Opis choroby *

Definicja

A rare genetic neurodevelopmental syndrome characterized by mild intellectual disability, developmental delay, dysmorphic facial features, growth- and feeding problems, hypotonia, epilepsy, behavioral problems and a variety of congenital abnormalities.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
SIN3A-related intellectual disability syndrome
WITKOS
SIN3A-related intellectual disability syndrome
WITKOS

Kod ORPHA
500163

Kod OMIM
-

Kod ICD10
Q87.8

Kod ICD11
-

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*Źródło

orphanet