## Opis choroby *

Definicja

A rare congenital anomaly of the kidney and urinary tract (CAKUT) in which one or both kidneys (unilateral or bilateral MCDK respectively) are large, distended by multiple cysts, and non-functional. Unilateral MCDK is typically asymptomatic if the other kidney is fully functional but may occasionally present with abdominal obstructive signs when the cysts become too large. Bilateral MCDK is considered a lethal entity and neonates present with features of the Potter sequence, severe pulmonary hypoplasia and severe renal failure, and generally die shortly after birth.

Dane

**Klasyfikacja**
Wada morfologiczna

Synonimy
MCDK
Dysplazja wielotorbielowata nerki
MCDK
Multicystic renal dysplasia

**Kod ORPHA**
1851

**Kod OMIM**
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**Kod ICD10**
Q61.4

**Kod ICD11**
LB30.9

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*Źródło

orphanet