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## The U-Net model application for retinal vessels segmentation using the machine learning library TensorFlow

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*Abstract*— In this article the implementation of neural network architecture based on a dense U-Net network is proposed. It is noted that retinal blood vessels are the basis for clinical diagnosis of some diseases. A review of the convolutional networks use for classification tasks and generalizion retinal vessel segmentation algorithms is performed. The general process of the neural network is presented. The differences between the real and the obtained results were evaluated. Evaluation of the neural network is carried out on several parameters. The figure with the recognized blood vessels as a result of the model is presented.

## Keywords — machine learning, neural network, machine learning library, retinal vessels segmentation.

#### I. INTRODUCTION

Retinal blood vessels are the basis for the clinical diagnosis of some diseases. Achieving automatic retinal vessel segmentation in fundus imaging is an important and challenging task. This paper proposes neural network architecture based on a dense U-Net network.

Convolutional networks have existed for a long time [27], their success was limited due to the size of the available training kits and the size of the considered networks. Kryzhevsky's breakthrough [1] was due to the controlled learning of a large network with 8 layers and millions of parameters of the ImageNet data set with 1 million training images. Since then, even larger and deeper networks have been taught [14].

The typical use of convolutional networks is used for classification tasks, where the source image is one class label. However, in many visual tasks, especially in biomedical image processing, the desired result should include localization, ie the class label should be assigned to each pixel. Moreover, thousands of educational images are usually inaccessible for Roksolana Milian

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biomedical tasks. Thus, Ciresan, Gambardella, Giusti and Schmidhuber [6] taught a network in a sliding window installation to predict the class label of each pixel by providing a local patch around that pixel. First, this network can be localized. Second, the training data in terms of patches is much larger than the number of training images.

#### II. MATERIAL AND METHODS

It is worth considering the network architecture shown in Figure 1, which is illustrated in [17]. It consists of a contracting path (left side) and an expansive path (right side). The laying path corresponds to the typical architecture of the convolutional network. It consists of repeated application of two 3x3 convolutions (unchanged convolutions), each of which is a rectified linear unit (ReLU) and a combination operation of max. 2x2 with step 2 to reduce the sample. At each sampling step, the number of functional channels is doubled. Each step in the expansive path consists of a resampling of the object map, followed by a 2x2 convolution ("convolution from above"), which halves the number of object channels, concatenation with a properly truncated object map from the contracting path and two 3x3 convolutions, each of which is accompanied by ReLU. Trimming is necessary due to the loss of border pixels in each gyrus. On the final layer, the 1x1 convolution is used to map each 64-component feature vector to the desired number of classes. In total, the network has 23 convolutional layers.



Figure 1. U-Net architecture (example for 32x32 pixels with the lowest resolution). Each blue box corresponds to a multi-channel feature map. The number of channels is indicated at the top of the window. Size x-y is located in the lower left edge of the box. White boxes represent copied object maps. Arrows indicate various operations.

Retinal blood vessels contain significant information regarding human health. Observation of the morphological structure of retinal vessels in fundus images is used not only for screening retinal vessel diseases, but also for auxiliary diagnosis of other diseases such as stroke [5], hypertension [21], diabetes-induced retinopathy [10] and glaucoma [9]. Images of the fundus have the following characteristics: low contrast between the vessel and the background, serious interventions in the affected area and a complex vessel structure, which creates many problems to achieve segmentation of retinal vessels [3]. Currently, the main method of retinal vessel segmentation is manual annotation by professional physicians.

However, with the constant development of medical imaging technology, more and more images of the fundus require segmentation. Manual annotation of retinal vessels requires a lot of time and energy from physicians, and different physicians always adopt different standards for retinal vessel segmentation. Therefore, many algorithms for automatic retinal vessel segmentation have been proposed for Computer-Aided Diagnosis (CAD). Achieving automatic segmentation of retinal vessels can not only reduce the workload of physicians, but also avoid the subjective influence of different physicians on the results of segmentation.

According to the generalized retinal vessel segmentation algorithms [11], [23], it can be divided into two categories: supervised and unsupervised algorithms. In general, unsupervised learning algorithms do not require manual annotation data and generally use some pre-established rules to extract vessel features and achieve segmentation, such as matching algorithms based on filters [16], deforming algorithms based on models [18], and tracking-based algorithms [24]. However, fixed rules of segmentation often cannot correspond to a variety of morphological distribution of vessels.

The main idea of supervised learning algorithms is to teach a segmentation model using fundus images with segmentation annotations, which allow models to automatically extract vessel features to achieve vessel segmentation, for example, algorithms based on the Bayesian model [8], support vector machine based on algorithms [7] and deep learning algorithms [19], [4], [22], [20]. However, supervised learning algorithms require huge data with manual marking, which is difficult to obtain.

In recent years, algorithms based on deep learning have continued to be developed and worked well in the field of retinal vessel segmentation, which have gradually become the main algorithm. Long, Shelhamer and Darrell [12], proposed a Fully Collapsed Neural (FCN) network, various retinal vessel segmentation algorithms based on the FCN structure are constantly coming out. Oliveira, Pereira and Silva [2] presented a retinal vessel segmentation algorithm that combines multiscale wavelet transform and multiscale FCN. Lu, Xu, Chen and Luo [13] used a Coarse-to-Fine Fully Convolutional Neural (CF-FCN) network to extract blood vessels in fundus images. The CF-FCN aimed to use the original data information and compensate for the output of the neural network by the spatial relationship between the pixels in the fundus image. However, the results of segmentation of the FCN-based structure did not have adequate results in detail. Therefore, it usually needed some processing methods, such as Conditional Random Field (CRF) and Markov Random Field (MRF).

Among the FCN-based segmentation algorithms is the Unet structure proposed by Ronneberger, Fischer and Brox [17], changed the method of connecting the transition to the map of functions in the FCN with the addition of concatenate, which is widely used in the field of segmentation of biomedical images. In addition, a large number of advanced U-Net-based algorithms have emerged for retinal vessel segmentation. For example, Xiao, Lian and Luo [26] propose a model based on a U-network with a weighted shutter of attention for segmentation of small retinal vessels. Gao, Cai and Qiu [25] combine Gaussian and U-Net filtering to achieve retinal vessel segmentation. Alom, Hasan and Yakopcic [15] offer two advanced models based on U-Net. The first is Recurrent Convolutional Neural Network (RCNN), and the second is the Recurrent Residual Convolutional Neural Network (RRCNN), both of which have been used successfully for retinal vessel segmentation.

#### III. RESULTS AND DISCUSSION

The implementation of a simple U-net model for segmentation of retinal blood vessels is presented below. The model is based on the Keras and Tensorflow libraries. The method was tested on a public DRIVE dataset. For the DRIVE data set, which consists of 40 images, we used 20 images to train the model and 20 images to test the model. Finally, complete content and organizational editing before formatting.

Contrast-constrained adaptive histogram alignment (CLAHE) is used to normalize the images. The image is divided into small blocks called "tiles" (the default tile size is 8x8 in OpenCV). Then each of these blocks is aligned. Therefore, in a small area, the histogram is limited to a small area (if there is no noise). If there is noise, it will be amplified. To avoid this, contrast restriction is applied. If the specified

contrast limit is exceeded (default 40 in OpenCV), these pixels are truncated and evenly distributed in other containers before applying histogram alignment. After alignment, bilinear interpolation is used to remove artifacts within the tile. Gamma correction using the lookup table is applied.

We used the Adam optimizer with *initial\_learning\_rate* = 0.0003, *first\_decay\_steps* = 12000, *t\_mul* = 1000, *m\_mul* = 0.5, *alpha* = 1e-5.

For method evaluation we use Precision, Recall, Confusion matrix, and the area under the Receiver Operating Characteristic Curve (AUC-ROC) (Figures 2 and 3).

Recall (also called True Positive Rate or Sensitivity) is defined as the ratio of truly classified vessel pixels. Precision (also called True Negative Rate or Specificity) is the ratio of truly classified non-vessel pixels.

$$Precision = \frac{TP}{TP + FP}$$
$$Recall = \frac{TP}{TP + FN}$$

Where TP is the number of pixels that are classified as vessels in the image, which are correctly classified. FP is the number of pixels classified as vessels which are incorrectly classified. FN is the number of pixels classified as non-vessels which are incorrectly classified.



Figure 2. Results of metrics calculation Dice Score, Precision, Recall, AUC



Figure 3. Confussion matrix for model evaluation

Another used metric is the AUC (Area under the curve) or ROC (Receiver Operating Characteristic Curve) (a graphical plot from (0;0) to (1;1) of the false positive rate (x-axis) versus the true positive rate (y-axis)). Figure. 4 shows the ROC curve.



Figure 4. ROC curve

To classify the images, we exported the images in the form of 25 tiles of the same size for deep learning.



Figure 5. Exported images

The forming tiles process for the deep learning application is presented in Figure 6.

Code + Text		RAM Disk Editing
plt.ins else;	ave(train_patch_dir+image_name+"-"+str(i)+"-groundtruth.jpg",(patch_groundtr	uth_list[i]/225.0).astype(np.uint8),cmap = plt.cm.gray)
plt.ims	ave(train_patch_dir+image_name+"_"+str(i)+"_val_img.jpg",patch_image_list[i]	)
plt.ims	<pre>ave(train_patch_dir+image_name+"_"+str(i)+"_val_groundtruth.jpg",(patch_grou</pre>	ndtruth_list[i]/225.0).astype(np.uint8),cmap = plt.cm.gray)
# delete origin	al patch images	
if not os.path.	exists(train_patch_dir):	
os.mkdir(trai	.n_patch_dir)	
else:	(teals eatch die)	
os.mkdir(trai	in_patch_dir)	
<pre>if not os.path.</pre>	exists(test_save_dir): save_dir)	
for i in todm(r	n images ange(len(train_image_path_list)),desc="Generate the training patches: "): rains (area ath light) atch our atch size to (see Two (beschick)) = co	t show-Town to visualize the sample spaces, which is much slower th
ruskesbaccu(c	rest_snege_peci_sist[i],pecci_non,pecci_siste,crestaing=free,snow=reste) = se	c show-rive to visualize the sample process, which is much slower to
for i in todm(r	<pre>ange(len(val_image_path_list)),desc="Generate the val patches: "): (a) image path_list[i].patch_pum_patch_size_training=False.show=False) # set</pre>	showTrue to visualize the sample process, which is much slower the
Generate the tr Generate the va	aining patches: 100%  14/14 [04:47<00:00, 20.51s/it] 1 patches: 50%  3/6 [00:59<00:59, 19.78s/it]	

Figure 6. The forming training tiles process in the Google Colab environment

Binary cross entropy was used as a loss function to determine the error between actual and obtained results and to minimize it.

Iteration Loss function		Dice coefficient	Iteration	Loss function	Dice coefficient
1	0.5218	0.4782	38	0.1181	0.8819
2	0.4937	0.5063	39	0.1152	0.8848
3	0.4742	0.5258	40	0.1131	0.8869
4	0.4552	0.5448			
5	0.4355	0.5645	124	0.0228	0.9772
6	0.4161	0.5839	125	0.0226	0.9774
7	0.3970	0.6030	126	0.0224	0.9776
8	0.3783	0.6217	127	0.0219	0.9781
9	0.3603	0.6397	128	0.0215	0.9785
			129	0.0214	0.9786
31	0.1410	0.8590	130	0.0214	0.9786
32	0.1368	0.8632	131	0.0207	0.9793
33	0.1334	0.8666	132	0.0206	0.9794
34	0.1293	0.8707	133	0.0205	0.9795
35	0.1262	0.8738	134	0.0201	0.9799
36	0.1230	0.8770	135	0.0200	0.9800
37	0.1201	0.8799	136	0.0198	0.9802

TABLE I. MINIMIZING THE LOSS FUNCTION VALUE IN THE LEARNING PROCESS

The calculating process of the loss function and the Dice coefficient took place directly in the software environment.

+ Cod	le + Text		
	epoch 36,	batch	93, train_loss:0.1230, train_dice:0.8770, val_loss:0.2505, val_dice:0.7495
(U	epoch 37,	batch	218, loss:0.1201, dice:0.8799
	epoch 38,	batch	93, train_loss:0.1181, train_dice:0.8819, val_loss:0.2486, val_dice:0.7514
	epoch 39,	batch	218, loss:0.1152, dice:0.8848
	epoch 40,	batch	93, train_loss:0.1131, train_dice:0.8869, val_loss:0.2474, val_dice:0.7526
	epoch 41,	batch	218, loss:0.1115, dice:0.8885
	epoch 42,	batch	93, train_loss:0.1096, train_dice:0.8904, val_loss:0.2432, val_dice:0.7568
	epoch 43,	batch	218, loss:0.1083, dice:0.8917
	epoch 44,	batch	93, train_loss:0.1070, train_dice:0.8930, val_loss:0.2451, val_dice:0.7549
	epoch 45,	batch	218, loss:0.1060, dice:0.8940
	epoch 46,	batch	93, train_loss:0.1052, train_dice:0.8948, val_loss:0.2415, val_dice:0.7585
	epoch 47,	batch	218, 1055:0.1042, d1ce:0.8958
	epoch 48,	batch	95, train_Ioss:0.1037, train_0120005, val_Ioss:0.2435, val_dice:0.7565
	epoch 49,	batch	218, 105510-1052, 0102-0.8908
	epoch 50,	batch	93, train_Ioss:0.1029, train_01c0:0.89/1, Vai_10ss:0.2422, Vai_01c0:0.7578
	epoch 51,	batch	210, 1055:0.1025, ULCE:0.0975 02 train loss:0.1025 train dica:0.9075 val loss:0.2412 val dica:0.7588
	epoch 52,	hatch	18 loc: 0 1023 dice 0 8077
	epoch 54	batch	03. train loss:0.102. train dice:0.8078, val loss:0.2414, val dice:0.7586
	epoch 55.	batch	218. los:0.1039. dice:0.8961
	epoch 56.	batch	93. train loss:0.1143. train dice:0.8857. val loss:0.2430. val dice:0.7570
	epoch 57.	batch	218, loss:0.1103, dice:0.8897
	epoch 58,	batch	93, train loss:0.1056, train dice:0.8944, val loss:0.2365, val dice:0.7635
	epoch 59,	batch	218, loss:0.1015, dice:0.8985
	epoch 60,	batch	93, train_loss:0.0983, train_dice:0.9017, val_loss:0.2283, val_dice:0.7717
	epoch 61,	batch	218, loss:0.0953, dice:0.9047
	epoch 62,	batch	93, train_loss:0.0916, train_dice:0.9084, val_loss:0.2229, val_dice:0.7771
	epoch 63,	batch	218, loss:0.0888, dice:0.9112
	epoch 64,	batch	93, train_loss:0.0860, train_dice:0.9140, val_loss:0.2139, val_dice:0.7861
	epoch 65,	batch	218, loss:0.0834, dice:0.9166
	epoch 66,	batch	93, train_loss:0.0811, train_dice:0.9189, val_loss:0.2211, val_dice:0.7789
	epoch 67,	Datch	101, 1055:0.0/90, d1Ce:0.9210
			Executing (2b 9m 15c) Cell 5 train step() 5 cell() 5 cell()

Figure 7. The calculating process of the loss function and the Dice coefficient in the Google Colab environment

The result of the neural network is well traced comparing, for example, the initial image of the fundus in the eleventh image (Figure 8) and the results of the U-Net model that presented in Figure 9 with recognized blood vessels for the eleventh image.



Figure 8. The initial image of the fundus in the eleventh image



Figure 9. The U-net model results for the eleventh image

#### IV. CONCLUSION

Automatic the retinal vessels segmentation in the fundus images is an important task, as images with detected blood vessels can diagnose some diseases. This paper proposes the implementation of a neural network architecture based on a dense U-Net network in Google Colab on a public DRIVE data set using the Tensorflow machine learning library for the retinal vessels segmentation. The general process of the neural network is presented. Binary cross-entropy is used as a loss function for calculation the error between the actual and obtained results and its minimization. The results are presented in the table. Precision, Recall, Confusion matrix, and the area under the Receiver Operating Characteristic Curve (AUC-ROC) were used evaluating the method. The results of the model, namely the recognized blood vessels are presented.

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