

The influence of molasses and yeast culture on the performance of growing bulls on grass silage-based diet

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ABSTRACT

The effects of dietary inclusion of yeast culture (YC) and of gradually increasing proportion of sugar beet molasses [0 (M_0), 100 (M_{100}) and 200 (M_{200}) g kg⁻¹] in the concentrate mixture on feed intake, live weight gain and carcass characteristics of growing Ayrshire bulls were examined in a 3 x 2 factorial design. Twenty-eight animals with a mean initial liveweight (LW) of 268 kg were used. Molasses replaced a part of concentrate, which was based on a mixture (1:1) of barley and sugar beet pulp. Grass silage was offered *ad libitum* and the concentrate supplements at the rate of 100 g per LW^{0.6} on air dry basis. YC (5 g day⁻¹) was top-dressed on the concentrate. Feed intake was not affected by the treatments but LW gain decreased linearly ($P < 0.05$) with increasing rate of molasses (1.21, 1.16 and 1.09 kg day⁻¹; SEM 0.029). As the proportion of molasses increased from 0 to 200 g/kg concentrate the efficiency of feed conversion declined (linear effect $P < 0.01$) from 6.10 to 6.75 kg DM kg⁻¹ LW gain. YC had no significant effect either on feed intake (7.26 vs 7.49 kg day⁻¹) or LW gain (1.14 vs 1.17 kg day⁻¹). Yeast interacted with concentrate so that with M_0 LW gain was increased and with M_{200} decreased by YC. The differences in carcass weight and carcass gain reflected a pattern similar to those in LW gain. Dressing proportion and fat grades were not affected by the treatments but carcass quality grade was better in animals receiving YC.

KEY WORDS: growing cattle, yeast culture, molasses, silage

INTRODUCTION

Yeast cultures (*Saccharomyces cerevisiae*) have been used as a dietary supplement for ruminants for many years. It has been thought to improve rumen

function, and therefore, production and feed efficiency. However, the responses in liveweight gain and milk production have been variable (Wallace, 1994). It appears that the effects of the products are dependent on the composition of the basal diet. The greatest responses have been observed with high-concentrate diets (Williams and Newbold, 1990; Carro et al., 1992a). This may be related to the stabilizing effect of yeast culture on rumen fermentation and to a reduction in postprandial concentration of lactic acid (Williams, 1989).

In Finland beef production is based on the utilization of large quantities of grass silage in the diet. Feeding grass silage-based diets is characterized by a high postprandial concentration of ammonia N in the rumen. Extensive proteolysis and fermentation of water soluble carbohydrates of the grass in the silo leads to unbalanced supply of energy and nitrogen for rumen microbes. Sugar supplements have been used to better match energy and N supply from silage for rumen microbes (Syrjälä, 1972; Chamberlain et al., 1985). Although sugar supplements were in these studies more efficient than starch supplements in reducing the concentration on ammonia N in the rumen, production responses to replacement of other carbohydrate sources with sugar-rich supplements have been disappointing in production experiments (e.g. Huhtanen, 1987). This may be associated with an increased production of lactic acid and low postprandial rumen pH on diets rich in sugars (Khalili and Huhtanen, 1991).

The objective of the present study was to evaluate the efficiency of a yeast culture supplement in growing cattle fed a grass silage-based diet. Another specific objective was to determine whether the adverse postprandial effects of a sugar supplement given as sugar beet molasses can be alleviated by yeast supplementation.

MATERIAL AND METHODS

Diets

Silage was prepared from a secondary growth timothy (*Phleum pratense*) sward harvested with a flail-harvester. A formic acid-based additive was applied at the rate of 4 litres t⁻¹ of grass at the time of harvesting. The grass was ensiled in a bunker silo. Grass silage was offered *ad libitum* in amounts of 0.05-0.10 in excess of previous day's intake. The silage was given once daily and the concentrates twice daily in two equal portions. The ingredients of the concentrate mixtures are shown in Table 1. Of the basal concentrate (M₀), a mixture of barley and unmolassed sugar beet pulp (1:1) was gradually replaced with beet molasses at the rates of 100 (M₁₀₀) or 200 g kg⁻¹ (M₂₀₀). Each concentrate mixture was given either without (Y-) or with (Y+) live yeast culture (Western Yeast Culture

Cel-Con; Western Yeast Company, U.S.A.) supplementation. Yeast culture (YC) was top-dressed to the concentrate at a rate of 5 g day⁻¹. The concentrates were given at a rate of 100 g per kg liveweight (LW)^{0.60} on fresh weight basis to maintain a constant forage to concentrate ratio in the diet throughout the experiment. Refusals were removed once daily before the afternoon feeding. The bulls had a free access to water.

Animals and design

Twenty-eight Finnish Ayrshire bulls, with an average liveweight of 268 kg, were divided according to liveweight into five blocks of six (3 blocks) or five (2 blocks) animals. Within the block the animals were allocated at random to the six experimental treatments in a 3 x 2 factorial design (three concentrate supplements without or with yeast culture supplementation). The number of animals for the concentrate treatments were 9-10 and for yeast culture treatments 14. Animals were held in individual stalls.

Experimental procedures

Daily silage and concentrate intakes were recorded individually. The liveweights of the animals were recorded before the morning feeding on two consecutive days at the beginning and at the end of the experiment. During the experiment the animals were weighed at 4-week intervals. The treatments were imposed for periods of 140 (2 blocks) or 168 days (3 blocks). The animals with a heavier initial liveweight remained five 28-d periods in the experiment and those with a lighter initial liveweight six 28-d periods to reach an average LW of 440 kg. Silage was sampled once a week and weekly samples were composited to provide a sample for a 4-week period. Each concentrate was mixed in one batch. The samples were collected during the experiment once a week and composited to provide two samples for each concentrate mixture. At the end of the experiment the animals were slaughtered. Carcass conformation and fat scores were assessed according to Finnish standards. Conformation was described on a 1 to 6 scale (6 best) and fat scores from 0 (very lean) to 5 (very fat).

Chemical and statistical analysis

The chemical analysis of the feed samples and silage fermentation quality were those as described by Jaakkola et al. (1990). Metabolizable energy (ME) contents of the feeds were calculated according to the Ministry of Agriculture, Food and Fisheries (MAFF, 1975). The treatments were arranged factorially and data was subjected to analysis of variance. The model included the effects of block, concentrate, yeast and yeast x concentrate interaction with 18 degrees of freedom

TABLE 1

Composition of concentrate mixtures, g kg⁻¹

Ingredient	Concentrate		
	M ₀	M ₁₀₀	M ₂₀₀
Barley	320.0	283.3	246.6
Molassed sugar beet pulp	320.0	283.3	246.6
Oat hulls	40.0	40.0	40.0
Wheat middlings	140.0	140.0	140.0
Rapeseed meal	105.0	105.0	105.0
Molasses	—	100.0	200.0
Dicalcium phosphate	10.0	8.0	8.0
Sodium chloride	3.0	3.0	3.0
Calcium carbonate	10.0	12.0	12.0
Calcium lignosulphonate	10.0	10.0	10.0
Meat and bone meal	25.0	25.0	25.0
Fat	15.0	15.0	15.0
Trace element premix ¹	1.0	1.0	1.0
Vitamin premix ²	1.0	1.0	1.0

¹ contained (mg g⁻¹): Fe 16.6, Mn 28.5, Cu 10.8, Zn 172, I 3.7 and Co 4.6

² vitamin A (14 000 IU g⁻¹), vitamin D₃ (2 000 IU g⁻¹), vitamin E (11.5 mg g⁻¹), vitamin C (97.5 mg g⁻¹) and Niasin (89.1 mg g⁻¹)

for error. The effects of the type of concentrate supplement were further separated using orthogonal polynomial contrasts into linear and quadratic effects of the rate of molasses inclusion. Because the experiment was not completely balanced, least square (LS) means are presented in the tables. Live-weight gain was calculated either as a difference or using a second degree polynomial regression. Carcass gain was estimated by assuming a dressing proportion of 500 g kg⁻¹ in the beginning of the experiment.

RESULTS

Feed composition

The quality of the silage was not very good as indicated by the low value of 0.627 for the *in vitro* organic matter digestibility and relatively high concentrations of ammonia N, butyric acid and other volatile fatty acids. The differences in the composition of concentrates were as expected (Table 2); NDF and starch contents decreased and sugar contents increased with increasing rate of molasses

TABLE 2

Chemical composition and calculated feeding values of experimental feeds

	Concentrates			
	Silage ¹	M ₀	M ₁₀₀	M ₂₀₀
Dry matter, g kg ⁻¹	236	880	872	868
In dry matter, g kg ⁻¹				
ash	113	81	87	95
crude protein	141	158	155	154
ether extract	54	53	50	48
crude fibre	303	112	105	97
NFE ²	389	596	603	606
NDF	557	368	335	307
ADF	314	133	120	116
ADL	30	25	24	22
cellulose	284	108	96	96
hemicellulose	243	235	215	191
starch	—	229	204	173
WSC ³	18	77	117	160
ME, MJ kg ⁻¹ DM	8.90	12.19	12.20	12.20

¹ in silage: pH 4.17; in DM (g kg⁻¹): lactic acid 87, acetic acid 27, propionic acid 2.8, butyric acid 7.0; in total N (g kg⁻¹): ammonia N 91, soluble N 517

² nitrogen free extractives

³ water soluble carbohydrates

inclusion. The starch content calculated as a difference of OM – (crude protein + ether extract + NDF + WSC) was on average 50 g kg⁻¹ higher than the starch content analysed chemically, probably reflecting a high pectin content of sugar beet pulp. There were no differences in crude protein content or in calculated ME values of the concentrates.

Feed intake and LW gain

No significant differences were observed in silage or total DM intake between the treatments (Table 3). However, the response to YC in daily silage DM intake tended ($P < 0.10$) to interact with concentrate being +0.86 kg day⁻¹ with M₀ and -0.22 kg day⁻¹ with M₂₀₀. Similar interactions were also noted for total DM and calculated ME intakes because the amounts of concentrate refusals were negligible. The average proportion of concentrate in the diet was 390 g kg⁻¹ on DM basis.

The average LW gain declined from 1.21 to 1.09 kg day⁻¹ (linear effect $P < 0.05$) as the proportion of molasses in the concentrate increased (Table 4). The difference was similar when LW gain was estimated using a regression

TABLE 3
Feed and nutrient intake of bulls receiving grass silage with three concentrates each given without (Y-) or with yeast culture (Y+)

Intake, kg DM day ⁻¹	Concentrate			Yeast			SEM ¹	Significance ²		
	M ₁	M ₁₀₀	M ₅₀₀	Y-	Y+	C		Y	Y x C	
silage	4.42	4.60	4.47	4.40	4.60	NS	0.127	NS	+	
concentrate	2.92	2.87	2.85	2.86	2.90	NS	0.019	NS	NS	
total	7.35	7.46	7.32	7.26	7.49	NS	0.138	NS	+	
DM, g kg ⁻¹ LW ^{0.75}	88.3	90.7	89.9	88.59	90.8	NS	1.17	NS	NS	
ME, MJ day ⁻¹	75.1	75.9	74.6	74.1	76.3	NS	1.26	NS	+	
DCP, g day ⁻¹	709	710	695	695	714	NS	11.6	NS	+	

¹ standard error of means
² significance: C = linear effect of the rate of molasses inclusion, Y = effect of yeast supplement, C x Y concentrate x yeast interaction
 NS = non-significance, + P < 0.10, * P < 0.05, ** P < 0.01

TABLE 4
Live weight, liveweight gain and feed conversion of bulls receiving grass silage with three concentrates each given without (Y-) or with yeast culture (Y+)

	Concentrate				Yeast			Significance ²		
	M ₁	M ₁₀₀	M ₃₀₀	M ₃₀₀	Y-	Y+	SEM ¹	C	Y	Y x C
Difference method										
initial LW, kg	267.9	268.1	268.3	268.3	267.7	268.5	1.57	NS	NS	NS
final LW, kg	457.2	449.3	440.0	440.0	446.0	451.7	5.37	NS	NS	*
LW gain, g day ⁻¹	1209	1157	1093	1093	1138	1169	29.1	*	NS	+
feed conversion rate										
DM, kg kg ⁻¹ LWG	6.10	6.45	6.75	6.75	6.41	6.46	0.093	**	NS	NS
ME, MJ kg ⁻¹ LWG	62.3	65.6	68.9	68.9	65.4	65.8	0.99	**	NS	NS
Regression method										
initial LW, kg	267.8	267.9	267.8	267.8	266.8	268.8	1.84	NS	NS	NS
final LW, kg	455.3	448.8	439.7	439.7	445.2	450.6	5.45	NS	NS	*
LW gain, g day ⁻¹	1198	1154	1094	1094	1139	1158	29.7	+	NS	+
feed conversion rate										
DM, kg kg ⁻¹ LWG	6.16	6.46	6.76	6.76	6.40	6.52	0.101	*	NS	NS
ME, MJ kg ⁻¹ LWG	63.0	65.7	69.0	69.0	65.4	65.4	1.08	*	NS	NS

for significance: see Table 3

Slaughter data of bulls receiving grass silage with three concentrates each given without (Y-) or with yeast culture (Y+)

TABLE 5

	Concentrate			Yeast		SEM ¹	Significance ²		
	M ₁	M ₁₀₀	M ₂₀₀	Y-	Y+		C	Y	Y x C
Carcass weight, kg	233.1	230.5	225.7	227.4	232.1	2.86	NS	NS	*
Quality grade ¹	3.6	3.2	3.4	3.2	3.6	0.11	NS	*	NS
Fatness grade ²	0.8	1.0	0.9	0.9	0.9	0.09	NS	NS	NS
Dressing proportion ¹	0.510	0.513	0.513	0.510	0.514	0.0034	NS	NS	NS
Dressing proportion ⁴	0.512	0.514	0.513	0.511	0.515	0.0033	NS	NS	NS
Carcass gain, g day ⁻¹ ³	633	618	582	598	624	16.4	+	NS	NS
Carcass gain, g day ⁻¹ ⁴	639	618	584	601	623	16.6	+	NS	NS

¹ For significance: see Table 3

² 0 = best quality

³ 0 = leanest

⁴ LW gain calculated by difference method

⁵ LW gain calculated by regression method

method. YC supplement had no effect on average LW gain. As it was noted for feed intake, the animals receiving M_0 were more responsive to YC (0.18 kg day^{-1}) than those receiving M_{200} ($-0.10 \text{ kg day}^{-1}$). Feed conversion rate declined ($P < 0.01$) both in terms of DM and ME used per kg LW gain as the proportion of molasses increased.

Carcass data

No significant dietary effects on the dressing proportion were observed between the treatments, and the differences in carcass weight reflected those in liveweight (Table 5). Animals receiving yeast had a better ($P < 0.05$) carcass quality but no significant dietary effects on the fatness grade were observed. Carcass gain declined (linear effect $P = 0.08$) with increasing rate of molasses inclusion.

DISCUSSION

Effect of molasses

Molasses and other sugar supplements have been used to improve the utilization of silage N for rumen microbial protein synthesis. There is a variable degree of proteolysis of plant protein during ensilage, and silage N is rapidly and extensively degraded in the rumen. Feeding grass silage is characterized by high peak values of rumen ammonia N concentration followed by low concentrations for extensive periods between the meals. In reducing rumen ammonia N concentration sugar supplements have been more efficient than starch supplements (Syrjälä, 1972; Chamberlain et al., 1985). This has been assumed to be related to the faster energy release from sucrose than from starch (Syrjälä, 1972). Later studies have, however, shown that starch and barley supplementations increase rumen protozoal numbers (Chamberlain et al., 1985) leading to increased recycling of nitrogen in the rumen. Reduced LW gain of bulls receiving molasses in the present study do not support the idea of synchronization of energy and N release with easily fermentable carbohydrates, rather the reverse was true.

In agreement with the present results, Drennan et al. (1994) reported reduced LW gain in bulls when cane molasses accounted for 400 g kg^{-1} of concentrate DM and for 180 g kg^{-1} total DM. In a Danish study (Andersen and Fredriksen, 1977), the net energy value for cane molasses was 0.70 of that of barley. Similarly, Huhtanen (1987) reported a negative response in dairy cows given a grass silage-based diet when 2 kg of molasses was used to replace on DM basis barley or unmolassed sugar beet pulp.

The reasons for adverse effect of molasses on LW gain are not clear. The rate of inclusion of molasses in the diet was too small to depress rumen cellulolytic activity. In the studies of Huhtanen (1987, 1988) proportions of molasses greater than those used in the present study did not decrease *in vivo* digestion of cell wall constituents, or the rate or extent of forage DM degradation in the rumen. Similar silage DM intake irrespective of the level of molasses in the diet also suggests that reduced LW gain was not related to differences in silage digestibility or in estimated ME intake. It is possible that the lower LW gain on molasses containing diets was related to reduced supply of amino acids from concentrate. The results of Drennan et al. (1994) indicated a greater reduction in LW gain due to molasses inclusion for low protein than for high protein diets. Molasses contains less rumen undegradable protein than barley and sugar beet pulp. Nitrogen in molasses is mainly in non-protein N that probably provides less preformed amino acids and peptides for rumen microbes than barley or sugar beet pulp. However, the contribution of molasses to diet DM (40 and 80 g kg⁻¹) is relatively small to decrease the supply of amino acids markedly. It should also be noted that all supplements contained heat moisture treated rapeseed meal which has sometimes (Aronen et al., 1992) but not always (Huhtanen et al., 1989; Aronen, 1990) improved LW gain in cattle receiving grass silage-based diets.

Because it is not very likely that the origin for reduced LW gain with molasses containing diets lies on inadequate supply of energy and protein yielding nutrients, reduced LW gain must reflect either changes in body composition or in the efficiency of the utilization of ME for growth. Carcass fat scores do not suggest increased fat deposition with molasses in the present study (Table 5), which agrees with results reported by Drennan et al. (1994). Earlier studies (e.g. Lofgreen and Otagaki, 1960) have shown that the net energy (NE) value of molasses depends on its contribution to dietary DM. In cattle the NE value of molasses when it comprised 300-400 g kg⁻¹ of dietary DM was only 50% of that observed at lower levels (100-150 g kg⁻¹). In the study of Heineman and Hanks (1977) the calculated energy value of molasses was very sensitive to the level of inclusion. Molasses DM had feeding value of 1.04 and 0.64 relative to barley DM at inclusion rates of 100 and 140 g kg⁻¹ of DM, respectively. However, in the present study the amounts of molasses used were below these levels. In the study of Huhtanen and Robertson (1988) a trend towards an increase in heat production with sucrose compared with other carbohydrate sources in the diets of sheep also suggests that the efficiency of energy utilization of diets rich in sucrose may be reduced. In their study, the level of sucrose was much higher than those used in the present study. From this discussion it would appear that despite of rather low rate of inclusion, a lower efficiency of energy utilization is the most likely explanation for reduced performance with molasses containing diets.

The effect of yeast supplement

In agreement with other studies in growing cattle (Mutsvangwa et al., 1992; Drennan and Moloney, 1993; Mir and Mir, 1994) YC had only a small effect on LW gain. In studies with cattle offered grass silage, YC supplementation increased LW gain and improved feed conversion efficiency significantly in one of the three trials (Drennan and Moloney, 1993). The effects of YC on digestibility, rumen fermentation pattern and microbial protein synthesis have been small (Huhtanen, 1991; Carro et al., 1992b), and are probably of little biological significance. The published literature shows large variation in response to YC inclusion in the diet, especially in the rumen fermentation pattern but also in ammonia N concentration, digestibility and ruminal forage degradation. The effects of yeast supplements on the supply of nutrients from the diet are discussed in more detail by Huhtanen (1991), Carro et al. (1992b) and Moloney and Drennan (1994). Recent results of Newbold et al. (1995) suggest that differences exist between strains of *S. cerevisiae* in their ability to modify rumen fermentation which could partly explain the variable responses to YC supplementation. It has also been proposed that increased bacterial viability is related to partial removal of oxygen entering the rumen (Wallace, 1994). In animals given silage-based diets the amount of oxygen entering the rumen is likely to be smaller than with dried forages which with large population of viable yeast in silage may explain the limited ability of yeast cultures to stimulate rumen fermentation with silage-based diets.

Concentrate x yeast culture interaction

An interaction ($P < 0.10$) between the inclusion rate of molasses and YC suggested that the effects of YC may depend on the composition of the basal diet. Results from several studies show that the responses to inclusion of YC culture may depend on the composition of the basal diet. Williams et al. (1991) reported that cows given YC increased milk yield when the diet contained 600 g kg⁻¹ of concentrate in the diet but not when the proportion of concentrate was 500 g kg⁻¹. *In vitro* study by Carro et al. (1992a) indicate that the positive effects of YC were greatest on the highest proportion of concentrate in the diet. In contrast to these studies, Flachowsky et al. (1993) reported a much greater depressive effect of YC on ruminal DM degradation of forages with a high concentrate than with low concentrate diets. Fiems et al. (1993) found more pronounced effects of YC with maize-silage than with sugar beet pulp. Moloney and Drennan (1994) observed that the effect of YC on ruminal N metabolism appeared to be dependant on crude protein content of the basal diet. Rumen ammonia concentration was not affected with a low protein diet but it was reduced with a high protein diet when

yeast was added. However, these studies do not provide any explanation for the apparent interaction between the composition of concentrate and dietary inclusion of YC in the present study, and therefore it might be worthy to investigate the mechanisms of interactions between diet composition and yeast. In the present study, the decreasing response to YC with increasing proportion of molasses suggest that a combination of molasses and YC caused changes in rumen metabolism which had an overall negative impact on animal performance.

In conclusion, the present results confirm the earlier findings that the utilization of grass silage-based diets can not be improved by increasing the sugar content in the supplement. In the present study, molasses caused a decreased LW gain at a lower level of inclusion than observed in previous studies. This was partly related to the adverse effect of YC with the highest proportion of molasses. Inclusion of YC had no significant effects on feed intake or LW gain. The effect of YC tended to be dependent on the composition of the concentrate. Because the responses to dietary inclusion of YC in nutrient supply and in animal performance appear to depend on the composition of the diet and probably also on the strain of *S. cerevisiae* used, more research is needed to elucidate conditions in which the use of YC can be economically justified.

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STRESZCZENIE

Wpływ dodatków melasy i kultur drożdży na wyniki opasu buhajków żywionych kiszonką z traw

Doświadczenie opasowe, trwające 168 dni, przeprowadzono na 28 rosnących buhajkach Ayrshire o średniej początkowej masie ciała 268 kg w układzie czynnikowym 3 x 2. Buhajki żywiono kiszonką z traw z dodatkiem pasz treściwych, uzupełnianych odpowiednio w grupach żywymi kulturami drożdży (YC, 5 g/dzień) oraz wzrastającą ilością melasy buraczanej [0 (M_0), 100 (M_{100}), i 200 (M_{200}), g/kg]. Mierzono pobranie paszy, przyrost masy ciała i oceniano jakość tuszy. Melasą zastępowano część paszy treściwej, złożonej ze śruty jęczmiennej i wysłodków buraczanych (1:1). Kiszonkę z traw podawano zwierzętom do woli, a paszę treściwą w ilości 100 g/kg masy ciała ($MC^{0,6}$).

Ilość pobieranej paszy nie zależała od podawanych dodatków melasy i drożdży, natomiast przyrost masy ciała obniżał się liniowo ($P < 0,05$) wraz ze wzrostem ilości podawanej melasy (1,21; 1,16 i 1,09 kg/dzień; SEM 0,029). W miarę wzrostu udziału melasy od 0 do 200 g/kg paszy treściwej, wykorzystanie paszy obniżało się (efekt liniowy – $P < 0,01$) z 6,10 do 6,75 kg SM/kg przyrostu. Dodatek drożdży nie wpłynął na ilość pobieranej paszy (7,26 vs 7,49 kg/dzień) i przyrosty masy ciała (1,14 vs 1,17 kg/dzień). Stwierdzono interakcję drożdże x pasza treściwa; pod wpływem dodatku drożdży przyrosty masy ciała w grupie M_0 , zwiększyły się, a w grupie M_{200} zmniejszyły się.

Różnice w masie tuszy i przyrostach masy tuszy miały podobny układ jak różnice w przyrostach żywej wagi zwierząt. Nie stwierdzono wpływu stosowanych dodatków na wydajność rzeźną i jakość tłuszczu, lecz jakość tuszy zwierząt otrzymujących dodatek drożdży była lepsza.