



Deep Vein Thrombosis in the Setting of Neurofibromatosis Type 1: Case Report

Trombose venosa profunda em neurofibromatose tipo 1: Relato de caso

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Abstract

Neurofibromatosis type 1 (NF1) is a genetic syndrome which typically presents with neurological manifestations. Some of the patients may also present with vasculopathies, among which arterial aneurysms and stenosis are the most common. Deep vein thrombosis (DVT) has rarely been described, and, to the best of our knowledge, the present is the first report of DVT due to venous compression by a neurofibroma in the setting of NF1.

This is the case of a 23-year-old male with NF1 who experienced DVT due to compression of the left posterior tibial veins by a large tumor arising from the tibial nerve. The DVT was acutely treated with enoxaparin and then with rivaroxaban. Two months after the diagnosis, Doppler ultrasonography showed partial recanalization and persistence of the DVT. The patient was then referred to neurosurgery for surgical resection of the tumor. There were no complications during the procedure, and the patient did not present postoperative neurological deficits. The final histopathological diagnosis was of a benign neurofibroma. After one year of follow-up with vascular surgery, the patient presented no more episodes of DVT.

In case there is a tumor compressing the deep vessels of the leg and promoting DVT, surgical resection with microsurgical techniques may be curative.

Keywords

- ▶ neurofibromatosis 1
- ▶ neurofibroma
- ▶ venous thrombosis

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Resumo

A neurofibromatose tipo 1 (NF1) é uma síndrome genética classicamente associada a manifestações neurológicas. Contudo, alguns dos pacientes apresentam também manifestações vasculares, dentre as quais aneurismas e estenoses arteriais são as mais comuns. Trombose venosa profunda (TVP) associada a NF1 raramente foi descrita, e, pelo que sabemos, este é o primeiro relato de TVP devido a compressão venosa por neurofibroma no contexto de NF1.

Este é o caso de um paciente masculino de 23 anos, com NF1, que apresentou um episódio de TVP devido a compressão das veias tibiais posteriores esquerdas por um grande neurofibroma que surgia do nervo tibial. A TVP foi tratada de forma aguda, com enoxaparina e rivaroxabana. Dois meses após o diagnóstico, a ultrassonografia com Doppler demonstrou recanalização parcial com persistência da TVP. O paciente foi então referido à neurocirurgia para ressecção da massa. Não houve complicações devido ao procedimento, e o paciente não apresentou déficits neurológicos. O diagnóstico histopatológico final foi de um neurofibroma. Após um ano de seguimento com a cirurgia vascular, o paciente não apresentou mais episódios de TVP.

Caso haja um tumor comprimindo os vasos profundos da perna e promovendo TVP, a ressecção cirúrgica com técnicas microcirúrgicas pode ser curativa.

Palavras-chave

- ▶ neurofibromatose 1
- ▶ neurofibroma
- ▶ trombose venosa

Background

Neurofibromatosis type 1 (NF1) is a complex genetic syndrome, in which the gene for neurofibromin (a tumor suppressor protein) suffers a wide range of mutations,¹ resulting in decreased neurofibromin synthesis. The prevalence of NF1 is around 1/3,000,^{2,3} and it presents as a systemic disease, with dermatological, cardiovascular, gastrointestinal, orthopedic, central, and peripheral nervous system manifestations.^{4,5} One of the cardinal features of NF1 is the predisposition toward the development of certain peripheral nerve sheath tumors (PNSTs). These may be benign, such as neurofibroma, or malignant, such as the malignant peripheral nerve tissue tumor (MPNST).¹ Among the systemic complications of NF1, there is a series of vascular abnormalities, of which arterial manifestations are the most common. Those mostly comprise aneurysms and stenoses of renal, aortic, and mesenteric arteries,⁶ with hypertension as the most common clinical feature secondary to these.^{4,6,7} Venous manifestations are, in turn, exceedingly rare, and may include venous thrombosis (VT).⁸⁻¹³

Literature Search

To investigate previous cases of NF1 with VT, we searched the MEDLINE database for articles written in English with the following Mesh keywords and Boolean operators: (“*Neurofibromatosis 1*” AND “*Venous thrombosis*”) OR (“*Neurofibromatosis 1*” AND “*Vascular disease*” AND “*Vein*”). While applying article-type filters to select case reports, clinical studies, observational studies, reviews, and systematic reviews, the search yielded 29 individual results, all case reports. In total, there were six case reports of NF1-associated VT.^{9-12,14,15} By searching the references of each of these studies, we were able to find another case.⁸ In these reports, five patients had aneurysms correlated to the formation of thrombus^{9,11,12,14,15} and two did not,^{8,10} one of which was secondary to compression by an exostosis.¹⁰

The other 23 articles reported arterial manifestations, vascular malformations, and NF1-associated vascular retinopathy. Some of these cases also presented with venous manifestations,^{13,16-23} yet none with thrombosis.

Case Presentation**History and Examination**

One year before surgery, a 22-year-old male with NF1 presented with swelling on the left leg, accompanied by pain on walking. Doppler ultrasonography (USG) was performed and showed a large mass over the course of the left tibial nerve. There was no DVT, and the left saphenous veins were tortuous. We decided not to resect the tumor at that moment. The patient was treated with 0.5 g of metamizole twice a day for 1 week, with improvement of the pain.

Ten months later, the patient presented with an acute episode of edema and pain in the left leg. Doppler USG was performed, and showed DVT of the posterior tibial veins and thrombophlebitis of one collateral of the saphenous vein. The patient was hospitalized for 3 days, and underwent treatment with enoxaparina (1.5 mg/kg once a day). A computed tomography angiogram (angio-CT) was performed, and it showed compression of the left fibular and posterior tibial veins by a large tumor arising from the tibial nerve (▶ **Fig. 1**). Given the setting of NF1, the lesion was presumed to be a neurofibroma. The patient continued treatment with 15 mg of rivaroxaban twice a day for 21 days, and then 20 mg once a day. Doppler USG was again performed two months after the initial diagnosis of DVT, and it showed partial recanalization with persistence of a thrombus (▶ **Fig. 2**). The saphenous veins were congested and tortuous. The patient was then referred to the authors' care for evaluation by vascular and peripheral nerve surgery.

Upon physical examination, he presented with pain on the left leg (visual analog scale [VAS] = 6), M4+ left foot flexion on the British Medical Research Council (BMRC) scale, and



Fig. 1 (A) Three coronal slices of a computed tomography angiogram (angio-CT) showing the compression of the posterior tibial veins by the tumor (T) arising from the tibial nerve. Note the deviation of the course of one of the posterior tibial veins (ptv), compressed against the muscles of the posterior compartment of the leg. One of the fibular veins (fv) appears to be compressed against the fibula. (B) 3D reconstruction of an angio-CT from a posterior-medial point of view showing the mass effect of the tumor onto the posterior tibial veins. The white arrows show one of the deviated fibular veins. The black arrows show one of the deviated posterior tibial veins.



Fig. 2 Doppler ultrasonography of the left posterior tibial veins, performed two months after hospitalization for the treatment of deep vein thrombosis. There is partial recanalization, yet the thrombus is still present. Abbreviation: V TIB POST ESQ, left posterior tibial veins.

plantar hypesthesia. The mass was not palpable, yet the left leg was edematous. There was also a positive Tinel sign over the course of the left tibial nerve.

Electroneuromyography was performed, and it showed a delay in motor conduction with a reduction of amplitude (3.7 μ V), and a minor delay in the distal sensitive response of the left sural nerve. Magnetic resonance imaging (MRI) was then also performed to better evaluate the tumor's relationship to neighboring structures and to help in the preoperative planning (→ Fig. 3). Even though the lesion was large (96 mm on its largest axis), it presented predominantly high ($> 1.3 \times 10^{-3} \text{ mm/s}^2$) apparent diffusion coefficient (ADC) values on diffusion-weighted imaging (DWI). This, along with the absence of features suggestive of malignancy (that is, peripheral enhancement, perilesional edema, intratumoral cystic changes), favored the hypothesis of a benign neurofibroma.^{24,25} There were no signs of muscle denervation.

Surgery

Given the presence of a large tumor inside the left tibial nerve, and the evidence of the persistence of DVT on Doppler USG, surgical resection of the tumor with decompression of the vessels was then decided. With the aid of loupes, a classic approach to the left tibial nerve through the medial aspect of the leg was performed, with mobilization of the gastrocnemius and soleus muscles. Gentle progressive dissection was then performed, with individualization of the compressed deep

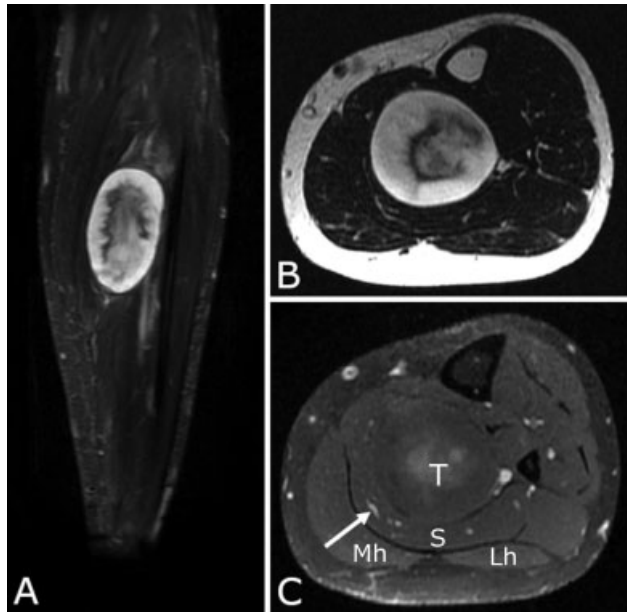


Fig. 3 (A) Coronal Short tau inversion recovery weighted (STIR-weighted) magnetic resonance imaging (MRI) without contrast of a large heterogeneous lesion (96 × 56 × 49 mm) arising from the left tibial nerve and compressing the deep vessels of the posterior compartment of the left leg. The lesion presents with well-defined borders, and there are no apparent signs of tissue invasion or perilesional edema. (B) Axial T2-weighted MRI without contrast of the lesion showing predominantly high signal intensity and a central region with lower signal intensity. (C) T1-weighted MRI with contrast showing discrete and heterogeneous central enhancement. The arrow shows one of the posterior tibial veins compressed against the soleus muscle fascia. Abbreviations: Lh, lateral head of the gastrocnemius; Mh, medial head of the gastrocnemius; MRI, magnetic resonance imaging; S, soleus muscle; T, tumor..

vessels of the leg. After the complete exposure of the tumor (and of the healthy tibial nerve proximally and distally to the mass), electrical mapping of its surface was conducted with the aid of intraoperative electrical stimulation to identify an area devoid of functional fascicles. With the aid of microsurgical techniques under microscopy, a sharp opening of the pseudotumoral capsule was performed in this “electrically-silent” area. An intraoperative biopsy was then performed, and it revealed a probable neurofibroma, without characteristics of malignancy. The pseudotumoral capsule could be differentiated from the true tumor capsule by its color and consistency (respectively white-grayish and tough, against yellowish and soft), and a cleavage plane was established in a fascicle-free corridor. A circumferential dissection was performed with complete isolation of the tumor toward its proximal and distal poles to identify the tumor’s fascicle of origin (in this case, only one). The fascicles that entered and exited the tumor were not functional, and they were sectioned. The huge tumoral mass was resected en bloc, with the preservation of the functional fascicles. The borders of the nerve were gently everted to look for any residual tumor inside the nerve (“open book” maneuver) (► Fig. 4).

Follow-up

No postoperative complications were observed. The patient left the hospital with the same deficit as before (M4+ foot flexion) and reduction of pain (VAS = 3). He was treated with pregabalin as soon as was discharged (75 mg 3 times a day during the first month; twice, during the second month; and once, during the third month), and was directed to physiotherapy treatment after the stitches were removed. He was followed up in the outpatient clinic, and, after 3 months, recovered to M5 foot flexion and presented with no more

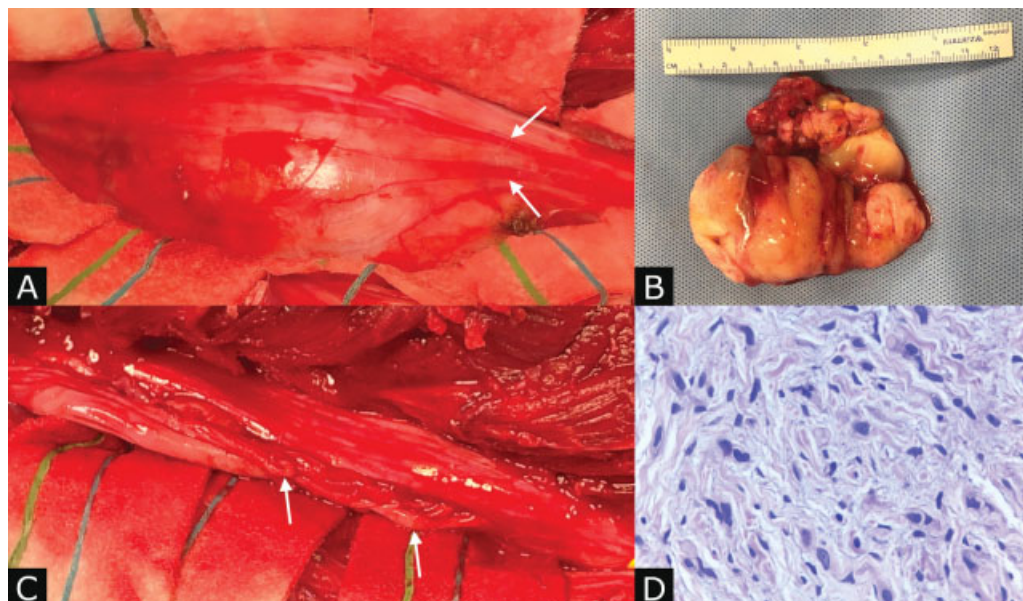


Fig. 4 (A) Exposure of the tumor. The arrows show functional fascicles dislocated by the mass. (B) Surgical aspect after resection of the lesion. The arrows show the preserved functional fascicles inside the tibial nerve. (C) Tumor resected en bloc. (D) Hematoxylin and eosin, 400x. Benign neurofibroma showing hypocellular proliferation of slightly elongated spindle cells, with wavy and hyperchromatic nuclei; in some, the nucleolus is evident. Collagen and a slight amount of mucoid material are observed among the neoplastic cells, as well as rare mononuclear inflammatory cells. The specimen presented no necrosis, and there were rare nuclear atypia.

pain, so that pregabalin was discontinued. Six months after the surgery, the patient was also evaluated by vascular surgery, and no further episodes of DVT were observed.

Discussion

In the present study, we report a case of a patient with NF1 who experienced DVT due to compression of posterior tibial veins by a neurofibroma in the posterior compartment of the leg. To the best of our knowledge, the present is the first report of VT secondary to external compression by neurofibroma in the setting of NF1.

The prevalence of NF1 vasculopathy has been estimated to be of at least 8% in a study²⁶ conducted with 181 pediatric patients with NF1. Its mechanisms are yet not completely understood.^{4,27-29} The vascular manifestations of NF1 appear to increase morbidity and mortality. A nationwide study³⁰ conducted in the United States with death certificates showed that NF1 patients younger than 30 years of age who died were more than twice as likely to have been diagnosed with vascular disease when compared with those without NF.

The vasculopathy pathogenesis of NF1 has been shown to involve a series of events caused by dysfunction of the synthesis of neurofibromin (the encoded protein of the NF1 gene), which physiologically works as a downregulator of the Ras cascade signaling. Without the downregulation, the Ras signaling pathway augments cell proliferation in the vascular endothelium (with enhanced expression of cyclin D and cells more frequently entering the cell cycle).²⁸ It also causes smooth-muscle hyperplasia⁷ and inflammation.²⁹ The cell lineage that appears to be key in this pathogenesis is the bone-marrow-derived cell (BMDC). It has been shown that the inactivation of the NF1 gene in this cell lineage in mice was both sufficient and necessary to cause neointima formation and evidence of vascular inflammation similar to that observed in NF1 knockout mice.²⁹

Although the manifestations of NF1 vasculopathy are mostly arterial, patients may also present with venous rupture, venous aneurysms, and/or VT.^{8-12,31} It has been shown that the absence of NF1 *in vitro* is sufficient for human venous endothelium cells to undergo autonomous proliferation.²⁸ This endothelial dysfunction has been extensively implicated in the pathogenesis of VT.³²⁻³⁵ In this setting, the venous endothelium also shows altered vascular morphogenesis, which helps to explain the vascular morphological alterations in NF1 patients (that is, stenosis and aneurysms),^{6,9,12,14,19,31} some of which may further increase the risk of VT (that is, venous aneurysm).³⁶⁻³⁸

We have found four reports of NF1 patients presenting with venous aneurysm and an associated VT. Seinturier et al.¹² presented a case in which a 64-year-old woman, with NF1, developed pulmonary embolism secondary to a thrombosed venous femoral aneurysm. There are other 3 case reports of NF1 patients younger than 50 years of age who presented with internal jugular thrombosis secondary to aneurysmal degeneration.^{9,11,14}

Lehrnbecher et al.⁸ reported a case in which a 4-year-old boy was diagnosed with NF1 systemic vasculopathy, including

thrombosis of the right posterior tibial vein, without evidence of a correlated aneurysm or compression/infiltration by a tumor.

None of the seven reports described compression or infiltration of venous structures by a neurofibroma. Nonetheless, DVT in NF1 patients may also be caused via extrinsic compression by these tumors, as blood stasis has also been implicated in thrombogenesis.^{33,39} This, we think, contributed to the process of thrombogenesis in the present case, given that the tumor arose from the tibial nerve, in a tight region in the posterior compartment of the leg.

Deep vein thrombosis secondary to external compression by tumors has been reported in other settings such as the superior vena cava, and pulmonary and iliofemoral veins.⁴⁰⁻⁴² It has been reported once in the setting of NF1, by an exostosis, in the popliteal vein.¹⁰ One study⁴³ conducted with patients harboring high-grade non-Hodgkin lymphoma showed that venous compression by the tumor was present in 51% of patients with DVT. Given the higher prevalence of neurofibromas in NF1 patients, it is thus reasonable to assume that DVT secondary to tumoral compression may also have a higher prevalence in this population.

Conclusion

Clinicians should be alert to signs and symptoms suggestive of DVT in NF1 patients, as NF1 seems to favor venous endothelium dysfunction. Tumors may also compress or infiltrate the veins, promoting VT through blood stasis, when there is venous compression; or through endothelial damage, when infiltration occurs. In case there is a tumor compressing venous structures and promoting DVT, surgical resection with microsurgical techniques may be curative and able to preserve neurological function.

Conflict of Interests

The authors have no conflict of interests to declare.

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