

A Macular Detachment Hiding an Optic Disk Pit

Ouidad Elbaz^{1*}, Younes Tlemcani¹, Sarah Belghmaidi¹, Ibtissam Hajji¹, Abdeljalil Moutaouakil¹

¹Ophthalmology Department, CHU Mohamed VI, av Ibn Sina, Marrakech, Maroc

DOI: <https://doi.org/10.36347/sjmcr.2025.v13i05.027>

| Received: 11.03.2025 | Accepted: 16.04.2025 | Published: 13.05.2025

*Corresponding author: Ouidad Elbaz

Ophthalmology Department, CHU Mohamed VI, av Ibn Sina, Marrakech, Maroc

Abstract

Case Report

Introduction: Optic disk pit is a rare congenital anomaly, susceptible to macular complications such as serous retinal detachment of the posterior pole. Its pathogenesis remains controversial, as does its treatment. **Case Report:** A 32-year-old patient presented with a progressive decline in visual acuity in her right eye over the past three years. Visual acuity was 3/10 in the right eye and counting fingers at 1 m in the left eye. Funduscopy revealed a temporal Optic disk pit on the right eye associated with a macular serous retinal detachment. On the left, a chorioretinal coloboma was found occupying the posterior pole. Macular optical coherence tomography (OCT) confirmed the subretinal elevation and its papillary origin. Our approach was to implement close surveillance. **Conclusion:** Optic disk pit is a rare congenital anomaly. The main complication is serous maculopathy, the pathophysiology of which remains incompletely understood and its treatment is not consensual. Optic disk pit should always be considered in cases of serous retinal detachment by fundus examination supplemented by OCT.

Keywords: Optic Disk Pit, Serous Macular Detachment, OCT.

Copyright © 2025 The Author(s): This is an open-access article distributed under the terms of the Creative Commons Attribution 4.0 International License (CC BY-NC 4.0) which permits unrestricted use, distribution, and reproduction in any medium for non-commercial use provided the original author and source are credited.

INTRODUCTION

Optic disk pit is a rare congenital anomaly, susceptible to macular complications such as serous retinal detachment of the posterior pole. Its pathogenesis remains controversial, as does its treatment.

CASE REPORT

A 32-year-old patient presented with a progressive decline in visual acuity in her right eye over

the past three years. Visual acuity was 3/10 in the right eye and counting fingers at 1 m in the left eye. Funduscopy revealed a temporal Optic disk pit on the right eye associated with a macular serous retinal detachment. On the left, a chorioretinal coloboma was found occupying the posterior pole. Macular optical coherence tomography (OCT) confirmed the subretinal elevation and its papillary origin (Figure1). Our approach was to implement close surveillance.

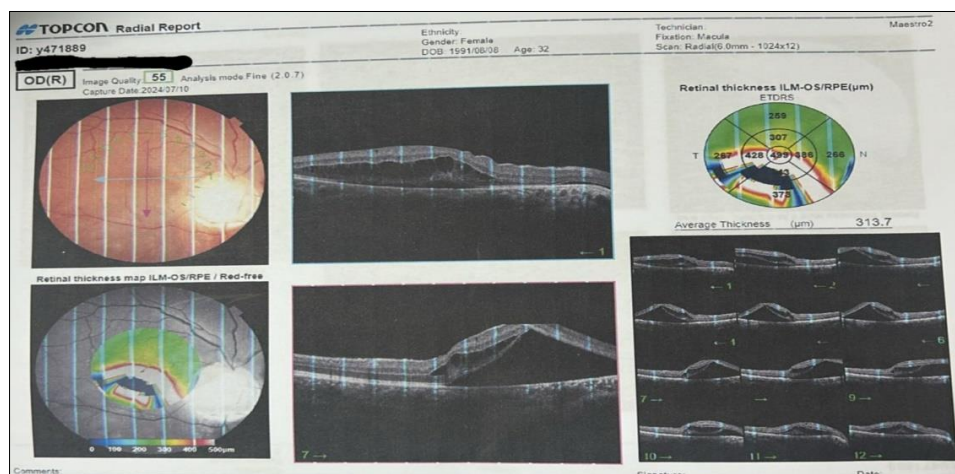


Figure 1: OCT confirms the subretinal elevation and its papillary origin in the right eye

DISCUSSION

Colobomatous optic disk pit are a congenital anomaly related to a dehiscence of the cribriform plate of the optic nerve, which causes a dysplastic retinal herniation, limited by a collagen wall [1]. This anomaly affects 1 in 11,000 eyes (according to the Blue Mountain Study) and is mostly unilateral (bilateral in 15% of cases). The dimples appear at the back of the eye as a grayish depression, most often temporal to the optic nerve [2]. Unlike a coloboma which generally affects the inferonasal part of the optic nerve.

Colobomatous optic disk pit is usually asymptomatic in the absence of complications and discovered during a systematic examination of the fundus. It can be complicated in 25 to 75% of cases by serous retinal detachment (SRD) and macular retinal schisis responsible for a decrease in visual acuity (VA). Then called maculopathy secondary to colobomatous dimple [3]. Papillary maculopathy is defined by the concentration of intraretinal and subretinal fluid at the level of the macula. Despite advances in fundus imaging, the origin of the fluid remains unknown (vascular, cerebrospinal fluid, dehiscence within the dimple) and the exact pathogenesis of maculopathy is not fully elucidated [4-6]. A recent study by the International Retina Group (IRG) shows that this maculopathy most often affects patients aged 30 to 40, although it is possible at any age [7], particularly in the elderly, and can be confused with other diagnoses such as exudative age-related macular degeneration (AMD) [3]. It is therefore particularly important to look for the presence of a colobomatous dimple in the assessment of RSD or intraretinal schisis [2]. The diagnosis is confirmed by optical coherence tomography (OCT) [8].

CONCLUSION

Optic disk pit is a rare congenital anomaly. The main complication is serous maculopathy, the pathophysiology of which remains incompletely understood and its treatment is not consensual. Optic disk pit should always be considered in cases of serous retinal detachment by fundus examination supplemented by OCT.

REFERENCES

1. Korobelnik J-F, Tadayoni R. *Décollement de la rétine, chirurgie maculaire*. Paris : Lavoisier, 2012.
2. Guillaume Le Guern, Aude Couturier. *Maculopathies associées aux fossettes colobomateuses : suivi et prise en charge* Les Cahiers d'Ophtalmologie Mai 2021; n°245:18-20.
3. Gabrielle P.-H. *Fossette colobomateuse : un diagnostic à ne pas manquer en cas de décollement séreux rétinien maculaire*. 29 janvier 2021
4. Dimitrios Kalogeropoulos. *Optic Disc Pit Maculopathy: A Review*, DOI: 10.22608/APO.2018473
5. Irini Chatziralli, *Optic disk pit maculopathy: current management strategies*, PMID: 30127591 PMCID: PMC6089607 DOI: 10.2147/OPHTH.S153711.
6. Steel DHW, Suleman J, Murphy DC, et al. *Optic disc pit maculopathy: A two-year nationwide prospective population-based study*. *Ophthalmology*. 2018; 125:1757–1764.
7. Friedman, *Prevalence of age-related macular degeneration in the United States*, *Arch Ophthalmol*, № 122, c. 564 DOI: 10.1001/archophth.122.4.564
8. Iglicki M, Busch C, Loewenstein A et al. *Underdiagnosed optic disk pit maculopathy: spectral domain optical coherence tomography features for accurate diagnosis*. *Retina*. 2019; 39(11):2161-6.