Liposarcomas represent the second most common type of soft tissue sarcomas after malignant fibrous histiocytoma. There are four histological subtypes according to the World Health Organization: well-differentiated (atypical lipomatous) liposarcoma, dedifferentiated, myxoid (round-cell) liposarcoma \(1\), and pleomorphic liposarcoma \(1\). Their primary anatomical sites of origin from liposarcomas are the retroperitoneum in 25-50\% and the lower limbs in 25-35\%. The most frequent distant sites for metastasis arising from ML include lungs, retroperitoneum, abdominal cavity, and chest wall \(1\). ML accounts for 30-50\% of all liposarcomas, with a peak incidence between the third and the fifth decade of life, being more frequent in males than females, exhibiting an overall metastatic rate of 30\%. Interestingly, it has an expansive growth pattern, with less aggressive infiltrative behavior, thus, causing frequently obstructive clinical syndromes at the main site of metastasis, however, metastatic intracardiac soft tissue sarcomas represent an extremely rare clinical entity \(1\). Some authors believe that in patients with a known history of liposarcoma, the time interval between diagnosis and the occurrence of distant metastatic disease ranges from 3 to 25 years \(1\).

Fulminant right ventricular (RV) failure from RV metastasis causing outflow obstruction, cardioembolic complications, such as ischemic stroke or significant massive pulmonary tumor embolism (PTE) could occur, carrying catastrophic consequences \(2-6\). PTE complicating RV metastasis from ML represents an extremely difficult premortem diagnostic challenge for clinicians. Herein, we describe, to be best of our knowledge, the first case, of a rapidly progressive hematogenous metastatic ML to the RV complicated with extensive PTE with catastrophic outcomes.

Case description

A 63-year-old Hispanic woman with medical history of hypertension and hyperlipidemia was transferred to...
our hospital, with a significant history of 5-week progressively worse dyspnea; in the previous hospital, she was diagnosed with pneumonia; however, despite antibiotics given, her condition started to deteriorate over the following weeks. The patient also reported unknown weight loss in the past 4 months, as well as progressively worse right hip pain and discomfort with subjective swelling for the past 2 months, provoking her a couple of falls.

On physical examination, her blood pressure was 105/70 mmHg, heart rate at 105 beats per minute, afebrile with an oxygen saturation of 94% at 4L/NC; cardiac auscultation revealed significant tachycardia; lung auscultation with bilateral lungs clear; right hip with pain on deep palpation, and exacerbation of pain on extension and flexion of it; and no skin discolorations noted.

Laboratory findings were notable for severe thrombocytopenia of 25,000 cells/mm³ (normal > 140,000 cells/mm³) and lactate dehydrogenase of 1042 U/L (normal < 250 U/L). 12-lead electrocardiogram showed sinus tachycardia. Chest radiography showed bilateral basal multifocal airspace opacities, suggesting possible infarcts or atelectasis. Bilateral lower extremity venous Doppler ultrasonography did not reveal any deep vein thrombosis.

Transthoracic echocardiography showed a “D”-shaped morphology of the left ventricle during systole, left ventricular ejection fraction between 60% and 65%; moderate right atrial enlargement, severe RV dilation, and hypokinesis with positive McConnell’s sign, with multiple intraventricular echodensities, suggestive of RV thrombi, some of them pending in the posterior leaflets of the tricuspid valve, and other attached to the free wall of the RV, biggest thrombi dimensions are 1.5 cm x 1.7 cm (Fig. 1A). Computed tomographic angiogram of the chest showing intracavitary RV thrombi with right main and left lobar bilateral pulmonary emboli extension, and absence of contrast enhancement within the filling defects (B). Please, also, see supplementary videoclip of the transthoracic echocardiography in the electronic version of this article.

Computed tomographic (CT) angiography of the chest for pulmonary embolism protocol showed central right and left main bilateral pulmonary emboli, extending into the lobar arteries; multiple thrombi filling the RV, with bilateral diffuse lower lobes ground glass opacities (Fig. 1B). CT with contrast of the abdomen and pelvis showed extensive peripherally enhancing fluid collections within the right hip and right thigh musculature, measuring 14.3 cm x 14.8 cm. Pathologic macroscopic and microscopic descriptive findings are shown (Fig. 2A-2F).

During her hospital course, she was managed with intravenous argatroban drip, given her severe consumptive thrombocytopenia. Orthopedic surgery performed a fine-needle aspiration biopsy on her right hip, given her extensive lesions concerning for malignancy, and given her very poor surgical candidacy and unacceptable
high risk for major bleeding events. The patient’s clinical condition rapidly deteriorated in the next 24-48 h, to the point that she became severely hypoxemic, hypotensive, requiring mechanical ventilatory support, and vasopressor therapies. Unfortunately, the patient passed away, after her family members decided to withdraw care through her 4th day of hospital stay.

Discussion

Although primary cardiac tumors are rare (between 0.01% and 0.1% in postmortem studies), the frequency of cardiac metastasis is more common, ranging between 0.5% and 3.7% in the general population;

however, its frequency in patients with known malignancy varies from 9.1% to 14.5% in patients with advanced metastatic malignancies. The symptomatology from cardiac metastasis is non-specific, and the clinical features are highly variant and dependent on the location and disseminated tumor burden.

Cardiac metastasis can spread to the heart mainly by four mechanisms: direct extension or contiguity, hematogenous spread, lymphatic spread, or by transvenous diffusion. Tumors such as renal cell carcinomas and hepatocellular carcinomas can extend into the inferior vena cava, and right atrium, through transvenous diffusion. Certain malignancies such as renal cell carcinomas and hepatocellular carcinomas can extend into the inferior vena cava and right atrium, through transvenous diffusion.

Pericardial metastasis is the most common site of invasion, with an estimated frequency of 60-69% of all necropsy studies; epicardial and myocardial involvement represent the second and third most common sites, in approximately 25-34% of cases; endocardial and intracavitary are rare, representing 3-5% on necropsy studies; however, they can present with exaggerated and life-threatening clinical features with immediate negative outcomes. Cardiogenic shock due to fulminant RV failure from RV metastasis causing outflow obstruction, significant cardioembolic complications from tumor embolism, causing ischemic stroke or pulmonary tumor emboli, like in our patient, can carry catastrophic outcomes.

Intracardiac metastatic solid sarcomas represent extremely rare clinical entities. From <40 cases of metastatic cardiac ML that has been reported in medical literature in English language since 1968, only 13 cases have been reported in the medical literature involving the RV since the first case reported by Godwin et al., in 1981.

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massive PTE, complicating RV metastasis from ML, and ending up with devastating consequences.

**Conflicts of interest**

None.

**Funding**

None.

**Ethical disclosures**

Protection of human and animal subjects. The authors declare that no experiments were performed on humans or animals for this study.

Confidentiality of data. The authors declare that no patient data appear in this article.

Right to privacy and informed consent. The authors declare that no patient data appear in this article.

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