# Familial hypocalciuric hypercalcemia

INSERM

## Source

*INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial hypocalciuric hypercalcemia. ORPHA:405*

Familial hypocalciuric hypercalcemia (FHH) is a generally asymptomatic genetic disorder of phosphocalcic metabolism characterized by lifelong moderate hypercalcemia along with normo- or hypocalciuria and elevated plasma parathyroid hormone (PTH) concentration.