Methane Production by Mixed Ruminal Cultures Incubated in Dual-Flow Fermentors

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Methane Production by Mixed Ruminal Cultures Incubated in Dual-Flow Fermentors

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ABSTRACT

This study evaluated the effects of dilution rate and forage-to-concentrate ratio on gas production by rumen microbes. Continuous cultures were used to monitor methane production at three liquid dilution rates (3.2, 6.3, or 12.5%/h) and three forage-to-concentrate ratios (70:30, 50:50, or 30:70). Filtered ruminal contents were allowed 6 d of adaptation to diets followed by 7 d of data collection. Forage consisted of pelleted alfalfa and the concentrate mix included ground corn, soybean meal, and a mineral and vitamin premix. The experiment was replicated in a split-plot design. Total volatile fatty acid production averaged 58.0 mmol/d and was not affected by treatment. Molar proportion of acetate increased with increasing forage-to-concentrate ratio. Molar proportion of propionate tended to decrease at dilution rate of 12.5%/h and increased with the medium and low forage-to-concentrate ratio. Culture pH tended to be greater at a dilution rate of 12.5%/h. Methane production that was calculated from stoichiometric equations was not affected by treatments. However, methane production based on methane concentration in fermentor headspace resulted in an interaction effect of treatments. Stoichiometric equations underestimated methane output at higher dilution rates and with high forage diets. Total diet fermentability was lowest at dilution rate of 3.2%/h. Increasing dilution rates increased microbial yield; increasing the proportion of concentrate improved microbial efficiency. Dilution rate and forage-to-concentrate ratio altered the partition of substrate by microbes. Methane production based on actual concentrations differed from values estimated using stoichiometry of end-product appearance.

(Key words: continuous culture, methane production, stoichiometric equation, microbial energetics)

INTRODUCTION

Volatile fatty acids, CO2, and methane (CH4) are the major end products of anaerobic fermentation of feeds in the rumen. Stoichiometric equations relating substrate degradation to VFA and gas production have been developed and are commonly used to estimate digestibility of ruminant feeds (Wolin, 1960; Russell and Baldwin, 1979; Menke et al., 1979; Van Soest, 1994). Total gas production is increasingly used as a measure of the fermentation potential of feeds (Beuvink and Spoelstra, 1992; Pell and Schofield, 1993). Because the composition of the end products influences the amount of gas produced, a close relationship between the two has been reported (Naga and Harmeyer, 1975; Taya et al., 1980; Menke and Steingass, 1988). However, rate of fermentation can vary with carbohydrate fraction; rapidly fermenting carbohydrates may not always result in larger amounts of total gas production (Beuvink and Spoelstra, 1992). Also, total gas yield can vary considerably due to the incorporation of carbons into microbial mass as well as the different metabolic pathways by which carbohydrate fractions can be degraded by rumen microbes (Krishnamoorthy et al., 1991; Beuvink and Spoelstra, 1992; Van Soest, 1994). The relationship between microbial biomass and gas yield can vary also with growth conditions (Krishnamoorthy et al., 1991; Blümmel et al., 1997). With short-duration incubation studies, fermentation rate as measured by gas production has been used as an index of net growth yield of rumen microorganisms (El-Din and El-Shazly, 1969). Extending the period of incubation reduced net growth of microorganisms even though gas production continued to increase (Raab, 1980). Increased lysis of microbial cells as a consequence of substrate exhaustion and uncoupled fermentation may contribute to reduced net growth at longer incubation times (Van Nevel and Demeyer, 1977). Changes in mi-
microbial yield or metabolism may alter the relationship between substrate digestion and gas production and could affect the estimation of digestion rate from gas measurements (Doane et al., 1997). Previous studies do not make any distinction between CO2 and CH4 and assume that the latter arises directly from the former (Beuvink and Spoelstra, 1992; Theodorou et al., 1998). There is evidence to suggest that more than one fermentation metabolite may be used for the production of CH4 (Miller, 1995). Methanogens have been shown to have variable affinities for hydrogen and are capable of altering their growth rates depending upon the availability of hydrogen and CO2 (Morgan et al., 1997). No study has compared actual CH4 production in mixed ruminal cultures to estimates based on stoichiometric equations. In addition, the generation of microbial biomass can be a sink for reducing equivalents and, if ignored, it can result in the overprediction of CH4 output (Wolin, 1960; Hungate, 1967).

The objective of the present study was to determine CH4 production by mixed ruminal microbes under different dilution rates and dietary treatments. Direct measurements of CH4 concentration in fermentor headspace were compared with estimates of CH4 production based on stoichiometric equations.

**MATERIALS AND METHODS**

**Incubation Conditions**

A mature lactating Holstein cow fitted with a rumen cannula was fed a diet consisting of 63% roughage and 37% concentrate. Grab samples of ruminal contents were taken from various sites within the reticulo-rumen, filtered through double-layered cheesecloth, and transported to the laboratory in sealed, preheated containers. In the laboratory, contents from the containers were filtered again through double-layered cheesecloth into a large, wide-mouth beaker and mixed thoroughly before pouring into the fermentors. The preparation time of the ruminal contents in the laboratory did not exceed 15 min. Approximately 700 ml of the strained ruminal fluid was transferred into each of three fermentors with a continuous dual-flow system (Teather and Sauer, 1988). The design of the fermentors allows for natural stratification of contents similar to the way it occurs in the rumen. The stratification of the fermentor contents is accomplished with the presence of a glass “T” that serves as the fermentor overflow. The entrance to the outlet “T” is near the bottom of the suspended particle phase, which allowed for differential liquid and solid turnover rates. The presence and maintenance of the “mat” in the cultures and the relatively slower rate at which culture contents are stirred (10 to 12 rpm) in these fermentors allow protozoal populations to remain in cultures over extended periods. We did not directly measure protozoal populations in the present study. However, protozoa were visually present in the fermentors for the duration of the experiment. Following inoculation, protozoa were consistently discernible as large white bands within the fermentors. Over a 1- to 2-d period, the protozoa tended to localize either at the base or the bottom of the overflow port, areas that seemed to have the slowest turnover rate. The close association between the methanogens and protozoa and the sustained production of methane at high levels in these fermentors for extended periods of operation also suggests that protozoa are maintained during the duration of the study. Several hours prior to the addition of the ruminal fluid, the system was purged with CO2 gas. To displace O2 and maintain anaerobic conditions in the vessels, the rate of CO2 flow through the fermentors was fixed at 20 mL/min throughout the experiments.

A circulating water bath was used to maintain the temperature of the fermentors at 39°C. Continuous stirring of fermentor contents was achieved with the aid of a central paddle set at a speed of 10 rpm. Artificial saliva was prepared as described by Slyter et al. (1966) and delivered continuously at 0.73 mL/min. Liquid turnover rate was increased or decreased by adjusting the saliva flow rate as described below.

**Dietary Treatments and Dilution Rates**

Experimental diets consisted of 3 forage-to-concentrate (F:C) ratios: high forage (HF), 70:30; medium forage (MF), 50:50; and low forage (LF), 30:70. The forage comprised 100% pelleted alfalfa (16.5% CP) and the concentrate (13.1% CP) consisted of 81.4% ground corn, 10.4% soybean meal (48% CP), 2.1% bentonite, 1.5% sodium bicarbonate, 1.3% phosphate, 1.1% lime-stone, 1.0% soybean oil, 1.0% salt, and 0.2% vitamin-mineral premix. The chemical composition of dietary treatments is as follows: HF: 31.8% NDF, 15.5% CP, 67.8% TDN; MF: 25.2% NDF, 14.8% CP, 72.8% TDN, and LF: 18.6% NDF, 14.1% CP, and 77.7% TDN. The digestible energy content of the diets was 3.0, 3.2, and 3.4 Mcal/kg and was estimated based on NRC (1989) values. A total of 12.8, 12.9, or 13.0 g DM of the diet was added daily in two equal amounts to the fermentor with HF, MF, or LF diets. All fermentors were stabilized on the HF diet for 2 d (stabilization period). Preliminary results have indicated that microbial VFA, CH4, and pH reach a stable steady-state level within 2 d following the addition of ruminal contents into the fermentors. At the end of d 2, one fermentor was maintained on the HF diet, and the other two received the MF diet for an additional 2 d. On d 5, one of the two fermentors receiving the MF diet was switched to the LF diet and was...
allowed to stabilize for an additional 2 d. By the end of
d 6, all three fermentors had been stabilized for at least
d 2 d on the respective dietary treatments. Data were
collected from d 7 to 13 (treatment period).

The effect of the 3 diets was tested at three different
dilution rates of 0.032, 0.063, and 0.125/h, approximat-
ing 0.8, 1.5, and 3.0 volume turnovers per day. These
dilution rates were chosen to cover the physiological
range of fluid turnover rates that are typically observed
in vivo. Throughout all experiments, fermentors were
allowed to stabilize for 2 d at a saliva flow rate of 0.73
mL/min. Following stabilization, saliva flow rate was
either maintained at 0.73 mL/min resulting in a frac-
tional dilution rate of 6.3%/h or it was adjusted, begin-
ning on d 3, to obtain fractional dilution rate of 3.2 or
12.5%/h (0.37 or 1.46 mL/min, respectively).

Sampling and Analyses

Five milliliters of thoroughly mixed fermentor con-
tents was taken 2 h after the a.m. and p.m. feeding
daily for 7 d and analyzed for VFA by GLC (model CP-
3380; Varian, Walnut Creek, CA) using a fused silica
capillary column (Nukol; Supelco Inc., Bellefonte, PA)
and for ammonia-N (NH₃-N) using a colorimetric assay
(Beecher and Whitten, 1970). Production of NH₃-N was
calculated as:

\[ \text{NH}_3-N \text{ (g/d)} = (\text{NH}_3-N \text{ concentration, mg/ml} \times \text{fermentor volume (700 ml)}) \times \text{turnover rate of fermentor/1,000.} \]

Ten microliters of headspace gas samples from the fer-
mentor was drawn into a gas-tight syringe (Hamilton
Co., Reno, NV) and analyzed for CH₄ by GLC (model
CP-3800; Varian) using a stainless steel column packed
with Molsieve 5A 45/60 mesh (Supelco Inc.). The pH of
the ruminal cultures was monitored continuously and
recorded when samples for CH₄ were taken.

Assuming that carbohydrates (hexoses) are the major
source of VFA in the rumen, partitioning of substrate
use was expressed as the amount of substrate fer-
mented to VFA, gas (CH₄ + CO₂ direct and indirect),
or microbial biomass. The direct source of CO₂ is the
fermentation of glucose by various pathways yielding
VFA, ATP, and CO₂. Because the in vitro methods use
bicarbonate-based buffering solutions, CO₂ is released
into the gas phase as VFA enter the medium (Beuvink
and Spoelstra, 1992). This source of CO₂ is considered
as indirect CO₂ production. The amount of substrate
fermented to VFA, CH₄, and CO₂ was calculated based
on the moles of individual VFA produced, daily methane
output, and CO₂ released from fermentation and buffer
addition (Wolin, 1960; Van Soest, 1994; Blümmel et al.,
1997). Total moles of ATP production were estimated
by assigning 2, 3, and 3, moles of ATP per mol of acetic,
propionic, and butyric acids, respectively, and 1 mol
of ATP per mol of CH₄ (Groot et al., 1998). Microbial
efficiency was reported to average 11.7 g of cells/mol of
ATP and was not affected significantly by the energy
concentration in mixed cultures of ruminal organisms
(Isaacson et al., 1975). However, changing dilution
rates did affect the efficiency of microbial protein syn-
thesis that was reported to be 7.5, 11.6, and 16.7 g of
cells/mol of ATP at fractional dilution rates of 2, 6, and
12%/h, respectively (Isaacson et al., 1975). The dilution
rates tested in the present study were similar to those
used by Isaacson et al. (1975) and, therefore, the yield
of microbial biomass per mol ATP (Y_ATP) was set to be
7.5, 11.6, and 16.7 mg/mmol ATP for dilution rates of
3.2, 6.3, and 12.5%/h, respectively. Also assumed was
that 80% of bacterial components were synthesized
from glucose skeletons (Groot et al., 1998). Conse-
quently, microbial biomass from glucose consumption
was calculated as:

\[ \text{Microbial biomass (g/d)} = 0.8 \text{Y_ATP (2 Acetate, mmol/d} + 3 \text{Propionate, mmol/d} + 3 \text{Butyrate, mmol/d} + \text{CH}_4, \text{ mmol/d)/1,000} \]

Energy contents of acetate, propionate, butyrate, valer-
ate, isobutyrate, and isovalerate were used to estimate
digestible energy. Similarly, energy content of CH₄ was
used to estimate contribution of energy in CH₄ to total
digestible energy. In addition to direct measurement of
CH₄, the production of fermentation gases, CO₂ and
CH₄, and the associated production of H₂O, was calcu-
lated using the equation outlined by Wolin (1960) and
Blümmel et al. (1997).

Experimental Design and Statistical Analyses

Within a run, an experimental period lasted 13 d,
which included 6 d for adaptation followed by 7 d for
data collection. Daily values were averaged across the
7 d within each run. A single run was composed of 3
fermentors that were inoculated simultaneously with
ruminal contents obtained from the same cow. Each
fermentor was randomly assigned to one of 3 diets with
different F:C ratios. Each run was replicated (n = 2) at
each of the three dilution rates.

Data were analyzed using the GLM procedure of SAS
(SAS Inst. Inc., Cary, NC). Split-plot design was used
with dilution rate as whole plot and F:C ratio as subplot.
The model used is described by:

\[ Y_{ijk} = \mu + DR_i + \text{Run}_j (DR_i) + FC_k + DR_i \times FC_k + e_{ijk} \]
Table 1. Concentration and production of VFA as affected by dilution rate (DR) and forage-to-concentrate ratio (F:C).1

<table>
<thead>
<tr>
<th>VFA</th>
<th>DR, %/h</th>
<th>HF</th>
<th>MF</th>
<th>LF</th>
<th>SE</th>
<th>Significance of effect</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>3.2</td>
<td>6.3</td>
<td>12.5</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total, mM</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Individual, mol/100 mol</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Acetate (A)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Propionate (P)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Butyrate</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Valerate</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Isobutyrate</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Isovalerate</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>A:P ratio</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total, mmol/d</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Individual, mmol/d</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

1HF = High forage (70% forage:30% concentrate); MF = medium forage (50% forage:50% concentrate); LF = low forage (30% forage:70% concentrate).

2DR \times F:C = Interaction between dilution rate and F:C ratio.

3Production parameters are based on 1 L of ruminal cultures.

4NS = Not significant (P > 0.10).

where

\[ Y_{ijk} = \text{individual response variable measured,} \]
\[ \mu = \text{overall mean,} \]
\[ \text{DR}_i = \text{fixed effect of dilution rate (i = 1 to 3),} \]
\[ \text{Run}_i (\text{DR}_i) = \text{whole plot error,} \]
\[ \text{FC}_k = \text{fixed effect of F:C ratio (k = 1 to 3),} \]
\[ \text{DR}_i \times \text{FC}_k = \text{fixed effect of interaction between dilution rate and F:C ratio, and} \]
\[ e_{ijk} = \text{subplot error.} \]

Dilution rates were compared with the whole plot error term. Forage-to-concentrate ratios and interaction between dilution rate and F:C ratio were tested using the subplot error term. Standard errors appropriate for comparisons among different means were calculated as follows (Steel et al., 1997):

\[ \text{for comparing dilution rate:} \]
\[ \text{SE (}\overline{y}_{1.} - \overline{y}_{2.}\text{)} = \sqrt{\frac{2 \cdot \text{MSE}_{\text{whole plot}}}{6}}, \]

\[ \text{for comparing F:C ratio:} \]
\[ \text{SE (}\overline{y}_{1.} - \overline{y}_{2.}\text{)} = \sqrt{\frac{2 \cdot \text{MSE}_{\text{subplot}}}{6}}. \]

Comparison of dilution rate and F:C ratio means was done by contrast test with Fisher’s protected LSD test when the effect of dilution rate or F:C ratio (P ≤ 0.10) was detected by the model. The level of significance accepted was P ≤ 0.05 and 0.10 for trend.

RESULTS

Total VFA concentrations (millimolar) and molar percentages of individual VFA are reported in Table 1. Increasing the dilution rate reduced (P < 0.01) the concentration of total VFA. Molar proportion of acetate ranged from 49 to 56 and was not affected by dilution rate. Propionate remained unchanged (P > 0.10) when dilution rate increased from 3.2 to 6.3%/h but tended to decrease (P < 0.07) at 12.5%/h. The acetate-to-propionate (A:P) ratio increased (P < 0.01) with increasing the dilution rate from 3.2 or 6.3 to 12.5%/h. Increasing the proportion of concentrate in the diet decreased (P < 0.01) the molar proportion of acetate. Molar proportions of propionate were lowest for the HF diet and increased (P < 0.03) for the MF and LF diets. Molar proportions of valerate were similar for the HF and MF diets but increased for the LF diet. The A:P ratio was highest for the HF diet and decreased (P < 0.01) for the LF diet.

There were dilution rate \times F:C interactions for molar percentages of butyrate and isovalerate. At 3.2%/h, bu-
Table 2. Ruminal pH, ammonia-N (NH3-N), and methane (CH4) production as affected by dilution rate (DR) and forage-to-concentrate ratio (F:C).

<table>
<thead>
<tr>
<th>Item²</th>
<th>3.2</th>
<th>6.3</th>
<th>12.5</th>
<th>SE</th>
<th>DR</th>
<th>F:C</th>
<th>DR × F:C³</th>
</tr>
</thead>
<tbody>
<tr>
<td>DR, %/h</td>
<td>HF</td>
<td>MF</td>
<td>LF</td>
<td>HF</td>
<td>MF</td>
<td>LF</td>
<td>HF</td>
</tr>
<tr>
<td>pH</td>
<td>5.3</td>
<td>5.0</td>
<td>5.0</td>
<td>5.7</td>
<td>5.5</td>
<td>5.6</td>
<td>6.7</td>
</tr>
<tr>
<td>NH3-N, mg/dl</td>
<td>15.6</td>
<td>14.7</td>
<td>12.1</td>
<td>27.3</td>
<td>31.1</td>
<td>25.5</td>
<td>18.2</td>
</tr>
<tr>
<td>NH3-N, g/d</td>
<td>88.5</td>
<td>89.4</td>
<td>104.2</td>
<td>102.8</td>
<td>109.7</td>
<td>99.9</td>
<td>106.5</td>
</tr>
<tr>
<td>Gas estimated, mmol/d</td>
<td>88.5</td>
<td>89.4</td>
<td>104.2</td>
<td>102.8</td>
<td>109.7</td>
<td>99.9</td>
<td>106.5</td>
</tr>
<tr>
<td>CH4 measured</td>
<td>13.1</td>
<td>13.1</td>
<td>13.4</td>
<td>16.1</td>
<td>14.9</td>
<td>12.1</td>
<td>19.1</td>
</tr>
<tr>
<td>NH3-N, g/d</td>
<td>9.6</td>
<td>6.1</td>
<td>4.2</td>
<td>25.5</td>
<td>19.6</td>
<td>11.2</td>
<td>29.1</td>
</tr>
<tr>
<td>mmol/g DM fed</td>
<td>0.8</td>
<td>0.5</td>
<td>0.3</td>
<td>2.0</td>
<td>1.5</td>
<td>0.9</td>
<td>2.3</td>
</tr>
</tbody>
</table>

¹HF = High forage (70% forage:30% concentrate); MF = medium forage (50% forage:50% concentrate); LF = low forage (30% forage:70% concentrate).
²Production parameters are based on 1 L of ruminal cultures.
³DR × F:C = Interaction between dilution rate and F:C ratio.
⁴NS = Not significant (P > 0.10).
⁵NH3-N (g/d) = (NH3-N concentration, mg/dl × fermentor volume (700 ml) × turnover rate of fermentor)/1,000. Turnover rate of fermentor is 0.8, 1.5, and 3.0 for 3.2, 6.3, and 12.5%/h, respectively.
⁶Fermentative CO2 + fermentative CH4 + buffering CO2. All gas productions were estimated.
⁷(Acetate, mmol/d) + (2 × butyrate, mmol/d) − (CO2, mmol/d).
Table 3. Amount of substrate used for fermentation end products and microbial growth as affected by dilution rate (DR) and forage-to-concentrate ratio (F:C).1

<table>
<thead>
<tr>
<th>Item</th>
<th>DR, %/h</th>
<th>3.2</th>
<th>6.3</th>
<th>12.5</th>
<th>Significance of effect</th>
</tr>
</thead>
<tbody>
<tr>
<td>DM fed, g/d</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>HF</td>
<td>12.8</td>
<td>12.9</td>
<td>13.0</td>
<td></td>
<td></td>
</tr>
<tr>
<td>MF</td>
<td></td>
<td></td>
<td></td>
<td></td>
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</tr>
<tr>
<td>LF</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Substrate used, g/d</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>For VFA4</td>
<td>3.3</td>
<td>3.4</td>
<td>4.2</td>
<td></td>
<td></td>
</tr>
<tr>
<td>For gas5</td>
<td>1.9</td>
<td>1.8</td>
<td>2.0</td>
<td></td>
<td></td>
</tr>
<tr>
<td>For microbial biomass6</td>
<td>1.0</td>
<td>1.0</td>
<td>1.2</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total7</td>
<td>6.1</td>
<td>6.2</td>
<td>7.5</td>
<td></td>
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</tr>
<tr>
<td>Fermentability, %8</td>
<td>48.0</td>
<td>48.4</td>
<td>57.3</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Microbial synthesis9</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>g/d</td>
<td>0.79</td>
<td>0.81</td>
<td>0.98</td>
<td></td>
<td></td>
</tr>
<tr>
<td>g DM/kg DM fermented10</td>
<td>132.7</td>
<td>133.5</td>
<td>136.4</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

1HF = High forage (70% forage:30% concentrate); MF = medium forage (50% forage:50% concentrate); LF = low forage (30% forage:70% concentrate).
2DR × F:C = Interaction between dilution rate and F:C ratio.
3NS = Not significant (P > 0.10).
4(Acetate, mol/d × 60.05) + (Propionate, mol/d × 74.08) + (butyrate, mol/d × 88.10).
5Substrate used for (CO2, mol/d × 44) + (CH4, mol/d × 16) + (2H2O, mol/d × 36).
6Substrate used for microbial biomass = ATP (mmol) × YATP (7.5, 11.6, and 16.7 mg for 3.2, 6.3, and 12.5%/h, respectively).
7Substrate used for VFA, CO2 + C H4 + 2 H2O, and microbial biomass.
8Total substrate fermented expressed as a percentage of DM fed.
9Microbial biomass (g/d) = (((0.8 × YATP ((2 × Acetate, mmol/d) + (3 × propionate, mmol/d) + (3 × butyrate, mmol/d) + (CH4, mmol/d))/1000). YATP is 7.5, 11.6, and 16.7 mg for 3.2, 6.3, and 12.5%/h, respectively.
10Microbial growth efficiency (g DM/kg DM fermented) = ((microbial biomass, g/DM fermented, g) × 1000).

the dilution rate to 12.5%/h did not increase methane output for the MF diet, but there was a substantial increase for the LF diet, which was numerically higher compared with the MF diet. When expressed as mmol of methane produced per gram of DM fed, the effects of dilution rate and F:C were similar to the total daily production rates described above.

The amount of substrate used for gas (CH4 + direct and indirect CO2) was not affected (P > 0.10) by either dilution rate or F:C (Table 3). The amount of substrate used for microbial biomass was 1.07 g/d at the lowest dilution rate and increased (P < 0.01) to 1.83 and 2.46 g/d at dilution rate of 6.3 and 12.5%/h, respectively (Table 3). Based on the substrate used for VFA, gas, and microbial biomass, total diet fermentability was similar (P > 0.10) at dilution rates of 6.3 and 12.5%/h, respectively (Table 3). Microbial yields increased (P < 0.01) with increasing dilution rate and averaged 0.86, 1.46, and 1.97 g/d at dilution rates of 3.2, 6.3, and 12.5%/h, respectively (Table 3). Increasing the proportion of concentrate in the diet did not affect microbial yield (P > 0.10) but improved (P < 0.02) microbial growth efficiency (Table 3). There was an interaction between dilution rate and F:C for microbial efficiency. At a dilution rate of 6.3%/h, microbial growth efficiency in cultures receiving LF diet was numerically higher than cultures receiving MF diet, but increasing dilution rate to 12.5%/h resulted in numerically higher efficiency of microbial growth in fermentors receiving MF diet compared with those receiving the LF diet.

The amount of energy produced daily in the form of VFA and as a percentage of digestible energy fed was not affected by dilution rate (Table 4). A higher proportion of concentrate tended to increase (P < 0.08) the amount of energy (kcal/d) captured in VFA due primarily to the increased digestible energy fed. The VFA energy as a percentage of digestible energy fed was not affected by F:C ratio. There was an interaction for the amount of energy released in CH4 (P < 0.01). At dilution rates of 3.2 and 6.3%/h, cultures receiving the LF diet had the lowest rate of methane production; however, increasing dilution rate to 12.5%/h resulted in the MF diet having the lowest methane output.

**DISCUSSION**

Altering liquid turnover rates had no effect on daily production of total VFA or in the proportion and production of acetate by ruminal cultures. These results are similar to those reported by Isaacson et al. (1975) and Hoover et al. (1984). Molar proportion of propionate
Table 4. Amount of digestible energy (DE) partitioned into VFA and methane (CH4) obtained by actual measurement as affected by dilution rate (DR) and forage-to-concentrate ratio (F:C).1

<table>
<thead>
<tr>
<th>Item</th>
<th>DR, %/h</th>
<th>Significance of effect</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>3.2</td>
<td>6.3</td>
</tr>
<tr>
<td>DE fed, kcal/d</td>
<td></td>
<td></td>
</tr>
<tr>
<td>VFA kcal/d4</td>
<td>16.6</td>
<td>17.1</td>
</tr>
<tr>
<td>% of DE</td>
<td>43.6</td>
<td>41.6</td>
</tr>
<tr>
<td>CH4 kcal/d5</td>
<td>2.0</td>
<td>1.3</td>
</tr>
<tr>
<td>% of DE</td>
<td>5.3</td>
<td>3.2</td>
</tr>
</tbody>
</table>

1HF = High forage (70% forage:30% concentrate); MF = medium forage (50% forage:50% concentrate); LF = low forage (30% forage:70% concentrate).
2DR × F:C = Interaction between dilution rate and F:C ratio.
3NS = Not significant (P > 0.10).
4(Acetate, mol/d × 209.4 kcal/mol) + (propionate, mol/d × 367.2 kcal/mol) + (butyrate, mol/d × 524.3 kcal/mol) + (valerate, mol/d × 681.6 kcal/mol) + (isobutyrate, mol/d × 524.3 kcal/mol) + (isovalerate, mol/d × 681.6 kcal/mol).
5CH4 (kcal/d) = (CH4, mol/d × 210.8 kcal/mol).

tended to decrease and that of butyrate increased with an increase in dilution rate, but production of both fatty acids remained unchanged. Ruminal pH and ruminal methane output increased with increasing dilution rate. The effect of dilution rate on VFA production and fermentation profile in other studies has been variable. Total organic acid production decreased (Carro et al., 1995) or increased (Fuchigami et al., 1989) when fractional dilution rates increased. Molar percentages of propionate in ruminal fluid were reported to be higher (Isaacson et al., 1975; Hoover et al., 1984) or lower (Thomson et al., 1978; Crawford et al., 1980) as a result of increasing dilution rates.

In the present study, dilution rate had a significant effect on the isoacids. Rapid liquid turnover lowered the proportion of ruminal valerate but increased that of the branched chain isoacids, isobutyrate, and isovalerate. Part of the valerate produced in the rumen comes from the fermentation of carbohydrates and part comes from the fermentation of AA. The branched-chain isoacids arise almost exclusively from the oxidative deamination of AA. The inability of the cellulolytic bacteria to transport preformed branched chain AA across their cell wall makes the branched chain isoacids essential for normal growth of fiber-digesting bacteria (Bryant, 1973). The decrease in valerate may be due to reduced retention time of the fermentable carbohydrates at higher liquid passage rates. An increase in branched-chain isoacids suggests enhanced deaminative activity at higher dilution rates, since a reduced uptake should have resulted in a lower acetate production.

Increasing liquid turnover increased NH3-N output. Ammonia-N concentration, similar to the isoacids, is a function of the rate of release and rate of uptake by microbial populations. Ruminal pH can also influence NH3-N production (Erflé et al., 1982), and increasing dilution rates increased NH3-N with a concomitant increase in culture pH similar to results reported earlier (Hoover et al., 1984). Ammonia-N and branched-chain isoacids increased at higher dilution rates, suggesting either enhanced rate of production or reduced utilization. Both are used predominantly by cellulolytic organisms and, since acetate production was not affected, higher liquid dilution rates seem to have enhanced rates of production rather than reduced rates of utilization.

Similar values for DM digestibility have been reported earlier with no effect on dilution rate (Hoover et al., 1984; Carro et al., 1995). In the study by Hoover et al. (1984), increasing dilution rates did not seem to reduce NDF digestibility or the digestibility of the more rapidly fermentable carbohydrate fraction. In fact, in some cases, increasing dilution rate seemed to increase cellulose and NDF digestibility (Hoover et al., 1984).

Increasing dilution rate did not affect total organic acid production, but actual methane output increased. In contrast, lowering F:C ratio reduced methane production and increased ruminal propionate as has been documented in several studies. Increasing dilution rates did not affect the amount of substrate used for VFA production but increased the amount used for gas and microbial biomass. Actual methane production increased significantly at each successive increase in dilution rate, but the proportion of substrate used for total gas output (CH4 + CO2) increased only when dilution rates were increased from 3.2 to 12.5%/h. The growth rate of methanogens is relatively slow, which results in reduced numbers during rapid rates of transit from...
the rumen (Wolin et al., 1997). The highest dilution rate in our study (12.5%/h) did not seem to have a negative impact on methane production. Calculated microbial yields and efficiencies increased with an increase in turnover rate suggesting that, at higher turnover rates, a greater proportion of substrate energy is used for bacterial synthesis (Hespell and Bryant, 1979).

According to Wolin (1960), the amount of total gas produced (\(\text{CH}_4 + \text{CO}_2\)) can be determined from the amount and molar proportion of acetate, propionate, and butyrate. Hence, based on the stoichiometric equation, variations in the molar proportion of acetate, propionate, and butyrate will have a direct influence on gas volumes. Blümmel et al. (1997) reported that the total substrate required for the production of equal amounts of gas from widely different VFA patterns was very similar. Accounting for the efficiency of ATP use by microbial populations, the amount of substrate required for microbial biomass can vary without changes in VFA proportions (Blümmel et al., 1997). It is known that the growth yields of ruminal microbes can be relatively high, and that microbial cells have a negative oxidation-reduction state (Van Kessel and Russell, 1996). Since stoichiometric equations used to estimate fermentation balance do not consider microbial cells as end products of ruminal fermentation, cell yields can have a significant impact on estimates of methane production.

The Cornell Net Carbohydrate and Protein System suggests a maximum incorporation of 40% of the fermented feed carbohydrate into microbial biomass (Russell et al., 1992). With substrates consisting predominantly of structural carbohydrates, microbial biomass yield was negatively correlated to gas production over a 24-h period of incubation (Blümmel et al., 1997). In another study (Krishnamoorthy et al., 1991), a curvilinear relationship was reported over a 2-h period between microbial protein synthesis and gas production. The curvature varied with the type of substrate, with cellulose producing a steep upward curve and starch producing a less steep curve. The relationship between microbial mass and gas volume is complex and can vary with the type of substrate as well as with the time of sampling.

In previous experiments, no distinction was made between \(\text{CO}_2\) and \(\text{CH}_4\), and it is assumed that the latter arises directly from the former. As per the stoichiometric equations outlined previously (Wolin, 1960; Blümmel et al., 1997), we included both the direct \(\text{CO}_2\) production from the fermentative route as well as the indirect contribution from the reaction of the VFA with the bicarbonate supplied in the saliva in estimating total gas production. Total gas production was not affected by the F:C ratio and dilution rates in the present study. Methane production, when estimated using stoichiometric equations, was also not altered by either the dilution rates or F:C ratio. However, actual measurements of methane production decreased significantly with an increase in the level of concentrate and increased with an increase in dilution rate. The decrease in methane in cultures receiving a higher proportion of concentrate is consistent with the shift in reducing equivalents toward propionate formation. However, the increased methane formation at higher dilution rates was unexpected.

Gas production and VFA formation are closely related processes, but gas production can vary without any change in total VFA production (Beuvink and Spoelstra, 1992). The amount of gas released indirectly is assumed to be a constant; therefore, the variation is attributed primarily to gas produced directly as an end product of ruminal fermentation. Direct gas production varies with the pattern of VFA and, given the complex nature of mixed feedstuffs and fermentative pathways of microbial metabolism, it can vary considerably. Most species of rumen microbes are capable of fermenting various substrates resulting in similar end products (Hungate, 1966), and some have switched end products depending on their growth rate (Russell and Wallace, 1997).

Hydrogen and \(\text{CO}_2\) are the major precursors of \(\text{CH}_4\) formation in the rumen (Hungate, 1967), and most methanogens can utilize these substrates to generate ATP (Thauer et al., 1977). The distribution of methanogenic species in the rumen is not known and whereas some species can occur in high concentration, others may be present in low concentration (Wolin et al., 1997). Methanogens have a slower growth rate compared with other rumen bacteria, but there are some species, Methanobrevibacter spp. in particular, that will grow more rapidly with \(\text{H}_2\) than other methanogens (Wolin et al., 1997). The success of microorganisms to survive depends on their ability to maximize growth rate or growth yield (Neijssel and de Mattos, 1994; Russell and Wallace, 1997). Culture of Methanobacter thermautotrophicum did not grow when \(\text{H}_2\) supply was low, but they continued to produce methane. At higher availability of \(\text{H}_2\), growth of \(M. \text{thermautotrophicum}\) occurred, but methanogenesis remained constant as long as growth was \(\text{H}_2\) limiting (Morgan et al., 1997). When \(\text{H}_2\) supply was not growth limiting, \(\text{CH}_4\) production increased rapidly (Morgan et al., 1997). Changing dilution rates has a profound impact on the growth rate of bacteria and alters the metabolic pathways of fermentation. Given the complex interactions between microbial growth and fermentation environment, altering dilution rates could interfere with growth by changing the concentration and/or shifting the metabolic pathways.
CONCLUSIONS

Methane production in continuous cultures was measured to determine whether it was altered by dilution rate and F:C ratio. Increasing dilution rate or the level of forage in the diet decreased propionate and increased methane output. Methane production estimated from stoichiometric equations remained unchanged, irrespective of the dilution rate or level of forage in the diet. Compared with actual measurements of methane concentration in fermentor headspace, stoichiometric estimations consistently underestimated methane output at higher dilution rates and with high forage diets. Methane production seems to be influenced by the stoichiometry of substrate fermentation to acetate and propionate as well as the proportion of substrate carbon fixed in microbial biomass.

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