Familial isolated hypoparathyroidism

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial isolated hypoparathyroidism. ORPHA:2238

Familial isolated hypoparathyroidism (FIH) is a rare heterogeneous group of metabolic disorders characterized by abnormal calcium metabolism due to deficient secretion of parathormone (PTH), without other endocrine disorders or developmental defects.