# EPNS Research Meeting: Alicante: 26-27 October 2018

**Program Neuromuscular working group, Friday 26 October, 13:30-16:30:**

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)