

Diabetic Retinopathy: Evaluation and Analysis using Computer Method

Anil Kumar¹, Shelly Kataria², Anita Singh Banafar³, Rashmi Mishra⁴, Sushil Shukla⁵

¹Department of Applied Sciences and Humanities (Mathematics)

IMS Engineering College Ghaziabad, UP India

²Chandigarh Group of Colleges, Landran, Mohali, Punjab, India

³Department of Applied Mathematics, Jabalpur Engineering College Jabalpur MP India

⁴Department of Applied Science and Humanities,

GL Bajaj Institute of Technology and Management Greater Noida UP India

⁵Department of Mathematics, Faculty of Engineering and Technology.

Veer Bahadur Singh Purvanchal University, Jaunpur, UP India

Email ID: dranilkumar73@rediffmail.com

Cite this paper as: Anil Kumar, Shelly Kataria, Anita Singh Banafar, Rashmi Mishra, Sushil Shukla, (2025) Diabetic Retinopathy: Evaluation and Analysis using Computer Method. *Journal of Neonatal Surgery*, 14 (32s), 4739-4747.

ABSTRACT

This study suggests novel approaches for solving retinopathy diabetes through computational methods. Diabetes is the primary cause of this widespread retinal infection, which is a leading cause of vision impairment in middle-aged and older adults. In order to successfully monitor the infection's course, early identification and detection by routine screening and appropriate treatment would be highly advantageous. Retinopathy is now detected via a labor-intensive, time-consuming process that mostly depends on a doctor's skill. Automated detection of diabetic retinopathy is required to overcome these problems. Diagnosis of diabetic retinopathy also depends on early detection because, with appropriate treatment, it can prevent blindness.

In order to recommend lone sick individuals to a master for extra care and supervision, a robotized prediction framework for diabetic retinopathy upgrades in the eye is necessary because there are more people with the condition than eye experts who can scan them. Shaded photographs of the retina are used to analyze the angles and stages of retinopathy. An great tool for compelling Diabetic Retinopathy screening is image recognition, which may be used to detect these many symptoms and phases of the disease in a robotized manner. Additionally, it can send the patient to a professional for aid. A unique approach that detects the early indicators of diabetic retinopathy and classifies them into distinct groups is proposed in this work.

This method can be used to confirm or disprove the detection in medical situations. The approach created here could also be used to combine detecting expertise for the benefit of humanity. The model gave the highest possible accuracy. Therefore, a plan that has been implemented may help people with diabetic retinopathy avoid becoming completely blind.

Keywords: Neural Networks, Convolutional Neural Networks, Retinopathy diabetes, Prediction, Analysis, CNN.

1. INTRODUCTION

An estimated 463 million people worldwide suffer from diabetes mellitus, an important health problem that is projected to impact 700 million by 2045 [18]. Diabetic retinopathy (DR), the most common diabetic eye circumstances, affects at least one-third of individuals with diabetes [19]. The global burden of diabetic retinopathy (DR) continues to worsen and DR remains a leading cause of vision loss worldwide. Here, we describe an algorithm to predict DR progression by means of deep learning (DL), using as input color fundus photographs (CFPs) acquired at a single visit from a patient with diabetic retinopathy (DR). Innovations in digital health solutions are increasingly acknowledging the critical role that computational platforms, sensors, software, and communication technologies play. In addition to making mass screening possible, these technologies may improve diagnostic precision. Technology in the field of digital health is developing quickly, particularly when it comes to managing diseases like diabetes. With the ability to identify complex correlations in incoming data and benchmark against performance standards, complex computational skills make pattern recognition possible, which is an important advancement. The one of the most prominent and deceptive micro vascular outcomes of diabetes is diabetic retinopathy (DR), which can worsen absent signs or symptoms until it suddenly results in blind. Within 20 years of the onset

of diabetes, retinopathy develops in almost all people who suffer from type 1 diabetes and around 60% of patients with type 2 diabetes. But until it reaches a point when it poses a serious threat to vision, DR frequently remains undetected.

Worldwide, diabetes affects one in twelve persons. As a result, managing the disease requires periodic choices that are frequently quite stressful. The most common problem in the world that can still be prevented from blindness is diabetic retinopathy. High blood sugar affects retinal blood vessels, causing bleeding or obstruction, which is the cause of this vision impairment. A major visual impairment risk factor is increased by neovascularization, a sign of proliferative diabetic retinopathy (PDR).

In order to identify and apply ocular diseases, Seth et al. [6] carried out computer analyses. The terms "NPDR" and "PDR" refer to two different types of diabetic retinopathy: PDR is an advanced stage of the disease that clearly shows neovascularization and is associated with a higher which causes abnormal blood vessels to form on the optic disc or other parts of the eye. Vision-impairing

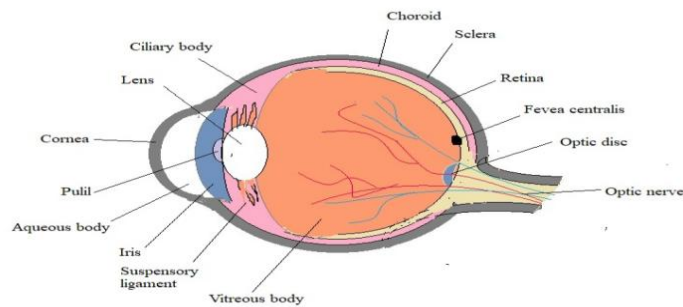


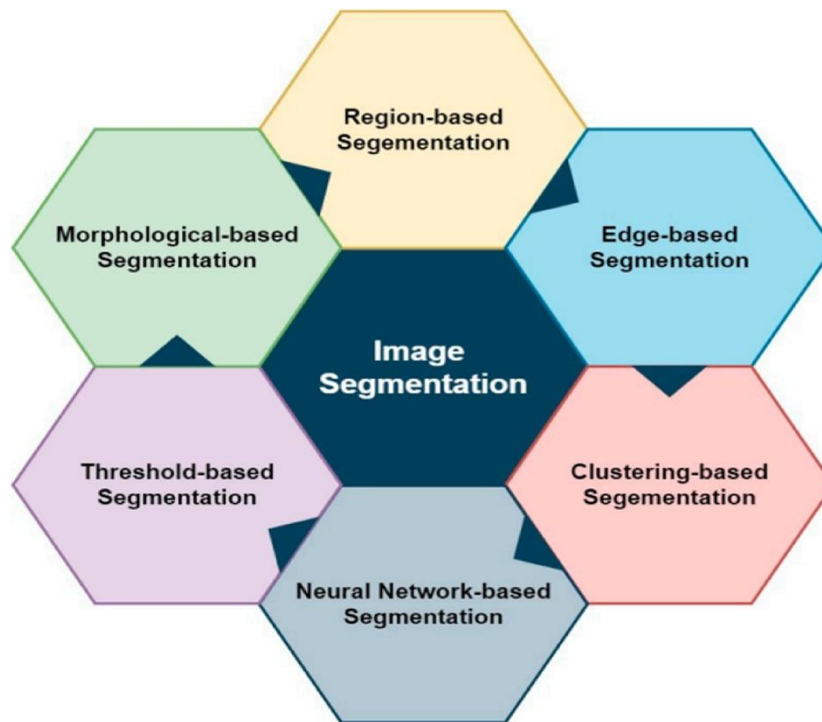
Fig1: Anatomical Structure of Eye risk of visual impairment. NPDR is defined as early stage retinal alterations without neovascularization.

The damage caused by hyperglycemia to the retinal vasculature begins with wall thickening, microaneurysm formation (which is the formation of tiny outpouchings of the artery lumens), and finally rupture, which is followed by intraretinal hemorrhages that are stopped by the internal limiting membrane (ILM). These bleedings, which are referred to as "dot and blot" bleedings because of their speckly appearance, cause the vessels to burst, letting fluid seep into the retina [3]. When people with diabetic retinopathy (DR) have macular edema, which is defined by fluid collection beneath the macula and disturbs its natural architecture, it frequently results in visual impairment.

Fluid accumulations may leave behind residues that resemble streams receding after a storm; these deposits are called hard exudates and are yellow, waxy deposits made of residual lipids. Cotton wool spots (CWS) are fluffy, white patches that appear as non-proliferative diabetic retinopathy (NPDR) worsens. Affected vessels receive chronic damage that may result in nerve fiber layer infarcts.

People with diabetes [4] are at risk for acquiring diabetic retinopathy, which includes proliferative diabetic retinopathy (PDR). People with metabolic syndrome and other pre-diabetic metabolic disorders may also be affected. The development of abnormal microvasculature, or neovascularization, within the eye is the defining feature of this condition. Fibrous proliferation is also present in the retina, the layer of the eye that is sensitive to light, and the surrounding vitreous humor, which is a gel-like substance that supports the retina and keeps it apart from the lens. In diabetic retinopathy, neovascularization results from retinal ischemia, or inadequate blood flow to the retina, intraocular hemorrhages may occur from these aberrant arteries leaking into the vitreous fluid or retina. Due to the disease's abnormal development of arteries characteristic, proliferative retinopathy carries an important chance of macular edema. Macular edema is the most common cause of visual impairment in people of working age. It is caused by fluid leaking from arteries and affects central vision in the macula, the part of the eye responsible for acceptable, detailed vision.

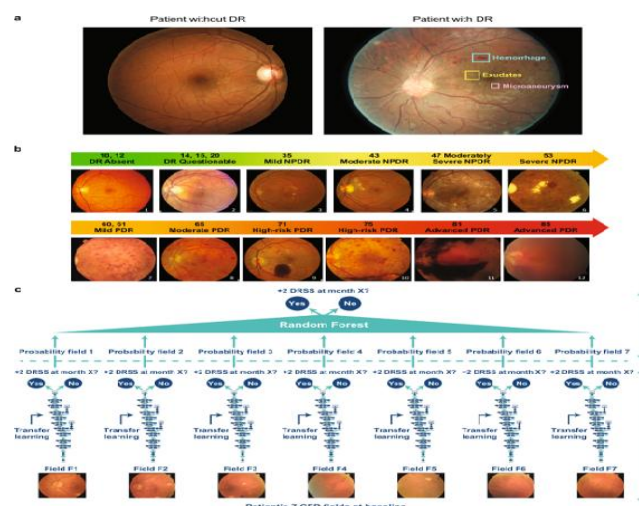
Additionally, the development of novel treatments designed for patients with mild and moderate NPDR may be aided by the identification of rapid DR-progressing individuals using a predicted DR progression algorithm. Due to their scope and/or duration, clinical trials that aim to improve or aggravate DR have been deemed excessively costly. Such an AI-based recruitment strategy would increase the likelihood of success for clinical trials of novel drugs envisioned to address the unmet need of those members of the early DR population at the greatest risk of progression and vision loss by enriching the clinical trial population with fast-progressing individuals. This is especially critical in light of the growing prevalence of DR worldwide and its potential impact on social structures and health care systems.



Flow chart of Segmentation Image

2. INVESTIGATIVE TECHNIQUES.

Detection, diagnosis, and analysis of diabetic retinopathy (DR) with computer assistance have become more common in recent years, reducing the workload for ophthalmologists and addressing diagnostic variability. Because of their superior performance in evaluating and interpreting medical imaging, especially ophthalmic pictures, neural networks—more specifically, Convolutional Neural Networks (CNNs)—have gained popularity. A CNN is a particular kind of feed-forward neural

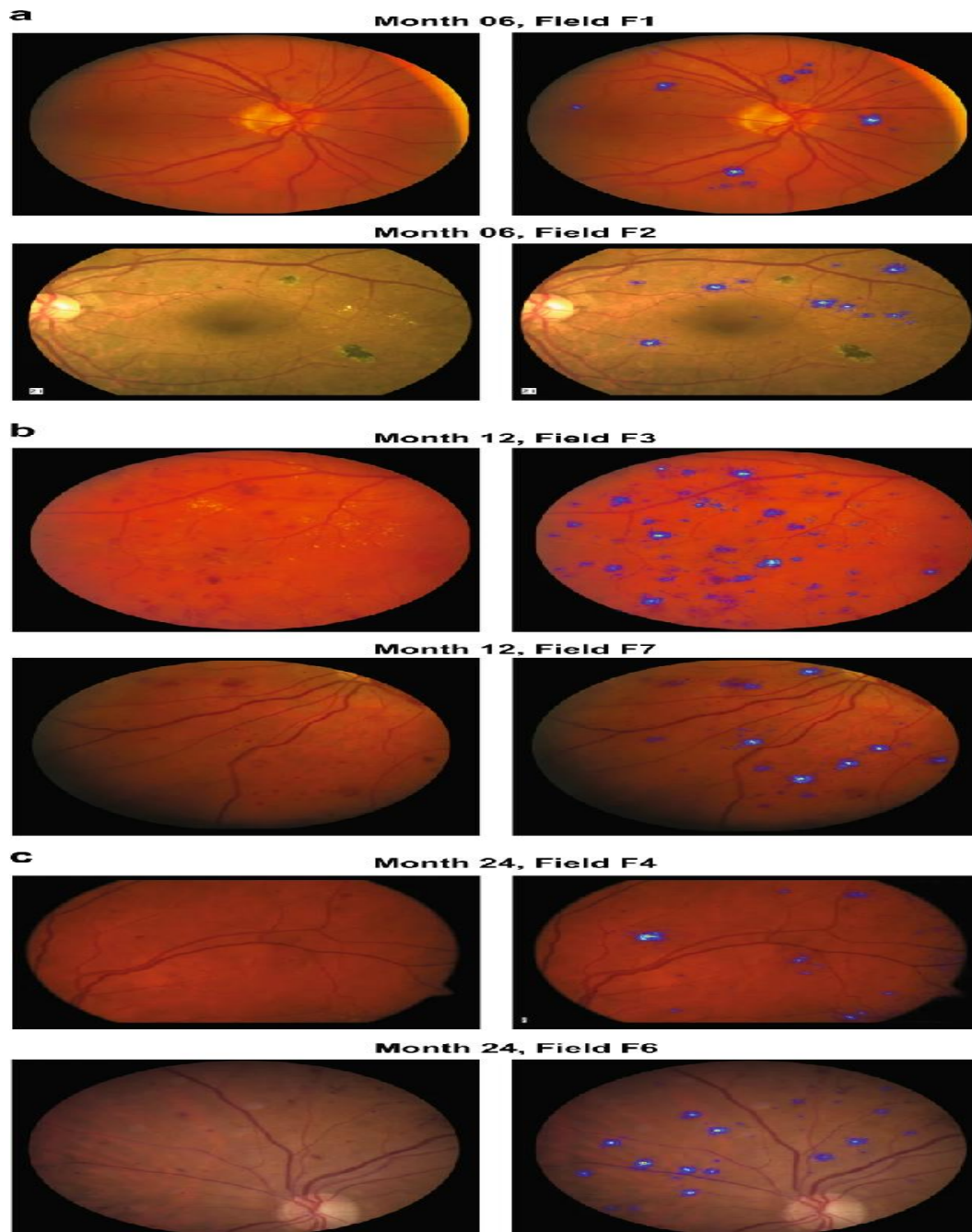


Conventional Neural Networks is the main architecture used for image recognition and detection functions.

network that consists of an input layer, several hidden layers (such pooling, convolutional relu, flatten, fully connected, and softmax layers), and a multi-label classification layer at the end [5]. These networks are especially good at executing challenging computer vision tasks with remarkable accuracy, which frequently involve many object classes. The number and arrangement of layers in a CNN's architecture can be changed to improve model performance. CNNs are convenient for programmers to create and customize for medical image analysis due to their versatility and comprehensibility. Clinical judgment and patient care results have been enhanced as a result of their ability to quickly interpret and categorize images,

greatly advancing automated systems for identifying and diagnosing diabetic retinopathy.

An illustration of an attribution map overlaying an original test color the fundus image. The attribution map is on the right in each collection, and the original image is on the left. The primary focus of deep convolutional neural networks' attribution is on exudates, hemorrhages, and microaneurysms. Additionally, there are two occurrences of attribution maps for the model that forecasts the progression of diabetic retinopathy (DR) at month six. b Two attribution map examples for the AI model predicting the progression of DR at month 12. c Two attribution map examples for the model that predicts the development of DR at month 24.



Images of various time in moths

3. DATA PRE PROCESSING

Data pre-processing is the first step in data analysis, and it is essential to prepare picture data for further analysis. Using the 'cv2.imread' function, images are first imported, and the pixel values of each image are saved in an array. Next, every picture is scaled to a consistent 50x50 pixel size and converted into a grayscale format. To guarantee consistency throughout the

collection, the image size must be standardized. Grayscale conversion decreases the number of pixels by condensing the image to a single channel, as opposed to RGB images, which contain three color channels. This reduction keeps pertinent visual information for analysis while reducing computing complexity. Each image's label data is kept in a different numpy array.

The dataset is then divided into training and testing subgroups. Thirty percent of the data are kept aside for testing, with the remaining seventy percent going toward the training set. This division keeps unseen data for assessing the model's capacity for generalization while ensuring that it is trained on a sizable enough dataset. The visual arrays' pixel values are normalized to improve computing speed and numerical stability. Although the original data's pixel values range from 0 to 255, each pixel value is scaled by dividing it by 255. Pixel values have been converted to a standard range between 0 and 1 by this transformation, making computation easier and more effective for later phases of data analysis and model training.

4. CNN DEVELOPMENT

A network architecture for deep learning that learns directly from data is called a convolutional neural network, sometimes known as a ConvNet or CNN. Whether it involves detecting objects, classes, and categories in photographs, CNNs are especially helpful in identifying patterns. Furthermore, they are quite good in distinguishing signal, time-series, and audio data.

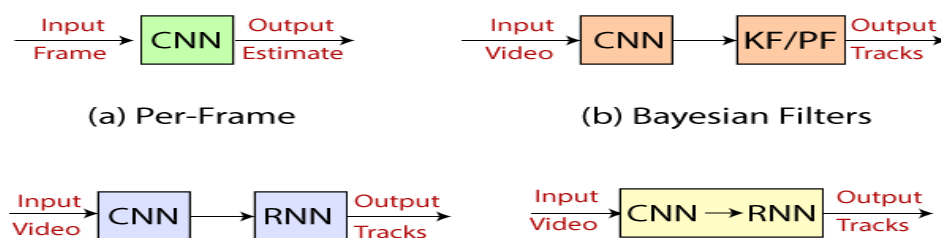
A structure with learnable weights and biases at each neuron distinguishes Convolutional Neural Networks (CNNs) from standard Neural Networks. Processing of grayscale photos scaled to 50x50 pixels is done at the input layer first. In order to extract features and preserve them in a multi-dimensional array, a 2D convolutional layer convolves over the input pictures using a 3x3 pixel window. After every convolutional layer, a rectified linear activation function is implemented. The computation of the weighted total of inputs is a critical step in selecting whether to activate a neuron. In order to preserve linearity and make optimization using gradient-based techniques simpler, ReLUs return the input value directly if it is positive or zero otherwise.

CNNs are able to effectively generalize while maintaining computational efficiency because of these qualities. Max-pooling layers are used to gradually reduce spatial dimensions and minimize parameter count. These layers provide an output matrix where each element represents the most significant value of its corresponding zone in the input by aggregating the highest values within specified filter zones (e.g., 2x2). Dropout layers are included to reduce over fitting. During training, they randomly eliminate about 45% of the connections among neurons, increasing the robustness of the model. The model architecture also makes substantial use of batch normalization (BN) layers[5,6,7,8,9,10,11, 12, 18].

By establishing the output distribution of neurons, BN enhances the performance, stability, and training speed of neural networks through allowing more seamless activation function operations. After the output has been flattened, it is processed via a number of thick layers, with a sigmoid activation function emerging in the final layer. This function is well for jobs requiring binary classification since it scales each output value between 0 and 1. Ultimately, binary cross-entropy loss is utilized to construct the model and align it with the sigmoid activation function that is employed in the output layer.

5. ALGORITHM OF CNN AND RNN MODEL

Pattern recognition and image processing both heavily depend on CNN, an effective recognition method. It includes a lot of features, including adaptability, an easy design, and few training parameters. It's now an established area in image and speech detection.



The model's algorithm performs in the following steps, which are outlined below:

Step 1 Layers Class

conv2d (Conv2D) (None, 48, 48, 256)
activation (Activation) (None, 48, 48, 256)
max_pooling2d (MaxPooling2D) (None, 24, 24, 256)

Step 2: Conv2d feature_map

We can now preserve down sampling through using max pooling global (48, 48, 256) on our 4x4 maps of functions.

Dropout: Dropout:

Conv2D=inputLayer

Activation=Conv2D

conv2d_1 (Conv2D) (None, 22, 22, 256)

activation_1 (Activation) (None, 22, 22, 256)

Step 3: We will now include a classification layer. Using 4x4 function maps, it's time to implement global max pooling.

dense (Dense) (None, 64)

dense_1 (Dense) (None, 1)

activation_2 (Activation) (None, 1)

6. SEGMENTATION OF BLOOD VESSELS

There are four stages that diabetic retinopathy passes through. In the first stage, the retina's tiny blood vessels acquire tiny swellings that resemble balloons and are known as microaneurysms. The veins delivering blood to the retina narrow and reduce blood flow in the moderate non-proliferative retinopathy (NPR) stage. Blood flow to certain regions of the retina is further restricted as the disorder develops and further blood vessels clog. The retina's afflicted regions send signals to encourage the formation of new blood vessels for nourishment in reaction to this reduction in blood flow.

The material that fills the interior of the eye, known vitreous humor, is filled with these new vessels as they proliferate over the surface of the retina. On the other hand, considerable vision loss and blindness may result if these newly formed blood vessels hemorrhaging blood.

(i) Splitting an optical disc

The vein is recognized as the optic plate's conclusion and passageway (OD). Constraints and division are essential duties for a robotized retinal imaging framework. In this sense, the area of OD is readily designated as the crucial center.

(ii) Spreading Red Lesions

Diabetic retinopathy (DR) is predominantly detected by red lesions, including hemorrhages and microaneurysms.

In order to preserve patients' vision, it is essential to identify these red lesions early on and automatically. First, color correction and then vein extraction are the three essential phases in determining the presence of red lesions. Multiple characteristics are obtained from fundus pictures of the eye in this research by using image capture and processing techniques. Direct retinal vision is possible with fundus photography. Detecting and extracting diabetic retinopathy features from RGB color-patterned fundus images is the first stage in the four-phase proposed method.

These images highlight significant characteristics such blood vessels, optic spinal discs, clustered vessels, hard exudates, and microaneurysms. These features are essential markers that determine whether diabetic retinopathy is present in an input image. The classification depends entirely on whether the input image contains hard exudates, clustered arteries, or microaneurysms. Diabetic retinopathy is the diagnosis made for the patient if any of the above signs are observed.

We utilized the reliable image processing features of MATLAB 2020a for this purpose. By making it easier to extract and analyze these crucial characteristics from fundus pictures, the Image Processing Toolbox involved into MATLAB 2020a enables for the efficient classification of normal and pathological retinal diseases using image processing techniques.

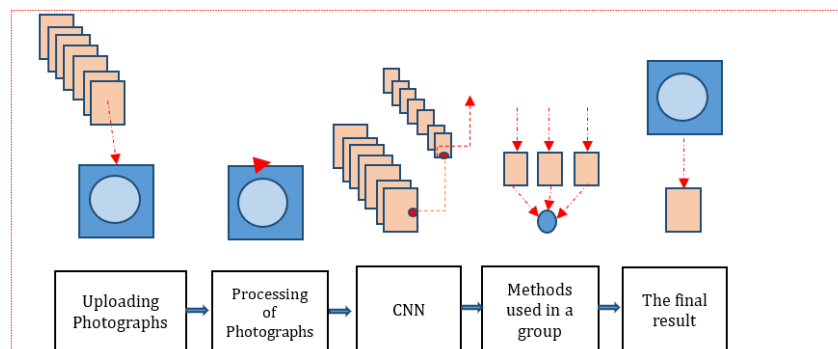
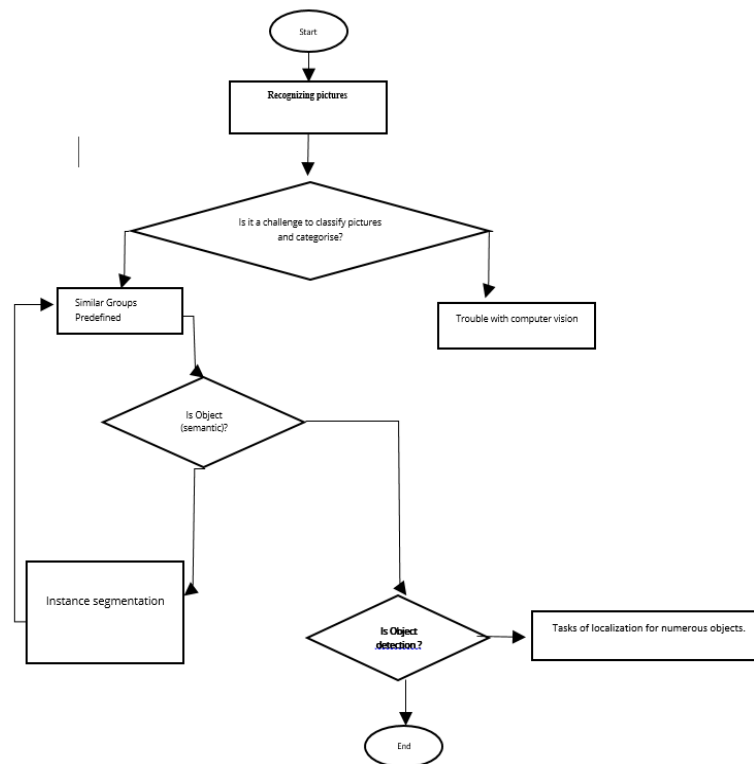


Fig 2: Diabetic retinopathy process flow

Figure 2 in image detection and recognition software, convolutional neural networks, or CNNs, are the most commonly employed architecture. Each of the several layers which make up these neural networks uses groups of neurons to process discrete areas of a picture. A complete representation of the full image is created by partially overlaying the outputs of each layer. The network is able to extract hierarchical information from the image structure by having successive layers repeat this process on the newly created representation. GPUs speed up the processing of intricate neural network operations, which is a major benefit for CNNs [3]. The number of photos and videos in training data sets has increased as digital cameras have rapidly advanced and become more accessible to everyone.

correspondingly.



7. RESULTS AND DISCUSSION

A rapid and efficient algorithm for automatically localizing the optic disc in retinal fundus images involves capturing typical and atypical images for analysis. Following image acquisition and processing, we performed statistical analysis by plotting histograms and calculating metrics such as mean, median, standard deviation, and others for the images under consideration

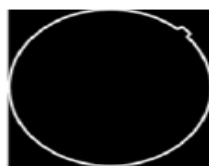


Fig3: Diabetic retinopathy is detected via edge detection.



Fig4: Detecting vessels to diagnose diabetic retinopathy



Fig 5: Diabetic Retinopathy can be detected using hard exudates.

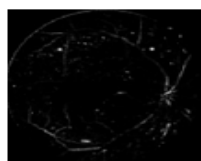


Fig 6: Diabetic Retinopathy Detection Using Micron Detection



Fig7: Diabetic Retinopathy Diagnosis using Optic Disc Detection

Based on the collected data, it has been observed that typical images exhibit comparable statistical characteristics, with mean values ranging approximately from 0.21 to 0.2165 and median values ranging from 8 to 15. Similarly, standard deviation values for typical images are also quite consistent, typically falling within the range of 0.12 to 0.13 for X-axis measurements of the plot and 30.58 to 30.48241 for Y-axis measurements. These findings indicate that typical images consistently demonstrate similar statistical attributes, which distinguish them significantly from the statistical attributes observed in irregular or anomalous eye images. This statistical analysis enables effective differentiation between typical and irregular images based on their distinct measurable characteristics.

8. CONCLUSION

Utilizing clinical trial data has the added advantage of offering high-quality, standardized imaging formats and methods, along with assessments performed by researchers in concealment at a centralized reading center. This indicates, however, that our work is now restricted to clinical trial populations falling within an extent of predefined eligibility criteria. Consequently, to guarantee that these findings are repeatable and relevant to a larger DR population, validation using real-world datasets will be crucial; the authors are now addressing this issue with further research. This study introduces novel computational approaches for addressing diabetic retinopathy (DR), focusing on image recognition and detection using Convolutional Neural Networks (CNNs). CNNs are structured with multiple layers, each analyzing small sections of an image through collections of neurons. This layered approach facilitates the creation of a comprehensive image representation by overlapping outputs from each collection within a layer. Subsequent layers then refine this representation, allowing the network to learn and discern complex structural features of the image.

This framework is tailored to automate the localization of the optic disc in retinal fundus images. The study employed a dataset comprising both typical and anomalous retinal images, subjected to rigorous analysis through histogram plotting and calculation of statistical metrics such as mean, median, and standard deviation. This study introduces novel computational approaches for addressing diabetic retinopathy (DR), focusing on image recognition and detection using Convolutional Neural Networks (CNNs) [12,13,14,15,16,17]. The CNNs are structured with multiple layers, each analyzing small sections of an image through collections of neurons. This layered approach facilitates the creation of a comprehensive image representation by overlapping outputs from each collection within a layer. Subsequent layers then refine this representation, allowing the network to learn and discern complex structural features of the image. Python programming language was utilized to implement the CNN architecture, which follows a basic structure:

of the optic disc in retinal fundus images.

By detecting initial signs of diabetic retinopathy (DR) categorized into various stages, this method holds potential for clinical validation and intervention. Moreover, the developed approach could be expanded to consolidate knowledge in DR detection, contributing to advancements in medical diagnostics and patient care. By detecting initial signs of diabetic retinopathy (DR) categorized into various stages, this method holds potential for clinical validation and intervention. Moreover, the developed approach could be expanded to consolidate knowledge in DR detection, contributing to advancements in medical diagnostics and patient care. The input data in a CNN model is processed through a number of layers that are designed to extract successively abstract features before completing the classification. Convolutional layers and pooling layers are the fundamental elements of a CNN. Convolutional layers use filters to extract features from the input data, while pooling layers down sample the convolutional layers' output to lower the dimensionality of the data.

REFERENCES

- [1] Shegdar, S., Bhatlavande, A., Patil, D., & Kadam, S. (2021): Automated Early Detection of Diabetic Retinopathy. In *Techno-Societal 2020*, pp. 897-904, Springer publication.
- [2] Oda, M., Yamaguchi, T., Fukuoka, H., Ueno, Y., & Mori, K. (2020): Automated eye disease classification method from anterior eye image using anatomical structure focused image classification technique, *Proceedings of SPIE Medical Imaging 2020: Computer-Aided Diagnosis*, Vol.11314, 1131446.
- [3] Meiburger, K. M., Acharya, U. R., & Molinari, F. (2018): Automated localization and segmentation techniques for B-mode ultrasound images: A review. *Computers in biology and medicine*, vol. 92, pp 210-235.
- [4] Qiao, L., Zhu, Y., & Zhou, H. (2020): Diabetic retinopathy detection using prognosis of micro aneurysm and early diagnosis system for non-proliferative diabetic retinopathy based on deep learning algorithms. *IEEE Access*, vol. 8, pp 104292-104302.
- [5] Abbas, S. M., & Singh, S. N. (2018): Region-based object detection and classification using faster R-CNN, 4th International Conference on Computational Intelligence & Communication Technology pp. 1-6, IEEE. 2018.
- [6] Yu, K.-H., Beam, A. L. & Kohane, I. S. Artificial intelligence in health care. *Nat. Biomed. Eng.* **2**, 719–731 (2018).
- [7] Ting DSW, Cheung CY, Lim G, et al.(2017): Development and Validation of a Deep Learning System for

Diabetic Retinopathy and Related Eye Diseases Using Retinal Images From Multiethnic Populations With Diabetes. *JAMA*, 318(22):2211–2223.

- [8] A. Kwasigroch, A. Mikolajczyk and M. Grochowski (2017): Deep convolutional neural networks as a decision support tool in medical problems - malignant melanoma case study, *Trends in Advanced Intelligent Control Optimization and Automation*, pp. 848-856.
- [9] Y. LeCun, K. Kavukcuoglu and C. Farabet (2010): Convolutional networks and applications in vision", *Proceedings of 2010 IEEE International Symposium on Circuits and Systems*, pp. 253-256.
- [10] V. Ramya, R. Jayaparvathy (2023): A New Technique using Deep Learning to Segment Retinal Blood Vessels, *2023 2nd International Conference on Automation, Computing and Renewable Systems (ICACRS)*, pp.1405-1410.
- [11] Jiajia Li, Chenhao Lin, Zikai Wang, Haitao Song, Feng Tan, Lun Hu, Pengwei Hu,(2023) : MLGL: Model-free Lesion Generation and Learning for Diabetic Retinopathy Diagnosis, *2023 IEEE International Conference on Bioinformatics and Biomedicine (BIBM)*, pp.1266-1271.
- [12] Dharsinala Harikrishna, Naluguru Udaya Kumar (2022): Artificial Intelligence System for Classification of Diabetic Retinopathy", *2022 Smart Technologies, Communication and Robotics (STCR)*, pp.1-5..
- [13] T. Rajasekar, Albert Joshwa A, M Deepa, A I Kaaviya(2022):Graphical User Interface for Classification of Diabetic Retinopathy using Image Processing", *2022 4th International Conference on Inventive Research in Computing Applications (ICIRCA)*, pp.1789-1796.
- [14] Marcello D. Giammarco, G. Iadarola, F. Martinelli, F. Mercaldo, A. S.(2022):Explainable Retinopathy Diagnosis and Localisation by means of Class Activation Mapping", *2022 International Joint Conference on Neural Networks (IJCNN)*, pp.1-8.
- [15] Sashank S., Sowmya S. (2020): Detection and Prognosis Evaluation of Diabetic Retinopathy using Ensemble Deep Convolutional Neural Networks", *2020 International Electronics Symposium (IES)*, pp.78-85.
- [16] Sheena P.; Sindhu K.; Vishalbalaji S., Saranya J.; Palanivelan M.(2023) :Efficient CNN based detection of diabetic retinopathy, *AIP Conf. Proc.* 2829, 060001.
- [17] Reguant, R., Brunak, S. and Saha, S. (2021): Understanding inherent image features in CNN-based assessment of diabetic retinopathy. *Sci Rep***11**, 9704.
- [18] Haq, N.U., Waheed, T., Ishaq, Hassan, MA. Safie. N , , NF and Shoaib M :Computationally efficient deep learning models for diabetic retinopathy detection : a systematic literature review. *Artif Intell Rev* **57**, 309 (2024)
- [19] Alicia J. Jenkins, Mugdha V. Joglekar, Anandwardhan A. Hardikar, Anthony C. Keech, David N. O'Neal, S. Andrzej, Januszewski Biomarkers in diabetic retinopathy, *Rev. Diabet. Stud.: Reg. Dev. Stud.*, 12 (1–2) (2015), p. 159.