Garrod Symposium 2007

"Neurometabolic Disorders and Newborn Screening"

Updated Programme

Saturday May 5th

Morning session

Barbara Plecko, Graz / Vancouver
Pyridoxine dependent epilepsy: a newly discovered gene defect in a well known condition

Ingrid Tein, Toronto
Metabolic myopathies: New insights in the carnitine transporter deficiency.

Nina Raben, Bethesda
Progress in understanding Pompe Disease

Marc Patterson, New York
Niemann Pick type C

Joseph Muenzer, Chapel Hill
ERT for MPSII – Hunter Disease

Free Communications

Afternoon session 13.30 – 17.00

Eduard Struijs, Amsterdam
The importance of innovative analytical methods (e.g. GC-MS and LC-MS/MS) for investigation of inborn errors of metabolism

Cary Harding, Portland
Challenge and importance of collecting clinical outcome information from newborn screening

Mark Montgomery, Calgary
How to implement a successful CF newborn screening program

Sandra Sirrs, Vancouver
Canadian Fabry Disease Initiative (CFDI) as a model for a national collaborative study on Inborn Errors of Metabolism

Free communications

Sunday May 6th

Garrod Association Board Meeting
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