A Case Report: Bilateral Optic Pit with Its Inferonasal Location in Right Eye

Mahmut Erkam Arslan, Emine Pangal, Cemal Özsaygılı, Süleyman Demircan and Ayşe Çiçek*
Department of Ophthalmology, Kayseri Training and Research Hospital, Kayseri, Turkey

Abstract
Optic Disc Pit (ODP) is a rare congenital anomaly which is seen approximately in 1/11,000. Optic disc pits are bilateral in up to 10% to 15% of cases. ODP has been observed temporally in 70%, centrally in 20% and inferioirly in 10% of cases. We present a bilateral ODP case with atypical inferonasal presentation at right eye.

Introduction
Purpose
To present clinical features of a bilateral optic pit case.

Methods
A 30-year-old woman with bilateral optic pit who was admitted to our clinic for a routine ophthalmologic examination has been presented with her fundus photos, Fundus Fluorescein Angiography (FFA), Optical Coherence Tomography (OCT) and perimetry results.

Case Presentation
In detailed ophthalmologic assessment; best-corrected visual acuity was 20/20 in both eyes, anterior segment examination was unremarkable, intraocular pressures were recorded as 22 mmHg OD and 21 mmHg OS (with Goldmann applanation tonometry). Central corneal thicknesses were 639 μm OD and 638 μm OS. Corrected intraocular pressures were 16 mmHg in right eye and 15 mmHg in left eye. Dilated fundus examination of right eye revealed an optic pit located inferonasal to the optic disc. Fundus examination of left eye revealed an optic pit located temporally. FFA showed hypofluorescence at nasal inferior of the right optic disc and at temporal of the left optic disc. FFA showed no hyperfluorescence at macula or other retinal areas on both eyes. Loss of retinal tissue at optic pit areas was observed in OCT image at the level of optic nevre head (this was correlated with optic disc pit). The patient was informed to apply for medical examination in case of vision loss. The examination findings of patient were stable at follow up examination after 2 months. The ophthalmologic examination was advised for every 6 months.

Result
Optic Disc Pit (ODP) is a rare congenital anomaly which is seen approximately in 1/11,000 [1] Histologically, ODP is a herniation of dysplastic retina into a collagen-rich excavation that extends into the subarachnoid space through a defect in the lamina cribrosa [2]. It effects males and females equally [1]. Optic disc pits are bilateral in up to 10% to 15% of cases. ODP has been observed temporally in 70%, centrally in 20% and inferioirly in 10% of cases [3,4]. Our case is interesting because of its bilaterality and inferonasal location of optic disc pit in right eye. The patients with ODP should be followed up for maculopathy. It should be noted that ODP maculopathy could effect both eyes at different times asymetrically.

References