## 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