Open Peer Review on Qeios

Multicystic dysplastic kidney

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Multicystic dysplastic kidney. ORPHA:1851

Multicystic dysplastic kidney (MCDK) is a congenital anomaly of the kidney and urinary tract (CAKUT) in which one or both kidneys (unilateral or bilateral MCDK respectively; see these terms) are large, distended by multiple cysts, and non-functional.