

## Clinical Image

## Cerulean Cataract

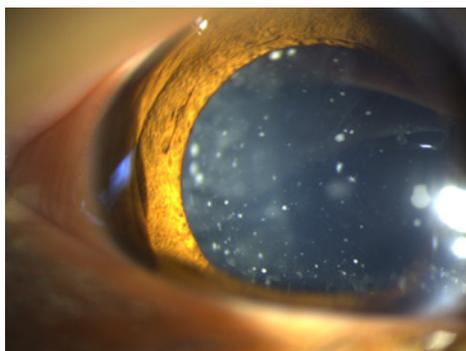
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## Clinical Image

A 14-year-old girl presented with a two years history of gradual decrease of vision in both eyes. The best corrected visual acuity was 0, 3 LogMAR in both eyes. The examination of the anterior segment on slit-lamp of both eyes revealed multiple tiny bluish-white opacities distributed in the lens nucleus and cortex in the form of concentric circles corresponding to a congenital cerulean cataract.



**Figure 1:** Diffuse light slit lamp photograph of the left eye showing multiple bluish-white opacities spread throughout the cortex of lens.

No other abnormality was observed in the slit lamp in both eyes. Phacoemulsification surgery was planned for each eye with a good evolution.

Cerulean cataract, also known as blue dot cataract, is a rare phenotypic variant of congenital cataract, first described by Vogt [1]. Cerulean cataracts are inherited as an autosomal dominant trait [2]. It is a developmental cataract characterized by bluish-white opacifications scattered in the nucleus and cortex of the lens [3]. Patients are usually asymptomatic until the age of 18–24 month.

## Disclosure of Interest

The authors declare that they have no competing interest.

## References

1. Francis PJ, Berry V, Bhattacharya SS, Moore AT. The genetics of childhood cataract. *J Med Genet.* 2000; 37: 481–8.
2. Litt M, Carrero-Valenzuela R, LaMorticella DM, et al. Autosomal dominant cerulean cataract is associated with a chain termination mutation in the human beta-crystallin gene CRYBB2. *Hum Mol Genet.* 1997; 6: 665–8.
3. Ram J, Singh A. Cerulean cataract. *QJM.* 2019; 37.