

## 1. INTRODUCTION.

Diabetic retinopathy (die-Uh-BET-ik ret-NOP-uh-thee) is a diabetes complication affects eyes. It's caused by damage to the blood vessels of the light-sensitive tissue at the back of the eye (retina). At first, diabetic retinopathy might cause no symptoms or only mild vision problems.

## 2. LITERATURE SURVEY

Diabetes is a globally prevalent disease that can cause visible microvascular complications such as diabetic retinopathy and macular in the human eye retina, the images of which are today used for manual disease screening and diagnosis. This intensive task could greatly benefit from automatic detection using deep learning technique. Here we present a deep learning system that identifies referable diabetic retinopathy

comparably or better than presented in the previous studies, although we use only a small fraction of images ( $<1/4$ ) in training but are aided with higher image resolutions. We also provide novel results for five different screening and clinical

grading systems for diabetic retinopathy and macular

classification including state-of-the-art results for accurately classifying images according to clinical five-grade diabetic retinopathy and for the first time for the four-grade diabetic macular scales. These results suggest that a deep learning system could increase the cost-effectiveness of screening and diagnosis, while attaining higher than recommended performance, and that the system could be applied in clinical

examinations requiring finer grading.

### Introduction

Diabetic retinopathy is the most common microvascular complication in diabetes, for the screening of which the retinal imaging is the most widely used method due to its high sensitivity in detecting retinopathy. The evaluation of the severity and degree of retinopathy associated with a person having diabetes is currently performed by medical experts based on the fundus or retinal images of the patient's eyes. As the number of patients with diabetes is rapidly increasing, the number of retinal images produced by the screening programmes will also increase, which in turn introduces a large intensive burden on the medical experts as well

as cost to the healthcare services. This could be alleviated with an automated system either as support for medical experts' work or as full diagnosis tool. There are two recent studies that have investigated the use of deep learning systems in automated detection of

diabetic retinopathy. Both show that an automated system, based on the deep learning artificial neural

network approach, can achieve high sensitivity with high specificity in detecting the referable diabetic retinopathy, defined as moderate or worse diabetic retinopathy. There are also other

referable eye complications that have recently been investigated with this approach, such as diabetic macular and possible glaucoma and age-related macular degeneration. Methods

#### Original fundus image dataset

The research of present study was done in collaboration with Digifundus Ltd, an ISO 9001:2015 certified provider of diabetic retinopathy screening and monitoring services in Finland. Digifundus Ltd provided an open, anonymized retinal image dataset of patients with diabetes, including 41122 graded retinal colour images from 14624 patients. The images were taken with Canon CR2 retinal camera after inducing with tropicamide 5 mg/ml eye drops. Two 45 degree colour fundus photographs, centered on fovea and optic disc were taken from the patient's both eyes. The output images were of variable resolutions, ranging from  $3888 \times 2592$  to  $5184 \times 3456$  pixels. The present study is a methodological study with anonymized medical data and without any intervention in the integrity of a person such as contact with a person. In Finnish law this is not considered as a medical study requiring approval by an ethics committee or a written consent of a person. Retinal image grading systems and gradability Each of the retinal images had been graded with respect to three different criteria, (i) diabetic retinopathy, (ii) macular edema, and (iii) gradability. Images are graded with the proposed international clinical diabetic retinopathy and macular edema disease severity scales denoted later as PIRC and PIMEC, respectively. The image gradability is a two-stage system, which considers an image to be either gradable or not. All

personnel participating in retinopathy assessment had

over 10 years' experience in diabetic retinopathy grading. Retinal images with no lesions or mild diabetic lesions were graded by an optometrist and an M.D. trained for retinopathy grading. All images with moderate or worse changes were graded by two ophthalmologists both with more than 10 years of experience in grading fundus images. If there was a disagreement in grading, such an image was not included in this study. PIRC and PIMEC grades were further used to obtain additional three types of grading systems: (i) a binary system of nonreferable / referable diabetic

retinopathy (NRDR/RDR), (ii) a binary system of nonreferable / referable diabetic macular

edema (NRDME/RDME), and (iii) three-class system of ungradable/NRDR/RDR. The NRDR/RDR system considers the cases with no diabetic retinopathy and mild diabetic retinopathy as nonreferable diabetic retinopathy, and the cases with moderate or worse diabetic retinopathy as referable diabetic retinopathy. This system has been used in recent works investigating automated detection of diabetic retinopathy. The NRDME/RDME system here is defined such that the absence of macular edema is defined as nonreferable diabetic macular edema and any level of macular edema as referable diabetic macular edema. Note that only the gradable images were graded for diabetic retinopathy and macular edema. Ungradable images were included in a single task, in combination with referable diabetic retinopathy classification, which constitutes the grading system QRDR, in which each image is considered to be either ungradable, depicting nonreferable diabetic retinopathy, or depicting referable diabetic retinopathy

(ungradable/NRDR/RDR). Image preprocessing and dataset division In the model training and subsequent primary validation, we used preprocessed versions of the original images. The preprocessing consisted of image cropping followed by resizing. Each image was cropped to a square

shape which included the most tightly contained circular area of fundus. The procedure removed most of the black borders and all of the patient related annotations from the image data. Each of the cropped images were then resized to five different standard input image sizes of  $256 \times 256$ ,  $299 \times 299$ ,  $512 \times 512$ ,  $1024 \times 1024$ , and  $2095 \times 2095$  pixels. The largest image size was the smallest native resolution of the retinal

cameras after the preprocessing steps. Here the creation of multiple resolutions was done for the purposes of analyzing the effect of the input image resolution on the classification performance. The obtained processed datasets were divided into three sets: training, tuning, and primary validation set in the 70%, 10% and 20% proportions of the whole image dataset, respectively, separately for each of the grading systems used in the experiments. In the division per a particular grading system, the different sets were to have similar grade distributions, and that the dataset data per patient to not reside in multiple but only in one of the three sets (of training, tuning, and primary validation), in order to prevent the possibility of

obtaining over-optimistic results due to data memorization. Table 1 shows the statistics of the resulting divisions that were used in the experiments. Note that the grade distributions across the different sets were similar, with respect to each grading system, for example, when we consider the NRDR/RDR-system, the proportion of images associated with referable diabetic retinopathy in the training, tuning and primary validation set 44%, 43.9% and 43.4%, respectively. Results

In the binary classification tasks, i.e. NRDR/RDR and NRDM/ RDME, our algorithm achieved the best results using the largest  $2095 \times 2095$  pixels input image size. In the NRDR/RDR classification on our primary validation set having 7118 images, our algorithm achieved the sensitivity of 0.896 (with 95% CI: 0.885–0.907) and specificity 0.974 (with 95% CI: 0.969–0.979) and AUC of 0.987 (with 95% CI: 0.984–0.989). Our model

performance was evaluated at the operating point where the tuning set achieved 0.900 sensitivity, in a similar manner to Ting et al. while Gulshan et al. had two operating points namely at a high specificity (0.980) point and at a high sensitivity (0.975) point. In Table we present the AUC values of our model, along with the AUC values reported by Gulshan et al. and Ting et al. The two other recent studies, Krause et al. and Guan et al.

also explored the NRDR/RDR classification, but as they do not report results close to the 0.900 sensitivity point, References

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3.IDEATION & PROPOSED SOLUTION:

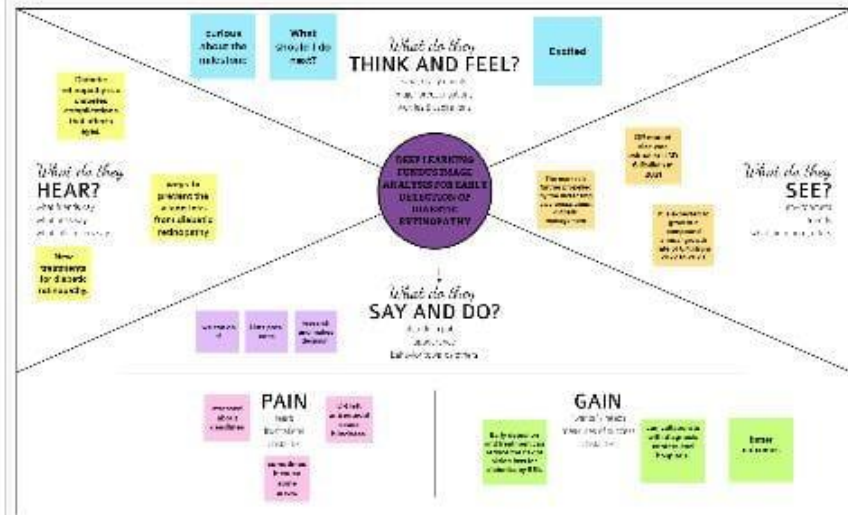
3.1 EMPATHY MAP CANVAS

# Empathy Map Canvas

Gain insight and understanding on solving customer problems.

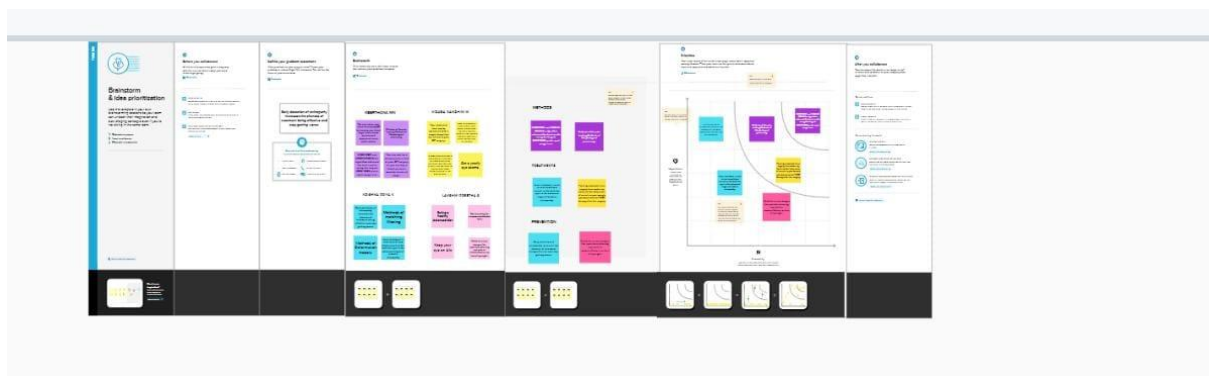
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Build empathy and keep your focus on the user by putting yourself in their shoes.



Share your feedback

## 3.2 IDEATION & BRAINSTORMING



### 3.3 PROPOSED SOLUTION:

#### Project Design Phase-I Proposed Solution

Date	19 September 2022
Team ID	PNT2022TMD33204
Project Name	DEEP LEARNING FUNDUS IMAGE ANALYSIS FOR EARLY DETECTION OF DIABETIC RETINOPATHY.
Maximum Marks	2 Marks

#### Proposed Solution Template:

Project team shall fill the following information in proposed solution template.

S.No	Parameter	Description
•	Problem Statement (Problem to be solved)	<ul style="list-style-type: none"><li>Diabetic retinopathy is a leading cause of vision loss globally. Early detection of retinopathy increases the chances of treatment being effective and stop getting worse.</li></ul>
•	Idea / Solution description	<ul style="list-style-type: none"><li>You can reduce your of developing DR by keeping your blood sugar levels, blood pressure and cholesterol levels under control and Pay attention to the vision changes.</li></ul>
•	Novelty / Uniqueness	<ul style="list-style-type: none"><li>IDX-DR is an AI diagnostic system that autonomously diagnosis patients for diabetic retinopathy.</li><li>No need for specialist overread or telemedicine call backs.</li><li>A Simple user interface.</li><li>customised workflow integration solution.</li></ul>
•	Social Impact / Customer Satisfaction	<ul style="list-style-type: none"><li>Helps in preventing the loss of visibility to the needed through CSR activities or through healthcare camps.</li></ul>
•	Business Model (Revenue Model)	<ul style="list-style-type: none"><li>Can collaborate with diagnosis centers and hospitals and government for health awareness camps.</li></ul>
•	Scalability of the Solution	<ul style="list-style-type: none"><li>Agreement was high, and exams containing more than minimal DR were detected. IDX-DR analyzes images for signs of diabetic retinopathy is accurate and providing results in 30seconds.</li></ul>

### 3.4 PROBLEM SOLUTION FIT:

Project Title: DEEP LEARNING FUNDUS IMAGE ANALYSIS FOR EARLY DETECTION OF DIABETIC RETINOPATHY.		Project Design Phase-I - Solution Fit Template		TEAM ID:PNT2022TMID33204	
Define CS, JT into CC	<b>1. CUSTOMER SEGMENT(S)</b> A 50years old man who has been diabetic for 10 years and thought him diabetes was under control but noticed some irregularities in his vision. He wants to find is he has a diabetic retinopathy (DR).	<b>6. CUSTOMER CONSTRAINTS</b> <ul style="list-style-type: none"> <li>Limited range of vision</li> <li>Get a yearly eye exam</li> <li>Control your blood sugar level</li> <li>Eat a balanced diet lower in fat and sugar.</li> </ul>	<b>5. AVAILABLE SOLUTIONS</b> Regular eye exams, good control of blood sugar and blood pressure, and early intervention for vision problems can help prevent severe vision loss.	Explore AS, differentiate	
	<b>2. JOBS-TO-BE-DONE / PROBLEMS</b> Diabetic retinopathy involves the growth of abnormal blood vessels in the retina. Complications can lead to serious vision problems: <ul style="list-style-type: none"> <li>Blindness or vision loss</li> <li>Retinal detachment</li> <li>Glaucoma</li> </ul>	<b>9. PROBLEM ROOT CAUSE</b> The real cause of this problem is poor control over diabetes, irregular blood sugar levels, Blood pressure and cholesterol.	<b>7. BEHAVIOUR</b> He seeks advices through the personal friends and family. He looks for any lifestyle changes that can be improve his condition.		Focus on AS, JT into BE
Identify CS, JT into BE	<b>3. TRIGGERS</b> Seeing other healthcare institutions produce better results when early detection of diabetic retinopathy saves more patient's vision. By seeing how many patient's vision can be saved by early detection of diabetic retinopathy.	<b>10. YOUR SOLUTION</b> In case of diabetes, reduce your risk of getting diabetic retinopathy by doing the following <ul style="list-style-type: none"> <li>Manage your diabetes</li> <li>Make a healthy eating and physical activity part of your daily routine.</li> <li>Smoking increases your risk of various diabetes complications, including DR.</li> <li>Contact your eye doctor right away if your vision suddenly changes or becomes blurry, spotty or hazy.</li> </ul>	<b>8. CHANNELS of BEHAVIOUR</b> <b>ONLINE</b> <ul style="list-style-type: none"> <li>Finding the cause of the problem</li> <li>Reading reviews of people having blurry vision</li> </ul> <b>OFFLINE</b> <ul style="list-style-type: none"> <li>Consulting a doctor taking prescribed medicine.</li> <li>Regular checkup for eyes.</li> </ul>	Identify CS, JT into BE	
	<b>4. EMOTIONS: BEFORE / AFTER</b> <b>BEFORE</b> <ul style="list-style-type: none"> <li>Poor vision</li> <li>Fear of permanent blindness</li> <li>Dark or empty areas in vision</li> <li>Blurred vision</li> </ul> <b>AFTER</b> <ul style="list-style-type: none"> <li>If treated early, vision will be corrected and regained.</li> <li>The possibility of getting blind is low.</li> </ul>				

## 4 REQUIRMENT ANALYSIS

### 4.1 FUNCTIONAL REQUIREMENT &

### 4.2 NON-FUNCTIONAL REQUIREMENTS

#### Project Design Phase-II Solution Requirements (Functional & Non-functional)

Date	05 October 2022
Team ID	PNT2022TMD33254
Project Name	Deep learning fundus image analysis for early detection of Diabetic retinopathy.
Maximum Marks	4 Marks

##### Functional Requirements:

Following are the functional requirements of the proposed solution:

FR No.	Functional Requirement [Epic]	Sub Requirement (Story / Sub-Task)
FR-1	Identifying and selecting dataset	The appropriate dataset to enhance the model's performance is necessary to select.
FR-2	Invitation and Information	Invite the full cohort for screening, supplying information tailored appropriately for different groups to enable informed choice to participate.
FR-3	Training	It is required to import the libraries needed for training the model.
FR-4	Diagnosis	Diagnose true cases and identify the false solution.
FR-5	Testing	Conduct screening tests with different data to test if the model is trained well to predict the medical condition.
FR-6	Reporting and treatment	Report the outcomes to identify false negatives and improve effectiveness of the screening program. The testing of the model helps us to identify the appropriate treatment.

##### Non-functional Requirements:

Following are the non-functional requirements of the proposed solution:

FR No.	Non-Functional Requirement	Description
NFR-1	Usability	Users with basic understanding of the medical condition and computer knowledge can operate the system. User friendly interface that can be accessed with ease by users.
NFR-2	Security	Deep learning AI can be more precise around sensitive organs and tissues, reduce blood loss, risk of infection and pain during detection.
NFR-3	Reliability	There is a chance of hardware failure or false positives when the testing data is more different.

NFR-4	Performance	than the training dataset. Permission granted only by the administrator of the system. If the system update fails or bugs in the code even though the system can roll back to its initial state. The performance of the model is meant to give speed results for the patients.
NFR-5	Availability	The treatment should be available at low cost so that everyone with DR can find it beneficial.
NFR-6	Scalability	By processing more datasets for the reference of DR detection.



## 5.PROJECT DESIGN

### 5.1 DATA FLOW DIAGRAMS &

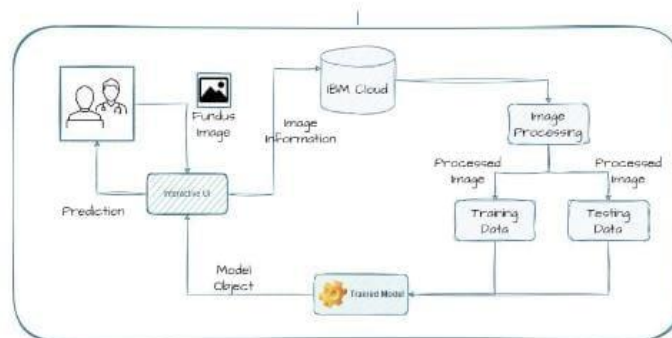
### 5.2 USER STORIES

#### Project Design Phase-II Data Flow Diagram & User Stories

Date	03October 2022
Team ID	PNT2022TMID33204
Project Name	Deep Learning Fundus Image Analysis for Early Detection of Diabetic Retinopathy
Maximum Marks	4 Marks

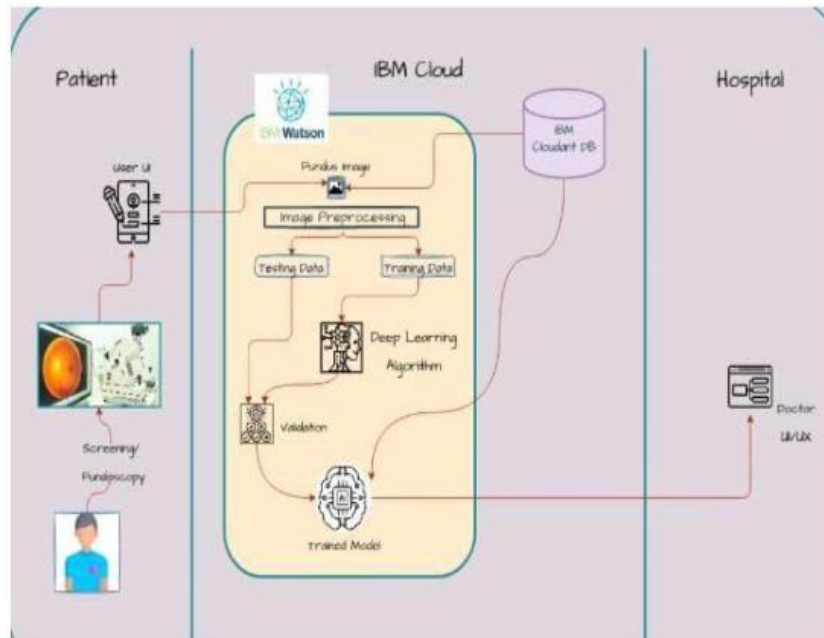
##### Data Flow Diagrams:

A Data Flow Diagram (DFD) is a traditional visual representation of the information flows within a system. A neat and clear DFD can depict the right amount of the system requirement graphically. It shows how data enters and leaves the system, what changes the information, and where data is stored.



### 5.3 SOLUTION & TECHNICAL ARCHITECTURE:

#### TECHNOLOGY ARCHITECTURE:

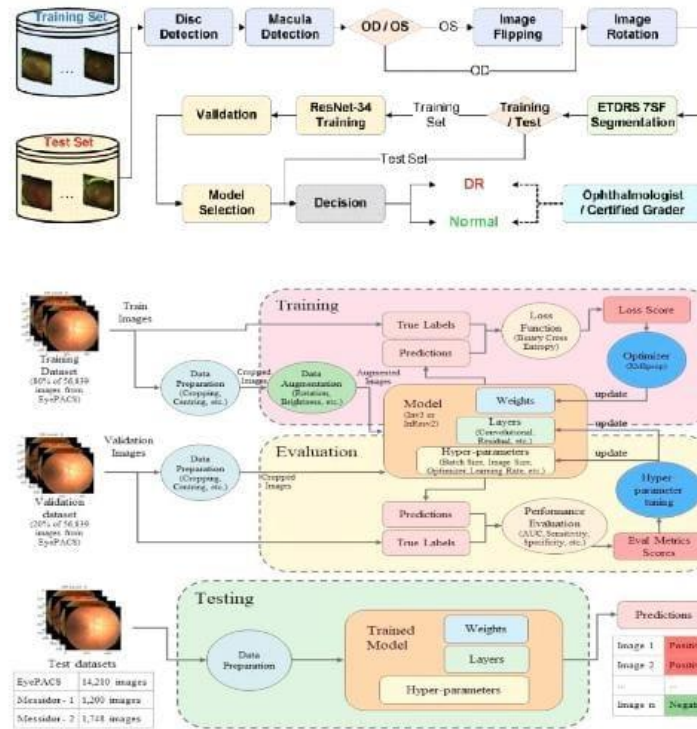


## PROJECT DESIGN PHASE –1

### SOLUTION ARCHITECTURE

DATE	17 OCTOBER 22
TEAM ID	PNT2022TMID33204
PROJECT NAME	DEEP LEARNING FUNDUS IMAGE ANALYSIS FOR EARLY DETECTION OF DIABETIC RETINOPATHY.
MAXIMUM MARKS	4 MARKS

#### SOLUTION ARCHITECTURE:



## 6. PROJECT PLANNING & SCHEDULING:

### 6.1 SPRINT PLANNING & ESTIMATION:

## 7. CODING & SOLUTIONING:

```
from cloudant.client import cloudant
client = cloudant.iam("d0963b0d-6a01-4b4d-ac1b-78ccef67e303-bluemix", "vRzNCfpRbo2Bf4UIdNaX8fP-vWRHGp5vr9at08fDs7")
my_database = client.create_database('diabetic-retinopathy')
```

## 8. ADVANTAGE

Early detection and timely treatment having been shown to prevent visual loss and blindness in patients with retinal complication of diabetes

## DISADVANTAGES

High capital setup costs

Need to provide regular training for grader

## 9.conclusion

Diabetic retinopathy is now recognized to be an inflammatory neurovascular complication of the systemic disease with neuronal injury preceding the current clinical microvascular the inflammatory tissue injury concurrent in other organs.