Cystic tumor of the atrioventricular node in a patient with intermittent complete heart block

Vineeta Ojha,1, Niraj Nirmal Pandey,1 Gautam Sharma,2 Priya Jagia1

DESCRIPTION

A 53-year-old man presented with occasional shortness of breath and palpitations for the last 6 months. Holter monitoring revealed intermittent complete heart block and junctional escape rhythm. Transthoracic echocardiography was unremarkable. Cardiac MR was performed to look for any scar or any other structural cause for palpitations. It revealed a 1.2×1 cm nodular lesion at the anterior mitral leaflet-interatrial septal junction. This lesion appeared iso intense on T1W images, hyperintense on T2W images and showed peripheral rim enhancement with a central dark core on late gadolinium enhancement (LGE) images (figure 1A–C). T2 mapping values were also raised (65 ms). On high T1 (Inversion Time) LGE images, the lesion was relatively bright. Subsequently, CT angiography was also performed for the assessment of coronaries and to look for any calcification within the lesion. CT demonstrated the lesion with an attenuation of ~70 HU (Hounsfield Units) on the non-contrast CT with no significant enhancement observed in the postcontrast scans (figure 1D–F). Coronary arteries were normal.

On the basis of morphological characteristics, the mass was presumed to be benign. Considering the unusual location of the mass, differential diagnosis included benign lesions like hamartoma, fibroma, cystic tumour of the atrioventricular node (CTAVN) and bronchogenic cyst. However, considering the signal characteristics and the location of the mass as well as patient’s symptoms, a likely diagnosis of CTAVN was made. Excision of the lesion was suggested; however, the patient did not opt for the same. Subsequently, pacemaker implantation was done and the patient is on the routine follow-up.

CTAVNs are extremely rare congenital lesions located at the base of the interatrial septum.1 Because of their location, these masses often cause complete heart block and arrhythmias and may result in sudden cardiac death. These tumours have mostly been diagnosed post-mortem with only a few antemortem reports in the literature.2 CTAVNs are generally small and the patients may remain asymptomatic or may present with palpitations, dizziness and dyspnoea. They may have variable signal intensity on CT as well as MRI depending on the composition of the microcysts and the amount of fibrous component. Fiset et al described a similar case of a 51-year-old woman with third-degree atrioventricular nodal block and junctional rhythm. Echocardiography revealed an echogenic mass at the same location as in our case, which showed a soft-tissue attenuation (63 HU) on CT. Although signal intensity on the non-contrast MRI was similar to our case, the lesion showed intense contrast enhancement.3 However, absence of postcontrast enhancement has also been reported in other similar cases.4 Although rare, a mass at this unusual location (at the base of the interatrial septum) should alert the radiologists to the possibility of cystic tumour of the atrioventricular node.

Learning points

► A mass at the base of the interatrial septum should alert the clinicians and radiologists to the possibility of cystic tumour of the atrioventricular node.
► Because of their location, these masses often cause complete heart block and arrhythmias and may result in sudden cardiac death.
for a definitive diagnosis of this lesion, which is not present in the current case.

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ORCID ID Vineeta Ojha http://orcid.org/0000-0003-4371-5615

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