Opis choroby *

Definicja

A rare genetic multiple congenital anomalies/dysmorphic syndrome characterized by the association of auricular abnormalities (such as external ear abnormalities and postauricular pits) and cleft lip with or without cleft palate. Additional manifestations include myopia, nystagmus, and retinal pigment abnormalities.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
77300

Kod OMIM
-

Kod ICD10
Q87.0

Kod ICD11
-

---

*Źródło

orphanet