



Rhinocerebral-Orbital Mucormycosis: Salvageable with Prompt Recognition and Treatment

Authors

Dr Vijay Bhaisare, Dr Amol Chaudhari, Dr Komal Khetwani

Corresponding Author

Dr Amol Chaudhari

18, Muktai Housing Society, Pavan Nagar, Nasik Maharashtra 452010

Email: amolplanet70@gmail.com

ABSTRACT

A case of 10 yrs. old type 1 diabetic male, who presented with left eye orbital cellulitis and lid necrosis. After prompt diagnosis of Rhinocerebral orbital mucormycosis the patient was debulked and was started with i.v amphotericin with renal monitoring along with blood sugar control, which resulted in patient survival. Hence we conclude that inspite of intracranial extension seen in MRI, high suspicion, prompt diagnosis, improvement of general condition and early intervention can be salvageable even in poor prognosis mucormycosis cases.

Keywords: *Mucormycosis, Rhizopus, Diabetic ketoacidosis.*

Introduction

Mucormycosis (also known as phycomycosis and zygomycosis) is an aggressive opportunistic fungal infection.

Although mucormycosis can affect other parts of the body such as lungs and gastrointestinal tract, this case study focuses on the rhinocerebral-orbital type.¹ It is common in especially diabetic patients. Other conditions associated are multiple blood transfusions, haematological malignancies, chronic steroid abuse.^{2,3} Rhinocerebral-orbital mucormycosis, if associated with lid gangrene is considered prognostically poor. It is a life-threatening infection associated with significant mortality.⁴ Mucormycosis is difficult to diagnose early, as patients often present with non-specific symptoms. By the time signs of orbital apex involvement develop, it is often too late to save the patient's vision, or even patient's eye or life.

Pathogenesis

Diabetic or immunocompromised patients presenting with sinus diseases. Organisms invade the Paranasal Sinus Mucosa. Organisms reach into the orbit or brain parenchyma (causing sino-orbital &/or rhinocerebral infections). Invasion of blood vessels leading to tissue infarction & massive necrosis with bone destruction.⁵⁻⁷ Involvement of orbital structures & orbital apex leading to orbital apex syndrome. Further posterior extension leading to cavernous sinus thrombosis & Brain parenchyma involvement.

Case Report

A case of 10 yr. old male, who was presented with Diminution of vision left eye since 2 months which was gradual in onset and progressive in nature. and Swelling over left eyelid with blackish discoloration. Patient is known case of diabetes

mellitus Type 1 since 2 yrs. On examination of right eye no abnormality detected. Left eye vision was light perception Negative. Lids swollen with black necrotic eschar having oedematous margin .Conjunctiva congested & chemosed Cornea hazy & oedematous. Ocular movements were restricted in all gazes. Digital tension was decreased .Rest details were not appreciated. (Figure.1)on oral examination, Tongue necrosis was present. (Figure.2) Nasal Cavity was obstructed. After history and examination of the patient we suspected it as a case of mucormycosis and we get MRI brain and orbit done. At the same time we started treatment.

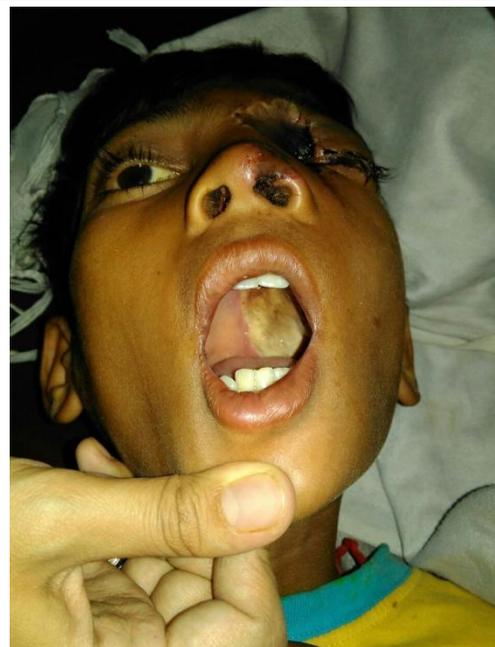


Figure 2.

Investigations

MRI shows B/L orbital cellulitis, more significant on left side with mild left proptosis. The abscess is seen in the left parasellar region with involvement of the cavernous sinus and orbital apex. The intracranial meningeal thickening and enhancement is seen in the bilateral paramedian basifrontal regions suggestive of meningitis. (Fig 3) Findings are suggestive of fungal etiology.



Figure1.

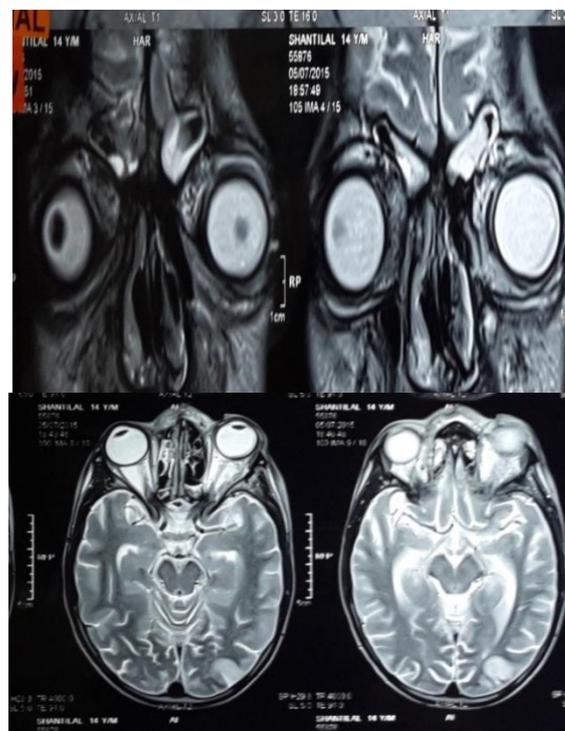


Figure 3

Treatment

After prompt diagnosis the patient was started with i.v liposomal amphotericin 4mg/kg i.v per day for 11 days with renal monitoring along with blood sugar control. The patient underwent orbital exenteration and radical debridement of involved adjacent structures. Almost all of the necrotic and infected tissue was excised and specimen was sent for histopathological examination. The surgical

defect was packed with amphotericin-soaked Surgical. Hence the combination of exenteration and antifungal therapy saved his life. Histopathological examination of scrapings of involved tissue were diagnostic .Broad, irregular, nonseptate hyphae with its sporangial capsule suggests Rhizopus. (Figure 4)

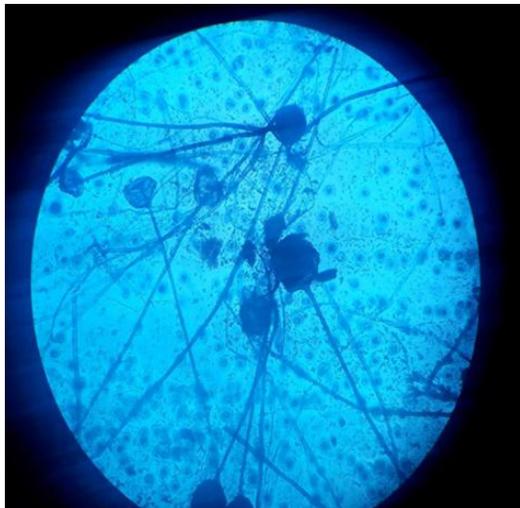


Figure 4

Discussion

Rhinocerebral mucormycosis still remains a poorly understood disease with high mortality rate. Presently, the triad of clinician's awareness, prompt initiation of treatment and timely surgical intervention represent the effective way of managing the disease.⁸ It is an acute opportunistic mycosis that predominantly occurs in the patients with diabetes. The clinic physician may see patients with Rhinocerebral mucormycosis in its earliest stages masquerading as other less serious diseases. Blitzer and Lawson found that in their review of 170 cases of Rhinocerebral mucormycosis, 63% of untreated diabetics died as compared with 17% mortality rate when therapy included aggressive surgery and amphotericin B administration. Absences of intracranial or orbital extension are indicators of good prognosis. Diabetic ketoacidosis at presentation is the single most important detrimental factor. The initial

clinical picture might cloud the aggressive nature of the disease.⁹

Conclusion

This case is noteworthy as the patient came to us in poor condition but with high suspicion, prompt diagnosis, improvement of general condition, blood sugar control and early intervention is salvageable in this poor prognosis mucormycosis case and we could discharge the patient. Hence we conclude that rhinocerebral-orbital mucormycosis should be considered in all the patients with orbital inflammation associated with retinal or orbital infarction, regardless of their immunologic status because prompt surgical and medical intervention is critical and is essential for cure.

References

1. Ribes JA et al. Clin Microbial Rev. 2000; 13(2):236-301.
2. Kasapoglu F et al. Otolaryngol Head Neck Surg. 2010; 143(5):614-620.
3. Roden MM et al. Clin Infect Dis. 2005; 41(5):634-653
4. Abril V, Ortega E, Segarra P, et al. Rhinocerebral mucormycosis in a patient with AIDS: a complication of diabetic ketoacidosis following pentamidine therapy. Clin Infect Dis 1996; 23:845-846.
5. Greenberg RN et al. Curr Opin Infect Dis. 2004; 17(6):517-525.
6. Yohai RA et al. Surv Ophthalmol. 1994; 39(1):3-22.
7. Gamaletsou MN et al. Curr Infect Dis Rep. 2012; 14(4):423-434.
8. Bray WH, Giangiacomo J, Ide CH. Orbital apex syndrome. Surv Ophthalmol 1987; 32:136-40.
9. Lehrer RI, Howard DH, Sypherd PS, et al. Mucormycosis. Ann Intern Med 1980; 33:93-1.