Opis choroby *

Definicja

A rare, genetic, multiple congenital anomalies syndrome characterized by urinary tract anomalies, nephrosis, conductive deafness, and digital malformations, including short and bifid distal phalanges of thumbs and big toes. There have been no further descriptions in the literature since 1962.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Braun-Bayer syndrome
Zespół Brauna i Bayera
Nephrosis-hearing loss-urinary tract-digital malformations syndrome

Kod ORPHA
2669

Kod OMIM
256200

Kod ICD10
Q87.8

Kod ICD11
LD2H.Y

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*Źródło

orphanet