Hypodense Regions (Holes) in the Retinal Nerve Fiber Layer in Frequency-Domain OCT Scans of Glaucoma Patients and Suspects

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PURPOSE. To better understand hypodense regions (holes) that appear in the retinal nerve fiber layer (RNFL) of frequency-domain optical coherence tomography (fdOCT) scans of patients with glaucoma and glaucoma suspects.

METHODS. Peripapillary circle (1.7-mm radius) and cube optic disc fdOCT scans were obtained on 208 eyes from 110 patients (57.4 ± 13.2 years) with glaucomatous optic neuropathy (GON) and 45 eyes of 45 controls (48.0 ± 12.6 years) with normal results of fundus examination. Holes in the RNFL were identified independently by two observers on the circle scans.

RESULTS. Holes were found in 33 (16%) eyes of 28 (25%) patients; they were not found in any of the control eyes. Twenty-four eyes had more than one hole. Although some holes were relatively large, others were small. In general, the holes were located adjacent to blood vessels; only three eyes had isolated holes that were not adjacent to a vessel. The holes tended to be in the regions that are thickest in healthy controls and were associated with arcuate defects in patients. Holes were not seen in the center of the temporal disc region. They were more common in the superior (25 eyes) than in the inferior (15 eyes) disc. Of the 30 eyes with holes with reliable visual fields, seven were glaucoma suspect eyes with normal visual fields.

CONCLUSIONS. The holes in the RNFL seen in patients with GON were probably due to a local loss of RNFL fibers and can occur in the eyes of glaucoma suspects with normal visual fields. (Invest Ophthalmol Vis Sci. 2011;52:7180 –7186) DOI:10.1167/iovs.11-7716

Glaucoma damages retinal ganglion cells (RGC) and their axons. This loss of RGC axons leads to thinning of the peripapillary retinal nerve fiber layer (RNFL) and to characteristic optic disc changes. Optical coherence tomography (OCT) permits in vivo imaging of the peripapillary RNFL; the recently improved resolution of frequency domain (fd) OCT allows imaging of the macular RGC layer as well.

Numerous fdOCT studies of glaucoma patients have documented thinning of the RNFL (see Refs. 1, 2 for review) and, more recently, of the RGC layer.3–6 Many of these studies have suggested ways to calculate the thickness of one or more layers of the fdOCT scan so as to improve the sensitivity and specificity of tests for glaucomatous damage. Because fdOCT is still a relatively new technique, additional information may be revealed in these scans that might improve our understanding of glaucomatous damage.

We have observed hypodense regions (holes) in the RNFL of some patients with glaucomatous optic neuropathy (GON). Our purpose was to better understand the prevalence and location of these regions.

METHODS

We examined 208 eyes of 110 patients (mean age, 57.4 ± 13.2 years) with GON and 45 eyes of 45 controls (48.0 ± 12.6 years) with normal disc examinations. Consecutive patients with GON referred for multifocal VEP testing were included if they had well-centered scans with quality scores better than 65 and little motion artifact. Written informed consent was obtained from all subjects. Procedures followed the tenets of the Declaration of Helsinki, and the protocol was approved by the Committee of the Institutional Board of Research of Columbia University.

Peripapillary circle (1.7-mm radius) and 6 mm × 6 mm cube scans of the optic disc (3D-OCT 1000 and 3D-OCT 2000; Topcon Inc, Paramus, NJ) were obtained. For the peripapillary circle scans, each of the 16 overlapping (averaged) B-scans had 1024 A-scans, whereas the cube scan had 512 A-scans for each of the 128 B-scans. Figure 1A shows the location of the circle scan for one of the patients. To identify hypodense regions (holes) in the RNFL, two of the authors examined the circle scans. When they disagreed, a third author adjudicated. The resolution of the cube scans was, of course, poorer than the resolution of the circle scans, and cube scans were only used to confirm that the holes extended beyond the circle scan image (see Fig. 3).

RESULTS

A typical peripapillary scan is shown in Figure 1B for a right eye. A magnified view of the region within the white box is presented in Figure 2A, where blood vessels can be identified by the shadows they cast. The red ellipses show a cluster of three small holes to the left of a large blood vessel, with a larger hole falling to the right of the same vessel. Twenty-eight (25%) of the 110 patients had holes in at least one eye. Of the 208 eyes with GON, 33 (16%) had at least one hole. Holes were not found in any of the healthy control eyes.
Figure 2 shows examples of the pattern of holes seen in the RNFL. Some holes were small (Figs. 2A, 2D, 2G), and some were relatively large (Figs. 2E, 2F) and sometimes in clusters (Fig. 2A). Of the 33 eyes with holes, 24 had more than one hole. These holes could be near each other (Figs. 2A, 2B, 2E) or they could be in another area of the RNFL, often the other half of the RNFL. The holes tended to occur immediately adjacent to a vessel, as shown in Figures 2A to 2F. In fact, 31 of the 33 eyes had at least one hole near a vessel. Further, only three eyes had a hole that was not immediately adjacent to a vessel; two of these are shown in Figures 2G and 2H.

Using the cube scans, it was often possible to follow the course of a hole across scan lines, confirming that the holes were not local artifacts. Figure 3 provides an example. Figure 3A (the same image as Figure 2G) is from the standard circle scan with a radius of 1.7 mm. The holes are present, although more difficult to see in the equivalent circle scan (Fig. 3B, green border) derived from the cube scan. The colored circles on the fdOCT image in Figure 3C show the locations of derived circle scans closer to (blue) and further from (red) the optic disc than the 1.7 mm scan. The equivalent portions of these derived scans are shown in Figure 3B, where a hole is present at radii of 1.2 and 1.7 mm but a depression is present at a radius of 2.4 mm. The symbols in Figure 3C mark the centers of the holes (magenta circles) and depressions (orange triangles). Note in Figure 3C that there are holes closer to the disc and a depression/thinning of the RNFL further from the disc and that together they trace a path consistent with that of an RNFL bundle.

The locations of the holes in all 33 eyes are shown in Figure 4. The location, in degrees, is referenced to a 360° scale, where 0° is the most temporal part of the disc (9 o’clock for the right eye and 3 o’clock for the left eye). (B) Circle scan of eye 33. White box: region with holes, enlarged in Figure 2A.

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Figure 2. Examples of holes and clusters of holes in eight eyes: eyes 33 (A), 7 (B), 2 (C), 32 (D), 12 (E), 22 (F), 9 (G), and 18 (H).
deviation (MD) of $-0.25 \text{ dB}$ and glaucoma hemifield test (GHT) results within normal limits; there was only one abnormal (5%) point. Seven eyes had MD and GHT scores within normal limits, and another two eyes had borderline GHT and normal MD values.

In general, the eyes with holes had mild to moderate field defects. The 30 eyes with holes and visual fields within 7 months of the fOCT scan had a median MD of $-3.78 \text{ dB}$. Twenty-six of these eyes had MD values better than $-5 \text{ dB}$ and only one had an MD worse than $-8.04 \text{ dB}$. More holes were seen in nine eyes with MD values within normal limits.

**DISCUSSION**

On peripapillary fOCT scans, hypodense regions, which looked like holes, were seen in the RNFL of the patients with glaucomatous optic discs. These holes were typically located adjacent to major blood vessels. We hypothesize that these holes were probably caused by local loss of RNFL fibers and that they might have been early signs of glaucomatous damage because they also occurred in eyes of glaucoma suspects with normal visual fields.

A number of findings support the first hypothesis. First, holes were not seen in the control eyes. Second, the holes were typically seen in the arcuate regions of the RNFL associated with glaucomatous damage and not in the regions (e.g., center of the temporal sector) less likely to be associated with damage. Third, when the holes were relatively large, a visual field defect was seen in the corresponding region. Finally, it is often possible to track a hole throughout the cube scan, where it follows a course consistent with an arcuate defect. These findings confirm that we were not dealing with a local phenomenon or artifact.
We do not have a satisfactory explanation for why the holes tended to appear near blood vessels. One might expect more holes near major vessels because these are also the regions where the RNFL is thickest and most susceptible to arcuate damage. However, this does not explain why nearly all the holes are adjacent to blood vessels because there are other thick regions of the RNFL that are not immediately adjacent to blood vessels. One possibility is that a loss of local axons may create a local mechanical force that pulls axons away from the nearest vessel. Alternatively, and probably less likely because vessels are known to constrict in patients with glaucoma, it may be the vessel itself is being pulled away from the axons. In any case, there are reasons to believe that the presence of small holes might be an early sign of glaucomatous damage to the RNFL. Again, we did not see holes in healthy controls, but we did see holes in seven eyes with normal visual fields (both MD and GHT within normal range). Further, holes were seen in eight hemifields without a cluster of abnormal points on the visual field. It remains to be seen whether the presence of holes
will be clinically useful in identifying early glaucomatous damage.

**References**


