Effect of nutrition on seasonal patterns of LH, FSH and testosterone concentration, testicular mass, sebaceous gland volume and odour in Australian cashmere goats

S. W. Walkden-Brown1*, B. J. Restall2, B. W. Norton1, R. J. Scaramuzzi3† and G. B. Martin4,5

1Department of Agriculture, University of Queensland, QLD 4072, Australia; 2Wollongbar Agricultural Institute, NSW Agriculture, Wollongbar, NSW 2477, Australia; 3CSIRO Division of Animal Production, Prospect, NSW 2148, Australia; 4Animal Science Group, Faculty of Agriculture, University of Western Australia, Nedlands, WA 6009, Australia; and 5CSIRO Division of Animal Production, Wembley, WA 6014, Australia

The effects of season and diet on LH, FSH and testosterone concentrations, testicular mass, sebaceous gland volume and male odour were examined in mature Australian cashmere goat bucks fed ad libitum with diets of low or high quality for 16 months under natural photoperiod at 29°S, 153°E (n = 6 per treatment). Each week plasma was sampled, the bucks were weighed, scored for male odour and assessed for testicular mass based on scrotal circumference. Each month a skin sample was taken from the occipital region for histological assessment of sebaceous gland volume. For each variable there was a clear circannual cycle that was significantly influenced by dietary treatment. In bucks fed the low-quality diet, the timing of seasonal changes in LH and testosterone concentration, sebaceous gland volume and odour score was similar, with a mid-autumn peak. In each case the high-quality diet advanced, extended the duration and increased the magnitude of the seasonal increase. FSH concentrations peaked in late spring (in bucks on the high-quality diet) or summer (in bucks on the low-quality diet), reaching a nadir in early winter. The high-quality diet significantly increased concentrations only in the last 2 months of the experiment (spring). There was no overall association between these variables and change in testicular mass; instead, it was strongly correlated with voluntary feed intake and change in body mass, themselves subject to seasonal variation with a winter or spring peak. The high-quality diet induced large increases in body mass and testicular mass during the first months of the experiment without influencing the seasonally low concentrations of FSH, LH and testosterone present at the time. These results demonstrate that the male, like the female, Australian cashmere goat, exhibits marked reproductive seasonality, and that nutrition is a powerful modulator of the seasonal cycle. They suggest that testosterone concentration, sebaceous gland volume and odour score are ultimately dependent upon LH secretion, which appears to be under strong seasonal (photoperiodic) control, with the effects of enhanced nutrition limited to periods when photoperiodic inhibition is waning. However, seasonal regulation of testicular mass, and therefore sperm production, appears to be primarily dependent on changes in voluntary feed intake and growth, with the seasonal cycle of testicular mass more a consequence of the seasonal appetite or growth cycle than of changing gonadotrophin concentrations.

Introduction

For all species, a balance must be achieved between reproductive strategies based on either opportunism or seasonal cuing (Bronson, 1985). Opportunistic breeders will mate whenever environmental conditions are good, while seasonal breeders restrict mating to a time that will result in optimal conditions for the rearing of the young – the most physiologically demanding event of the reproductive cycle. Changing photoperiod is the most common environmental cue used to regulate reproduction in seasonal breeders (Clarke, 1981), but many species and strains do not fit neatly into a seasonal or opportunistic classification because, for them, the timing of...
reproductive events is determined by complex interactions between photoperiodic, social and nutritional cues. While most studies investigating these interactions to date have used small rodent species more suited to opportunistic strategies (e.g. Irby et al., 1984; Wayne and Rissman, 1990; Wayne et al., 1991), few have been carried out on larger, more strongly photoperiodic species.

The goat, like the sheep, is classified as seasonally polyoestrous and photoperiod is the primary environmental cue used to regulate reproduction (Bissonnette, 1941). This seasonality is strongly present in female Australian cashmere goats and the feral goats from which they are derived (Harrington, 1982; Restall, 1992). Nevertheless, it is increasingly evident that social and nutritional cues are also important regulators of the seasonal reproductive cycle in this species. For example, seasonally anovulatory does can be induced to ovulate in response to the sudden introduction of males (the so-called male effect), with the magnitude of the response influenced by both season (Chemineau, 1987; Restall, 1992) and the nutritional status of the male (Walkden-Brown et al., 1993). Bucks exposed to females in oestrus also exhibit an increase in gonadotrophin and testosterone secretion (the so-called female effect); the response depends upon season (Howland et al., 1985). In addition, drought-breaking rain and the associated increase in the availability of feed can initiate reproductive activity in Australian feral goats (Harrington, 1982), and prevailing nutritional conditions also appear to influence reproductive activity in less seasonal tropical goats (Gonzalez-Stagnaro, 1983).

On the basis of this knowledge, we hypothesized that mature bucks maintained under ambient photoperiod and fed diets of different quality would exhibit a circannual pattern of variation in reproductive activity, the timing and magnitude of which would be influenced by both the quality of diet and exposure to females in oestrus. These hypotheses were tested in an experiment using mature Australian cashmere goats. In this paper, we report the effects of nutrition on gonadotrophin concentrations, testicular function and some secondary sexual characteristics.

Materials and Methods

Location and animals

The experiment was carried out at the Wollongbar Agricultural Institute, Wollongbar (28°48' S, 153°25'E). Hours of daylight (sunrise to sunset) vary from 10.3 h at the winter solstice (June 21) to 14.0 h at the summer solstice (December 22), while mean minimum and maximum temperatures range from 8.9 and 17.6°C in July to 18.5 and 26.4°C in January. The experimental animals (n = 12) were drawn from a random-bred line of domesticated feral goats. They were 34 months old and had all previously been used for breeding. The history and management of the goat population at Wollongbar has been described by Restall and Pattie (1989).

Experimental procedure

Bucks were individually housed in 2.0 m x 2.5 m pens under natural photoperiod for 16 months between 1 July and 13 October of the following year. The experiment commenced when body mass was at its seasonal nadir for bucks at pasture in this environment. All bucks were offered an introductory diet of 1.2 kg pelleted lucerne (Medicago sativa) and coarsely hammermilled pangola grass (Digitaria decumbens) hay day⁻¹ in equal parts for 10 days, before they were divided into two treatment groups stratified on body mass and libido. One group (Low) was fed a low-quality diet of hammermilled pangola grass hay (crude protein: 6.9%; metabolizable energy: 6.6 MJ kg⁻¹) ad libitum, while the other group (High) was fed a high-quality diet of pelleted lucerne (crude protein: 17.6%; metabolizable energy: 8.3 MJ kg⁻¹) ad libitum for the remainder of the experiment. The diets and their digestibility, together with data on voluntary feed intake and growth, have been described by Walkden-Brown et al. (1994a).

Each week the bucks were sampled for plasma, weighed and the scrotal circumference was measured. Each month a skin biopsy was taken to assess sebaceous gland activity. Every second month blood samples were collected at intervals of 20 min over two 8 h periods, with each buck exposed to a female in oestrus throughout the second period. The acute effects of females in oestrus on pulsatile hormone secretion have been reported separately by Walkden-Brown et al. (1994b). In early October it was noted that the bucks in the High group began exhibiting a strong buck odour, while those in the Low group did not; therefore, in each week thereafter the odour was evaluated subjectively. This was done by smelling the dorsum of the neck 10–15 cm immediately posterior to the base of the horns, where the male odour was strongest, and allocating a score of 0 (neutral odour, no different from a female or a castrated male), 1 (mild male odour), 2 (moderate male odour) or 3 (strong male odour). These measurements were highly repeatable within operators and were carried out by a single operator for the entire experiment.

Weekly plasma samples for LH, FSH and testosterone assay comprised pooled aliquots from three jugular blood samples collected at intervals of 40 min. The scrotal circumference was measured to the nearest 0.1 cm in the standing animal using a light elastic band placed around the neck of the scrotum to seat the testes in the scrotum. The scrotal skin was clipped every 6 weeks. Skin samples were collected with a 1 cm trephine at random from a region 5 cm wide on the dorsal aspect of the neck, between 5 and 10 cm posterior to the base of the horns.

Radioimmunoassay

The LH and testosterone assays are described in detail by Walkden-Brown et al. (1994b). LH was assayed in duplicate using a heterologous double-antibody radioimmunoassay with caprine LH standards (Henniawati, 1993; immunopotency: 0.208 × NIH-oLH-S20) and rabbit antiserum directed against ovine LH. The mean (±SEM) assay sensitivity was 1.33 ± 0.14 µg l⁻¹. The mean intra-assay coefficients of variation were 12.3, 8.1 and 6.3% for samples containing 3.0, 12.0 and 36.0 µg l⁻¹, respectively, while the mean interassay coefficients of variation were 21.9, 7.8 and 6.6%, respectively. Testosterone was assayed in duplicate using a single antibody radioimmunoassay after extraction with hexane. The mean (±SEM) assay sensitivity was 0.08 ± 0.01 µg l⁻¹. The mean
intra-assay coefficients of variation for quality controls containing 0.38, 2.89 and 8.19 µg l\(^{-1}\) were 16.9, 9.4 and 8.1%, respectively, while the mean interassay coefficients were 12.7, 7.2 and 6.7%, respectively.

FSH samples were assayed in duplicate in a single assay, using an ovine radioimmunoassay described and validated for goats by Miller and Martin (1993). The standard used was NIAMDD-oFSH- RP-1, and the limit of assay sensitivity was 0.13 µg l\(^{-1}\). The mean intra-assay coefficients of variation for quality controls containing 1.45, 2.61 and 4.24 µg l\(^{-1}\) were 13.9, 8.6 and 9.0%, respectively.

**Determination of sebaceous gland volume**

Skin samples were fixed in buffered 10% formalin before being dehydrated, cleared and embedded with paraplast in an automated sample processor. Sections, 10 µm thick, were cut and stained with haematoxylin and eosin. The total volume of sebaceous gland in a single section, as a percentage of the total volume of the section, was determined by volumetric analysis using differential point counting as described by Weibel et al. (1966). Sebaceous gland tissue was readily identified on the basis of staining and morphological characteristics (Maddocks and Jackson, 1988). Counts were made on five different fields selected at random within each of five distinct zones on the circular section (lower, central, upper, right, left), and covered a minimum of 16.7% of the total area of the section. To estimate the total sebaceous gland volume in a skin sample, sections at 180 µm depth in the skin and at 200 µm intervals thereafter were examined as described above. The total volume of sebaceous gland in each sample was then determined using the following formula:

\[
V_{\text{tot}} = (\pi r^2) \times (D_1 - D_0) \times (\sum V_{1...n}/100s) \times (R^3/r^3),
\]

where \(V_{\text{tot}}\) is the total volume of sebaceous gland in sample (mm\(^3\)), corrected for shrinkage due to fixation, \(r\) is the mean radius of \(n\) sections (mm), \(D_1...n\) is the depth in mm, of sections 1...\(n\), \(V_{1...n}\) is the percentage of sebaceous gland tissue in equally spaced sections 1...\(n\), \(s\) is the number of counted sections containing sebaceous tissue, and \(R\) is the radius of sections prior to fixing (0.5 cm).

The value \((R^2/R^2)\) is a correction factor for shrinkage in cross-sectional area and depth for the sample.

The measurement of \(V_{\text{tot}}\) is time consuming, so an attempt was made to identify an accurate indirect measure of this variable in a preliminary study. \(V_{\text{tot}}\) was measured in 16 skin samples selected to provide maximum variation (Fig. 1). The samples were from four individual bucks (two from each diet group): a sample from October, January, April and July was taken from each buck. The relationship between total sebaceous gland volume in a sample and the relative volume of sebaceous gland at various depths in the sample was then examined. The percentage of sebaceous tissue at a skin depth of 780 µm \(V_{780}\) was the best fixed depth predictor of \(V_{\text{tot}}\), with a strong linear relationship between the two variables \(V_{\text{tot}} = -5.04 + 1.26 \times V_{780}; R^2 = 0.874; P < 0.001\). Consequently, all further samples were measured for sebaceous gland volume at a skin depth of 780 µm.

**Derived variables**

**Testicular mass.** Testicular mass was estimated from scrotal circumference measurements, using the equation:

\[
PTM = 21.5SC - 323.7,
\]

where \(PTM\) is the paired testicular mass in g, and \(SC\) is the scrotal circumference in cm \((R^2 = 0.88; n = 75; P < 0.001; Walkden-Brown et al., 1994c)\). The transformed variable was used because it is more informative, and enables direct comparisons of the magnitude of change in body mass and testicular mass to be made.
Daily change in testicular mass. The daily change in testicular mass was derived from weekly data smoothed using the Lowess smoothing algorithm to reduce error associated with measurement (Cleveland, 1981). The smoothed data provided an excellent fit of the data within bucks and removed the majority of the apparently random noise in the series, accounting for 86.1 ± 0.05% of the variation within bucks, and 97% of the total experimental variation in testicular mass. However, the unsmoothed data were used for the analysis of testicular mass.

Statistical analyses

Weekly data (n = 67) were grouped into monthly means (calendar months; n = 16) for each buck. These means, and records of any monthly measurements, were then subjected to repeated measures analysis of variance (Wilkinson, 1990). Log transformation of the hormone data to correct for heterogeneity of variance did not alter the conclusions of the analysis, so the analysis of the untransformed data is presented for ease of interpretation. Associations between variables over time were examined with standard linear correlation methods, using monthly treatment means (within treatments: n = 16; pooled: n = 32). In the text, means are presented with the standard error of the mean.

Results

Plasma concentrations of LH, FSH and testosterone

For all three hormones, there was a significant effect of month (P < 0.001) and a significant interaction between the effects of month and diet (P < 0.05; Fig. 2a,b,c). In the case of testosterone, the overall effect of diet was also significant (P < 0.05).

FSH concentrations varied seasonally in both treatments, reaching a peak value in late spring (High) or summer (Low) and reaching a nadir in late autumn or early winter (Fig. 2a). Diet only affected FSH concentration in the final 2 months of the experiment, with significantly higher concentrations in bucks in the High group (P < 0.05).

LH concentrations showed significant seasonal variation, rising in both treatments in late spring and falling during late autumn to a winter nadir (Fig. 2b). Diet affected the magnitude of the seasonal rise in bucks in the High group, with these bucks having significantly higher concentrations in November and May (P < 0.05). The concentrations of LH and FSH were correlated in bucks on the low-quality (r = 0.79, P < 0.001) but not the high-quality diet (r = 0.46, not significant).

Fig. 2. Monthly means (± SEM) of plasma concentrations of (a) FSH, (b) LH and (c) testosterone, (d) the percentage of sebaceous gland tissue at a skin depth of 780 μm in samples taken from the occipital region and (e) odour score for 3-year-old cashmere bucks fed diets of low (□; n = 6) or high (♦; n = 6) quality ad libitum under natural photoperiod at 29°S for 16 months. The dashed curve represents the annual curve of photoperiod (units not shown; range: 10.3–14.0 h).
Testosterone concentrations showed a marked seasonal profile in both dietary treatments, closely paralleling, but of greater amplitude than, that of LH (Fig. 2c). The effect of diet on the seasonal pattern was also more pronounced, with significantly higher concentrations occurring in bucks on the high-quality diet from November to February ($P < 0.05$). The concentration of testosterone was strongly correlated with LH concentration within both treatments (Low: $r = 0.82$; High: $r = 0.91$, $P < 0.001$) and overall ($r = 0.87$, $P < 0.001$), but was not correlated with FSH concentration (Low: $r = 0.33$; High: $r = 0.23$; not significant).

Sebaceous gland volume

The volume of sebaceous gland tissue was significantly influenced by diet and month of measurement, with significant interaction between the two effects ($P < 0.001$; Fig. 2d). There was marked seasonal variation in both treatments that was similar to that observed for LH and testosterone. Bucks on the high-quality diet had significantly higher sebaceous gland volumes than did those on the low-quality diet from November to February and from August to October in the second year ($P < 0.05$). Sebaceous gland volume was strongly and positively correlated with LH and testosterone concentrations within dietary treatments (Low: $r = 0.73$ and 0.84, respectively; $P < 0.001$; High: $r = 0.84$ and 0.83, respectively; $P < 0.001$) and overall ($r = 0.80$ and 0.84, respectively; $P < 0.001$), but was not correlated with FSH concentration in either treatment.

Odour score

Both diet and month of measurement influenced the odour score significantly, with significant interaction between these effects ($P < 0.001$; Fig. 2e). There was a clear seasonal pattern in odour score in both treatments, with scores peaking in autumn and reaching a nadir in late winter or early spring. Bucks fed the high-quality diet had significantly higher odour scores ($P < 0.05$) than did those fed the low-quality diet in all months except June and July of the second year. Odour score was positively correlated with testosterone concentration and sebaceous gland size within treatments (Low: $r = 0.65$ and 0.73, respectively; $P < 0.05$; High: $r = 0.88$ and 0.82, respectively; $n = 13$; $P < 0.001$) and overall ($r = 0.71$ and 0.79, respectively; $n = 26$; $P < 0.001$).

Body mass

There were significant effects of both diet and month on body mass and the change in body mass, with significant interaction between these effects ($P < 0.001$; Fig. 3a,c). Bucks in

![Fig. 3. Monthly means (± SEM) for (a) body mass, (b) paired testicular mass, (c) change in body mass, (d) change in testicular mass and (e) the ratio of testicular to body mass for 3-year-old cashmere bucks fed diets of low (○; $n = 6$) or high (*) quality ad libitum under natural photoperiod at 29°S for 16 months. The dashed curve represents the annual curve of photoperiod (units not shown; range: 10.3–14.0 h).](image)
both treatment groups exhibited a seasonal growth cycle characterized by weight loss during autumn, despite free access to the diets. The effect of diet was seasonally dependent, with the high-quality diet enhancing growth significantly only from July to October in both years.

**Testicular mass**

Testicular mass was influenced by both diet ($P < 0.05$) and month of measurement ($P < 0.001$), with significant interaction between these effects ($P < 0.001$; Fig. 3b). Seasonal change was evident in both treatment groups, with the pattern of change closely following that of body mass, peaking in summer and reaching a nadir in winter. Testicular mass was greater in bucks fed the high-quality diet for all but the first 2 months of the experiment and during April ($P < 0.05$).

Testicular mass was strongly correlated with body mass both within treatments (Low: $r = 0.74$; High: $r = 0.92$; $P < 0.001$) and overall ($r = 0.96$; $P < 0.001$; Fig. 4a). Correlations with gonadotrophin concentrations varied with treatment (LH: Low: $r = 0.80$, $P < 0.001$; High: $r = 0.52$, $P < 0.05$; FSH: Low: $r = 0.62$, $P < 0.05$; High: $r = 0.89$, $P < 0.001$) and the overall association was only moderate (LH: $r = 0.59$; FSH: $r = 0.62$; $P < 0.001$; Fig. 4b,c).

Change in testicular mass was significantly higher in bucks on the high-quality diet compared with those on the low-quality diet during the first 4 months of the experiment (July–October), but over the following 3 months (November–January) the reverse was true ($P < 0.05$; Fig. 3d). Beyond January, there were no significant differences between the treatments in any month.

Daily change in testicular mass was positively correlated with digestible organic matter intake and daily change in body mass within treatments (Low: $r = 0.86$, $P < 0.001$ and $r = 0.56$, $P < 0.05$, respectively; High: $r = 0.89$ and $0.99$, $P < 0.001$, respectively) and overall ($r = 0.85$ and $0.89$, $P < 0.001$ respectively; Fig. 5a,b). Overall, there was no association between change in testicular mass and the concentration of LH, FSH or testosterone, but significant associations were found within treatment groups (Fig. 5c,d). In bucks on the low-quality diet, change in testicular mass was positively correlated with FSH ($r = 0.50$, $P < 0.05$), but not LH or testosterone concentrations. In bucks fed the high-quality diet, it was negatively correlated with LH and testosterone concentrations ($r = -0.58$ and $-0.74$, respectively; $P < 0.05$) and there was no association with FSH.

Because paired testicular mass was so closely associated with body mass, the ratio of the two variables was examined to determine whether the association was modulated by season or diet. It was significantly influenced by month of measurement ($P < 0.001$), and there was a significant interaction between the effects of month and diet ($P < 0.001$) (Fig. 3e). There was a significant seasonal shift in the ratio in bucks on the low-quality diet with a peak in late summer and autumn, while in bucks fed the high-quality diet the ratio was stable over the first 4 months before declining continuously throughout the rest of the experiment. The ratio was not significantly correlated with any hormone in bucks on the high-quality diet, but was...
correlated with LH \( (r = 0.65; \ P < 0.05) \) and testosterone \( (r = 0.69; \ P < 0.01) \), but not with FSH \( (r = 0.43; \) not significant) concentrations in bucks on the low-quality diet.

**Discussion**

The effects of season and diet on hormone concentrations and secondary sexual characteristics in this study are best considered in relation to two fundamental seasonal cycles, both of them subject to modulation by prevailing nutritional conditions. The first of these cycles is a reproductive cycle driven by changing secretion of gonadotrophins (Walkden-Brown *et al.*, 1994b), while the second is an appetite or growth cycle driven by unknown factors (Walkden-Brown *et al.*, 1994a). Most of the variables examined appeared to be driven by secretion of pituitary gonadotrophins, in particular LH, forming a cascade of dependent effects through testosterone secretion, sebaceous gland size and odour score. However, testicular mass did not fit into this cascade and appeared to be influenced primarily by changes in the appetite or growth cycle with some modulation by gonadotrophins.

The seasonal profile of FSH, LH and testosterone concentrations observed in bucks on the constant low-quality diet is in broad agreement with that observed in other studies in goats (Muduili *et al.*, 1979; Howland *et al.*, 1985; Ritar, 1991), and seasonal breeds of sheep (Lincoln and Short, 1980; D’Occhio and Brooks, 1983; Lincoln *et al.*, 1990). In rams, such changes are driven by changing photoperiod and mediated primarily by changes in the frequency of pulses of GnRH released from the hypothalamus (Lincoln and Short, 1980). In goats, photoperiod also drives reproductive seasonality (Bissonnette, 1941; Delgadillo and Chemineau, 1992), and seasonal changes in LH pulse frequency suggest that the effects of photoperiod are at least partly driven by changing GnRH secretion (Howland *et al.*, 1985; Ritar, 1991; Walkden-Brown *et al.*, 1994b). That the seasonal testosterone cycle closely followed that of LH is not surprising given the dependence of testosterone on LH secretion. The greater amplitude of the testosterone cycle compared with that of LH is probably due to seasonal changes in testicular responsiveness to LH, with greatly enhanced responsiveness during the autumn rut (Walkden-Brown *et al.*, 1994b). The seasonal pattern of sebaceous gland volume closely matched that of testosterone since these glands are regulated by androgens (Ebling, 1957; Jenkinson *et al.*, 1967). During the period when the sebaceous gland was large it was noticeable that the fleece of the bucks, particularly in the neck region, became greasy with sebum. This sebum had an extremely strong, unpleasant buck odour and changes in its secretion were almost certainly responsible for the seasonal changes in odour in the bucks (Jenkinson *et al.*, 1967; Sasada *et al.*, 1983). The precise functions of this seasonal pattern of sebaceous gland size in the buck, and the associated changes in odour, remain obscure. There is evidence that the odour of buck fleece contributes to the ovulatory response of does to the introduction of bucks (Shelton, 1980; Claus *et al.*, 1990), and it has been postulated that combinations of odours from male urine and fleece may signal the metabolic status of one male to
others (Coblentz, 1986). It is also possible that, in common with many other ungulates, these odours play a role in the marking of territory (Müller-Schwarze, 1991) or in the recognition of individuals (Beauchamp et al., 1976).

The effects of diet on testosterone secretion, sebaceous gland size and odour score all appear to flow from the season-dependent effect of diet on LH secretion. For each of these variables, the effect of the high-quality diet was to advance, extend the duration of and increase the magnitude of the seasonal rise, with the greatest effect occurring during the summer months. Nutritional effects on the secretion of LH have been reported in many species, and most studies have demonstrated an inhibitory effect of undernutrition at the hypothalamus (l’Anson et al., 1991). Fewer studies have examined the effects of enhanced nutrition. In Merino rams, supplementation with lupin grain results in a rapid increase in the frequency of LH pulses; these pulses then dissipate in 3–4 weeks, despite continued testicular and body growth (Martin et al., 1994a, b). The effect of improved nutrition in the present experiment was delayed rather than acute, and the dependence upon season contrasts with the situation in Merino rams in which the response of LH to enhanced nutrition has been observed at all times of the year (GB Martin, unpublished). This difference in the role of season in determining the LH response to improved nutrition may reflect differences in the extent of photoperiod regulation of the gonadotrophic axis. Cashmere goats are more strongly seasonal than are Merino sheep and it is possible that the extent of photoperiodic inhibition of the gonadotrophic axis during winter and spring precludes responses to nutrition, while in the relatively nonseasonal Merino, photoperiodic inhibition is weak and able to be overridden by nutritional stimuli. The effect of diet on testosterone secretion was not related to the effect on testicular size, suggesting that diet-induced testicular growth does not alter Leydig cell function, as appears to be the case in the Merino, ram (Martin et al., 1987, 1994b). The effects of diet on odour score were of greater magnitude and duration than those on sebaceous gland size. This probably results from greater accumulation of sebum in the fleece of bucks fed the high-quality diet, given the longer period of sebaceous gland enlargement in response to this treatment, and the absence of weathering by sunlight and rainfall.

The effect of diet on FSH concentration was less pronounced than for LH, although there was evidence of an earlier peak in FSH concentration in bucks on the high-quality diet, consistent with the advancement of the seasonal rise in LH concentration. However, FSH concentration only differed significantly between diets in the later stages of the experiment at a time when LH concentrations were not affected, suggesting differential regulation of FSH and LH secretion by nutrition. It is not clear why significant differences in FSH concentration were restricted to this period. While there was no significant response to nutrition during the early months of the experiment, there was a trend towards higher FSH concentrations in the High treatment group, consistent with the observation that supplemental nutrition increases FSH concentrations in plasma of rams (Martin et al., 1994b).

Although the effects of both season and diet on testosterone concentration were closely associated with changes in LH concentration, the effects on testicular growth appeared to be less dependent on changes in gonadotrophin concentrations. The only consistently strong correlates with testicular growth, within and across treatments, were body growth and voluntary intake of food, indicating a dependence upon the metabolic status of the animal. The acute effect of nutrition on testicular mass during the first few months of the experiment was not related to changes in either LH or FSH concentration. Similarly, the long-term difference between nutritional treatments in testicular mass was maintained during periods when concentrations of these hormones did not differ. It would appear, then, that the effects of diet on testicular mass are mediated primarily by metabolic signals acting independently of changes in gonadotrophin concentrations. The close association between seasonal changes in body and testicular mass suggests that the seasonal testicular cycle could be similarly mediated, being a secondary response to the seasonal appetite or growth cycle. However, the seasonal cycle of testicular growth was also associated with FSH concentration in bucks fed the low-quality diet and, for the latter part of the experiment, in bucks fed the high-quality diet (the early part of the cycle being clearly disrupted by the acute testicular response to this diet). This finding is consistent with the important role that FSH plays in regulating spermatogenesis in rams (Courot and Ortantav, 1981; Kilgour et al., 1994) and indicates that FSH may be involved in regulating the seasonal testicular cycle in goat bucks, as appears to be the case in rams (Lincoln, 1989). Nevertheless, the clear response of the testis to metabolic changes, independently of changes in gonadotrophin concentrations, suggests that the main factor driving the seasonal testicular cycle is the seasonal appetite or growth cycle.

Testicular size is a good predictor of sperm production in most species (Amann, 1970), including goats (Walkden-Brown et al., in press), and increases in testicular mass induced by improved nutrition in rams are associated with even greater increases in sperm production (Oldham et al., 1978). If we assume that changes in testicular size during the experiment were associated with changes in the spermatogenic capability of the bucks, then there was clear dissociation between the effects of diet on the testosterone-secreting and the spermatogenic functions of the testis during winter and spring (Figs 2c, 3b). Nutritionally induced dissociation between testicular growth and testosterone secretion also occurs in Merino rams (Ritar et al., 1984; Martin et al., 1987; Martin et al., 1994a, b), and may be a characteristic of male ruminants living in environments where food supply is erratic or out of phase with normal photoperiodic responses. In cashmere bucks, derived from feral animals living in semi-arid central Australia, such dissociation may allow testicular development in response to changing nutrition to precede behavioural responses induced by testosterone, ensuring that the testes are well developed by the time of mating.

The results support our original hypotheses that Australian cashmere bucks exhibit considerable reproductive seasonality, and that the expression of this seasonality is modified by the nutritional environment. Of the reproductive variables examined, testosterone concentration, sebaceous gland volume and odour score appeared to be ultimately dependent upon LH secretion and under fairly rigid seasonal control (probably photoperiodic), with the effects of enhanced nutrition limited to periods when seasonal inhibition is waning. However, changes
in testicular mass (and therefore sperm production) appeared to be driven primarily by changes in the voluntary intake of feed and in growth, often independently of changes in gonado-
trophin concentration. Enhanced nutrition is thus able to induce rapid testicular growth, but not increased testosterone production, during winter and spring when photoperiodic inhibition of the hypothalmo–pituitary–testicular axis is maximal.

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