Rhino facial zygomycosis: case report
Zigomicose rino facial: relato de caso

Juliana Miguita e Souza¹, Antonio José Sproesser Junior¹, Alexandre Felippu Neto¹, Florencia Barbero Fuks¹, Carlos Augusto Cardim de Oliveira¹

ABSTRACT
Zygomycosis is an invasive disease that affects both immunocompetent and immunocompromised, depending on the type of strain. This disease diagnosis is clinical and histopathological, and its treatment is based on antifungal therapy and surgical cleaning. This paper reports a case of a boy with invasive zygomycosis rinofacial who final treatment was successful after underwent antifungal and surgical therapies.

Keywords: Zygomycosis/therapy; Zygomycosis/diagnosis; Magnetic resonance imaging; Tomography, X-ray computed; Case reports

RESUMO
A zigomicose é uma doença invasiva, que acomete tanto imunocompetentes como imunocomprometidos, dependendo do tipo da cepa. O diagnóstico é clínico e histopatológico, e o tratamento é baseado em antifúngico e em limpeza cirúrgica. O presente relato de caso é sobre um menino com zigomicose rinofacial invasiva com tratamento final bem-sucedido, após terapias antifúngicas e limpezas cirúrgicas.

Descritores: Zigomicose/terapia; Zigomicose/diagnóstico; Imagem por ressonância magnética; Tomografia computadorizada por raios-x; Relatos de casos

INTRODUCTION
Zygomycosis is a rare fungal, severe, invasive and lethal disease with rapid progress. This disease is prominent in countries with warm and humid climate.¹,² Its incidence in Brazil is unknown because of the scarcity of reports in the literature, but it is well known that several cases occur in north and northeast region of the country.³ There is a tendency in international literature, mainly in immunocompromised patients, of increase in the incidence of fungal invasive disease. An incidence of zygomycosis of 2-4% was found in autopsies of cancer patients both in Japan and the United States.³

The zygomycosis has several clinical presentations (cutaneous, pulmonary, systemic and gastrointestinal), however, the rhino-orbital-cerebral is the most common.⁴,⁵

Zygomycosis etiologic agents originate from Mucorales that primarily affects immunocompromised patients and base disease carriers, and from the Entomophthorales that is commonly found in immunocompetent individuals and rarely causes angioinvasive disease.¹,⁵

The diagnosis is based on a combination of mycologic and histopathological tests, and clinical presentation.⁶ The fungal infection could be determined by direct examination with potassium hydroxide (KOH) and culture on Sabouraud’s medium. The histopathological analysis shows nonseptate hyphae with branches at 90° angle, which usually does not invade tissues and blood vessels.⁷

Common symptoms include nasal congestion, nasal discharge and chronic sinusitis,⁸ however, the infection may also present fever, lethargy, headache, retro-orbital pain, sudden vision loss, proptosis, periorbital cellulitis, epistaxis and seizure.⁹

At physical examination necrotic crusts can appear on the nasal septum, turbinates and palate. Initially, the orbital involvement can be show by proptosis and periorbital cellulitis with subsequent ophthalmpolegy and amaurosis.⁶ Later, the spread of the infection to the central nervous system can occur by the ethmoid bone and sphenoid sinus after bone destruction.

¹Hospital Israelita Albert Einstein, São Paulo, SP, Brazil.
Corresponding author: Juliana Miguita e Souza – Hospital Israelita Albert Einstein, Avenida Albert Einstein, 627/701 – Jardim Leonor – Zip code: 06652-900 – São Paulo, SP, Brazil – Phone: (55 11) 2151-1233
E-mail: msjuliana@uol.com.br
Received on: Feb 26, 2013 – Accepted on: Dec 11, 2013
DOI: 10.1590/S1679-45082014RC2579
Treatment success depends on rapid diagnosis and the combined approach of surgical procedures and antifungal therapy. The amphotericin B is the drug of choice for initial treatment. There are studies reporting increase of up to 79% in patient survival after adoption of this drug. For the effective control of the disease, high doses of amphotericin B must be used. These doses range from 0.8 to 1.5mg/kg daily, which correspond to doses very close to nephrotoxic levels. 

The use of hematopoietic as stimulating factors and hyperbaric oxygen therapy may be beneficial, but few studies had established the efficacy of these procedures. There are also studies reporting success by the oral administration of potassium iodide. Surgical debridement, which main goal is to remove as much devitalized tissue as possible, in addition to establish adequate sinus drainage, has a significant impact on the disease morbidity and mortality.

Prevention of the disease consists in environmental control, avoidance or reduction of direct contact with fungal propagules, which are found in plants, flowers and house dust.

Poor prognosis factors are delayed initiation of treatment, intracranial (hemiplegia and hemiparesis), palate or orbital involvement as well as bilateral facial sinus involvement, and facial necrosis.

This paper reports a case of immunocompetent patient with late diagnosis and who initially had unfavorable clinical evolution. Treatments were based on the experience of the responsible physician and case reports previously published in the literature. The result was total remission of the disease and no sequelae.

CASE REPORT

A 9-year-old boy from city of Belém (PA) was referred to the pediatric unit of the Hospital Israelita Albert Einstein (HIAE) in São Paulo, for treatment of rhino facial zygomycosis. He was a previously healthy and had a habit of smelling roses. In March 2010, the patient presented tumors inside his left nostril that increased progressively. The child was diagnosed with cellulitis and treated with antibiotics, topic and systemic corticoids, however, the lesions did not improve but got worse. The computed tomography of the skull revealed sinusopathy of left paranasal sinuses (Figure 1). Almost 3 months later, the patient was submitted to surgical cleaning of facial sinus along with resection of lesion on nasal wall. The histopathological exam showed zygomycosis. After two weeks, he become to present painful cellulitis in orbital region (Figure 2). Treatment was initiated with amphotericin B and caspofungin. Despite the introduction of medication, the disease continued to progress and the pain worsened, being also observed a decrease in ocular motility and proptosis. The magnetic resonance test showed periorbital invasion with probable contiguity to the skull base (Figure 3). With this clinical picture, the child was submitted again to two surgical cleanings, and the antifungal treatment was extended. Subsequently, the patient condition improved and lesions and periorbital edema reduced. The Conidiobollus sp was isolated from paranasal secretions culture.

After stabilization of patient’s condition, he returned to Belém to complete the treatment with amphotericin B. The appearance of side effects such as hypocalcemia, hyperthermia and body pain caused his hospitalization at intensive care unit for hemodynamic and hydroelectrolytic stability. After his discharge from intensive care unit, the treatment was replaced by oral antifungal posaconazole that caused a worsening in clinical picture.
The patient returned to our service in August 2010 and a new magnetic resonance exam of the skull showed progression of infiltrative lesions affecting the left hemiface, orbits and large extension of ethmoid and sphenoid sinus, besides affect the pachymeningeal (Figure 4). Because of clinical picture worsening, persistence fever and complications related to hospitalization a therapeutic scheme was used with amphotericin B, caspofungin, terbinafine, itraconazole, corticoids, acyclovir, erythropoietin, ceftriaxone, cefepime and ciprofloxacin at different times and also hyperbaric oxygenotherapy. Fifteen days after the last session of hyperbaric chamber, a magnetic resonance showed lesions but without signs of invasion of healthy tissues (Figure 5). After almost 2 months, the patients underwent another surgical cleaning and at that time there were improvement on obstructive symptoms and practically, the cure of the disease.

DISCUSSION

The Conidiobollus sp is a saprophytic organism that grows on the floor and propagules in vegetation. This parasite often affect arthropods; human infection is accidental. Fungal infections can be acquired by inhalation or direct inoculation, being this latter the case of our patient.

The most common presentation of the disease is an initial clinical picture with nasal lesion of rapid growing that provoke obstructive signs that, within few weeks, spread to periorbital region and cause edema, ptosis and proptosis.

The late diagnosis, about 3 months, justifies the high morbidity of this disease and it can be explained by several types of differential diagnosis for the initial symptoms. In our case, the skin lesion was first treated as impetigo and other obstructive symptoms such as sinusitis. The lack of therapeutic response resulted in surgical cleaning and biopsy.

It is important to mention that intravenous amphotericin B has been considered the treatment of choice. In our patient, this fact was well registered because in the period that he received this medication by the adequate route the response was more satisfactory. However for a completely adequate treatment it is necessary to be careful with drug administration in order to avoid side effects, which were also observed in the child in our report.

The hyperbaric oxygenotherapy might contributed to the improvement, although there is no evidences of its use for treatment of zygomycosis. The surgical cleaning is an essential part of the therapeutic strategy, because it removes as much infected material as possible and enables to observe the sinus drainage, which was observed in our case. For this reason, the use of surgical cleaning should be emphasized.
CONCLUSION
The zygomycosis is a severe fungal disease that can affect both immunocompetent and immunocompromised individuals.

The literature shows that rapid diagnosis and early adequate treatment are fundamental for the disease prognosis. The patient described in this case, although with little delay in the diagnosis, had a favorable progress due to treatments used.

REFERENCES
1. Prabhu RM, Patel R. Mucormycosis and entomophthoramycosis: a review of the clinical manifestations, diagnosis and treatment. Clin Microbiol Infect. 2004;10 Suppl 1:31-47. Review.
2. Hoogendijk CF, van Heerden WF, Pretorius E, Vismer HF, Jacobs JF. Rhino-orbitocerebral entomophthoramycosis. Int J Oral Maxillofac Surg. 2006;35(3):277-80.
3. Tadano T, Pain NF, Hueb M, Fontes CJ. [Entomophthoramycosis (zygomycosis) caused by Conidiobolus coronatus in Mato Grosso (Brazil): case report]. Rev Soc Bras Med Trop. 2005;38(2):188-90. Portuguese.
4. Mohanty D, Dhar M, Dwivedi S. Mucormycosis. Trop Doct. 2010;40(2):127-8.
5. Marques SA, Camargo RM, Abbade LP, Marques ME. Mucormicose: infecção oportunística grave em paciente imunossuprimido. Relato de caso. Diagn Tratamento. 2010;15(2):64-8.
6. Severo CB, Giazzelli LS, Severo LC. Zigomicose Curso de Atualização-micoses. J Bras Pneumol. 2010;36(1):134-41. Review.
7. Dannasui E, Meletiadis J, Mouton JW, Meis JF, Verweij PE; Eurofung Network. In vitro susceptibilities of zygomycetes to conventional and new antifungals. J Antimicrob Chemother. 2003;51(1):45-52.
8. Michael RC, Michael JS, Mathews MS, Rupa V. Unusual presentation of entomophthoramycosis. Indian J Med Microbiol. 2009;27(2):156-8.
9. Xavier SD, Korn GP, Granato L. Mucormicose rinocerebral: apresentação de caso com sobrevida e revisão de literatura. Rev Bras Otorrinolaringol. 2004;70(5):710-14.
10. Lithander J, Louon E, Worthing E, Ganesh A, Al-Lawatia YM, Elamin A, et al. Orbital entomophthoramycosis in an infant: recovery following surgical debridement, combination antifungal therapy and use of hyperbaric oxygen. Br J Ophthalmol. 2001;85(3):374-5.