

Scientific letters

Acute Ischemia in Upper Limb Secondary to Myxoma

Isquemia aguda en extremidad superior secundaria a un mixoma

To the Editor,

Cardiac tumors are an unusual cause of acute limb ischemia. Myxomas are the most frequent benign tumors and most are symptomatic. Embolization occurs in 30% to 40% of patients with myxomas, with cerebral arteries being the most common destination.¹

We present the case of a 73-year-old woman with a history of hypertension who came to the emergency department due to pain and sudden onset of coldness in the left limb, which developed over 3 h. On examination, she had a brachial pulse in the bicipital groove, without distal pulse or deficits in sensitivity and motility. Cardiac auscultation was normal and the electrocardiogram showed no abnormalities in rhythm, just an incomplete blockage of the right branch. With the diagnosis of acute limb ischemia, the patient was urgently operated on. A thrombectomy of the brachial, radial, and ulnar artery was performed, obtaining abundant material that was gelatinous and friable to the touch, very different from a typical thrombus (Fig. 1A). After surgery, the patient recovered distal pulses. Suspecting a cardiac tumor, an urgent transthoracic echocardiogram was ordered, which showed a mobile mass with a wide attachment base, anchored to the interatrial septum. The tumor was approximately 36 mm × 25 mm, which prolapsed into the left ventricle, producing probable traumatic severe mitral regurgitation associated with the traction of the anterior mitral leaflet (Fig. 2). The size and function of the left ventricle were normal. These findings were confirmed by a transesophageal echocardiogram. The study was completed with a cardiac catheterization that resulted in a 70% to 80% stenosis in the middle part of the anterior descending artery. After anesthesia assessment, the patient was taken to the cardiac surgery department for intervention. We proceeded with bi-atrial resection of the tumor with coronary bypass of the left mammary

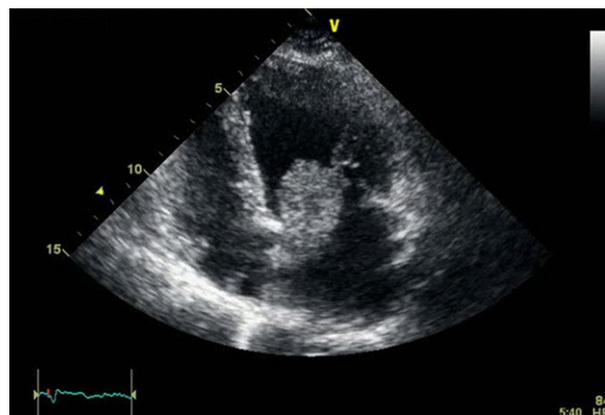


Figure 2. Image of the transthoracic echocardiogram that shows a tumor attached to the septum prolapsing into the ventricle.

artery to the anterior descending artery, and mitral valve repair using annuloplasty with Edwards ring. The resected tumor was 55 mm × 40 mm (Fig. 1B), and the interatrial septum defect was repaired with a bovine pericardium patch. The diagnosis of myxoma was confirmed after pathological study of the tumor. After 1 year of follow-up, the patient was asymptomatic and had no relapses.

Myxomas originate in the mesenchymal cells in the endocardium of the septum. They have an incidence rate of 0.5% per million inhabitants/year. They are more frequent in women by a 3:1 margin. Most are located in the left atrium (87%) in the septum near the fossa ovalis. Other less frequent locations include the mitral valve, the right atrium, and the ventricles.²

There are familial forms, called the Carney complex, that are inherited in an autosomal dominant manner, characterized by

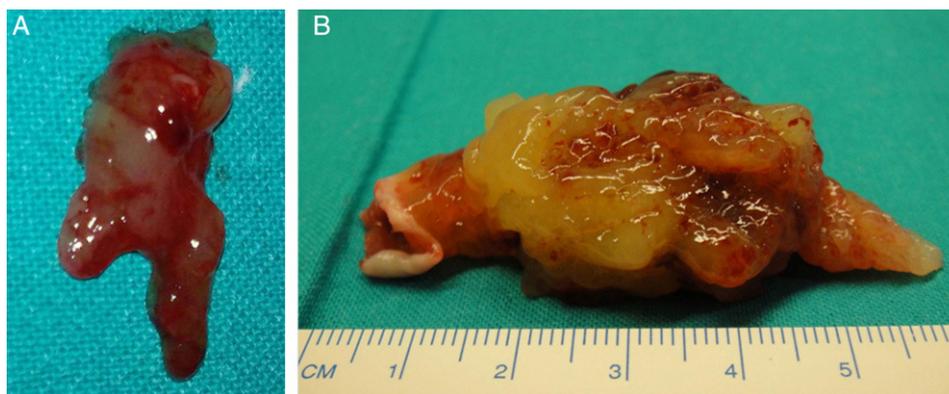


Figure 1. A, embolus extracted from the brachial bifurcation of the arm. B, image of the resected tumor.

mesenchymal tumours, cutaneous lentiginosis, hyperfunctional endocrine disorders, and tumors of the peripheral nerves.³ Symptoms depend on location, size, and mobility of the tumor. Symptoms may be due to obstruction of the ventricle, with the onset of dyspnea, secondary to pulmonary edema, syncope, and even sudden death. Systemic embolization occurs in 25% to 40% of cases due to tumor fragmentation, with the brain being the most frequent destination. Limb embolisms, as in our case, in visceral, coronary, and renal arteries are rare. Complete embolization of the tumor has also been reported. Patients may present with symptoms including fever, skin rash, joint pain, and weight loss, as a result of the production of cytokines (interleukin-6) and growth factors by the myxoma.⁴

Transthoracic echocardiography is the method of choice for the diagnosis of cardiac myxomas. However, transesophageal ultrasound has greater sensitivity (almost 100%) than transthoracic ultrasound (90%).⁵ The treatment of choice is surgical resection even in asymptomatic patients. It must be performed very soon after diagnosis due to the risk of embolism and sudden death. Surgical mortality varies from 2% to 5%, and the prognosis after resection is excellent. Recurrence of these tumors varies from 0.4% to 5% at 22 years of treatment. Therefore, follow-up with echocardiography is recommended at least every 5 years. Cardiac myxomas have greater recurrence in young males, multifocal cases, and familial forms.⁶ In these patients, follow-up must therefore be more stringent.

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REFERENCES

1. Klaus R. Cardiac myxomas. *New Engl J Med*. 1995;333:1610-7.
2. Mac Gowan SW, Sidhy P, Aherne T, Luke D, Wood AE, Neligan MC, et al. Atrial myxoma: national incidence, diagnosis and surgical management. *Ir J Med Sci*. 1993;162:223-6.
3. Goldstein MM, Casey M, Carney JA, Basson CT. Molecular genetic diagnosis of the familial myxoma syndrome (Carney complex). *Am J Med Genet*. 1999;86:62-5.
4. Sakamoto H, Sakamati T, Kanda T, Tsuchiya Y, Sato M, Sato H, et al. Vascular endothelial growth factor is an autocrine growth factor for cardiac myxoma cells. *Circ J*. 2004;68:488-93.
5. Manning WJ, Weintraub RM, Waksmonski CA, Haering JM, Rooney PS, Maslow AD, et al. Accuracy of transesophageal echocardiography for identifying left atrial thrombi. A prospective intraoperative study. *Ann Intern Med*. 1995;123:817-22.
6. Keeling IM, Oberwalder P, Anelli-Monti M, Schuchlenz H, Demel U, Tilz GP, et al. Cardiac myxomas 24 years of experience in 49 patients. *Eur J Cardiothorac Surg*. 2002;22:971-7.

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Severe Community-Acquired Methicillin-Resistant *Staphylococcus aureus* Endocarditis in a Child With Structurally Normal Heart: a Case Report

Endocarditis bacteriana grave por *Staphylococcus aureus* resistente a metilina adquirida en la comunidad en un niño con corazón estructuralmente normal

To the Editor,

A 2-year-old male infant was admitted to the hospital with a history of rapid decline in general health and persistent hyperthermia (38.5 °C to 39 °C) over the previous 6 days. No history of congenital heart disease or previous hospitalization was reported by parents. General inspection revealed eutrophic child, easily irritable, and small pustular lesions over both legs. Rapid pulse (120 bpm), dry and pale mucosa, and bilateral subconjunctival hemorrhagic spots were noted. Large erythematous area over the right hypochondrium, diffuse petechial exanthema over the lower limbs, and left proximal forefinger interphalangeal joint swelling were also observed (Fig. 1).

Cardiac examination revealed hyperdynamic precordium and 3+/4+ systolic murmur was heard over the apex irradiating to the left axilla and posterior torso.

Laboratory findings on admission were white blood cell count 24 500/μL with 25% rod-like neutrophil, platelet 21 000/μL, hematocrit 24.7%, creatinine 0.4 mg/dL, GGT 54 IU/L, alkaline phosphatase 518 IU/L, sodium 130 mEq/L, and potassium 2.9 mEq/L. Bidimensional color Doppler echocardiogram with parasternal short axis and apical long axis views showed left chambers enlargement (left ventricular diameter 37 mm diastolic and 23 mm systolic). A filamentary vegetation was observed extending from the ventricular aspect of the anterior mitral valve leaflet to

the left ventricular outflow tract adherent to the noncoronary aortic cusp (Figs. 2A and B; Movies 1 and 2). Severe mitral and mild aortic regurgitations were also present (Movie 3).

An acute infective endocarditis (IE) was thus considered and before starting an empiric antibiotic schedule (ceftriaxone, oxacillin, and vancomycin), two blood specimens were cultured. A surgical intervention was considered but disapproved by the cardiac team.

In 48 h, metabolic acidosis, respiratory distress with sign of spontaneous bleeding from lung parenchyma, and anisocoria evolved. New echocardiogram showed that sub-aortic filamentary vegetation was no longer present and right ventricle was dilated. A new image adhered to the septal tricuspid valve leaflet was observed (Figs. 2C and D; Movies 4 and 5). The child evolved with bilateral mydriasis, progressing to death.

Staphylococcus aureus (*S. aureus*) was isolated from all blood samples, and was sensitive to vancomycin, clindamycin, and sulfamethoxazole-trimethoprim and resistant to both oxacillin and penicillin.

Current report describes a case of aggressive IE caused by *S. aureus* involving 3 cardiac valves in an otherwise healthy male infant. In children, IE is commonly associated with congenital or rheumatic heart disease.¹ In healthy children, bacteremia secondary to skin infection or due to central catheters invasion, particularly in neonates, is associated with IE, and *S. aureus* has been the most frequent infective agent isolated from blood cultures.^{1,2} Community-acquired methicillin-resistant *S. aureus* (CA-MRSA) is a rapidly growing worldwide problem.³ Reassuringly, the prevalence of CA-MRSA colonization in a community's young children remains very low.⁴ Milstone et al. reported a 61% CA-MRSA prevalence among all MRSA-colonized children in a pediatric intensive care unit.⁵ Jaggi et al. reported that invasive CA-MRSA represented 12.7% of all MRSA strain isolated in pediatric ICU, but none related to IE.⁶ Children with CA-MRSA infections