Unusual presentation of giant schwannoma in the sciatic nerve

Julia Brasileiro de Faria Cavalcante, Pedro Nogarotto Cembraneli, Renata Brasileiro de Faria Cavalcante, Volmer Fernandes Valente Junior, José Edison da Silva Cavalcante

ABSTRACT

Introduction: Schwannomas are benign, encapsulated, slow-growing, generally solitary tumors originating from Schwann cells in the peripheral nerve heath. Schwannoma of the sciatic nerve is a rare condition and, therefore, little documented in the literature. Case Report: A 77-year-old female patient arrived at the clinic complaining of pain and the appearance of a mass in the middle third of her left thigh. She presented with a well-defined mass, horizontally movable, hyperalgesia to the touch at the site of the mass, and positive Tinel’s sign. Magnetic resonance imaging showed multiple lesions suggestive of neurinomas, with intense nodular enhancement after gadolinium contrast injection, diffuse, in the right thigh and in the middle third of the left thigh. Surgical treatment was proposed by our team, but the patient preferred to continue the outpatient follow-up, because the injury did not interfere in her daily activities. Conclusion: Although schwannoma in the sciatic nerve is rare, it should be considered in the differential diagnosis of patients presenting with conditions that affect this nerve.

Keywords: Schwannoma, Sciatic nerve, Tumor

INTRODUCTION

Peripheral nerve tumors are a heterogeneous group of benign tumors, which are rare in the general population [1]. Schwannomas are primary tumors of the neural sheath originating from Schwann cells [2, 3]. The occurrence of these tumors in the sciatic nerve is very rare (1%) [4]. These tumors grow slowly and since most of them are asymptomatic, they can be detected incidentally. It is also possible that they become symptomatic due to mechanical compression. Magnetic resonance imaging (MRI) is the diagnostic tool of choice for diagnosis of most soft-tissue tumors, including schwannomas, because of its high diagnostic predictability [5].

CASE REPORT

A 77-year-old female patient sought care due to persistent low-intensity pain and mass in the middle
region of the left thigh for two years. She reported that the mass had been progressively growing in the last few months associated with paresthesia of the left lower limb.

Physical examination showed preserved strength and hyperalgesia at the site of the mass. The patient described the pain as intermittent, radiating to the distal part of the left lower limb, with burning and electric shock-like sensations. Deep palpation of the left thigh region revealed a well-defined mass, approximately 8 cm in diameter (Figure 1), with regular contours, not hardened, horizontally movable, painful to the touch.

Magnetic resonance imaging showed multiple lesions suggestive of neurinomas, with intense nodular enhancement after gadolinium contrast injection, from 1.0 to 5.0 cm in diameter, diffuse, in the right thigh and in the middle of the left thigh, with heterogeneous hypersignal in T2 and foci of intrallesional cystic degeneration (Figure 2). A tumor mass was visualized in the middle third of the sciatic nerve in the left thigh, measuring $9.0 \times 7.0 \times 6.0$ cm, with cystic solid content (Figure 3).

The diagnosis of peripheral nerve schwannoma was established based on the characteristics of the MRI and the clinical history of the patient, who did not present other semiological findings. The surgical treatment was suggested by our team, but the patient refused it and chose to be followed up in the outpatient clinic, because she affirmed that the injury did not interfere in her daily activities.

**DISCUSSION**

Schwannomas of peripheral nerves in the lower limbs most frequently affect the posterior tibial nerve. The occurrence of this type of tumor in the sciatic nerve is very rare, with a frequency below 1%, and the presence of a painful palpable mass is the most reported clinical presentation [4, 6, 7].

Schwannomas grow very slowly over years. The lesions stretch slowly and lengthen the fascicles, still preserving the neurological function [8, 9]. Microscopically,
the tumor is composed of spindled cells arranged in interlocking bundles. The nuclei tend to be arranged parallel to each other, showing an aspect that is called palisade arrangement [10].

Although most of these tumors are asymptomatic, they may be associated with pain and paresthesia in the corresponding neural territory. Hypoesthesia and motor deficits are more common in patients presenting with tumors larger than 4 cm in diameter [11, 12]. On physical examination, the tumor can be moved laterally, but not longitudinally toward the tumor trajectory, and percussion can produce positive Tinel’s sign [8, 9].

The exam of choice for the diagnosis and management of the lesion is MRI [5, 13], whose findings are: homogeneous mass, with medium to low signal intensity in T1-weighted sequences, while presenting hyperintense signal in T2 after gadolinium contrast infusion [9].

The treatment of schwannomas is surgical. The procedure aims at the complete resection of the tumor employing microsurgical techniques [8, 9, 13].

CONCLUSION

Although schwannoma in the sciatic nerve is a rare tumor, it should be considered in the differential diagnosis of patients presenting with conditions that affect this nerve.

REFERENCES


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Author Contributions

Julia Brasileiro de Faria Cavalcante – Conception of the work, Design of the work, Acquisition of data, Analysis of data, Interpretation of data, Drafting the work, Revising the work critically for important intellectual content, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved

Pedro Nogarotto Cembraneli – Conception of the work, Design of the work, Acquisition of data, Analysis of data, Interpretation of data, Drafting the work, Revising the work critically for important intellectual content, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved

Renata Brasileiro de Faria Cavalcante – Conception of the work, Design of the work, Acquisition of data, Analysis of data, Interpretation of data, Drafting the work, Revising the work critically for important intellectual content, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved

Volmer Fernandes Valente Junior – Conception of the work, Design of the work, Acquisition of data, Analysis of data, Interpretation of data, Drafting the work, Revising the work critically for important intellectual content, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved

José Edison da Silva Cavalcante – Conception of the work, Design of the work, Acquisition of data, Analysis of data, Interpretation of data, Drafting the work, Revising the work critically for important intellectual content, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that
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Authors declare no conflict of interest.

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