Opis choroby *

Definicja

A rare, genetic renal tubular disease characterized by tubular damage and interstitial fibrosis in absence of glomerular lesions and clinically manifesting with chronic kidney disease (CKD) and slow progression to end-stage kidney disease (ESKD).

Dane

Klasyfikacja
Choroba

Synonimy
ADTKD
Autosomalna dominująca kanalikowo-śródmiąższowa choroba nerek
Familial juvenile hyperuricemic nephropathy
MCKD
Medullary cystic kidney disease

Kod ORPHA
34149

Kod OMIM
174000

Kod ICD10
Q61.5

Kod ICD11
GB82

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*Źródło

orphanet