsalehi H, Hoseinzadeh Z, Ghaderzadeh M, Rafiee M. A mobile application based on efficient lightweight CNN model for classification of B-ALL cancer from non-cancerous cells: a design and implementation study. *Informat Med Unlocked.* 2023;39:101244.
39. Elhassan TA, Rahim MSM, Zaiton MHS, Swee TT, Alhaj TA, Ali A, Aljurf M. Classification of atypical white blood cells in acute myeloid leukemia using a two-stage hybrid model based on deep convolutional autoencoder and deep convolutional neural network. *Diagnostics.* 2023;13(2):196.
40. Tahiri MA, Bencherqui A, Karmouni H, Amakdouf H, Sayyouri M, Qjidaa H. White blood cell automatic classification using deep learning and optimized quaternion hybrid moments. *Biomed Signal Process Control.* 2023;86:105128.
41. Yentrapragada D. Deep features based convolutional neural network to detect and automatic classification of white blood cells. *J Ambient Intell Humaniz Comput.* 2023;14(7):9191–205.
42. Olayah F, Senan EM, Ahmed IA, Awaji B. Blood slide image analysis to classify WBC types for prediction haematology based on a hybrid model of CNN and handcrafted features. *Diagnostics.* 2023;13(11):1899.

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