Bronchial Necrosis from Mucormycosis: A Case Report and Review

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Abstract
Disseminated pulmonary mucormycosis is an uncommon and aggressive life threatening condition associated with a high mortality rate. Pulmonary mucormycosis can spread to contralateral lung before it is even detected, involving bronchial necrosis leading to a bilobectomy with tissue flap coverage. Although our report describes a fatal outcome, this case evaluates the role for surgery in the treatment of difficult infections.

Keywords: Pulmonary mucormycosis; Bronchial necrosis

Introduction
Mucormycosis is a lethal infection among immunocompromised hosts. Pulmonary mucormycosis can spread to contralateral lung before it is even detected. We present a case of pulmonary and disseminated mucormycosis in a young diabetic female after a tooth extraction. A 22-year-old female was admitted to an outside hospital after she was found unresponsive in her home. She was taken to an outside hospital where she was found to have diabetic ketoacidosis, septic shock, acute respiratory failure, and acute renal failure. Two days prior to her admission she underwent a tooth extraction. She began complaining of fevers, chills, headaches, changes in vision, cough, shortness of breath, chest pain, abdominal pain, nausea, vomiting, diarrhea, and a rash soon after this procedure. She did not present to a physician until her significant other was unable to awake her. Her past medical history is significant for poorly controlled diabetes mellitus type I, with recent Hemoglobin A1c of greater than 14 percent and occasional methamphetamine use. She has had no prior operations nor other significant family or social history.

Upon admission at the outside hospital she was found to be in septic shock with lactate of 11 mMol/L, diabetic ketoacidosis, acute respiratory failure requiring mechanical ventilation, and acute renal failure. After cultures were obtained, antibiotics were started. A CT scan was performed on the abdomen and pelvis revealing pneumatisis intestinalis and pneumobilia. General surgery was involved and no acute intervention was pursued as her exam was benign. She improved and was extubated after a few days. Later the same day, she had a cardiac arrest and was re-intubated. An echocardiogram was done revealing an ejection fraction of 65-75 percent with mild tricuspid and pulmonary valve regurgitation. A bronchoscopy was performed concerning for necrotizing infection of the right middle and lower lobes. A CT scan of the chest was performed, demonstrating diffuse necrosis of the right lower lobe and peri-bronchiolar necrosis of the right middle lobe (Figure 1). Blood cultures produced positive results for Escherichia coli and Candida glabrata. A sputum culture produced an unspecified organism, concerning for mucormycosis. She was transferred to our institution for further care.

Upon transfer, she remained intubated, on 40 percent oxygen with saturations measuring 99 percent. She was awake with a residual right sided foot drop after her recent cardiac arrest. Her white blood cell count was 10.6 bil/L on admission with hemoglobin 9.6 g/dL, Creatinine 0.6 mg/dL, albumin 2.9 g/dL, and transthyretin 8.6 mg/dL. HIV rapid screen was negative. Other labs were within normal limits. Her vital signs were within normal limits for her age and she did not require vasopressor support. A repeat CT scan of the head, chest, abdomen, and pelvis was performed. The CT scan of the chest revealed increasing opacities and the presence of a small pneumothorax (Figure 2). The CT scan of the abdomen revealed a loculated fluid collection measuring 9 cm × 7 cm, concerning for an abscess and a small lesion in the spleen concerning for infarct or abscess. The CT scan of the head showed mild ventricular predominance, concerning for early communicating hydrocephalus.

Infectious disease, pulmonary medicine, thoracic surgery, and critical care physicians were involved with her care. She was given supportive measures, including Intravenous fluids, insulin, nutrition, antibiotics, and antifungals. She was given piperacillin-tazobactam 4.5 g intravenous every 6 h, vancomycin 1000 mg intravenous every 12 h, and voriconazole 200 mg intravenous every 12 h. Voriconazole was started because initial fungal cultures showed Candida krusei and Aspergillus species, not A. fumigatus. After further discussion with *Corresponding authors: Jessica Heimes, Department of Thoracic Surgery, Loma Linda University, 11234 Anderson St. Schuman Pavilion, Suite 1617, Loma Linda, CA, USA, Tel: 909-558-4354; Fax: 909-558-0348; E-mail: jheimes@llu.edu
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with patchy involvement of the middle lobe parenchyma was present. After resection bronchoscopy was performed and the bronchial stump appeared viable at the staple line. This bronchial staple line was covered with an intercostal pedicled muscle flap.

The specimen was sent to pathology for evaluation and the patient had invasive mucormycosis, involving the vascular margins, with extensive parenchymal necrosis and hemorrhage. Broad based hyphae were found in the lung vasculature and parenchyma (Figure 4). Fungal cultures had been sent but were negative for growth after 4 weeks.

She was extubated on postoperative day 2 and received aggressive pulmonary treatments. Antifungal medications were adjusted and the patient was started on liposomal amphotericin B at 10 mg/kg intravenous every 24 h, caspofungin 100 mg intravenous every 24 h, and posaconazole 300 mg intravenous every 12 h. Ophthalmology and ENT were consulted for dilated eye exam and exam for invasive Mucor to the sinuses and Candida glabrata. No evidence of invasive Mucor was identified in these areas. Interventional radiology drained the pelvic fluid collection and cultures were sent with no evidence of mucormycosis. She was unable to swallow without aspiration after extubation and a nasojejunal feeding tube was placed for nutrition. Her chest tubes were removed on postoperative day 6. She was reintubated for respiratory failure on postoperative day 12 and was unable to be weaned from the ventilator. A percutaneous tracheostomy was placed on postoperative day 16 for prolonged intubation. On postoperative day 21 she arrested and was given multiple rounds of epinephrine and chest compressions. She was resuscitated but air and fluid was seen coming from her chest tube sites. Her tracheostomy tube was exchanged over a bronchoscope for an endotracheal tube which was placed in the left mainstem bronchus for exclusion of the right lung. Upon evaluation of the right bronchial stump, a large bronchopleural fistula was identified. A chest tube was replaced on the right side for drainage of air and fluid.

After further discussions with her family and the poor prognosis associated with her disease, the family decided to move to comfort care and withdraw life support on postoperative 22. The patient expired and no autopsy was performed.

Discussion

Mucormycosis is a rare fungal infection from Rhizopus, Mucor, Lichtheimia, Cunninghamella berthollethiae species [1,2]. The spores are generally inhaled or enter through open wounds and then germinate in the host forming hyphal elements [1-4]. Rhinocerebral and pulmonary forms are the most common manifestations of mucormycosis encountered at 55 and 30 percent respectively [5]. Disseminated mucormycosis occurs in 9 percent of cases [5]. Pulmonary mucormycosis infections usually occur in patients who are immunocompromised, such as transplant recipients (especially lung and liver transplant patients), chemotherapy patients, chronic corticosteroid therapy, renal failure, or even diabetics [2].

In healthy individuals, mononuclear and polymorphonuclear phagocytes eliminate fungal spores and hyphae by oxidative and non-oxidative killing mechanisms [2,6,7]. Defects in phagocytic cell activity permit unrestricted growth of the hyphal form and lead to invasive infection, such as those seen with hyperglycemia and acidosis [2,6-8]. Ibrahim et al. [9] discovered a glucose regulated protein 78 (GRP78) as a novel host receptor that interacts with Rhizopus, mediating invasion and damage of human endothelial cells by the fungus [2,10]. Elevated glucose and iron levels upregulate GRP78 expression and promote endothelial cell invasion and damage [2]. The immunosuppressed patient has an increased risk of infection with Mucor species.
Corticosteroids impair migration, ingestion, and phagolysosome fusion in human macrophages, increasing the susceptibility to *Mucor* infections [2,6,7]. Patients in iron overload states, including those patients undergoing chelation therapy with deferoxamine are predisposed to *Mucor* infections. Deferoxamine eliminates the fungicidal effect of serum and increases *in vitro* fungal growth by acting as a siderophore for *Mucorales* species [2,11-13]. Iron chelators such as deferiprone and deferasirox which lack xenosiderophore activity and have been shown to be protective in animal models of mucormycosis [2,12]. Some case reports show a potential beneficial effect of deferasirox therapy in *Mucor* patients [2,14]. Diabetic individuals have a tendency for increased serum iron secondary to impaired transferrin binding, leading to increased infection [4].

Pulmonary mucormycosis is the second most common presentation of *Mucor* in diabetics, while the rhinocerebral form is most common in this population [4]. *Mucor* species flourish in acidic environments, therefore renal failure and diabetics have increased susceptibility [1]. In addition, this species has a ketone reductase enzyme that allows it to thrive in high glucose environments [4]. Prolonged treatments with voriconazole has been associated with an increased incidence of these infections because it is not active against these fungi [3,4].

*Mucor* hyphae are well known to be angioinvasive, causing hemorrhage, thrombosis, infarction, and tissue necrosis of invaded tissues. Pulmonary mucormycosis spreads rapidly when the spores spread into bronchioles and alveoli, causing pulmonary parenchymal necrosis and a high mortality rate [1]. Endobronchial lesions are found in a third of pulmonary mucormycosis patients leading to high mortality rates secondary to obstruction of the airways and vascular invasion leading to hemorrhage [4]. The in-hospital mortality rate with isolated pulmonary mucormycosis is 65 percent. If the disease is disseminated the mortality rate reaches 98 percent [1]. The most common cause of death is fungal sepsis related to disseminated disease. *Mucor* can spread rapidly to the contralateral lung and distal organs when treatment is not started in a timely fashion [2,15]. Pulmonary mucormycosis patients usually die from disseminated disease before respiratory failure occurs. However, dissemination is rarely detected antemortem [2,15,16].

High resolution CT scan is the best method of determining the extent of pulmonary mucormycosis and typically shows evidence of infection earlier than standard chest radiographs [2]. Nodular opacities without an air bronchogram are the most common findings on CT scan and may be accompanied by multiple nodules or pleural effusions [2,17,18]. Halo and air-crescent signs are associated with mucormycosis, but in centrally located lesions there is increased risk of pulmonary artery erosion and hemoptysis [2]. Mucormycosis has been associated with a reverse halo sign, described as a focal round area of ground-glass attenuation surrounded by a ring consolidation [2,16,17].

Diagnosis is difficult in mucormycosis because tissue swabs, sputum, and bronchial alveolar lavage fluid are usually nondiagnostic [2,19]. Despite invasion of blood vessels, blood cultures rarely grow *Mucorales* species [2]. Mucorales appear as broad nonseptate hyphae with branches occurring at right angles [20].

Mucormycosis is treated with antifungals, surgery, reducing immunosuppression, control of blood glucose, and discontinuation of deferoxamine treatment [2,13,21]. Amphotericin B is the initial treatment of choice and is the only approved drug for the treatment of mucormycosis [3,4]. The largest reported case series of mucormycosis included 24 patients who received a lipid complex of amphotericin B with overall response rate of 71 percent without significant toxic effects, even in those with pre-existing renal disease [2,22]. Doses have been suggested from 5 mg/kg/day to 10 mg/kg/day up to 15 mg/kg/day [2,4,23-25]. Further trials are ongoing to determine appropriate doses of amphotericin B.

Posaconazole may be an alternative therapy for intolerance, or used in conjunction with amphotericin B [3]. The dosage of posaconazole was 800 mg daily in four divided doses which showed an overall success rate of 70 percent in 24 patients with minimal toxicity [2,26-28]. A retrospective review of posaconazole based salvage therapy in 91 patients with refractory mucormycosis revealed an overall success rate of 61 percent [2,29]. Posaconazole is an oral medication with limitations associated with absorption, especially those with poor oral intake [2,7,8]. The steady-state plasma concentrations of the drug are not reached until 1 week of therapy has started [2,7,8]. However, the data is not complete. Some studies have found combination therapy beneficial while others have found superior results with amphotericin monotherapy [5]. Antifungals such as voriconazole and caspofungin are not effective against *Mucor* and prophylaxis with these drugs has been shown to be associated with *Mucor* infections [4]. However, a combination of amphotericin B and caspofungin was associated with improved therapy in 41 patients with biopsy-proven rhino-orbital cerebral mucormycosis [2,30]. Duration of therapy should be individualized.

Surgical debridement is indicated in patients with isolated mucormycosis involving one lung and should be performed early in those who can tolerate the procedure [1,31]. Antifungal agents usually have poor penetration at the site of infection secondary to thrombosis and tissue necrosis, therefore surgical debridement is of paramount importance [2]. Tedder et al. [31] found a benefit to aggressive surgical resection in pulmonary mucormycosis. They found a 9.4 percent mortality rate in patients treated with surgical resection compared to 50 percent mortality in those treated with medical therapy alone [1,31]. An additional review found the mortality rate in those who received antifungal therapy alone for pulmonary mucormycosis was 55 percent, compared with those who received both antifungals and surgery [2,15]. Lobectomy, and sometimes pneumonectomy, may be required for improvement. The benefit of pulmonary resection decreases after the disease disseminates.

Other treatments have been proposed to combat Mucorales. One such therapy mentioned in the literature include hyperbaric oxygen, especially for rhinocerebral disease [2,32]. Cytokine therapy using interferon gamma and granulocyte-macrophage-colony stimulating factor have been proposed to improve the ability of phagocytic cells to kill Mucorales species [2,6-8,13]. Iron chelators to starve the fungus of iron stores have been proposed using drugs such as deferiprone and deferasirox [2,7].

**Conclusion**

A high index of suspicion for mucormycosis is necessary as less than 50 percent of cases are diagnosed antemortem [4]. This case is unique because mucormycosis was diagnosed antemortem and the bronchoscopic findings reveal a classic description of the damage *Mucor* species can inflict in susceptible patients. Although rare, mucormycosis can cause devastating effects in the immunocompromised host, as seen in this case. In this patient, surgical resection of the necrotic tissue would give this patient this best chance of survival. Resection was performed because it was felt she would have the best chance of survival at such a young age. Unfortunately, her disease was diffuse and she succumbed to her infection. Further research is needed to improve both diagnosis treatment of this troublesome disease.
