Constrictive Pericarditis – A Rare but Important Cause of Cardiac Failure: A Case Report and Review of Literature.


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Constrictive pericarditis (CP) is an uncommon cardiac disease which is often difficult to diagnose because of its vague, various and nonspecific clinical presentations. An awareness and understanding of this condition is very important, as it is a progressive condition and the likelihood of cure depends very much on its early identification and treatment.

In our report, we presented a middle-aged female patient had non-specific symptoms and signs for two years, observing echocardiographic and hemodynamic findings of constrictive pericarditis, she was eventually diagnosed to have idiopathic constrictive pericarditis. In this report etiologies, classical clinical history and physical examination findings of constrictive pericarditis were described. Radiologic, electrocardiographic, echocardiographic and angiographic findings are discussed and the hemodynamics of CP were reviewed.

**Keywords:** Congestive cardiac failure, constrictive pericarditis; echocardiography; pericardiectomy

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Özetté: Konstruktif perikardit (KP) klinik bulguların belirsiz, çeşitli ve özgül olmadığınından tanısı güçlükle konulabilen seyrek bir kalp hastalığıdır. KP ilerleyici bir hastalık olup erken tanı konulup müdahalede edildiğinde tedavi olası çok yüksek olduğundan hastalığın farkındalığı ve durumun anlaşılması önemlidir. Biz yazımızda iki yıl spesifik olmayan semptom ve bulgulara sahip, KP’ın ekokardiyografik ve hemodinamik bulgularının izlendiği ve sonunda idiopatik konstruktif perikardit tanı koyduğuımız orta yaşlı kadın hastanın sunduk. Bu yazında konstruktif perikarditin etyolojisi, klasik klinik öykü ve fizik muayene bulgularına degradirdik. Radyolojik; elektrokardiyografik, ekokardiyografik, anjiyografik bulgulan tarsıktık ve hemodinamik bulgulan gözden geçirdik.

Anahtar Kelimeler: konjestif kalp yetmezliği, konstruktif perikardit, ekokardiyografi, perikardiektomi
**Giriş**

There are a few truly curable cardiac diseases. One of these is constrictive pericarditis (CP), a rare cardiac condition that usually manifests itself with nonspecific symptoms and signs spread out over many months or even years. Furthermore, clinical signs and symptoms of right heart failure accompanied by existing risk factors for pericardial disease should raise suspicion for constrictive pericarditis. The diagnosis of constrictive pericarditis remains a challenge because its physical findings and hemodynamics mimic other disease. A better appreciation of this disease is important, if one is to suspect its presence in a clinical scenario, which is an essential step that may lead to its diagnosis.1-3

**Case Report**

A 34-year-old female presented with worsening dyspnea on exertion, fatigue and palpitation for two years. On examination, her blood pressure and pulse were 100/60 mmHg, 89 bpm respectively. She hadn’t jugular venous distention, ascites, an enlarged liver and pitting edema. On cardiac auscultation, there was a regular rhythm without any murmur and pericardial knock. She hadn’t history of tuberculosis, surgery, trauma, irradiation or other disease. Investigations looking for a possible cause of CP, such as viral markers, rheumatoid factor and antinuclear antibodies, were either negative or inconclusive. Electrocardiography showed sinus rhythm with non-specific diffuse T wave inversions. Chest X-ray did not show any sign of previous pulmonary tuberculosis, although the left lateral telecardiogram revealed thick intense calcification of the pericardium enclosing the heart (Fig. 1). Echocardiography showed mildly dilated left atrium with normal-sized left ventricle and particular thickening of the pericardium in the neighborhood of posterior left ventricular wall (Fig. 2a). A septal bounce (inspiratory septal shift to the left) was readily visualized in the apical four-chamber view. There was an increased respiratory variation of the early diastolic wave through the mitral valve (>25%) (Fig. 2b). The inferior vena cava was dilated (24.7 mm) and noncollapsing in the subcostal views (Fig. 2c). Tricuspid regurgitation was trivial with a pulmonary artery pressure of 35 mmHg. Tissue Doppler echocardiography showed an early diastolic mitral annular velocity E’of 14.7 cm/sec at the septal mitral annulus. Computed tomography of the thorax showed diffuse, incomplete calcification of the pericardium with a thickness ranging from 6 mm (Fig. 3a-b). Coronary angiography showed normal coronary arteries. However, pericardial calcification was seen near the apex on fluoroscopy. Simultaneous left and right heart catheterisations were performed and demonstrated elevation and equalisation of the diastolic pressures of all cardiac chambers to within 5 mmHg. The mean right atrial pressure was 18 mmHg with a prominent Y descent, right ventricular end diastolic pressure (RVEDP) 18 mmHg, left ventricular end diastolic pressure (LVEDP) 17 mmHg and pulmonary capillary wedge pressure 20 mmHg. The ventricular tracing showed a dip-and-plateau pattern (“square root sign”). The diagnosis was calcific CP. The council of cardiology and cardiovascular surgery decided to close clinical follow-up of patients, because of there is no serious symptoms of heart failure yet.

![Figure 1: Lateral chest x-ray demonstrates pericardial calcification](image1)

![Figure 2a: 2-D Echocardiography showed particular thickening of the pericardium on the posterior of left ventricle, 2b: Pulsed-wave Doppler at the mitral valve shows increased respiratory variation and decreased deceleration time of the E](image2)
wave. 2c: The inferior vena cava was dilated and noncollapsing in the subcostal views.

Figure 3a-b: Chest computed tomography scan shows generalized incomplete thickening of the pericardium

Discussion
CP is caused by adhesions between the visceral and parietal layers of the pericardium and progressive pericardial fibrosis that restricts diastolic filling of the heart which results in fluid overload or diminished cardiac output in response to exertion. The diagnosis of CP remains a challenge because its physical findings and hemodynamics mimic other disease. CP should be suspected in patients with clinical features of right-sided heart failure. A high level of suspicion is required to confirm the diagnosis. 1

In the past, the commonest aetiology was tuberculosis, and this remains true in developing countries, but not for developed nations. In a recent study, idiopathic CP is a common cardiac disease and the three most common identifiable causes were cardiac surgery (18%), pericarditis (16%) and mediastinal irradiation (13%). Other possible causes include connective tissue diseases, malignancy, trauma and infections. Other cardiac diseases, in particular right atrial myxoma, tricuspid valve dysfunction, and restrictive cardiomyopathy must be ruled out. 2

Presentations are often insidious and non-specific, such as reduced effort tolerance, orthopnoea or fatigue. In a recent European study, the average duration of symptoms before definitive diagnosis was 20 months. The most important clinical finding is that of a raised jugular venous pressure with elevation on inspiration (Kussmaul’s sign), though this in itself is not specific for CP, as it can occur in any condition with elevated right sided pressures. There may be a diastolic pericardial knock due to the sudden cessation of ventricular filling, which may be heard as a “third heart sound”. Other clinical findings such as pulmonary congestion, hepatomegaly, ascites and peripheral oedema are a reflection of the elevated diastolic pressures causing heart failure. Our patient was admitted to our clinic with presented us dyspnea on exertion, fatigue and palpitation without congestive cardiac failure sign and did not have a particular etiology for pericardial constriction. 3

The diagnosis of CP can be elusive and initial clinical suspicion often helps. Unfortunately, there is no single diagnostic test. Electrocardiography is almost always abnormal and often shows nonspecific low QRS voltages with generalised T wave inversion or flattening. Atrial fibrillation occurs in less than 50% of patients. Chest X-ray occasionally shows some pericardial calcification and though it may suggest CP, it is not diagnostic in itself, as stated by Lorrell, “Calcified pericardium is not necessarily a constricted one.” 4

Among the available diagnostic tests, echocardiography is a class I indication in patients with suspected pericardial disease. 5 Computed tomography (CT), magnetic resonance imaging (MRI) or cardiac catheterisation may require for diagnosing. However, pericardial thickening detected on CT or MRI is not
diagnostic for CP and, on the other hand, normal pericardial thickness does not rule out the diagnosis of CP.6 Talreja et al. found that the normal thickness of the pericardium in 18% of patients with CP.7 Normal pericardial thickness is 2 mm or less. Pericardial thickness > 4 mm is suggestive of CP and > 6mm specificity is very high.8 In addition, the diagnosis of pericardial thickening is not enough by itself to make the diagnosis.9

On echocardiography, actual pericardial thickening is often not seen, but the inferior vena cava is often dilated. Increased pericardial thickness on echocardiography have been reported 95% of the sensitivity and specificity of 86% for CP.10 On M-mode echocardiography, diastolic “septal bounce” may be seen, as well as abrupt checking of the left ventricular posterior wall diastolic movement by the rigid pericardium (coinciding with the “pericardial knock”).11 Another suggestive sign is early opening of the pulmonary valve in late diastole, due to an elevated RVEDP. Echocardiographic detection of excessive respiratory flow changes through the mitral valves, such as an inspiratory decrease of more than 25%, right ventricular systolic pressure less than 50 mmHg, and an early diastolic mitral annular velocity (E’) of ≥8 cm/sec by tissue Doppler echocardiography enabled us to diagnosis CP.6,10

Invasive hemodynamic evaluation may be necessary in some patients for the diagnosis.12 Simultaneous recording of left and right heart pressures is helpful in documenting the presence of a constrictive physiology. The characteristic finding is elevation and equilibration of diastolic filling pressure in all cardiac chambers to within 5 mmHg. There is often a “dip and plateau” or “square root sign” for both right and left ventricle diastolic pressure curves, with a low early diastolic pressure and rapid rise to a high plateau, though this sign is not specific in itself. It is important to avoid prior diuresis before this test, as the characteristic findings may be obscured in the presence of hypovolaemia. Sometimes a rapid volume infusion to increase intravascular volume can help to unmask the characteristic findings.13

Finally, Chronic CP is a progressive disease. Some patients may survive for many years controlled with diuretic therapy, but the majority will become increasingly symptomatic and disabled. The treatment of constrictive pericarditis consists of diuretics, salt restriction, and supportive therapy for the underlying condition. Surgical pericardiectomy, which should be as complete as possible, is the definitive therapy which offers the possibility of a cure, with a low in-hospital mortality rate if performed in patients with good condition and without advanced disease. Higher pressures (central venous pressure of 12 mmHg to 15 mmHg) and liver dysfunction secondary to passive congestion are indications for urgent surgery.14 Predictors of poor outcome include radiation related constrictive pericarditis, New York Heart Association class III or IV at presentation, myocardial atrophy, inflammation or scarring.15,16 Survival of the patients following pericardiectomy is better than that of individuals treated without surgery.17
References