

Opis choroby *

Definicja

A rare, genetic, cardiac rhythm disease characterized by ventricular fibrillation in the absence of any structural or functional heart disease, or known repolarization abnormalities. The presence of J waves is associated with a higher risk of nocturnal ventricular fibrillation events and a higher risk of recurrence.

Dane

Klasyfikacja

Choroba

Synonimy

Familial paroxysmal ventricular fibrillation, non
Brugada type

Kod ORPHA

228140

Kod OMIM

612956

Kod ICD10

I49.0

Kod ICD11

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*Źródło

orphanet