Acrorenal syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Acrorenal syndrome. ORPHA:971

Acrorenal syndrome comprises a wide spectrum of congenital malformative disorders characterized by the co-occurrence of distal limb anomalies (usually bilateral cleft feet and/or hands) and renal defects (e.g. unilateral or bilateral agenesis), that can be associated with a variety of other anomalies such as those of genitourinary tract (genital anomalies, ureteral hypoplasias, vesicoureteral reflux), abdominal well defects, intestinal atresias, and lung malformations. Familial cases have been reported in which an autosomal recessive inheritance was suspected.