

**CASE OF INTRAMEDULLARY SPINAL CORD METASTASIS OF BREAST
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ABSTRACT

The intramedullary spinal cord metastasis is extremely rare and represents 8,5% of central nervous system metastasis. We report a case of intramedullary spinal cord metastasis, in a 50 years-old patient with metastatic breast carcinoma, diagnosed by magnetic resonance imaging and treated with Systemic corticosteroid therapy and palliative radiation. Through this paper, we aim to bring to literature a new case of intramedullary spinal cord metastasis of breast cancer, while developing the clinical presentation, diagnostic modalities and the management of this rare lesion.

KEYWORDS: Intramedullary spinal cord metastasis, breast carcinoma.**INTRODUCTION**

Breast cancer is the most common form of cancer in Moroccan females. Distant metastasis is common. But intramedullary spinal cord metastasis remains an unusual type. Incidence is reported as 8,5% of all central nervous system metastasis and 0,9%–5% of all spinal cord metastasis,^[1,2] in the literature.

Most often discovered at an advanced stage with several metastases, symptoms of intramedullary spinal cord metastasis are rapid and aggressive, the diagnosis is based on magnetic resonance imaging. However, because of their rarity, no standard therapeutic regime has been developed and the prognosis remains poor.

Through an observation of intramedullary spinal cord metastasis in a patient with breast cancer, we present a recent systematic literature review regarding the clinical presentation, diagnostic modalities and the management of this rare metastasis.

CASE REPORT

We present a case of a 50 year-old female, who was under palliative chemotherapy for a history of a right breast carcinoma, with lung and bone metastasis approximately two years previously with stable disease.

After she completed three cycles of chemotherapy with Docetaxel (100 mg/m² every three weeks), she was

admitted with numbness and aching pain of both lower extremities associated with weakness and a intense headache that had been progressive over the previous two months and development of paraparesis one week before presentation, Physical examination revealed paraparesis (left lower extremity: 3/5, right lower extremity: 3/5) in a conscious patient with OMS 3 with no other physical symptoms.

On her cerebral and spinal Magnetic resonance imaging, we detected an intramedullary and highly gadolinium-enhancing lesion with regular borders (Figure1) associated with two cerebral metastatic lesions.

Systemic corticosteroid therapy in parallel to a palliative antalgic Radiotherapy (5 fractions of a total of 20 Gray) to both of the brain and intramedullary lesion was performed, with no relief from pain.

The patient died three months after the progression of her disease.



Figure 1: Spine magnetic resonance imaging showing solitary intramedullary nodular lesion.

DISCUSSION

Intramedullary spinal cord metastasis is a rare complication of cancer, affecting 0.1%–0.4% of all cancer patients. It accounts only 8.5% of all central nervous system metastasis and 0.9%–5% of all spinal cord metastasis.^[1,2] Most commonly, this rare complication is secondary to lung cancer 48%, but it can also be seen in breast cancer 16%, kidney cancer, and lymphoma.^[3]

First described by Buchholz and al. in 1897; Intramedullary spinal cord metastasis is most often discovered at an advanced stage with several metastases like in our patient with pulmonary, bone and cerebral localizations.^[5]

Many pathogenetic mechanisms are suggested, haematogenous spread is believed to account for most cases; it is developed through systemic route (arterial or venous) or through the vertebral venous plexus; Although the spread of tumors to the spinal cord can be related to lepto-meningeal dissemination by the cerebrospinal fluid or by direct invasion from contiguous structures.^[5]

In our case as in the literature, it is commonly seen on thoracic area (42%) and then on cervical area (31%).^[6] In 2013, Rostami et al. reported in a review article that so far there are 85 cases that have breast malignancy as the primary cancer. The localization is detectable in 52 cases (62%). 45 patients have solitary lesions and 17 of them have cervical (38%), 17 have thoracic (38%), and 11 have lumbar (24%) lesions.^[7,8]

Clinically, Rostami et al reported that symptoms are rapid onset in less than a month, the most common are sensory loss (22.1%), pain (21%), weakness (21%) as well as sphincter disorders (12%).^[7]

Cerebrospinal fluid examinations are frequently abnormal, with an elevated protein value, but they have much less value in the diagnosis of intramedullary spinal cord metastasis and it infrequently contains malignant cells.^[7,10]

Nowadays, Magnetic resonance imaging is the gold standard method and the most sensitive and specific test for detection of intramedullary spinal cord metastasis,^[11] it gives information on the nature of the lesion and its relation with adjacent tissues. The use of gadolinium-enhanced Magnetic resonance images has further improved sensitivity; Magnetic resonance imaging also helps for the planning of the surgery.

Treatment options for intramedullary spinal cord metastasis include radiation with or without steroid therapy, surgery, chemotherapy, immunotherapy, and targeted therapy. Because of the blood–brain barrier, chemotherapy is regarded as inappropriate for intramedullary spinal cord metastasis.^[2,12] Immunotherapy lacks efficacy and produces serious adverse reactions; further clinical trials are needed.^[12]

Moreover, the choice of the treatment had to take into account the general state of the patient, stage of the disease as well as the location and extent of the metastases.

Surgical resection is indicated in case of localized intramedullary spinal cord metastasis with a controlled primitive, with the aim at decompression of functional neural tissue, improvement of the neurological symptomatology and histological confirmation of tumor.^[13]

Regarding radiation therapy, there are some reports of long-lasting remission after irradiation, but these are limited to patients in whom a very early diagnosis was made and/or who had highly radiosensitive tumors such as small-cell carcinoma or lymphoma.^[14,15,16]

The place of chemotherapy remains unknown but depends on the primary tumor,^[3] Weissman and Grossman,^[17] have recommended treatment combining intrathecal chemotherapy and radiotherapy in patients

with intramedullary spinal cord metastasis from small cell carcinoma.

Prognosis of intramedullary spinal cord metastasis remains poor. The majority of patients have a poor life expectancy due to synchronous brain metastases and a poor performance status secondary to neurological deficits.

Like in our case, Germ *et al.*^[14] found that more than 80% of patients died within 3 months.

CONCLUSION

Because of their rarity, intramedullary spinal cord metastasis is generally associated with unduly delayed diagnosis and poor survival. Treatment options include radiation therapy, surgery, chemotherapy, immunotherapy, and targeted therapy; however, there is no consensus on optimal treatment. More reports are needed to enable exploration of the mechanisms of this metastasis and the optimal forms of therapy. Waiting that, the primary objectives still is maintaining quality of life of these patients by using adequate supportive care.

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