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Optic Disc and Cup Segmentation Through Fuzzy Broad Learning System for Glaucoma Screening

Riaz Ali, Bin Sheng, Ping Li, *Member, IEEE*, Yan Chen, Huating Li, Po Yang, Younhyun Jung, Jinman Kim, and C. L. Philip Chen, *Fellow, IEEE*

Abstract—Glaucoma is an ocular disease that causes permanent blindness if not cured at an early stage. Cup-to-disc ratio (CDR), obtained by dividing the height of optic cup (OC) with the height of optic disc (OD), is a widely adopted metric used for glaucoma screening. Therefore, accurately segmenting OD and OC is crucial for calculating a CDR. Most methods have employed deep learning methods for the segmentation of OD and OC. However, these methods are very time-consuming. We present a new fuzzy broad learning system-based technique for OD and OC segmentation with glaucoma screening. We comprehensively integrated extracting a region of interest (ROI) from RGB images, data augmentation, extracting red and green channel images, and inputting them to the two separate fuzzy broad learning system-based neural networks for segmenting the OD and OC, respectively, and then calculated CDR. Experiments show that our fuzzy broad learning system-based technique outperforms many state-of-the-art methods.

Index Terms—Ocular disease, optic disc and cup, segmentation, broad learning system, neural networks, fuzzy system.

I. INTRODUCTION

GLAUCOMA is an ophthalmic disease that causes damage to the optic disc (also called the optic nerve head (ONH)). It is the second prime cause of blindness worldwide and can affect everyone from newborn babies to older people [1]. Glaucoma is usually assessed in three ways, which are: measuring the intraocular pressure (IOP), conducting a visual field test, and examining ONH [2]. However, ONH assessment is preferred and practiced widely by experts [3]. Cup-to-disc ratio (CDR) is a clinical measurement that is commonly

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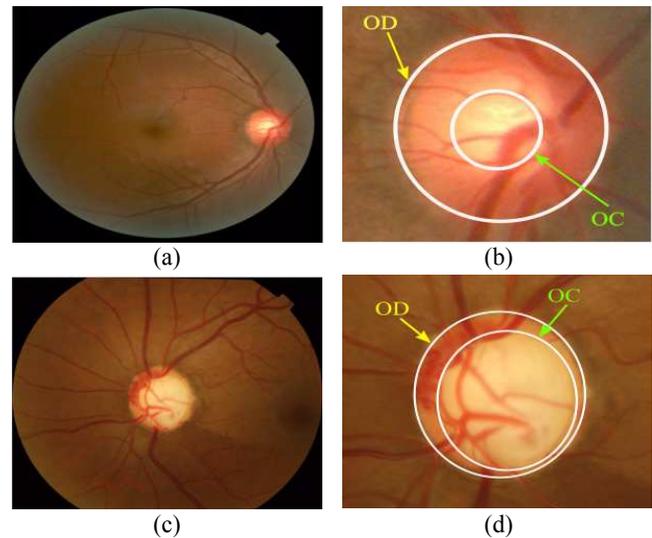


Fig. 1: (a) Image of a normal eye. (b) Boundaries of OD and OC in a normal eye. (c) Image of a glaucomatous eye. (d) Boundaries of OD and OC in a glaucoma affected eye.

used by ophthalmologists for ONH assessment [3]. CDR is calculated as the ratio of the vertical diameter of optic cup (OC), a yellowish brighter area at the center of optic disc (OD), to the vertical diameter of OD. Because internal structures of an eye are affected by the increased pressure caused by glaucoma, the size of OC increases with respect to the size of OD; a phenomenon also called cupping [4]. Thence, usually a larger CDR indicates the presence of glaucoma. Fig. 1 shows a significant difference in sizes of OD and OC in a healthy and glaucomatous eye. Traditionally, clinicians calculated CDR by manually segmenting OD and OC. Yet, manual segmentation takes huge time and is prone to the subjective judgment of ophthalmologists. Hence, there is a need for automatic and fast segmentation of OD and OC for large-scale and efficient diagnosis of glaucoma.

Several works have been proposed for OD and OC segmentation. Some of them use deep learning, and some employ non-deep learning-based approaches. The non-deep learning-based methods are not much time-consuming, but they usually lack in providing as promising results as deep learning approaches do. These methods mainly consist of color and contrast thresholding, region segmentation, and boundary detection techniques [5], [6]. Most of the modern techniques, which rely on the features like texture, color, and gradient, do not

using pseudoinverse. This system has the combined advantages of both BLS and fuzzy systems.

Therefore, this paper targets at finding out a practically effective and efficient glaucoma screening solution by studying the state-of-the-art OD and OC segmentations. Due to the efficiency and performance capability of FBLS, we propose a new method of OD and OC segmentation, which adopts this learning model. We comprehensively integrated extracting a region of interest (ROI) from RGB images, data augmentation, extracting red and green channel images, and inputting them to the two separate fuzzy broad learning system-based neural networks for segmenting the OD and OC, respectively, and then calculated CDR. Experiments show that our fuzzy broad learning system-based technique outperforms state-of-the-art methods, and our approach significantly accelerates the computational time and maintains high accuracy of OD/OC segmentation and glaucoma detection.

The contributions of our work presented in this paper are:

- A novel broad learning-based fuzzy neural network is proposed for effective and efficient OD and OC segmentation for glaucoma screening, where the mapped feature nodes of the original BLS are replaced with Takagi-Sugeno fuzzy subsystems. Each of these fuzzy subsystems processes the input data, hence this method has the advantage of an ensemble structure.
- This approach extracts features more effectively from the input image because it uses the k -means algorithm to decide the count of fuzzy rules in every fuzzy subsystem and Gaussian membership functions' centers. The application of the k -means technique produces different centers from the input data for every subsystem, that assures the generation of more distinct features and thus results in the extraction of abundant discriminative information from the input image as compared to BLS.
- An enhanced least square method is designed in the proposed BLS-based fuzzy model to get the final weights with smaller training errors. Also, an l_1 -norm regularization is employed to prohibit the model from overfitting. This makes our approach maintain high accuracy of OD/OC segmentation and glaucoma detection.
- An extensive in-depth and thorough experimental evaluation is carried out to show the robustness and usefulness of our technique. The obtained results exhibit that our method sufficiently outperforms state-of-the-art algorithms in OD and OC segmentation and glaucoma classification. Additionally, for evaluation of glaucoma, we construct a new dataset of retinal images called Shanghai Chinese Retinal Images Dataset (SCRID) comprising 566 retinal fundus images.

II. RELATED WORK

A. Morphological Operations and Reconstruction

Some of the earlier works use morphological operations and reconstruction for segmenting OD and OC. Nugroho *et al.* [4] introduced morphological reconstruction and convex hull for segmenting OC, while OD is segmented by combining morphological reconstruction and active contour. However,

active contour models depend on a proper initialization of OD contour for providing optimal results [14]. In [23], the dilation operation was used to overcome the problem of the misclassifying area around the retina after applying thresholding. They also use opening followed by dilation to remove the irregularity of segmented OD and to give proper disc size. For segmenting cup, they use a closing operation to remove blood vasculature in the green color channel. The work of [24] involved dilation and erosion operations to eliminate blood vessels in OD, and to smoothen the intensity around OD. Similarly, [25] also used morphological operations for removing retinal vasculature and other pathologies and enhancing OD for getting more accurate segmentation.

B. K -Means Clustering, Superpixel Classification, and Shape-Based Models

Ayub *et al.* [26] detected glaucoma via k -means clustering for color-based segmentation of OD and OC after applying preprocessing and extracting ROI. K is assigned the value 2 because the entire image is divided into two regions by preprocessing. Arumugam and Nivedha [27] also segmented OD by applying k -means clustering after preprocessing and removing blood vessels. Cheng *et al.* [2] applied the superpixel classification technique for segmenting OD and OC. The former two methods use a simple linear iterative clustering (SLIC) algorithm to combine neighboring pixels into superpixels. The algorithms applied here are more efficient than the conventional k -means algorithm as the size of the search region is limited here, which results in less number of distance calculations. In the shape-based techniques, OD is modeled as an elliptical or circular object. These methods try to segment OD by Hough transforms [28] or ellipse fitting [29]. However, these shape-based methods are unsuitable for images of different color intensities and containing blood vessels.

C. Learning-Based Methods

Many methods like [30]–[32] used machine learning techniques to segment OD and OC. The work of [30] segmented OD and OC from color fundus images of the retina using a fully automatic regression-based technique. Initially, it applies a circular Hough transform to roughly segment OD. The preliminary shapes of OD and OC are calculated from the approximated OD obtained from the previous step. The method repeatedly learns the final shape of OD and OC from the preliminary shape by a cascaded shape regression technique. For better performance, a data augmentation technique is employed to produce artificial training data. Abramoff *et al.* [31] used stereo color photographs for the evaluation of automatic OD segmentation. They obtained a reference segmentation of the neuroretinal rim and OC by glaucoma experts manually. Then, they evaluated stereo pairs and corresponding reference standard by pixel feature classification. However, these approaches rely on the extracted features, which may help them to perform well on specific datasets but not on others. Moreover, manual feature extraction makes these techniques time-consuming. In the recent past, many methods have used deep learning for OD and OC segmentation

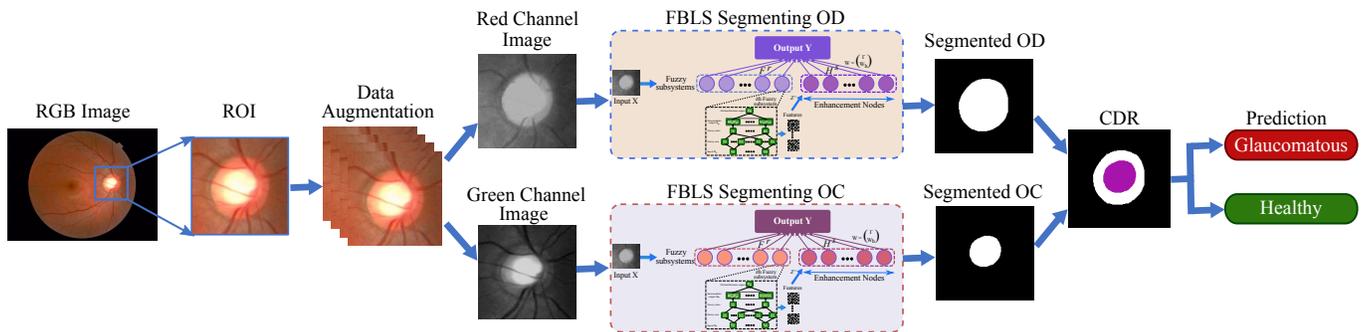


Fig. 3: Overview of our proposed approach of OD and OC segmentation for glaucoma detection using FBS.

because of its promising results. Lim *et al.* [33] applied a CNN to segment OD and OC. First, they roughly confine the ROI. Then the ROI is transformed into visual features. After that, the method generates a pixel-level probability map of retinal structures through CNN to classify the transformed image. Finally, boundaries of OD and OC are predicted by the segmentation of that map. The work of [3] segmented OD and OC jointly by employing a deep CNN called M-Net. The essential four parts of M-Net are a multi-scale input layer, a U-shaped CNN, a side-out layer, and a multi-label loss function. An image pyramid is built by the input layer. The U-shaped CNN is the primary structure that learns the hierarchical features. The side-out layer works as an initial classifier and generates local prediction maps. Lastly, the final segmentation map is created by the multi-label loss function. Additionally, their method applies polar transformation to convert fundus images to a polar coordinate system that produces better-quality segmentation results. Many other methods have also utilized deep learning for the segmentation of OD and OC. Although deep learning-based methods provide excellent results, they take a long time to train the models.

D. Fuzzy Logic

The authors of [24] employ fuzzy *c*-means and morphological operations to segment OD and OC but lack in providing exceptional results as compared to state-of-the-art approaches. Some authors, like [34] and [35], have obtained competent results in regression and classification by using FNNs. Nonetheless, FNNs consume much time in training the parameters of fuzzy rules because FNNs are trained like usual neural networks, which take a lot of time for training. Some researchers have proposed improved algorithms to overcome the problems present in FNNs. The work of [36] proposes a hierarchical hybrid FNN in which a few fuzzy subsystems randomly combine various input features to form an intermediary output, and then a neural network handles these intermediate outputs and the remaining input attributes together to decrease the number of fuzzy rules and input dimension. An online sequential fuzzy extreme learning machine is proposed in [37], where membership functions are assigned the antecedent parameters randomly, and then corresponding subsequent parameters are determined, which reduces the learning time of the algorithm. Sun *et al.* [38] have developed a fuzzy learning system that aggregates the

data through *k*-means clustering and derives the membership functions in every fuzzy rule by using an extreme learning machine.

Recently, Feng *et al.* [16] have proposed FBS, which comprises fuzzy subsystems and enhancement nodes. In this approach, every fuzzy subsystem processes the incoming data, and the features of input data are preserved because the output of total fuzzy subsystems is dispatched to enhancement nodes to be transformed nonlinearly. Finally, the weights connecting the coefficients of every fuzzy subsystem and output of enhancement nodes with the final output layer are computed through pseudoinverse. The training process in FBS is significantly reduced because of its flat network. Keeping the need for fast computation and accuracy in mind, we propose a method of OD and OC segmentation, which is less time-consuming and provides decent results.

III. PROPOSED METHOD

Fig. 3 shows an overview of our proposed method of glaucoma detection using OD and OC segmentation. The steps involved are discussed below:

A. Extracting the Region of Interest

As shown in Fig. 3, the first step of our segmentation approach is the extraction of an ROI from the original RGB fundus image. We have used the technique of [39] to get the ROI. In addition to the OD area, the ROI also contains some background information to help the network discriminate the OD from the background.

B. Data Augmentation

Machine learning and deep learning methods need to be trained on the large datasets for providing optimal performance. However, most of the specialized tasks, such as medical image classifications or segmentation tasks, have insufficient data. Most of the retinal datasets also contain only a few hundred images. The models trained with small datasets do not generalize well and are less accurate in the testing. Researches such as [40] demonstrated that data augmentation could prevent this problem. Hence, we have also applied data augmentation in our method to increase the training data size artificially. Our data augmentation consists of flipping left-to-right, flipping top-to-bottom, rotating 90° anti-clockwise,

and adding noise (consisting of Gaussian noise, and salt and pepper noise). Data augmentation has helped us achieve better results. Fig. 4 shows a sample segmentation without data augmentation, which lacks in providing promising results. Similarly, results without data augmentation are also shown in Fig. 6, Fig. 7, Fig. 10, and Fig. 11.

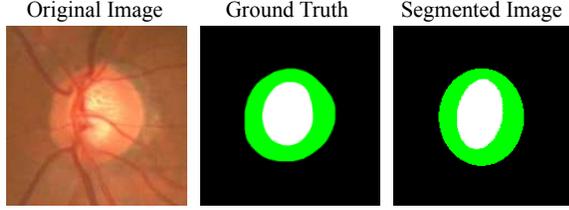


Fig. 4: A sample of segmentation results without applying data augmentation.

C. Extracting the Red and Green Channel Images

Compared to other channels, the red channel of an RGB fundus image is less influenced by blood vessels and shows more differences between the OD area and background (non-OD area). Hence, it is a suitable choice for OD segmentation. Likewise, the green channel is more appropriate for OC segmentation as the brightness and contrast of OC pixels is very high in this channel. Therefore, we have split the RGB image and fed the red channel and green channel images into two separate FBLSS for segmenting OD and OC, respectively. Before inputting the red and green channel images to the FBLSS, the mean intensity value is subtracted from the original intensity value of each of the channel images, which exposes the OD and OC areas more clearly for better segmentation.

D. OD and OC Segmentation

Here, we input the Red/Green channel images obtained from the previous step to FBLSS and receive the segmented OD/OC as an output. Let n denotes the quantity of fuzzy subsystems in FBLSS and m represents the enhancement nodes in FBLSS. So, the input data is represented as $X = (x_1, x_2, \dots, x_n)^T \in \mathbb{R}^{n \times m}$. Initially, the input $x_q = (x_{q1}, x_{q2}, \dots, x_{qm})$, $q = 1, 2, \dots, n$ is mapped to the i th fuzzy system by using the first-order Takagi-Sugeno fuzzy model. x_{qt} denotes a first-order polynomial in the Takagi-Sugeno fuzzy model, $t = 1, 2, \dots, m$. It can be calculated as Eq. (1):

$$z_{qk}^i = \sum_{t=1}^m \alpha_{kt}^i x_{qt} \quad (1)$$

where α_{kt}^i is the coefficient and $k = 1, 2, \dots, K_i$ denotes the fuzzy rule's number in the i th fuzzy model. We have chosen the Gaussian function as the membership function, which can be defined as shown in Eq. (2):

$$\mu_{kt}^i(x_{qt}) = e^{-\left(\frac{x_{qt} - c_{kt}^i}{\sigma_{kt}^i}\right)^2} \quad (2)$$

where c_{kt}^i and σ_{kt}^i represent, respectively, width and center of the Gaussian membership function. We have applied the k -means technique to get K_i clustering centers. Because of the

k -means method's randomness in the initial conditions, distinct centers are selected from the input in every fuzzy subsystem that helps extract more features due to the working of all fuzzy subsystems in this ensemble manner. After that, each rule's weighted activation level can be calculated as Eq. (3):

$$\omega_{qk}^i = \frac{\prod_{t=1}^m \mu_{kt}^i(x_{qt})}{\sum_{k=1}^{K_i} \prod_{t=1}^m \mu_{kt}^i(x_{qt})} \quad (3)$$

where $\prod_{t=1}^m \mu_{kt}^i(x_{qt})$ denotes fire strength of i th fuzzy subsystem's k th fuzzy rule. After that, the intermediary output of q th training sample in the i th fuzzy subsystem is defined as Eq. (4):

$$Z_{qi} = (\omega_{q1}^i z_{q1}^i, \omega_{q2}^i z_{q2}^i, \dots, \omega_{qK_i}^i z_{qK_i}^i) \quad (4)$$

Eq. (5) shows the output vector for the total training samples in an i th fuzzy subsystem.

$$Z_i = (Z_{1i}, Z_{2i}, \dots, Z_{ni}), i = 1, \dots, r \quad (5)$$

The Eq. (6) shows the intermediate output matrix for r fuzzy subsystems.

$$Z^r = (Z_1, Z_2, \dots, Z_r) \in \mathbb{R}^{n \times (K_1 + K_2 + \dots + K_r)} \quad (6)$$

After that, for preserving the characteristics of inputs, Z^r is nonlinearly transformed by the enhancement nodes, which is represented in Eq. (7):

$$H_j = \Psi(Z^r \omega_j + \beta_j), j = 1, \dots, s \quad (7)$$

where H_j denotes the enhancement nodes transformed from Z^r , ω_j and β_j are, respectively, weights and bias terms that connect the fuzzy subsystems' outputs to their correspondent enhancement layers and are selected randomly between values from 0 to 1, $\Psi(\cdot)$ is Sigmoid activation function. The groups of all the enhancement nodes are computed as Eq. (8):

$$H^s = (H_1, H_2, \dots, H_s) \quad (8)$$

The output obtained from each fuzzy subsystem is transmitted to the top layer, along with the output of enhancement nodes. Every fuzzy subsystem will be a multi-output model because the training target $Y \in \mathbb{R}^{N \times C}$ consists of C components. The output vector for i th fuzzy subsystem in q th training sample is calculated as denoted by Eq. (9):

$$\begin{aligned} F_{qi} &= \left(\sum_{k=1}^{K_i} \lambda_{k1}^i \omega_{qk}^i z_{qk}^i, \dots, \sum_{k=1}^{K_i} \lambda_{kC}^i \omega_{qk}^i z_{qk}^i \right) \\ &= \left(\sum_{k=1}^{K_i} \lambda_{k1}^i \omega_{qk}^i \left(\sum_{t=1}^m \alpha_{kt}^i x_{qt} \right), \dots, \right. \\ &\quad \left. \left(\sum_{k=1}^{K_i} \lambda_{kC}^i \omega_{qk}^i \left(\sum_{t=1}^m \alpha_{kt}^i x_{qt} \right) \right) \right) \\ &= \sum_{t=1}^m \alpha_{kt}^i x_{qt} (\omega_{q1}^i, \dots, \omega_{qK_i}^i) \begin{pmatrix} \lambda_{11}^i & \dots & \lambda_{1C}^i \\ \vdots & & \vdots \\ \lambda_{K_i1}^i & \dots & \lambda_{K_iC}^i \end{pmatrix} \end{aligned} \quad (9)$$

Eq. (10) shows the output vector for i th fuzzy subsystem's total training samples.

$$F_i = (F_{1i}, F_{2i}, \dots, F_{ni}) = D\Omega^i \lambda^i \quad (10)$$

where $D = \text{diag} \sum_{t=1}^m \alpha_{kt}^i x_{1t}, \dots, \sum_{t=1}^m \alpha_{kt}^i x_{nt}$, and

$$\Omega^i = \begin{pmatrix} \omega_{11}^i & \dots & \omega_{1K_i}^i \\ \vdots & & \vdots \\ \omega_{n1}^i & \dots & \omega_{nK_i}^i \end{pmatrix}, \lambda^i = \begin{pmatrix} \lambda_{11}^i & \dots & \lambda_{1C}^i \\ \vdots & & \vdots \\ \lambda_{K_i 1}^i & \dots & \lambda_{K_i C}^i \end{pmatrix}.$$

The aggregate output of r fuzzy subsystems is calculated as shown in Eq. (11):

$$F^r = \sum_{t=1}^r F_t = \sum_{t=1}^r D \Omega^i \lambda^i = D (\Omega^1, \dots, \Omega^r) \begin{pmatrix} \lambda^1 \\ \vdots \\ \lambda^r \end{pmatrix} = D \Omega \Gamma \quad (11)$$

where $\Omega = (\Omega^1, \dots, \Omega^r)$ is the matrix containing the fire strengths ω_{qk}^i and $\Gamma = ((\lambda^1)^T, \dots, (\lambda^r)^T)^T$ is comprised of the parameters to be computed subsequently.

In the end, we connect the output F^r of all fuzzy subsystems and H^s of the enhancement nodes to the output of FBLs denoted as Y , which is defined as Eq. (12):

$$\begin{aligned} Y &= F^r W_f + H^s W_h \\ &= D \Omega \Gamma + H^s W_h \\ &= (D \Omega, H^s) \begin{pmatrix} \Gamma \\ W_h \end{pmatrix} \\ &= P W \end{aligned} \quad (12)$$

where W_f and W_h , respectively, are the weights of F^r and H^s , and their value is set to 1. $P = (D \Omega, H^s)$ and W denotes the parameter matrix of fuzzy BLS which is comprised of Γ and W_h and can be calculated through the training targets $Y : W = P^+ Y$.

The vector P^+ can be computed by Eq. (13):

$$P^+ = \arg \min_W \|PW - Y\|_2^2 + \lambda \|W\|_1 \quad (13)$$

An enhanced least square method is used here to get W with smaller errors, and λ represents the additional restraints on the summation of squares of weights in [15]. The former part is an l_2 -norm regularization and shows the training errors. The latter part is an l_1 -norm regularization. The use of l_1 -norm regularization makes the proposed model robust against outliers [19] and averts our network from overfitting. The FBLs learn the discriminative features of images and distinguishes OD/OC from the background of retinal images and generates the segmentation maps.

E. Area Opening and Morphological Closing

In a few images, the segmentation result generated by FBLs contains small pixels outside OD and OC regions, which degrade the segmentation accuracy when the morphological closing operation is performed on these images. Therefore, we first perform the area opening operation on the segmented result obtained from the FBLs to remove those small objects outside the actual OD and OC area and then perform the morphological closing operation. We have used the closing operation to fill the holes in the segmented OD caused by the presence of blood vessels in that region. Fig. 5 shows the closed OD images with and without performing an area opening operation.

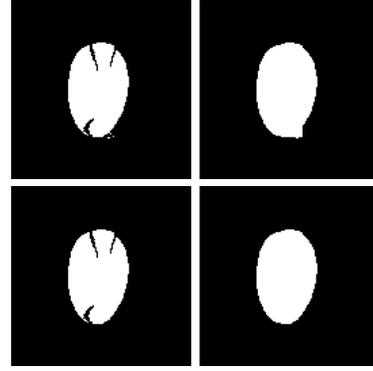


Fig. 5: The top and bottom rows show the segmentation results without and with the area opening operation, respectively.

IV. EXPERIMENTAL RESULTS

A. Datasets

We have performed experiments on one publicly available dataset RIM-ONE-r3 [41] and one dataset (SCRID) from the Shanghai Sixth People's Hospital. We could not get the ORIGA dataset (used by [3] and [14]) due to the clinical policies of its owner. The RIM-ONE-r3 dataset consists of 159 fundus images having a resolution of 2144×1424 pixels. The SCRID dataset contains 566 images of 2048×1536 pixels resolution. We divided the images in training and testing set with a ratio of 80% and 20%, respectively, and applied the data augmentation on only the training images. So, the number of training images became 635 and 2265 for the RIM-ONE-r3 and the SCRID dataset, respectively.

B. Training Time

One of the significant contributions of our method is the efficient training of the network model. Table I shows the training time of our proposed method, along with the training times of the state-of-the-art methods [3], [11]–[14]. We re-ran the methods of [3] and [13] with the same division of training and testing of 80% and 20% respectively, and present the consumed time in Table I. The reported times of other methods are taken from the respective papers as we could not get the codes of other methods to run them on our machine. Some methods have not provided their setting information. Hence, we have put “N/A: Not Available” in place of their computing mode. It can be observed that our method outperforms state-of-the-art methods by a significant margin in terms of training time. We ran our method on a workstation having Intel Xeon E5-2630 v4 CPU with NVIDIA Tesla K80 GPU. The training times of our method were 127.113 and 231.164 seconds for the RIM-ONE-r3 and the SCRID datasets, respectively.

C. Quantitative Results

We evaluated the segmentation performance of our technique using the Dice coefficient (F1 score), which is used by popular methods like [42], [43]. Dice coefficient, represented here as DC, is calculated using Eq. (14):

$$DC = \frac{2|A \cap B|}{|A| + |B|} \quad (14)$$

TABLE I: The training time of our method compared with some of the state-of-the-art methods. (N/A: Not Available)

Method	Training Time (seconds)	Computing Mode
Fu <i>et al.</i> [3]	5241 [¶] / 18759 *	GPU
Sevastopolsky [13]	4623 [¶] / 16247 *	GPU
Maninis <i>et al.</i> [11]	11200 [‡]	N/A
Zilly <i>et al.</i> [12]	3296 [‡]	N/A
Al-Bander <i>et al.</i> [14]	≈ 54000 [†]	GPU
Our Proposed	127.113 [¶] / 231.164 *	GPU

The [†], [‡], [¶] symbols represent ORIGA, DRIONS-DB, and RIM-ONE-r3 datasets, respectively. The * symbol represents the SCRID dataset.

where, A is the predicted segmentation map, and B is the ground truth segmentation map.

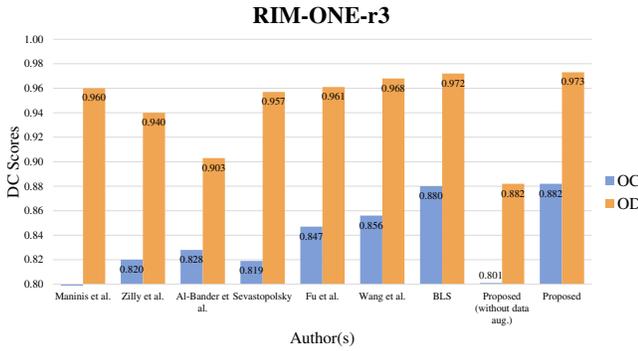


Fig. 6: The DC scores of our method, [11], [13], [3], [12], [14], and [32] for OD and OC segmentation on the RIM-ONE-r3.

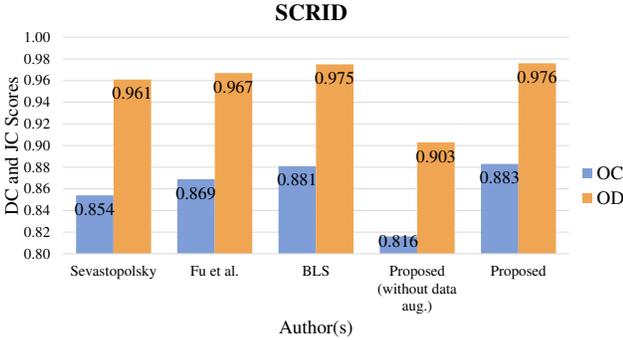


Fig. 7: The DC scores of our method, [13], and [3] for OD and OC segmentation on the SCRID dataset.

Fig. 6 and Fig. 7 show the comparison of OD and OC segmentation results of our method with other works in the literature. The comparison consists of the DC scores. As discussed earlier, we re-ran the codes of [3] and [13]. The data for other methods is taken from [13], [14], [32]. The DC score of our method for OD segmentation for the RIM-ONE-r3 dataset is 0.973. In addition to this, our proposed method achieves good results on OC segmentation as we have employed an entire FBLS-based neural network to segment OC. As shown in Fig. 6, the DC score of our approach for OC segmentation for the RIM-ONE-r3 dataset is 0.882. Some

methods have not reported the DC scores for OC segmentation in their papers; that is why DC scores for OC segmentation are not shown at some places in Fig. 6. The DC scores of our method for OD and OC segmentation for the SCRID dataset are 0.976 and 0.883, respectively, as presented in Fig. 7. It is evident from these results that our method outperforms state-of-the-art methods in OD and OC segmentation.

D. Qualitative Results

Fig. 8 and Fig. 9 show the segmentation results of our method in comparison to [3], [13] on RIM-ONE-r3 and SCRID datasets, respectively. The first column contains the original ROI images; the second column shows the images segmented by our method, the third column shows the result of Fu *et al.* [3], the fourth column contains the segmentation result of [13], and the fifth column shows the ground truth images. These results exhibit that our method provides better results of OD and OC segmentation when compared to the other two techniques.

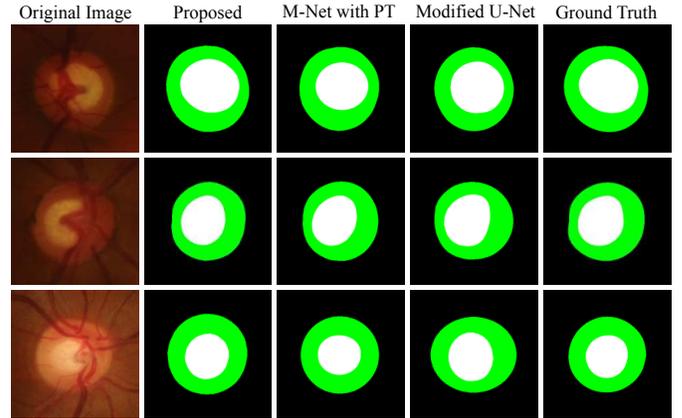


Fig. 8: Segmentation results of our method compared with [3], [13], and the ground truth images of the RIM-ONE-r3 dataset.

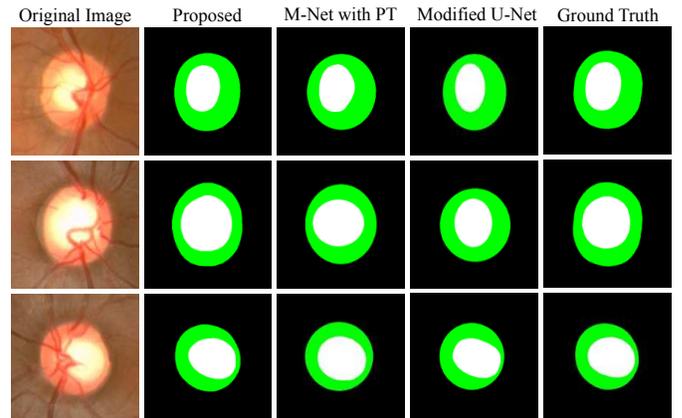


Fig. 9: Segmentation results of our method compared with [3], [13], and the ground truth images of the SCRID dataset.

E. Glaucoma Screening

We also performed a binary classification on images as being healthy or glaucoma affected with the help of computed

CDRs because CDR is an important clinical indicator of glaucoma progression. $CDR = VD_{cup}/VD_{disc}$ where, VD_{cup} and VD_{disc} respectively, represent the vertical diameters of OC and OD. Due to better segmentation of OC, we calculate the CDR more accurately, and therefore, achieve better results of glaucoma screening. We compared our classification results with the experts' evaluation results provided with the datasets and computed the accuracy of our method. Fig. 10 and Fig. 11 show the glaucoma diagnosis performance of our method by reporting the Receiver Operating Characteristic (ROC) curve with AUC scores for the RIM-ONE-r3 and the SCRID datasets, respectively. Our method achieves the best AUC scores of 0.906 and 0.923 for glaucoma screening on the RIM-ONE-r3 and the SCRID datasets, respectively.

TABLE II: Classification errors on RIM-ONE-r3 and SCRID

Dataset	M-Net+PT	Modified U-Net	BLS	Proposed
RIM-ONE-r3	0.156	0.218	0.125	0.093
SCRID	0.088	0.123	0.079	0.061

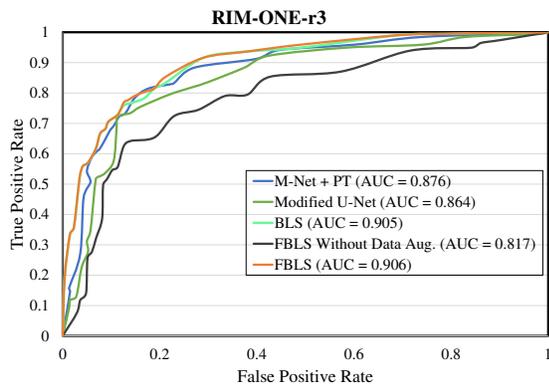


Fig. 10: The ROC curves with AUC scores for glaucoma diagnosis on the RIM-ONE-r3 dataset.

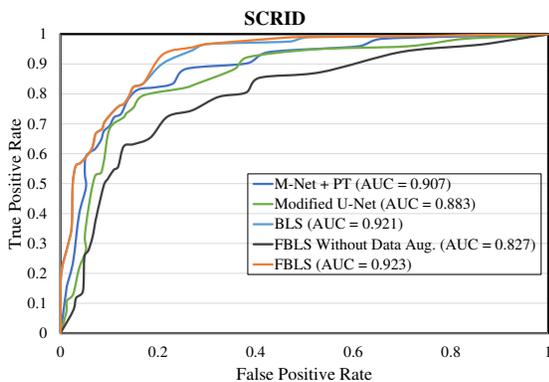


Fig. 11: The ROC curves with AUC scores for glaucoma diagnosis on the SCRID dataset.

We also present the classification errors (calculated as incorrect predictions / total predictions) in the Table II. It is

obvious that our method has lesser errors, 0.093 and 0.061 for the RIM-ONE-r3 and the SCRID datasets, respectively, than the other methods and, hence, outperforms them here as well.

F. Significance Test

To show the statistical significance in the calculated CDRs, we have conducted a non-parametric statistical test, called the Friedman test [44] with $\alpha = 0.05$, on the CDRs calculated by M-Net+PT [3], Modified U-Net [13], and our proposed method FBLs. We applied the Friedman test on the test set images, which are 20% of each dataset. Hence, the number of values becomes 32 from the RIM-ONE-r3 and 113 from the SCRID dataset. Our method FBLs achieved significantly better results by reporting $p = 0.009$ on the RIM-ONE-r3 dataset.

However, the Friedman test only tells that CDRs are statistically different, but it does not tell which group is exactly different. Hence, to see where the difference exists, we need to perform a post hoc test. We performed a post hoc test by running the Wilcoxon signed-rank test [45]. The Wilcoxon signed-rank test is a non-parametric test that finds the difference between two groups of data. So, we performed this test on different combinations of CDRs computed by these three methods as: i) FBLs to M-Net+PT ii) FBLs to Modified U-Net iii) M-Net+PT to Modified U-Net, to find which one of them is significantly different. Nevertheless, before looking at the results of Wilcoxon signed-rank, we need to consider one more thing. As we are performing multiple tests for comparison, some of these tests will give a p -value less than 0.05 by chance [46]. To overcome this, we need to apply a Bonferroni adjustment to the results obtained from the Wilcoxon tests. The Bonferroni adjustment is computed by dividing the initial significance level (0.05) by the number of tests performed. In our case, the new significance level is obtained as $0.05/3 = 0.017$ because we are performing three comparison tests. When our method is compared with M-Net+PT, the Wilcoxon signed-rank test gave $p = 0.006$, which is less than 0.017. Similarly, $p = 0.003$ is obtained when we compare our method with Modified U-Net. The comparison results of M-Net+PT and Modified U-Net were not statistically significant because $p = 0.66$ was achieved when these two methods were compared.

In the same manner, we performed the Friedman test on the SCRID dataset. Here, also, our method obtained statistically significant results by reporting $p = 0.001$. Besides, we also conducted a post hoc test with the Wilcoxon signed-rank test with a Bonferroni adjustment applied. There was no significant difference between M-Net+PT and Modified U-Net $p = 0.761$. Nevertheless, there was significant difference in the results of FBLs to M-Net+PT $p = 0.002$ and FBLs to Modified U-Net $p = 0.003$.

G. Using SCRID for Training and RIM-ONE-r3 for Testing

To further testify the robustness of our technique, we also experimented the cross-training by using the SCRID dataset for training and the RIM-ONE-r3 dataset for testing. The obtained results are shown in Table III. In these cross-training results, our approach has gotten the best DC score of 0.856

for OC segmentation, along with the second-best DC score for OD segmentation and the second-best AUC score of 0.953 and 0.860, respectively. By looking at these results, we can claim that our method provides competent results on unseen images when compared with the other state-of-the-art methods.

TABLE III: DC and AUC scores of cross-training.

Technique	DC Score for OD Segmentation	DC Score for OC Segmentation	AUC
Sevastopolsky [13]	0.939	0.813	0.845
Fu <i>et al.</i> [3]	0.948	0.838	0.864
Wang <i>et al.</i> [32]	0.959	0.847	0.853
BLS	0.951	0.849	0.856
Proposed	0.953	0.856	0.860

H. Results on the REFUGE and the DRISHTI-GS1 Datasets

We also performed the experiments on the REFUGE [42] and DRISHTI-GS1 [43] datasets to show the strength and potential of our method. The REFUGE dataset consists of 1200 images equally divided into three parts: training set, validation set, and test set (each of the three parts contains 400 images). The resolution of training set images is 2124×2056 pixels, and the validation and test images are of the 1634×1634 pixels size. We trained our technique on the training set and evaluated it on the test set images. Then, we compared our results with the results of the top three teams from the REFUGE challenge [42] organized in conjunction with MICCAI 2018. The DC scores compared with the best three methods in terms of OD and OC segmentation are reported in Table IV. Our method obtained the best DC scores of 0.9743 and 0.8845 for OD and OC segmentation, respectively.

TABLE IV: Comparison of DC scores on the REFUGE.

Technique	For OD Segmentation	For OC Segmentation
Team CUHKMED	0.9602	0.8826
Team Masker	0.9464	0.8837
Team BUCT	0.9525	0.8728
BLS	0.9621	0.8823
Proposed	0.9743	0.8845

We also compared the AUC scores of our method with the top three methods in terms of the best AUC scores. Those results are presented in Table V. Here, our approach obtained the third-best AUC score of 0.9721. These results also demonstrate a satisfactory performance of our method on the REFUGE dataset when compared with state-of-the-art algorithms.

TABLE V: AUC scores on the REFUGE dataset.

Technique	AUC Score
Team VRT	0.9885
Team SDSAIRC	0.9817
Team CUHKMED	0.9644
BLS	0.9673
Proposed	0.9721

The DRISHTI-GS1 dataset comprises 101 images having a resolution of 2896×1944 pixels. The dataset is split into training and test set containing 50 and 51 images, respectively.

The DC scores for OD and OC segmentation and AUC score of our approach compared with [13], [3] and [32] are shown in Table VI. As seen from the results in Table VI, our method has achieved the best DC scores of 0.968 and 0.880 for segmenting OD and OC, respectively. Moreover, our technique has also gotten the best AUC score of 0.887. These results also show that our method performs better than some of the best algorithms on the DRISHTI-GS1 dataset.

TABLE VI: Comparison of our method's DC and AUC scores with state-of-the-art methods on the DRISHTI-GS1 dataset.

Technique	DC Score for OD Segmentation	DC Score for OC Segmentation	AUC
Sevastopolsky [13]	0.947	0.850	0.849
Fu <i>et al.</i> [3]	0.953	0.861	0.864
Wang <i>et al.</i> [32]	0.965	0.858	0.858
BLS	0.964	0.865	0.870
Proposed	0.968	0.880	0.887

I. Segmentation in Noisy Images

The presence of noise in the retinal images makes it hard to perform an accurate segmentation of OD/OC. We tested the performance of our technique on noisy images, and one sample segmentation is shown in Fig. 12. These results also indicate that our approach generates decent results on noisy images.

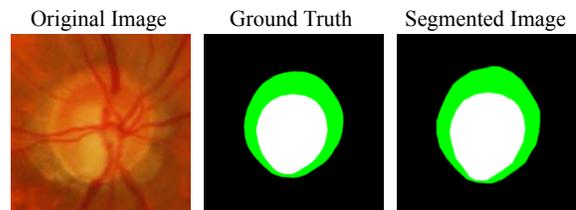


Fig. 12: A sample of the performance on a noisy image.

J. Limitation

One of the limitations of our method is that it requires pre-processing and post-processing for providing optimal results. Another limitation may be the need to extract individual channels to perform the segmentation (red channel for OD and green channel for OC segmentation), which is unlike some of the other methods which directly work on RGB fundus images.

V. CONCLUSION

In this paper, we have presented a novel technique of OD and OC segmentation, which adopts the fuzzy broad learning system. The proposed approach outperforms the state-of-the-art techniques in terms of network training time, OD and OC segmentation, and glaucoma screening results. The main advantage of our method is the efficient training process due to which it can be trained even on machines without having a GPU. Because of the fast training, our proposed work uses an entire FBLN-based neural network for OC

segmentation, which helps it attain more accurate OC segmentation results. The experiments performed on the RIM-ONE-r3 and the SCRID datasets show promising results of OD and OC segmentation. Additionally, CDR was calculated with the help of the segmented OD and OC. With the help of the calculated CDR, our method provides better performance on glaucoma diagnosis as compared to many contemporary glaucoma screening algorithms.

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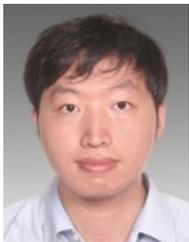
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