

Intracranial Migration Of Ventriculo—Peritoneal Shunt Catheter

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Abstract

Shunt migration into the cranium is a rare complication of surgical treatment for hydrocephalus. The authors report a case of shunt migration in a child and discuss common physiopathology and treatment.

INTRODUCTION

The migration of ventriculo—peritoneal shunt catheter is a rare complication of surgical treatment for hydrocephalus. The migration may be in either direction and is more frequent with hard and spring loaded shunt tubes ^{1,2}.

The authors report a case of intracranial dislocation of a ventriculo—peritoneal shunt catheter. The pathophysiological mechanism of the catheter migration is discussed.

CASE REPORT

A male child was born by breech delivery at the 30th week of gestation. Four months later a macrocephalia and a tense fontanel developed.

Computed tomography showed an increase of ventricular volume. The child underwent a ventriculoperitoneal shunt (medium pressure). On the 9th postoperative day, the child was discharged. Some days later, the child returned to João Alves Filho Hospital (Aracaju – Sergipe – Brazil), with signs and symptoms of increased intracranial pressure. Dehydration, macrocephalia, tense fontanel and tumor in right posterior parietal region were present.

Computed tomography showed an increase of ventricular volume, air in right temporal horn and shunt tube in the cranium. (Figure 1). The child underwent surgical removal of the tube and a new tube was implanted in contralateral side. On the 7th postoperative day, the child discharged. Nowadays, the child had been treated with anticonvulsant drugs and ambulatory follow – up.

Figure 1

Figure 1: Computed tomography showing intracranial shunt catheter



DISCUSSION

Shunt complications have been frequently reported in literature. The intracranial migration of ventriculo—peritoneal shunt is the most rare complication and constitutes 0,1% to 0,4% of all shunt procedures ¹. Distal migration of the shunt has often been reported ^{1,2,3,4,11}. The pressure gradient between the cranial and peritoneal cavities decides the direction of migration ^{6,12}. The mechanism of shunt migration involves adhesion, necrosis, penetration, perforation, migration and extrusion ^{6,8,13}.

Two principal causes have been suggested to explain the shunt migration into the cranium: the mechanic force

moving the shunt catheter into the cranium and the low resistance^{6,13}. In childhood, vigorous flexion–extension movements of head may act as a windlass, facilitating upward migration of the shunt catheter¹⁰. Over and above that, the distance between the ventricular and the peritoneal ends of the catheter is smaller than in adults, and proximal migration is easier¹³.

Shimizu et al¹³ reported a case with visual shortage and seizures. Those authors suggested that the cause of migration shunt catheter had been related to stress due to seizures, constipation and osteolysis insulted by craniotomy. Gupta & Mann⁸ reported a case of shunt migration in a child with Dandy Walker cyst. Absence of raised intracranial pressure in this case suggests equilibration of cerebrospinal fluid pressure gradient. The diagnosis was incidental and the patient can be followed up expectantly. Cerrón-Rojas et al⁵ reported a case of simultaneous cephalic migration into the intraventricular and subdural spaces. Those authors concluded that some factors are necessary: such as detachment of the shunt of the distal end (technical fault), underlying disease (porencephaly), dynamic factors causing expulsion (abdominal peristaltic movements), dynamic translocation factor (neck movements), dynamic attraction factor (increased cerebrospinal fluid reabsorption) and unishunt catheter (offering no resistance to passage through the trepanation orifice)⁵. Technical fault is reported such a cause of migration by others authors^{6,8}.

The treatment consists of removing the migrated shunt and implantation of a new shunt, preferably with a reservoir^{1,9}. Migration of shunt is not prevented by locks and slip clips

^{7,8}.

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