

## Opis choroby \*

### Definicja

A rare congenital non-syndromic heart malformation characterized by a communication between the ascending aorta and the pulmonary trunk in the presence of two normally formed semilunar valves. It may be an isolated finding or occur in association with other anomalies. Severe clinical manifestations, such as congestive heart failure or pulmonary hypertension, typically develop in early life.

### Dane

#### Klasyfikacja

Wada morfologiczna

#### Synonimy

Congenital aortopulmonary artery fistula  
Okienko aortalne  
Wrodzona przetoka tętnicza aortalno-płucna  
Wrodzony ubytek przegrody aortalno-płucnej  
Congenital aortopulmonary septal defect

#### Kod ORPHA

2037

#### Kod OMIM

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#### Kod ICD10

Q21.4

#### Kod ICD11

LA8B.0

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#### \*Źródło

orphanet