

Citation:

Sinclair, LA and Johnson, DL and Wilson, S and Mackenzie, AM (2017) Added dietary sulfur and molybdenum has a greater influence on hepatic copper concentration, intake, and performance in Holstein-Friesian dairy cows offered a grass silage- rather than corn silage-based diet. Journal of Dairy Science. ISSN 0022-0302 DOI: https://doi.org/10.3168/jds.2016-12217

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### **INTERPRETIVE SUMMARY**

Added dietary sulfur and molybdenum has a greater influence on hepatic copper concentration, intake and performance in Holstein-Friesian dairy cows offered a grass silage than a corn silage based diet By Sinclair. The objectives of our study were to determine the effect of different forages on the copper status and milk performance in dairy cows when fed without or with antagonists to copper absorption. We found that, only in the high inclusion grass silage based diet did the addition of dietary sulphur and molybdenum reduce intake and milk yield and increase somatic cell count. Liver copper concentration also declined more rapidly in cows offered a grass silage diet with added sulfur and molybdenum, but blood copper levels were unaffected. We advise that the basal forage should be taken into account when supplementing copper, particularly if sulfur and molybdenum levels are high. 

26	RUNNING HEAD: COPPER METABOLISM IN DAIRY COWS
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28	Added dietary sulfur and molybdenum has a greater influence on hepatic
29	copper concentration, intake and performance in Holstein-Friesian dairy
30	cows offered a grass silage than a corn silage based diet
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46	Key words: copper, dairy cow, forage, liver,
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### ABSTRACT

51 To test the hypothesis that the metabolism of Cu in dairy cows is affected by basal forage and 52 added S and Mo, 56 dairy cows that were 35 (SE +/- 2.2) days post calving and yielding 38.9 53 kg milk/d (SE +/- 0.91) were offered one of four diets in a 2 x 2 factorial design for a 14 wk 54 period. The four diets contained approximately 20 mg Cu/kg DM, and had a corn silage to grass silage ratio of 0.75:0.25 (C) or 0.25:0.75 (G) and were either unsupplemented (-) or 55 56 supplemented (+) with an additional 2g S/kg DM and 6.5 mg Mo/kg DM. There was an 57 interaction between forage source and added S and Mo on DM intake, with cows offered G+ 58 having a 2.1 kg DM lower intake than those offered G-, but there was no effect on the corn 59 silage based diets. Mean milk yield was 38.9 kg/d, and there was an interaction between basal 60 forage and added S and Mo, with yield being decreased in cows offered G+, but increased on 61 C+. There was no effect of dietary treatment on milk composition or live weight, but body 62 condition was lower in cows fed added S and Mo irrespective of forage source. There was an 63 interaction between forage source and added S and Mo on milk somatic cell count, which was 64 higher in cows offered G+ compared to G-, but not in cows fed the corn silage based diets, 65 although all values were low (mean values of 1.75, 1.50, 1.39 and 1.67 log<sub>10</sub>/mL for C-, C+, G- and G+ respectively). Mean plasma Cu, Fe and Mn concentrations were 13.8, 41.3 and 0.25 66 67 µmol/L respectively and were not affected by dietary treatment, whereas plasma Mo was 0.2 µmol/L higher in cows receiving added S and Mo. The addition of dietary S and Mo decreased 68 69 liver Cu balance over the study period in cows fed either basal forage, but the decrease was 70 considerably greater in cows receiving the grass silage based diet. Similarly, hepatic Fe 71 decreased more in cows receiving G than C when S and Mo were included in the diet. It is 72 concluded that added S and Mo reduces hepatic Cu reserves irrespective of basal forage source, 73 but this decrease is considerably more pronounced in cows receiving grass silage than corn

silage based rations, and is associated with a decrease in intake, milk performance and increasein milk somatic cell count.

- 76 Key words: copper, corn silage, dairy cow, grass silage, liver
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# INTRODUCTION

79 It has long been recognized that Cu is an important trace element for normal health and 80 performance in dairy cattle, principally due to its requirement in approximately 300 different 81 proteins with functions ranging from efficient iron metabolism, hair pigmentation, 82 antioxidants, release of hormones and synthesis of connective tissue (Suttle, 2010). As a 83 consequence, Cu responsive disorders result in production and economic losses due to effects 84 on fertility, performance and health (NRC, 2005). Clinical signs in dairy cows can be caused 85 by a dietary deficiency of Cu, but are often are related to interactions with dietary antagonists 86 such as S and Mo, Fe and Zn that inhibit Cu absorption and/or metabolism (Suttle, 2010), with 87 S and Mo receiving the most research attention. It has been proposed that dietary sulfates 88 present in feed or water are reduced in the rumen to sulfides which then react with molybdate 89 to form thiomolybdates (Dick et al., 1975). Gould and Kendall (2011) discussed that 90 thiomolybdates may be present in the rumen as di, tri or tetrathiomolybdates, with trimolybdate 91 predominant at a ruminal pH of 6.5, whereas tetrathiomolybdate is most prevalent at lower pH 92 values. Thiomolybdates form insoluble complexes with Cu rendering it unabsorbable (Suttle, 93 1991), resulting in Cu responsive disorders. At high Mo intakes (e.g. >8 mg Mo/kg DM) and 94 very low Cu:Mo ratios (less than 1:1) thiomolybdates may also leave the rumen and be 95 absorbed (Suttle 2010), subsequently binding to Cu containing enzymes such as caeruloplasmin (Cp), impairing their function (Gould and Kendall, 2011). 96

It is recognized that the degree of thiomolybdate formation in the rumen can also be
affected by the basal forage and method of preservation (Suttle 1974; Suttle 1983; Suttle 2010),

99 although our understanding of the mechanism remains poor. For example, in grass hays, the 100 inhibitory effect of Mo on Cu absorption is less than that of S, whereas in fresh grass Cu 101 absorption is greatly affected by small additions of S and Mo, with semi-purified diets being 102 intermediate (Suttle, 1983). There is a large body of literature comparing the effect of grass 103 silage with corn silage on dairy cow intake and performance (e.g. Hart et al., 2015; Phipps et 104 al., 1995), and in general, replacing grass silage with corn silage results in an increase in DM 105 intake, milk yield and milk protein content. There is however, little information on the relative 106 effects of either of these forages on Cu metabolism in Holstein-Friesian dairy cows, despite 107 their importance in contemporary dairy cow rations. A lack of understanding of the influence 108 of S and Mo on Cu metabolism in dairy cows fed different forages may be contributing to the 109 unnecessary over-supplementation of Cu. Indeed, recent surveys of commercial trace-element 110 feeding rates in the USA and UK (e.g. Castillo et al., 2013; Sinclair and Atkins, 2013) have 111 reported that dietary Cu is frequently fed at levels well above that recommended by national 112 feed standards such as ARC (1980) or NRC (2001). Feeding Cu above nutritional requirements 113 can result in chronic Cu poisoning, whereby there is a gradual increase in hepatic Cu 114 concentrations, ultimately leading to rupture of lysomes, hepatic necrosis, haemoglobinuria, 115 methnaemoglobinaemia and rapid death (Bidewell et al., 2000). The objectives of our current 116 study were to determine the effect of level of inclusion of corn silage and grass silage fed either 117 without or with added sulfur and molybdenum on indicators of copper status, performance and 118 health in Holstein-Friesian dairy cows.

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## MATERIALS AND METHODS

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122 *Animals, Management and Treatments.* The procedures involving animals were 123 conducted in accordance with the UK Animals (Scientific Procedures) Act 1986, and were

124 approved by the Harper Adams Animal Welfare and Ethical Review Board. Fifty-six Holstein-125 Friesian dairy cows (8 primiparous and 48 multiparous) that were 35 (SE +/- 2.2) days post calving and yielding 38.9 kg/d (SE +/- 0.91) of milk were used. From calving until wk 5 of 126 127 lactation the cows were group housed and fed a diet containing (g/kg DM) grass silage 95, 128 alfalfa silage 90; corn silage 324; chopped wheat straw 20; urea treated wheat 100; soy hulls 129 80; molasses 50; soybean meal 66; rapeseed meal 64; distillers grains 64; palm kernel meal 18; 130 protected fat 14; minerals and vitamins 15. Based on recordings taken in wk 4 of lactation the 131 animals were blocked and allocated to one of four dietary treatments according to lactation 132 number (prima or multi), calving date, milk yield, milk composition, BCS (using a 1-5 scoring 133 system on a quarter point scale; Lowman et al., 1976) and live weight. Cows remained on 134 study for 14 wks.

135 Based on the mineral analysis of the forages (Table 1) and NRC (2001) values for the 136 other feeds, fours diets were formulated to contain approximately 20 mg Cu/kg DM and a corn 137 silage to grass silage ratio of 0.75:0.25 (C) or 0.25:0.75 (G: DM basis; Table 2). To evaluate 138 the effects of dietary antagonists on Cu metabolism, the diets were either unsupplemented (-) or supplemented (+) with additional S and Mo, to result in a total dietary concentration of 139 140 approximately 3.5 g S/kg DM or 7.5 mg Mo/kg DM (an increase of approximately 2 g S/kg 141 DM (+160%) and 6.5 mg Mo/kg DM (+ 500%). There were therefore 4 dietary treatments: C-142 (0.75 corn silage:0.25 grass silage (DM basis), no additional antagonists); C+ (0.75 corn 143 silage:0.25 grass silage, with additional S and Mo); G- (0.25 corn silage:0.75 grass silage, no 144 additional antagonists) and G+ (0.25 corn silage:0.75 grass silage, with additional S and Mo). 145 Additional Cu was supplied as CuSO<sub>4</sub>.5H<sub>2</sub>O, sulfur as ammonium sulfate (TG Tennants, West 146 Bromwich, UK) and molybdenum as sodium molybdate (Acros Organics, Geel, Belgium). 147 Feed grade urea was added to G- and C- to provide an equivalent amount of rumen degradable 148 N as supplied by the ammonium sulphate. The diets were supplemented with other feed

149 ingredients to support a milk production of approximately 38 kg/d according to Thomas (2004; 150 Table 2). All dietary ingredients were mixed and fed as a TMR using a forage mixer calibrated 151 to  $\pm 1$  kg, and fed through Insentec roughage intake feeders fitted with an automatic animal 152 identification and forage weighing system calibrated to  $\pm 0.1$  kg (Sinclair et al., 2005). Fresh 153 feed was offered daily at 1.05 of *ad libitum* intake with refusals collected twice weekly on a 154 Tuesday and Friday. The cows were housed in the same portion of a free stall building containing Super Comfort free stalls fitted with foam mattresses. The passageways were 155 156 scraped using automatic scrapers and the stalls bedded twice weekly with sawdust. All cows 157 had continual access to fresh bore-hole water which contained a concentration of S, Fe, Cu and 158 Mo of 19.3 mg/L, 6.5, 2.9 and 0.5  $\mu$ g/L respectively.

159 Experimental routine. Cows were milked twice daily at approximately 0530 h and 160 1530 h, with yield recorded at each milking and samples taken fortnightly at consecutive am 161 and pm milkings for subsequent composition and somatic cell count (SCC) analysis. The cows 162 were weighed and BCS recorded after the evening milking in the wk prior to allocation and 163 then fortnightly. Forage samples were collected weekly: half the sample was oven dried at 70°C 164 to constant weight, and the amount of corn silage to grass silage adjusted to achieve the desired 165 ratio. The other sample was frozen and bulked for subsequent analysis. Samples of each of the 166 four diets were collected immediately following feeding once per wk and stored at -20°C prior 167 to subsequent analysis. During wks 0, 1, 2, 4, 8 and 14 of the study blood samples were 168 collected at 1000 h via jugular venipuncture into vacutainers (Becton Dickinson Vacutainer 169 Systems, Plymouth, UK) containing, silica (for samples used to determine Cp), or lithium 170 heparin (for samples used to determine superoxide dismutase (SOD) activity) and sodium 171 heparin (for samples used to determine mineral concentrations and metabolites). During wk 0 and 14 of the study liver biopsy samples were collected from all cows through the 11<sup>th</sup> 172

intercostal space as described by Davies and Jebbett (1981), and stored at -80°C prior to
subsequent analysis.

Chemical analysis. Weekly forage and TMR samples were bulked within month and 175 176 analyzed according to AOAC (2012) for DM (934.01), CP (990.03) and starch (920.40). In addition, forage samples were analyzed for pH, ammonia-N, water soluble carbohydrates 177 178 (MAFF, 1986), and VFA based on the method of Jones and Kay (1976). The analysis of NDF 179 and ADF were conducted according to Van Soest et al. (1991) with the use of a heat-stable  $\alpha$ -180 amylase (Sigma, Gillingham, UK), and expressed exclusive of residual ash. The ME content 181 of the forages was determined by near infra-red reflectance spectroscopy (Eurofins 182 Laboratories, Wolverhampton, UK) using a system approved by the UK advisory services 183 (Offer et al., 1996). Forage and TMR minerals were extracted using the DigiPREP digestion 184 system (Qmx Laboratories, Essex, UK), and analyzed as described by Cope et al. (2009) by 185 inductively coupled plasma-mass spectrometry (ICP-MS; Thermo Fisher Scientific Inc., 186 Hemel Hempstead, UK). Serum samples were analyzed for Cp according to Henry et al. (1974) 187 and plasma samples for superoxide dismutase (SOD; Randox Laboratories, kit catalogue no. 188 SD 125), BHBA and urea (Randox Laboratories, County Antrim, UK; kit catalogue no. RB 189 1007, and UR221 respectively) using a Cobas Miras Plus autoanalyser (ABX Diagnostics, 190 Bedfordshire, UK). Plasma and liver samples were analyzed for Cu, Fe, Mn and Mo by ICP-191 MS as described by Sinclair et al., (2013). Milk samples were analyzed using a Milkoscan 192 Minor (FOSS, Warrington, UK) calibrated by the methods of AOAC (2012), and SCC was 193 determined by Eurofins Laboratories (Wolverhampton, UK).

194 *Statistical analysis.* Performance, plasma minerals and metabolites were analyzed by
195 repeated measures ANOVA as a 2 x 2 factorial design. Milk SCC was transformed to log<sub>10</sub>
196 prior to analysis. Treatment degrees of freedom were split into main effects of forage source

(corn versus grass silage), antagonist (Ant; without; (-) versus with; (+)) and their interaction
(Int) and analyzed as:

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$$Y_{ijkl} = \mu + B_i + F_j + A_k + T_l + F.A_{jk} + F.T_{jl} + A.T_{kl} + F.A.T_{jkl} + \varepsilon_{ijkl}$$
  
200 Where  $Y_{ijkl} =$  dependent variable;  $\mu =$  overall mean;  $B_i =$  fixed effect of blocks;  $F_j =$  effect of  
201 forage (j = corn or grass silage);  $A_k =$  effect of S and Mo (k = -, +);  $T_l =$  effect of time; F.A\_{jk} =  
202 interactions between forage and antagonist; F.T\_{jl} = interaction between forage and time; A.T\_{kl}  
203 = interaction between forage and time; F.A.T<sub>jkl</sub> = interaction between forage antagonist and  
204 time, and  $\varepsilon_{ijkl} =$  residual error.

205 Hepatic mineral concentration was analyzed by ANOVA as a 2 x 2 factorial design as:

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$$Y_{ijk} = \mu + B_i + F_j + A_k + F A_{jk} + \varepsilon_{ijk}$$

207 Where  $Y_{ijk}$  = dependent variable;  $\mu$  = overall mean;  $B_i$  = fixed effect of blocks;  $F_j$  = effect of 208 forage (j = corn or grass silage);  $A_k$  = effect of S and Mo (k = -, +); F.A<sub>jk</sub> = interactions between forage and antagonist; and  $\varepsilon_{ijk}$  = residual error. For hepatic mineral concentrations the 209 210 concentration during wk 0 was used where appropriate as a covariate to determine the final and 211 rate of mineral deposition or mobilization. All statistical analysis was conducted using Genstat 212 version 17.1 (VSN Int. Ltd., Oxford, UK) and is presented as means with standard error of the 213 mean (SEM); P < 0.05 was used as the significant threshold and a trend was considered when 214 *P* < 0.1.

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### RESULTS

217 *Diet Analysis, Intake and Animal Performance.* Compared to the corn silage, the grass 218 silage contained 85 g/kg less DM, and was 82 g/kg DM higher in CP and 0.4 MJ/kg DM higher 219 in ME (Table 1). The two forages had a similar fiber content, but the grass silage was 43.2 g/kg 220 DM higher in lactic acid than the corn silage. Compared to the corn silage, the mean content of Ca, P, Mg and S was 5.0, 1.0, 0.2 and 2.2 g/kg DM higher respectively, and Cu, Mo, Fe and
Zn 3.3, 0.84, 94 and 14.2 mg/kg DM higher respectively in the grass silage.

223 The DM content of the corn based diets (C- and C+) was 47 g/kg higher than the grass 224 silage based diets (G- and G+), whereas CP was on average 11 g/kg DM higher in the grass 225 than the corn silage based diets (Table 2). The content of NDF was higher in the corn than the 226 grass silage based diets, but ADF concentration was similar across all four diets, averaging 225 227 g/kg DM. All four diets had a similar P and Mg concentration, but the grass silage based diets 228 (G- and G+) contained approximately 2 g/kg DM more Ca. The mean concentration of Cu was 229 20 mg/kg DM, and the two diets with added antagonists (C+ and G+) had concentrations of S 230 and Mo of 3.3 g/kg DM and 7.8 mg/kg DM respectively, which were close (P > 0.05) to the 231 predicted values of 3.5 g/kg DM and 7.5 mg/kg DM respectively. In contrast, the two diets 232 with no added S and Mo (Corn- and Grass-) had low concentrations of S and Mo at 1.3 g/kg 233 DM and 1.3 mg/kg DM respectively that were also close (P > 0.05) to predicted.

234 Cows offered the corn silage based diets had a daily DM intake that was 2.2 kg/d higher 235 (P < 0.001) than those offered the grass silage based diets (Table 3), an effect that was evident 236 from wk 1 of the study (Fig 1). There was an interaction (P < 0.05) between forage source and 237 Cu antagonists; adding S and Mo reduced DM intake by 2.1 kg/d in cows fed the grass silage 238 but not the corn silage based diet. We also found an interaction between forage source and 239 antagonist on Cu intake, which was lowest (P < 0.05) in cows fed G+ compared to the other 3 240 treatments. There was an interaction (P < 0.05) between forage source and Cu antagonists on 241 milk yield, with yield decreasing with the addition of S and Mo in cows fed the grass silage 242 based diet, but increasing in those offered the corn silage based diet. In contrast, there was no 243 effect (P > 0.05) of dietary treatment on milk fat, protein or lactose content or daily fat yield, 244 but we found that daily milk protein yield was 0.05 kg/d higher (P < 0.05) in cows fed the corn 245 silage based diet. We found no effect (P > 0.05) of dietary treatment on live weight or daily

246 live weight change, but there was an effect of antagonist on BCS and BCS change (P < 0.05), 247 with cows fed added S and Mo (C+ and G+) having a lower score and gained less BCS over 248 the study period than those not supplemented with S and Mo (C- and G-; Fig 2). There was an 249 interaction (P < 0.05) between forage source and Cu antagonists on milk SCC count, with the 250 addition of S and Mo increasing SCC in cows fed the grass but not the corn silage based diet.

251 Plasma Mineral Profile, Cu Mediated Enzymes and Metabolites. We found no effect (P > 0.05) of dietary treatment on plasma Cu concentration, with a mean value of 13.7  $\mu$ mol/L 252 253 (Table 4). There was an effect of time on plasma Cu, with the concentration increasing in the 254 first wk of the study, and then fluctuating in subsequent wks (Fig 3). We also found an effect 255 (P < 0.001) of dietary treatment on mean plasma Mo concentrations, which were higher in 256 cows fed added S and Mo, but there was no effect (P > 0.05) of basal forage. There was no 257 effect (P > 0.05) of dietary treatment on plasma Fe or Mn concentrations. Serum Cp 258 concentrations were higher (P < 0.01) in cows fed the grass silage based diets or with added S 259 and Mo (P < 0.05). In contrast, we found no effect of dietary treatment on blood Cp:Cu ratio, 260 although there was a trend (P < 0.1) for a lower ratio in cows fed the corn silage based diets, or in animals receiving added S and Mo. There was no effect (P > 0.05) of dietary treatment 261 on plasma SOD, BHBA or BUN concentrations, with mean values of 2918 U/gHb, 0.43 262 mmol/L and 5.44 mmol/L respectively. 263

264 Hepatic Mineral Concentration. There was no difference between dietary treatments 265 (P > 0.05) in initial hepatic Cu concentration, which averaged 443 mg/kg DM (Table 5). We did find an effect of forage source on final Cu concentration, which was higher (P < 0.05) in 266 267 cows fed the corn compared to the grass silage based diets. There was also an effect of Cu 268 antagonists on final hepatic Cu concentration, which was 142 mg/kg DM lower (P < 0.01) in cows fed added S and Mo. There was a trend (P < 0.1) for an interaction between forage source 269 270 and Cu antagonists on the rate of change in hepatic Cu concentration, with a decrease of 61

271 mg/kg DM over the 14 wk study period in cows fed added S and Mo in combination with grass 272 silage (G+), but an increase of 11 mg/kg DM in cows offered the corn silage based diet (C+). 273 We found no difference between treatments in initial hepatic Mo concentration (P >274 0.05), whereas final Mo concentration was higher (P < 0.05) in cows fed added S and Mo (C+ 275 and G+). Initial hepatic Fe concentration did not differ between treatments (P > 0.05), whereas 276 final concentration was lower (P < 0.01) in cows fed added S and Mo, and there was a trend 277 (P < 0.1) for final hepatic Fe concentration to be higher in cows offered the corn compared to 278 the grass silage based diet. The addition of S and Mo resulted in a net decrease in hepatic Fe 279 concentration over the study period of 19 mg/kg DM compared to an increase in cows that 280 were not supplemented with S and Mo of 50 mg/kg DM, although most of this difference could 281 be attributed to cows fed the grass silage based ration with added S and Mo (G+) decreasing in 282 hepatic Fe concentration (P < 0.1) compared to an increase in cows fed any of the other dietary treatments. Finally, we found no effect (P < 0.05) of dietary treatment on hepatic Mn 283 284 concentrations, although cows fed the grass silage with Cu antagonists (G+) tended (P < 0.1) 285 to decrease by the greatest amount.

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## DISCUSSION

288 Intake and Performance. Our study is the first to determine Cu status and metabolism 289 in high yielding dairy cows when fed corn or grass silage based rations at different S and Mo 290 concentrations. Corn silage is generally regarded as having a lower Cu concentration than grass 291 silage (NRC, 2001), but we supplemented the diets to ensure that levels were similar across all 292 treatments, averaging 20.0 mg Cu/kg DM. The dietary level of 20 mg Cu /kg DM was lower 293 than the mean value of 27.9 mg/kg DM/d that was reported in the diet of early lactation cows 294 in the UK (Sinclair and Atkins, 2013), but similar to the 18 mg/kg DM reported on 39 295 Californian dairy units by Castillo et al., (2013). Additionally, we added S and Mo at a rate to

ensure that the supplemented diets (C+ and G+) had similar concentrations which would be 296 297 expected to substantially reduce Cu absorption and subsequent metabolism. Differences in 298 dietary S and Mo concentration between diets within the same level of antagonist was small (P 299 > 0.05), and therefore the main effect was the difference between the unsupplemented and 300 supplemented diets. Using the equations of Suttle and McLauchlan (1976), we predicted that 301 the C- and G- diets would result in an apparent digestibility co-efficient of Cu of approximately 302 0.054, whereas the C+ and G+ diets would be two-thirds lower at approximately 0.018. As a 303 consequence, we predicted that animals receiving C- or G- had a similar Cu supply but were 304 over supplied by approximately 220 mg Cu/d whereas those receiving C+ or G+ were 305 undersupplied by approximately 200 mg Cu/d. However, the use of the current equations did 306 not predict any interaction between forage source and antagonist on Cu status or performance.

307 Similar to other studies that have investigated the effect of replacing grass silage with 308 corn silage (Phipps et al., 1995; Hart et al., 2015), we found that DM intake was increased at 309 the higher corn inclusion rate, although it is accepted that the change in forage composition 310 from the pre-study diet was greater for cows on G than C diets. However, we also found an 311 interaction between forage inclusion level and Cu antagonists on intake, with added S and Mo 312 having little effect in cows fed the corn silage based diet, but reduced intake by 2.1 kg DM/d 313 in those receiving the grass silage based diet. Our diets were supplemented with both S and 314 Mo, and it is therefore not possible to determine the effects of each element independently. 315 Some authors have reported a decrease in DMI in cattle when dietary S exceeded 2 g/kg DM 316 (Spears et al., 2011), although others have reported little effect of dietary S concentration up to 317 6 g/kg DM (Richter et al., 2012). Under acidic ruminal conditions most of the S would be 318 present as H<sub>2</sub>S, which may be eructated and absorbed by the lungs or absorbed across the rumen 319 epithelium (Bray and Till, 1975; Drewnoski et al., 2012). High circulating concentrations of 320 H<sub>2</sub>S can have neurological effects including polioenchalomalacia that is associated with a

321 reduced intake (Gould, 1998). The large role that ruminal pH plays in the form of sulphide 322 present in the rumen has been suggested as a possible explanation for the differences observed 323 in sulfur tolerance between concentrate and roughage fed cattle (Drewnoski et al., 2012), and 324 could explain the reduced DMI of cows offered G+ in our study. However, we did not monitor 325 ruminal H<sub>2</sub>S or pH levels, and the influence of level of inclusion of corn and grass silage on 326 ruminal pH is difficult to predict as it is dependent on a number of factors including initial 327 forage pH, buffering capacity of the diet, forage particle length, and supplementary feed level, 328 composition, and degree of processing (Krause and Oetzel, 2006).

329 Molybdenum interacts with S in the rumen resulting in the formation of various 330 isomers of thiomolybdate, a reaction which is reversible and pH dependent, with the 331 formation of tetra-thiomolybdate being favored at lower ruminal pH values (Gould and 332 Kendall, 2011). Indeed, the dietary addition of Mo has been proposed as a potential sink for 333 H<sub>2</sub>S in the rumen (Kessler et al., 2012), potentially reducing the negative effects of excess 334 dietary S on intake, although this approach has not been supported by recent studies with beef 335 animals (Kessler et al., 2012). An alternative hypothesis for the effect of added S and Mo on 336 intake may be related to the absorption of tetra-thiomolybdates as these can have a direct 337 effect on Cu containing enzymes such as peptidylglycine  $\alpha$ -amidating monooxygenase which 338 exerts an influence on the appetite-regulating hormones cholecystokinin and gastrin (Suttle, 339 2010), although studies in this area in ruminants are scarce. Ruminal absorption of tetra-340 thiomolybdates is increased at lower ruminal pH values, and it is possible that differences in 341 the ruminal pH in cows fed the different forages affected uptake. The conditions under which 342 thiomoybdates are absorbed is, however, a controversial subject area, and it was proposed by 343 Suttle et al., (2010) that absorption was unlikely unless dietary Cu:Mo rations were below 344 1:1, well below the 2.5:1 in our C+ and G+ diets. It is also possible that the added Mo 345 resulted in molybdenosis, however, no characteristic signs such as scouring were noted and

346 dietary values were well below that reported in other studies that have also reported no signs347 (Raisbeck et al. 2006).

348 Studies that have fed varying levels of Cu to dairy cows in the absence of high levels 349 of dietary antagonists have reported little effect on DM intake (see review of Sinclair and 350 Mackenzie 2013), and it therefore appears unlikely that a lower tissue supply of Cu per se was 351 responsible for the differences in DM intake reported here. It is of interest to note that the 352 inclusion of S and Mo reduced BCS in the cows in our study, irrespective of basal forage level. 353 This effect may be attributed to different mechanisms for each of the forage treatments, as milk 354 yield was higher in cows fed C+ compared to C-, whereas intake was lower in cows fed G+ 355 compared to G-.

356 The interaction between basal forage source and Cu antagonists on milk SCC in our 357 study is difficult to explain, although all values were low. The role of Cu on milk SCC has been 358 demonstrated in dairy cattle in some but not all studies. For example, increasing dietary Cu 359 concentration from a sub-optimal level of 6.5 mg/kg DM to 26.5 mg/kg DM was shown to 360 reduce the peak increase in milk SCC following a challenge with E. Coli which was attributed 361 to a greater ability of neutrophils to kill invading bacteria, although the duration of the infection 362 was unaffected (Scaletti et al., 2003). In contrast, dietary Cu concentration was not shown to have an effect on milk SCC concentration following a challenge with E. Coli in the studies of 363 364 Scaletti and Harmon (2012), or when different levels of dietary Cu were fed (Chase et al., 365 2000). In our study, cows receiving G+ were in negative Cu balance as evidenced by the 366 depletion of hepatic Cu reserves, whereas all other treatments were in positive balance. It is 367 therefore possible that this lower Cu status contributed to the increased milk SCC, although 368 other indicators of Cu status such as plasma Cu and plasma Cu:Cp were unaffected by dietary treatment. The lower DM intake that we observed in cows receiving G+ may also have 369 370 contributed to a greater metabolic stress and indirectly increased milk SCC.

372 Plasma Mineral Profile, Cu Mediated Enzymes and Metabolites. We found that plasma Cu 373 concentrations were unaffected by dietary treatment, with all values being above the 9 mmol/L 374 considered to be adequate (Laven and Livesey, 2005). Our finding is consistent with others that 375 have supplemented Cu at different levels (Chase et al., 2000), with different levels of dietary S 376 and Mo (Sinclair et al., 2013), or with different dietary sources of Cu (Scaletti and Harmon, 377 2012; Sinclair et al., 2013). In a meta-analysis of the relationship between dietary concentration 378 of Cu, S and Mo and plasma Cu in growing cattle, Dias et al., (2013) concluded that any 379 prediction equation would be limited, and that it is only when animals have either very low or 380 high hepatic Cu reserves that plasma values can be usefully employed as an indicator of Cu 381 status (Laven and Livesey, 2005). The plasma Cu:Cp ratios reported in our study were 382 generally low, and unaffected by dietary treatment. Similarly, we found that plasma SOD, a Cu 383 containing enzyme involved in the defense against free radicals (Suttle, 2010), was unaffected 384 by dietary treatment. Our findings therefore support Suttle (2010) who suggested that the 385 dietary ratio of Cu:Mo needed to be close to 1:1 before there is a risk of thiomolybdates causing 386 a systemic impairment of Cu containing enzymes.

387 Hepatic Mineral Concentration. One of the first biochemical changes observed under 388 Cu deprivation is a decrease in hepatic concentration (Suttle, 2010), as the liver is generally 389 regarded as the principal storage organ for Cu (Laven and Livesey, 2005). In our study initial 390 hepatic Cu levels were high and variable at  $443 \pm 29.2$  (SE) mg/kg DM, although most (68%) 391 animals were below the upper limit of 510 mg/kg DM suggested to pose a risk of toxicity 392 (Livesey et al., 2002). The initial mean hepatic Cu concentration that we found was also lower 393 than that reported by Kendall et al., (2015), where almost 40% of cull dairy cows in the UK 394 were reported to have a concentration above 500 mg Cu/kg DM. As we anticipated, there was 395 a significant reduction in hepatic Cu concentration following the addition of dietary S and Mo,

396 but the greater reduction in cows fed a grass silage compared to the corn silage based diet was 397 unexpected, although the difference failed to reach full statistical significance. Suttle (2013) discussed that changes in hepatic Cu concentration are an exponential function of initial hepatic 398 399 Cu concentration, most probably due to a greater rate of biliary excretion at higher liver 400 concentrations. We therefore log<sub>e</sub> transformed and re-analyzed the initial and final hepatic Cu 401 concentrations to more accurately determine the influence of diet on hepatic Cu reserves. 402 Similar to the untransformed data, we found no difference (P > 0.1) between treatments in 403 initial liver Cu concentration, but we did now find an interaction (P < 0.05) between forage 404 source and Cu antagonist on daily liver Cu balance (log<sub>e</sub> final – log<sub>e</sub> initial), confirming that 405 high dietary concentrations of S and Mo have a greater effect on Cu metabolism in cows 406 receiving a grass silage than a corn silage based diet.

407 The influence of forage source on the absorption of Cu is well demonstrated in sheep 408 (e.g. Suttle 1983; Suttle 2010), and in the absence of high Mo concentrations, the absorption 409 coefficient of Cu was reported to be 0.014 in grazed grass, 0.049 in grass silage, 0.073 in hay 410 and 0.128 in leafy brassicas. This is however, the first study to report a substantial difference 411 in Cu status in dairy cows fed corn or grass silage based rations, but only when S and Mo 412 concentrations were high. Dietary Fe may interact with added S reducing hepatic Cu 413 concentration (Suttle, 2010). However, the low dietary concentration of Fe in all of our diets 414 compared to that reported for typical dairy cow rations in the UK (Sinclair and Atkins, 2013) 415 or California (Castillo et al., 2013), in combination with the similarity in dietary Fe and S 416 concentration between C+ and G+, does not support Fe as having a major influence in our 417 study. Consideration should also be given to the lower DM intake of cows receiving G+ which 418 resulted in a lower Cu intake of 49 mg/d than G-. Nevertheless, at the rate of decline in hepatic 419 Cu concentration in cows receiving G+, concentrations would reduce and eventually approach 420 the 25 mg Cu/kg DM threshold considered to deficient (Laven and Livesey, 2005). In contrast, in cows fed C- or G-, feeding 20 mg Cu/kg DM would result in a rapid increase in hepatic Cu
concentration, whereas those receiving C+ would be relatively unchanged. Given such large
differences in Cu status when fed the same dietary level, we recommend that forage source as
well as dietary S and Mo concentration should be taken into account when supplementing dairy
cows with Cu.

426 Similar to our previous study (Sinclair et al. 2013), liver Mo concentrations were little 427 affected by dietary treatment, despite a 6.5 mg/kg DM difference in dietary concentration 428 between (-) and (+) treatments, and we can conclude that the liver does not appear to be either 429 a major store or a sensitive indicator of Mo status. Ferritin is the main storage form of Fe in the 430 body, and is particularly concentrated in the liver where concentrations of between 100 to 1000 431 mg Fe/kg DM are considered to be normal in cattle (Suttle, 2010). Hepatic Fe concentrations 432 at the beginning and end of our study were within this range, but similar to Cu, hepatic Fe 433 concentrations were negatively affected by the addition of S and Mo, particularly in the grass 434 silage based diet. In contrast, Phillippo et al., (1987) reported in growing calves fed a barley-435 straw based diet that an additional 5 mg Mo/kg DM increased liver Fe concentrations, which 436 was associated with a decrease in plasma Fe concentrations.

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### CONCLUSIONS

We found that the addition of S and Mo had no effect on DM intake or milk yield in cows fed a corn silage based ration, but were reduced and milk SCC increased when a grass silage based diet was fed. In the absence of additional S and Mo, a diet containing 20 mg Cu/kg DM whether based on grass or corn silage, contains well in excess of requirements as evidenced by the net increase in hepatic Cu concentration. In contrast, in the presence of high levels of S and Mo, feeding 20 mg Cu/kg DM will result in a rapid depletion of hepatic Cu concentrations in cows fed grass silage, but not corn silage based diets. Within the limits of this study we also

446	found that there was little effect of added Cu antagonists on plasma Cu or indicators of plasma
447	Cu enzyme activity, even at the high levels of S and Mo, and suggest that use of these
448	parameters to predict Cu status is limited. Reasons for the differences in Cu metabolism in
449	cows when fed grass or corn silage based rations is unclear and require further investigation,
450	but our results highlight the importance of taking account of forage source when formulating
451	diets for dairy cows, particularly when dietary S and Mo levels are high.
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453	ACKNOWLEDGMENTS
454	The authors are grateful to AHDB Dairy for funding the work. The technical support of G.
455	Vince and the assistance of N. Atkins at Harper Adams University are also gratefully
456	acknowledged.
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Table 1. Chemical composition of com and grass shage								
	Corn silage	Grass silage						
DM, g/kg	341	256						
CP, g/kg DM	75	157						
Ash, g/kg DM	46							
Ammonia-N, g/kg total N	9.03	8.39						
рН	3.6	3.9						
ME, MJ/kg DM	10.8	11.2						
Water soluble carbohydrate, g/kg DM	26.2	68.8						
NDF, g/kg DM	449	439						
ADF, g/kg DM	229	246						
Volatile fatty acids								
Lactic, g/kg DM	62.1	105.3						
Acetic, g/kg DM	16.1	22.6						
Propionic, g/kg DM	0.92	1.06						
Butyric, g/kg DM	<0.6	<0.6						
Ethanol, g/kg DM	1.84	28.1						
Minerals								
Ca, g/kg DM	2.3	7.3						
P, g/kg DM	2.3	3.3						
Mg, g/kg DM	1.5	1.7						
S, g/kg DM	0.9	3.1						
Cu, mg/kg DM	4.7	8.0						
Mo, mg/kg DM	0.59	1.43						
Fe, mg/kg DM	65.0	159.4						
Zn, mg/kg DM	23.6	37.8						
Mn, mg/kg DM	15.6	34.8						

 Table 1. Chemical composition of corn and grass silage

	C-	C+	G-	G+
Ingredient, g/kg DM				
Grass silage	133	134	398	399
Corn silage	400	401	133	134
Urea-treated wheat	111	111	167	167
Soy hulls	89	89	89	89
Rapeseed meal	58	58	31	31
Soybean meal	96	96	31	31
Distillers dark grains with solubles	58	58	31	31
Sopralin <sup>1</sup>			58	58
Molasses	33	33	33	33
Protected fat	13	13	20	20
Urea	2		2	
Mins/vits <sup>2</sup>	7	7	7	7
Total	1000	1000	1000	1000
Chemical analysis				
DM, g/kg	404	421	364	368
Ash, g/kg DM	71	71	92	93
CP, g/kg DM	181	185	193	194
NDF, g/kg DM	407	403	381	387
ADF, g/kg DM	222	224	228	224
Ca, g/kg DM	5.40	5.45	7.84	7.49
P, g/kg DM	3.57	3.82	3.96	3.69
Mg, g/kg DM	2.72	2.84	2.92	2.79
S, g/kg DM	1.20	3.15	1.32	3.45
Cu, mg/kg DM	19.9	19.5	20.7	20.5
Mo, mg/kg DM	1.17	7.94	1.48	7.70
Fe, mg/kg DM	183	226	287	252
Zn, mg/kg DM	49.2	46.3	51.8	48.8
Mn, mg/kg DM	61	68	70	60

**Table 2**. Diet composition and chemical analysis of diets high in corn silage (C) or grass silage (G) fed without (-) or with (+) added S and Mo.

<sup>1</sup>Formaldehyde treated soybean meal, Frank Wright Trouw, Ashbourne, UK <sup>2</sup>Mineral/vitamin premix (Rumenco, Staffordshire, UK). Major minerals (g/kg): Ca 240, P 80, Mg 120; Trace minerals (mg/kg): Cu 0, Zn 7,000, Mn 2,000, I 400, Co 80, and Se 50; vitamins (mg/kg) were: retinol 105, cholecalciferol 1.75, and all *rac*  $\alpha$ -tocopherol acetate 5,000. <sup>3</sup>SEM for differences between dietary concentrations (n = 8 per treatment) for S and Mo was 0.11 and 0.29 respectively.

C+ and G+ diets also received additional ammonium sulfate and sodium molybdate dihydrate.

		Diets				Significance, <i>P</i> -value <sup>1</sup>			
	C-	C+	G-	G+	SEM	F	А	Int	
Intake									
DM, kg/d	23.5	24.0	22.6	20.5	0.48	< 0.001	0.111	0.012	
Cu, mg/d	467	466	467	418	9.6	0.022	0.007	0.015	
Mo, mg/d	27.4	190.2	33.5	157.5	2.69	< 0.001	< 0.001	< 0.001	
S, g/d	28.1	74.9	29.9	70.4	1.25	0.302	< 0.001	0.013	
Fe, g/d	4.30	5.42	6.43	5.17	0.121	< 0.001	0.564	< 0.001	
Milk yield, kg/d	38.1	40.6	38.9	37.9	0.77	0.225	0.373	0.034	
Fat, g/kg	37.8	36.6	38.2	37.4	1.37	0.656	0.475	0.889	
Protein, g/kg	32.5	32.6	31.6	32.5	0.80	0.173	0.901	0.646	
Lactose, g/kg	46.5	46.4	46.7	46.3	0.32	0.975	0.328	0.680	
Fat yield, kg/d	1.43	1.43	1.47	1.39	0.059	0.944	0.484	0.468	
Protein yield, kg/d	1.23	1.30	1.22	1.21	0.024	0.049	0.242	0.142	
Lactose yield, kg/d	1.77	1.93	1.80	1.73	0.063	0.185	0.434	0.060	
Lwt, kg	651	653	646	639	7.9	0.237	0.818	0.587	
Lwt change, kg/d	0.43	0.30	0.20	0.25	0.131	0.309	0.738	0.518	
Condition score	2.49	2.35	2.49	2.31	0.047	0.803	0.001	0.744	
Condition score change	0.35	0.13	0.27	0.09	0.081	0.470	0.019	0.801	
Milk SCC (log <sub>10</sub> /mL)	1.72	1.50	1.39	1.67	0.086	0.381	0.714	0.017	

**Table 3.** Intake and performance of early lactation dairy cows fed diets high in corn silage (C) or grass (G) silage fed without (-) or with (+) added S and Mo.

 $^{1}$ F= main effect of forage source, A = main effect of antagonists, Int = interaction between forage and antagonists

	Diets					Significance, P-		-value <sup>1</sup>	
	C-	C+	G-	G+	SEM	F	А	Int	
Plasma Cu, µmol/L	13.3	13.7	14.3	13.7	0.51	0.340	0.889	0.332	
Plasma Mo, µmol/L	0.33	0.50	0.27	0.50	0.029	0.271	< 0.001	0.375	
Plasma Fe, µmol/L	43.2	40.5	40.7	40.9	1.61	0.519	0.446	0.384	
Plasma Mn, µmol/L	0.25	0.24	0.27	0.25	0.010	0.124	0.239	0.740	
Caeruloplasmin, mg/dL	17.9	15.9	20.3	18.1	0.79	0.006	0.010	0.909	
Caeruloplasmin:Cu	1.37	1.22	1.41	1.36	0.057	0.096	0.090	0.377	
SOD <sup>2</sup> U/gHb	2960	2841	2954	2915	89.8	0.710	0.387	0.657	
BHBA, mmol/L	0.42	0.38	0.44	0.48	0.048	0.210	0.963	0.406	
BUN, mmol/L	5.22	5.44	5.70	5.39	0.189	0.265	0.802	0.172	

**Table 4.** Plasma mineral concentration and metabolites and serum caeruloplasmin in early lactation dairy cows fed diets high in corn silage (C) or grass silage (G) fed without (-) or with (+) added S and Mo. Blood samples were collected during wks 0, 1, 2, 4, 8 and 14 of the study.

<sup>1</sup>F= main effect of forage source, A = main effect of antagonists, Int = interaction between forage and antagonists. There was a time x treatment effect on plasma Mo (P < 0.05), which increased with time in animals receiving C+ and G+ compared to C- and G-<sup>2</sup>Superoxide dismutase

	Diets					Signifi	-value <sup>1</sup>	
	C-	C+	G-	G+	SEM	F	А	Int
Initial Cu, mg/kg DM	522	426	407	418	47.0	0.201	0.372	0.262
Final Cu, mg/kg DM	587	437	490	357	41.0	0.038	0.002	0.837
Cu change, mg/kg DM per day	0.66	0.11	0.84	-0.62	0.253	0.275	0.001	0.078
Initial Mo, mg/kg DM	3.90	3.50	3.39	4.12	0.356	0.878	0.636	0.120
Final Mo, mg/kg DM	3.92	4.19	3.79	4.71	0.221	0.377	0.011	0.149
Mo change, µg/kg DM per day	0.20	6.94	4.08	6.02	4.622	0.750	0.356	0.600
Initial Fe, mg/kg DM	378	313	288	295	36.6	0.150	0.422	0.334
Final Fe, mg/kg DM	411	319	352	253	31.8	0.057	0.005	0.908
Fe change, µg/kg DM per day	336	61	653	-429	222.4	0.690	0.005	0.079
Initial Mn, mg/kg DM	10.20	9.60	9.15	10.41	0.565	0.839	0.560	0.109
Final Mn, mg/kg DM	10.18	10.38	9.96	9.84	0.305	0.223	0.895	0.610
Mn change, µg/kg DM per day	-0.20	7.96	8.26	-5.82	5.704	0.641	0.605	0.060

**Table 5**. Liver mineral concentrations in early lactation dairy cows fed diets high in corn silage (C) or grass silage (G) fed without (-) or with (+) added S and Mo.

 ${}^{1}F$ = main effect of forage source, A = main effect of antagonists, Int = interaction between forage and antagonists



**Figure 1.** Weekly DM intake in early lactation dairy cows fed diets high in corn silage and fed without ( $\blacksquare$ ) or with ( $\Box$ ) added S and Mo, or diets high in grass silage fed without ( $\blacktriangle$ ) or with ( $\triangle$ ) added S and Mo. Pooled SEM = 0.72. Forage, *P* < 0.001; Forage x Ant, *P* = 0.012; Time, *P* < 0.001; Forage x time, *P* = 0.003.



**Figure 2.** Fortnightly BCS in early lactation dairy cows fed diets high in corn silage and fed without ( $\blacksquare$ ) or with ( $\square$ ) added S and Mo, or diets high in grass silage fed without ( $\blacktriangle$ ) or with ( $\triangle$ ) added S and Mo. Pooled SEM = 0.067. Ant, *P* < 0.001; Time, *P* < 0.001; Time x ant, *P* = 0.077.



**Figure 3.** Plasma Cu concentrations in early lactation dairy cows fed diets high in corn silage and fed without ( $\blacksquare$ ) or with ( $\square$ ) added S and Mo, or diets high in grass silage fed without ( $\blacktriangle$ ) or with ( $\triangle$ ) added S and Mo. Pooled SEM = 0.87. Time, *P* < 0.001.