Prenatal benign hypophosphatasia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Prenatal benign hypophosphatasia. ORPHA:247638

Prenatal benign hypophosphatasia (PB-HPP) is a very rare form of hypophosphatasia (see this term) characterized by prenatal skeletal manifestations (limb shortening and bowing) that slowly resolve spontaneously and later develop into the milder infantile, childhood or adult forms of the disease.