Osteoblastoma is a rare benign primary bone tumor. It accounts for 3% of benign and 1% of all primary bone tumors. The treatment goal is complete surgical resection. This treatment limits the risk of recurrence. As osteoblastoma is a highly vascular tumor, complete resection is often difficult. This report describes the case of a 19-year-old male patient who presented severe right-sided neck and shoulder pain. The computed tomography scan revealed a mass lesion on C7 compatible with osteoblastoma. Preoperative embolization and tumor resection were performed. At the 3-year follow-up, the patient had no restrictions on daily activities, and, to date, there has been no evidence of recurrence.

Abstract
Osteoblastoma is a rare benign primary bone tumor that account for 3% of benign and 1% of all primary bone tumors. The treatment goal is complete surgical resection. This treatment relates to its low incidence, non-specific symptoms and normal radiological study during the initial course of disease. The treatment goal is complete surgical resection which allows the complete regression of complaints and decrease the likelihood of relapse.

Case Report
A 19-year-old boy was referred to Orthopedic Consult for neck pain. The past medial history was unremarkable. He presented with a year history of severe right-sided neck and shoulder pain, worst in the night. The pain was progressive, not relieved with medical treatment and exacerbated in the

Introduction
Osteoblastoma is a rare and benign primary bone tumor that account for 3% of benign and 1% of all primary bone tumors. Approximately 32–46% involve the spine. These lesions occur predominately in second and third decade of life and in male with a 2:1 male to female ratio. Pain is the most commonly cited symptom and often present for greater than a year prior to diagnosis. Late diagnosis of these lesions relates to its low incidence, non-specific symptoms and normal radiological study during the initial course of disease.
last two months with paresthesias. On examination, cervical range of motion was restricted and painful. Neurological examination was unremarkable. Cervical spine X-ray was normal. A cervical CT scan revealed a mass lesion with well-defined limits involving facet joint and the right pedicle of C7, with about 18.9 × 15.2 mm larger diameter without spinal cord compression, compatible with osteoblastoma (►Fig. 1). We completed study with angio-CT and neck vessels Doppler revealing stenosis of vertebral artery with cleavage plan with the same (►Fig. 2). It was decided surgical resection.

The patient underwent preoperative embolization one day prior to surgery (►Fig. 3). On the following day, the patient underwent tumor resection by posterior approach followed by C6-C7 intersomatic arthrodesis with “cage” filled with iliac crest autograft for anterior approach (►Fig. 4).

The postoperative period was uneventful. A CT scan was obtained on a first postoperative day and demonstrated a complete surgical resection (►Fig. 5). He was discharged on the fifth day postoperatively. Histopathologic examination confirmed the diagnosis. Present at 3 years postoperatively, the patient presents no complaints and with normal daily life activity without evidence of recurrence (►Fig. 6).

**Discussion**

Osteoblastoma and osteoid osteoma were described in 1935 by Jaffe apud Samdani et al. They are both hypervascular tumors benign clinically and histologically similar. However, osteoblastoma demonstrates more aggressive characteristics. It can be locally aggressive and cause neurologic impairment due the expansible growth. Furthermore, it can undergo malignant transformation to osteosarcoma.

In spine osteoblastoma usually involves the posterior elements. Although it is considered a benign tumor, spinal lesions recurrence is not uncommon and have been reported in 9.7–15% of series of spinal osteoblastomas.
The most common presentation is neck pain and it is not relieved by nonsteroidal anti-inflammatory drugs. Other common signs and symptoms include torticollis, stiffness and local tenderness.

The diagnosis is usually made by CT scan. It will delineate the location and osseous involvement of the mass. Surgical excision is the treatment of choice for osteoblastoma. This treatment limits the risk of recurrence. The main reason for high rate of recurrence is incomplete resections. Careful pre-operative planning is essential.

Because osteoblastoma is a high vascular tumor, complete resections is often impeded by extensive intraoperative bleeding. Some authors recommend pre-operative embolization. It has been described since 1979 by Dick et al. and apud Samdani et al. as an adjunctive therapy for benign bone tumors. More recently, it has been success- fully applied to highly vascular spinal tumors. It reduces intraoperative bleeding, and help to enhance visualization to perform a complete resection. It also reduces post-operative complications.

The authors report a well-succeeded case of C7 osteoblastoma. In the current case, the authors preferred a two stage approach. The posterior approach allowed tumor resection. The anterior approach allowed anterior fusion for spine stabilization, with interbody cage and instrumentation with anterior plate and screws. We chose anterior arthrodesis to facilitate imaging follow-up, since it diminishes the imaging artifact in the posterior elements and that could hinder the interpretation of a possible local recurrence.

We perform a pre-operative embolization that facilitated tumor resection because it decrease bleeding and improve visualization. The complete resection of this lesion allowed the complete regression of complaints and decreases the likelihood of recurrence (without it, the recurrence rate has been reported between 9.8% and 15%).

Fig. 3  Preoperative embolization images.

Fig. 4  Left: intraoperative picture of tumor resection; right: postoperative X-ray (lateral view and AP).
Confl icts of Interest
The authors declare no confl icts of interest.

References