

Revista Portuguesa de **Cardiologia**Portuguese Journal of **Cardiology**

www.revportcardiol.org



CASE REPORT

Carcinoid heart - Case report*

Ana Rita Santos^{a,*}, Bruno Stuart^b, Carlos Cotrim^b, Ana Rita Almeida^b, Paula Fazendas^b, Liliana Lopes^b, Inês Cruz^b, Hélder Pereira^b

- ^a Servico de Cardiologia, Hospital do Espírito Santo, EPE, Évora, Portugal
- ^b Serviço de Cardiologia, Hospital Garcia de Orta, Almada, Portugal

Received 3 December 2011; accepted 26 January 2012 Available online 13 September 2012

KEYWORDS

Carcinoid heart disease; Transthoracic echocardiography; Three-dimensional echocardiography Abstract We describe the case of a 76-year-old woman, diagnosed with a neuroendocrine tumor of the cecum in 2004, with liver metastases and carcinoid syndrome since September 2010. The patient had been treated intermittently with chemotherapy cycles, and remained symptomatic, with worsening secondary lesions. In June 2011 she began to present signs and symptoms of right heart failure and was hospitalized in September 2011. Transthoracic two- and three-dimensional echocardiography revealed enlarged right atrium and ventricle and thickened and fixed tricuspid and pulmonary valve leaflets, causing severe tricuspid regurgitation and mild pulmonary regurgitation and stenosis, suggestive of carcinoid heart disease. The authors discuss the clinical importance of transthoracic echocardiography, and the more recent three-dimensional echocardiography as the diagnostic exam of choice in these cases, as it is especially suitable for assessing the valves and subvalvular apparatus.

© 2011 Sociedade Portuguesa de Cardiologia. Published by Elsevier España, S.L. All rights reserved.

PALAVRAS-CHAVE

Doença carcinóide cardíaca; Ecocardiografia transtorácica; Ecocardiografia tridimensional

Coração carcinóide - a propósito de um caso clínico

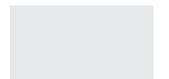
Resumo Os autores apresentam o caso de uma doente de 76 anos, com o diagnóstico de tumor neuroendócrino do cego desde 2004 com metastização hepática, e com síndrome carcinóide desde setembro de 2010. A doente realizou ciclos de quimioterapia de forma intermitente, pelo que manteve sintomas e agravamento das lesões secundárias. Desde junho de 2011 apresenta sinais e sintomas compatíveis com insuficiência cardíaca direita, ficando internada em setembro de 2011. Nesta sequência realizou ecocardiograma transtorácico bi e tridimensional que revelou cavidades direitas dilatadas e folhetos das válvulas tricúspide e pulmonar espessados, fixos, condicionando insuficiência tricúspide grave e estenose e insuficiência pulmonar ligeiras, sugestivas de doença carcinóide cardíaca. Os autores discutem a relevância clínica da ecocardiografia transtorácica, e, atualmente, da ecocardiografia tridimensional, como o

E-mail address: anarita_asantos@hotmail.com (A.R. Santos).

^{*} Please cite this article as: Santos, A.R. Coração carcinóide – a propósito de um caso clínico. Rev Port Cardiol 2012. http://dx.doi.org/10.1016/j.repc.2012.01.020.

^{*} Corresponding author.

662 A.R. Santos et al.



método de investigação de escolha nestes casos, ao permitir avaliar de forma única as válvulas e o aparelho subvalvular.

© 2011 Sociedade Portuguesa de Cardiologia. Publicado por Elsevier España, S.L. Todos os direitos reservados.

Introduction

Carcinoid tumors, with an incidence in the general population of 1/100 000 individuals, are rare malignant neoplasms that arise from enterochromaffin cells located in the gastrointestinal tract or lungs.^{1,2}

At the time of diagnosis, 20–30% of patients have disseminated disease and present with carcinoid syndrome, characterized by flushing, diarrhea and bronchospasm. This syndrome is caused by tumor release of serotonin and other vasoactive substances.³

Approximately 50–60% of patients with the syndrome present carcinoid heart disease, which is caused by vasoactive substances secreted by metastatic tumor cells in the liver reaching the right heart, leading to deposition of fibrous tissue in the endocardium. Studies suggest that the prevalence of carcinoid heart disease is decreasing following the introduction of somatostatin analogs and other antitumor therapies designed to reduce the tumor load and the production of tumor secretory products.^{4,5}

Carcinoid heart disease is initially well tolerated, and patients may remain in New York Heart Association (NYHA) functional class I, even with severe valvular lesions in the right heart. However, typical signs and symptoms of right heart failure appear as the disease progresses.

The echocardiographic features are well defined: classically the tricuspid and pulmonary subvalvular apparatus and valve leaflets are thickened and fixed, limiting leaflet excursion and preventing coaptation, which can cause a combination of valve regurgitation and stenosis. In most patients with carcinoid heart disease, the tricuspid valve is affected, with or without pulmonary valve involvement. Around 15% of patients have valve lesions in the left heart, also characterized by diffuse leaflet thickening, although less severe than in the right heart. Mild pericardial effusion occurs in 10% of patients, but cardiac metastases are rare. 4

We present the case of a female patient with carcinoid syndrome in whom transthoracic two- and three-dimensional echocardiography provided sufficiently accurate assessment of valve structures to establish a diagnosis of carcinoid heart disease without the need for transesophageal echocardiography.

Case report

We describe the case of a 76-year-old woman, with no relevant personal history, diagnosed with a neuroendocrine tumor of the cecum with an insular pattern in January 2004 following right hemicolectomy, and followed since then in medical oncology consultations.

Abdominal computed tomography and ¹¹¹Inpentetreotide scintigraphy (octreoscan) revealed metastasization to the liver, and palliative treatment was begun with five million units (MU) interferon-alfa three times a week. In April 2004 the patient underwent excision of five liver metastases, bilateral oophorectomy and excision of a peritoneal nodule. Histological study of all fragments revealed carcinoid tumor, and so interferon-alfa therapy was maintained.

In March 2005, on repeat octreoscan no liver lesions were observed, and the patient continued to be monitored. In January 2007, although clinically asymptomatic, she presented liver uptake on octreoscan and elevated 5-hydroxy-indolacetic acid, an indication for interferon or chemotherapy, but the patient refused any treatment.

In September 2010, she presented sporadic episodes of flushing, diarrhea and bronchospasm and octreotide therapy was begun for carcinoid syndrome, but despite slight symptomatic improvement, the therapy was discontinued in December 2010 due to intolerance (nausea, vomiting and abdominal pain).

In January 2011, the patient complained of fatigue on progressively less exertion and showed disease progression with worsening liver lesions; oral chemotherapy was begun with capecitabine, but was discontinued after four cycles due to intolerance (nausea, vomiting and diarrhea). She still had symptoms of carcinoid syndrome and worsening secondary lesions, but refused directed therapy.

In June 2011, the patient began to present lower limb edema, increased abdominal volume and worsening fatigue, and was admitted in September 2011 with a provisional diagnosis of decompensated heart failure.

Physical examination showed pale and dry mucous membranes, liver palpable 4cm below the right costal margin, positive fluid wave sign and lower limb edema up to the hip joint.

Diagnostic exams included an electrocardiogram on admission (Figure 1), which showed sinus rhythm, heart rate of 70 bpm, and low QRS voltage in all leads.

The admission chest X-ray revealed obliteration of both costophrenic angles, compatible with bilateral pleural effusion

Transthoracic two- and three-dimensional echocardiography (Figures 2 and 3) showed normal left ventricular dimensions and wall thickness, with no wall motion abnormalities and preserved global systolic function. The right chambers were enlarged (Figure 2A), with good right ventricular global systolic function (tricuspid annular plane systolic excursion: 22 mm), with no significant gradient between the right ventricle and right atrium (Figure 2D). The tricuspid valve leaflets were visibly thickened, preventing coaptation in systole and causing severe tricuspid regurgitation (Figure 2B), which was more evident in three-dimensional images (Figure 2C1 and C2).

The pulmonary valve cusps were thickened, causing mild pulmonary stenosis (Figure 3C) and regurgitation (Figure 3B).

Download English Version:

https://daneshyari.com/en/article/3020605

Download Persian Version:

https://daneshyari.com/article/3020605

<u>Daneshyari.com</u>