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## Effect of landscape connectivity on plant communities a review of response patterns

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1 **Title:**

2 Effect of landscape connectivity on plant communities: a review of response patterns

3

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14 **Abstract:**

15 Context. Fragmentation in agricultural landscapes is considered as a major threat to  
16 biodiversity. Thus, ecological corridors are deployed at multiple scales to increase connectivity.

17 However, there is limited consensus about their efficiency, especially for plants.

18 Objectives. We assimilated existing knowledge to assess whether and how landscape  
19 connectivity impacts plant communities.

20 Methods. We reviewed published literature across more than 20 years, providing an overview  
21 on the influence of connectivity on plant communities.

22 Results. We found that landscape connectivity has a varying and complex influence on the  
23 composition and diversity of plant communities (*i.e.* community taxonomic structure), due to  
24 the multiplicity of factors that modulate its effect. Our understanding of how of landscape  
25 connectivity impacts the dispersal of plants is improved by using biological traits (*i.e.*  
26 community functional structure). Finally, we showed that landscape connectivity promotes  
27 actual dispersal between connected communities.

28 Conclusions. This review emphasises the pertinence of trait-based and actual dispersal  
29 approaches to improve our understanding and ability to predict the effect of connectivity loss  
30 on plant communities, allowing us to identify new prospects for future research.

31

32 **Keywords:** plant communities, traits, seed dispersal, actual dispersal, ecological corridors

### 33 **Introduction**

34 Anthropogenic activities, including urbanization and agricultural intensification, present major  
35 threats to biodiversity, notably through their effects on landscape fragmentation (Stoate et al.  
36 2001; Kato et al. 2009; Haddad et al. 2015). Landscape fragmentation impacts the populations  
37 of plants through two distinct effects: habitat loss (*i.e.* a decrease in habitat area) and  
38 connectivity loss (*i.e.* decreased plant dispersal among remnant patches; Taylor et al. 1993)  
39 (Fahrig 2003; Fischer and Lindenmayer 2007). Because of their sessile way of life, plants are  
40 particularly vulnerable to these effects, and many studies have reported a strong decline in  
41 floristic diversity in fragmented agricultural landscapes (Andreasen et al. 1996; Kleijn and  
42 Verbeek 2000; Luoto et al. 2003; Baessler and Klotz 2006; Kleijn et al. 2009). The effect of  
43 habitat area on plant diversity is well established (*e.g.* Scanlan 1981; Dzwonko and Loster 1988;  
44 Zacharias and Brandes 1990; Honnay et al. 1999; Godefroid and Koedam 2003; Gignac and  
45 Dale 2007). Small patch size increases extinction risk by reducing carrying capacity.  
46 Consequently, populations in smaller habitats are more susceptible to extinction, due to  
47 demographic stochasticity (MacArthur and Wilson 1967). Only in the last two decades have  
48 studies begun investigating how connectivity loss affects plant diversity and the corresponding  
49 mechanisms; consequently, there is a low level of synthesis of existing knowledge.

50         Depending on the intensity of connectivity loss, the spatial distribution of species might  
51 range from a continuous population to a set of isolated populations with a metapopulation  
52 structure (*i.e.* a network of local populations linked by dispersal fluxes, Hanski 1994) as an  
53 intermediate case. More recently, the metapopulation concept has been extended to the  
54 community level (*i.e.* metacommunity concept, Leibold et al. 2004). A metacommunity refers  
55 to a set of local communities that are linked by the dispersal of multiple potentially interacting  
56 species (Wilson 1992). The strong interest in metacommunity theory has given rise to four  
57 paradigms to describe metacommunities: i) species sorting (*i.e.* habitat patches are

58 environmentally heterogeneous with high enough dispersal to enable species to fill niches  
59 within habitat patches because of niche diversification); ii) mass effects (*i.e.* habitat patches are  
60 environmentally heterogeneous and high dispersal enough override local dynamics); (iii) patch  
61 dynamics (*i.e.* habitat patches are environmentally homogeneous and species differ in their  
62 ability to disperse. Along a colonisation–competition trade-off, successful colonisers  
63 outcompete poor competitors and *vice versa*); and iv) neutral models (*i.e.* species do not differ  
64 in their fitness or niche) in the case of ecological drift (Leibold et al. 2004; Logue et al. 2011).  
65 These models formulate different hypotheses on the relative importance of dispersal, habitat  
66 heterogeneity and interactions between species in the structuring mechanisms of local  
67 communities (Leibold et al. 2004; Logue et al. 2011). Plant disperse passively between habitat  
68 patches via pollen and/or seeds. However, this process is only successful if habitat patches are  
69 sufficiently connected (Fahrig and Merriam 1985; Bowne and Bowers 2004); in other words,  
70 plant dispersal among source patches must be facilitated by suitable landscape elements (Taylor  
71 et al. 1993). Landscape connectivity might be promoted by the occurrence of corridors between  
72 favourable patches, such as continuous, relatively narrow, habitat bands connecting two patches  
73 (Burel and Baudry 1999). However, connectivity might also include nonlinear or discontinuous  
74 landscape features that are permeable to plant dispersal (Beier and Noss 1998; Chetkiewicz et  
75 al. 2006). Thus, high connectivity between habitat patches might reduce the adverse effects of  
76 fragmentation by facilitating genetic fluxes among local populations through plant dispersal  
77 (Wilson and Willis 1975), which contrasts with animals for which all movements is concerned.

78         However, the ability of plants to disperse in fragmented landscapes might depend on  
79 their dispersal strategy, with some being more or less able to promote frequent long-distance  
80 dispersal events. For instance, if wind-dispersal or water-dispersal is generally achieved over a  
81 long distance, depending on the intensity of the wind or current and the adaptations of the seeds,  
82 gravity-dispersed plants (*i.e.* barochory) are dispersed more locally (Nathan et al. 2002;

83 Tackenberg 2003; Vittoz and Engler 2007; Pollux et al. 2009; Nilsson et al. 2010). Animal-  
84 dispersed plants are dispersed over short distances of <100 m (*e.g.* by small animals, such as  
85 insects, small mammals and some bird species) to long distances of several kilometres (*e.g.* by  
86 large or migratory animals; Vittoz and Engler 2007). These vectors might be influenced  
87 differently by landscape elements and, in turn, might influence plant responses to connectivity  
88 loss. Within a certain type of dispersal mode, plant species might represent a large range of  
89 capacities to disperse with respect to quantity (*i.e.* number of potential dispersers) and distance  
90 (*i.e.* dispersal range) (Johst et al. 2002; Nathan 2006; Auffret et al. 2017), which might decrease  
91 or increase their chance to disperse. These responses involve a selection of particular biological  
92 traits (*i.e.* morphological, biochemical, physiological, structural, phenological or behavioural  
93 characteristics of organisms that influence performance or fitness, Violle et al. 2007). At the  
94 plant community scale, analyses of how these traits respond to connectivity might help us to  
95 understand the processes that are involved, which would facilitate a more mechanistic analysis  
96 of plant assembly in fragmented landscapes.

97         Because plants are sessile, the effect of connectivity on plant assemblages might be  
98 time-dependent, with responses reflecting a relaxation time (*i.e.* the time taken for a community  
99 of species to reach a new equilibrium after an environmental disturbance, Diamond 1972;  
100 Kuussaari et al. 2009). By extension, the response of plant to connectivity might also reflect an  
101 extinction debt, which is defined as the number or proportion of extant species predicted to  
102 become extinct as the species community reaches a new equilibrium after an environmental  
103 disturbance (Tilman et al. 1994; Ovaskainen and Hanski 2002; Kuussaari et al. 2009). The  
104 response of plant assemblages to connectivity might also reflect an immigration lag, *i.e.* the  
105 time that has elapsed between an immigration-committing forcing event (*e.g.* increasing  
106 connectivity) and the establishment of an immigrating species (Jackson and Sax 2010). By  
107 extension, this phenomenon would lead to immigration credit (Jackson and Sax 2010), which

108 is defined as the number of species committed to eventual immigration following a forcing  
109 event (Kuussaari et al. 2009). In addition, temporal lags in extinction and immigration are  
110 mediated by variation in traits across species under experimental situations (Haddad et al.  
111 2015). Thus, considering the role of historical connectivity on the composition and diversity of  
112 plant communities and how traits mediate the delayed response of plant might improve our  
113 understanding of how plant species respond to connectivity loss.

114 Metacommunity theory recognizes that established local communities may additionally  
115 be driven by stochasticity (which is eventually spatially correlated), local patch conditions and  
116 species traits (*i.e.* colonisation vs. competition trade-offs) (Mouquet and Loreau 2003). As a  
117 result, recent studies argued that assessing how connectivity affects established communities in  
118 habitat patches is subject to bias. Indeed, once a propagule arrives in a patch, its ability to  
119 establish depends on a large range of local factors linked to local habitat quality (*e.g.*  
120 management, land-use history, Brudvig 2016) and biotic interactions (*e.g.* plant-plant, plant-  
121 animal, plant-soil interactions, Archer and Pyke 1991; Pyke and Archer 1991; Fukami and  
122 Nakajima 2013). Therefore, it might be difficult to disentangle the effects of landscape  
123 connectivity and local filters. In addition, the increase of connectivity might indirectly affect  
124 the establishment and coexistence of plants in the community by promoting the displacement  
125 of: i) organisms that might predate on (*e.g.* herbivores and granivorous species, Orrock et al.  
126 2003; Orrock and Damschen 2005; Rossetti et al. 2017) or parasitize seeds (*e.g.* Sullivan et al.  
127 2011), or ii) strong competitors that once, they arrived in a local patch, drive local biotic  
128 interactions and outcompete less competitive species (Simberloff and Cox 1987; Mouquet and  
129 Loreau 2003, *e.g.* exotic invasive plant species, Minor et al. 2009; Minor and Gardner 2011).  
130 To assess the direct effect of connectivity on seed dispersal, Calabrese and Fagan (2004)  
131 proposed the concept of actual functional connectivity (Fig. 1 and Box 1). This concept is based  
132 on the direct assessment of seed fluxes depending on habitat connectivity. This form of direct

133 assessment might reflect the effect of connectivity on the process of dispersal alone,  
134 independent of other local factors that are difficult to standardise.

135         The number of studies analysing the influence of landscape connectivity on biodiversity  
136 has increased considerably since the early part of the 21st century (Ayram et al. 2016). To  
137 facilitate the identification of general patterns, reviews have been compiled on how connectivity  
138 affects global biodiversity (Murphy and Lovett-Doust 2004; Haddad and Tewksbury 2006;  
139 Gilbert-Norton et al. 2010; Haddad et al. 2011, 2015; Fletcher et al. 2016). These reviews have  
140 provided evidences of the effect of connectivity at different levels of organization (individuals,  
141 populations and communities) essentially through experimental approaches (Haddad and  
142 Tewksbury 2006; Haddad et al. 2011, 2015) or sometimes coupled with correlatives ones  
143 (Murphy and Lovett-Doust 2004; Gilbert-Norton et al. 2010; Fletcher et al. 2016). However,  
144 these reviews mostly focus on animal biodiversity. When these reviews consider existing  
145 literature on plants, the response of plants to connectivity is usually often assessed at the  
146 population level and under experimental conditions and does not take into account the traits of  
147 plant species (Murphy and Lovett-Doust 2004; Gilbert-Norton et al. 2010; Haddad et al. 2011,  
148 2015; Fletcher et al. 2016). As a result, our understanding of the role of connectivity for plants  
149 remains limited, especially at the community level. The effect of connectivity on plant  
150 communities cannot be predicted from its effect on plant populations and individuals, as  
151 communities incorporate distinct properties linked to their dynamics such as priority effects  
152 (Fukami 2015) and successional changes in species diversity over time. This lack of knowledge  
153 may be particularly problematic for the conservation of plant biodiversity, which aims to go  
154 beyond single-species approaches (Groves et al. 2002).

155         In the present review, we aimed to provide a synthesis of existing scientific knowledge  
156 on how connectivity loss influences plant communities. To achieve this synthesis, we only  
157 considered studies that assessed connectivity and plant species responses at the community



158 level. Connectivity was analysed under experimental approaches through the presence or  
159 absence of an experimentally-created corridor between patches (Fig. 1 and Box 1) or under  
160 correlative approaches using quantitative indices. Quantitative connectivity assessments were  
161 based on isolation metrics such as Euclidean distance, structural connectivity indices (see  
162 various examples in the main text), least-cost distance and resistance distance (respectively  
163 derived from graph and circuit theory, potential functional connectivity) (Fig. 1 and Box 1).  
164 This approach included: i) analysis of individual species responses across most species of the  
165 community and, ii) aggregated measures across most species of a given community (for more  
166 details, see Appendix S1).

167 Specifically, we present how landscape connectivity influences the richness and  
168 composition of plant communities (section 1). We also review how plants respond to  
169 connectivity from a mechanistic perspective (related to dispersal vector mode and plant  
170 dispersal traits), to demonstrate that traits may help to define species sensitivity to connectivity  
171 loss (section 2). We also evaluate how time affects plant taxonomic and functional response to  
172 connectivity (section 3). Finally, we review how connectivity drives actual dispersal (section  
173 4). We conclude that landscape connectivity influences plant communities as a filter for species  
174 and traits. We also emphasize that trait-based and actual dispersal approaches are the most  
175 relevant for predicting how connectivity loss affects plant communities. We use our findings to  
176 suggest further research needs.

177

### 178 **Contrasted effect of landscape connectivity on plant community taxonomic structure**

179 Higher connectivity facilitates plant dispersal and thus decreases the probability of local  
180 extinction. Thus, high connectivity might influence the taxonomic structure of communities  
181 (*i.e.* the composition and diversity of plant communities), increasing species diversity and  
182 similarity in composition between connected patches.

183

184 *Plant community diversity*

185 How connectivity affects species diversity has been a central question in landscape ecology  
186 over the last two decades. Early studies tested this question by comparing pairs of patches that  
187 were (or not) connected through a corridor. The effect of the presence/absence of a corridor on  
188 plant diversity has been, for instance, addressed through the experimental manipulation of  
189 connectivity between habitat patches. One of the most well-known designs is based at Savannah  
190 River Site (South Carolina, USA), which supports a range of landscape designs embedded in a  
191 forest matrix. In this experimental design, each landscape is composed of a central patch that is  
192 connected to one peripheral patch by a corridor, and a peripheral patch that is not connected to  
193 the central patch (Box 2). In this design, patches connected by a corridor displayed a richness  
194 higher than 20% compared to the non-connected patches (Damschen et al. 2006). Plant species  
195 richness also increases from 10 to 18% in the matrix surrounding the connected patches by a  
196 mass spatial effect (Brudvig et al. 2009); thus, the presence of corridors might positively impact  
197 adjacent landscape elements.

198 Another set of studies used connectivity measurements based on the Euclidean distance  
199 or on the permeability of the landscape elements to dispersal. In the latter case, these  
200 measurements incorporate the physical attributes of the landscape with or without information  
201 about the organism of interest (structural or potential functional connectivity) and assess the  
202 degree of connectivity for each habitat patch (Fig. 1 and Box 1). Depending on the ecosystem  
203 type, landscape connectivity has a contrasting effect on community diversity. For instance, local  
204 heathland species richness increases with structural connectivity [measured through the  
205 Incidence Function Model index (IFM index) based on the area and the Euclidean distance to  
206 other patches of similar habitat type, Hanski 1994; Moilanen and Nieminen 2002)] (Piessens et  
207 al. 2004), whereas structural connectivity does not influence the richness and the density of

208 species (*i.e.* average number of species per square meter) of local plant communities in semi-  
209 natural grasslands (Lindborg and Eriksson 2004; Helm et al. 2006; Cousins et al. 2007).  
210 Lindborg and Eriksson (2004) demonstrated that this absence of a relationship with  
211 connectivity was maintained at different spatial scales (1 km or 2 km radius). Only one study  
212 demonstrated a negative influence of connectivity on local plant community richness in riverine  
213 wetlands. Specifically, Bornette et al. (1998) reported that high connectivity decreases species  
214 richness when assessing structural surficial connectivity between cut-off channels and the river  
215 by quantifying the overflows and backflows of rivers into cut-off channels. Under these  
216 conditions, high connectivity impedes recruitment due to over-frequent flood scouring or due  
217 to nutrient-rich and turbid surficial waters being supplied to the wetland, which reduces species  
218 richness.

219 Most studies investigating how landscape connectivity affects plant species  
220 communities measure species richness at the alpha-scale (*i.e.* local diversity) (Bennett et al.  
221 2006). In comparison, studies carried out at a gamma-scale (*i.e.* landscape diversity) remain  
222 limited. For instance, Favre-Bac et al. (2014) reported that the structural connectivity of ditch  
223 networks (*i.e.* number of disconnected network units) decreases species richness of ditch bank  
224 communities when analysing 27 square sites of 500 x 500 m. This result might be attributed to  
225 the influence of landscape heterogeneity, which might mitigate the negative effects of habitat  
226 fragmentation (Tschardt et al. 2012).

227 Overall, no general pattern has been detected on the positive effect of corridors on plant  
228 diversity through correlative approaches, contrary to experimental approaches (Haddad et al.  
229 2015). Two possible explanations why correlative studies have failed to demonstrate a general  
230 pattern are: i) the too low suitability and number of available habitat patches for plants to  
231 colonise and establish, and ii) the lack of consideration of the quality (*e.g.* management  
232 practice) of the landscape elements. These studies assessed connectivity through the presence

233 and identity of landscape elements to determine their permeability to dispersal, overlooking the  
234 potential effect of the quality of these elements on their permeability.

235

### 236 *Plant community composition*

237 In fragmented landscapes, remnant habitat patches might be highly heterogeneous, causing  
238 strong divergence in the composition of local communities (Tscharntke et al. 2012). In addition,  
239 the amount of dispersal between remnant patches might strongly impact these dynamics  
240 (Mouquet and Loreau 2003; Mouquet et al. 2006). When isolated, seed limitation is expected  
241 to enhance divergence between local communities at the metacommunity scale, further  
242 reducing their similarity (Mouquet and Loreau 2003; Tscharntke et al. 2012). A particular case  
243 of dissimilarity in species composition, the nested subset (*i.e.* a community in which the species  
244 present at species-poor sites constitute subsets of those from species-rich sites, Patterson and  
245 Atmar 1986) was highlighted in heathland communities by Piessens et al. (2004). In these  
246 systems, dissimilarity increases as structural connectivity decreases. The higher the landscape  
247 structural connectivity, the higher the similarity of the community. Similar results were also  
248 reported for communities of linear landscape elements, such as field margins, road verges and  
249 ditches (Thiele et al. 2017). For instance, Thiele et al. (2017) tested three types of connectivity  
250 metrics (Euclidian distance, potential functional connectivity: least-cost distance and resistance  
251 distance, see Fig. 1 and Box 1.), and demonstrated that the results were independent of the way  
252 that connectivity was evaluated.

253         Within local communities, fragmentation theoretically decreases species richness  
254 because it acts as a strong ecological filter, selecting for species that are able to disperse and  
255 survive in isolated patches. Thus, Haddad and Tewksbury (2006) theorised that specialist  
256 species should be more affected by connectivity loss than generalist species, because the latter  
257 group might perceive corridors as being of lower quality compared to the surrounding habitats.

258 Isolated patches might contain more generalist than specialist species in plant communities  
259 compared to connected patches. In grassland communities, specialist species decrease to a  
260 greater extent compared to generalist species in response to the loss of structural connectivity  
261 (Adriaens et al. 2006; Brückmann et al. 2010; Evju et al. 2015). Adriaens et al. (2006) reported  
262 that only specialist species are impacted by a change to the structural connectivity (assessed  
263 with the IFM index) of grasslands, with the mean number of species per site decreasing by  
264 about 37%. Similar results were obtained by Brückmann et al. (2010) and Evju et al. (2015).  
265 For instance, Brückmann et al. (2010) recorded a decrease of 24-37% in the richness of  
266 specialist species in grasslands in response to a total loss of structural connectivity (IFM index),  
267 whereas generalists were not affected. Therefore, the effect of connectivity on plant species  
268 depends on the degree of specialisation. Thus, different types of ecological corridors might  
269 interact in a landscape. For instance, grassland corridors between forest patches might reinforce  
270 the effect of forest corridors in connected forest patches only when considering generalist  
271 species in forests. These interactions between corridor types might rank from potential  
272 complementarity for generalist species to potential antagonism for specialist species, in which  
273 one ecological corridor type is favourable while the other constitutes a barrier. Thus, it is  
274 important to consider how complementary between the different types of ecological corridors  
275 are in studies linking connectivity to plant communities. This consideration could be coupled  
276 with the improved modelling of ecological corridor types through the use of SDMs (Species  
277 Distribution Models; Guisan and Thuiller 2005), which represent a promising approach to  
278 assess the permeability of landscape features to dispersal (*e.g.* Morato et al. 2014), rather than  
279 expert knowledge (Rayfield et al. 2010; Spear et al. 2010; Zeller et al. 2012; Koen et al. 2012).  
280  
281 **Effect of landscape connectivity on the functional structure of plant communities**

282 Because of the contrasting results provided by taxonomic approaches, there is a growing need  
283 to predict how plant communities respond to changes in connectivity. Analyses based on the  
284 functional structure of communities (*i.e.* the composition and diversity of traits) might help to  
285 improve our understanding of species responses by providing a mechanistic explanation (for a  
286 detailed synthesis of these studies, see Table 1).

287

### 288 *Dispersal vector mode*

289 Studies using trait-based approaches initially focused on how plant dispersal vectors influence  
290 their responses to connectivity. Plant dispersal vectors determine plant dispersal distance. For  
291 instance, dispersal is close to the parent plant for gravity-dispersed species, and up to several  
292 hundred metres or even kilometres for animal-, wind- and water- dispersed species (Sorensen  
293 1986; Nathan et al. 2002; Tackenberg 2003; Vittoz and Engler 2007; Pollux et al. 2009; Nilsson  
294 et al. 2010).

295 Animal-dispersal is oriented, and depends on animal behaviour and movement in the  
296 landscape. There is broad consensus that animal movement is facilitated by landscape  
297 connectivity (Murphy and Lovett-Doust 2004; Haddad and Tewksbury 2006; Gilbert-Norton et  
298 al. 2010; Haddad et al. 2011, 2015; Fletcher et al. 2016); thus, animal-dispersed plant species  
299 might be largely dependent on landscape connectivity. It is more difficult to predict how  
300 connectivity affects wind-dispersed plants, as this type of dispersal is made at random or occurs  
301 in relation to the dominant wind direction. Differences in these two plant dispersal vectors  
302 might lead to different responses to connectivity. Damschen et al. (2008) surveyed the species  
303 richness of bird-dispersed and wind-dispersed species over a seven-year period in the Savannah  
304 River Site (Box 2). The authors demonstrated that, for both functional groups, the richness of  
305 communities was lower in unconnected versus connected patches. However, this difference in  
306 richness reached an asymptote for bird-dispersed species after five years, but not for wind-

307 dispersed species. This asymptote is due to the behaviour of birds that regularly visit all patches  
308 of the experimental design. As a result, even isolated patches occasionally receive new  
309 colonists, reducing the difference in species richness between connected and unconnected  
310 patches over time. By contrast, the presence of an open corridor affects wind dynamics by  
311 redirecting and promoting airflow and “ejection hotspots” (*i.e.* locations in which seeds have a  
312 relatively high probability of being uplifted) in connected patches. These processes increase the  
313 likelihood of seed uplift, and, therefore, the likelihood of seed redirection and bellow among  
314 connected patches (Damschen et al. 2014). As a result, the difference in species richness  
315 between connected and unconnected patches continues to increase over time. Based on the same  
316 design, Brudvig et al. (2009) showed that the richness of animal-dispersed species increased by  
317 13–19% per 100 m<sup>2</sup> in the surrounding matrix in response to connectivity, whereas no mass  
318 spatial effect was found for wind-dispersed species. This result was due to a barrier effect of  
319 the surrounding matrix of dense pine plantations on wind-dispersed species (Damschen et al.  
320 2014). In contrast, animals were not restricted to the experimental design, moving in and out of  
321 patches to the matrix, resulting in their contributing to the dispersal of plants. When assessed  
322 in more correlative approaches, no particular influence of landscape connectivity (Euclidean  
323 distance, structural and potential functional connectivity) was demonstrated on animal-  
324 (external, internal and by ants) and wind-dispersed species (Piessens et al. 2005; Lindborg  
325 2007; Evju et al. 2015; Thiele et al. 2017, but see contrasting results provided by Verheyen et  
326 al. (2004) who only considered vertebrate-animal dispersal and wind dispersal), despite their  
327 potential for long-distance dispersal. This finding suggests that, in real systems, the effect of  
328 connectivity on the functional structure of plant communities is overshadowed by other  
329 characteristics of the landscape, such as patch shape, which influences the quantity of the  
330 interface with other habitats (*e.g.* Arellano-Rivas et al. 2016) or the fine-scale structuring of

331 corridors. In turn, this phenomenon influences the degree of permeability for animal- (*e.g.*  
332 animal vector behavioural changes) and wind- (*e.g.* wind dynamics changes) dispersed species.

333 Another dispersal mode contributing to long-distance dispersal is hydrochory. Many  
334 studies have investigated the role of water dispersal in structuring plant communities, in both  
335 flowing and stagnant systems (for a review, see Nilsson et al. 2010). However, very few studies  
336 present the effect of connectivity on water-dispersed plant communities. Favre-Bac et al. (2014)  
337 showed that the richness of water-dispersed species at the landscape scale was independent of  
338 ditch network connectivity, regardless of the measures of connectivity considered (*i.e.* total  
339 ditch length, number of culverts, number of intersections and number of disconnected ditch  
340 network subunits). However, similarity within local water-dispersed communities decreased  
341 with the number of intersections, suggesting that these intersections act as barriers to dispersal  
342 and cause a decrease of seed fluxes between communities (Favre-Bac et al. 2014). Similar  
343 results were obtained by Piessens et al. (2005).

344 Other modes have been less studied, mostly because they were supposed to contribute  
345 less to long-distance dispersal and, therefore, to plant sensitivity to connectivity. Nonetheless,  
346 several studies have demonstrated a positive response of unassisted species (*i.e.* species without  
347 dispersal structures, mostly gravity-dispersed species) in response to connectivity by both  
348 experimental (Damschen et al. 2008) and correlative (Kolb and Diekmann 2005; Thiele et al.  
349 2017) approaches. Damschen et al. (2008) demonstrated that the difference in unassisted  
350 species richness between connected and non-connected patches was been six-fold greater after  
351 seven years. A similar response was obtained by Thiele et al. (2017). The authors obtained a  
352 positive relationship between connectivity (Euclidean distance and potential functional  
353 connectivity) and similarity for unassisted species communities of linear landscape elements  
354 (field margins, ditches, road verges). Despite the increase in unassisted species richness  
355 (Damschen et al. 2008), the relative proportion of these species does not seem to be influenced



356 by connectivity within communities. When using a correlative approach, Lindborg (2007)  
357 reported that the proportion of unassisted species in semi-natural grasslands was independent  
358 of structural connectivity (IFM index). These results corroborate the work of Piessens et al.  
359 (2005), who demonstrated that unassisted species are not sensitive to structural connectivity.  
360 However, the dispersal mechanisms explaining the differential responses of unassisted species  
361 in terms of richness and relative proportion to landscape connectivity have yet to be clarified.  
362 As suggested by Damschen et al. (2008), these results challenge the notion that unassisted  
363 species are truly unassisted in their dispersal, corroborating evidence from other studies  
364 (Vellend et al. 2003; Higgins et al. 2003).

365

#### 366 *Plant dispersal traits*

367 Beyond the dispersal mode, precise combinations of traits might favour the ability of plants to  
368 produce a large number of potential dispersers (*e.g.* seed number, vegetative fragmentation),  
369 and to be dispersed over long distances (*e.g.* seed mass, low terminal velocity, high gut  
370 retention), with both characteristics corresponding to the definition of species dispersal ability  
371 (Johst et al. 2002; Nathan 2006; Auffret et al. 2017). Thus, species with traits promoting high  
372 dispersal ability might be less sensitive to connectivity loss.

373

#### 374 Emergent groups to connectivity

375 Initially, studies that analysed the dispersal traits of plants in response to connectivity were  
376 based on a functional group approach similar to the one proposed by Lavorel et al. (1997);  
377 namely, emergent groups are groups of species that reflect natural correlation of biological  
378 attributes. By coupling changes to plant occurrence or abundance along the gradient of  
379 connectivity and trait-data on plant species, these studies analysed the syndrome of traits that  
380 characterise plant sensitivity or resistance to connectivity.

381 One of the very first studies on emergent groups focused on forest communities (Kolb  
382 and Diekmann 2005). These authors analysed species biological traits for two groups of plants:  
383 one corresponding to sensitive plants (*i.e.* the occurrence of which decreases with structural  
384 connectivity loss, evaluated through IFM index) and one corresponding to resistant plants (*i.e.*  
385 the occurrence of which is not impacted by structural connectivity loss). Sensitive plants were  
386 characterised by high clonality, few and heavy seeds and unassisted dispersal. Resistant plants  
387 displayed assisted dispersal, no or little clonal growth and produced many and thin seeds. Traits  
388 promoting local establishment instead of long-distance dispersal contributed to plant sensitivity  
389 to connectivity loss. Using a similar approach, Adriaens et al. (2006) classified specialist  
390 species from calcareous grasslands, using 16 life-history traits, and detected four different  
391 groups. Higher structural landscape connectivity increased species richness in only one of the  
392 groups, whereas the other groups were not affected. This group contains species that are  
393 characterised by a perennial rosette with no vegetative multiplication and no long-distance  
394 dispersal mode (*i.e.* ballistochory, ant-dispersed seeds, barochory). Such species are more  
395 sensitive to connectivity loss, as they are less likely to recolonise and establish in patches after  
396 extinction. The three other groups were not impacted by connectivity loss. These groups  
397 included orchids (characterised by short flowering, low mass seeds and wind-dispersal), half-  
398 rosette species (with fruity dispersules and seeds, with optional vegetative reproduction  
399 mechanisms) and annual species (early and long flowering small autogamous annuals, low mass  
400 and no long-distance dispersal mode). However, the use of a large set of many traits, including  
401 ecological traits (*i.e.* linked with species niche), makes it difficult to disentangle the effect of  
402 dispersal traits from the other traits when evaluating how species respond to connectivity loss.  
403  
404 Response traits to connectivity

405 Studies over the last two decades have focused on traits considered individually to define  
406 individual response traits (*i.e.* traits reflecting the response of organisms to a given  
407 environmental factor, Lavorel et al. 1997) to landscape connectivity. This approach differs from  
408 the emergent group approach by analysing one trait at a time to characterise species sensitivity  
409 to connectivity. Species that displayed similar trait values in response to connectivity are  
410 designed as response groups of plants (*i.e.* group of species that respond in similar ways to a  
411 given environmental factor for a given trait, Lavorel et al. 1997).

412 Verheyen et al. (2004) characterised species sensitivity for 17 species of temperate  
413 forest communities to connectivity loss through the regression coefficient derived from the  
414 regression model, using patch occupancy as the dependent variable and structural connectivity  
415 (modified version of IFM index that incorporates patch age, Verheyen et al. 2004) as the  
416 independent variable. Species characterised by lower numbers of seeds (and thus, a low number  
417 of potential dispersers) were sensitive to connectivity loss. In ruderal (brownfields)  
418 communities, Schleicher et al. (2011) demonstrated that wind-dispersed species sensitive to  
419 structural connectivity loss are those characterised by lower seed number and higher terminal  
420 velocity (*i.e.* rate at which a seed can fall in still air). High terminal velocity corresponds to  
421 short dispersal range by reducing the probability of dispersal by air uplift, while reduced seed  
422 number corresponds to a low number of potential dispersers. Consequently, both trait  
423 expressions are indicative of low dispersal ability. Sensitive and resistant plants were the most  
424 differentiated using a quotient of the seed number divided by terminal velocity. Moreover, the  
425 authors found that the clearest differentiation between the two connectivity response groups  
426 was obtained when combining seed number and terminal velocity in a quotient. These results  
427 highlight a compensatory relationship between these two traits; high dispersal distance might  
428 counterbalance low numbers of dispersers and *vice versa*. Favre-Bac et al. (2017a) used traits  
429 directly linked to water-dispersal to analyse how ditch communities respond to connectivity

430 loss. The authors demonstrated that water-dispersed species that are highly sensitive to  
431 structural connectivity loss are characterised by short and round seeds with low mass and low  
432 buoyancy (Favre-Bac et al. 2017a). Low buoyancy does not favour long-distance dispersal in  
433 low current systems because species do not tolerate long or frequent retention events when  
434 being dispersed. The authors also demonstrated the role of seed form on the probability of being  
435 retained by obstacles in stagnant ditches during dispersal events. In this case, long and thin  
436 seeds might be better aligned to flow direction, and succeed better in passing obstacles, such as  
437 culverts and intersections (De Ryck et al. 2012). This type of study advanced our understanding  
438 of the factors that induce dispersal in the studied system, providing some predictions on the  
439 mechanisms causing species resistance in less connected networks.

440         Some studies demonstrated that connectivity responses are associated to traits linked  
441 with the ability of plants to establish in habitat patches. For instance, Favre-Bac et al. (2017a)  
442 reported that, in highly branched ditch networks (*i.e.* where intersections constitute obstacles to  
443 dispersal), the most resistant plants had higher seed germination rates. This higher seed  
444 germination rate increased their ability to develop when blocked by obstacles during dispersal.  
445 Other traits linked to further steps of plant growth, such as competition (for instance, plant  
446 height) and duration of growth (life-cycle duration), were also investigated. It was found that  
447 species with lower competition capacity and short life-cycles (in temperate forest communities,  
448 Verheyen et al. 2004, in heathland communities, Piessens et al. 2004, 2005; in dry calcareous  
449 grasslands communities, Evju et al. 2015) are more sensitive to the loss of structural  
450 connectivity.

451  
452         Therefore, it is important to develop approaches that take dispersal traits into account.  
453 Such approaches are expected to complement and fill knowledge gaps on existing work based  
454 on vector-types. Transposing the conceptual framework of response traits developed by Lavorel

455 et al. (1997) to landscape ecology represents a promising avenue towards understanding and  
456 predicting plant community responses to changes in connectivity.

457

#### 458 *Community weighted mean traits and functional diversity*

459 Studies over the last 15 years on community ecology have developed other methods to describe  
460 the functional structure of plant communities. These methods include aggregated measures  
461 quantifying the community the mean (*i.e.* community weighted mean traits) and range of trait  
462 values (functional diversity, for review, see Mouchet et al. 2010). In response to local factors,  
463 environmental filtering leads to convergent trait values that are adapted to environmental  
464 characteristics (van der Valk 1981; Weiher and Keddy 1995; Grime 2006). In contrast, biotic  
465 filtering can lead to: i) divergent trait values within communities (limiting similarity theory,  
466 MacArthur and Levins 1967; Grime 2006), leading to niche differentiation, or ii) convergent  
467 trait values within communities (competitive hierarchy theory, Herben and Goldberg 2014) due  
468 to the competitive exclusion of less competitive species. Transposition of these theories to the  
469 landscape scale is a promising research avenue in functional ecology. In response to reduced  
470 connectivity, plant communities at the landscape level should be filtered toward trait syndromes  
471 that promote high dispersal ability (*i.e.* low functional diversity and weighted mean trait values  
472 converge toward high seed number or traits promoting long-distance dispersal).

473         Research assessing how the functional structure of communities respond at the gamma  
474 scale depending on landscape characteristics is still at an early stage. To our knowledge, only  
475 one study has used this highly promising approach (Favre-Bac et al. 2017b). The authors  
476 demonstrated that lower ditch network connectivity induces convergent trait values toward  
477 small seed production and high seed buoyancy. Higher seed buoyancy is particularly  
478 favourable for successfully passing culverts, which constitute barriers (Soomers et al. 2010),  
479 because seeds might persist until water level changes, wind strength or wind direction enables

480 them to continue moving downstream. In comparison, lower seed production indicates that  
481 species invest heavily in vegetative growth, consolidating isolated local populations (Stöcklin  
482 and Winkler 2004). By contrast, higher connectivity in ditch networks reduces this filtering  
483 effect, or, even, leads to divergent trait values in seed mass. Consequently, in widely connected  
484 landscapes, several strategies coexist across local communities constituting the  
485 metacommunity. Higher seed weight provides more resources to guarantee growth during the  
486 early stages of establishment (Cornelissen et al. 2003), which mostly depend on competitive  
487 interactions with other plants within the local community. This local biotic filter favours the  
488 dissimilarity of plant traits involved in harvesting resources through the niche-partitioning  
489 effect (limiting similarity, MacArthur and Levins 1967; Pacala and Tilman 1994). These results  
490 demonstrate that, at the gamma scale, reduced connectivity acts as a filter on the dispersal traits  
491 of plant species involved in resistance to connectivity loss.

492

#### 493 *Contribution of functional traits towards improving landscape connectivity models*

494 Trait-based studies provide a better understanding of the mechanisms involved in how plants  
495 respond to connectivity. Thus, by integrating plant traits in landscape connectivity modelling,  
496 a more accurate prediction of the responses of species to connectivity loss should be obtained.  
497 This approach has been facilitated by the development of potential functional connectivity  
498 models (Fig. 1 and Box 1) that account for species dispersal distance in addition to the cost of  
499 species to cross over landscape features such as those of Pascual-Hortal and Saura (2006,  
500 Integral Index of Connectivity, IIC) and Saura and Pascual-Hortal (2007, Probability of  
501 Connectivity, PC). The two indices – IIC and PC – are based on different assumptions about  
502 the probability of connection between two considered patches (binary or probabilistic,  
503 respectively). However, only a few studies have used this type of graph-based connectivity

504 index for plants (see García-Feced et al. 2011; Aavik et al. 2014). Thus, these approaches need  
505 to be generalised.

506

### 507 **Effect of time on the response of plants to connectivity**

#### 508 *Historical landscape connectivity drives the taxonomic structure of plant communities*

509 Most studies have overlooked the potential effect of relaxation time when analysing the  
510 influence of connectivity on plant communities. However, this effect has been detected in  
511 grassland communities. For example, Helm et al. (2006) showed that the alpha richness of  
512 specialist calcareous grasslands is independent of their current structural connectivity, but is  
513 dependent on their connectivity 70 years ago (*i.e.* dependent on the state of the landscape before  
514 drastic habitat loss and connectivity loss). These results strongly support the concept that some  
515 plant communities are mostly composed of species that have the ability to resist to  
516 fragmentation by persisting without completing the whole life cycle (*i.e.* with no sexual  
517 reproduction stages, Eriksson 1996). Grassland plant communities are indeed mostly composed  
518 of perennial plants with long-lived vegetative life-cycle, and may then survive for decades after  
519 environmental changes (Eriksson 1996). Lindborg and Eriksson (2004) also detected a response  
520 of alpha diversity (richness and density) to landscape structural connectivity older than 50 years  
521 and 100 years ago in semi natural grasslands. The authors demonstrated an interactive effect  
522 between time and spatial resolution. A positive effect of connectivity 50 years ago was only  
523 detected for alpha diversity at the smaller scale (1 km radius); however, a positive effect of  
524 connectivity 100 years ago was detected at both scales considered (1 km radius and 2 km  
525 radius). The presence of a relaxation time following an environmental perturbation might have  
526 indirect consequences on the biological scale of the response. For instance, Cousins and  
527 Vanhoenacker (2011) demonstrated that the gamma diversity of semi-grassland ecosystems  
528 decreases more slowly after a decline in grasslands abundance in the landscape compared to  
529 alpha diversity. Although no study has yet investigated the difference in the response of alpha

530 and gamma diversity on historical connectivity, we assume that such results could be transposed  
531 to connectivity loss. The time required to establish an equilibrium between spatial connectivity  
532 and species richness or diversity is longer at larger spatial and biological scales compared to  
533 smaller scales (O'Neill et al. 1986; Allen and Starr 1988); thus, the relaxation time of plant  
534 assemblages occurs at different spatial scales.

535 To our knowledge, only Naaf and Kolk (2015) and Haddad et al. (2015) have  
536 investigated how connectivity affects the magnitude of immigration credit. By studying newly-  
537 established forest patches, Naaf and Kolk (2015) found that the magnitude of immigration credit  
538 mainly depends on the structural connectivity of forests (IFM index). In connected forest  
539 patches, immigration lag affected five forest specialist species compared to nine species in  
540 highly isolated forest patches. Haddad et al. (2015) obtained similar results by studying  
541 successional vegetation of pine plantations over a decade in the Savannah River Site  
542 experimental design (Box 2). More specifically, immigration lags resulted in 15% fewer species  
543 after a decade in unconnected patches compared to connected patches. Future studies should  
544 investigate this effect in other community types.

545

#### 546 *Dispersal traits promoting time-dependent responses to connectivity*

547 Because plants might respond to connectivity time-dependently (either with relaxation time or  
548 with immigration credit), some biological traits might promote such effects. For instance, the  
549 life duration of species or seed-bank persistence might contribute to the delayed response to  
550 changes in connectivity. The work of Lindborg (2007) supported this assumption,  
551 demonstrating that the proportion of short-lived plant species is influenced by current structural  
552 connectivity, whereas that of clonal long-lived and long seed-bank persistence species is  
553 influenced by historical structural connectivity (Table 1). Conversely, some traits might  
554 influence the effect of immigration lag. For instance, species with poor dispersal abilities might



555 contribute strongly to immigration credit. This assumption was verified by Naaf and Kolk  
556 (2015). The authors demonstrated that immigration credit is higher in species that are  
557 characterised by low seed production and dispersal potential (*sensu* Vittoz and Engler 2007)  
558 compared to species with high seed production and dispersal potential in forest specialist  
559 communities. However, few studies have used trait-based approaches to determine how  
560 connectivity changes over time.

561

### 562 **Effect of landscape connectivity on actual seed dispersal**

563 Studies assessing actual dispersal (Fig. 1) in response to connectivity have developed over the  
564 last 15 years, but remain limited. Many papers have assessed seed fluxes at the community  
565 level, particularly for water-dispersed species (*e.g.* Andersson and Nilsson 2002; Boedeltje et  
566 al. 2003, 2004; Moggridge et al. 2009), but very few studies have considered the effect of  
567 connectivity on these seed fluxes. Only a few studies have taken connectivity into account, with  
568 the response of plants being almost exclusively assessed at the species level, which are  
569 presented in the next section.

570

#### 571 *Number of dispersed seeds*

572 The effect of connectivity on the abundance of dispersed seeds has mostly been investigated  
573 under experimental conditions, focusing on particular species that were selected for their  
574 abundance or representativeness of the community process. Seed fluxes for three plant species  
575 dispersed by birds were assessed in connected and unconnected patches using the Savannah  
576 River Site experimental design. Seeds were trapped in peripheral (*i.e.* receiver) patches, of  
577 which just one was connected to the central patch and the others were not (Box 2). To provide  
578 confidence that all seeds or fruits found in seed traps were exclusively derived from the central  
579 patch, two methods were carried out, depending on the abundance of the species considered: i)

580 the removal of all naturally occurring individuals from the peripheral patches, and ii) the use of  
581 marked seeds or fruits in the central patch by fluorescence. Based on this approach, Tewksbury  
582 et al. (2002) reported that twice the number of *Ilex vomitoria* seeds was trapped in connected  
583 patches compared to unconnected patches. The authors also reported that a greater proportion  
584 (an increase of 18%) of *Myrica cerifera* were present in faecal samples collected in connected  
585 patches that contained fluorescent powder compared to unconnected patches. Similar results  
586 were obtained by Haddad et al. (2003) for both *Myrica cerifera* and *Rhus copallina*, with five  
587 and two times more seeds, respectively, being transported from the central patch in connected  
588 compared to unconnected patches. Using an individual-based model, Levey et al. (2005)  
589 demonstrated that the distribution of *Myrica cerifera* is explained by the movement of birds  
590 that were 31% more likely to be found in connected patches compared to unconnected patches.  
591 Thus, it is important that future studies extend these analyses to consider other modes of  
592 dispersal.

593

#### 594 *Seed flux composition*

595 Studies analysing the seed communities of plants that actually disperse in response to landscape  
596 connectivity remain scarce. To our knowledge, only one study investigated this type of  
597 relationship (Suárez-Esteban et al. 2013). The authors assumed that unpaved roads act as  
598 corridors, and analysed actual dispersal along unpaved roads in comparison to adjacent  
599 scrubland, which was not considered to be a corridor. By analysing the seeds of all fleshy-fruit  
600 shrubs that were contained in the faecal samples of frugivorous mammals, the authors  
601 demonstrated that the composition of seed communities in unpaved roads differed to that  
602 observed in scrublands, due to the effect of corridors on animal behaviour. However, this  
603 finding was dependent on the animal-vector being considered (rabbit, carnivore or ungulate).

604 Such studies analysing actual dispersal are expected to improve the accuracy of assessing  
605 current seed dispersal, allowing us to determine how connectivity influences dispersal *per se*.

606

### 607 **New prospects for future research**

608 Analysing the response of the plant community to connectivity loss has not been as extensively  
609 studied as for animal guilds, despite the dispersal specificity of plants. Through our review, we  
610 demonstrated that: i) landscape connectivity does promote the actual dispersal of plants  
611 between communities (section 4), but ii) its influence on community taxonomic structure does  
612 not follow a general pattern due to the many factors that modulate its effect (sections 1 and  
613 3). We also found that: iii) the use of functional traits provides a better understanding of the  
614 mechanisms involved in plant responses to connectivity (sections 2 and 3).

615         These three key-findings should be considered in light of several limitations related to  
616 the studies selected for this review. First, this review did not compare the relative role of  
617 landscape connectivity *vs.* other landscape factors (*e.g.* habitat size or habitat amount) in  
618 structuring plant communities. Thus, it is beyond the scope of this review to shed light on the  
619 SLOSS (Single Large Or Several Small) debate, derived from island biogeography or to discuss  
620 on the relative importance between SLOSS and the Habitat Amount Hypothesis (Fahrig 2013)  
621 (but see Lindgren and Cousins 2017). Secondly, the dark diversity (*i.e.* all species that are  
622 absent from a habitat patch but that could disperse to and establish there, Pärtel et al. 2011) may  
623 result from dispersal limitation (Riibak et al. 2015). However, the reviewed studies investigated  
624 the role of landscape connectivity on expressed plant communities (*i.e.* potential dispersal, Fig.  
625 1), rather than on the absence of species, neglecting the role of landscape connectivity on dark  
626 diversity. Third, the consideration of traits in the response to connectivity raises a question for  
627 some authors (see for example Haddad et al. 2015). These authors suggest that plant dispersal  
628 might be better described by statistical probabilities and stochastic factors rather than traits.

629 Within these limitations, we identified research prospects from the three key-findings identified  
630 here:

631

632 1) Landscape connectivity promotes actual dispersal between connected communities. Because  
633 of the scarcity and the animal-dispersed species focus of the studies on this topic, we can only  
634 encourage more studies to use this approach, particularly for other dispersal modes.

635

636 2) The influence of landscape connectivity on plant taxonomic structure do not follow a general  
637 pattern because of the multiplicity of factors that modulate its effect. To better disentangle the  
638 effect of landscape connectivity from the other factors, explicit consideration of both spatial  
639 and temporal scales is necessary to define a relevant resolution of the landscape. An assessment  
640 of landscape connectivity at different spatial scales should be developed to detect the dispersal  
641 range at which plant species respond to changes in connectivity. Because some species exhibit  
642 a time lag in their response to connectivity changes, efforts to incorporate historical data to  
643 model historical connectivity are necessary. In addition, the presence of a corridor *per se* is not  
644 sufficient to connect habitats. Indeed, the quality of the corridors (or landscape elements) should  
645 be considered and determined by their management practice. For instance, in hedgerows, the  
646 corridor quality is determined by the vegetation structure and vertical organisation at a fine  
647 scale. The assessment of connectivity might account for these fine-scale characteristics by using  
648 recent remote sensing techniques (Betbeder et al. 2014) to better predict species abundance  
649 such as for carabids (Betbeder et al. 2015). Future studies should investigate this new  
650 methodology for plants. Connectivity might be evaluated while considering the potential  
651 interaction between habitats, to better account for the degree of ecological specialisation of  
652 species. Complementarity between the different ecological corridor types (*i.e.* provided by  
653 different habitat types) might explain the lowest sensitivity of generalist species to connectivity

654 loss. In contrast, antagonistic effects might occur for specialist species. The deployment of such  
655 multi-habitat landscape connectivity models (which independently consider different  
656 ecological corridor types) could enhance our understanding of the responses of plant species to  
657 connectivity at a community scale.

658

659 3) Using functional traits provides a better understanding of the mechanisms involved in plant  
660 responses to connectivity. Some trait syndromes characterise species that are highly sensitive  
661 to connectivity loss. Determining trait combinations that might predict, at best, plant responses  
662 in time and space to changes in connectivity is a key challenge of forthcoming research. Using  
663 such functional indicators would allow these species to be detected in a manner that is  
664 reproducible and independent of ecosystems and regional pools (Lavorel et al. 1997). This  
665 approach is promising for defining general functional indicators of sensitivity. Such indicators  
666 may, for instance, help to identify species that are less likely to colonise and establish in patches  
667 after connectivity loss. In addition, the use of functional connectivity indices, which considers  
668 the dispersal distance of species, should be generalized in studies investigating plants,  
669 especially at the community level. The dispersal traits that promote time-dependent responses  
670 to connectivity that we have identified should be integrated in these indices to provide an  
671 adequate understanding of connections between plant communities over time.

672

673 Beyond highlighting needs for future research, this synthesis provides information of use for  
674 land-use planning (green and blue infrastructure implementation, for instance, Sandström 2002;  
675 Tzoulas et al. 2007). Such information could provide new methods and decision-making tools  
676 to promote the operational establishment of ecological corridors, taking plant communities into  
677 account.

678

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682 **References**

- 683
- 684 Aavik T, Holderegger R, Bolliger J (2014) The structural and functional connectivity of the  
685 grassland plant *Lychnis flos-cuculi*. *Heredity* 112:471–478. doi: 10.1038/hdy.2013.120
- 686 Adriaens D, Honnay O, Hermy M (2006) No evidence of a plant extinction debt in highly  
687 fragmented calcareous grasslands in Belgium. *Biol Conserv* 133:212–224. doi:  
688 10.1016/j.biocon.2006.06.006
- 689 Adriaensen F, Chardon JP, De Blust G, Swinnen E, Villalba S, Gulinck H, Matthysen E (2003)  
690 The application of ‘leastcost’ modelling as a functional landscape model. *Landsc Urban Plan*  
691 64:233–247. [https://doi.org/10.1016/S0169-2046\(02\)00242-6](https://doi.org/10.1016/S0169-2046(02)00242-6)
- 692 Allen TFH, Starr TB (1988) *Hierarchy: Perspectives for Ecological Complexity*. University of  
693 Chicago Press, Chicago
- 694 Andersson E, Nilsson C (2002) Temporal variation in the drift of plant litter and propagules in  
695 a small boreal river. *Freshw Biol* 47:1674–1684. doi: 10.1046/j.1365-2427.2002.00925.x
- 696 Andreasen C, Stryhn H, Streibig JC (1996) Decline of the Flora in Danish Arable Fields. *J Appl*  
697 *Ecol* 33:619–626. doi: 10.2307/2404990
- 698 Archer S, Pyke DA (1991) Plant-animal interactions affecting plant establishment and  
699 persistence on revegetated rangeland. *J Range Manag* 44:558–565. doi: 10.2307/4003036
- 700 Arellano-Rivas A, De-Nova JA, Munguía-Rosas MA (2016) Patch isolation and shape predict  
701 plant functional diversity in a naturally fragmented forest. *J Plant Ecol* 1–11. doi:  
702 10.1093/jpe/rtw119
- 703 Auffret AG, Rico Y, Bullock JM, Hooftman DAP, Pakeman RJ, Soons MB, Suárez-Esteban A,  
704 Traveset A, Wagner H, Cousins SAO (2017) Plant functional connectivity – integrating  
705 landscape structure and effective dispersal. *J Ecol*. doi: 10.1111/1365-2745.12742
- 706 Ayram CAC, Mendoza ME, Etter A, Salicrup DRP (2016) Habitat connectivity in biodiversity  
707 conservation: A review of recent studies and applications. *Prog Phys Geogr Earth Environ*  
708 40:7–37. doi: 10.1177/0309133315598713
- 709 Baessler C, Klotz S (2006) Effects of changes in agricultural land-use on landscape structure  
710 and arable weed vegetation over the last 50 years. *Agric Ecosyst Environ* 115:43–50. doi:  
711 10.1016/j.agee.2005.12.007
- 712 Beier P, Noss RF (1998) Do Habitat Corridors Provide Connectivity? *Conserv Biol* 12:1241–  
713 1252. doi: 10.1111/j.1523-1739.1998.98036.x
- 714 Bennett AF (2003) *Linkages in the Landscape: The Role of Corridors and Connectivity in*  
715 *Wildlife Conservation*. IUCN-The World Conservation Union, Gland, Cambridge

716 Bennett AJ, Radford JQ, Haslem A (2006) Properties of Land Mosaics: Implications for Nature  
717 Conservation in Agricultural Environments. *Biol Conserv* 133:250–264. doi:  
718 10.1016/j.biocon.2006.06.008

719 Betbeder J, Hubert-Moy L, Burel F, Corgne S, Baudry J (2015) Assessing ecological habitat  
720 structure from local to landscape scales using synthetic aperture radar. *Ecol Indic* 52:545–557.  
721 doi: 10.1016/j.ecolind.2014.11.009

722 Betbeder J, Nabucet J, Pottier E, Baudry J, Corgne S, Hubert-Moy L (2014) Detection and  
723 Characterization of Hedgerows Using TerraSAR-X Imagery. *Remote Sens* 6:3752–3769. doi:  
724 10.3390/rs6053752

725 Boedeltje G, Bakker JP, Bekker RM, Van Groenendael JM, Soesbergen M (2003) Plant  
726 dispersal in a lowland stream in relation to occurrence and three specific life-history traits of  
727 the species in the species pool. *J Ecol* 91:855–866. doi: 10.1046/j.1365-2745.2003.00820.x

728 Boedeltje G, Bakker JP, Ten Brinke A, Van Groenendael JM, Soesbergen M (2004) Dispersal  
729 phenology of hydrochorous plants in relation to discharge, seed release time and buoyancy of  
730 seeds: the flood pulse concept supported. *J Ecol* 92:786–796. doi: 10.1111/j.0022-  
731 0477.2004.00906.x

732 Bornette G, Amoros C, Lamouroux N (1998) Aquatic plant diversity in riverine wetlands: the  
733 role of connectivity. *Freshw Biol* 39:267–283. doi: 10.1046/j.1365-2427.1998.00273.x

734 Bowne DR, Bowers MA (2004) Interpatch movements in spatially structured populations: a  
735 literature review. *Landsc Ecol* 19:1–20. doi: 10.1023/B:LAND.0000018357.45262.b9

736 Brückmann SV, Krauss J, Steffan-Dewenter I (2010) Butterfly and plant specialists suffer from  
737 reduced connectivity in fragmented landscapes. *J Appl Ecol* 47:799–809. doi: 10.1111/j.1365-  
738 2664.2010.01828.x

739 Brudvig LA (2016) Interpreting the effects of landscape connectivity on community diversity.  
740 *J Veg Sci* 27:4–5. doi: 10.1111/jvs.12365

741 Brudvig LA, Damschen EI, Tewksbury JJ, Haddad NM, Levey DJ (2009) Landscape  
742 connectivity promotes plant biodiversity spillover into non-target habitats. *Proc Natl Acad Sci*  
743 106:9328–9332. doi: 10.1073/pnas.0809658106

744 Burel F, Baudry J (1999) *Ecologie du paysage. Concepts, méthodes et applications*. Editions  
745 TEC et DOC, Paris

746 Calabrese JM, Fagan WF (2004) A comparison-shopper's guide to connectivity metrics. *Front*  
747 *Ecol Environ* 2:529–536. doi: 10.1890/1540-9295(2004)002[0529:ACGTCM]2.0.CO;2



748 Chetkiewicz C-LB, Clair CCS, Boyce MS (2006) Corridors for Conservation: Integrating  
749 Pattern and Process. *Annu Rev Ecol Evol Syst* 37:317–342. doi:  
750 10.1146/annurev.ecolsys.37.091305.110050

751 Cornelissen JHC, Lavorel S, Garnier E, Díaz S, Buchmann N, Gurvich DE, Reich PB, Ter  
752 Steege H, Morgan HD, Van Der Heijden MGA, Pausas JG, Poorter H (2003) A handbook of  
753 protocols for standardised and easy measurement of plant functional traits worldwide. *Aust J*  
754 *Bot* 51:335–380. doi: 10.1071/BT02124

755 Cousins SAO, Ohlson H, Eriksson O (2007) Effects of historical and present fragmentation on  
756 plant species diversity in semi-natural grasslands in Swedish rural landscapes. *Landsc Ecol*  
757 22:723–730. doi: 10.1007/s10980-006-9067-1

758 Cousins SAO, Vanhoenacker D (2011) Detection of extinction debt depends on scale and  
759 specialisation. *Biol Conserv* 144:782–787. doi: 10.1016/j.biocon.2010.11.009

760 Damschen EI, Baker DV, Bohrer G, Nathan R, Orrock JL, Turner JR, Brudvig LA, Haddad  
761 NM, Levey DJ, Tewksbury JJ (2014) How fragmentation and corridors affect wind dynamics  
762 and seed dispersal in open habitats. *Proc Natl Acad Sci* 111:3484–3489. doi:  
763 10.1073/pnas.1308968111

764 Damschen EI, Brudvig LA, Haddad NM, Levey DJ, Orrock JL, Tewksbury JJ (2008) The  
765 movement ecology and dynamics of plant communities in fragmented landscapes. *Proc Natl*  
766 *Acad Sci* 105:19078–19083. doi: 10.1073/pnas.0802037105

767 Damschen EI, Haddad NM, Orrock JL, Tewksbury JJ, Levey DJ (2006) Corridors increase  
768 plant species richness at large scales. *Science* 313:1284–1286. doi: 10.1126/science.1130098

769 De Ryck DJR, Robert EMR, Schmitz N, Van der Stocken T, Di Nitto D, Dahdouh-Guebas F,  
770 Koedam N (2012) Size does matter, but not only size: Two alternative dispersal strategies for  
771 viviparous mangrove propagules. *Aquat Bot* 103:66–73. doi: 10.1016/j.aquabot.2012.06.005

772 Diamond JM (1972) Biogeographic kinetics: estimation of relaxation times for avifaunas of  
773 southwest pacific islands. *Proc Natl Acad Sci* 69:3199–3203. doi: 10.1073/pnas.69.11.3199

774 Dzwonko Z, Loster S (1988) Species richness of small woodlands on the western Carpathian  
775 foothills. *Vegetatio* 76:15–27. doi: 10.1007/BF00047384

776 Eriksson O (1996) Regional Dynamics of Plants: A Review of Evidence for Remnant, Source-  
777 Sink and Metapopulations. *Oikos* 77:248–258. doi: 10.2307/3546063

778 Evju M, Blumentrath S, Skarpaas O, Stabbetorp OE (2015) Plant species occurrence in a  
779 fragmented grassland landscape: the importance of species traits. *Biodivers Conserv* 24:547–  
780 561. doi: 10.1007/s10531-014-0835-y

781 Fahrig, L. (2013). Rethinking patch size and isolation effects: the habitat amount hypothesis.  
782 *Journal of Biogeography*, 40(9), 1649–1663. doi:10.1111/jbi.12130

783 Fahrig L (2003) Effects of Habitat Fragmentation on Biodiversity. *Annu Rev Ecol Evol Syst*  
784 34:487–515. doi: 10.1146/annurev.ecolsys.34.011802.132419

785 Fahrig L, Merriam G (1985) Habitat Patch Connectivity and Population Survival. *Ecology*  
786 66:1762–1768. doi: 10.2307/2937372

787 Favre-Bac L, Ernoult A, Mony C, Rantier Y, Nabucet J, Burel F (2014) Connectivity and  
788 propagule sources composition drive ditch plant metacommunity structure. *Acta Oecologica*  
789 61:57–64. doi: 10.1016/j.actao.2014.10.006

790 Favre-Bac L, Lamberti-Raverot B, Puijalón S, Ernoult A, Burel F, Guillard L, Mony C (2017a)  
791 Plant dispersal traits determine hydrochorous species tolerance to connectivity loss at the  
792 landscape scale. *J Veg Sci* 28:605–615. doi: 10.1111/jvs.12518

793 Favre-Bac L, Mony C, Burel F, Seimandi-Corda G, Ernoult A (2017b) Connectivity drives the  
794 functional diversity of plant dispersal traits in agricultural landscapes: the example of ditch  
795 metacommunities. *Landsc Ecol* 32:2029–2040. doi: 10.1007/s10980-017-0564-1

796 Fischer J, Lindenmayer DB (2007) Landscape modification and habitat fragmentation: a  
797 synthesis. *Glob Ecol Biogeogr* 16:265–280. doi: 10.1111/j.1466-8238.2007.00287.x

798 Fletcher RJ, Burrell NS, Reichert BE, Vasudev D, Austin JD (2016) Divergent Perspectives on  
799 Landscape Connectivity Reveal Consistent Effects from Genes to Communities. *Curr Landsc*  
800 *Ecol Rep* 1:67–79. doi: 10.1007/s40823-016-0009-6

801 Fukami T (2015) Historical Contingency in Community Assembly: Integrating Niches, Species  
802 Pools, and Priority Effects. *Annu Rev Ecol Evol Syst* 46:1–23. doi: 10.1146/annurev-ecolsys-  
803 110411-160340

804 Fukami T, Nakajima M (2013) Complex plant–soil interactions enhance plant species diversity  
805 by delaying community convergence. *J Ecol* 101:316–324. doi: 10.1111/1365-2745.12048

806 García-Feced C, Saura S, Elena-Rosselló R (2011) Improving landscape connectivity in forest  
807 districts: A two-stage process for prioritizing agricultural patches for reforestation. *For Ecol*  
808 *Manag* 261:154–161. doi: 10.1016/j.foreco.2010.09.047

809 Gignac LD, Dale MRT (2007) Effects of size, shape, and edge on vegetation in remnants of the  
810 upland boreal mixed-wood forest in agro-environments of Alberta, Canada. *Can J Bot* 85:273–  
811 284. doi: 10.1139/B07-018

812 Gilbert-Norton L, Wilson R, Stevens JR, Beard KH (2010) A meta-analytic review of corridor  
813 effectiveness. *Conserv Biol* 24:660–668. doi: 10.1111/j.1523-1739.2010.01450.x

814 Godefroid S, Koedam N (2003) How important are large vs. small forest remnants for the  
815 conservation of the woodland flora in an urban context? *Glob Ecol Biogeogr* 12:287–298. doi:  
816 10.1046/j.1466-822X.2003.00035.x

817 Grime JP (2006) Trait convergence and trait divergence in herbaceous plant communities:  
818 Mechanisms and consequences. *J Veg Sci* 17:255–260. doi: 10.1111/j.1654-  
819 1103.2006.tb02444.x

820 Groves CR, Jensen DB, Valutis LL, Redford KH, Shaffer ML, Scott JM, Baumgartner JV,  
821 Higgins JV, Beck MW, Anderson MG (2002) Planning for Biodiversity Conservation: Putting  
822 Conservation Science into Practice. A seven-step framework for developing regional plans to  
823 conserve biological diversity, based upon principles of conservation biology and ecology, is  
824 being used extensively by the nature conservancy to identify priority areas for conservation.  
825 *BioScience* 52:499–512. doi: 10.1641/0006-3568(2002)052[0499:PFBCPC]2.0.CO;2

826 Guisan A, Thuiller W (2005) Predicting species distribution: offering more than simple habitat  
827 models. *Ecol Lett* 8:993–1009. doi: 10.1111/j.1461-0248.2005.00792.x

828 Haddad NM, Bowne DR, Cunningham A, Levey DJ, Sargent S, Spira T (2003) Corridor use by  
829 diverse taxa. *Ecology* 84:609–615. doi: 10.1890/0012-9658(2003)084[0609:CUBDT]2.0.CO;2

830 Haddad NM, Brudvig LA, Clobert J, Davies KF, Gonzalez A, Holt RD, Lovejoy TE, Sexton  
831 JO, Austin MP, Collins CD, Cook WM, Damschen EI, Ewers RM, Foster BL, Jenkins CN,  
832 King AJ, Laurance WF, Levey DJ, Margules CR, Melbourne BA, Nicholls AO, Orrock JL, Son  
833 D-X, Townshend JR (2015) Habitat fragmentation and its lasting impact on Earth’s ecosystems.  
834 *Sci Adv* 1:e1500052. doi: 10.1126/sciadv.1500052

835 Haddad NM, Hudgens B, Damschen EI, Levey DJ, Orrock JL, Tewksbury JJ, Weldon AJ  
836 (2011) Assessing positive and negative ecological effects of corridors. In: Liu J, Hull V,  
837 Morzillo AT, Wiens JA (eds) *Sources, Sinks and Sustainability*. Cambridge University Press,  
838 Cambridge, pp 475–503

839 Haddad NM, Tewksbury JJ (2006) Impacts of corridors on populations and communities. In:  
840 Crooks KR, Sanjayan M (eds) *Connectivity Conservation: maintaining connections for nature*.  
841 Cambridge University Press, Cambridge, pp 390–415

842 Hanski I (1994) A Practical Model of Metapopulation Dynamics. *J Anim Ecol* 63:151–162.  
843 doi: 10.2307/5591

844 Helm A, Hanski I, Pärtel M (2006) Slow response of plant species richness to habitat loss and  
845 fragmentation. *Ecol Lett* 9:72–77. doi: 10.1111/j.1461-0248.2005.00841.x

846 Herben T, Goldberg DE (2014) Community assembly by limiting similarity vs. competitive  
847 hierarchies: testing the consequences of dispersion of individual traits. *J Ecol* 102:156–166.  
848 doi: 10.1111/1365-2745.12181

849 Higgins SI, Nathan R, Cain ML (2003) Are Long-Distance Dispersal Events in Plants Usually  
850 Caused by Nonstandard Means of Dispersal? *Ecology* 84:1945–1956. doi: 10.1890/01-0616

851 Honnay O, Hermy M, Coppin P (1999) Effects of area, age and diversity of forest patches in  
852 Belgium on plant species richness, and implications for conservation and reforestation. *Biol*  
853 *Conserv* 87:73–84. doi: 10.1016/S0006-3207(98)00038-X

854 Jackson ST, Sax DF (2010) Balancing biodiversity in a changing environment: extinction debt,  
855 immigration credit and species turnover. *Trends Ecol Evol* 25:153–160. doi:  
856 10.1016/j.tree.2009.10.001

857 Johst K, Brandl R, Eber S (2002) Metapopulation persistence in dynamic landscapes: the role  
858 of dispersal distance. *Oikos* 98:263–270. doi: 10.1034/j.1600-0706.2002.980208.x

859 Katoh K, Sakai S, Takahashi T (2009) Factors maintaining species diversity in satoyama, a  
860 traditional agricultural landscape of Japan. *Biol Conserv* 142:1930–1936. doi:  
861 10.1016/j.biocon.2009.02.030

862 Kleijn D, Kohler F, Báldi A, Concepción ED, Clough Y, Díaz M, Gabriel D, Holzschuh A,  
863 Knop E, Kovács A, Marshall EJP, Tschamntke T, Verhulst J (2009) On the relationship between  
864 farmland biodiversity and land-use intensity in Europe. *Proc Biol Sci* 276:903–909. doi:  
865 10.1098/rspb.2008.1509

866 Kleijn D, Verbeek M (2000) Factors affecting the species composition of arable field boundary  
867 vegetation. *J Appl Ecol* 37:256–266. doi: 10.1046/j.1365-2664.2000.00486.x

868 Koen EL, Bowman J, Walpole AA (2012) The effect of cost surface parameterization on  
869 landscape resistance estimates. *Mol Ecol Resour* 12:686–696. doi: 10.1111/j.1755-  
870 0998.2012.03123.x

871 Kolb A, Diekmann M (2005) Effects of Life-History Traits on Responses of Plant Species to  
872 Forest Fragmentation. *Conserv Biol* 19:929–938. doi: 10.1111/j.1523-1739.2005.00065.x

873 Kuussaari M, Bommarco R, Heikkinen RK, Helm A, Krauss J, Lindborg R, Öckinger E, Pärter  
874 M, Pino J, Rodà F, Stefanescu C, Teder T, Zobel M, Steffan-Dewenter I (2009) Extinction debt:  
875 a challenge for biodiversity conservation. *Trends Ecol Evol* 24:564–571. doi:  
876 10.1016/j.tree.2009.04.011

877 Lavorel S, McIntyre S, Landsberg J, Forbes TDA (1997) Plant functional classifications: from  
878 general groups to specific groups based on response to disturbance. *Trends Ecol Evol* 12:474–  
879 478. doi: 10.1016/S0169-5347(97)01219-6

880 Leibold MA, Holyoak M, Mouquet N, Amarasekare P, Chase JM, Hoopes MF, Holt RD, Shurin  
881 JB, Law R, Tilman D, Loreau M, Gonzalez A (2004) The metacommunity concept: a  
882 framework for multi-scale community ecology. *Ecol Lett* 7:601–613. doi: 10.1111/j.1461-  
883 0248.2004.00608.x

884 Levey DJ, Bolker BM, Tewksbury JJ, Sargent S, Haddad NM (2005) Effects of landscape  
885 corridors on seed dispersal by birds. *Science* 309:146–148. doi: 10.1126/science.1111479

886 Lindborg R (2007) Evaluating the distribution of plant life-history traits in relation to current  
887 and historical landscape configurations. *J Ecol* 95:555–564. doi: 10.1111/j.1365-  
888 2745.2007.01232.x

889 Lindborg R, Eriksson O (2004) Historical Landscape Connectivity Affects Present Plant  
890 Species Diversity. *Ecology* 85:1840–1845. doi: 10.1890/04-0367

891 Lindgren JP, Cousins SAO (2017) Island biogeography theory outweighs habitat amount  
892 hypothesis in predicting plant species richness in small grassland remnants. *Landsc Ecol*  
893 32:1895–1906. doi: 10.1007/s10980-017-0544-5

894 Logue JB, Mouquet N, Peter H, Hillebrand H (2011) Empirical approaches to  
895 metacommunities: a review and comparison with theory. *Trends Ecol Evol* 26:482–491. doi:  
896 10.1016/j.tree.2011.04.009

897 Luoto M, Rekolainen S, Aakkula J, Pykälä J (2003) Loss of Plant Species Richness and Habitat  
898 Connectivity in Grasslands Associated with Agricultural Change in Finland. *AMBIO J Hum*  
899 *Environ* 32:447–452. doi: 10.1579/0044-7447-32.7.447

900 MacArthur R, Levins R (1967) The Limiting Similarity, Convergence, and Divergence of  
901 Coexisting Species. *Am Nat* 101:377–385. doi: 10.1086/282505

902 MacArthur RH, Wilson E (1967) *The Theory of Island Biogeography*. Princeton University  
903 Press, Princeton

904 McRae BH, Dickson BG, Keitt TH, Shah VB (2008) Using Circuit Theory to Model  
905 Connectivity in Ecology, Evolution, and Conservation. *Ecology* 89:2712–2724. doi:  
906 10.1890/07-1861.1

907 Minor ES, Gardner RH (2011) Landscape connectivity and seed dispersal characteristics inform  
908 the best management strategy for exotic plants. *Ecol Appl Publ Ecol Soc Am* 21:739–749

909 Minor ES, Tessel SM, Engelhardt KAM, Lookingbill TR (2009) The role of landscape  
910 connectivity in assembling exotic plant communities: a network analysis. *Ecology* 90:1802–  
911 1809. doi: 10.1890/08-1015.1

912 Moggridge HL, Gurnell AM, Mountford JO (2009) Propagule input, transport and deposition  
913 in riparian environments: the importance of connectivity for diversity. *J Veg Sci* 20:465–474.  
914 doi: 10.1111/j.1654-1103.2009.05498.x

915 Moilanen A, Nieminen M (2002) Simple Connectivity Measures in Spatial Ecology. *Ecology*  
916 83:1131–1145. doi: 10.2307/3071919

917 Morato RG, Ferraz KMPM de B, de Paula RC, de Campos CB (2014) Identification of Priority  
918 Conservation Areas and Potential Corridors for Jaguars in the Caatinga Biome, Brazil. *PLoS*  
919 *ONE* 9:. doi: 10.1371/journal.pone.0092950

920 Mouchet MA, Villéger S, Mason NWH, Mouillot D (2010) Functional diversity measures: an  
921 overview of their redundancy and their ability to discriminate community assembly rules. *Funct*  
922 *Ecol* 24:867–876. doi: 10.1111/j.1365-2435.2010.01695.x

923 Mouquet N, E. Miller T, Daufresne T, M. Kneitel J (2006) Consequences of varying regional  
924 heterogeneity in source–sink metacommunities. *Oikos* 113:481–488. doi: 10.1111/j.2006.0030-  
925 1299.14582.x

926 Mouquet N, Loreau M (2003) Community patterns in source-sink metacommunities. *Am Nat*  
927 162:544–557

928 Murphy HT, Lovett-Doust J (2004) Context and connectivity in plant metapopulations and  
929 landscape mosaics: does the matrix matter? *Oikos* 105:3–14. doi: 10.1111/j.0030-  
930 1299.2004.12754.x

931 Naaf T, Kolk J (2015) Colonization credit of post-agricultural forest patches in NE Germany  
932 remains 130–230years after reforestation. *Biol Conserv* 182:155–163. doi:  
933 10.1016/j.biocon.2014.12.002

934 Nathan R (2006) Long-distance dispersal of plants. *Science* 313:786–788. doi:  
935 10.1126/science.1124975

936 Nathan R, Katul GG, Horn HS, Thomas SM, Oren R, Avissar R, Pacala SW, Levin SA (2002)  
937 Mechanisms of long-distance dispersal of seeds by wind. *Nature* 418:409–413. doi:  
938 10.1038/nature00844

939 Nilsson C, Brown RL, Jansson R, Merritt DM (2010) The role of hydrochory in structuring  
940 riparian and wetland vegetation. *Biol Rev Camb Philos Soc* 85:837–858. doi: 10.1111/j.1469-  
941 185X.2010.00129.x

942 O'Neill RV, Deangelis DL, Waide JB, Allen TFH (1986) *A Hierarchical Concept of*  
943 *Ecosystems*. Princeton University Press, Princeton

944 Orrock JL, Damschen EI (2005) Corridors Cause Differential Seed Predation. *Ecol Appl*  
945 15:793–798. doi: 10.1890/04-1129

946 Orrock JL, Danielson BJ, Burns MJ, Levey DJ (2003) Spatial Ecology of Predator–Prey  
947 Interactions: Corridors and Patch Shape Influence Seed Predation. *Ecology* 84:2589–2599. doi:  
948 10.1890/02-0439

949 Ovaskainen O, Hanski I (2002) Transient Dynamics in Metapopulation Response to  
950 Perturbation. *Theor Popul Biol* 61:285–295. doi: 10.1006/tpbi.2002.1586

951 Pacala SW, Tilman D (1994) Limiting similarity in mechanistic and spatial models of plant  
952 competition in heterogeneous environments. *Am Nat* 143:222–257. doi: 10.1086/285602

953 Pärtel M, Szava-Kovats R, Zobel M (2011) Dark diversity: shedding light on absent species.  
954 *Trends Ecol Evol* 26:124–128. doi: 10.1016/j.tree.2010.12.004

955 Pascual-Hortal L, Saura S (2006) Comparison and development of new graph-based landscape  
956 connectivity indices: towards the prioritization of habitat patches and corridors for conservation.  
957 *Landsc Ecol* 21:959–967. doi: 10.1007/s10980-006-0013-z

958 Patterson BD, Atmar W (1986) Nested subsets and the structure of insular mammalian faunas  
959 and archipelagos. *Biol J Linn Soc* 96:65–82. doi: 10.1007/BF00317508

960 Piessens K, Honnay O, Hermy M (2005) The role of fragment area and isolation in the  
961 conservation of heathland species. *Biol Conserv* 122:61–69. doi: 10.1016/j.biocon.2004.05.023

962 Piessens K, Honnay O, Nackaerts K, Hermy M (2004) Plant Species Richness and Composition  
963 of Heathland Relics in North-Western Belgium: Evidence for a Rescue-Effect? *J Biogeogr*  
964 31:1683–1692. doi: 10.2307/3554767

965 Pollux BJA, Luteijn A, Van Groenendael JM, Ouborg NJ (2009) Gene flow and genetic  
966 structure of the aquatic macrophyte *Sparganium emersum* in a linear unidirectional river.  
967 *Freshw Biol* 54:64–76. doi: 10.1111/j.1365-2427.2008.02100.x

968 Pyke DA, Archer S (1991) Plant–plant interactions affecting plant establishment and persistence  
969 on revegetated rangeland. *J Range Manag* 44:550–557. doi: 10.2307/4003036

970 Rayfield B, Fortin M-J, Fall A (2010) The sensitivity of least-cost habitat graphs to relative cost  
971 surface values. *Landsc Ecol* 25:519–532. doi: 10.1007/s10980-009-9436-7

972 Riibak K, Reitalu T, Tamme R, Helm A, Gerhold P, Znamenskiy S, Bengtsson K, Rosén E,  
973 Prentice HC, Pärtel M (2015) Dark diversity in dry calcareous grasslands is determined by  
974 dispersal ability and stress-tolerance. *Ecography* 38:713–721. doi: 10.1111/ecog.01312

975 Rossetti MR, Tschardt T, Aguilar R, Batáry P (2017) Responses of insect herbivores and  
976 herbivory to habitat fragmentation: a hierarchical meta-analysis. *Ecol Lett* 20:264–272. doi:  
977 10.1111/ele.12723

978 Sandström UG (2002) Green Infrastructure Planning in Urban Sweden. *Plan Pract Res* 17:373–  
979 385. doi: 10.1080/02697450216356

980 Saura S, Pascual-Hortal L (2007) A new habitat availability index to integrate connectivity in  
981 landscape conservation planning: Comparison with existing indices and application to a case  
982 study. *Landsc Urban Plan* 83:91–103. doi: 10.1016/j.landurbplan.2007.03.005

983 Scanlan M (1981) Biogeography of forest plants in the Prairie-Forest Ecotone in Western  
984 Minnesota. In: Burges RL, Sharpe DM (eds) *Forest Island Dynamics in Man-dominated*  
985 *Landscapes*. Springer, Berlin, pp 97–124

986 Schleicher A, Biedermann R, Kleyer M (2011) Dispersal traits determine plant response to  
987 habitat connectivity in an urban landscape. *Landsc Ecol* 26:529–540. doi: 10.1007/s10980-011-  
988 9579-1

989 Simberloff D, Cox J (1987) Consequences and Costs of Conservation Corridors. *Conserv Biol*  
990 1:63–71. doi: 10.1111/j.1523-1739.1987.tb00010.x

991 Soomers H, Winkel DN, Du Y, Wassen MJ (2010) The dispersal and deposition of  
992 hydrochorous plant seeds in drainage ditches. *Freshw Biol* 55:2032–2046. doi: 10.1111/j.1365-  
993 2427.2010.02460.x

994 Sorensen AE (1986) Seed Dispersal by Adhesion. *Annu Rev Ecol Syst* 17:443–463. doi:  
995 10.1146/annurev.es.17.110186.002303

996 Spear SF, Balkenhol N, Fortin M-J, McRae BH, Scribner K (2010) Use of resistance surfaces  
997 for landscape genetic studies: considerations for parameterization and analysis. *Mol Ecol*  
998 19:3576–3591. doi: 10.1111/j.1365-294X.2010.04657.x

999 Stoate C, Boatman N, Borralho R, Carvalho CR, de Snoo GR, Eden P (2001) Ecological  
1000 impacts of arable intensification in Europe. *J Environ Manage* 63:337–365. doi:  
1001 10.1006/jema.2001.0473

1002 Stöcklin J, Winkler E (2004) Optimum reproduction and dispersal strategies of a clonal plant  
1003 in a metapopulation: a simulation study with *Hieracium pilosella*. *Evol Ecol* 18:563–584. doi:  
1004 10.1007/s10682-004-5144-6

1005 Suárez-Esteban A, Delibes M, Fedriani JM (2013) Barriers or corridors? The overlooked role  
1006 of unpaved roads in endozoochorous seed dispersal. *J Appl Ecol* 50:767–774. doi:  
1007 10.1111/1365-2664.12080

1008 Sullivan LL, Johnson BL, Brudvig LA, Haddad NM (2011) Can dispersal mode predict corridor  
1009 effects on plant parasites? *Ecology* 92:1559–1564. doi: 10.1890/10-1116.1

1010 Tackenberg O (2003) Modeling Long-Distance Dispersal of Plant Diaspores by Wind. *Ecol*  
1011 *Monogr* 73:173–189. doi: 10.2307/3100012



1012 Taylor PD, Fahrig L, With KA (2006) Landscape connectivity: a return to the basics. In:  
1013 Connectivity Conservation: maintaining connections for nature. Cambridge University Press,  
1014 Cambridge, pp 29–43

1015 Taylor PD, Fahrig L, Henein K, Merriam G (1993) Connectivity Is a Vital Element of  
1016 Landscape Structure. *Oikos* 68:571–573. doi: 10.2307/3544927

1017 Tewksbury JJ, Levey DJ, Haddad NM, Sargent S, Orrock JL, Weldon A, Danielson BJ,  
1018 Brinkerhoff J, Damschen EI, Townsend P (2002) Corridors affect plants, animals, and their  
1019 interactions in fragmented landscapes. *Proc Natl Acad Sci* 99:12923–12926. doi:  
1020 10.1073/pnas.202242699

1021 Thiele J, Buchholz S, Schirmel J (2017) Using resistance distance from circuit theory to model  
1022 dispersal through habitat corridors. *J Plant Ecol* rtx004. doi: 10.1093/jpe/rtx004

1023 Tilman D, May RM, Lehman CL, Nowak MA (1994) Habitat destruction and the extinction  
1024 debt. *Nature* 371:65–66. doi: 10.1038/371065a0

1025 Tischendorf L, Fahrig L (2000) On the usage and measurement of landscape connectivity.  
1026 *Oikos* 90:7–19. doi: 10.1034/j.1600-0706.2000.900102.x

1027 Tschardt T, Tylianakis JM, Rand TA, Didham RK, Fahrig L, Batáry P, Bengtsson J, Clough  
1028 Y, Crist TO, Dormann CF, Ewers RM, Fründ J, Holt RD, Holzschuh A, Klein AM, Kleijn D,  
1029 Kremen C, Landis DA, Laurance W, Lindenmayer D, Scherber C, Sodhi N, Steffan-Dewenter  
1030 I, Thies C, van der Putten WH, Westphal C (2012) Landscape moderation of biodiversity  
1031 patterns and processes - eight hypotheses. *Biol Rev* 87:661–685. doi: 10.1111/j.1469-  
1032 185X.2011.00216.x

1033 Tzoulas K, Korpela K, Venn S, Yli-Pelkonen V, Kaźmierczak A, Niemela J, James P (2007)  
1034 Promoting ecosystem and human health in urban areas using Green Infrastructure: A literature  
1035 review. *Landsc Urban Plan* 81:167–178. doi: 10.1016/j.landurbplan.2007.02.001

1036 van der Valk AG (1981) Succession in Wetlands: A Gleasonian Approach. *Ecology* 62:688–  
1037 696. doi: 10.2307/1937737

1038 Vellend M, Myers JA, Gardescu S, Marks PL (2003) Dispersal of Trillium seeds by deer:  
1039 implications for long-distance migration of forest herbs. *Ecology* 84:1067–1072. doi:  
1040 10.1890/0012-9658(2003)084[1067:DOTSBD]2.0.CO;2

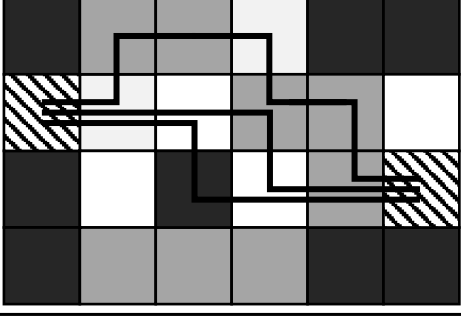
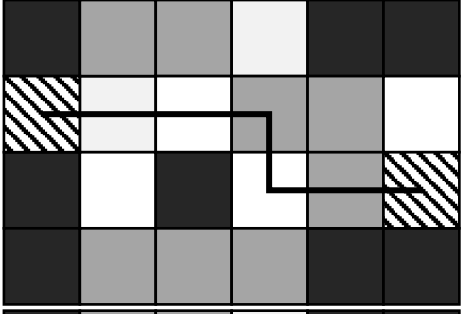
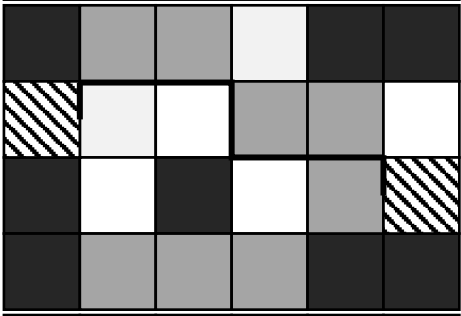
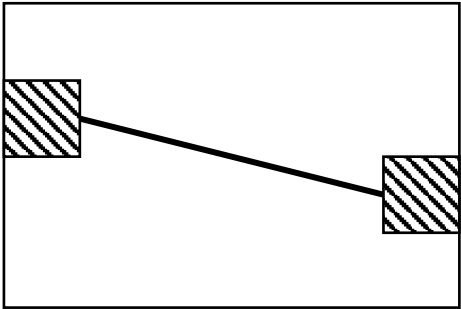
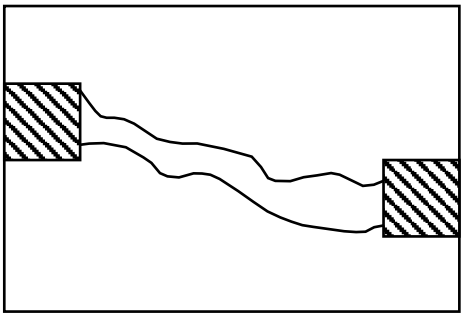
1041 Verheyen K, Vellend M, Calster HV, Peterken G, Hermy M (2004) Metapopulation dynamics  
1042 in changing landscapes: a new spatially realistic model for forest plants. *Ecology* 85:3302–  
1043 3312. doi: 10.1890/04-0395

1044 Violle C, Navas M-L, Vile D, Kazakou E, Fortunel C, Hummel I, Garnier E (2007) Let the  
1045 concept of trait be functional! *Oikos* 116:882–892. doi: 10.1111/j.0030-1299.2007.15559.x

1046 Vittoz P, Engler R (2007) Seed dispersal distances: a typology based on dispersal modes and  
1047 plant traits. *Bot Helvetica* 117:109–124. doi: 10.1007/s00035-007-0797-8  
1048 Weiher E, Keddy PA (1995) The Assembly of Experimental Wetland Plant Communities.  
1049 *Oikos* 73:323–335. doi: 10.2307/3545956  
1050 Wilson DS (1992) Complex Interactions in Metacommunities, with Implications for  
1051 Biodiversity and Higher Levels of Selection. *Ecology* 73:1984–2000. doi: 10.2307/1941449  
1052 Wilson EO, Willis EO (1975) Applied Biogeography. In: Cody ML, Diamond JM (eds)  
1053 Ecology and Evolution of Communities. Harvard University Press, Cambridge, pp 522–534  
1054 Zacharias D, Brandes D (1990) Species Area-Relationships and Frequency: Floristical Data  
1055 Analysis of 44 Isolated Woods in Northwestern Germany. *Vegetatio* 88:21–29. doi:  
1056 10.2307/20038634  
1057 Zeller KA, McGarigal K, Whiteley AR (2012) Estimating landscape resistance to movement:  
1058 a review. *Landsc Ecol* 27:777–797. doi: 10.1007/s10980-012-9737-0  
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1062 **Fig. 1** Methods used to determine how landscape connectivity influences plant dispersal. (A)  
1063 Study of potential dispersal (*i.e.* results of seed bank expression in habitat patches, integrating  
1064 hence the local establishment process). White squares with diagonal lines represent focal habitat  
1065 patches (patches of interest). Solid squares, triangles, circles and rhombus represent different  
1066 plant species. Four measures of connectivity are employed: (1) *Presence/absence of corridors*;  
1067 (2) *Euclidian distance*; (3) *Structural connectivity* and (4) *Potential functional connectivity*.  
1068 White squares represent landscape elements of the same habitat type than the focal habitat  
1069 patches. Grey and black squares represent two different types of habitat patches. To determine  
1070 potential functional connectivity, two different modelling approaches are commonly employed:  
1071 (4\*) *least-cost path* and (4\*\*) *circuit theory*. Both methods require parameterising resistance  
1072 surfaces, with cost value reflecting the cost of species to traverse landscape features. In these  
1073 cases, assigned costs to landscape features increase from white (highly permeable to dispersal)  
1074 to black (slightly permeable to dispersal). (B) Study of actual dispersal (*i.e.* seed rain) White  
1075 squares with diagonal lines represent habitat patches. Solid squares, triangles, circles and  
1076 rhombus represent the seeds of different plant species. One measure of connectivity is  
1077 employed: (5) *Actual functional connectivity*. Thicker arrows represent a higher rate of dispersal  
1078 (*i.e.* higher connectivity), while thinner arrows represent a lower rate of dispersal (*i.e.* lower  
1079 connectivity).

A



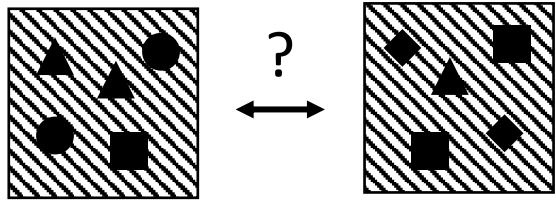
(1)

(2)

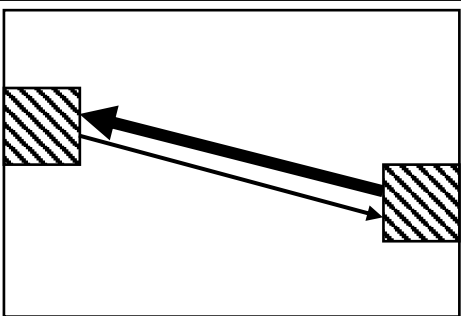
(3)

(4\*)

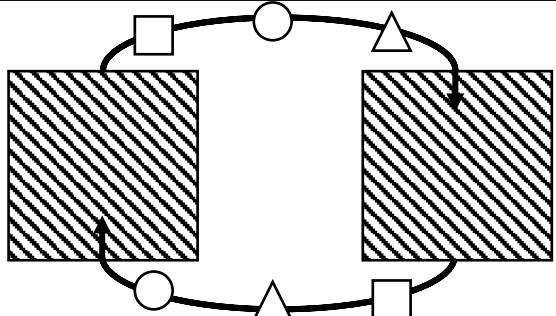
(4\*\*)



B



(5)



### **Box 1** Glossary

**Structural connectivity:** measure of connectivity based on the spatial position of patches independent of the attributes of the organism of interest (Tischendorf and Fahrig 2000; Bennett 2003; Taylor et al. 2006).

**Potential functional connectivity:** measure of connectivity that combines the physical attributes of the landscape with limited information about dispersal ability (Calabrese and Fagan 2004). This measure explicitly considers the behavioural responses of organisms to the physical structure of the landscape (Tischendorf and Fahrig 2000; Bennett 2003; Taylor et al. 2006).

**Least-cost paths:** modelling method to determine potential functional connectivity that assumes a unique path between two patches (represented by a solid line in Fig. 1), hypothesising that individuals perceive their environment in a way that leads them to “select” the optimal path (Adriaensen et al. 2003). The connectivity measure that results from this modelling method is an isolation measure called least-cost distance: the higher the least-cost distance, the lower connectivity.

**Circuit theory:** modelling method to determine to determine potential functional connectivity that evaluates all possible paths (represented by the three different solid lines in Fig. 1), assuming that dispersal follows random walks. The connectivity measure that results from this modelling method is an isolation measure called resistance distance: the higher the resistance distance, the lower connectivity (McRae et al. 2008).

**Actual functional connectivity:** measure of connectivity based on the observation of individuals moving through a landscape (Calabrese and Fagan 2004). This measure explicitly considers the behavioural responses of organisms to the physical structure of the landscape (Tischendorf and Fahrig 2000; Bennett 2003; Taylor et al. 2006).

**Box 2** Example of the Savannah River Site (SRS) experimental design for connectivity studies.

This experimental design is located in the Savannah River Site, a National Environmental Research Park in South Carolina, USA.

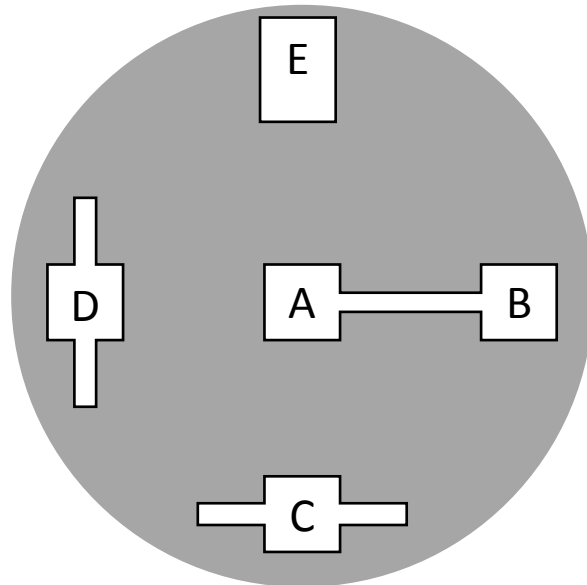
In 2000, eight 50-hectares landscapes were selected. These landscapes were composed of mature (40- to 50-year-old) forest, dominated by loblolly pine (*Pinus taeda*) and longleaf pine (*Pinus palustris*).

Within each landscape, five early-

successional habitat patches were created by cutting and removing all trees, and then burning the cleared areas. The patches included (Fig. 2): one central patch of 1 ha (A) and four peripheral patches (B, C, D, E) that were equal in distance (150 m) from the central patch, but with different structures (see below).

The experiment focused on two corridor functions. First, how a corridor impact connectivity through the following configuration: the central patch is connected to one peripheral patch (B) by a 150 m long and 25 m wide corridor and is not connected to the three other peripheral patches (B, C, D). The three other peripheral patches are equal in area to a patch plus a corridor (1.375 ha). Second, how a corridor impacts patch shape through the following manipulation of peripheral patches: in four landscapes, two of the remaining three peripheral patches were winged (C and D) and one was rectangular (E); in the other four landscapes, two peripheral patches were rectangular and one was winged.

The number of experimental landscapes used varied from six (Damschen et al. 2006) to eight (Tewksbury et al. 2002; Haddad et al. 2003, 2015; Levey et al. 2005; Damschen et al. 2008; Brudvig et al. 2009). This number was not clarified in Haddad et al. (2003).



**Fig. 2** One of the eight experimental landscapes in the SRS.

In our review, only the connectivity function of the corridor is discussed; thus, we only refer to connected patches *versus* unconnected patches.

1082

1083

1084 **Table 1** Synthesis of studies dealing with the effect of current and historical landscape connectivity on functional structure of plant communities.

1085 See Fig.1 for detailed explanations on the connectivity measures.

Approach	Community type	Connectivity measure	Community variable	response	Main results	Reference
Experimental	Early successional vegetation of large pine plantations	Connected vs. unconnected patches (during 7 years)	Difference in species richness	alpha	↗→ for animal dispersed species ↗ for wind dispersed species ↗ for unassisted species	Damschen et al. (2008)
Correlative	Field margins, road verges & ditches	Euclidian distance Potential connectivity: least-cost and resistance distance	Jaccard similarity		∅ for animal and wind dispersed species ↘ for unassisted species	Thiele et al. (2017)
Correlative	Ditches	Structural connectivity: Total ditch length (TDL) Number of culverts (NoC) Number of intersections (NoI) Number of subunits (NoS)	Gamma richness similarity Species trait values: Seed mass Seed buoyancy Seed morphology Seed velocity Seed germination	and	∅ for water-dispersed species  ↘ (TDL); ∅ (NoC); ↗ (NoI) ∅ (TDL); ↗ (NoC, NoI) ↗ species with round seeds (TDL); ↗ species with long seeds (NoC, NoI) ↘ (TDL); ∅ (NoC); ↗ (NoI) ↘ (TDL); ∅ (NoC); ↗ (NoI)	Favre-Bac et al. (2014) Favre-Bac et al. (2017a)
			Standard effect size values calculated for Rao's quadratic entropy and gamma scale weighed mean trait values Seed production Seed buoyancy Seed mass		∅ for water-dispersed species  ↘ (TDL); ∅ (NoC); ↗ (NoI) ∅ (TDL); ↗ (NoC, NoI) ↗ species with round seeds (TDL); ↗ species with long seeds (NoC, NoI) ↘ (TDL); ∅ (NoC); ↗ (NoI) ↘ (TDL); ∅ (NoC); ↗ (NoI) ∅ (TDL, NoC); ↘ Convergence toward the production of small seeds (NoI); ↗ Convergence (NoS) toward the production of small seeds ∅ (TDL; NoI; NoS); ↗ Convergence toward a high buoyancy (NoC) ↗ Divergence (TDL); ∅ (NoC, NoI, NoS);	Favre-Bac et al. (2017b)



Correlative	Temperate forests	Structural connectivity (IFM index)	Trait values of emergent groups to connectivity loss	Sensitive emergent group: Unassisted dispersal Few and heavy seeds High clonality Short seed bank longevity Small height	Resistant emergent group: Assisted dispersal Many and thin seeds No or little clonality Long seed bank longevity Tall	Kolb and Diekmann (2005)
Correlative	Grasslands	Structural connectivity (IFM index)	Species richness of emergent groups	↗ for perennial rosette ∅ for orchids, half-rosette and annuals species		Adriaens et al. (2006)
Correlative	Dry calcareous grasslands	Structural connectivity (IFM index)	Probability of occurrence in response to connectivity	↗ for species with low seed number ∅ for species whatever their dispersal mode (wind-dispersal or not), seed mass or seed bank longevity ↗ for short-living species ∅ for clonal species		Evju et al. (2015)
Correlative	Heathlands	Structural connectivity (IFM index)	Species richness in the different categories of the seed longevity index	↗ for species with short seed bank longevity		Piessens et al. (2004)
Correlative	Heathlands	Structural connectivity (IFM index)	Regression coefficient resulting from the logistic regression between patch occupancy for each species as explained by connectivity intensity	↗ for species with short seed bank longevity ∅ for species whatever their mean plant height, seed mass, dispersal mode, growth form, self-compatibility, clonal growth form and seed number		Piessens et al. (2005)

Correlative	Temperate forests	Structural connectivity (modified version of IFM index that incorporates patch age)	Regression coefficient resulting from the logistic regression between patch occupancy for each species as explained by connectivity intensity	↗ for vertebrate and wind-dispersed species ↗ for species with low seed production ↗ for species with small height ∅ for species whatever their life cycle duration	Verheyen et al. (2004)
Correlative	Urban - ruderals	Structural connectivity (IFM index)	Trait values of response groups to connectivity: Terminal velocity Number of seeds Seed mass	Sensitive > resistant species Sensitive < resistant species Sensitive = resistant species	Schleicher et al. (2011)
Correlative	Grasslands	Current (C) and historical (H) structural connectivity (IFM index) 1 km radius 2 km radius	Proportion of species	∅ for animal-, wind- dispersed and unassisted species ∅ for species whatever their seed size ∅ (C); ↘ (H) for long seed bank persistence species ↗ (C); ∅ (H) for annual plants ∅ (C); ↘ (H) for perennials with or without clonality at 1 km scale. ∅ (C) for perennials with or without clonal ability at 2 km scale ↘ (H) for perennials without clonal ability at 2 km scale	Lindborg (2007)

1087 **Effect of landscape connectivity on plant communities: a review of response patterns**

1088

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1094 **Appendix S1 Methods**

1095

1096 We systematically reviewed articles to assess whether and how landscape connectivity affects  
1097 the assembly of plant communities. Using Web of Science and Google Scholar to gather  
1098 publications using the following keywords: landscape connectivity, habitat isolation, corridors  
1099 and plants. Because ecosystem functioning is completely different in tropical systems, we  
1100 excluded these studies, only considering studies conducted in temperate systems. We also  
1101 excluded studies that focus on fragmentation (*i.e.* that are conducted in a fragmented landscape  
1102 context); that is, when it is not possible to disentangle the effects of decreased connectivity  
1103 (variable of interest) and decreased habitat area. We only considered studies that: i) manipulated  
1104 connectivity through the presence (or absence) of an experimentally-created corridor between  
1105 two habitat patches for experimental approaches, and ii) measured connectivity in a quantitative  
1106 manner (*via* the use of indices) for correlative approaches. Quantitative connectivity  
1107 assessments were based on isolation metrics such as Euclidean distance, structural connectivity  
1108 indices (see various examples in the main text), least-cost distance or resistance distance. We  
1109 selected studies that focused on how landscape connectivity affected plants at the community  
1110 level. Within this framework, we included studies with taxonomic approaches (*i.e.* looking at  
1111 the composition and diversity of plant communities) and trait-based approaches (*i.e.* looking at  
1112 the composition and diversity of traits). These studies assessed plant community response  
1113 through: i) analysis of individual species responses across most species at the community level,  
1114 and ii) aggregated measures across most species of a community. More specifically, only the  
1115 following measures were extracted from articles that provided: i) taxonomic and ii) trait-based  
1116 approaches:

1117

1118 i) Taxonomic approaches: alpha diversity (species richness, species density or species  
1119 abundance), beta diversity (Jaccard similarity index) and gamma diversity (gamma  
1120 species richness)

1121 ii) Trait-based approaches:

1122 a. Dispersal mode approach: analyses similar to the taxonomic approach but per  
1123 dispersal mode, proportion of species per dispersal mode and indicators for  
1124 species sensitivity to connectivity (probability of occurrence in response to  
1125 connectivity, regression coefficient resulting from the logistic regression  
1126 between patch occupancy for each species as explained by connectivity  
1127 intensity)

1128 b. Emergent groups approach: trait values of emergent groups and species richness  
1129 of emergent groups

1130 c. Response traits approach: indicators for species sensitivity to connectivity  
1131 (probability of occurrence in response to connectivity, regression coefficient

1132 resulting from the logistic regression between patch occupancy for each species  
1133 as explained by connectivity intensity), species richness in the different  
1134 categories of the seed longevity index, proportion of species characterised by a  
1135 trait, trait value of response groups to connectivity and species trait values  
1136 d. Community weighted mean traits and functional diversity approach: functional  
1137 gamma diversity (standard effect size values calculated for Rao's quadratic  
1138 entropy traducing functional divergence or convergence) and gamma aggregated  
1139 mean trait values (gamma scale weighed mean trait values, indicating the mean  
1140 value towards which traits converged)

1141

1142 Concerning actual dispersal approaches, we included studies at the species level because very  
1143 few studies have assessed the actual dispersal of communities in relation to connectivity.

1144