COVID-19 associated mucormycosis in head and neck region of children during current pandemic: Our experiences

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ABSTRACT

Aim: Mucormycosis is a rare but severe fungal infection, particularly found in immunocompromised patients. Currently this dreaded mucormycosis is rising among COVID-19 paediatric patients during their treatment period or after their discharge from hospital. It is also called as black fungus. The aim of this study is to evaluate the COVID-19 associated mucormycosis (CAM) in head and neck region of the paediatric patients with its clinical manifestations and management.

Material and methods: This is a descriptive and prospective study of paediatric patients with COVID-19 associated mucormycosis (CAM) carried out at a postgraduate teaching hospital. This study was conducted between March 2020 to April 2021. Patient profile such as age, sex, comorbidities, clinical presentations, diagnosis and treatment of the CAM were analysed. There were 12 paediatric patients of CAM were enrolled in this study.

Results: Out of 12 paediatric patients of CAM, there were eight male and four female patients, aged from 3 years to 16 years. Out of the 12 patients, 3 were diabetic (25%). Three patients (25%) were taking prolonged systemic steroids with prolonged hospital ICU stay. Two children (16.66%) were under chemotherapy for acute leukaemia. The common clinical symptoms were facial swelling, facial pain, nasal block and nasal discharge. Diagnosis was confirmed by histological examination. All were treated with endoscopic surgical debridement and amphotericin B. Two patients were passed away; one was due to cerebral involvement and another was due to respiratory failure by pneumonia.

Conclusions: Early identification and prompt treatment in paediatric patients with CAM are required. Aggressive endoscopic surgical debridement for local control and appropriate systemic antifungal treatment will help to improve the prognosis and survival of the patients.

KEY WORDS:
COVID-19 associated mucormycosis, pediatric patient, endoscopic surgical debridement, amphotericin B.

INTRODUCTION

Mucormycosis is a dreaded fungal infection with characteristic feature of angioinvasion [1]. It is caused by a fungus belong to the order Mucorales of the class Zygomycetes. This fungus is saprophytic in nature and often found in soil, vegetables, respiratory and digestive system of the human. Rhizopus species (approximately 44%) and Mucor species (approximately 15%) are most commonly seen in the mucormycosis [1]. The critical ill COVID-19 patients those were admitted to the intensive care unit (ICU) and required mechanical ventilation or had prolonged duration hospital stays and those are taking systemic steroid for prolonged period are likely to get co-fungal infections such as mucormycosis, called COVID-19 associated mucormycosis (CAM) [2, 3].
The histopathological study, direct, microscopy and culture from the clinical samples are the important diagnostic modalities for the CAM [4]. The CAM in children is less frequently documented in the literature. The objective of this study is to analyse the detail of clinical profile and management of the CAM in children.

MATERIAL AND METHODS

PATIENTS

This is a prospective study conducted in 12 children of either sex in the age group of 3 to 16 years suffering from CAM during March 2020 to April 2021. Our Institutional Ethics Committee (IEC) approved this study with the reference number IEC/IMS/SOA/13/06.03.2020. COVID-19 paediatric patients infected with mucormycosis during treatment period at COVID hospital or after discharge from the COVID hospital were included in this study. All of them reverse transcription polymerase (RT-PCR) positive for viral RNA and diagnosed COVID-19 at the time of hospitalization. The COVID-19 paediatric patients without mucormycosis or Non-COVID-19 paediatric patients with mucormycosis were excluded from this study. Patients of CAM older than 18 years of age were excluded in this study.

METHODS

For each registered child, the detail clinical record included demographic data, predisposing factors, clinical presentations, investigations, treatment and treatment outcome. The clinical presentations, diagnosis, treatment and outcome of the CAM among children were studied.

DIAGNOSIS OF MUCORMYCOSIS

All the enrolled paediatric patients underwent diagnostic nasal endoscopy for assessing the bilateral nasal cavity and nasopharynx. Direct microscopic examination with 10% potassium hydroxide (KOH) was utilised for confirming the broad aseptate hyphae. Culture was done in Sabouraud dextrose agar, Sabouraud dextrose with chloramphenicol agar and yeast extract agar. Biopsy was done in all cases and the histopathological study included haematoxylin and eosin, periodic acid-Schiff and Grocott-Gomori’s methenamine silver (GMS) staining. Computed tomography (CT) scan of the nose and paranasal sinus and magnetic resonance imaging (MRI) were done to find out the extent of the diseases into orbit and brain. During nasal endoscopy, the tissue from the nasal cavity sent for microscopy, culture and histopathological examination showing broad non-Septate fungal hyphae with right-angled hyphae branches. All paediatric patients underwent endoscopic debridement of the mucormycosis along with extermination of the orbit in two cases, followed by parenteral infusion amphotericin B (1-1.5 mg/kg/day) and total dose of 2.5-3 gm.

STATISTICAL METHODS

SPSS Statistics for Windows, version 20, was used for all statistical analyses (IBM-SPSS Inc., Chicago, IL, USA).

RESULTS

There were 12 paediatric patients of CAM were enrolled in this study with age range from 3 to 16 years. Out of 12 paediatric patients of mucormycosis, there were 8 boys (66.66%) and 4 girls (33.33%) with male to female ratio of 2:1. Out of the 12 patients, 3 (25%) were diabetic. All the 3 diabetic mellitus patients were under treatment with oral hypoglycaemic agents/insulins regularly, but their blood sugar was poorly controlled. Two children (16.66%) were diagnosed with acute leukaemia, three patients (25%) were taking high dose of steroids and one (8.33%) had taken tocilizumab during the treatment of the COVID-19 infection. Out of the 12 patients, 6 (50%) were diagnosed with sinonasal mucormycosis, 4 (33.33%) had rhino-orbital mucormycosis, 1 (8.33%) had oronasal (nasal and palatal involvement) type of the mucormycosis and 1(8.33%) had rhino-orbital-cebral mucormycosis (Table 1). The children presented with foul smell nasal discharge, nasal block, headache, eye pain, proptosis, facial swelling and blurring of vision or loss of vision. All the 12 paediatric patients presented with foul smelling nasal discharge and nasal block. Out of the 12 patients, 7 (58.33%) of them were presenting with facial pain but 6 (50%) were presenting with facial swelling (Figure 2). Three (25%) children were presenting with headache, two (16.66%) had proptosis, one (8.33%) had nasal septal perforation and one (8.33%) had altered sensorium. Before the surgical debridement, the nasal swab sent for KOH mount where all patients showed asceptate hyphae. Culture of the nasal discharge showed *Rhizopus oryzae* in nine patients and the rest showed no growth. All patients underwent endoscopic surgical debridement under general anaesthesia. All the patients were also administered intravenous infusion of amphotericin B. In this study, out of two patients (16.66%) of death, one was due to cerebral involvement and another case died because of the respiratory failure by
pneumonia during treatment at COVID hospital. Patient follow-up was done after 6 months’ interval after surgery.

DISCUSSION

The current COVID-19 pandemic originated in Wuhan, China, in December 2019 and became global pandemic because of its rapid spread [5]. The spectrum of clinical presentations of symptomatic COVID-19 patient ranges from mild to critical [6]. COVID-19 patients usually show higher levels of inflammatory cytokines (interleukin (IL)-2R, IL-6, IL-10 and tumour necrosis factor-alpha), impaired cell-mediated immune response, affect both CD4+ T and CD8+ T cells [7, 8]. COVID-19 patients often treated with steroids and immunomodulators which impair the immune system of the patient. So, COVID-19 patients have susceptibility towards fungal co-infections such as mucormycosis [9]. Mucormycosis is a rare, fatal, angio-invasive and opportunistic fungal infection among children which can affect any organ of the body [10]. In the head and neck region, rhino-orbital-cerebral infection is common type of the mucormycosis infections [11]. The mucormycosis infection accounts for approximately 10% of all mycotic infections [12]. Although immune deficiency is an important risk factors for co-fungal infections, there are several other diseases also provide chance for mucormycosis in children. Hematological malignancy is an important risk factor for mucormycosis [13]. In one study, the incidence of mucormycosis among children with acute lymphoblastic leukemia (ALL) was 2.2% [14]. Acute lymphoblastic leukemia and increasing age were two important risk factors for mucormycosis.

| Patient’s serial | Age (years) | Sex | Affected part | Clinical presentations | Co-morbid diseases | Treatment | Outcome |
|------------------|------------|-----|---------------|------------------------|-------------------|-----------|---------|
| 1                | 3          | M   | Sinonasal     | Facial swelling, facial pain, nasal discharge, nasal block | Acute lymphoblastic leukemia | Endoscopic surgical debridement plus amphotericin B | Cured |
| 2                | 5          | F   | Naso-orbital  | Facial pain, nasal block, nasal discharge, facial swelling, nasal septal perforation | Prolonged use of steroids in COVID hospital | Endoscopic surgical debridement plus amphotericin B | Cured |
| 3                | 7          | M   | Sinonasal     | Facial pain, nasal block, nasal discharge | Acute lymphoblastic leukemia | Endoscopic surgical debridement plus amphotericin B | Cured |
| 4                | 10         | M   | Oronasal      | Facial pain, palatal black eschar, nasal discharge, nasal block | Uncontrolled diabetes | Endoscopic surgical debridement plus amphotericin B | Cured |
| 5                | 11         | F   | Sinonasal     | Facial pain, nasal discharge, nasal block | Non-Hodgkin’s lymphoma | Endoscopic surgical debridement plus amphotericin B | Death due to respiratory failure |
| 6                | 12         | F   | Naso-orbital-cerebral | Facial swelling, headache, altered sensorium, proptosis, nasal discharge, nasal block | Prolonged use of steroids in COVID hospital | Endoscopic surgical debridement plus amphotericin B | Death due to rapid spread to brain |
| 7                | 12         | M   | Naso-orbital  | Headache, orbital pain, nasal discharge, nasal block | Uncontrolled diabetes | Endoscopic surgical debridement plus amphotericin B | Cured |
| 8                | 14         | M   | Sinonasal     | Facial swelling, facial pain, nasal discharge, nasal block | Non-Hodgkin’s lymphoma | Endoscopic surgical debridement plus amphotericin B | Cured |
| 9                | 15         | M   | Naso-orbital  | Facial swelling, facial pain, proptosis, nasal discharge, nasal block | Treated with tocilizumab | Endoscopic surgical debridement plus amphotericin B | Cured |
| 10               | 15         | M   | Sinonasal     | Facial swelling, numbness over face, nasal discharge, nasal block | Uncontrolled diabetes mellitus | Endoscopic surgical debridement plus amphotericin B | Cured |
| 11               | 16         | M   | Sinonasal     | Headache, Numbness over face, nasal discharge, nasal block | Use of systemic steroids | Endoscopic surgical debridement plus amphotericin B | Cured |
| 12               | 16         | F   | Naso-orbital  | Headache, nasal discharge, nasal block | No co-morbidity | Endoscopic surgical debridement plus amphotericin B | Cured |
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Factors for developing mucormycosis [14]. In our study, 16.66% of the children were diagnosed with acute lymphoblastic leukemia. Other risk factors are hematopoietic stem cell transplants or solid organ transplant recipient. Solid organ cancer (without transplant) is not usually associated with mucormycosis [15]. Diabetes mellitus is also another important risk factor with 9% to 36% of the cases of mucormycosis found in diabetes [16]. In our study, 25% cases were diabetic. Injudicious use of steroids during treatment of COVID-19 infections lower the immunity is also an important risk factor for getting mucormycosis infection. Systemic steroid not only lowers the immunity and but also increases the blood glucose level in the blood and so act as risk for getting fungal infection [5]. In this study, 25% of the children were taking systemic steroid for longer period. In some children, no specific underlying risk factors are associated with mucormycosis [17]. In our study, one (8.33%) patient had taken tocilizumab during the treatment of the COVID-19 infection. Tocilizumab is an immunomodulator which impair the immunity of the patient. In our study, one case (8.33%) had no risk factors for causing mucormycosis. Iron overload, deferoxamine therapy, intravenous drug use, kidney diseases are less common in children but remain well recognized risk factors for causing mucormycosis in children [18]. Burns and traumatic ulcers/wounds may be associated with cutaneous mucormycosis [18].

The diagnosis of the CAM may be hampered by their non-specific symptoms and signs such as bloody nasal discharge, facial swelling, facial pain, conjunctival suffusion, blurry vision which may mimic to common infectious etiologies [19]. The important point behind successful management is early identification, elimination of the predisposing factors, aggressive surgical debridement and infusion of systemic antifungal agents. Diagnosis may be delayed because of its non-specific clinical manifestation. Timely diagnosis is the key behind the prompt treatment of the CAM in children. Clinical suspicion and early diagnosis with prompt treatment are key steps for preventing the morbidity and mortality of this dreaded clinical condition like rhino-orbital-cerebral mucormycosis [20]. Proper history taking, physical examination and imaging are key component for diagnosis of the suspected mucormycosis. In CAM, computed tomography (CT) scan will often reveal the bone destruction. Brain magnetic resonance imaging (MRI) is helpful to rule out any intracranial or orbital involvement [21]. MRI of the brain may show multiple areas of infarction and ischemia indicating invasive fungal disease. In case of unstable hemodynamic and poor respiratory status with inability to keep the patient in supine position without oxygen desaturation made unfeasibility for performing MRI. Bedside diagnostic nasal endoscopy can be done in a timely manner and histopathological processing in case of active COVID-19 infection are useful for starting the treatment for rhino-orbital mucormycosis. Mucor is usually demonstrated via a nasal biopsy and subsequent culture. Tissue is sent for histopathological examination and KOH mount which confirm the mucormycosis [22]. Fresh tissue examination with KOH/Calcofluor-white is helpful to establish a rapid diagnosis. Frozen section is shortening the time of the diagnosis and improving
the outcome this dreaded infection. Direct microscopy, histopathology and culture from the clinical samples are the major diagnostic modalities for the mucormycosis [23].

Mucormycosis requires an urgent intervention, because of rapid progression and destructive nature of this infection [24]. Delayed initiation of treatment results in higher chance of mortality. Aggressive surgical debridement of the necrotic tissue till getting healthy bleeding tissue is useful for better drug penetration to the affected sites. Histopathological examination post-operatively are advised to ensure complete clearance of the lesion. The standard medical treatment for CAM is amphotericin B in a dose of 1-1.5 mg/kg/day for weeks to months on the basis of the clinical response [25]. However, the exact duration of the treatment is not well established [26]. The less toxic type of amphotericin B is liposomal form, colloidal dispersable form and lipid complex are administered in greater dose (3-5 mg/kg/day with little side effects. Topical use of amphotericin B is also useful but it has limited availability and high cost. Posaconazole is an important alternative antifungal drug but its use has not been established in pediatric age group [25]. In few selected patients of mucormycosis, hyperbaric oxygen is helpful [26]. In this study, a combined endoscopic surgical debridement and medical treatment with amphotericin B was the mainstay of the treatment. Antifungal treatment is usually insufficient to cure mucormycosis as angioinvasion and thrombosis of the vessels in mucormycosis prevent optimal penetration of the antifungal agents to the site of infection. Surgical debridement of the infected necrotic tissue is usually required along with systemic antifungal agents [27].

Despite intensive use of broad-spectrum antifungal and adequate surgical debridement, the mortality rate of these patients is more than 40% since many decades [28]. The mortality rate in the pediatric patients of CAM is high, particularly in immunocompromised children. The high mortality in pediatric patients with CAM is associated with untreated cases, disseminated infection at age less than one year [29]. A study showed six cases of invasive mucormycosis during treatment of malignancies where all received liposomal amphotericin B and four received surgical excision along with antifungal treatment and two of them died where 1 (17%) was due to mucormycosis [30]. The outcome of invasive mucormycosis in non-transplant children resulted in approximately 90% success rate [31]. The use of liposomal amphotericin B contributed to positive outcome and survival in invasive mucormycosis. Invasive opportunistic fungal diseases are very important causes for morbidity and mortality in children with cancer and those with allogenic haemopoietic stem cell transplantation [32]. Apart from different underlying co-morbidities in adult age group, invasive fungal diseases like mucormycosis in infants, children and adolescents are unique with respect to epidemiology, diagnostic techniques, pharmacology and dosing antifungal medications absence of phase 3 clinical trials for evidence based decisions [32].

Poor oral hygiene during oxygen administration and use of unsterile, unclean oxygen providing medical devices are also responsible for co-fungal infections to the COVID-19 children. So, the oral care should be done and devices giving oxygen must be maintained with optimum sterility [11]. High index of clinical suspicion, early diagnosis and prompt treatment can improve the survival of the children. Sometimes the diagnosis is delayed by the clinicians those are not much familiar to this disease COVID-19 children with mucormycosis.

CONCLUSIONS

Mucormycosis is a fatal fungal infection resulting in vascular invasion by the hyphae leading to thrombosis and necrosis of the host tissue. Paediatric patients of haematological malignancies, diabetes mellitus or taking systemic steroids or under any immunosuppressive medication with COVID-19 infection are at higher susceptibility to mucormycosis. In COVID-19 children, the severity of the mucormycosis is due its rapid progression and angio-invasive nature. Paediatrician or paediatric otorhinolaryngologists should act promptly to identify the mucormycosis particularly in immunocompromised children. The widely accepted treatment for mucormycosis is amphotericin B along with surgical debridement. The rising of mucormycosis or black fungus in COVID-19 paediatric patients can be managed effectively if identified early with adequate treatment with amphotericin B, surgical debridement and controlling of the associated risk factors.

STUDY LIMITATION

This study has a relatively small sample size and may limit the outcome of the above interpretation. However, the outcome of this study will definitely encourage the future research work in this fatal clinical entity called COVID-19 associated mucormycosis.

DISCLOSURE

The authors declare no conflict of interest.

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