

Opis choroby *

Definicja

A rare renal tract malformation characterized by dilated malformation of the medullary collecting ducts (typically bilateral), and associated with stone formation, renal colic, hematuria, urinary tract infection, nephrocalcinosis, calcium nephrolithiasis, pyelonephritis, hypercalciuria and hypocitraturia. The disease is associated with abnormal distal tubular functions.

Dane

| Klasyfikacja | Synonimy |
|--------------------|---------------------------------|
| Wada morfologiczna | Cacchi-Ricci disease |
| | Choroba Cacchi i Ricciego |
| | MSK |
| | MSK |
| | Precalicial canalicular ectasia |

| Kod ORPHA | Kod OMIM | Kod ICD10 |
|-----------|----------|-----------|
| 1309 | - | Q61.5 |

| Kod ICD11 |
|-----------|
| LB30.8 |

*[Źródło](#)

orphanet