Unexplained extensive calcification of the venae cavae extending into the right atrium causing partial obstruction of the tricuspid valve

Hashir Kareem,¹ Tom Devasia,¹ Krishnananda Nayak,² Sumit Agarwal¹

DESCRIPTION
A 23-year-old woman, who had been diagnosed recently with hypothyroidism, was referred to us for an echocardiogram. She was apparently asymptomatic when she developed fatigue and mild generalised oedema. She did not have any history of chest pain, breathlessness, palpitation or syncope. Her echocardiogram revealed the following very unusual finding (figure 1, video 1). A large, irregular, highly echogenic mass was seen in the right atrium (RA), partially prolapsing through the tricuspid valve (TV) in diastole. There was mild obstruction of the TV as evidenced by increased velocity gradients across the valve. The mass seemed to be in continuity with a highly echogenic structure lining the inferior vena cava (IVC) (figure 2). Rest of the cardiac structures were normal. There was no cardiac enlargement or pericardial effusion. No evidence of pulmonary hypertension was seen. There was no history suggestive of pulmonary embolism. She underwent a CT scan which revealed extensive calcification of the IVC extending from the level of the renal veins up to the RA and also extending into the superior vena cava (SVC) (figure 3). The CT scan showed clearly that the right atrial structure was in direct continuity with the calcification in the IVC (figure 3). Moreover, the density of the structure on CT was compatible with calcium, further proving that the structure was indeed an extension of the calcification in the IVC. In addition to this, there was a small thrombus in the RA which was not calcified. There was no evidence of deep vein thrombosis of the leg. She was investigated for any evidence of thrombophilic conditions since thrombus calcification is a known cause for this type of presentation.¹ Proteins C and S were normal, so were antithrombin levels. There was no evidence of antiphospholipid antibody syndrome (APLAS). Serum homocysteine was normal. Screening for connective tissue diseases was also normal. There was no evidence of parathyroid abnormality. Her serum calcium and phosphate levels were normal. Renal function was also normal.

Figure 1 (A and B) Four-chamber view of echocardiogram showing the calcific structure (black arrows) in the right atrium partially prolapsing through the tricuspid valve (white arrow) into the right ventricle (RV) during diastole (B) (LV, left ventricle).

Figure 1

To cite: Kareem H, Devasia T, Nayak K, et al. BMJ Case Rep Published online: [please include Day Month Year] doi:10.1136/bcr-2014-204070

Images in...

CrossMark

BMJ Case Reports: first published as 10.1136/bcr-2014-204070 on 5 June 2014. Downloaded from http://casereports.bmj.com/ on 16 September 2023 by guest. Protected by copyright.
IVC calcification has been described in neonates and is associated with disseminated intravascular coagulation, placentofetal embolus, hypotensive shock, dehydration, focal infection, septicaemia and structural anomalies.\(^1\) It may also be associated with malignant disease. However, the aetiology in most cases remains unclear.\(^1\) It is extremely rare in adults and only a few cases have been reported in literature. Most patients are asymptomatic and the calcification is usually detected incidentally on X-ray or CT. Chetwood et al described a patient with IVC calcification who presented with recurrent pulmonary embolism.\(^2\) IVC calcification has been described in association with APLAS.\(^3\)

The most extraordinary aspect of this case is the extension of calcification into the RA and partial prolapse of the structure through the TV. It is possible that the calcification may have extended to the RA along a pre-existing abnormal Eustachian valve. However, this is just conjecture based on the echocardiographic appearance and the exact mechanism remains a mystery. We believe that such extensive IVC calcification with extension into the RA, TV and SVC in an adult patient has not been reported before.

The patient was started on thyroid hormone supplementation and oral anticoagulation in view of the risk for pulmonary embolism. Surgical intervention was considered. However, in view of the extensive calcification, it was deemed too risky. She is currently on regular follow-up.

**Learning points**

- Inferior vena caval (IVC) calcification is usually found in neonates and children and is extremely rare in adults.
- IVC calcification is usually benign and detected incidentally but may occasionally be associated with pulmonary embolism.

**Competing interests** None.

**Patient consent** Obtained.

**Provenance and peer review** Not commissioned; externally peer reviewed.

**REFERENCES**


**Figure 2** Subcostal view of echocardiogram showing the calcific structure in the inferior vena cava (black arrows) that is continuous with the structure in the right atrium (white arrows).

**Figure 3** (A) CT scan—coronal section showing extensive calcification of the inferior vena cava (IVC) (black arrows) starting just above the renal veins with calcification of the superior vena cava (white arrow); (B) sagittal view showing the calcific structure extending from the IVC in to the right atrium (black arrows).