Response to comment on: Bilateral idiopathic spontaneous filtering bleb with ectopia lentis: A case report and review of literature

Dear Sir,

We thank for the interest shown in our article and for sharing the details of your patient with similar clinical picture.^[1,2] We agree that the facial dysmorphic features are subtle in Traboulsi syndrome or facial dysmorphism, lens dislocation, anterior-segment abnormalities, and spontaneous filtering blebs, which is due to mutation in aspartyl/ asparaginyl β -hydroxylase.^[3] We compared our patient's facial features [Figs. 1a and b] with the pictures of patients diagnosed with Traboulsi syndrome,^[4-6] and our patient's facial feature is not as typical as the ones mentioned in the literature. Taking the entire clinical picture into account, he may be a patient with Traboulsi syndrome but needs genetic testing to confirm the diagnosis.

Declaration of patient consent

The authors certify that they have obtained all appropriate patient consent forms. In the form the patient(s) has/have given his/her/their consent for his/her/their images and other clinical information to be reported in the journal. The patients understand that their names and initials will not be published



Figure 1: Profile picture of the patient, front (a) and side (b) view

and due efforts will be made to conceal their identity, but anonymity cannot be guaranteed.

Financial support and sponsorship Nil.

Conflicts of interest There are no conflicts of interest.

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Access this article online	
Quick Response Code:	Website: www.ijo.in
	DOI: 10.4103/ijo.IJO_249_18

Cite this article as: Chandran P, Khairnar AS, Aboobacker N, Raman GV. Response to comment on: Bilateral idiopathic spontaneous filtering bleb with ectopia lentis: A case report and review of literature. Indian J Ophthalmol 2018;66:605-6.

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