

## 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

<b>Klasyfikacja</b>	<b>Synonimy</b>
Choroba	LPG LPG

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
329481	611771	N07.8

**Kod ICD11**  
MF82

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### \*Źródło

orphanet