Unilateral high myopia: optical components, associated factors, and visual outcomes

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Aim: To elucidate the optical basis for unilateral high myopia and to identify the factors associated with its development.

Methods: Medical records of 48 children (aged 4 months to 17 years; mean age 6.8 years) with unilateral high myopia (5 dioptres or more) seen consecutively by the author during a 15 year period were reviewed. 45 (94%) of the 48 patients had unilateral axial myopia.

Results: The mean refractive difference between paired eyes was 9.4 (SD 3.6) dioptres and the more myopic eye was on average 3.3 (1.8) mm longer than the less myopic eye. All but three of the patients had an ocular disorder associated with reduced acuity, central nervous system abnormality, or family history of high myopia.

Conclusion: Clinical conditions associated with unilateral high myopia can be identified in the majority of patients and often account for the associated visual impairment.

Unilateral high myopia is an uncommon condition, unlike myopia in general which is found in 15% of children by age 15 years. Although the eye is too long for its optical power in myopia, the two eyes typically grow by the same amount resulting in symmetric myopia. The presence of asymmetric myopia indicates that either the axial growth or the combined refractive power of the cornea and lens is different between eyes. Axial and refractive myopia can be distinguished on the basis of measurements of corneal power and total axial length. Few published reports have addressed the optical basis for anisometropic myopia derived from these measurements. Sorsby et al inferred from measurements of refractive power in 68 patients that increased axial length was the major cause of anisometric myopia.

Most other studies of anisometropic myopia have focused on treatment and visual outcomes. Clinical findings have not been emphasised except by Rosenthal et al who reported fundus abnormalities (staphyloma and pigmentary disturbances) or cataract in seven of 53 patients. But conversely, numerous studies have stressed the increased incidence of axial myopia in monocularly deprived animals and in human infants with the unilateral presence of blepharoptosis, corneal opacification, cataract, or vitreous haemorrhage. A higher incidence has also been reported with medullated nerve fibres and optic nerve hypoplasia. Whether there is a significant prevalence of these predisposing factors in unselected patients with anisometropic myopia is unknown.

A consecutive series of patients with unilateral high myopia were evaluated with the intent of identifying associated clinical conditions and learning more about the mechanisms that underlie unilateral axial myopia and visual impairment.

PATIENTS AND METHODS

The study sample consisted of all children with unilateral high myopia, seen consecutively over a 15 year period. Unilateral high myopia was defined as relative myopia of at least 5 dioptres with absolute myopia of at least 3.5 dioptres. Since measurements of total axial length (TAL) and corneal power have been a routine part of my examination, the majority of patients with unilateral high myopia were entered into this study. Patients who had undergone cataract removal or scleral buckling procedures were excluded.

All patients had complete eye examinations. Refractions were determined 40 minutes after instilling one or two drops of cyclopentolate. (Cyclopentolate 0.5% was used in infants less than 6 months of age, 1.0% was used in children between the ages of 6 months and 2 years, and 2% was used in older children.) Total axial length (TAL) was measured by A-scan ultrasonography (Ophthalmic; Biophysics Medical, Clermont-Ferrand, France). Corneal power was measured with a table mounted keratometer (CLC Ophthalmometer; American Optical, Buffalo, NY, USA), or hand held keratometer (MK 500; Nidek, Tokyo, Japan). All measurements of TAL and corneal power were done in awake children in the outpatient clinic.

Axial myopia was defined as myopia which was due to an interocular difference in TAL. Refractive myopia was defined as myopia which was due to an increased refractive power of the cornea, lens, or both. Axial anisometropic myopia was considered to be absolute when the TAL of the more myopic eye was above the normal range and relative when the TAL of the more myopic eye was within the normal range or below.

Visual acuities were measured using optotypes, Allen cards, or Teller acuity cards (TACs) in 40 patients depending upon age and ability level. To provide a common measure and linear scale, acuities were transformed into the logarithm of the maximal angle of resolution in minutes of arc (logMAR) where 20/20 = 0.0, 20/200 = 1.0, etc. Acuities of 20/2000 or less were arbitrarily assigned a value of 2.0.

Ten patients in this series were treated for amblyopia with occlusion therapy. The remaining 38 patients were not treated for amblyopia because they were visually mature (8 years of age or older) or the underlying ocular abnormality was severe enough to account for the reduced vision.

RESULTS

Forty-eight patients (23 males, 25 females) ranging from 4 months to 17 years of age (mean age 6.8 years) satisfied the selection criteria. The more myopic eye was the right eye in 21 patients. Forty-five of the 48 patients had axial myopia and three had refractive myopia.

Anisometropic myopia was also categorised on the basis of the refraction in the fellow eye. Twelve of the patients had anisotropic bilateral myopia where both eyes are myopic; 19
had mixed anisometropic myopia where one eye is myopic and the other eye is hyperopic (1 dioptre or more); and 17 had anisometropic unilateral myopia where one eye is myopic and the other eye is emmetropic. Figure 1 shows the bilateral refractions in dioptres for the 42 patients who were aged 10 years or younger.

Twenty seven (56%) of the patients had esotropia (16) or exotropia (11). The remaining 21 patients were orthophoric or had microstrabismus. Binocularity was reduced or absent in all but one patient in whom it was measured.

Corneal power was measured bilaterally in 24 (50%) of the 48 patients. Corneal power (in spherical equivalent) ranged from 40.0 to 49.4 dioptres (mean 44.4 dioptres). The mean (SD) for the interocular difference in corneal power was 0.1 (0.1) dioptres.

Figure 2 shows the bilateral refractions in dioptres for the 42 patients who were aged 10 years or younger.

Figure 1 Refractions in dioptres for 42 patients with unilateral high myopia. Hyperopic refractions are represented by unsigned values and myopic refractions by minus (–) values.

Figure 2 Total axial length (TAL) values in 42 patients with unilateral high myopia. For each pair of eyes the circle represents the total axial length of the more myopic eye and the square represents the total axial length of the less myopic eye. The mean total axial length values for normal subjects (middle curve) and upper and lower 95% confidence bands (outer curves) are shown for comparison.

Figure 3 The refractive difference between the more myopic eye and less myopic eye in dioptres is compared with the difference in their total axial length in millimetres.

Three (6%) of the patients had unilateral refractive myopia associated with lens abnormalities. One of these three patients had a previous corneal laceration complicated by lens subluxation and segmental cataract. A second patient had lens subluxation following blunt ocular trauma. A third patient had a unilateral malformation of the anterior segment with a thickened lens compared to the other eye (7.9 mm versus 4.1 mm). The mean refractive difference between the myopic eye and the less myopic eye was 8.4 dioptres (range 7.5–9.7). In each patient the interocular refractive difference was due to an increase in refractive power of the lens. The mean interocular difference in TAL was 0.1 mm (range 0.0–0.2) and corneal power of both eyes was equal.

Ocular disorders, systemic abnormalities, or relevant family history which were found to be associated with unilateral high myopia are listed in Table 1. Relevant factors were identified in all but three patients. Fifteen (31.3%) patients had optic nerve abnormalities including hypoplasia (10), myelinated nerve fibres (three), atrophy (one), and coloboma (one). Optic nerve hypoplasia was segmental in four patients, diffuse in six patients. Ten (20.8%) of the patients had CNS abnormalities in (86.7%) patients or below in six patients. In other words, the anisometropic axial myopia was absolute in 57.8% of patients and relative in the remaining 42.2%.

Figure 3 compares the refractive difference in dioptres between eyes with the corresponding difference in TAL in millimetres. The mean refractive difference was 9.4 (SD 3.6) dioptres (range 4.5–18.2 dioptres). The more myopic eye was an average 3.3 (1.8) mm (range 0.8–8.9 mm) longer than the less myopic eye. A slope of 0.34 for the regression line indicates that there is a 2.9 dioptre refractive difference for each 1 mm of TAL difference. Thus, for all patients, the interocular refractive difference was predominantly accounted for by the interocular difference in TAL ($r^2 = 0.62; p < 0.001$).

Table 1 Factors associated with development of unilateral high myopia

<table>
<thead>
<tr>
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<th>Number of patients (%)</th>
</tr>
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<tr>
<td>Optic nerve disorder</td>
<td>15 (31.3)</td>
</tr>
<tr>
<td>CNS abnormality</td>
<td>10 (20.8)</td>
</tr>
<tr>
<td>Lens abnormality</td>
<td>6 (12.5)</td>
</tr>
<tr>
<td>Retinopathy of prematurity</td>
<td>5 (10.4)</td>
</tr>
<tr>
<td>Family history of high myopia</td>
<td>3 (6.3)</td>
</tr>
<tr>
<td>Buphthalmos</td>
<td>3 (6.3)</td>
</tr>
<tr>
<td>Macular scar/chorioretinal coloboma</td>
<td>2 (4.1)</td>
</tr>
<tr>
<td>Congenital ptosis</td>
<td>1 (2.0)</td>
</tr>
<tr>
<td>None*</td>
<td>3 (6.3)</td>
</tr>
</tbody>
</table>

*One of these patients had nasomaxillary dysplasia (Binder syndrome).

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early life related to perinatal hypoxic ischaemic injury (four), meningitis (two), trauma (two), congenital hydrocephalus (one), or bilateral occipital lobe infarctions (one). Lens abnormalities were implicated in six (12%) patients, three with axial myopia and three with refractive myopia (see preceding paragraph). Of those with axial myopia, two had fetal nuclear cataracts and one had posterior lenticonus. Three (6.6%) of the patients had buphthalmos, two with unilateral or asymmetric glaucoma and one with neurofibromatosis. There was a family history of bilateral high myopia in three patients, one with hemifacial microsomia (case 3) and one with a presumed defect in unilateral emmetropisation (case 1). One of the patients without a known associated condition had nasomaxillary dysplasia (Binder syndrome).

Five patients were born prematurely (mean gestational age 26.6 weeks) and had a history of retinopathy of prematurity (ROP). The TAL of the longer eye was within the normal range in two patients and below in three patients.25 The more myopic eye was on average 2.8 mm (range 1.4–5.3 mm) longer than the less myopic eye. Corneal powers of paired eyes were within 1.5 dioptres of each other. All of the patients had ROP residua in the posterior pole. The relation of anisometric myopia to total axial length and ROP residua is shown in Table 2.

Acuities were bilaterally measured with optical correction in 40 patients using the Snellen test (32), Allen cards (six), or the Teller acuity card procedure (two). Data from the remaining eight patients in whom acuities were not measured objectively were excluded. Figure 4 is a scatter plot of the acuity data. For 34 of 40 patients, acuity is worse in the more myopic eye with 30 of those having acuities between 20/20 and 20/40 (0.0 to 0.3 logMAR) in the less myopic eye and 20/200 to 20/2000 (1.0 to 2.0 logMAR) in the more myopic eye.

The following case reports are presented because each implicates a novel mechanism associated with asymmetric eye growth.

**Case 1**

A 2 month old girl enrolled in a longitudinal study of visual development was noted to have anisometric hyperopia (+0.50 + 0.50 × 60 right eye; +3.75 + 0.75 × 130 left eye). Visual acuity was 20/130 right eye and 20/180 left eye using

### Table 2 Relation of anisometric myopia to visual acuity, total axial length, and ROP residua

<table>
<thead>
<tr>
<th>Case</th>
<th>Age (years)</th>
<th>Sex</th>
<th>Eye</th>
<th>Visual acuity</th>
<th>Refraction (dioptres)</th>
<th>TAL (mm)</th>
<th>ROP residua</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>0.8</td>
<td>F</td>
<td>R</td>
<td>CSM</td>
<td>–8.50 + 1.50 × 45°</td>
<td>18.4</td>
<td>Macular heterotopia</td>
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<tr>
<td></td>
<td></td>
<td></td>
<td>L</td>
<td>UCUSUM</td>
<td>–1.50 + 1.00 × 180°</td>
<td>15.0</td>
<td>Retinal fold</td>
</tr>
<tr>
<td>B</td>
<td>2.3</td>
<td>M</td>
<td>R</td>
<td>CSM</td>
<td>–5.75 + 1.00 × 180°</td>
<td>19.2</td>
<td>Normal</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>L</td>
<td>UCUSUM</td>
<td>+0.50</td>
<td>17.8</td>
<td>Macular heterotopia</td>
</tr>
<tr>
<td>C</td>
<td>4.5</td>
<td>F</td>
<td>R</td>
<td>UCUSUM</td>
<td>+1.00 + 1.00 × 105°</td>
<td>15.8</td>
<td>Retinal fold</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>L</td>
<td>CSM</td>
<td>–4.00 + 1.00 × 90°</td>
<td>17.8</td>
<td>Normal</td>
</tr>
<tr>
<td>D</td>
<td>5.0</td>
<td>F</td>
<td>R</td>
<td>5/30</td>
<td>–9.50</td>
<td>23.1</td>
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<td></td>
<td></td>
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<td>+0.75</td>
<td>21.2</td>
<td>Normal</td>
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<tr>
<td>E</td>
<td>3.5</td>
<td>F</td>
<td>R</td>
<td>10/30</td>
<td>–4.00</td>
<td>23.8</td>
<td>Macular heterotopia</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>L</td>
<td>1/30</td>
<td>+4.50</td>
<td>18.3</td>
<td>Previous exudative retinal detachment</td>
</tr>
</tbody>
</table>

*CSM = central, steady, and maintained fixation. UCUSUM = uncentral, unsteady, and unmaintained fixation.

![Figure 4](http://bjophthalmol.com)

**Comment**

Unilateral high myopia was presumably due to a primary defect of emmetropisation in one eye.

**CASE 2**

A previously healthy 6 month old child was initially evaluated for a bilateral occipital lobe infarction following blunt head trauma. Falling between two bedposts, the infant crushed his
Case 3

A healthy 13 year old child with congenital esotropia was referred for evaluation of residual esotropia and anisometropic myopia. She was born at term; growth and development were normal. Family history was strongly positive for high myopia. She was born at term; growth and development were age appropriate. A maternal great grandmother had bilateral high myopia. Acuity was 20/20 at both eyes. Refraction was +2.00 + 1.00 at 90° right eye and −6.00 + 1.25 at 90° left eye. Motility examination revealed a 40 dioptre esotropia. Slit lamp examination was normal. Fundus examination showed peripapillary retinal tissue. Keratometry readings were 45.0 dioptres bilaterally. Pattern reversal VEPs were approximately 21.0 mm right eye and 24.9 mm left eye. Pattern reversal VEPs were 22.9 mm right eye and 24.5 mm left eye. Keratometry readings were 44.00 by 45.75 bilaterally.

Comment

Unilateral high myopia in a child with a strong family history of bilateral high myopia suggests that hemifacial microsoma may have offset the genetic propensity of the ipsilateral eye to grow excessively.

Case 4

An 8 year old boy was seen in consultation for an abnormal appearing left optic disc with reduced acuity. He was born at term, and development was age appropriate. A maternal great grandmother had bilateral high myopia. Acuity was 20/20 right eye and 20/400 left eye. Refraction was +0.50 right eye and −6.62 left eye. There was a relative left afferent pupillary defect. Fundus examination demonstrated segmental optic nerve hypoplasia left eye and normal sized right optic disc (see Fig 5). TAL values were 21.9 mm right eye and 24.1 mm left eye. Keratometry readings were 45.0 dioptres bilaterally.

Comment

The association of optic nerve hypoplasia with unilateral high myopia implicates local retinal and/or central brain mechanisms in the control of ocular growth.

DISCUSSION

We found that 94% of children with unilateral high myopia (5 dioptres or more) had axial elongation of the more myopic eye as the cause of myopic anisometropia confirming the results of Sorsby. Comparing the TAL of the more myopic eye with age matched normative data, we found that the anisometropia was absolute in 57.8% of patients and relative in the remaining 42.2%. Only three (6%) of the patients had unilateral high myopia on a refractive basis. Two of the three patients had lens subluxation following blunt trauma. None of these patients had unilateral high myopia resulting primarily from asymmetric increases in corneal power. Excluding patients with eye trauma, all but one patient had unilateral high myopia due to axial length differences.

The higher prevalence of high myopia in children born to highly myopic parents and Mendelian inheritance pattern in some families with high myopia suggest there is an underlying genetic defect in this condition. Why a genetic defect in the growth plan of the eye would lead to unilateral high myopia is puzzling. One possible explanation for the dissimilar growth between eyes is skewed lyonisation in females heterozygous for X linked myopia. Significantly, each of the three patients in this series with a family history of high myopia were females. Selective inactivation of one parental chromosome in each eye progenitor may lead to predominant expression of the myopia gene in one eye but not in the fellow eye. Precedence for skewed X chromosome inactivation in the eye can be found in monozygotic twins and eye pairs of individuals discordant for colour vision deficiency. Alternatively, local environmental factors may be important. For example the patient (case 3) with unilateral hemifacial microsoma may have inherited the propensity to high myopia bilaterally but the facial deformation may have influenced the growth of the ipsilateral eye, thereby cancelling the myopia.

Beginning with the work of Wiesel and Raviola on axial myopia induced by form deprivation, environmental factors have been shown to influence how the eye grows. Experimental animals deprived of form vision by suturing the eyelid shut or optical defocusing and human counterparts with ptosis, media opacities can develop ipsilateral axial myopia. Reduction in the quality of visual inputs presumably provides a weak feedback signal to the central growth control mechanism, thereby resulting in excessive axial elongation under certain conditions. That over 40% of our patients had an obstruction within the visual axis (ptosis, cataract) or anatomic defect of the lens, optic nerve, or macula implicates a vision dependent mechanism.
Although the results presented here focus on the relation between monocular visual deprivation and axial myopia, degradation of visual input is also associated with axial hyperopia. These data suggest that the sign of the induced refractive error is influenced by severity of visual deprivation and postnatal age. For example, Bradley et al have shown in rhesus monkeys that treatment of one eye with a diffuser contact lens retards the growth of the treated eye. In this experiment, the diffuser lens results in a loss of contrast sensitivities across all spatial frequencies relative to severe form deprivation, which differentially degrades high spatial frequency inputs. Moreover, this study highlights the impact of age by showing that visual deprivation from birth induces axial hyperopia. In comparison, unilateral form deprivation during the infantile phase of ocular growth or adolescence tends to be associated with excessive axial elongation.

It is important to mention that non-visually dependent mechanisms may be involved in the regulation of ocular growth. Local post-receptor mechanisms seem to be important. For example, with disorders of the optic nerve and macula the loss of ganglion cells or other retinal cell types have been implicated in the disturbance of local growth mechanisms.

A large percentage (31%) of patients with anisometropic myopia had abnormalities of the optic nerve of which hypoplasia was the most frequent. The high prevalence of unilateral optic nerve hypoplasia in this study compared to previous studies suggests that in the presence of high axial myopia, the concurrent presence of optic nerve hypoplasia is underappreciated. Distinguishing the hypoplastic disc from the optic disc distorted by high myopia can be difficult, especially when the optic disc margin is poorly demarcated. Furthermore, the hypoplastic disc appears larger when viewed through the ophthalmoscope, owing to the magnification effect that was species dependent. Despite these inconsistencies, they concluded that axial myopia is caused by an alteration of visual inputs and mediated by the nervous system. By the same mechanism, distortion of visual inputs by the CNS abnormalities described in our study may lead to axial myopia.

One patient (case 1) with longitudinal data provided important insights into the mechanism that regulates emmetropisation during visual development. Initially this infant was found to have anisometropic hyperopia in the left eye (+0.50 right eye + 3.75 left eye) which evolved within a few months without treatment into anisometropic myopia in the left eye (−0.50 right eye −3.50 left eye). One interpretation of this observation is that growth of the left eye lagged initially and then accelerated, overshooting the amount required for emmetropisation. That is, there seemed to be a primary defect in the process of emmetropisation in the affected eye. Why this growth disturbance occurred unilaterally is unclear but the fact that concurrent acuity differences paralleled the refractive differences suggest the underlying mechanism is vision dependent. The same scenario might apply to other cases of "idiopathic" unilateral axial myopia where the eye is otherwise normal and there is no history of visual deprivation.

Congenital glaucoma was readily identified as another predisposing factor in two patients. In the immature eye where VEPs are reported to be abnormal. These visual deficits suggest that the nerve axons and/or visual connections are functionally defective. During development axon signals induce progenitor cells invading the optic nerve to differentiate into type 2 astrocytes or oligodendrocytes. Developmental abnormalities of nerve axons may lead to faulty differentiation of these precursor cells into oligodendrocytes resulting in ectopic production of myelin within the retinal ganglion cell layer. Myelinated nerve fibres are probably an epiphenomenon caused by the same axonal abnormality that underlies reduced visual function and leads to axial myopia.

Central nervous system abnormalities were the most frequent predisposing factor after optic nerve abnormalities. This association suggests the central nervous system influences eye growth but the literature on this subject is sparse. Raviola and Wiesel hypothesised that striate cortex receives visual inputs and provides feedback signals to subcortical visual mechanisms which, in turn, exert control over eye growth. However, in monkeys, surgical removal of both occipital lobes had no abortive effect on lid suture induced myopia, and pharmacological paralysis of accommodation (with atropine) or stimulation (with isofluorophate) had an inconsistent effect that was species dependent. Despite these inconsistencies, they concluded that axial myopia is caused by an alteration of visual inputs and mediated by the nervous system. By the same mechanism, distortion of visual inputs by the CNS abnormalities described in our study may lead to axial myopia.
the cornea and sclera are still distensible, one eye may differentially elongate due to unilateral or asymmetric increases in intracocular pressure. However, the severity of myopia may not reflect the full amount of axial elongation as the myopia is partially reduced by corneal flattening or posterior displacement of the lens. The possibility that corneal haze or glaucomatous optic nerve damage predisposed to axial elongation on the basis of degraded visual inputs can not be excluded.

Anisometropic myopia is a common sequela of ROP. The incidence of anisometropia (defined as an interocular refractive difference of 2 dioptres or more in spherical equivalence) in the CRYO-ROP study was as high as 49% (57 of 117) among the subset of infants with myopia and active ROP in zone 2 (stage 3) or zone 1. Previous studies including axial length measurements have reported shortened, normal or elongated lengths of ROP eyes. Since TAL is usually normal or abnormally short in ROP eyes, the myopia has been primarily attributed to increases in lens power. For five patients with ROP in this series the more myopic eye was normal size or small but still longer than the fellow eye, indicating that TAL difference was the principal cause of anisometropic myopia. Thus, the axial elongation of the more myopic eye was only relative to the fellow eye which was abnormally small and included ROP residue in about five of five patients. Taken together these findings suggest that the severe anisometropic myopia sometimes associated with ROP is primarily due to the reduced growth of the less myopic eye. Furthermore, the more myopic and longer eye tended to have the better visual acuity and more normal appearing macula.

Similar to previous studies which reported visual acuities of 20/100 or worse in 50–85% of patients with anisometropic myopia, 75% of the patients in this study had acuities of 20/200 or worse in the affected eye. Reduced acuities were usually attributed to anisometropic amblyopia and coexisting ocular abnormalities. Since the mean age of our patients' parents (7 years) placed them close to visual maturity, this study cannot adequately address the potential benefits of spectacle correction and amblyopia treatment. Pollard et al reported visual acuities of 20/40 or better in 22 of 40 patients with unilateral high myopia after amblyopia treatment. By comparison other investigators have reported that visual loss is refractory to treatment in the majority of patients.

In summary, nearly 90% of the patients with axial myopia had an associated abnormality of the eye, central nervous system or family history of high myopia. In previous studies, excluding those devoted to specific conditions, little attention has been given to the clinical background in which unilateral high myopia appears. Such a high prevalence of associated pathology in the present series suggests that axial myopia is usually a consequence of a pre-existing abnormality. For patients with a history of a traumatic lens injury, congenital glaucoma, or ROP, the cause for the unilateral high myopia is usually clear. Finding unilateral axial myopia in a patient without a relevant ocular history should prompt the ophthalmologist to look critically for predisposing ocular defects (especially optic nerve disorders) and to carefully consider the perinatal and family history. In an attempt to systematize the diagnostic evaluation of the patient with unilateral high myopia, I propose an algorithm based on my experience (Fig 6).

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