

Opis choroby *

Definicja

A rare, congenital, non-syndromic, heart malformation characterized by the persistence of the embryonic right valve of the sinus venosus which results in a subdivision of right atrium into two chambers. Clinical manifestations depend on the degree of right atrial septation and the size of sinoatrial orifice and vary from asymptomatic to symptoms of tricuspid valve stenosis, atrial fibrillation, cyanosis, syncope, elevated central venous pressure and right heart failure. The anomaly may be isolated or associated with other congenital heart anomalies.

Dane

Klasyfikacja	Synonimy	
Wada morfologiczna	Cor triatriatum dextrum Prawy przedsionek podzielony Divided right atrium	
Kod ORPHA 99098	Kod OMIM -	Kod ICD10 Q24.2
Kod ICD11 LA8F		

*Źródło

orphanet