Multiple ossified spinal meningiomas in the thoracic spine: A case report and literature review

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Background: Ossified spinal meningioma (OSM) is a rare form of a spinal tumor. The surgical strategies and pathologic findings related to OSM have been investigated in recent years. However, multiple OSMs are rarely reported. Here, we intend to present a rare case of multiple OSMs and review the relevant published literature.

Case Presentation: A 76-year-old woman experienced a progressive sensorimotor disturbance in her bilateral lower limbs for the past 2 years. She complained of inability to walk, urinary incontinence, and chronic constipation when referred to our hospital. A neurological examination revealed a diminished sensation below the bilateral T7, and her neurological status was Nurick Grade 6. Magnetic resonance imaging (MRI) revealed multiple intradural-extramedullary neoplasms at the T7–T11 level. Computed tomography (CT) scans showed five high-density masses of varying sizes in the spinal canal at the T7–T12 level. The patient underwent tumor resection through T7–T11 laminectomy. A histopathological examination revealed multiple OSMs.

Conclusion: We reported a rare case of multiple OSMs in an elderly patient. After one-stage complete resection, the patient recovered with satisfactory curative effect. Although elderly patients will face various postoperative complications due to their poor physical condition, we still recommend one-stage complete resection of multiple OSMs to reduce recurrence.

Keywords: multiple meningiomas, spinal meningioma, ossification, calcification, surgery

Introduction

Meningioma is the second most common of all primary spine tumors, accounting for 25%–45% (1). Multiple meningiomas usually occur inside the cranium and occasionally in both the skull and the spinal canal (2). While intracranial meningiomatosis presents as solid, cystic, or solid-cystic lesions, ranging between WHO Grade I and III, multiple spinal meningiomas are usually homogeneously solid...
WHO Grade I lesions (3). Ossified spinal meningioma (OSM), a rare form of meningioma, accounts for 0.7%–5.5% of all spinal meningiomas (4–6). However, only one case of multiple OSMs has been reported to date (7). Here, we present the first exclusive case of multiple OSMs with more than two independently situated meningiomas in a 76-year-old woman and a review of the literature.

Case presentation

Medical history

A 76-year-old woman reported a history of progressive sensorimotor disturbance in both lower limbs for the past 2 years. Initially, the patient’s condition was misdiagnosed as “lumbar disc herniation” in the local hospital, and she received conservative treatment. However, her symptoms did not improve and became more serious. When referred to our hospital, she complained of inability to walk and bladder and bowel control loss. A neurological examination revealed a diminished sensation below the T7 level. The strength of both lower limb muscles was 1/5. Her neurological status was diagnosed as Nurick Grade 6. Magnetic resonance imaging (MRI) revealed multiple intradural extramedullary tumors at the T7–T11 level compressing the spinal cord (Figure 1A). Computed tomography (CT) scans showed five high-density masses of varying sizes in the spinal canal at the T7–T12 level, and the dural sac was significantly compressed (Figures 1B,C). Three large high-density masses (13.4 mm*8.4 mm, 21.7 mm*13.0 mm, and 14.9 mm*12.5 mm) were observed in the T7–T9 segment, located in the dorsal left rear spinal cord with clear boundaries and ossification signals (Figures 1D–F). The other two high-density masses (8.7 mm*5.3 mm and 11.1 mm*6.73 mm) of different sizes were found in the T10–T12 segment.

Surgical management

The patient underwent tumor resection through T7–T11 laminectomy. The tumors occupied >90% of the transverse diameter of the spinal canal and were entirely ossified. As the vertebral fenestration was too small, it was challenging to remove the tumors. Hence, the bilateral facet joints were removed to enlarge the window. To maintain the stability of...
the spine, we performed T7–T12 long-segment pedicle screw fixation and posterolateral bone graft fusion simultaneously. Bilateral facetectomy provides advantages over simple laminectomy or laminoplasty in terms of width of the operative corridor and long-term preservation of the spinal alignment. As the dura mater had severely adhered, a fine right-angled hook was used to dissect the neural tissue dorsally away from the neoplasms, which were then resected en bloc, together with parts of the dura mater and arachnoid. After a complete separation of multiple OSMs, the dura mater became normal and it was sutured. The drainage was routinely placed, and the wound was closed layer by layer.

**Histopathological outcomes**

The paraffin sections were stained with hematoxylin and eosin (H&E). A histopathological examination revealed a large number of psammoma bodies in the stroma with significant signs of ossification [World Health Organization (WHO) Grade I]. In addition, there were many mature bone tissues around the tumor cells, including trabecular bone, as well as bone marrow with hematopoiesis (Figures 2A–C). Immunohistochemical findings revealed EMA (+), Vim (+), P53 (−), S-100 (+), and Ki67 (approximately 5%) (Figures 3D–F).
Postoperative and follow-up results

Postoperative neurological improvement was significant, and no complications were found. The patient was discharged 2 weeks after surgery. After the 2-year follow-up, she was able to walk without assistance. Her neurological status recovered to Nurick Grade 3. CT scans showed a total resection of the tumors, and there was no recurrence 2 years after surgery (Figures 3A–E).

Discussion

Including our current report, 33 articles (4, 7–37) containing 43 cases of OSMs have been published as of this year, according to PubMed. (In Table 1, the search terms are (((ossified [Title/Abstract]) OR osteoblastic [Title/Abstract])) OR (osseous metaplasia [Title/Abstract])) OR (psammomatous [Title/Abstract]) OR (calciﬁed [Title/Abstract])) AND (spinal [Title/Abstract]) AND (meningioma [Title/Abstract]), and 76 potential studies are identiﬁed. We also checked the reference lists of all, including articles, to add 10 other articles.) For this condition, female predominance has been clearly noted (female, 38; male, 5), and the average age is 58.4 years, ranging from 15 to 90. Most tumors are located in the thoracic spine, except four in the cervical region and one in the lumbar region. Bone formation and hematopoiesis have been found in 7 cases, and only 2 cases of multiple OSMs have been identiﬁed, including our report.

Multiple meningiomas are deﬁned as more than two independently situated meningiomas arising simultaneously or sequentially (2). The pathogenesis of multiple meningiomas remains elusive. In fact, at present, two kinds of hypotheses are proposed by the relevant literature: one supports the theory of monoclonal spreading and the other suggests that of clonally unrelated onset for lesions arising sequentially from two clearly distinct spinal regions (38). A genomic proﬁling study reveals that multiple meningiomas can be of both mono- and multiclonal origin. Even monoclonal multiple meningiomas can acquire intertumor heterogeneity through branched evolution resulting in pathology, and the landscape of one tumor may not be representative of the others. Thus, multiple meningiomas should be tailored individually if feasible (39).

The concept of ossiﬁcation of meningioma should be distinguished from calciﬁcation. Calcification is more a radiologic description than a histopathological diagnosis and is commonly seen in psammomatous meningioma. Based on the classiﬁcation by the World Health Organization (WHO), ossiﬁcation in meningioma is histologically classiﬁed as a subtype of metaplastic meningioma and is characterized by the expression of mesenchymal components (40). The occurrence of OSM also remains unclear. One hypothesis is that ossiﬁcation results from the repeated accumulation of hydroxyapatite crystals in psammoma bodies (41). However, some reports (4, 8, 10, 15, 21) have occasionally found that ossiﬁcation can occur without psammoma bodies, which may not support the hypothesis. Hence, most studies prefer to believe that ossiﬁcation is secondary to the metaplasia of arachnoid cells and interstitial cells, which induce the synergistic effect of osteoblast, ﬁbroblast, and angiogenesis components in bone tissue formation (4, 8, 10, 23). Up to now, seven studies have mentioned the formation of hematopoietic tissue in OSM. This unexpected ﬁnding seems to conﬁrm the mesenchymal potential of meningiomatous cells, which may undergo bone metaplasia (9, 11–13, 19, 24).

Although OSM was previously reported to grow very slowly and experience a long asymptomatic period, the symptoms may appear at an early stage in intraspinal tumors compared with intracranial tumors due to a smaller space in the intraspinal canal (15). As most commonly in WHO Grade I, multiple spinal meningiomas can be considered benign lesions with a good prognosis, especially if the lesion and affected dura are macroscopically complete removed (Simpson grade I) (42). Mirimanoff et al. (43) found that the incidence rates of secondary operation after a total resection for follow-up periods of 5, 10, and 15 years were 6%, 15%, and 20%, respectively. In contrast, after a subtotal resection, the probability was 25%, 44%, and 84%, respectively.

Compared with the common meningiomas, OSM closely adheres to the dura and arachnoid (7). If the tumor and the dura mater are not separated, they could be removed together and repaired with an artiﬁcial dura mater covered with gelatin sponge. Ruggeri et al. (44) proposed that a poor surgical outcome in patients with ossiﬁed tumors results from a more “invasive” surgical removal of an ossiﬁed mass: an internal debulking is not feasible for a hard tumor. In our case, the ossiﬁed tumors occupied the entire transverse diameter of the spinal canal and could be approached more safely with laminectomy by removing the bilateral facet joints from T7 to T12 instead of with hemilaminectomy: it offers a wider operative ﬁeld for a better manipulation of the neural tissue, especially the thoracic spinal cord, which is more susceptible to damage than the cervical spinal cord and lumbar spinal roots. Therefore, we performed a T7–T12 long-segment pedicle screw ﬁxation and posterolateral bone graft fusion to prevent the occurrence of iatrogenic kyphosis and maintain the stability of the spine. After a 2-year follow-up, no complications such as internal ﬁxation failure and kyphosis occurred. The use of intraoperative ultrasound represents a valuable surgical aid for real-time neuronavigation, allowing the operating team to evaluate the decompression of the spinal cord and rule out subdural blood clots at the time of dural closure/reconstruction (45). For spinal intradural extreemullary lesions, the sensitivity, speciﬁcity, and positive
and negative predicted values of intraoperative neurophysiological monitoring (IONM) are reported to be 75, 100, 100, and 97%, respectively. This indicates that IONM predicts neurological deficits with high accuracy, although its role in preventing new neurological deficits in spinal meningiomas has yet to be proved (46). The current trends in the use of adjuvant radiotherapy and radiosurgery for spinal meningiomas have increased; nonetheless, this has not led to

### TABLE 1 Summary of ossified meningioma cases.

| Study               | Age/Sex | Level     | Tumor number | Symptoms     | Histological characteristics                       |
|---------------------|---------|-----------|--------------|--------------|-----------------------------------------------------|
| Roger (18)          | 16/F    | T9        | 1            | Myelopathy   | Psammoma bodies, bone cells                         |
| Friedberg (29)      | 69/F    | T1–T2     | 1            | Myelopathy   | Psammoma bodies, mature cancellous bone             |
| Kandel et al. (30)  | 17/F    | T8        | 1            | Myelopathy   | Meningotheliomatous, psammoma bodies, bone spicule  |
| Nijima et al. (4)   | 75/F    | T8–T9     | 1            | Myelopathy   | Psammoma bodies, bone spicule                       |
| Kitagawa et al. (32)| 75/F    | T9–T10    | 1            | Myelopathy   | Psammoma bodies, bone tissue                        |
| Nakayama et al. (33)| 74/F    | T9        | 1            | Myelopathy   | Matured lamellar bone tissue                        |
| Huang et al. (34)   | 73/F    | T5        | 1            | Myelopathy   | Psammoma bodies, bone marrow                        |
| Saito et al. (31)   | 54/F    | T11       | 1            | NA           | Metaplastic (osseous)                               |
| Naderi et al. (35)  | 15/M    | T4        | 1            | Myelopathy   | Psammoma bodies, mature bone tissue                 |
| Liu et al. (36)     | 70/F    | T11       | 1            | Myelopathy   | Psammoma bodies, bone tissue                        |
| Hirabayashi et al.  | 82/F    | L3        | 1            | Cauda equina syndrome osseous                       |
| Tahir et al. (8)    | 40/F    | T6        | 1            | Myelopathy   | Mineralized bone                                    |
| Uchida et al. (7)   | 76/F    | T8 and T12| 2            | Myelopathy   | Psammoma bodies, mature bone                        |
| Lici et al. (9)     | 58/F    | T6        | 1            | Myelopathy   | Psammoma bodies, lamellar bone tissue, hematopoiesis|
| Chotai et al. (12)  | 61/F    | T4–T5     | 1            | Myelopathy   | Psammoma bodies, mature lamellar bone, hematopoiesis|
| Ju et al. (10)      | 61/F    | T9–T10    | 1            | Myelopathy   | Heterotopic ossification                            |
| Taneoka et al. (11) | 78/F    | T9        | 1            | Myelopathy   | Psammoma bodies, mature bone, hematopoiesis         |
| Yamane et al. (14)  | 61/F    | T12       | 1            | Myelopathy   | Psammoma bodies, cancellous bone with bone marrow   |
| Chan et al. (13)    | 64/F    | T9–T10    | 1            | Myelopathy   | Psammoma bodies, bone marrow, hematopoiesis         |
| Alafaci et al. (15) | 45/M    | T2–T3     | 1            | Myelopathy   | Seven cases of osseous component in association with psammoma bodies, Two cases of immature bone trabeculae |
|                      | 75/F    | T3–T4     | 1            | Myelopathy   | Psammoma bodies, lamellar bone tissue, hematopoiesis|
|                      | 86/F    | T3–T4     | 1            | Myelopathy   | Psammoma bodies, mature bone tissue                 |
|                      | 65/F    | T7        | 1            | Myelopathy   | Psammoma bodies, bone trabeculae                    |
|                      | 72/F    | C7        | 1            | Myelopathy   | Psammoma bodies, bone trabeculae                    |
|                      | 40/F    | T1–T2     | 1            | Myelopathy   | Psammoma bodies, mature bone tissue                 |
|                      | 65/F    | T7–T8     | 1            | Myelopathy   | Psammoma bodies, mature bone tissue                 |
|                      | 40/F    | C7        | 1            | Myelopathy   | Psammoma bodies, bone trabeculae                    |
|                      | 41/F    | T2–T3     | 1            | Myelopathy   | Psammoma bodies, bone trabeculae                    |
| Demir et al. (17)   | 26/F    | T9–T11    | 1            | Myelopathy   | Psammoma bodies                                    |
| Cochran et al. (19) | 47/F    | T8        | 1            | Radiculopathy | Psammoma bodies, bone marrow, hematopoiesis         |
| Xia and Tian (16)   | 90/M    | T10–T11   | 1            | Spinal cord injury after fall                       | Psammoma bodies, bone trabeculae                    |
| Prakash et al. (20) | 60/F    | T7–T8     | 1            | Myelopathy   | Psammoma bodies, immature bony trabeculae          |
| Sakamoto et al. (21)| 57/F    | C7        | 1            | Myelopathy   | Osseous core, fibrous                              |
| Murakami et al. (23)| 29/F    | T12       | 1            | Back pain, leg numbness                            | Psammoma bodies, mature bone tissue                 |
| Taha et al. (22)    | 22/F    | T4–T5     | 1            | Myelopathy   | Psammoma bodies, bone trabeculae                    |
| Wang et al. (24)    | 52/F    | T4        | 1            | Back pain                                           | Psammoma bodies, immature trabecular bone, hematopoiesis|
| Xu et al. (25)      | 85/F    | T11       | 1            | Back pain, leg pain                                | Psammoma bodies                                    |
| Buchanan et al. (26)| 64/M    | T4        | 1            | Myelopathy   | Psammoma bodies, bone formation, osseous metaplasia |
| Wong et al. (27)    | 75/F    | T10–T11   | 1            | Myelopathy   | Psammoma bodies, immature trabeculae bone          |
| Thakur et al. (28)  | 74/F    | T8        | 1            | Tingling paresthesia                               | Psammoma bodies, bony hard-tissue fragments        |
| Present case        | 76/F    | T7–T12    | 5            | Myelopathy   | Psammoma bodies, bone trabecular bone, hematopoiesis|
a significant increase in overall survival rates (47). Larger tumor size and borderline or malignant behavior are reported to be associated with increased radiation use; the introduction of multimodal adjuvant technologies such as radioenhancers has yet to provide evidence for superior outcomes (48). To sum up, surgical en bloc resection of OSM is difficult, especially in multiple OSMs, but it is still the best treatment.

Although ossified meningioma could be identified in a plain radiograph, CT and MRI are complementary methods of diagnosing a calcified spinal meningioma, especially in cases replaced entirely by calcification (49). On MR images, the signal intensities of calcifications within the masses were variable, and the extent of signal intensities suggesting calcification did not concur with that of calcified foci within a mass, as seen on CT images (49).

Conclusion

In the current report, we present a rare case of multiple OSMs in the thoracic spine. Although the mechanism of occurrence is not clear, a total resection is generally required, and a satisfying prognosis and a low recurrence rate can be expected. The surgical strategy for OSMs differs from that for other meningiomas. We suggest that preoperative CT be used to accurately locate ossification. Intraoperative use of a wide surgical corridor with total laminectomy, combined with bilateral facet joint resection, identification of upper and lower poles, and early CSF drainage, is helpful in decreasing neural retraction.

Data availability statement

The original contributions presented in the study are included in the article/Supplementary Material, further inquiries can be directed to the corresponding author.

Ethics statement

Ethical review and approval was not required for the study on human participants in accordance with the local legislation and institutional requirements. The patients/participants provided their written informed consent to participate in this study. Written informed consent was obtained from the individual(s) for the publication of any potentially identifiable images or data included in this article.

Author contributions

CD and YL contributed to the conception and design of the study. CD wrote the first draft of the manuscript. YL and HW wrote sections of the manuscript. YZ did the literature review. YM was involved in the concept development, quality control of the data, and interpretation of the manuscript. All authors contributed to the article and approved the submitted version.

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Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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