Antimetropia in a 10-year-old boy with unilateral tilted disc syndrome

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Abstract: Antimetropia is a condition in which one eye is myopic, while the fellow eye is hyperopic. This report describes a case of antimetropia associated with unilateral tilted disc syndrome. A 10-year-old boy presented with the complaint of diminished vision for distant objects in his right eye for several months. His uncorrected distance visual acuity was 0.15 in the right eye and 1.0 in the left eye. Best corrected visual acuity was 1.0, with −6.00 D/−2.00 D 170° in the right eye and +1.50 D in the left eye. The cover test revealed right esotropia and hypertropia. Bifoveal fixation was achieved using a 14 prism diopter (PD) base at 220° in the right eye. Fundus examination revealed a tilted disc with inferior staphyloma in the right eye, but no abnormal findings were observed in the left eye. In addition, the patient occasionally experienced diplopia under casual viewing conditions. A cycloplegic refraction revealed −6.00 D/−2.00 D 170° in the right eye and +1.50 D/−2.00 D 5° in the left eye. Thus, prism glasses were prescribed. While wearing the prism glasses, the patient has shown no diplopia and maintains good binocular function.

Keywords: amblyopia, strabismus, anisometropia, staphyloma

Introduction
Anisometropia is a common condition characterized by an interocular difference in refractive error. Small degrees of anisometropia are inconsequential, but higher degrees are often accompanied by amblyopia in the more ametropic eye. In contrast, a significant degree of antimetropia, in which one eye is myopic and the fellow eye is hyperopic, is quite rare.1–4

Tilted disc syndrome is a relatively common congenital anomaly, occurring in 0.4% to 3.5% of the population, which consists of the rotation of the disc as a clockwise or counterclockwise rotation.5,6 Associated findings typically include an obliquely directed long axis of the disc, a posterior staphyloma, and situs inversus. The most important correlate of these lesions is a myopic astigmatism, most pronounced in the region of the staphyloma, and superotemporal or bitemporal visual field depression.5–8 The aim of this study was to report a case of a 10-year-old boy with antimetropia associated with unilateral tilted disc syndrome.

Case report
A 10-year-old boy presented with the complaint of diminished vision for distant objects in his right eye for several months. There was no significant ocular or medical history, and birth history was normal. His uncorrected distance visual acuity was 0.15 in the right eye and 1.0 in the left eye. Best corrected visual acuity was 1.0 with −6.00 D/−2.00 D 170° in the right eye and +1.50 D in the left eye. The cover test revealed right esotropia and hypertropia. Bifoveal fixation was achieved using a 14 prism diopter (PD) base at 220° in the right eye. Fundus examination revealed a tilted disc with inferior staphyloma in the right eye, but no abnormal findings were observed in the left eye. In addition, the patient occasionally experienced diplopia under casual viewing conditions. A cycloplegic refraction revealed −6.00 D/−2.00 D 170° in the right eye and +1.50 D/−2.00 D 5° in the left eye. Thus, prism glasses were prescribed. While wearing the prism glasses, the patient has shown no diplopia and maintains good binocular function.
D 170° in the right eye and +1.50 D in the left eye. The keratometry readings were 43.00 D at 172°/45.75 D at 82° in the right eye and 43.25 D at 107°/46.75 D at 100° in the left eye. The cover test revealed right esotropia and hypertropia. Bifoveal fixation was achieved using a 14 prism diopter (PD) base at 220° in the right eye. Fundus examination revealed a counterclockwise rotated disc with inferior staphyloma in the right eye (Figure 1A), but abnormal emergence of the retinal vessels (situs inversus) was not detected in the right optic disc. In contrast, no abnormal findings were observed in the left eye (Figure 1B). Ultrasonic B-scan echography (UD-8000; Tomey Corp, Nagoya, Japan) revealed inferior staphyloma in the right eye (Figure 2A), but no abnormal findings were observed in the left eye (Figure 2B). The axial length was 25.8 mm in the right eye and 23.2 mm in the left eye. The condition was diagnosed as antimetropia due to tilted disc syndrome. In addition, the patient occasionally experienced diplopia under casual viewing conditions. A cycloplegic refraction using 1% solution of atropine sulfate (three times per day for 7 days) disclosed −6.00 D/−2.00 D 170° in the right eye, and +1.50 D/−2.00 D 5° in the left eye. Prism glasses, with 7 PD base at 220° in the right eye and 7 PD base at 40° in the left eye, were prescribed. While wearing the prism glasses, the patient has shown no diplopia and maintains good binocular function.

**Discussion**

This report describes a patient with antimetropia, diplopia, and strabismus with tilted disc syndrome. To our knowledge, there are no cases in the literature presenting an antimetropia with tilted disc syndrome.

Tilted disc syndrome is characterized by an oval optic disc with an oblique axis, an inferonasal crescent, situs inversus of the retinal vessels, myopic astigmatism, and visual field defects. Dehghani et al. evaluated the ocular, refractive, and biometric characteristics in patients with tilted disc syndrome and compared them with a normal population. According to their report, there were no significant differences in spherical equivalent and total astigmatism between groups. However, mean best corrected visual acuity was significantly worse in the tilted disc syndrome patients. Furthermore, the lenticular astigmatism was greater in the syndrome group, whereas the corneal component was greater in controls. The measured biometric features were the same in both groups, except for the lens thickness and lens–axial length factor, which were greater in the tilted disc syndrome group.

Although other biometric data, including anterior chamber depth, lens thickness, vitreous chamber depth, and relative lens position were unavailable in this case, it is likely that the antimetropia was due to elongation of the axial length in unilateral tilted disc syndrome.

A significant degree of antimetropia with strabismus is extremely rare. Richards showed that motility evaluation revealed monofixation and that a small angle esotropia was common, occasionally accompanied by a vertical element. Additionally, in the same study, it was demonstrated that a switch in fixation sometimes occurs in strabismus with antimetropia when the preferred, less hyperopic eye evolves into a slightly myopic eye. A patient with compound hyperopic anisometropia most often prefers fixation with the less hyperopic eye and is usually free of diplopia because of the habitual suppression of the more hyperopic eye. However, when the less ametropic hyperopic preferred eye becomes myopic, the patient may switch to the previously suppressed and still hyperopic eye.

Although the previously preferred eye, ocular alignment, and changes in refractive error in this patient were unknown, the possible explanation for his condition is as follows: as the patient did not complain of diminished vision for far and near objects until the initial visit, gradual increasing myopic change in the right eye resulted in diminished vision for distant objects, and as a result of the deteriorated binocular function with the normal ocular alignment, strabismus with diplopia manifested.

Although the current findings are based on a single case, it is suggested that early detection and optical correction might be essential for amblyopia prevention and to attain good binocular function in patients with antimetropia.

Finally, most eyes with tilted disc syndrome reportedly have a good visual prognosis. However, when the upper temporal border of an inferior staphyloma lies across the macula, it is occasionally accompanied by macular complications that may lead to visual loss. Therefore, careful observation is
also necessary, for the early detection of several types of macular complications, on an inferior staphyloma associated with tilted disc syndrome.

**Disclosure**
The author reports no conflicts of interest in this work.

**References**