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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.

Qeios ID: GT3ZO5 · https://doi.org/10.32388/GT3ZO5