Central nervous system blastomycosis presenting as a year-long chronic headache

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This case describes a posterior fossa mass due to blastomycotic infection in a non-immunocompromised 41-year-old male presenting with a chronic headache for over one year. Given the risk of herniation, no lumbar puncture could be performed. A full work-up found no evidence of systemic infection. Surgical resection helped identify the mass as a blastomycotic abscess. Magnetic resonance imaging characteristics of the mass were helpful in the identification of the mass as a fungal abscess.

Key words: CNS blastomycosis; blastomycotic abscess; chronic headache; posterior fossa mass; fungal abscess

INTRODUCTION

Blastomycosis isolated to the central nervous system (CNS) is exceedingly rare. In the absence of other more common systems’ involvement (such as lungs), blastomycosis is rarely considered high on the differential. This case describes a nonspecific presentation of a blastomycotic abscess as a chronic headache and discusses the difficulties associated with the prompt diagnosis of this disease. The scope of this case is to demonstrate that when presenting as a posterior fossa mass, blastomycotic abscesses have a vast differential and ultimately require a biopsy or resection for the correct diagnosis. MRI imaging may be helpful in helping distinguish fungal abscesses from their neoplastic or bacterial counterparts.

CASE REPORT

A 41 years old male from rural Illinois with a history of hypertension and alcohol abuse presented for evaluation of 16 months of progressive dull occipital headaches, unresponsive to pain medications. Over the prior 6 months the patient developed nausea with rapid head movements as well as dysarthria and right hand clumsiness. Non-contrast computed tomography (CT) of the brain demonstrated an ill-defined mixed density lesion in the cerebellum [Figure 1]. Magnetic resonance imaging (MRI) revealed a 3.5 cm × 2.6 cm intra-axial mass in the midline superior cerebellum abutting the tentorium and compressing the 4th ventricle. The mass was wildly hyperintense on T1-weighted images and heterogeneous but predominately hypointense on T2/FLAIR [Figure 2]. It demonstrated significant internal susceptibility on susceptibility weighted images compatible with vascularity and/or hemorrhage.
was avid internal enhancement of the lesion on post-contrast images. Additionally there was a thin tract of contrast enhancement extending from the lesion along the right lateral margin of the brainstem. There was mild edema of the surrounding cerebellum.

Differential diagnosis of the posterior fossa mass in a 41 year old male included neoplastic (high- or low-grade gliomas, medulloblastoma, hemangioblastoma, lymphoma and infectious etiologies. Pilocytic astrocytomas (WHO grade I) and medulloblastomas (WHO grade IV) occur more frequently in the pediatric population. CNS lymphomas occur more frequently in the elderly or HIV+ population. Hemangioblastomas often occur in association with Von Hippel-Lindau syndrome. Other than the suspicion of alcohol abuse, patient was otherwise not immunocompromised. He was HIV negative, lacked peripheral leukocytosis, fevers, and had an unremarkable CT of the chest. Given the location, the size, and the mass effect of the lesion, no lumbar puncture could be pursued due to the risk of herniation. The patient underwent a suboccipital craniotomy achieving a complete resection. Pathology showed granulomas with fungal organisms. Periodic acid-Schiff and Giemsa (GMS) stains revealed rounded yeast forms consistent with *Blastomyces dermatitidis* [Figure 3]. Post-operative cerebrospinal fluid (CSF) revealed > 1,000 White blood cells with neutrophilic predominance, normal glucose at 64 and high protein at 300.

**DISCUSSION**

Blastomycosis is a pyogranulomatous infection that occurs predominantly by inhalation of the microconidia of the dimorphic fungus *B. dermatitidis*. The fungus is prevalent in the areas along the Mississippi and Ohio river basins as well as Midwestern regions bordering the Great Lakes. About 91% of infections are pulmonary with subsequent dissemination to the skin, bone or genitourinary system. CNS involvement occurs only rarely 5-10% of cases.[1] Isolated CNS blastomycosis is rare. Infection occurs in both immunocompetent and immunosuppressed hosts. For example, in a case series of 22 patients with CNS blastomycosis, only 12 patients were immunocompromised (i.e. HIV, chronic steroid use, anti-tumor necrosis factor therapy for more than 6 months).[2] Clinical and experimental evidence (predominantly animal studies) suggests that chronic alcohol consumption significantly alters many lines of immune system and predisposes alcoholics to an increased risk of infection, increased morbidity, and mortality.[3]

CNS blastomycotic infection can present as either acute or indolent meningoencephalitis. At times, the only symptoms are intractable headaches. In a recent study that evaluated outcomes of 16 patients with CNS blastomycosis, the most frequent symptoms at presentation were headaches or a focal neurologic deficit in 63 and 56% of patients respectively.[4] Other manifestations include leptomeningeal involvement, single or multiple abscesses intracranially or in the spinal cord, as well as in the epidural space causing cord compression. On the MRI these may present as a single or multiple lesions

![Figure 1: Axial head computed tomography image revealing a predominantly hypodense mass lesion in the cerebellum, centered at the vermis. The lesion compresses the fourth ventricle (arrow pointing to the mass)](image1)

![Figure 2: Axial magnetic resonance imaging images of fluid-attenuated inversion recovery (a) and T1 postcontrast (b) at initial presentation and then postoperatively (c and d) revealing complete radiographic resection of the lesion](image2)
producing mass effect (particularly in the cerebellum), diffuse leptomeningeal enhancement, cerebritis or obstructive hydrocephalus. Restricted diffusion is frequently one of the earliest MRI findings with fungal abscesses. This occurs due to an increased cellularity and viscosity of the pus associated with the infection and frequently precedes gadolinium enhancement. Reduced diffusion signal may frequently be heterogeneous. In smaller lesions, it may be punctate. When compared with fungal infections, bacterial abscesses tend to have a highly restricting homogeneous center. In contrast to their marked diffusion abnormality, fungal abscesses may demonstrate only a weak ring enhancement. This is thought to be secondary to a weak peripheral immune response. A combination of ring enhancement and diffusion signal can help differentiate fungal abscesses from bacterial abscesses or brain metastases. Brain metastases tend to have a thicker ring enhancement and a reduced diffusion in the necrotic center. Brain metastases frequently have a thicker ring enhancement, but typically have no reduced diffusion in the necrotic center.[5]

Definitive diagnosis is established either by isolation of the fungus from a culture or direct visualization on the histological slides. Isolation from the CSF is uncommon. In a case series of 22 patients with CNS blastomycosis, CSF cultures were positive only in 2 patients.[2] Serologic testing is generally considered not to be useful in blastomycosis due to high cross-reactivity with other endemic mycoses. Antigen testing may be positive in the urine and serum. PCR is rarely used and typically not commercially available.

Thus, most cases require a biopsy and a histopathologic examination of the tissue to arrive at the correct diagnosis. The case described above had negative serology, CSF culture and required a tissue sample obtained during resection to diagnose it as a blastomycosis abscess.

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There are no conflicts of interest.

REFERENCES