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Cerebellopontine angle pilocytic astrocytoma mimicking acoustic schwannoma

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Abstract We describe a case of pilocytic astrocytoma of the cerebellum mimicking an acoustic schwannoma. The tumour protruded into the porus acusticus and enlarged the internal auditory meatus, which is a quite unusual characteristic of glial tumours.

Key words Angle, cerebellopontine · Astrocytoma · Schwannoma, acoustic

Introduction

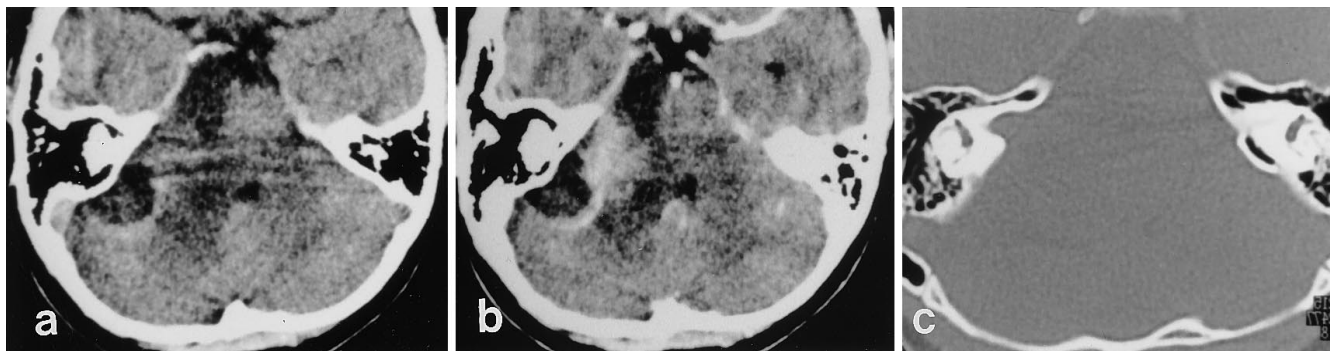
Juvenile pilocytic astrocytomas of the cerebellum, usually originating in the cerebellar hemisphere and vermis, rarely occur in the cerebellopontine angle (CPA) [1]. A few CPA glial tumours causing internal auditory meatus (IAM) enlargement have been re-

ported [2–5]. We present a patient with a CPA pilocytic astrocytoma showing IAM enlargement.

Case report

An 8-year-old girl with a 1-year history of hearing difficulty presented with right mild facial palsy and horizontal nystagmus. CT demonstrated a CPA tumour and enlargement of the right IAM (Fig. 1). MRI revealed a tumour surrounded by multiple cysts in the right CPA. It enhanced heterogeneously with Gd-DTPA, and had a lateral portion protruding into the IAM (Fig. 2). Pure-tone audiometry showed sensorineural hearing loss in the right ear and

Fig. 1 a, b CT before and after contrast medium demonstrates a cerebellopontine angle tumor. **c** A bone window reveals enlargement of right internal auditory meatus



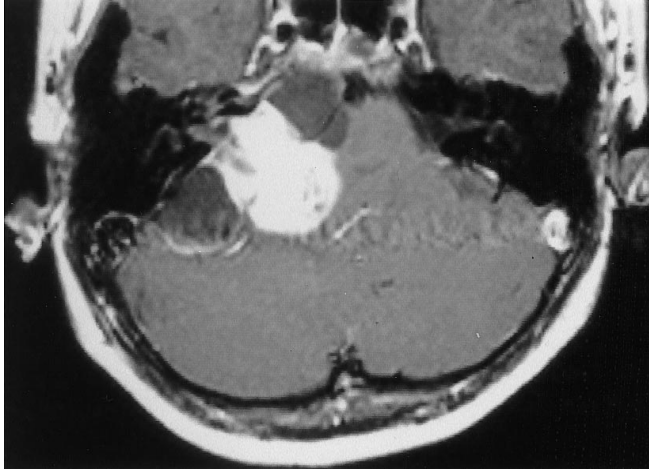


Fig. 2 Contrast-enhanced T1-weighted axial image shows a cerebellopontine angle mass extending into the widened internal auditory meatus

auditory brain stem response studies found no response to right ear stimulation.

The tumor, preoperatively diagnosed as an acoustic schwannoma, was resected through a suboccipital craniectomy. The pathological diagnosis was pilocytic astrocytoma.

Discussion

Pilocytic astrocytomas of the cerebellum are relatively common in children and young adults. They usually originate in the cerebellar hemisphere and often affect the vermis. Although they may mimic an acoustic schwannoma when they grow eccentrically into the CPA [1], neuroradiological differential diagnosis is not difficult.

In our case, CT using a bone algorithm revealed enlargement of the IAM. Bony changes of the IAM are frequent occur with acoustic schwannoma, but rare with CPA glial tumours [3–5]. To our knowledge, only one case of CPA pilocytic astrocytoma with IAM enlargement has been reported [2]. The clinical course and neuro-otological testing were also indicative of acoustic schwannoma. Although acoustic schwannoma is rare in children [6, 7], it should be added to the list of differential diagnostic possibilities.

In this case, the lateral part of the tumour protruded into the anterior part of the widened IAM and a small eccentric cyst, thought to be entrapped cerebrospinal fluid, filled the rest of it. These findings, not characteristic of acoustic schwannomas, may indicate the possibility of a glial tumour.

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