Carcinoid heart disease from ovarian primary presenting with acute pericarditis and biventricular failure

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Abstract

A case is described of a 54 year old woman who had acute pericarditis with large exudative effusion accompanied by severe right and left ventricular failure. The patient was finally diagnosed with carcinoid heart disease from an ovarian carcinoid teratoma. She was treated with octreotide—a somatostatin analogue—followed by radical surgical resection of the neoplasm. At one year follow up only mild carcinoid tricuspid regurgitation remained. Only 16 cases of carcinoid heart disease from an ovarian primary have been described in literature. Moreover clinically manifest acute, non-metastatic pericarditis and left heart failure are not considered as possible presentations of carcinoid heart disease, whatever the origin. In a recent series a small pericardial effusion was considered an infrequent and unexpected echocardiographic finding in carcinoid heart patients. One case of “carcinoid pericarditis” has previously been described as a consequence of pericardial metastasis. Left sided heart involvement is usually caused by bronchial carcinoids or patency of foramen ovale; both were excluded in the case presented. (Heart 1998; 80: 623–626)

Keywords: carcinoid heart disease; ovarian tumour; acute pericarditis; heart failure

Primary ovarian carcinoid tumours are rare, accounting for 0.3% of all carcinoid tumours.1 Carcinoid syndrome and carcinoid heart disease are believed to complicate fewer than 10% of these rare ovarian neoplasms.2 Only 16 cases of carcinoid heart disease from an ovarian primary are described in the literature.3–4 Such tumours are usually diagnosed following detection of a pelvic mass, carcinoid syndrome (diarrhoea, flushing), or typical carcinoid heart disease (right sided valvar involvement). We describe a case in which an ovarian carcinoid teratoma was found in a patient who presented with acute pericarditis and biventricular failure. To our knowledge this is the first report of acute, non-metastatic, “myopericarditis” as the first clinical manifestation of carcinoid disease.

Case report

A 54 year old woman was admitted to our division complaining of a dry, irritating cough and slight evening fever for 15 days and more recent mild dyspnoea and dull chest pain. She was in physiological menopause and a check up three years ago had excluded cardiovascular disease. A mild, grade 2/6 holosystolic murmur could be heard on the left lower sternal border and pericardial friction rubs persisted for a few days after admission. The liver was palpable three fingerbreadths below the right costal margin. Prominent jugular veins, abdominal–jugular reflux, and mild bilateral ankle oedema could be appreciated. ECG showed non-specific ST-T changes. Raised serum lactic dehydrogenase and aminotransferase were found on admission; creatine kinase was repeatedly normal. Chest radiography showed pronounced enlargement of the cardiac silhouette and normal lungs. Echocardiography showed a large pericardial effusion without signs of cardiac tamponade (fig 1), moderate to severe tricuspid regurgitation with right atrial enlargement, and reduced left ventricular systolic function.

Figure 1 Echocardiogram on first admission. A large pericardial effusion is seen around the cardiac chambers with a “swinging heart” appearance.
Fully evacuative pericardiocentesis produced 250 ml of serous, haematic exudate negative for neoplastic cells and mycobacterium. A persistent butterfly-shaped erythematous malar rash had been present since admission, therefore systemic lupus erythematosus was initially suspected, but laboratory tests and a search for specific autoantibodies proved negative. Pericardiocentesis and medical treatment resolved the symptoms. The patient was diagnosed with acute exudative pericarditis and discharged on maintenance treatment (diuretics, aspirin, antibiotic) and close follow up.

Unexpectedly she returned 11 days later because of worsening dyspnoea, peripheral oedema, cough, and recurrent evening fever. She was admitted to the intensive care unit owing to critical congestive heart failure. ECG showed worsening of ST-T changes consistent with ventricular systolic overload (fig 2A). A left posterobasal pleural effusion was evident on chest x rays. Echocardiography showed no pericardial effusion but indicated severely reduced left ventricular systolic function (ejection fraction 26%). Right heart Swan-Ganz monitoring showed raised central venous (34 mm Hg) and pulmonary wedge pressures (36 mm Hg) and a reduced cardiac index (2.5 litres/min). Intravenous diuretics and nitrates, and oral angiotensin converting enzyme inhibitors led to rapid improvement, and the patient was moved from intensive care after a few days.

The patient had no history of intravenous drug abuse. Blood cultures were repeatedly negative. No endocardial vegetations were evident using transoesophageal echocardiography, but we could distinctly see diffuse thickening, shortening, and stiffening of the tricuspid leaflets (fig 3) without commissural

Figure 2  (A) ECG on admission to the intensive care unit for severe congestive heart failure. Severe ST-T alterations consistent with ventricular systolic overload are evident. (B) At one year follow up there is complete normality of the tracing, except for first degree atrioventricular block.

Figure 3 (A) Tricuspid valve aspect on transoesophageal echocardiography. (B) Typical dagger-shaped regurgitant signal of the tricuspid valve. Both suggest carcinoid heart disease.
Carcinoid heart disease from ovarian primary

Discussion

Up to 90% of carcinoids arise from the small bowel and the appendix. The remainder usually originate from other gastrointestinal sites or the bronchi. Ovary, thymus, and breast are very rare sites of origin. Carcinoid syndrome and carcinoid heart disease are caused by the offending action of many humoral substances produced by the tumour, usually inactivated by liver and lung enzymes. Gastrointestinal carcinoids tributary to portal vein circulation therefore give rise to carcinoid syndrome only when liver metastases secrete downstream from the hepatic filter. Ovarian carcinoids, on the contrary, release their products directly into the caval system and may give rise to carcinoid syndrome and carcinoid heart disease before the development of any metastatic lesion. Surgical removal can therefore be a radical treatment, and prevent the progression of heart lesions. For this reason, prognosis of carcinoid heart disease from ovarian carcinoid tumours was considered good since the very first reports. Nevertheless, other authors subsequently regarded the prognosis uncertain in the same cases and reported the possibility of progression of heart lesions despite surgical removal of the tumour. In such cases, cardiac surgery is indicated and can be life-saving.

In our case, the patient presented with a myopericarditis-like clinical picture—not a typical presentation of carcinoid disease. The diagnosis was suspected on the basis of the tricuspid valve’s peculiar aspect at ultrasound. Transoesophageal echocardiography was useful in magnifying this aspect, thus confirming the importance of both transthoracic and transoesophageal ultrasounds in diagnosing carcinoid heart disease as already mentioned in literature.

Pericarditis as a presentation of carcinoid heart disease was described in 1973 in one case of pericardial metastasis from a bronchial carcinoid tumour, but the pericardial effusion in that case was clearly related to a direct involvement from the contiguous neoplasm. A small pericardial effusion has been reported as a possible echocardiographic finding in carcinoid heart disease, and was present in 14% of cases in the largest recent series. Nevertheless, pericardial effusion remains an uncommon aspect of the disease and is an unexpected finding at echocardiography.

In contrast, we observed definite signs and symptoms of pericardial involvement as an initial manifestation of the disease, so that the diagnosis of acute pericarditis of unknown origin was initially formulated. Origin of the heart failure was also unclear. Right heart failure could have been related to the tricuspid valve disease whereas left heart failure had no evident causes. Moreover, left sided heart involvement in carcinoid heart disease is present in only 7% of patients and occurs in bronchial carcinoids or when a patent foramen ovale coexists, or perhaps with very prolonged and severe disease (a unique case requiring quadruple valve replacement has been described). Reduced left ventricular systolic
function is described in fewer than 4% of the largest published series.\(^5\)

We could therefore clinically suspect an accompanying or complicating myocarditis of unknown origin. The heart failure could otherwise be related directly to the carcinoid heart disease as hypothetical “toxic” damage by substances produced by the tumour. In this regard, an endomyocardial biopsy could have been interesting, but was deemed of no practical benefit in management and was not performed.

Finally, we emphasise the efficacy of octreotide in resolving all major symptoms of the carcinoid syndrome in our case, and we believe that the uncomplicated intraoperative course that ensued was largely the result of pretreatment with this long acting somatostatin analogue.\(^7\)

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