Meeting of Baltic metabolic specialists

14.11.2007. - 15.11.2007.
Venue: CH "Gaiļezers"
Juglas str. 20

Preliminary Program

14th of November, 2007

11.00 Welcome by Ass.Prof. R. Lugovska and dr. Z.Krumina
11.15 Congenital disorders of glycolisation. RA. Wevers, Professor in Clinical Chemistry, Nijmegen, The Netherlands (45 min)
12.00 Mitochondrial diseases. K. Joost, K. Ounap, Tartu, Estonia,

12.30 – 13.00 coffee break
13.00 Neurotransmitter diseases. Prof. RA. Wevers (45 min)
14.00 Creatine transporter defect –screening among XMR patients. K.Ounap, Tartu, Estonia
14.15 Laboratory diagnostic of creatine metabolism. K.Kall, Tallin, Estonia
14.30 A case of Fanconi – Bickel syndrome due to previously not described homozygous 1 bp deletion in SLC2A2 gene. B.Tumiene, Vilnius, Lithuania
14.45 Speaker from Lithuania will be specified
15.00-16.00 Lunch
16.00 Experience of Lithuanian paediatricians in diagnosing and treating IME in patients with acute encephalopathies of unknown reason.B. Skerliene, Vilnius, Lithuania (30 min)
16.30 The most common inherited liver diseases. Dr.B.Lace, Latvia (30 min)

Social dinner

15th of November, 2007
9.00 "From bed to the bench" – Prof. RA. Wevers (45 min)
10.00 Laboratory practice in Latvia – P.Vēvere (Latvia) 30 min
10.30 Lysinuric protein intolerance: case report – dr. Z.Krūmiņa (Latvia) 15 min
10.45-11.15 coffee break
11.15 Unsolved cases (3-4 cases, presentation time 10 min) Estonia
13.00-14.00 Lunch
14.00 Unsolved cases (3-4 cases, presentation time 10 min) Lithuania
15.00 Unsolved cases (3-4 cases, presentation time 10 min) Latvia
16.00 Baltic metabolic group meeting – web site, the representative in SSIEM etc. Closing remarks dr.Z.Krūmiņa, prof.K.Ounap