Hydatid cyst of the quadrigeminal cistern. A case report for unusual location with literature review

Saif Saood Abdelrazaq¹, Abdullah H. Al Ramadan², Ali Adnan Dolachee³, Mohammed Maan AbdulAzeez⁴, Ali Saud Abdulrazzaq⁵, Amar Saeed Rashid⁶, Samer S. Hoz⁵

¹ Neurosurgery Department, Al Sadar Teaching Hospital, Najaf, IRAQ
² Department of Neurosurgery, King Faisal Specialist Hospital and Research Centre, Riyadh, SAUDI ARABIA
³ Department of Neurosurgery, College of Medicine, Al Qadisiyah University, IRAQ
⁴ College of Medicine, University of Baghdad, Baghdad, IRAQ
⁵ Neurosurgery Department, Neurosurgery Teaching Hospital, Baghdad, IRAQ
⁶ Department of Neurosurgery, College of Medicine, Kufa University, Baghdad, IRAQ

ABSTRACT
Intracranial hydatid cyst involves supratentorial area and mainly affecting the middle cerebral artery territory with the predilection of the partial lobe. It can be single - which is the most common - or multiple up to 35 cysts. They tend to be huge at the time of symptomatic presentation especially when they are presented as a solitary lesion with a slow growth rate around 1.5 cm/year, however, it is variable and it can be up to 10 cm/year. Surgical treatment is mandatory for all patients once the correct diagnosis is made, except for patients with multiple organ involvement in poor general conditions and deep-located cysts. The existence of hydatidosis in the cisternal spaces must not be neglected given the capacity of E. granulosus larvae to disseminate via the CSF. In this case report; two and half years’ male child presented with a history of 2 attacks of generalized seizure for the last 72 hours with the head circumference at the upper normal limit for his age. This paper presents the first case report demonstrating a primary single hydatid cyst located in the quadrigeminal cistern in a child. We concluded that in spite of the feasibility of the imaging and the high suspension of cerebral hydatid cyst, still, the reports show more locations which can be described as unusual although for a head to toe suspected distribution of hydatid disease is already understood. An eminent medical and surgical (if indicated) treatment of the primary cerebral hydatid cyst are always effective and recommended.

INTRODUCTION
Choroid Echinococcosis could be characterized as a zoonotic infestation caused by cestode species of the genus Echinococcus (1).
It is most commonly involving the liver then lungs. (2) Intracranial involvement occurs in 1-2% of Echinococcosis and 50-70% of these cases affect the paediatric age group. (3) It showed slightly male preference in most of the case series. (4,5,6,7,8) Geographically endemic area of hydatid disease includes some of South America countries like Argentina, Brazil, Chile, Peru and Uruguay, France, Italy and Greece in Europe, Turkey, Israel, Lebanon, Syria and Jordan in the Middle East, China, Iran, and India in Southeast Asia and Tunisia in Africa. (9)

Intracranial hydatid cyst involves supratentorial area and mainly affecting the middle cerebral artery territory with the predilection of the partial lobe. It can be single - which is the most common - or multiple up to 35 cysts. (10,11,12) They tend to be huge at the time of symptomatic presentation especially when they are presented as a solitary lesion with a slow growth rate around 1.5 cm/year, however, it is variable and it can be up to 10 cm/year. (13) Surgical treatment is mandatory for all patients once the correct diagnosis is made, except for patients with multiple organ involvement in poor general conditions and deep-located cysts. (14) The existence of hydatidosis in the cisternal spaces must not be neglected given the capacity of E. granulosus larvae to disseminate via the CSF. (15)

This paper presents the first case report demonstrating a primary single hydatid cyst located in the quadrigeminal cistern in a child.

**CASE SCENARIO**

Two and half years’ male child presented with history of 2 attacks of generalized seizure for the last 72 hours with head circumference at the upper normal limit for his age.

Initial lab workup was done to exclude metabolic derangements the patient was sent for neuroimaging. The general lab tests were within normal ranges and the brain MRI showed a midline rounded cystic lesion, 5x3 cm in size, located in extra-axially in the quadrigeminal cistern and extend infra-tentorially to compress the cerebellum downward along with bilateral dilated lateral ventricles (Figure 1).

**FIGURE 1.** Pre-operative brain MRI showing a midline rounded cystic lesion, 5x3 cm in size, located in extra-axially in the quadrigeminal cistern and extend infra-tentorially to compress the cerebellum downward along with bilateral dilated lateral ventricles. The lesion was hypo-intense and homogenous in T1 with intensity higher than CSF, hyper-intense in T2 and showed no enhancement in contrasted image, also there was some vivid inclusions within the upper part of the cyst.

**FIGURE 2.** the delivered intact hydatid cyst.
The lesion was hypo-intense and homogenous in the T1 sequence of the MRI with intensity higher than the CSF, hyperintense in the T2, and showed no enhancement in the contrasted image, also there were some vivid inclusions within the upper part of the cyst.

At that point, the top differential diagnosis included were epidermoid, dermoid and arachnoid cysts only, because that location was an unusual location for a hydatid cyst and the age of the patient was too young for a hydatid cyst.

The decision was to do a surgical resection or fenestration according to the intra-operative findings. we planned to delay the CSF shunting option as the sulci were not effaced, the young age of the patient and the surgery would remove the cause of the obstruction of the CSF pathways.

The surgery conducted at the next day, an intratentorial supra-cerebellar approach was done via midline occipital craniotomy. The cyst was a typical hydatid cyst. The cyst resected as one piece via initial dissection and then delivered using Dowling's hydrodissection technique (Figure 2,3).

The postoperative course went uneventfully. The histopathology confirms the diagnosis, further workup excluded other organs involvement.

![Figure 3. Early postoperative axial brain CT scan showing resection cavity](image)

<table>
<thead>
<tr>
<th>No.</th>
<th>Authors</th>
<th>Age/Gender</th>
<th>Complain</th>
<th>Diameter</th>
<th>Location</th>
<th>Treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Rumboldt et al (27)</td>
<td>50 Year/Male</td>
<td>Headache</td>
<td>30 mm</td>
<td>Premedullary Cistern</td>
<td>Surgery</td>
</tr>
<tr>
<td>2</td>
<td>Beskonakli et al (28)</td>
<td>33 Year/Male</td>
<td>Headache and impairment</td>
<td>N/A</td>
<td>Interpeduncular Cistern</td>
<td>Surgery</td>
</tr>
<tr>
<td>3</td>
<td>Sanli et al (29)</td>
<td>7 Year/Male</td>
<td>Symptoms of high Intracranial pressure</td>
<td>25 x 25 mm</td>
<td>Ambient Cistern</td>
<td>Surgery</td>
</tr>
<tr>
<td>4</td>
<td>Kizilca et al (37)</td>
<td>45 Female</td>
<td>Symptoms of high Intracranial pressure</td>
<td>N/A</td>
<td>Premedullary Cistern</td>
<td>Surgery</td>
</tr>
<tr>
<td>5</td>
<td>The authors of this case report</td>
<td>2.5 Year/Male</td>
<td>Seizure</td>
<td>50 x 30 mm</td>
<td>Quadrigeminal Cistern</td>
<td>Surgery</td>
</tr>
</tbody>
</table>
Hydatid cyst of the quadrigeminal cistern

**DISCUSSION**

Variant locations of hydatid cyst were reported in the literature including aqueduct of Sylvius (16), Frontal lobe associated with massive edema and raise in the intracranial pressure (17), temporal lobe with midbrain herniation. (18)

Thalamic hydatid cyst was managed surgically through the transcallosal approach in a child who presented with left-sided weakness. (19)

An interosseous lesion involving the skull estimated to represent 3-4% of the lesion involving the skeleton which evident in 2% of all hydatid cyst. (20)

Spinal hydatid cysts are estimated at less than 1%, they are usually extradural and tend to rupture intraoperatively more than other location. (21) Meckel's cave can be involved with extension to the Cerebellopontine angle and the middle fossa. (22)

Brainstem hydatid cyst was reported in middle age woman presented with a headache and progressive hemiparesis. (23) both cases of Meckel's cave and Brainstem hydatid cysts were treated with decompression first prior to the removal to decrease the chances of intraoperative rupture. (22,23)

Cerebellar lesion with mass effect causing obstructive hydrocephalus was encountered in a child with a progressive headache followed by signs and symptoms of high intracranial pressure. (24)

Intraventricular involvement was estimated between 1.3 - 16.6% of multiple case series for intracranial hydatid disease with overall good outcome after surgical excision. (25) Other unusual locations include the sellar and the para-sellar regions were reported. (26)

Cisternal hydatid cysts were reported to involve the pre-midullary, the interpeduncular and the ambient cisterns. (27,28,29) (Table 1)

An intermittent headache can be the only presenting symptoms even with a huge cyst. (30) In a Large series from China, headache and vomiting were the most common symptoms, this finding was supported in the literature as well.

Seizure attacks, focal neurological deficits, and skull deformities were reported. Papilledema was frequent in the ophthalmological examination especially in patients with high intracranial pressure, optic atrophy was seen as well. (4,31,32)

The Magnetic Resonance Imaging (MRI) thought to be more sensitive than the computed tomography (CT) to identify the pericystic layer and to differentiated the hydatid cyst from the other similar cystic lesions like the epidermoid cyst, however, CT scan is more sensitive to identify the calcified lesions. (33)

Serious complications related to the rupture of the hydatid cyst include recurrence, cerebral vasospasm and infarction, hydrocephalus, anaphylactic shock, chemical meningitis, and even death. (34)

In two series of long-term follow up for cerebral hydatidosis in children, it can be concluded that the improvements of the imaging facilities lead to early recognition and improve the outcome. Poor prognostic factors can be identified including delay in the treatment, the rupture of the cyst at the presentation or intraoperatively, and poor control of the systemic disease. Single versus multiple cyst and age at the presentation was not identified as prognostic factors, however, some authors consider the multiplicity as a malignant form of the disease. (35,36)

Review of all the available literature reveals that our case is the first reported case of hydatid cyst located in the quadrigeminal cistern which was removed successfully.

**CONCLUSION**

In spite of the feasibility of the imaging and the high suspension of cerebral hydatid cyst, still, the reports show more locations which can be described as unusual although for a head to toe suspected distribution of hydatid disease is already understood. An eminent medical and surgical (if indicated) treatment of the primary cerebral hydatid cyst are always effective and recommended.

**ABBREVIATIONS**

CT: computed tomography
MRI: magnetic resonance imaging
CSF: cerebrospinal fluid

**REFERENCES**


