

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

*Źródło

orphanet