




Review Article

Microvascular damage assessed by optical coherence tomography angiography for glaucoma diagnosis: a systematic review of the most discriminative regions

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ABSTRACT.

A growing number of studies have reported a link between vascular damage and glaucoma based on optical coherence tomography angiography (OCTA) imaging. This multitude of studies focused on different regions of interest (ROIs) which offers the possibility to draw conclusions on the most discriminative locations to diagnose glaucoma. The objective of this work was to review and analyse the discriminative capacity of vascular density, retrieved from different ROIs, on differentiating healthy subjects from glaucoma patients. PubMed was used to perform a systematic review on the analysis of glaucomatous vascular damage using OCTA. All studies up to 21 April 2019 were considered. The ROIs were analysed by region (macula, optic disc and peripapillary region), layer (superficial and deep capillary plexus, avascular, whole retina, choriocapillaris and choroid) and sector (according to the Garway–Heath map). The area under receiver operator characteristic curve (AUROC) and the statistical difference (p-value) were used to report the importance of each ROI for diagnosing glaucoma. From 96 screened studies, 43 were eligible for this review. Overall, the peripapillary region showed to be the most discriminative region with the highest mean AUROC (0.80 ± 0.09). An improvement of the AUROC from this region is observed when a sectorial analysis is performed, with the highest AUROCs obtained at the inferior and superior sectors of the superficial capillary plexus in the peripapillary region (0.86 ± 0.03 and 0.87 ± 0.10 , respectively). The presented work shows that glaucomatous vascular damage can be assessed using OCTA, and its added value as a complementary feature for glaucoma diagnosis depends on the region of interest. A sectorial analysis of the superficial layer at the peripapillary region is preferable for assessing glaucomatous vascular damage.

Key words: glaucoma – microvascular analysis – multilayer analysis – ocular blood flow – optical coherence tomography angiography – regions of interest – vascular damage

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Introduction

Glaucoma is the leading cause of irreversible blindness worldwide, with primary open-angle glaucoma (POAG) as its most prevalent form (Van Melkebeke et al. 2018). Glaucoma is a multifactorial disease characterized by the loss of neural retinal ganglion cells. Classic theories attribute glaucomatous neuronal damage to mechanical trauma caused by elevated intraocular pressure (IOP) or to dysfunction of vascular perfusion and subsequent optic nerve ischaemia (Halpern & Grosskreutz 2002). Although elevated IOP remains the only confirmed modifiable risk factor for development and progression of glaucoma (Kass & Gordon 2000; Heijl et al. 2002; Jesus et al. 2017), differences in vascular parameters have been continuously reported between glaucoma and healthy individuals, at both ocular and systemic level (Barbosa-Breda et al. 2019a,b).

A number of techniques, such as fluorescein angiography, colour Doppler imaging, laser speckle flowgraphy and laser Doppler flowmetry, have been used in the evaluation of ocular and retinal blood perfusion (Michelson

et al. 1996; Sugiyama et al. 2010; Stalmans et al. 2011; Spaide et al. 2015; Abegão Pinto et al. 2016; Barbosa-Breda et al. 2019a,b). The application of these modalities to glaucoma has contributed to a more comprehensive assessment of the role of vascular supply in the disease pathophysiology. With the introduction of new imaging modalities such as optical coherence tomography angiography (OCTA), standard OCT devices are now capable of analysing retinal blood flow, which is witnessed by extensive literature already published in ocular diseases (Fang et al. 2016; Koustenis et al. 2017). Due to the early stage of the technology, different strategies have been proposed and are currently used to retrieve the angiographic data from the OCT scans (e.g. the split-spectrum amplitude-decorrelation angiography (Jia et al. 2012), the OCT-based microangiography (Zhang & Wang 2015), the OCTA ratio analysis (OCTARA) (Stanga et al. 2016) and the speckle variance OCTA (Xu et al. 2014)), which may lead to variability between results. Besides that, the quality of the OCTA scans may change significantly according to the acquisition parameters (e.g. number of images to be averaged) or artefacts such as eye movements. A number of studies have also developed and used different image processing algorithms to measure the vessel density from the OCTA images (e.g. percentage of vessel pixels in the respective region (Yip et al. 2019), mean intensity from the grayscale image (Jesus et al. 2019) or fractal analysis (Gadde et al. 2016)). Despite the differences that may exist between the OCTA imaging strategies and the algorithms to compute the vessel density, significantly lower vessel density and blood flow index in the macula (Akil et al. 2017; Chen et al. 2017; Chung et al. 2017; Alnawaiseh et al. 2018), optic disc (Bojikian et al. 2016; Chen et al. 2016; Cennamo et al. 2017; Chen et al. 2017; Chihara et al. 2017) and peripapillary region (Akil et al. 2017; Alnawaiseh et al. 2018; Lin et al. 2019) have been observed in glaucoma eyes in comparison with healthy ones. For all these regions, the diagnostic abilities increased with the severity of glaucoma (Chen et al. 2017; Chihara et al. 2017; Chung et al. 2017). Current results achieved with OCTA have presented it as a potential alternative or

complementary technology for assisting glaucoma diagnosis. In comparison with the current imaging examinations used for the diagnosis and follow-up, OCTA has shown to be less affected by the floor effect observed on structural OCT analysis and to require less patient cooperation than visual field testing (Van Melkebeke et al. 2018).

As in many medical imaging technologies at their early development stage, a number of approaches for estimating the microvascular density based on different regions of interest (ROIs) have been proposed. However, data reported in these approaches are often conflicting and/or arising from small-scale studies, hindering the development of a general methodology to study glaucomatous vascular damage. Microvascular density measured from OCTA has shown to be device-dependent, artefact-dependent (e.g. eye motion, vitreous floaters, and media opacities) (Spaide et al. 2016; Sánchez Brea et al. 2019) and, more importantly, dependent on the imaged ROI. Since OCTA imaging is restricted to a narrow field of view, and the acquisition of a single image with good quality (i.e. no movement artefacts and good contrast) often requires a long exposure time (in patients known to have poor ocular surface and sometimes poor fixation capacities), it is important to ensure an efficient image acquisition, focusing first in the ROIs that yield more relevant information. Moreover, the distribution of the vascular glaucomatous damage among retinal, choriocapillaris and choroid layers is still under research. It is not clear yet whether the significant changes observed at the choriocapillaris and choroid are due to imaging artefacts or due to an actual disease mechanism (Sousa et al. 2019).

The aim of this systematic review was to contribute to the understanding of the role of vascular damage in glaucoma. To that end, the review focuses on the vascular density retrieved from the different ROIs that have been studied so far in the literature, reporting which ROIs have been found to be the most promising for studying glaucoma.

Methods

This research adhered to the Preferred Items for Systematic Reviews and Meta-analyses (PRISMA) guidelines.

Study selection

A literature search was carried out in the PubMed database. The search query can be found in Appendix A. All studies that were published from the 1 January 2014 to the 21 April 2019 were included. The inclusion criteria for each study were as follows: (i) primary study, (ii) mention how the vessel density was computed, (iii) English language, (iv) conducted in humans, (v) investigate glaucomatous eyes in comparison with a healthy control group and (vi) reports at least: the area under the receiver operating characteristic curve (AUROC) or the statistical difference between the control and glaucoma groups. Four authors (A.B., N.B., V.E. and E.F.) screened all the titles and abstracts independently. A full-text screening was carried out by two authors (N.B. and E.F.) independently. In case of disagreement, a third author (A.B. or V.E.) was consulted to reach consensus.

Data collection

The extracted data included the following: study characteristics, AUROC values for different ROIs, microvascular density mean and standard deviation, and p-values from the statistical comparison between healthy and glaucoma groups. If no statistical comparison or p-values were provided, only the AUROC values were collected and *vice versa*.

For every study, the following characteristics were extracted: sample size including number of patients and eyes for each group, average age in years, and statistical difference (p-value) between groups, glaucoma severity, OCT device brand and respective light-source wavelength, cut-off value for the signal strength index (SSI) (or similar image quality measure) used to exclude patients/eyes and field of view of the OCTA image.

Data collection included the different layers: retina (including superficial, deep and avascular), choriocapillaris and choroid (Fig. 1A); the different regions: macula, optic disc (OD) and peripapillary or circumpapillary (when a circular band around the optic disc was considered instead of whole image) (Fig. 1B); and the sectors according to the Garway–Heath map (Garway–

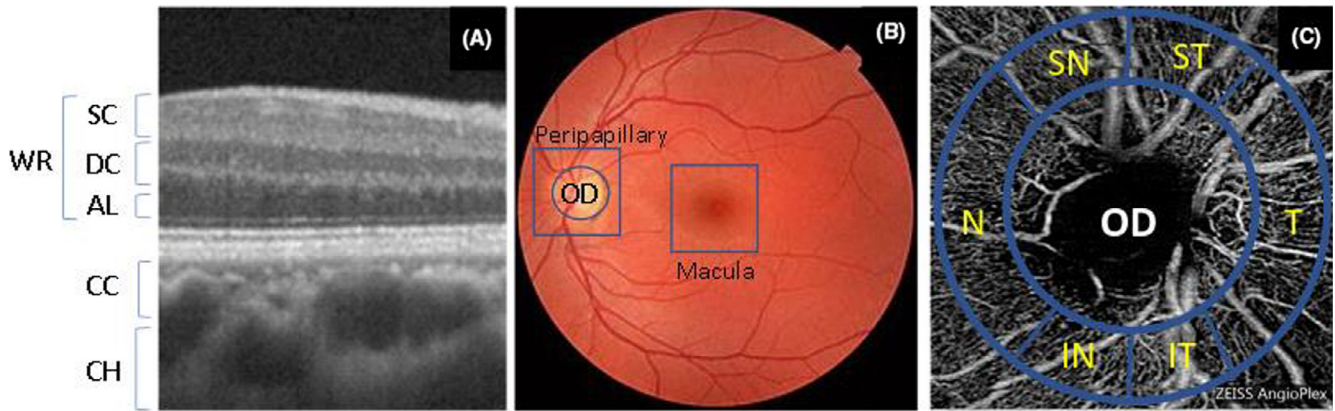


Fig. 1. Regions of interest (ROIs) considered in this review for the analysis of the glaucomatous vascular damage. (A) Optical coherence tomography image showing the retinal superficial capillary plexus (SC), deep capillary plexus (DC), avascular layer (AL), whole retina (WR), choriocapillaris (CC) and choroid (CH). (B) Fundus image highlighting the macula, optic disc (OD) and peripapillary regions. (C) Optical coherence tomography angiography image with a circumpapillary representation of the Garway–Heath sectors: superonasal (SN), superotemporal (ST), temporal (T), inferotemporal (IT), inferonasal (IN) and nasal (N).

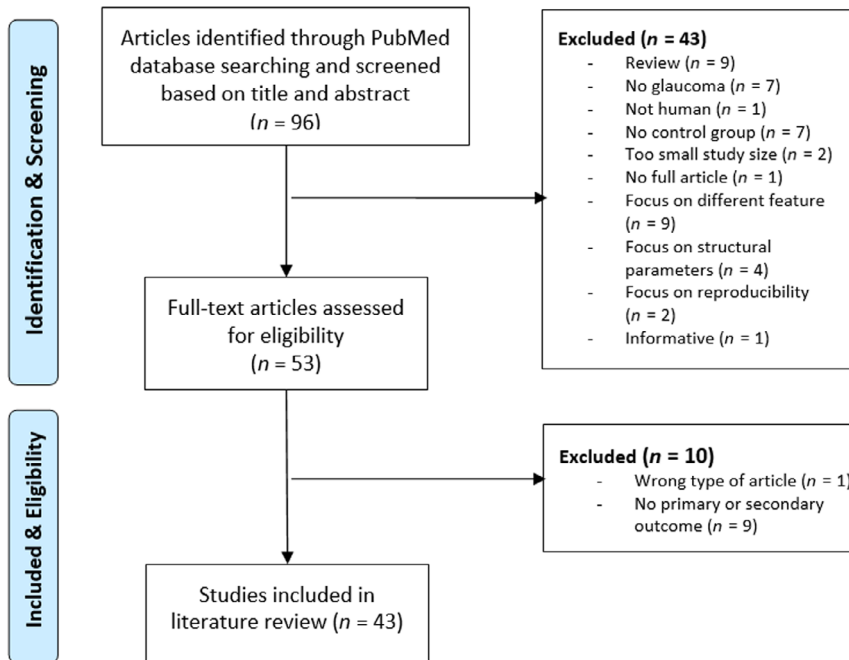


Fig. 2. Flowchart of the study selection.

Heath et al. 2000): superonasal (SN), superotemporal (ST), temporal (T), inferotemporal (IT), inferonasal (IN), nasal (N) and the inside disc (D) (Fig. 1C). The ‘whole retina’ was treated as a layer (Fig. 1A). Moreover, the ‘whole region’ (all the sectors combined) and the ‘fovea’ (centre of the macular region) were considered as sectors for the purpose of this review.

Data analysis

The collected data were used to find which ROIs (region, layer and sector)

have been studied and their discriminating power between glaucoma and healthy controls. Since the data analysis was oriented towards the clinical interpretation, the microvascular density was treated as a generic feature, not taking into account the different mathematical approaches used to estimate it.

The AUROC was considered as the most relevant metric for evaluating which ROIs are the most promising for studying glaucoma, since it provides the performance measurement for classification problem at various

thresholds settings. The AUROC of all ROIs included in all reviewed studies was averaged to determine a threshold for selecting the studies that would undergo a qualitative assessment (described in section Qualitative assessment). The statistically significant differences (given as p-values) were used to complement the information provided by the AUROCs and assess whether a ROI was relevant for differentiating glaucoma from healthy controls.

For those studies that did not report an AUROC, the decision to perform a qualitative assessment was based on the statistical comparison between the glaucoma and the healthy group. Hence, the ROIs that presented significant statistical differences (p-value < 0.05) were also qualitatively evaluated.

Qualitative assessment

There are several characteristics in a study that have been reported in the literature as potentially impacting the outcome of OCTA-based glaucoma assessment and thus introducing bias. Thus, despite the high AUROC a method may have, it does not dismiss a careful qualitative analysis to identify these potential sources of bias. Hence, in this review, all the studies that reported an AUROC value above the threshold (mean AUROC values for all studied ROIs), and the ability to significantly differentiate glaucoma from healthy subjects, were qualitatively assessed. The qualitative assessment

Table 1. Area under the receiver operating characteristic curve (AUROC) for differentiating glaucoma eyes from healthy controls according to the region, layer and sector for all reviewed studies.

Sector	Region		
	Optic disc	Macula	Peripapillary region
1. Superficial capillary plexus			
Whole image/region	0.92 (Yip et al. 2019) 0.78 (Shin et al. 2017) 0.77 (Rao et al. 2017c) 0.76 (Rao et al. 2017b) 0.75 (Alnawaiseh et al. 2018) 0.73 (Rao et al. 2017a) 0.57 (Chung et al. 2017) 0.57 (Chen et al. 2016)	0.96 (Takusagawa et al. 2017) 0.94 (Rabiolo et al. 2018) 0.94 (Chung et al. 2017) 0.94 (Chen et al. 2017) 0.92 (Rabiolo et al. 2018) 0.84 (Yip et al. 2019) 0.84 (Rabiolo et al. 2018) 0.84 (Rabiolo et al. 2018) 0.82 (Rabiolo et al. 2018) 0.80 (Kuryшева et al. 2018) 0.79 (Rabiolo et al. 2018) 0.78 (Lommatzsch et al. 2018) 0.77 (Rabiolo et al. 2018) 0.75 (Kuryшева et al. 2018) 0.71 (Rao et al. 2017c) 0.70 (Triolo et al. 2017) 0.70 (Rao et al. 2017d) 0.69 (Alnawaiseh et al. 2018) 0.67 (Rao et al. 2017c)* 0.52 (Kwon et al. 2017)	0.96 (Akil et al. 2017) 0.96 (Akil et al. 2017) 0.94 (Yarmohammadi et al. 2016a) 0.93 (Chen et al. 2017) 0.90 (Akil et al. 2017) 0.90 (Rolle et al. 2019) 0.88 (Triolo et al. 2017) 0.85 (Rao et al. 2017d) 0.85 (Rao et al. 2017b) 0.83 (Rao et al. 2017a) 0.82 (Akil et al. 2017) 0.81 (Chung et al. 2017) 0.80 (Cennamo et al. 2017) 0.80 (Kuryшева et al. 2018) 0.78 (Lommatzsch et al. 2018) 0.76 (Akil et al. 2017) 0.76 (Akil et al. 2017) 0.69 (Alnawaiseh et al. 2018)
Inside disc	0.91 (Rao et al. 2017b) 0.81 (Alnawaiseh et al. 2018) 0.72 (Rolle et al. 2019) 0.60 (Kiyota et al. 2018)		
Superior	0.95 (Geyman et al. 2017) 0.78 (Shin et al. 2017) 0.73 (Rao et al. 2017d)	0.98 (Takusagawa et al. 2017) 0.79 (Kuryшева et al. 2018) 0.69 (Lommatzsch et al. 2018) 0.67 (Lommatzsch et al. 2018) 0.65 (Rao et al. 2017b) 0.65 (Rao et al. 2017d) 0.63 (Rao et al. 2017a) 0.56 (Triolo et al. 2017)	1.00 (Akil et al. 2017) 0.98 (Akil et al. 2017) 0.95 (Chung et al. 2017) 0.86 (Chung et al. 2017) 0.82 (Rao et al. 2017d) 0.77 (Rao et al. 2017b) 0.74 (Triolo et al. 2017)
Inferior	0.89 (Geyman et al. 2017) 0.84 (Shin et al. 2017) 0.67 (Rao et al. 2017d)	0.98 (Takusagawa et al. 2017) 0.69 (Kuryшева et al. 2018) 0.69 (Rao et al. 2017d) 0.68 (Lommatzsch et al. 2018) 0.68 (Lommatzsch et al. 2018) 0.61 (Rao et al. 2017a) 0.54 (Triolo et al. 2017)	0.89 (Chung et al. 2017) 0.88 (Rao et al. 2017d) 0.86 (Chung et al. 2017) 0.80 (Triolo et al. 2017)
Nasal	0.74 (Rao et al. 2017d) 0.70 (Rao et al. 2017a) 0.54 (Shin et al. 2017)	0.70 (Kuryшева et al. 2018) 0.68 (Lommatzsch et al. 2018) 0.68 (Lommatzsch et al. 2018) 0.65 (Rao et al. 2017d) 0.56 (Rao et al. 2017a)	0.86 (Kuryшева et al. 2018) 0.85 (Chung et al. 2017) 0.84 (Rao et al. 2016) 0.82 (Chung et al. 2017) 0.78 (Rao et al. 2017d) 0.73 (Triolo et al. 2017) 0.72 (Rao et al. 2017b) 0.70 (Rao et al. 2017a) 0.59 (Rolle et al. 2019) 0.86 (Chung et al. 2017)
Temporal	0.71 (Shin et al. 2017) 0.70 (Rao et al. 2017d)	0.74 (Kuryшева et al. 2018) 0.72 (Lommatzsch et al. 2018) 0.71 (Lommatzsch et al. 2018) 0.67 (Rao et al. 2017d) 0.64 (Rao et al. 2017a)	0.83 (Chung et al. 2017) 0.79 (Kuryшева et al. 2018) 0.75 (Rolle et al. 2019) 0.70 (Rao et al. 2017a) 0.68 (Triolo et al. 2017) 0.68 (Rao et al. 2017d) 0.48 (Rao et al. 2016)
Temporal superior	0.71 (Rao et al. 2017a)	0.58 (Triolo et al. 2017)	0.83 (Rao et al. 2017b) 0.81 (Kuryшева et al. 2018) 0.76 (Rao et al. 2017a) 0.71 (Rao et al. 2017c)* 0.71 (Rao et al. 2017c) 0.68 (Rao et al. 2016) 0.56 (Rolle et al. 2019)

Table 1. (Continued)

Sector	Region		
	Optic disc	Macula	Peripapillary region
Nasal superior	0.83 (Geyman et al. 2017) 0.61 (Rao et al. 2017a) 0.59 (Rao et al. 2017a)	0.62 (Triolo et al. 2017)	0.78 (Kuryшева et al. 2018) 0.78 (Rao et al. 2017b) 0.72 (Rao et al. 2016) 0.70 (Rao et al. 2017a) 0.65 (Rolle et al. 2019) 0.94 (Kuryшева et al. 2018) 0.89 (Rao et al. 2017a) 0.88 (Rao et al. 2016) 0.84 (Rao et al. 2017b) 0.83 (Rao et al. 2017c) 0.75 (Rao et al. 2017c)*
Temporal inferior	0.61 (Rao et al. 2017a)	0.61 (Triolo et al. 2017)	0.88 (Kuryшева et al. 2018) 0.81 (Rao et al. 2017a) 0.78 (Rao et al. 2017b) 0.77 (Rao et al. 2016) 0.70 (Rolle et al. 2019)
Nasal inferior		0.59 (Triolo et al. 2017)	0.88 (Kuryшева et al. 2018) 0.81 (Rao et al. 2017a) 0.78 (Rao et al. 2017b) 0.77 (Rao et al. 2016) 0.70 (Rolle et al. 2019)
Circumpapillary Whole image			0.89 (Jesus et al. 2019) 0.89 (Chen et al. 2017) 0.87 (Kwon et al. 2017) 0.53 (Kiyota et al. 2018)
Nasal			0.78 (Jesus et al. 2019)
Temporal			0.77 (Jesus et al. 2019)
Temporal superior			0.85 (Jesus et al. 2019)
Nasal superior			0.79 (Jesus et al. 2019)
Temporal inferior			0.87 (Jesus et al. 2019)
Nasal inferior			0.86 (Jesus et al. 2019)
2. Deep capillary plexus Whole image/region	0.67 (Shin et al. 2017)	0.99 (Rabiolo et al. 2018) 0.99 (Rabiolo et al. 2018) 0.99 (Rabiolo et al. 2018) 0.97 (Rabiolo et al. 2018) 0.92 (Rabiolo et al. 2018) 0.86 (Yip et al. 2019) 0.79 (Rabiolo et al. 2018) 0.70 (Rabiolo et al. 2018) 0.70 (Lommatzsch et al. 2018) 0.70 (Alnawaiseh et al. 2018) 0.63 (Rao et al. 2017a)	0.70 (Lommatzsch et al. 2018)
Superior	0.63 (Shin et al. 2017)	0.69 (Lommatzsch et al. 2018) 0.69 (Lommatzsch et al. 2018)	
Inferior	0.74 (Shin et al. 2017)	0.71 (Lommatzsch et al. 2018) 0.69 (Lommatzsch et al. 2018)	
Nasal	0.52 (Shin et al. 2017)	0.71 (Lommatzsch et al. 2018) 0.71 (Lommatzsch et al. 2018)	
Temporal	0.66 (Shin et al. 2017)	0.72 (Lommatzsch et al. 2018) 0.67 (Lommatzsch et al. 2018)	
3. Whole retina Whole image/region	0.96 (Yip et al. 2019) 0.93 (Akil et al. 2017) 0.91 (Rao et al. 2017c) 0.90 (Kuryшева et al. 2018) 0.86 (Akil et al. 2017) 0.84 (Kuryшева et al. 2018) [†] 0.82 (Rao et al. 2017c)* 0.77 (Rao et al. 2017c) [†] 0.74 (Alnawaiseh et al. 2018) 0.74 (Rao et al. 2017c) [†]	0.91 (Takusagawa et al. 2017)	
4. Choriocapillaris Whole image/region		0.84 (Alnawaiseh et al. 2018)	0.83 (Yarmohammadi et al. 2016a)
5. Choroid Whole image/region	0.76 (Yip et al. 2019)		

The bold font highlights all numerical values above the selected threshold (AUROC > 0.77). No values were reported for the avascular layer. The whole image/region is defined as all sectors combined.

* With Disc Haemorrhage.

[†] Inside disc.

to measure the risk of bias was performed independently by two authors (A.B. and E.F.). Criteria were composed in cooperation with experienced ophthalmologists (J.B.B. and I.S.; Appendix B). The following six aspects, ordered by relevance, were considered:

1 *Age*. The age should not differ significantly between the glaucoma and the healthy groups. If there is an age difference between groups, an adjustment should be executed. Otherwise, the outcomes are considered as less reliable, because the microvascular density decreases with age (Lin et al. 2019).

2 *Eye*. Measurements obtained from both eyes of a subject are likely correlated. Hence, unless proper statistical methods are employed, there is a higher risk of bias if both eyes are included in the study.

3 *Type and severity of glaucoma*. Studies have a higher risk of bias when they report combined results of primary and secondary types of glaucoma, because of the difference in pathophysiology. Furthermore, the more severe the glaucoma, the more advanced the damage, not allowing to accurately infer the sensitivity of the studied feature. This means that a classification problem with high AUROC values for severe glaucoma may not be a good predictor for early diagnosis, even though they could still be good features for follow-up.

4 *OCT specifications*. Different hardware specifications play a role in OCT image quality, especially in deeper layers such as the choriocapillaris and the choroid. Results from studies using different hardware should not be compared to each other but rather discussed separately.

5 *Image quality*. Studies that included images with SSI values (or similar quality measures) below the suggested inclusion value provided by the manufacturer are at higher risk of bias. Suggested values by manufacturers: for Angioplex[®], include if >6 (out of 10); for AngioVue[®], include if >45 (out of 100) (Spaide et al. 2016).

6 *Fovea-disc axis correction*. If a sectorial analysis is performed, fovea-disc axis correction should be executed for all the OCTA images to assure that the features are computed for the same ROIs between subjects

(e.g. using a Panomap[®] image; or any other reference of the relative position of the fovea and the optic disc (Mwanza et al. 2015; Jesus et al. 2019)).

Results

Study selection

Ninety-six studies were identified using the search query in Appendix A. From those, 53 studies were considered eligible after screening the titles and abstracts. Full-text screening resulted in 43 studies that met all inclusion criteria and, hence, were eligible for the data analysis (Fig. 2). All the included studies provided a statistical analysis of the quantitative vascular evaluation for different ROIs. Twenty-four studies provided AUROC as an outcome. The complete table with the characteristics of the reviewed studies can be found in Appendix B.

AUROC analysis

The AUROCs presented in the reviewed studies are summarized in Table 1 organized per layer, region and sector. All studies calculated AUROC values based on the microvascular density, despite using different image processing techniques for intensity quantification or binarization. Although the macular region showed the highest AUROC values (considering all studies individually), when taking the mean of all ROIs, the peripapillary region had the highest AUROC of 0.80 ± 0.09 , whereas the macula and the optic disc both had AUROC of 0.74 ± 0.12 . The mean AUROC values for all studied ROIs are shown in Fig. 3. The average of all AUROC values in Table 1 is 0.77, which was set as the threshold for deciding whether a study or ROI should be further analysed in the qualitative assessment. All three regions (optic disc, macular and peripapillary) yielded values above this threshold, as shown in Table 1. Table 1 also shows that the avascular layer was not mentioned in any study, the choriocapillaris only in two studies (Yarmohammadi et al. 2016a; Alnawaiseh et al. 2018) and the choroid in one study (Yip et al. 2019). On the other hand, the whole retina, and the superficial and deep capillary plexuses have been investigated frequently.

Macular region

For the whole image of the macula in the superficial layer, 10 out of 20 values were reported above the threshold (Chung et al. 2017; Takusagawa et al. 2017; Kuryшева et al. 2018; Lommatzsch et al. 2018; Rabiolo et al. 2018; Yip et al. 2019), and 6 out of 11 values were above the threshold in the deep layer (Rabiolo et al. 2018; Yip et al. 2019). Only one value above the threshold was reported for the macula in the whole retina (Takusagawa et al. 2017) and choriocapillaris (Alnawaiseh et al. 2018).

Optic disc

The inside disc (Rao et al. 2017b; Alnawaiseh et al. 2018) and the inferior sector (Geyman et al. 2017; Shin et al. 2017) in the superficial layer were the ROIs with the highest AUROC, based on the reports of two studies. The whole region of the optic disc in the whole retina layer had 7 out of 10 values above the threshold (Akil et al. 2017; Rao et al. 2017c; Kuryшева et al. 2018; Yip et al. 2019).

Peripapillary and circumpapillary region

The whole region (Yarmohammadi et al. 2016a; Akil et al. 2017; Cenamo et al. 2017; Chung et al. 2017; Triolo et al. 2017; Rao et al. 2017a; Rao et al. 2017b; Rao et al. 2017d; Kuryшева et al. 2018; Lommatzsch et al. 2018; Rolle et al. 2019), superior (Akil et al. 2017; Chung et al. 2017; Rao et al. 2017d), inferior (Chung et al. 2017; Triolo et al. 2017; Rao et al. 2017d) and temporal inferior sectors (Rao et al. 2016; Shin et al. 2017; Rao et al. 2017a; Rao et al. 2017b; Kuryшева et al. 2018) in the superficial layer often presented AUROC values above the threshold. Also, for the whole region of the circumpapillary ROI (circular band in the peripapillary region) in the superficial layer, multiple values were reported above the threshold (Chen et al. 2017), (Kwon et al. 2017), (Jesus et al. 2019). Only one AUROC for the whole region in the choriocapillaris above the threshold (Yarmohammadi et al. 2016a) was reported.

p-value analysis

The results for the vascular density differed greatly between and within

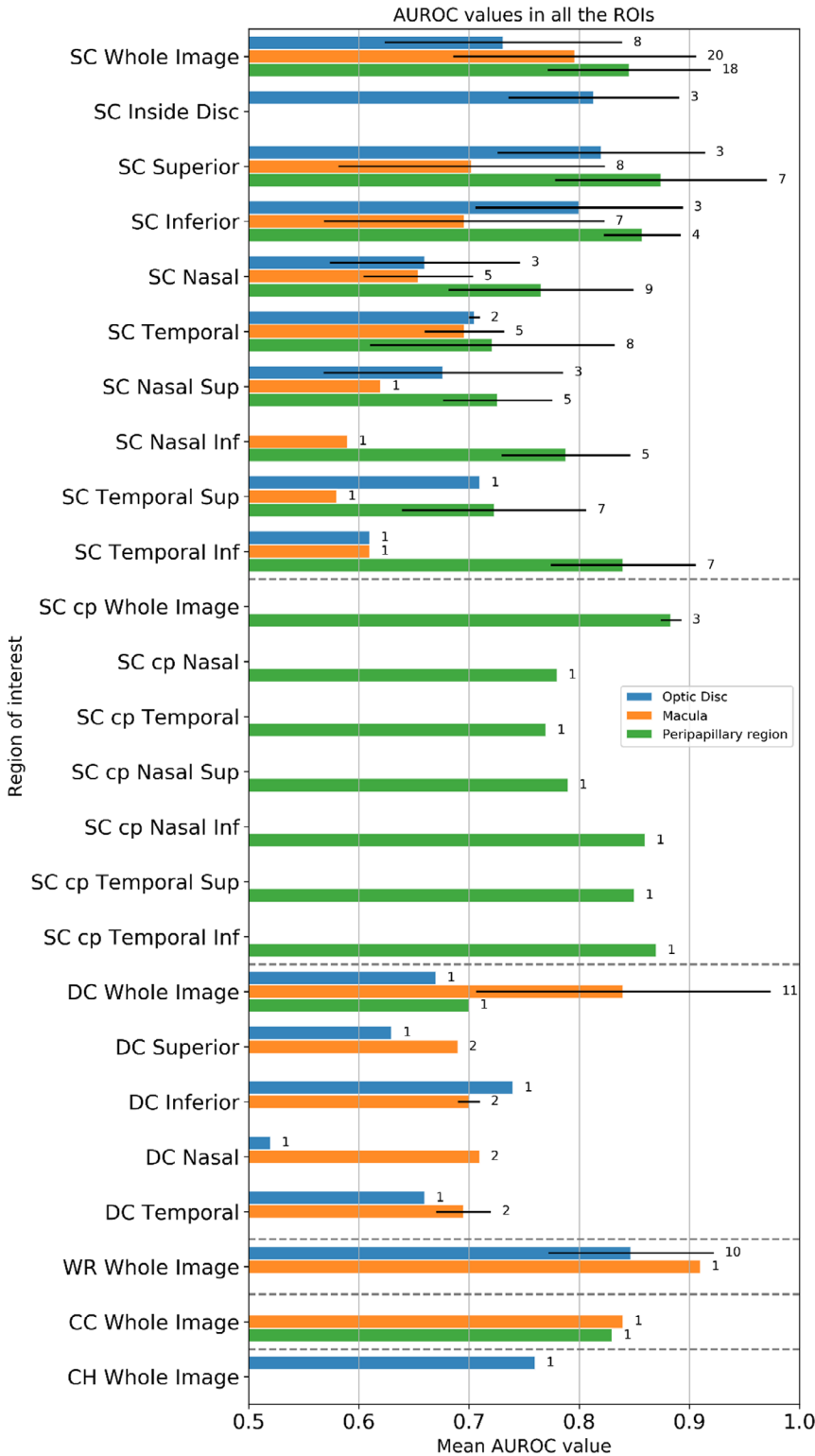


Fig. 3. Mean AUROC and standard deviation value/number of observations for each ROI. AUROC = area under the receiver operating characteristic curve, CC = choriocapillaris, CH = choroid, cp = circumpapillary, DC = deep capillaris plexus, ROI = Regions of interest, SC = superficial capillaris, WR = whole retina.

ROIs, as shown in Appendix C. Nevertheless, a statistically significant difference between control and glaucoma groups was observed for all the

analysed ROIs. The number of statistically significant differences is summarized in Fig. 4 (and detailed in Table C1 in Appendix C).

Macular region

The whole image of the macula in the superficial layer included 15 out of 17 significant values. Five out of six values reported for the whole image in the deep layer were significant. Only one value, however significant, was reported for the choriocapillaris and none for the choroid.

Optic disc

The inside disc sector in the superficial layer included nine significant values and only one non-significant. The inferior segment in the superficial layer included only three values; however, all of them are significant. Only one out of 17 values reported a non-significant difference for the whole image of the optic disc in the superficial layer. No values were reported for the whole image in the whole retina.

Peripapillary and circumpapillary region

The whole image in the superficial layer included 17 out of 19 significant values. The superior and inferior sectors of the peripapillary region in the superficial layer were not represented as much in the literature. However, all the studies that analysed these regions reported a significant difference between the groups (four and three values, respectively). Seven out of eight values for the temporal inferior sector in the superficial layer were significant. No values were reported for the whole region in the choriocapillaris. Five out of five values were reported as significant for the whole region of the circumpapillary ROI in the superficial layer. One value was reported for the temporal superior, temporal inferior and the nasal inferior sectors in the superficial layer, and all three of them were significant.

Qualitative assessment

The bold font in Table 1 highlights the studies that provided one (or more) AUROC values above the threshold. The complete qualitative assessment was performed in the 22 studies that met the requirement of having an AUROC > 0.77 (Appendix D). From these study characteristics, it was possible to draw the following observations:

1 Age. Six studies (Rao et al. 2016; Yarmohammadi et al. 2016a; Geyman et al. 2017; Rao et al. 2017a; Rabiolo

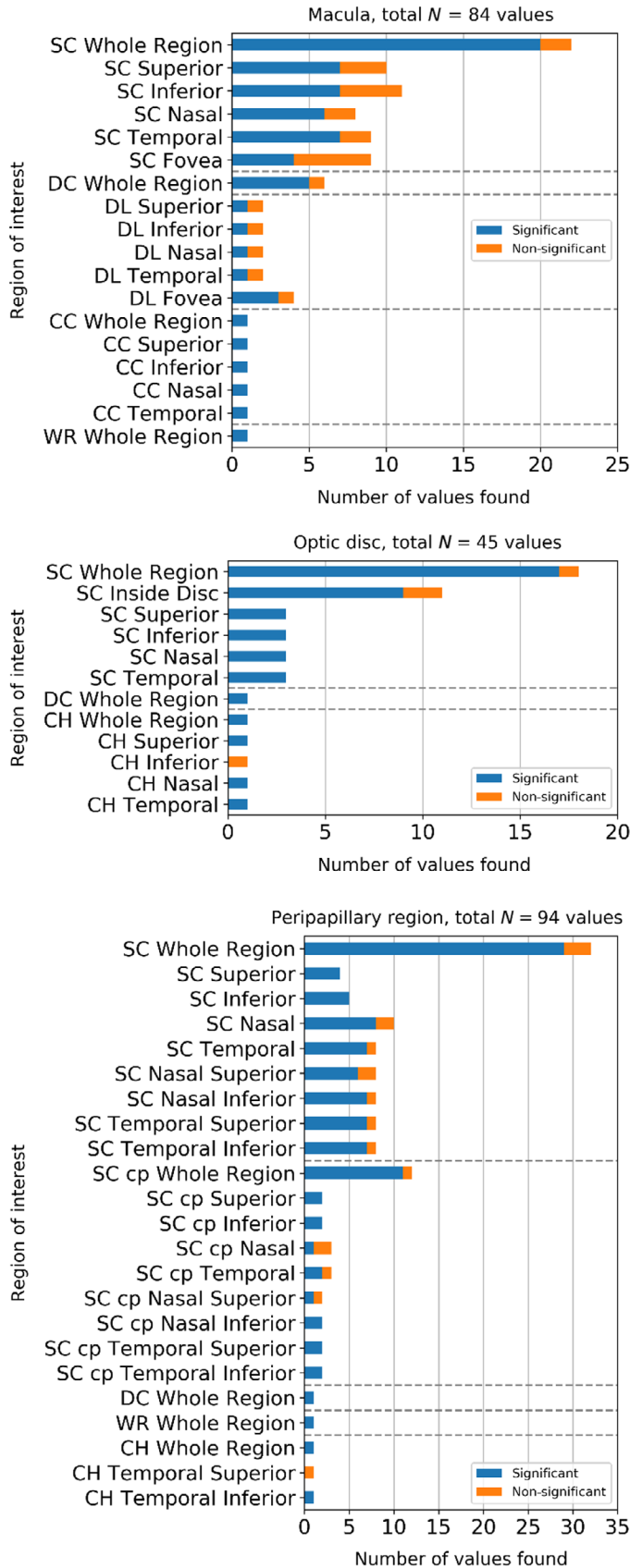


Fig. 4. Number of studies with AUROC values > 0.77 for each ROI or that presented a significant (blue) and non-significant (orange) statistical difference between healthy and glaucoma groups for the three regions; optic disc, macula and the peripapillary. AUROC = area under the receiver operating characteristic curve, CC = choriocapillaris, CH = choroid, cp = circumpapillary, DC = deep capillaris plexus, ROI = Regions of interest, SC = superficial capillaris, WR = whole retina.

et al. 2018; Yip et al. 2019) reported a significant difference in age. However, all of them performed age correction. 2 *Eye*. Ten studies (Rao et al. 2016; Yarmohammadi et al. 2016a; Cennamo et al. 2017; Rao et al. 2017a; Rao et al. 2017b; Rao et al. 2017c; Rao et al. 2017d; Alnawaiseh et al. 2018; Rabiolo et al. 2018; Yip et al. 2019) included both eyes from the same subject. All of these studies except for Alnawaiseh et al. (Alnawaiseh et al. 2018), Cennamo et al. (Cennamo et al. 2017) and Yip et al. (Yip et al. 2019) mentioned to have performed a correction for this. Rolle et al. (Rolle et al. 2019) only mentioned the number of eyes and not the number of subjects included in the study.

3 *Type and glaucoma severity*. No patients with secondary glaucoma were included in any of the studies. All studies used a study population with different levels of glaucoma severity, however, Chen et al. (Chen et al. 2017), Jesus et al. (Jesus et al. 2019) and Yip et al. (Yip et al. 2019) used a patient group with a relatively low visual field mean deviation (MD) (respectively, -8.8 ± 6.2 dB, -7.8 ± 6.5 dB, and -11.07 ± 8.25 dB) when compared to the other studies with an average MD of -6.36 dB.

4 *OCT specifications*. All studies acquired the images with an OCT device with a light-source wavelength of 840 nm, except Akil et al. (Akil et al. 2017), Rabiolo et al. (Rabiolo et al. 2018) and Triolo et al. (Triolo et al. 2017) which used an OCT system with a wavelength of 1040–1060 nm.

5 *Image quality*. Akil et al. (Akil et al. 2017) and Shin et al. (Shin et al. 2017) did not report whether they used a cut-off value for exclusion due to image quality. However, they did report that 5 and 10 images, respectively, were not analysed because of poor OCTA image quality. Eight studies (Rao et al. 2016; Geyman et al. 2017; Rao et al. 2017a; Rao et al. 2017b; Rao et al. 2017d; Lommatzsch et al. 2018; Jesus et al. 2019; Yip et al. 2019) differed from the manufacturer's suggested cut-off value. All these studies used a lower cut-off value than the standard recommended value (Spaide et al. 2016).

6 *Fovea-disc axis correction*. In none of the studies that performed a sectorial analysis, it was mentioned that a fovea-disc axis correction was performed, except for Jesus et al. (Jesus et al. 2019).

Discussion

This systematic review gives an insight into which ROIs have been studied so far in literature and which ones seem to contribute the most to an accurate diagnosis of glaucoma using microvascular density computed from OCTA. The ROIs in OCTA imaging were defined by three arguments: region of acquisition, layer and sector.

The region of acquisition (macula, optic disc or the peripapillary region) should be the first argument to be considered in OCTA imaging, since it is related to the ability to detect glaucomatous vascular damage. Although the highest AUROCs (considering all studies individually) were observed at the macula, the peripapillary region showed the highest AUROCs when averaging all values per region of acquisition. As mean AUROC is a more reliable indicator than its maximum, we may conclude that the peripapillary region is the most relevant for studying glaucomatous vascular damage.

The second argument to be considered is the layer. Overall, the highest AUROCs were obtained for the superficial layer. Nonetheless, the deeper layers presented in some cases similar classification values to the superficial layer. However, the limited number of studies that have covered these deeper layers does not allow to draw conclusions on their added value for the diagnosis. These layers have been avoided due to the difficulty to explain the physical meaning of the imaged content. As light travels deeper through retinal tissue, it becomes more susceptible to refraction and diffraction. Moreover, given the heterogeneity of retinal tissue, light reflection and absorption occur at different levels depending on the region of acquisition and respective refraction index. As a consequence, shadows are projected to deeper layers, creating what is known as projection artefacts. Therefore, and despite the significant differences observed at the choriocapillaris and the choroid, it is difficult to conclude whether these differences arise from the pathology itself or are a consequence of imaging artefacts. Further research needs to be done in order to understand to what extent the information imaged by OCTA at deeper layers is reliable.

The last argument, and the smallest area, is the sector. A sectorial analysis is not always performed in glaucomatous vascular studies. A number of studies have opted for analysing the retinal layers, mainly the superficial vascular plexus, without any sector discrimination. However, for those that performed sectorial analysis, it was shown that microvascular density is affected differently depending on the sector. Taking the most studied region of acquisition and layer as reference (the superficial layer of the peripapillary region), it can be concluded from this review that the inferior sector (AUROC = 0.86 ± 0.03) and the superior sector (AUROC = 0.87 ± 0.10) are the most promising at discriminating glaucoma. Moreover, Fig. 3 shows that a sectorial circumpapillary analysis (with a fixed distance from the optically hollow) seems to provide a better discrimination than a sectorial peripapillary configuration (which takes into account the entire scan). Such a difference may be explained by the reduced variability present in the circumpapillary region, a specific circular ROI with fixed dimensions around the optic disc.

Overall, looking at the number of studies that used OCTA information to infer glaucomatous vascular damage and the respective AUROCs, it can be concluded that the whole region at the superficial layer of the peripapillary ROI is the most accurate measurement for glaucoma assessment, which could be even further improved by a sectorial circumpapillary analysis. This result was somehow expected, since glaucoma is characterized by a loss of optic nerve axons, which traverse the retina superficially in an anatomical area included in the OCTA's superficial layer. Moreover, all the axons meet at the optic nerve which makes a circumpapillary analysis at the peripapillary ROI the best option to capture information from all of them at the same time. Macular scans are indeed relevant but can miss damage that falls outside the macular scan area (Van Melkebeke et al. 2018).

Nevertheless, a certain discrepancy and conflicting results have also been observed between sectors at different layers and regions of acquisition. Possible reasons for such a variability are related to the data and respective study design, and were qualitatively

evaluated. Although no significant differences were observed in terms of age (except for one study which did not provide information (Rolle et al. 2019)), it was noted that three studies (Cennamo et al. 2017; Rao et al. 2017b; Alnawaiseh et al. 2018) used both eyes of the same subject, without mentioning any correction (Appendix D). Age and inclusion of both eyes can constitute a source of bias in the results, since the microvascular density decreases with age, and the data from both eyes are highly correlated. No secondary types of glaucoma were included in any of the reviewed studies. However, three of them (Chen et al. 2017; Jesus et al. 2019; Yip et al. 2019) used a glaucoma group with a relatively low visual field MD (-7.8 dB and lower) which may lead discrepancy between results. The comparison of different regions with data from groups with different severity groups may contribute to the misinterpretation of the data, as the more severe the glaucoma, the more advanced the damage is. Furthermore, three studies (Akil et al. 2017; Triolo et al. 2017; Rabiolo et al. 2018) used a 1040 nm OCT device and achieved a high diagnostic accuracy. All of these devices were Swept-Source OCT (SSOCT) which could potentially indicate that a SSOCT may provide a better OCTA image quality and, consequently, may result in higher AUROCs. Further research is recommended to confirm the advantages of using SSOCT for OCTA imaging in assessing glaucomatous microvascular damage. A high risk of bias was identified in eight studies that included images with an image quality below the threshold suggested by the manufacturer (see Appendix D). Two other studies did not report which threshold was used. Only one study performed a fovea-disc axis correction (Jesus et al. 2019). Due to eye motion or slight differences in position during image acquisition, OCTA images from different subjects might not match the same sectors at the same location. Therefore, sectorial analysis requires images to be previously corrected, for instance taking the fovea-disc axis into account. This way all subjects will have the same reference point for the sectorial analysis.

Another reason for the current variability between studies is related to the method employed to extract vascular density. Although it is not the focus of

this review, different image processing approaches can lead to different vascular interpretations within the same subject data. A popular method among the community is the OCTA image binarization based on thresholding techniques. The ratio of white or black pixels over a specific area is used to estimate the microvascular dropout. In general, the threshold is chosen based on an empirical analysis using an image processing programme such as ImageJ (Abramoff et al. 2004). The separation of micro- from macrovasculature is another source of variability between studies. In some studies, the macrovasculature is segmented and removed from the region of acquisition. Other authors have opted for estimating vascular density based on all the information presented on the OCTA image. Macrovasculature is not expected to be affected by glaucoma, and it is a subject-dependent anatomical feature. Thus, an analysis on image pixel intensity including macrovasculature is not desirable, as it may bias the results. Similarly, the optically hollow area inside the optic disc, as well as the foveal avascular zone (FAZ), is subject-dependent. Therefore, it is desirable to segment and exclude these areas from the ROI before the microvascular density estimation is performed. Nevertheless, further research is needed for a better understanding of the variability between mathematical approaches and to understand which is the most appropriated for glaucoma diagnosis. Although a few research lines have already considered more complex procedures, such as fractal analysis (Gadde et al. 2016), replication studies are still needed to evaluate such advanced/complex methods.

The superior and inferior sectors of the superficial layer of the peripapillary region may be suitable for the diagnosis. However, the averaged AUROC reported in the reviewed articles is still lower than the values obtained with retinal nerve fibre layer thickness (measured through standard OCT imaging) and lower than the optic disc features (extracted from fundus imaging (Hemelings et al. 2020)), which usually result in AUROC values higher than 0.9. Nevertheless, recent studies have shown that vascular density assessed by OCTA seems to perform better than the gold standard biomarkers at discriminating advanced cases of

glaucoma (Barbosa-Breda et al. 2018; Van Melkebeke et al. 2018). Hence, follow-up of (advanced) glaucoma using OCTA imaging may be a window of opportunity to establish OCTA as a common practice in the clinical environment. Thus, new studies will be required to infer which OCTA ROI is the best at glaucoma follow-up.

Conclusions

This review provides a comprehensive summary of the research on glaucomatous microvascular damage based on the analysis of different ROIs imaged with OCTA. The collected data show that the superficial layer in the peripapillary region is the most informative to infer vascular damage. Furthermore, at this location and layer, the inferior and superior sectors have been found as the most discriminative ROIs to study glaucomatous vascular damage with OCTA.

References

Abegão Pinto L, Willekens K, Van Keer K, Shibesh A, Molenberghs G, Vandewalle E & Stalmans I (2016): Ocular blood flow in glaucoma – the Leuven Eye Study. *Acta Ophthalmol* **94**: 592–598.

Abramoff MD, Magalhães PJ & Ram SJ (2004): Image processing with ImageJ. *Biophotonics Int* **11**: 36–42.

Akil H, Huang AS, Francis BA, Satta SR & Chopra V (2017): Retinal vessel density from optical coherence tomography angiography to differentiate early glaucoma, preperimetric glaucoma and normal eyes. *PLoS ONE* **12**: 1–12.

Alnawaiseh M, Lahme L, Müller V, Rosenreter A & Eter N (2018): Correlation of flow density, as measured using optical coherence tomography angiography, with structural and functional parameters in glaucoma patients. *Graefe's Arch Clin Exp Ophthalmol* **256**: 589–597.

Barbosa-Breda J, Andrade de Jesus D, Van Keer K et al. (2018): AngioOCT peripapillary microvascular density outperforms standard OCT parameters as a discriminant between different glaucoma severity levels – The Leuven Eye Study. *Invest Ophthalmol Vis Sci* **59**: 4478.

Barbosa-Breda J, Abegão-Pinto L, Van Keer L et al. (2019a): Improved discrimination between normal-tension and primary open-angle glaucoma with advanced vascular examinations – the Leuven Eye Study. *Acta Ophthalmol* **97**: e50–e56.

Barbosa-Breda J, Abegão-Pinto L, Van Keer K, Jesus DA, Lemmens S, Vandewalle E,

Rocha-Sousa A & Stalmans I (2019b): Heterogeneity in arterial hypertension and ocular perfusion pressure definitions: Towards a consensus on blood pressure-related parameters for glaucoma studies. *Acta Ophthalmol* **97**: e487–e492.

Bojikian KD, Chen CL, Wen JC et al. (2016): Optic disc perfusion in primary open angle and normal tension glaucoma eyes using optical coherence tomography-based microangiography. *PLoS ONE* **11**: e0154691.

Cennamo G, Montorio D, Velotti N, Sparnelli F, Reibaldi M & Cennamo G (2017): Optical coherence tomography angiography in pre-perimetric open-angle glaucoma. *Graefe's Arch Clin Exp Ophthalmol* **255**: 1787–1793.

Chen CL, Zhang A, Bojikian KD et al. (2016): Peripapillary retinal nerve fiber layer vascular microcirculation in glaucoma using optical coherence tomography-based microangiography. *Investig Ophthalmol Vis Sci* **57**: 475–485.

Chen HSL, Liu CH, Wu WC, Tseng HJ & Lee YS (2017): Optical coherence tomography angiography of the superficial microvasculature in the macular and peripapillary areas in glaucomatous and healthy eyes. *Investig Ophthalmol Vis Sci* **58**: 3637–3645.

Chihara E, Dimitrova G, Amano H & Chihara T (2017): Discriminatory power of superficial vessel density and prelaminar vascular flow index in eyes with glaucoma and ocular hypertension and normal eyes. *Investig Ophthalmol Vis Sci* **58**: 690–697.

Chung JK, Hwang YH, Wi JM, Kim M & Jung JJ (2017): Glaucoma diagnostic ability of the optical coherence tomography angiography vessel density parameters. *Curr Eye Res* **42**: 1458–1467.

Fang PP, Lindner M, Steinberg JS, Müller PL, Gliem M, Charbel Issa P, Krohne TU & Holz FG (2016): Klinische Anwendungen der OCT-Angiographie. *Der Ophthalmologe* **113**: 14–22.

Fard MA, Suwan Y, Moghimi S, Geyman LS, Chui TY, Rosen RB & Ritch R (2018): Pattern of peripapillary capillary density loss in ischemic optic neuropathy compared to that in primary open-angle glaucoma. *PLoS ONE* **13**: e0189237.

Gadde SGK, Anegondi N, Bhanushali D, Chidambara L, Yadav NK, Khurana A & Sinha Roy A (2016): Quantification of vessel density in retinal optical coherence tomography angiography images using local fractal dimension. *Investig Ophthalmology Vis Sci* **57**: 246.

Garway-Heath DF, Poinosawmy D, Fitzke FW & Hitchings RA (2000): Mapping the visual field to the optic disc in normal tension glaucoma eyes. *Ophthalmology* **107**: 1809–1815.

Geyman LS, Garg RA, Suwan Y et al. (2017): Peripapillary perfused capillary density in primary open - angle glaucoma across disease stage: An optical coherence tomography angiography study. *Br J Ophthalmol* **101**: 1261–1268.

- Halpern DL & Grosskreutz CL (2002): Glaucomatous optic neuropathy: mechanisms of disease. *Ophthalmol Clin North Am* **15**: 61–68.
- Heijl A, Leske MC, Bengtsson B, Hyman L, Bengtsson B, Hussein M & Early Manifest Glaucoma Trial Group (2002): Reduction of Intraocular Pressure and Glaucoma Progression. *Arch Ophthalmol* **120**: 1268.
- Hemelings R, Elen B, Barbosa-Breda J et al. (2020): Accurate prediction of glaucoma from colour fundus images with a convolutional neural network that relies on active and transfer learning. *Acta Ophthalmol* **98**: e94–e100. <https://doi.org/10.1111/aos.14193>
- Jesus DA, Majewska M, Krzyzanowska-Berkowska P. & Iskander DR (2017): Influence of eye biometrics and corneal microstructure on noncontact tonometry. *PLoS ONE* **12**: e017718.
- Jesus DA, Barbosa Breda J, Van Keer K, Rocha Sousa A, Abegão Pinto L & Stalmans I (2019): Quantitative automated circumpapillary microvascular density measurements: a new angioOCT-based methodology. *Eye* **33**: 320–326.
- Jia Y, Tan O, Tokayer J et al. (2012): Split-spectrum amplitude-decorrelation angiography with optical coherence tomography. *Opt Express* **20**: 4710.
- Jia Y, Wei E, Wang X, et al. (2014): Optical coherence tomography angiography of optic disc perfusion in glaucoma. *Ophthalmology* **121**: 1322–1332.
- Kass MA & Gordon MO (2000): Intraocular pressure and visual field progression in open-angle glaucoma. *Am J Ophthalmol* **130**: 490–491.
- Kim SB, Lee EJ, Han JC & Kee C (2017): Comparison of peripapillary vessel density between preperimetric and perimetric glaucoma evaluated by OCT-angiography. *PLoS ONE* **12**: 1–12.
- Kiyota N, Kunikata H, Shiga Y, Omodaka K & Nakazawa T (2018): Ocular microcirculation measurement with laser speckle flowgraphy and optical coherence tomography angiography in glaucoma. *Acta Ophthalmol* **96**: e485–e492.
- Koustenis A, Harris A, Gross J, Januleviciene I, Shah A & Siesky B (2017): Optical coherence tomography angiography: an overview of the technology and an assessment of applications for clinical research. *Br J Ophthalmol* **101**: 16–20.
- Kromer R, Glusa P, Framme C, Pielan A & Junker B (2019): Optical coherence tomography angiography analysis of macular flow density in glaucoma. *Acta Ophthalmol* **97**: e199–e206.
- Kumar RS, Anegondi N, Chandapura RS, Sudhakaran S, Kadambi SV, Rao HL, Aung T & Roy AS (2016): Discriminant function of optical coherence tomography angiography to determine disease severity in glaucoma. *Investig Ophthalmol Vis Sci* **57**: 6079–6088.
- Kuryshcheva NI, Maslova EV, Zolnikova IV, Fomin AV & Lagutin MB (2018): A comparative study of structural, functional and circulatory parameters in glaucoma diagnostics. *PLoS ONE* **13**: 1–21.
- Kwon J, Choi J, Shin JW, Lee J & Kook MS (2017): Glaucoma diagnostic capabilities of foveal avascular zone parameters using optical coherence tomography angiography according to visual field defect location. *J Glaucoma* **26**: 1120–1129.
- Lin Y, Jiang H, Liu Y, Rosa Gameiro G, Gregori G, Dong C, Rundek T & Wang J (2019): Age-related alterations in retinal tissue perfusion and volumetric vessel density. *Invest Ophthalmol Vis Sci* **60**: 685–693.
- Liu L, Jia Y, Takusagawa HL et al. (2015): Optical coherence tomography angiography of the peripapillary retina in glaucoma. *JAMA Ophthalmol* **133**: 1045.
- Liu C-H, Wu W-C, Sun M-H, Kao L-Y, Lee Y-S & Chen HS-L (2017): Comparison of the retinal microvascular density between open angle glaucoma and nonarteritic anterior ischemic optic neuropathy. *Invest Ophthalmol Vis Sci* **58**: 3350–3356.
- Lommatzsch C, Rothaus K, Koch JM, Heinz C & Grisanti S (2018): OCTA vessel density changes in the macular zone in glaucomatous eyes. *Graefes Arch Clin Exp Ophthalmol* **256**: 1499–1508.
- Mansoori T, Sivaswamy J, Gamalapati JS & Balakrishna N (2017): Radial peripapillary capillary density measurement using optical coherence tomography angiography in early glaucoma. *J Glaucoma* **26**: 438–443.
- Michelson G, Schmauss B, Langhans MJ, Harazny J & Groh MJ (1996): Principle, validity, and reliability of scanning laser Doppler flowmetry. *J Glaucoma* **5**: 99–105.
- Mwanza J, Lee G & Budenz D (2015): Effect of adjusting retinal nerve fiber layer profile to fovea-disc angle axis on the thickness and glaucoma diagnostic performance. *Am J Ophthalmol* **161**: 12–21.
- Nascimento E Silva R, Chiou CA, Wang M et al. (2019): Microvasculature of the optic nerve head and peripapillary region in patients with primary open-angle glaucoma. *J Glaucoma* **28**: 281–288.
- Poli M, Cornut P-L, Nguyen A-M, De Bats F & Denis P (2018): Accuracy of peripapillary versus macular vessel density in diagnosis of early to advanced primary open angle glaucoma. *J Fr Ophtalmol* **41**: 619–629.
- Rabiolo A, Gelormini F, Sacconi R et al. (2018): Comparison of methods to quantify macular and peripapillary vessel density in optical coherence tomography angiography. *PLoS ONE* **13**: 1–20.
- Rao HL, Pradhan ZS, Weinreb RN et al. (2016): regional comparisons of optical coherence tomography angiography vessel density in primary open-angle glaucoma. *Am J Ophthalmol* **171**: 75–83.
- Rao HL, Kadambi SV, Weinreb RN et al. (2017a): Diagnostic ability of peripapillary vessel density measurements of optical coherence tomography angiography in primary open-angle and angle-closure glaucoma. *Br J Ophthalmol* **101**: 1066–1070.
- Rao HL, Pradhan ZS, Weinreb RN et al. (2017b): Vessel density and structural measurements of optical coherence tomography in primary angle closure and primary angle closure glaucoma. *Am J Ophthalmol* **177**: 106–115.
- Rao HL, Pradhan ZS, Weinreb RN et al. (2017c): Optical coherence tomography angiography vessel density measurements in eyes with primary open-angle glaucoma and disc hemorrhage. *J Glaucoma* **26**: 888–895.
- Rao HL, Pradhan ZS, Weinreb RN et al. (2017d): A comparison of the diagnostic ability of vessel density and structural measurements of optical coherence tomography in primary open angle glaucoma. *PLoS ONE* **12**: 1–13.
- Rolle T, Dallorto L, Tavassoli M & Nuzzi R (2019): Diagnostic ability and discriminant values of oct-angiography parameters in early glaucoma diagnosis. *Ophthalmic Res* **61**: 143–152.
- Sánchez Brea L, Andrade De Jesus D, Shirazi MF, Pircher M, van Walsum T & Klein S (2019): Review on retrospective procedures to correct retinal motion artefacts in OCT imaging. *Appl Sci* **9**: 2700.
- Scripsema NK, Garcia PM, Bavier RD et al. (2016): Optical coherence tomography angiography analysis of perfused peripapillary capillaries in primary open-angle glaucoma and normal-tension glaucoma. *Investig Ophthalmology Vis Sci* **57**: OCT611.
- Shin JW, Sung KR, Lee JY, Kwon J & Seong M (2017): Optical coherence tomography angiography vessel density mapping at various retinal layers in healthy and normal tension glaucoma eyes. *Graefes Arch Clin Exp Ophthalmol* **255**: 1193–1202.
- Sousa D, Marques-Neves C, Kayat K & Barbosa-Breda J (2019): Optical coherence tomography angiography quantitative assessment of choriocapillaris blood flow in central serous chorioretinopathy. *Am J Ophthalmol* **200**: 250.
- Spaide RF, Klancnik JM & Cooney MJ (2015): Retinal vascular layers imaged by fluorescein angiography and optical coherence tomography angiography. *JAMA Ophthalmol* **133**: 45.
- Spaide RF, Fujimoto JG, Waheed NK & Science C (2016): Image artifacts in optical coherence angiography. *Retina* **35**: 2163–2180.
- Stalmans I, Vandewalle E, Anderson DR et al. (2011): Use of colour Doppler imaging in ocular blood flow research. *Acta Ophthalmol* **89**: 609–630.
- Stanga PE, Tsamis E, Papayannis A, Stringa F, Cole T & Jalil A (2016): Swept-source optical coherence tomography Angio™ (Topcon Corp, Japan): technology review. *Dev Ophthalmol* **56**: 13–17.
- Sugiyama T, Araie M, Riva CE, Schmetterer L & Orgul S (2010): Use of laser speckle flowgraphy in ocular blood flow research. *Acta Ophthalmol* **88**: 723–729.
- Takusagawa H, Liang L, Ma KN et al. (2017): Projection-resolved optical coherence tomography angiography of macular retinal circulation in glaucoma. *Ophthalmology* **124**: 1589–1599.
- Triolo G, Rabiolo A, Shemonski ND et al. (2017): Optical coherence tomography angiography macular and peripapillary

vessel perfusion density in healthy subjects, glaucoma suspects, and glaucoma patients. *Investig Ophthalmol Vis Sci* **58**: 5713–5722.

Van Melkebeke L, Barbosa-Breda J, Huygens M & Stalmans I (2018): Optical coherence tomography angiography in glaucoma: a review. *Ophthalmic Res* **60**: 139–151.

Xu J, Wong K, Jian Y & Sarunic MV (2014): Real-time acquisition and display of flow contrast using speckle variance optical coherence tomography in a graphics processing unit. *J Biomed Opt* **19**: 026001.

Xu H, Zhai R, Zong Y, Kong X, Jiang C, Sun X, He Y & Li X (2018): Comparison of retinal microvascular changes in eyes with high-tension glaucoma or normal-tension glaucoma: a quantitative optic coherence tomography angiographic study. *Graefes Arch Clin Exp Ophthalmol* **256**: 1179–1186.

Yarmohammadi A, Zangwill LM, Diniz-Filho A et al. (2016a): Optical coherence tomography angiography vessel density in healthy, glaucoma suspect, and glaucoma eyes. *Investig Ophthalmology Vis Sci* **57**: OCT451.

Yarmohammadi A, Zangwill LM, Diniz-Filho A et al. (2016b): Relationship between optical coherence tomography angiography vessel density and severity of visual field loss in glaucoma. *Ophthalmology* **123**: 2498–2508.

Yarmohammadi A, Zangwill LM, Diniz-Filho A et al. (2017): Peripapillary and macular vessel density in patients with glaucoma and single-hemifield visual field defect. *Ophthalmology* **124**: 709–719.

Yip VCH, Wong HT, Yong VKY et al. (2019): Optical coherence tomography angiography of optic disc and macula vessel density in glaucoma and healthy eyes. *J Glaucoma* **28**: 80–87.

Zhang A & Wang RK (2015): Feature space optical coherence tomography based microangiography. *Biomed Opt Express* **6**: 1919.

Zhu L, Zong Y, Yu J, Jiang C, He Y, Jia Y, Huang D & Sun X (2018): reduced retinal vessel density in primary angle closure glaucoma: a quantitative study using optical coherence tomography angiography. *J Glaucoma* **27**: 322–327.

Zivkovic M, Dayanir V, Kocaturk T et al. (2017): Foveal avascular zone in normal tension glaucoma measured by optical coherence tomography angiography. *Biomed Res Int* **2017**: 1–7.

Appendix A

Search query used in the PubMed database:

("Glaucoma*" [Mesh] OR "Glaucoma*" [tiab]) AND ("optical coherence tomography angiography" [tiab] OR "OCTA" [tiab] OR "OCT-A" [tiab] OR "OCT angiography" [tiab] OR "optical coherence tomography based microangiography" [All Fields] OR "angio-OCT" [tiab] OR "OCT-angio" [tiab]) AND ("Glaucoma/diagnostic imaging" [Mesh] OR "Glaucoma/diagnosis" [Mesh] OR "Glaucoma/analysis" [Mesh] OR "Image Analysis" [tiab] OR "Image Processing" [tiab] OR "Image Enhancement" [Mesh] OR "Image Enhancement" [tiab] OR "Image processing, Computer

Assisted" [Mesh] OR "Image processing, Computer Assisted" [tiab] OR "Computer-Assisted Image Processing" [Mesh] OR "Computer-Assisted Image Processing" [tiab] OR "Computer Assisted Image Processing" [Mesh] OR "Computer Assisted Image Processing" [tiab] OR "Image Reconstruction" [Mesh] OR "Image Reconstruction" [tiab] OR "Image Reconstructions" [Mesh] OR "Image Reconstructions" [tiab] OR "Reconstruction, Image" [Mesh] OR "Reconstruction, Image" [tiab] OR "Reconstructions, Image" [Mesh] OR "Reconstructions, Image" [tiab] OR "Image Analysis, Computer-Assisted" [tiab] OR "Image Analysis, Computer Assisted" [Mesh] OR "Image Analysis, Computer Assisted" [tiab] OR "Computer-Assisted Image Analysis" [Mesh] OR "Computer-Assisted Image Analysis" [tiab] OR "Computer Assisted Image Analysis" [Mesh] OR "Computer Assisted Image Analysis" [tiab] OR "Analysis, Computer-Assisted Image" [Mesh] OR "Analysis, Computer-Assisted Image" [tiab] OR "Computer-Assisted Image Analyses" [Mesh] OR "Computer-Assisted Image Analyses" [tiab] OR "Image Analyses, Computer-Assisted" [Mesh] OR "Image Analyses, Computer-Assisted" [tiab] OR "Algorithm*" [tiab] OR "microvascular density" [tiab])

Appendix B

Table B1. Characteristics of included studies. The publications in bold font were qualitatively assessed in this review.

Author	Number of patients/eyes	Eyes in control group (%)	Eyes in glaucoma group (%)	Age (years) (p-value)	Type of glaucoma	OCT device	OCT light-source wavelength (nm)	Image quality cut-off value	Field of view (mm)
Akil et al. (2017)	56/56	16 (28.6)	20 mild POAG (35.7) 20 PPG (35.7)	62.2 versus 65.38 versus 63.13 p = 0.7	Mild POAG, PPG	DRI OCT, Triton, TOPCON	1050	NA	3 × 3
Alnawaiseh et al. (2018)	36/69	34 (49.3)	35 (50.7)	62 versus 63.09 p = 0.661	OAG	AngioVue	840	<50	3 × 3 pf 4.5 × 4.5 OD 6 × 6
Bojtkian et al. (2016)	89/89	28 (31.5)	61 (68.5)	68.8 versus 66.2 versus 64.6 p = 0.38	POAG, NTG	Cirrus-HD-OCT-5000, Zeiss AngioPlex AngioVue	840	<6	
Cennamo et al. (2017)	58/86	48 (55.8)	38 (44.2)	59.20 versus 65.05 p = 0.119	PPOAG	AngioVue	840	<50	7 × 7
Chen et al. (2016)	88/88	20 (22.7)	26 GS (29.5) 21 POAG (23.9) 21 NTG (23.9)	68.3 versus 68 versus 65.7 p = 0.51	GS, POAG, NTG	Cirrus-HD-OCT-5000, Zeiss AngioPlex AngioVue	840	<6	6 × 6
Chen et al. (2017)	53/53	27 (50.9)	26 (49.1)	57 versus 57 p = 0.84	POAG	AngioVue	840	<45	4.5 × 4.5 pp 6 × 6 macula 4.5 × 4.5
Chihara et al. (2017)	105/105	25 (23.8)	66 (76.2)	56.2 versus 60.4 p = 0.258	POAG	AngioVue	840	<40	
Chung et al. (2017)	253/253	113 (44.7)	80 Early (31.6) 35 Moderate (13.8) 25 Severe (9.9)	49.8 versus 50.5 versus 52.9 versus 54.5 p = 0.133	Early, moderate, severe	AngioVue	840	<50	4.5 × 4.5
Fard et al. (2018)	78/125	80 (64.0)	45 (36.0)	48.4 versus 60.2 p = 0.08	POAG	AngioVue	840	<40	4.5 × 4.5
Geyman et al. (2017)	84/84	24 (28.6)	22 Mild (26.2) 20 Moderate (23.8) 18 Severe (21.4)	52 versus 66 versus 63 versus 62 p < 0.001	Mild, moderate, severe POAG	AngioVue	840	<40	4.5 × 4.5
Jesus et al. (2018)	122/122	40 (32.8)	82 (67.2)	63 versus 66 p-value NA	PAOG, NTG	Cirrus-HD-OCT, Zeiss AngioPlex	840	<6	3 × 3
Jia y et al. (2014)	35/35	24 (68.6)	8 PG (22.9) 3 PPG (8.6)	52 versus 68 p = 0.000	PG, PPG	Ultrahigh speed swept-source OCT imaging device AngioVue	1050	NA	3 × 3
Kim et al. (2000)	22/44	9 (20.5)	13 (79.5)	35 versus 55.3 p < 0.001	PNTG	AngioVue	840	<48	4.5 × 4.5
Kiyota et al. (2017)	102/102	20 (19.6)	82 (80.4) - 28 Mild (34.1) - 25 Moderate (30.5) - 29 Severe (35.4)	59 versus 60 versus 61 versus 61 p = 0.79	OAG, mild, moderate, severe	Swept-source (SS-OCT) Angio (Topcon)	1000	NA	4.5 × 4.5
Kromer et al. (2017)	51/51	21 (41.2)	30 (58.8)	70.3 versus 72.6 p = 0.298	POAG	SPECTRALIS, Heidelberg Engineering	870	NA	5 × 3.5

Table B1. (Continued)

Author	Number of patients/eyes	Eyes in control group (%)	Eyes in glaucoma group (%)	Age (years) (p-value)	Type of glaucoma	OCT device	OCT light-source wavelength (nm)	Image quality cut-off value	Field of view (mm)
Kumar et al. (2019)	183/273	74 (27.1)	93 POAG (34.1) 70 PACG (25.6) Of which: - 83 Early (30.4) - 43 Moderate (15.8) - 45 Severe (16.5) 28 PPG (10.3) 48 Early POAG (38.4) 42 Moderate to severe POAG (33.6)	58.2 versus 57.04 versus 61.2 versus 62.6 versus 61.1 p < 0.001	PPG, early, moderate, severe	AngioVue	840	<40	4.5 × 4.5
Kuryshva et al. (2016)	125/125	35 (28.0)	48 Early POAG (38.4) 42 Moderate to severe POAG (33.6)	62.4 versus 53.7 versus 65.1 age-matched	Early POAG, moderate to severe POAG	AngioVue	840	<50	4.5 × 4.5 OD 6 × 6 macula
Kwon et al. (2018)	125/125	45 (36.0)	45 PVFD (36.0) 35 CVFD (28.0)	49 versus 48 versus 51 p = 0.644	OAG, with PVFD, OAG with CVFD	Cirrus-HD-OCT, Zeiss AngioPlex	840	<8	3 × 3
Lommatzsch et al. (2017)	115/115	50 (43.5)	85 (56.5)	62 versus 61 p = 0.45	POAG, PEG, NTG, AAG	AngioVue	840	<40	6 × 6
Lommatzsch et al. (2017)	54/54	24 (44.4)	30 (55.6)	52 versus 62 p = 0.02	POAG, PEG, NTG, AAG	Cirrus-HD-OCT, Zeiss AngioPlex	840	<8	6 × 6
Liang et al. (2015)	24/24	12 (50.0)	12 (50.0)	67 versus 70 age-matched	PG, PPG	AngioVue	840	<40	3 × 3
Liu et al. (2015)	NA/43	27 (62.8)	16 (37.2)	55.8 versus 53.8 p = 0.46	OAG	AngioVue	840	<45	4.5 × 4.5 pp 6 × 6 macula
Mansoori et al. (2018)	76/76	52 (68.4)	24 (31.5)	49.89 versus 52.12 p = 0.57	POAG	AngioVue	840	<60	4.5 × 4.5
Poli et al. (2019)	36/52	15 (28.8)	15 PPG (28.8) 6 Mild (11.5) 16 Moderate to advanced (30.8)	39 versus 60 versus 63 versus 74 p = 0.002	PPG, POAG, mild moderate to advanced	AngioVue	840	<40	6 × 6 FAZ 4.5 × 4.5 OD
Rabiolo et al. (2018)	53/88	27 (30.7)	26 (69.3)	47.7 versus 60.5 p = 0.008	POAG	SS-OCTA PLEX Elite 9000, Zeiss	1040-1060	<7	6 × 6
Rao et al. (2018)	92/142	78 (54.9)	64 (45.1)	58 versus 66 p = 0.01	POAG	AngioVue	840	<35	4.5 × 4.5 OD 3 × 3 macula 4.5 × 4.5
Rao et al. (2016)	104/160	48 (30.0)	63 POAG (39.4) 49 PACG (30.7)	52 versus 65 versus 64 p < 0.001 and p = 0.002	POAG, PACG	AngioVue	840	<40	4.5 × 4.5 OD 3 × 3 macula
Rao et al. (2017b)	122/163	66 (40.5)	34 POAG with DH (20.9) 63 POAG without DH (38.7)	59.7 versus 65.6 versus 66.0 p = 0.93	POAG with DH, POAG without DH	AngioVue	840	<45	4.5 × 4.5 OD 3 × 3 macula
Rao et al. (2017c)	117/195	78 (40.0)	117 (60)	60.7 versus 62.8 p = 0.3	POAG	AngioVue	840	<35	4.5 × 4.5 OD 3 × 3 macula
Rao et al. (2017a)	117/173	77 (44.5)	31 PAC (17.9) 65 PACG (37.6)	60.7 versus 60.3 versus 62.0 p = 0.86 and p = 0.44	PAC, PACG	AngioVue	840	<35	4.5 × 4.5 OD 3 × 3 macula
Rolle et al. (2017d)	NA/71	13 (18.3)	39 PPG (54.9) 19 POAG (26.8)	60.0 versus 63.4 versus 64.5 p = 0.24	PPG, POAG	AngioVue	840	<50	3 × 3

Table B1. (Continued)

Author	Number of patients/eyes	Eyes in control group (%)	Eyes in glaucoma group (%)	Age (years) (p-value)	Type of glaucoma	OCT device	OCT light-source wavelength (nm)	Image quality cut-off value	Field of view (mm)
Scripsema et al. (2019)	92/92	26 (28.3)	40 POAG (43.5) 26 NTG (28.3)	63.72 versus 66.3 versus 67.92 p = 0.34	POAG, NTG	AngioVue	840	NA	3.5 × 3.5 4.5 × 4.5
Shin et al. (2016)	93/93	51 (54.8)	42 (45.2)	50.7 versus 51.8 p = 0.642	NTG	Cirrus-HD-OCT, Zeiss AngioPlex	840	NA	6 × 6
Silva et al. (2019)	46/46	16 (34.8)	30 (65.7)	67.9 versus 65.4 p = 0.37	POAG	DRI-OCT, Triton, Topcon	1050	<48	3 × 3
Takusagawa et al. (2010)	60/60	30 (50.0)	30 (50.0)	65 versus 65 age-matched	PG	AngioVue	840	<50	6 × 6
Triolo et al. (2017)	120/120	40 (30.0)	40 GS (30.0) 40 POAG (30.0)	60.9 versus 61.7 versus 63.6 p = 0.38	GS, POAG	SS-OCTA PLEX Elite 9000, Zeiss	1040–1060	<7	6 × 6
Xu et al. (2014)	127/127	51 (40.2)	43 HTG (33.9) 33 NTG (26.0)	48.96 versus 50.91 versus 52.8 p = 0.467 and p > 0.99	HTG, NTG	AngioVue	840	<60	6 × 6 F 4.5 × 4.5 OD
Yarmohammadi et al. (2016a)	164/261	23 (14.0)*	37 GS (14.2)* 104 OAG (39.8)*	53.5 versus 68.2 versus 72.4 p < 0.001	GS, OAG	AngioVue	840	<48	4.5 × 4.5
(Yarmohammadi et al. 2016a)	152/152	26 (17.1)	52 GS (34.2) Mild 46 (30.3) Moderate and severe 28 (18.4)	55.6 versus 68.7 versus 72.9 versus 75.7 p < 0.001	GS, POAG, mild, moderate and severe	AngioVue	840	<48	4.5 × 4.5
Yarmohammadi et al. (2016b)	86/86	28 (32.6)	58 (67.4)	69.91 versus 71.37 p = 0.190	OAG	AngioVue	840	<48	4.5 × 4.5 pp 3 × 3 macula
Yip et al. (2017)	53/90	58 (64.4)	15 POAG (16.7) 14 NTG (15.6) 1 juvenile OAG (1.1)	51.17 versus 57.5 p = 0.051	POAG, NTG, CAG	AngioVue	840	<40	3 × 3
Zhu et al. (2015)	44/78	39 (50.0)	2 CAG (2.2) 39 (50.0)	57.83 versus 59.93 p = 0.095	PACG	AngioVue	840	<40	4.5 × 4.5
Zivkovic et al. (2018)	51/51	21 (41.2)	30 (58.8)	69.5 versus 70.1 p = 0.70	NTG	Cirrus-HD-OCT, Zeiss AngioPlex	840	<6	3 × 3 FAZ

CVFD = central visual field defect, F = fovea, FAZ = foveal avascular zone, GS = glaucoma suspect, NA = not available, that is, not mentioned in the article, NTG = normal tension glaucoma, OAG = open-angle glaucoma, OD = optic disc, P = perimetric, PACG = primary angle closure glaucoma, pf = parafoveal, POAG = primary open-angle glaucoma, pp = peripapillary, PP = peripapillary, PVFD = peripheral visual field defect.

In the work published by Lommatzsch et al. (2017), two studies were performed. In each study, a different number of eyes were examined using one of two devices (Optovue or the Zeiss Cirrus).

Appendix C

Table C1. Vessel density values (%) for healthy control group (bold) versus glaucoma patients according to the ocular layer, region and sector. Some studies provided multiple values for the same area due to comparison between approaches or different glaucoma types. The radial peripapillary capillary plexus is shown here as sublayer of the superficial capillary plexus.

Sector	Region		
	Optic Disc	Macular Region	Peripapillary region
1.a Radial peripapillary capillary plexus			
Whole image	14.6 versus 5.3 (Yip et al. 2017)		61.9 versus 54.4 (Rao et al. 2017c)*
	51.8 versus 47.3 (Rabiolo et al., 2018)		47 versus 40 versus 46 (Chen et al. 2016)
	49.3 versus 46.7 (Rabiolo et al. 2018)		63.62 versus 56.15 (Zhu et al. 2015)
	49.5 versus 43.4 (Rabiolo et al. 2018)		48.03 versus 46.09 versus 45.34 versus 42.36 (Chung et al. 2017)
	48.1 versus 41.9 (Rabiolo et al. 2018)		62 versus 55 (Rao et al. 2016)
	49.7 versus 37.1 (Rabiolo et al. 2018)		53.3 versus 43.8 (Chen et al. 2017)
	40.8 versus 39.3 (Rabiolo et al. 2018)		40.7 versus 37.9 (Nascimento et al., 2019)
	55.42 versus 50.58 (Alnawaiseh et al. 2018)†		63.07 versus 59.99 (Alnawaiseh et al. 2018)*
	53 versus 54 (Triolo et al. 2017)		60.8 versus 52.5 versus 52.7 (Rao et al. 2018)*
	53 versus 50 (Triolo et al. 2017)		58.0 versus 56.9 versus 46.3 (Kuryшева et al. 2016)
	55.8 versus 52.9 versus 43.1 (Kuryшева et al. 2016)		61.9 versus 53.3 (Rao et al. 2017a)*
	53.9 versus 45.2 (Rao et al. 2017c)*		65.28 versus 60.34 (Rolle et al. 2017d)
	58.8 versus 55.4 versus 48.6 versus 43.1 versus 36.5 (Kumar et al. 2019)		93 versus 80.55 (Liu et al. 2019)
	54.4 versus 50.4 (Rao et al. 2016)		49.8 versus 38.2 versus 42.1 (Chihara et al. 2017)
	53.9 versus 45.1 (Rao et al. 2017a)*		86.6 versus 86.83 versus 80.03 (Akil et al. 2017)
	58.09 versus 54.01 (Alnawaiseh et al. 2018)†		42.2 versus 35.1 versus 30.8 versus 28.3 (Geyman et al. 2017)‡
	86.6 versus 78.04 versus 70.1 (Akil et al. 2017)		42.3 versus 30.2 (Fard et al. 2018)‡
			53.01 versus 44.73 (Liu et al. 2015)
			56.7 versus 51.6 (Yarmohammadi et al. 2016a)
			56.7 versus 48.3 (Yarmohammadi et al. 2016a)
			56.7 versus 41.7 (Yarmohammadi et al. 2016a)
			41.32 versus 36.49 (Scripsema et al. 2019)‡
			41.32 versus 33.13 (Scripsema et al. 2019)‡
			41.88 versus 36.48 (Scripsema et al. 2019)‡
			41.88 versus 33.31 (Scripsema et al. 2019)‡
			55.70 versus 51.87 (Poli et al. 2019)
			55.70 versus 46.21 (Poli et al. 2019)
			55.70 versus 37.45 (Poli et al. 2019)

Table C1. (Continued)

Sector	Region		
	Optic Disc	Macular Region	Peripapillary region
Inside Disc	<p>20.9 versus 11.0 (Yip et al. 2017)</p> <p>55.45 versus 49.09 (Rolle et al. 2017d)</p> <p>48.1 versus 40.1 (Rao et al. 2017c)*</p> <p>48.1 versus 46.5 (Rao et al. 2017a)*</p> <p>46,78 versus 34.07 (Alnawaiseh et al. 2018)*</p> <p>47.4 versus 40.2 (Rao et al. 2016)</p> <p>77 versus 68 versus 70 (Bojikian et al. 2016)</p> <p>50.2 versus 45.3 versus 37.7 (Kuryshcheva et al. 2016)</p> <p>37.9 versus 39.8 versus 29.8 versus 29.8 versus 26.7 (Kumar et al. 2019)</p> <p>Not significant (Nascimento et al., 2019; Rolle et al. 2017d)</p>		<p>not significant (Rao et al. 2017b; Poli et al. 2019; Rolle et al. 2017d)</p>
Inferior	<p>23.1 versus 13.3 (Yip et al. 2017)</p> <p>64.7 versus 53.9 (Rao et al. 2017c)*</p> <p>48.7 versus 43.7 (Rao et al. 2017a)*</p>		<p>64.7 versus 53.9 (Rao et al. 2017c)*</p> <p>46 versus 37 versus 46 (Chen et al. 2016)</p> <p>63.26 versus 59.72 (Zhu et al. 2015)</p> <p>44.3 versus 36.7 versus 30.5 versus 24.7 (Geyman et al. 2017)‡</p> <p>43.9 versus 36.1 (Fard et al. 2018)‡</p>
Superior	<p>21.4 versus 12.7 (Yip et al. 2017)</p> <p>49.1 versus 40.8 (Rao et al. 2017c)*</p> <p>49.1 versus 49.5 (Rao et al. 2017a)*</p>		<p>63.6 versus 55.2 (Rao et al. 2017c)*</p> <p>45 versus 39 versus 46 (Chen et al. 2016)</p> <p>43.6 versus 35.2 versus 30.3 versus 26.3 (Geyman et al. 2017)‡</p> <p>43.7 versus 29.1 (Fard et al. 2018)‡</p>
Nasal superior			<p>63.26 versus 59.72 (Zhu et al. 2015)</p> <p>63.91 versus 60.71 versus 57.84 versus 49.42 (Chung et al. 2017)</p> <p>62.9 versus 55.3 (Rao et al. 2016)</p> <p>59.1 versus 51.8 versus 53.0 (Rao et al. 2018)*</p> <p>55.6 versus 54.7 versus 44.8 (Kuryshcheva et al. 2016)</p> <p>59.4 versus 51.8 (Rao et al. 2017a)*</p>
Nasal inferior			<p>not significant (Mansoori et al. 2018; Rao et al. 2017b)‡</p> <p>64.40 versus 55.85 (Zhu et al. 2015)</p> <p>65.80 versus 60.94 versus 59.11 versus 45.07 (Chung et al. 2017)</p>

Table C1. (Continued)

Sector	Region		
	Optic Disc	Macular Region	Peripapillary region
Nasal			63.4 versus 55.2 (Rao et al. 2016)
			62.0 versus 53.8 versus 53.1 (Rao et al. 2018)*
			62.1 versus 59.2 versus 54.4 (Rao et al. 2017b)
			55.8 versus 57.6 versus 45.0 (Kurysheva et al. 2016)
			61.2 versus 54.3 (Rao et al. 2017a)*
			not significant (Mansoori et al. 2018)‡
		17.3 versus 8.2 (Yip et al. 2017)	59.5 versus 53.2 (Rao et al. 2017c)*
		48.9 versus 41.2 (Rao et al. 2017c)*	61.39 versus 54.61 (Zhu et al. 2015)
		48.9 versus 44.2 (Rao et al. 2017a)*	60.23 versus 59.21 versus 55.80 versus 48.77 (Chung et al. 2017)
			59.7 versus 56 (Rao et al. 2016)
Temporal superior			58.0 versus 51.2 versus 51.9 (Rao et al. 2018)*
			56.9 versus 54.8 versus 46.5 (Kurysheva et al. 2016)
			42.8 versus 35.5 versus 33.2 versus 29.8 (Geyman et al. 2017)‡
			43.3 versus 32.2 (Fard et al. 2018)‡
			not significant (Chen et al. 2016; Rao et al. 2017b)
			66.25 versus 58.36 (Zhu et al. 2015)
			68.82 versus 64.15 versus 56.55 versus 43.31 (Chung et al. 2017)
			66.6 versus 59.4 (Rao et al. 2016)
			63.3 versus 56.5 versus 53.5 (Rao et al. 2018)*
			60.4 versus 59.0 versus 49.2 (Kurysheva et al. 2016)
Temporal inferior			64.9 versus 56.9 (Rao et al. 2017a)*
			20.2 versus 10.8 (Mansoori et al. 2018)‡
			not significant (Rao et al. 2017b)
			66.95 versus 60.11 (Zhu et al. 2015)
			68.52 versus 64.93 versus 61.04 versus 49.15 (Chung et al. 2017)
			66 versus 55.5 (Rao et al. 2016)
			64.6 versus 49.7 versus 52.8 (Rao et al. 2018)*
			60.5 versus 59.8 versus 44.4 (Kurysheva et al. 2016)
			20.7 versus 13.4 (Mansoori et al. 2018)‡
			63.0 versus 51.8 (Rao et al. 2017a)*

Table C1. (Continued)

Sector	Region		
	Optic Disc	Macular Region	Peripapillary region
Temporal	17.3 versus 11.0 (Yip et al. 2017) 44.3 versus 34.9 (Rao et al. 2017c)* 44.3 versus 36.9 (Rao et al. 2017a)*		not significant (Rao et al. 2017b) 60.3 versus 56.5 (Rao et al. 2017c)* 68.83 versus 64.16 versus 62.13 versus 50.55 (Chung et al. 2017) 60.4 versus 57 (Rao et al. 2016) 59.0 versus 55.0 versus 53.6 (Rao et al. 2018)* 59.4 versus 58 versus 47 (Kuryshva et al. 2016) 45.2 versus 41.3 versus 37.5 versus 36.2 (Geyman et al. 2017)‡ 44.6 versus 28.2 (Fard et al. 2018)‡
Circumpapillary Whole image			not significant (Rao et al. 2017b) 34 versus 24 (Jesus et al. 2017)* 61.5 versus 53.3 (Chen et al. 2017) 51.9 versus 39.5 versus 34.4 (Kwon et al. 2018) 62.4 versus 59 versus 54.7 (Yarmohammadi et al. 2016b) 64.2 versus 60.3 versus 55.1 (Yarmohammadi et al. 2018) 65.1 versus 57.5 (Yarmohammadi et al. 2016a) 65.1 versus 49.6 (Yarmohammadi et al. 2016a) 42.45 versus 37.20 (Scripsema et al. 2019)‡ 42.45 versus 33.40 (Scripsema et al. 2019)‡ 42.99 versus 37.75 (Scripsema et al. 2019)‡ 42.99 versus 34.24 (Scripsema et al. 2019)‡
Temporal			not significant (Yarmohammadi et al. 2016a) 30 versus 23 (Jesus et al. 2017)* 62.1 versus 55.3 (Chen et al. 2017)
Temporal inferior			not significant (Liu et al. 2015) 66.4 versus 46.2 (Chen et al. 2017) 67.29 versus 46.39 (Liu et al. 2017)
Temporal superior			66.0 versus 55.9 (Chen et al. 2017) 64.78 versus 54.47 (Liu et al. 2015)
Superior			34 versus 23 (Jesus et al. 2017)* 62.01 versus 55.37 (Liu et al. 2015)
Nasal			30 versus 24 (Jesus et al. 2017)* not significant (Chen et al. 2017; Liu et al. 2015)
Nasal inferior			63.6 versus 51.8 (Chen et al. 2017)

Table C1. (Continued)

Sector	Region		
	Optic Disc	Macular Region	Peripapillary region
Nasal superior			62.23 versus 54.66 (Liu et al. 2015)
Inferior			59.2 versus 54.7 (Chen et al. 2017) not significant (Liu et al. 2015) 38 versus 23 (Jesus et al. 2017)* 64.76 versus 50.53 (Liu et al. 2015)
1. Superficial capillary plexus			
Whole image	Not significant (Nascimento et al., 2019)	46.89 versus 42.31 (Lommatzsch et al. 2017) 44 versus 40 (Lommatzsch et al. 2017) 43.2 versus 38.5 (Chen et al. 2017) 46.21 versus 39.55 (Zivkovic et al. 2018) 52.51 versus 48.89 (Alnawaiseh et al. 2018)* 58.02 versus 52.37 (Rolle et al. 2017d) 58.02 versus 49.09 (Rolle et al. 2017d) 48.3 versus 44.7 (Rao et al. 2016)* 49.8 versus 47.5 (Rao et al. 2016) 11.5 versus 8.8 (Yip et al. 2017) 37.3 versus 31.8 (Kromer et al. 2017) 60.5 versus 47.2 (Takusagawa et al. 2010) 53.77 versus 51.63 (Alnawaiseh et al. 2018)* 50.7 versus 45.9 versus 42.2 (Kurysheva et al. 2016) 49.5 versus 46.9 (Rao et al. 2017c)* 53.8 versus 51.1 versus 48.3 (Yarmohammadi et al. 2016b) 50.83 versus 46.99 (Zhu et al. 2015) 52.27 versus 50.83 versus 47.15 versus 45.46 (Chung et al. 2017) 49.5 versus 46.6 (Rao et al. 2017a)* 43.11 versus 38.46 (Liu et al. 2015) not significant (Chen et al. 2017; Triolo et al. 2017)	56.6 versus 51.3 versus 46.2 (Yarmohammadi et al. 2018)
Superior		50.3 versus 47.5 (Rao et al. 2017c)* 51.60 versus 47.62 (Zhu et al. 2015) 49.2 versus 49.7 (Rao et al. 2017a)* 52.65 versus 51.60 versus 47.64 versus 45.74 (Chung et al. 2017) 50.8 versus 47.6 (Rao et al. 2016) 51.60 versus 47.62 (Zhu et al. 2015) 53.2 versus 47.3 versus 47.4 (Kurysheva et al. 2016) not significant (Liu et al. 2015; Triolo et al. 2017; Kromer et al. 2017)	
Nasal		48.3 versus 45.9 (Rao et al. 2017c)* 48.5 versus 47 (Rao et al. 2016) 50.57 versus 45.99 (Zhu et al. 2015) 51.2 versus 46.8 versus 46.3 (Kurysheva et al. 2016) 52.24 versus 50.58 versus 47.87 versus 46.85 (Chung et al. 2017) 56.2 versus 43.58 (Kromer et al. 2017) not significant (Chen et al. 2017; Liu et al. 2015)	
Temporal		49.6 versus 46.8 (Rao et al. 2017c)* 52.64 versus 51.13 versus 47.36 versus 46.05 (Chung et al. 2017) 59.7 versus 56.0 (Rao et al. 2016) 49.8 versus 47.3 (Rao et al. 2016) 51.95 versus 48.18 (Zhu et al. 2015) 52.4 versus 47.2 versus 46.1 (Kurysheva et al. 2016) 43.06 versus 39.71 (Liu et al. 2015) not significant (Chen et al. 2017; Kromer et al. 2017)	
Inferior		51.0 versus 46.6 (Rao et al. 2017c)* 49.3 versus 48.9 (Rao et al. 2017a)*	

Table C1. (Continued)

Sector	Region		
	Optic Disc	Macular Region	Peripapillary region
		51.38 versus 50.29 versus 46.17 versus 44.11 (Chung et al. 2017) 60.4 versus 57 (Rao et al. 2016) 48.5 versus 47 (Rao et al. 2016) 50.57 versus 45.99 (Zhu et al. 2015) 51.2 versus 46.8 versus 46.3 (Kuryшева et al. 2016) not significant (Chen et al. 2017; Liu et al. 2015; Triolo et al. 2017; Kromer et al. 2017) 36.4 versus 32.1 versus 30.8 (Kuryшева et al. 2016) 53.61 versus 47.23 (Poli et al. 2019) 53.61 versus 41.98 (Poli et al. 2019) 53.61 versus 39.89 (Poli et al. 2019) not significant (Chen et al. 2017; Alnawaiseh et al. 2018; Lommatzsch et al. 2017; Poli et al. 2019; Kromer et al. 2017)*	
Fovea			
2. Deep capillary plexus			
Whole image	44.1 versus 39.4 (Nascimento et al., 2019)	53.28 versus 48.23 (Lommatzsch et al. 2017) 56.97 versus 54.95 (Alnawaiseh et al. 2018)* 23.3 versus 13.6 (Yip et al. 2017) 30.6 versus 30.2 (Kromer et al. 2017) 57.3 versus 52.2 versus 46.6 (Kuryшева et al. 2016) not significant (Takusagawa et al. 2010) 62.1 versus 57 versus 54.1 (Kuryшева et al. 2016) not significant (Kromer et al. 2017)	59.93 versus 57.07 (Alnawaiseh et al. 2018)*
Superior		60 versus 55.8 versus 54.1 (Kuryшева et al. 2016) not significant (Kromer et al. 2017)	
Temporal		59.3 versus 56 versus 52 (Kuryшева et al. 2016) not significant (Kromer et al. 2017)	
Inferior		59.6 versus 56.3 versus 50.7 (Kuryшева et al. 2016) not significant (Kromer et al. 2017)	
Nasal		33.3 versus 30.11 (Lommatzsch et al. 2017) 29.45 versus 31.75 (Alnawaiseh et al. 2018)* 35.4 versus 32.5 versus 26.3 (Kuryшева et al. 2016) not significant (Kromer et al. 2017)	
Fovea			
3. Choriocapillaris			
Whole image		22.5 versus 15.3 (Yip et al. 2017)	
Inferior		22.2 versus 15.8 (Yip et al. 2017)	
Superior		22.8 versus 15.1 (Yip et al. 2017)	
Nasal		22.9 versus 15.3 (Yip et al. 2017)	
Temporal		24.3 versus 15.1 (Yip et al. 2017)	
4. Choroid			
Whole image	22.8 versus 16.5 (Yip et al. 2017)		63.2 versus 64.2 versus 57.9 (Kim et al. 2000) 68.4 versus 64.7 versus 48.37 (Kim et al. 2000) not significant (Kim et al. 2000)
Temporal inferior			
Temporal superior			
Superior	26 versus 17.8 (Yip et al. 2017)		
Nasal	23.6 versus 17.3 (Yip et al. 2017)		
Temporal	19.2 versus 13.1 (Yip et al. 2017)		
Inferior	Not significant (Yip et al. 2017)		
5. Whole retina			
Whole image		51.58 versus 47.27 versus 45.56 (Xu et al. 2014) [§]	63.29 versus 55.57 versus 49.78 (Xu et al. 2014) [§]

All values are in mean, except when specified differently. Value were considered as significant at a cut-off of <0.05

* Values given in median.

† Whole enface.

‡ Large vessels extracted, capillary density.

§ Not further specified which layer was used.

Appendix D

Table D1. Qualitative assessment for the studies with an AUROC value above the threshold of 0.77.

Author	Age	Eye	Types and glaucoma severity (VF MD, dB)	OCT specifications	Image Quality cut-off	Fovea-Disc axis
Akil et al. (2017)	No	No	POAG: 2.8 ± 1.9 PPG: 1.65 ± 2.25	1050	NA*	Yes
Alnawaiseh et al. (2017)	No	Yes	$-2.74 (-5.17, -1.25)^\dagger$	840	<50	No, NA
Cennamo et al. (2017)	No	Yes	0.15 ± 1.17	840	<50	No, NA
Chen et al. (2017)	No	No	-8.8 ± 6.2	840	<45	Yes
Chung et al. (2017)	No	No	Early: -2.42 ± 1.69 Moderate: -8.35 ± 1.77 Severe: -20.12 ± 5.31	840	<50	Yes
Geyman et al. (2017)	Yes, corrected	No	Mild: -3.2 ± 1.7 Moderate: -8.0 ± 2.0 Severe: -21.4 ± 7.1	840	<40	Yes
Jesus et al. (2019)	No	No	-7.8 ± 6.5	840	<6[‡]	No
Kuryshcheva et al. (2016)	No	No	Early POAG: -2.1 ± 3.4 Moderate/Severe POAG: -11.8 ± 6.1	840	<50	Yes
Kwon et al. (2018)	No	No	PVFD: -4 ± 4 CVFD: -3.2 ± 2.2	840	<8 [‡]	No, NA
Lommatsch et al. (2017)	Yes, corrected	no	AngioVue group: -4.8 ± 6.31 AngioPlex group: -4.63 ± 5.7	840	<40	Yes
Rabiolo et al. (2018)	Yes, corrected	yes, corrected	NA	1040–1060	<7 [§]	No, NA
Rao et al. (2018)	Yes, corrected	yes, corrected	$-5.3 (-9.6, -3.1)^\dagger$	840	<35	Yes
Rao et al. (2016)	Yes, corrected	yes, corrected	POAG: $-6.3 (-13.5, -3.1)^\dagger$ PACG: $-9.2 (-16.0, -3.3)^\dagger$	840	<40	Yes
Rao et al. (2017c)	NO	Yes, corrected	$-6.3 (-12.5, -3.5)^\dagger$	840	<35	Yes
Rao et al. (2017a)	No	Yes, corrected	PAC: $-1.9 (-3.6, -0.8)^\dagger$ PACG: $-8.2 (-16.0, -4.0)^\dagger$	840	<35	Yes
Rao et al. (2017b)	No	Yes, corrected	POAG with DH: $-3.7 (-6.3, -2.5)^\dagger$ POAG without DH: $-3.8 (-7.5, -2.8)^\dagger$	840	<45	Yes
Rolle et al. (2017d)	No	Number of patients NA	PPG: -0.96 ± 1.15 POAG: -4.63 ± 3.53	840	<50	Yes
Shin et al. (2016)	No	No	-5.88 ± 6.01	840	NA [‡]	Yes
Takusagawa et al. (2010)	No	No	-5.32 ± 3.50	840	<50	No, NA
Triolo et al. (2017)	No	No	-5.45 ± 2.27	1040–1060	<7 [§]	Yes
Yarmohammadi et al. (2018,b)	Yes, corrected	Yes, corrected	$-3.9 (-8.8, -1.8)^\dagger$	840	<48	No, NA
Yip et al. (2017)	Yes, corrected	Yes	-11.07 ± 8.25	840	<40	

If a risk of bias for age or eye existed, yes was filled in. All values for visual field mean deviation are given in mean \pm SD, unless indicated otherwise. All images were acquired using AngioVue OCTA device, unless indicated otherwise. Corrections for age and/or the use of both eyes for one subject are presented in the table when executed (corrected). If no sector analysis was executed in the study, a fovea-disc axis correction was considered as unnecessary or not applicable (NA). Study characteristics that may bias the results are given in bold.

CFVD = central visual field defect, DH = disc haemorrhage, MD = mean deviation, PACG = primary angle closure glaucoma, PVFD = peripheral visual field defect, POAG = primary open-angle glaucoma, PPG = preperimetric glaucoma, VF = visual field.

* DRI OTC, Triton, Topcon.

[†] Values in median (IQR).

[‡] Cirrus-HD-OCT-5000, Zeiss Angioplex.

[§] SS-OCTA PLEX Elite 9000, Zeiss.

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