## MORE THAN HALF OF PATIENTS

with HPP may not have overt bone manifestations\*1



Rethink HPP disease burden especially the impact on daily function and QoL, and the potential for progression in children and adults<sup>3–5</sup>

\*Skeletal manifestations were present in 44.2% (N=53/120) of paediatric (<18 years) and 48.2% (N=66/137) of adult (≥18 years) patients with HPP.¹ Data from the Global HPP Registry, sponsored by Alexion Pharmaceuticals, an observational, longitudinal, multinational, long-term study collecting data on HPP diagnosis, history, clinical course, symptoms (including multisystemic aspects of disease) and burden of illness from patients who have a diagnosis of HPP.¹⁵ †Includes craniosynostosis, developmental delay, increased intracranial pressure and seizures.¹ †Includes bone deformity, pseudofractures, recurrent and poorly healing fractures, rickets-like changes (by radiograph).¹ HPP, hypophosphatasia; QoL, quality of life.

- 1. Högler W, et al. BMC Musculoskelet Disord. 2019;20(1):80. 2. Dahir KM, et al. Orphanet J Rare Dis. 2022;17(1):277.
- **3.** Rush ET, et al. *Orphanet J Rare Dis*. 2019;14(1):201. **4.** Seefried L, et al. *J Bone Miner Res*. 2020;35(11):2171–2178.
- **5.** Szabo SM, et al. *Orphanet J Rare Dis.* 2019;14(1):85.

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