Ozone Decreased Enteric Methane Production by 20% in an *in vitro* Rumen Fermentation System

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Ozone (O₃) is volatile, highly oxidative, and has theoretical potential to reduce ruminant enteric methanogenesis by interactions between archaea and bacteria, and substrate and oxygen. The effects of O₃ on the rumen microbiota, fermentation parameters, and CH₄ emissions were studied through *in vitro* fermentation using a RUSITEC apparatus with O₃ dissolved in the salivary buffer. The substrate consisted of maize silage or grain concentrates, and the treatments were (1) control (no O₃) and (2) O₃ at 0.07 ± 0.022 mg/L in the buffer. A 4-day adaptation period followed by a 6-day experimental period was used for measuring gas production and composition, as well as fermentation characteristics, which included ruminal volatile fatty acids (VFA) and liquid- and solid-associated microbial communities. Ozone treatment decreased total gas production by 15.4%, most notably CH₄ production by 20.4%, and CH₄ gas concentration by 5.8%, without compromising dry matter digestibility (DMD) of either maize silage or grain concentrates. There were no significant effects of O₃ treatment on VFA production or pH. Ozone treatment reduced the relative abundance of methanogens, particularly *Methanomicrobium*. This study demonstrates the potential use of O₃ as a method to reduce ruminant enteric methanogenesis.

**Keywords:** enteric, methanogenesis, rumen, cattle, O₃, CH₄ (methane)

**INTRODUCTION**

Ruminant CH₄ production is notoriously one of the largest contributing factors toward greenhouse gas (GHG) emissions, accounting for 50% of emissions from the agricultural sector in Australia, and 7% to total Australian GHG emissions (Hook et al., 2016; Duarte et al., 2017b). Research toward methods to reduce CH₄ emissions is increasingly valuable with increasing worldwide demand for meat and dairy products; however, minimal progress has actually been achieved (Martin et al., 2010).
Studies have shown that manipulation of feed directly or through additives has been an effective strategy (Beauchemin et al., 2008; Hristov et al., 2013; Teoh et al., 2019) in reducing CH₄ production. The rumen microbiome produces volatile fatty acids (VFA) from otherwise non-digestible plant polysaccharides and these VFA are the main source of energy to ruminants (Dijkstra, 1994; Henderson et al., 2015). Ruminal archaea, namely, the methanogens, are sensitive to changes in their environment, particularly oxygen availability, and produce CH₄ as a metabolic by-product under hypoxic conditions (Issazadeh et al., 2013; Patra et al., 2017). Indeed, Moate et al. (2019) have recently shown that in comparison to non-fistulated cows, rumen-fistulated cows have a lower CH₄ yield, presumably due to inhibition of the methanogenic populations by oxygen.

One common method of disinfecting water is ozonation (also known as O₃ disinfection), using O₃ gas to destroy bacteria and viruses (Remondino and Valdenassi, 2018). Ozone gas (O₃) is formed upon collision of oxygen gas (O₂) with free oxygen atoms. Currently, O₃ is widely used in the food industry (Brodowska et al., 2018); however, it has not yet been considered as a dietary additive to mitigate CH₄ production in ruminants. Methanogens should be inhibited more by O₃ compared to O₂, because they are extremely sensitive to oxygen and O₃ is a more powerful oxidant than O₂ (Wang et al., 2004). The majority of the rumen microbiome is composed of bacteria and protozoa by mass, with archaea accounting for less than 4% of the microbial community (Matthews et al., 2019). When compounds such as hemiacetal of chloral and starch analogs such bromochloro-methane were included in the diets of ruminants, rumen methanogens and CH₄ production were observed to be severely reduced without negative effects on dry matter intake, dry matter digestibility (DMD) or liveweight gain (Trei et al., 1972; Tomkins et al., 2009; Martinez-Fernandez et al., 2018).

Thus, we hypothesized that (1) if O₃ was included in an in vitro ruminal fermentation process, it would decrease CH₄ production and (2) would have a CH₄-inhibiting effect without affecting feed digestibility and other ruminal fermentation parameters. As such, the objective of this study was to evaluate the effect of including O₃ in an in vitro ruminal fermentation system on CH₄ production, DMD, the rumen microbiota, and other fermentation parameters.

**MATERIALS AND METHODS**

This study was performed in accordance with The University of Sydney Animal Ethics Committee (Approved Protocol Number 2015/835). The cow used for ruminal contents was housed at Corstorphine Dairy, The University of Sydney (Camden Farm Dairy, Cobbitty, NSW, Australia, 34° 04′ S; 150° 81′ 69′ E).

**Experimental Design and Treatments**

A feed consisting of maize silage and concentrates was fed through a RUSITEC apparatus mimicking in vitro ruminal fermentation, to investigate the effects of O₃ supplementation on fermentation parameters, the rumen microbiota and methanogen activity ultimately leading to CH₄ production. The experiment consisted of a completely randomized design with two treatments: a control and an O₃-buffered treatment, and three replicates of each over the course of each 10-day experiment (Klevenhusen et al., 2014; Ertl et al., 2015; Sarnataro et al., 2019). The O₃ treatment contained O₃ at 0.07 mg/L that was pumped into McDougall’s buffer and then synthesized into the feed, whereas the control consisted of maize silage and concentrates only.

The study was comprised of 2 × 10-day incubation runs, whereby fermentation parameters (including pH, oxidation–reduction potential, total gas production, CH₄ gas production, and effluent volume production) were measured daily for 6 days following a 4-day adaptation period. These are similar procedures used by Soliva and Hess (2007), Klevenhusen et al. (2014), Ertl et al. (2015), Soliva et al. (2015), and Sarnataro et al. (2019) in Rusitec studies. Effluent samples were collected from days 5–10 for analysis. Solid- and liquid-associated microbes were sampled at days 0, 5, and 10 in each incubation run. DMD was measured on days 6–9 using retained nylon bags. The VFA production was calculated using results from daily (days 0–10) sampling of effluent contents and effluent volume production.

**Preparation of Substrate, O₃, and Rumen Inoculum**

The basis for each feed (control and treatment) consisted of maize silage (50% w/w, 5.5 g DM-basis) and concentrate (“grain”; Table 1) (5.5 g DM-basis for each substrate), weighed into individual porous nylon bags (11 cm × 6.5 cm, pore size 50 μm)—the daily total mass being 11 g (DM basis). Substrate components were derived from Corstorphine farm, The University of Sydney. Details of climate, harvest, processing, and substrate feed analysis are detailed by Teoh et al. (2019).

The ruminal fluid donor, a non-lactating Holstein dairy cow, was fed pasture ad libitum (including oaten hay at 6 kg DM/d). The donor was ruminally fistulated and some of the rumen inoculum was collected on the first day of each experimental run, 3 h post-morning feeding. Using cheesecloth for separation, the solid and liquid portions of the ruminal contents were separated and stored in preheated thermos containers, to be immediately transported to the laboratory (Teoh et al., 2019).

The O₃ was produced by a commercial ozonator (model O3 Series BO3, CSN Global Pty Ltd., Buderim, QLD 4556, Australia) and pumped into the buffer (McDougall’s buffer) using a fish tank aerator (Aqua One Precision 750—3.5 W; 19 kPa). The daily average concentration of O₃ across the experiment was 0.07 mg/L, measured twice daily for each 10-day run with a

| TABLE 1 | Chemical composition of substrates used. |
| % in the DM |
| Crude protein | Neutral detergent fiber | Crude fat | Ash | Non-fibrous carbohydrates |
| Maize silage | 8.4 | 38.3 | 3.79 | 3.23 | 46.3 |
| Concentrate | 15.5 | 20.8 | 2.70 | 8.83 | 52.2 |
RUSITEC Fermentation

For each run, the RUSITEC apparatus was set up with six separate fermentation vessels (800 mL each) submerged in a water bath at 39°C. Three of these vessels were intended for the control, and three for the O3 treatment. On day 1, each vessel contained three smaller, porous nylon bags in a smaller vessel submerged in approximately 780 mL (until the entire vessel was filled) of the collected liquid ruminal fluid. Of the nylon bags, one contained approximately 70 g wet weight of the collected solid ruminal contents used unprocessed to inoculate the system and another two substrate bags as described above. Each fermentation vessel was equipped with an input port for O3-buffered infusion, and an outlet port for the effluent. The fermentation vessels were continuously infused with McDougall’s buffer at a dilution rate of 33 mL/h (control), while the three treatment vessels were continuously infused with the same buffer plus O3. To mimic rumination, electric motors moved the contents of the fermentation vessels up and down so as to mix solid and liquid particles and their associated microbes (Teoh et al., 2019).

The nylon bag containing ruminal solids was allowed to incubate for 24 h, before being removed and replaced with two bags: one containing maize silage and one containing concentrates. On day 1, each vessel contained three nylon bags, and on each subsequent day there were four at any given time—and each of these substrate bags was allowed to remain in the fermentation vessel for 48 h (Teoh et al., 2019). The last two bags that were placed in the vessel at day 9 were not used in DM digestibility calculations on day 10 as they had only undergone 24 h of incubation.

Sample Collection and Analysis

Total Gas and CH4 Production

Air-tight bags (Plastigas, Linde AG, Munchen, Germany) were connected to the effluent output ports of each fermentation vessel. Each day, the total volume of gas produced by each vessel was recorded using a drum-type gas meter (Terry et al., 2018). Each effluent gas bag was manually pressurized to produce a reading on the meter.

Methane (CH4) production was determined daily from day 5 until the end of each 10-day run. To measure daily gas production, 34 mL of gas was collected from each gas bag. Some gases from the previous day’s fermentation may potentially be trapped in the tube connecting the fermentation vessel to the gas sampling bag. To prevent these gases from contaminating the gas samples, the first 17 mL from each sample was evacuated, and the remaining 17 mL used for analysis. This retained 17 mL gas sample was transferred into a 10 mL evacuated extainer (Labco Ltd., High Wycombe, United Kingdom). A 3 mL subsample was taken from the extainer and CH4 percentage was determined by gas chromatography (GC; Agilent model 7890A). The GC conditions used were as described by Teoh et al. (2019). Methane production was calculated by multiplying the total gas volume by the percentage of CH4, with correction for temperature and pressure (Duarte et al., 2017b). Results were expressed as mg CH4/g DM.

Dry Matter Digestibility (DMD), pH, Oxidation–Reduction Potential

Dry matter digestibility was calculated using the contents of the nylon bags (maize silage and concentrates) that had been fermenting for 48 h. The collection of these bags took place on days 6–9, when DNA extraction was not required. Each of the bags was washed in a washing machine set to cold-delicate for 30 min, before being dried at 60°C until a constant weight was obtained (approximately 48 h). The residue weight was recorded and used in the calculation of DMD which was expressed as a percentage (%) of the original DM substrate.

The pH of each vessel was measured daily (during nylon bag exchange) using a pH meter (TPS pH-mV-Temp Meter, Model WP-80) calibrated at 39°C. Redox potential was also measured during this time using an oxidation–reduction potential meter calibrated at 39°C (MW 500 ORP meter range of ±999 mV, accuracy of ±5 mV, Milwaukee Instruments, Inc., Rocky Mount, NC, United States; Ramos et al., 2018).

Effluent Volume and Volatile Fatty Acids (VFA) Production

Effluent volume production was measured daily using 2 L glass flasks, connected to the effluent output of each fermentation vessel and submerged in ice to halt fermentation and microbial growth. Daily effluent was measured using a measuring cylinder and expressed as mL/day. From days 5 to 10, a 2 mL sample of effluent contents was collected for determination of concentrations of VFA. This sample was centrifuged (13,500 × g for 2 min at 5°C) and a 1.5 mL subsample of the supernatant was transferred into a 2 mL centrifuge tube and acidified with 0.3 mL metaphosphoric acid (0.20; wt/v). The subsample was then frozen at −20°C until it was analyzed for VFA concentrations using GC (Agilent model 7820A). Crotonic acid was used as internal standard. The settings used were FID set up at 250°C, air flow 300 mL per min, makeup flow (N2) ran at 30 mL per min with a capillary column (DB-FFAP, 30 m × 0.32 mm ID × 1 µm). Helium was used as a carrier gas with a flow rate of 30 mL per min. The split inlet was heated to 225°C, and set at 9.526 PSI, with He constant flow 1.5 mL per min, and split ratio 50:1. The oven temperature was programmed to 150°C (held at that temperature for 1 min) and then ramped at 5°C per min to 195°C (held 3 min). Total and individual VFA were estimated daily by multiplying VFA concentration by the volume of effluent.

Sequencing and Analysis of the Archaeal and Bacterial 16S rRNA Gene

The V4 hypervariable region of archaea and bacteria was sequenced as previously described (Terry et al., 2018; Teoh et al., 2019) using an Illumina MiSeq instrument and the MiSeq Reagent Kit with 500 cycles (Illumina, Inc., San Diego, CA, United States) according to manufacturer’s instructions. DADA2 v. 1.12.1 (Callahan et al., 2016) in R v. 3.6.1 was used to process and quality-filter all sequences. Briefly, the forward and reverse reads were trimmed to 220 and 200 bp, respectively, merged
with a minimum overlap of 75 bp, and chimeras removed. Taxonomy was assigned to the remaining sequences, referred to here as operational taxonomic units (OTUs) at 100% similarity, using the RDP naïve Bayesian classifier and the SILVA SSU database release 132 (Quast et al., 2012). The OTUs classified as chloroplasts, mitochondria, or eukaryotic in origin were removed prior to analyses.

The number of OTUs per sample, Shannon diversity, and inverse Simpson’s diversity indices were calculated in R using Phyloseq 1.28.0 (McMurdie and Holmes, 2013). The bacterial structure was assessed using Bray–Curtis dissimilarities which were calculated with vegan 2.5.6 and the effect of O3 treatment was determined using PERMANOVA (adonis2 function) in R with 10,000 permutations. The LAM and SAM samples were assessed separately as they are quite different from each other in terms of microbial diversity and structure (Duarte et al., 2017b).

All 16S rRNA gene sequences were submitted to the Sequence Read Archive under BioProject number PRJNA590488.

Statistical Analysis

Fermentation data were analyzed using the MIXED procedure of SAS (SAS, Inc., 2020; SAS Online Doc 9.1.4). The model included the fixed effects of O3, day, and O3 × day interaction. Therefore, the individual fermenter was used as the experimental unit for statistical analysis. The minimum values of Akaike's information criterion were used to select the covariance structure among compound symmetry, heterogeneous compound symmetry, autoregressive, heterogeneous autoregressive, Toeplitz, unstructured, and banded for each parameter. Treatment means were compared using the least squares mean linear hypothesis test (LSMEANS/DIFF). Significance was declared at P ≤ 0.05 and a trend was discussed when 0.05 < P ≤ 0.10.

Associations between the relative abundance of the 10 most relatively abundant genera, as well as three methanogenic taxa, and fermentation parameters were assessed using Spearman’s rank correlation coefficient (r). P-values were adjusted for multiple comparisons the Benjamini–Hochberg method. Fermentation parameters were fit to the non-metric multidimensional scaling (NMDS) ordinations of the Bray–Curtis dissimilarities using the envfit function in vegan with 10,000 permutations. Differentially abundant OTUs between the O3 and control treatments were determined using DESeq2 v. 1.24.0 (Love et al., 2014) in R. To account for unequal sequencing depth, the 16S rRNA gene samples were randomly subsampled to 23,900 sequences per sample prior to analysis of the diversity measures and Bray–Curtis dissimilarities. DESeq2 uses a negative binomial generalized linear model and the Wald test to identify differentially abundant OTUs. For the DESeq2 analysis, samples were not randomly subsampled and only OTUs present in ≥25% of the samples were included.

RESULTS

Effect of O3 on Fermentation Parameters

In comparison to the control, O3 treatment approximately doubled the measured oxidation–reduction potential (P ≤ 0.01) in the buffer solutions in the AM and PM, but there was no difference in the oxidation–reduction potential in the incubation fluid of the control and O3 treatments (Table 2). The addition of O3 caused a 15.4% decrease in total gas production (P ≤ 0.01), and a 20.4% decrease in CH4 production (P ≤ 0.01), without compromising DMD of either the maize silage (P = 0.81) or concentrates (P = 0.44), nor total VFA production (P = 0.49) (Table 2). The concentration of CH4 (%) was decreased by 5.8% in the O3-buffered treatments (P = 0.01). Additionally, O3 supplementation had no effect on the pH (P = 0.87). There were no effects of O3 treatment on acetate, propionate, branched-chain VFA, valerate, or caproate production (P ≥ 0.23); however, butyrate production (mmol/d) was 7.14% greater (P = 0.04) with O3 supplementation compared to the control treatment.

Effect of O3 on the Rumen Microbiota

A total of 3491 archaeal and bacterial OTUs were identified among all samples with an average sequencing depth of 53,411 ± 1454 SEM sequences per sample. The structure of the rumen solid-associated microbiota (SAM) was strongly affected by substrate (PERMANOVA: R2 = 0.63; P < 0.01; Supplementary Figure S1) and less so by sampling time (R2 = 0.048; P < 0.01; Figures 1A,B). However, the solid-associated rumen samples were not affected by O3 treatment within each substrate type (P > 0.05). The microbial community structure of the liquid-associated rumen samples (LAM) was also not affected by O3 treatment (P > 0.05) but there was a time effect (R2 = 0.25; P < 0.01; Figure 1C). Additionally, no differentially abundant OTUs were identified between the control and O3 treatments for either the LAM or SAM samples.

### Table 2

|                  | Control | O3     | SEM   | P-value |
|------------------|---------|--------|-------|---------|
| pH               | 6.82    | 6.82   | 0.04  | 0.87    |
| Redox buffer, AM| 149.0   | 288.4  | 12.64 | < 0.01  |
| Redox buffer, PM| 160.3   | 305.2  | 12.75 | < 0.01  |
| Redox vessel     | 317.4   | 319.0  | 6.83  | 0.87    |
| Gas production, mL| 1790.8 | 1515.2 | 50.06 | < 0.01  |
| CH4, %           | 5.87    | 5.53   | 0.07  | 0.01    |
| CH4, mg/d        | 75      | 59.7   | 2.28  | < 0.01  |
| DMD concentrate, %| 75.9    | 74.9   | 0.89  | 0.44    |
| DMD silage, %    | 60.1    | 60.0   | 0.52  | 0.81    |
| Total VFA, mmol/d| 9.38    | 9.75   | 0.36  | 0.49    |
| Acetate (A), mmol/d| 4.51   | 4.45   | 0.16  | 0.81    |
| Propionate (P), mmol/d| 5.37   | 5.19   | 0.13  | 0.31    |
| Butyrate, mmol/d | 1.68    | 1.80   | 0.04  | 0.04    |
| BCVFA, mmol/d    | 0.26    | 0.25   | 0.01  | 0.40    |
| Valerate, mmol/d | 2.36    | 2.27   | 0.08  | 0.37    |
| Caproate, mmol/d | 0.52    | 0.47   | 0.03  | 0.23    |
| Ratio A:P        | 0.84    | 0.86   | 0.09  | 0.07    |

BCVFA, branched-chain VFA (iso-butyrate + iso-valerate); SEM, standard error of the means.
Among the fermentation parameters, butyrate ($R^2 = 0.36; P < 0.01$) and total VFA ($R^2 = 0.28; P < 0.05$) were significantly associated with the NMDS ordination of the Bray–Curtis dissimilarities for the SAM samples from the grain concentrates substrate. Methane ($R^2 = 0.31; P < 0.05$) and propionate ($R^2 = 0.33; P < 0.05$) concentrations as well as pH ($R^2 = 0.27; P < 0.05$) were significantly correlated with NMDS ordination for the maize silage SAM samples. There were no differences on the alpha diversity indices identified between the O$_3$ and control for the LAM (Supplementary Table S1).

Further analysis of the SAM microbiota (Table 3) demonstrated that on day 10 the supplementation with O$_3$ decreased the number of OTUs (richness) and the Shannon diversity index ($P = 0.02$) compared to the control. On day 5, none of the alpha-diversity metrics were affected by O$_3$ treatment ($P \geq 0.63$).

Five unique archaeal genera along with unclassified *Methanomethylophilaceae* members belonging to the *Methanobacteria* and *Thermoplasmata* classes were identified in the rumen samples. All of these taxa are methanogens. The large
majority of archaeal 16S rRNA gene sequences were identified as unclassified members of the Methanomethylophilaceae family (data not shown). The overall relative abundance of these methanogenic taxa was 5.15 ± 0.28% (SEM) and 0.43 ± 0.04% in the LAM and SAM samples, respectively. Ozone treatment decreased the relative abundance of the total methanogen population in the SAM microbiota on day 10 (P = 0.02; Table 4), as well as Methanomicrobium (P = 0.05) and unclassified Methanomethylophilaceae (P = 0.01). There was no effect of O₃ treatment (P ≥ 0.14) on any of the methanogenic genera in the LAM microbiota (Table 5).

The effect of O₃ treatment on the 10 most relatively abundant bacterial genera on days 5 and 10 in the SAM samples is displayed in Table 6 and Figure 2. Nearly all of these genera differed in relative abundance between the grain concentrates and silage diets; however, only the Schwartzia genus was significantly affected by the O₃ treatment, and only on day 5 in the grain diet. None of the other relatively abundant genera were affected by O₃ supplementation. There were no significant differences in the relative abundance of the 10 most relatively abundant bacterial genera in the LAM samples by O₃ treatment or between the grain concentrates and maize silage diets (P > 0.05) (Figure 3). Prevotella and Bifidobacterium were the most relatively abundant genera in the LAM and SAM samples, respectively. Associations between the 10 most relatively abundant genera as well as the three most relatively abundant methanogenic taxa and fermentation parameters were determined using Spearman’s rank correlation coefficient (Figur 4). Only the relative abundance of Methanomicrobium and Ruminobacter was significantly associated with any of the parameters (P < 0.05) and both were negatively correlated with pH. Some of the positive associations of note (Spearman’s rank correlation coefficient ρ > 0.50; P < 0.10) were Prevotellaceae YAB2003 with CH₄ and propionate, Megaphera with BCVFA and caproate, and Methanobrevibacter with BCVFA.

### DISCUSSION

It is known that manipulation of diet through changes in composition or additives is an effective strategy to reduce enteric methanogenesis in the ruminant (Beauchemin et al., 2008; Hristov et al., 2013; Teoh et al., 2019). It has been suggested

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**Table 3** | Effect of ozone (O₃) on richness (number of OTUs) and diversity indexes of the solid-associated microbe (SAM) samples for each sampling days 5 and 10.

| SAM | Treatment | Substrate | P-value |
|-----|-----------|-----------|---------|
|     | Day       | Control   | O₃      | SEM     | Grain | Maize silage | SEM |
| Number of OTUs | 5 | 249 | 234 | 22.6 | 171 | 312 | 22.0 | 0.63 | <0.01 | 0.07 |
| Shannon diversity | 5 | 2.5 | 2.5 | 0.13 | 2.37 | 2.54 | 0.133 | 0.99 | 0.39 | 0.10 |
| Inverse Simpson's diversity | 5 | 5.09 | 5.12 | 0.539 | 5.66 | 4.54 | 0.514 | 0.97 | 0.14 | 0.14 |
| Number of OTUs | 10 | 302 | 226 | 21.1 | 193 | 336 | 21.1 | 0.02 | <0.01 | 0.80 |
| Shannon diversity | 10 | 3.15 | 2.57 | 0.149 | 2.51 | 3.21 | 0.144 | 0.02 | 0.01 | 0.20 |
| Inverse Simpson's diversity | 10 | 11.68 | 5.77 | 3.008 | 6.30 | 11.16 | 2.914 | 0.20 | 0.25 | 0.30 |

SEM, standard error of the means.

**Table 4** | Effects of ozone (O₃) treatment on the percent relative abundance of methanogenic genera on days 5 and 10 in solid-associated microbes (SAM) samples.

| SAM | Treatment | Substrate | P-value |
|-----|-----------|-----------|---------|
|     | Day | Control | O₃ | SEM | Grain | Maize silage | SEM | O₃ Substrate | O₃ × substrate |
| Methanobrevibacter | 5 | 0.149 | 0.142 | 0.0308 | 0.052 | 0.239 | 0.0308 | 0.88 | <0.01 | 0.72 |
| Methanosphaera | 5 | ND | ND | ND | ND | ND | ND | N/A | N/A |
| Methanomicrobium | 5 | 0.014 | 0.018 | 0.0054 | 0.008 | 0.023 | 0.0041 | 0.58 | <0.01 | 0.34 |
| Methan茵micrococcus | 5 | ND | ND | ND | ND | ND | ND | N/A | N/A |
| Methanomethylophilus | 5 | 0.002 | 0.002 | 0.0015 | 0.002 | 0.002 | 0.0015 | 0.78 | 0.78 | 0.10 |
| UC Methanomethylophilaceae | 5 | 0.229 | 0.220 | 0.0337 | 0.136 | 0.313 | 0.037 | 0.91 | 0.03 | 0.37 |
| Total methanogens | 5 | 0.393 | 0.383 | 0.0712 | 0.198 | 0.578 | 0.0712 | 0.92 | <0.01 | 0.38 |
| Methanobrevibacter | 10 | 0.077 | 0.079 | 0.0192 | 0.034 | 0.122 | 0.0192 | 0.95 | <0.01 | 0.76 |
| Methanosphaera | 10 | ND | ND | ND | ND | ND | ND | N/A | N/A |
| Methanomicrobium | 10 | 0.058 | 0.079 | 0.0061 | 0.030 | 0.068 | 0.0061 | 0.05 | <0.01 | 0.14 |
| Methan茵micrococcus | 10 | ND | 0.0014 | 0.0009 | 0.0002 | 0.0012 | 0.00086 | 0.25 | 0.44 | 0.44 |
| Methanomethylophilus | 10 | 0.002 | 0.002 | 0.0015 | 0.005 | 0.002 | 0.0015 | 0.23 | 0.18 | 0.10 |
| UC Methanomethylophilaceae | 10 | 0.433 | 0.268 | 0.0431 | 0.222 | 0.479 | 0.0431 | 0.01 | <0.01 | 0.13 |
| Total methanogens | 10 | 0.570 | 0.392 | 0.0512 | 0.290 | 0.672 | 0.0512 | 0.05 | <0.01 | 0.17 |

SEM, standard error of the means; ND, not detected; N/A, non-applicable. UC, unclassified.
TABLE 6 | Effects of ozone (O₃) treatment on the percent relative abundance of the methanogenic genera on days 5 and 10 in solid-associated microbe (SAM) samples.

| SAM                        | Treatment | Substrate | O₃  | Substrate | O₃    | O₃ × substrate |
|----------------------------|-----------|-----------|-----|-----------|-------|----------------|
|                            | Day       | Control   |    | SEM       |       |                |
| Bifidobacterium            | 5         | 0.319     |    | 0.600     | 0.1323| 0.17           |
| Methanospirillum            | 5         | 0.001     |    | 0.009     | 0.0036| 0.15           |
| Methanocorpus              | 5         | 0.049     |    | 0.042     | 0.0129| 0.72           |
| Methanobrevibacter         | 5         | 0.004     |    | 0.002     | 0.0016| 0.48           |
| Methanosphaera             | 5         | 0.070     |    | 0.067     | 0.0283| 0.94           |
| Methanobrevibacter         | 5         | 0.71      |    | 4.74      | 0.422 | 0.14           |
| Methanosphaera             | 5         | 6.15      |    | 5.46      | 0.363 | 0.21           |
| Methanocorpus              | 5         | 0.237     |    | 0.326     | 0.0539| 0.27           |
| Methanocorpus              | 5         | 0.001     |    | 0.001     | 0.0010| 0.59           |
| Methanocorpus              | 10        | 0.291     |    | 0.384     | 0.1452| 0.66           |
| Methanocorpus              | 10        | 0.004     |    | 0.019     | 0.0084| 0.22           |
| Methanocorpus              | 10        | 0.050     |    | 0.051     | 0.0211| 0.97           |
| Methanomethylophilaceae    | 5         | 4.28      |    | 3.36      | 0.544 | 0.25           |
| Methanomethylophilaceae    | 5         | 10.47     |    | 4.14      | 0.592 | 0.40           |

SEM, standard error of the means; UC, unclassified.

that certain bacterial and archaeal species in the rumen are more susceptible than others to unfavorable conditions (Wintsche et al., 2018). In the present study, inclusion of O₃ in the buffer during in vitro fermentation led to a 15.4% decrease in total gas production, and a 20.4% decrease in CH₄ production, confirming our first hypothesis. As far as we can ascertain, this is the first study to demonstrate in vitro, that ruminal methanogenesis can be inhibited by O₃.

The O₃ treatment reduced enteric CH₄ production without compromising DMD. Although treatment with O₃ did not significantly alter the bacterial and archaeal microbiota in the LAM samples and had no effect on the SAM bacterial populations, it was associated with a reduction in relative abundance of methanogens in the SAM samples and an increase in the production of butyric acid. In particular, significant effects were observed on both the SAM methanogenic microbiota and bacterial diversity and richness on day 10. Feng et al. (2013) demonstrated similar findings with a decreased methanogen diversity in rice-paddy soil exposed to elevated ground-level O₃. While the most relatively abundant genera differ between the present study and that of Feng et al. (2013), similar patterns of increasing richness (number of OTUs) over time were observed.

The relative abundance of the Methanocorpus genus, unclassified Methanomethylophilaceae, and total methanogenic 16S rRNA gene sequences was significantly reduced on day 10 in the O₃-treated SAM samples (P ≤ 0.05). Similarly, Zhang et al. (2016) indicated a decrease in the abundance of methanogens in both O₃-sensitive and O₃-tolerant rice cultivars subjected to elevated ground-level O₃. In the present study, none of the methanogenic genera in either the rumen solid or liquid samples were affected by the O₃ treatment on day 5. However, by day 10, the relative abundance of methanogens in the solid samples was reduced by approximately 31%, indicating that the effect of O₃ on methanogens increased with time. Methanocorpus and other hydrogenotrophs including Methanobrevibacter utilize recycled hydrogen and carbon dioxide for methanogenesis. Ozone molecules are highly reactive and when present in the

TABLE 5 | Effects of ozone (O₃) treatment on the percent relative abundance of the methanogenic genera on days 5 and 10 in liquid-associated microbe (LAM) samples.

| LAM                         | Day | Control | O₃ | SEM | P-value |
|-----------------------------|-----|---------|----|-----|---------|
| Methanobrevibacter          | 5   | 0.319   | 0.600 | 0.1323 | 0.17 |
| Methanosphaera              | 5   | 0.001   | 0.009 | 0.0036 | 0.15 |
| Methanocorpus               | 5   | 0.049   | 0.042 | 0.0129 | 0.72 |
| Methanocorpus               | 5   | 0.004   | 0.002 | 0.0016 | 0.48 |
| Methanomethylophilaceae     | 5   | 0.070   | 0.067 | 0.0283 | 0.94 |
| Methanomethylophilaceae     | 5   | 5.71    | 4.74 | 0.422 | 0.14 |
| Methanobrevibacter          | 5   | 6.15    | 5.46 | 0.363 | 0.21 |
| Methanosphaera              | 5   | 0.237   | 0.326 | 0.0539 | 0.27 |
| Methanocorpus               | 5   | 0.001   | 0.001 | 0.0010 | 0.59 |
| Methanocorpus               | 10  | 0.291   | 0.384 | 0.1452 | 0.66 |
| Methanocorpus               | 10  | 0.004   | 0.019 | 0.0084 | 0.22 |
| Methanocorpus               | 10  | 0.050   | 0.051 | 0.0211 | 0.97 |
| Methanomethylophilaceae     | 10  | 4.28    | 3.36 | 0.544 | 0.25 |
| Methanomethylophilaceae     | 10  | 4.87    | 4.14 | 0.592 | 0.40 |

SEM, standard error of the means; UC, unclassified.

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**FIGURE 2** Box and whisker plots of the 10 most relatively abundant bacterial genera in the solid-associated microbe (SAM) rumen samples by ozone treatment and day (D5 or D10) for the grain (G) and maize silage (MS) substrates. The box in the box plots indicates the interquartile range (IQR) (middle 50% of the data), the middle line represents the median value, and the whiskers represent 1.5 times the IQR.

The rumen will preferentially utilize H$_2$ for conversion into H$_2$O, reducing hydrogen ion availability for methanogens to synthesize CH$_4$ (Zabranska and Pokorna, 2018). Competition for hydrogen thus has the ability to reduce methanogen activity, as seen in Zhao et al. (2018). A similar reduction in methanogenesis by using this hydrogen-competition mechanism has been achieved with other feed supplements such as nitrate (Zhao et al., 2018). However, this hydrogen competition mechanism is unlikely to be the only mechanism whereby ozone inhibits methanogenesis. As referred to in the introduction, ozone has a general biocidal effect against many types of microorganisms, and given the sensitivity of methanogens to oxygen, it seems that ozone may also have a specific biocidal effect against methanogens, or at least a direct inhibitory effect on methanogenesis (Brodowska et al., 2018; Remondino and Valdenassi, 2018).

*Methanomethylophilaceae* is a newly proposed family within the *Methanomassiliicoccales* order (Gaci et al., 2014). The characterized taxa in this order are hydrogen-dependent methylotrophs that produce CH$_4$ through the reduction of methanol or methylamine (Lang et al., 2015) and are relatively abundant in the bovine rumen (Henderson et al., 2015; Holman and Gzyl, 2019). Interestingly, we recently reported a decrease in *Methanomethylophilaceae* members in response to hardwood biochar supplementation in a similar RUSITEC experiment (Teoh et al., 2019). We did not find any strong associations between any of the three most relatively abundant methanogenic taxa and CH$_4$ production although the PCR primers used likely do not amplify all 16S rRNA genes found in methanogens. *Methanobrevibacter* was positively associated with BCVFA concentration while *Methanomicrobium* was negatively correlated. This may be related to the fact that *Methanobrevibacter* spp. and *Methanomicrobium* spp. have opposite growth requirements for BCVFA (Tanner and Wolfe, 1988; Miller and Lin, 2002). *Schwartzia* was the only bacterial genus among the relatively abundant genera that was significantly affected by O$_3$ treatment and only in the grain substrate diet at day 5. The only species in this genus is *Schwartzia succinivorans* which was originally isolated from the bovine rumen and utilizes succinate as a sole energy source producing propionate.
Schwartzia has also been previously reported to be negatively associated with CH$_4$ emissions in cattle (Cunha et al., 2017).

Ozone is a highly reactive free radical with strong positive redox potential enabling it to attack glycoproteins and glycolipids in the bacterial cell membrane resulting in rupture and destruction of the cell (Greene et al., 2012; Megahed et al., 2018). In the present research, O$_3$ treatment almost doubled the redox potential in the buffer solution added to the "O$_3$" incubations. Methanogenic archaea have unique cell walls composed of pseudopeptidoglycan, or pseudomurein, components that make their cell wall similar to those of Gram-positive bacteria and this may contribute to their sensitivity to antimicrobials (Varnava et al., 2017; Doyle et al., 2019) and destructive agents such as O$_3$ that target glycosidic bonds. Friedman et al. (2016) suggested that the redox potential effect is caused by a reduction in the number of anti-reactive oxygen species proteins in the genome of the most abundant methanogens, particularly Methanomicrobium spp. Additionally, O$_3$ is able to cause a complete loss of function by disruption of essential enzymatic activity, as the disturbance of membrane-bound enzymes, proteins, and DNA leads to cell wall lysis (Komanapalli and Lau, 1996) and ultimately a reduction in methanogenesis and total gas production.

Differences in individual and herd CH$_4$ emissions can be attributed to intrinsic factors such as the rumen microbiota and variances in particle retention time in the rumen (Martin et al., 2010), as well as extrinsic factors such as manipulation of feed components including silage derivatives, concentrates, and additives. Methanogenesis is largely dependent on the metabolic properties and function of certain members of the rumen bacterial microbiota, which are in turn dependent on substrate type and availability (Evans et al., 2019). Although forage substrate preparation has been shown to have little effect on fermentation parameters (Duarte et al., 2017a), forage quality is known to impact the rumen microbiota, with cattle having a higher feed conversion efficiency producing approximately 20% less CH$_4$ (Nkrumah et al., 2006; Hegarty et al., 2007). Feed used in the present study is based on a combination of maize silage and grain concentrates at 50%
FIGURE 4 | Heatmap of the association between the 10 most relatively abundant bacterial genera and three methanogenic taxa and fermentation parameters based on Spearman’s rank correlation coefficient. Correlations that are statistically significant (P < 0.05) are indicated with **” and those with P-values less than 0.10 are denoted with “*”.

w/w on a 5.5 g DM basis, consisting of 11.95% CP and 29.56% NDF. Similar studies have seen a correlation between the use of relatively high-quality forage and a reduction in ruminant CH$_4$ production (Ramos et al., 2018); though further investigation is required to demonstrate a causal relationship as other experimental conditions are highly variable (Benchaar et al., 2001).

High concentrate diets with a greater presence of readily fermentable substrates (e.g., starch) compared to high forage diets have resulted in a decrease in CH$_4$ emissions (Popova et al., 2011). Starch in grain silages favors propionate production rather than acetate as propionate acts as a sink for hydrogen ions and reduces the availability of H$_2$ for methanogens (Beauchemin et al., 2008). Thus, an increase in the abundance of propionate-producing bacteria should reduce methanogenesis through the diversion of hydrogen (Myer et al., 2015). Indeed, a decreased acetate: propionate (A:P) ratio has been observed in numerous feed manipulation studies that have resulted in decreased CH$_4$ production (Ramos et al., 2018; Williams et al., 2019). It should also be noted that the A:P ratio tended to be greater (P = 0.07) in the O$_3$ treatment; this despite a high relative abundance of Prevotella spp., some of which are propionate producers (McCabe et al., 2015; Emerson and Weimer, 2017). This also corresponds with the positive association observed between the related Prevotellaceae YAB2003 group and propionate concentration. Taken together, our findings that ozone had no effect on the A/P ratio, no effects on the relative abundance of ruminal bacteria,
but reduced the total number of methanogens and reduced production of methane indicate that ozone treatment specifically inhibits methanogenesis.

While the present study has shown O$_3$ treatment can reduce methanogenesis in bovine ruminal fluid, there are many questions to be answered before this could be implemented as a strategy to reduce CH$_4$ emissions from the livestock industries. For example, what would be the best and cheapest way to administer O$_3$ to ruminants? Ozonation of drinking water offers a potential solution since substantial production benefits have been realized when using clean drinking water in cattle (Schütz, 2012). It is also unknown whether O$_3$ administration reduces enteric CH$_4$ production in vivo. Results from this study indicate the importance and promise of in vivo testing, although certain factors including rumen transit time, individual rumen microbiota composition, and fermentation characteristics will differ between individual cattle. Additionally, does O$_3$ administration to ruminants have adverse impacts on their health and production? It is hoped that this experiment may stimulate further research designed to answer these questions and that O$_3$ may eventually play a role in reducing enteric CH$_4$ emissions from livestock industries.

CONCLUSION

The inclusion of O$_3$ into a fermenter vessel via buffer decreased total gas production, notably CH$_4$ production by 20.4%. It is important to note that neither DMD nor total VFA production was affected. Ozone, upon interaction with pre-existing oxygen in the buffer, doubled the buffer redox potential. There was no effect by O$_3$ on other fermentation parameters, with the exception of a 7% increase in butyrate production. Ozone treatment decreased both the richness (number of OTUs) and diversity (Shannon diversity) in the solid rumen samples on day 10 ($P = 0.02$). Also, on day 10, O$_3$ treatment decreased the relative abundance of the total methanogen population in the SAM samples, particularly Methanomicrobium spp. The decrease in CH$_4$ gas production may be explained by disruption of the members of the methanogenic genera; however, further research is required in vivo to ensure the safety and efficiency of O$_3$ for commercial agriculture.

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DATA AVAILABILITY STATEMENT

The datasets presented in this study can be found in online repositories. The names of the repository/repositories and accession number(s) can be found in the article/Supplementary Material.

ETHICS STATEMENT

The animal study was reviewed and approved by The University of Sydney Animal Ethics Committee (Approved Protocol Number 2015/835).

AUTHOR CONTRIBUTIONS

AC designed the studies. LZ, EC, DH, KG, PM, and AC acquired data, read and critically revised drafts for intellectual contents, and approved the final manuscript. LZ, EC, DH, PM, and AC conducted laboratory analysis. DH and KG conducted bioinformatics. AC and DH ran statistical analysis. LZ, EC, DH, PM, and AC wrote the manuscript. All authors contributed to the article and approved the submitted version.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmicb.2020.571537/full#supplementary-material

Supplementary Figure 1 | Non-metric multidimensional scaling (NMDS) of the Bray-Curtis dissimilarities for the archaeal and bacterial solid-associated rumen samples by substrate and treatment.
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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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