Insidious onset of intermittent claudication as the primary manifestation of infective endocarditis

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Abstract: Musculoskeletal manifestations of infective endocarditis are well-described in the literature. However, insidious onset of localized calf pain is an uncommon presentation of embolization and may represent a diagnostic challenge owing to the nonspecific nature of the symptoms. This study reviewed the literature and reports a case of infective endocarditis in a patient who presented with bilateral calf pain as the primary complaint and reason for seeking medical attention.

Keywords: musculoskeletal pain, embolization, ischemia, endocarditis

Introduction

Infective endocarditis (IE) presents with variable symptoms, including systemic or cardiovascular manifestations.¹ Some of these symptoms are related to either embolic seeding of the organism or an immunological phenomenon associated with it. Although musculoskeletal symptoms are described in the literature, these are frequently overlooked by physicians.²⁻³ In most reports, patients present with musculoskeletal manifestations resultant from acute limb ischemia,⁴⁻⁶ but more insidious presentations are rare. Failure to recognize this scenario may delay medical attention, contributing to a larger vegetation burden and increasing the mortality rates. Herein, a case is reported in which subacute bilateral calf pain was the primary manifestation of IE.

Case presentation

A 48-year-old man presented to the emergency department with 5 weeks of progressive intermittent claudication that was associated with low-grade fever and worsening dyspnea. His medical history was notable for tobacco use (five packs per day), heavy alcohol consumption, and cannabis use. He denied intravenous drug abuse and reported antibiotic use (amoxicillin) in the week prior to admission.

His vital signs were remarkable – a heart rate of 120 bpm, respiratory rate of 40 rpm, and oral temperature of 39°C. On physical examination, breath sounds were decreased bilaterally and a 3/6 holosystolic murmur was perceived in the apex. Along with this, he had erythematous and swollen gingiva. No erythema or swelling was noted in the lower limbs, but capillary filling time was bilaterally delayed. Complete blood count revealed a white blood cell count of 15.4×10^3 cells/ μ L (22% bands) and normocytic anemia with a hemoglobin level of 7.7 g/dL. Erythrocyte sedimentation rate and C-reactive protein were both increased at 94 mm/h and 185 mg/L, respectively.

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Figure 1 (A) Apical four-chamber view. (B) Long axis view.

Notes: (A) The red arrow shows the mass in the mitral valve during diastole. (B) The blue arrow shows a mass in the anterior leaflet of the mitral valve.

A chest X-ray showed bilateral pleural effusion, and the remaining biochemistry panel, urinalysis, and coagulation tests were within normal limits. Furthermore, serologic tests for HIV, viral hepatitis, and syphilis, in addition to immunologic studies (including antinuclear antibodies and anti-dsDNA) were negative. During evaluation, transthoracic echocardiography revealed a 2.1×2 cm hyperechoic lesion on the anterior leaflet of the mitral valve (Figure 1A: a red arrow shows a mass in the apical four-chamber view. Figure 1B: a blue arrow shows the mass in the long axis view), which was associated with severe mitral regurgitation (regurgitant volume of 73 mL and vena contracta of 19 mm) and an eccentrically directed posterior jet (despite jet angle, the effective regurgitation orifice was $>0.4 \text{ cm}^2$). In addition, the ejection fraction was preserved. Similar findings were confirmed by transesophageal echocardiography. Furthermore, Doppler ultrasound of both inferior limbs showed bilateral arterial thrombosis in the right popliteal artery and in the left common femoral artery, with patency of collateral circulation. A subsequent computed tomography (CT) angiography of the lower extremities showed intraluminal filling defects in left common femoral and iliac artery (Figure 2A) and in the distal popliteal artery of both limbs (Figure 2B). The patient was started empirically on penicillin and gentamicin for possible infectious endocarditis. Blood culture results demonstrated no growth on the day of admission or during the follow-up (day 1, 3, and 5).

Despite the early use of broad-spectrum antibiotics for 4 days, he remained febrile and developed amaurosis fugax in the right eye, but CT scan of the head did not show evidence of an acute cerebrovascular event. His renal function declined, and the patient showed signs of hemodynamic instability with a mean arterial pressure of less than 60 mmHg. Thus, he was taken to the operating room on day 5 of hospitalization for

Figure 2 (A) CT angiography of lower extremities show intraluminal filling defects with occlusion of the left common iliac artery (red arrow) and left common femoral artery (blue arrow). (B) Intraluminal filling defect with occlusion in the distal popliteal artery of both limbs (red and blue arrows).

Abbreviation: CT, computed tomography.

mitral valve debridement and valve replacement with a 27 mm size Carbomedics Orbis metallic valve (Sorin Group, Milan, Italy). After surgery, his fever subsided and clinical condition improved gradually. Histopathological and microbiologic studies (gram stain, bacterial and fungal cultures [Chocolate Agar, Mac Conkey agar, Heart–Brain Infusion agar, Czapek–Dox agar, and CHROMagar]) failed to find any organism. Considering his previous exposure to β -lactam medications and given the absence of etiologic risk factors other than his extensive gingival inflammation, he continued empiric treatment with ceftriaxone for a total of 4 weeks to cover *Streptococci viridans*, *Staphylococcus aureus*, and HACEK group organisms. Peripheral vascular procedures were not pursued due to the improvement of his bilateral calf pain and capillary filling time. Finally, the patient was placed

on anticoagulation for his metallic valve and was enrolled in physical therapy. During follow-up (1, 3, and 6 months), he showed mild improvement in his bilateral claudication, and his overall status was stable.

Discussion

Musculoskeletal manifestations of IE, such as intermittent claudication, are described in the literature but are still poorly defined in medical practice. In a retrospective analysis including 192 cases of IE, 44% of them presented one or more types of musculoskeletal manifestations (mainly arthralgia and low back pain);³ however, only 5.7% of the patients presented myalgia localized to the thigh or calf. In addition, 27% of the patients reported musculoskeletal complaints as one of the first symptoms of this disease. Furthermore, myalgia may be the sole manifestation of IE and can precede the diagnosis by as much as 6 weeks.² No association has been noted between the severity of the rheumatic symptoms and laboratory test abnormalities.³

A number of embolic complications in IE have been reported in the literature. Previous reports have addressed acute embolic effects that may compromise limbs,^{5–7} spleen, or the central nervous system.⁸ Nonetheless, a subtle onset of musculoskeletal complaints, such as in this case, is rare and may lead to an adverse outcome if left unrecognized. The delayed clinical presentation of the patient may be partially explained by the compromise of small and medium-sized arteries as well as the progressive development of collateral circulation. In addition, it is possible that a certain degree of vascular damage was already present given his long-standing history of smoking.

Despite performing several blood cultures (6), no organism was isolated, possibly due to previous antibiotic use, which is the main cause of culture-negative endocarditis.⁹ Clinical guidelines recommend treatment with broad-spectrum antibiotic coverage.⁹ However, on the basis of the local epidemiology,¹⁰ international data of IE,¹¹ and taking into consideration the patient's risk factors (eg, extensive gingival inflammation and the lack of intravenous drug use), it was decided to prescribe ceftriaxone to treat *Streptococci viridans*. Appropriate antibiotic coverage is pivotal to prevent severe embolic events that may occur particularly during the first 2 weeks of the disease,¹² especially if organisms such as *Staphylococcus aureus*,¹³ *Streptococcus bovis*,¹⁴ or fungi are involved. Additional factors such as vegetation characteristics (mobile vegetation length >10 mm, determined by transesophageal echocardiography)^{9,15} may also be useful to estimate the real risk of further embolism, as in the patient (Box 1). Taking into account the aforementioned factors, a careful evaluation is needed when estimating the

Box 1 High risk for embolic events

Anterior mitral leaflet vegetation, particularly with size 10 mm.
One embolic event during first 2 weeks of antimicrobial therapy.
Increase in vegetation size despite appropriate antimicrobial therapy.
Infection with particular microorganisms (*Staphylococcus aureus*, *Streptococcus bovis*, *Candida* spp.)
Previous embolism (after diagnosis)
Multivalvular IE (bivalvular) involvement

Note: Data from Baddour et al⁹ and Habib et al.¹⁶

Abbreviation: IE, infective endocarditis.

real embolic burden as early surgical intervention may be warranted (recently acknowledged in the 2015 European guidelines for IE).¹⁶

Musculoskeletal manifestations of IE may appear early in the course of disease and may be a subtle clue to diagnosis. This case emphasizes the necessity of an early comprehensive examination and a broad differential diagnosis in patients with subacute vascular ischemic symptoms.

Patient consent

Written informed consent was obtained from the patient for publication of this case report and any accompanying images.

Author contributions

George Vasquez-Rios and Maria T Gamero wrote the manuscript and conducted the literature search. George Vasquez-Rios, Maria T Gamero, Jesus de la Cruz, and Roy Dueñas were part of the medical team taking care of the patient. Gabriel A Hernandez and Eduardo Hernandez were consulted for expert opinion. George Vasquez-Rios, Maria T Gamero, Jesus de la Cruz, Gabriel A Hernandez, Eduardo Hernandez, and Roy Dueñas critically revised the manuscript for important intellectual consent. All authors contributed toward data analysis, drafting and critically revising the paper and agree to be accountable for all aspect of the work.

Disclosure

The authors report no conflicts of interest in this work.

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