Opis choroby *

Definicja

A rare genetic renal disease characterized by the formation of intraglomerular lipoprotein thrombi due to lipid deposition in severely dilated glomerular capillaries. Laboratory examination reveals abnormal serum lipid profiles, in particular markedly elevated apolipoprotein E. Clinical manifestations include proteinuria or nephrotic syndrome with hypertension and potential progression to chronic renal failure. Systemic complications of dyslipidemia are not observed.

Dane

Klasyfikacja
Choroba

Synonimy
LPG
LPG

Kod ORPHA
329481

Kod OMIM
611771

Kod ICD10
N07.8

Kod ICD11
MF82

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*Źródło

orphanet