**Case report**

**Extraocular myositis as an initial manifestation of polyangiitis granulomatosis**

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**A B S T R A C T**

Granulomatosis with polyangiitis (GPA) is a systemic vasculitis with multiorgan involvement. Although ocular involvement has frequently been described, there are few reported cases of extraocular myositis in this disease.  
The case is presented of a Mexican woman with GPA who debuted with extraocular myositis.

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**Miositis extraocular como manifestación inicial de granulomatosis con poliangeitis**

**R E S U M E N**

La granulomatosis con poliangeitis (GPA) es una vasculitis sistémica con afectación multiorganica. El compromiso ocular ha sido descrito frecuentemente, sin embargo, existen pocos casos reportados de miositis extraocular en esta enfermedad.  
Presentamos el caso de una mujer mexicana con GPA que comenzó con miositis extraocular.

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Introduction

Granulomatosis with polyangiitis (GPA) is a systemic small-vessel vasculitis which affects almost any organ. Ocular manifestations occur in 30–50% of patients, and in 8% of cases can lead to permanent blindness. The orbital commitment occupies the second place after the scleral involvement, occurring in 5–30% of patients with GPA. The orbital inflammation can occur primarily as a granulomatous lesion (orbital pseudotumor) or secondarily, either by diffuse infiltration of a nasal disease or by extraocular myositis. Although the orbital pseudotumor is the most frequent cause of ocular proptosis in GPA representing 40–60% of orbital tumors in this disease; we should not forget that there are some other secondary ocular manifestations with orbital involvement in GPA.

The clinical and prognostic relevance is that the orbital disease in this vasculitis has been associated with functional loss of the eye and serious systemic disease.

Below we present the case of a woman with GPA who started with extraocular myositis, the rarest form of orbital involvement in this vasculitis.

Case presentation

A 56-year-old woman who started with pain and right ocular proptosis in 2013 was treated with systemic steroids, with which the episode remitted. In 2015 she presented again pain and right ocular proptosis and was treated by the ophthalmologist who documented by computerized axial tomography (CAT) of the orbits, extraocular myositis; in order to obtain a diagnosis, the patient underwent a biopsy of the upper rectus muscle and the lacrimal gland. The histopathological result was nonspecific inflammation; (Fig. 1A and B) and she received prednisone at a dose of 1 mg/kg/day for 2 months with which there was a remission. In March 2016 she presented weight loss, fever, neuropathic pain in the lower extremities, painful right ocular proptosis; and extraocular myositis was documented by CAT scan (Fig. 2A and B). The diagnosis of GPA was integrated due to the ocular involvement, constitutional symptoms, sensory neuropathy in lower extremities and positive antineutrophil cytoplasmic antibodies (ANCA) (c-ANCA: 1:40 [reference value: <1:40] and PR3: 5.4 [reference value: <4.3]). The patient received 3 bolus of methylprednisolone 1g and cyclophosphamide 1g monthly for 6 months; subsequently, a CAT scan of the orbits was performed in November 2016 (Fig. 3A and B) in which a remission of the ocular manifestation was found.

Case discussion

GPA is a systemic vasculitis in which any structure of the eye can be compromised, however, the most frequent are the episclera in 20–25% and the sclera in 15–20%. Orbital involvement occurs in 15% of patients with this vasculitis and occupies the second place in ocular compromise.

Due to the nature of the vasculitis itself, there are few series that report ophthalmologic findings in this disease. The series in which the orbital involvement is classified and which describes in more detail is that of Ismailova et al., in which 74 patients with orbital manifestations are described in a series of 226 patients with GPA followed-up for 4 years. Of the 74 patients with orbital involvement, 45 (60.8%) had a granulomatous lesion (orbital pseudotumor), 26 (35.2%) developed rhinosinusitis with orbital involvement and 3 (4%) developed extraocular myositis. The 3 patients who developed extraocular myositis had localized GPA with negativity for ANCA, 2 of them had recurrence of the manifestation and none of them developed systemic disease.

The case presented shows a woman with painful unilateral proptosis with a final diagnosis of GPA, but who started...
steroids and cyclophosphamide showed significant clinical and radiological improvement. Although the biopsy of the extraocular muscle did not show data compatible with vasculitis; the response to treatment, as well as the extraocular clinical manifestations and the positivity of ANCAs allowed us to obtain the diagnosis. The role of biopsy in this orbital manifestation has not been studied.

According to our search, it is the first case of systemic GPA that began with unilateral and recurrent extraocular myositis.

When the ophthalmologist faces a patient with painful unilateral proptosis, he must consider GPA within the etiological possibilities; and the rheumatologist who faces a patient with an established diagnosis of GPA, and who develops painful unilateral proptosis must consider, in addition to the orbital pseudotumor, extraocular myositis as another diagnostic possibility.

**Conclusions**

The ocular manifestations in GPA can represent, from the functional point of view, a catastrophe for the patient, even leading him/her to blindness; it is important that both rheumatologists and ophthalmologists are familiar with the ocular clinical spectrum of this systemic vasculitis, since the timely diagnosis definitely changes the prognosis.

**Conflict of interest**

The authors declare that they do not have any conflict of interest.

**REFERENCES**