Bilateral Brown’s syndrome in a mother and her son: case report

Síndrome de Brown bilateral em mãe e filho: relato de caso

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ABSTRACT

This case report describes clinical data from mother and son with bilateral Brown’s syndrome and highlights possible genetically determined predispositions.

Keywords: Ocular motility disorders/genetics; Ocular motility disorders/pathology; Strabismus/congenital; Syndrome; Tendons/physiopathology; Oculomotor muscles/physiopathology; Case reports [Publication type]

INTRODUCTION

Brown’s syndrome (or superior oblique tendon sheath syndrome) was first described by Brown in 1950(1). This motility defect consists of the inability to raise the adducted eye above the midhorizontal plane, with a positive forcedduction test in supra-adduction. Less elevation restriction can be found in the midline and a smaller elevation deficiency is detectable in abduction. Exodeviation usually increases as the eyes move upwards above the midline.

In 1973 Brown redefined this syndrome(2) and divided it into two subtypes, i.e., true and simulated sheath syndrome. The true syndrome is congenital, permanent and invariably associated with a positive traction test of the involved adducted eye in the field of elevation. The true syndrome is the most common form(2-3). Many reports have tried to elucidate the true form and its etiologies. A taut superior oblique tendon(1-3) is a possible cause. Katz et al.(4) reported Brown’s syndrome in monozygotic twin girls. Association with autosomal dominant pathology(5) and genetic inheritance have also been reported(6-7).

We describe here the eye examination of two cases of bilateral Brown’s syndrome occurring in mother and son.

CASE 1

A seven-year-old boy was referred to us because his mother noticed that he could not elevate his eyes, with an exodeviation occurring when he tried to elevate them, since the age of 8 months. The medical history of the boy and his family was unremarkable and the child had no complaints. Eye examination revealed 1.0 (logMAR=0) visual acuity in each eye, cycloplegic refraction was +1.00 spheric diopter in each eye, both pupils were reactive to light, and slit lamp examination and indirect fundoscopy were within normal limits. Motility examination (Figure 1) showed an abnormal head position with a mild chin elevation. In the cover test, he showed orthophoria in an abnormal head position and a small exophoria in the primary position. In dextroversion, he had right hypertropia and in levoversion he had left hy-
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The mother of the boy (case 1) had the same motility defect (Figure 2), observed by her husband some years ago. She was 32 years old and reported no previous inflammatory diseases. Her visual acuity was 1.0 (logMAR=0) in each eye and slit lamp examination and fundoscopy were equally normal. When submitted to the cover test, she showed orthophoria in the primary position and in infraversion. In supraversion, there was important exotropia, as observed in her son, showing a V pattern. She was also unable to elevate both eyes in adduction. In dextro- and levoversion she showed right and left hypertropia, respectively. The forced duction test could not be done. The Titmus test revealed stereopsis of 40”.

The boy appears to have a true Brown’s syndrome. Although the mother has the same clinical findings, she was not examined previously. A forced duction test was not performed on her, but it appears that she has also a Brown’s syndrome.

Figure 1 - The nine positions of gaze, boy

Figure 2 - The nine positions of gaze, mother
DISCUSSION

Little is known about the pathogenesis of true Brown’s syndrome. Although the superior oblique tendon is usually found to be abnormally taut, the cause of this anatomic dysfunction is still unknown, but some previously published case reports have suggested a genetically determined predisposition in this syndrome, consistent with recessive inheritance at the DURS1 locus and dominant inheritance with reduced penetrance at the DURS1, DURS2, and FEOM1 loci. The almost perfectly symmetrical bilateral Brown’s syndrome found in the here reported mother and child supports this hypothesis.

RESUMO

Este relato de caso descreve achados clínicos de mãe e filho com síndrome de Brown bilateral e discute a possibilidade de predisposição genética.

Descritores: Transtornos da motilidade ocular/genética; Transtornos da motilidade ocular/patologia; Estrabismo/congênito; Síndrome; Tendões/fisiopatologia; Músculos oculomotores/fisiopatologia; Relatos de casos [Tipo de publicação]