Opis choroby *

Definicja

A rare genetic, intellectual disability syndrome characterized by intellectual disability, childhood hypotonia, severe expressive speech delay, autism spectrum disorder, and a distinctive facial appearance with a spectrum of additional clinical features.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
261494

Kod OMIM
610253

Kod ICD10
Q87.8

Kod ICD11
LD2F.1Y

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*Źródło

orphanet