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## Validation of Ovine Excreta Using *in vitro* Gas to Assess Feed for Ruminants

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### ABSTRACT

Ovine excreta as inoculum for biogasification was used to assess *in vitro* and *in sacco* nutritive value of tropical pastures for ruminants. Pastures under study were: *Paspalum notatum*, *Cynodon nlemfuensis*, *Panicum maximum*, *Sporobolus indicus*, and *Desmodium* sp. Excreta were collected on grassy pastures areas. CO<sub>2</sub> systematic injections prevented strict anaerobic microorganisms contact with oxygen. A positive and significant correlation resulted from biogasification using either ovine excreta or ovine ruminal fluid. Determinant coefficient was 0,79. *Panicum maximum* and *Cynodon nlemfuensis* reached the highest nutritive value, while *Sporobolus indicus* showed the lowest one.

**Key Words:** *ovine excreta, digestibility, nutritive value*

### INTRODUCTION

Ruminants are an important source of food, as they turn products (forages and legumes) into high quality food like milk and meat, a process humans and other single-stomach organisms cannot do (Ørskov, 1999; Panin, 2000; Moore y Jung, 2001; Preston, 2003; FAO, 2008).

Food evaluation methods have been developed to serve several purposes, including animal yielding improvements (France *et al.*, 2000).

Some of the nutritional assessing methods used nowadays are *in situ* or *in sacco* ruminant degradability, and *in vitro* gas production.

The use of feces as inoculum to perform *in vitro* feed evaluation leads to similar reports by other authors, using rumen fluid, thus preventing stress and animal physical damage in the process of drawing the rumen fluid (Akhter y Hossain, 1998; Mauricio *et al.*, 2001; Van Thu, 2003).

Research done by Martínez (2005), Hernández (2006) and Resillez (2008) in the province of Camagüey clearly show the potential of bovine droppings as inoculum to carry out nutritional assessment of forages, using *in vitro* gas production. However, higher ruminants cause more expenses in terms of maintenance as lab animals. Hence, lower ruminants, especially ovine, are used. The aim of this paper is to validate ovine excreta using *in vitro* production of gas to assess ruminant feed.

### MATERIALS AND METHODS

*Experiment 1. Dynamics of in vitro gas production with ovine rumen fluid*

Several pasture samples were analyzed: Texan (*Paspalum notatum*), Star (*Cynodon nlemfuensis*), Guinea (*Panicum maximum*), Straw (*Sporobolus indicus*) and Crawling legume (*Desmodium* sp.). Some dried samples (200 mg) were put in syringes, and carefully shaken when heated in warm bath at 39 °C. A siphon was used to draw rumen fluid from the fistulate animals. The dilution was used with softened mineral medium (1 + 2) and the samples were incubated by triplicate. The general procedure used to produce gas *in vitro* stems from the principles by Menke *et al.* (1979), using 100 ml glass syringes (1 ml minimum). Readings were done every two hours for the first 24 h; then every 24 h, until 96 h.

*Experiment 2. Dynamics of in vitro gas production from ovine excreta, using dilution 1 + 3*

The samples were analyzed just like in the previous experiment, only the inoculum was changed. The ovine excreta for the inoculum were collected within the first three hours of dropped, indoors, where all animals sleep, and early in the morning, as suggested by Akhter *et al.* (1996). Then they were mixed with the softened mineral medium (1 + 3), in home blender for about a minute. The solution was filtered through inert sieve material (Dederon), in order to remove the solid particles (Martínez, 2005).

The procedure was done by systematic injection of CO<sub>2</sub> to keep oxygen from only anaerobic microorganisms.

The procedure used for *in vitro* gas production was the same as before. According to Pedraza (1998), the procedure was adjusted to modifica-

tions in LABCA conditions (Martínez, 2005). A volume of 30 ml of the inoculum was added to each syringe and the samples were incubated by triplicate. The gas volume was measured every three hours during the first 24 h and at 24, 48 and 72 h after the run.

## RESULTS AND DISCUSSION

The *in vitro* production of gas with rumen fluid is already established (Ramachandra and Krishnamoorthy, 2000; Van Thu, 2003).

Figure 1 shows the highest production of gas reaching up to 25 ml and more for legumes from 24 hours on. Straw pasture (*Sporobolus indicus*) has less nutritional value than star (*Cynodon nlemfuensis*), guinea (*Panicum maximum*), and Texan (*Paspalum notatum*). The similarity of forage behavior for both methods is apparent. The curves are monotones for the *in vitro* production of gas with rumen fluid.

Different behavior in the dynamics for the *in vitro* production of gas among the inocula from several animals corroborates observations by (Mertens *et al.*, 1997; Rymer *et al.*, 1999; Nagadi *et al.*, 2000; Cone *et al.*, 2002; Tscherning *et al.*, 2002), reporting that the offspring may be related to the characteristics of the animal diet. It has a significant influence on the rumen's microbial composition. Some variations may be observed for similar animals, even with inoculum prepared identically (Bueno *et al.*, 2005).

The substrate that goes to the large intestine is different from the one that enters the rumen, since most easily-digested nutrients have been removed and, also, because other endogenous materials have added, like mucopolysaccharides and enzymes. During microbial digestion of the large intestine volatile fatty acids are produced and then absorbed. Gases are also produced, but in general terms, digestion in the intestine is less efficient than in the rumen (McDonald *et al.*, 2002).

Figure 2 shows that dynamic behavior for the *in vitro* production of gas will depend, like every microbial growth process, on the substrate, the medium and the inoculum (Menke *et al.*, 1979; Mauricio *et al.*, 2001; Martínez, 2005). The volume of gas produced by *in vitro* incubation of a substrate is closely related to its digestibility and; therefore, with its energy value (Menke and Steingass, 1988; Getachew *et al.*, 2004)

The little difference in the production of gas during the first hours owes to the necessary adjustment phase of microorganisms to the new substrate (Menke *et al.*, 1979; Mauricio *et al.*, 2001). When adjustment is over, the gas production dynamics will basically depend on the amount of acid produced by microorganisms from the source of carbon supplied.

In that case, it is like in Figure 1, in relation to the production of gas from pastures; that is, straw (*Sporobolus indicus*), with the lowest volume of gas. All the other pastures produce more than 5 ml of gas after 48 h. The graphic with drop-pings tends to be sigmoidal.

When excreta is used the volume of gas produced after 72 h it is lower than when rumen fluid is used, which is the expected behavior (Mauricio *et al.*, 2001; Cone *et al.*, 2002; Martínez, 2005; Martínez, 2008).

An explanation for lower productions of gas using excreta given by Mould *et al.* (2005) claims that fecal micro flora has hydrolytic profiles different from those of rumen. These authors suggest that there are more qualitative (metabolic) than quantitative differences among the inocula. Cone *et al.* (2002) reported similar conclusions. All this may be influenced by the animal diet, both in composition and under processing. In this work only animals fed with poor-quality and quantity pastures. Tract microorganisms, closely associated with wastes from rumen, are also excreted with feed residues in the excreta (Theodorou *et al.*, 1994). The fecal matter basically remains anaerobic after dropped and the micro flora is alive for several hours (Holter, 1991).

The cultures inoculated with fecal matter need more time to achieve degradation potential than the ones using rumen fluid, but when they reach certain threshold, the growing speed is similar in both cases. Dhanoa *et al.* (2004), suggest the addition of more microorganisms is inoperative when the maximum degradation speed is reached.

In the excreta, microorganisms are under "suspended animation", due to the lower number of substrates: they try to survive and their metabolic activity is decreased. However, the microorganisms in the rumen, growing in a substrate-rich medium have a different behavior (Cone *et al.*, 2002).

Figure 3 shows the positive correlation among the volumes of cumulated gas when rumen's fluid

and ovine droppings are used, thus strengthening the idea that the latter might be used as inoculum (Martínez, 2008).

Aiple *et al.* (1992) described a modification in the method of Tilley and Terry (1963), by using an ovine excreta (feces) suspension as inoculum to produce *in vitro* digestion by gas production. Later on, Akhter *et al.* (1999) in comparative works between bovine feces and ovine rumen fluid concluded that the droppings may be an alternative to evaluate *in vitro* digestibility of forages.

## CONCLUSIONS

Ovine feces may be used as inoculum to produce gases for *in vitro* and *in sacco* nutritional assessments of forages for ruminant feed.

Positive and significant correlations were observed between the production of gas with ovine feces and using ovine rumen samples. The dynamics for *in vitro* production of gas using ovine rumen fluid and ovine droppings indicate that the forages assessed are arranged from higher to lower nutritional value.

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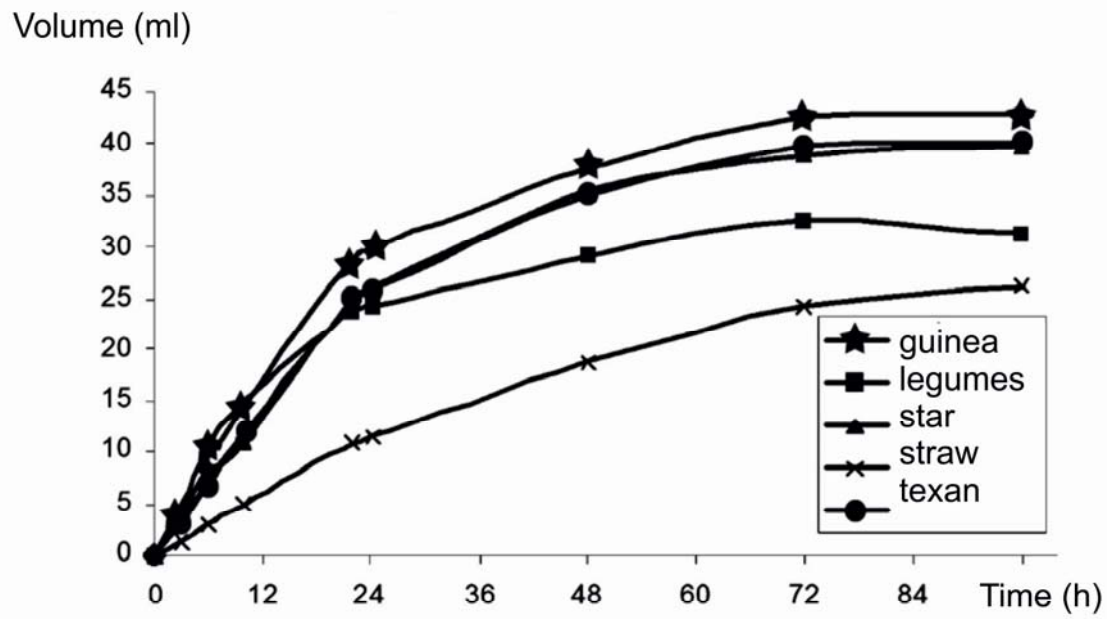
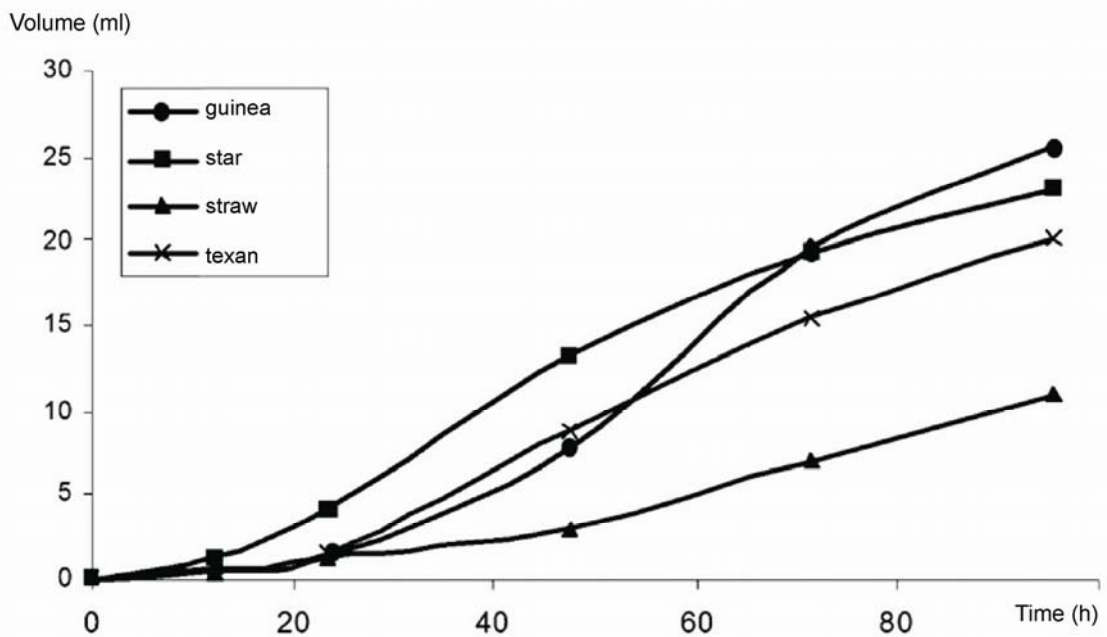
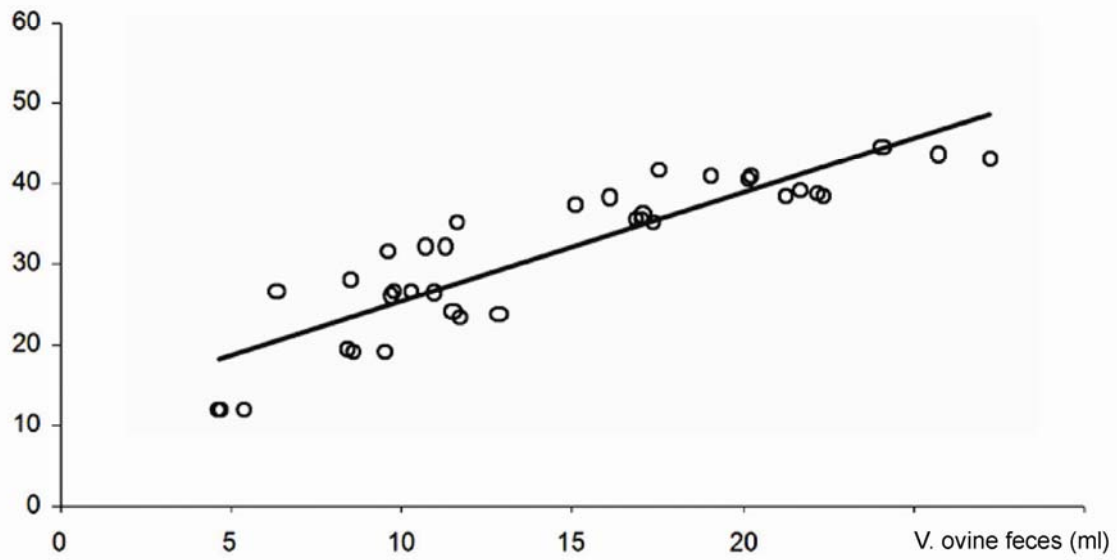


Fig. 1. Production of gas at different times using ovine rumen fluid as inoculum



**Fig. 2. Production of gas at different times using ovine feces as inoculum**

V. rumen fluid (ml)



$n = 36$ ;  $r^2 = 0,78$ ;  $P < 0,01$

**Fig. 3. Relation between the volume of gas using ovine feces and using ovine rumen fluid**