Opis choroby *

Definicja

A rare, systemic amyloidosis characterized by slowly progressive renal disease presenting with proteinuria, hypertension and decreased glomerular filtration rate leading to progressive renal failure. Histology reveals amyloid deposits of leukocyte chemotactic factor-2 protein in the renal cortical interstitium, tubular basement membranes, glomeruli and the vessel walls. Extra-renal deposits can be seen in the liver, lungs, spleen and adrenal glands.

Dane

Klasyfikacja
Choroba

Synonimy
Leukocyte chemotactic factor-2 amyloidosis
Amyloidoza czynnika-2 chemotaktycznego leukocytów

Kod ORPHA
439224

Kod OMIM
-

Kod ICD10
E85.8

Kod ICD11
5D00.0

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*Źródło

orphanet