

## 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