

Ultradian, circadian and seasonal rhythms in cortisol secretion and adrenal responsiveness to ACTH and yarding in unrestrained red deer (*Cervus elaphus*) stags

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Abstract

Seasonal changes in the activity and responsiveness of the adrenal gland in red deer (*Cervus elaphus*) stags were quantified by measuring 24 h endogenous cortisol secretory profiles and plasma cortisol responses to either administration of exogenous ACTH or a standardised stressor during November (period of velvet growth), February (pre-rut), April (mid-rut) and July (post-rut) (southern hemisphere) using a remote blood sampling device (DracPac).

Ultradian rhythms in the concentration of plasma cortisol were observed resulting from the episodic secretion of cortisol from the adrenal cortex at a mean rate of 0.8 pulses/h. Circadian rhythms in plasma cortisol concentrations were also found in 11 out of the 20 complete 24 h profiles (mean amplitude, 3.8 ± 1.4 ng/ml).

Seasonal rhythms in mean 24 h plasma cortisol concentrations and cortisol pulse parameters were also observed. Mean 24 h plasma cortisol concentrations were higher in November (12.5 ± 1.0 ng/ml) than in February (6.3 ± 1.0 ng/ml), April (4.0 ± 1.0 ng/ml) or July (4.2 ± 1.0 ng/ml). Cortisol pulse height, nadir and

amplitude were all significantly higher in November than at other times of the year ($P < 0.01$).

Peak cortisol concentrations following infusion of ACTH₁₋₂₄ (0.04 IU kg⁻¹) were higher ($P < 0.05$) in November (55.8 ± 2.7 ng/ml) and lower ($P < 0.001$) in April (33.7 ± 1.8 ng/ml) than those in February and July (48.7 ± 2.0 ng/ml and 45.4 ± 2.0 ng/ml respectively). The area under the cortisol response curve was significantly smaller ($P < 0.05$) in April (266.6 ± 15.3 ng/ml/190 min) than at other times of the year (February, 366.1 ± 15.3 ng/ml/190 min; July, 340.7 ± 15.3 ng/ml/190 min and November, 387.8 ± 21.2 ng/ml/190 min).

These data demonstrate that the adrenal gland of the red deer stag exhibits ultradian, circadian and seasonal rhythms in activity, and that its responsiveness to ACTH varies with season. November, a period of reproductive quiescence in the southern hemisphere, with new antler growth and rapid weight gain, is associated with higher mean plasma cortisol concentrations and a greater responsiveness to exogenous ACTH. In contrast, the breeding season is associated with lower adrenal activity and responsiveness.

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Introduction

Changes in the activity and function of the hypothalamic–pituitary–adrenal (HPA) axis are routinely used to assess welfare in farm animals (Barnett & Hemsworth 1990, Fraser & Broom 1990). Studies in a variety of species including human (Veldhuis *et al.* 1990), cattle (Ladewig & Smidt 1989), sheep (Kennaway *et al.* 1981) and deer (Monfort *et al.* 1993) have shown that the HPA axis exhibits variations over time in basal activity and responsiveness, distinct from that following the imposition of a stressor. This variation can form a consistent rhythm which fluctuates within or about 24 h (ultradian and circadian rhythms respectively) or can be of longer

duration, with variations occurring on an annual basis (seasonal or circannual rhythms).

Ultradian rhythms in plasma concentrations of glucocorticoids are the result of episodic secretion of glucocorticoid pulses from the adrenal cortex. Such pulsatile secretion is thought to prevent down-regulation of the HPA axis while maintaining its ability to respond to stress (Monfort *et al.* 1993). In deer, episodic secretion of cortisol, the predominant glucocorticoid (van Mourik *et al.* 1985, Smith & Bubenik 1990), has been investigated in Eld's deer (*Cervus eldi thamin*) (Monfort *et al.* 1993) but there appears to be no information on cortisol secretory dynamics in red deer (*Cervus elaphus*).

Circadian rhythms in plasma glucocorticoid concentrations are well characterised for humans (Weitzman *et al.* 1971) and rodents (Keller-Wood & Dallman 1984), with peak concentrations preceding the activity phase of the daily cycle. This rhythm is entrained by feeding (Saito *et al.* 1989) and activity/sleep wake cycles (Born *et al.* 1997). There is no evidence from studies of deer for a circadian rhythm in HPA axis activity (white-tailed deer (*Odocoileus virginianus*) (Bubenik *et al.* 1983); rusa deer (*Cervus timorensis*) (van Mourik & Stelmasiak 1984); Eld's deer (Monfort *et al.* 1993)). However, in these studies the deer were subjected to either handling or indoor housing, factors known to attenuate circadian rhythms in other species (Irvine & Alexander 1994).

Seasonal changes in activity and responsiveness of the HPA axis have been reported in a number of species including humans (Walker *et al.* 1997), primates (Schiml *et al.* 1996), rodents (Boswell *et al.* 1994) and fish (McLeese *et al.* 1994). Evidence for a seasonal rhythm in adrenal activity or responsiveness in deer varies between species and studies, with seasonal changes reported in white-tailed deer (Bubenik *et al.* 1983), reindeer (*Rangifer tarandus*) (Nilssen *et al.* 1985), axis deer (*Axis axis*) (Chapple *et al.* 1991) and red deer (Suttie *et al.* 1995, Cassidy 1996) but not in Eld's deer (Monfort *et al.* 1993) or in another study on axis deer (Bubenik & Brown 1989).

Red deer stags undergo dramatic seasonal changes in physiology and behaviour associated with the breeding season (rut). These include changes in secretion of hormones involved in the reproductive (Lincoln & Kay 1979, Suttie *et al.* 1992) and growth and metabolic (Suttie *et al.* 1989) axes. Changes in behaviour at this time include increased aggression (Suttie 1985) and a reduced voluntary food intake (Loudon *et al.* 1989) resulting in pronounced weight loss (Kay 1979). It is well documented in other mammalian species that HPA axis activity can be modulated by changes in reproductive function (increased activity and responsiveness in castrates compared with entire males (Bass *et al.* 1982, Verkerk & Macmillan 1997)) and changing metabolic and growth demands (Yanovski *et al.* 1997) as well as by social factors (Lyons *et al.* 1988). This has yet to be demonstrated for red deer.

Changes in responsiveness of the HPA axis as assessed by the adrenal response to physiological (e.g. adrenocorticotrophin (ACTH) infusion, insulin induced hypoglycaemia) or psychological (e.g. handling, open field test) challenges are commonly used in assessing the normal functioning of the HPA axis in animal stress research. Seasonal changes in adrenal responsiveness to ACTH challenge have recently been demonstrated in red deer stags (Suttie *et al.* 1995, Cassidy 1996) but there have been no studies of seasonal variation in response to stress or basal activity of the HPA axis.

The aim of the current study was to identify and quantify ultradian, circadian and seasonal rhythms in basal cortisol secretion of red deer stags. In addition, we sought

to measure seasonal changes in adrenal and HPA axis responsiveness following challenges with ACTH and a standardised management stressor.

Routine handling of red deer (e.g. yarding, drafting and restraint) has been shown to result in activation of the HPA axis (Ingram *et al.* 1994, 1997, Carragher *et al.* 1997). To avoid the confounding effects of handling stress on basal HPA axis activity, a remote blood sampling device (Ingram *et al.* 1994) was used to obtain blood samples from undisturbed animals at pasture.

Materials and Methods

Animals

Twelve 3-year-old red deer stags were maintained at the Ruakura Agricultural Centre, Hamilton, New Zealand (37°46' S, 175°20' E). Two groups of 6 animals, balanced for weight, were formed in July 1995. Each group was kept in a separate 0.25 hectare paddock during the experiment and in 0.25–0.5 hectare paddocks between experiments, with free access to pasture (ryegrass-white clover) and water.

Procedure and remote blood sampling

On four separate occasions throughout the year (1/11/95, 7/2/96, 10/4/96 and 10/7/96), blood samples were collected from the same three animals in each group over a 24 h period (24 h profile) and following two types of challenge (ACTH and yarding). Animals from two separate groups were used so that data were obtained from two replicates, thereby reducing the chances that features unique to one group would influence the results obtained. The experimental procedure was similar for each season, as detailed below.

On day 1 all animals were weighed and the animals to be blood sampled were sedated with an i.m. injection of 1 mg kg⁻¹ xylazine hydrochloride (Rompun, Bayer, Auckland, NZ) and a modified double lumen catheter (Cavafix Duo16/18G 32 cm, B. Braun, Melsungen, Germany) (see Fig. 1) was inserted into the jugular vein. At the same time all cannulated animals were fitted with a canvas and leather backpack. Sedation was reversed with an i.v. injection of 0.25 mg kg⁻¹ yohimbine hydrochloride (Revercyl, Aspiring Animal Services Ltd, Wanaka, NZ) and the animals were then reunited with the remaining three animals in their group and returned to their respective paddocks.

The following day (day 2) the 24 h profile blood samples were collected. Both groups were penned between 0600 and 1030 h and each of the six experimental animals was fitted with a battery powered remote blood sampling device (DracPac, Engineering Development Group, HortResearch and ABWRC, AgResearch, Ruakura Agricultural Centre, Hamilton, NZ).

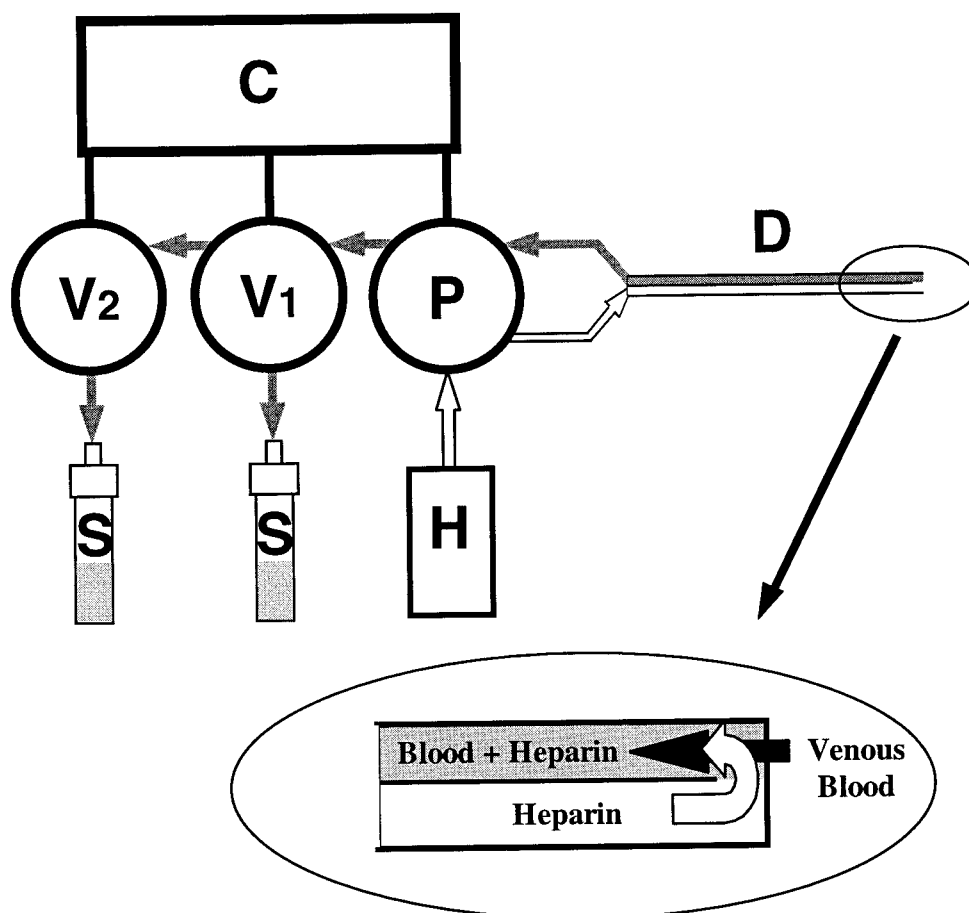


Figure 1 Schematic representation of the DracPac remote blood sampling system used for automatic continuous sampling of blood from free ranging animals. The device comprises: a microprocessor control box (C), a peristaltic pump unit (P), a plastic bag containing concentrated heparin (H), two 38-port rotary switching valves (V_{1-2}), 74 separate blood collection tubes (S) (4.5 ml monovette syringe) and a double lumen catheter (D) (Cavafix Duo16/18G 32 cm) which is modified by shortening the end of the catheter by ~5 cm and removing 1–2 mm of the wall between the two lumina at the tip to facilitate optimal mixing of the outflowing heparin and inflowing blood (see Ladewig & Stribny 1988).

The DracPac device (Fig. 1) consists of a pump unit containing two small peristaltic pumps for pumping heparin and drawing blood (length 150 mm, diameter 50 mm, weight 395 g), two 38-position rotary switching valves (length 200 mm, diameter 60 mm, weight 950 g), a 6 volt battery supply (Duracel, MN908) and a control box (length 160 mm, width 80 mm, height 60 mm, weight 580 g) containing a programmable microprocessor which allows programming of the start time, duration and rate of sample collection. The heparin pump delivers heparinised saline (5000 IU/ml) down one side of the double lumen catheter to the tip where it mixes with jugular blood being continuously drawn up the second lumen by the action of the blood pump. Heparin was delivered at a rate of 1.1 ml/h on days 2–3 and 2.6 ml/h on day 4. The heparinised blood passed through one or both rotary

switching valves and was collected into 1 of 74 separate blood collection tubes (4.5 ml monovette syringe, Sarstedt Ltd, Numbrecht, Germany). The samples were stored on ice pads (Dry Chill, Warragul, Vic., Australia) in an insulated pouch within the backpack. Cortisol concentrations remain stable in heparinised whole blood samples collected from red deer and stored in this fashion for at least 24 h (J R Ingram, unpublished data). Cortisol concentrations are also stable in heparinised whole blood of cattle stored at room temperature for up to 72 h (Reimers *et al.* 1983).

The DracPac was programmed to pump blood to waste until 1400 h (3.5 h after release from the yarding facility) to allow cortisol concentrations to return to baseline after handling (Ingram *et al.* 1994). The device was programmed to collect blood continuously at a rate of

15 ml per hour for 74 samples, each of 20 min duration. Following collection of the last blood sample (1440 h day 3), the animals were returned to the yards and the DracPac and blood samples were removed from the backpack.

On day 4, blood samples were collected before, during and after two types of challenge (ACTH₁₋₂₄ infusion and a yarding stressor). The 6 experimental animals were fitted with the remote blood sampling devices as on day 2, and returned to their respective paddocks. The DracPacs were programmed to pump blood to waste until 1230 h (3.5 h after release from the yarding facility), then to collect two 10 min baseline samples (30 ml/h). Immediately following the second baseline sample the DracPac was programmed to infuse 2 ml physiological saline containing 0.04 IU ACTH₁₋₂₄/kg live weight (Synacthen; Ciba NZ, Auckland, NZ) over 5 min into the jugular vein. Fresh blood was then drawn from the animal through the blood lines to a waste tube for 5 min to clear the blood lines of saline. Following the ACTH infusion and flushing of blood lines (total time 10 min) eighteen 10 min blood samples (30 ml/h) were collected over 180 min. Three hours after the ACTH challenge (1600 h) the animals were subjected to the yarding procedure. This comprised bringing the animals in their respective groups of six from their experimental paddocks into the yarding facility and keeping them in an indoor pen (2.4 × 2.4 m) for 10 min before returning them to their paddock (total time from leaving the paddock, yarding and return to the paddock was 30 min). Eight 10 min blood samples (30 ml/h) were collected during yarding and for 50 min subsequently at pasture. Following collection of the last blood sample, the experimental animals were returned to the yards and the catheter, blood samples and all equipment removed. A prophylactic dose of antibiotic (Tripen LA, Ethical Agents Ltd, Auckland, NZ) was administered i.m. after removal of the catheter. The experimental protocol was approved by the Ruakura Animal Ethics Committee.

Plasma cortisol analyses

All blood samples were centrifuged (1200 g for 15 min) after removal from the backpacks and the plasma stored at -20 °C until assayed for cortisol. Plasma cortisol concentrations were determined in duplicate, after extraction in ethyl acetate, by an ¹²⁵I radioimmunoassay method with polyethylene glycol separation (Ingram *et al.* 1994). A standard curve, using charcoal stripped deer plasma, was constructed and used to determine the concentration of cortisol in individual plasma samples. The cross-reactivity of the antiserum with 11-deoxycortisol was 5.7% and with cortisone was 1.2%. The mean inter- and intra-assay coefficients of variation for spiked deer plasma controls of known low, medium and high cortisol concentrations were 13.1 and 13.8% respectively. Assay sensitivity was 0.9 ng/ml.

Statistical analysis

Pulse detection Discrete pulses in cortisol secretion were quantified using cluster analysis (Veldhuis & Johnson 1986), a statistically based peak detection algorithm. A cortisol pulse was defined as a statistically significant increase in a cluster of cortisol concentrations which preceded a significant decrease. A cluster size of one point for the nadir and one point for a peak was used along with a constant 14% coefficient of variation and a pooled *t* statistic of 1 to limit the false peak detection rate to approximately 10% (Veldhuis & Johnson 1986). The pulse parameters measured for each 24 h profile were pulse frequency, pulse height, pulse amplitude, nadir and peak width.

Circadian rhythm detection Circadian rhythms in plasma cortisol concentrations were detected using a single cosinor model developed for Microsoft Excel (Microsoft Corp., USA) (Bourdon *et al.* 1995). The amplitudes and times of peak concentrations (acrophase) were obtained using this model in which the data were represented by the best fitting cosine function using the least squares calculation.

Statistical analyses Plasma cortisol concentrations are represented as means ± s.e.m. Statistical comparisons were performed using analysis of variance (MINITAB for Windows, Release 11.21, Minitab Inc., PA, USA). A blocking structure was used to separate the between- and within-animal variations. When significant (*P* < 0.05) trends were present, between-month comparisons were made using Student's *t*-test.

For the ACTH and yarding cortisol responses, the baseline values were calculated as the average of the 2 samples collected before the imposition of the treatment. Peak heights were defined as the maximum cortisol concentration observed after treatment initiation. Blood sample collection using the DracPac device was by continuous withdrawal over the 10 min sampling period. Therefore, cortisol concentrations for each sample represented an integrated value for that sampling period allowing the area under the cortisol response curve to be calculated by summing the measured concentrations observed after imposition of the treatments (190 min and 70 min for ACTH challenge and yarding stressor respectively).

Results

The percentage of samples collected successfully using the remote blood sampling device was 90% for the 24 h sampling (days 2–3) and 77% during the challenges. Loss of samples was mainly due to severing of blood collection lines between the catheter and the blood sampler.

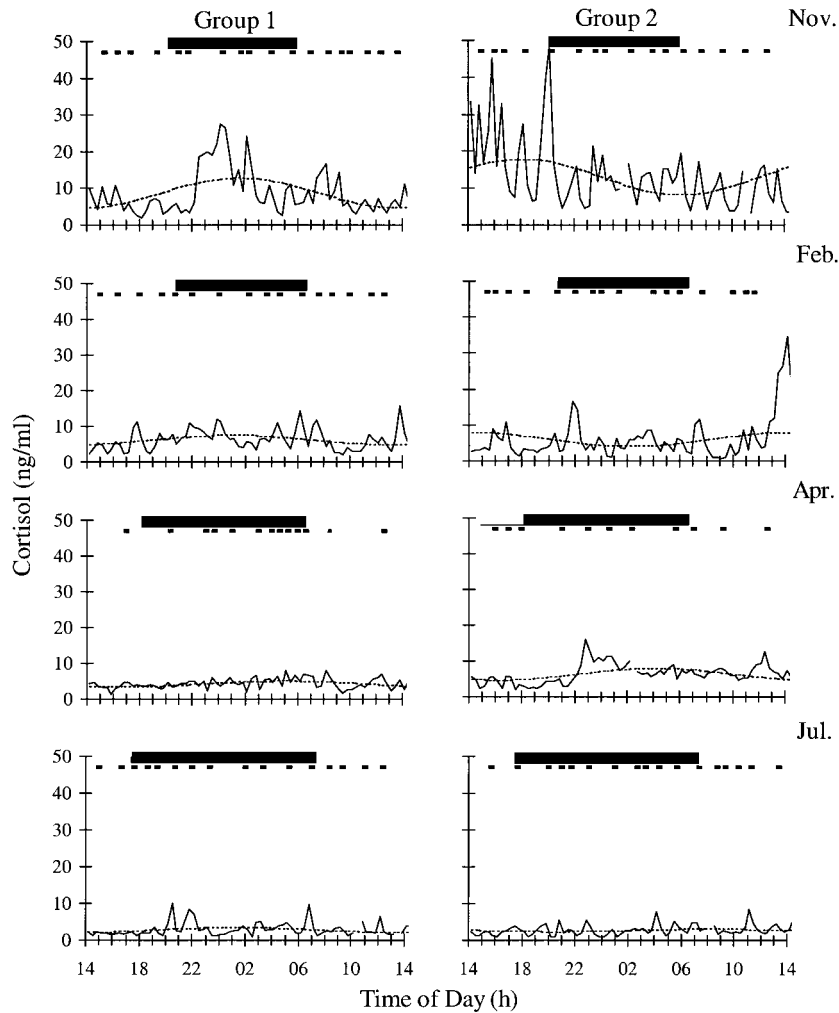


Figure 2 Seasonal effects on 24 h profiles of plasma cortisol collected from a representative stag from each group (1 and 2) at four different times of the year November (Nov.), February (Feb.), April (Apr.) and July (Jul.). The occurrences of cortisol pulses as detected by cluster analysis are shown as points at the top of each graph. Significant circadian rhythms are represented by the best fitting sine curve (dashed line) for each data series derived from COSINOR analysis. The black bars at the top of each graph represent the period of darkness for each month.

Animal live weights increased significantly ($P < 0.001$) from a mean of 116.7 ± 2.3 kg in November 1995 to a peak of 135.3 ± 2.3 kg in February 1996. Mean live weights tended to decline in April (131.8 ± 2.3 kg) and again in July (127.2 ± 2.3 kg), although these differences were not statistically significant ($P > 0.05$).

Twenty-four hour cortisol profiles

Representative profiles of cortisol concentration over 24 h for 2 stags (one from each replicate group) for each month are presented in Fig. 2. Cortisol profiles were characterised by a pulsatile secretory pattern and described

by the following parameters: pulse frequency, mean pulse height, mean nadir, mean pulse amplitude, and mean pulse width. The mean values for the six experimental animals for each of the four months are shown in Table 1. The frequency of cortisol pulses did not differ significantly between November, February and July, although pulse frequencies obtained in April were significantly lower ($P < 0.01$) than those in November and July. On average, cortisol pulses occurred once every 72 min, a frequency of 0.8 peaks/h. The height, nadir and amplitude of the cortisol pulses were significantly higher ($P < 0.001$) in November than at other times of the year. Cortisol pulse duration was significantly longer ($P < 0.05$) in April

Table 1 Seasonal effects on mean plasma cortisol concentrations (ng/ml) and cortisol pulse parameters (means \pm S.E.M.) derived from cluster analysis of 24 h cortisol profiles collected from six red deer stags at four different times of the year

	November	February	April	July
Cortisol (24 h mean)	12.5 \pm 1.0 ^a	6.3 \pm 1.0 ^b	4.0 \pm 1.0 ^b	4.2 \pm 1.0 ^b
Pulse				
Frequency (pulses/h)	0.89 \pm 0.03 ^a	0.82 \pm 0.03 ^{ab}	0.72 \pm 0.03 ^b	0.90 \pm 0.03 ^a
Height (ng/ml)	17.1 \pm 1.3 ^a	9.0 \pm 1.3 ^b	5.7 \pm 1.3 ^b	6.5 \pm 1.3 ^b
Nadir (ng/ml)	7.5 \pm 0.6 ^a	3.4 \pm 0.6 ^b	2.8 \pm 0.6 ^b	2.3 \pm 0.6 ^b
Amplitude (ng/ml)	9.7 \pm 0.8 ^a	5.3 \pm 0.8 ^b	2.9 \pm 0.8 ^b	4.1 \pm 0.8 ^b
Width (min)	61.1 \pm 2.6 ^a	62.6 \pm 2.6 ^{ab}	69.5 \pm 2.6 ^b	57.2 \pm 2.6 ^a

Values with different superscripts are significantly different ($P \leq 0.05$) between months.

compared with November and July. Pulses in February were of intermediate duration and not significantly different.

Individual mean 24 h plasma cortisol concentrations ranged from 1.9 to 22.5 ng/ml over seasons and were significantly higher ($P < 0.001$) in November than at other times of the year.

Significant ($P < 0.05$) circadian rhythms were found in 9 of the 20 cortisol profiles suitable for analysis (Table 2). Another two profiles in February approached significance ($P = 0.051$ and $P = 0.061$) and were included in the overall analysis. Parameters of the fitted sine curves are also given in Table 2. The amplitude of the circadian rhythm when present tended to be higher in November than in February, April and July, although this difference was not significant ($P = 0.146$, overall mean 3.8 ± 1.4 ng/ml). The acrophase (highest point) of the circadian rhythm in cortisol secretion occurred at significantly different ($P < 0.05$) times of the day in April and February, while the times of the acrophase in November and July were not significantly different ($P > 0.05$) from either April or February.

ACTH challenge

The mean cortisol concentrations following ACTH challenge for each of the four months are presented in

Fig. 3. The preinfusion cortisol concentrations were not significantly different ($P > 0.05$) between months (mean 4.6 ± 0.4 ng/ml). Peak cortisol concentrations following ACTH infusion were significantly higher ($P < 0.05$) in November (55.8 ± 2.7 ng/ml) and lower ($P < 0.001$) in April (33.7 ± 1.8 ng/ml) than those in February and July (48.7 ± 2.0 ng/ml and 45.4 ± 2.0 ng/ml respectively).

The area under the cortisol response curve was significantly smaller in April (266.6 ± 15.3 ng/ml/190 min) ($P < 0.05$) than at other times of the year (February, 366.1 ± 15.3 ng/ml/190 min; July, 340.7 ± 15.3 ng/ml/190 min and November, 387.8 ± 21.2 ng/ml/190 min).

Yarding stressor

The mean cortisol concentrations in response to the yarding stressor for each of the four months are presented in Fig. 4. The concentration of cortisol in baseline samples taken before the yarding stressor did not differ significantly between months (mean 4.2 ± 0.5 ng/ml). There were no significant seasonal differences in the peak concentrations of cortisol following yarding (November 23.1 ± 5.6 ng/ml; February 20.7 ± 4.9 ng/ml; April 10.9 ± 5.6 ng/ml and July 9.7 ± 4.6 ng/ml), or in the area under the cortisol response curve (November 81.5 ± 32.2 ng/ml/70 min; February 101.6 ± 27.9 ng/ml/70 min; April 47.9 ± 32.2 ng/ml/70 min and July 40.1 ± 26.6 ng/ml/70 min).

Table 2 Circadian effects on plasma cortisol concentrations derived from COSINOR analysis of 24 h cortisol profiles at four different times of the year (means \pm S.E.M.)

	November	February	April	July
No. of animals*	4/6	2 [†] /4	3/5	2/5
Amplitude (ng/ml) ⁺	8.4 \pm 2.5 ^a	1.5 \pm 2.5 ^a	1.1 \pm 2.5 ^a	2.1 \pm 2.5 ^a
Acrophase (h:min) [§]	00:07 \pm 2:29 ^{ab}	17:44 \pm 2:29 ^a	06:42 \pm 2:29 ^b	01:36 \pm 2:29 ^{ab}

*No. of animals indicates stags exhibiting a significant ($P < 0.05$) circadian rhythm in plasma cortisol concentrations out of those stags with a complete 24 h profile (i.e. ≥ 72 samples). ⁺Amplitude is the mean amplitude of the fitted sine curve expressed as ng/ml. [§]Acrophase is the mean time of day that the sine wave is maximal, expressed in hours and minutes. [†]In February the two animals had rhythms that approached significance ($P = 0.051$ and $P = 0.061$ respectively). Values with different superscripts are significantly different ($P \leq 0.05$) between months.

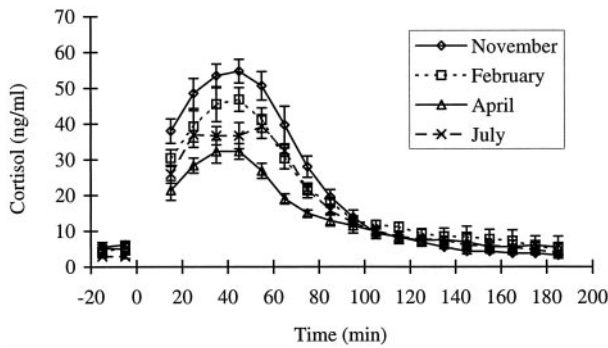


Figure 3 Seasonal effects on adrenocortical responses to ACTH₁₋₂₄ challenge (0.04 IU kg⁻¹) administered at time 0, to free-ranging red deer stags at pasture in November ($n=4$), February ($n=5$), April ($n=5$) and July ($n=5$). Values are shown as the monthly means \pm S.E.M.

Discussion

This study has shown that basal cortisol secretion in red deer stags is characterised by an episodic pattern of release from the adrenal gland, a feature common in other species (humans (Krieger *et al.* 1971); primates (Sarnyai *et al.* 1995); sheep (Fulkerson & Tang 1979)). Significant pulses in cortisol concentration occurred on average every 72 min, a frequency (0.8 peaks/h) similar to that reported for Eld's deer (0.6 peaks/h (Monfort *et al.* 1993)), and other ruminant species including cattle (0.5 peaks/h (Ladewig & Schmidt 1989, Lefcourt *et al.* 1993)), sheep (0.8–1.2 peaks/h (Fulkerson & Tang 1979)) and non ruminant species such as the horse (0.6 peaks/h (Irvine & Alexander 1994)). This pulsatile release of cortisol during periods of normal undisturbed activity may function to prevent down regulation of the adrenal axis, thereby maintaining the ability of the axis to respond maximally to stressors.

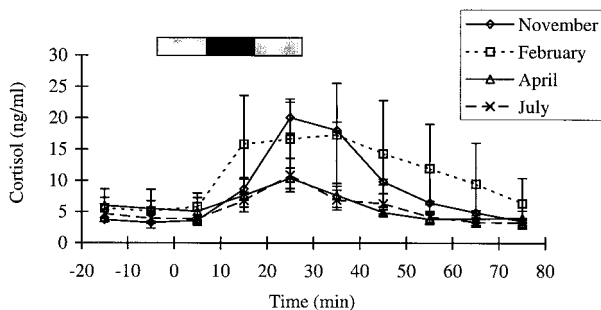


Figure 4 Seasonal effects on the adrenocortical response of free ranging red deer stags to a yarding stressor and subsequent recovery at pasture in November ($n=4$), February ($n=5$), April ($n=4$) and July ($n=5$). Grey bars indicate time spent moving to and from yards, the black bar indicates time spent in yards. Values are shown as the monthly means \pm S.E.M.

Individual mean 24 h concentrations of plasma cortisol found in the present study (1.9 to 22.5 ng/ml) are similar to basal cortisol concentrations reported for hand-reared Eld's deer (5.4–14.5 ng/ml (Monfort *et al.* 1993)) and in previous studies using the DracPac remote sampling device on undisturbed red deer stags (5.4–22 ng/ml (Ingram *et al.* 1994, 1997, Matthews *et al.* 1994, Carragher *et al.* 1997, Waas *et al.* 1997)) as well as in stag blood sampled immediately after being shot dead while undisturbed at pasture (5.7 ± 3.7 ng/ml (Smith & Dobson 1990)). These basal concentrations are generally lower than those reported for red deer in studies where animals have been physically or chemically restrained prior to blood sampling (6–27 ng/ml (Jopson *et al.* 1990); 18–27 ng/ml (Goddard *et al.* 1994); 17–55 ng/ml (Bubenik & Bartos 1993)). This indicates that in previous studies, restraint artefacts have been a potential confounding factor in the assessment of adrenal activity and function in red deer.

The presence of a circadian rhythm in plasma glucocorticoid concentrations is well established in humans (Weitzman *et al.* 1971) and rodents (Keller-Wood & Dallman 1984) and involves an anticipatory rise in cortisol prior to the activity phase, which is typically entrained by circadian rhythms in feeding (Saito *et al.* 1989) and resting (Born *et al.* 1997). In ruminants, contradictory reports exist about the presence of circadian rhythms in basal plasma cortisol concentrations. For example, in cattle several authors have reported a circadian rhythm in cortisol concentrations (Hays *et al.* 1975, Fulkerson *et al.* 1980, Thun *et al.* 1981, Lefcourt *et al.* 1993) whereas Hudson *et al.* (1975) failed to detect any circadian rhythm. Such rhythms in cortisol concentrations have not been identified in white-tailed deer (Bubenik *et al.* 1983), rusa deer (van Mourik & Stelmasiak 1984) or Eld's deer (Monfort *et al.* 1993), although the existence of a circadian rhythm in the latter study could not be ruled out as the animals were blood sampled for only 10 h per day. The deer in these studies were all subjected to either handling or indoor housing, factors known to attenuate circadian rhythms in horses (Irvine & Alexander 1994). In the present study, a significant circadian rhythm in plasma cortisol concentrations was found in half of the 24 h profiles suitable for analysis. Amplitudes were generally small, the exception being November, when a sustained period of agonistic behaviour in one group (data not shown) may have contributed to increased amplitudes.

Where reported, circadian rhythms are typically of low amplitude in ruminants (e.g. cattle, 1–1.4 ng/ml (Thun *et al.* 1981, Lefcourt *et al.* 1993)). At pasture, ruminants such as sheep, cattle and deer graze and rest in bouts throughout the day and night (Gates & Hudson 1983, Kilgour & Dalton 1984). Thus, they may not have the same cues available for entrainment of a circadian rhythm compared with non-ruminants or ruminants maintained indoors on fixed feeding schedules. While there is some

evidence from our study for circadian variability in cortisol concentrations in red deer at pasture, this rhythm is of low amplitude and potentially easily disrupted by stress associated with handling and blood sampling or social and environmental factors (Irvine & Alexander 1994). Further studies would be required to determine conclusively if circadian rhythms were a fundamental feature of HPA axis activity in red deer stags.

Seasonal rhythms in activity, behaviour, metabolism and secretion of hormones are well documented in cervid species (Lincoln *et al.* 1970, Loudon & Brinklow 1992, Suttie *et al.* 1992, Asher *et al.* 1996). In the present study significant seasonal rhythms were observed in the dynamics of 24 h cortisol secretion from the adrenal cortex of red deer stags. The frequency of cortisol pulses was significantly lower during the rut in red deer stags which contrasts with the lack of a seasonal pattern found in Eld's deer (Monfort *et al.* 1993) and may reflect fundamental differences between the two species of deer (temperate vs tropical habitat, autumn vs spring breeding). The role of androgens might play in modulating the pulsatile nature of HPA axis activity in red deer has yet to be investigated; however, androgens have been shown to suppress corticotrophin releasing hormone levels within the paraventricular nucleus (Bingaman *et al.* 1994) suggesting a central site of action (Handa *et al.* 1994).

November (southern hemisphere) is a unique time of year for red deer stags. It represents a degree of sexual quiescence uncommon in many other domestic ruminants. Secretion of luteinizing hormone (LH) from the pituitary and testosterone from the testes is negligible (i.e. pulse frequency of LH and testosterone is less than one pulse per 24 h and basal concentrations of LH and testosterone are below detectable concentrations (Suttie *et al.* 1992)). With no significant testicular activity, stags behave similarly to castrated animals (Lincoln *et al.* 1970). Major changes also occur in appetite, feed utilisation and fat accretion (Asher *et al.* 1994). The increase in cortisol pulse height, pulse nadir and pulse amplitude in November indicates that there is a shift in the regulation of the HPA axis during spring. Changes in height and amplitude reflect to some extent the increased responsiveness of the adrenal gland to ACTH seen at this time of year (Suttie *et al.* 1995, Cassidy 1996). However, the threefold increase in cortisol pulse parameter values compared with that seen during the rut would suggest that the adrenal gland changes are also accompanied by changes in endogenous ACTH secretion. Due to its rapid degradation in blood, ACTH could not be measured in remotely collected samples. However, male Soay sheep which display similar seasonal rhythms in basal cortisol concentrations and adrenal responsiveness to ACTH also show a marked seasonal variation in endogenous ACTH secretion with the peak concentrations occurring during the seasonal peak in cortisol concentrations (Ssewanyana *et al.* 1990). The increase in nadir also implies an up regulation of the central drive to the

HPA axis in conjunction with a down regulation of the glucocorticoid negative feedback system controlling the axis.

The increase in cortisol pulse parameters in November translated into greater mean 24 h cortisol concentrations which were significantly higher than those at other times of the year. Seasonal changes in cortisol secretion have been reported in males of some other seasonally breeding deer species. Cortisol concentrations have been reported to decline during the rut in white-tailed deer (Bubenik *et al.* 1983), reindeer (Nilssen *et al.* 1985) and axis deer (Chapple *et al.* 1991), while other studies have reported increased concentrations during the rut for red deer (Feher *et al.* 1994), or no change in reindeer (Ringberg 1979) and Eld's deer (Monfort *et al.* 1993). The metabolic and reproductive implications of a seasonal rhythm in basal cortisol concentrations in red deer stags are discussed below.

Alternatively, the decline in basal cortisol concentrations over the course of the experiment may result from a process of habituation to the remote blood sampling procedure. However, subsequent experiments (J R Ingram, unpublished data) conducted in November and April using red deer stags naive to the blood sampling procedure on both occasions, have revealed similar seasonal differences in cortisol concentrations to those found in the present study.

Cortisol concentrations and the temporal pattern of release from the adrenal gland in response to ACTH challenge were similar to those previously reported for red deer stags using the DracPac technique (Ingram *et al.* 1997). The maximum cortisol concentrations were within the range reported for red deer following ACTH challenge (males: 26 to 100 ng/ml (Bubenik & Bartos 1993, Ingram *et al.* 1997); females: 40 to 90 ng/ml (Jopson *et al.* 1990, Goddard *et al.* 1994)) and following routine handling procedures in our facility (males: 25 to 70 ng/ml (Matthews & Cook 1991, Ingram *et al.* 1994, 1997, Matthews *et al.* 1994, Carragher *et al.* 1997)). In the present study, red deer stags displayed significant seasonal variation in response to ACTH challenge. Maximal responses were higher in November (spring) and declined to a minimum in April (mid rut, autumn). Recent studies of adrenal responsiveness in red deer stags (Suttie *et al.* 1995, Cassidy 1996) have shown a similar seasonal rhythm with greater plasma cortisol responses to ACTH challenge during spring/summer compared with the breeding season and winter months.

Significant elevations in plasma cortisol concentrations were seen in response to a standardised acute stress (yarding) at all times of the year. The amplitude of the cortisol response to yarding in November and February was twice that seen in April and July, although this seasonal variation was not statistically significant due to the large variability seen in individual responses. Peak cortisol concentrations in November and February, however,

were similar to concentrations reported in an earlier study on stags, carried out in December (23 ng/ml (Carragher *et al.* 1997)), using the DracPac blood sampling technique and a similar yarding challenge. The generally lower cortisol responses in April and July would suggest either a reduced capacity of the HPA axis to respond to stress as supported by the ACTH challenge data, or a reduction in the perceived stressfulness of the yarding procedure possibly mediated via either habituation to the challenge or the attenuating effects of testosterone on fearfulness (Boissy & Bouissou 1994).

Compared with other routine handling procedures that have been evaluated using the DracPac technique, such as restraint (Ingram *et al.* 1994, Carragher *et al.* 1997), removal of growing antler (Matthews *et al.* 1994) and transport (Waas *et al.* 1997), the increase in cortisol secretion following yarding was of lesser amplitude and of short duration indicating that the procedure was potentially less stressful.

It is evident from this study that red deer stags have a strong seasonal rhythm in HPA axis activity and responsiveness. The mechanism by which this change occurs has yet to be determined in red deer, although these changes closely follow seasonal rhythms in testicular function (Suttie *et al.* 1992) and growth rates (Kay 1979). Testosterone concentrations which peak in stags during the rut (Suttie *et al.* 1992), can inhibit cortisol secretion directly by influencing steroidogenic pathways involved in the synthesis of cortisol in the adrenal cortex (Hornsby 1982, Miller 1988). It is possible that the hypertrophy and hyperplasia of the adrenal cortex seen in rutting wild red deer stags (Kapp 1989) and white-tailed deer stags (Hoffman & Robinson 1966) is in response to the reduced ability of the adrenal to produce cortisol at this time.

In addition, androgens can influence HPA axis activity by competing with cortisol for binding sites on carrier proteins such as corticosteroid binding globulin (CBG) (Bradley & Stoddart 1992) and plasma albumin (Ward *et al.* 1992). Androgens can also suppress hepatic synthesis of CBG (Gala & Westphal 1965) effectively increasing unbound concentrations of cortisol. This, in turn, would increase negative feedback to the HPA axis, reducing HPA axis activity and trophic drive to the adrenal cortex. A further mechanism by which androgens can influence HPA axis activity is by binding to androgen receptors in regions of the central nervous system such as the hippocampus known to modulate the HPA axis (Handa *et al.* 1994).

The seasonal rhythm in plasma cortisol concentrations found in the present study also correlates with the seasonal cycle in growth in the red deer stag, with higher concentrations of cortisol during the period of weight gain in November and reduced concentrations during periods of weight loss during the rut. Interestingly, humans suffering from Cushing's syndrome (chronic hypercortisolaemia)

show clinical signs of obesity whereas adrenocortical deficiency (chronic hypocortisolaemia) in humans is characterised by weight loss (Hauner *et al.* 1987).

The dramatic increase in weight during the spring and early summer and the higher levels of plasma cortisol coincide with elevated insulin concentrations and the development of insulin resistance in red deer stags (McMahon *et al.* 1997). It is well established that glucocorticoids stimulate feeding behaviour and insulin secretion (Dallman *et al.* 1995), and that cortisol excess can cause insulin resistance and obesity in humans (Brandes 1977). Cortisol promotes differentiation of adipocyte precursors into adipocytes and stimulates lipogenesis in the presence of insulin (Grégoire *et al.* 1991), and may have a role to play in the increased appetite, feed utilisation and fat accretion at this time of year in stags.

The period of the rut, with its increased aggression (Suttie 1985) and pronounced weight loss (Kay 1979), could be considered a period of increased social and nutritional stress. Chronic social stress has been reported to reduce plasma cortisol concentrations in female red deer (Goddard *et al.* 1994), yearlings (Hanlon *et al.* 1995) and stags (J R Ingram, unpublished data), while prolonged periods of weight loss in humans (over 26 weeks) resulted in a significant decrease in concentrations of cortisol and cortisol-binding globulin but no change in the concentration of free cortisol (Yanovski *et al.* 1997). In addition, nutritional stress is often associated with a metabolic disorder termed fatty liver syndrome. Red deer stags during the rut in the wild have a high incidence (80%) of fatty liver syndrome (Kapp *et al.* 1989) which may impair liver function reducing the synthesis of CBG (Veldman & Meinders 1996) and can directly inhibit steroidogenesis by limiting the availability of cholesteryl esters used in the synthesis of steroid hormones (Morrow *et al.* 1979, Nakagawa *et al.* 1997).

Glucocorticoid excess stimulates bone resorption and inhibits bone formation resulting in accelerated bone loss (Kleerekoper *et al.* 1997). Deer undergo skeletal bone loss during antler growth in spring (Hillman *et al.* 1973). The elevated concentrations of cortisol during the period of new antler growth in the present study and during antler mineralisation in late summer (Suttie *et al.* 1995) suggest a role for this steroid in red deer new antler growth.

The inhibitory influence of the HPA axis upon the reproductive axis (reviewed in Rivest & Rivier 1995) and immune function (reviewed in Munck *et al.* 1984) is well established in many animals. Therefore, it may be advantageous for stags during the breeding season, when inter-male competition for females is intense, to have reduced activity and responsiveness of the HPA axis as a strategy to maintain reproductive and immune competence in the face of increased physical and psychological stress.

In summary, the current data demonstrate that there is episodic activity of the HPA axis. There are also strong

seasonal effects on HPA axis activity and responsiveness to ACTH challenge in red deer stags. November, a period of reproductive quiescence, new antler growth and rapid weight gain, is associated with higher mean plasma cortisol concentrations and a greater responsiveness to ACTH. In contrast, the breeding season is associated with lower adrenal activity and responsiveness. The mechanisms involved have yet to be elucidated for red deer; however the inhibitory actions of androgens, variations in metabolic demand and exogenous factors such as chronic social/nutritional stress may be important factors. This study also highlights the advantages of remote blood sampling technology in obtaining undisturbed measurements of HPA axis activity and functioning without the confounding effects of repeated handling and restraint stress, information which is essential before reliable physiological measures of stress can be determined.

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