Alterations in Energy Partitioning and Methane Emissions in Murciano-Granadina Goats Fed Orange Leaves and Rice Straw as a Replacement for Beet Pulp and Barley Straw

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Simple Summary: Reducing methane emissions in ruminants with the recycling of agro-industrial by-products is of great importance today. Pruning waste from citrus trees is currently burned or incorporated into soil. Regarding rice straw, this waste is traditionally eliminated through controlled burning, releasing into the atmosphere large amounts of greenhouse gases as well. The aim of this study was to convert this recovered waste into a new animal feed capable of reducing methane emissions in ruminants. The interest in using waste by-products for ruminant nutrition is increasing. Therefore, we replace the beet pulp and cereal straw from dry-nongravid goats’ diet with orange leaves and rice straw with the objective of studying their effect upon intake, digestibility, energy efficiency, carbon and nitrogen balance, and methane emissions.

Abstract: Considering the huge quantities of crops by-products and pruning waste such as rice straw and citrus leaves produced annually worldwide, and their potential pollution capacity, recycling as feed for livestock is an alternative. The objective was to study these by-products effect on energy balance and methane emissions in 10 Murciano-Granadina goats at maintenance. The control diet (CTR) included barley straw and beet pulp while the experimental diet (ORG) consisted of rice straw and orange leaves. Differences were found for energy intake (248 kJ/kg of BW 0.75 greater for CTR than ORG). The intake of metabolizable energy was 199 kJ/kg of BW 0.75 lower in ORG than CTR, and the energy efficiency was higher with CTR (0.61) than ORG (0.48). Protein retained in the body was 9 g/goat greater with CTR than ORG, and fat retention in the body was approximately 108 g/goat greater with CTR than ORG. Despite more unfavorable energy balance in response to feeding ORG than CTR, the retention of body energy was always positive. Reductions in CH4 emissions were detected when goats were fed ORG diet (from 22.3 to 20.0 g/d). Overall results suggested that feeding orange leaves and rice straw was effective in reducing CH4 emissions without adversely affecting energy balance.

Keywords: orange leaves; rice straw; methane emissions; goats; maintenance

1. Introduction

Ecological leftovers such as crop residues, food wastes, and agro-industrial by-products constitute human-inedible feed biomass [1]. Huge quantities of waste biomass are generated as agricultural and food industry by-products, explaining around 30% of global agricultural production [2]. Usage of such “less food-competing” biomass as feedstuff in animal diets is a potential approach to diminish food feed competition that can also help mitigate environmental effects of livestock.

The combination of livestock with crop production is a manner of creating sustainable farming systems that purpose to optimize resource usage. Crop residues and fibrous...
agro-industrial by-products do not enclose the nutrient balance needed to support efficient ruminal fermentation and animal performance. In countries with more specialized livestock systems, rice straw is considered of low nutritional value and is burned, however in developing countries where livestock are integrated with cropping, it is a valuable resource [3]. Despite this, in many areas such as Southeast Asia and Mediterranean countries, cereal straw is used as fodder providing fiber.

Rice (488 million tons of husked rice) is the world’s third major cereal crop after corn and wheat but produces the greatest amount of crop residues in the form of rice straw [4,5]. For instance, Spain produces 525,504 tons/year of rice straw [6]. Other important horticultural by-products in Spain come from citrus trees. Spain is one of the most important citrus production regions in the world and is acclaimed for cultivation of oranges and mandarins. Consistent with [7], Spain generates 1.87 million tons per year of pruning waste (on dry matter basis), of which almost 500 g/kg is leaves and 500 g/kg wood [7].

The main limitation of rice straw and orange leaves as feedstuff is the low nitrogen content and digestibility; these by-products are composed of cell wall constituents with slight soluble cell components, thus have to be degraded via microbial fermentation. The productivity of animals fed these waste by-products could increase with supplementation with a source of protein or N. Thus, these crops and pruning wastes could be converted into a valuable feed product for ruminants [3,8]. Furthermore, due to its high essential oils content, orange leaves could be advantageous for reducing methane productions from ruminants [9].

Studies carried out in livestock are scarce. With the use of citrus pulp, there are many studies, but not with citrus leaves, and as an example we can point to some work in broilers [10] and goats [11]. With rice straw there are more studies, mainly from Southeast Asia [12–14]. However, there are no studies that combine citrus leaf and rice straw in the same ration.

Considering the large amount of rice straw and citrus leaves produced annually worldwide, and their potential pollution capacity, especially when burned, revalorization and reutilization of these by-products as a complementary feed sources for livestock is a relevant subject in the exploration of the circular economy. Thus, the objective of this study was to replace beet pulp and cereal straw with orange leaves and rice straw in the diet to fed dry-non-pregnant goats with the aim of studying their effect upon intake, digestibility, energy efficiency, carbon (C) and nitrogen (N) balance, and CH₄ emissions.

2. Materials and Methods

2.1. Ethics Statement

Experimental procedures were approved (with reference 2017/VSC/PEA/00182) by the Animal Use and Care Committee of the Universitat Politècnica de València (UPV, Valencia, Spain). Besides, the procedures followed the practice codes for animals in experimental work advised by [15].

2.2. Animals and Diets

The experiment was managed at the Experimental Farm belonging to the UPV Animal Science Department (UPV), Valencia (Spain). Ten dry, non-pregnant multiparous Murciano-Granadina goats were chosen and assigned to two homogenous groups; five goats per group based on alike body weight (BW; 44.1 ± 3.65 kg) in a cross-over design (2 treatments crossed with 2 periods). The breed used was the Murciano-Granadina as it is a typical native breed of this geographical area of Spain. Females have been chosen because the production system for this breed revolves around milk production and the mothers are in standardized lactations at 210 days. During the drying phase it is important to have a maintenance diet for the animals, and ideally it is to be able to use crop residues and fibrous agro-industrial by-products. Treatments involved two different pelleted diets (Table 1). The control diet (CTR) included barley straw and beet pulp and the experimental diet (ORG) consisted of rice straw and orange leaves. Both diets included sunflower meal as
protein source and sugarcane molasses to get a good pellet texture. Diets followed nutrient recommendations by Calsamiglia et al. [16] for goats at maintenance. The feed offered was 2 kg, which was distributed twice a day; at 0900 h and at 1600 h. Water and mineral vitamin blocks were free for all goats.

Table 1. Ingredients and chemical composition of diets.

| Item                              | Diet 1 | Diet 2 |
|-----------------------------------|--------|--------|
| **Ingredients, % DM**             |        |        |
| barley straw                      | 27     | 0      |
| rice straw                        | 0      | 24     |
| beet pulp                         | 45     | 0      |
| orange leaves                     | 0      | 45     |
| sunflower meal 28                 | 20     | 23     |
| sugarcane molasses                | 7      | 7      |
| calcium carbonate                 | 0.7    | 0.5    |
| sodium chloride                   | 0.3    | 0.3    |
| **Chemical composition, % of DM** |        |        |
| Dry matter                        | 96     | 96     |
| Organic matter                    | 90     | 80     |
| Ash                               | 6      | 16     |
| Crude Protein                     | 12     | 14     |
| Ether rextract                     | 1.1    | 1.1    |
| Neutral detergent fiber           | 50     | 44     |
| Acid detergent fiber              | 31     | 29     |
| Acid detergent lignin             | 5      | 7      |
| Hemicellulose                     | 19     | 15     |
| Cellulose                         | 26     | 23     |
| NFC 2                             | 31     | 26     |
| Carbon                            | 45     | 42     |
| Nitrogen                          | 2      | 2      |
| Gross energy, MJ/kg DM 3          | 17     | 16     |

1 CTR = control; ORG = orange leaves and rice straw. 2 NFC = non fibrous carbohydrate; 100 – (NDF + ash + CP + EE). 3 DM = Dry matter.

2.3. Experimental Procedure and Sampling

Apparent total-tract digestibility, partition of energy, balance of C and N, and CH4 emission were determined. The crossover design experiment consisted of two 38-day periods and at the beginning and end of the experimental period the BW were obtained (TruTest FastWeight Crate, Datamars Livestock, AFB Farm supplies, Cheshire, UK). Goats were allocated in pens and fed the experimental diets for 14 days during adaptation, then allocated for an additional 7 days to individual metabolism crates at thermoneutrality (20–23 °C; Hobo probe, ONSET data loggers, Cape Cod, MA, USA). During the next 5 consecutive days, daily feed offered, orts, and total fecal and urine output were recorded and collected for each animal. Plastic buckets for urine collection contained 100 mL 10% (v/v) of H2SO4. Representative samples (10%) of diet, feces, and urine were stored at −20 °C and pooled for chemical analysis. On the last day of the digestibility trial and before the morning feeding, ruminal liquor samples were sampled using a stomach tube and pH was immediately determined (portable pH meter Model 265A, Orion Research Inc., Beverly, MA, USA). A ruminal liquor sub-sample was acidified with 50% H2SO4 and frozen until ammonia-N (NH3-N) determination. The other ruminal liquor sub-sample was mixed with H3PO4 and kept frozen until volatile fatty acids (VFA) analysis. Jugular blood was sampled (10 mL tubes treated with EDTA) and centrifuged for plasma separation and then stored at −20 °C. Afterward, goats were moved for 2 days to pens prior to gas exchange procedures.
Gas exchange determination (oxygen: \( \text{O}_2 \); carbon dioxide: \( \text{CO}_2 \) and methane: \( \text{CH}_4 \)) was quantified for each goat during a 24 h period. The system used for gas exchange quantification was based on an indirect calorimetric system designed for small ruminants. The respirometry system was armed with 2 ventilated head-hoods, 2 flow-meters (Thermal Mass Flowmeter Sensyflow VT-S, ABB, Alzenau, Germany), and 2 air suctions afforded by centrifugal fans (CST60 Soler Palau Inc., Parets del Vallès, Barcelona, Spain). Gas analyzer measured \( \text{O}_2 \) following paramagnetic principle, and \( \text{CH}_4 \) and \( \text{CO}_2 \) were measured following the infrared principle (Easyflow Gas Analyzer, model 3020, ABB, Alzenau, Germany). Reference gases were used to calibrate the gas analyzers before each test, and for a deep and detailed explanation of the portable open-circuit respirometry system used in this study it is advisable to consult the following works: Fernández et al. [17–19]. Further, the entire system was gravimetrically calibrated by injecting pure gas \( \text{N}_2 \) and \( \text{CO}_2 \) into the head-hood [20] (precision scale MOBBA mini-SP 0.2–30 kg, Industrial Weighing System, Barcelona, Spain). With the calibration of the entire system, the calibration factors were obtained according to Brockway et al. [21]. The \( \text{O}_2 \) consumption and the \( \text{CH}_4 \) and \( \text{CO}_2 \) production were calculated as described by Aguilera [22]. Atmospheric air sample (the blank for calculations) was sampled and analyzed before and after each determination made in goats.

2.4. Chemical Analysis

Feed, feed refusals, and fecal samples were first dried in a forced-air-oven at 55 °C for 48 h and urine was lyophilized. The dry matter (DM) of diets, orts, and feces was obtained by oven-drying at 102 ± 2 °C for 24 h (no. 934.01, AOAC [23]). Ash concentration (no. 942.05, AOAC [18]) and organic matter (OM) determination were attained by incineration of a sample in an electric muffle furnace at 550 °C for 6 h. Feed, refusals, and feces were analyzed for neutral detergent fiber (aNDF) and acid detergent fibre (ADF) using the ANKOM Fiber Analyzer (A220, ANKOM Technologies, Fairport, NY, USA). The aNDF was NDF assayed with amylase (heat-stable) and expressed inclusive of residual ash. The lignin content was obtained by solubilization of cellulose with \( \text{H}_2\text{SO}_4 \), following procedures of Van Soest [24]. The sample was subjected to an acid hydrolysis before the ether extract (EE) was detached with petroleum ether (Soxhlet System HT Tecator, Hillerød, Denmark; 1047 Hydrolyzing Unit and 1043 Extraction Unit) using method no. 920.39 AOAC [23]. The Dumas principle (method no. 968.06, AOAC [23]) were used to determine the C and N amounts with the analyzer TruSpec CN; LECO Corporation, St. Joseph, MI, USA. The estimation of non-fibrous carbohydrate (NFC) of diets was obtained by difference, according to [25]. The gross energy (GE) content of the dried samples (feed, feces, and urine) was obtained by combustion in an adiabatic bomb calorimeter (Gallenkamp Autobomb; Loughborough, UK).

Determination of VFA in ruminal liquor samples were described by Jouany [26] using a gas chromatograph (Fisons 8000 series; Fisons Instruments SpA, Milan, Italy) and the fatty acid (FA) methyl esters of orange leaf lipids were followed as described by O’Fallon et al. [27]. A Focus Gas Chromatograph (Thermo, Milan, Italy) was used to analyze FA methyl esters. Both equipment’s were armed with a split/splitless injector and a flame ionization detector.

Metabolites in ruminal liquor, urine and blood plasma samples were analysed as follow. Briefly, glutamate and free amino groups and glutamate were determined according to Larsen [28]. Urine and plasma ammonium, total protein, urea, uric acid, albumin and glucose were analyzed agreeing to standard protocols (Siemens Diagnostics® Clinical Methods for ADVIA 1800). Plasma non-esterified fatty acids (NEFA) were attained using the NEFA C ACS-ACOD assay method and, \( \beta \)-hydroxy butyrate (BOHB) was realized as recommended by Harano et al. [29]. Both NEFA and BOHB analyses were performed on the ADVIA 1800 System.
2.5. Calculations

The metabolizable energy (ME) intake (MEI) was the difference between intake and excretion. Therefore, the difference among GE intake (GEI) and energy losses in feces (E_{feces}), urine (E_{urine}) and CH\textsubscript{4} (E_{methane}). The CH\textsubscript{4} energy value was 39.5 kJ/L CH\textsubscript{4}, as reported by Brouwer [30].

Heat production (HP; in kJ) was calculated from indirect calorimetry recording the gas exchange (O\textsubscript{2}, CO\textsubscript{2} and CH\textsubscript{4}; all gases measured in L), and urine N (N_{urine}; g), using the Brouwer’s equation [30]:

\[ \text{HP (kJ)} = 16.18 \times O_2 + 5.02 \times CO_2 - 2.17 \times CH_4 - 5.99 \times N_{urine}. \]

The energy retention (RE) in the body was calculated as MEI – HP. The non-protein respiratory quotient (RQ_{np}) was obtained following the next expression: RQ_{np} = (CO_2 - (N_{urine} \times 6.25 \times 0.774))/(O_2 - (N_{urine} \times 6.25 \times 0.957)).

Efficiency of use of ME was determined according to [31]; km = 0.287 \times q + 0.554, kf = 0.78 \times q + 0.006, kmf = (km \times kf \times 1.5)/(kf + 0.5 \times km) with q being the energy metabolisability (the ratio ME/GE in the diet). The km was the ME efficiency for maintenance, kf the ME efficiency for growth and fattening, and kmf the ME efficiency for maintenance and growth. Although goats were not lactating and not pregnant, we considered a production level of 1.5 due to the fact they were fed beyond maintenance to reach a positive energy balance with the objective to enter mating when photoperiod was adequate. Digestible energy, ME, and NE of the diets were also calculated. From C and N balance the protein (R_{protein}) and fat (R_{fat}) retained in the body were calculated (g) following the method proposed by McLean [20].

2.6. Statistical Analysis

Effects of replacement of barley straw and beet pulp with rice straw and orange leaves on intake, digestibility, ruminal fermentation, energy and C-N balances, and CH\textsubscript{4} emission were studied using a mixed model (nlme library and lme function) in R [32].

The statistical model used was:

\[ Y = \mu + D + T + D \times T + \text{goat} + \epsilon \]

where Y was the dependent variable, \(\mu\) was the overall mean, and D and T were the fixed effects of diet and period of time, and their interaction; goat was the random effect of goat; and \(\epsilon\) was the random error. Least squares mean were obtained and differences were considered significant at \(p < 0.05\).

3. Results

No significant effect was observed for period of time and their interaction with diet in the crossover design; thus, the effect of diet was only reported in tables. The calibration factor from indirect calorimetry was 1.0043 ± 0.00126 (\(n = 4\)) for O\textsubscript{2}, 0.9951 ± 0.00982 (\(n = 4\)) for CO\textsubscript{2} and 0.9655 ± 0.00623 (\(n = 4\)) for CH\textsubscript{4}.

3.1. Feed Intake, Digestibility, and Ruminal Fermentation

A difference in DMI (\(p < 0.05\)) was observed between diets (0.34 kg/d, lower in CTR than ORG) (Table 2). Despite greater intake in the group of goats fed ORG, the apparent digestibility was lesser (\(p < 0.05\)) compared with CTR except for EE and CP, where no differences were observed. No differences were observed in EE digestibility between ORG and CTR, but ORG diet was richer in essential oils and polyunsaturated fatty acids (PUFA) due to the orange leaf (Table 3).
Table 2. Body weight, feed intake, and apparent total tract digestibility (% of DM) of Murciano-Granadina dried goats (n = 10) by the type of diet.

| Item 1 | Diet 2 | SEM 3 | p-Value |
|--------|--------|-------|---------|
|        | CTR    | ORG   |         |
| BW     | 46.6   | 41.5  | 0.94    | 0.0049  |
| DMI, kg/d | 1.41 | 1.74  | 0.273   | 0.0001  |
| DM     | 67     | 46    | 2.1     | 0.0001  |
| OM     | 69     | 51    | 1.8     | 0.0001  |
| EE     | 45     | 46    | 1.4     | 0.7261  |
| CP     | 59     | 62    | 0.9     | 0.2073  |
| NDF    | 59     | 37    | 2.3     | 0.0001  |
| ADF    | 58     | 36    | 2.3     | 0.0001  |
| ADL    | 25     | 21    | 1.9     | 0.2710  |
| hemicellulose | 62   | 40    | 2.3     | 0.0001  |
| cellulose | 64  | 41    | 2.4     | 0.0001  |
| NFC    | 89     | 67    | 1.9     | 0.0001  |
| GE     | 68     | 50    | 1.8     | 0.0001  |

1 BW = body weight; BW<sup>0.75</sup> = metabolic body weight; DMI = dry matter intake; DM = dry matter; OM = organic matter; CP = crude protein; EE = ether extract; NDF = neutral detergent fiber; ADF = acid detergent fiber; NFC = non fibrous carbohydrate: 100 − (NDF + ash + CP + EE); GE = gross energy. 2 CTR = control; ORG = orange leaves and rice straw. 3 SEM = Standard error of the mean.

Table 3. Fatty acid profile from orange leaves (mg/100 mg).

| Fatty Acids | Orange Leaves |
|------------|--------------|
| C4:0       | 0.000        |
| C6:0       | 0.000        |
| C8:0       | 0.000        |
| C10:0      | 0.001        |
| C11:0      | 0.000        |
| C12:0      | 0.017        |
| C13:0      | 0.122        |
| C14:0      | 0.034        |
| C14:1      | 0.000        |
| C15:0      | 0.002        |
| C16:0      | 0.274        |
| C16:1      | 0.001        |
| C17:0      | 0.010        |
| C17:1      | 0.000        |
| C18:0      | 0.039        |
| C18:1n9t   | 0.000        |
| C18:1n9c   | 0.057        |
| C18:1n7    | 0.010        |
| C18:2n6t   | 0.000        |
| C18:2n6c   | 0.326        |
| C20:0      | 0.013        |
| C18:3n6    | 0.006        |
| C20:1      | 0.000        |
| C18:3n3    | 0.294        |
| C18:1n9t   | 0.000        |
| C18:1n9c   | 0.057        |
| C18:1n7    | 0.010        |
| C18:2n6t   | 0.000        |
| C18:2n6c   | 0.326        |
| C20:0      | 0.013        |
| C18:3n6    | 0.006        |
| C20:1      | 0.000        |
| C18:3n3    | 0.294        |
| C24:0      | 0.022        |

1 CLA = conjugated linoleic acid.
Average ruminal pH never fell below 6.2 (Table 4), suggesting that values obtained were sufficiently high to maintain normal ruminal fermentation [33]. Although stomach tube is a suitable non-invasive technique for ruminal fluid sampling, it is prone to saliva contamination, which would increase the pH in the sample [34]. The NH₃-N was higher in ORG than CTR and no differences were found for total VFA between diets (51.3 mM/L on average). However, differences were observed \((p < 0.05)\) for propionic, isobutyric, and isovaleric acid.

Table 4. Ruminal parameters: pH, NH₃-N, and volatile fatty acids (VFA) of Murciano-Granadina dried goats \((n = 10)\) by the type of diet.

| Item                        | CTR | ORG | SEM 3 | \(p\)-Value |
|-----------------------------|-----|-----|-------|-------------|
| pH                          | 6.5 | 6.9 | 0.05  | 0.2155      |
| NH₃-N, mg/dL                | 12.3| 15.1| 0.44  | 0.0125      |
| Total VFA, mM/L             | 54.2| 48.4| 3.11  | 0.6212      |
| Individual VFA, mM/L        |     |     |       |            |
| Acetic acid                 | 32.7| 31.9| 1.73  | 0.8214      |
| Propionic acid              | 11.8| 8.1 | 0.87  | 0.0245      |
| iso Butyric acid            | 0.30| 0.66| 0.065 | 0.0001      |
| Butyric acid                | 8.30| 5.93| 0.748 | 0.1169      |
| Isovaleric acid             | 0.24| 0.79| 0.103 | 0.0004      |
| n-Valeric acid              | 0.79| 0.90| 0.058 | 0.3376      |
| n-Caproic acid              | 0.08| 0.10| 0.012 | 0.5665      |
| Heptanoic acid              | 0.000| 0.002| 0.0008 | 0.1479      |

\(^1\)NH₃-N = ammonia nitrogen. \(^2\)CTR = control; ORG = orange leaves and rice straw. \(^3\)SEM = Standard error of the mean.

3.2. Energy Balance

Due to the differences found between DMI, differences \((p < 0.05)\) were found for GEI (1406 and 1654 kJ/kg of BW\(^{0.75}\) for CTR and ORG, respectively) (Table 5). Feces and urine energy losses were also bigger \((p < 0.05)\) in response to feeding ORG compared with CTR. At the same time, this effect might partly explain the reduction \((p < 0.05)\) in energy losses as CH₄ (69 vs. 66 kJ/kg of BW\(^{0.75}\) for CTR and ORG, respectively).

Due to greater losses in feces and urine with the ORG diet, the MEI was 199 kJ/kg of BW\(^{0.75}\) lower \((p < 0.05)\) compared with CTR. Despite the differences in GEI between the diets, no significant differences were found for HP \((482 \text{ kJ/kg of BW}^{0.75}, \text{ on average})\) and RE was positive with both treatments. Greater \((p < 0.05)\) retention was detected in CTR compared with ORG \((224 \text{ kJ/kg of BW}^{0.75}\) greater in CRT than ORG). Change was not found for RQnp, with an averaged value of 1.01.

Significant differences \((p < 0.05)\) were observed for ME efficiencies. Efficiency of ME for maintenance is defined as km and, following the INRA [31] equations, a greater value was observed in response to feeding CTR compared with ORG \((0.73 \text{ vs. 0.67 for CTR and ORG, respectively})\). The ME efficiency for growth and fattening \((kf)\) according to INRA [31], was 0.47 and 0.31 for CTR and ORG, respectively. Because goats were in positive energy balance, we obtained the combined efficiency of maintenance, growth, and fattening \((kmf)\); 0.61 and 0.48 was observed for CTR and ORG, respectively.

When energy balance was expressed as percentage of GE intake, differences \((p < 0.05)\) were found. Greater values were observed with CTR than ORG. The DEI/GEI was 18 points greater in CTR compared with ORG, and MEI/GEI was 21 points greater in CTR compared with ORG. When expressed per kg of DM, differences between diets \((p < 0.05)\) were also detected for DE, ME, and NE.
Table 5. Energy balance (kJ/kg of BW\(^{0.75}\) and day) of Murciano-Granadina dried goats (\(n = 10\)) by the type of diet.

| Item \(^1\) | Diet \(^2\) | SEM \(^3\) | \(p\)-Value |
|------------|------------|----------|-------------|
| GEI        | 1406       | 1654     | 46.4        | 0.0060      |
| \(E_{\text{feces}}\) | 447        | 824      | 38.5        | 0.0001      |
| \(E_{\text{urine}}\) | 51         | 124      | 7.4         | 0.0001      |
| DEI        | 959        | 830      | 31.3        | 0.0385      |
| \(E_{\text{methane}}\) | 69         | 66       | 2.0         | 0.0224      |
| MEI        | 839        | 640      | 36.0        | 0.0043      |
| HP         | 469        | 494      | 8.8         | 0.1653      |
| RE         | 370        | 146      | 36.5        | 0.0013      |
| RQ\(\text{np}\) | 1.03    | 0.99     | 0.015       | 0.1254      |

**ME efficiencies**

\[ \begin{align*}
\text{km} & = 0.73 \quad 0.67 \quad 0.014 \quad 0.0302 \\
\text{kf} & = 0.47 \quad 0.31 \quad 0.012 \quad 0.0001 \\
\text{kmf} & = 0.69 \quad 0.45 \quad 0.011 \quad 0.0001 \\
\% \text{GEI} & \\
\text{DEI} & = 68 \quad 50 \quad 2.8 \quad 0.0001 \\
\text{MEI} & = 60 \quad 39 \quad 2.3 \quad 0.0001 \\
\text{RE} & = 26 \quad 9 \quad 2.6 \quad 0.0001 \\
\end{align*} \]

**Diet energy value, MJ/kg DM**

\[ \begin{align*}
\text{DE} & = 12 \quad 8 \quad 0.3 \quad 0.0001 \\
\text{ME} & = 11 \quad 6 \quad 0.1 \quad 0.0001 \\
\end{align*} \]

\(^1\) GEI = gross energy intake; \(E_{\text{feces}}\) = energy losses in feces; \(E_{\text{urine}}\) = energy losses in urine; \(E_{\text{methane}}\) = energy losses in methane; DEI = digestible energy intake; ME = metabolizable energy; MEI = metabolizable energy intake; HP = heat production; RE = energy retention; RQ\(\text{np}\) = respiration quotient corrected to nitrogen; km = ME efficiency for maintenance; kf = ME efficiency for fattening; kmf = ME efficiencies for maintenance and fattening.

\(^2\) CTR = control; ORG = orange leaves and rice straw.

\(^3\) SEM = standard error of the mean.

3.3. Carbon and Nitrogen Balance

With the exception of \(C_{\text{CO2}}\), differences (\(p < 0.05\)) were found in C balance (Table 6). Following the trend observed for intake and apparent digestibility, C in feces and urine was 11.4 and 1.6 g/kg of BW\(^{0.75}\) greater with ORG compared with CTR, respectively. Losses in C from \(\text{CH}_4\) were lower with ORG compared with CON (0.03 g/kg of BW\(^{0.75}\)), and total excretion of C was 14 g/kg of BW\(^{0.75}\) greater with ORG than CTR. With respect to N, goats consumed 0.8 g/kg of BW\(^{0.75}\) more with the ORG than CTR diet. However, excretion was 0.86 g/kg of BW\(^{0.75}\) greater with ORG than CTR.

3.4. Metabolites

Ruminal, urine, and plasma metabolites primarily related to energy and protein metabolism did not differ greatly in response to diets, in part due to high variability (Table 7). Urea was greater with ORG than CTR in ruminal fluid and urine but non-significant, and difference (\(p < 0.05\)) was found in plasma. Uric acid in urine was higher (\(p < 0.05\)) in response to feeding ORG than CTR, and glutamate was higher (\(p < 0.05\)) with CTR than ORG. Greater (\(p < 0.05\)) branched-chain amino acid concentrations were observed in CTR than ORG and, no differences were detected for BOHB and NEFA.

3.5. Methane Emissions

The \(\text{CH}_4\) emission was shown in Table 8. Compared with the CTR diet, goats fed ORG emitted significantly (\(p < 0.05\)) fewer \(\text{CH}_4\) emissions (22.3 vs. 20.0 g/d for CTR and ORG, respectively). The Ym for both diets (\(\text{CH}_4\) conversion factor defined as \(E_{\text{methane}}/\text{GEI}\)) was 4.9 and 4.0% with CTR and ORG, respectively (\(p < 0.05\)). In the current study, when \(\text{CH}_4\) emission was expressed over DM, OM intake, and fiber, statistical differences (\(p < 0.05\)) remained.
Table 6. Daily balance of carbon and nitrogen (g/kg de BW\(^{0.75}\)) of Murciano-Granadina dried goats (\(n = 10\)) by the type of diet.

| Item                  | Diet | SEM  | \(p\)-Value |
|-----------------------|------|------|-------------|
|                       | CTR  | ORG  |             |
| \(C_{\text{intake}}\) | 35.69| 45.13| 1.296       | 0.0001      |
| \(C_{\text{fees}}\)  | 11.93| 23.29| 1.129       | 0.0001      |
| \(C_{\text{urine}}\) | 1.21 | 2.85 | 0.172       | 0.0001      |
| \(C_{\text{CO2}}\)   | 13.35| 14.41| 0.258       | 0.1383      |
| \(C_{\text{CH4}}\)   | 0.94 | 0.91 | 0.008       | 0.0256      |
| \(C_{\text{excretion}}\) | 27.44| 41.46| 1.389       | 0.0001      |
| \(C_{\text{retained body}}\) | 8.25 | 3.67 | 0.555       | 0.0001      |
| \(N_{\text{intake}}\) | 1.56 | 2.36 | 0.085       | 0.0001      |
| \(N_{\text{fees}}\)  | 0.64 | 0.90 | 0.032       | 0.0001      |
| \(N_{\text{urine}}\) | 0.63 | 1.22 | 0.085       | 0.0001      |
| \(N_{\text{excretion}}\) | 1.27 | 2.12 | 0.100       | 0.0001      |
| \(N_{\text{retained body}}\) | 0.30 | 0.24 | 0.067       | 0.0123      |
| \(R_{\text{protein}}, \text{g/goat}\) | 33  | 25   | 3.3         | 0.0158      |
| \(R_{\text{fat}}, \text{g/goat}\)  | 169 | 62   | 12.6        | 0.0024      |
| \(R_{\text{protein}}, \text{g/goat}\) | 27.44| 41.46| 1.389       | 0.0001      |
| \(R_{\text{fat}}, \text{g/goat}\) | 8.25 | 3.67 | 0.555       | 0.0001      |

\(^1\) C = carbon; \(N = \) nitrogen; \(R = \) retention. \(^2\) CTR = control; ORG = orange leaves and rice straw. \(^3\) SEM = standard error of the mean.

Table 7. Metabolites in rumen liquor, urine, and plasma of Murciano-Granadina dried goats (\(n = 10\)) by the type of diet.

| Item                       | Diet | SEM  | \(p\)-Value |
|----------------------------|------|------|-------------|
|                           | CTR  | ORG  |             |
| \(\text{Rumen liquor}\)   |      |      |             |
| Free amino groups, mEq/L   | 1.28 | 1.02 | 0.099       | 0.2121      |
| Glutamate, mM              | 117  | 107  | 9.1         | 0.5945      |
| Urea, mM                   | 27.8 | 33.1 | 3.49        | 0.4826      |
| \(\text{Urine}\)          |      |      |             |
| Protein, mg/L              | 304  | 215  | 48.0        | 0.3844      |
| \(\text{Free amino groups, mM/L}\) | 13.7 | 17.0 | 1.99        | 0.4349      |
| Glutamate, mM/L            | 74   | 41   | 6.4         | 0.0024      |
| Urea, mM/L                 | 601  | 621  | 44.6        | 0.8345      |
| Uric acid, mM/L            | 224  | 471  | 80.8        | 0.0452      |
| Malate, mM/L               | 21.8 | 28.2 | 2.81        | 0.2787      |
| \(\text{Plasma}\)         |      |      |             |
| Protein, g/L               | 81   | 81   | 1.8         | 0.9836      |
| Albumin, g/L               | 35   | 36   | 0.8         | 0.8414      |
| Glutamate, µM/L            | 65   | 49   | 4.9         | 0.0351      |
| \(\text{Free amino groups, mEq/L}\) | 2.8  | 2.4  | 0.3         | 0.5040      |
| \(\text{Free Branched Chain Amino acids, mM/L}\) | 0.78 | 0.56 | 0.056       | 0.0423      |
| Urea, mM/L                 | 5.8  | 7.3  | 0.29        | 0.0020      |
| Glucose, mM/L              | 4.0  | 4.0  | 0.21        | 0.9449      |
| \(\text{BOHB}^{3}, \text{mM/L}\) | 0.34 | 0.31 | 0.026       | 0.5431      |
| \(\text{NEFA}^{4}, \text{mEq/L}\) | 111  | 159  | 23.4        | 0.3328      |

\(^1\) CTR = control; ORG = orange leaves and rice straw. \(^2\) SEM = standard error of the mean. \(^3\) BOHB = Beta-Hydroxybutyrate. \(^4\) NEFA = nonesterified fatty acids.
Table 8. Methane emission of Murciano-Granadina dried goats \((n = 10)\) by the type of diet.

| Item 1                                                                 | Diet 2 | SEM 3 | p-Value |
|------------------------------------------------------------------------|--------|-------|---------|
| CH\(_4\), g/d                                                         | CTR    | ORG   |         |
| ratio CH\(_4\)/CO\(_2\) in breath                                      | 22.3   | 20.0  | 0.482   | 0.035   |
| Ym                                                                    | 4.9    | 4.0   | 0.07    | 0.0018  |
| CH\(_4\)/DMi, g/kg                                                   | 16.0   | 11.6  | 0.577   | 0.0001  |
| CH\(_4\)/OMi, g/kg                                                   | 17.1   | 13.8  | 0.564   | 0.0025  |
| CH\(_4\)/NDF, g/kg                                                   | 32.4   | 26.7  | 1.055   | 0.0010  |
| CH\(_4\)/ADFi, g/kg                                                  | 52.1   | 40.0  | 1.788   | 0.0003  |
| CH\(_4\)/hemicellulose intake, g/kg                                  | 3.0    | 1.7   | 0.134   | 0.0001  |
| CH\(_4\)/cellulose intake, g/kg                                       | 61.9   | 51.6  | 1.998   | 0.0089  |

1 Ym = methane energy/gross energy intake; DMi = dry matter intake; OMi = organic matter intake; NDFi = neutral detergent fiber intake; ADFi = acid detergent fiber intake. 2 CTR = control; ORG = orange leaves and rice straw. SEM = standard error of the mean.

4. Discussion

Higher intake in the ORG diet was accompanied by lower digestibility. One of the reasons for the reduction in digestibility could be the higher ash content (16 vs. 6\%) in the ORG diet. Another reason was that EE in diet ORG was richer in PUFA and could affect the digestibility of the fiber. Thus, PUFA could be responsible for the reduction in fiber digestibility. Classical work from Palmquist [35] indicated that dietary PUFA were more likely to inhibit fiber degradability with a concomitant reduction in fermentation, possibly due to coating food particles and preventing bacterial attachment.

Average ruminal pH never fell below 6.2 suggesting that values obtained maintained normal ruminal fermentation [33]. The greater values of NH\(_3\)-N in response to feeding ORG indicated poor utilization of dietary protein and extra N in the rumen. Regarding VFA, propionic acid was greater in response to feeding CTR and isobutyric and isoalveval were greater in goats fed ORG. As this VFA is mainly generated during degradation of branched-chain amino acids, the greater isoalvevalic acid concentration observed in goats fed ORG suggested greater ruminal protein degradation. The greater NH\(_3\)-N in ORG diet could indicate an inefficient use of amino groups for ruminal protein synthesis [36]. Thus, the probable asynchrony between protein degradation and fiber carbohydrates appeared to be more pronounced with the ORG diet.

Greater GEI was found in ORG than CTR due that the higher DMI observed in ORG. The smaller digestibility with the ORG diet indicated greater energy losses in feces, suggesting that the bigger high intake is accompanied by a higher rate of digesta transit. It could be possible that feeding ORG sustained the positive effect of lipids, essential oils, and tannins in the diet on CH\(_4\) reduction [9]. Due to greater energy losses with the ORG diet, the MEI was lower (640 vs. 839 kJ/kg of BW\(^{0.75}\) for ORG and CTR, respectively). The average value proposed by AFRC [37] using 17 estimates of ME for maintenance (MEm) derived from feeding trials with goats was 438 kJ/kg BW\(^{0.75}\) per day. Agreeing with other studies, the variation between estimates was substantial; ranging from 365 to 530 kJ/kg BW\(^{0.75}\) per day. Thus, Luo et al. [38] reviewing studies conducted with goats (dairy, meat, indigenous and Angora goats breeds, and several animal categories; dry and no pregnant, lactating, male, growing, and wethers) estimated MEm and it varied extensively; from 422 to 501 kJ/kg BW\(^{0.75}\) per day. The NRC [39] adopted for goats the maintenance requirements estimated from [38]. INRA [31] with a q of 0.64 suggested 441 kJ/kg BW\(^{0.75}\) per day. Prieto et al. [40] working with adult and castrated male goats attained a value of 443 kJ MEm/kg BW\(^{0.75}\) per day, and Aguilera et al. [41] obtained 401 kJ/kg BW\(^{0.75}\) per day. The model obtained by Fernández [42] had values of 460 kJ MEm/kg BW\(^{0.75}\) per day. Averaging the studies mentioned above, the MEm would be 445 kJ/kg BW\(^{0.75}\) per day. As the MEI obtained was 839 and 640 kJ/kg BW\(^{0.75}\) per day for the CTR and ORG diet, respectively, diet CTR had more energy available for weight
recovery (395 and 196 kJ ME/kg BW\(^{0.75}\) per day for CTR and ORG, respectively). The value of ME expressed over kg of DM was 11 and 6 MJ ME/kg DM in response to feeding CTR and ORG, respectively. Therefore, the RE was 370 kJ/kg BW\(^{0.75}\) for CTR and 146 kJ/kg BW\(^{0.75}\) for ORG, suggesting that poor microbial protein synthesis coupled with lower content of glucogenic nutrients with the ORG diet (i.e., CTR diet contain beet pulp) did not favor the partitioning of ME into body tissue and, hence, the body fat deposition [43].

Maintenance costs represent the main component of total energy requirements, which underscores the importance for an accurate estimation. The km obtained was 0.73 vs. 0.67 for CTR and ORG, respectively. The AFRC [37] and NRC [39] systems for goats estimated ME efficiency for maintenance using the same equation, which varies from 0.64 (low-quality diets) to 0.75 (high-quality diets). Using indirect calorimetry and regression techniques, Prieto et al. [40] obtained a value of 0.73 with adult Granadina castrated males’ goats and Aguilera et al. [41] obtained a value of 0.67 with lactating Granadina goats. Thus, CTR appears to have been a better-quality diet with the km obtained and the effectiveness found being within the ranges that literature reported. The kmf was 0.61 and 0.48 for CTR and ORG, respectively. If calculating a ratio between RE and MEI from Table 5, the values obtained were 0.44 and 0.23 for CTR and ORG, respectively. The ME efficiency for maintenance and recovery of body reserves followed the same trend, and was greater in CTR than ORG. The ME requirements for body gain differs due to changes from ME to RE, depending on even if energy derived from feed (lipogenic or glucogenic nutrient as Van Knegsel [43] reported) or body fat mobilization [44]. Thus, with both diets, animals recovered energy despite the fact that the effectiveness of the diet was greater for CTR.

Efficiency of C retained over C ingested was 23% and 8% when feeding CTR and ORG, respectively. In another study with Murciano-Granadina dry and non-pregnant goats, feeding close to maintenance the value obtained was 5% [17]. Another study with lactating Muciano-Granadina goats generated a value ranging from 4% to 9% [45]. The ratio between retained N and ingested N was again lower when feeding ORG than CTR (19% vs. 10% for CTR and ORG, respectively), indicating that the amount of protein in the ORG diet was above the animal’s needs. In the study of Fernández et al. [17], the value obtained was 38%, and Romero et al. [45] reported values ranging from 12% to 15%. From the C and N retained, retention of protein and fat were estimated [20]. Differences were detected between diets and the R\(_{\text{protein}}\) was 9 g/goat greater in response to feeding CTR than ORG. The R\(_{\text{fat}}\) was approximately 108 g/goat greater with CTR than ORG.

Urea was greater with ORG than CTR in plasma, indicating lower feed protein efficiency [46]. Urine uric acid concentration was higher in response to feeding ORG than CTR, again indicating that feeding ORG led to poor microbial protein synthesis and greater excretion of N compounds. The higher glutamate with CTR than ORG likely contributed to glutamine synthesis across the animal tissues [28]. Branched-chain amino acids may be used for synthesis of muscle or milk protein, preserved into the cell for structural protein synthesis, used as precursor for different metabolic and catabolic processes, or passes unaltered into milk, blood, or lymph [47]. The greater branched-chain amino acid concentrations observed in response to feeding CTR than ORG could be indicative of greater utilization by other peripheral tissues such as adipose [48]. Additionally, the lack of differences for BOHB or NEFA suggested little effect of diets in whole-body energy balance [49].

The reduction in CH\(_4\) when feeding ORG was partly caused by decreasing overall carbohydrate digestion. Goats fed ORG emitted 2.3 g/d less CH\(_4\) than CTR, a response that agrees with the fact that FA from orange leaves have inhibitory effect on protozoa and cellulolytic bacteria that can cause shifts in fermentation patterns that reduce CH\(_4\) production [3,11]. In addition to the known negative effects of PUFA on CH\(_4\) production via direct toxic effects on ruminal microorganism and protozoa [50,51], secondary metabolites compounds in plants like tannins and essential oils from orange leaves in the ORG diet also could explain the mitigation of CH\(_4\) observed with this diet [9,52,53]. Thus, although the reduction in CH\(_4\) averaged 10%, differences in fat percentage between CTR and ORG averaged 1.1%. Because ruminants waste between 2 and 12% of their dietary GEI as CH\(_4\),
a diminution in CH$_4$ production represents an improvement in feed efficiency [54]. The Ym was 4.9 and 4.0% with CTR and ORG, respectively. Concerning the ORG diet, the FA profile of ORG (Table 3) probably had a negative effect on methanogens and fiber degradation, which was reflected in the lower NDF and ADF digestibility (Table 2). While CH$_4$ emissions were most commonly expressed in the literature over GEI, the other meaningful indicator was over DM or OM intakes. The CH$_4$ reduction was observed also when it was expressed relative to DM, OM, and different fiber fractions. Patra et al. [55] reported that some plant secondary metabolites such as saponins and tannins, which are present in orange leaves, may show an inhibitory effect on methanogenic activity. Thus, further research should be performed to better understand how secondary compounds in citrus residue directly impact ruminal microorganisms.

5. Conclusions

The present study provides data for energy, C-N balance, and CH$_4$ emissions in Murciano-Granadina goats to support maintenance when crops and pruning waste by-products were fed. The diet replaced barley straw and beet pulp with rice straw and orange leaves, keeping sunflower meal as the same source of protein. Despite the greater feed intake when supplementing orange leaves and rice straw, the digestibility and energy balance were lower. The MEI was 199 kJ/kg of BW$^{0.75}$ lower in response to feeding orange leaves and the value of ME was 6 and 11 MJ ME/kg DM for that diet and the control, respectively. Retention of energy was 224 kJ/kg of BW$^{0.75}$ greater with the control and positive with both diets. Greater efficiency of maintenance, growth, and fattening was detected when feeding the control (0.61) compared with orange leaves diet (0.48). The efficiency of C retained over C ingested was 15 points greater with the control diet, and the N efficiency 9 points greater with the control diet. Reduction in CH$_4$ emissions were detected with the orange diet (from 22.3 to 20.0 g/d). Thus, despite the lower energy, C and N balance efficiencies when orange leaves were supplemented, goats recovered body reserves, and, consequently, these fibrous by-products were utilized without detectable unfavorable effects on energy metabolism. The economic advantages and environmental impact of feeding orange leaves should be assessed.

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