Central nervous system Aspergillus fumigatus infection after near drowning


Aims: To report the case of a 26 year old white man, who developed chronic meningitis and intracerebral granulomata 15 days after an episode of near drowning in a swamp.

Methods: Aspergillus fumigatus was isolated from cerebrospinal fluid cultures.

Results: The patient died 70 days after the symptoms were first noticed, and seven days after a subarachnoid haemorrhage. Aspergillus has never been reported before as a cause of intracranial infection after near drowning.

Conclusions: Physicians must be aware of this possibility when confronted with such a situation, because there are now effective therapeutic options for systemic aspergillosis.

Aspergilli are probably the most common group of fungi in our environment. Most aspergillus species are found in the soil. Aspergillosis occurs worldwide, with no apparent age, sex, or race differentials. The major pathogen, Aspergillus fumigatus, is found in the soil, but is seen most abundantly in decomposing organic materials. Aspergillus hyphae can be seen in clinical specimens using direct microscopic examination, and show dichotomous branching of septate hyaline hyphae with a typical acute angle. Aspergillus spp grow well on a variety of conventional agar media, produce spore chains, conidiophores, and conidial heads (L Renon. Recherches cliniques et experimentales sur la pseudotuberculose aspergillare. Thesis. Paris: 1893: No. 89).

Abbreviations: ALT, alanine aminotransferase; AST, aspartate aminotransferase; CAT, computerised tomography; CNS, central nervous system; CSF, cerebrospinal fluid; IV, intravenous; b.i.d., twice a day; t.i.d., three times a day; q.i.d., four times a day; MRI, magnetic resonance imaging.

CASE REPORT

A 26 year old white man suffered near drowning in a mangrove swamp on the seacoast of São Paulo, in eastern Brazil. After being taken to a nearby hospital, he was treated for aspirating pneumonia and was released after two days. Two weeks later he started complaining of headache of progressive intensity, and after 14 days he was submitted to a computerised tomography (CAT) scan and to magnetic resonance imaging (MRI) of the head, which revealed several hypodense lesions suggestive of granulomata. He was then referred and admitted to our clinic. On admission, he was still complaining of headaches, but his physical and neurological examinations were unremarkable, except for the presence of a mild fever (37.2°C) and mild meningismus. His red blood cell count was normal (4.85 × 10¹²/litre; haemoglobin, 132 g/litre) but there was pronounced neutrophilic leucocytosis (20 000 × 10⁶ cells/litre; 84% neutrophils (80% segmented, 4% bands), 10% lymphocytes, and 6% monocytes). His erythrocyte sedimentation rate was 27 mm/h, blood glucose and creatinine were normal. An x ray of the lungs disclosed interstitial micronodular lesions. An electrocardiogram showed no abnormalities. Cerebrospinal fluid (CSF) examination revealed neutrophilic meningitis (165 × 10⁶ cells/litre; 69% neutrophils, 31% lymphocytes), with a normal glucose (3.55 mmol/litre) but high total proteins (0.778 g/litre). Pseudallescheriasis was suspected, and broad spectrum antibiotics, in addition to antifungal treatment were started, with 1 gceftriaxone given intravenously (IV) twice a day (b.i.d.), 500 mg metronidazole IV b.i.d., 2 ml cotrimoxazole IV four times a day (q.i.d.), 200 mg fluconazole IV b.i.d., 4 mg dexamethasone IV three times a day (t.i.d.), and 100 mg ketoconazone IV b.i.d. A stereotactic biopsy was planned but cancelled because a CAT scan performed five days later showed amelioration of the brain lesions. At this time, his blood count was normal, but the leucocytosis had increased (28 500 × 10⁶ cells/litre; 91% neutrophils (83% segmented, 8% bands), 6% lymphocytes, and 3% monocytes). In addition, transaminases were slightly increased (aspartate aminotransferase (AST), 2.32 ukat/litre, alanine aminotransferase (ALT), 2.08 ukat/litre). After 10 days of treatment, the low grade fever and headache recurred. The number of neutrophils in the peripheral blood had increased (32 400 × 10⁶ cells/litre; 82% neutrophils—76% segmented, 6% bands, 10% lymphocytes, and 8% monocytes), and CSF examination showed 280 × 10⁶ cells/litre, mostly (64%) neutrophils, an increased total protein (0.89 g/litre), and a normal glucose (3.05 mmol/litre). Serum AST had returned to normal values but ALT remained somewhat high (1.73 ukat/litre). Serum anti-human immunodeficiency virus enzyme linked immunosorbent assay of blood was negative. Blood cultures and a search for fungi in CSF stains and cultures yielded negative results. After 12 days of treatment the patient became confused and disoriented. Although his CSF findings had slightly improved, 150 mg itraconazole IV b.i.d., together with 30 mg/day amphotericin B IV, was given instead of fluconazole. There was a partial response to treatment, but after two days 300 mg/day IV pentamidine was added to the treatment regimen because an amoebic infection was suspected. The dose of amphotericin B was...
increased to 70 mg/day and given with 0.5 mg amphotericin B intrathecally every other day. Haloperidol (5 mg) was given at bedtime for night agitation. His condition continued to deteriorate over the following days. A few small papular skin lesions ensued, and he suddenly presented a pronounced meningismus, associated with a left conjugate gaze palsy and a left sided weakness, involving mainly his left arm. CSF examination showed a haemorrhage superimposed on the meningitis, and his CAT scan disclosed blood in the subarachnoid spaces, enlarged ventricles, and an almost complete resolution of the granulomata. CSF findings revealed an increase in the number of leucocytes (148.8 x 10^6 cells/litre; 93% neutrophils), an increase in CSF protein (0.29 g/litre), and a decrease in CSF glucose (1.16 mmol/litre). His blood creatinine had risen to 1768 μmol/litre. He was transferred to the intensive care unit. At this time, A fumigatus was isolated from the CSF culture. Ceftriaxone, pentamidine, and cotrimoxazole were discontinued and 3 g ceforoxime IV t.i.d., 600 mg clindamycin IV q.i.d., and 500 mg/day methylprednisolone were added to metronidazole and itraconazole. However, the patient’s condition continued to deteriorate and he died 56 days after being admitted for treatment.

**DISCUSSION**

CNS aspergillosis is a rare condition in immunocompetent hosts, and is favoured by working in agriculture, craftwork, and by a tropical climate. In patients with few risk factors, the entire disease period can last from 9.5 months to four years. Diabetes mellitus type II seems to be a predisposing condition. Primary sites of infection are the lungs in the immunodepressed, and the paranasal sinus in immunocompetent individuals. Less frequently, gastrointestinal and skin infection preceding CNS infection can occur, and more rarely CNS primary infection without an extracranial source. Immunoocompetent patients may present with multiple cranial nerve disorders, a cavernous sinus syndrome, or with an orbital apex syndrome, reflecting local invasion from paranasal sinuses. Isolated or multiple granulomata may also occur and present focal signs, associated or not associated with meningitis. These presentations differ from that seen in the immunodepressed, in whom there are multiple areas of brain infarction, haemorrhage, encephalomalacia, and isolated or multiple abscesses, leading to infectious and/or stroke syndromes. CSF smears should be carefully examined with India ink for fungi, but will rarely be positive. In our patient, special care was taken to avoid confusion with Pseudallescheria boydii, a fungus that may invade the CNS after near drowning, and that may resemble an aspergillosis infection. Another important differential diagnosis is CNS infection by free living amoebae, such as Acanthamoeba spp, and particularly Naegleria spp, which is related to swimming in, or contact with, polluted waters or lakes.

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Because Aspergillus spp are difficult to detect in CSF smears and cultures, the determination of serum aspergillus galactomannan, with two positive results, coupled with typical radiological findings, is highly sensitive and specific to support the diagnosis. In immunodepressed individuals, typical MRI and CAT scan findings in the paranasal sinus and lungs, respectively, may suggest the diagnosis. The CSF determination of galactomannan is also a promising diagnostic tool. A case in which the diagnosis was clarified through MRI findings and CSF polymerase chain reaction to Aspergillus spp has been reported.

Treatment for invasive aspergillosis has changed since our case was studied. Amphotericin B and other clinical and surgical alternatives showed few encouraging results. Although intravenous amphotericin B has been the mainstay treatment for CNS aspergillosis, two new drugs for IV use—voriconazole and caspofungin—are promising agents, with a good tolerability profile.

Pseudallescheria boydii CNS infections have been described after near drowning in dirty and polluted waters or in manure reservoirs. Amoebic CNS infections are reported to occur after swimming in dirty pools or reservoirs. However, before our case, Aspergillus spp had not been suspected as a possible aetiology after near drowning. Because infection with aspergillus can be diagnosed and treated, it should always be considered when involvement of the CNS begins after an episode of near drowning.

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