Opis choroby *

Definicja

A group of rare renal diseases, characterized by amyloid fibril deposition of apolipoprotein A-I or A-II (AApoAI or AApoAII amyloidosis), lysozyme (ALys amyloidosis) or fibrinogen A-alpha chain (AFib amyloidosis) in one or several organs. Renal involvement leading to chronic renal disease and renal failure is a common sign. Additional manifestations depend on the organ involved and the type of amyloid fibrils deposited.

Dane

Klasyfikacja
Choroba

Synonimy
Amyloidosis, Ostertag type
Amyloidoza, typ Ostertaga
Dziedziczna amyloidoza nerek
Dziedziczna nefropatia amyloidowa
Rodzinna amyloidoza nerkowa
Rodzinna nefropatia amyloidowa
Familial amyloid nephropathy
Familial renal amyloidosis
Hereditary amyloid nephropathy
Hereditary renal amyloidosis

Kod ORPHA
85450

Kod OMIM
105200

Kod ICD10
E85.0

Kod ICD11
5D00.2Y

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*Źródło

orphanet