## YPSP01-02 - EFFECTIVENESS, QUALITY OF LIFE AND CHANGES IN BURDEN OF DISEASE IN CHILDREN AND ADOLESCENTS WITH ADHD TRANSITIONING TO OROS MPH

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**Objective:** To explore effectiveness, quality of life outcomes, burden of disease in children and adolescents with ADHD transitioning from ER MPH or Atomoxetine onto PR OROS MPH.

**Methods:** Twelve week open label study including 224 patients (aged 6-18) with ADHD (ICD-10) transitioning from ER MPH (N=180), Atomoxetine (N=42) or both (N=2) onto flexibly dosed PR OROS MPH. Starting dose was based on clinical judgement. Assessments included <u>C</u>hildren's <u>G</u>lobal <u>Assessment Scale</u>, IOWA <u>Conners' parent rating scale</u>, quality of life (ILC), and open question related to late afternoon or evening activities.

**Results:** 224 patients (85.3% boys, median age 11 yrs) were documented. 81% completed the study. Median starting and final dose of PR OROS MPH was 36 mg/d. Mean C-GAS improved from 58  $\pm$  15 (previous ER MPH group) and 54  $\pm$  11 (previous ATX group) to 71 $\pm$ 16 (12 $\pm$ 15; p< 0,01) and 64 $\pm$ 18 (9 $\pm$ 16; p< 0.001), respectively.

Playing with other children, doing household chores, behavior towards visitors and doing homework were improved after switching from ATX to OROS MPH (all p < 0.05), but not for going to bed (p = 0.57). All items improved in the previous ER MPH group (p < 0.0001). Symptoms measured on IOWA Connor´s rating scale as well as burden of disease (ILC) improved in children, adolescents as well as their care givers (p < 0.005).

**Conclusion:** Transition from ER MPH and ATX onto PR OROS MPH was associated with improved functionality, social interaction and decreased burden of disease in children and adolescents with ADHD.