Pulmonary cryptococcosis in a patient with ulcerative colitis treated with prednisolone and azathioprine

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INTRODUCTION

Ulcerative colitis (UC) is a chronic disease characterized by diffuse inflammation of the colonic mucosa that affects the rectum and proximally extends in a variable, symmetrical and circumferential manner throughout the colon. Its cardinal symptom is bloody diarrhea associated with fecal urgency and rectal tenesmus, and its clinical course is intermittent, with exacerbations and remissions (1). Initial treatment is usually pharmacological, and surgery may be performed in only a few cases. The use of some drugs, such as corticosteroids, immunomodulators and antitumor necrosis factor (TNF) causes immunosuppression, which in some cases may be associated with opportunistic infections (2).

The following is the case of a male patient diagnosed with UC, who received escalation therapy with salicylates, immunomodulators and biological therapy, without achieving adequate control of the disease. As a result, the patient was taken to surgery and had a successful postoperative recovery. However, during treatment, he presented with pulmonary cryptococcosis that was studied clinically, radiologically, and histologically, and appropriate treatment was provided, achieving a satisfactory response.

Abstract

Inflammatory bowel disease comprises Crohn’s disease (CD) and ulcerative colitis (UC), the latter being a chronic disease characterized by diffuse inflammation of the colonic mucosa that affects the rectum and extends proximally. Its clinical course is intermittent with exacerbations and remissions and its treatment is generally pharmacological, with steroids, immunomodulators, and anti-tumor necrosis factor inhibitors (TNF), which cause the patient to be in a state of immunosuppression associated, in some cases, with opportunistic infections. The literature describes the occurrence of pulmonary cryptococcosis in patients with human immunodeficiency virus (HIV) infection, in cases associated with drug treatment of patients with CD, as well as with other opportunistic infections such as tuberculosis and herpes. This is one of the first cases of pulmonary cryptococcosis reported in a patient diagnosed with ulcerative colitis, who received step therapy with salicylates, immunomodulators, and biological therapy. The infection was documented clinically, radiologically, and histologically. The patient received the appropriate treatment and had a satisfactory evolution.

Keywords

Inflammatory bowel disease, Ulcerative colitis, Immunosuppression, Opportunistic infections, Pulmonary cryptococcosis.
CLINICAL CASE

A 48-year-old male consulted in November 2011 due to an 8-month history of diarrhea with mucus and blood as well as a 10-kg weight loss; no other relevant symptoms were informed by the patient. On physical examination, mild tenderness over the colonic area was reported. A colonoscopy was performed, finding extensive colitis, and biopsies, which later confirmed the diagnosis of UC, were performed; thus, treatment with mesalazine 3 g/P.O./day and a corticosteroid cycle (prednisolone) of 40 mg/P.O./day was started, achieving clinical improvement. The patient was discharged with a permanent prescription of mesalazine, as well as corticosteroids use, with a gradual decrease in their dose until suspension.

Three months later, and after an early and voluntary suspension of both drugs, the patient had a relapse. A colonoscopy revealed ulcerative pancolitis with a Mayo severity score of 2 and E3 S2 severity degree according to the Montreal classification. Mesalazine was restarted at 3 g/P.O./day, achieving symptoms resolution.

However, in October 2013, the patient had a new relapse. A stool test showed amebiasis, which was treated with metronidazole plus nifuroxazide every 12 hours, for 7 days, without any improvement. Therefore, considering the reactivation of the disease, azathioprine 150 mg/P.O./day (2.5 mg/kg/day) was added to the prescription of mesalazine, achieving adequate remission.

The patient consulted again in November 2014 due to bloody diarrhea, malaise, asthenia and adynamia, and weight loss of 4 kg. He was taking mesalazine 3 g/P.O./day and azathioprine 50 mg/P.O./day (2.5 mg/kg/day). The patient was diagnosed with anemic heart disease, lower gastrointestinal tract bleeding and active steroid-dependent UC, with a Montreal classification S3 severity. Corticosteroid pulse treatment was initiated (prednisolone 40 mg/day). Moreover, 2 units of packed red blood cells and iron sucrose were transfused.

During an endoscopic check-up, ulcerative pancolitis and pseudopolyps were found, which, compared with the previous colonoscopy, showed a progression of the disease, thus the initiation of biological therapy was considered. Tests for tuberculin (purified protein derivative, PPD), hepatitis A, B, and C, and HIV were performed. A chest x-ray was also done, revealing a nodule in the left upper lobe and a nodule in the lower lobe, 6 mm each (Figure 1). A computed tomography of the chest confirmed these findings (Figure 2).

Suspecting tuberculous granuloma, a wedge biopsy of the lower lobe nodule was performed. The pathology report showed lung parenchyma with chronic caseating granulomatous disease, compatible with cryptococcosis, and negative for acid-fast bacilli (AFB) (Figure 3). Fungal cultures confirmed a moderate growth of Cryptococcus neoformans, while negative PPD allowed ruling out latent tuberculosis.

Considering the diagnosis of pulmonary cryptococcosis without systemic involvement, the infectious disease service started treatment with fluconazole at 400 mg/P.O./day and contraindicated biologic therapy for at least the
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Thus, total proctocolectomy, J-pouch surgery, ileoanal anastomosis, and protective ileostomy were performed. The ileostomy was closed during a second operation in February 2016. Currently, the patient is asymptomatic, has a good quality of life and does not require medication.

DISCUSSION

In recent decades, the incidence of opportunistic infections, such as tuberculosis and cryptococcosis, has increased especially due to the emergence of diseases such as acquired immunodeficiency syndrome (AIDS), where cryptococcosis can occur in 5 to 10% of cases. These opportunistic infections have also been reported in patients undergoing immunosuppression following organ transplantation, in those with inflammatory diseases, and even

Figure 2. A pulmonary nodule with thick calcifications of 6 mm in diameter was observed in the left upper lobe apicoposterior segment, as well as another subpleural nodular image of 6 mm in diameter in the left lower lobe apical segment with soft tissue density. Source: Own elaboration.

Figure 3. Using hematoxylin-eosin stain, a necrotizing granuloma with debris and histiocytes was observed in the lung parenchyma. Higher magnification (10 X) made it possible to appreciate in detail the cryptococcus spores, indicated with arrows. Source: Own elaboration.

first eight weeks of fluconazole treatment. Furthermore, a monthly serum cryptolatex test was indicated during the first year, obtaining negative results.

Two months after starting treatment with fluconazole, adalimumab was added to the mesalazine and azathioprine treatment scheme. Initially, the patient had an adequate response, but after eight months, he was readmitted due to a new relapse. Reactivation of UC was then considered based on the disease progression (S3 type according to the Montreal severity scale) and the presence of severe colitis as measured with the Truelove and Witts severity index.

However, given the inadequate response to adalimumab, treatment with infliximab was started, achieving an initial improvement of symptoms. However, he had another relapse in October 2015, which was considered a failed response to biological treatment, so surgical treatment was decided.
in immunocompetent persons. It has been estimated that up to one million new cases of cryptococcosis are diagnosed worldwide, with about 650,000 deaths each year (3, 4).

Pulmonary cryptococcosis is caused by an encapsulated yeast that is acquired after inhaling spores from the environment (5). There are several species, but the most common is *Cryptococcus neoformans*, which can be found anywhere in the world; the *C. gattii* species is found in tropical and subtropical regions (6).

In most cases, it is possible to establish exposure to pigeon, poultry, dog and cat droppings, or other environmental spores. However, the source of infection is not identified in up to 15% of cases (7). In our patient, exposure was not determined.

Patients may experience mild symptoms, such as headaches, coughs, and fevers, or systemic involvement manifested by chills, hemoptysis, and weight loss. Although the percentage may vary, according to several publications, individuals may be asymptomatic in 17-35% of cases. Diagnosis is often made incidentally through imaging findings of pulmonary nodules, which can be solitary or multiple, as well as areas of consolidation or reticulonodular pattern (8, 9).

Our patient had no cryptococcosis-related symptoms. Therefore, diagnosis was incidental and based on the initial finding of pulmonary nodules, which were later confirmed through CT scan. This technique is one of the available diagnostic methods, and it usually shows few or multiple subpleural nodules or masses with or without halo. In addition, in a smaller number of cases, consolidation, pleural effusions, or an increase in the size of mediastinal nodes may be observed (10, 11).

Other diagnostic techniques, such as the serum measurement of capsular antigen, are also used. This technique has the advantage of being non-invasive, although false negative and positive results have been reported, so it is necessary to be cautious when using it (7). Currently, molecular biology tests are also used, which have the advantage of being fast but take more time for proper assessment.

The diagnostic tools considered to be the most specific are culture and tissue biopsy; the latter is the most important and can be collected through puncture or surgically; also, Chinese ink, Gomory, periodic acid-Schiff (PAS) and mucicarmine are the stains used (12). In the case of our patient, culture and biopsy were performed, both confirming the presence of the cryptococcal spores.

Treatment of cryptococcosis in immunocompromised patients with fluconazole was introduced in the 1990s, achieving a marked decrease in the presence of this disease (13). Currently, in mild or moderate lung infections, treatment with fluconazole (400 mg/day) is prescribed for 6-12 months, as it was the case of our patient.

On the other hand, in severe infections or cases when the central nervous system is involved, induction therapy with liposomal amphotericin B (3-4 mg/kg/day IV) or amphotericin B lipid complex (5 mg/kg/day IV) is used for 2 weeks, followed by the administration of fluconazole (400 mg/day) for 8 weeks (14, 15).

Recently, resistance to fluconazole has been observed, especially in eastern countries. According to various reports, this resistance increased from 5.1% to 22.6% over a period of 7 years due to the excessive use of fluconazole for prophylaxis in immunocompromised patients or its administration in primary or maintenance treatments (16).

Few cases of patients with inflammatory bowel disease and cryptococcosis have been reported. However, in all those cases, it has been associated with the administration of anti TNFα agents, whereas in most scenarios, it has been associated in patients with celiac disease (17). So far, no cases of cryptococcosis have been reported in people with UC treated with corticosteroids and azathioprine.

**CONCLUSIONS**

To the best of our knowledge, this is the first case reported in the literature of a patient with UC treated with corticosteroids and azathioprine. The patient had pulmonary cryptococcosis, which is an opportunistic disease that particularly affects people with AIDS. However, it has also been observed in transplanted patients and in those who have undergone treatment for inflammatory diseases. Since these patients are considered immunocompromised and cryptococcal infection could be asymptomatic in them, further studies should be carried out prior to starting biologic therapy.

**Conflict of interest**

None declared by the authors.