

Opis choroby *

Definicja

A rare, life-threatening, congenital, non-syndromic, conotruncal heart malformation disease characterized by absent or severely undeveloped pulmonary valve leaflets (with a restrictive ring of thickened tissue at the place of the pulmonary valve annulus), associated with an intact ventricular septum and a patent ductus arteriosus, manifesting with marked respiratory insufficiency. Additional features include dilated main pulmonary artery (with or without dilatation of pulmonary artery branches), to-and-fro flow at site of the dysplastic pulmonary valve, and systolic pressure gradient across narrowed pulmonary valve. Tricuspid atresia and variable extra-cardiac anomalies (e.g. diaphragmatic hernia or cleft lip/palate), may be present.

Dane

Klasyfikacja

Zespół wad wrodzonych APV/PDA, non-Fallot type

Brak zastawki płucnej - ubytek przegrody

międzykomorowej - przetrwały przewód tętniczy

APV/PDA, non-Fallot type

PVA/PDA, non-Fallot type

Kod ORPHA

99048

Kod OMIM

-

Kod ICD10

Q22.2

Kod ICD11

-

*Źródło

orphanet