Opis choroby *

Definicja

A very rare multiple congenital anomalies syndrome described in three brothers of one South-African family, and characterized by hypospadias and intellectual deficit, in association with microcephaly, craniofacial dysmorphism, joint laxity and beaked nails.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Goldblatt-Wallis syndrome
Zespół Goldblatta i Wallisa

Kod ORPHA
2261

Kod OMIM
241760

Kod ICD10
Q87.8

Kod ICD11
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*Źródło

orphanet