Invitation

Metabolic Symposium
Reykjavik, 29 September 2016

Could it be an inborn error of metabolism?

13:30 – 19:00
Hilton Reykjavik Nordica, Sudurlandsbraut 2, Reykjavik

Please sign up by sending an email mentioning:

• Name
• Title
• Hospital/institution
• Address
• and if you are staying for dinner (please mention any dietary requirements you may have)

to: linda.olsson@bmrn.com

Please reply by Wednesday 31 Aug 2016 at the latest
Metabolic Symposium

The incidence of each inherited metabolic disease is low, but since there are many different diseases the total number of patients is relatively large.

The diseases can appear at any age and with symptoms from different organs. Many are treatable with a good prognosis if the diagnosis is made in time.

With this symposium we aim to increase the awareness regarding how and when to suspect that a patient suffers from a treatable metabolic disease. The aim is also to guide clinicians regarding the diagnostic and laboratory work-up when investigating a patient for metabolic diseases.

This symposium is organized in collaboration with the Department of Genetics and Molecular Medicine, Landspítali, Reykjavik, Iceland and is supported by unrestricted educational grants from Actelion, Alexion, BioMarin, Genzyme, Orphan Europe and Shire.

Program

13:30 – 14:00  Registration and welcome coffee

14:00 – 14:20  Welcome and introduction
Dr Leifur Franzson, Landspitali University Hospital

14:20 – 15:00  Mucopolysaccharide disorders – overview and management
Prof Frits Wijburg, Academic Medical Centre, Amsterdam, the Netherlands

15:00 – 15:40  Eye symptoms in connection with metabolic disorders
Adjunct Prof Kristina Teär Fahnehjelm, Gothenburg University, Sweden

15:40 – 16:20  Homocysteinuria
Dr Rolf Zetterström, Karolinska University Hospital, Solna

16:20 – 16:35  Break

16:35 – 17:15  Niemann-Pick Disease type C – overview and management
Prof Frits Wijburg, Academic Medical Centre, Amsterdam, the Netherlands

17:15 – 17:55  Fabry Disease – overview and management
Dr Jukka Saarinen, Vaasa Central Hospital, Finland

17:55 – 18:35  Hypophosphatasia (HPP)
Prof Outi Mäkitie, Karolinska Institute, Stockholm

18:35 – 19:00  General discussion and closure
Dr Leifur Franzson, Landspitali University Hospital

19:00  Dinner