

Opis choroby *

Definicja

A rare non-amyloid monoclonal immunoglobulin deposition disease characterized by deposition of abnormal immunoglobulin light chains in the kidneys, resulting in nephrotic syndrome and renal failure. Symptomatic extrarenal deposition is uncommon, although hepatic, cardiac, and neural deposits have been reported. The condition frequently occurs in association with multiple myeloma or in patients with M protein and marrow plasma cells at monoclonal gammopathy of undetermined significance levels.

Dane

Klasyfikacja

Podtyp kliniczny

Synonimy

LCDD

LCDD

Kod ORPHA

93558

Kod OMIM

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Kod ICD10

D89.8

Kod ICD11

2A83.52

*Źródło

orphanet