A Rare Case of Primary Malignant Melanoma Presenting with Lower Extremity Weakness

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The authors report a pathologically confirmed case of primary malignant melanoma located in the intradural lumbosacral area, which is extremely rare due to its atypical location and manifestation. However rare, primary malignant melanoma could present as an intradural tumor and cause neurological deficit.

Abstract

Keywords
► intramedullary tumor
► melanoma
► retroperitoneum

Introduction

Primary malignant melanomas of the central nervous system (CNS) are extremely rare and account for about one percent of all melanomas.1

Intradural primary malignant melanoma in lumbosacral area is a very rare entity and the exact incidence is still unclear. We report a case of primary malignant intradural melanoma in the lumbosacral area which extended into the retroperitoneal space through the L5 and S1 foramina.

Case Presentation

A 47-year-old woman was admitted to our hospital with a complaint of progressive worsening of severe back pain from approximately 4 weeks ago and weakness in the right lower extremity from a week ago. No remarkable finding was noted in her past medical history. A neurological examination revealed moderate paresis and numbness on the right lower extremity. Motor strength of proximal and distal muscles of the right lower extremity was assessed to be 3/5, whereas the left lower extremity had 5/5 strength. Magnetic resonance imaging (MRI) of the lumbosacral spine showed an intradural mass in the level of L5 and S1 vertebrae which extended into the retroperitoneal space through the L5 and S1 foramina. The lesion was iso- to hypointense on T2 and slightly hypointense on T1-weighted images. Diffuse contrast enhancement was also prominent on T1-weighted images after gadolinium (GD) injection (►Fig. 1–3). The lumbosacral computed tomography (CT) scan showed an isodense lesion at the same levels that enlarged the foramina and eroded the bone (►Fig. 4).

During the operation, laminectomies were performed from L5 to S1 levels. The first interesting thing, which was seen was the severe enlargement of L5 and S1 nerve roots. A dark black vascular tumor was observed immediately after dural incision (►Fig. 5, 6). This pigmented tumor showed clear invasion of L5 and S1 roots under the operative microscope. The tumor was dissected from the rootlets and it was excised grossly as much as safely possible. The tumor extended into the retroperitoneal space through the enlarged L5 and S1 roots. The postoperative course was eventful. She had recovery in motor strength in the early postoperative period.

Histopathological sections demonstrated a highly cellular lesion composed of clusters of atypical cells (►Fig. 7). The cells had pleomorphic nuclei with irregular borders and prominent nucleoli. Some of the nuclei contained vacuoles. There was some mitosis and abundant intra- and extracellular pigments. Immunohistochemical staining revealed positive immunoreactivity for S100 protein and human melanoma black-45.

After confirmation of the diagnosis, the patient underwent a thorough systemic examination and survey. Any other foci of melanoma, including skin and orbit, could not be found.
and primary malignant melanoma was confirmed. Palliative chemotherapy was begun by the oncology service, and she died 6 months later.

Discussion

Skin is the most common origin for melanoma, but other places such as gastrointestinal system and orbit were also reported. About 10% of all cases arise from noncutaneous regions. Primary and metastatic intradural melanomas are rare.

According to the study by Fuld et al, the thoracic region is the most common place of primary spinal melanoma and about half of the cases were observed in this area. The lumbosacral region is the rare place for melanoma presentation as a primary or metastatic tumor. Metastasis from cutaneous melanoma is the most common differential diagnosis of primary melanoma. In approximately 10% of patients with secondary melanoma, the primary site is unknown. Other differential diagnoses are melanotic schwannoma and other neuroectodermal tumors, which could be excluded by negative immunohistochemical (IHC) stains for neuroendocrine markers such as (neuron-specific enolase) NSE, chromogranin, and synaptophysin.

Cutaneous and extracutaneous melanomas are histopathologically similar. S-100 protein, HMB-45, melanin A, and vimentin are used as immunohistochemical markers which help diagnosis of melanoma.

Hence S-100 has the highest sensitivity for the differentiation of melanocytes, it is used as a screening tool. However, S-100 is also found in several other tissues, including nerve sheath tumors and gliomas.

HMB-45 and melanin A are the two most common melanocyte-specific monoclonal antibodies used for the diagnosis of melanoma. The sensitivity of HMB-45 and melanin A for melanoma is approximately 93 to 100%. Vimentin is almost always present in malignant melanoma with a sensitivity of 93%.
The interesting point about melanoma histopathology is the absence of melanin pigment in approximately 50% of cases.\(^7\)

The best modality for the diagnosis of spinal cord tumors is MRI. Spinal cord melanomas are usually slightly hyperintense on T1, iso- to hypointense on T2 weighted images, and show homogeneous mild enhancement after intravenous GD injection. These features are prominently related to paramagnetic features of melanocytes.\(^{14,15}\) However, it is difficult to diagnose melanoma on imaging and the definite way to diagnose melanoma involves histopathological studies. Hence, survival of primary and secondary melanoma is different, and differentiation of primary from secondary melanoma is important.

The life expectancy of metastatic melanoma is about one year, while survival of primary melanoma is reported to be approximately 6 years.\(^2,16\) The prognosis of extracutaneous melanoma is worse than cutaneous melanoma. According to Hayward’s criteria for the diagnosis of primary CNS melanoma, the following two conditions are needed: Absence of malignant melanoma outside the CNS and histopathological confirmation.\(^2,17\)

The clinical course of melanoma is variable. Surgical resection is the first modality for primary melanomas; however, complete resection of the lesion is palliative and does not prevent further metastasis. Postoperative radiotherapy or chemotherapy was recommended by several authors. However, no evidence establishes the role of adjuvant therapy on the improvement of survival. Recently, the application of immunotherapy and molecular-targeted therapy in the treatment of malignant melanoma have improved.\(^5,6\)

We reported a pathologically confirmed case of primary malignant melanoma located in the intradural lumbosacral area, which is extremely rare due to its atypical location and manifestation. She did not show any signs and symptoms of melanoma at admission. During surgery, after laminectomy, the interesting point was the severe enlargement of L5 and S1 foramina in the affected side which enlarged the foramina. After opening the dura along with the root, the dark black vascular tumor was observed, which extended into the retroperitoneal space. The origin of the tumor is exactly unclear for us. This could originate intradurally and extended to retroperitoneal space or might be originated from retroperitoneal space and extend to intradural space through L5 and S1 foramina. Both of these possibilities are extremely rare.

**Fig. 3** Sagittal T1 MRI with GD enhancement. The tumor was enhanced diffusely. GD, gadolinium; MRI, magnetic resonance imaging.

**Fig. 4** Axial CT scan view at L5 and S1 level. The tumor expanded the right S1 foramina and extended into the retroperitoneal space. Bone erosion was seen. CT, computed tomography.
Conclusions

However rare, primary malignant melanoma could present as an intradural tumor and cause neurological deficit.

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Conflict of Interest
None declared.

References