X-linked hypophosphatemia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. X-linked hypophosphatemia. ORPHA:89936

X-linked hypophosphatemia (XLH) is a hereditary renal phosphate-wasting disorder characterized by hypophosphatemia, rickets and/or osteomalacia, and diminished growth.