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# Familial primary hypomagnesemia

INSERM

## Source

INSERM. (1999). *Orphanet: an online rare disease and orphan drug data base*. Familial primary hypomagnesemia. ORPHA:34526

A rare mineral absorption and transport disorder characterized by a selective defect in renal or intestinal magnesium (Mg) absorption, resulting in a low Mg concentration in the blood.