Opis choroby *

Definicja

A rare genetic neurodevelopmental disorder characterized by early-onset drug-resistant seizures and severe neurodevelopmental impairment with major motor development delay.

Dane

Klasyfikacja
Choroba

Synonimy
CDD
Zaburzenie z niedoboru CDKL5

Kod ORPHA
505652

Kod OMIM
300672

Kod ICD10
G40.4

Kod ICD11
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*Źródło

orphanet