Effect of condensed tannins from *Leucaena leucocephala* on rumen fermentation, methane production and population of rumen protozoa in heifers fed low-quality forage

Angel T. Piñeiro-Vázquez1,2,*, Jorge R. Canul-Solis1, Guillermo O. Jiménez-Ferrer3, José A. Alayón-Gamboa4, Alfonso J. Chay-Canul3, Armin J. Ayala-Burgos1, Carlos F. Aguilar-Pérez1, and Juan C. Ku-Vera1

**Objective:** The aim of the experiment was to assess the effect of increasing amounts of *Leucaena leucocephala* forage on dry matter intake (DMI), organic matter intake (OMI), enteric methane production, rumen fermentation pattern and protozoa population in cattle fed *Pennisetum purpureum* and housed in respiration chambers.

**Methods:** Five crossbred heifers (*Bos taurus* × *Bos indicus*) (BW: 295±6 kg) were fed chopped *P. purpureum* grass and increasing levels of *L. leucocephala* (0%, 20%, 40%, 60%, and 80% of dry matter [DM]) in a 5×5 Latin square design.

**Results:** The voluntary intake and methane production were measured for 23 h per day in respiration chambers; molar proportions of volatile fatty acids (VFAs) were determined at 6 h postprandial period. Molar concentration of VFAs in rumen liquor were similar (p>0.05) between treatments. However, methane production decreased linearly (p<0.005), recording a maximum reduction of up to ~61% with 80% of DM incorporation of *L. leucocephala* in the ration and no changes (p>0.05) in rumen protozoa population were found.

**Conclusion:** Inclusion of 80% of *L. leucocephala* in the diet of heifers fed low-quality tropical forages has the capacity to reduce up to 61.3% enteric methane emission without affecting DMI, OMI, and protozoa population in rumen liquor.

**Keywords:** Methane; Crossbred Heifers; Volatile Fatty Acids; Forages; Respiration Chambers

**INTRODUCTION**

Methane (CH₄) gas is a byproduct of anaerobic fermentation of carbohydrates in the rumen. It is considered a greenhouse gas with a global warming potential twenty-three times greater than that of CO₂ [1] and every year 80 million tons are generated from anthropogenic activities. It has been estimated that enteric methane contributes 39.1% of greenhouse gas emissions generated by the livestock sector [2]. Additionally, CH₄ emissions represent an energy loss which ranges from 2% to 12% of the gross energy consumed by ruminants [3].

In tropical regions, ruminants are usually fed low-quality forages which are characterized by their low concentration of crude protein (CP) and digestible energy, and their high content of neutral detergent fiber (NDF) and lignin, which induces a higher methane convention rate value [4] decreasing the efficiency of energy utilization [5]. Nonetheless, feeding ruminants with legumes favors a lower CH₄ production [6].

In the tropical regions, there is a large diversity of legume species which contain secondary metabolites such as condensed tannins (CT), which decreases the protozoa population, methanogenic archaea [7], and the synthesis of enteric CH₄ [8,9]. *Leucaena leucocephala* (*L. leucocephala*) is one of the main tropical shrub species which contains CT with potential...
to reduce CH₄ emissions [10], improve CP digestibility [11], N retention [8,9] and improve the efficiency of energy intake by ruminants [12]. Harrison et al [13] have recently reported higher levels of productivity with cattle fed leucaena in Australia as well as lower enteric CH₄ emissions to the atmosphere.

The aim of the work hereby described was to assess the effect of supplying increasing concentrations of CT from L. leucocephala in the ration on feed intake, molar proportions of volatile fatty acids (VFAs), changes in rumen protozoa population and enteric CH₄ emissions by heifers fed a basal ration of tropical grass.

MATERIALS AND METHODS

Animal care
The animals were treated in accordance with guidelines and regulations for animal experimentation of the Faculty of Veterinary Medicine and Animal Science (FMVZ), University of Yucatan, Merida, Mexico.

Location
The experiment was carried out at the FMVZ, University of Yucatan in Merida, Mexico. Climate in the region is warm with an average temperature of 26.8°C and average rainfall of 984.4 mm per year [14].

Experimental animals
Five crossbreds (Bos taurus×Bos indicus) heifers with an average body weight (BW) of 295±6.8 kg were housed in metabolic crates in a roofed building with concrete floor without walls. Before the experiment heifers were internally dewormed with Ivermectin (AGROVET, S.A de C.V. Mexico, Federal District, Mexico) (1%; 1 mL/50 kg BW) and injected intramuscularly with ADE vitamins (1 mL/10 kg BW).

Experimental design and treatments
A 5×5 Latin square design [15] was used; each period lasted 19 days (13 days for adaptation to management and rations, and six days for measurements of response variables). During the adaptation period (13 days), the animals remained outside the chambers and only during the measurement period (6 days) the intake, digestibility and enteric methane production were measured. Prior to the experiment, animals were adapted to enter the respiration chambers for a period of ~3 weeks to reduce the effect of stress on voluntary intake of dry matter (DM) and behavior in and out of the chambers. Basal ration consisted of fresh forage of Taiwan (Pennisetum purpureum [P. purpureum]) grass and increasing levels of chopped forage of L. leucocephala as a replacement for the P purpureum forage which was homogenously mixed (Forage-legume) to avoid selection. The chemical composition of the forages is shown in Table 1. Experimental treatments were increasing levels of L. leucocephala (0%, 20%, 40%, 60%, and 80% of DM) in the ration. Additionally, heifers were supplemented with a commercial mineral mixture Fogysal Cattle (FOGYSA S.A de C.V. Merida, Yucatan, Mexico) (250 g/heifers/d).

Response variables

Determination of rumen pH and molar proportions of volatile fatty acid: Samples of rumen liquor were taken by an esophageal tube according to Ramos-Morales et al [16]. Rumen liquor was obtained during six days before measurements in each period, six hours postprandial according to recommendations by Hales et al [17] and Bathha et al [18] in order to analyze rumen pH and pattern of VFAs in the rumen. Samples of rumen liquor were filtered through double layers of cheesecloth to retain large particles. Rumen pH was determined immediately after taking the sample with a portable potentiometer (HANNA Instruments, Woonsocket, RI, USA), previously calibrated with buffers at pH’s 4, 7 and 10. For the determination of molar proportions of VFAs in rumen liquor, 4 mL samples were taken and added 1 mL of a deproteinizing solution consisting of metaphosphoric and 3-methylvaleric acids. For VFA analysis, the technique proposed by Ryan [19] was employed using a gas chromatograph (Hewlett-Packard, 5890 series III), equipped with a flame ionization detector and a column HP-FFAP measuring 30 m×0.53 mm, temperature of the injector and the detector was in both cases 200°C.

Measurement of enteric CH₄
Measurement of CH₄ was carried out while heifers were housed in respiration chambers [20], with dimensions 2.10 m×1.60 m×3.10 m (height, width and length, respectively) and a total internal volume of 9.38 m³. Mass flow generators (Sable Systems International, Las Vegas, NV, USA) pulled out the air from the chamber at a rate of 500 L/min creating a negative pressure of 475 Pa inside the chamber relative to the outside environment.

Measurement of CH₄ in air samples was carried out with

Table 1. Chemical composition of the forages (g/kg DM)

| Item                        | Pennisetum purpureum | Leucaena leucocephala |
|-----------------------------|-----------------------|-----------------------|
| Organic matter              | 930.32                | 940.5                 |
| Crude protein               | 60.64                 | 160.29                |
| NDF                         | 670.62                | 580.85                |
| ADF                         | 410.99                | 420.01                |
| Lignin                      | 70.24                 | 160.0                 |
| Secondary metabolites       |                       |                       |
| Total phenols               | 0                     | 7.7                   |
| Condensed tannins           | 0                     | 21.0                  |

DM, dry matter; NDF, neutral detergent fiber; ADF, acid detergent fiber.

1) Expressed as g tannic acid eq/kg DM.

2) Expressed as g leuconidin eq/kg DM.
an infrared analyzer (MA-10, Sable Systems International, USA). High purity CH₄ gas (99.9999% CH₄; Praxair, Mexico) was released from a small cylinder into the chambers and the recovery rate was assessed to be 100.2% which is comparable to recovery rates obtained in other laboratories [21]. The methane analyzer was zeroed by infusing pure N₂ and the linearity (r² = 0.9999) of the analyzer response was assessed by infusing increasing concentrations of methane (1,000; 2,500; 5,000; 7,500 ppm) diluted in N₂, before each run. Heifers were kept inside the respiration chambers at a temperature of 23°C and relative humidity of 55%. Fresh water was available at all times from an automatic drinking bowl and a small fan mixed the air inside the chambers. Methane measurements were carried out during three consecutive days for 23 h continuously during the measurement period of the response variables, inside the respiration chambers intake and digestibility were also measured [20,4]. Data obtained were adjusted to 24 h runs with ExpeData software (Sable Systems International, USA) and converted to liters of methane produced per day.

**Voluntary intake**

Heifers were fed _ad libitum_ allowing a refusal of at least 15% of the dry matter offered the previous day. Feed refusals were weighed at 0900 h the following day. Voluntary intake was determined by the difference between the amount offered and refused. Feed offered covered an estimated intake of 7.0 kg DM/d.

**Rumen protozoa population**

Protozoa in samples of rumen liquor were counted according to the procedures described by Rosales [22]. One mL of rumen liquor was mixed with 1 mL of methylgreen formalin saline solution (35 mL/L formaldehyde, 0.14 mM NaCl, 0.92 mM methylgreen) and centrifuged at 2,000 rpm for 20 min. Then, an aliquot was taken and introduced to a modified Fusch-Rosenthal chamber (Electron Microscopy Science, Hatfield, PA, USA) (IVD 98/79 CE) and observed under the microscope (Mikon-YS100) (Nikon Instruments Inc, Melville, NY, USA) at 40×. Number of ciliate protozoa was reported as log 10 from an aliquot was taken and introduced to a modified Fusch-Rosenthal solution (35 mL/L formaldehyde, 0.14 mM NaCl, 0.92 mM methylgreen) to the procedures described by Rosales [22]. One mL of rumen liquor was mixed with 1 mL of methylgreen formalin saline solution (35 mL/L formaldehyde, 0.14 mM NaCl, 0.92 mM methylgreen) and centrifuged at 2,000 rpm for 20 min. Then, an aliquot was taken and introduced to a modified Fusch-Rosenthal chamber (Electron Microscopy Science, Hatfield, PA, USA) (IVD 98/79 CE) and observed under the microscope (Mikon-YS100) (Nikon Instruments Inc, Melville, NY, USA) at 40×. Number of ciliate protozoa was reported as log 10 from an aliquot was taken and introduced to a modified Fusch-Rosenthal solution (35 mL/L formaldehyde, 0.14 mM NaCl, 0.92 mM methylgreen) to the procedures described by Rosales [22]. One mL of rumen liquor was mixed with 1 mL of methylgreen formalin saline solution (35 mL/L formaldehyde, 0.14 mM NaCl, 0.92 mM methylgreen) and centrifuged at 2,000 rpm for 20 min. Then, an aliquot was taken and introduced to a modified Fusch-Rosenthal chamber (Electron Microscopy Science, Hatfield, PA, USA) (IVD 98/79 CE) and observed under the microscope (Mikon-YS100) (Nikon Instruments Inc, Melville, NY, USA) at 40×. Number of ciliate protozoa was reported as log 10 from an aliquot was taken and introduced to a modified Fusch-Rosenthal solution (35 mL/L formaldehyde, 0.14 mM NaCl, 0.92 mM methylgreen) to the procedures described by Rosales [22].

**Chemical analysis**

Dry matter was determined in a forced air oven at 55°C for 48 h (constant weight) [25]. Nitrogen (CP = N×6.25) determinations were carried out with a LECCO CN-2000 series 3740 instrument (LECCO, Corporation, Saint Joseph, MI, USA) [25]. Organic matter was assessed by incineration in muffle furnace at 550°C for 6 h (AOAC Method #923.03) and the content of NDF and acid detergent fiber (ADF) were determined using the methods described by Van Soest et al [26]. Concentration of CT in forage samples was performed by the HCL-butanol method described by Makkar [27].

**Statistical analysis**

Data on voluntary intake, apparent digestibility, nitrogen balance, protozoa population and CH₄ production were subjected to analysis of variance for a 5×5 Latin square design [15] using the procedure PROC ANOVA from SAS (SAS Inst. Inc., Cary, NC, USA) with the model Yᵢⱼᵏ = μ+Pᵢ+Tⱼ+Tkᵢ+eᵢⱼᵏ, where Yᵢⱼᵏ is the dependent variable, μ is the general mean, Pᵢ is the effect of period, Tⱼ is the effect of animal, Tkᵢ is the effect of treatment, and eᵢⱼᵏ is the residual error. Statistical differences were declared significant at p≤0.05. Additionally, surface response analysis was carried out to assess the linear, quadratic or cubic effects [28] of the response to treatments (0%, 20%, 40%, 60%, and 80% of _L. leucocephala_ in the ration).

**RESULTS**

**Voluntary intake**

Dry matter intake (DMI, kg/d and g/kg⁰.⁷⁵) was similar among treatments with different levels of _L. leucocephala_ forage, reporting an average of 7.0 (kg/d) and 98.7 (g/kg⁰.⁷⁵), respectively (p>0.05). Organic matter intake (OMI) followed the same trend as DMI (Table 2). However, crude protein intake (CPI) increased linearly as the level of incorporation of _L. leucocephala_ in the ration was augmented (p<0.0001).

**Enteric methane production**

The average of methane production in this study was 20.1 L/kg of DMI with the control treatment and as _L. leucocephala_ increased in the diet there was a reduced CH₄ production; with incorporation of 80% of _L. leucocephala_ in the ration methane production reduced by 61.6% respect to 0% of legume (linear reduction p = 0.0005). Also, methane emissions expressed as L CH₄/d, kg of DM, OM, NDF, ADF intake, kg of digestibility dry matter, and organic matter digestibility (OMD), neutral detergent fiber digestibility (NDFD), an acid detergent fiber digestibility (ADFD) was linearly reduced (p<0.01) as the percentage incorporation of _L. leucocephala_ in the ration was increased (Table 3).

**pH, NH₃-N, and molar proportions of volatile fatty acid in the rumen**

No differences were observed in rumen pH as the inclusion of _L. leucocephala_ in the ration of cattle was increased (p>0.05). Although rumen NH₃-N was different between treatments and linearly increased as the level of the foliage of the legume
was increased (p<0.01). As from 80% of *L. leucocephala* (36 mg/dL) in the ration, rumen NH₃-N was significantly increased (138%) with respect to the treatment without incorporation of *L. leucocephala* (15.3 mg/dL) (p<0.05) (Table 4).

Molar proportions of acetic, propionic and butyric acids in rumen liquor were not affected (p>0.05) by the incorporation of *L. leucocephala* in the ration, registering averages of 53.8, 12.9, and 5.7 mmol/L, respectively. When the molar proportion of acetic acid was expressed as percentage, a linear increase (p = 0.037) was detected as the level of incorporation of *L. leucocephala* was increased. Conversely, the molar proportion of propionic acid maintained a linear reduction (p = 0.032) as *L. leucocephala* was increased in the ratio (p<0.0002) (Table 4).

## Rumen protozoa population

Total protozoa numbers, holotrichs, and entodiniomorph in the rumen of cattle were not affected at 6 h after intake of *L. leucocephala* forage (p>0.05) (Table 5).

## DISCUSSION

### Voluntary intake

Intake of tropical foliage that contains CT at moderate concentrations (3% to 6% of DM) improves the nutrient intake of ruminant rations, particularly DM, OM, and CP [29]. In the present work, the incorporation of up to 80% of *L. leucocephala* equivalent to a supply of 16.4 g/kg DMI of CT only had an influence on the greater CP intake (kg/d, CPI/kg DMI Table 2).

### Methane (CH₄) production

| Items                  | Percent of *L. leucocephala* supplemented (% of DM) | SE | Contrast |
|------------------------|-----------------------------------------------------|----|----------|
| BW (kg)                |                                                     |    |          |
| Intake                 |                                                     |    |          |
| DM (kg/d)              | 7.03                                                 | 0.60| ns       |
| OM (kg/d)              | 6.54                                                 | 0.55| ns       |
| CP (kg/d)              | 0.50                                                 | 0.10|          |
| NDF (kg/d)             | 4.63                                                 | 0.40|          |
| ADF (kg/d)             | 2.85                                                 | 0.25|          |
| Digestible intake (kg/d) |                                              |    |          |
| DDM                    | 3.90                                                 | 0.54| ns       |
| OMD                    | 3.72                                                 | 0.52| ns       |
| NDFD                   | 2.60                                                 | 0.33|          |
| ADFD                   | 1.55                                                 | 0.22|          |

DM, dry matter; SE, standard error; L, linear contrast; BW, body weight; OM, organic matter; CP, crude protein; NDF, neutral detergent fiber; ADF, acid detergent fiber; DDM, digestible dry matter; OMD, digestible organic matter; NDFD, digestible neutral detergent fiber; ADFD, digestible acid detergent fibre.

* p < 0.05; ** p < 0.01; ns, non-significant (p > 0.05). No significant effects were found for the quadratic and cubic contrasts.

**Table 3.** Methane (CH₄) production in crossbred heifers fed *Pennisetum purpureum* grass and increasing levels of *Leucaena leucocephala*

| Items                  | Percent of *L. leucocephala* supplemented (% of DM) | SE | Contrast |
|------------------------|-----------------------------------------------------|----|----------|
| BW (kg)                |                                                     |    |          |
| Enteric methane        |                                                     |    |          |
| L CH₄/d                | 137.3                                               | 14.8| **        |
| L CH₄/kg of DMI        | 20.1                                                | 2.1 | **        |
| L CH₄/kg of OMI        | 21.6                                                | 2.3 | **        |
| L CH₄/kg of NDFI       | 30.4                                                | 3.4 | **        |
| L CH₄/kg of ADFI       | 49.1                                                | 5.3 | ** **     |
| L of methane/kg of digestible fractions intake |                                             |    |          |
| L CH₄/kg of DDM        | 37.7                                                | 5.1 | **        |
| L CH₄/kg of OMD        | 39.6                                                | 5.4 | **        |
| L CH₄/kg of NDFD       | 56.5                                                | 11.5| ns        |
| L CH₄/kg of ADFD       | 92.0                                                | 33.2| ns        |

DM, dry matter; SE, standard error; L, linear contrast; Q, quadratic contrast; C, cubic contrast; BW, body weight; DMI, dry matter intake; OMI, organic matter intake; NDFI, neutral detergent fiber intake; ADFI, acid detergent fiber intake; DDM, digestible dry matter; OMD, digestible organic matter; NDFD, digestible neutral detergent fiber; ADFD, digestible acid detergent fibre.

* p < 0.05; ** p < 0.01; ns, non-significant (p > 0.05).
gestibility organic matter intake [DOMI]) without affecting DMI, OMI, and acid detergent fiber intake (Table 2). Additionally, reduction in NDF intake might have been influenced by the small amount of NDF contained in the foliage of *L. leucocephala* (580.8 g/kg DM) and as the level of incorporation of tree foliage was increased, the amount of NDF was reduced [6]. These results differ from those of Delgado et al [30] when they included 27% of *L. leucocephala* in sheep rations and observed an increase in the DMI and OMI, similar to results reported by Wahyuni et al [11] who observed an increase in DMI when they included increasing levels of *L. leucocephala*, registering an increase of 30%, when 60% *leucaena* was incorporated in the ration compared to a control treatment.

It has been pointed out that doses below 2% CT do not affect DMI [31,32]. Grainger et al [33] found that 1.46% CT in the ration of dairy cows did not affect DMI and Beauchemin et al [32] using 2% CT in cattle rations did not find an effect on nutrient intake. In this study, levels of up to 80% *L. leucocephala* only produces 1.64% of CT per kg DMI in the cattle ration, thus it is unlikely that this level influenced DM and OM intakes.

**Enteric methane production**

In this experiment, with heifers fed foliage of *L. leucocephala*, the methane production was lower (11.5 L/kg of DM) [6,34,8] in respect to the heifers without inclusion of *L. leucocephala* (20.1 L/kg of DM) in the ration. In this context, 80% of *L. leucocephala* foliage in the ration, decreased by ~61% the rumen methane production with respect to the control treatment, these results were similar to those obtained [35,8,34].

In the present work, a linear reduction in ruminal methane production was observed as the level of foliage of *L. leucocephala* in the ration was increased (Table 3) and could be associated with CT content. Several studies have demonstrated that inclusion of CT using *L. leucocephala*, reduces ruminal methane production. Tan et al [36] under *in vitro* conditions

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### Table 4. Effects of CT on ruminal pH, N-NH₃, and volatile fatty acids (VFA's) in crossbred heifers fed *Pennisetum purpureum* grass and increasing levels of *Leucaena leucocephala*

| Items                                    | Percent of *L. leucocephala* supplemented (% of DM) | SE  | Contrast |
|------------------------------------------|----------------------------------------------------|-----|----------|
|                                          | 0        | 20   | 40   | 60   | 80   | L   | Q    | C    |
| Rumen pH                                 | 6.8      | 6.9  | 6.7  | 6.8  | 6.8  | 0.10| ns   | ns   | ns   |
| Rumen N-NH₃ (mg/dL)                      | 15.3     | 22.0 | 27.3 | 34.0 | 36.5 | 0.07| ns   | **   | ns   |
| Volatile fatty acids concentration (mmol/L) |                                    |     |      |      |      |     |      |
| Acetate                                  | 52.6     | 54.8 | 53.4 | 52.1 | 54.4 | 5.31| ns   | ns   | ns   |
| Propionate                               | 13.6     | 13.1 | 12.8 | 12.2 | 12.8 | 1.14| ns   | ns   | ns   |
| Butyrate                                 | 6.1      | 6.1  | 5.6  | 5.3  | 5.4  | 0.47| ns   | ns   | ns   |
| Isobutyric                               | 0.4      | 0.36 | 0.4  | 0.37 | 0.3  | 0.04| ns   | ns   | ns   |
| Valeric                                  | 0.3      | 0.34 | 0.4  | 0.4  | 0.4  | 0.04| *    | ns   | ns   |
| Isovaleric                               | 0.2      | 0.2  | 0.2  | 0.2  | 0.2  | 0.04| ns   | ns   | ns   |
| Total VFA                                | 73.5     | 75.0 | 72.9 | 70.7 | 73.8 | 6.90| ns   | ns   | ns   |
| A:P                                      | 3.8      | 4.1  | 4.1  | 4.2  | 4.2  | 0.10| *    | ns   | ns   |
| Molar proportions of VFA (%)             |          |      |      |      |      |     |      |
| Acetate                                  | 71.4     | 72.9 | 73.1 | 73.5 | 73.5 | 0.67| *    | ns   | ns   |
| Propionate                               | 18.6     | 17.6 | 17.7 | 17.4 | 17.5 | 0.33| *    | ns   | ns   |
| Butyrate                                 | 8.5      | 8.3  | 7.7  | 7.6  | 7.5  | 0.41| ns   | ns   | ns   |
| Isobutyric                               | 0.5      | 0.5  | 0.5  | 0.5  | 0.5  | 0.05| ns   | ns   | ns   |
| Valeric                                  | 0.4      | 0.4  | 0.5  | 0.6  | 0.6  | 0.04| **   | ns   | ns   |
| Isovaleric                               | 0.3      | 0.3  | 0.3  | 0.3  | 0.3  | 0.06| *    | ns   | ns   |

CT, condensed tannins; NH₃, ammonia; DM, dry matter; SE, standard error; L, linear contrast; Q, quadratic contrast; C, cubic contrast. * p < 0.05; ** p < 0.01; ns, non-significant (p > 0.05).

### Table 5. Population of rumen protozoa (log₁₀ cells/mL) in crossbred heifers fed *Pennisetum purpureum* grass and increasing levels of *Leucaena leucocephala*

| Items                        | Percent of *L. leucocephala* supplemented (% of DM) | SE  | Contrast |
|------------------------------|----------------------------------------------------|-----|----------|
|                              | 0        | 20   | 40   | 60   | 80   | L   | Q    | C    |
| Population of rumen protozoa |                                    |     |      |      |      |     |      |
| Holotrich (log₁₀)            | 4.0      | 4.1  | 4.1  | 3.9  | 4.1  | 0.18| ns   | ns   | ns   |
| Entodiniomorph (log₁₀)       | 5.5      | 5.3  | 5.5  | 5.5  | 5.5  | 0.07| ns   | ns   | ns   |
| Total protozoa (log₁₀)       | 9.5      | 9.4  | 9.6  | 9.3  | 9.6  | 0.19| ns   | ns   | ns   |

DM, dry matter; SE, standard error; L, linear contrast; Q, quadratic contrast; C, cubic contrast. * p < 0.05; ** p < 0.01; ns, non-significant (p > 0.05).
found a linear reduction in CH₄ production when they included 20%, 30%, 40%, 50%, and 60% of L. leucocephala, reporting a reduction of 63% in CH₄ production when the level of incorporation of the legume was 60% compared to the control treatment (0% L. leucocephala). Rira et al [10] included 75% and 100% of L. leucocephala in an in vitro system and found reductions of 27.0% and 31.5% in CH₄ emissions, while Molina-Botero et al [37] incubated 100% of L. leucocephala and found a reduction of 31% in CH₄ emissions when expressed as g/kg DM. Delgado et al [30] included 27% of L. leucocephala in sheep rations and observed a reduction of 15.6% in CH₄ emissions (L/kg DMI).

In the experiment hereby described, a reduction of 39.4% was recorded in CH₄ production with 40% of DM ration of L. leucocephala fed to growing heifers housed in open-circuit respiration chambers. Recent work by Soltan et al [8] included 35% of L. leucocephala in sheep rations and found a reduction in CH₄ production of 14.3% when expressed as L/kg of digestible organic matter, this result is lower than that found in the present experiment with 40% of inclusion of L. leucocephala (32.1% less CH₄). Also, these results differ from results reported by Kennedy and Chamrley [4] who registered a reduction (L CH₄/kg DMI) in emissions of 13.4% and 20.5% when they included 22% and 44% of L. leucocephala in cattle rations.

A higher reduction in methane production per kg of DMI was found in the experiment hereby described with 20% and 40% incorporation of L. leucocephala (26.7% and 39.4% less CH₄/kg DMI, respectively) and when results were expressed as L CH₄/kg DOMI, our results were similar to the values reported by Kennedy and Chamrley [4] with 22% and 44% of L. leucocephala in the ration (19.2% and 28.2% less CH₄/kg DOMI). However, our results with 80% of inclusion of L. leucocephala in the diet differed from that reported by Dias-Moreira et al [34] who added 82% of L. leucocephala (40 g CT/kg DM) to sheep rations and found a reduction of 26% in rumen methane production. This could be associated with the CT concentrations on the diet. Grainger et al [33] included 1.46% CTs per kg dry matter and found a reduction of 22.3%, and in our present work the inclusion of 1.64% CT per kg of ration DM (80% of L. leucocephala), induced a reduction in methane production of 61.3%.

Min et al [31] found that the inclusion of 2% of CT as DM exerted a reduction of 31% in ruminal CH₄ production. In several experiments using plants containing CTs, similar reductions in CH₄ emissions to those found in the experiment hereby described have been registered. Puchala et al [35] found that goats fed Lespedeza cuneata containing 153 g CT/kg DM reduced enteric CH₄ emissions by 57.1%. Similar data were reported by Animut et al [38] in goats eating 200, 447, and 613 g DM of Lespedeza striata and found a reduction in CH₄ emissions of 32.8%, 47.3%, and 58.4%, respectively.

This lower digestibility [4,8] could reduce CH₄ synthesis because the fibrous component on the diet with legume was lower due to their lower amounts of lignified components compared with tropical grasses, so the time of residence in the rumen is shorter [39,40,7]. In this experiment, a reduction in rumen protozoa population was not observed, probably due to the low concentration of CT (<20 g/kg DM; [41]), however, several workers have mentioned that CT can reduce protozoa population but furthermore, they can also reduce the enzymatic activity of protozoa and bacteria [10].

**Rumen pH and molar proportions of volatile fatty acid**

Rumen pH was unaffected by the level of inclusion of L. leucocephala due to the fact that the forages had a high content of cellulose. The increase in the concentration of rumen NH₃-N is related to the increase in CP digestibility and it was observed that rumen ammonia concentration in the treatments with inclusion of L. leucocephala was above values considered as normal (15 to 20 mg/dL). This is contrary to that reported by Traiyakun et al [42], who included 0%, 25%, 50%, and 75% of L. leucocephala in goat rations and found no effect on rumen NH₃-N concentrations. Osakwe and Steingass [43] observed a reduction in rumen NH₃-N concentrations as the level of L. leucocephala was increased from 0%, 25%, and 50% of ration dry matter, claiming this was due to the reduction in rumen degradation of CP [44]. In the present study, rumen concentrations of acetic, propionic and butyric acids (mmol/L) were unaffected; however, when acetic acid is expressed as a percentage relative to the other VFA, its contribution increased as the level of L. leucocephala in the ration was increased. On the contrary, percentage of propionic acid was linearly reduced as L. leucocephala in the ration was increased, and the same trend was observed for the proportion of butyric acid. It was observed that the ratio acetic:propionic acid was increased as the level of L. leucocephala in the ration was augmented. These results agree with those reported by Tan et al [36] who under in vitro conditions incorporated 0%, 20%, 30%, 40%, 50%, and 60% L. leucocephala, and observed a linear increase in the concentration of acetic acid, but a reduction in the proportion of propionic acid and an increase in the acetic:propionic ratio in the rumen. Our results are also supported by Tiemann et al [45], who evaluated the molar proportions of VFA under in vitro conditions when they mixed L. leucocephala with P. purpureum and found that L. leucocephala resulted in a higher amount of acetic acid but reduced the concentration of propionic acid. Rira et al [10] evaluated in vitro inclusions of 25%, 50%, 75%, and 100% L. leucocephala and they did not find any effect on the molar proportions (mol/100 mol) of acetic and propionic acids. However, when they included 44% of L. leucocephala in Texel sheep rations, a reduction of 5.6% in acetic acid and an increase of 22.8% in the concentration of propionic acid were found. The results of other studies also differ; Soltan
et al [44] found that at levels of 50% of *L. leucocephala*, total concentration of VFA was increased by 17.3%, contrary to that found in the present experiment where no effect was observed on any of the main VFA in the rumen. Authors such as Tiemann et al [45] pointed out that the source of CT is an important factor which determines the effect on ruminal fermentation, however, the concentrations, molecular weight and the chemical structure of CTs are all important factors which determine their effect on the molar proportions of VFA [45,46]. It is still possible that the low concentrations of CT (1.64% CT/kg DMI) at the maximum level (80% of ration DM as *L. leucocephala*) of incorporation of the legume, had no effect in the concentrations of VFA when expressed as mmol/L.

**Rumen protozoa population**
The precise mechanism whereby CTs reduce CH$_4$ production is still uncertain, however CTs have the capacity to reduce protozoa [46] and methanogenic archaea populations [7]. CT could also reduce CH$_4$ production by means of indirect mechanisms as the reduction in digestibility of dietary components [42] by forming chemical complexes with carbohydrates and proteins as it was found in this experiment in which a reduction in OMD was recorded as the level of incorporation of *L. leucocephala* was increased as well as by linearly reduced NDFD and ADFD (Table 2).

Several studies indicate that CTs of tropical trees or legumes have the capacity of reducing rumen protozoa [46,47,36,35], and bacteria [7] populations. It is known that protozoa populations are responsible for 37% of CH$_4$ emissions [48] and they can degrade up to 50% of the fibrous fraction of feedstuffs [49,50]. Working with protozoa-free animals, Yañez-Ruiz et al [48] found a reduction in CH$_4$ emissions. In the experiment hereby described, no effect was detected of CT contained in *L. leucocephala* on rumen populations of holotrich and entodiniomorph, similar data was reported by Sliwinski et al [51] who fed 0.5% and 1.0% CT/kg DM and did not find any effect on these populations in the rumen. These findings are similar to those reported by Rira et al [10] who incorporated 44% of *L. leucocephala* as a source of CT and did not find any effect on rumen protozoa population. However, under *in vitro* conditions, Tan et al [26] included 20%, 30%, 40%, 50%, and 60% of *L. leucocephala* and found a linear reduction in total and ciliate protozoa. Also, Galindo et al [47] found that at inclu- sions of 30% of *L. leucocephala*, rumen protozoa population was reduced by 39.4%. Puchala et al [35] supplemented goats with *Lespedeza cuneata* (20% CT/kg DM) and observed a reduction of 65% in ciliate protozoa compared to the treatment when only grass was fed. These findings suggest that small doses of CT in *L. leucocephala* (1.64% CT/kg DMI; 80% of *L. leucocephala* in ration DM) have no effect on rumen protozoa population.

**CONCLUSION**
As a source of CT, the tropical legume tree *L. leucocephala* has the capacity to reduce enteric methane emissions in cattle without affecting DM and OM intake, while providing a considerably improved intake of rumen fermentable N for the growth of the microbial population. Inclusion of 80% of *L. leucocephala* in the diet of heifers fed low-quality tropical forages has the capacity to reduce up to 61.3% the methane emission without affecting DMI and OMI, protozoa population and the molar concentration of VFAs.

**CONFLICT OF INTEREST**
We certify that there is no conflict of interest with any financial organization regarding the material discussed in the manuscript.

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