13:30-13:40 Introduction (Thomas and Nina)

13:40-14:10 Ethical dilemmas in the treatment of SMA (Thomas + discussion all)

14:10-14:40 Small molecules and genetic and epigenetic modifiers in the treatment of neuromuscular disorders (Nina Barisic)

14:40-15:00 P4HTM gene associated with neuromuscular symptoms and developmental delay or intellectual disability (Johanna Uusimaa)

15:00-15:15 COFFEE BREAK

15:15-15:30 SMA and DMD clinical and epidemiological overview in Latvia (Signe Setiere)

15:30-16:00 Transition to adult care (Thomas + discussion all)

16.00-16.15 Next generation sequencing in patients presenting with double phenotypes myopathy and cardiomyopathy (Thomas)

16:15-16:30 Summarizing remarks/discussion (Thomas, Nina)