Widespread Cutaneous Necrosis as the First Clinical Manifestation of Secondary Antiphospholipid Syndrome

Necrosis cutánea extensa como la primera manifestación clínica de un síndrome antifosfolípidico

To the Editor:

Antiphospholipid syndrome (APS) is an acquired prothrombotic state characterized by recurrent thromboses, pregnancy loss, thrombocytopenia, and the presence of antiphospholipid antibodies, antiphospholipid antibodies, lupus anticoagulant and/or anti-β2-glycoprotein antibody. The original clinical and laboratory criteria for APS, called the Sapporo criteria, were first published in 1999. These were replaced by the Sydney criteria in 2006 when patients were required to have at least 1 clinical criterion and 1 laboratory criterion for a diagnosis of APS to be made.

A wide variety of dermatologic manifestations have been described in patients with APS, including livedo reticularis, livedoid vasculitis, digital gangrene, erythematous macules, skin ulcerations, and, on rare occasions, extensive cutaneous necrosis. We report the case of a patient with widespread cutaneous necrosis as the initial manifestation of APS.

A 48-year-old woman presented at the emergency room with a 2-week history of fever, arthralgia, malaise, and chest pain. On examination she was noted to have 3 bullous erythematous violaceous plaques on her right leg (Fig. 1). Her medical history was remarkable for hypothyroidism, systemic lupus erythematosus (SLE), and obsessive-compulsive disorder. Medications included thyroxine, olanzapine, risperidone, and sertraline. Laboratory tests revealed thrombocytopenia (platelet count 25,000) and normal prothrombin and partial thromboplastin times. Lupus anticoagulant was present and the antinuclear antibody titer was 1:320. C3 and C4 levels were normal.

The lesions deteriorated rapidly despite initiation of corticosteroid therapy with prednisolone at a dosage of 1 mg/kg/d. The dermatology service was called to evaluate the patient on day 7 of hospitalization. Examination revealed extensive cutaneous necrosis on the right leg (Fig. 2). Further laboratory data revealed an elevated anticardiolipin antibody immunoglobulin (Ig) G titer (30; normal <23). Anti-β2-glycoprotein antibody IgG levels were also elevated, but cryoglobulins and cryofibrinogens were normal. A 4-mm punch biopsy from the periphery of a lesion revealed hemorrhage throughout the dermis with organized thrombi in many dermal vessels, with no evidence of vasculitis (Fig. 3). No further lesions developed after the addition of intravenous heparin. At discharge, the heparin was replaced by long-term oral anticoagulation therapy and prednisone was tapered over 8 weeks. At the 3-month follow-up, only residual scarring remained (Fig. 4). There have been no new thrombotic events in 1 year of follow-up.

Figure 2 Cutaneous lesions on day 7 of hospitalization. Extensive cutaneous necrosis on the right leg.

Figure 3 Skin biopsy. Extensive epidermal basal vasculopathy, hemorrhage throughout the dermis with organized thrombi in many dermal vessels and no evidence of vasculitis.

Figure 1 Cutaneous lesions on admission. Bullous erythematous violaceous plaques on the right leg.
**Metastatic Melanoma of the Tongue: A Rare Case**

**Melanoma metastásico de la lengua: un caso raro**

To the Editor,

Cutaneous melanoma is an extremely aggressive malignant tumor arising from melanocytes and accounting some 15% of all cancers. Incidence varies from 3–5 cases per 100,000 inhabitants and year in the Mediterranean region to 12–20 cases per 100,000 inhabitants and year in Nordic countries, and is still increasing worldwide. Cutaneous melanoma is believed to be a cancer that primarily affects white skin, and the risk of developing these tumors is 10 times higher in white-skinned populations than in those with darker skin. Melanoma is known for its aggressiveness, and metastases to bones, lungs, brain, liver, or lymph nodes are expected. However, there have been very few reports of oral metastases, which mainly affect the gingiva, tongue, tonsils, and mandible.

We describe the case of an 86-year-old white man admitted with a black lesion on the tongue that had persisted for about 2 months. During the consultation, the patient rejected the idea that he had a disease and complained about the appearance, concurrently with the lingual lesions, of**

**Bibliografía**


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