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