Opis choroby *

Definicja

X-linked microcephaly-growth retardation-prognathism-cryptorchidism syndrome is a rare syndromic intellectual disability characterized by hypotonia, microcephaly, severe developmental delay, seizures, intellectual disability, growth retardation, cardiac septal defects, cryptorchidism, hypospadias, and dysmorphic features - prominent ears, prognathism, thin upper lip, dental crowding.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
435938

Kod OMIM
300998

Kod ICD10
Q87.8

Kod ICD11
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*Źródło

orphanet