Opis choroby *

Definicja

A rare non-immune-mediated glomerular disease characterized by abnormal accumulation of type III collagen within the mesangium and subendothelial space of the glomerulus. Clinically it usually manifests with proteinuria (often in the nephrotic range), microscopic hematuria, peripheral edema and/or hypertension. Progression to end-stage kidney failure is possible.

Dane

Klasyfikacja
Choroba

Synonimy
Collagenofibrotic glomerulopathy
Collagenofibrotic glomerulopathy

Kod ORPHA
84087

Kod OMIM
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Kod ICD10
N07.6

Kod ICD11
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*Źródło

orphanet