Biochemistry, hormones and adipocytokines in prepubertal children born with IUGR evoke metabolic, hepatic and renal derangements

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Children born with IUGR develop features of the metabolic syndrome and exhibit deranged markers of hepatorenal physiology. Metabolic and hepatorenal biochemistry and the rs9939609 FTO polymorphism were investigated in prepubertal children born with IUGR. Ninety-eight prepubertal children (46 IUGR and 52 AGA), subdivided in <5 years and >5 years old groups were included. Anthropometry; creatinine, eGFR, urea, AST, ALT, triglycerides, uric acid, total cholesterol, HDL-c, LDL-c, glucose, C-peptide, insulin and glucagon z-scores; HOMA-IR; leptin and adiponectin concentrations; rs9939609 FTO polymorphism frequency were measured. In males, weight and ALT were higher and adiponectin was lower, in IUGR <5 years; C-peptide, insulin and leptin were higher in IUGR >5 years; C-peptide was higher in all IUGR, than the respective AGA. In females, creatinine and triglycerides were higher in IUGR <5 years old; creatinine was higher and eGFR was lower in all IUGR, than the respective AGA. In males and females, creatinine was higher in all IUGR, than the respective AGA; C-peptide, insulin and HOMA-IR were lower, and AST was higher in IUGR <5 than in IUGR >5 years old. FTO rs9939609 frequency did not differ between IUGR and AGA. In conclusion prepubertal males born with IUGR increased weight, insulin and leptin and decreased adiponectin, as compared to males born AGA, emerge as early metabolic syndrome characteristics. The concentrations of these hormones do not differ between prepubertal males and females born with IUGR. Weight control, healthy nutrition and physical exercise should be recommended to these children. The deranged renal (particularly evident in females below the age of 5) and liver biochemistry in prepubertal children born with IUGR suggests that hepatorenal derangements might commence in utero. Regular checkup of biochemical and lipid profile is recommended for all children born with IUGR.

In intrauterine growth restriction (IUGR) fetus fails to achieve its growth potential due to genetic or environmental factors in utero. Intrauterine growth restriction is defined as estimated birth weight below the 3rd centile. The term small for gestational age (SGA) refers to an infant whose birth weight is below the 10th centile for the appropriate gestational age. This clinical problem affects approximately 8% of pregnancies and is associated with perinatal mortality and morbidity. Several risk factors of maternal, fetal and placental origin have been...
possibly related to adrenal steroid hormones increase in adrenarche. As previously reported, the concentrations 5 years represents the lower limit of initiation of adrenarche. Thus, it was chosen to discriminate for differences old; males born IUGR born AGA divided in 8 groups according to their gender and age (males born IUGR female ratio between the IUGR and the AGA groups did not differ significantly. Both IUGR and AGA were years in both male and female children, after which a gradual increase was observed. The Institutional Review of the adrenal androgen DHEAS in peripheral circulation were maintained at a minimum level until the age of 5 years in both male and female children, after which a gradual increase was observed. Preterm infants with extremely low birth weight exhibit markedly decreased glomerulogenesis. In low birthweight preterm infants, active glomerulogenesis occurs in postnatal period, but it does not suffice to compensate the decrease of the number of nephrons in utero. Children born with IUGR are reported to have fewer nephrons but the same glomerular volume, whereas children with birth weight less than 2.5 kg have fewer nephrons and larger glomerular volume than children of higher birth weight. Preterm infants with extremely low birth weight exhibit markedly decreased glomerulogenesis. In low birthweight preterm infants, active glomerulogenesis occurs in postnatal period, but it does not suffice to compensate the decrease of the number of nephrons in utero. Children born with IUGR are reported to have fewer nephrons but the same glomerular volume, whereas children with birth weight less than 2.5 kg have fewer nephrons and larger glomerular volume than children of higher birth weight. 

Intrauterine growth restriction, as originally been suggested in the “thrifty phenotype” hypothesis by Barker et al., has been causally associated with the development of metabolic syndrome in adulthood and with long-term metabolic sequelae such as obesity, impaired glucose tolerance, diabetes mellitus type 2 (T2DM), hypertension, dyslipidemia, cardiovascular disease and polycystic ovary syndrome (PCOS). In a rat model, catch-up growth in IUGR was associated with the development of an insulin resistant phenotype of adipose tissue which in its turn contributes to the development of obesity. Girls born with low birth weight, followed by postnatal catch up growth, exhibit increased visceral fat and insulin resistance in pre-school age. Young adults born SGA demonstrate hyperinsulinemia. Furthermore, animal studies demonstrated the association between low birth weight and reduced number of nephrons. Preterm infants with extremely low birth weight exhibit markedly decreased glomerulogenesis. In low birthweight preterm infants, active glomerulogenesis occurs in postnatal period, but it does not suffice to compensate the decrease of the number of nephrons in utero. Children born with IUGR are reported to have fewer nephrons but the same glomerular volume, whereas children with birth weight less than 2.5 kg have fewer nephrons and larger glomerular volume than children of higher birth weight. 

Intrauterine growth restriction has been associated with the development of obesity in the future. Obesity in childhood and adulthood has emerged as one of the most serious public health concerns. The rs9939609 polymorphism of the human fat mass and obesity associated (FTO) gene was the first robust identification of a common gene variant to be associated with increased body mass index (BMI) and obesity. In other studies, FTO rs9939609 polymorphism was associated with insulin resistance, T2DM, metabolic syndrome and cardiovascular disease. 

To investigate the early development of features of metabolic syndrome as well as the biochemical, hormonal and adipokines profiles in prepubertal children born with IUGR a cohort of prepubertal children born with IUGR was studied cross-sectionally for anthropometric, biochemical, hormonal and adipokines parameters in comparison with a cohort of prepubertal healthy children born appropriate for gestational age (AGA). The possible association of the rs9939609 polymorphism of FTO gene with the development of obesity was also investigated.

**Subjects and Methods**

**Subjects.** Ninety-eight healthy Caucasian children from a Pediatric Endocrinology Center were enrolled in this cross-sectional study [46 (17 male, 29 female; median age: 3.750 years) born full-term IUGR and 52 (26 male, 26 female; median age: 5.125 years) born full-term AGA]. Both IUGR and AGA groups were homogenous by age when a statistical comparison of the means with Factors ANOVA followed by an LSD post-hoc test was performed (Table 1). Children were characterized as IUGR or AGA when their birth weight was below the 3rd customized centile or between the 10th and 90th customized centile for gestational age, respectively. Children between the 3rd and 10th customized centile were excluded. The classification of IUGR and AGA was done according to the Fenton 2013 growth charts.

All participants were prepubertal (Tanner I) and of normal weight (Table 2). The male to female ratio for all subjects taken as a whole was 43/55 = 0.78; for the children born with IUGR was 17/29 = 0.58 and for the children born AGA was 26/26 = 1. When these figures were tabulated in a 2 × 2 contingency table, the calculated Pearson’s chi-square test was 1.686 with a p-value 0.194, which is not significant at p < 0.05, indicating that the male to female ratio between the IUGR and the AGA groups did not differ significantly. Both IUGR and AGA were divided in 8 groups according to their gender and age (males born IUGR < 5 years old; males born AGA < 5 years old; males born IUGR > 5 years old; males born AGA > 5 years old; females born IUGR < 5 years old; females born AGA < 5 years old; females born IUGR > 5 years old; females born AGA > 5 years old). The cut-off point of 5 years represents the lower limit of initiation of adrenarche. Thus, it was chosen to discriminate for differences possibly related to adrenal steroid hormones increase in adrenarche. As previously reported, the concentrations of the adrenal androgen DHEAS in peripheral circulation were maintained at a minimum level until the age of 5 years in both male and female children, after which a gradual increase was observed. The Institutional Review Board of the National and Kapodistrian University of Athens approved the study which was conducted in accordance with the Declaration of Helsinki on human experimentation as revised in 1996. Informed written consent was obtained from the parent/guardians of each child.

|                | Male (n = 43) | Female (n = 55) |
|----------------|--------------|-----------------|
| Mean Age ± SE  | 5.20 ± 0.46  | 4.44 ± 0.38     |
| Median Age ± SD| 5.25 ± 3.03  | 3.67 ± 2.83     |

Table 1. Comparison of age between male and female children born IUGR and AGA at the observation time. Statistical comparisons were performed with multiple factors ANOVA followed by an LSD post-hoc test for the means and a non-parametric median test for the medians. Neither analysis revealed any statistical differences among the groups.
determination, while the remaining serum samples were stored frozen in multiple aliquots (at −80 °C) for further analyses. Serum samples were thawed once before analysis and were protected from light and auto-oxidation. All assays were performed in duplicate and the mean value was recorded.

### Assays

Biochemical parameters including serum glucose, total cholesterol, high density lipoprotein HDL-and low density lipoprotein LDL- cholesterol (HDLC and LDLc, respectively), triglycerides (TG), uric acid, creatinine, urea, aspartate aminotransferase (AST), alanine aminotransferase (ALT) were determined by using the Siemens Advia 1800 Clinical Chemistry Analyzer (Siemens Healthcare Diagnostics, Tarrytown, NY, USA).

Glomerular Filtration Rate (GFR) was estimated according to the Bedside Schwartz formula (2009): eGFR = 0.413 × height (cm)/creatinine (mg/dL).39

Serum insulin concentrations were measured with an electrochemiluminescence immunoassay using the automated analyser Cobas e411 and the Elecsys Insulin Kit (Roche Diagnostics, Basel, CH); serum C-peptide concentrations were measured with a solid phase two-site chemiluminescent immunometric assay using a chemiluminescence autoanalyzer Immulite 2000 (Siemens Healthcare Diagnostics, UK); serum glucagon concentrations were measured with a radioimmunoassay kit (Euro-Diagnostica, Malmo, Sweden); serum leptin concentrations were determined by using a Milliplex Map Human Metabolic Panel (Millipore, USA), which is based on the Luminex xMAP technology; serum adiponectin concentrations were determined by using an enzyme-linked immunoabsorbent assay (ELISA) purchased from Orgenium Laboratories, Finland. The intra- and inter-assay coefficients of variation (CV) and the sensitivity limits of these assays were, respectively: 2.0%, 2.8% and 0.2 μU/mL; 3.5%, 6.2% and 0.09 ng/ml; 2.5%, 8.8% and 3 pmol/L; <7%, <10% and 0.184 ng/ml; <10%, <12% and 0.185 ng/ml.

### Indices of carbohydrate metabolism

Insulin resistance was estimated by the homeostasis model assessment (HOMA-IR) = Glucose × Insulin/405 using fasting concentrations (glucose and insulin in mg/dl and mU/L, respectively).30

Glucose-to-insulin ratio (G/I) was calculated using fasting concentrations (glucose and insulin in mg/dl and mU/L, respectively).31

The quantitative insulin sensitivity check index (QUICKI) was calculated as follows: 1/[log(fasting insulin μU/mL) + log(fasting glucose mg/dl)].32

β-cell secretion of insulin was estimated by the HOMA-β = [(360 × Insulin)/(Glucose-63)]% using fasting concentrations (glucose and insulin in mg/dl and mU/L, respectively).33

### FTO Genotyping

Genotyping of FTO was performed as previously described.34 DNA was extracted from whole blood samples using the Qiamp Blood Kit (Qiagen, Germany) and then it was stored at −20 °C until analysis. The FTO single nucleotide polymorphism (SNP) (rs9939609) was genotyped by polymerase chain reaction (PCR) and restriction fragment length polymorphism analyses. Genomic deoxyribonucleic acid (DNA) (20 ng) was incubated in a 10 μl solution containing 1 X NH4 buffer, 2.5 mmol/L Mg, 200 μmol/L each deoxyribose nucleoside triphosphate (dNTP), 20 pmol forward (5-AACTGGCTCTTTGAAATGAGATCGA-3) and reverse (5-AGAGTACAGAGCTATCAAAGTGCTAGAC-3) oligonucleotide primers, and 0.5 U Taq DNA polymerase (Bioline Ltd., London, UK). The PCR mix was incubated at 94 °C for 5 min followed by 20 cycles of 94 °C for 45 sec, 61 °C for 45 sec, and 72 °C for 45 sec. After this, the PCR mix was incubated for 15 cycles of 94 °C for 45 sec, 51 °C for 45 sec, and 72 °C for 45 sec, followed by a final incubation at 72 °C for 10 min. Then it was incubated at 37 °C for 16 h with 2 U Scal (New England Biolabs, Hitchin, UK). Upon running the final products

### Table 2

|                          | IUGR(n = 46) | AGA(n = 52) |
|--------------------------|-------------|-------------|
| Birth weight(g)          | 2272.56 ± 60.03a | 2965.10 ± 57.09 |
| Birth length(cm)         | 46.61 ± 0.46a  | 50.03 ± 0.25  |
| Waist/hip ratio          | 0.90 ± 0.01   | 0.91 ± 0.01   |
| Weight z-score           | 0.39 ± 0.21   | 0.68 ± 0.18   |
| Height z-score           | 0.13 ± 0.20   | 0.54 ± 0.21   |
| BMI z-score              | 0.37 ± 0.25   | 0.56 ± 0.15   |

**Table 2.** Comparison of anthropometric parameters between children born IUGR and AGA at birth and at the observation time expressed as mean ± S.E. Statistical comparisons were performed with multiple factors ANOVA. * denotes statistically significant difference from the respective AGA group.

### Protocol

At the first visit, all children had an initial clinical examination by the same physician (E.S.). Their birth weight and length were recorded (Tables 2 and 3). Anthropometric measurements were recorded (the mean value of two measurements for each subject) and a venous blood sampling was performed at fasting. Body weight was measured to the nearest 0.1 kg (Beam Balance 710, Seca, Birmingham, UK), while children were wearing their underclothes. Height was measured to the nearest 0.1 cm using a stadiometer (Seca 208, Hanover, MD), while children were barefoot. BMI (in kilograms per square meter) was calculated. Weight, height and BMI z-scores were calculated from the respective data for the Greek population (Tables 2 and 3)28. Maximum hip and waist circumferences were measured in cm in duplicate with a flexible tape and waist-to-hip ratio was calculated (Table 2). Blood was collected into tubes containing serum separating (SST)-gel and was centrifuged for serum separation. Serum was immediately analyzed after sample collection, for biochemical and hormonal parameters determination, while the remaining serum samples were stored frozen in multiple aliquots (at −80 °C) for further analyses. Serum samples were thawed once before analysis and were protected from light and auto-oxidation. All assays were performed in duplicate and the mean value was recorded.
Table 3. Comparison of anthropometric parameters between children born IUGR and AGA at birth (birth weight and length) and at the observation time (weight z-score, height z-score, BMI z-score) expressed as mean ± S.E. Statistical comparisons were performed with multiple factors ANOVA. * denotes statistically significant difference from the respective AGA group. # denotes statistically significant difference from the respective group aged >5 years old. † denotes statistically significant difference from the respective female group.

| Parameter         | Male (n = 43) | Female (n = 55) |
|-------------------|---------------|-----------------|
|                  | IUGR (n = 17) | AGA (n = 26)    | IUGR (n = 29) | AGA (n = 26) |
| Weight z-score    | 1.14 ± 0.41 ± | 0.70 ± 0.30     | -0.045 ± 0.18± | 0.65 ± 0.21  |
|                   | 0.31 ± 0.33   | 0.28 ± 0.25     | 0.42 ± 0.28    | 0.61 ± 0.19  |
| Height z-score    | 1.10 ± 0.41±  | 0.50 ± 0.23     | -0.058 ± 0.29  | 0.61 ± 0.19  |
| BMI z-score       | 1.10 ± 0.41±  | 0.50 ± 0.23     | -0.058 ± 0.29  | 0.61 ± 0.19  |

Mean weight and length were significantly lower in subjects born with IUGR than in those born AGA (p < 0.05) (Table 2). In female subjects mean weight z-score was significantly lower in those born with IUGR than in those born AGA (p < 0.05). In all female subjects mean estimated GFR value was significantly lower in subjects born with IUGR than in those born AGA (p < 0.05) (Fig. 1). Mean urea z-score was not significantly different among the studied groups (p > 0.05). In all female subjects mean AST z-score was significantly higher in those aged <5 years old than in those aged >5 years old and also than in subjects born AGA aged <5 years old (p < 0.05). In all female subjects

on a 3% agarose gel, the wild-type T allele produced a 182-bp band and the polymorphic A allele produced 154- and 28-bp bands.

**Statistical analysis.** Results are reported as mean ± standard error (SE) or median and interquartile range (25th–75th percentile). Statistical significance was set at p < 0.05. Standard deviations scores (z-scores) of biochemical and glucose metabolism parameters were calculated from the respective mean and SD values for age from data of our laboratory. Anthropometric measurements, biochemical parameters, glucose metabolism parameters and adipokines were compared between IUGR and AGA groups by employing multiple factors analysis of variance (ANOVA). Significant main effects were revealed by Fischer’s post-hoc test.

Comparison of the frequencies of the FTO genotypes in IUGR and AGA children for each polymorphism was performed with a 2 × 3 Pearson’s chi-square (3 genotypes for the 2 studied groups) test as previously described35. All polymorphisms were tested for Hardy-Weinberg-Equilibrium (HWE) 35. Allele frequencies comparison was also performed in the same way (wild-type T allele versus polymorphic A allele for the two studied groups) 35.

**Results**

**Anthropometry in children born with IUGR and in children born AGA.** Mean birth weight and mean birth length were significantly lower in subjects born with IUGR than in subjects born AGA (p < 0.05) (Table 2). In female subjects mean weight z-score was significantly lower in those born with IUGR than in those born AGA (p < 0.05). In subjects born with IUGR mean weight and BMI z-scores were significantly higher in males than in females (p < 0.05) (Table 3).

In male subjects aged <5 years old mean weight z-score was significantly higher in those born with IUGR than in those born AGA (p < 0.05). In male subjects born with IUGR mean BMI z-score was significantly lower in those aged <5 than in those aged >5 years old (p < 0.05). In female subjects aged <5 years old mean BMI z-score was significantly lower in those born with IUGR than in those born AGA (p < 0.05). In subjects born with IUGR and aged <5 years old mean height z-score was significantly higher in males than in females (p < 0.05) (Table 3).

**Biochemical parameters in children born with IUGR and in children born AGA.** The measurements of biochemical parameters in the studied groups at the observation time, expressed in z-scores, are presented in Table 4.

Mean creatinine z-score was significantly higher in subjects born with IUGR than in those born AGA (p < 0.05). In female subjects aged <5 years old mean creatinine z-score was significantly higher in subjects born with IUGR than in those born AGA (p < 0.05). In all female subjects mean estimated GFR value was significantly lower in subjects born with IUGR than in those born AGA (p < 0.05) (Fig. 1). Mean urea z-score was not significantly different among the studied groups (p > 0.05). In subjects born with IUGR mean AST z-score was significantly higher in those aged <5 years old than in those aged >5 years old and also than in subjects born AGA aged <5 years old (p < 0.05). In all female subjects
Table 4. Comparison of biochemical parameters between children born IUGR and AGA at the observation time expressed as mean ± S.E. Statistical comparisons were performed with multiple factors ANOVA. * denotes statistically significant difference from the respective AGA group. # denotes statistically significant difference from the respective group aged >5 years old. * denotes statistically significant difference from the respective female group.

|                  | Male (n=43) | AGA (n=26) | Female (n=55) |
|------------------|-------------|------------|---------------|
|                  | IUGR (n=17) | AGA (n=14) | IUGR (n=29)   | AGA (n=26) |
| Creatinine z-score | 0.33 ± 0.24* | −0.29 ± 0.21 | 0.36 ± 0.11* | −0.35 ± 0.22 |
| Urea z-score | 0.26 ± 0.19 | 0.04 ± 0.17 | −0.14 ± 0.22 | −0.06 ± 0.21 |
| AST z-score | 0.04 ± 0.25 | −0.09 ± 0.19 | 0.32 ± 0.20* | −0.31 ± 0.17 |
| ALT z-score | 0.55 ± 0.34* | 0.05 ± 0.21 | −0.13 ± 0.14 | −0.26 ± 0.17 |
| Triglycerides z-score | −0.31 ± 0.25 | −0.23 ± 0.15* | 0.49 ± 0.20* | −0.13 ± 0.18 |
| Uric acid z-score | 0.01 ± 0.25 | −0.08 ± 0.20 | 0.10 ± 0.19 | −0.13 ± 0.21 |
| Total choI z-score | −0.05 ± 0.22 | −0.15 ± 0.21 | 0.31 ± 0.17 | −0.17 ± 0.21 |
| HDL-c z-score | 0.32 ± 0.28 | 0.09 ± 0.18 | −0.04 ± 0.17 | −0.27 ± 0.20 |
| LDL-c z-score | −0.12 ± 0.18 | −0.16 ± 0.20 | 0.35 ± 0.18 | −0.16 ± 0.21 |

Below 5 years | Above 5 years | Below 5 years | Above 5 years |
|--------------|--------------|--------------|--------------|
| Creatinine z-score | 0.37 ± 0.34 | −0.22 ± 0.24 | 0.28 ± 0.37 | −0.34 ± 0.34 | 0.23 ± 0.13* | −0.70 ± 0.31 | 0.71 ± 0.08 | −0.02 ± 0.30 |
| Urea z-score | 0.39 ± 0.23 | 0.05 ± 0.31 | 0.11 ± 0.32 | 0.03 ± 0.18 | 0.01 ± 0.25 | −0.06 ± 0.28 | −0.59 ± 0.41 | −0.06 ± 0.33 |
| AST z-score | 0.53 ± 0.24* | −0.38 ± 0.23 | −0.46 ± 0.38 | 0.15 ± 0.29 | 0.61 ± 0.23* | −0.48 ± 0.24 | −0.42 ± 0.29 | −0.17 ± 0.24 |
| ALT z-score | 0.71 ± 0.27* | −0.36 ± 0.28 | 0.38 ± 0.65 | 0.37 ± 0.28* | 0.02 ± 0.16 | −0.13 ± 0.30 | −0.53 ± 0.23 | −0.37 ± 0.18 |
| Triglycerides z-score | −0.69 ± 0.13* | −0.08 ± 0.17 | 0.13 ± 0.47 | −0.38 ± 0.24 | 0.60 ± 0.27* | −0.36 ± 0.18 | 0.22 ± 0.21 | 0.09 ± 0.29 |
| Uric acid z-score | −0.28 ± 0.22 | 0.04 ± 0.33 | 0.31 ± 0.45 | −0.03 ± 0.26 | −0.20 ± 0.21* | 0.04 ± 0.35 | 0.86 ± 0.20* | −0.27 ± 0.24 |

Figure 1. Estimated Glomelural Filtration Rate (eGFR) in the IUGR (hatched bars) and AGA (white bars) groups. Significant differences were assessed by One-Way factors ANOVA (p < 0.05). * denotes statistically significant difference from the respective female group. # denotes statistically significant difference from the respective group aged >5 years old. * denotes statistically significant difference from the respective female group.

Mean AST z-score was significantly higher in those born with IUGR than in those born AGA (p < 0.05). The mean ALT z-score was significantly higher in male subjects born with IUGR and aged <5 years old than in their respective control AGA subjects (p < 0.05). Mean ALT z-score was significantly higher in male subjects born with IUGR than in female subjects born with IUGR (p < 0.05).

Mean total cholesterol, HDL-c and LDL-c z-scores were not significantly different among the studied groups (p < 0.05). In all female subjects mean TG z-score was significantly higher in those born with IUGR than in those born AGA (p < 0.05), while it was significantly higher in female subjects born with IUGR and aged <5 years old than in their respective control AGA subjects (p < 0.05). Mean TG z-score was significantly lower in male subjects born with IUGR and aged <5 years old than in their respective female subjects born with IUGR (p < 0.05).
In female subjects born with IUGR mean z-score of uric acid was significantly lower in those aged <5 than in those aged >5 years old (p < 0.05), while it was significantly higher in female subjects born with IUGR and aged >5 years old than in their respective control AGA subjects (p < 0.05).

**Glucose metabolism parameters and insulin resistance indexes in children born with IUGR and in children born AGA.** The measurements of glucose metabolism parameters of the studied groups at the observation time are presented in Table 5. Mean glucose z-score was significantly lower in male born with IUGR and aged <5 years old than in their respective control AGA and than in subjects born with IUGR and aged >5 years old (p < 0.05). Mean glucose z-score was significantly higher in male subjects born with IUGR and aged >5 years old than in their respective female subjects born with IUGR (p < 0.05).

Mean C-peptide z-score was significantly higher in all male subjects born with IUGR than in all male subjects born AGA (p < 0.05). Mean C-peptide and insulin z-scores as well as mean HOMA-IR value were significantly lower in male subjects born with IUGR and aged <5 years old than in male subjects born with IUGR and aged >5 years old (p < 0.05). The latter had mean C-peptide and insulin z-scores higher than in their respective male subjects born AGA as well as than in female subjects born with IUGR and aged >5 years old (p < 0.05).

Mean HOMA-β, QUICKI and glucose/insulin ratio values did not differ significantly among the studied groups (p < 0.05) (data not shown).

**Adipocytokines in children born with IUGR and in children born AGA.** Adipocytokines concentrations of the studied groups at the observation time are presented in Figs 2 and 3. Mean leptin concentration was significantly lower in male subjects born with IUGR and aged <5 years old than in male subjects born with IUGR and aged >5 years old (p < 0.05), while the latter had significantly higher mean leptin concentration than in their respective control male subjects born AGA (p < 0.05). Mean adiponectin concentration was significantly lower in male subjects born with IUGR and aged <5 years old than in their respective control male subjects born AGA (p < 0.05). Male subjects born with IUGR and aged >5 years old had significantly higher mean leptin/waist and mean leptin/(waist/hip) ratios than their respective control male subjects born AGA (p < 0.05), while these parameters were significantly lower in female subjects born with IUGR and aged >5 years old than in their respective control female subjects born AGA (p < 0.05).

**Genetic Analysis of FTO rs9939609 polymorphism frequencies in children born with IUGR and in children born AGA.** The frequencies of the genotypes of the rs9939609 polymorphism of the FTO gene in the AGA and IUGR groups are presented in Table 6. When chi-square analysis followed by Fisher’s post hoc was applied, the frequencies of the genotypes did not differ statistically between the IUGR and AGA groups (p < 0.05). When the frequencies of the rs9939609 polymorphism (T replaced by A) were tested for Hardy-Weinberg equilibrium, in the AGA and the IUGR groups, both were found to be in Hardy-Weinberg equilibrium (Table 7).

When the FTO allele frequencies in the IUGR and AGA groups were calculated in a 2 × 2 contingency table no statistically significant difference was revealed between the two groups, for the T and A alleles. The frequencies of the genotypes of the rs9939609 polymorphism were also tested for the presence of a dominant genetic model (A allele increases risk) in both IUGR and AGA groups. No statistically significant difference was revealed regarding the presence of the A allele. Finally, when the frequencies of the genotypes of the rs9939609 polymorphism were tested for the presence of a recessive genetic model (two A alleles increase risk) in both IUGR and AGA groups, no statistically significant difference was found.

Table 5. Comparison of glucose metabolism and hormone parameters between children born IUGR and AGA at the observation time expressed as mean ± S.E. Statistical comparisons were performed with multiple factors ANOVA. * denotes statistically significant difference from the respective AGA group, † denotes statistically significant difference from the respective group aged >5 years old. ‡ denotes statistically significant difference from the respective female group.
We found that birth weight and length were lower in all subjects born with IUGR compared to those born AGA, as expected\(^1,4\). Interestingly, males born with IUGR below 5 years old had higher weight z-score than their respective subjects born AGA, as well as their respective females born with IUGR. Of note, the latter did not differ in weight from their corresponding females born AGA, indicating that both males and females born with IUGR demonstrate weight catch-up before 5 years of age. However, in this study, males do so in a more prominent

### Table 6

| Polymorphism | Genotype | AGA (n = 52) | IUGR (n = 45) | χ² (P) | Fischer’s (Pₐ, Pₐ, N. of tables) |
|--------------|----------|--------------|---------------|--------|---------------------------------|
| rs9939609    | TT       | 0.21 (n = 11) | 0.33 (n = 15) | χ² = 2.6 | Pₐ = 0.271                      |
| rs9939609    | TA       | 0.56 (n = 29) | 0.53 (n = 24) | P = 0.27 | Pₐ = 0.271                      |
| rs9939609    | AA       | 0.23 (n = 12) | 0.13 (n = 6)  |        | N of tables evaluated = 513     |

**Discussion**

We found that birth weight and length were lower in all subjects born with IUGR compared to those born AGA, as expected\(^1,4\). Interestingly, males born with IUGR below 5 years old had higher weight z-score than their respective subjects born AGA, as well as their respective females born with IUGR. Of note, the latter did not differ in weight from their corresponding females born AGA, indicating that both males and females born with IUGR demonstrate weight catch-up before 5 years of age. However, in this study, males do so in a more prominent
Born AGA, albeit within normal range. Previous studies suggested that rapid catch-up growth in early postnatal children born SGA in their first and fourth year of age they found that the former experienced a greater weight gain than those born AGA, while genetic predisposition in leptin production by adipose tissue might alter leptin sensitivity and adiponectin concentrations to appear as well between female children born with IUGR and their mothers. The hyperinsulinemia observed at an early age suggests that the insulin resistance developing in children born with IUGR during adulthood might have its origins even before childhood, i.e. in utero. In the past we have shown that during normal pregnancy in non-obese, non-diabetic women the increase of the adipose tissue from the first trimester of pregnancy is accompanied by a significant progressive increase of maternal insulin sensitivity. Furthermore, in this study, leptin and adiponectin concentrations in males born with IUGR older and younger than 5 years old, respectively, were higher and lower, respectively, than their respective control AGA groups. Other authors reported lower concentrations of adiponectin in children born SGA as compared to children born AGA, while others suggested the development of leptin resistance in utero. Previously, we measured these adipocytokines in the serum of mothers of babies born with IUGR and babies born AGA. The postnatal maternal concentrations of leptin and adiponectin were higher and lower, respectively, in the mothers of IUGR compared to those of AGA babies. Thus, we suggested that children born with IUGR are associated to a specific high-leptin and low-adiponectin maternal profile. Also, in this study, leptin and adiponectin concentrations in males born with IUGR were higher and lower, respectively, than their control AGA groups. Note, leptin and adiponectin concentrations of males born with IUGR at the time of the study, were not statistically different from leptin and adiponectin concentrations of female children born with IUGR. It is possible that the small number of subjects included in this study did not allow the differences in leptin and adiponectin concentrations to appear as well between female children born with IUGR and those born AGA. The physiopathologic relationship of leptin with the development of insulin resistance together with the reported hyperleptinemia in children born with IUGR and their mothers suggest a possible genetic and/or epigenetic predisposition for the development of insulin resistance in children born with IUGR. In a rat model study the exposure to prenatal undernutrition led to the development of leptin resistance in adulthood, independently of diet-induced obesity. These findings suggest that prenatal conditions such as undernutrition can lead to future changes in energy metabolism and development of metabolic syndrome in adulthood through alterations in leptin sensitivity in utero, while genetic predisposition in leptin production by adipose tissue might also contribute to this phenomenon. Other authors, though, reported no difference in leptin concentrations and higher adiponectin concentrations in children born SGA as compared to children born AGA. The inclusion of children born SGA instead of those born with IUGR, as well as the differences in the number and age of the examined cases in that study compared to the present study might explain this discrepancy. Furthermore, we also reported a positive relationship between birth weight and the number of glomeruli, as well as a negative correlation between birth weight and glomerular volume in humans was also reported.

Table 7. Hardy-Weinberg Equilibrium analysis for children born IUGR and AGA.

| Observed | Expected | x² | P
|----------|----------|----|----
| AGA      |          |    |    
| TT       | 11       | 12.5| 0.697
| TA       | 29       | 26.0| 0.556
| AA       | 12       | 13.5| 0.404
| IUGR     |          |    |    
| TT       | 15       | 16.2| 0.556
| TA       | 24       | 21.6|         
| AA       | 6        | 7.2 | 0.456

Legend: TT = homozygous for the wild type allele, TA = heterozygous, AA = homozygous for the mutant allele, x² = chi-square, P = probability.

As compared to children born AGA beyond 12 months of age.

Regarding insulin secretion, in this study, all males born with IUGR as well as males born with IUGR older than 5 years old had higher C-peptide and insulin z-scores, respectively, than their respective control subjects born AGA. In addition, HOMA-IR index was found lower in males born with IUGR below 5 years old than in males born with IUGR above 5 years old. The hyperinsulinemia observed at an early age suggests that the insulin resistance developing in children born with IUGR during adulthood might have its origins even before childhood, i.e. in utero. In the past we have shown that during normal pregnancy in non-obese, non-diabetic women the increase of the adipose tissue from the first trimester of pregnancy is accompanied by a significant progressive increase of maternal insulin sensitivity. Furthermore, in this study, leptin and adiponectin concentrations in males born with IUGR older and younger than 5 years old, respectively, were higher and lower, respectively, than their respective control AGA groups. Other authors reported lower concentrations of adiponectin in children born SGA as compared to children born AGA, while others suggested the development of leptin resistance in utero. Previously, we measured these adipocytokines in the serum of mothers of babies born with IUGR and babies born AGA. The postnatal maternal concentrations of leptin and adiponectin were higher and lower, respectively, in the mothers of IUGR compared to those of AGA babies. Thus, we suggested that children born with IUGR are associated to a specific high-leptin and low-adiponectin maternal profile. Also, in this study, leptin and adiponectin concentrations in males born with IUGR were higher and lower, respectively, than their control AGA groups. Of note, leptin and adiponectin concentrations of males born with IUGR at the time of the study, were not statistically different from leptin and adiponectin concentrations of female children born with IUGR. It is possible that the small number of subjects included in this study did not allow the differences in leptin and adiponectin concentrations to appear as well between female children born with IUGR and those born AGA. The physiopathologic relationship of leptin with the development of insulin resistance together with the reported hyperleptinemia in children born with IUGR and their mothers suggest a possible genetic and/or epigenetic predisposition for the development of insulin resistance in children born with IUGR. In a rat model study the exposure to prenatal undernutrition led to the development of leptin resistance in adulthood, independently of diet-induced obesity. These findings suggest that prenatal conditions such as undernutrition can lead to future changes in energy metabolism and development of metabolic syndrome in adulthood through alterations in leptin sensitivity in utero, while genetic predisposition in leptin production by adipose tissue might also contribute to this phenomenon. Other authors, though, reported no difference in leptin concentrations and higher adiponectin concentrations in children born SGA as compared to children born AGA. The inclusion of children born SGA instead of those born with IUGR, as well as the differences in the number and age of the examined cases in that study compared to the present study might explain this discrepancy. Also, in that study no age group below 5 years old was included.

Furthermore, the higher and lower concentrations of creatinine and eGFR, respectively, in all female subjects born with IUGR and the higher concentrations of creatinine in the younger than 5 years old children born with IUGR might indicate that subjects born with IUGR might be at risk for developing renal derangement, particularly those of female sex. Previous studies reported reduced nephron number in children born with IUGR at birth. A positive relationship between birth weight and the number of glomeruli, as well as a negative correlation between birth weight and glomerular volume in humans was also reported.

Also, in this study, the higher concentrations of AST and ALT in the younger than 5 years old children born with IUGR than in their respective control subjects born AGA might indicate a developing liver dysfunction in subjects born with IUGR that starts in early childhood. In the past, liver biopsies associated IUGR with fatty liver in rats as well as with non-alcoholic fatty liver disease (NAFLD) in children. It is possible that children born...
with IUGR might be predisposed to the development of liver dysfunction and NAFLD. The higher values of AST and ALT reported in this study might be associated to an early development of fatty liver in children born with IUGR. In addition, in this study, TG and uric acid z-scores were, respectively, higher in all females born with IUGR and in females born with IUGR older than 5 years old than in their respective control groups of children born AGA.

Regarding the frequency of the FTO gene rs9939609 polymorphism we found no statistically significant difference among the groups of children born with IUGR and AGA. This polymorphism was in Hardy-Weinberg equilibrium in both groups.

In summary, we found that male children born with IUGR younger than 5 years old present with higher weight than their respective subjects born AGA. This finding associates enhanced weight increase to the “catch-up growth” phenomenon observed in children born with IUGR. Prepubertal boys born with IUGR, older and younger than 5 years old present, respectively, with increased concentrations of insulin and leptin and lower concentrations of adiponectin. This early observed hyperinsulinemia, hyperleptinemia and hypo adiponectinemia indicates that the known development of insulin and leptin resistance in adults born with IUGR manifests early in childhood. These findings should be corroborated to the increased prevalence of metabolic syndrome, including obesity, in adults born with IUGR. Creatinine is increased in all subjects born with IUGR, while hepatic enzymes AST and ALT are increased in all children born with IUGR younger than 5 years old and in male children born with IUGR younger than 5 years old, respectively. An adverse prenatal environment may induce early derangement of hepatic and renal function in IUGR fetuses, although the underlying mechanisms remain unclear.

In conclusion, this study highlights the importance of the early development of insulin and leptin resistance in children born with IUGR possibly associated to long-term metabolic consequences. Furthermore, it seems that there is an early development of hepatic and renal derangement in these children. More studies are needed to investigate the underlying mechanisms (possibly of epigenetic participation) which lead to changes in insulin and leptin sensitivity, as well as in derangements in the physiology of liver and kidneys. These events might occur in utero and in the early postnatal period. It is intriguing to investigate prospectively whether the suggested expression of features of the metabolic syndrome in these children during adulthood has its origins in prepubertal period.

References
1. Wollmann, H. A. Intrauterine growth restriction: definition and etiology. *Horm Res* 49, 1–6 (1998).
2. Clayton, P. E. et al. Management of the child born small for gestational age through to adulthood: a consensus statement of the International Societies of Pediatric Endocrinology and the Growth Hormone Research Society. *J Clin Endocrinol Metab* 92, 804–810 (2007).
3. Figuera, F. & Gratacos, E. Update on the diagnosis and classification of fetal growth restriction and proposal of a stage-based management protocol. *Fetal Diagn Ther* 36, 86–98 (2014).
4. Mandruzatto, G. et al. Intrauterine restriction (IUGR). *J Perinat Med* 36, 277–281 (2008).
5. Kanaka-Gantenbein, C., Mastorakos, G. & Chrousos, G. P. Endocrine-related causes and consequences of intrauterine growth retardation. *Ann NY Acad Sci* 997, 150–157 (2003).
6. Crume, T. L. et al. The long-term impact of intrauterine growth restriction in a diverse U.S. cohort of children: the EPOCH study. *Obesity (Silver Spring)* 22, 608–615 (2014).
7. Hales, C. N. Metabolic consequences of IUGR. *Pediatr Dev Pathol* 27, 755–777 (2003).
8. Rogers, I. The influence of birthweight and intrauterine environment on adiposity and fat distribution in later life. *Int J Obes Relat Metab Disord* 27, 92 (2003).
9. Nobili, V., Alisi, A., Panera, N. & Agostoni, C. Low birth weight and catch-up-growth associated with metabolic syndrome: a ten year systematic review. *Pediatr Endocrinol Rev* 6, 241–247 (2008).
10. Ibanez, L., Lopez-Bermejo, A., Diaz, M. & Marcos, M. V. Endocrinology and gynecology of girls and women with low birth weight. *Fetal Diagn Ther* 30, 243–249 (2011).
11. Berends, L. M., Fernandez-Twinn, D. S., Martin-Gronert, M. S., Cripps, R. L. & Otsanne, S. E. Catch-up growth following intrauterine growth-restriction programmes an insulin-resistant phenotype in adipose tissue. *Int J Obes (Lond)* 37, 1051–1057 (2013).
12. Leger, J. et al. Reduced final height and indications for insulin resistance in 20 year olds born small for gestational age: regional cohort study. *BMJ* 315, 341–347 (1997).
13. Vehaskari, V. M., Aixles, D. H. & Manning, I. Prenatal programming of adult hypertension in the rat. *Hypertension* 47, 889–894 (2006).
14. Mandruzzato, G. et al. Intrauterine growth restriction and blood pressure development in prepubertal children born with IUGR. *Arch Dis Child* 92, 345 (2007).
15. Lappalainen, T. et al. Association between the FTO rs9939609 polymorphism and the metabolic syndrome in a non-Caucasian multi-ethnic sample. *Cardiovasc Diabetol* 7, 5 (2008).
16. Faa, G. et al. Marked individual variability in renal maturation of preterm infants: lessons from autopsy. *J Matern Fetal Neonatal Med* 23(Suppl 3), 129–133 (2010).
17. Hinchliffe, S. A., Lynch, M. R., Sargent, P. H., Howard, C. V. & Van Velzen, D. The effect of intrauterine growth retardation on the development of renal nephrons. *Br J Obstet Gynaecol* 99, 296–301 (1992).
18. Manalich, R., Reyes, L., Herrera, M., Melendi, C. & Fundora, I. Relationship between weight at birth and the number and size of renal glomeruli in humans: a histomorphometric study. *Pediatr Dev Pathol* 27, 17–25 (2004).
19. Faa, G. et al. Marked individual variability in renal maturation of preterm infants: lessons from autopsy. *J Matern Fetal Neonatal Med* 23(Suppl 3), 129–133 (2010).
20. Weiss, R. et al. Obesity and the metabolic syndrome in children and adolescents. *N Engl J Med* 350, 2362–2374 (2004).
21. Frayling, T. M. et al. A common variant in the FTO gene is associated with body mass index and predisposes to childhood and adult obesity. *Science* 316, 889–894 (2007).
22. Freathy, R. M. et al. Common variation in the FTO gene alters diabetes-related metabolic traits to the extent expected given its effect on BMI. *Diabetes* 57, 1419–1426 (2008).
23. Al-Attar, S. A. et al. Association between the FTO rs9939609 polymorphism and the metabolic syndrome in a non-Caucasian multi-ethnic sample. *Cardiovasc Diabetol* 7, 5 (2008).
24. Lappalainen, T. et al. Association of the FTO gene variant (rs9939609) with cardiovascular disease in men with abnormal glucose metabolism—the Finnish Diabetes Prevention Study. *Nutr Metab Cardiovasc Dis* 21, 691–698 (2011).
25. Fenton, T. R. et al. Validating the weight gain of preterm infants between the reference growth curve of the fetus and the term infant. *BMC Pediatr* 13, 92 (2013).
26. Autsch, R. J. The physiology and biochemistry of adrenarche. *Endocr Dev* 20, 20–27 (2011).
Author Contributions
E.J.S. conceptualized and designed the study, performed initial data collection, contributed to the analysis and interpretation of data, drafted the initial manuscript, and approved the final manuscript as submitted. G.P. carried out the statistical analysis and interpretation of data, reviewed and revised the manuscript, and approved the final manuscript as submitted. G.V. contributed to the conceptualization and design of the study, the interpretation of data, reviewed and revised the manuscript, and approved the final manuscript as submitted. A.I.M. performed laboratory analyses, reviewed and revised the manuscript, and approved the final manuscript as submitted. A.I.M. performed laboratory analyses, reviewed and revised the manuscript, and approved the final manuscript as submitted. I.P. had the responsibility of all laboratory analyses, contributed to the interpretation of data, reviewed and revised the manuscript, and approved the final manuscript as submitted. D.H. followed pregnant mothers of IUGR, carried out the characterization of IUGR babies in utero, contributed to the conceptualization of the study, critically reviewed the manuscript, and approved the final manuscript as submitted. N.I. contributed to the conceptualization and the design of the study, followed-up the prepubertal IUGR children, contributed to the collection of the data, critically reviewed the manuscript, and approved the final manuscript as submitted. G.M. conceptualized and designed the study, supervised data collection and proper execution of the protocol and laboratory analyses, revised statistical analysis, interpreted statistical analysis, designed manuscript writing, critically reviewed the manuscript, and approved the final manuscript as submitted.

Additional Information
Competing Interests: The authors declare no competing interests.

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