Cystotoperitoneal Shunt Surgery during infancy in Porencephalic cyst located in Frontal region led to regaining of developmental milestone

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Abstract: Porencephalic cyst is considered as an extremely uncommon developmental disorder of the central nervous system, being characterized by the presence of a fluid-filled cysts or cavities located within the cerebral hemispheres. It can be associated with varied aetiology and can present with a spectrum of clinical presentation varying from asymptomatic to grossly spastic limbs, mental retardation, cognitive impairment and intractable seizure. Extensive Pubmed and Medline search did not yield any result when searched for term “infancy, porencephalic cyst, cystoperitoneal shunt.” However, clear guideline for management is still lacking. Authors report an interesting case of giant porencephalic cyst located in the right frontal region in infancy and underwent cystoperitoneal shunt surgery, which lead to good outcome with remarkable recovery of delayed milestones with adequate scholastic performances along with marked diminution in the size of porencephalic cyst.

Key words: porencephaly, raised intracranial pressure, cystoperitoneal shunt surgery

Introduction

Porencephalic cyst is considered as an extremely rarer encephalomalatic disorder of the central nervous system characterized by the presence of a fluid-filled cavities or cysts. ¹ It usually represents the end result of various types of the cerebral parenchymal injury, but can also be congenital developmental disorder, or consequences of intracerebral hemorrhage. ²,³ Author reports an interesting case of porencephalic cyst located in the right frontal region, who underwent cystotoperitoneal shunt surgery, led to recovery of regressive milestones and currently child is extremely well scholastically at four years following CSF diversion surgery.
Case illustration

A one-year-old male infant was brought to our neuroscience emergency services with complaint of failure to thrive, delayed developmental milestones with repeated episodes of vomiting and progressive head enlargement for the last six months. On general physical examination at admission, vitals were stable, with bulging anterior fontanelle and head circumference was 54 cm. On neurological examination, he was alert, crying. Evaluation of milestones, he was unable to can sit up without support and not able to crawl and could not use the pincer grasp. He was not able to begin to make recognizable syllables like "ma" and also had regression of language development. Fundi showed bilateral papilloedema. He had bilateral up gaze paresis with presence of sunset sign.

The routine hematological and biochemical parameters were normal limit. The computerized cranial tomography which was carried out after admission (Figure 1) revealed presence of a giant porencephalic cyst in the right frontal region crossing across the midline and causing mass effect obstruction along the CSF pathway leading to dilated lateral ventricles, the third ventricle and the periventricular ooze was present. (Figure 2)

He underwent cysto-peritoneal shunt surgery under general anaesthesia. CSF study revealed normal biochemical parameters with normal cell count and further, CSF culture did not show growth of microorganism. After surgery he regained milestones like crawling, pincer grip and started telling Monosyllabic words over two months after the surgery. (Figure 3)

The computerized tomography carried out at follow-up at four years after shunt surgery revealed well decompressed ventricles (Figure 4) with ventricular catheter well placed, (Figure 5) and marked reduction in the size of porencephalic cyst with well-functioning shunt. Follow-up at three year, he was doing extremely well with developmental milestone appropriate to age and the body weight also appropriate to the age.

Figure 1: Noncontrast cranial computed scan showing large porencephalic cyst

Figure 2: Tomogram showing shut in situ
Discussion

In 1859, Heschl coined the term “porencephaly” to describe presence of a cavity within the human brain. The porencephalic cavities can result from focal cerebral degeneration secondary to haemorrhages. Further de novo or inherited heterozygous mutations in COL4A1 coding the type IV α1 collagen chain, which is essential for vascular basement membranes structural integrity is observed in individuals with porencephaly by Yonder et al. Yoneda et al postulated abnormalities of the α1α1α2 heterotrimers of type IV collagen causes porencephaly. During embryonic life. The large craters develop on the brain surface and gets lined with smooth tissue, filled up with fluid and causes local mass effect and focal neurological deficit and regression of milestones. Usually caused by antenatal or perinatal insult prior to
development of the cerebral gyri due to various pathology i.e. Infective, infarction ischemic or intracerebral haemorrhage.

Depending on the location, size and approximate to the vital region and severity of the defects, symptoms are highly variable and may cause only minor neurological deficit to severe mental retardation and paralysis in extreme cases. The patients with a porencephalic cyst also frequently suffer with seizure disorder.\textsuperscript{[4,7]}

The diagnosis can be also made even during antenatal period with ultrasound and considered as preferred imaging modality during fetal life. In infant, trans-illumination of the skull can also aid in the diagnosis of the porencephalic cyst. Cranial computed tomography reveals location, size; volume, degree of mass effect, associated hydrocephalus and associated developmental pathology and effect of previous treatment and very useful in serial follow-up in assessing the size and evaluating the effect of cystoperitoneal shunt and recurrence of porencephalic cyst.\textsuperscript{[8]} Currently, the magnetic resonance imaging is considered as modalities of choice as it clearly depicts the pathology and associated obstruction along the CSF pathway and associated congenital hydrocephalus, agenesis of corpus callosum or other developmental pathology.

Porencephalic cyst diagnosis requires detailed clinical and neurological evaluation, families history, clinical observations, or based on neuroimaging findings. Neuroimaging establishes the diagnosis and further assessment of intelligence, memory and speech evaluation can further aid in the planning of the holistic management and rehabilitation measures.\textsuperscript{[4]}

Currently, there is no curative therapy for porencephalic cyst and various treatments currently offered included physiotherapy for spastic limbs, rehabilitation, providing antiepileptic medication for those suffering with seizures. Surgical procedure may include cerebrospinal fluid diversion surgery like cystoperitoneal shunt, VP shunt surgery and surgery for epilepsy with intractable epilepsy.\textsuperscript{[1]}

According to the location of porencephalic cyst, extent of the mass effect, size of cavities, and severity of the disorder, combinations of treatment modalities of treatment is usually made tailor made to suit the case.

Prognosis of porencephaly depending on various factors i.e. The location of the cyst and extent of the damage of the brain. Cases with mild neurological deficit can lead normal lives and self-care is possible, however more seriously disabled cases require lifelong support in nursing care and rehabilitation measures.\textsuperscript{[8]} Early diagnosis, medication, participation in rehabilitation activity related to fine-motor control, and communication skill can significantly improve the symptoms and ability to cope with the severe disability and prompt to lead to normal life.

Infants, with proper treatment, can develop good locomotor control as our case, as he was offered surgical treatment at infancy continued antiepileptic medication, led to good recovery of milestones and improvement in cognitive and scholastic performance.
Conclusion
Early diagnosis and management of symptomatic raised intracranial pressure in case of porencephalic cyst, appropriate surgical intervention at early age and antiepileptic medication and rehabilitation measure and regular follow-up cases may aid in not only good neurological recovery and acquiring good motor and cognitive skill and can lead independent life as our cases underwent Cystoperitoneal shunt at one year age.

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