Quantitatively Measured Anatomic Location and Volume of Optic Disc Drusen: An Enhanced Depth Imaging Optical Coherence Tomography Study

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Quantitatively Measured Anatomic Location and Volume of Optic Disc Drusen: An Enhanced Depth Imaging Optical Coherence Tomography Study

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PURPOSE. Optic disc drusen (ODD) are found in up to 2.4% of the population and are known to cause visual field defects. The purpose of the current study was to investigate how quantitatively estimated volume and anatomic location of ODD influence optic nerve function.

METHODS. Anatomic location, volume of ODD, and peripapillary retinal nerve fiber layer and macular ganglion cell layer thickness were assessed in 37 ODD patients using enhanced depth imaging optical coherence tomography. Volume of ODD was calculated by manual segmentation of ODD in 97 B-scans per eye. Anatomic characteristics were compared with optic nerve function using automated perimetric mean deviation (MD) and multifocal visual evoked potentials.

RESULTS. Increased age (P = 0.015); larger ODD volume (P = 0.002); and more superficial anatomic ODD location (P = 0.007) were found in patients with ODD visible by ophthalmoscopy compared to patients with buried ODD. In a multivariate analysis, a worsening of MD was significantly associated with larger ODD volume (P < 0.0001). No association was found between MD and weighted anatomic location, age, and visibility by ophthalmoscopy. Decreased ganglion cell layer thickness was significantly associated with worse MD (P = 0.025) and had a higher effect on MD when compared to retinal nerve fiber layer thickness.

CONCLUSIONS. Large ODD volume is associated with optic nerve dysfunction. The worse visual field defects associated with visible ODD should only be ascribed to larger ODD volume and not to a more superficial anatomic ODD location.

Keywords: optic disc drusen, optic nerve head drusen, 3D segmentation, visual field defects
investigated whether RNFL and macular ganglion cell layer (GCL) thickness work as anatomic correlates to optic nerve dysfunction in ODD patients.

**PATIENTS AND METHODS**

The study was a prospective observational study approved by the scientific ethics committee of the Capital Region, Denmark (H-4-2013-040).

**Patient Selection**

Patients diagnosed with ODD from January 1, 2009, to January 1, 2016, were asked to participate in the study. All patients were seen at the Department of Ophthalmology at Rigshospitalet-Glostrup, Denmark. The patient exclusion criteria were best corrected visual acuity (BCVA) >logMAR 0.2, age <18 years, and presence of systemic disease that could affect optic nerve function. Exclusion criteria for individual eyes were localized eye or optic nerve disease other than ODD (e.g., optic neuritis, glaucoma, etc.) or ODD complications (e.g., drusen-associated anterior ischemic optic neuropathy, central retinal artery, vein occlusion, etc.) that could affect optic nerve function.

Informed consent was obtained from the subjects after explanation of the nature and possible consequences of the study. All procedures adhered to the tenets of Declaration of Helsinki.

**Data Acquisition**

All examinations were performed by a single examiner (LM). All included participants were asked about medication use as well as ophthalmic and medical history. Best corrected visual acuity was determined using Early Treatment Diabetic Retinopathy Study (ETDRS) charts (4-meter original series; Precision-Vision, La Salle, IL, USA). Patients were examined using slit lamp biomicroscopy and intraocular pressure was measured by applanation tonometry. Spectral domain EDI-OCT (Spectralis HRA+OCT; Heidelberg Engineering, Heidelberg, Germany) was performed using the following protocol: (1) dense optic nerve head scan for identification and quantification of ODD with EDI-OCT in both vertical and horizontal directions with 30 μm between each B-scan (97 scans), averaging 30 B-scans; (2) peripapillary evaluation of RNFL thickness with a 12° circumferential scan; and (3) macula overview in vertical direction with 240 μm between each B-scan for evaluation of macular GCL thickness. All scans were performed in high resolution with averaging of B-scans using the built-in eye tracking feature. Patients were dilated with 2.5% phenylephrine before OCT acquisition.

Recording and analysis of mfVEP data was performed as previously described. Briefly, patients were stimulated in a viewing distance of 30 cm to a screen (22-inch, high-resolution LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a cortically scaled 56-segment LCD display, 90% brightness and 65% contrast; Hitachi, Ltd., Tokyo, Japan) containing a corn.
eye with worst MD on automated perimetry was used in patients with bilateral ODD.

Mean and standard deviations or median and interquartile ranges (skewed distributions) were reported for continuous variables. Student’s t-test or Wilcoxon signed-rank test (skewed distribution) was used to compare patients with visible and buried ODD. We used \( \chi^2 \) or Fisher’s exact tests (expected count < 5) for categorical data.

The assumptions of linearity, homoscedasticity, and normal distribution of residuals were tested when performing multiple regressions. The contribution of each predictor in the multiple regression analysis was found by assessing standardized parameter estimates (the change in \( Y \), measured in units of its standard deviation, associated with a 1 standard deviation change in \( X \)). The assumptions of linearity, homoscedasticity, related pairs, and normality of variables were tested when performing correlation analysis. Adjustment for multiple testing in the correlation analyses was performed using the Holm-Bonferroni method. The predetermined level of statistical significance for the comparisons was \( P \leq 0.05 \).

**RESULTS**

We included 37 patients (30 women and 7 men) in this study. All included patients were Caucasian. Bilateral ODD were found in 95% of the patients. All eyes with ODD had one or more hyporeflective structures with a full or partial hyperreflective margin using OCT. Differences in clinical, mfVEP, and EDI-OCT findings were compared between patients with visible (visible by ophthalmoscopy) and buried (only visible by EDI-OCT) ODD (Table 1). Patient with buried ODD were significantly younger (median age: 21 years) than patients with visible ODD (median age: 33 years; \( P = 0.015 \)). Significantly thinner peripapillary RNFL thickness and macular GCL thickness (3–6 mm from fovea) were found in patients with visible ODD (\( P < 0.001, P = 0.002 \)). A tendency toward larger scleral canal size in patients with visible ODD was found (\( P = 0.05 \)). A worse MD was found in patients with visible ODD (\( -4.3 \) dB) when compared to patients with buried ODD (\( -1.9 \) dB; \( P = 0.025 \)). The quantitative measure of anatomic location was significantly different between the two groups with the center of weighted ODD mass being 172 \( \mu \)m below the reference level in patients with visible ODD and 306 \( \mu \)m below the reference level in patients with buried ODD (\( P = 0.046 \)). Volume of ODD was larger in patients with visible ODD (0.29 \( \text{mm}^3 \)) than in patients with buried ODD (0.01 \( \text{mm}^3 \); \( P = 0.002 \)).

A multiple linear regression was calculated to predict MD based on ODD volume, visibility by ophthalmoscopy, anatomic ODD location, and age. A significant regression equation was found (\( R^2 = 0.52, P < 0.0001 \)). Larger ODD volume was associated with worse MD (\( P < 0.0001 \)). For every 1 \( \text{mm}^3 \) increase in ODD volume, the MD decreased by 18.1 dB (CI 95% -25.6 to -10.7 dB). Anatomic ODD location, age, and visibility by ophthalmoscopy were not found significantly associated with MD when adjusted for the other variables. When looking at standardized parameter estimates, ODD volume had a higher effect on MD when compared to weighted ODD location. Figure 2 illustrates the ODD segmentation, en face overview, and corresponding visual field and RNFL thickness map in three selected patients.

Another multiple linear regression was performed to estimate the relative effect of RNFL and macular GCL thickness on MD. A significant regression equation was found (\( R^2 = 0.58 \), \( P < 0.0001 \)).
TABLE 1. Differences in Clinical, mVEP, and EDI-OCT Findings in Patients With Visible and Buried ODD

<table>
<thead>
<tr>
<th>Variable</th>
<th>Visible ODD (n = 32)</th>
<th>Buried ODD (n = 5)</th>
<th>P Values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Female sex, n</td>
<td>26</td>
<td>4</td>
<td>1.0*</td>
</tr>
<tr>
<td>Age, y</td>
<td>63 (26)</td>
<td>21 (1)</td>
<td>0.015‡</td>
</tr>
<tr>
<td>Refractive error, D</td>
<td>−0.75 ± 3.1</td>
<td>0.95 ± 2.8</td>
<td>0.28‡</td>
</tr>
<tr>
<td>IOP, mm Hg (applanation tonometry)</td>
<td>13 (5)</td>
<td>13 (2)</td>
<td>0.98‡</td>
</tr>
<tr>
<td>BCVA, (ETDRS, letters)</td>
<td>88 (6)</td>
<td>87 (1)</td>
<td>0.79‡</td>
</tr>
<tr>
<td>Ishihara</td>
<td>16/16</td>
<td>16/16</td>
<td>0.44‡</td>
</tr>
<tr>
<td>RAPD</td>
<td>8</td>
<td>0</td>
<td>0.56*</td>
</tr>
<tr>
<td>MD, DB</td>
<td>−4.5 (10.3)</td>
<td>−1.9 (2.6)</td>
<td>0.025‡</td>
</tr>
<tr>
<td>Peak-to-peak mVEP amplitude, nV</td>
<td>122 ± 55</td>
<td>131 ± 48</td>
<td>0.74‡</td>
</tr>
<tr>
<td>Signal-to-noise ratio mVEP amplitude</td>
<td>3.8 ± 1.1</td>
<td>3.3 ± 0.6</td>
<td>0.29‡</td>
</tr>
<tr>
<td>Second peak mVEP latency, ms</td>
<td>154±9</td>
<td>153 ± 2</td>
<td>0.81‡</td>
</tr>
<tr>
<td>Retinal macular thickness, µm</td>
<td>277.8 ± 20.7</td>
<td>279.2 ± 10.6</td>
<td>0.83‡</td>
</tr>
<tr>
<td>Macular GCL thickness 3–6 mm, µm</td>
<td>29.5 ± 5.8</td>
<td>36.4 ± 3.0</td>
<td>0.002‡</td>
</tr>
<tr>
<td>Scleral canal diameter, µm</td>
<td>1631 (250)</td>
<td>1477 (114)</td>
<td>0.05‡</td>
</tr>
<tr>
<td>ODD volume, mm³</td>
<td>0.29 (0.41)</td>
<td>0.01 (0.02)</td>
<td>0.002‡</td>
</tr>
<tr>
<td>Mean anatomic location below reference level, µm</td>
<td>49 (211)</td>
<td>295 (114)</td>
<td>0.007‡</td>
</tr>
</tbody>
</table>

*χ² test. †Wilcoxon rank sum test. ‡Student’s t-test.

P < 0.0001). Worse MD was significantly associated only with GCL thickness (P = 0.025) when adjusted for the other variable. Mean deviation increased by 0.65 dB (CI 95% 0.08–1.2 dB) for every 1 µm increase in GCL thickness. When looking at standardized parameter estimates, GCL thickness had a higher effect on MD when compared to RNFL thickness.

Table 2 summarizes the correlation between anatomic and functional markers of optic nerve dysfunction. Macular GCL thickness had the highest degree of correlation when compared to MD (ρ = 0.76, P < 0.0001), while macular GCL thickness and peripapillary RNFL thickness were comparable when looking at mVEP parameters. The unadjusted correlation coefficient for the correlation between ODD volume and MD was −0.66 (Fig. 3).

DISCUSSION

Our study is, to the best of our knowledge, the first to quantitatively assess both the ODD volume and anatomic location of ODD. Using visibility by ophthalmoscopy to classify ODD in superficial or buried might be obsolete due to technical advances in imaging techniques. We therefore calculated ODD volume and anatomic location quantitatively and applied it in a multivariate model for a better understanding of their relative contribution to optic nerve dysfunction.

We found that a larger ODD volume resulted in worse MD when adjusted for age, visibility by slit lamp, and anatomic location. Other studies have quantitatively assessed ODD size,11,12 and similar results were found in a recent case-series including five patients,12 where an excellent correlation between ODD volume and MD using automated perimetry was found. We suspect the increasing optic nerve dysfunction caused by larger ODD volume might be a result of either direct compression of adjacent ganglion cell axons, leading to ganglion cell death or secondary to compromised vascular flow.21

No association between weighted anatomic ODD location and MD was found in the current study. This is interesting as several studies have found worse MD in patients with ophthalmoscopically visible ODD.8,22,25 Other studies have further found more abnormal visual fields in patients with visible ODD when compared to patients with buried ODD.24–26 The results from this study suggest that age, ODD volume, and ODD location all contribute to ODD volume visibility. This means that equating ODD visibility on ophthalmoscopy with superficial anatomic ODD location only, incorrectly leads one to believe that there is an association between superficial anatomic ODD location and worse MD. Our findings suggest that solely larger ODD volume, and not a more superficial ODD location, results in higher degrees of visual field defects.

Based on our finding that visibility by ophthalmoscopy did not have an effect on MD when adjusted for age, ODD volume and ODD location, we argue that the classification using ODD visibility by ophthalmoscopy is not ideal to estimate optic nerve dysfunction. Furthermore, the term “superficial”, often used in the classification, is misleading as several factors, such as age and ODD volume, influence the visibility. In this regard, the term “visible” ODD might therefore be more appropriate to use than superficial ODD.

We found differences in age, MD, RNFL, and GCL thickness that are supported by several studies when using the classification of ODD as visible or buried.8–11,23,27 Our results of the multivariate analysis suggest that GCL thickness is a better anatomic correlate to optic nerve dysfunction in ODD patients than RNFL thickness. While the macular GCL thickness has not been explored extensively in ODD literature, macular ganglion cell–inner plexiform layer thickness has proven to be a predictor of early glaucoma with the same sensitivity as RNFL thickness.28,29 It has even been suggested that individually segmented GCL thickness could be a better predictor for the presence of preperimetric glaucoma than RNFL thickness.30

Conflicting results have been published about the role of scleral canal size in ODD etiology.31–34 A larger scleral canal in patients with superficial ODD has been previously reported,35,36 and in this study, the same tendency was found. In unpublished data (Malmqvist L, unpublished poster presentation, 2016), we have found smaller scleral canal in ODD children when compared to healthy children and we therefore suggest the finding of this study is due to a displacement of Bruch’s membrane caused by distending ODD. This was originally proposed as an alternative explanation for similar findings in a study by Floyd et al.34 Our proportion of patients
with bilateral ODD (95%) is the highest reported in the literature. Most studies have reported bilateral ODD in 62% to 76% of patients. By using EDI-OCT in our study, we were able to diagnose eyes with small and deeply buried ODD overlooked by ophthalmoscopy, autofluorescence, B-scan ultrasound, and even conventional spectral-domain OCT. Based on these results, we propose that bilateral ODD are more common than previously believed.

In this study, we included mfVEP amplitude and latency as an objective measure of optic nerve function as previous studies have found significantly decreased amplitude and latency delays in patients with optic disc drusen when compared to control subjects. Hence an association between ODD volume and mfVEP parameters was expected. Parameters of mfVEP were significantly correlated with RNFL thickness, but not with ODD volume. That mfVEP amplitude was not correlated with ODD volume is likely a result of the high intersubject variability and in this case, we assume that the variability was too high to describe the more subtle changes in optic nerve dysfunction when only comparing ODD patients.

The major limitation of the study was the use of multiple regression analysis with the limited amount of patients. By using the covariate “visibility by ophthalmoscopy,” including only five eyes with buried ODD, the estimation was not strong. The fact that we, in multiple regression analyses, did not find an association between ODD volume and mfVEP parameters was expected. However, the mean spherical equivalent refraction was not significantly different between patients with visible and buried ODD. In this study, we exclusively measured the volume of ODD defined as hyporeflective structures with a full or partial

### TABLE 2. Correlation Coefficients for Correlations Between Anatomic and Functional Markers of Optic Nerve Dysfunction

<table>
<thead>
<tr>
<th>Variable</th>
<th>Automated Perimetry MD</th>
<th>Multifocal Visual Evoked Potentials</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>P2P</td>
</tr>
<tr>
<td>ODD volume</td>
<td>-0.66*</td>
<td>-0.43</td>
</tr>
<tr>
<td>RNFL thickness</td>
<td>0.72*</td>
<td>0.51*</td>
</tr>
<tr>
<td>GCL thickness</td>
<td>0.76*</td>
<td>0.56*</td>
</tr>
</tbody>
</table>

Pearson’s correlation coefficient between anatomic and functional markers of optic nerve dysfunction.

* Holm-Bonferroni adjusted significant (P < 0.05). Latency was measured as second peak latency. Global RNFL thickness was measured peripapillary. Thickness of GCL was measured as the mean of a 3 to 6 mm ring around the fovea.

**Figure 2.** En face view and horizontal 3D view of optic nerve head and segmented ODD, as well as corresponding visual fields and RNFL thickness map in three different patients. Different colors symbolize the individual drusen. Colors in the RNFL thickness map indicate if the thickness in different regions is within or outside the statistical limits of normality: green, normal; yellow, borderline below normal limits; red, below normal limits; blue, borderline above normal limits. (A) Right eye of a patient with total ODD volume of 0.07 mm³. Visual field testing revealed a near-normal mean deviation of −2.9 dB and global RNFL thickness was decreased to 77 μm. (B) Right eye of a patient with total ODD volume of 0.23 mm³. Visual field testing revealed a mean deviation of −12.3 dB and global RNFL thickness was decreased to 57 μm. (C) Left eye of a patient with total ODD volume of 0.69 mm³. Visual field testing revealed a mean deviation of −24.3 dB and global RNFL thickness was decreased to 40 μm.
hyperreflective margin. However, other studies have reported ODD as either hyperreflective, granular, or hyporeflective when using OCT.11,39 The conflicting descriptions of ODD morphology are a limitation in this as well as other ODD studies, and should be addressed in future research.

In conclusion, this study suggests that ODD volume is significantly associated with optic nerve dysfunction. Even though a worse MD is often found in patients with visible ODD, a more superficial anatomic ODD location is not necessarily associated with worse MD. The current classification using visibility by ophthalmoscopy is an unspecific marker of optic nerve dysfunction compared to quantitative measurements of ODD volume using EDI-OCT.

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References


