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Case Report

Racemose neurocysticercosis

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Abstract

Background: Neurocysticercosis (NCC) is an invasive parasitic infection of the central nervous system caused by the larval stage of the tapeworm Taenia solium. The clinical manifestations of NCC depend on the parasitic load and location of infection, as well as the developmental stage of the cysticerci and host immune response, with symptoms ranging from subclinical headaches to seizures, cerebrovascular events, and life-threatening hydrocephalus. Racemose NCC represents a particularly severe variant of extraparenchymal NCC characterized by the presence of multiple confluent cysts within the subarachnoid space and is associated with increased morbidity and mortality, as well as a decreased response to treatment. Albendazole is the preferred drug for the treatment of racemose NCC due to its superior cerebrospinal fluid penetration compared to praziquantel and the ability to be used concomitantly with steroids.

Case Description: In this report, we describe a 39-year-old man recently emigrated from Mexico with racemose NCC and hydrocephalus successfully treated with prolonged albendazole treatment, high-dose dexamethasone, and ventriculoperitoneal shunt placement for the relief of obstructive hydrocephalus.

Conclusions: Treatment of racemose NCC represents a significant clinical challenge requiring multimodal intervention to minimize infectious- and treatment-related morbidity. We review the clinical, diagnostic, and therapeutic features relevant to the management of this aggressive form of NCC.

Key Words: Neurocysticercosis, racemose, Taenia solium, treatment

INTRODUCTION

Neurocysticercosis (NCC) is a common parasitic infection of the central nervous system (CNS) caused by the tapeworm Taenia solium. Infection is typically characterized by the presence of intraparenchymal cysts within the cerebral hemispheres at the gray-white junction and NCC is the most frequent cause of acquired epilepsy in the developing countries.6,14,23 Parasitic infestation is endemic to South and Central America, Southeast Asia, China, and Sub-Saharan Africa23 and is of increasing importance in the United States due to high volumes of immigration from these endemic areas.6,14,24 Extraparenchymal NCC is less common than the intraparenchymal form and is defined as NCC involving the subarachnoid, meningeal, and intraventricular space. Extraparenchymal NCC presents
unique diagnostic and therapeutic challenges compared to the intraparenchymal form. The racemose variant of extraparenchymal NCC represents a particularly aggressive infection and is associated with relatively increased morbidity and mortality and decreased responsiveness to medical treatment. The variant is characterized by the presence of multilobulated confluent cysts within the basal subarachnoid space that appear “grape-like” both radiographically and pathologically. In this report, we describe the successful management of a patient with extensive racemose NCC and hydrocephalus treated with cerebrospinal fluid (CSF) diversion and prolonged administration of albendazole and corticosteroids with complete radiographic and clinical resolution of infection.

**CASE REPORT**

A 39-year-old male presented to the emergency room with a complaint of worsening headaches and visual decline. The patient emigrated from Mexico 13 months prior to presentation and reported no other significant medical history. He described diffuse headaches that had progressed over a 1-month period with new-onset bilateral visual impairment and nausea for the last week. His neurologic exam was nonfocal, however, the ophthalmologic evaluation revealed bilateral papilledema and diminished visual acuity. He was afebrile on presentation and without meningeal irritation signs. Initial laboratory findings indicated a normal white blood cell count of 7400/mm³ with 73% polymorphonuclear leukocytes, 21% lymphocytes, and 1% eosinophils. A head computed tomography (CT) scan demonstrated hydrocephalus with transependymal flow and a multilobulated hypodense mass in the basal cisterns. Subsequent magnetic resonance imaging (MRI) was performed which demonstrated the presence of too numerous to count nonenhancing, T2-hyperintense cystic lesions in the lateral ventricles and subarachnoid spaces including the bilateral sylvian fissures and cerebellopontine angles, as well as the suprasellar, interpeduncular, and prepontine cisterns with the largest cyst measuring 21 mm in diameter and exerting mild mass effect upon the brainstem [Figure 1]. Minor sylvian fissure enhancement was noted bilaterally, however, magnetic resonance angiography (MRA) showed no large vessel stenosis or occlusion. A screening MRI of the spine demonstrated a small subarachnoid cystic lesion at the level of the conus and multiple septations at the S1–S2 level. A lumbar puncture was performed, and an elevated opening pressure was documented. CSF analysis revealed a mild lymphocytic pleocytosis with 60 white blood cells/mm³ (96% lymphocytes, 21% polymorphonuclear leukocytes, 4% eosinophils). The initial CSF protein level was 106 mg/dL with a glucose level of 54 mg/dL. CSF cysticercus antibody ELISA testing was positive, and a diagnosis of racemose NCC was made based on the immunoassay results, neuroimaging findings, and a history of recent emigration from Mexico.

After extensive discussion between the neurosurgical and infectious diseases teams, we elected to proceed with placement of a ventriculoperitoneal shunt to ameliorate his elevated intracranial pressure and communicating hydrocephalus. High-dose dexamethasone (4 mg QID) was initiated followed by administration of prolonged albendazole (10 mg/kg/day) treatment. The patient experienced immediate relief of his visual deficit following the CSF diversion treatment. He maintained a mild headache disorder that responded well to steroids, and he was able to be slowly weaned off of this medication over a 12 month period. MRI scan performed at that time revealed complete radiographic resolution of his subarachnoid, spinal, and intraventricular cystic lesions with very mild persistent meningeal enhancement [Figure 2]. A serum cysticercus antigen ELISA test was nonreactive, and albendazole was discontinued.

**DISCUSSION**

Cysticercosis is the most common helminthic infection of the CNS with infection occurring primarily through the ingestion of eggs of the parasitic tapeworm *T. solium*. The eggs hatch within the digestive system and the released oncospheres penetrate the intestinal mucosa to enter the circulation. The oncospheres are then passively transported to various tissues including the brain,
Similarly, Colli recently described that extension of the cysts from the ventricles via the CSF may lead to infection within the basal cisterns, sylvian fissures, and cortical and spinal subarachnoid spaces. Excessive basal subarachnoid infection involving numerous parasitic membranes and enlarged vesicular cysts accompanied by a relative lack of scolices is termed racemose NCC, although no definitive diagnostic criteria for this rare but severe variant have been described.\[6\]

The diagnosis of racemose NCC is typically based on neuroimaging, serum and CSF laboratory evaluation, and clinical, epidemiologic data including immigration from or frequent travel to disease-endemic areas. Contrast-enhanced MRI is the imaging modality of choice for the detection of extraparenchymal NCC as CT has limited sensitivity for identification of intraventricular and small subarachnoid cysts.\[16\] Lesions typically appear as hyperintense on T2-weighted sequences and provide a clear delineation of the cyst wall and the presence of obstructive or communicating hydrocephalus. Contrast evaluation is important to determine the extent of the host inflammatory response including ependymitis and basal arachnoiditis, as well as the evolutionary stage of the parasite. While the clinical and neuroimaging findings are of paramount importance, serologic testing may provide additional data, especially in patients without an appropriate epidemiologic history. Peripheral eosinophilia is a common but nonspecific finding with NCC.\[8\] Similarly, the presence of CSF eosinophils is frequently indicative of NCC and is a helpful in differentiating NCC from other forms of inflammatory chronic meningitis, although this finding has been reported to occur in only 15% of patients.\[9\] The most common CSF abnormalities observed with NCC are a mild-moderate lymphocytic pleocytosis and increase in protein with the levels varying according to parasitic burden and location of the infection. Positive serum and CSF immunoassays for detection of anticyticercal antibodies are highly suggestive of NCC infection in the appropriate clinical setting, although false-negative results are a well-recognized limitation.\[8\] The sensitivity of these immunoassays has not been established for the racemose variant but would be expected to be increased given the robust inflammatory reaction typically elicited with subarachnoid infection.

The intraventricular and cisternal forms of NCC typically manifest in a more clinically aggressive manner than parenchymal infection and frequently cause intracranial hypertension due to CSF outflow obstruction or inflammatory basal arachnoiditis.\[21\] Cysts within the ventricular cavities may be free floating and cause obstruction at the foramina of Monro, sylvian aqueduct, or fourth ventricle and may present with rapid clinical deterioration secondary to acute hydrocephalus. These cysts may also become attached to the ependymal wall of the ventricle and result in ependymitis following cyst degeneration that may lead to intraventricular loculation and make CSF diversion more problematic. Similarly, cyst degeneration within the subarachnoid space may elicit an accompanying host inflammatory response with basal arachnoiditis, vasculitis, elevated intracranial pressure, and subacute to chronic hydrocephalus requiring CSF diversion. In 2002, DeGiorgio et al. demonstrated that patients with heavy cyst burdens and hydrocephalus carried the highest risk of mortality with NCC, primarily because of elevated intracranial pressure and shunt-related problems.\[7\] Similarly, Colli et al. reported a 30.8% mortality in patients who required placement of a ventriculoperitoneal shunt and a >80% rate of shunt failure secondary to shunt obstruction due to proteinaceous debris.\[4\] In addition to hydrocephalus, the active inflammatory process may lead to vasculitis and cerebrovascular complications. Angiography or MRA may reveal segmental narrowing or occlusion of large vessels within the basal cisterns and sylvian fissures, with the majority of infarcts occurring in the lenticulostrate artery distributions.\[13\] Basal meningitis may also cause cranial nerve dysfunction due to fibrous entrapment.\[16\] Callacondo et al. recently described that extension of infection to the spinal subarachnoid space is very common in racemose NCC, with spinal involvement documented in 17 of 28 patients in their prospective study leading the authors to recommend that screening MRI of the spine be performed in basal subarachnoid disease to document spinal involvement, prevent complications, and monitor for recurrent infection.\[5^\]
No consensus exists regarding the optimal treatment of racemose NCC including the duration of anthelmintic treatment and the role of surgical intervention. In patients presenting with hydrocephalus, the priority is relief of raised intracranial pressure. Obstructive hydrocephalus due to intraventricular cysts is best managed endoscopically via cyst removal or drainage and often obviates the need for permanent shunt placement. Intraventricular cysts may be freely mobile and imaging immediately prior to neuroendoscopic intervention is recommended to confirm the location of the cyst. A relative contraindication for endoscopic removal of intraventricular cysts is the presence of significant ependymal enhancement as this typically indicates dense adhesion between the parasitic cyst and the ependymal wall with attempted cyst removal associated with neurologic injury and intraventricular hemorrhage. A role for endoscopic or microsurgical removal of subarachnoid cysts within the basal cisterns and sylvian fissures is less established due to frequently high cyst burden and widespread subarachnoid distribution and inflammatory response. Giant (>5 cm) subarachnoid cysts with associated mass effect and intracranial hypertension are most often managed surgically, although Proaño et al. demonstrated complete resolution of giant cysts in 19 of 33 patients treated medically with prolonged multicourse anthelmintic and corticosteroid therapy. Couldwell et al. suggested that surgical therapy be reserved for cisternal forms of NCC if initial medical management fails or experiences neurologic decline as conventional microsurgical approaches have been associated with significant morbidity, likely due to the presence of arachnoiditis with adherence of the cyst walls to cranial nerves and arteries. More recently, several authors have described minimally invasive keyhole or endoscopic approaches to the basal cisterns to achieve cyst drainage, although the utility of these approaches for extensive subarachnoid disease has yet to be established. Fortunately, spillage of cyst contents with these approaches has not proven to exacerbate the inflammatory response in the postoperative period, although careful attention to this concern and copious intraoperative irrigation are recommended.

In our patient with hydrocephalus and relatively rapid progression of visual loss, given the bilateral sylvian fissure involvement and absence of any dominant cystic lesion or focal neurologic deficits, we elected to proceed with placement of a ventriculoperitoneal shunt prior to initiating therapy directed against the invasive infection. Delayed exacerbation of intracranial hypertension frequently occurs several days following administration of cysticidal agents as a result of cyst degeneration and subsequent host inflammatory response with the release of proinflammatory cytokines. For this reason, steroids should be administered concurrently with anthelmintic therapy to reduce the inflammatory response. Subarachnoid cysts are considered less responsive to pharmacologic therapy compared to intraparenchymal lesions, with albendazole the preferred drug for treatment of racemose NCC due to its superior CSF penetration compared to praziquantel. Additionally, the CNS distribution of albendazole is less affected by steroid administration. The dose and duration of treatment must be individually tailored to each patient depending on their parasitic burden and clinical and radiographic response to treatment with the treatment course frequently longer than that prescribed for the parenchymal disease. Aggressive medical treatment has been shown to decrease the incidence of shunt malfunction, which has been directly linked to poorer clinical outcome. For patients with giant cysts, Proaño et al. achieved excellent results with a 4 weeks course of albendazole, although more than half of their patients required more than one treatment course to achieve cyst resolution. 4 of the 33 patients in their study required 3 or more months of therapy and 10 patients also received praziquantel following a failure of the cysts to respond to albendazole.

CONCLUSION

In our patient, we reported the successful clinical resolution of racemose NCC following CSF diversion and prolonged anthelmintic therapy and corticosteroid administration to prevent the development of chronic subarachnoid inflammation. No shunt complications or revisions occurred, and a complete radiographic response was observed. Racemose NCC represents a, fortunately, rare but aggressive form of extraparenchymal NCC often resulting in basilar arachnoiditis, vasculitis, and cranial neuropathy. Proper recognition of this infection and appropriate management with careful consideration of the deleterious effects of the attendant host inflammatory response to cyst degeneration is necessary to reduce patient morbidity and mortality. Racemose NCC accompanied by intracranial hypertension and hydrocephalus is associated with poorer clinical outcomes, often related to ventriculoperitoneal shunt infection and malfunction. Vigilant clinical and radiographic monitoring of response to treatment is imperative to reduce neurologic sequelae and infection relapse as prolonged, or multicourse medical therapy is frequently necessary.

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Conflicts of interest
There are no conflicts of interest.
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