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
-

Kod ICD10
D89.8

Kod ICD11
2A83.52

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*Źródło

orphanet