

## **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 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