Radio-renal syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Radio-renal syndrome. ORPHA:3015

Radio-renal syndrome is a rare developmental defect during embryogenesis characterized by variable upper limb reduction defects and renal anomalies. Patients typically present absence/hypoplasia of digits, radii and/or ulnae, short stature and mild external ear malformation, as well as kidney agenesis or ectopia. There have been no further descriptions in the literature since 1983.