Case Report

Immunoglobulin G4-related disease of the cavernous sinus with orbit invasion – A case report

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INTRODUCTION

Immunoglobulin G4-related disease (IgG4-RD) is a rare systemic inflammatory disease that is characterized by tissue infiltration caused by IgG4 plasma cells and sclerosing inflammation of various body organs. At present, there are very few reported cases of IgG4-RD invasion of cavernous sinus and the orbit.

ABSTRACT

Background: Immunoglobulin G4-related disease (IgG4-RD) is a rare systemic disease of unknown etiology. It is characterized by tissue infiltration caused by IgG4 plasma cells and sclerosing inflammation of various body organs. At present, there are very few reported cases of IgG4-RD invasion of cavernous sinus and the orbit.

Case Description: A 56-year-old female with a history of rheumatoid arthritis was presented with a gradual onset of right oculomotor, abducens, and trigeminal nerve deficits. Four weeks after the onset of symptoms, the patient developed gradual visual deficit. Following this, a trial of steroids was administered to the patient. However, the treatment did not work as expected and patient's condition worsened. She progressed on to suffer complete visual loss in the right eye. Extensive work-up conducted on her turned out to be nondiagnostic. After this, the patient was referred to us for our evaluation. Neuroimaging revealed a right-sided cavernous sinus and orbital apex lesion. Given the lack of diagnosis and response to steroid treatment, we recommended surgical intervention and performed a modified pterional and pretemporal approach with extradural anterior clinoidectomy and transcavernous approach. We performed a lesion biopsy and cavernous sinus decompression, which helped in the partial recovery of visual function. The pathology report was consistent with IgG4-RD.

Conclusion: IgG4-RD is a rare disease that occurs even less in combination with cavernous sinus and orbit invasion. The rarity of the disease and the diverse presentation of symptoms have sometimes caused delayed diagnosis and intervention. Patients who failed to respond to conservative management and patients in the fibrotic stage of the disease without other organ involvement may benefit from surgical intervention if amenable. Early suspicion, diagnosis, and intervention can facilitate better prognosis.

Keywords: IgG4-related diseases, IgG4-related disease with intracranial invasion, Immunoglobulin G4-related disease of cavernous sinus and/or orbit invasion

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management of the disease.\textsuperscript{[2-4,12,13,19]} In this report, we will describe the unique case of a patient with IgG4-RD of cavernous sinus and orbit invasion, which was treated through surgical intervention after conservative management and steroid therapy failed to work.

**CASE PRESENTATION**

**Clinical presentation**

A 56-year-old female with a history of rheumatoid arthritis was managed chronically with immunosuppression with etanercept. She developed a preoperative progression of visual decline and ophthalmoplegia for 7 weeks. Initially, she was admitted to the emergency department with right-sided eye pain and diplopia of 2 weeks duration and was hospitalized and managed by neurology, rheumatology, oncology, and ophthalmology. Her vision was initially intact with visual acuity of 20/25 on the right eye and 20/20 on the left eye with complete visual fields. She presented with the right pupil sparing ophthalmoplegia, also with trochlear, and abducens palsies. Specifically, she had right-sided ptosis, limited adduction/abduction, limited down/up gaze, having also ophthalmic (V1) and maxillary (V2) trigeminal nerve sensory branches dysfunction with hypesthesia. Extensive work-up was conducted, including lumbar puncture with cerebrospinal fluid analysis and cytology studies, which returned normal. The meningitis panel and cultures were all negative. Additional tests such as human immunodeficiency virus, angiotensin-converting enzyme, venereal disease research laboratory, antineutrophil cytoplasmic antibody, antinuclear antibodies, serum protein electrophoresis, immunofixation electrophoresis, and lymphoma profile were all negative. Computed tomography of chest, abdomen, and pelvis did not show any signs of malignancy. Neuroimaging showed enhancing soft tissue within the right cavernous sinus extending into the right orbital apex and posterior aspect of the right lateral rectus muscle, with enlargement of the superior ophthalmic vein [Figure 1].

Differential diagnosis suggested retro-orbital granulomatous pseudotumor such as Tolosa-Hunt syndrome, granulomatous infection such as sarcoidosis, infiltrating neoplasm such as lymphoma, metastasis, and other neoplastic etiologies. Cavernous sinus thrombosis and carotid cavernous fistula were considered unlikely, given the lack of conjunctival injection and papilledema.

The patient was initially administered with intravenous steroids for 7 days that improved her ptosis and V1-V2 hypesthesia, with stable visual function. A repeat brain and orbit MRI 1 week after treatment did not show changes on the right orbital apex/cavernous sinus lesion, but given mild improvement, she was then discharged home with etanercept and methotrexate weekly.

She had outpatient follow-up with ophthalmology 3 weeks after discharge from the hospital showing similar visual examination. One week later, the patient experiences complete right visual loss, then she was referred to neurosurgery clinic and she was evaluated for possible urgent surgical intervention.

The patient underwent to a right-sided skull base pretemporal approach with orbitotomy at the lateral wall and roof of the orbital apex, and extradural anterior clinoidectomy. Mini-rongeurs were used to decompress the optic canal, and a 1 mm diamond drill was used at the core of anterior clinoid process and optic strut. Mini-rongeurs and skull base dissectors were used to remove the cortical bone during the anterior clinoid process. The lateral wall of the cavernous sinus was then explored to allow the dissection of the cranial nerves III, IV, V1, and V2. Samples were taken from the posterior orbital lesion, the orbital apex lesion, and the inflamed tissue around clinoid carotid. The lateral wall of cavernous sinus was opened at the supratrochlear triangle and tissue sample was extracted from there as well. Durotomy was performed for intradural exploration to identify the entire oculomotor nerve through the intradural cistern segment and cavernous sinus and the infiltrated dura.

![Figure 1](image_url)
in the middle fossa was resected. Fibrin glue, nonsuturable synthetic dura substitute, and fat graft harvested from the abdomen were used for dura reconstruction. Finally, during postoperative period, the patient was administered a short course of steroids.

The resected tissue was submitted to the department of surgical pathology for examination. The dura and soft tissues showed dense fibrosis and a brisk chronic inflammatory infiltrate with numerous lymphocytes, plasma cells, and occasional histiocytes. During immunohistochemistry, numerous plasma cells were highlighted on IgG staining. Approximately half of the plasma cells also expressed IgG4. The combined histomorphology and immunophenotype were consistent with IgG-related sclerosing disease involving orbital region soft tissues and dura mater. No granulomatous inflammation or evidence of a neoplastic process was seen [Figure 2].

During the 6 months follow-up, extraocular movement deficits from the oculomotor and abducens nerve were resolved. Visual function had also improved; the patient was now able to count fingers from a distance of 3 feet (since she had right complete visual loss preoperatively). Improvement was also noted in the patient's visual field [Figure 3]. Imaging studies showed postsurgical changes related to right-sided skull base approach, orbital apex mass resection, and fat packing. There was no significant abnormal enhancement of or soft tissue at the right cavernous sinus or posterior orbital apex. There were also no signs of disease reoccurrence [Figure 4].

**DISCUSSION**

IgG4-RD is multiple organs disorder that occurs most frequently in the pancreatic and hepatobiliary system. Isolated head, neck, and brain involvement in IgG4-RD is uncommon. The first case of lateral skull base IgG4-RD was reported in 2012. Isolated IgG4-RD invasion of cavernous sinus is even more rare. In this report, we have discussed about treatment given to a patient who was presented with cavernous sinus and orbit invasion IgG4-RD, and required surgical intervention to attain adequate functional improvement.

So far, very few population-based studies of IgG4-related disease have been conducted. Therefore, information about the epidemiology of the disease is only sparsely available. In fact, the current nomenclature of IgG4-RD was accepted as recent as in 2011, while the first international consensus on the pathological findings that currently define IgG4-RD was reached in 2012. The current studies have reported that IgG4-RD is more likely to occur in patients with certain striking demographic features. For instance, IgG4-RD is reported to occur predominantly in men between the sixth and seventh decades of life. It is said to be frequently associated with lymphadenopathy and expected to respond well to steroid therapy. However, these characteristics were not there in the case of our patient. Studies indicate that the pancreas is usually the most affected organ. However, it has also been documented that IgG4-RD can affect nearly every organ in the human body, including the skull base. Some IgG4-RD cases were also reported in pediatric populations. IgG4-RD cases are currently more frequently reported in Asian countries. In our case, the patient was a 54-year-old female with IgG4-RD invasion of the right cavernous sinus and orbit.

IgG4-RD is diagnosed based on clinical, radiographic, biochemical, and histopathologic evidence. In most cases, IgG4-RD with intracranial invasion occurs with nonspecific symptoms. Neuroimaging findings also mimic other more common intracranial disorders, which can lead to subsequent delays in its diagnosis and management. There are, however, some clinical presentations that are unique to IgG4-RD, such as subacute presentation of clinical symptoms and lack of constitutional illness. Hence, when patients are presented with these conditions, clinicians are advised to include IgG4-RD in the differentials. Fevers and elevations of C-reactive protein levels have also been reported to be unusual in the IgG4-RD patient cohort. This characteristic was consistent with that of our patient who presented a gradual onset of clinical symptoms without constitutional illness, as shown by the unremarkable extensive work-up results.

**Figure 2:** (a) High-power view showing dense fibrous tissue with lymphocyte and plasma cell-rich inflammatory infiltrate (H&E, ×400); (b) CD138 immunohistochemical stain highlights plasma cells within the inflammatory infiltrate (IHC, ×200); (c) IgG4 immunohistochemical stain stains approximately half of the plasma cells (IHC, ×200).
The most commonly presented symptoms of IgG4-RD are similar to that of any other intracranial space-occupying lesions. The effects of local pressure of the invasion, such as progressive headache, cranial nerve palsies without evidence of systemic disease, nausea, and vomiting, are some of the common symptoms of IgG4-RD. Depending on the location of the disease, isolated lesions often mimicked a diverse spectrum of skull/skull base pathologies including lymphoma, sarcoma, nasopharyngeal carcinoma, neurosarcoïdosis, granulomatosis with polyangiitis, sarcoma, giant cell arteritis, Langerhans cell histiocytosis, and benign skull base tumors.\cite{6,12,32,38}

Radiologically, IgG4-related mass lesions, as in our case, present as homogenously enhanced lesions, which are isointense to hypointense on T1-weighted and T2-weighted images.\cite{6,12,32,38,40}

Three histopathological findings characterize the disease in the affected organ: (1) The presence of a storiform pattern of sclerosis; (2) a dense lymphoplasmacytic infiltrate; and (3) an increased proportion of IgG4-positive cells with respect to IgG-positive cells according to immunohistochemical evidence. Although elevated concentrations of IgG4 in tissue and serum are helpful in diagnosing IgG4-RD, neither one is a specific diagnostic marker. In fact, misdiagnoses of IgG4-RD are increasingly common because of the excessive emphasis that physicians tend to place on moderate elevations of serum IgG4 concentration and their overreliance on the finding of IgG4-positive plasma cells in tissues. Histopathological analysis of biopsy specimens remains the cornerstone in the diagnosis of IgG4-RD. Due to the scarcity, ambiguities, and uncommon nature of intracranial IgG4-RD, its diagnosis can be a challenge and may require several biopsies, which can often lead to delays in diagnosis and management of the disease. Corticosteroids and immunosuppressive agents may exacerbate similar appearing lesions such as osteomyelitis or alter the diagnostic yield of a biopsy and therefore should not be started unless there is a suspected diagnosis of IgG4-RD.\cite{5-25,29,32-37}

There are currently a few accepted criteria for the diagnosis of IgG4-RD, such as the criteria put forth by Okazaki et al.\cite{23} and Umehara et al.\cite{33} Overall, diagnosis of this disease is often based on highly suggestive clinical and radiological findings, one or more organs with diffused or localized edema or a mass characteristic of IgG4, serological diagnosis with IgG4 greater than 135 mg/dl, and pathology diagnosis.\cite{23}

Our patient met the diagnostic criteria of focal lesion in the cavernous sinus and orbit, and had histopathology confirmed through biopsy.

At present, corticosteroids are the first line of treatment administered for IgG4-RD. Proposed regimens include prednisone 40 mg/day with adjustments based on disease response, or prednisolone 0.6 mg/kg/day for 2–4 weeks, followed by a 3–6 months taper to 5 mg/
day and maintenance dose of 2.5–5 mg for up to 3 years. Symptomatic improvement, decrease in IgG4 levels, and decrease in lesion load can be expected within weeks of treatment commencement. Biological therapy combined with radiotherapy has also been employed successfully in the induction and maintenance of clinical response to IgG4-RD, and has been proposed as second-line therapy. There is currently no evidence supporting the use of steroid sparing agents to treat this disease.

Successful surgical intervention for treating IgG4-RD has been retrospectively reported in a few studies. However, prospective data on the surgical treatment of IgG4-RD still remain limited. Studies on the natural history of IgG4-RD show that if left untreated, the disease can progress to extensive organ fibrosis. Patients who are in the fibrotic stage of IgG4-RD often respond poorly to steroid therapy, which makes surgical resection a necessary intervention. Our patient exhibited bone and dense fibrous tissue with fibrosis and severe chronic inflammation of IgG4-RD at the orbital apex and cavernous sinus, which severely compressed the optic canal and orbital apex contents. This patient pupil-sparing ophthalmoplegia is an interesting phenomenon. It has been suggested that slow enlarging masses in the cavernous sinus can preserve this function due to the topographic relation and different degrees of compression between parasympathetic pupillomotor fibers, which are not evenly distributed on the surface of the nerve but rather form a distinct bundle in its dorsomedial aspect. Therefore, any compressive force from a mass within the cavernous sinus spares this function and mainly damages the somatosensory axons. Other explanation is based on ischemic changes also due to compression, different to the microvascular ischemic etiology as seen in diabetes mellitus.12,27 We believe that the patient had reached the fibrotic stage of the disease by the time she was brought to us, which is why she was not responding well to steroid therapy. Thus, we had to go in for a surgical intervention, to decompress the bone structures at the orbital apex and optic canal. Another subset of patients who may benefit from surgical intervention is those who exhibit disease relapse, as well as those who cannot tolerate or face an increased risk of adverse side effects from long-term use of steroids. Patients with solitary IgG4-RD mass without other organ involvement can also benefit from surgical intervention, provided the benefits outweigh the associated risks. It is to be noted that postoperative nonrecurrence of IgG4-RD has been documented as an additional advantage of surgical intervention, in patients who are good candidates for surgical management.9,10,17,27,28,36 Our case also highlights the pitfalls of steroid therapy. Our patient had poor response to initial steroid therapy, which led to the need for surgical intervention. Postoperatively, the patient showed improved vision, full recovery of extraocular movements, and nonrecurrence during the 6-month follow-up.

CONCLUSION

IgG4-RD is an uncommon disease that should be included in the differential diagnosis for cavernous sinus lesions. Early suspicion and initiation of therapy can significantly improve the outcome of the treatment for the disease. A delay in treatment for any reason or lack of response can result in serious and irreversible sequelae. As of today, IgG4-RD is treated with systemic corticosteroids at high doses that are subsequently tapered. In cases of IgG4-RD involving cavernous sinus and orbital apex, we strongly recommend that surgical decompression be performed, to prevent permanent deficit that can be caused by inflammatory tissue compression at the compact space between the orbital apex and the optic canal. We believe that more studies should be done in this field, to provide more insight and algorithm on the surgical management of IgG4-RD, especially with skull base involvement.

Declaration of patient consent

Patient’s consent not required as patients identity is not disclosed or compromised.

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Conflicts of interest

There are no conflicts of interest.

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