Perlman syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Perlman syndrome. ORPHA:2849

Perlman syndrome is characterized principally by polyhydramnios, neonatal macrosomia, bilateral renal tumours (hamartomas with or without nephroblastomatosis), hypertrophy of the islets of Langerhans and facial dysmorphism.