Resolution of aspergillosis in neuroimaging of an immunocompromised patient with pulmonary and cerebral lesions

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**ABSTRACT**

Central nervous system (CNS) aspergillosis is uncommon and considered the most lethal form of aspergillosis. Indeed, current therapeutic strategies such as combination antifungal regimen, neurosurgical resection of infected tissue, and removal of infection source fail to improve the unsatisfactory prognosis of CNS aspergillosis in the majority of the patients. The authors describe a case of chronic pulmonary aspergillosis with concomitant CNS lesions that dramatically responded to antifungal therapy and the CNS lesions resolved in follow-up imaging.

Key words: Aspergillosis, Aspergillus, central nervous system, voriconazole

**Introduction**

Aspergillosis of central nervous system (CNS) is uncommon. It occurs mainly in the setting of disseminated infection (essentially from a pulmonary source), as well as from direct extension, from the paranasal sinuses or direct inoculation through trauma and neurosurgery.\textsuperscript{[1]} CNS aspergillosis is seen almost exclusively in immunocompromised patients and considered the most lethal form of aspergillosis. Indeed, current therapeutic strategies such as combination antifungal regimen, neurosurgical resection of infected tissue, and removal of infection source fail to improve the unsatisfactory prognosis of CNS aspergillosis in the majority of the patients.\textsuperscript{[2,3]}

In this paper, the authors describe a case of chronic pulmonary aspergillosis (CPA) with concomitant CNS lesions that dramatically responded to antifungal therapy and the CNS lesions resolved in follow-up imaging.

**Case Report**

A 53-year-old smoker man was admitted to our hospital with a 5 months history of productive cough associated with intermittent episodes of hemoptysis. The patient was under treatment with prednisone for chronic headache since 2 years before. On admission, physical examination was not remarkable. Laboratory study showed mild leukocytosis and high levels of serum erythrocyte sedimentation rate (ESR) and C-reactive protein (CRP) (188 mm and 118 mg/dL, respectively). In chest X-ray, a cavitary lesion with air fluid level in left upper lobe was seen which was better demonstrated in following thorax computed tomography (CT) scan [Figures 1 and 2].

Microscopic study of sputum specimen did not show any microorganism in different staining. As the patient complained of headache, brain imaging was also performed. Brain magnetic resonance imaging (MRI) revealed multiple ring enhancement lesions with peripheral edema in right temporooroccipital lobe and left cerebellar hemisphere.

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Echocardiogram and blood culture did not show any evidence of endocarditis.

In next step, the patient underwent a CT-guided percutaneous biopsy of the lung lesion. Considering clinical manifestation, laboratory data, and imaging findings, an empiric antimicrobial regimen (intravenous meropenem, vancomycin, and voriconazole) was started. Histopathological study of the lung specimen revealed septated hyaline hyphae with dichotomous acute angle branching among the areas of necrosis surrounded by fibrotic tissue and inflammation. The findings were compatible with the diagnosis of CPA with concomitant CNS lesions. Other microscopic studies for detecting further microorganism, include gram staining, acid-fast staining, and tissue culture, were negative. Hence, the administration of meropenem and vancomycin was stopped after 3 weeks, and treatment with voriconazole was continued. Control chest X-ray and brain MRI were done 2 weeks and also 4 weeks later, which revealed complete resolution of cerebral and pulmonary lesions [Figures 5 and 6]. At the end of the 4th week of treatment, patient’s symptoms include productive cough and headache were resolved, and the serum level of ESR and CRP diminished to 19 mm and 4 mg/dL, respectively.

Intravenous antifungal therapy continued for 12 weeks with careful clinical and radiologic follow-up.

Discussion

In this report, a patient with CPA and concomitant cerebral lesions was described. CPA implies a pattern of aspergillosis in immunocompetent patients that usually progresses slowly over weeks or months. It is characterized by the formation of at least one pulmonary cavity, positive related microscopic or serologic evidence, and the presence of clinical symptoms. Clinical symptoms include weight loss, fatigue, chronic productive cough, hemoptysis, and/or shortness of breath.

Figure 1: Upright chest X-ray showed a consolidation with cavitation in left upper lobe

Figure 2: Axial contrast-enhanced chest computed tomography scan showed a consolidation with cavitation in left upper lobe

Figure 3: Axial contrast-enhanced T1-weighted image showed a ring enhancement lesion in right temporooccipital lobe

Figure 4: Axial contrast-enhanced T1-weighted image showed a ring enhancement lesion in left cerebellar hemisphere
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The major predisposing factors for developing CPA are underlying lung disease (such as tuberculosis and chronic obstructive pulmonary disease) and various degrees of immune dysfunction secondary to the conditions such as diabetes mellitus, alcoholism, and corticosteroid consumption. Our patient was under long-term treatment with prednisone for chronic headache. Corticosteroids have been introduced as a risk factor for both invasive aspergillosis and CPA. CNS involvement of aspergillosis is a diagnostic and therapeutic challenge; since the clinical presentation is nonspecific and the neuroradiologic findings are similar to other CNS fungal, bacterial, and mycobacterial infections, and also brain tumors.

In our patient, multiple ring enhancement lesions were seen in CT scan and MRI of brain. Histopathological study of specimens obtained from biopsy of lung lesion was leading for the diagnosis of aspergillosis. ESR and CRP are nonspecific inflammatory markers in CPA and are usually elevated. Their main role is in the monitoring during treatment.

In this patient, intravenous antifungal therapy alone and without surgical intervention resulted in significant clinical improvement and decrease inflammatory markers. We believe that if the fungal abscesses were secondarily infected with bacteria, the lesions would relapse or would not resolve primarily with pure antifungal treatment. Duration of standard antibiotic regimen for bacterial brain abscess (without surgery) in an immunocompromised patient is at least 8 weeks, while we stopped meropenem and vancomycin within 3 weeks. The course of resolution of lesions in brain MRI was dramatic. All cerebral and cerebellar lesions were resolved within a month. We continued intravenous treatment for 12 weeks and then followed the patients with oral antifungal treatment. After 6 months, no recurrence was observed.

We described a case that was unusual in three respects: First, the patient had minimal neurologic symptom despite multiple cerebral and cerebellar lesions. This can be due to the lack of inflammatory response because of the state of immune dysfunction and receiving high cumulative dose of corticosteroid. Second, the involvement of cerebellum in CNS aspergillosis is very rare. Aspergillosis has a predilection for the involvement of anterior and middle cranial fossa. Third, this patient showed a dramatic response to antifungal therapy alone. In today's medicine, neurosurgical resection of infected tissue is frequently performed as an adjuvant to antifungal therapy of CNS aspergillosis. However, in this case surgical intervention was not required.

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Conflicts of interest
There are no conflicts of interest.

References