Familial primary hypomagnesemia with hypercalciuria and nephrocalcinosis without severe ocular involvement

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial primary hypomagnesemia with hypercalciuria and nephrocalcinosis without severe ocular involvement. ORPHA:31043

Familial primary hypomagnesemia with hypercalciuria and nephrocalcinosis without severe ocular involvement (FHHN) is a form of familial primary hypomagnesemia (FPH; see this term), characterized by recurrent urinary tract infections, nephrolithiasis, bilateral nephrocalcinosis, renal magnesium (Mg) wasting, hypercalciuria and kidney failure.