



Case Report

Dorsal intraspinal B-cell Non-Hodgkin lymphoma in two patients

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Abstract

Non-Hodgkin Lymphoma is rarely diagnosed as spinal cord compression syndrome. Caused by an epidural mass, this complication is often encountered in the late stage of the disease. We report two cases presenting symptoms of low thoracic spinal cord compression due to an epidural tumor on the MRI. Possible complications that may occur and how to prevent adverse events during surgery are essential. This case is of special interest for being a low-incidence pathology in a rare location. The case is of high interest because of the importance of resecting the lesion to prevent sequels and the complexity of its management and highlights the importance of considering the diagnosis of this event.

Introduction

Non-Hodgkin lymphoma is a group of malignant neoplasms originating from the lymphoid tissues, mainly the lymph nodes. It originates from B cell precursors, mature B cells, T cell precursors, and mature T cells. This tumor may result due to chromosomal translocation, various toxins, infections, and chronic inflammation. It comprises various subtypes, each with different clinical features and responses to therapy. It can be mainly divided into two groups, indolent and aggressive, based on the disease's prognosis. Indolent lymphomas present with waxing and waning lymphadenopathy for many years, whereas aggressive lymphomas have specific B symptoms such as weight loss, night sweats, and fever and can result in death within a few weeks if untreated. Non-Hodgkin lymphoma arises either from B cells, T cells, or natural killer cells due to chromosomal translocation or mutation/deletion. Proto-oncogenes are activated by chromosomal translocation, and tumor suppressor genes are inactivated by chromosomal deletion or mutation. The t (14;18) translocation is the most common chromosomal abnormality in Non-Hodgkin lymphoma. This translocation results in the overexpression of cyclin D1, a cell cycle regulator [1,2]. We present two cases of intramedullary non-Hodgkin lymphoma. The comparison between the two cases is of great interest because it occurs despite being of the same cell lineage with different clinical manifestations and radiological characteristics [3]. Only 0.6% - 6.5% produce spinal neurological complications, the most prone being those with large B cells, affecting only 2/3 of the vertebral level [4-7]. Few patients with intramedullary

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involvement have been described in the literature, with epidural infiltration being the most common mechanism of spinal neurological complication [8-10].

Case presentations

They are a 77-year-old patient and another 78-year-old. The first presented with right cervicobrachialgia and paresis of the upper limbs. The spinal MRI shows a lesion with the invasion of the C7-T1-T2 vertebral bodies, involvement of the spinal canal, and foraminal stenosis, as well as injury to the pelvis and para-aortic retroperitoneal lymph nodes. The second presented gait instability and paresis in the right leg with allodynia in the left leg and urinary retention. The spinal MRI revealed a well-defined intramedullary lesion suggestive of ependymoma. In both cases, decompressive laminectomies and flavectomy were performed with biopsies, resulting in the first-grade IV large B-cell non-Hodgkin lymphoma, which received radiotherapy and R-CHOP chemotherapy with a significant reduction in tumor mass, and the second highgrade B-cell lymphoma. of malignancy/Intramedullary mantle lymphoma T4-T5. Receiving R-CHOP chemotherapy with a favorable response.

Patient 1

The patient was admitted to the hospital due to cervicaldorsal pain for a year. In the latest weeks, it has worsened,

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being of great intensity and irradiating mainly towards the right arm. Associated weakness to mobilize both arms, more marked at the distal level. The MRI of the spine shows a lesion invading the C7-D1-D2 vertebral bodies in its anterior region, with involvement of the medullary canal and marked foraminal stenosis at the exit of the C8 roots. A Total Body CT study was carried out where a pelvic lesion was observed, after ruling out the possibility of a biopsy for bone marrow study without significant findings, it was decided to transfer the lesion for a neurosurgical biopsy (Figure 1).

Dorsolumbar MRI

Two bone lesions with a soft tissue mass: one in the bodies in the first and second thoracic vertebrae (blue arrow), showing soft tissue mass centered at C7, D1, D2 and axial T1 showing epidural mass with extension into the posterior elements and lateral masses (Figure 2; blue arrow). Another lesion was found in the pubis and right ischiopubic ramus. Compatible with myeloma or metastasis. Another small lytic lesion on the left scapula, more non-specific, is of doubtful value. Three infrarenal left paraaortic retroperitoneal pathological nodes are related to the same process. Mediastinal nodule, probable ganglion.

Pulmonary nodules: One of 10 mm in the right lung, it can be a hamartoma, granuloma, or metastasis; and several millimeters in the lingula, probable granulomas and less probable metastasis.



Figure 1: Patient 1: Sagital MRI T2 STIR

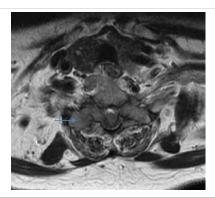


Figure 2: Patient 1: Axial MRI T2

Patient 2

The patient was admitted for scheduled surgery. For months he had presented gait instability with clumsiness in the legs and a sensation of numbness in the distal MMII, diagnosed with polyneuropathy and at the time he was evaluated by Rehabilitation. Progressive symptoms of two weeks of evolution, of leg weakness plus the right with impossibility to walk, in the last week allodynia of the left leg. No bulbar clinic or evident in lower limbs, no back pain. He wasn't capable to stand up for ten days. Urinary retention since 24h. Partial excision of intramedullary lesion T4-T5 was performed under NFS monitoring (Figure 3).

Cervicothoracic MRI

Intramedullary lesion at T4-T5 level (blue arrow), hyperintense with contrast (Figure 4; blue arrow).

Discussion

Low back pain and radicular pain are frequent complaints, with the most common cause herniated intervertebral disc. Other causes include overgrowth of bone intruding into the spinal canal, spinal stenosis, fracture-dislocation of the spine, infections and neoplasm.

The two patients reported in this paper, who were admitted with neurological signs, suffered from a malignant epidural non-Hodgkin's lymphoma. Lymphomas are a heterogeneous group of malignancies of B cells or T cells that



Figure 3: Patient 2: Sagital MRI STIR.

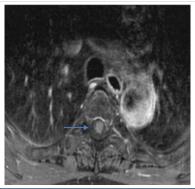


Figure 4: Patient 2: Axial MRI T1 with contrast.



usually originate in the lymph nodes but may originate in any organ of the body. Extranodal disease is an adverse prognostic factor, particularly the involvement of the Central Nervous System (CNS). Between 5% and 10% of patients with nodal presentation of lymphoma may develop CNS involvement. The intermediate or high histological type of non-Hodgkin's lymphoma and the presence of an underlying immune deficiency are the most significant risk factors for secondary CNS involvement. CNS presentations may include spinal cord compression, leptomeningeal spread, or intracerebral mass lesions. In addition, other mechanisms of neuropathy should be considered, such as the effects of chemotherapy [11-13].

Spinal cord compression is a rare presentation of non-Hodgkin's lymphoma, occurring in 0.1% to 3.3% of patients. It is commonly caused by extradural disease, either because of an isolated deposit within the spinal canal or by the extension from an adjacent nodal mass or bone involvement. Less commonly, non-Hodgkin's lymphoma may arise subdurally or within the spinal cord. Spinal cord compression typically presents with back pain, leg numbness and tingling, radicular pain followed by extremity weakness, paresis and paralysis, but can also be asymptomatic at presentation.

Conclusion

Despite the availability of new techniques for early diagnosis, it can be difficult to differentiate initially serious conditions as an isolated tumoral involvement of the CNS in the absence of widespread disease. Although CNS involvement is most frequently described in intermediate and high-grade non-Hodgkin's lymphoma, patients with a medical history of low-grade lymphoma can casually develop this kind of involvement. In patients with a history of lymphoma and neurological signs, an MRI of the spine must be performed and is the technique of choice as it is able to identify spinal cord lesions not visible by other radiographic techniques. Although spinal epidural involvement is a rare presentation of non-Hodgkin's lymphoma, it can be the only symptom of lymphoma and must be considered. Late diagnosis, location of non-Hodgkin's lymphoma in the cervical and upper thoracic region, rapid progression of neurological deficit and the appearance of paraplegia, and immunosuppression are associated with a worse prognosis. The objective of surgery is decompression and taking biopsies for anatomopathological study [14,15]. Corticosteroids, chemotherapy with a CHOP regimen, local radiotherapy and immunotherapy with Rituximab are used.

Conflict of interest

The authors declare no conflicts of interest and have adhered to the accepted ethical standards by obtaining ethical approval.

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