Opis choroby *

Definicja

A group of rare arthrogryposis syndromes characterized by fetal akinesia, multiple congenital contractures, anterior horn cell degeneration, skeletal muscle atrophy, and other features, depending on the subtype. All types are lethal in the fetal or neonatal period.

Dane

Klasyfikacja
Grupa fenomenów

Synonimy
LCCS
LCCS

Kod ORPHA
294965

Kod OMIM
-

Kod ICD10
Q68.8

Kod ICD11
-

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*Źródło

orphanet