Opis choroby *

Definicja

Peroxisomal acyl-CoA oxidase deficiency is a rare neurodegenerative disorder that belongs to the group of inherited peroxisomal disorders and is characterized by hypotonia and seizures in the neonatal period and neurological regression in early infancy.

Dane

Klasyfikacja
Choroba

Synonimy
Pseudo-NALD
Pseudoadrenoleukodystrofia
Pseudo-NALD
Pseudo-noworodkowa adrenoleukodystrofia
Pseudo-neonatal adrenoleukodystrophy
Pseudoadrenoleukodystrophy

Kod ORPHA
2971

Kod OMIM
264470

Kod ICD10
E71.3

Kod ICD11
5C57.1

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*Źródło

orphanet