Intra-adrenal factors are not involved in the differential control of cortisol and adrenal androgens in human adrenals

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Abstract

The differential control of adrenal androgens and cortisol may be due to intra-adrenal factors, which may be age- or sex-related, or due to extra-adrenal factors, such as circulating hormones. The purpose of this study was to identify any intrinsic differences that may exist in steroidogenic production occurring within adrenals obtained from males and females, and any maturational differences that may evolve with age. Using human adrenals from 48 transplant donors (32 males, 16 females; ages 5–60 years), the influences of age and sex on basal production of and ACTH-stimulated cortisol, androstenedione and dehydroepiandrosterone (DHEA) were examined in freshly prepared adrenal cell suspensions. Basal and ACTH-stimulated cortisol, androstenedione and DHEA production were similar in adrenals from males and females and did not correlate significantly with age when the whole group was examined. When steroidogenesis in male and female adrenals was examined separately against age, a significant correlation was observed only for basal and ACTH-stimulated androstenedione in adrenals from males in the younger age group, 5–30 years (basal: \( r = 0.84, P = 0.0001 \); ACTH-stimulated: \( r = 0.52, P = 0.007 \)). Examination of the relationships between the steroids disclosed that the basal and ACTH-stimulated cortisol/androgen ratios did not correlate significantly with age, but the androstenedione/DHEA ratio showed a significant direct relationship with age in males only (basal: \( r = 0.53, P = 0.006 \); ACTH-stimulated: \( r = 0.5, P = 0.01 \)).

These data suggest that the influences of sex and age are minor in the modulation of adrenal steroidogenesis and support the concept that extra-adrenal factors dominate in the differential modulation of adrenal androgens and cortisol. The relationship between the androstenedione/DHEA ratio and increasing age in men is consistent with the recently reported stimulatory effect of testosterone on adrenal steroidogenesis by induction of the conversion of DHEA to androstenedione.

European Journal of Endocrinology 138 567–573

Introduction

Whereas it is well established that cortisol production is under the control of adrenocorticotropic hormone (ACTH), it has been known for some time that adrenal androgens are regulated in a more complex manner (1). Adrenal androgens are controlled partly, but not exclusively, by ACTH. Several physiological and pathological situations exist in which there is a divergence of adrenal androgen and cortisol secretion (1–3). Cortisol concentrations are increased in stress and anorexia nervosa, whereas androgen concentrations are suppressed (4). In some patients with partial hypopituitarism, cortisol concentrations are normal and androgen concentrations are suppressed (5). Patients with ectopic ACTH syndrome have increased cortisol concentrations, whereas androgen concentrations are increased, normal or decreased (6). Circulating adrenal androgens demonstrate a striking and unexplained relationship with age. Adrenarche is characterized by an increase in blood concentrations of adrenal androgens at the end of the first decade of life. Serum concentrations of adrenal androgens peak at 20–30 years of age and remain at this level throughout the 4th and 5th decade. With ageing, the concentration of adrenal androgens and their response to ACTH decline (7, 8). This pattern of adrenal androgen concentrations is seen in both sexes (9) and differs from that of adrenal glucocorticoid, as serum cortisol concentrations and cortisol responsiveness to ACTH remain relatively constant throughout life (8). When ACTH is administered to hypophysectomized chimpanzees, it is adequate to maintain cortisol production but, given alone, it is unable to maintain a normal cortisol/androgen ratio (10). Therefore the evidence suggests that some non-ACTH mechanism, either alone or in combination with ACTH, is involved in the control of adrenal androgens (5).

The control of adrenal androgens may be through the agency of ‘intra-adrenal’ or ‘extra-adrenal’ factors. The potential endogenous adrenal gland mechanisms regulating adrenal androgen secretion include maturational and zonation of the adrenal gland (11), adrenal...
blood flow that may expose the inner zone to high concentrations of cortisol (12), and changes in the properties of the adrenal enzymes and their co-factors (13–15). Alternatively, factors outside the adrenal gland have been proposed, such as prolactin (16), growth hormone (17), insulin (18), sex steroids (19, 20) and fragments of the ACTH precursor molecule, pro-opiomelanocortin, such as β-endorphin (21) and joining peptide (21, 22). The existence of such a cortical androgen stimulating hormone has been disputed (23–25). The present study was designed to examine the influence of ageing and sex on the relative production of cortisol, androstenedione and dehydroepiandrosterone (DHEA), basally and in response to ACTH, and thus enable us to examine the influence of possible endogenous maturational changes that may influence differential production of cortisol and androgens.

**Materials and methods**

**Subjects**

Human adrenals were obtained at the time of kidney resection from 48 brain dead transplant donors, 32 males aged 28.8 ± 11.3 years and 16 females aged 37.3 ± 21.7 years (mean ± S.D.). The large majority of the donors had suffered subarachnoid haemorrhage or head injuries. Before kidney resection, some donors had been pretreated with dexamethasone and desmopressin (DDAVP) or with dopamine and antibiotics. None of the donors had suffered subarachnoid haemorrhage or head injuries. Before kidney resection, some donors had been pretreated with dexamethasone and desmopressin (DDAVP) or with dopamine and antibiotics. None of the donors had had chronic underlying disease that might be identified as arising from the zona reticularis. Cells from the zona fasciculata were recognized by their irregular shape and abundant lipid droplet content; these cells comprised approximately 50% of the adrenal cell count. Aliquots of cell suspension, 800 µl, were added to plastic culture tubes (NUNC 0–43141) that contained 100 µl EMEM (basal) or ACTH(1–39) (Peninsula Laboratories, Liverpool, England). The final volume was made up to 1 ml with EMEM. The tubes containing cells and additions were incubated in a shaking water-bath under 100% oxygen for 2 h at 37°C. Cells were prepared on 48 separate occasions from 48 pairs of human adrenals. On each occasion, cells were incubated in the absence of ACTH (basal conditions) and in the presence of ACTH 10⁻¹⁰ mol/l (ACTH-stimulated conditions) in three to five replicate tubes. When steroid production was assessed in these replicate tubes, the variability was usually less than 15%. If large discrepancies occurred, the results of the assay were rejected. Preincubation tubes that contained cells without any added peptide were removed before incubation and stored at 4°C. At the end of incubation, the cell suspensions were frozen and the intracellular steroids were released into the medium by repeated freezing and thawing. The steroids were measured by specific radioimmunoassay. The protocol was similar to that previously used for guinea-pig adrenal cells (26). The cells were capable of responding to ACTH with a significant increase in production of cortisol, androstenedione and DHEA. In order to standardize steroid production from one cell preparation to another, the values obtained were adjusted for the number of cells present by expressing steroid production as pmol/10⁶ cells.

**Steroid measurement**

Cortisol was measured by direct radioimmunoassay using antiserum supplied by Guildhay (Guilford, Surrey, England), cortisol-3-(O-carboxymethyl)-oximo [2⁻¹²⁵ I]-iodohistamine (Amersham Int., Amersham, England) and phosphate buffer 0.5 mol/l, pH 7.4. Aliquots of cell lysates and standards were incubated in the presence of antibody and radiolabelled cortisol overnight at 4°C. Unbound cortisol was separated from antibody-bound cortisol using 1% dextran–gelatin-coated charcoal, followed by centrifugation at 3100 r.p.m. for 15 min. The supernatant was decanted and the radioactivity in the free fraction (the charcoal pellet), was measured using a gamma counter. Androstenedione and DHEA concentrations were...
measured by radioimmunoassay after extraction of steroids with diethyl ether and further purification by chromatography over celite (27). The androstenedione assay used androstenedione-10-[125I]-iodine derivative (Diagnostic Products UK Ltd, Abingdon, Oxfordshire, England) as the radiolabel and the antiserum was obtained from Radioassay Systems Labs (RSL, Carson, USA; Lot No. 1440 R1/P). The DHEA assay used dehydro-[1, 2, 6, 7-3H]-epiandrosterone, (Amersham; code TRK 157) as radiolabel; the antiserum was obtained from ICN Biomedicals Inc. (Diagnostics Division, Costa Mesa, CA 92629, USA, Lot R1–46–21). The specificities, sensitivities and precision of the assays have been reported previously by us (26).

To identify changes in the production of the steroids relative one to another, we utilized androstenedione/cortisol, DHEA/cortisol and androstenedione/DHEA ratios in both basal and ACTH-stimulated cell suspensions.

**Statistical analysis**

Results are reported as mean ± S.E. Steroid production in the male and female groups was compared using the Mann–Whitney U non-parametric test. The correlation of each group with age was assessed by using regression analysis. A P value < 0.05 was considered to be statistically significant.

**Results**

**Cortisol and androgen production in human adrenals in vitro**

Cortisol, androstenedione and DHEA production was similar in cells from adrenal glands obtained from males and females under basal or ACTH-stimulated conditions (Fig. 1). However, when the basal and ACTH-stimulated androstenedione/cortisol, DHEA/cortisol and androstenedione/DHEA ratios were compared between males and females, the ACTH-stimulated androstenedione/cortisol ratio (male 0.31 ± 0.13, female 0.07 ± 0.06; P < 0.05) and androstenedione/DHEA ratio (male 2.7 ± 0.05, female 1.5 ± 0.88; P < 0.05) were significantly greater in the males. When basal and ACTH-stimulated cortisol, androstenedione and DHEA production was examined against age in the total group, no significant correlation was observed (Table 1). However, analysis of values obtained with male and female adrenals separately revealed that, in the male glands, basal and ACTH-stimulated androstenedione concentrations correlated significantly with age (r = 0.41, P = 0.02; r = 0.47, P = 0.007; Fig. 2), although cortisol or DHEA did not correlate significantly with age in the group (Table 1). Basal and ACTH-stimulated cortisol, androstenedione and DHEA did not significantly correlate with age in the female group (Table 1). When the male group was divided into almost
equal numbers of specimens from younger (less than 30 years, \( n = 17 \)) and older (30 years and over, \( n = 15 \)) individuals, it was observed that the significant correlation of androstenedione with age was confined to the younger subgroup (basal: \( r = 0.84, P = 0.0001 \); ACTH-stimulated: \( r = 0.50, P = 0.0089 \); Fig. 3). No significant correlation was observed when androstenedione was examined against age in adrenals from subjects aged 30–60 years (basal: \( r = 0.29, P = 0.36 \); ACTH-stimulated: \( r = 0.21, P = 0.417 \)).

The possible effects of pretreatment of the donors on the subsequent in vitro steroidogenesis of adrenal cells was examined by subdividing the donors for whom adequate information was available into three groups: those treated with dexamethasone and DDAVP (\( n = 14 \)), those treated with dopamine and antibiotics (\( n = 15 \)) and those who received no pretreatment (\( n = 7 \)). There were no significant differences in basal or ACTH-stimulated cortisol, androstenedione or DHEA production by adrenal cells between the three groups of donors. Furthermore, there were no significant correlations of steroid production with age in any of the three groups.

**Cortisol, androstenedione and DHEA interrelationships**

In the male group, basal and ACTH-stimulated androstenedione/DHEA ratios correlated significantly with age (\( r = 0.53, P = 0.006; r = 0.50, P = 0.001 \) respectively; Fig. 4), but the DHEA/cortisol or androstenedione/cortisol ratios did not (Table 2). In the female adrenals, the basal and ACTH-stimulated androstenedione/DHEA, DHEA/cortisol and androstenedione/cortisol ratios all failed to correlate significantly with age (Table 2). When the androstenedione/DHEA ratio in males was examined for the two age categories (5–30 years and 30 years and over), the basal androstenedione/DHEA ratio in the younger group correlated significantly with age (\( r = 0.6, P = 0.0175 \)), but the ACTH-stimulated androstenedione/

<table>
<thead>
<tr>
<th>Correlation coefficient (r)</th>
<th>Cortisol</th>
<th>Androstenedione</th>
<th>DHEA</th>
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<tbody>
<tr>
<td></td>
<td>Basal</td>
<td>ACTH-stimulated</td>
<td>Basal</td>
</tr>
<tr>
<td>All</td>
<td>0.23</td>
<td>0.21</td>
<td>0.22</td>
</tr>
<tr>
<td>Male</td>
<td>0.36</td>
<td>0.189</td>
<td>0.41*</td>
</tr>
<tr>
<td>Female</td>
<td>0.114</td>
<td>0.346</td>
<td>0.135</td>
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</tbody>
</table>

Significant correlations: * \( P = 0.02 \), ** \( P = 0.007 \).

Figure 2 Basal and ACTH-stimulated androstenedione production plotted against age in male human adrenal cells in vitro (\( n = 32 \)). There was a statistically significant correlation of both basal (correlation coefficient \( r = 0.41, P = 0.02 \)) and ACTH-stimulated androstenedione production (\( r = 0.47, P = 0.007 \)) with age in male adrenal cells which was not seen in female adrenal cells.
DHEA in the younger group and basal and ACTH-stimulated androstenedione/DHEA in the older group did not correlate with age.

Discussion

Some authors have suggested that the increase in adrenal androgen production at adrenarche is due to some maturational change within the adrenal gland itself (11–13). It has been reported that the zona reticularis, the innermost zone of the adrenal cortex that predominantly secretes adrenal androgens, appears at the time of adrenarche (11). However, no reciprocal decrease in the size of the zona reticularis occurs during the 5th decade of life, when the adrenal androgen concentrations begin to decline. Another
Table 2 Correlation coefficients for the relationship of basal and ACTH-stimulated androstenedione (Andro)/DHEA, DHEA/cortisol and androstenedione/cortisol ratios with age (years) in male and female human adrenal cells.

<table>
<thead>
<tr>
<th>Group</th>
<th>Correlation coefficient (r)</th>
<th>Andro/DHEA</th>
<th>Andro/cortisol</th>
<th>DHEA/cortisol</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Basal ACTH-stimulated Basal ACTH-stimulated Basal ACTH-stimulated</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>0.53* 0.50**</td>
<td>0.383</td>
<td>0.355</td>
<td>0.23 0.199</td>
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<tr>
<td>Female</td>
<td>0.297 0.269</td>
<td>0.253</td>
<td>0.022</td>
<td>0.46 0.04</td>
</tr>
</tbody>
</table>

Significant correlations: ** P = 0.001, * P = 0.006.

possible mechanism is that of long-term exposure of the zona reticularis to cortisol as a consequence of blood flow through the adrenal from the outer cortex to the inner, which may bring about an alteration in the enzymes involved in adrenal androgen production (23). It has also been suggested that maturational changes in the activities of some adrenal enzymes (17-hydroxylase, 17–20-lyase and the 3β-hydroxysteroid dehydrogenase. Δ4–5 isomerase system) may account for alterations in steroidogenesis favouring androgens (13–15). Alternatively, factors extrinsic to the adrenal may specifically control adrenal androgen biosynthesis and secretion – for example, the putative cortical androgen stimulating hormone. Several studies support this concept. Using our human adrenal cell preparations, we have reported that joining peptide and β-endorphin, peptides arising from the ACTH precursor pro-opiomelanocortin, have androgen-stimulating properties (21). Parker et al. (22) also reported stimulatory effects of joining peptide on adrenal androgen production, but others have not confirmed this (24, 25). Evidence also exists that gonadal steroids influence adrenal glucocorticoid and androgen secretion: for example, testosterone increases the adrenal response to ACTH stimulation (19, 20). A 60 000 molecular weight glycoprotein isolated from human pituitary has been reported to stimulate DHEA from males younger than 30 years, as a result of increased 3β-hydroxysteroid-dehydrogenase, Δ4–5 isomerase enzyme system activity. The normal increase in blood concentrations of testosterone from testicular origin in young men appears to enhance adrenal conversion of DHEA to androstenedione.

While it is theoretically possible that the pretreatment of the donors might effect subsequent steroidogenesis in the adrenal cells, we observed no differences between cortisol or androgen production in adrenal cells from donors who had no pretreatment and those pretreated with dexamethasone and DDAVP or dopamine and antibiotics. The lack of any effect of donor pretreatment on subsequent steroidogenesis suggests that any such influence on the adrenal is mediated via an acute extra-adrenal mechanism. Furthermore, the specific steroidogenic pattern seen in the adrenals from young men only was not obscured either by stress or by pretreatment of donors. Thus the model used provides useful new insights into the differential control of adrenal glucocorticoid and androgen production. Adrenal androgen production did not correlate with age or sex, with the exception of androstenedione in...
younger male adrenals, suggesting that factors exogenous to the adrenal gland are more important in the control of adrenal androgens than are intrinsic changes. Indeed, it is possible that the correlation of androstenedione production with age in young male adrenals was due to an increasing in vivo effect of testicular testosterone on adrenal 3β-hydroxysteroid dehydrogenase. Δ4→3 isomerase enzyme system activity. Thus our data support the concept that it is ‘extra-adrenal’ rather than ‘intra-adrenal’ factors that dominate in the differential modulation of androgens and cortisol production. However, the identity of this ‘extra-adrenal’ factor has yet to be elucidated.

Acknowledgements

The authors wish to thank the Transplant Co-Ordination Team in Beaumont Hospital, Dublin, Ireland, for their assistance in obtaining human adrenal tissue. This work was performed with the aid of a grant from the Health Research Board of Ireland.

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Received 29 June 1997
Accepted 22 January 1998