

1 **Small rocky outcrops: natural features to promote biodiversity in oak**
2 **wood-pastures**

3 Chozas, Sergio^{1*}, Tapia, Susana¹, Palmeirim, Jorge¹, Alegria, Carla², Correia, Otilia¹

4 ¹ cE3c – Centro de Ecologia, Evolução e Alterações Ambientais, Faculdade de Ciências,
5 Universidade de Lisboa. Campo Grande, 1749-016 Lisboa, Portugal.

6 ² SFCOLAB - Associação Smart Farm COLAB Laboratório Colaborativo para a Inovação
7 Digital na Agricultura, Rua Cândido dos Reis nº1, Espaço SFCOLAB, 2560-312 Torres Vedras,
8 Portugal

9
10 **Note:** This is the preprint version of the manuscript. The journal published article can be
11 accessed in the [website](https://doi.org/10.1111/avsc.12634) of the Applied Vegetation Science journal.
12 <https://doi.org/10.1111/avsc.12634>

13

14 **Running Head:** Rocky outcrops promote wood-pasture biodiversity

15 * Corresponding autor

16 Sergio Chozas: <https://orcid.org/0000-0001-6741-1259>

17 Susana Tapia: <https://orcid.org/0000-0002-6645-8159>

18 Jorge Palmeirim: <https://orcid.org/0000-0003-4734-8162>

19 Carla Alegria: <https://orcid.org/0000-0002-9461-4569>

20 Otilia Correia: <https://orcid.org/0000-0002-1053-0561>

21 **Abstract**

22 **Aims:** The Iberian oak wood-pastures are unique agroforestry systems supporting high
23 levels of biodiversity and ecosystem services. Small rocky outcrops are geological
24 features commonly found in these systems and constitute biodiversity reservoirs,
25 protecting sensitive species from grazing and farming activities. We aimed to assess the
26 relevance of including rocky outcrop conservation within wood-pastures to increase
27 biodiversity. To achieve this goal, we study the plant communities occurring within the
28 outcrops and in the wood-pasture matrix to evaluate the impact of rocky outcrops on the
29 overall plant taxonomic and functional diversities of these systems.

30 **Location:** Montemor-o-Novo (Alentejo, Portugal)

31 **Methods:** We sampled 102 plant communities occurring in outcrops and in the adjacent
32 wood-pasture matrix and analysed alpha, beta, gamma and functional diversities. We
33 identified the main intrinsic factors affecting outcrop plant composition and their
34 functional groups using Linear (LM) and Generalised Linear Mixed Models (GLMMs) and
35 characterised the effect of outcrop size throughout Generalised Additive Models (GAMs).

36 **Results:** We found plant richness to be similar in wood-pasture matrix and outcrops.
37 However, beta diversity analysis revealed a high species turnover between both
38 communities. Functional indices indicated a higher plant functional diversity in outcrops
39 and trait analyses identified three functional groups dissimilarly distributed in both
40 communities: i) perturbation and stress-sensitive plants, with outcrops constituting an
41 important refuge for this group, and ii) grazing-tolerant and iii) weedy herbs dominating
42 the wood-pastures. Finally, we also found increased plant richness in outcrops with a
43 higher length of their minor axis, i.e. wider outcrops, and higher rock cover area.

44 **Conclusions:** Our results indicate that the presence of small rocky outcrops in
45 evergreen oak wood-pastures highly increase their gamma and functional diversities.
46 Consequently, outcrop protection strongly impacts overall wood-pasture biodiversity and

47 underline the suitability of including outcrop conservation as a cost-effective solution
48 capable of increasing biodiversity in these agroforestry systems.

49

50 **Keywords:** Agroforestry, Conservation, Environmental heterogeneity, Functional
51 diversity, Montado/Dehesa, Nature-based Solutions, Plant diversity, Small habitat
52 patches, SLOSS debate, Sustainable development.

53

54 **Introduction**

55 Global biodiversity has been declining for the last decades (Pereira et al. 2013; Davis et
56 al. 2018). Among the global change factors responsible for biodiversity loss, land-use
57 change, habitat fragmentation, and climate change are the most detrimental (Tilman et
58 al. 2017). Biodiversity loss linked to climate change strongly impacts ecosystem
59 functioning (EF) and the delivery of multiple ecosystem services (ES) (Allan et al. 2015).
60 Thus, a paradigm shift is needed to develop strategies that conciliate economic
61 development, biodiversity conservation and ecosystem functioning (Oliver and Morecroft
62 2014).

63 In this context, Nature-based Solutions (NbS), defined as actions to protect,
64 sustainably manage and restore natural or modified ecosystems (Cohen-Shacham et al.
65 2016), constitute cost-effective means to safeguard biodiversity and EF while mitigating
66 climate change (Keesstra et al. 2018). For example, increasing green space and planting
67 trees in urban areas can mitigate the impact of the urban heat island effect (Grilo et al.
68 2020). Agroforestry systems are identified as promising systems to implement NbS,
69 since they support high biodiversity levels (Torralba et al. 2016), substantially contribute
70 to climate change strategies (Verchot et al. 2007), and foster EF (Jose 2009) while
71 generating socio-economic benefits (Pavlidis and Tsihrintzis 2018).

72 The Iberian evergreen oak wood-pastures, known as *Montados* in Portugal and
73 *Dehesas* in Spain, occupy a 3.5 – 4 million ha area and emerge as key semi-natural
74 systems to implement NbS promoting climate change mitigation and supporting
75 biodiversity in the Iberian Peninsula (Olea and San Miguel-Ayanz 2006). This system is
76 structurally similar to savannah type ecosystems, with cork oak (*Quercus suber*) and
77 holm oak (*Quercus rotundifolia*) coexisting with pastures and crops (Pinto-Correia et al.
78 2011). It results from the transformation of ancient evergreen oak woodland areas by
79 human activity over hundreds of years (Bugalho et al. 2011). This low-intensive farming
80 regime maintain ecosystems that deliver a wide range of Ecosystem Services. These
81 good and services range from direct provisioning services (e.g., cork production,
82 livestock) to indirect regulation and supporting services (e.g., carbon sequestration, soil
83 conservation) and cultural services (recreation or conservation of rare species) (Branco
84 et al. 2010). Besides its socio-economic relevance, when adequately managed, this
85 system maintains its internal structural diversity (Bugalho et al. 2011). Due to its role in
86 the biodiversity conservation in Iberian landscapes, these wood-pastures are considered
87 High Nature Value Farming Systems (Paracchini et al. 2008). Despite its significance,
88 this habitat endures various negative pressures being classified as Near Threatened by
89 the European Red List of Habitats (Janssen et al. 2016) while its conservation status
90 was assessed as Unfavourable/Bad by the Habitats Directive (EEA 2013).

91 Underlining the importance of small patches for conservation in the context of the
92 'single large or several small' (SLOSS) debate (Wintle et al. 2019; Fahrig 2020; Deane
93 et al. 2020), especially in countryside ecosystems (Pereira and Daily 2006; Mendenhall
94 et al. 2014), several authors have demonstrated that habitat heterogeneity resulting from
95 small landscape discontinuities can substantially improve the overall biodiversity wealth
96 in wood-pastures (Moreno et al. 2015; Leal et al. 2016; Concepción et al. 2020). Previous
97 research showed that Small Natural Features (according to Hunter et al. 2017) such as
98 olive orchards and riparian galleries enhance environmental heterogeneity and,

99 consequently, species diversity of mammals and birds in oak wood-pastures (Rosalino
100 et al. 2009; Leal et al. 2011). Recently, Oksuz et al. (2020) identified small shrubby
101 patches as promising NbS to increase *Montado* biodiversity, finding them frequently, but
102 not always, associated with small rocky outcrops. Rocky outcrops are geological
103 formations that are common in *Montados* and contribute to increasing their structural
104 heterogeneity (Martín and Lopez 2002). These features constitute reservoirs of
105 biodiversity, protecting sensitive species from grazing and farming activities (Fitzsimons
106 and Michael 2017), developing small forest habitat fragments embedded in a grassland
107 matrix (Plieninger et al. 2004). The buffering capacity of outcrops towards environmental
108 changes, e.g., by conducting and retaining run-off water (Speziale and Ezcurra 2015),
109 make them potential “stepping-stones” within metacommunity systems, that may allow
110 the persistence and migration of species in response to global change impacts (Ottaviani
111 et al. 2019). Outcrops also provide additional features for ecosystem services of wood-
112 pastures such as refuges and food for birds, mammals and pollinators, including small
113 and big game species (Bauer et al. 2017); they enhance carbon sequestration and
114 storage and contribute to local water balance control (Centeno et al. 2010) and they
115 constitute recreation and tourism areas (Twidale 2000).

116 Given these facts, outcrops enhance the structural heterogeneity of wood-pastures
117 and can, therefore, support a higher variety of communities (Benton et al. 2003).
118 However, there is a knowledge gap in the literature addressing the conservation value
119 of rocky outcrops. In this work, we continue the line of research initiated by Oksuz (2020)
120 regarding the impact of Small Natural Features on biodiversity in Portuguese *Montados*.
121 That study included several taxa, however, with a limited reach due to the small number
122 of surveyed patches compliant to all the taxa. Free of that constraint, in the present study,
123 we analyse the impact of rocky outcrops on plant taxonomic and functional diversity and
124 the influence of the outcrop’s characteristics (e.g., vegetation and rock cover, outcrop
125 size) on that diversity. We hypothesise that these small geologic elements are occupied

126 by distinct plant communities enhancing both plant diversity and plant functional diversity
127 of the total landscape in Iberian oak wood-pastures and that this distinctiveness is
128 dependent on the outcrop characteristics. The results will provide information about the
129 effectiveness of including rocky outcrop as cost-effective NbS capable of increasing
130 biodiversity and promoting Ecosystem services of wood-pastures.

131

132 **Methods**

133 *Study system*

134 The study area, comprising about 33.5 km², is located in Montemor-o-Novo, Alentejo,
135 Portugal (Figure 1), in the southern margin of the Tagus river (38°46'N, 8°14'W; 38°41'N,
136 8°20'W), with elevation varying from 130 to 180 m a.s.l. The climate is mild
137 Mediterranean with an oceanic influence, characterised by a warm, dry summer and
138 strong seasonal and interannual variability in precipitation and temperature (mean
139 annual precipitation of 660 mm and mean annual air temperature of 15.4 °C (SNIRH
140 2019). A total of 9 farmsteads were sampled. These farmsteads hold large oak wood-
141 pasture areas used for cork extraction, livestock raising (cattle, sheep and pig) and
142 hunting. Dominant soil types are dry acidic soils, and small rocky outcrops cover less
143 than 0.5% of the total wood-pasture area (Oksuz et al. 2020). The vegetation of these
144 patches usually includes holm and cork oaks, olive trees and a mixed-species shrub and
145 lianoid understory composed by typical Mediterranean evergreen oak woodland species.
146 Fieldwork was conducted in May and June 2013.

147 *Community surveys and explanatory variables*

148 We sampled plant communities from 32 identified outcrops within the nine-farmstead
149 area. On each outcrop, we placed one 10×10 m plot assuming that: i) the generalised
150 shape of the outcrops is an ellipse; and ii) the central axis of the plot includes the centre

151 of the ellipse and is perpendicular to the major axis of the ellipse (Figure 2).
152 Simultaneously, we placed 32 10×10 m plots on pastures in the vicinity of the sampled
153 outcrops. Plots were located randomly but always in flat areas, avoiding slopes and
154 significant topographic variations. Onwards, "outcrop" or "matrix" are used to identify the
155 respective plots, for short.

156 The composition and cover of plant species in the plots were estimated using an
157 extended scale adapted from the Braun-Blanquet cover-abundance scale (5: 75-100%;
158 4: 50-75%; 3: 25-50%; 2: 5-25%; 1: few individuals; 0.5: very few individuals; 0.1: one
159 individual) (Damgaard 2014). In each plot, we registered cover of rocks, mosses, lichens,
160 bare soil, and litter using the same scale (Appendix S1). The height of trees, shrubs,
161 herbs, and rocks was also measured.

162 Plant nomenclature follows the Checklist da Flora de Portugal (ALFA -Associação
163 Lusitana de Fitossociologia. 2010), and species identities determined using *Flora Iberica*
164 (Castroviejo 1986-2015) and *Nova Flora de Portugal* (Franco 1971-1984; Franco &
165 Afonso 1994-2003).

166 *Comparative analyses*

167 First, we assessed the plant richness in both communities. Variation in species
168 composition among outcrops and wood-pasture matrices, i.e beta diversity, was
169 calculated using presence and absence data accounting for the spatial turnover –
170 species replacement between both communities – and the nestedness – species loss
171 from community to community – components (Baselga and Leprieur 2015). Pair-wise
172 Wilcoxon tests comparing richness, plant life form, namely herbaceous, shrub, climber
173 and tree, were performed to assess differences in community structure between outcrops
174 and matrix. We also compared the cover between the two communities for all tree
175 species, both adult and shrubby stages. As a measure of regeneration, we considered
176 trees when individuals were ≥ 2 m height.

177 Indicator species are used as ecological indicators of communities, and they
178 ultimately represent qualitative characteristics of the ecosystem (Cáceres et al. 2010).
179 We identified the indicator species in both communities by calculating all species'
180 indicator values (IndVal; Legendre and Legendre 1998). This index quantifies the fidelity
181 and specificity of each species to a given type of community We used the "labdsv" R
182 Package to calculate the IndVal values (Roberts 2015).

183 The description of the vegetation composition was achieved by a non-metric
184 multidimensional scaling (NMS) ordination of the study plots based on the cover of the
185 plant species (van der Maarel 2007), using the "function metaMDS" of R Package vegan
186 (Oksanen et al. 2013). Braun-Blanquet cover-abundance scores were converted to a
187 percentage scale ranging from 2.5% to 87.5% (1 = 2.5%, 2 = 15%, 3 = 37.5%, 4 = 62.5%,
188 5 = 87.5%). We used Bray-Curtis clustering to measure the dissimilarity between plots
189 and assessed the goodness-of-fit of the ordination through the percentage of variance
190 represented by each consecutive axis (see McCune & Grace 2002 for details). NMS
191 axes resulting from these analyses represent the dissimilarity in plant composition.

192 We explored the relationships of outcrop and wood-pasture matrix
193 characteristics, namely cover and height of trees, shrubs, herbs (maximum) and rocks
194 (maximum and average) and bare soil cover, with their plant communities by correlating
195 these characteristics with the NMS ordination using the "envfit" function of vegan. The
196 strength of those relationships was evaluated through the squared correlation
197 coefficient (r^2).

198 To assess the functional diversity of both outcrops and the oak wood-pasture
199 matrix, we classified plant species regarding ten selected traits (Appendix S2, Table S1).
200 We considered seven traits associated with plant responses to grazing (tolerant or not),
201 disturbance (weed behaviour or not), drought (drought-tolerant, indifferent or drought-
202 avoiders), edaphic conditions (acidophilous, indifferent or basophilous; rupicolous or not;

203 and nitrophilous or not) and light (sciophilous or heliophilous). We also included dispersal
204 strategy (short or long dispersal, following Vittoz and Engler, 2007) and life and growth
205 forms as generalist traits informing about climate, disturbance, competitive ability and
206 defence responses of plants (Wright et al. 2006). Together, the ten traits allow the
207 characterisation of the adaptive responses of sampled species to outcrops and matrix
208 (Cadotte et al. 2011) by calculating: i) functional diversity indices, namely functional
209 richness, evenness, divergence, dissimilarity and Rao's quadratic entropy (Villéger et al.
210 2008); and ii) identifying functional groups. Plant functional groups were defined using a
211 dendrogram of species based on the ten trait values and built according to Ward's
212 hierarchical agglomerative clustering method (Murtagh and Legendre 2014). Functional
213 diversity analyses were performed using the "FD" package (Laliberté et al. 2014).

214 Finally, Pair-wise Wilcoxon tests comparing functional indices and groups of both
215 communities were performed to assess differences in the functional diversity between
216 outcrops and the matrix.

217 *Outcrop characteristics' effect*

218 To assess the effect of rocky outcrop characteristics in plant composition and functional
219 diversity, we included 19 additional outcrops and we measured the area and the
220 perimeter of the 51 outcrops using orthophotos extracted from Google Earth and
221 analysed with ArcGis software (ESRI 2019). We registered the composition and cover of
222 plant species in the outcrops along the transect defined by the semi-minor axis of the
223 ellipse, identified as outcrop axis, using the point-intercept method (Nunes et al. 2015).
224 At each (intercepted) point, spaced every 50 cm along the transect, a 5 mm diameter rod
225 was stuck in the ground making a 90° angle. All plant species, rocks, litter and bare soil
226 touching the rod were recorded. We calculated cover estimates as the proportion of
227 points intercepted per transect. The same measurements were taken in the matrix plots,

228 using as sampling transect the segment linking the centre of two opposite sides of the
229 plot (matrix transect).

230 We tested the effects of i) outcrop size (area, perimeter and length of the outcrop
231 axis), ii) cover of rocks, plants, trees, shrubs, herbs, bare soil and litter and iii) the height
232 of trees, shrubs and herbs on the outcrop taxonomical diversity (richness, Simpson and
233 Shannon indices) and functional diversity (functional group cover). This was done
234 performing Linear (LM) and Generalized Linear Mixed Models (GLMMs). Multicollinearity
235 among potential explanatory variables was handled by dropping collinear covariates
236 when correlated at $|\text{Spearman } r| > 0.8$ (Zuur et al. 2010). We modelled outcrop size, cover
237 and height variables as fixed effects and farmsteads as a random effect to control their
238 potential variability using the restricted maximum likelihood method (REML). To compare
239 the fits of LM and GLMMs models, we performed ANOVAs with the regression objects
240 as two separate arguments. These analyses were performed using “nlme” (Pinheiro et
241 al. 2021) and “lm4” (Bates et al. 2015) packages. When necessary, data were
242 transformed for normality and beta regression was performed using the R packages
243 “glmmADMB” (Skaug et al. 2013) and glmmTMB packages (Brooks et al. 2017) when the
244 dependent variables were beta-distributed. Additionally, when dependent variable
245 assumed the extremes 0 and 1, the transformation $(y^*(n - 1) + 0.5)/n$ where n is the
246 sample size, was performed to allow the application of beta-regression analyses
247 (Smithson and Verkuilen 2006). Marginal and conditional coefficients of determination
248 for the models were calculated using “MuMIn” package and the fit for each model was
249 validated using the DHARMA-package (Hartig 2020). Then, we characterise the
250 relationship between the cover of the functional groups and the length of both the outcrop
251 axis and the matrix transect using Generalised Additive Models (GAMs) from the “mgcv”
252 software package (Wood 2006). DHARMA-package was also used to validate GAM
253 models.

254 Finally, we studied the effect of spatial distance among sites on the existing plant
255 communities. To do this, we performed Mantel tests considering the geographic
256 distances and species cover matrix to determine if plant communities were spatially
257 structured, using the “ecodist” package (Goslee and Urban 2007).

258 All statistical analyses were performed using the computing environment R
259 version 4.0.3 (R Core Team 2021).

260

261 **Results**

262 *Comparative analyses*

263 From the 64 characterised plots, we sampled a total of 205 species: 63 exclusively in
264 outcrops, 53 in wood-pastures and 89 species occurring in both plot types (Appendix S3,
265 Table S1). For each species, the values of the ten selected traits were compiled
266 (Appendix S2: Table S2). Overall beta diversity between wood-pasture matrices and
267 outcrops was 0.398, clearly dominating the turnover component (0.375) over nestedness
268 (0.023).

269 Pair-wise Wilcoxon tests indicated that plant richness was not significantly different
270 between outcrop and wood-pasture plots (Table 1). Same tests comparing the species
271 number regarding life form found significant differences among the number of
272 herbaceous species, shrubs, climbers and trees occurring in outcrops and woody
273 pastures, respectively. Regarding the abundance of tree species of both communities,
274 pair-wise Wilcoxon tests indicated significant differences between outcrops and matrix
275 for holm oak cover for tree (and shrubby stages. Moreover, olive trees (*Olea europaea*),
276 either in adult or shrubby stages, occur only in outcrops. Similarly, kermes oak (*Quercus*
277 *coccifera*) occurs only in tree form within the outcrops. No differences between outcrops
278 and matrix were found regarding cork oak cover stages. Only a shrubby cork oak

279 individual was registered in a wood-pasture plot, probably indicating cork oak
280 regeneration limitations.

281 Indicator value (IndVal) analyses identified 9 and 14 species with indicator values
282 higher than 0.7 for outcrops and wood-pastures, respectively (Appendix S4). Outcrop
283 species with the highest IndVal values were *Geranium robertianum* and *Umbilicus*
284 *rupestris* (0.968 and 0.867, respectively), two rupicolous species (SPBotanica 2013).
285 However, typical ancient oak woodland species such as holm oak (0.769) and shrubs
286 (0.791 for both *Ruscus aculeatus* and *Rhamnus alaternus*) also had high IndVal values.
287 Regarding the matrix species with high IndVal, all were herbaceous species,
288 characteristic of grazing pastures such as *Agrostis pourretii* and *Echium plantagineum*
289 (0.975 and 0.916, respectively).

290 The 2-dimensional NMS ordination based on the plant species cover data, with
291 final stress of 21.84% (Figure 3), described the main differences in vegetation
292 composition. The first axis accounted for the most variance (28.15% out of 39.21%), and
293 it clearly separates the communities occurring in outcrops and the wood-pasture matrix.
294 Correlation analyses show that the main characteristics separating outcrops and wood-
295 pasture matrix were the herb cover, the tree maximum height and the cover of rocks
296 ($r^2 = 0.89, 0.83, \text{ and } 0.73, p < 0.001$, respectively), then the tree cover and the height of
297 shrubs and rocks ($r^2 = 0.60, 0.56 \text{ and } 0.55, p < 0.001$, respectively) and, to a lower extent,
298 the height of herbs and the cover of bare soil ($r^2 = 0.45 \text{ and } 0.26, p < 0.001$, respectively)
299 (Appendix S5).

300 Regarding functional diversity, the average of functional richness, functional
301 evenness, functional divergence and Rao's quadratic entropy (were significantly higher
302 in outcrops than in the matrix while functional dissimilarity presented no significant
303 differences between both communities (Table 1). Trait analyses clearly identified
304 combinations of traits linked to one of the communities, namely annual and grazing-

305 tolerant herbaceous species for the wood-pasture matrix, and rupicolous and sciophilous
306 woody species for outcrops. It also defined three large functional groups: i) species with
307 low tolerance towards environmental stress and perturbation (sensitive species),
308 including woody, rupicolous and sciophilous species, and two perturbation and stress-
309 tolerant species groups, namely ii) grazing-tolerant herbs and iii) weedy herbs. The three
310 functional groups had a significant dissimilar cover in outcrops and wood-pastures
311 (Table 1, Figure 3).

312 *Outcrop characteristics' effect*

313 Outcrop areas ranged from 75 to 5000 m² (969.9 m² on average), perimeter from 40 to
314 300 m (135 m on average) and the outcrop axis (outcrops edge to centre distance)
315 ranged from 2.5 to 10 m, with most between 4.5 and 5 m (4.8 m on average). A total of
316 94 plant species were registered in the 51 outcrops using the point-intercept method
317 (Appendix S3, Table S2).

318 Linear (LM) and Generalized Linear Mixed Models (GLMMs) identified a positive
319 relationship of plant richness with the length of outcrop axes and with their overall plant
320 cover (Table 2). Likewise, we found an increasing trend of taxonomical diversity
321 associated with rock cover. However, rock cover was also related to lower plant cover.
322 Additionally, outcrops with higher tree cover showed higher levels of uniformity
323 (i.e. higher Simpson's diversity index values). Regarding functional diversity, sensitive
324 species benefited from the presence of higher rocks, while weedy and grazing-tolerant
325 herbs respond negatively to them. Finally, differences among farmsteads were only
326 verified for the overall plant cover and weedy herb cover (Table 3).

327 Generalised Additive Models (GAMs) characterised the relationships of the
328 functional groups cover with the length of the outcrop axis (Figure 4a), identifying a sharp
329 gradient from the margins of the outcrop to the centre for grazing-tolerant and weedy
330 herbs, while sensitive species showed a slightly increase to about the middle of the

331 outcrop axis. Moreover, only sensitive species, the dominant functional group in the
332 outcrops, were present in the centre of the outcrop. As expected, GAM models did not
333 identify significant variations of sensitive species and grazing-tolerant herb distributions
334 along the matrix axis but, surprisingly, weedy herbs decreased slightly at the end of the
335 transect (Figure 4b). Furthermore, sensitive species cover was very low (8% on
336 average), while grazing-tolerant herb cover (105% on average) and weedy herbs (67 %
337 on average) dominated the system.

338 Mantel tests analyses indicated a low but significant spatial autocorrelation among
339 the plant communities at a taxonomical level. These similarities were higher among
340 wood-pasture plots ($r = 0.27$, $p < 0.001$) than among outcrops ($r = 0.15$, $p < 0.001$).

341 **Discussion**

342 According to our results, small rocky outcrops in the wood-pastures significantly impact
343 *Montados* by highly increasing their overall plant richness and plant functional
344 biodiversity. Overall plant richness, i.e. gamma diversity, in *Montados* is deeply affected
345 by the presence of outcrops: about 30.7% of the sampled species were recorded
346 exclusively in outcrops, whereas 25.85% only occurred in the surrounding matrix. Both
347 communities had significant differences regarding the cover of dominant life forms. As
348 expected, herbaceous species dominate wood-pasture matrix while trees, shrubs and
349 climbers play an essential role in the outcrops' plant community. Additionally, olive trees
350 and kermes oaks occur only in the outcrops, and the holm oak cover is also significantly
351 higher in the outcrops. However, we found no significant differences in the cover of adult
352 cork oaks.

353 The main factor shaping the differences between both communities was the
354 presence of rocks, in fact, most of the indicator species of the outcrop were rupicolous
355 or tolerant to rocky habitats. While preventing mechanical clearing and limiting grazing,
356 rocks allow the growth of well-developed shrubs and trees. Moreover, these formations

357 determine a vegetation gradient development, from the outcrops margin to the centre,
358 ultimately intensifying the dissimilarities between both communities. Sensitive species,
359 including rupicolous, sciophilous and oak woodland species, dominate the outcrops
360 communities, while matrix communities are mostly composed of grazing-tolerant and
361 weedy herbs. Among the functional indices, the significantly higher functional
362 dissimilarity in outcrops points to a more functionally diverse community with a higher
363 niche differentiation and a lower competition between species (Morcillo et al. 2019). It is
364 precisely the contrast between both communities that points outcrops as a critical factor
365 in maintaining high diversity levels in this managed habitat. Outcrops provide
366 microhabitats and harbour species not found in the surrounding vegetation matrix.
367 Therefore, outcrops constitute local *refugia* for light, heat-intolerant plants and other
368 organisms such as lichens and beetles (Oksuz 2020). The structural complexity of
369 outcrops influences the number and types of species and this variation clearly
370 determines a profound impact in the provision of ecosystem services (Plas 2019),
371 including refuges for small and big game species (Pia et al. 2013), food provision for
372 birds, mammals and reptilians (Ferber et al. 2014) and cultural services (Barroso et al.
373 2012).

374 Analysing the distribution and cover of the functional groups along the outcrop axis
375 allows us to understand how these formations affect plant wood-pasture diversity. In
376 wood-pastures, the functional group's distribution is reasonably homogenous; however,
377 the distribution follows a spatial gradient from the margins to the centre in the outcrops.
378 Two drivers appear to shape plant distribution inside the outcrops: light availability and
379 disturbance. On the one hand, shrubs and trees limit the amount of light inside the
380 outcrops, enabling colonisation by sciophilous species and preventing the occurrence of
381 light-tolerant species. On the other hand, outcrops constitute grazing and farming
382 refuges (Milchunas and Noy-Meir 2002), protecting species without grazing avoidance
383 or disturbance-tolerance traits.

384 Furthermore, outcrop buffering capacity towards environmental changes (Ottaviani
385 et al. 2019) and their widespread distribution throughout the wood-pastures of the study
386 area, make them potential “stepping-stone” systems. This fact is corroborated by the low
387 levels of autocorrelation found among outcrops, confirming that dispersal ability is not a
388 limiting factor in the assembly mechanisms of plant communities in the studied outcrops.
389 These systems also lead to the establishment of source-sink dynamics between outcrops
390 and wood-pastures since they foster the persistence and, when disturbance intensity
391 lowers, the migration of species with different levels of tolerance to disturbance (Ron et
392 al. 2018). Likewise, by increasing the connectivity in the matrix of these agroforestry
393 systems, outcrops promote recolonisation and higher abundances of local populations
394 through immigration like for example for birds (Renjifo 2001). This fact also stands for
395 other taxa such as reptiles, lichens and beetles (Martín and Lopez 2002; Oksuz 2020).

396 Similarly to Milchunas and Noy-Meir (2002), we did not find a significant effect of
397 the full outcrop extent and perimeter in the plant diversity of these geologic formations.
398 These authors found that 86% of small refuge studies reported positive effects on plant
399 diversity compared to 50% for larger refuges. However, we found a clear spatial-
400 dependent effect; taxonomical and functional plant diversity significantly varied from the
401 margin to the centre of the outcrops. These results point to an "edge effect" along the
402 length of the outcrop axis. This buffer area constitutes an ecotone between the matrix
403 and the outcrops with a high ecological value, not only for plants, as our results show,
404 but also for animals which require these boundaries to fulfil their requirements (Leal et
405 al. 2011).

406 Regarding the impact of the farmsteads on outcrop diversity, this effect was only
407 verified for the overall plant and weedy herb covers, probably resulting from the owner's
408 different management practices, such as shrub cutting and grazing intensities. Moreover,
409 the higher spatial autocorrelation found in the wood-pastures suggests that these
410 communities respond similarly to the management strategies of the different farmsteads.

411 However, our data did not allow us to test this hypothesis, and more studies are needed
412 to explore and understand this topic.

413 To summarise, our results indicate that even small rocky outcrops have a strong
414 effect on overall (gamma) plant and functional diversities in evergreen oak wood-
415 pastures, in line with the authors supporting the importance of small habitat patches for
416 biodiversity (Wintle et al. 2019; Fahrig 2020; Deane et al. 2020). This effect is especially
417 relevant since the Portuguese government classified these agroforestry systems as of
418 high priority for biodiversity conservation. We propose that minor changes in
419 management practices, such as the protection of small rocky outcrops or shrub patches,
420 may significantly impact the biodiversity of agroforestry landscapes. Therefore,
421 preserving, enlarging and reshaping pre-existing outcrops or creating “artificial” outcrops
422 where rocks are naturally available, may represent a valuable biodiversity-friendly
423 agroforestry practice (i.e. a nature-based solution), since they contribute to increasing
424 heterogeneity in agroforestry systems without significant impacts on the economic
425 activity of oak wood-pastures. Moreover, it is imperative to plan for protected area
426 networks that promote local environmental heterogeneity, including the protection of
427 rocky outcrops as *refugia* for mesic-adapted species and as potential stepping stones
428 that allow the dispersal of these species between adjacent environments. Additionally,
429 under projected climate-change scenarios, rocky outcrops may provide micro-
430 climatically diverse habitats, distinct from those of the surrounding vegetation matrix,
431 serving as climatic refugia and thereby facilitating the persistence of specialist species.
432 Finally, instead of adopting high-cost-low-return strategies, it would be important for
433 landowners and decision-makers to include rocky outcrop preservation in the land
434 management planning of *Montados*, since they constitute a cost-effective element for the
435 promotion of biodiversity in this agroforestry system.

436

437 **Appendices**

438 Appendix S1 - Plot characteristics

439 Appendix S2 - Plant trait data

440 Appendix S3- Primary data

441 Appendix S4 - Indicator values

442 Appendix S5 - Squared correlation coefficient of environmental factors and functional
443 groups

444 **Credit authorship contribution statement**

445 All authors conceived the idea, discussed the results and commented on the manuscript.

446 SC analysed the data, produced the figures and led the writing; ST designed and
447 conducted fieldwork, and identified the species; CA made substantial contributions to the
448 writing.

449 **Data accessibility**

450 Primary data are presented as Supporting information.

451 **Funding information**

452 This study received Portuguese funds from the Fundação para a Ciência e Tecnologia
453 (FCT) through the PTDC/AGR-AAM/108448/2008 project. SC was supported by the FCT
454 PORBIOTA project through a postdoctoral fellowship. ST was supported by the
455 Associação para a Investigação e Desenvolvimento de Ciências (FCiências.ID). CA was
456 also supported by FCT through an individual postdoctoral fellowship (grant number
457 SFRH/BPD/126703/2016).

458 **Acknowledgements**

459 We are thankful to the landowners, who kindly allowed us to work in their farmlands and
460 to Catarina Costa for the beautiful sketches of the wood-pastures and the rocky outcrops.

461 **Conflict of Interests**

462 The authors declare no conflict of interests.

463 **References**

- 464 ALFA -Associação Lusitana de Fitossociologia. (2010) Checklist da Flora de Portugal
465 (Continental, Açores e Madeira). http://www3.uma.pt/alfa/checklist_flora_pt.html.
466 Accessed 20-sept-2016
- 467 Allan E, Manning P, Alt F, et al (2015) Land use intensification alters ecosystem
468 multifunctionality via loss of biodiversity and changes to functional composition.
469 *Ecol Lett* 18:834–843. <https://doi.org/10.1111/ele.12469>
- 470 Barroso FL, Pinto-Correia T, Ramos IL, et al (2012) Dealing with landscape fuzziness
471 in user preference studies: Photo-based questionnaires in the Mediterranean
472 context. *Landsc Urban Plan* 104:329–342.
473 <https://doi.org/10.1016/J.LANDURBPLAN.2011.11.005>
- 474 Bates D, Mächler M, Bolker BM, Walker SC (2015) Fitting Linear Mixed-Effects Models
475 Using lme4. *J Stat Softw* 67:1–48. <https://doi.org/10.18637/JSS.V067.I01>
- 476 Bauer DM, Bell KP, Nelson EJ, Calhoun AJK (2017) Managing small natural features:
477 A synthesis of economic issues and emergent opportunities. *Biol Conserv* 211:80–
478 87. <https://doi.org/10.1016/J.BIOCON.2017.01.001>
- 479 Benton TG, Vickery JA, Wilson JD (2003) Farmland biodiversity: is habitat
480 heterogeneity the key? *Trends Ecol Evol* 18:182–188.
481 [https://doi.org/10.1016/S0169-5347\(03\)00011-9](https://doi.org/10.1016/S0169-5347(03)00011-9)
- 482 Branco O, Bugalho M, Silva LN, et al (2010) Hotspot areas for biodiversity and
483 Ecosystem Services in montados – HABEaS. Lisbon
- 484 Brooks ME, Kristensen K, van Benthem KJ, et al (2017) Modeling zero-inflated count
485 data with glmmTMB. *bioRxiv* 132753. <https://doi.org/10.1101/132753>
- 486 Bugalho MN, Caldeira MC, Pereira JS, et al (2011) Mediterranean cork oak savannas

487 require human use to sustain biodiversity and ecosystem services. *Front Ecol*
488 *Environ* 9:278–286. <https://doi.org/10.1890/100084>

489 Cáceres M De, Legendre P, Moretti M (2010) Improving indicator species analysis by
490 combining groups of sites. *Oikos* 119:1674–1684. <https://doi.org/10.1111/J.1600->
491 [0706.2010.18334.X](https://doi.org/10.1111/J.1600-0706.2010.18334.X)

492 Cadotte MW, Carscadden K, Mirotchnick N (2011) Beyond species: functional diversity
493 and the maintenance of ecological processes and services. *J Appl Ecol* 48:1079–
494 1087. <https://doi.org/10.1111/j.1365-2664.2011.02048.x>

495 Castroviejo S (ed) *Flora Ibérica* 1-18, 21. Real Jardín Botánico, CSIC, Madrid

496 Centeno JD, García Rodríguez M, Moya Palomares ME (2010) Influence of granite
497 landforms on water balance in semi-arid and humid climates. *Cad do Lab*
498 *xeolóxico Laxe* 35:99–108

499 Cohen-Shacham E, Walters G, Janzen C, Maginnis S (2016) Nature-based solutions to
500 address global societal challenges. IUCN International Union for Conservation of
501 Nature, Gland, Switzerland

502 Concepción ED, Aneva I, Jay M, et al (2020) Optimizing biodiversity gain of European
503 agriculture through regional targeting and adaptive management of conservation
504 tools. *Biol Conserv* 241:108384. <https://doi.org/10.1016/J.BIOCON.2019.108384>

505 Damgaard C (2014) Estimating mean plant cover from different types of cover data: a
506 coherent statistical framework. *Ecosphere* 5:art20. <https://doi.org/10.1890/ES13->
507 [00300.1](https://doi.org/10.1890/ES13-00300.1)

508 Davis M, Faurby S, Svenning J-C (2018) Mammal diversity will take millions of years to
509 recover from the current biodiversity crisis. *Proc Natl Acad Sci U S A* 115:11262–
510 11267. <https://doi.org/10.1073/pnas.1804906115>

511 Deane DC, Nozohourmehrabad P, Boyce SSD, He F (2020) Quantifying factors for
512 understanding why several small patches host more species than a single large
513 patch. *Biol Conserv* 249:108711. <https://doi.org/10.1016/J.BIOCON.2020.108711>

514 EEA (2013) Dehesas with evergreen *Quercus* spp - Report under the Article 17 of the
515 Habitats Directive. Brussels

516 ESRI (2019) ArcGIS Desktop: Release 10.7.1 Redlands, CA: Environmental Systems
517 Research Institute.

518 Fahrig L (2020) Why do several small patches hold more species than few large
519 patches? *Glob Ecol Biogeogr* 29:615–628. <https://doi.org/10.1111/GEB.13059>

520 Fergar SW, Schleuning M, Hemp A, et al (2014) Food resources and vegetation
521 structure mediate climatic effects on species richness of birds. *Glob Ecol Biogeogr*
522 23:541–549. <https://doi.org/10.1111/GEB.12151>

523 Fitzsimons JA, Michael DR (2017) Rocky outcrops: A hard road in the conservation of
524 critical habitats. *Biol Conserv*. <https://doi.org/10.1016/j.biocon.2016.11.019>

525 Franco JA Nova Flora de Portugal, continente e Açores 1-2 (1971-1984). Sociedade
526 Astória, Lisbon

527 Franco JA, Afonso MLR Nova Flora de Portugal, continente e Açores (1994-2003).
528 Escolar Editora, Lisbon

529 Goslee SC, Urban DL (2007) The ecodist Package for Dissimilarity-based Analysis of
530 Ecological Data. *J Stat Softw* 22:1–19

531 Grilo F, Pinho P, Aleixo C, et al (2020) Using green to cool the grey: Modelling the
532 cooling effect of green spaces with a high spatial resolution. *Sci Total Environ*
533 724:138182. <https://doi.org/10.1016/J.SCITOTENV.2020.138182>

534 Hartig F (2020) DHARMA: Residual Diagnostics for Hierarchical Regression Models. In:

535 Compr. R Arch. Netw. <http://florianhartig.github.io/DHARMa/>[https://cran.r-](https://cran.r-project.org/package=DHARMa)
536 [project.org/package=DHARMa](https://cran.r-project.org/package=DHARMa). Accessed 13 Dec 2021

537 Hunter ML, Bauer DM, Bell KP, et al (2017) Conserving small natural features with
538 large ecological roles: A synthetic overview. *Biol Conserv* 211:88–95.
539 <https://doi.org/10.1016/J.BIOCON.2016.12.020>

540 Janssen JAM, Rodwell JS, Garcia Criado M, et al (2016) European Red List of Habitats
541 – Part 2. Terrestrial and freshwater habitats. Luxemburgo. 38 pp.
542 [[https://forum.eionet.europa.eu/european-red-list-habitats/library/terrestrial-](https://forum.eionet.europa.eu/european-red-list-habitats/library/terrestrial-habitats/e.-grasslands/e7.3-mediterranean-wooded-pastu)
543 [habitats/e.-grasslands/e7.3-mediterranean-wooded-pastu](https://forum.eionet.europa.eu/european-red-list-habitats/library/terrestrial-habitats/e.-grasslands/e7.3-mediterranean-wooded-pastu)]. Downloaded October
544 2020

545 Jose S (2009) Agroforestry for ecosystem services and environmental benefits: an
546 overview. *Agrofor Syst* 76:1–10. <https://doi.org/10.1007/s10457-009-9229-7>

547 Keesstra S, Nunes J, Novara A, et al (2018) The superior effect of nature based
548 solutions in land management for enhancing ecosystem services. *Sci Total*
549 *Environ* 610–611:997–1009. <https://doi.org/10.1016/J.SCITOTENV.2017.08.077>

550 Laliberté E, Legendre P, Shipley B (2014) FD: measuring functional diversity from
551 multiple traits, and other tools for functional ecology. R package version 1.0-12.

552 Leal AI, Correia RA, Granadeiro JP, Palmeirim JM (2011) Impact of cork extraction on
553 birds: Relevance for conservation of Mediterranean biodiversity. *Biol Conserv*
554 144:1655–1662. <https://doi.org/10.1016/j.biocon.2011.02.021>

555 Leal AI, Rainho A, Martins RC, et al (2016) Modelling future scenarios to improve
556 woodland landscapes for birds in the Mediterranean. *J Nat Conserv* 30:103–112.
557 <https://doi.org/10.1016/J.JNC.2016.02.001>

558 Legendre P, Legendre L (1998) Numerical ecology, Second Eng. Elsevier Science BV,
559 Amsterdam

560 Martín J, Lopez P (2002) The effect of Mediterranean dehesa management on lizard
561 distribution and conservation. *Biol Conserv* 108:213–219.
562 [https://doi.org/10.1016/S0006-3207\(02\)00107-6](https://doi.org/10.1016/S0006-3207(02)00107-6)

563 McCune B, Grace JB (2002) *Analysis of Ecological Communities*. MjM Software
564 Design, Gleneden Beach, Oregon

565 Mendenhall CD, Karp DS, Meyer CFJ, et al (2014) Predicting biodiversity change and
566 averting collapse in agricultural landscapes. *Nat* 2014 5097499 509:213–217.
567 <https://doi.org/10.1038/nature13139>

568 Milchunas DG, Noy-Meir I (2002) Grazing refuges, external avoidance of herbivory and
569 plant diversity. *Oikos* 99:113–130. [https://doi.org/10.1034/j.1600-](https://doi.org/10.1034/j.1600-0706.2002.990112.x)
570 [0706.2002.990112.x](https://doi.org/10.1034/j.1600-0706.2002.990112.x)

571 Morcillo L, Camacho-Garzón A, Calderón JS, Bautista S (2019) Functional similarity
572 and competitive symmetry control productivity in mixtures of Mediterranean
573 perennial grasses. *PLoS One* 14:e0221667.
574 <https://doi.org/10.1371/journal.pone.0221667>

575 Moreno G, Gonzalez-Bornay G, Pulido F, et al (2015) Exploring the causes of high
576 biodiversity of Iberian dehesas: the importance of wood pastures and marginal
577 habitats. *Agrofor Syst* 2015 901 90:87–105. [https://doi.org/10.1007/S10457-015-](https://doi.org/10.1007/S10457-015-9817-7)
578 [9817-7](https://doi.org/10.1007/S10457-015-9817-7)

579 Nunes A, Tápia S, Pinho P, et al (2015) Advantages of the point-intercept method for
580 assessing functional diversity in semi-arid areas. *iForest - Biogeosciences For*
581 *8:471–479*. <https://doi.org/10.3832/ifor1261-007>

582 Oksanen J, Guillaume Blanchet, F. Roeland Kindt, Pierre Legendre, Peter R. Minchin,
583 R. B. O’Hara, Simpson GL, Solymos P, et al (2013) *vegan: Community Ecology*
584 *Package*. R package version 2.0-7

585 Oksuz DP (2020) Taxonomic and functional diversity patterns of multi-taxa in
586 Mediterranean wood-pastures. Lisbon University

587 Oksuz DP, Aguiar CAS, Tápia S, et al (2020) Increasing biodiversity in wood-pastures
588 by protecting small shrubby patches. For Ecol Manage 464:118041.
589 <https://doi.org/10.1016/j.foreco.2020.118041>

590 Olea L, San Miguel-Ayanz A (2006) The Spanish dehesa. A traditional Mediterranean
591 silvopastoral system linking production and nature conservation. In: 21st General
592 Meeting of the European Grassland Federation. pp 3–13

593 Oliver TH, Morecroft MD (2014) Interactions between climate change and land use
594 change on biodiversity: attribution problems, risks, and opportunities. Wiley
595 Interdiscip Rev Clim Chang 5:317–335. <https://doi.org/10.1002/wcc.271>

596 Ottaviani G, Keppel G, Marcantonio M, et al (2019) Woody species in resource-rich
597 microrefugia of granite outcrops display unique functional signatures. Austral Ecol
598 44:575–580. <https://doi.org/10.1111/aec.12745>

599 Paracchini M, Petersen J, Hoogeveen Y, et al (2008) High nature value farmland in
600 Europe: an estimate of the distribution patterns on the basis of land cover and
601 biodiversity data.

602 Pavlidis G, Tsihrintzis VA (2018) Environmental Benefits and Control of Pollution to
603 Surface Water and Groundwater by Agroforestry Systems: a Review. Water
604 Resour Manag 32:1–29. <https://doi.org/10.1007/s11269-017-1805-4>

605 Pereira H, Daily G (2006) Modeling biodiversity dynamics in countryside landscapes.
606 Ecology 87:1877–1885

607 Pereira HM, Ferrier S, Walters M, et al (2013) Essential biodiversity variables. Science
608 339:277–8

609 Pia M V., Renison D, Mangeaud A, et al (2013) Occurrence of top carnivores in relation
610 to land protection status, human settlements and rock outcrops in the high
611 mountains of central Argentina. *J Arid Environ* 91:31–37.
612 <https://doi.org/10.1016/J.JARIDENV.2012.11.004>

613 Pinheiro J, Bates D, DebRoy S, et al (2021) nlme: Linear and Nonlinear Mixed Effects
614 Models. R package version 3.1-152

615 Pinto-Correia T, Ribeiro N, Sá-Sousa P (2011) Introducing the montado, the cork and
616 holm oak agroforestry system of Southern Portugal. *Agrofor Syst* 82:99–104.
617 <https://doi.org/10.1007/s10457-011-9388-1>

618 Plas F van der (2019) Biodiversity and ecosystem functioning in naturally assembled
619 communities. *Biol Rev* 94:1220–1245. <https://doi.org/10.1111/BRV.12499>

620 Plieninger T, Pulido FJ, Schaich H (2004) Effects of land-use and landscape structure
621 on holm oak recruitment and regeneration at farm level in *Quercus ilex* L.
622 dehesas. *J Arid Environ* 57:345–364. [https://doi.org/10.1016/S0140-](https://doi.org/10.1016/S0140-1963(03)00103-4)
623 [1963\(03\)00103-4](https://doi.org/10.1016/S0140-1963(03)00103-4)

624 Podani J, Csányi B (2010) Detecting indicator species: Some extensions of the IndVal
625 measure. *Ecol Indic* 10:1119–1124.
626 <https://doi.org/10.1016/J.ECOLIND.2010.03.010>

627 R Core Team (2021) R: A language and environment for statistical computing.

628 Renjifo LM (2001) Effect of natural and anthropogenic landscape matrices on the
629 abundance of subandean bird species. *Ecol Appl* 11:14–31.
630 [https://doi.org/10.1890/1051-0761\(2001\)011\[0014:EONAAL\]2.0.CO;2](https://doi.org/10.1890/1051-0761(2001)011[0014:EONAAL]2.0.CO;2)

631 Roberts DW (2015) Package “labdsv”. Ordination and Multivariate Analysis for
632 Ecology. <https://cran.r-project.org/web/packages/labdsv/labdsv.pdf>. Accessed 11
633 May 2015

634 Ron R, Fragman-Sapir O, Kadmon R (2018) The role of species pools in determining
635 species diversity in spatially heterogeneous communities. *J Ecol* 106:1023–1032.
636 <https://doi.org/10.1111/1365-2745.12840>

637 Rosalino LM, Rosário J do, Santos-Reis M (2009) The role of habitat patches on
638 mammalian diversity in cork oak agroforestry systems. *Acta Oecologica* 35:507–
639 512. <https://doi.org/10.1016/J.ACTAO.2009.03.006>

640 Skaug H, Fournier D, Nielsen A, et al (2013) Generalized Linear Mixed Models using
641 AD Model Builder. R package version 0.7.5

642 Smithson M, Verkuilen J (2006) A better lemon squeezer? Maximum-likelihood
643 regression with beta-distributed dependent variables. *Psychol Methods* 11:54–71.
644 <https://doi.org/10.1037/1082-989X.11.1.54>

645 SNIRH (2019) SNIRH (Sistema Nacional de Informação de Recursos Hídricos–
646 Portugal). Agência Nacional de Àguas - ANA

647 SPBotanica (2013) Flora on - Flora digital de Portugal. <http://www.flora-on.pt/>.
648 Accessed 23 Dec 2019

649 Speziale KL, Ezcurra C (2015) Rock outcrops as potential biodiversity refugia under
650 climate change in North Patagonia. *Plant Ecol Divers* 8:353–361.
651 <https://doi.org/10.1080/17550874.2014.983200>

652 Tilman D, Clark M, Williams DR, et al (2017) Future threats to biodiversity and
653 pathways to their prevention. *Nature* 546:73–81.
654 <https://doi.org/10.1038/nature22900>

655 Torralba M, Fagerholm N, Burgess PJ, et al (2016) Do European agroforestry systems
656 enhance biodiversity and ecosystem services? A meta-analysis. *Agric Ecosyst*
657 *Environ* 230:150–161. <https://doi.org/10.1016/J.AGEE.2016.06.002>

658 Twidale CR (2000) Granite outcrops: their utilisation and conservation. *J R Soc West*
659 *Aust* 115

660 van der Maarel E (2007) Transformation of cover-abundance values for appropriate
661 numerical treatment - Alternatives to the proposals by Podani. *J Veg Sci* 18:767–
662 770. <https://doi.org/10.1111/j.1654-1103.2007.tb02592.x>

663 Verchot L V., Van Noordwijk M, Kandji S, et al (2007) Climate change: linking
664 adaptation and mitigation through agroforestry. *Mitig Adapt Strateg Glob Chang*
665 12:901–918. <https://doi.org/10.1007/s11027-007-9105-6>

666 Villéger S, Mason NWH, Mouillot D (2008) New multidimensional functional diversity
667 indices for a multifaceted framework in functional ecology. *Ecology* 9:2290–2301

668 Vittoz P, Engler R (2007) Seed dispersal distances: A typology based on dispersal
669 modes and plant traits. *Bot Helv* 117:109–124. [https://doi.org/10.1007/s00035-](https://doi.org/10.1007/s00035-007-0797-8)
670 [007-0797-8](https://doi.org/10.1007/s00035-007-0797-8)

671 Wintle BA, Kujala H, Whitehead A, et al (2019) Global synthesis of conservation
672 studies reveals the importance of small habitat patches for biodiversity. *Proc Natl*
673 *Acad Sci* 116:909–914. <https://doi.org/10.1073/PNAS.1813051115>

674 Wood SN (2006) *Generalized Additive Models: An Introduction with R*. Chapman and
675 Hall/CRC, Boca Raton, Florida

676 Wright SJ, Bunker D, Dalling J, et al (2006) Towards a functional trait based research
677 program within the Center for Tropical Forest Science

678 Zuur AF, Ieno EN, Elphick CS (2010) A protocol for data exploration to avoid common
679 statistical problems. *Methods Ecol Evol* 1:3–14. [https://doi.org/10.1111/j.2041-](https://doi.org/10.1111/j.2041-210X.2009.00001.x)
680 [210X.2009.00001.x](https://doi.org/10.1111/j.2041-210X.2009.00001.x)

681

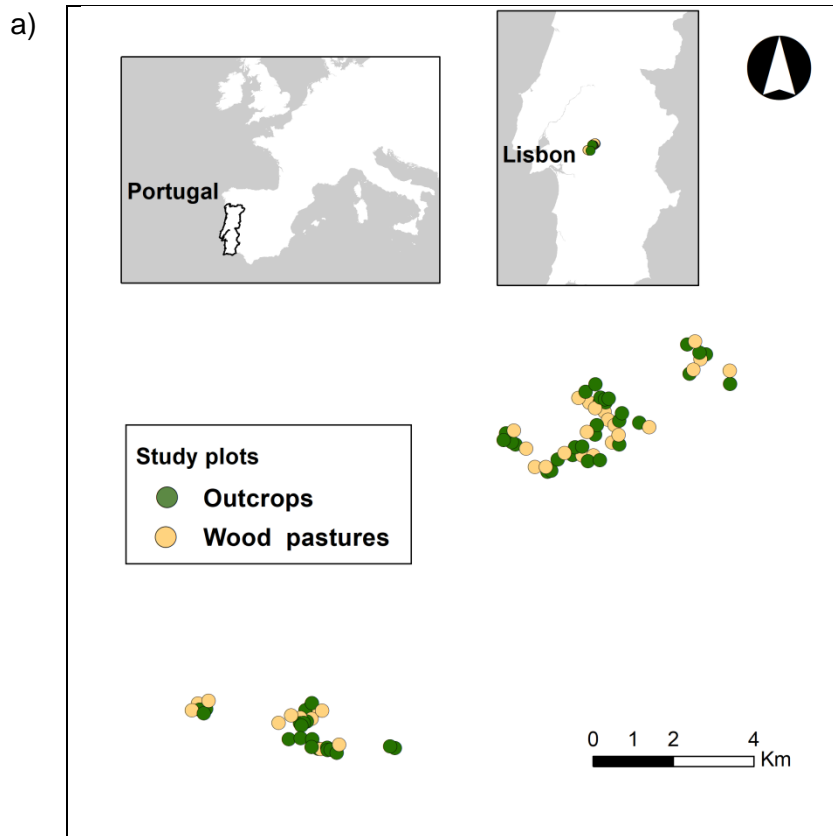
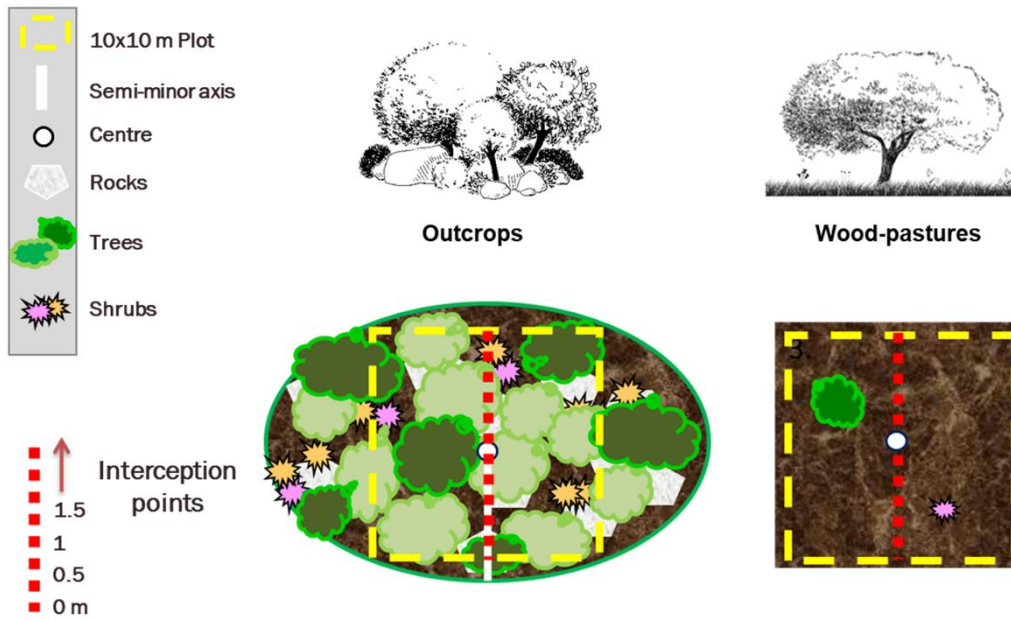


Figure 1. a) 102 10x10 m plots, 51 located in rocky outcrops (32 for analyses comparing outcrops and wood-pasture communities, and 19 more for analysing the effect of size and spatial distribution of outcrops) and 51 (32 + 19) in wood-pastures, were studied in Alentejo, Portugal, examples of b) outcrop and c) wood-pasture plots sampled.

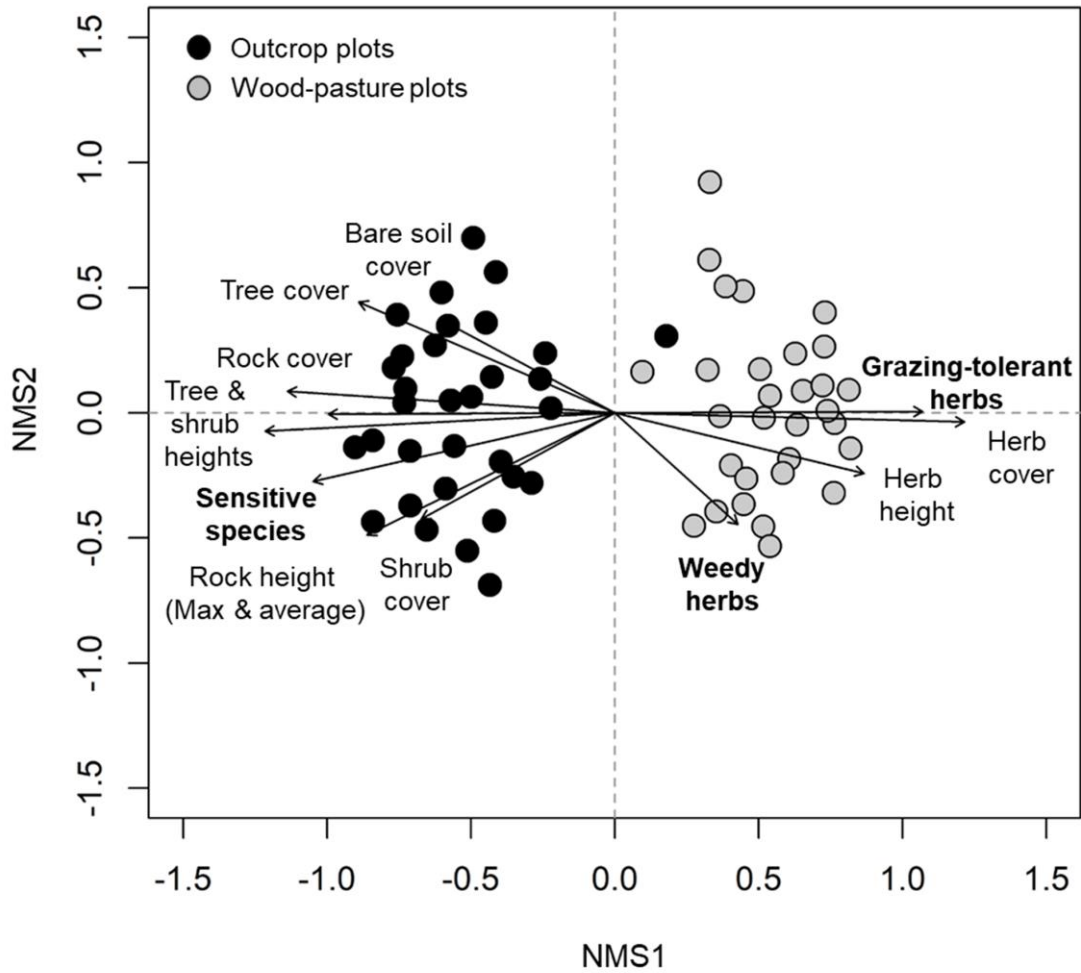




684

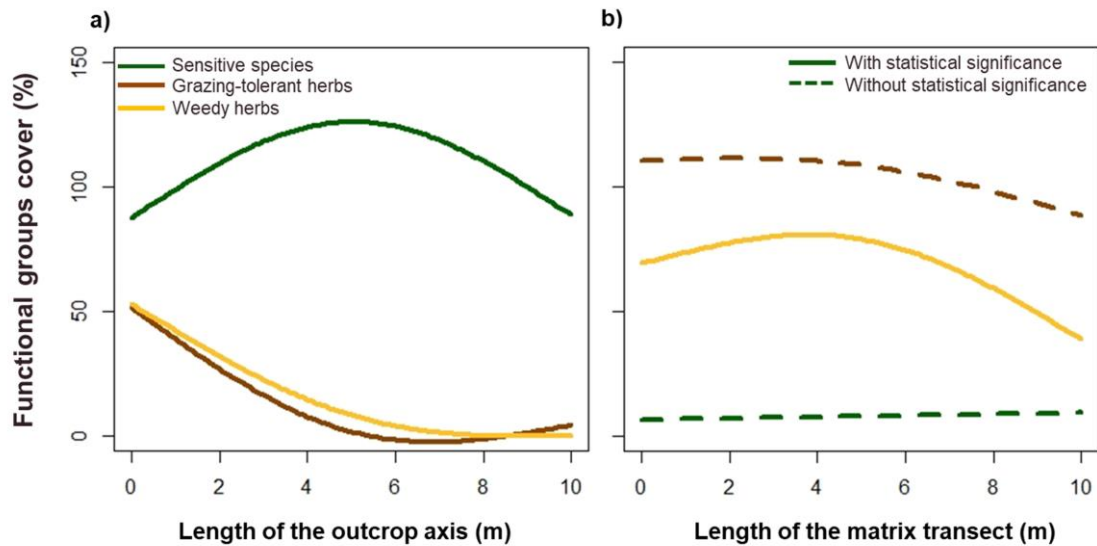
685 **Figure 2.** Sampling design. Cover of plants, rocks, mosses, lichens, bare soil and litter was
 686 registered in the studied 10x10 m plots, first in the whole plot area (n=64: 32 outcrops + 32 wood-
 687 pastures) and then, using the point-intercept method, in the outcrop (n= 51) and wood-pastures
 688 (n=51) axes (refer to material and methods for details).

689



690

691 **Figure 3.** Axes 1 and 2 of the 2-dimensional nonmetric multidimensional scaling ordination (NMS)
 692 of study plots based on plant species cover (final stress 21.84%). Vectors represent significant
 693 correlations between species composition and i) the environmental characteristics separating
 694 outcrop and matrix communities and ii) the plant functional groups identified (in bold).



695

696 **Figure 4.** Relationships between a) the outcrop axis length (from edge to centre) and
 697 b) matrix transect length and the cover of functional groups, namely sensitive species
 698 and grazing-tolerant and weedy herbs. Lines represent the main trend of a generalised
 699 additive model (GAM). Percentage of variance explained: a) 59.4%^{***}, 80.3%^{***} and
 700 97.6%^{***}, respectively; Effective degrees of freedom (k): 1.96, 1.95 and 1.98,
 701 respectively; b) Percentage of variance explained 63.5%^{***} and Effective degrees of
 702 freedom (k): 1.93.

703

704 **Tables**

705 Table 1. Pair-wise Wilcoxon test summary on the differences between outcrops and
 706 wood-pastures for variables and indices regarding species and functional diversities.
 707 Values ar mean and standard deviation (SD) for each variable and index and the Statistic
 708 value, number of pairs (n) and *p*-value returned by the tests.

	Outcrops	Wood-pastures	Pair-wise Wilcoxon tests		
	Mean (SD)	Mean (SD)	Statistic	n	p
<u>Species diversity</u>					
Richness	26.25 (7.49)	28.59 (9.00)	322	32	ns
Herbaceous Richness	19.94 (8.68)	27.81 (6.85)	424	32	0.003
Shrubs Richness	3.62 (1.66)	0.37 (0.87)	0	32	7.33x10 ⁻⁷
Climbers Richness	1.15 (1.08)	0 (0)	0	32	8.09x10 ⁻⁶
Trees Richness	1.5 (0.88)	0.4 (0.61)	19	32	2.32x10 ⁻⁵
Holm oak cover – tree (%)	12.98 (21.26)	1.01 (2.76)	5.5	32	4.62x10 ⁻⁴
Holm oak cover – shrub (%)	2.75 (10.96)	0.47 (2.65)	16	32	0.02
Olive tree cover – tree (%)	19.14 (28.57)	0 (0)	0	32	6.81x10 ⁻⁴
Olive tree cover – shrub (%)	0.7 (2.71)	0 (0)	0	32	ns
Kermes oak cover - tree (%)	4.84 (16.81)	0 (0)	0	32	ns
Cork oak cover – tree (%)	5.94 (18.49)	2.97 (11.47)	15	32	ns
Cork oak cover – shrub (%)	0 (0)	0.003 (0.018)	1	32	ns
<u>Functional diversity</u>					
Functional richness	0.08 (0.02)	0.05 (0.02)	97	32	0.001
Functional evenness	0.55 (0.08)	0.48 (0.08)	105	32	0.002
Functional divergence	0.90 (0.10)	0.81 (0.13)	120	32	0.006
Rao’s quadratic entropy	0.064 (0.03)	0.046 (0.02)	140	32	0.019
Functional dissimilarity	0.20 (0.08)	0.19 (0.07)	208	32	ns
Sensitive species	78.54 (35.46)	7.31 (18.36)	5	32	1.34*10 ⁻⁶
Grazing-tolerant herbs	5.42 (7.09)	85.71 (40.89)	528	32	4.66*10 ⁻¹⁰
Weedy herbs	8.98 (11.47)	46.57 (54.10)	455	32	1.62*10 ⁻⁴

709

710 Table 2 Generalised Linear Mixed Models (GLMMs) results testing the influence of the
 711 outcrop size, the cover and height of vegetation and rocks, and the farmsteads (random
 712 effect) on the outcrop species and functional diversities. Values correspond to the
 713 estimate, standard error (Std. Error), t value, p-value and the marginal coefficient of
 714 determination (R^2_m) returned by the models and the sample size (n).

	Estimate	Std. Error	t value	p-value	R^2_m	n
Log (Plant richness)						
Intercept	0.921	0.204	4.508	0.000		
Length of the outcrop axis	0.061	0.028	2.198	0.033	0.37	51
Plant cover	0.004	0.001	5.041	0.000		
Sqrt (Shannon index)						
(Intercept)	1.255	0.054	23.441	0		
Rock cover	0.001	0.000	2.025	0.050	0.22	51
Herb cover	-0.002	0.001	-2.969	0.005		
Simpson index						
(Intercept)	0.379	0.049	7.772	<0.001		
Tree cover	0.001	0.000	2.137	0.038	0.18	51
Plant cover	-0.001	0.000	-2.985	0.004		
Plant cover						
(Intercept)	204.271	9.907	20.618	<0.001		
Rock cover	-0.984	0.240	-4.095	0.000	0.29	51
Bare soil	-1.348	0.514	-2.621	0.012		
Sensitive species cover						
(Intercept)	91.636	11.733	7.810	0		
Rock height	18.088	7.414	2.440	0.019	0.11	51
Grazing tolerant herb cover						
(Intercept)	-1.897	0.486	-3.900	0.000		
Richness	0.196	0.051	3.850	0.000	0.57	51
Maximum rock height	-0.504	0.164	-3.080	0.002		
Weedy herb cover						
(Intercept)	11.873	6.838	1.736	0.089		
Rock height	-8.208	3.210	-2.558	0.014	0.49	51
Herb cover	0.387	0.067	5.739	0.000		

715

716 **Table 3**

717 Influence of farmsteads (random effect) on plant cover and taxonomical and functional
 718 diversities of outcrops as described by the models' log-likelihood ratio test (LRT). Values
 719 correspond to the log-likelihood value (log-Lik), *p* value and sampling size (n). In bold *p*
 720 < 0.05. When the impact of farmsteads was significant, the conditional coefficient of
 721 determination (R^2c) returned by the models was calculated.

	log-Lik	<i>p</i>-value	n	R^2c
Log (Plant richness)	-16.819	0.513	51	
Sqrt (Shannon index)	18.14	0.315	51	
Simpson index	33.029	0.385	51	
Plant cover	-250.930	0.003	51	0.61
Sensitive species cover	-254.434	0.225	51	
Grazing tolerant herb cover	45.366	0.902	51	
Weedy herb cover	-217.2243	0.039	51	0.64

722