Scanning laser Doppler flowmeter study of retinal blood flow in macular area of healthy volunteers

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Aim: To compare the interocular and intraocular differences of capillary perfusion, and the intraocular regional differences of retinal blood flow in the macular area of healthy volunteers.

Methods: Tissue blood flow in the macula was examined in both eyes of 20 healthy volunteers with the Heidelberg retinal flowmeter. Blood flow measurements were made in a 10˚x2.5˚ area superior and inferior to the macula. The mean blood flow (MBF) was calculated by an automatic full field perfusion image analyser program. The MBF in the right and left eyes and in the superior and inferior macular areas of the same eye were compared.

Results: The ratios of the MBF in the right eye to the left eye in the macular areas were 1.00, and 1.03, respectively. The ratio of the MBF in the superior macular area to the inferior area was 1.01 for the right eyes and 1.04 for the left eyes.

Conclusions: Because no significant differences were found in the MBF between the two eyes and between the superior and inferior macular areas in the same eye, interocular (for example, affected eye versus fellow eye) and intraocular (superior versus inferior macular areas) comparisons of MBF can be made to determine if changes in retinal perfusion have occurred.

More and more attention has been focused on the impairment of tissue blood flow in the pathogenesis of retinal diseases and optic nerve diseases. Therefore, quantification of ocular blood flow, especially by non-invasive methods, has become essential in assessing and managing patients with retinal diseases and impaired ocular tissue perfusion.

There have been a number of investigations on ocular circulation using various techniques, each of which has advantages and disadvantages. Among these, the Heidelberg retinal flowmeter (HRF, Heidelberg Engineering, Heidelberg, Germany) has been used to measure retinal tissue flow from which accurate haemodynamic properties can be obtained. This technique combines a laser Doppler flowmeter with a scanning laser ophthalmoscopic system. Using a two dimensional measurement of the perfusion determined by the Doppler shift, it is possible to obtain reproducible and quantitative values of the capillary blood flow in distinct areas of a capillary meshwork of the retina and optic nerve head.

A recently developed computer program, called the automatic full field perfusion image analyser (AFFPIA), can be used to perform a full field analysis of images obtained with the HRF. This program incorporates a method of deleting recordings containing saccades from the analysis which then reduces the rate of image rejections.

Clinical studies have shown significant impairment of ocular blood flow in some retinal or optic nerve diseases. In some of these studies, the ocular blood flow measurements in the involved eye were compared with the flow in the fellow eye. This method is based on the hypothesis that the interocular blood flow in the two eyes is not statistically different. The blood flow in the ophthalmic artery and the central retinal artery has been reported to be symmetrical during the neonatal period and in adults, and several studies have shown that no significant interocular differences in perfusion are present.

The aim of this study was to compare the retinal capillary perfusion of the right and left eyes in the macular area of healthy volunteers using the HRF with AFFPIA. In addition, we compared the retinal capillary perfusion in the superior and inferior macular regions.

SUBJECTS AND METHODS

Twenty healthy subjects, five men and 15 women aged 18–42 years (mean 23.5 (SD 5.9)) years were studied. The experiments were performed according to the guidelines of the Declaration of Helsinki and, after explaining the purpose of the study, a signed informed consent was obtained from all subjects. All subjects underwent a thorough ophthalmological examination, including visual acuity, intraocular pressure (IOP) measurements, and slit lamp biomicroscopy. All had corrected visual acuity of at least 20/20, IOP below 21 mm Hg, and a normal retina and optic disc in both eyes. Subjects were excluded if they had undergone intraocular or laser surgery, had evidence of an eye infection, any history of systemic or ophthalmological disease or family history of diabetes or glaucoma, taken any medications within 2 weeks of the study, had a distance refractive error (spherical equivalent) of more than 6.0 dioptres, an astigmatism of more than 2.0 dioptres, and an interocular IOP difference >3 mm Hg, or refractive error difference >2.0 dioptres.

After the IOPs were measured by applanation tonometry, blood flow measurements were performed on both eyes by scanning laser Doppler flowmetry (SLDF) using the HRF. The areas measured were the superior and inferior areas around the nasal macula (Fig 1). The mean blood flow (MBF) was calculated by AFFPIA.

The principle, validity, and reliability of the SLDF technique to measure ocular blood flow have been published. The SLDF data included the measurement of a 10˚x2.5˚ retinal area with 256 points x64 lines. The pixelation of the instrument was 10 μm with a wavelength of 795 nm, an intensity of 100 μW, and a data acquisition time 2.048 seconds. Every line was scanned 128 times with a line sampling rate of 4000 Hz. The final data set contained 128 discrete recordings at each pixel, and the data were converted into a continuous wave function for each pixel.

The signals were fast Fourier transformed to obtain the power spectrum of the multiple frequency shift components. Images were included automatically by AFFPIA if the direct current (DC) value within the measurement area was between 40 and 228 arbitrary units. With confocal scanning
RESULTS

The IOPs in all subjects ranged from 10–16 (mean 12.3 (SD 1.7)) mm Hg in the right eye, and from 10–15 (12.4 (1.7)) mm Hg in the left eye. The IOPs in the two eyes were highly correlated ($r = 0.796$, $p = 0.0005$).

The MBF in each area is shown in Table 1. No significant difference was found in the MBF between superior and inferior areas in each eye ($p > 0.05$ in both eyes). The MBF in the right eye was highly correlated with the MBF in the left eye in the superior ($r = 0.747$, $p = 0.0011$; Fig 2A) and the inferior ($r = 0.797$, $p = 0.0005$; Fig 2B) nasal paramacular areas. In addition, the correlation of the MBF between the superior and inferior nasal paramacular areas in each eye was also highly significant ($r = 0.720$, $p = 0.0017$ in the right eye; Fig 3A, and $r = 0.877$, $p = 0.0001$, in the left eye; Fig 3B).

The MBF in each eye was not significantly correlated with the IOP (Fig 4) or with the age (Fig 5). The coefficients of variation demonstrated good reproducibility of the measurements—that is, 4.92% in superior areas and 7.21% in inferior areas in right eye, and 7.74% and 7.74% in left eye.

DISCUSSION

Attention has been focused on the changes of retinal blood flow in some retinal diseases to try to determine the aetiology and severity of the disease process, and also to evaluate the outcome of medical and/or surgical therapy. Attention has been focused on the changes of retinal blood flow in some retinal diseases to try to determine the aetiology and severity of the disease process, and also to evaluate the outcome of medical and/or surgical therapy.5–10,19–24 The perfusion in different areas was found to vary considerably, and was dependent on the user defined measurement window.25,26 In addition, the pulse phase analysed is an important source of bias even when the images are obtained by a skilled operator. In most reports using this experimental protocol, parts of the images, up to around 50%, were excluded from the analysis because of poor quality caused by eye movements or media opacities even when the images are obtained by a skilled operator. In most reports using this experimental protocol, parts of the images, up to around 50%, were excluded from the analysis because of poor quality caused by eye movements or media opacities.8,10 Moreover, a small (10×10) pixel frame might not be a large enough sampling window to overcome the physiological variability in blood flow across the retina. This sentiment has been expressed in previous reports.7,27

We used AFFPIA to minimise the bias caused by the heart beat associated pulsation and the spatial heterogeneity. In addition, we were careful in our exclusion criteria. Thus, we

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Mean blood flow (MBF) values</th>
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<tbody>
<tr>
<td></td>
<td>Right eye</td>
</tr>
<tr>
<td>Superior area</td>
<td>283.81 (60.40) [AU]</td>
</tr>
<tr>
<td>Inferior area</td>
<td>282.01 (54.12) [AU]</td>
</tr>
<tr>
<td>S/I ratio</td>
<td>1.01 (0.15)</td>
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AU = arbitrary units.

Data are expressed as mean (SD). R/L = ratio of the right eye to the left eye. S/I = ratio of the superior area to inferior area. No significant difference was found in the MBF between both eyes in both superior ($p = 0.627$) and inferior ($p = 0.351$) areas. No significant difference was found in the MBF between superior and inferior areas in each eye ($p = 0.765$ in right eye and $p = 0.316$ in left eye).
have evaluated the capillary perfusion around the nasal macula in the right and left eyes of healthy volunteers using the SLDF with AFFPIA, and the reproducibility at all locations was quite satisfactory and comparable to that of previous reports.92 8

Some investigators have reported that the blood flow velocities of the ophthalmic artery, central retinal artery, and central retinal vein were weakly or not correlated with age.91 41 92 42 9 In addition, when the retinal or optic nerve head microcirculation was evaluated using SLDF, there showed no significant correlation between MBF and age. This may be because our subjects were younger than 42 years and the range of ages was limited. Further studies with a large number of healthy subjects of different ages will be needed to determine whether age affects the MBF.

Studies examining the correlation of ocular circulation between superior and inferior retinal region have produced varying results.31 53 03 1 Yoshida et al noted that the blood flow in the superior temporal artery was higher than the inferior temporal artery. Feke and associates found the blood flow of the inferior retina was, on average, 6% greater than the flow of the superior retina but this difference was not significant. Schwartz et al reported that the blood flow in the inferior temporal artery was significantly greater than that in the superior temporal artery, but no significant differences were found for the inferior nasal artery compared to the superior nasal artery by fluorescein angiography with scanning laser opthalmoscopy. Squirrell et al reported that blood flow in the superior and inferior areas obtained by SLDF was significantly different around the macula in the three of 10 normal subjects. Our study is basically about microcirculation and free from previous findings on arterial blood flow. Our results strongly suggest that microcirculation of the superior area did not differ significantly from that in the inferior area around the macula. The discrepancy between these findings and those of Squirrell et al might be partly explained by the differences in the area analysed. As they suggested, the significant difference in blood flow between the two areas of normal retinas supports the idea that SLDF scans deeper than these has been assumed and may actually analyse both the retinal and choroidal capillary beds. In that case, both their data and our blood flow data around the macula reflect the contributions of the rich choriocapillaris, and the findings are dependent on any thinning of the retina in the analysed region.

As the sampling depth of the HRF is 300 μm, thinning of the retina, such as in the foveal and peripapillary area and of the neuroretinal rim, can cause the instrument to measure the deeper layers (for example, choriocapillaris) and not exclusively the retinal capillary bed.31 03 1 Measuring the retinal thickness at the location of interest might help resolve this problem.

In the foveola (about 350 μm in diameter), where the retina is very thin (about 130 μm) and avascular, the contribution from the choroidal circulation and/or the offset value of the HRF instrument might be involved. In the current study, we placed the centre of foveola at the edge of

Figure 2 (A) Correlation of the MBFs of the right and left eyes in the superior paramacular area. The MBFs in both eyes are significantly correlated. (B) Correlation of the MBFs between the right and left eyes in the inferior paramacular area. The MBFs in both eyes are significantly correlated.

Figure 3 (A) Correlation between the MBF in the superior and inferior areas of the right eye. The MBFs in both areas are significantly correlated. (B) Correlation between the MBF in the superior and inferior areas of the left eye. The MBFs in both areas are significantly correlated.
the measurement area ($10 \times 2.5\degree$) as shown in Figure 1. Assuming that the measurement area of $10 \times 2.5\degree$ is equal to $2.7 \times 0.7$ mm, and the diameter of foveola is $350 \mu m$, the percentage of foveola in the measurement area is about 1.27%. Thus, the contribution of the foveola in this study is very small.

Doppler ultrasound studies of the blood velocities in the ophthalmic artery and central retinal artery and vein in healthy subjects showed that no significant difference exists between the right and left eyes. Recently, Rawji and Flanagan and Boehm et al reported that there were no significant interocular differences in retinal and peripapillary capillary perfusion using SLDF. Our data confirmed their findings. All of the values in our study showed a difference between the two eyes of 11% to 13% which can be used as an index of the interocular variability in healthy eyes of young individuals without significant ametropia. However, the exact value of the technique should be interpreted in the light of the concerns mentioned. The mean flow from the entire image is significantly less variable compared with that from a particular area—for example, $10 \times 10$ pixels. Therefore, AFFPIA is a useful tool to evaluate the tissue MBF with less bias, and thus the present data can serve as a useful starting point for studies of the relation between ocular circulation and some retinal diseases.
Ocular microcirculation of healthy volunteers

1473

References

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