Post typhoid fever neuroretinitis with serous retinal detachment and choroidal involvement - A case report

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ABSTRACT

Purpose: To report post typhoid fever neuroretinitis with Serous Retinal Detachment and choroidal involvement.
Observation: Patients with diminished vision post typhoid fever can present with neuroretinitis with serous retinal detachment.
Conclusion and importance: With help from noninvasive imaging such as optical coherence tomography angiography (OCTA) and Deep Range Imaging (DRI), we were able to conclude choroidal involvement - which has not been discussed in literatures yet. OCTA and choroidal thickness both served as good indicators for monitoring the response of treatment in this case.

1. Introduction

Salmonella typhi is a common gastrointestinal pathogen. It causes typhoid—a waterborne infectious disease. However, on rare instances, the organism has been reported to cause ocular pathologies, ranging from uveal complications such as iritis, retinal hemorrhage, choroiditis, endophthalmitis and panophthalmitis to retinal complications such as retinitis and vasculitis. The hypothesized mechanism is either via a direct invasion by the organism or via an immune-complex mediated hypersensitivity reaction.

Here, we report a patient who presented with neuroretinitis with serous retinal detachment following a typhoid fever four weeks prior to presentation.

2. Case report

A 21-year-old female presented at our retina clinic, complaining of sudden onset painless diminution of vision in her left eye since 27 days. After seven days of the onset, she started experiencing the same symptoms on her right eye. She initially visited a local eye clinic, from where she was referred to our comprehensive care center. On questioning, she has a recent history of typhoid fever, onset approximately 45 days ago and resolved after taking oral azithromycin 500mg and cefixime 400mg BID for 10 days. Diagnosis was established by both blood culture a positive Widal test showing S. typhi O antigen titer at 1:160 and H antigen at 1:80. Her AH and BH antigens were negative.

Her visual acuities were 5/60 OD and 4/60 OS. The slit-lamp biomicroscopic evaluation of the anterior chamber was normal. Dilated fundus examination of RE showed clear vitreous with hyperemic disc with blurred margin. Multiple whitish fluffy lesion extending over the superotemporal arcade and over macular area with formation of ‘macular star’ (Fig. 1a, Fig. 2a). Optical coherence tomography (OCT) and OCT-angiography (OCTA) were done using Topcon Medical Systems Triton Deep Range Imaging (DRI) PLUS SS-OCT which revealed thickened and detached neurosensory retina over the macular region of both eyes. Additionally, OCTA of both eyes for choroidal vasculature revealed

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abnormal “patchy” flow voids in the choriocapillaris-likely suggestive of a sluggish blood flow or ischemia. Deep range imaging (DRI) of the choroid revealed increased choroidal thickness and dilated choroidal vasculature, indicating a concurrent choroidal inflammation (Fig. 1a, b, Fig. 2a, b).

Baseline blood, biochemical, and serological investigations were done to rule out pathologies mimicking a similar clinical picture. Then, a working diagnosis of Post typhoid fever neuroretinitis with serous

![Figures 1-2](image_url)
retinal detachment was made. The patient was started on oral prednisolone 1mg/kg/day for 10 days, oral ciprofloxacin 500mg twice daily for a week, topical flurbiprofen 0.03% four times a day, and topical tropicamide 1% twice a day. Adjunct antibiotics were added as the patient had defaulted the previously prescribed medication on day 7. She was advised to follow-up in 10 days.

At follow-up, the patient’s visual acuity had improved to 6/24 in both eyes with no further improvement on refraction. The OCTA evaluation revealed that the central serous detachment had decreased with minimal residual subretinal fluid (Figs. 1b and 2b). OCTA imaging of the

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Fig. 2. Response of left eye to treatment. 2a. Day 1 (Top) Left eye had less numbers of cotton wool spot and an incomplete macular star. (Middle) Central serous retinal detachment was with increased choroidal thickness. (Lower and bottom) OCTA and vascular density map of choriocapillaris as heat map showing coarse pattern. 2b. Day 10 (Top) Fundus photos showing decreased areas of cotton wool spots. (Middle) Decreased sub-retinal fluid level but choroidal thickness increased. (Lower and Bottom) Choriocapillaris signals and vascular density map returning to more fine signals but still with presence of flow voids. 2c. Day 24 (Top) Normal fundus with sharp foveolar reflex. (Middle) Complete resolution of subretinal fluid with decreased macular thickness with normalization of choroidal thickness. (Lower and Bottom) OCTA and vascular density map of the choroid showing near normal choriocapillaris pattern.
retina was normal. Although the choroid’s OCTA also revealed reduction in flow voids, DRI still revealed a persistent and increased choroidal thickness in both eyes (Figs. 1b and 2b). Considering the possibility of an ongoing choroidal inflammation despite improvement in retinal imaging, the patient was advised to continue the steroid at the same dose for 10 more days while discontinuing the antibiotic. Topical medications were unchanged.

The next follow up was on day 24 after the initial presentation. The patient had vision of 6/12 and best corrected vision of 6/9 in BE. On OCT, the SRF had completely resolved with thinning of macular area – likely suggestive of atrophy. The choroidal thickness was normalizing with architecture of the choriocapillaris returning to near normal (Figs. 1c and 2c). Her vitals were stable. We tapered the steroid at the rate of 10mg/week while continuing the topical medication. She was advised to follow-up after 4 weeks for visual field but was lost to follow up.

3. Discussions

In this case report, we described a patient with a recent history of typhoid fever who developed bilateral neuroretinitis with star-shaped maculopathy associated with bilateral serous detachment.

Prabhushanker et al reported a case of bilateral retinitis following typhoid fever. The fundus examination of the right eye showed white fluffy lesions and superficial hemorrhages around macula with macular star suggestive of retinitis. The OCT showed macular serous retinal detachment. The patient was treated with steroids and followed every 2 weeks for 3 months. The final OCT of the right eye revealed complete resolution of serous detachment. However, due to minimal disc edema, neuroretinitis was ruled out.

There are very few case reports on this rare condition. Retinal infiltration is the assumptive pathophysiology behind the condition. Neuroretinitis of other origins like cat scratch and Lyme’s diseases has been reported more frequently. A pathology that incites immune-mediated response can disturb the blood-retinal barrier. It has been hypothesized that a micro emboli can obstruct microcirculation, eventually leading to sluggish blood flow and disc edema. A similar pathological cascade is thought to result in retinal vasculature leakage, causing exudative retinopathy.

It has been also been postulated that immunological complexes are behind the pathogenesis of immune-mediated vasculitis secondary to typhoid infection. The complexes, in return, inflicts further damage by affecting self-antigens via homology or molecular mimicry. Case reports by Laul R et al. and Relhan N et al. found that typhoid manifests as vasculitis and neuroretinitis with OCT revealing macular sensory detachment. Cases were successfully treated with oral steroids. However, the role of steroid has always been controversial in the treatment of infective neuroretinitis due to the lack of literature on this topic. Mild cases may show spontaneous resolution but severe cases require treatment.

In our case, oral ciprofloxacin and oral steroid were started on the day of presentation and baseline investigation was done to rule out other causes. On the fourth week of follow up, complete resolution of disc edema and SRF with faint impression of macular star and improved visual acuity were noticeable. In addition to OCT, we used OCTA and choroidal imaging (DRI) to evaluate the status of both retina and choroid and to monitor the response to treatment. So far, many literatures have reported only about the insult to the optic disc and retina. Using DRI and OCTA of the choroid, we were able to report the involvement of the choroid. We also found that the inflammation of the choroid continued to persist even after the clinical signs suggested a resolution — such as decrease in the SRF and improvement of the visual acuity. OCTA which initially revealed a disrupted architecture of the choriocapillaris vasculature; gradually regained its much more “granular” pattern after initiating the treatment. This also further coincided with improvement in visual acuity.

Various reports have indicated that choroidal thickness and assessment of its vascularity can help in decision making and monitoring disease progression. In our previous report of a sympathetic ophthalmia which had a similar clinical features in the posterior segment, we had expressed how choroidal thickness and its vasculature pattern/architecture analysis using OCTA could be helpful in monitoring the disease progression. The same imaging technique was found to be important in the management of this case as well. This, we believe, further adds to the evidence that OCTA of the retina and the choroid and choroidal involvement have the potential to be used as vital indicators in monitoring the pathologies of the uvea and retina.

4. Conclusion

A rare case of noninfectious, immune-mediated, neuroretinitis and vasculitis with macular neurosensory detachment that occurred after the resolution of typhoid fever can be treated effectively with oral steroid. Noninvasive imaging modalities such as OCT, OCTA and DRI indicate choroidal involvement. These imaging techniques also play important roles in monitoring both the progression of the disease and the response to treatment.

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Conflict of interest

All of the authors do not have any conflicts of interest to declare.

Ethical consideration/patient consent

Written consent to publish this case has not been obtained. This report does not contain any personal identifying information.

Authorship

All authors attest that they meet the current ICMJE criteria for Authorship.

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