

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 -			

*Źródło

orphanet