Opis choroby *

Definicja

A rare genetic intellectual disability characterized by the association of intellectual disability with variable other anomalies in the absence of a well-characterized syndrome. Associated abnormalities may include facial dysmorphism, neurological signs and symptoms, behavioral problems, and abnormalities of various other organ systems.

Dane

Klasyfikacja
Choroba

Synonimy
Complex neurodevelopmental disorder
Złożona choroba neurorozwojowa

Kod ORPHA
528084

Kod OMIM
620065

Kod ICD10
F84.8

Kod ICD11
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*Źródło

orphanet