We report an intracerebral abscess caused by a recently identified nocardial species, in an immunocompetent individual without extraneural involvement.

A 63-year-old non-smoking Caucasian male with an unremarkable history of hypertension, presented with a 2-week history of personality changes including apathy, some difficulty walking, a tendency to veer to the left and a possible seizure. Preliminary assessment, including laboratory investigations were unremarkable. Computed tomography (CT) of his brain revealed a right frontal lobe multi-loculated, ring-enhancing lesion with vasogenic edema and associated mass effect (Figure 1). He was then referred to our institution.

We found the patient afebrile, with no other focal neurological deficits. Repeat investigations including white blood cell count and chest x-ray, were normal. With a presumptive diagnosis of brain tumor, the patient was started on oral steroids and anti-seizure medication. A right frontal craniotomy was performed. Using bipolar cautery, a small corticectomy was made in the abnormal hyperemic brain surface overlying the lesion. An odorless, yellowish, creamy pus-like discharge emanated from just underneath the cortex. Further dissection revealed a thick walled and multi-loculated cavity. Specimens were collected and the site was copiously irrigated to remove the pus-like material. The lateral ventricle was carefully avoided. Broad-spectrum coverage with ceftriaxone and metronidazole was commenced. Tissue histopathology returned consistent with *Nocardia*. Therefore, the antibiotics were changed to intravenous (i.v.) trimethoprim/sulfamethoxazole (TMP/SMX) and metronidazole. Metronidazole was discontinued when the anaerobic cultures returned negative.

Tissue and fluid specimens were cultured on blood and chocolate agars. Gram-positive, non-hemolytic, catalase positive, weakly acid-fast, filamentous, branching bacilli were observed within 72 hours. The organism was presumptively identified as a *Nocardia* species. The Canadian Science Centre for Human and Animal Health confirmed it as *N. paucivorans*, based on 16S rDNA gene sequencing and conventional biochemical tests.

Post-operatively, the patient made a rapid recovery and remained without focal deficits. The CT scan demonstrated a residual cavity behind the area of resection. Further investigations did not reveal any evidence of immunocompromise, or of nocardiosis elsewhere in the body. After a week in hospital, the patient was discharged home on i.v. TMP/SMX.

Trimethoprim/sulfamethoxazole was changed to oral route when a CT, done three weeks postoperatively, showed marked attenuation of the residual abscess. Complete resolution was documented on the CT performed three months postoperatively (Figure 2). Antibiotics were discontinued. When last evaluated nine months after discharge, the patient continued to remain well, with no recurrent abscess on CT.

Nocardiosis is primarily a pulmonary disease. It causes up to 2% of cerebral abscesses, mostly in the immunocompromised. It is a potentially lethal infection, with high mortality unless rapidly diagnosed and treated. The diagnosis is usually made by the demonstration of a characteristic fluffy or ring-enhancing lesion on imaging and by isolation of the organism from clinical specimens.
compromised. Increasingly, it is being recognized in immunocompetent individuals with cerebral abscesses. Primary cerebral abscesses without evidence of pulmonary infection are unusual. The immune status and the timeliness of treatment influence outcome, so that the immunocompetent patient receiving early treatment intervention will do better. To date, 73 Nocardia species have been identified, including 42 additions since 2000. Only a few species are neuroinvasive, with N. asteroides being the most frequent. There are three previous reports of N. paucivorans infection. Two patients were immunocompromised and had intracranial abscesses. The third had chronic pulmonary disease and N. paucivorans was isolated from sputum. In our case, the patient was immunocompetent and without any chronic debilitating disorders.

Based on history and radiological findings, we made a presumptive diagnosis of a neoplastic process, with abscess considered less likely. A slow, afebrile presentation with radiological imaging consistent with brain tumor, has been reported in cerebral nocardiosis. In previous reports, diagnosis of a neoplastic process was revised following operative findings and pathology results. Preoperative magnetic resonance imaging or MR spectroscopy most probably would have made the distinction between an abscess and a tumor. However, that information would not have altered the intent of early surgical intervention.

Figure 2: Computed tomography scan with contrast. Complete resolution of the abscess following 3 months of treatment.

Compared with poor outcomes in other cases, our plan of early surgery and prolonged antibiotics administration, led to a complete recovery. Unlike previous reports, we were able to administer steroids for a week post-operatively without any adverse effects. The duration of antibiotic therapy remains empirical. We used clinical, laboratory and radiological evidence to determine the duration. Antibiotics were administered i.v. for a month and orally for 3 months (total=4 months), effecting a cure. In vitro data supports the efficacy of TMP/SMX in N. paucivorans infections.

This case highlights nocardial cerebral abscesses in immunocompetent patients, where the residual nocardial abscess can be successfully treated with antibiotics. Further studies are needed to better understand the pathogenesis, improve management and establish preventive measures.

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References