Mucormycosis Of The Paranasal Sinus With Gas Forming Maxillary Osteomyelitis – A Case Report

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Citation


Abstract

Mucormycosis (MM) is a deep, opportunistic mycosis caused by filamentous fungi included in the order Mucorales that belong to the class Zygomycetes of fungi. It is recognized as one of the most rapidly progressive lethal form of fungal infection in human beings with a high mortality of 70–100%. Among the conditions predisposing to MM are Diabetes Mellitus, malnutrition, haematological malignancies, neutropenia, burns, surgical procedures, occlusive dressings, antibiotics, long term steroid therapy and immunosuppressive therapy.

INTRODUCTION

Mucormycosis (MM) is a deep, opportunistic mycosis caused by filamentous fungi included in the order Mucorales that belong to the class Zygomycetes of fungi. It is recognized as one of the most rapidly progressive lethal form of fungal infection in human beings with a high mortality of 70–100%. Among the conditions predisposing to MM are Diabetes Mellitus, malnutrition, haematological malignancies, neutropenia, burns, surgical procedures, occlusive dressings, antibiotics, long term steroid therapy and immunosuppressive therapy.

The term osteomyelitis designates inflammation of the bone and its marrow cavity. Osteomyelitis and Mucormycosis have been known to be associated with intraosseous gas collection. Intraosseous gas collection is usually observed in various pathologic conditions, including osteomyelitis due to certain gas-forming organisms (known as emphysematous osteomyelitis), osteonecrosis, postoperative and posttraumatic conditions.

Bone involvement as a part of Mucormycosis is most commonly seen with the rhino cerebral form. Involvement of other bones such as femur, spine and temporal bones has also been reported in literature. However emphysematous changes associated with Mucormycosis induced osteomyelitis is thought to be extremely rare. The presence of gas forming osteomyelitis of the maxillary bones due to Mucormycosis of the paranasal sinuses has not yet been reported in literature.

We present in this report a case of Mucormycosis of the naso-orbital region in a known diabetic with gas forming osteomyelitis of the facial bones with a review of relevant literature.

CASE REPORT

A 50 year old male farmer from a low socioeconomic, rural background presented to the emergency department of a tertiary referral centre in Bangalore, South India with a 10 days history of pain, swelling and progressive loss of vision in the right eye associated with a week long history of headache and facial pain. He was a diagnosed case of, poorly controlled, non insulin dependent diabetes mellitus (NIDDM) since 8 years on irregular treatment with oral hypoglycemic drugs.

On clinical examination there was facial fullness over the cheek bones bilaterally. They were tender on palpation along with facial soft tissue crepitus. On Anterior rhinoscopy extensive foul smelling blackish crusts over the floor of the nasal cavity along with septal perforation involving the
anterior part of the cartilage was seen. The oral cavity showed extensive erosion of the hard palate which was covered with foul smelling slough. A detailed ophthalmological examination confirmed total internal ophthalmoplegia with complete loss of vision and proptosis of the right eye. The left eye however maintained a normal visual acuity.

Blood investigations revealed elevated blood sugars (280mg/dl) and low serum potassium levels (3.4Meq/dl). Urine ketone bodies were present significantly and the Arterial Blood Gas (ABG) analysis indicated a state of severe uncompensated metabolic acidosis. Nasal crusts were sent for microbiological investigation. Potassium hydroxide (KOH) preparation of the sample revealed thick aseptate fungal filaments strongly suggestive of zygomycosis (Fig-1).

Figure 1
Fig-1. Potassium hydroxide (KOH) preparation of the nasal crusts showing thick aseptate fungal filaments suggestive of zygomycosis.

A contrast enhanced computerized tomography (CECT) of the paranasal sinuses (PNS) revealed bilateral extensive PNS infection with features suggestive of bilateral maxillary bony osteomyelitis with multiple air filled pockets. A nasal septal perforation with mucosal ulcerations was noted on the nasal septum and the hard palate. Extensive inflammation of buccal mucosa on both sides was present along with involvement of the soft tissue of the pterygopalatine fossa. Orbital cuts revealed bilateral orbital cellulitis with the right side being more affected than the left along with right optic nerve sheath inflammation (Fig-2).

In view of multiple air pockets a possibility of an anaerobic infection was suspected along with the working diagnosis of Mucormycosis with facial osteomyelitis. He was posted for an endoscopic debridement of the crusts within the nasal cavity under general anesthesia. Intra-operatively, abundant black foul smelling crusts and necrotic tissue was found over the floor of nasal cavity, lateral wall and the septum. The cartilaginous part of nasal septum showed extensive necrosis along with a perforation. Minimal cheesy material was evacuated from within the maxillary and ethmoidal sinuses on both sides. A bilateral Maxillary antrostomy, anterior and posterior ethmoidectomy was performed endoscopically. The Lamina propria was opened on both sides. Bilateral inferior turbinectomy was also carried out in view of the extensive necrosis and a formal Septoplasty was also executed. Following this the hard palate was also debrided. Pus and mucosal tissue from the maxillary antrum was sent for a repeat microbiological and histopathological evaluation. The Gram’s stain revealed numerous gram negative bacteria and a few gram positive cocci. Aerobic culture yielded Escherichia coli sensitive to carbapenem and amino glycosides. No pathogenic organisms were isolated on anaerobic culture.

A Mycological workup included a KOH staining, calcofluor-white staining and culture of the tissue and pus sample. Broad aseptate fungal filaments were seen on KOH/calcofluor examination. Culture on Sabarauks Dextrose agar yielded grayish white cotton candy like colonies. These isolates were identified as Rhizopus species. Histopathological findings confirmed evidence of fungal

Figure 2
Fig-2. A contrast enhanced computerized tomography (CECT) of the paranasal sinuses (PNS) revealed bilateral extensive PNS infection with features suggestive of bilateral maxillary bony osteomyelitis with multiple air filled pockets
infection especially zygomycosis associated with vascular invasion and extensive osteomyelitis of the bony fragments (Fig-3).

**Figure 3**
Fig-3. 400X Haematoxylin and eosin stain of the tissue obtained after endoscopic debridement shows features of fungal infection by zygomycosis with evidence of vascular invasion and osteomyelitis. Also seen are abundant gram positive cocci and gram positive bacilli.

Post operatively patient was treated with intravenous Amphotercin infusion in 5 percent dextrose solution at 50 milligrams per day for a total dose of 2 grams which was completed in 6 weeks. Care was taken to avoid exposure of Amphotercin to sunlight and the intravenous drip set was adequately covered. Serum potassium levels were monitored to look out for hypokalemia. Blood sugars were maintained with subcutaneous insulin administration and oral hypoglycemic agents. Weekly nasal endoscopy and removal of crusts was performed along with saline nasal douching. Antibiotic coverage for gram negative bacteria and anaerobes was added and he was administered intravenous metronidazole along with third generation cephalosporins.

He responded remarkably to the treatment and attained complete recovery after 2 months of monitoring and medications. Vision improved immediately after surgery and reached normal visual acuity by one month. He was followed up monthly over the next 6 months to observe for any recurrence of disease. Follow up included monthly diagnostic nasal endoscopy, vision tests and three monthly KOH for nasal secretions which were subsequently negative for fungal disease.

**DISCUSSION**
Mucormycosis (MM) is a deep, opportunistic mycosis caused by filamentous fungi included in the order Mucorales that belong to the class Zygomycetes of fungi. It is recognized as one of the most rapidly progressive lethal form of fungal infection in human beings with a high mortality of 70–100%. Among the conditions predisposing to MM are Diabetes Mellitus, malnutrition, haematological malignancies, neutropenia, burns, surgical procedures, occlusive dressings, antibiotics, long term steroid therapy and immunosuppressive therapy. The site of Mucormycosis can be determined by the nature of predisposing infection. (Table 1)

**Figure 4**
Table 1- Relationship between predisposing condition and the resulting predominant site of infection.

<table>
<thead>
<tr>
<th>Predisposing condition</th>
<th>Predominant site of infection</th>
</tr>
</thead>
<tbody>
<tr>
<td>Diabetic ketoacidosis</td>
<td>Rhinocerebral</td>
</tr>
<tr>
<td>Neutropenia</td>
<td>Pulmonary and disseminated</td>
</tr>
<tr>
<td>Corticosteroids</td>
<td>Pulmonary, disseminated, or rhinocerebral</td>
</tr>
<tr>
<td>Dextrose</td>
<td>Disseminated</td>
</tr>
<tr>
<td>Malnutrition</td>
<td>Gastrointestinal</td>
</tr>
<tr>
<td>Trauma, catheter/injection site, skin maceration</td>
<td>Cutaneous/subcutaneous</td>
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</table>

Our patient had poorly controlled non insulin dependent diabetes mellitus which is a very common predisposing factor for a fungal infection such as Mucormycosis. This microbe causing Mucormycosis may be present in the oral cavity, nasal passages, throat and stools of healthy patients without any apparent clinical signs of infection. Although several genera are associated with this disease, the most common are Rhizopus, Rhizomucor and Absidia. Rhizopus is the predominant pathogen accounting for 90% of the cases of rhino cerebral Mucormycosis. Mucormycosis of the rhino cerebral type (39%)presents most commonly as an acute infection followed by the pulmonary type (22%), the disseminated and cutaneous type (16% each), the gastrointestinal type (4%), and type at other locations (3%).

Fungal infection usually originates from the paranasal sinuses. This then invades the blood vessels and subsequently spread through them. Once fungal hyphae enter into the blood stream they can disseminate to other organs such as cerebrum or lungs which can be fatal. Mucor hyphae form thrombi within the blood vessels that reduce vascularity to the tissues and cause necrosis. Disseminated involvement of Mucormycosis is observed in diabetics with ketoacidosis, which favours rapid proliferation of fungus and its invasion into the orbit and cerebrum. (Fig- 4)
The initial symptoms of rhino cerebral Mucormycosis are consistent with either sinusitis or periorbital cellulitis that include eye or facial pain and facial numbness, followed by the onset of conjunctival suffusion, blurry vision, and soft tissue swelling. Fever is variable and may be absent in up to half of the cases. If untreated, infection usually spreads from the ethmoidal sinus to the orbit, resulting in loss of extra ocular muscle function and proptosis. Marked chemosis may also be seen. The infection may rapidly extend into the neighboring tissues if not treated at the earliest. Our patient presented with acute symptoms of headache, facial pain, swelling and progressive loss of vision. The examination findings of blackish slough in nasal cavity and oral cavity were suggestive of fungal infection. The bilateral tenderness and crepitus found over cheeks was unusual and indicated some form of gas accumulation within the bones. Since zygomycosis is not gas-forming, this infection might not have been from a direct result of a fungal infection. An associated infection with a gas forming bacteria was considered at the time of diagnosis as bone necrosis caused by secondary bacterial infection in necrotic tissue is a possibility.

Zygomycosis (ZM) is reported to cause osteomyelitis of skull bones, tibia, cuboidal bones and vertebrae in immunocompromised patients. As it spreads through the perivascular spaces, bone destruction is uncommon and late finding in ZM even when the surrounding tissue is extremely involved. In our patient histopathological examination of the tissue biopsy revealed evidence of fungal infection by Zygomycetes associated with vascular invasion and extensive osteomyelitis of the bony fragments along with abundant gram positive cocci and gram positive bacilli. The pus Culture indicated the presence of abundant Escherichia Coli. Till date Mucor has been associated with gas formation only in case of emphysematous gastritis, where it occurred along with other gas forming bacteria such as Escherichia coli.

The presence of air collections in the soft tissue could be due to infectious and non infectious aetiology. It is predominantly associated with infections due to anaerobic pathogens like Clostridium, Bacteroides, and anaerobic streptococci or mixed infections with aerobic and anaerobic bacteria such as Klebsiella, Escherichia coli, Salmonella, Streptococcus, and Staphylococcus. Therefore gas-forming osteomyelitis can be associated with Clostridium, Bacteroides, E. coli, Klebsiella, Streptococcus, and Mycobacterium.

The imaging findings of rhino cerebral Mucormycosis on CT and MR imaging are diagnostic. These include soft-tissue opacification of sinuses with hyper dense material, nodular mucosal thickening, and an absence of fluid levels in the maxillary, ethmoidal, frontal, and sphenoid sinuses, in decreasing order of incidence. Sinus contents have a variety of MR signal characteristics, including T2 hyper intensity or marked hypo intensity on all sequences, possibly secondary to the presence of iron and manganese in the fungal elements. Typically, proptosis occurs because of enhancing soft-tissue masses crowding the orbital apex and the cavernous sinuses. Thickening and lateral displacement of the medial rectus muscle are characteristic of orbital invasion from disease in the ethmoidal sinuses. Lack of enhancement of the superior ophthalmic vein or ophthalmic and internal carotid arteries may be seen and is related to vasculitis and thrombosis. In our patient a contrast enhanced computerized tomography (CECT) of the paranasal sinuses (PNS) revealed bilateral extensive PNS infection with features suggestive of bilateral maxillary bone osteomyelitis with multiple air filled pockets. This finding is not common for Mucormycosis of the paranasal sinuses.

The diagnosis of zygomycosis almost always requires a histopathological evidence of fungal invasion of the tissues. Obtaining a fungus in culture from a potentially infected site is rarely sufficient to establish the diagnosis of
Mucormycosis because the causative agent is ubiquitous, may colonize normal persons, and is a relatively frequent laboratory contaminant.

However, a microbiological study including culture of the lesion is important to identify the organism. With the emergence of many new species as pathogens, culture based diagnosis will help establish a data. Histology of zygomycosis characteristically shows hyphae that are wide, non-septate, branching at right angles, and look empty when haematoxylin and eosin stained. Three principles in the patient management were followed. Firstly, control of diabetes for which the patient was advised insulin therapy and dietary restrictions. Secondly, removal of the necrotic bone, which acted as a nidus of infection and prevented the action of systemically, administered antifungal drugs (due to thrombosis of blood vessels). Finally, an early empirical parenteral antibiotic therapy with a broad range of antibiotics covering Gram-negative and anaerobic bacteria. The antifungal drug of choice is amphotericin, preferably the lipid formulation because adequate dosage can be delivered with reduced renal toxicity. Newer antifungal agents such as voriconazole and caspofungin have not shown significant activity against Mucor in clinical and in vitro studies. The new triazole antifungal agent, posaconazole, has been compared with amphotericin B, voriconazole, and itraconazole in vitro studies. This drug could have potential for clinical development in the treatment of zygomycosis. Surgical resection of the involved tissue along with amphotericin improves survival and removes the nidus of infection. Conventional treatment regimes with amphotericin and newer antifungal medications do not usually include simultaneous coverage specifically for gram negative gas forming bacteria and on many occasions surgical debridement is deferred if patient is mildly symptomatic. If the osteomyelitis is left untreated, it allows for the fungal infection to invade deeply as necrotic bony tissue is a rich nidus for further infection leading to dreaded complications at a much faster rate. Hence the need for evaluation for coexistent gram negative bacterial infection and its subsequent management with surgical debridement along with a multidisciplinary team approach to identify early complications and sequelae is strongly recommended.

CONCLUSION

This report highlights a rare case of Mucormycosis of the paranasal sinuses in a relatively young diabetic with associated gas forming osteomyelitis of the facial bones in a very short duration of just 10 days. It most importantly emphasizes the need for an immediate intervention and management in such a case keeping in mind the possibility of a rapidly progressing bony osteomyelitis that can augment the precipitation to fatal intracranial and intraorbital complications in a much unexpected and rampant manner. Conventional treatment regimes with amphotericin and newer antifungal medications do not usually include simultaneous coverage specifically for gram negative gas forming bacteria and on many occasions surgical debridement is deferred if patient is mildly symptomatic. If the osteomyelitis is left untreated, it allows for the fungal infection to invade deeply as necrotic bony tissue is a rich nidus for further infection leading to dreaded complications at a much faster rate. Hence the need for evaluation for coexistent gram negative bacterial infection and its subsequent management with surgical debridement along with a multidisciplinary team approach to identify early complications and sequelae is strongly recommended.

References

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