Spinal Osteoid Osteoma Revealed by Radiculopathy: Case Report and Literature Review

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Spinal Osteoid Osteoma Revealed by Radiculopathy: Case Report and Literature Review

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ABSTRACT

Osteoid osteoma (OO) is a benign tumor that usually occurs in long bones of young males. We report a rare case of spinal OO in a 25-year-old woman, revealed by a sciatica. Spinal radiographs and computed tomography scan were normal, although performed at 6 months of symptom evolution. On magnetic resonance imaging, however, an important edema of the right transverse process of L5 vertebrae was depicted but was inconclusive. The diagnosis of OO was finally retained on a second computed tomography scan with thinner slices focused on the edematous area. The patient had an en-bloc excision of the tumor with complete regression of symptoms. Due to the atypical clinical presentation and the absence of common findings in imaging, the diagnosis was delayed by 12 months. Radiculopathy caused by spinal OO is a rare condition with no more than 30 cases reported in the literature. In fact, spinal OO usually presents with inflammatory back pain or painful scoliosis. This case emphasizes the importance of early suspicion and diagnostic interventions in the detection and treatment of OO.

Keywords: osteoma, osteoid, spine, radiculopathy, sciatica

BACKGROUND

Osteoid osteoma (OO) represents about 2% to 3% of all bone tumors.1 It usually occurs in the metaphysis and diaphysis of long bones.2 The spine is involved in up to 25% of OOs, particularly posterior elements of the lumbar segment.3 The clinical presentation varies depending on the location of the tumor.

We herein report the case of an OO of the lumbar spine revealed by sciatica. Given the atypical clinical presentation, the infrequency of such a location, and the imaging diagnosis difficulty, our case is worthy to report. The management of this entity was discussed based on a literature review of spinal OO revealed by radiculopathy. The search of articles was performed in Medline, Embase, Scopus, and Cochrane library databases. The following key words were used: (“Osteoma, Osteoid”) AND (“spine” OR “radiculopathy” OR “sciatica” OR “brachial neuralgia”). Key words referred to medical subject heading (MeSH).

CASE PRESENTATION

A 25-year-old woman presented with low back pain and right L5 sciatica evolving for 6 months. She had no trauma or any relevant family or medical history. There was no history of fever or night sweats. She was experiencing morning stiffness and night pain that interfered with sleep. Pain was not relieved by paracetamol and opioids. Clinical examination revealed lumbar stiffness and no signs of neurological compromise except for the sciatica. Laboratory tests showed an erythrocyte sedimentation rate at 33 mm and C-reactive protein level at 5.3 mg/L.

All laboratory and imaging results, including plain radiography and a lumbar computed tomography (CT) scan, were initially normal. Magnetic resonance imaging (MRI) performed 10 months after symptom onset revealed a bone marrow edema of the right L5 transverse process, centered by a linear low-intensity image on T1 and T2 weighting (Figures 1A and 1B). However, no diagnosis could be established.

Since pain kept worsening despite analgesics, a second CT scan was performed after 12 months of symptom evolution, using thinner slices on the right transverse apophysis of L5. This CT scan showed a 4 mm cortico-subcortical osteolytic lesion, centered...
by a hyperdense millimetric nidus, surrounded by an area of osteosclerosis (Figure 1C). These findings led to a diagnosis of OO of the right transverse process of L5.

However, pain was only partially and temporarily relieved with aspirin. Therefore, the patient underwent an en-bloc excision with resection of the right L5 transverse process under general anesthesia. The patient was placed in the prone position on a radiolucent table. We used a posterior midline approach to expose the L5 transverse process under microscopic control. An L5 transverse resection was performed with a bone surgical chisel with preservation of the articular capsule of the right L4–L5 articular mass. Histopathology of the intraoperative excised specimens (Figure 2A) confirmed the diagnosis of OO. Lumbar x rays and CT scan were performed the first postoperative day and confirmed the absence of residual nidus (Figures 2B and 2C). Immediate pain relief was reported, and sciatica disappeared within the first postoperative hours, with a persistent remission at 30 months follow up. No complication associated with surgery occurred, and no muscle weakness was observed.

**DISCUSSION**

We report the case of a 25-year-old woman with an OO located in the L5 right transverse process revealed by right L5 sciatica. Our case is particular by the atypical clinical presentation, the tumor’s location in the transverse process, and the occurrence in a female patient.

In fact, OO affects mainly males with a sex ratio of 3:1.4,5 Tumor distribution in spine regions varies among authors. For most authors, lumbar spine is the most commonly affected segment, followed by cervical and thoracic spine and sacrum.6 In a case series of 84 spinal OOs, posterior elements of the
spine were 3 times more affected than vertebral body. Lamina and pedicles on the vertebral arch are most commonly affected, but cases involving the transverse processes have also been reported.

The classic presentation of OO in a spinal location is a painful scoliosis or an isolated spinal night pain alleviated with the use of aspirin or nonsteroidal anti-inflammatory drugs. Radicular pain, as in our case, appears as a rare clinical picture in spinal OO. To date, there are 19 other reported cases of spinal OO revealed by radicular pain (Table). To our knowledge, only 2 cases of OO of the transverse process revealed by radicular pain have been reported.

Despite advances in diagnostic technology, the average duration before establishing a diagnosis is 15 to 20 months. Depicting spinal OO on x rays is not an easy task, especially when x rays are performed in early stages of symptoms evolution, where radiographs are often normal. In more advanced conditions, we might identify the common aspect of OO, which is an area of bone sclerosis surrounding a small radiolucent central region corresponding to the nidus. However, in our case, radiographs were still normal after a year of disease evolution. This might be explained by the complex anatomy of the spine and the small size of the tumor, making the osteolytic lesion almost not visible on plain radiographs.

MRI and CT are more sensitive than x rays for detecting soft tissue and bone marrow abnormalities adjacent to OO. However, in our case, MRI could not individualize the nidus, which was drowned in edema and osteosclerosis. Moreover, when the nidus is not identified, images of perilesional edema provided with MRI might be confused with an aggressive tumor-evoking aspect or with images of bone infection. In such cases, CT scan might redress the diagnosis. CT scan remains the most specific diagnostic imaging modality to analyze the nidus with the varying degrees of perinidal sclerosis.

Our case emphasizes the value of CT scan in the diagnosis of OO by giving thin cross-sectional images of the suspected area. In a case reported by Ono et al., CT with 1-mm slices showed the nidus with a clear, round osteolytic lesion ossification inside, leading to a diagnosis of OO, although the nidus had been overlooked in CT with 5-mm slices because of its small size, and the patient was diagnosed with lumbar spondylolysis on initial MRI and CT.

Spinal OO can be treated either with surgical intervention or minimally invasive interventions. Surgical options include intralesional curettage or marginal or en-bloc excision. Minimally invasive interventions consist of radiofrequency thermocoagulation (RFA) or CT-guided core excision. RFA has been proven to be effective and safe, with less complications and shorter hospitalization than surgical treatment. However, RFA cannot be performed safely if there is neurological impairment, if cortical bone around CT lesions is not intact, or if the cerebrospinal fluid between the lesion and nerve root or spinal cord is not sufficient (<1 mm) on MRI. In cases like ours in which osseous tumor infiltration produces radicular nerve root irritation and radicular symptoms, wide surgical excision of the tumor and surrounding tissue is the treatment of choice. Moreover, en-bloc excision is a guarantor of lower risk of recurrence than piecemeal excision, curettage, and...
<table>
<thead>
<tr>
<th>Ref No.</th>
<th>Author</th>
<th>Year of Publication</th>
<th>No. Patients</th>
<th>Age at Diagnosis (y)</th>
<th>Duration of Symptoms Before Diagnosis (mo)</th>
<th>Radiculopathy Was a Revealing Symptom</th>
<th>Imaging Tool Used for Diagnosis</th>
<th>Localization</th>
<th>Treatment</th>
<th>Treatment Complications</th>
</tr>
</thead>
<tbody>
<tr>
<td>6</td>
<td>Etemadifar et al</td>
<td>2015</td>
<td>6 (among 19 spinal OO)</td>
<td>F 26</td>
<td>52</td>
<td>No</td>
<td>NP</td>
<td>L3 (pedicle)</td>
<td>Open intralesional excision with posterior approach</td>
<td>No recurrence or neurologic deficit after surgical excision among the 19 cases</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>M 23</td>
<td>F 22</td>
<td>5</td>
<td>No</td>
<td></td>
<td>T11 (pedicle)</td>
<td></td>
<td></td>
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<td></td>
<td></td>
<td></td>
<td>F 33</td>
<td>M 23</td>
<td>6</td>
<td>No</td>
<td></td>
<td>C6 (lamina)</td>
<td></td>
<td></td>
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<td></td>
<td></td>
<td></td>
<td>M 13</td>
<td>F 13</td>
<td>5</td>
<td>No</td>
<td></td>
<td>L4 (facet)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>Janin et al</td>
<td>1981</td>
<td>2</td>
<td>F 15</td>
<td>1.5</td>
<td>Yes (C6 radiculopathy)</td>
<td>CT scan</td>
<td>C5 (pedicle)</td>
<td>Excision of pedicle and adjacent facets</td>
<td>No recurrence</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>M 35</td>
<td>Yes (sciatica)</td>
<td>CT scan</td>
<td>L3 (superior articular facet)</td>
<td></td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>10</td>
<td>Gokce et al</td>
<td>2013</td>
<td>1</td>
<td>M 18</td>
<td>6</td>
<td>Yes (C7-C8 brachial neuralgia)</td>
<td>X rays, MRI and CT scan</td>
<td>T1 (transverse process-lamina junction)</td>
<td>NP</td>
<td></td>
</tr>
<tr>
<td>11</td>
<td>Zenmyo et al</td>
<td>2011</td>
<td>2</td>
<td>M 15</td>
<td>NP</td>
<td>Yes (left sciatica)</td>
<td>MRI and CT scan</td>
<td>S1 (lamina)</td>
<td>Surgical curettage of the lesion</td>
<td>NP</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>M 35</td>
<td>Yes (T10 radiculopathy)</td>
<td>Radionuclide bone scan, MRI and CT-scan (x rays revealed only few distinct abnormalities in the lumbarosacral spine)</td>
<td>T11 (superior articular process)</td>
<td>En-bloc resection of superior articular process</td>
<td>NP</td>
<td></td>
<td></td>
</tr>
<tr>
<td>12</td>
<td>Nebreda et al</td>
<td>2017</td>
<td>1</td>
<td>M 20</td>
<td>18</td>
<td>Yes (left sciatica)</td>
<td>MRI; technetium-99m total body bone scan; CT scan</td>
<td>L4 (transverse process)</td>
<td>Surgical tumor resection of the transverse process</td>
<td>No recurrence at 6-mo follow up</td>
</tr>
<tr>
<td>13</td>
<td>Sheng et al</td>
<td>2018</td>
<td>1</td>
<td>M 38</td>
<td>120</td>
<td>Yes (left sciatica)</td>
<td>CT scan and MRI</td>
<td>L5 (appendix)</td>
<td>Pedicle and lamina resection with internal fixation of L5.</td>
<td>No follow up</td>
</tr>
<tr>
<td>14</td>
<td>Ozaki T et al</td>
<td>2002</td>
<td>2 (among a group of 9 patients with spinal OO)</td>
<td>M 12</td>
<td>24</td>
<td>NP (radiating pain in the upper arm)</td>
<td>NP</td>
<td>C5 (lamina)</td>
<td>Resection</td>
<td>No recurrence at 123-mo follow up</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>M 14</td>
<td>NP (pain in the buttock and thigh)</td>
<td>NP</td>
<td>L5 (facet)</td>
<td>Resection and spondylodiscitis</td>
<td>No recurrence at 35-mo follow up</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ref No.</td>
<td>Author</td>
<td>Year of Publication</td>
<td>No. Patients</td>
<td>Sex</td>
<td>Age at Diagnosis (y)</td>
<td>Duration of Symptoms Before Diagnosis (mo)</td>
<td>Radiculopathy Was a Revealing Symptom</td>
<td>Imaging Tool Used for Diagnosis</td>
<td>Localization</td>
<td>Treatment</td>
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<tr>
<td>15</td>
<td>Gadiya</td>
<td>2019</td>
<td>2 M</td>
<td>29</td>
<td>NP</td>
<td>10</td>
<td>Yes (C5 radiculopathy)</td>
<td>X rays, CT scan, and MRI.</td>
<td>C4 (facet and lamina)</td>
<td>Hemilaminectomy by posterior approach with complete excision of nidus and posterior instrumented fusion of C4-C5 segment with the help of the left C4-C5 lateral mass fixation.</td>
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<td></td>
<td></td>
<td></td>
<td>F 29</td>
<td>10</td>
<td>Yes (C7 radiculopathy)</td>
<td>CT scan and MRI (x rays were normal)</td>
<td>C6 (pedicle)</td>
<td></td>
<td></td>
<td>CT-guided excisional biopsy with excision of nidus. 13 mo later, due to recurrence of symptoms, an hemilaminectomy was performed. No fusion was needed.</td>
</tr>
<tr>
<td>16</td>
<td>Wang et al</td>
<td>2016</td>
<td>1 M</td>
<td>20</td>
<td>2</td>
<td>Yes (sciatica)</td>
<td>CT scan and MRI (x rays showed indistinct lesion of L5)</td>
<td>L5 (lateral recess and the intervertebral foramen)</td>
<td>NP</td>
<td>NP</td>
</tr>
<tr>
<td>17</td>
<td>Dhaliwal et al</td>
<td>2014</td>
<td>1 F</td>
<td>33</td>
<td>18</td>
<td>Yes (T11 radiculopathy)</td>
<td>CT scan and MRI (plain radiographs were normal)</td>
<td>T11 (postero-inferior angle of the vertebra)</td>
<td>NP</td>
<td>NP</td>
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<tr>
<td>18</td>
<td>Dietzfelbinger et al</td>
<td>1994</td>
<td>1 M</td>
<td>21</td>
<td>18</td>
<td>Yes (sciatica)</td>
<td>Scintigraphy and CT scan</td>
<td>L5 (pedicle)</td>
<td>NP</td>
<td>NP</td>
</tr>
</tbody>
</table>

Abbreviations: CT, computed tomography; F, female; M, male; MRI, magnetic resonance imaging; No. patients, number of patients with spinal OO with radiculopathy; NP, not precised; OO, osteoid osteoma.
RFA, since it leads to a complete removal of the nidus.\textsuperscript{5,22}

**CONCLUSIONS**

Radiculopathy is an uncommon clinical presentation in spinal OO that might be responsible for a prevalent diagnosis delay. Although not frequent, OO should be suspected in all young patients with inflammatory spinal symptoms even if x rays are normal.

Our case emphasizes the better accuracy of CT scan in detecting OO compared to MRI and x rays. In case of any doubt, it is necessary to resort to the thin section CT scan which could be guided with MRI. Treatment depends on location. Surgical removal can be performed safely in spinal OO within posterior elements. RFA can be a good alternative if there is no neurological impairment.

**KEY MESSAGES**

- Although radicular pain revealing spinal osteoid osteomas is clinically uncommon, the diagnosis should always be suspected and imaging findings interpreted with caution especially in young patient with inflammatory and chronic symptoms.
- Osteoid osteoma may display misleading imaging findings, and it can be difficult to differentiate osteoid osteoma from other conditions such as infection or other benign or malignant tumors.
- To make the correct diagnosis, it is necessary to identify the nidus, and it is important to be familiar with the radiologic findings of osteoid osteoma and its mimics.

**REFERENCES**


**Disclosures and COI:** All authors declare that they have participated in the design and analysis of the paper and that they have approved the final version. Additionally, there are no conflicts of interest in connection with this paper. We certify that the manuscript is original, not published previously, and not under concurrent consideration elsewhere.

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