Opis choroby *

Definicja

A rare congenital non-syndromic heart malformation characterized by an imperforate or absent mitral valve. In most cases, there is a univentricular atrioventricular connection to a dominant right ventricle via a tricuspid valve, and a hypoplastic left ventricle. Morphologic heterogeneity is considerable, and hemodynamic picture and clinical manifestation depend on the type and severity of associated cardiovascular anomalies (such as ventricular septal defect or aortic atresia).

Dane

Klasyfikacja
Wada morfologiczna

Kod ORPHA
1205

Kod OMIM
-

Kod ICD10
Q23.2

Kod ICD11
LA89.2

---

*Źródło

orphanet