Likewise, 1 year-old Juan For Polymyositis have demonstrated that the rate of placental transfer is very low.\(^8\) Mahadevan et al.,\(^8\) showed that concentrations of infliximab and adalimumab, but not CZP, are greater in umbilical cord than in maternal serum; mean levels of infliximab and adalimumab in cord reach concentrations between 150% and 160% greater than the concentrations in maternal serum.\(^9\)

There are other circumstances in which placental transfer of antibodies after weeks 20–22 can be detrimental to the fetus/neonate. Typically, in cases of rheumatoid arthritis and systemic lupus erythematosus, the passage of anti-Ro/SSA and/or anti-La/SSB autoantibodies of IgG isotype can cause an eruption due to exposure to the sun or the development of congenital heart block, which affects roughly 2% of fetuses/neonates of patients who have these autoantibodies.\(^6\)

With the lack of data on the safety of mAb, the performance of clinical trials in pregnant patients is ruled out for ethical considerations, and the decision to use them would depend in each case on the clinical situation, as well as that of the potential benefits and risks for the mother, fetus or newborn. Long-term observational studies would enable us to confirm the efficacy and safety of category B mAb during pregnancy, to determine whether gestational exposure to mAb involves a long-term risk for the immune system being developed in the newborn, or should this vary depending on the trimester for exposure. It is important to consider that the administration of vaccines with alive or attenuated virus or bacteria, an indication that is present in certain vaccine calendars for newborns, for example, with bacillus Calmette-Guérin (BCG), with which, infection can have a fatal outcome.\(^5\) For these reasons, it is recommendable that they be postponed until the sixth month of life.

**Conflicts of Interest**

LV: Abbvie, Roche Farma, Bristol-Myers Squibb, Pfizer, UCB, MSD and GSK; JGOb: nothing to declare; DHF: nothing to declare;

**Polymyositis in a Patient With Ulcerative Colitis**

**Polimiositis en un paciente con colitis ulcerosa**

Dear Editor,

Ulcerative colitis (UC) is an inflammatory disease that affects the colorectal mucosa in a diffuse form, continuously and superficially.\(^1\) Polymyositis (PM) belongs to the spectrum of idiopathic inflammatory myopathies, and is differentiated from dermatomyositis (DM) because of the absence of the characteristic skin rash.\(^2\) Muscle involvement in the form of PM has rarely been described in UC. We report the case of a patient who was initially diagnosed as having UC, and developed PM over the course of his disease. We also review cases published to date. Our patient was a 46-year-old man, an ex-smoker, who presented with persistent bloody diarrhea with no other associated symptoms, and was diagnosed with UC. At the age of 58 years, he presented with an episode of pain and swollen hands, and was too weak to climb stairs. Laboratory tests detected increases in creatine kinase (CK) (1578 U/L) and lactate dehydrogenase (LDH) (506 U/L), although all other inflammatory parameters were normal. The rest of the biochemical study, including complete blood count, showed normal values. The immunological study demonstrated the presence of antinuclear antibodies, at a titer of 1/160, and negative tests for anti-extractable nuclear antigens (ENA), anti-DNA antibodies, antineutrophil cytoplasmic antibodies (ANCA), anti-centromere antibodies, anti-Scl-70 and antibodies specific for myositis (Jo-1, PL7, PL12, Oj, Ej, SRP, Pm-Scl, Mi2 and Ku). A muscle biopsy of quadriceps was performed (Fig. 1), and evidenced variability in the size of the muscle fibers, with some necrotic fibers, regenerative basophilic fibers and frequent mononuclear inflammatory cell infiltrates, mostly lymphocytic and located in the endomysium. The expression of the major histocompatibility complex class I antigens was positive in the muscle fibers, all of which was compatible with the diagnosis of PM. Treatment was begun with a tapering dose of prednisone starting at 20 mg/day, to be followed by methotrexate (maximum dose 25 mg/week), and normalization of the CK and LDH levels was achieved after 16 months of treatment.

Fig. 1. Image of the muscle biopsy: (A) There are muscle fibers with regenerative peripheral basophilic areas (hematoxylin–eosin stain ×63). (B) Muscle expression of the class I antigens of the human leukocyte antigen (HLA) system. (C) There is an endomysial lymphocyte cell infiltrate located in the (hematoxylin–eosin stain ×40). (D) Variability of the size of the muscle fibers and periendothymal (hematoxylin–eosin stain ×10).

Table 1
Characteristics of the Patients With Polymyositis Associated With Ulcerous Colitis.

<table>
<thead>
<tr>
<th>Authors [Ref]</th>
<th>Sex</th>
<th>Age (years)</th>
<th>Associated defects</th>
<th>AAB</th>
<th>Time between the diagnosis of UC and the development of PM</th>
<th>Muscle groups affected</th>
<th>Extramacular manifestations</th>
<th>Treatment</th>
<th>Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hernández et al.3</td>
<td>M</td>
<td>11</td>
<td>HLA-B27 (−) Sacroiliitis</td>
<td>No</td>
<td>Simultaneous</td>
<td>Regions proximal to extremities and neck flexors</td>
<td>No</td>
<td>Oral prednisone 1.5 mg/kg/day</td>
<td>Improvement of muscle manifestations</td>
</tr>
<tr>
<td>Evrard et al.4</td>
<td>M</td>
<td>33</td>
<td>Not associated chronic liver disease</td>
<td>No</td>
<td>4 years earlier</td>
<td>Lower extremities</td>
<td>Fever</td>
<td>Oral sulfasalazine 1.5 g/day Betamethasone enemas</td>
<td>Improvement of bowel and muscle manifestations</td>
</tr>
<tr>
<td>Kaneoka et al.5</td>
<td>W</td>
<td>57</td>
<td>Not associated liver cirrhosis</td>
<td>ANA 1/160 RF (+)</td>
<td>11 years earlier</td>
<td>Torso and regions proximal to extremities and neck flexors</td>
<td>No</td>
<td>Oral prednisolone 40 mg/day</td>
<td>UC quiescent, PM with moderate activity. Death due to liver failure</td>
</tr>
<tr>
<td>Chugh et al.6</td>
<td>W</td>
<td>78</td>
<td>Esophagitis and erosive gastritis</td>
<td>No</td>
<td>15 years earlier</td>
<td>Torso and regions proximal to extremities and neck flexors</td>
<td>Polyarthritis</td>
<td>Oral prednisolone 40 mg/day Oral 5-aminosalicylic acid 400 mg/8 h Prednisolone enemas</td>
<td>UC in remission, improvement of muscle manifestations</td>
</tr>
<tr>
<td>Voigt et al.7</td>
<td>W</td>
<td>33</td>
<td>−</td>
<td>ANA 1:2560 Anti-DNA (+)</td>
<td>4 years earlier</td>
<td>Regions proximal to extremities</td>
<td>No</td>
<td>IV prednisolone 200 mg/day for 7 days, followed by oral prednisone 60 mg/day</td>
<td>UC quiescent, PM in remission</td>
</tr>
</tbody>
</table>

AAB, autoantibodies; ANA, antinuclear antibodies; CK, creatine kinase; IV, intravascular; M, man; PM, polymyositis; Ref, reference; RF, rheumatoid factor; UC, ulcerative colitis; W, woman.
Muscle involvement in inflammatory bowel disease is uncommon; in fact, only 7 cases associating PM and UC have been published to date1-9 (Table 1). Only in the patient we describe was the diagnosis reached because of the increase in muscle enzymes. The response to glucocorticoid treatment was so good, in general, except in an individual who was refractory to several immunosuppressive agents and whose disease entered remission with methotrexate.9 It has been suggested that the basis for the development of PM in UC results from the common immune-mediated mechanism, in which bowel inflammation and the damage to the mucosa would lead to a release of antigens, stimulating the production of antibodies that would damage the muscle. In conclusion, the development of PM in UC should be taken into account, especially when these patients describe symptoms such as myalgia, muscle weakness or present an increase in muscle enzymes. Although there are few cases reported in the literature, it may be underdiagnosed given that the symptoms may be few and not very specific, and can be attributed to other causes.

Conflict of Interest

The authors declare they have no conflict of interest directly or indirectly related to the contents of this manuscript.

References


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Spondylodiscitis Without Endocarditis Caused by Streptococcus mitis

To the Editor,

Spondylodiscitis is an infection of an intervertebral disc and the vertebrae. Its incidence is on the rise in recent years. The causative microorganism in most adult patients is Staphylococcus aureus. To date, Streptococcus species have had little relevance as causative agents of vertebral osteomyelitis. However, the number of cases caused by these microorganisms has increased in recent years.1 We report the case of a patient with spondylodiscitis caused by Streptococcus mitis.

The patient was a 49-year-old man, an ex-smoker, with nothing else remarkable in his clinical history. He presented with a 4-week history of low back pain with inflammatory features and pain that radiated down his left leg. He did not have associated fever or metabolic syndrome. There was no evidence of skin lesions, either, and he did not consume farmouse dairy products. He had not undergone any dental procedures in the preceding 2 years. Notable findings in the results of laboratory tests were the absence of leukocytosis in the complete blood count and a normal differential leucocyte count, but elevation of acute-phase reactants was detected (erythrocyte sedimentation rate [ESR], 80 mm; C-reactive protein [CRP], 42 mg/L). Serological tests for human immunode-