

## Opis choroby \*

### Definicja

A rare, genetic, renal tubular disease characterized by progressive outgrowths of fluid-filled cysts from the renal epithelium, which can manifest with hematuria, urinary tract infections, hypertension, and abdominal or flank pain. The slowly progressive loss of kidney function may evolve to end stage kidney disease (ESKD).

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

ADPKD

ADPKD

#### Kod ORPHA

730

#### Kod OMIM

600666

#### Kod ICD10

Q61.2

#### Kod ICD11

GB81

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### \*Źródło

orphanet