# BRIEF REPORTS

# Infected Left Atrial Myxoma

Antonio García-Quintana,ª Pedro Martín-Lorenzo,ª Javier Suárez de Lezo,ª Marta Díaz-Escofet,ª Rafael Llorens,<sup>b</sup> and Alfonso Medina<sup>a</sup>

<sup>a</sup>Servicio de Cardiología, Hospital Universitario de Gran Canaria Dr. Negrín, Las Palmas de Gran Canaria, Spain.

<sup>b</sup>Servicio de Cirugía Cardíaca, Hospiten Rambla, Santa Cruz de Tenerife, Spain.

Myxoma is the most common primary tumor of the heart. It is uncommon for these tumors to become infected and, at times, clinical presentation is no different from that of an uninfected myxoma. We describe the case of a 58year-old woman with a previous pharyngeal infection that developed into infection of a left atrial myxoma and which was complicated by systemic embolism affecting the lower limbs. Streptococcus oralis was identified in blood cultures and embolic material. The tumor was resected and the patient's subsequent clinical evolution was uneventful. The incidence of infected myxoma is very low. A review of the literature based on individual case reports is presented.

Key words: Myxoma. Endocarditis. Tumor. Echocardiography.

#### Mixoma auricular izquierdo infectado

El mixoma es el tumor primario más frecuente que afecta al corazón. La infección de estos tumores es infrecuente y, en ocasiones, su presentación clínica es indistinguible de un mixoma no infectado. Describimos el caso de una mujer de 58 años con una infección faríngea previa que desarrolló infección sobre un mixoma auricular izquierdo, complicado con embolia sistémica en los miembros inferiores. Se aisló Streptococcus oralis en los hemocultivos y en el material embólico. El tumor se resecó y la evolución clínica posterior resultó sin incidencias. La frecuencia del mixoma infectado es muy baja y una revisión de la bibliografía se ha de basar en los casos aislados publicados.

Palabras clave: Mixoma. Endocarditis. Tumor. Ecocardiografía.

# INTRODUCTION

Primary tumors of the heart are rare, with an incidence of 0.0017%-0.19% in autopsy series.<sup>1</sup> Three quarters of the tumors are benign; half of these are myxomas and the rest mostly lipomas, papillary fibroelastomas, and rhabdomyomas. Myxomas are more common among women and can affect both atria,<sup>2</sup> the ventricles,<sup>3</sup> or the mitral valve, although the left atrium is most commonly involved. The clinical presentation is characterized by obstruction of the mitral valve,<sup>4</sup> embolism, and constitutional symptoms, in addition to fever, anemia, or an elevated erythrocyte

E-mail: antoniogarcia@secardiologia.es

Received November 18, 2004. Accepted for publication March 14, 2005. sedimentation rate; nevertheless, infection of these tumors is rare. The differential diagnosis is difficult, particularly with uninfected myxoma and mural endocarditis and therefore, diagnostic criteria have been proposed.5

## **CASE STUDY**

A 58-year-old woman with a history of bilateral breast cancer treated by surgery and adjuvant radiotherapy and chemotherapy in complete remission was admitted for symptoms of confusion, fever, and abdominal pain of 3 days' evolution. She reported constitutional syndrome and fever during the preceding 6 months, following an episode of acute pharyngitis. In the physical examination, she presented cachexia, a temperature of 40°C, and tachycardia with no murmurs. The analyses disclosed leukocytosis (22 300 leukocytes/mL), normocytic anemia, and an erythrocyte sedimentation rate (ESR) of 107 mm/h.

Correspondence: Dr. A. García-Quintana.

Servicio de Cardiología. Hospital Universitario de Gran Canaria Dr. Negrín. Bco. La Ballena, s/n. 35020 Las Palmas de Gran Canaria. España.



**Figure 1.** Echocardiogram: mass with pedicle, of heterogeneous density and joined to the interatrial septum (A). The mass is adhering to the interatrial septum and has prolapsed through the mitral valve (B). Ao indicates aorta; LV, left ventricle; RA, right atrium; RV, right ventricle; asterisk, myxoma.

The echocardiogram showed a vegetative mass with pedicle in the left atrium,  $5.4 \times 1.5$  cm in size, of heterogeneous density and adhering to the interatrial septum, with prolapse in the left ventricle, but no significant mitral valve obstruction or regurgitation (Figure 1). Empirical antibiotic therapy with ampicillin and gentamicin was started and blood was drawn for cultures, which were positive for *Streptococcus oralis*.

TABLE. Criteria for the Diagnosis of Infected Atrial Myxoma

Definitive	<ol> <li>Myxoma documented by histology, and</li> <li>A. Microorganisms observed in the sample, or</li> <li>b. Positive blood cultures and evidence of</li> </ol>
	inflammation in the sample
Probable	1. Myxoma documented by histology, and
	<ol><li>Positive blood cultures and evidence of inflammation in the sample</li></ol>
Possible	<ol> <li>Characteristic appearance on transthoracic or transesophageal echocardiography, and</li> <li>Positive blood cultures</li> </ol>

Taken from Horstkotte et al.8



**Figure 2.** Arteriography: occlusion of left common iliac and right deep femoral artery. Arrow: area of occlusion (A). Echocardiogram: a decrease in the size of the mass (B) is observed after the embolic event. LV indicates left ventricle; RA, right atrium; RV, right ventricle; arrowhead, myxoma after embolism.

Forty-eight hours after admission, the patient presented sudden pain and pallor in the lower left limb, with no femoral pulse. Arteriography showed embolism of the left common and external iliac arteries and the right deep femoral artery (Figure 2), and surgical embolectomy was performed. The embolic material showed *Streptococcus oralis* on its surface. Echocardiographic follow-up revealed a pronounced decrease in tumor size (3×2 cm) (Figure 2).

Due to the risk of further embolism formation, the mass was surgically resected. Histological study disclosed the presence of fusiform cells surrounded by a lax mucopolysaccharide-rich stroma, as well as the presence of Gram-positive cocci. Antibiotic therapy was continued for one week, and the patient's clinical evolution was uneventful.

### DISCUSSION

Criteria have been proposed to aid in the diagnosis of infected myxoma (Table). Our literature review revealed 35 definitive cases, 5 probable and 1 case of possible infected myxoma,<sup>5</sup> with 45% having risk factors that could have contributed to the infection: dental work (22%), recent infections (10%), invasive procedures (5%), use of intravenous drugs, long-term use of corticoids, and poor dental health. The microorganisms involved were *Streptococcus viridans* (44%) and *Staphylococcus aureus* (15%), a microbiological spectrum similar to that of native valve endocarditis. Bacteremia does not prove that the myxoma is infected, as there have been reports of positive blood cultures while the tumors show no inflammation or infection.

The differential diagnosis of infected myxoma<sup>6</sup> mainly includes uninfected myxoma, since fever can appear in the absence of infection, as well as mural endocarditis and infected intracardiac thrombus. Myxoma is associated with systemic embolism,<sup>7</sup> although the risk appears to be greater when the myxoma is infected.

Surgery usually resolves the condition but should be done early; operative mortality is low. There is some tendency to maintain the standard antibiotic regimen for endocarditis,<sup>8</sup> although patients treated for less than two weeks do not appear to experience more complications.<sup>5</sup>

Since the clinical presentation of infected myxoma may be similar to that of uninfected myxoma, blood cultures should be done whenever a patient with myxoma presents fever, and echocardiography should be performed in patients with fever of unknown origin when the initial techniques are not conclusive.<sup>9,10</sup>

#### REFERENCES

- 1. Reynen, K. Cardiac myxomas. N Engl J Med. 1995;333:1610-7.
- Jiménez-Navarro MF, Carlos Gavilan J, María Melero J, Rodríguez Bailón I, Bermúdez F, Porras C, et al. Mixoma de gran tamaño en la aurícula derecha. Rev Esp Cardiol. 2001;54:399-401.
- Ramírez Moreno A, Anguita Sánchez M, Castillo Domínguez JC, Siles Rubio JR, Franco Zapata M, Casares Mediavilla J, et al. Mixoma ventricular izquierdo aislado descubierto casualmente por ecocardiografía. Rev Esp Cardiol. 1998;51:763-5.
- Ulecia Martínez MA, Torres Ruiz JM, Chamorro Santos CE, Moreo Herrero T. Cirugía emergente por mixoma auricular izquierdo. Rev Esp Cardiol. 2000;53:1279-80.
- Revankar SG, Clark RA. Infected cardiac myxoma: case report and literature review. Medicine (Baltimore). 1998;77:337-44.
- Gabe ED, Rodríguez Correa C, Vigliano C, San Martino J, Wisner JN, González P, et al. Mixomas cardíacos: correlación anatomoclínica. Rev Esp Cardiol. 2002;55:505-13.
- Luaces Mendez M, Vilacosta I, Sarria C, Fernández C, San Román JA, Sanmartín JV, et al. Endocarditis infecciosa y embolias del eje hepatoesplenorrenal. Rev Esp Cardiol. 2004;57:1188-96.
- Horstkotte D, Follath F, Gutschik E, Lengyel M, Oto A, Pavie A, et al. Guía de Práctica Clínica sobre prevención, diagnóstico y tratamiento de la endocarditis infecciosa. Rev Esp Cardiol. 2004;57: 952-62.
- 9. Dekkers P, Elbers HR, Morshuis WJ, Jaarsma W. Infected left atrial myxoma. J Am Soc Echocardiogr. 2001;14:644-5.
- 10. Gregory SA, O'Byrne WT, Fan P. Infected cardiac myxoma. Echocardiography. 2004;21:65-7.