Opis choroby *

Definicja

A rare primary cutaneous amyloidosis characterized by macular or reticulate hyperpigmentation with symmetrically distributed guttate hypo- and hyperpigmented lesions which progress gradually over the years to involve almost the entire body (with relative sparing of the face, hands, feet and neck). Patients are usually asymptomatic, however mild pruritus may be associated. Amyloid deposition in the papillary dermis is observed on skin biopsy. Systemic amyloidosis is not present and association with generalized morphea, atypical Parkinsonism, spasticity, motor weakness or colon carcinoma is rare.

Dane

Klasyfikacja
Choroba

Synonimy
Amyloidosis cutis dyschromica
Amyloidosis cutis dyschromica

Kod ORPHA
319635

Kod OMIM
617920

Kod ICD10
L99.0*

Kod ICD11
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*Źródło

orphanet