Axenfeld-Rieger Syndrome

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Received date: May 21, 2018; Accepted date: May 24, 2018; Published date: May 31, 2018

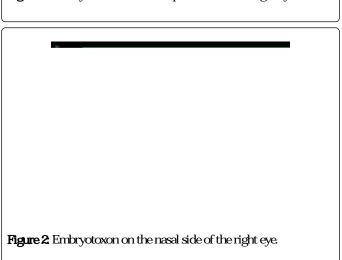
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We report the case of a patient aged 47 who consult for a decrease in visual acuity. Examination of the anterior segment spotting unilateral irrido-trabecular dysgenesis of the right eye with abnormal visibility of the schwalbe line corresponding to a posterior embryotoxon and associated angular abnormalities (Figures 1-3).

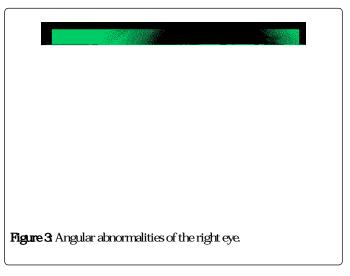
e eye tone measured with Goldmann tonometer showed 14 mmHg in the right eye and 15 mmHg in the 'e eye e examination of the fundus of the eye bds symmetrical morphology of the optic discs without pathological papillary excavation.



Figure 1: Embryotoxon on the temporal side of the right eye.



e remainder of the somatic examination reveals no abnormality associated especially the absence of dental malformation. Chronic glaucoma is seen in 50% of patients [1]. e diagnosis of Axenfeld-Reiger syndrome uncomplicated of chronic glaucoma has been established, despite the absence of signs of Rieger [1]. No treatment has been established. Regular checks have been proposed to detect any complications including glaucoma.



Is syndrome is inherited as an autosomal dominant manner. It is found that 2 genes are mainly involved in the transmission; *PITX2* gene in 4q25, present in 10-60% of patients, mainly associated with systemic alterations such as dental malformations [2,3].

e other gene responsible is FOXC1 located in 6q25, present in 50% of cases and manifested by ocular alterations, especially glaucoma [2-4]. e d] erebt]U diagnosis arises with the Peters anomaly which consists of a defect of the posterior surface of the cornea associated with a stromal opacity. Currently, it is sugge all