The relationships between potassium intakes, transmural potential difference of the rumen epithelium and magnesium absorption in wethers

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(Received 21 May 2003 – Revised 6 October 2003 – Accepted 15 October 2003)

In vitro studies with isolated sheep rumen epithelium have shown that an increase in the lumen K concentration induces an increase in the transmural potential difference across the rumen epithelium (serosal side: positive), which is associated with a decrease in Mg transport. However, at lumen K concentrations >80 mmol/l, Mg transport across the epithelium became independent of the lumen K concentration. The present study was carried out to determine whether this observation also occurs in vivo. Four ruminally fistulated wethers were fed four rations supplemented with KHCO₃ (15.7, 37.6, 59.4 or 77.4 g K/kg DM) in a 4 × 4 Latin square design. Increased K intakes significantly increased the rumen K concentration. For all data combined, Mg absorption expressed as % intake was negatively correlated with the transmural potential difference (blood side: positive) (Pearson’s r = −0.709; R² = 0.468, P = 0.002, n = 16). It is concluded that in wethers apparent Mg absorption becomes independent of the dietary K concentration when the K concentration is >60 g/kg DM or equivalent to a postprandial rumen K concentration of about 125 mmol/l.

Magnesium absorption: Potassium intake: Transmural potential difference: Sheep

Pastures in areas with intensive livestock production are generally rich in K (Fisher et al. 1994; Schonewille et al. 1997) due to frequent fertilization with manure, which may be injected directly into the soils (Schonewille et al. 1997). Consequently, K concentrations of ≥35 g/kg DM are common in fresh grass. Grazing cattle on such pastures entails the risk of hypomagnesaemia (Kemp, 1960), primarily due to a K-induced decrease in Mg absorption (Fontenot et al. 1989). In ruminant animals, Mg absorption mainly takes place in the rumen (Rogers & Van’t Klooster, 1969; Tomas & Potter, 1976) and consists of a K-insensitive, carrier-mediated process and a K-sensitive, electrogenic transport component (Leonhard et al. 1989). Mg uptake through the electrogenic transport component depends on the concentration of Mg in the soluble fraction of rumen contents and the apical membrane potential (PDa) across the rumen epithelial cells (intracellular contents are negative relative to rumen contents). At low rumen K concentrations, the PDa provides a driving force for Mg uptake by rumen epithelial cells, but high rumen K concentrations cause a depolarization of the PDa; this reduces Mg uptake (Leonhard-Marek & Martens, 1996). Thus, the observed decrease in Mg absorption at high rumen K concentration is caused by the K-induced change in the PDa. Since the transmural potential difference (PDT) is directly related to PDa, it follows that a change in PDa causes an alteration of PDT (Leonhard-Marek & Martens, 1996). Indeed, it was shown that the K-induced decrease in Mg absorption was highly correlated with an increased PDT (Martens & Blume, 1986; Martens et al. 1987).

It has been demonstrated (Ferreira et al. 1966; Leonhard-Marek & Martens, 1996) that, under in vitro conditions with isolated sheep rumen epithelium, both PDT and PDa are correlated with the logarithm of the lumen K concentrations. This means that the change in PDt and PDa is much greater when the lumen K concentration increases from 20 to 40 mmol/l compared with the same increase from a concentration of 80 mmol/l. Indeed, net Mg fluxes became independent of lumen K at concentrations >80 mmol/l (Leonhard-Marek & Martens, 1996), which implies that the K-induced reduction in Mg absorption might be non-linear under practical feeding conditions. Balance data from sheep indicate that the depressant effect of supplemental K on Mg absorption becomes lower at higher initial dietary K concentrations (Greene

Abbreviations: PDa, apical membrane potential; PDT, transmural potential difference.
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et al. 1983a,b; Grings & Males, 1987). Furthermore, in an experiment with sheep it was shown that extra K did not affect Mg absorption when the dietary K content was increased from the high initial value of 39 to 82 g/kg DM (Rahnema & Fontenot, 1986). Because neither rumen K concentrations nor PDt were measured in those feeding studies, the proposed mechanism of the K-induced inhibition of Mg absorption described earlier remains to be validated in vivo. Thus, we studied the effect of increasing K intakes on the PDt and Mg absorption to check the linearity of the K-induced decrease in Mg absorption under feeding conditions. K intake was increased by supplementing the basal concentrate with KHCO₃, which is as effective as intrinsic K salts in depressing Mg absorption (Schonewille et al. 1999b). Therefore, KHCO₃ was used to obtain a range of 15–75 g K/kg DM, which is beyond the practical range of 23–49 g K/kg DM (Centraal Veevoederbureau, 2003), but it was anticipated that the chosen range of dietary K concentrations would enhance the interpretation of the data.

Materials and methods

Animals and experimental design

The experimental protocol was approved by the Animal Experiments Committee of the Utrecht Faculty of Veterinary Medicine. Four ruminally fistulated Zwartbles wethers were used. They were 1·5 years old and weighed 61 (SE 1·22) kg. The wethers had been fistulated at least 1 month before the start of the experiment. They were housed in individual pens (1·9 × 1·2 m) with a layer of wood shavings, or in metabolism crates with slatted floors. The trial had a 4 Latin square design and was preceded by 14 d pre-experimental period. The wethers were randomly assigned to each sequence of feeding of the four experimental rations. Each experimental period lasted 28 d. Tap water was available ad libitum.

Experimental diets

The experimental diets were formulated by the addition of KHCO₃ to the basal concentrate. The ingredient composition of the pelleted concentrates (diameter 3 mm) is shown in Table 1. During the pre-experimental period, all wethers were offered 582 g commercial concentrate and 129 g hay/d. Thereafter, the sheep were fed one of the four experimental diets. The amount of feed offered and the analysed composition of the experimental diets was shown in Table 2. The wethers were fed a restricted amount of the experimental diets so as to maintain constant intakes of non-variable nutrients; the calculated metabolizable energy intake was 7·5 MJ/d for each diet (Centraal Veevoederbureau, 2003). In each 24 h interval the diets were offered three times per d in three equal portions; the portions were given at 00.00, 08.00 and 16.00 hours.

Sample collection

The experimental concentrates and hay were sampled from days 18 to 28 of each period, were ground, and subsequently stored in a sealed jar at room temperature (18°C). From days 18 to 25 of each experimental period, faeces were collected quantitatively. The 24 h faeces collections were stored at −18°C in plastic bags. At the end of each experimental period, the faeces collections were pooled for each wether and mixed thoroughly. Two samples each representing 10% total faeces collection from each wether for each experimental period were dried for 5 d at 60°C, ground, and stored in a sealed jar at room temperature.

On day 26 of each experimental period, before the morning meal, 70 ml Cr-EDTA solution (100 g Cr-EDTA/l, pH 6·5–6·7) was introduced into the rumen by the cannula as a marker to assess rumen volume and passage rate of the rumen liquid phase. Samples of the rumen liquid was subsequently stored in a sealed jar at room temperature.

### Table 1. Compositions of the experimental concentrates

<table>
<thead>
<tr>
<th>Ingredients (g)</th>
<th>Low K</th>
<th>Medium K</th>
<th>High K</th>
<th>Extreme K</th>
</tr>
</thead>
<tbody>
<tr>
<td>KHCO₃</td>
<td>994</td>
<td>994</td>
<td>994</td>
<td>994</td>
</tr>
<tr>
<td>MgO</td>
<td>6</td>
<td>6</td>
<td>6</td>
<td>6</td>
</tr>
</tbody>
</table>

*The constant components consisted of (g): sugar beet pulp 250, soyabean meal 200, sugarcane molasses 37, linsed meal 120, maize gluten feed 150, maize meal 230, NaCl 5, premix 2. The premix consisted of (mg/g): ZnSO₄·H₂O 67·5, MnSO₄·H₂O 57·5, CuSO₄·H₂O 25·0, KIO₃ 0·6, CoSO₄·7·H₂O 0·4, Na₂SeO₃·5H₂O 0·4, retinyl acetate preparation 0·9, cholecalciferol preparation 4·2, α-tocopherol acetate preparation 25·0, CaCO₃ 818·5.

### Table 2. Intake and analysed compositions of the experimental rations

<table>
<thead>
<tr>
<th>Experimental ration</th>
<th>Low K</th>
<th>Medium K</th>
<th>High K</th>
<th>Extreme K</th>
</tr>
</thead>
<tbody>
<tr>
<td>Intake (g DM per sheep per d)</td>
<td>535</td>
<td>571</td>
<td>603</td>
<td>638</td>
</tr>
<tr>
<td>Nutrient composition of total rations (g/kg DM)</td>
<td>166</td>
<td>167</td>
<td>162</td>
<td>154</td>
</tr>
</tbody>
</table>

*For details of the composition of the concentrate, see Table 1.
†The analysed DM contents were as follows (g/kg concentrate): low K: Mg 8·7, K 14·8, Na 2·9, Ca 5·6, P 5·3; medium K: Mg 6·8, K 41·3, Na 2·8, Ca 5·4, P 5·1; high K: Mg 6·4, K 67·4, Na 2·6, Ca 5·2; P 4·8; extreme K: Mg 5·9, K 88·3, Na 2·4, Ca 4·9, P 4·5.
‡The analysed DM content of hay was 935 g/kg and the analysed composition was (g/kg DM): crude protein (N × 6·25) 104, crude fat 16, crude fibre 312, Mg 1·4, K 19·7, Na 1·6, Ca 4·1, P 2·1.
Magnesium absorption in sheep

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Measurement of potential difference

The PDa is of primary interest to explain the K-induced inhibition of Mg uptake by rumen epithelial cells, but it cannot be measured under feeding conditions. PDa is strongly correlated with PDi (Leonhard-Marek & Martens, 1996). Therefore, we measured PDi in the present study. The measurement of the potential difference between blood (jugular vein) and rumen contents has been described by Schonewille et al. (1999). PDi was measured continuously (09.00 to 15.00 hours) on day 28 of each period. Briefly, the PDi was monitored twice per s using an analogue to digital converter (Data Acquisition Board, PCI 20428W-1; Intelligent Instrumentation Inc., Tucson, AZ, USA) built into a computer. The analogue to digital converter was connected with two separate reference electrodes (REF201; Radiometer, Paris, France). Each electrode was dipped into 500 ml saturated KCl solution (3.5 M) and the rumen and blood electrodes were connected by saturated KCl–agar bridges.

Chemical analyses

N was determined by the macro-Kjeldahl method (International Dairy Federation, 1986); a factor of 6.25 was used to convert N mass into crude protein. Dithethyl ether extracts of the feedstuffs were prepared according to the Association of Official Analytical Chemists (1984); the solvent was evaporated and the crude fat residue was weighed. The crude fibre content of feedstuffs was estimated according to the method of the Association of Official Analytical Chemists (1984). Feedstuffs and faeces samples were ashed at 480°C for 6 h to enable analysis for Mg, K, Na, Ca and P. The ash samples were dissolved in 15 ml 4M-HCl and the acidified solution was analysed for Mg, Ca and K by atomic absorption spectroscopy and for Na by atomic emission spectroscopy (Perkin Elmer 3110; Perkin Elmer Corp., Norwalk, CT, USA). Total P in feedstuffs was measured with a spectrophotometer following the method of Quinlan & DeSesa (1955). The concentrations of Mg and K in rumen contents were determined directly by atomic absorption spectroscopy and those of Na and Cr(III) in rumen contents were measured directly by atomic emission spectroscopy (Ram et al. 1998). The accuracy of each assay run was found to be within 3% as the variation of a commercial hay powder (CRM 129; Community Bureau of Reference, Brussels, Belgium) or from that of in-house reference samples. Mg concentration in plasma was measured directly by atomic absorption spectroscopy (Perkin-Elmer, 1982). The combined within- and between-run precision of the determinations (CV) was ≤3.0%.

Calculations and statistical analyses

Before statistical analysis, the PDi data for each experimental animal were pooled for each measuring day. The calculations to obtain rumen volume and the passage rate of rumen contents have been described by Schonewille et al. (1999). All data were subjected to ANOVA with animal, experimental period and dietary treatment as factors (Wilkinson, 1990). When the influence of treatment was significant, Tukey’s t test was used to identify diets with different effects on the variable involved. For the data from each sheep (n = 4) and for each diet (n = 4), linear correlations were calculated between rumen variables, PDi and Mg absorption. For the calculations it was assumed that the sixteen data points could be considered to be independent. To detect rumen variables that were related to Mg absorption, multiple regression analysis was performed (Wilkinson, 1990) with animal and period as factor, Mg absorption as dependent variable and rumen content concentrations of Mg and K, rumen pH, rumen volume, passage rate of rumen contents and PDi as independent variables. Forward stepwise regression was performed by incorporating into the model the rumen variable showing the highest significant partial correlation coefficient for its relationship with Mg absorption. Throughout, the level of significance was set at P<0.05.

Results

Feed intake and body weight

The wethers consumed all feed supplied. The experimental rations did not significantly influence the body weight of the sheep. For all treatments combined, body weight was 63 (SE 0.3) kg (n = 4).

Magnesium absorption and plasma magnesium

Intakes of Mg were similar for all treatments (Table 3). Faecal Mg excretions were significantly increased after feeding the rations containing the two highest levels of K; the difference in faecal Mg excretion between these two treatments was not significant. Consequently, both absolute Mg absorption (g/d) and Mg absorption expressed as % intake were significantly decreased on the high- and extreme-K rations. Because Mg intake was essentially constant, absolute Mg absorption (g/d) and Mg absorption expressed as % intake were highly correlated (Pearson’s r 0.998, P<0.001, n = 16).

The difference in K intakes among treatments did not significantly affect plasma Mg concentrations. For all treatments combined, plasma Mg concentration was 1.0 (SE 0.03) mmol/l (n = 4).
Mineral concentrations in rumen contents, rumen pH and transmural potential difference

Mg concentrations in rumen contents both before and after feeding were not significantly affected by the level of K intake (Table 4). Although ANOVA indicated a borderline significant influence of K intake on rumen pH, significant differences between specific rations could not be identified by Tukey's test. There was a significant correlation between the mean postprandial rumen pH and the mean postprandial Mg concentration in rumen contents (Pearson's $r = 0.663$, $P = 0.005$, $n = 16$).

The level of K intake significantly increased the rumen K concentrations and decreased the rumen Na concentrations in a dose-dependent fashion (Table 4). A strong negative correlation was observed between the mean postprandial K and Na concentrations in rumen contents (Pearson's $r = 0.909$, $P < 0.001$, $n = 16$).

The increase in K intake was associated with an increase in PDt (blood side: positive) (Table 4). The increase in the PDt after feeding the medium- v. low-K intake was not significant ($P = 0.090$), but PDt was significantly increased by the two highest levels of K intake when compared with the low-K ration. There was a significant, positive association between the mean postprandial rumen K concentrations and the PDt; the linear regression was $y = 43.82 + 0.17x$ ($R^2_{adj} = 0.468$, $P = 0.002$, $n = 16$).

Rumen volume and passage rate

Rumen volume was significantly increased when dietary K intake was increased from the low to extreme level (Table 5). The fractional outflow (%/h) of the liquid phase was increased after feeding of the medium-K ration, but was not significantly different for the low, high and extreme levels of K intake. The absolute outflow rate of the liquid phase was not influenced by the level of K intake.

Multiple regression analysis

The multiple regression model with a constant, the factors (animal and experimental period) and the independent rumen variables (Mg concentration, PDt, rumen volume and passage rate of rumen contents) explained 78.8 % of the observed variance in Mg absorption ($P = 0.025$). After

<table>
<thead>
<tr>
<th>Table 3. Intake, faecal excretion and absorption of magnesium in wethers fed the experimental rations*</th>
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<tbody>
<tr>
<td>(Mean values for four sheep per group)</td>
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<table>
<thead>
<tr>
<th>Experimental ration</th>
<th>Statistical significance of effect: $P_{disc}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Low K</td>
<td>Medium K</td>
</tr>
<tr>
<td>Intake (g/d)</td>
<td>3.8</td>
</tr>
<tr>
<td>Faeces (g/d)</td>
<td>2.2&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Absorption</td>
<td>0.006</td>
</tr>
<tr>
<td>% intake</td>
<td>1.6&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

ND, not determined.
<sup>a,b,c</sup> Mean values within a row with unlike superscript letters were significantly different (Tukey's $t$ test; $P < 0.05$).
* For details of diets and procedures, see Tables 1 and 2 and p. 184.
† Values were ND because the wethers were fed a restricted amount of feed.

<table>
<thead>
<tr>
<th>Table 4. Rumen contents magnesium, potassium and sodium concentrations, and pH and transmural potential difference (PDt) in wethers fed the experimental rations*</th>
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<tr>
<td>(Mean values for four sheep per group)</td>
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<table>
<thead>
<tr>
<th>Experimental ration</th>
<th>Statistical significance of effect: $P_{disc}$</th>
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<tbody>
<tr>
<td>Mg (mM)</td>
<td></td>
</tr>
<tr>
<td>07.30 hours</td>
<td>8.7</td>
</tr>
<tr>
<td>Post-feeding†</td>
<td>9.6</td>
</tr>
<tr>
<td>K (mM)</td>
<td>42.9&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td>07.30 hours</td>
<td>43.0&lt;sup&gt;d&lt;/sup&gt;</td>
</tr>
<tr>
<td>Post-feeding†</td>
<td>100.4&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Na (mM)</td>
<td>94.7&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>07.30 hours</td>
<td>6.58</td>
</tr>
<tr>
<td>Post-feeding†</td>
<td>6.47</td>
</tr>
<tr>
<td>PDt (mV)&lt;sup&gt;‡&lt;/sup&gt;</td>
<td>48.2&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

<sup>a,b,c</sup> Mean values within a row with unlike superscript letters were significantly different (Tukey's $t$ test; $P < 0.05$).
* For details of diets and procedures, see Tables 1 and 2 and p. 184.
† Geometrical mean values of samples taken at 09.00, 11.00, 13.00, 15.00 and 17.00 hours.
‡ Geometrical mean values of PDt values measured from 09.00 to 15.00 hours.
Table 5. Rumen volume (RV) and passage rate (PR) of rumen contents in wethers fed the experimental rations* (Mean values for four sheep per group)

| Experimental ration | Low K | Medium K | High K | Extreme K | SEM | Statistical significance of effect: $P_{d\text{iet}}$
<table>
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<tr>
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</thead>
<tbody>
<tr>
<td>RV (litres)</td>
<td>4.0&lt;sup&gt;ab&lt;/sup&gt;</td>
<td>4.3&lt;sup&gt;b&lt;/sup&gt;</td>
<td>5.0&lt;sup&gt;ab&lt;/sup&gt;</td>
<td>6.7&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.33</td>
<td>0.008</td>
</tr>
<tr>
<td>PR %/h</td>
<td>4.6&lt;sup&gt;ab&lt;/sup&gt;</td>
<td>6.9&lt;sup&gt;b&lt;/sup&gt;</td>
<td>3.6&lt;sup&gt;a&lt;/sup&gt;</td>
<td>3.5&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.48</td>
<td>0.040</td>
</tr>
<tr>
<td>RV/m²/h</td>
<td>196</td>
<td>242</td>
<td>207</td>
<td>231</td>
<td>22.4</td>
<td>0.500</td>
</tr>
</tbody>
</table>

<sup>a,b</sup>Mean values within a row with unlike superscript letters were significantly different (Tukey's $t$ test; $P<0.05$).
* For details of diets and procedures, see Tables 1 and 2 and p. 184.

Discussion

The present study confirms earlier work showing that the addition of KHCO₃ to the ration of ruminant animals inhibits Mg absorption (Newton *et al.*, 1972; Poe *et al.*, 1985; Wylie *et al.*, 1985; Khorasani & Armstrong, 1990; Schonewille *et al.*, 1999a) and that the depressant effect of high K intake on Mg absorption is associated with an increase in the PDt (Schonewille *et al.*, 1999a). In the current feeding trial the linear relationship between the observed PDt and Mg absorption was calculated to be:

$$y = 65.27 - 0.52x \quad (R^2_{\text{adjusted}} = 0.239, P = 0.031, n = 16),$$

which supports the mechanism proposed on the basis of studies with isolated rumen models. Furthermore, Leonhard-Marek & Martens (1996) demonstrated that Mg absorption under *in vitro* conditions became independent from the lumen K concentration at a concentration $>80$ mmol/l. Indeed, when the PDt values measured after feeding the ration with $15.7$ g K/kg DM are omitted from the dataset, PDt becomes independent from the lumen K concentration at a concentration above values of approximately $125$ mmol/l.

In the present study, Mg absorption became independent from the dietary K concentration when it was increased from $37.6$ (medium K) to $59.4$ g/kg DM (high K), which was associated with an increase of the mean postprandial rumen K concentrations from $88$ to $127$ mmol/l (Table 4). Thus, on the basis of both *in vitro* observations (Leonhard-Marek & Martens, 1996) and the present *in vivo* PDt measurements, Mg absorption should not be further depressed at dietary K concentrations $>37.6$ g/kg DM (medium K) because the rumen K level then becomes $>80$ mmol/l (Table 4). However, it appeared that Mg absorption was significantly reduced (Table 3) when the dietary K concentration increased from $37.6$ (medium K) to $59.4$ g/kg DM (high K), which was associated with an increase of the mean postprandial rumen K concentrations from $88$ to $127$ mmol/l (Table 4). In the present study, Mg absorption became independent from the dietary K concentration when it was increased from $59.4$ (high K) to $77.4$ g/kg DM (extreme K). The associated increase in Mg absorption dropped non-significantly from $41.9$ to $35.4$ % intake when the dietary K concentration was increased from $15.7$ to $37.6$ g/kg DM (Table 3).

Table 6. Multiple regression model accounting for 83.3% of the observed variance in magnesium absorption*†

| Independent variable | Unit | $r$ ‡ | Statistical significance of effect: $P_{\text{Variable}}$
<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Constant</td>
<td>% Mg intake</td>
<td>66·3</td>
<td>6·90</td>
</tr>
<tr>
<td>Experimental period</td>
<td>Rank order</td>
<td>NG§</td>
<td>$&lt;0.001$</td>
</tr>
<tr>
<td>PDt</td>
<td>mV</td>
<td>$-0.61$</td>
<td>0.11</td>
</tr>
</tbody>
</table>

* For details of diets and procedures, see Table 1 and p. 184.
† The regression coefficient equals the change in average response of Mg absorption (% intake) if the independent variable increases by one unit.
‡ The regression coefficient was calculated by the lack of correlation between PDt and rumen K concentrations from the dataset, PDt becomes independent from the lumen K concentration at a concentration $>80$ mmol/l. Indeed, when the PDt values measured after feeding the ration with $15.7$ g K/kg DM are omitted from the dataset, PDt becomes independent from the mean postprandial K concentrations in rumen contents (Table 4). This independence was also illustrated by the lack of correlation between PDt and rumen K concentrations (Pearson’s $r = 0.310$, $P = 0.326$, $n = 12$). Furthermore, a significant linear relationship was calculated between the logarithmically transformed rumen K concentration and PDt, the regression formula being PDt (mV) $= 36.7 \log$ (rumen K) $- 10.7$ (Pearson’s $r = 0.760$, $P = 0.001$, $n = 16$). The slope of this regression equation corresponds to the slopes reported by Scott (1966), Ferreira *et al.* (1966) and Martens & Blume (1986), who reported values of $43, 24.4–39.7$ and $37.1$ respectively.

Thus, on the basis of both *in vitro* observations (Leonhard-Marek & Martens, 1996) and the present *in vivo* PDt measurements, Mg absorption should not be further depressed at dietary K concentrations $>37.6$ g/kg DM (medium K) because the rumen K level then becomes $>80$ mmol/l (Table 4). However, it appeared that Mg absorption was significantly reduced (Table 3) when the dietary K concentration increased from $37.6$ (medium K) to $59.4$ g/kg DM (high K), which was associated with an increase of the mean postprandial rumen K concentrations from $88$ to $127$ mmol/l (Table 4). In the present study, Mg absorption became independent from the dietary K concentration when it was increased from $59.4$ (high K) to $77.4$ g/kg DM (extreme K). The associated increase in Mg absorption dropped non-significantly from $41.9$ to $35.4$ % intake when the dietary K concentration was increased from $15.7$ to $37.6$ g/kg DM (Table 3).

In the current experiment Mg absorption dropped by $0.3$ percentage units when the dietary K concentration was increased by $1$ g/kg DM. This value agrees with the outcome of previous experiments using wethers fed similar amounts of KHCO₃ (Ram *et al.*, 1998; Schonewille *et al.*, 1999a). However, the observed decrease in Mg absorption of 6.5 percentage units in the current experiment was not statistically significant, which is probably related to the small number of animals used. In keeping with the work of McLean *et al.* (1985), the rumen Na concentration was linearly decreased when K intake was increased. Na absorption may be augmented through an increase in rumen K concentration as mediated by the difference in PDa (Martens *et al.*, 1998).

In addition to the electrogenic component of Mg uptake by rumen epithelial cells, the chemical gradient of soluble Mg between the lumen and cell content is also of importance. The soluble Mg concentration depends on the rumen conditions.
pH (Dalley et al. 1997). The Mg concentration in ultracentrifuged rumen contents can be affected by various factors including Mg intake, dietary Mg source, Mg solubility, and rumen volume and outflow (Van’t Klooster, 1967). The soluble Mg concentration depends on the rumen pH. At pH values >6.0, a sharp fall in the concentration of soluble Mg was observed by Dalley et al. (1997); this may relate to the formation of insoluble magnesium (ammonium) phosphates (Axford et al. 1982). In our present study, rumen pH was positively correlated with rumen K concentrations. Because mean postprandial rumen pH and mean postprandial rumen Mg concentrations were negatively associated, it could be suggested that an increase in K intake also reduces Mg absorption by lowering the concentration of Mg in the rumen contents. Furthermore, rumen volume was increased with increasing K intakes (Table 5); this may have interfered with Mg absorption. However, on the basis of stepwise regression the variance in rumen Mg concentrations did not significantly contribute to the explained variance in Mg absorption. Thus, it seems that the observed changes in either rumen pH or rumen volume did not significantly affect rumen Mg concentrations and Mg absorption in our wethers fed different concentrations of KHCO₃.

In conclusion, the present investigation indicates that raising the dietary K concentration from the low to high level (15.7 to 59.4 g K/kg DM) produced an increase in PDt and a marked decrease in apparent Mg absorption. However, apparent Mg absorption was not further depressed when dietary K concentration was increased above 59.4 g K/kg DM. This observation is important in relation to practical feeding of dairy cows where the amount of supplemental Mg is considered to be dependent on K intake. However, caution is warranted when quantitatively extrapolating the current findings to dairy cattle feeding. Contrary to what would be expected, urinary excretion of Mg in cows did not become marginally depressed when the dietary K content was increased from 48.0 to 75.5 g/kg DM (Jittakhot et al. 2003). However, in that study with cows, rumen K concentrations were 104 mm after the feeding of the ration containing 75.5 g K/kg DM, whereas in the current study with sheep fed a ration with identical K level, the rumen K concentration was as great as 150 mm.

Acknowledgements
We thank Jan van der Kuijlen for laboratory assistance and Anton Uijttewaal for biotechnical help. This study was supported by the Product Board Animal Feed (Productschap Diervoeder), The Hague, The Netherlands.

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