

Opis choroby *

Definicja

A rare multiple congenital anomalies syndrome characterized by the association of uni- or bilateral radial defects, uni- or bilateral Duane anomaly (congenital limited horizontal eye movement accompanied by globe retraction which results in narrowing of the palpebral fissure), renal abnormalities, sensorineural and/or conductive hearing loss, and, less frequently, imperforate anus and scoliosis.

Dane

Klasyfikacja	Synonimy
Zespół wad wrodzonych	Duane-radial ray syndrome Zespół anomalii Duana i kości piszczelowej

Kod ORPHA	Kod OMIM	Kod ICD10
93293	607323	Q87.8

Kod ICD11
LD2F.1Y

*Źródło

orphanet