CASE REPORT

Polymorphic Clinical Presentation of Cerebellar Arachnoid Cyst: A Case Report. Etiopathogenetic Theory and Review of the Literature

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Arachnoid cysts (AC) are developmental collections of cerebrospinal fluid covered by layers of arachnoidal epithelium and are usually located in the middle cranial fossa. Localizations in the posterior fossa are uncommon and generally remain asymptomatic or cause vague and non-specific symptoms. We describe the unusual case of a patient with an AC suffering from recurrent polymorphic vertigo with atypical nistagmus and transient sensorineural hearing loss. The clinical-diagnostic features are discussed, etiopathogenic theories are proposed and a review of the literature on posterior cranial fossa AC is reported.

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Introduction

Arachnoid (AC) or leptomeningeal cysts are benign, intra-arachnoid cystic lesions that are filled with cerebrospinal fluid (CSF). Likely developmental in origin and generally asymptomatic, these lesions can become symptomatic as a result of enlargement or intracystic hemorrhage\(^1\). Although their exact etiology is unknown, various hypotheses have been considered. These hypotheses include active fluid secretion from the cystic wall, fluid accumulation secondary to changes in the osmotic gradient, and vascular pulsation causing CSF flow between the cyst and subarachnoid space\(^1\). AC occur throughout the central nervous system (CNS), and while generally no communication is demonstrable with the subarachnoid space, a more recent hypothesis has been suggested based on intraoperative reports of apparent one-way, slit-valve communications between the cyst and subarachnoid space. AC represent approximately 1% of all intracranial space-occupying lesions. They are typically located in the middle cranial fossa, but other locations including the cerebellopontine angle, cerebellar hemispheres and posterior fossa have been described.\(^2,4\)

They may be associated with other clinical situations, such as a carotid-jugular fistula, dissection of the contralateral carotid artery and hemifacial atrophy and thus represent a generalized connective tissue disorder\(^5\).

AC may remain asymptomatic and undiagnosed until the compression of adjacent anatomical structures caused by the growth of the cyst results in various clinical symptoms that are dependent on the location of the cyst. When an AC is located in the posterior fossa, the presenting symptoms are frequently otologic.\(^6\)
We here report the case of a patient with a cerebellar AC suffering from recurrent polymorphic vertigo with atypical nystagmus and transient sensorineural hearing loss. We describe the clinical-diagnostic features together with a review of the literature on such a topic.

**Case Report**

This 73-year-old woman, who had been in good health previously, presented to the Department of Otolaryngology (tertiary referral center) of the Department of Otolaryngology, University of Perugia, with a 2-day history of positional vertigo attacks associated with neurovegetative symptoms and dizziness. She did not report any aural fullness or tinnitus.

Current medications included oral hypoglycemic agents for type II diabetes mellitus and amlodipine for hypertension. She did not report any previous vertigo attack or dizziness, migraine or other neurological symptoms. Pure tone audiometry revealed only a mild, symmetrical, high-frequency, sloping, sensorineural hearing loss, compatible with presbycusis.

Vestibular examination confirmed a vestibular paroxysmal positional vertigo (BPPV) for a canalolithiasis of the posterior semicircular canal. Semont’s liberatory maneuver was performed with immediate resolution of patient’s symptoms. After 2 weeks of complete health, the patient presented a new BPPV attack, which was successfully treated with liberatory maneuvers.

Two weeks later the patient presented at our Department with substantially different complaints: objective vertigo unrelated to head position, associated with hearing symptoms. Bedside vestibular examination showed rotatory horizontal right-beating nystagmus unrelated to head movements, slight left-side index deviation and normal cerebellar vestibular tests. Audiometry showed a low-medium frequency, climbing-shape, sensorineural hearing loss on the right ear. Osmotic diuretics were administered with complete resolution of patient’s vestibular symptoms after one day of treatment. In particular, vestibular examination (including head-shaking and caloric tests) was normal and a resolution of right ear hearing loss was noticed. Such presentation was suggestive of a newly onset Meniere-like attack. Computerized tomography showed a posterior cranial fossa AC, occupying the median-right paramedian cerebellum, associated with hypoplasia of the right cerebellar lobe and vermis, without any focal lesion or recent bleeding image (Figure 1). Such findings were confirmed by magnetic resonance imaging (Figures 2a, b). During the next month the patient presented 2 more BPPV episodes and 1 more hydropic attack, with residual dizziness and low tolerance to head movements and optokinetic stimuli. She also developed a persistent, spontaneous, down-beating nystagmus. In agreement with our neurologist, we considered such nystagmus related to the AC. A neurosurgical consultation was carried out, which however excluded any surgical approach in relation to patient’s age and absence of endocranial hypertension symptoms.

**Discussion**

Arachnoid cysts, firstly described by Bright in 1831, are fluid-filled spaces generated by congenital anomalous splitting of the arachnoid membrane; they may communicate with the subarachnoid space or may be partially or completely separated from it.\(^{1,8}\) AC may form in any position within the skull, more commonly in the left hemisphere, in the middle and
anterior cranial fossa; in a lower percentage of cases (from 20 to 30 per cent) they are located in the posterior cranial fossa, where they may occupy a retrocerebellar area, or grow in the cerebello-pontine angle.\[9\]

The male/female ratio ranges from 2/1 to 3/1 in the different reports and average duration of symptoms before the definitive diagnosis may vary from six months to five years.\[10\]

Signs and symptoms related to the presence of AC depend on their size and location: while supratentorial cysts may produce mild untreatable headache, difficulties in concentration, more rarely temporal lobe epilepsy and, in paediatric patients, bulging of parietal bone, posterior fossa AC tend to have a protean clinical manifestation, with hypoacusis, tinnitus, dysmetria, ataxia, together with the cerebello-pontine angle syndrome, and sometimes vomiting, headache and papillar edema for endocranial hypertension secondary to fourth ventricle obstruction.\[8\]

Interestingly, vestibular symptoms (vertigo and dizziness) predominate over cochlear symptoms (tinnitus and hearing loss) and nystagmus is by far the most common clinical sign.\[8\]

Atypical presentation of AC, especially in case of posterior fossa location, have also been reported (Table 1)\[11-24\].

The case we present is didactic because of the polymorphic atypical symptoms presented by the patient: firstly, BPPV attacks resolved with liberatory maneuvers; secondly, a Meniere-like attack (objective vertigo with rotatory horizontal right-beating nystagmus unrelated to head movements, sand right sensorineural hearing loss) responding to osmotic diuretics; finally, a persistent, spontaneous, down-beating nystagmus, with BPPV and Meniere-like episodes. Since no surgical procedure was carried out, it is not possible to have a definite confirmation of a cause-effect relation between patient’s clinical presentation and her posterior cranial fossa AC. However, AC polymorphic clinical presentation reported in the literature suggests a role of AC in the development of our patient’s signs and symptoms. The mechanisms proposed to explain the atypical clinical presentation of the AC are the generation of a neurovascular conflict or an impairment of ‘vasa nervorum’ due to the stretching of the involved cranial nerves. This may determine an alteration in the normal endolymphatic circulation and an impaired terminal vascularisation of the macular neuroepithelium with detachment of otoconial debris that could explain respectively the relapsing episodes of Menieriform and positional vertigo. Down-beating nystagmus may be secondary to AC-related vermian and cerebellar lobe hypoplasia with consequent disinhibition of the vestibulo-ocular reflex.
Table 1. Review of the literature on arachnoid cyst presenting otologic symptoms

<table>
<thead>
<tr>
<th>Author (year)</th>
<th>Arachnoid cyst location</th>
<th>Clinical presentation</th>
<th>Proposed pathophysiology</th>
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</thead>
<tbody>
<tr>
<td>Jallo et al (1997) [11]</td>
<td>Cerebellopontine angle (5 patients)</td>
<td>Refractory headaches associated with nausea and vomiting (3 cases); cerebellar signs (2 cases)</td>
<td>Cyst-related compression on neurovascular structures of the cerebellopontine angle</td>
</tr>
<tr>
<td>Babu et al (1991) [12]</td>
<td>Cerebellopontine angle (1 patient)</td>
<td>Contralateral trigeminal neuralgia; symptoms resolution after cyst excision</td>
<td>Stretching of brainstem and cranial nerves</td>
</tr>
<tr>
<td>Sakata et al (1987) [14]</td>
<td>Vermis (20 patients)</td>
<td>Positioning nystagmus with augmented latency and reduced fatigability</td>
<td>Incomplete inhibition of the vestibulo-ocular system including the cerebellar flocculonodular lobe or vestibulo-cerebellum</td>
</tr>
<tr>
<td>O’reilly et al (2003) [4]</td>
<td>Posterior fossa (2 patients)</td>
<td>Aural fullness, tinnitus, vertigo, nausea and emesis (Meniere-like syndrome); symptoms resolution in 1 patient after cysto-peritoneal shunt.</td>
<td>Cyst-related vascular compromise, altered cerebro-spinal fluid dynamics, displacement of the VIII nerve, or compression of the endolympathic sac</td>
</tr>
<tr>
<td>Haberkamp et al (1990) [6]</td>
<td>Posterior cranial fossa (1 case)</td>
<td>Fluctuating sensorineural hearing loss (1 case); vague dizziness and ataxia with no vertigo (1 case)</td>
<td>Cyst-related compressive effect on neurological structures</td>
</tr>
<tr>
<td>Police et al (1997) [15]</td>
<td>Posterior cranial fossa (1 case)</td>
<td>Bilateral progressive sensorineural hearing loss and episodic vertigo; the hearing loss was more pronounced in the right than the left ear; bilateral vestibular weakness was found with a &quot;severe&quot; right-sided loss.</td>
<td>Cyst-related compression of neurological structures</td>
</tr>
<tr>
<td>Samii et al (1999) [16]</td>
<td>Cerebellopontine angle (12 patients)</td>
<td>Vertigo or dizziness (8 cases), tinnitus (5 cases), hearing loss (6 cases); most symptoms resolution after cyst surgical removal</td>
<td>Cyst-related compression on neurovascular structures of the cerebellopontine angle</td>
</tr>
<tr>
<td>Pappas et al (1981) [17]</td>
<td>Internal auditory canal (3 cases), cerebellum (1 case)</td>
<td>Progressive right-sided hearing loss, tinnitus, and experiences of dizziness; diminished caloric response in the right ear</td>
<td>Compression on neurovascular structures</td>
</tr>
<tr>
<td>Hadley et al (1985) [18]</td>
<td>Posterior cranial fossa (5 cases)</td>
<td>Tinnitus, lightheadedness, incoordination, and occasional vertigo with normal audiometry</td>
<td>Compression on neurovascular structures</td>
</tr>
<tr>
<td>Erdinger et al (1999) [19]</td>
<td>Posterior cranial fossa (12 patients)</td>
<td>Vertigo, hypoacusis and gait instability (all cases) resolved after surgical treatment; V and VIII nerve palsies (1 case)</td>
<td>Compression on neurovascular structures</td>
</tr>
<tr>
<td>Gosepath et al (1994) [20]</td>
<td>Posterior cerebellum (1 case)</td>
<td>Episodic vertigo with nausea, emesis, left-sided hearing loss, and bilateral tinnitus</td>
<td>Compression on neurological structures</td>
</tr>
<tr>
<td>Chan et al (1991) [21]</td>
<td>Cerebellar vermis</td>
<td>Dizziness and intermittent vertical oscillopsia after oblique extension of the head and neck, intermittent downbeat nystagmus lasting about 50 seconds and elicited by head extension and rotation; nystagmus resolution after cyst removal</td>
<td>Obstructive hydrocephalus</td>
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<td>Aggouri et al (2010) [22]</td>
<td>Posterior fossa</td>
<td>Two- month history of headaches, nausea, and vertigo associated with walking instability; symptoms resolution after cyst removal.</td>
<td>Compression on neurological structures</td>
</tr>
<tr>
<td>Ottaviani et al (2002) [23]</td>
<td>Cerebellar convexity</td>
<td>Left-sided hypoacusia and tinnitus</td>
<td>Compression on the cerebellopontine angle</td>
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</table>
In conclusion, posterior fossa arachnoid cysts must be considered in the differential diagnosis of patients presenting with atypical audiovestibular or Meniere-like signs and symptoms. Neuroimaging (magnetic resonance and computerized tomography) is mandatory for AC diagnosis.

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References


