

Opis choroby *

Definicja

A rare multiple congenital anomalies/dysmorphic syndrome characterized by brachydactyly, nystagmus, and cerebellar ataxia. Intellectual deficit and strabismus have also been reported. There have been no further descriptions in the literature since 1934.

Dane

Klasyfikacja

Zespół wad wrodzonych
Biemond syndrome
Zespół Biemonda

Synonimy

Kod ORPHA

1246

Kod OMIM

113400

Kod ICD10

Q87.8

Kod ICD11

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*Źródło

orphanet