REN-related autosomal dominant tubulointerstitial kidney disease

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. REN-related autosomal dominant tubulointerstitial kidney disease. ORPHA:217330

Familial juvenile hyperuricemic nephropathy type 2 is a rare autosomal dominantly inherited disease of childhood characterized by hypoproliferative anemia, hyperuricemia and slowly progressing kidney failure due to dysregulation of the renin-angiotensin system (RAS).