Opis choroby *

Definicja

A rare, genetic muscular dystrophy affecting female carriers and characterized by variable degrees of muscle weakness due to progressive skeletal myopathy, sometimes associated with dilated cardiomyopathy or left ventricle dilation.

Dane

Klasyfikacja
Choroba

Kod ORPHA
206546

Kod OMIM
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Kod ICD10
G71.0

Kod ICD11
-

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*Źródło

orphanet