Case Study

Bilateral Subdural Hygroma in a Case with Sylvian Arachnoid Cyst

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Abstract
Arachnoid cysts are congenital lesions arising from the arachnoid membrane. The incidence of arachnoid cysts in the general population is around 0.1% and their most common location is the middle fossa. Arachnoid cysts usually present during the childhood and are identified incidentally. When symptomatic, the most common presentations include headache and seizure. The popular treatment for this lesion is surgery. In this case report, we describe a 12 year-old boy with a middle cranial fossa arachnoid cyst presenting with progressive headache, and bilateral subdural hygroma following a trivial head trauma.

Introduction
Arachnoid cysts are congenital lesions developing within the arachnoid membrane (1). They are non-neoplastic, benign lesions filled with cerebrospinal fluid (CSF) or a fluid similar to it (1). Their most commonly present in the middle fossa and are enclosed by the arachnoid membrane (2,3). Arachnoid cysts are relatively rare. Their incidence in the general population is around 0.1% based on the autopsy studies, with an estimated incidence between 0.5% and 1.6% (4).

Arachnoid cysts most commonly present during childhood and are usually identified incidentally. Almost 75% of intracranial arachnoid cysts were found to present before 3 years of age in one series (5). The most common symptoms include headache and seizures. When there is hydrocephalus or intracranial hypertension the common treatment is surgery (6). Intracranial hypertension secondary to arachnoid cyst rupture especially in the subdural space is a rare clinical entity [7, 8].

In this case report, we describe a 12 year-old boy with bilateral subdural hygroma and a middle cranial fossa arachnoid cyst presenting with progressive headache following a trivial head trauma. To the best of our knowledge this is not yet reported in the literature.

Case Presentation
A 12 year-old boy had a falling down while playing about 40 days before his admission to the hospital. His occiput was injured in the event. He had only dizziness immediately after the trauma for 2-3 minutes with no laceration, bleeding, nausea, vomiting, loss of consciousness (LOC), Seizure or post-traumatic amnesia (PTA), etc.
He had mild, left parietal beating headache which gradually generalized and became associated with nausea and progressive non-bloody vomiting, 3 days after the trauma. Vomiting was mainly postprandial.

He was then referred to the neurosurgery clinic for evaluations. On neurological examination, his optic disk was flat and cranial nerves were normal.

His meningeal signs were positive. Computed tomographic (CT) scan of brain (Figure 1) showed a hypo-dense lesion in left middle cranial fossa partially squeezing the ventricular system without midline shift. Brain magnetic resonance imaging (MRI) showed the lesion in the same place, widening Sylvian fissure, compressing temporal lobe and squeezing lateral ventricles.

The lesion had the similar intensity to cerebrospinal fluid (CSF). There was also fluid collection both in frontal subdural spaces hypo-intense on T1 (Figure 2) and hyper-intense on T2 images (Figure 3). The findings were suggestive of Sylvian fissure arachnoid cyst and bilateral subdural hygroma.

He was subsequently admitted to hospital for craniotomy, fenestration and marsupialization of the cyst into subarachnoid spaces, and evacuation of the hygroma.

Postoperative imaging (Figure 4) showed resolution of the subdural hygroma with small extracranial collection of CSF. He discharged from hospital in good condition with disappearance of all of his complains. In follow-up visit, he did no meningeal signs. His neurologic examination was also normal.
Discussion

Intracranial arachnoid cysts are congenital malformations formed through the splitting of arachnoid membranes due to an increased pulsation of the CSF (9).

They are considered intra-arachnoidal in location and account for 1% of the intracranial mass lesions (10). They are located in the middle cranial fossa in 25-80% of cases. Almost all of these cysts are unilateral, with a slight predilection for the left side (11).

A progressive increase in cyst volume may occur in many cases (12). Intra-cystic hemorrhage, an osmotic gradient allowing a passive fluid-diffusion into the cyst, a ball-valve mechanism or an active secretion from the cyst wall and pulsation of intracystic fluid of venous or arterial origin are some theories to explain the cyst growth (10, 13, 14).

Surgical indications and techniques should be based on the type of arachnoid cyst (communicating or non-communicating), growth of the cyst, compression or displacement of the surrounding neurovascular structures, and most importantly, the patient’s symptomatology and on-going clinical course.

Sixty to ninety percent of arachnoid cysts present during childhood (15, 16). Clinical presentation varies primarily by age and location of the lesion. Some arachnoid cysts remain asymptomatic throughout life.

The most common presenting symptom is headache, which may be due to local mass effect, high intracranial pressure (ICP), or hydrocephalus (17). Arachnoid cysts of the middle cranial fossa may be associated with hemorrhage, however, this complication has been considered very infrequent.

Hemorrhage into or around an arachnoid cyst is primarily precipitated by minor trauma and, very rarely, can be spontaneous. Several reports have been published on the association of arachnoid cysts with subdural hygroma and subdural hematoma after minor head injuries (7,8,16,20).

Arachnoid cysts may cause cognitive disorders and deficits of different aspects of the patient’s psychological profile (18).

Even a trivial trauma can cause rupture of the cyst as seen in the present case, where a simple falling down while playing resulted in the formation of hygroma.

The indication for surgery and the mode of surgical treatment are still a matter of debate. Different types of surgical approaches can be performed. Cyst fenestration, stereotactic puncture, endoscopic cyst fenestration, cystoperitoneal shunt, cyst marsupialization into the subarachnoid space and complete or partial resection of the cyst wall are among the main options (19).

Although the risk of hemorrhage of arachnoid cyst is very low, when the hemorrhage occurs it is treated by hematoma evacuation in most cases. Sometimes, there remains a need for fenestration of the cyst into basal cisterns by endoscopy, microsurgical or cystoperitoneal shunt (20).

In our case, patient’s age, minority of trauma and symptomatology were similar to what reported in the literature. Meanwhile, formation of the subdural hygroma in both sides of the brain where falx cerebri acts as a barrier cannot be explained. Such an

Figure 4. Post-op brain CT scan revealed resolution of bilateral subdural hygroma with small extracranial collection of the CSF
interesting finding in the present case prompted us to report this clinical vignette.

**Conclusion**

In patients with arachnoid cyst it is important to keep in mind that even a trivial event may result in a significant intracranial complication, potentially in both hemispheres.

**References**

20. meshkini A, Meshkini M. Two case series reports: 8 cases of Arachnoid Temporoparietal cysts (Middle Fossa & Sylvian fissure) and 2 cases of chronic subdural hematoma: Injury & violence journal. 2012;4(51)