Opis choroby *

Definicja

A very rare multiple congenital anomalies syndrome characterized by limb deficiencies and renal anomalies that include split hand-split foot malformation, renal agenesis, polycystic kidneys, uterine anomalies and severe mandibular hypoplasia. An autosomal recessive mode of inheritance has been suggested.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Split hand/split foot-mandibular hypoplasia syndrome
Rozszczep dłoni/rozszczep stopy - hipoplazja żuchwy

Kod ORPHA
958

Kod OMIM
200980

Kod ICD10
Q87.8

Kod ICD11
LD2F.1Y

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*Źródło

orphanet