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In humans, early cortisol biosynthesis provides a mechanism to safeguard female sexual development

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In humans, sexual differentiation of the external genitalia is established at 7–12 weeks post conception (wpc). During this period, maintaining the appropriate intrauterine hormone environment is critical. In contrast to other species, this regulation extends to the human fetal adrenal cortex, as evidenced by the virilization that is associated with various forms of congenital adrenal hyperplasia. The mechanism underlying these clinical findings has remained elusive. Here we show that the human fetal adrenal cortex synthesized cortisol much earlier than previously documented, an effect associated with transient expression of the orphan nuclear receptor nerve growth factor IB-like (NGFI-B) and its regulatory target, the steroidalogenic enzyme type 2 3β-hydroxysteroid dehydrogenase (HSD3B2). This cortisol biosynthesis was maximal at 8–9 wpc under the regulation of ACTH. Negative feedback was apparent at the anterior pituitary corticotrophs. ACTH also stimulated the adrenal gland to secrete androstenedione and testosterone. In concert, these data promote a distinctive mechanism for normal human development whereby cortisol production, determined by transient NGFI-B and HSD3B2 expression, provides feedback at the anterior pituitary to modulate androgen biosynthesis and safeguard normal female sexual differentiation.

Introduction
The differentiation of male or female phenotype consistent with chromosomal sex is a key component of development. In humans, sexual dimorphism of the external genitalia is established during a critical phase of the first trimester in response to male hormones from the fetal testis. During this period, overexposure of the 46,XX female fetus to these androgens causes virilized genitalia at birth. This balance between cortisol biosynthesis and potential androgenesis after 12 wpc (3), earlier immunoreactivity in the human fetal anterior pituitary (6) has never been correlated to adrenal function. Indeed, anencephalic fetuses, which lack the anterior pituitary, have normal appearing adrenals during the first trimester (7).

For these reasons, we sought to clarify the potential for the adrenal cortex to participate in cortisol and androgen biosynthesis during the first trimester of human development. In this study, we determined that adrenocortical activity correlated to that of the developing anterior pituitary. Our findings, linked to transient expression of the key steroidalogenic enzyme HSD3B2 and its regulatory orphan nuclear receptor, nerve growth factor IB-like (NGFI-B) (8), support a distinctive developmental mechanism for cortisol biosynthesis in humans that protects normal female sexual differentiation.

Results
The human adrenal cortex is distinct from the bipotential gonad at 33 days post conception (dpc) (9, 10), after which it can be dissected free from surrounding structures, including the kidney (Figure 2A). The adrenal cortex grew rapidly during the early fetal period. Its wet weight increased almost 10-fold from 8 wpc to 10 wpc (Figure 2B). At 41 dpc CD34-positive vascular channels
penetrated the embryonic gland between the aorta and mesonephros, and vessel density later increased at the adrenocortical periphery (Figure 2, C and D).

The precise onset of adrenocortical function has not been defined, either by steroid secretion or expression profile of the biosynthetic enzymes (Figure 1). By immunohistochemistry (IHC) as shown in Figure 3, steroid acute regulatory protein (StAR), CYP11A, CYP17, and CYP21 were first detected robustly at 50–52 dpc within the nascent inner fetal zone (Figure 3, B, G, L, and Q). Immunoreactivity in this location was also detected using an antibody that recognizes both CYP11B1 and CYP11B2 (Figure 3V). Using specific primers and sequencing, RT-PCR amplified products from both CYP11B1 and CYP11B2 transcripts (data not shown). Within the outer definitive zone, StAR, CYP11A, CYP21, and CYP11B1/CYP11B2 were more weakly detected and, in line with our previous investigations, CYP17 appeared largely absent (Figure 3L) (10). In later specimens up to 14 wpc, these profiles persisted in the fetal zone; however, the definitive zone also stained weakly positive for CYP17 (Figure 3, M, N, and O).

For de novo cortisol biosynthesis from cholesterol, the expression of HSD3B2 is necessary (Figure 1). In all previous studies restricted to later material, this protein has only been detected reliably after 22 weeks’ gestation (3, 5), where it is regulated by the orphan nuclear receptor NGFI-B (8). Like other adrenocortical enzymes, HSD3B immunoreactivity was absent at 41 dpc; yet surprisingly, positive cells were apparent at 50–52 dpc, mostly at the interface between the definitive and fetal zones (Figure 4, A and B). Expression was particularly abundant and more widespread in specimens at 8–9 wpc, with many cells demonstrating HSD3B immunoreactivity that declined at later time points until no protein could be detected at 14 wpc (Figure 4, C–F). IHC for NGFI-B on neighboring sections revealed a remarkably similar profile of nuclear expression (primary antibodies to HSD3B and NGFI-B were both raised in rabbit prohibiting colocalization studies; Figure 4, G–L).

For all the IHC experiments, expression was indistinguishable between male and female specimens. The data were also supported by sequenced RT-PCR products for StAR, CYP11A, CYP17, CYP21, and HSD3B isoforms using intron-spanning primers (Figure 5A). The HSD3B immunoreactivity correlated solely to expression of the type 2 isoform HSD3B2 (Figure 5B).

Our findings on enzyme expression predicted cortisol biosynthesis. Compared with the kidney, immunoaassays for cortisol, validated by gas chromatography/mass spectrometry (Supplemental Methods and Supplemental Figure 1; supplemental material available online with this article; doi:10.1172/JCI25091DS1), revealed 9- to 18-fold higher cortisol content in the adrenal gland during the first trimester (Figure 6A). Although this steep gradient could reflect differential metabolism, adrenal expression of type 11β-hydroxysteroid dehydrogenase (HSD11B2), which inactivates cortisol to cortisone, exceeded that in the early kidney (data not shown). Cortisol content per adrenal weight dropped by approximately 50% between 8 and 10 wpc (Figure 6A), consistent with the waning profile of HSD3B2 expression (Figure 4).

The response of paired first-trimester adrenal glands to agonist was investigated in male and female specimens by overnight cul-
tue and media assay for cortisol. Ex vivo culture of human adrenocortical cells at mid-gestation has artificially induced HSD3B2 (3). RT-PCR before and after culture ensured that this was not the case in our experiments (Supplemental Figure 2). Forskolin, which mimics the effect of ACTH, stimulated an 11-fold increase in adrenal cortisol content at 8 and 9 wpc compared with only 7.5-fold at

Figure 3
Steroidogenic enzyme and STAR expression in the developing human adrenal gland. Brightfield IHC at sequential developmental ages counterstained by toluidine blue with antibodies to STAR (A–E), CYP11A (F–J), CYP17 (K–O), CYP21 (P–T), and CYP11B1/CYP11B2 (U–Y). Dotted ring in A illustrates extent of the adrenal cortex in left panels (A, F, K, P, and U). In other images, the definitive zone (DZ) is oriented to the left and the fetal zone (FZ) to the right. At 14 wpc, these 2 zones were separated by an additional transitional zone (TZ). Scale bar: 300 μm.

Figure 4
HSD3B and NGFI-B immunoreactivity in the developing human adrenal gland. Brightfield IHC with antibodies to HSD3B (A–F) and NGFI-B (G–L) counterstained with toluidine blue. Dotted rings in A and G illustrate extent of the adrenal cortex. In other images, the definitive zone is oriented to the left and fetal zone to the right. Scale bar: 300 μm.
10 wpc (data not shown). Very similar results were obtained from the media, showing significantly diminished cortisol secretion at 10 wpc (Figure 6B). These data defined maximal cortisol production at 8–9 wpc, when the adrenal gland was also responsive to ACTH (Figure 6C) and expressed its receptor, the type 2 melanocortin receptor (MC2R; Figure 6D).

Collectively, these experiments confirmed ACTH-responsive cortisol biosynthesis by the early human adrenal cortex. We used the same fetal collection to ask whether there was coincident maturity of anterior pituitary corticotrophs to secrete ACTH. The developing pituitary was negative for ACTH immunoreactivity at 41 dpc (data not shown). However, identical to the onset of adrenocortical enzyme expression, cytoplasmic ACTH was detected at 50–52 dpc (Figure 7B), which overlapped with the more widespread expression of nuclear glucocorticoid receptor (GR; Figure 7C; note that colocalization studies were not possible as both antibodies were raised in mice). Increased ACTH expression was detected in the anterior pituitary at 8 wpc (Figure 7D), with similar findings at 10 wpc (data not shown). Overnight in vitro culture at 8 wpc demonstrated that these early corticotrophs secreted abundant ACTH, which was suppressible by dexamethasone (Table 1).

Finally, with the onset of major differentiation of the external genitalia at 8 wpc (Figure 8), we determined the capacity of the fetal adrenal gland to secrete androgens. Androstenedione and testosterone were assayed from the media of paired adrenal

![Figure 5](http://www.jci.org/articles/view/25091)

**Figure 5**
RT-PCR analysis of adrenocortical enzymes and StAR at 8 wpc. (A) RT followed by 22 and 28 cycles of PCR for transcripts encoding steroidogenic enzymes. (B) Specific identification of HSD3B2 and HSD3B1 isoforms in the presence (+) and absence (−) of RT. No HSD3B1 transcript was detected in the adrenal sample after 42 cycles of PCR. Positive controls were fetal testis (HSD3B2) and skin (HSD3B1); negative controls were H2O and genomic DNA (G).

![Figure 6](http://www.jci.org/articles/view/25091)

**Figure 6**
Cortisol content and secretion from the early human adrenal cortex. (A) Cortisol content (mean ± SEM) per mg adrenal tissue at 8, 9, and 10 wpc, with 8 wpc kidney as control. (B) Cortisol secretion (mean ± SEM) of paired adrenal glands stimulated by 10 μM forskolin at 8, 9, and 10 wpc. (C) Cortisol secretion (mean ± SEM) of paired adrenal glands at 8 wpc in response to varying doses of ACTH(1–24). Statistical analyses of stimulated secretion compared to basal secretion achieved the same level of significance (***P < 0.005) for each ACTH dose. *P < 0.05; **P < 0.02. (D) PCR amplification of type 2 melanocortin receptor (MC2R) in the presence (+) and absence (−) of RT in the adrenal gland, testis, and ovary at 8 wpc with GAPDH control.
glands cultured overnight in the presence or absence of ACTH, forskolin, or dexamethasone. Dexamethasone failed to alter androstenedione and testosterone secretion significantly, both of which were stimulated approximately 3-fold in male and female adrenal glands by forskolin and, to a lesser extent, ACTH (Table 2). In contrast, neither ACTH-stimulated coculture of adrenal with fetal testis or ovary nor ACTH stimulation of gonad alone augmented androgen production (data not shown). The conversion of androstenedione to testosterone relied on 17β-hydroxysteroid dehydrogenase (HSD17B) activity. By RT-PCR, the type 3 isoform, HSD17B3, responsible for testosterone biosynthesis in the testis, was only very weakly detected in the first-trimester adrenal cortex (Figure 8C). The type 5 enzyme, encoded by the AKR1C3 gene, was more readily detected. The detection of both androstenedione and testosterone was validated by mass spectrometry (Supplemental Figure 1).

Discussion

Sexual differentiation of the external genitalia is a fundamental aspect of human development, tied by the phenotype of CYP21 deficiency to the fetal adrenal cortex. The nature and timing of this event are important. From the second trimester onward, adrenocortical steroids such as dehydroepiandrosterone can act as precursors for androgen biosynthesis (3, 11). However, by this stage, the female fetus is relatively well protected, as high placental aromatase activity converts this androgen to estrogen (12). In contrast, sexual dimorphism is established during a critical period in the first trimester, when a relative lack of aromatase permits normal androgen-mediated male sexual differentiation (see Figure 8) but conversely renders the female fetus susceptible to virilization (11). We have investigated this key period of human endocrine development.

Adrenal enzyme expression by 50–52 dpc included those that are precursors for androgen biosynthesis, notably CYP17, CYP11A, and CYP11B (3, 11). These enzymes are expressed in the fetal adrenal cortex at 50 dpc and are upregulated by ACTH stimulation (3). However, the relative lack of ACTH production in the first trimester suggests that adrenal androgen biosynthesis is not regulated by the anterior pituitary. The transient presence of NGFI-B expression at 50 dpc suggests that NGFI-B would determine steroidogenic preference by regulating fetal adrenal androgen biosynthesis (3). This hypothesis has been supported by the expression of NGFI-B in fetal adrenal glands at 50 dpc (3). In contrast, neither ACTH-stimulated coculture of adrenal with fetal testis or ovary nor ACTH stimulation of gonad alone augmented androgen production (data not shown). The conversion of androstenedione to testosterone relied on 17β-hydroxysteroid dehydrogenase (HSD17B) activity. By RT-PCR, the type 3 isoform, HSD17B3, responsible for testosterone biosynthesis in the testis, was only very weakly detected in the first-trimester adrenal cortex (Figure 8C). The type 5 enzyme, encoded by the AKR1C3 gene, was more readily detected. The detection of both androstenedione and testosterone was validated by mass spectrometry (Supplemental Figure 1).

Table 1

<table>
<thead>
<tr>
<th>ACTH content</th>
<th>Control</th>
<th>Basal</th>
</tr>
</thead>
<tbody>
<tr>
<td>ACTH secretion in vitro from the anterior pituitary at 8 wpc</td>
<td>&lt; 1.1 pmol/l/16 h</td>
<td>410 ± 7.4 pmol/l/16 h</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Inhibition by dexamethasone</th>
<th>5 nM</th>
<th>1 μM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Percentage decrease in media ACTH upon exposure to dexamethasone</td>
<td>32.34% ± 13.0%</td>
<td>51.9% ± 2.5%</td>
</tr>
</tbody>
</table>

Control, media alone; Basal, after overnight culture of anterior pituitary. Percentage decrease in media ACTH upon exposure to dexamethasone was standardized against untreated anterior pituitary explant. Normal upper limit of circulating ACTH in human adults is <9 pmol/l. Results shown as mean ± SEM from at least 2 experiments.
Taken together, these data reveal what we believe to be novel functioning of the early adrenal cortex and reinforce distinctive aspects of human development (perhaps including other higher primates). Our observations suggest a delicate balance to early female differentiation that is vulnerable to androgen before the protective appearance of placental aromatase (12). However, the early cortisol biosynthesis, facilitated by transient expression of adrenocortical NGFI-B and HSD3B2, would inhibit ACTH production by the anterior pituitary corticotroph. This negative feedback would minimize ACTH-driven androgen secretion to produce a transient mechanism that safeguards the major period of female sexual development (Figure 9). In support of this model, loss of cortisol feedback by inactivation of the GR has previously been associated with disrupted female sexual differentiation and ambiguous genitalia at birth (although the patient also carried 1 disrupted allele of the CYP21 gene, this would be insufficient by itself to generate an intersex phenotype; ref. 24).

Our future experiments will expand upon this developmental model by seeking to understand the regulation and identify the molecular consequences of this early adrenal steroid biosynthesis.

**Methods**

Collection of human embryonic and fetal material. Ethical approval for these studies was granted by the Southampton and South West Hampshire Local Research Ethics Committee. The collection and staging of human embryonic and fetal material was carried out with informed consent as described previously (25–27) using the Carnegie classification and fetal foot length to provide a direct assessment of gestational age as dpc or wpc. Organs from 121 fetuses were analyzed. Tissue preparation for gene expression studies has been described previously (10).

IHC. IHC was performed as described previously using trypsin or sodium citrate antigen exposure techniques (25). Sections were incubated with primary antibody (optimal dilution shown in Table 3) overnight at 4°C and with biotinylated secondary antibodies for 2 hours at 4°C according to the manufacturer’s instructions (Vector Laboratories). Following incubation for 1 hour at room temperature with streptavidin–horseradish peroxidase conjugate (1:200; Vector Laboratories), the color reaction was developed with diaminobenzidine containing 0.1% H2O2, and slides were counterstained with toluidine blue. For brightfield dual IHC, the Vector Red kit was used according to the manufacturer’s instructions (Vector Laboratories). All antibodies to steroidogenic enzymes have been reported previ-

### Table 2

<table>
<thead>
<tr>
<th>Androstenedione secretion</th>
<th>Mean ± SEM</th>
<th>Fold increase</th>
<th>P</th>
</tr>
</thead>
<tbody>
<tr>
<td>Basal</td>
<td>29.3 ± 2.77 nmol/l</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Forskolin</td>
<td>117 ± 13.5 nmol/l</td>
<td>4.00 ± 0.11</td>
<td>&lt; 0.005</td>
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<tr>
<td>ACTH</td>
<td>35.0 ± 1.73 nmol/l</td>
<td>1.20 ± 0.03</td>
<td>&lt; 0.05</td>
</tr>
<tr>
<td>DEX</td>
<td>22.9 ± 3.40 nmol/l</td>
<td>0.78 ± 0.08</td>
<td>NS</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Testosterone production</th>
<th>Mean ± SEM</th>
<th>Fold increase</th>
<th>P</th>
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<tbody>
<tr>
<td>Basal</td>
<td>1.15 ± 0.10 nmol/l</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>Forskolin</td>
<td>3.33 ± 0.33 nmol/l</td>
<td>2.90 ± 0.29</td>
<td>&lt; 0.002</td>
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<tr>
<td>ACTH</td>
<td>2.50 ± 0.10 nmol/l</td>
<td>2.17 ± 0.09</td>
<td>&lt; 0.001</td>
</tr>
<tr>
<td>DEX</td>
<td>1.05 ± 0.23 nmol/l</td>
<td>0.91 ± 0.20</td>
<td>NS</td>
</tr>
</tbody>
</table>

Androstenedione and testosterone were not detected in control media. The response to 10 μM forskolin, 100 nM ACTH (1–24), and 1 μM dexamethasone (DEX) was tested.
overnight in the same media. Anterior pituitary was cultured explants. Following initial over-gland, dissected from the development, glucocorticoid feedback, the cell line NCI-H295R (37, 38). Human adrenocortical tumor (basal analysis). For analysis of cross-reactivity was limited to human adrenocortical tumor, androgenic tumors. For analysis of included omission of primary or secondary antibody. The ACTH antibody was raised against amino acids 1–24 of ACTH [ACTH(1–24)]. Potential cross-reactivity was minimal: 5.4% for testosterone with 5α-dihydrotestosterone, 1.49% for androstenedione with testosterone, and less than 1% for other known androgenic steroids. This media composition has proved optimal in our experience for the main maintenance of phenotype of the human adrenocortical tumor cell line NCI-H295R (37, 38). Anterior pituitary was cultured overnight in the same media (basal analysis). For analysis of glucocorticoid feedback, the gland, dissected from the developing sella turcica, was divided into the midline for culture as 2 explants. Following initial overnight culture and media ACTH assay, fresh media was added containing either vehicle or dexamethasone (Sigma-Aldrich) for a second overnight incubation. Media ACTH assay results were expressed as percent inhibition following the addition of dexamethasone standardized against the untreated sample.

**Steroid extraction, hormone assay, and verification.** For organ steroid analysis, weighed adrenal glands were ground in liquid nitrogen, suspended in phosphate-buffered saline (pH 7.2), and extracted in dichloromethane (2.0 ml). After brief centrifugation, the organic phase was isolated and extracts were dried under air at 25°C and resuspended in assay diluent (250 μl). Media steroid and ACTH assays were performed directly on 0.5 ml aliquots from culture. Hormones were assayed according to the manufacturer’s instructions using Immulite 2000 (ACTH, androstenedione and cortisol; Diagnostic Products Corporation) and ADVIA Centaur (testosterone; Bayer) automated chemiluminescent immunoassays. The ACTH assay targets 2 sites and only detects intact ACTH (amino acids 1–39). Cortisol, testosterone, androstenedione, and ACTH were undetected in control media samples. Assay cross-reactivity was minimal: 5.4% for testosterone with 5α-dihydrotestosterone, 1.49% for androstenedione with testosterone, and less than 1% for other known androgenic steroids. Gas chromatography/mass spectrometry was used to verify the steroid immunoassays (Supplemental Figure 1).

**Statistics.** Steroid assays were analyzed using the 2-tailed Student’s t test. Adrenal steroid experiments were conducted at least 3 times at each dose and age on either paired or pooled organs. ACTH was measured on 2 (dexamethasone studies) or 3 occasions (basal studies). Results are presented as mean ± SEM.

**Acknowledgments**

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**Table 3**

<table>
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<tr>
<td>Polyclonal anti-CYP11B1/2</td>
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**Figure 9**

Schematic diagram of human early adrenal function and its implications for androgen-mediated development.

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