A 75-year-old male, a resident of a rural area and a farmer by occupation, visited our outpatient clinic with the symptoms of poor cognition and memory decline over 2 weeks. He denied any history of fever, headache, blurred vision, vomiting or seizure. He was afebrile and his vital signs were stable. There were no lab- oratory abnormalities including leukocytosis or C-reactive protein rising. Upon the neurologic examination, he was conscious and there were no neurologic deficits except intermittent expressive dysphasia and disorientation. Brain magnetic resonance imaging (MRI) was performed because of suspicion of some type of dementia. It showed a 20 mm sized nodular enhancing mass.
with peritumoral edema in the left frontal lobe. The mass had a central cystic portion with diffusion restriction (Fig. 1). High-grade glioma or metastatic tumor was initially presumed based on his age and progressive symptoms. Excisional biopsy was performed for tissue diagnosis. The lesion appeared to be white and took the form of a relatively hard mass with a clear boundary, permitting radical excision of the mass (Fig. 2). Pathological examination revealed multiple necrotizing granulomas with brown pigmented fungal hyphae. Septated hyphae and melanin pigments were confirmed at Fontana-Masson stain consistent with CP (Fig. 3).

The patient was started on intravenous amphotericin B at a dose of 68 mg daily. After 10 days, he was switched to 270 mg of intravenous voriconazole twice a day because of the elevation of serum creatinine. He took the injection for 8 weeks, followed by oral voriconazole 200 mg twice a day for 2 months. A follow-up brain MRI 3 weeks after surgical excision demonstrated a significant resolution of the edema. Ongoing resolution of the lesion was found on the latest follow-up MRI (Fig. 4). He showed dramatic improvement in his symptoms including disorientation and memory disturbance after completion of surgery and antifungal therapy.

**DISCUSSION**

CP is a rare infection caused by darkly pigmented fungi, namely dematiaceous fungi. Dematiaceous fungi represent a group of filamentous molds that contain melanin pigment in their cell walls. *Cladophialophora bantiana* is the most frequently isolated species. *Rhinocladiella mackenziei* (formerly *Ramichloridium mackenziei*) is the second most common cause of CP, which is exclusively endemic in the Middle East area. Most agents are found in soil. Because of this occupational predisposition has been reported in agricultural workers, especially farmers due to risk of soil exposure. CP commonly occurs in the second and third decades of life with male predominance, except *Rhinocladiella mackenziei* which affects adults with a median age of 62 years without male predominance.

The most unique characteristic of CP is its occurrence irrelevant to the immune status of the host. Even though immunodeficiency may play a role as a risk factor, there are many reports of this infection in immunocompetent individuals similar to the patient in this report. The portal entry to brain

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**Fig. 2.** Intraoperative photography demonstrates a white-yellowish and hard mass with well-defined capsule.

**Fig. 3.** Gross preparation shows white, well-demarcated, round masses in brain parenchyma (A). Hematoxylin and eosin stain reveals necrotizing granulomas (red stared, ×100) and inflammatory infiltrates (yellow stared, ×100) (B), as well as brown colored septated hyphae (×1000) (C). Black colored melanin pigments are present in branched fungal hyphae on Fontana-Masson stain (×400) (D).

**Fig. 4.** Magnetic resonance imaging 1 year after surgery depicts complete resolution of abscess and edema in the frontal lobe.
is unclear, although several possible routes have been suggested, such as hematogenous dissemination of inhaled spores or accidental skin inoculation as well as direct extension from adjacent paranasal sinuses or ears\(^{1,3,6,10,11,13,15,17,19,23,25}\). The authors were unable to ascertain the route of infection in the present case. Pathogenesis of CP is associated with the presence of melanin as a virulence factor that provides advantages in evading host defense and crossing the blood-brain barrier by binding to hydrolytic enzyme\(^{1,3,4,16,17}\).

Clinical spectrum of phaeohyphomycosis was listed as a variable, ranging from solitary subcutaneous nodules to a life-threatening infection\(^{3,11,16-18}\). In the central nervous system (CNS) manifestation, brain abscess is a classic clinical presentation\(^ {1,2,3}\). Patients can also present meningoitis, encephalitis, myelitis or arachnoiditis\(^ {17,20}\). Hemiparesis and headache are the most common symptoms followed by various clinical manifestations\(^ {17}\). About 70-80% of cases typically manifest as a single brain abscess particularly on the frontal lobe (52%) like in our case, while multiple brain abscesses can be seen in immunocompromised patients\(^ {6,17,19}\).

The diagnosis of CP can be difficult because dematiaceous fungi are often considered contaminants when identified in culture. Furthermore, the pathogen can not always be cultured and isolated from the serum or cerebrospinal fluid (CSF)\(^ {3,14,17,23}\). Molecular techniques are available to speedily identify these fungi even to the genus level\(^ {17}\). Therefore, diagnosis is made by surgical biopsy. Only the tissue examination can be useful to identify irregularly swollen hyphae with yeast-like structure and to confirm the presence of dematiaceous hyphae in melanin-specific Fontana-Masson stain\(^ {6,17,19}\). Unfortunately in this case, fungus was not identified in the culture of surgical specimen, therefore, the species that causes CP could not be detected. Meanwhile, the brain MRI reveals a ring-enhancing lesion with a low-attenuation core, suggesting the presence of necrosis or pus\(^ {3,17,20}\). In cases where high-grade glioma or metastasis is mimicked by irregular and variably contrast-enhancing lesions, magnetic resonance spectroscopy may be used to differentiate the entities\(^ {6,17,19}\). Imaging findings of this patient were more suggestive of a glioma than an abscess, because nodular heterogeneity on contrast injection mimicked the images seen in high-grade tumors. Consequently, surgical biopsy is essential for the diagnosis of CP.

Because of the rarity of the cases, there is no standard treatment guidelines for CP. A combination of surgical and medical treatments is generally recommended\(^ {1,2,3}\). Complete excision of brain lesions may provide better results than simple aspiration unless the lesion is multiple or is located within the eloquent area of the brain\(^ {1,17}\). Antifungal agents are generally used in combination of amphotericin B, 5-flucytosine and itraconazole because it is associated with improved survival rates\(^ {3,14,17,20}\). Voriconazole can be used as alternative to itraconazole because of its good penetration into both CSF and brain tissue\(^ {3,17,20}\). Duration of taking the medications is still unknown because most reported patients expired during treatment except a few survivors who received voriconazole for about 12 months\(^ {6,19}\). In addition, posaconazole may be a potent drug when pathogen is Rhinocladiella mackenziei\(^ {1,3,4,19}\). In this case, amphotericin B was replaced by voriconazole because of serum creatinine elevation. Amphotericin B has fatal side effects such as nephrotoxicity, therefore, close observation on kidney function is needed.

The prognosis of CP is poor. Mortality rate approaches 100% in untreated patients, while that of treated cases as high as 65% to 73% despite the aggressive treatment\(^ {1,3,6,10,17,19,23}\). Interestingly, mortality rate did not differ significantly between immunocompromised and immunocompetent patients (75% vs. 71%)\(^ {12,17}\). Multiple brain abscesses are associated with worse prognosis than solitary lesion\(^ {6,20}\). Fortunately, the patient reported here had a good response to surgery and chemotherapy and showed fine recovery without any sequela. Solitary lesion and the good general condition of the patient, together with an aggressive therapeutic approach, are therefore inferred to contribute to a favorable outcome. Further studies are necessary to find more potentially useful antifungal regimen for these refractory infections and to investigate more detailed pathophysiology and prognostic factors to increase the survival rate.

**CONCLUSION**

CP is rare disease, but challenging one with high mortality rate, particularly when the CNS is affected. As shown in this report, complete resection and adequate antifungal therapy are the most recommended modality for patients with CP-related abscess to this time.

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