Opis choroby *

Definicja

Congenital megacalycosis is a rare renal malformation, characterized by non-obstructive dilation of the renal calyces as well as an increased calyceal number (12-20), with a normal renal pelvis, ureter, and bladder. It may be unilateral or bilateral and is usually asymptomatic unless complicated by nephrolithiasis and urinary tract infection.

Dane

Klasyfikacja
Wada morfologiczna

Kod ORPHA
93109

Kod OMIM
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Kod ICD10
Q63.8

Kod ICD11
LB31.Y

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*Źródło

orphanet