THEMATIC MEETING

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PKU BREAKING THE DOGMA

17TH INTERNATIONAL SYMPOSIUM OF THE PORTUGUESE SOCIETY FOR METABOLIC DISORDERS

HYBRID EVENT

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HOTEL LUX , FÁTIMA - PORTUGAL

9TH SEPTEMBER 2021

18:00 - 19:30 (GMT)

Phenylketonuria (PKU) is a rare, inherited, chronic and life-long condition. Advances in the treatment of this disease led to different perspectives on its management, explored in this meeting.

AGENDA (GMT)

18:00 - 18:05 **Welcome**

18:05 - 18:20 **Kuvan in Infants |** *María Bueno Delgado*

18:20 - 18:35 **The problem with self-awareness in PKU |** *Cary O. Harding*

18:35 - 18:50 **New road map for nutritional management |** *Júlio César Rocha*

18:50 - 18:20 **Open Discussion**

Moderator: Júlio César Rocha
Álvaro Hermida Ameijeiras, María Bueno Delgado

19:20 - 19:25 **Patient Experience**

19:25 - 19:30 **Conclusions**

SPEAKERS

MARÍA BUENO DELGADO
Paediatrician. Hospital Universitario Virgen del Rocío.
Sevilla, Spain

CARY O. HARDING
Clinical Geneticist. Oregon Health & Science University.
Portland, OR, USA

JÚLIO CÉSAR ROCHA
Nutritionist. Centro Hospitalar Universitário de Lisboa Central.
Lisbon, Portugal

ÁLVARO HERMIDA AMEIJEIRAS
Internist. Hospital Clínico Universitario de Santiago de Compostela.
Santiago de Compostela, Spain