

Rare Disease Research Forum- Challenges and Solutions

Preliminary programme

9.00-9.20	Introduction	Désirée Gavhed/ Annika Larsson
<u>Session 1. RD Research</u>		
9.20-9.40	Genetic research	Magnus Nordenskjöld
9.40-10.00	Metabolic disorders	Anna Wedell
10.00-10.20	Hematological disorders	Jan-Inge Henter
10.20-10.50 Break		
10.50-11.10	Endocrinological disorders	Anna Nordenström
11.10-11.30	Neuromuscular disorders	Thomas Sejersen
11.30-11.50	Patient organizations/foundations and research	Barbro Westerholm
11.50-12.10	Rare diseases in the primary care	Kjartan Thorarinsson/ Jonas Lindbäck
12.10-13.30 LUNCH		
<u>Session 2. Research support and services</u>		
13.30-13.45	Core facilities at KI	Nancy Pedersen
13.45-14.00	Biobank	Mark Divers
14.00-14.15	Support and needs of industry	Stephen James, Swedish Orphan Biovitrum
14.15-14.30	Support from EMA (COMP)	Kerstin Westermark
14.30-14.45	Making the most of patient registries	Elizabeth Hernberg-Ståhl, Late Phase Solutions Europe
14.45-15.15 Break		
<u>Session 3. Panel discussion</u>		
15.15-16.00	Panel Discussion	