Opis choroby *

Definicja

An extremely rare genetic syndromic intellectual disability described in less than 20 families to date and characterized by total or partial alopecia associated with intellectual deficit. The syndrome can be associated with other anomalies such as seizures, sensorineural hearing loss, delayed psychomotor development, and/or hypertonia.

Dane

Klasyfikacja Synonimy

Choroba Perniola-Krajewska-Carnevale syndrome

Zespół Perniola, Krajewskiej i Carnevale

Kod ORPHA Kod OMIM Kod ICD10

2850 613930 Q87.8

Kod ICD11

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*Źródło

orphanet