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Differences in peripartal plasma parameters related to calcium homeostasis of dairy sheep and goats in comparison with cows

Mirja R. Wilkens*, Annette Liesegang, Julia Richter, David R. Fraser, Gerhard Breves, Bernd Schröder

aDepartment of Physiology, University of Veterinary Medicine Hannover, Foundation, Hannover, Germany

bInstitute of Animal Nutrition, Vetsuisse Faculty Zurich, University of Zurich, Zurich, Switzerland

cCenter for Applied Biotechnology and Molecular Medicine, University of Zurich, Zurich, Switzerland

dFaculty of Veterinary Science, University of Sydney, Sydney, Australia

*Corresponding author: Mirja R. Wilkens, Department of Physiology, University of Veterinary Medicine Hannover, Bischofsholer Damm 15/102, 30173 Hannover, Germany.

Tel.: + 49 511 856 7628; fax: + 49 511 856 7687.

E mail address: mirja.wilkens@tiho-hannover.de

Key words: transition; calcium homeostasis; close-up; dry period; ruminants; bone metabolism;

Short title: Peripartal calcium homeostasis in ruminants
Summary:

Recently it has been demonstrated that there are differences between sheep and goats in respect to the adaptation to a calcium restricted diet. It was the aim of the present study to evaluate whether species-specific peculiarities also occur when calcium homeostasis is challenged by lactation. Therefore, we investigated the time courses of plasma parameters related to calcium homeostasis (calcium, phosphate, calcitriol, the bone resorption marker CrossLaps® and the bone formation marker osteocalcin) during the transition period in multiparous animals of both species and compared the results to data from a former study carried out with dairy cows. Like in cows, plasma calcium and the ratio of bone formation to bone resorption decreased at parturition in goats while plasma calcitriol increased. On the 10th day postpartum the bone parameters of goats reached prepartum values again which was not the case in cows. Sheep were found to experience a challenge of calcium homeostasis already 10 days before parturition, reflected by a very low ratio of bone formation to bone resorption which was not accompanied by an increase in plasma calcitriol. Additionally, sheep and goats which had been in milk for 3 months were sampled, dried-off and sampled again 6 weeks later. In dried-off animals there were no differences in parameters of bone metabolism detectable. In conclusion we could show that the contribution of bone mobilisation to the compensation for the enhanced calcium demand due to lactation differs between the three ruminant species.
Introduction

Small ruminants are often used as models for cows to study basic aspects of calcium (Ca) homeostasis. However, there are differences in homeostatic mechanisms between goats and sheep that need to be considered before any results can be extrapolated to dairy cows, especially with respect to the transition period.

On a body weight basis the milk yield of dairy goats is comparable to that of dairy cows and hypocalcemia occurs in both species usually at the onset of lactation, while sheep develop hypocalcemia more often during late gestation (Oetzel, 1988). But when comparing feeding behaviour, cattle and sheep can be regarded as grazers, while goats in their natural habitat select high energy (concentrate) feed (Hofmann, 1989). These differences in feed type might influence gastrointestinal mineral absorption as it has been shown that goats adapt more efficiently to dietary Ca restriction by up-regulating intestinal Ca transport than sheep (Wilkens et al., 2011; Wilkens et al., 2012b). Interestingly, it has also been reported that a reduction in Ca supply did not lead to an increase in apparent digestibility of Ca in lactating cows (Taylor et al., 2008).

Comparative data on the response of different ruminant species to the challenge of Ca homeostatic mechanisms at the onset of lactation are limited. In studies investigating bone metabolism in goats and sheep it has been demonstrated that around parturition sheep tend to have greater concentrations of bone resorption markers and more pronounced depression of bone formation markers in blood, compared to goats (Liesegang et al., 2006, 2007).

As milk fever in dairy cows is associated very closely with parturition, it was the aim of the present study to investigate changes in plasma parameters of Ca homeostasis during the peripartal period of sheep and goats by taking samples in shorter intervals than it has been done in the studies mentioned above and compare them to respective data from cows. Based on our former studies we hypothesise that there are differences in the response to the peripartal challenge of Ca homeostasis which cannot be simply explained with the higher milk yield usually observed in goats and cows.
Besides ionised and total Ca and phosphate plasma, concentrations of calcitriol, the biologically active metabolite of vitamin D, were determined. Furthermore, the bone markers were addressed. Bone turnover or remodelling means concurrent bone formation and resorption (Allen, 2003). As absolute values of biochemical markers of bone metabolism are influenced by various factors like age, circadian rhythms and season (Hannon & Eastell, 2000), the ratio of osteocalcin to CrossLaps® was used to assess bone turnover. The ratio of bone formation markers to bone resorption markers has been used before to estimate the status of bone remodelling in horses and cows (Lepage et al., 1998; Liesegang et al., 2000).

Material and methods

Animals, diets, milking pattern and blood sampling

The study was carried out with 5 to 6 multiparous animals of each species (sheep, goats and cows), aged between 2 and 5 years. All animals were of breeds used for milk production in Central Europe: East Friesian dairy sheep, Saanen type goats and Holstein-Friesian cows. Average body weights were 66.3 ± 1.6 kg (sheep), 49.1 ± 1.7 kg (goats) and 679 ± 25 kg (cows). Sheep and goats were housed in a free-stall barn at the Department of Physiology of the University of Veterinary Medicine, Foundation, Hanover, Germany. During the entire observation period, the animals had access to hay (Table 1), minerals and water ad libitum. The animals were group-fed prepartum, receiving 75 g concentrate (Table 1) per 10 kg body weight per day.

Immediately after parturition, the kids and lambs were removed and milk yield was recorded for 20 consecutive days. The ewes and does were milked and provided with concentrate three times daily at 0800, 1400 and 2000 h. The amount of concentrate was calculated for each individual animal according to the milk yield of the previous milking (800 g per kg milk) to meet the requirements for dairy sheep and goats recommended by the Society of Nutrition Physiology (GfE).
Data from cows were obtained from a previous study (Wilkens et al., 2012a). The cows were kept on a commercial dairy farm located in Saxony, Germany. They were fed a total mixed ration ad libitum before and after parturition (Ca content a.p.: 7.8 g/kg DM; Ca content p.p.: 10.0 g/kg DM). The calves were removed within 24 h postpartum, after which milking was done twice daily. In the present study, the peripartal plasma parameters of 6 cows of the former control group were compared with respective values from the small ruminants.

From calculated gestational day 140 (sheep and goats) or day 175 (cows), two blood samples (lithium heparin and EDTA) were collected from the jugular (sheep and goats) or coccygeal vein (cows), every other day at 0900 h until day 10 postpartum. Additional samples were taken immediately after parturition and at 6, 12 and 24 h postpartum. The prepartum samples were re-classified according to the exact day of parturition. Samples taken on day 2 and 1 prepartum were classified as sample day -2, samples taken on day 4 and 3 as sample day -4, etc.

A further 6 goats and 5 sheep, of the same age and breed were sampled after three months of lactation. They were then dried off and sampled again six weeks later.

The protocol of animal treatment was approved by the Animal Welfare Commissioner of the University of Veterinary Medicine, Foundation, Hanover, Germany, and its conduct was supervised according to German Animal Welfare Law.

Sample analysis

After measuring ionised Ca in whole blood with a blood gas analyser (Chiron Diagnostics, RapidLab™ 348), samples were centrifuged (2,000 g, 15 min) at ambient temperature and plasma was collected and stored at -18°C.

The concentrations of phosphate and total Ca in plasma of sheep and goats were measured colorimetrically by standard spectrometric techniques (Sarkar & Chauhan, 1967; Kruse-Jarres, 1979), while total Ca and phosphate concentrations in the plasma samples from cows were
determined by standard diagnostic methods in the laboratory of the Clinic for Cattle of the University of Veterinary Medicine, Foundation, in Hannover. Plasma calcitriol of sheep and goats was measured with a commercial radioreceptor assay (Immundiagnostik AG, Bensheim, Germany). Quantification of the bone formation marker osteocalcin and the bone resorption marker CrossLaps® was done using commercial enzyme-linked immunosorbent assay kits according to the manufacturer’s instructions (MicroVueTM Osteocalcin EIA Kit, Quidel® Corporation, Santa Clara, USA; Serum CrossLaps®, IDS Ltd, Frankfurt am Main, Germany).

Data presentation and statistical analysis

Data are presented as mean ± SEM. The daily milk yield of sheep and goats is corrected for the animals’ body weights. All statistical analyses were performed using GraphPad Prism® 5.0 (GraphPad Software, San Diego, USA). Gaussian distribution according to Kolmogorov-Smirnov was confirmed for all data. Comparison of peripartal ionised and total Ca, phosphate, CrossLaps® and osteocalcin plasma concentrations was carried out by analysis of variance (ANOVA) for repeated measurements with a saturated model of the fixed factor species. Bonferroni post-test was applied to reveal species differences at each time point and differences over time within each species. In all cases, $P$ values < 0.05 were considered statistically significant.

Results

Milk yield

Three of the 5 ewes and 5 of the 6 does produced twins, while the remaining small ruminants and all the cows had single offspring. Milk yield was not affected by the number of offspring. There was no significant difference in milk yield between the sheep and goats over the first few
days after parturition. However, daily milk production of the goats then increased to a maximum of 0.085 l/kg BW, whereas milk yield of the ewes remained more or less constant at a level of 0.04 l/kg BW per day (Fig. 1).

Macro minerals in blood and plasma

Whole blood and plasma concentrations of ionised and total Ca and phosphate measured in sheep, goats and cows around parturition are given in Figures 2, 3 and 4. Compared to day 2 prepartum, ionised Ca measured in whole blood of goats showed a significant decrease at parturition and also on the 10th day postpartum. There was no similar change in ionised Ca associated with lambing in ewes. The transient decrease observed in cows with a minimum at 12 hours postpartum was not statistically significant because of high individual variances, but compared to sheep and goats postpartum ionised Ca in cows was significantly lower. However, the great challenge of Ca homeostasis in cows at parturition becomes obvious when the plasma concentrations of total Ca are considered. In all species, a small, but not statistically significant decrease in plasma phosphate was observed at parturition. Intriguingly, a pronounced increase in plasma phosphate was found on the 10th day postpartum in goats.

Calcitriol

The profile of plasma concentrations showed that in goats, calcitriol was highest at parturition while sheep had no significant changes and cows showed the highest calcitriol levels on the 2nd day postpartum (Fig. 5).
The ratio of the bone formation marker osteocalcin to the bone resorption marker CrossLaps® is given in Figure 6. In sheep plasma, a very low ratio was found throughout the entire observation period, while goats and cows had significantly greater ratios during the last gestational days which then decreased at parturition. On the 10th day postpartum basal values were found again in goats, while the cows’ osteocalcin to Crosslapps® ratios remained as low as those of sheep.

Effects of terminating lactation

Comparisons of plasma concentrations of total Ca, phosphate, calcitriol and the ratio of osteocalcin to CrossLaps® in ewes and does after three months of lactation and six weeks after the end of lactation are presented in Table 2. Plasma concentrations of total Ca were not affected by lactation status. However, lower concentrations were found in both groups of sheep compared to goats. Both species showed a small increase in plasma phosphate after lactation had ceased. While no effects of lactation status were found with calcitriol concentrations, the ratio of osteocalcin to CrossLaps® increased markedly in both species after lactation had ceased. In contrast to peripartal values, there were no species differences in these bone marker ratios in non-lactating animals.

Discussion

There is a sudden increase in the outflow of Ca from blood at the onset of lactation compared to the last days of gestation (Horst et al., 2005). The resulting transient decrease in plasma Ca at parturition and shortly after has been described for cows in numerous studies (Reinhardt et al., 2011). The discrepancy between ionised and total Ca in cows after parturition in the present study could be explained by factors caused by the negative energy balance almost always reported for cows in early lactation (Butler & Smith, 1989). For example, alterations of plasma albumin
or pH might have affected the ratio between ionised and total Ca. In contrast to the group fed
dairy cows, the small ruminants were fed individually according to their milk yield.
The absence of such a depression of Ca at parturition in sheep has already been reported
(Liesegang, 2008). This might suggest that in contrast to goats and cows there is no particular
challenge to the mechanisms of Ca homeostasis at lambing due to the secretion of Ca in milk.
However, as there were only minor differences in milk production between sheep and goats
immediately after parturition, it is unlikely that the Ca demand for milk production during the
first 24 hours p.p. was very much higher in goats compared to sheep. The daily placental transfer
in ewes between day 133 and 140 of gestation has been shown to amount for 255 mg per kg
foetal body weight (Durand et al., 1983). Assuming a Ca concentration of 2.5 g/L in colostrum
(Liesegang 2008), for the sheep in the present study the loss of Ca into the milk has been
approximately 4 g on the first day p.p. compared to an estimated daily demand of 2 g for late
gestation.
But in contrast to does and cows, the osteocalcin to CrossLaps® ratio in ewes was already very
low at the end of gestation. Since in non-lactating animals there were no differences between
goats and sheep, this low formation to resorption marker ratio is most probably not
characteristic for sheep in general nor caused by variations in the sensitivities of the ELISA
kits. The time course of the osteocalcin to CrossLaps® ratio around parturition more probably
suggests that in ewes the Ca homeostatic mechanisms have already been challenged before
lambing and there may have been a greater contribution of the skeleton to maintenance of Ca
homeostasis than in the other two ruminant species. This is in line with the clinical observation
that ewes often develop hypocalcemia during the last days before lambing when the foetal
skeleton is mineralised (Oetzel, 1988) and that the peripartal plasma Ca concentrations are
lower in twin-bearing than in single-bearing ewes (Raoofi et al., 2013).

However, all of the goats except one in the present study also produced twins. As it has already
been demonstrated that goats have a greater ability to adapt to dietary Ca restriction than sheep
(Wilkens et al., 2012b), it may be that the does were adapted better than sheep to the enhanced Ca demand of foetal bone growth in late pregnancy. The onset of lactation induced a small decrease in ionised Ca accompanied by a concurrent increase in calcitriol in does. A decrease in ionised Ca leads to a secretion of parathyroid hormone (PTH) which than stimulates the renal \(1\alpha\)-hydroxylase, the enzyme needed for the transformation of 25-hydroxyvitamin D to 1,25-dihydroxyvitamin D, the active hormone calcitriol (Fraser & Kodicek, 1970).

Assuming that Ca homeostasis in ewes had already been severely challenged before parturition, one would expect a calcitriol response occurring even earlier in these animals. However, this was not the case. Lower calcitriol concentrations in sheep compared to goats experiencing the same dietary Ca restriction have already been demonstrated (Wilkens et al., 2010; Wilkens et al., 2012b). As there is no suitable assay available for the determination of PTH in small ruminant species, it is difficult to reveal the reasons for this phenomenon.

In cows and goats bone resorption seemed to be induced not before parturition. But while goats reached pre-kidding values at the 10th day postpartum again, cows’ and sheep’s osteocalcin to CrossLaps® ratios remained very low. This might point to are more rapid stimulation of gastrointestinal Ca absorption in goats allowing the animals do reduce bone mobilisation again.

For sheep it has been shown that the maximal efficiency of gastrointestinal Ca absorption was reached 2 months after parturition, although bone resorption had already been stimulated before lambing (Braithwaite et al., 1970). Comparable balance studies carried out with goats are not available. In cows a significant increase of gastrointestinal Ca absorption was found to take place one week postpartum (van’t Klooster, 1976) or between the 10th and the 20th day after parturition, respectively (Ramberg et al., 1970). It should be taken into account that these data were obtained some time ago when milk production in dairy cows was not as high as nowadays.

Against the background of the low osteocalcin to CrossLaps® ratios found in cows on the 10th day postpartum it might be speculated that bone mobilisation was more prolonged in cows of the present study.
The reason for the slight decrease in plasma phosphate observed in all species is usually assumed to represent a loss of phosphate to the milk (Robertson et al., 1956). Furthermore, it has been shown that a decrease in plasma Ca induces a concurrent increase in cortisol. Cortisol was demonstrated to have an impact on plasma phosphate (Horst & Jorgensen, 1982). No satisfying explanation could be found for the increase in goats’ plasma phosphate concentrations on the 10th day postpartum. As the blood concentration of ionised Ca showed a second decrease on the 10th day, too, these alterations might be associated with the steep rise in milk production observed in does between the 7th and the 12th day after parturition. The role of phosphate in lactating animals is not yet fully understood. In contrast to calcium, plasma concentrations of phosphate are not that strictly controlled and the exchange between the different pools is more complicated. Phosphate is not only mobilised from bone, but also from soft tissues and in ruminants large amounts of phosphate secreted via the salivary glands and reabsorbed again in the gastrointestinal tract have to be taken into account (Braithwaite, 1983). However, the effect of drying-off on plasma phosphate underlines that it might be influenced by the hormonal regulation mechanisms involved in lactation and bone metabolism.

Conclusion

We could demonstrate that in line with former studies sheep seem not to respond with an adequate stimulation of calcitriol synthesis to challenges of Ca homeostasis. Furthermore, striking differences regarding peripartal bone metabolism were observed between the three ruminant species investigated. To evaluate whether sheep or goats are the better model for studying Ca homeostasis in the dairy cow, further studies under more controlled feeding regimes of the cows and a larger number of animals are needed.

The financial support of the Deutsche Forschungsgemeinschaft (SCHR 342/8-2) is gratefully acknowledged.
Table 1

Ingredients of hay and concentrate fed to sheep and goats expressed as percentage of original substance; n.d.: not detectable using standard procedures.

<table>
<thead>
<tr>
<th></th>
<th>Hay</th>
<th>Concentrate</th>
</tr>
</thead>
<tbody>
<tr>
<td>Crude protein</td>
<td>8.21</td>
<td>20.0</td>
</tr>
<tr>
<td>Crude fat</td>
<td>1.52</td>
<td>2.60</td>
</tr>
<tr>
<td>Crude fibre</td>
<td>26.5</td>
<td>11.0</td>
</tr>
<tr>
<td>Crude ash</td>
<td>7.30</td>
<td>8.70</td>
</tr>
<tr>
<td>Calcium</td>
<td>0.41</td>
<td>0.90</td>
</tr>
<tr>
<td>Phosphorus</td>
<td>0.32</td>
<td>0.55</td>
</tr>
<tr>
<td>Sodium</td>
<td>0.11</td>
<td>0.20</td>
</tr>
<tr>
<td>Energy [MJ per kg]</td>
<td>7.50</td>
<td>10.2</td>
</tr>
<tr>
<td>Vitamin D [IU/kg]</td>
<td>n.d.</td>
<td>1,000</td>
</tr>
</tbody>
</table>
Table 2

Total calcium (mmol/l), phosphate (mmol/l), calcitriol (pg/ml) and the ration between osteocalcin (ng/ml) and Crosslaps® (ng/ml) in plasma of goats and sheep measured after 3 months in milk and 6 weeks after having been dried-off, respectively. Results of 2way ANOVA (species, effect of lactation and interaction). Means ± SEM.

<table>
<thead>
<tr>
<th></th>
<th>Total calcium</th>
<th>Phosphate</th>
<th>Calcitriol</th>
<th>osteocalcin to CrossLaps®</th>
</tr>
</thead>
<tbody>
<tr>
<td>Goats</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Lactating (6)</td>
<td>2.56 ± 0.06</td>
<td>1.69 ± 0.09</td>
<td>12.8 ± 0.86</td>
<td>46 ± 5.8</td>
</tr>
<tr>
<td>Dried-off (6)</td>
<td>2.63 ± 0.03</td>
<td>1.87 ± 0.14</td>
<td>12.8 ± 0.82</td>
<td>62 ± 5.2</td>
</tr>
<tr>
<td>Sheep</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Lactating (5)</td>
<td>2.20 ± 0.06</td>
<td>1.59 ± 0.08</td>
<td>13.4 ± 3.35</td>
<td>35 ± 5.3</td>
</tr>
<tr>
<td>Dried-off (5)</td>
<td>2.18 ± 0.09</td>
<td>2.01 ± 0.16</td>
<td>8.1 ± 0.90</td>
<td>62 ± 12</td>
</tr>
</tbody>
</table>

ANOVA

<table>
<thead>
<tr>
<th></th>
<th>Species</th>
<th>Lactation</th>
<th>Interaction</th>
<th></th>
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<tbody>
<tr>
<td></td>
<td>P &lt; 0.001</td>
<td>n.s.</td>
<td>n.s.</td>
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</table>
Figures

Figure 1
Daily milk yield of goats and sheep normalised to the animals’ body weight. RM ANOVA revealed significant effects of time ($P < 0.001$), species ($P < 0.001$) and an interaction of time and species ($P < 0.001$). Significant differences between species revealed by Bonferroni post-test for each day are marked with asterisks ($P < 0.05$: *; $P < 0.01$: **; $P < 0.001$: ***). Means ± SEM.

Figure 2
Blood concentrations of ionised calcium measured in goats, sheep and cows around parturition. RM ANOVA revealed significant effects of species ($P < 0.01$) and an interaction of time and species ($P < 0.05$). Different letters represent significant differences within one species. The symbols # and $ indicate significant differences between cows and goats (#) or sheep ($), respectively. Means ± SEM.

Figure 3
Plasma concentrations of total calcium measured in goats, sheep and cows around parturition. RM ANOVA revealed significant effects of time ($P < 0.001$) and species ($P < 0.001$). Different letters represent significant differences within one species. The symbols # and $ indicate significant differences between cows and goats (#) or sheep ($), respectively. Means ± SEM.

Figure 4
Plasma concentrations of phosphate measured in goats, sheep and cows around parturition. RM ANOVA revealed significant effects of time ($P < 0.001$) and an interaction of time and species
(P < 0.001). Different letters represent significant differences within one species. The symbols # and § indicate significant differences between goats and cows (#) or sheep (§), respectively. Means ± SEM.

**Figure 5**

Plasma concentrations of calcitriol measured in goats and sheep around parturition. RM ANOVA revealed significant effects of time (P < 0.05) and species (P < 0.05). Different letters represent significant differences within one species. Means ± SEM.

**Figure 6**

Ratio between plasma concentrations of osteocalcin and CrossLaps® measured in goats, sheep and cows around parturition. RM ANOVA revealed significant effects of time (P < 0.001), species (P < 0.001) and an interaction of time and species (P < 0.001). Different letters represent significant differences within one species. The symbols #, $ and § indicate significant differences between goats and cows (#), sheep and cows ($) or sheep and goats (§), respectively. Means ± SEM.
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Fig. 1
**Ca\(^{2+}\) in mmol/L**

- **Goats (6)**
- **Sheep (5)**
- **Cows (6)**

**Fig. 2**
Fig. 3
Fig. 4
Calcitriol in pg/mL, goats and sheep

**Fig. 5**

- Goats (6)
- Sheep (5)