Differences in expression and activity of 11β-hydroxysteroid dehydrogenase type 1 and 2 in human placentas of term pregnancies according to birth weight and gender

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Abstract

Background: Fetal exposure to maternal glucocorticoids may determine fetal growth and the programing of later disorders. Availability of the glucocorticoids in the placenta is regulated by the 11β-hydroxysteroid dehydrogenase (11β-HSDs) enzymes. To date, there are discrepancies with regard to cortisol (F) cord blood levels in fetuses with intrauterine growth retardation in different species.

Objective: To study the expression and activity of 11β-HSDs in placentas from full term small for gestational age (SGA), appropriate for gestational age (AGA) and large for gestational age (LGA) newborns, and cortisol cord blood concentration.

Methods: Twenty-five placentas from AGA, 24 SGA and 25 LGA were collected.

Results: SGA newborns had significantly lower and LGA newborns had significantly higher birth weight, birth length, head circumference, and placental weight than AGA counterparts. We observed a direct correlation between placental weight and birth weight, birth length and head circumference, and higher cord F levels in SGA newborns. The 11β-HSD1 expression was similar among the SGA, AGA, and LGA placentas. However, within the placentas of SGA newborns, the 11β-HSD1 mRNA levels were significantly reduced in the chorionic plate compared with basal plate. An inverse correlation between cord F levels and activity of 11β-HSD1 in the chorionic plate of the SGA placentas was detected. The 11β-HSD2 activity was seven- to eightfold higher compared with 11β-HSD1 in the placentas, and there was a lower 11β-HSD2 activity in females’ SGA placentas compared with the male SGA placentas.

Conclusion: We observed a lower expression and activity of 11β-HSD1 in the chorionic plate of the SGA placentas, suggesting a possible compensatory mechanism to diminish the higher cortisol fetal concentrations observed in fetuses with intrauterine growth restriction.

Introduction

Intrauterine growth retardation (IUGR) is associated with an increased incidence of cardiovascular disease and alterations in glucose homeostasis in later life. Many hypotheses have been advanced to explain these features (1). Intrauterine alterations in adrenocortical function or cortisol (F) metabolism, which occur during gestation, may have long-lasting effects leading to these derangements in adulthood. Two different isoenzymes of 11β-hydroxysteroid dehydrogenase type 1 (11β-HSD1) and type 2 (11β-HSD2) participate in F metabolism and both have been identified in human placentas (2, 3). The 11β-HSD1 is a bidirectional enzyme that acts primarily as an 11-oxoreductase forming active cortisol from inactive cortisone (4). Its activity in human placentas has been found to be lower than those of 11β-HSD2. Changes in these enzymes during gestation may be important in regulating fetal birth weight and in the mechanisms involved in human parturition (5, 6). Placental 11β-HSD2 mRNA levels have been reported to be significantly decreased in intrauterine growth restriction pregnancies when compared with gestationally matched, appropriately grown placentas (7). Decreased placental 11β-HSD2 activity has been suggested to play a role in IUGR and to its metabolic consequences (8). Mutations in the 11β-HSD2 gene invariably have reduced birth weight, and recently it has been shown that there is a reduced placental 11β-HSD2 activity in preterm pregnancies also present with IUGR (9). Interestingly, placental architecture and blood flow are not uniform across chorioallantoic human placental disk. Proximity to the umbilical cord, basal plate or chorionic plate may influence perfusion and protein and gene expression (10). Despite these limitations, it seems plausible that
differences in sampling site contribute to variability in
gene and protein expression across the placental disk
(10). We hypothesize that the expression and/or the
activity of 11β-HSDs differ between the chorionic and
basal plates of the placentas according to the newborn
birth weight in human term pregnancies. Because
currently there is no information regarding the
expression and placental activity of 11β-HSDs type 1
and 2 from newborns born small for gestational age
(SGA) compared with those with macrosomia and
whether there are differences according to gender in
human term pregnancies, we designed a study to
determine the mRNA expression and the activity of
11β-HSDs in full-term placentas from newborns born
appropriate for gestational age (AGA), SGA, and large
for gestational age (LGA) as well as to assess F
concentrations in cord blood.

Material and methods

Sample collection
The placental tissue was collected immediately after
delivery. We selected pregnancies from full-term preg-
nancies (37–40 week of gestation) defined by standard
clinical criteria and with a birth weight between the 10th
and the 90th centiles for gestational age defined as AGA;
below the 10th centile was defined as SGA and over the
90th centile as LGA using the Chilean birth weight
reference (11). Exclusion criteria were maternal hyper-
tension, diabetes, use of glucocorticoids during preg-
nancy, or a reduced amount of amniotic fluid at term. All
mothers gave their informed consent and this protocol
was approved by the Institutional Review Board of the
San Borja Arriaran Clinical Hospital in Santiago, Chile.

Each placenta was inspected by a pathologist (EK) for
every visible abnormalities and was excluded in the
presence of macroscopic alterations. Placental samples
were rapidly processed as described by Wyatt et al. (10).
Briefly, each placenta was sectioned transversally
(∼ 5 cm) using a sterile scalpel near the cord insertion
site. The tissue was subsequently divided into three
horizontal segments from the basal towards the
chorionic surface. Each sample was rinsed thoroughly
in cold sterile saline solution (NaCl 9.0 g/l), and samples
from the different sections were frozen in liquid nitrogen
and stored at −80 °C for RNA and protein analysis.

The cord blood was processed by the laboratory within
20 min, centrifuged (2000 × g for 10 min at 4 °C), and the
sera were aliquoted and stored at −20 °C until assayed.

RNA purification, RT, and semi-quantitative
PCR
Total RNA was isolated from the frozen placentas
(chorionic and basal plate samples) as described previously (12)
using TRIzol Reagent (Invitrogen). cDNA was synthesized from 2 μg of total RNA, previously
digested by DNase I (Fermentas AB, Vilnius, Lithuania),
using random primers (Invitrogen) and 200 U
RevertAid H Minus M-MuLV reverse transcriptase
(Fermentas) following the manufacturer’s instructions.
The gene amplifications were assessed using specific pairs of primers: 11β-HSD1 upstream, 5’-AGGAAAG-
CTCATGGGAGACTAG-3’ and downstream, 5’-ATGG-
TGAATATCATCATGAAAAATTC-3’; 11β-HSD2
upstream, 5’-TCAGACAGAAGTCAAGTGAGAAGC-3’
and downstream, 5’-GGAATGCCATGAAGTG-3’;
18s rRNA upstream, 5’-GTAACCGTTGAACCCATT-3’
and downstream, 5’-CCATCCAATCGGATGTACCC-3’.

Two μl of cDNA were added to the reaction mix (total
volume of 25 μl) containing PCR buffer, 1.5 mmol/l
MgCl₂, 0.625 U Taq DNA polymerase (Invitrogen),
0.25 mmol/l nucleotide mix (Fermentas), and
0.4 μmol/l of each specific human primer (Integrated
DNA Technologies, Miami, FL, USA). The PCR was
performed in a P × 2 Thermal Cycler (Thermo Electron
Corporation, Waltham, MA, USA) at 94 °C for 45 s
denaturation), 55 °C for 60 s (annealing), and 72 °C
for 90 s (extension), and repeated for 30 cycles. As an
internal control, 18s rRNA was amplified in each
sample following the same conditions described above,
and repeated for 18 cycles. To determine that the
amplification of all the genes was within a linear range,
we evaluated the linearity of amplification of the
corresponding transcripts in human placenta and we
subsequently selected the number of cycles.

Amplified fragment products were visualized on a
1.0% agarose gel using ethidium bromide staining.
Semi-quantification of PCR products was performed by
image analysis (Kodak EDAS 290 Electrophoresis
Documentation and Analysis System, Kodak 1D Image
Analysis Software).

Protein extraction
The frozen placental tissue was powdered in a ceramic
mortar with liquid N₂ and homogenized for 30 s with
a mechanical homogenizer (Kontes Glass Company,
Vineland, NJ, USA) in ice-cold 0.1 M PBS pH 7.5
supplemented with anti-proteases (Complete, Mini,
EDTA-free Protease Inhibitor Cocktail Tablets, Roche
Applied Science). The tissue homogenate was centri-
figuted at 10 000 g for 30 min at 4 °C. The resulting
supernatant was collected and assayed for protein
concentration using the BCA protein assay kit (Pierce,
Rockford, IL, USA) with BSA as a standard.

11β-HSD activity studies: type 1 and type 2
isoforms
Tissue homogenates were centrifuged for 10 min at
1000 g to sediment large tissue fragments, and a protein
assay was performed on the supernatant (BCA, Pierce).
To measure the activities of 11β-HSD1 and 11β-HSD2, placental homogenates from chorionic and basal plates (0.40 mg of total protein) were incubated in 1.0 ml phosphate buffer (0.1 mol/l; pH 7.6) in the presence of 50,000 c.p.m. of 3H cortisol or 3H F (Amersham International). 0.1 μM cortisone or 0.25 μM F, and 400 nmol/l NADP or NADP respectively, for 30 min at 37 °C in a shaking water bath. Aliquots were extracted into 10 volumes of dichloromethane, and steroids were separated by high-performance thin layer chromatography (HPTLC, Merck), using methanol-chloroform (5:95) as a mobile phase. The bands containing the labeled F and cortisol were identified by u.v. light of the cold carriers, cut out into scintillation vials, and measured in scintillation counter (Tracor Analytic Delta 300). The rates of F to cortisol or cortisone to F conversion were calculated from the specific activity of the labeled F or cortisone and the radioactivity of cortisone or F, and results expressed as the amount of F or cortisol (picograms) formed per min per mg protein.

Cortisol assay

Cord blood cortisol was determined using a commercial RIA (Diagnostic Products Co., DPC, Los Angeles, CA, USA). The sensitivity of this assay is 1 μg/dl. The intra- and inter-assay coefficient of variations are 3.1 and 5.2% respectively.

Statistical analyses

Results are shown as mean ± S.E.M. Differences within each group (SGA, AGA, and LGA) were assessed by one-way ANOVA or Kruskal–Wallis, followed by Bonferroni test for multiple comparisons. To compare the results of the basal and chorionic plates in each placenta, we used the paired t-test or Wilcoxon test. The studies of correlations were performed using the Pearson or Spearman test. Statistics were performed using SPSS version 11.5; a value of P < 0.05 was considered significant.

Results

Twenty-four placentas from SGA newborns (14 males), 25 from LGA newborns (14 males), and 25 from AGA newborns (14 males) were collected. Of the total deliveries, 24.7% were by cesarean section. The median Apgar score after 1 and 5 min was nine. The anthropometrics and clinical data of all neonates are shown in Table 1. As expected, the SGA newborns had significantly lower birth weight SDS (P < 0.001), birth length SDS (P < 0.001), head circumference SDS (P < 0.001), and placental weight (P < 0.01) than the AGA and LGA newborns. Likewise, the LGA newborns had significantly higher birth weight SDS, birth length SDS, head circumference SDS, and placental weight than the AGA newborns. In addition, the cortisol cord blood levels were significantly higher in the SGA group (mean: 16.0, range: 8.6–31.2 μg/dl) compared with the AGA (mean: 11.0, range: 4.7–15.4 μg/dl) and LGA (mean: 10.3, range 5.1–17.3 μg/dl) newborns (Fig. 1).

Table 1 Anthropometric characteristics of the small for gestational age (SGA), appropriate for gestational age (AGA), and large for gestational age (LGA) newborns.

<table>
<thead>
<tr>
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<th>SGA (24)</th>
<th>AGA (25)</th>
<th>LGA (25)</th>
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<tbody>
<tr>
<td>Gestational age (weeks)</td>
<td>38.1 ± 0.3</td>
<td>39.2 ± 0.2</td>
<td>39.5 ± 0.2</td>
</tr>
<tr>
<td>Birth weight (SDS)</td>
<td>−1.71 ± 0.10†</td>
<td>0.25 ± 0.13</td>
<td>2.50 ± 0.14†</td>
</tr>
<tr>
<td>Birth length (SDS)</td>
<td>−1.04 ± 0.25†</td>
<td>0.65 ± 0.20</td>
<td>1.53 ± 0.15†</td>
</tr>
<tr>
<td>Head circumference (SDS)</td>
<td>−0.10 ± 0.03†</td>
<td>0.01 ± 0.03</td>
<td>0.15 ± 0.04†</td>
</tr>
<tr>
<td>Placental weight (g)</td>
<td>495 ± 15*</td>
<td>631 ± 19</td>
<td>741 ± 23†</td>
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</table>

Data are shown as mean ± S.E.M. *P < 0.05 SGA versus AGA and LGA; †P < 0.05 LGA versus AGA.

Figure 1 Cortisol cord blood levels in SGA, AGA, and LGA newborns. The concentrations are expressed as mean ± S.E.M. Data were analyzed by two-way ANOVA followed by a post hoc LSD test. *P < 0.05; SGA versus AGA and LGA newborns.
11β-HSD1 and 11β-HSD2 activities in the placenta

We observed a lower activity of 11β-HSD1 in both chorionic and basal plates of the SGA placentas, compared with the activity observed in the AGA and LGA placentas. This activity was significantly lower in the chorionic plate of the SGA placentas compared with the chorionic plate of the AGA placentas (0.89 ± 0.03 vs 1.32 ± 0.14 pg/min per mg protein, \( P = 0.020 \)). The activity of the 11β-HSD2 enzyme in placenta was seven- to eightfold higher compared with the activity of 11β-HSD1, \( (P!0.01) \) and there were no differences among the three groups of newborn weight placenta.

The 11β-HSD1 and 11β-HSD2 activities were studied according to the newborn gender. There were no differences of enzyme activities between females \( (n=32) \) and males \( (n=42) \) in the whole group of placentas. However, when placentas were separated according to the newborns’ birth weight, we observed a lower activity of the 11β-HSD1 (Fig. 3A) in the chorionic plate of female SGA placentas compared with the chorionic plate of the female AGA placentas \( (0.89 \pm 0.03 \text{ vs } 1.32 \pm 0.14 \text{ pg/min per mg protein, } P=0.029) \), as well as in male SGA placentas compared with the chorionic plate of the male AGA placentas \( (0.94 \pm 0.07 \text{ vs } 1.31 \pm 0.20 \text{ pg/min per mg protein, } P=0.037) \). However, within the AGA placentas, we observed differences in the 11β-HSD2 activity according to the newborn gender. In the females’ AGA placentas \( (n=11) \), a lower activity for 11β-HSD2 (Fig. 3B) was detected compared with the male AGA placentas \( (n=14) \) in the chorionic plate \( (5.2 \pm 0.9 \text{ vs } 9.5 \pm 1.0 \text{ pg/min per mg protein, } P=0.007) \) as well as in the basal plate \( (7.6 \pm 1.0 \text{ vs } 9.5 \pm 1.0 \text{ pg/min per mg protein, } P=0.091) \). These differences in 11β-HSD2 activity were not observed in SGA or LGA placentas.

Correlation studies

We found a direct correlation between the weight of the placenta with birth weight SDS \( (r=0.749, P<0.001) \), birth length SDS \( (r=0.647, P<0.001) \), and birth head circumference SDS \( (r=0.693, P<0.001) \) for the whole group. A direct correlation was found between placental weight with 11β-HSD1 activity in the chorionic plate \( (r=0.359, P=0.006) \) for the whole group. Conversely, there was no significant correlation between placental weight and 11β-HSD1 or 11β-HSD2 expression (data not shown). Cord blood levels of cortisol in the whole group correlated inversely with birth weight \( (r=-0.369, P=0.005) \), placental weight \( (r=-0.426, P=0.001) \), and with 11β-HSD1 activity in the chorionic plate \( (r=-0.315, P=0.015) \).

Discussion

In the present study, we analyzed the expression and activity of 11βHSD type 1 and 2 in human placentas of term pregnancies according to birth weight and gender. We observed a direct correlation between placental
weight and birth weight SDS, birth length SDS and head circumference SDS. In addition, we found higher cord F levels in SGA at term newborns. However, in this regard, there are some discordant observations (13, 14). No differences in F from mixed umbilical cord serum in SGA newborns compared with matched AGA controls have been detected (14). Of note, in this later report, ten subjects were born preterm and three postterm, and since gestational age influences adrenal function, this result might be biased. In addition, recently Strinic et al., including only term newborns have found lower F levels in IUGR fetuses in umbilical venous blood (13). The latter observation has been also reported in animals (15). In mammals, glucocorticoids are essential for normal fetal organ growth and maturation (16). However, excessive exposure to glucocorticoids in utero leads to IUGR (16, 17), giving support to our results of increased F cord blood levels in SGA newborns. It has been suggested that the rise in plasma F concentrations in SGA fetuses (18) is a mechanism to combat hypoglycemia and may contribute to the maturation of fetal tissues. However, in human fetuses, there is no plasma F surge before 36 week’s gestation by which time most fetal organs are mature (19).

The cause of increased blood F in fetuses with suspected IUGR is debatable. Fetal plasma ACTH has been reported to be decreased, which might be due to the negative feedback from increased fetal F concentrations or the direct consequence of reduced placental size and possibly reduced placental production of ACTH. Corticotrophin-releasing hormone (CRH), or both (20). Therefore, one likely explanation is increased adrenal blood flow and adrenal hyperplasia, the consequence of redistribution in fetal circulation in response to hypoxemia. Increased plasma CRH–ACTH in the umbilical cord of 28 growth-retarded fetuses has been reported, strongly suggesting that placental CRH may be stimulated in conditions of chronic stress, and in turn modulates fetal pituitary–adrenal function in high risk pregnancies (21).

The 11β-HSD1 expression was similar among the SGA, AGA, and LGA placentas in both the chorionic and basal plates of the placentas. However, only within the placentas of SGA newborns, the 11β-HSD1 mRNA levels were significantly reduced in the chorionic plate compared with the basal plate of these placentas. Interestingly, we observed an inverse correlation between cord F levels with activity of 11β-HSD1 in the chorionic plate of the SGA placentas. In addition, the 11β-HSD1 activity in the SGA placentas was lower compared with the enzyme activity observed in the AGA and LGA placentas, and this difference was strengthened when comparing the chorionic plate of SGA placentas with the chorionic plate of AGA placentas and in placentas from female newborns. These findings have not been reported previously. In human pregnancies with IUGR, a decreased expression of 11β-HSD1 in placenta has been reported (22). Nevertheless, this finding must be interpreted with caution since it included term and preterm neonates. In fetal sheep, a specific increase in the expression of 11β-HSD1 mRNA in the liver of growth-restricted fetuses in late gestation has been observed (23). The authors suggest that an increased hepatic exposure to F in the growth-restricted fetus may be important in the reprogramming of hepatic physiology that occurs after growth restriction. This possible regulation of the 11β-HSD1 enzymes in the placenta by F is also supported by the direct correlation between the placental weight and activity of 11β-HSD1 in the chorionic side of the placentas and an inverse correlation with cord blood F. These findings point towards a compensatory mechanism of the fetal side of the placental to higher levels of fetal F. Several in vivo and ex vivo studies have shown that the GH–insulin-like growth factor 1 (IGF1) system inhibits the expression and activity of 11β-HSD1 resulting in reduced local regeneration of F. Recently, the mechanism through which GH inhibits the 11β-HSD1 gene promoter activity has been shown to be via IGF1 in HepG2 cells (24). Interestingly, in the chorionic plate of the SGA placentas, we have found an increased expression of IGF1, and thus this could be a mechanism through which the 11β-HSD1 gene is decreased in the chorionic plate of the SGA placentas (25).

As has been previously reported, the 11β-HSD2 enzyme activity was seven- to eightfold higher compared with 11β-HSD1 activity from the three
groups of placentas. Decreased placental 11β-HSD2 activity has been suggested as a reason for IUGR and its metabolic consequences. In the present study, there were no significant differences in the mRNA expression and activity of 11β-HSD2 among the three groups of placentas from newborns with different birth weights. Moreover, no differences in the 11β-HSD2 mRNA expression and protein activities from the chorionic plate compared with the basal plate were observed among the SGA, AGA, and LGA placentas. Our findings are in contrast to other recent reports. In one study in 19 IUGR placentas of different gestational ages, a decreased 11β-HSD2 gene expression was observed in intrauterine growth restriction pregnancies compared with placentas from gestational matched AGA newborns (7). Importantly, the numbers of term placentas with IUGR in this study were only four and with very severe documented IUGR, not only a low birth weight newborn. In small preterm infants, reduced placental 11β-HSD2 activity is associated with low relative birth weight (24) and severe fetal distress (9). It is important to point out that in the latter study, the placentas included were from pregnancies between 22 and 32 weeks of gestational age and the 11β-HSD2 enzyme expression increases after 28 weeks by 12-fold. Deficiency of the barrier to maternal glucocorticoids may represent a common pathway between the maternal environment and feto-placental programing of later disease. However, we did not find significant variations in the expression or in the activity of this enzyme among the different groups. This finding is in agreement with Rogerson et al. (26) who reported that the enzymatic activity and mRNA levels of 11β-HSD2 measured in 111 human placentas taken from normal births were not correlated with either fetal or placental weight. These studies suggest that variations in placental 11β-HSD type 2 activities do not influence fetal or placental weight in humans. The expression and activity of this enzyme in fresh frozen human placenta increase progressively from first to second to third trimester (term) pregnancies at a time when maternal circulating levels of glucocorticoid are rising. The finding of attenuated placental 11β-HSD2 activity in IUGR in some reports, but not others, must be carefully interpreted due to the differences in expression and activity in this enzyme throughout gestation (24). One of the strengths of our report is that gestational ages were within a narrow range of term pregnancies. On the other hand, severe IUGR is usually interrupted earlier than term and thus may have precluded us to find significant differences of expression and activity of this enzyme in term placentas (27). The capacity of 11β-HSD2 is massive, allowing maintenance of fetal F concentrations several times lower than maternal F, and, at least in term infants, three-fourths of F concentrations originate from the fetal adrenals (28).

Nevertheless, we observed a lower 11β-HSD2 activity in AGA female placentas compared with AGA male placentas. These gender differences were not found in the other birth weight groups of placentas, and yet we do not have a plausible explanation about the meaning of this observation. In a previous study by Murphy et al. (29), the authors document a different effect by fetus gender in asthmatic pregnant women in placental 11β-HSD2 activity, but not within the placentas collected from mothers without asthma. The differences with our findings might be due to tissue extracted from total placenta and the rather small sample size. We hypothesize that the lower activity of 11β-HSD2 in AGA females might be a consequence of differences in estrogen levels (30).

In summary, we observed in full-term pregnancies, a lower 11β-HSD1 expression and activity in the chorionic plate of the SGA placentas, suggesting a possible compensatory mechanism to diminish the higher F fetal concentrations observed in fetuses with intrauterine growth restriction. The finding of different expression and/or the activity of 11β-HSDs between the chorionic and basal plate of the placentas according to the newborn birth weight and according to gender have not been reported previously and point towards a possible different regulation of this enzyme within the placenta.

**Declaration of interest**

There is no conflict of interest that could be perceived as prejudicing the impartiality of the research.

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