Case Report

**Suboccipital osteoblastoma: Microsurgical resection of a rare entity**

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Abstract

**Background:** Osteoblastomas are rare lesions comprising 1% of all bone tumors. The occipital bone is one of the rarest affected bone, with only 11 cases reported during the last 40 years.

**Case Description:** Here, we describe the clinical presentation and the radiological features of a suboccipital osteoblastoma that was successfully resected in a 30-year-old man. A short video shows the microsurgical removal of the lesion. There was no recurrence during a 12-month follow-up.

**Conclusions:** Even if osteoblastomas are benign tumors, a complete removal has to be achieved to reduce the risk of recurrences. This makes necessary an appropriate monitoring of the patient.

**Key Words:** Occipital bone, osteoblastoma, park bench position, suboccipital approach

**INTRODUCTION**

Osteoblastomas are rare lesions comprising 1% of all bone tumors. Skull bones are affected in only approximately 3% of the cases, and the occipital bone is far less involved than facial, temporal, frontal, and skull base bones. Here, we present the 12th case of a patient with an occipital osteoblastoma reported in the literature during the last 40 years, with a detailed description of its main features as well as of our surgical strategy.

**CASE REPORT**

A 30-year-old man, without significant medical history, was admitted to our department for a painful left occipitocervical swelling, progressively worsening over the past several months. A physical examination revealed an occipital hard mass covered by the normal skin, without any neurological deficit.

A computerized tomography (CT) scan showed an osteolytic lesion inside the diploic space surrounded by sclerotic tissue expanded to the left squamous portion of the occipital bone around the foramen magnum. In addition, the CT angiography with three-dimensional (3D) reconstruction revealed a close relation between the tumor and the vertebral and occipital arteries [Figure 1]. Eventually, a craniocervical...
magnetic resonance imaging (MRI) showed the soft extracranial tissues partially affected by the tumor, without intracranial extension [Figure 1].

The patient underwent a left suboccipital approach in the park bench position. After the dissection of the soft tissues, the lesion was targeted according to anatomic landmarks. Under the operating microscope, the tumor was progressively removed, using high speed drills and rongeurs, up to the normal dura and surrounding normal bone. However, the left occipital condyle was left undisturbed [Figure 2; Video 1] (Supplemental Digital Content, illustrating the microsurgical removal of the lesion).

The postoperative course was uneventful, and the patient’s symptoms disappeared. The histopathological examination was consistent with benign osteoblastoma, and the postoperative CT scan showed the complete resection of the lesion [Figure 3]. There was no recurrence during a 12-month follow-up.

**DISCUSSION**

Osteoblastomas comprise 1% of all bone tumors. Skull bones are affected in only 3% of the cases, and occipital osteoblastomas are even rarer. To the best of our knowledge, this is the 12th case reported in the last 40 years.\(^2,5,14,15\)

The presentation of the tumor is more frequent in men less than 30 years old; calvarian osteoblastomas essentially share the same characteristics of the osteoblastomas in other locations,\(^10\) and they are treated for intractable pain, increase in size, and bone destruction.\(^20\)

Occipital osteoblastomas present as circumscribed expansive intraosseous lesions between the inner and outer table of the occipital bone, with lytic and sclerotic appearance in plain radiographs and CT scans.\(^2,16\) CT angiography may help to describe the vascular supply of the tumor, thus estimating the necessity of a...
presurgical embolization. The MRI is too variable to define the diagnosis of osteoblastoma, although it is very useful to determine the involvement of the adjacent soft tissues, as in our case.[14,16]

Even though giant osteoid osteomas are reported, unlike osteoblastomas,[9] they frequently have less than 15 mm in their maximal diameter, and the severe pain produced by them during the night is usually relieved by acetylsalicylic acid.[12,20] On the other hand, osteosarcomas, dermoid cysts, intradiploic meningiomas, as well as eosinophilic granulomas may also mimic osteoblastomas.[17,19] Only cysts, intradiploic meningiomas, as well as eosinophilic granulomas may also mimic osteoblastomas.[17,19] Only the histopathological examination, which is characterized by irregular bony trabeculae surrounded by osteoblasts and osteoid tissue, will define the diagnosis.[5,14]

The recurrence of osteoblastomas after surgery is approximately 9.8–15%.[20] They may show sarcomatous changes, local extension into the extraskeletal soft tissues, and may even metastasize.[13,17] Their clinical outcome is basically related with the completeness of the resection and with their location. Apparently, osteoblastomas confined to short and flat bones have a more aggressive behavior, which leads us to conclude that the resection of osteoblastomas must be as large as possible. Hence, in the case of a tumor located in the occipitocervical junction, a stabilization may be required, as well as the transposition of the vertebral artery.[5,7] However, occipitocervical stability is not compromised if less than one-half of the C0–C1 joint is removed.[6,8,18] Some osteoblastomas may be hypervascularized, thus requiring a preoperative embolization by using particles or some another kind of endovascular agents.[5] Because of the risk of recurrence, a careful follow-up of the patient is essential.[7]

In summary, occipital osteoblastomas are extreme rare benign bone tumors with typical histological features, sometimes hypervascularized, and with a high tendency to recur when they are not completely resected. At times, their treatment may require presurgical embolization, occipitocervical fusion, and/or transposition of the vertebral artery. Their high risk of recurrence makes necessary an appropriate monitoring of the patient.

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**Conflicts of interest**

There are no conflicts of interest.

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