Right ventricular thrombi – suspicion of arrhythmogenic right ventricular dysplasia: a case report

Livija Sušić1
Vedrana Baraban2
Josip Vincelj3
Jasmina Ćatić3
Robert Blažeković3
1Health Centre Osijek, Osijek, Croatia
2Josip Juraj Strossmayer University of Osijek School of Medicine, University Hospital Centre Osijek, Osijek, Croatia
3Dubrava University Hospital, Zagreb, Croatia

KEYWORDS: dyspnea, syncope, cardiac arrhythmia, thrombus, arrhythmogenic right ventricular cardiomyopathy.

CITATION: Cardiol Croat. 2015;10(3-4):81-82. DOI: http://dx.doi.org/10.15836/ccar.2015.81


*ADDRESS FOR CORRESPONDENCE: Livija Sušić, Dom zdravlja Osijek, Park kralja Petra Krešimira IV/6, HR-31000 Osijek, Croatia. Phone: +385-31-225-304. E-mail: livija.susic@gmail.com

CASE REPORT: 61-year-old woman with long history of hypertension presented with progressive dyspnea and chest pain that she has been experiencing at minor physical exertion for several months. Current medical records confirmed T-wave inversion in right precordial leads, attacks of supraventricular and ventricular arrhythmia, including attacks of non-sustained ventricular tachycardia and recurrent syncopal episodes from the age of 23. Dilated right heart chamber are detected by transthoracic echocardiogram year 2006, presence of the shunt was excluded by scintigraphy.

IMAGING STUDIES: 2-dimensional transthoracic echocardiogram revealed one large (Figure 1) and two smaller hyperechogenic masses (Figure 2) inside extremely dilated (Figure 3) and globally hypokinetic right ventricle with moderate tricuspid regurgitation. 3-dimensional transesophageal echocardiogram confirmed mobile pedunculated mass by free wall of right ventricle (Figure 4). Magnetic resonance imaging describe dilated right heart chambers, hypertrophic septomarginal trabeculae of the mid-apical part of right ventricular and mass within the same chamber appears to be a tumour. CT pulmonary angiogram, PET CT and coronarography ruled out specific pathomorphological substrate.

TREATMENT: Three large individual masses were surgically removed from the chamber of right ventricle (Figure 5). Annuloplasty of the tricuspid valve was performed.

PATHOHISTOLOGICAL DIAGNOSIS: focal-organizing thrombi.
CONCLUSION: Although final diagnosis has not been made, it is possible that the patient suffers from arrhythmogenic right ventricular dysplasia. This conclusion is based on the presence of three big McKenna's criteria1 (T-wave inversion in right precordial leads, ECHO-PLAX RVOT >32 mm and non-sustained ventricular tachycardia detected during Holter monitoring) and the fact that literary sources describe several cases of thrombotic masses in right heart chambers in patients with the same diagnosis.2,3

LITERATURE


Right ventricular thrombi – suspicion of arrhythmogenic right ventricular dysplasia: a case report