Diabetic Retinopathy Screening using Deep Learning for Multi-class Imbalanced Datasets

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Abstract. Screening and diagnosis of diabetic retinopathy disease is a well known problem in the biomedical domain. The use of medical imagery from a patient's eye for detecting the damage caused to blood vessels is a part of the computer-aided diagnosis that has immensely progressed over the past few years due to the advent and success of deep learning. The challenges related to imbalanced datasets, inconsistent annotations, less number of sample images and inappropriate performance evaluation metrics has caused an adverse impact on the performance of the deep learning models. In order to tackle the effect caused by class imbalance, we have done extensive comparative analysis between various state-of-the-art methods on three benchmark datasets of diabetic retinopathy: - Kaggle DR detection, IDRiD and DDR, for classification, object detection and segmentation tasks. This research could serve as a concrete baseline for future research in this field to find appropriate approaches and deep learning architectures for imbalanced datasets.

Keywords: Diabetic retinopathy, Deep learning, Image classification, Object detection, Segmentation, Transfer learning

1 Introduction

Diabetes is a well-known chronic disease which has been known to contribute to an alarming rise in the number of deaths worldwide. The adverse effects of diabetes are usually seen in the retina of eyes, which can lead to the loss of vision problems to a great extent at later stages of the disease. Diabetic Retinopathy (DR) is the condition where eyes are affected due to the occurrence of diabetes [1] [2]. This problematic condition occurs due to the presence of high levels of sugar in the blood vessels of the retina. Hence, detection of DR at early stages of the disease plays a crucial role in saving the life of the patient from this life-threatening situation. According to the collaborative research study conducted by early treatment Diabetic Retinopathy research group (ETDS) and International Clinical Diabetic Retinopathy, there are several DR levels which have been categorized, such as: stage 0 which apparently signifies no retinopathy, whereas stages 1, 2 and 3 indicate mild, moderate and severe non-proliferative diabetic

retinopathy conditions, respectively [3]. Stage 4 is the dangerous and advanced stage which leads to the proliferative diabetic retinopathy condition. The last stage has an overall serious impact on the health of the patient as it might lead to different problems related to vision, such as blurriness or blindness. Automatic computer-aided diagnosis screening of DR can speeden up the process in a much more effortless manner as compared to the manual time consuming screening task. The use of such automated methods would result in early and reliable detection of the disease, which can also be easily applied for mass screening of populations. With the recent developments in computing hardware, there has been a tremendous ease in applying deep learning techniques extensively, which has greatly benefited the biomedical domain. Thus, the application of deep learning has resulted in significant improvement in the diagnosis of DR in comparison to the traditional approaches.

The objective of this work is to conduct extensive experimental comparative analysis using computer vision techniques to find out the appropriate pre-trained neural network for performing classification, object detection and segmentation on diabetic retinopathy multi-class imbalanced datasets. An imbalanced dataset is one in which the population of the majority classes is much higher than the population of the minority classes, due to which the classification results are biased towards the majority classes [4]. It is observed that most real world datasets in the biomedical domain are imbalanced in nature, and it was found that datasets associated with diabetic retinopathy have a highly skewed class distribution. For our work, we specifically focus on multi-class imbalanced datasets where the samples are not equally distributed in the respective classes. We have emphasized on the challenging aspects of creation of deep learning models for multiclass imbalanced datasets, as the mystifying aspect is to identify the classes which will fall into majority or minority categories based on the samples in each class distribution [5][6]. Along with that, we also emphasize on the significance of the right evaluation metrics that are to be used for measuring the performance of different deep learning models for respective computer vision tasks. The multi-class distribution of samples is more challenging in comparison to the binary class problem (having two classes). However, in order to have an efficient diagnostic method, it is equally important to recognize both the under-represented classes (minority classes) and majority classes equally. There is a need to have effective models which can diagnose diabetes accurately, and also detect the type of diabetes, based on automated diagnosis using classification tasks. Further, it is also important to locate anomalies in the retinal images using object detection and segmentation techniques, in order to provide accurate treatment at an early stage to save the life of the patient suffering from diabetes and avoid the high risk of vision loss. It is important to automatically count and detect the presence of the lesions in the retina such as exudates, microaneurysms and hemorrhage, for appropriate diagnosis for diabetic retinopathy. The microaneurysms are red dots having 12-100 μm diameters, which is usually due to minute blood clots. Hard exudates are shiny white or cream spots which are present in the retina of the eve due to presence of an exudate fluid. Excess accumulation of this fluid might be lifethreatening if it appears near to the center of

the macula of the patient's eye. Hemorrhages occur due to the release of blood from broken blood vessels, and can be of different shapes and sizes. In this paper, we have shown extensive experiments using various pre-trained networks to demonstrate their efficacy for various computer vision tasks on DR images. To the best of our knowledge we have not found any paper related to combining the comparative analysis of various pre-trained networks for classification, object detection and segmentation of diabetic retinopathy datasets in a single paper, which also notably comprises of a curated set of the most popular problem statements in computer vision. [7] [8]. Classification is the supervised learning technique to classify unknown test samples into categories [9], while object detection is used to locate objects in an image by drawing the bounding box around the object region [10]. Whereas, image segmentation is the process of dividing an image into multiple connected regions based on some similarity criteria [11]. Computer vision problems can be broadly classified into dense, sparse and global labeling tasks. Classification is regarded as a global labeling task where a single image is assigned a single class label; object detection is considered a sparse labeling task since only a few pixels of the image are labeled as there are bounding boxes which contain regional information of sparse locations; Segmentation, on the other hand, is a dense labeling task as each and every pixel of the image has an assigned class label.

Nowadays, in the computer vision domain, most of the problems has increased the resolution capacity with the support of Convolutional Neural Networks (CNNs) that is quite populary used in various application areas. CNNs have proved to be revolutionary in the field of computer vision. CNNs can perform automatic feature extraction as well as classification in a unit, which outperforms traditional approaches [10]. They have been widely used in the current era for classification, segmentation and object detection tasks. In case of object detection, CNN is used by modeling the object detection as a regression problem, or in other words, it can be used as a backbone network for the object detection task. Training of CNNs can be broadly categorized into two categories: either training from scratch or reusing pretrained networks to train the model on a target dataset. The pre-trained models are the ones

which are already trained on a larger dataset comprising of millions of images [12]. There are several advantages of using pre-trained models over CNNs that are trained from scratch, since they are based upon the transfer learning concept. Another advantage of using pretrained networks is that there are less requirements of computational resources, and training time would also be significantly lesser [13]. So considering the various advantages of the pre-trained models, in this paper, we have conducted a comprehensive performance analysis of various state-of-the art deep learning pretrained models on three popular biomedical diabetes datasets of varied sizes: Kaggle DR detection or Kaggle DRD [14], IDRiD [15] and DDR [16] for image classification, object detection and segmentation tasks. It is notable to mention that the availability of pre-trained networks for segmentation and object detection tasks is comparatively lesser in comparison to those for classification tasks. The major contributions of our work are highlighted as follows: (1) We have done extensive comparative experimental analysis between various state-ofthe art pre-trained networks using three popular well known imbalanced diabetes datasets for image classification, using small, medium and large multi-class imbalanced datasets. (2) Comparison is also conducted between various state-of-the art methods for object detection and segmentation tasks using two available multi-class imbalanced datasets. (3) Emphasis is laid upon the relevance of using the transfer learning approaches instead of training the CNN from scratch, as training the network from scratch will involve massive computational power and resources for small, medium and larger diabetes datasets. The rest of the paper is organized as follows: literature review is described in section 2; in sections 3, 4 and 5, we have discussed the methodology, datasets and performance evaluations metrics, respectively, which is followed by the implementation details, results and discussion in sections 6 and 7. Finally, conclusion is presented in section 8.

2 Literature Review

We have conducted a literature survey compiled set of works regarding the usage of deep learning in diabetic retinopathy detection. There are readily available datasets related to diabetes, to which researchers have applied advancement in computational resources, deep learning has been extensively used in various applications. Healthcare is one of the domains where deep learning is prevalent nowadays. (2019) had emphasized on Raghu et al. the importance of transfer learning using CNNs for medical imaging datasets [17]. The experiments were conducted on RETINA and CHEXPERT datasets, and the Receiver Operating Characteristics (ROC) - Area Under Curve (AUC) performance metric was used for the evaluation. Various researchers have worked on different pre-trained networks for diabetic retinopathy datasets. Hagos and Shri Kant (2019) had highlighted the application of Inception-v3 pre-trained model on a smaller subset of diabetic retinopathy detection dataset by considering accuracy as the evaluation measure, and they achieved 90.9% accuracy [18]. For an imbalanced dataset, there are many other reliable evaluation metrics other than accuracy which should be considered, as well, while measuring the performance of the model, which are missing in many works. Thota and Reddy (2020)had proposed a variant of VGG16 pre-trained model for performing the classification task on EyePACS dataset [19] whose performance was evaluated using metrics such as sensitivity, specificity and AUC along with accuracy. Lam et al. (2018) had used GoogleNet and AlexNet pre-trained models on the Messidor-1 diabetic retinopathy dataset [20]. They took concatenated features from different pre-trained networks and passed them as input to machine learning classifiers. Kassani et al. (2019) had shown improved performance by considering Xception as the pre-trained network for feature extraction, and further processed the extracted features using a multi-layer perceptron neural network [21]. Wan et al. (2018) had obtained high performance on the publicly available Kaggle DRD dataset by using VggNet that outperformed other pre-trained networks such as AlexNet, GoogleNet and ResNet, and they had also emphasized upon the role of fine tuning and transfer learning [22]. Saini and Susan proposed deep learning based DCGAN approach for tackling class imbalance through synthetic data generation for Breast Cancer detection [23]. A modified VGG16 network architecture was proposed for the purpose, that

deep learning techniques for automatic detec-

tion of diabetic retinopathy. With the recent

outperformed various other networks including CNNs trained from scratch. Additionally, the authors proposed VGGIN-Net network which was based upon transfer learning and essentially constructed using a combination of VGG16 and Inception networks [24]. It was found that fine-tuning pre-trained models in the biomedical domain yield much better results than training models from scratch. There are numerous other works in literature related to object detection and segmentation using various diabetes datasets. Zhang et al. (2020) had proposed a convolutional neural network formulated by combining few convolutional and deconvolutional layers along with the Fully Connected (FC) and softmax layers to distinguish the presence of microaneurysms in retinal fundus images [25]. Oliveira et al. (2021) had proposed a variant of Faster R-CNN, with data augmentation, for Diabetic foot Ulcer Detection [26]. Porwal et al. (2018) reported results compiled by various researchers who had participated in the grand diabetic retinopathy segmentation and grading challenge on Indian retinopathy Image dataset (IDRiD) [27]. From the analysis, it was interpreted that the top performance approaches involved some form of data augmentation or ensemble models. Another major finding was that solving the class imbalance problem would lead to a tremendous improvement in the overall performance of the model. So dealing with class imbalance is an important aspect, along with the overfitting problem, in order to generally enhance the performance of the network. Data Augmentation is a regularization method [28] which helps to resolve the overfitting problem generally occurring in deep learning models. It is equally important to select the correct set of data augmentation operations to overall improve the performance of the model. However, selecting the correct set of data augmentation operations is a tedious and manual process and also it is difficult to design the correct pipeline of operations. So certain automated approaches are proposed in literature to select the correct set of data augmentation operations such as AutoAugment [29] and RandAugment [30]. AutoAugment is a computationally expensive process and it is formulated by the Proximal Policy Optimization (PPO) algorithm with a large search space of different augmentation operations and their magnitudes. However,

the RandAugment approach proposed by the same authors removed the PPO algorithm resulting in a much smaller search space. A simple grid search based tuning approach reduces the computational complexity as well as the search space, deeming it to be one of the most effective automated approaches for data augmentation of CNNs.

Hu et al. proposed a model based on neutroscopic enhancement using neural networks to perform segmentation and extract the region of colorectal polyps found in tissue scans while comparing their approach with various other state-of-the-art models for the given biomedical area and presented the results in their work [31]. Wu et al. presented a cloud-based electronic medical record system highlighting the importance of cloud technology, related systems and specialized hardware that apart from dealing with challenges in data handling also takes care of the security aspects including confidentiality of patient information [32]. Their work also describes the different scalability challenges that come with large volumes of biomedical datasets. Advancements in digital technology especially ones involving edge devices and specialized hardware for small scale systems have increased the resolution capacity of different multimedia applications. Yang had presented a novel string matchet al. ing algorithm with extremely low complexity suitable for lossless compression for image and video content streaming [33]. Similarly, advent of hardware accelerators in this field have greatly benefited the biomedical domain for deep learning based approaches. Dai et al. proposed a deep LSTM based model trained using EMG signals which helps to rehabilitate musculoskeletal framework of the body which is another area of research in the biomedical community [34]. Tang et al. proposed a novel segmentation algorithm based on retinal fundus images using a back propagation neural network with the help of morphological processing, histogram equalization and gaussian filtering [35]. Additionally, He et al. explored the application of artificial neural networks for detecting lesion regions in lung cancer images and comparing different approaches for diagnosis of cancer [36]. Hence, deep learning models can be generalized to be applied to various areas from the biomedical domain, and contemporarily there have been multiple advancements in the field. Some of the major areas where biomedical applications have benefited from deep learning based architectures include musculoskeletal rehabilitation, rectal, breast, cervical, retinal and lung cancer, etc. For our work, we focus exclusively on deep learning based architectures for classification, segmentation and object detection on diabetic retinopathy datasets.

3 Methodology

In this section, we explain our methodology for the extensive comparative analysis of various state-of-the-art pre-trained deep learning networks for image classification, segmentation and object detection on challenging diabetic retinopathy datasets of varied sample sizes. We have projected an automated diagnosis system for diabetic retinopathy that performs classification, object detection and segmentation on diabetic retinopathy fundus images, as depicted in Figure 1. The different modules of our framework are explained next.

3.1 Classification

Classification is a supervised learning approach where the class label is known in advance. The model is trained using the training dataset having multiple images so that if any unknown sample images from the test dataset are passed to the trained model, it can predict the category or class to which the test sample belongs. The predicted output is matched with the target output in order to check how well the model has learned. With the recent advancements in high performance computing devices, deep learning is widely used in image classification tasks nowadays [37]. Here we have performed image classification on three varied size diabetic retinopathy image datasets to detect the degree of severity of diabetic retinopathy for the retina image of a given patient. Various pre-trained deep learning CNN architectures have been applied on three diabetic retinopathy datasets: - Kaggle DRD, IDRiD and DDR, in order to perform classification: VGG16, VGG19, ResNet50, ResNet101, ResNet152, Inception v3, ResNet50 v2, ResNet101 v2, ResNet152 v2, Xception, InceptionResNet v2, MobileNet v2, DenseNet169, DenseNet201 and Efficient-NetB0.

3.2 Object Detection

Object detection is used to determine the presence of an object in an image by drawing the bounding box around the object [38] [39]. Presence of multiple objects in an image determines whether it comes under single or multiclass object detection category. Object detection and localization is the procedure of detecting the presence of objects in the image with the coordinates of the location where the object is located in the image. Pathak et al. (2018) had emphasized the role of the wide usage of CNN-based deep learning in object detection tasks in various fields such as robotics, surveillance and transportation, autonomous driving, [40] [41] [42] [43].medical domain etc. In our current work, we have applied EfficientDet-D0, ResNet50 based Faster RCNN, SSD using MobileNet v1 and MobileNet v2, as well as ResNet50 based RetinaNet pre-trained networks on two popular datasets: IDRiD Dataset for Fovea and Optical Disc Detection along with DDR Dataset for lesion detection.

3.3 Segmentation

In image segmentation, the input image is divided into distinguished regions or segments. Each region has pixels having similar characteristics. In other words, segmentation is the process used to find boundaries to locate objects. Labels are assigned to each pixel in an image such that the pixels having the same characteristics are assigned the same label. In the segmentation process, the foreground is separated from the background. To perform segmentation we can use multiple pre-trained networks. In case of CNN architectures, we have encoder and decoder inside the model, both combined together. Image segmentation reduces the complexity and further makes the analysis of the image much simpler. We have applied DeepLab v2, DeepLab v3, PSPNet with cross-entropy loss and focal loss, respectively, on two popular datasets: DDR dataset for Lesion Detection and IDRiD dataset for both lesion as well as organ (fovea and optic disc) segmentation.

4 Datasets

While conducting the experiments, we have considered three well-known multi-class imbalanced diabetic datasets of varying sample sizes



Figure 1: Automated system for diabetic retinopathy screening using classification, object detection and segmentation of fundus images.

for classification, segmentation and object detection, that can be categorized based on their sizes into small, medium and large dataset categories as described below.

4.1 Kaggle DRD Dataset

It is a very large publicly available dataset that was originally published in a Kaggle competition. It consists of high-resolution images captured in varied imaging conditions, belonging to four classes: class 0 (no DR), class 1 (mild DR), class 2 (moderate DR), class 3 (severe DR) and class 4 (proliferative DR). Figure 2 depicts the illustration of a few random samples taken from different diabetic retinopathy grades from the Kaggle DRD dataset. In total 53,576 and 35,126 images are present in the test and train datasets, respectively [14]. The Kaggle DRD dataset is imbalanced in nature, and the test dataset has reasonably high number of image samples existing in each class. Our experiments use the same train-test split as that of the original competition.: Class 0 has 31,403 samples, class 1 consists of 3042 samples, class 2 has 6282, and further classes 3 and 4 have 977 and 966 samples, respectively. It was observed that the minority class comprises 2.016% of the entire dataset. For pre-processing, the images were resized from their very high initial resolution so that the radius of an eyeball is 300 pixels, then they were cropped to 90% of the radius, and encoded into 72 JPEG quality. [44]

4.2 DDR Dataset

We have also used the DDR dataset proposed by Li et al [16] for the experimentation. The dataset consists of three categories of annotations: - bounding box, pixel and DR grading level annotations. There are a very large number of sample images (13,673 fundus images) presented in the dataset, which were collected from 9,598 patients consisting of 6

Dataset	Segmentation	Object Detection	Classification
Kaggle DRD	-	-	 Diabetic Retinopathy Grading Class 0 - No DR Class 1 - Mild DR Class 2 - Moderate DR Class 3 - Severe DR Class 4 - Proliferative DR
DDR	 Lesion Segmentation Hemorrhages (HA) Microaneurysms (MA) Soft Exudates (SE) Hard Exudates (EX) 	 Detection of Lesion Hemorrhages (HA) Microaneurysms (MA) Soft Exudates (SE) Hard Exudates (EX) 	 Diabetic Retinopathy Grading Class 0 - No DR Class 1 - Mild DR Class 2 - Moderate DR Class 3 - Severe DR Class 4 - Proliferative DR Class 5 - Ungradable
IDRiD	 Lesion Segmentation Hemorrhages (HA) Microaneurysms (MA) Soft Exudates (SE) Hard Exudates (EX) Organ Segmentation Optical Disc (OD) 	Detection of Organ Centroids • Optical Disc • Fovea Centralis	 Diabetic Retinopathy Grading Class 0 - No DR Class 1 - Mild DR Class 2 - Moderate DR Class 3 - Severe DR Class 4 - Proliferative DR

Table 1: Distribution of classes across different tasks (Segmentation, Object Detection and Classification) for the three diabetic retinopathy datasets.

classes. Out of 13,673 fundus images, 6835 and 2733 images, respectively, are kept for training and validation purposes, and the remaining 4105 images are taken out for testing respectively. The images were originally collecte from a closer clinical perspective from 147 hospitals, covering 23 provinces in China. In our experiments we use the same split as that of the original authors. Figure 3 depicts a few random samples taken from different diabetic retinopathy grades from the DDR dataset. We found that in this dataset, the most underrepresented class is around 1.726% of the total samples. In the case of segmentation, there were a few samples (very minimal, around 5) with corrupted masks in the dataset that had to be removed before training.

4.3 Indian Diabetic Retinopathy image (IDRiD) Dataset

IDRiD is the first available diabetic retinopathy images database based on the Indian population and it consists of three parts: segmentation, disease grading (classification) and localization (object detection). The dataset consists of 4288×2848 size jpg file images [17]. During collection of the images, pupils of all subjects were dilated with one drop of tropicamide at 0.5% concentration such that they were captured with 50 degrees field of view. In both the cases of disease grading and localization, a total of 516 images are present in the dataset, out of which 413 images are considered for training and the remaining 103 images are kept for testing. Figure 4 illustrates a few random samples taken from different diabetic retinopathy grades from the IDRiD dataset. The ground truth labels available for localization are for Optic Disc Center location and Fovea Center location. The number of samples in the minority class is around 4.986% of the complete dataset, rendering it to be a case of severe class imbalance. In case of localization task for this dataset, the centroids of each of the objects to be localized i.e either lesion or fovea or optic disc, and we used the centroid information to generate bounding boxes of fixed sizes using a preset radius.

Table 1 reflects the overall distribution of samples in various classes for all the datasets, shown for different tasks such as segmentation, object detection and classification. Kaggle DRD dataset has classification task only, but IDRiD and DDR dataset have segmentation and object detection tasks available besides classification. Table 1 gives a clear depiction of the multiple classes associated with different computer vision tasks. Further, Figure 5 illustrates the histogram of the number of image samples present in each class of the three diabetes datasets, while Figure 6 depicts the histogram of the number of pixels belonging to each segmentation class for the respective diabetes datasets. As concluded from the uneven class population profiles in the two figures, there is a high degree of class imbalance associated with all three datasets for both the classification and segmentation task. In our work, we analyze the performance of the state-of-the-art deep neural networks for the DR screening task in the current scenario of high class imbalance.

5 Performance Evaluation Measures

In this paper, we have done comparative analysis between different state-of-the-art pretrained networks by using various performance evaluation metrics. For the classification task, various performance evaluation parameters are considered such as Cohen's Kappa (unweighted, linear, quadratic weighted), Accuracy, ROC-AUC (weighted and macro average), F1-score, Index Balanced Accuracy (IBA) and Geometric Mean (GMean) [45] [23] [46] [47] [48]. In the case of object detection, we have considered the following evaluation parameters for measuring the performance of the object detection models: Mean Average Precision (mAP), mAP @ 0.5IoU, mAP @ 0.75IoU, mAP (small, medium, large) and Average Recall (AR): AR @ 1,10,100 [49] [50]. Further, Intersection over Union (IoU) and Dice Score evaluation metrics are used for segmentation [51] [52].

6 Implementation Details

For conducting all our experiments, the TensorFlow v2.3 framework was used [53] with Python 3.8. For the classification and segmentation tasks, the CNNs were trained with the help of Keras models, while for object detection the TensorFlow Object Detection API was used. The experiments were accelerated using Google Cloud TPU hardware, access to which was available through the TensorFlow Research Cloud (TRC) program. In all the experiments, TPU v3-8 cores were used for training the deep learning models. For our experimental setup, the batch size of 512 is considered for all the datasets. The total training steps taken was 12,000 (400 epochs x 30 steps per epoch) for DDR [16] and for Kaggle DRD Dataset [14], while for IDRiD dataset it was set to 1000 (200 epochs x 5 steps per epoch) [15].

Various data augmentation operations were applied on all the images present in the train datasets while conducting the classification task, such as horizontal shifts, vertical shifts, rotation, flip etc., to reduce the over-fitting problem [46]. In all the classification datasets, the original class distributions of samples were imbalanced in nature. The target distribution was achieved such that each class has an equal number of samples represented during the training of the pipeline in all the batches. This was achieved using rejection resampling technique which is also popularly known as random undersampling. Hence undersampling technique was applied on-the-fly during the training process at the time of selection of each mini-batch, and it was also ensured that the samples are shuffled and randomly picked from the original distribution. The training process is made to last for sufficiently large numbers of epochs such that all training samples are covered irrespective of the fact that sample rejection has been applied at batch level. It was observed that without the application of this rejection resampling strategy, each classifier would face the adverse effects of class imbalance. However, data augmentation techniques were applied in all the experiments corresponding to the classification and object detection task. Data augmentation is a regularization technique that helps to enhance the overall performance of deep learning models [54]. Some data augmentation learning policies had also automated the data augmentation process. We have used the RandAugment technique while training the network [30]. There are several advantages of using RandAugment over other augmentation techniques as it will lead to the reduction of the search space or removal of separate search space because more computational expensive resources were required while training which will help to achieve optimal performance. The RandAugment approach works well for numerous datasets as well as tasks such as object detection and classification. Due to all these unique properties it results in better performance in comparison to other augmentation methods. The various data augmentation operations that we have applied are random grid shuffle, rescale, horizontal and vertical flips in horizontal and vertical flips, shear in x and y direction, translate in x and y direction, rotate, posterize, contrast, sharpness, and cutout. Cutout [55] is like dropout at



Figure 2: Illustration depicting a few random samples taken from different diabetic retinopathy grades for kaggle DRD dataset.

data level, which creates random black square patches in the images; it was found to be an essential augmentation operation.

After many iterations through our experimental setup, we were able to use the following hyperparameters for all of our classification experiments. The training process was intended at fine-tuning each of the classification networks using pre-trained weights in order to adapt the models for diabetic retinopathy tasks. A standard categorical cross-entropy loss and adaptive momentum optimization (Adam) variant of SGD was used while train-The choice of learning rate was set to ing. lower values in order to ensure that our models continually learn from retinal images without completely exterminating the knowledge on the large-scale ImageNet. Similar to fine tuning of CNNs in other works, we used an exponentially decaying learning rate with warmup in the range 1e-5 and 2e-4. Under this setting, the learning rate was initially ramped up linearly to 2e-4 for the first 160 epochs, sustained at that rate for another 80 epochs, and finally decayed exponentially for the remaining epochs using a decay of 0.8. Evaluation on validation

was performed each 10 epochs to ensure models can train well for longer. Each of the models were trained for a net budget of 400 sweeps (epochs) and the largely set value of number of warmup epochs is solely due to the random undersampling technique which significantly reduces samples from each epoch in order to obtain a balanced class distribution at train time. The warmup and sustain phase of the learning rate schedule is slightly longer to ensure that the complete dataset can be iterated across during the training phase by the models before the value of learning rate drops significantly. The batch size for the experiments were set to 512 and images were rescaled to $224 \ge 224$ which were found to be ideal for training on TPU v3-8 accelerators. TPU v3-8 hardware are ASIC chips designed by Google and operated on Google Cloud Platform and are specifically developed for fast machine learning and deep learning workloads. It can consume up to 128GB of on-chip high bandwidth memory (HBM) for up to 8 TPU cores which was helpful for accelerating our experiments, and the higher batch sizes significantly helped consume the large amount of available on-chip memory



Figure 3: Illustration depicting a few random samples taken from different diabetic retinopathy grades for DDR dataset.

for which learning rate was scaled accordingly [56]. For data augmentation along with the rejection resampling technique (random undersampling approach), as discussed, we slightly modified the RandAugment technique [57] to incorporate random flips, random grid shuffle and rescale transforms. A few set of randomly augmented samples generated using this technique are shown in Figure 7. After a series of manual tuning trials, RandAugment was applied with m=8 and n=2 and it was found that adding the extra image operations would greatly improve the classification performance for the diabetes datasets.

In case of lesion segmentation on the DDR and IDRiD datasets, we fine-tuned the segmentation models that are pre-trained on the PAS-CAL VOC 2012 dataset. The training process was carried out on TPUs using focal loss with gamma set to 2.0 and batch size of 64. The exponentially decayed learning rate schedule was similar to the classification experiments and in the range 1e-5 and 5e-4 with 80 warmup epochs. The focal loss models were able to significantly perform better than categorical cross-entropy loss due to the very high number of background pixels which causes imbalance. Random flip based data augmentation was used during training on images of size 384 x 384, validation performed every 25 epochs and each model was trained for a total of 250 epochs. Similar to segmentation and classification, we used Adam optimizer for training the object detection models as well. Images of size $512 \ge 512$ were used to train the object detectors for around 10000 steps for each dataset and with each step having a batch size of 32. The training process also involved data augmentation using random flips, random square crop, random padding with different combinations of random hue, saturation, contrast and brightness. Due to the stochastic nature of neural networks in all of the deep learning models, we trained all classification, segmentation as well as object detection models for a total of five runs with different random seed set each time, and the results are represented with the help of mean and standard deviation for each evaluation metric.



Figure 4: Illustration depicting a few random samples taken from different diabetic retinopathy grades for IDRiD dataset.



Figure 5: Histogram depicting number of image samples present for classification task for respective diabetic datasets.



Figure 6: Histogram depicting number of pixels belonging to each segmentation class for respective diabetic datasets.



Figure 7: Random samples generated with the help of modified RandAugment and after applying normalization pre-processing.

Madala		Cohen's Kappa	1	A	ROC A	UC
wodels	Unweighted	Linearly Weighted	Quadratic Weighted	Accuracy	Weighted-average	Macro-average
VGG16 [58]	0.4305 ± 0.0360	0.6516 ± 0.0175	0.5547 ± 0.0281	0.7706 ± 0.0394	0.8087 ± 0.0030	0.8251 ± 0.0038
VGG19 [58]	0.4514 ± 0.0053	0.6638 ± 0.0026	0.5718 ± 0.0034	0.7874 ± 0.0029	0.8078 ± 0.0038	0.8284 ± 0.0019
InceptionV3 [59]	0.4134 ± 0.0042	0.6357 ± 0.0050	0.5374 ± 0.0048	0.7814 ± 0.0024	0.8046 ± 0.0020	0.8261 ± 0.0010
ResNet50 [60]	0.4336 ± 0.0046	0.6514 ± 0.0048	0.5561 ± 0.0045	0.7906 ± 0.0015	0.8101 ± 0.0021	0.8302 ± 0.0016
ResNet50V2 [61]	0.4127 ± 0.0028	0.6341 ± 0.0034	0.5361 ± 0.0034	0.7782 ± 0.0029	0.8004 ± 0.0020	0.8224 ± 0.0021
ResNet152 [60]	0.4316 ± 0.0035	0.6465 ± 0.0014	0.5526 ± 0.0020	0.7914 ± 0.0012	0.8103 ± 0.0011	0.8305 ± 0.0005
ResNet101 [60]	0.4317 ± 0.0050	0.6488 ± 0.0061	0.5539 ± 0.0057	0.7913 ± 0.0022	0.8097 ± 0.0018	0.8306 ± 0.0017
ResNet152V2 [61]	0.4266 ± 0.0051	0.6456 ± 0.0042	0.5493 ± 0.0043	0.7846 ± 0.0017	0.8069 ± 0.0018	0.8252 ± 0.0012
ResNet101V2 [61]	0.4229 ± 0.0028	0.6435 ± 0.0011	0.5464 ± 0.0017	0.7827 ± 0.0031	0.8026 ± 0.0015	0.8236 ± 0.0007
Xception [62]	0.4371 ± 0.0019	0.6587 ± 0.0033	0.5618 ± 0.0027	0.7906 ± 0.0014	0.8139 ± 0.0010	0.8310 ± 0.0005
InceptionResNetV2 [63]	0.4324 ± 0.0048	0.6513 ± 0.0049	0.5558 ± 0.0044	0.7962 ± 0.0010	0.8140 ± 0.0015	0.8320 ± 0.0009
MobileNetV2 [64]	0.4013 ± 0.0050	0.6345 ± 0.0049	0.5309 ± 0.0048	0.7608 ± 0.0058	0.8010 ± 0.0011	0.8227 ± 0.0013
DenseNet121 [65]	0.4465 ± 0.0023	0.6678 ± 0.0030	0.5716 ± 0.0027	0.7911 ± 0.0029	0.8189 ± 0.0018	0.8340 ± 0.0021
DenseNet169 [65]	0.4462 ± 0.0045	0.6655 ± 0.0044	0.5705 ± 0.0044	0.7963 ± 0.0020	0.8198 ± 0.0020	0.8334 ± 0.0017
DenseNet201 [65]	0.4408 ± 0.0071	0.6567 ± 0.0062	0.5628 ± 0.0065	0.7986 ± 0.0012	0.8199 ± 0.0016	0.8344 ± 0.0013
EfficientNetB0 [66]	0.4288 ± 0.0035	0.6579 ± 0.0046	0.5565 ± 0.0042	0.7713 ± 0.0028	0.8134 ± 0.0018	0.8317 ± 0.0013

Table 2: Illustration of classification results of various pre-trained network on Kaggle Dataset using Cohen's Kappa and ROC AUC evaluation metrics.

Table 3: Illustration of classification results of various pre-trained network on DDR Dataset using Cohen's Kappa and ROC AUC evaluation metrics.

Madala		Cohen's Kappa	1	A	ROC A	UC
wodels	Unweighted	Linearly Weighted	Quadratic Weighted	Accuracy	Weighted-average	Macro-average
VGG16 [58]	0.6228 ± 0.0099	0.8451 ± 0.0059	0.7499 ± 0.0078	0.7589 ± 0.0062	0.9165 ± 0.0029	0.9013 ± 0.0025
VGG19 [58]	0.6330 ± 0.0067	0.8490 ± 0.0031	0.7574 ± 0.0042	0.7646 ± 0.0041	0.9135 ± 0.0036	0.9048 ± 0.0029
InceptionV3 [59]	0.5662 ± 0.0050	0.8205 ± 0.0039	0.7092 ± 0.0045	0.7249 ± 0.0026	0.8936 ± 0.0037	0.8840 ± 0.0034
ResNet50 [60]	0.5950 ± 0.0048	0.8314 ± 0.0056	0.7287 ± 0.0052	0.7427 ± 0.0030	0.9089 ± 0.0020	0.8956 ± 0.0049
ResNet50V2 [61]	0.5826 ± 0.0133	0.8261 ± 0.0049	0.7198 ± 0.0091	0.7350 ± 0.0077	0.8989 ± 0.0014	0.8894 ± 0.0036
ResNet152 [60]	0.5934 ± 0.0140	0.8320 ± 0.0088	0.7282 ± 0.0118	0.7422 ± 0.0081	0.9093 ± 0.0041	0.8939 ± 0.0019
ResNet101 [60]	0.5909 ± 0.0081	0.8285 ± 0.0055	0.7251 ± 0.0066	0.7408 ± 0.0049	0.9119 ± 0.0026	0.8961 ± 0.0037
ResNet152V2 [61]	0.5943 ± 0.0056	0.8325 ± 0.0023	0.7292 ± 0.0037	0.7424 ± 0.0037	0.9112 ± 0.0032	0.8977 ± 0.0017
ResNet101V2 [61]	0.5909 ± 0.0091	0.8307 ± 0.0056	0.7261 ± 0.0072	0.7398 ± 0.0055	0.9086 ± 0.0017	0.8965 ± 0.0013
Xception [62]	0.5856 ± 0.0199	0.8273 ± 0.0082	0.7221 ± 0.0137	0.7365 ± 0.0120	0.8981 ± 0.0033	0.8826 ± 0.0013
InceptionResNetV2 [63]	0.5903 ± 0.0137	0.8296 ± 0.0067	0.7256 ± 0.0099	0.7403 ± 0.0082	0.9012 ± 0.0021	0.8817 ± 0.0049
MobileNetV2 [64]	0.5064 ± 0.0290	0.7965 ± 0.0151	0.6689 ± 0.0215	0.6898 ± 0.0170	0.8880 ± 0.0041	0.8694 ± 0.0072
DenseNet121 [65]	0.6102 ± 0.0061	0.8393 ± 0.0023	0.7402 ± 0.0033	0.7514 ± 0.0038	0.9139 ± 0.0027	0.8985 ± 0.0065
DenseNet169 [65]	0.6043 ± 0.0068	0.8376 ± 0.0043	0.7367 ± 0.0051	0.7485 ± 0.0039	0.9119 ± 0.0029	0.8969 ± 0.0029
DenseNet201 [65]	0.6172 ± 0.0045	0.8433 ± 0.0032	0.7458 ± 0.0031	0.7568 ± 0.0026	0.9129 ± 0.0021	0.8907 ± 0.0019
EfficientNetB0 [66]	0.5910 ± 0.0059	0.8364 ± 0.0031	0.7306 ± 0.0042	0.7374 ± 0.0040	0.9091 ± 0.0042	0.8962 ± 0.0058

Table 4: Illustration of classification results of various pre-trained network on IDRiD Dataset using Cohen's Kappa and ROC AUC evaluation metrics.

Models		Cohen's Kappa	L	Accuracy	ROC AUC		
Models	Unweighted	Linearly Weighted	Quadratic Weighted	Accuracy	Weighted-average	Macro-average	
VGG16 [58]	0.3938 ± 0.0500	0.5662 ± 0.0600	0.4914 ± 0.0538	0.5728 ± 0.0357	0.8114 ± 0.0048	0.7939 ± 0.0074	
VGG19 [58]	0.3793 ± 0.0303	0.5920 ± 0.0378	0.4979 ± 0.0243	0.5592 ± 0.0244	0.8136 ± 0.0032	0.7987 ± 0.0129	
InceptionV3 [59]	0.4327 ± 0.0371	0.6007 ± 0.0488	0.5319 ± 0.0341	0.5961 ± 0.0280	0.7841 ± 0.0110	0.7245 ± 0.0187	
ResNet50 [60]	0.3965 ± 0.0421	0.6079 ± 0.0485	0.5227 ± 0.0471	0.5709 ± 0.0278	0.8013 ± 0.0135	0.7561 ± 0.0226	
ResNet50V2 [61]	0.3921 ± 0.0481	0.5616 ± 0.0251	0.4888 ± 0.0298	0.5689 ± 0.0340	0.7838 ± 0.0132	0.7335 ± 0.0332	
ResNet152 [60]	0.4208 ± 0.0563	0.5986 ± 0.0306	0.5212 ± 0.0432	0.5883 ± 0.0404	0.7968 ± 0.0140	0.7539 ± 0.0141	
ResNet101 [60]	0.4447 ± 0.0472	0.6071 ± 0.0274	0.5389 ± 0.0366	0.6058 ± 0.0334	0.8037 ± 0.0123	0.7577 ± 0.0217	
ResNet152V2 [61]	0.3421 ± 0.0520	0.5373 ± 0.0264	0.4530 ± 0.0383	0.5340 ± 0.0370	0.7720 ± 0.0193	0.7134 ± 0.0288	
ResNet101V2 [61]	0.4066 ± 0.0341	0.5916 ± 0.0214	0.5120 ± 0.0131	0.5786 ± 0.0244	0.7735 ± 0.0153	0.7125 ± 0.0271	
Xception [62]	0.4182 ± 0.0276	0.5939 ± 0.0189	0.5116 ± 0.0222	0.5903 ± 0.0199	0.8006 ± 0.0091	0.7474 ± 0.0138	
InceptionResNetV2 [63]	0.3834 ± 0.0429	0.5632 ± 0.0341	0.4867 ± 0.0390	0.5650 ± 0.0310	0.7881 ± 0.0029	0.7560 ± 0.0084	
MobileNetV2 [64]	0.3768 ± 0.0347	0.5608 ± 0.0268	0.4838 ± 0.0275	0.5592 ± 0.0354	0.7759 ± 0.0096	0.7396 ± 0.0082	
DenseNet121 [65]	0.3882 ± 0.0243	0.6049 ± 0.0410	0.5092 ± 0.0318	0.5612 ± 0.0160	0.7878 ± 0.0099	0.7352 ± 0.0189	
DenseNet169 [65]	0.4244 ± 0.0241	0.6206 ± 0.0115	0.5352 ± 0.0127	0.5922 ± 0.0182	0.8021 ± 0.0060	0.7489 ± 0.0144	
DenseNet201 [65]	0.4089 ± 0.0285	0.5844 ± 0.0210	0.5102 ± 0.0244	0.5806 ± 0.0210	0.8034 ± 0.0065	0.7555 ± 0.0081	
EfficientNetB0 [66]	0.4089 ± 0.0057	0.6376 ± 0.0299	0.5375 ± 0.0192	0.5767 ± 0.0053	0.8059 ± 0.0096	0.7633 ± 0.0178	

Medels	F1 Score								
wodels	Class 0	Class 1	Class 2	Class 3	Class 4	Weighted Average			
VCC16 [58]	$0.8781~\pm$	$0.0935~\pm$	$0.5527~\pm$	$0.3429~\pm$	$0.5364~\pm$	$0.7543~\pm$			
VGG10 [56]	0.0275	0.0384	0.0286	0.0371	0.0086	0.0208			
VCC10 [58]	$0.8905~\pm$	$0.0804~\pm$	0.5701 \pm	0.3171 \pm	$0.5337~\pm$	$0.7644 \pm$			
VGG19 [56]	0.0019	0.0129	0.0033	0.0196	0.0194	0.0014			
IncontionV3 [50]	$0.8871~\pm$	$0.0659~\pm$	0.5171 \pm	$0.3257~\pm$	$0.5066~\pm$	$0.7526~\pm$			
inception v 5 [55]	0.0016	0.0054	0.0067	0.0133	0.0088	0.0009			
$\mathbf{P}_{os}\mathbf{N}_{ot}$ 50 [60]	0.8928 \pm	$0.0606~\pm$	$0.5366~\pm$	$0.3446~\pm$	0.5232 \pm	$0.7601 \pm$			
Residet30 [00]	0.0010	0.0046	0.0067	0.0164	0.0148	0.0010			
DecNetFOV2 [61]	$0.8850~\pm$	$0.0701~\pm$	$0.5150~\pm$	0.3335 \pm	$0.5252~\pm$	$0.7517~\pm$			
Residet 50 v 2 [01]	0.0019	0.0031	0.0057	0.0254	0.0056	0.0014			
DogNot152 [60]	0.8927 \pm	0.0544 \pm	0.5331 \pm	0.3494 \pm	$0.5357~\pm$	$0.7595~\pm$			
	0.0007	0.0028	0.0045	0.0053	0.0050	0.0005			
$\mathbf{P}_{os}\mathbf{N}_{ot}101$ [60]	0.8931 \pm	0.0470 \pm	$0.5348~\pm$	0.3444 \pm	0.5248 \pm	$0.7591~\pm$			
Residential [00]	0.0013	0.0090	0.0046	0.0219	0.0106	0.0019			
ResNet152V2 [61]	$0.8891~\pm$	0.0710 \pm	$0.5280~\pm$	0.3535 \pm	$0.5280~\pm$	$0.7572~\pm$			
	0.0010	0.0108	0.0060	0.0202	0.0070	0.0009			
PosNot101V2 [61]	$0.8879~\pm$	0.0701 \pm	$0.5260~\pm$	0.3408 \pm	$0.5290~\pm$	$0.7557~\pm$			
Residential v 2 [01]	0.0020	0.0039	0.0044	0.0101	0.0045	0.0011			
Vantian [62]	0.8928 \pm	$0.0657~\pm$	$0.5421~\pm$	0.3436 \pm	$0.5396~\pm$	$0.7616~\pm$			
Aception [62]	0.0011	0.0044	0.0035	0.0127	0.0095	0.0009			
Incention Dec Not V2 [62]	$0.8960~\pm$	$0.0506~\pm$	0.5310 \pm	$0.3350~\pm$	0.5315 \pm	$0.7609~\pm$			
inceptionResiverv2 [03]	0.0008	0.0076	0.0052	0.0184	0.0148	0.0012			
MabileNetV2 [64]	0.8748 \pm	0.1043 \pm	0.5180 \pm	0.3421 \pm	0.5077 \pm	$0.7468~\pm$			
MobileNet V 2 [04]	0.0032	0.0062	0.0031	0.0189	0.0251	0.0031			
DongoNot121 [65]	$0.8935~\pm$	$0.0745~\pm$	$0.5527~\pm$	$0.3530~\pm$	0.5474 \pm	$0.7647~\pm$			
Denservet121 [05]	0.0018	0.0068	0.0025	0.0158	0.0094	0.0013			
Domao Not 160 [65]	0.8964 \pm	$0.0525~\pm$	0.5494 \pm	$0.3488~\pm$	0.5511 \pm	$0.7648~\pm$			
Denservetros [05]	0.0010	0.0040	0.0063	0.0050	0.0064	0.0017			
Domao Not 201 [65]	$0.8970~\pm$	$0.0541~\pm$	0.5431 \pm	0.3430 \pm	0.5343 \pm	$0.7639~\pm$			
Denservet201 [05]	0.0008	0.0068	0.0077	0.0081	0.0158	0.0020			
Efficient Not B0 [66]	$0.8806~\pm$	0.1043 \pm	0.5411 \pm	0.3623 \pm	$0.5590~\pm$	$0.7562~\pm$			
EncientivetB0 [00]	0.0018	0.0028	0.0042	0.0071	0.0044	0.0018			

Table 5: Illustration of classification results of various pre-trained network on Kaggle Dataset using F1 Score evaluation metrics.

Table 6: Illustration of classification results of various pre-trained network on Kaggle Dataset using Indexed Balanced Accuracy(IBA) evaluation metrics.

N(I	ndex Bala	nced Accu	ıracy (IBA	A)
Models	Class 0	Class 1	Class 2	Class 3	Class 4	Weighted Average
VCC16 [F9]	0.5312 \pm	$0.0695~\pm$	$0.4990~\pm$	$0.2582~\pm$	0.4026 \pm	$0.5201 \pm$
VGG10 [58]	0.0058	0.0540	0.0118	0.0776	0.0563	0.0045
VCC10 [F9]	0.5426 \pm	0.0487 \pm	0.5182 \pm	0.2152 \pm	$0.3806~\pm$	$0.5304~\pm$
VGG19 [58]	0.0040	0.0101	0.0101	0.0241	0.0210	0.0032
IncontionV2 [50]	0.4914 \pm	$0.0375~\pm$	0.4350 \pm	0.2372 \pm	0.3632 \pm	$0.4930~\pm$
inception v 3 [59]	0.0110	0.0035	0.0161	0.0158	0.0079	0.0068
PosNot50 [60]	$0.5036~\pm$	0.0332 \pm	0.4531 \pm	0.2503 \pm	0.3758 \pm	$0.5041~\pm$
Residet50 [00]	0.0116	0.0033	0.0161	0.0178	0.0180	0.0074
BosNot50V2 [61]	$0.4975~\pm$	$0.0407~\pm$	$0.4400~\pm$	$0.2467~\pm$	0.3812 \pm	$0.4961~\pm$
itesivet50v2 [01]	0.0080	0.0028	0.0078	0.0262	0.0125	0.0047
ResNet152 [60]	0.4962 \pm	0.0291 \pm	0.4470 \pm	0.2525 \pm	$0.3850~\pm$	$0.4997~\pm$
	0.0092	0.0017	0.0113	0.0078	0.0039	0.0056
BesNet101 [60]	$0.4980~\pm$	0.0252 \pm	0.4494 \pm	$0.2507~\pm$	0.3729 \pm	$0.5006 \pm$
	0.0100	0.0054	0.0123	0.0256	0.0158	0.0064
$\mathbf{R}_{os}\mathbf{N}_{ot}15\mathbf{2V2}$ [61]	$0.5059~\pm$	$0.0405~\pm$	$0.4504~\pm$	$0.2637~\pm$	0.3812 \pm	$0.5038~\pm$
	0.0081	0.0075	0.0154	0.0178	0.0093	0.0054
BosNot101V2 [61]	$0.5045~\pm$	$0.0403~\pm$	$0.4499~\pm$	$0.2528~\pm$	$0.3789~\pm$	$0.5022 \pm$
	0.0075	0.0027	0.0126	0.0107	0.0054	0.0044
Xception [62]	0.5092 \pm	0.0371 \pm	$0.4582~\pm$	$0.2487~\pm$	0.3964 \pm	$0.5080 \pm$
Acception [02]	0.0028	0.0029	0.0055	0.0145	0.0092	0.0019
Incention ResNet V2 [63]	$0.4859~\pm$	0.0264 \pm	0.4319 \pm	0.2401 \pm	0.3762 \pm	$0.4942~\pm$
inceptionitesi (ci v 2 [00]	0.0084	0.0045	0.0130	0.0195	0.0186	0.0055
MobileNetV2 [64]	$0.5167~\pm$	$0.0758~\pm$	$0.4517~\pm$	0.2614 \pm	$0.3666~\pm$	$0.5045~\pm$
	0.0070	0.0072	0.0095	0.0241	0.0261	0.0033
DenseNet121 [65]	$0.5251 \pm$	$0.0434 \pm$	$0.4752 \pm$	$0.2613 \pm$	$0.4020 \pm$	$0.5190 \pm$
	0.0044	0.0048	0.0043	0.0224	0.0138	0.0023
DenseNet169 [65]	$0.5105~\pm$	$0.0283~\pm$	$0.4623~\pm$	$0.2541~\pm$	0.4016 \pm	$0.5106 \pm$
	0.0028	0.0025	0.0068	0.0101	0.0091	0.0024
DenseNet 201 [65]	$0.4924 \pm$	$0.0282 \pm$	$0.4450 \pm$	$0.2421 \pm$	$0.3860 \pm$	$0.4997 \pm$
	0.0118	0.0039	0.0142	0.0091	0.0185	0.0078
EfficientNetB0 [66]	0.5384 \pm	$0.0708~\pm$	$0.4852~\pm$	0.2804 \pm	$0.4309~\pm$	$0.5234 \pm$
	0.0018	0.0025	0.0039	0.0062	0.0053	0.0012

Modela	Geometric Mean (GMean)								
Models	Class 0	Class 1	Class 2	Class 3	Class 4	Weighted Average			
VCC16 [58]	0.7170 \pm	0.2632 \pm	0.7192 \pm	0.5228 \pm	$0.6521~\pm$	$0.7176 \pm$			
VGG10 [58]	0.0078	0.0901	0.0090	0.0708	0.0416	0.0018			
VCC10 [58]	0.7235 \pm	0.2304 \pm	0.7323 \pm	0.4820 \pm	$0.6358~\pm$	$0.7239~\pm$			
VGG19 [58]	0.0028	0.0246	0.0064	0.0258	0.0171	0.0022			
IncontionV2 [50]	$0.6861~\pm$	0.2034 \pm	0.6745 \pm	$0.5057~\pm$	0.6218 \pm	$0.6966 \pm$			
inception v 5 [59]	0.0084	0.0095	0.0115	0.0163	0.0065	0.0052			
DecNetE0 [60]	0.6945 \pm	0.1914 \pm	$0.6879~\pm$	0.5191 \pm	0.6320 \pm	$0.7043~\pm$			
Residet 50 [00]	0.0088	0.0096	0.0113	0.0180	0.0145	0.0056			
DogNot50V2 [61]	0.6910 \pm	0.2117 \pm	$0.6780~\pm$	$0.5150~\pm$	0.6364 \pm	$0.6991~\pm$			
Itesivet50 v 2 [01]	0.0061	0.0072	0.0056	0.0267	0.0099	0.0037			
PosNot152 [60]	$0.6890~\pm$	0.1792 \pm	$0.6836~\pm$	0.5215 \pm	$0.6395~\pm$	$0.7009~\pm$			
itesivet152 [00]	0.0069	0.0052	0.0080	0.0078	0.0031	0.0042			
ResNet101 [60]	$0.6903~\pm$	$0.1661~\pm$	$0.6853~\pm$	$0.5192~\pm$	0.6297 \pm	$0.7016~\pm$			
	0.0075	0.0180	0.0087	0.0259	0.0128	0.0048			
ResNet152V2 [61]	$0.6967~\pm$	0.2108 \pm	$0.6857~\pm$	0.5325 \pm	$0.6365~\pm$	$0.7045~\pm$			
	0.0060	0.0188	0.0109	0.0175	0.0074	0.0040			
PosNot101V2 [61]	$0.6958~\pm$	0.2107 \pm	$0.6853~\pm$	0.5218 \pm	0.6346 \pm	$0.7034~\pm$			
Residential v 2 [01]	0.0058	0.0071	0.0088	0.0107	0.0044	0.0034			
Vantion [62]	$0.6987~\pm$	0.2023 \pm	0.6917 \pm	$0.5176~\pm$	$0.6485~\pm$	$0.7072~\pm$			
Aception [02]	0.0022	0.0079	0.0039	0.0147	0.0072	0.0015			
Incontion Dec Not V2 [62]	$0.6809~\pm$	0.1704 \pm	$0.6729~\pm$	$0.5086~\pm$	0.6323 \pm	$0.6965~\pm$			
inceptionitesivet v 2 [05]	0.0063	0.0144	0.0094	0.0207	0.0149	0.0041			
MobileNetV2 [64]	$0.7066~\pm$	0.2878 \pm	$0.6862~\pm$	$0.5299~\pm$	0.6243 \pm	$0.7066 \pm$			
Mobilei (et v 2 [04]	0.0056	0.0133	0.0065	0.0242	0.0215	0.0028			
DonsoNot121 [65]	0.7103 \pm	0.2184 \pm	$0.7036~\pm$	$0.5299~\pm$	$0.6528~\pm$	$0.7153~\pm$			
Denserveti21 [05]	0.0034	0.0121	0.0028	0.0217	0.0107	0.0018			
DonsoNot169 [65]	0.6992 \pm	$0.1767~\pm$	0.6948 \pm	0.5231 \pm	0.6526 \pm	$0.7087~\pm$			
Denservetros [05]	0.0020	0.0077	0.0048	0.0100	0.0070	0.0017			
DonsoNot201 [65]	$0.6856~\pm$	$0.1764~\pm$	$0.6826~\pm$	0.5110 \pm	$0.6402~\pm$	$0.7005~\pm$			
Denserver201 [00]	0.0088	0.0119	0.0103	0.0094	0.0146	0.0058			
Efficient Not B0 [66]	0.7219 \pm	$0.2786~\pm$	0.7097 \pm	0.5487 \pm	$0.6749~\pm$	$0.7199~\pm$			
EfficientNetB0 [66]	0.0014	0.0048	0.0027	0.0059	0.0040	0.0008			

Table 7: Illustration of classification results of various pre-trained network on Kaggle Dataset using GMean evaluation metrics.

Table 8: Illustration of classification results of various pre-trained network on DDR Dataset using F1 Score evaluation metrics.

Madala	F1 Score							
Models	Class 0	Class 1	Class 2	Class 3	Class 4	Class 5	Weighted Average	
VCC16 [F9]	$0.8487~\pm$	0.0610 \pm	$0.6592~\pm$	$0.2698~\pm$	0.7188 \pm	$0.8606~\pm$	$0.7327 \pm$	
VGG10 [58]	0.0044	0.0294	0.0097	0.0191	0.0220	0.0086	0.0069	
VCC10 [59]	$0.8588~\pm$	$0.0609~\pm$	0.6741 \pm	$0.2879~\pm$	0.6971 \pm	$0.8561~\pm$	$0.7407~\pm$	
VGG19 [58]	0.0027	0.0153	0.0071	0.0322	0.0207	0.0080	0.0044	
InceptionV3 [59]	0.8292 \pm	$0.0360~\pm$	$0.5780~\pm$	$0.2967~\pm$	$0.6470~\pm$	$0.8671~\pm$	$0.6922 \pm$	
	0.0016	0.0093	0.0098	0.0348	0.0133	0.0066	0.0044	
$\mathbf{P}_{oc}\mathbf{N}_{ot}$	$0.8392~\pm$	0.0316 \pm	0.6208 \pm	0.3520 \pm	0.6774 \pm	$0.8636~\pm$	$0.7133 \pm$	
Residet50 [60]	0.0041	0.0066	0.0104	0.0383	0.0193	0.0062	0.0037	
PogNot50V2 [61]	0.8338 \pm	$0.0507~\pm$	$0.5976~\pm$	0.2816 \pm	$0.6895~\pm$	$0.8611~\pm$	$0.7035~\pm$	
ResNet50V2 [61]	0.0062	0.0060	0.0202	0.0672	0.0122	0.0038	0.0102	
PocNot152 [60]	$0.8393~\pm$	$0.0387~\pm$	0.6182 \pm	$0.2760~\pm$	$0.6683~\pm$	0.8622 \pm	$0.7108~\pm$	
ResNet152 [60]	0.0064	0.0117	0.0207	0.0316	0.0106	0.0070	0.0100	
ResNet101 [60]	$0.8381~\pm$	$0.0362~\pm$	$0.6145~\pm$	$0.2750~\pm$	$0.6700~\pm$	$0.8666~\pm$	$0.7093~\pm$	
	0.0028	0.0139	0.0118	0.0288	0.0078	0.0105	0.0060	
PocNot152V2 [61]	$0.8386~\pm$	0.0423 \pm	$0.6131~\pm$	$0.2682~\pm$	$0.7081~\pm$	$0.8651~\pm$	$0.7118 \pm$	
11031(0110272 [01]	0.0037	0.0171	0.0101	0.0389	0.0155	0.0067	0.0039	
BesNet101V2 [61]	$0.8393~\pm$	$0.0599~\pm$	$0.6087~\pm$	0.3112 \pm	$0.6877~\pm$	$0.8550~\pm$	$0.7100 \pm$	
itesitetiti v 2 [01]	0.0067	0.0119	0.0133	0.0598	0.0241	0.0045	0.0065	
X ception [62]	$0.8342~\pm$	$0.0263~\pm$	$0.6031~\pm$	$0.3615~\pm$	0.6794 \pm	$0.8643~\pm$	$0.7054~\pm$	
Acception [02]	0.0091	0.0064	0.0295	0.0366	0.0059	0.0037	0.0146	
Inception ResNet V2 [63]	$0.8371~\pm$	$0.0260~\pm$	$0.6083~\pm$	$0.2460~\pm$	$0.6867~\pm$	$0.8728~\pm$	$0.7076 \pm$	
	0.0062	0.0148	0.0227	0.0265	0.0115	0.0038	0.0106	
MobileNetV2 [64]	$0.8136 \pm$	$0.0220~\pm$	$0.5256 \pm$	0.3024 \pm	$0.3582 \pm$	$0.8069 \pm$	$0.6429 \pm$	
	0.0131	0.0185	0.0488	0.0669	0.0788	0.0184	0.0226	
DenseNet121 [65]	$0.8473~\pm$	$0.0482~\pm$	$0.6322~\pm$	$0.3278~\pm$	$0.7048~\pm$	$0.8663 \pm$	$0.7231 \pm$	
Denservetizi [00]	0.0044	0.0054	0.0105	0.0314	0.0183	0.0037	0.0043	
DenseNet169 [65]	$0.8425~\pm$	$0.0431~\pm$	$0.6253~\pm$	$0.3165~\pm$	$0.7038~\pm$	$0.8679 \pm$	$0.7183 \pm$	
Demserverroo [00]	0.0031	0.0148	0.0069	0.0084	0.0162	0.0027	0.0049	
DenseNet201 [65]	$0.8489~\pm$	0.0223 \pm	$0.6461~\pm$	$0.3029~\pm$	$0.7010~\pm$	$0.8686~\pm$	$0.7268 \pm$	
Denserver201 [00]	0.0039	0.0123	0.0058	0.0281	0.0178	0.0055	0.0027	
EfficientNetB0 [66]	$0.8399~\pm$	$0.0830~\pm$	$0.5928~\pm$	$0.3961~\pm$	$0.7205~\pm$	$0.8668~\pm$	$0.7107~\pm$	
EnicientNetB0 [66]	0.0035	0.0125	0.0068	0.0174	0.0112	0.0041	0.0035	

	Index Balanced Accuracy (IBA)									
Models	Class 0	Class 1	Class 2	Class 3	Class 4	Class 5	Weighted Average			
VCC16 [59]	$0.7278~\pm$	0.0342 \pm	$0.5072~\pm$	$0.1780~\pm$	0.6238 \pm	0.9103 \pm	$0.6384 \pm$			
VGG10 [58]	0.0084	0.0176	0.0118	0.0111	0.0312	0.0087	0.0081			
VCC10 [F9]	0.7494 \pm	0.0341 \pm	$0.5303~\pm$	0.1912 \pm	0.6148 \pm	$0.8866~\pm$	$0.6499~\pm$			
VGG19 [58]	0.0059	0.0085	0.0110	0.0146	0.0397	0.0092	0.0057			
Incontion V2 [F0]	$0.6842~\pm$	0.0199 \pm	0.4181 \pm	$0.1966~\pm$	$0.5329~\pm$	0.9194 \pm	$0.5908~\pm$			
Inception v 3 [59]	0.0046	0.0052	0.0112	0.0302	0.0218	0.0066	0.0045			
PosNot50 [60]	$0.7065~\pm$	0.0170 \pm	0.4654 \pm	0.2393 \pm	0.5574 \pm	0.9098 \pm	$0.6142~\pm$			
itesivet50 [00]	0.0091	0.0042	0.0136	0.0360	0.0255	0.0062	0.0047			
ResNet50V2 [61]	0.6936 \pm	0.0275 \pm	0.4373 \pm	0.1914 \pm	0.5735 \pm	0.9193 \pm	$0.6035~\pm$			
	0.0150	0.0039	0.0232	0.0514	0.0094	0.0064	0.0118			
BogNot152 [60]	$0.7051~\pm$	0.0199 \pm	0.4612 \pm	0.1808 \pm	$0.5552~\pm$	$0.9106~\pm$	$0.6127~\pm$			
ResNet152 [60]	0.0153	0.0062	0.0238	0.0321	0.0137	0.0158	0.0124			
ResNet101 [60]	0.7017 \pm	0.0190 \pm	0.4554 \pm	$0.1859~\pm$	$0.5587~\pm$	$0.9081~\pm$	$0.6104~\pm$			
	0.0070	0.0075	0.0144	0.0239	0.0068	0.0131	0.0068			
ResNet152V2 [61]	$0.7038~\pm$	0.0228 \pm	0.4544 \pm	$0.1729~\pm$	0.5979 \pm	0.9139 \pm	$0.6129~\pm$			
	0.0072	0.0091	0.0103	0.0302	0.0193	0.0079	0.0048			
BosNot101V2 [61]	$0.7057~\pm$	$0.0332~\pm$	$0.4501~\pm$	0.2074 \pm	$0.5761~\pm$	$0.9081~\pm$	$0.6112 \pm$			
itesivetioiv 2 [01]	0.0151	0.0067	0.0164	0.0502	0.0201	0.0050	0.0086			
Xception [62]	$0.6953~\pm$	0.0142 \pm	$0.4437~\pm$	$0.2473~\pm$	$0.5651~\pm$	0.9294 \pm	$0.6062 \pm$			
Aception [02]	0.0209	0.0034	0.0342	0.0315	0.0128	0.0074	0.0172			
Inception ResNet V2 [63]	$0.7000~\pm$	0.0133 \pm	$0.4498~\pm$	$0.1622~\pm$	$0.5794~\pm$	$0.9238~\pm$	$0.6094 \pm$			
inceptionitesi (et v 2 [00]	0.0146	0.0078	0.0260	0.0237	0.0134	0.0136	0.0120			
MobileNetV2 [64]	0.6464 \pm	0.0114 \pm	$0.3680~\pm$	$0.2759~\pm$	0.2112 \pm	$0.9336~\pm$	$0.5470 \pm$			
widdhei ver v 2 [04]	0.0353	0.0104	0.0556	0.1484	0.0589	0.0044	0.0237			
DenseNet121 [65]	$0.7230~\pm$	$0.0265~\pm$	$0.4766~\pm$	$0.2179~\pm$	$0.5991~\pm$	0.9220 \pm	$0.6275~\pm$			
Denserveti21 [00]	0.0094	0.0026	0.0142	0.0286	0.0301	0.0122	0.0059			
DenseNet169 [65]	$0.7115 \pm$	$0.0228~\pm$	$0.4662 \pm$	$0.2177 \pm$	$0.6017 \pm$	$0.9190~\pm$	$0.6212 \pm$			
Denserverios [05]	0.0073	0.0085	0.0090	0.0109	0.0259	0.0096	0.0059			
DenseNet201 [65]	$0.7250~\pm$	0.0114 \pm	0.4912 \pm	$0.2045~\pm$	$0.5939~\pm$	0.9168 \pm	$0.6324 \pm$			
Denserver201 [00]	0.0091	0.0064	0.0072	0.0197	0.0291	0.0050	0.0042			
EfficientNetB0 [66]	$0.7079~\pm$	$0.0537~\pm$	$0.4285~\pm$	$0.3139~\pm$	0.6233 \pm	$0.9316~\pm$	$0.6124 \pm$			
Encientivet Do [00]	0.0066	0.0086	0.0076	0.0315	0.0108	0.0072	0.0047			

Table 9: Illustration of classification results of various pre-trained network on DDR Dataset using Indexed Balanced Accuracy(IBA) evaluation metrics.

Table 10: Illustration of classification results of various pre-trained network on DDR Dataset using GMean evaluation metrics.

			Ge	ometric M	lean (GM	ean)	
Models	Class 0	Class 1	Class 2	Class 3	Class 4	Class 5	Weighted Average
NGG16 [F9]	0.8431 \pm	$0.1876~\pm$	$0.7250~\pm$	$0.4397~\pm$	$0.8032~\pm$	$0.9562~\pm$	$0.8026 \pm$
VGG16 [58]	0.0051	0.0561	0.0080	0.0133	0.0191	0.0042	0.0050
MOG10 [F9]	$0.8569~\pm$	$0.1931~\pm$	$0.7398~\pm$	$0.4553~\pm$	$0.7974~\pm$	$0.9449~\pm$	$0.8099~\pm$
VGG19 [58]	0.0038	0.0233	0.0069	0.0169	0.0236	0.0045	0.0035
Incention V2 [50]	$0.8153~\pm$	$0.1472~\pm$	$0.6611~\pm$	$0.4608~\pm$	$0.7458~\pm$	$0.9605~\pm$	$0.7725~\pm$
Inception v 3 [59]	0.0034	0.0200	0.0083	0.0336	0.0144	0.0032	0.0030
ResNet50 [60]	$0.8295~\pm$	$0.1366~\pm$	$0.6956~\pm$	$0.5070~\pm$	$0.7619~\pm$	$0.9560~\pm$	$0.7874~\pm$
	0.0059	0.0163	0.0094	0.0378	0.0166	0.0031	0.0030
ResNet50V2 [61]	0.8211 \pm	0.1741 \pm	$0.6756~\pm$	0.4520 \pm	0.7724 \pm	0.9604 \pm	$0.7805~\pm$
	0.0098	0.0124	0.0170	0.0630	0.0059	0.0030	0.0077
ResNet152 [60]	0.8282 \pm	0.1468 \pm	0.6927 \pm	0.4418 \pm	$0.7605~\pm$	$0.9563~\pm$	$0.7863~\pm$
	0.0099	0.0258	0.0170	0.0389	0.0089	0.0075	0.0080
ResNet101 [60]	$0.8259~\pm$	0.1425 \pm	$0.6889~\pm$	$0.4486~\pm$	0.7628 \pm	$0.9553~\pm$	$0.7849~\pm$
	0.0046	0.0296	0.0101	0.0283	0.0043	0.0064	0.0043
ResNet152V2 [61]	0.8274 \pm	$0.1557~\pm$	$0.6881~\pm$	0.4323 \pm	$0.7876~\pm$	$0.9580~\pm$	$0.7865~\pm$
	0.0043	0.0345	0.0075	0.0381	0.0118	0.0038	0.0030
BooNot101V2 [61]	$0.8287~\pm$	$0.1908~\pm$	$0.6849~\pm$	$0.4708~\pm$	$0.7739~\pm$	$0.9550~\pm$	$0.7855~\pm$
	0.0097	0.0189	0.0118	0.0594	0.0128	0.0024	0.0056
X contion [62]	0.8224 \pm	$0.1247~\pm$	$0.6800~\pm$	$0.5156~\pm$	$0.7669~\pm$	$0.9652~\pm$	$0.7822~\pm$
Aception [02]	0.0135	0.0150	0.0248	0.0310	0.0081	0.0035	0.0110
Incontion ResNot V2 [63]	$0.8249~\pm$	$0.1170~\pm$	$0.6846~\pm$	0.4194 \pm	$0.7760~{\pm}$	$0.9627~\pm$	$0.7842~\pm$
inceptionitesivet v 2 [00]	0.0094	0.0361	0.0188	0.0305	0.0084	0.0064	0.0077
MobileNetV2 [64]	$0.7907~\pm$	$0.1042~\pm$	$0.6203 \pm$	$0.5276~\pm$	$0.4742~\pm$	$0.9656 \pm$	$0.7437 \pm$
	0.0237	0.0469	0.0434	0.1402	0.0664	0.0019	0.0161
DenseNet121 [65]	$0.8396 \pm$	$0.1711 \pm$	$0.7038~\pm$	$0.4847 \pm$	$0.7881 \pm$	$0.9617~\pm$	$0.7958~\pm$
	0.0059	0.0085	0.0096	0.0326	0.0186	0.0057	0.0037
DenseNet169 [65]	$0.8321~\pm$	$0.1565 \pm$	$0.6968~\pm$	$0.4852~\pm$	$0.7897~\pm$	$0.9604~\pm$	$0.7917~\pm$
Denser(etrop [00]	0.0047	0.0300	0.0063	0.0118	0.0159	0.0045	0.0038
DenseNet201 [65]	0.8404 \pm	$0.1083~\pm$	$0.7142~\pm$	$0.4703~\pm$	$0.7849~\pm$	0.9594 \pm	$0.7987~\pm$
	0.0058	0.0335	0.0048	0.0220	0.0179	0.0023	0.0027
EfficientNetB0 [66]	$0.8303~\pm$	0.2426 \pm	$0.6697~\pm$	$0.5790~\pm$	$0.8031~\pm$	$0.9662~\pm$	$0.7865 \pm$
	0.0039	0.0196	0.0056	0.0273	0.0065	0.0033	0.0029

Madala	F1 Score								
Wodels	Class 0	Class 1	Class 2	Class 3	Class 4	Weighted Average			
VCC16 [58]	0.7112 \pm	$0.0000~\pm$	$0.5901~\pm$	$0.4909~\pm$	$0.2259~\pm$	$0.5371~\pm$			
VGG10 [58]	0.0537	0.0000	0.0266	0.0176	0.1669	0.0385			
VCC10 [58]	$0.6759~\pm$	$0.0000~\pm$	$0.5764~\pm$	$0.4809~\pm$	0.3474 \pm	$0.5347~\pm$			
V GG19 [56]	0.0274	0.0000	0.0388	0.0758	0.0419	0.0202			
IncontionV3 [59]	0.7322 \pm	$0.0000~\pm$	$0.6058~\pm$	0.5716 \pm	$0.2500~\pm$	$0.5669 \pm$			
inception vo [00]	0.0336	0.0000	0.0288	0.0249	0.0859	0.0283			
\mathbf{B} osNot50 [60]	$0.7461~\pm$	$0.0000~\pm$	$0.5911~\pm$	0.4537 \pm	0.1732 \pm	$0.5355~\pm$			
	0.0275	0.0000	0.0306	0.0808	0.1196	0.0323			
\mathbf{B} osNot50V2 [61]	$0.6768~\pm$	$0.0000~\pm$	$0.5963~\pm$	$0.5574~\pm$	$0.2089~\pm$	$0.5378~\pm$			
	0.0250	0.0000	0.0207	0.1054	0.1419	0.0379			
BesNet152 [60]	0.7228 \pm	$0.0000~\pm$	0.6128 \pm	$0.5330~\pm$	$0.2599~\pm$	$0.5601 \pm$			
	0.0201	0.0000	0.0587	0.0670	0.0483	0.0380			
BesNet101 [60]	$0.7287~\pm$	$0.0500~\pm$	$0.6263~\pm$	$0.5679~\pm$	0.2704 \pm	$0.5764~\pm$			
itesitetioi [00]	0.0331	0.1118	0.0397	0.0602	0.0375	0.0326			
ResNet152V2 [61]	$0.6520~\pm$	$0.0000~\pm$	$0.5840~\pm$	0.4406 \pm	$0.1948~\pm$	$0.5025~\pm$			
	0.0385	0.0000	0.0517	0.0693	0.0450	0.0346			
BosNot101V2 [61]	$0.6897~\pm$	$0.0000~\pm$	$0.5941~\pm$	0.5423 \pm	0.3076 \pm	$0.5511~\pm$			
	0.0186	0.0000	0.0362	0.0698	0.0483	0.0253			
Xception [62]	0.7218 \pm	$0.0000~\pm$	$0.5945~\pm$	$0.5507~\pm$	0.2732 \pm	$0.5590~\pm$			
Keeption [02]	0.0355	0.0000	0.0218	0.0266	0.0644	0.0172			
Incontion RosNotV2 [63]	$0.6947~\pm$	$0.0000~\pm$	$0.6088~\pm$	0.4599 \pm	$0.1665~\pm$	$0.5243~\pm$			
inceptionitesi (et v 2 [00]	0.0343	0.0000	0.0294	0.0629	0.0457	0.0263			
MobileNetV2 [64]	$0.7046~\pm$	0.0421 \pm	$0.5600~\pm$	0.4197 \pm	0.2943 \pm	$0.5232~\pm$			
MODIFILIEU Z [04]	0.0396	0.0942	0.0335	0.0462	0.0487	0.0164			
DenseNet121 [65]	$0.7039~\pm$	$0.0500~\pm$	$0.5759~\pm$	0.4787 \pm	0.3117 \pm	$0.5413~\pm$			
Denser(et121 [00]	0.0300	0.1118	0.0229	0.0609	0.0923	0.0198			
DenseNet169 [65]	$0.7349~\pm$	$0.0000 \pm$	$0.6012 \pm$	$0.5069 \pm$	$0.3032 \pm$	$0.5611 \pm$			
Denser(erros [00]	0.0175	0.0000	0.0200	0.0692	0.0616	0.0171			
DonsoNot201 [65]	0.7147 \pm	$0.0000~\pm$	$0.6057~\pm$	0.5272 \pm	0.2191 \pm	$0.5490 \pm$			
Denservet201 [00]	0.0309	0.0000	0.0205	0.0368	0.0531	0.0209			
Efficient Net B0 [66]	0.7148 \pm	$0.0000~\pm$	0.5450 \pm	$0.5358~\pm$	0.4210 \pm	$0.5573~\pm$			
Emclentiverbo [00]	0.0179	0.0000	0.0100	0.0748	0.0347	0.0058			

Table 11: Illustration of classification results of various pre-trained network on IDRiD Dataset using F1 Score evaluation metrics.

Table 12: Illustration of classification results of various pre-trained network on IDRiD Dataset using Indexed Balanced Accuracy (IBA) evaluation metrics.

		I	ndex Bala	nced Accu	iracy (IBA	A)
Models	Class 0	Class 1	Class 2	Class 3	Class 4	Weighted Average
VCC16 [58]	$0.6088~\pm$	$0.0000~\pm$	0.4857 \pm	0.3579 \pm	0.1395 \pm	$0.4579~\pm$
VGG10 [58]	0.0748	0.0000	0.0353	0.0206	0.1110	0.0368
VCC10 [F9]	$0.5667~\pm$	$0.0000~\pm$	0.4711 \pm	$0.3559~\pm$	0.2227 \pm	$0.4453~\pm$
VGG19 [58]	0.0354	0.0000	0.0471	0.0739	0.0324	0.0216
IncontionV3 [50]	0.6451 \pm	$0.0000~\pm$	0.5078 \pm	0.4544 \pm	0.1520 \pm	$0.4869 \pm$
inception v 5 [59]	0.0450	0.0000	0.0375	0.0257	0.0583	0.0276
BogNotf0 [60]	0.6644 \pm	$0.0000~\pm$	0.4873 \pm	0.3497 \pm	0.1093 \pm	$0.4623~\pm$
Residet30 [00]	0.0423	0.0000	0.0374	0.0853	0.0784	0.0313
ResNet50V2 [61]	0.5779 \pm	$0.0000~\pm$	0.4935 \pm	0.4229 \pm	0.1370 \pm	$0.4559~\pm$
	0.0344	0.0000	0.0261	0.1064	0.0983	0.0339
ResNet152 [60]	0.6175 \pm	$0.0000~\pm$	$0.5185~\pm$	0.4143 \pm	0.1525 \pm	$0.4784~\pm$
	0.0299	0.0000	0.0765	0.0646	0.0316	0.0420
$\mathbf{R}_{os}\mathbf{N}_{ot}101$ [60]	0.6328 \pm	0.0361 \pm	$0.5360~\pm$	0.4466 \pm	$0.1536~\pm$	$0.4958~\pm$
Itesivetion [00]	0.0448	0.0808	0.0533	0.0733	0.0309	0.0367
DocNot152V2 [61]	$0.5381~\pm$	$0.0000~\pm$	0.4811 \pm	0.3029 \pm	0.1220 \pm	$0.4206 \pm$
Resiver152 v 2 [01]	0.0456	0.0000	0.0640	0.0575	0.0304	0.0380
BosNot101V2 [61]	$0.5895~\pm$	$0.0000~\pm$	$0.4929~\pm$	$0.4244~\pm$	$0.1939~\pm$	$0.4658 \pm$
Itesivetioi v 2 [01]	0.0284	0.0000	0.0460	0.0656	0.0320	0.0252
X contion [62]	0.6282 \pm	$0.0000~\pm$	0.4934 \pm	$0.3957~\pm$	$0.1662~\pm$	$0.4749~\pm$
Aception [02]	0.0510	0.0000	0.0279	0.0221	0.0399	0.0215
Incontion Ros Not V2 [63]	$0.5958~\pm$	$0.0000~\pm$	$0.5098~\pm$	$0.3366~\pm$	$0.0958~\pm$	$0.4512~\pm$
inceptionitesivet v 2 [05]	0.0520	0.0000	0.0362	0.0750	0.0366	0.0320
MabilaNatV2 [64]	0.6228 \pm	$0.0669~\pm$	0.4490 \pm	0.2604 \pm	$0.1803~\pm$	$0.4444 \pm$
Mobilet (et v 2 [04]	0.0595	0.1495	0.0407	0.0426	0.0380	0.0271
DenseNet121 [65]	$0.5930~\pm$	$0.0361~\pm$	0.4710 \pm	0.3701 \pm	$0.2188~\pm$	$0.4529 \pm$
	0.0354	0.0808	0.0288	0.0730	0.0897	0.0181
DenseNet169 [65]	0.6450 \pm	$0.0000~\pm$	0.5014 \pm	0.3781 \pm	$0.1936~\pm$	$0.4804 \pm$
Denservetros [05]	0.0225	0.0000	0.0264	0.0914	0.0576	0.0176
DonsoNot 201 [65]	$0.6089~\pm$	$0.0000~\pm$	0.5082 \pm	0.4118 \pm	$0.1247~\pm$	$0.4692~\pm$
Denservet201 [05]	0.0429	0.0000	0.0268	0.0389	0.0314	0.0214
Efficient Net B0 [66]	0.6247 \pm	$0.0000~\pm$	0.4319 \pm	0.4296 \pm	0.2929 \pm	$0.4664~\pm$
Encienti (etbo [00]	0.0266	0.0000	0.0103	0.0887	0.0559	0.0042

M	Geometric Mean (GMean)								
wodels	Class 0	Class 1	Class 2	Class 3	Class 4	Weighted Average			
VCC16 [F9]	0.7834 \pm	$0.0000~\pm$	0.6915 \pm	0.6151 \pm	0.3376 \pm	$0.6847~\pm$			
VGG10 [58]	0.0453	0.0000	0.0254	0.0164	0.2145	0.0270			
VCC10 [59]	$0.7565~\pm$	$0.0000~\pm$	$0.6838~\pm$	0.6107 \pm	$0.4892~\pm$	$0.6760 \pm$			
VGG19 [58]	0.0223	0.0000	0.0326	0.0586	0.0332	0.0157			
Incention V2 [50]	0.8040 \pm	$0.0000~\pm$	0.7122 \pm	$0.6890~\pm$	0.3994 \pm	$0.7061~\pm$			
inception v 3 [59]	0.0273	0.0000	0.0243	0.0185	0.0822	0.0194			
DecNetE0 [60]	0.8155 \pm	$0.0000~\pm$	$0.6980~\pm$	0.6033 \pm	0.3028 \pm	$0.6886~\pm$			
Residet50 [00]	0.0240	0.0000	0.0275	0.0728	0.1835	0.0228			
ResNet50V2 [61]	$0.7600~\pm$	$0.0000~\pm$	0.7035 \pm	0.6625 \pm	0.3396 \pm	$0.6835~\pm$			
	0.0210	0.0000	0.0180	0.0797	0.2023	0.0246			
BosNot152 [60]	$0.7907~\pm$	$0.0000~\pm$	0.7142 \pm	$0.6577~\pm$	0.4057 \pm	$0.6997~\pm$			
itesivet152 [00]	0.0176	0.0000	0.0533	0.0506	0.0389	0.0304			
$\mathbf{P}_{os}\mathbf{N}_{ot}101$ [60]	$0.7983~\pm$	$0.0885~\pm$	$0.7281~\pm$	$0.6819~\pm$	$0.4075~\pm$	$0.7120 \pm$			
Resinet101 [60]	0.0269	0.1979	0.0345	0.0527	0.0378	0.0253			
ResNet152V2 [61]	0.7373 \pm	$0.0000~\pm$	$0.6891~\pm$	$0.5661~\pm$	$0.3617~\pm$	$0.6571~\pm$			
	0.0307	0.0000	0.0455	0.0522	0.0498	0.0294			
BosNot101V2 [61]	$0.7694~\pm$	$0.0000~\pm$	0.7016 \pm	$0.6655~\pm$	$0.4566~\pm$	$0.6909 \pm$			
Itesivetioi v 2 [01]	0.0164	0.0000	0.0312	0.0477	0.0397	0.0178			
X contion [62]	$0.7945~\pm$	$0.0000~\pm$	$0.6983~\pm$	0.6465 \pm	0.4224 \pm	$0.6971~\pm$			
Aception [02]	0.0298	0.0000	0.0189	0.0170	0.0493	0.0153			
Incontion Ros Not V2 [63]	$0.7734~\pm$	$0.0000~\pm$	0.7107 \pm	0.5938 \pm	0.3193 \pm	$0.6800 \pm$			
inceptionitesivet v 2 [05]	0.0302	0.0000	0.0263	0.0648	0.0596	0.0232			
MobileNetV2 [64]	0.7823 \pm	$0.1185~\pm$	0.6734 \pm	$0.5276~\pm$	$0.4400~\pm$	$0.6751~\pm$			
	0.0335	0.2650	0.0281	0.0434	0.0467	0.0194			
DenseNet121 [65]	$0.7755~\pm$	$0.0885~\pm$	$0.6850~\pm$	$0.6217~\pm$	$0.4769~\pm$	$0.6821~\pm$			
	0.0227	0.1979	0.0202	0.0587	0.0951	0.0133			
DenseNet169 [65]	$0.8049~\pm$	$0.0000~\pm$	$0.7058~\pm$	$0.6277~\pm$	$0.4535~\pm$	0.7014 \pm			
Denservetros [00]	0.0136	0.0000	0.0149	0.0729	0.0654	0.0125			
DenseNet201 [65]	$0.7850~\pm$	$0.0000~\pm$	$0.7086~\pm$	$0.6567~\pm$	$0.3660~\pm$	$0.6935~\pm$			
Denserver201 [00]	0.0259	0.0000	0.0176	0.0289	0.0513	0.0152			
Efficient Net B0 [66]	$0.7909~\pm$	$0.0000~\pm$	0.6610 \pm	$0.6673~\pm$	0.5573 \pm	$0.6918~\pm$			
Encienti (etb) [00]	0.0155	0.0000	0.0080	0.0658	0.0523	0.0030			

Table 13: Illustration of classification results of various pre-trained network on IDRiD Dataset using GMean evaluation metrics.

Table 14: Illustration of lesion detection results of various pre-trained networks on DDR Dataset using mAP, AR evaluation metrics.

Madala		Detection Boxes Precision							Detection Boxes Recall				
Models	mAP	\mathbf{mAP}	\mathbf{mAP}	\mathbf{mAP}	\mathbf{mAP}	mAP	AR@1	A B @10	AB@100	AR@100	AR@100	AR@100	
	mai	@0.5 IoU	@0.75 IoU	(small)	(medium)	(large)	Anei	Aneit	Aneito	(small)	(medium)	(large)	
EfficientDat D0 [67]	$0.0065~\pm$	$0.0189~\pm$	$0.0047~\pm$	0.0015 \pm	$0.0129~\pm$	0.0296 \pm	0.0070 \pm	0.0123 \pm	$0.0175~\pm$	$0.0037~\pm$	$0.0401 \pm$	0.0770 \pm	
Enicient Det-Do [07]	0.0012	0.0210	0.0011	0.0001	0.0121	0.0001	0.0018	0.0007	0.0017	0.0000	0.0014	0.0010	
Faster RCNN	0.0042 \pm	$0.0139~\pm$	$0.0001~\pm$	$0.0007~\pm$	$0.0098~\pm$	0.0441 \pm	$0.0083~\pm$	0.0151 \pm	0.0172 \pm	$0.0016~\pm$	0.0318 \pm	0.1199 \pm	
(ResNet-50) [68]	0.0023	0.0021	0.0001	0.0000	0.0000	0.0021	0.0001	0.0012	0.0102	0.0007	0.0024	0.0171	
SSD	0.0206 \pm	$0.0425~\pm$	$0.0312~\pm$	0.0035 \pm	$0.0482~\pm$	0.1038 \pm	$0.0180~\pm$	$0.0387~\pm$	$0.0501~\pm$	$0.0192~\pm$	$0.1201~\pm$	$0.1753~\pm$	
(MobileNetV1)[69]	0.0101	0.0201	0.0001	0.0009	0.0013	0.0012	0.0023	0.0102	0.0012	0.0103	0.0027	0.0102	
RetinaNet	0.0163 \pm	$0.0381~\pm$	$0.0152~\pm$	0.0012 \pm	0.0327 \pm	0.1319 \pm	$0.0199~\pm$	0.0402 \pm	0.0513 \pm	$0.0191~\pm$	$0.1298~{\pm}$	0.1645 \pm	
(ResNet50)[70]	0.0064	0.0016	0.0230	0.0102	0.0121	0.0014	0.0101	0.0078	0.0018	0.0012	0.0201	0.0019	
SSD	0.0192 \pm	$0.0283~\pm$	$0.0089~\pm$	0.0012 \pm	0.0284 \pm	0.0934 \pm	0.0132 \pm	0.0277 \pm	$0.0359~\pm$	0.0132 \pm	$0.0792~\pm$	0.1537 \pm	
(MobileNetV2)[71]	0.0209	0.0023	0.1023	0.0029	0.0023	0.0012	0.0012	0.0100	0.0080	0.0012	0.0010	0.0208	

Table 15: Illustration of fovea and optic disc detection results of various pre-trained networks on IDRiD Dataset using mAP, AR evaluation metrics.

	Ι	Detection B	oxes Precisio	Detection Boxes Recall				
Models	mAP	\mathbf{mAP}	\mathbf{mAP}	\mathbf{mAP}	AR@1	∆ R@10	AR@100	AR@100
	1117 11	@0.5 IoU	@0.75 IoU	(large)	meer	me	meioo	(large)
Efficient Dot D0 [67]	0.7221 \pm	$0.9810~\pm$	$0.9353~\pm$	0.7419 \pm	0.8161 \pm	0.8121 \pm	0.8243 \pm	$0.8254~\pm$
EncientDet-D0 [07]	0.0076	0.0010	0.0019	0.0121	0.0094	0.0191	0.0201	0.0092
Faster RCNN	0.7828 \pm	$0.9532~\pm$	0.9021 \pm	0.7292 \pm	0.8181 \pm	0.8914 \pm	$0.8500~\pm$	0.8510 \pm
(ResNet-50) [68]	0.02319	0.0167	0.0143	0.0132	0.0129	0.0132	0.0103	0.0190
\mathbf{SSD}	0.7822 \pm	$0.9790~\pm$	$0.9289~\pm$	0.7821 \pm	0.8241 \pm	0.8332 \pm	0.8212 \pm	0.8190 \pm
(MobileNetV1)[69]	0.0101	0.0230	0.0012	0.0023	0.0102	0.0121	0.0121	0.0120
${f RetinaNet}$	0.6912 \pm	$0.9690~\pm$	$0.8791~\pm$	0.7133 \pm	0.7123 \pm	0.7277 \pm	0.7278 \pm	0.7245 \pm
$({ m ResNet50})[70]$	0.0012	0.0102	0.0561	0.0131	0.0012	0.0121	0.0121	0.0234
\mathbf{SSD}	0.7612 \pm	$0.9756~\pm$	0.9144 \pm	0.7624 \pm	0.8012 \pm	0.8004 \pm	0.8190 \pm	0.8037 \pm
(MobileNetV2)[71]	0.0076	0.0129	0.0075	0.0034	0.0121	0.0103	0.0013	0.0612

	Segmentation - Dice score									
Models	Dicescore (Background)	$egin{array}{c} { m Dicescore} \ ({ m EX}) \end{array}$	Dicescore (HA)	Dicescore (MA)	$egin{array}{c} { m Dicescore} \ ({ m SE}) \end{array}$	mDicescore				
PSPNet [72]										
(w/ Focal Loss)	0.9910 ± 0.0001	0.0978 ± 0.0098	0.1319 ± 0.0138	0.0022 ± 0.0007	0.0324 ± 0.0028	0.0661 ± 0.0058				
DeepLab v2 [73]										
(w/ Focal Loss)	0.9901 ± 0.0002	0.0295 ± 0.0185	0.0575 ± 0.0403	0.0000 ± 0.0000	0.0078 ± 0.0123	0.0237 ± 0.0165				
DeepLab v3 [74]										
(w/ Focal Loss)	0.9912 ± 0.0002	0.1909 ± 0.0155	0.1569 ± 0.0255	0.0180 ± 0.0053	0.0323 ± 0.0053	0.0995 ± 0.0101				
PSPNet [72]										
(w/ Cross-entropy Loss)	0.9908 ± 0.0003	0.0993 ± 0.0052	0.1260 ± 0.0063	0.0015 ± 0.0004	0.0265 ± 0.0032	0.0634 ± 0.0029				
DeepLab v2 [73]										
(w/ Cross-entropy Loss)	0.9898 ± 0.0002	0.0307 ± 0.0194	0.0236 ± 0.0398	0.0000 ± 0.0000	0.0043 ± 0.0047	0.0146 ± 0.0134				
DeepLab v3 [74]										
(w/ Cross-entropy Loss)	0.9912 ± 0.0001	0.1907 ± 0.0116	0.1699 ± 0.0256	0.0142 ± 0.0087	0.0314 ± 0.0070	0.1016 ± 0.0063				

Table 16: Illustration of lesion segmentation results of various pre-trained networks on DDR Dataset using Dice Score evaluation metric.

Table 17: Illustration of lesion segmentation results of various pre-trained networks on DDR Dataset using Intersection over Union (IoU) evaluation metric.

Madala	Segmentation - Intersection over Union (IoU)									
Wodels	IoU	IoU	IoU	IoU	IoU	mIoII				
	(Background)	(\mathbf{EX})	(HA)	(MA)	(SE)	miou				
PSPNet [72]										
(w/ Focal Loss)	0.9954 ± 0.0001	0.1582 ± 0.0147	0.2096 ± 0.0197	0.0040 ± 0.0014	0.0458 ± 0.0032	0.1044 ± 0.0084				
DeepLab v2 $[73]$										
(w/ Focal Loss)	0.9950 ± 0.0001	0.0502 ± 0.0313	0.0947 ± 0.0654	0.0000 ± 0.0000	0.0117 ± 0.0176	0.0391 ± 0.0265				
DeepLab v3 [74]										
(w/ Focal Loss)	0.9955 ± 0.0001	0.2886 ± 0.0203	0.2439 ± 0.0352	0.0312 ± 0.0095	0.0457 ± 0.0075	0.1524 ± 0.0136				
PSPNet [72]										
(w/ Cross-entropy Loss)	0.9953 ± 0.0002	0.1599 ± 0.0077	0.2008 ± 0.0085	0.0029 ± 0.0008	0.0390 ± 0.0038	0.1007 ± 0.0040				
DeepLab v2 $[73]$										
(w/ Cross-entropy Loss)	0.9948 ± 0.0001	0.0530 ± 0.0333	0.0393 ± 0.0650	0.0000 ± 0.0000	0.0071 ± 0.0078	0.0249 ± 0.0223				
DeepLab v3 [74]										
(w/ Cross-entropy Loss)	0.9955 ± 0.0000	0.2887 ± 0.0149	0.2630 ± 0.0374	0.0250 ± 0.0147	0.0445 ± 0.0092	0.1553 ± 0.0082				

Table 18: Illustration of lesion, fovea and optic disc segmentation results of various pre-trained networks on IDRiD Dataset using Dice Score evaluation metric.

Madala	Segmentation - Dice score										
Models	Dicescore	Dicescore	Dicescore	Dicescore	Dicescore	Dicescore	mDiagaana				
	(Background)	(\mathbf{EX})	(HA)	(MA)	(SE)	(OD)	mDicescore				
PSPNet [72]											
(w/ Focal Loss)	0.9611 ± 0.0002	0.1921 ± 0.0160	0.3011 ± 0.0053	0.0038 ± 0.0018	0.1346 ± 0.0137	0.8857 ± 0.0073	0.3035 ± 0.0065				
DeepLab v2 $[73]$											
(w/ Focal Loss)	0.9502 ± 0.0018	0.0031 ± 0.0043	0.0287 ± 0.0406	0.0000 ± 0.0000	0.0000 ± 0.0000	0.8158 ± 0.0119	0.1695 ± 0.0049				
DeepLab v3 [74]											
(w/ Focal Loss)	0.9633 ± 0.0005	0.1213 ± 0.0351	0.3634 ± 0.0183	0.0298 ± 0.0270	0.0570 ± 0.0233	0.8986 ± 0.0044	0.2940 ± 0.0029				
PSPNet [72]											
(w/ Cross-entropy Loss)	0.9616 ± 0.0013	0.1876 ± 0.0029	0.3026 ± 0.0019	0.0012 ± 0.0020	0.1154 ± 0.0001	0.8914 ± 0.0041	0.2997 ± 0.0101				
DeepLab v2 [73]											
(w/ Cross-entropy Loss)	0.9509 ± 0.0011	0.0000 ± 0.0000	0.0898 ± 0.0418	0.0000 ± 0.0000	0.0000 ± 0.0000	0.8138 ± 0.0358	0.1807 ± 0.0140				
DeepLab v3 [74]											
(w/ Cross-entropy Loss)	0.9631 ± 0.0012	0.1859 ± 0.0657	0.3460 ± 0.0276	0.0033 ± 0.0022	0.0595 ± 0.0056	0.8990 ± 0.0055	0.2988 ± 0.0184				

Table 19:	Illustratio	on of lesion,	fovea and	optic disc	e segmentation	n results of	various	pre-trained
networks	on IDRiD	Dataset us	ing Interse	ction over	Union (IoU)	evaluation	metric.	

Madala	Segmentation - Intersection over Union (IoU)										
Models	IoU	IoU	IoU	IoU	IoU	IoU	mIoII				
	(Background)	(\mathbf{EX})	(HA)	(MA)	(SE)	(OD)	miou				
PSPNet [72]											
(w/ Focal Loss)	0.9800 ± 0.0001	0.3009 ± 0.0232	0.4457 ± 0.0069	0.0075 ± 0.0035	0.1919 ± 0.0152	0.9386 ± 0.0040	0.3769 ± 0.0074				
DeepLab v2 [73]											
(w/ Focal Loss)	0.9742 ± 0.0010	0.0057 ± 0.0080	0.0462 ± 0.0654	0.0000 ± 0.0000	0.0000 ± 0.0000	0.8957 ± 0.0058	0.1895 ± 0.0103				
DeepLab v3 [74]											
(w/ Focal Loss)	0.9811 ± 0.0002	0.1997 ± 0.0545	0.5140 ± 0.0200	0.0551 ± 0.0496	0.0830 ± 0.0381	0.9460 ± 0.0024	0.3596 ± 0.0056				
PSPNet [72]											
(w/ Cross-entropy Loss)	0.9803 ± 0.0027	0.2940 ± 0.0017	0.4461 ± 0.0120	0.0024 ± 0.0012	0.1696 ± 0.0105	0.9423 ± 0.0011	0.3709 ± 0.0206				
DeepLab v2 $[73]$											
(w/ Cross-entropy Loss)	0.9745 ± 0.0006	0.0000 ± 0.0000	0.1505 ± 0.0644	0.0000 ± 0.0000	0.0000 ± 0.0000	0.8938 ± 0.0238	0.2089 ± 0.0163				
DeepLab v3 [74]											
(w/ Cross-entropy Loss)	0.9810 ± 0.0007	0.2908 ± 0.0929	0.4970 ± 0.0289	0.0064 ± 0.0041	0.0839 ± 0.0124	0.9459 ± 0.0030	0.3648 ± 0.0240				



Figure 8: Results of segmentation model from PSPNet trained using focal loss for (i) DDR and (ii) IDRiD datasets.

7 Results and Discussion

We have conducted extensive experimentation using the three imbalanced publicly available diabetic retinopathy datasets of varied sizes for image classification, object detection and segmentation tasks. Tables 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13 illustrate the comparative analysis between various pretrained models, for the classification task for the three diabetic retinopathy datasets: - Kaggle DRD, DDR and IDRiD. From the analvsis it was found that DenseNet121 proves to be an effective model in case of all three datasets, in comparison to InceptionV3, MobileNetV2, Xception, ResNet50, EfficientNet-B0, VGG16, VGG19, ResNet152, ResNet101, ResNet152V2, ResNet101V2, DenseNet169, DenseNet201, ResNet50V2 and Inception-ResNetV2 pre-trained networks. After DenseNet121, the second-most effective pretrained network is Xception for all the three datasets with respect to various classification evaluation metrics: Cohen's Kappa (unweighted, linear, quadratic weighted), Accuracy, ROC-AUC (weighted and macro average) and F1-score, Index Balanced Accuracy (IBA) and Geometric Mean (GMean). Another observation found was that Class 1, Early stage diabetes is difficult to detect irrespective of whether that class falls under minority category or not, in case of all the three datasets. Third observation inculcated after conducting the experiments was that data augmentation used for training is suitable for longer training (400/200 epochs). Fourth observation found was that samples are equally distributed during training using rejection resampling that had a significant impact on the overall performance of the classification task. DenseNet and Xception architectures are both simple architectures amongst other prevalent architectures present in the study. DenseNet works well in evading the vanishing gradient problem and enabling feature reuse and has achieved state of the art (SOTA) results on the ImageNet, CIFAR and SVHN datasets [12] [18]. It is composed of dense blocks where every layer receives concatenated data from all the previous layers, thereby simplifying the residual network pattern. Moreover, it requires fewer parameters than the other convolutional networks. In contrast to ResNets, which also use skip connection. DenseNets concatenate the output and

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the input feature maps. Due to input concatenation, ease of access to learned maps for the corresponding layers is increased. Fine tuning on the pre-trained DenseNet helped in preserving the salient image features, along with learning the DR classification features.

Tables 14 and 15 depict the Detection Boxes Precision (mean average precision (mAP)) and Detection Boxes recall (average recall (AR)) metrics for object detection tasks for both DDR and IDRiD datasets. In this mAP metric, we also have an overlap criterion that lays down the minimum value of the intersection over union (IoU), which is used for correct detection. The value was taken as 0.5, 0.75I for IoU criterion. The results illustrates that in case of DDR dataset, SSD (MobileNetV1) is more efficient in lesion detection in comparison to EfficientDet-D0, Faster RCNN (ResNet-50), RetinaNet (ResNet50) and SSD (MobileNetV2) pre-trained networks. However, in the case of the IDRiD dataset for Fovea and Optic Disc Detection, EfficientDet-D0, Faster RCNN (ResNet-50), SSD (MobileNetV1) works well in comparison to other pre-trained networks (RetinaNet (ResNet50) and SSD (MobileNetV2)) while considering mAP without IoU evaluation metrics. But after taking consideration of mAP with IoU value we can see that EfficientDet-D0 is showing better results in comparison to other pre-trained networks in all aspects (when considering small and large objects). Tables 16, 17, 18, 19 show a comparative analysis between various stateof-the-art pre-trained networks for segmentation in case of DDR and IDRiD datasets, respectively, using Dice score and IoU (intersection over union) evaluation metrics between various pre-trained networks: DeepLabV2 and DeepLabV3 and PSPNet with cross-entropy loss and focal loss. After experimentation it was observed that PSPNet (with Focal Loss) is working best in comparison to other pretrained networks taken into consideration, results from which are shown in Figure 8.

8 Conclusion

In this paper, we have emphasized research on the medical diabetic retinopathy imbalanced datasets for classification, object detection and segmentation tasks in a unified framework. We have done extensive comparative analysis between various state-of-the-art pre-

trained deep models on varied size datasets. Throughout our study, we have focused on the transfer learning approaches for improving the performance of models. After conducting the analysis it was found that the DenseNet121 pre-trained model is the best suited for diabetic retinopathy image classification. The EfficientDet-D0 and SSD (MobileNetV1) are best suited for object detection on diabetic retinopathy datasets. In case of segmentation, PSPNet (with focal loss) performs best in comparison to other pre-trained networks. In future works, we will emphasize on the creation of architectures using this baseline study for lesion detection and segmentation, and explore ensemble learning with different pre-trained networks for more effective diagnosis of diabetic retinopathy as well as other challenging areas from the biomedical domain [31] [34] [35] [36] in order to be able to derive more generic approaches.

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