Using high quality forages to improve *in vitro* rumen degradability and fermentation of low quality forages

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Introduction Low quality forages (LQ), like cereal straws, are the main basal feeds for ruminants in developing countries. However, animal production is low when ruminants are reared on nutrient deficient LQ as the main diet. To get more production from ruminants it is necessary to enhance the utilization of these LQ which could be achieved by using small amounts of high quality forages (HQ) as supplements with LQ. This study compared the effect of incubating *in vitro* different amounts of various HQ with LQ on the DM degradability (IVD) and fermentation (e.g. ammonia) profiles of rice straw (Straw) and grass hay (Hay) as LQ.

Materials and methods A 3x4x2 factorial arrangement in duplicate was used to assess the effect of 3 supplements (rye grass=RG, silage=Si and rape seed plant=RP) each at four levels of 0, 250, 500 and 750 mg/g DM on IVD and ammonia (AL) of 2 LQ (Straw and Hay) during 96h of *in vitro* incubations. Rumen fluid (RF) was obtained from 2 fistulated sheep just before feeding, strained through a cheese cloth, and then it was mixed with a pre-warmed buffer at 1:4 ratio to prepare inoculum. The incubations of forages were conducted in 50-ml polypropylene tubes containing 0.4g of ground LQ to which relevant HQ were added according to the experimental design but by enclosing these HQ individually in small polyester bags (45μ). These bags permitted the mixing of solubles but not the HQ with LQ and so helped observe the IVD of LQ without their direct mixing with HQ. About 40 ml of the inoculum was added under CO_2 to each tube which was sealed with a rubber stopper containing pressure release valve and incubated at $39^{\circ}C$ for the pre-determined times. After incubation the tubes were submerged in ice to stop fermentation. The liquids and residues were separated by centrifuging the tubes at 3000 rpm for 10 minutes. The residues were washed with distilled water and dried at $60^{\circ}C$ for 48h to determine IVD. The supernatants were acidified with 1N HCl and analysed for AL by using a colorimeter at 660 nm. The IVD and AL data were analyzed by using the General Linear Model of Minitab to study the main effects of LQ, HQ and HQ level=S and their interactions at P<0.05 for the IVD and AL.

Results Main effects of LQ, HQ and S were significant (P<0.003) for IVD and AL. However, due to some significant (P<0.05) interactions between these variables, the mean IVD and AL for each treatment combination are shown in Table 1. IVD was higher for Hay than Straw with all HQ. Mean IVD of LQ was higher with RG than other HQ where IVD increased with RG and Si but reduced with RP for Straw but not for Hay, hence LQ x HQ interaction (P<0.002). While IVD of both LQ did not increase with the increase in HQ after 250 mg , AL continued to increase with increase in HQ from 0 to 500mg. In fact AL was significantly (P<0.001) higher for Hay than Straw and it was highest with 500 mg RP than other HQ.

Table 1 Effects of different level of HO on the mean IVD and AL for LO after 96h of incubation

Table 1 Effects of different level of free of the off the fineal TVD and AE for EQ after 30h of fineabation									
LQ	HQ	IVD (g/kg)				AL (mg/L)			
		Amount of HQ (S) (mg/g) LQ				Amount of HQ (S) (mg/g) LQ			
		0	250	500	750	0	250	500	750
Straw	RG	607	666	656	652	92	165	187	188
	RP	604	583	567	563	96	175	209	205
	Si	608	633	625	562	93	135	155	168
Hay	RG	740	765	756	675	124	167	210	212
	RP	732	738	721	740	120	173	225	220
	Si	725	808	753	761	128	163	181	182
Pooled SEM		11.1				5.97			
P <for hq,="" lq,="" s<="" td=""><td colspan="4">LQ=0.001;HQ=0.002; S=0.003;</td><td colspan="4">LQ=0.001;HQ=0.001;S=0.001;LQxHQ=0.3;</td></for>		LQ=0.001;HQ=0.002; S=0.003;				LQ=0.001;HQ=0.001;S=0.001;LQxHQ=0.3;			
		LQxHQ=0.002; LQxS=0.8; HQxS=0.09;				LQxS=0.07; HQxS=0.001; LQxHQxS=0.5			
LQxHQxS=0.02									

Conclusions The effect of HQ on IVD and AL varied with the type and level of HQ and the LQ during *in vitro* incubation for 96h. While RP showed negative effect on IVD of LQ, other HQ showed maximum IVD at 250 mg. Possible antinutritional factors in RP might have affected the microbial activity which resulted in reduced IVD of LQ. Conversely, more crude protein might have increased AL when HQ were used at higher levels RG and Si can be used as supplements to increase the utilization of LQ but their amounts needs to be optimised in association with the target LQ

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