

**EFFECT OF MATURITY ON CHEMICAL COMPOSITION, PHENOLICS
AND AMINO ACIDS CONTENT AND RUMEN DEGRADABILITY OF
*Sorghum almum***

[EFECTO DE LA MADUREZ EN LA COMPOSICIÓN QUÍMICA,
CONTENIDO DE FENOLES Y AMINO ÁCIDOS Y DEGRADABILIDAD
RUMINAL DE *Sorghum almun*]

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SUMMARY

The effect of advancing maturity on chemical composition and *in sacco* rumen degradability of *Sorghum almum* was investigated. The grass was established and harvested at 6, 10 and 14 weeks-old. Herbage samples were analyzed for dry matter and chemical constituents following standard procedures. *In sacco* (Nylon bag) rumen degradability was done using 6 wethers (Age: 8 mo., 20.6 kg LW) and fed a ration comprising of 3:1 fresh *S. almum* (DM: 148.9 g DM and CP: 109.8 g kg⁻¹ DM) and *Medicago sativa* hay (DM: 903.1 g DM and CP: 188.6 g kg⁻¹ DM). Nylon bags containing 5 g DM were incubated (0, 12, 24, 36 and 48 h incubation periods). Dry matter, NDF, ADF and ADL content increased by 72.98, 62.64 and 56.5% respectively between 6 and 14 weeks. In contrast, CP, EE, total phenolics and RFV decreased by 66.67, 73.42, 25.25 and 38.65% respectively over the same period. Concentration of arginine, methionine, lysine and total amino acids declined by 66.02, 78.26, 46.44 and 63.41% respectively. DM, OM and CP degradability also declined with advancing age of the grass (P < 0.01; P < 0.0001 and P < 0.0001, respectively). It was therefore concluded that advancing maturity adversely affected the quality of *Sorghum almum*.

Key words: Sheep, Amino acids, Total phenolics, Effective degradability.

INTRODUCTION

Though it is classified as a terrestrial weed in some countries (Minnesota DNR, 1999), Columbus grass (*Sorghum almum*) is an important source of nourishment for ruminant livestock in many production systems. Due to its high forage characteristics (Kallah et al., 1999; Muhammad, 1993;

RESUMEN

Se estudio el efecto de la madurez sobre la composición química y la degradabilidad ruminal de *Sorghum almum*. El pasto fue cosechado a la 6, 10 y 14 semanas de crecimiento. La degradabilidad ruminal fue medida en 6 borregos (8 meses edad, PV 20.6 kg) con canula ruminal y alimentados con *S. almum* fresco (148.9 g MS y PC: 109.8 g kg⁻¹ MS) y heno de *Medicago sativa* (903.1 g MS y PC: 188.6 g kg⁻¹ MS) en proporción 3:1. Se incubaron bolsas de nylon con 5 g de MS (0, 12, 24, 36 y 48 h). El contenido de MS, FDN, FDA y ADL incrementó en 72.98, 62.64 y 56.5% respectivamente entre la 6^a. y 14^a. semana. En contraste, PC, EE, fenoles y RFV disminuyeron en 66.67, 73.42, 25.25 y 38.65% respectivamente. Las concentraciones de arginina, metionina, lisina y total de amino ácidos declinaron en 66.02, 78.26, 46.44 y 63.41% respectivamente. La degradabilidad de la MS, MO y PC se redujo con la edad (P < 0.01; P < 0.0001 y P < 0.0001, respectivamente). Se concluye que la madurez afecta de manera adversa la calidad del *Sorghum almum*.

Palabras clave: Ovinos, Amino ácidos, polifenoles totales, degradabilidad efectiva.

Muhammad et al., 1994), drought (Gohl, 1981) and disease resistance (Rich et al., 2004; Reed et al., 2002), *Sorghum almum* is particularly an attractive forage crop for ruminant livestock production in many tropical and subtropical regions. In these regions, *Sorghum almum* is widely recognized and valued for its high yield of palatable herbage (Reeds, 1976; Bogdan, 1977; Myoya et al., 1987) suitable for

ensiling (Kallah *et al.*, 1997; Kallah *et al.*, 1999; Aminah *et al.*, 2000; Chin, 2002), haymaking (Duke, 1983) and grazing (Catchpoole, 1972). In Argentina where it was first developed, *Sorghum almum* is used almost exclusively for grazing (Duke, 1983). Its chemical composition (Kallah *et al.*, 1999; Chin, 2002) and digestibility (Rodriquez, 2005) is closely comparable to those of commonly cultivated high quality tropical grass fodder crops (Muia, 2000; Keftasa, 1990). Excellent performance of ruminant livestock on *Sorghum almum* based diets has also been recorded (Sheridan *et al.*, 2003; Aminah *et al.*, 2000), which in turn suggests that, with a sound feeding management, the reported presence of prussic acid (Gohl, 1981) poses no major risk to ruminants. Its economic contribution in smallholder peri-urban dairy production systems has also been documented (Agyemang *et al.*, 1998). Despite its long history in the country (Orodho, 2006), cultivation of *Sorghum almum* on smallholder farms in Kenya has remained low. The scanty literature on its chemical composition, total phenolic and amino acid profile and rumen degradability characteristics at different morphological stages and under different edaphic factors under Kenyan farming condition is presently being seen as the major drawback, to its adoption by smallholder farmers. This paper reports the results of the study conducted to determine the effect of advancing maturity on these quality aspects and relative feed value. The overall objective was to generate technical information that would guide *Sorghum almum* utilization in Kenya and other tropical and sub-tropical regions.

MATERIAL AND METHODS

Forage establishment and sampling

The study was conducted at the forage experimental unit of the National Animal Husbandry Research Centre in Naivasha Kenya (soil: 216 g Ca kg⁻¹ DM, 113 mg P kg⁻¹ DM and pH = 7.38). The centre is located on grid 0° 40' S and 36° 26' E at an altitude of 1940 metres above sea level (Jaezhold and Schmidt, 1982) with a mean temperature of 18° C. To study the effect of advancing maturity on concentration of chemical constituents and rumen degradability of Columbus grass (*Sorghum almum*), 36 plots (size: 2 x 2 sq. m separated by 30 cm weed free guard rows) were demarcated and further divided into 3 similar units (N = 12). In each plot, 5 rows of 2 m length, 30 cm apart and about 4 cm deep were drilled. A total of 25 g of seed (viability approx: 80%) were then sown with each line receiving 5 g of seed (evenly sprinkled). The plots were later harvested at the age of 6, 10 and 14 weeks each time at 5 cm above soil. Representative grabs of the freshly harvested forage were made per plot, chopped to pieces of 2 cm length, mixed thoroughly and 2 composite samples (500 g each)

were taken for dry matter (DM) determination, chemical analyses, amino acids profiling, total phenolic assaying and rumen degradability study.

Animals and diets

Six rumen fistulated wethers aged about 8 months with a mean live weight (LW) of 20.6 kg were used. Prior to the study, all the wethers were drenched with a broad-spectrum anthelmintic, Valbazen (Smith-kline, Switzerland) and sprayed with an acaricide to eradicate both internal and external parasites respectively, they were then weighed, ear-tagged and then placed into well-ventilated individual metabolic cages (size: 120 cm long x 50 cm wide x 90 cm high) raised 60 cm above a concrete floor and equipped with feed and water troughs. A ration comprising of 3:1 fresh 8 – 10 weeks old Columbus grass (*Sorghum almum*: 148.97 g DM; 109.87 g CP kg⁻¹ DM, 652.97 g NDF kg⁻¹ DM, 346.6 g ADF kg⁻¹ DM and Lucerne hay (*Medicago sativa*: 903.1 g DM; 188.6 g CP kg⁻¹ DM, 426.8 g NDF kg⁻¹ DM, 318.1 g ADF kg⁻¹ DM), both chopped to a particle size of 2 cm, was offered *ad libitum* twice daily in equal amounts at 07.30 h and 16.30 h beginning 14 days before the start of the trial. They also had a free access to a balance mineral lick and clean drinking water.

In sacco degradability

Dry matter (DM), Organic matter (OM) and crude protein (CP) degradation characteristics were determined using the nylon bag technique as described by Ørskov and McDonald (1979) and Ørskov *et al.* (1980). The feed materials were dried at 65 °C for 24 hours and thereafter grounded in a Christy and Norris mill to pass a 2.5 mm sieve (Abdulrazak and Fujihara, 1999). Representative samples of 5 g each were weighed (in duplicates) into clean well-labeled and weighed nylon bags of pore size 40 µm and inner size of 6.5 cm x 12 cm. The bags were inserted into the rumen (12 bags per wether) all at the same time and withdrawn in a sequentially at: 12, 24, 36 and 48 h. Immediately after withdrawal, the bags were dipped into cold water to stop further microbial activity and then rinsed by cold tap water to remove the rumen matter from the outside of the bags. Thereafter, the bags were washed gently under slow running tap water for 30 minutes, till the water was clear. Finally, they were dried at 65°C for 48 hours in a forced air oven, desiccated for 30 minutes and then weighed. To determine the washing loss 2 additional bags (0 h) containing 5 g each for each of the test feeds were soaked in water bath kept at 39°C for 1 h and thereafter underwent the same washing and drying procedures as the incubated bags. The *in sacco* dry DM, OM and CP digestibility at different incubation periods were determined as described by Ørskov *et al.* (1980).

Laboratory analyses

The nylon bags (containing residue) were dried in a forced-air oven at 65°C for, 48 h after which residue weights were calculated. Both forage samples and residues were grounded to pass 1 mm screen and stored in airtight plastic bottles while awaiting chemical analyses. Ash was determined (AOAC, 1990, ID 942.05) and micro-Kjeldahl N in feed and residues (AOAC, 1990, ID 954.01). Crude protein was calculated as Kjeldahl N x 6.25. Neutral-detergent fibre (NDF), acid-detergent fibre (ADF) and acid-detergent lignin (ADL) were determined by the procedures of Goering and Van Soest (1970) and Van Soest and Robertson (1985). Ether extract (EE) was determined by extracting the sample with petroleum ether using a Gerhart Soxtherm 2000 Automated (AOAC, 1990, ID 920.39). Representative samples taken from the various harvests (dried at 40°C and grounded to pass 1 mm screen) were also assayed for total phenolics (TP) by the Folin assay method (Folin – Denis method) as described by Waterman and Mole (1994) and Palm and Rowland (1997). Linear regression was used to construct a standard curve relating absorbance (760 nm) of the Folin assay reaction mixture to the known concentration of tannic acid (μg) (standard). Similarly, representative dry samples (approx. 5 μg) from forage harvested at different maturity stages were also assayed for both essential and non-essential amino acids (AA) concentration according to AOAC (1990, ID 994.12 Llamas and Fontaine 1994).

Statistical analysis and calculations

The chemical constituents in the 3 grass harvests were analysed as a randomised complete block design using SAS (2002). The statistical model used was: $Y_i = \mu + A_i + e_i$; where Y_i = the concentration of the chemical constituents measured; μ = overall mean, A_i = age at harvest and e_i = random experimental error. The Proc GLM was used to perform the analysis of variance (ANOVA) and means were contrasted with significance assumed when $P = 0.05$ (SAS, 2002). Linear and quadratic regression functions were fitted using SPSS (2003) to describe the exhibited change patterns. Data for *in sacco* degradation were fitted to the first order kinetics defined by the exponential equation of the form $p = a + b(1 - e^{-ct})$; (Ørskov and McDonald 1979; McDonald, 1981): where p represents degradation of DM, OM and CP at time t ; 'a' – readily soluble fraction; 'b' – slowly degradable fraction; 'c' is the rate of degradation of 'b' per h and 'a + b' is potential degradation. Effective rumen degradation of the forage components was calculated as: $ED = a + (b \times c) / (c + k)$, assuming the rumen fractional outflow rates (k) of 2, 5 and 8% h^{-1} (Ørskov and Shand, 1997; Ikhimioya et al., 2005). Degradation

constants a , b and c were derived using neway excel (version 5.0) as described by Chen (1997) and the accruing data was analyzed using SPSS (2003) and SAS (2002).

RESULTS AND DISCUSSION

Dry matter (DM) content increased by 72.98% ($r^2 = 0.9341$; $p < 0.0001$; SEM = 5.87) from 125.5 g DM at 6 to 217.1 g DM at 14 weeks, representing a gain of 1.6 g DM d^{-1} (table 1). Neutral detergent fibre (NDF), acid detergent fibre (ADF) and acid detergent lignin (ADL) increased by 33.05 ($p < 0.0001$), 62.64 ($P < 0.0001$) and 56.5% ($p < 0.0001$) respectively. Concentration of cellulose also increased ($p < 0.0001$). In contrast, crude protein (CP) and ether extract (EE) content and relative feed value (RFV) decreased by 66.67 ($r^2 = 0.9921$; $p < 0.0001$), 73.42 ($r^2 = 0.9419$; $p < 0.0001$) and 38.65% ($r^2 = 0.9927$; $p < 0.0001$) respectively. Concentration of hemicellulose, nitrogen free extract (NFE), non-structural carbohydrates (NSC) and total carbohydrates (CHO) exhibited inconsistent change pattern with advancing maturity of the grass. A high correlation (negative) was founded between age at harvest and CP ($r = -0.994$; $p < 0.0001$), EE ($r = -0.920$; $p < 0.0001$) and RFV ($r = -0.930$; $p < 0.0001$)(table 1). The pattern of concentration change for hemicellulose, NFE and NSC were not correlated with advancing maturity of the grass. DM, OM, NDF, ADF, ADL and CHO recorded strong (positive) and significant correlation with age of the grass. From the results, it is also clear that quadratic function achieved a better fit for the prediction of the chemical nutrients' concentration change pattern over the study period as evidenced by the comparatively higher p - values and coefficients of determination (table 1). The chemical (table 1) values recorded are generally comparable to those reported for *Sorghum alnum* (Kallah et al., 1999; Chin, 2002; Gohl, 1981) and other tropical grasses (Keftasa, 1990; Muia, 2000; Wouters, 1985; Snijders et al., 1992). At 10 weeks however, concentration of NDF surpassed the range of 600 – 650 g kg^{-1} DM suggested as the critical limit above which efficiency of utilization of tropical forages by ruminants would be impaired (Van Soest, 1982; Muia, 2000). At the same maturity stage, concentration of CP was at the threshold of the range of 70 – 80 g kg^{-1} DM below which intake of tropical feed resources would be severely depressed (Larbi et al., 1992; Minson, 1990; Leng, 1990). The observed increase in concentration of fibre components was consistent with Arthington and Brown (2005). The decline in protein concentration with advancing maturity of the grass was attributed to the decrease in CP concentration on leaf swards, stem and increased proportion of stem in the mature grass (Buxton, 1996). The recorded CP decline rate of 1.86 g kg^{-1} DM d^{-1} was slightly above the mean of 1 g kg^{-1} DM d^{-1} in data

reported by Minson (1990) for several tropical forages and falls within the 1.0 – 1.7 g kg⁻¹ DM d⁻¹ reported by Keftasa (1990) in Rhodes grass (*Chloris gayana*). Another important feature of the chemical composition of tropical grasses, which may in part account for their lower digestibility (Keftasa, 1990), is the relatively high content of phenolic compounds covalently bound in the cell wall (Lowry *et al.*, 1993; Paul *et al.*, 2003). In this study, presence of total phenolic compounds was recorded. The mean concentration however decreased by 25.25% between 6 and 14 weeks from 38.49 to 28.77 g kg⁻¹ DM, which agreed with Martin (2006). These mean values fell within the range of 10 – 50 g kg⁻¹ DM reported by Lowry *et al.* (1993) in 19 species of native and introduced grasses in North Queensland.

Dietary supply of rumen by-pass protein and microbial synthesis of free amino acids are recognized as the 2 main sources of amino acids available for absorption from the small intestine of ruminants under normal feeding conditions (Von Keyserlingk *et al.*, 1998; Taghizadeh *et al.*, 2005); hence the need to continue developing sufficient knowledge on amino acid profiles of important ruminant feedstuffs particularly forages. The amino acids concentration values obtained in this study (table 2) were consistent with those reported in literature (Tedeschi *et al.*, 2001; Von

Keyserlingk *et al.*, 1998; Taghizadeh *et al.*, 2005). Concentration of all amino acids assayed decreased with advancing maturity of the grass.

The study revealed that concentration of Arginine, Methionine, Lysine and total amino acids (TAA) in the grass herbage decreased by 66.02, 78.26, 46.44 and 63.41% respectively between 6 and 14 weeks (table 2). Decline in concentration of amino acids with advancing plant maturity has been recorded (Yao *et al.*, 2000; Arthington and Brown, 2005). Leucine, Alanine, Aspartic acid and Glutamic acid showed the highest concentration, whereas Methionine, Serine and Cysteine recorded lowest concentration in *Sorghum almum* herbage. Rate of decrease in concentration of individual amino acids with age of the grass, differed markedly. Results further showed that rumen degradability of *Sorghum almum* herbage is strongly influenced by growth stage at harvest (tables 3, 4 and 5). This was evidenced by the observed marked differences in rumen degradability (table 3) and fermentation parametric (table 4) values between the various harvests. At 48 h incubation period, the DM and CP degradability of grass herbage harvested at the age of 6 weeks were 71.17 and 71.99%, whereas those of 14 weeks old herbage were 59.49 and 53.68% respectively.

Table 1. Chemical composition and relative feed value of *Sorghum almum* as influence by advancing maturity (g kg⁻¹ DM except where stated).

Component	Age at harvest (Weeks)			SEM	ANOVA		Contrasts		Correlation	
	6	10	14		r ²	P ^a	L	Q	r	P
Air DM (g kg ⁻¹)	125.5 ^a	153.9 ^b	217.1 ^c	5.87	0.9341	****	****	****	0.944	****
Analytical DM (g kg ⁻¹)	904.9 ^a	930.2 ^b	926.9 ^b	4.36	0.6885	**	*	**	0.663	*
Organic matter	753.1 ^a	781.7 ^b	803.5 ^c	5.48	0.8252	***	****	****	0.903	****
Crude protein	156.3 ^c	97.8 ^b	52.1 ^a	2.19	0.9921	****	****	****	-0.994	****
Crude fibre	195.8 ^a	346.3 ^b	356.6 ^b	6.72	0.9755	****	****	****	0.882	****
Neutral detergent Fibre	519.9 ^a	664.3 ^b	691.7 ^c	3.96	0.9918	****	****	****	0.957	****
Acid det. Fibre	259.6 ^a	377.6 ^b	422.2 ^c	5.89	0.9784	****	****	****	0.927	****
Acid det. Lignin	35.4 ^a	36.8 ^a	55.4 ^b	2.74	0.7881	***	**	**	0.794	**
Ether extract	39.5 ^c	16.5 ^b	10.5 ^a	1.79	0.9419	****	****	****	-0.920	****
Hemicellulose	260.4 ^a	286.5 ^b	269.5 ^a	4.91	0.6191	*	NS	*	0.271	NS
Cellulose	224.2 ^a	340.9 ^b	366.8 ^c	5.43	0.9775	****	****	****	0.928	****
N free extract ¹	361.5 ^b	321.1 ^a	384.3 ^b	7.11	0.8187	***	NS	****	0.322	NS
NSC ²	132.5 ^b	72.9 ^a	122.4 ^b	3.79	0.9398	****	NS	****	-0.153	NS
CHO ³	557.4 ^c	667.3 ^b	740.9 ^a	7.31	0.9731	****	****	****	0.980	****
RFV ⁴	122.9 ^c	83.3 ^b	75.4 ^a	1.03	0.9927	****	****	****	-0.930	****

¹NFE (g kg⁻¹ DM) = DM – (EE + CP + Ash + CF) (Van Soest, 1982); ²NSC was calculated as 100% - (CP% + NDF% + EE% + Ash%); ³CHO was determined as as described by Arieli *et al.* (1999), where CHO (g kg⁻¹ DM) = OM content – (CP + EE); ⁴Relative feed value = [(88.9 – (0.78 x ADF%)) x (120/NDF%)]/ 1.29 (Agric-facts, 2006); **** p < 0.0001; *** p < 0.001; ** p < 0.01; * p < 0.5 and NS – Not significant (p > 0.05); L – linear function; Q – quadratic function; P^a – ANOVA p-value; N =12 in all cases.

Table 2. Amino acid composition (mg g⁻¹ DM) of *Sorghum alnum* harvested at different maturity stages.

Amino Acid	Grass age (Weeks)			Regression equations				Correlation Coefficients ²	
	6	10	14	r ²	P	r ²	P	r	P
<i>Essential (mg/g DM)</i>									
Arginine	5.15	2.65	1.75	0.901	**	0.968	**	- 0.949	**
Histidine	1.90	1.05	0.75	0.886	**	0.953	*	- 0.941	**
Isoleucine	4.11	2.05	1.41	0.890	**	0.970	**	- 0.944	**
Leucine	8.81	4.21	2.85	0.879	**	0.967	**	- 0.938	**
Lysine	4.50	3.05	2.41	0.929	**	0.974	**	- 0.964	**
Methionine	1.15	0.55	0.25	0.740	*	0.767	NS	- 0.860	*
Phenylalanine	5.91	3.05	1.65	0.898	**	0.933	*	- 0.948	**
Threonine	3.05	1.60	1.15	0.882	**	0.963	**	- 0.939	**
Valine	6.61	3.50	2.41	0.907	**	0.975	**	- 0.952	**
<i>None essential (mg/g DM)</i>									
Alanine	9.65	3.80	2.35	0.931	**	0.973	**	- 0.965	**
Aspartic acid ¹	8.35	5.80	3.65	0.950	***	0.952	*	- 0.975	***
Glutamic acid ¹	10.80	6.12	4.11	0.912	**	0.961	**	- 0.955	**
Glycine	5.30	2.55	1.82	0.871	**	0.968	**	- 0.934	**
Serine	1.15	0.65	0.51	0.792	*	0.869	*	- 0.890	*
Tyrosine	3.10	1.65	1.01	0.902	**	0.946	*	- 0.951	**
Proline	3.85	1.95	1.45	0.890	**	0.991	***	- 0.944	**
Cysteine	0.65	0.50	0.41	0.915	**	0.927	*	- 0.956	**
Total AA	81.31	44.55	29.75	0.912	**	0.967	**	- 0.955	**

1 – The Aspartic acid peak contained Asparagines, and the Glutamic acid peak contained Glutamine;

2 – Correlation between individual amino acid and advancing plant maturity.

*** p < 0.001; ** p < 0.01; * p < 0.5 and NS – Not significant (p > 0.05);

Table 3. Influence of maturity on dry matter, organic matter and crude protein disappearance of *Sorghum alnum*.

Incubation time (h)	N	Grass age (Weeks)			SEM	P	ANOVA	
		6	10	14			r ²	CV
DM disappearance (%)								
0	12	28.18 ^c	25.31 ^b	21.69 ^a	0.865	***	0.6529	8.45
12	18	49.75 ^c	40.21 ^b	36.67 ^a	0.881	****	0.6985	8.86
24	18	59.15 ^c	51.51 ^b	46.21 ^a	0.935	****	0.6547	7.59
36	18	66.97 ^c	57.86 ^b	53.26 ^a	0.991	****	0.6604	7.08
48	18	71.17 ^c	65.22 ^b	59.49 ^a	0.609	****	0.7832	3.95
OM disappearance (%)								
0	12	12.95 ^b	11.07 ^{ab}	9.27 ^a	0.951	*	0.3327	20.98
12	18	39.18 ^b	29.69 ^a	27.55 ^a	0.955	****	0.6225	12.61
24	18	51.12 ^c	44.04 ^b	39.69 ^a	1.037	****	0.5484	9.78
36	18	60.19 ^c	51.97 ^b	47.99 ^a	0.968	****	0.6183	7.69
48	18	64.39 ^c	59.22 ^b	54.07 ^a	0.749	****	0.6506	5.36
CP disappearance (%)								
0	12	19.91 ^b	21.88 ^b	11.23 ^a	0.877	****	0.8476	12.16
12	18	47.23 ^c	36.45 ^b	26.88 ^a	0.844	****	0.8508	9.72
24	18	57.63 ^c	50.41 ^b	42.13 ^a	0.815	****	0.7801	6.91
36	18	67.34 ^c	57.44 ^b	51.59 ^a	0.828	****	0.7837	5.98
48	18	71.99 ^c	65.26 ^b	53.68 ^a	0.577	****	0.9101	3.84

N – Number of Nylon bags; SEM – Standard error mean; CV – Coefficient of variation;

Means with the same superscript (^{a, b, c}) within the same row are not significantly different (p > 0.05);

**** p < 0.0001; *** p < 0.001; * p < 0.5

Organic matter degradability followed the same pattern. At the same incubation period, the quickly ('a'), slowly ('b'), rate of degradation ('c') and potential ('a + b') DM fermentation fractions of herbage harvested at 6 weeks were 13.63, 6.3, 48.08 and 8.74% respectively higher than those recorded for herbage harvested at 14 weeks (table 4). Similarly, the 'a', 'b', 'c' and 'a + b' CP fermentation fractions were 51.45, 9.95, 45.27 and 20.79% respectively higher for grass herbage harvested at 6 weeks than those recorded for herbage harvested at 14 weeks. The

effective degradability ($5\%h^{-1}$) of DM, OM and CP for herbage harvested at 6 weeks was 22.86, 21.30 and 32.51% respectively higher than those recorded for grass herbage harvested at 14 weeks (table 5). Effective degradability of DM, OM and CP decreased progressively with increasing rate of passage (table 5). ED of DM for grass herbage harvested at 6, 10 and 14 weeks decreased by 25.38, 32.41 and 34.4% respectively with an increase of rumen outflow rate from 2 to $8\%h^{-1}$.

Table 4. Fermentation characteristics of *Sorghum alnum* as influence by age at harvest.

Fraction	N	Grass age (Weeks)			SEM	P	ANOVA	
		6	10	14			R ²	CV
Dry matter								
a	12	23.41 ^c	21.68 ^b	20.22 ^a	0.394	***	0.6862	4.43
b	18	46.99 ^b	47.46 ^b	44.02 ^b	0.937	*	0.3459	4.97
c	18	0.052 ^b	0.029 ^a	0.027 ^a	0.005	**	0.5364	32.45
a + b	18	70.39 ^b	69.14 ^b	64.24 ^a	1.041	**	0.5659	3.75
Organic matter								
a	12	13.13 ^b	10.99 ^{ab}	9.33 ^a	0.976	*	0.3363	21.43
b	18	57.68 ^b	54.09 ^b	44.54 ^a	1.262	****	0.7944	5.94
c	18	0.049 ^b	0.034 ^a	0.031 ^a	0.005	*	0.3319	32.69
a + b	18	70.82 ^c	65.09 ^b	53.87 ^a	1.431	****	0.8288	5.54
Crude protein								
a	12	21.69 ^b	19.19 ^b	10.53 ^a	0.709	****	0.8607	10.25
b	18	56.77 ^b	54.56 ^b	51.12 ^a	1.155	*	0.4473	5.22
c	18	0.055 ^b	0.033 ^a	0.029 ^a	0.004	***	0.6867	22.04
a + b	18	77.82 ^b	73.76 ^b	61.64 ^a	0.986	****	0.9015	3.34

N – Number of Nylon bags; SEM – Standard error mean; CV – Coefficient of variation;

Means with the same superscript (^{a,b,c}) within the same row are not significantly different ($p > 0.05$);

**** $p < 0.0001$; *** $p < 0.001$; ** $p < 0.01$; * $p < 0.5$

Table 5. Mean effective degradability values of nutrients in *Sorghum alnum* as influenced by age at harvest.

Outflow rate	N	Grass age (Weeks)			SEM	ANOVA		Regression	
		6	10	14		r ²	P	L	Q
Dry matter									
$k = 0.02$	18	61.47 ^c	56.77 ^b	51.63 ^a	0.486	0.9715	****	****	****
$k = 0.05$	18	51.23 ^c	43.97 ^b	39.52 ^a	0.466	0.9817	****	****	****
$k = 0.08$	18	45.87 ^c	38.37 ^b	33.87 ^a	0.507	0.9795	****	****	****
Organic matter									
$k = 0.02$	18	53.31 ^c	48.81 ^b	44.37 ^a	0.573	0.9529	****	****	****
$k = 0.05$	18	41.13 ^c	35.13 ^b	32.37 ^a	0.481	0.9674	****	****	****
$k = 0.08$	18	34.63 ^c	28.57 ^b	25.64 ^a	0.501	0.9656	****	****	****
Crude protein									
$k = 0.02$	18	60.32 ^c	55.77 ^b	45.52 ^a	0.526	0.9857	****	****	****
$k = 0.05$	18	48.61 ^c	42.37 ^b	32.81 ^a	0.462	0.9899	****	****	****
$k = 0.08$	18	41.54 ^c	36.36 ^b	26.73 ^a	0.402	0.9915	****	****	****

N – Number of Nylon bags; SEM – Standard error mean; CV – Coefficient of variation;

Means with the same superscript (^{a,b,c}) within the same row are not significantly different $p > 0.05$);

k – Rumen outflow rate; L and Q – Linear and quadratic equations;

**** $p < 0.0001$

The obtained *in sacco* degradability and fermentation values in this study were consistent with those reported in literature for tropical grass (Keftasa, 1990; Muia, 2000; Rodriguez et al., 2005). A decline in rumen degradability of nutrients with advancing grass maturity has been recorded (Arthington and Brown, 2005; Mislevy and Martin, 1998; Bogoro et al., 2006; Rodriguez et al., 2005). Decrease of rumen fermentation parameters with advancing grass maturity and increased outflow rates has also been reported (Rodriguez et al., 2005; Ikhimiya et al., 2005). The observed variation in degradability between the different harvests could be due to the decreasing concentration of the more degradable cell contents (Van Soest, 1994) and the accumulation of lingo-cellulose fractions (Smith et al., 1989; Akin, 1989) compounded by the decreased leaf to stem ratio (Hides et al., 1983) as the grass matured. Lignified cells reduce the penetration of rumen fungal rhizoids (Latham et al., 1987), which results in reduced degradability. Though presence of total phenolics has been associated with decreased rumen degradability (Waghorn et al., 1994), no relationship between degradability and phenolic content was observed in this study.

CONCLUSION

Advancing maturity adversely affects the concentration of essential chemical nutrients, degradability and relative feed value of *Sorghum almum*. The results of this study favored between 6 and 10 weeks as the appropriate stage to feed *Sorghum almum* herbage. Caution should however be observed when feeding young because this study has shown that in immature herbage, concentration of phenolic compounds is high.

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