Opis choroby *

Definicja

A rare familial cardiomyopathy characterized by the dilation of left ventricle and progressively impairing of systolic ventricular function, in the absence of abnormal loading conditions or coronary artery disease sufficient to cause global systolic impairment. The disease may cause heart failure or arrhythmia. The disease is isolated when no additional atypical cardiac or extracardiac manifestations are present.

Dane

Klasyfikacja
Choroba

Synonimy
Familial or idiopathic dilated cardiomyopathy
Rodzinna izolowana kardiomiopatia rozstrzeniowa

Kod ORPHA
154

Kod OMIM
613642

Kod ICD10
I42.0

Kod ICD11
BC43.00

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*Źródło

orphanet