Perinatal lethal hypophosphatasia

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Perinatal lethal hypophosphatasia. ORPHA:247623

Perinatal lethal hypophosphatasia (PL-HPP) is a very rare form of hypophosphatasia (see this term) characterized by markedly impaired bone mineralization in utero due to reduced activity of serum alkaline phosphatase (ALP) and causing stillbirth or respiratory failure within days of birth.