Opis choroby *

Definicja

An extremely rare genetic congenital heart disease characterized by the presence of atrial septal defect, mostly of the ostium secundum type, associated with conduction anomalies like atrioventricular block, atrial fibrillation or right bundle branch block.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
1479

Kod OMIM
108900

Kod ICD10
Q21.1

Kod ICD11
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*Źródło

orphanet