Oligomeganephronia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Oligomeganephronia. ORPHA:2260

Oligomeganephronia is a developmental anomaly of the kidneys, and the most severe form of renal hypoplasia (see this term), characterized by a reduction of 80% in nephron number and a marked hypertrophy of the glomeruli and tubules.