The influence of sex and gonadectomy on the hypothalamo–pituitary–adrenal axis of the sheep

B J Canny, K A O'Farrell, I J Clarke¹ and A J Tilbrook

Department of Physiology, Monash University, Clayton, Victoria, 3168 Australia

¹Prince Henry's Institute of Medical Research, PO Box 5152, Clayton, Victoria, 3168, Australia

(Requests for offprints should be addressed to B J Canny)

Abstract

There is a sex difference in the hypothalamo–pituitary–adrenal (HPA) axis of many species, although there are sparse data on the sheep. In the present study we have compared the HPA axes of intact and gonadectomised adult male and female sheep at the level of the median eminence, pituitary and adrenal glands using a variety of *in vitro* approaches. The concentration of arginine vasopressin (AVP) was higher ($P < 0.01$) in the median eminence of male than female sheep, and was also elevated by gonadectomy of either sex ($P < 0.01$). The concentration of corticotrophin-releasing factor (CRF) in the median eminence did not differ between the sexes, but was also elevated in both sexes following gonadectomy ($P < 0.01$). Anterior pituitary pro-opiomelanocortin mRNA concentrations were higher ($P < 0.05$) in intact male sheep than in intact females, with the levels in gonadectomised animals of both sexes being intermediate. In contrast to this finding, basal ACTH secretion from anterior pituitary cells was higher ($P < 0.05$) in cultures derived from female sheep than those from males, but gonadectomy was without effect. There was no effect of sex or gonadectomy on *in vitro* ACTH secretion in response to AVP, CRF or the combination of AVP and

CRF, and in all cases the combination of AVP and CRF generated greater ($P < 0.0001$) ACTH secretion than AVP alone. AVP alone was more effective ($P < 0.01$) than CRF alone as an ACTH secretagogue. The adrenal glands were larger ($P < 0.05$) in female than male sheep, with no effect of gonadectomy. Basal cortisol production was greatest ($P < 0.05$) in cultures of adrenal cells from intact male sheep, though ACTH- and 8Br cAMP-induced cortisol production was greater in the cultures of cells from females ($P = 0.05$); there were no effects of gonadectomy. Cultures of adrenocortical cells from male sheep had greater ($P < 0.05$) basal cAMP production, but ACTH-stimulated cAMP production did not differ between any of the groups of animals. These findings show a range of differences in the HPA axis of male and female sheep. Furthermore, they suggest that the heightened activity of the axis in the female occurs primarily due to differences at the level of the adrenal gland, and that greater adrenal responsiveness of female animals is due to differences in the latter stages of steroidogenesis, rather than an effect on ACTH signal transduction at its receptor.

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Introduction

It is well established for various species that there are sex differences in the hypothalamo–pituitary–adrenal axis (HPA), and that these differences occur at virtually all levels of the axis. There are sex-related differences in: the levels of Type I and II glucocorticoid receptors in the hippocampus and hypothalamus (Turner & Weaver 1985, Carey *et al.* 1995, MacLusky *et al.* 1996); the expression of corticotrophin-releasing factor (CRF) and arginine vasopressin (AVP) in the hypothalamus (Vamvakopoulos & Chrousos 1993, Paulmyer-Lacroix *et al.* 1996, Viau & Meaney 1996); the secretion of adrenocorticotrophin (ACTH) from the anterior pituitary (Gallucci *et al.* 1993); and the size and steroidogenic potential of the adrenal

glands (Kitay 1961, Gaskin & Kitay 1970, Roelfsma *et al.* 1993, El-Migdadi *et al.* 1995). It is thought that the differences observed between males and females reflect differential effects of the sex steroids on the HPA axis, as the activity of many proteins which regulate the axis are directly regulated by the steroids and sex steroid receptors are present at many levels of the axis (Hirst *et al.* 1992, Herbison 1995, Bethea *et al.* 1996, Madigou *et al.* 1996). In particular, oestrogen appears to have a stimulatory action on the HPA axis, and hence the axis of the female exhibits generally greater reactivity than that of the male (Handa *et al.* 1994, Young 1995). Precise physiological or pathological explanations for this increased activity in the female are, however, lacking, but it has been recently hypothesised that the differences in the

HPA axes may explain sex-related differences in psychological and immunological phenomena (Da Silva 1995, Young 1995).

Although it is clear that the HPA axis of the sheep may be regulated by sex-related factors (Bell *et al.* 1991, Keller-Wood & Wood 1991, Pecins-Thompson & Keller-Wood 1994, Schwartz & Rose 1998), few formal comparisons have been made between males and females of this species. Despite the availability of sophisticated *in vitro* culture techniques for a number of years, these approaches have rarely been used to dissect the differences between the HPA axes of the two sexes of any species. We are also unaware of any studies where *in vitro* techniques have been used to simultaneously examine the effects of sex and gonadectomy (GDX) on different levels of the HPA axis. Accordingly, our knowledge of the way that sex and sex hormones regulate the HPA axis is limited, and few studies have examined the underlying cellular mechanisms that may be the determining factors. The aim of the current study was to compare the HPA axes of male and female sheep at the level of the median eminence, anterior pituitary and adrenal gland. Furthermore, we examined the effects of long-term GDX in both sexes.

Materials and Methods

Materials

Unless otherwise stated, all materials were from Sigma Chemical Co. (St Louis, MO, USA).

Experimental design

This experiment compared male and female sheep that had either been left intact or had undergone GDX up to 10 months previously. The concentrations of immunoreactive (ir)-AVP and ir-CRF in the median eminence, and pro-opiomelanocortin (POMC) mRNA in the anterior pituitary were measured by RIA and Northern blot analysis respectively. Primary cultures of anterior pituitary and adrenocortical cells were established, and the following parameters were measured: basal and AVP-, CRF- and AVP plus CRF-induced ir-ACTH secretion; basal and ACTH- and 8-Br cAMP-induced cortisol production; and basal and ACTH-induced cAMP production.

Animals

All animal procedures were conducted with prior institutional ethical approval under the requirements of the Australian Prevention of Cruelty to Animals Act 1986 and the NH&MRC/CSIRO/AAC *Code of Practice for the Care and Use of Animals for Scientific Purposes*.

Seven rams and nine ewes of the Romney Marsh breed were used, and the animals were gonad-intact, 3 years of

age and weighed between 47 and 69 kg. Animals were held in paddocks at a latitude of 37° 49' south, and had free access to pasture and water. Ten months prior to the experiment, four rams and three ewes underwent GDX using standard surgical techniques, with another two ewes undergoing GDX 6 weeks before the experiment. GDX and tissue collection were conducted during the breeding season for this breed and latitude (Bremner *et al.* 1984). Between 7 and 10 days prior to the experiment, the animals were transported to the Monash University Department of Physiology Animal House (approximately 50 km), where they were kept in individual pens with free access to food and water (ambient temperature 22 °C, 12 h:12 h light:darkness cycle, lights on 0700 h). Animals were killed between 0900 and 1000 h, using a lethal i.v. injection of pentobarbitone sodium (Lethobarb, 80 mg/kg; Virbac (Australia), Peakhurst, Australia). At autopsy the median eminence, anterior pituitary gland and the left adrenal gland were removed; the median eminence and a longitudinal hemisection of the anterior pituitary were immediately frozen in liquid nitrogen and stored at -70 °C until further analysis, while the other section of the pituitary and the left adrenal gland were collected into ice-cold PBS containing 5×10^{-4} M MgCl₂.

Pituitary culture

Primary culture of anterior pituitary cells was performed using established techniques (Familar *et al.* 1989). In brief, single pituitary sections (approximately one half of an anterior pituitary) were diced to small cubes (approximately 1 mm³), and treated with trypsin (25 mg/hemipituitary) (Gibco, Grand Island, NY, USA) and DNase (40 µg/ml) (Boehringer Mannheim, Mannheim, Germany) in Dulbecco's modified essential medium (DMEM) (JRH Biosciences, Lenexa, KS, USA). After a single cell suspension was obtained, cells were washed in DMEM, containing 10% foetal calf serum (FCS) (CSL Pty Ltd, Melbourne, Victoria, Australia), 2×10^{-2} M Hepes, 100 µg/ml streptomycin and 100 U/ml penicillin (Gibco), and plated at a density of 2×10^5 cells/well in 1 ml of this medium in 24-well culture dishes (Linbro, Flow Laboratories, McLean, VA, USA). This procedure routinely yielded $7-10 \times 10^6$ cells/hemipituitary, with a cell viability of greater than 90%, as assessed by the trypan blue exclusion test. Cells were kept at 37 °C in a humidified 5% CO₂ incubator for 72 h prior to treatment. On the morning of treatment, cells were washed extensively with serum-free DMEM containing 0.1% Polypep and allowed to equilibrate for 1 h prior to treatment with various concentrations of AVP (Auspep, West Melbourne, Victoria, Australia), ovine CRF (Auspep) or AVP in the presence of CRF (10^{-9} M) for 3 h. All treatments were conducted in triplicate. Medium was collected at the end of this period, frozen and stored at -20 °C for later

ACTH analysis. After the medium was removed the plates were frozen on dry ice, then kept for subsequent DNA measurement.

Adrenal culture

Primary cultures of adrenal cortical cells were obtained using established techniques (Canny *et al.* 1998). Following removal of the adrenal capsule, the cortex was separated from the medulla, and diced finely. The cortical cells were dispersed in medium 199 (Gibco) containing 2×10^{-2} M Hepes, 100 µg/ml streptomycin, 100 U/ml penicillin and 0.1% Polypep (treatment medium) and 0.2% collagenase I (Worthington, Freehold, NJ, USA) in a waterbath at 37 °C for 2 h, with gentle agitation. Every 30 min the suspension was passed through a siliconised wide-bore Pasteur pipette for 5 min. The final suspension was put through a Teflon mesh filter (70 µm), washed repeatedly in medium 199 containing 2×10^{-2} M Hepes, 100 µg/ml streptomycin, 100 U/ml penicillin, and 10% FCS, and the cells then plated at a density of 2×10^5 cells/well in 1 ml of this medium in 24-well tissue culture plates. The cell viability was greater than 90% as assessed by trypan blue exclusion. Cells were cultured for 72 h at 37 °C, in 5% CO₂ in a humidified incubator and were extensively washed with treatment medium prior to treatment. The cells were preincubated for 1 h prior to treatment with the appropriate experimental reagent for 3 h. All treatments were conducted in triplicate. When cAMP responses to ACTH₁₋₂₄ (Synacthen; Ciba-Geigy, Pendle Hill, NSW, Australia) were being assessed, 10^{-3} M 1-methyl-3-isobutylxanthine was added to the treatment medium. After 3 h the medium was removed and stored at -20 °C until assay. The plates were snap frozen on dry ice, and then stored at -20 °C for subsequent DNA determination. Cortisol production from these cultures in response to angiotensin II (AII) and phorbol myristate acetate has recently been reported (Canny *et al.* 1998).

Extraction of peptides from the median eminence

AVP and CRF were extracted from the median eminence by homogenisation in 2 ml ice-cold 2 M acetic acid. After sitting on ice for 20 min, the homogenate was spun at 15 000 g for 30 min at 4 °C. The supernatant was removed, divided into aliquots and lyophilised and stored at -20 °C until assay. At this time, an aliquot was taken and resuspended in the appropriate buffer. Extraction efficiency, as assessed by recovery of ¹²⁵I-labelled peptides was greater than 90%.

AVP assay

AVP was measured using a heterologous double-antibody RIA with a sheep antiserum (S45) raised against AVP

conjugated to keyhole limpet haemocyanin. The antiserum was used at a final dilution of 1:240 000 in a 5×10^{-2} M PBS containing 0.25% BSA and 0.02% NaN₃, using ¹²⁵I-AVP as tracer and AVP as a standard. The assay demonstrates a crossreactivity of 100% for lysine vasopressin, and less than 0.01% for AVP-gly, Arg8 vasotocin and oxytocin, when compared with AVP. Extracts of median eminence reconstituted in assay buffer diluted in parallel to the standard curve for AVP. The sensitivity of the assay was routinely 3.9 pg/tube. Donkey anti-sheep IgG serum (1:800 dilution in 0.5 ml 2% polyethylene glycol (PEG)) was used as the precipitating antibody. All the samples were assayed in one assay and the intra-assay coefficient of variation of five quality control standards was 6.3%.

CRF assay

CRF was measured using a heterologous double-antibody RIA with a rabbit antiserum (R3) raised against a peptide (Cys-His-Ser-Asn-Arg-Lys-Leu-Leu-Asp-Ile-Ala-NH₂; Chiron Mimotopes, Clayton, Victoria, Australia) conjugated to keyhole limpet haemocyanin via the N-terminal cysteine residue. This peptide is based upon the ten C-terminal amino acids of ovine CRF. The antiserum was used at a final dilution of 1:120 000 in 10^{-1} M PBS containing 0.5% BSA and 0.1% NaN₃, using ¹²⁵I-[Tyr⁰]-ovine CRF (Auspep) as tracer and ovine CRF as standard. The assay crossreacts 100% with [Tyr⁰]-ovine CRF, and less than 0.1% with ACTH₁₋₃₉, AVP and rat/human CRF. Extracts of median eminence reconstituted in assay buffer diluted in parallel to the standard curve for CRF. The sensitivity of the assay was routinely 3.9 pg/tube. Goat anti-rabbit IgG serum (1:150 dilution in 0.5 ml 2% PEG) was used as the precipitating antibody. All the samples were assayed in one assay and the intra-assay coefficient of variation of five quality control standards was 11.9%.

Anterior pituitary mRNA assays

Total RNA was extracted from the hemipituitary glands as previously described (Mercer *et al.* 1989) and 25 µg total RNA were run for Northern blotting. The RNA was hybridised with a ³²P-labelled ovine POMC probe (van der Pavert *et al.* 1997). The RNA was also hybridised with a ³²P-labelled glyceraldehyde 3-phosphate dehydrogenase GAPDH probe and the expression of the POMC mRNA was standardised against the density of the GAPDH mRNA probe using laser densitometry.

ACTH assay

ACTH was measured using a heterologous double-antibody RIA with a rabbit antiserum (R7) raised against ACTH₁₋₂₄ conjugated to keyhole limpet haemocyanin. This antiserum was used at a final dilution of 1:80 000 in

a buffer containing 5×10^{-2} M Tris-HCl, 5×10^{-4} M EDTA, 10^{-2} M *n*-ethyl maleimide, 2% PEG, 0.01% Triton X-100, 0.2% gelatine, pH 7.4, using ^{125}I -ACTH₁₋₃₉ as tracer and ACTH₁₋₃₉ as standard. The sensitivity of the assay was routinely 2 pg ACTH₁₋₃₉/tube. The antiserum crossreacts 500% with ACTH₁₋₂₄, 10% with POMC, 5.2% with CLIP (ACTH₁₈₋₃₉) and less than 0.1% with α -melanocyte-stimulating hormone (ACTH₁₋₁₃.NH₂) and β -endorphin, when compared with ACTH₁₋₃₉ (100%). Goat anti-rabbit IgG serum (1:150 dilution in 0.5 ml 2% PEG) was used as the precipitating antibody. Medium from anterior pituitary primary culture diluted in parallel to the standard curve. All samples were assayed in duplicate, and the assay had intra- and interassay coefficients of variation of 5.6 and 8.7% respectively ($n=7$).

cAMP assay

Following freezing of the cells, cAMP was extracted using acid/ethanol (10^{-2} M HCl in 95% ethanol). The cAMP RIA was performed as described by Marley *et al.* (1991) with the exception that the bound cAMP was separated utilising a second antibody (goat anti-rabbit) at 1/150 dilution in 2% PEG in a 4 °C overnight incubation, followed by centrifugation ($3000 \times g$ 4 °C, 30 min), aspiration and the pellet counted using a Packard Bell γ -counter. The sensitivity of the assay was routinely less than 5 fmol/tube, the ED₅₀ 133.9 ± 9.4 fmol/tube ($n=6$) and the intra- and interassay coefficients of variation were 8.6% ($n=10$) and 9.8% ($n=6$) respectively.

Cortisol assay

Cortisol production by adrenal cultures was measured by RIA as previously described (Bocking *et al.* 1986), with the exception that samples were not extracted as medium alone did not interfere with the assay. In brief, samples were assayed in duplicate using [$1,2,6,7\text{-}^3\text{H}(\text{N})$]cortisol (10 000 c.p.m., Amersham, Boston, MA, USA) and a sheep antiserum raised against 4-pregnen-11 β ,17 α , 21-triol-3,20-dione-21-acetate 3 CMO-BSA (#3368, donated by Dr R I Cox, CSIRO, NSW, Australia). Bound radioactivity was separated following the addition of 20% PEG and centrifugation (1800 g 15 min), and counted by liquid scintillation counting using a Beckman β -counter. The crossreactivity of various steroids (relative to cortisol) are as follows: cortisone (20.5%), 17 α -hydroxyprogesterone (3.9%), corticosterone (1.0%), progesterone (0.6%) and 11-deoxycortisol (<0.1%). The sensitivity of the assay was 2.1 ± 0.1 ng/ml ($n=6$) and the intra- and interassay coefficients of variation were 9.9 and 13.8% ($n=6$).

DNA and protein assays

DNA was extracted from the culture plates by sonication for 15 min in 2 M NaCl, 2×10^{-3} M EDTA,

5×10^{-2} M Na₂PO₄ buffer. The extracts were measured using the fluorochrome Hoechst 33258 at a concentration of 0.11 $\mu\text{g}/\text{ml}$ as previously described (Labarca & Paigen 1980). The interassay coefficient of variation was 12.6% ($n=7$). Median eminence protein content was determined using a BioRad (Hercules, CA, USA) assay, with BSA as a standard.

Statistics

CRF and AVP concentrations were corrected for the protein concentration of the median eminence, while ACTH secretion and cortisol and cAMP production by pituitary and adrenal cultures, respectively, were corrected for DNA content of the cells. In all cases, the results were compared using a multifactorial ANOVA procedure, with sex, GDX and the interaction term sex by GDX as main factors, with an error term of animal effect nested within sex and GDX. Data were analysed following assessment of homogeneity of variances by Bartlett's and Cochran's tests; where necessary, data were log or square root transformed prior to analysis. A repeated measures procedure was used for the *in vitro* secretion studies. As significant between-animal effects were present in all analyses involving agonist-induced secretion *in vitro*, secretion was expressed as a ratio of basal secretion for that culture. In all cases results are expressed as means $\pm$ S.E.M., and a comparison of the four groups is presented. Differences between individual means were, however, not determined unless a statistically significant interaction between the main factors (sex and GDX) was present. A probability level of 5% was considered significant.

Results

Hypothalamus

The concentrations of ir-AVP and ir-CRF in the median eminence of gonad-intact and GDX male and female sheep are shown in Fig. 1. Concentrations of ir-AVP were higher ($P<0.01$) in the median eminence of male sheep than in female sheep (Fig. 1A), and were also elevated ($P<0.01$) in both sexes following GDX (Fig. 1A). In contrast, the concentrations of ir-CRF were similar in the median eminence of male and female sheep (Fig. 1B), but there was a significant elevation ($P<0.01$) in the concentration of the peptide following GDX of both sexes (Fig. 1B).

Anterior pituitary

The levels of the mRNA for POMC, relative to that of GAPDH, in the anterior pituitaries of intact and GDX male and female sheep are shown in Fig. 2. There was a

Figure 1 The concentration of ir-AVP (A) and ir-CRF (B) in the median eminence of intact female ($n=4$), GDX female ($n=5$), intact male ($n=3$) and GDX male ($n=4$) sheep. ir-AVP and ir-CRF concentrations were higher ($P<0.01$) in GDX than intact sheep of both sexes, while ir-AVP concentrations were higher ($P<0.01$) in male than in female sheep.

significant ($P<0.05$) interaction between the sex and GDX with respect to POMC expression, such that the expression was higher ($P<0.05$) in intact males than intact females, with that of GDX animals of both sexes being intermediate between, and not different from either of the gonad-intact groups.

Basal secretion of ir-ACTH *in vitro*, after correction for DNA content of the cultures, was higher ($P<0.05$) in cultures of cells obtained from female sheep when compared with those from males (Table 1); there was, however, no effect of GDX. Secretion of ir-ACTH in response to CRF (10^{-11} – 10^{-7} M, Fig. 3A), AVP (10^{-11} – 10^{-7} M, Fig. 3B) and AVP (10^{-10} – 10^{-7} M) in combination with CRF (10^{-9} M, Fig. 3C) was determined relative to basal secretion in each culture. Following this correction, there were no effects of the sex or gonadal status of the animal from which the culture was derived on ir-ACTH responses to any treatment. For each treatment (CRF or AVP alone, and AVP plus CRF), there was a significant ($P<0.01$) concentration effect of the treatment. AVP alone was more effective ($P<0.01$) than CRF alone in inducing ir-ACTH secretion (Fig. 3A and

Figure 2 Levels of expression of mRNA for POMC in the anterior pituitary gland in intact female ($n=4$), GDX female ($n=5$), intact male ($n=3$) and GDX male ($n=4$) sheep. The relative optical density was determined by laser densitometry after correction for the intensity of hybridisation of a probe for mRNA of GAPDH. Bars which do not share a letter (a or b) are significantly ($P<0.05$) different from each other.

B), with the combination AVP plus CRF being more effective ($P<0.001$) than either peptide alone (Fig. 3C). This hierarchy of potencies was not affected by sex or GDX.

Adrenal gland

The adrenal glands were proportionately heavier ($P<0.05$) in female sheep than males, although there was no effect of GDX (Table 1). Basal cAMP production was higher ($P<0.05$) in cultures from male sheep (Table 1), but not different between the animals of different gonadal status. ACTH_{1–24}-induced increases in cAMP production, relative to basal production, were not different between cultures from male or female, or gonad-intact and GDX sheep (Fig. 4). Basal cortisol production was highest in adrenal cultures derived from gonad-intact male sheep, being significantly ($P<0.05$) greater from these cultures than those from either gonad-intact females or GDX males (Table 1). Basal cortisol production from the cultures obtained from GDX females was not different from that of any other group. Overall, the relative increase in cortisol production in response to ACTH_{1–24} was greater ($P=0.05$) in cultures from female than male sheep, and tended ($P=0.06$) to be greater in cultures from gonad-intact compared with GDX animals of both sexes (Fig. 5). In all groups of animals, ACTH_{1–24} induced

Table 1 Differences in adrenal weight, basal pituitary ACTH secretion and basal adrenal cortisol and cAMP production in intact and GDX male and female sheep

	Intact female	GDX female	Intact male	GDX male
Basal ACTH secretion (pg/ng DNA) ^A	24.1 ± 4.0 (n=4)	24.6 ± 2.4 (n=5)	15.6 ± 4.4 (n=3)	7.3 ± 1.7 (n=4)
Left adrenal mass (mg/kg body weight) ^A	44.3 ± 3.9 (n=4)	48.5 ± 3.2 (n=5)	32.4 ± 3.7 (n=3)	40.7 ± 3.2 (n=4)
Basal cortisol production (ng/ng DNA)	7.4 ± 0.7 ^a (n=4)	8.4 ± 0.6 ^{a,b} (n=4)	10.4 ± 0.4 ^b (n=3)	7.5 ± 0.8 ^a (n=4)
Basal cAMP production (pmol/ng DNA) ^B	0.54 ± 0.05 (n=4)	0.59 ± 0.07 (n=4)	0.94 ± 0.08 (n=3)	0.76 ± 0.07 (n=4)

^ASignificant effect of sex on this parameter ($P<0.05$), with females being greater.
^BSignificant effect of sex on this parameter ($P<0.05$), with males being greater.
^a and ^b show the statistical significance of interaction between the sex and gonadal status of the animals; values which do not share a superscript are significantly different ($P<0.05$) from each other.

Figure 3 ACTH secretion from anterior pituitary cultures from intact female ($n=4$), GDX female ($n=5$), intact male ($n=3$) and GDX male ($n=3$) sheep, in response to ovine CRF (A), AVP (B), or AVP in the presence of CRF (10^{-9} M) (C), expressed as a ratio of the basal ACTH secretion in individual cultures. The sex or presence of gonads had no effect on the ACTH secretory response to any secretagogue.

concentration-dependent increases ($P<0.001$) in cortisol production (Fig. 5). The relative increase in cortisol production in response to 8BrcAMP was greater ($P<0.05$) in cultures from female sheep, but there were no significant effects of the GDX on the ability of 8BrcAMP to induce cortisol production (Fig. 6).

Discussion

We have shown that there are differences between female and male sheep at all levels of the HPA axis, and that long-term GDX can have various effects on the differences between the sexes. These findings represent, to our

Figure 4 Increases in cAMP production above basal by sheep adrenal cortical cultures in intact female ($n=4$), GDX female ($n=4$), intact male ($n=3$) and GDX male ($n=4$) sheep in response to ACTH. There was no significant effect of the sex or gonadal status of the animal from which the culture was derived on cAMP production.

knowledge, the first comparison of the effects of both sex and gonadal status on stimulated secretion *in vitro* of ACTH by the anterior pituitary cells and cortisol production by adrenocortical cells, and suggest that sex differences exist at the level of the adrenal gland but not the pituitary gland. Our observations that adrenocortical cells of female animals have a greater steroidogenic potential and that this increase appears to occur at a post-receptor level, extend the findings of increased adrenal function in females of a number of other species (for review see Handa *et al.* 1994, El-Migdadi *et al.* 1995, Nowak *et al.* 1995). AVP and CRF concentration in the median eminence, anterior pituitary POMC mRNA concentration and basal ACTH secretion *in vitro*, and basal adrenocortical cell cortisol and cAMP production, were variously affected by sex and GDX, and these observations reinforce the value of examining a number of levels of the HPA axis and the complex regulation of this system.

The concentration of both AVP and CRF in the median eminence was influenced by GDX, and in the case of AVP differences existed between male and female animals. AVP concentrations were found to be higher in males, and also following GDX. These findings are consistent with some, but not all observations, in other

Figure 5 ACTH-induced increases in cortisol ('F') production above basal in adrenal cortical cultures of intact female ($n=4$), GDX female ($n=4$), intact male ($n=3$) and GDX male ($n=4$) sheep. Cortisol production was greater in cultures from female sheep ($P=0.05$) than male sheep, and tended to be greater in cultures from intact sheep ($P=0.06$) than GDX sheep.

species. While some studies have shown no effect of sex or gonadal status on hypothalamic AVP mRNA concentrations or the number of AVP-immunoreactive neurones in the rat (Crowley & Amico 1993, Bingaman *et al.* 1994, Chisari *et al.* 1995, Patchev *et al.* 1995), it has been shown that median eminence AVP concentrations are higher in vehicle-treated GDX male rats when compared with testosterone-replaced animals (Viau & Meany 1996). In contrast, testosterone may have a stimulatory effect on the number of AVP neurones in the hypothalamus of rats and pigs (van Eerdenburg *et al.* 1991, De Vries *et al.* 1994). The reason for the difference in the findings of these studies is not readily apparent, though variations in the physiological state of the animals may play a role. It has been demonstrated that while GDX *per se* did not affect the concentration of AVP mRNA in male or female rat hypothalami, the presence of sex steroids was essential to allow the normal increase in expression seen in response to an increase in plasma osmolarity (Crowley & Amico 1993).

It is also clear that the gonads can regulate the levels of CRF in the hypothalamus, though the effect again appears to depend upon the specific experimental paradigm. The human CRF gene has putative oestrogen

Figure 6 8BrcAMP-induced increases in cortisol (F) production above basal in adrenal cortical cultures of intact female ($n=4$), GDX female ($n=4$), intact male ($n=3$) and GDX male ($n=4$) sheep. Cortisol production was greater in cultures from female sheep ($P<0.05$) than male sheep, but there was no effect of the gonadal status of the animal from which the culture was derived.

response elements in the 5' promoter region, and these have been associated with an increase in CRF gene expression by oestradiol *in vitro* (Vamvakopoulos & Chrousos 1993, Paulmyer-Lacroix *et al.* 1996). The effects of oestradiol *in vivo* on CRF expression, however, appear to depend on complex interactions involving adrenal glands. Paulmyer-Lacroix *et al.* (1996) have shown that oestradiol treatment *in vivo* inhibits CRF expression in intact rats, but increases the expression in adrenalectomised animals. These observations most probably reflect the fact that oestradiol has stimulatory actions on adrenal glucocorticoid production (*vide infra*), which, in turn, will inhibit CRF production (Vamvakopoulos & Chrousos 1994, Jingami *et al.* 1985). The effects of testosterone, as deduced from experiments involving GDX and/or *in vivo* replacement regimens, on hypothalamic CRF appear somewhat variable. Inhibitory (Haas & George 1988, Bingaman *et al.* 1994) or no effects (Almeida *et al.* 1992, Shanks *et al.* 1994, Hadid *et al.* 1995, Viau & Meaney 1996) of testosterone on CRF neurones have both been reported, without any clear explanation of the differences between the studies. In the specific case of the sheep, CRF mRNA content in the paraventricular nucleus is not altered during pregnancy, lactation or exogenous oestradiol and progesterone treatment of the female sheep (Broad *et al.* 1995). The amount of CRF mRNA was

elevated immediately after parturition, though it is not clear if this was due to changing levels of gonadal steroids or a reflection of the stress of parturition. While it has been argued that the CRF gene represents an important point of regulation of the sexual dimorphism of the HPA axis (Vamvakopoulos & Chrousos 1993), the available evidence does not unequivocally support this contention.

The content of POMC mRNA was greater in the anterior pituitaries of intact males than intact females, with GDX animals of both sexes having intermediate levels of POMC mRNA. There is a substantial literature describing the regulation of POMC mRNA in the hypothalamus by gonadal steroids, but few reports of the anterior pituitary exist. In the hypothalamus of the rat, oestradiol and testosterone have variable effects on POMC expression (Wilcox & Roberts 1985, Blum *et al.* 1989, Chowen-Breed *et al.* 1989), which appear to depend upon the precise anatomical location that is being studied. In a study of male sheep, testosterone decreased POMC mRNA levels in some, but not all, hypothalamic sites (Hileman *et al.* 1996). In ewes, no difference in anterior pituitary POMC mRNA was reported between intact and GDX animals (Mercer *et al.* 1989), while progesterone and oestradiol treatment appeared to increase anterior pituitary POMC mRNA (Whisnant *et al.* 1992) in GDX ewes. Our present observations may be at odds with a number of studies which report higher ACTH content in the anterior pituitaries of female rats (Lesniewska *et al.* 1990, Chisari *et al.* 1995), though it is not clear if such differences reflect the species studied, or the methodology (mRNA vs peptide content) employed. It is worth noting in the present study, that basal ACTH secretion was greater from the cultures of the anterior pituitary cells of female animals, and this parameter may be a better reflection of anterior pituitary ACTH content.

The sex or gonadal status of the animals from which anterior pituitary cultures were derived had no effect on ACTH secretion in response to CRF, AVP or their combination. As has been previously reported (Familar *et al.* 1989), AVP alone is a more potent ACTH secretagogue than is CRF alone, in sheep anterior pituitary cultures, and this order of effectiveness was not altered by differing sex or gonadal status. AVP has also been reported to be an effective *in vivo* ACTH secretagogue in the sheep (Smith *et al.* 1989, Pecins-Thompson & Keller-Wood 1994), though these findings have been recently called into dispute (McFarlane *et al.* 1995). A number of previous studies have examined the *in vitro* ACTH responses of anterior pituitaries from rats of the two sexes and various gonadal statuses to a range of ACTH secretagogues, and these have also been unable to elucidate any differences (Hary *et al.* 1993, Handa *et al.* 1994).

At the level of the adrenal gland, the most striking difference observed was that cortisol production in

response to both ACTH and 8BrcAMP was greater from cultures of female animals, with GDX appearing to have little effect. This observation is consistent with an extensive and long-standing *in vivo* literature which demonstrates that the female adrenal cortex produces more glucocorticoids than does the male, an effect which has been attributed to the stimulation of the adrenal gland by oestradiol (Kitay 1961, Coyne & Kitay 1969, for review see Handa *et al.* 1994). In contrast, it does not appear in the sheep that oestradiol is the single regulator of adrenocortical responsiveness, as the effect of long-term GDX on either ACTH- or 8BrcAMP-induced cortisol production was the same in both sexes. Somewhat surprising, however, was our observation that basal cortisol production was greatest by cells derived from intact male sheep. This finding differs from a number of previous studies (Vinson *et al.* 1978, Lesniewska *et al.* 1990, Wiesenfeld & Michaelis 1994), where glucocorticoid production from female animals appears to be greater under basal conditions. In two studies examining basal cortisol production by sheep fetuses, one found increased production by females (Ayromloo & Essman 1978), while the other suggested that males produced more cortisol (Schwartz & Rose 1998). The *in vitro* production of cortisol by adrenocortical cells from female sheep was greater by cells from late-pregnant ewes than by cells from non-pregnant ewes (Glickman *et al.* 1980), and it is possible that the relatively oestrogenic environment of late pregnancy underlies this difference.

Our observation that both ACTH and 8BrcAMP induce greater cortisol production in cultures from female animals suggests that the differences in adrenocortical cells occurs distal to the production of cAMP, which is considered the principal intercellular mediator of ACTH action at the adrenal (Simpson & Waterman 1988). Consistent with this notion is our observation that stimulated cAMP production was not affected by sex or GDX. The expression or activity of a number of steroidogenic enzymes in the adrenal has been reported to be increased in females, compared with males, or stimulated by oestrogens and inhibited by androgens (Belanger *et al.* 1991, Gallant *et al.* 1991, Perry & Stalvey 1992, Issacson *et al.* 1993, El-Migdadi *et al.* 1995, Nowak *et al.* 1995). The observation of increased steroidogenic potential in the female may, however, be somewhat secretagogue-specific, as we have recently shown that AII-induced cortisol production is greater in adrenocortical cultures from intact male sheep, than it is in GDX male or intact female sheep (Canny *et al.* 1998).

In conclusion, the results of this study have demonstrated that significant differences in the HPA axes exist between male and female sheep, and that some of these differences appear to directly depend on gonadal factors. Importantly, isolated adrenocortical cells of female sheep exhibit enhanced steroidogenesis in response to ACTH, and this increase appears to occur at a site following the

production of cAMP. This finding confirms, *in vitro*, the response of many species *in vivo*, and suggests a specific cellular mechanism of action. These observations therefore provide a model system to further investigate the differences between the HPA axes of male and female animals.

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