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1	The effect of varying proportion and chop length of lucerne silage in a maize
2	silage-based total mixed ration on diet digestibility and milk yield in dairy
3	cattle
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10	
11	Short title: Inclusion rate and chop length of lucerne silage
12	
13	Abstract
14	The objective was to assess the effects of inclusion rate and chop length of lucerne
15	silage, when fed in a total mixed ration (TMR), on milk yield, dry matter (DM) intake
16	(DMI) and digestion in dairy cows. Diets were formulated to contain a 50:50
17	forage:concentrate ratio (DM basis) and to be isonitrogenous (170 g/kg_CP). The
18	forage portion of the offered diets was comprised of maize and lucerne silage in
19	proportions (DM basis) of either 25:75 (HL) or 75:25 (LL). Lucerne was harvested
20	and conserved as silage at either a long (L) or short (S) chop length. These variables
21	were combined in a 2x2 factorial arrangement to give four treatments (HLL, HLS,
22	LLL, LLS) which were fed in a Latin square design study to Holstein dairy cows in
23	two separate experiments. Sixteen and 8 multiparous, mid-lactation, cows were used
24	in experiments 1 and 2, respectively. To ensure sufficient silage for both
25	experiments, different cuts of lucerne silage (taken from the same sward) were used

26 for each experiment: first cut for experiment 1 (which was of poorer quality) and 27 second cut for experiment 2. Dry matter intake, milk yield and milk composition 28 where measured in both experiments, and total tract digestibility and nitrogen (N) 29 balance were assessed using four cows in experiment 2. In experiment 1 cows fed LL had increased DMI (+3.2 kg/day), compared with those fed HL. In contrast, there 30 31 was no difference in DMI due to lucerne silage inclusion rate in experiment 2. A 32 reduction in milk yield was observed with the HL treatment in both experiment 1 and 33 2 (-3.0 and -2.9 kg/day, respectively). The HL diet had reduced digestibility of DM 34 and organic matter (OM) (-3 and -4%, respectively), and also reduced the efficiency 35 of intake N conversion into milk N (-4%). The S chop length increased total tract 36 digestibility of DM and OM (both +4%), regardless of inclusion rate. Inclusion of 37 lucerne silage at 25% of forage dry matter increased milk yield relative to 75% 38 inclusion, but a S chop length partially mitigated adverse effects of HL on DMI and 39 milk yield in experiment 1 and on DM digestibility in experiment 2.

40

41 Keywords: lucerne, chop length, intake, milk yield, digestibility

42

43 Implications

A high inclusion rate of lucerne at 75% of forage dry matter (DM) within a total mixed
ration (TMR) negatively affected diet digestibility and milk yield relative to a low
inclusion rate. However, a short chop length increased diet digestibility at both
lucerne inclusion rates, and therefore could be used to partly mitigate the negative
effects of high lucerne silage inclusion in diets.

49

50 Introduction

Lucerne (*mendicago sativa*) is widely utilised as a forage legume in dairy cow diets in semi-arid environments including parts of the US, Eastern Europe and Australia. Reduced requirement for inorganic N fertilisation may make it more economical to grow than well fertilised grasses depending on fertiliser price (Phelan *et al.*, 2015) and therefore shows potential for greater use in intensive Northern European dairy systems. Establishing guidelines for the feeding of lucerne in such systems is critical for efficient utilisation.

58 Lucerne and maize silages in the diet are complementary to each other with 59 the former providing rumen degradable protein and the latter providing fermentable 60 energy from starch to drive microbial protein synthesis using ammonia and amino 61 acids from lucerne protein degradation. Previous work has shown that the milk yield 62 obtained from lucerne-maize forage combinations can equal that of grass-maize 63 combinations (Sinclair et al., 2015). However, the optimum inclusion rate of each is 64 not certain. In one study where inclusion rates of chopped lucerne hay to maize 65 silage were varied between 25% and 75% lucerne inclusion within forage dry matter (DM), milk production decreased by 3.3 kg/d with the high rate of lucerne hay 66 67 inclusion (Akbari-Afjani et al., 2014).

68 Lucerne is also a source of physically effective neutral detergent fibre 69 (peNDF) in diets for lactating dairy cows as it has a highly lignified stem that can 70 encourage rumination. Physically effective NDF has been defined as the NDF 71 present in longer particles within a feed (Mertens, 1997), typically considered to be 72 particles greater than 4 mm using the Penn State Particle Separator (PSPS) system 73 (Maulfair and Heinrichs, 2012). Previous research has shown short chop lengths (5 74 mm theoretical length) of lucerne haylage (Kononoff and Heinrichs, 2003) and silage 75 (Beauchemin et al., 1994) can increase DM Intake (DMI) and improve energy

balance relative to long chop lengths of 22 and 10 mm respectively. Therefore, the objective of our study was to examine the effects of lucerne silage chop length on diet DMI and milk yield. A second objective was to investigate how chop length may interact with the inclusion rate of lucerne silage when substituted for maize silage in a TMR. We hypothesised that a lower inclusion rate of lucerne silage and a shorter chop length will increase intake and milk yield in line with previous studies discussed above and that these effects will relate to increased digestibility.

83

84 Material and methods

85 Forage harvesting and clamp sampling

86 This study involved two separate experiments carried out simultaneously at the 87 Centre for Dairy Research (CEDAR), University of Reading, between June and 88 September 2015. The lucerne silage for both experiments was made on-site in the 89 year prior to the start of the trial and conserved in concrete-walled clamps sheeted 90 with a layer of oxygen-barrier film, two layers of plastic sheeting and a weighted top 91 sheet. Experiment 1 utilised a first cut, which was ensiled on 31 May 2014 92 (estimated 10% bloom). The harvested material was windrowed and wilted for 24 h. 93 Alternate swaths originating from the same field area were used to create the two 94 chop lengths, long (L) and short (S), by altering the knife arrangement of the 95 precision chop forage harvester (Claas Jaguar 840 model, Claas Group, Harsewinkel, Germany) from a theoretical chop length of 14 mm (shortest setting) to 96 97 19 mm (longest setting). The long chopped material was collected from the field first 98 followed by short chopped material and each were placed in identical adjacent 99 clamps. The resulting silage was ensiled using Axphast Gold additive containing 100 Lactobacillus Plantarum (Biotal, Cardiff) for low DM silages. The silage produced for

101 Experiment 2 was created on 11 July 2014 in the same way, from the same sward, 102 at second cut (also at an estimated 10% bloom). A longer wilting period of 48 h was 103 allowed, and Axcool Gold additive containing Lactobacillus Buchneri (Biotal, Cardiff, 104 UK) for high DM silages was applied. Following fermentation, core samples for all 105 cuts were taken for chemical composition analyses (Sciantec Analytical Services, 106 Cawood, UK). Maize silage for the study was taken from a commercial crop of mixed 107 varieties harvested in autumn 2014 and ensiled in a concrete-walled clamp with no 108 additive and sheeted as described for the lucerne clamps. The average particle size 109 for the maize silage was determined to be 10mm using a PSPS.

110

111 Diets

112 A TMR with 50:50 ratio of forage:concentrate on a DM basis was fed. The forage 113 was comprised of maize and lucerne silage in proportions (DM basis) of either 25:75 (high lucerne; HL) or 75:25 (low lucerne; LL), respectively. The two inclusion rates 114 115 and the two chop lengths (L or S) were combined in a 2x2 factorial design to give 116 four treatments (HLL, HLS, LLL, LLS). Diets were formulated to be isonitrogenous 117 (170g CP/kg DM) and contain similar levels of NDF (330 and 320 g/kg DM for 118 Experiments 1 and 2 respectively) based on an analysis of core samples from the 119 silage clamps used. Maize meal was included at higher rates in the HL diet to partly 120 offset the reduction in maize starch associated with lower maize silage inclusion in 121 these diets (Table 1), however there was still a significant difference between starch 122 concentration in the resulting TMRs (Table 2).

123

124 *Table 1*

125 *Table 2*

126

127 Animals

128 For Experiment 1, 16 multiparous Holstein-Friesian dairy cows in mid lactation (144 129 d in milk, s.e.m. \pm 4.3) weighing 701 kg and in fourth parity on average, were blocked 130 (4 cows per block) according to milk yield and randomly assigned to one of four initial 131 treatments within each block in a replicated 4x4 Latin square design experiment with 132 three week periods. Cows were housed in a cubicle yard, bedded on sand and 133 individually fed using CALAN gates (American Calan, Northwood, NH, USA). 134 Continual access to water was given. Fresh feed was offered for ad libitum intakes 135 (10% refusals per day) once daily at 1000 h. Refusals were removed on Mondays, 136 Wednesday and Fridays.

137 For Experiment 2, eight multiparous Holstein-Friesian dairy cows in mid 138 lactation (141 d in milk, s.e.m. \pm 13.4) weighing 704 kg and in fourth parity on 139 average, in two blocks (of which one block contained four cows fitted in a previous 140 lactation with Bar Diamond rumen cannula (Parma, Idaho, USA)) were randomly assigned within each block to one of four initial treatments according to a 4x4 Latin 141 142 square design with three week periods as in experiment 1. The block of four non-143 fistulated cows were used for measurements of total tract diet digestion. All 144 procedures carried out in experiment 2 were licensed and monitored by the UK 145 government Home Office under the Animal (Scientific Procedures) Act 1986. 146 Animals were housed in a cubicle yard and individually fed once daily for ad libitum 147 intake through Insentec RIC feeders (Insentec B.V., Marknesse, The Netherlands) 148 during weeks one and two of each period. Cubicles were bedded with wood 149 shavings and continuous access to water was provided. In the final week of each 150 period animals were housed and milked in individual tie stalls situated adjacent to

the cubicle yard to facilitate sampling. Animals were given two days to acclimatise to
the stalls before sampling began. While in tie stalls, animals were fed twice daily at
1000 and 1600 hours for *ad libitum* intake (10% refusals). Refusals were taken daily
at 0930 h.

155

156 Experimental routine

157 Intake and diet analysis. Weights of feed offered and refused were taken during the 158 final week of each period. For the four animals used for digestibility measurements 159 (experiment 2) only measurements from five days were statistically analysed. The 160 DM of the feed offered and refused was measured in a forced air oven at 100°C for 161 24 hours. Bulked daily grab samples of the TMR and diet components were also 162 taken and frozen at -20°C until analysed. Samples of the constituents of the TMR 163 were analysed for DM, N (using the macro kjeldahl method), ash (by combustion at 164 500°C for 16 hours), NDF (assayed with heat-stable amylase, inclusive of residual 165 ash), ADF (inclusive of residual ash), starch, and water soluble carbohydrates (WSC) as described previously (Reynolds et al., 2014). Starch was converted to 166 167 glucose by treatment of the hot water extract with amyloglucosidase followed by acid 168 hydrolysis (Macrae and Armstrong, 1968). Total reducing sugars were measured 169 calorimetrically and the result was corrected for cold water soluble reducing sugars 170 (Fuller, 1967). Crude protein (CP) concentration was calculated by multiplying N 171 (g/kg DM) by 6.25. Concentrations (g/kg DM) of CP, NDF, ADF, ash, starch and 172 WSC in each TMR were calculated based on constituent inclusion rates. 173 Furthermore, TMR and diet components were analysed in triplicate for particle size 174 distribution using a PSPS with holes measuring 4 mm, 8 mm and 19 mm in diameter 175 and a bottom pan. Material from each sieve was collected and dried (at 60°C for 72

h) to give a DM correction. Average particle size of the sample was calculated asdescribed previously (Heinrichs, 2013).

178 Degradability of DM and N in each forage was measured using an *in situ* method 179 with rumen cannulated lactating Holstein dairy cattle (Ørskov and McDonald, 1979). 180 These cattle were housed in cubicles, in a dedicated metabolism unit, fed a 181 commercial grass-maize based TMR diet once daily. Samples (not dried or further 182 chopped) of each silage were placed in polyester bags (40 µm pore size) that were 183 then incubated sequentially in the rumen of three different animals for six time 184 intervals (3, 6, 12, 24, 48, and 72h). Three replicate '0' hour bags were soaked in a 185 tub of cold tap water with agitation for 5 minutes before being refrozen alongside the 186 bags that were incubated in the rumen. Residue was subsequently analysed for DM 187 and N concentration as described above.

188

Milk yield and composition. Cows were milked twice daily at 0630h and 1630h. In experiment 1 separate milk samples were taken during each of the last four consecutive milkings in each period and analysed for fat, crude protein, casein, lactose, urea, and somatic cell count (SCC) by mid infra-red spectroscopy on a CombiFoss machine (National Milk Laboratories, Chippenham, Wiltshire, UK). In Experiment 2 milk samples obtained throughout the third week of each period were analysed as for experiment 1.

196

Diet apparent digestibility and N balance. Beginning at 1000 h (prior to morning
feeding) on day 17 of each period cows used for digestion trials were fitted with a
harness and chute allowing total collection of faeces and urine for five consecutive
24 h periods (Reynolds *et al.*, 2014). Urine was collected into containers containing 1

201 L of 5 molar sulphuric acid. In addition, 200ml spot urine samples were collected 202 twice daily in each of the five consecutive 24h periods, immediately acidified using 203 10ml of 5 molar sulphuric acid, and bulked. At the end of the collection period a 204 representative subsample of the bulked spot samples was obtained and stored 205 frozen until analysed for N. The bulked spot samples were used to determine urinary 206 N concentration to account for any volatilised N losses. At the end of each 24 h 207 period the total faeces and urine collected were weighed. Faeces were mixed, and 208 subsampled as a fixed proportion of total volume to produce a representative bulk 209 sample and stored at -20°C for subsequent analysis. Faecal and feed samples were 210 analysed for DM, N, OM, Starch, NDF, and ADF concentration and urine samples 211 analysed for N concentration as described above for feed samples (Reynolds et al., 212 2014).

213

214 Statistical analysis

215 For silage degradability, an exponential curve fitted to percentage degradation at 216 each time point was used to obtain fractions termed 'a', 'b' and 'c' as described 217 previously (Ørskov and McDonald, 1979). Rumen outflow rate (k) was assumed to 218 be 0.05 hr⁻¹. Feed efficiency was calculated as estimated milk energy yield (Tyrrell 219 and Reid, 1965) divided by DMI. Data from each experiment were analysed 220 separately. Experiment 1 was analysed as four simultaneous Latin Squares. 221 Averages for each cow and treatment combination were analysed to determine fixed 222 effects of square, period, lucerne inclusion rate (IR), chop length (CL), and their 223 interaction (IRxCL) and random effects of cow within square using mixed models 224 procedures of SAS (version 9.1). For experiment 2, data obtained within two 225 simultaneous Latin Squares were analysed in the same way. For each variable the

covariance structure giving the best fit was selected. Data from one cow (not one
used for the digestion trial) in experiment 2 in period four was removed as her DMI
and milk yield did not fully recover following mastitis that occurred during the
adaptation period.

230

231 Results

232

233 Forage quality

234 The first cut silage used for experiment 1 had lower DM (-354 g/kg), and a higher pH 235 (+1.1), than the second cut silage used for experiment 2 (Table 3). Higher DM 236 (second vs first cut) and shorter chop length were associated with lower pH and 237 greater lactic acid concentration but reduced acetic, butyric and propionic acid 238 concentrations. Crude protein concentrations were similar for the first and second cut 239 silages (174 g/kg DM). Of particular note, NDF and ADF were higher in the first cut 240 silages then the second cut, suggesting greater maturity in the first cut silages. 241 The degradability fraction a was smaller in the experiment 1 lucerne silages 242 than in the experiment 2 silages (-13%) and there was also reduced effective 243 degradability and total degradation of DM (fractions a + b = 64% vs 75% for 244 experiment 1 and experiment 2 silages, respectively). Degradation profiles for N 245 showed that the lucerne silages had a higher EPD than that of maize. The rate of 246 degradation of N (c) in the rumen was faster for the short chopped silages for both 247 cuts but the difference was greater within the experiment 2 silages (0.04/h for L and 248 0.09/h for S; *P* < 0.001).

249

250 *Table 3*

251

252 Forage and diet particle size

253 The average silage particle length was 12.6 mm and 9.4 mm in first cut silages (P <254 0.006), and 14.3 mm and 9 mm in second cut silages (P = 0.001) for the long and 255 short chop silages respectively. For both experiments the long chop increased 256 particles retained on the 8 mm sieve and reduced particles on the 4 mm sieve and 257 the bottom pan relative to the short chop silages (P < 0.01; Figure 1). The long chop 258 length increased the proportion of particles on the 19 mm sieve for the lucerne silage 259 used in experiment 2 (P < 0.001), but not the lucerne silage used for experiment 1 260 (Figure 1).

261

262 *Figure 1*

263

264 In both experiments, average particle size of the diets fed (Table 4) increased with 265 both greater lucerne inclusion (P < 0.001) and chop length (P < 0.05, < 0.001 in 266 experiments 1 and 2 respectively). In experiment 1, the proportion of particles 267 retained on the 19 mm screen increased (P < 0.02) with increased chop length. The 268 proportion of particles retained on the 4 mm screen in experiment 1 was decreased 269 (P < 0.03) by increased chop length for the HL, but not the LL diet (inclusion rate by 270 chop length interaction, P = 0.03). In experiment 2, there were greater effects of 271 chop length on particle distribution on the 19 and 4 mm screens for the HL than the 272 LL diets (inclusion rate by chop length interaction, P < 0.01) and a greater difference 273 on the 8 mm screen for the LL than the HL diet (inclusion rate by chop length 274 interaction, P < 0.05).

276 *Table 4*

277

278 Intake, milk yield and milk composition

279 The effect of lucerne silage inclusion rate on DMI varied between experiments with a 280 DMI reduction of 3.2 kg/d where HL diets were fed in experiment 1 (P < 0.001), 281 whereas there was no difference in DMI between treatments in experiment 2 (P >282 0.22). In both experiments feeding the HL diets decreased milk yield (-3.0 and -2.9 283 kg/d in Experiments 1 and 2 respectively; P < 0.02; Table 4). In experiment 1, a 284 longer chop length decreased milk yield relative to using a shorter chop length by -285 1.6 kg/d (P < 0.001), although this effect was not observed in experiment 2. As a 286 result, the estimated conversion efficiency of feed DM into milk energy also differed 287 between experiments, with HL diets tending to produce greater conversion efficiency 288 in experiment 1 (P < 0.08) and LL diets increasing feed efficiency in experiment 2 (P289 = 0.001).

290

291 *Table 5*

292

293 Milk fat concentration was not affected by treatment in either experiment (Table 5), 294 however, in experiment 1, milk fat yield was greater (P < 0.017) when LL diets were 295 fed. In experiment 1, milk protein concentration was increased by 0.7 g/kg (P < 0.001) 296 when HL diets were fed, although, due to increased milk yield, milk protein yield was 297 highest (P < 0.001) when LL diets were fed. In experiment 2, feeding HL diets led to a 298 decrease in milk protein concentration of 1.0 g/kg (P < 0.04) although there were no 299 differences in total protein yield between treatments. Milk protein yield in experiment 300 1 was reduced by chop length, where a 45 g/d reduction with longer chop length (P < 301 0.003) was observed. Milk urea concentration was higher (P < 0.001) when HL diets 302 were fed in experiment 1.

303

304 Apparent digestibility and N balance

305 Increasing lucerne inclusion rate decreased DM and starch intake and increased

- ADF intake of cows used for measurements of digestibility and N-balance (P < 0.04,
- 307 < 0.003, and < 0.006 respectively, Table 6). Digestibility of DM was lower for the HL</p>
- diets by 3.6% relative to the LL diets (P < 0.05). Increasing chop length also reduced

309 DM digestibility by 4.3% (P < 0.02). Greater inclusion rate of lucerne and longer chop

- length both decreased the digestibility of organic matter by 3.7% and 3.2% (P < 0.03
- and *P* < 0.006, respectively). There were no differences in the digestibility of starch,
- NDF or ADF between HL and LL diets, although NDF digestibility tended (P < 0.10)

to be lower for longer lucerne chop length diets.

314

315 *Table 6*

316

317 Intakes of N were greater for LL diets (P < 0.01) as a result of higher DMI (Table 7). 318 There was a tendency forere increaseds in faecal N concentration (P < 0.064) and total manure (faecal plus urine) N excretion (P < 0.03) when HL diets were fed. 319 320 Faecal N also tended to increased when the chop length was increased (P < 321 0.0709). There was greater partitioning of intake N into the milk for the LL diets with 322 an increase in N use efficiency of 3.3% (P < 0.009) and N digestibility was also greater (P < 0.02). Shortening the chop length of the lucerne reduced faecal N 323 324 excretion by 57 g/d (P < 0.01). 325

326 *Table 7*

327

328 Discussion

329 Forage quality and particle size

330 The nutritive value of the four lucerne silages used in the study was variable.

Although crude protein levels were similar at 172 g/kg, the second cut (experiment 2)

332 silages were lower in NDF and ADF than the first cut (experiment 1) silages,

\$33 suggesting increased maturity in the first cut relative to the second cut forage. In the

334 <u>silage fed in experiment 1, high acetic acid and low lactic acid concentrations</u>

\$35 indicated poor fermentation, although pH reduction was adequate. High levels of

WSC in the experiment 2 silage may indicate that increased time spent wilting this

337 crop (48h vs. 24h for the first cut) resulting in a higher DM reduced fermentation

activity, or that the original concentration of sugar in this crop was higher than for the

first cut crop. These results collectively indicate increased silage quality in the

340 second cut silage with higher DM concentration. The effective degradation of DM

and protein in the lucerne silages ranged from 37.8-56.7% and 72.6-78.6%

respectively which are similar to previously published figures (56% for EDMD and

343 72% EPD for mid-bloom fresh lucerne (Hoffman *et al.*, 1993)).

Variation between the long and short chop silages within each experiment was observed despite care being taken at harvest to control variables other than chop length. Notably, pH and acetic, propionic and butyric acids were reduced for both short cut silages relative to long cut silages while lactic acid was increased. This may be explained by increased silage density through better compaction achieved with the short chop which helps to create the necessary anaerobic environment in the silo. Short cut silage was also collected from the field after long chopped silage leading to a small increase in wilting time which may also have increased theconcentration of sugars available for fermentation.

353 The differences in physical structure achieved by varying chop length of the 354 silages were similar for both experiments. The theoretical difference between 355 average particle size according to the settings of the forage harvester was 5 mm 356 which was relatively close to the achieved differences of 3.6 mm and 5.3 mm for 357 experiments 1 and 2 respectively. The differences in mean particle lengths achieved 358 by varying chop length in this study are similar to those used in previous research 359 (e.g. 5mm, Beauchemin et al. (1994); and 7mm, Bhandari et al., (2007)). Although 360 the difference in mean particle length is small, there were larger differences in the 361 relative quantities of particle size fractions measured using a PSPS.

362

363 Intake, milk yield and nutrient digestibility

364 Effects of lucerne inclusion rate. The effects of a higher dietary inclusion rate of 365 lucerne on DMI differed in the two experiments. In experiment 1, feeding the first cut 366 silage at the higher inclusion rate decreased DMI, and milk yield. In contrast, 367 feeding the second cut silage at the same rate to a smaller number of cows had no 368 effect on DMI, but a reduction in milk yield was still observed. In a similar UK study 369 where grass silage was replaced with lucerne silage in the diet a reduction in intake 370 was also seen when 60% of forage DM was comprised of lucerne silage (Sinclair et 371 al., 2015). Rumen fill can be limiting factor on DMI depending on the the extent to 372 which the diet is comprised of forage (Beauchemin et al., 2003). In this case the 373 second cut silages showed 11% greater total DM degradability (a+b) over 72 hours 374 than the first cut silage indicating that the first cut silage would have contained a 375 greater mass of forage dry matter within the rumen during this time. The differences

in forage DM degradability might explain conflicting effects on DMI seen between
experiments. Furthermore, the silage used in experiment 1 had a high concentration
of acetic acid which <u>h</u>was been linked with reduced intake in some studies where
dietary concentrations were 25-50 g/kgDM (Daniel *et al.*, 2013), and this may have
also contributed to lower intakes on HL diets in this instance.

381 Feeding HL diets led to a reduction in the digestibility of both DM-, and OM and N. 382 This indicates that the lucerne silages used in this study were less digestible than the 383 maize silage, reflecting greater ADF concentration. Decreases in milk yield observed 384 for the HL diets in both experiments could be related to this reduction in DM and OM 385 digestibility, and therefore ME, and also the lower starch concentration of the HL 386 diets (Table 2). Furthermore, there would have been a greater imbalance between 387 supply of metabolisable protein to ME in HL diets which would contribute to the 388 reduction in milk yield observed. These findings align with previous studies which 389 show that lucerne typically has a lower ME content than many other forage legumes 390 or grasses (Steinshamn, 2010). Efficiency of N utilisation was also reduced when HL diets were fed, which is partly attributable to lower milk yield seen on HL diets. Also, 391 392 since diets were not balanced for rumen degradable N supply in this study, there 393 was a surplus of rapidly degradable N for the HL diets which would have contributed 394 to reduced N utilisation and high milk urea values. There was greater partitioning of 395 N into faeces than urine in this study, which may in part reflect low N digestibility, 396 particularly on HL diets. In addition, the spot sampling method adopted in this study 397 may not have fully accounted for diurnal changes in urine concentration. 398

Effects of lucerne chop length. In experiment 1, DMI increased when a short chop
silage was fed which was also observed with lucerne haylage (Kononoff and

401 Heinrichs, 2003) but was contrary to the results of experiment 2 and a numerous 402 previous studies in which reducing lucerne chop length did not affect DMI 403 (Beauchemin et al., 1994 and 2003; Bhandari et al., 2007). Increased DMI with 404 shorter chop length might suggest increased speed of particle breakdown and/or 405 rumen outflow and lower rumen fill allowing higher rates of intake relative to the 406 longer chop length. Some studies found this to be the case where short and long 407 chop lucerne lengths were compared (using mean particle lengths of 1 and 8 mm; 408 Yansari et al., 2004) although others noted no change in passage rate even when an 409 effect on DMI was observed (Kononoff and Heinrichs, 2003). The short chop length 410 also increased DM digestibility, and the magnitude of the effect was greater than that 411 of inclusion rate. This may be explained by smaller particles exhibiting a greater 412 surface area for microbial attachment, although this is only one of many factors that 413 govern the rate of cellulolysis in the rumen (Mason and Stuckey, 2016). The increase 414 in digestibility might also explain why the short chop mitigated some of the negative 415 effect of the HL diet on milk yield (+1.6 kg/d milk produced on LL diets; Table 6). 416 Using shorter chop lengths with high lucerne silage diets shows potential as a 417 strategy to partly mitigate reduced nutrient use efficiency when lucerne is included at 418 higher rates in the diet. Further research into lucerne agronomy and variety 419 development for delayed plant lignification to increase the acceptable harvest 420 window is an approach which may improve prospects for feeding lucerne in the 421 future.

422

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- 430

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487 **Table 1** Ingredients used to create experimental total mixed rations in two separate

488 experiments.

	Diet						
-	Experi	ment 1	Experi	ment 2			
Item	LL	HL	LL	HL			
Ingredients, g/kg DM							
Lucerne silage	125	375	125	375			
Maize silage	375	125	375	125			
Concentrate blend							
Cracked Wheat	80	80	80	80			
Maize Meal	61	70	54	97			
Unmolassed Sugar Beet Feed	40	40	40	40			
Soy Hulls	79	88	82	108			
Soybean Meal	98	89	100	65			
Rapeseed Meal	98	89	100	65			
Molasses	10	10	10	10			
Dicalcium phosphate	5	5	5	5			
Salt	5	5	5	5			
Dairy Mineral	10	10	10	10			
Megalac ¹	15	15	15	15			

489 LL = low lucerne diet; HL = high lucerne diet;

490 ¹ Megalac rumen protected fat supplement (Volac International ltd., Royston, UK)

492 **Table 2** The chemical composition of four total mixed rations containing a high (HL)

493 or la	<i>эw (LL)</i> -	concentration	of lucerne	silage at a	long (L)	or short (S)	chop	length f	fed
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		D	iet				P value	;
Item ¹	LLS	LLL	HLS	HLL	SEM	IR	CL	IRxCL
Experiment 1, g/kg DM								
DM, g/kg	467	424	358	334	8.8	0.001	0.011	0.366
Ash	65	66	86	86	0.8	0.001	0.844	0.844
СР	181	179	163	171	3.4	0.138	0.367	0.343
NDF	334	337	329	348	6.0	0.689	0.164	0.280
ADF	269	244	227	222	4.4	0.096	0.163	0.283
Starch	256	256	165	155	7.9	0.044	0.589	0.609
WSC	37	36	27	27	2.5	0.009	0.993	0.896
n	4	4	4	4				
Experiment 2, g/kg DM								
DM, g/kg	553	572	611	635	3.3	0.001	0.002	0.386
Ash	61	62	77	78	0.4	0.001	0.350	0.946
СР	170	170	171	174	2.5	0.115	0.435	0.535
NDF	318	321	327	338	2.8	0.002	0.026	0.206
ADF	204 ^a	208 ^a	234 ^b	236°	1.7	0.001	0.004	0.042
Starch	242	242	162	164	2.1	0.001	0.195	0.597
WSC	37	36	34	32	0.4	0.001	0.001	0.105
n	7	8	8	8				

494 *in two separate experiments.*

495 IR = Inclusion rate; CL = chop length; IRxCL = interaction between IR and CL; DM = dry matter; OM =

496 organic matter; WSC = water soluble carbohydrate.

497 ^{a,b} Where there is a significant interaction, values within a row with different superscripts differ

498 significantly at *P*<0.05.

499 Table 3 Analysis of the chemical composition and degradability characteristics of

500 four lucerne silages harvested at first cut (used in experiment 1) or second cut (used

		L	_				
	Maize	Exp	o. 1	Ex	р. 2	SEM	<i>p</i> value
Item	silage	L	S	L	S		
Chemical composition ¹ , g/kgDM							
DM, g/kg	384 ^a	218 ^b	225 ^b	587°	559°	10.0	0.001
СР	73 ^a	176 ^b	175 ^b	170 ^b	175 [⊳]	6.4	0.001
ОМ	965 ^a	874 ^b	875 ^b	892°	893°	2.1	0.001
NDF	368 ^a	513 ^b	498°	408 ^d	390 ^{ad}	7.6	0.001
ADF	215 ^a	441 ^b	418 ^c	355 ^d	328 ^e	4.7	0.001
Starch	376	-	-	-	-	-	-
WSC	3 ^a	1 ^a	1 ^a	10 ^b	16 ^b	1.1	0.001
n	4	4	4	4	4		
Fermentation characteristics ²							
рН	-	6.2	5.6	4.9	4.7	-	-
Ethanol, g/kgDM	-	22.8	6.13	0.05	1.36	-	-
Lactic acid, g/kgDM	-	<5.0	7.29	27.1	43.1	-	-
Acetic acid, g/kgDM	-	56.9	40.7	7.77	0.97	-	-
Propionic acid, g/kgDM	-	8.07	4.80	0.51	0.39	-	-
Butyric acid, g/kgDM	-	41.3	15.6	1.02	0.72	-	-
Degradability parameters ³							
DM degradability							
a, %	44.1 ^a	26.5 ^b	16.5°	32.7 ^d	34.6 ^d	0.43	0.001
b, %	37.6 ^a	38.2 ^a	46.5 ^b	42.8 ^{ab}	40.4 ^a	1.76	0.037
c, %/h	3.26 ^a	4.34 ^a	4.24 ^a	3.82 ^a	6.02 ^b	0.495	0.037
EDMD, %	58.6 ^a	44.2 ^b	37.7°	51.3 ^d	56.5 ^e	0.72	0.001
Protein degradability							
a, %	62.3 ^a	67.2 ^b	63.0 ^a	59.0°	61.9 ^{ac}	0.91	0.003
b, %	24.2	24.2	21.2	30.6	26.4	4.04	0.593
c, %/h	1.95 ^a	2.95 ^{ab}	4.19 ^b	3.97 ^b	8.63 ^c	0.614	0.001
EPD, %	67.4 ^a	74.8 ^b	72.6 ^c	72.5 ^c	78.6 ^d	0.76	0.001
n	3	3	3	3	3		

501 in experiment 2) at either a long (L) or short (S) chop length.

502 DM = dry matter; OM = organic matter; WSC = water soluble carbohydrates; ME = metabolisable

503 energy; EDMD = effective dry matter degradability; EPD = effective protein degradability.

504 ¹ Average chemical composition from analyses of bulk samples taken in each period of the study

505 analysed using mixed models with fixed effect of silage and period.

506 ² The analysis from clamp core samples taken at 3 separate points in the clamp and bulked.

507 ³ Degradability parameters determined by *in sacco* incubation in the rumen, using the model of

508 (Ørskov and McDonald, 1979) where a = rapidly soluble material; b = non-soluble but degradable

- 509 material; c = rate f degradation of b; effective degradability = a+b[c/(c+k)] (Ørskov and McDonald,
- 510 1979) where k = an assumed outflow rate of 0.05/hr. Mean values from each of 3 cows were analysed
- 511 using mixed models with fixed effects of silage and random effects of cow.
- 512 ^{a,b} Where there is a significant interaction, values within a row with different superscripts differ
- 513 significantly at *P*<0.05.
- 514

515 Table 4 The distribution of particle size (DM basis) in four total mixed rations

516 containing a high (HL) or low (LL) concentration of lucerne silage at a long (L) or

		D	iet				P value	;
Item ¹	LLS	LLL	HLS	HLL	SEM	IR	CL	IRxC
Experiment 1								
Material retained, %DM								
19mm	8.4	9.6	21.7	30.0	1.67	0.001	0.018	0.10
8mm	39.9	32.8	41.3	39.9	0.60	0.001	0.421	0.22
4mm	17.3ª	17.2ª	21.4 ^b	16.2ª	0.86	0.148	0.025	0.03
Bottom pan	35.4	31.1	24.7	20.6	1.40	0.074	0.189	0.96
Mean particle size, cm ¹	0.62	0.67	0.82	0.97	0.415	0.001	0.046	0.23
n	4	4	4	4				
Experiment 2								
Material retained, %DM								
19mm	3.2ª	5.0ª	5.3ª	12.1 ^b	0.75	0.001	0.001	0.00
8mm	36.4 ^a	41.9 ^b	37.4 ^{ac}	39.1°	0.50	0.129	0.012	0.02
4mm	16.5ª	13.5 ^b	18.7°	12.6 ^b	0.24	0.033	0.001	0.00
Bottom pan	43.8	39.8	37.9	36.3	0.50	0.001	0.010	0.09
Mean particle size, cm	0.50	0.56	0.54	0.65	0.014	0.001	0.001	0.09
n	3	4	4	4				

short (S) chop length in two separate experiments. 517

518

519 ¹ Mean particle size was determined using the recommended equation of Penn State University 520 (Heinrichs, 2013).

521 ^{a,b} Where there is a significant interaction, values within a row with different superscripts differ 522 significantly at P<0.05.

523

525 **Table 5** Dry matter intake, milk yield, milk composition and feed conversion

- 526 efficiency of lactating dairy cows fed a total mixed ration containing a high (HL) or
- 527 low (LL) concentration of lucerne silage at a long (L) or short (S) chop length in two
- 528 separate experiments.

	Diet					P value			
Item ¹	LLS	LLL	HLS	HLL	SEM	IR	CL	IRxCL	
Experiment 1									
DMI, kg/d	26.4	26.0	23.7	22.3	0.74	0.001	0.017	0.172	
Milk yield, kg/d	35.2	33.9	32.5	30.6	1.04	0.001	0.001	0.449	
Est. Milk energy, MJ/d ¹	101.4	100.0	93.6	89.7	4.12	0.001	0.073	0.379	
Energy efficiency, MJ/kg ²	3.84	3.85	3.99	4.00	0.119	0.079	0.926	0.970	
Milk composition									
Milk fat, g/kg	36.9	37.7	37.6	38.3	1.37	0.263	0.242	0.705	
Milk protein, g/kg	30.2	30.5	31.2	30.9	0.68	0.001	0.962	0.066	
Milk urea, mg/kg	292	311	424	432	14.4	0.001	0.088	0.469	
Fat yield, kg/d	1.29	1.28	1.21	1.21	0.065	0.017	0.844	0.954	
Protein yield, kg/d	1.10	1.06	1.00	0.95	0.036	0.001	0.003	0.706	
n	16	16	16	16					
Experiment 2									
DMI, kg/d	23.0	23.0	23.8	23.7	0.75	0.227	0.994	0.916	
Milk yield, kg/d	31.5 ^{ab}	33.7 ^b	30.8ª	28.7ª	2.21	0.013	0.953	0.043	
Est. Milk energy, MJ/d ¹	85.5 ^{ab}	91.9 ^a	86.1 ^{ab}	82.1 ^b	5.95	0.074	0.636	0.047	
Energy efficiency, MJ/kg ²	3.73	3.95	3.59	3.54	0.249	0.002	0.409	0.068	
Milk composition									
Milk fat, g/kg	35.0	33.9	35.1	35.9	1.53	0.357	0.907	0.378	
Milk protein, g/kg	30.1	30.6	29.7	29.0	0.62	0.034	0.768	0.146	
Milk urea, mg/kg	291	306	324	333	23.0	0.105	0.508	0.862	
Fat yield, kg/d	1.03	1.08	1.09	1.12	0.831	0.104	0.216	0.795	
Protein yield, kg/d	0.92	0.95	0.92	0.90	0.667	0.261	0.842	0.387	
n	7	8	8	8					
IR = Inclusion rate; CL = chop ler	ngth; IRx(CL = inte	raction be	etween I	R and C	L; DMI =	dry matte	er	

530 intake.

529

531 ¹ Estimated milk energy = Milk yield, kg*((fat concentration, g/kg *0.0384+protein concentration, g/kg

532 *0.0223+lactose concentration, g/kg *0.0199)-0.108)

533 ² Energy Efficiency calculated as Estimated milk energy in MJ/d divided by DMI in kg

534 ^{a,b} Where there is a significant interaction, values within a row with different superscripts differ

535 significantly at *P*<0.05.

536

Table 6 The apparent DM, OM, NDF, ADF and starch digestibility of four total mixed
rations containing a high (HL) or low (LL) concentration of lucerne silage at a long (L)
or short (S) chop length when fed to lactating dairy cows (in experiment 2).

		D	iet		P valu			<u>;</u>
Item ¹	LLS	LLL	HLS	HLL	SEM	IR	CL	IRxCL
Dry matter								
DMI, kg/d	23.8	25.0	23.1	22.1	0.57	0.040	0.862	0.116
Faecal DM, kg/d	6.70	7.90	7.70	8.89	0.445	0.022	0.010	0.992
DM digestibility, %	70.4	67.2	67.9	62.5	1.40	0.043	0.015	0.424
Organic Matter								
OM intake, kg /d	20.9	22.7	22.0	22.0	0.91	0.846	0.376	0.368
Faecal OM, kg/d	5.66	6.69	6.38	7.42	0.378	0.029	0.007	0.984
OM digestibility,%	73.1	70.5	71.0	66.2	1.22	0.021	0.006	0.292
Starch								
Starch intake, kg/d	5.28	5.73	3.98	4.00	0.257	0.002	0.407	0.438
Faecal starch, kg/d	0.16	0.24	0.14	0.16	0.034	0.015	0.019	0.081
Starch digestibility, %	96.7	95.8	96.9	96.3	0.78	0.668	0.250	0.858
Fibre								
NDF intake, kg/d	6.99	7.69	7.85	8.07	0.313	0.107	0.208	0.492
Faecal NDF, kg/d	3.05	3.56	3.43	3.77	0.200	0.095	0.032	0.564
NDF digestibility, %	56.0	52.8	56.9	53.8	2.04	0.572	0.095	1.000
ADF Intake, kg/d	4.51	5.01	5.59	5.86	0.207	0.005	0.135	0.628
Faecal ADF, kg/d	2.30	2.69	2.56	2.93	0.355	0.616	0.490	0.985
ADF digetsibility, %	50.4	47.3	52.3	50.0	2.67	0.381	0.309	0.863
n	4	4	4	4				

541 IR = Inclusion rate; CL = chop length; IRxCL = interaction between IR and CL; DM = dry matter; DMI

542 = dry matter intake; OM = organic matter.

Table 7 The apparent digestibility of N and N balance in lactating dairy cows fed total
mixed rations containing a high (HL) or low (LL) concentration of lucerne silage at a
long (L) or short (S) chop length (in experiment 2).

	Diet						P value	
Item ¹	LLS	LLL	HLS	HLL	SEM	IR	CL	IRxCL
N intake, g/d	674	692	654	635	10.7	0.010	0.953	0.067
Faecal N, g/d (a)	<u>209</u> 197	2 <u>41</u> 51	2 <u>42</u> 33	2 <u>78</u> 92	1 <u>9.2</u> 7.6	0.0 <u>54</u> 39	0.0 <u>61</u> 09	0. <u>921</u> 877
N digested, g/d	<u>389^{ab}430</u>	4 <u>53^{ab}28</u>	4 <u>65^b</u> 47	3 <u>80ª</u> 77	<u>10.4</u> 22.3	0. <u>820</u> 481	0. <u>263</u> 164	0. <u>041</u> 185
N digestibility, %	70. <u>3</u> 2	6 <u>5.6</u> 4.5	6 <u>2.3</u> 4.0	5 <u>6.4</u> 5.4	<u>2.37</u> 4 .26	0. <u>019</u> 286	0. <u>053</u> 302	0.7 <u>06</u> 92
Urinary N, g/d (b)	157	168	187	166	12.5	0.171	0.551	0.127
Excreted N, g/d (a+b)	3 <u>97</u> 4 9	4 <u>07</u> 13	4 <u>1628</u>	4 <u>27</u> 60	<u>19.3</u> 24.6	0. <u>270</u> 029	0. <u>497</u> 067	0. <u>988</u> 489
Milk N, g/d ¹	160	170	150	144	5.8	0.028	0.773	0.247
N use efficiency, % $^{\rm 2}$	25.7	25.1	22.3	21.9	0.78	0.008	0.572	0.886
n	4	4	4	4				

546 IR = Inclusion rate; CL = chop length; IRxCL = interaction between IR and CL; N = Nitrogen.

547 ¹ Milk N = milk protein yield / 6.25

²N use efficiency calculated as the percentage of ingested N found as milk protein N.

549

Figure captions

- **Figure 1** The effect of Short (S) or Long (L) chop length of lucerne silage on the
- distribution of particles (dry matter corrected) across the sieves of a Penn State
- 555 Particle Separator for first cut silage (experiment 1) and second cut lucerne silage
- 556 (experiment 2). Values are the means of measurements taken in each period (n=4).