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Systemic lupus erythematosus and Crohn's disease: An uncommon association of two autoimmune diseases

Sirs,

The association of systemic lupus erythematosus (SLE) with inflammatory bowel disease is rare (1-3). Moreover, the differentiation between digestive manifestations of SLE and extra-digestive manifestations of Crohn's disease (CD) may be particularly difficult (2). Also, sulphasalazine therapy has been implicated in the development of lupus-like syndrome in patients with inflammatory bowel disease (4,5). We report a new case of the association between these two autoimmune diseases. In our case, SLE manifestations preceded the clinical manifestations of CD and, consequently, no drug therapy implication in SLE manifestations

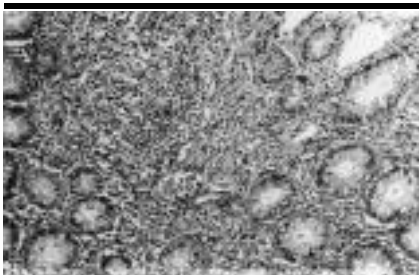


Fig. 1. Infiltration of lymphocytes and plasmatic cells, lymphoid hyperplasia and multinucleate giant cell granuloma formation.

was found.

A 41-year-old Caucasian woman presented at the hospital because of abdominal pain and diarrhea. She had been diagnosed 2 years earlier as having SLE because of a photosensitivity malar rash, oral ulcers, peripheral synovitis, positive antinuclear antibodies at 1/1280 (by indirect immunofluorescence using as substrate Hep2 cells), and positive anti-nDNA (by indirect immunofluorescence using as substrate crithidia luciliae) at 1/80. At the time of the onset of diarrhea she was also on treatment with non-steroidal anti-inflammatory drugs and chloroquine (250 mg/day). She recalled her symptoms having begun with abdominal pain and bloody diarrhea (8 times a day) 1 week before admission.

Physical examination showed abdominal distention, but no clinical features of SLE flare-up were observed. Apart from anemia (hemoglobin 9.04 g/dl) no other hematological abnormalities were found. Hepatic and renal function tests were normal. Antinuclear antibodies and anti-native DNA were positive at 1/320 and 1/40, respectively. Serum levels of C3 and C4 were within normal ranges. IgG and IgM anticardiolipin antibodies and ANCA tests were negative. Negative results for both stool cultures and parasitologic examinations of the feces ruled out enteric bacterial pathogens or amebiasis. Chest radiograph and electrocardiogram were also normal.

A colonoscopy showed edematous and hyperemic areas with aphthous ulcers, pseudopolyps and deep transversal and longitudinal fissures. These features were compatible with a moderately active CD. Biopsy specimens confirmed the endoscopic findings. Microscopic examination revealed a chronic inflammation affecting all layers of the colon, which was more severe in the lamina propria of the mucosa with infiltration of lymphocytes and plasmatic cells, aphthous ulcers in areas with lymphoid hyperplasia, fissures extending into the serosa, epithelial cells and multinucleate giant cell granulomas. Figure 1 shows the inflammatory infiltrate with granuloma formation. Neither thrombi in vessels nor any other evidence of ischemic colitis were observed. Once the diagnosis was established steroid therapy (prednisone 1 mg/Kg) and azathioprine (2 mg/kg) was started and a rapid improvement of symptoms was observed. At present, 6 months after the diagnosis the patient is free of symptoms.

The diagnosis of SLE in our patient was based on the criteria proposed by the American College of Rheumatology (formerly the American Rheumatism Association) (6). All parts of the gastrointestinal tract may be involved in SLE (7). However, the development of CD in patients with lupus unrelated

to drugs is exceptional (2, 3). Although vasculitis in the bowel due to SLE may be difficult to distinguish from the onset of an inflammatory bowel disease, the onset of diarrhea and abdominal pain in our patient was not associated with a flare of her SLE. In this case, histological findings are the cornerstone to making the differential diagnosis between the two conditions. Evidence of immunofluorescence deposits of immunoglobulins and complement on the capillary wall and electron-dense deposits on electronic microscopy are needed to establish a diagnosis of lupus vasculitis involving the gut.

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