# Choroid automatic segmentation and thickness quantification on swept-source optical coherence tomography images of highly myopic patients

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**Background:** Thinning of the choroid has been linked with various ocular diseases, including high myopia (HM), which can lead to visual impairment. Although various artificial intelligence (AI) algorithms have been developed to quantify choroidal thickness (ChT), few patients with HM were included in their development. The choroid in patients with HM tends to be thinner than that of normal patients, making it harder to segment. Therefore, in this study, we aimed to develop and implement a novel deep learning algorithm based on a group-wise context selection network (GCS-Net) to automatically segment the choroid and quantify its thickness on swept-source optical coherence tomography (SS-OCT) images of HM patients.

**Methods:** A total of 720 SS-OCT images were obtained from 40 HM eyes and 20 non-HM eyes and were used to develop a GCS-Net to segment the choroid. The intersection-over-union (IoU), Dice similarity coefficient (DSC), sensitivity, and specificity were used to assess the performance in relation to manually segmented ground truth. The independent test dataset included 3,192 images from 266 HM eyes. The ChT in the test dataset was measured manually and automatically at 9 different regions within the choroid. The average difference in the ChT between the 2 methods was calculated. The intraclass correlation coefficient (ICC) was calculated to evaluate the agreement between the 2 measurements.

**Results:** Our method reached an IoU, DSC, sensitivity, and specificity of 87.89%, 93.40%, 92.42%, and 99.82% in HM, respectively. The average difference in the ChT between the 2 measurements was  $5.54\pm4.57 \mu m$ . The ICC was above 0.90 (P<0.001) for all regions of the choroid, indicating a very high level of agreement.

**Conclusions:** The GCS-Net proposed in our study provides a reliable and fast tool to quantify ChT in HM patients and could potentially be used as a tool for monitoring ChT in ocular diseases related to the choroid.

**Keywords:** Deep learning algorithm; choroidal thickness (ChT); high myopia (HM); swept-source optical coherence tomography (SS-OCT)

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#### Introduction

The prevalence of high myopia (HM) continues to increase (1-3). It is estimated that there will be 938 million people living with HM by 2050, accounting for 9.8% of the world's population (1). Pathological myopia, which is a major cause of visual impairment and blindness worldwide, often occurs in eyes with HM (4-6).

The choroid is a vascular structure located between the retina and sclera, which supplies nutrients to the outer retina and serves critical physiological functions such as regulation of intraocular pressure and light absorption (7,8). Several studies on non-HM patients have shown that age can lead to thinning of the choroid (9-12); however, choroidal thinning is also related to many diseases, including myopia and age-related macular degeneration (AMD) (13-16).

The choroidal thickness (ChT) decreases with increasing levels of myopia (8,17). The ChT in HM patients can be about half to a quarter thinner when compared with that of patients with normal vision and correlates closely with the refractive error, axial length (AL), and posterior staphyloma height (8,18-20). The accurate quantification of the ChT is essential in the study of ocular diseases associated with the choroid. However, the manual measurement of ChT is time-consuming, limiting its use as a potential indicator for monitoring disease progression in HM patients.

The use of artificial intelligence (AI), especially deep learning algorithms, in ophthalmology has increased considerably in the past few years for the diagnosis, classification, prediction, and prognosis of ocular diseases (21-23). Optical coherence tomography (OCT) is often used to acquire high-resolution ocular images as part of a clinical examination and has an important role in the development of AI algorithms. As evidenced by previous studies, the use of AI models based on OCT images results in highly accurate detection of various pathological conditions, including retinal lesions, AMD, macular edema, retinoschisis, retinal detachment, and macular hole (24-26).

Although previous studies have applied traditional algorithms to segment the choroid, they have had limitations such as being applicable only to segment the normal choroid or requiring high-quality images (27-29). Regarding the thin choroid in HM, deformation of the retina caused by elongation of the AL, posterior staphyloma, or the possible existence of pathological atrophy regions makes the detection of the retinal and choroidal structure inaccurate and difficult.

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Therefore, in this study, we aimed to develop a novel deep learning algorithm based on a group-wise context selection network (GCS-Net) to automatically segment the choroid region and quantify the ChT on swept-source optical coherence tomography (SS-OCT) images of HM patients. The accuracy of the algorithm was validated against a manually segmented choroid as ground truth. We present the following article in accordance with the TRIPOD reporting checklist (available at https://atm. amegroups.com/article/view/10.21037/atm-21-6736/rc).

#### **Methods**

#### Participants

We conducted a training, validation, and external testing study on an AI model using SS-OCT images. The data source of this study was a section of the Shanghai Eye Study for Older People, which was a population-based, crosssectional study including individuals aged 50 years and older in Shanghai, China, between 2016 and 2018. Patients were excluded from the study if they had a history of eye surgery (except cataract surgery), corneal opacity, severe cataract, glaucoma, systemic diseases with ocular involvement, and fundus lesions unrelated to myopia (e.g., AMD, diabetic retinopathy, and optic neuropathy).

The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013). The study was approved by the Ethics Committee of the Shanghai General Hospital, Shanghai Jiao Tong University School of Medicine (No. 2015KY153), and informed consent was provided by all individual participants.

#### **Ophthalmic examinations**

All patients enrolled in this study underwent comprehensive clinical interviews and ophthalmic examinations, including assessments of the refractive error using an autorefractor instrument (model KR-8900; Topcon, Tokyo, Japan), measurement of intraocular pressure (Full Auto Tonometer TX-F; Topcon), slit-lamp biomicroscopy, color fundus examination, and AL measurement using IOL Master (Carl Zeiss Meditec, Jena, Germany). Subjective refraction was performed by an experienced optometrist for all participants. The spherical equivalent (SE) was obtained as follows: SE = sphere power + (cylinder power/2). Eyes with an ocular AL greater or equal to 26 mm were defined as HM (13). The Topcon Atlantis DRI-1 SS-OCT scanner

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was used to acquire 12-line radial B-scans centered on the fovea. The SS-OCT parameters were as follows: wavelength of the light source =1,050 nm, scan rate =100,000 A-scans per second, depth resolution =8  $\mu$ m, and lateral resolution =10  $\mu$ m. The image size of each B-scan was 1,024 (width of B-scans) × 992 (depth of B-scans) pixels, which corresponds to a total area of 9×2.6 mm<sup>2</sup>.

The ChT, defined as the vertical distance between Bruch's membrane and the choroidal-scleral interface, was measured using a specialized grid focused on the macula as described in the Early Treatment of Diabetic Retinopathy Study (ETDRS). The diameters for the foveal circle, parafoveal circle, and perifoveal circle of the ETDRS grid were set at 1, 3, and 6 mm, respectively. The parafoveal and perifoveal regions were further divided into temporal, superior, nasal, and inferior quadrants. The average ChT was measured at 9 different regions, including the inner and outer parts of each of the 4 quadrants and the center of the choroid.

#### Datasets

The development dataset included a total of 720 SS-OCT B-scans obtained from 60 eyes, comprising 40 HM eyes and 20 non-HM eyes. Each B-scan was considered an independent image and exported as a jpeg to use in algorithm development. All images were labeled by 1 welltrained grader (ML) and supervised by a retina specialist (YF), who checked the manual segmentation at least once. Subsequently, these images were subdivided into a training dataset (80% of the images) for model development and a validation dataset (20% of the images) for validating the model. To facilitate calculation of the AI model, we adjusted the OCT image size to 512×256 pixels. We then applied online augmentation of left-right flip, which simulated the symmetry of right and left eyes, to increase the size of the training set and improve the generalization ability and robustness of the model. The AI system was verified using an independent real test clinical dataset consisting of 3,192 images obtained from 266 HM eyes according to the same criteria listed above. None of the latter images had been used previously in the training and validation datasets.

#### GCS-Net

A GCS-Net can effectively select multiscale context information to achieve accurate segmentation of choroid regions with different thicknesses (30). Extraction of multiscale features corresponding to different perception fields also gives the model strong ability in distinguishing choroid and other retinal structures. The novelty in designing GCS-Net lies in 2 modules, i.e., the group-wise channel dilation (GCD) module, the group-wise spatial dilation (GSD) module, and a boundary optimization subnetwork (BON). Both GCD and GSD adopt the selfattention mechanism, where feature maps are recalibrated with adaptive weights. In this way, features with higher discriminating ability are emphasized. The GCD module can select multiscale information under the guidance of channel information, while the GSD module can use spatial information to guide the fusion of multiscale context information. The BON uses deep supervision to solve the problem of choroidal boundary blur.

In the GCD and GSD modules, the input feature maps are divided into groups and each group goes through dilated convolutions with different rates, thus obtaining multiscale features. The GCD module downsamples the feature matrix to obtain the channel weights and multiplies each convoluted group with the corresponding weight. Finally, the weighted multiscale features are added to the input features as the output. The difference between the GSD module and the GCD module is that the weights obtained by the GSD module are spatial weights.

The GCS-Net is a U-shaped network with 4-layer encoder and decoder. The GCD module is embedded between the encoder and decoder, where the output of each layer of the encoder is connected to the decoder through the GCD module. The GSD module is added after each deconvolution layer of the decoder. Except for the bottom layer, the sum of the outputs of the GCD and the GSD module forms the input of the next deconvolution layer. The boundary optimization network adopts a deep supervision strategy. That is, the output of each GSD module is up-sampled and convoluted, and based on the result, an edge loss is calculated compared with the ground truth edge map. The 4 edge losses obtained from each layer are added to the region loss to obtain the total loss of the GCS-Net. The overview of GCS-Net and the illustration of GCD and/or GSD module are shown in Figure 1.

#### **Evaluation** metrics

In the validation dataset, the performance of the AI model in segmenting the choroid was quantitatively assessed using 4 evaluation metrics: the intersection-over-union (IoU), the Dice similarity coefficient (DSC), sensitivity, and specificity. Page 4 of 11



Figure 1 The proposed deep learning-based algorithm framework. (A) Overview of the GCS-Net. (B) The illustration of GCD and GSD module. GCS-Net, group-wise context selection network; GCD, group-wise channel dilation; GSD, group-wise spatial dilation.

The formulas used to calculate IoU and DSC are as follows:

$$IoU = \frac{TP}{TP + FP + FN}$$
[1]

$$DSC = \frac{2 \times TP}{2 \times TP + FP + FN}$$
[2]

The TP, FP, and FN represent the number of true positive, false positive, and false negative predictions. For image segmentation, TP represents the number of pixels predicted as foreground by automatic segmentation and labeled as foreground in the ground truth, FP represents the number of pixels predicted as foreground but labeled as background in the ground truth, and FN represents the number of pixels predicted as background but labeled as foreground in the ground truth (*Figure 2*). The IoU and DSC are used to measure the ratio of overlap, which reflects the similarity of the 2 samples. Sensitivity reflects the proportion of correctly segmented foreground parts in the ground truth. Specificity reflects the proportion of correctly segmented background parts in the ground truth. We used Python (v3.7; Python Software Foundation, Wilmington, DE, USA) and the Pytorch (v1.7) deep learning framework of Pycharm (2019, JetBrains, Prague, Czech Republic) software to perform model experiments and to calculate



**Figure 2** The definition of TP, FP, and FN for image segmentation. (A1 and B1) Choroid boundaries delineated by ground truth (green outline) and the prediction of GCS-Net (red outline). (A2 and B2) Mask of ground truth and GCS-Net. The foreground is the white part of the mask output and the background is the black part. (C) Diagram of overlapping mask. TP (blue area) represents the number of pixels predicted as foreground by automatic segmentation and labeled as foreground in the ground truth, FP (pink area) represents the number of pixels predicted as foreground but labeled as background in the ground truth, and FN (yellow area) represents the number of pixels predicted as background but labeled as foreground in the ground truth, FP, false positive; FN, false negative; GCS-Net, group-wise context selection network.

these evaluation metrics.

# Comparison with clinical ophthalmologist and statistical analysis

After choroid automatic segmentation, the ChT was subsequently calculated by converting the pixel counts into µm, and the results of automatic segmentation were compared with manual segmentation calculated by the built-in Topcon software with its caliper in the ETDRS grid. The lateral magnification was adjusted using AL by Littmann's formula (31). In the test dataset, the performance of the algorithm to measure the ChT was assessed by calculating the difference between the automated and manual measurements for all 9 measured regions. The intraclass correlation coefficient (ICC) was used to analyze the similarity between the 2 methods. The patients' characteristics were shown as means ± standard deviation for continuous data and as counts or proportions for categorical data. Data distribution was examined using the Kolmogorov-Smirnov test. The Mann-Whitney U and chi-square tests were used to assess whether there were

statistically significant differences between the training and validation datasets. A generalized estimation equation was used to account for internal correlation for binocular data. All statistical analyses were performed using the software SPSS 26.0 (IBM Corp., Armonk, NY, USA), and a 2-tailed P value below 0.05 was deemed statistically significant.

#### Results

The generalized estimation equation models showed no significant differences in ocular parameters between the 2 eyes; thus, there was no need to adjust for associations between them. A total of 326 eyes of 215 participants were included in this study after comprehensive ophthalmic examinations, comprising 60 eyes in the development dataset and 266 eyes in the test dataset. The development image dataset was subsequently divided into a training set (576 scans) and a validation set (144 scans). *Table 1* shows the demographic and clinical characteristics of the training and validation sets, and no significant difference was found in these parameters (P>0.05).

The performance of GCS-Net was evaluated in both

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Variables	Training dataset		Validatio	Durshus	
variables –	High myopia	Non-high myopia	High myopia	Non-high myopia	P value
No. of eyes	32	16	8	4	
No. of images	384	192	96	48	
Age, y	70.03±5.96	65.00±6.79	67.75±6.41	68.50±3.79	0.956
Gender, male/female	17/15	7/9	4/4	1/3	0.605
SE, diopter	-11.00±3.93	-0.57±2.01	-10.94±3.43	-1.69±3.26	0.919
IOP, mmHg	13.66±2.87	14.33±2.33	14.19±2.72	13.63±3.35	0.892
AL, mm	28.13±0.86	23.38±0.48	27.50±1.25	23.12±1.57	0.405

Table 1 Demographic an	d ocular characteristics	of training and	l validation datasets
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SE, spherical equivalent; IOP, intraocular pressure; AL, axial length.

Table 2 The performance of GCS-Net in high myopia and non-high myopia eyes

Variables	IoU (%)	DSC (%)	Sensitivity (%)	Specificity (%)
Total	87.89±6.93	93.40±4.10	92.81±6.34	99.66±0.52
High myopia	87.89±7.10	93.40±4.22	92.42±6.56	99.82±0.11
Non-high myopia	87.88±6.57	93.42±3.84	93.59±5.79	99.33±0.80

GCS-Net, group-wise context selection network; IoU, intersection-over-union; DSC, Dice similarity coefficient.

HM and non-HM eyes (*Table 2*). In HM eyes, the IoU, DSC, sensitivity, and specificity were 87.89%, 93.40%, 92.42%, and 99.82%, respectively, thus supporting the ability of GCS-Net in choroidal segmentation. The visual assessment also showed a good agreement between the automated segmented choroid and the manual ground truth (*Figure 3*).

In the test dataset, the algorithm was used for automatic ChT calculation and compared with manual segmentation to further verify the accuracy of the calculation. In this HM test dataset, the average SE was  $-10.52\pm3.50$  D and the average AL was 27.76 $\pm$ 0.99 mm. The average center foveal ChT was 69.6 $\pm$ 39.1 µm, and a decreasing trend was found horizontally from the temporal to nasal region (*Figure 4*). The average difference between the 2 methods was 5.54 $\pm$ 4.57 µm with a maximum error of 24.07 µm (*Table 3*). The ICC was above 0.90 (P<0.001) for all choroid regions. The lowest ICC (0.944, P<0.001) was found in the outer nasal region of the ETDRS grid. With the exception of this region, the ICCs in all other sectors were above 0.97 (P<0.001).

Furthermore, the GCS-Net algorithm was shown to be time-saving. It took about 10 minutes to segment and calculate ChT for 3,192 B-scans in the test dataset, while the same task took approximately 72 hours to complete manually by 1 clinician alone.

#### Discussion

To our knowledge, this was the first study to use HM eyes as test objects for evaluating automated choroid segmentation and thickness calculation. The GCS-Net shows great agreements with manual segmentation and provides a fast and reliable tool for calculating the thin choroid in HM eyes.

Manual image analysis is time-consuming and subjective. Moreover, automatic segmentation using an internal algorithm in SS-OCT is sometimes inaccurate, which may be due to errors caused by over-smoothing or artifact interference. Therefore, it is necessary to develop credible, objective automated methods to segment the choroid and measure its thickness. There are some studies that included the quantification of retinal features and ChT in AMD eyes (32-34), investigations on optic disc changes and their association with ChT in young myopic patients (35), and the use of algorithms to determine changes in ChT and



**Figure 3** Example B-scans with choroid segmentation results overlaid. The green and red outline represents the ground truth and the prediction, respectively. (A1-A3) Images of a 73-year-old female (axial length, 27.72 mm). (B1-B3) Images of a 70-year-old male (axial length, 28.28 mm). (C1-C3) Images of a 63-year-old female (axial length, 29.13 mm). A perforating scleral vessel toward the subfoveal choroid was observed. (A1, B1 and C1) Original B-scans. (A2, B2 and C2) Ground truth. (A3, B3 and C3) The proposed GCS-Net. GCS-Net, groupwise context selection network.

volume of choroidal vessels (34,36,37). However, previous studies of automated ChT calculations were designed based on mostly non-HM eyes (38-40). Automated identification and calculation of ChT in HM eyes still warrant further study, and our team has research experience in this field (30,41).

Our proposed GCS-Net algorithm to automatically segment the choroid showed good agreement with manual segmentation. The thickness of the choroid was subsequently calculated and its results were compared with manual segmentation calculated by the built-in Topcon software. The average ChT in the test dataset composed of images with HM was  $75.9\pm34.4 \mu$ m. As expected, the ChT was thinner than that reported in previous studies, as they had tended to focus on non-HM participants (38,39). By analyzing the difference of ICCs in the 9 regions, the ChT calculated in the outer nasal region was less consistent with the manual. The possible reason is that this region is close to the optic disc, which increases the segmentation difficulty. However, ICC was above 0.90 for all regions of the choroid, and we believe this slight inaccuracy in the outer nasal region is clinically acceptable.

In this study, we measured the average ChT at 9 different regions, and a decreasing trend of ChT was



**Figure 4** The distribution of choroidal thickness for high myopia eyes calculated by GCS-Net and manual adjustment in horizontal (A) and vertical (B) direction. GCS-Net, group-wise context selection network.

Table 3 Average, minimum, maximum, and mean error difference between the manual and GCS-Net automated choroidal thickness measurements for all nine ETDRS regions in the test dataset. The intraclass correlation coefficient between the two measurements is also presented

Regions (ETDRS)	GCS-Net thickness (µm)	Manual thickness (µm)	Minimum error (µm)	Maximum error (µm)	Error (µm)	ICC (P value)
Center	69.6±39.1	71.6±35.5	0.02	28.86	5.27±5.13	0.981 (P<0.001)
Inner_temporal	78.6±40.5	79.4±38.1	0.02	35.58	4.79±4.81	0.985 (P<0.001)
Inner_superior	76.8±41.2	77.9±38.2	0.08	47.81	5.42±5.90	0.980 (P<0.001)
Inner_nasal	69.6±35.8	69.4±33.0	0.05	24.76	5.03±4.07	0.982 (P<0.001)
Inner_inferior	77.4±37.1	77.6±35.0	0.01	46.42	5.14±5.22	0.979 (P<0.001)
Outer_temporal	86.5±39.9	88.7±38.7	0.03	31.70	5.92±5.26	0.980 (P<0.001)
Outer_superior	84.3±38.3	86.3±36.4	0.02	26.53	5.19±5.20	0.981 (P<0.001)
Outer_nasal	59.4±26.6	60.7±23.7	0.02	39.14	6.59±5.32	0.944 (P<0.001)
Outer_inferior	82.1±37.6	81.8±36.2	0.02	23.75	4.44±3.71	0.988 (P<0.001)
Average	75.9±34.4	78.5±30.6	<0.01	24.07	5.54±4.57	0.976 (P<0.001)

GCS-Net, group-wise context selection network; ETDRS, Early Treatment of Diabetic Retinopathy Study; ICC, intraclass correlation coefficient.

found from the temporal to the nasal region of the macula, which corresponded with some previous research (42,43). In contrast, in the vertical direction, the ChT was quite symmetrical (*Figure 4*). The superior and inferior regions were thicker, and the fovea was thinnest. Most manual ChT measuring methods in previous studies were at the subfoveal location and at certain distances of the temporal, superior, nasal, and inferior quadrants from the fovea (42,44-46). However, the selected points may not necessarily represent all choroidal pathologies, and variations in the selection of the point used by the observers may lead to interobserver variation. Our study has provided an effective method to manage these problems, and the data could be used to facilitate the differential diagnosis of ocular pathologies that may cause variations in thickness at specific zones within the choroid.

The deep learning algorithm-based GCS-Net is much more efficient and robust than traditional methods, and our

previous study compared it to other algorithms (30). Using traditional methods, preprocessing is often needed, such as denoising, enhancement, and retina layer segmentation (27-29), while the GCS-Net is an end-to-end method that produces results from the original B-scans. While many traditional methods such as graph-based ones require the choroid region to be continuous, the GCS-Net can accurately detect the discontinuity caused by the optic disc, pathological atrophy, or retinal folding artifacts. In addition, GCS-Net is a lightweight network with a relatively small number of parameters. This allows successful learning even with a medium-sized training dataset.

Our study had several limitations that have to be acknowledged. First, the algorithm was learnt and tested in the same environment (i.e., the same clinical dataset), thus limiting its application. Second, the algorithm was developed and tested mainly on SS-OCT images centered on the fovea, which highlights the need to validate the algorithm's performance for the choroid located near the optic disc. Finally, the ChT maps were not evaluated in this study as the automatic topographic map reconstruction from 12-radial scans would lead to large errors. The assessment of the thickness of the choroid on volumetric scans was beyond the scope of this study. To achieve choroid segmentation on images acquired with other scanning protocols, the model has to be retrained with the specific images, or transfer learning techniques need to be applied. The current model segments each B-scan independently. Further extensions that use multiple adjacent B-scans or the whole volume as input should further explore context information to improve accuracy.

In conclusion, the GCS-Net algorithm proposed in our study provides a reliable and fast method to automatically segment and calculate the ChT in HM eyes. Therefore, this tool could be used as a monitoring tool to assess the ChT in HM patients. Moreover, it could also provide a powerful tool for further research on ocular diseases related to thinning of the choroid that could lead to visual impairment.

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#### Footnote

*Reporting Checklist:* The authors have completed the TRIPOD reporting checklist. Available at https://atm. amegroups.com/article/view/10.21037/atm-21-6736/rc

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*Ethical Statement:* The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013). The study was approved by the Ethics Committee of the Shanghai General Hospital, Shanghai Jiao Tong University School of Medicine (No. 2015KY153), and informed consent was provided by all individual participants.

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