Case report of bilateral Intermediate uveitis

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Introduction

Multiple sclerosis (MS) is a chronic inflammatory disorder of the central nervous system (CNS) white matter and a common cause of neurological disability in young adults . Optic neuritis is the most frequent ophthalmic manifestation of MS, however intraocular inflammation may also occur.

Methods

A case report of a 37 year-old man presenting with recent onset of photophobia, in addition to gradually progressing vision deterioration in both eyes. Clinical findings on presentation summarized in table (1) with OCT findings in figure (1). On questioning, a medical history of Tonsillectomy & adenoidectomy, Total splenectomy & Multiple sclerosis as reported.

	Right eye	Left eye
Visual acuity	6/60	6/60
Anterior chamber	Mild reaction (cells +1)	Mild reaction (cells +1)
Pupils	RRR, no RAPD	RRR, no RAPD
Intraocular pressure	10mmHg	10mmHg
Anterior vitreous	Severe vitritis	Severe vitritis
Posterior vitreous	Snow balls	Snow balls
Central fundus	Cystoid macular oedema	Cystoid macular oedema
Peripheral retina	Vasculitis, fibrosis, snowbanking	Vasculitis, fibrosis, snowbanking
Table (1): Clinical findings		



Figure (1): OCT scans of right and left eye on presentation



Results

A diagnosis of Intermediate uveitis caused by Multiple sclorosis was made, with Oral steroid started at 60 mg daily with a gradually tapered dose. Visual outcome 2 weeks after starting treatment: 6/12 right eye, and 6/24 left eye, and 4 weeks after starting treatment: 6/12 right eye, and 6/18 left eye., with displayed OCT results, figure (2).

Discussion

Optic neuritis is the most frequent ophthalmic manifestation of MS, however intraocular inflammation may also occur (1). The most common types of uveitis were intermediate (pars planitis) and panuveitis. These findings are in line with Markomichelakis's and Biousse 's findings (2,3)

About 3-27% of patients with multiple sclerosis (MS) develop IU/pars planitis, (4,5) and 7.8-14.8% of patients with IU/pars planitis develop MS (6,7)

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