Case report of subcutaneous entomophthoromycosis with retroperitoneal invasion

Relato de caso de entomofthoromicose subcutânea com invasão retroperitoneal

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ABSTRACT
The authors describe a case of entomophthoromycosis in a previously healthy patient, who presented with an abscess in the right buttock. After surgical drainage it evolved into a retroperitoneal tumor. The patient improved clinically after resection of the mass and ketoconazole treatment. The histopathological analysis showed the Splendore-Hoeppli phenomenon, suggesting Basidiobolus ranarum infection, a zygomycosis generally restricted to the subcutaneous tissue, with rare gastrointestinal involvement.


RESUMO
Os autores descrevem um caso de entomofthoromicose em paciente previamente saudável, que apresentou abscesso em nádega direita, evoluindo, após drenagem cirúrgica, para tumoração retroperitoneal. Após ressecção da massa, o paciente obteve melhora clínica, em uso de cetoconazol. A análise histopatológica evidenciou fenômeno de Splendore-Hoeppli, sugerindo infecção por Basidiobolus ranarum, uma zigomicose geralmente restrita ao tecido subcutâneo, com raro comprometimento gastrointestinal.


We refer to human zygomycosis as polymorphic diseases caused by fungi of the Zygomycetes class. This class is subdivided into two orders: Mucorales and Entomophthorales. Unlike the Mucorales order, which has the ability to cause systemic disease, this last order is mainly represented by Conidiobolus coronatus, the etiologic agent of subcutaneous infections confined to the nasal mucosa and adjacent tissues, and Basidiobolus ranarum (also known as Basidiobolus haptosporus), which involves the trunk and limbs, and rarely the gastrointestinal tract.

We report a rare form of entomophthoromycosis suggesting Basidiobolus ranarum infection, which extended to the retroperitoneum, after subcutaneous infection in the right buttock.

CASE REPORT

A 43-year-old male farmer from the rural area of Unai, in State of Minas Gerais, developed an abscess in the right buttock, without previous local trauma, and was admitted to the Hospital of Sobradinho, Brasília, Brazil, in July 1989. The patient’s medical history was unremarkable and he was previously healthy. No
systemic illness was observed. Roentgenographs of the pelvis showed a soft-tissue abscess without bone or joint lesions. The abscess was surgically drained and the lesion improved without specific medication. The contents of the drainage were not sent for culture. Histopathologic analysis of a subcutaneous tissue sample showed nonspecific chronic inflammation.

In February 1990, seven months after discharge, the patient started to complain of arthralgia in the right knee, abdominal pain, malaise, uncharacteristic fever, vomiting and weight loss. He was admitted to the same hospital, where a diagnostic procedure was performed.

On physical examination, the patient had a poor nutritional state, weighing only 45kg, with normal vital signs. No lymphadenopathy was found. The lungs and heart and neurologic examination were normal. His abdomen was tender, particularly in the right flank, where a palpable, nodular mass was located. The liver extended 5cm below the right costal border.

Hematologic laboratory data obtained on admission were as follows: hematocrit 0.41 and white-cell count 10.8 x 10^9/L, with a normal differential count. The blood chemistry values were normal, as well as the coagulation profile and urinalysis. The tuberculin skin test was negative at 48 hours. Ultrasound of the abdomen confirmed the presence of a thick mass in the right flank (405cm^3).

He subsequently underwent an exploratory laparotomy, where a mass was found in the retroperitoneum, upon the right kidney. It was excised and sent for histopathological analysis. The resected material consisted of an irregular mass, which was whitish, smooth, firm and measured approximately 10cm in the greatest diameter. Microscopically, hematoxylin and eosin (H&E)-stained sections showed a dense eosinophilic infiltrate. Many noncaseating granulomas were present (Figure 1) and multiple areas demonstrating Splendore-Hoeppli phenomenon were observed, with a thick, eosinophilic, hyalinized sheath surrounding the thin-walled fungal hyphae (Figures 2 and 3). No vascular invasion or fungal thrombosis were seen. A culture of the specimen was not performed. Based on the localization of lesions, the agent aspect in the tissues, and histopathologic picture, Basidiobolus ranarum was suggested.

The patient received ketoconazole (400mg/d) for 35 days with a complete clinical improvement.

**DISCUSSION**

Mucormycosis (infection with fungi of the order Mucorales) is characterized by a rapidly progressive and usually fatal course, primarily in immunocompromised patients. Common clinical manifestations include rhinocerebral, pulmonary gastrointestinal and cutaneous involvement as well as dissemination. Due to its vasculotropism, the hyphal infiltrations found near blood vessels and septic thromboembolism are common. The infection. In 1988, of 33 Latin American cases reported, 84% were described in Brazil. The tissue reaction is almost pathognomonic, consisting of a chronic granulomatous reaction with thin-walled, often septate hyphae or hyphal fragments (4-10μm in diameter) with a thick eosinophilic sheath (Splendore-Hoeppli phenomenon) staining intensely pink in H&E sections and bright red in PAS-stained sections.
The infection due to *B. ranarum* is widely prevalent in tropical and subtropical regions and is often misdiagnosed or not reported. Infection usually begins as small papules following a scratch or puncture by an insect, plant, or other fungus-bearing object. The lesion commonly develops in the perineum, buttocks, trunk and thighs, probably due to the habit of using decaying plant leaves lying on the ground as toilet paper after defecating in the open⁷. Since the fungus cannot withstand refrigeration, biopsied material should be cultured immediately after collection. This probably contributes to the absence of the definitive diagnosis in most of the cases where there is no previous suspicion of fungal infection⁷ ¹⁰.

Visceral disease caused by *B. ranarum* is rare, but cases of gastrointestinal infection have been reported worldwide⁵ ⁸ ¹³ ¹⁵. It is not clear how the fungus gains access to the gastrointestinal tract of patients. It is attributed to ingestion, with the portal of entry being the stomach, or to direct inoculation via minor rectal trauma.

Possibly due to protease enzymes produced by *B. ranarum* and other Entomophthorales¹², the fungus is able to invade soft tissues and eventually evolve beyond a subcutaneous infection. One patient with a voluminous lesion resembling a soft tissue tumor has been reported elsewhere². Two patients similar to ours have been reported, one with retroperitoneal lesions and one with pelvic cavity lesions. A surgical procedure prior to the clinical detection of the internal masses was observed in both¹¹ ¹⁴. We consider that the rare forms of intra-abdominal entomophthoromycosis without involvement of the gastrointestinal tract are probably related to invasive procedures, which possibly implant the fungus more deeply.

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REFERENCES