Case Report

A Case Report on Rhinocerebral Mucormycosis of a Post COVID-19 Diabetic Patient

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Abstract
Mucormycosis is an invasive fungal infection caused by different saprophytic environmental fungus occurring predominantly among immunosuppressed patients. Coronavirus disease 2019 (COVID-19) itself and its treatment with immunosuppressive drugs and oxygen delivery systems are setting the scenes for opportunistic and co-infections with fungus. An elderly Bangladeshi woman with background of diabetes mellitus, hypertension and COVID-19, presented with the complaints of nasal obstruction, yellowish nasal discharge, hairing impairment and loosening of teeth. An early diagnosis combined with medical and surgical intervention can decrease mortality and morbidity. This is one of the few cases of post-COVID mucormycosis reported from Bangladesh.

Key word: Rhinocerebral Mucormycosis, COVID-19

Introduction
Mucormycosis (zygomycosis) is a rare opportunistic fungal infection caused by fungi belonging to the order Mucorales and the family Mucoraceae¹. It is the third most common angioinvasive fungal infection after candidiasis and aspergillosis². It usually affects the immunocompromised individuals and rarely causes disease in apparently healthy individuals³.

In the immunocompromised host, mucormycosis infection results from rapid proliferation and invasion of fungal organisms in deeper tissues⁴. The major risk factors for mucormycosis include uncontrolled diabetes mellitus with ketoacidosis, other metabolic acidosis, corticosteroid therapy, organ or bone marrow transplantation, neutropenia, trauma and burns, hematologic malignancy (leukaemia, lymphoma, etc) and iron chelation therapy in patients receiving hemodialysis⁵⁶⁷.

Coronavirus disease 2019 (COVID-19) itself, its treatment with corticosteroids and their immunomodulators, invasive and non-invasive ventilator supports and other oxygen delivery systems, prolonged hospital stay and comorbidities, all are setting the scenarios for opportunistic infections and co-infections with fungus. In case of diabetic ketoacidosis, in high blood glucose and acidic PH, phagocytes become dysfunctional and their chemotaxis and intracellular killing mechanisms become impaired⁸. Infection starts with inhalation of spores through the nose or mouth or even through a skin laceration followed by spread to the paranasal sinuses and subsequently to the orbit, meninges and brain by direct extension⁹. Early identification of the disease and aggressive and prompt medical and surgical interventions are required to prevent the high morbidity and mortality associated with this disease process¹⁰.

Case report
A 56-year-old Muslim housewife, a known case of uncontrolled diabetes mellitus, hypertension and previous SARS-CoV-2 infection with pneumonia reported to ENT department of Combined Military Hospital, Dhaka Cantonment on 27 May, 2021 with the complaints of nasal obstruction and yellowish nasal discharge for last 01 month. These complaints developed after 15 days of recovery from Covid-19 infection. She also complained of gradual loosening of teeth and impairment of hearing for same duration. On examination, she was ill looking and moderately anaemic. Nasal endoscopy revealed left sided DNS along with blackish crusts found in the mucous membrane of the nasal cavity, nasopharynx, posterior
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Hoque et al

Urine R/M/E showed: Albumin ++, Glucose++, ketone

body: absent.

Other biochemical markers were FBS 12.7 mmol/l, HbA1c 8.1%. Her Serum creatinine was 2.2 mg/dl. Her Serum TSH was 1.14 µIU/ml, Serum iron profile: S. iron 31 µgm/dl, S. Ferritin 342.5 ngm/ml, TIBC 285 µgm/dl. Her Serum electrolytes were Na+ 126 mmol/l, K+ 2.3
mmol/l, Cl1100 mmol/l, HCO3 22.1 mmol/l.

Aseptically collected nasopharyngeal swab was examined
under microscope. Wet film preparation, KOH preparation
and Lactophenol cotton blue preparation showed sporangium
with sporangiospores, broad (10-20 µm), nonseptated,
ribbon like hyphae with wide angle branching at irregular
intervals morphologically resemble Mucor species.
Culture in SDA media yielded the growth of fluffy, whitish
to yellow colony of Mucor species after three days of
incubation at 22°С in aerobic condition with optimum PH.
(Fig:1,2,3,4)

Fig 1: Broad, nonseptated, hyphae with sporangium (KOH
preparation 40X)

Fig 2: Whitish to yellow colony of Mucor species in SDA media

Fig 3: Wet film preparation from culture shows broad,
nonseptated, ribbon like hyphae with wide angle branching
at irregular intervals

Fig 4: Staining by Lactophenol Cotton Blue shows wide
angle branching hyphae with spore
Histopathology of tissue from posterior nasal cavity revealed necroinflammatory exudates with fungal ball composed of hyphae and spores arranged in compact clusters which were positive for PAS stain (Fig 5).

![Fig 5: Positive PAS stain shows fungal ball composed of hyphae and spores](image)

Spiral CT Scan of P.N.S were suggestive of fungal acute pan sinusitis with bony erosions at right and left alveolar process of maxilla and bilateral chronic mastoiditis (Fig 6). CT Scan of brain shows no significant changes.

![Fig 6: Spiral CT Scan of PNS shows pansinusitis with bony erosion](image)

She was planned for surgical debridement under general anaesthesia followed by IV antifungal Liposomal Amphotericin B with other supportive therapy.

**Discussion**

Though mucormycosis is an uncommon disease, its prevalence in India is 70 times the world wide estimated rate\(^{11}\). So far 187 cases of COVID-19 associated mucormycosis have been reported in India during this pandemic\(^{12}\). No specific data is available in our country.

Studies have shown that diabetes mellitus alters the immunologic capability to resist mucormycosis through reduction of the phagocytic ability of granulocytes\(^{13}\). In addition, Rhizopus species thrive best in an acidic and glucose-rich environment. Differential diagnosis includes aspergillosis, histoplasmosis, paracoccidioidomycosis. A definitive diagnosis of mucormycosis can only be made by a biopsy and culturing the fungus in the laboratory\(^{14}\). Histologically, mucormycosis is characterized by extensive tissue necrosis, large fungal hyphae, which are nonseptate and have a ribbon-like appearance, with budding and dichotomous branching\(^{15}\). However, in our case, the diagnosis was done immediately by wet film microscopy with aseptically collected specimen and further confirmed by culture in SDA media.

If mucormycosis is suspected, initial empirical therapy with antifungal drugs Amphotericin B should begin. Surgical debridement of the infected and necrotic tissue is the standard treatment along with the medical treatment. In our case we have started antifungal therapy with Amphotericin-B without delay which retard the further angioinvasion. Combined treatment increases the survival to 78% as compared to 57.5% with medical treatment alone. In the presence of intracranial extension, surgical debridement may not be possible and almost all the cases are fatal\(^{16}\). Several surgical procedures have been described ranging from debridement of the mucosa, Caldwell-Luc surgery, medial maxillectomy, ethmoidectomy, sphenoidectomy, radical maxillectomy with orbital exenteration. The standard medical therapy is Amphotericin B in a dose of 1-1.5 mg/kg/day for several weeks depending upon the clinical response and degree of nephrotoxicity\(^{17}\). Other modalities of treatment tried are hyperbaric oxygen therapy and nasally nebulized Amphotericin B, oral Posaconazole\(^{18}\).

**Conclusion**

Early diagnosis by experienced microscopist, surgical debridement and antifungal therapy with Amphotericin B can reduce the mortality and morbidity of mucormycosis cases.

**Conflict of Interest**

The authors stated that there is no conflict of interest in this study.
organisms in deeper tissues. The major risk factors for mucormycosis infection in the immunocompromised host, mucormycosis infection in patients receiving hemodialysis, lymphoma, etc, and iron chelation therapy in patients with corticosteroids and their immunomodulators, invasive trauma and burns, hematologic malignancy (leukaemia, and brain by direct extension. Early identification of the killing mechanisms become impaired. Infection starts with sporangiospores, broad (10-20 µm), nonseptated, and have a ribbon-like appearance, with budding and dichotomous branching. However, in our case, the diagnosis of mucormycosis was confirmed by microscopic examination of the clinical specimens obtained from the patient’s nasal cavity.

**Culture and Identification**

Culture in SDA media yielded the growth of fluffy, whitish and co-infections with fungus. In case of diabetic ketoacidosis, amphotericin B with other supportive therapy. Amphotericin-B without delay which retard the further reduction of the phagocytic ability of granulocytes. In our case we have started antifungal therapy with posaconazole.

**Histopathology**

Histopathologically, mucormycosis is characterized by extensive necroinflammatory exudates with fungal ball clusters which were positive for PAS stain. Histopathology of tissue from posterior nasal cavity revealed necroinflammatory exudates with fungal ball.

**Clinical Management**

Early diagnosis by experienced microscopist, surgical debridement and antifungal therapy with Amphotericin B anesthesia followed by IV antifungal Liposomal amphotericin B. She was planned for surgical debridement under general anesthesia followed by IV antifungal Liposomal amphotericin B. She was planned for surgical debridement under general anesthesia followed by IV antifungal Liposomal amphotericin B. The authors stated that there is no conflict of interest in this study.

**References**