Zespół skrzelowo-uszny
Kod Orpha: 52429 Kod OMIM: 608389

Opis choroby *

Definicja
Branchiootic syndrome is a rare, genetic multiple congenital anomalies syndrome characterized by second branchial arch anomalies (branchial cysts and fistulae), malformations of the outer, middle and inner ear associated with sensorineural, mixed or conductive hearing loss, and the absence of renal abnormalities. Typical ear findings consist of malformed auricles (e.g. lop or cupped ears), preauricular pits and/or tags, and middle and/or inner ear dysplasias (inculding cochlear, vestibular and semicircular channel hypoplasia, malformation of the ossicles and of middle ear space).
Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
52429

Kod OMIM
608389

Kod ICD10
Q87.0

Kod ICD11
LD2F.1Y

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.