

# More salt, please: global patterns, responses, and impacts of foliar sodium in grasslands

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| 52       | 30<br>37 | sites and collected data used in these analyses; EB, ES, and EL also coordinated the experiment. EB wrote the first draft of the manuscript, and all other authors contributed substantially to revisions. See |
| 53       | 38       | Table SI 11 for details.   |
| 54<br>55 | 39       | <b>Data information:</b> All data contributors are listed in SI Table 12.  |
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| 40 | ABSTRACT   |  |  |
|----|--|--|--|
| 41 | Sodium is unique among abundant elemental nutrients, because most plant species do not require it for      |  |  |
| 42 | growth or development, whereas animals physiologically require sodium. Foliar sodium influences            |  |  |
| 43 | consumption rates by animals and can structure herbivores across landscapes. We quantified foliar          |  |  |
| 44 | sodium in 201 locally-abundant, herbaceous species representing 32 families and, at 26 sites on four       |  |  |
| 45 | continents, experimentally manipulated vertebrate herbivores and elemental nutrients to determine          |  |  |
| 46 | their effect on foliar sodium. Foliar sodium varied taxonomically and geographically, spanning five        |  |  |
| 47 | orders of magnitude. Site-level foliar sodium increased most strongly with site aridity and soil sodium;   |  |  |
| 48 | nutrient addition weakened the relationship between aridity and mean foliar sodium. Within sites, high     |  |  |
| 49 | sodium plants declined in abundance with fertilization, whereas low sodium plants increased. Herbivory     |  |  |
| 50 | provided an explanation: herbivores selectively reduced high nutrient, high sodium plants. Thus,           |  |  |
| 51 | interactions among climate, nutrients, and the resulting nutritional value for herbivores determine foliar |  |  |
| 52 | sodium biogeography in herbaceous-dominated systems.   |  |  |
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| 54 | INTRODUCTION   |  |  |
| 55 | Sodium is an essential nutrient for herbivores (Michell 1989; Snell-Rood et al. 2014) that can determine   |  |  |
| 56 | animal foraging preferences and movement patterns in space and time (McNaughton 1988; Prather et           |  |  |
| 57 | al. 2018). In contrast, sodium is not used for physiological function in most plants, and at high          |  |  |
| 58 | concentrations sodium can be toxic for plants (Mäser et al. 2002; Pardo & Quintero 2002; Marschner         |  |  |
| 59 | 2011; Maathuis 2014). Because of this key difference in the mineral nutrition of herbivores and the        |  |  |

60 plants they eat, herbivores must use natural salt licks and seek out and efficiently use the sodium

- 61 present in plants to meet physiological demands for sodium (Michell 1989). In spite of the essential role
- 62 of plant sodium content for wild herbivores (Seastedt & D. A. Crossley 1981), there is little
- 63 understanding of the relative importance of the many factors that may control foliar sodium in plants.
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| 64 | For example, abiotic factors including soil sodium content, soil fertility, or climate may determine                         |
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| 65 | sodium availability, whereas biotic constraints such as plant species phylogeny and lifeform or                              |
| 66 | palatability to herbivores may determine the capacity for sodium exclusion and whole tissue losses that                      |
| 67 | may occur with preferential herbivory. Further, these factors may interact and operate globally or                           |
| 68 | regionally to influence foliar sodium, and context may determine whether foliar sodium is likely to                          |
| 69 | interact with herbivory to determine the composition of plant communities in future environments.                            |
| 70 | Plants access sodium through leaf uptake from atmospheric deposition (Benes <i>et al.</i> 1996) or root                      |
| 71 | uptake from soil water (Epstein 1973). Because of the similarity of sodium to the potassium ion that is                      |
| 72 | physiologically critical for plants, cation transporters of roots will transport both sodium and potassium                   |
| 73 | across cell membranes (Pardo & Quintero 2002; Maathuis 2014). Although a relatively small group of                           |
| 74 | plants – mostly C <sub>4</sub> grasses – requires sodium (Brownell & Crossland 1972; Furumoto <i>et al.</i> 2011), the       |
| 75 | sodium cation is present in the foliage of many species and can be used for a variety of critical plant                      |
| 76 | functions, including stomatal opening and closing, particularly when potassium is in short supply                            |
| 77 | (Subbarao et al. 2003). However, terrestrial sodium is geographically variable (Kaspari et al. 2008;                         |
| 78 | Kaspari <i>et al.</i> 2009; Wicke <i>et al.</i> 2011; Vet <i>et al.</i> 2014; Doughty <i>et al.</i> 2016) because of mineral |
| 79 | acquisition from sources such as ocean spray, terrestrial salinization, or road salting practices                            |
| 80 | (Ramakrishna & Viraraghavan 2005; Vet <i>et al.</i> 2014), urine (Kaspari <i>et al.</i> 2017), loss from leaching            |
| 81 | (Vitousek & Sanford 1986), and climatic influences, particularly aridity (Raheja 1966). In spite of these                    |
| 82 | general associations, it remains unclear whether foliar sodium varies predictably among plant taxonomic                      |
| 83 | lineages or biogeographically with e.g., distance to coast or site aridity and whether there are site or                     |
| 84 | plant species characteristics that effectively predict the foliar sodium content of the most abundant                        |
| 85 | plants.  |
| 86 | Although plant sodium is often assumed to simply track soil sodium supply, at biogeographic scales, a                        |

Although plant sodium is often assumed to simply track soil sodium supply, at biogeographic scales, a
 growing body of evidence suggests that plant sodium content may not be determined solely via soil

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88 sodium supply. Like other soil cations, sodium uptake by plants can be reduced in high pH soils (Tyler & 89 Olsson 2001; Bolan & Brennan 2011), and aridity can lead to increased soil pH (Slessarev et al. 2016), 90 suggesting that aridity may either increase foliar sodium via increased soil sodium or reduce it via 91 increased soil pH. Evidence also is accumulating that the supply of macronutrients such as nitrogen can 92 reduce the availability of mineral cations to plants (Lucas et al. 2011). Thus, anthropogenic activities that 93 are altering soil pH or increasing macronutrient supply to ecosystems (Franklin et al. 2016) may 94 interactively alter the sodium content of foliage and quality of foliage for herbivores (Kaspari et al. 95 2017). Furthermore, herbivores may themselves alter the sodium concentration in plant tissue either by 96 promoting the availability of sodium through recycling (McNaughton et al. 1997; Doughty et al. 2016), 97 by promoting saline soil conditions (McLaren & Jefferies 2004), or selectively consuming plant species 98 with elevated salt levels in their foliage (Seastedt & D. A. Crossley 1981; Welti et al. 2019). These 99 conditions may, alternatively, promote plant species with relatively high foliar sodium that have traits, 100 such rapid regrowth, basal meristems, or use of sodium to modify osmotic potential under drought, that 101 are beneficial under both saline soil conditions and high grazing intensity (Coughenour 1985; Veldhuis et 102 al. 2014; Griffith et al. 2017). 103 Here, we use existing and experimentally-created environmental gradients to address the following 104 questions (1) Patterns of foliar sodium: Which site (10<sup>4</sup> m<sup>2</sup>), plot (10<sup>0</sup> m<sup>2</sup>), and species characteristics 105 predict foliar sodium content? For example, does foliar sodium vary predictably among plant taxa, with 106 distance to coast, or along a gradient of soil pH or site aridity? (2) Responses of foliar sodium to a 107 changing environment: Do selective herbivory or elevated nutrient supply reduce foliar sodium at the 108 local (plot) scale? (3) Effects of foliar sodium on grassland species composition: Does a grassland species' 109 foliar sodium content predict changes in the species' relative abundance in response to herbivory or 110 elevated nutrients?

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| 3<br>4         | 112 | METHODS  |
| 5<br>6<br>7    | 113 | Experimental design and locations. Samples for this study were collected at 26 sites that are part of a  |
| 8<br>9         | 114 | long-term, nutrient-addition and herbivore-fencing experiment being performed in herbaceous-   |
| 10<br>11       | 115 | dominated sites around the world, the Nutrient Network distributed experiment (NutNet,   |
| 12<br>13       | 116 | www.nutnet.org). The subset of the NutNet sites that were able to collect tissue samples that comprise   |
| 14<br>15<br>16 | 117 | the data used in this study spanned Africa, Australia, Europe, and North America (SI Table 1).   |
| 17<br>18<br>19 | 118 | Each site had three experimental blocks composed of 10 – 5 x 5 m plots, each assigned randomly to one  |
| 20<br>21       | 119 | of 10 unique treatment combinations. Treatments included a factorial addition of N (10 g N m $^{-2}$ yr $^{-1}$ as   |
| 22<br>23       | 120 | timed-release urea [(NH <sub>2</sub> ) <sub>2</sub> CO]), P (10 g P m <sup>-2</sup> yr <sup>-1</sup> as triple-super phosphate [Ca(H <sub>2</sub> PO <sub>4</sub> ) <sub>2</sub> ]), and K |
| 24<br>25       | 121 | (10 g K m <sup>-2</sup> yr <sup>-1</sup> as potassium sulphate [K <sub>2</sub> SO <sub>4</sub> ]) plus micronutrients ( $\mu$ , a mix of Fe (15%), S (14%), Mg                             |
| 26<br>27<br>28 | 122 | (1.5%), Mn (2.5%), Cu (1%), Zn (1%), B (0.2%) and Mo [0.05%]), for a total of 8 plots/block. Importantly,  |
| 28<br>29<br>30 | 123 | no sodium (Na) was added in any treatment. N, P, and K were applied annually at each site for 2-4 years  |
| 31<br>32       | 124 | (SI Table 1); the micronutrient mix, $\mu$ , was applied once in the first experimental year to avoid toxicity.  |
| 33<br>34<br>35 | 125 | For the focal fence and fertilization experiment, fence treatments were crossed with the control and the   |
| 36<br>37       | 126 | all nutrient treatment (N+P+K $\mu$ ), adding two fenced plots to each block. Fences were built to exclude   |
| 38<br>39<br>40 | 127 | medium and large mammals and had been in place for 2-4 years at the time of sampling. Fences were  |
| 41<br>42       | 128 | 230 cm tall with four strands of barbless wire suspended at equal vertical distances above the lower   |
| 43<br>44       | 129 | 90 cm which was surrounded by 1-cm woven wire mesh with a 30-cm outward-facing flange stapled to   |
| 45<br>46       | 130 | the ground. At some sites, logistical considerations required slight modifications of the fence design   |
| 47<br>48<br>49 | 131 | (Fence exceptions table, SI Table 2). All sampling plots were separated by at least 1 m wide walkways to   |
| 50<br>51       | 132 | reduce the impact of treatments on adjacent plots. For additional methods details, see (Borer et al.   |
| 52<br>53       | 133 | 2014).   |
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134 Pre-treatment soil collection. Before applying the experimental treatments, three 2.5 x 10cm soil cores 135 were collected from each experimental plot, combined, homogenized into a single sample for each 5 x 5 136 m plot (roughly 500 g of soil), and dried. Percent soil C and N from each plot were analyzed in a single 137 analytical laboratory using a Costech ECS 4010 CHNSO Analyzer on pulverized soil (Knops lab, University 138 of Nebraska, USA). Extractable soil P, K, and micronutrients, including Na, and pH for every soil sample 139 also were quantified in a single analytical laboratory using standard methods (Borer et al. 2014) (A&L 140 Laboratories, Memphis, Tennessee, USA). Across our study sites, plot-level soil sodium ranged from 21 141 ppm (at Val Mustair in Switzerland) to 150 ppm (at Elliott Chaparral, USA). 142 Plant abundance and biomass estimation. To determine the most abundant plant species in each plot

and the change in cover of species in response to the experimental treatments, the percent areal cover
 of each species was estimated to the nearest 1 percent for each species within a permanently marked 1 m<sup>2</sup> subplot of each treatment unit.

A metric of site-level net herbivore impact was estimated as the average difference in live mass inside and outside of fences within a block during the first year of the treatment. To estimate this, we clipped the aboveground biomass of all plants rooted within a 0.2 m<sup>2</sup> area of each fenced and control plot. Each sample was divided into growth from the current year and litter from previous years. We used the first year of treatment to estimate herbivore impact on vegetation mass, prior to species-level selection and turnover in response to long-term herbivore exclusion.

152 Foliar sampling & sodium analysis. Within each plot, the most abundant species were determined as a
 153 function of percent cover, and a single healthy leaf was collected from five unique individuals of the
 154 species with the greatest cover at the site. Most sites had three to five dominant species present in most
 155 plots; however, one site collected 8 different species (Val Mustair), because there were not clearly
 156 dominant species. All leaves were transported in a cooler, and then dried at 60°C for 48 hours (Firn *et al.*

| 2<br>3<br>4    | 157 | 2019). The collected species represented 5.3% (Val Mustair, Switzerland, a high elevation, highly diverse       |
|----------------|-----|---|
| 5<br>6         | 158 | (25 species/plot) site; this is the site that sampled 8 species) to 52.1% (Saline, KS, USA) of the total plot   |
| 7<br>8         | 159 | cover with an average representation of 26% of the total cover across all plots and sites (SI Table 1). All     |
| 9<br>10<br>11  | 160 | leaves were then sent to Queensland University of Technology (Dr. J. Firn) for sodium analysis. Dried           |
| 12<br>13       | 161 | leaves were ground to a fine powder, then analyzed for sodium content with an Agilent 8800 Laser                |
| 14<br>15       | 162 | Ablation Inductively Coupled Plasma Mass Spectrometer (LA-ICP-MS), following Duodu et al. (Duodu et             |
| 16<br>17       | 163 | al. 2015) with two exceptions: C, the most abundant naturally occurring element, was used as a                  |
| 18<br>19<br>20 | 164 | standard, and no additional pulverizing was performed beyond that required for C analysis. The                  |
| 20<br>21<br>22 | 165 | reference material for sodium was NIST SRM 1570a Trace elements in spinach leaves (USA National                 |
| 23<br>24       | 166 | Institute of Standards and Technology 2014). Elemental quantification followed the method of Longerich          |
| 25<br>26       | 167 | et al. (1996), using Iolite, a data reduction software (Paton <i>et al.</i> 2010).                              |
| 27<br>28<br>29 | 168 | <u>Climate data.</u> The WorldClim database provided comparable long-term climate data for all sites (version   |
| 29<br>30<br>31 | 169 | 1.4; http://www.worldclim.org/bioclim). These global climate data were interpolated at high-resolution          |
| 32<br>33       | 170 | from data stations with 10 to 30 years of data (Hijmans <i>et al.</i> 2005). We used these data to test whether |
| 34<br>35       | 171 | foliar sodium in the most abundant taxa declined with mean annual precipitation (MAP in mm per year)            |
| 36<br>37       | 172 | or increased with a site-level index of aridity (MAP divided by potential evapotranspiration in mm per          |
| 38<br>39<br>40 | 173 | year)(Barrow 1992). Site-level MAP ranged from 14 at Sheep Station, USA to 1898 mm of annual                    |
| 41<br>42       | 174 | precipitation at HJ Andrews LTER; Lookout, USA and the index of aridity ranged from 0.2 at Mount                |
| 43<br>44       | 175 | Caroline, Western Australia to 2.4 mm at Val Mustair, Switzerland (SI Table 1).                                 |
| 45<br>46       |     |   |
| 47<br>48       | 176 | Analyses. We explored the relative importance and interactions among the many factors that we                   |
| 49<br>50<br>51 | 177 | hypothesized to constrain foliar sodium. Many of these factors could covary (e.g., annual precipitation,        |
| 52<br>53       | 178 | distance to coast, and soil pH), and it was possible that there could be multiple models that were              |
| 54<br>55       | 179 | similarly informative (i.e., had similar AICc values). For this reason, we used a multi-model approach,         |
| 56<br>57       | 180 | which does not try to identify a single best model (Grueber et al. 2011). This information theoretic            |
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approach starts by calculating all possible subsets of the parameters in the full model, and then uses Akaike's information criterion (AICc) to determine the subset of models sharing similarly high levels of parsimony (Grueber et al. 2011). In our case, we included in our high parsimony set all models that fell within 4 AICc units of the model with the lowest AICc value (Grueber et al. 2011). Parameter estimates and significance are based on a weighted average of the set of high parsimony models. We present the weighted average parameter value estimate, significance, and the summed AIC weights for all models in which the parameter is included, or *importance*. We used the *dredge* function in the MuMIn R library to calculate the AICc of all possible models and the model.avg function in the MuMIn R library to calculated the weighted parameter and statistics.

All models used a random effect structure with site and species within site treated as random intercepts to account for the hierarchical nature of the sampling. To examine biogeographic predictors of foliar sodium, we examined only control plot values, but for the effects of environmental change, we used data from all experimental plots. Experimental treatments were retained in all models. Because of missing soil data, one site (Mt. Caroline) is excluded from experimental analyses. In addition, to avoid bias from having rare species that were found only in one treatment driving the results, for the analysis of the fence and fertilization experiment (shown in Fig 4), we include species that are present in Control plots and at least two other treatments (e.g., Control, Fence, and Fertilized or Control, Fence, and Fence + Fertilized). Similarly, for analysis of the factorial nutrient experiment (SI Figure 1), we include only species present in Control plots and at least 5 other treatments. Finally, in analyses of abiotic factors associated with foliar sodium, we tested the leverage of two outlier sites. In particular, we examined the role of a single site (Sheep Station, USA) in driving the association of foliar sodium with soil pH and another site (Lancaster, UK) in determining the importance of distance from the coast in foliar sodium content.

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| 2<br>3<br>4          | 204 | In addition to assessing foliar sodium, we also used multi-model inference to examine the cover                |    |
| 5<br>6<br>7<br>8     | 205 | response of each plant for which sodium was measured in each plot as a function of the sodium                  |    |
|                      | 206 | concentration of that species. For assessing the effects of foliar sodium on plant cover in response to t      | he |
| 9<br>10<br>11        | 207 | experimental treatments, species with less than 0.1% cover in a plot were removed (23 out of 1,828             |    |
| 12<br>13             | 208 | records or 1.3%).  |    |
| 14<br>15<br>16<br>17 | 209 | All analyses were performed in R (version 3.3; R Foundation for Statistical Computing).                        |    |
| 18<br>19             | 210 | RESULTS  |    |
| 20<br>21<br>22       | 211 | Patterns of foliar sodium  |    |
| 23<br>24<br>25       | 212 | Foliar sodium in 201 of the most abundant grassland plant species from 26 sites on four continents,            |    |
| 25<br>26<br>27       | 213 | including representatives of 32 plant families, varied across five orders of magnitude among sites and         |    |
| 28<br>29<br>30<br>31 | 214 | the most abundant plant taxa in unmanipulated plots. Foliar sodium ranged from 0.5 ppm in Phleum               |    |
|                      | 215 | pratense (Poaceae) to 28,271 ppm in Epaltes australis (Asteraceae, SI Table 1), and average site-level         |    |
| 32<br>33<br>34       | 216 | plant sodium across the most abundant species ranged from 2.7 ppm (at Konza Prairie in the North               |    |
| 35<br>36             | 217 | American Great Plains) to 9,715 ppm (at Burrawan in southeastern Australia). Foliar sodium of the mo           | st |
| 37<br>38             | 218 | abundant species in control plots was similar across grasses with C4 (463 $\pm$ 201 ppm) and C3 (624 $\pm$ 159 | )  |
| 39<br>40             | 219 | ppm) photosynthetic pathways ( $P$ = 0.10). However, across all taxa in unmanipulated (control) plots,         |    |
| 41<br>42<br>43       | 220 | foliar sodium varied spatially both within and among sites (Fig. 1); mean foliar sodium content also           |    |
| 44<br>45             | 221 | varied substantially among plant families (Fig. 1, <i>P</i> <0.001, SI Table 3).                               |    |
| 46<br>47             | 222 | We found that among sites, mean site-scale foliar sodium in control plots increased with soil sodium           |    |
| 48<br>49<br>50       | 223 | (Fig. 2, P=0.015; t=2.68), whereas within sites, foliar sodium did not co-vary with plot-scale soil sodium     | ۱  |
| 51<br>52             | 224 | (P=0.51; t=0.64). In a model that included multiple candidate predictors (site aridity, distance from          |    |
| 53<br>54             | 225 | coast, soil pH, photosynthetic pathway, and soil sodium), foliar sodium declined with increasing site-         |    |
| 55<br>56<br>57       | 226 | level water availability (increasing AI; P=0.001) and soil pH (Fig. 3, P=0.04, SI Table 4). However, our       |    |
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model selection criteria did not retain soil sodium or photosynthetic pathway in final models. The decline in foliar sodium was similar across both coastal and inland sites except for a single site in the UK with high precipitation and exceptionally high sodium ion deposition relative to most locations on Earth (Vet et al. 2014) (Fig 3b, Lancaster, UK). In contrast, for sites with neutral to acidic soils (all except one in this study, Sheep Station, USA), there was no relationship between foliar sodium and soil pH (Fig. 3). Thus, the biogeographic variation in foliar sodium content is explained, in part, by a combination of local conditions, including soil sodium availability and aridity. *Responses of foliar sodium to a changing environment* 

Nutrients and herbivory interacted to determine the foliar sodium of the most abundant plants, and the strength of this effect depended on aridity but not soil pH (SI Table 5). In particular, at mesic sites, when herbivores were present, nutrient addition favored abundant plants with high foliar sodium compared to plants in ambient (control) plots (Fig. 4a, SI Table 5). As a result, the addition of the full suite of nutrients (N+P+K $\mu$ , but not Na) outside of fences weakened the negative effect of increasing water availability (increasing AI) on foliar sodium content (Fig. 4b). The factorial nutrient addition experiment clarified that the interaction between aridity and nutrient supply was primarily driven by the effects of potassium and micronutrients (Kµ) and to a lesser extent the effects of nitrogen and phosphorus addition (SI Table 6, SI Fig. 1).

We examined the subset of species that were sampled multiple times among plots and sites to explore the role of intraspecific variability of sodium content in determining these observed responses. Of the 245 246 201 species in this experiment, 41 were among the most abundant (and therefore sampled) in plots at more than one site, and 94 were sampled in both control and treatment plots within sites. Models of the subset of species present among sites and in both control and treatment plots were qualitatively similar to models of the larger dataset for both experiments (SI Tables 7 and 8), suggesting that some of

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the observed variation in foliar chemistry is attributable to intraspecific change in foliar sodium contentin response to the biotic and abiotic environment.

252 Effects of foliar sodium on grassland species composition

253 The sodium content of foliage and plot-scale nutrient supply contributed to the effects of herbivores on 254 changes in the relative abundance of grassland plant species. Fertilization (with NPK $\mu$ ) increased the 255 cover of the most abundant species, and in the presence of herbivores, the abundance of species low in 256 foliar sodium increased in response to fertilization, whereas high sodium species became less abundant 257 when fertilized (Fig. 5). However, in the absence of herbivores, fertilization had no consistent effects on 258 species abundances in relation to their foliar sodium concentration (SI Table 9, SI Fig. 2). These effects 259 on foliar sodium were independent of the intensity of herbivory among sites (measured as the site-level 260 log ratio of live biomass inside and outside of herbivore exclusion fences (P > 0.57 for all main effects 261 and interactions; importance < 0.40 [model not shown]). The factorial nutrient addition experiment 262 clarified that, in the presence of herbivores, the addition of any elemental nutrient caused dominant 263 plant species with relatively high foliar sodium content to decline more than species with lower foliar 264 sodium (SI Table 10); this effect was greatest in response to fertilization with P (SI Table 10). These 265 results point to selective consumption by herbivores of high nutrient, high sodium plants.

266 DISCUSSION

267 This multi-continent, biogeographic study demonstrated that foliar sodium in dominant grassland plants 268 is highly variable among sites and even plots within a site, and there also is significant variation in foliar 269 sodium among families and taxa within families, regardless of geographic location. These patterns likely 270 reflect variation in long-term environmental conditions (e.g., aridity, grazing) that have selected for 271 species with differing strategies for environmental sodium uptake. While there is evidence for 272 phylogenetic conservation of cation transport proteins that can influence sodium uptake (Schachtman &

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273 Liu 1999) with predictable differences across photosynthetic pathways (Brownell & Crossland 1972), 274 photosynthetic pathway was not a predictor of foliar sodium in grasses. Nonetheless, the very highest 275 foliar sodium content recorded in this study was 9% (91,818 ppm) in *Eragrostis curvula* (Poaceae, 276 commonly called African Lovegrass) found at Burrawan, Australia. This species has a C4 photosynthetic 277 pathway, indicating a physiological requirement for sodium, and this site is among the more arid sites in 278 the experiment, suggesting that both photosynthetic pathway (Brownell & Crossland 1972; Furumoto et 279 al. 2011) and aridity (Raheja 1966) can be strongly associated with foliar sodium, in some cases. 280 However, while individual species supported this hypothesis, as a group, C<sub>4</sub> grasses were not 281 consistently high in foliar sodium. 282 The results of this globally-extensive study demonstrate that the relative abundance of plant species in 283 grasslands is altered by herbivores as a function of sodium content and elemental nutrient supply. In 284 particular, herbivores in grasslands spanning four continents with a variety of herbivore types and 285 densities consistently reduced the cover of plants with high foliar sodium only in high nutrient 286 conditions. The reduction in abundance of sodium-rich plants in fertilized plots is evidence of targeted 287 herbivory of high sodium, protein-rich plants. In particular, herbivores are attracted to plots with 288 elevated nutrients (Mattson 1980), and selective consumption reduces the abundance those species 289 with the highest sodium. These plants are not likely extirpated from the community, since the same 290 species are generally found at higher abundance inside herbivore exclosures, rather they are likely to be 291 in a constant state of regrowth from having their aboveground foliage selectively consumed. Such 292 selective foraging is common in many ecosystems (Belovsky 1981; Jefferies et al. 1994; Wallis de Vries & 293 Schippers 1994; Bartolome et al. 1998; Doughty et al. 2016). Related to this, the impact of herbivores on 294 sodium content of the most abundant plant species was contingent on aridity, with foliar sodium 295 content high and indistinguishable among experimental treatments at arid sites, but declining with 296 increasing water availability. Our arid region results are consistent with previous work that found

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| 3<br>4                           | 297 | positive feedbacks generating and maintaining high sodium content grazing lawns because of high          |
| 5<br>6                           | 298 | evaporation rates under the cropped vegetation (McNaughton 1988). By examining herbivore impa            |
| 7<br>8<br>9                      | 299 | across a much broader precipitation gradient, we demonstrate that both aridity and herbivory             |
| 9<br>10<br>11                    | 300 | determine foliar sodium biogeography across the world's grasslands, with declining sodium conten         |
| 12<br>13                         | 301 | under increased precipitation and preferential feeding by herbivores.                                    |
| 14<br>15<br>16                   | 302 | Our experimental work also demonstrated that the sodium content of locally abundant plants increased     |
| 17<br>18                         | 303 | with soil sodium at the site-scale; however, when included in models, site aridity was a much more       |
| 19<br>20                         | 304 | effective predictor of biogeographic variation in foliar sodium than soil sodium. At broad spatial sca   |
| 21<br>22<br>23                   | 305 | foliar sodium is positively related to soil sodium as has been observed in previous work (Sutcliffe 1    |
| 23<br>24<br>25                   | 306 | Epstein 1973; Pardo & Quintero 2002; Maathuis 2014), but foliar sodium was not strongly predicte         |
| 26<br>27                         | 307 | distance to coast, a common a surrogate for sodium ion deposition (Vet et al. 2014). However, bec        |
| 28<br>29                         | 308 | arid regions are characterized by high evapotranspiration relative to precipitation, these sites tend    |
| 30<br>31<br>32                   | 309 | accumulate salts over time (Raheja 1966). In contrast, coastal sites may have both high ion input a      |
| 33<br>34                         | 310 | high precipitation (Vet et al. 2014), reducing the environmental pools of ions, including sodium, an     |
| 35<br>36                         | 311 | causing a mismatch between salt deposition and the location of sodic soils (Wicke et al. 2011). In the   |
| 37<br>38                         | 312 | study, the coastal site with exceptionally high foliar sodium relative to site-scale precipitation (Land |
| 39<br>40<br>41                   | 313 | UK) is also situated in a location on Earth with an exceptionally high rate of sodium ion input (Vet e   |
| 42<br>43                         | 314 | 2014), suggesting that site aridity combined with direct measures of site-level sodium ion input rat     |
| 44<br>45                         | 315 | likely provide even better predictions of site-level foliar sodium in the most abundant plant taxa. In   |
| 46<br>47                         | 316 | addition, although we found a decline in foliar sodium with increasing soil pH, this pattern was o       |
| 48<br>49<br>50                   | 317 | by a single, arid site in the intermountain west of the USA. While this pattern is consistent with       |
| 50<br>51<br>52                   | 318 | expectations of reduced cation uptake in higher pH soils (Tyler & Olsson 2001; Bolan & Brennan 2         |
| 53<br>54                         | 319 | we have only a single site with a pH above neutral. Because soil pH is intimately associated with        |
| 55<br>56<br>57<br>58<br>59<br>60 | 320 | aridity (Slessarev et al. 2016), disentangling the roles of soil pH and aridity in determining grass     |

r experimental work also demonstrated that the sodium content of locally abundant plants increases h soil sodium at the site-scale; however, when included in models, site aridity was a much more ective predictor of biogeographic variation in foliar sodium than soil sodium. At broad spatial scales, ar sodium is positively related to soil sodium as has been observed in previous work (Sutcliffe 1959; tein 1973; Pardo & Quintero 2002; Maathuis 2014), but foliar sodium was not strongly predicted by tance to coast, a common a surrogate for sodium ion deposition (Vet *et al.* 2014). However, because l regions are characterized by high evapotranspiration relative to precipitation, these sites tend to umulate salts over time (Raheja 1966). In contrast, coastal sites may have both high ion input and h precipitation (Vet et al. 2014), reducing the environmental pools of ions, including sodium, and sing a mismatch between salt deposition and the location of sodic soils (Wicke *et al.* 2011). In this dy, the coastal site with exceptionally high foliar sodium relative to site-scale precipitation (Lancaster, is also situated in a location on Earth with an exceptionally high rate of sodium ion input (Vet et al. 4), suggesting that site aridity combined with direct measures of site-level sodium ion input rate will ly provide even better predictions of site-level foliar sodium in the most abundant plant taxa. In lition, although we found a decline in foliar sodium with increasing soil pH, this pattern was driven a single, arid site in the intermountain west of the USA. While this pattern is consistent with pectations of reduced cation uptake in higher pH soils (Tyler & Olsson 2001; Bolan & Brennan 2011), have only a single site with a pH above neutral. Because soil pH is intimately associated with dity (Slessarev et al. 2016), disentangling the roles of soil pH and aridity in determining grassland

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|               | 321 | plant sodium biogeography will require more thorough sampling, particularly at sites with basic soils       |
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|               | 322 | spanning a range of aridity. Nonetheless, the strong spatial variation in foliar sodium suggests that       |
|               | 323 | environmental context is key in determining foliar sodium which, by extension, implies that future          |
| )             | 324 | environmental changes may alter foliar sodium for herbivores. Given the importance of dietary sodium        |
| 2             | 325 | for herbivores (Seastedt & D. A. Crossley 1981; McNaughton 1988; McNaughton et al. 1997; Kaspari et         |
| <br>          | 326 | al. 2008; Doughty et al. 2016), biogeographic patterns of foliar sodium in abundant grassland plants may    |
| ,<br>,        | 327 | arise from interactions with wild herbivores, and likely have significant implications for the distribution |
| )             | 328 | and impacts of consumers in grassland ecosystems.   |
| )             | 329 | The strong difference in the physiological importance of sodium to grassland plants and wild herbivores     |
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|               |     | has gained increasing attention in ecology, with recent calls for a greater understanding of the            |
| ,             | 331 | biogeography of sodium (Kaspari <i>et al.</i> 2008). The current study of both patterns and responses to    |
| )             | 332 | experimental manipulation, performed at 26 sites spanning wide biotic and abiotic gradients,                |
| )             | 333 | demonstrates that aridity, soil acidity, nutrient supply, and herbivory, interact to influence              |
| -<br>;<br>;   | 334 | biogeographic patterns of foliar sodium and its effect on plant abundance. In future environments,          |
| 5             | 335 | climate change is expected to impact global patterns of soil salinity via changes in precipitation and      |
| ,<br>;        | 336 | evapotranspiration (Schofield & Kirkby 2003). The current results suggest that the impact of these          |
| )             | 337 | changes on grassland plant composition will depend on the interactive effects of large-scale changes in     |
| 2             | 338 | aridity and elemental nutrient (N, P) supply and the resulting nutritional value for consumers.             |
| ,<br>         | 339 | ACKNOWLEDGMENTS   |
| ,<br>,        | 340 | This work was generated using data from the Nutrient Network experiment ( <u>http://www.nutnet.org</u> ),   |
| ;<br>)        | 341 | funded at the site-scale by individual researchers. Coordination and data management have been              |
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| 3<br>4         | 345 | Minnesota Supercomputer Institute for hosting project data and the Institute on the Environment and          |
| 5<br>6         | 346 | sDiv at the University of Leipzig for hosting Network meetings. We thank QUT's Central Analytical            |
| 7<br>8<br>9    | 347 | Facilities (CARF), part of the Institute of Future Environments (IFE) for use of their facilities to analyze |
| 9<br>10<br>11  | 348 | leaf nutrient concentrations. Author contributions are listed in SI Table 11 and data contributors are       |
| 12<br>13<br>14 | 349 | listed in SI Table 12.   |
| 15<br>16       | 350 | Code availability R code of all analyses will be made available via GitHub (https://github.com/).            |
| 17<br>18       | 351 | Data availability Data supporting the findings of this study will be made available on Dryad                 |
| 19<br>20<br>21 | 352 | (http://datadryad.org).  |
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Fig. 1. Foliar Na variation across taxonomic and spatial scales: Variance components analysis of foliar sodium in the 85 locally-abundant plant taxa from control plots at 26 sites across nested taxonomic and spatial scales. Foliar sodium for 41 species was measured at two or more sites. Variation in foliar sodium associated with plant location is shown in orange; variation associated with taxonomic groups is shown in blue. Variance explained by genus is extremely small, but non-zero (<3x10<sup>-6</sup>), thus is barely visible in this graph. SI Table 3 provides the full statistical model associated with this figure.





Fig. 3. Predictors of foliar Na: The foliar sodium of the most abundant plant species declined across a gradient of plot-scale pH (z=2.03, P=0.04) and site-scale water availability (MAP/PET; z=3.24, P=0.001). Data include the 85 taxa across 22 sites that were growing in control plots. Coastal (orange) and Inland (blue) are divided at 100km from a coast. The dashed yellow line shows the model with all sites included; the solid yellow line shows these relationships without a single site in the UK (Lancaster, orange circled site) with high precipitation and coastal salt input. Similarly, the dashed blue line shows the model with all sites included; the solid blue line shows the relationships without the only site with basic soil pH found in US Intermountain West (Sheep Station, blue circled site). Error bars represent ±SE. SI Table 4 provides the full statistical model associated with the solid lines shown in this figure. 



Fig. 4. Responses of foliar sodium to changes in herbivory and nutrient supply: Response of foliar Na in
153 locally abundant plants to a factorial combination of fencing to reduce vertebrate herbivory and
fertilization by a suite of micro- and macronutrients (not including Na<sup>+</sup>) (a) across a gradient in plot-scale
pH and (b) across a gradient in site-scale water availability. Foliar sodium is higher than expected from
control plots where precipitation is relatively high and nutrients are added (z=3.49, P=0.0005). Error bars
represent ±SE. SI Table 5 provides details of the full statistical model.



## 557 SI: DATA AND MODEL TABLES UNDERLYING RESUTS TEXT

558 SI Table 1. Sites, locations, mean annual precipitation (MAP), index of aridity, modeled nitrogen deposition (N Dep.), measured plot-scale soil

559 pH, and measured foliar sodium in each of the most abundant species at the site (leaf Na (ppm).

| Site name                 | Continent     | Country | Latitude   | Longitude  | MAP  | AI     | Leaf Na (ppm) | Soil pH | Soil Na (ppm) |
|---------------------------|---------------|---------|------------|------------|------|--------|---------------|---------|---------------|
| Mt Gilboa                 | Africa        | ZA      | -29.28424  | 30.29174   | 943  | 0.7797 | 233.13        | 5.07    | 35.58         |
| Summerveld                | Africa        | ZA      | -29.81161  | 30.71573   | 944  | 0.7324 | 125.01        | 5.15    | 43.58         |
| Bogong                    | Australia     | AU      | -36.874    | 147.254    | 1678 | 1.9159 | 228.35        | 4.47    | 22.10         |
| Burrawan                  | Australia     | AU      | -27.734896 | 151.139517 | 643  | 0.4335 | 9715.51       | 5.55    | 59.53         |
| Kinypanial                | Australia     | AU      | -36.2      | 143.75     | 408  | 0.3224 | 751.22        | 6.04    | 148.43        |
| Mt. Caroline              | Australia     | AU      | -31.782138 | 117.610853 | 324  | 0.2186 | 7628.36       | 5.29    | 38.19         |
| Fruebuel                  | Europe        | СН      | 47.113187  | 8.541821   | 1546 | 2.0892 | 3.86          | 5.46    | 25.50         |
| Val Mustair               | Europe        | СН      | 46.631345  | 10.372252  | 681  | 2.4389 | 38.31         | 5.66    | 26.70         |
| Companhia das Lezirias    | Europe        | РТ      | 38         | -8         | 564  | 0.4532 | 65.69         | 5.93    | 25.81         |
| Lancaster                 | Europe        | UK      | 53.9856247 | -2.6284176 | 1522 | 2.2003 | 2478.44       | 4.77    | 41.56         |
| Cowichan                  | North America | СА      | 48.46      | -123.38    | 762  | 1.0743 | 112.67        | 5.63    | 48.60         |
| Boulder South Campus      | North America | US      | 39.972022  | -105.23354 | 487  | 0.3701 | 2358.66       | 6.82    | 58.39         |
| Bunchgrass (Andrews LTER) | North America | US      | 44.2766854 | -121.96802 | 1618 | 1.9348 | 38.65         | 5.54    | 23.71         |

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|     | Chichaqua Bottoms          | North America | US | 41.7850667 | -93.385383 | 871  | 0.849  | 22.15  | 6.11 | 21.94  |
|-----|----------------------------|---------------|----|------------|------------|------|--------|--------|------|--------|
|     | Duke Forest                | North America | US | 36.00828   | -79.020423 | 1157 | 0.9121 | 70.05  | 5.27 | 19.07  |
|     | Elliott Chaparral          | North America | US | 32.875     | -117.05224 | 344  | 0.2565 | 459.89 | 5.69 | 145.46 |
|     | Hopland REC                | North America | US | 39.0127534 | -123.06031 | 1065 | 0.8593 | 346.67 | NA   | 22.99  |
|     | Konza LTER                 | North America | US | 39.070856  | -96.582821 | 889  | 0.7608 | 2.67   | NA   | 20.56  |
|     | Lookout (Andrews LTER)     | North America | US | 44.2051771 | -122.12845 | 1877 | 2.3085 | 246.88 | 5.07 | 20.83  |
|     | Mclaughlin UCNRS           | North America | US | 38.8642721 | -122.40641 | 936  | 0.6615 | 316.49 | NA   | 42.48  |
|     | Sagehen Creek UCNRS        | North America | US | 39.43      | -120.24    | 831  | 0.8579 | 307.13 | 5.93 | 63.67  |
|     | Saline Experimental Range  | North America | US | 39.05      | -99.1      | 608  | 0.491  | 41.99  | NA   | 23.67  |
|     | Sheep Experimental Station | North America | US | 44.242989  | -112.19839 | 246  | 0.2689 | 14.02  | 7.98 | 23.54  |
|     | Shortgrass Steppe LTER     | North America | US | 40.81667   | -104.76667 | 369  | 0.3244 | 36.65  | 6.16 | 21.88  |
|     | Sierra Foothills REC       | North America | US | 39.2355096 | -121.2837  | 936  | 0.6932 | 42.19  | 5.96 | 36.04  |
|     | Smith Prairie              | North America | US | 48.2065807 | -122.62475 | 605  | 0.7796 | 421.54 | 6.09 | 43.53  |
| 560 |                            |               |    |            |            |      |        |        |      |        |

| 5                    |     | Site name  | Fence Typ       | pe Exce      | eption description   |  |  |  |  |  |
|----------------------|-----|--|-----------------|--------------|----------------------|--|--|--|--|--|
| 7<br>8<br>9          |     | Lancaster  | Sheep           | Sim          | ilar to NutNet stand | ard but top strand at 1.2 m  |  |  |  |  |
| 10<br>11             |     | Sheep Experimental   | Sheep           | Sim          | ilar to NutNet stand | ard but top strand at 1.2 m  |  |  |  |  |
| 12<br>13             |     | Station  |                 |              |                      |  |  |  |  |  |
| 14<br>15<br>16<br>17 |     | Val Mustair  | Val Musta       |              | 4                    | 5 cm diameter) driven 70 cm into ground, 3 m apart, covered with 5<br>high and with extra cabling and supports to prevent snow damage. |  |  |  |  |
| 18<br>19<br>20       |     |  |                 | Fen          | ces enclose 6 m x 7  | m area.  |  |  |  |  |
| 21<br>22             | 565 |  |                 |              |                      |  |  |  |  |  |
| 23<br>24             | 566 | SI Table 3. Patterns of foliar Na: Analysis of spatial and taxonomic variance components in foliar sodium of 85 locally abundant grassland species |                 |              |                      |  |  |  |  |  |
| 25<br>26<br>27       | 567 | found in the unmanipulate  | ed control plot | s of 26 site | 5.                   |  |  |  |  |  |
| 27<br>28<br>29       | 568 | Random effects:  |                 |              |                      |  |  |  |  |  |
| 30                   | 569 | Groups   | Name            | Variance     | Std.Dev.             | Number of obs for group  |  |  |  |  |
| 31<br>32             | 570 | Taxon:(genus:Family)   | (Intercept)     | 2.389e-02    | 4.888e-01            | 85   |  |  |  |  |
| 33                   | 571 | genus:Family   | (Intercept)     | 7.450e-12    | 2 2.729e-06          | 66   |  |  |  |  |
| 34<br>35             | 572 | plot:site_code   | (Intercept)     | 8.214e-02    | 2 2.866e-01          | 60   |  |  |  |  |
| 36                   | 573 | site_code  | (Intercept)     | 8.052e-02    | 8.973e-01            | 22   |  |  |  |  |
| 37<br>38             | 574 | Family   | (Intercept)     | 2.924e-02    | 2 1.710e-01          | 17   |  |  |  |  |
| 39<br>40             | 575 | Residual   |                 | 4.623e-02    | 2 2.150e-01          |  |  |  |  |  |
| 41                   | 576 |  |                 |              |                      |  |  |  |  |  |
| 42<br>43             |     |  |                 |              |                      |  |  |  |  |  |
| 44                   |     |  |                 |              |                      | 26   |  |  |  |  |
| 45<br>46             |     |  |                 |              |                      |  |  |  |  |  |
| 47                   |     |  |                 |              |                      |  |  |  |  |  |

| 1<br>2   |     |  |                   |                 |              |            |                       |                     |                          |  |  |  |
|----------|-----|--|-------------------|-----------------|--------------|------------|-----------------------|---------------------|--------------------------|--|--|--|
| 3<br>4   | 577 | SI Table 4. Predictors of foliar   | Na:               |                 |              |            |                       |                     |                          |  |  |  |
| 5<br>6   | 578 | Variation of site-level mean foliar sodium with distance to coast, aridity (MAP/PET), and soil pH for the 85 dominant grassland species found in |                   |                 |              |            |                       |                     |                          |  |  |  |
| 7<br>8   | 579 | the control plots of the 26 stuc   | ly sites. Model s | hows the condit | ional averag | e estimat  | es of model param     | eters for all sites | except the very high     |  |  |  |
| 9<br>10  | 580 | precipitation, very high sodium influx site (Lancaster; see Figure and legend in main text).   |                   |                 |              |            |                       |                     |                          |  |  |  |
| 11<br>12 | 581 | Estimate Std. Error Adjusted SE z value $Pr(> z )$ Importance Num models   |                   |                 |              |            |                       |                     |                          |  |  |  |
| 13<br>14 | 582 | (Intercept)  | 1.7957            | 0.1884          | 0.1897       | 9.466      | < 2e-16 ***           |                     |                          |  |  |  |
| 15<br>16 | 583 | c.coastal  | 0.3700            | 0.4098          | 0.4125       | 0.897      | 0.36975               | 0.78                | 6                        |  |  |  |
| 17<br>18 | 584 | z.AI   | -1.3337           | 0.4089          | 0.4116       | 3.240      | 0.00119 **            | 1.00                | 8                        |  |  |  |
| 19<br>20 | 585 | z.pH   | -0.4880           | 0.2392          | 0.2406       | 2.028      | 0.04252 *             | 1.00                | 8                        |  |  |  |
| 21<br>22 | 586 | c.coastal:z.pH   | -1.3458           | 0.5913          | 0.5954       | 2.260      | 0.02379 *             | 0.66                | 4                        |  |  |  |
| 23<br>24 | 587 | z.soil.na.lg   | 0.2049            | 0.2199          | 0.2213       | 0.926      | 0.35448               | 0.35                | 4                        |  |  |  |
| 25       | 588 | c.coastal:z.AI   | -0.6275           | 1.7061          | 1.7181       | 0.365      | 0.71492               | 0.12                | 1                        |  |  |  |
| 26<br>27 | 589 | c.coastal:z.soil.na.lg   | -0.7663           | 0.5640          | 0.5677       | 1.350      | 0.17712               | 0.13                | 2                        |  |  |  |
| 28<br>29 | 590 |  |                   |                 |              |            |                       |                     |                          |  |  |  |
| 30<br>31 | 591 | Signif. codes: 0 '***'   | 0.001 '**'        | 0.01 '*' 0.0    | )5 '.' 0.1   | ' ' 1      |                       |                     |                          |  |  |  |
| 32<br>33 | 592 |  |                   |                 |              |            |                       |                     |                          |  |  |  |
| 34<br>35 | 593 | SI Tables 5 & 6. Responses of f  | oliar sodium to   | a changing envi | ronment:     |            |                       |                     |                          |  |  |  |
| 36<br>37 | 594 |  |                   |                 |              |            |                       |                     | ulation of houting and   |  |  |  |
| 38       |     | SI Table 5. Response of foliar s   |                   |                 |              |            |                       |                     |                          |  |  |  |
| 39<br>40 | 595 | nutrients. Regression table sho  | ws conditional a  | average model r | esults witho | ut Lancast | ter; when this site i | s included, the r   | esults are qualitatively |  |  |  |
| 41<br>42 |     |  |                   |                 |              |            |                       |                     |                          |  |  |  |
| 43<br>44 |     |  |                   |                 |              |            |                       |                     | 27                       |  |  |  |
| 45<br>46 |     |  |                   |                 |              |            |                       |                     | 21                       |  |  |  |
| 40       |     |  |                   |                 |              |            |                       |                     |                          |  |  |  |

| 2        |     |                          |                   |                |                   |            |             |        |                   |                         |           |
|----------|-----|--------------------------|-------------------|----------------|-------------------|------------|-------------|--------|-------------------|-------------------------|-----------|
| 3<br>4   | 596 | similar but the effect o | of nutrient addit | ion across the | water availabilit | y gradient | is somewha  | t wea  | aker due to the o | extreme outlier. The re | gression  |
| 5<br>6   | 597 | table shows the condi    | tional average v  | alues across m | odels in which p  | arameters  | were includ | ed, tl | ne number of m    | odels in which parame   | ters were |
| 7<br>8   | 598 | included, and their im   | portance in the   | models.        |                   |            |             |        |                   |                         |           |
| 9<br>10  | 599 |                          |                   |                |                   |            |             |        |                   |                         |           |
| 11<br>12 | 600 |                          | Estimate          | Std. Error     | Adjusted SE       | z value    | Pr(> z )    |        | Importance        | Num models              |           |
| 13<br>14 | 601 | (Intercept)              | 1.6286948         | 0.1348117      | 0.1350290         | 12.062     | < 2e-16     | ***    |                   |                         |           |
| 15<br>16 | 602 | z.AI                     | -0.6601019        | 0.3111010      | 0.3116025         | 2.118      | 0.034140    | *      | 1.00              | 5                       |           |
| 17<br>18 | 603 | z.pH                     | -0.3787517        | 0.0792799      | 0.0794058         | 4.770      | 1.8e-06     | ***    | 1.00              | 5                       |           |
| 19<br>20 | 604 | c.Fnc                    | -0.0144227        | 0.0345935      | 0.0346491         | 0.416      | 0.677226    |        | 1.00              | 5                       |           |
| 21<br>22 | 605 | c.NPK                    | 0.0816837         | 0.0347964      | 0.0348523         | 2.344      | 0.019093    | *      | 1.00              | 5                       |           |
| 23<br>24 | 606 | c.Fnc:c.NPK              | 0.0006431         | 0.0686576      | 0.0687668         | 0.009      | 0.992538    |        | 1.00              | 5                       |           |
| 25       | 607 | c.Fnc:z.AI               | -0.0134223        | 0.0699543      | 0.0700666         | 0.192      | 0.848083    |        | 1.00              | 5                       |           |
| 26<br>27 | 608 | c.NPK:z.AI               | 0.2581523         | 0.0739461      | 0.0740549         | 3.486      | 0.000490    | ***    | 1.00              | 5                       |           |
| 28<br>29 | 609 | c.NPK:z.pH               | 0.1199696         | 0.0799398      | 0.0800687         | 1.498      | 0.134047    |        | 0.58              | 3                       |           |
| 30<br>31 | 610 | c.Fnc:c.NPK:z.AI         | -0.4660938        | 0.1361232      | 0.1363347         | 3.419      | 0.000629    | ***    | 1.00              | 5                       |           |
| 32<br>33 | 611 | c.Fnc:z.pH               | 0.0060036         | 0.0802725      | 0.0804007         | 0.075      | 0.940477    |        | 0.35              | 3                       |           |
| 34<br>35 | 612 | c.Fnc:c.NPK:z.pH         | 0.2183428         | 0.1549416      | 0.1551920         | 1.407      | 0.159451    |        | 0.12              | 1                       |           |
| 36       | 613 |                          |                   |                |                   |            |             |        |                   |                         |           |
| 37<br>38 | 614 | Signif. codes: (         | 0 '***' 0.00      | 1 '**' 0.01    | '*' 0.05 '.       | ′ 0.1 ′    | ' 1         |        |                   |                         |           |
| 39<br>40 |     |                          |                   |                |                   |            |             |        |                   |                         |           |
| 41<br>42 |     |                          |                   |                |                   |            |             |        |                   |                         |           |
| 43<br>44 |     |                          |                   |                |                   |            |             |        |                   |                         | 28        |
| 45<br>46 |     |                          |                   |                |                   |            |             |        |                   |                         | 20        |

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 SI Table 6. Response of foliar sodium in 179 dominant grassland plant species growing in plots treated with a factorial addition of elemental nutrients (but not sodium). Model excludes one site (Lancaster) which was a substantial outlier for AI and pH. Models are qualitatively similar with Lancaster included. The regression table shows the conditional average values across models in which parameters were included; the number of models in which parameters were included are shown below the table.

| 13<br>14 | 620 |              | Estimate | Std. Error | Adjusted SE | z value | Pr(> z )    | Importance | Num models |
|----------|-----|--------------|----------|------------|-------------|---------|-------------|------------|------------|
| 15<br>16 | 621 | (Intercept)  | 1.68363  | 0.13936    | 0.13947     | 12.071  | < 2e-16 *** | -          |            |
| 17       | 622 | z.AI         | -0.64993 | 0.31441    | 0.31465     | 2.066   | 0.03887 *   | 1.00       | 40         |
| 18<br>19 | 623 | z.pH         | -0.24259 | 0.05384    | 0.05388     | 4.503   | 6.7e-06 *** | 1.00       | 40         |
| 20<br>21 | 624 | c.K          | -0.01129 | 0.02289    | 0.02291     | 0.493   | 0.62225     | 1.00       | 40         |
| 22       | 625 | c.N          | 0.06474  | 0.02302    | 0.02304     | 2.810   | 0.00495 **  | 1.00       | 40         |
| 23<br>24 | 626 | c.P          | 0.02553  | 0.02282    | 0.02283     | 1.118   | 0.26360     | 1.00       | 40         |
| 25<br>26 | 627 | c.K:c.N      | 0.03479  | 0.04543    | 0.04546     | 0.765   | 0.44413     | 0.95       | 37         |
| 27       | 628 | c.K:z.AI     | 0.24490  | 0.05063    | 0.05066     | 4.834   | 1.3e-06 *** | 1.00       | 40         |
| 28<br>29 | 629 | c.K:z.pH     | 0.10960  | 0.05034    | 0.05038     | 2.176   | 0.02958 *   | 0.92       | 35         |
| 30<br>31 | 630 | c.N:z.AI     | 0.10176  | 0.04981    | 0.04984     | 2.042   | 0.04120 *   | 1.00       | 40         |
| 32       | 631 | c.P:z.AI     | 0.11270  | 0.04591    | 0.04595     | 2.453   | 0.01417 *   | 1.00       | 40         |
| 33<br>34 | 632 | c.K:c.N:z.AI | 0.23943  | 0.09389    | 0.09396     | 2.548   | 0.01083 *   | 0.95       | 37         |
| 35<br>36 | 633 | c.N:z.pH     | -0.07174 | 0.05135    | 0.05139     | 1.396   | 0.16272     | 0.57       | 24         |
| 37       | 634 | c.N:c.P      | 0.05290  | 0.04522    | 0.04525     | 1.169   | 0.24239     | 0.45       | 21         |
| 38<br>39 | 635 | c.K:c.N:z.pH | 0.10350  | 0.09983    | 0.09990     | 1.036   | 0.30019     | 0.18       | 9          |
| 40<br>41 | 636 | c.K:c.P      | 0.03394  | 0.04511    | 0.04515     | 0.752   | 0.45217     | 0.33       | 16         |
|          |     |              |          |            |             |         |             |            |            |

| 1        |     |                        |            |            |         |           |      |    |  |
|----------|-----|------------------------|------------|------------|---------|-----------|------|----|--|
| 2<br>3   | 637 | c.P:z.pH -0.01439      | 0.05017    | 0.05021    | 0.287   | 0.77439   | 0.18 | 10 |  |
| 4<br>5   | 638 | c.K:c.P:z.AI 0.10380   | 0.08934    | 0.08941    | 1.161   | 0.24564   | 0.10 | 5  |  |
| 6<br>7   | 639 | c.N:c.P:z.AI 0.01774   | 0.08995    | 0.09002    | 0.197   | 0.84379   | 0.05 | 3  |  |
| 8        | 640 | c.N:c.P:z.pH -0.12625  | 0.09267    | 0.09274    | 1.361   | 0.17343   | 0.03 | 2  |  |
| 9<br>10  | 641 | c.K:c.N:c.P -0.08298   | 0.09077    | 0.09084    | 0.913   | 0.36103   | 0.03 | 2  |  |
| 11<br>12 | 642 |                        |            |            |         |           |      |    |  |
| 13       | 643 | Signif. codes: 0 '***' | 0.001 '**' | 0.01 '*' 0 | .05 '.' | 0.1 ' ' 1 |      |    |  |
| 14<br>15 | 644 |                        |            |            |         |           |      |    |  |
| 16<br>17 |     |                        |            |            |         |           |      |    |  |
| 18       |     |                        |            |            |         |           |      |    |  |
| 19<br>20 |     |                        |            |            |         |           |      |    |  |
| 21<br>22 |     |                        |            |            |         |           |      |    |  |
| 23       |     |                        |            |            |         |           |      |    |  |
| 24<br>25 |     |                        |            |            |         |           |      |    |  |
| 26       |     |                        |            |            |         |           |      |    |  |
| 27<br>28 |     |                        |            |            |         |           |      |    |  |
| 29<br>30 |     |                        |            |            |         |           |      |    |  |
| 31       |     |                        |            |            |         |           |      |    |  |
| 32<br>33 |     |                        |            |            |         |           |      |    |  |
| 34<br>35 |     |                        |            |            |         |           |      |    |  |
| 36       |     |                        |            |            |         |           |      |    |  |
| 37<br>38 |     |                        |            |            |         |           |      |    |  |
| 39       |     |                        |            |            |         |           |      |    |  |
| 40<br>41 |     |                        |            |            |         |           |      |    |  |
| 42<br>43 |     |                        |            |            |         |           |      |    |  |
| 44       |     |                        |            |            |         |           |      |    |  |
| 45       |     |                        |            |            |         |           |      |    |  |









| 1<br>2      |     |                         |                  |                   |                 |              |                     |                   |                             |       |
|-------------|-----|-------------------------|------------------|-------------------|-----------------|--------------|---------------------|-------------------|-----------------------------|-------|
| 2<br>3<br>4 | 655 | SI Tables 7 and 8. Res  | ponses of folia  | r sodium to a ch  | anging enviro   | onment:      |                     |                   |                             |       |
| 5<br>6      | 656 | SI Table 7. Response of | of foliar sodium | to experimenta    | l manipulatio   | n of herbiv  | vores and nutrients | for the subset c  | f 60 species present in co  | ntrol |
| 7<br>8      | 657 | plots and at least 3 ex | perimentally tr  | eated plots of th | ne fence x fert | ilization ex | operiment. The reg  | ression table sho | ows the conditional average | ge    |
| 9<br>10     | 658 | values across models,   | relative import  | tance values are  | shown below     | the table.   |                     |                   |                             |       |
| 11<br>12    | 659 |                         |                  |                   |                 |              |                     |                   |                             |       |
| 13<br>14    | 660 |                         | Estimate St      | td. Error Ad      | justed SE :     | z value      | Pr(> z )            | Importance        | Num models                  |       |
| 15<br>16    | 661 | (Intercept)             | 1.60760          | 0.15658           | 0.15690         | 10.246       | < 2e-16 ***         |                   |                             |       |
| 17<br>18    | 662 | z.AI                    | -0.90293         | 0.35730           | 0.35803         | 2.522        | 0.01167 *           | 1.00              | 6                           |       |
| 19<br>20    | 663 | z.pH                    | -0.42239         | 0.08360           | 0.08377         | 5.042        | 5e-07 ***           | 1.00              | 6                           |       |
| 21<br>22    | 664 | c.Fnc                   | -0.01764         | 0.03728           | 0.03736         | 0.472        | 0.63688             | 1.00              | 6                           |       |
| 23<br>24    | 665 | C.NPK                   | 0.09444          | 0.03743           | 0.03751         | 2.518        | 0.01181 *           | 1.00              | 6                           |       |
| 25          | 666 | c.Fnc:c.NPK             | -0.01041         | 0.07499           | 0.07514         | 0.139        | 0.88984             | 1.00              | 6                           |       |
| 26<br>27    | 667 | c.Fnc:z.AI              | 0.01691          | 0.07840           | 0.07856         | 0.215        | 0.82958             | 0.92              | 5                           |       |
| 28<br>29    | 668 | c.Fnc:z.pH              | 0.05260          | 0.08338           | 0.08354         | 0.630        | 0.52896             | 0.55              | 4                           |       |
| 30<br>31    | 669 | c.NPK:z.AI              | 0.28640          | 0.08119           | 0.08134         | 3.521        | 0.00043 ***         | 1.00              | 6                           |       |
| 32<br>33    | 670 | c.NPK:z.pH              | 0.12557          | 0.08425           | 0.08442         | 1.487        | 0.13690             | 0.69              | 4                           |       |
| 34<br>35    | 671 | c.Fnc:c.NPK:z.AI        | -0.48307         | 0.15742           | 0.15771         | 3.063        | 0.00219 **          | 0.92              | 5                           |       |
| 36<br>37    | 672 | c.Fnc:c.NPK:z.pH        | 0.36304          | 0.17535           | 0.17566         | 2.067        | 0.03876 *           | 0.34              | 2                           |       |
| 38          | 673 |                         |                  |                   |                 |              |                     |                   |                             |       |
| 39<br>40    | 674 | Signif. codes:          | 0 '***' 0.00     | 0.01              | '*' 0.05        | ··' 0.1      | ' ′ 1               |                   |                             |       |
| 41<br>42    |     |                         |                  |                   |                 |              |                     |                   |                             |       |
| 43<br>44    |     |                         |                  |                   |                 |              |                     |                   |                             | 33    |
| 45<br>46    |     |                         |                  |                   |                 |              |                     |                   |                             | 33    |
| 40<br>47    |     |                         |                  |                   |                 |              |                     |                   |                             |       |

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| 3<br>4      | 675 | SI Table 8. Respo | onse of foliar soo | dium to a factoria | al addition of e | lemental | nutrients (b | out not sodiu | im for the subse | t of 62 species present in control |
|-------------|-----|-------------------|--------------------|--------------------|------------------|----------|--------------|---------------|------------------|------------------------------------|
| 5           | 676 | plots and at leas | t 6 experimenta    | lly treated plots  | in the factorial | ΝΡΚμ exp | periment. T  | he regressio  | n table shows th | e conditional average values       |
| /<br>3<br>5 | 677 | across models, r  | elative importar   | nce values are sh  | own below the    | table.   |              |               |                  |                                    |
| 10          | 678 |                   |                    |                    |                  |          |              |               |                  |                                    |
| 11<br>12    | 679 |                   | Estimate S         | td. Error Ad       | justed SE z      | value    | Pr(> z )     |               | Importance       | Num models                         |
| 13<br>14    | 680 | (Intercept)       | 1.650418           | 0.156368           | 0.156525         | 10.544   | < 2e-16      | * * *         |                  |                                    |
| 15<br>16    | 681 | z.AI              | -0.713841          | 0.363376           | 0.363740         | 1.963    | 0.04970      | *             | 1.00             | 78                                 |
| 17<br>18    | 682 | z.pH              | -0.258238          | 0.060821           | 0.060879         | 4.242    | 2.22e-05     | * * *         | 1.00             | 78                                 |
| 19          | 683 | c.K               | -0.006721          | 0.025683           | 0.025709         | 0.261    | 0.79378      |               | 1.00             | 78                                 |
| 20<br>21    | 684 | c.N               | 0.073374           | 0.025576           | 0.025601         | 2.866    | 0.00416      | * *           | 1.00             | 78                                 |
| 22<br>23    | 685 | c.P               | 0.021680           | 0.025634           | 0.025660         | 0.845    | 0.39818      |               | 1.00             | 78                                 |
| 24<br>25    | 686 | c.K:c.N           | 0.044384           | 0.050890           | 0.050941         | 0.871    | 0.38360      |               | 0.76             | 57                                 |
| 26<br>27    | 687 | c.K:z.AI          | 0.303815           | 0.056138           | 0.056193         | 5.407    | 1.00e-07     | ***           | 1.00             | 78                                 |
| 28<br>29    | 688 | c.K:z.pH          | 0.138719           | 0.056652           | 0.056708         | 2.446    | 0.01444      | *             | 0.99             | 77                                 |
| 30<br>31    | 689 | c.N:c.P           | 0.075999           | 0.050956           | 0.051006         | 1.490    | 0.13622      |               | 0.72             | 57                                 |
| 32          | 690 | c.N:z.AI          | 0.081498           | 0.056873           | 0.056926         | 1.432    | 0.15225      |               | 0.89             | 68                                 |
| 33<br>34    | 691 | c.N:z.pH          | -0.112398          | 0.056589           | 0.056645         | 1.984    | 0.04723      | *             | 0.89             | 67                                 |
| 35<br>36    | 692 | c.P:z.AI          | 0.131357           | 0.053185           | 0.053237         | 2.467    | 0.01361      | *             | 0.99             | 77                                 |
| 37<br>38    | 693 | c.K:c.N:z.AI      | 0.247463           | 0.109837           | 0.109940         | 2.251    | 0.02439      | *             | 0.70             | 50                                 |
| 39<br>40    | 694 | c.P:z.pH          | -0.018531          | 0.057573           | 0.057628         | 0.322    | 0.74778      |               | 0.43             | 38                                 |
| 41<br>42    | 695 | c.N:c.P:z.pH      | -0.213430          | 0.106099           | 0.106205         | 2.010    | 0.04447      | *             | 0.31             | 25                                 |

| 1<br>2   |     |                              |                 |                   |                 |             |           |           |                |                         |          |
|----------|-----|------------------------------|-----------------|-------------------|-----------------|-------------|-----------|-----------|----------------|-------------------------|----------|
| 3<br>4   | 696 | c.K:c.N:z.pH 0.142           | 729 0.11        | .13412 0.1        | .13525 1.       | .257 0.     | 20866     |           | 0.27           | 20                      |          |
| 5        | 697 | c.K:c.P 0.051                | 492 0.05        | 50758 0.0         | 50809 1.        | .013 0.     | 31085     |           | 0.40           | 38                      |          |
| 6<br>7   | 698 | c.N:c.P:z.AI 0.023           | 955 0.11        | 0.1               | .12340 0.       | .213 0.     | 83115     |           | 0.12           | 12                      |          |
| 8<br>9   | 699 | c.K:c.P:z.AI 0.076           | 209 0.10        | 0.1582 0.1        | .01683 0.       | .749 0.     | 45357     |           | 0.07           | 7                       |          |
| 10<br>11 | 700 | c.K:c.N:c.P -0.051           | 459 0.10        | 02002 0.1         | .02104 0.       | .504 0.     | 61427     |           | 0.03           | 4                       |          |
| 12<br>13 | 701 | c.K:c.P:z.pH -0.013          | 211 0.10        | 0.1               | .03284 0.       | .128 0.     | 89822     |           | 0.01           | 2                       |          |
| 14       | 702 |                              |                 |                   |                 |             |           |           |                |                         |          |
| 15<br>16 | 703 | Signif. codes: 0 '           | ***' 0.001      | · ** · 0.01       | ·*' 0.05        | .' 0.1      | ' ' 1     |           |                |                         |          |
| 17<br>18 | 704 |                              |                 |                   |                 |             |           |           |                |                         |          |
| 19<br>20 | 705 |                              |                 |                   |                 |             |           |           |                |                         |          |
| 21<br>22 |     |                              |                 |                   |                 |             |           |           |                |                         |          |
| 23       | 706 | SI Tables 9 & 10. Effects of | of foliar sodiu | im on changes     | in species ab   | undance     | n respons | e to a ch | anging environ | ment:                   |          |
| 24<br>25 | 707 | SI Table 9. Response of p    | ot scale cove   | r of focal specie | es as a functio | on of folia |           |           | -              | ntal manipulation of he | rbivores |
| 26<br>27 | 708 | and nutrients. (N=153 spe    | ecies).         |                   |                 |             |           |           |                |                         |          |
| 28<br>29 | 709 |                              |                 |                   |                 |             |           |           |                |                         |          |
| 30<br>31 | 710 | (conditional average)        |                 |                   |                 |             |           |           |                |                         |          |
| 32       | 711 |                              | Estimate        | Std. Error A      | djusted SE      | z value     | Pr(> z )  |           | Importance     | Num models              |          |
| 33<br>34 | 712 | (Intercept)                  | 1.17346         | 0.05116           | 0.05125         | 22.898      | < 2e-16   | ***       |                |                         |          |
| 35<br>36 | 713 | z.lf.na.lg                   | -0.02289        | 0.04719           | 0.04727         | 0.484       | 0.6282    |           | 1.00           | 5                       |          |
| 37<br>38 | 714 | c.Fnc                        | -0.01806        | 0.02609           | 0.02613         | 0.691       | 0.4895    |           | 1.00           | 5                       |          |
| 39       | 715 | c.NPK                        | 0.06037         | 0.02624           | 0.02628         | 2.297       | 0.0216    | *         | 1.00           | 5                       |          |
| 40<br>41 | 716 | c.Fnc:c.NPK                  | -0.06492        | 0.05079           | 0.05087         | 1.276       | 0.2019    |           | 0.61           | 3                       |          |
| 42<br>43 |     |                              |                 |                   |                 |             |           |           |                |                         |          |
| 44       |     |                              |                 |                   |                 |             |           |           |                |                         | 35       |
| 45       |     |                              |                 |                   |                 |             |           |           |                |                         | 5.       |

| 1<br>2   |     |                   |              |               |                  |             |             |             |                 |               |                 |               |     |
|----------|-----|-------------------|--------------|---------------|------------------|-------------|-------------|-------------|-----------------|---------------|-----------------|---------------|-----|
| 3        | 717 | c.Fnc:z.lf.na.    | lg           | 0.06108       | 0.05457          | 0.05465     | 1.118       | 0.2637      | 0.56            | 3             |                 |               |     |
| 4<br>5   | 718 | c.NPK:z.lf.na.    | lg ·         | -0.23515      | 0.05410          | 0.05418     | 4.340       | 1.43e-05    | *** 1.00        | 5             |                 |               |     |
| 6<br>7   | 719 | c.Fnc:c.NPK:z.    | lf.na.lg     | 0.20691       | 0.10431          | 0.10448     | 1.980       | 0.0477      | * 0.30          | 1             |                 |               |     |
| 8        | 720 |                   |              |               |                  |             |             |             |                 |               |                 |               |     |
| 9<br>10  | 721 | Signif. codes:    | 0 '***'      | 0.001 '**'    | 0.01 '*' 0.0     | 05 '.' 0.   | 1 ' ' 1     |             |                 |               |                 |               |     |
| 11<br>12 | 722 |                   |              |               |                  |             |             |             |                 |               |                 |               |     |
| 13<br>14 | 723 |                   |              |               |                  |             |             |             |                 |               |                 |               |     |
| 15<br>16 | 724 | SI Table 10. Resp | onse of plo  | t scale cover | of focal species | as a functi | ion of foli | ar sodium i | n response to a | factorial add | lition of eleme | ental nutrier | nts |
| 17<br>18 | 725 | (but not sodium). | . The regres | sion table sh | ows the condit   | ional avera | ige values  | across mo   | dels. (N=179 sp | ecies)        |                 |               |     |
| 19<br>20 | 726 |                   |              |               |                  |             |             |             |                 |               |                 |               |     |
| 21<br>22 | 727 | (conditional a    | verage)      |               |                  |             |             |             |                 |               |                 |               |     |
| 23<br>24 | 728 |                   | Estimate     | Std. Error    | Adjusted SE      | z value :   | Pr(> z )    |             | Importance      | Num model     | ls              |               |     |
| 25<br>26 | 729 | (Intercept)       | 1.11276      | 0.05274       | 0.05278          | 21.083      | < 2e-16     | ***         |                 |               |                 |               |     |
| 27       | 730 | z.lf.na.lg        | -0.10085     | 0.03770       | 0.03773          | 2.673       | 0.00752     | **          | 1.00            | 11            |                 |               |     |
| 28<br>29 | 731 | c.K               | 0.02774      | 0.01732       | 0.01734          | 1.600       | 0.10951     |             | 1.00            | 11            |                 |               |     |
| 30<br>31 | 732 | c.N               | 0.02403      | 0.01748       | 0.01749          | 1.374       | 0.16944     |             | 1.00            | 11            |                 |               |     |
| 32       | 733 | c.P               | 0.03991      | 0.01723       | 0.01724          | 2.315       | 0.02063     | *           | 1.00            | 11            |                 |               |     |
| 33<br>34 | 734 | c.K:c.P           | 0.08592      | 0.03427       | 0.03429          | 2.505       | 0.01223     | *           | 1.00            | 11            |                 |               |     |
| 35<br>36 | 735 | c.K:z.lf.na.lg    | -0.06344     | 0.03453       | 0.03455          | 1.836       | 0.06639     |             | 0.71            | 7             |                 |               |     |
| 37       | 736 | c.N:z.lf.na.lg    | -0.06115     | 0.03468       | 0.03471          | 1.762       | 0.07810     | •           | 0.68            | 7             |                 |               |     |
| 38<br>39 | 737 | c.P:z.lf.na.lg    | -0.10016     | 0.03469       | 0.03471          | 2.885       | 0.00391     | * *         | 1.00            | 11            |                 |               |     |
| 40<br>41 | 738 | c.K:c.N           | 0.02331      | 0.03446       | 0.03448          | 0.676       | 0.49899     |             | 0.26            | 4             |                 |               |     |
| 42       |     |                   |              |               |                  |             |             |             |                 |               |                 |               |     |
| 43<br>44 |     |                   |              |               |                  |             |             |             |                 |               |                 |               | 36  |
| 45<br>46 |     |                   |              |               |                  |             |             |             |                 |               |                 |               |     |

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|--------|------------|---------|
| 1<br>2 |            |         |
| 3      | 739        | c.N:c.P |
| 4<br>5 | 740        |         |
| 6<br>7 | 741        | Signif. |
| 8      | 742        |         |

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| 739 | c.N:c.P         | 0.01356         | 0.03428         | 0.03430        | 0.395            | 0.69253     | 0.23        | 4           |                                       |
|-----|-----------------|-----------------|-----------------|----------------|------------------|-------------|-------------|-------------|---------------------------------------|
| 740 |                 |                 |                 |                |                  |             |             |             |                                       |
| 741 | Signif. code    | s: 0 '***' (    | 0.001 '**' 0    | .01 '*' 0.05   | ' <b>.</b> ' 0.1 | ''1         |             |             |                                       |
| 742 |                 |                 |                 |                |                  |             |             |             |                                       |
| 743 |                 |                 |                 |                |                  |             |             |             |                                       |
| 744 |                 |                 |                 |                |                  |             |             |             |                                       |
| 745 | SI Table 11. Au | thor contributi | ons and site-le | evel acknowled | lgments ta       | able.       |             |             |                                       |
|     |                 |                 | Developed       |                | 0                | Contributed |             | Nutrient    |                                       |
|     |                 | Contributed     | research        | Analyzed       | Wrote            | to paper    | Site        | Network     | Site-level acknowledgments            |
|     | Name            | samples         | question        | data           | paper            | writing     | coordinator | coordinator | (funding, access, etc)                |
|     |                 |                 |                 |                |                  |             |             |             | i i i i i i i i i i i i i i i i i i i |

|               | Continuation |          | /    |       |         | 0.00        |             |                        |
|---------------|--------------|----------|------|-------|---------|-------------|-------------|------------------------|
| Name          | samples      | question | data | paper | writing | coordinator | coordinator | (funding, access, etc) |
| Borer,        | x            | х        | x    | x     | 4       | x           | x           |                        |
| Elizabeth T.  |              |          |      |       |         | 0           |             |                        |
| Lind, Eric M. | х            | х        | x    |       | x       |             | ×           |                        |
| Seabloom,     | x            |          | x    |       | х       | x           | х           |                        |
| Eric W.       |              |          |      |       |         |             |             |                        |
| Firn,         | x            |          |      |       | x       | x           |             |                        |
| Jennifer      |              |          |      |       |         |             |             |                        |

| Anderson,    |   |   |   | х  | х             |                              |
|--------------|---|---|---|----|---------------|------------------------------|
| T. Michael   |   |   |   |    |               |                              |
| Bakker,      |   |   |   | x  | x             |                              |
| Elisabeth S. |   |   |   |    |               |                              |
| Biederman,   | Х | ~ |   | x  | x             |                              |
| Lori         |   |   |   |    |               |                              |
| La Pierre,   | Х |   | C | x  | x             | Funding: Konza Prairie LTER  |
| Kimberly J   |   |   | 0 |    |               |                              |
| MacDougall,  | x |   |   | x  | x             | Funding: NSERC Discovery     |
| Andrew S     |   |   |   | -4 |               | Grant; In-kind site support: |
|              |   |   |   |    | $O_{h}$       | Nature Conservancy of        |
|              |   |   |   |    | $\neg \gamma$ | Canada; sampling processing  |
|              |   |   |   |    |               | Carly Ziter                  |
| Joslin       | x |   |   | x  | x             |                              |
| Moore        |   |   |   |    |               |                              |

| 2                          |                |  |     |                        |               |             |            |  |  |  |    |
|----------------------------|----------------|--|-----|------------------------|---------------|-------------|------------|--|--|--|----|
| 3<br>4                     |                | Risch, Anita   | х   |                        |               | х           | x          |  |  |  |    |
| 5<br>6<br>7                |                | С.   |     |                        |               |             |            |  |  |  |    |
| 7<br>8<br>9                |                | Schütz,  | x   |                        |               | х           | x          |  |  |  |    |
| 9<br>10<br>11              |                | Martin   |     |                        |               |             |            |  |  |  |    |
| 12<br>13                   |                | Stevens,   | x   | $\sim$                 |               | х           | x          |  |  |  |    |
| 14<br>15<br>16             |                | Carly J.   |     |                        |               |             |            |  |  |  |    |
| 17<br>18                   | 746            |  |     |                        |               |             |            |  |  |  |    |
| 19<br>20                   | 747            |  |     |                        |               |             |            |  |  |  |    |
| 21<br>22<br>23<br>24<br>25 | 748            | SI Table 12. All data contributors listed by site; site names match those in SI Table 1. Their effort in providing samples was key to this work. |     |                        |               |             |            |  |  |  |    |
|                            |                | Site Pl  |     | Site name(s) from whi  | ich trait dat | a were co   | ontributed |  |  |  |    |
| 26<br>27                   |                | Peter Adler  |     | Sheep Experimental St  | tation        |             |            |  |  |  |    |
| 28<br>29<br>30             |                | Jonathan Bakl  | ker | Smith Prairie          |               |             |            |  |  |  |    |
| 31<br>32                   |                | Lori Biederma  | in  | Chichaqua Bottoms      |               |             |            |  |  |  |    |
| 33<br>34                   |                | Dana Blumenthal  |     | Shortgrass Steppe LTER |               |             |            |  |  |  |    |
| 35<br>36<br>37             |                |  |     | .TER), Sierra F        | oothills REC  | Hopland REC | C, Lookout |  |  |  |    |
| 38<br>39                   | (Andrews LTER) |  |     |                        |               |             |            |  |  |  |    |
| 40<br>41                   |                |  |     |                        |               |             |            |  |  |  |    |
| 42<br>43                   |                |  |     |                        |               |             |            |  |  |  |    |
| 44<br>45                   |                |  |     |                        |               |             |            |  |  |  | 39 |
| 46<br>47                   |                |  |     |                        |               |             |            |  |  |  |    |

| 1                                    | 0   |  |
|--------------------------------------|---|--|
| 1<br>1<br>1<br>1<br>2                | 2345678901                                |  |
| 2<br>2<br>2<br>2<br>2<br>2<br>2<br>2 | 1<br>2<br>3<br>4<br>5<br>6<br>7           |  |
| 2<br>2<br>3<br>3                     | 8<br>9<br>0<br>1                          |  |
| 3<br>3                               | 2<br>3<br>4<br>5<br>6<br>7<br>8<br>9<br>0 |  |
| 4<br>4<br>4<br>4                     | -1<br>-2<br>-3<br>-4<br>-5                |  |

| Cynthia Brown             | Shortgrass Steppe LTER   |
|---------------------------|--|
| Miguel Bugalho            | Companhia das Lezirias   |
| Maria Caldeira            | Companhia das Lezirias   |
| Elsa Cleland              | Elliott Chaparral  |
| Kendi Davies              | Boulder South Campus   |
| Jennifer Firn             | Burrawan   |
| Daniel Gruner             | Sagehen Creek UCNRS  |
| Sabine Güsewell           | Fruebuel   |
| W. Stanley Harpole        | Hopland REC, Chichaqua Bottoms, Mclaughlin UCNRS, Sierra Foothills REC |
| Yann Hautier              | Fruebuel   |
| Andy Hector               | Fruebuel   |
| Janneke Hille Ris Lambers | Smith Prairie  |
| Kirsten Hofmockel         | Chichaqua Bottoms  |
| Julia Klein               | Shortgrass Steppe LTER   |
| Alan Knapp                | Shortgrass Steppe LTER   |
|                           |  |

| 1              |  |  |  |  |
|----------------|--|--|--|--|
| 2<br>3<br>4    | Kimberly La Pierre   | Konza LTER, Saline Experimental Range  |  |  |
| 5<br>6<br>7    | Andrew MacDougall  | Cowichan   |  |  |
| 8<br>9         | Brett Melbourne  | Boulder South Campus   |  |  |
| 10<br>11       | Charles Mitchell   | Duke Forest  |  |  |
| 12<br>13<br>14 | Joslin Moore   | Bogong   |  |  |
| 15<br>16       | John Morgan  | Bogong, Kinypanial   |  |  |
| 17<br>18<br>19 | Suzanne Prober   | Mt. Caroline   |  |  |
| 20<br>21       | Anita Risch     Val Mustair       Martin Schuetz     Val Mustair |  |  |  |
| 22<br>23<br>24 | Martin Schuetz   | Val Mustair  |  |  |
| 24<br>25<br>26 | Eric Seabloom  | Hopland REC, Lookout (Andrews LTER), Mclaughlin UCNRS, Bunchgrass (Andrews LTER), Sierra |  |  |
| 27<br>28<br>29 |  | Foothills REC  |  |  |
| 30<br>31       | Melinda Smith  | Konza LTER, Saline Experimental Range  |  |  |
| 32<br>33       | Carly Stevens  | Lancaster  |  |  |
| 34<br>35<br>36 | Lauren Sullivan  | Chichaqua Bottoms  |  |  |
| 37<br>38       | Peter Wragg  | Mt Gilboa, Summerveld  |  |  |
| 39<br>40<br>41 |  |  |  |  |
| 41<br>42       |  |  |  |  |

| 1<br>2   |     |               |                     |
|--|-----|---------------|---------------------|
| 3<br>4   |     | Justin Wright | Duke Forest         |
| 5<br>6<br>7<br>8<br>9<br>10<br>11<br>12<br>13  |     | Louie Yang    | Sagehen Creek UCNRS |
|  | 749 |               |                     |
|  | 750 |               |                     |
|  | 751 |               |                     |
| 14<br>15   | 752 |               |                     |
| 17   | 753 |               |                     |
| <ol> <li>16</li> <li>17</li> <li>18</li> <li>19</li> <li>20</li> <li>21</li> <li>22</li> <li>23</li> <li>24</li> <li>25</li> <li>26</li> <li>27</li> <li>28</li> <li>29</li> <li>30</li> <li>31</li> <li>32</li> <li>33</li> <li>34</li> <li>35</li> <li>36</li> <li>37</li> <li>38</li> <li>39</li> <li>40</li> <li>41</li> <li>42</li> <li>43</li> <li>44</li> <li>45</li> <li>46</li> <li>47</li> </ol> |     |               |                     |



SI Figure 3. Site-level soil pH declines as a function of site-level water availability (MAP/PET); this relationship does not vary as a function of
 distance from coast. Coastal and Inland are divided at 100km from a coast. The Lancaster site in the UK, shown with a black circle and cross-hairs
 in this figure, falls along this line, but has very high coastal sodium influence in its precipitation, leading to exceptionally high site-level sodium
 (see main text).