Septo-optic dysplasia - Case report

Displasia septo-optica - Relato de caso

ABSTRACT

The clinical triad of septo-optic dysplasia (SOD) comprises the absence of the septum pellucidum, congenital optic nerve dysplasia, and multiple endocrine disorders. When any two of these factors are present, the condition is defined as an incomplete form of SOD. The authors report the case of an incomplete form of SOD in a 9-year-old boy with low vision and nystagmus present from birth. The bilateral ophthalmoscopic examination revealed small papillae with double contour images. Magnetic resonance imaging showed hypoplasia of the optic nerve bilaterally, chiasm and absence of the septum pellucidum.

Keywords: Optic disc/pathology; Optic disc/abnormalities; Optic nerve/pathology; Optic nerve/abnormalities; Optic nerve diseases/diagnosis; Septo-optic dysplasia/congenital; Septum pellucidum/abnormalities; Case report

INTRODUCTION

Septo-optic dysplasia (SOD) is a rare congenital syndrome characterized by an absent septum pellucidum, a primitive optical vesicle, and dysplasia of the optic nerve, chiasm and tract. This syndrome was described by De Morsier in 1956 who found in nine of 36 patients with agenetic septum pellucidum an association with optic nerve hypoplasia (1). Optic nerve hypoplasia is mandatory for the diagnosis and results in a variable degree of decreased vision.

The clinical association of SOD with pituitary hormone deficiency was established in 1970 (2). Its pathogenesis is still unknown but it has been associated with low maternal age and nulliparity (3-5).

We report the case of an incomplete form of SOD and describe the methods by which the diagnosis was documented.

CASE REPORT

Male, born at 40 weeks of gestation, with a weight of 3.4 kg after a normal pregnancy and labor. He was the first child of nonconsanguineous parents. The mother was 18-years-old at delivery and denied exposure to alcohol or drugs. The infant had an uncomplicated hospital course and was discharged after two days.

He was referred to our hospital at the age of nine due to complaints of low visual acuity and nystagmus from birth. Physical examination revealed a 130-cm-tall boy with normal findings at a general medical examination. His visual acuity (Snellen) was 20/100 on the right and 20/200 on the left that could not be improved with refraction. Both pupils responded normally to direct and consensual light. Versions were normal and rotatory nystagmus was present. Ophthalmologic examination disclosed pale, small-sized optic
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discs, encircled by a hypopigmented avascular scleral crescent, characterizing a double contour of the papilla (double ring sign) (Figure 1). The retinal vessels and macula were normal.

Magnetic resonance imaging revealed absence of the septum pellucidum (Figure 2), hypoplasia of the optic nerves (Figure 3) and chiasm. Endocrinologic test results (basal cortisol, adrenocorticotropic hormone, growth hormone, thyroid stimulating hormone, serum thyroxine, testosterone) were normal. A diagnosis of septo-optic dysplasia was established.

**DISCUSSION**

Septo-optic dysplasia has been described as a malformation of the central nervous system characterized by the absence of the septum pellucidum, and hypoplasia of the chiasm and optic nerves. It was first described by De Morsier in 1956(1). The full syndrome was described in 1970(2) by Kaplan et al. who presented six patients with pituitary hormone deficiency, optic nerve hypoplasia and midline abnormality of the prosencephalon. Optic nerve hypoplasia may occur in normal individuals or in those with disorders of the central nervous system. It is generally bilateral but it can be asymmetrical or unilateral(6).

The cause of septo-optic dysplasia remains uncertain. Interestingly, as reported in other reports(3-5), our patient was associated with low maternal age and nulliparity.

There is a wide variability of the clinical picture. Usually patients with SOD come to medical attention for signs related to decreased visual acuity, such as congenital nystagmus, or small stature.

In some cases, intellectual and neurological development as well as language and behavior of SOD patients have been described as normal. Williams et al.(7) studied a group of children with SOD and noted that an absent septum pellucidum was not associated with important neurological and intellectual deficiencies. Approximately two thirds of the SOD patients suffer from hypothalamic-pituitary dysfunction. The most commonly affected hypopituitary hormone is growth hormone. Hypopituitarism may become apparent at any age.
Although early onset is most common, absence of endocrine deficiencies does not preclude the development of endocrine deficiencies in later life(8).

A fundoscopic examination reveals a small optic disc encircled by a halo of hypopigmented tissue caused by hypoplasia of the retinal epithelial pigment. The retinal vessels are normal(6).

The wide spectrum of anatomical malformations of the brain and optic apparatus in SOD can be visualized by magnetic resonance imaging or computed tomography.

A diagnosis of septo-optic dysplasia should be suspected in patients who have a visual deficiency and a congenital nystagmus as well as a fundoscopic examination that reveals a double image of the papilla, an imaging examination showing agenesis of the septum pellucidum, and hypophyseal or hypothalamic hormonal deficiencies.

Our case reinforces the notion that SOD may be a relatively benign entity and remain unsuspected. Our patient’s intelligence was normal. He had no evidence of pituitary dysfunction. It is important to note that patients with SOD need long-term neurological, ophthalmological and endocrinological investigation.

REFERENCES