

## Opis choroby \*

### Definicja

A rare congenital anomaly of the coronary sinus characterized by its stenosis at the ostium, lumen, or origin, typically leading to dilation of the vessel. Symptoms are variable and can include palpitations, tachypnea, dyspnea, chest pain, fatigue, and cyanosis. The malformation may be associated with other cardiac anomalies, such as coronary artery-coronary sinus fistula, unroofed coronary sinus, atrial septal defect, coronary sinus-left atrium fistula, total anomalous pulmonary venous connection, and ventricular septal defect.

### Dane

### Klasyfikacja

Wada morfologiczna

#### Kod ORPHA

99117

#### Kod OMIM

-

#### Kod ICD10

Q21.1

#### Kod ICD11

LA86.Y

---

### \*Źródło

orphanet