Early Detection of Diabetic Retinopathy through Machine Learning Techniques

Manjula L, G T Raju

Abstract: Diabetic Retinopathy (DR) is progressive syndrome that leads to loss of vision if not detected and treated. Retina is inner tunic of the eyeball which is capillary and delicate transparent membrane. It is high developed tissue of eye which plays a major role for vision. Retina is the source for detection of many disorders. Part of retina with optic disc can be viewed through optamoloscope and termed as fundus image which is a basis of diagnosis for DR. DR can be categorized as Proliferative Diabetic Retinopathy (PDR), Diabetic Maculopathy, Non-proliferative Diabetic Retinopathy (NPDR) and Advanced Diabetic Eye Disease. Machine Learning (ML) techniques play a vital role in early detection of DR. In this paper a review on the existing techniques with open issues to be addressed is presented for diagnosing DR and model is proposed to consider the features namely Microaneurysms, Retinal Hemorrhages, Hard exudates, Cotton wool Spots, Neovascularization for classification of DR. These features can be combined with hypertension to predict other disorders like stroke, chronic heart disease, renal dysfunction, cardiovascular mortality and so on which overcome the need of other preliminary checkup. The complete profile of disorders for a diabetic patient can be deduced by the retinal fundus image.

Keywords: Diabetic Retinopathy, Machine Learning, Retinal Fundus Images.

I. INTRODUCTION

Retina is the innermost tunic of eyeball and the only part of Central Nervous System which can be viewed through optamoloscope. Retina contains light sensitive cells in the eyes, driving from neural tube which is derived from optic cup in turn from optic vesicle. Optical vesicle originates from diencephalon which imparts sensory information connecting brain regions in turn control several autonomic functions of peripheral nervous system. So Retina is the source for detection of many disorders.

Retina comprises of two distinct regions namely Posterior Pole and Peripheral Retina. The Posterior pole of the retina includes the Optic Disc and Macula Lutea which can be viewed through fundus images. Optic Disc is vascular and Macula Lutea is avascular.

Fundus images are the source for diagnosis of many vascular disorders of retina namely Retinal occlusions, Diabetic Retinopathy (DR), Hypertensive Retinopathy, Retinopathy of prematurity, Cardio vascular Risk, Stroke and so on. Early diagnosis is the key to slow down the progression of these diseases.

II. CLASSIFICATION OF DIABETIC RETINOPATHY

Hyperglycemia with uncontrolled DM is starting point of the development of DR [9]. Hyperglycemia is the increased levels of glucose in blood which leads to Microangiopathy (small vessel disease) which affects pre-capillary arterioles, capillaries and venules are the basic pathology in diabetic retinopathy. Effects of Microangiopathy producing DR include:

- **Breakdown of Blood** – Retinal Barrier: Retinal Oedema, Hemorrhages and leakage of lipids (hard Exudates).
- **Weakened Capillary Wall**: Aneurysms and Hemorrhages.
- **Micro vascular Oclusions**: Produce ischemia which is the limit in blood supply to the tissues, causing a lack of oxygen that is preferred for cellular metabolism) and arteriovenous shunts (intra retinal micro vascular abnormalities).

Fig. 1. Retina with DR features

DR is one of the major causes for loss of vision which refers to a retinal changes seen in patients with Diabetic Mellitus (DM) and the features are as shown in figure 1. Earlier detection is a key factor to prevent loss of vision for people with DM. The longer a person with DM has higher chance of developing DR. Other Risk factors include Heredity, Hypertension, Smoking, Obesity, Anemia and Hyperlipidaemia.

Revised Manuscript Received on December 15, 2019.

Manjula L, Department of Computer Science and Engineering, RNS Institute of Technology, Bangalore, India. Email: manjularagh10@gmail.com

Dr. GT Raju, Department of Computer Science and Engineering, RNS Institute of Technology, Bangalore, India. Email: gtraju1990@gmail.com

Published By:
Blue Eyes Intelligence Engineering & Sciences Publication
Early Detection of Diabetic Retinopathy through Machine Learning Techniques

DR has been variously classified as shown in figure 2:

![Fig. 2. Classification of DR](image)

A. Non-Proliferative Diabetic Retinopathy:
NPDR is common in Type 2 DM patients. They are identified by Microaneurysms, Retinal Hemorrhages, Retinal oedema, hard exudates, Venous Abnormalities, Cotton wool Spots, Intraretinal micro vascular abnormalities. NPDR is further classified as:
- Mild NPDR - One Microaneurysms at least must be there.
- Moderate NPDR - Microneurysms/interretinal hemorrhages in 2 or 3 quadrants, early mild Intraretinal Microvascular Abnormality (IRMA), arduous or fleecy exudates may be or not be present.
- Severe NPDR - 4-2-1 Rule i.e. Four Quadrants of Microaneurysms and extensive internal hemorrhages, Two quadrants of venous bleeding and One quadrant of IRMA changes. In which anyone of 4-2-1 is present.
- Very Severe - 4-2-1 Rule in which any two of them is present.

B. Proliferative Diabetic Retinopathy:
PDR develops with many of 50% cases after about 20 years of onset of the DM. Juvenile onset diabetes patients are more prone. Occurrence of neo vascularization at optic disk or elsewhere is the hallmark of PDR. Neo vascularization is referred to as a state in which firm pathologic vessels originates from the active retinal veins and widen along the interior surface of retina. Later on results in the formation of Fibrovascularneiretinal membrane or vitreous hemorrhage and vitreous detachment.

C. Diabetic Maculopathy:
It refers to changes in Macular region and adversely affects the vision. It involves thickening of retina or arduous exudates at or within the 500 micron of Fovea.

D. Advanced Diabetic Eye Disease:
It is the anarchic PDR. It is noticeable by complications such as continual vitreous hemorrhage, Tractional retinal separation and Neo vascular Glaucoma. Earlier detection of DR does not involve Advanced Diabetic Eye disease as it is final stage for vision loss.

III. MACHINE LEARNING TECHNIQUES
ML is programming computing device to improve a performance measure by using example data or by past experience. They are broadly categorized into Unsupervised, Supervised and Reinforcement learning. ML approaches are applied to many fields in which health applications have an extensive benchmark. Diagnosis of DR is a major yardstick.

A. Supervised Learning:
Supervised methods have great importance for medical image classification. Supervised ML is a task of data acquisition function that corresponds input to output supported on pattern input-output twain. It infuses a task from the labeled training data which consists of examples of training set.
Supervised ML algorithms include grouping and regression. When the outputs are confined to limited primed of values classification algorithms are used. Some of the approaches for diagnosis of DR are tabulated as shown in Table-I. As deduced in the table-I, the CNN (Convolution Neural Network) is considered as the best approach for multiple feature selection and for classification, CNN or any other ML techniques can be used.

B. Unsupervised Learning:
Unsupervised Learning uses the unlabeled data where the training set is not provided. So the common diagnostic features are considered among them by organizing the data, searching and changing supported on internal knowledge. Algorithm tries to place related things in a cluster and non related in a different cluster. The idea of similarity is inferred from the similarity measure. Some of the approaches for diagnosis of DR are tabulated as shown in Table-II.
The table I and II present different machine learning approaches. The open challenges to be addressed are:
- Multi-Feature Selection namely Microaneurysms, Retinal, Hemorrhages, Retinal oedema, Cotton woolen Spots, Arduous exudates, Venous Retardation, Intraretinal micro vascular abnormalities is a challenging task.
- The different classification techniques PDR, NPDR and Diabetic Maculopathy.
- The Performance and time efficient method selection for the feature extraction and classification.
Table- I: Supervised learning Techniques to diagnose DR

<table>
<thead>
<tr>
<th>Authors/Reference</th>
<th>Approach</th>
<th>Data Set</th>
<th>Classification</th>
<th>Classification Parameters</th>
<th>Performance Measures</th>
<th>Merits</th>
<th>Demerits</th>
</tr>
</thead>
<tbody>
<tr>
<td>Handayani Tjandrastra [3], 2013</td>
<td>Soft Margin SVM</td>
<td>Messidor database</td>
<td>1. Moderate NPDR 2. Severe NPDR</td>
<td>Hard Exudates</td>
<td>- - 90.54%</td>
<td>Improved speed with high accuracy</td>
<td>NPDR is constructed only based on exudates features.</td>
</tr>
<tr>
<td>ShuangYu [5], 2017</td>
<td>CNN</td>
<td>E-Ophtha EX</td>
<td>1. DR 2. Non-DR</td>
<td>Pixel level Exudates detection</td>
<td>96% 88.85% 91.92%</td>
<td>Fair generalization capability even with varying the contrast and image quality.</td>
<td>Considers only exudates feature.</td>
</tr>
<tr>
<td>Devvi Sarwinda [7], 2018</td>
<td>SVM and Random Forest learning</td>
<td>DiaretDB0</td>
<td>1. Normal 2. Mild-NPDR 3. Moderate-NPDR 4. Severe-NPDR</td>
<td>Texture analysis</td>
<td>- 91% 85%</td>
<td>Features are extracted from HOG method where no explicit training is required</td>
<td>Constrained only for NPDR</td>
</tr>
<tr>
<td>Maroof Ahmad [8], 2019</td>
<td>DEEP LEARNNG</td>
<td>Messidor-2</td>
<td>1. NPDR 2. Normal 3. (PDR)</td>
<td>1. Micro aneurysms 2. Hard exudates 3. Hemorrhages 4. Neo-Vascularisation</td>
<td>- - 0.90-0.95 (Granular level) 0.20 - 0.45 (Average Localization)</td>
<td>1. Eliminates the need for feature selection explicitly designed 2. Classification performance is high</td>
<td>Average Localization score is less</td>
</tr>
<tr>
<td>Dinial [9], 2019</td>
<td>CNN and SVM</td>
<td>Messidor database - base12 &amp; base13</td>
<td>1. Normal NPDR 2. Severe NPDR</td>
<td>- - 95.83% (base 12) 95.24% (base 13)</td>
<td>Required time for computation is reduced by fine-tuning with CNN</td>
<td>Results with low efficiency when applied to large datasets</td>
<td></td>
</tr>
</tbody>
</table>

* Results reported in the database Messidor-2

Table- II: Unsupervised learning Techniques to diagnose DR

<table>
<thead>
<tr>
<th>Authors/Reference</th>
<th>Approach</th>
<th>Data Set</th>
<th>Classification</th>
<th>Classification Parameters</th>
<th>Performance Measures</th>
<th>Merits</th>
<th>Demerits</th>
</tr>
</thead>
<tbody>
<tr>
<td>Akara Sopharak [1], 2009</td>
<td>Fuzzy C- means Clustering</td>
<td>non-mydrastic retinal camera images</td>
<td>1. DR 2. Non-DR</td>
<td>Exudates</td>
<td>99.24 87.28 -</td>
<td>Improved execution speed</td>
<td>Considers only exudates feature.</td>
</tr>
<tr>
<td>Carla Agurto [2], 2010</td>
<td>Multiscale Amplitude-modulation frequency-modulation (AM-FM) &amp; K-Means Cluster</td>
<td>Messidor &amp; ETDRS database</td>
<td>1. Normal and 2. Pathological retinal images</td>
<td>Characterizatio n of retinal structures</td>
<td>0.71 0.97 -</td>
<td>It considers only global classification of images without having explicitly trained database which infers amnotation of each lesion.</td>
<td>Does not consider different classification of DR</td>
</tr>
<tr>
<td>Anupama. [6], 2017</td>
<td>Hessian Matrix &amp; Kmeans clustering</td>
<td>Retinal images</td>
<td>1. DR 2. Non-DR</td>
<td>Blood Vessel Segmentation</td>
<td>98% 84% 96%</td>
<td>Improved performance</td>
<td>Minimal feature consideration</td>
</tr>
</tbody>
</table>
IV. RESULTS AND DISCUSSIONS

The proposed model given in figure 3 considers the retina as the base for classification of DR and prediction of other disorders like stroke, chronic heart disease, renal dysfunction, cardiovascular mortality. current standards for cardiovascular disease mortality screening requires numerous multivariate derived from blood samples, age, gender, smoking status, hypertension, body mass index, glucose, and cholesterol levels [10]. Most cardiovascular risk calculators use some aggregation of these variants to determine patients at the risk of experiencing either a major cardiovascular event or cardiac-related mortality within a pre-determined period of time.

It can be hypothesized that hemorrhages, hard exudates, and micro aneurysms combined with hypertension indicate hefty association with heart failure, stroke, renal non-function and cardiovascular risk [12].

Fig. 3. Proposed Model for classification of DR

V. CONCLUSION

The proposed model considers features to be extracted from fundus image for the diagnosis of DR with different classification. So feature selection and extraction must be carefully defined and when combined with hypertension give the association with many other disorders namely Stroke, Congestive Heart Failure and Cardiovascular Risk. It eliminates the need of many preliminary tests and provides complete predictions of disorders to be referable or not.

The future scope includes the implementation of the model using CNN for feature selection as it considers many parameters and the suitable classifier for offline must be analyzed for time and performance efficiency.

REFERENCES


AUTHORS PROFILE

Manjula L is a Research Scholar at RNSIT. She earned Mtech and BE in Computer Science and Engineering from Visvesvaraya Technological University. Her research interests are Machine Learning, Artificial Intelligence, Internet of Things and Ophthalmology.

Dr G T Raju currently working as Vice-Principal and Head of CSE Dept. RNSIT. He has 25+ years of teaching and 12+ years of research experience. He has published 55 papers in international Journals, conferences. He has organized 28 workshops/FDP/conferences & delivered 30 invited talks. He has received 21 lakhs research grants from AICTE and VTU. He is life member of ISTE and CSI. His research interests are pattern recognitions, web mining, image processing & information retrieval, machine learning and artificial intelligence. He has held various positions at university.