Intradural Retroclival Chordoma

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A 43-year-old woman presented with dizziness, ataxia and right hearing difficulty. Her magnetic resonance images demonstrated an inhomogeneously contrast-enhanced large tumor growing into right cavernous sinus and Meckel’s cave located totally within intradural retroclival region. She underwent retromastoid suboccipital craniotomy to resect the tumor mass and adjuvant gamma knife radiosurgery for remnant tumor at 1 month after operation. Adjuvant radiosurgery after surgical excision seems to be effective for the treatment of intradural extraosseous chordomas.

KEY WORDS: Chordoma - Intradural - Retroclival.

INTRODUCTION

Chordomas are rare tumors of central nervous system (about 0.2% of 6000 brain tumors) that arise from remnants of the primitive notochord. They can arise elsewhere, but tend to cluster at the two ends of the primitive notochord, approximately 35% cranially in sphenoccipital region (clivus), and 50% in the spine at the sacrococcygeal region. This neoplasm is thought to be essentially extradural in nature and generally associated with extradural extension and bone destruction. However, if allowed to grow for long periods, it may invade the dura and extend intradurally as well as extradurally.

Intradural chordomas are very rare tumors that should be distinguished from neurinomas and classic chordomas, because of their different biological behaviors.

We describe the rare case of a patient with intradural chordoma located in retroclival and cerebellopontine angle region, which was near totally resected via retromastoid suboccipital approach, followed by adjuvant gamma-knife radiosurgery for remnant tumor.

CASE REPORT

This 43-year-old woman was healthy until she suffered an episode of dizziness, ataxia, mild right hearing difficulty and right facial numbness.

Examination

On initial evaluation, the patient had ataxic gait. Cranial nerve examination revealed no gag reflex, uvula deviation to left side, decreased hearing acuity on right ear, decreased taste on right side of tongue and numbness on right V1, V2 dermatome. The remainder of the neurological examination was within normal ranges. Magnetic resonance imaging demonstrated that the tumor located in the retroclival and right cerebellopontine angle, growing into right cavernous sinus and Meckel’s cave and compressing the brainstem from medulla to midbrain (Fig. 1). The tumor appeared as a low-intensity area on the T1-weighted image and as a high-intensity area on the T2-weighted image. The tumor was enhanced inhomogeneously after administration of gadolinium-diethylenetriamine pentaacetic acid (Gd-DTPA).

Operation

Under the impression of acoustic schwannoma, the patient...
underwent operation. We chose the retromastoid suboccipital approach. Crainotomy was made until transverse sinus and part of a sigmoid sinus were exposed. A soft, friable and grayish mass was seen. The tumor was clearly distinguished from the brain cortex and was easily removed by means of tumor forceps and suction tools. Also, this tumor contained a necrotic materials. Some areas of tumor was attached to right trigeminal nerve and located near the right facial and vestibulocochlear nerve complex. After near total resection of tumor (Fig. 2), these nerves (trigeminal, facial and vestibulocochlear nerves), superior cerebellar artery and posterior inferior cerebellar artery were well preserved. Rest of operation was uneventful.

Postoperative course
The patient complained dizziness, but other symptoms, such as ataxia, right facial numbness without paralysis, right hearing disturbance, were much more improved. In particular, the hearing disturbance was more improved after operation. Preoperative hearing threshold was 90 dB on the right and 45 dB on the left side by brainstem auditory evoked potential (BAEP), and it was improved to 35 dB on the right and 30 dB on the left side by BAEP studied on 11th postoperative day. Postoperative MR imaging indicated residual tumor in right cerebellopontine area (Fig. 3). Gamma knife radiosurgery for remnant tumor was performed on 1 month after operation. The remnant tumor volume was 8.3 cc and marginal dose of 15 Gy was administered. MR imaging, performed 14 months after gamma knife radiosurgery, revealed decreased size of remnant tumor compared with one before gamma knife radiosurgery (Fig. 4).

Histological findings
The tumor tissue was histologically characterized by lobules composed of typical physaliphorous cells with abundantly vacuolated cytoplasm. Immunohistochemical analysis showed the positive expression for cytokeratin, epithelial membrane antigen, vimentin and S-100 protein (Fig. 5). The histological features and antigen expression were consistent with the diagnosis of chordoma.

DISCUSSION
Including the this case, only 18 previous cases of primary
intradural and intradural extraosseous chordoma have been reported (Table 1)\textsuperscript{1,4,6,8,13}). It can be seen that many are from the pre-MRI era, and many do not provide histological or immunocytochemical evidence, raising the possibility that some may not be entirely extraosseous and some may be not actually be chordoma. Their locations included the prepontine region in ten cases, the suprasellar region in two cases and the intrasellar, foramen magnum, tentorium and ponto-cerebellar region in one case, respectively.

Ectopic notochordal tissue was found in an intradural location anterior to the pons in up to 2\% at autopsy; these have been given name “ecchordosis physaliphora”\textsuperscript{2,3,20}). The differential diagnosis of ecchordosis physaliphora and chordoma based on histological and radiological features is likely to be difficult. The MIB-1 proliferating cell index may be useful in the histological differential diagnosis of chordomas, which are neoplastic tumors, and ecchordoses, which are not true tumor\textsuperscript{13}.

Intradural extraosseous chordoma has clearly different features from those of typical chordoma\textsuperscript{17}). These entities show different biological behavior, with gradient evolution of growth and malignancy from the usually asymptomatic ecchordosis, to the slowly evolving intradural chordoma, to the highly malignant and invasive chordoma\textsuperscript{10}). Classic chordomas in bone frequently have ill-defined margins, and complete resection is usually not feasible even with extensive surgery because a few tumor cells are often left behind in the bone or dura. In contrast, the intradural type of tumor characteristically has a slower growth pattern, sharply circumscribed margins compared with classic chordomas. However, stereotactic radiosurgery is valuable as an adjuvant or primary treatment for selected patients with chordoma and has potential advantages over standard fractionated irradiation, although the length of follow-up review has been insufficient\textsuperscript{7}). For tumor control, conventional radiotherapy needs very high doses of radiation. This dose may carry a high risk of severe radiation injury to the normal brain structures\textsuperscript{9}). In contrast, radiosurgery has the advantage of high dose single-session irradiation and efficacy for small sized tumor mass\textsuperscript{12}). For this reason, we selected stereotactic radiosurgery for remnant tumor in our case. Also, proton-beam irradiation is reported to be efficacious for the treatment of chordomas\textsuperscript{10}).

| Author(Year) | Age (yrs), Sex | Tumor location | Surgery extent | Radio-surgery | Recurrence |
|--------------|---------------|---------------|---------------|--------------|------------|
| Dahlin and MacCarty (1952) | ND | Prepontine | None | None | No |
| Bartolini (1974) | 57, M | Prepontine | None | None | No |
| Mathews and Wilson (1974) | 41, M | Intrasellar | Subtotal | Yes | ND |
| Stam and Kamphorst (1982) | 75, M | Prepontine | None | None | ND |
| Mapstone et al. (1983) | 26, M | Prepontine | Subtotal | ND | ND |
| Vaquero et al. (1983) | 39, F | Suprasellar | Total | ND | ND |
| Yuhi et al. (1986) | 28, M | Prepontine | ND | None | No |
| Katayama et al. (1991) | 30, F | Foramen magnum | Total | None | No |
| Warnick et al. (1991) | 58, M | Fentorium | Subtotal | ND | ND |
| Hardie (1992) | 28, F | Prepontine | ND | ND | ND |
| Tashiro et al. (1994) | 56, F | Prepontine | ND | ND | ND |
| Wanihusi et al. (1994) | 54, M | Suprasellar | Total | None | No |
| Nishigaya et al. (1998) | 56, M | Prepontine | Subtotal | None | No |
| Korinth et al. (1998) | 48, F | Ponto-cerebellar | Subtotal | Yes | ND |
| Danilewicz et al. (2000) | 19, F | Prepontine | Total | None | No |
| Dow et al. (2003) | 9, F | Cerebellar | Total | None | No |
| Masui et al. (2006) | 63, M | Prepontine | Total | None | No |
| Present case | 43, F | Prepontine | Nearly total | Yes | No |

ND : not described

ND: not described
CONCLUSION

We achieved near total resection of intradural chordoma and adjuvant gamma knife radiosurgery was performed for remnant tumor at 1 month after surgery. Although there are no sufficient data for efficacy of an adjuvant gamma-knife radiosurgery, the tumor of our case gradually decreased in size based on follow-up MR imaging.

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