

## 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-  
śródmieższowa 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