McKusick-Kaufman syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. McKusick-Kaufman syndrome. ORPHA:2473

McKusick-Kaufman syndrome is a very rare, genetic developmental disorder presenting in the neonatal period characterized by genitourinary malformations, polydactyly, and more rarely, congenital heart disease or gastrointestinal malformations.