

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 | Synonimy |
| Zespół wad wrodzonych LMS | LMS |

| | | |
|------------------|-----------------|------------------|
| Kod ORPHA | Kod OMIM | Kod ICD10 |
| 69085 | 603543 | Q82.4 |

Kod ICD11
LD27.0Y

[*Źródło](#)

orphanet