

## Opis choroby \*

### Definicja

A rare familial cardiomyopathy characterized by left ventricular enlargement and/or reduced systolic function preceded or accompanied by significant conduction system disease and/or arrhythmias including bradyarrhythmias, supraventricular or ventricular arrhythmias. Disease onset is usually in early to mid-adulthood. Sudden cardiac death may occur and may be the presenting symptom. In some cases, it is associated with skeletal myopathy.

### Dane

### Klasyfikacja

Choroba

#### Kod ORPHA

300751

#### Kod OMIM

115200

#### Kod ICD10

I42.0

#### Kod ICD11

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### \*Źródło

orphanet