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