INTRAMEDULLARY THORACIC MENINGIOMA: A RARE CASE REPORT AND REVIEW OF THE LITERATURE

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ABSTRACT

Only a few cases of intramedullary meningiomas in the cervical and thoracic region have been reported in the literature. We present a rare case of intramedullary meningioma in the lower thoracic region.

A 68-year-old woman with complaints of back pain, gait disturbance, and paresthesias in both lower extremities for 4 months was admitted to our institution. Initially, she presented with weakness in her legs that gradually deteriorated over time. The patient had been operated on for breast cancer 2 years prior to the onset of current complaints. The neurological examination revealed spastic inferior paraparesis (McCormick grade IV), hypoesthesia below Th10 dermatome, bowel and bladder disturbances. Magnetic resonance imaging of the thoracic spine demonstrated intramedullary tumor that was hypointense on T1 and T2-weighted images but showed homogenous enhancement after gadolinium administration. After gross total resection, the histopathological examination confirmed the diagnosis of atypical intramedullary meningioma.

CONCLUSION: The intramedullary meningiomas are extremely rare, but they should not be excluded from the differential diagnosis, especially in cases of intramedullary tumors with homogenous enhancement on magnetic resonance imaging. The best treatment strategy is gross total removal of the tumor.

Keywords: Intramedullary meningioma; atypical meningioma, MRI, surgery thoracic spine,

INTRODUCTION

Spinal meningiomas can occur in each segment of the spine, but they are most common in the thoracic region. Intradural extramedullary meningiomas account for 25% to 50% of all intradural extramedullary tumors, and the incidence is 0,5-2/100 000 per year [1]. Intra-extradural spinal meningiomas represent 5% - 6% of the cases [2]. Only a few cases of intramedullary meningiomas have been reported in the literature [3].

We present a rare case of intramedullary meningioma in the lower thoracic spine and put stress on its clinical presentation, imaging findings, surgical techniques, and pathological results.
Fig. 2. MRI before and after gadolinium enhancement: A/ T1 MRI without enhancement - isointense lesion. B/ T1 after enhancement – homogenous enhancement of the tumor mass.

Taking into account the history of oncological disease, the lesion was initially interpreted as intramedullary metastasis. The patient underwent a T11 – T12 laminectomy, midline durotomy and ultrasonic localization of the lesion. After posterior midline myelotomy, a pale grayish tumor was visualized that had rubbery consistency (Fig. 3 A). Using microsurgical technique, gross total resection of the tumor was achieved (Fig. 3 B).

Fig. 3. Intraoperative photos. A/ Posterior myelotomy B/ Post resection of the tumor.

The histopathological examination confirmed a tumor with a presence of small, atypical, hyperchrome cells expressing high mitotic index, macronucleoli, and hypercellularity. The diagnosis of atypical meningioma (Grade II WHO) was confirmed (Fig. 4).

Fig. 4. Atypical meningioma X-EX400 (WHO Grade II). Increased cellularity (white arrow); Small atypical cells with high N/C ratio (black-white arrow); Large and prominent nucleoli (white arrowhead).
In the postoperative period, the patient demonstrated favorable recovery of the neurological deficit. At the second postoperative week, she was able to walk assisted. At the 8th postop month, she died due to progression of the systemic breast cancer disease.

**DISCUSSION**

Intradural extramedullary meningiomas account for 25% – 46% of the spinal tumors [3]. The peak incidence of spinal meningiomas is between the 5th and the 7th decade, but they can occur at any age. Women are commonly affected (75-80% of the cases), and the female: male ratio is 4:1-8:1 [2].

Spinal meningiomas demonstrate homogenous enhancement after gadolinium administration and appear mildly hyperintense on T1- and T2-weighted MRI images [4]. Another typical characteristic is the so-called “dural tail sign” observed on post-contrast T1 MRI series but in the case of intradural extramedullary localization [3].

According to Klekamp, the intramedullary tumors represent about 5% of all spinal tumors, while intramedullary meningiomas are under 1% [5, 6].

Because of their rarity, the imaging characteristics of intramedullary meningiomas are not well described. In most cases, they are iso- or hypointense on T1 sequences and mildly hyperintense on T2 sequences (with homogeneous enhancement?) [7]. This is also confirmed in our clinical case.

The pathogenetic mechanism of development is still unknown. According to Moriuchi et al., meningocytes are modified fibroblasts deriving from mesenchymal cells. Therefore, intramedullary meningiomas could develop in the mesenchymal cells lining the perivascular spaces of the neuraxis [8].

There are 16 case reports of intramedullary meningiomas in the literature (Table 1) [3, 7-21]. 9 of them were men, and 7 were women with an average age of 49 years (7 – 69). According to their localization, 8 of these intramedullary meningiomas were located in the cervical region (50%), 1 in the cervicothoracic region (6,2%), and seven cases in the thoracic region (43.8%). As far as we know, our case is the first described in the lower thoracic region. This confirms that intramedullary meningiomas could affect the spinal cord at any level. In contrast to other meningiomas, the intramedullary meningiomas occur more often in the younger patient, male gender and are located predominantly in the cervical region.

**CONCLUSION**

Surgery should be the primary treatment. Total tumor resection is usually possible using a meticulous microsurgical technique. Interestingly, satisfactory results are reported after gross total or partial resections [3, 8, 10, 11, 18, 19].

The histological types of reported intramedullary meningiomas are presented in Table 1. We can summarize that most of them are not benign, consistent with WHO Grade II, a fact also confirmed by our case. Therefore, cautious MRI follow-up is needed to detect early signs of recurrence. In such cases, adjuvant radiotherapy should also be considered.

<table>
<thead>
<tr>
<th>Authors</th>
<th>Sex/Age</th>
<th>Localization</th>
<th>Pathohistological type</th>
</tr>
</thead>
<tbody>
<tr>
<td>Salvati et al. [10]</td>
<td>F/67</td>
<td>C_2 - C_4</td>
<td>Fibroblastic</td>
</tr>
<tr>
<td>Moriuchi et al. [8]</td>
<td>F/54</td>
<td>C_2 - C_4</td>
<td>Transitional</td>
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<tr>
<td>Covert, et al. [12]</td>
<td>F/69</td>
<td>T_2 - T_3</td>
<td>Anaplastic</td>
</tr>
<tr>
<td>Raza, et al. [13]</td>
<td>F/65</td>
<td>T_2 - T_6</td>
<td>Atypical</td>
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<tr>
<td>Park, et al. [14]</td>
<td>F/65</td>
<td>T_9 - T_10</td>
<td>Clear cell</td>
</tr>
<tr>
<td>Salehpour, et al. [15]</td>
<td>M/21</td>
<td>C_2</td>
<td>Syncytial type</td>
</tr>
<tr>
<td>Sahni, et al. [7]</td>
<td>M/42</td>
<td>C_3 - T_2</td>
<td>Atypical</td>
</tr>
<tr>
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<td>M/33</td>
<td>T_1 - T_3</td>
<td>Low grade</td>
</tr>
<tr>
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<td>F/7</td>
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<td>Atypical</td>
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<tr>
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<td>M/15</td>
<td>T_7 - T_8</td>
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<tr>
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<td>M/52</td>
<td>C_5</td>
<td>Whorling-sclerosing</td>
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<tr>
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<td>M/64</td>
<td>T_3 - T_4</td>
<td>Low-grade</td>
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</table>
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