

**Clinical Image**

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**Lens coloboma secondary to neglected buphthalmos****\*Corresponding Author: Mousumi Banerjee**Dr Rajendra Prasad Centre for Ophthalmic Sciences,  
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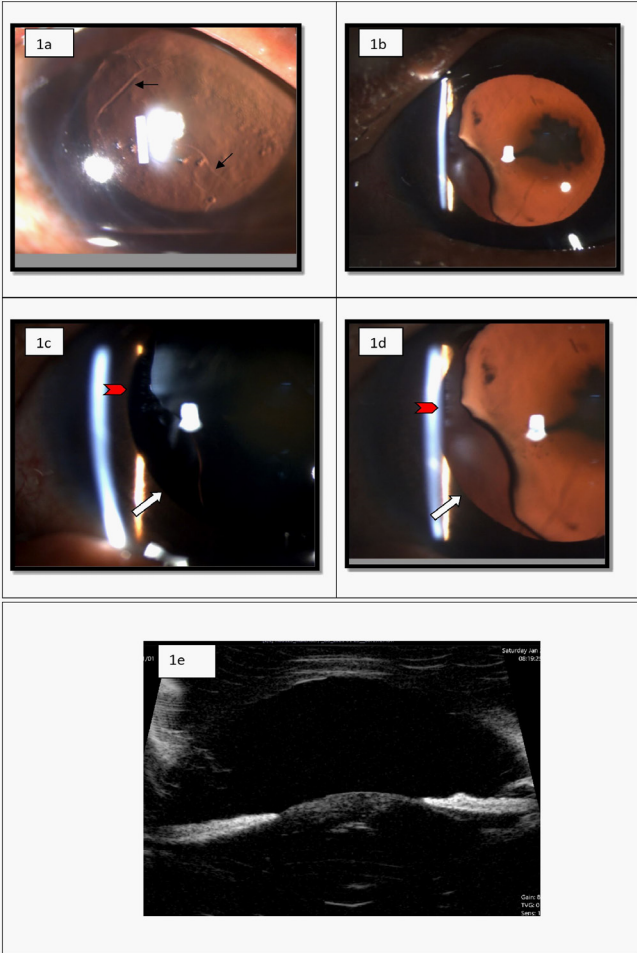
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**Clinical image description**

A 36-year-old male presented with progressive diminution of vision OS for 3 months. He was a known case of bilateral congenital glaucoma who underwent trabeculectomy at 6 months and 5 years of age OD and OS respectively. Best corrected visual acuity was 6/24 OD and 1/60 OS. Slit lamp examination revealed bilateral Haab striae (Figure 1a) with lens subluxation OS evident by the scalloped border of the lens with broken zonules in the superonasal quadrant and lens coloboma in the inferonasal quadrant with the absence of zonules (Figure 1b,c,d). A posterior subcapsular cataract was also noted OS. Advanced glaucomatous optic nerve cupping was noted OU. An intraocular pressure of 10 mm Hg OD and 16 mm Hg OS was noted. Biometry documented an axial length of 26.30 mm OD and 28.75 mm OS with a keratometry of 42.50D/46.50D @20°/110°OD and 37.75D/40.00D @ 45°/135°OS. Ultrasound bio-microscopy depicted increased sphericity of the lens with broken zonules OS (Figure 1e).

Lens coloboma is a misnomer as it does not indicate actual absence or defect in lenticular substance. It develops due to absence of zonules which lead to localized loss of tension on lens capsule, resulting in retraction of the lens and formation of a notch or fissure [1,2]. It is technically a coloboma of the zonule and not of the lens [3]. Lens coloboma is a congenital anomaly associated with ocular anomalies including iris, choroid or optic disc coloboma [4] or might be one of the manifestations of systemic diseases, including Marfan syndrome and Marshall syndrome [5]. In this case, coloboma is more likely to be an acquired condition secondary to uncontrolled globe enlargement OS as evident by the neglected buphthalmos till 5 years of age leading to excessive zonular stretching. Zonular discontinuity led to the scalloped border of the lens superonasally and absence of zonules inferonasally led to flattening of the lens giving a coloboma like appearance. Thus, lens coloboma can also be a sign of uncontrolled primary congenital glaucoma apart from extreme buphthalmos. Routine ophthalmic screening is required in such cases to prevent lenticular astigmatism and subsequent amblyopia.



**Figure 1:** Clinical photograph of Haab striae OS depicted by black arrows (1a). Figure 1 b depicts the superonasal scalloped border of the lens with broken zonules and inferonasal lens coloboma with absent zonules on retro –illumination. A central opacity depicting posterior subcapsular cataract can also be noted. Figure 1 c, d show the enlarged view of the lens anomaly with broken zonules superonasally (red arrowhead) and absent zonules inferonasally (white arrow) on indirect and retro-illumination respectively. Ultrasound bio-microscopy reveals increased sphericity of the lens with localised absent zonules nasally (1e).

### Learning points

1. Lens coloboma can be a sign of uncontrolled primary congenital glaucoma.
2. Uncontrolled enlargement of the globe secondary to neglected glaucoma can lead to excessive zonular stretching and formation of lens coloboma.
3. Surgical treatment is required if coloboma causes significant lenticular astigmatism leading to amblyopia, associated cataract or if the lens edge bisects the pupil.

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