Case Report

Cutaneous leukocytoclastic vasculitis accompanied by pulmonary tuberculosis*

Vasculite leucocitoclástica cutânea associada à tuberculose pulmonar

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Abstract
We report the case of a 50-year-old male with a rare combination: pulmonary tuberculosis and cutaneous leukocytoclastic vasculitis. The patient sought emergency treatment presenting with headache, arthralgia, cutaneous rash, and weight loss (4 kg) in the last 20 days. A chest X-ray, performed in a previous outpatient visit, revealed cavitation in the middle and upper lobes of the right lung, as confirmed by computed tomography. Sputum smear microscopy (Ziehl-Neelsen staining) was positive in three consecutive samples, and the result of the skin lesion biopsy was consistent with cutaneous leukocytoclastic vasculitis. The patient was therefore diagnosed with cutaneous leukocytoclastic vasculitis accompanied by pulmonary tuberculosis. Our objective was to describe a combination rarely reported in the medical literature and to discuss the possible pathogenic mechanisms of this combination.

Keywords: Vasculitis, hypersensitivity; Tuberculosis; Hypersensitivity.

Introduction
Vasculitis is a clinicopathologic process characterized by blood vessel wall inflammation.¹ Vasculitis can be primary or secondary to other systemic diseases. Cutaneous leukocytoclastic vasculitis, also known as hypersensitivity vasculitis, is the type of vasculitis most commonly found in clinical practice, and can have several causes, including infections, neoplasms, and use of medications.²

The combination of tuberculosis and vasculitis was first described by Parish & Rhodes in 1967.³ There are three main forms of the combination of pulmonary tuberculosis and vasculitis: pulmonary tuberculosis/cutaneous leukocytoclastic vasculitis (as in the case reported here); pulmonary tuberculosis/Henoch-Schönlein purpura; and pulmonary tuberculosis/vasculitis secondary to rifampicin.⁴⁻⁸

Case report

A previously healthy 50-year-old male driver sought emergency treatment at the Federal University of Paraná Hospital de Clínicas presenting with a history of headache, arthralgia, and skin rash. Arthralgia, which had started 15 days prior, affected the wrist, elbow, ankle, and knee joints. One week later, a maculopapular rash appeared, being distributed in the abdomen as well as in the upper and lower

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lobes of the right lung. The blood workup revealed the following: globular volume, 36.8%; hemoglobin, 12.4 g/dL; leukocytes, 9,130 × 10^3/µL, with 5% rods; platelet count, 294,000 × 10^3/µL; serum creatinine, 0.9 mg/dL; aspartate aminotransferase, 18 mg/dL; alanine aminotransferase, 16 mg/dL; adenosine triphosphate (international normalized ratio), 1.09; anti-HBc, negative; anti-HCV, negative; serological test for syphilis (Venereal Disease Research Laboratory test), negative; and anti-HIV, negative.

After the initial evaluation in the emergency room, the patient was admitted for investigation. The additional laboratory tests performed presented the following results: antineutrophil cytoplasmic antibody, negative; cryoglobulin, negative; antinuclear factor, nonreactive; erythrocyte sedimentation rate, 86 mm in the first hour, and lactate dehydrogenase, 165 U/L. A computed tomography scan of the chest revealed cavitated and centrilobular opacities—presenting a “tree-in-bud” pattern—predominantly in the upper portions of the right lung, suggesting a chronic granulomatous process (Figure 3). Sputum smear microscopy (Ziehl-Neelsen staining) was positive in three consecutive samples, and the patient was classified as a weak reactor on the purified protein derivative test.

The patient was diagnosed with active pulmonary tuberculosis on post-admission day 4. Respiratory isolation was maintained, treatment with regimen I—rifampicin (600 mg/day), isoniazid (400 mg/day), and pyrazinamide (2,000 mg/day)—was instituted, and the patient remained hospitalized. A few days after the initiation of treatment, the patient developed persistent arthralgia in the upper and lower limbs. In addition, the rash was still present, with periods of improvement and periods of exacerbation, sometimes accompanied by pruritus.

Consequently, the hypothesis of hypersensitivity vasculitis secondary to tuberculosis persisted. A skin lesion biopsy revealed cutaneous leukocytoclastic vasculitis. Therefore, the diagnosis of active pulmonary tuberculosis accompanied by hypersensitivity vasculitis secondary to tuberculosis was confirmed.

The treatment with regimen 1 was maintained. Analgesia with tramadol (50 mg/day) and dipyrone (1 g/day) was introduced, as was dexchlorpheniramine (2 mg/day) for relief of the pruritus. The patient showed improvement of the arthralgia and a decrease in the number of skin lesions. We opted...
for outpatient follow-up treatment, and the patient was discharged from the hospital after 20 days.

Discussion

Vasculitis can affect blood vessels of all sizes in any organ, and this results in a wide variety of signs and symptoms in the clinical presentation.\textsuperscript{1,2}

There are different ways of classifying vasculitis. It can be classified as localized, such as isolated cutaneous vasculitis, or systemic, affecting various organs and systems. These can be designated as primary or idiopathic. They can also be designated as secondary when within the context of another pathological process, such as infections, including tuberculosis (as is the case of our patient), collagenosis, neoplasms, and drug hypersensitivity. Currently, the most widely used classification system for vasculitis is the Chapel Hill Consensus Conference system.\textsuperscript{1,2}\textsuperscript{1} According to this classification, cutaneous leukocytoclastic vasculitis is classified as small vessel vasculitis. Histologically, the characteristic lesion is an angiocentric inflammatory process associated with leukocytoclasia (neutrophil fragmentation) and fibrinoid necrosis. It occurs more frequently in small blood vessels, and the pathophysiological mechanism involved is the deposition of immune complexes, usually immunoglobulin M and G, that activate the complement pathways, leading to the production of chemotactic factors and the expression of adhesion molecules. The involvement is usually cutaneous and limited. It can be idiopathic or secondary to infections, neoplasms, collagenosis, or the use of medications. However, in order to diagnose this type of vasculitis, it is necessary to rule out systemic manifestations and glomerulopathies.\textsuperscript{1,2,11}

Since pulmonary tuberculosis is an infectious process, it can manifest as vasculitis, although this combination is rare.

In cases of cutaneous leukocytoclastic vasculitis, \textit{Mycobacterium tuberculosis} is not found in the vessel wall, which differentiates it from cutaneous tuberculosis, in which microorganisms are seen in biopsy samples. Considering that the existence of circulating immune complexes in pulmonary tuberculosis has previously been demonstrated, and that their levels are related to disease activity, the mechanism of injury proposed for this type of vasculitis is deposition of immune complexes (formed by antibodies against antigens of the bacillus) in the vascular wall rather than direct aggression of the bacillus.\textsuperscript{3,12,13}

Clinically, this type of vasculitis can appear as the first symptom of the disease or it can be part of the overall clinical profile. In our patient, the most pronounced profile was that of vasculitis, accompanied by systemic symptoms (fever, headache, arthralgia, and weight loss) that are common to all types of tuberculosis and that also occur in vasculitis. It is more common in young individuals whose immune system is unimpaired, and here we emphasize that our patient had no history of recurrent infections suggestive of immunodeficiency, and that he tested negative for HIV. In addition, no gender-related differences in incidence have been reported.\textsuperscript{14}

A confounding factor is the appearance of vasculitis following tuberculosis treatment with rifampicin, since there are reports of cases demonstrating the relationship between the use of rifampicin and the onset of cutaneous leukocytoclastic vasculitis. In cases of tuberculosis-related vasculitis, the skin lesions improve with the introduction of the treatment regimen, without the need for specific anti-inflammatory treatment, only medication for symptoms, as demonstrated in the case reported here, in which the skin lesions persisted for some days, showing gradual remission until complete resolution. There are host-related factors that facil-
itate the onset of vasculitis, such as blood stasis, hypercoagulability, and, in cases of tuberculosis, the degree of chronic inflammation, the rapid resolution of which leads to remission of vasculitis.[5,7]

Another possible manifestation, though even rarer, is the combination of pulmonary tuberculosis and Henoch-Schönlein purpura: a type of renal involvement in the form of glomerulonephritis, as part of a bout of systemic vasculitis triggered by tuberculosis.[8]

Cutaneous leukocytoclastic vasculitis is treated by treating the underlying disease; in this case, tuberculosis, as previously mentioned. In vasculitis caused by medications (for example, rifampicin), it is recommended that the medication be discontinued and replaced by an alternative regimen, which results in clear lesion improvement. Antihistamines and nonsteroidal anti-inflammatory drugs can be used for symptomatic treatment. Severe skin manifestations can be treated with corticosteroids, the use of which was not necessary in our patient.[2]

This case illustrates one of the multiple clinical presentations of tuberculosis, all of which make it difficult to rapidly identify and control the disease in order to break the transmission cycle.[14,15] It is of note that, in developing countries, the differential diagnosis in cases of weight loss and persistent fever should always be made with tuberculosis, indicating the need for thorough investigation, including pulmonary imaging, sputum smear microscopy, and even lesion biopsy using specific staining. Further studies are needed to elucidate the specific mechanism of the combination of pulmonary tuberculosis and cutaneous leukocytoclastic vasculitis.

References