Familial juvenile hyperuricemic nephropathy type 1

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial juvenile hyperuricemic nephropathy type 1. ORPHA:209886

Familial juvenile hyperuricemic nephropathy type 1 (FJHN1) is a rare kidney disorder characterized by hyperuricemia, progressive nephropathy, and gout occurring at an early age.