Regional Relationship Between Retinal Nerve Fiber Layer Thickness and Corresponding Visual Field Sensitivity in Glaucomatous Eyes

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Objective: To establish the structure-function relationship between peripapillary retinal nerve fiber layer (RNFL) thickness and visual field (VF) test points in standard automated perimetry.

Methods: We included 213 eyes with open-angle glaucoma and VF loss in this cross-sectional study. Correlations between individual VF sensitivity at 52 test points and peripapillary RNFL thickness divided into 16 sectors were calculated. The RNFL thickness was measured by Stratus optical coherence tomography. A new VF cluster map corresponding to RNFL sectors was generated by grouping the VF test points with the highest relation to each RNFL sector.

Results: The VF sensitivity at each test point was significantly correlated with the sectoral RNFL thickness. The highest coefficient of determination ($R^2$) for a superotemporal RNFL sector and VF sensitivity at an inferotemporal test point (9° temporal and 15° inferior from the center) in standard automated perimetry was 0.500 ($P < .001$). Clustered VF test points most highly related to the RNFL sectors were asymmetrically located between the upper and lower hemifields. A newly developed map revealed significant structure-function relationships.

Conclusions: We describe an association between VF sensitivity at test points and sectoral RNFL thickness. Nine clustered VF test points corresponding to 9 RNFL regions were demonstrated from the structure-function relationships.

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GLAUCOMA IS AN OPTIC NEUROPATHY characterized by progressive injury of the optic nerve and retinal nerve fiber layer (RNFL) that results in visual field (VF) deficits. It is of critical importance to detect the relationship between structural (RNFL or optic disc configuration) and functional (VF) damage for the diagnosis and management of glaucomatous optic neuropathy.

Various ocular imaging instruments, such as confocal scanning laser ophthalmoscopy, scanning laser polarimetry, and optical coherence tomography (OCT), have become available. Confocal scanning laser ophthalmoscopy can analyze optic disc configuration, whereas scanning laser polarimetry and OCT can measure RNFL thickness with objectivity and reproducibility.

In routine clinical practice, the VF is usually assessed via fixed test points with 6° distance with standard automated perimetry (SAP). A recent article claimed that RNFL thickness, measured by Stratus OCT, showed the strongest structure-function relationship among the 3 instruments.

Gardiner et al examined the relationships between VF sensitivity (VFS) at 52 tested locations using SAP and optic disc configuration using confocal scanning laser ophthalmoscopy. According to this study, VFS at each test point was significantly correlated with neuroretinal rim narrowing of the optic disc. As another way of analyzing structure and function, dividing the VF by grouping VFS points into several areas and averaging the threshold values of the clustered VFS points was documented. For example, Garway-Heath et al proposed a map of SAP test locations relating to positions of entry into the optic disc using RNFL defects recorded by photographs. However, to our knowledge there have been no studies evaluating the association of VFS at individual test points or at clustered points with peripapillary RNFL thickness measured with OCT technology.

The purposes of this study are to investigate the relationship between VFS at each
Figure 1. Schematic diagram of 16 retinal nerve fiber layer sectors in right eyes measured with optical coherence tomography. ST indicates superotemporal; SN, superonasal; IN, inferonasal; and IT, inferotemporal.

tested location in SAP and RNFL thickness measured with Stratus OCT and to evaluate the strength of the functional and structural association when the map described by Garway-Heath and colleagues is adopted. In addition, we developed a new structure-function map relating clustered VF test points to RNFL sectors, which is based on the RNFL thickness measured with OCT, and compared the relationships between the 2 maps.

In this retrospective study, nonrandomized subjects in the Department of Ophthalmology, Kobe University Hospital, Kobe, Japan, were consulted between April 1, 2005, and November 30, 2006. This study was conducted in accordance with the ethical standards stated in the 1964 Declaration of Helsinki and with the approval of the Medical Ethical Committee of the Kobe University Graduate School of Medicine. The committee determined that this study did not need subjects’ approval.

A total of 213 glaucomatous eyes from 131 Japanese adults (60 men and 65 women) younger than 70 years were enrolled in this study. No subjects had a history of diabetes mellitus. All of the eyes underwent comprehensive ophthalmic examinations and had a best-corrected visual acuity of at least 20/40. The SAP was performed with the Swedish Interactive Threshold Algorithm standard and central 30-2 program. The VF reliability criteria included fixation losses and false-positive and false-negative rates of less than 25%. The SAP and OCT examinations were performed within a maximum period of 3 months.

The SAP was performed with the Humphrey Field Analyzer 750 (Humphrey-Zeiss Instruments, Dublin, California) using the Swedish Interactive Threshold Algorithm standard and central 30-2 program. The VF reliability criteria included fixation losses and false-positive and false-negative rates of less than 25%. The SAP and OCT examinations were performed within a maximum period of 3 months.

RNFL THICKNESS

The RNFL thickness was measured with the peripapillary Fast RNFL program of the Stratus OCT (Carl Zeiss Meditec, Inc, Dublin) after a pupillary dilation. The basic principles and technical characteristics of the OCT have been described previously. The RNFL thickness was calculated using the system software (Stratus version 3.0 software; Carl Zeiss Meditec, Inc). One well-focused scan with a best signal length of at least 7 was included. Images with inappropriate RNFL borders, such as those that penetrated to another layer, were excluded.

The RNFL thickness was determined at 256 circumpapillary points (1.40625°) on a circle with a default diameter of 3.4 mm around the center of the optic disc. The data were exported and then combined into 16 sectors (22.5°) as shown in Figure 1. When the temporal margin was designated as 0°, superotemporal sectors 1 through 4 were 0° to 22.5°, 22.5° to 45°, 45° to 67.5°, and 67.5° to 90°, respectively. Superonasal sectors 1 through 4 were 90° to 112.5°, 112.5° to 135°, 135° to 157.5°, and 157.5° to 180°, respectively. Inferonasal sectors 1 through 4 were 180° to 202.5°, 202.5° to 225°, 225° to 247.5°, and 247.5° to 270°, respectively. Inferotemporal sectors 1 through 4 were 270° to 292.5°, 292.5° to 315°, 315° to 337.5°, and 337.5° to 360°, respectively.

STATISTICAL ANALYSES

All of the data were exported into a personal computer. Left-eye data were converted into right-eye format. All of the statistical analyses were performed using StatView version 5.0 statistical software (SAS Institute, Inc, Cary, North Carolina). The VFS at the 52 nonblind test points located within the 24-2 SAP VF was used and expressed in decibels. The relationships between RNFL thickness in 16 sectoral regions and VFS at the individual 52 points, as well as clusters of VFS as described in detail later, were subjected to a quadratic regression analysis. Second-order polynomial regression analysis has been demonstrated to better fit structure-function relationships than linear analysis when VFS is expressed in a logarithmic form (eg, in decibels), whereas this fit does not hold when VFS is expressed in a nonlogarithmic form (eg, 1/lambert). The coefficient of determination ($R^2$) was found by the quadratic regression. $P < .05$ was considered statistically significant.

The structure-function relationships from 6 regions were adopted from the map (Figure 2) designated previously by Garway-Heath and colleagues. In addition, a different map of the VFS clusters related to RNFL thickness was generated as follows. The RNFL sector with the highest relationship with VFS at each VF test point was extracted. Based on these data, the contiguous VF test points that had the highest $R^2$ with an identical RNFL sector could be grouped into a specific area of these contiguous VFS test points. Among the RNFL sectors, superotemporal sectors 1 and 2, inferotemporal sectors 3 and 4, and superonasal sectors 2 to 4 and inferonasal sectors 1 to 3 had significant relationships with a few VF test points. Thus, these 3 regions of RNFL sectors were combined. By doing this, 9 regional areas were constructed as the new map. Correlations between structure-function relationships in the new map were calculated.
The mean (SD) subject age was 51.5 (11.4) years (range, 22-69 years). The mean deviation in SAP ranged from −23.15 to −0.94 dB with a mean (SD) of −6.40 (5.45) dB. The mean (SD) VFS at the 52 nonblind locations in the 24-2 VF was 23.97 (5.57) dB.

The VFS at individual test points was significantly related to the thickness of more than 3 RNFL sectors. Among the 832 (521100316) relationships, the highest $R^2$ was 0.500 between RNFL superotemporal sector 3 and VFS at a test point located 6° temporally and 9° inferiorly from the center. Figure 3 shows pie charts representing graded $R^2$ between VFS at individual test points and RNFL sectors allocated according to the 24-2 test point locations. As earlier, each pie corresponds to a defined RNFL sector. The darker a sector is shaded, the stronger the $R^2$ relationship is. A white sector represents no significant relationship. The RNFL superonasal sector 4 had no significant relationship with VFS at any tested points. When a pair of VFS test points located in the mirror position at the upper and inferior hemifields was compared, the RNFL sector with the highest $R^2$ tended to be oriented more vertically in the upper hemifield. The highest $R^2$ at the 52 locations on SAP are shown in Figure 4. Overall, VFS in the nasal VF exhibited a higher $R^2$ than that in the temporal VF.

Figure 5 shows the RNFL sector that had the highest association with VFS at each VF test point from the pie chart shown in Figure 3. A bisecting line is drawn at the edge of the circle for each location in the sector corresponding to the entry points of RNFL defect by Garway-Heath and colleagues, and their degrees are shown for comparison with our map. The locations were determined using the temporal margin (9-o’clock position) set as 0° in right eyes. The correspondence rate between the 2 maps was 54% (28 of 52 test points) for the VF test points. The VF test points in nasal retina tend to have associations with RNFL sectors oriented slightly more toward the temporal side (0°) compared with the map by Garway-Heath and colleagues.

The 52 VF test points were clustered into regional groups of VFs that corresponded to RNFL sectors in 2 ways. First, the arrangement from Garway-Heath and colleagues was adopted (Figure 2). Coefficients of determination ($R^2$) between these 6 VFS clusters and the corresponding 6 sectors are listed in Table 1. Second, we generated a different grouped structure-function map that had 9 regional corresponding areas as mentioned in the “Methods” section (Figure 6). As shown in Table 2, the strength of correspondence between the RNFL sectors and the clustered VF test points in this map was similar to that in the map from Garway-Heath and colleagues in which the association between the sectoral optic disc margin and the clustered VF test points was evaluated. There was an asymmetrical correspondence of the VF loci and RNFL sectors between the upper and lower hemifields.

There were significant correlations between the VFS at individual test points and in several RNFL sectors. Grouping VF test points into 9 areas enabled us to generate the OCT-based structure-function correspondence map. The correspondence was somewhat different from a map generated by Garway-Heath and colleagues. This difference was rooted primarily with regions of structural parameters measured. Garway-Heath and colleagues used the optic disc margin, whereas OCT measured the RNFL thickness away from the optic disc margin. Because retinal nerve fibers do not project radially from the optic disc, the retinotopic orientation is different between a site on the optic disc margin and that on the concentric RNFL.
On the other hand, both maps show an asymmetrical correspondence of VFS loci and RNFL sectors between the upper and lower hemifields. Inferior RNFL sectors govern broader VF clusters in the upper hemifield than superior RNFL sectors in the lower hemifield. A previous investigation\textsuperscript{11} using a cluster analysis of static VFs described horizontal asymmetry. Another study\textsuperscript{12} also showed horizontal asymmetry in the long-term progression of glaucomatous VF loss. Duke-Elder\textsuperscript{13} noticed the asymmetry in the entry of ganglion cell axons into the optic disc. He noted that axons from the inferior retina preferentially entered into the nasal side of the optic disc more frequently than those from the upper retina. From another point of view, clockwise rotation in the right eye and counterclockwise rotation in the left eye can be used to describe RNFL sector orientation relative to VF loci. Similar asymmetry of the structure-function relationships was also demonstrated in a previous study\textsuperscript{2} that analyzed optic disc configuration by confocal scanning laser ophthalmoscopy and a recent study\textsuperscript{14} that analyzed RNFL thickness by Stratus OCT. The degree of rotation in our map may consist of more than

![Figure 3. The relationship between retinal nerve fiber layer sectors and visual field sensitivity at 52 test points in standard automated perimetry with a 24-2 central program. Each circle shows the relation map for 1 test point in the visual field. The retinal nerve fiber layer sectors are shaded according to the strength of the correlation of determination ($R^2$), with darker shading indicating a larger correlation.](image)

![Figure 4. The highest correlations of determination between visual field sensitivity and sectoral retinal nerve fiber layer thickness at 52 tested locations. $P$ values are also shown.](image)
1 RNFL sector (equal to 22.5°). Garway-Heath and colleagues demonstrated that the location of the optic nerve head was a mean (SD) of 15.5° (0.9°) nasal to and 1.9° (1.0°) above the fovea. It could be speculated that the rotation angle was 7.0° (tangent $\tan^{-1} \frac{1.9}{15.5}$) from their data. Another study reported this angle to be 5.6°. We suggest that the degree of rotation in our map is bigger than that of the anatomical rotation. Whether this asymmetry affects or is responsible for the development and progression of glaucomatous optic neuropathy is a matter of particular interest and requires further investigation.

The $R^2$ values at VF locations temporal to the blind spot were relatively low compared with those at locations nasal to the blind spot. Additionally, the superonasal sector 4 of the RNFL showed no significant relationship with VFS at any tested location in SAP. These results may arise owing to the small number of eyes with low VFS in this area. Only 6 of 213 eyes (2.8%) had a VFS less than 10 dB at the test point 6° temporal to the blind spot, whereas 31 of 213 eyes (14.6%) had a VFS less than 10 dB at the test point 6° nasal to the blind spot. It has been suggested that the thickness of the nasal RNFL sector has higher individual variability and less reproducibility. It is very difficult to estimate the structure-function relationship at the nasal region adjacent to the optic disc because of the paucity of test points on SAP at the temporal regions adjacent to the blind spot.

There are several limitations to this type of study. First, there may be a selection bias. In this study, glaucoma was defined on the basis of structure and VF loss. We wanted to generate a corresponding map between OCT-based RNFL thickness and VF test points. To facilitate this procedure,
we intentionally used eyes with glaucomatous optic neuropathy with VF loss. Therefore, eyes with so-called preperimetric glaucomatous optic neuropathy were excluded from this study. This may cause some bias in establishing the strength of the structure-function associations.

Second, we included both eyes in the partial population. It is known that optical and biochemical characteristics are similar between bilateral eyes. The use of both eyes may therefore inflate the structure-function association values.

Third is the aging effect. Because RNFL thickness measured by OCT is known to be significantly reduced in eyes from an older population, participants older than 70 years were excluded. Additionally, we reported that RNFL thickness was affected by both the size of the optic disc and refraction. We cannot completely rule out the influence of these issues on the present results because corrected measurements (eg, total deviation at tested locations in SAP or age-adjusted RNFL thickness) were not used.

Last but not least, the structure-function relationship may be skewed. The range of defect depth at specific VF test points affects the strength of the associations. If there is greater loss in some RNFL or VF locations, this may result in a stronger apparent association with these locations even if the true association is with correlated but less damaged RNFL or VF locations. Also, if some RNFL sectors tend to have less interindividual variability than others, the apparent (statistical) association may be stronger with a correlated but less variable RNFL sector. This may distort the true structure-function relationship. These effects may skew associations to the inferotemporal and superotemporal (arcuate) regions of the RNFL. As reviewed by Hood and Kardon, the actual correspondence between structure and function cannot be fully revealed.

In conclusion, the OCT-measured sectoral RNFL thickness had significant associations with VFS on SAP. The correspondence map between clustered VF test points and the grouped RNFL sectors was different from the map proposed by Garway-Heath and colleagues, essentially owing to differences of measurement of structural parameters. Both maps showed an asymmetrical correspondence of VFS loci and RNFL sectors between the upper and lower hemifields.

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**REFERENCES**


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