Tricuspid valve agenesis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Tricuspid valve agenesis. ORPHA:95457

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.