

The nutritive value of silages Digestion of nitrogenous constituents in sheep receiving diets of grass silage and grass silage and barley.

BY P. C. THOMAS, D. G. CHAMBERLAIN, N. C. KELLY
AND M. K. WAIT

The Hannah Research Institute, Ayr KA6 5HL

(Received 16 July 1979 – Accepted 4 October 1979)

1. Two experiments were conducted to study the digestion of nitrogenous constituents in the rumen, small intestine and caecum and colon of sheep given diets of grass silage or grass silage and barley. Three silages were used. One was made from first-harvest grass in the spring and the others from regrowth grass cut in either early autumn or late autumn. All were of perennial ryegrass (*Lolium perenne*) and were preserved with formic acid.

2. Expt 1 involved a comparison between the spring silage given alone (644 g dry matter (DM)/d) and the spring silage supplemented with barley (151 g DM/d). The intakes (g/d) of total nitrogen for the silage diet and for the supplemented diet were 14.89 and 17.36. Corresponding values (g/d) for N passage were 15.55 and 18.53 ($P < 0.01$) at the duodenum, 6.01 and 7.09 at the ileum and 5.06 and 5.52 in the faeces. The barley supplement had no significant ($P < 0.05$) effect on rumen ammonia-N concentration.

3. Expt 2 involved a comparison between the two autumn-cut silages each offered at a level of feeding of approximately 700 g DM/d. The intakes (g/d) of total N for the early-cut silage and for the late-cut silage were 21.67 and 15.62 respectively. Corresponding values (g/d) for N passage were 17.10 and 16.96 at the duodenum, 6.65 and 6.80 at the ileum and 4.5 and 5.22 in the faeces. The concentration of $\text{NH}_3\text{-N}$ in the rumen was significantly ($P < 0.001$) higher with the early-cut silage than with the late-cut silage.

4. In both experiments the rates of bacterial crude protein ($\text{N} \times 6.25$) synthesis in the rumen, estimated using α, ϵ -diaminopimelic acid as a marker, were low, 142 and 161 g crude protein/kg organic matter apparently digested in the rumen for the spring silage and the spring silage and barley diets respectively, and 68 and 103 g crude protein/kg organic matter apparently digested in the rumen for the early-cut autumn silage and the late-cut autumn silage respectively. For all diets there was a relatively low contribution of bacterial crude protein to the duodenal passage of crude protein and the amounts of individual amino acids ingested in the diets had a marked influence on the amino acids passing to the duodenum and as a consequence on the mixture of amino acids taken up from the small intestine.

5. The results are discussed in relation to the nutritive value of silage N for ruminants.

Ensilage of grass leads to extensive hydrolysis of the plant proteins and to a partial degradation of the amino acids formed (see Oshima & McDonald, 1978). In sheep and cows given highly-digestible perennial ryegrass (*Lolium perenne*) silages prepared using formic acid as an additive a large proportion of the apparently-digested nitrogen is excreted in the urine and although the proportion is reduced when the silage is supplemented with barley (Kelly & Thomas, 1978; McDonald & Thomas, 1978) dairy-cow-feeding experiments have indicated that the nutritive value of dietary crude protein in silage-barley diets is low (Castle, 1975).

Thomas *et al.* (1979) studied the sites of digestion of organic matter, gross energy and carbohydrates in sheep receiving three formic-acid-preserved silages. One silage was made from first-harvest grass in the spring and offered either alone or with a barley supplement, the others were from regrowth grass cut in either early autumn or late autumn. In the present paper results pertaining to the digestion of nitrogenous constituents obtained in the experiments of Thomas *et al.* (1979) are reported, and they are discussed in relation to the nutritive value of silage N for ruminant animals.

Table 1. *The concentration of total nitrogen, non-protein-N and ammonia-N in the diets and the dietary contents of digestible organic matter and rumen-digestible organic matter*

	Expt 1		Expt 2	
	Spring silage	Spring silage and barley*	Early-cut autumn silage	Late-cut autumn silage
Total N (g/kg DM)	23.4	22.0	31.9	22.1
Non-protein-N (g/kg total N)	644	555†	395	538
NH ₃ -N (g/kg total N)	139	120	44	89
Digestible organic matter (g/kg DM)	643	682	744	678
Rumen-digestible organic matter (g/kg DM)‡	421	427	537	506

DM, dry matter.

* For proportions, see p. 470.

† Barley N assumed to be protein.

‡ Digestible organic matter and dietary organic matter apparently digested in the rumen, from Thomas *et al.* (1979).

MATERIALS AND METHODS

Details of the experimental designs, animals, collection procedures and statistical analyses were given by Thomas *et al.* (1979). Two experiments were conducted. Expt 1 involved a comparison between a spring silage (644 g DM/d) and the same silage supplemented with rolled barley (644 g silage DM/d plus 151 g barley DM/d). Expt 2 involved a comparison between two silages (each offered at approximately 700 g DM/d) prepared from the same regrowth grass sward cut at either an early or late stage of maturity in the autumn. Methods of chemical analysis were as described by Kelly & Thomas (1978) and Thomas *et al.* (1979). Samples for amino acid analysis were prepared by hydrolysis with 6 M-hydrochloric acid in sealed tubes at 105° for 24 h. Samples for cystine analysis were prepared by oxidation with 0.86 M-performic acid before hydrolysis. Individual amino acids were separated and determined using an automatic amino acid analyser (LKB Instruments, Croydon, Surrey) following the procedure of Moore *et al.* (1958).

RESULTS

The composition of the diets

The total N contents of the diets and the concentrations of non-protein-N and ammonia-N in the total nitrogen are given in Table 1 together with the dietary contents of digestible organic matter and rumen-digestible organic matter derived from the results of Thomas *et al.* (1979). In both Expt 1 and Expt 2 a large proportion of the total N in each diet was in non-protein form, although the concentrations of NH₃-N in the diets was low. In Expt 1, for the spring silage and barley diet the N content was lower and the digestible organic matter content was higher than for the diet of silage alone; the values for total N:digestible organic matter for the diets were 0.0322 and 0.0364 respectively. In Expt 2 the early-cut silage had a higher total N content and digestible organic matter content than the late-cut silage; the values for total N:digestible organic matter for the diets were 0.0429 and 0.0326 respectively.

Table 2 shows the results for the amino acid composition of the diets. In addition to the amino acids shown the silages contained α,ϵ -diaminopimelic acid in low concentrations, less than 100 mg/kg DM. In Expt 1 both diets had a similar total amino acid (TAA)-N content and amino acid composition.

Table 2. The amino acid composition (g amino acid/kg determined amino acids) of the diets and the dietary content of determined total amino acid nitrogen (TAA-N)

Amino acid	Expt 1		Expt 2	
	Spring silage	Spring silage and barley*	Early-cut autumn silage	Late-cut autumn silage
Aspartic acid	109	102	114	130
Threonine	57	54	59	63
Serine	63	62	61	69
Glutamic acid	160	181	149	136
Proline	97	101	67	69
Glycine	63	60	68	68
Alanine	99	88	88	84
Cystine	11	15	12	10
Valine	70	65	61	63
Methionine	16	17	11	15
Isoleucine	50	45	42	42
Leucine	67	64	92	85
Tyrosine	16	17	25	28
Phenylalanine	36	37	53	49
Histidine	16	18	17	22
Lysine	36	37	34	30
Arginine	34	33	46	39
TAA-N (g/kg DM)	10.3	10.5	17.5	9.5

DM, dry matter.

* For proportions, see p. 470.

Table 3. The mean quantities (g/24 h) of total nitrogen in the food and faeces and of total N and non-ammonia-N (NAN) entering and leaving the small intestine of sheep given diets of spring silage and spring silage and barley (Expt 1) and early-cut autumn silage and late-cut autumn silage (Expt 2) and values for the porportion of apparently-digested N 'disappearing' before the small intestine, in the small intestine and in the caecum and colon

(Mean values with their standard errors; no. of animals/treatment in parentheses)

	Expt 1 (6)			Expt 2 (5)		
	Spring silage	Spring silage and barley	SEM	Early-cut autumn silage	Late-cut autumn silage	SEM
In food: Total N	14.89	17.36	—	21.67	15.62	—
At duodenum:						
Total N	15.55	18.53	0.61**	17.10	16.96	0.99
NAN	14.51	17.16	0.53**	15.82	15.93	0.88
At ileum:						
Total N	6.01	7.09	0.40	6.65	6.80	0.23
NAN	5.57	6.64	0.35	6.17	6.05	0.19
In faeces: Total N	5.06	5.52	0.24	4.50	5.22	0.29
Apparent digestibility of N	0.657	0.683	0.015	0.792	0.666	0.015***
Disappearance of digestible N:						
Before small intestine	-0.079	-0.103	0.050	0.265	-0.149	0.073**
In small intestine	0.979	0.972	0.065	0.610	0.999	0.078**
In caecum and colon	0.098	0.132	0.040	0.125	0.150	0.027

Statistical significance of difference between treatments: ** $P < 0.01$, *** $P < 0.001$.

Table 4. *The amino acid composition (g/kg determined amino acids) of duodenal digesta of sheep given diets of spring silage and spring silage and barley (Expt 1) and early-cut autumn silage and late-cut autumn silage (Expt 2)*

(Mean values with their standard errors; no. of animals/treatment in parentheses)

Amino acids	Expt 1 (6)			Expt 2 (5)		
	Spring silage	Spring silage and barley	SEM	Early-cut autumn silage	Late-cut autumn silage	SEM
Aspartic acid	146	133	3.6*	105	128	3.8**
Threonine	69	75	3.9	53	65	1.9**
Serine	57	52	0.9***	49	56	1.9*
Glutamic acid	138	139	3.8	97	126	7.3*
Proline	47	43	3.0†	77	38	2.7**
Glycine	67	70	1.8	52	76	5.8*
Alanine	80	69	4.1	72	72	5.6
Cystine	13	16	1.1	14	18	1.3*
Valine	66	63	1.2	64	69	4.5
Methionine	16	17	0.9	20	15	2.1
α,ϵ -diaminopimelic acid	4	5	0.4	2	3	0.2***
Isoleucine	45	47	1.3	59	48	2.4*
Leucine	59	66	3.1	104	93	6.7
Tyrosine	35	42	3.0	36	39	4.3
Phenylalanine	46	50	2.6	66	42	4.2**
Histidine	17	19	1.1	6	9	1.2
Lysine	63	60	2.0	76	73	4.6
Arginine	32	37	1.9	50	32	3.6**

Statistical significance of difference between treatments: * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$.

† SEM for spring silage, n 4; for spring silage and barley multiply SEM by 1.07.

Digestion of nitrogenous constituents in the rumen and intestines

Rumen NH₃-N concentrations. With each of the four diets used there were pronounced post-feeding 'peaks' in the NH₃-N content of the rumen fluid; concentrations were maximum in samples taken 1 h or 2 h after feeding. In Expt 1 supplementation of the silage diet with barley had no significant ($P < 0.05$) effect on NH₃-N concentration at any of five sampling times over the period 0-6 h after feeding; mean 'daily' values (mg/l) were 202 for the silage diet and 198 for the silage and barley diet (SEM 17.9; $P > 0.05$). In contrast, in Expt 2 there were significant ($P < 0.05$) differences with diet in rumen NH₃-N concentration at each sampling; mean 'daily' values were (mg/l) 242 (SEM 14.7) for the early-cut autumn silage and 168 (SEM 4.0) for the late-cut autumn silage ($P < 0.001$).

Sites of digestion of N. The spring silage diet in Expt 1 provided 14.89 g total N/d whilst for the spring silage and barley diet the value was 17.36 g total N/d (Table 3). With both diets there was a net gain in total N from the food to the duodenum and the duodenal passage of total N for the barley-supplemented diets was significantly ($P < 0.01$) greater than for the silage diet. This difference closely paralleled the difference between the diets in the intakes of rumen-digestible organic matter which were 273 and 339 g/d for the spring silage diet and the spring silage and barley diet respectively. The passage of total N to the ileum and faeces for each of the diets reflected the passage at the duodenum; the mean digestibilities of duodenal N before the ileum were 0.611 and 0.617 (SEM 0.02; $P > 0.05$), and before the faeces were 0.674 and 0.701 (SEM 0.008; $P < 0.05$) for the spring silage and spring silage and barley diets respectively. There were significant differences between diets

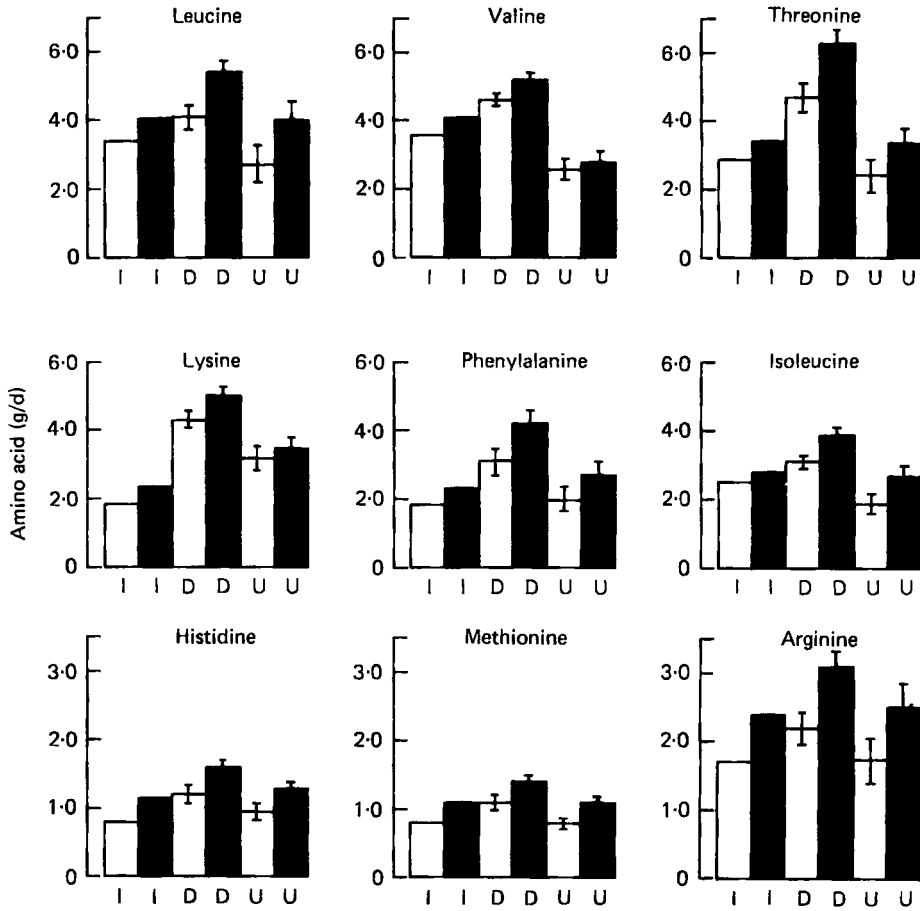


Fig. 1. Expt 1. The amounts of individual 'essential' amino acids in the food (I), passing to the duodenum (D) and taken up between the proximal duodenum and terminal ileum (U) in sheep given diets of (□) spring silage and of (■) spring silage and barley. Values are means with their standard errors represented by vertical bars, for six animals.

in the passage of $\text{NH}_3\text{-N}$ to the duodenum or ileum and the results for NAN closely followed those for total-nitrogen.

In Expt 2, the intake of total N was greater with the early-cut silage than with the late-cut silage (Table 3) but the intakes of rumen-digestible organic matter, 366 and 360 g/d, were similar for both diets. For the early-cut silage there was a loss of N before the duodenum whilst for the late-cut silage there was a gain in N. There were no significant ($P < 0.05$) differences between silages in the passage of total N at the duodenum, at the ileum or in the faeces. The mean apparent digestibilities of duodenal N between the duodenum and the ileum were 0.601 and 0.601 (SEM 0.03) and between the duodenum and the faeces were 0.729 and 0.703 (SEM 0.02) for the early-cut silage and the late-cut silage respectively. There was a large and significant ($P < 0.001$) difference between the silages in the apparent digestibility of dietary N which was due entirely to events within the rumen.

Amino acid composition of duodenal digesta. Despite the differences between diets in the duodenal flow of N in Expt 1, differences between diets in the amino acid composition of the duodenal digesta (Table 4) were small and significant only for aspartic acid ($P < 0.05$) and serine ($P < 0.001$). In contrast in Expt 2, where duodenal flows of N were similar for

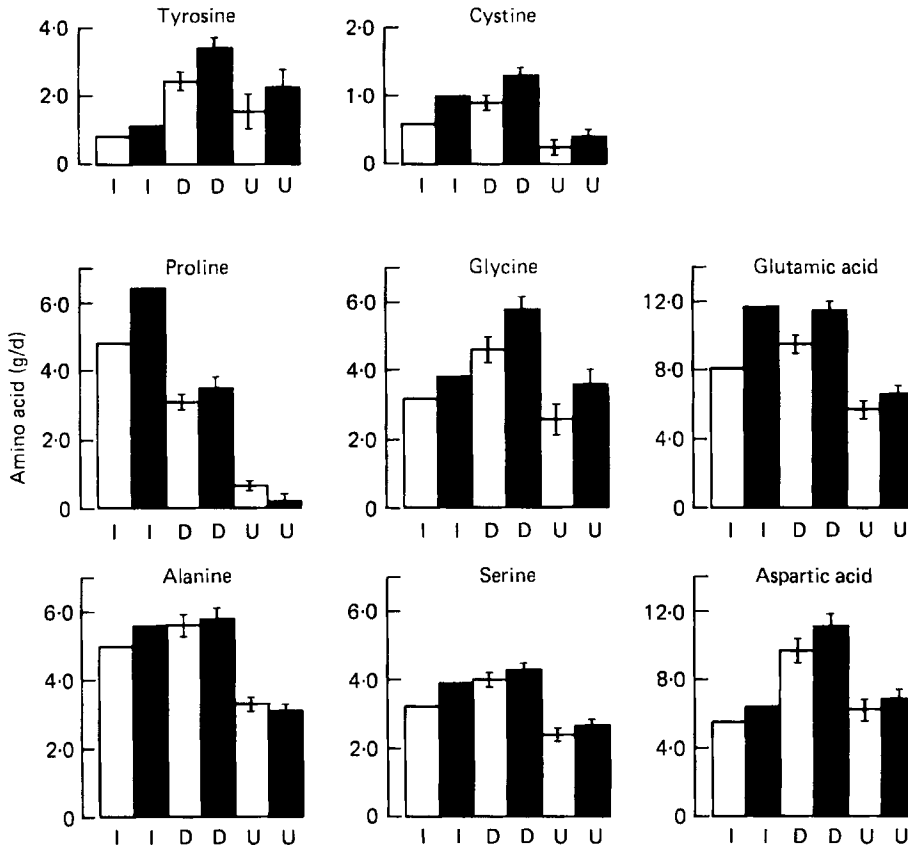


Fig. 2. Expt 1. The amounts of individual 'non-essential' amino acids in the food (*I*), passing to the duodenum (*D*) and taken up between the proximal duodenum and terminal ileum (*U*) in sheep given diets of (□) spring silage and of (■) spring silage and barley. Values are means with their standard errors represented by vertical bars, for six animals.

both diets, there were significant differences between diets in the duodenal concentration of aspartic acid, threonine, serine, glutamic acid, proline, glycine, cystine, α,ϵ -diamino pimelic acid, isoleucine, phenylalanine and arginine (Table 4).

Amino acid intake, duodenal passage and uptake from the small intestine. Figs. 1 and 2 summarize the results from Expt 1 for the dietary intake, passage to the duodenum and net uptake between the duodenum and terminal ileum of individual amino acids. Excepting proline and cystine, the duodenal passage of each amino acid was greater than the amount consumed in the diet and for lysine, phenylalanine, histidine, arginine, tyrosine and aspartic acid the amount taken up from the small intestine was also greater than the intake. Supplementation of the silage diet with barley increased the intake of each amino acid, and there were corresponding and consistent increases for each amino acid in the amount passing to the duodenum and in the amount taken up between the duodenum and ileum. In most instances the differences between diets in the duodenal passage of amino acids were statistically significant ($P < 0.05$) but the standard errors of estimates of amino acid uptake were relatively large and differences between diets in uptake generally did not reach significance at $P < 0.05$.

The results for Expt 2 (Figs. 3 and 4) showed more variable relationships between the

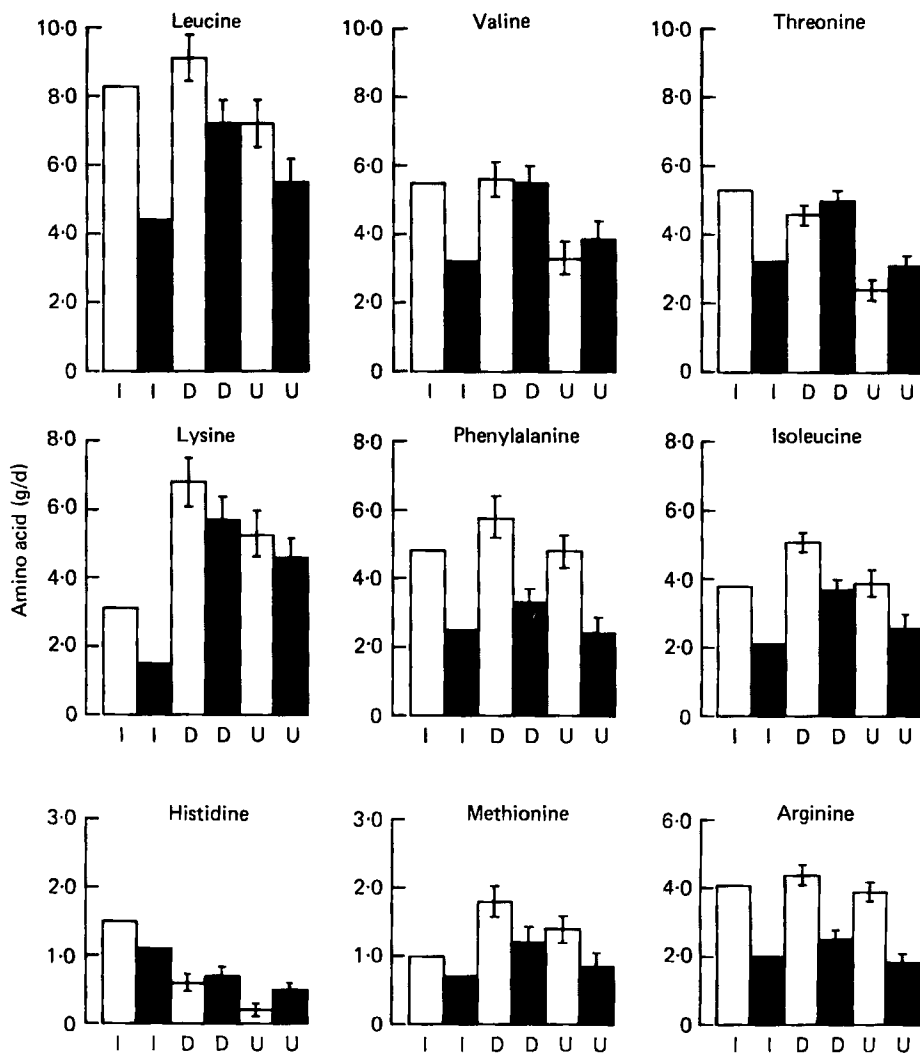


Fig. 3. Expt 2. The amounts of individual 'essential' amino acids in the food (I), passing to the duodenum (D) and taken up between the proximal duodenum and terminal ileum (U) in sheep given diets of (□) early-cut autumn silage and of (■) late-cut autumn silage. Values are means with their standard errors represented by vertical bars, for five animals.

duodenal passage of amino acids and the dietary intake of amino acids than were observed in Expt 1. For some acids there was a net gain from the food to the duodenum with one silage and a net loss from the food to the duodenum with the other, for example see valine, threonine, proline, glycine, glutamic acid, alanine, serine and aspartic acid (Figs 3 and 4). There were significant differences between diets in the duodenal passage of proline ($P < 0.001$), glycine and isoleucine ($P < 0.05$) and phenylalanine and arginine ($P < 0.01$). Differences between diets in intestinal uptake were significant for isoleucine ($P < 0.05$), phenylalanine ($P < 0.01$), glycine ($P < 0.05$) and arginine ($P < 0.01$).

Bacterial crude protein synthesis in the rumen. The passage of bacterial N to the duodenum was calculated from the passage of α, ϵ -diaminopimelic acid (Hogan & Weston, 1970) on the basis that bacteria contain 45 mg α, ϵ -diaminopimelic acid/g total N (Chamberlain &

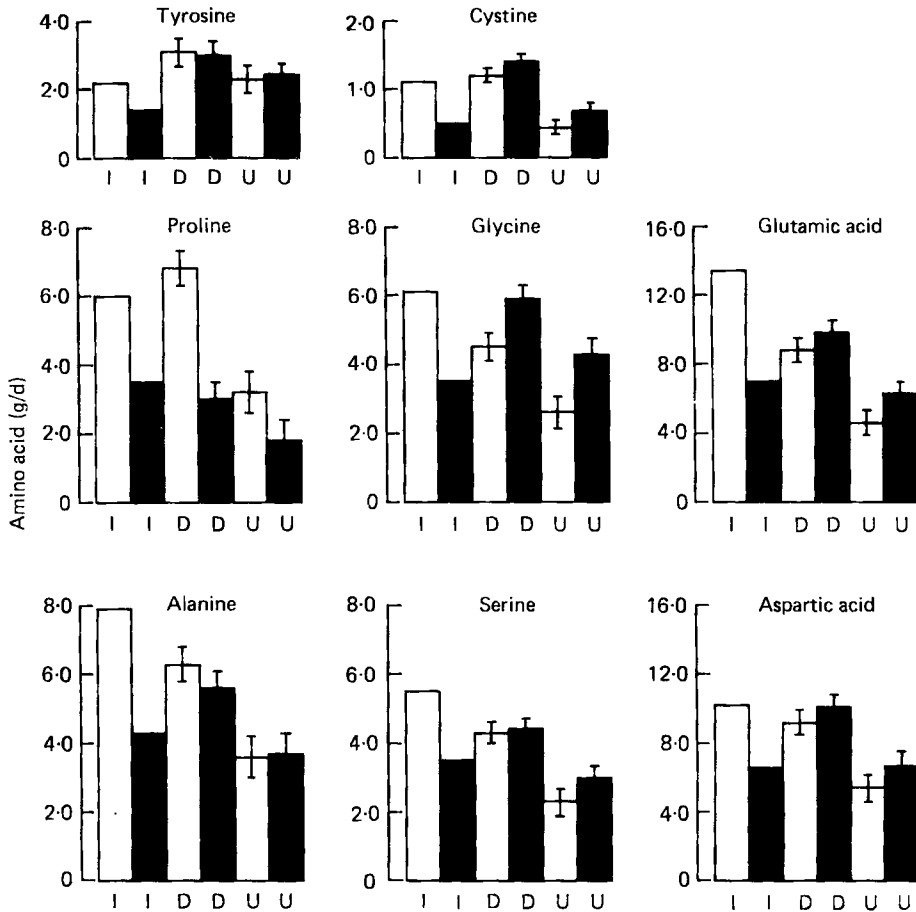


Fig. 4. Expt 2. The amounts of individual 'non-essential' amino acids in the food (*I*), passing to the duodenum (*D*) and taken up between the proximal duodenum and terminal ileum (*U*) in sheep given diets of (□) early-cut autumn silage and of (■) late-cut autumn silage. Values are means with their standard errors represented by vertical bars, for five animals.

Thomas, 1979). Since dead bacterial cells are degraded in the rumen (Nolan & Leng, 1972) and α, ϵ -diaminopimelic acid is metabolized by rumen micro-organisms (Lane & Ling, 1979) it was assumed that the α, ϵ -diaminopimelic acid ingested in the silages was degraded in the rumen, and that the bacterial N passing to the duodenum represented bacterial crude protein synthesized in the rumen.

In Expt 1 the two diets had a similar rumen-digestible organic matter content (Table 1) so that the intake of rumen-digestible organic matter varied with the DM intake; there was a significant ($P < 0.001$) difference between diets in the duodenal passage of bacterial N (Table 5). Rates of bacterial crude protein synthesis per kg organic matter digested in the rumen did not, however, differ significantly between diets (Table 5).

In Expt 2 dietary intakes of rumen-digestible organic matter were similar for both diets but there were significant ($P < 0.01$ and $P < 0.05$) differences between diets in the duodenal passage of bacterial nitrogen (Table 5) and in the rate of bacterial crude protein synthesis (Table 5).

Table 5. The mean quantities of bacterial nitrogen passing to the duodenum and the rates of bacterial crude protein ($N \times 6.25$) synthesis in the rumen in sheep given diets of spring silage and spring silage and barley (Expt 1) and early-cut autumn silage and late-cut autumn silage (Expt 2)

(Mean values with their standard errors; no. of animals/treatment in parentheses)

	Bacterial N (g/24 h)	Bacterial crude protein synthesis (g crude protein/ kg organic matter apparently digested in the rumen)
Expt 1 (6):		
Spring silage	5.91	142
Spring silage and barley†	8.55	161
SEM	0.55***	15
Expt 2 (5):		
Early-cut autumn silage	3.78	68
Late-cut autumn silage	5.92	103
SEM	0.31**	7.5*

Statistical significance of difference between treatments: * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$.

† For proportions, see p. 470.

Table 6. The relationships between the passage of amino acids to the duodenum (g/d) and the intake of amino acids in the diet (g/d) and between the uptake of amino acids in the small intestine (g/d) and the passage of amino acids to the duodenum (g/d) in sheep receiving diets of silage or silage and barley

(The relationships are derived from the mean values observed with the four diets used in Expt 1 and Expt 2)

Amino acid	Relationship between duodenal passage (D) and dietary intake (I)	Correlation coefficient	Relationship between uptake in the small intestine (U) and duodenal passage (D)	Correlation coefficient
Aspartic acid	$D = 11.52 - 0.204I$	-0.54	$U = -1.22 + 0.751D$	0.89
Threonine	$D = 5.95 - 0.209I$	-0.30	$U = -0.02 + 0.549D$	0.87
Serine	$D = 3.88 + 0.090I$	0.48	$U = -0.85 + 0.814D$	0.51
Glutamic acid	$D = 9.95 - 0.005I$	-0.01	$U = -0.59 + 0.646D$	0.86
Proline	$D = 0.42 + 0.707I$	0.51	$U = -0.99 + 0.592D$	0.80
Glycine	$D = 6.33 - 0.271I$	-0.48	$U = -2.31 + 1.060D$	0.95
Alanine	$D = 4.74 + 0.190I$	0.96	$U = 1.55 + 0.320D$	0.33
Cystine	$D = 1.10 + 0.097I$	0.11	$U = -0.37 + 0.690D$	0.91
Valine	$D = 4.38 + 0.203I$	0.43	$U = -1.64 + 0.915D$	0.76
Methionine	$D = 0.13 + 1.339I$	0.74	$U = -0.21 + 0.912D$	1.00
Isoleucine	$D = 1.04 + 1.037I$	0.86	$U = -1.16 + 0.994D$	1.00
Leucine	$D = 2.03 + 0.879I$	0.89	$U = -1.00 + 0.908D$	1.00
Tyrosine	$D = 2.53 + 0.320I$	0.43	$U = -0.08 + 0.743D$	0.81
Phenylalanine	$D = 1.55 + 0.889I$	0.94	$U = -1.10 + 0.994D$	0.97
Histidine	$D = 1.97 - 0.842I$	-0.52	$U = -0.22 + 0.952D$	0.98
Lysine	$D = 3.39 + 0.930I$	0.61	$U = -0.99 + 0.938D$	0.98
Arginine	$D = 0.78 + 0.878I$	0.99	$U = -0.68 + 1.050D$	1.00

DISCUSSION

The results of the experiments described here highlight several features of the digestion of the nitrogenous constituents in formic-acid silages which may have significance in determining the nutritive value of the silage N. In the first instance, it is clear from the results of Expt 1 and Expt 2 that the duodenal passage of total N and NAN does not directly reflect the N intake and varies with the quantity of organic matter digested in the rumen. Relationships between duodenal N passage, dietary N intake and rumen-digested organic matter, similar to those observed here, have been widely reported for dried forage diets. In a summary of results of experiments with Australian forages, Hogan (1975) described the relationship by the equation $Y = 0.33 + 0.0288X$, where Y is the total N reaching the intestine (g/g total N intake) and X is the intake of digestible organic matter (g/g total N intake). For the diets used in the present experiments this equation provides estimates of duodenal total N passage 3–28% higher than those actually observed. Similar over estimates of 3–11% are obtained for three diets containing red-clover silage reported by Thomas *et al.* (1976), and of 20% for the control grass silage used by Beever *et al.* (1977).

In the present experiments the rates of bacterial crude protein synthesis in the rumen were low (Table 5) especially with the early-cut and late-cut autumn silages in Expt 2; rates of bacterial protein synthesis for dried-forage diets are typically 19–23 g crude protein/kg organic matter apparently digested in the rumen (Hogan & Weston, 1970). In both Expt 1 and Expt 2 there was a much lower proportion of bacterial crude protein in the protein passing to the duodenum than is normally observed with dried-forage diets. A similar low proportion of bacterial crude protein in the duodenal crude protein was reported by Thomas *et al.* (1976) for sheep given diets containing red-clover silage. With silage diets much of the crude protein at the duodenum must be of dietary, protozoal or endogenous origin. In sheep endogenous secretions in the abomasum could contribute approximately 2 g N/d to the N passing to the duodenum (Harrop, 1974). Thus of the NAN passing to the duodenum in Expt 1 and Expt 2 dietary plus protozoal N was approximately 6.6, 6.6, 10.0 and 8.0 g/d for the spring silage, spring silage and barley, early-cut autumn silage and late-cut autumn silage diets respectively. The amount of protozoal N was probably small, possibly 1–2 g/d (Weller & Pilgrim, 1974) and this suggests that a substantial part of the N at the duodenum was in undigested dietary crude protein. The intakes (g/d) of true protein-N in the spring silage and spring silage and barley diets in Expt 1 were 5.3 and 7.7 (5.3 from silage) whilst in Expt 2 the corresponding values were 13.2 for the early-cut autumn silage and 7.2 for the late-cut autumn silage. The evidence suggests that the ruminal degradability of the true protein in the silage was low.

There is support for this suggestion elsewhere in the results; there were consistent relationships between the dietary intake and duodenal passage of certain of the amino acids (Table 6). Moreover the uptake of individual amino acids in the small intestine was generally closely related to the entry of the amino acid into the duodenum (Table 6) so that for certain amino acids the intake in the diet had an important influence on the uptake from the small intestine.

As a consequence of the low rate of bacterial protein synthesis and of the low degradability of silage true protein in the rumen, the mixture of amino acids passing to the duodenum and absorbed in the intestine was low in methionine and lysine. In sheep given hay, and hay and cereal diets the duodenal amino acid mixture contained 30–40 g methionine/kg total amino acids and 75–95 g lysine/kg total amino acids (Chamberlain & Thomas, 1979); corresponding values in the present experiments were 15–20 g methionine/kg total amino acids and 60–76 g lysine/kg total amino acids. Intravenous infusion experiments have shown that methionine is limiting for tissue synthesis in sheep given a diet of formic-acid silage

(Kelly & Thomas, 1975) and a limited supply of methionine and lysine to the tissues may be an important factor contributing to the low nutritive value of silage N for dairy cows (see Castle, 1975).

In conclusion, the present results indicate that with formic acid silages of the type used here, the duodenal passage of total N is slightly lower than that which would be predicted for a dried-forage diet of corresponding N and digestible organic matter content. Furthermore, the silage diets appear to have a characteristically-low ruminal synthesis of bacterial crude protein and high duodenal passage of silage true protein which has an important influence on the composition of the amino acid mixture passing to and taken up from the small intestine. The reasons for the resistance of the silage true protein to microbial attack in the rumen and for the low ruminal rates of bacterial protein synthesis require further study; the importance of the dietary supply of essential nutrients such as sulphur to the rumen bacteria remains to be investigated.

The authors are grateful to Mrs A. Maxwell, Mrs A. McLaughlin and Miss A. G. Wilson and her staff for skilled technical assistance, to Mr S. Robertson for care of the experimental animals and to Mr J. N. Watson for making available the grasses and the silage. N. C. K. is grateful to the Meat and Livestock Commission for a postgraduate award.

REFERENCES

- Beever, D. E., Thomson, D. J., Cammell, S. B. & Harrison, D. G. (1977). *J. agric. Sci., Camb.* **88**, 61.
Castle, M. E. (1975). *Agric. Prog.* **50**, 53.
Chamberlain, D. G. & Thomas, P. C. (1979). *J. Sci. Fd Agric.* **30**, 677.
Harrop, C. J. F. (1974). *J. agric. Sci., Camb.* **83**, 249.
Hogan, J. P. (1975). *J. Dairy Sci.* **58**, 1164.
Hogan, J. P. & Weston, R. H. (1970). In *Physiology of Digestion and Metabolism in the Ruminant*, p. 474 [A. T. Phillipson, editor]. Newcastle-upon-Tyne: Oriol Press.
Kelly, N. C. & Thomas P. C. (1978). *Br. J. Nutr.* **40**, 205.
Lane, H. A. & Ling, J. R. (1979). *Proc. Nutr. Soc.* **38**, 80A.
McDonald, L. & Thomas, P. C. (1978). *Proc. 10th int. Dairy Congr.*, p. 63.
Moore, S., Spackman, D. M. & Stein, W. M. (1958). *Analyt. Chem.* **30**, 1185.
Nolan, J. V. & Leng, R. A. (1972.) *Br. J. Nutr.* **27**, 177.
Oshima, M. & McDonald P. (1978). *J. Sci. Fd Agric.* **29**, 497.
Thomas, P. C., Chamberlain, D. G. & Alwash, A. H. (1976). *J. Br. Grassld Soc.* **31**, 123.
Thomas, P. C., Kelly, N. C., Chamberlain, D. G. & Wait, M. K. (1979). *Br. J. Nutr.* **43**, 481.
Weller, R. A. & Pilgrim, A. F. (1974). *Br. J. Nutr.* **32**, 341.