

Opis choroby *

Definicja

A rare, genetic, multiple congenital anomalies/dysmorphic syndrome characterized by the triad: congenital, bilateral, symmetrical, subtotal, external auditory canal atresia, bilateral vertical talus and increased interocular distance.

Dane

Klasyfikacja **Synonimy**

Zespół wad wrodzonych Rasmussen-Johnsen-Thomsen syndrome

Kod ORPHA

3023

Kod OMIM

133705

Kod ICD10

Q87.8

Kod ICD11

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*Źródło

orphanet