Strongyloides stercoralis: a rare and severe presentation in a pregnant woman


1Department of Microbiology, Lozano Blesa University Hospital, Zaragoza, Spain; 2Department of Infectious Diseases, Lozano Blesa University Hospital, Zaragoza, Spain; 3Department of Obstetrics and Gynecology, Lozano Blesa University Hospital, Zaragoza, Spain; 4Department of Pathology, Lozano Blesa University Hospital, Zaragoza, Spain; 5Department of Family and Community Medicine, Corralejo Health Center, Las Palmas, Spain

CASE REPORT

A 23-year-old primigravid Colombian woman in her 21st week of pregnancy presented with acute abdominal pain and oral intolerance. She had arrived to Spain two months before, worked as a saleswoman, and did not report any contact with toxic substances.

At admission, she showed a poor general condition, weight loss (BMI 16), dehydration, low-grade fever, and diffuse abdominal pain with tenderness. Laboratory tests showed an elevated white cell count (16 x 10^9 cells/L), hemoglobin of 11.3 g/dL, normal eosinophil count, and an elevated C-reactive protein. Abdominal ultrasound revealed no pathological findings. Stool and blood cultures were negative. Due to the worsening clinical situation and constant weight loss, a nasogastric tube was inserted.

Further laboratory findings showed a positive fecal occult blood test, and negative serology tests for Toxoplasma gondii, Rubella, Measles, EBV, Syphilis, HIV, and hepatitis. These findings led to the performance of an oral endoscopy, which showed ulcers with mild spontaneous bleeding in the second half of the duodenum and progressive narrowing of the lumen. The histopathological examination of the duodenal biopsy revealed larvae compatible with a parasitic infection: there was presence of several stages of the worm embedded in mucosal crypts (Figure 1). Additional findings included villous flattening, crypt hyperplasia, and an inflammatory reaction consisting of lymphocytes, plasma cells, and eosinophils.

A stool examination (formalin-ethyl acetate technique) revealed both Charcot-Leyden crystals and a huge number of rhabditoid (but not filariform) larvae of Strongyloides stercoralis. Unusually, eggs were also detected (Figure 2). Despite the parasite burden, the eosinophil count remained within normal limits, so we decided to check the HTLV status. After a positive Enzyme-Linked Immunosorbent Assay (ELISA) screening test, an HTLV-I infection was confirmed by Line Immunoassay (LIA).

Therapy with oral ivermectin was initiated with a dose of 200 µgr/kg/day for two days, with resolution of the digestive symptoms and resumed oral tolerance. Another fecal sample was submitted just prior to delivery and, again, a very high load of rhabditoid larvae was detected, which led to the prescription of another course of ivermectin.

Finally, the patient underwent a normal vaginal birth and, a month later both mother and child were parasite-free (tested by PCR).

DISCUSSION

The patient presented with intestinal obstruction due to S. stercoralis. This condition, as well as severe strongyloidiasis in pregnant women, is seldom reported. This parasite has a complex life cycle consisting of a free-living and a parasitic cycle. In both cycles, the parasite reaches the intestinal tract, where the adult female lives and reproduces. The larvae can either be passed in the stool or cause autoinfection. S. stercoralis is known to be one of the only two helminths with this ability, where the infective larvae can penetrate the mucosa or the perianal skin and reach any other tissue (Page et al., 2018).
Strongyloidiasis is endemic in tropical and subtropical regions but can also be found in non-endemic areas due to increases in travelling and migration (Montes et al., 2010). In some cases, the infection can evolve to a hyperinfection syndrome due to an increase in the larval burden in the traditional reproductive route (skin, gut, and lungs), or can cause a disseminated infection (when larvae are found outside this route, e.g., heart, liver, central nervous system). The risk of suffering from this severe form of the illness is increased in immunosuppressed patients, such as those under corticosteroid treatment or with positive HIV or HTLV-I status (Schrär et al., 2013).

In our case, the histopathological results of the biopsies pointed to the final diagnosis. Finding a prominent eosinophilic infiltrate and Charcot-Leyden crystals should prompt a search for parasitic infestation. The small intestine is the most commonly affected site, but in rare cases the stomach and colon can also be involved (Montgomery and Voltaggio, 2012).

Previously, Malézieux-Picard (2017) reported a similar case in a pregnant woman (Malézieux-Picard et al., 2017). Nevertheless, pregnancy alone does not seem to be the major cause, since only very few reports of intestinal obstruction due to *S. stercoralis* in pregnant women have been published and pregnancy has not been described as a risk factor for severe strongyloidiasis. Thus, we speculate that the underlying HTLV infection was the only trigger in our case. Indeed, the clinical condition of our patient dramatically improved once the anthelminthic therapy began.

Oral ivermectin remains the treatment of choice for strongyloidiasis (Nutman, 2017). In case of oral intolerance, malabsorption syndromes or in worsening clinical situa-
ations, the possibility of using ivermectin either through the rectal or subcutaneous route has been described (Tarr et al., 2003).

HTLV-1 infection is a well-known risk factor for a more severe presentation of strongyloidiasis, including disseminated infections and intestinal obstruction (Toledo et al., 2015). HTLV-1 decreases the Th2 type of immune response (Toledo et al., 2015). This explains both the increase in prevalence and the severity of strongyloidiasis in these patients. Mother to child is one of the most important routes for HTLV transmission, thus, breastfeeding should be avoided (Carneiro-Proietti et al., 2014). In this way, strongyloidiasis helped us expose an underlying viral infection. We suggest that HTLV status should be screened in every severe S. stercoralis infection or when, despite a correct treatment, a relapse is observed.

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References


