Nutrient Digestibility and Gas Production of Some Tropical Feeds Used in Ruminant Diets Estimated by the in vivo and in vitro Gas Production Techniques

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Abstract: Some feedstuffs which used in ruminants diet (corn grain, soybean meal, wheat bran and alfalfa) were analyzed for chemical composition, apparent in vivo nutrient digestibility, in vitro fermentation gas production and metabolizable energy. Chemical composition of test feeds differed in nutrient contents. Initially apparent in vivo digestibility of alfalfa nutrients were obtained then digestibility of nutrients for the other test feeds were determined by difference method, using 16 Ghezel mature rams (mean weight of 43.9±4 kg). In vivo DM, CP, NDF and OM apparent digestibility were different among the test feeds (p<0.05). Regarding to the results, corn grain had a high DM and OM digestibility between test feeds and soybean meal had a high CP and NDF digestibility between test feeds (p<0.05). Cumulative gas production was recorded at 2, 4, 8, 12, 16, 24, 36, 48, 72 and 96 h of incubation and the equation of p = A (1-e-ct) was used to describe the kinetics of gas production. Potential gas production (A) and rates of gas production (c) differed (p<0.01) among feeds. Corn grain showed higher potential gas production (A) (326.5 mL g⁻¹ DM) and wheat bran had higher rate of gas production (c) (0.097 h⁻¹) than the other feeds, inverses alfalfa (257.6 mL g⁻¹ DM) and corn grain (0.048 h⁻¹) had lower potential gas production and rate of gas production than the other test feeds, respectively. The metabolizable energy (MJ kg⁻¹ DM) content of feeds was calculated using in vivo organic matter digestibility and gas production data. According to in vivo organic matter digestibility data, the ME values ranged from 9.2 in alfalfa to 13.3 MJ kg⁻¹ DM in corn grain. It was concluded that regarding to different chemical composition of test feeds, the in vivo digestibility, in vitro gas production and ME of feeds showed different values.

Key words: Feed evaluation, in vivo digestibility, gas production, metabolizable energy

INTRODUCTION

The nutritive value of a ruminant feed is determined by the concentrations of its chemical components, as well as their rate and extent of digestion. Determining the digestibility of feeds in vivo is laborious, expensive, requires large quantities of feed and is largely unsuitable for single feedstuffs thereby making it unsuitable for routine feed evaluation. In vitro methods provide less expensive and more rapid alternatives[17]. Digestibility may be directly determined in vivo or estimated by using in vitro procedures, which are cheaper and more convenient[5].

There are number of in vitro techniques available to evaluate the nutritive value of feeds at relatively low cost such as in vitro gas production technique. The gas measuring technique was considered to be a routine method of feed evaluation after the work of Menke et al.[20], where a high correlation between gas production in vitro and in vivo apparent digestibility was reported. Gas production techniques are based on the principle that anaerobic microbial digestion of carbohydrates releases gas (primarily CO₂ and CH₄) and VFA[21].

Metabolizable energy represents that portion of the feed energy that can be utilized by the animal[11]. In vivo Organic Matter Digestibility (OMD) is defined as the proportion of feed OM apparently digested in the total digestive tract. Organic matter digestibility is a measure of energy available to ruminants and is used in protein evaluation system[18]. Menke and Steingass[27] reported a strong correlation between Metabolizable Energy (ME) values measured in vivo and predicted from 24 h in vitro gas production and chemical composition of feeds. The in vitro gas production method has also been widely used to evaluate the energy value of several classes of feeds[15], particularly straws[22], agro-industrial by-products[19], compound feeds[23] and various tropical feeds[20].

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The objective of the present study was to determine apparent in vivo digestibility, gas production function and ME of some feedstuffs (corn grain, soybean meal, wheat bran and alfalfa).

MATERIALS AND METHODS

Experimental feeds: The samples of Corn Grain (CG), soybean meal (SBM), Wheat Bran (WB) and alfalfa (AA) were collected from dairy farm in northwestern of Iran. Alfalfa samples were not identified by maturity or variety. Samples of all test feeds for the gas production technique were milled through a 2.0 mm sieve and oven-dried at 80°C until constant weight and for chemical analyses they were milled through a 1.0 mm sieve.

Chemical analysis: Samples of feeds and feces were dried in an oven at 105°C for 24 h and the DM content calculated. Ground samples were analyzed for ash[4]. Determinations of N were conducted using the Kjeldahl method in an automated Kjelfoss apparatus (Foss Electric, Copenhagen, Denmark). Neutral-detergent fiber and ADF were determined by the detergent procedures of Van Soest et al.[66]. Acid-Detergent Insoluble Nitrogen (ADIN) was determined as nitrogen in Acid-detergent residue. Ether Extract (EE) was determined by extracting the sample with ether[4].

Apparent in vivo digestibility: Initially apparent in vivo digestibility of alfalfa nutrients were obtained then digestibility of nutrients for the other test feeds were determined by difference method[8]. Sixteen Ghezel rams (43.9±4 kg body weight) were used to measure the apparent in vivo digestibility of feeds. The animals were kept in metabolism stalls. Before the studies, all animals were sheared, dewormed. The alfalfa digestibility was measured in four rams fed alfalfa as ad libitum and other feeds digestibility were measured in twelve ram at the maintenance of body weight (38 g DM/W0.75; NRC [30]) with a diet comprising 600 g kg⁻¹ test feeds (CG, SBM and WB) and 400 g kg⁻¹ basal feed (alfalfa). The diet was offered twice a day at 08:30 and 15:30 h in equal amounts after collecting the refusals. All animals were given free access to mineral salt lick and water and throughout the experiment. Experimental period lasted for 21 days, comprising 14 days for adaptation to the diet and 7 days for total feces collection. Samples of feeds offered and feces were taken daily and bulked over the trial period. The dried and milled samples were used for the determination of dry matter, organic matter, crude protein, Neutral-detergent fiber and acid-detergent fiber. Digestibility of DM, OM, CP, NDF and ADF were then determined for each diet using equations given by Pond et al.[8].

In vitro gas production: Samples (300 mg) were weighed into 100 mL serum vial. Mc Dougall[25] buffer solution was prepared and placed in a water bath at 39°C. Rumen liquor samples were obtained from the two wethers that were fed on a diet comprising (DM basis), 550 g kg⁻¹ alfalfa hay, 400 g kg⁻¹ barely grain, 50 g kg⁻¹ wheat bran and 2 g kg⁻¹ lime stone at maintenance level[30]. Rumen fluid was collected after the morning feeding. Rumen fluid was pumped with a manually operated vacuum pump and transferred into pre-warmed thermos flask, combined, filtered through four layers of cheesecloth and flushed with CO₂. Each feed sample was incubated in triplicate with 20 mL of rumen liquor and buffer solution (1:2). Three vials containing only the rumen fluid/buffer solution and no feed sample was included with each test and the mean gas production value of these vials was termed the blank value. The vials were sealed immediately after loading and were affixed to a rotary shaker platform (lab-line instruments Inc Melors dark, USA) set at (120 rpm) housed in an incubator. Gas production was measured in each vial after 2, 4, 8, 12, 16, 24, 36, 48, 72 and 96 h of incubation using a water displacement apparatus[12].

Calculations and statistical analysis: Rate and extent of gas production was determined for each feed by fitting gas production data to the one component McDonald model: Y = A (1-e⁻<sup>ct</sup>), where y is the volume of gas produced at time t, A the potential gas production, c the fractional rate of gas production was determined for each feed by fitting gas production data to the one component McDonald model: Y = A (1-e⁻<sup>ct</sup>), where y is the volume of gas produced at time t, A the potential gas production, c the fractional rate of gas production. Parameters A and c were estimated by an iterative least square method using a non-linear regression procedure of the statistical analysis systems[33].

The metabolizable energy (MJ/kg DM) content of feeds and short chain fatty acid (SCFA) was calculated using equations of McDonald et al.[25], Menke and Steingass[27] and Menke et al.[26] as:

for all feeds,
ME (MJ/kg DM) = 0.016 DOMD
for forage feeds,
ME (MJ/kg DM) = 2.20 + 0.136 GP + 0.057 CP + 0.0029 CF²
for concentrate feeds,
ME (MJ/kg DM) = 1.06 + 0.157 GP + 0.084 CP + 0.22 CF - 0.081 CA
for all feeds,
SCFA (mmol/200 mg DM) = 0.0222 GP - 0.00425

where, DOMD is \textit{in vivo} digestible organic matter in dry matter; GP is 24 h net gas production (mL/200 mg DM); CP, CF and CA are crude protein, crude fat and crude ash (%DM), respectively.

Data on apparent \textit{in vivo} digestibility and gas production parameters were subjected to one-way analysis of variance using the analysis of variation model (ANOVA) of SAS\textsuperscript{[33]}. Multiple comparison tests used Duncan’s multiple-range test\textsuperscript{[34]}.

RESULTS AND DISCUSSION

**Chemical composition:** The chemical composition of test feeds is presented in the Table 1. The CP content of feeds ranged from 11.8% in corn grain to 49.2% in soybean meal. The NDF content of feeds ranged from 10% in corn grain to 53.3% in wheat bran. Corn grain contained substantially higher OM level than the other feeds.

**Apparent \textit{in vivo} digestibility:** Values for digestibility of DM, CP, OM, NDF and ADF for each feedstuff are given in Table 2. There were differences between levels of disappearance for DM, CP, OM and NDF among feeds (p<0.05). The lower and higher extent of DM digestibility was observed in AA (67%) and CG (87%), respectively (p<0.05). For CP, the level of digestibility varied from 72% for AA to 98% for SBM. In AA, the digestibility of OM was lower than that of the other feedstuffs, whereas digestibility of OM from CG was higher than from the other feedstuffs. In addition, digestibility of NDF in CG was lower than in the other feedstuffs, whereas values for SBM were higher than the other feedstuffs (p<0.05).

**In vitro gas production:** There was a difference (p<0.05) in gas production among feeds (Table 3). Potential gas production (A) and rates of gas production (c) differed (p<0.01) among feeds. The pattern of fermentation of test feeds was distinctly different, particularly at first times of incubation (Fig. 1). Wheat bran fermented faster and corn grain fermented slower than other test feeds.

**Metabolizable energy (ME) and Short chain fatty acid (SCFA):** According to studies that reported by Menke and Steingass\textsuperscript{[27]}, Menke at al\textsuperscript{[26]} and McDonald et al\textsuperscript{[25]}, SCFA and ME could be evaluated by 24 h in vitro gas production data and in vivo organic matter digestibility. These results are shown in Table 4.
The DM and CP digestibility values for CG, SBM, WB and AA were slightly lower than the values reported by Taghizadeh et al.\[35\]. This could be expected, because the mobile nylon bag technique gives an estimate of true digestibility\[13,16\] rather than apparent digestibility, which is obtained using conventional in vivo digestibility determinations. The digestibility values of DM, CP, OM, NDF and ADF for SBM and WB were similar to reported values by Milis et al.\[28\].

High DM and OM digestibility for BG can be predicted due to high non structural carbohydrates that supply available energy as ATP for microbial growth. Low digestibility OM in AA compared to CG was due to high containing of structural carbohydrates. The digestibility value of OM for SBM was different with reported values by MAFG\[29\] and woods et al.\[38\], but the digestibility value of OM for CG was similar to reported value by Cooper et al.\[9\]. The variation in digestibility values for test feeds compared to other study results can be related to differences in chemical composition, techniques of feed processing for SBM and WB and cut stage for AA.

The strong correlation between extent of gas production and chemical composition and the poor correlation between rate of gas production and chemical composition, is consistent with Nsahlai et al.\[32\]. Low gas yield for corn grain in initial incubation times compared to the other test feeds was resulted due to high content of slowly fermented carbohydrates in corn grain. The high level of wheat bran and soybean meal gas yield in several incubation times can be assumed that degradable nitrogen was not limiting microbial activity allowing the SBM and WB carbohydrate fractions be degraded according to their potential. Giasi-Boubaker et al.\[14\] reported the positive correlation between CP and gas production at 24 h in Mediterranean browse species. Getachew et al.\[17\] reported that feed CP level was negatively correlated with gas production. However other studies with different types of feeds (i.e., CP ranging from 32 to 487 g kg\(^{-1}\) DM; Blümml et al.\[15\]) have shown no effect of CP level on gas production.

The lower extent of gas production occurred in alfalfa, also it had the lower fractional rate of gas production (c), potential gas production (A). These species were high in cell wall and lignin, which have been widely reported to decrease rate and extent of gas production\[13,16\]. This suppressing effect probably results from a reduction in attachment of ruminal microbes to feed particles\[24\].

The gas production values for corn grain, soybean meal and alfalfa were different those reported by Getachew et al.\[16\], which could be due to differences in the chemical composition of the feeds.

The high positive correlation among GP, DM and OM digestibility has been reported\[17\]. Al-Masri\[19\] reported a very highly significant (p<0.0001) relationship between gas production and the true and apparent fermented organic matter.

According to in vivo organic matter digestibility data, the ME values ranged from 9.2 in alfalfa to 13.3 MJ kg\(^{-1}\) DM in corn grain. This could be expected, because the corn grain had the highest organic matter digestibility among the other test feeds. On the contrary, alfalfa had the lower (p<0.05) organic matter digestibility because of its high containing of NDF and ADF. The negative effect of NDF on organic matter digestibility and metabolizable energy is in close agreement with al-Masri\[3\]. NRC\[11\] was reported that ME for corn grain, soybean meal, wheat bran and alfalfa were 13.04, 13.83, 10.65 and 8.19 MJ kg\(^{-1}\) DM, respectively. The difference between the NRC\[3\] data and current study for ME values of test feeds can be predicted due to variation in estimated assay, nutrient

**Table 3**: In vitro gas production characteristics of feed samples incubated in buffered rumen fluid

| Feeds      | Gas production (mL g\(^{-1}\) DM) | Gas production constants |
|------------|----------------------------------|--------------------------|
|            | 2 h | 4 h | 8 h | 12 h | 16 h | 24 h | 36 h | 48 h | 72 h | 96 h |
| CG         | 173 | 455 | 887 | 1265 | 1695 | 2235 | 2685 | 2975 | 3135 | 3205 |
| SBM        | 465 | 925 | 1575 | 1965 | 2265 | 2575 | 2875 | 3045 | 3195 | 3335 |
| WB         | 555 | 1215 | 1915 | 2235 | 2485 | 2745 | 2985 | 3165 | 3265 | 3325 |
| AA         | 365 | 745 | 1295 | 1615 | 2185 | 2185 | 2405 | 2525 | 2585 | 2605 |
| SEM (n=3)  | 2.96 | 2.83 | 4.48 | 5.23 | 5.58 | 6.71 | 7.99 | 8.21 | 8.63 | 9.16 |

**Table 4**: Evaluated metabolizable energy by in vivo digestibility and gas production results (MJ kg\(^{-1}\) DM)

| Feedstuffs | According to in vivo digestibility data | According to in vitro gas production data |
|------------|----------------------------------------|----------------------------------------|
|            | DOMD ME (MJ kg\(^{-1}\) DM) | ME (MJ kg\(^{-1}\) DM) | SCFA (m mol /200 mg DM) |
| Corn grain | 848 | 13.3 | 10.0 | 0.98 |
| Soybean meal | 771 | 12.1 | 13.4 | 1.13 |
| Wheat bran | 685 | 10.7 | 11.5 | 1.21 |
| Alfalfa | 591 | 9.2 | 7.9 | 0.96 |

**DOMD = In vivo digestible organic matter in dry matter (g kg\(^{-1}\) DM)**

The gas production results (MJ kg\(^{-1}\) DM) obtained for test feeds were different from the values reported by Cooper et al.\[29\] and Woods et al.\[30\], but the digestibility value of OM for CG was similar to reported value by Cooper et al.\[9\]. The variation in digestibility values for test feeds compared to other study results can be related to differences in chemical composition, techniques of feed processing for SBM and WB and cut stage for AA.

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composition of feeds and processing techniques. According to in vitro gas production data, the ME values ranged from 7.9 in alfalfa to 13.4 MJ kg$^{-1}$ DM in soybean meal.

The difference between two methods' results, especially for CG, arises from their kinetic of fermentation and rate of gas production in in vitro gas production technique. Fermentative gas is produced mainly when feedstuffs are fermented to acetate and butyrate, with propionate yielding gas only due to buffering of the acid. Thus feeds that produce high amounts of propionate yield lower gas volumes.$^6$ Low determination of corn grain's metabolizable energy in gas production method can be resulted from its low rate of gas production and extent of gas production at 24 h. The high non-fiber carbohydrate content of corn grain leads to proportionally higher propionate production, thereby reducing the acetate to propionate ratio.$^{17}$ Highly significant correlation has been observed between SCFA and gas production.$^6$ The molar proportions of different SCFA (acetate, propionate and butyrate) produced is dependent on the type of substrate.$^6$

Protein degradation leads to a proportionally smaller amount of SCFA. The extent of SCFA production from proteins is dependent upon on the amino acid composition of the feeds and the extent of rumen deamination of these amino acids. The carbon skeleton arising from deamination gives rise to a variety of VFA. For example, fermentation of glycine can lead to ammonia and acetic acid without the release of CO$_2$ and that of leucine, isoleucine and valine to isovaleric acid, 2-methylbutyric acid and isobutyric acids, respectively.$^{17}$

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