Opis choroby *

Definicja

Laurin-Sandrow syndrome (LSS) is characterised by complete polysyndactyly of the hands, mirror feet and nose anomalies (hypoplasia of the nasal alae and short columella), often associated with ulnar and/or fibular duplication (and sometimes tibial agenesis). It has been described in less than 20 cases. Some cases with the same clinical signs but without nasal defects have also been reported, and may represent the same entity. The etiology of LSS is unknown. Different modes of inheritance have been suggested.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Mirror hands and feets-nasal defects syndrome
Lustrzane odbicie dłoni i stóp - wady nosa
Zespół Sandrowa
Sandrow syndrome

Kod ORPHA
2378

Kod OMIM
135750

Kod ICD10
Q87.2

Kod ICD11
LD26.2

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*Źródło

orphanet