

Intravascular Leiomyomatosis With Cardiac Extension: Simultaneous Abdominal and Cardiac Surgery[☆]



Leiomiomatosis intravascular con extensión intracardiaca: cirugía abdominal y cardíaca en un solo tiempo

Intravenous leiomyomatosis is a histologically benign, rare disease in which smooth muscle cells proliferate intravascularly without invading other structures. They reach the intrauterine veins and, occasionally, can reach the retroperitoneal vessels and even the right cardiac chambers. It usually affects women aged 40–50.

Few case reports have been reported. We present an uncommon case and briefly comment on a review of the literature.

The patient is a 53-year-old woman who was in follow-up by cardiology due to episodes of syncope. Upon physical examination, a pelvic mass was palpated that reached the navel region, which was hard in consistency and painful to the touch. Thoracoabdominal-pelvic CT scan with intravenous contrast (Fig. 1) showed evidence of a multilobulated mass with heterogenous densitometry corresponding with a myomatous/sarcomatous uterus, thrombosis of the right ovarian vein and involvement of the inferior vena cava up to the right atrium. Echocardiogram demonstrated a dilated right ventricle; the right atrium presented a mobile homogeneous mass measuring 26 mm×35 mm that prolapsed towards the right ventricle; the inferior vena cava was also occupied. Biopsy results defined the lesion as a leiomyoma. The patient study was completed with PET/CT, which showed a mildly hypermetabolic tumour, suggesting uterine leiomyoma, and hypometabolism in the inferior vena cava and right atrium, not suggestive of tumour thrombosis.

On 6 May 2015, surgery was performed by 2 surgical teams (General and Digestive Tract Surgery together with Cardiac Surgery) in 1 single operation (410 min), which included: resection of the uterine leiomyoma by hysterectomy with bilateral adnexectomy; excision of the intercaval and right interatrial tumour expansion, requiring cardiopulmonary bypass without cardiac arrest. The surgical specimens were sent for pathology study, which confirmed intravenous leiomyomatosis (Fig. 2).

The patient was admitted to the Recovery Unit for 5 days and was later transferred to the Abdominal Transplantation Unit, where the patient's condition improved satisfactorily. She was discharged 23 days after surgery.

Intravenous leiomyomatosis affects women, mainly after hysterectomy due to uterine leiomyomata.¹ In this condition, these benign smooth muscle tumours demonstrate intravascular expansion and can even reach the heart chambers through the inferior vena cava, obstructing

venous return. In 2013, fewer than 300 cases were registered in the English-speaking medical literature.²

Some theories for the origin and growth mechanisms of these tumours include³:

- Intravenous local extension of leiomyomata
- Direct carcinogenesis from the vascular walls within the myometrium; the tumour can float freely within the lumen of the vessel or adhere to it.

The diagnosis should be considered in cases of deep vein thrombosis, mass within the inferior vena cava or atrium, or non-specific symptoms associated with a history of pelvic surgery, such as hysterectomy.⁴

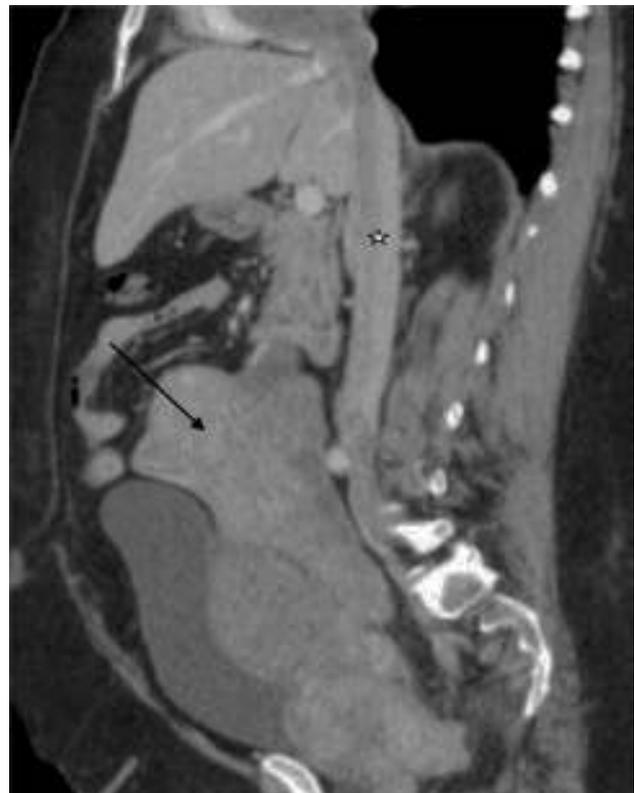


Fig. 1 – CT scan with a large uterine tumour (arrow) and tumour growth in the iliac, ovarian and caval veins (star), up to the right atrium.

[☆] Please cite this article as: Calvo-Rodríguez D, Narváez Muñoz A, Fernández Pérez A, Durán D, Varo Pérez E. Leiomiomatosis intravascular con extensión intracardiaca: cirugía abdominal y cardíaca en un solo tiempo. Cir Esp. 2016;94:306-307.



Fig. 2 – Surgical specimen with a large uterine mass and tumour growth in the iliac veins, cava and right atrium.

Symptoms are variable, including: pelvic pain and vaginal bleeding; respiratory and cardiac symptoms (dyspnoea, cough, right heart failure). On certain occasions, the intracardiac extension of these tumours can lead to sudden cardiac death.⁵

Diagnostic studies involve:

- Tomography: provides evaluation of the tumour extension.
- Magnetic resonance: shows low to intermediate signal intensity in T1.
- Pelvic ultrasound: studies the uterine adnexa and the presence of intravascular thrombus; also assesses intracardiac involvement.⁶

The treatment of choice is surgery with complete tumour excision to avoid recurrences. This can be done in two operations (less risk for haemorrhage due to anticoagulation during cardiopulmonary bypass) or in one (good results have been observed).⁷ There have been reports of cases in

which cardiac arrest was necessary for correct tumour excision.

In cases of incomplete resection or tumour recurrence, the use of anti-oestrogen drugs such as tamoxifen is controversial. Recurrence rates of up to 30% have been reported in incomplete resections, and recurring intravenous leiomyomatosis may appear 6 months to 15 years later; therefore, long-term follow-up is essential.⁸

REFERENCES

1. Demirkiran F, Sal V, Kaya U, Alham C, Tokgozoglu N. Intravenous leiomyoma with extension to the heart: a case report and review of the literature. *Case Rep Obstet Gynecol.* 2013;2013. ID 602407.
2. Guzman L, Dergal E, Capurso A, Hernández Valencia JA, Garmilla Espinosa J. Leiomiomatosis intravenosa: informe de una paciente. *Cir Gen.* 2000;22.
3. Cruza I, Joãoa I, Stuarda B, Lala M, Bento L, Cotrim C, et al. Intravenous leiomyomatosis: a rare cause of intracardiac mass. *Rev Port Cardiol.* 2014;33. 735.e1-5.
4. Morales M, Anacleto A, Leal J, Carvalho S, Derl'Arco J. Intravascular leiomyoma with heart extension. *Clinics.* 2012;67:83-7.
5. Jung Ch, Myung S, Moon S. A case of huge intravenous leiomyomatosis with intracardiac extension. *J Women Med.* 2010;3:29-31.
6. Clay T, Dimitriou J, McNally O, Rusell PA, Newcomb AE, Wilson AM. Intravenous leiomyomatosis with intracardiac extension: a review of diagnosis and management with an illustrative case. *Surg Oncol.* 2013;22:e44-52.
7. Qiang L, Jie M, Bin H, Xiatao P, Hao L. One-stage surgical removal of intravenous leiomyomatosis with right heart extension is safe. *Heart Surg Forum.* 2010; 115014.
8. Moniaga N, Randall L. Uterine leiomyomatosis with intracaval and intracardiac extension. *Gynecol Oncol Rep.* 2012;2:130-2.

Dania Calvo-Rodríguez^{a,*}, Adrián Narváez Muñoz^b, Aquilino Fernández Pérez^a, Darío Durán^b, Evaristo Varo Pérez^a

^aUnidad de Trasplante Abdominal, Hospital Clínico Universitario de Santiago de Compostela, Santiago de Compostela, Spain

^bServicio de Cirugía Cardíaca, Hospital Clínico Universitario de Santiago de Compostela, Santiago de Compostela, Spain

*Corresponding author.

E-mail address: daniamjcr@gmail.com (D. Calvo-Rodríguez).

2173-5077/

© 2015 AEC. Published by Elsevier España, S.L.U. All rights reserved.