Serum glucocorticoids and adiponectin associate with insulin resistance in children born small for gestational age

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Abstract

Objectives: Altered glucocorticoid activity is one possible mechanism linking fetal growth restriction with later insulin resistance (IR) and type 2 diabetes. We aimed to investigate whether serum glucocorticoid parameters are related to IR in children born small for gestational age (SGA).

Design: A total of 110 children (55 age- and gender-matched pairs born SGA or appropriate for gestational age (AGA) in a case–control setting) were studied at the mean age of 12.2 (±0.2) years.

Methods: Serum cortisol, corticosteroid-binding globulin (CBG), free cortisol index (FCI = cortisol/CBG), and glucocorticoid bioactivity (GBA, transactivation assay) were analyzed and related to serum adiponectin and insulin-like growth factor-binding protein 1 (IGFBP1) concentrations and homeostasis model assessment for IR (HOMA-IR) and QUICKI indices.

Results: In the pooled study population, GBA correlated well with cortisol and FCI (r = 0.681 and 0.586 respectively; P < 0.001 for both). Serum cortisol, CBG, FCI, GBA, HOMA-IR, or QUICKI did not differ between the SGA and AGA subjects, but the SGA children had lower body mass index (P = 0.005) and waist circumference (WC) (P = 0.001). The mean GBA in the highest GBA quartile was higher among the SGA subjects than among the AGA subjects (138.6 vs 96.4 nmol/l cortisol equivalents, P < 0.001). In the SGA children, GBA correlated positively with HOMA-IR (r = 0.522, P < 0.001) and inversely with adiponectin (r = −0.278, P = 0.042) (WC/height ratio adjustments), and in logistic regression analysis, higher GBA (odds ratio (OR) 1.3; P = 0.013), lower adiponectin (OR 1.4; P = 0.038), and lower IGFBP1 (OR 1.9; P = 0.010) associated independently with higher HOMA-IR.

Conclusions: These findings suggest that increased glucocorticoid activity and low serum adiponectin concentrations associate with IR in SGA children.

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transactivation assay (17), differs between children born small for gestational age (SGA) and appropriate for gestational age (AGA), and whether GBA is related to insulin resistance (IR) and serum adiponectin in SGA children. Serum adiponectin concentrations have been reported to decrease in obesity and insulin-resistant states (reviewed in (18)).

Subjects and methods

Definitions

SGA was defined as birth weight or length or ponderal index > 2 S.D. scores below the respective mean for the gestational age and sex (19). The ponderal index was calculated as (weight (g)/length^3 (cm)) × 100. AGA was defined as birth weight, birth length, and ponderal index ≥ −2 S.D. scores and ≤ 2 S.D. scores of the respective mean for the gestational age and sex. Homeostasis model assessment for IR (HOMA)-IR index was calculated as (\((\text{fasting insulin, } \mu \text{U/ml}) \times (\text{fasting glucose, mmol/l})/22.5\) and QUICKI as \(1/\log\) (fasting insulin, \(\mu \text{U/ml}\) + log (fasting glucose, mg/dl)) (see (20)). Free cortisol index (FCI) was calculated as serum cortisol (nmol/l)/corticosteroid-binding globulin (CBG) (mg/l) (21).

Subjects

The study population consisted of all full-term children who were born SGA at Kuopio University Hospital, Finland, between 1984 and 1986: 73 SGA children (70 singletons and three twins) were included in the study. Each SGA child had the next born full-term AGA child matched for sex as the control subject. At the age of 12 years, 55 SGA children (20 boys and 35 girls) and 55 AGA control subjects participated in this study. Nine mothers (16%) of the SGA subjects were diagnosed to have arterial hypertension (measured blood pressure more than 140/90 mmHg at least two times) during pregnancy; five of these (9.1% of the SGA mothers) had preeclampsia. None of the SGA subjects had chromosomal abnormalities, recognizable genetic diseases, or congenital anomalies. The mean age in both the SGA and AGA groups was 12.2 (s.d. 0.2) years. The study protocol was approved by the Research Ethics Committee of Kuopio University Hospital. Informed written consent was obtained from the child and the parents.

Table 1

<table>
<thead>
<tr>
<th>Variable</th>
<th>SGA (n = 55)</th>
<th>AGA (n = 55)</th>
<th>P value^a</th>
</tr>
</thead>
<tbody>
<tr>
<td>Birth weight (g)</td>
<td>2452 (314)</td>
<td>3455 (475)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Birth weight (SDS)</td>
<td>−2.44 (0.53)</td>
<td>−0.24 (0.99)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Birth length (cm)</td>
<td>46.2 (2.1)</td>
<td>50.4 (2.3)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Birth length (SDS)</td>
<td>−2.27 (0.95)</td>
<td>−0.08 (1.16)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Height 12 years (SDS)</td>
<td>−0.16 (0.90)</td>
<td>0.60 (1.0)</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Weight for height 12 years (%)</td>
<td>99 (17)</td>
<td>109 (22)</td>
<td>0.021</td>
</tr>
<tr>
<td>BMI 12 years (kg/m^2)</td>
<td>17.5 (2.9)</td>
<td>19.8 (4.2)</td>
<td>0.005</td>
</tr>
<tr>
<td>Waist circumference 12 years (cm)</td>
<td>62.5 (7.1)</td>
<td>69.6 (11.1)</td>
<td>0.001</td>
</tr>
<tr>
<td>Waist circumference/height 12 years</td>
<td>0.42 (0.05)</td>
<td>0.45 (0.06)</td>
<td>0.013</td>
</tr>
<tr>
<td>S-GBA (nmol/l cortisol equivalents)</td>
<td>78.0 (44.6)</td>
<td>64.9 (22.9)</td>
<td>0.127</td>
</tr>
<tr>
<td>S-cortisol (nmol/l)</td>
<td>292.5 (137.4)</td>
<td>272.1 (82.8)</td>
<td>0.624</td>
</tr>
<tr>
<td>S-CBG (mg/l)</td>
<td>64.0 (13.9)</td>
<td>65.3 (14.1)</td>
<td>0.795</td>
</tr>
<tr>
<td>Free cortisol index (nmol/mg)</td>
<td>4.68 (2.19)</td>
<td>4.27 (1.48)</td>
<td>0.456</td>
</tr>
<tr>
<td>HOMA-IR index</td>
<td>1.91 (0.74)</td>
<td>2.03 (0.96)</td>
<td>0.763</td>
</tr>
<tr>
<td>QUICKI</td>
<td>0.35 (0.02)</td>
<td>0.35 (0.02)</td>
<td>0.987</td>
</tr>
<tr>
<td>S-adiponectin (mg/l)</td>
<td>10.3 (5.5)</td>
<td>11.1 (6.7)</td>
<td>0.388</td>
</tr>
<tr>
<td>S-IGFBP1 (µg/l)</td>
<td>70.0 (34.8)</td>
<td>58.8 (30.3)</td>
<td>0.090</td>
</tr>
</tbody>
</table>

^aWilcoxon matched-pairs signed rank test for differences. Pubertal development stages (no difference between the study groups), S-cortisol concentrations (14) and IGFBP1 data (31) have been reported previously.
parameters of glucocorticoid and insulin metabolism in SGA and AGA subjects

In the combined SGA and AGA groups, serum GBA values correlated strongly with serum cortisol concentrations measured by the immunoassay ($r = 0.681$, $P < 0.001$) as well as with the FCI derived from the cortisol/CBG ratio ($r = 0.586$, $P < 0.001$). The means of serum GBA, cortisol, CBG, or FCI did not differ between the SGA and AGA children (Table 1), but the distributions of GBA, cortisol, CBG, or FCI were skewed to high values in the SGA group (demonstrated by higher S.D. values in the SGA group than in the AGA group, $P < 0.001$). Consequently, the mean GBA in the highest GBA quartile was significantly higher among the SGA subjects than among the AGA subjects (138.6 vs 96.4 nmol/l cortisol equivalents, $P < 0.001$). No significant differences were found in the means of HOMA-IR or QUICKI index between the SGA and AGA groups.
blood pressure levels did not differ when compared to Waist circumference/height (%)

K vs 0.43,
Pm

Low IGFBP1 (10

Low adiponectin level (mg/l) 0.37 0.038 1.4 1.0–2.0

High GBA level (10 nmol/l

Covariate Regression coefficient Significance Odds ratio Confidence interval (95%)

HOMA-IR index 2.89 (0.62) 1.58 (0.41) <0.001
QUICKI 0.33 (0.01) 0.36 (0.02) <0.001
S-adiponectin (mg/l) 7.2 (3.8) 11.4 (5.6) 0.002
S-IGFBP1 (µg/l) 44.4 (16.9) 78.7 (35.2) 0.001
S-GBA (nmol/l cortisol equivalents) 107.7 (60.9) 67.9 (32.7) 0.010
S-cortisol (nmol/l) 360.5 (177.1) 269.2 (114.5) 0.061
S-CBG (mg/l) 63.7 (14.7) 64.1 (13.7) 0.772
Free cortisol index (nmol/mg) 5.92 (3.04) 4.26 (1.66) 0.089
24-h systolic BP (mmHg) 120.4 (7.5) 116.4 (9.0) 0.024
BMI (kg/m²) 18.4 (2.8) 17.3 (2.9) 0.164
Waist circumference (cm) 63.6 (7.3) 62.1 (7.1) 0.434
Waist circumference/height 0.43 (0.05) 0.41 (0.05) 0.306

*The Mann–Whitney test for differences. Sex distribution and pubertal development according to the Tanner B/G staging did not differ between the HOMA-IR subgroups (P=0.483 and 0.218 respectively; χ² test). The ambulatory blood pressure values of the whole study population have been reported previously (32).

Factors associating with HOMA-IR index in the SGA children

The SGA subjects in the highest HOMA-IR quartile had significantly higher GBA, lower adiponectin and IGFBP1 concentrations, and higher 24-h systolic blood pressure than the SGA subjects in the lower HOMA-IR quartiles. Interestingly, serum cortisol concentrations (measured by the immunoassay) or FCI did not differ significantly between the high and lower HOMA-IR SGA subgroups (Table 2). When similar comparisons were performed in the control group, the AGA subjects in their highest HOMA-IR quartile had lower mean IGFBP1 (35.5 vs 65.8 µg/l, P=0.001) and higher BMI (23.9 vs 18.4 kg/m², P<0.001) and WC/height ratio (0.50 vs 0.43, P=0.001), but adiponectin (P=0.33), GBA (P=0.440), and other glucocorticoid parameters or blood pressure levels did not differ when compared to those of the AGA subjects in the lower HOMA-IR quartiles (detailed data not shown). For the multiple logistic regression analysis, the dependent variable HOMA-IR was dichotomized using the 75th percentile level as the cut-off point. In this analysis, higher serum GBA and lower adiponectin and IGFBP1 levels associated independently with higher HOMA-IR index in the SGA group (Table 3). An increase of 10 nmol/l cortisol equivalents in serum GBA increased by 1.3-fold the risk of higher HOMA-IR, while a decrease of 1 mg/l in serum adiponectin associated with a 1.4-fold and a decrease of 10 µg/l in serum IGFBP1 with a 1.9-fold risk of high HOMA-IR. The model was adjusted for WC/height ratio, sex, and pubertal stage, none of which was independently associated with high HOMA-IR. The model used explained 61.1% of the variation of HOMA-IR (Table 3).

Discussion

The present cohort study showed no differences in the mean serum GBA levels or other glucocorticoid parameters between the SGA children and their sex- and age-matched AGA controls. However, in the SGA group, serum GBA values exceeding the 75th percentile were significantly higher than those in the AGA group. Furthermore, the SGA subjects with the highest

Table 3 Factors predicting high homeostasis model assessment for insulin resistance (HOMA-IR) index in the small for gestational age children. Multiple logistic regression analysis (n=55)*.

<table>
<thead>
<tr>
<th>Covariate</th>
<th>Regression coefficient</th>
<th>Significance</th>
<th>Odds ratio</th>
<th>Confidence interval (95%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>High GBA level (10 nmol/l cortisol equivalents)</td>
<td>0.27</td>
<td>0.013</td>
<td>1.3</td>
<td>1.1–1.6</td>
</tr>
<tr>
<td>Low adiponectin level (mg/l)</td>
<td>0.37</td>
<td>0.038</td>
<td>1.4</td>
<td>1.0–2.0</td>
</tr>
<tr>
<td>Low IGFBP1 (10 µg/l)</td>
<td>0.67</td>
<td>0.010</td>
<td>1.9</td>
<td>1.2–3.1</td>
</tr>
<tr>
<td>Waist circumference/height (%)</td>
<td>-0.05</td>
<td>0.657</td>
<td>1.0</td>
<td>0.8–1.2</td>
</tr>
</tbody>
</table>

*Adjusted for sex (P=0.116) and pubertal stage (P=0.310). The model explained 61.1% of HOMA-IR index variation.
HOMA-IR index had higher serum GBA and lower adiponectin and IGFBP1 levels than those with lower HOMA-IR index. These findings suggest that there is an SGA subgroup with a more insulin-resistant state detectable already in childhood, and that increased glucocorticoid activity associates with it.

Previous studies have revealed increased plasma cortisol concentrations or their reactivity, IR, and type 2 diabetes in adult subjects born with low birth weight (5, 6, 10, 11). Cortisol excess in humans, e.g. in Cushing’s syndrome or during pharmacologic glucocorticoid treatment, may induce adverse effects including the components of the metabolic syndrome (hypertension, IR, dyslipidemia, and visceral obesity) (reviewed in (23)). Reynolds et al. (10) reported that men with low birth weight had enhanced responses of plasma cortisol to ACTH and increased total urinary cortisol metabolite excretion. In these subjects, features of metabolic syndrome such as raised blood pressure, glucose intolerance, and hypertriglyceridemia were associated with enhanced adrenal responsiveness to ACTH (10). The present study showed an association between GBA, measured by the transactivation assay, and IR indices in SGA children. Accordingly, intrauterine growth restriction may lead to alterations in glucocorticoid activity, which may contribute to IR.

Serum GBA associated more clearly than immunologically measured cortisol concentrations or FCI with HOMA-IR index in the SGA children. More than 90% of circulating cortisol in human serum is bound to proteins (CBG and albumin). FCI, calculated as a ratio of serum total cortisol to CBG, eliminates the effect of CBG variation in total cortisol values, and FCI has thus been shown to correlate with free serum cortisol, which is considered the biologically active hormone (21). Calculation of FCI does not eliminate the influence of variable albumin concentrations on total cortisol levels. We did not measure serum albumin concentration in our study, but we could speculate that the SGA subjects might have had slightly lower serum albumin levels as they were quite slim and their mean BMI was significantly lower than that of the AGA controls (Table 1, (22)). At least theoretically, variation in serum albumin concentrations could explain the better association of GBA than that of immunologically measured cortisol concentrations or FCI with HOMA-IR index in our study. We assume that our GBA measurement, based on a transactivation assay, estimates the circulating bioavailable cortisol concentrations better than the total cortisol measurement or even the calculated FCI. Thus, GBA measurements can give useful additional information (compared to immunological cortisol assays) on the endogenous activity of the HPA axis and cortisol metabolism. Apparently, GBA measurements have even greater value in conditions where the combined effects of exogenous synthetic glucocorticoids and endogenous cortisol on circulating glucocorticoid milieu are evaluated (24).

Adiponectin is an adipocyte-specific protein with insulin-sensitizing, anti-atherogenic, and anti-inflammatory properties. Circulating adiponectin is decreased in obesity, IR, and type 2 diabetes (reviewed in (18)). In our study, low adiponectin levels associated with higher HOMA-IR and lower QUICKI indices, i.e. decreased insulin sensitivity, in the SGA children. In addition, serum adiponectin showed a negative correlation with WC and WC/height ratio, suggesting that some SGA children might have extra visceral fat explaining reduced adiponectin levels even though they were not overweight or obese. This type of association was recently reported in somewhat younger SGA children (25). Glucocorticoids have a negative effect on adiponectin (26, 27). Dexamethasone inhibited adiponectin release from human adipocytes (26). Furthermore, decreased circulating adiponectin levels were found in nonobese patients with Cushing’s syndrome and in healthy subjects after i.v. hydrocortisone administration (27). Cianfarani et al. reported that SGA children had lower adiponectin levels than their AGA control children (28), and that the adiponectin levels were even lower in those SGA children with good postnatal catch-up growth than in those without it (28, 29). In contrast, Evagelidou et al. (30) found that adiponectin levels had a trend to be slightly higher in SGA children than in their control children. The present study could not find any difference in adiponectin concentrations between SGA and AGA children, and no difference in adiponectin levels was found between the SGA children with good or poor catch-up growth. However, in our SGA subjects, serum adiponectin concentrations correlated inversely with GBA levels even when adjusted for WC/height ratio, which suggests an independent association between GBA and adiponectin. This finding is consistent with the previous reports on the relationship between adiponectin and glucocorticoids (26, 27).

The reasons for being born SGA are numerous and include fetal, uteroplacental, and maternal factors. The possible mechanisms explaining increased cardiovascular morbidity in adults born SGA (Barker’s hypothesis) may also vary depending on the etiology of the intrauterine growth restriction. Thus, it is not surprising that it has been difficult to find conclusive evidence for a single mechanism explaining the association between small birth size and cardiovascular morbidity in adulthood. The current study showed that the subgroup of SGA children with the highest HOMA-IR index had significantly higher GBA than the SGA subjects with lower HOMA-IR index, and that GBA turned out to associate independently of obesity (BMI or WC/height ratio) with high HOMA-IR index. Low serum adiponectin level also associated independently of obesity with high HOMA-IR index. Furthermore, high GBA levels associated independently of obesity with low adiponectin levels. Thus, increased...
glucocorticoid activity may increase IR in SGA subjects by several mechanisms. The strengths of this study were the carefully matched SGA and AGA children with a similar age at examination. Moreover, the blood samples were taken exactly between 0900 and 1000 h, minimizing the effect of diurnal variation on both cortisol and adiponectin concentrations. The transactivation-based GBA assay turned out to give additional information compared with the traditional cortisol assay and FCI. Obviously, it is desirable that our findings will be replicated in other (hopefully larger) studies.

In conclusion, there is an SGA subgroup with a more insulin-resistant state detectable already in childhood. Increased glucocorticoid activity and low adiponectin associate independently with this increased IR. GBA measurement, based on a transactivation assay, seems to estimate the circulating bioavailable cortisol concentrations better than the total cortisol measurement or the calculated FCI.

Declaration of interest
The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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References


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