Neuronavigation-guided Endoscopic Endonasal Excision of Schwannoma-like Chordoma of Meckel’s Cave: A Case Report

Ali AKAY¹, Sercan GODE², M. Sedat CAGLI³

¹Kent Hospital, Department of Neurosurgery, Izmir, Turkey
²Ege University, School of Medicine, Department of Otolaryngology, Izmir, Turkey
³Ege University, School of Medicine, Department of Neurosurgery, Izmir, Turkey

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Corresponding author: Ali AKAY  dr.aliakay@gmail.com

ABSTRACT

Chordomas are locally aggressive malignant tumors due to their recurrence potential and originate from embryonic notochord remnants. Chordomas can originate anywhere on the axial skeleton. They are extradural and spread by bone destruction. Chordomas are locally aggressive tumors that invade the dura mater, and may also present with secondary intradural growth. The Meckel’s cave location of chordomas has been very rarely reported in the literature. Chordomas located in Meckel's cave can be radiologically confused with trigeminal schwannomas. Herein, we report a case of Meckel’s cave chordoma that was successfully excised through neuronavigation-guided endoscopic endonasal excision, a technique commonly used in skull base surgeries.

KEYWORDS: Chordoma, Endoscopic endonasal, Meckel's cave, Skull base

INTRODUCTION

Chordomas are midline tumors that develop from primitive notochord remnants (2). The incidence of these tumors is 1% in intracranial tumors and 4% in tumors of the bone (5,8). The most frequent localizations are the sacrococcygeal region (30 to 50%), spheno-occipital region (30 to 35%) and vertebral region (15 to 30%). Chordomas usually develop in the cranium from the spheno-occipital synchondrosis of the clivus, but may also develop from the lower clivus and dorsum sellae. They are extradural and are spread by bone destruction. Chordomas are locally aggressive tumors that invade the dura mater, and they may also present with secondary intradural growth (10,14). However, chordomas have been also shown in the literature to present with primary intradural development, although rarely (4,15). Intradural chordomas most commonly demonstrate prepontine localization (1,6,13,15). The prognosis of intradural chordomas is better than that of extradural chordomas. Recurrence is rare due to the absence of bone invasive and the possibility of total (en-bloc) removal (1,3,6,13). One of the rarely reported localizations for intradural or extradural chordomas is Meckel's cave (MC). Herein, we present a rare case of extradural chordoma of Meckel’s cave.

CASE REPORT

A 41-year-old female patient presented with double vision for the past four months. Cranial magnetic resonance imaging (MRI) showed a massive lesion destroying the petrous bone and the left lateral wall of the clivus with heterogeneous contrast in the left Meckel’s cave (Figure 1A-C). Neurological examination demonstrated an outward visual limitation in the left eye, and sixth cranial nerve palsy. The preliminary radiological diagnosis of the mass lesion was schwannoma. The patient underwent neuronavigation-guided endoscopic endonasal excision. The Meckel’s cave location of chordomas has been very rarely reported in the literature. Chordomas located in Meckel's cave can be radiologically confused with trigeminal schwannomas. Herein, we report a case of Meckel’s cave chordoma that was successfully excised through neuronavigation-guided endoscopic endonasal excision, a technique commonly used in skull base surgeries.
DISCUSSION

In 1984, Kapila et al. reported Meckel’s cave as a dural sheath on the gasser ganglion, located on the posterior inferior surface of the cavernous sinus, and on the medial posterior of the medial fossa (7). Primer tumors of Meckel’s cave make up less than 0.5% of all intracranial tumors (11). The majority of Meckel’s cave tumors are trigeminal schwannomas (33%). The other pathologies include meningiomas, invasive pituitary macroadenomas, metastases, epidermoid cysts, lipoma, lymphoma, and other skull base tumors (12). The incidence of trigeminal schwannomas among all intracranial tumors is 0.07 to 0.28%; however, the incidence of all intracranial schwannomas is reported to be between 0.8 to 8%. Trigeminal schwannomas enlarge, spreading from the middle fossa to the posterior fossa and present as a characteristic dumbbell shape on radiological images. However, this may not be the case with smaller lesions as in this case. On MRI, trigeminal schwannomas on T1-weighted imaging appear as hyperintense images, whereas they appear as hyperintense and heterogeneous contrast enhancements on T2- and contrast-enhanced images.

The location of chordomas at the skull base is generally in the neighborhood of the midline of the clivus. In the literature, chordomas are reported to be rarely observed in places such as the intrasellar, suprasellar, tentorium cerebelli, foramen magnum, cervical and thoracic region. Chordomas of the Meckel’s cave are rarely observed in the literature. Their radiological characteristics are similar to those of schwannomas, and may be confused with trigeminal schwannoma especially in this region. Although similar cases have been reported in the literature, they are very rare (4,9). The classical means of accessing tumors of Meckel’s cave is by the microscopic subtemporal extradural approach.

Figure 1: A) Matching of endoscopic image with cranial CT bone window image of Meckel’s cave localization lateral to the septum within the sphenoid sinus. B) Matching of tumor tissue access with contrast axial cranial MRI image following removal of the window-shaped bone in the sphenoid sinus. C) Matching of endoscopic image of tumor tissue with coronal plane of contrast axial cranial MRI image.
CONCLUSION

With recent developments in endoscopic technology, there is increasingly frequent use of endoscopic endonasal skull base interventions. In particular, nowadays, endoscopic endonasal skull base surgery has become the gold standard method for accessing sellar, clival, and parasellar masses. As it is particular in this case, we suggest that the use of navigation also continuously improves the reliability of endoscopic surgery by eliminating the misleading effect of anatomical variations.

REFERENCES