SAVE THE DATE

HPP in focus: Genetics, manifestations, and diagnosis

TELEBRIDGE SERIES

Series objectives

- Describe the genetics, clinical manifestations, diagnostic criteria and burden of disease of HPP in children and adults
- Illustrate the patient journey in HPP and identify barriers to diagnosis
- Highlight low ALP level as a key marker to trigger further examinations in clinical practice

Thursday 9 October
17:00–19:00 (CEST)

Central Europe

Czech Republic
Slovakia
Hungary

Dr. Izabela Michalus
and Dr. Jordi Pérez-López

Thursday 16 October
18:00–20:00 (EEST)

Baltics

Lithuania
Estonia
Latvia

Dr. Adela Chirita-Emandi
and Dr. Jordi Pérez-López

Thursday 23 October
16:00–18:00 (CEST)

Balkans

Croatia
Serbia
Slovenia
Bulgaria

Dr. Adela Chirita-Emandi
and Dr. Jordi Pérez-López

Thursday 30 October
17:00–19:00 (EET)

Romania

Dr. Izabela Michalus
and Dr. Jordi Pérez-López

We look forward to welcoming you to one of these webinars!

ALP, alkaline phosphatase; HPP, hypophosphatasia.
This meeting is organised and funded by Alexion, AstraZeneca Rare Disease and is intended for healthcare professionals only.
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