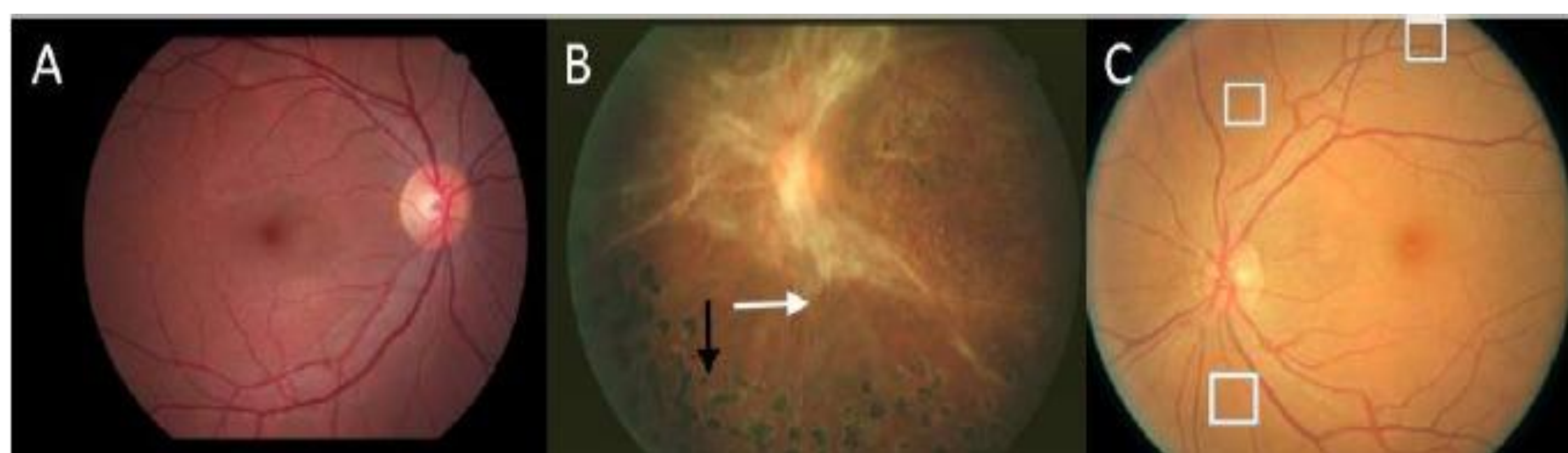


### Introduction

- Diabetic retinopathy (DR) is the damage to retinal blood vessels that eventually leads to blindness if not treated.
- Of an estimated 285 million people with diabetes mellitus worldwide, approximately one third have signs of DR. [1]
- While robust algorithms exist to diagnose late stage DR, early detection is still unsolved problem because it is difficult to detect micro-aneurysms (Figure 1). [1]



**Figure 1:** (A) Normal fundus photograph (B) Severe DR with white arrows pointing towards flame shaped hemorrhages (C) Early stage DR with white boxes showing micro-aneurysms

**Project goal: Develop machine learning algorithms for early DR detection**

### Previous Work

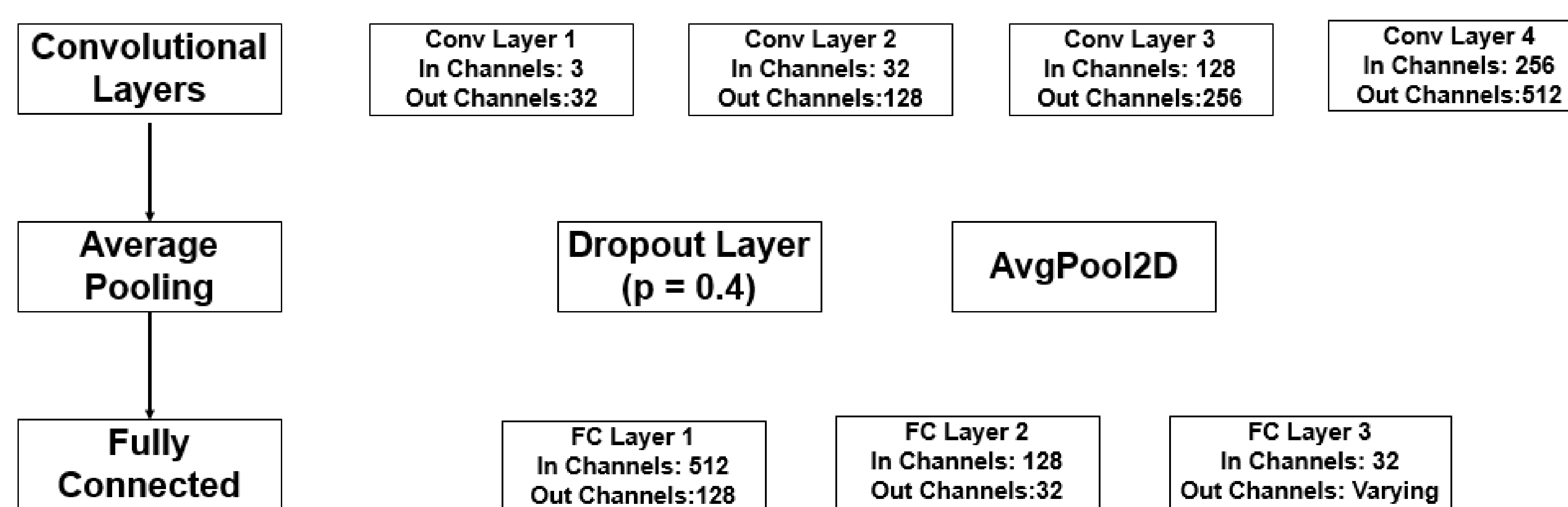
- Previous literature models have been able to perform binary classification of diabetic retinopathy at accuracies upwards of 90%. [2]
- Literature methods have not shown impressive results in the graded classification of DR, especially in terms with early diagnosis. [3]
- The first neural network that tried to differentiate between normal patches of retina from patches with micro-aneurysms achieved an accuracy of 74%. [4, 5]

### Dataset

- The APTOS 2019 blindness dataset was used to train all the models in this study. [6]
- A clinician has rated each image for the severity of diabetic retinopathy on a scale of 0 (no DR) to 4 (proliferative DR).
- The APTOS dataset was used to run three experiments:

- Binary classification:** labels changed to “0” if the image had no DR, or “1” otherwise.
- Early detection:** images with no DR were labelled “0,” “1” if original label was “1” or “2”, or “2” if original label was “3” or “4”.
- Severity classification:** original labels maintained.

### Model Architecture and Hyperparameters



**Figure 2:** Convolutional neural network architecture. Last fully connected layer had variable out channels depending on experiment

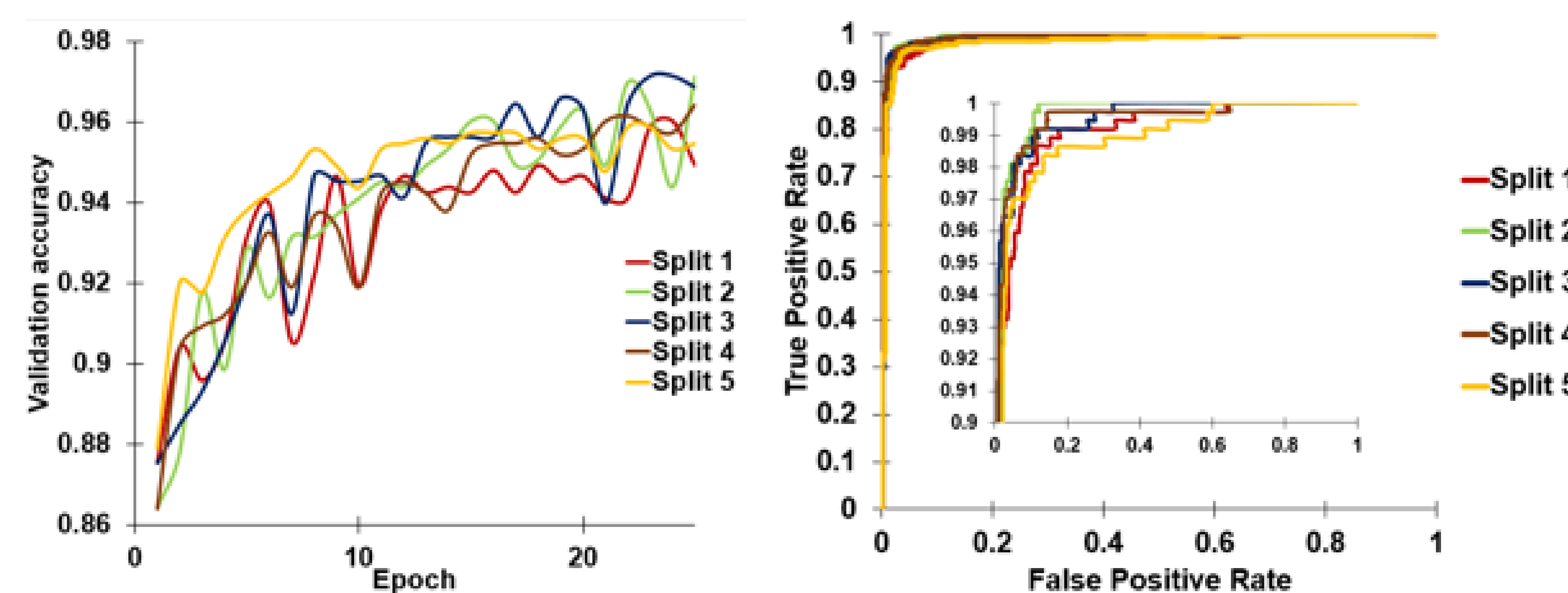
- Batch size = 8
- Epochs: 25 (binary classifier), 40 (early detector and severity classifier)
- Learning rate: 0.0001
- Optimizer: Adam optimizer
- Kernel size: 3x3 for all convolutional layers

### Training and validation protocol

- Five-fold stratified cross validation study performed for all experiments. Four splits for training and one split for validation.
- For binary classification, the true positive and false positive rates were calculated for each split. For the early detection and severity classifying experiments, confusion matrices were created for each split.

### Results

- The binary classifier model gave very strong validation accuracies (average 97%) and strong ROC curves (Figure 3).



**Figure 3:** (A) Validation curves and (B) ROC Curves for the five splits from cross-validation study of binary classification experiment.

- The early detector and severity classifier gave validation accuracies surpassing literature results (Table 1 and 2). [4, 5]

**Table 1:** Validation accuracy for early detector

Split	Validation Accuracy
1	0.86
2	0.87
3	0.83
4	0.84
5	0.86
Average	0.852

**Table 2:** Validation accuracy for severity classifier

Split	Validation Accuracy
1	0.79
2	0.78
3	0.83
4	0.80
5	0.77
Average	0.794

### Discussion

- The binary classifier’s sensitivity was calculated as 0.932 and the specificity as 0.989, which are comparable to literature models [2].
- The specificity being higher than sensitivity means that the model is better at predicting healthier cases than DR cases.
- Confusion matrices (Figure 4) show that model good at predicting early DR, but also confuses it with late DR.

		Model Prediction			Model Prediction						
		Split 2	0	1	2	Split 3	0	1	2	3	4
True Label	0	350	8	2	354	2	5	0	0		
	1	12	243	17	7	40	24	2	0		
	2	0	53	44	3	10	183	3	1		
	3	0	0	26	11	1	0	0	26	11	1
4	0	8	31	3	17	0	8	31	3	17	

**Figure 4:** Confusion matrices for (A) early detector (B) severity classifier

- This could happen because deeper in the model, the aperture is larger than the micro-aneurysms.

### Future Work

- Create residual connections so that learning earlier in the model can be connected with deeper layers.
- Move to a 3-1-1 cross validation study.
- To avoid overfitting, stop training the model when training and validation loss diverge.

### References

[1] Lee, R., et al. Eye and vision 2.1 (2015): 17.  
 [2] Lam, Carson, et al. AMIA 2018 (2018): 147.  
 [3] Gardner G., et al. British journal of Ophthal. 1996;80(11):940–944.  
 [4] Gargeya R, et al. Elsevier. 2017  
 [5] Gulshan V., et al. JAMA, 2016;316(22):2402–2410.  
 [6] <https://www.kaggle.com/c/aptos2019-blindness-detection/data>