

## 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