

## Clinical Image

# Stellar Neuroretinitis as Presenting Sign of Behcet's Disease

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## Observation

A 40-year-old One-eyed patient (right eye) acuity history of post-traumatic burst in his left eye at the age of 6 years. Consult for a decrease of vision on his right eye that has progressed for 3 weeks. Ophthalmic examination found visual acuity: 2/10 P10 in the right eye, the left eye presents enophthalmos with an opacified cornea; the anterior segment was normal in the right eye, fundus examination shows hyalitis graded 1+, papillary edema, peripapillary hemorrhages and stellar macular exudates, cottony nodules, copper arteries with vascular sheathing of the superior temporal branch and areas of retinal ischemia (Figure 1). The inflammatory and serological assessment is negative. Fluorescein retinal angiography shows papillary diffusion of fluorescein and vasculitis (Figure 2). We find serous retinal detachment on OCT (Figure 3). The general examination found bipolar aphthosis. The diagnosis of Behçet's disease is retained. Treatment with immunosuppressants allowed the regression of papillary edema with persistence of some macular exudates and great improving visual acuity.

## Discussion

Neuroretinitis can be due to several diseases; infectious diseases being the most frequent in young subjects [1,2]. More rarely, the inflammatory origin especially Behçet's disease can produce neuroretinitis which is more frequently accompanied by a vitreous reaction [2,3]. Behçet's disease is an idiopathic systemic inflammatory disease characterized by intraocular inflammation, oral and genital ulcers, skin lesions and many other visceral damage [4]. Ocular involvement is characterized by anterior hypopyonic uveitis, posterior involvement being rarer, especially if it is isolated, which is the case in our patient [5,6]. OCT is used to check for posterior segment edema and/or Serious Retinal Detachment. Retinal angiography has a threefold benefit: confirming papillary edema, directing towards its ischemic or inflammatory aetiology and looking for vasculitis. The treatment of Behçet's disease is controversial between corticosteroid therapy and immunosuppressants. The prognosis depends on the speed and precocity of the treatment, making collaboration between ophthalmologists and internists particularly important [7].

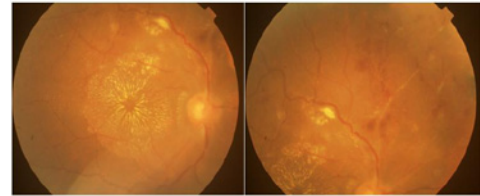


Figure 1: Fundus (Right eye) shows: Left: Neuroretinitis. Right: Occlusive vasculitis.

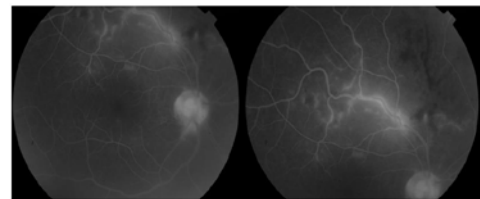


Figure 2: Retinal angiography with fluorescein in late stages: On the left: papillary diffusion. On the right: vasculitis with vascular occlusion.

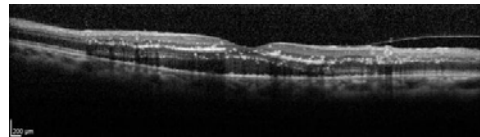


Figure 3: Optical Coherence Tomography (OCT): macular exudates especially at the level of the outer plexiform layer, serous retinal detachment.

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