

## **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	Synonimy
Choroba	ADTKD
	Autosomalna dominująca kanalikowo-śródmiąższowa choroba nerek
	Familial juvenile hyperuricemic nephropathy
	MCKD
	Medullary cystic kidney disease
<b>Kod ORPHA</b>	<b>Kod OMIM</b>
34149	174000
<b>Kod ICD11</b>	<b>Kod ICD10</b>
GB82	Q61.5

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\*[Źródło](#)

orphanet