Opis choroby *

Definicja

A rare syndrome with limb malformations as a major feature characterized by congenital scalp defects and postaxial polydactyly type A. There is a wide variability of expression, with some patients showing only one of the typical manifestations. There have been no further descriptions in the literature since 1985.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
1003

Kod OMIM
181250

Kod ICD10
Q87.2

Kod ICD11
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*Źródło

orphanet