Intestinal Histoplasmosis with *Histoplasma duboisii* in a Patient Infected by HIV-1 in Abidjan (Ivory Coast)

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Abstract

We report a case of intestinal histoplasmosis with *Histoplasma duboisii* in a 39-year-old patient infected with HIV-1, admitted to the emergency department due to peritonitis with fever and weight loss. He underwent a right hemicolectomy, and the pre-perforation and ulcerated macroscopic aspect during surgery suggested a malignant tumor. The anatomopathological examination of the specimen removed revealed the presence of inflammatory granulomas containing *Histoplasma duboisii* yeasts. The evolution was rapidly unfavorable, culminating in the patient’s death 12 days after treatment onset with intravenous amphotericin B associated with an antiseptic bi-antibiotherapy. The authors point out the rarity of intestinal localization of histoplasmosis in patients infected with HIV in Abidjan and the contribution of anatomopathological examination in the diagnosis of this condition.

Keywords: Abidjan; Amphotericin B; Intestinal histoplasmosis; HIV-AIDS; West Africa

Introduction

PAfrican histoplasmosis, or histoplasmosis with *Histoplasma duboisii*, is a profound and rare mycosis, mostly found in West and Central Africa. It rarely occurs in disseminated form because of its particular tropism for the skin, the lymph nodes and the skeleton, and is rarely associated with HIV [1]. In the Ivory Coast, this condition is rarely described in patients infected with HIV [2]. However, cases of cutaneous and bone localizations of histoplasmosis with *Histoplasma duboisii* were been recently reported [3-5]. The clinical presentation of histoplasmosis in an HIV patient that we describe in this case has never been reported before in the Ivory Coast.

Observation

Mr. K.B., age 39, and an Ivory Coast native living continuously in Abidjan, was admitted to the Sainte Anne Marie International Polyclinic of Abidjan on March 10, 2007, for severe abdominal pain and emesis lasting for five days. His temperature was 39.6°C, his blood pressure at 110/60 mmHg, his pulse at 104 beats/min, and his general condition was altered, associated with conjunctival pallor and dehydration. He presented with diffuse abdominal contracture, evoking peritonitis. The pleuropulmonary, cardiovascular, musculoskeletal and mucocutaneous systems did not reveal any particularity. The lung radiography showed a minimal bilateral pleural effusion, and the unprepared abdomen radiography showed some fluid level in the periphery. Moreover, the patient presented with normochromic normocytic anemia (hemoglobin at 10.8g/dl), bicytopenia (leukocytes at 2,200/mm3, red blood cells at 2,840,000/mm3), and thrombocytosis at 502,000/mm3. The other biological analyses, including hematuria, creatinine, natrema, kalemia, transaminases, glycemia, hemocultures, Widal and Félix serodiagnosis, were normal. An emergency right hemicolectomy was performed. The specimen removed showed an ulcerated and pre-perforation aspect. Colonic continuity was restored during the same operation. Histopathological examination of the specimen revealed an inflammatory granuloma consisting of multinucleated macrophages containing yeast and surrounded by a clear halo, evoking the *duboisii* variety of Histoplasma. It presented lymphoplasmocytes and polynuclear eosinophils, without histological evidence of malignancy [Figure 1]. The patient was infected with HIV-1 with a CD4 count of 49/mm3. The serology of histoplasmosis was not performed. Treatment included a bi-antibiotherapy (ceftriaxone + metronidazole) used as an antiseptic and intravenous amphotericin B (at 1 mg/kg every other day). The evolution became rapidly unfavorable, with exitus of the patient on day 12 due to multiple organ failure.

Figure 1: Inflammatory granuloma consisting of lymphoplasmocytes and macrophages containing numerous Histoplasma with a clear halo (hematoxylin-eosin staining, ×400).

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Received March 06, 2011; Accepted September 20, 2011; Published September 25, 2011


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Discussion

In the Ivory Coast, histoplasmosis with *Histoplasma duboisii* has been rarely described in patients with AIDS [3-5]. This confirms the idea that *Histoplasma duboisii*, which is mainly found in Africa, is rarely associated with HIV, while *Histoplasma capsulatum*, which, in its disseminated form, is a definition criterion for AIDS, is common in America [1]. Its rarity in our country is due to many reasons: its clinical polymorphism, which may mislead the clinical diagnosis, as in many observations described in Ivory Coast [3-5]; but also to the unavailability and/or non-systematic use of anatopathological examination and serological tests. Indeed, the gastrointestinal manifestations (abdominal pain, diarrhea and sometimes blood emesis) are not specific for intestinal histoplasmosis, and can be found in 3% to 12% of patients with the disseminated form [6]. Sometimes, it is possible to observe gastrointestinal bleeding, intestinal perforation or even peritonitis, as in our case. For this reasons, Loulergue et al. already suggested that diagnostic of histoplasmosis in African patients should not be discounted because of the HIV status of the patients based on their cases and previous literature review [7]. The reports of many authors support the idea that the low incidence of African histoplasmosis could be due to a problem of under-reporting or under-recognition in Africa [7-9]. In resource limited situation, empirical anti-fungal treatment could be considered in patients with suspected gastro-intestinal histoplasmosis in order to diagnose and to treat this potentially life-threatening infection. The histopathological appearance of histoplasmosis is an inflammatory granuloma rich in epithelial and giant cells circumscribed by lymphplasmonocytes. At the center of the granuloma, there is fibrinohemorrhagic necrosis, or necrosis caseosa, which in our tropical context can guide the diagnosis to a tuberculose lesion [6]. On a morphological level, *Histoplasma* may be confused with *Candida glabrata; Penicillium marneffei*, *Cryptococcus neoformans*, and the amastigote form of trypanosomes, hence the interest in the research of a blood or urine antigen for *Histoplasma* to facilitate definitive diagnosis. Unfortunately, these diagnostic tests are not available yet in limited resources setting. Many authors have reported cases of disseminated histoplasmosis under antiretroviral therapy related to immune restoration [10-14]. None of these cases have been still reported in Africa. Perhaps, we could then fear a resurgence of this disease in Africa as a result of the current generalization of antiretroviral therapy. According to international recommendations, Amphotericin B is the preferred treatment for disseminated and intestinal histoplasmosis [6] and Itraconazole, for the consolidation phase; but in our context, Itraconazole is not always available. These treatments should improve the prognosis of the disease, whose lethality can in some cases reach 60% [9,15,16].

Conclusion

Intestinal histoplasmosis with *Histoplasma duboisii* is rarely associated with AIDS in the Ivory Coast. This rarity can be due to under-reporting or under-diagnostic, and diagnostic difficulties because of the lack of specific clinical sign, and the unavailability of sensitive and specific diagnostic test. In this context, empirical antifungal treatment should be founded in front of evocative clinical signs of intestinal histoplasmosis.

References