(CARS) while administering the WPSSI-III. GARS and CARS scores were collapsed into autistic versus non-autistic categories and used to analyse the data.

**Results** Parent’s GARS identified a significantly higher number of children (38%) with ASD than did the examiner’s ratings (CARS, 10%). GARS’ groups did not differ significantly in NICU length of stay, incidence of sepsis, or severe IVH. At 6, 15, and 24 months the ASD group scored significantly lower on the BSID-III and at four years they had a lower mean IQ (89 v. 105, p < 0.0001). Executive functioning for the ASD group on the Brief-P was poor (70.03 v. 47.4, p < 0.0001).

**Conclusion** Parent’s perception of NICU morbidities and cognitive delays affect the identification of ASD in extremely preterm infants.

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**Oral abstracts**

**THE ROLE OF FTO RS17817449 AND RS 9939609 SNP (SINGLE NUCLEOTIDE POLYMORPHISM) IN CHILDREN’ OBESE**

**O-050**

Obesity is a problem with great importance and certain actuality in the field of paediatric pathology because of its increased incidence in children and its complications. Fat mass and obesity associated (FTO) genes are related with weight gain and obesity during childhood, being correlated with body mass index (BMI) and leptin levels.

The aim of our study was to establish the role of the polymorphisms of rs9939609 and rs17817449 FTO genes in determining obesity in a child population from Romania.

**Methods** We assessed 225 hospitalised children in a tertiary emergency paediatric hospital divided into: control group - 110 patients with normal nutritional status and obese group -105 patients. The two groups underwent the evaluation of FTO rs17817449 and rs 9939609 SNP, and the measurement of anthropometric parameters [body mass index (BMI), middle upper arm circumference (MUAC), tricipital skinfold thickness (TST)].

**Results** We observed that the AA genotype of R9939609 SNP gene, is more frequent in obese group [p = 0.01, OR 2.778 (95% CI 1.245–6.201)] and the A alleles is heigher in obese group [p = 0, 0148, OR 1.692 95%]. According to the RS17817449 gene mutation no association was found. We observed correlations in AA genotype, between BMI and MUAC (p = 0.0011) and TST, while A allele carriers was associated only with MUAC.

**Conclusion** Rs9939609 SNP AA genotype and A allele carriers are associated with obesity in children, and are both correlated with BMI and MUAC, while AA genotype is also correlated with TST.

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**25-HYDROXY VIT D, ADIPONECTIN LEVELS AND CARDIOMETABOLIC RISK FACTORS IN A SAMPLE OF OBESE CHILDREN**

**O-052**

Association between vitamin D, adiponectin and obesity is a matter of debate, as they play important role in linking obesity with different cardiometabolic risk factors.

**Objectives** Evaluation of the association between metabolic risk factors with both adiponectin and vitamin D levels and that between adiponectin and vitamin D among obese Egyptian children.

**Subjects and methods** This case-control cross sectional study consisted of 65 obese and 30 healthy children, aged 8–11 years. 25-Hydroxy vitamin D [25(OH) D] level, serum adiponectin, total cholesterol (TC), triglycerides (TG), high-density lipoprotein-cholesterol (HDL-C) and low-density lipoprotein-cholesterol (LDL-C) were measured.

**Results** The mean 25(OH) D levels in the obese and control groups were 29.9 ± 10.3 and 39.7 ± 12.7 ng/mL, respectively (p < 0.0001). The mean 25(OH)D and adiponectin levels in the obese were lower than that in the control group (p < 0.0001). 25(OH)D were inversely correlated with body mass index (BMI), triglyceride, total cholesterol and LDL-cholesterol (LDL-C), while adiponectin level were inversely correlated with