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