Opis choroby *

Definicja

A clonal B-cell disorder characterized by the aggregation and deposition of insoluble amyloid fibrils derived from misfolding of monoclonal immunoglobulin light chains. It usually presents as systemic AL amyloidosis with involvement of one or more parenchymal organ(s) and, less frequently, as localized amyloidosis with usually nodular deposits restricted to a single organ and/or system.

Dane

Klasyfikacja
Choroba

Synonimy
Light-chain amyloidosis
Amyloidoza immunoglobulinowa
Amyloidoza łańcuchów lekkich
Amyloidoza pierwotna
Primary amyloidosis

Kod ORPHA
85443

Kod OMIM
254500

Kod ICD10
E85.9

Kod ICD11
5D00.0

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*Źródło

orphanet