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
Zespół wad wrodzonych

Synonimy
Rasmussen-Johnsen-Thomsen syndrome

Kod ORPHA
3023

Kod OMIM
133705

Kod ICD10
Q87.8

Kod ICD11
-

---

*Źródło

orphanet