Dural sinus thrombosis in pediatric age

A trombose do seio dural é uma situação clínica rara, que resulta normalmente da complicação de processos infecciosos dos seios perinasais. Os sintomas e sinais são extremamente variados e inespecíficos sendo o diagnóstico feito através da ressonância magnética nuclear. Esse trabalho relata a ocorrência de trombose do seio dural em um paciente com idade pediátrica. Paciente com 10 anos de idade, sexo masculino foi enviado ao serviço de urgência devido à diplopia e endotropia no olho esquerdo. No exame oftalmológico foi detectado papiledema bilateral, diplopia binocular e endotropia do olho esquerdo. Visual acuity was 10/10 bilateralmente. Given the suspected space occupying lesion of the central nervous system, the MRI was performed and confirmed the diagnosis of DST. For evaluating the intracranial pressure (IP), a lombar puncture (LP) with manometry was carried out and revealed IP of 20 mmHg (normal values: <15mmHg). Towards this, the child’s was admitted for medical treatment (low molecular weight enoxaparin subcutaneous 1.5 mg/kg/day (60 mg/day), prednisolone 35 mg/per day and acetazolamide 250 mg/per day) over 10 days. After 1 month of follow-up there was deterioration of the ophthalmologic condition. A new LP was made and showed IP of 40 mmHg resilient to medical treatment. After multidisciplinary discussion of the case, it was decided for conducting lumboperitoneal shunt. The need for a great deal of suspicion for both the initial diagnosis and for monitoring complications make DST approach a special process.

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INTRODUCTION

Dural sinus thrombosis (DST) comprises the sagittal, transverse and sigmoid sinuses, and in about one third of the cases it comprises multiple sinuses(1,2).

The estimated incidence of DST in pediatric age is approximately 0.67 per 100,000 children per year(3).

TSD typically results from a complication of infectious processes of the paranasal sinuses and the middle ear, but there may be aseptic forms associated to predisposing medical conditions for thromboembolic events such as: haematological disorders, trauma, anemia and dehydration(3-5).

The clinical presentation is nonspecific and may manifest as coma, headaches, vomiting, lethargy with ocular paresis and diplopia, which often leads to a late diagnosis.

The diagnosis is made by nuclear magnetic resonance (NMR) alone or in association with cerebral angiography (6).

This paper reports the case of occurrence of DST in a pediatric patient.

CASE REPORT

A 10-year old male patient is sent to the emergency room due to endothropy of the left eye (LE) and binocular diplopia. As relevant systemic background there was a flu syndrome since 1 month before, and binocular diplopia with a development of about fifteen days.

The ophthalmological exam showed intermittent binocular diplopia, endothropy of the left eye of about 15° (Hirschberg Method) and limitation of abduction in levoversion of the LE. It presented best visual acuity (VA) corrected bilaterally of 10/10. Biomicroscopy without changes. Goldmann applanation tonometry: 12mmHg in both eyes, and in fundoscopic observation: optical discs with undefined boundaries in both eyes by exuberant bilateral papilledema. Macular regions were normal in both eyes (Figure 1).

The decision was to perform a retinal fluorescein angiography which showed late hyperfluorescence with dye leak in both eyes.

An MRI was performed in association with the cerebral angiography (Figure 2) which revealed: exclusion, in the venous study, of all the right sigmoid sinus, the distal portion of the transverse sinus and the proximal segment of the internal jugular vein, which is followed by isosignal on T1 and a discrete hypersignal on the remaining pondersation, and after the use of contrast an enhancement that is predominantly peripheral was observed. The expansion of the optic nerve sheath is also observed. These aspects are suggestive of venous thrombosis.

Given the suspicion of DST, the decision was to request the Neurology to observe, and no neurological disorders were detected. Nevertheless, a lumbar puncture was performed and the result revealed an intracranial pressure (PIC) of 20 mmHg.

Before the clinical profile, the child was hospitalized for medical treatment with low molecular weight enoxaparin 1.5 mg/kg/day subcutaneously (60 mg/day), prednisolone 35 mg/day orally and acetazolamide 250 mg/day orally.

During hospitalization the tests of blood count, biochemistry, microbiology, serology, a study of prothrombotic factors, autoimmunity and a study of the cerebrospinal fluid (CSF) were performed.

The child was discharged after ten days of hospitalization, with no complaints, treated with enoxaparin 60 mg/day and a regressive scheme of oral corticosteroids. The follow-up was performed at the end of 1 month and showed: VA RE: 4/10 and VA LE: 1/10. In the fundoscopy: less exuberant papilledema, but associated with bilateral pallor of the optic disc. The performance of a control NMR showed a profile similar to the previous one. The performance of visual fields showed severe bilateral loss of the visual field, more intense in the LE, with preservation of central islands of vision (Figure 3). Further evaluation has been requested by the Neurology, and a new lumbar puncture was performed with manometry, whose result revealed PIC of 40 mmHg. The clinical case was then discussed with the Neurosurgery which indicated external lumbar drainage with monitoring of the CSF pressure by continuous manometry. Despite the decrease in the PIC, the decision was to have a lumboperitoneal shunt. The child was discharged after 48 hours, with no complaints, PIC between 8-12 mmHg and decreased bilateral papilledema.

After 1 month, the child was asymptomatic and with the following eye exam results: VA RE: 9/10 and VA LE: 5/10. The funduscopy showed an improvement of the edema of the optic disc, however associated to pallor in the RE and optic atrophy in LE (Figure 4). The performance of new visual fields showed am improvement (Figure 5).
DISCUSSION

The DST usually involves the sagittal, transverse and sigmoid sinuses, showing an incidence of 0.67 cases per 100,000 children per year(10).

The clinical presentation is nonspecific, often leading to a late diagnosis or even to its omission. Symptoms can range from headache, decreased VA, diplopia to lethargy, coma and death, having Sébire et al.(7) concluded that headaches, fever and changes of the cranial nerves were the most common form of presentation.

This clinical situation typically results from infectious processes of the paranasal sinuses or prothrombotic conditions, being acquired such as anemia and dehydration, or genetic such as increased factor VIII, increased fibrinogen, decreased proteins C and S, factor V Leiden mutation, homozygosity of thermolabile gene variant of methylenetetrahydrofolate reductase (t-MTHFR). In the same study of Sébire et al.(7), anemia and increased factor VIII were the most commonly found laboratory findings. In about 25% of the cases there is no predisposing factor identified(9).

In the present clinical case, a comprehensive study of infectious causes was performed by serology to screen adenovirus, mycoplasma, pneumonia, cytomegalovirus, herpes simplex 1 and 2, Epstein-Barr virus, and all the results were negative. Blood cultures for anaerobic, aerobic and fungi also presented negative results. The research of prothrombotic risk factors including fibrinogen, protein C and S, anti-thrombin, anti-cardiolipin, lupus anticoagulant, plasminogen, homocysteine, prothrombin, factor V Leiden, homozygosity for t-MTHFR, factor VIII and factor XII have all had negative results.

Electrocardiogram, echocardiogram, carotid Doppler were also performed - all without changes.

Given the results of the supplementary diagnostic exams performed, it was concluded that this is an idiopathic condition, fitting in the 25% of cases without an etiologic diagnosis. It could also be supposed that the flu-like illness occurred a month before has triggered the current situation, but there is no clinical data to prove this relationship. There is some controversy regarding the use of enoxaparin, but several studies have shown that its use is associated with a decreased mortality and an increased recovery rate(9,10).

The diagnosis of DST requires a large initial clinical suspicion, and the confirmation is established through the use of NMR angiography (6). This method allows the identification of the thrombus and the involved sinus completing deficit, currently being considered as the gold standard.

The incidence of intracranial hypertension arises in the work of Sébire et al. in 62% of cases, yet the convulsions and the appearance of new thrombotic events are the most frequent complications(7). As there was no initial response to the medical therapy and there was a worsening of PIC, there was a need for neurosurgical therapy with the placement of lumboperitoneal shunt.

Permanent complications such as blindness and focal neurological deficits are described in about 6-20% of the cases(1,11,12), and a mortality rate between 4.3 and 30%. The prognosis seems to be related to the extent and location of the parenchymal lesion, hemoglobin level, age, and most likely the rapid diagnosis and treatment.
**CONCLUSION**

The dural sinus thrombosis is a rare but fatal clinical situation in some cases. Often the ophthalmologist becomes the key element both in the diagnosis and monitoring of these patients, therefore being important to alert the ophthalmic community to its importance.

**REFERENCES**


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