



2019

programme

Advancing knowledge in rare diseases

Mitochondrial medicine 30 years on: state of the art

11 – 13 April, Nice, France

Pascal Laforêt, Paris
Shamima Rahman, London
Angeles Garcia-Cazorla, Barcelona

Inborn errors of metabolism in neonatology: clues for successful diagnosis and treatment

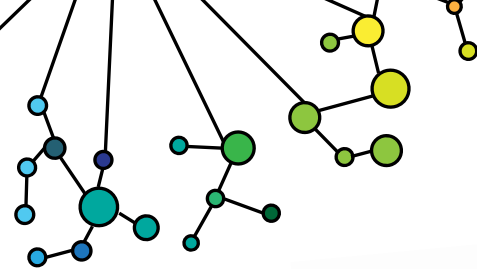
13 – 15 June, Santiago de Compostela, Spain

Ivo Barić, Zagreb
Maria-Luz Couce, Santiago de Compostela

The life course of inherited metabolic diseases: from childhood to adulthood

07 – 09 November, Hong Kong

Grace Poon, Hong Kong
Sheng Bun, Hong Kong
CW Fung, Hong Kong
Robin Lachmann, London



The foundation develops and delivers tailored solutions in training and education, for healthcare professionals, to promote accurate diagnosis of rare diseases and care of patients.

Contact :

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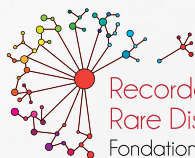
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