



ORAL PRESENTATION

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# Early markers of the metabolic syndrome in children born post-term

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From 7th APPES Biennial Scientific Meeting  
Nusa Dua, Bali. 14-17 November 2012

We recently showed from a Swedish cohort that nearly half of boys born post-term ( $\geq 42$  weeks gestation) were overweight or obese at 16 years of age. We hypothesized that post-term children would display features of insulin resistance and the metabolic syndrome even in their pre-pubertal years.

90 healthy pre-pubertal children aged 4–11 years, with birth weight appropriate-for-gestational-age were studied: 36 children born post-term (18 boys and 18 girls) and 54 children (36 boys and 18 girls) born at term (38–40 weeks). Insulin sensitivity was measured using Bergman's minimal model. Other assessments included fasting lipid and hormonal profiles, body composition using whole-body dual-energy x-ray absorptiometry, and 24-hour ambulatory blood pressure monitoring.

Insulin sensitivity was reduced in post-term children ( $8.44 \pm 0.74$  vs  $13.55 \pm 0.89 \times 10^{-4} \cdot \text{min}^{-1} \cdot (\text{mU/L})$ ;  $p < 0.0001$ ). Post-term children had an adverse lipid profile, with higher total cholesterol ( $4.26 \pm 0.17$  vs  $3.92 \pm 0.11 \text{ mmol/l}$ ;  $p = 0.023$ ) and LDL ( $2.51 \pm 0.13$  vs  $2.25 \pm 0.08 \text{ mmol/l}$ ;  $p = 0.016$ ) concentrations. Further changes suggestive of the metabolic syndrome among post-term children included an increased android to gynoid fat ratio ( $0.74 \pm 0.03$  vs  $0.61 \pm 0.02$ ;  $p = 0.026$ ), and a reduction in the normal nocturnal systolic blood pressure dip ( $8.2 \pm 1.0$  vs  $13.8 \pm 1.0$ ;  $p = 0.016$ ). Post-term children also had higher serum leptin ( $7.18 \pm 0.91$  vs  $3.67 \pm 0.41 \text{ ng/ml}$ ;  $p = 0.011$ ), lower adiponectin ( $8226 \pm 693$  vs  $10536 \pm 556 \text{ ng/ml}$ ;  $p = 0.046$ ), and lower IGFBP1 ( $9.56 \pm 1.06$  vs  $18.03 \pm 1.59 \text{ ng/ml}$ ;  $p = 0.032$ ) concentrations.

Our study shows for the first time that post-term children have early features of the metabolic syndrome,

including reduced insulin sensitivity, adverse lipid profile, and increased abdominal adiposity.

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Published: 3 October 2013

doi:10.1186/1687-9856-2013-S1-O22

Cite this article as: Ayyavoo et al.: Early markers of the metabolic syndrome in children born post-term. *International Journal of Pediatric Endocrinology* 2013 2013(Suppl 1):O22.

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