

**Calling  
All Doctors!**

## Join the Hypophosphatasia Conversation on HPP Connect

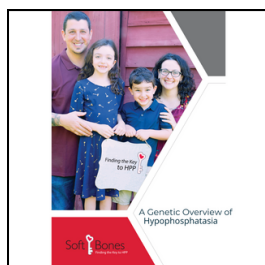
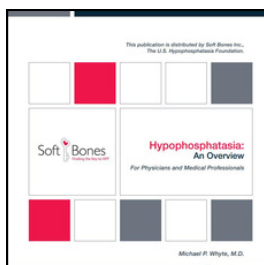
HPP Connect is an online collaborative community for ongoing, multi-specialty, peer-to-peer exchange, dialogue and education focused on hypophosphatasia (HPP). It connects clinicians with experience evaluating and treating HPP to one another to foster knowledge-sharing and collaboration.

## View the Soft Bones TeleECHO and Second International Scientific Meeting On-Demand

The HPP TeleECHO curriculum was designed to provide continuing education for the HPP healthcare team. Through "telementoring" presentations and discussions on different HPP-related topics, it also aimed to improve patient care. From August 2021 - March 2023, specialists presented on various topics to reflect the multidisciplinary nature of HPP. View our on-demand recordings from the HPP TeleECHO and 2nd International Scientific Meeting on HPP Connect.

## Access Patient & Physician Resources

From diagnosis to treatment, resources are available on a variety of topics:



## About Soft Bones

Soft Bones, Inc., The US Hypophosphatasia Foundation, is a patient advocacy group founded in 2009 to provide information and to establish a forum to educate, empower, and connect patients living with hypophosphatasia, their families, and caregivers. The Foundation also promotes research of rare bone disease through awareness and fundraising efforts. For more information, contact [Info@SoftBones.org](mailto:Info@SoftBones.org).

