Eye pain in 13-year-old teenager: a case report of idiopathic orbital myositis

Renata Barbosa Paolilo 1D Paula da Cunha Pinho 1D

Department of Neurology, Samaritano Hospital of São Paulo, São Paulo, Brazil

A previously healthy 13-year-old male presented with sporadic headache that evolved to continuous right eye pain and blurred vision within a week. There was no previous infection or relevant family history. Neurological examination revealed restricted abduction of right eye and horizontal diplopia. Contrast-enhanced orbital magnetic resonance imaging (MRI) is shown in Figures 1 and 2. Cerebrospinal fluid analysis and extensive laboratory workup were normal. He was treated with intravenous corticosteroid for 5 days and oral tapering for 2 weeks. Orbital MRI performed within 2 months resulted normal. After a three-year follow-up, the patient keeps asymptomatic.

Orbital myositis is a non-infectious inflammatory condition of unknown etiology. Pediatric cases account for 6-17% of all orbital inflammatory disorders. Besides being rare, prompt recognition, diagnosis, and treatment are important. Axial and coronal orbital MRI is a sensitive diagnostic tool.

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Figure 1. Coronal fat-suppressed contrast-enhanced T1 shows enlarged right medial muscle and mild infiltration of the surrounding fat.

Figure 2. Coronal fat-suppressed T2 shows high signal in the right medial muscle. The classic appearance of extraocular muscle (EOM) myositis includes a unilateral thickening of one or two EOMs, often also involving the surrounding fat, tendon and myotendinous junction.