

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 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