Intraventricular Tuberculoma with Profound Visual Loss: A Case Report and Literature Review

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Abstract

Intracranial tuberculoma is a rare manifestation of tuberculosis involving the central nervous system. The involvement of the ventricular system is extremely uncommon. We describe a young woman with bilateral papilledema secondary to intraventricular tuberculoma with hydrocephalus. She was treated with anti-tuberculosis therapy and intravenous dexamethasone. Her visual acuity deteriorated after one month of treatment. We provide a literature review of this uncommon ocular sequelae.

Categories: Neurology, Ophthalmology, Infectious Disease
Keywords: tuberculosis, intraventricular tuberculoma, visual loss

Introduction

Intracranial tuberculoma is a rare manifestation of tuberculosis. Recent data showed that cerebral tuberculoma accounted for 39%-46% of intracranial space occupying lesions [1-2]. It can spread to various portions of the brain and is commonly found in the cerebellum, basal ganglia, and cerebral hemisphere [3]. We report a rare case of intraventricular tuberculoma with hydrocephalus in a young woman who developed profound visual loss during the anti-tuberculosis treatment.

Case Presentation

A 27-year-old woman presented with a progressive painless visual loss in both eyes for one week prior to consultation. It was associated with preceding headache, nausea, and vomiting of two months duration. There was no history of prolonged fever, chronic cough, reduced weight, or loss of appetite. However, her father was treated for pulmonary tuberculosis two years ago and he had completed his anti-tuberculosis therapy.

Her best corrected visual acuity was 3/60 (OD) and 4/60 (OS). The optic nerve function tests were impaired bilaterally and included light brightness, contrast sensitivity, and color vision; however, the right sight was more affected than the left. There was a relative afferent papillary defect present on her right eye. The confrontation test revealed generalized haziness with dense central scotoma on her bilateral vision. The anterior segment examination and intraocular pressure were essentially normal. The fundoscopy examination revealed bilateral papilledema. The optic discs were swollen and elevated with peripapillary flame-shaped
hemorrhages as well as the presence of macular exudates (Figure 1). However, no sign of vitritis or panuveitis was observed.

![Fundus photography of bilateral eyes](image)

**FIGURE 1: Fundus photography of bilateral eyes**
Bilateral papilledema with flame-shaped hemorrhages (blue arrow) and macular exudates (green arrow)

On presentation, she was alert and orientated to time, place, and person. Her vital signs were stable. She was also afebrile. There were no signs of meningism or localizing signs. A respiratory examination revealed crepitation in the right lung. The other cranial nerves examinations were normal. The rest of the systemic examinations were unremarkable.

Her baseline blood investigations were unremarkable except for an increased erythrocyte sedimentation rate (79 mm/h) and C-reactive protein (23 mg/l). The human immunodeficiency virus antibody test was negative. The Mantoux test was positive (16 mm area of induration). The sputum microscopic examination detected the presence of acid-fast bacilli. The chest radiograph showed cavitations at the right lower zone (Figure 2).
The computed tomography (CT) scan of the brain and orbit revealed the presence of multiple ring-enhancing hyperdense lesions with central hypodensity and perilesional edema. The lesions were located at the anterior horn of the right lateral ventricle (Figure 3A-3B). There were also multiple small lesions at the cerebrum and cerebellum (Figure 3A).
Solitary tuberculoma at the anterior horn of the right lateral ventricle (blue arrow). Multiple small parenchymatous discs were observed in the cerebrum (green arrow) with evidence of obstructive hydrocephalus.

However, there was no traction over the septum pellucidum. The lateral and third ventricles were dilated on both sides. She declined further diagnostic tests, including lumbar puncture and magnetic imaging resonance (MRI) of the brain and orbit. A presumptive diagnosis of bilateral papilledema secondary to intraventricular tuberculoma with hydrocephalus was considered. However, the patient refused the ventriculoperitoneal shunt procedure.

Anti-tuberculosis therapy was commenced immediately with oral isoniazid 300 mg, oral rifampicin 600 mg, oral pyrazinamide 1500 mg, intramuscular streptomycin 1000 mg, and intravenous dexamethasone 4 mg. Her symptoms of headache, nausea, and vomiting subsided.

However, visual acuity of both eyes deteriorated progressively to non-perception of light after four weeks of treatment. She had no new emerging neurological symptoms or signs associated with the visual loss. A repeat funduscopy examination showed atrophy of the optic discs in both eyes (Figure 4).

FIGURE 4: Fundus photography of bilateral eyes
Bilateral pale optic disc (blue arrow) with regressing macular exudates (green arrows) after four weeks on anti-tuberculosis therapy.

We planned to extend her oral dexamethasone while continuing the same anti-tuberculosis regime. Unfortunately, the patient defaulted follow-up after one month of treatment.

Discussion
A well-developed intraventricular tuberculoma is not common in tuberculosis infections. The effective blood circulation of the ventricular system acts as a barrier to infection despite evidence of tubercles in the choroid [3–6]. Based on the PubMed search, we encountered 18 published cases of intraventricular tuberculoma from 1987 to 2018 [5–14], including the current case report in Table 1. The age ranged from two years to 62 years. There were eight males and 11 females, including our patient.

<table>
<thead>
<tr>
<th>Author</th>
<th>Year</th>
<th>Sex/Age</th>
<th>Tuberculoma Location</th>
<th>Initial VA</th>
<th>Ocular Finding</th>
<th>Treatment</th>
<th>Follow-Up</th>
<th>Final</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tengku-Fatishah et al.</td>
<td>2018</td>
<td>4 of 8</td>
<td></td>
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<tbody>
<tr>
<td>F/5</td>
<td></td>
<td></td>
<td>Right lateral ventricle and right capsular region</td>
<td>Not available</td>
<td>Not available</td>
<td>Anti-tuberculosis</td>
<td>3 months</td>
<td>Good recovery</td>
</tr>
<tr>
<td>Right lateral ventricle with traction of the septum pellucidum. Another lesion seen in the cerebral hemisphere. Presence of asymmetrical hydrocephalus</td>
<td>Not available</td>
<td>Bilateral papilledema with choroiditis</td>
<td>Anti-tuberculosis, Dexamethasone, VP shunt</td>
<td>Defaulted</td>
<td>After the VP shunt, the patient neurologically improved. However, he defaulted follow-up</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>F/2</td>
<td></td>
<td></td>
<td>Left lateral ventricle, thalamus, cerebral, cerebellar, basal ganglia, and brainstem. Presence of hydrocephalus</td>
<td>Not available</td>
<td>No papilledema</td>
<td>Anti-tuberculosis</td>
<td>12 months</td>
<td>Good recovery</td>
</tr>
<tr>
<td>Hsu, et al. [6]</td>
<td>2004</td>
<td>F/19</td>
<td>Right lateral ventricle. Presence of hydrocephalus</td>
<td>Not available</td>
<td>Mild papilledema</td>
<td>Anti-tuberculosis, Surgical excision, VP shunt</td>
<td>12 months</td>
<td>Good recovery</td>
</tr>
<tr>
<td>Sonmez, et al. [7]</td>
<td>2008</td>
<td>M/22</td>
<td>Right lateral ventricle. Presence of asymmetrical hydrocephalus</td>
<td>Not available</td>
<td>Not available</td>
<td>Surgical excision, Anti-tuberculosis</td>
<td>Not available</td>
<td>Good recovery, however, final VA outcome was not mentioned</td>
</tr>
<tr>
<td>N’da, et al. [8]</td>
<td>2013</td>
<td>F/10</td>
<td>Third ventricle. Presence of obstructive hydrocephalus</td>
<td>Bilateral VA loss</td>
<td>Not available</td>
<td>Surgical excision, Anti-tuberculosis</td>
<td>6 months</td>
<td>Good recovery, however, final VA outcome was not mentioned</td>
</tr>
<tr>
<td>Coulibaly, et al. [9]</td>
<td>2013</td>
<td>M/26</td>
<td>Right lateral ventricle. Presence of asymmetrical hydrocephalus</td>
<td>OU 4/60</td>
<td>Not available</td>
<td>Surgical excision, External ventricular drainage, Anti-tuberculosis, Steroid</td>
<td>3 months</td>
<td>Total disappearance of the lesion. Final VA outcome was not mentioned</td>
</tr>
<tr>
<td>Udayakumaran,</td>
<td>2014</td>
<td>F/27</td>
<td>Third ventricle and thalamus. Presence of Not available</td>
<td>Not available</td>
<td>Anti-tuberculosis, Ofloxacin,</td>
<td></td>
<td>12 months</td>
<td>Good recovery</td>
</tr>
</tbody>
</table>
TABLE 1: Published cases of interventricular tuberculoma from 1987 to 2018

<table>
<thead>
<tr>
<th>Study</th>
<th>Year</th>
<th>Sex</th>
<th>Site</th>
<th>VA, visual acuity</th>
<th>Treatment</th>
<th>Follow-up</th>
<th>Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>present study</td>
<td>2018</td>
<td>F/27</td>
<td>Right lateral ventricle, cerebral, and cerebellum. Presence of obstructive hydrocephalus</td>
<td>3/60 OD, 4/60 OS</td>
<td>Bilateral papilledema Anti-tuberculosis Dexamethasone</td>
<td>1 month</td>
<td>Bilaterally, no perception of light. Patient defaulted follow-up</td>
</tr>
<tr>
<td>Li et al. [14]</td>
<td>2017</td>
<td>M/47</td>
<td>Right lateral ventricle. Presence of hydrocephalus</td>
<td>Not available</td>
<td>Not available Intrahepal isoniazid, Dexamethasone, Anti-tuberculosis</td>
<td>15 months</td>
<td>Disappearance of tuberculoma</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Not available</td>
<td>Not available Intrahepal isoniazid, Dexamethasone, Anti-tuberculosis</td>
<td>7 months</td>
<td>Disappearance of tuberculoma</td>
</tr>
<tr>
<td></td>
<td></td>
<td>M/23</td>
<td>Right lateral ventricle</td>
<td>Not available</td>
<td>Not available Intrahepal isoniazid, Dexamethasone, Anti-tuberculosis</td>
<td>3.5 months</td>
<td>Tuberculoma size decreased</td>
</tr>
<tr>
<td></td>
<td></td>
<td>F/31</td>
<td>Fourth ventricle and right lateral ventricle</td>
<td>Not available</td>
<td>Not available Intrahepal isoniazid, Dexamethasone, Anti-tuberculosis</td>
<td>9 months</td>
<td>Tuberculoma size decreased</td>
</tr>
<tr>
<td></td>
<td></td>
<td>F/82</td>
<td>Fourth ventricle</td>
<td>Not available</td>
<td>Not available Intrahepal isoniazid, Dexamethasone, Anti-tuberculosis</td>
<td></td>
<td>Good recovery</td>
</tr>
<tr>
<td>Sadashiva et al. [13]</td>
<td>2017</td>
<td>M/33</td>
<td>Left lateral ventricle with obstructive hydrocephalus</td>
<td>Not available</td>
<td>Not available Anti-tuberculosis, Endoscopic biopsy, VP shunt</td>
<td>16 months</td>
<td>Good recovery</td>
</tr>
<tr>
<td>Sharma et al. [12]</td>
<td>2017</td>
<td>F/21</td>
<td>Third ventricle with obstructive hydrocephalus</td>
<td>Not available</td>
<td>Bilateral papilledema VP shunt, Surgical excision, Anti-tuberculosis</td>
<td>1 month</td>
<td>Good recovery</td>
</tr>
<tr>
<td>Sachdeva et al. [11]</td>
<td>2017</td>
<td>M/7</td>
<td>Foramen of Monro</td>
<td>Not available</td>
<td>Bilateral papilledema Surgical excision, Anti-tuberculosis</td>
<td>15 months</td>
<td>Good recovery</td>
</tr>
</tbody>
</table>

The hematogenous spread of Mycobacterium tuberculosis through the choroidal plexus or ependyma is the common route of tubercle bacilli entry [3,5-7]. Berthier et al. reported that three of their patients had a lateral ventricle tuberculoma without choroidal plexus involvement [3]. These findings were similar to our patient who developed a right lateral ventricle tuberculoma without signs of choroidal plexus involvement. The tuberculoma in these patients probably arose from the subependymal tuberculare.

Our patient developed a deterioration of visual acuity after four weeks of anti-tuberculosis therapy commencement. We postulate that this could be due to a paradoxical expansion of tuberculoma and the worsening of the pre-existing hydrocephalus. The paradoxical phenomenon has been described in multiple works of literature [14-19]. It is described as an excessive immune response by the host cell to the mycobacterial antigen that causes the...
expansion of the initial lesion or the growing of a new tuberculoma during the anti-tuberculosis therapy [17].

Another possible mechanism is due to poor drug penetration into the ventricular system. Udayakumaran et al. had performed an external third ventriculostomy in a case with a third intraventricular tuberculoma and obstructive hydrocephalus [10]. Despite multiple cerebral spinal fluid redirection procedures that had been performed along with the anti-tuberculosis regime, the tuberculoma was still being intractable. Thus, endoscopic third ventriculostomy was performed to excise the lesion and further cleared the cerebrospinal fluid route simultaneously [10]. Li and colleagues demonstrated a favorable outcome with the intrathecal isoniazid therapy [14].

A variety of treatment modalities had been attempted, as shown in Table 1. Five authors opted for prior surgical excision especially when the diagnosis was doubted [7-9,11-12]. This was due to the rarity of the tumor location or the absence of the systemic manifestation of tuberculosis infection. The lesion may also be imitating malignancy, as there were no specific clinical signs suggesting intraventricular tuberculoma. These patients were started on anti-tuberculosis treatment after a histopathology study confirmed tuberculoma [7-9,11].

Most of the published cases presented with features of hydrocephalus [3-10,12-14]. Six of these patients underwent cerebral spinal fluid redirection procedures, which included a ventriculoperitoneal shunt and external ventricular drainage [3-4,6,9,13-14]. Our patient was treated conservatively with an anti-tuberculosis regime and intravenous dexamethasone because she declined a ventriculoperitoneal shunt and further surgical intervention.

**Conclusions**

Progressive visual loss during the treatment of an intraventricular tuberculoma with hydrocephalus is a challenging clinical scenario. Apart from poor drug penetration into the cerebrospinal fluid, the paradoxical reaction may be attributed to the recalcitrant nature of the disease. A standard guideline on the treatment regime is mandatory. This is very essential to prevent permanent visual loss and further neurological disability.

**Additional Information**

**Disclosures**

**Human subjects:** Consent was obtained by all participants in this study. **Conflicts of interest:** In compliance with the ICMJE uniform disclosure form, all authors declare the following: **Payment/services info:** All authors have declared that no financial support was received from any organization for the submitted work. **Financial relationships:** All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. **Other relationships:** All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

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