

## Opis choroby \*

### Definicja

An intermediate form of lichen myxedematosus (LM) (a form of mucin dermal deposit) which does not meet the criteria for either scleromyxedema or the localized form. Three clinical subtypes have been described and include scleromyxedema without monoclonal gammopathy; localized forms with monoclonal gammopathy and/or systemic symptoms; localized forms with mixed features of the 5 subtypes of localized LM (discrete form, acral persistent papular mucinosis, self-healing papular mucinosis, papular mucinosis of infancy, and a pure nodular form). The course of atypical LM is unpredictable because only a few cases have been reported.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Intermediate lichen myxedematosus

Liszaj śluzowaty średniozaawanowany

#### Kod ORPHA

86797

#### Kod OMIM

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#### Kod ICD10

L98.5

#### Kod ICD11

EB90.1Y

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#### \*Źródło

orphanet