Mucormycosis of the hard palate masquerading as carcinoma

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Abstract

A growing number of medically compromised patients are encountered by dentists in their practices. Opportunistic fungal infections such as mucormycosis usually occur in immunocompromised patients but can infect healthy individuals as well. Mucormycosis is an acute opportunistic, uncommon, frequently fatal fungal infection, caused by a saprophytic fungus that belongs to the class of phycomycetes. Among the clinical differential diagnosis we can consider squamous cell carcinoma. Such cases present as chronic ulcers with raised margins causing exposure of underlying bone. There is a close histopathological resemblance between mucormycosis and aspergillosis. Microscopically, aspergillosis has septate branching hyphae, which can be distinguished from mucormycotic hyphae by a smaller width and prominent acute angulations of branching hyphae. A definitive diagnosis of mucormycosis can be made by tissue biopsy that identifies the characteristic hyphae, by positive culture or both. The culture demonstration of the organism in affected tissue is essential for early diagnosis and management.

Case Report

A 50-year-old male patient was referred to the Oral and Maxillofacial Surgeon for a progressive non-healing wound with a foul odor and ulceration of the palate and left maxilla of about 2 months duration. The patient had undergone extraction of left maxillary lateral incisor and canine 3 months earlier due to poor peri-odontal status. Following extractions the socket never healed completely. Later, he developed headache, sinusitis and noticed a non-healing ulcer on the palate. The patient was moderately built and nourished with normal gait. He also gave a medical history of uncontrolled diabetes mellitus and hypertension. Over the past 2 months the patient was advised various antibiotics and analgesics by different dental practitioners as well as general physicians, and had very limited benefit. The patient also gave a history of tobacco chewing from past 15 years.

Extraorally no swelling was noted and the overlying skin appeared normal with no local rise in temperature or any other secondary changes.

On intraoral examination, an infiltrating ulcer approximately 3×4.5 cm² with irregular borders was appreciated over the hard palate. Ulcer was covered over by the necrotic slough and on the anterior part of the ulcer; the underlying bone was also exposed. Ulcer was not tender with signs of erythema over the margins (Figure 1). Maxillary left lateral incisor and canine were missing.

A provisional diagnosis of squamous cell carcinoma was made. The differential diagnosis of midline lethal granuloma and fungal infection were thought. Later the patient was sent for blood investigations, which revealed raised ESR (34 mm/1st hour) and random blood glucose level 460 mg/100mL. Tridot, HBs test and chest X ray were non-contributory. The computed tomography (CT) scan revealed focal destruction of the bone of anterior palate and in the left anterior part of the maxilla (Figure 2).

The patient underwent a partial maxillectomy (Figure 3) and the specimen was creamish white, non-vital, measured about 5×3 cm with rough surface (Figure 4). Microscopic exami-
nation of the specimen revealed bony spicules and dead lamellated bone around the fatty marrow showing ischemic necrosis. Numerous non-septate, broad, branching hyphae were seen within the necrotic marrow tissue (Figures 5 and 6). Thus, a final diagnosis of mucormycosis was arrived. A strict diabetic control and a course of amphotericin B were advised. He was rehabilitated with an obtur- tor for the oro-nasal fistula (Figures 7 and 8) and was being followed up for the past 6 months and has no further complaints.

Discussion

Mucormycosis is a saprophytic aerobic fungus commonly found in the environment in bread moulds or decaying vegetation. This organism is frequently found to colonize the oral mucosa, nasal mucosa, paranasal sinuses and pharyngeal mucosa of asymptomatic patients. These fungi do not usually cause dis- ease in healthy people with intact immune sys- tems, but patients with a number of conditions can be predisposed to the development of invasive fungal disease.12,13

In 1885, Paltauf14 documented the first histologic description of generalized mucormyco- sis in a 52-year-old patient. Since then, oppor- tunistic infections caused by fungi of the order Mucorales have been recognized in associa- tion with diabetes, hematologic malignant dis- eases, immunosuppressive therapy, thermal burns and surgery.15

Infection arises through inhalation of spores and contamination of the traumatized tissue, ingestion and direct inoculation.15,16 The hallmarks of disease caused by these organisms are angioinvasion, thrombosis, ischemia and necrosis of involved tissue.17-19

High incidence of mucormycosis is seen in uncontrolled diabetic patients, because they produce the enzyme ketoreductase, which allows them to utilize the patient’s ketone bodies. It is also likely that the hyperglycemia stimulates fungal growth and the reduced chemotaxis and phagocytic efficiency permit innocuous organisms to proliferate.4

Oral manifestations of mucormycosis are frequently the first clinical signs to arise, probably because of the highly vascularized struc- ture of oral soft tissues. Furthermore, it has been suggested that vascular ruptures and bleeding due to dental extractions may create a portal of entry for fungi into the maxillofacial regions.20 Intraorally, the hard palate is usually affected because of its proximity to the infection of the nasal fossa and paranasal sinuses. In addition, isolated intraoral involve- ment is extremely rare.21

The diagnosis of mucormycosis is usually based on a clinical picture revealing the inva-
sive course of the disease and is confirmed by biopsy, where the specimen will show broad, non-septate hyphae with branching at right angles and invading tissue. However, negative culture results often appear despite positive histologic findings.

Differential diagnosis of destructive lesions of the palate and maxilla should include squamous cell carcinoma, nasopharyngeal carcinoma, peripheral T-cell lymphoma (formerly termed midline lethal granuloma), chronic granulomatous diseases like tuberculosis, tertiary syphilis, Wegener’s granulomatosis and other deep fungal infection.

The present case was provisionally diagnosed as squamous cell carcinoma of the palate which presented as chronic necrotizing ulcer with raised margins and exposure of the underlying bone. Similar cases are reported in the literature by few authors recently.

On the contrary, a recent report suggests that mucormycosis confined to the oral tissues without any systemic involvement is very rare. When mucormycosis presents as an isolated intraoral infection, the ulcers may also have a white necrotic-appearing tissue without black scab which may be due to a necrotic infarction of the palate. The lesion will be usually deep and causes denudation of the palate along with osteomyelitis of the affected bone. In such situations; the differential diagnoses should also include acute necrotizing gingivitis or periodontitis, herpetic infections and traumatic ulcers. In more severe cases, the extensive mucormycosis may resemble malignancies, cancrum oris, pyogenic orbital cellulitis and other systemic mycoses.

Treatment of mucormycosis consists of surgical debridement; systemic antifungal therapy and treatment of any underlying condition are most effective methods. Control and prevention of opportunistic fungal infection in patients suffering from debilitating diseases such as diabetic ketoacidosis, immunodepression, blood dyscrasia, solid organ transplant, patients on long term steroids and bone marrow transplant is very important. Once mucormycosis infection diagnosed in debilitated patients, it must be treated propitiously, without any delay, by different modalities, medically and surgically.

Conclusions

Mucormycosis is a life-threatening fungal infection that frequently occurs in immunocompromised patients. These infections are becoming increasingly common, yet survival remains poor. Since mucormycosis occurs rarely in the oral cavity, it may pose a diagnostic and therapeutic dilemma for those who are not familiar with its clinical presentation. A greater understanding of the pathogenesis of the disease may lead to future therapies. Early diagnosis and management of the lesion will localize and lessen the chances of spreading to the adjacent tissue and may have great advantages.

The case presented here emphasizes the concept that simple procedures such as dental extractions can cause catastrophic complications in susceptible patients. In the present scenario, dental professionals may encounter patients whose general health is compromised. We must remain cautious in our efforts to perform procedures and follow such patients to be certain that appropriate healing occurs.

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