Effect of dietary supplementation or cessation of magnesium-based alkalizers on milk fat output in dairy cows under milk fat depression conditions

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ABSTRACT

We aimed to evaluate the effects of dietary supplementation with magnesium oxide and calcium-magnesium dolomite on milk fat synthesis and milk fatty acid profile or persistency in milk fat synthesis after their cessation in dairy cows under milk fat depression conditions. Twenty-four multiparous dairy cows in early lactation (mean $\pm$ standard deviation; 112 ± 14 d in milk) were used in a randomized complete block design. Milk fat depression was induced in all cows for 10 d by feeding a diet containing 35.2% starch, 28.7% neutral detergent fiber, and 4.8% total fatty acid (dry matter). The experiment was conducted in 2 periods. During the Mg-supplementation period (d 1–20), cows were randomly assigned to (1) the milk fat depression diet used during the induction phase (control; $n = 8$), (2) the control diet plus 0.4% magnesium oxide (MG; $n = 8$), or (3) the control diet plus 0.8% calcium-magnesium dolomite (CMC; $n = 8$). Compared with the control group, feeding the magnesium-supplemented diets increased milk fat concentration and yield by 12% within 4 d. During the 20-d Mg-supplementation period, both the MG and CMC diets increased milk fat concentration and yield, as well as 3.5% fat-corrected milk and energy-corrected milk yield, without affecting dry matter intake, milk yield, and milk protein and lactose concentrations. In the Mg-cessation period (d 21–30), all cows received the control diet, which resulted in a greater milk fat concentration and yield in the cows that had already received the MG and CMC diets in the Mg-supplementation period. Whereas, milk fat concentration and yield remained high after discontinuation of the magnesium-containing alkalizer until d 27. The difference in milk fat synthesis was associated with

lower *trans*-10 C18:1 (–22%) and higher *trans*-11 C18:1 (+12.5%) concentrations in milk during the Mg-supplementation period. Furthermore, it was evident that within 2 d of supplementation, the *trans*-10:*trans*-11 ratio was lower in MG and CMC cows compared with cows receiving the control. This suggested that the effect of magnesium-based alkalizers on milk fat synthesis was mediated via a shift in ruminal biohydrogenation of *cis*-9,*cis*-12 C18:2 in the rumen. In conclusion, abrupt addition of magnesium oxide and calcium-magnesium dolomite increased milk fat synthesis, which persisted for 7 d after cessation of magnesium-based alkalizers. A similar ability to recover milk fat synthesis and normal fatty acid biohydrogenation pathways was observed for magnesium oxide and calcium-magnesium dolomite.

Key words: magnesium, milk fat depression, *trans*-fatty acid, performance

INTRODUCTION

Increasing dietary NFC is a common approach to maximizing milk production in high-producing dairy cows. Typically, lactation diets for high-producing cows contain more than 40% NFC, mainly starch, which accounts for 70 to 80% of most cereal grains (NRC, 2001). Feeding highly fermentable carbohydrates results in increased concentration of VFA in the rumen and decreased rumen pH, which can affect the microbial composition of the rumen (Calsamiglia et al., 1999; Dijkstra et al., 2012). These changes in rumen pH and microbiota composition are associated with shifts in rumen biohydrogenation (BH) pathways (Fuentes et al., 2009; Sandri et al., 2020), leading to increased rumen synthesis of BH intermediates capable of inducing milk fat depression (MFD), such as *trans*-10,*cis*-12 CLA and *trans*-9,*cis*-11 CLA (Baumgard et al., 2000; Perfield et al., 2007).

Magnesium oxide is the most common supplemental source of magnesium for dairy cows; however, feeding magnesium oxide in excess of the magnesium requirement established by the NRC (2001) can increase ru-

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men pH and milk fat synthesis in dairy cows (Erdman et al., 1982; Bach et al., 2018). This additional magnesium oxide supplementation may increase feeding costs, whereas alternative magnesium sources may have similar effects while being less expensive.

In vivo studies have shown that regulation of rumen pH by the addition of buffer reduces the production of BH intermediates associated with MFD (Kalscheur et al., 1997; Cabrita et al., 2009). Similarly, different types of magnesium-based alkalizers have been proposed to stabilize rumen pH and thus reduce the occurrence of MFD (Neville et al., 2019; Razzaghi et al., 2020). Bach et al. (2018) reported that a magnesium-based product can neutralize rumen pH and maintain milk fat content more efficiently than sodium bicarbonate when cows are challenged with 3 kg/d of additional barley in the ration. In addition, feeding potassium carbonate sesquihydrate decreased *trans*-10 C18:1 content in milk [0.68 vs. 0.40% of total fatty acids (FA)] and increased milk fat synthesis (Harrison et al., 2012). Rauch et al. (2012) reported no difference in milk fat yield and feed efficiency between cows fed sodium bicarbonate or calcium-magnesium dolomite. Recently, we examined the diurnal rumen pH pattern by adding magnesium-based alkalizers when TMR formulated as potentially acidotic was fed to dairy cows and found that calcium-magnesium dolomite had a similar ability to magnesium oxide to shorten the time that rumen pH remained below 5.8 (from 12.5 h/d in control group to 9.1 and 8.7 h/d, respectively; Razzaghi et al., 2021). The acid-neutralizing ability of magnesium oxide probably reflects adequate solubility in rumen fluid (Le Ruyet and Tucker, 1992; Schaefer et al., 1982), which allows alkalizer to be effective at low pH when cows are fed high-starch diets. A similar mechanism may be at work with calcium-magnesium dolomite, which provides a novel alkalizer that achieves similar effects to commonly used minerals, such as magnesium oxide, but at a lower cost.

Although the beneficial effect of magnesium sources on rumen pH and milk fat content has been studied, the effect of supplementation or cessation of dietary magnesium sources on milk fat synthesis under MFD conditions is not known. In our previous study, we investigated the effect of magnesium sources (magnesium oxide or calcium-magnesium dolomite) on rumen fermentation and daily rumen pH patterns; however, that study was not designed to measure changes in milk fat synthesis due to experimental conditions (Razzaghi et al., 2021). To the best of our knowledge, no study has evaluated the use of calcium-magnesium dolomite and magnesium oxide as the main alkalizers for the recovery from MFD. We hypothesized that calcium-magnesium dolomite acts as a buffer for rumen pH that favors nor-

mal BH pathways and thus may promote recovery of milk fat synthesis. The main objective of the current study was to determine how rapidly magnesium oxide and calcium-magnesium dolomite supplementation may affect recovery of milk fat synthesis and ruminal BH intermediates in high-yielding dairy cows fed a milk fat-depressing diet. In addition, we aimed to evaluate the potential residual effects of magnesium-based supplements on milk fat synthesis after treatment cessation.

MATERIALS AND METHODS

Cows, Experimental Design, and Treatments

All experimental procedures and animal manipulations were approved by the Animal Care and Use Committee of Ferdowsi University of Mashhad (IACUC #A20175) as described in Iranian Council of Animal Care (1995). Twenty-four multiparous Holstein cows (660 ± 14 kg of BW, 112 ± 14 DIM and 45 ± 1.7 kg of milk/d; mean $\pm$ SD) were used in a randomized complete block design. Cows were blocked according to parity, pre-experimental milk yield, and DIM. Sample size and power analyses were used to calculate (Morris, 1999) the minimum number of replicates needed per treatment ($n = 8$) to detect a 10% level of observed mean differences for the primary outcome variables, including DMI and lactation performance, with a power of 0.80 and $\alpha = 0.05$, using 8 replicates per treatment. Estimates of variation for these variables were based on previously reported values (Yu et al., 1998; Alfonso-Avila et al., 2017; Malekkhahi et al., 2021).

All cows were housed in individual tiestalls in the Rayner Dairy Teaching and Research Facility (Ferdowsi University of Mashhad, Mashhad, Iran) with free access to water at all times. Animals were fed the normal herd ration during a 7-d pre-study period with forage-to-concentrate ratio of 40:60, formulated with 31.5% NDF, 16.7% CP, 31.6% starch (DM basis), and minerals and vitamins as recommended by the NRC (2001). Corn silage, alfalfa silage, and chopped alfalfa hay were used as forage components. Alfalfa hay was chopped to a theoretical length of 30 mm before feeding using a harvesting machine equipped with a sieve size controller (Golchin Trasher Hay Co.). The primary starch source during the pre-study period was dry ground corn.

Thereafter, during the 10-d MFD period, dairy cows were fed the milk fat-depressing diet with forage-to-concentrate ratio of 35:65 as a TMR, formulated with 35.5% starch, 29.0% NDF, and 4.8% total FA. The MFD diet contained 3.2% roasted soybeans (cracked) as a source of UFA and 38.5% of a processed corn product (DM basis) as a primary starch source (Table

1). Corn grain processing (i.e., super-conditioning process) was carried out by placing the ground corn in the super-conditioner (model 680 Dabbel, Asiab Machine Iranian) and injecting steam to increase the moisture content to 18 to 20% along with cooking at 95°C for 6 min as explained by Malekhhahi et al. (2021).

After the MFD-induction period (for 10 d), the cows began the experimental phase, which consisted of 2 periods, period 1 (Mg supplementation, d 1–20) and period 2 (Mg cessation, d 21–30). In the Mg-supplementation period, cows were randomly assigned to 1 of 3 treatments as follows: (1) MFD diet (**CTRL**), (2) the MFD diet + 0.4% DM magnesium oxide (including 67% magnesium oxide; **MG**), and (3) the MFD diet + 0.8% DM calcium-magnesium dolomite (a mixture of 53% calcium dolomite and 44% magnesium carbonate; **CMC**). Part of the wheat bran was replaced with 0.4% of diet DM as magnesium oxide, and calcium-magnesium dolomite replaced the limestone completely and wheat bran partially with 0.8% (diet DM basis). Magnesium oxide and calcium-magnesium dolomite doses were determined using the neutralization-capacity method proposed by Bach et al. (2018) and recently validated in dairy cows (Razzaghi et al., 2021). Accordingly, the magnesium oxide dose was set at half of the calcium-magnesium dolomite dose based on the expected neutralizing activity (the neutralizing capacity of magnesium oxide and calcium-magnesium dolomite was 36 and 20 mEq/g, respectively). Feeding rates of magnesium oxide and calcium-magnesium dolomite were consistent with recommended rates used in commercial diets (Rauch et al., 2012; Bach et al., 2018). In addition, the neutralizing activity of magnesium sources in 5% acetic acid was evaluated by measuring pH change according to Goff (2018; data not shown). Rations were formulated to be isoenergetic, isonitrogenous, and meet the nutrient requirements (NRC, 2001) of lactating dairy cows (weighing 660 kg, milk yield = 45 kg, fat concentration = 3.5%, true protein concentration = 3.0%, and intake = 27 kg of DM/d). During the Mg-cessation period, all cows were returned to the MFD diet for d 10. Cows were fed individually at 0800, 1600, and 2400 h to achieve a refusal rate of 10% and to allow ad libitum intake. Twice-weekly forage samples were collected, and DM content was determined to allow adjustment of diets to maintain forage-to-concentrate ratio. All cows were milked 3 times daily at 0700, 1500, and 2300 h.

Experimental Measurements and Sampling

Feed intake and milk yield and components were recorded daily throughout the experiment. Feed refusals were collected before the morning feeding and weighed daily. Samples of TMR were collected daily and pooled

at 5-d intervals, then weighed, oven-dried at 60°C for 48 h, and ground in a Wiley mill (standard model 4; Arthur H. Thomas Co.) to pass a 1-mm sieve. The procedures of AOAC International (2005) were used to measure DM by oven drying at 100°C for 24 h (method 934.01). Ash was determined by combustion at 600°C for 2 h (method 942.05; AOAC International, 2005), and ether extract content (method 920.39; AOAC International, 2005) was determined using a Soxhlet Gerhardt apparatus (model SE 416; Gerhardt). Crude protein (Kjeldahl N $\times$ 6.25) was determined by the block digestion method using a copper catalyst and steam distillation in boric acid (method 2001.11; AOAC International, 2005) with a 2100 Kjeltac distillation unit (Foss Inc.). Neutral detergent fiber and ADF were analyzed using the Fibertec System (1010 Heat Extractor) as described by Van Soest et al. (1991). The NDF and ADF values are reported inclusive of residual ash. Sodium sulfite and heat-stable α -amylase (Sigma A3306, Sigma-Aldrich) were used for NDF analysis. Nonfiber carbohydrate content (% DM) of the diets was calculated as $100 - (\text{NDF} + \text{CP} + \text{ether extract} + \text{ash})$ (NRC, 2001). Total starch was determined according to the procedure of Hall et al. (1999). The total content of FA in the diet was determined by GC using non-adeanoic acid (C19:1) as an internal standard (Sukhija and Palmquist, 1988). Subsamples of the TMR were subjected to digestion in HCO_3 (70%) + H_2O_2 (30%) using a digestion block for mineral measurements according to the procedure described by Mills and Jones (1996). Concentrations of calcium, magnesium, phosphorus, potassium, sodium, and sulfur in the diet were determined by inductively coupled plasma optical emission spectrometry (ICP-OES, Optima 4300DV, Perkin Elmer). Chloride in TMR was determined by silver nitrate titration after extraction with 0.5% nitric acid.

Milk was collected daily from all 3 daily milkings, and samples were pooled proportionately based on milk weights. Daily pooled milk samples were stored at 4°C with a preservative (potassium bichromate) until the further determination of milk fat, protein, and lactose using the Milko-Scan 605 analyzer (Foss Electric). Additional milk samples were collected on the first 5 consecutive days of the Mg-supplementation period without preservative and stored at -20°C until the end of the experiment for subsequent determination of the milk FA profile. Fat-corrected milk was estimated as $3.5\% \text{ FCM} = (0.434 \times \text{kg of milk}) + (16.216 \times \text{kg of milk fat})$ (Erdman, 2011). Energy-corrected milk was calculated as $\text{ECM} = (0.327 \times \text{kg of milk}) + (12.95 \times \text{kg of milk fat}) + (7.2 \times \text{kg of milk protein})$ (Orth, 1992). Feed efficiency was determined using $3.5\% \text{ FCM/DMI}$ (kg of 3.5% FCM per kg of DMI) and ECM/DMI (kg of ECM per kg of DMI).

Table 1. Ingredient and nutrient composition of the experimental total mixed ration¹

Item	Dietary treatment ²		
	CTRL	MG	CMC
Ingredient, % of DM			
Corn silage	26.0	26.0	26.0
Alfalfa silage	5.00	5.00	5.00
Alfalfa hay	4.00	4.00	4.00
Processed corn ³	38.5	38.5	38.5
Soybean meal, 44% CP	10.0	10.0	10.0
Canola meal	6.00	6.00	6.00
Fish meal	1.30	1.30	1.30
Meat meal	2.00	2.00	2.00
Roasted soybeans, cracked	3.20	3.20	3.20
Wheat bran	2.50	2.10	2.00
Vitamin and mineral premix ⁴	1.00	1.00	1.00
Limestone	0.30	0.30	—
Salt	0.20	0.20	0.20
Magnesium oxide ⁵	—	0.40	—
Calcium-magnesium dolomite ⁶	—	—	0.80
Chemical composition (% of DM, unless otherwise noted)			
NE _L , ⁷ Mcal/kg DM	1.62	1.62	1.62
DM	61.2	61.7	61.5
Ash	6.53	7.45	7.26
NDF	28.7	28.5	28.5
ADF	17.8	17.7	17.7
CP	16.6	16.7	16.6
Ether extract	5.06	5.10	5.01
Total fatty acids	4.80	4.85	4.83
Nonfiber carbohydrates	46.7	46.6	46.6
Starch	35.2	35.4	35.5
Minerals, % of DM			
Sodium	1.20	1.21	1.21
Potassium	1.21	1.20	1.22
Chloride	0.33	0.33	0.33
Sulfur	0.24	0.24	0.23
Calcium	0.73	0.74	0.76
Magnesium	0.23	0.42	0.32
Phosphorus	0.52	0.51	0.51
DCAD, ⁸ mEq/kg of DM	126	125	120
Fatty acids, g/100 g of FA			
C12:0	0.54	0.58	0.60
C14:0	0.60	0.64	0.63
C16:0	25.2	24.8	24.9
C16:1	0.44	0.48	0.45
C18:0	2.10	2.20	2.32
<i>cis</i> -9 C18:1	23.0	22.8	22.9
<i>cis</i> -11 C18:1	0.72	0.80	0.80
<i>cis</i> -9, <i>cis</i> -12 C18:2	37.0	37.2	37.1
<i>cis</i> -9, <i>cis</i> -12, <i>cis</i> -15 C18:3	10.4	10.5	10.3

¹Samples of the TMR were collected daily and pooled by 5-d intervals in the entire experiment.

²CTRL = control (milk fat-depressing diet; no magnesium-based alkalizers); MG = 0.4% DM of magnesium oxide; CMC = 0.8% DM of calcium-magnesium dolomite.

³Super-conditioned corn (Dordaneh Khorasan-e-Razavi Animal Feed Co.). The geometric mean and SD of the particle size in processed corn was 1.50 ± 0.23 mm.

⁴Each kilogram of the vitamin–mineral premix contained the following (DM basis): vitamin A (50,000 IU), vitamin D₃ (10,000 IU), vitamin E (4,000 IU), calcium (196 g), phosphorus (96 g), sodium (71 g), magnesium (19 g), iron (3 g), copper (0.3 g), manganese (2 g), zinc (3 g), cobalt (0.1 g), iodine (0.1 g), and selenium (0.001 g).

⁵Contained 48% magnesium, 1.7% sodium, and 2.9% calcium.

⁶Contained 15% magnesium, 0.8% sodium, and 22.5% calcium.

⁷According to NRC (2001).

⁸According to NRC (2001).

For the analysis of milk FA, milk fat was extracted using the centrifugation technique described by Luna et al. (2005). The refrigerated raw milk sample was kept

at 20°C for 20 min and centrifuged at $17,800 \times g$ for 30 min at 4°C. The fat layer was transferred to a micro-tube and centrifuged at $19,300 \times g$ for 20 min at room

temperature. After the second centrifugation, the top layer was kept for FA analysis of milk fat. Subsequently, the extracted lipids were subjected to base-catalyzed transmethylation with sodium methoxide to give FAME following the procedure described by Christie (1982) with modifications by Chouinard et al. (1999). Fatty acid methyl esters were then recovered in hexane and separated in a gas chromatograph (3400 Varian Star instrument, Varian Inc.) equipped with a CP-SIL-88 capillary column (Chrompack, 60 m $\times$ 0.25 mm, Varian) and a flame ionization detector with helium as the carrier gas. The column temperature was initially set at 50°C for 1 min and increased by 10°C/min to 190°C for another 130 min. The injector temperature was 280°C, and that of the detector was set at 300°C. The FAME peaks were identified by comparing their retention time with that of a standard mixture of 37 components FAME (Sigma-Aldrich, Supelco 18919-1AMP) and 60 individual FAME standards (Sigma-Aldrich).

Statistical Analysis

Before statistical analysis, the normality of data distribution was tested by the Shapiro-Wilk test using the UNIVARIATE procedure and evaluated visually by plotting residuals (PROC PLOT). Of the tested parameters, feed intake and milk yield and components were not normally distributed. Nonnormally distributed values were scaled using a $\log_{10}$ transformation to correct for heteroscedasticity and meet assumptions of normality (reported in Supplemental Table S1; <https://data.mendeley.com/datasets/ngszd7jrtz/1>; Razzaghi, 2021). After the log-transformation, the distribution of the data was tested again, and the data were normally distributed. Least squares means were back-transformed to original units for interpretation of tables and figures (Table 2 and Figure 1). Data were analyzed as a randomized complete block design with the REPEATED statement in the MIXED procedure of SAS (version 9.4, SAS Institute Inc.) using the following model:

$$Y_{ijkl} = \mu + T_i + D_j + (T \times D)_{ij} + A_{(k)l} + B_l + \varepsilon_{ijkl},$$

where Y_{ijkl} is the variable observed, μ is the overall mean, T_i is the fixed effect of treatment ($i = 1-3$), D_j refers to fixed effect of time (d), $(T \times D)_{ij}$ is the fixed effect of the interaction, $A_{(k)l}$ is the random effect of cow (k) within block (l), B_l is the random effect of block, and ε_{ijkl} denotes the residual error. Feed intake and milk production or component values were analyzed for 3 discrete periods as Mg supplementation (d 1–20), Mg cessation (d 21–30), and overall (d 1–30). However, we reported P -values for the entire period, although

data on milk FA profile were analyzed for Mg supplementation period (d 1–5). Three variance-covariance structures (autoregressive type 1, compound symmetry, and Toeplitz) were tested, and the first-order autoregressive covariance structure was determined as the most appropriate covariance structure for all repeated statements according to the Akaike and Bayesian information criteria. Main effects were declared significant at $P \leq 0.05$, and tendencies were declared at $0.05 < P \leq 0.10$. Interactions were declared significant at $P \leq 0.10$, and tendencies were declared at $0.10 < P \leq 0.15$. The PDIFF option was used with Tukey adjustment to test for significance of multiple comparisons. The SLICE option in the LSMEANS statement was used to compare the treatments at a specific day when treatment by time interaction was significant.

RESULTS

Dry matter intake, milk yield, milk components, and milk efficiency during periods of Mg supplementation, Mg cessation, and overall are shown in Table 2, and transformed data are reported in Supplemental Table S1. The addition of magnesium oxide or calcium-magnesium dolomite did not affect DMI and milk yield, and there was no interaction between treatment and time for these variables throughout the period. The MG and CMC diets resulted in a greater ($P = 0.01$) milk fat concentration and yield, whereas the interaction between treatment and time was significant ($P < 0.01$) for milk fat concentration, but only a trend for milk fat yield ($P = 0.11$). Such an increase in milk fat concentration was observed within the first 4 d of the Mg-supplementation period (Figure 1, panel A). It remained high throughout the rest of the Mg-supplementation period and remained elevated after discontinuation of the magnesium-containing alkalizers until d 27. In addition, a similar pattern was observed for milk fat yield (Figure 1, panel B).

Treatment did not affect milk protein and lactose concentrations or yields throughout the period. The MG and CMC groups had greater yields of 3.5% FCM ($P = 0.03$) and ECM ($P = 0.04$) throughout the period compared with CTRL, but we observed no interaction between treatment and time for these variables. Feed efficiency (kg of FCM/kg of DMI) was greater ($P < 0.01$) in cows fed MG and CMC than in cows fed CTRL for the entire period. In addition, ECM efficiency (kg of ECM/kg of DMI) was lower ($P < 0.01$) for CTRL than for MG group.

The effects of treatments on the profile of milk FA during the period of Mg supplementation are summarized in Table 3. Treatment differences were observed

Table 2. Feed intake and milk production and composition in cows fed a milk fat-depressing (MFD) diet supplemented or not with magnesium-based alkalisers¹

Item	Dietary treatment ²			P-value		
	CTRL	MG	CMC	SEM	Treatment	Time
DML, kg/d						
Mg supplementation (d 1 to 20)	26.4 (25.3–27.5)	26.7 (25.6–27.9)	27.0 (25.7–28.0)	0.44		
Mg cessation (d 21 to 30)	26.0 (24.9–27.4)	26.3 (25.3–27.9)	26.8 (25.5–28.0)	0.90		
Overall (d 1 to 30)	26.3 (25.4–27.3)	26.6 (25.8–27.6)	26.8 (25.9–27.7)	0.45	0.41	0.03
Milk yield, kg/d						
Mg supplementation (d 1 to 20)	44.8 (43.0–46.5)	45.0 (43.2–46.8)	45.2 (43.4–47.0)	0.50		
Mg cessation (d 21 to 30)	44.5 (43.8–45.6)	44.5 (43.8–45.6)	44.8 (44.0–45.9)	1.26		
Overall (d 1 to 30)	44.7 (43.6–46.0)	45.0 (43.8–46.1)	45.1 (43.9–46.2)	0.56	0.67	0.53
ECM ³ , kg/d						
Mg supplementation (d 1 to 20)	40.9 (38.7–43.2)	43.4 (41.2–45.6)	43.9 (41.6–46.0)	0.61		
Mg cessation (d 21 to 30)	40.1 (39.2–42.6)	42.1 (40.4–43.8)	43.0 (41.3–44.7)	0.96		
Overall (d 1 to 30)	40.5 (39.4–42.4)	42.6 (41.4–44.5)	43.5 (42.0–45.1)	0.76	0.04	<0.01
3.5% FCM ⁴ , kg/d						
Mg supplementation (d 1 to 20)	39.8 (37.3–42.4)	42.9 (40.3–45.4)	43.4 (40.8–46.0)	0.68		
Mg cessation (d 21 to 30)	39.9 (37.8–42.0)	41.4 (39.2–43.6)	42.4 (40.3–44.5)	1.18		
Overall (d 1 to 30)	39.7 (38.0–41.6)	42.5 (40.6–44.2)	42.8 (41.3–44.9)	0.88	0.03	<0.01
Milk fat, %						
Mg supplementation (d 1 to 20)	2.72 (2.43–3.11)	3.16 (2.81–3.50)	3.22 (2.90–3.57)	0.053		
Mg cessation (d 21 to 30)	2.83 (2.45–3.20)	3.02 (2.61–3.44)	3.12 (2.74–3.50)	0.098		
Overall (d 1 to 30)	2.78 (2.48–3.08)	3.11 (2.81–3.41)	3.20 (2.90–3.50)	0.150	0.01	<0.01
Milk protein, %						
Mg supplementation (d 1 to 20)	3.08 (2.94–3.21)	3.09 (2.95–3.22)	3.07 (2.94–3.21)	0.022		
Mg cessation (d 21 to 30)	3.09 (2.94–3.24)	3.07 (2.91–3.23)	3.07 (2.94–3.21)	0.029		
Overall (d 1 to 30)	3.08 (2.97–3.20)	3.09 (2.97–3.19)	3.07 (2.96–3.18)	0.054	0.48	<0.01
Milk lactose, %						
Mg supplementation (d 1 to 20)	4.61 (4.51–4.70)	4.58 (4.48–4.68)	4.61 (4.51–4.71)	0.029		
Mg cessation (d 21 to 30)	4.61 (4.49–4.73)	4.60 (4.48–4.72)	4.59 (4.47–4.71)	0.052		
Overall (d 1 to 30)	4.61 (4.53–4.69)	4.59 (4.51–4.67)	4.60 (4.52–4.68)	0.032	0.18	<0.01
Milk fat, kg/d						
Mg supplementation (d 1 to 20)	1.26 (1.05–1.47)	1.44 (1.23–1.64)	1.48 (1.25–1.68)	0.031		
Mg cessation (d 21 to 30)	1.27 (1.07–1.46)	1.35 (1.14–1.57)	1.41 (1.21–1.60)	0.042		
Overall (d 1 to 30)	1.26 (1.09–1.42)	1.41 (1.24–1.58)	1.45 (1.28–1.62)	0.082	0.01	<0.01
Milk protein, kg/d						
Mg supplementation (d 1 to 20)	1.38 (1.26–1.51)	1.40 (1.27–1.52)	1.39 (1.27–1.51)	0.016		
Mg cessation (d 21 to 30)	1.37 (1.29–1.47)	1.37 (1.28–1.46)	1.38 (1.29–1.47)	0.015		
Overall (d 1 to 30)	1.38 (1.30–1.47)	1.39 (1.30–1.48)	1.39 (1.31–1.48)	0.041	0.89	0.01
Milk lactose, kg/d						
Mg supplementation (d 1 to 20)	2.07 (1.93–2.21)	2.07 (1.93–2.21)	2.10 (1.95–2.22)	0.024		
Mg cessation (d 21 to 30)	2.06 (1.97–2.15)	2.06 (1.97–2.14)	2.07 (1.98–2.15)	0.020		
Overall (d 1 to 30)	2.07 (1.98–2.16)	2.07 (1.97–2.16)	2.08 (1.99–2.17)	0.046	0.74	0.03
3.5% FCM/DMI						
Mg supplementation (d 1 to 20)	1.51 (1.47–1.54)	1.60 (1.56–1.63)	1.61 (1.58–1.64)	0.031		
Mg cessation (d 21 to 30)	1.52 (1.49–1.55)	1.56 (1.53–1.58)	1.59 (1.56–1.62)	0.061		
Overall (d 1 to 30)	1.51 (1.49–1.54)	1.58 (1.56–1.61)	1.61 (1.58–1.63)	0.012	<0.01	<0.01
ECM/DMI						
Mg supplementation (d 1 to 20)	1.55 (1.52–1.58)	1.62 (1.59–1.65)	1.63 (1.60–1.66)	0.030		
Mg cessation (d 21 to 30)	1.56 (1.53–1.59)	1.58 (1.55–1.61)	1.61 (1.59–1.64)	0.056		
Overall (d 1 to 30)	1.55 (1.53–1.58)	1.60 (1.58–1.63)	1.63 (1.60–1.65)	0.011	<0.01	<0.01

¹In the Mg-supplementation period, cows were fed with control diet (n = 8), whereas 8 cows remained on the control diet plus magnesium oxide and 8 cows were fed with the control diet plus calcium-magnesium dolomite for 20 d. In the Mg-cessation period, all cows were again fed the control diet for 10 d. Values reported as least squares means (95% CI are given in parentheses) and pooled SEM. All data were transformed using $\log_{10}$ transformation for statistical analysis and back-transformed for interpretation.

²CTRL = control (MFD diet with no magnesium-based alkalisers; MG = 0.4% DM of magnesium oxide; CMC = 0.8% DM of calcium-magnesium dolomite. The MG and CMC diets were fed only during the Mg-supplementation period.

³ECM yield = (kg of milk $\times$ 0.3246) + (kg of milk protein $\times$ 7.04), as described by Orth (1992).

⁴3.5% FCM yield = (0.432 $\times$ kg of average milk yield) + (16.216 $\times$ kg of fat); Erdman (2011).

for the concentrations of C6:0 (0.97 vs. 1.32, CTRL vs. MG and CMC; $P = 0.02$), C18:0 (11.5 vs. 12.4, CTRL and MG vs. CMC; $P = 0.02$), *trans*-10 C18:1 (1.49 vs. 1.16, CTRL vs. MG and CMC; $P < 0.01$), and *trans*-11 C18:1 (1.54 vs. 1.78, CTRL vs. MG; $P = 0.01$) in milk fat. Although the concentration of C16:0 tended to increase ($P = 0.10$) in cows fed the MG diet (30.1 vs. 31.2 g/100 g of FA, CTRL vs. MG), the concentration of *trans*-9 C18:1 tended to decrease ($P = 0.06$) with feeding of the CMC diet (0.20 vs. 0.17 g/100 g of FA, CTRL vs. CMC). The proportion of C6:0 in milk fat increased (+26%) in cows fed magnesium-based supplements compared with CTRL, whereas the proportion of C18:0 increased (+7%) in cows fed the CMC diet compared with all other diets. Furthermore, *trans*-10 and *trans*-11 C18:1 concentrations in milk fat decreased (−22%) and increased (+12.5%), respectively, in cows receiving magnesium-based supplements compared with the CTRL group. In addition, dietary supplementation with magnesium-based alkalizers did not alter *cis*-9,*trans*-11 CLA milk fat concentrations. Magnesium supplementation reduced total *trans*-FA by 10.5% in cows fed the MG and CMC diets compared with those fed the CTRL diet. However, the proportions of milk de novo ($P = 0.10$) and mixed origin FA ($P = 0.06$) tended to increase in cows fed the MG diet compared with those fed the CTRL and CMC diets, but the concentration of preformed FA in milk was not affected by the treatments. The ratio of *trans*-10:*trans*-11 was also greater ($P < 0.01$) in CTRL compared with those fed MG and CMC diets.

An effect of time on milk FA profile was observed for C12:0, C14:0, *trans*-C14:1, *cis*-1C4:1, C18:0, and *trans*-C18:1 isomers ($P < 0.05$). Significant treatment $\times$ time interactions were observed for C18:0 and total *trans*-FA ($P < 0.05$) and the ratio of *trans*-10:*trans*-11 ($P < 0.01$). It was evident that within 48 h of supplementation, the ratio of *trans*-10:*trans*-11 was lower in cows fed MG and CMC compared with cows fed CTRL (Figure 2), which persisted in the remaining period of Mg supplementation (until d 5).

DISCUSSION

Diets, Performance, and Milk Fat Synthesis

In the present experiment, supplementation of magnesium in MFD-inducing diets attenuated the decrease in milk fat, an effect that gradually disappeared after cessation of magnesium supplementation. These changes were associated with improved BH characteristics and not with changes in DMI or milk yield. In fact, the addition of 0.4% magnesium oxide and 0.8% calcium-magnesium dolomite to the diet did not affect feed in-

take in our study. Other studies in lactating dairy cows showed that buffering or alkalizing agents had no effect on lactation performance, either in typical (Cabrita et al., 2009; Rauch et al., 2012) or high-concentrate diets (Kalscheur et al., 1997; Khorasani and Kennelly, 2001); however, other studies (Neville et al., 2019; Razzaghi et al., 2020) reported increased feed intake. Rauch et al. (2012) reported that there was no difference in feed intake and milk yield between control and calcium-magnesium dolomite diets on commercial dairy farms, but differences in dietary starch and NDF contents compared with our study were evident (starch = 15.7 vs. 35.2% and NDF = 33.9 vs. 28.7%, on DM basis). In addition, there are divergent results regarding the effects of feeding buffers on milk yield. Some studies have documented no effects (Bach et al., 2018; Razzaghi et al., 2020), whereas others have reported increased milk yield (Razzaghi et al., 2021) with the addition of buffering or alkalizing agents.

In our study, MFD was intentionally induced using 38.5% DM of processed corn and 3.2% DM of cracked roasted soybeans in the experimental diets. The expectation was that greater ruminal starch degradability would lower rumen pH, and this could account for any observed differences in the pattern of BH intermediates (Mohammed et al., 2010; Dewanckele et al., 2019). In addition, UFA was supplied in all diets via a slowly available FA from cracked roasted soybeans. The experimental diets contained an average of 28.6% NDF, 35.3% starch, and 4.8% total FA to induce MFD. Previous time course work has shown that changes in BH occur rapidly and a maximum MFD was observed within 10 to 14 d (Rico and Harvatine, 2013); therefore, the onset of MFD was expected within the duration of the MFD-induction period. In our study, the average milk fat concentrations at the beginning and end of the MFD-induction period were 3.82 and 2.75%, and milk fat yields were 1.74 and 1.22 kg/d, respectively. This reflected a 28 and 30% reduction in milk fat concentration and yield, respectively, during the MFD-induction period. Similarly, Razzaghi et al. (2021) reported a greater milk fat concentration (+11%) and yield (+21%) when similar doses of calcium-magnesium dolomite and magnesium oxide (0.8 and 0.4% of the diet DM) were used compared with a nonsupplemented diet when cows were fed a 34.2% starch diet.

The objective of this experiment was to investigate the role of magnesium oxide and calcium-magnesium dolomite in relation to changes in milk fat synthesis after supplementation or cessation of dietary magnesium-based alkalizers. In addition, the experimental diet was formulated to increase the risk of MFD, and thus CTRL cows had greater *trans*-10 C18:1 concentration in milk (1.49% of FA), similar to a recent meta-analysis

Figure 1. Milk fat concentration (%; panel A) and milk fat yield (kg/d; panel B) during the Mg-supplementation period (d 1–20) for cows fed either a milk fat-depressing (MFD) diet alone (CTRL; ● and solid line), the MFD diet + 0.4% magnesium oxide (MG; ▲ and solid line), or the MFD diet + 0.8% calcium-magnesium dolomite (CMC; ■ and dotted line) and during the Mg-cessation period (d 21–30) when all cows received the MFD diet. Treatment × time interaction was $P < 0.01$ for milk fat concentration and $P = 0.11$ for milk fat yield. Different letters (a,b) indicate the significant difference ($P < 0.05$) between treatments within a time point. Milk fat values are back-transformed LSM, and error bars represent SEM.

of more than 500 treatment means (mean = 1.39% of FA; Matamoros et al., 2020). Moreover, milk fat yield of cows fed MG or CMC was greater after withdrawal of magnesium oxide and calcium-magnesium dolomite

supplements than in the CTRL group (Figure 1, panel B). This suggested that the effects of the magnesium supplements on milk fat synthesis may have an effect on fat metabolism via altered BH pathways (i.e., *trans*-11

C18:1 and *trans*-10 C18:1 shift). The timing (within 96 h) of the increase in milk fat concentration induced by the addition of magnesium-based alkalizers is consistent with a previous report (Rico and Harvatine, 2013) of the timing of changes in milk fat concentration during the induction and recovery of diet-induced MFD. However, in the Mg cessation period, the positive effect of magnesium sources on milk fat synthesis disappeared by 7 d after cessation of magnesium supplementation until the end of the experiment. During the period of Mg supplementation, milk fat concentration and yield

increased over time in both MG and CMC diets, but the recovery rate was slower than that of the potassium source in the study of Ma et al. (2017). They reported that recovery of MFD in dairy cows occurred within 2 d after supplementation of an MFD diet (contained 1.8% soybean oil and 22% starch, DM-based) with 0.59% potassium carbonate sesquihydrate.

The magnesium-supplemented diets resulted in greater yields of ECM and 3.5% FCM compared with the CTRL group, which is in agreement with the results of Razzaghi et al. (2021) and Khorasani and Kennelly

Table 3. Milk fatty acid profile in the first consecutive 5 d of Mg-supplementation period for cows fed a milk fat-depressing (MFD) diet supplemented or not with magnesium-based alkalizers¹

Fatty acids (g/100 g of FA)	Dietary treatment ²			SEM	P-value		
	CTRL	MG	CMC		Treatment	Time	Treatment × time
C4:0	1.41	1.54	1.65	0.083	0.16	0.02	0.17
C6:0	0.97 ^b	1.36 ^a	1.28 ^a	0.091	0.02	0.95	0.16
C8:0	1.05	1.02	1.03	0.045	0.89	0.92	0.69
C10:0	2.20	2.28	2.18	0.119	0.83	0.63	0.89
C10:1	0.30	0.34	0.29	0.016	0.18	0.21	0.50
C12:0	2.86	2.78	2.92	0.102	0.65	<0.01	0.73
C14:0	11.0	11.3	10.8	0.31	0.60	<0.01	0.63
<i>cis</i> -9 C14:1	0.88	0.94	0.92	0.014	0.13	<0.01	0.11
<i>trans</i> -9 C14:1	0.35	0.34	0.38	0.021	0.24	<0.01	0.27
C15:0	0.85	0.83	0.86	0.032	0.96	0.22	0.33
C15:1	0.09	0.08	0.07	0.010	0.51	0.41	0.18
C16:0	30.1 ^y	31.2 ^x	30.6 ^{xy}	0.33	0.10	0.24	0.97
C16:1	1.76	1.81	1.66	0.144	0.76	0.78	0.11
C17:0	0.53	0.49	0.41	0.039	0.14	0.11	0.24
C18:0	11.3 ^b	11.8 ^b	12.4 ^a	0.27	0.02	<0.01	0.02
<i>trans</i> -8 C18:1	0.25	0.22	0.23	0.016	0.12	<0.01	0.70
<i>trans</i> -9 C18:1	0.20 ^x	0.18 ^{xy}	0.17 ^y	0.011	0.06	<0.01	0.11
<i>trans</i> -10 C18:1	1.49 ^a	1.19 ^b	1.13 ^b	0.053	<0.01	0.05	0.48
<i>trans</i> -11 C18:1	1.54 ^b	1.78 ^a	1.74 ^a	0.076	0.01	0.02	0.83
<i>cis</i> -9 C18:1	23.1	22.6	22.2	0.34	0.52	0.85	0.27
<i>cis</i> -9, <i>cis</i> -12 C18:2	2.50	2.45	2.52	0.123	0.92	0.58	0.95
<i>cis</i> -9, <i>trans</i> -11 C18:2	0.79	0.81	0.86	0.044	0.57	0.34	0.99
<i>trans</i> -10, <i>cis</i> -12 C18:3	ND	ND	ND	ND	—	—	—
<i>cis</i> -9, <i>cis</i> -12, <i>cis</i> -15 C18:3	0.61	0.57	0.65	0.036	0.36	0.69	0.67
C20:0	0.73	0.73	0.66	0.034	0.12	0.58	0.52
<i>cis</i> -9 C20:1	0.15	0.16	0.18	0.010	0.19	0.31	0.13
C22:0	0.13	0.18	0.15	0.011	0.22	0.17	0.82
<i>cis</i> -9 C22:1	0.13	0.15	0.13	0.008	0.55	0.46	0.15
C24:0	0.71	0.76	0.85	0.033	0.32	0.25	0.15
Sums							
De novo fatty acids ⁴	21.1 ^y	21.9 ^x	21.5 ^{xy}	0.28	0.10	<0.01	0.52
Mixed fatty acids ⁵	31.9 ^y	33.1 ^x	32.2 ^{xy}	0.32	0.06	0.28	0.87
Preformed fatty acids ⁶	46.8	45.0	46.3	0.95	0.25	0.05	0.51
Total <i>trans</i> -FA ⁷	3.91 ^a	3.59 ^b	3.42 ^b	0.107	<0.01	0.04	0.03
<i>trans</i> -10: <i>trans</i> -11 18:1 ratio	0.98 ^a	0.69 ^b	0.66 ^b	0.030	<0.01	<0.01	<0.01

^{a,b}Means with different superscripts within a row differ ($P < 0.05$).

^{x,y}Means with different superscripts within a row differ ($P \leq 0.10$).

¹In the Mg-supplementation period, cows were fed with control diet ($n = 8$), whereas 8 cows remained on the control diet plus magnesium oxide and 8 cows were fed with the control diet plus calcium-magnesium dolomite for 20 d.

²CTRL = control (milk fat depression diet with no magnesium-based alkalizers); MG = 0.4% DM of magnesium oxide; CMC = 0.8% DM of calcium-magnesium dolomite.

³*Trans*-10,*cis*-12 CLA was not detected in milk fat (ND).

⁴Sum of straight even-chain fatty acids from 4 to 14-carbon.

⁵Sum of C16 and C16:1.

⁶Sum of odd-chain fatty acids (C15:0), and all fatty acids with a chain length of 17-carbon and greater.

⁷Sum of *trans*-9 C14:1, *trans*-8 C18:1, *trans*-9 C18:1, *trans*-10 C18:1, *trans*-11 C18:1.

(2001) that show higher milk fat yield. In addition, there was a difference in feed efficiency between magnesium-supplemented diets and CTRL. Although MFD diets may provide more energy for adipose tissue instead of milk fat synthesis (Harvatine et al., 2009b), it appears that in our study, the improved feed efficiency may be a result of greater ECM yield and, ultimately, less energy being deposited in adipose tissue with similar feed intake. The specific mechanism of how magnesium-based alkalizer supplementation leads to increased milk fat synthesis could be attributed to enhanced triglyceride uptake by the mammary gland (Thomas and Emery, 1969) or changes in ruminal fermentation pattern (Erdman et al., 1982; Razzaghi et al., 2021). In the current study, increased milk fat synthesis may be related to the decreased milk concentrations of *trans*-10 C18:1 reported here, the shorter time in which rumen pH remains below a threshold (i.e., 5.8), and the greater acetate-to-propionate ratio as reported by Razzaghi et al. (2021). It is important to note that both the MG and CMC diets had lower concentrations of *trans*-10 C18:1 than the CTRL group (−20 and −24% in MG and CMC, respectively), as both supplements could decrease the alternative BH pathway during MFD.

In one of the previous studies in which calcium-magnesium dolomite was added to lactating dairy cow diet (0.76% of diet DM), Rauch et al. (2012) reported that milk fat concentration was similar to the control treatment. Based on a few experiments, it appears

that the addition of alkalizer to diets high in starch and corn silage in lactating cows may increase milk fat concentration due to the greater rumen solubility of alkalizer at lower rumen pH. Recently, Razzaghi et al. (2020) reported that a commercial buffer blend containing cation sources of calcium and magnesium increased milk fat concentration and yield in cows fed high-concentrate diets. The acid-neutralizing capacity of alkalizers is associated with some physical and chemical properties that explain the different rates of solubilization in rumen fluid (Le Ruyet and Tucker, 1992). In the present study, the neutralization capacity of magnesium oxide and calcium-magnesium dolomite were 36 and 20 mEq H⁺/g, respectively, which is comparable to other studies (Crawford et al., 2008; Bach et al., 2018). Schaefer et al. (1982) suggested that the acid-neutralizing capacity of magnesium oxide probably depends on its solubility in rumen fluid. They observed that alkalizers were effective at low pH when cows were fed starch-rich diets. The results of an in vitro study with the buffer and alkalizer showed that the buffering capacity of the rumen fluid increased for magnesium oxide and stabilized over 24 h, whereas this duration was 0 to 12 h for sodium bicarbonate, which can be attributed to a gradual dissolution of the alkalizer (Le Ruyet and Tucker, 1992). Fuentes et al. (2009) showed that the content of intermediates from incomplete BH in the rumen (*trans*-10 C18:1 and *trans*-10, *cis*-12 CLA) increased dramatically when the rumen pH decreased

Figure 2. *Trans*-10:*trans*-11 ratio in milk fat during the first 5 d of the Mg-supplementation period for cows fed either a milk fat-depressing (MFD) diet alone (CTRL; ● and solid line), the MFD diet + 0.4% magnesium oxide (MG; ▲ and solid line), or the MFD diet + 0.8% calcium-magnesium dolomite (CMC; ■ and dotted line). Treatment × time interaction was $P < 0.01$. Different letters (a,b) indicate the significant difference ($P < 0.05$) between treatments within a time point. Error bars represent SEM.

from 6.4 to 5.6. Similarly, the decreased rumen pH in the CTRL group in the present study may have inhibited the activity of some rumen bacterial species believed to be critical for rumen BH (Jenkins et al., 2008) and consequently for the regulation of milk fat synthesis.

Changes in Milk FA Profile

Magnesium oxide and calcium-magnesium dolomite supplementation increased *trans*-11 C18:1 in milk fat by 12.5% compared with CTRL group, supporting the hypothesis that feeding magnesium-based alkalizers promote the normal (*trans*-11 C18:1) BH pathway (Bauman and Griinari, 2001; Rico and Harvatine, 2013) rather than the altered pathway (*trans*-10 C18:1). In the current study, it appeared that magnesium-based alkalizers promoted lower shifts in BH, probably by controlling rumen pH and stimulating fiber-digesting rumen bacteria associated with normal BH (Jenkins et al., 2008). Among factors promoting the alternate pathway, a starch-rich diet may alter the rumen environment and promote a shift in BH pathways in favor of *trans*-10 C18:1 FA, regardless of dietary PUFA content (Rico et al., 2015; Sandri et al., 2020), in line with our results.

The concentration of *cis*-9,*cis*-12 C18:2 in milk fat was not affected by the treatments, suggesting that magnesium-based supplements do not affect the initial rate of isomerization, but rather shift BH pathways, as suggested by the increased concentrations of *trans*-10 C18:1. *Trans*-10,*cis*-12 CLA is one of the most potent inhibitors of milk fat synthesis (Shingfield and Griinari, 2007), but was not detected in milk fat in the present study. Importantly, this does not necessarily mean that *trans*-10,*cis*-12 CLA was not synthesized in the rumen, but only that these levels were not sufficient to be above the detection limit under our conditions. Furthermore, the levels of *trans*-10 C18:1 (average 1.27% of FA) were very low compared with other studies, where they can reach levels of up to 4.0% of FA under MFD conditions (Rico and Harvatine, 2013), suggesting that less substrate was available for BH via this pathway. This is likely the result of the lower dietary levels of PUFA in our experiment (sum of C18:2 and C18:3 FA in the diet; 2.2% of DM) compared with the study of Rico and Harvatine (2013; 3.8% of diet FA). In addition, others have reported a transient reduction in milk fat synthesis in the absence of *trans*-10,*cis*-12 CLA in milk fat when fed high-concentration diets low in dietary *cis*-9,*cis*-12 C18:2 (Sandri et al., 2020). It cannot be ruled out that other mechanisms were at play, including the effect of unidentified isomers capable of MFD (Perfield et al., 2007).

Typically, concentration of de novo FA (i.e., from mammary origin) decrease during MFD, whereas concentration of preformed FA (i.e., from the circulation) increase (Ramirez Ramirez et al., 2015; Rico et al., 2015). The relationship between *trans*-10,*cis*-12 CLA or *trans*-10 C18:1 in milk and de novo FA concentration in milk is negative, whereas the relationship with the proportion of preformed FA is positive (Bauman and Griinari, 2001). In our study, a trend for increased concentration of de novo and mixed FA (16-carbon; half derived from de novo synthesis in the mammary gland and the other half from the circulation) was observed when MG was added to the diets compared with the CTRL group. Increased rumen accumulation of *trans*-10 isomers was reported in some experiments when feeding a high fat and starch diet to dry and lactating cows (Zened et al., 2013; Ramirez Ramirez et al., 2015). The increase in *trans*-11 C18:1 during Mg-supplementation period in the magnesium-fed animals, and also the increased de novo FA in the MG diet, suggested that the rumen shifts back to the normal BH pathway, whereas a greater concentration of *trans*-10 C18:1, a key marker of the altered BH pathway, was observed in the CTRL group. In addition, supplementation of magnesium-based alkalizers resulted in a lower *trans*-10:*trans*-11 ratio within 48 h of supplementation, which remained low during the rest of the Mg-supplementation period (the first 5 d of this period). Alterations in *trans*-10 C18:1 are commonly associated with cases of MFD and altered BH in the rumen, with *trans*-10 C18:1 acting as an intermediate in the alternative BH pathway (Lock et al., 2007; Lascano et al., 2016). Although this isomer is not a direct cause of MFD, it is often used as a proxy for cases of MFD and persistence of the altered BH pathway (Lock et al., 2007). The *trans*-11 C18:1 isomer associated with the normal BH pathway is a substrate of the desaturation of *cis*-9,*trans*-11 CLA (Harvatine et al., 2009a). In our previous study, a tendency to decrease the yield of total *trans*-FA in milk was observed when cows were fed magnesium oxide and calcium-magnesium dolomite, whereas we observed a decrease in the yield of *trans*-10 C18:1 and an increase in the yield of *trans*-11 C18:1 in milk fat compared with nonsupplemented diet (Razzaghi et al., 2021). Remarkably, the concentrations of *cis*-9,*cis*-12 C18:2 and *cis*-9,*trans*-11 CLA were not affected by magnesium-based alkalizers, in contrast to previous observations in the recovery of MFD, where the concentrations of *trans*-11 C18:1 and *cis*-9,*trans*-11 CLA were similarly affected (Rico et al., 2015). In the current study, the reason for the lack of simultaneous changes in these 2 isomers is unclear, but it could be related to the lower PUFA loading compared with previous MFD recovery studies.

CONCLUSIONS

Overall, a diet containing magnesium-based alkalizers improved milk fat concentration and yield within 4 d of supplementation in lactating dairy cows that experienced biohydrogenation-induced MFD. Responses to milk fat synthesis remained greater for 7 d after cessation of supplementation, whereas this positive effect disappeared from 8 d after cessation of magnesium supplementation until the end of the experiment. The difference in milk fat concentration and yield was associated with lower and greater concentrations of *trans*-10 and *trans*-11 C18:1, respectively, during the experiment. This suggested that the effect of magnesium-based alkalizers on milk fat synthesis was mediated by a shift in the ruminal C18:2 biohydrogenation pathway. The shifts in FA profile indicated that dietary magnesium oxide and calcium-magnesium dolomite restored milk fat synthesis by promoting the normal BH pathway and reducing flux through the alternative pathway. Therefore, calcium-magnesium dolomite and magnesium oxide have a similar ability to improve milk fat synthesis in dairy cows under MFD conditions.

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