Hypophosphatasia commonly known as “HPP” disease is a genetic disorder, characterized by: low alkaline phosphatase (ALP), high B6 (PLP), skeletal deformities, fractures, premature tooth loss, and chronic pain. Decreased renal function, chiari malformation, and craniosynostosis may also be present.

Please handle with caution.

Symptoms of fractures are not always obvious; pain often indicates a fracture. X-rays may not always indicate a fracture.

For more information contact:
www.softbones.org or (866) 827-9937

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