Bone Remodeling in Choroidal Osteoma Monitored by Fundus Photography and Spectral-Domain Optical Coherence Tomography

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Key Words
Choroidal osteoma · Tumor progression · Bone remodeling · Fundus photographs · Spectral-domain optical coherence tomography

Abstract
Choroidal osteoma is a benign ossifying tumor of the choroid, consisting of mature bone tissue. It has been described to enlarge and evolve at varying rates over time. Here, we report and quantify the progression of a unilateral choroidal osteoma in a 7-year-old boy by fundus photography, and document tumor remodeling by spectral domain optical coherence tomography images.

Introduction
Choroidal osteoma is a benign rare tumor consisting of normal cancellous bone located between the inner third and outer third of the choroidal layers [1]. The typical characteristics of choroidal osteomas are their unilaterality, juxta- or peripapillary location, predominance in young females [2], rapid or slow tumor progression [3–5], tumor involution [6] and decalcification [7, 8]. Diagnosis, by convention, is made by clinical examination and fundus angiography, and confirmed by the presence of choroidal calcification on ultrasonography or other modes of imaging such as computed tomography. Spectral-domain optical coherence tomography (SD-OCT) has provided high-resolution cross-sectional images of the tumor and overlying retina [9–11], as well as a 3-dimensional view of the tumor [12]. Remodeling of the...
tumor has been suggested to explain the varying degrees of tumor reflectivity found on SD-OCT [10]. In this case report we objectively quantify the progression of a choroidal osteoma in a young boy, and demonstrate the effects of tumor remodeling documented by SD-OCT.

Case Report

An otherwise healthy 7-year-old boy, diagnosed with a right submacular choroidal osteoma as an incidental finding, was followed up over a period of 5 years. He had neither a preceding history of systemic illness nor of ocular trauma. His vision remained 20/20 in both eyes with minimal refractive correction throughout the years. There was varying central metamorphopsia. Anterior segment examination was unremarkable. Examination revealed a large round and well-demarcated orange-red-colored submacular lesion with central patchy depigmented areas with pigment mottling on the tumor surface (fig. 1a). Ultrasound B scan showed a highly reflective choroidal lesion with accompanying acoustic shadowing forming a pseudo-optic nerve appearance, classical of choroidal osteoma (fig. 2). SD-OCT images revealed disruption of the normal foveal architecture, with irregular RPE elevations with overlying thinning (c). The left eye showed normal appearance on examination and SD-OCT scanning (b, d).

Fig. 1. Fundus appearance on presentation showing a well-demarcated choroidal osteoma in the right macula with punctate dots of hyperpigmentation and central RPE thinning (a). SD-OCT images showed an irregularly elevated RPE layer with loss of normal foveal architecture. The RPE overlying the tumor showed some thinning. There was fragmentation of the photoreceptor inner and outer segment junction (arrow) suggesting photoreceptor damage, with irregular RPE elevations with overlying thinning (c). The left eye showed normal appearance on examination and SD-OCT scanning (b, d).
Discussion

Choroidal osteoma is a benign bone tumor, and like all living mature bone tissues, undergoes an active remodeling process. This is supported by the histopathological findings of osteocytes, osteoclasts and osteoblasts by Williams et al. [14] in enucleated eyes with choroidal osteoma. SD-OCT findings of cavernous or sponge-like features with variable internal reflectivity [9, 10] corroborate the presence of bony trabeculae and are suggestive of the various stages of the bone formation process occurring in choroidal osteomas. A common SD-OCT finding in choroidal osteoma is tumor elevation into the vitreous cavity [10–12, 15]. In our case report, we used fundus images to monitor tumor surface area changes and SD-OCT to monitor changes in tumor thickness. We report a similar finding of tumor elevation in the early stages, progressively evolving to central excavation as the tumor edge enlarges. Central...
tumor depression has also been observed previously [1, 6], and is believed to be caused by osteoclastic activity causing bone resorption. The increase in size and the formation of a central excavation suggests active bone remodeling in the underlying tumor in this case.

This remodeling process may or may not be associated with tumor progression over time. Progression of choroidal tumors has been documented to occur in 40–64% of cases [1, 3, 16]. In a series of 36 patients followed up for up to 22 years, Aylward et al. [3] observed that enlargement of choroidal tumors occurred in all directions in a uniform manner. No association was detected between growth of tumor and puberty. From our observation, the tumor growth spurt in this case occurred as the patient approached adolescence, suggesting that puberty may be a contributory factor. Possible reasons to consider hormonal influence in choroidal osteoma are its predominance in young females in the reproductive age group [1, 3, 6].

Fig. 4. Fundus photos taken annually documented tumor progression, which showed the tumor size increasing in all directions. Over the period of 5 years, SD-OCT images showed gradual flattening and eventual depression over the area of RPE elevation (arrows), which resulted in a crater formation in the central macula, suggesting that tumor remodeling was taking place. a–e Fundus photographs of the right eye taken yearly. f–j Corresponding SD-OCT images of the right macula. k–o RPE layer images from SD-OCT scanning of the right eye.
2], and reports of tumor occurrences or progression during pregnancy [6, 17] and in patients with thyroid diseases [18]. Transient elevation of serum parathyroid hormone and alkaline phosphatase levels has been reported during active growth of choroidal osteomas [19].

Choroidal osteoma is typically described as a slow-growing tumor [1, 3, 16], although there are several reported cases of rapid tumor growth [1, 4, 20]. Shields et al. [16] documented tumor growth at 0.37 mm per year by measuring the mean basal tumor diameter using indirect ophthalmoscopy. The tumor in this case showed a mean progression in surface area of 4,396.2 mm² per year. The rate of tumor growth reported here is more reliable as it is based upon serial fundus photos and calculated using an easily available and widely accepted image processing and analysis software (ImageJ). This method allows a more reliable, reproducible and comparable measurement of tumor growth. The main limitation of this method is that it only measures change in tumor surface area but not tumor depth, and therefore may not necessarily reflect the true tumor growth.

### Table 1. Calculation of tumor surface area growth

<table>
<thead>
<tr>
<th>Age, years</th>
<th>Surface area estimation, mm²</th>
<th>Surface area growth per year, mm²</th>
<th>Percentage of increase</th>
</tr>
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<tr>
<td>7</td>
<td>26,334</td>
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<tr>
<td>8</td>
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<td>36,400</td>
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<td>10</td>
<td>40,966</td>
<td>4,566</td>
<td>155.6</td>
</tr>
<tr>
<td>11</td>
<td>48,315</td>
<td>7,349</td>
<td>183.5</td>
</tr>
</tbody>
</table>

The tumor surface area estimation was measured by calibrating the vertical diameter of the optic disk to 1.5 mm in each fundus photo using the ImageJ software. The surface area of the tumor from each fundus photo taken during annual visits is calculated in square millimeters. The percentage of increase in tumor surface area was measured by comparing the yearly increase to the baseline in the year 2009. The mean tumor surface area growth over the period of 5 years was 4,396.2 mm²/year.

### References