To the Editor:

Cryptosporidiosis is an infection caused by a protozoan (Cryptosporidium spp.) that infects the epithelial cells of the gastrointestinal tract of humans and animals.(1)

Clinical manifestations depend on the immune status of the patient. In immunocompetent patients, cryptosporidiosis causes episodes of self-limiting diarrhea, especially in children in Latin America. In AIDS patients with advanced immunodeficiency, cryptosporidiosis is the most common parasitic cause of prolonged diarrhea, associated with pronounced weight loss, and can progress to severe dehydration and electrolyte disturbance.(1,2)

In the pre-antiretroviral therapy (ART) era, intestinal cryptosporidiosis was responsible for diarrheal symptoms in 10-30% of AIDS patients in developed countries and in 30-50% of those in developing countries.(3) Currently, in developed countries in which the rates of environmental contamination are low and in which potent ART is widely available, intestinal cryptosporidiosis occurs at an incidence rate of < 1 per 100 person-years among AIDS patients.(4)

Pulmonary impairment is a rare complication of intestinal cryptosporidiosis, having been reported in immunocompromised patients, most of whom have AIDS and severe immunodeficiency.(2) The prevalence of pulmonary cryptosporidiosis, however, might be underestimated, given that it is not systematically investigated.(10) In a retrospective study conducted in Spain and investigating 275 AIDS patients, 43 had enteritis caused by Cryptosporidium spp.; in 7 of those 43 patients, sputum examination identified Cryptosporidium spp. oocysts.(9)

The pathogenesis of pulmonary cryptosporidiosis has yet to be fully clarified.(2,6) The possibility that pulmonary involvement is due to inhalation of oocysts during an episode of vomiting or results from hematogenous dissemination has been a point of discussion.(2) Although Cryptosporidium oocysts do not usually invade the intestinal mucosa, they have been found within macrophages, leading to reduced phagocytic ability. In addition, this parasite can multiply in macrophages in vitro, which suggests that extraintestinal parasites spread via circulating phagocytes.(10) This hypothesis is supported by the presence of Cryptosporidium spp. within the blood vessels in the intestinal and pulmonary submucosa, as revealed by autopsy studies.(2,7)

The clinical manifestations of pulmonary cryptosporidiosis are nonspecific and typically include chronic cough, fever, and dyspnea as the most common symptoms, and radiological changes may or may not occur.(2) Although there have been no reports of pathognomonic radiological findings in pulmonary cryptosporidiosis, interstitial opacity has been described in some case reports.(2,3,7,8)

In a survey conducted in referral hospitals for the treatment of HIV/AIDS patients with clinically suspected pulmonary tuberculosis in the city of Recife, Brazil, Cryptosporidium spp. was the only agent identified in 2 (1.5%) of the 130 sputum examinations performed during the survey period; diagnosis was based on the finding of Cryptosporidium spp. oocysts on Ziehl-Neelsen-stained sputum smear (Figure 1). In both cases, the patients had evening fever, cough, weight loss, and general malaise, all of which were suggestive of pulmonary tuberculosis. In the 2 cases studied, the patients were severely immunocompromised, with very low CD4+ T-cell counts (25 and 37 cells/µL). Although the presence of Pneumocystis jirovecii was not investigated, sulfamethoxazole-trimethoprim was prescribed for prophylaxis against that agent. Both patients had recently been diagnosed with AIDS. Only one of the patients was initially suspected
of having intestinal cryptosporidiosis (which was confirmed by stool examination for ova and parasites), whereas the other had clinical signs suggestive of pulmonary tuberculosis but no digestive complaints. Chest X-ray findings (Figure 2) were normal in the 2 patients. Sputum culture (on Löwenstein-Jensen medium and in Middlebrook 7H9 medium) yielded no bacterial growth.

The patients showed progressive clinical, immunological, and virological improvement after ART initiation. There is no specific drug with proven efficacy in the treatment of cryptosporidiosis, and clinical improvement is related to the immune status of the patient.\(^{(2)}\)

We emphasize the need for investigating this parasitosis in AIDS patients with clinically suspected pulmonary tuberculosis, especially in those with prolonged diarrhea.

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