Case report
Peripheral neuropathy in systemic vasculitis and other autoimmune diseases – a report of five cases emphasizing the importance of etiologic characterization

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ABSTRACT
Introduction: Peripheral neuropathies may present in the context of systemic vasculitis and other autoimmune diseases. The etiologic characterization is crucial to define the treatment and prognosis in secondary vasculitis. The purpose of this study is to describe the pathway of etiologic investigation including the role of nerve biopsy.

Methods: Retrospective analysis of patients seen in the neuromuscular outpatient clinic during the last four years with peripheral neuropathy in the context of systemic vasculitis or other autoimmune diseases.

Results: We present five patients with stepwise progressive sensorimotor deficits of upper and lower limbs. All patients presented with systemic features and one of them had an established diagnosis of systemic vasculitis. They underwent an extended blood panel, including autoimmune and serologic tests. Electromyography and nerve conduction studies revealed asymmetric axonal sensorimotor polyneuropathies in four patients, and an axonal sensorimotor multiple mononeuropathy in one. Four patients underwent nerve biopsy and the other performed a skin biopsy, with findings suggestive of possible vasculitic processes. The etiologies identified included microscopic polyangiitis, HBV-related polyarteritis nodosa and two eosinophilic granulomatosis with polyangiitis. In the last patient a specific etiology could not be established.

Conclusion: This series reveals the etiologic and phenotypic diversity of peripheral neuropathies related with systemic vasculitis. The therapeutic approach and prognosis were distinct in each patient, emphasizing the importance of a prompt diagnosis and appropriate treatment.

1. Introduction
Peripheral nervous system impairment in systemic vasculitis is often related with poor functional prognosis and increased morbidity, with sensorimotor deficits that are sometimes irreversible [1,2].

Either primary or secondary to autoimmune diseases, vasculitis affecting small arteries or large arterioles are more likely to produce a peripheral neuropathy [3,4]. The primary vasculitis most commonly associated with neuropathy includes polyarteritis nodosa (PAN), essential mixed cryoglobulinemic and anti-neutrophil cytoplasmic antibodies (ANCA) associated vasculitides [5–8]. Among the autoimmune diseases, rheumatoid arthritis, hepatitis B virus (HBV)-associated PAN, and hepatitis C virus (HCV)-related cryoglobulinemic vasculitis are the most frequent [5,7,8].

The treatment and prognosis of these neuropathies is variable and depend on a correct and individualized etiologic characterization.

2. Methods
We conducted a retrospective study of patients with peripheral neuropathy in the context of systemic vasculitis, seen in the last four years in our neuromuscular outpatient clinic. All patients gave informed consent for publication.
3. Results

3.1. Patient 1

A 57-year-old male presented with a four-month history of progressive lower limbs livedo and myalgias, plantar dysesthesias and bilateral foot drop. He also had a productive cough, bilateral ankle edema and a constitutional syndrome with involuntary weight loss of 9 kg in two months. A week before the admission he noticed tactile and thermalgia hypoesthesia in the left ulnar nerve territory.

Blood panel showed mild normocytic-normochromic anemia, erythrocyte sedimentation rate (ESR) of 92 mm/h, creatinine of 1.4 mg/dl. He was positive for ANA, ANCA anti-myeloperoxidase, anti-SSA and anti-SSB. HIV, HBV and HCV serologies were negative. Upper and lower gastrointestinal endoscopy, abdominal and renal ultrasound were unremarkable. Chest CT-scan revealed only discrete pleuroparenchymal fibrocicatricial densities in both apices. Cerebrospinal fluid (CSF) analysis was normal. Nerve conduction studies (NCS) revealed an absence or reduction in amplitude of sensory nerve action potentials (SNAP) in the lower limbs. Electromyography (EMG) of lower limb muscles was normal, in keeping with a mild sensory axonal polyneuropathy. A skin biopsy identified an intense inflammatory infiltrate mainly lymphocytic into a skin arteriole wall (Fig. 1A), with internal elastic lamina disruption and fibrinoid necrosis (Fig. 1B). These changes were compatible with a leukocytoclastic vasculitis, supporting the diagnosis of a clinically probable vasculitic neuropathy associated with microscopic polyanangiitis, by Peripheral Nerve Society (PNS) criteria [9].

Patient was started on intravenous methylprednisolone 1 g/day during 3 days and then oral prednisolone 1 mg/kg/day, with a clear improvement of pain and constitutional syndrome. Three months later, he was almost completely recovered of motor deficit and corticotherapy was tapered off. Although neurologically stabilized, cyclophosphamide 100 mg/day and then azathioprine 75 mg/day were needed due to renal impairment.

After two years, he reported a relapse of motor deficits and neuropathic pain. He was restarted on oral prednisolone 1 mg/kg/day and azathioprine dose was increased to 100 mg/day, with complete recovery of symptoms within two weeks. The dosages were gradually tapered off, and he is currently under azathioprine 75 mg/day and prednisolone 5 mg/day, without new neurological symptoms.

3.2. Patient 2

A 53-year-old male presented with progressive bilateral lower limb edema, plantar dysesthesias and hypesthesia in bilateral sural nerves territory, beginning four months earlier, evolving to left foot drop a month later, and right median nerve territory hypesthesia (thermalgic, vibratory and light touch) and lower limb rash two months after the onset. Blood panel showed an active HBV infection. NCS revealed an absence in SNAP in the lower limbs and a reduction in amplitude of compound motor action potentials (CMAP) in the tibial and peroneal nerves. EMG showed signs of chronic reinnervation and active denervation in distal muscles of the lower limbs, with a slight asymmetric distribution. These findings are consistent with chronic, moderately severe, asymmetric sensorimotor axonal polyneuropathy. A sural nerve biopsy showed perivascular inflammatory infiltrate in a vessel of the perineurium (Fig. 2A) with asymmetric nerve fibre loss (Fig. 2B), suggesting a vasculitic process and allowing the diagnosis of a clinically probable vasculitic neuropathy due to HBV-related polyarteritis nodosa by PNS criteria [9].

He was treated with oral prednisolone 60 mg/day with partial response and was started on tenofovir 2 months after with complete remission of the clinical picture. Three years later, he remains under tenofovir with undetectable viral load and neurologically asymptomatic.

3.3. Patient 3

A 54-year old female, with eosinophilic granulomatosis with polyangiitis (EGPA) with pulmonary involvement and peripheral eosinophilia, presented with progressive lower limbs paresthesias, left foot drop and bilateral painful plantar dysesthesias, beginning four weeks before. One week before, she noticed pretibial skin lesions, bilateral ankle edema and bilateral lower limb tactile hypesthesia. The day before, she reported involvement of the right foot with bilateral foot drop, and hypesthesia in the right cubital nerve territory associated with weakness in the fifth finger flexion. She was under a prednisolone tapering regimen, with a decrease to 20 mg/day 2 months before and to 10 mg/day fifteen days before the presentation.

Blood panel revealed eosinophilia of 9200, slightly elevated IgE levels and ANCA anti-myeloperoxidase positive. Chest CT-scan showed patchy upper lobe ground-glass opacities, septal thickening and centrilobular micronodules suggestive of chronic eosinophilic pneumonia. NCS revealed an absence or reduction in SNAP in the lower and upper limbs and an absence or reduction of CMAP in the lower limbs. EMG showed signs of chronic reinnervation and active denervation in distal muscles of the lower limbs, with a symmetric distribution, which is in keeping with a severe sensorimotor axonal polyneuropathy with active denervation. Sural nerve biopsy revealed asymmetric nerve fibre loss (Fig. 3A) and cicatricial changes in the vessel wall with internal elastic lamina disruption (Fig. 3B), supporting a clinically probable vasculitic neuropathy by PNS criteria [9].

The diagnosis of peripheral neuropathy associated with EGPA was...
established, and intravenous cyclophosphamide was started with re-
solution of neurological symptoms after 3 months. She is free of neu-
rological symptoms since then. However, because of the severe pul-
monary impairment, she is currently waiting for a lung transplant.

3.4. Patient 4

A 67-year-old female, with past medical history of asthma and
breast cancer HER2+ treated with extended lumpectomy, che-
motherapy and radiotherapy, and under trastuzumab each three weeks,
presented with progressive distal sensorimotor deficits. She reported
bilateral plantar paresthesias and dysesthesias beginning nine days
before, evolving to right hand along with a weakness of the right thumb
opposition, and then lower limb petechiae, progressive distal tactile and
 thermoalgic hypoesthesia and bilateral foot drop, losing the ability to
walk three days before the admission. She had started montelukast due
to uncontrolled asthma a week before the symptoms onset.

Blood panel revealed mild normocytic-normochromic anemia, eo-
sinophilia of 13,800, IgE levels higher than 5000 U/l and ESR of
102 mm/h. Autoimmunity tests were negative. Chest CT-scan revealed
mild bilateral fibrocicatricial changes. CSF analysis was unremarkable.
NCS revealed an absence or reduction in SNAP in the lower and upper
limbs and an absence or reduction of CMAP in the lower limbs. EMG
showed signs of chronic reinnervation and active denervation in distal
muscles of the lower limbs, with a symmetric distribution, which is
concordant with a severe sensorimotor axonal polyneuropathy with
active denervation. A skin biopsy showed neutrophilic infiltrate into a
skin vessel wall (Fig. 4A) suggesting a leukocytoclastic vasculitis. A
sural nerve biopsy revealed cicatricial aspects in a perineurive vessel,
including disruption of internal elastic lamina, circumferential intimal
hyperplasia (Fig. 4B and C) and intraluminal angiogenesis suggestive of
vessel recanalization (Fig. 4B), compatible with a previous vasculitic
process. These findings supported the diagnosis of a clinically probable
vasculitic neuropathy associated with EGPA by PNS criteria [9].

Montelukast was discontinued and intravenous methylprednisolone
1 g/day during 3 days was started, followed by oral prednisolone 1 mg/
kg/day. Patient was discharged after a month slightly better, but still
unable to walk.

3.5. Patient 5

A 34-year-old female, with past medical history of rosacea, pre-
sented with progressive sensorimotor deficits beginning three weeks
before, with tactile hypoesthesia and dysesthesia in the left sural nerve
territory. Two weeks after, she developed hypoesthesia and dysesthesia in
the right median nerve territory and weakness of the right hand. Three
days prior to admission she refers the onset of moderate right hand arthralgia, mostly in the morning, without inflammatory signs.

Blood panel was unremarkable, except for a mild hypocomple-
mentemia and low levels of IgA. CSF analysis showed mild pleocytosis
(38 leukocytes with 51.7% of mononuclear cells) with normal glucose and proteins. The neurophysiological findings were compatible with a multiplex mononeuropathy, revealing an axonal compromise of the right median nerve and the left sural nerve, with a normal EMG. Sural nerve biopsy revealed severe axonal neuropathy, without evidence of regeneration, but with an asymmetric involvement of nerve fibres (Fig. 5), suggesting a clinically probable vasculitic neuropathy by PNS criteria [9].

Considering the presence of a peripheral neuropathy associated with a suspected autoimmune disease of unknown etiology, patient was started on oral prednisolone 20 mg/day. However, she did not report any improvement and stopped it because of adverse effects. The symptoms resolved spontaneously one year later and she remains asymptomatic for two years even with no medication.

4. Conclusion

Vasculitic neuropathy has a characteristic clinical presentation with progressive sensorimotor symptoms developing over weeks to months [5,7]. Most patients experience one or more acute attacks, with a stepwise installation of the deficits, but one-third have a chronic slowly progressive course [5,7,8]. Peripheral nervous system vasculitis exhibits three patterns of clinical involvement [3,7]. The commonest and most distinctive one is a multiple mononeuropathy, occurring in 35–65% of the patients with vasculitic neuropathy [3,5]. However, these patients may also present with a distal asymmetric polyneuropathy or, more rarely, a distal symmetric polyneuropathy [3,8]. In this series, all the patients presented with an asymmetric sensorimotor deficit, with a stepwise progression, featuring a pattern of overlapping multiple mononeuropathies, evolving to distal asymmetric polyneuropathies in the first four patients. The constitutional symptoms, as asthenia, anorexia and involuntary weight loss, the skin lesions and/or the specific organ involvement suggested an underlying systemic vasculitis and, although nonspecific, provided clues to the etiological diagnosis.

Despite the clinical similarities, the extensive diagnostic evaluation revealed distinct underlying aetiologies which determined different therapeutic approaches and prognosis. In the first patient, the peripheral nerve involvement was evident at the time of initial presentation. In addition to systemic symptoms, the stepwise progression of the neurological deficits raised the hypothesis of a vasculitic etiology. The diagnosis of a microscopic polyangiitis was supported by laboratory and pathological tests, and the treatment with corticosteroids was started.

Fig. 4. Eosinophilic granulomatosis with polyangiitis (A) Skin histology, H&E, Inflammatory infiltrate (arrow) mainly composed of small lymphocytes (CD3+, CD10+, CD20-) into a skin arteriole wall (B and C) Perineve vessels from sural nerve histology, toluidine blue, Perineve vessels with cicatrical aspects including disruption of internal elastic lamina, circumferential intimal hyperplasia (B and C) and intraluminal angiogenesis (B). Scale bar: 50 μm (A, B and C). (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

Fig. 5. Undetermined systemic autoimmune disease. Histology of sural nerve, toluidine blue. Severe and active axonal neuropathy with an asymmetric involvement of nerve fibres. Scale bar: 200 μm (A) and 100 μm (B and C). (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)
with a complete resolution of the peripheral neuropathy. The second patient also presented with a stepwise progressive peripheral neuropathy but with more subtle systemic symptoms. The diagnosis of a HBV-related polyarthritis nodosa was decisive to start a directed antiviral treatment besides corticotherapy, emphasizing the importance of a specific etiological diagnosis for initiating the proper treatment. In the third patient, the diagnosis of EGPA was already known and the peripheral neuropathy was not a presenting symptom. However, in this case, the confirmation of a probable peripheral nerve vasculitis supported the escalation of therapy allowing a better neurological prognosis. In the fourth patient, it was the investigation of the neurological symptoms that led to the diagnosis of EGPA. Some authors have reported EGPA as a potential side effect in individuals with moderate to severe asthma on leukotriene receptor antagonists, such as montelukast [10,11]. Because of this reported association, we decided to stop montelukast in addition to the beginning of corticotherapy. This patient also had a past medical history of breast cancer and was under trastuzumab, a monoclonal antibody that was previously reported to cause cutaneous vasculitis [12]. Even though we cannot establish causality between this therapy and the peripheral neuropathy, we believe that this vasculitis must be multifactorial and the chemotherapy may contributed to the clinical picture. To our best knowledge this is the first case of peripheral neuropathy and EGPA described in a patient under trastuzumab. Finally, the last patient had an atypical presentation and course of a possible vascular or vasculitic neuropathy that was assumed to be a part of a systemic autoimmune disease of unknown etiology. The pathological findings in the perinerve were crucial to support this belief and to guide the treatment.

In fact, the pathological studies were very important for the management of our patients, because even without definitive criteria of vasculitic neuropathy, the suggestive neuropathological findings supported our clinical suspicion and supported the use of immunosuppressive therapies. In the first patient, the nerve biopsy was not performed because the patient exhibited typical skin lesions that were more accessible and allowed a prompt diagnosis. In fact, in patients with skin involvement, given its greater accessibility, the skin biopsy can anticipate the decision-making process when the nerve biopsy is delayed, as occurred in two of the patients presented. However, our cases emphasize that the nerve biopsy may be useful to clarify the mechanism of the neuropathy, and it should be considered whenever a vasculitic neuropathy is suspected or as the final step in the diagnostic investigation of a neuropathy of unknown origin [3]. The nerve selection for biopsy was based on careful neurological and neurophysiological examination and a combined muscle biopsy was not performed. Although, it has been suggested that the addition of muscle biopsy improves the yield of definite vasculitis, we also know that one of the most important decisions regarding diagnostic yield is related to the selection of the biopsy site, being crucial to biopsy an affected sensory nerve (with or without muscle) [13,14]. In our experience, the nerve biopsy alone proved to be effective in establishing the diagnosis.

In conclusion, we believe that the etiologic characterization of peripheral neuropathies in the context of systemic vasculitis and other autoimmune diseases is essential to define the natural history and the prognosis of these patients, to prevent potential complications and to optimize the therapeutic approaches. Considering the broad etiologic and phenotypic diversity of peripheral neuropathies, the management of these patients must be individualized in order to identify the underlying etiology.

Declarations

The authors declare that there is no conflict of interest regarding the publication of this paper.

All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional research committee and with the 1964 Helsinki Declaration and its later amendments or comparable ethical standards.

All patients gave written informed consent for publication.

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