



HAL
open science

Towards the assessment of biodiversity and management practices in mountain pastures using diagnostic species?

Corentin Nicod, Bérangère Leys, Yorick Ferrez, Vincent Manneville, Arnaud Mouly, Brendan Greffier, Christophe Hennequin, Yvette Bouton, Nicolas Chemidlin Prévost-Bouré, François Gillet

► To cite this version:

Corentin Nicod, Bérangère Leys, Yorick Ferrez, Vincent Manneville, Arnaud Mouly, et al.. Towards the assessment of biodiversity and management practices in mountain pastures using diagnostic species?. Ecological Indicators, 2019, 107, pp.1-9. 10.1016/j.ecolind.2019.105584 . hal-02270642

HAL Id: hal-02270642

<https://institut-agro-dijon.hal.science/hal-02270642v1>

Submitted on 20 Dec 2021

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution - NonCommercial 4.0 International License

1 Towards the assessment of biodiversity and management practices in mountain
2 pastures using diagnostic species?

3 Corentin Nicod ^{a,*}, Bérangère Leys ^a, Yorick Ferrez ^b, Vincent Manneville ^c, Arnaud Mouly ^a,
4 Brendan Greffier ^b, Christophe Hennequin ^b, Yvette Bouton ^d, Nicolas Chemidlin Prévost-
5 Bouré ^e, François Gillet ^a

6 ^a *Université Bourgogne Franche-Comté, UMR CNRS 6249 Chrono-Environnement, 16 route de Gray,*
7 *25030 Besançon, France*

8 ^b *Conservatoire Botanique National de Franche Comté – Observatoire Régional des Invertébrés, 7 rue*
9 *Voirin, 25000 Besançon, France*

10 ^c *Institut de l'Élevage, 9 allée Pierre de Fermat, 63170 Aubière, France*

11 ^d *Comité Interprofessionnel de Gestion du Comté, Unité R&D, Bât. INRA-URTAL 30 route de*
12 *Versailles, 39800 Poligny, France*

13 ^e *Université Bourgogne – Franche Comté, INRA, AgroSup Dijon, UMR 1347 Agroécologie, 17 rue de*
14 *Sully, 21000 Dijon, France*

15 * corresponding author

16 Email addresses: corentin.nicod@univ-fcomte.fr (C. Nicod), berangereleys@gmail.com (B. Leys),
17 yorick.ferrez@cbnfc.org (Y. Ferrez), Vincent.Manneville@idele.fr (V. Manneville), arnaud.mouly@univ-
18 fcomte.fr (A. Mouly), brendan.greffier@cbnfc.org (B. Greffier), christophe.hennequin@cbnfc.org (C.
19 Hennequin), y.bouton@comte.com (Y. Bouton), n.chemidlin@agrosupdijon.fr (N. Chemidlin Prévost-
20 Bouré), francois.gillet@univ-fcomte.fr (F. Gillet).

21 **Abstract**

22 Diversity of plant communities in semi-natural grasslands is strongly dependent on management
23 practices. To characterize community diversity and agricultural practices, we need easy-to-use
24 indicators. The diagnostic species are used to characterize and differentiate vegetation types, but are
25 they useful to assess the various facets of community diversity and the agricultural practices applied to
26 managed grasslands?

27 Based on the fidelity analysis of 311 phytosociological relevés, considered as references for mesic
28 grassland classification in the Franche-Comté region (eastern France), we identified eight diagnostic
29 species of mesic pastures (*Cynosurion cristati*). We then compared the number of diagnostic species
30 observed in 45 mountain pastures with taxonomic, phylogenetic and multi-trait functional diversity
31 indices, as well as with agronomic variables describing stocking rate and fertilization. To account for
32 the role of diagnostic species identity, we compared Spearman rank correlations obtained for these
33 diagnostic species with correlations calculated for 1000 random samples of eight species from the
34 regional species pool of mesic grasslands.

35 The number of diagnostic species in the grassland plot was significantly positively correlated with most
36 taxonomic, phylogenetic and functional diversity metrics. However, only the positive correlations with
37 Simpson taxonomic diversity and Rao functional diversity were significantly different from those
38 obtained from random samples. Moreover, the number of diagnostic species was significantly
39 negatively correlated with stocking rate and fertilization intensity, although only correlations with
40 industrial fertilizer input were significantly different from results obtained from random samples.

41 These results show that the number of diagnostic species observed in a pasture may be a good
42 indicator of a high taxonomic and functional diversity and of a low fertilization. Thereby, a special
43 attention should be given to diagnostic species to assess the status of grassland biodiversity, including
44 functional aspects.

45 **Key-words:** diagnostic species; grasslands; mountain pastures; biodiversity assessment;
46 management.

47 1. Introduction

48 Oligotrophic to mesotrophic semi-natural grasslands with a long history of management through
49 grazing or mowing are among the most species-rich ecosystems in the world and display very high
50 small-scale plant species richness (Cousins and Eriksson 2002; Wilson *et al.* 2012). Their high
51 biodiversity, resulting from a long evolution through extensive management (Mašková *et al.*, 2009;
52 Schermer *et al.*, 2016), provides high levels of many provisioning ecosystem services (Mauchamp *et al.*
53 *et al.*, 2013; Schirpke *et al.*, 2017) and support their stability. It is recognized that plant diversity reduces
54 inter-annual variability of biomass production (Hector *et al.*, 2010; Cadotte *et al.*, 2012), species
55 richness improves resistance of grassland productivity to climatic extreme events (Isbell *et al.*, 2015),
56 and the presence of subordinate species increases the adaptability of the grassland ecosystem to
57 climatic hazard (Mariotte *et al.*, 2013). Beyond conservation concerns, maintaining or enhancing
58 grassland biodiversity is therefore of first interest to promote resilience and maintain ecosystem
59 functions in a context of global change (Craine *et al.*, 2011; Oliver *et al.*, 2015). However, temperate
60 grasslands are among the most endangered ecosystems by land-use changes (Hoekstra *et al.*, 2005),
61 which are considered as one of the primary drivers of biodiversity loss (Foley *et al.*, 2005; Flynn *et al.*,
62 2009).

63 In Europe, significant changes in grassland management occurred in the last decades, aiming at
64 improving grassland productivity (Wesche *et al.* 2012). Two main opposite tendencies affect grassland
65 vegetation (Plantureux *et al.*, 2005; Buttler *et al.*, 2009): less productive parcels that are located far
66 away from the farms and not easily accessible are progressively neglected and tend to be afforested;
67 most productive parcels, with an easy access for machines, are intensified, often with an associating
68 increase in defoliation and nutrient inputs (Gaujour *et al.*, 2012). In relation to changes in grassland
69 management, recent studies recorded a decrease in plant diversity and a shift in species composition
70 over the last decades in different grassland types across European regions. In central Europe, an
71 important decrease of species richness and functional diversity was observed in resampled plots
72 compared to historical ones (Wesche *et al.*, 2012). In the French Jura Mountains, resampled plots
73 showed a lower phylogenetic and functional diversity and became dominated by nitrogen-demanding
74 competitive grasses (Gillet *et al.*, 2016). Similar studies in Scottish grasslands highlighted a loss of
75 functional richness (Pakeman *et al.*, 2017) and taxonomic diversity, associated with an increased
76 cover of nitrogen-demanding species (Mitchell *et al.*, 2017). Research in the Swiss Alps indicated a
77 recent homogenization of plant diversity at a regional scale (Homburger and Hofer 2012).

78 In response to the current loss of biodiversity in Europe, the Habitat Directive was adopted by the
79 European Union to protect the most seriously threatened habitats and to ensure their resilience to
80 global changes (OJEC, 1992; Gaston *et al.*, 2008; Louette *et al.*, 2015). Each EU member state is
81 required by the Habitat Directive to ensure the favorable conservation status of natural habitats
82 belonging to the Natura 2000 network through conservation measures (Ostermann *et al.*, 1998).
83 However, the Habitat Directive does not give a precise definition of the conservation status, nor
84 indicators to evaluate it (Kovač *et al.*, 2016). Consequently, although the concept of favorable
85 conservation status is still subject to intense discussion and research (Velázquez *et al.*, 2010; Louette

86 *et al.*, 2015), recent research was conducted to identify cost-effective indicators to evaluate it, but
87 mainly for forest habitats (Cantarello and Newton, 2008; Hernando *et al.*, 2010; Kovač *et al.*, 2016).

88 In this context, we stress the need to determine easy-to-use indicators in the field, allowing the
89 monitoring of plant diversity and agricultural practices in mountain pastures. Specifically, the large
90 traditionally pastoral region of the French Jura Mountains, devoted to dairy farming and the production
91 of protected designation of origin (PDO) cheese, was subject to a decline in grassland biodiversity
92 over the past decades due to management intensification (Gillet *et al.*, 2016). Thus, the PDO cheese
93 sector requires indicators to monitor the trends of biodiversity and management practices in
94 agricultural grasslands to be able to prevent further decline in grassland biodiversity. Indeed, in
95 addition to the provision and the maintenance of high levels of many ecosystem services, high
96 grassland biodiversity is one of the key components of the “terroir”, the basis of the specificity of PDO
97 products. In this study, we investigated the use of diagnostic species as indicators of biodiversity and
98 management practices for pastures grazed by dairy cows in the French Jura Mountains. As it is
99 generally possible to detect diagnostic species by non-destructive and inexpensive means, it could be
100 possible to use them to assess the conservation status of a local grassland habitat under the Habitat
101 Directive, provided they are good indicators of biodiversity and management practices of this habitat.

102 Diagnostic species can be defined as species that show a preference for a given habitat or biotic
103 community type (De Cáceres *et al.* 2008), which are commonly used in field survey for the
104 identification of community types. It is acknowledged that diagnostic species are taxa that indicate
105 habitat conditions where they are found because they are particularly responsive to habitat
106 modifications (Chiarucci *et al.*, 2008; Santoro *et al.*, 2012). In pastures, the intensification of
107 management practices leads to a decrease of taxonomic, functional and phylogenetic diversity
108 (Carmona *et al.*, 2012; Mauchamp *et al.*, 2014; Dainese *et al.*, 2015). As the condition of the habitat is
109 a limiting factor controlling the population of specialist species (Evans and Arvela, 2011), diagnostic
110 species are supposed to be among the first species to disappear in response to the intensification of
111 management practices. Thus, it may be possible to use diagnostic species to assess the trends of
112 biodiversity and management practices in pastures. Indeed, the disappearance of diagnostic species
113 should indicate a change in management practices towards an intensification in association with a
114 decrease in plant diversity. Based on the hypothesis that species restricted to one or a few habitats
115 are potentially better indicators of environmental change than generalist species (De Cáceres and
116 Legendre, 2009), we addressed the following questions:

- 117 1) Are diagnostic species good indicators of biodiversity and management practices of the
118 vegetation type they characterize?
- 119 2) Are diagnostic species better indicators of biodiversity and management practices than other
120 grassland species?

121 2. Material and methods

122 2.1. Study area and vegetation sampling

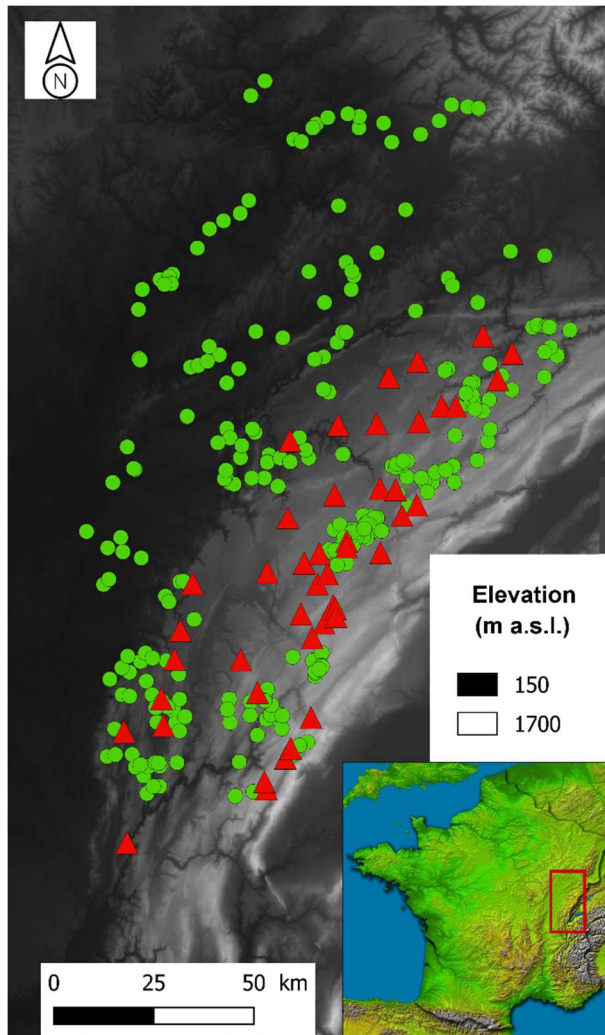
123 Our study was carried out in the French Jura Mountains. This mountain range is composed of three
124 main structural units across an elevation gradient: first plateau (500-800 m a.s.l.), second plateau
125 (800-950 m a.s.l.) and high range (950-1700 m a.s.l.). Climate is suboceanic (fully humid, warm
126 summer), ranging from warm temperate (Cfb) to boreal (Dfb, at the highest elevations) according to
127 Köppen-Geiger climate classification (Kottek *et al.*, 2006; Rubel *et al.*, 2017). Predominant soils are
128 cambisols developed on limestone with a variable superficial cover of silt. In the French Jura
129 Mountains, permanent grasslands represent 22% of the area and support dairy farming and PDO
130 cheese production, mainly Comté. Such production implies constraining specifications for agricultural

131 practices; e.g. in the PDO Comté area, the average nutrient inputs must not exceed 120 kg N ha⁻¹ yr⁻¹
132 and the stocking rate is limited to 1.3 livestock units per hectare (one livestock or adult bovine unit LU
133 is equivalent to one dairy cow).

134 In Ferrez *et al.* (2007), 525 phytosociological relevés, realized between 1973 and 2007 in eastern
135 France, were analyzed to elaborate a typology for mesic grassland classification. To obtain
136 homogeneous groups of relevés for the typology, the authors excluded the phytosociological relevés
137 not well discriminated by cluster analysis, thus retaining 371 relevés. In our study, we selected 311
138 among the 371 phytosociological relevés presented in Ferrez *et al.* (2007) to compute a list of
139 diagnostic species of mesic pastures (*Cynosurion cristati* Tüxen 1947). When selecting these relevés,
140 we considered only those belonging to phytosociological associations present in the French Jura
141 Mountains, which explains the exclusion of 60 phytosociological relevés.

142 45 permanent grasslands were then chosen in the French Jura Mountains across an elevation
143 gradient, from which both vegetation relevés and detailed agricultural data were available (Fig. 1).
144 They cover a wide range of agricultural practices but are all primarily used as pastures for dairy cows.
145 According to the regional phytosociological classification (Ferrez *et al.*, 2011), the plant communities in
146 these 45 grasslands belong to the *Cynosurion cristati* Tüxen 1947 (mesic pastures).

147 It has been shown that, for a same plot size, sampling units with more elongated shapes present
148 higher species richness than squares or circles (Heegaard *et al.*, 2007; Güler *et al.*, 2016). Thus, to
149 account for a maximum of plant species, the vegetation of each grassland was sampled from May
150 2017 to July 2017 in a rectangular plot of 120 m² (60 m × 2 m), on the flattest area located inside the
151 grassland and far from parcel margin. The choice of a plot size close to 100 m² is common in
152 phytosociology (Chytrý and Otýpková, 2003) and is consequently widely applied to permanent
153 grasslands (Hejcman *et al.*, 2010; Homburger and Hofer 2012; Gillet *et al.*, 2016), because it is more
154 efficient than a square plot of 16 m² to account for most of the species present. All observed vascular
155 plant species were listed and the cover of each species was estimated using the seven degrees of the
156 dominance Braun-Blanquet scale (r, +, 1, 2, 3, 4, 5). These codes were converted into absolute
157 percentage cover (van der Maarel, 1979) and then adjusted to relative percentage cover by summing
158 to 100% for each plot. We considered relative cover because its estimation is more reliable than the
159 one of absolute cover, and because only relative dominance was considered for the computation of
160 diversity metrics.



161

162 Fig. 1: Location of the 311 phytosociological relevés used to compute the diagnostic species list of
 163 mesic pastures (green circles) and of the 45 grassland plots sampled in 2017 (red triangles).

164 **2.2. Establishment of the list of diagnostic species**

165 The geographical area, the number of phytosociological relevés and the vegetation units taken into
 166 account strongly influence the selection of diagnostic species (Chytrý *et al.*, 2002; Tichý and Chytrý,
 167 2006; De Cáceres and Legendre, 2009). To determine diagnostic species of a given vegetation unit,
 168 the most appropriate dataset should be selected in relation with the objectives of the study, as there is
 169 no better way to proceed (De Cáceres and Legendre, 2009). As this study focused on mesic pastures
 170 in the French Jura Mountains, we used only phytosociological relevés of mesic grasslands realized in
 171 the study area. This choice is supported by the fact that extracting a common list of diagnostic species
 172 of larger geographical area is difficult due to the absence of shared species in remote areas (De
 173 Cáceres *et al.*, 2015).

174 The 311 selected phytosociological relevés were assigned to four vegetation units (alliances)
 175 according to the mesic grassland classification of the Franche-Comté region proposed in Ferrez *et al.*
 176 (2007): 29% (88/311) belonged to the *Arrhenatherion elatioris* W. Koch 1926 (lowland mesic hay
 177 meadows), 35% (108/311) to the *Trisetum flavescens-Polygonum bistortae* Braun-Blanquet & Tüxen ex
 178 Marschall 1947 (mountain mesic hay meadows), 33% (105/311) to the *Cynosurion cristati* Tüxen 1947
 179 (mesic pastures), and 3% (10/311) to the *Lolium perennis-Plantaginion majoris* G. Sissingh 1969 (mesic
 180 trampled grasslands).

181 To compute a diagnostic species list of mesic pastures, we used the phi coefficient of association
182 (Chytrý *et al.*, 2002; De Cáceres and Legendre, 2009), a measure of species fidelity to vegetation
183 types. Applied to species cover data, this index is the Pearson correlation computed between a
184 quantitative vector (*i.e.*, the vector containing the species cover values in the various phytosociological
185 relevés) and a binary vector (*i.e.*, “1” for phytosociological relevés belonging to the alliance of interest
186 and “0” for phytosociological relevés belonging to other alliances). The calculation was performed
187 using the `multipatt()` function from the `indicspecies` package (De Cáceres and Legendre, 2009) in R
188 3.4.2 (<http://www.R-project.org>), setting 5000 permutations. Because differences in the size of the
189 target vegetation units are known to influence the determination of diagnostic species (Tichý and
190 Chytrý, 2006) and the variation in cover of different species can be an important source of
191 heterogeneity among vegetation units (Willner *et al.*, 2009), we used a group-equalized phi coefficient
192 of association and considered species cover values instead of presence-absence data (see De
193 Cáceres and Legendre, 2009). We retained only the species that showed significant fidelity ($P < 0.05$)
194 for 10 runs of the `multipatt()` function, to avoid selecting species close to the threshold of significance.

195 2.3. Diversity metrics

196 The taxonomic, functional and phylogenetic facets of community diversity were acknowledged to be
197 complementary aspects of community structure, responding differently to environmental changes and
198 not always positively correlated (Cadotte *et al.*, 2013; Pavoine *et al.*, 2013). Thus, to measure
199 community diversity, we computed taxonomic, functional and phylogenetic alpha diversities at the plot
200 level (Devictor *et al.*, 2010; Cadotte *et al.*, 2013; Dainese *et al.*, 2015). Taxonomic diversity was
201 measured through species richness (N0), inverse Simpson diversity (N2) based on Rényi general
202 entropy (Hill, 1973), and Pielou evenness (J1) according to Jost (2010). Functional and phylogenetic
203 facets of diversity were computed using the Rao quadratic diversity with Jost correction, as it
204 corresponds to a generalization of the inverse Simpson index (Jost, 2007; de Bello *et al.*, 2010).

205 Multi-trait Rao quadratic diversity was computed from a standardized Euclidean distance matrix using
206 four selected functional traits, weighted by the relative cover of individual plant species: leaf dry matter
207 content (LDMC), leaf nitrogen content per dry mass (LNC), specific leaf area (SLA) and maximum
208 plant height (MH). These traits are widely used in functional ecology of grasslands (Louault *et al.*,
209 2005; Ansquer *et al.*, 2009; Garnier and Navas, 2012; Lososová *et al.*, 2016; Mauchamp *et al.*, 2016).
210 Maximum plant height is an important morphological factor determining plant competitive ability
211 (Tilman, 1988; Westoby *et al.*, 2002). LDMC, LNC and SLA are functional traits linked to resource
212 acquisition and growing strategy (Garnier *et al.*, 2004). These traits were extracted from two
213 databases: LEDA Traitbase for LDMC and SLA (Kleyer *et al.*, 2008) and TRY database for LNC and
214 MH (Kattge *et al.*, 2011). Species with missing information were removed, so that multi-trait Rao
215 quadratic diversity was sometimes computed with a bit less than 90% of the relative percentage cover.
216 All computations were performed using R v 3.4.2 (<http://www.R-project.org>), `vegan` package (Oksanen
217 *et al.*, 2018), and `Rao()` function (de Bello *et al.*, 2010).

218 To compute phylogenetic diversity, we constructed a phylogenetic tree composed of the vascular plant
219 species recorded during the field work. The phylogenetic tree was built following the same procedure
220 described in different studies conducted in the French Jura Mountains (Perronne *et al.*, 2014;
221 Mauchamp *et al.*, 2014 and Mauchamp *et al.*, 2016). The sequences of two genes encoding
222 chloroplast proteins (*rbcL* and *matK*) were obtained using the GenBank database (Benson *et al.*,
223 2006). Their complementarity allows a good resolution of the tree for the recorded species (Hilu *et al.*,
224 2003). We replaced species not yet sequenced for the selected genes by their closest available
225 relative, according to the APG III classification (APG 2009). All sequences were first aligned using
226 ClustalX 2.1 software (Larkin *et al.*, 2007) and then adjusted manually for both genes. A combined

227 analysis of these two datasets was then performed following the Bayesian Monte Carlo Markov Chain
228 (MCMC) approach under BEAST 1.8.4 software (Drummond and Rambaut, 2007). We used
229 MrModeltest2 software (v. 2.0, Evolutionary Biology Center, Uppsala University, SE) to choose the
230 model of nucleotides substitution that best fits the data according to Akaike's information criterion. The
231 general time reversible model (GTR) with among-site substitution rate heterogeneity described by a
232 gamma distribution (Yang, 1994) was then selected. To calibrate the rates of molecular evolution of
233 each lineage for angiosperms clades (Bell *et al.*, 2005), several family relationships were constrained
234 in BEAST following the APG III phylogeny (APG 2009). An uncorrelated relaxed molecular clock that
235 takes into account evolutionary heterogeneity between lineages was used for branch length
236 calculation (Drummond *et al.*, 2006). Finally, the ultrametric tree topology was saved in a newick
237 format. We used the Rao quadratic entropy, weighted by the ultrametric distances of the phylogenetic
238 tree, to compute phylogenetic diversity (RaoPD), utilizing Rao() function (de Bello *et al.*, 2010).

239 2.4. Management practices

240 For each of the 45 selected pastures, farmers provided detailed information about grazing
241 management and fertilization level. For the grazing management, the obtained information was the
242 number of dairy cows or livestock units (LU) grazing the parcel, their time of presence at each rotation
243 and the time elapsed between two rotations in 2017. For the fertilization practices, the collected data
244 were the type, the quantity and the spreading date of the different fertilizers applied (farmyard
245 manures and industrial fertilizers). Moreover, general characteristics of the farm, building type, herd
246 performance, manure and slurry production, and spreading management of fertilizers were collected at
247 the farm level. Based on this survey, we computed different agronomic variables.

248 The stocking rate in 2017, traducing the animal pressure on the pasture and expressed in LU day ha⁻¹,
249 was calculated by multiplying the number of dairy cows with the time they spent on the parcel, divided
250 by the parcel area. To account for variations in grazing intensity along the year, we calculated the
251 stocking rate in spring, summer and autumn separately. The studied pastures were located close to
252 the farm buildings and aimed to be grazed by dairy cows according to a same rotation plan applied by
253 the farmers each year. Since the dairy cows herd was not expected to change significantly in recent
254 years, the values calculated for 2017 were assumed to be similar to those of previous years.

255 The data given by farmers relative to the N input from farmyard manures were sometimes imprecise.
256 Thus, we decided to estimate the total manure amount produced in the farm per year F_{man} . To do so,
257 we used the following formula:

$$258 F_{man} = \frac{T}{12} N Q$$

259 T is the time spent by the cattle inside the cowshed, expressed in month. N is the number of dairy
260 cows constituting the herd (LU) and Q is the farmyard manure amount produced by a dairy cow
261 present 12 months in a cowshed (expressed in m³ LU⁻¹ yr⁻¹ or t LU⁻¹ yr⁻¹). The Q value depends on the
262 type of building and the kind of produced manure. According to the Institut de l'Elevage (2012), the
263 retained values were 13.5 t LU⁻¹ yr⁻¹ for solid manure production and 23 m³ LU⁻¹ yr⁻¹ for liquid manure
264 production. The calculated total manure amount produced in the farm per year F_{man} was then divided
265 by the total surface of spreading in order to obtain an estimate per hectare and per year. This quantity
266 was multiplied by the mean nitrogen content of each manure type: 5 kg N t⁻¹ for solid manure, 5 kg N
267 m⁻³ for liquid manure and 3 kg N m⁻³ for diluted liquid manure (Institut de l'Elevage, 2012). The given
268 result was the estimated N input spread on the parcel from farmyard manure, expressed in kg N ha⁻¹
269 yr⁻¹. As the organic part of the nitrogen present in farmyard manure is not directly available for plants
270 because it requires a post-spreading mineralization by soil microorganisms (Kirkham *et al.*, 2008;

271 Gaujour *et al.*, 2012), a last calculation was done to consider only the N input available for plant
272 growth during the field work in 2017. Its aim was to exclude the part of the N input not yet mineralized.
273 To do so, we multiplied the N input by the proportion of available nitrogen depending of the time
274 elapsed from the manure spread to the vegetation sampling period. The utilized values were 0.5 for
275 liquid manure spread during the sampling year, 0.2 for liquid manure spread one year before the field
276 work, 0 for liquid manure spread two years before the field work, 0.15 for solid manure spread during
277 the sampling year, 0.1 for solid manure spread one year before the field work and 0.05 for solid
278 manure spread two years before the field work (Institut de l'Élevage, 2012). Thereby, the calculated
279 available N input from farmyard manure (N_{man}), expressed in kg N ha⁻¹ yr⁻¹, relied on the fertilization
280 practices of the past three years.

281 As nutrients present in industrial fertilizers are directly available for plants (Cavalli *et al.*, 2014), we
282 considered the values given by farmer, expressed in kg N ha⁻¹ yr⁻¹, as the available N input on the
283 parcel from industrial fertilizers (N_{ind}).

284 2.5. Statistical analyses

285 We first performed a principal component analysis (PCA) on the Hellinger-transformed species-cover
286 matrix of the 45 grassland plots (Borcard *et al.*, 2018). To get an overview of the relationships between
287 all computed variables and the number of observed diagnostic species, we fitted these variables on
288 the PCA plot using the `envfit()` function from the `vegan` R package (Oksanen *et al.*, 2018). To address
289 the question of whether diagnostic species are reliable biodiversity indicators in mountain pastures, we
290 tested Spearman rank correlation between all diversity metrics and the number of diagnostic species
291 observed in each plot. We also tested Spearman rank correlation between agronomic variables and
292 the number of diagnostic species to address of whether diagnostic species are good management
293 indicators or not. It is a major difference, especially in field surveys, whether a species is diagnostic
294 *per se* or only when it exceeds a certain cover value (Willner *et al.*, 2009). Here, we decided to
295 consider all diagnostic species present in the grassland plots regardless of their cover. Indeed,
296 defining a cover threshold to take into account or not a diagnostic species would have been subjective
297 and different for each species, as some species naturally show low or high cover values.

298 To address the question of whether the selected diagnostic species are better indicators compared to
299 any other grassland species, we generated 1000 samples of the same number of randomly selected
300 species. The species taken into account to do so were all the species that occurred in more than 5%
301 of the 311 phytosociological relevés describing mesic grasslands in the study area, so that casual taxa
302 were avoided. We then compared Spearman rank correlation calculated using the diagnostic species
303 with correlations calculated using the randomly selected species samples, for all diversity and
304 agronomic variables. We set the correlation value obtained for the diagnostic species as a reference
305 and computed a non-parametric *P*-value based on the number of greater and lower correlations
306 obtained for the 1000 random samples:

$$307 \quad P = 1 - \frac{|A - B|}{1000}$$

308 *A* is the number of higher correlation values and *B* the number of lower correlation values. All analyses
309 were performed using R v 3.4.2 (<http://www.R-project.org>).

310 3. Results

311 3.1. Diagnostic species

312 The set of diagnostic species was composed of eight species (Table 1): *Cynosurus cristatus* L.,
 313 *Prunella vulgaris* L., *Achillea millefolium* L., *Agrostis capillaris* L., *Plantago media* L., *Veronica*
 314 *serpyllifolia* L., *Scorzoneroides autumnalis* (L.) Moench, and *Pimpinella saxifraga* L. The phi values
 315 ranged from 0.241 to 0.422 (after 5000 permutations). Six of these species were frequent in the 45
 316 grassland plots whereas *Scorzoneroides autumnalis* (L.) Moench was present at low frequency and
 317 *Pimpinella saxifraga* L. was absent.

318 A significant positive phi coefficient indicated that the cover of the species in phytosociological relevés
 319 belonging to the *Cynosurion cristati* Tüxen 1947 (mesic pastures) was higher than its cover in
 320 phytosociological relevés not belonging to it. Larger values (close to 1) indicated a greater degree of
 321 joint fidelity between the species and the group of relevés belonging to the *Cynosurion cristati* Tüxen
 322 1947. As the phi values were rather low (close to 0) for the eight diagnostic species, it indicated a low
 323 degree of joint fidelity.

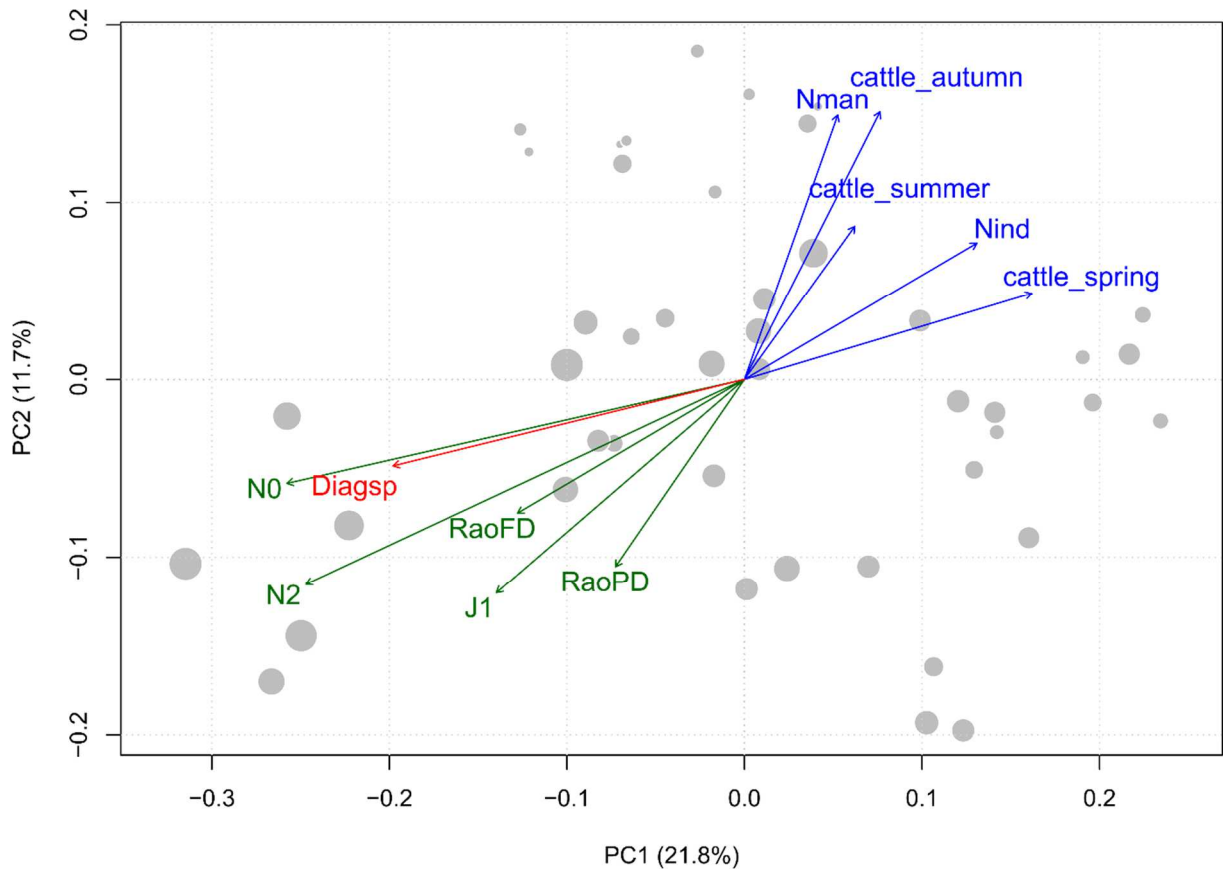
324 Table 1. Degree of joint fidelity of the eight diagnostic species of mesic pastures (*Cynosurion cristati*
 325 Tüxen 1947), obtained after 5000 permutations on the 311 phytosociological relevés. The last column
 326 presents the frequency of these diagnostic species in the 45 grassland plots.

Species name	phi	P	Frequency
<i>Cynosurus cristatus</i> L.	0.422	< 0.001 ***	0.60
<i>Prunella vulgaris</i> L.	0.383	0.002 **	0.58
<i>Achillea millefolium</i> L.	0.324	0.007 **	0.78
<i>Agrostis capillaris</i> L.	0.324	0.007 **	0.76
<i>Plantago media</i> L.	0.298	0.007 **	0.44
<i>Veronica serpyllifolia</i> L.	0.268	0.022 *	0.62
<i>Scorzoneroides autumnalis</i> (L.) Moench	0.261	0.016 *	0.09
<i>Pimpinella saxifraga</i> L.	0.241	0.033 *	0.00

327

328 3.2. Relationships between diversity, management and number of diagnostic species

329 The fitted variables on the PCA first axes showed positive correlations between all agronomic
 330 variables, as well as positive correlations between all diversity metrics (Fig. 2). Axes 1 and 2
 331 represented 33.5% of the variance of the Hellinger-transformed species matrix. The number of
 332 diagnostic species observed in the grassland plot (Diagsp) was positively correlated with all displayed
 333 diversity metrics and negatively correlated with all displayed agronomic variables. This means that
 334 when a high number of diagnostic species was found in a given grassland, it indicated a high species
 335 richness (N0), inverse Simpson diversity (N2), Pielou evenness (J1), functional diversity (RaoFD) and
 336 phylogenetic diversity (RaoPD). At the opposite, when a low number of diagnostic species was found
 337 in a given grassland, it indicated a high fertilization intensity through the use of farmyard manure
 338 (Nman) and industrial fertilizers (Nind), and a high stocking rate during spring (cattle_spring), autumn
 339 (cattle_autumn), and summer in a lesser extent (cattle_summer).



340

341

342 Fig. 2: Principal component analysis of the Hellinger-transformed species composition of the 45
 343 grasslands plots, with fitted diversity metrics (green arrows), agronomic variables (blue arrows) and
 344 number of diagnostic species observed in the grassland plot (Diagsp, red arrow). Diversity metrics: N0
 345 species Richness, N2 inverse Simpson Diversity, J1 Pielou evenness, RaoPD Rao phylogenetic
 346 diversity, RaoFD Rao functional diversity. Agronomic variables: cattle_spring stocking rate during
 347 spring, cattle_summer stocking rate during summer, cattle_autumn stocking rate during autumn,
 348 Nman available N input from farmyard manures, Nind available N input from industrial fertilizers. Sites
 349 are represented by grey point whose size is proportional to elevation. Species are not shown.

350 These results were confirmed by Spearman rank correlations (Table 2). Indeed, we found significant
 351 positive correlations between the number of observed diagnostic species in the grassland plot and all
 352 diversity metrics (N0, N2 and J1 for taxonomic diversity, RaoPD for phylogenetic diversity and RaoFD
 353 for functional diversity). Significant negative correlations were also found between the number of
 354 observed diagnostic species and almost all agronomic variables (cattle_spring and cattle_autumn for
 355 stocking rate, and Nman and Nind for fertilization). Only the negative correlation with stocking rate
 356 during summer (cattle_summer) was not significant. Thereby, the number of diagnostic species in a
 357 pasture revealed to be a good indicator of high plant diversity and extensive management practices
 358 (low stocking rate and low fertilization).

359 Table 2: Spearman rank correlations between diversity metrics or agronomic variables and the number
 360 of diagnostic species observed in the grassland plot.

		<i>rho</i>	<i>P</i>
Diversity metrics			
N0	Species richness	0.741	< 0.001 ***

N2	Inverse Simpson diversity	0.689	< 0.001 ***
J1	Pielou evenness	0.302	0.044 *
RaoPD	Rao phylogenetic diversity	0.395	0.007 **
RaoFD	Rao functional diversity	0.540	< 0.001 ***
Agronomic variables			
cattle_spring	stocking rate (LU·day·ha ⁻¹ during spring)	- 0.521	< 0.001 ***
cattle_summer	stocking rate (LU·day·ha ⁻¹ during summer)	- 0.168	0.269
cattle_autumn	stocking rate (LU·day·ha ⁻¹ during autumn)	- 0.371	0.012 *
Nman	Available N on the parcel from farmyard manures (kg N·ha ⁻¹ ·yr ⁻¹)	- 0.328	0.028 **
Nind	Available N on the parcel from industrial fertilizers (kg N·ha ⁻¹ ·yr ⁻¹)	- 0.599	< 0.001 ***

361

362 3.3. Diagnostic species in comparison to randomly selected species

363 The comparison of the Spearman rank correlations calculated using the diagnostic species list with
364 correlations calculated using the 1000 randomly selected species lists (Table 3) revealed that
365 diagnostic species were better indicators than other grasslands species for inverse Simpson diversity
366 (N2), Rao functional diversity (RaoFD), and the available N input from industrial fertilizers (Nind).
367 Although results obtained for stocking rate during spring (cattle_spring) showed $P = 0.058$, the eight
368 diagnostic species were not better indicators than randomly selected species for the other variables. In
369 particular, species richness (N0), which presented a high correlation with the number of diagnostic
370 species ($\rho = 0.741$), was also highly correlated to the number of species issued from random
371 samples.

372 Table 3: Number of greater and lower Spearman rank correlations between diversity metrics or
373 agronomic variables and the number of species in 1000 random samples, as compared to the
374 correlations calculated using the set of eight diagnostic species.

	diagnostic species	1000 random samples		<i>P</i>
	<i>rho</i>	<i>Number of greater rho</i>	<i>Number of lower rho</i>	
Diversity metrics				
N0	0.741	105	895	0.210
N2	0.689	15	985	0.030 *
J1	0.302	133	867	0.266
RaoPD	0.395	58	942	0.116
RaoFD	0.540	10	990	0.020 *
Agronomic variables				
cattle_spring	- 0.521	973	27	0.054
cattle_summer	- 0.168	695	305	0.610
cattle_autumn	- 0.371	849	151	0.302
Nman	- 0.328	911	89	0.178
Nind	- 0.599	993	7	0.014 *

375

376 4. Discussion

377 4.1. Establishment of the list of diagnostic species

378 In this study, we identified a set of eight diagnostic species of mesic pastures (*Cynosurion cristati*
379 Tüxen 1947) based on 311 phytosociological relevés realized in eastern France. Our results revealed
380 that the degree of joint fidelity between each diagnostic species and the group of relevés belonging to
381 the alliance of mesic pastures was low. Indeed, the diagnostic species of pastures were sometimes
382 found with low cover in relevés belonging to other alliances of mesic grasslands (Ferrez *et al.*, 2007).
383 These results were not surprising, regarding the fact that it is unattainable to find species with absolute
384 fidelity to a given vegetation type (Willner *et al.*, 2009). Moreover, we considered only the class of
385 mesic grasslands (*Arrhenatheretea elatioris* Braun-Blanquet 1949) for the establishment of the list of
386 diagnostic species. Thus, we ignored the affinity of our diagnostic species for vegetation units
387 belonging to other classes, such as the dry grasslands (*Festuco valesiacaе-Brometea erecti* Braun-
388 Blanquet & Tüxen ex Braun-Blanquet 1949). Consequently, the diagnostic species *Pimpinella*
389 *saxifraga*, commonly found in dry grasslands, was not present in the 45 grassland plots that we
390 analysed. In the study area, this species is present in lowland pastures whereas it occurs only in dry
391 grasslands and fringes in mountain areas.

392 4.2. Diagnostic species as biodiversity and management indicators

393 The number of diagnostic species present in a grassland revealed to be a good indicator of
394 biodiversity and management practices, as almost all tested variables showed significant correlations
395 with it. The combined presence of several diagnostic species of the *Cynosurion cristati* Tüxen 1947
396 (mesic pastures) in a grassland belonging to this vegetation type was associated with a high plant
397 taxonomic, functional and phylogenetic diversity and an extensive management (low stocking rate and
398 low fertilization). These findings were predictable as an increase in the number of plant species is
399 often associated with an increase of plant diversity (Cadotte *et al.*, 2011; Pavoine and Bonsall, 2011),
400 and an extensive management in grassland is necessary to maintain plant diversity (Mašková *et al.*,
401 2009; Wilson *et al.*, 2012; Mauchamp *et al.*, 2014).

402 The nitrogen input from industrial fertilizers showed to be significantly negatively correlated with the
403 number of observed diagnostic species. Moreover, the correlation obtained for the diagnostic species
404 was higher than correlations obtained for randomly selected species. These results have to be
405 considered regarding the fact that industrial fertilizers are the most detrimental source of nutrients for
406 plant diversity in mountain grasslands (Mauchamp *et al.*, 2016), because nutrients are directly
407 available for plants (Cavalli *et al.*, 2014). Diagnostic species are able to react to modification of the
408 environmental conditions of their habitat (Bazzichetto *et al.*, 2016). A study in coastal dunes
409 ecosystems showed that diagnostic species can be indicators of soil conditions and provide useful
410 indications for habitat monitoring, because they have narrow edaphic requirement (Angiolini *et al.*,
411 2018). These previous results are similar to our observation concerning the nitrogen input from
412 industrial fertilizers. Our results support the idea that diagnostic species are the first species to
413 disappear when the nitrogen input increases through the use of industrial fertilizers, because they
414 have narrow requirement concerning the nitrogen supply (Online Appendix A). In comparison to
415 industrial fertilizers, the impact of the application of farmyard manures on mountain grassland plant
416 diversity is less pronounced (Mauchamp *et al.*, 2016). Indeed, the organic forms of the nitrogen
417 present in farmyard manure are not directly available for plants and requires a post-spreading
418 mineralization by soil microorganisms (Kirkham *et al.*, 2008; Gaujour *et al.*, 2012). The fact that the
419 impact of this type of fertilizer on plant diversity is less important compared to industrial fertilizers could
420 explain why the response of diagnostic species was not different from the response of other grassland
421 species. The number of diagnostic species present in a grassland showed to be negatively correlated
422 with the stocking rate, particularly with the stocking rate in spring. This can be explained by the fact
423 that an important stocking rate in spring is the most detrimental for plant diversity (Critchley *et al.*,
424 2007). However, the response of the eight diagnostic species to the stocking rate was not different

425 from the response of randomly selected species. Diagnostic species of pastures are species that are
426 well adapted to the disturbances induced by cattle activity. Thus, it is likely that diagnostic species of
427 pastures do not react particularly to variations in stocking rate in comparison to other species.

428 The significant positive correlation obtained between the number of observed diagnostic species and
429 species richness is rather tautological, as the number of diagnostic species is not independent from
430 species richness. This also explains why the correlation obtained using the set of eight diagnostic
431 species was not different from correlations obtained using randomly selected species. Indeed,
432 relationships between these two indices are positive regardless of the identity of the concerned
433 species. Although Pielou evenness showed significant positive correlation with the number of
434 diagnostic species, indicating that cover differences between species in the grassland plot were
435 smaller when a high number of diagnostic species was observed, diagnostic species were not better
436 indicators than randomly selected species for this diversity metric. However, in comparison to
437 randomly selected species, diagnostic species were more reliable to assess inverse Simpson index.
438 The relationships between these taxonomic diversity metrics and the number of observed diagnostic
439 species could be related to the nitrogen input from industrial fertilizers. Indeed, in the French Jura
440 Mountains, Mauchamp *et al.* (2016) recorded that the nitrogen input from industrial fertilizer had a
441 negative impact on inverse Simpson index, but not on species evenness. As diagnostic species
442 revealed to be better indicators than randomly selected species to assess the nitrogen input from
443 industrial fertilizers, they were also indirectly better indicators to assess inverse Simpson index.

444 According to our results, diagnostic species were better indicators than randomly selected species for
445 functional diversity. This can be explained by the fact that the range of the trait values of the diagnostic
446 species matched the range of the trait values of the species pool of the 45 grassland plots, at least for
447 SLA and LDMC (Online Appendix B). Functional traits are known to be particularly responsive to
448 environmental changes (Díaz *et al.*, 2013). Therefore, in relation to a change in environmental
449 conditions, a loss of species with particular values for SLA or LDMC in the plant community is likely to
450 be followed by a loss of the diagnostic species with the same traits values. Thus, the combined
451 presence of many diagnostic species in a given grassland indicated a wide range of the trait values for
452 SLA and LDMC, consequently a high functional diversity.

453 Our list of diagnostic species was constituted of species from different lineages. Thus, the combined
454 presence of many diagnostic species revealed that different species lineages were present in the plant
455 community, which can explain the significant positive correlation obtained between the number of
456 observed diagnostic species and phylogenetic diversity. Moreover, according to Villalobos and Vamosi
457 (2016), the alteration in grassland plant community composition increases the prevalence of
458 zygomorphic flower species lineages. Thus, a plant indicator list equilibrated in actinomorphic and
459 zygomorphic lineages like our list of diagnostic species, consequently representative of a large
460 phylogenetic diversity, reinforces the potential for the assessment of phylogenetic diversity. However,
461 results obtained for phylogenetic diversity were not different between diagnostic species and randomly
462 selected species. Indeed, it is likely that many random species subsets were also equilibrated in
463 species from different lineages.

464 4.3. Future direction for the use of diagnostic species in European grasslands

465 In this study, we have shown that we can use diagnostic species of the *Cynosurion cristati* to assess
466 biodiversity and management practices in mesic pastures of the French Jura Mountains. Given the
467 fact that it is easy to count for a number of diagnostic species in a local grassland, the use of
468 diagnostic species to assess biodiversity and management practices in agricultural grasslands should
469 be further investigated. In Europe, permanent grasslands are ecosystems whose floristic composition
470 is strongly dependent on the agricultural practices used for their management (Králóvec *et al.*, 2009).

471 Moreover, permanent grasslands belonging to the alliance of the *Cynosurion cristati* Tüxen 1947 are
472 widely represented across temperate Europe at low and mid-elevations (Mucina *et al.*, 2016).
473 Consequently, we believe that the use of diagnostic species as biodiversity and management
474 practices indicators in pastures of the French Jura Mountains belonging to the *Cynosurion cristati*
475 Tüxen 1947 can be applied to other European grasslands belonging to this vegetation type. More
476 globally, we believe that the methods that we used in the French Jura Mountains can be extended to
477 other alliances of anthropogenic managed grasslands in Europe, because their high biodiversity is the
478 result of a relatively similar long evolution through extensive management (Mašková *et al.*, 2009). In
479 addition, the results of a recent study conducted in the Jura Mountains support the potential use of
480 diagnostic species as biodiversity and management practices indicators in other habitat types. Indeed,
481 in relation to land-use abandonment and nitrogen deposition in wetlands, Rion *et al.* (2018) reported a
482 decrease of taxonomic diversity associated with a decrease in frequency and cover of diagnostic
483 species. However, the fact that we only investigated the use of diagnostic species as biodiversity and
484 management practices indicators in pastures for dairy cows have to be considered regarding the
485 possible generalization of the results. Moreover, our results seem difficult to generalize to grassland
486 ecosystems that are not managed, such as alpine grasslands, continental steppes or tallgrass prairie.

487 Our results support the idea that the concept of diagnostic species matches the concept of typical
488 species mentioned, but not defined, in the Habitat Directive (OJEC, 1992). According to Evans and
489 Arvela (2011), typical species should be species only found in a habitat and sensitive to changes in
490 the condition of this habitat. They should also be selected to reflect favorable structure and functions
491 of the habitat. In our study, we showed that diagnostic species were good indicator of management
492 practices in pastures of the Jura Mountains, especially fertilization. Our results also revealed that the
493 combined presence of many diagnostic species in a grassland habitat was associated with a high
494 plant diversity, mainly taxonomic and functional diversity. Thereby, we argue that the number of
495 diagnostic species in a given grassland could be used as one of the indicators of the favorable
496 conservation status of the local grassland habitat. This statement is supported by a study realized in
497 Mediterranean coastal dunes, which highlighted that diagnostic species can provide helpful indications
498 for conservation, monitoring and restoration of these habitats (Angiolini *et al.*, 2018). However, despite
499 the fact they can present a high plant diversity (Mauchamp *et al.*, 2014), pastures belonging to the
500 *Cynosurion cristati* Tüxen 1947 are not integrated in the habitat directive (OJEC, 1992). In order to use
501 the number of diagnostic species in the evaluation of the conservation status of a local grassland
502 habitat, future research have to be conducted, focusing on the assessment of biodiversity using
503 diagnostic species in grassland types integrated in the Natura 2000 network, such as mesic hayfields,
504 dry grasslands or wet grasslands.

505 5. Conclusion

506 This study provides a first insight about the use of diagnostic species as biodiversity and management
507 indicators. In the study area, the number of diagnostic species in a pasture revealed to be a good
508 indicator of taxonomic and functional diversity of the plant community and of management practices,
509 especially fertilization. As it is easy to count a number of diagnostic species in a grassland plot, a
510 special attention should be given to diagnostic species to assess the status of grassland biodiversity,
511 including functional aspects. However, future research is needed to show if the results of this study
512 can be extended to others grassland types such as hayfields, dry grasslands or wet grasslands, and to
513 other geographical areas.

514 Acknowledgment

515 This research was supported by the Comité Interprofessionnel de Gestion du Comté (CIGC) and the I-
516 SITE Bourgogne Franche-Comté program and took place in the long-term ecological research site
517 Jurassian Arc (<http://zaaj.univ-fcomte.fr/?lang=en>). We thank the farmers who gave their time to
518 collaborate to this project and Cédric Letourneur for collecting survey data. We thank the two
519 anonymous reviewers who helped us improving the manuscript.

520 References

521 Angiolini, C., Bonari, G., Landi, M., 2018. Focal plant species and soil factors in Mediterranean coastal
522 dunes: an undisclosed liaison? *Estuarine, Coastal and Shelf Science* 211, 248-258.

523 Ansquer, P., Duru, M., Theau, J.P., Cruz, P., 2009. Convergence in plant traits between species within
524 grassland communities simplifies their monitoring. *Ecological Indicators* 9, 1020-1029.

525 APG (Angiosperm Phylogeny Group), 2009. An update of the Angiosperm Phylogeny Group
526 classification for the orders and families of flowering plants: APG III. *Botanical Journal of the Linnean
527 Society* 161, 105-121.

528 Bazzichetto, M., Malavasi, M., Acosta, A.T.R., Carranza, M.L., 2016. How does dune morphology
529 shape coastal EC habitats occurrence? A remote sensing approach using airborne LiDAR on the
530 Mediterranean coast. *Ecological Indicators* 71, 618-626.

531 Bell, C.D., Soltis, D.E., Soltis P.A., 2005. The age of the angiosperms: a molecular timescale without a
532 clock. *Evolution* 59, 1245-1258.

533 Benson, D.A., Karsch-Mizrachi, I., Lipman, D.J., Ostell, J., Wheeler, D.L., 2006. GenBank. *Nucleic
534 Acids Research* 34, D16–D20.

535 Borcard, D., Gillet, F., Legendre, P., 2018. *Numerical Ecology with R, Second Edition*. Use R! series.
536 Springer, New York.

537 Buttler, A., Kohler, F., Gillet, F., 2009. *The Swiss mountain wooded pastures: patterns and processes*.
538 pp. 377-396 in Rigueiro-Rodriguez, A., McAdam J. & Mosquera-Losada, M.R., eds. *Agroforestry in
539 Europe: current status and future prospects*, Advances in Agroforestry series. Springer, New York.

540 Cadotte, M.W., Carscadden, K., Mirotnick, N., 2011. Beyond species: functional diversity and the
541 maintenance of ecological processes and services. *Journal of Applied Ecology* 48, 1079-1087.

542 Cadotte, M.W., Dinnage, R., Tilman, D., 2012. Phylogenetic diversity promotes ecosystem stability.
543 *Ecology* 93, 223-233.

544 Cadotte, M.W., Albert, C.H., Walker, S.C., 2013. The ecology of differences: assessing community
545 assembly with trait and evolutionary distances. *Ecology Letters* 16, 1234-1244.

546 Cantarello, E., Newton, A.C., 2008. Identifying cost-effective indicators to assess the conservation
547 status of forested habitats in Natura 2000 sites. *Forest Ecology and Management* 256, 815-826.

548 Carmona, C.P., Azcárate, F.M., de Bello, F., Ollero, H.S., Lepš, J., Peco, B., 2012. Taxonomical and
549 functional diversity turnover in Mediterranean grasslands: interactions between grazing, habitat type
550 and rainfall. *Journal of Applied Ecology* 49, 1084-1093.

551 Cavalli, D., Cabassi, G., Borrelli, L., Fuccella, R., Degano, L., Bechini, L., Marino, P., 2014. Nitrogen
552 fertiliser value of digested dairy cow slurry, its liquid and solid fractions, and of dairy cow slurry. *Italian
553 Journal of Agronomy* 9, 71-78.

554 Chiarucci, A., Bacaro, G., Rocchini, D., 2008. Quantifying plant species diversity in a Natura 2000
555 network: old ideas and new proposals. *Biological Conservation* 141 (10), 2608-2618.

556 Chytrý, M., Tichý, L., Holt, J., Botta-Dukát, Z., 2002. Determination of diagnostic species with
557 statistical fidelity measures. *Journal of Vegetation Science* 13, 79-90.

558 Chytrý, M., Otýpková, Z., 2003. Plot sizes used for phytosociological sampling of European vegetation.
559 *Journal of Vegetation Science* 14: 563-570.

560 Cousins, S.A.O., Eriksson, O., 2002. The influence of management history and habitat on plant
561 species richness in a rural hemiboreal landscape, Sweden. *Landscape Ecology* 17, 517-529.

562 Craine, J.M., Nippert, J.B., Towne, E.G., Tucker, S., Kembel, S.W., Skibbe, A., McLauchlan, K.K.,
563 2011. Functional consequences of climate change-induced plant species loss in a tallgrass prairie.
564 *Oecologia* 165, 1109-1117.

565 Critchley, C.N.R, Fowbert, J.A., Wright, B., 2007. Dynamics of species-rich upland hay meadows over
566 15 years and their relation with agricultural management practices. *Applied Vegetation Science* 10,
567 307-314.

568 Dainese, M., Lepš, J., de Bello, Francesco., 2015. Different effects of elevation, habitat fragmentation
569 and grazing management on the functional, phylogenetic and taxonomic structure of mountain
570 grasslands. *Perspectives in Plant Ecology, Evolution and Systematics* 17, 44-53.

571 de Bello, F., Lavergne, S., Meynard, C., Lepš, J., Thuiller, W., 2010. The partitioning of diversity:
572 showing Theseus a way out of the labyrinth. *Journal of Vegetation Science* 21, 992-1000.

573 De Cáceres, M., Font, X., Oliva, F., 2008. Assessing species diagnostic value in large data sets: a
574 comparison between phi coefficient and Ochiai index. *Journal of Vegetation Science* 19, 779-788.

575 De Cáceres, M., Legendre, P., 2009. Associations between species and groups of sites: indices and
576 statistical inference. *Ecology* 90(12), 3566-3574.

577 De Cáceres, M., Chytrý, M., Agrillo, E., Attorre, F., Botta-Dukát, Z., Capelo, J., Czúcz, B., Dengler, J.,
578 Ewald, J., Faber-Langendoen, D., Feoli, E., Franklin, S.B., Gavilán, R., Gillet, F., Jansen, F., Jiménez-
579 Alfaro, B., Krestov, P., Landucci, F., Lengyel, A., Loidi, J., Mucina, L., Peet, R.K., Roberts, D.W.,
580 Roleček, Schaminée, J.H.J., Schmidtlein, S., Theurillat, J.P., Tichý, L., Walker, D.A., Wildi, O., Willner,
581 W., Wiser, S.K., 2015. A comparative framework for broad-scale plot-based vegetation classification.
582 *Applied Vegetation Science* 18, 543-560.

583 Devictor, V., Mouillot, D., Meynard, C., Jiguet, C., Thuiller, W., Mouquet, N., 2010. Spatial mismatch
584 and congruence between taxonomic, phylogenetic and functional diversity: the need for integrative
585 conservation strategies in a changing world. *Ecology letters* 13, 1030-1040.

586 Díaz, S., Purvis, A., Cornelissen, J.H.C., Mace, G.M., Donoghue, M.J., Ewers, R.M., Jordano, P.,
587 Pearse, W.D., 2013. Functional traits, the phylogeny of function, and ecosystem service vulnerability.
588 *Ecology and Evolution* 3(9), 2958-2975.

589 Drummond, A.J., Ho, S.Y.W., Phillips, M.J., Rambaut, A., 2006. Relaxed phylogenetics and dating
590 with confidence. *PLoS Biology* 4: e88.

591 Drummond, A.J., Rambaut, A., 2007. BEAST: Bayesian evolutionary analysis by sampling trees. *BMC*
592 *Evolutionary Biology* 7: 214.

- 593 Evans, D., Arvela, M., 2011. Assessment and Reporting under Article 17 of the Habitats Directive.
594 European Topic Centre on Biological Diversity, Paris, France. 123p.
- 595 Ferrez, Y., 2007. Contribution à l'étude phytosociologique des prairies mésophiles de Franche-Comté.
596 *Les Nouvelles Archives de la Flore Jurassienne* 5, 59-151.
- 597 Ferrez, Y., Bailly, G., Beaufils, T., Collaud, R., Caillet, M., Fernez, T., Gillet, F., Guyonneau, J.,
598 Hennequin, C., Royer, J.M., Schmitt, A., Vergon-Trivaudey, M.J., Vadam, J.C., Vuilleminot, M., 2011.
599 Synopsis des groupements végétaux de Franche-Comté. *Les Nouvelles Archives de la Flore*
600 *Jurassienne et du nord-est de la France*, numéro spécial 1. 282p.
- 601 Flynn, D.F.B., Gogol-Prokurat, M., Nogeire, T., Molinari, N., Trautman Richers, B., Lin, B.B., Simpson,
602 N., Mayfield, M.M., DeClerck, F., 2009. Loss of functional diversity under land use intensification
603 across multiple taxa. *Ecology Letters* 12, 22-33.
- 604 Foley, J.A., DeFries, R., Asner, G.P., 2005. Global Consequences of Land Use. *Science* 309, 570–
605 574.
- 606 Garnier, E., Cortez, J., Billès, G., Navas, M.L., Roumet, C., Debussche, M., Laurent, G., Blanchard, A.,
607 Aubry, D., Bellmann, A., Neill, C., Toussaint, J.P., 2004. Plant functional markers capture ecosystem
608 properties during secondary succession. *Ecology* 85, 2630-2637.
- 609 Garnier, E., Navas, M.L., 2012. A trait-based approach to comparative functional plant ecology:
610 concepts, methods and applications for agroecology. A review. *Agronomy for Sustainable*
611 *Development* 32, 365-399.
- 612 Gaston, K.J., Jackson, S.F., Nagy, A., Cantu-Salazar, L., Johnson, M., 2008. Protected areas in
613 Europe. Principle and practice. *Annals of the New York Academy of Sciences* 1134, 97-119.
- 614 Gaujour, E., Amiaud, B., Mignolet, C., Plantureux, S., 2012. Factors and processes affecting plant
615 biodiversity in permanent grasslands. A review. *Agronomy for Sustainable Development* 32, 133-160.
- 616 Gillet, F., Mauchamp, L., Badot, P.M., Mouly, A., 2016. Recent changes in mountain grasslands: a
617 vegetation resampling study. *Ecology and Evolution* 8, 2333-2345.
- 618 Güler, B., Jentsch, A., Bartha, S., Bloor, J.M.G., Campetella, G., Canullo, R., Házi, J., Kreyling, J.,
619 Pottier, J., Szabó, G., Terziyska, T., Uğurlu, E., Wellstein, C., Zimmermann, Z., Dengler, J., 2016. How
620 plot shape and dispersion affect plant species richness counts: implications for sampling design and
621 rarefaction analyses. *Journal of Vegetation Science* 27: 692-703.
- 622 Hector, A., Hautier, Y., Saner, P., Wacker, L., Bagghi, R., Joshi, J., Scherer-Lorenzen, M., Spehn,
623 E.M., Bazeley-White, E., Weilenmann, M., Caldeira, M.C., Dimitrakopoulos, P.G., Finn, J.A., Huss-
624 Danell, K., Jumpponen, A., Mulder, C.P.H., Palmberg, C., Pereira, J.S., Siamantziouras, A.S.D., Terry,
625 A.C., Troumbis, A.Y., Schmid, B., Loreau, M., 2010. General stabilizing effects of plant diversity on
626 grassland productivity through population asynchrony and overyielding. *Ecology* 91, 2213-2220.
- 627 Heegaard, E., Okland, R.H., Bratli, H., Dramstad, W.E., Engan, G., Pedersen, O., Solstad, H., 2007.
628 Regularity of species richness relationships to patch size and shape. *Ecography* 30: 589-597.
- 629 Hejman, M., Schellberg, J., Pavlů, V., 2010. Long-term effects of cutting frequency and liming on soil
630 chemical properties, biomass production and plant species composition of Lolio-Cynosuretum
631 grassland after the cessation of fertilizer application. *Applied Vegetation Science* 13: 257-269.

632 Hernando, A., Tejera, R., Velázquez, J., Victoria Núñez, M., 2010. Quantitatively defining the
633 conservation status of Natura 2000 forest habitats and improving management options for enhancing
634 biodiversity. *Biodiversity Conservation* 19, 2221-2233.

635 Hill, M.O., 1973. Diversity and evenness: a unifying notation and its consequence. *Ecology* 54, 427-
636 432.

637 Hilu, K.W., Borsch T., Müller, K., Soltis, D.E., Soltis, P.S., Savolainen, V., Chase, M.W., Powell, M.P.,
638 Alice, L.A., Evans, R., Sauquet, H., Neinhuis, C., Slotta, T.A.B., Rohwer, J.G., Campbell, S.C.,
639 Chatrou, L.W., 2003. Angiosperm phylogeny based on matK sequence information. *American Journal*
640 *of Botany* 90, 1758-1776.

641 Hoekstra, J.M., Boucher, T.M., Ricketts, T.H., Roberts, C., 2005. Confronting a biome crisis: global
642 disparities of habitat loss and protection. *Ecology Letters* 8, 23-29.

643 Homburger, H., Hofer, G., 2012. Diversity change of mountain hay meadows in the Swiss Alps. *Basic*
644 *and Applied Ecology* 13, 132-138.

645 Institut de l'Elevage, 2012. *Fertiliser avec les engrais de ferme. Ouvrage collectif*. Institut de l'Elevage,
646 ITCF, ITAVI, ITP, Technipel 250, 1-104.
647 [http://idele.fr/no_cache/recherche/publication/idelesolr/recommends/fertiliser-avec-les-engrais-de-](http://idele.fr/no_cache/recherche/publication/idelesolr/recommends/fertiliser-avec-les-engrais-de-ferme.html)
648 [ferme.html](http://idele.fr/no_cache/recherche/publication/idelesolr/recommends/fertiliser-avec-les-engrais-de-ferme.html) [accessed January 30, 2019].

649 Isbell, F., Craven, D., Connolly, J., Loreau, M., Schmid, B., Beierkuhnlein, C., Bezemer, T.M., Bonin,
650 C., Bruelheide, H., de Luca, E., Ebeling, A., Griffin, J.N., Guo, Q., Hautier, Y., Hector, A., Jentsch, A.,
651 Kreyling, J., Lanta, V., Manning, P., Meyer, S.T., Mori, A.S., Naeem, S., Niklaus, P.A., Wayne Polley,
652 H., Reich, P.B., Roscher, C., Seabloom, E.W., Smith, M.D., Thakur, M.P., Tilman, D., Tracy, B.F., van
653 der Putten, W.H., van Ruijven, J., Weigelt, A., Weisser, W.W., Wilsey, B., Eisenhauer, N., 2015.
654 Biodiversity increases the resistance of ecosystem productivity to climate extremes. *Nature* 526, 574-
655 577.

656 Jost, L., 2007. Partitioning diversity into independent alpha and beta components. *Ecology* 88, 2427-
657 2439.

658 Jost, L., 2010. The Relation between Evenness and Diversity. *Diversity* 2, 207-232.

659 Kattge, J., Diaz, S., Lavorel, S., Prentice, C., Leadley, P., Boenisch, G., Garnier, E., Westoby, M.,
660 Reich, P.B., Wright, I.J., Cornelissen, J.H.C., Violle, C., Harrison, S.P., van Bodegom, P.M.,
661 Reichstein, M., Enquist, B.J., Soudzilovskaia, N.A., Ackerly, D.D., Anand, M., Atkin, O., Bahn, M.,
662 Baker, T.R., Baldocchi, D., Bekker, R., Blanco, C.C., Blonder, B., Bond, W.J., Bradstock, R., Bunker,
663 D.E., Casanoves, F., Cavender-Bares, J., Chambers, J.Q., Chapin, F.S. III, Chave, J., Coomes, D.,
664 Cornwell, W.K., Craine, J.M., Dobrin, B.H., Duarte, L., Durka, W., Elser, J., Esser, G., Estiarte, M.,
665 Fagan, W.F., Fang, J., Fernandez-Mendez, F., Fidelis, A., Finegan, B., Flores, O., Ford, H., Frank, D.,
666 Freschet, G.T., Fyllas, N.M., Gallagher, R.V., Green, W.A., Gutierrez, A.G., Hickler, T., Higgins, S.I.,
667 Hodgson, J.G., Jalili, A., Jansen, S., Joly, C.A., Kerkhoff, A.J., Kirkup, D., Kitajima, K., Kleyer, M.,
668 Klotz, S., Knops, J.M.H., Kramer, K., Kuehn, I., Kurokawa, H., Laughlin, D., Lee, T.D., Leishman, M.,
669 Lens, F., Lenz, T., Lewis, S.L., Lloyd, J., Llusia, J., Louault, F., Ma, S., Mahecha, M.D., Manning, P.,
670 Massad, T., Medlyn, B.E., Messier, J., Moles, A.T., Mueller, S.C., Nadrowski, K., Naeem, S.,
671 Niinemets, U., Noellert, S., Nueske, A., Ogaya, R., Oleksyn, J., Onipchenko, V.G., Onoda, Y.,
672 Ordonez, J., Overbeck, G., Ozinga, W.A., Patino, S., Paula, S., Pausas, J.G., Penuelas, J., Phillips,
673 O.L., Pillar, V., Poorter, H., Poorter, L., Poschlod, P., Prinzing, A., Proulx, R., Rammig, A., Reinsch, S.,
674 Reu, B., Sack, L., Salgado-Negre, B., Sardans, J., Shiodera, S., Shipley, B., Siefert, A., Sosinski, E.,

675 Soussana, J.F., Swaine, E., Swenson, N., Thompson, K., Thornton, P., Waldram, M., Weiher, E.,
676 White, M., White, S., Wright, S.J., Yguel, B., Zaehle, S., Zanne, A.E., Wirth, C., 2011. TRY – a global
677 database of plant traits. *Global Change Biology* 17, 2905-2935.

678 Kirkham, F.W., Tallowin, J.R.B., Sanderson, R.A., Bhogal, A., Chambers, B.J., Stevens, D.P., 2008.
679 The impact of organic and inorganic fertilizers and lime on the species-richness and plant functional
680 characteristics of hay meadow communities. *Biological Conservation* 141, 1411-1427.

681 Kleyer, M., Bekker, R.M., Knevel, I.C., Bakker, J.P., Thompson, K., Sonnenschein, M., Poschlod, P.,
682 van Groenendael, J.M., Klimeš, L., Klimešová, J., Klotz, S., Rusch, G.M., Hermy, R., Adriaens, D.,
683 Boedeltje, G., Bossuyt, B., Dannemann, A., Endels, P., Götzenberger, L., Hodgson, J.G., Jackel, A.K.,
684 Kühn, I., Kunzmann, D., Ozinga, W.A., Römermann, C., Stadler, M., Schlegelmilch, J., Steendam,
685 H.J., Tackenberg, O., Wilmann, O., Cornelissen, J.H.C., Eriksson, O., Garnier, E., Peco, B., 2008. The
686 LEDA Traitbase: a database of life-history traits of Northwest European flora. *Journal of Ecology* 96,
687 1266-1274.

688 Kottek, M., Grieser, J., Beck, C., Rudolf, B., Rubel, F., 2006. World Map of the Köppen–Geiger climate
689 classification updated. *Meteorologische Zeitschrift* 15 (3), 259–263.

690 Kovač, M., Kutnar, L., Hladnik, D., 2016. Assessing biodiversity and conservation status of the Natura
691 2000 forest habitat types: Tools for designated forestlands stewardship. *Forest Ecology and*
692 *Management* 359, 256-267.

693 Larkin, M.A., Blackshields, G., Brown, N.P., Chenna, R., McGettigan, P.A., McWilliam, H., Valentin, F.,
694 Wallace, I.M., Wilm, A., Lopez, R., Thompson, J.D., Gibson, T.J., Higgins D.G., 2007. Clustal W and
695 Clustal X version 2.0. *Bioinformatics* 23, 2947-2948.

696 Lososová, Z., Čeplová, N., Chytrý, M., Tichý, L., Danihelka, J., Fajmon, K., Láníková, D., Preislerová,
697 Z., Řehořek, V., 2016. Is phylogenetic diversity a good proxy for functional diversity of plant
698 communities? A case study from urban habitats. *Journal of Vegetation Science* 27, 1036-1046.

699 Louault, F., Pillar, V.D., Aufrère, J., Garnier, E., Soussana, J.F., 2005. Plant traits and functional types
700 in response to reduced disturbance in a semi-natural grassland. *Journal of Vegetation Science* 16,
701 151-160.

702 Louette, G., Adriaens, D., Paelinckx, D., Hoffmann, M., 2015. Implementing the Habitats Directive:
703 How Science can support decision making. *Journal for Nature Conservation* 23, 27-34.

704 Mariotte, P., Vandenberghe, C., Kardol, P., Hagedorn, F., Buttler, A., 2013. Subordinate plant species
705 enhance community resistance against drought in semi-natural grasslands. *Journal of Ecology* 101,
706 763-773.

707 Mašková, Z., Doležal, J., Květ, J., Zemek, F., 2009. Long-term functioning of a species-rich mountain
708 meadow under different management regimes. *Agriculture, Ecosystems and Environment* 132, 192-
709 202.

710 Mauchamp, L., Gillet, F., Mouly, A., Badot, P.M., 2013. *Les prairies : biodiversité et services*
711 *écosystémiques*. Presses universitaires de Franche-Comté – Conseil National des Appellations
712 d'Origine Laitières, Besançon, France. 130p.

713 Mauchamp, L., Mouly, A., Badot, P.M., Gillet, F., 2014. Impact of management type and intensity on
714 multiple facets of grassland biodiversity in the French Jura Mountains. *Applied Vegetation Science* 17,
715 645-657.

716 Mauchamp, L., Mouly, A., Badot, P.M., Gillet, F., 2016. Impact of nitrogen inputs on multiple facets of
717 plant biodiversity in mountain grasslands: does nutrient source matter? *Applied Vegetation Science*
718 19, 206-217.

719 Mitchell, R.J., Hewison, R.L., Britton, A.J., Brooker, R.W., Cummins, R.P., Fielding, D.A., Fisher, J.M.,
720 Gilbert, D.J., Hester, A.J., Hurskainen, S., Pakeman, R.J., Potts, J.M., Riach, D., 2017. Forty years of
721 change in Scottish grassland vegetation: Increased richness, decreased diversity and increased
722 dominance. *Biological Conservation* 212, 327-336.

723 Mucina, L., Bültmann, H., Dierssen, K., Theurillat, J.P., Raus, T., Čarni, A., Šumberová, K., Willner, W.,
724 Dengler, J., Gavilán García, R., Chytrý, M., Hájek, M., Di Pietro, R., Iakushenko, D., Pallas, J.,
725 Daniëls, F.J.A., Bergmeier, E., Santos Guerra, A., Ermakov, N., Valachovič, M., Schaminée, J.H.J.,
726 Lysenko, T., Didukh, Y.P., Pignatti, S., Rodwell, J.S., Capelo, J., Weber, H.E., Solomeshch, A.,
727 Dimopoulos, P., Aguiar, C., Hennekens, S.M., Tichý, L., 2016. Vegetation of Europe: hierarchical
728 floristic classification system of vascular plant, bryophyte, lichen, and algal communities. *Applied*
729 *Vegetation Science* 19 (Suppl. 1), 3-264.

730 OJEC, 1992. Council Directive 92/43/EEC of 21 May 1992 on the conservation of natural habitats and
731 of wild fauna and flora. *Official Journal of the European Community*, L, 206.

732 Oksanen, J., Blanchet, F.G., Friendly, M., Kindt, R., Legendre, P., McGlinn, D., Minchin, P.R., O'Hara,
733 R.B., Simpson, G.L., Solymos, P., Stevens, M.H.H., Szoecs, E., Wagner H., 2018. vegan: Ordination
734 methods, diversity analysis and other functions for community and vegetation ecologists [online].
735 Available from <http://cran.r-project.org/web/packages/vegan/index.html> [accessed October 9, 2018].

736 Oliver, T.H., Heard, M.S., Isaac, J.B., Roy, D.B., Procter, D., Eigenbrod, F., Freckleton, R., Hector, A.,
737 Orme, C.D.L., Petchey, O.L., Proença, V., Raffaelli, D., Suttle, K.B., Mace, G.M., Martin-López, B.,
738 Woodcock, B.A., Bullock, J.M., 2015. Biodiversity and Resilience of Ecosystem Functions. *Trends in*
739 *Ecology and Evolution* 30, 673-684.

740 Ostermann, O.P., 1998. The need for management of nature conservation sites designated under
741 Natura 2000. *Journal of Applied Ecology* 35(6), 968-973.

742 Pakeman, R., Hewison, R.L., Riach, D., Fisher, J.M., Hurskainen, S., Fielding, D.A., Mitchell, R.J.,
743 2017. Long-term functional structure and functional diversity changes in Scottish grasslands.
744 *Agriculture, Ecosystems and Environment* 247, 352-362.

745 Pavoine, S., Bonsall, M.B., 2011. Measuring biodiversity to explain community assembly: a unified
746 approach. *Biological Reviews* 86, 792-812.

747 Perronne, R., Mauchamp, L., Mouly, A., Gillet, F., 2014. Contrasted taxonomic, phylogenetic and
748 functional diversity patterns in semi-natural permanent grasslands along an altitudinal gradient. *Plant*
749 *Ecology and Evolution* 147, 165-175.

750 Plantureux, S., Peeters, A., McCracken, D., 2005. Biodiversity in intensive grasslands. Effect of
751 management, improvement and challenges. *Agronomy Research* 3, 153-164.

752 Rion, V., Gallandat, J.D., Gobat, J.M., Vittoz, P., 2018. Recent changes in the plant composition of
753 wetlands in the Jura Mountains. *Applied vegetation Science* 21, 121-131.

754 Rubel, F., Brugger, K., Haslinger, K., Auer, I., 2017. The climate of the European Alps: Shift of very
755 high resolution Köppen-Geiger climate zones 1800-2100. *Meteorologische Zeitschrift* 26, 115-125.

- 756 Santoro, R., Carboni, M., Carranza, M.L., Acosta, A.T.R., 2012. Focal species diversity patterns can
757 provide diagnostic information on plant invasions. *Journal for Nature Conservation* 20 (2), 85-91.
- 758 Schermer, M., Darnhofer, I., Daugstad, K., Gabillet, M., Lavorel, S., Steinbacher, M., 2016. Institutional
759 impacts on the resilience of mountain grasslands: an analysis based on three European case studies.
760 *Land Use Policy* 52, 382-391.
- 761 Schirpke, U., Kohler, M., Leitinger, G., Fontana, V., Tasser, E., Tappeiner, U., 2017. Future impacts of
762 changing land-use and climate on ecosystem services of mountain grassland and their resilience.
763 *Ecosystem Services* 26, 79-94.
- 764 Tichý, L., Chytrý, M., 2006. Statistical determination of diagnostic species for site groups of unequal
765 size. *Journal of Vegetation Science* 17, 809-818.
- 766 Tilman, D., 1988. *Plant Strategies and the Dynamics and Structure of Plant Communities*. Princeton
767 University Press, Princeton, New Jersey, United States. 366pp.
- 768 van der Maarel, E., 1979. Transformation of cover-abundance values in phytosociology and its effect
769 on community similarity. *Vegetatio* 39, 97-114.
- 770 Velázquez, J., Tejera, R., Hernando, A., Núñez, M.V., 2010. Environmental diagnosis: Integrating
771 biodiversity conservation in management of Natura 2000 forest spaces. *Journal for Nature
772 Conservation* 18, 309-317.
- 773 Villalobos, S., Vamosi, J.C., 2016. Increasing land use drives changes in plant phylogenetic diversity
774 and prevalence of specialists. *PeerJ* 4, e1740.
- 775 Wesche, K., Krause, B., Culmsee, H., Leuschner, C., 2012. Fifty years of change in Central European
776 grassland vegetation: Large losses in species richness and animal-pollinated plants. *Biological
777 Conservation* 150, 76-85.
- 778 Westoby, M., Falster, D.S., Moles, A.T., Vesk, P.A., Wright, I.J., 2002. Plant ecological strategies:
779 some leading dimensions of variation between species. *Annual Review of Ecology and Systematics*
780 33, 125-159.
- 781 Willner, W., Tichý, L., Chytrý, M., 2009. Effects of different fidelity measures and contexts on the
782 determination of diagnostic species. *Journal of Vegetation Science* 20, 130-137.
- 783 Wilson, J.B., Peet, R.K., Dengler, J., Pärtel, M., 2012. Plant species richness: the world records.
784 *Journal of Vegetation Science* 23, 796-802.
- 785 Yang, Z., 1994. Maximum likelihood phylogenetic estimation from DNA sequences with variable rates
786 over sites, approximate methods. *Journal of Molecular Evolution* 39, 306-314.