She reported neck stiffness without associated neurological symptoms. Physical examination revealed a soft, non-tender, uniform mass without surface abnormalities. Computed tomography (CT) and magnetic resonance imaging (MRI) showed a circumscribed mass involving the posterior neck muscles in the midline between the C2 and C6 spinous processes with a large calcified irregular component. The mass with dual components was totally removed under general anesthesia and no recurrence was observed after 6 months of follow-up. We also reviewed the clinicopathologic features of previously reported osteolipomas in the literature and suggest that although osteolipoma is a rare variant of lipoma, it should be considered in the differential diagnosis when a lipoma of the posterior neck mixed with a bony component is encountered.

Key Words: Lipoma · Osteolipoma · Ossified lipoma · Ossifying lipoma · Lipomatous tumor.

INTRODUCTION

Lipoma is a common benign soft tissue neoplasm that sometimes may have mixed tissue components. Lipomas with mixed components are named according to the type of tissue. Ossification of a lipoma was first described in 1959, and it is rarely reported. Several names have been used to describe ossification of lipomas, including secondary calcification, ossified lipoma, ossifying lipoma, or osteolipoma, and some authors have used these terms interchangeably. As with classic lipomas, lipomas with ossifications may be found in any part of the body, but are usually found in the head, neck, oral cavity, and extremities, adjacent to bone. Only two cases of osteolipoma arising in connection with vertebrae have been described. Here, we present a case of an intramuscular osteolipoma on the posterior neck, independent of vertebrae, and review the clinicopathologic features of osteolipoma previously published in the literature.

CASE REPORT

A 51-year-old female presented with a 5-year history of a painless, progressively enlarging mass on the posterior neck. She reported neck stiffness without associated neurological symptoms. Physical examination revealed a soft, non-tender, uniform mass without surface abnormalities. Computed tomography (CT) and magnetic resonance imaging (MRI) showed a circumscribed mass involving the posterior neck muscles in the midline between the C2 and C6 spinous processes with a large calcified irregular component. The mass with dual components was totally removed under general anesthesia and no recurrence was observed after 6 months of follow-up. We also reviewed the clinicopathologic features of previously reported osteolipomas in the literature and suggest that although osteolipoma is a rare variant of lipoma, it should be considered in the differential diagnosis when a lipoma of the posterior neck mixed with a bony component is encountered.

Key Words: Lipoma · Osteolipoma · Ossified lipoma · Ossifying lipoma · Lipomatous tumor.
The patient was diagnosed with an intramuscular osteolipoma of the posterior neck and no recurrence was observed at the 6-month follow-up.

DISCUSSION

A lipoma is a common soft tissue neoplasm, but lipomas with distinct osseous components are rare. As noted by Heffernan et al., in Allen’s review of 635 cases, less than 1% of lipomas were ossified. Further, as reported by de Castro, et al., when Furlong analyzed histological findings from a group of 125 lipomas of the oral and maxillofacial region, there was no variant that presented as osseous metaplasia. Lipomas presenting with osseous components have been reported as secondary calcification of lipoma or ossified lipoma. Secondary calcification of lipoma involves calcification that progresses from the periphery inwards as a result of necrosis after impaired blood supply or repeated microtrauma. An ossified lipoma is defined as a histologic variant of lipoma that has undergone osseous metaplasia, and can be classified into 2 groups according to the predominance of the osseous component: ossifying lipoma and osteolipoma. If the adipose component of the lipoma is more dominant than the osseous component, it is an ossifying lipoma, whereas an osteolipoma presents with a dominant osseous component. The terms have been used interchangeably.

Osteolipomas have been reported in middle aged or elderly patients presenting with large painless masses with a long duration, even years, and they may be found incidentally. There are no reports indicating gender predilection. They have been reported at various sites located adjacent to bone or periosteum, including upper and lower extremity, oral mucosa, soft tissue of neck, and intracranial regions, including tuber cinereum, hypothalamus, suprasellar cistern, and the interhemispheric area. Only 2 cases of spinal osteolipoma have been described. One patient was an 8-year-old female who presented with a progressively enlarging swelling over the lumbar region with a dermal sinus tract, and other was 20-year-old female with a cervical osteolipoma contained within the spinal canal. Both lesions had some continuity to adjacent bone. In the case presented here, the osseous lipoma was located in the deep neck musculature, independent of bony vertebral structures.

There are several hypotheses regarding the pathogenetic mechanisms that influence the osseous metaplasia of lipomas, but the details are still not clear. There are two generally accepted theories for osseous metaplasia. First, these tumours may originate from multipotent mesenchymal cells that can differentiate into lipoblasts, chondroblasts, fibroblasts, angio blasts, and osteoblasts. This could account for the several lipomatous variations of classic lipomas, i.e. lipoblastoma, lipomatosis, fibrolipoma, angiolipoma, myolipoma, myelolipoma, hibernoma, and atypical lipoma, and mesenchymoma is defined as a rare soft tissue lesion that is composed of fibrous tissue associated with two or more types of well differentiated cell lines.

Fig. 1. Preoperatively CT (A : sagittal) and sagittal MRI (B : T1-weighted, C : T2-weighted) revealing an irregular calcification involving posterior neck within soft tissue mass compatible with fat tissue.

Fig. 2. Whole body bone scan (A : anterior, B : posterior) shows an amorphous calcification mass with increased uptake, and no bone metastases.

Fig. 3. Grossly, the tumor consists largely of fat and calcification presented with the red bone marrow (black arrow).
mesenchymal cells that would not normally be found in the same area\textsuperscript{2-3}. Second, ossification may also have been induced by poor nutritional supply in the centre of a large lipoma after repetitive trauma, metabolic changes, or ischemia leading to transformation of fibroblasts into osteoblasts\textsuperscript{4,5}. Fritchie et al\textsuperscript{6} reported cytogenetic analyses of three osteolipomas and reported that the translocations in all cases were consistent with the karyotypic features of lipoma. Thus, osteolipoma is a variant of classic lipoma.

Other benign tumors that may contain bone including teratomas or dermoids, masses with secondary ossification due to trauma, liposarcomas with metaplastic changes, or congenital malformations, should be considered in the differential diagnosis. In addition, tumour calcinosis, calcification in a bursa, and other conditions such as ossifying fibromas, myositis ossificans, and osteosarcomas should be taken into consideration. The use of CT scanning provides excellent visualization of the calcified or ossified components of a lipoma and confirmation of proximity to adjacent bone, and MR imaging can also provide detailed information that is useful for further evaluation\textsuperscript{2,4,6-11}.

Definitive diagnosis of osteolipoma can be made with histopathologic findings after surgical excision, which is usually the recommended treatment. A histopathologic appearance of diffuse, mature ossification within adipose tissue and gross features demonstrating a dominant osseous component confirm the diagnosis. Histologically confirmed osteolipomas are benign neoplasms, as with classic lipomas, and do not recur\textsuperscript{2,4-6,11}.

CONCLUSION

To the best of our knowledge, it is the first case report of an intramuscular osteolipoma within the posterior neck independent of adjacent bone. We suggest that although osteolipoma is a rare variant, it is important to keep it in mind when a soft tissue mass of the posterior neck mixed with osseous component is encountered.

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