

Supplementary information

The results of biodiversity–ecosystem functioning experiments are realistic

In the format provided by the authors and unedited

1 **Supplementary Information** for Jochum et al. 2020 Nature Ecology and Evolution paper entitled
2 “The results of biodiversity-ecosystem functioning experiments are realistic”

3 **Supplementary Methods**

4 **1. Details of ecosystem function measurement in the Jena Experiment and BioDIV**

5 **Jena and BioDIV plant aboveground biomass** In Jena, aboveground plant biomass was harvested bi-
6 annually (late May and late August), just prior to mowing. Here, we used only the first harvest, which
7 represents peak standing biomass in most years, from years 2006–2015. All vegetation was clipped at 3
8 cm above ground in up to four rectangles of 0.2 m × 0.5 m per plot with the location of these rectangles
9 being randomly assigned each year. For BioDIV, aboveground peak plant biomass was harvested
10 annually in August by clipping 0.1 m × 6 m strips (see above) each year from 2006–2015. For both
11 studies, harvested target-species biomass was sorted into individual species, dried to constant weight at
12 70 °C for at least 48 h and weighed. Target plant community biomass was then calculated as the sum of
13 the biomass of the individual sown species (g m⁻²).

14 **Jena and BioDIV aboveground plant biomass C:N ratio** In Jena, the combined target species
15 material from the spring biomass harvest (May) was shredded (Analysenmühle, Kinematica, Littau,
16 Switzerland). A subsample of the shredded material was milled to fine powder in a ball-mill (mixer mill
17 MM2000 Retsch, Haan, Germany) and 5–10 mg was used for CN analysis with an elemental analyzer.
18 C and N content were calculated as percentage elemental concentration of dry material and C:N ratios
19 as the ratio between those percentages for years 2007-2012.

20 In BioDIV, two strips of 0.1 m × 6 m were clipped, typically in late July or early August with
21 clip strip locations rotated each year. Unsorted biomass was air-dried at 40 °C. Dried biomass samples
22 were ground (standard Thomas Wiley mill) and the resulting sample homogenized. A sub-sample was

23 re-ground in a Wiley Mini-Mill, stored in glass scintillation vials and re-dried prior to lab analysis.
24 Percent C and N content in dry matter were determined using an elemental analyzer (NA1500, Carlo-
25 Erba Instruments or ECS 4010, COSTECH Analytical Technologies Inc., Valencia, CA, USA) at
26 University of Minnesota or at the Ecosystems Analysis Lab, University of Nebraska, Lincoln. Ratios of
27 dry mass elemental content were then calculated from these results for year 2006.

28 **BioDIVsoil organic C** Soil C samples were taken at all BioDIV plots during summer 2006 at 0–20 cm
29 depth on nine sites per plot ¹. Samples were sieved to remove roots, combined for each plot, mixed and
30 ground. Subsequently, soil samples were dried at 40 °C for 5 days. For each plot, two soil samples were
31 analysed for total C by combustion and gas chromatography (Costech Analytical ECS 4010 instrument,
32 Costech Analytical Technologies Inc., Valencia, CA). Because of the acidic sandy soil at the BioDIV
33 experiment, soil organic C can be considered a close approximation of total C here. We used the
34 average of the two measurements of C in % total carbon of dry weight.

35 **Jena soil organic C** Soil organic C in the Jena “main” experiment was determined in 2008, 2011 and
36 2014. Using a split-tube sampler (4.8 cm diameter), three soil cores per plot were taken to a depth of 30
37 cm ². Soil cores were segmented into 5 cm depth sections and pooled per depth sections and plot. Soil
38 was then dried, sieved and milled. Subsequently, total C was determined by combustion with an
39 elemental analyzer at 1,150 °C (Elementaranalysator vario Max CN, Elementar Analysensysteme
40 GmbH, Hanau, Germany). Because of the calcareous bedrock, Jena soil contains high proportions of
41 inorganic carbon (e.g., in 2014, the proportion of inorganic C was on average 39 % and ranged from 18
42 % to 73 %). To account for this, inorganic C concentration was measured after oxidative removal of
43 organic C for 16 h at 450 °C in a muffle furnace. Finally, organic C concentration was calculated as the

44 difference between total and inorganic C for each 5-cm-layer ² and we averaged over the two
45 uppermost layers to get organic C content for 0–10 cm depth. Subsequently, we averaged over the three
46 samples to get soil organic C content per plot in g kg⁻¹ soil for each year.

47 **Jena and BioDIV root biomass** In Jena, standing root biomass was sampled down to 40 cm depth in
48 all plots in June 2011 and 2014. On each plot, three cores of 3.5 cm diameter were taken and
49 immediately stored at 4 °C until further handling. The total sample was washed to determine root
50 biomass. Bulk samples were carefully washed by hand over a sieve of 0.5 mm mesh size. Remaining
51 soil particles and stones were removed with tweezers. Roots were dried at 60–70 °C and weighed
52 subsequently ³. Unit: g m⁻²

53 In BioDIV, root biomass was sampled in 2010 after aboveground biomass clipping by collecting
54 three 5 cm diameter × 30 cm depth cores per clipped strip ¹. Roots were washed free of soil, sorted
55 from other organic material, dried and weighed. Unit: g m⁻²

56 **Jena herbivory rate** In Jena, invertebrate herbivory rates were assessed as proportional damage for
57 every plant species × plot-combination. Herbivory rates of individual plant species were used to
58 calculate community herbivory rates based on four different types of invertebrate herbivory: chewing,
59 rasping, sap sucking and leaf mining. Samples of the Jena biomass harvest were used after sorting to
60 species. For a maximum of 20 randomly chosen leaves per plant species, damage area was estimated in
61 mm² as total value of the four damage types and total leaf area of every leaf was measured with an area
62 meter (LI-3000C Area Meter equipped with a LI3050C transparent belt conveyor accessory, LI-COR
63 Biosciences, Lincoln, USA). For details on the methods used see ⁴. Here, we used percentage herbivory
64 of the target species community from the late harvest, as this was available for three years from 2010–

65 2012. Unit: % damage

66 **Jena soil microbial biomass C** Soil sampling and measurement of basal and substrate-induced
67 microbial respiration with an oxygen-consumption apparatus was done on each plot in September 2010
68 ⁵. Oxygen consumption of soil microorganisms in a fresh-soil equivalent to 3.5 g dry weight was
69 measured at 22 °C. Substrate-induced respiration was determined by adding D-glucose to saturate
70 catabolic enzymes of microorganisms according to preliminary studies (4 mg g⁻¹ dry soil solved in 400
71 µl deionized water; ^{6,7}). Maximum initial respiratory response (µl O₂ g⁻¹ dry soil h⁻¹) was calculated as
72 mean of the lowest three oxygen consumption values within the first 10 h after glucose addition.
73 Microbial biomass C (µg C g⁻¹ dry soil) was calculated as 38 × maximum initial respiratory response as
74 suggested by preliminary studies ⁸. Previous work has shown that the 2010 microbial biomass data are
75 representative for long-term plant diversity effects ⁷.

76 **Jena phosphatase activity** Nine soil cores (diam. 2 cm, 0–5 cm depth) were combined to one
77 composite sample per plot to assess phosphatase activity in 2013 ⁹. Because of the alkaline pH of the
78 soil, we measured alkaline phosphomonoesterase activity (phosphatase activity) according to the assay
79 by ¹⁰. For each soil sample, one replicate and one blank value were included. One gram of field moist
80 soil was mixed with toluene, modified universal buffer (MUB) and p-nitrophenylphosphate (pNP), and
81 incubated at 37 °C for 1 hour. Subsequently, we added CaCl₂ and NaOH. To blanks, pNP was added
82 after incubation. The solution was filtered through P-free filters (MN 619 G ¼, Macherey-Nagel GmbH
83 & Co. KG, Düren, Germany). Directly after filtration, pNP concentrations [µg ml⁻¹] were measured at
84 400 nm with a spectrophotometer (PU 8675 VIS spectrophotometer, Philips GmbH, Hamburg,
85 Germany). The soil moisture was determined gravimetrically, i.e. by weighing before and after drying

86 at 105 °C to convert phosphatase activities to dry matter ($\mu\text{g pNP g}^{-1} \text{ h}^{-1}$).

87 **Jena pollinator abundance** In 2010 and 2012, hymenopterans were sampled by suction sampling
88 using a modified commercial vacuum cleaner (Kärcher A2500, Kärcher GmbH, Winnenden, Germany).
89 In each year, within each plot, two random subplots of 0.75 m x 0.75 m were chosen, covered with a
90 gauze-coated cage of the same size, and arthropods within cages were sampled. The sampling was
91 carried out between 9 a.m. and 4 p.m. within two 4-day sampling periods. The overall abundance of
92 hymenopterans across the two samples per plot was used as a proxy of pollinator abundance and thus
93 potential for pollination on each plot in the respective year. Unit: number of individuals

94 **2. Processing TRY and other plant-trait data to generate species-level values**

95 For each of the geographical species subsets, TRY trait data were processed separately following a
96 standardized protocol: i) Removal of duplicate observations (e.g. duplicate entries of leaf mass from the
97 same individual). ii) Removal of non-open data and removal of data obtained from outside the
98 respective target continents. iii) Calculation of outliers for each trait-species combination (trait mean
99 ± 1.96 SD as outlier definition). iv) Removal of observations with TRY ErrorRisk > 4 . v) Averaging
100 over trait-species values per TRY dataset. vi) Removal of TRY datasets with more than 5% of values
101 identified as outliers. vii) Averaging over trait-species mean values of the remaining datasets. For the
102 US species, TRY data was combined with additional trait data collected in naturally occurring
103 polycultures at Cedar Creek (personal communication with J.A. Catford¹¹ now available on TRY as
104 dataset 354, P.B. Reich, J. Cavender-Bares). These Cedar Creek trait averages per dataset were
105 included into the averaging process at step v). Finally, trait values of synonyms and accepted species
106 names were averaged and assigned to the accepted plant-species names where necessary.

107 **Supplementary Information on sensitivity analyses I.**

108 To test how robust our results are to key methodological decisions in our analysis, we performed a
109 number of sensitivity analyses. Specifically, we tested how methodological decisions regarding the
110 PCA analysis of comparing plant-community properties affected the number and identity of
111 experimental plots selected as realistic. These methodological choices include the selection of
112 community properties entering the PCA, the method of calculating multidimensional overlap, and a set
113 of other methodological details such as, for example, the number of PCA axes used for calculating
114 multidimensional overlap or the choice of real-world datasets (see below). If not noted otherwise, we
115 used the 12 vif-selected community properties and the convex-hull volume method to calculate
116 intersections (main analysis) for both the German and US comparison. The different analyses are
117 described below. Please refer to Supplementary Tables S2 and S3 below for a comparison of selected
118 realistic plots for the main analysis and the sensitivity analyses presented here.

119 We assessed the impact of using different subsets of community properties in the PCAs by re-running
120 our analysis with the following subsets: i) all 21 community properties available (“All21”), ii) the 12
121 vif-selected community properties (“Full12”, main analysis), iii) four subsets defined by excluding one
122 class of community properties (taxonomic, phylogenetic, functional diversity, or CWM functional
123 traits; called “Allbuttax”, “Allbutphyl”, “Allbutfun”, and “AllbutCWM”), and iv) all 21 properties
124 except the 5 properties showing the biggest differences between experimental and real-world datasets
125 in the two regions (“remove5”, sensitivity A), respectively. Supplementary Table 1 below summarizes
126 which community properties are used in which analysis.

127 To test what impact the methodology used for calculating multidimensional overlap had on our
128 findings, we combined the above-described community-property subsets with three overlap calculation
129 methods: i) three-dimensional convex hull volumes (main analysis), ii) three-dimensional

130 hypervolumes, and iii) two-dimensional 95% confidence interval ellipses (see main text methods for
131 details).

132 Performing all possible combinations of subsets and methods would lead to an unmanageable number
133 of results and, given the relative robustness of our results to most methodological decisions, seems
134 unnecessary. Hence, we present results for a selection of combinations. For two community-property
135 subsets, namely the All21 and the Full12 subsets, all three overlap calculation methods were run. For
136 the remaining subsets, only the convex hull volume method was used. The resulting overlap (realistic
137 plots) is presented for all these combinations in Supplementary Tables S2 and S3. Alternative versions
138 of Fig. 1 (PCA and overlap calculation) are shown in Supplementary Fig. 1 for all above-described
139 combinations. Alternative versions of Fig. 2 (BEF) relationships are shown in Supplementary Fig. 3 for
140 all combinations of the All21 and Full12 subsets and the three overlap calculation methods and
141 additionally for the species-abundance based NMDS, for the USA dataset.

142 Below, we provide additional details for these analyses and succinctly summarize their results.

143 **Full12:** Here, we used the same community property subset as in the main analysis (see Supplementary
144 Table 1), but tested two additional overlap calculation methods. For the Jena Experiment, this resulted
145 in 24 and 21 plots to be chosen as realistic (instead of 23) with 91 and 87 % of overlap with the main
146 analysis realistic subset for the “hyper” and “ellipse” method, respectively (Supplementary Table 2 and
147 Supplementary Fig. 1). For BioDIV, 109 and 133 plots were deemed realistic, with 87 and 96 %
148 overlap with the main analysis realistic subset, for the “hyper” and “ellipse” methods, respectively
149 (Supplementary Table 3 and Supplementary Fig. 1). For the BEF relationships (Supplementary Fig. 3),
150 these changes resulted in a slight increase of the number of relationships that changed significantly
151 after subsetting (confidence intervals not containing each other’s slope estimate) from the
152 unconstrained to the constrained dataset (3 and 4 BEF relationships compared to 2 in the main analysis

153 for “hyper” and “ellipse”, respectively).

154 **All 21:** Here, instead of removing community properties based on the vif-based selection of
155 multicollinear variables, all 21 community properties entered the PCA's. For the Jena Experiment, this
156 resulted in 33, 32, and 34 of 82 (40, 39, and 41 %) plots to be selected as realistic which included 91,
157 96, and 96 % of the plots selected as realistic in the main analysis for the “chull”, “hyper”, and
158 “ellipse” methods, respectively. For BioDIV, 121, 104, and 136 of 159 (76, 65, and 86 %) were selected
159 as realistic, containing 95, 85, and 91 % of the main analysis realistic plots for the “chull”, “hyper”, and
160 “ellipse” methods, respectively. For the BEF relationships, these slight changes translated into 2, 1 and
161 1 BEF relationships changing significantly between the unconstrained and the constrained dataset
162 (compared to 2 changes in the main analysis) for the “chull”, “hyper”, and “ellipse” methods,
163 respectively (see Supplementary Fig. 3 for details). Given this very strong similarity to the main
164 analysis, we conclude that using all 21 community properties would not change our conclusions. Thus,
165 we stick to choosing community properties based on the vif selection to reduce multicollinearity among
166 variables entering the PCA.

167 **Allbuttax:** Here, we used all community properties except for the taxonomic diversity properties (see
168 Supplementary Table 1). For the Jena Experiment, 44 instead of 23 of 82 plots were deemed realistic,
169 including all plots chosen as realistic in the main analysis. For BioDIV, 122 instead of 121 of 159 plots
170 were deemed realistic, including 94 % of the ones chosen as realistic in the main analysis.

171 **Allbutphyl:** Here, we used all community properties except for the phylogenetic diversity properties
172 (see Supplementary Table 1). For the Jena Experiment, 41 instead of 23 of 82 plots were deemed

173 realistic, including all plots chosen as realistic in the main analysis. For BioDIV, 125 instead of 121 of
174 159 plots were deemed realistic, including 96 % of the ones chosen as realistic in the main analysis.

175 **Allbutfun:** Here, we used all community properties except for the functional diversity properties (see
176 Supplementary Table 1). For the Jena Experiment, 30 instead of 23 of 82 plots were deemed realistic,
177 including 96% of the plots chosen as realistic in the main analysis. For BioDIV, 124 instead of 121 of
178 159 plots were deemed realistic, including 96 % of the ones chosen as realistic in the main analysis.

179 **AllbutCWM:** Here, we used all community properties except for the CWM functional traits (see
180 Supplementary Table 1). For the Jena Experiment, 19 instead of 23 of 82 plots were deemed realistic,
181 including 83 % of the plots chosen as realistic in the main analysis. For BioDIV, 126 instead of 121 of
182 159 plots were deemed realistic, including 90 % of the ones chosen as realistic in the main analysis.

183 Below, we describe the additional sensitivity analyses (A-F) for which results are shown in
184 Supplementary Table 1, but not in Supplementary Fig. 1 and Fig. 3.

185 **A: Remove only 5 community properties:** Here, we removed community properties based on the
186 proportion of biodiversity experiment communities falling within the real-world range of those
187 properties (see Supplementary Tables S4 and S5). As the similarity of biodiversity experiment and real-
188 world communities regarding these properties differed between the German and US dataset, we
189 removed different variables for these two data sets. For each dataset, we removed the 5 community
190 properties with the lowest proportion of biodiversity experiment communities falling within the real-
191 world range for these properties. For the German PCA, we removed SEve, S, PD, FRic, and MNTD.

192 For the US PCA; we removed LDMC, FEve, SLA, leaf N, and FRic. For the Jena Experiment, this
193 resulted in 49 of 82 (60 %) plots to be selected as realistic with all of the main analysis realistic plots
194 included in this subset. For BioDIV, 112 of 159 (70 %) were selected as realistic, containing 83 % of
195 the main analysis realistic plots. Given this very strong overlap with the main analysis, we conclude
196 that using this subset of community properties would not change our conclusions. Thus, we stick to
197 choosing community properties based on the vif selection to reduce multicollinearity among variables
198 entering the PCA.

199 **B: Change number of axes in PCA used for convex hull intersection:** In the main analysis, we used
200 the first three PCA axes for 3-dimensional convex hull and hypervolumes to determine the intersection
201 between biodiversity experiment and real-world communities in multidimensional space (see Extended
202 Data Fig. 6 for an overview of the variance explained by all PCA axes). Here, we changed this number
203 of axes to the number of axes explaining above-average variance. All axes explaining more than 1/12
204 (above average) of the total variance were included. For the German dataset, these were the first five
205 axes, explaining a summed total of 73 % of the total variance (compared to the first three axes
206 explaining 53 %). When using these five axes, 19 instead of 23 (of 82, 23 %) Jena Experiment plots
207 were selected as realistic (Supplementary Table 2 and 3), containing 83 % of the plots selected as
208 realistic in the main analysis. For the US dataset, it was the first four axes, together explaining 64 % of
209 the total variance (compared to the first three axes explaining 55 %). Due to an R “geometry” error in
210 the convex hull calculation when using four axes in the US dataset (most likely some PCA points too
211 close to each other), the intersection could not be calculated. This was probably caused by a few
212 communities being too similar (nearly coincident points) in these four axes. The error did not occur at
213 three axes and is more likely to occur at 4-D and more according to the package documentation. Given

214 the broadly comparable number and identity of realistic plots between the 2-D ellipse, 3-D convex hull
215 and 3-D hypervolume intersection calculation (see Supplementary Table 3), we assume that adding the
216 fourth dimension (explaining 9 % of the total variance) to the convex hull is unlikely to heavily
217 influence the selection of realistic plots for BioDIV. Overall, although changing the number of axes
218 changed the number and identity of selected realistic plots, there is still a strong overlap of realistic
219 plots selected by this sensitivity analysis and the main analysis, so we conclude that, qualitatively, our
220 results would not change.

221 **C: Use all available non-experiment plots as real world rather than just the most**

222 **methodologically comparable:** In the main analysis, we used only a subset of the available real-world
223 datasets to identify realistic biodiversity experiment plots in the intersection calculations. This decision
224 was made because the vegetation survey methodology of some real-world datasets differed quite
225 substantially from the methodology in the biodiversity experiments (transects, subplots, very different
226 vegetation survey area). Here, we tested the impact of using all available real-world datasets (see
227 Figure 1 a and e) in the intersection calculation on the selection of realistic experimental plots. For the
228 Jena Experiment, this resulted in 42 of 82 (52 %) plots to be selected as realistic with all of the main
229 analysis realistic plots included in this subset. For BioDIV, 122 of 159 (77 %) were selected as realistic,
230 containing all of the main analysis realistic plots. Looking at Figure 1 a and e, these numbers could
231 have been expected given that, in the German PCA, for example the Jena invasion communities are
232 shifted towards the real-world communities relative to the Jena Experiment main communities. In the
233 US comparison, however, most the additional real-world communities (Old field succession
234 chronosequence and Oak savannah communities) do not extend further towards the real-world
235 communities than the Fertilization 1 and 2 communities already used in the main analysis. Given this

236 very strong overlap with the main analysis, we conclude that using all real-world data sets available
237 would not change our conclusions. Thus, we stick to only using the most comparable datasets in our
238 main analysis.

239 **D: Change overlap criterion, i.e. explore number of years in intersection for each plot:** In the main
240 analysis, we selected biodiversity experiment plots as realistic if their plant communities fell in the
241 PCA-based intersection of experiment and real-world data in at least one year. Here, we show for how
242 many years which of the “realistic” plots had plant communities falling within the intersection.
243 For the Jena Experiment, 6 of 23 realistic plots were included in all years, all but three plots were at
244 least included in more than one year. For BioDIV, 74 of 122 realistic plots were included in all years,
245 and another 21 plots were included in all but one year. All other plots were included in fewer years.
246 This indicates that, for BioDIV, changing the criteria for being defined realistic (e.g. having at least 50
247 % of the annual communities in the intersection) would not change the conclusions a lot. In Jena,
248 however, choosing e.g. 50 % of annual communities as a threshold would lead to only very few plots
249 being selected as realistic. The vast majority of main-analysis realistic plots are not only included in
250 only a single year. Consequently, even the most realistic Jena Experiment plots are only real-world
251 comparable in a subset of all years considered here. However, these are the most real-world comparable
252 subset of the Jena Experiment plots. Our aim was to remove the most unrealistic plots from each
253 experiment and then compare BEF relationships. While it is interesting to see how the two biodiversity
254 experiments differ in their relation to the real-world communities, comparing BEF relationships is only
255 really possible if there are enough plots for each BEF model to fit a relatively reliable model. We
256 conclude that choosing a different threshold here would reduce the number of realistic plots, but mainly
257 impact our analysis by not providing enough data points to fit the Jena BEF relationships for the subset

258 of realistic plots only.

259 **E: Calculate realistic plots using species-abundance NMDS instead of community-property**

260 **PCA's:** In the main analysis, selection of realistic experimental plots is always based on the
261 community-property PCA's. Here, we test how the selection of realistic plots changes when using the
262 first three axes of a species-abundance based NMDS (function "metaMDS" in R package "vegan"¹²).
263 As visualized in the NMDS biplots of Supplementary Figure 2, the plant community composition of
264 biodiversity experiments and real-world systems differs far more in the German than in the US
265 comparison. In the US comparison, there is a good overlap in plant community NMDS space, resulting
266 in 53 of the 159 plots (33 %) to be selected as realistic, containing 43 % of plots selected as realistic
267 based on the community property PCA. In the US comparison, the biodiversity experiment and real-
268 world plots are all situated in a relatively small area, thus we expect some similarity in species
269 composition even though the management of these sites differs. Constraining data to only those 53
270 plots led to two of the four US BEF relationships changing significantly from the unconstrained to the
271 constrained dataset, but in both cases, the slopes of the realistic subsets were higher (see
272 Supplementary Fig. 3), thus suggesting that experiments might underestimate BEF relationships for
273 certain functions and thus strengthening our conclusion that experimental findings are generally robust.
274 In the German comparison, the Jena Experiment community NMDS space contains the real-world
275 plots, but, interestingly, very few Jena Experiment plots actually fall within the real-world NMDS
276 space (see Supplementary Figure 2). As such, only 2 of the Jena Experiment plots (2 %) would be
277 selected as realistic based on this methodology. This indeed is a very low percentage of realistic sites,
278 but this is not unexpected. The German real-world plots are spread out across three geographically
279 distant regions across Germany with a strong gradient of land-use intensity and the Jena Experiment

280 was established in a single old field site with (obviously and intentionally) narrow environmental
281 gradients and plant species being selected from a deliberately regional species pool fitting for the given
282 environmental and geographical context. Consequently, we never expected the plant communities
283 themselves to be very similar in their composition. This is exactly the reason why our analysis is based
284 on comparing the properties of these plant communities irrespective of the identity of species involved.

285 **F: Only Germany: Use only real-world plots with management similar to Jena Experiment:** In the
286 main analysis, we compare plant communities from biodiversity experiments to those of real-world
287 plots covering a broad gradient of land-use intensities, especially in the German dataset (Biodiversity
288 Exploratories). In addition to visualizing the different land-use intensity and its components (mowing,
289 grazing, fertilization) in an alternative version of Figure 1 (Supplementary Figure 5), we tested the
290 impact of using only Biodiversity Exploratories plots with land use roughly comparable to the Jena-
291 Experiment for the German comparison. The Jena Experiment plots are mown twice a year and neither
292 fertilized nor grazed. Specifically, we removed all Biodiversity Exploratories plots that are grazed or
293 fertilized and those that are never mown, leaving only the mown (but neither grazed nor fertilized)
294 plots in the analysis. This procedure resulted in only 10 Exploratories plots left in the dataset (9 plots
295 from the North-East region Schorfheide-Chorin and one plot from the South-West region Schwäbische
296 Alb). Expectedly, this resulted in only a very small subset of the Jena Experiment plots being selected
297 as realistic, namely 10 of 82 plots (12 %), containing 43 % of the originally selected realistic plots. The
298 aim of our study was to compare the biodiversity experiments to real-world grasslands in the same
299 countries. These real-world grasslands are managed differently than the biodiversity experiments. Still,
300 we find experimental plots with plant communities comparable in their properties to those of the real-
301 world plots. While it would be interesting to compare the Jena Experiment to real-world plots

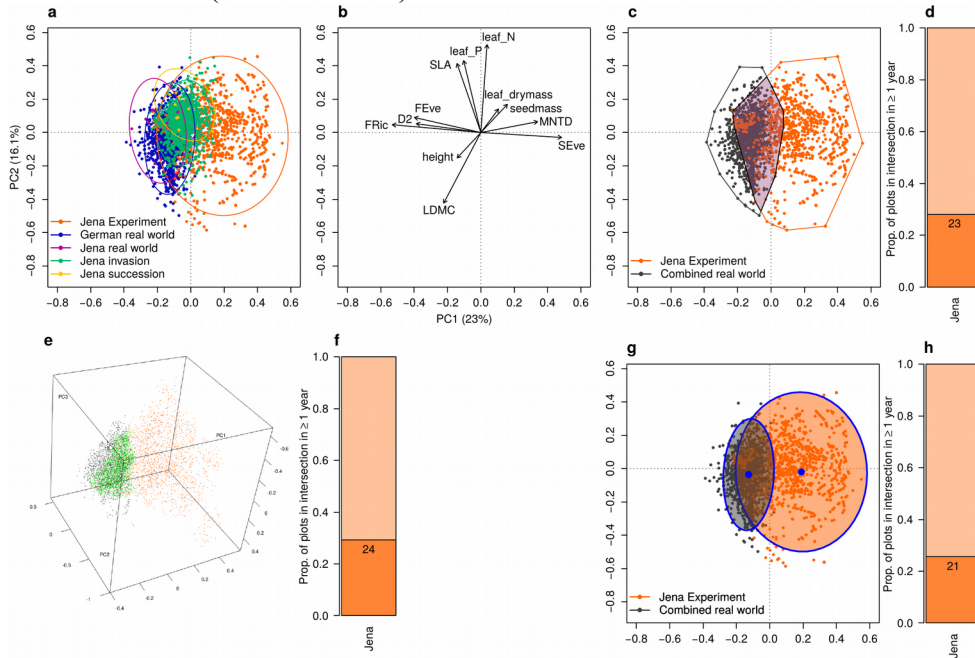
302 undergoing similar land use, this is not feasible with our dataset and we thus stick to using all real
303 world datasets sampled in a relatively comparable manner to the biodiversity experiments.

304 Please note that, for the US data, when leaving all 21 or keeping 16 (removing the five most different)
305 community properties in the PCA, results of the intersection calculation should be taken with caution
306 due to a slight onset of horseshoe-like patterns in the PCA. We take this as another reason to stick to
307 our vif-based selection of community properties to enter the PCA for the main analysis.

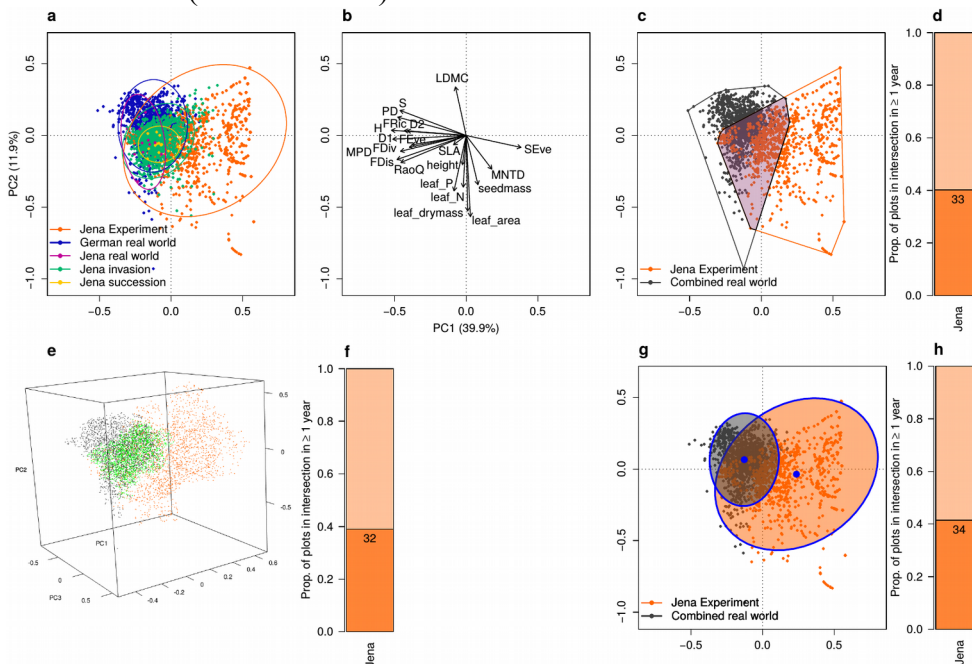
308 **Supplementary Table 1.** Community properties used in the different main and sensitivity subsets.
309 Column “Class” specifies which class of community properties a given property belongs to (taxonomic,
310 phylogenetic, and functional diversity, and CWM functional traits). “Full 12 vif” is the vif-selected
311 subset used in the main analysis. The other subsets are used in sensitivity analyses.

Class		All 21	Full 12 vif	All but tax	All but phyl	All but fun	All but CWM	GER remove 5	USA remove 5
fun	FRic		1	1	1	0	1	0	0
fun	FEve		1	1	1	0	1	1	0
fun	FDiv		0	1	1	0	1	1	1
fun	FDis		0	1	1	0	1	1	1
fun	RaoQ		0	1	1	0	1	1	1
CWM	leaf_area		0	1	1	1	0	1	1
CWM	SLA		1	1	1	1	0	1	0
CWM	leaf_drymass		1	1	1	1	0	1	1
CWM	LDMC		1	1	1	1	0	1	0
CWM	leaf_N		1	1	1	1	0	1	0
CWM	leaf_P		1	1	1	1	0	1	1
CWM	height		1	1	1	1	0	1	1
CWM	seedmass		1	1	1	1	0	1	1
tax	S		0	0	1	1	1	0	1
tax	H		0	0	1	1	1	1	1
tax	D1		0	0	1	1	1	1	1
tax	D2		1	0	1	1	1	1	1
tax	SEve		1	0	1	1	1	0	1
phyl	PD		0	1	0	1	1	0	1
phyl	MPD		0	1	0	1	1	1	1
phyl	MNTD		1	1	0	1	1	0	1

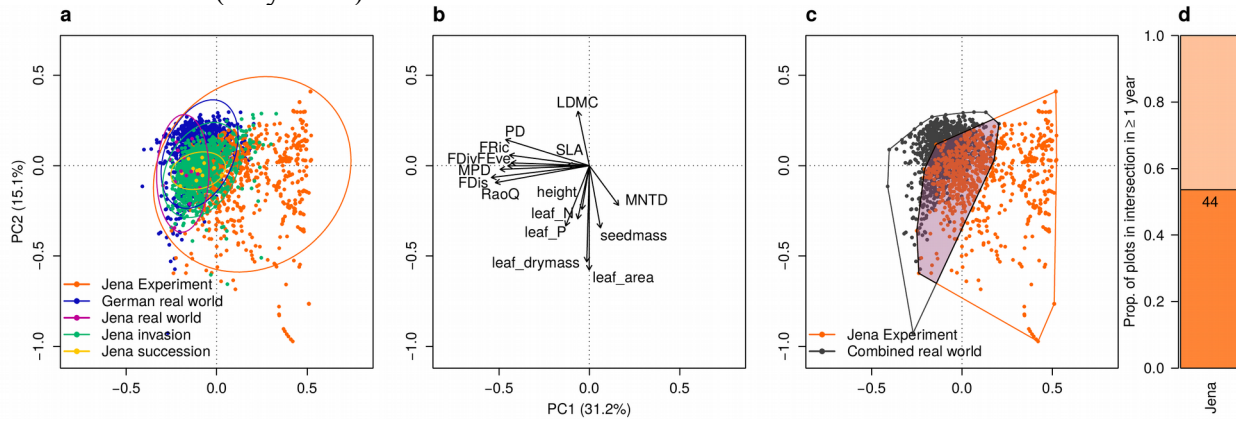
312 **Supplementary Figure 1.** Alternative versions of Fig. 1 based on alternative intersection scenarios
 313 (community- property subsets and overlap calculation methods). a & b: PCA and factor loadings; c &
 314 d: 3D convex hull volume, e & f: 3D hypervolume, g & h: 2D ellipse. 12 different versions: For subsets
 315 Full12 (vif selected) and All21, three methods are shown. For subsets excluding one of the community-
 316 property classes (Allbuttax = all properties except taxonomic div. properties), only the convex hull
 317 method is presented. All scenario's shown for both geographical datasets. See Supplementary Table 1
 318 for community properties included per subset and Supplementary Fig. 3 and Supplementary Tables 2
 319 and 3 for overlap for species-abundance based NMDS.
 320 German Full12 (three methods):



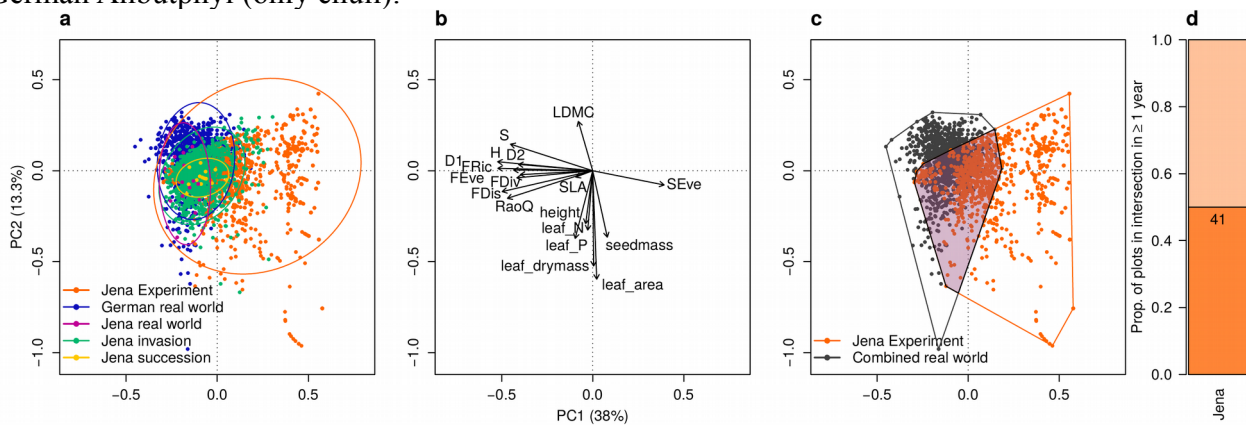
321 German All21 (three methods):



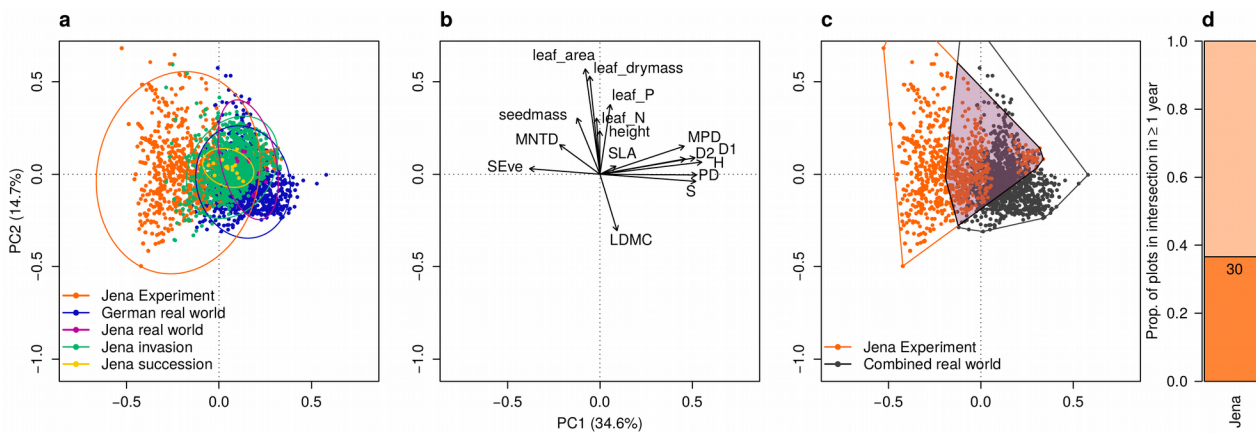
322 German Allbuttax (only chull):



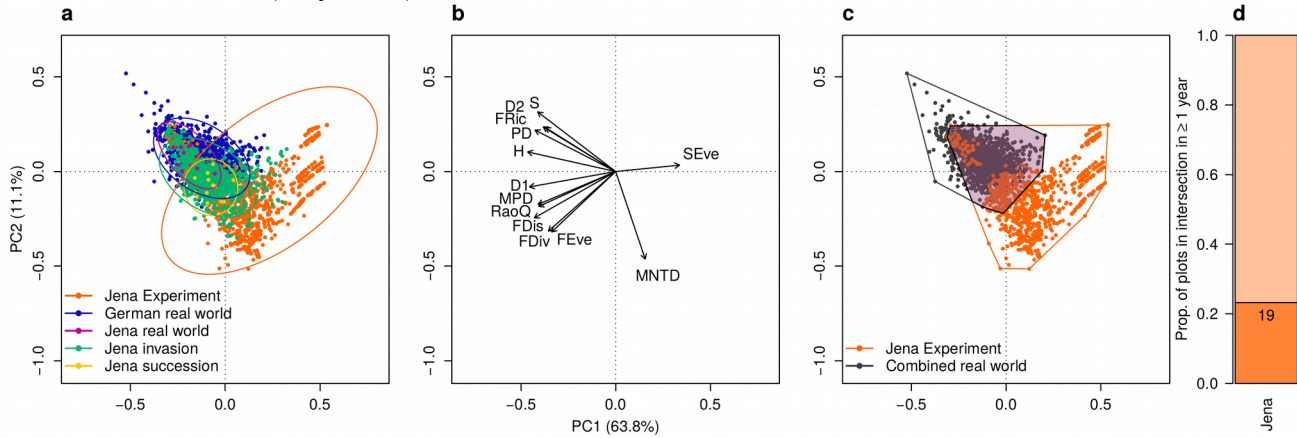
323 German Allbutphyl (only chull):



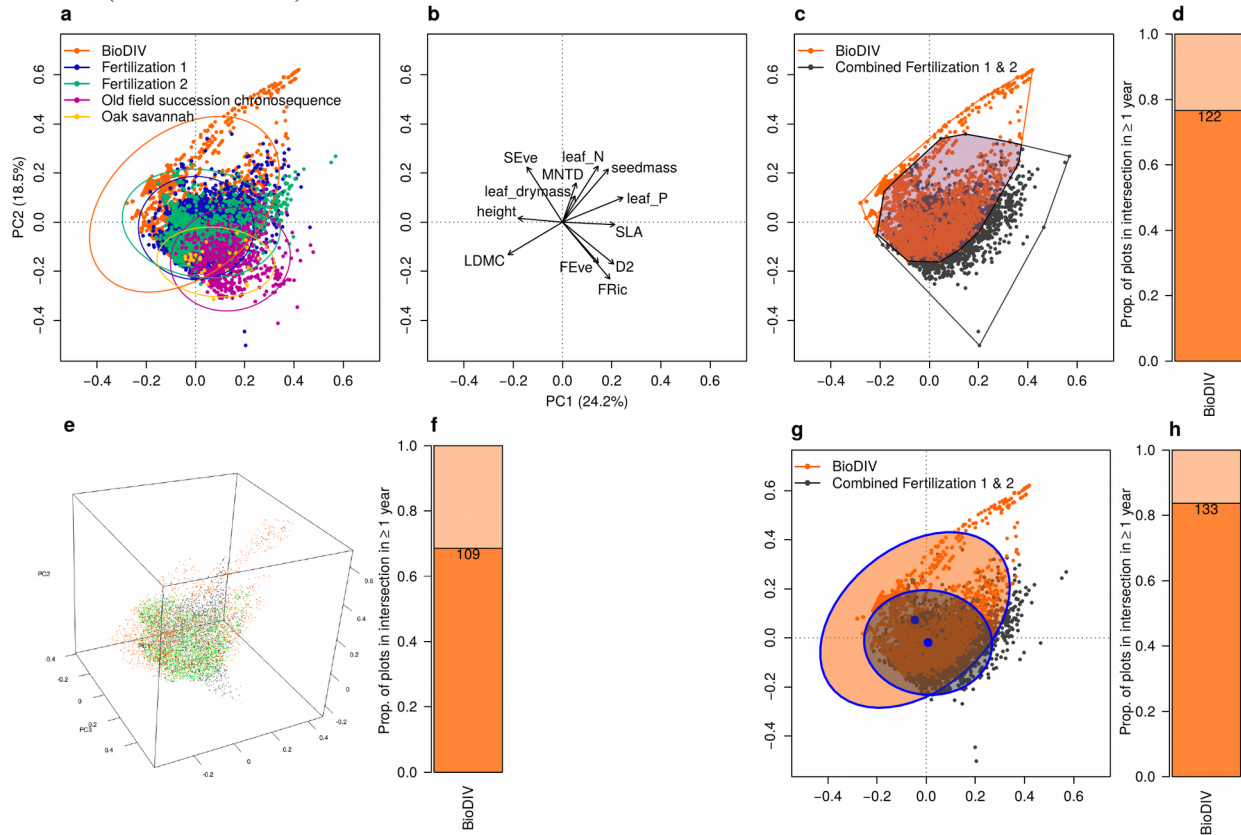
324 German Allbutfun (only chull):



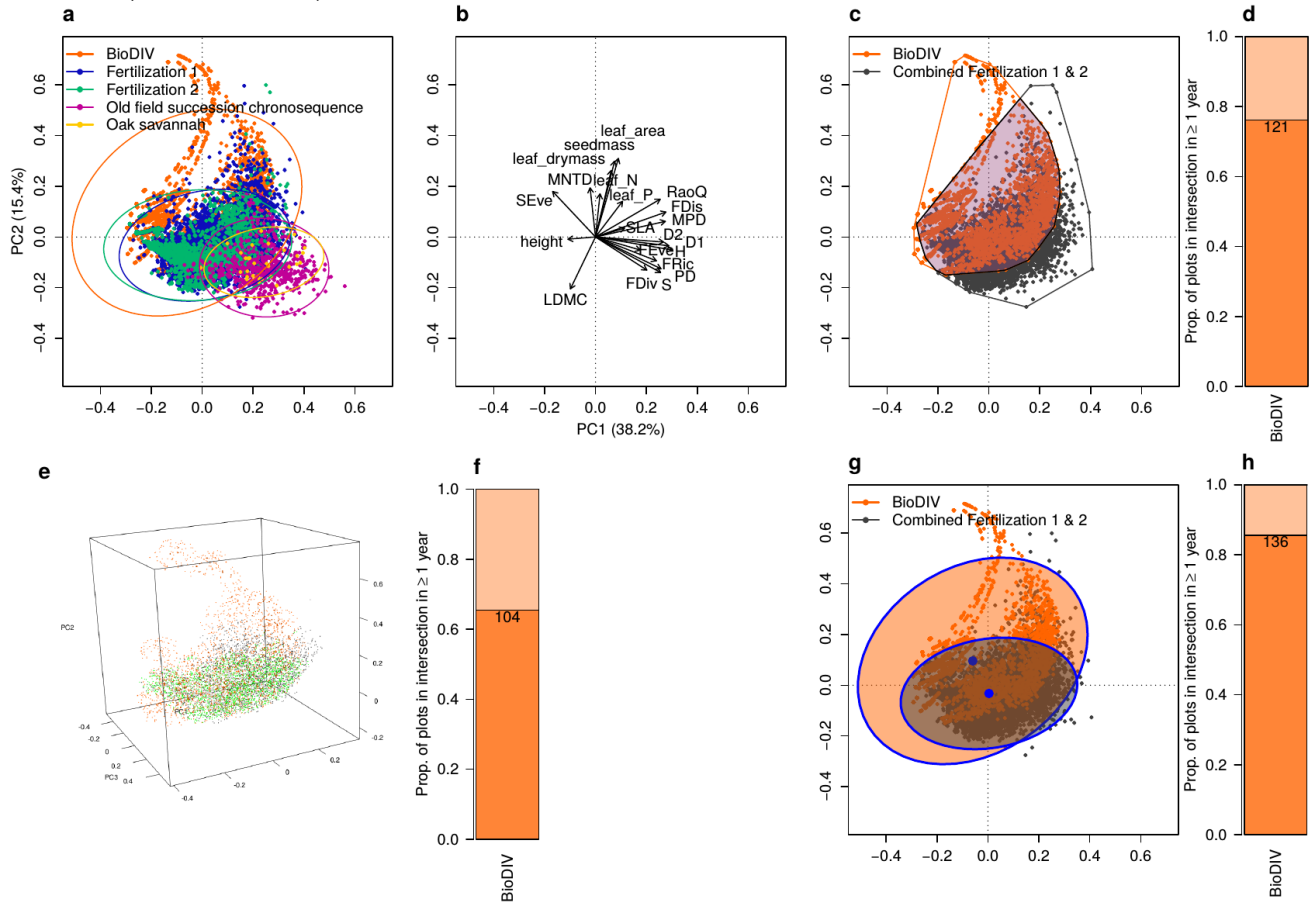
325 German AllbutCWM (only chull):



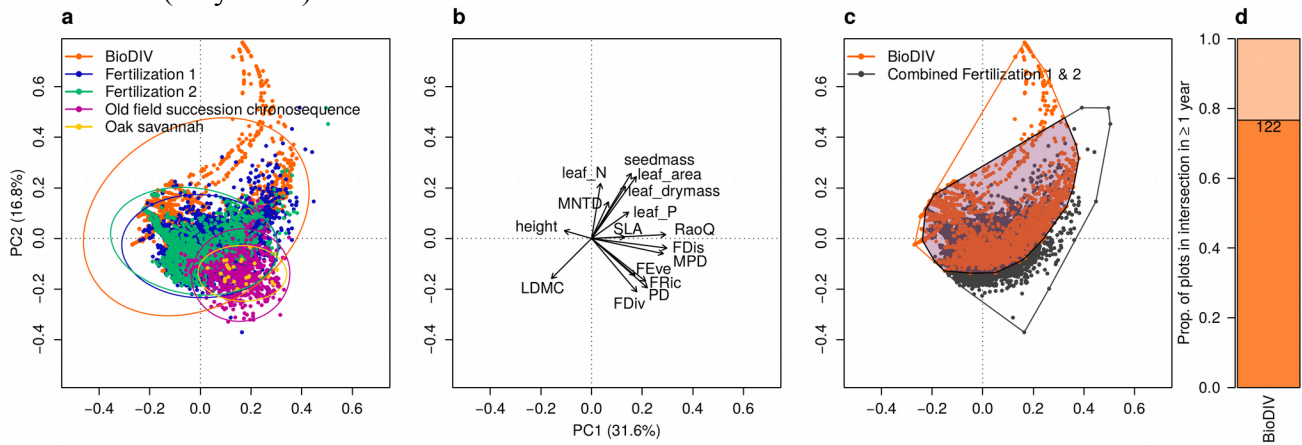
326 US Full12 (three methods):



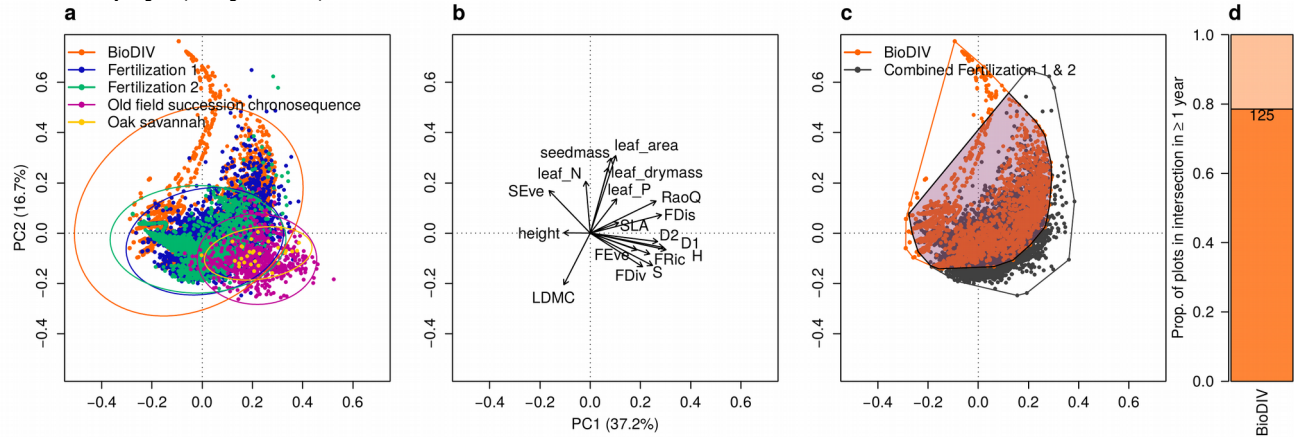
327 US All21 (three methods):



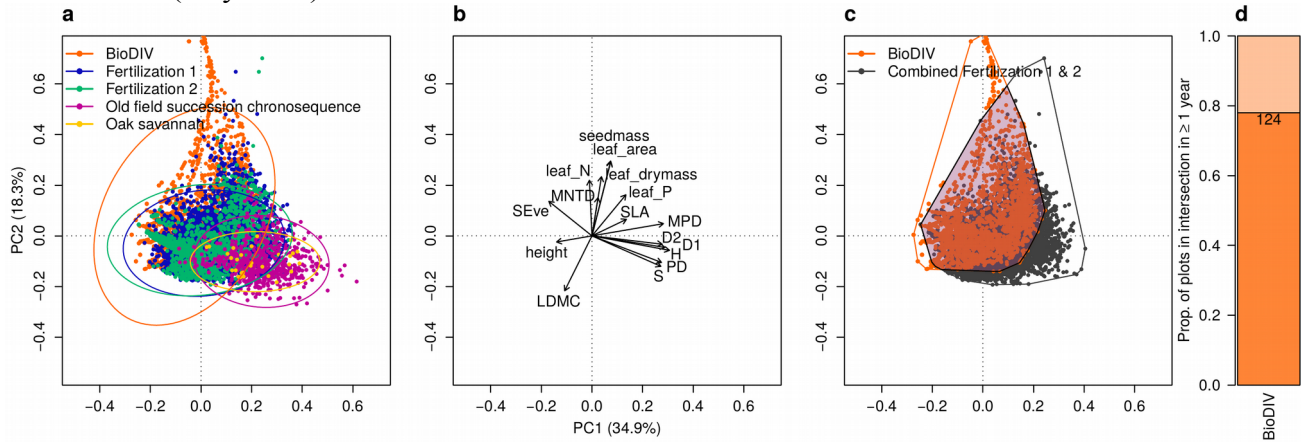
328 US Allbuttax (only chull):



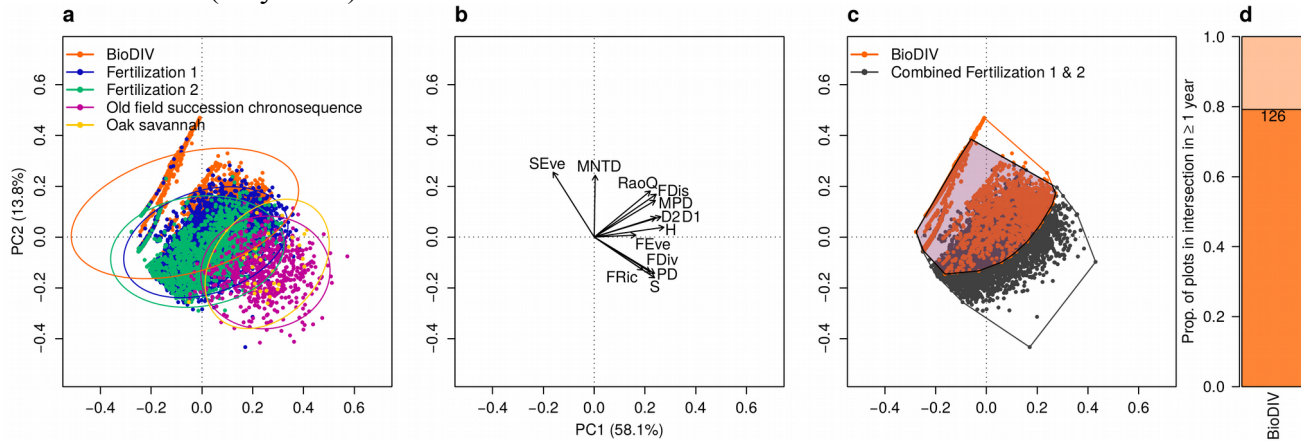
329 US Allbutphyl (only chull):



330 US Allbutfun (only chull):



331 US AllbutCWM (only chull):



332 **Supplementary Table 2.** Jena Experiment plots included in the different sensitivity overlap scenarios
333 vs. all experimental plots. Jena Experiment plots with their sown diversity (sown_div), number of
334 functional groups (num_fg) and their selection as realistic plots (1) based on various sensitivity
335 analyses described in Supplementary Information on sensitivity analyses I and Supplementary Table 1.
336 The selection of “realistic” plots is compared to the main analysis (Full12_chull). Note that column D
337 shows the number of years with communities falling inside the intersection of 3-D convex hull volumes
338 (13 = all years). The last four rows show the number of chosen realistic plots (sum), the percentage of
339 total Jena Experiment plots (82) chosen as realistic (percent_tot), if there are more plots selected as
340 realistic than in the main analysis (yes, same, or no), and the proportion of realistic plots chosen in the
341 main analysis (Full12_chull) also selected in each sensitivity analysis (percent_12vif). Plots are sorted
342 by sown diversity levels.

plot	sown_div	num_fg	Full12_chull	Full12_hyper	Full12_ellipse	AI121_chull	AI121_hyper	AI121_ellipse	Albutax_chull	Albutphyl_chull	Albutfun_chull	AlbutCWM_chull	A_removes5	B_moreaxes	C_allreal	D_count	E_NMDS	F_LUrem
B1A22	60	4	1	1	1	1	1	1	1	1	1	1	1	1	1	13	0	1
B2A03	60	4	1	1	1	1	1	1	1	1	1	1	1	1	1	13	0	1
B3A14	60	4	1	1	1	1	1	1	1	1	1	1	1	1	1	13	0	1
B4A01	60	4	1	1	1	1	1	1	1	1	1	1	1	1	1	13	0	1
B1A01	16	4	1	1	1	1	1	1	1	1	1	1	1	1	1	3	0	0
B1A06	16	2	1	1	1	1	1	1	1	1	1	1	1	1	1	6	0	0
B1A11	16	1	1	1	1	1	1	1	1	1	1	1	1	1	1	2	0	0
B1A20	16	3	1	0	0	1	1	1	1	1	1	1	1	0	1	1	0	0
B2A10	16	2	1	1	1	1	1	1	1	1	1	1	1	1	1	13	0	1
B2A18	16	4	1	1	1	1	1	1	1	1	1	1	1	1	1	8	0	1
B2A22	16	3	1	1	1	1	1	1	1	1	1	1	1	1	1	7	0	0
B3A09	16	1	1	1	1	1	1	1	1	1	1	1	1	1	1	11	1	1
B3A16	16	2	1	0	0	1	1	1	1	1	1	1	1	0	1	1	0	0
B3A22	16	4	1	1	1	1	1	1	1	1	1	1	1	1	1	6	0	0
B3A24	16	3	1	1	1	1	1	1	1	1	1	1	1	0	1	12	0	0
B4A02	16	3	1	1	1	1	1	1	1	1	1	1	1	1	1	6	0	0
B4A18	16	4	1	1	1	1	1	1	1	1	1	1	1	1	1	2	0	0
B4A20	16	2	1	1	1	1	1	1	1	1	1	1	1	0	1	4	0	1
B1A02	8	2	1	1	1	1	1	1	1	1	1	1	1	1	1	3	0	1
B1A03	8	3	0	1	0	1	1	1	1	1	0	1	0	1	0	0	0	0
B1A12	8	1	0	0	0	0	0	0	1	1	0	0	1	0	1	0	0	0
B1A14	8	4	0	1	0	1	1	1	1	1	1	0	1	0	1	0	0	0
B2A12	8	1	0	0	0	1	1	1	1	1	1	0	1	0	1	0	0	0
B2A14	8	4	0	0	0	1	1	1	1	1	1	0	1	0	1	0	0	0

B2A17	8	2	0	0	0	0	0	1	1	1	0	0	1	0	1	0	0	0
B2A21	8	3	0	0	0	1	1	1	1	1	1	0	1	0	1	0	0	0
B3A04	8	1	1	1	1	1	1	1	1	1	1	0	1	1	1	13	1	1
B3A05	8	3	1	1	1	1	1	1	1	1	1	0	1	1	1	3	0	0
B3A07	8	4	0	0	0	1	0	0	1	1	1	0	1	0	1	0	0	0
B3A20	8	2	0	0	0	0	0	0	1	1	0	0	1	0	1	0	0	0
B4A06	8	1	0	0	0	0	0	1	0	0	0	0	1	0	1	0	0	0
B4A08	8	2	0	1	0	1	1	1	1	1	1	0	1	0	1	0	0	0
B4A10	8	3	0	0	0	1	1	1	1	1	1	0	1	0	1	0	0	0
B4A16	8	4	1	1	0	0	1	1	1	1	1	0	1	1	1	3	0	0
B1A04	4	4	0	0	0	0	0	0	1	1	0	0	1	0	0	0	0	0
B1A13	4	1	0	0	0	0	0	0	1	0	0	0	1	0	1	0	0	0
B1A19	4	3	0	0	1	1	0	1	1	1	0	0	1	0	1	0	0	0
B1A21	4	2	0	0	0	1	1	0	1	1	0	0	1	0	1	0	0	0
B2A01	4	4	0	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0
B2A06	4	2	0	0	0	0	0	0	1	0	0	0	1	0	0	0	0	0
B2A09	4	1	0	0	0	0	1	0	0	0	0	0	1	0	0	0	0	0
B2A16	4	3	0	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0
B3A03	4	3	0	0	0	1	0	1	1	1	0	0	1	0	1	0	0	0
B3A11	4	2	0	0	0	1	1	1	1	1	0	0	1	0	1	0	0	0
B3A13	4	1	1	1	1	0	0	0	1	1	0	0	1	1	1	1	0	0
B3A23	4	4	0	0	0	0	0	0	1	0	0	0	1	0	0	0	0	0
B4A04	4	4	0	0	0	0	0	0	1	1	0	0	1	0	1	0	0	0
B4A07	4	2	0	0	0	0	0	0	0	0	0	0	1	0	1	0	0	0
B4A11	4	3	0	0	0	0	0	0	1	1	0	0	1	0	0	0	0	0
B4A22	4	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B1A05	2	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B1A07	2	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B1A16	2	2	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B1A17	2	2	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B2A02	2	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B2A08	2	2	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B2A19	2	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B2A20	2	2	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B3A02	2	2	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B3A08	2	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B3A19	2	2	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B3A21	2	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B4A14	2	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B4A15	2	2	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B4A17	2	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0

B4A21	2	2	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B1A08	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B1A09	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B1A15	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B1A18	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B2A04	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B2A05	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B2A13	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B2A15	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B3A01	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B3A06	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B3A12	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B3A17	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B4A03	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B4A09	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B4A12	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
B4A13	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
sum			23	24	21	33	32	34	44	41	30	19	49	19	42	23	2	10
percent_tot			0.28	0.29	0.26	0.4	0.39	0.41	0.54	0.5	0.37	0.23	0.6	0.23	0.51	0.28	0.02	0.12
more			same	yes	no	yes	yes	yes	yes	yes	yes	no	yes	no	yes	same	no	no
percent_12vif				0.91	0.87	0.91	0.96	0.96	1	1	0.96	0.83	1	0.83	1	1	0.09	0.43

343 **Supplementary Table 3.** BioDIV plots included in the different sensitivity overlap scenarios vs. all
344 experimental plots. BioDIV plots with their sown diversity (sown_div), number of functional groups
345 (num_fg) and their selection as realistic plots (1) based on various sensitivity analyses described in
346 Supplementary Information on sensitivity analyses I and Supplementary Table 1. The selection of
347 “realistic” plots is compared to the main analysis (Full12_chull). Note that column D shows the number
348 of years with communities falling inside the intersection of 3-D convex hull volumes (19 = all years).
349 The last four rows show the number of chosen realistic plots (sum), the percentage of total BioDIV
350 plots (159) chosen as realistic (percent_tot), if there are more plots selected as realistic than in the main
351 analysis (yes, same, or no), and the proportion of realistic plots chosen in the main analysis
352 (Full12_chull) also selected in each sensitivity analysis (percent_12vif). Plots are sorted by sown
353 diversity levels.

	plot	sown_div	num_fg	Full12_chull	Full12_hyper	Full12_ellipse	All21_chull	All21_hyper	All21_ellipse	Allbuttax_chull	Allbutphy_chull	Allbutfun_chull	AllbutCWM_chull	A_removes5	C_allreal	D_count	E_NMDS
9		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
27		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
30		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
34		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
35		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	0
46		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
68		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
82		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
89		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
107		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
108		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
136		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	0
156		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	0
160		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	0
164		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
169		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
174		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	0
186		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
202		16	4	1	1	1	1	1	1	1	1	1	1	1	1	19	1
220		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
227		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
235		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
239		16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
242		16	5	1	1	1	1	1	1	1	1	1	1	1	1	18	1

253	16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
257	16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
273	16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
299	16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
301	16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
318	16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
328	16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
329	16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
331	16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	0
336	16	5	1	1	1	1	1	1	1	1	1	1	1	1	19	0
339	16	5	1	1	1	1	1	1	1	1	1	1	1	1	18	1
12	8	4	1	1	1	1	1	1	1	1	1	1	1	1	19	1
15	8	4	1	1	1	1	1	1	1	1	1	1	1	1	19	1
22	8	4	1	1	1	1	1	1	1	1	1	1	1	1	18	0
50	8	4	1	1	1	1	1	1	1	1	1	1	1	1	19	1
57	8	4	1	1	1	1	1	1	1	1	1	1	1	1	19	1
67	8	5	1	1	1	1	0	1	1	1	1	1	1	1	18	0
74	8	5	1	1	1	1	1	1	1	1	1	1	1	1	15	0
81	8	4	1	1	1	1	1	1	1	1	1	1	1	1	19	1
98	8	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
104	8	5	1	1	1	1	1	1	1	1	1	1	1	1	19	0
111	8	4	1	1	1	1	1	1	1	1	1	1	1	1	19	1
115	8	4	1	1	1	1	1	1	1	1	1	1	1	1	11	1
118	8	5	1	1	1	1	1	1	1	1	1	1	1	1	17	0
130	8	5	1	1	1	1	1	1	1	1	1	1	1	1	19	0
146	8	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
170	8	4	1	1	1	1	1	1	1	1	1	1	1	1	18	0
177	8	5	1	1	1	1	1	1	1	1	1	1	1	1	19	0
178	8	3	1	1	1	1	1	1	1	1	1	1	1	1	19	0
206	8	4	1	1	1	1	1	1	1	1	1	1	1	1	19	1
208	8	4	1	1	1	1	1	1	1	1	1	1	1	1	19	1
210	8	4	1	1	1	1	1	1	1	1	1	1	1	1	19	1
213	8	5	1	1	1	1	1	1	1	1	1	1	1	1	19	1
232	8	5	1	1	1	1	1	1	1	1	1	1	1	1	18	0
266	8	5	1	1	1	1	1	1	1	1	1	1	1	1	19	0
283	8	3	1	1	1	1	1	1	1	1	1	1	1	1	19	0
292	8	3	1	1	1	1	1	1	1	1	1	1	1	1	19	1
293	8	4	1	1	1	1	1	1	1	1	1	1	1	1	19	0
303	8	3	1	1	1	1	1	1	1	1	1	1	1	1	18	1

307	8	4	1	1	1	1	1	1	1	1	1	1	1	1	18	0
313	8	4	1	1	1	1	1	1	1	1	1	1	1	1	18	0
3	4	2	1	1	1	1	1	1	1	1	1	1	1	1	18	0
24	4	3	1	1	1	1	1	1	1	1	1	1	1	1	18	0
26	4	3	0	0	1	0	0	1	0	0	0	1	0	0	0	0
28	4	2	1	1	1	1	1	1	0	1	1	0	0	1	19	0
33	4	4	1	0	1	1	0	1	1	1	1	1	1	1	3	0
44	4	3	1	0	1	1	0	0	1	1	1	1	0	1	19	0
45	4	4	1	0	1	1	0	1	1	1	1	1	1	1	1	0
53	4	4	1	0	1	1	1	1	1	1	1	1	1	1	19	1
58	4	3	1	0	1	1	1	1	1	1	1	1	1	1	8	1
62	4	3	1	1	1	1	1	1	1	1	1	1	1	1	11	0
70	4	3	1	1	1	1	1	1	1	1	1	1	1	1	19	0
93	4	2	1	0	0	1	0	0	1	1	1	1	0	1	2	0
110	4	3	1	0	1	1	0	1	1	1	1	1	1	1	18	0
133	4	3	1	1	1	1	1	1	1	1	1	1	1	1	19	0
138	4	2	1	1	1	1	1	1	1	1	1	1	1	1	19	0
139	4	2	1	1	1	1	1	1	1	1	1	1	1	1	18	0
149	4	4	0	0	0	1	0	1	1	1	1	1	1	0	0	0
176	4	3	1	1	1	1	1	1	1	1	1	1	1	1	19	1
190	4	3	1	1	1	1	1	1	1	1	1	1	1	1	19	0
199	4	3	1	1	1	1	1	1	1	1	1	1	1	1	14	0
201	4	3	0	0	0	1	0	1	1	1	1	1	1	0	0	0
223	4	3	1	1	1	1	1	1	1	1	1	1	1	1	15	1
225	4	4	1	1	1	1	1	1	1	1	1	1	1	1	3	0
229	4	3	1	1	1	1	1	1	1	1	1	1	1	1	18	1
233	4	4	1	1	1	1	1	1	1	1	1	1	1	1	19	1
286	4	3	1	1	1	1	1	1	1	1	1	1	1	1	19	1
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325	4	4	1	1	1	1	1	1	1	1	1	1	1	1	15	0
6	2	1	1	1	1	1	1	1	1	1	1	1	1	1	17	0
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32	2	2	0	0	0	0	0	0	0	0	0	0	0	0	0	0
48	2	2	1	1	0	1	1	1	1	1	1	1	1	1	9	0
56	2	2	1	1	1	1	1	1	1	1	1	1	1	1	16	0
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127	2	2	1	1	1	1	1	1	0	1	1	0	0	1	12	0
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165	2	2	0	0	1	0	0	1	0	0	0	1	1	0	0	1
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236	2	2	1	0	1	1	0	0	1	1	1	1	1	1	8	0
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311	2	2	1	0	1	1	0	1	1	1	1	1	1	1	18	0
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16	1	1	1	0	0	0	0	0	0	0	0	0	0	1	18	0
20	1	1	1	1	1	0	0	0	1	0	0	0	0	1	18	0
29	1	1	1	1	1	1	1	1	0	1	1	0	0	1	19	0
31	1	1	0	0	1	0	0	1	0	0	0	0	0	0	0	0
69	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0
83	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0
87	1	1	0	0	0	0	0	1	0	0	0	0	0	0	0	0
92	1	1	0	0	0	0	0	1	0	0	0	0	0	0	0	0
94	1	1	1	1	1	1	1	1	0	1	1	0	0	1	19	0
109	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0
129	1	1	0	1	1	0	0	1	0	0	0	0	0	0	0	0

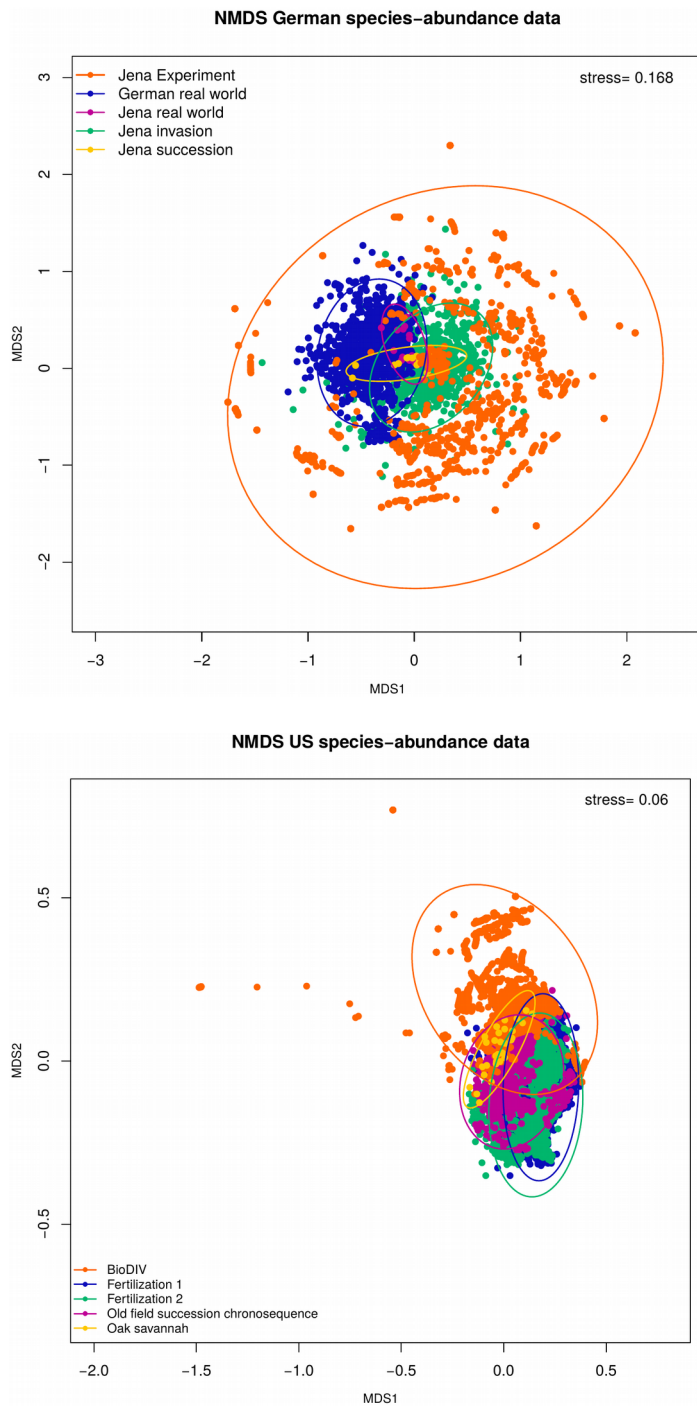
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142	1	1	0	0	0	0	0	1	0	0	0	1	1	0	0	0
153	1	1	1	1	1	1	1	1	1	1	1	0	0	1	10	0
163	1	1	0	0	0	0	0	1	0	0	0	0	0	0	0	0
167	1	1	0	1	1	0	0	1	0	0	0	0	0	0	0	0
230	1	1	1	0	1	0	0	0	1	0	0	0	0	1	18	0
237	1	1	1	1	1	1	1	1	1	1	1	1	1	1	19	1
256	1	1	1	1	1	1	1	1	1	1	1	0	0	1	11	0
265	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0
267	1	1	0	1	1	0	0	1	0	0	0	0	0	0	0	0
268	1	1	0	0	0	0	0	1	0	0	0	1	1	0	0	0
280	1	1	0	0	1	0	0	1	0	0	0	0	0	0	0	0
282	1	1	0	0	0	0	0	1	0	0	0	0	0	0	0	0
290	1	1	1	1	1	1	1	1	1	1	1	1	0	1	19	0
308	1	1	1	0	0	0	0	0	0	0	0	0	0	1	19	0
333	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0
338	1	1	1	1	1	1	1	1	1	1	1	1	0	1	19	0
sum			122	109	133	121	104	136	122	125	124	126	112	122	122	53
percent_tot			0.77	0.69	0.84	0.76	0.65	0.86	0.77	0.79	0.78	0.79	0.7	0.77	0.77	0.33
more			same	no	yes	no	no	yes	yes	yes	yes	yes	no	yes	same	no
percent_12vif				0.87	0.96	0.95	0.85	0.91	0.94	0.96	0.96	0.9	0.83	1	1	0.43

354 **Supplementary Table 4.** T-test results for differences between German experimental and real-world
355 plots. Welsh t-tests with unequal variances. Full set of 21 community properties averaged across all
356 years per plot for Jena Experiment (82 plots) and combined real-world data (German real world: 150
357 plots, Jena real world: 14 plots). T-statistic, degrees of freedom (df), experimental (Exp) data mean and
358 real world (RW) data mean are rounded to two, p-values to 5 decimal places. The last column gives the
359 proportion of Jena plots falling within the community property range of the real-world plots.

model	t_statistic	df	mean_Exp	mean_RW	p_value	in_rw_range
FRic	-19.04	192.57	2.74	12.84	0	0.24
FEve	-5.57	83.98	0.34	0.52	0	0.5
FDiv	-5.34	85.44	0.46	0.68	0	0.59
FDis	-8.5	88.07	1.18	1.95	0	0.56
RaoQ	-9.51	105.73	2.39	4.61	0	0.55
leaf_area	1	106.43	1820.3	1648.03	0.31776	0.82
SLA	-4.54	140.03	24.26	26.44	0.00001	0.94
leaf_drymass	2.12	93.49	61.03	48.28	0.03662	0.82
LDMC	-5.27	98.28	0.22	0.25	0	0.61
leaf_N	1.31	96.71	25.66	24.81	0.19315	0.74
leaf_P	-1.44	109.38	2.19	2.26	0.15242	0.93
height	-2.36	115.5	0.45	0.5	0.01988	0.76
seedmass	3.95	84.7	2.76	1.52	0.00016	0.72
S	-18.8	195.11	6.45	28.05	0	0.11
H	-14.22	99.02	0.97	2.3	0	0.43
D1	-10.9	84.47	0.46	0.83	0	0.51
D2	-10.17	133.06	3.23	7.42	0	0.51
SEve	16.35	86.69	0.65	0.27	0	0.05
PD	-20.34	163.66	720.24	1704.8	0	0.12
MPD	-8.5	101.21	105.59	188.92	0	0.66
MNTD	6.62	82.65	109.29	31.2	0	0.29

360 **Supplementary Table 5.** T-test results for differences between US experimental and real-world plots.
 361 Welsh t-tests with unequal variances. Full set of 21 community properties averaged across all years per
 362 plot for BioDIV (159 plots) and combined real-world data (Fertilization 1 & 2; 207 and 162 plots,
 363 respectively). T-statistic, degrees of freedom (df), experimental (Exp) data mean and real world (RW)
 364 data mean are rounded to two, p-values to 5 decimal places. The last column gives the proportion of
 365 BioDIV plots falling within the community property range of the real-world plots.

model	t_statistic	df	mean_Exp	mean_RW	p_value	in_rw_range
FRic	-7.59	243.07	1.4	2.62	0	0.51
FEve	-2.73	176.06	0.29	0.35	0.00693	0.28
FDiv	-8.42	182.59	0.37	0.63	0	0.59
FDis	0.79	187.27	1.02	0.97	0.43191	0.59
RaoQ	3.46	182.45	2.22	1.61	0.00068	0.6
leaf_area	8.82	176.29	1429.56	724.02	0	0.77
SLA	-8.01	204.34	14.77	16.5	0	0.4
leaf_drymass	10	193.36	74.23	45.67	0	0.73
LDMC	-2.57	169.1	0.3	0.32	0.01105	0.22
leaf_N	-4.9	237.85	16.97	18.77	0	0.5
leaf_P	-6.68	248.46	1.55	1.71	0	0.94
height	3.37	193.75	0.77	0.7	0.0009	0.56
seedmass	5.21	169.51	3.88	2.18	0	0.82
S	-12.12	291.42	4.03	7.85	0	0.76
H	-4.21	222.01	0.72	0.96	0.00004	0.73
D1	-4.18	215.21	0.36	0.47	0.00004	0.7
D2	-1.83	226.06	2.23	2.46	0.06831	0.71
SEve	18.02	188.98	0.71	0.38	0	0.8
PD	-13.45	338.91	605.02	861.86	0	0.67
MPD	-0.3	239.47	92.55	94.68	0.76228	0.68
MNTD	4.71	169.23	97.1	50.95	0.00001	0.6



366 **Supplementary Figure 2.** NMDS biplots of species-abundance data for German and US dataset.
 367 First two axes of 3-dimensional NMDS for German (upper panel) and US (lower panel) species-
 368 abundance data as calculated with function “metaMDS” and plotted with function “biplot” in R
 369 package *vegan*¹². Note that, for the US comparison, communities with 100 % relative abundance of
 370 *Elymus smithii* are not shown in this biplot since they show extreme values in the first axis.

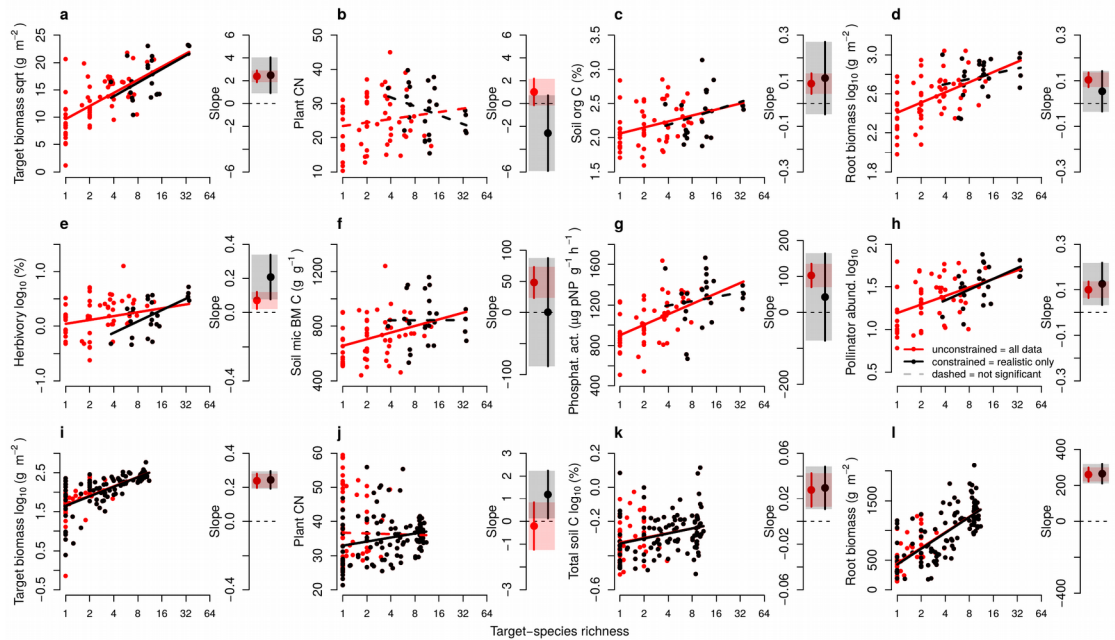
371 **Supplementary Table 6.** T-test results for differences between realistic and unrealistic plots for the
372 Jena Experiment. Welsh t-tests with unequal variances. Realistic plots were calculated based on the 12
373 vif-selected community properties and the convex hull volume method. All properties were averaged
374 across all available years per plot (23 realistic and 59 unrealistic plots). T-statistic, degrees of freedom
375 (df), means of realistic (real) and unrealistic communities (unreal) are rounded to two, p-values to four
376 decimal places.

model	t_statistic	df	mean_real	mean_unreal	p_value
FRic	8.25	24.88	7.32	0.96	0
FEve	6.46	74.97	0.54	0.27	0
FDiv	7.82	66.62	0.75	0.34	0
FDis	9.61	77.96	1.98	0.87	0
RaoQ	9.22	55.01	4.49	1.58	0
leaf_area	0.13	75.23	1842.7	1811.57	0.8971
SLA	0.63	49.5	24.64	24.11	0.5334
leaf_drymass	-0.55	78.6	57.26	62.5	0.5845
LDMC	1.75	42.61	0.23	0.21	0.0865
leaf_N	0.3	78.09	25.88	25.58	0.7679
leaf_P	2.24	77.01	2.31	2.14	0.0281
height	1.87	61.58	0.5	0.42	0.0663
seedmass	-0.35	79.48	2.64	2.81	0.7303
S	5.66	22.7	15.18	3.05	0
H	9.17	34.51	1.92	0.6	0
D1	9.9	72.8	0.78	0.34	0
D2	5.06	22.72	6.65	1.9	0
SEve	-10.33	78.5	0.45	0.74	0
PD	5.7	25.31	1093.58	574.71	0
MPD	5.47	38.3	176.42	77.97	0
MNTD	-3.56	77.95	64.82	126.63	0.0006
sown diversity	4.77	22.34	21.74	3.46	0.0001
no funct groups	3.54	36.95	2.83	1.85	0.0011

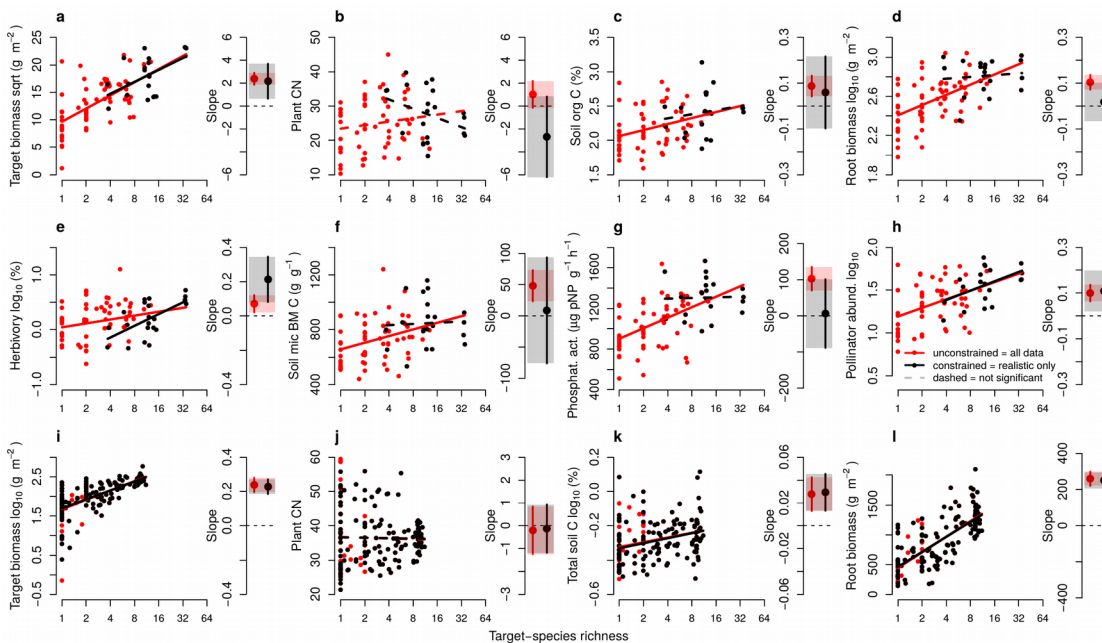
377 **Supplementary Table 7.** T-test results for differences between realistic and unrealistic plots for
378 BioDIV. Welsh t-tests with unequal variances. Realistic plots were calculated based on the full set of
379 community properties and the convex hull volume method. All properties were averaged across all
380 available years per plot (122 realistic and 37 unrealistic plots). T-statistic, degrees of freedom (df),
381 means of realistic (real) and unrealistic communities (unreal) are rounded to two, p-values to four
382 decimal places.

model	t_statistic	df	mean_real	mean_unreal	p_value
FRic	10.7	121	1.82	0	0
FEve	14.65	121	0.38	0	0
FDiv	14.71	121	0.48	0	0
FDis	9.44	92.96	1.26	0.22	0
RaoQ	7.92	95.01	2.75	0.49	0
leaf_area	-0.11	44.52	1423.77	1448.65	0.9168
SLA	1.04	46.22	14.91	14.3	0.3043
leaf_drymass	-1.27	46.88	71.95	81.76	0.2119
LDMC	-0.66	44.55	0.3	0.31	0.5149
leaf_N	1.2	42.28	17.27	15.99	0.2366
leaf_P	1.42	46.02	1.57	1.48	0.1633
height	-1.81	40.78	0.75	0.87	0.0773
seedmass	0.15	41.7	3.92	3.76	0.8815
S	11.53	130.41	4.87	1.25	0
H	12.76	155.85	0.91	0.09	0
D1	12.7	145.19	0.45	0.06	0
D2	11.23	133.74	2.58	1.09	0
SEve	-10.43	102.79	0.64	0.92	0
PD	12.13	154.37	660.65	421.61	0
MPD	10.42	111.91	115.44	17.08	0
MNTD	1.95	55.87	107.8	61.82	0.056
sown diversity	11.51	137.13	7.81	1.65	0
no funct groups	10.95	122.28	3.49	1.54	0

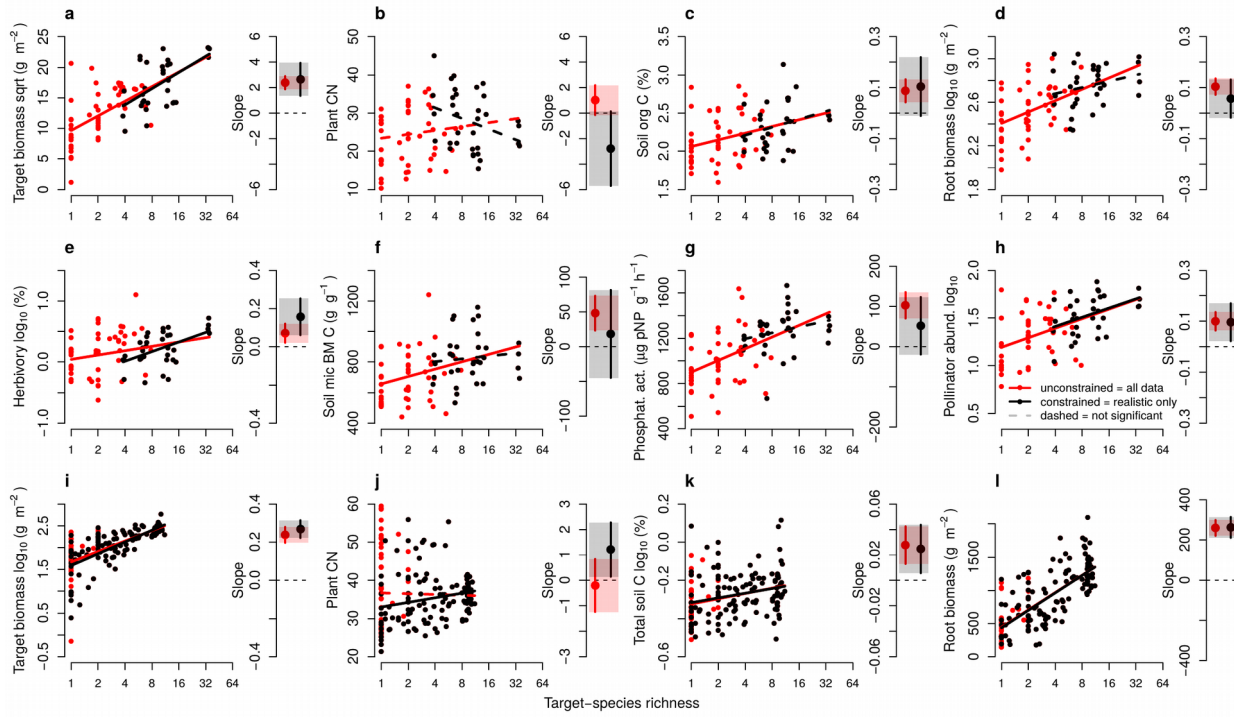
383 **Supplementary Figure 3.** Alternative versions of Fig. 2 based on alternative intersection scenarios
 384 (community- property subsets and overlap calculation methodology). Panels a-h Jena Experiment,
 385 panels i-l BioDIV (see main text Fig. 2). 6 different versions: 3 methods (chull, hyper, ellipse) and 2
 386 community property subsets (Full 12 vif selected and All 21 community properties; see Supplementary
 387 Table 1 for details on included community properties) plus one USA-only version based on overlap
 388 from species-abundance based NMDS with the chull method. Note that convex hull method with 12
 389 vif-selected properties is main text Fig. 2.
 390 Full 12 – hyper:



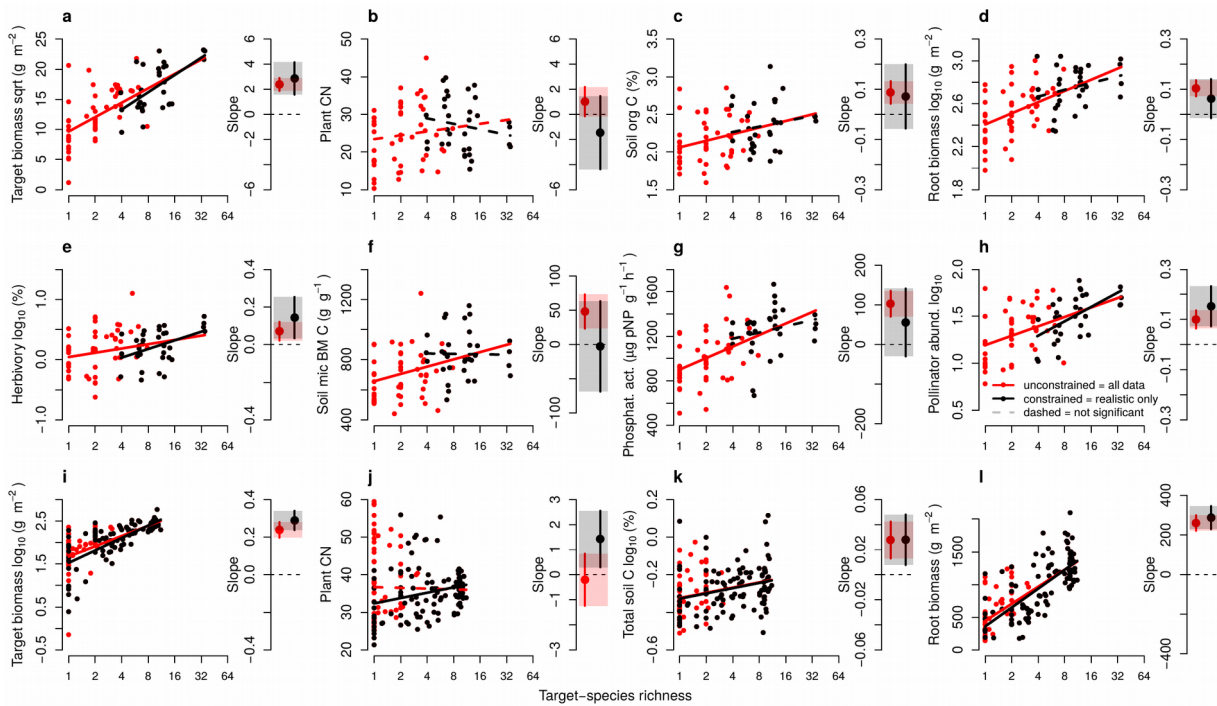
391 Full 12 – ellipse:



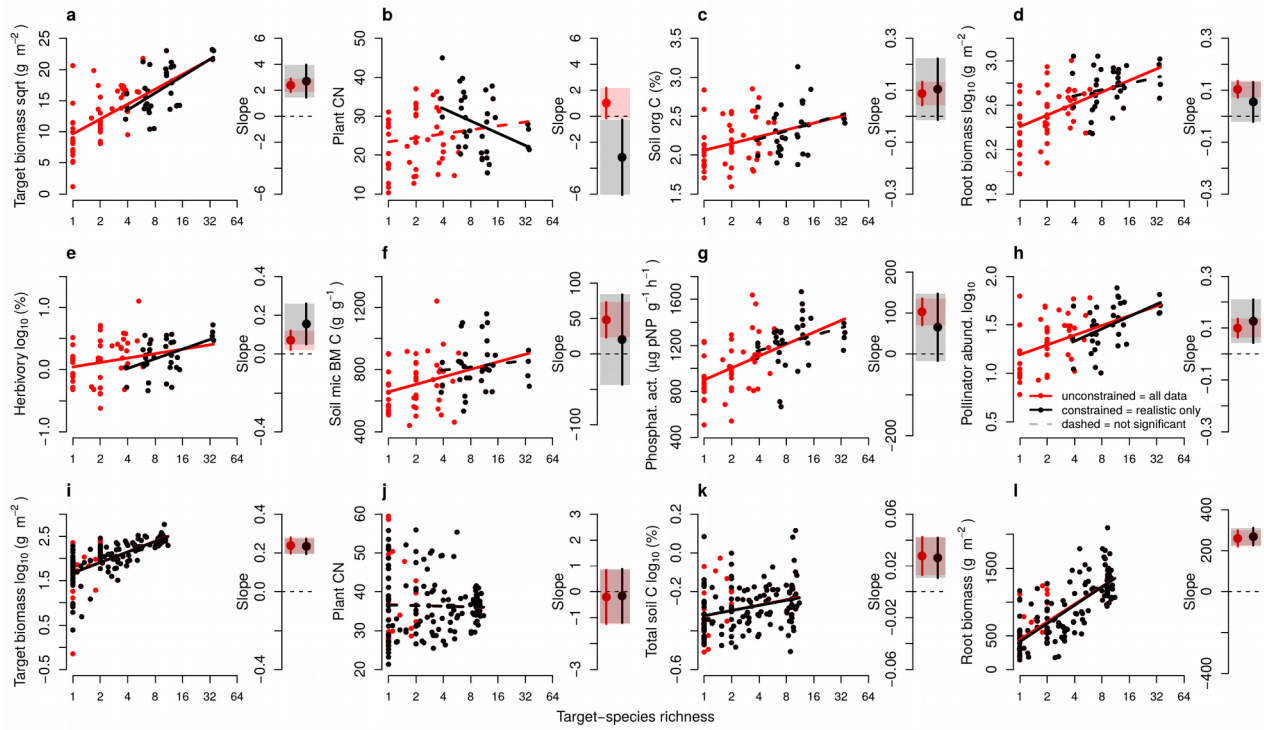
392 All 21 – chull:



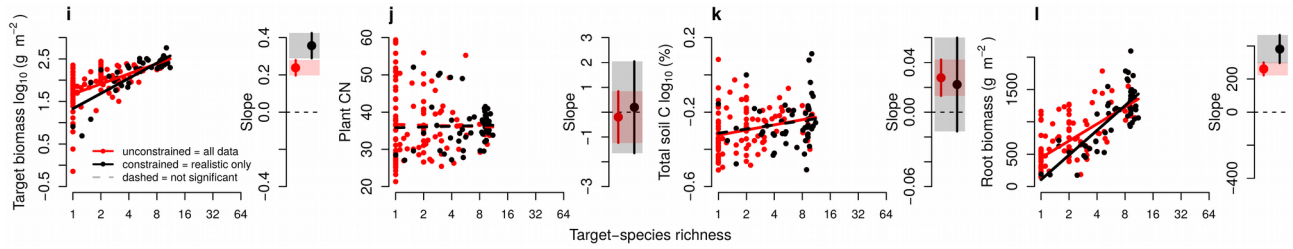
393 All 21 – hyper:



394 All 21 – ellipse:



395 USA – NMDS – chull:



396 **Supplementary Table 8.** Constraining-related change in functioning at maximum species richness.
 397 For each of the 12 BEF relationships from the Jena Experiment (J) and BioDIV presented in Fig. 2, the
 398 table shows the constraining-related percentage change in the model-predicted function variable at
 399 maximum species richness (the proportional difference in the un-transformed function value at the
 400 right-hand tip of the black and red lines in Fig. 2). The average absolute percentage function change is
 401 10.3% (SE: 4%).

function	% change in predicted functioning
J_biomass	-3.9
J_plantCN	-18.75
J_soilorgC	-0.46
J_rootbiomass	-20.24
J_herbivory	46.24
J_micBMC	-7.8
J_phosphatase	-8.17
J_pollinators	9.74
BioDIV_biomass	3.78
BioDIV_plantCN	3.59
BioDIV_soilC	0.39
BioDIV_rootbiomass	-0.05

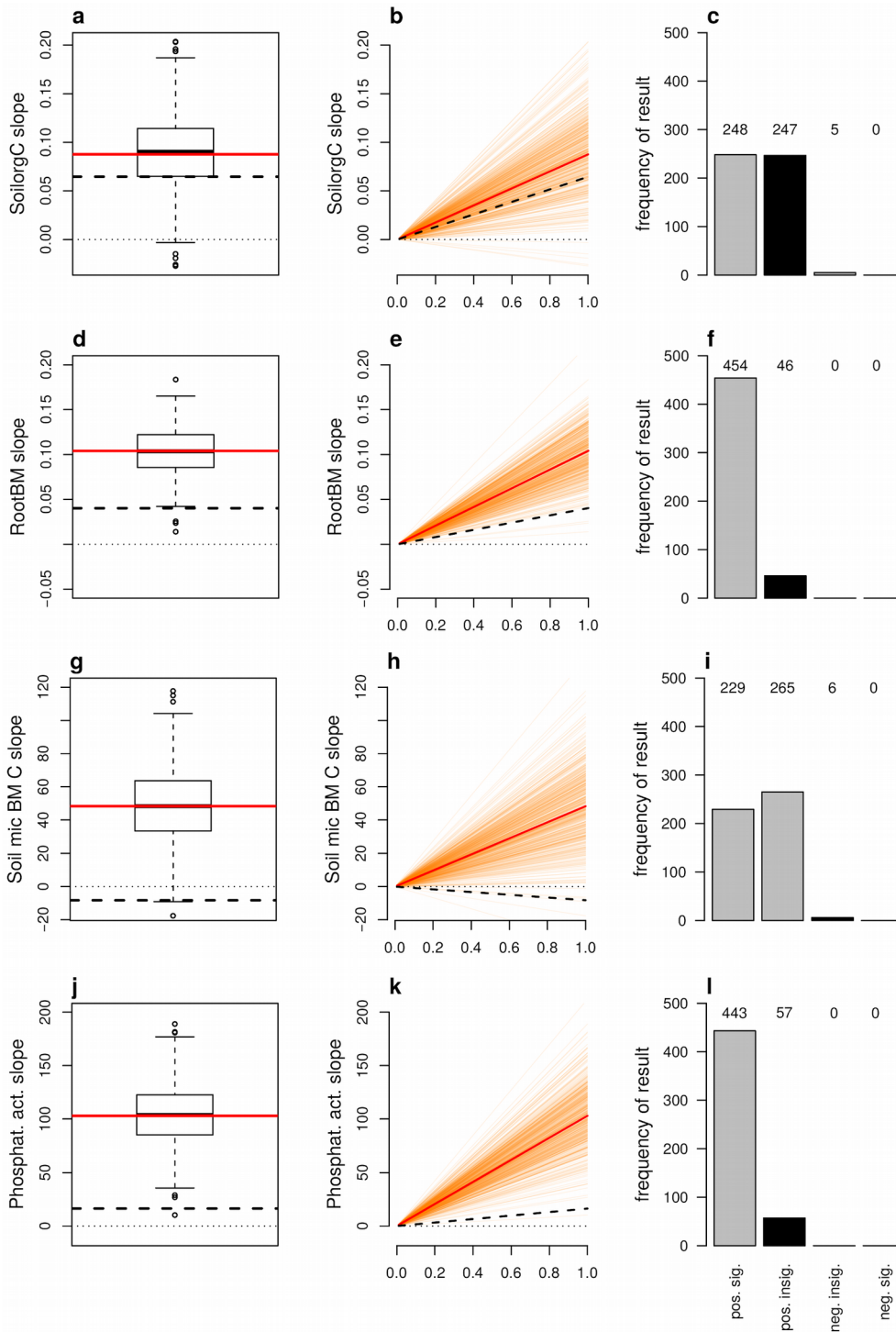
402 **Supplementary Information on sensitivity analyses II**

403 We performed two very similar sensitivity analyses testing if changes in BEF relationships from being
404 significant (all communities) to non-significant (realistic plots only) were likely caused by the related
405 reduction in sample size or the reduced species richness gradient of the constrained Jena plots or if a
406 randomly-selected reduced number of plots was still likely to result in a significant relationship.
407 Therefore, for each of the four BEF relationships found to switch significance (Jena soil organic C (a-
408 c), root biomass (d-f), soil microbial biomass C (g-i) and phosphatase activity (j-l)), we repeatedly (500
409 times), randomly selected 23 Jena plots and re-ran the model testing for the BEF relationship and
410 saving the slope estimates and p-values (Supplementary Fig. S4a, fully random choice of plots).
411 Supplementary Fig. 4 shows the distribution of these 500 random-selection slopes (boxplots in first
412 column and orange lines in middle column) in comparison to the unconstrained (all plots, red lines) and
413 constrained (PCA-selection based realistic plots only, black dashed lines) slopes from Fig. 2. Dotted
414 black lines indicate zero slopes. The right column shows the frequency of positive significant, positive
415 insignificant, negative insignificant and negative significant relationships obtained by the 500 random
416 subsets of 23 plots with the black bar highlighting the PCA-based realistic result from Fig. 2.

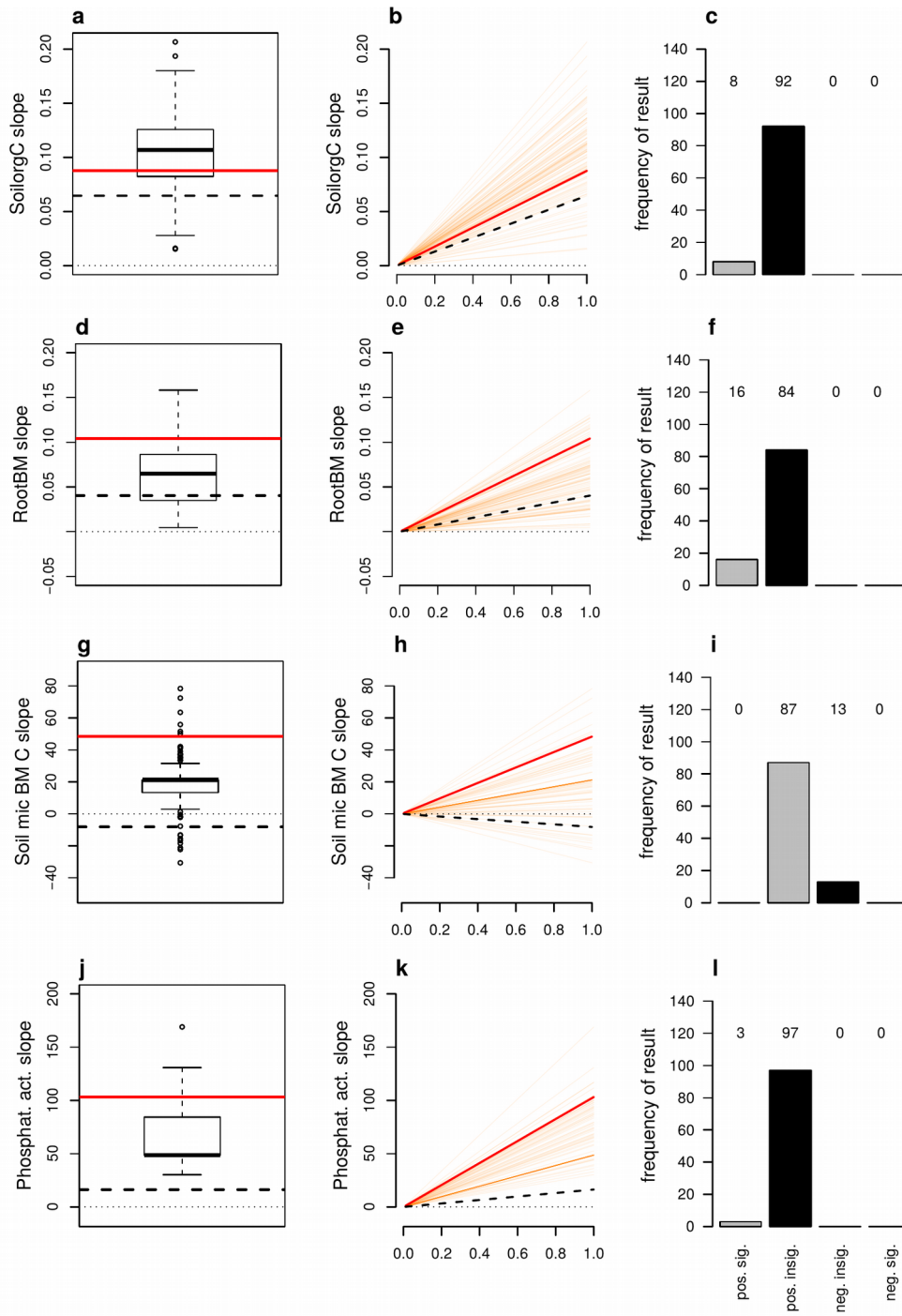
417 In a second version of this analysis, we restricted the random choice of Jena plots to only draw
418 from the 42 plots within the species richness range of the realistic plots in the main analysis (S equal to
419 or larger than 3.7). Here, we repeated the random draw only 100 times. Otherwise, the analysis and
420 figure are identical.

421 Supplementary Fig. 4 shows that black dashed lines and the results of the PCA-based realistic
422 subset divert relatively strongly from the 500 random-selection results. Specifically, the PCA-based
423 realistic subset resulted in strikingly shallower slopes than the random choices and non-significantly
424 positive or even negative relationships while a big part of the random subsets resulted in significant
425 positive or at least non-significantly positive relationships. As such, our PCA-based selection of
426 realistic plots is highly non-random in comparison to the random-selection of plots, thus indicating that
427 our methodology is successful in finding a subset of plots based on prior knowledge (realistic plots
428 based on the multidimensional, multivariate comparison of communities) and does not simply create a
429 random subset of plots. Furthermore, these results show that, for these four Jena soil processes,
430 experiment-derived BEF relationships might not be as important or strong in real-world systems, at
431 least as long as plant communities in those real-world systems deviate from those in experiments, e.g.
432 in their species richness gradients. Future developments of real-world plant communities due to global
433 change drivers and increasing anthropogenic pressure might change this conclusion by rendering less
434 diverse communities realistic, thus aligning the species richness gradients of biodiversity experiments
435 and related real-world systems and increasing the slope of the BEF relationships.

436 Interestingly, Supplementary Fig. 5 shows that, when restricting the species richness gradient of
437 randomly drawn realistic plots to being comparable to the realistic plots chosen in the main analysis,
438 the picture changes. The vast majority of random slopes is now not significant anymore, as can be seen
439 from the barplots in the right-hand column. This shows that the change from significant to non-
440 significant BEF relationships is not caused by the reduced sample size but primarily driven by the
441 truncated species-richness gradient in more real-world comparable biodiversity experiment data
442 subsets.



443 **Supplementary Figure 4.** Random selection sensitivity analysis for Fig. 2 relationships turning
 444 insignificant. 500 random draws of Jena plots and the respective BEF slopes for four selected
 445 functions.



446 **Supplementary Figure 5.** Random selection sensitivity analysis for Fig. 2 relationships turning
 447 insignificant. 100 restricted random draws of Jena plots and the respective BEF slopes for four selected
 448 functions, random draws confined to the species richness gradient also covered by the realistic Jena
 449 plots in the main analysis (equal to or larger than 3.7 species).

450 **Supplementary Table 9.** Differences between range in function for unconstrained and constrained data
 451 underlying the BEF relationships in Fig. 2. Values are presented for unconstrained (uncon) and
 452 constrained (con) datasets of Jena (J) and BioDIV BEF relationships. Constraining was done using the
 453 12 vif-selected community properties and the convex hull method. Ranges were calculated based
 454 maximum and minimum function performance in unconstrained and constrained datasets. Range
 455 changes were calculated as the proportion of unconstrained functioning still covered by constrained
 456 functioning. Changes are caused by the removal of unrealistic plots which alters the distribution of
 457 function values for a given species richness level, but also by the reduction of the species-richness
 458 gradient that is caused by the removal of plots. The across-year species-richness gradient in Jena
 459 changed from 1-35.2 species (unconstrained) to 3.7-35.2 species (constrained). The BioDIV species
 460 richness gradient was 1-11.1 species and did not change from unconstrained to constrained datasets.

model_name	uncon_range	con_range	range_change
J_biomass	22	9.7	0.44
J_plantCN	34.59	24.33	0.7
J_soilorgC	1.54	1.26	0.82
J_rootbiomass	1.06	0.66	0.63
J_herbivory	1.72	1.05	0.61
J_micBMC	800.54	624.56	0.78
J_phosphatase	1159.23	956.63	0.83
J_pollinators	1.1	0.82	0.74
BioDIV_biomass	2.91	2.38	0.82
BioDIV_plantCN	38.04	34.59	0.91
BioDIV_soilC	0.62	0.62	1
BioDIV_rootbiomass	1952.87	1952.87	1
Jena_avg			0.69
BioDIV_avg			0.93
Overall_avg			0.77

461 **Supplementary Table 10.** Drivers of ecosystem functioning in all vs. realistic communities of
462 biodiversity experiments. In order to assess how the importance of function drivers changes, we
463 analysed relative importance of community properties in explaining functions for the complete set of
464 biodiversity experiment plots and realistic-only plots. As functions, we chose plant aboveground
465 biomass and soil organic carbon as they cover different types of functions (unlike e.g. plant
466 aboveground and root biomass) and are available for both experiments. For each of these functions
467 (response), in each of the two experiments (experiment), we compared the relative importance of 8
468 selected community properties (predictor), two taxonomic diversity metrics, two functional diversity
469 metrics, two phylogenetic diversity metrics and two community weighted mean functional traits (S,
470 SEve, FRic, FEve, PD, MNTD, seed mass, SLA). These pairs of community properties (from each of
471 the community property types (pred_type): tax, phyl, funct, CWM) were selected based on relatively
472 low correlation within the type of properties.
473 For each function, we set up a full model with the function variable as the response and the 8 selected
474 community properties as predictors (simple linear model, no interactions, just additive effects). All
475 variables were standardized to zero mean and unit variance before model runs to compare relative
476 importance. Subsequently, we used the function “dredge” in R package “MuMIn” to compute models
477 with all possible variable combinations and rank them by AICc. We then used the function
478 “importance” from the same R package to extract the summed akaik weight of all models that each
479 variable was present in as a predictor (weight_all). This procedure was repeated for the constrained
480 dataset to obtain predictor importance in the constrained (realistic plots only) datasets (weight_con;
481 based on the main-text analysis using convex hull volume and the vif-selected variable set for the
482 PCA’s). Finally, for each of the eight predictors of both functions in both experiments, we calculated
483 the absolute difference in variable importance between the analyses with all plots and realistic plots
484 only (abs_change).

experiment	response	predictor	pred_type	weight_all	weight_con	abs_change
Jena	biomass	seedmass	CWM_trait	1	0.99	-0.01
Jena	biomass	SEve	taxonomic	0.9	0.91	0.01
Jena	biomass	S	taxonomic	0.9	0.99	0.09
Jena	biomass	FEve	functional	0.46	0.1	-0.36
Jena	biomass	PD	phylogenetic	0.32	0.79	0.47
Jena	biomass	MNTD	phylogenetic	0.28	0.17	-0.11
Jena	biomass	FRic	functional	0.25	0.3	0.05
Jena	biomass	SLA	CWM_trait	0.25	0.59	0.34
Jena	soilorgC	SEve	taxonomic	0.6	0.2	-0.4
Jena	soilorgC	FRic	functional	0.56	0.24	-0.32
Jena	soilorgC	PD	phylogenetic	0.33	0.2	-0.13
Jena	soilorgC	S	taxonomic	0.32	0.19	-0.13
Jena	soilorgC	FEve	functional	0.29	0.19	-0.1
Jena	soilorgC	seedmass	CWM_trait	0.29	0.46	0.17
Jena	soilorgC	MNTD	phylogenetic	0.27	0.26	-0.01

Jena	soilorgC	SLA	CWM_trait	0.24	0.24	0
BioDIV	biomass	FRic	functional	0.99	0.99	0
BioDIV	biomass	MNTD	phylogenetic	0.88	0.75	-0.13
BioDIV	biomass	PD	phylogenetic	0.58	0.62	0.04
BioDIV	biomass	S	taxonomic	0.42	0.39	-0.03
BioDIV	biomass	seedmass	CWM_trait	0.32	0.28	-0.04
BioDIV	biomass	FEve	functional	0.31	0.28	-0.03
BioDIV	biomass	SEve	taxonomic	0.27	0.28	0.01
BioDIV	biomass	SLA	CWM_trait	0.26	0.3	0.04
BioDIV	soilC	SEve	taxonomic	0.79	0.88	0.09
BioDIV	soilC	seedmass	CWM_trait	0.73	0.25	-0.48
BioDIV	soilC	PD	phylogenetic	0.52	0.54	0.02
BioDIV	soilC	FEve	functional	0.43	0.37	-0.06
BioDIV	soilC	S	taxonomic	0.31	0.32	0.01
BioDIV	soilC	FRic	functional	0.29	0.29	0
BioDIV	soilC	SLA	CWM_trait	0.27	0.27	0
BioDIV	soilC	MNTD	phylogenetic	0.27	0.27	0

485 **Supplementary Table 11.** Correlation coefficients for CWM's versus functional, phylogenetic metrics
 486 and evenness, German dataset. Pearson correlation coefficients for Jena Experiment (upper part) and
 487 combined German real world community properties (lower part). Bold values are mean absolute
 488 correlation coefficients for the columns, the overall mean is the absolute mean across all column
 489 averages.

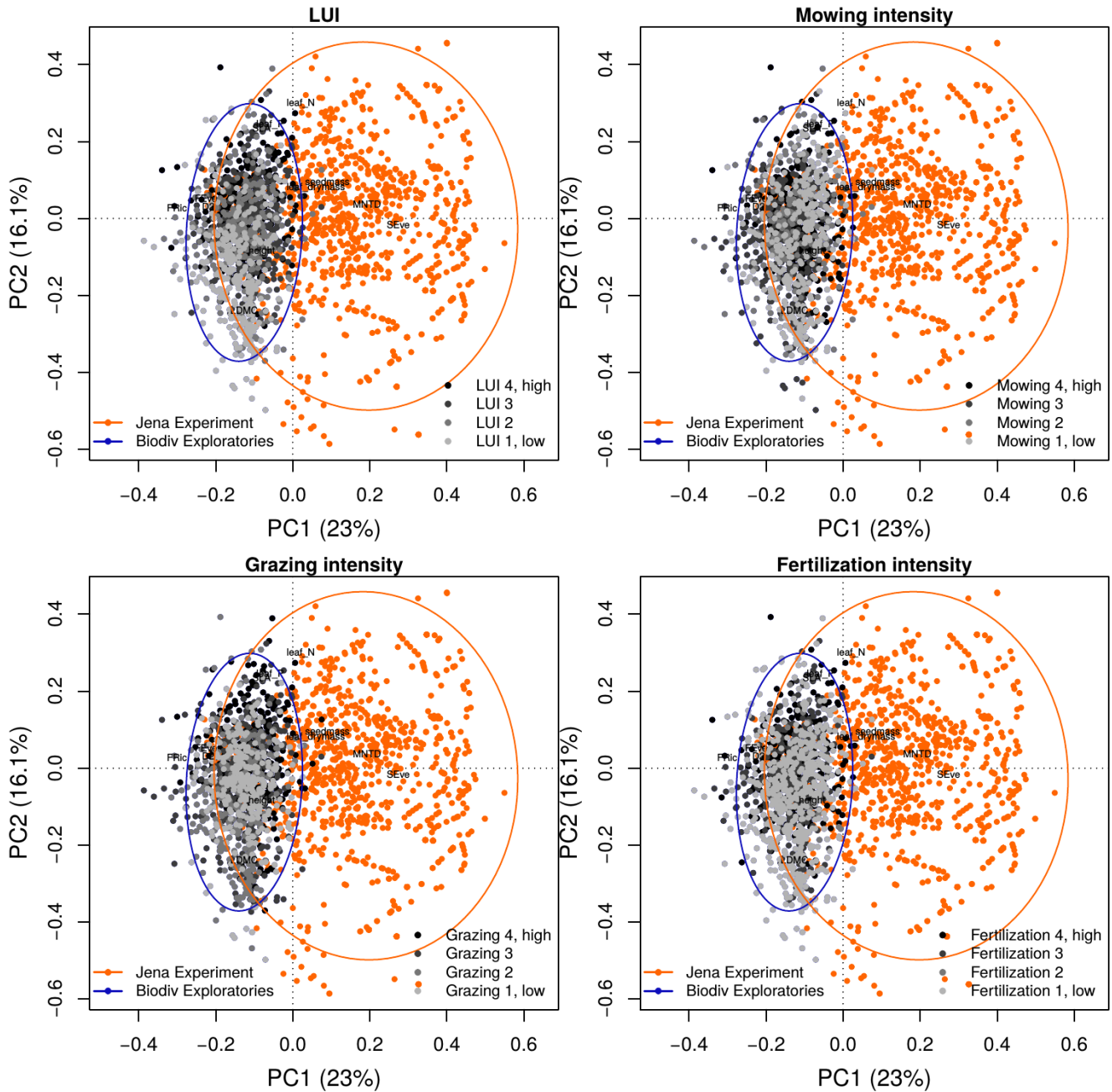
Jena	FEve	FDiv	FDis	RaoQ	SEve	MPD	MNTD
leaf_area	-0.02	-0.03	0.02	0.07	-0.01	0	0.12
SLA	0	-0.09	0.05	0.06	0.09	0.09	-0.02
leaf_drymass	0.05	0.11	0.1	0.08	-0.12	0.08	0.23
LDMC	0	0.03	0.02	0.06	-0.07	-0.08	-0.15
leaf_N	0.09	0.07	0.14	0.13	-0.09	0.05	-0.03
leaf_P	0.12	0.16	0.22	0.21	-0.16	0.18	0.02
height	0.04	0.05	0.04	0.1	-0.08	-0.08	-0.12
seedmass	-0.01	0.02	-0.02	0.03	0.01	-0.1	-0.13
avg_abs	0.04	0.07	0.08	0.09	0.08	0.08	0.1
overall mean	0.08						

German RW	FEve	FDiv	FDis	RaoQ	SEve	MPD	MNTD
leaf_area	-0.08	-0.09	0.34	0.47	0.19	0.04	0.12
SLA	-0.21	-0.21	0.07	0.07	0.14	-0.13	0.03
leaf_drymass	-0.03	0.12	0.52	0.57	0.25	0.36	0.18
LDMC	0.1	-0.09	-0.23	-0.2	-0.25	-0.5	-0.06
leaf_N	-0.15	-0.05	0.29	0.26	0.12	0.25	0.05
leaf_P	-0.19	-0.24	0.06	0.11	0.06	-0.16	0.03
height	-0.03	-0.22	-0.04	0.05	-0.07	-0.5	-0.07
seedmass	0.14	0.14	0.35	0.42	0.04	0.16	0.08
avg_abs	0.12	0.15	0.24	0.27	0.14	0.26	0.08
overall mean	0.18						

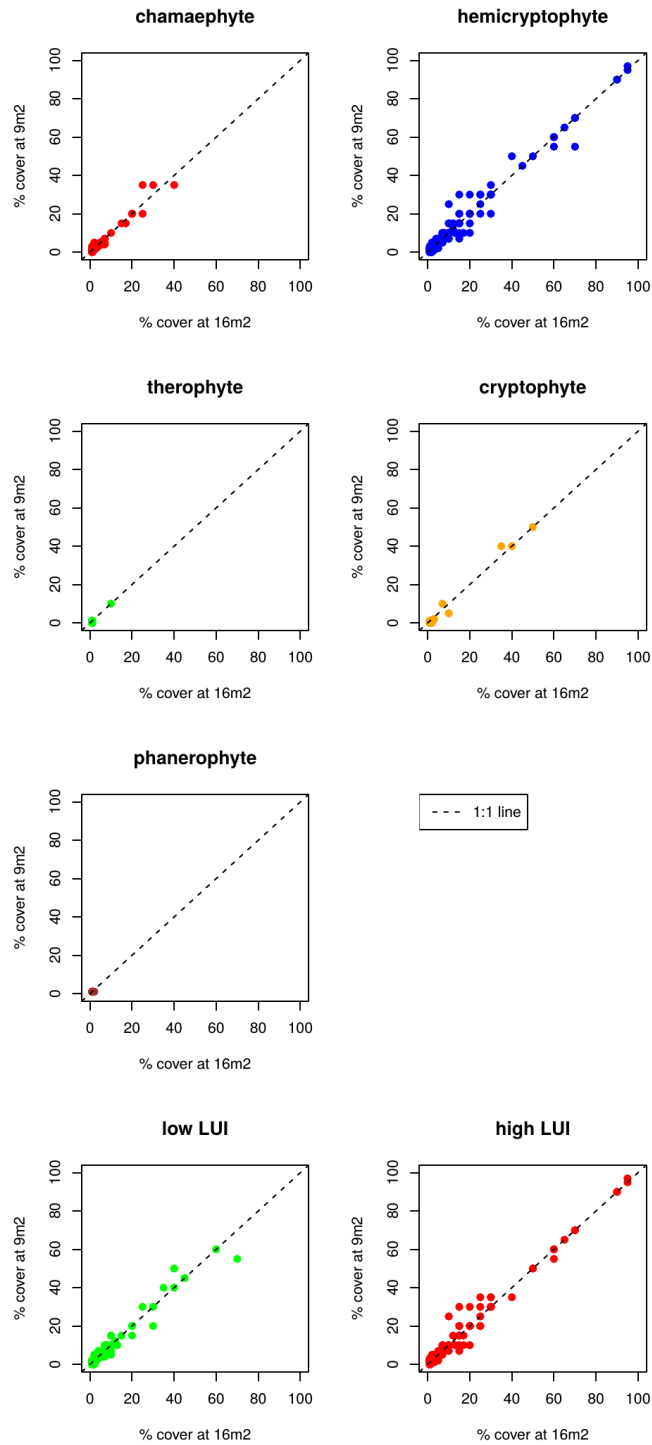
490 **Supplementary Table 12.** Correlation coefficients for CWM's versus functional, phylogenetic metrics
 491 and evenness, US dataset. Pearson correlation coefficients for BioDIV (upper part) and combined US
 492 real world community properties (lower part). Bold values are mean absolute correlation coefficients
 493 for the columns, the overall mean is the absolute mean across all column averages.

BioDIV	FEve	FDiv	FDis	RaoQ	SEve	MPD	MNTD
leaf_area	0.13	0.2	0.4	0.47	-0.17	0.24	0.1
SLA	0.15	0.18	0.22	0.26	-0.19	0.15	-0.08
leaf_drymass	0.04	0.1	0.26	0.32	-0.1	0.12	0.07
LDMC	-0.11	-0.14	-0.3	-0.35	0.1	-0.25	-0.12
leaf_N	0.05	0.03	0.22	0.26	0.04	0.23	0.13
leaf_P	-0.02	-0.02	0.13	0.17	0.08	0.09	0.13
height	0.01	0.08	0.01	0.02	-0.15	-0.06	-0.07
seedmass	0.11	0.15	0.37	0.44	-0.13	0.27	0.14
avg_abs	0.08	0.11	0.24	0.29	0.12	0.18	0.1
overall mean	0.16						

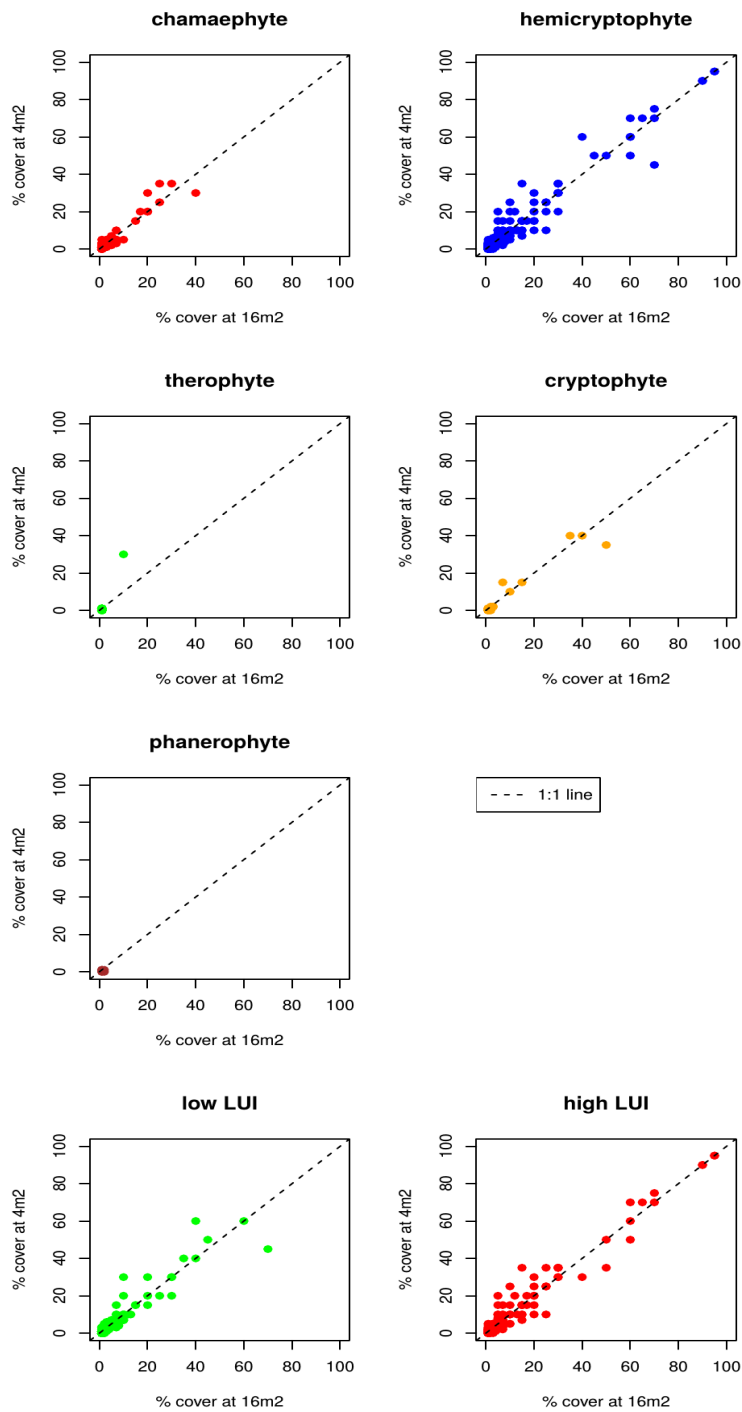
USA RW	FEve	FDiv	FDis	RaoQ	SEve	MPD	MNTD
leaf_area	0.09	0.13	0.5	0.6	-0.07	0.38	0.12
SLA	0.06	0.08	0.15	0.14	-0.21	0.33	0.23
leaf_drymass	0.07	0.11	0.51	0.65	-0.04	0.32	0.1
LDMC	-0.11	-0.06	-0.27	-0.23	0.06	-0.33	-0.13
leaf_N	-0.18	-0.36	-0.29	-0.2	0.18	-0.2	0.13
leaf_P	0.13	-0.03	0.33	0.29	-0.01	0.48	0.38
height	-0.23	-0.18	-0.41	-0.28	0.08	-0.55	-0.24
seedmass	0.04	-0.03	0.29	0.36	0.01	0.29	0.29
avg_abs	0.11	0.12	0.34	0.34	0.08	0.36	0.2
overall mean	0.22						



494 **Supplementary Figure 6.** Alternative versions of Fig. 1a showing Exploratories land-use intensity
 495 gradients. PCA biplot of Jena Experiment (orange dots and ellipse) and Biodiversity Exploratories
 496 communities (gray shaded dots, blue circle) showing land-use intensity (LUI), mowing, grazing and
 497 fertilization intensity of Exploratories plots. Land use intensity indices have been standardized across
 498 years 2008-2015 and across all three Exploratories regions¹³. Gray shades from light to dark depict
 499 increasing land-use intensity using categorical variables calculated based on the quantiles of the
 500 different land-use data.

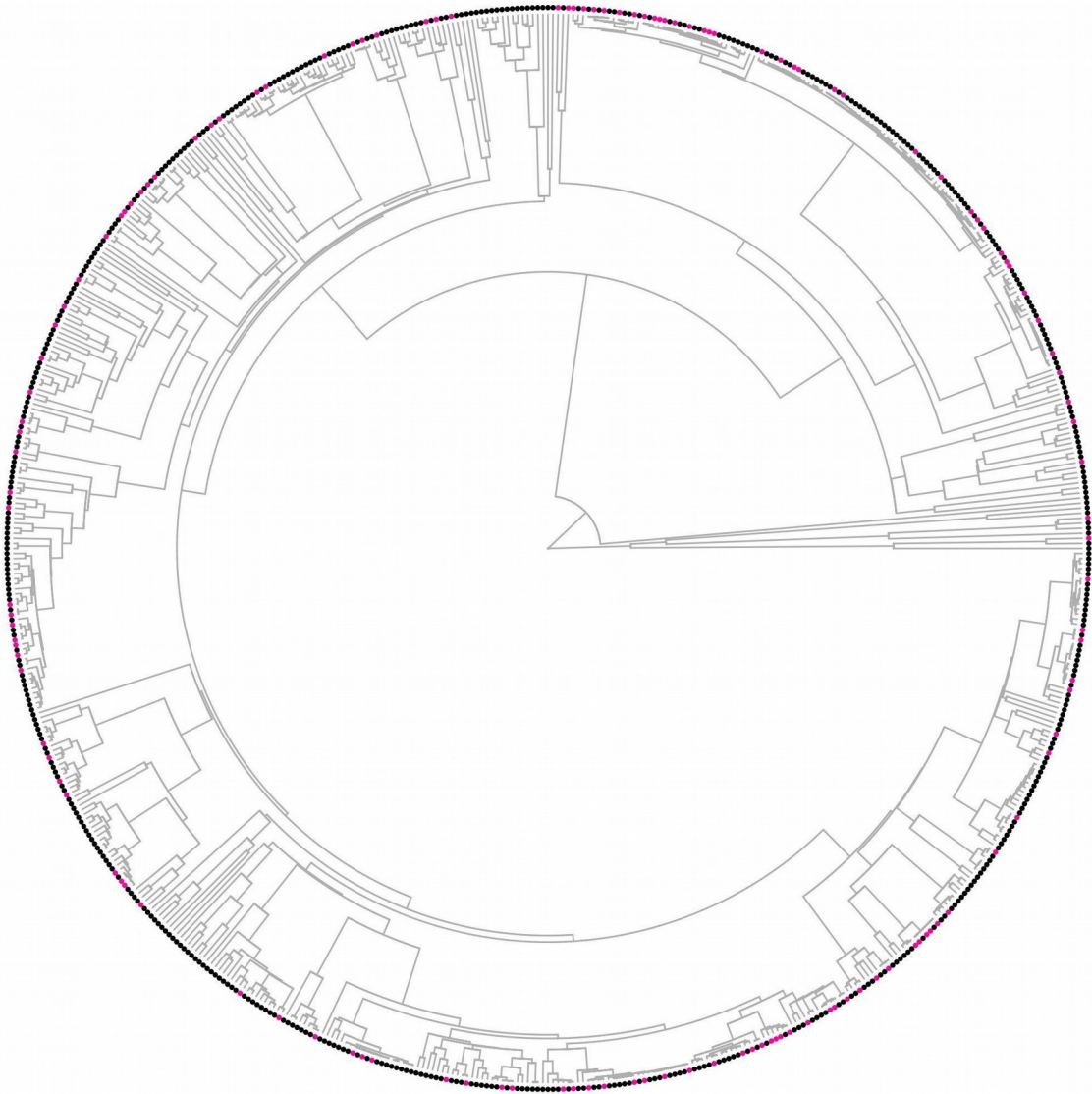


501 **Supplementary Figure 7.** Cover versus vegetation survey size scaling sensitivity check for
 502 Biodiversity Exploratories (German real world). Here, 16 to 9 m², the latter being the vegetation survey
 503 area of the Jena main and Jena real world plots. For this figure, species were sorted into lifeforms using
 504 the R package “TR8”¹⁴ and information from The Ecological Flora Database¹⁵.



505 **Supplementary Figure 8.** Cover versus vegetation survey size scaling sensitivity check for
 506 Biodiversity Exploratories (German real world). Here, 16 to 4 m², the latter resembling the vegetation
 507 survey area of the Jena invasion plots. For this figure, species were sorted into lifeforms using the R
 508 package “TR8”¹⁴ and information from The Ecological Flora Database¹⁵.

509 **Supplementary Figure 9.** Phylogenetic backbone tree (one example of the 50 replicates).
510 Overall 664 species. 132 species (19.9%, pink dots) that were not present in the backbone phylogeny
511 used to build this tree were randomly inserted into their genera (see methods for details).



512 **Supplementary Table 13.** TRY references for plant species trait data from TRY ¹⁶ requests 2968 and
513 4106. Data sources are sorted by the region their trait data have been used for (Germany=GER or
514 USA). Note that, as mentioned in the main text, trait data for the USA dataset have been complemented
515 by data from Cedar Creek plant trait assessments by Jane Catford, Peter Reich and Jeannine Cavender-
516 Bares ^{11,17}.

Region	TRY_Dataset	Reference
GER	Altitudinal Vicariants Spain	89
GER	ArtDeco Database	92
GER	BASECO: a floristic and ecological database of Mediterranean French flora	100
GER	BiolFlor Database	110
GER	BiolFlor Database	28
GER	BiolFlor Database	29
GER	BiolFlor Database	30
GER	BiolFlor Database	31
GER	BiolFlor Database	18
GER	BiolFlor Database	23
GER	BiolFlor Database	24
GER	BiolFlor Database	25
GER	BiolFlor Database	26
GER	BiolFlor Database	27
GER	BiolFlor Database	37
GER	BiolFlor Database	38
GER	BiolFlor Database	39
GER	BROT Plant Trait Database	40
GER	BROT Plant Trait Database	41
GER	Cedar Creek prairie plants (leaf, seed, dispersule, height, plant, root)	unpub.
GER	Climbing plants trait dataset	90
GER	Ecological Flora of the British Isles	24
GER	Functional traits explaining variation in plant life history strategies	47
GER	GLOPNET - Global Plant Trait Network Database	91
GER	GLOPNET - Global Plant Trait Network Database	53
GER	Grassland Plant Trait Database	54
GER	Grassland Plant Trait Database	55
GER	Harze Trait Intravar: SLA, LDMC and Plant Height for Calcareous Grassland Species in South Belgium	unpub.
GER	Herbs Water Relations on Soil Moisture Gradients	56
GER	Hydrophytes Traits Database	93
GER	Italian Alps Plant Traits Database	94
GER	Italian Alps Plant Traits Database	95

GER	KEW Seed Information Database (SID)	58
GER	KEW Seed Information Database (SID)	59
GER	Leaf Allometry Dataset	96
GER	Leaf Allometry Dataset	97
GER	Leaf and Whole Plant Traits Database	98
GER	Leaf and Whole Plant Traits Database	99
GER	Leaf and Whole Plant Traits Database	101
GER	Leaf and Whole Plant Traits Database	102
GER	Leaf and Whole Plant Traits Database	103
GER	Leaf and Whole Plant Traits Database	104
GER	Leaf and Whole Plant Traits Database	105
GER	Leaf and Whole Plant Traits Database	106
GER	Leaf and Whole Plant Traits Database	107
GER	Leaf and Whole Plant Traits Database	108
GER	Leaf and Whole Plant Traits Database	109
GER	Leaf and Whole Plant Traits Database	unpub.
GER	Leaf and Whole Plant Traits Database	111
GER	Leaf Area, Dry Mass and SLA Dataset	unpub.
GER	Leaf Economic Traits Across Varying Environmental Conditions	61
GER	Leaf N-Retention Database	62
GER	Leaf Physiology Database	112
GER	Leaf Physiology Database	unpub.
GER	Leaf Structure and Economic Spectrum	66
GER	Leaf Structure and Economic Spectrum	67
GER	Leaf Structure and Economic Spectrum	68
GER	Leaf Structure, Venation and Economic Spectrum	69
GER	Leaf Structure, Venation and Economic Spectrum	70
GER	Leaf Structure, Venation and Economic Spectrum	60
GER	Leaf Structure, Venation and Economic Spectrum	71
GER	Leaf traits from Baltic Island species	113
GER	Leaf Traits in Central Apennines Beech Forests	114
GER	Northern mixed-grass prairie species traits - Wyoming, USA	unpub.
GER	Nutrient Resorption Efficiency Database	115
GER	Nutrient Resorption Efficiency Database	116

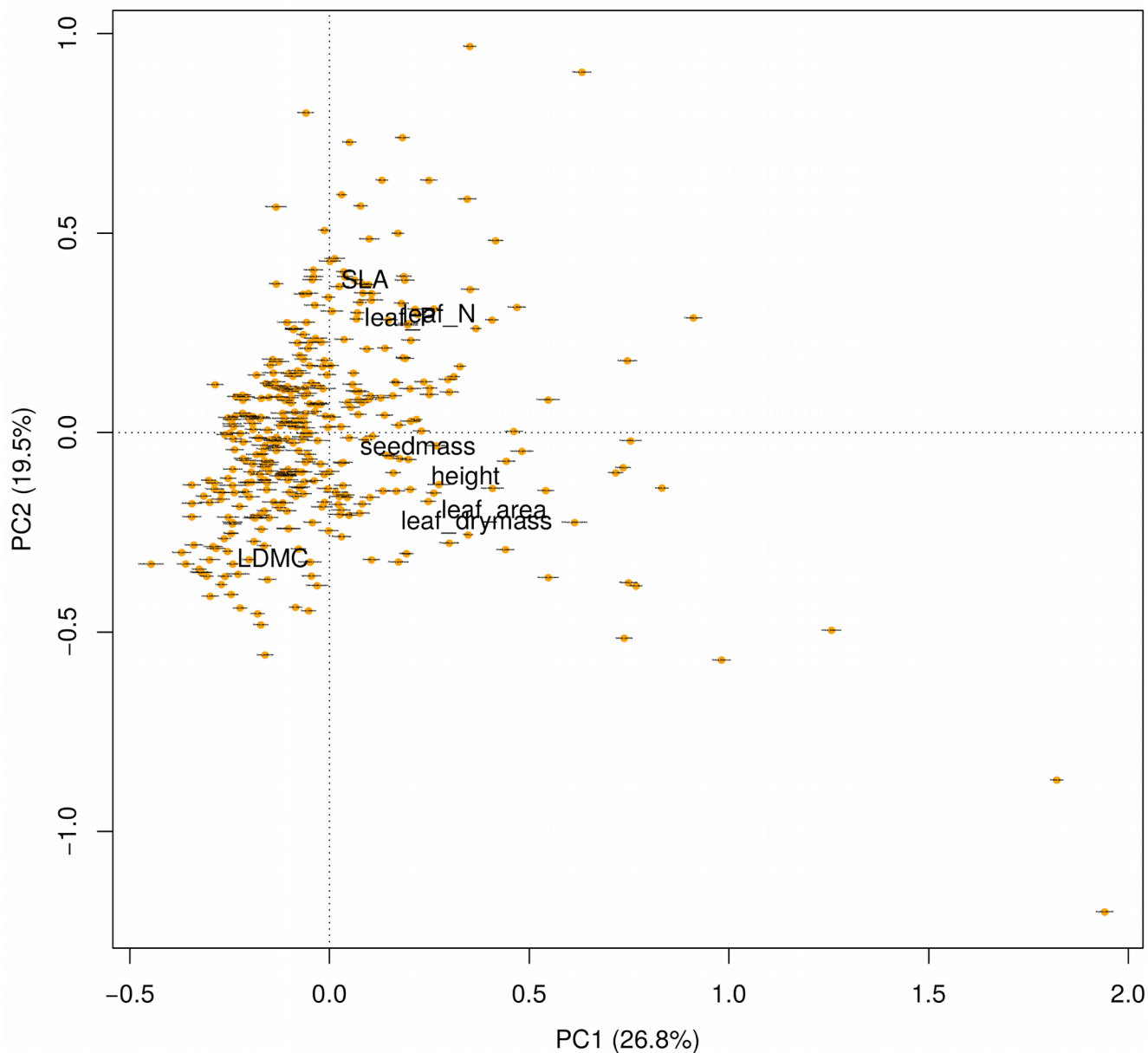
GER	Photosynthesis Traits Worldwide	74
GER	PLANTATT - Attributes of British and Irish Plants	117
GER	Plant Coastal Dune Traits (France, Aquitaine)	unpub.
GER	Plant Traits of Acidic Grasslands in Central Spain	118
GER	Plant traits of grassland species	81
GER	Reich-Oleksyn Global Leaf N, P Database	51
GER	Reproductive Allocation	82
GER	Seed Information Database (SID) Seed Mass 2010	83
GER	Sheffield & Spain Woody Database	119
GER	Sheffield & Spain Woody Database	120
GER	Sheffield & Spain Woody Database	121
GER	Sheffield & Spain Woody Database	122
GER	Specific leaf area responses to environmental gradients through space and time	123
GER	The LEDA Traitbase	84
GER	The Netherlands Plant Traits Database	124
GER	The Netherlands Plant Traits Database	125
GER	The VISTA Plant Trait Database	85
GER	The VISTA Plant Trait Database	86
GER	The VISTA Plant Trait Database	87
GER	The VISTA Plant Trait Database	88
GER	The Xylem/Phloem Database	126
GER	The Xylem/Phloem Database	127
GER	Traits of the Hungarian flora	128
GER	UV-B Radiation Sensitivity of Hieracium Pilosella	129
GER	Wetland Dunes Database	130
GER	Wetland Dunes Database	131
GER	Wetland Dunes Database	unpub.
GER	Wetland Dunes Database	132
GER	Wetland Dunes Database	133
GER	Whole Plant Hydraulic Conductance	134
USA	ArtDeco Database	135
USA	BiolFlor Database	110
USA	BiolFlor Database	19
USA	BiolFlor Database	20
USA	BiolFlor Database	21

USA	BiolFlor Database	22
USA	BiolFlor Database	18
USA	BiolFlor Database	23
USA	BiolFlor Database	24
USA	BiolFlor Database	25
USA	BiolFlor Database	26
USA	BiolFlor Database	27
USA	BiolFlor Database	28
USA	BiolFlor Database	29
USA	BiolFlor Database	30
USA	BROT Plant Trait Database	31
USA	BROT Plant Trait Database	32
USA	California Coastal Grassland Database	33
USA	Cedar Creek Savanna SLA, C, N Database	17
USA	Cold Tolerance, Seed Size and Height of North American Forest Tree Species	unpub.
USA	ECOCRAFT	34
USA	ECOCRAFT	35
USA	ECOCRAFT	36
USA	Floridian Leaf Traits Database	37
USA	Functional traits explaining variation in plant life history strategies	38
USA	Functional Traits of Graminoids in Semi-Arid Steppes Database	39
USA	Functional Traits of Graminoids in Semi-Arid Steppes Database	40
USA	Global 15N Database	41
USA	Global A, N, P, SLA Database	42
USA	GLOPNET - Global Plant Trait Network Database	43
USA	GLOPNET - Global Plant Trait Network Database	44
USA	Grassland Plant Trait Database	45
USA	Grassland Plant Trait Database	46
USA	Herbs Water Relations on Soil Moisture Gradients	47
USA	Jasper Ridge leaf chemistry data	48
USA	KEW Seed Information Database (SID)	49
USA	KEW Seed Information Database (SID)	50
USA	Leaf Area, Dry Mass and SLA Dataset	unpub.
USA	Leaf economics spectrum and venation networks in <i>Populus tremuloides</i>	51
USA	Leaf Economic Traits Across Varying Environmental Conditions	52
USA	Leaf N-Retention Database	53
USA	Leaf Photosynthesis and Nitrogen at Oak Ridge Dataset	54
USA	Leaf Structure and Chemistry	55

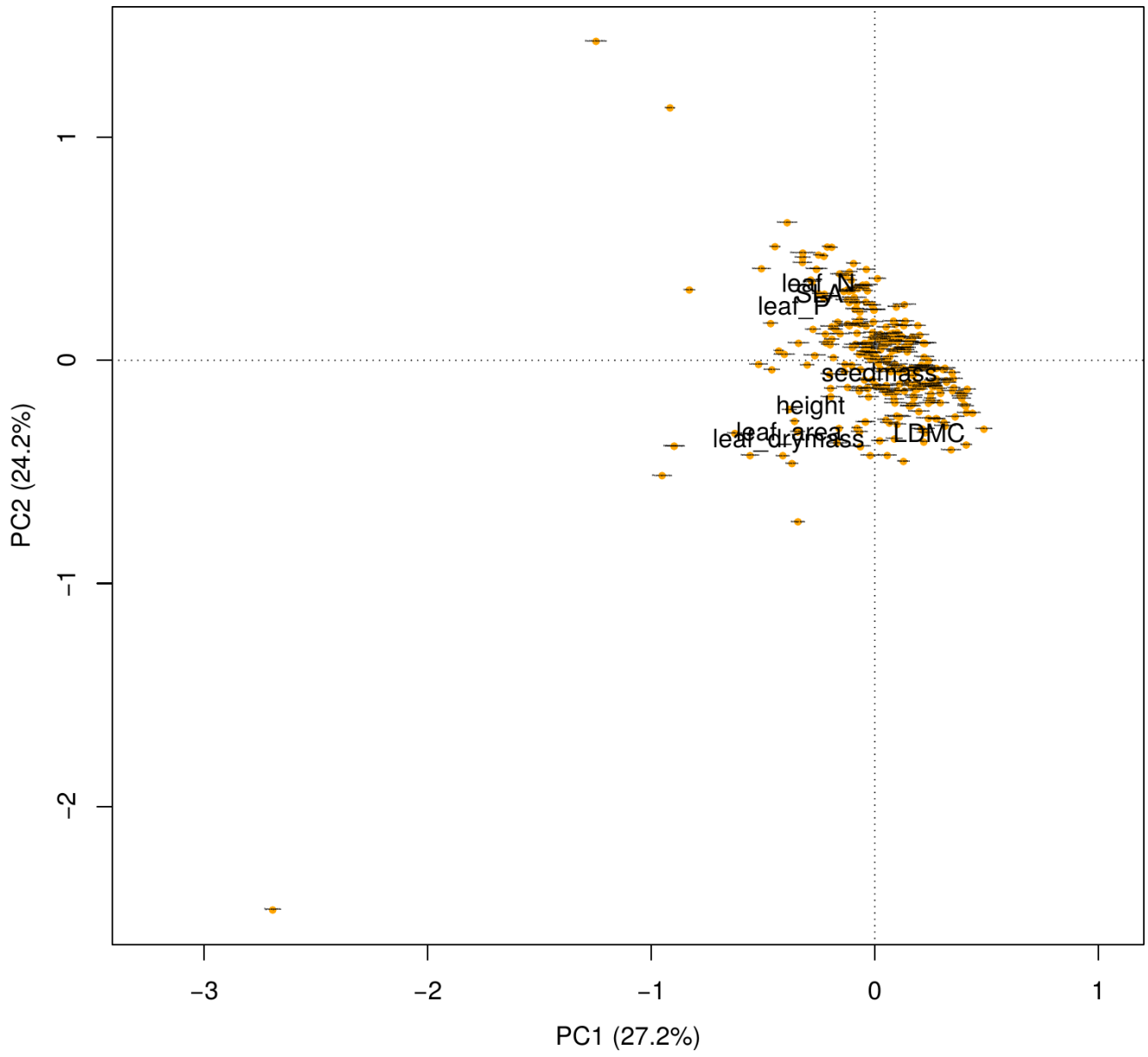
USA	Leaf Structure and Chemistry	56
USA	Leaf Structure and Economics Spectrum	57
USA	Leaf Structure and Economics Spectrum	58
USA	Leaf Structure and Economics Spectrum	59
USA	Leaf Structure, Venation and Economic Spectrum	60
USA	Leaf Structure, Venation and Economic Spectrum	61
USA	Leaf Structure, Venation and Economic Spectrum	61
USA	Leaf Structure, Venation and Economic Spectrum	62
USA	Midwestern and Southern US Herbaceous Species Trait Database	unpub.
USA	New York Old Field Plant Traits Database	63
USA	Northern mixed-grass prairie species traits - Wyoming, USA	unpub.
USA	Old fields of Eastern US (Siefert Data)	64
USA	Photosynthesis and Leaf Characteristics Database	unpub.
USA	Photosynthesis Traits Worldwide	65
USA	Plant Traits for Pinus and Juniperus Forests in Arizona	66
USA	Plant Traits for Pinus and Juniperus Forests in Arizona	67
USA	Plant Traits of Canadian Forests	68
USA	Plant Traits of Canadian Forests	78
USA	Plant Traits of Canadian Forests	79
USA	Plant Traits of Canadian Forests	80
USA	Plant traits of grassland species	81
USA	Reich-Oleksyn Global Leaf N, P Database	61
USA	Reproductive Allocation	82
USA	Rocky Mountain Biological Laboratory WSR/gradient plant traits	unpub.
USA	Seed Information Database (SID) Seed Mass 2010	83
USA	The LEDA Traitbase	84
USA	The VISTA Plant Trait Database	85
USA	The VISTA Plant Trait Database	86
USA	The VISTA Plant Trait Database	87
USA	The VISTA Plant Trait Database	88
USA	Tundra Plant Traits Database	unpub.

517 **Supplementary Figure 10.** PCA of plant species and their traits for German and US comparison.
518 Each point represents the traits of a single species in the German or US dataset. For obvious outliers,
519 the ability of each species to score such extreme values was individually confirmed e.g. by checking
520 that certain species have unusually large leaf area or leaf nitrogen content. Note that since most of the
521 calculated community properties are relative-abundance weighted, these outliers have little impact on
522 the community properties of a given plant community.

523 Germany:



524 USA:



525 **Supplementary Table 14.** Percentage cover of species with trait information for the German and US
526 datasets. For each region (Germany = GER and USA = US) and each of the eight functional traits used
527 in the analysis, the table shows the percentage of total cover occupied by species with available,
528 unimputed trait information across all plots and all datasets of the respective region. As an example, the
529 first row shows that in the German dataset, 97.6 % of the total cover across all plots and all datasets
530 was occupied by species we had original (TRY or personal communication) trait data for. After genus-
531 based inference and imputation of trait data to fill all gaps, 2 % of species in the German dataset (8 of
532 373 species) and 8 % of species in the US dataset (23 of 291 species) had identical trait information
533 across all traits.

region	trait	summed cover %
GER	leaf_area	97.6
GER	SLA	89.7
GER	leaf_drymass	61.6
GER	LDMC	65.0
GER	leaf_N	85.2
GER	leaf_P	78.3
GER	height	99.9
GER	seedmass	99.0
US	leaf_area	99.0
US	SLA	98.4
US	leaf_drymass	99.5
US	LDMC	97.7
US	leaf_N	92.7
US	leaf_P	73.2
US	height	99.4
US	seedmass	99.4

534 **Supplementary Table 15.** Species with altered trait values to avoid Gower dissimilarity zeros.
 535 Species are sorted by region (GER=Germany, US=USA) and by the percentage shift that their trait
 536 values were subject to. In two cases in the US dataset, there were three same-genus species with
 537 identical trait values and here two of them needed different shifts in order to obtain non-zero Gower
 538 dissimilarity values.

GER, 0.001% shift up

Acinos arvensis
Arabidopsis thaliana
Chenopodium sp
Clinopodium acinos
Echinochloa crus-galli
Epilobium sp
Listera ovata
Mentha aquatica
Sesleria albicans
Orobanche caryophyllacea
Rubus sp
Rumex thyrsiflorus
Poa angustifolia
Potentilla neumanniana
Veronica spicata

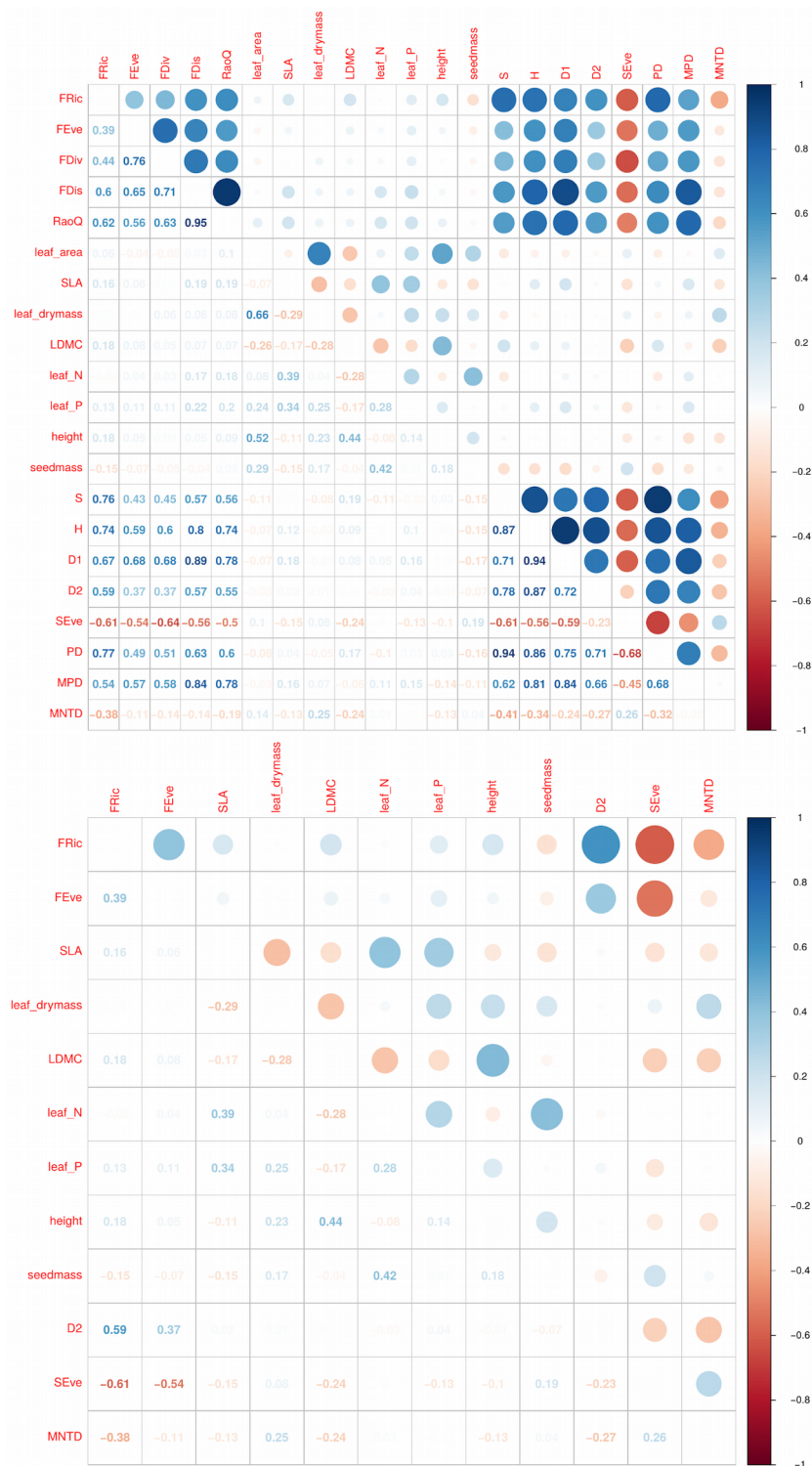
US 0.001% shift up

Achillea sp
Agrostis sp
Allium stellatum
Antennaria plantaginifolia
Calamagrostis sp
Echinacea serotina
Euphorbia geyeri
Galium sp
Gnaphalium sp
Melilotus sp
Parthenocissus inserta
Polygala sp
Polygonatum sp
Rhus sp
Rumex sp
Salix humilis
Solidago altissima
Stachys sp
Taraxacum sp
Tradescantia bracteata

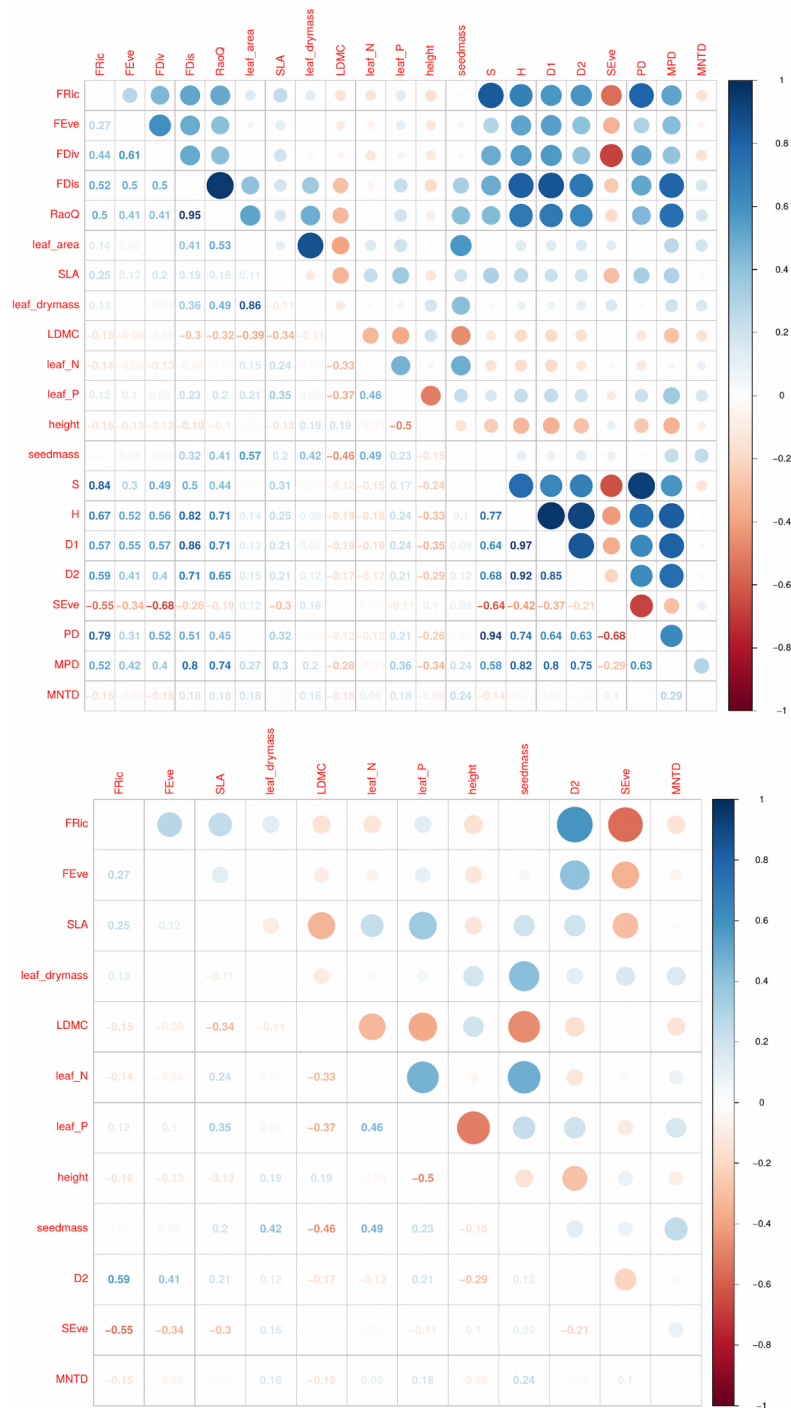
US 0.002% shift up

Antennaria sp
Tradescantia sp

539 **Supplementary Table 16.** Correlation coefficients for 21 plant community properties for the German
 540 dataset. Pearson correlation coefficients and color code (see legend) for all 21 properties (upper
 541 diagram) and the subset of 12 community properties retained after stepwise removal due to variance
 542 inflation factors above 3 (lower diagram). Diagrams were created using the “corrplot” package¹³⁶ in R.



543 **Supplementary Table 17.** Correlation coefficients for 21 plant community properties for the US
 544 dataset. Pearson correlation coefficients and color code (see legend) for all 21 properties (upper
 545 diagram) and the subset of 12 community properties retained after stepwise removal due to variance
 546 inflation factors above 3 (lower diagram). Diagrams were created using the “corrplot” package¹³⁶ in R.



547 **Supplementary References**

- 548 1. Fornara, D. A. & Tilman, D. Plant functional composition influences rates of soil carbon and
549 nitrogen accumulation. *J. Ecol.* **96**, 314–322 (2008).
- 550 2. Lange, M. *et al.* Plant diversity increases soil microbial activity and soil carbon storage. *Nat.*
551 *Commun.* **6**, 6707 (2015).
- 552 3. Ravenek, J. M. *et al.* Long-term study of root biomass in a biodiversity experiment reveals shifts in
553 diversity effects over time. *Oikos* **123**, 1528–1536 (2014).
- 554 4. Meyer, S. T. *et al.* Consistent increase in herbivory along two experimental plant diversity gradients
555 over multiple years. *Ecosphere* **8**, e01876 (2017).
- 556 5. Scheu, S. Automated measurement of the respiratory response of soil microcompartments: Active
557 microbial biomass in earthworm faeces. *Soil Biol. Biochem.* **24**, 1113–1118 (1992).
- 558 6. Eisenhauer, N. *et al.* Plant diversity effects on soil microorganisms support the singular hypothesis.
559 *Ecology* **91**, 485–496 (2010).
- 560 7. Strecker, T., Mace, O. G., Scheu, S. & Eisenhauer, N. Functional composition of plant communities
561 determines the spatial and temporal stability of soil microbial properties in a long-term plant
562 diversity experiment. *Oikos* **125**, 1743–1754 (2016).
- 563 8. Beck, T. *et al.* An inter-laboratory comparison of ten different ways of measuring soil microbial
564 biomass C. *Soil Biol. Biochem.* **29**, 1023–1032 (1997).
- 565 9. Hacker, N. *et al.* Plant diversity shapes microbe-rhizosphere effects on P mobilisation from organic
566 matter in soil. *Ecol. Lett.* **18**, 1356–1365 (2015).
- 567 10. Eivazi, F. & Tabatabai, M. A. Phosphatases in soils. *Soil Biol. Biochem.* **9**, 167–172 (1977).
- 568 11. Catford, J. A. *et al.* Traits linked with species invasiveness and community invasibility vary with
569 time, stage and indicator of invasion in a long-term grassland experiment. *Ecol. Lett.* **22**, 593–604
570 (2019).
- 571 12. Inouye, R. *et al.* Old-field succession on a Minnesota sand plain. *Ecology* **68**, 12–26 (1987).
- 572 13. Tilman, D. Community Invasibility, Recruitment Limitation, and Grassland Biodiversity. *Ecology*
573 **78**, 81–92 (1997).
- 574 14. Roscher, C., Schumacher, J. & Baade, J. The role of biodiversity for element cycling and trophic
575 interactions: an experimental approach in a grassland community. *Basic Appl. Ecol.* **121**, 107–121
576 (2004).
- 577 15. Fischer, M. *et al.* Implementing large-scale and long-term functional biodiversity research: The
578 Biodiversity Exploratories. *Basic Appl. Ecol.* **11**, 473–485 (2010).
- 579 16. Roscher, C. *et al.* Convergent high diversity in naturally colonized experimental grasslands is not

- 580 related to increased productivity. *Perspect. Plant Ecol. Evol. Syst.* **20**, 32–45 (2016).
- 581 17. Weisser, W. W. *et al.* Biodiversity effects on ecosystem functioning in a 15-year grassland
582 experiment: patterns, mechanisms, and open questions. *Basic Appl. Ecol.* **23**, 1–73 (2017).
- 583 18. Tilman, D. *et al.* The influence of functional diversity and composition on ecosystem processes.
584 *Science (80-.).* **277**, 1300–1302 (1997).
- 585 19. Tilman, D. Secondary succession and the pattern of plant dominance along experimental nitrogen
586 gradients. *Ecol. Monogr.* **57**, 189–214 (1987).
- 587 20. Oksanen, J. *et al.* Vegan: community ecology package. *R Packag. version 2.3-4* (2016).
- 588 21. Adler, D. & Kelly, T. vioplot: violin plot. R. *R Packag. version 0.3.0* (2018).
- 589 22. Blüthgen, N. *et al.* A quantitative index of land-use intensity in grasslands: Integrating mowing,
590 grazing and fertilization. *Basic Appl. Ecol.* **13**, 207–220 (2012).
- 591 23. Bocci, G. TR8: An R package for easily retrieving plant species traits. *Methods Ecol. Evol.* **6**, 347–
592 350 (2015).
- 593 24. Fitter, A. & Peat, H. The Ecological Flora Database. *J. Ecol.* **82**, 415–425 (1994).
- 594 25. Kattge, J. *et al.* TRY - a global database of plant traits. *Glob. Chang. Biol.* **17**, 2905–2935 (2011).
- 595 26. Willis, C. G. *et al.* Phylogenetic community structure in Minnesota oak savanna is influenced by
596 spatial extent and environmental variation. *Ecography (Cop.)*. **33**, 565–577 (2010).
- 597 27. Klotz, S. & Kühn, I. Indikatoren zum anthropogenen Einfluss auf die Vegetation. *Schriftenr. für Veg.*
598 **38**, (2002).
- 599 28. Durka, W. Blüten- und Reproduktionsbiologie. *Schriftenr. für Veg.* **38**, (2002).
- 600 29. Durka, W. Chromosomenzahlen, Ploidiestufen und DNA-Gehalt. *Schriftenr. für Veg.* **38**, (2002).
- 601 30. Durka, W. Phylogenie der Farn- und Blütenpflanzen Deutschlands. *Schriftenr. für Veg.* **38**, (2002).
- 602 31. Klotz, S. & Kühn, I. Blattmerkmale. *Schriftenr. für Veg.* **38**, (2002).
- 603 32. Klotz, S. & Kühn, I. Ökologische Strategietypen. *Schriftenr. für Veg.* **38**, (2002).
- 604 33. Klotz, S. & Kühn, I. Soziologische Bindung der Arten. *Schriftenr. für Veg.* **38**, (2002).
- 605 34. Krumbiegel, A. Morphologie der vegetativen Organe (außer Blätter). *Schriftenr. für Veg.* **38**, (2002).
- 606 35. Kühn, I. & Klotz, S. Angaben zu den Arealen. *Schriftenr. für Veg.* **38**, (2002).
- 607 36. Kühn, I. & Klotz, S. Floristischer Status und gebietsfremde Arten. *Schriftenr. für Veg.* **38**, (2002).
- 608 37. Kühn, I., Durka, W. & Klotz, S. BiolFlor - A new plant-trait database as a tool for plant invasion
609 ecology. *Divers. Distrib.* **10**, 363–365 (2004).

- 610 38. Otto, B. Merkmale von Samen, Früchten, generativen Germinulen und generativen Diasporen.
611 *Schriftenr. für Veg.* **38**, (2002).
- 612 39. Trefflich, A., Klotz, S. & Kühn, I. Blühphänologie. *Schriftenr. für Veg.* **38**, (2002).
- 613 40. Paula, S. & Pausas, J. G. Burning seeds: Germinative response to heat treatments in relation to
614 resprouting ability. *J. Ecol.* **96**, 543–552 (2008).
- 615 41. Paula, S. *et al.* Fire-related traits for plant species of the Mediterranean Basin. *Ecology* **90**, 1420
616 (2009).
- 617 42. Sandel, B., Corbin, J. D. & Krupa, M. Using plant functional traits to guide restoration: A case study
618 in California coastal grassland. *Ecosphere* **2**, 1–16 (2011).
- 619 43. Medlyn, B. E. & Jarvis, P. G. Design and use of a database of model parameters from elevated [CO₂]
620 experiments. *Ecol. Modell.* **124**, 69–83 (1999).
- 621 44. Medlyn, B. E. *et al.* Stomatal conductance of forest species after long-term exposure to elevated
622 CO₂ concentration: A synthesis. *New Phytol.* **149**, 247–264 (2001).
- 623 45. Medlyn, B. E. *et al.* Effects of elevated [CO₂] on photosynthesis in European forest species: A meta-
624 analysis of model parameters. *Plant, Cell Environ.* **22**, 1475–1495 (1999).
- 625 46. Cavender-Bares, J., Keen, A. & Miles, B. Phylogenetic structure of Floridian plant communities
626 depends on taxonomic and spatial scale. *Ecology* **87**, 109–122 (2006).
- 627 47. Adler, P. B. *et al.* Functional traits explain variation in plant lifehistory strategies. *Proc. Natl. Acad.*
628 *Sci. U. S. A.* **111**, 740–745 (2014).
- 629 48. Adler, P. A comparison of livestock grazing effects on sagebrush steppe, USA, and Patagonian
630 steppe, Argentina. (Colorado State University, 2003).
- 631 49. Adler, P. B., Milchunas, D. G., Lauenroth, W. K., Sala, O. E. & Burke, I. C. Functional traits of
632 graminoids in semi-arid steppes: A test of grazing histories. *J. Appl. Ecol.* **41**, 653–663 (2004).
- 633 50. Craine, J. M. *et al.* Global patterns of foliar nitrogen isotopes and their relationships with climate,
634 mycorrhizal fungi, foliar nutrient concentrations, and nitrogen availability. *New Phytol.* **183**, 980–
635 992 (2009).
- 636 51. Reich, P. B., Oleksyn, J. & Wright, I. J. Leaf phosphorus influences the photosynthesis-nitrogen
637 relation: A cross-biome analysis of 314 species. *Oecologia* **160**, 207–212 (2009).
- 638 52. Wright, I. J. *et al.* The worldwide leaf economics spectrum. *Nature* (2004). doi:10.1038/nature02403
- 639 53. Wright, I. J. *et al.* Irradiance, temperature and rainfall influence leaf dark respiration in woody
640 plants: Evidence from comparisons across 20 sites. *New Phytol.* **169**, 309–319 (2006).
- 641 54. Takkis, K. Changes in plant species richness and population performance in response to habitat loss
642 and fragmentation. *Diss. Biol. Univ. Tartu.* **255**, (2014).

- 643 55. Takkis, K., Saar, L., Pärtel, M. & Helm, A. Effect of environment and landscape on the traits of six
644 plant species in fragmented grasslands. *Prep.*
- 645 56. Sheremetev, S. *Herbs on the soil moisture gradient (water relations and the structural-functional
646 organization)*. KMK, Moscow (2005).
- 647 57. Dahlin, K. M., Asner, G. P. & Field, C. B. Environmental and community controls on plant canopy
648 chemistry in a Mediterranean-type ecosystem. *Proc. Natl. Acad. Sci. U. S. A.* **110**, 6895–6900 (2013).
- 649 58. Royal Botanical Gardens KEW. Seed Information Database (SID). (2008).
- 650 59. Royal Botanical Gardens KEW. Seed Information Database (SID). (2011).
- 651 60. Blonder, B., Violle, C. & Enquist, B. J. Assessing the causes and scales of the leaf economics
652 spectrum using venation networks in *Populus tremuloides*. *J. Ecol.* **101**, 981–989 (2013).
- 653 61. Wright, J. P. & Sutton-Grier, A. Does the leaf economic spectrum hold within local species pools
654 across varying environmental conditions? *Funct. Ecol.* **26**, 1390–1398 (2012).
- 655 62. de Vries, F. T. & Bardgett, R. D. Plant community controls on short-term ecosystem nitrogen
656 retention. *New Phytol.* **210**, 861–874 (2016).
- 657 63. Wilson, K. B., Baldocchi, D. D. & Hanson, P. J. Spatial and seasonal variability of photosynthetic
658 parameters and their relationship to leaf nitrogen in a deciduous forest. *Tree Physiol.* **20**, 565–578
659 (2000).
- 660 64. Auger, S. L'importance de la variabilité interspécifique des traits fonctionnels par rapport à la
661 variabilité intraspécifique chez les jeunes arbres en forêt mature. (Université de Sherbrooke,
662 Sherbrooke (Quebec), 2012).
- 663 65. Auger, S. & Shipley, B. Inter-specific and intra-specific trait variation along short environmental
664 gradients in an old-growth temperate forest. *J. Veg. Sci.* **24**, 419–428 (2013).
- 665 66. Pierce, S., Brusa, G., Vagge, I. & Cerabolini, B. E. L. Allocating CSR plant functional types: The use
666 of leaf economics and size traits to classify woody and herbaceous vascular plants. *Funct. Ecol.* **27**,
667 1002–1010 (2013).
- 668 67. Pierce, S., Ceriani, R. M., De Andreis, R., Luzzaro, A. & Cerabolini, B. The leaf economics
669 spectrum of Poaceae reflects variation in survival strategies. *Plant Biosyst.* **141**, 337–343 (2007).
- 670 68. Pierce, S., Luzzaro, A., Caccianiga, M., Ceriani, R. M. & Cerabolini, B. Disturbance is the principal
671 α -scale filter determining niche differentiation, coexistence and biodiversity in an alpine community.
672 *J. Ecol.* **95**, 698–706 (2007).
- 673 69. Blonder, B. *et al.* The shrinkage effect biases estimates of paleoclimate. *Am. J. Bot.* **99**, 1756–1763
674 (2012).
- 675 70. Blonder, B. *et al.* Testing models for the leaf economics spectrum with leaf and whole-plant traits in

- 676 *Arabidopsis thaliana*. *AoB Plants* **7**, plv049 (2015).
- 677 71. Blonder, B., Violle, C., Bentley, L. P. & Enquist, B. J. Venation networks and the origin of the leaf
678 economics spectrum. *Ecol. Lett.* **14**, 91–100 (2011).
- 679 72. Siefert, A. Spatial patterns of functional divergence in old-field plant communities. *Oikos* **121**, 907–
680 914 (2012).
- 681 73. Siefert, A., Fridley, J. D. & Ritchie, M. E. Community functional responses to soil and climate at
682 multiple spatial scales: When does intraspecific variation matter? *PLoS One* **9**, e111189 (2014).
- 683 74. Maire, V. *et al.* Global effects of soil and climate on leaf photosynthetic traits and rates. *Glob. Ecol.*
684 *Biogeogr.* **24**, 706–717 (2015).
- 685 75. Laughlin, D. C., Leppert, J. J., Moore, M. M. & Sieg, C. H. A multi-trait test of the leaf-height-seed
686 plant strategy scheme with 133 species from a pine forest flora. *Funct. Ecol.* **24**, 493–501 (2010).
- 687 76. Laughlin, D. C., Fulé, P. Z., Huffman, D. W., Crouse, J. & Laliberté, E. Climatic constraints on trait-
688 based forest assembly. *J. Ecol.* **99**, 1489–1499 (2011).
- 689 77. Bond-Lamberty, B., Wang, C. & Gower, S. T. Aboveground and belowground biomass and sapwood
690 area allometric equations for six boreal tree species of northern Manitoba. *Can. J. For. Res.* **32**,
691 1441–1450 (2002).
- 692 78. Bond-Lamberty, B., Wang, C., Gower, S. T. & Norman, J. Leaf area dynamics of a boreal black
693 spruce fire chronosequence. *Tree Physiol.* **22**, 993–1001 (2002).
- 694 79. Bond-Lamberty, B., Wang, C. & Gower, S. T. The use of multiple measurement techniques to refine
695 estimates of conifer needle geometry. *Can. J. For. Res.* **33**, 101–105 (2003).
- 696 80. Bond-Lamberty, B., Wang, C. & Gower, S. T. Net primary production and net ecosystem production
697 of a boreal black spruce wildfire chronosequence. *Glob. Chang. Biol.* **10**, 473–487 (2004).
- 698 81. La Pierre, K. J. & Smith, M. D. Functional trait expression of grassland species shift with short- and
699 long-term nutrient additions. *Plant Ecol.* **216**, 307–318 (2015).
- 700 82. Manning, P., Houston, K. & Evans, T. Shifts in seed size across experimental nitrogen enrichment
701 and plant density gradients. *Basic Appl. Ecol.* **10**, 300–308 (2009).
- 702 83. Royal Botanical Gardens KEW. Seed Information Database (SID). (2014).
- 703 84. Kleyer, M. *et al.* The LEDA Traitbase: A database of life-history traits of the Northwest European
704 flora. *J. Ecol.* **96**, 1266–1274 (2008).
- 705 85. Fortunel, C. *et al.* Leaf traits capture the effects of land use changes and climate on litter
706 decomposability of grasslands across Europe. *Ecology* **90**, 598–611 (2009).
- 707 86. Garnier, E. *et al.* Assessing the effects of land-use change on plant traits, communities and ecosystem
708 functioning in grasslands: A standardized methodology and lessons from an application to 11

- 709 European sites. *Ann. Bot.* **99**, 967–985 (2007).
- 710 87. Pakeman, R. J. *et al.* Impact of abundance weighting on the response of seed traits to climate and
711 land use. *J. Ecol.* **96**, 355–366 (2008).
- 712 88. Pakeman, R. J. *et al.* Relative climatic, edaphic and management controls of plant functional trait
713 signatures. *J. Veg. Sci.* **20**, 148–15 (2009).
- 714 89. Milla, R. & Reich, P. Multi-trait interactions , not phylogeny , fine-tune leaf size reduction with
715 increasing altitude. *Ann. Bot.* **107**, 455–465 (2011).
- 716 90. Gallagher, R. V. & Leishman, M. R. A global analysis of trait variation and evolution in climbing
717 plants. *J. Biogeogr.* **39**, 1757–1771 (2012).
- 718 91. Wright, I. J. *et al.* The worldwide leaf economics spectrum. *Nature* **428**, 821–827 (2004).
- 719 92. Cornwell, W. K. *et al.* Plant species traits are the predominant control on litter decomposition rates
720 within biomes worldwide. *Ecol. Lett.* **11**, 1065–1071 (2008).
- 721 93. Pierce, S., Bruse, G., Sartori, M. & Cerebolini, B. Combined use of leaf size and economics traits
722 allows direct comparison of hydrophyte and terrestrial herbaceous adaptive strategies. *Ann. Bot.* **109**,
723 1047–1053 (2012).
- 724 94. Bragazza, L. Conservation priority of Italian alpine habitats: A floristic approach based on potential
725 distribution of vascular plant species. *Biodivers. Conserv.* **18**, 2823–2832 (2009).
- 726 95. Dainese, M. & Bragazza, L. Plant traits across different habitats of the Italian Alps: A comparative
727 analysis between native and alien species. *Alp. Bot.* **122**, 11–21 (2012).
- 728 96. Price, C. A. & Enquist, B. J. Scaling mass and morphology in leaves: An extension of the wbe
729 model. *Ecology* **88**, 1132–1141 (2007).
- 730 97. Price, C. A., Enquist, B. J. & Savage, V. M. A general model for allometric covariation in botanical
731 form and function. *Proc. Natl. Acad. Sci. U. S. A.* **104**, 13204–13209 (2007).
- 732 98. Kazakou, E., Vile, D., Shipley, B., Gallet, C. & Garnier, E. Co-variations in litter decomposition, leaf
733 traits and plant growth in species from a Mediterranean old-field succession. *Funct. Ecol.* **20**, 21–30
734 (2006).
- 735 99. McKenna, M. F. & Shipley, B. Interacting determinants of interspecific relative growth: Empirical
736 patterns and a theoretical explanation. *Ecoscience* **6**, 286–296 (1999).
- 737 100. Gachet, S., Véla, E. & Tatoni, T. BASECO: A floristic and ecological database of Mediterranean
738 French flora. *Biodivers. Conserv.* **14**, 1023–1034 (2005).
- 739 101. Meziane, D. & Shipley, B. Interacting components of interspecific relative growth rate: Constancy
740 and change under differing conditions of light and nutrient supply. *Funct. Ecol.* **13**, 611–62 (1999).
- 741 102. Meziane, D. & Shipley, B. Interacting determinants of specific leaf area in 22 herbaceous species:

- 742 Effects of irradiance and nutrient availability. *Plant, Cell Environ.* **22**, 447–459 (1999).
- 743 103. Pyankov, V. I., Kondratchuk, A. V. & Shipley, B. Leaf structure and specific leaf mass: The alpine
744 desert plants of the Eastern Pamirs, Tadjikistan. *New Phytol.* **143**, 31–142 (1999).
- 745 104. Shipley, B. The Use of Above-Ground Maximum Relative Growth Rate as an Accurate Predictor of
746 Whole-Plant Maximum Relative Growth Rate. *Funct. Ecol.* **3**, 771–775 (1989).
- 747 105. Shipley, B. Structured Interspecific Determinants of Specific Leaf Area in 34 Species of Herbaceous
748 Angiosperms. *Funct. Ecol.* **9**, 312–319 (1995).
- 749 106. Shipley, B. Trade-offs between net assimilation rate and specific leaf area in determining relative
750 growth rate: Relationship with daily irradiance. *Funct. Ecol.* **16**, 682–689 (2002).
- 751 107. Shipley, B. & Lechowicz, M. J. The functional co-ordination of leaf morphology, nitrogen
752 concentration, and gas exchange in 40 wetland species. *Ecoscience* **7**, 183–194 (2000).
- 753 108. Shipley, B. & Parent, M. Germination Responses of 64 Wetland Species in Relation to Seed Size,
754 Minimum Time to Reproduction and Seedling Relative Growth Rate. *Funct. Ecol.* **5**, 111–118
755 (1991).
- 756 109. Shipley, B. & Vu, T. T. Dry matter content as a measure of dry matter concentration in plants and
757 their parts. *New Phytol.* **153**, 359–364 (2002).
- 758 110. Briemle, G., Nitsche, S. & Nitsche, L. Nutzungswertzahlen für Gefäßpflanzen des Grünlandes.
759 *Schriptr. für Veg.* **38**, (2002).
- 760 111. Vile, D. Significations fonctionnelle et ecologique des traits des especes vegetales: exemple dans une
761 succession post-cultural mediterraneenne et generalisations. (2005).
- 762 112. Kattge, J., Knorr, W., Raddatz, T. & Wirth, C. Quantifying photosynthetic capacity and its
763 relationship to leaf nitrogen content for global-scale terrestrial biosphere models. *Glob. Chang. Biol.*
764 **15**, 976–991 (2009).
- 765 113. Hattermann, D., Elstner, C., Bernhardt-Römermann, M. & Eckstein, L. *Measurements from the*
766 *project 'Relative effects of local and regional factors as drivers for plant community diversity,*
767 *functional trait diversity and genetic structure of species on Baltic uplift islands' DFG: BE 4143/5-1*
768 *and EC 209/12-1.*
- 769 114. Campetella, G. *et al.* Patterns of plant trait-environment relationships along a forest succession
770 chronosequence. *Agric. Ecosyst. Environ.* **145**, 38–48 (2011).
- 771 115. Vergutz, L., Manzoni, S., Porporato, A., Novalis, R. F. & Jackson, R. B. Global resorption efficiencies
772 and concentrations of carbon and nutrients in leaves of terrestrial plants. *Ecol. Monogr.* **82**, 205–220
773 (2012).
- 774 116. Vergutz, L., Manzoni, S., Porporato, A., Novalis, R. & Jackson, R. A Global Database of Carbon and
775 Nutrient Concentrations of Green and Senesced Leaves. Data set. Available on-line

- 776 [http://daac.ornl.gov] from Oak Ridge National Laboratory Distributed Active Archive Center, Oak
777 Ridge, Tennessee. (2012).
- 778 117. Hill, M. O., Preston, C. D. & Roy, D. B. Plant Attributes of British and Irish Plants: Status, Size,
779 Life history, Geography and Habitats. *Centre for Ecology and Hydrology* (2004).
- 780 118. Peco, B., De Pablos, I., Traba, J. & Levassor, C. The effect of grazing abandonment on species
781 composition and functional traits: The case of dehesa grasslands. *Basic Appl. Ecol.* **6**, 175–183
782 (2005).
- 783 119. Castro-Díez, P., Puyravaud, J. P. & Cornelissen, J. H. C. Leaf structure and anatomy as related to leaf
784 mass per area variation in seedlings of a wide range of woody plant species and types. *Oecologia*
785 **116**, 57–66 (2000).
- 786 120. Castro-Díez, P., Puyravaud, J. P., Cornelissen, J. H. C. & Villar-Salvador, P. Stem anatomy and
787 relative growth rate in seedlings of a wide range of woody plant species and types. *Oecologia* **116**,
788 57–66 (1998).
- 789 121. Cornelissen, J. H. C. A triangular relationship between leaf size and seed size among woody species:
790 Allometry, ontogeny, ecology and taxonomy. *Oecologia* **118**, 248–255 (1999).
- 791 122. Cornelissen, J. H. C. *et al.* Functional traits of woody plants: Correspondence of species rankings
792 between field adults and laboratory-grown seedlings? *J. Veg. Sci.* **14**, 311–322 (2003).
- 793 123. Dwyer, J. M., Hobbs, R. J. & Mayfield, M. M. Specific leaf area responses to environmental
794 gradients through space and time. *Ecology* **95**, 339–410 (2014).
- 795 124. Ordoñez, J. C. *et al.* Leaf habit and woodiness regulate different leaf economy traits at a given
796 nutrient supply. *Ecology* **91**, 3218–3228 (2010).
- 797 125. Ordoñez, J. C. *et al.* Plant strategies in relation to resource supply in mesic to wet environments:
798 Does theory mirror nature? *Am. Nat.* **175**, 225–239 (2010).
- 799 126. Schweingruber, F. & Landolt, W. The Xylem Database. Swiss Federal Research Institute WSL
800 (2005).
- 801 127. Schweingruber, F. H. & Poschlod, P. Growth rings in herbs and shrubs: Life span, age determination
802 and stem anatomy. *For. Snow Landsc. Res.* **79**, 195–415 (2005).
- 803 128. Lhotsky, B., Csecserits, A., Kovács, B. & Botta-Dukát, Z. New plant trait records of the Hungarian
804 flora.
- 805 129. Beckmann, M., Hock, M., Bruelheide, H. & Erfmeier, A. The role of UV-B radiation in the invasion
806 of *Hieracium pilosella*-A comparison of German and New Zealand plants. *Environ. Exp. Bot.* **75**,
807 173–180 (2012).
- 808 130. Bakker, C., Rodenburg, J. & Van Bodegom, P. M. Effects of Ca- and Fe-rich seepage on P
809 availability and plant performance in calcareous dune soils. *Plant Soil* **275**, 111–122 (2005).

- 810 131. Bakker, C., Van Bodegom, P. M., Nelissen, H. J. M., Ernst, W. H. O. & Aerts, R. Plant responses to
811 rising water tables and nutrient management in calcareous dune slacks. *Plant Ecol.* **185**, 19–28
812 (2006).
- 813 132. Van Bodegom, P. M., Sorrell, B. K., Oosthoek, A., Bakker, C. & Aerts, R. Separating the effects of
814 partial submergence and soil oxygen demand on plant physiology. *Ecology* **89**, 193–204 (2008).
- 815 133. Van Bodegom, P. M., De Kanter, M., Bakker, C. & Aerts, R. Radial oxygen loss, a plastic property of
816 dune slack plant species. *Plant Soil* **271**, 351–364 (2005).
- 817 134. Mencuccini, M. The ecological significance of long-distance water transport: Short-term regulation,
818 long-term acclimation and the hydraulic costs of stature across plant life forms. *Plant, Cell Environ.*
819 **26**, 163–182 (2003).
- 820 135. Cornwell, W. K. *et al.* Plant species traits are the predominant control on litter decomposition rates
821 within biomes worldwide. *Ecol. Lett.* **11**, 1065–1071 (2008).
- 822 136. Wei, T. & Simko, V. R package ‘corrplot’: Visualization of a Correlation Matrix. *R Packag. version*
823 *0.84* (2017).