Autosomal dominant polycystic kidney disease type 1 with tuberous sclerosis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Autosomal dominant polycystic kidney disease type 1 with tuberous sclerosis. ORPHA:88924

Polycystic kidney disease with tuberous sclerosis (PKD-TSC) is characterised by early-onset and severe polycystic kidney disease with various manifestations of tuberous sclerosis (multiple angiomyolipomas, lymphangioleiomyomatosis and periventricular calcifications of the central nervous system).