Opis choroby *

Definicja

A rare, genetic, ectodermal dysplasia syndrome characterized by severe hand/foot anomalies, breast and/or nipple hypoplasia, and ectodermal dysplasia (principally teeth and nail anomalies). Cleft lip/palate may be variably present.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
LMS
LMS

Kod ORPHA
69085

Kod OMIM
603543

Kod ICD10
Q82.4

Kod ICD11
LD27.0Y

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*Źródło

orphanet