Opis choroby *

Definicja

Intellectual disability-severe speech delay-mild dysmorphism syndrome is a rare, genetic, syndromic intellectual disability disorder, with highly variable phenotype, typically characterized by mild to severe global development delay, severe speech and language impairment, mild to severe intellectual disability, dysphagia, hypotonia, relative to true macrocephaly, and behavioral problems that may include autistic features, hyperactivity, and mood lability. Facial gestalt typically features a broad, prominent forehead, hypertelorism, downslanting palpebral fissures, ptosis, a short bulbous nose with broad tip, thick vermilion border, wide, and open mouth with downturned corners. Brain, cardiac, urogenital and ocular malformations may be associated.

Dane

Klasyfikacja Synonimy

Zespół wad wrodzonych FOXP1 syndrome

FOXP1 syndrome

 Kod ORPHA
 Kod OMIM
 Kod ICD10

 391372
 613670
 Q87.0

Kod ICD11

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<u>*Źródło</u>

orphanet