Opis choroby *

Definicja

A group of rare, genetic, progressive muscular dystrophies, including Duchenne muscular dystrophy (DMD), Becker muscular dystrophy (BMD) and a symptomatic form in female carriers. The diseases represent a spectrum of severity ranging from progressive skeletal and cardiac muscle wasting and weakness (DMD, BMD) to less severe muscle weakness or isolated cardiomyopathy affecting carrier females.

Dane

Klasyfikacja
Grupa fenomenów

Synonimy
Severe dystrophinopathy, Duchenne and Becker type
Ciężka dystrofinopatia typu Duchenne'a i Beckera

Kod ORPHA
262

Kod OMIM
-

Kod ICD10
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Kod ICD11
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*Źródło

orphanet