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2	Diversity-induced plant history and soil history effects
3	modulate plant responses to global change
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15 Abstract

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17 Global change has dramatic impacts on grassland diversity. However, little is known about how fast species can adapt to these changes and how this affects their responses to global change. 18 19 To close this gap, we performed a common garden experiment testing whether plant responses to global change are influenced by the selection history of the plants and the conditioning 20 21 history of soil at different levels of plant diversity. Therefore, we collected seeds and took soil samples from 14-year old plant communities of a biodiversity experiment. Offspring of plants 22 23 from low- and high-diversity communities were either grown in their own soil or in soil of a 24 different community, and were either exposed to drought, increased nitrogen input, or a combination of both. Results show that, under nitrogen addition, offspring of plants selected at 25 high diversity produced more biomass than those selected at low diversity, while drought 26 27 neutralized differences in biomass production. Moreover, under the influence of global change drivers, mainly soil, and to a lesser extent plant history, influenced the expression of plant traits. 28 29 Our results show that plant diversity modulates plant-soil interactions and growth strategies of plants, which feedback on the eco-evolutionary pathways of the plants and thus their responses 30 31 to global change.

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Key words: plant-soil interaction, plant-soil feedback, drought, fertilization, micro-evolution,
eco-evolutionary feedback, nutrient enrichment, climate change

35 Introduction

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Human activities, such as the combustion of fossil fuels and the intensification of agriculture, are leading to global environmental changes, causing increased air temperatures, altered precipitation patterns, and rising amounts of nitrogen to ecosystems (IPCC, Pörtner et al. 2021). The consequences are more frequent extreme weather events such as droughts (Dai et al. 2018) and a growing accumulation of nitrogen in the soils (Holland et al. 2005). Both, drought and increased nitrogen input, in turn, further influence ecosystems and climatic conditions; hence, they are known as major global change drivers (Sage 2020).

44 Some of the most tremendous negative effects of global change are changes in 45 ecological communities (Dornelas et al. 2014) and the extinction of species (Sage 2020), whereby plant species are particularly concerned due to their low mobility, with drastic 46 47 consequences for the functioning of ecosystems. Studies in grassland biodiversity experiment have shown that low- and high-diversity plant communities significantly differ in their 48 productivity and stability (Isbell et al. 2015; Marquard et al. 2009; Tilman et al. 2006). Low-49 diversity communities were shown to lose productivity over time, while high-diversity 50 51 communities are more stable, so that plant diversity-productivity relationships become more 52 positive over time (Cardinale et al. 2007; Meyer et al. 2016; Reich et al. 2012). The different development of plant-soil and plant-plant interactions at low and high diversity are assumed to 53 be the important drivers of this strengthening biodiversity effect (Eisenhauer et al. 2019; Thakur 54 55 et al. 2021). At low plant diversity, an accumulation of soil-borne pathogens might be responsible for lower plant community productivity (Mommer et al. 2018; Thakur et al. 2021), 56 57 while in high-diversity communities, complementarity effects among plants inhibit such negative processes, causing a higher productivity of these plant communities (Cardinale et al. 58 2007; Reich et al. 2012). Consequently, these findings raise the question, whether populations 59 60 of the same plant species develop differently over time when growing at high or low diversity due to differences in eco-evolutionary feedbacks (Bailey et al. 2006; Linhart 1988; Post and Palkovacs 2009; terHorst and Zee 2016). Indeed, there is empirical evidence that plant individuals at high diversity are selected for greater niche complementarity among species leading to a more complete use of available resources (Zuppinger-Dingley et al. 2014). At low plant diversity, however, the accumulation of soil-borne pathogens may cause persistent species to adapt to this increase by producing more defense compounds, so that over time selection favors individuals that invest more in defense and less in growth (Eisenhauer et al. 2019).

Taken together, low and high plant diversity may differently affect eco-evolutionary 68 feedbacks and thus the microevolution of plants, which could have the effect that plants selected 69 70 at low diversity respond differently to global change drivers than plants selected at high diversity, due to differences in the phenotype and/or growth strategies. In a previous transplant 71 experiment (Lipowsky et al. 2011), it was shown that some of the studied grassland species 72 73 showed difference in their phenotype depending on plant history (monoculture or mixture) and soil environment (home or away soil). For example, it was shown that the species *Cirsium* 74 oleraceum (L.) Scop. had a higher number of leaves, when originated from mixture 75 communities, and were taller, when grown in home soil. Furthermore, several greenhouse 76 77 studies showed similar results, i.e., that plants selected at low or high diversity or grown with 78 "own" or different soil biota vary in their productivity and trait expression (Hahl et al. 2020; van Moorsel et al. 2018b; Zuppinger-Dingley et al. 2014). Such diversity-induced differences 79 in the phenotype could lead to different responses of plants to global change drivers. For 80 81 example, it is possible that, due to differences in leaf number or root structure, plants selected at low diversity have a lower resistance against drought than plants selected at high diversity. 82 Such changes would contribute to a faster extinction of species, which makes research into 83 these processes an essential frontier. 84

In summary, differences in plant-plant and plant-soil interactions at low and high diversity may lead to differences in eco-evolutionary feedbacks; however, little is known about

how rapidly and pervasively these differences occur (terHorst and Zee 2016). Moreover, it is 87 88 not known whether these differences affect the response of plants to global change drivers (Pugnaire et al. 2019), such as drought, nitrogen input or a combination of both, which is 89 assumed to be a common scenario in the future (Craven et al. 2016; Sage 2020). To address 90 91 these knowledge gaps, we performed a common garden experiment using plant and soil material from a long-running biodiversity experiment (Jena Experiment). For our study, we collected 92 93 seeds of four grass species and took samples of soil biota (soil samples), which both had been selected for 14 years, either at low or high diversity (communities with two or six plant species 94 95 and different plant species composition). Plants were grown either in soil inoculated with their 96 home soil biota, i.e. soil biota of the community, where the seeds had been collected, or in soil 97 inoculated with away soil biota, i.e. with soil biota of a different plant community (differing in plant diversity or composition). The aim of the study was to test, whether plant history (origin 98 99 of plants), soil history (origin of soil biota), and soil treatment (home/away) influence the response of the plants to global change. Therefore, plants were either non-treated (control), or 100 exposed to drought, increased nitrogen input, or a combination of both, drought and nitrogen 101 input, in a full factorial design. We hypothesized that 102

103 (I) plant and soil communities develop differently at low and high diversity over time. 104 Therefore, we expected that offspring of plants (Ia) selected at high diversity generally shows higher biomass production compared to offspring selected at low diversity in the 105 control, and that plants under control conditions produce more biomass (Ib) in home than 106 107 in away soil and (Ic) in high-diversity than in low-diversity soil. Further, (Id) we supposed effects of plant history, soil history, and soil treatment on trait expression of control 108 109 plants. For example, we expected that offspring of plants from high-diversity communities show higher values for traits related to relative growth rates (e.g., leaf 110 greenness, specific leaf area) and nutrient economy (e.g., shoot nitrogen concentrations) 111 than offspring of plants from low-diversity communities. 112

global change drivers have a strong impact on biomass production and trait expression of 113 (II) 114 plants. We expected that (IIa) drought reduces, and (IIb) nitrogen input increases plant biomass, while (IIc) the combination of both global change drivers has no impact on plant 115 biomass production, because drought and nitrogen input compensate each other's impact. 116 (III) because of different development of plants and soil communities at low and high 117 diversity, offspring of plants (IIIa) selected at different diversity and grown in different 118 119 soil (IIIb: home vs. away soil, IIIc: soil from low- vs. high-diversity communities) respond differently to global change drivers regarding performance and trait expression. 120

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122 **Results**

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124 *Hypothesis 1: offspring of plants selected at different diversity and grown in different soil (high*

125 vs. low diversity, home vs. away) show differences in productivity and trait expression

126 Biomass production

Plants grown in soil of six-species communities tended to produce more root biomass 127 than plants in soil of two-species communities in the control (Table 3; Fig. 2). At species-level, 128 129 A. elatius produced more root biomass and had higher root-shoot ratio, and D. glomerata 130 produced more shoot and total biomass in soil of six-species than two-species communities (Fig. 2, 3a; Appendix S3: Table S8, S10). The other two species, P. trivialis and A. pratensis 131 did not differ significantly in biomass production dependent on soil or plant history (Fig. 2, 3a; 132 133 Appendix S3: Table S9, S11). Initial shoot number showed no influence on later biomass production except for shoot biomass of D. glomerata and root biomass of A. elatius, which, 134 however, did not change the general patterns. 135

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137 Plant traits and pathogen infestation

Legacy treatments had no consistent effects across the four species on the expression of 138 139 shoot, leaf, or root traits in the control (Appendix S3: Table S1). At species-level, legacy 140 treatments did not affect trait expression in A. elatius (Fig. 3a; Appendix S3: Table S1). Plants of A. pratensis were taller in home than in away-different soil and had thicker roots (higher root 141 diameter) in six- than in two-species soil (Fig. 3a; Appendix S3: Table S3). Plants of D. 142 *glomerata* had higher leaf greenness and stomatal conductance, when seeds originated from 143 two-species communities (Fig. 3a; Appendix S3: Table S4). Plants of P. trivialis had lower 144 shoot nitrogen concentration and root diameter, and higher SRL in home soil than in away soil 145 (Fig. 3a; Appendix S3: Table S5). 146

We found a low pathogen infestation of *A. elatius* and *A. pratensis* ($0.8\% \pm 1.9\%$ (SD) and $0.1\% \pm 0.5\%$, respectively), mainly by the rust species *Puccinia graminis* Pers. and *Puccinia coronata* Corda. Plants of *D. glomerata* and *P. trivialis*, in contrast, were strongly infested by the mildew *Blumeria graminis* (DC.) Speer ($3.1\% \pm 4.2\%$ and $8.6\% \pm 16.5\%$, respectively). Regarding legacy treatments, *D. glomerata* plants had a lower infestation when grown in home soil than in away soil, while mildew infestation of *P. trivialis* plants did not differ between legacy treatments (Fig 3a; Appendix S3: Table S6).

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Hypothesis 2: global change drivers have a strong impact on biomass production and trait
expression.

157 Biomass production

Overall, global change drivers had a strong impact on almost all response variables (Table 2; Fig. 3b-d; Appendix S2: Table S1-9). Compared to control plants, drought reduced shoot biomass production, which was found across all study species and at species-level (Fig. 4a, d). In contrast, drought did not have consistent effects on root biomass (Fig. 4a, d). Drought had positive impact on root biomass of *A. elatius* and *D. glomerata*, while root biomass of *A. pratensis* decreased under drought and did not change significantly in *P. trivialis* (Fig. 4d).

Total biomass production was decreased, when plants were exposed to drought (Fig. 4a, d) except for *D. glomerata*, where it was not different from the control (Fig. 4d). Root-shoot ratios increased under drought (Fig. 4a, d), which was found for all species except for *P. trivialis* (no significant change; Fig. 4d).

168 Nitrogen input increased shoot, root, and thus also total biomass across the four species
169 (Fig. 4b) as well as in separate analyses of *A. elatius* and *A. pratensis* (Fig. 4e). Plants of *D.*170 *glomerata* and *P. trivialis* did not change in root biomass when fertilized (Fig. 3e). Nitrogen
171 input caused a decrease in root-shoot ratio in all species (Fig. 4a, e).

When plants were treated with both global change drivers in combination, the negative 172 173 impact of drought on shoot biomass was cancelled out by the positive impact of nitrogen input 174 leading to an overall slight increase of shoot biomass (compared to control plants) that was also significant at the species-level except for A. elatius (Fig. 4c, f). Consistent with this, the positive 175 176 impact of nitrogen input on root biomass was also cancelled out by drought when plants were treated with both global change drivers, i.e. control plants and plants treated with both global 177 change drivers did not differ in root biomass production, across all study species (Fig. 4c). At 178 species-level, the combination of both global change drivers had an additive effect on root 179 180 biomass production of A. elatius and D. glomerata, i.e. plants of both species showed highest 181 root biomass when treated with both global change drivers (Fig. 4f). In A. pratensis and P. trivialis, both global change drivers in combination decreased root biomass production (Fig. 182 3f). Taken together, the combination of both global change drivers led to a slight increase in 183 184 total biomass production, across all study species and for the high-productive species A. elatius and D. glomerata, while plants of the low-productive species A. pratensis and P. trivialis had a 185 similar total biomass production as in the control (Fig. 4c, f). Root-shoot ratios were as low as 186 in fertilized plants, across all species and in P. trivialis (Fig. 4c, f). Plants of A. elatius and D. 187 glomerata increased root-shoot ratios, similar to plants under drought (Fig. 4f). In contrast, A. 188

pratensis strongly decreased root-shoot ratios resulting in the lowest values compared to theother treatments (Fig. 4f).

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192 Plant traits and pathogen infestation

Across all study species, drought did not significantly alter growth height, but nitrogen 193 input increased height (Appendix S2: Fig. S1). When treated with both global change drivers, 194 195 drought canceled out the positive nitrogen input effect, leading to similar height of plants treated with both global change drivers and control plants. Further, drought and nitrogen input 196 increased shoot nitrogen concentrations and leaf greenness, with additive effects when both 197 198 global change drivers were applied together (Appendix S2: Fig. S1). Drought did not influence 199 LDMC and SLA, while nitrogen input decreased LDMC and increased SLA (Appendix S2: Fig. S2). When treated with both global change drivers, drought mitigated the decrease of 200 201 LDMC under nitrogen input, while the increase of SLA under nitrogen input did not change with drought (Appendix S2: Fig. S2). Stomatal conductance was increased, when plants were 202 203 treated with drought, but did not change when fertilized irrespective of the drought treatment (Appendix S2: Fig. S2). In terms of root traits, we found a decrease of RLD under drought 204 205 (irrespective of fertilization) and an increase in root diameter under nitrogen input (irrespective 206 of drought; Appendix S2: Fig. S3). Results of species-specific trait expression changes under 207 global change drivers can be found in Figure 3b-d and Appendix S2.

In *D. glomerata*, mildew infestation remained unchanged when treated with drought, but increased with nitrogen input. When treated with both global change drivers, mildew infestation was as high as in fertilized plants (Fig. 3b-d; Appendix S2: Fig. S4). In *P. trivialis*, mildew infestation was increased under drought and when fertilized, while the combination of both global change drivers led to the highest mildew infestation (Fig. 3b-d; Appendix S2: Fig. S4).

215 *Hypothesis 3: offspring of plants selected at different diversity and grown in different soil (high*

vs. low diversity, home vs. away) respond differently to global change drivers

217 Biomass production

Plants from two- and six-species communities did not differ in shoot biomass production
when treated with drought, but plants from six-species communities treated with drought tended
to produce more root biomass than plants from two-species communities across all study
species (Table 3; Fig. 5a, d, g). At species-level, we found no significant effects of legacy
treatments under drought (Fig. 3b; Appendix S3: Table S8-S11).

When plants were fertilized, we found an impact of plant history across all study species: 223 224 fertilized plants originated from six-species communities had a higher root and total biomass 225 production than plants from two-species communities (Table 3; Fig. 5a, g). This was also found in D. glomerata plants, which tended to produce more shoot and total biomass when originated 226 227 from six-species communities (Fig. 3c; Appendix S3: Table S10). In A. elatius, total biomass production of fertilized plants was significantly higher (and shoot biomass marginally 228 significantly higher), when plants were grown in home and away-same than in away-different 229 soil (Fig. 3c; Appendix S3: Table S8). In A. pratensis, fertilized plants grown in two-species 230 231 community soil tended to produce more total biomass than in six-species community soil (Fig. 232 3c; Appendix S3: Table S9), while fertilized *P. trivialis* showed no significant differences (Fig. 3c; Appendix S3: Table S11). 233

When plants were treated with both global change drivers, the effects of nitrogen input were cancelled out or changed by drought, i.e. there was no significant impact of legacy treatments on biomass production across all study species and for *A. elatius* (Table 3; Fig. 3d; Appendix S3: Table S8). In *D. glomerata*, the significant influence of plant history disappeared, but plants in home and away-different soil showed higher root-shoot ratios than plants in awaysame soil (Fig. 3d; Appendix S3: Table S10). Plants of *P. trivialis* treated with both global change drivers tended to have higher root biomass and root-shoot ratios when grown in home

than in away-same soil (Fig. 3d; Appendix S3: Table S11). In contrast to the overall trend, A. 241 242 pratensis was the only species which showed a similar response to nitrogen input and treatment with both global change drivers: the biomass production was higher in two- than in six-species 243 community soil (for both global change drivers: significant higher root biomass and root-shoot 244 245 ratios; Fig. 3d; Appendix S3: Table S9).

- 246
- 247 Plant traits and pathogen infestation

Shoot nitrogen concentration was not influenced by plant or soil history when treated 248 with drought, but fertilized plants in six-species soil had higher shoot nitrogen concentrations 249 than in two-species soil (soil history effect; Appendix S3: Table S1; Fig. S1). Moreover, 250 fertilized plants had lower shoot nitrogen concentrations in home than in away-different soil 251 252 (soil treatment effect; Appendix S3: Table S1; Fig. S1). When plants were treated with both global change drivers, the nitrogen input effect on soil history was cancelled out by drought, 253 while the impact of soil treatment did not: plants in home soil still had lower shoot nitrogen 254 255 concentration than plants in away soil (Appendix S3: Table S1; Fig. S1). Other plant traits 256 (growth height, leaf greenness, LDMC, SLA, stomatal conductance, root traits) did not significantly differ depending on legacy treatments when plants were treated with nitrogen or 257 258 drought (Appendix S3: Table S1). At species level, we found a large number of different responses depending on legacy treatments and type of global change driver, which can be found 259 in Figure 3b-d and Appendix S3. 260

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Mildew infestation of *D. glomerata* plants exposed to drought was higher in home than in away soil, while this drought effect was cancelled out by nitrogen input (Appendix S3: Table 262 263 S6). Mildew infestation of *P. trivialis* plants was not significantly influenced by legacy treatments, neither with nor without global change drivers (Appendix S3: Table S6). 264

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Tables 266

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Table 1 Summary list of response variables and experimental factors of the common garden

269 experiment

Variable	Abbreviation	Unit	Description
Response variables			
Biomass production			
Total biomass	Total Bm	g _{total}	Shoot and root biomass per pot
Shoot mass	Shoot Bm	g _{shoot}	Shoot biomass per pot
Root mass	Root Bm	g _{root}	Root biomass per pot
Root-shoot ratio Aboveground traits	-	$g_{root} g_{shoot}^{-1}$	Root biomass divided by shoot biomass per pot
Growth height	-	cm	Stretched shoot length of longest vegetative shoot ^a
Shoot nitrogen concentration	N _{Shoot}	mg N g _{shoot} ⁻¹	Nitrogen mass per dry shoot mass
Leaf greenness	-	-	Unitless estimate of leaf chlorophyll concentration ^a
Specific leaf area	SLA	mm _{leaf} ² mg _{leaf} ⁻¹	Leaf area per dry leaf mass ^a
Leaf dry matter content	LDMC	mg _{leaf} g _{leaf} ⁻¹	Dry leaf mass per water-saturated fresh leaf mass ^a
Stomatal conductance Belowground traits	g₅	mmol m ⁻² s ⁻¹	Stomatal conductance per leaf area ^a
Root diameter	Dia	mm	Average root diameter of the root subsample
Specific root length	SRL	$m_{root} g_{root}^{-1}$	Root length per dry root biomass (subsample)
Root length density	RLD	cm _{root} cm _{soil} ⁻³	Root length (extrapolated) per soil volume (pot)
Pathogen infestation	-	%	Percentage of infested leaf area (estimated) ^a
Experimental factors			
Species identity	Species ID	-	Study species
Legacy treatments			
Plant history	PH	-	Species richness of the plant community, where the
· · · · · · · · · · · · · · · · · · ·			seeds were collected – two or six plant species
Soil history	SH	-	Species richness of plant community, where the soil for
Contributory	011		inoculation was taken – two or six plant species
			Origin of seed and soil in one pot:
			 same plot origin = home soil treatment
Coil trootmont	ст.		 different plot origin, but same species richness =
Soli treatment	31	-	away-same soil treatment
			 different plot origin, different species richness =
			away-different soil treatment
	global change		-
Global change driver treatments	driver / global		
- <u>-</u>	change / GC		
	Drought / D	_	30% instead of 60% water saturation
Drought treatment		-	30/0 instead of $00/0$ water saturation

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272

^aaveraged per pot

273**Table 2** Summary of mixed-effect model analyses testing the effects of species identity (N = 4),274legacy treatments (plant history, soil history, soil treatment), global change treatments (drought,275nitrogen input), and their interactions on plant performance (total biomass, shoot biomass, root276biomass). Shown are degrees of freedom (Df), Chi² and P-values (P). Significant effects (P <</td>2770.05) are given in bold, marginally significant effects (P < 0.1) in italics.</td>

	Total biomass				Shoot bion	nass	Root biomass		
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Species identity (ID)	3	73.25	<0.001	3	80.17	<0.001	3	121.30	<0.001
Plant history	1	3.48	0.062	1	1.36	0.244	1	3.40	0.065
Soil history	1	0.01	0.915	1	0.04	0.851	1	0.49	0.484
Soil treatment	2	2.17	0.338	2	1.20	0.548	2	3.66	0.161
Drought (D)	1	83.05	<0.001	1	110.26	<0.001	1	2.81	0.094
Nitrogen input (N)	1	257.26	<0.001	1	425.93	<0.001	1	15.89	<0.001
Species ID x Plant history	3	0.71	0.872	3	1.77	0.621	3	0.63	0.890
Species ID x Soil history	3	1.68	0.642	3	0.18	0.980	3	3.64	0.303
Species ID x Soil treatment	6	4.29	0.638	6	6.64	0.355	6	2.30	0.891
Species ID x D	3	52.00	<0.001	3	43.11	<0.001	3	98.61	<0.001
Species ID x N	3	30.46	<0.001	3	33.73	<0.001	3	18.28	<0.001
DxN	1	35.27	<0.001	1	27.47	<0.001	1	10.90	0.001
Species ID x Plant history x D	4	0.92	0.922	4	4.42	0.353	4	0.72	0.948
Species ID x Soil history x D	4	1.17	0.883	4	5.33	0.255	4	0.54	0.969
Species ID x Soil treatment x D	8	2.81	0.946	8	4.78	0.781	8	3.30	0.914
Species ID x Plant history x N	4	2.66	0.617	4	5.75	0.219	4	1.69	0.792
Species ID x Soil history x N	4	6.59	0.159	4	3.47	0.482	4	5.26	0.262
Species ID x Soil treatment x N	8	9.35	0.314	8	4.62	0.797	8	15.48	0.050
Species ID x Plant history x D x N	4	14.85	0.005	4	27.25	<0.001	4	12.61	0.013
Species ID x Soil history x D x N	4	13.14	0.011	4	14.39	0.006	4	11.81	0.019
Species ID x Soil treatment x D x N	8	6.19	0.626	8	7.91	0.442	8	4.81	0.778

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Table 3 Summary of mixed-effect model analyses testing the effects of species identity (N = 4), legacy treatments (plant history, soil history, soil treatment), and their interactions on plant performance (total biomass, shoot biomass, root biomass and root-shoot ratio), when non-treated (control) or treated with global change drivers (drought, nitrogen input, drought and nitrogen input [D x N]). Shown are degrees of freedom (Df), Chi² and P-values (P). Significant effects (P < 0.05) are given in bold, marginally significant effects (P < 0.1) in italics.

						Total bi	omass					
	Control			Drought			Nitrogen			D x N		
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Species ID	3	57.93	<0.001	3	37.43	<0.001	3	60.10	<0.001	3	27.83	<0.001
Plant history (PH)	1	0.04	0.840	1	2.08	0.149	1	4.86	0.027	1	1.17	0.280
Soil history (SH)	1	2.60	0.107	1	0.44	0.507	1	1.15	0.283	1	0.10	0.756
Soil treatment (ST)	2	0.80	0.670	2	0.46	0.795	2	3.78	0.151	2	3.28	0.194
Species ID x PH	3	1.05	0.790	3	3.44	0.328	3	1.37	0.712	3	0.04	0.998
Species ID x SH	3	3.06	0.382	3	2.48	0.478	3	1.61	0.657	3	2.48	0.479
Species ID x ST	6	3.44	0.752	6	3.55	0.737	6	4.91	0.555	6	4.04	0.672
						Shoot bi	iomass	;				
		Contro	I		Drough	t		Nitroge	n	D x N		
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Species ID	3	45.01	<0.001	3	47.33	<0.001	3	64.80	<0.001	3	43.66	<0.001
Plant history (PH)	1	0.03	0.859	1	0.45	0.502	1	2.56	0.110	1	0.77	0.381
Soil history (SH)	1	1.57	0.211	1	0.11	0.743	1	1.97	0.161	1	0.06	0.799
Soil treatment (ST)	2	0.24	0.886	2	2.51	0.286	2	4.39	0.112	2	1.91	0.385
Species ID x PH	3	0.18	0.980	3	6.79	0.079	3	4.34	0.227	3	0.60	0.900
Species ID x SH	3	7.50	0.058	3	2.08	0.556	3	0.06	0.996	3	0.67	0.881
Species ID x ST	6	6.46	0.374	6	2.67	0.849	6	7.67	0.263	6	2.27	0.893
						Root bio	omass					
Control		Drought			Nitrogen			D x N				
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Species ID	3	107.40	<0.001	3	93.04	<0.001	3	101.11	<0.001	3	81.40	<0.001
Plant history (PH)	1	<0.01	0.957	1	2.79	0.095	1	3.34	0.068	1	1.20	0.274
Soil history (SH)	1	2.79	0.095	1	1.27	0.259	1	0.05	0.828	1	0.11	0.742
Soil treatment (ST)	2	0.60	0.740	2	0.74	0.691	2	2.08	0.354	2	4.40	0.111
Species ID x PH	3	2.74	0.434	3	1.34	0.720	3	0.78	0.855	3	0.76	0.860
Species ID x SH	3	3.68	0.299	3	3.00	0.391	3	3.11	0.375	3	4.02	0.259
Species ID x ST	6	7.55	0.273	6	5.43	0.490	6	3.05	0.803	6	9.25	0.160

288 Figures

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Figure 1 Overview of experimental design. In 2016, ripe seeds of four grass species were 292 293 collected in two- and six-species plots of the Dominance Experiment (Jena Experiment), stored in a freezer and allowed to germinate in spring 2017. After germination, soil samples were 294 collected from the plots and mixed with sterilized background soil (5% + 95%), filled in pots 295 and planted with two seedlings (12 pot replicates per plot). In four pots per plot, plant and soil 296 had the same plot origin (home soil); in four pots, species richness of plant and soil origin were 297 the same, but plant species composition was different (away-same soil) and in four pots, species 298 299 richness of plant and soil origin were different (= different origin of plant and soil; awaydifferent soil; total $Nr_{pots} = 576$). Plants were exposed to global change drivers: drought, 300 301 nitrogen input, or the combination of drought and nitrogen input, or were not treated (control).



Figure 2 Shoot and root biomass production (a), and total biomass production (b) of plants grown either in soil originated from two-species or six-species communities across all four study species and separately for each species. Bars show mean values (± 1 SE); stars above bars indicate significant differences (P < 0.05), stars in brackets indicate marginally significant differences (P < 0.1).



Figure 3 caption on next page

Figure 3 Schematic overview of the results of the common garden experiment testing how plants 310 311 with a different origin (plant history) or grown in different soil (soil history, soil treatment) differ in performance and trait expression (a) and respond to global change drivers like drought (b), 312 nitrogen input (c), and the combination of both (c). Illustrated is the impact of legacy treatments 313 (= "legacy effect") and global change treatments (= "global change effect") on shoot and root 314 biomass production as well as on plant traits (growth height ("Height"), shoot nitrogen 315 concentration ("N_{shoot}"), leaf greenness ("Greenness"), leaf dry matter content ("LDMC"), specific 316 leaf area ("SLA"), stomatal conductance ("g_s"), mildew infestation ("Mildew"), root diameter 317 ("Dia"), specific root length ("SRL"), root length density ("RLD")) of the four study species. For 318 319 legacy effects, schematic illustrations of plants indicate differences in shoot and/or root biomass, when originated from two-species ("2") or six-species ("6") communities (= plant history (PH)), 320 when grown in two-species ("2") or six-species ("6") community soil (= soil history (SH)), or 321 322 when grown in away ("a") or home ("h") soil (= soil treatment; "a_s" = away-same soil). Arrows behind traits (for legacy effects) indicate, in which treatment group the value was significant higher 323 (arrow up) or lower (arrow down), e.g. "- SH: SLA ⁶ indicate that SLA in plants grown in six-324 species soil was higher than in two-species soil and "- LDMC \h" indicate that LDMC was higher 325 in plants grown in home than in away soil. For global change effects, schematic illustrations of 326 327 plants indicate whether shoot and/or root biomass of plants increased (blue arrow up) or decreased (blue arrow down) due to the impact of the respective global change driver (blue horizontal line 328 indicate no change). Arrows behind traits (for global change effects) indicate and increase (arrow 329 330 up) or decrease (arrow down) of the trait value due to the impact of the respective global change driver. 331



Figure 4 Response of plants treated with drought, nitrogen input, or a combination of both relative to non-treated plants (control) for total biomass, shoot biomass, root biomass, and root-shoot ratio across four study species (a-c) and separately for each species (d-f). Points are means and error bars are standard deviation. No symbol indicates significant differences between plants treated with global change driver and control plants, "n.s." indicate no significant difference.



Figure 5 caption on next page

- **Figure 5** Total biomass (a-c), shoot biomass (d-f), and root biomass (g-i) of plants (across all
- four study species) originated from two- or six-species communities (plant history; a, d, g);
- grown in soil originated from two-species or six-species communities (soil history; b, e, h); or
- 342 grown in home, away-same or away-different soil (soil treatment; c, f, i) and were either non-
- treated (control) or treated with drought, nitrogen input (N input) or a combination of both (D
- + N). Bars show mean values (\pm 1 SE); stars above bars indicate significant differences (P <
- 0.05), stars in brackets indicate marginally significant differences (P < 0.1).

347 Discussion

348

349 Hypothesis 1: offspring of plants selected at different diversity and grown in different soil (high
350 vs. low diversity, home vs. away) show differences in productivity and trait expression

351 Our findings that A. elatius and D. glomerata plants in soil of high-diversity communities produce more biomass than in soil of low-diversity communities are in line with 352 several greenhouse studies showing that soil conditioned by multiple plant species has a more 353 positive impact on plant growth than soil conditioned by only one or two plant species 354 (Guerrero-Ramírez et al. 2019; Yang et al. 2015). Plants probably suffered more from 355 pathogens when grown in soil of low-diversity communities and/or benefitted more from 356 interactions with soil mutualists in soil of high-diversity communities (Eisenhauer et al. 2019; 357 Guerrero-Ramírez et al. 2019; Schnitzer et al. 2011). Interestingly, this soil legacy effect was 358 359 only found in A. elatius and D. glomerata, which were both highly-productive species in the long-term field experiment. In contrast, the low-productive species A. pratensis and P. trivialis 360 showed no significant difference when grown in differently conditioned soils. This is an 361 indication that A. elatius and D. glomerata interact and/or benefit more, and A. pratensis and 362 P. trivialis less, with soil mutualists, which are more abundant at high plant diversity, explaining 363 364 our findings and the species-specific performance in the field experiment.

In contrast to biomass production, we did not find any significant influence of soil 365 history on plant trait expression of A. elatius and D. glomerata. Nevertheless, we detected some 366 other legacy treatment effects on plant trait expression, which was also found in related studies 367 368 (van Moorsel et al. 2018a; van Moorsel et al. 2018b). The impact of soil history on root diameter of A. pratensis, and the impact of soil treatment (home/away) on the growth height of A. 369 370 pratensis, on shoot nitrogen concentrations and root traits of P. trivialis, and on mildew infestation of D. glomerata indicate that plant-soil interactions influencing growth, defense, and 371 resource use strategies of plants (Xi et al. 2021), while this impact is species-specific. Moreover, 372

D. glomerata plants had higher leaf greenness and stomatal conductance, when originated from
low-diversity than from high-diversity plant communities. This could be an adaptation to higher
light availability and lower soil moisture in low-diversity communities due to lower shading
(Bachmann et al. 2018; Fischer et al. 2019; Lorentzen et al. 2008).

377

378 Hypothesis 2: global change drivers have a strong impact on the productivity and trait
379 expression of plants.

In accordance with our second hypothesis, we found that drought reduced total biomass 380 production. This was mainly caused by a loss of shoot biomass, while drought differently 381 382 affected root biomass production of the studied grass species. Individuals of A. elatius and D. 383 glomerata increased in root biomass at the expense of shoot biomass, leading to higher rootshoot ratio under drought. This is a commonly observed strategy to avoid dehydration, which 384 385 enables plants to tap water from deeper soil layers (in the field) and at the same time minimizes the water loss caused by transpiration (Eziz et al. 2017). In contrast, the low-productive species 386 either decreased instead of increased root biomass (A. pratensis) or did not change root biomass 387 production (P. trivialis) under drought. Interestingly, the low-productive species had a three 388 times higher loss of total biomass under drought (A. pratensis: -17.1%; P. trivialis: -15.3%) 389 390 than the highly-productive species (A. elatius: -6.4%; D. glomerata: -5.7%, no significant loss 391 of total biomass in D. glomerata). Presumably, the drought resistance strategy of A. elatius and D. glomerata is more effective, which is possibly a competitive advantage under the field 392 393 conditions of the Jena Experiment, explaining the dominance of these species.

The influence of drought on the expression of plant traits was plant species-specific, except for shoot nitrogen concentrations and leaf greenness, which increased under drought in three species (except *P. trivialis*). Similar results were found in previous studies (Kocoń and Staniak 2014; Rolando et al. 2015) and indicate a general strategy against drought stress: plants decrease the cell density of shoot tissues, in line with the reduction of shoot biomass to minimize

the water loss, leading to an increase in the concentration of nitrogen compounds and 399 400 chlorophyll (strong correlation between leaf greenness and chlorophyll concentration were 401 found in Bachmann et al., 2018). At species-level, the low-productive species showed trait expression changes similar to biomass loss under drought, while the highly-productive species 402 403 D. glomerata decreased in LDMC and increased in stomatal conductance, which is contrary to recent studies showing the opposite strategy to resist drought (high LDMC, low stomatal 404 405 conductance) (Bristiel et al. 2018; Jaballah et al. 2008; Lozano et al. 2020). The results may 406 differ because *D. glomerata* in our study was infested by the mildew *Blumeria graminis*, which may have changed the leaf structure, and thus also trait expression changes under drought. 407

408 Furthermore, our second hypothesis was confirmed by showing that nitrogen input 409 increased biomass production. At species-level, shoot biomass was increased in all four species, while root biomass was enhanced only in A. elatius and A. pratensis. In D. glomerata and P. 410 411 trivialis, there was also a slight, but non-significant increase in root biomass. Both species showed a strong increase in mildew infestation when fertilized. This confirms the nitrogen-412 disease hypothesis indicating that nitrogen supply increases infection severity by altering leaf 413 properties and resources for pathogens (Dordas 2008). In D. glomerata and P. trivialis, severe 414 415 infestation by powdery mildew Blumeria graminis may have led to a decrease in rates of net 416 photosynthesis (Hibberd et al. 1996; Mandal et al. 2009), so that the reduced amount of energy 417 was mainly invested in shoot biomass, e.g. for a higher leaf turnover, and less in root biomass. We found consistent changes in plant trait expression over all four species in response 418 419 to nitrogen input: growth height (except A. elatius), shoot nitrogen concentrations, and leaf greenness increased in all four species when fertilized, confirming an earlier study by Siebenkäs 420 et al. (2015). Further nitrogen-induced changes in trait expression were likely affected by 421 mildew infestation: the highly-infested species (D. glomerata, P. trivialis) showed lower 422 LDMC and higher SLA, while LDMC and SLA of non-infested species did not change. 423 424 Probably, D. glomerata and P. trivialis plants responded to the increase in mildew infestation

with a change in the leaf architecture (Cappelli et al. 2020), which could enable plants to turn 425 426 over their leaves more quickly and thus produce constantly new and unaffected leaves. With 427 regard to root traits, the non-infested species decreased in specific root length (and A. pratensis also in root diameter), while root traits remained unchanged in the highly-infested species. The 428 429 decrease in SRL and increase in diameter (i.e. thicker and shorter roots) in combination with the increase in root biomass of the fertilized A. elatius and A. pratensis plants indicate that these 430 431 plants changed the root architecture building fewer fine roots when nutrient availability is 432 enhanced, which is in line with similar research (Siebenkäs et al. 2015).

Finally, we hypothesized that global change drivers cancel out each other's impact when 433 434 applied together. This was true for the low-productive species A. pratensis and P. trivialis, 435 which did not change in total biomass production compared to control plants as also found in other research (Carlsson et al. 2017). However, the strong decrease in root-shoot ratios indicates 436 437 that A. pratensis and P. trivialis plants changed their growth strategies. Interestingly, the highproductive species A. elatius and D. glomerata slightly increased in total biomass, which is 438 mainly explainable by the additive positive impact of drought and nitrogen input on root 439 biomass, resulting in increased root-shoot ratios. Obviously, dominant (or highly-productive) 440 species in our study benefitted more strongly from the combined application of the global 441 442 change drivers in comparison to subordinate (or low-productive) species. Assuming that dry periods are becoming more frequent (Ruosteenoja et al. 2018) and nitrogen deposition may 443 steadily rise (Reay et al. 2008), our results suggest that competitive interactions change under 444 445 the impact of multiple global change drivers, and subordinate species may become more severely threatened by extinction (Pugnaire et al. 2019). 446

Moreover, our results show that the combined effects of the two global change drivers on plant trait expression may differ from the effect of drought or nitrogen input alone, with strong negative effects for some plant species (e.g. highest mildew infestation of *P. trivialis* under combined impact of global change drivers). This suggests that plants change in

451 physiology and morphology and thus in their response to global change, when a combined 452 impact becomes more frequent, with an unknown influence on community composition and 453 ecosystem functioning in the long term. This finding underlines the need for studies 454 investigating multiple, interacting global change drivers (Rillig et al. 2019; Thakur et al. 2018). 455

456 *Hypothesis 3: offspring of plants selected at different diversity and grown in different soil (high*457 *vs. low diversity, home vs. away) respond differently to global change drivers*

The soil history effect, i.e. the beneficial effect of soil biota from high-diversity plant 458 communities on biomass production of control plants, disappeared in treatments with global 459 460 change drivers, which may be explainable by a change in soil community structure under drought (Kaisermann et al. 2017; Pugnaire et al. 2019) and/or nitrogen input (Wei et al. 2018). 461 In line with our result, similar studies have shown that drought (Fry et al. 2018; Wilschut and 462 463 van Kleunen 2021) and nitrogen input (in 't Zandt et al. 2019) can interrupt or change plantsoil interactions. As a result, the beneficial effect of soil biota from high-diversity communities 464 in A. elatius and D. glomerata could have been lost due to the reduction of soil mutualists (Yang 465 et al. 2021) and/or an increase in soil-borne pathogens caused by global change drivers 466 467 (Delgado-Baquerizo et al. 2020; Tylianakis et al. 2008; van der Putten et al. 2016).

468 Next to soil history, we also found altered plant responses to global change drivers when plants originated from low- or high-diversity communities (plant history). When treated with 469 drought, there was no significant difference, but nitrogen input had a more positive impact on 470 471 plants originated from high-diversity than from low-diversity communities. One possible explanation is that plants at high diversity were selected for greater niche complementarity 472 (Zuppinger-Dingley et al. 2014), while plants at low diversity were selected for increased 473 defense against species-specific pathogens (Eisenhauer et al. 2019), that are often accumulate 474 in low-diversity environments (Eisenhauer et al. 2012). Consequently, the offspring of 475 476 individuals originated from high-diversity communities may be more efficient in allocating

additional resources in increased growth, explaining our results. Interestingly, we did not find
any significant plant history effect in plants treated with both global change drivers, indicating
that drought had a strong impact on the growth strategy of the plants and can counteract positive
diversity effects.

481 Finally, we found that plants in home and away soil may respond differently to global change drivers; however, this was only true for the high-productive species A. elatius: plants 482 483 benefitted more from fertilization in home and away-same than in away-different soil. The home advantage supports the idea that a decrease of plant diversity can lead to changes in plant-484 soil interactions and thus to differences in eco-evolutionary feedbacks at low and high diversity 485 486 (terHorst and Zee 2016). With our data in hand, we cannot determine the exact reason why we 487 found the home advantage under fertilization but not under control conditions; however, our results show that plants may respond differently to global change drivers depending on the soil 488 489 community with which they interact.

Similar to the biomass production results, almost all differences in trait expression found 490 in control plants disappeared when treated with global change drivers. Instead, many other 491 changes in trait expression occurred depending on the type of global change driver treatment 492 493 and plant species identity. Taken together, these results indicate that mainly soil biota (soil 494 history and soil treatment) and only to a lesser extent plant history play an important role in the 495 expression of traits under the influence of global change drivers. This suggests that the soil biota composition is strongly associated with the physiology and morphology of the plants. 496 497 Therefore, shifts in soil biota composition due to plant species loss and/or global change driver impact can have strong effects on the response of plants to global change, which could further 498 499 accelerate plant community change and species loss (Pugnaire et al. 2019; Yang et al. 2021).

500

501 Conclusion

In the present study, we showed for the first time that offspring of plants selected at low 503 504 and high plant diversity differently respond to global change and that plant-soil interactions play a significant role in this process. This suggests that not only external influences (i.e. global 505 change drivers), but related changes within the community (i.e. changes in eco-evolutionary 506 feedbacks) could promote a further loss of species and thus an acceleration of global change 507 effects. To confirm this assumption, future research should test the long-term influence of 508 509 global change drivers on soil biota and plants selected at low and high diversity under more 510 realistic conditions, such as plants growing in communities under field conditions.

511

512 Materials and methods

513

514 The Jena Experiment

515 Seed and soil material for our common garden experiment was collected from a longterm biodiversity experiment, the Jena Experiment, which is located in the floodplain of the 516 Saale river near Jena (Thuringia, Germany, 50° 55'N, 11° 35'E, 130 m a.s.l.) (Roscher et al. 517 2004; Weisser et al. 2017). Before the establishment of the Jena Experiment in 2002, the site 518 519 was a highly fertilized arable field, which had been used for growing wheat and vegetables from 520 the early 1960s until 2000. Mean annual air temperature recorded from 2007 to 2016 at the 521 experimental site (Weather Station Jena-Saaleaue, Max-Planck-Institute for Biogeochemistry Jena, https://www.bglobal change-jena.mpg.de/wetter/) was 9.7°C, and mean annual 522 523 precipitation was 587 mm. The soil of the study site is a Eutric Fluvisol, whereas soil texture changes from sandy loam to silty clay with increasing distance from the river Saale. Thus, four 524 525 blocks were arranged parallel to the riverside (Roscher et al., 2004).

526 Material for our study was collected in a sub-experiment of the Jena Experiment, the so-527 called Dominance Experiment. The species pool of this experiment included nine species, 528 which often reach dominance in Central European mesophilic grasslands of the Arrhenatherion

type (Ellenberg 1988): five grasses, two legumes, and two herbs. Sown plant species richness 529 530 levels were 1, 2, 3, 4, 6, and 9 species. Each species occurred eight times in the different compositions of each species-richness level. Moreover, each possible two-species combination 531 was present with equal frequency at each species-richness level of the mixtures (i.e. 2-9 species; 532 more information about the design can be found in Roscher et al. 2004). In May 2002, seeds 533 were sown with a density of 1000 viable seeds per m^2 . Seeds from all species were purchased 534 535 from a commercial supplier (Rieger-Hoffman GmBH, Blaufelden-Raboldshause, Germany). From 2002 to 2009, plants were grown in plots of 3.5×3.5 m; from 2010 onwards, plot size 536 was reduced to 1×1 m. Plots were mown every year in June and September and mown plant 537 538 material was removed. All plots were regularly weeded and never fertilized.

539

540 Seed collection, selection of study species, and experimental plots

541 In summer 2016, we collected seed material from the nine species in all Dominance Experiment plots (as bulk sample per species and plot) and stored them in a freezer (at -20° C) 542 until further use. We chose four grass species (Alopecurus pratensis L., Arrhenatherum elatius 543 (L.) P. Beauv. ex J. Presl et C. Presl, Dactylis glomerata L., Poa trivialis L.) as study species. 544 545 Furthermore, we selected 12 plots per species (six two-species and six six-species plots, i.e. 48 546 plots in total), where sufficient seed material was available. The selected plots were evenly distributed in the four blocks of the experiment (Roscher et al. 2004). The study species differed 547 strongly in their biomass production in the Dominance Experiment plots. In the two-species 548 549 plots, all four species showed a high biomass production; however, in the six-species plots, only A. elatius and D. glomerata were highly-productive, while A. pratensis and P. trivialis showed 550 551 intermediate levels and decreased in biomass production over the years (Clark et al. 2019; Roscher et al. 2007). For simplification, from here onwards, A. elatius and D. glomerata are 552 referred to as "highly-productive" species, while A. pratensis and P. trivialis are referred to as 553 "low-productive" species. 554

555

556 Preparation of background substrate and study plants

For the pot substrate, we used a sterilized sand-soil mix (= background substrate), which 557 was then inoculated with fresh living soil (5% of the total substrate by weight) from the selected 558 plots. This inoculation method is a common procedure to investigate plant-soil interactions and 559 has the advantage that only low amounts of living soil are needed and that potential abiotic 560 561 feedbacks are eliminated (Pernilla Brinkman et al. 2010). To produce sterile background substrate, we collected 1.6 m³ soil substrate from the Jena Experiment in May 2017. This soil 562 substrate was a mix of excavated soil material from different experimental plots, which was 563 564 stored for several years at the experimental area. The soil substrate was sieved to 10 mm, homogenized, and mixed with 0.4 m³ quartz sand (WF 33, Quarzwerke GmbH, Walbeck, 565 Germany). Afterwards, the soil-sand mix was steam-sterilized twice for 150 minutes at ~80°C. 566 567 More information about the steam-sterilization method and changes of abiotic and biotic soil properties can be found in Dietrich et al. (2020). 568

For the preparation of study plants, QuickPotTM trays of 20 cm³ volume (Hermann Meyer 569 KG, Rellingen, Germany) were sterilized with a potassium hypochlorite solution (Eau de Javel: 570 2.6 g KClO to 100 ml water; 1:1) and filled with an autoclaved mixture of sand and soil from 571 572 the Jena Experiment (1:1; sterilized twice for 40 min at 121°C) in June 2017. Each species and origin (i.e. plot) was sown with two or three seeds per pot plate cell. QuickPotTM trays were 573 placed in an open greenhouse (Research Station Bad Lauchstädt, UFZ) to promote germination 574 575 by natural daily temperature fluctuations. Trays were regularly watered (with demineralized water). On 29 June 2017, A. pratensis seeds were reseeded because of low germination rate. 576 577 For the other three species, one seedling per pot plate cell was removed if more than two seeds were germinated. 578

579

580 *Common garden experiment*

In July 2017, 12 soil cores (5 cm diameter, 10 cm depth) were taken in a grid of 20 x 20 581 582 cm in each Dominance Experiment plot selected for the study and stored in a cooling chamber (4°C). Soil cores were pooled per plot and sieved through a sieve with 5 mm mesh size to 583 remove stones and coarse roots. Then, 2800 cm³ steam-sterilized background substrate was 584 thoroughly mixed with 150 cm³ fresh-sieved living soil and filled in a heat-cleaned pot (3 L, 585 diameter 14.9 cm, height 18 cm) with 12 replicates per plot. Seedlings per pot plate cell were 586 587 separated, and two seedlings per species with same plot origin were transplanted into one pot (Fig. 1). In four pots per plot, we transplanted plants, which had the same plot origin as the 588 inoculated soil (home soil treatment); in the other eight pots, plant and soil origin were different 589 590 (away soil treatment). In four of these away pots, species richness of plant and soil origin was the same, but plant species composition was different (away-same soil treatment), and in the 591 other four away pots, species richness of plant and soil origin was different (away-different soil 592 593 treatment; Fig. 1). Seedlings of D. glomerata were transplanted on 18 July 2017, followed by A. elatius (20 July 2017), P. trivialis (20 and 24 July 2017), and A. pratensis (26 – 28 July 594 2017). Seedlings were immediately watered with 200 ml demineralized water after 595 transplantation, and the initial number of shoots was counted. In total, the experiment consisted 596 597 of 576 pots, each with two plants. The pots were placed in an open greenhouse with a roof, 598 which automatically closes at rain, and ambient temperatures (Research Station Bad Lauchstädt, UFZ). Pots were distributed in six blocks placing the 12 pots filled with soil from 599 one plot in one block, i.e. in each block there were 12 pots with soil of one two-species and one 600 601 six-species plot per species. The position of the pots within the blocks was randomly chosen and changed once a month to avoid potential side effects by neighboring pots and edge effects 602 of the tables. 603

During the first week after planting, plants were watered every day with 200 ml demineralized water. From week two to four, all pots were watered every other day with 380 ml demineralized water without further treatments to allow the establishment of plants and soil

607	biota i	n the pots (380 ml were used to achieve a water saturation of the soil of 60%; calculation							
608	can be found in Appendix S1). On 23 August 2017, treatments with the global change drivers								
609	were started. For every treatment (control, drought, nitrogen input, combination of drought and								
610	nitrogen input), we used three of the 12 pots per plot (one home, one away-same, and one away-								
611	differe	nt pot, respectively; Fig. 1).							
612	(I)	For control, pots were watered as before (380 ml; every other day) and were not							
613		fertilized.							
614	(II)	Drought was simulated by reduced water saturation (= 30% water saturation = 225 ml;							
615		calculation can be found in Appendix S1). Pots were still watered every other day but							
616		with 225 ml instead of 380 ml demineralized water.							
617	(III)	Nitrogen input was applied once a week with 95 mg NH_3NO_4 (33.125 mg nitrogen)							
618		resulting in a total nitrogen amount of 265 mg after eight fertilization events, which is							
619		equivalent to a nitrogen input of 150 kg ha ⁻¹ year ⁻¹ nitrogen (medium value for managed							
620		grasslands in Germany; Häußermann et al 2019). Fertilized plants were watered as							
621		before (380 ml; every other day).							
622	(IV)	For the combination of drought and nitrogen input, pots were watered with a reduced							
623		amount (225 ml) and were fertilized once a week (in the same way as for the nitrogen							
624		input treatment alone).							

Once a month, all pots were weighted before watering. The measured weight per pot was subtracted from dry soil weight plus the assigned amount of water (380 or 225 ml). The difference revealed the amount of water which was then used to water the pot to keep the anticipated levels of water saturation for the drought and control treatment.

629

630 *Data collection*

After 11 weeks of growth with global change driver treatments, plants were harvested 631 632 block-wise (between 16 October and 8 November 2018). Before harvest, aboveground traits and leaf fungal pathogen infestation were measured (Table 1). For growth height (in cm), we 633 measured the stretched shoot length of the longest vegetative shoot per plant. Only 15% of the 634 635 plants had flowered, which was neglected due to the small case number. For leaf greenness (unitless estimate of foliar chlorophyll content), three fully expanded leaves from vegetative 636 637 shoots of each plant were measured with a SPAD 502 Plus Chlorophyll Meter (Spectrum Technologies, Inc.) and values were averaged per plant. Stomatal conductance (g_s ; mmol m⁻² s⁻ 638 ¹) was measured at one fully expanded leaf per plant (i.e. two leaves per pot) with a SC-1 Leaf 639 640 Porometer (Decagon Devices Inc.). This was done block-wise and always one day after watering, between 10 a.m. and 3 p.m. Shortly before harvest, the percentage of total leaf area, 641 which was infested by fungal pathogens was estimated for each plant. A subsample of leaves 642 643 per species was taken to identify pathogens morphologically at the species level under a light microscope. Moreover, three fully expanded leaves per individual were cut, packed in wet paper 644 towels to achieve water saturation, and stored overnight in a cooling chamber at 4°C. On the 645 next day, leaves were weighed as bulk sample per pot (i.e. six leaves) after removing water 646 647 droplets with tissue paper. Afterwards, total leaf area was measured with a leaf area meter (LI-648 3000C Area Meter equipped with LI3050C transparent conveyer belt accessory, LICOR, USA). LDMC was calculated as the ratio of dry weight to fresh weight ($mg_{leaf} g_{leaf}^{-1}$) and SLA as the 649 ratio of leaf area to dry weight $(mm_{leaf}^2 mg_{leaf}^{-1})$. 650

For biomass harvest, plants were cut at ground level, and roots were cleaned by rinsing
off all soil over a 0.5 mm sieve. The fresh root biomass was weighed and a subsample of around
1-2 g fresh weight was stored at -20°C. At a later point, roots were thawed and scanned on a
flatbed scanner at 800 dpi (Epson Expression 10000 XL scanner, Regent Instruments, Quebec,
Canada), and root diameter and root length of the subsample were measured with an image
analysis software (WinRHIZO; Regent Instruments, Quebec City, Canada). Specific root length

657 (SRL) was calculated as the ratio of root length to root dry biomass (of the subsample; m_{root} 658 g_{root}^{-1}) and root length density (RLD) as the ratio of root length to soil volume in the pot (root 659 length was extrapolated from the ratio of dry root biomass of the measured subsample to total 660 dry root mass per pot; $cm_{root} cm_{soil}^{-3}$).

All biomass and leaf samples were dried at 70°C for 48 h and then weighed. To calculate total shoot biomass per pot (each with two individuals), dry shoot biomass and dry leaf mass of the sample used for leaf area measurements were added. To calculate total root biomass, dry biomass of the scanned subsample was extrapolated from the ratio of fresh root biomass to dry root biomass per pot and added to the weighed dry root biomass per pot.

666 For chemical analysis, shoot biomass of each pot was chopped, and a subsample was 667 ground with a ball mill. Then, 10 mg milled material was used to determine shoot nitrogen concentration with near-infrared spectroscopy (MPA Multi Purpose FT-NIR Analyzer, Bruker 668 669 GmbH, Ettlingen, Germany). The calibration models used to predict shoot nitrogen concentrations were derived from laboratory data generated from previous samples of grass 670 species. The accuracy of the predictions was verified by a repeated nitrogen concentration 671 analysis of 45 randomly selected samples with an elemental analyzer (Vario EL Element 672 673 Analyzer, Elementar, Hanau, Germany). Significant positive correlation (p < 0.001, r = 0.97, N 674 = 45) between concentrations resulted from near-infrared spectroscopy and analysis with the elemental analyzer demonstrate high accuracy of our predictions. 675

676

677 Data analysis

To test whether the plants performed differently depending on legacy treatments (plant history, soil history, soil treatment [home/away]), or type of global change treatment, linear mixed-effects models were fitted for all measured response variables per pot as summarized in Table 1. Furthermore, some variables were transformed to meet the assumptions of normality and variance homogeneity: if necessary, root biomass and RLD were square root-transformed

and root-shoot ratio, SLA, stomatal conductance, SRL, and pathogen infestation were logtransformed. Furthermore, outlier values of LDMC of three *P. trivialis* pots (extremely low values), and LDMC and SLA of one *A. elatius* pot (extremely low LDMC, high SLA), were excluded from the analysis.

687 For mixed-effect model analysis, we started with a null model with the random effects only. We used seed plot identity (plot, where the seeds had been collected) and soil plot identity 688 689 (plot, from which the inoculation soil had been taken) as random effects. Then, we successively added the fixed effects with species identity first, followed by the legacy treatments: plant 690 history (species richness of the plant community, where the seeds had been collected: two or 691 692 six), soil history (species richness of the plant community, where the soil for inoculation had 693 been taken: two or six), and soil treatment (home, away-same, away-different), followed by the global change driver treatments: drought (control or drought) and nitrogen input (control or 694 695 nitrogen), and finally all interactions between species identity and the other fixed effects to check whether species differ in their responses. For analysis of stomatal conductance, we used 696 daytime and air temperature as covariates, which were entered before adding the experimental 697 factors to account for possible effects of the measurement time. 698

699 Because of multiple significant interactions between species identity and other fixed 700 effects (Table 2), we further analyzed the response variables separately per species. Therefore, 701 we used the same fixed effect structure as explained above, but without species identity and additionally with the interactions between legacy treatments and global change driver 702 703 treatments (which was not done in the first model, because otherwise, it would have become too complex). For pathogen infestation, we only analyzed data of D. glomerata and P. trivialis, 704 705 because of very low infestation rates of A. elatius and A. pratensis plants. To test whether initial size influenced the performance of the phytometers later in the experiment, we added initial 706 707 shoot number as a fixed effect before the other fixed effects in separate models for analysis of 708 shoot and root biomass production.
Because of multiple significant interactions between legacy treatments and global change driver treatments (Appendix S2: Table S1-S10), we further analyzed the data for each global change driver treatment separately. We used plant history, soil history, and soil treatment as fixed effects for species-specific analysis, and for analyses across all four species, we extended the models by fitting species identity first and all possible interactions between species identity and legacy treatments in the end.

All models were fitted with maximum likelihood (ML), and likelihood ratio tests were used to decide on the significance of the fixed effects. Tukey's HSD test was used to test differences among soil treatment groups. All calculations and statistical analyses were done in R (version 3.6.1, R Development Core Team, http://www.R-project.org) including the package *lme4* (glmer and lmer) (Bates et al. 2015) and *multcomp* (Tukey HSD) (Hothorn et al. 2016) for mixed-effects model analysis.

721

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731

732 Data availability

The data that support the findings of this study are openly available in BExIS at
https://jexis.idiv.de/ddm/data/Showdata/238

735

736 **References**

- Bachmann D., Roscher C., Buchmann N. (2018) How do leaf trait values change spatially and
 temporally with light availability in a grassland diversity experiment? Oikos 127:935-
- 739 948. doi: 10.1111/oik.04533
- Bailey J.K., Wooley S.C., Lindroth R.L., Whitham T.G. (2006) Importance of species
 interactions to community heritability: a genetic basis to trophic-level interactions.
- 742 Ecology Letters 9:78-85. doi: 10.1111/j.1461-0248.2005.00844.x
- 743 Bates D. et al. (2015) Package 'lme4'. Convergence 12:2
- Bristiel P., Gillespie L., Østrem L., Balachowski J., Violle C., Volaire F. (2018) Experimental
 evaluation of the robustness of the growth–stress tolerance trade-off within the perennial
 grass *Dactylis glomerata*. Functional Ecology 32:1944-1958. doi: 10.1111/13652435.13112
- Cappelli S.L., Pichon N.A., Kempel A., Allan E. (2020) Sick plants in grassland communities:
 a growth-defense trade-off is the main driver of fungal pathogen abundance. Ecology
 Letters 23:1349-1359. doi: 10.1111/ele.13537
- Cardinale B.J. et al. (2007) Impacts of plant diversity on biomass production increase through 751 752 time because of species complementarity. Proceedings of the National Academy of Sciences of United States of America 104:18123-18128. 753 the doi: 754 10.1073/pnas.0709069104
- Carlsson M., Merten M., Kayser M., Isselstein J., Wrage-Mönnig N. (2017) Drought stress
 resistance and resilience of permanent grasslands are shaped by functional group
 composition and N fertilization. Agriculture, Ecosystems & Environment 236:52-60.
 doi: 10.1016/j.agee.2016.11.009

- 759 Clark A.T. et al. (2019) Predicting species abundances in a grassland biodiversity experiment:
- 760 Trade-offs between model complexity and generality. Journal of Ecology 108:774-787.

761 doi: 10.1111/1365-2745.13316

- Craven D. et al. (2016) Plant diversity effects on grassland productivity are robust to both
 nutrient enrichment and drought. Philosophical Transactions of the Royal Society B:
 Biological Sciences 371:20150277. doi: 10.1098/rstb.2015.0277
- Dai A.G., Zhao T.B., Chen J. (2018) Climate change and drought: a precipitation and
 evaporation perspective. Current Climate Change Reports 4:301-312. doi:
 10.1007/s40641-018-0101-6
- Delgado-Baquerizo M. et al. (2020) The proportion of soil-borne pathogens increases with
 warming at the global scale. Nature Climate Change 10:550–554. doi: 10.1038/s41558020-0759-3
- Dietrich P., Cesarz S., Eisenhauer N., Roscher C. (2020) Effects of steam sterilization on soil
 abiotic and biotic properties. Soil Organisms 92:99–108. doi: 10.25674/so92iss2pp99

773 Dordas C. (2008) Role of nutrients in controlling plant diseases in sustainable agriculture. A

review. Agronomy for Sustainable Development 28:33-46. doi: 10.1051/agro:2007051

775 Dornelas M. et al. (2014) Assemblage time series reveal biodiversity change but not systematic

776 loss. Science 344:296-299. doi: 10.1126/science.1248484

Eisenhauer N. et al. (2019) Biotic interactions, community assembly, and eco-evolutionary
 dynamics as drivers of long-term biodiversity–ecosystem functioning relationships.

Research Ideas and Outcomes 5:e47042. doi: 10.3897/rio.5.e47042

- Eisenhauer N., Reich P.B., Scheu S. (2012) Increasing plant diversity effects on productivity
 with time due to delayed soil biota effects on plants. Basic and Applied Ecology 13:571578. doi: 10.1016/j.baae.2012.09.002
- 783 Ellenberg H. (1988) Vegetation ecology of Central Europe, 4th edn. Cambridge University
 784 Press, Cambridge

- Eziz A., Yan Z., Tian D., Han W., Tang Z., Fang J. (2017) Drought effect on plant biomass
 allocation: A meta-analysis. Ecology and evolution 7:11002-11010. doi:
 10.1002/ece3.3630
- Fischer C. et al. (2019) Plant species richness and functional groups have different effects on
 soil water content in a decade-long grassland experiment. Journal of Ecology 107:127141. doi: 10.1111/1365-2745.13046
- Fry E.L., Johnson G.N., Hall A.L., Pritchard W.J., Bullock J.M., Bardgett R.D. (2018) Drought
 neutralises plant–soil feedback of two mesic grassland forbs. Oecologia 186:1113-1125.
 doi: 10.1007/s00442-018-4082-x
- Guerrero-Ramírez N.R., Reich P.B., Wagg C., Ciobanu M., Eisenhauer N. (2019) Diversity dependent plant–soil feedbacks underlie long-term plant diversity effects on primary
 productivity. Ecosphere 10:e02704. doi: 10.1002/ecs2.2704
- Hahl T., van Moorsel S.J., Schmid M.W., Zuppinger-Dingley D., Schmid B., Wagg C. (2020)
 Plant responses to diversity-driven selection and associated rhizosphere microbial
 communities. Functional Ecology:707-722. doi: 10.1111/1365-2435.13511
- 800 Häußermann U., Bach M., Klement L., Breuer L. (2019) Stickstoff-Flächenbilanzen für
- Bo1 Deutschland mit Regionalgliederung Bundesländer und Kreise—Jahre 1995 bis 2017;
 Methodik, Ergebnisse und Minderungsmaßnahmen. Abschlussbericht TEXTE
 131:2019
- Hibberd J., Richardson P., Whitbread R., Farrar J. (1996) Effects of leaf age, basal meristem
 and infection with powdery mildew on photosynthesis in barley grown in 700 μmol
 mol⁻¹CO₂. New Phytologist 134:317-325. doi: 10.1111/j.1469-8137.1996.tb04636.x
- Holland E.A., Braswell B.H., Sulzman J., Lamarque J.-F. (2005) Nitrogen deposition onto the
 United States and Western Europe: synthesis of observations and models. Ecological
 Applications 15:38-57. doi: 10.1890/03-5162

- Hothorn T. et al. (2016) Package 'multcomp'. Simultaneous inference in general parametric
 models. Project for Statistical Computing, Vienna, Austria
- in 't Zandt D., van den Brink A., de Kroon H., Visser E.J. (2019) Plant-soil feedback is shut
- down when nutrients come to town. Plant and Soil 439:541-551. doi: 10.1007/s11104019-04050-9
- Isbell F. et al. (2015) Biodiversity increases the resistance of ecosystem productivity to climate
 extremes. Nature 526:574-577. doi: 10.1038/nature15374
- Jaballah S., Gribaa A., Volaire F., Ferchichi A. (2008) Ecophysiological responses of perennial
 grasses *Stipa lagascae* and *Dactylis glomerata* under soil water deficit. Options
 Méditerranéennes Serie A 79:303-307
- 820 Kaisermann A., de Vries F.T., Griffiths R.I., Bardgett R.D. (2017) Legacy effects of drought
- 821 on plant–soil feedbacks and plant–plant interactions. New Phytologist 215:1413-1424.
 822 doi: 10.1111/nph.14661
- Kocoń A., Staniak M. (2014) Productivity and gas exchange parameters of selected pasture
 grasses under drought stress. Journal of Food, Agriculture and Environment 12:131-135
- 825 Linhart Y.B. (1988) Intrapopulation differentiation in annual plants. III. The contrasting effects
- 826 of intra-and interspecific competition. Evolution 42:1047-1064. doi: 10.1111/j.1558827 5646.1988.tb02523.x
- Lipowsky A., Schmid B., Roscher C. (2011) Selection for monoculture and mixture genotypes
 in a biodiversity experiment. Basic and Applied Ecology 12:360-371. doi:
 10.1016/j.baae.2011.03.005
- Lorentzen S., Roscher C., Schumacher J., Schulze E.D., Schmid B. (2008) Species richness and
 identity affect the use of aboveground space in experimental grasslands. Perspectives in
 Plant Ecology, Evolution and Systematics 10:73-87. doi: 10.1016/j.ppees.2007.12.001

Lozano Y.M., Aguilar-Trigueros C.A., Flaig I.C., Rillig M.C. (2020) Root trait responses to
 drought are more heterogeneous than leaf trait responses. Functional Ecology 34:2224-

836 2235. doi: 10.1111/1365-2435.13656

- Mandal K., Saravanan R., Maiti S., Kothari I.L. (2009) Effect of downy mildew disease on
 photosynthesis and chlorophyll fluorescence in *Plantago ovata* Forsk. Journal of Plant
 Diseases and Protection 116:164-168. doi: 10.1007/Bf03356305
- Marquard E. et al. (2009) Plant species richness and functional composition drive overyielding
 in a six-year grassland experiment. Ecology 90:3290-3302. doi: 10.1890/09-0069.1
- Meyer S.T. et al. (2016) Effects of biodiversity strengthen over time as ecosystem functioning
 declines at low and increases at high biodiversity. Ecosphere 7:e01619. doi:
 10.1002/ecs2.1619
- Mommer L. et al. (2018) Lost in diversity: the interactions between soil-borne fungi,
 biodiversity and plant productivity. New Phytologist 218:542-553. doi:
 10.1111/nph.15036
- Pernilla Brinkman E., Van der Putten W.H., Bakker E.J., Verhoeven K.J. (2010) Plant–soil
 feedback: experimental approaches, statistical analyses and ecological interpretations.
 Journal of Ecology 98:1063-1073. doi: 10.1111/j.1365-2745.2010.01695.x
- Pörtner H.O. et al. (2021) IPBES-IPCC co-sponsored workshop report on biodiversity and
 climate change; IPBES and IPCC. doi: 10.5281/zenodo.4782538.
- Post D.M., Palkovacs E.P. (2009) Eco-evolutionary feedbacks in community and ecosystem
 ecology: interactions between the ecological theatre and the evolutionary play.
 Philosophical Transactions of the Royal Society B: Biological Sciences 364:1629-1640.
 doi: 10.1098/rstb.2009.0012
- Pugnaire F.I. et al. (2019) Climate change effects on plant-soil feedbacks and consequences for
 biodiversity and functioning of terrestrial ecosystems. Science advances 5:eaaz1834.
- doi: 10.1126/sciadv.aaz1834

- Reay D.S., Dentener F., Smith P., Grace J., Feely R.A. (2008) Global nitrogen deposition and
 carbon sinks. Nature Geoscience 1:430-437. doi: 10.1038/ngeo230
- Reich P.B. et al. (2012) Impacts of biodiversity loss escalate through time as redundancy fades.

863 Science 336:589-592. doi: 10.1126/science.1217909

- Rillig M.C. et al. (2019) The role of multiple global change factors in driving soil functions and
 microbial biodiversity. Science 366:886-890. doi: 10.1126/science.aay2832
- Rolando J.L., Ramirez D.A., Yactayo W., Monneveux P., Quiroz R. (2015) Leaf greenness as
 a drought tolerance related trait in potato (*Solanum tuberosum* L.). Environmental and

868 Experimental Botany 110:27-35. doi: 10.1016/j.envexpbot.2014.09.006

- 869 Roscher C. et al. (2004) The role of biodiversity for element cycling and trophic interactions:
- an experimental approach in a grassland community. Basic and Applied Ecology 5:107-

871 121. doi: 10.1078/1439-1791-00216

- Roscher C., Schumacher J., Weisser W.W., Schmid B., Schulze E.D. (2007) Detecting the role
 of individual species for overyielding in experimental grassland communities composed
 of potentially dominant species. Oecologia 154:535-549. doi: 10.1007/s00442-007-
- 875 0846-4
- Ruosteenoja K., Markkanen T., Venalainen A., Raisanen P., Peltola H. (2018) Seasonal soil
 moisture and drought occurrence in Europe in CMIP5 projections for the 21st century.
 Climate Dynamics 50:1177-1192. doi: 10.1007/s00382-017-3671-4
- 879 Sage R.F. (2020) Global change biology: A primer. Global Change Biology 26:3-30. doi:
 880 10.1111/gcb.14893
- Schnitzer S.A. et al. (2011) Soil microbes drive the classic plant diversity–productivity pattern.
 Ecology 92:296-303. doi: 10.1890/10-0773.1
- Siebenkäs A., Schumacher J., Roscher C. (2015) Phenotypic plasticity to light and nutrient
 availability alters functional trait ranking across eight perennial grassland species. AoB
- 885
 Plants 7:plv029. doi: 10.1093/aobpla/plv029

886	terHorst C.P.,	Zee P.C.	(2016)	Eco-evolu	utionary	dynamics	in plan	t–soil f	feedbacks.	Functional

Ecology 30:1062-1072. doi: 10.1111/1365-2435.12671

- Thakur M.P. et al. (2018) Reduced feeding activity of soil detritivores under warmer and drier
 conditions. Nature Climate Change 8:75-78. doi: 10.1038/s41558-017-0032-6
- 890 Thakur M.P. et al. (2021) Plant-soil feedbacks and temporal dynamics of plant diversity-
- productivity relationships. Trends in Ecology & Evolution. doi:
 10.1016/j.tree.2021.03.011
- Tilman D., Reich P.B., Knops J.M. (2006) Biodiversity and ecosystem stability in a decadelong grassland experiment. Nature 441:629-632. doi: 10.1038/nature04742
- Tylianakis J.M., Didham R.K., Bascompte J., Wardle D.A. (2008) Global change and species
 interactions in terrestrial ecosystems. Ecology Letters 11:1351-1363. doi:
 10.1111/j.1461-0248.2008.01250.x
- van der Putten W.H., Bradford M.A., Pernilla Brinkman E., van de Voorde T.F., Veen G. (2016)
- Where, when and how plant-soil feedback matters in a changing world. Functional
 Ecology 30:1109-1121. doi: 10.1111/1365-2435.12657
- 901 van Moorsel S.J. et al. (2018a) Community evolution increases plant productivity at low
 902 diversity. Ecology Letters 21:128-137. doi: 10.1111/ele.12879
- van Moorsel S.J., Schmid M.W., Hahl T., Zuppinger-Dingley D., Schmid B. (2018b) Selection
 in response to community diversity alters plant performance and functional traits.
 Perspectives in Plant Ecology, Evolution and Systematics 33:51-61. doi:
 10.1016/j.ppees.2018.05.002
- 907 Wei W. et al. (2018) Fertilizer N application rate impacts plant-soil feedback in a sanqi
 908 production system. Science of the Total Environment 633:796-807. doi:
 909 10.1016/j.scitotenv.2018.03.219

- 910 Weisser W.W. et al. (2017) Biodiversity effects on ecosystem functioning in a 15-year
- 911 grassland experiment: Patterns, mechanisms, and open questions. Basic and Applied
- 912 Ecology 23:1-73. doi: 10.1016/j.baae.2017.06.002
- Wilschut R.A., van Kleunen M. (2021) Drought alters plant-soil feedback effects on biomass
 allocation but not on plant performance. Plant and Soil 462:285-296. doi:
 10.1007/s11104-021-04861-9
- Xi N. et al. (2021) Relationships between plant-soil feedbacks and functional traits. Journal of
 Ecology. doi: 10.1111/1365-2745.13731
- 918 Yang G., Roy J., Veresoglou S.D., Rillig M.C. (2021) Soil biodiversity enhances the persistence
- 919 of legumes under climate change. New Phytologist 229:2945-2956. doi:
 920 10.1111/nph.17065
- Yang L., Maron J.L., Callaway R.M. (2015) Inhibitory effects of soil biota are ameliorated by
 high plant diversity. Oecologia 179:519-525. doi: 10.1007/s00442-015-3351-1
- 223 Zuppinger-Dingley D., Schmid B., Petermann J.S., Yadav V., De Deyn G.B., Flynn D.F. (2014)
- 924 Selection for niche differentiation in plant communities increases biodiversity effects.
- 925 Nature 515:108-111. doi: 10.1038/nature13869

926

928 Appendix S1

929 Journal: eLife

930 Article: Diversity-induced plant history and soil history effects modulate plant responses to

- 931 global change
- 932 Authors: Peter Dietrich, Jens Schumacher, Nico Eisenhauer, Christiane Roscher
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- 934

935 Calculation of irrigation water quantity per pot

1) After one week of growing, pots were watered until 100% saturation and then weighted (=

937 Weight wet soil).

2) To determine the amount of water, which is needed to get 60% water saturation (control

939 value), we used the following equations:

940 (I)
$$\frac{22\% \text{ (water holding capacity)} \times 60\% \text{ saturation}}{100\% \text{ saturation}} = 13.2\%$$

941 (II) Weight_{wet soil} - $\frac{\text{Weight}_{wet soil} \times 100\% \text{ saturation}}{13.2\% + 100} = \text{Weight}_{water control}$

First, we multiplied the water holding capacity of the Jena Experiment soil-sand mix (22%)
times 60% saturation and then divided the result by 100% saturation. Second, Weight wet soil was
multiplied with 100 and then divided by 113.2. Third, the calculated weight for a 60% saturation
was subtracted from Weight wet soil per pot and averaged over all pots, which resulted in 380 ml
water.

3) Drought was simulated by 50% lower water saturation (30% saturation), while the amountof water was calculated as followed:

949 (I)
$$\frac{22\% \text{ (water holding capacity)} \times 30\% \text{ saturation}}{100\% \text{ saturation}} = 6.6\%$$

950 (II) Weight_{wet soil} - $\frac{\text{Weight}_{\text{wet soil}} \times 100\% \text{ saturation}}{6.6\% + 100} = \text{Weight}_{\text{water drought}}$

Appendix S2 951

Journal: eLife 952

Article: Diversity-induced plant history and soil history effects modulate plant responses to 953

- 954 global change
- Authors: Peter Dietrich, Jens Schumacher, Nico Eisenhauer, Christiane Roscher 955
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- 957

958 Hypothesis 2: global change drivers have a strong impact on biomass production and trait expression.

- 959
- Plant traits (separately for each species) 960

The grass P. trivialis was the only species which growth height decreased with drought, all 961 962 other species showed no significant change under drought. Under nitrogen input, the species P. 963 trivialis, A. pratensis, and D. glomerata (marginally significant) increased in growth height, 964 while under the combined impact of both global change drivers, no species significantly 965 changed in growth height (D. glomerata marginally significantly increased in height; Fig. S1). The species A.elatius, D. glomerata, and A. pratensis increased in shoot nitrogen concentration 966 and leaf greenness under the impact of drought and/or nitrogen input (similar to analysis across 967 all species; Fig. S1). In P. trivialis, drought did not affect shoot nitrogen concentration or leaf 968 greenness, and there was no additive impact of both global change drivers on leaf greenness 969 970 (leaf greenness was as high as in fertilized plants; Fig. S1).

Global change drivers had no significant influence on LDMC or SLA of A. elatius and A. 971 pratensis except for LDMC decrease and SLA increase of A. elatius plants when treated with 972 973 both global change drivers (Fig. S2). Plants of D. glomerata decreased in LDMC and increased in SLA when treated with single global change drivers, while nitrogen input had a stronger 974 impact than drought (Fig. S2). When treated with both global change drivers, D. glomerata 975 976 plants had still a significantly lower LDMC and higher SLA compared to control plants. In P. trivialis, drought had no significant influence on LDMC and SLA, while nitrogen input 977

- 978 decreased LDMC and increased SLA (Fig. S2). When treated with both global change drivers,
- 979 LDMC and SLA were as high as in fertilized plants.
- 980 In *D. glomerata*, stomatal conductance was increased, when plants were treated with drought,
- and in *A. pratensis* decreased, when treated with both global change drivers (Fig. S2). Stomatal
- 982 conductance in *A. elatius* and *P. trivialis* did not change with global change treatments (Fig.
- 983 S2).
- 984 In A. elatius, SRL decreased when fertilized, irrespective of drought, while other root traits did
- 985 not change significantly (Fig. S3). In *A. pratensis*, drought, nitrogen input, and both global
- 986 change drivers together had similar negative impacts on SRL and RLD (except for RLD under
- 987 nitrogen input, which did not change; Fig. S3). Root diameter of *A. pratensis* plants increased
- 988 under single global change drivers with additive effects under the combined application (Fig.
- 989 S3). In D. glomerata, RLD increased and in P. trivialis RLD decreased and root diameter
- 990 increased, when treated with both global change drivers (Fig. S3).

992



Figure S1 Response of plants treated with drought, nitrogen input or a combination of both (D+N) relative to non-treated plants (control) for growth height, shoot nitrogen concentrations and leaf greenness across all four study species and separately for each species. Points are means and error bars are standard deviation. Stars indicate significant differences (P < 0.05) between plants treated with GC driver and control plants.



Figure S2 Response of plants treated with drought, nitrogen input or a combination of both (D+N) relative to non-treated plants (control) for LDMC, SLA and stomatal conductance across all four study species and separately for each species. Points are means and error bars are standard deviation. Stars indicate significant differences (P < 0.05) between plants treated with GC driver and control plants.



Figure S3 Response of plants treated with drought, nitrogen input or a combination of both (D+N) relative to non-treated plants (control) for root diameter, SRL and RLD across all four study species and separately for each species. Points are means and error bars are standard deviation. Stars indicate significant differences (P < 0.05) between plants treated with GC driver and control plants.



Figure S4 Response of plants treated with drought, nitrogen input or a combination of both (D+N) relative to non-treated plants (control) for mildew infestation for *D. glomerata* and *P. trivialis*. Stars indicate significant differences (P < 0.05) between plants treated with GC driver and control plants.

1010 Tables

1011

1012**Table S1** Summary of mixed-effect model analyses testing the effects of species identity (N = 4),1013legacy treatments (plant history, soil history, soil treatment), global change treatments (drought,1014nitrogen input) and their interactions on root-shoot ratio. Shown are degrees of freedom (Df), Chi²1015and P-values (P). Significant effects (P < 0.05) are given in bold, marginally significant effects (P <</td>10160.1) in italics.

1017

		Root-shoot	ratio
	Df	Chi ²	Р
Species identity (ID)	3	133.41	<0.001
Plant history	1	1.11	0.292
Soil history	1	1.08	0.300
Soil treatment	2	1.81	0.404
Drought (D)	1	60.01	<0.001
Nitrogen input (N)	1	89.83	<0.001
Species ID x Plant history	3	0.87	0.832
Species ID x Soil history	3	4.07	0.254
Species ID x Soil treatment	6	2.79	0.835
Species ID x D	3	95.53	<0.001
Species ID x N	3	9.31	0.025
D x N	1	2.19	0.139
Species ID x Plant history x D	4	2.02	0.733
Species ID x Soil history x D	4	1.58	0.812
Species ID x Soil treatment x D	8	4.97	0.760
Species ID x Plant history x N	4	2.91	0.573
Species ID x Soil history x N	4	3.18	0.528
Species ID x Soil treatment x N	8	18.18	0.020
Species ID x Plant history x D x N	4	10.42	0.034
Species ID x Soil history x D x N	4	11.14	0.025
Species ID x Soil treatment x D x N	8	5.20	0.736

Table S2 Summary of mixed-effect model analyses testing the effects of species identity (N = 4),1020legacy treatments (plant history, soil history, soil treatment), global change treatments (drought,1021nitrogen input) and their interactions on plant performance (total biomass, shoot biomass, root1022biomass and root-shoot ratio). Shown are degrees of freedom (Df), Chi² and P-values (P). Significant1023effects (P < 0.05) are given in bold, marginally significant effects (P < 0.1) in italics.</td>

	Total biomass				Shoot bio	mass		Root bior	mass	Root-shoot ratio			
	Df	Chi	Р	Df	Chi	Р	Df	Chi	Р	Df	Chi	Р	
Species ID	3	73.25	<0.001	3	80.17	<0.001	3	121.30	<0.001	3	133.41	<0.001	
Plant history	1	3.48	0.062	1	1.36	0.244	1	3.40	0.065	1	1.11	0.292	
Soil history	1	0.01	0.915	1	0.04	0.851	1	0.49	0.484	1	1.08	0.300	
Soil treatment	2	2.17	0.338	2	1.20	0.548	2	3.66	0.161	2	1.81	0.404	
Drought (D)	1	83.05	<0.001	1	110.26	<0.001	1	2.81	0.094	1	60.01	<0.001	
Nitrogen input (N)	1 257.26 <0.001			1	425.93	<0.001	1	15.89	<0.001	1	89.83	<0.001	
D x N	1	29.23	<0.001	1	23.02	<0.001	1	8.50	0.004	1	1.75	0.185	
Plant history x D	1	0.22	0.639	1	0.21	0.643	1	0.01	0.916	1	<0.01	0.977	
Soil history x D	1	<0.01	0.944	1	0.07	0.786	1	0.10	0.746	1	0.23	0.635	
Soil treatment x D	2	1.79	0.409	2	0.77	0.681	2	1.37	0.503	2	1.29	0.526	
Plant history x N	1	1.48	0.224	1	1.59	0.207	1	0.60	0.437	1	0.35	0.553	
Soil history x N	1	3.44	0.064	1	1.33	0.249	1	2.46	0.116	1	0.83	0.363	
Soil treatment x N	2	1.43	0.489	2	1.40	0.496	2	0.43	0.806	2	0.49	0.782	
Plant history x D x N	1	2.12	0.146	1	0.84	0.358	1	1.78	0.183	1	1.27	0.260	
Soil history x D x N	1	0.95	0.330	1	2.78	0.095	1	0.08	0.780	1	0.03	0.864	
Soil treatment x D x N	0 x N 2 1.37 0.504			2	1.93	0.381	2	0.91	0.635	2	0.73	0.693	

1027**Table S3** Summary of mixed-effect model analyses testing the effects of species identity (N = 4),1028legacy treatments (plant history, soil history, soil treatment), global change treatments (drought,1029nitrogen input) and their interactions on plant trait expression. Shown are degrees of freedom (Df),1030Chi² and P-values (P). Significant effects (P < 0.05) are given in bold, marginally significant effects</td>1031(P < 0.1) in italics.</td>

	Growth height			Sho	oot nitroaei	1 conc	Leaf greenness		
	Df	Chi ²	P	Df	Chi ²	P	Df	Chi ²	P
Species ID	3	71 45	<0.001	3	57.20	<0.001	3	79.55	<0.001
Plant history	1	0.15	0.694	1	~0.01	0.960	1	0.05	0.830
Soil history	1	1 60	0.004	1	0.64	0.000	1	0.00	0.683
Soil treatment	2	3 08	0.207	2	2 27	0.420	2	0.17	0.000
Drought (D)	1	19 71	<0.157	4	65.46	<0.021	1	66 15	<0.742
Nitrogon input (N)	1	22.02	<0.001	1	772.20	<0.001	1	500.15	<0.001
	1	32.93	<0.001	1	112.20	<0.001	1	0.01	<0.001
D X N Diant history y D	1	2.00	0.294	1	40.00		1	<0.01	0.997
Plant history x D	1	2.99	0.064	1	0.06	0.606	1	< 0.01	0.950
Soli history x D	1	0.51	0.477	1	0.11	0.735	1	1.57	0.210
Soil treatment x D	2	3.54	0.171	2	0.02	0.990	2	0.69	0.707
Plant history x N	1	0.50	0.478	1	1.34	0.246	1	0.91	0.341
Soil history x N	1	1.41	0.235	1	0.19	0.666	1	1.54	0.215
Soil treatment x N	2	1.87	0.392	2	3.30	0.192	2	2.42	0.299
Plant history x D x N	1	0.83	0.364	1	0.21	0.645	1	0.79	0.373
Soil history x D x N	1	0.69	0.407	1	3.06	0.080	1	<0.01	0.977
Soil treatment x D x N	2	4.94	0.085	2	1.56	0.458	2	0.04	0.983
		LDMC			SLA		Stor	natal cond	uctance
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Air temperature	-	-	-	-	-	-	1	5.34	0.021
Daytime	-	-	-	-	-	-	1	38.25	<0.001
Species ID	3	80.52	<0.001	3	124.00	<0.001	3	47.15	<0.001
Plant history	1	0.80	0.373	1	0.06	0.805	1	1.25	0.264
Soil history	1	0.10	0.750	1	1.22	0.270	1	0.37	0.543
Soil treatment	2	1.13	0.570	2	1.64	0.441	2	3.38	0.185
Drought (D)	1	0.94	0.333	1	0.11	0.743	1	0.90	0.343
Nitrogen input (N)	1	62.84	<0.001	1	61.63	<0.001	1	8.16	0.004
D x N	1	6.69	0.010	1	0.01	0.904	1	9.33	0.002
Plant history x D	1	0.04	0.841	1	0.34	0.559	1	0.06	0.806
Soil history x D	1	0.49	0.484	1	0.02	0.883	1	0.65	0.420
Soil treatment x D	2	0.24	0.887	2	0.23	0.889	2	0.18	0.914
Plant history x N	1	0.65	0.421	1	0.16	0.688	1	0.69	0.406
Soil history x N	1	0.12	0.734	1	1.07	0.300	1	0.63	0.428
Soil treatment x N	2	0.66	0.719	2	2.92	0.232	2	0.08	0.960
Plant history x D x N	1	0.16	0.687	1	1.77	0.183	1	0.87	0.351
Soil history x D x N	1	<0.01	0.962	1	0.95	0.331	1	0.02	0.887
Soil treatment x D x N	2	2.27	0.322	2	1.33	0.514	2	3.73	0.155
		Root diam	eter		SRL			RLD	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Species ID	3	165.58	<0.001	3	174.84	<0.001	3	125.84	<0.001
Plant history	1	0.03	0.872	1	0.32	0.569	1	1.14	0.286
Soil history	1	0.37	0.544	1	0.36	0.546	1	0.25	0.617
Soil treatment	2	1.50	0.473	2	2.80	0.246	2	4.97	0.083
Drought (D)	1	11.19	0.001	1	7.67	0.006	1	16.09	<0.001
Nitrogen input (N)	1	19.83	< 0.001	1	6.68	0.010	1	1.29	0.257
D x N	1	0.25	0.619	1	1 27	0.261	1	2 14	0 144
Plant history x D	1	0.37	0 544	1	0.34	0.559	1	0.67	0 4 1 4
Soil history x D	1	0.12	0.725	1	0.48	0.491	1	0.07	0.798
Soil treatment x D	2	1 67	0.434	2	0.40	0.723	2	0.44	0.802
Plant history x N	1	0.40	0.528	1	1 91	0 167	1	<0.11	0.944
Soil history x N	1	0.40	0.520	1	0.15	0.702	1	10.07 A R N	0.252
Soil treatment v N	י 2	0.42	0.010	י כ	1 60	0.100	י 2	0.00	0.000
Diant history v D v N	∠ 1	0.21	0.012	<u>۲</u>	1.09	0.430	∠ 1	0.00	0.909
Soil history V D V N	1	1 / 2	0.002	1	1.22 2.17	0.210	1	2 0/	0.734
Soil treatment v D v N	2	0.75	0.224	2	0.8/	0.003	2	1 02	0.600

Table S4 Summary of mixed-effect model analyses testing the effects legacy treatments (plant history, soil history, soil treatment), global change treatments (drought, nitrogen input) and their interactions on plant performance (total biomass, shoot biomass, root biomass and root-shoot ratio) of *A. elatius* and *A. pratensis*. Shown are degrees of freedom (Df), Chi² and P-values (P). Significant effects (P < 0.05) are given in bold, marginally significant effects (P < 0.1) in italics.

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		Total biom	ass		Shoot bion	nass		Root biom	ass	R	oot-shoot	ratio
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.26	0.609	1	0.05	0.827	1	0.10	0.747	1	0.11	0.738
Soil history	1	0.39	0.533	1	0.03	0.865	1	1.02	0.312	1	1.28	0.258
Soil treatment	2	2.06	0.357	2	1.59	0.452	2	1.31	0.520	2	0.30	0.861
Drought (D)	1	21.54	<0.001	1	50.79	<0.001	1	6.13	0.013	1	67.84	<0.001
Nitrogen input (N)	1	125.48	<0.001	1	128.72	<0.001	1	31.68	<0.001	1	13.70	<0.001
DxN	1	36.23	<0.001	1	45.06	<0.001	1	1.86	0.173	1	0.13	0.715
Plant history x D	1	1.01	0.315	1	2.37	0.123	1	0.05	0.823	1	1.28	0.258
Soil history x D	1	0.27	0.606	1	2.01	0.156	1	0.71	0.399	1	2.11	0.146
Soil treatment x D	2	1.21	0.545	2	3.22	0.200	2	0.13	0.939	2	1.21	0.545
Plant history x N	1	0.92	0.337	1	2.00	0.157	1	0.02	0.879	1	0.46	0.497
Soil history x N	1	0.87	0.352	1	0.05	0.832	1	1.37	0.242	1	2.29	0.130
Soil treatment x N	2	3.07	0.215	2	0.80	0.669	2	6.25	0.044	2	5.64	0.060
Plant history x D x N	1	0.07	0.792	1	<0.01	0.980	1	0.15	0.696	1	0.02	0.884
Soil history x D x N	1	0.61	0.434	1	0.05	0.822	1	0.89	0.344	1	1.17	0.279
Soil treatment x D x N	2	3.61	0.165	2	2.25	0.326	2	1.33	0.515	2	0.56	0.757
						∆ nrat	oncie					

						1							
		Total biom	ass		Shoot bion	nass		Root biom	ass	Root-shoot ratio			
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	
Plant history	1	0.57	0.452	1	0.42	0.518	1	0.43	0.512	1	<0.01	0.985	
Soil history	1	0.68	0.408	1	<0.01	0.945	1	1.47	0.225	1	0.80	0.371	
Soil treatment	2	0.34	0.845	2	0.29	0.865	2	0.23	0.892	2	0.07	0.967	
Drought (D)	1	71.43	<0.001	1	38.06	<0.001	1	60.92	<0.001	1	0.15	0.696	
Nitrogen input (N)	1	74.74	<0.001	1	162.92	<0.001	1	9.71	0.002	1	55.50	<0.001	
DxN	1	26.47	<0.001	1	3.98	0.046	1	24.94	<0.001	1	16.49	<0.001	
Plant history x D	1	0.08	0.772	1	0.51	0.477	1	0.48	0.488	1	1.07	0.301	
Soil history x D	1	0.43	0.512	1	0.37	0.546	1	0.20	0.653	1	0.01	0.912	
Soil treatment x D	2	1.17	0.557	2	0.19	0.911	2	2.12	0.346	2	3.60	0.165	
Plant history x N	1	0.40	0.529	1	1.26	0.261	1	0.02	0.875	1	0.14	0.709	
Soil history x N	1	5.45	0.020	1	1.19	0.275	1	4.53	0.033	1	1.24	0.265	
Soil treatment x N	2	2.78	0.249	2	2.50	0.287	2	1.21	0.547	2	0.13	0.938	
Plant history x D x N	1	0.55	0.458	1	0.02	0.881	1	0.59	0.442	1	0.08	0.771	
Soil history x D x N	1	0.28	0.595	1	0.30	0.585	1	0.78	0.376	1	1.44	0.230	
Soil treatment x D x N	2	0.91	0.634	2	0.05	0.975	2	1.45	0.485	2	2.41	0.300	

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Table S5 Summary of mixed-effect model analyses testing the effects legacy treatments (plant1042history, soil history, soil treatment), global change treatments (drought, nitrogen input) and their1043interactions on plant performance (total biomass, shoot biomass, root biomass and root-shoot ratio)1044of *D. glomerata* and *P. trivialis*. Shown are degrees of freedom (Df), Chi² and P-values (P).1045Significant effects (P < 0.05) are given in bold, marginally significant effects (P < 0.1) in italics.</td>

	D. glomerata											
		Total biom	ass		Shoot bion	nass		Root biom	ass	R	oot-shoot	ratio
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	1.51	0.219	1	1.32	0.251	1	1.12	0.289	1	0.19	0.662
Soil history	1	0.00	0.957	1	0.01	0.912	1	0.07	0.787	1	0.05	0.829
Soil treatment	2	0.79	0.673	2	0.11	0.948	2	2.65	0.266	2	2.94	0.230
Drought (D)	1	0.98	0.323	1	12.71	<0.001	1	20.48	<0.001	1	58.54	<0.001
Nitrogen input (N)	1	82.06	<0.001	1	124.42	<0.001	1	8.87	0.003	1	16.79	<0.001
DxN	1	0.07	0.790	1	0.04	0.843	1	0.61	0.434	1	0.53	0.467
Plant history x D	1	0.05	0.821	1	0.55	0.458	1	0.24	0.623	1	1.40	0.236
Soil history x D	1	0.56	0.453	1	2.20	0.138	1	0.14	0.706	1	0.27	0.601
Soil treatment x D	2	0.09	0.955	2	0.55	0.758	2	1.09	0.579	2	3.01	0.222
Plant history x N	1	1.55	0.213	1	1.85	0.174	1	0.62	0.432	1	0.29	0.592
Soil history x N	1	1.42	0.234	1	2.24	0.135	1	0.26	0.612	1	0.25	0.618
Soil treatment x N	2	0.05	0.976	2	0.72	0.699	2	1.94	0.378	2	3.83	0.147
Plant history x D x N	1	4.64	0.031	1	3.35	0.067	1	4.09	0.043	1	3.81	0.051
Soil history x D x N	1	4.21	0.040	1	3.68	0.055	1	2.87	0.090	1	1.64	0.200
Soil treatment x D x N	2	1.70	0.428	2	3.03	0.220	2	0.66	0.718	2	0.32	0.853
						P. triv	vialis					

		Total biom	nass		Shoot bion	nass		Root bioma	ass	Root-shoot ratio			
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	
Plant history	1	0.91	0.340	1	0.03	0.870	1	1.49	0.222	1	1.36	0.244	
Soil history	1	0.26	0.611	1	0.08	0.781	1	2.43	0.119	1	4.29	0.038	
Soil treatment	2	1.23	0.540	2	1.18	0.556	2	0.62	0.732	2	0.09	0.956	
Drought (D)	1	23.05	<0.001	1	22.42	<0.001	1	8.93	0.003	1	0.00	0.988	
Nitrogen input (N)	1	27.28	<0.001	1	87.31	<0.001	1	1.12	0.290	1	45.86	<0.001	
D x N	1	3.81	0.051	1	2.16	0.141	1	2.81	0.094	1	2.10	0.147	
Plant history x D	1	0.08	0.775	1	1.03	0.311	1	0.03	0.874	1	0.20	0.656	
Soil history x D	1	<0.01	0.969	1	0.21	0.649	1	0.15	0.696	1	0.21	0.646	
Soil treatment x D	2	0.80	0.670	2	0.69	0.708	2	0.38	0.828	2	1.04	0.594	
Plant history x N	1	<0.01	0.972	1	0.87	0.350	1	0.32	0.569	1	0.73	0.391	
Soil history x N	1	<0.01	0.984	1	0.01	0.936	1	0.01	0.920	1	0.03	0.857	
Soil treatment x N	2	4.20	0.123	2	1.87	0.392	2	6.33	0.042	2	7.28	0.026	
Plant history x D x N	1	0.25	0.614	1	<0.01	0.978	1	0.17	0.680	1	0.00	0.972	
Soil history x D x N	1	0.02	0.890	1	1.11	0.292	1	1.09	0.296	1	2.88	0.089	
Soil treatment x D x N	2	0.35	0.838	2	0.49	0.782	2	1.16	0.559	2	1.97	0.373	

1049**Table S6** Summary of mixed-effect model analyses testing the effects of legacy treatments (plant1050history, soil history, soil treatment), global change treatments (drought, nitrogen input) and their1051interactions on plant trait expressions of *A. elatius*. Shown are degrees of freedom (Df), Chi² and P-1052values (P). Significant effects (P < 0.05) are given in bold, marginally significant effects (P < 0.1) in</td>1053italics.

A. elatius	Growth height				ot nitroge	n conc.	Leaf greenness		
	Df	Chi ²	P	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.24	0.625	1	0.12	0.725	1	0.07	0.795
Soil history	1	0.61	0.436	1	0.36	0.547	1	0.67	0.413
Soil treatment	2	2.01	0.365	2	0.80	0.670	2	0.19	0.907
Drought (D)	1	2.11	0.146	1	36.64	<0.001	1	30.19	<0.001
Nitrogen input (N)	1	5.35	0.021	1	142.97	<0.001	1	153.54	<0.001
D x N	1	0.02	0.881	1	32.71	<0.001	1	0.27	0.604
Plant history x D	1	4.68	0.030	1	1.41	0.236	1	0.48	0.487
Soil history x D	1	0.01	0.904	1	0.26	0.612	1	0.06	0.813
Soil treatment x D	2	3.10	0.212	2	0.38	0.827	2	1.58	0.453
Plant history x N	1	1.15	0.284	1	1.08	0.300	1	3.76	0.053
Soil history x N	1	0.61	0.434	1	0.20	0.656	1	1.09	0.295
Soil treatment x N	2	3.03	0.220	2	0.27	0.874	2	2.37	0.305
Plant history x D x N	1	0.59	0.443	1	1.85	0.174	1	0.37	0.545
Soil history x D x N	1	0.93	0.334	1	0.03	0.854	1	0.06	0.813
Soil treatment x D x N	2	7.64	0.022	2	0.26	0.877	2	1.95	0.377
		LDMC			SLA		Stor	natal cond	uctance
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Air temperature	-	-	-	-	-	-	1	<0.01	0.948
Daytime	-	-	-	-	-	-	1	8.05	0.005
Plant history	1	0.46	0.500	5	1.69	0.194	1	0.49	0.486
Soil history	1	0.19	0.666	6	1.83	0.176	1	0.05	0.823
Soil treatment	2	1.37	0.504	8	1.14	0.565	2	3.38	0.184
Drought (D)	1	7.57	0.006	9	12.37	<0.001	1	4.58	0.032
Nitrogen input (N)	1	1.05	0.307	10	0.05	0.832	1	2.00	0.158
DxN	1	0.02	0.889	11	1.87	0.171	1	0.17	0.681
Plant history x D	1	1.48	0.224	12	1.94	0.164	1	1.08	0.298
Soil history x D	1	0.36	0.549	13	0.79	0.373	1	0.05	0.830
Soil treatment x D	2	< 0.01	0.998	15	1.73	0.420	2	0.73	0.693
Plant history x N	1	0.01	0.904	16	0.08	0.782	1	0.04	0.836
Soil history x N	1	0.01	0.936	17	1.69	0.193	1	0.36	0.549
Soil treatment x N	2	2.16	0.339	19	2.01	0.367	2	0.24	0.886
Plant history x D x N	1	<0.01	0.999	20	1.96	0.162	1	0.42	0.518
Soil history x D x N	1	0.10	0.752	21	0.15	0.696	1	1.48	0.224
Soil treatment x D x N	2	0.35	0.840	23	0.50	0.781	2	1.99	0.369
		Root diame	ter	- D (SRL			RLD	
D	Df	Chi ²	۲	Df	Chi ²	Ч	Df	Chi ²	<u>۲</u>
Plant history	1	0.08	0.783	1	0.31	0.576	1	0.09	0.767
Soil history	1	0.23	0.629	1	0.22	0.639	1	0.82	0.364
Soil treatment	2	2.89	0.236	2	5.30	0.071	2	3.35	0.187
Drought (D)	1	0.32	0.572	1	5.25	0.022	1	0.04	0.851
Nitrogen input (N)	1	3.46	0.063	1	13.72	<0.001	1	0.13	0.723
D X N	1	0.01	0.932	1	1.62	0.204	1	<0.01	0.989
Plant history x D	1	0.39	0.531	1	0.11	0.740	1	0.77	0.380
Soil history x D	1	0.01	0.938	1	0.95	0.329	1	0.29	0.590
Soil treatment x D	2	2.11	0.349	2	0.51	0.775	2	0.45	0.797
Plant history x N	1	0.09	0.764	1	1.41	0.235	1	1.29	0.256
Soil history x N	1	1.35	0.246	1	0.32	0.573	1	3.53	0.060
Soil treatment x N	2	0.68	0./11	2	1.06	0.590	2	1.76	0.416
Plant history x D x N	1	1.68	0.194	1	2.73	0.099	1	3.70	0.054
Soil history x D x N	1	4.45	0.035	1	0.52	0.469	1	1.46	0.227
Soli treatment x D x N	- 2	2.00	0.369	2	2.75	0.253	2	2.26	0.324

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Table S7 Summary of mixed-effect model analyses testing the effects of legacy treatments (plant history, soil history, soil treatment), global change treatments (drought, nitrogen input) and their interactions on plant trait expressions of *A. pratensis*. Shown are degrees of freedom (Df), Chi² and P-values (P). Significant effects (P < 0.05) are given in bold, marginally significant effects (P < 0.1) in italics.

A. pratensis		Growth hei	ght	Sho	oot nitrogei	n conc.	L	eaf green	ness
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	1.35	0.246	1	0.16	0.687	1	0.49	0.485
Soil history	1	0.71	0.400	1	<0.01	0.967	1	0.11	0.745
Soil treatment	2	8.50	0.014	2	1.38	0.501	2	0.20	0.903
Drought (D)	1	1.07	0.300	1	15.42	<0.001	1	16.09	<0.001
Nitrogen input (N)	1	10.63	0.001	1	246.65	<0.001	1	143.35	<0.001
DxN	1	1.40	0.236	1	17.58	<0.001	1	0.86	0.353
Plant history x D	1	0.16	0.692	1	<0.01	0.979	1	0.58	0.446
Soil history x D	1	0.31	0.577	1	0.52	0.471	1	3.04	0.081
Soil treatment x D	2	1.11	0.575	2	0.50	0.778	2	3.39	0.183
Plant history x N	1	0.28	0.597	1	0.17	0.681	1	< 0.01	0.994
Soil history x N	1	0.01	0.919	1	0.10	0.747	1	1.10	0.293
Soil treatment x N	2	2.42	0.299	2	6.58	0.037	2	0.19	0.911
Plant history x D x N	1	0.18	0.672	1	0.87	0.352	1	1.06	0.304
Soil history x D x N	1	0.10	0.072	1	0.07	0.332	1	0.03	0.304
Soil treatment x D x N	י ר	0.45	0.501	י 2	2.09	0.405	2	0.03	0.003
			0.034	2	2.00	0.555	Stor	0.52	0.004
	Df	Chi ²	D	Df	Chi ²	Р	Df		
Air tomporatura		CIII	Г	DI	CIII	F	1	0.16	F
Air temperature	-	-	-	-	-	-	1	0.10	0.000
Dayume Diant history	-	-	-	-	- 0.40	-	1	1.70	0.102
Plant history	1	2.82	0.093	1	0.19	0.000	1	0.43	0.513
Soli history	1	1.80	0.180	1	0.94	0.332	1	0.41	0.520
Soil treatment	2	3.57	0.168	2	5.69	0.058	2	3.67	0.159
Drought (D)	1	4.02	0.045	1	1.29	0.255	1	6.17	0.013
Nitrogen input (N)	1	0.75	0.388	1	2.93	0.087	1	3.64	0.056
DXN	1	0.33	0.566	1	0.41	0.524	1	3.45	0.063
Plant history x D	1	0.13	0.715	1	0.27	0.604	1	0.03	0.862
Soil history x D	1	0.16	0.685	1	<0.01	0.980	1	0.64	0.423
Soil treatment x D	2	1.40	0.497	2	1.39	0.499	2	0.01	0.993
Plant history x N	1	1.03	0.311	1	1.02	0.313	1	0.58	0.447
Soil history x N	1	<0.01	0.950	1	0.78	0.377	1	0.18	0.669
Soil treatment x N	2	0.64	0.726	2	2.56	0.278	2	0.27	0.874
Plant history x D x N	1	0.80	0.372	1	1.67	0.197	1	2.57	0.109
Soil history x D x N	1	4.17	0.041	1	1.01	0.315	1	0.23	0.634
Soil treatment x D x N	2	0.18	0.912	2	1.09	0.581	2	15.71	<0.001
		Root diame	eter		SRL			RLD	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.01	0.935	1	0.28	0.597	1	0.06	0.809
Soil history	1	0.18	0.676	1	0.01	0.934	1	0.92	0.337
Soil treatment	2	0.54	0.763	2	0.97	0.615	2	0.12	0.940
Drought (D)	1	39.31	<0.001	1	5.25	0.022	1	82.01	<0.001
Nitrogen input (N)	1	51.80	<0.001	1	5.33	0.021	1	0.34	0.560
D x N	1	0.09	0 767	1	5 57	0.018	1	4.32	0.038
Plant history x D	1	0.00	0.906	1	0.30	0.587	1	0.26	0.611
Soil history x D	1	0.09	0.000	1	0.00	0.001	1	0.02	0.877
Soil treatment v D	י כ	2 58	0.700	י 2	0.01 ∕/ 22	0.010	י 2	0.02	0.0/1
Blant history x N	- 1	2.00	0.270	- 1	4.00	0.007	4	0.11	0.340
Soil history x N	1	6 20	0.009	1	0.19 Q 1/I	0.000	1	0.17	0.002
Soil treatment v N	1	1.00	0.011	ו ס	0.14	0.004	1	0.03	0.420
Diant history v D v N	∠	1.02	0.402	<u>ک</u>	3.21 0.45	0.190	<u>ک</u>	1.24	0.539
Fiant history X D X N	1	0.54	0.401	ן ג	1.15	0.700	ן א	0.20	0.094
Soil history X D X N	1	1.82	0.178	1	1.87	0.172	1	0.27	0.605
Soli treatment X D X N	2	3.23	0.199	2	1.63	0.443	2	0.70	0.703

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1064**Table S8** Summary of mixed-effect model analyses testing the effects of legacy treatments (plant1065history, soil history, soil treatment), global change treatments (drought, nitrogen input) and their1066interactions on plant trait expressions of *D. glomerata*. Shown are degrees of freedom (Df), Chi² and1067P-values (P). Significant effects (P < 0.05) are given in bold, marginally significant effects (P < 0.1)</td>1068in italics.

D. glomerata		Growth he	ight	Sho	pot nitroger	n conc.	L	eat green	ness
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.05	0.831	1	0.58	0.444	1	0.22	0.640
Soil history	1	1.56	0.212	1	1.35	0.245	1	0.27	0.606
Soil treatment	2	5.25	0.073	2	0.75	0.687	2	0.55	0.760
Drought (D)	1	<0.01	0.976	1	19.10	<0.001	1	29.41	<0.001
Nitrogen input (N)	1	11.51	0.001	1	183.85	<0.001	1	172.91	<0.001
D x N	1	<0.01	0.949	1	3.72	0.054	1	0.08	0.781
Plant history x D	1	0.01	0.920	1	0.05	0.828	1	2 75	0.097
Soil history x D	1	0.82	0.366	1	0.08	0.774	1	0.22	0.639
Soil treatment v D	2	0.02	0.000	2	0.00	0.880	2	0.22	0.000
Plant history v N	∠ 1	0.40	0.700	<u>ح</u> 1	2 06	0.000	∠ 1	0.21	0.099
Soil bistory X N	1	0.91	0.341	1	2.90 0.22	0.000	1	1 75	0.437
Soil fisioly X IN	1	0.23	0.033	1	0.32	0.071	1	1.75	0.100
Soil treatment X IN	2	0.35	0.840	2	0.29	0.000	2	4.92	0.085
Plant history X D X N	1	0.12	0.733	1	1.62	0.204	1	0.54	0.462
Soil history x D x N	1	0.71	0.400	1	5.07	0.024	1	< 0.01	0.998
Soil treatment x D x N	2	0.06	0.969	2	2.15	0.341	2	0.33	0.846
		LDMC			SLA		Stor	natal cond	uctance
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Air temperature	-	-	-	-	-	-	1	0.39	0.531
Daytime	-	-	-	-	-	-	1	20.31	<0.001
Plant history	1	0.12	0.727	1	0.80	0.371	1	5.08	0.024
Soil history	1	0.58	0.445	1	0.32	0.573	1	<0.01	0.944
Soil treatment	2	0.58	0.749	2	1.20	0.548	2	0.54	0.765
Drought (D)	1	0.07	0.798	1	0.54	0.461	1	9.01	0.003
Nitrogen input (N)	1	55.57	<0.001	1	57.43	<0.001	1	2.72	0.099
D x Ň	1	20.69	<0.001	1	6.61	0.010	1	6.34	0.012
Plant history x D	1	0.04	0.842	1	0.46	0.498	1	0.07	0.793
Soil history x D	1	0.01	0.926	1	0.09	0.762	1	< 0.01	0.991
Soil treatment x D	2	1 43	0.490	2	0.09	0.958	2	0 19	0.907
Plant history x N	1	0 90	0.320	1	0.00	0.803	1	0.10	0.571
Soil history x N	1	2 48	0.020	1	2 10	0.000	1	0.62	0.406
Soil treatment v N	י 2	2.40 0.13	0.110	י 2	1 56	0.155	י 2	0.03	0.400
Diant history y D y M	∠ ₄	0.13	0.930	4	0.00	0.409	4	0.09	0.300
Fidfit filstory X D X N	T A	2.00	0.157	T A	0.09	0.768	T A	0.33	000.0
	1	1.30	0.254	1	4.99	0.026	1	5.98	0.014
Soil treatment x D x N	2	3.56	0.169	2	1.09	0.579	2	1.57	0.456
		Root diam	eter		SRL	_		RLD	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.60	0.438	1	0.96	0.326	1	2.61	0.107
Soil history	1	0.06	0.805	1	0.07	0.791	1	0.01	0.933
Soil treatment	2	0.07	0.967	2	0.58	0.749	2	2.44	0.296
Drought (D)	1	0.93	0.335	1	1.16	0.281	1	9.45	0.002
Nitrogen input (N)	1	1.22	0.270	1	0.37	0.545	1	7.05	0.008
DxN	1	0.80	0.370	1	1.73	0.189	1	0.08	0.773
Plant history x D	1	3.60	0.058	1	0.64	0.425	1	0.25	0.614
Soil history x D	1	1.41	0.235	1	0.62	0.430	1	0.23	0.632
Soil treatment x D	2	0.65	0.721	2	1.95	0.377	2	2.43	0.297
Plant history x N	1	0.19	0.667	1	2.05	0.152	1	0.03	0.854
Soil history x N	1	0.60	0.437	1	<0.01	0.994	1	0.00	0.646
Soil treatment v N	2	0.00	0.457	2	0.07	0.004	2	1 76	0 4 1 4
Plant history v D v N	<u>د</u> 1	1 /0	0.000	<u>د</u> 1	0.37	0.010	<u>د</u> 1	2 11	0.79
Soil bistory X D X N	1	0.49	0.222	1	0.14	0.712	1	1.07	0.070
Soil filstory X D X N	1	0.49	0.483	1	3.54	0.000	1	1.07	0.301
Soli li eatment x D x N	2	CO. I	0.438	2	1.10	0.559	2	0.20	0.907

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Table S9 Summary of mixed-effect model analyses testing the effects of legacy treatments (plant history, soil history, soil treatment), global change treatments (drought, nitrogen input) and their interactions on plant trait expressions of *P. trivialis*. Shown are degrees of freedom (Df), Chi² and Pvalues (P). Significant effects (P < 0.05) are given in bold, marginally significant effects (P < 0.1) in italics.

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P. trivialis	Growth height		Sho	oot nitrogei	n conc.	L	_eaf green	ness	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.06	0.800	1	0.00	0.997	1	0.93	0.334
Soil history	1	2.29	0.131	1	0.05	0.824	1	1.10	0.294
Soil treatment	2	1.66	0.435	2	0.51	0.776	2	1.15	0.563
Drought (D)	1	30.17	<0.001	1	5.46	0.019	1	1.42	0.233
Nitrogen input (N)	1	12.16	<0.001	1	297.03	<0.001	1	108.82	<0.001
DxN	1	1.72	0.190	1	17.06	<0.001	1	1.09	0.296
Plant history x D	1	0.22	0.637	1	0.11	0.736	1	3.08	0.079
Soil history x D	1	2.28	0.131	1	0.53	0.469	1	0.06	0.806
Soil treatment x D	2	3.11	0.211	2	1.03	0.598	2	0.18	0.916
Plant history x N	1	5.16	0.023	1	0.05	0.821	1	0.13	0.719
Soil history x N	1	3.49	0.062	1	0.04	0.842	1	0.36	0.549
Soil treatment x N	2	2.08	0.354	2	1.04	0.594	2	1.98	0.371
Plant history x D x N	1	0.92	0.336	1	0.03	0.865	1	0.11	0.738
Soil history $x D x N$	1	0.13	0.718	1	0.18	0.669	1	0.00	0.967
Soil treatment x D x N	2	2 11	0.348	2	5.57	0.062	2	1 74	0.418
			0.040	2	SI A	0.002	Stor	matal cond	uctance
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	P
Air temperature		-		-	-	-	1	38.70	<0.001
Davtime	_	_	_	_	-	_	1	18 64	<0.001
Plant history	1	<0.01	0 965	1	0.62	0 431	1	0.18	0.675
Soil history	1	0.08	0.300	1	0.02	0.485	1	0.10	0.070
Soil treatment	2	1.64	0.441	2	2 12	0.405	2	3 25	0.000
Drought (D)	1	2.85	0.001	1	2.12	0.040	1	0.20	0.107
Nitrogen input (N)	1	57 72	-0 001	1	2.75 A1 AA	~0.001	1	0.22	0.000
	1	0.30	0.534	1	0.62	0.431	1	2.87	0.000
Plant history x D	1	1.00	0.004	1	0.02	0.40	1	2.07	0.030
Soil bistony x D	1	2.26	0.230	1	0.30	0.540	1	2.00	0.091
Soil treatment x D	י ר	2.20	0.155	י ר	1 22	0.502	י ר	0.01	0.900
Bloot bistory x N	4	0.19	0.900	4	1.55	0.010	4	0.40	0.019
	1	5.57	0.000	1	0.04	0.212	1	0.33	0.004
Soil fristory x N	1	0.54	0.401	1	0.21	0.040	1	2.40	0.110
Soli treatment x N	2	1.09	0.300	2	3.10	0.213	2	1.30	0.506
Plant history X D X N	1	0.13	0.720	1	0.58	0.446	1	0.14	0.704
Soll history x D x N	1	1.15	0.283	1	1.01	0.315	1	7.44	0.006
Soli treatment x D x N	2	3.30	0.192	2	0.99	0.610	2	2.20	0.333
	- Df	Root diam	eter	D 4	SRL		D 4	RLD	
Diant history	Dr		P	Dr		P	Dr		P
Plant history	1	2.10	0.147	1	2.30	0.123	1	0.04	0.040
	1	0.08	0.781	1	0.30	0.581	1	1.31	0.253
Soll treatment	2	0.13	0.938	2	0.31	0.856	2	1.13	0.568
Drought (D)	1	14.18	<0.001	1	0.89	0.347	1	18.25	<0.001
Nitrogen input (N)	1	0.17	0.677	1	3.49	0.062	1	0.03	0.872
DXN	1	0.88	0.349	1	0.25	0.618	1	1.16	0.282
Plant history x D	1	0.40	0.525	1	0.27	0.602	1	0.16	0.692
Soil history x D	1	0.48	0.487	1	0.20	0.655	1	1.36	0.244
Soil treatment x D	2	5.85	0.054	2	0.50	0.777	2	0.43	0.808
Plant history x N	1	1.28	0.258	1	0.07	0.795	1	0.28	0.594
Soil history x N	1	1.21	0.271	1	0.36	0.549	1	0.65	0.418
Soil treatment x N	2	2.99	0.225	2	9.11	0.011	2	0.33	0.846
Plant history x D x N	1	0.33	0.566	1	0.05	0.821	1	0.02	0.878
Soil history x D x N	1	4.11	0.043	1	9.74	0.002	1	2.06	0.151
Soil treatment x D x N	2	0.52	0.772	2	1.40	0.495	2	1.43	0.488

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Table S10 Summary of mixed-effect model analyses testing the effects of legacy treatments (plant history, soil history, soil treatment), global change treatments (drought, nitrogen input) and their interactions on mildew infestation of *D. glomerata* and *P. trivialis*. Shown are degrees of freedom (Df), Chi² and P-values (P). Significant effects (P < 0.05) are given in bold, marginally significant effects (P < 0.1) in italics.

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Mildew infestation		D. glomer	ata	P. trivialis				
	Df	Chi ²	Р	Df	Chi ²	Р		
Plant history	1	0.29	0.588	1	0.01	0.939		
Soil history	1	0.24	0.622	1	4.16	0.041		
Soil treatment	2	0.22	0.896	2	3.36	0.187		
Drought (D)	1	2.44	0.119	1	10.69	0.001		
Nitrogen input (N)	1	42.75	<0.001	1	38.76	<0.001		
DxN	1	1.05	0.305	1	0.98	0.321		
Plant history x D	1	0.03	0.855	1	0.02	0.889		
Soil history x D	1	2.25	0.134	1	0.07	0.788		
Soil treatment x D	2	5.79	0.055	2	0.25	0.884		
Plant history x N	1	<0.01	0.953	1	0.25	0.614		
Soil history x N	1	0.21	0.643	1	0.50	0.477		
Soil treatment x N	2	0.32	0.854	2	1.22	0.544		
Plant history x D x N	1	3.00	0.083	1	0.09	0.770		
Soil history x D x N	1	1.69	0.193	1	0.93	0.335		
Soil treatment x D x N	2	7.15	0.028	2	0.62	0.734		

1085

1087 Appendix S3

1088 Journal: eLife

1089 Article: Diversity-induced plant history and soil history effects modulate plant responses to global1090 change

1091 Authors: Peter Dietrich, Jens Schumacher, Nico Eisenhauer, Christiane Roscher

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1094 *Hypothesis 3: offspring of plants selected at different diversity and grown in different soil (high vs.*

1095 *low diversity, home vs. away) respond differently to global change drivers*

1096

1097 Plant traits and pathogen infestation (across species and for each species)

Growth height did not differ depending on soil or plant history when plants were treated with global 1098 change drivers across all study species and for D. glomerata (Table S1, S4). Plants of A. elatius in 1099 home soil were smaller than plants in away-same soil (Table S2). Nitrogen input had no influence, 1100 1101 while plants were tallest in home soil and smallest in away-different soil when treated with both global change drivers (Table S2). Plants of A. pratensis exposed to drought were taller when grown 1102 1103 in home than in away-different soil; however, this positive home effect was also only found in control 1104 plants (marginal significant; Table S3). When fertilized, the positive home effect on growth height 1105 disappeared (Table S3). Plants of P. trivialis were taller in two- than in six-species community soil when treated with both global change drivers, but they were not different when treated separately 1106 1107 with drought or nitrogen input (Table S5).

Leaf greenness and shoot nitrogen concentrations were not influenced by legacy treatments when exposed to drought. When fertilized, plants still did not differ in leaf greenness but had higher shoot nitrogen concentrations in six-species than in two-species soil, found across all study species and for *D. glomerata* (Table S1, S4). Moreover, fertilized plants had a lower shoot nitrogen concentration in home than in away-different soil, found across all species and for *A. pratensis* (Table S1, S3). When plants were treated with both global change drivers, the nitrogen input effect on soil history was

cancelled out by drought (across all species and for *D. glomerata*), while the impact of soil treatment
did not: plants in home soil still had lower shoot nitrogen concentration than plants in away soil
(across all species and for *A. pratensis*).

Plants treated with global change drivers did not differ significantly in LDMC or SLA dependent on legacy treatments, across all study species and in *A. elatius* (Table S1, S2). Drought resulted in higher LDMC of *A. pratensis* plants grown in six-species soil, and the combined application of drought and nitrogen input resulted in lower SLA in home than in away soil (Table S3). Fertilized *D. glomerata* plants had higher SLA in six- than in two-species community soil (Table S4). Plants of *P. trivialis* treated with both global change drivers had lower LDMC in two- than in six-species community soil (Table S5).

Stomatal conductance (g_s) did not differ significantly depending on legacy treatments when plants were treated with global change drivers across all study species and for *A. elatius* and *P. trivialis* (Table S1, S2, S5). In *A. pratensis*, fertilized plants showed a lower g_s when grown in home than in away soil. This effect was cancelled out by drought, when treated with both global change drivers (Table S3). In *D. glomerata*, plants had higher g_s when originated from six-species communities and treated with both global change drivers; however, this was also found in control plants (Table S4).

1130 Across all study species, root diameter, SRL and RLD were not influenced by legacy treatments 1131 when treated with global change drivers (Table S1). In A. elatius, root traits also did not differ, when 1132 treated with single global change drivers, but under the combined influence of both global change drivers, plants grown in away-different soil showed the highest SRL, and plants in away-same soil 1133 1134 had the lowest SRL (Table S2). In A. pratensis, plants exposed to drought had higher SRL and RLD in two- than in six-species soil. When fertilized, we did not find an effect of legacy treatment, but the 1135 combination of both global change drivers led to higher SRL and lower root diameter when plants 1136 1137 were grown in away-same than in away-different or home soil (Table S3). In D. glomerata, RLD of plants exposed to drought was higher when originated from six-species than from two-species 1138 communities. This positive diversity impact disappeared when fertilized (Table S4). In P. trivialis, 1139

1140 SRL were lower in plants grown in six-species community soil, when exposed to drought. When

1141 fertilized, this difference disappeared (Table S5).

1142 Mildew infestation of *D. glomerata* plants exposed to drought was higher in home than in away soil,

1143 while this drought effect was cancelled out by nitrogen input (Table S6). Mildew infestation of *P*.

- 1144 trivialis plants was not significantly influenced by plant or soil history, neither with nor without
- 1145 global change drivers (Table S6).
- 1146
- 1147 Figures



1148

Figure S1 Shoot nitrogen concentrations (mg N g_{shoot}^{-1}) across all four species (a) grown in soil from two- or six-species communities and (b) grown in away-different (away_dif), away-same or home soil, and either non-treated ("Control"), treated with drought ("Drought"), with nitrogen ("N input") or a combination of drought and nirogen input ("D + N"). Bars show mean values (± 1 SE); stars above bars indicate significant differences (P < 0.05).

1155 Tables

1156

Table S1 Summary of mixed-effect model analyses testing the effects of species identity, legacy treatments (plant history, soil history, soil treatment) and their interactions on plant trait expressions, when non-treated (control) or treated with GC drivers (drought, nitrogen input, drought and nitrogen input (D x N)). Shown are degrees of freedom (Df), Chi² and P-values (P). Significant effects (P < 0.05) are given in bold, marginally significant effects (P < 0.1) in italics.

1162

	Growth height											
		Contro	I		Drough	t		Nitroger	า		D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Species ID	3	36.51	<0.001	3	46.47	<0.001	3	26.45	<0.001	3	53.85	<0.001
Plant history (PH)	1	1.76	0.185	1	1.08	0.299	1	0.06	0.812	1	0.75	0.387
Soil history (SH)	1	0.48	0.488	1	0.86	0.354	1	1.52	0.217	1	1.40	0.237
Soil treatment (ST)	2	3.99	0.136	2	5.49	0.064	2	2.68	0.262	2	4.37	0.113
Species ID x PH	3	4.12	0.249	3	4.53	0.210	3	2.62	0.455	3	0.17	0.982
Species ID x SH	3	3.65	0.301	3	1.16	0.762	3	1.14	0.766	3	6.66	0.084
Species ID x ST	6	8.19	0.224	6	13.52	0.035	6	6.01	0.423	6	7.18	0.305
					Sho	ot nitrogen	conce	ntration				
		Contro			Drough	t		Nitroger	۱		D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Species ID	3	49.63	<0.001	3	23.08	<0.001	3	73.52	<0.001	3	30.02	<0.001
Plant history (PH)	1	0.94	0.333	1	0.08	0.775	1	0.50	0.480	1	0.03	0.871
Soil history (SH)	1	<0.01	0.963	1	1.50	0.221	1	4.67	0.031	1	<0.01	0.953
Soil treatment (ST)	2	2.94	0.230	2	1.32	0.517	2	7.52	0.023	2	8.53	0.014
Species ID x PH	3	2.80	0.424	3	5.03	0.170	3	4.00	0.262	3	2.20	0.533
Species ID x SH	3	1.14	0.767	3	2.99	0.392	3	7.02	0.071	3	0.31	0.958
Species ID x ST	6	12.36	0.054	6	6.88	0.332	6	6.13	0.409	6	4.73	0.579
						Leaf gree	ennes	S				
		Contro			Drough	t		Nitroger	١		D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Species ID	3	45.88	<0.001	3	44.96	<0.001	3	54.85	<0.001	3	71.04	<0.001
Plant history (PH)	1	1.61	0.204	1	0.11	0.740	1	0.43	0.514	1	0.02	0.876
Soil history (SH)	1	0.18	0.675	1	1.84	0.175	1	1.04	0.308	1	0.11	0.738
Soil treatment (ST)	2	2.10	0.350	2	1.62	0.444	2	0.41	0.813	2	1.62	0.445
Species ID x PH	3	4.39	0.222	3	3.98	0.264	3	1.88	0.600	3	2.78	0.427
Species ID x SH	3	4.45	0.216	3	3.44	0.329	3	0.89	0.829	3	0.35	0.950
Species ID x ST	6	3.54	0.739	6	3.92	0.688	6	8.79	0.186	6	3.38	0.759
						LDN	1C					
		Contro			Drough	t		Nitroger	า		D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Species ID	3	32.76	<0.001	3	22.47	<0.001	3	78.30	<0.001	3	43.04	<0.001
Plant history (PH)	1	0.33	0.565	1	2.01	0.156	1	0.03	0.861	1	0.03	0.870
Soil history (SH)	1	0.02	0.887	1	0.56	0.456	1	0.06	0.808	1	0.17	0.680
Soil treatment (ST)	2	2.83	0.243	2	1.27	0.529	2	1.34	0.511	2	0.80	0.670
Species ID x PH	3	1.71	0.635	3	0.26	0.967	3	1.00	0.802	3	4.79	0.188
Species ID x SH	3	1.69	0.638	3	4.04	0.257	3	5.48	0.140	3	2.91	0.405
Species ID x ST	6	3.52	0.742	6	1.10	0.981	6	5.73	0.454	6	11.22	0.082
						SL	A					
		Contro			Drough	t		Nitroger	<u>ו</u>		D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	P	Df	Chi ²	Р
Species ID	3	86.36	<0.001	3	57.20	<0.001	3	101.71	<0.001	3	73.53	<0.001
Plant history (PH)	1	0.19	0.661	1	0.39	0.530	1	1.55	0.214	1	0.33	0.567
Soil history (SH)	1	0.64	0.425	1	0.01	0.926	1	3.35	0.067	1	0.26	0.607
Soil treatment (ST)	2	4.38	0.112	2	1.43	0.488	2	2.32	0.313	2	1.50	0.472
Species ID x PH	3	1.58	0.663	3	1.26	0.738	3	0.96	0.810	3	4.38	0.223
Species ID x SH	3	2.26	0.521	3	1.47	0.690	3	3.69	0.297	3	1.90	0.592
Species ID x ST	6	2.38	0.882	6	2.88	0.824	6	4.08	0.666	6	14.22	0.027

1163

Table S1 continued

		Stomatal conductance											
		Contro	I		Drough	t		Nitroge	า		D x N		
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	
Temperature	1	0.75	0.388	1	1.40	0.237	1	3.18	0.074	1	0.18	0.670	
Daytime	1	18.95	<0.001	1	13.20	<0.001	1	5.72	<0.001	1	16.06	<0.001	
Species ID	3	45.36	<0.001	3	24.61	<0.001	3	42.88	<0.001	3	21.71	<0.001	
Plant history (PH)	1	0.60	0.438	1	0.01	0.910	1	0.48	0.490	1	2.95	0.086	
Soil history (SH)	1	0.10	0.757	1	0.05	0.818	1	1.15	0.283	1	0.07	0.797	
Soil treatment (ST)	2	0.08	0.963	2	2.67	0.263	2	4.85	0.088	2	0.20	0.905	
Species ID x PH	3	4.59	0.204	3	3.18	0.365	3	4.89	0.180	3	4.89	0.180	
Species ID x SH	3	2.60	0.457	3	3.53	0.317	3	3.23	0.358	3	3.36	0.340	
Species ID x ST	6	8.36	0.213	6	4.47	0.614	6	3.82	0.701	6	4.76	0.575	
						Root dia	ameter						
		Contro			Drough	t		Nitroge	า		D x N		
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	
Species ID	3	97.02	<0.001	3	103.81	<0.001	3	93.37	<0.001	3	106.66	<0.001	
Plant history (PH)	1	0.87	0.352	1	0.02	0.883	1	<0.01	0.951	1	0.08	0.775	
Soil history (SH)	1	0.17	0.680	1	0.22	0.643	1	1.41	0.235	1	0.03	0.873	
Soil treatment (ST)	2	2.42	0.298	2	0.93	0.629	2	1.28	0.528	2	0.46	0.793	
Species ID x PH	3	0.79	0.852	3	0.19	0.979	3	4.53	0.291	3	3.28	0.350	
Species ID x SH	3	6.10	0.107	3	3.40	0.334	3	5.40	0.145	3	0.31	0.959	
Species ID x ST	6	9.36	0.155	6	2.06	0.914	6	1.41	0.965	6	13.49	0.036	
						SR	L						
		Contro	l		Drough	t		Nitroge	า		D x N		
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	
Species ID	3	125.58	<0.001	3	123.96	<0.001	3	117.21	<0.001	3	144.90	<0.001	
Plant history (PH)	1	0.31	0.579	1	0.04	0.833	1	2.81	0.094	1	0.05	0.830	
Soil history (SH)	1	<0.01	0.986	1	1.17	0.279	1	1.37	0.242	1	1.48	0.224	
Soil treatment (ST)	2	1.46	0.483	2	0.67	0.717	2	4.01	0.135	2	0.28	0.869	
Species ID x PH	3	5.15	0.161	3	2.11	0.550	3	2.96	0.397	3	2.31	0.510	
Species ID x SH	3	3.89	0.274	3	6.14	0.105	3	3.40	0.334	3	1.93	0.586	
Species ID x ST	6	13.23	0.040	6	2.92	0.819	6	2.90	0.821	6	14.70	0.023	
						RL	D						
		Contro	l		Drough	t		Nitroge	า		D x N		
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	
Species ID	3	99.14	<0.001	3	101.33	<0.001	3	91.27	<0.001	3	75.25	<0.001	
Plant history (PH)	1	0.00	0.956	1	3.36	0.067	1	0.11	0.742	1	0.98	0.323	
Soil history (SH)	1	2.93	0.087	1	0.14	0.710	1	0.67	0.413	1	0.55	0.460	
Soil treatment (ST)	2	2.50	0.286	2	2.56	0.279	2	0.03	0.983	2	4.98	0.083	
Species ID x PH	3	1.35	0.716	3	5.11	0.164	3	2.59	0.459	3	0.59	0.900	
Species ID x SH	3	5.42	0.144	3	2.89	0.409	3	0.45	0.929	3	0.49	0.921	
Species ID x ST	6	2.77	0.838	6	4.44	0.617	6	0.91	0.989	6	6.27	0.393	

Table S2 Summary of mixed-effect model analyses testing the effects of legacy treatments (plant history, soil history, soil treatment) on plant trait expressions of *A. elatius*, when non-treated (control) or treated with GC drivers (drought, nitrogen input, drought and nitrogen input (D x N)). Shown are degrees of freedom (Df), Chi² and P-values (P). Significant effects (P < 0.05) are given in bold, marginally significant effects (P < 0.1) in italics.

A. elatius						Growth	height					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.32	0.569	1	2.94	0.087	1	1.01	0.314	1	0.13	0.719
Soil history	1	1.50	0.221	1	0.07	0.787	1	0.14	0.706	1	0.29	0.593
Soil treatment	2	2.67	0.263	2	10.64	0.005	2	1.55	0.461	2	7.58	0.023
					Shoot	nitrogen	concer	ntration				
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.52	0.472	1	3.46	0.063	1	<0.01	0.974	1	0.06	0.802
Soil history	1	0.89	0.347	1	0.04	0.843	1	1.64	0.200	1	0.06	0.803
Soil treatment	2	1.40	0.497	2	1.54	0.462	2	1.99	0.369	2	2.07	0.354
	_					Leaf gree	enness	5				
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	1.19	0.275	1	0.60	0.438	1	1.13	0.288	1	0.22	0.636
Soil history	1	1.50	0.221	1	0.99	0.321	1	0.03	0.862	1	0.15	0.699
Soil treatment	2	5.20	0.074	2	0.44	0.801	2	3.64	0.162	2	0.84	0.656
	_					LDN	/IC					
		Control			Drought			Nitrogen			DxN	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.01	0.942	1	1.15	0.284	1	<0.01	0.987	1	1.02	0.313
Soil history	1	0.07	0.798	1	0.13	0.718	1	0.04	0.837	1	0.31	0.580
Soil treatment	2	0.03	0.985	2	0.34	0.844	2	2.00	0.369	2	2.44	0.295
						SL	A					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.44	0.507	1	0.61	0.435	1	0.48	0.488	1	1.63	0.202
Soil history	1	0.04	0.836	1	0.22	0.638	1	0.88	0.348	1	1.08	0.300
Soil treatment	2	0.59	0.744	2	0.13	0.936	2	2.74	0.254	2	3.10	0.212
					Sto	omatal co	nducta	nce				
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Temperature	1	0.05	0.827	1	0.53	0.465	1	0.91	0.340	1	0.09	0.763
Daytime	1	6.15	0.013	1	3.92	0.048	1	0.68	0.408	1	0.37	0.544
Plant history	1	0.49	0.484	1	0.05	0.824	1	1.23	0.267	1	0.18	0.670
Soil history	1	0.83	0.361	1	0.13	0.718	1	0.92	0.336	1	<0.01	0.998
Soil treatment	2	0.96	0.618	2	1.69	0.429	2	2.99	0.224	2	0.33	0.846
						Root dia	ameter					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.24	0.627	1	<0.01	0.972	1	0.46	0.497	1	0.45	0.503
Soil history	1	1.37	0.242	1	0.53	0.467	1	2.59	0.108	1	0.10	0.754
Soil treatment	2	4.85	0.089	2	0.52	0.770	2	1.00	0.605	2	3.86	0.145
						SR	L					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.80	0.371	1	0.16	0.686	1	2.32	0.128	1	0.54	0.462
Soil history	1	0.06	0.807	1	0.02	0.884	1	2.66	0.103	1	0.21	0.649
Soil treatment	2	2.94	0.230	2	1.81	0.404	2	4.63	0.099	2	9.49	0.009
						RL	D					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	1.02	0.313	1	0.03	0.859	1	2.42	0.120	1	1.44	0.230
Soil history	1	2.51	0.113	1	1.14	0.286	1	1.03	0.310	1	0.46	0.500
Soil treatment	2	4.52	0.104	2	1.24	0.539	2	0.26	0.878	2	1.40	0.497

Table S3 Summary of mixed-effect model analyses testing the effects of legacy treatments (plant history, soil history, soil treatment) on plant trait expressions of *A. pratensis*, when non-treated (control) or treated with GC drivers (drought, nitrogen input, drought and nitrogen input (D x N)). Shown are degrees of freedom (Df), Chi² and P-values (P). Significant effects (P < 0.05) are given in bold, marginally significant effects (P < 0.1) in italics.

A. pratensis						Growth	height					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	1.50	0.221	1	0.56	0.454	1	0.03	0.868	1	0.94	0.332
Soil history	1	0.44	0.508	1	0.15	0.700	1	0.03	0.874	1	0.82	0.365
Soil treatment	2	5.77	0.056	2	6.56	0.038	2	3.00	0.223	2	0.26	0.879
					Shoot	nitrogen	concer	ntration				
		Control			Drought	maogen	0011001	Nitrogen				
	Df	Chi2	D	Df	Chi2	D	Df		П	Df	Chi2	D
Disatistation			F			F						
Plant history	1	1.75	0.186	1	0.17	0.680	1	0.10	0.755	1	0.84	0.358
Soil history	1	0.37	0.544	1	0.96	0.328	1	0.01	0.939	1	0.00	0.966
Soil treatment	2	4.61	0.100	2	1.74	0.419	2	9.05	0.011	2	6.83	0.033
						Leaf gree	enness	5				
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.07	0.786	1	1.03	0.311	1	0.58	0.445	1	0.18	0.673
Soil history	1	0.03	0.869	1	1.85	0.174	1	0.90	0.343	1	0.19	0.661
Soil treatment	2	1.16	0.560	2	0.60	0.743	2	0.61	0.737	2	2.21	0.332
		-				LDN	ЛС					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.24	0.561	1	0.28	0.529	1	0.40	0.527	1	2 17	0.140
Fiant history	1	0.34	0.301	1	0.00	0.057	1	0.40	0.327	1	2.17	0.140
Soll treatment	1 2	0.11	0.730	1	3.02	0.007	1 2	2.32	0.120	1 2	0.05	0.021
Son treatment	Z	0.30	0.635	Z	1.42	0.492	2	1.10	0.555	Z	3.91	0.141
		0 / 1			D	SL	A	N 111				
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi	Р
Plant history	1	0.07	0.786	1	0.32	0.572	1	0.00	0.984	1	1.28	0.259
Soil history	1	0.20	0.654	1	2.81	0.094	1	0.23	0.632	1	0.05	0.828
Soil treatment	2	2.21	0.331	2	0.70	0.704	2	1.18	0.555	2	8.59	0.014
					Sto	omatal co	nducta	nce				
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Temperature	1	1.17	0.279	1	0.22	0.642	1	0.44	0.507	1	0.17	0.678
Daytime	1	0.77	0.379	1	0.07	0.786	1	1.13	0.289	1	8.38	0.004
Plant history	1	0.05	0.824	1	0.16	0.690	1	0.66	0.415	1	0.61	0.436
Soil history	1	1.30	0.255	1	0.14	0.706	1	0.79	0.373	1	0.53	0.466
Soil treatment	2	2.35	0.308	2	4.41	0.110	2	2.55	0.002	2	1.59	0.452
						Root dia	ameter					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	P	Df	Chi ²	P	Df	Chi ²	P	Df	Chi ²	P
Plant history	1	0.28	0.505	1	0.18	0.673	1	0.20	0.653	1	0.09	0.770
Soil history	1	5.61	0.000	1	0.10	0.073	1	1.24	0.000	1	0.00	0.770
Soll treatment	1 2	1.00	0.010	1	0.95	0.331	1 2	1.04	0.240	1 2	0.01	0.942
Soli treatment	Z	1.02	0.002	2	0.29	0.000	2	1.20	0.555	2	0.00	0.040
		O a ra fra a l			Davashi	36	L	NP1			DUN	
		Control			Drought			Nitrogen				
	Df	Chi ²	P	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi	P
Plant history	1	0.42	0.515	1	0.24	0.623	1	0.61	0.435	1	0.01	0.916
Soil history	1	0.33	0.567	1	7.10	0.008	1	0.17	0.677	1	2.73	0.098
Soil treatment	2	5.24	0.073	2	0.88	0.644	2	0.11	0.945	2	6.03	0.049
						RL	D					
		Control			Drought			Nitrogen			DxN	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.28	0.595	1	0.12	0.729	1	0.08	0.781	1	0.09	0.763
Soil history	1	0.75	0.387	1	4.79	0.029	1	0.13	0.716	1	0.03	0.861
Soil treatment	2	0.28	0.869	2	2.39	0.303	2	0.19	0.909	2	3.02	0.221

1181**Table S4** Summary of mixed-effect model analyses testing the effects of legacy treatments (plant1182history, soil history, soil treatment) on plant trait expressions of *D. glomerata*, when non-treated1183(control) or treated with GC drivers (drought, nitrogen input, drought and nitrogen input (D x N)).1184Shown are degrees of freedom (Df), Chi² and P-values (P). Significant effects (P < 0.05) are given in</td>1185bold, marginally significant effects (P < 0.1) in italics.</td>

D. glomerata	Growth height											
		Contro	I		Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.73	0.394	1	0.11	0.741	1	0.06	0.802	1	0.01	0.912
Soil history	1	0.69	0.405	1	0.91	0.340	1	1.25	0.263	1	0.18	0.675
Soil treatment	2	1.66	0.436	2	1.06	0.589	2	2.37	0.306	2	1.09	0.581
					Shoo	t nitrogen	conce	ntration				
		Contro			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	1.13	0.289	1	0.56	0.455	1	2.38	0.123	1	0.56	0.453
Soil history	1	<0.01	0.952	1	2.18	0.140	1	8.44	0.004	1	0.05	0.818
Soil treatment	2	2.72	0.257	2	2.46	0.293	2	3.07	0.215	2	0.71	0.701
						Leaf gre	enness	5				
		Contro	l		Drought			Nitrogen			DxN	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	4.93	0.026	1	0.02	0.886	1	0.17	0.680	1	0.13	0.723
Soil history	1	1.23	0.267	1	1.17	0.279	1	0.15	0.703	1	0.01	0.908
Soil treatment	2	2.33	0.313	2	3.58	0.167	2	1.16	0.560	2	0.68	0.713
		Caratas			Drevekt	LDN	/IC	Nitrease			DuN	
	Df	Contro Chi ²		Df	Chi ²	D	Df	Chi ²	D	Df		D
Plant history		0.86	0.252	1	1 19	0.279	1	0.27	0.540	1	0.64	0 422
Soil history	1	2.00	0.555	1	0.12	0.270	1	2 21	0.340	1	0.04	0.423
Soil treatment	2	2.03	0.154	2	0.12	0.727	2	1 74	0.137	2	3.05	0.394
Soli treatment	2	2.50	0.307	2	0.20	0.303 SI	Δ	1.74	0.410	2	5.05	0.210
		Contro			Drought	01	~	Nitrogen			DxN	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	1.41	0.235	1	0.01	0.904	1	1.50	0.220	1	0.14	0.706
Soil history	1	2.29	0.130	1	0.28	0.595	1	3.86	0.050	1	0.02	0.888
Soil treatment	2	2.60	0.272	2	1.88	0.392	2	0.09	0.956	2	0.89	0.641
					St	omatal co	nducta	ince				
		Contro			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Temperature	1	1.12	0.289	1	<0.01	0.951	1	0.04	0.843	1	0.08	0.782
Daytime	1	24.06	<0.001	1	12.16	<0.001	1	4.04	0.044	1	4.37	0.037
Plant history	1	3.77	0.052	1	1.05	0.304	1	1.79	0.181	1	4.89	0.027
Soil history	1	1.44	0.231	1	1.55	0.214	1	0.47	0.493	1	2.34	0.126
Soil treatment	2	0.43	0.805	2	1.62	0.445	2	0.27	0.872	2	1.04	0.595
						Root dia	ameter					
		Contro			Drought			Nitrogen			DxN	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.64	0.422	1	0.02	0.876	1	1.83	0.176	1	2.43	0.119
Soil history	1	0.33	0.567	1	2.50	0.114	1	0.34	0.559	1	0.16	0.691
Soil treatment	2	0.60	0.741	2	3.21	0.201	2	0.16	0.924	2	2.03	0.363
		Orantas			December	SR	L	NP and a second			DuN	
		Contro		Df	Drought		D1	Nitrogen		D4		
Disathister			P	Dī		P	Dī		P	Df		P
Plant history	1	2.55	0.111	1	0.54	0.462	1	0.08	0.777	1	0.36	0.548
Soll history	1	1.73	0.188	1	1.42	0.233	1	0.32	0.570	1	0.22	0.643
Soli treatment	2	2.23	0.329	2	0.24	0.888	2	2.28	0.320	Z	2.38	0.304
		Contro	1		Drought	RL	J	Nitrogen				
	Df	Chi ²	Р	Df	Chi ²	P	Df	Chi ²	P	Df	Chi ²	P
Plant history	1	0.01	0 923	1	7 58	0.006	1	0.77	0,380	1	0.03	0.862
Soil history	1	0.27	0.602	1	0.02	0.901	1	0.54	0.464	1	0.18	0.673
Soil treatment	2	0.36	0.835	2	4.51	0.105	2	0.96	0.619	2	5.25	0.073

Table S5 Summary of mixed-effect model analyses testing the effects of legacy treatments (plant history, soil history, soil treatment) on plant trait expressions of *P. trivialis*, when non-treated (control) or treated with GC drivers (drought, nitrogen input, drought and nitrogen input (D x N)). Shown are degrees of freedom (Df), Chi² and P-values (P). Significant effects (P < 0.05) are given in bold, marginally significant effects (P < 0.1) in italics.

P. trivialis						Growth	height					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	2.81	0.094	1	0.32	0.571	1	0.98	0.323	1	0.16	0.688
Soil history	1	0.62	0.429	1	0.92	0.338	1	1.12	0.289	1	5.02	0.025
Soil treatment	2	4.77	0.092	2	1.59	0.452	2	2.99	0.224	2	1.14	0.566
					Shoot	nitrogen	concer	ntration				
		Control			Drought	_		Nitrogen	_		D x N	
	Df	Chi ²	Р	Df		P	Dt	Chi ²	P	Df	Chi ²	P
Plant history	1	<0.01	0.986	1	0.01	0.934	1	0.07	0.785	1	0.01	0.915
Soil history	1	0.15	0.695	1	0.57	0.452	1	0.06	0.802	1	0.45	0.503
Soll treatment	Z	9.66	0.008	2	2.33	0.313	2	1.18	0.554	2	3.80	0.145
	. <u> </u>	Control			Drought	Lear gree	enness	Nitrogon				
	Df	Chi ²	D	Df	Chi ²	D	Df	Chi ²	D	Df		D
Plant history	1	0.14	0 708	1	2 /1	Г 0 120	1	0.04	Г 0.845	1	2 35	Г 0 126
Soil history	1	1 41	0.700	1	1 10	0.120	1	0.04	0.043	1	0.00	0.120
Soil treatment	2	0.13	0.200	2	0.37	0.200	2	5.22	0.0074	2	0.00	0.616
	2	0.10	0.000	2	0.01	LDN		0.22	0.014	2	0.07	0.010
		Control			Drought			Nitrogen			DxN	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.81	0.369	1	0.24	0.627	1	0.05	0.826	1	1.34	0.247
Soil history	1	0.08	0.776	1	0.01	0.927	1	0.47	0.492	1	4.25	0.039
Soil treatment	2	3.34	0.188	2	0.72	0.696	2	3.01	0.222	2	2.64	0.268
						SL	A					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.26	0.611	1	0.21	0.643	1	1.44	0.231	1	0.80	0.372
Soil history	1	0.41	0.522	1	0.40	0.528	1	1.47	0.226	1	0.33	0.565
Soil treatment	2	2.29	0.319	2	0.53	0.769	2	3.35	0.187	2	4.08	0.130
					Sto	omatal co	nducta	ince				
		Control			Drought			Nitrogen			D x N	
-	Df	Chi ²	P	Df	Chi ²	P	Df	Chi ²	P	Df	Chi ²	P
Temperature	1	10.96	0.001	1	8.08	0.004	1	7.25	0.007	1	4.31	0.038
Daytime	1	3.93	0.047	1	1.12	0.289	1	1.22	0.270	1	6.35	0.012
Plant history	1	<0.01	0.949	1	0.60	0.439	1	2.96	0.085	1	0.29	0.589
Soll history	1	0.68	0.410	1	0.95	0.330	1	2.72	0.099	1	0.14	0.704
Son treatment	Z	2.40	0.293	Z	0.54	0.763	Z	0.95	0.622	Z	1.49	0.474
		Control			Drought	RUULUIA	ameter	Nitrogen			DxN	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.16	0.686	1	0.07	0 794	1	0.55	0.458	1	2 91	0.088
Soil history	1	3.06	0.000	1	0.07	0.579	1	0.95	0.329	1	0.06	0.800
Soil treatment	2	7.48	0.024	2	0.28	0.870	2	0.07	0.967	2	2.00	0.369
				_		SR	L					
		Control			Drought	-		Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	2.10	0.147	1	0.94	0.332	1	1.82	0.178	1	1.04	0.308
Soil history	1	1.83	0.177	1	3.68	0.055	1	2.26	0.133	1	0.19	0.660
Soil treatment	2	5.73	0.057	2	0.56	0.755	2	1.97	0.374	2	1.83	0.401
	_					RL	D					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.23	0.632	1	0.01	0.904	1	0.01	0.920	1	0.54	0.463
Soil history	1	3.38	0.066	1	0.07	0.792	1	0.01	0.926	1	0.16	0.685
Soil treatment	2	0.63	0.731	2	0.61	0.739	2	0.16	0.924	2	3.25	0.197

Table S6 Summary of mixed-effect model analyses testing the effects of legacy treatments (plant history, soil history, soil treatment) on mildew infestation of *D. glomerata* and *P. trivialis*, when nontreated (control) or treated with GC drivers (drought, nitrogen input, drought and nitrogen input (D x N)). Shown are degrees of freedom (Df), Chi² and P-values (P). Significant effects (P < 0.05) are given in bold, marginally significant effects (P < 0.1) in italics.

1198

Mildew infestation	D. glomerata												
		Control			Drought	0		Nitrogen			D x N		
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	
Plant history	1	0.58	0.447	1	1.18	0.277	1	0.88	0.348	1	0.26	0.613	
Soil history	1	0.41	0.522	1	2.63	0.105	1	<0.01	0.946	1	0.11	0.746	
Soil treatment	2	6.01	0.049	2	7.65	0.022	2	0.93	0.628	2	0.09	0.958	
	P. trivialis												
		Control			Drought			Nitrogen	D x N				
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	
Plant history	1	<0.01	0.996	1	<0.01	0.973	1	0.03	0.860	1	0.21	0.647	
Soil history	1	1.20	0.274	1	2.66	0.103	1	1.68	0.195	1	0.05	0.817	
Soil treatment	2	3.94	0.139	2	1.78	0.412	2	0.16	0.921	2	2.10	0.350	
1200 Biomass production

1201

1202**Table S7** Summary of mixed-effect model analyses testing the effects of species identity (N = 4),1203legacy treatments (plant history, soil history, soil treatment) and their interactions on root-shoot ratio,1204when non-treated (control) or treated with global change drivers (drought, nitrogen input, drought1205and nitrogen input (D x N)). Shown are degrees of freedom (Df), Chi² and P-values (P). Significant1206effects (P < 0.05) are given in bold, marginally significant effects (P < 0.1) in italics.</td>

1207

	Root-Shoot ratio												
	Control			Drought				Nitroge	n	D x N			
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	
Species ID	3	115.37	<0.001	3	116.36	<0.001	3	101.12	<0.001	3	108.37	<0.001	
Plant history (PH)	1	0.02	0.880	1	1.48	0.225	1	1.64	0.200	1	0.46	0.496	
Soil history (SH)	1	1.81	0.178	1	1.60	0.206	1	0.24	0.622	1	<0.01	0.992	
Soil treatment (ST)	2	0.46	0.793	2	1.96	0.376	2	1.19	0.551	2	3.54	0.170	
Species ID x PH	3	3.88	0.275	3	1.47	0.690	3	0.86	0.836	3	2.77	0.428	
Species ID x SH	3	5.98	0.113	3	3.99	0.263	3	2.53	0.471	3	3.71	0.295	
Species ID x ST	6	10.54	0.104	6	6.76	0.344	6	1.85	0.933	6	14.79	0.022	

1208

1209

1210 Table So Summary of mixed-effect model analyses testing the effects of legacy treatments (p	1210	Table S8 Summary of mixed-effect model	analyses testing the effects	of legacy treatments (pla
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1211 history, soil history, soil treatment) on plant performance (total biomass, shoot biomass, root

1212 biomass and root-shoot ratio) of *A. elatius*, when non-treated (control) or treated with GC drivers

1213 (drought, nitrogen input, drought and nitrogen input (D x N)). Shown are degrees of freedom (Df),

- 1214 Chi² and P-values (P). Significant effects (P < 0.05) are given in bold, marginally significant effects
- 1215 (P < 0.1) in italics.

A. elatius						Total bi	omass					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.35	0.557	1	0.82	0.364	1	0.71	0.401	1	0.26	0.613
Soil history	1	1.08	0.298	1	0.76	0.383	1	0.06	0.811	1	0.47	0.494
Soil treatment	2	0.10	0.949	2	2.91	0.233	2	6.44	0.040	2	0.99	0.610
						Shoot bi	iomass					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.12	0.726	1	3.36	0.067	1	1.27	0.260	1	0.01	0.904
Soil history	1	0.35	0.556	1	0.24	0.621	1	0.55	0.460	1	0.63	0.428
Soil treatment	2	2.08	0.354	2	2.89	0.236	2	5.24	0.073	2	0.98	0.613
						Root bio	omass					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.03	0.860	1	0.15	0.701	1	0.01	0.916	1	0.36	0.551
Soil history	1	3.81	0.051	1	0.62	0.433	1	0.17	0.676	1	0.22	0.636
Soil treatment	2	2.05	0.359	2	2.38	0.304	2	2.25	0.325	2	1.68	0.432
						Root-sho	oot ratio)				
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.07	0.797	1	1.62	0.203	1	0.16	0.691	1	0.31	0.576
Soil history	1	4.86	0.027	1	0.24	0.626	1	0.50	0.479	1	0.07	0.787
Soil treatment	2	3.11	0.211	2	2.39	0.302	2	0.18	0.915	2	1.88	0.391

Table S9 Summary of mixed-effect model analyses testing the effects of legacy treatments (plant1219history, soil history, soil treatment) on plant performance (total biomass, shoot biomass, root biomass1220and root-shoot ratio) of *A. pratensis*, when non-treated (control) or treated with GC drivers (drought,1221nitrogen input, drought and nitrogen input (D x N)). Shown are degrees of freedom (Df), Chi² and P-1222values (P). Significant effects (P < 0.05) are given in bold, marginally significant effects (P < 0.1) in</td>1223italics.

A. pratensis						Total bio	biomass						
		Control			Drought		Nitrogen				D x N		
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	
Plant history	1	0.05	0.820	1	0.27	0.603	1	1.63	0.202	1	1.44	0.230	
Soil history	1	0.02	0.879	1	1.05	0.306	1	2.97	0.085	1	2.07	0.151	
Soil treatment	2	3.43	0.180	2	0.17	0.917	2	1.29	0.525	2	2.80	0.247	
						Shoot bi	omass						
		Control		Drought				Nitrogen	I	D x N			
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	
Plant history	1	0.11	0.741	1	0.29	0.590	1	0.65	0.421	1	2.23	0.135	
Soil history	1	0.14	0.710	1	0.33	0.564	1	0.86	0.354	1	<0.01	0.971	
Soil treatment	2	0.15	0.927	2	1.84	0.398	2	1.03	0.596	2	1.35	0.509	
						Root bio	omass						
		Control		Drought			Nitrogen				D x N		
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	
Plant history	1	0.13	0.719	1	0.23	0.629	1	0.97	0.324	1	0.47	0.495	
Soil history	1	0.15	0.703	1	1.16	0.281	1	1.83	0.176	1	3.98	0.046	
Soil treatment	2	2.78	0.250	2	1.38	0.501	2	0.47	0.789	2	3.16	0.206	
						Root-sho	ot ratio)					
		Control			Drought			Nitrogen	I		D x N		
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	
Plant history	1	0.13	0.719	1	0.01	0.920	1	0.30	0.584	1	0.90	0.342	
Soil history	1	0.20	0.654	1	0.31	0.579	1	0.42	0.517	1	4.57	0.033	
Soil treatment	2	1.33	0.514	2	4.94	0.084	2	0.04	0.982	2	0.37	0.832	

1227 Table S10 Summary of mixed-effect model analyses testing the effects of legacy treatments (plant

1228 history, soil history, soil treatment) on plant performance (total biomass, shoot biomass, root

biomass and root-shoot ratio) of *D. glomerata*, when non-treated (control) or treated with GC

1230 drivers (drought, nitrogen input, drought and nitrogen input (D x N)). Shown are degrees of

1231 freedom (Df), Chi^2 and P-values (P). Significant effects (P < 0.05) are given in bold, marginally

1232 significant effects (P < 0.1) in italics.

D. glomerata						Total bi	omass					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.56	0.456	1	2.22	0.136	1	3.09	0.079	1	0.13	0.715
Soil history	1	6.28	0.012	1	0.76	0.384	1	0.73	0.394	1	<0.01	0.978
Soil treatment	2	1.52	0.467	2	0.94	0.626	2	1.26	0.533	2	0.73	0.693
						Shoot bi	omass					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.02	0.885	1	1.28	0.259	1	3.18	0.075	1	0.22	0.640
Soil history	1	8.27	0.004	1	0.81	0.369	1	0.33	0.567	1	0.15	0.700
Soil treatment	2	3.06	0.216	2	0.44	0.801	2	3.34	0.188	2	0.14	0.932
						Root bio	omass					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	1.40	0.236	1	2.55	0.111	1	1.98	0.160	1	0.04	0.848
Soil history	1	0.90	0.343	1	0.45	0.501	1	0.99	0.319	1	0.21	0.644
Soil treatment	2	2.49	0.288	2	2.06	0.358	2	0.02	0.992	2	3.16	0.206
						Root-sho	ot ratio)				
		Control			Drought			Nitrogen	I		D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	1.65	0.199	1	1.71	0.191	1	0.93	0.335	1	0.01	0.936
Soil history	1	<0.01	0.983	1	0.44	0.505	1	0.43	0.514	1	0.75	0.387
Soil treatment	2	3.14	0.208	2	2.84	0.242	2	0.20	0.906	2	7.72	0.021

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Table S11 Summary of mixed-effect model analyses testing the effects of legacy treatments (plant history, soil history, soil treatment) on plant performance (total biomass, shoot biomass, root biomass

and root-shoot ratio) of *P. trivialis*, when non-treated (control) or treated with GC drivers (drought,

- 1239 nitrogen input, drought and nitrogen input (D x N)). Shown are degrees of freedom (Df), Chi² and P-
- values (P). Significant effects (P < 0.05) are given in bold, marginally significant effects (P < 0.1) in
- 1241 italics. 1242

P. trivialis												
	Control				Drought			Nitrogen		D x N		
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.12	0.732	1	1.25	0.264	1	0.28	0.599	1	0.43	0.513
Soil history	1	0.12	0.731	1	0.14	0.704	1	0.07	0.796	1	0.05	0.826
Soil treatment	2	0.01	0.995	2	1.82	0.404	2	1.69	0.430	2	4.06	0.131
						Shoot bi	iomass					
		Control			Drought			Nitrogen		D x N		
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.01	0.920	1	1.91	0.167	1	0.39	0.532	1	0.01	0.943
Soil history	1	<0.01	0.973	1	0.47	0.492	1	0.46	0.499	1	0.19	0.663
Soil treatment	2	1.34	0.511	2	0.81	0.667	2	1.22	0.545	2	2.96	0.227
						Root bio	omass					
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.21	0.647	1	0.66	0.417	1	1.48	0.224	1	1.45	0.229
Soil history	1	0.33	0.566	1	1.24	0.266	1	0.74	0.389	1	0.03	0.870
Soil treatment	2	1.36	0.506	2	1.10	0.577	2	1.99	0.370	2	5.03	0.081
	Root-shoot ratio											
		Control			Drought			Nitrogen			D x N	
	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р	Df	Chi ²	Р
Plant history	1	0.23	0.630	1	0.14	0.708	1	2.00	0.158	1	2.25	0.134
Soil history	1	0.23	0.630	1	3.19	0.074	1	1.57	0.211	1	0.15	0.697
Soil treatment	2	3.61	0.164	2	0.68	0.711	2	2.16	0.340	2	5.12	0.077

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Drought + N input:









in away-different soil