BALTIC METABOLIC GROUP MEETING: INHERITED NEUROMETABOLIC DISORDERS

31st of March - 1st of April, 2011

Vilnius

Program

***Meeting venue**: Conference hall (red auditorium), Vilnius University Hospital Santariskiu Clinics, Santariskiu str.2, Vilnius*

31st of March (Thursday)

10.30 – 12.00	***Registration***
12.00 – 12.15	Welcome addresses: prof. Vaidutis Kučinskas
12.15 – 12.55	Monique Williams (Rotterdam): Neonatal presentation of neurometabolic disorders
13.00 – 13.15	K. Kall (Tartu): Biochemical changes in patients with PDH deficiency
13.15 – 14.00	***Lunch***
14.00 – 14.45	Nico Abeling (Amsterdam): Neurotransmitter defects: a diagnostic challenge
14.55 – 15.10	K. Õunap (Tartu): Unusual genetic cause of OTC deficiency in 9-year old girl
15.15 – 15.30	D. Bauze (Riga): 2-Hydroxyglutaric aciduria, L-form – case report
15.35 – 15.50	J. Songailienė (Vilnius): Formiminoglutamate aciduria – case report
15.55 – 16.15	***Coffee break***
16.15 – 16.30	D. Lomele (Riga): Riboflavin responsive Multiple acyl-Coa dehydrogenase deficiency? – case report.
16.35 – 16.50	Nico Abeling (Amsterdam): Brown-Vialetto-Van Laere syndrome mimicking Riboflavin - responsive MADD is caused by riboflavin transporter defect. A new treatable inborn error of metabolism.
16.55 – 17.05	L. Cimbalistienė (Vilnius): Non-ketotic hyperglycinemia – case report
17.05 – 17.15	I. Grinfelde (Riga): Infant with sialidosis - case report
19.00	***Diner: Restaurant JUVENTUS, Universiteto str.7, Vilnius***

1st of April (Friday)

9.00 – 10.45	Workshop
10.45 – 11.05	***Coffee break***
11.05-11.20	B. Skerlienė (Vilnius): 3-methylglutaconic aciduria (MGA) type II - Barth syndrome; Case report.
11.25-11.40	M-L. Uudelepp (Estonia): First diagnosed case of MCAD in Estonia
11.45- 12.00	K. Joost (Estonia): The prevalence of long-chain 3-hydroxyacyl-CoA dehydrogenase deficiency in Estonia"
12.05 – 12.20	T-M. Laht (Estonia): Dairy products - use in diets for children with metabolic disorders.
12.25 – 13.25	***Lunch***
13.25 – 13.40	A.Matulevičienė (Vilnius): Case of Duchenne muscular dystrophy and succinic semialdehyde dehydrogenase deficiency.
13.40 - 13.55	K. Krabbi (Estonia) :"The prevalence of mucopolysaccharidoses in Estonia"
13.55 – 14.30	Unknown cases: 3-4 presentations form each country.
14.30	Closing remarks, discussions, future plans.