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Sex-specific associations between cortisol and birth weight in pregnancies complicated by asthma are not due to differential glucocorticoid receptor expression

Nicolette A Hodyl,1 Hayley Wyper,2 Annette Osei-Kumah,1 Naomi Scott,2 Vanessa E Murphy,3 Peter Gibson,3 Roger Smith,2 Vicki L Clifton1

ABSTRACT
Background Fetal growth inhibition is a known sequela of in utero glucocorticoid exposure and has long-term consequences for adult health. Sex-specific fetal growth patterns are observed in pregnancies with maternal asthma and may be due to differential sensitivity of the placenta to glucocorticoids. It is currently unknown whether expression of the placental glucocorticoid receptor (GR) becomes altered with asthma or the use of inhaled corticosteroids.

Methods Pregnant women with mild asthma (n=52), moderate-severe asthma (n=71) and without asthma (n=51) were recruited at John Hunter Hospital, Newcastle, Australia. At delivery, placentae and cord blood were collected, and fetal sex and birth weight were recorded. Placental GR heterogeneous nuclear RNA (hnRNA), mRNA and protein were measured and cord blood cortisol concentrations were assessed.

Results Placental GR gene activity increased with cortisol exposure but decreased with increased inhaled corticosteroid treatment (p=0.05). With maternal asthma, female birth weight centiles were inversely associated with cortisol (r=-0.286, p=0.017) and, despite a decrease in placental GR mRNA (p=0.003), placental GRα protein levels were unchanged. In males, no change to cortisol, birth weight or placental GR were evident in pregnancies with asthma. Together, these results indicate that in pregnancies complicated by asthma, placental GR gene activity, but not mRNA expression or protein levels, is dependent on cortisol and inhaled corticosteroid treatment.

Conclusions The sex-specific associations between cortisol and birth weight observed in pregnancies with asthma are not due to altered GR expression; however, they may be due to differential glucocorticoid sensitivity via preferential transcription of GR isoforms or post-translational modifications.

INTRODUCTION
With the increasing prevalence of childhood allergy and atopic disease worldwide, an urgent need has arisen to understand in utero mechanisms that predispose infants towards this outcome. These mechanisms are currently unknown; however, the contribution of maternal asthma and atopy during pregnancy to wheeze and atopic symptoms in childhood is in the preliminary stages of investigation.1 2 Retrospective studies have indicated that maternal atopy alters feto-placental immune mechanisms, with cord blood cytokine concentrations skewed towards a type 2 T helper profile.3 5 Ongoing work from our laboratory has demonstrated that the effects of maternal asthma during pregnancy on feto-placental immunity are dependent on asthma severity and use of inhaled corticosteroids (ICS) for asthma treatment, and differ according to fetal sex.6 9 Further, maternal asthma is associated with a reduction in female birth weight,9 and low birth weight has been independently linked to other adverse health outcomes in both children and adults.10

Animal and human studies indicate that fetal growth is reduced following excess exposure to glucocorticoids (for a review, see Seckl and Holmes11). This appears to be an important factor driving the sex-specific reduction in birth weight that we have previously reported in pregnancies complicated by asthma. Specifically, an asthma-induced reduction in the placental cortisol-metabolising enzyme, 11β hydroxysteroid dehydrogenase type 2 (11β-HSD2) was associated with reduced female birth weights.9 This reduction in 11β-HSD2 activity increases exposure of the female fetus to cortisol. In these studies, the use of ICS to manage asthma restored female birth weight centiles to control levels and normalised 11β-HSD2 activity.12 The protective effect on female birth weight conferred by ICS treatment for maternal asthma may be due to sexually dimorphic alterations in placental glucocorticoid responsiveness.

Responsiveness to glucocorticoids is mediated both centrally and peripherally by the glucocorticoid receptor (GR), of which there are multiple isoforms.13 We have previously identified the presence of four GR isoforms in the placenta: GRα, GRβ, GRγ and GR-ß, although the latter was only identifiable in a subset of placentae.14 The most predominant mature isoform expressed was GRα, consistent with its ubiquitous expression throughout the body. This isoform mediates the functional biological effects of glucocorticoids, including cell growth, proliferation and differentiation (for a review, see Pujols et al15), and our studies have indicated that its expression in the placenta becomes inhibited following in vitro glucocorticoid exposure.14 The actions of other GR isoforms also determine functional responses to glucocorticoids: GRβ is a dominant inhibitor of GRα binding,16 GR-ß enhances GRα transcription17 and GRγ plays a role in mediating glucocorticoid resistance in childhood acute lymphoblastic leukaemia.18

The placental response to glucocorticoids appears crucial in determining fetal growth outcomes. Exposure to synthetic glucocorticoids during...
pregnancy is associated with reductions in birth weight in a number of animal models, which often appear sex specific; however, data from comparable human studies remain inconclusive. This is an important consideration in the study of placental GR regulation in pregnant women with asthma, as circulating endogenous cortisol increases with asthma severity and asthma is often managed with inhaled or oral steroids. This study will therefore assess the effects of endogenous cortisol and ICS on in vivo GR expression in the delivered placenta and its relationship to birth weight. The aims of this study were (1) to identify whether maternal asthma was associated with differences in transcription and translation of placental GR mRNA and GR protein expression; (2) to identify whether ICS treatment affects regulation of the placental GR; and (3) to identify sex-specific relationships between birth weight centiles and GR protein expression in asthmatic and control pregnancies.

METHODS
Participants
Ethics approval was gained from the Hunter New England Health and the University of Newcastle Human Research Ethics Committees. We have previously reported maternal and fetal characteristics of a cohort of pregnant women recruited into our asthma studies and have reported an adherence rate to ICS treatment of 80%. The current data correspond to a subset of this cohort, where we were able to collect placenta and venous cord blood with 45 min of delivery. Power analyses were conducted using Power and Precision Software, ensuring a power of 80% and of 0.05, based on variability observed in our previous findings. Control women without asthma (n=51) and women with asthma (n=123) were recruited at their first antenatal visit to the John Hunter Hospital (Newcastle, NSW, Australia). Asthma severity was assessed as described previously, and graded as mild, moderate or severe at 12, 18 and 30 weeks gestation. Participants were grouped according to the most severe grade of asthma experienced during pregnancy, as previously reported.

Given that cigarette smoking can affect birth weight and attenuate cortisol levels, participant self-report of cigarette smoking was recorded from antenatal records and following interview at recruitment. Birth weight and length were recorded and converted into centiles using growth charts and a birth centile calculator (http://www.gestation.net/; Perinatal Institute, UK), with birth centiles <10 defined as small for gestational age (SGA).

Placental tissue preparation, RNA isolation and real-time reverse transcription–PCR (RT–PCR) analysis
Placental tissue samples were pooled from six cotyledons, blotted to remove blood, snap-frozen and stored at –80°C. Pooling of tissue removes variability across sites on the placenta. Total RNA was extracted from 0.2 g of crushed frozen placenta using the TRizol method (Invitrogen, Mt Waverley, Australia), as described previously. Quantitative real-time RT–PCR was used to determine mRNA abundance of total GR, GRα, GR exons 1A5 and GR heterogeneous nuclear RNA (hnRNA) in placental samples compared with levels of the housekeeping gene, β-actin. GR hnRNA is the precursor of GR mRNA and thus serves as a reliable surrogate marker of gene activity in both fresh tissue and cultured cells. The real-time RT–PCR protocol, primer sequences and optimal concentrations were identical to those reported previously. All samples were assayed in triplicate and variability in mRNA expression was <5%.

Protein extraction and western blot analysis
Placental tissue was homogenised and protein extracted as described previously. Extracted protein samples were prepared by combining extract (20 μg) with 5 μl of NuPAGE lithium dodecyl sulfate (LDS) sample buffer. Samples were heated for 12 min at 70°C and 2 μl of NuPAGE reducing agent (Invitrogen, Life Technologies, Carlsbad, California, USA) was added. Protein samples (20 μg/lane) were separated by electrophoresis through 7% Tris-acetate precast gels on an Xcell SureLock Electrophoresis cell (Invitrogen). Gels were run at 150 V for 1 h in a Tris-acetate running buffer with 500 μl of NuPAGE antioxidant (Invitrogen) in the inner chamber. Proteins were electrothermally transferred to 0.45 μm pore nitrocellulose membranes (Hybond-C Extra, Amersham Biosciences, UK) at 50 V for 1 h. Transfer membranes were incubated overnight with 5% skim milk in blocking solution (Tris-buffered saline (TBS) with 5% skim milk powder). Blots were incubated for 2 h at room temperature with 1:200 dilution of the primary antibody in blocking solution. For detection of GRα protein, an affinity-purified rabbit polyclonal immunoglobulin G (IgG) antibody was used (Santa Cruz Biotechnology, Santa Cruz, California, USA). Blots were incubated for 1 h at room temperature with a horseradish peroxidase-labelled goat antirabbit secondary antibody provided with the ECL kit (Pierce Biotechnology, Rockford, Illinois, USA). Signal detection of antibody complexes was performed using Pierce Supersignal Dura Extended Duration Substrate (Pierce Biotechnology) as per the manufacturer’s instructions. Densitometric analysis (Multi Gauge V2.4) was used to quantify the intensity of the protein signals relative to a 38 kDa band to which the secondary antibody bound, to adjust for loading (figure 1). We identified that the 38 kDa band changed with protein concentration in our optimisation studies and we have
previously used this method as a control for loading variations. Each blot contained a positive control (rat brain protein extract) and an interassay control (pooled placental sample).

Plasma collection and cortisol radioimmunoassay (RIA)
Plasma was separated from cord blood via centrifugation. Cortisol concentrations were determined in duplicate in plasma collected from the placental umbilical vein, using a commercial RIA kit (Diagnostic Systems Laboratories, Webster, Texas, USA) following the manufacturer’s instructions. The detection limit was 4.7 nmol/l; the intra-assay variation was 5.1% and cross-reactivity of cortisol antiserum with other steroids was <0.1%.

Statistical analysis
Statistics were performed using the Statistical Package for the Social Sciences (SPSS v 17; SPSS, Chicago, Illinois, USA). An a priori decision was made to analyse the data separately by sex when comparing the effects of asthma severity and treatment, as past research indicates that placental cortisol metabolism differs between males and females in the presence of asthma. The abundance of GR hnRNA and mRNA transcripts was log (natural) transformed to normalise the data. To assess the effect of asthma severity and asthma treatment on protein and mRNA expression abundance, one-way analysis of variance (ANOVA) was used with Tukey post hoc testing where indicated. Student t tests and Mann–Whitney U tests were used for comparisons made between two normally and non-normally distributed variables, respectively. Pearson correlations were performed to assess associations between the log-transformed expression data, birth weight centiles and cortisol concentrations. Frequencies of maternal smoking and SGA births in the control and asthma groups were analysed using Chi². p Values ≤0.05 were considered statistically significant.

RESULTS
Maternal and neonatal characteristics
The maternal and neonatal characteristics are reported according to both asthma severity and asthma treatment in tables 1 and 2, respectively. Maternal age was significantly higher in the control population than both the populations with asthma (F(2,171)=8.715, p<0.001). Maternal forced expiratory volume in 1 s (FEV1) was significantly lower in the moderate–severe asthma group compared with both the control and mild asthma groups (F(2,171)=3.225, p=0.043), but did not differ according to asthma treatment (p=0.100). More women with asthma smoked (53.8%) compared with controls (5.9%; χ²=13.564, p=0.001).

Male birth weight centiles were significantly greater than those of females (t(175)=2.78, p=0.006). Female birth weight, birth weight centile, length, and abdominal and head circumferences were not different between the groups. In females, cord blood cortisol was significantly increased in the moderate–severe asthma group compared with the controls and the mild asthma groups (F(2,67)=3.2, p=0.047), and negatively correlated with birth weight centiles (r=-0.286, p=0.017, n=69). Significantly more females were born SGA in the group with moderate–severe asthma (22.5%) than in both the control (9.5%) and the mild asthma (5.2%) groups (χ²=6.018, p=0.049).

In male neonates, no difference was observed in birth weight centiles or cortisol between the control and asthma treatment groups, and the two were not correlated. Frequency rates of male infants born SGA did not differ between the control and asthma groups. Length centiles were significantly reduced in the moderate–severe asthma group compared with the control and mild asthma groups (F(2,54)=4.159, p=0.021).

Table 1  Maternal characteristics of the study populations according to asthma treatment and asthma severity

<table>
<thead>
<tr>
<th></th>
<th>Asthma treatment</th>
<th>Asthma severity</th>
<th>p Value</th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Control</td>
<td>No treatment</td>
<td>Inhaled corticosteroids</td>
<td>p Value</td>
<td>Mild</td>
<td>Moderate–severe</td>
</tr>
<tr>
<td>n</td>
<td>51</td>
<td>52</td>
<td>71</td>
<td>&lt;0.001</td>
<td>58</td>
<td>65</td>
</tr>
<tr>
<td>Age</td>
<td>29.8 (27.4 to 30.9)*</td>
<td>25.7 (23 to 27)</td>
<td>26.0 (25 to 28)</td>
<td>&lt;0.001</td>
<td>26.3 (24.3 to 26.8)</td>
<td>26.5 (25.0 to 28.0)</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>164.2 (162.2 to 165.9)</td>
<td>164.1 (161.8 to 166.1)</td>
<td>164.0 (162.7 to 165.8)</td>
<td>0.99</td>
<td>163.4 (162.0 to 166.0)</td>
<td>164.4 (162.6 to 165.8)</td>
</tr>
<tr>
<td>Weight (early) (kg)</td>
<td>73.9 (65.0 to 77.4)</td>
<td>76.9 (69.2 to 81.1)</td>
<td>75.3 (69.8 to 81.3)</td>
<td>0.80</td>
<td>77.4 (68.9 to 80.7)</td>
<td>74.7 (70.0 to 81.6)</td>
</tr>
<tr>
<td>Weight (late) (kg)</td>
<td>83.0 (76.8 to 87.9)</td>
<td>88.3 (81.7 to 92.9)</td>
<td>85.4 (80.5 to 91.2)</td>
<td>0.47</td>
<td>88.1 (82.2 to 92.8)</td>
<td>85.3 (80.2 to 91.2)</td>
</tr>
<tr>
<td>BMI</td>
<td>27.4 (24 to 29)</td>
<td>28.3 (26 to 30)</td>
<td>27.7 (26 to 30)</td>
<td>0.82</td>
<td>28.3 (25 to 30)</td>
<td>27.7 (26 to 30)</td>
</tr>
<tr>
<td>Gravide (median, range)</td>
<td>2 (1–10)</td>
<td>2 (1–8)</td>
<td>2 (1–8)</td>
<td>0.11</td>
<td>2 (1–8)</td>
<td>2 (1–8)</td>
</tr>
<tr>
<td>Parity (median, range)</td>
<td>1 (0–6)*</td>
<td>0 (0–4)</td>
<td>1 (0–6)</td>
<td>0.011</td>
<td>0 (0–4)</td>
<td>1 (0–6)</td>
</tr>
<tr>
<td>FEV1</td>
<td>3.2 (3.0 to 3.4)</td>
<td>3.2 (3.0 to 3.3)</td>
<td>3.0 (2.9 to 3.2)</td>
<td>0.10</td>
<td>3.2 (3.0 to 3.4)</td>
<td>3.0 (2.8 to 3.2)</td>
</tr>
<tr>
<td>Vital capacity</td>
<td>3.9 (3.8 to 4.1)</td>
<td>3.8 (3.6 to 4.0)</td>
<td>3.8 (3.6 to 3.9)</td>
<td>0.70</td>
<td>3.8 (3.6 to 3.9)</td>
<td>3.8 (3.6 to 3.9)</td>
</tr>
<tr>
<td>GC dose, third trimester</td>
<td>0</td>
<td>0</td>
<td>1054.2 (890.7 to 1233.2)</td>
<td>&lt;0.001</td>
<td>234.5 (122.9 to 350.3)</td>
<td>928.0 (705.9 to 1171.5)</td>
</tr>
<tr>
<td>Smoking (%)</td>
<td>3 (5.9%)</td>
<td>14 (26.9%)</td>
<td>20 (28.2%)</td>
<td>0.006</td>
<td>12 (20.7%)</td>
<td>22 (33.8%)</td>
</tr>
</tbody>
</table>

Significance values represent a comparison of each asthma treatment and the control group and of each asthma severity and the control group. Data represent mean (95% CI) unless indicated.
*Significantly higher than the asthma groups.
†Significantly lower than the mild group.
‡Significantly higher than the control and no treatment groups.
§Significantly higher than the control group.
BMI, body mass index; FEV1, forced expiratory volume in 1 s; GC, glucocorticoid.
The effect of glucocorticoid treatment on GR gene activity, mRNA expression and protein levels

The treatment of asthma varied according to asthma severity ($\chi^2=26.73, df=1, p<0.001$), with ICS treatment used by 33% (n=19) of the group with mild asthma and 79% (n=52) of the group with moderate–severe asthma, while the remainder did not treat their asthma with ICS during pregnancy. The use of ICS did not affect the already heightened GR hnRNA expression in the mild asthma group. However, within the moderate–severe asthma group, ICS was associated with significantly lower hnRNA expression compared with the no-treatment group ($t(45)=2.08, p=0.050$), but hnRNA levels comparable with the controls (figure 3). ICS was not associated with changes to GR total mRNA, GRz mRNA expression or GRz protein levels (figure 3).

In the ICS group, both cord blood cortisol and GR hnRNA were positively associated with exon 1A3 expression ($r=0.362, p=0.011$ and $r=0.816, p<0.001$, respectively). In this group, the dose of ICS used in the third trimester was negatively correlated with GR hnRNA expression ($r=-0.578, p<0.001$). In the no-treatment group, no associations were observed between cortisol and hnRNA or exon 1A5 expression. Further, the expression of total GR

Table 2 Characteristics of female and male neonates in each study population according to asthma treatment and asthma severity

<table>
<thead>
<tr>
<th></th>
<th>Control</th>
<th>Mild</th>
<th>Moderate–severe</th>
<th>p Value</th>
<th>Asthma treatment</th>
<th>No treatment</th>
<th>Inhaled corticosteroids</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Females</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>BW (g)</td>
<td>3391.4 (2916.8 to 3520.3)</td>
<td>3468.3 (3233.9 to 3517.8)</td>
<td>3309.0 (3135.4 to 3157.8)</td>
<td>0.34</td>
<td>3483.4 (3321.0)</td>
<td>3262.6 (3079.5 to 3445.8)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>BW centile</td>
<td>47.6 (33.0 to 58.2)</td>
<td>54.5 (40.7 to 62.3)</td>
<td>46.4 (37.2 to 55.6)</td>
<td>0.25</td>
<td>50.2 (44.7 to 64.1)</td>
<td>46.3 (34.7 to 54.1)</td>
<td>0.05</td>
<td></td>
</tr>
<tr>
<td>Length (cm)</td>
<td>51.4 (49.0 to 54.2)</td>
<td>50.9 (49.8 to 52.7)</td>
<td>50.7 (49.4 to 51.7)</td>
<td>0.76</td>
<td>51.4 (50.2 to 52.7)</td>
<td>50.3 (49.1 to 51.5)</td>
<td>0.25</td>
<td></td>
</tr>
<tr>
<td>Length centile</td>
<td>70.2 (51.1 to 100.0)</td>
<td>65.2 (54.9 to 84.6)</td>
<td>63.3 (50.5 to 72.0)</td>
<td>0.76</td>
<td>71.0 (69.1 to 82.8)</td>
<td>59.6 (48.4 to 70.8)</td>
<td>0.18</td>
<td></td>
</tr>
<tr>
<td>Head circumference (cm)</td>
<td>34.6 (33.3 to 36.9)</td>
<td>34.2 (33.2 to 35.3)</td>
<td>34.5 (34.0 to 34.9)</td>
<td>0.79</td>
<td>34.3 (33.7 to 35.2)</td>
<td>34.3 (34.0 to 34.9)</td>
<td>0.74</td>
<td></td>
</tr>
<tr>
<td>Abdominal circumference (mm)</td>
<td>271.6 (238.4 to 264.0)</td>
<td>245.0 (229.0 to 261.2)</td>
<td>284.0 (256.0 to 272.4)</td>
<td>0.37</td>
<td>247.1 (230.6 to 263.6)</td>
<td>264.0 (256.5 to 271.5)</td>
<td>0.37</td>
<td></td>
</tr>
<tr>
<td>Cord blood cortisol (mmol/l)</td>
<td>51.2 (49.0 to 53.7)</td>
<td>52.0 (49.8 to 52.7)</td>
<td>50.7 (49.4 to 51.7)</td>
<td>0.76</td>
<td>51.4 (50.2 to 52.7)</td>
<td>50.3 (49.1 to 51.5)</td>
<td>0.25</td>
<td></td>
</tr>
<tr>
<td>Appar 5</td>
<td>9.3 (8.9 to 9.6)</td>
<td>9.1 (8.8 to 9.4)</td>
<td>9.3 (9.1–9.4)</td>
<td>0.60</td>
<td>9.2 (8.9 to 9.4)</td>
<td>9.3 (8.9 to 9.4)</td>
<td>0.82</td>
<td></td>
</tr>
<tr>
<td>SGA n (%)</td>
<td>2 (9.5)</td>
<td>1 (3.2)</td>
<td>9 (22.5)*</td>
<td>0.049†</td>
<td>3 (9.7)</td>
<td>7 (17.5)</td>
<td>0.54</td>
<td></td>
</tr>
<tr>
<td>Males</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>BW (g)</td>
<td>3642.0 (3374.9 to 3803.1)</td>
<td>3782.7 (3555.4 to 4010.1)</td>
<td>3501.2 (3323.1 to 3706.9)</td>
<td>0.06</td>
<td>3816.7 (3633.6 to 4007.9)</td>
<td>3565.8 (3217.3 to 3637.5)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>BW centile</td>
<td>61.9 (50.1 to 72.4)</td>
<td>64.2 (50.0 to 75.6)</td>
<td>55.3 (45.4 to 65.2)</td>
<td>0.31</td>
<td>61.8 (53.2–75.2)</td>
<td>56.4 (41.6 to 63.5)</td>
<td>0.72</td>
<td></td>
</tr>
<tr>
<td>Length (cm)</td>
<td>52.3 (50.7 to 53.7)</td>
<td>53.2 (52.9 to 54.9)</td>
<td>51.3 (50.0 to 52.5)*</td>
<td>0.038‡</td>
<td>53.5 (52.0 to 55.0)</td>
<td>51.9 (50.7 to 53.1)</td>
<td>0.49</td>
<td></td>
</tr>
<tr>
<td>Length centile</td>
<td>83.1 (70.3 to 93.7)</td>
<td>94.5 (90.8 to 98.2)</td>
<td>88.5 (65.9 to 82.4)*</td>
<td>0.015‡</td>
<td>92.0 (85.4 to 98.5)</td>
<td>75.2 (62.6 to 86.3)</td>
<td>0.36</td>
<td></td>
</tr>
<tr>
<td>Head circumference (cm)</td>
<td>34.1 (33.3 to 35.3)</td>
<td>35.3 (34.6 to 36.4)</td>
<td>34.9 (34.1 to 35.7)</td>
<td>0.86</td>
<td>35.2 (33.9 to 36.1)</td>
<td>35.3 (34.5 to 35.9)</td>
<td>0.90</td>
<td></td>
</tr>
<tr>
<td>Abdominal circumference (mm)</td>
<td>261.6 (256.1 to 267.1)</td>
<td>264.1 (255.7 to 282.5)</td>
<td>286.7 (263.8 to 281.6)</td>
<td>0.16</td>
<td>264.1 (256.3 to 272.0)</td>
<td>271.1 (262.9 to 279.2)</td>
<td>0.21</td>
<td></td>
</tr>
<tr>
<td>Cord blood cortisol (mmol/l)</td>
<td>50.4 (47.0 to 53.7)</td>
<td>52.0 (49.8 to 52.7)</td>
<td>50.7 (49.4 to 51.7)</td>
<td>0.76</td>
<td>51.4 (50.2 to 52.7)</td>
<td>50.3 (49.1 to 51.5)</td>
<td>0.25</td>
<td></td>
</tr>
<tr>
<td>Appar 5</td>
<td>9.5 (9.2 to 9.9)</td>
<td>9.1 (8.9 to 9.4)</td>
<td>9.1 (9.1–9.4)</td>
<td>0.17</td>
<td>9.2 (9.0 to 9.5)</td>
<td>8.1 (8.8 to 9.4)</td>
<td>0.11</td>
<td></td>
</tr>
<tr>
<td>SGA n (%)</td>
<td>2 (6.6)</td>
<td>1 (3.7)</td>
<td>2 (7.7)</td>
<td>0.82</td>
<td>1 (4.5)</td>
<td>2 (6.5)</td>
<td>0.94</td>
<td></td>
</tr>
</tbody>
</table>

Reported p values represent significance level of univariate analysis of variance of asthma severity or asthma treatment group compared with the control group. Data represent mean (95% CI).

*Significant compared with both the control and mild asthma groups (p<0.05).

†Significant main effect of asthma severity.

BW, birth weight; SGA, small for gestational age.
GR mRNA, GRα mRNA and GRα protein levels did not differ between the ICS and no-treatment groups. The dose of ICS used in the third trimester did not correlate with cortisol, GRα mRNA expression, GRα protein levels or birth weight centiles.

**The effect of fetal sex on GR gene activity, mRNA expression and protein levels**

Expression levels of total GR mRNA and GRα mRNA in control and asthmatic pregnancies differed according to fetal sex. In females, a significantly higher abundance of total GR mRNA was demonstrated in placentae from the control compared with the asthmatic groups irrespective of asthma severity (F(1,54)=4.99, p=0.003). A similar pattern of abundance was observed with GRα mRNA in the placentae from females; however, this failed to reach statistical significance (p=0.08) (figure 4). In males, the expression of the total placental GR mRNA and GRα mRNA transcripts was unchanged with asthma. Protein levels of GRα were significantly higher in placentae from males compared with females (t(54)=2.385, p=0.022; figure 4), but did not change with asthma. Placental GR hnRNA and GR 1A3 expression were not independently affected by fetal sex.

The associations between placental expression of hnRNA, 1A3 and cortisol that were observed with ICS were demonstrated to be sex specific. In placentae from males, hnRNA expression was positively correlated with cortisol in both the control and ICS groups (r=0.594, p=0.005 and r=0.474, p=0.003, respectively) but not the no-treatment group. Expression of exon 1A3 was also correlated with cortisol in the ICS group (r=0.630, p=0.002). Expression of both hnRNA and exon 1A3, however, was unaffected by the dose of ICS used in the third trimester. In placentae from females, no correlation was observed between hnRNA expression, exon 1A3 and cortisol in any group. However, third
trimester dose of ICS was negatively associated with hRNA and exon 1A3 expression ($r=-0.523$, $p=0.004$ and $r=-0.415$, $p=0.028$, respectively).

**The effect of cigarette smoking on placental GR gene activity, mRNA expression and protein levels**

Smoking was associated with a significant decrease in female birth weight centiles in pregnancies complicated by asthma (smoker, mean=29.24, SEM=6.00; non-smoker, mean=55.82, SEM=5.76; $t(69)=3.808$, $p<0.001$); smoking did not affect male birth weight centiles. Cigarette smoking was, however, not associated with changes in GR gene activity, mRNA transcripts or protein levels in placenta from either male or female births (data not shown).

**DISCUSSION**

The findings of this work indicate sex-specific alterations to placental GR regulation, cortisol and growth in the presence of asthma. Specifically, female but not male birth weight centiles were inversely associated with cord blood cortisol levels in the presence of maternal asthma; however, concurrent changes to placental GR protein were not evident. These data suggest that decreased female fetal growth is not due to an increase in glucocorticoid sensitivity via increased GR protein levels per se.

The comparable GR protein levels observed in placenta from males and females in both the asthmatic and control groups should predict equal sensitivity to cortisol and similar downstream functional responses. In the males, this held true, with no differences in birth weight centiles apparent between the asthmatic or control groups. However, in the female placentae, GR protein levels were not predictive of birth weight outcomes. In females, low birth weight centiles occurred with the highest cortisol levels that characterised the group with untreated, moderate−severe asthma. Maternal smoking further reduced female birth weight centiles in this group, while not affecting placental GR protein levels. These findings suggest a role for an alternative cortisol-mediated pathway in the regulation of female fetal growth. A likely candidate is the insulin-like growth factor (IGF) axis, as increased cortisol is associated with decreased female fetal growth in a high cortisol environment, as do the downstream functional effects typically associated with cortisol binding (such as growth and placental cytokine inhibition).

Despite significant reductions in GR mRNA in female placentae in the asthma compared with the control groups, no change in GR protein levels were observed. This may be accounted for by differences in GR mRNA stability or, alternatively, may result from preferential transcription of other GR isoforms, resulting in differential glucocorticoid sensitivity. We have previously demonstrated decreased mRNA expression of two placental GR isoforms, GR$_A$ and GR$_B$, following in vitro treatment of placentae with the synthetic glucocorticoid dexamethasone, while GR$_B$ was unaltered.

Given that the GR-P isoform appears to exist in upregulating GR activity, decreased GR-P expression in the presence of asthma-induced increases in cortisol may result in reduced placental glucocorticoid sensitivity. The evidence for glucocorticoid-specific effects on GR isoform-specific transcription is conflicting and appears to be tissue specific. An increased ratio of GR$_B$ to GR$_A$ protein has been demonstrated in dexamethasone-treated nasal polyp tissue in vitro, while decreased ratios are observed following glucocorticoid exposure of bronchial epithelial cells. Further, increased GR$_B$ to GR$_A$ protein ratios have been demonstrated in cases of glucocorticoid insensitivity in a range of inflammatory conditions (asthma, nasal polyps and ulcerative colitis).

The sex from which the donor tissue was derived was not included in these studies, but may in part explain the varied results.

The present study has provided insight into sex-specific regulation of the GR in placentae of pregnancies complicated by asthma, characterised by increased endogenous cortisol and the use of exogenous glucocorticoids. This provides a platform to examine further the regulation of placental GR and its downstream effects on cortisol signalling, and may further elucidate determinants of glucocorticoid sensitivity, contributing to our understanding of sex-specific differences in fetal growth in utero. Scientifically, these findings necessitate that any further studies addressing tissue responses to corticosteroids should take into consideration the sex from which the tissue was derived in interpretation of the results. Clinically,
the findings of this work indicate that male and female growth responses to maternal pathology and/or corticosteroid exposure are sex specific. This in itself warrants further study, as it may have important implications for the clinical management of observed fetal growth alterations in pregnancies complicated by asthma.

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Competing interests PG has received expense reimbursement for lectures given in educational symposia sponsored by GlaxoSmithKline, AstraZeneca and Novartis. He has conducted research sponsored by GlaxoSmithKline and Pharmaxis. The other authors declare no competing interests.

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