CASE REPORT

Recurrent Photoreceptor Loss with Spontaneous Recovery as a Presenting Sign of Syphilitic Outer Retinopathy

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Abstract: A 64-year-old male presented with sudden onset of decreased vision with a central blind spot and shimmering visual symptoms without significant inflammatory findings, and was noted on optical coherence tomography to have localized loss of photoreceptors, which spontaneously resolved after two weeks. This recurred subsequently three times spaced by months of recovery with normal vision and normal photoreceptors. In the fourth episode, the patient developed an arcuate-shaped area of outer retinitis more typical for acute syphilitic posterior placoid chorioretinitis, and the diagnosis was made serologically. With subsequent treatment for syphilis, there was not any recurrence after the treatment and vision recovered to 20/20 with a normal photoreceptor layer.

Keywords: Syphilis, Outer retinopathy, Acute syphilitic, Posterior placoid chorioretinitis, Achoroidal thickening, AZOOR.

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1. INTRODUCTION

Sudden vision loss associated with localized damage to the photoreceptor layer on Optical Coherence Tomography (OCT) without significant inflammatory findings is an unusual clinical presentation, especially when there is spontaneous resolution with the recovery of normal vision and a normal photoreceptor layer on OCT. Acute Zonal Occult Outer Retinopathy (AZOOR) is an idiopathic disease with acute damage to photoreceptors with minimal funduscopic findings. It is associated with photopsias and is usually found in young females. Acute cone photoreceptor vision loss was recently described as a presenting sign of AZOOR, but the findings persisted and were associated with decreased signal in the photoreceptor layer on OCT [1]. Differential diagnosis includes other acute inflammatory diseases, such as acute multifocal placoid pigment epitheliopathy or unilateral acute idiopathic maculopathy, and infectious diseases, such as tuberculosis and syphilis, sarcoidosis, and intraocular lymphoma.

2. CASE REPORT

A 64-year-old Caucasian male presented with shimmering visual phenomena and a central blind spot in the left eye. Visual acuity was 20/25 in the right eye and 20/50 in the left eye. Slit-lamp examination was normal without iritis, and a dilated fundus exam revealed a focal intraretinal hemorrhage in the macula in the left eye without vitritis. The OCT of the left eye showed diffuse loss of the photoreceptor layer and increased chorioidal thickness (Fig. 1A). At two weeks, vision in the left eye spontaneously improved to 20/25 with spontaneous recovery of the photoreceptor layer and vision recovered to 20/20 with a normal photoreceptor layer.

Eleven weeks later, he complained of decreased vision in the left eye with vision decreased to 1/200. The funduscopic exam showed retinal whitening and fundus autofluorescence, fluorescein angiography, and OCT angiography findings, which are noted in Fig. (3). OCT displayed subretinal fluid, disruption of the subfoveal cone photoreceptor layer, and chorioidal thickening (Fig. 1C). At one month follow up, there was spontaneous recovery of vision to 20/30 with the recovery...
of the photoreceptor layer and resolution of subretinal fluid and outer retinal changes without intervention (Fig. 1D).

Five months later, he complained of central blurring in his left eye with visual acuity of 20/25. Slit-lamp examination and posterior examination were unremarkable with OCT in the left eye showing attenuation of photoreceptor mostly nasally and choroidal thickening (Fig. 1E-1G). The patient’s vision spontaneously recovered again with the recovery of the photoreceptor layer.

Fourteen months after the initial presentation, he reported blurry vision in the left eye with visual acuity of 20/40. Dilated fundus examination revealed an arcuate pattern of retinal whitening (Fig. 4). OCT showed disruption of the outer retina with marked loss of the photoreceptor layer. Due to a suspicion of syphilitic posterior placoid chorioretinitis with this more significant presentation, serology was performed for syphilis, which showed a positive FTA-ABS and VDRL (1:64). HIV testing by the infectious disease consultant was negative. He was treated with two weeks of intravenous penicillin G followed by 2.4 million units of intramuscular Bicillin. Two years after the last episode, his vision is 20/20 in both eyes with no recurrences in the left eye and recovery of a normal photoreceptor layer (Fig. 1H). His VDRL decreased from 1:64 to 1:4 after treatment.

3. DISCUSSION

This case demonstrates the unusual finding of symptomatic vision loss without clinical signs of active inflammation and with loss of the ellipsoid layer on OCT, which rapidly and spontaneously resolved without treatment on three separate episodes with the recovery of good vision and a normal ellipsoid layer. Additional findings included choroidal thickening and decreased flow on OCT angiography isolated to the left eye. Specifically, in the first episode, there is prominent choroidal thickening and an intact External Limiting Membrane (ELM) and disruption of the ellipsoid zone on OCT. While the posterior segment examination did not fit the clinical picture of Typical Acute Syphilitic Posterior Placoid Chorioretinitis (ASPPC), which usually presents as a large yellowish placoid outer retinal lesion, it does suggest an outer retinopathy associated with choroidal inflammation targeting the outer retina. The isolated clinical picture of loss and recovery of the photoreceptor layer on multiple occasions is unique in this case and not previously reported with ocular syphilis. OCT angiography in the second episode showed flow voids in the choriocapillaris, suggesting associated choroidal inflammation. The last episode presented a yellowish-white outer retinal lesion, which was very suggestive of ASPPC, leading to a definitive diagnosis. After the final episode, there was positive serology for syphilis and ASPPC was diagnosed and treated through infectious disease consultation with intravenous penicillin and intramuscular penicillin with no recurrent episodes following appropriate antibiotic treatment over two years.

The imaging, in this case, highlights the acute damage resulting in outer photoreceptor dysfunction, photopsias, and decreased vision. The associated choroidal thickening is suggestive of acute inflammation that may play a role in the pathogenesis of this disease. Evidence in this case for choroidal inflammation includes increased choroidal thickness and punctate hyperreflective dots on OCT and flow voids on OCT angiography at the level of the choriocapillaris. There is one case report by Tsui et al. that supports this notion that flow voids may be seen in ASPPC on OCT angiography [2]. In another case report by Brito et al., they observed choroidal thickening with loss of the ELM and photoreceptors on OCT in unilateral ASPPC [3]. Pichi et al. specifically looked at the OCT findings in ASPPC and found that there was choroidal hyperreflectivity in 14/30 eyes (46.6%) again supporting the involvement of the choroid in the pathogenesis [4].

Spontaneous recovery of ASPPC without initial treatment for syphilis has been previously reported [5, 6]. However, acute loss and then spontaneous recovery of the photoreceptor layer spaced by months of asymptomatic periods with normal vision and photoreceptor recovery are unique presentations of ocular syphilis demonstrated by this case, which mimicked findings seen in acute zonal occult outer retinopathy [1]. The uniqueness of the ocular immune privilege environment may play a role in the ability of the pathogen to elicit an immune response and then resolve spontaneously [4]. Additionally, the pathogen itself, Treponema pallidum, is characterized by phases of latency and its systemic clinical manifestations follow a path defined by current immune status, site of infection, and timing/duration of disease [7]. Similar is the case with the eye that this pathogen is capable of exacerbations and periods of inactivity.
Fig (2). Color fundus photograph (A) and OCT (B) of the uninvolved right eye. Note the lack of inflammation or edema in the color photograph. Note the normal and intact photoreceptor layer on the OCT.

Fig (3). Multimodality imaging from the second flare demonstrates retinal whitening and intraretinal hemorrhage on fundus photo (A) with hyperautofluorescence of the nasal macula on fundus autofluorescence (B) with patchy chroidal filling and delayed perfusion along the inferotemporal vein on fluorescein angiography (C) and flow voids in the left eye on optical coherence tomography angiography (D).

Fig (4). Fundus photo (A) of the fourth flare displays retinal whitening extending from the optic nerve and into the macula with corresponding fundus autofluorescence (B) showing hyperautofluorescence and stippled autofluorescence in the superotemporal macula of the left eye.

CONCLUSION

This case highlights the unusual presentation of outer retinal damage with loss of the photoreceptor layer on OCT with complete spontaneous recovery for extended periods between episodes. Prior to the diagnosis of syphilis, the presenting findings were considered an unusual variant of acute zonal occult outer retinopathy with central involvement. Understanding the potential presenting clinical spectrum of recurrent and spontaneous resolving outer retinopathy as a possible presenting pattern of syphilis is important, as the treatment of syphilis has now prevented any recurrence with follow-up for over two years while preventing other systemic complications of syphilis.

ETHICS APPROVAL AND CONSENT TO PARTICIPATE

Not applicable.

HUMAN AND ANIMAL RIGHTS

Not applicable.

CONSENT FOR PUBLICATION

Informed consent was taken from all the patients when they were enrolled.

STANDARDS FOR REPORTING

CARE guidelines and methodology were followed to conduct the study.

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CONFLICT OF INTEREST

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REFERENCES


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