PARIS MEETING PROGRAM (December 5th)			
9:00	9:15	Marco and Zeynep	
SESSION 1	- METABERN (25' + 5' discussion)		
9:15	9:45	Information on the Learning&Training Programme of the MetabERN; Research initiatives of the subnetwork Peroxisomal (Peroxisomal Biogenesis defects) and Lipid related Disorders (Smith Lemli Opitz syndrome)	Francois Eyskens
9:45	10:15	Beyond the Genome: (Cellular) Metabolomics and Proteomics as Functional Readouts of Genetic Variation	Dr. Irene Körver-Keularts
10:15	10:45	Making Sense of Genetic Variants: The Role of Fluxomics in Functional Diagnostics	Dr. Marek Noga
10:45	11:00	Metabolic disorders around the EU & Norway	Sally Ann Lynch
11:00	12:00	Brunch	
SESSION 2A: COLLABORATIVE RESEARCH TALKS (10' + 5' discussion)			
12:00	12:15	Shprintzen-Goldberg syndrome: long term follow-up of the cardiovascular features in an international cohort of 29 patients	Faivre Laurence
12:15	12:30	Impact and natural history of metabolic and endocrine complications in a large RATARS cohort	Francesca Clementina Radio
12:30	12:45	Novel variants and DNA methylation alteration on pathologic tissues from a cohort of Italian patients with spinal dysraphisms	Valeria Capra
12:45	13:00	Delineation of the adult phenotype of Nicolaides-Baraitser, Coffin-Siris, Costello and CFC syndrome	Dagmar Wieczorek
13:00	13:30	Break	
SESSION 2B: COLLABORATIVE RESEARCH TALKS (10' + 5' discussion)			
13:30	13:45	Functional stratification of YWHAZ variants explains distinct neurodevelopmental disorders	Marco Tartaglia
13:45	14:00	RYR2-catecholaminergic polymorphic ventricular tachycardia (CPVT) and neurodevelopmental disorders	Luisa MARSILI
14:00	14:15	Vitamin D and calcium metabolism in adults with Williams-Beuren syndrome	Laura de Graaff
14:15	14:30	Expanding the phenotypic and genotyping spectrum of NAA15 related disorder	Alessandra Renieri
14:30	15:00	Closing remarks	