

Opis choroby *

Definicja

A rare, congenital, non-syndromic heart malformation characterized by partial or complete absence of tricuspid valve tissue and its apparatus, with an existing orifice. It can be isolated or associated with other heart anomalies. Clinical presentation is variable and may include syncope, arrhythmias, cyanosis, right heart dilatation and failure.

Dane

Klasyfikacja	Synonimy
Wada morfologiczna	Congenital unguarded tricuspid orifice Wrodzony brak zastawki trójdzielnej

Kod ORPHA	Kod OMIM	Kod ICD10
95457	-	Q22.4

Kod ICD11
LA87.0Y

*Źródło

orphanet