Rare Case of Optic Nerve Glioma Revealed by Russell Syndrome

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Abstract

Optic nerve glioma (or optic glioma), is a rare form of glioma which affects the optic nerve. There is very little data available in the literature that describes this pathology. It represents 5% of brain tumors in children. The diagnosis is always made on the occasion of ophthalmological clinical signs (nystagmus, loss of visual acuity). In some cases, its mode of revelation is uncommon, delaying the diagnosis and the treatment. Through our work, we have reported the case of a 17-month-old infant with optic chiasm glioma, revealed by diencephalic cachexia evolving since the age of 5 months. The child was put on enteral nutrition at constant flow rate and polychemotherapy with an improvement in the clinical state and radiological stabilization after a 3-month follow-up.

Introduction

Diencephalic cachexia (Russell syndrome) was first described in 1951 by Russel [1]. It affects young children under the age of one with an average age of 6 months. It includes all of the clinical signs (severe weight loss despite an adequate caloric intake and absence of neurological abnormalities) linked to hypothalamic dysfunction, related to lesions occupying the space of the optic chiasm region (low grade gliomas) or tumors near the fourth ventricle [2] and whose mechanism remains unexplained. We report a case of optic chiasm glioma, diagnosed at 17 months of age with severe undernutrition and a facial dysmorphia explained by Russell syndrome.

Observation

It’s about a 17-month-old infant of non-consanguineous parents. He was the second of two siblings (a healthy 4-year-old sister). He was admitted to the pediatric gastroenterology department for moderate undernutrition. He was on exclusive breastfeeding from birth until the age of 2 months before turning to artificial milk. It was diversified at the age of 6 months with an adequate caloric intake. Its symptomatology goes back at the age of 5 months with severe anorexia, weight-loss stagnation and involuntary movements of the two eyeballs. The clinical admission examination showed a conscious, overactive child, a delay in weight with an estimated weight of 6.8 kg (-3DS) and an estimated size of 75 cm (-2DS), while the cranial perimeter was normal at 48cm. Mild to moderate undernutrition with a pasty fold of undernutrition and sign of slump were observed. We observed also a facial dysmorphia with a triangular face, a pointed chin and a bulging forehead evoking the Russell syndrome (Figure 1 & 2). The rest of the somatic examination did not find any signs in favour of neurofibromatosis. Ophthalmologic examination showed an accentuated bilateral rotary nystagmus in the left eye and a unilateral oedema of the left eye in stage I. The malabsorption and hormonal balance were normal. The eso-gastro-duodenal fibroscopy was normal and the jejunal biopsy objectified the absence of villous atrophy, the absence of metaplasia or dysplasia and pathogenic microorganism. Brain magnetic resonance imaging showed the presence of an optic chiasma glioma. In fact, the MRI has shown the presence of a medial suprasellar lesion process lateralized to the left, poorly limited, in T1 hypo signal (Figure 3), T2 hypersignal (Figure 4 & 5) and flair, without translation on the diffusion sequences, enhanced heterogeneously after injection of gadolinium (Figure 6 & 7), measuring 35X37, 5X26cm (TX AP XH).

On the topographic plan:

a) From behind, it fills the inter-pedunde cistern, with discreet mass effect on the left mesencephalic peduncle.
b) Forward, it encompasses the optic chiasm and extends along the path of the left optic nerve without endo-orbital extension.

c) At the top, comes into contact with the 3rd ventricle.

d) At the bottom, she widens the sella turcica with respect for the physiological T1 hypersignal of the post-pituitary gland.

e) Laterally, it includes the middle and anterior cerebral arteries which remain permeable. (Figure 3-7)

The biopsy was not performed on our patient due to the risk of blindness. Regarding the therapeutic issue, the child was “re-typed” nutritionally where he was put on enteral nutrition at constant flow for one month. Due to the achievement of optic chiasm, a chemotherapy combining Carboplatin and Vincristine was established following the LGG protocol. This recommended treatment was accompanied by regular quarterly pediatric, ophthalmological and radiological monitoring in order to look for signs of progression of this pathology. After a 3-month follow-up, the patient presented an improvement in the clinical state marked by an increase in weight estimated at 4 kg and radiological stabilization (image 8) as the cerebral control MRI was in favor of a chiasma glioma optical appearance stable compared to the MRI of 9/30/2019.

Discussion

Glioma of the optic nerve is rare and poorly described in the literature. It represents 5% of childhood brain tumors. Its frequency is less than 5% [3] and its prevalence in the general population is 1 per 100,000 [4]. Most childhood gliomas are sporadic, but in 30 to 50% of cases, they are associated with neurofibromatosis type 1 [5]. These sporadic cases are more aggressive with rapid chiasmatic and retro-chiasmatic expansion whereas cases linked to neurofibromatosis remain confined to the optic nerve and have a slower progression [4,6]. Gliomas of the optic nerve are usually revealed around the age of 2 to 7 years.
Chiasma glioma revealed by blindness. A rare case of optic nerve glioma, a 4-month-old child. J Fr Ophtalmol 31(6 Pt 1): 618-621.

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Conclusion

The diagnosis of optic chiasm glioma in a young infant is rare, especially in case of atypical clinical picture such as diencephalic cachexia, which constitutes a rare mode of revelation that can delay the diagnosis and management until the blindness stage. Chemotherapy is the first-line treatment given the risk of blindness.

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