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
Wada morfologiczna

Synonimy
Congenital unguarded tricuspid orifice
Wrodzony brak zastawki trójdzielnej

Kod ORPHA
95457

Kod OMIM
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Kod ICD10
Q22.4

Kod ICD11
LA87.0Y

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*Źródło

orphanet