**INTRODUCTION**

Primary central nervous system lymphoma (PCNSL) accounts for 4 to 7% of brain tumors. The onset of this lymphoma is generally in the 5th and 6th decade of life. Primarily involves the brain, eyes, and leptomeninges without other systemic findings. It occurs mostly in the immunosuppressed population but its incidence has increased in normal patients as well such as this patient (Chan & Wallace, 2004; Hormigo & De Angelis, 2003; Smith *et al.*, 2000). The most common ophthalmic manifestations are posterior uveitis or vitritis, combined anterior and posterior uveitis, or subretinal pigment epithelial infiltrates (Hochberg & Miller, 1988). The anterior or posterior uveitis in 75% of patients and sub retinal pigment epithelium (RPE) infiltrates in 20% of patients. Sub RPE infiltrates from accumulation of lymphocytes may be intermittently present leaving geographic atrophy on resolution. The appearance of ischemic retinopathy or optic neuropathy is from accumulation of lymphocytes around the retinal vessels.

If the patient with uveitis has a negative or partial response to steroids, then one should consider samples from the aqueous and vitreous should be done for cytology and complete diagnostic testing. Other tests such as Immunophenotyping to determine clonality sharing same surface light chain, gene rearrangement to identify clonality within the cells, neuroimaging and cerebrospinal fluid study should be done.

Chemotherapy such as rituximab, a chimeric antibody, induces apoptosis in CD20 positive cells-B cells. Intravitreal methotrexate and/or rituximab with or without vitrectomy are indicated treatment of the vitritis. The five-year survival rate ranges from less than 20% to 34% with the best therapy (Panageas *et al.*, 2005; DeAngelis *et al.*, 2002). The differential diagnosis between chronic inflammation and intraocular lymphoma become very difficult (Wilson *et al.*, 1992). The suspicion of a masquerade syndrome, particularly a primary CNS lymphoma, should always be considered in the elderly like this case (Zamiri *et al.*, 1997).

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CASE REPORT

A 65-year-old Vietnamese male had one month of progressive blurred vision in both eyes and no other eye complaint. Other medical history revealed hypertension and high cholesterol. Through a careful history taking with the suspicion of the etiology revealed that one year previously, he had confusion and difficulty walking. The Imaging test was conducted and demonstrated a mass lesion of (5.2x5.0x3.5cm lesion) in the right frontal lobe with edema, involvement of corpus callosum, and 15 mm of L to R midline shift was noticed. After steroid treatment of the brain edema, he had stealth image guided left frontal brain biopsy showing diffuse large B-cell lymphoma (DLBCL), LDH 312, and confirmed the diagnosis of primary diffuse large B cell lymphoma. He was treated with chemotherapy of Methotrexate 3.5 g/m² and cytarabine 2 mg/m² and whole brain radiation therapy. There was a complete remission after the therapy.

Vision was OD 20/60, OS 20/200. Refraction: -0.25+1.00x118 OD, +0.50 +0.75x34 OS. Intraocular pressure was 8.8/8.7 measured by auto, non-contact tonometry.

Slit lamp exam each eye demonstrated a quiet anterior segment with no conjunctival injection and no anterior chamber cells or flare. There was
1+ nucleus sclerotic cataract both eyes. Fundus of both eyes was partly obscured by cells mainly in the posterior vitreous. There appeared to be a few drusen in the right eye. Fundus photography and Fluorescein angiography showed no other abnormalities. (Figure 1) Optical Coherence Tomography (OCT) of each macula was normal. (Figure 2). It was felt that the patient had B cell lymphoma intraocular recurrence. We did not treat the patient with local chemotherapy using intravitreal Methotrexate (IMTX) (Helbig et al., 2003; Frenkel et al., 2008). Instead, as the hazy vision affected the patient’s functional life style, sequential bilateral vitrectomies were done on 5/1/17 and 7/17/17 without complication. The vitreous sample from the first eye having vitrectomy was sent for study which demonstrated large B cell lymphoma. Post-op vision improved to 20/25 without the complaint of hazy vision. (Figure 3)

DISCUSSION

This case was diagnosed from a good history taking due to the suspicion of the masquerade syndrome. However, because of the malignant nature of this tumor and to differentiate from other causes vitritis, vitrectomy was indicated for the confirmation of the intraocular recurrence of lymphoma.

The vitrectomy dramatically improved the vision in one eye. Subsequently the second eye was also operated as requested by patient. There was no recurrent retina lesion detected after the vitreous were cleared in both eye. Frequent future oncological and ophthalmological follow up are essential to save this patient’s eye sight and life.

In this patient recent follow up examination on 12/1/2017 showed clear vitreous both eyes with 2+ nucleus sclerosis cataracts. Vision best corrected was 20/50 OD, 20/30 OS. He will be followed every three months.

CONCLUSION

Good history taking assisted in the diagnosis of intraocular lymphoma in this case with bilateral vitritis. However, bilateral vitrectomy restored the vision and further confirmed the diagnosis of lymphoma for future follow up care.


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