Case report

Resistant orbital pseudotumor treated with rituximab in a patient with systemic lupus erythematosus. A case presentation

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ARTICLE INFO

Article history:
Received April 27, 2009
Accepted July 8, 2009

Keywords:
Exophthalmos
Systemic lupus erythematosus
Orbital pseudotumor
Rituximab

ABSTRACT

Ocular manifestations in Systemic Lupus Erythematosus (SLE) are relatively frequent, with a major prevalence of the Keratoconjunctivitis sicca. Nevertheless, the appearance of unilateral exophthalmos secondary to orbital pseudotumor in patients with SLE is extremely rare,1-7 and on occasion it can be refractory to conventional pharmacological treatment (glucocorticoids and immunosuppressants). We present the case of a patient with SLE and orbital pseudotumor refractory to treatment with Cyclophosphamide (CF) and an excellent clinical response, with disappearance of the ophthalmological condition after the beginning of therapy with Rituximab (1 g × 2), continuing after the infusion of two complete cycles without incidents.

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Pseudotumor orbitario refractario tratado con rituximab en paciente con lupus eritematoso sistémico. A propósito de un caso

RESUMEN

Las manifestaciones oculares en el lupus eritematoso sistémico (LES) son relativamente frecuentes, con mayor prevalencia de la queratoconjuntivitis seca. Sin embargo, la aparición de exoftalmos unilateral secundario a pseudotumor orbitario en pacientes con LES es extremadamente rara,1-7 y en ocasiones puede presentarse refractaria al tratamiento farmacológico convencional (glucocorticoides e inmunosupresores). A continuación presentamos el caso de una paciente con LES y pseudotumor orbitario refractario a tratamiento con ciclofosfamida y excelente respuesta clínica con desaparición de la clínica oftalmológica tras el inicio de la terapia con rituximab (1 g × 2), eficacia mantenida tras la infusión de dos ciclos completos sin incidencias.

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Introduction

Ocular manifestations related to Systemic Lupus Erythematosus (SLE) have an incidence estimated between 3%-30%2,6 according to different series, affecting both the anterior and posterior segments of the eye. However, the appearance of exophthalmos secondary to inflammatory orbital pathology in patients with SLE, of which there are few published cases, is extremely rare.1-7 When faced with a unilateral exophthalmos, one must consider a differential diagnosis that includes infectious, neoplastic and endocrine processes (Graves disease), connective tissue diseases and systemic pathologies (vasculitis, SLE), among others.9,10 We present the case of a patient with SLE and unilateral exophthalmos which was secondary to a steroid and immunosuppressant resistant orbital pseudotumor, treated satisfactorily with rituximab (RTX).

Case report

Forty-nine year old woman with a history of hypertension, diabetes type 1 under insulin treatment with good glycemic control, extrinsic asthma and intermittent episodes of unilateral amaurosis of the right eye. In November 2006 she presented...
headache, diplopia and unilateral exophthalmos, and was evaluated and treated at the Neurology department. A magnetic resonance imaging (MRI) test displayed a cranial space-occupying lesion compatible with right intraorbital or dilated ophthalmic or orbital vein and a partial occupation of the right cavernous sinus. Suspecting a cavernous sinus syndrome, the start of anticoagulation was decided upon with acenocoumarol and testing for hypercoagulability and autoimmunity was carried out, with these findings: aCL IgG and IgM +, ANA + and anti-native DNA+. She was referred to our consultation in September 2007 due to data suggesting an autoimmune disease. The patient reported a history of rythmic inflammatory polyarthralgia for 2-3 years prior which improved after taking NSAIDs, as well as fever, fatigue and recurrent oral ulcers with an onset 6-7 months prior and a progressive increase of dyspnea in the last two months going to present with mild-moderate efforts to occurring at rest. Physical examination revealed mucocutaneous pallor, right exophthalmos, purpuric lesions on the lower limbs, residual oral ulcers, tachycardia (110 bpm) and synovitis of the wrists. We performed a biopsy of the lesions, and indicated the need for hospitalization to complete her study. The complementary tests revealed microcytic hypochromic anemia with a positive direct antiglobulin test, elevated acute phase reactants, hypocomplementemia, positive ANA with a high titer (>1/320) and positive anti-native DNA, positive IgG and IgM aCL, a urinary sediment without evidence of renal disease in a high percentage of cases. The authors have no disclosures to make.

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Discussion

Patients with SLE may have a variety of ocular manifestations, the most common being keratoconjunctivitis sicca. The appearance of exophthalmos as a manifestation in patients with SLE is extremely rare, occurring as secondary to orbital pseudotumor in most cases. The first description of orbital pseudotumor dates back to 1905, and was published by Birch-Hirschfeld, who classified it as a non-neoplastic inflammatory entity of unknown etiology in a large percentage of cases. It was composed of partly mature lymphocytes, plasma cells, macrophages and leukocytes, with varying degrees of fibrosis. This inflammatory process can affect the orbital tissues diffusely or specifically leading to different clinical variables: lacrimal gland (dacracyoanitis), muscle (miosisitis), Tenon capsule (tenositis) and, less frequently, orbital vessels and nerves affection. For its etiologic diagnosis it is necessary to perform a proper history and physical examination and additional laboratory and imaging tests (mainly CT or MRI). Sometimes it is necessary to perform a biopsy of the lesion to determine the underlying pathology.

In patients with unilateral exophthalmos and orbital pseudotumor, one should consider diseases such as Graves’ disease, neoplasms, infectious or less common causes such as cavernous sinus thrombosis, amyloid deposition, retrobulbar hematoma, hemangioma, sarcoidosis and histiocytosis in association with connective tissue diseases as part of their differential diagnosis.

In conclusion, with treatment, high efficiency has been described with the use of corticosteroids administered as high dose intravenous pulses as induction therapy and half the dose later on as a maintenance dose to prevent recurrence. Immunosuppressants such as azathioprine, CF and antimalariais (chloroquine / hydroxychloroquine) have been used as adjutants for remission or glucocorticoid-sparing therapy. TNF-inhibiting therapies have been shown to be effective in patients with inflammatory orbital affection resistant to the drugs mentioned above. In our case, because the case was a patient with a condition in which the B cell has a predominant role, and the resistance of the ocular problem to treatment with CF, we decided to initiate treatment with a monoclonal antibody directed against CD 20+cells (RTX), with an excellent clinical response and control of ocular symptoms after infusion of the first dose of the drug. This response persisted two months after the infusion of the second full cycle. Although we did not entirely rule out the probability of a lymphoproliferative process, the clinical presentation, with intense and very painful proptosis and conjunctival injection (findings found with greater prevalence in orbital pseudotumors), and the presence of clinical and laboratory data suggesting the diagnosis of SLE, made us decline the possibility of a diagnostic biopsy of the lesion. Experience has shown that the use of RTX in the treatment of patients with SLE resistant to various immunosuppressive drugs, including CF, shows promising results, especially for the control of renal disease in a high percentage of cases. There is little evidence for the treatment of orbital inflammatory disease with this molecule in the literature, but in secondary ophthalmopathy due to Graves’ disease, it appears to be effective. In conclusion, RTX could be an alternative to conventional treatments when they show an insufficient response.

Disclosures

The authors have no disclosures to make.
References