

# Unique Image Characteristics of an Occipital Primary Chondroblastic Osteosarcoma: A Rare Case Report and a Brief Literature Review

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## Abstract

### Keywords

- chondroblastic osteosarcoma
- occipital bone
- skull base
- computed tomography
- magnetic resonance imaging

Primary osteosarcomas of the skull and skull base are rare and comprise < 2% of all skull tumors. In head and neck osteosarcomas, the chondroblastic subtype occurs most frequently, which has an exceedingly poor outcome, but its image characteristic remains unknown. Herein, we report a case in the right occipital bone of the skull base and the unique characteristics of image. Pathologic examination of the surgical specimens led to the diagnosis of chondroblastic osteosarcomas. We believe those image characteristics can improve the understanding of skull chondroblastic osteosarcoma and the preoperative diagnosis.

## Introduction

Osteosarcoma (OS) develops most frequently in the extremities, and it is the most common histologic form of the primary bone cancers.<sup>1,2</sup> Head and neck OSs are rare, comprising only 6 to 10% of all OSs.<sup>3,4</sup> They typically present in the third or fourth decade of life and comprise only 1% of all pediatric head and neck malignancies. The most common craniofacial sites affected by OSs are the mandible and maxilla, followed by the calvaria and then the skull base.<sup>4–6</sup> On cytology, OS can be divided into several pathologic types, including the pleomorphic, epithelioid, chondroblastic, small cell, mixed, and osteoclast-like giant cell types.<sup>6</sup> In head and neck OSs, the chondroblastic type occurs most frequently.<sup>7</sup>

Skull base OSs can be challenging to resect and an aggressive surgical approach can result in poor cosmetic outcome.<sup>8</sup> Imaging plays a crucial role in the diagnosis of each subtype of OS and ultimately in patients' survival because the diagnosis is based on a combination of histopathologic and imaging features. The therapeutic options

and prognoses for different types of OS differ from each other, so correct diagnosis is essential.<sup>9,10</sup> Magnetic resonance imaging (MRI) or computed tomographic (CT) scan should be used to assess the extent of the primary tumor.<sup>11</sup>

In this case report, we describe a pediatric patient of occipital OS of the chondroblastic type. The chondroblastic type of OS has an exceedingly poor outcome.<sup>12</sup> However, the detailed imaging description of such cases have not been reported in the previous literatures. We present the CT, MRI, and enhanced MRI features of this case, followed by a brief review of the related cases reported in the previous literatures

## Case Report

A 9-year-old boy was admitted to our hospital with a major complaint of a growing mass on his head. Physical examination found a firm and tough mass on the right occipital that showed no tenderness upon palpation. CT scan showed the right occipital bone to be irregularly thickened with fluffy and cloudy calcification, with a mass deriving from the

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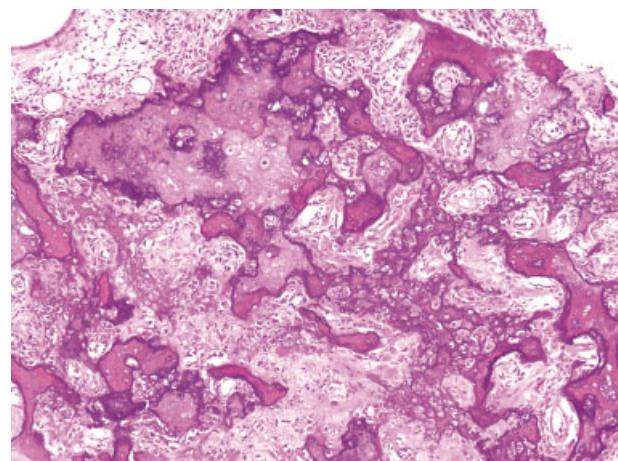
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internal occipital protuberance extending toward the basilar part of the occipital bone, invading the neighboring jugular foramen, the sublingual neural tube, and the mamillary process. On MRI, the lesion was 4.5- × 5.5- × 6.5-cm in size with calcifications areas of hypointensity in T1- and heterogeneous in T2-weighted series. Contrast MRI showed peripheral and septal enhancement in the interior side of the tumor (**►Fig. 1**). Significant mass effect was present, distorting the cerebellar hemisphere, pons, and the forth ventricle, which led to hydrocephalus, and the oppression of the sigmoid sinus and the transverse sinus. Histopathology examination reported lace-like osteoid material abutting the neoplastic cells (**►Fig. 2**), corresponding to the features of chondroblastic OS, and occipital bone chondroblastic OS was the final definitive diagnosis. A subtotal resection of the tumor was performed followed by radiation therapy. The patient died after half a year of local recurrence.

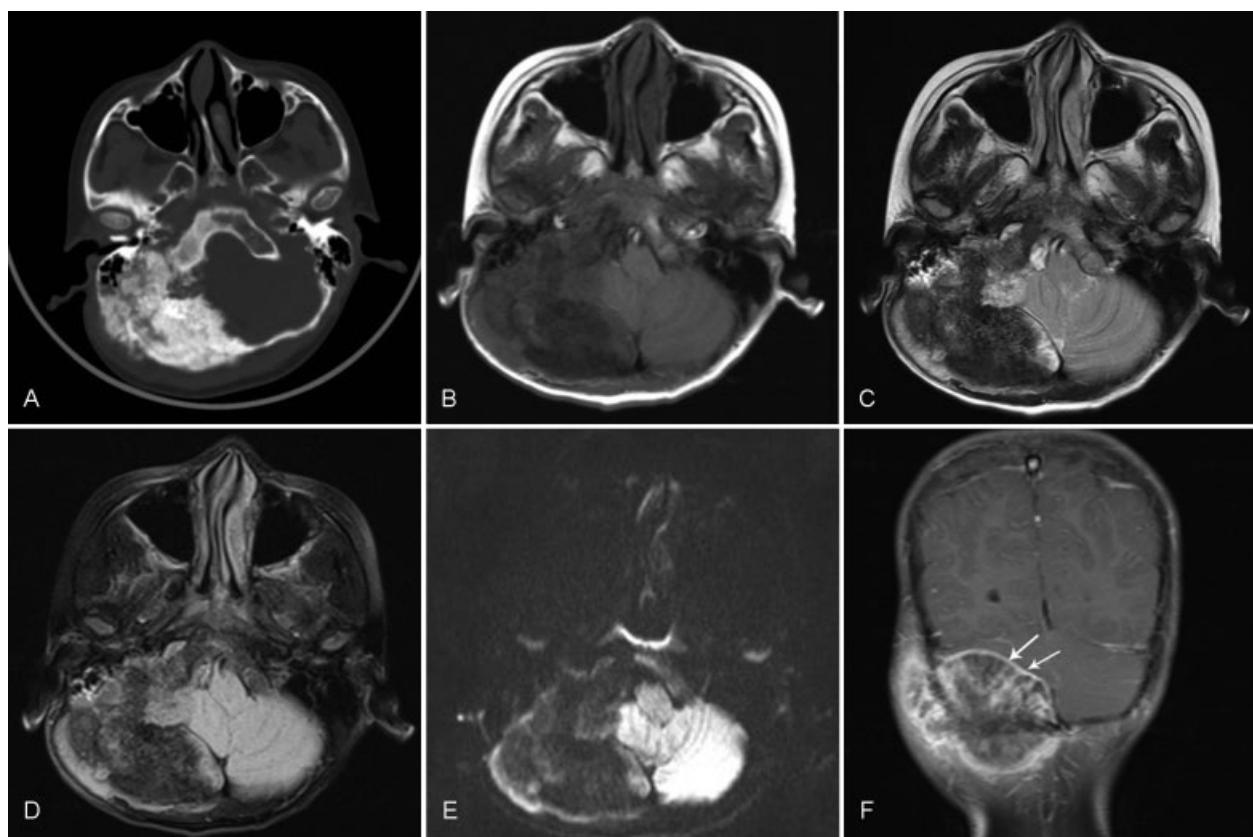
## Discussion

Craniofacial OSs are rare. They typically present in the third or fourth decade of life, account for fewer than 5% of OSs in children, and comprise only 1% of all pediatric head and neck malignancies. The most common craniofacial sites are the mandible and maxilla, followed by the calvaria and then the skull base.<sup>13–15</sup> Our case in the right occipital bone of skull



**Fig. 2** Histopathologic examination (hematoxylin and eosin,  $\times 200$ ) shows lace-like osteoid material abutting the neoplastic cells.

base is a very rare location. A search of the English language literature revealed 22 cranial OSs previously reported in children (**►Table 1**): 12 calvarial tumors and 10 tumors of the skull base. The mean age of the pediatric patients with cranial OS was 12.2 years old in this table. The patient in our case suffered at a younger age. On cytology, OS can be divided into pleomorphic, epithelioid, chondroblastic, small cell,



**Fig. 1** (A) Computed tomography of the skull shows fluffy calcification. (B) T1-weighted image shows a 4- × 8- × 10-cm mass lesion, isointense to the skull. (C, D) The mass is hypointense in most areas in the T2-weighted series, with focal high signals in the T2-weighted series and reduced signal in FLAIR series. (E) In Gd-enhanced MRI, most areas show no enhancement or heterogeneous enhancement, with peripheral and atypical septal enhancement on the coronal plane (white arrows). (F) No hyperintensity was observed in both intra- and peritumoral areas in the DWI series.

**Table 1** Summary of previously reported cases of calvarial and skull base osteosarcomas in pediatric patients

Author and year	Age at diagnosis	Location	Extent of resection	Adjuvant therapy	Follow-up	Outcome
Garland, 1945	17, M	Occipital	NR	RT	NR	NR
Reddy et al, 1973	8, F	Occipital	Biopsy	RT	NR	Dead, progressive disease
Goodman and McMaster, 1976	15, F	Parietal-occipital	NR	Chemotherapy and RT	6	Alive, disease free
Wang et al, 1981	17, M	Frontal-parietal-occipital	NR	RT	6	Dead, progressive disease
Benson et al, 1984	11, M	Frontal	NR	Chemotherapy	12	Alive, disease status
Sundaresan et al, 1985	11, M 13, F	Parietal Skull base	STR STR	Chemotherapy Chemotherapy	36 66	Alive, progressive disease Alive, disease free
Kornreich et al, 1988	12, F	Parietal	NR	Chemotherapy	144	Alive, disease free
Mark et al, 1991	14, M	Anterior skull base	NR	Chemotherapy and RT	12	Dead, progressive disease
Shramek et al, 1992	8, M	Parietal-occipital	GTR	Chemotherapy and RT	16	Alive, progressive disease
Salvati et al., 1993	11, M	Frontal-temporal	STR	RT	9	Dead, progressive disease
Chander et al, 2003	15, F	Frontal	GTR	NR	NR	NR
Author and Year	Age at diagnosis	Location	Extent of resection	Adjuvant therapy	Follow-up	Outcome
Ellison et al, 1996	11, F	Skull base	STR	Chemotherapy and RT	NR	NR
Gadwal et al, 2001	9, M 1, M	Sphenoid Sphenoid	NR NR	RT NR	9 NR	Dead, progressive disease NR
Chennupati et al, 2008	14, F	Skull base	Biopsy	Chemotherapy and RT	12	Alive, progressive disease
Kirby et al, 2011	16, M	Parietal	GTR	Chemotherapy	5	Alive, disease free
Oakley et al, 2011	15, M	Anterior skull base	GTR	Chemotherapy	NR	NR
Ohno et al, 2011	14, F	Anterior skull base	STR	Chemotherapy	26	Dead, progressive disease
Meel et al, 2012	10, M	Sphenoid	Biopsy	Chemotherapy and RT	18	alive, disease free
Caroline et al, 2014	14, M 12, M	Parietal skull base	GTR GTR	Chemotherapy Chemotherapy	16 12	Alive, disease free
He et al, 2016	9, M	Occipital	STR	Chemotherapy and RT	6	Dead, progressive disease

Abbreviations: GTR, gross total resection; NR, not reported; RT, radiation therapy; STR, subtotal resection.

mixed, and osteoclast-like giant cell types.<sup>6</sup> Our case is a chondroblastic subtype, which occurs most frequently in head and neck OSs.

The etiology of OS is unknown, but the major risk factors for development of OS in craniofacial bones may be similar to those of the long skeletal bones, consisting of exposure to radiation, retinoblastoma, Li-Fraumeni syndrome, and Paget's disease. The skull is a favored site for OS arising out of Paget's disease. Other bone abnormalities, such as fibrous dysplasia, multiple osteochondromatosis, chronic osteomyelitis, myositis ossificans, and trauma, have also been proposed as risk factors.<sup>7,15,16</sup> The presenting symptoms varied with the location of the tumors. The maxillary or cranial lesions usually produced no pain, which was in accordance with our case; however, mandibular tumors frequently presented with focal painful swelling.<sup>17,18</sup> Other common presenting symptoms include headache, cranial nerve palsies, exophthalmos, and visual impairment due to different location of the tumor.<sup>5,13</sup>

CT best demonstrates tumor mineralization, especially when minimal, and it is usually able to demonstrate tumor extension into the soft tissues. Hemorrhage, necrosis, and unmineralized, chondroblastic, or fibroblastic components of the tumor will appear as areas of low attenuation on CT if present. Unlike any other conventional OSs, we see fluffy calcification in our case, and we believe it is a characteristic of OS. The osteoblastic subtype is most common with nearly 90% containing variable amounts of cloudlike opacities.<sup>19</sup> Bose<sup>20</sup> reported an osteoblastic OS that appears as a large soft tissue density mass with a few bony densities. Compared with our case, the soft tissue mass is prominent and the calcification is less and diffuse.

MRI is the preferred modality for locally staging OS, and it should be performed before percutaneous biopsy because it can help identify areas of viable tumor and mineralized matrix. In our case of gadolinium (Gd)-enhanced MRI, we found no enhancement or heterogeneous enhancement in most areas of the tumor, with septonodular and rim enhancement, which is

in accordance with the current literature. Areas that demonstrate either a heterogeneous enhancement pattern or lack enhancement are the preferred sites for biopsy because they are more likely to contain both chondroid and osteoid elements that are necessary for the correct diagnosis.<sup>21,22</sup> Chondrosarcomas shows similar image characteristic, but they occur in an older age with a mean age of 57 years old. DWI can also help identify chondroblastic OS. Chondroblastic OSs also have significantly higher minimum and maximum apparent diffusion coefficient (ADC) values compared with other conventional OS subtypes, but they have a lower minimum ADC and similar maximum ADC value compared with chondrosarcoma.<sup>23</sup>

Skull base OSs can be challenging to resect, and an aggressive surgical approach can result in poor cosmetic outcome. Thus, skull base tumors have a poorer prognosis than mandibular or maxillary tumors.<sup>3</sup> Complete surgical excision is the mainstay of treatment of the primary tumor. Local recurrence is the main reason of treatment failure and mortality in head and neck OSs. Positive margins and a high tumor grade correlate with a statistically significant decrease in survival.<sup>11</sup> In our case, the tumor could not be completely removed because it invades significant neighboring bone structures, including the jugular foreman and the sublingual neural tube. The patient died after 6 months as a result of local recurrence.

In summary, chondroblastic OS has been shown to be associated with a poor preoperative chemotherapy response and has a worse prognosis than other variants.<sup>24</sup> However, this subtype has some particular image characteristic, which helps surgeons identify before surgery and set early therapeutic regimens.

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