Dural ectasia of the optic nerve sheath: is it always benign?

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Abstract: A 36-year-old woman with a 3-month history of progressive visual loss had papilledema, dilatation of the optic nerve sheaths and normal cerebrospinal fluid pressures. She was diagnosed as dural ectasia of the optic nerve sheaths and surgical decompression was performed. In this case report, severe visual loss is described as a serious complication of this rare disease and the importance of early surgical intervention is emphasized.

Keywords: optic nerve, dural ectasia, meningocele

Introduction

Optic nerve dural ectasia is saccular dilatation of the optic nerve sheath. It is characterized by expansion of the cerebrospinal fluid (CSF) around the optic nerve without orbital or cerebral neoplasm and inflammation.¹⁻³ The terms; optic hydrops, primary cyst of the optic nerve sheath, patulous subarachnoid space, cystic hygroma, arachnoid cyst, perioptic subdural hygroma, and dural ectasia of the optic nerve sheath all have been used to describe this entity since its first description in 1918.^{1,2,4} The term optic nerve sheath meningocele was introduced by Garrity et al in 1990.¹ We prefer the term dural ectasia to describe the optic nerve sheath dilatation. Imaging of the orbit using magnetic resonance imaging (MRI) in patients with dural ectasia reveals tube-like enlargement of the optic nerve/sheath complex.^{1,2} We present a patient with rapidly progressive visual loss which was related to dural ectasia of the optic nerve sheath and the vision remained unchanged despite surgical treatment.

Case report

A 36-year-old female was admitted (to the Department of Neurology, Hacettepe University Medical Faculty, Department of Neurology, Neuro-ophthalmology Unit, Ankara, Turkey) with the complaints of headache and severe bilateral visual loss that had progressed rapidly in 3 months. She was found to have papilledema on examination and had been treated with acetazolamide 250 mg qid for 8 weeks elsewhere. Idiopathic intracranial hypertension (IIH) was considered in diagnosis although opening pressure was recorded normal in the lumbar puncture (LP). On our examination her visual acuity (VA) was light perception OD and 20/100 OS. There was a right afferent pupillary defect. Fundoscopy revealed pale optic nerve head with blurred margins on the right and chronic disc edema on the left. She was not able to identify any of the Ishihara color plates OU. LP was performed with an opening pressure of 22 cm of H₂O. CSF analysis showed protein level of 40 mg/dL and no cells. MRI studies revealed a fluid-filled dilated optic nerve

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sheath surrounding the optic nerves (Figure 1). There was no evidence of a space-occupying lesion. Her left VA decreased to counting fingers in a week. The CSF pressure at the repeat LP was significantly low which could not be recorded. Dural ectasia of the optic nerve is considered in diagnosis. Due to progressive dysfunction of the optic nerve despite treatment with carbonic anhydrase inhibitors, optic nerve sheath decompression via medial transconjunctival approach was performed bilaterally by an ophthalmic surgeon (HK). The distal part of the orbital portion of the optic nerve was found to be significantly enlarged and had a bulb-like appearance. The dura was stretched but otherwise looked normal. Overlying dural vessels were also enlarged and tortuous. Three parallel longitudinal incisions, each measuring 2 to 2.5 mm in length, were made into the dura along the enlarged optic nerve. Large amount of clear fluid rapidly egressed with pressure following the first incision. The optic nerves deflated within seconds while cerebrospinal fluid continued to ooze for a while. No intraoperative complications were observed. The left optic disc edema resolved within one week and the patient remained stable over the next 3 months with the same visual acuity.

Discussion

A dilatation of the optic nerve sheath could be secondary to neoplasms and malformations of the orbit, like gliomas, meningiomas and arachnoid cysts involving the optic nerve sheath.^{1,2} Dural ectasia of the optic nerve is a rare cause

of optic nerve sheath enlargement due to the accumulation of CSF around the optic nerve with no associated pathology. The radiological examination of choice is MRI with fat suppression and high spatial resolution which shows tube-like enlargement of the optic nerve sheath isointense with cerebrospinal fluid.² Optic nerve glioma can be associated with thickening and distention of the optic nerve which is in contrast with the appearance of meningocele. Optic nerve sheath meningioma should also be considered in the differential diagnosis because of its “railroad track” appearance of the optic nerve.⁴ Clinical examination, CSF analysis, MRI, investigations including full blood count, anti-nuclear antibodies, erythrocyte sedimentation rate, folate, serum B12 and serum angiotensin converting enzyme levels in our patient revealed no evidence of any other systemic disease or abnormalities around the optic canal.

Papilledema and optic nerve sheath enlargement can also be seen with IIH.⁵⁻⁷ There are some common features of both diseases; however there is no significant evidence suggesting the role of raised intracranial pressure in the etiology of the dural ectasia of the optic nerve.^{1,8} Although optic nerve sheath dilatation is one of the radiological features of IIH, the enlargement of the optic nerve sheath to this degree is unusual. The clue for differential diagnosis is the LP opening pressure which should be more than 250 mm H₂O in IIH.⁹ At no time our patient had an elevated CSF pressure, including 3 lumbar punctures.

Patients with optic nerve meningocele may present with headache, and transient or permanent visual loss.¹ Ocular findings other than papilledema have been reported as progressive hyperopia, choroidal folds and cystoid macular edema.^{1,8,10}

More than 30 patients with optic nerve sheath meningocele have been described in the literature.^{1-4,11,12} The largest series included 13 cases.¹ All patients presented with visual complaints and three patients had visual acuity worse than 20/200.¹ Three out of 13 patients had neurofibromatosis type 1 (NF1). Two other case reports also described the ectasia of the perioptic dural sheath as an unusual and benign manifestation of NF1.^{4,12} Thus, most common association with dural ectasia of optic nerve sheath appears to be NF1. A case report and the review of the literature in 1997, disclosed total of 31 cases with optic nerve meningocele.² Optic nerve decompression surgery have been performed in 13 of these cases. Although visual acuities were not recorded in all patients, 7 cases showed no visual improvement despite the decompression surgery.² Spooler et al described a child with multiple congenital abnormalities and optic nerve meningocele that

Figure 1 Fluid-filled dilated sheath surrounds normal optic nerves on T2-weighted axial MRI.

progressively expanded and caused displacement of the orbit and complete blindness.¹¹

The most reasonable explanation for the development of the optic nerve sheath meningoceles comes from Hayreh's studies who investigated the relationship of the optic nerve sheath to the optic nerve and stated that congenital narrowing of the optic or congenital cranio-orbital junction anomalies could cause optic nerve sheath meningocele.¹³

Treatment of dural ectasia of the optic nerve sheath depends on optic nerve functions. Carbonic anhydrase inhibitors may be beneficial in some cases.¹⁰ The effects of corticosteroids are not mentioned in the literature. In cases with persistent papilledema, surgical decompression should be considered in the treatment.⁸

In summary, dural ectasia of the optic nerve is an enlargement of the orbital optic nerve sheath, diagnosed by MRI studies and can follow benign or sometimes an unfavorable course. The striking feature of our patient was rapidly progressive visual loss without any mass lesion or intracranial hypertension. The surgical decompression did not improve but preserved the remaining vision. Early surgical decompression of the optic nerve sheath in these patients who present with progressive optic nerve dysfunction may improve vision or prevent the progression.

Disclosures

The authors report no conflicts of interest.

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