Methane prediction based on individual or groups of milk fatty acids for dairy cows fed rations with or without linseed

Stefanie W. Engelke,¹ Gürbüz Daş,¹ Michael Derno,¹ Armin Tuchscherer,² Klaus Wimmers,³ Michael Rychlik,⁴ Hermine Kienberger,⁵ Werner Berg,⁶ Björn Kuhla,¹ and Cornelia C. Metges^{1,7*}

¹Institute of Nutritional Physiology “Oskar Kellner,” Leibniz Institute for Farm Animal Biology (FBN), Wilhelm-Stahl-Allee 2, 18196 Dummerstorf, Germany

²Institute of Genetics and Biometry, Leibniz Institute for Farm Animal Biology (FBN), Wilhelm-Stahl-Allee 2, 18196 Dummerstorf, Germany

³Institute of Genome Biology, Leibniz Institute for Farm Animal Biology (FBN), Wilhelm-Stahl-Allee 2, 18196 Dummerstorf, Germany

⁴Analytical Food Chemistry, Technical University of Munich, Maximus-von-Imhof-Forum, 85354 Freising, Germany

⁵Bavarian Center for Biomolecular Mass Spectrometry, Gregor-Mendel-Strasse 4, 85354 Freising, Germany

⁶Department of Technology Assessment and Substance Cycles, Leibniz Institute for Agricultural Engineering and Bioeconomy (ATB), Max-Eyth-Allee 100, 14469 Potsdam, Germany

⁷Nutritional Physiology and Animal Nutrition, Faculty of Agriculture and Environmental Sciences, University of Rostock, 18059 Rostock, Germany

ABSTRACT

Milk fatty acids (MFA) are a proxy for the prediction of CH₄ emission from cows, and prediction differs with diet. Our objectives were (1) to compare the effect of diets on the relation between MFA profile and measured CH₄ production, (2) to predict CH₄ production based on 6 data sets differing in the number and type of MFA, and (3) to test whether additional inclusion of energy-corrected milk (ECM) yield or dry matter intake (DMI) as explanatory variables improves predictions. Twenty dairy cows were used. Four diets were used based on corn silage (CS) or grass silage (GS) without (L0) or with linseed (LS) supplementation. Ten cows were fed CS-L0 and CS-LS and the other 10 cows were fed GS-L0 and GS-LS in random order. In feeding wk 5 of each diet, CH₄ production (L/d) was measured in respiration chambers for 48 h and milk was analyzed for MFA concentrations by gas chromatography. Specific CH₄ prediction equations were obtained for L0-, LS-, GS-, and CS-based diets and for all 4 diets collectively and validated by an internal cross-validation. Models were developed containing either 43 identified MFA or a reduced set of 7 groups of biochemically related MFA plus C16:0 and C18:0. The CS and LS diets reduced CH₄ production compared with GS and L0 diets, respectively. Methane yield (L/kg of DMI) reduction by LS was higher with CS than GS diets. The concentrations of C18:1 *trans* and n-3 MFA differed among GS and CS diets. The LS diets

resulted in a higher proportion of unsaturated MFA at the expense of saturated MFA. When using the data set of 43 individual MFA to predict CH₄ production (L/d), the cross-validation coefficient of determination (R^2_{CV}) ranged from 0.47 to 0.92. When using groups of MFA variables, the R^2_{CV} ranged from 0.31 to 0.84. The fit parameters of the latter models were improved by inclusion of ECM or DMI, but not when added to the data set of 43 MFA for all diets pooled. Models based on GS diets always had a lower prediction potential (R^2_{CV} = 0.31 to 0.71) compared with data from CS diets (R^2_{CV} = 0.56 to 0.92). Models based on LS diets produced lower prediction with data sets with reduced MFA variables (R^2_{CV} = 0.62 to 0.68) compared with L0 diets (R^2_{CV} = 0.67 to 0.80). The MFA C18:1 *cis*-9 and C24:0 and the monounsaturated FA occurred most often in models. In conclusion, models with a reduced number of MFA variables and ECM or DMI are suitable for CH₄ prediction, and CH₄ prediction equations based on diets containing linseed resulted in lower prediction accuracy.

Key words: dairy cow, methane emission prediction, methane mitigation, methane proxy, milk fatty acids

INTRODUCTION

Methane is a greenhouse gas and a product of rumen fermentation (Hristov et al., 2013). Methane production (L/d) is a heritable trait, and genetic selection for low-emitting cows is a promising mitigation option (Pickering et al., 2015; Negussie et al., 2017). Methane emission quantification in respiration chambers is considered as the gold standard but unsuitable for large-scale individual animal measurements (Hammond et al., 2016; Patra, 2016). One promising proxy for the prediction of CH₄ production is the concentration of

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*Corresponding author: metges@fbn-dummerstorf.de

milk fatty acids (**MFA**; van Gastelen and Dijkstra, 2016; Negussie et al., 2017). Dietary composition affects the relationship between MFA and CH₄ emission parameters (Mohammed et al., 2011; Dijkstra et al., 2016; Rico et al., 2016). Milk fatty acids synthesized de novo are predominantly generated from rumen acetate and BHB (Shingfield et al., 2013), which are derived from fiber fermentation, and they are positively associated with ruminal CH₄ production (Shingfield et al., 2013; Castro-Montoya et al., 2016a). Forage type in ruminant diets can affect CH₄ production as well as MFA pattern (Beauchemin et al., 2008; Hart et al., 2015; van Gastelen et al., 2015). Likewise, a high starch content favors propionate synthesis, which consumes fermentative hydrogen and can thus reduce CH₄ production (Hook et al., 2010; Knapp et al., 2014). The MFA groups of long-chain MUFA and PUFA and the individual C18:0 originate from plant fat-derived UFA (Ferlay et al., 2013; Kliem and Shingfield, 2016; Meignan et al., 2017) are either directly transferred into milk fat (e.g., C18:3n-3) or influence rumen fermentation, thereby altering the pattern and level of precursors for MFA synthesis and affect the postabsorptive lipid metabolism (de novo MFA synthesis, saturation or desaturation rates, or FA elongation; Angulo et al., 2012; Lanier and Corl, 2015; Meignan et al., 2017). In addition, dietary fat-associated decreases of fiber degradability and toxic effects of PUFA on archaea reduce CH₄ production (Maia et al., 2007; Benchaar et al., 2015). Furthermore, fat supplements can also affect DMI and, thus, net production of short-chain MFA.

Several authors have used MFA concentrations to develop CH₄ prediction equations that differ in type and numbers of explanatory MFA (e.g., van Lingen et al., 2014; Castro-Montoya et al., 2016b; Rico et al., 2016). For example, van Lingen et al. (2014) developed CH₄ predictions based on 21 individual and 3 groups of MFA concentrations, whereas Rico et al. (2016) included 83 individual MFA and sums of different MFA proportions. van Gastelen and Dijkstra (2016) pointed out that only C17:1 *cis*-9 and C18:1 *cis*-11 appeared in 2 or more of the prediction equations published, indicating that explanatory MFA are extremely diverse, which likely is a consequence of differences in diet composition (Dijkstra et al., 2011; Mohammed et al., 2011; Rico et al., 2016). We were interested to know how prediction equations change when the MFA data changes in the number of variables and whether individual MFA or groups of related MFA are used.

It has been also discussed whether the combination of MFA concentrations with additional explanatory parameters would improve prediction (Castro-Montoya et al., 2016b; Negussie et al., 2017; van Gastelen et al.,

2017). The DMI of cows is known as the main determinant for CH₄ production (Knapp et al., 2014), and DMI is associated with milk yield (Hristov et al., 2013). Consequently, milk yield could be a potential alternative for DMI as a variable in prediction equations (Hristov et al., 2013; Negussie et al., 2017). We have reported that a prediction equation based on MFA, determined by mid-infrared spectroscopy, which contained DMI as an additional explanatory variable, showed a similar coefficient of determination when DMI was replaced by ECM (Engelke et al., 2018).

Thus, our hypothesis was that CH₄ prediction equations based on MFA depend on the number of MFA or MFA group variables, and that inclusion of DMI or ECM together with MFA improves prediction equations. Hence, the objectives of this study were (1) to determine the effect of dietary composition on the relationship between MFA concentrations and CH₄ production, with special emphasis on the effect of linseed supplementation; (2) to compare CH₄ prediction models based on differently sized and composed MFA data sets; and (3) to test whether the inclusion of DMI or ECM in addition to MFA concentrations as variable improves prediction.

MATERIALS AND METHODS

Animals, Experimental Design, and Diets

The procedures performed in our study were in agreement with the German Animal protection law and approved by the relevant authority (Landesamt für Landwirtschaft, Lebensmittelsicherheit und Fischereiwesen Mecklenburg-Vorpommern, Germany; permission no. 7221.3-1-014/14). Twenty lactating German Holstein cows (106 ± 28 DIM, 29.5 ± 7.7 kg of ECM/d, 580 ± 57 kg of BW; mean $\pm$ SD), of which 15 cows were in second and 5 cows were in third lactation, were purchased from a dairy farm located in the region of Mecklenburg-Western Pomerania, Germany. All cows were sired by 1 bull (Omega 802670; Rinderzucht Mecklenburg-Vorpommern GmbH, Woldegk, Germany). Cows were kept in tiestalls, had free access to water, and were offered a TMR for ad libitum intake. Milking occurred at 0630 and 1630 h. To produce a wide range of CH₄ production values, we used 2 basal TMR, which were composed close to what is commonly used in German dairy farming. The major forage component was either corn silage (**CS**) or grass silage (**GS**), supplemented with (**LS**) or without linseed (**L0**; Table 1). Diets contained grass silage and corn silage at DM levels of 130 and 450 g/kg (CS) or 360 and 190 g/kg (GS), respectively. The LS diets contained 60 g of fat/

kg of DM, whereas the L0 diets had 30 g of fat/kg of DM. The diets had the same target energy level of 7 MJ of NE_L/kg of DM, which was achieved by replacing the linseed product with an isoenergetic amount of starch in the TMR. Forage-to-concentrate ratio differed from 66:34 (GS-L0), 73:27 (GS-LS), 65:35 (CS-L0), to 68:32 (CS-LS). We randomly selected 5 of the 20 cows to be fed the CS-L0 diet in the first 5-wk period (period A) and the CS-LS diet in the second 5-wk period (period B), whereas we randomly selected 5 other cows to be fed the CS-LS diet during the first 5-wk period and CS-L0 diet during the second 5-wk period. Another 5

cows were fed GS-L0 diet first and GS-LS diet second, whereas the remaining 5 cows were given the opposite treatments. The transition from the standard diet to an experimental diet or from one experimental diet to the other was made over 5 d in experimental week (EW) 0 and 6, respectively, in a step-wise fashion by replacing 20, 40, 60, 80, and 100% of DM of the standard diet by the experimental diet. The study was conducted in 5 experimental blocks. Each block ran over a total of 12 EW and included 4 cows, all 4 diets, and 2 periods (A and B). That means 2 cows per each diet were measured within each block. Cows assigned to the CS

Table 1. Ingredients and chemical composition of the experimental TMR consisting of basal rations based largely on grass silage (GS) or corn silage (CS) without (L0) and with (LS) linseed supplementation (means $\pm$ SD; n = 5)

Item	GS		CS	
	L0	LS	L0	LS
Ingredient (g/kg of DM)				
Grass silage	343 $\pm$ 31.0	380 $\pm$ 15.0	122 $\pm$ 21.5	142 $\pm$ 34.3
Corn silage	181 $\pm$ 33.2	195 $\pm$ 37.9	445 $\pm$ 46.7	452 $\pm$ 44.0
Straw, barley	52.1 $\pm$ 7.0	60.8 $\pm$ 23.2	39.8 $\pm$ 7.2	34.0 $\pm$ 10.3
Grass hay	80.8 $\pm$ 12.0	93.5 $\pm$ 22.0	43.2 $\pm$ 3.1	43.5 $\pm$ 3.0
Corn, ground	—	—	92.7 $\pm$ 9.2	25.4 $\pm$ 27.2
Soy extract meal	25.4 $\pm$ 26.8	40.1 $\pm$ 19.2	114 $\pm$ 14.2	87.1 $\pm$ 9.2
Barley, ground	99.5 $\pm$ 17.9	—	—	—
Wheat, ground	101 $\pm$ 9.8	—	—	—
Linseed product ¹	—	143 $\pm$ 11.4	—	137 $\pm$ 9.9
Concentrate ²	109 $\pm$ 19.7	78.4 $\pm$ 13.0	129 $\pm$ 7.3	65.0 $\pm$ 32.6
Mineral/vitamin mix ³	9.1 $\pm$ 0.5	9.5 $\pm$ 0.6	10.1 $\pm$ 0.6	10.1 $\pm$ 0.6
Calcium carbonate ⁴	—	—	4.1 $\pm$ 0.3	4.1 $\pm$ 0.3
Grass to corn silage ratio	1.90	1.95	0.27	0.31
Forage:concentrate ratio	66:34	73:27	65:35	68:32
DM (g/kg of FM ⁵)	468 $\pm$ 33.0	452 $\pm$ 45.0	446 $\pm$ 32.0	439 $\pm$ 33.0
Nutrients (g/kg of DM)				
Crude ash	69.9 $\pm$ 2.8	75.8 $\pm$ 2.5	66.1 $\pm$ 3.0	65.7 $\pm$ 4.2
CP	161 $\pm$ 9.0	173 $\pm$ 14.4	169 $\pm$ 11.7	163 $\pm$ 11.1
Crude fiber	166 $\pm$ 7.9	186 $\pm$ 7.1	155 $\pm$ 3.3	161 $\pm$ 4.5
Crude fat	28.1 $\pm$ 1.7	58.3 $\pm$ 7.8	29.8 $\pm$ 2.4	56.4 $\pm$ 6.5
Sugar	51.3 $\pm$ 15.7	50.1 $\pm$ 16.2	31.4 $\pm$ 7.9	30.3 $\pm$ 11.1
Starch	218 $\pm$ 12.0	110 $\pm$ 18.7	261 $\pm$ 25.4	216 $\pm$ 34.5
aNDF ⁶	371 $\pm$ 18.1	410 $\pm$ 12.9	330 $\pm$ 10.9	352 $\pm$ 9.2
ADF	198 $\pm$ 9.8	224 $\pm$ 7.9	186 $\pm$ 5.4	194 $\pm$ 5.0
NE _L (MJ/kg of DM)	6.9 $\pm$ 0.2	6.9 $\pm$ 0.1	7.0 $\pm$ 0.1	7.1 $\pm$ 0.1

¹Omegalin 60 (Spezialfutter Neuruppin GmbH und Co.KG, Neuruppin, Germany; per kg feed; 88% DM): 60% extruded linseed Tradilin and 40% bran. Composition: 19.5% CP, 25% crude fat, 8.5% crude fiber; fatty acids (% of fat): 6% palmitic acid (C16:0), 18.4% oleic acid (C18:1 *cis*-9), 18.5% linoleic acid (C18:2 *cis*-9,*cis*-12), 55% linolenic acid (C18:3 *cis*-9,*cis*-12,*cis*-15), and 12.8 MJ of NE_L/kg of DM.

²Concentrate MF 2000 (Vollkraft Mischfutterwerke GmbH, Güstrow, Germany; per kg feed; 88% DM): 33% extracted soy meal, 20% corn, 17% wheat gluten, 8% extracted rapeseed meal, 5% sugar beet pulp, 2% sodium hydrogen carbonate, 1.3% calcium carbonate, 0.2% sodium chloride. Composition: 24% CP, 3.3% crude fat, 6.2% crude fiber, 8.4% crude ash, 0.7% calcium, 0.5% phosphorus, 0.65% sodium, and 7.1 MJ of NE_L/kg of DM.

³Rinderstolz 9522 Salvana (Tierernährung, Kl.-O. Sparrieshoop, Germany; per kg feed; 88% DM): 39.3% calcium carbonate, 21.7% monocalcium phosphate, 21% sodium chloride, 11.9% magnesium oxide, 2% sugar beet molasses. Composition: 92% crude ash, 20% calcium, 8% sodium, 6% magnesium, 5% phosphorus, 1,000,000 IU of vitamin A, 200,000 IU of vitamin D₃, and 4500 mg of vitamin E.

⁴Kreidekalk (Spezialfutter Neuruppin GmbH und Co.KG): calcium carbonate; 37% calcium.

⁵Fresh matter.

⁶aNDF = (amylase) neutral detergent fiber.

Table 2. Performance and methane emission parameters of cows fed basal rations based on grass silage (GS) or corn silage (CS) with or without linseed supplementation (LS or L0, respectively)

Item	GS ¹		CS ¹		SE	P-value ²		
	L0	LS	L0	LS		B	L	B × L
DMI (kg/d)	15.94	14.80	18.91	17.76	0.985	0.024	0.061	0.990
ECM (kg/d)	22.61	22.01	29.04	28.28	1.359	0.002	0.334	0.910
Milk fat (%)	4.48	4.50	4.08	3.83	0.202	0.035	0.364	0.292
CH ₄ emission parameters								
CH ₄ production (L/d)	506.8	453.0	587.7	500.6	30.40	0.089	0.002	0.373
CH ₄ intensity (L/kg of ECM)	23.3	21.3	20.2	17.7	1.27	0.050	0.006	0.750
CH ₄ yield (L/kg of DMI)	32.7	31.9	31.1	28.2	1.84	0.289	0.011	0.138

¹LSM.²ANOVA *F*-test for the effects of basal ration (B), linseed supplementation (L), or their interaction (B × L).

and GS diets did not differ in their mean baseline CH₄ production (506 and 477 L/d, respectively; *P* = 0.441).

To keep feed composition as constant as possible over time, experimental diets were mixed once a week, conserved with 1% granulated propionic acid (BERGO TMR-stabil G, Bergophor Futtermittelfabrik Dr. Berger GmbH & Co.KG, Kulmbach, Germany), and vacuum-packaged in thirty 40-kg plastic bags (NeuRo Planen GmbH, Neuendorf, Germany). Cows were fed ad libitum from these bags twice daily at 0730 and 1730 h. Feed intake was recorded daily. Three cows had to be removed in the GS-LS period because of illness or feed refusal. The diet composition was calculated in accordance to recommendations of the German Society of Nutritional Physiology (Gesellschaft für Ernährungsphysiologie, 2001).

Feed Sampling and Analyses

Feed samples were collected during TMR mixing and stored at −20°C. At the end of each experimental block, feed samples were pooled to determine DM. The DM content was determined by drying at 60°C for 24 h and then at 103°C for 4 h (Naumann et al., 1976). Analyses of nutrient composition in feedstuffs (Table 1) were performed according to the Weender standard analysis (Naumann et al., 1976), with modifications by Van Soest et al. (1991), by the accredited feed laboratory of Landwirtschaftliche Untersuchungs-und Forschungsanstalt der LMS Agrarberatung GmbH (LUFA, Rostock, Germany). The energy content (NE_L) was calculated according to the German Society of Nutritional Physiology (Gesellschaft für Ernährungsphysiologie, 2001), except for the linseed product Omegalin 60 (Spezialfutter Neuruppin GmbH und Co. KG, Neuruppin, Germany) and concentrate MF 2000 (Vollkraft Mischfutterwerke GmbH, Güstrow, Germany), which were analyzed according to German feed regulations (Ger-

man Federal Law Gazette, 1981) §13 by LUFA. Nutrient intake of cows was calculated from the DMI and analyzed nutrient contents in feedstuffs (Supplemental Table S1; <https://doi.org/10.3168/jds.2018-14911>).

Methane Measurements

Methane production (L/d) was recorded in EW 5 and 11 for 2 subsequent 24-h periods, each using 4 open-circuit respiration chambers as described by Derno et al. (2009) and Bielak et al. (2016). Cows were fed with respective TMR diets at 0730 and 1730 h and feed intake was recorded continuously by feed troughs placed on balances and summarized over 24 h. Milking occurred at 0630 and 1630 h. Drinking water was freely available. The concentrations of CH₄ were measured at 6-min intervals throughout 23.9 h; the data were normalized to 24 h. The temperature and relative humidity in the chambers were 15°C and 65%, respectively. The mean recovery rate of the chambers was 99.9 ± 0.96%. Methane production parameters were calculated as CH₄ production (L/d), CH₄ yield defined as liters of CH₄ per kilogram of DMI, and CH₄ intensity defined as liters of CH₄ per kilogram of ECM (Table 2), where ECM (kg/d) = [1.05 + 0.38 × milk fat (%) + 0.21 × milk protein (%)]/3.28 × milk yield (kg/d) (Spiekens et al., 2009).

Milk Sampling and Analyses

During the CH₄ measurements, aliquots of milk from the evening of the first and the morning of the second 24-h period were pooled proportionally according to milk yield at each milking (0.25% of milk yield). Proximate milk composition was analyzed by the state control association Mecklenburg-Western Pomerania (Landeskontrollverband für Leistungs-und Qualitätsprüfung Mecklenburg-Vorpommern e.V.,

Güstrow, Germany) for milk fat and protein content using mid-infrared spectroscopy (MilkoScan FT6000 and MilkoScan FT+, Foss, Hillerød, Denmark). Aliquots of the pooled milk samples were stored at -20°C until MFA analysis using GC (Firl et al., 2014) performed by the Bavarian Biomolecular Mass Spectrometry Center (Freising, Germany). Milk lipid was extracted by a 1:1 chloroform:methanol mixture and esterified with trimethylsulfonium hydroxide to form FAME. A GC-flame ionization detector instrument (Hewlett Packard 6890, Palo Alto, CA) with a 7683 autosampler was equipped with a $100\text{ m} \times 0.25\text{ mm}$ column (CP 7420; $0.25\text{ }\mu\text{m}$ film thickness; Agilent Technologies, Böblingen, Germany; Firl et al., 2014). Identifications of peaks were made by comparison with known FAME standards. The GC analyses resulted in the separation and quantification (% of total lipids) of 46 individual MFA. Because not all individual MFA were detected in all milk samples, we excluded those which occurred in less than 85% of the milk samples (C18:1 *trans*-10, C18:2 *trans*-10, *cis*-12, and C21:0), which resulted in a total of 43 individual MFA (Table 3). Because SFA have been shown to be positively associated and UFA, MUFA, PUFA, C18:1 *cis* and *trans* isomers, and n-3 MFA have been shown to be negatively associated with CH_4 emission (Chilliard et al., 2009; Castro-Montoya et al., 2016a; van Gastelen and Dijkstra, 2016), we were interested to know the predictive power of equations containing groups of biochemically related MFA instead of individual MFA. Thus, MFA concentrations were summed as groups of SFA, UFA, MUFA, PUFA, C18:1 *cis* and *trans* isomers, and n-3 MFA (Supplemental Table S2; <https://doi.org/10.3168/jds.2018-14911>) according to our previous study in which MFA were predicted by infrared spectroscopy (Engelke et al., 2018).

Calculations and Statistical Analysis

ANOVA. The dependent variables CH_4 production, DMI, and ECM yield were measured on 2 consecutive days in EW 5 and 11 and were averaged per day in each EW. Data were analyzed with repeated measures ANOVA using PROC MIXED (SAS/STAT 9.3; SAS Institute Inc., Cary, NC). The model contained the fixed effects of basal ration (CS, GS), linseed supplementation (LS, L0), the interaction effect between basal ration and linseed supplementation, as well as effects of experimental blocks (1–5), periods (A or B), and the order (LS in period A or B first). The covariance structure was set to be compound symmetry. Effects were considered significant at $P < 0.05$ and least squares means were compared using the Tukey test with the

SLICE statement for performing a partitioned analysis of the least squares means for the interaction. Data are presented as least squares means $\pm$ standard error if not given otherwise.

Regression Models and Validation. Correlations were calculated between CH_4 production (L/d) values and concentrations of individual MFA and MFA groups using PROC CORR of SAS. Regression equations for the dependent variable CH_4 production (L/d) were constructed from MFA data of each diet separately using PROC REG of SAS with the STEPWISE variable selection method. Regression models for MFA data from combined data categorized by basal diets or linseed supplementation, as well as for data from all 4 diets collectively, were also estimated. Six data sets of variables were used for the stepwise variable selection to enter in the regression equations. Data sets 1 to 3 included 43 different individual MFA, whereas data sets 4 to 6 were generated according to the MFA grouping of Engelke et al. (2018) and contained 7 groups of MFA (SFA, UFA, MUFA, and PUFA, C18:1 *trans*, C18:1 *cis*, and n-3 MFA) as summed concentrations plus the individual MFA C16:0 and C18:0 as key MFA in milk fat metabolism (Supplemental Table S2; <https://doi.org/10.3168/jds.2018-14911>). Data set 1 included the 43 individual MFA only and data set 2 included 43 individual MFA and ECM data, whereas data set 3 included 43 individual MFA and DMI data. Data set 4 included the reduced data set of 7 groups of MFA and the 2 individual MFA only. Data set 5 included the same reduced data set of MFA variables as well as ECM, whereas data set 6 included the same reduced data set of MFA as well as DMI.

To have the most relevant explanatory variables in the regression equations and to avoid over-fitted regression models, we preselected candidate explanatory variables through their correlation coefficients with the dependent variable (i.e., CH_4 production). For this purpose, only variables that had a significant correlation ($P < 0.05$) with CH_4 production were allowed to potentially enter in the models. Thereafter, the preselected variables were allowed to enter and remain in the regression equations with a significance threshold of $P \leq 0.05$ (i.e., entry and stay levels of the stepwise variable selection method). Subsequently, explanatory variables in the final regression equations derived from data set 1 and 4 were made available to potentially enter in the regression models for data sets 2 and 5 together with ECM or 3 and 6 together with DMI, respectively. In this way, ECM or DMI was used to additionally improve regression models derived from data sets 1 and 4 or could replace 1 or more of the explanatory variables in the final regression models. In the final regression models

of each data set, multicollinearity was assessed through variation inflation factor, and explanatory variables were allowed to have a variation inflation factor <10, as suggested by Kaps and Lamberson (2004).

As cows were used rotationally with 2 different diets (L0 and LS) at 2 different periods, the regression equations based on data specific to basal ration and all diets included pooled data of periods (e.g., EW 5 and 11). This was possible because period did not affect CH₄ production and MFA profiles. The regression

equations based on data specific to linseed supplementation were not concerned with pooled data due to the experimental setup. The performance of the developed CH₄ prediction equations was assessed by an internal cross-validation with the existing data set. A cross-validation was performed by leaving 1 animal out at a time and performing a calibration with the remaining animals per model (each diet separately, basal diets, linseed supplementation or not, all diets collectively; Moraes et al., 2014). The root mean square error of

Table 3 Individual milk fatty acid (MFA) composition (% of total lipids) of cows fed basal rations based on grass silage (GS) or corn silage (CS) with (LS) or without (L0) linseed supplementation

MFA (% of total lipids)	GS ¹		CS ¹		SE	P-value ²		
	L0	LS	L0	LS		B	L	B × L
C4:0	3.81	3.86	3.78	4.14	0.173	0.467	0.123	0.248
C6:0	2.46	2.11	2.50	2.22	0.094	0.472	0.001	0.577
C8:0	1.45	1.08	1.17	1.48	0.056	0.368	0.001	0.410
C10:0	3.25	2.04	3.34	2.32	0.122	0.165	0.001	0.265
C10:1	0.41	0.25	0.39	0.24	0.019	0.681	0.001	0.734
C11:0	0.13	0.03	0.10	0.05	0.016	0.869	0.001	0.125
C12:0	3.92	2.20	3.97	2.57	0.147	0.184	0.001	0.135
C12:1	0.12	0.06	0.11	0.06	0.008	0.885	0.001	0.651
C13:0	0.16	0.06	0.15	0.09	0.015	0.728	0.001	0.069
C14:0	11.13	8.79	11.55	9.45	0.239	0.038	0.001	0.510
C14:0 <i>iso</i>	0.10	0.09	0.12	0.09	0.013	0.398	0.019	0.128
C14:1 <i>cis</i> -9	1.36	0.91	1.32	0.91	0.068	0.791	0.001	0.637
C15:0	1.37	0.80	1.25	0.89	0.095	0.851	0.001	0.142
C15:0 <i>iso</i>	0.19	0.17	0.24	0.18	0.015	0.082	0.012	0.165
C15:0 <i>anteiso</i>	0.42	0.38	0.44	0.44	0.036	0.269	0.473	0.518
C16:0	34.40	24.05	33.07	22.93	1.007	0.312	0.001	0.860
C16:0 <i>iso</i>	0.25	0.21	0.28	0.24	0.030	0.430	0.031	0.936
C16:1 <i>cis</i> -9	2.06	1.39	1.89	1.11	0.137	0.120	0.001	0.593
C16:1 <i>trans</i> -9	0.03	0.03	0.03	0.04	0.004	0.090	0.058	0.558
C17:0	0.58 ^a	0.45 ^c	0.56 ^a	0.51 ^b	0.020	0.306	0.001	0.015
C17:0 <i>iso</i>	0.28	0.26	0.30	0.27	0.021	0.420	0.132	0.747
C17:0 <i>anteiso</i>	0.45	0.36	0.40	0.47	0.028	0.363	0.001	0.547
C17:1 <i>cis</i> -9	0.26	0.17	0.23	0.16	0.017	0.244	0.001	0.373
C18:0	7.50	12.81	8.52	14.22	0.495	0.022	0.001	0.632
C18:0 <i>iso</i>	0.05	0.03	0.05	0.04	0.004	0.127	0.010	0.338
C18:1 <i>cis</i> -9	17.33 ^b	26.20 ^{a*}	17.21 ^b	23.91 ^{a*}	0.889	0.306	0.001	0.013
C18:1 <i>cis</i> -11	0.61	0.58	0.48	0.62	0.095	0.585	0.588	0.377
C18:1 <i>cis</i> -12	0.23	0.63	0.29	0.74	0.034	0.017	0.001	0.455
C18:1 <i>trans</i> -9	0.37	1.06	0.57	1.29	0.125	0.014	0.001	0.936
C18:1 <i>trans</i> -11	0.97	2.80	1.05	3.02	0.162	0.283	0.001	0.640
C18:2 <i>cis</i> -9, <i>cis</i> -12	2.15	2.92	2.24	1.87	0.576	0.396	0.666	0.219
C18:2 <i>cis</i> -9, <i>trans</i> -11	0.52	1.21	0.52	1.18	0.073	0.828	0.001	0.836
C18:3n-6	0.03	0.03	0.03	0.03	0.002	0.655	0.309	0.843
C18:3n-3	0.62	1.15	0.38	1.12	0.068	0.021	0.001	0.120
C20:0	0.11	0.14	0.13	0.15	0.007	0.126	0.001	0.429
C20:1 <i>cis</i> -11	0.08	0.11	0.05	0.06	0.023	0.117	0.175	0.474
C20:2n-6	0.02	0.02	0.03	0.02	0.006	0.944	0.652	0.764
C20:3n-6	0.07 ^b	0.04 ^c	0.10 ^a	0.05 ^c	0.006	0.016	0.001	0.003
C20:4n-6	0.07	0.07	0.07	0.06	0.008	0.822	0.385	0.416
C20:5n-3	0.05	0.07	0.04	0.05	0.012	0.082	0.055	0.819
C22:0	0.03	0.02	0.02	0.02	0.003	0.100	0.495	0.784
C22:5n-3	0.08 ^a	0.06 ^b	0.07 ^a	0.08 ^{ab}	0.007	0.642	0.168	0.043
C24:0	0.03	0.03	0.03	0.03	0.003	0.251	0.916	0.077

^{a-c}Values within a row with differing superscripts denote B × L interactions ($P < 0.05$).

¹LSM.

²ANOVA F-test for the effects of basal ration (B), linseed supplementation (L), or their interaction (B × L).

* $P < 0.10$.

cross-validation and the cross-validation coefficient of determination (R^2_{CV}) were estimated (Moraes et al., 2014). In addition, we report adjusted coefficient of determination (R^2_{Adj}) to account for the number of variables in the final models. We further evaluated the models using the concordance correlation coefficient (CCC) as reported earlier (Dijkstra et al., 2016; van Gastelen et al., 2017). Due to the low number of observations ($n = 7-10$) and variables with multicollinearity, we refrained from reporting predictions for individual diets.

RESULTS

Animal Performance and Methane Production

Cows fed the CS ratios had a higher DMI ($P = 0.024$) and ECM yield ($P = 0.002$) than those fed GS ratios (Table 2). Accordingly, CP, crude fat, starch, and NE_L intakes ($P < 0.05$) were higher with CS than with GS ratios (Supplemental Table S1; <https://doi.org/10.3168/jds.2018-14911>). The intake of sugar was higher ($P = 0.005$) with GS ratios but fiber intake was similar with both basal diets. The percentage of milk fat was higher ($P = 0.035$) with GS compared with CS feeding (Table 2). Linseed supplementation had no effect on ECM and milk fat ($P > 0.3$), but we noted a tendency toward a lower DMI ($P = 0.061$) with LS compared with L0 diets (Table 2). Feeding LS ratios resulted in twice the crude fat intake ($P = 0.001$) as with L0 ratios (Supplemental Table S1).

Cows fed the CS basal diets tended to have higher CH_4 production compared with those cows fed the GS ratios ($P = 0.089$; Table 2). The CH_4 yield ranged from a maximum 50 L/kg of DMI in cows fed the GS-L0 ration to a minimum of 21 L/kg of DMI with the CS-LS diet (Supplemental Table S3; <https://doi.org/10.3168/jds.2018-14911>). Basal diet had no effect on CH_4 yield ($P = 0.289$), but CH_4 intensity was lower ($P = 0.050$) by about 15% with CS than with GS ratios. Furthermore, LS decreased the level of all CH_4 emission parameters as compared with cows fed L0. The reduction of enteric CH_4 emission levels due to LS amounted to 10 and 13% for CH_4 intensity and production, respectively ($P < 0.01$). Methane yield (L/kg of DMI) was reduced ($P = 0.011$) by approximately 6% when diets with LS were fed.

MFA Composition and Correlations with Methane Production

With CS compared with GS diets, C14:0, C18:0, C18:1 *cis*-12, C18:1 *trans*-9, C20:3n-6, and the sum of C18:1 *trans* MFA concentrations were higher whereas C18:3n-3 and the sum of n-3 MFA were lower (Tables 3 and 4; $P < 0.04$). The LS diets resulted in lower concentrations of individual even-chain SFA from C6:0 to C16:0, the group of SFA ($P < 0.001$), the off-chain SFA from C11:0 to C17:0 ($P < 0.001$), and the individual MUFA C10:1, C12:1, C14:1 *cis*-9, C16:1 *cis*-9, C17:1 *cis*-9, as well as C18:0 *iso* and C20:3n-6 ($P < 0.05$; Table 3). Most of C18 PUFA and C20:0 MFA

Table 4. Milk fatty acid (MFA) groups (% of total lipids) as sums of concentrations of individual MFA of cows fed basal rations based on grass silage (GS) or corn silage (CS), with (LS) or without (L0) linseed supplementation

MFA (% of total lipids)	GS ¹		CS ¹		SE	P-value ²		
	L0	LS	L0	LS		B	L	B × L
Σ SFA ³	72.06	60.32	72.32	62.41	1.453	0.457	0.001	0.369
Σ UFA ⁴	27.43	39.09	27.10	36.60	1.440	0.377	0.001	0.279
Σ MUFA	23.81	33.88	23.63	32.15	1.044	0.470	0.001	0.153
Σ PUFA	3.62	5.55	3.47	4.45	0.652	0.332	0.011	0.357
Σ C18:1 <i>cis</i> ⁵	18.17	27.32	17.99	25.26	0.948	0.362	0.001	0.053
Σ C18:1 <i>trans</i> ⁶	1.33	3.72	1.62	4.31	0.193	0.014	0.001	0.395
Σ n-3 FA ⁷	0.76	1.29	0.49	1.25	0.079	0.031	0.001	0.128
Σ n-6 FA ⁸	2.34	3.07	2.47	2.02	0.585	0.098	0.001	0.092

¹LSM.

²ANOVA F-test for the effects of basal ration (B), linseed supplementation (L), or their interaction (B × L).

³Sum of C4:0, C6:0, C8:0, C10:0, C11:0, C12:0, C13:0, C14:0, C14:0 *iso*, C15:0, C15:0 *iso*, C15:0 *anteiso*, C16:0, C16:0 *iso*, C17:0, C17:0 *iso*, C17:0 *anteiso*, C18:0, C18:0 *iso*, C20:0, C22:0, and C24:0.

⁴Unsaturated fatty acids (FA) represent the sum of MUFA (C10:1, C12:1, C14:1 *cis*-9, C16:1 *cis*-9, C16:1 *trans*-9, C17:1 *cis*-9, C18:1 *cis*-9, C18:1 *cis*-11, C18:1 *cis*-12, C18:1 *trans*-9, C18:1 *trans*-11, and C20:1 *cis*-11) and PUFA (C18:2 *cis*-9, *trans*-11, C18:2 *cis*-9, *cis*-12, C18:3 *cis*-6, *cis*-9, *cis*-12, C18:3 *cis*-9, *cis*-12, *cis*-15, C20:2n-6, C20:3n-6, C20:4n-6, C20:5n-3, and C22:5n-3).

⁵Sum of C18:1 *cis*-9, *cis*-11, and *cis*-12.

⁶Sum of C18:1 *trans*-9 and *trans*-11.

⁷Sum of C18:3n-3, C20:5n-3, and C22:5n-3.

⁸Sum of C18:2n-6, C18:3n-6, C20:2n-6, C20:3n-6, and C20:4n-6.

concentrations were higher with LS-containing diets ($P < 0.05$; Table 3). A basal ration $\times$ linseed interaction was found for 4 of the 43 individual MFA ($P < 0.05$; C17:0, C18:1 *cis*-9, C20:3n-6, and C22:5n-3; Table 3). The concentrations of the MFA groups UFA, MUFA, PUFA, sums of C18:1 *cis* or *trans* isomers, and the sum of n-3 FA were higher with LS compared with L0 diets ($P < 0.01$; Table 4).

For combined data of all diets collectively, positive correlations were found between CH_4 production and concentrations of selected individual SFA, the sum of all SFA, as well as C18:3n-6 ($P \leq 0.05$; Figure 1 and Supplemental Table S4; <https://doi.org/10.3168/jds.2018-14911>). We observed negative correlations between CH_4 production and the levels of individual MFA of C15:0 *iso*, C17:1 *cis*-9, C18:1 *cis*-9, C18:1 *cis*-11, C18:1 *cis*-12, C18:1 *trans*-9, C18:3n-3, and C22:0 and groups of UFA, MUFA, C18:1 *cis* and *trans* isomers, and n-3 MFA ($P \leq 0.05$). Correlations of CH_4 production with the levels of individual MFA differed for data of the various pooled diets (Figure 1 and Supplemental Table S4). For example, correlations between C10:0 and CH_4 production ranged from 0.07 ($P = 0.76$) for the L0 diets to 0.63 ($P = 0.006$) for the LS diets, and for C18:3n-3 from -0.17 ($P = 0.509$) for the GS diets to -0.70 ($P = 0.002$) for the LS diets; correlations for C24:0 ranged from 0.14 ($P = 0.605$) for the LS diets to 0.83 ($P = 0.001$) for the CS diets.

Methane Prediction Models Using the Complete Set of MFA Variables

The prediction of CH_4 production based on data sets 1 to 3 resulted in coefficients of determination of the models (R^2_{Model}) between 0.36 and 0.91 ($P < 0.01$), R^2_{Adj} between 0.32 and 0.89, R^2_{CV} between 0.47 and 0.92, and root mean square error between 38.78 and 97.41, whereas CCC was between 0.53 and 0.95 (Table 5). The R^2_{Adj} values were slightly but systematically lower than that for R^2_{Model} . The additional inclusion of ECM (data set 2) resulted in higher R^2_{Model} values, ranging from 0.61 to 0.91, and R^2_{CV} , from 0.71 to 0.92, as compared with data set 1. Nevertheless, for all diets combined, the R^2_{Model} (0.69) and R^2_{CV} (0.74) of the equations for data set 2 were lower compared with that of data set 1 ($R^2_{\text{Model}} = 0.81$ and $R^2_{\text{CV}} = 0.85$). The inclusion of DMI (data set 3) instead of ECM (data set 2) resulted in comparable prediction values, with an R^2_{Model} range from 0.62 to 0.91 and R^2_{CV} from 0.70 to 0.92 (Table 5). With the exception of the GS diets, the R^2_{CV} and CCC values based on data sets 1 to 3 were at least 0.74 and 0.82, respectively, or higher. The R^2_{CV} based on GS diets was always lower compared with that of CS diets

Figure 1. Heatmap of correlations between individual and groups of milk fatty acids (% of total lipids) and methane production (L/d) calculated from data of the combined basal diets [grass silage (GS), corn silage (CS)], each basal diet with (L0) or without (LS) linseed supplementation, and all diets collectively.

Table 5. Summary of quality parameters and validation results of the multiple regression equations predicting methane production (CH_4 , L/d) using the complete data set of 43 milk fatty acid (MFA) variable (data sets 1–3), or a reduced number of MFA variables and groups of MFA (data sets 4–6)¹

Pooled diets	N	R^2_{Model}	P_{Model}	R^2_{Adj}	CCC	RMSE	R^2_{CV}
Data set 1							
GS	17	0.36	0.011	0.32	0.53	97.41	0.47
CS	20	0.86	0.001	0.84	0.93	45.54	0.89
L0	20	0.85	0.001	0.83	0.92	60.74	0.87
LS	17	0.78	0.001	0.75	0.88	41.77	0.84
All diets	37	0.81	0.001	0.78	0.90	56.38	0.85
Data set 2							
GS	17	0.61	0.001	0.58	0.75	77.84	0.71
CS	20	0.91	0.001	0.89	0.95	41.79	0.91
L0	20	0.91	0.001	0.89	0.95	48.25	0.92
LS	17	0.78	0.001	0.75	0.88	41.77	0.84
All diets	37	0.69	0.001	0.67	0.82	75.28	0.74
Data set 3							
GS	17	0.62	0.001	0.60	0.77	79.74	0.70
CS	20	0.91	0.001	0.89	0.95	39.55	0.92
L0	20	0.90	0.001	0.88	0.95	50.25	0.91
LS	17	0.81	0.001	0.79	0.90	38.78	0.87
All diets	37	0.74	0.001	0.73	0.85	66.86	0.80
Data set 4							
GS	17	0.26	0.035	0.21	0.42	106.50	0.31
CS	20	0.42	0.002	0.39	0.59	83.24	0.56
L0	20	0.57	0.001	0.52	0.72	93.39	0.67
LS	17	0.49	0.002	0.46	0.66	60.61	0.62
All diets	37	0.30	0.001	0.28	0.46	93.75	0.48
Data set 5							
GS	17	0.61	0.001	0.58	0.75	77.67	0.71
CS	20	0.74	0.001	0.71	0.85	61.84	0.79
L0	20	0.70	0.001	0.69	0.82	74.30	0.80
LS	17	0.49	0.002	0.46	0.66	60.61	0.62
All diets	37	0.69	0.001	0.67	0.82	64.57	0.80
Data set 6							
GS	17	0.62	0.001	0.60	0.77	79.56	0.70
CS	20	0.80	0.001	0.78	0.89	54.31	0.84
L0	20	0.73	0.001	0.72	0.85	73.34	0.80
LS	17	0.56	0.001	0.53	0.72	57.01	0.68
All diets	37	0.73	0.001	0.71	0.84	62.45	0.81

¹Each row summarizes validation results for a regression equation specific to data of diets based on combinations of basal ration [grass silage (GS), corn silage (CS)] and linseed supplementation [without (L0), with (LS)] or all diets collectively. Data set 1 included MFA only. Data sets 2 and 3 included individual MFA and ECM or DMI, respectively, as independent variables. Data set 4 included MFA variables (reduced number of variables) only, whereas data sets 5 and 6 additionally included ECM or DMI, respectively. The R^2_{Model} and P_{Model} values, the adjusted R^2_{Adj} , the concordance correlation coefficient (CCC) of the model, the root mean square error (RMSE) of cross-validation as well as the cross-validation coefficient of determination (R^2_{CV}) are given

(R^2_{CV} from 0.47 to 0.71 vs. 0.89 to 0.92). We found no discernible pattern of predictive MFA as explanatory variables in the prediction equations. However, C18:1 *cis*-9 and C24:0 appeared more often in the prediction equations than others (Supplemental Table S5; <https://doi.org/10.3168/jds.2018-14911>).

Methane Prediction Models Using a Reduced Number of MFA Variables

Three data sets (data sets 4 to 6) comprising the concentrations of a reduced number of MFA and biochemically related groups of summed MFA (Supplemental Table S2; <https://doi.org/10.3168/jds.2018-14911>)

were used to develop multiple regression equations. Using data set 4, prediction of CH_4 production resulted in R^2_{Model} between 0.26 and 0.57, R^2_{Adj} between 0.21 and 0.52, CCC between 0.42 and 0.72, and a R^2_{CV} between 0.31 and 0.67 (Table 5). The additional inclusion of ECM (data set 5) or DMI (data set 6) increased the R^2_{Model} and R^2_{CV} values, respectively, compared with data set 4. The R^2_{Adj} values were similar to R^2_{Model} but systematically lower. Models for CS diets always resulted in a higher prediction (R^2_{CV} from 0.56 to 0.84) and higher CCC values (from 0.59 to 0.89) compared with GS diets. Similarly, models of data from L0 diets (R^2_{CV} from 0.67 to 0.80, CCC from 0.72 to 0.85) gave higher predictions as compared with LS diets (Table 5).

However, for the LS diets, the inclusion of ECM (data set 5) did not improve the model. Predictions based on data from all 4 diets collectively including ECM as an explanatory variable in the model enhanced the predictive power by 130% ($R^2_{\text{Model}} = 0.69$) and 69% ($R^2_{\text{CV}} = 0.80$) compared with data set 4 ($R^2_{\text{Model}} = 0.30$ and $R^2_{\text{CV}} = 0.48$). The dominant predictive MFA variable in the regression equations of data set 4 to 6 was the MFA group MUFA (Supplemental Table S5; <https://doi.org/10.3168/jds.2018-14911>).

DISCUSSION

Animal Performance, Methane Production Parameters, and MFA Composition

The CS diets resulted in a higher DMI and ECM yield than the GS diets, which was comparable to what was reported previously for diets with higher proportions of corn silage than grass silage (Kliem et al., 2008; Sterk et al., 2011; Livingstone et al., 2015). In contrast to earlier studies, the type of the basal ration did not affect CH_4 yield (L/kg of DMI) of cows (Beauchemin et al., 2008; Hart et al., 2015; van Gastelen et al., 2015). This was likely due to similar crude fiber, ADF, and NDF intakes with grass- and corn silage-based diets. To make the diets isoenergetic, wheat and barley were added to the GS-L0 diet, and this added starch, diluted the fiber content, and aligned the fiber intake of the 2 basal diets. Furthermore, effects of starch and sugar in the GS-L0 diets might have reduced CH_4 production, which otherwise was expected to be higher with diets containing higher proportions of grass silage (van Gastelen et al., 2015). Although the NDF/DMI value was still higher in GS-based diets, the DMI and energy intake of cows fed GS diets was, on average, 3 kg and 17% MJ of NE_L less per day, respectively, as compared with the CS-based diets. This is a limitation of our study and possibly affected the width of range of CH_4 production. Methane production (L/d) tended to be higher with CS compared with GS diets due to the higher DMI, which is known to explain 52 to 64% of CH_4 production (Knapp et al., 2014). In contrast, CH_4 intensity tended to be lower in the CS than in the GS group due to higher ECM yield, which illustrates the importance of a high milk performance to mitigate CH_4 intensity (Yan et al., 2010; Zehetmeier et al., 2012; Gerber et al., 2013).

We observed only few differences in MFA concentrations between basal rations, although others reported more diverse MFA patterns with different silage types (Kliem et al., 2008; Livingstone et al., 2015; van Gastelen et al., 2015). For example, diets with different

proportions of grass and corn silage differed in the MFA concentrations of C18:1 isomers, total CLA, C18:2n-6, and C18:3n-3 (Kliem et al., 2008; van Gastelen et al., 2015). The small differences in concentrations of only a few MFA among basal diets found here were due to similar crude fiber, ADF, and NDF intakes with the basal diets, as discussed above. This can limit the range of MFA concentrations, and thus may affect CH_4 prediction.

In the present study, dietary supplementation of linseed decreased the level of all CH_4 emission parameters by 6, 10 and 13% for CH_4 yield, intensity, and production, although the reductive effects were greater with CS diets, as was observed earlier (Benchaar et al., 2015; Martin et al., 2016). Others found reductions in CH_4 yield between 8 and 20% by feeding various diets supplemented with extruded linseed (5 to 10% of dietary DM) or linseed oil (4% of dietary DM; Benchaar et al., 2015; Martin et al., 2016; Bayat et al., 2018). A 10 g/kg of DM increase in dietary fat resulted in a lower CH_4 yield by 1 g/kg of DMI in cattle (Grainger and Beauchemin, 2011). An increase of 1% in dietary lipid content led to a 4 to 5% reduction of CH_4 production (Clark, 2013). In our study, the LS diets contained 28 g/kg of DM more crude fat than the L0 diets, which corresponds to a lower CH_4 yield by 2.8 g/kg of DM, or 12 to 15%. The actual reduction of CH_4 yield by LS supplementation of GS and CS diets was 0.7 and 2.1 g/kg of DM or 2.7 and 9.4%, respectively. Thus, in our study, the linseed supplementation reduced CH_4 yield less than predicted by Grainger and Beauchemin (2011) or Clark (2013), but the reduction was larger with the corn silage-based diet. The lesser mitigation observed in our study compared with Grainger and Beauchemin (2011) might be due to the fact that we did not add linseed oil but linseeds to the diets. This might reduce or slow down the release of linseed oil in the rumen and is thus less suppressive to methanogens. As argued by Benchaar et al. (2015), the type of basal diet might influence the effect of linseed oil on CH_4 mitigation. For example, Chung et al. (2011) observed a mitigation effect of linseed only in combination with feeding barley silage (33% of CH_4 yield) but not with grass hay. This is similar to what was found in our study, which might be due to the opposite effect of linseed and grass, grass hay, or grass silage on CH_4 production.

Linseed supplementation tended to reduce DMI. Others observed no decrease of DMI when linseed supplementation was moderate (≤ 60 g of fat/kg of DM) as compared with a control diet (Ferlay et al., 2013; Kliem et al., 2017; Meignan et al., 2017). In the present study, the higher concentrate content in L0 diets was used to compensate for the higher energy density of linseed,

which might have led to a slightly higher DMI because concentrate is known as a favored feedstuff of cows when forage is provided *ad libitum* (Reynolds, 2006; Allen, 2014). The LS-supplemented diets contained twice the amount of fat than the nonsupplemented diets, which affected the concentrations of the majority of MFA. Almost all SFA were decreased with linseed supplementation through the direct inhibitory effect of long-chain SFA and UFA on fiber digestibility and *de novo* synthesis of MFA (Maia et al., 2007; Glasser et al., 2008; Shingfield et al., 2013). In contrast, C4:0 in milk fat was not affected by LS-containing diets. Effects of linseed on C4:0 concentrations are inconsistent in the literature (Ferlay et al., 2013; van Lingen et al., 2014; Kliem et al., 2017). Concentrations of off- and branched-chain MFA, which are derived from microbial synthesis (Vlaeminck et al., 2015), decreased when diets were supplemented by linseed. Lower levels of off- and branched-chain MFA are presumably related to the toxic effects of dietary PUFA on rumen microbiota (Enjalbert et al., 2017). The intake of increased levels of C18:3n-3 MFA with LS-supplemented diets increased the proportion of C18 MFA in milk fat due to biohydrogenation (Buccioni et al., 2012; Meignan et al., 2017). Diets containing linseed decreased SFA and increased UFA, *trans* MFA, n-3 MFA, and C18:0 (Chilliard et al., 2009; Ferlay et al., 2013; Meignan et al., 2017); the magnitude of changes in MFA pattern depended on the amount and form of linseed supply (Chilliard et al., 2009; Shingfield et al., 2013). Ferlay et al. (2013) showed that the C18:1 *cis/trans* isomer profile was specific for the forage type. This was similar to our study, where the C18:1 *trans* isomers were forage type-dependent, whereas the addition of linseed increased both C18:1 *cis* and *trans* isomers.

Diet Effects on the Relationship Between MFA and CH₄ Production and CH₄ Prediction

Each diet produces a characteristic MFA pattern (Kliem et al., 2008; Ferlay et al., 2013; Shingfield et al., 2013). We compared the correlations of individual MFA with CH₄ production among pooled diets (GS, CS, L0, LS, all diets) and found—depending on the dietary group—positive, negative, or no correlations. When using MFA data of all 4 diets collectively, we observed moderate correlations between CH₄ production and individual MFA (maximum $r = 0.52$ vs. -0.62). In a meta-analysis combining data from studies with a variety of different diets, weak to moderate correlations between CH₄ yield and intensity and MFA were detected (maximum $r = 0.36$ vs. -0.56 ; van Lingen et al., 2014). This suggests that direction and strength of

correlations between individual MFA and CH₄ emission parameters can be explained by the interaction of different dietary constituents on MFA patterns and CH₄ production. Thus, when MFA data derived from different diets were combined to determine the overall relationship between individual MFA and CH₄ production, the average correlation was weak to moderate. In general, our results confirm earlier observations that correlations between CH₄ production and some of the *de novo*-synthesized MFA and groups of SFA are positive, but negative for C18:1 *cis* isomers, UFA, MUFA, and n-3 FA (Chilliard et al., 2009; Mohammed et al., 2011; van Lingen et al., 2014).

When comparing CH₄ prediction among the tested diets, the coefficients of determination were lower for GS diets compared with CS diets. The effect of basal diet (GS and CS) on the CH₄ prediction potential was likely limited in our study because of few differences in MFA concentrations between basal rations, as discussed above. The prediction equations based on MFA data of diets containing linseed (LS vs. L0) produced 15 to 23% lower R^2_{CV} values compared with data from diets with no linseed, irrespective of the data set used. That linseed supplementation deteriorates CH₄ prediction confirms earlier observations (Williams et al., 2014; Dijkstra et al., 2016; Rico et al., 2016). Williams et al. (2014) concluded that the CH₄ prediction model by Chilliard et al. (2009) is a result of linseed oil supplementation and cannot accurately predict CH₄ when cows are fed other diets. Furthermore, Dijkstra et al. (2016) adopted the CH₄ prediction equations by van Lingen et al. (2014) and concluded that MFA profile and CH₄ yield of cows fed grass- or grass silage-based diets differ from those fed other diet types, especially for diets containing fat additives. In this context, Rico et al. (2016) pointed out that interaction effects of basal rations and linseed supplementation may influence prediction equations. Thus, as outlined above, we concluded that the high intake of n-3 FA affected interrelated biochemical pathways on several tissue and metabolic levels but in opposite directions for CH₄ production and MFA concentrations. Furthermore, we cannot exclude that tissue FA synthesis, desaturation, and elongation, which are unrelated to changes in rumen fermentation and thus CH₄ production, play a more dominant role when higher amounts of n-3 FA were fed.

Methane Prediction Models Using Full and Reduced Data Sets for MFA Variables

Our second objective was to compare CH₄ prediction models based on differently sized and composed data sets of MFA. Therefore, we compared CH₄ prediction

equations based on all 43 quantified MFA (full set; data sets 1 to 3) to equations constructed from a reduced number of MFA variables containing groups of biochemically related MFA (data sets 4 to 6). In previous studies it has been shown that the selected MFA groups as well as the individual MFA are positively (SFA, C16:0) or negatively (UFA, MUFA, PUFA, n-3 MFA, C18:0) associated with CH₄ emission (Castro-Montoya et al., 2016a; van Gastelen and Dijkstra, 2016). Models based on all 43 MFA showed a better prediction than those based on the reduced number of MFA group variables. In other studies, the prediction for CH₄ emission based on 24 to 83 MFA variables ranged between coefficients of determination of 0.47 and 0.95 (Chilliard et al., 2009; van Lingen et al., 2014; Rico et al., 2016), which is in the same range as with our equations, except for the GS diets in data sets 1 and 4.

The MFA entered in the equations as explanatory variables using data sets 1 to 3 differed within the data sets. Explanatory individual MFA frequently (4 to 7 times) occurring in our prediction equations were also found in other CH₄ prediction equations [i.e., C17:1 *cis*-9 (Mohammed et al., 2011; Castro-Montoya et al., 2016b; Rico et al., 2016), C18:1 *cis*-9 (van Lingen et al., 2014), and C24:0 (van Gastelen et al., 2017)]. van Gastelen and Dijkstra (2016) pointed out that only C17:1 *cis*-9 and C18:1 *cis*-11 appeared in several published equations, but in our study C18:1 *cis*-11 did not play a major role. When ECM or DMI were additionally included, C18:1 *cis*-11 was not considered. The explanatory MFA variable C17:1 *cis*-9 is derived from rumen microbial membrane lipids or from propionate (Vlaeminck et al., 2006) and is negatively associated with CH₄ production (g/d; Mohammed et al., 2011; Castro-Montoya et al., 2016b; Rico et al., 2016). In contrast, C18:1 *cis*-9 is derived directly from plant feed or is a desaturation product of C18:0, one of the major products when diets rich in forage and plant oils are fed (Alves et al., 2013; Meignan et al., 2017). Long-chain MFA ($\geq$ C20) are scarcely reported because they seem to be less frequently analyzed in CH₄-prediction studies (Dijkstra et al., 2011; van Lingen et al., 2014). Only a few publications reported long-chain MFA (C20:1 *cis*-9, C20:1 *cis*-11, C20:4n-3, C22:0, C22:6n-3, C24:0) as explanatory variables for CH₄ prediction (Mohammed et al., 2011; Castro-Montoya et al., 2016b; van Gastelen et al., 2017). Models based on a reduced number of MFA variables using concentration sums of groups of biochemically related MFA showed MUFA as dominating explanatory variable. The group of MUFA includes biohydrogenation intermediates and was therefore reported to be negatively associated with CH₄ production

(Mohammed et al., 2011; Castro-Montoya et al., 2016a; Vanrobays et al., 2016). Although SFA could not be identified as important explanatory variable for CH₄ production in our study, Weill et al. (2009) proposed SFA <C16 as predictors for CH₄ intensity.

Additional Inclusion of ECM and DMI in Prediction Models

Recently, it was suggested that the combination of MFA data sets with other CH₄ proxies could improve CH₄ prediction (van Gastelen and Dijkstra, 2016; Negussie et al., 2017). Rico et al. (2016) predicted CH₄ production (g/d) for cows with models including diet components, MFA, and DMI as variables in the data sets, but it turned out that in the best-fit model diet components as explanatory variables did not play a role; this indicates that the inclusion of DMI or ECM could be more meaningful as explanatory variables for the prediction models than dietary composition. The best equation reported by Chilliard et al. (2009) also included forage DMI as a parameter, indicating its potential importance. Thus, our third objective was to test the effect of additional inclusion of ECM and DMI. Dry matter intake is the major determinant for CH₄ production (Knapp et al., 2014) and ECM can reflect DMI (Hristov et al., 2013). Inclusion of ECM or DMI improved CH₄-prediction equations, resulting in an increase of the R²_{CV} by approximately 30% when based on pooled data from all diets. This was only true for the data set 5 and 6 with a reduced number of MFA variables, showing that CH₄ prediction by the complete set of 43 MFA variables could explain more variability of CH₄ production than DMI or ECM. van Gastelen et al. (2017) reported improved prediction of CH₄ production by 29% when using equations containing MFA plus volatile and nonvolatile metabolites of milk as compared with equations based on 42 MFA variables alone, indicating that accuracy of prediction can be gained from inclusion of CH₄ proxies other than only MFA as explanatory variables. Interestingly, with MFA data based on the GS diet with addition of ECM (data sets 2 and 5) or DMI (data sets 3 and 6) as a variable excluded MFA as explanatory variable from the equation. This might explain why the predictions for GS diets resulted in the lowest-quality parameters, because GS diets combine feedstuffs and nutrients with opposite effects on CH₄ production (Maia et al., 2007; Shingfield et al., 2013; Castro-Montoya et al., 2016a). In contrast, based on data of the LS diets, inclusion of ECM (data sets 2 and 5) did not improve the prediction as compared with predictions with data sets 1 and 4,

respectively, containing MFA variables only, suggesting a dominant effect of linseed (i.e., n-3 MFA) on the MFA pattern and thus CH₄ prediction.

CONCLUSIONS

Our study shows that the effect of the 2 basal diets containing either a large proportion of grass silage or corn silage on CH₄ production and MFA profile was small due to only moderate intake differences in crude fiber, ADF, and NDF, nutrients known to have a large effect on both CH₄ production and MFA profile. However, with CS diets DMI was higher and resulted in a tendency for higher CH₄ production. In contrast, the supplementation of the basal diets with linseed strongly reduced CH₄ emission parameters, and modified the MFA profile considerably. The developed CH₄ prediction models for dairy cows based on a data set of 43 individual MFA provided good prediction (R^2_{Adj} 0.32 to 0.89; R^2_{CV} 0.47 to 0.92), which was reduced by 10 to 15% for models based on a reduced data set with 9 MFA parameters (biochemically related MFA groups and C16:0 and C18:0 MFA). Prediction of CH₄ production based on a reduced data set can be improved when ECM or DMI are added as variables, indicating a gain of prediction accuracy when combining CH₄ proxies. In this study, prediction equations based on GS diets resulted in the lowest prediction accuracy, which might be due to fiber and starch as TMR ingredients in our GS diets with opposite effects on methane production. Prediction equations based on diets with linseed supplements had a lower prediction accuracy compared with equations based on data from diets without linseed. Therefore, caution is needed when prediction equations are derived from data sets including fat-supplemented diets or when diet-specific prediction equations are applied to cows fed nonmatching diets.

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