NPHP3-related Meckel-like syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. NPHP3-related Meckel-like syndrome. ORPHA:3032

NPHP3-related Meckel-like syndrome is a rare, genetic, syndromic renal malformation characterized by cystic renal dysplasia with or without prenatal oligohydramnios, central nervous system abnormalities (commonly Dandy-Walker malformation), congenital hepatic fibrosis, and absence of polydactyly.