A Rare Case of a Thoracic Spinal Hemangiopericytoma

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Abstract

Spinal hemangiopericytomas are rare tumors. To date, only 80 cases of spinal hemangiopericytomas have been reported in the literature. The postoperative evolution of this condition in rare cases can be complicated by a symptomatic compressive epidural hematoma that can be the source of serious neurological complications requiring emergency surgery. We report a rare case of intradural and extramedullary spinal hemangiopericytomas with favorable evolution after treatment in an 82 years old woman.

Keywords

Hemangiopericytoma, Spine, Intradural, Extramedullary, Radiotherapy, Emergency, Surgery

1. Introduction

The hemangiopericytoma is a rare and malignant tumor of the central nervous system classically developed from the meninges and able to metastasize outside the neuraxis [1]. It represents less than 1% of all the central nervous system tumors and 2% of the primary meningeal tumors [2]. Hemangiopericytomas have been first considered to be angioblastic meningiomas before OMS classified them as non-meningothelial tumors of mesenchymal origin in 1993 [1]. Since the 2016 central nervous tumors classification of the WHO, they are grouped with solitary fibrous tumors (TFS) under the same entity: TFS/hemangiopericytoma [3]. Most primary hemangiopericytomas of the central nervous system are intracranial and rarely located in the spine. The secondary spinal location is most often related to an intracranial origin [4]. The diagnosis is histologic and mole-
cular [2]. The complete surgical resection remains the standard treatment and the interest of radiotherapy remains controversial [5]. The outcome in case of an incomplete surgical resection is generally marked by a high rate of the tumor relapse in the local site [6]. To date, only about 80 cases of spinal hemangiopericytomatas have been reported in the literature [1]. We report a rare case of an intradural and extra-medullary thoracic hemangiopericytoma.

2. Case Report

An 82-year-old patient was hospitalized for a motor deficit of the lower right limb that was disturbing her gait and evolving for three weeks. The clinical examination of this patient showed a deficit in the right thigh and in the levator muscle of the right foot respectively at 3/5 and 2/5. There were no ascending fibers disorders. MRI of the thoracic spine revealed an intradural and extramedullary T9 - T10 tumor with a homogenous enhancement after gadolinium injection (Figure 1). The diagnosis of meningioma and neuroma was evoked on the basis of the characteristics of the tumor on MRI. A surgical resection was performed. This surgical procedure consisted of a laminectomy of T9 and an almost complete tumor resection with a capsular residue which represented approximately 10% of the tumor’s size. This capsular residue was very adherent to the spinal cord. The anatomopathological diagnosis of the lesion was Grade III hemangiopericytoma according to the WHO classification (Figure 2). The tumor exhibited a typical histological appearance of hemangiopericytoma with nuclear positivity of the STAT6 marker in immunohistochemistry. Grade III was proposed because of a high mitotic index. The immediate postoperative follow-up was marked by a compressive epidural hematoma of the operative site which worsened the motor deficit of the patient who passed to 0/5. This hematoma was evacuated in emergency with none complication during the immediate post operative follow-up. The patient’s discharge was allowed a few days later with a beginning of recovery of the motor strength regarding the right lower

Figure 1. MRI sequence T1 sagittal sections ((a) and (b)) and T2 axial section (c). (a) and (b): without and with injection showing an intradural and extramedullary tumor with homogeneous enhancement after injection of gadolinium, (c): Intradural and extramedullary lesion. Blues arrows show the tumor.
Figure 2. (a) Histological section showing a proliferation of monomorphic cells irrigated by many capillaries and encompassed in a thin fibrous network; (b): histological section after anti-STAT6 immunohistochemical staining, ×400: tumor nuclei positivity for STAT6, confirming the diagnosis of hemangiopericytoma

limb (3/5 proximal and 1/5 distal). A functional reeducation prescription has been made and is still going on. A brain MRI performed in order to the research for a primary tumor site was negative. A complementary radiotherapy, more precisely a volumetric arc therapy with intensity modulation has been carried out. It has consisted of a 52 grays administered in 26 fractions. At the postoperative consultation five months after the surgery the patient had an improvement in motor strength. This one was rate at 3/5. The research for distant metastatic foci after radiotherapy was negative.

3. Discussion

Hemangiopericytomas are rare vascular and aggressive tumors developed at the expense of Zimmermann’s pericytes [1] [7]. The 2016 WHO classification considers hemangiopericytomas and solitary fibrotic tumors of the nervous system as the same pathological entity due to immunohistochemical and molecular lesions similarities [3]. The primary spinal location of hemangiopericytomas has been rarely reported in the literature [2]. To date, about 80 cases of spinal hemangiopericytomas have been reported and the cervico-dorsal spine is the most frequent site [1]. The clinical manifestation can be summarized as a motor deficit, radiculalgia and sensory disorders [1]. The sphincter disorders are relatively rare and late [8]. The case we report was clinically manifested only by a deficit in the right lower limb. The radiological figures of spinal hemangiopericytomas are nonspecific and have similarities with other spinal tumors, including meningiomas and neuromas [5]. The tumor, most often, has an intradural, extramedullary localization, well circumscribed with a dural attachment [1]. Purely intramedullary or purely intradural locations are rare [8]. The tumor is most often heterogeneously enhanced on MRI in T1 sequence with injection of gadolinium [9]. The presence of voids signals zones in T2 sequence is indicative of the hypervascularized nature of the tumor [5]. The case we report had rather a homogeneous enhancement with T1 contrast MRI and no T2 void signal. The differential radiological diagnosis of spinal hemangiopericytomas is mainly dis-
Spinal Hemangiopericytomas are rare tumors which are usually difficult to be totally removed. The relapse of the tumors locally or metastases appearance is very common regarding the outcome of this disease. A close post-operative monitoring could be useful to detect such complications.

4. Conclusion

Spinal hemangiopericytomas are rare tumors. The diagnosis is relatively difficult because of the rarity of the tumor and the nonspecific nature of the clinico-radiological manifestations. Complete surgical resection is the optimal treatment. Adjuvant radiotherapy should be given in the case of a high grade...
tumor and incomplete tumor resection. Close monitoring would detect local recurrence and possible metastases.

Conflicts of Interest

The authors do not declare any conflict of interest.

References


