Solitary Osteochondroma of the Skull Base: A Case Report and Literature Review

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Abstract
We report a case of an osteochondroma in the posterior clinoid process that occurred in a 43-year-old man with trochlear nerve palsy. Although the potential preoperative diagnoses based on computed tomography and magnetic resonance imaging included other intracranial tumors such as calcified meningioma, thallium-201 single-photon emission computed tomography effectively differentiated osteochondroma from those possibilities. Via an orbitozygomatic approach, a subtotal resection was achieved with a good relief of symptoms. Twenty-two cases of solitary osteochondromas in the skull base have been reported that have demonstrated little risk of recurrence or malignant transformation. However, surgery for skull base osteochondromas does carry a significant risk with a reported mortality > 10%. Although some previous reports advocate complete resection as the only curative method for skull base osteochondromas, the risks of total resection should be weighed against the chance for recurrence; our review of the literature demonstrated a relatively high mortality and an extremely low incidence of recurrence.

Keywords
► abducens nerve palsy
► bone tumor
► osteochondroma
► posterior clinoid
► skull base

Introduction
An osteochondroma is a benign tumor composed of mature hyaline cartilage with focal ossification.1 It constitutes 10 to 15% of all bone tumors and 20 to 50% of benign bone tumors2 and is frequently found at the epiphysis of long bones such as the distal femur, proximal tibia, and proximal humerus. However, intracranial osteochondromas are relatively rare, and the literature consists only of sporadic case reports.1,3–23
We report a rare case of an intracranial osteochondroma of the posterior clinoid process that caused right abducens nerve palsy along with a review of the literature.

Case Report
A 43-year-old man with no past medical history developed diplopia 3 years ago. He consulted an ophthalmologist, who found no ophthalmologic disease. Because the symptom gradually subsided over several months, he was treated conservatively. Approximately 18 months after the onset, his diplopia recurred; this time, it did not subside and gradually deteriorated. He developed a continuous medial excursion of the right eye. He visited a general hospital, where he was diagnosed with right abducens nerve palsy. Magnetic resonance (MR) images obtained at the hospital revealed an intracranial mass in the right skull base region, and he was referred to our hospital. A detailed neurologic examination on the initial visit revealed complete abducens nerve palsy with no other cranial nerve deficit. He had no muscle weakness or sensory disturbance. A noncontrast computed tomography scan demonstrated a 3.0 × 2.5 × 2.5 cm well-circumscribed extra-axial mass lesion in the right posterior clinoid that contained numerous irregular calcified spots (►Fig. 1A, B). The lesion was located in the right posterior clinoid process of
the sphenoid bone and caused the destruction of the petrous apex of the temporal bone. MR imaging showed that the lesion had heterogeneous signals of intermediate to low intensity on the T1-weighted images (►Fig. 1C) and high intensity on the T2-weighted images (►Fig. 1D). The T1-weighted images with gadolinium enhancement showed strong enhancement of the noncalcified areas (►Fig. 1E). The dural tail sign was not apparent, and there was no perifocal edema. Cerebral angiography showed no tumor stain or arterial stenosis. Thallium-201 single-photon emission computed tomography (SPECT) revealed an extremely low uptake for meningothelial tumors. The preoperative differential diagnosis included meningioma, cavernous hemangioma with intralesional calcification, miscellaneous osteomatous lesions, sarcoma, fibrous dysplasia, and metastasis.

To relieve the abducens nerve palsy and make a definitive diagnosis, a surgical resection via right orbitozygomatic craniotomy was performed while monitoring the extraocular muscles. The right optic nerve, right internal carotid artery, and medial right incisura of the tentorium were exposed by opening the proximal sylvian fissure. The surface of tentorium bulged upward significantly (►Fig. 2A). The tentorium was cut, and the tumor was readily identified; it mildly adhered to the oculomotor nerve. Because the tumor was composed of white brittle tissue with multiple small calcified masses, we removed it piecemeal (►Fig. 2B). There was a substantial amount of bleeding from the cavernous sinus, but it was controlled with a small piece of Gelfoam. Intraoperative frozen sections were unavailable because the specimen was essentially calcified; nevertheless, we were convinced the lesion was benign rather than malignant from our intraoperative findings. Therefore, we intentionally left the thin tumor capsule to prevent the leakage of blood from the cavernous sinus to the prepontine cistern. Although the trochlear and abducens nerves were not found inside the tumor and the oculomotor nerve was preserved anatomically, we were unable to detect extraocular muscle movement during the intraoperative monitoring probably because of compression to the nerves by the surgical manipulation.

Postoperative MR images confirmed near total resection (►Fig. 2C). The pathologic findings demonstrated that the mass was macroscopically composed of bone and white substantial tissue like cartilage. On microscopic findings, there was bony tissue including bone marrow, trabecular bone, and fragments of hyaline cartilage comprising clusters of benign mature and degenerating chondrocytes with foci of irregular calcification and ossification (►Fig. 2D). The bone marrow was occupied by adipose tissue and fibrosis. These findings were consistent with the characteristics of osteochondroma.

Right oculomotor nerve palsy was newly observed after surgery, and the palsy completely resolved in 4 months. Although the abducens nerve palsy had persisted for > 18 months preoperatively, there was evidence of an almost full recovery at the 12-month follow-up after surgery.

Discussion
Osteochondromas are benign tumors composed of mature hyaline cartilage with focal ossification. It is the most common tumor of the bone arising in any part of the body, but it is rare in the skull. We conducted a comprehensive literature search using the online database and identified 22 cases of osteochondromas of the skull including our case (►Table 1). Nineteen of the 22 patients (86.4%) were men. Eleven cases (50.0%) arose in the skull base, whereas the
Others were found in the convexity and the falx. Three cases including ours occurred in the posterior clinoid. As shown in Table 1, ~80% of the osteochondromas in the skull base presented with focal cranial nerve deficits depending on the regions affected by the tumor, whereas non–skull base osteochondromas frequently caused epilepsy and headache as an initial symptom. Apart from visual disturbances, abducens nerve palsy was the most common cranial nerve deficit among skull base osteochondromas. This is probably related to the origin of the tumor cells, the cartilage rests in the synchondrosis. As shown in Table 1, skull base osteochondromas frequently arise in the parasellar region including the posterior clinoid and the middle cranial fossa, which is close to the confluence where the sphenopetrosal, spheno-occipital, and petro-occipital synchondrosis meet. We therefore speculate that the abducens nerve palsy is most frequently observed in skull base osteochondromas because the nerve runs closer to these synchondroses than other cranial nerves.

Some previous reports mentioned that osteochondroma can mimic meningioma. Osteochondromas are usually heterogeneously enhanced on T1-weighted images with gadolinium, indicating that contrast-enhanced MR images are not useful in completely discerning osteochondroma from meningiomas. Angiography has a diagnostic value because osteochondroma appears as avascular masses. In our experience, the findings of an extremely low uptake on thallium-201 SPECT were useful in differentiating osteochondroma from other intracranial tumors such as meningioma.

As for treatment, complete surgical removal is considered the only curative treatment for extracranial osteochondromas. Venkata et al mentioned that skull base osteochondromas may recur from incomplete resection. However, although gross total resection was achieved in only 11 of 23 cases (47.8%), only two recurrent cases (8.7%) have been reported in the literature. In addition, in our literature search, we did not find any reports of malignant transformation of solitary intracranial osteochondroma except for one case reported in 1935. Based on these facts, skull base osteochondromas are basically benign lesions, especially for solitary lesions; the risks of radical resection should be carefully weighed against the future risk of recurrence or malignant transformation. When a small asymptomatic bony lesion showed similar radiologic characteristics, a conservative management strategy could be used. Skull base osteochondromas are often located close to critical structures such as the internal carotid artery, the cavernous sinus, and the cranial nerves. Given that the mortality after surgery occurred in three patients (13.6%) in the literature, subtotal or even partial resections can be an important option to relieve the symptoms and achieve good long-term tumor control.

![Fig. 2](A) Intraoperative photograph showing the tumor covered by the tentorium. (B) The tumor was composed of brittle small calcified particles and hemorrhagic soft tumor components. (C) Postoperative T1-weighted magnetic resonance image with gadolinium demonstrating subtotal resection of the tumor. (D) Photomicrograph showing that the tumor consisted of bony tissue including bone marrow, trabecular bone, and benign hyaline cartilage. Original magnification: ×100.
### Table 1: Summary of cases of intracranial osteochondroma

<table>
<thead>
<tr>
<th>Authors, Year</th>
<th>Age, Sex</th>
<th>Location</th>
<th>Size (cm)</th>
<th>Symptoms, cranial nerve deficits</th>
<th>Treatment</th>
<th>Improvement of symptoms</th>
<th>Follow-up (months)</th>
<th>Recurrence</th>
</tr>
</thead>
<tbody>
<tr>
<td>Alpers, 1935</td>
<td>49, M</td>
<td>Convexity</td>
<td>ND</td>
<td>Epilepsy</td>
<td>Multiple operations</td>
<td>No</td>
<td>68</td>
<td>Yes (chondrosarcoma)</td>
</tr>
<tr>
<td>Forsythe, 1947</td>
<td>51, M</td>
<td>Convexity</td>
<td>ND</td>
<td>Headache</td>
<td>Removed in a piecemeal fashion</td>
<td>Yes</td>
<td>6</td>
<td>ND</td>
</tr>
<tr>
<td>Richards, 1960</td>
<td>49, M</td>
<td>Sella turcica</td>
<td>ND</td>
<td>Visual disturbance</td>
<td>PR</td>
<td>Yes</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>Herskovitz, 1973</td>
<td>57, F</td>
<td>Convexity</td>
<td>ND</td>
<td>Gait disturbance, memory disturbance</td>
<td>GTR</td>
<td>Yes</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>Ito, 1974</td>
<td>24, M</td>
<td>Posterior clinoid</td>
<td>6</td>
<td>CN6</td>
<td>PR</td>
<td>NAa</td>
<td>NAa</td>
<td>NAa</td>
</tr>
<tr>
<td>Himuro, 1977</td>
<td>52, M</td>
<td>Parasellar region</td>
<td>ND</td>
<td>Visual disturbance</td>
<td>PR</td>
<td>No</td>
<td>1</td>
<td>ND</td>
</tr>
<tr>
<td>Ikeda, 1980</td>
<td>41, M</td>
<td>Parasellar region</td>
<td>ND</td>
<td>CN6</td>
<td>STR</td>
<td>No</td>
<td>ND</td>
<td>ND</td>
</tr>
<tr>
<td>Matz, 1981</td>
<td>20, M</td>
<td>Convexity</td>
<td>8</td>
<td>Headache, vomiting, Visual disturbance</td>
<td>GTR</td>
<td>No</td>
<td>3</td>
<td>ND</td>
</tr>
<tr>
<td>Yamaguchi, 1983</td>
<td>24, M</td>
<td>Middle fossa</td>
<td>2</td>
<td>CN9, 10, 11 left hemiparesis</td>
<td>Multiple operations</td>
<td>No</td>
<td>45</td>
<td>Yes</td>
</tr>
<tr>
<td>Crawford, 1987</td>
<td>48, M</td>
<td>Falx</td>
<td>ND</td>
<td>Headache, gait disturbance</td>
<td>GTR</td>
<td>Yes</td>
<td>3</td>
<td>ND</td>
</tr>
<tr>
<td>Hatayama, 1989</td>
<td>15, M</td>
<td>Parasellar region</td>
<td>ND</td>
<td>CN5, CN6</td>
<td>STR</td>
<td>Yes</td>
<td>ND</td>
<td>ND</td>
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<td>Masuyama, 1994</td>
<td>28, F</td>
<td>Convexity</td>
<td>5</td>
<td>Epilepsy</td>
<td>GTR</td>
<td>Yes</td>
<td>1</td>
<td>ND</td>
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<tr>
<td>Sato, 1996</td>
<td>38, M</td>
<td>Posterior clinoid</td>
<td>4</td>
<td>Epilepsy</td>
<td>PR</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
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<tr>
<td>Nagai, 1998</td>
<td>45, F</td>
<td>Convexity</td>
<td>4.5</td>
<td>Headache</td>
<td>GTR</td>
<td>Yes</td>
<td>ND</td>
<td>ND</td>
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<tr>
<td>Haddad, 1998</td>
<td>25, M</td>
<td>Convexity</td>
<td>6</td>
<td>Epilepsy</td>
<td>GTR</td>
<td>Yes</td>
<td>12</td>
<td>No</td>
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<tr>
<td>Lin, 2002</td>
<td>15, M</td>
<td>Falx</td>
<td>14</td>
<td>Headache, epilepsy, sleep apnea</td>
<td>GTR</td>
<td>Yes</td>
<td>36</td>
<td>No</td>
</tr>
<tr>
<td>Omalu, 2003</td>
<td>53, M</td>
<td>Falx</td>
<td>8</td>
<td>Epilepsy</td>
<td>GTR</td>
<td>NAa</td>
<td>NAa</td>
<td>NAa</td>
</tr>
<tr>
<td>Bonde, 2007</td>
<td>20, M</td>
<td>Petrous bone</td>
<td>ND</td>
<td>Hemiplegia, CN10</td>
<td>PR</td>
<td>NAa</td>
<td>NAa</td>
<td>NAa</td>
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<tr>
<td>Inoue, 2009</td>
<td>29, M</td>
<td>Sella turcica</td>
<td>ND</td>
<td>Headache, visual disturbance</td>
<td>STR</td>
<td>Yes</td>
<td>60</td>
<td>No</td>
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<tr>
<td>Somerset, 2010</td>
<td>33, F</td>
<td>Convexity</td>
<td>9.5</td>
<td>Headache, hemiplegia</td>
<td>GTR</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
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<td>Venkata, 2011</td>
<td>24, M</td>
<td>Convexity</td>
<td>7</td>
<td>Headache, epilepsy</td>
<td>GTR</td>
<td>Yes</td>
<td>ND</td>
<td>No</td>
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<td>Lotfinia, 2012</td>
<td>73, M</td>
<td>Foramen magnum</td>
<td>ND</td>
<td>Quadriplegia, headache</td>
<td>GTR</td>
<td>Yes</td>
<td>18</td>
<td>No</td>
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<tr>
<td>Present case</td>
<td>43, M</td>
<td>Posterior clinoid</td>
<td>3</td>
<td>CN6</td>
<td>STR</td>
<td>No</td>
<td>6</td>
<td>No</td>
</tr>
</tbody>
</table>

Abbreviations: CN, cranial nerve; GTR, gross total resection; NA, not applicable; ND, no data available; PR, partial resection; STR, subtotal resection.

*Death due to postoperative complications.*
Conclusion

Intracranial osteochondromas are rare but should be included in the differential diagnosis of intracranial extra-axial tumors demonstrating atypical patterns of calcification. Thallium-201 SPECT might be useful in differentiating osteochondromas from other intracranial tumors. Although the long-term treatment outcome of intracranial osteochondromas is still unknown because of the scarcity of previous data, the risk of the goal of total resection should be carefully weighed against the chance of recurrence given the benign nature of the disease.

References