

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	Synonimy
Zespół wad wrodzonych	Biemond syndrome
	Zespół Biemonda

Kod ORPHA	Kod OMIM	Kod ICD10
1246	113400	Q87.8

Kod ICD11
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*Źródło

orphanet