

# GENERATIVE ADVERSARIAL NETWORKS FOR RETINAL IMAGE ENHANCEMENT WITH PATHOLOGICAL INFORMATION

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## ABSTRACT

Age-related macular degeneration (AMD) is a disease of the central retina, which is one of the main reasons for vision loss of elderly people. To monitor the level of AMD, the doctors mainly use the retinal fundus images. However, the quality of retinal images can be affected during the imaging process. It leads to low contrast and blurry images. Those bad quality images cannot be used for analyzing and diagnosis. For that reason, there are many studies about image enhancement in order to improve the quality of retinal photography. However, previous methods could not guarantee to keep the disease information after the enhancement process. Therefore, we introduce a generative adversarial model for AMD retinal image enhancement with additional factors to preserve the disease information. By exploiting drusen segmentation masks, our proposed model can enhance retinal photography quality and keep the pathological information.

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## INTRODUCTION

Age-related macular degeneration (AMD) encompasses a variety of diseases and conditions. It is a group of optic nerve diseases, with 'characteristic' progressive structural changes leading to loss of visual function in a 'characteristic' way. AMD is the second leading cause of blindness worldwide. The retina is the innermost layer in the eye and the retinal nerve fiber transmits the visual signal from the photoreceptors in the eye to the brain via the bundle going out of the eye, known as the optic nerve. AMD leads to continuous and speedy damage of the retinal nerve fiber layer and hence can lead to permanent blindness. The progression of the nerve fiber layer loss can be effectively stopped by treatment consisting of medication or surgery to reduce the intraocular pressure. Hence the diagnosis of AMD at an earlier stage is very important for its treatment.

A major concern with AMD detection is that the disease has no particular set of physical causes or symptoms that doctors can recognize to detect the disease in an early stage. The main focus in AMD diagnosis is to detect changes in the visual functioning of the eye at early stages of the disease so that vision can be protected and preserved through medical treatment. It has been proved that the development of visual field defects is preceded by RNFL damage in Age-related macular degeneration (AMD).

Studies show that as much as 40% of retinal nerve fiber in the eye can be lost without the detection of characteristic visual defect in AMD patients. Hence it is believed that the detection of damage in nerve fiber layer can lead to an early detection of AMD. Several computer-assisted imaging technologies for detecting the structural changes in the retinal nerve fiber layer have been developed. The assessment of the ganglion cell structure is based on measuring the thickness of the retinal nerve fiber layer.

One such device, the Scanning Laser Polarimeter (GDx-VCC, Laser Diagnostic Technologies, Inc., San Diego, CA) is based on the principle of measuring the change in the polarization of light exiting the eye. The retinal nerve fiber layer consists of parallel structures of diameter smaller than the wavelength of light, which makes it a birefringent structure. A birefringent structure has the ability to change the polarization of polarized light

doublepassingit.<sup>7</sup>Theamountofchangeinthestateofpolarization(retardation)canbeThe device provides a large array of points corresponding to retinal.

This is found to be proportional to the thickness of the nerve fiber layer. This retardation(degrees)informationisconvertedtothickness(microns)throughtheconversion factor based on the histologic comparison with monkey eyes.Anew version of the device (GDx-VCC)isdesignedtoindividuallycompensatefortheeffectsofbirefringentproperties of other parts of the eye like the cornea. The device provides a large array of points correspondingto retinal neverfiberlayerthicknessateachrespectivepoint across theback of the eye. The scans thus available are in the form of 128 X 256 images or gray-level thickness maps.The goal of this thesis is to analyze these scans and develop classification techniques for them.



**Photograph of a Scanning Laser Polarimetry Device (Courtesy: Laser Diagnostics, San Diego, CA)**

The first step in a classification problem is to extract useful features from the set of given data. The original scan is used to derive patterns or features that can separate the two classes. The feature extraction step is to obtain a feature vector, a set of different useful features, which reduce the dimension of the original data while keeping all the essential information contained in the data. The next step after obtaining a distinguishable and reliable set of features is to make them statistically independent. A very effective way to achieve this end is to perform Principal component analysis on the feature set. This will help in reducing feature dimension by eliminating redundancy caused by interdependencies in the feature vector. The last step is the classification of the data set into the two classes. Fisher's Linear Discriminant Analysis (LDF) provides an easy and robust way.

## **OBJECTIVE**

The primary objective of this research is to design and implement a Generative Adversarial Network (GAN)-based approach for retinal image enhancement with a special emphasis on preserving pathological information. The proposed system aims to:

1. **Enhance Retinal Image Quality** – Improve the visual clarity, contrast, and resolution of retinal images affected by noise, blur, or low illumination.
  2. **Preserve Pathological Features** – Ensure that clinically significant features such as lesions, exudates, hemorrhages, and microaneurysms remain intact during enhancement for accurate diagnosis.
  3. **Assist Medical Diagnosis** – Provide ophthalmologists and automated diagnostic systems with high-quality images that support effective detection and classification of retinal diseases.
  4. **Develop Robust GAN Architecture** – Utilize adversarial training strategies to generate realistic, high-fidelity images while minimizing information loss.
  5. **Evaluate Performance** – Validate the enhanced images using both quantitative metrics (PSNR, SSIM, etc.) and qualitative expert evaluations to confirm diagnostic reliability.
- Dey, N., Roy, A. B.; Das, A.; Chaudhuri, S. S: Optical cup to disc ratio measurement for glaucoma diagnosis using Harris corner

In this paper, Glaucoma is physiologically described as the deterioration of optic nerve cells, and is characterized by alterations in the optic nerve head and visual field. The measurement of neuro-retinal optic cup-to-

disc ratio (CDR) is an important index of Glaucoma, as the increased excavation of the optic cup occurs because of Glaucomatous neuropathy increasing the CDR. Currently, CDR evaluation is manually performed by ophthalmologists.

Eleesa Jacob, R. Venkatesh: A Method of Segmentation for Glaucoma Screening Using Superpixel Classification

In this paper, an optic disc and optic cup segmentation is used to identify the glaucoma disease in time. In optic disc and optic cup segmentation, super pixel classification for glaucoma screening is proposed. In optic disc segmentation, histograms and centre surround statistics are used to classify each super pixel as disc or non-disc. A self-assessment reliability score is computed to evaluate the quality of the automated optic disc segmentation. In optic cup segmentation, the location information is also included into the feature space for better performance in addition to the histograms and centre surround statistics. The segmented optic cup and optic disc is then used to compute the cup to disc ratio for glaucoma screening. From the cup to disc ratio, analysis is performed to identify whether the given image is glaucomatous or not. The segmentation can be analyzed using the MATLAB.

## PROPOSED SYSTEMS

### Retinal Nerve Fiber Layer and Scanning Laser Polarimetry

The retinal nerve fiber layer is composed of about 1.2-1.5 million ganglion cell axons originating in the retina. The axons are distributed in a characteristic pattern. The axons originating in the region nasal to the optic disc as well as in the macular area run directly toward the optic nerve head, while the axons originating in the temporal section run towards the superior or inferior poles of the optic disc before converging to the nerve head. These fibers are known to be most susceptible to early AMD retinal image damage.

The peripheral ganglion cell axons travel to the optic nerve head in the peripheral position while the central axons take a more superficial path and follow the innermost part within the optic nerve head. Due to the characteristic pattern of the nerve fiber layer axons, the thickness of the nerve fiber layer on the vertical poles of the optic disc is much higher than in the nasal and temporal optic disc poles.

The importance of the detection of RNFL damage as an early sign of Age-related macular degeneration (AMD) has been confirmed by numerous studies. Hoyt and Newman first described it in 1987<sup>25,26</sup>. Histological studies show that as much as 40% of retinal nerve fiber in the eye can be lost without the detection of characteristic visual defect in AMD patients<sup>6</sup>. The findings of Sommer and colleagues showed that RNFL damage could precede visual field loss by up to 5 years<sup>27</sup>. Hence it is believed that the detection of damage in nerve fiber layer can lead to an early detection of AMD.

RNFL defects related to Age-related macular degeneration (AMD) can be either diffused or localized. Localized defects generally include slit-like or groove-like defects in the RNFL. When these slit-like defects extend to the disc margin or the wedge-shaped defects are seen as notches in the neuroretinal rim in inferior or superior regions, it is judged as a sign of AMD retinal image abnormality.

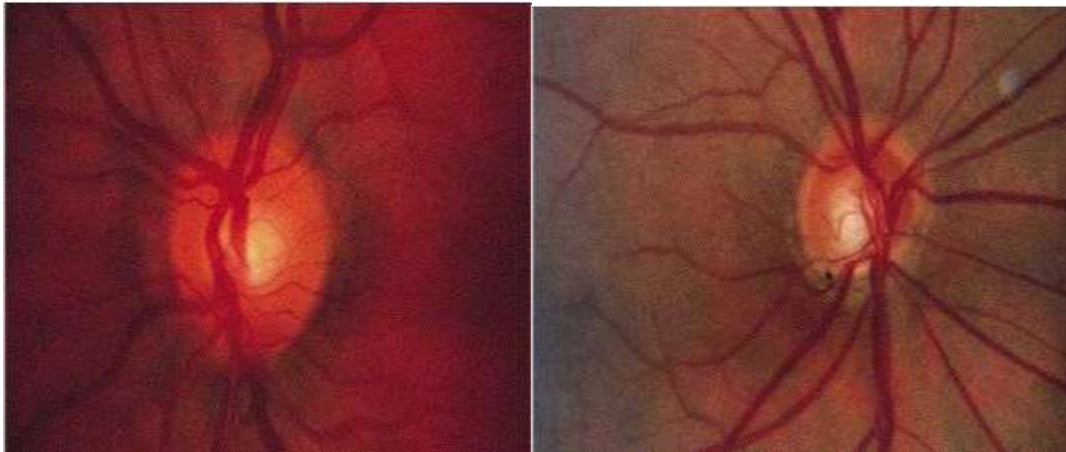
Although localized defects are easier to detect, diffuse RNFL loss is more common and difficult to diagnose. The second order retinal vessels, which are normally well concealed by the retinal nerve fiber layer, start to be seen in this kind of defects. The progressive loss of RNFL thickness in the superior and inferior poles is a sign of AMD retinal image damage.

There are several different techniques for the qualitative examination and quantitative measurement of nerve fiber damage caused by AMD. The qualitative techniques include examination of the retina through a dilated pupil using an ophthalmoscope or by using a red-free camera or using high-resolution black and white photographs. These are all, however, limited by the pupil size and media optics and tend to have high intra- and interobserver variability. To reduce these difficulties and provide more quantitative measurements of the nerve fiber layer, different devices have been developed. Several instruments have been developed that focus on imaging of the fundus (a mirror-like structure just behind the retina which acts as a light amplifier) and analyzing the topography of the retinal surface. These methods attempt to quantify the three-dimensional size and shape of the optic disc, which is considered to represent the bulk of the retinal nerve

fibers (Figure 3.1). Stereoscopic fundus photography uses photographs of fundus under different angles to obtain topographic information of the disc.

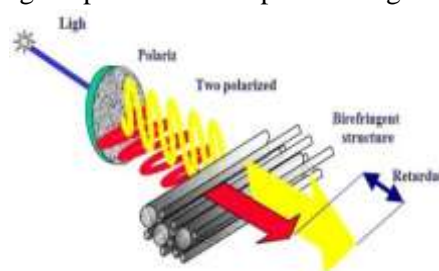
Confocal scanning laser ophthalmoscopy tries to obtain optical section images of the retina by scanning a laser beam across the eye fundus in two dimensions and provides video images on a monitor. These methods include instruments like the Topcon Imagenet, the Rodenstock Optic Nerve Analyzer (Rodenstock Instruments, Munich, Germany) and Heidelberg Retina Tomograph (Heidelberg Engineering, Heidelberg, Germany). Although these methods are a reasonable indicator of the condition of the optic disc, the analysis of the topography of the fundus is an indirect measure of the nerve fiber layer and is only suggestive.

Furthermore, the ultimate resolution of these methods is limited by the properties of the ocular media. Hence these kinds of imaging systems are not suitable for accurate measurement of the retinal nerve fiber layer thickness.



**Example of optic disc photography (a) normal disc (b) notching in optic disc (Courtesy: Handbook of Age-related macular degeneration (AMD) (Azura-Blanco Augusto))**

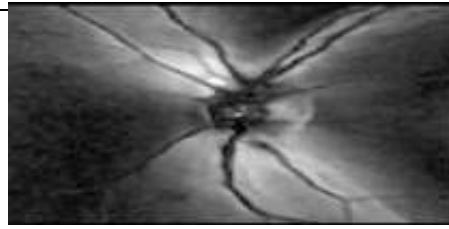
Scanning Laser Polarimetry (SLP) is a technique of providing a more quantitative measure of the thickness of the RNFL. The method is based on the principle of using imaging polarimetry to detect the birefringence of the retinal nerve fiber layer<sup>7,9</sup> (Figure 3.2). This technique utilizes the polarization properties of the retinal nerve fiber layer. The nerve fiber layer and other regions of the retina have been known to have polarization properties or birefringent properties. Form birefringence occurs when a medium consists of parallel cylindrical structure with diameters smaller than the wavelength of light passing through it. A birefringent structure has the ability to change the polarization of a polarized light double passing it<sup>29</sup>.



**Scanning Laser Polarimetry Device-Principle (Courtesy: Laser Diagnostics, San Diego, CA)**

The Scanning Laser Polarimeter uses this principle to scan the thickness of the retinal nerve fiber by employing a low power near infrared laser beam to illuminate the human retina.

Among the number of software-generated parameters provided by the company, the main ones are 'the number' and the 'NFI' (nerve fiber indicator). The parameter "number" is obtained through a neural network, which is fed with around 100-200 features from the scanned image. The NFI is obtained through a support vector machines recognizer and is available in the newer versions of the device. The latest version of the device implements the correction for the birefringent properties of the part of the eye other than the nerve fiber layer and is called GDx VCC. Currently doctors use the above-mentioned factors from the device along with other tests and gauge.



Grayscale representation of the RNFL thickness map image

## DESIGN APPROACHES ARCHITECTURE

### 1. Overall Architecture

The proposed system is built on a **Generative Adversarial Network (GAN)** framework, consisting of two main components:

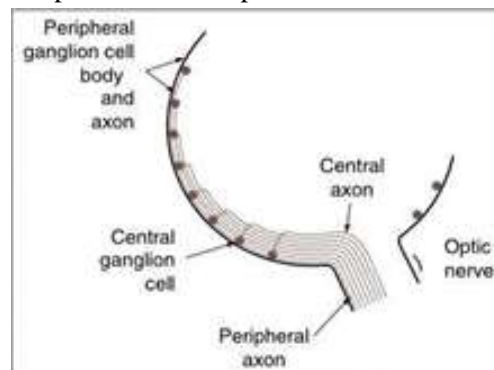
- **Generator Network**
  - Enhances low-quality retinal images.
  - Improves sharpness, illumination, and contrast.
  - Ensures pathological features such as lesions, exudates, hemorrhages, and microaneurysms remain intact.
- **Discriminator Network**
  - Differentiates between real high-quality images and GAN-enhanced images.
  - Provides feedback to improve the generator's accuracy.

### 2. Design Approach

1. **Data Input & Preprocessing**
  - Retinal fundus images acquired from scanning laser polarimetry and fundus photography.
  - Noise reduction, normalization, and augmentation techniques are applied.
2. **Feature Extraction**
  - Segmentation masks (e.g., drusen masks) highlight pathological regions.
  - Feature maps are extracted to guide GAN training.
3. **GAN Training Strategy**
  - **Adversarial Loss:** Encourages realistic image generation.
  - **Content Loss (MSE/MAE):** Maintains structural integrity.
  - **Perceptual Loss:** Preserves pathological information important for diagnosis.
4. **Validation & Testing**
  - Enhanced outputs compared using **PSNR, SSIM, and VIF**.
  - Ophthalmologists verify the medical significance of preserved features.

### 3. System Workflow (Architecture Diagram Explanation)

Although the PDF contains textual description, the conceptual architecture can be summarized as



Example of optic disc photography (a) normal disc (b) notching in optic disc (Courtesy: Handbook of Age-related macular degeneration (AMD) (Azuara-Blanco Augusto))

The ganglion axons that constitute the nerve fiber layer are essentially cylindrical rod-

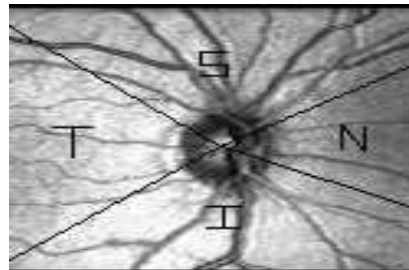


like structures that are parallel to the retinal surface and have extremely small diameters. When a light beam, perpendicular to its surface, is impinged on the retina, the reflected light is split into two rays that travel at different velocities.

## WORKING

### 1. Input Acquisition

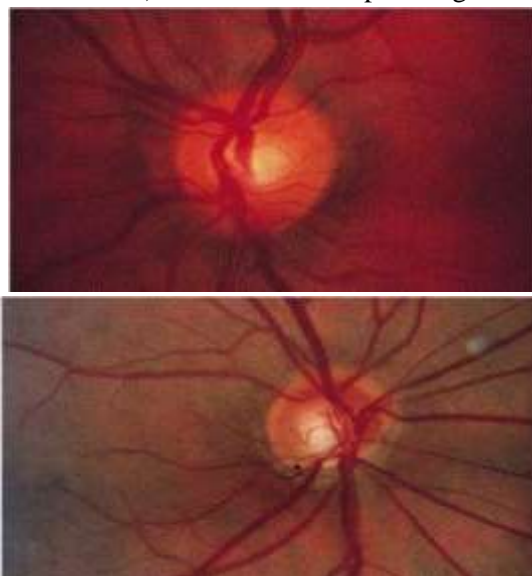
- Retinal fundus images are collected using scanning laser polarimetry and other retinal imaging devices.
- These images often suffer from low contrast, blur, and noise.



**Photograph of the optic nerve with optic disc in the center and the areas around it divided into the four sectors – Temporal, Superior, Nasal, and Inferior**

### Preprocessing

- The images are digitized and normalized.
- Segmentation masks (like drusen masks) are used to mark pathological regions for preservation



**Example of optic disc photography (a) normal disc (b) notching in optic disc (Courtesy: Handbook of Age-related macular degeneration (AMD) (Azua-Blanco Augusto))**

Scanning Laser Polarimetry (SLP) is a technique of providing a more quantitative measure of the thickness of the RNFL. The method is based on the principle of using imaging polarimetry to detect the birefringence of the retinal nerve fiber layer<sup>7,9</sup> (Figure 3.2). This technique utilizes the polarization properties of the retinal nerve fiber layer. The nerve fiber layer and other regions of the retina have been known to have polarization properties or birefringent properties. Form birefringence occurs when a medium consists of parallel cylindrical structures with diameters smaller than the wavelength of light passing through it.

### GAN Model Implementation

- **Generator:** Enhances low-quality retinal images (sharpness, contrast, clarity).
- **Discriminator:** Differentiates between real high-quality images and GAN-enhanced images.
- Training uses adversarial loss + perceptual/content loss to ensure enhancement **without losing pathological details**.

This section provides an introduction to MATLAB's desktop tools. You can also use MATLAB functions to

perform most of the features found in the desktop tools. The tools are:

- CurrentDirectoryBrowser
- WorkspaceBrowser
- ArrayEditor
- Editor/Debugger
- CommandWindow
- CommandHistory
- Launch Pad
- Help Browser

CommandWindow

Use the Command Window to enter variables and run functions and M-files.

CommandHistory

Lines you enter in the Command Window are logged in the Command History window. In the Command History, you can view previously used functions, and copy and execute selected lines. To save the input and output from a MATLAB session to a file, use the diary function.

### Processing and Enhancement

- GAN iteratively improves images while referencing segmentation masks so that disease features (lesions, hemorrhages, microaneurysms) are preserved.
- Enhanced images are visually and statistically compared with original

### □ Evaluation

- Metrics: PSNR, SSIM, VIF used for quantitative analysis.
- Ophthalmologists verify whether pathological information is retained.

### □ Output

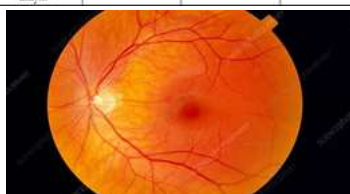
- High-quality retinal images with preserved pathological regions ready for diagnosis and computer-aided systems

### Healthy retina

### RESULT

Our present sample consists of SLP scans of 92 AMD retinal image eyes and 92 normal eyes, obtained from 47 Age-related macular degeneration (AMD) patients and 45 healthy people. Dr. Michael Sinai of

Method	Sensitivity	Specificity	Area under ROC curve
FT of 90° projection	75.12%	99.15%	0.7789
Wavelet analysis with FT of 90° projection	58.7%	89%	0.8235
Ring data from 2D FT	74.92%	91.5%	0.7813
Correlation coefficient with pattern image	67.5%	82%	0.8287
Combined Feature set	84.78%	91.3%	0.8433
FFTA – based on previously performed analysis	76.09	91.3	0.8588



### Performance evaluation of the different feature sets based on the ROC Analysis

Laser Diagnostics made the data set and its diagnosis (classification based on clinical findings) available to us for research purposes. This set was divided randomly and uniformly into two subsets. One subset was used as a training set for the classifier while the other was used as a test set. After the classification results were obtained, the ROC

analysis for performance evaluation.

## APPLICATIONS

**Detection and diagnosis of Age-related Macular Degeneration (AMD):** Utilizing computer-assisted imaging technologies to detect structural changes in the retinal nerve fiber layer for early AMD diagnosis.

**Glaucoma screening and diagnosis:** Developing automated systems for glaucoma detection using image processing and analysis of retinal images.

**Retinal image analysis:** Processing retinal images to assist clinical diagnosis and treatment, with a focus on automated diagnosis of retinal fundus images.

## ADVANTAGES

- It discusses techniques that “aid a physician in detecting possible subtle abnormalities.”
- The application of digital imaging to ophthalmology offers the possibility of processing retinal images to assist clinical diagnosis and treatment.
- The development of an automated system for analyzing the images of the retina will facilitate computer-aided diagnosis of eye diseases.
- The automated system can detect glaucoma efficiently and in less time.

## REFERENCES

- [1] J. Kim, M. Kim, H. Kang, and K. H. Lee, “U-gat-it: Unsupervised generative attentional networks with adaptive layer-instance normalization for image-to-image translation,” in International Conference on Learning Representations, 2020.
- [2] H. Fu, B. Wang, J. Shen, S. Cui, Y. Xu, J. Liu, and L. Shao, “Evaluation of retinal image quality assessment networks in different color-spaces,” Medical Image Computing and Computer Assisted Intervention MICCAI 2019, p. 4856, 2019.
- [3] H. Zhao, B. Yang, L. Cao, and H. Li, Data-Driven Enhancement of Blurry Retinal Images via Generative Adversarial Networks, 10 2019, pp. 75–83.
- [4] M. Zhou, K. Jin, S. Wang, J. Ye, and D. Qian, “Color retinal image enhancement based on luminosity and contrast adjustment,” IEEE Transactions on Biomedical Engineering, vol. 65, no. 3, pp. 521–527, 2018.
- [5] D. B. Russakoff, A. Lamin, J. Oakley, A. Dubis, and S. Sivaprasad, “Deep learning for prediction of amd progression: A pilot study.” Investigative ophthalmology and visual science, vol. 60 2, pp. 712–722, 2019.
- [6] G. Bhupendra and M. Tiwari, “Color retinal image enhancement using luminosity and quantile based contrast enhancement,” Multidimensional Systems and Signal Processing, 01 2019.
- [7] M. Zhou, K. Jin, S. Wang, J. Ye, and D. Qian, “Color retinal image enhancement based on luminosity and contrast adjustment,” IEEE Transactions on Biomedical Engineering, vol. 65, no. 3, pp. 521–527, 2018.
- [8] J.-Y. Zhu, T. Park, P. Isola, and A. A. Efros, “Unpaired image-to-image translation using cycle-consistent adversarial networks,” 2017 IEEE International Conference on Computer Vision (ICCV), Oct 2017.
- [9] C. Szegedy, S. Ioffe, V. Vanhoucke, and A. A. Alemi, “Inception-v4, inception-resnet and the impact of residual connections on learning,” in AAAI, 2017.
- [10] P. Dai, H. Sheng, J. Zhang, L. Li, J. Wu, and M. Fan, “Retinal fundus image enhancement using the normalized convolution and noise removing,” International Journal of Biomedical Imaging, vol. 2016, pp. 1–12, 01 2016.