Opis choroby *

Definicja

A rare congenital heart malformation of unknown etiology that is characterized by an extremely dilated right atrium, and that is usually asymptomatic and fortuitously discovered by echocardiography or chest radiography, and can be sometimes associated with other anomalies such as atrial arrhythmias (e.g. atrial flutter, atrial fibrillation, supraventricular tachycardia), severe tricuspid regurgitation, or atrial thrombus that could lead to potentially life-threatening thromboembolic complications.

Dane

Klasyfikacja
Wada morfologiczna

Kod ORPHA
1677

Kod OMIM
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Kod ICD10
Q20.8

Kod ICD11
LA8F

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*Źródło

orphanet