



HAL
open science

Kin selection, kin recognition and kin discrimination in plants revisited: A claim for considering environmental and genetic variability

Lucas Mazal, Alex Fajardo, Irène Till-Bottraud, Dov Corenblit, Boris Fumanal

► **To cite this version:**

Lucas Mazal, Alex Fajardo, Irène Till-Bottraud, Dov Corenblit, Boris Fumanal. Kin selection, kin recognition and kin discrimination in plants revisited: A claim for considering environmental and genetic variability. *Plant, Cell and Environment*, 2023, 46 (7), pp.2007 - 2016. 10.1111/pce.14584 . hal-04243901

HAL Id: hal-04243901

<https://hal.science/hal-04243901v1>

Submitted on 16 Oct 2023

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

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

1 **Kin selection, kin recognition and kin discrimination in plants revisited: a**
2 **claim for considering environmental and genetic variability**

3 *Lucas Mazal¹, Alex Fajardo², Irène Till-Bottraud¹, Dov Corenblit¹, Boris Fumanal³*

4 *¹Université Clermont Auvergne, CNRS, GEOLAB, F-63000 Clermont-Ferrand, France.*

5 *² Instituto de Investigación Interdisciplinaria (I³), Vicerrectoría Académica, Universidad de Talca,*
6 *Campus Lircay, Talca 3460000, Chile*

7 *³ Université Clermont Auvergne, INRAE, UMR 547 PIAF, F-63000 Clermont-Ferrand, France.*

8
9 Corresponding author Lucas Mazal, lucas.mazal@uca.fr

10 ORCID-ID: 0000-0002-1991-5986

11
12 **Funding:** The first author was funded for his PhD by the French Ministry of National
13 Education, Higher Education and Research. This research was supported by the French
14 government IDEX-ISITE initiative 16-IDEX-0001 (CAP 20-25).

15 **Conflicts of interest/Competing interests:** The authors declare that they have no conflict of
16 interest.

17 **Authors' contributions:** All authors contributed to the writing of the paper and all authors read
18 and approved the final manuscript.

19 **Keywords:** plant-plant interactions, cooperative behaviour, long-lived species, intraspecific
20 variation, context-dependency.

21 **Abstract**

22 Considering kin selection in the study of interacting conspecific plants broadens our vision of
23 plant behaviour and brings arguments to explain plant-plant positive interactions. These
24 interactions are the subject of abundant research in community ecology and the role of
25 relatedness in interactions between individuals has become a hot topic. Indeed, the past decade
26 has seen a steady accumulation of exciting but also controversial results regarding the behaviour
27 of plants in the presence of genetically related neighbours, i.e., kin discrimination. In addition
28 to the existence of some methodological and semantic aspects that should be strengthened to
29 accurately reveal the occurrence of kin recognition or selection, we claim here that some key
30 aspects must be considered to increase our ability to detect kin recognition and selection in
31 plants. These aspects include intraspecific variability, the fact that plant-plant interactions are
32 context-dependent, and the lifespan of species (particularly long-lived ones). Additionally, we
33 note that the study of the population spatial genetic structure (SGS) could be a useful tool to
34 identify candidate settings for the study of kin recognition and selection in plants.

35

36 **Introduction**

37 Plants are sessile organisms and cannot chose or move away from their neighbours, but are tied
38 to them throughout their life. In this context, identifying the factors that can influence the type
39 of interactions (positive, neutral or negative) between individuals is of prime importance. One
40 such factor that has become increasingly popular is genetic relatedness between individuals.
41 Recent research has shown compelling evidence that plants are able to detect their close
42 neighbours and actively interact with them. Indeed, plants exhibit complex recognition and
43 communication mechanisms (Bilas, Bretman & Bennett 2021). Classically studied between
44 different plant species, these interactions also apply at the intraspecific level (Bilas, Bretman &
45 Bennett 2021). Remarkably, plants can distinguish their conspecific neighbours based on their
46 genetic relatedness (i.e., kin recognition) and adjust their 'behaviour' accordingly (kin
47 discrimination) (Ehler & Bilde 2019, Anten & Chen, 2021a). Being able to identify related
48 individuals could be advantageous to reduce competition among relatives (File, Murphy &
49 Dudley 2012), a cooperative behaviour that can be promoted through kin selection. However,
50 traits that reduce competition can be costly for the individual (e.g., limiting resource acquisition;
51 Dudley, Murphy & File, 2013). Kin selection theory (Hamilton 1964) states that an altruistic
52 trait can be favoured by natural selection when the altruist and the recipient are related. This is
53 based on the concept of indirect fitness, i.e., indirect benefits accrued by the altruistic individual
54 through the fitness of the recipient, as the two individuals share genes. If individuals are
55 sufficiently closely related, altruism can favour the reproduction of the set of genes they share
56 despite losses in direct fitness by the altruist individual (Axelrod & Hamilton 1981). Note that
57 kin recognition and kin discrimination are not necessary for kin selection to occur (e.g., in
58 viscous populations, cooperation with neighbours is mostly directed towards kin; West et al.,
59 2007).

60 In a recent review, Anten & Chen (2021a) provide a critical and detailed overview of the
61 conceptual bases and latest results regarding kin recognition, kin discrimination and kin
62 selection in plants. In summary, they state that most of the studies have so far explored plant-
63 plant interactions at the root level. A frequently observed result is that plants growing with
64 related individuals show a reduction in belowground biomass compared to plants growing with
65 strangers (usually seeds from different mothers), which is interpreted as a reduction in
66 competition towards relatives, possibly leading to positive interactions (File, Murphy & Dudley
67 2012). The possible mechanisms of belowground kin recognition are root-exudates and
68 mycorrhizal networks. Although some methodological concerns are discussed by Anten &

69 Chen (2021a), the authors conclude that there is sufficient evidence for supporting kin
70 recognition in plants.

71 Anten & Chen (2021a) accomplished the best update so far regarding kin discrimination
72 and kin selection in plants. We think, nevertheless, that some aspects were not considered in
73 their review that are of great complementary importance to broaden the kin recognition and
74 selection framework: intraspecific variability, environmental context dependency, plant
75 longevity and spatial patterns of plant interactions. We expand on these underexplored aspects,
76 which we consider essential to build a testable framework about kin discrimination and kin
77 selection and expose two case studies that exemplify these new aspects. We then develop two
78 additional points:(i) several papers confuse (or use alternatively) the concepts of kin
79 recognition, kin discrimination and kin selection for which we offer a clarification; (ii) this
80 confusion often leads to methodological issues and we propose methodological guidelines for
81 experimental designs to detect kin discrimination and assess kin selection.

82

83 **Should we all cooperate? The neglected role of genetic diversity for kin interaction** 84 **mechanisms**

85 Several published studies, including Anten & Chen's (2021a) review, have concluded that kin
86 recognition is either present or absent in a given species. However, do we expect species to be
87 entirely homogeneous with regard to kin discrimination or even kin selection? In other words,
88 is it reasonable to assume that all individuals of the same species exhibit the same 'behaviour'?
89 Probably not, as already suggested by several authors (Clark 2010; Bolnick et al., 2011;
90 Westerband, Funk & Barton 2021). For example, it is recognised that intraspecific diversity in
91 plants shapes biotic and abiotic interactions and plays a key role in determining the performance
92 of individuals (Westerband, Funk & Barton 2021). Intraspecific diversity is the rule for a wide
93 range of traits (Siefert et al., 2015), it is therefore conceivable that not all individuals within a
94 species possess the same traits associated with kin recognition, and that not all individuals have
95 the (same) ability to distinguish their related neighbours. Moreover, our understanding of the
96 genetic and molecular bases underlying the natural variation of plant-plant interactions remains
97 limited and needs to be considered in further research (Subrahmaniam et al., 2018). We assert
98 here that intraspecific variability for kin recognition has been poorly addressed thus far,
99 although it represents a critical issue.

100 Theoretical models exploring kin recognition in various organisms (mainly on animals,
101 but the results are valid for all species capable of kin recognition) predict a wide range of
102 intraspecific variation in kin recognition mechanisms between and within populations (see
103 review by Penn & Frommen 2010), although other models do not predict kin recognition
104 mechanisms at equilibrium because these mechanisms are costly (Rousset & Roze 2007). We
105 argue that kin recognition in plants could still be an advantageous local strategy in certain
106 conditions while not being advantageous when the population reaches equilibrium. Thus,
107 intraspecific diversity for kin interaction mechanisms should be particularly important to
108 consider. Anten & Chen (2021a) and also Fréville et al. (2021) discussed this point in the case
109 of crops, which exhibit lower genetic diversity than wild species and where individuals interact
110 mostly with kin and proposed that studies include sampling replication at the genotypic level to
111 capture sufficient intraspecific variation in kin discrimination. Most studies have used only a
112 few different kin groups (usually two to four) and thus captured only a small part of the existing
113 variability in the species. Yet, some recent studies have made the effort to include a larger
114 number of kin groups (eight-ten and up to 28 kin groups) (Semchenko, Saar & Lepik 2017;
115 Torices, Gómez & Pannell 2018; Takigahira & Yamawo 2019). To overcome this important
116 issue, which can lead to erroneous conclusions at the species level, we advocate the use of at
117 least five to ten kin groups originating from several populations. The logic is as follows: the
118 probability of detection depends on the frequency of the trait. If one in five individuals in a
119 population can discriminate kin, using five kin groups there is still a 33% chance of missing the
120 trait (i.e., reaching erroneous conclusions), and this value is reduced to 11% with ten kin groups.
121 If the frequency is one in ten, the chances are 60% and 35%, respectively.

122

123 **Under which conditions is it beneficial to cooperate with relatives?**

124 Because plants are sessile organisms, they cannot move to avoid their neighbours' effects nor
125 environmental stress. Local adaptation (i.e., selection for traits that increase the fitness of
126 individuals in their local environment) and phenotypic plasticity (i.e., the ability of a given
127 genotype to express different phenotypes according to some environmental cues or
128 circumstances) are the two common processes to overcome these threats (e.g., de Villemereuil,
129 Mouterde, Gaggiotti & Till-Bottraud 2018; Fox, Donelson, Schunter, Ravasi & Gaitán-Espitia
130 2019). Phenotypic variability can thus be induced by variation in the local conditions and could
131 influence the nature of plant-plant interactions (Callaway, Pennings & Richards 2003). As a
132 consequence, should we expect the same type of interactions between individuals living under

133 different conditions? The stress gradient hypothesis (SGH, Bertness & Callaway 1994) predicts
134 that the level of stress will influence the type of interaction between individuals shifting from
135 positive (i.e., facilitation) in stressful environments to negative (i.e., competition) in favourable
136 environmental conditions (Bertness & Callaway 1994). Facilitation is defined as ‘an interaction
137 in which the presence of one species alters the environment in a way that enhances the growth,
138 survival and reproduction of a second species’ (Bronstein 2009, p.1160). Interactions show
139 ‘conditionality’ when costs and benefits, and thus outcomes, are affected in predictable ways
140 by ecological conditions (Bronstein 1994). Plant interactions thus depend on the ecological
141 settings in which they take place, and in particular, plants adjust to the level of stress (Bilas,
142 Bretman & Bennett 2021). It has been implicitly assumed that facilitation only occurs between
143 individuals of different species, but as shown and discussed by Fajardo & McIntire (2011),
144 facilitation can be found between individuals of the same species and was indeed suggested in
145 several other cases (Proença et al., 2019; Reijers, Akker, Cruijsen, Lamers & Heide 2019).
146 Facilitation allows, or even favours, the coexistence of individuals of the same species in close
147 proximity, i.e., within their zone of interaction (ZOI *sensu* Stoll & Weiner 2000). If the ZOI of
148 related individuals overlaps this would allow kin recognition, kin discrimination and kin
149 selection mechanisms to operate. Following SGH predictions, harsh environmental conditions
150 should lead to stronger positive interactions (i.e., cooperation) if related individuals are present.
151 We can thus expect contrasting responses for individuals living in stressful versus benign
152 environments and thus, in a changing environment, the same closely related individuals could
153 cooperate or compete depending on the level of stress.

154 So far, few studies have shown evidence that kin recognition depends on the level or the
155 type of environmental stress. In these studies, kin recognition and discrimination responses
156 were triggered by several abiotic factors including nutrient availability, for example, in *Glycine*
157 *max*, *Caesalpinia pulcherrima*, *Populus tomentosa* (Li, Xu & Liu 2018), and *Pisum sativum*
158 (Pezzola, Pandolfi & Mancuso 2020), heavy metal concentrations in *Sorghum vulgare* (Li, Xu
159 & Feng, 2018) and biotic factors such as the presence of another species in *Medicago* genera
160 (Ehlers, David, Damgaard, & Lenormand 2016). On the other hand, water availability, did not
161 trigger kin recognition responses in *Glechoma hederacea* (Goddard, Varga, John & Soulsbury
162 2020). Thus, it is possible that kin recognition is only expressed in a given kin group under
163 certain ecological conditions, which can only be identified by modulating the type of stress and
164 its intensity. We note then that it is essential to include both several stressors and several

165 genotypes (kin groups) in any experimental effort to elucidate kin recognition/selection in
166 plants.

167 Kin discrimination also depends on whether cooperation is obligate or facultative. In the
168 case of obligate cooperation, kin recognition is ultimately not useful as individuals always
169 cooperate whatever the identity of their neighbour is and there is no discrimination. In contrast,
170 facultative cooperation implies that individuals exhibit phenotypic plasticity for cooperation
171 and that they are able to distinguish relatives (i.e., present kin recognition). Both phenotypic
172 plasticity (Ghalambor et al., 2007) and kin recognition (Rousset & Roze 2007) have associated
173 costs. To identify cases where obligate or facultative cooperation is favoured requires thus to
174 estimate the costs and benefits of facultative cooperation (plasticity) in addition to those of
175 Hamilton's equation (Hamilton 1964), weighted by the probability of interacting with a relative.
176 Therefore, in the case of facultative cooperation, the question is not only with whom but also
177 under what conditions should individuals cooperate? In a stable environment, the phenotype is
178 constant and obligate cooperators would be favoured. In a changing environment, SGH suggests
179 that interactions shift between positive and negative and being a facultative cooperator would
180 be advantageous. In the latter case, cooperation will depend on the relatedness between
181 individuals. The important point here is that environmental conditions in which the experiment
182 takes place, but which are seldom considered in kin studies, condition the detection of kin
183 recognition/selection.

184

185 **Cooperating throughout life? Interactions in long-lived species**

186 For practical reasons, most of the studies on kin recognition thus far have focused on annual or
187 short-lived plants (Subrahmaniam, Roby & Roux 2021). Few studies have investigated long-
188 lived species such as perennial herbs, shrubs or even trees (Till-Bottraud, Fajardo & Rioux
189 2012; Fajardo, Torres-Díaz & Till-Bottraud 2016; Pickles et al., 2017), and mostly addressed a
190 short period of their lives (e.g., the seedling stage; Takigahira & Yamawo 2019; He et al., 2021).
191 Some authors reported positive interactions between mature related individuals following
192 herbivory, through discrimination of volatile chemicals emitted from relatives (Karban et al.,
193 2014; Hussain, Rodriguez-Ramos & Erbilgin 2019). Other showed that related seedlings grow
194 better in the presence of kin (Assay et al., 2020) and benefit more from the mycorrhizal network
195 shared with related neighbours (Pickles et al. 2017).

196 However, long-lived species exhibit several interesting characteristics that have not
197 been much considered: (i) the long lifespan means that an individual will likely experience
198 different environmental situations during its life; (ii) different life stages may have very
199 different ecologies (ontogenetic stages, e.g., seedling *vs.* mature tree, related to the regeneration
200 niche, Grubb 1977), kin discrimination and kin selection might thus be advantageous at a given
201 developmental stage but not at others. For example, at early stages when seedling mortality is
202 important because they are more sensitive to harsh conditions, positive interactions between
203 related individuals would ensure better survival of related genotypes; (iii) some long-lived
204 species, particularly trees, strongly affect their physical environment by improving their living
205 conditions (i.e., the ecosystem engineering effect *sensu* Jones, Lawton & Shachak 1994),
206 modulating the level of stress and improving resource availability through their effects on the
207 local environment (McIntire & Fajardo 2014). As this effect is incremental through time it will
208 be all the more important in long-lived species (Hortobágyi, Corenblit, Steiger & Peiry 2018).
209 As a given individual can experience different conditions through its lifetime, the chance of
210 detecting kin recognition could strongly depend on the life stage considered. This particular
211 point is essential to consider when conducting field studies where local conditions can change
212 or even be improved by engineering species. All these three characteristics are linked to the
213 SGH, although the gradient here is temporal as long-lived species interact with their neighbours
214 through decades or even centuries in a fluctuating or changing micro- and macro-environment.

215 Moreover, in seasonal environments, the physiological activity is reduced or potentially
216 halted by dormancy periods. Is this interaction between two long-lived individuals just one
217 absolute interaction or is it repeated interactions (e.g., in the different growing seasons), which
218 would lead to radically different outcomes (Axelrod & Hamilton 1981)? This is still an open
219 question, for which evolutionary game theory (Maynard Smith 1982) could provide a
220 theoretical framework.

221

222 **Spatial patterns of plant interactions: the added value of spatial genetic structure**

223 One potentially valuable tool to search for positive interactions among kin in natural
224 populations is the spatial genetic structure (SGS). The SGS, defined as the non-random spatial
225 distribution of genotypes, characterizes the relationship of relatedness estimates between pairs
226 of individuals and their physical proximity (Loiselle, Sork, Nason & Graham 1995; Hardy
227 2003). The genetic composition of a population has an influence on the type of biotic interaction

228 that can occur among immediate neighbours (Fajardo, Torres-Díaz & Till-Bottraud 2016).
229 Using a spatial genetic approach can thus offer complementary insights, particularly for *in situ*
230 studies. A significant SGS indicates that related individuals coexist close to each other, and
231 thus, the possibility of kin interactions increases. This makes SGS an interesting tool to detect
232 the presence of related individuals and to consider potential positive interactions in the context
233 of kin recognition and selection. Moreover, this SGS approach seems particularly relevant to
234 us as kin selection can evolve in viscous populations without the need for kin recognition or
235 discrimination (e.g., obligate cooperation) (Waldman 1988, Gardner & West, 2004). However,
236 the fact that individuals are “close to each other” does not mean that they will necessarily
237 interact. Plant–plant interactions are inherently local in nature. They mostly occur among
238 immediate neighbours, in which case an individual’s direct effect on another (negative or
239 positive) will have a delimited influence in space, i.e., the ZOI (Stoll & Weiner 2000). The ZOI
240 will grow as individuals grow because individuals, in particular trees, will occupy a larger area
241 with their aerial and belowground organs. Thus, over their lifetime, individuals will interact
242 with other individuals located further away as their ZOI expands, and thus allowing more or
243 different individuals to interact through time. Biotic interactions will be influenced by the local
244 SGS (i.e., the presence of related individuals within the ZOI) but can also significantly influence
245 the SGS within the ZOI of individuals through the survival of related individuals in local
246 patches, which can be monitored through time. However, very few studies have linked the SGS
247 and the ZOI when interactions between related individuals take place (Till-Bottraud, Fajardo &
248 Rioux 2012; Segovia, Vásquez, Faugeron & Haye 2015; Fajardo, Torres-Díaz & Till-Bottraud
249 2016).

250

251 **The relationship between the SGS and intraspecific interactions: two case studies in long-** 252 **lived species**

253 To illustrate how all the aspects we have outlined above could help us assess the ultimate
254 importance of kin selection, we describe here two examples of positive interactions between
255 related individuals in tree species.

256 First, in *Nothofagus pumilio* (Nothofagaceae) second-growth stands located in the
257 Chilean Patagonia, multi-stemmed trees are formed by related individuals. They are restricted
258 to forest edges where stress (wind and desiccation) was very high at the regeneration stage, and

259 where facilitation increased seedling survival (Fajardo & McIntire 2011) and growth in multi-
260 stemmed adult trees (Fajardo & McIntire 2010; McIntire & Fajardo 2011). Trees from different
261 multi-stemmed clusters are unrelated, and no group dispersal mechanism is reported for this
262 species (Till-Bottraud, Fajardo & Rioux 2012; Fajardo, Torres-Díaz & Till-Bottraud 2016)
263 indicating better survival when related seedlings happen to grow together. The ecological
264 mechanism involved here may be an altruistic or cooperative fusion of trunks, which would be
265 physiologically and biomechanically selectively advantageous for resource acquisition or
266 stability against wind (Tarroux & DesRochers 2011; Tarroux, DesRochers & Tremblay 2014;
267 Fajardo, Torres-Díaz & Till-Bottraud 2016), but fusion may also involve some costs (e.g.,
268 resource sharing, pathogen transmission). Stem merging scales up the level of organization to
269 the group and therefore competition does not occur among individuals but among groups of
270 trees (McIntire & Fajardo, 2011). In a kin selection framework, fusions would be selected and
271 would only occur between related individuals, as indirect fitness gains might offset these costs.

272 The other example is the Black Poplar (*Populus nigra* L. Salicaceae) in a riparian
273 context. At the establishment phase, *P. nigra* seedlings grow in high densities on exposed
274 alluvial bars of rivers, and facilitation is the major occurring interaction (Corenblit et al. 2018).
275 Young individuals form dense patches and mechanically protect each other from destruction
276 (Corenblit et al., 2014). During colonisation, *P. nigra* is known to enhance its local conditions
277 (i.e., ecosystem engineer species) when growing at high seedling densities by trapping fine
278 sediments and thus increasing nutrient availability and decreasing drought stress in its natural
279 river reach habitat (Corenblit et al., 2014; Hortobágyi, Corenblit, Steiger & Peiry 2018). The
280 modification of the environment by individuals can prevent, delay, or accelerate changes in the
281 types of interactions between individuals (Corenblit et al., 2018). Based on the evolution of the
282 SGS in a natural population, a shift in the type of interactions from positive to negative has
283 been proposed in *P. nigra* stands along riparian corridors (Mazal et al., 2021; Mazal, Corenblit,
284 Fumanal & Till-Bottraud 2022) (Fig.1). The case of *P. nigra* shows that, in long-lived species,
285 shifts in the type of interaction are possible and could be monitored by studying SGS (Mazal,
286 Corenblit, Fumanal & Till-Bottraud 2021). A cohort monitoring approach from establishment
287 to maturity in long-lived species could be an efficient approach to follow the evolution of
288 interactions between individuals and ultimately to estimate the survival component of fitness
289 (not just reproduction). In addition, cohorts should also be followed in contrasted conditions, if
290 possible along a gradient of environmental conditions, as the type of interaction may vary with
291 life stages and environmental conditions. An increase in SGS over time could be an indication

292 of positive intraspecific interactions between relatives and thus would warrant further
293 validation.

294 These examples open new research avenues to consider the impact of genetic variability,
295 environmental conditions, life-span and SGS in kin selection studies. However, we believe that
296 some semantic and methodological aspects need clarification to accurately address kin
297 recognition, discrimination or selection in such studies.

298

299 **Kin recognition, kin discrimination and kin selection**

300 Kin recognition, kin discrimination and kin selection are often used alternatively in articles,
301 however, they correspond to different concepts (see the review on kin recognition definitions
302 by Penn & Frommen 2010). The confusion between kin recognition and kin discrimination
303 probably comes from the fact that the only way to detect kin recognition in plants is through its
304 expression (trait or behaviour variation between kin and non-kin neighbour treatments), i.e., kin
305 discrimination. Indeed, many studies that showed the expression of a different phenotype
306 depending on the genetic identity of the neighbours used the term kin recognition but *de facto*
307 showed kin discrimination (Dudley & File 2007a, 2007b; Milla, Forero, Escudero & Iriondo
308 2009; Murphy & Dudley 2009; Lepik, Abakumova, Zobel & Semchenko 2012; Semchenko,
309 Saar & Lepik 2014; Yang, Li, Xu & Kong 2018; Li, Xu & Liu 2018; Goddard, Varga, John &
310 Soulsbury 2020). Although kin discrimination can be captured by measuring any kind of trait
311 that shows plasticity between kin and non-kin neighbours, its implication for kin selection can
312 only be inferred if it translates into fitness differences between the two treatments (Ehlers &
313 Bilde 2019). However, fitness is very rarely or only approximatively measured in “kin
314 recognition” studies and some authors tend to extrapolate their results on kin recognition or
315 discrimination and conclude without proof that there is evidence of kin selection in plants
316 (Crepy & Casal 2015; Zhang et al., 2016; Pezzola, Pandolfi & Mancuso 2020).

317 Studies using biomass-related metrics (Fig. 2a) often suggest that reducing below-
318 ground biomass is a cooperative behaviour indicative of kin selection. Increasing below-ground
319 biomass when growing with neighbours (a competitive trait) diverts resources from producing
320 more above-ground biomass. If the above-ground biomass is a good proxy for fitness
321 (Younginger, Sirová, Cruzan & Ballhorn 2017), a cooperative decrease in below-ground
322 biomass of all interacting individuals could thus increase their fitness (both direct and indirect
323 if kin are involved). However, the relationship between biomass and fitness is not always

324 correct, especially in long-lived species such as trees, where biomass provides little information
325 on how individuals will reproduce or may reflect many confounding factors, making their
326 interpretation difficult (see the methodological issues discussed below and in Fig.2). In fact,
327 results based on the observation of biomass differences can be interpreted in opposite ways
328 (Subrahmaniam, Roby & Roux 2021). To our knowledge, the only article that came close to
329 demonstrate a possible gain in fitness for closely related individuals (through the study of
330 reproductive traits) is Torices et al. (2018). They showed that *Moricandia moricandioides*
331 (Brassicaceae) individuals growing with kin produced larger floral displays than those growing
332 with strangers. The production of a larger floral display is costly but attracts more pollinators
333 that will also benefit their neighbours. The authors argue that individuals compensate for the
334 decrease in direct fitness caused by the production of large flowers with a gain in indirect fitness
335 by attracting more pollinators who will visit their related neighbours (Torices, Gómez & Pannell
336 2018). In this case, related neighbours benefit from the attraction of pollinators that will
337 pollinate their stigmas and export their pollen. These results were further supported by a second
338 study which showed that plants growing with kin invest more in petal pigments, nectar volume
339 and sugar content (Torices, DeSoto, Narbona, Gómez & Pannell, 2021).

340 Following the approach proposed by Ehlers & Bilde (2019), we recommend to move on
341 to the next step and to study kin selection *per se* generalizing the use of Hamilton's rule ($rB >$
342 C), and the measurement of the cost (C) and benefit (B) components of fitness separately on
343 each plant that take part in the interaction (note that, contrarily to what is stated in Ehlers &
344 Bilde (2019), only *interacting* related individuals should be included for the estimation of
345 indirect fitness), as well as their true (realized) relatedness (r), which can now be reliably
346 estimated using next-generation sequencing techniques (de Villemereuil, Gaggiotti, Mouterde
347 & Till-Bottraud 2016). Two particular cases need to be detailed: if $r = 0$, no indirect fitness can
348 be obtained. Conversely, if $r = 1$, Hamilton's rule is no longer needed to explain the outcome
349 of the interactions because each interactant is actually a clone of the same individual and the
350 indirect fitness is actually a direct fitness. This is the case in studies where clones are used as
351 kin groups (Yamawo, Sato & Mukai 2017). The case of highly self-pollinating species (i.e.,
352 same accession or variety) (Simonsen, Chow & Stinchcombe 2014; Yang, Li, Xu & Kong 2018;
353 Pezzola, Pandolfi & Mancuso 2020) is slightly different because although very close to 1, r is
354 actually slightly lower than 1. In a literature survey Subrahmaniam et al. (2021) showed that
355 44% and 10% of the studies that investigated interactions between related and unrelated
356 individuals were conducted in self-pollinating or clonal species, respectively. Can we extend

357 the results of these studies to other species that are not self-pollinated, and how general is a
358 mechanism if it is only shown to work for clonal or self-pollinating species? Although the cases
359 where r is very close to 1 are convenient to study, we think that they are rather extreme cases
360 and that they are therefore difficult to generalize to other species. In natural populations without
361 inbreeding, the expected relatedness full-sibs is 0.25, and values measured for ants and bees
362 where kin selection has been demonstrated range from 0.5 to 0.7 (Gadagkar et al. 2019; Kay,
363 Lehmann & Keller 2019). We argue that results obtained from highly-selfing or clonal species
364 should be interpreted in the same way as studies on self-recognition (i.e. recognition of
365 individual genotype between two ramets, be they connected or disconnected) (see also Bilas,
366 Bretman & Bennett 2021).

367 As underlined by Hamilton's rule, the level of relatedness between individuals is critical
368 to predict the outcome of the interactions. In a local population, varying levels of genetic
369 relatedness can co-exist and thus patterns of interactions between individuals can vary based on
370 their relatedness (Subrahmaniam, Roby & Roux 2021). Following Hamilton's rule, the potential
371 benefits of interaction should decrease linearly or display a threshold (Till-Bottraud & de
372 Villemereuil 2016) beyond which it is no longer advantageous to cooperate ($C > rB$). However,
373 the gradient of relatedness between individuals is rarely addressed in studies and this raises the
374 question of the generalized use of studies that tested interactions between individuals
375 originating from a single population (Subrahmaniam, Roby & Roux 2021). Few studies thus
376 far have used different levels of relatedness between individuals. Karban et al. (2014), for
377 example, used a gradient of relatedness and found that *Artemisia tridentata* individuals
378 responded more effectively to volatile cues from close relatives than from distant relatives when
379 subjected to herbivory. Another study using rice cultivars with different levels of relatedness
380 showed that plants altered their root behaviour and biomass allocation in response to relatedness
381 level (Yang, Li, Xu & Kong 2018). Moreover, using a relatedness gradient, Xu et al., (2021)
382 showed that plants discriminate their neighbours (related or unrelated) and adjust their growth,
383 competitiveness and chemical defence accordingly to improved weed suppression in rice
384 (through allelopathic processes) (see also commentary from Anten & Chen 2021b). Following
385 on the authors' discussion, we also believe that a relatedness gradient is an important factor that
386 can potentially trigger responses when plants interact with neighbouring individuals and thus
387 must be considered in future studies.

388

389

390

391

392 **Methodological challenges in studying kin recognition and selection**

393 Whether studies did (see the review by Anten & Chen 2021a) or did not find evidence of kin
394 recognition (e.g., Milla, Forero, Escudero & Iriondo 2009; Lepik, Abakumova, Zobel &
395 Semchenko 2012), much caution in their interpretations must be taken. When comparing
396 biomass of plants growing with kin or strangers, alternative hypotheses and several
397 confounding factors could explain the results and should be tested. A common difficulty to find
398 evidence of kin recognition in experiments is the lack of appropriate control treatments (Fig.2b).
399 In pot experiments, the use of a control treatment where plants are grown alone is mandatory
400 to determine if plants do compete or not (compared to plants grown alone). This control
401 treatment is also necessary to detect a potential genotype that has a better intrinsic growth.
402 Genotypes with better intrinsic growth capacities can lead to size inequality in groups of plants
403 (Simonsen, Chow & Stinchcombe 2014; Li, Xu & Liu 2018). Another confounding factor is
404 competitive ability, i.e. the relative growth of individuals grown in competition compared to
405 when grown alone (Masclaux et al., 2010; Zhang et al., 2019). Although it is usually argued
406 that a species can win either through being better at suppressing the performance of others
407 (effect) or being better at tolerating competition (response), Hart et al. (2018) showed that only
408 the competitive response is relevant to a species' competitive ability. Following early works on
409 kin recognition and discrimination in plants, several papers encouraged testing for competitive
410 ability (Masclaux et al., 2010; Chen, During & Anten 2012). However, this is still not a standard
411 procedure across studies and the lack of appropriate control treatments remains common in the
412 literature (Semchenko Saar & Lepik 2014; Crepy&Casal 2015; Murphy et al., 2017; Murphy,
413 Swanton, Van Acker & Dudley 2017; Li, Xu & Feng 2018; Pezzola, Pandolfi & Mancuso
414 2020). Here, we would like to emphasise that this lack of control treatment may result not only
415 in misinterpretation but also in the failure to reveal kin discrimination in studies (that have
416 therefore not been published). To test for competitive ability, it is necessary to set all the kin
417 groups in competition with each other and test whether a kin group always performs better than
418 the others prior to wrongly conclude to kin discrimination.

419 When plants grown with relatives were found to exhibit greater belowground biomass
420 than plants grown with strangers, the results were mostly interpreted using the niche partitioning
421 framework (File, Murphy & Dudley 2012; Subrahmaniam, Roby & Roux 2021). Niche

422 partitioning predicts that relatives overlap more in their niche use and compete more with each
423 other compared with unrelated conspecifics (Young 1981; Milla, Forero, Escudero & Iriondo
424 2009). However, stronger competition between relatives can be a by-product of forced
425 competition for a few dimensions of the niche that occurs when plants are grown at high density
426 in pots. The niche is a multidimensional concept (Hutchinson 1957; González, Dézerald,
427 Marquet, Romero & Srivastava 2017) with each component (e.g., light, nutrient, water, wind,
428 space, microbiota) occupying one dimension. Pots strongly constrain the growth of roots for
429 example in space or for the types of nutrients available. In pots, individuals share and compete
430 in a restricted number of dimensions and there may be no possible compensation in other
431 dimensions whereas in nature, individuals can compete in some dimensions of their niche but
432 not in others (Barker et al., 2017). Pot size can thus obscure the effects of positive interactions
433 between individuals (Hess & de Kroon 2007; Chen, Daring & Anten 2012) because root growth
434 responds to the available volume (McConnaughay & Bazzaz 1991), but also to nutrient amounts
435 (O'Brien, Gersani & Brown 2005) and concentrations (Maina, Brown & Gersani 2002).
436 Therefore, experiments conducted in restricted conditions (i.e., few niche dimensions) will bias
437 the conclusions towards niche partitioning versus kin discrimination and would not be relevant
438 in natural conditions where individuals growing with relatives can exploit all the dimensions of
439 their niche. On the other hand, when plants are supplemented with nutrients and resources are
440 not limiting, one can expect to find little or no difference between the stranger and kin
441 treatments because every individual has access to sufficient resources for its growth. Finally,
442 kin cooperation (kin selection) and competition (niche partitioning) can occur simultaneously,
443 possibly acting on different traits and thus different niche components (File, Murphy & Dudley
444 2012). Thus, depending on the observed traits and growth conditions it is possible to observe
445 one process but to miss the second. Therefore, although *in situ* experiments are challenging, we
446 share Anten & Chen's (2021a) appreciation that *in situ* experiments could solve some of the
447 problems associated with experiments under controlled conditions, in particular the constrained
448 niche aspects.

449

450 **Conclusion and final notes**

451 Over the last decade the scientific literature has continuously explored the role of relatedness
452 in the interactions between plants. However, we assert that some points have been generally
453 neglected and are worth considering in kin recognition/selection studies. The major aspects are
454 the underestimated role of intraspecific diversity for kin recognition and kin selection traits and

455 the fact that local conditions can influence the outcome of interactions. The concept of
456 "conditionality" and the "stress gradient hypothesis" suggest that intra-specific interactions, and
457 in particular interactions between relatives, are dependent on environmental conditions,
458 especially under various stress types. The case of long-lived species, where individuals face
459 different environmental conditions and stresses throughout their life, is especially promising.
460 In addition, studying the spatial genetic structure of populations for *in situ* studies would be of
461 great help in pointing out settings in which different types of interactions might take place. We
462 also discuss the challenges of distinguishing kin discrimination from kin selection, which are
463 to accurately measure the costs and benefits of the interaction as well as the relatedness of the
464 interacting individuals, and the importance of appropriate control treatments to test for all
465 alternative hypotheses (Fig. 2). We also strongly recommend several key-points to be
466 considered in future studies (Fig. 2b): to capture a large intraspecific diversity of the studied
467 species, to include different environmental conditions and to include a relatedness gradient
468 between individuals. This would help to take care of various confounding effects that could
469 interfere with the interpretation of the results.

470

471 **References**

472

473 Anten N.P.R. & Chen B.J.W. (2021a) Detect thy family: Mechanisms, ecology and agricultural
474 aspects of kin recognition in plants. *Plant, Cell & Environment* **44**, 1059–1071. doi:
475 10.1111/pce.14011

476 Anten N.P.R. & Chen B.J.W. (2021b) Kin discrimination in allelopathy and consequences for
477 agricultural weed control. *Plant, Cell & Environment*, **44**, 3705– 3708.
478 <https://doi.org/10.1111/pce.14173>

479 Axelrod R. & Hamilton W.D. (1981) The Evolution of Cooperation. The evolution of
480 cooperation. *Science* (New York, N.Y.) **211**, 1390–1396. doi: 10.1126/science.7466396

481 Barker J.L., Bronstein J.L., Friesen M.L., Jones E.I., Reeve H.K., Zink A.G. & Frederickson
482 M.E. (2017) Synthesizing perspectives on the evolution of cooperation within and between
483 species. *Evolution* **71**, 814–825.

484 Bertness M.D. & Callaway R. (1994) Positive interactions in communities. *Trends in Ecology*
485 *& Evolution* **9**, 191–193.

486 Bilas R.D., Bretman A. & Bennett T. (2021) Friends, neighbours and enemies: an overview of
487 the communal and social biology of plants. *Plant, Cell & Environment* **44**, 997–1013.

488 Bolnick D.I., Amarasekare P., Araújo M.S., Bürger R., Levine J.M., Novak M., ... Vasseur
489 D.A. (2011) Why intraspecific trait variation matters in community ecology. *Trends in Ecology*
490 *& Evolution* **26**, 183–192.

491 Bronstein J.L. (1994) Conditional outcomes in mutualistic interactions. *Tree* **9**, 214–217

492 Bronstein J.L. (2009) The evolution of facilitation and mutualism. *Journal of Ecology* **97**, 1160–
493 1170.

494 Callaway R.M., Pennings S.C. & Richards C.L. (2003) Phenotypic Plasticity and Interactions
495 among Plants. *Ecology* **84**, 1115–1128.

496 Chen, B.J.W., During, H.J. & Anten, N.P.R. (2012) Detect thy neighbor: Identity recognition
497 at the root level in plants. *Plant Science*, **195**, 157–167.
498 <https://doi.org/10.1016/j.plantsci.2012.07.006>

499 Clark J.S. (2010) Individuals and the Variation Needed for High Species Diversity in Forest
500 Trees. *Science* **327**, 1129–1132.

501 Corenblit D., Garófano-Gómez V., González E., Hortobágyi B., Julien F., Lambs L., ... Till-
502 Bottraud I. (2018) Niche construction within riparian corridors. Part II: The unexplored role of
503 positive intraspecific interactions in Salicaceae species. *Geomorphology* **305**, 112–122

504 Corenblit D., Steiger J., González E., Gurnell A.M., Charrier G., Darrozes J., ... Voldoire O.
505 (2014) The biogeomorphological life cycle of poplars during the fluvial biogeomorphological
506 succession: a special focus on *Populus nigra*. *Earth Surface Processes and Landforms* **39**, 546–
507 563.

508 Crepy, M.A. & Casal, J.J. (2014) Photoreceptor - mediated kin recognition in plants. *New*
509 *Phytologist*, **205**(1), 329–338. <https://doi.org/10.1111/nph.13040>

510 Dudley, S.A. & File, A.L. (2007a) Kin recognition in an annual plant. *Biology*, **3**(4), 435–438.
511 <https://doi.org/10.1098/rsbl.2007.0232>

512 Dudley, S.A. & File, A.L. (2007b) Yes, kin recognition in plants! *Biology Letters*, **4**(1), 69–70.
513 <https://doi.org/10.1098/rsbl.2007.0585>

514
515 Dudley S.A., Murphy G.P. & File A.L. (2013) Kin recognition and competition in plants.
516 *Functional Ecology* **27**, 898–906.

517 Ehlers B.K. & Bilde T. (2019) Inclusive fitness, asymmetric competition and kin selection in
518 plants. *Oikos* **128**, 765–774.

519 Ehlers B.K., David P., Damgaard C.F. & Lenormand T. (2016) Competitor relatedness, indirect
520 soil effects and plant coexistence. *Journal of Ecology*, **104**, 1126–1135.
521 <https://doi.org/10.1111/1365-2745.12568>

522 Fajardo A. & McIntire E.J.B. (2011) Under strong niche overlap conspecifics do not compete
523 but help each other to survive: facilitation at the intraspecific level: Intraspecific facilitation in
524 *Nothofagus pumilio*. *Journal of Ecology* **99**, 642–650. [https://doi.org/10.1111/j.1365-
525 2745.2010.01771.x](https://doi.org/10.1111/j.1365-2745.2010.01771.x)

526 Fajardo A., Torres-Díaz C. & Till-Bottraud I. (2016) Disturbance and density-dependent
527 processes (competition and facilitation) influence the fine-scale genetic structure of a tree
528 species' population. *Annals of Botany* **117**, 67–77.

529 File A.L., Murphy G.P. & Dudley S.A. (2012) Fitness consequences of plants growing with
530 siblings: reconciling kin selection, niche partitioning and competitive ability. *Proceedings of*
531 *the Royal Society B: Biological Sciences* **279**, 209–218.

532 Fox R.J., Donelson J.M., Schunter C., Ravasi T. & Gaitán-Espitia J.D. (2019) Beyond buying
533 time: the role of plasticity in phenotypic adaptation to rapid environmental change.
534 *Philosophical Transactions of the Royal Society B: Biological Sciences* **374**, 20180174.

535 Fréville H., Roumet P., Rode N.O., Rocher A., Latreille M., Muller M. & David J. (2019)
536 Preferential helping to relatives: A potential mechanism responsible for lower yield of crop
537 variety mixtures? *Evolutionary Applications* **12**, 1837–1849.

538 Gadagkar R., Gordon D., Keller L., Michod R., Queller D., Robinson G.E., ... West-Eberhard
539 M.J. (2019) Insights and opportunities in insect social behavior. *Current Opinion in Insect*
540 *Science* **34**, ix–xx. doi: 10.1016/j.cois.2019.08.009.

541 Gardner, A. & West, S.A. (2007) Social Evolution: the Decline and Fall of Genetic Kin
542 Recognition. *Current Biology*, **17**(18), R810–R812. <https://doi.org/10.1016/j.cub.2007.07.030>

543
544 Ghalambor C.K., McKay J.K., Carroll S.P., Reznick D.N. (2007) Adaptive versus non-adaptive
545 phenotypic plasticity and the potential for contemporary adaptation in new environments.
546 *Functional Ecology* **21**, 394–407.

547 Goddard E.L., Varga S., John E.A. & Soulsbury C.D. (2020) Evidence for Root Kin
548 Recognition in the Clonal Plant Species *Glechoma hederacea*. *Frontiers in Ecology and*
549 *Evolution* **8**, 578141.

550 González A.L., Dézerald O., Marquet P.A., Romero G.Q. & Srivastava D.S. (2017) The
551 Multidimensional Stoichiometric Niche. *Frontiers in Ecology and Evolution* **5**, 110.

552 Grubb, P.J. (1977) The maintenance of species richness in plant communities: The importance
553 of the regeneration niche. *Biological Reviews* **52**, 107–45.

554 Hamilton W.D. (1964) The genetical evolution of social behaviour. I. *Journal of Theoretical*
555 *Biology* **7**, 1–16.

556 Hardy O.J. (2003) Estimation of pairwise relatedness between individuals and characterization
557 of isolation-by-distance processes using dominant genetic markers. *Molecular Ecology* **12**,
558 1577–1588.

559 Hart S.P., Freckleton R.P. & Levine J.M. (2018) How to quantify competitive ability. Journal
560 of Ecology **106**, 1902–1909.

561 He Y., Xu H., Liu H., Luo M., Chu C. & Fang S. (2021) Sexual competition and kin recognition
562 co-shape the traits of neighboring dioecious *Diospyros morrisiana* seedlings. Horticulture
563 Research **8**, 162.

564 Hess L. & De Kroon H. (2007) Effects of rooting volume and nutrient availability as an
565 alternative explanation for root self/non-self-discrimination. Journal of Ecology **95**, 241–251.

566 Hortobágyi B., Corenblit D., Steiger J. & Peiry J.-L. (2018) Niche construction within riparian
567 corridors. Part I: Exploring biogeomorphic feedback windows of three pioneer riparian species
568 (Allier River, France). Geomorphology **305**, 94–111.

569 Hussain A., Rodriguez-Ramos J.C. & Erbilgin N. (2019) Spatial characteristics of volatile
570 communication in lodgepole pine trees: Evidence of kin recognition and intra-species support.
571 Science of the Total Environment **692**, 127–135. doi:
572 <https://doi.org/10.1016/j.scitotenv.2019.07.211>

573 Hutchinson G.E. (1957) Concluding remarks. Cold Spring Harbor Symposia. Quantitative
574 Biology **22**, 415–427.

575 Jones C.G., Lawton J.H. & Shachak M. (1994) Organisms as Ecosystem Engineers. Oikos **69**,
576 373.

577 Karban R., Shiojiri K., Ishizaki S., Wetzel W.C. & Evans R.Y. (2014) Kin recognition affects
578 plant communication and defence. Proceedings of the Royal Society B: Biological Sciences
579 **280**, 20123062–20123062.

580 Kay T., Lehmann L. & Keller L. (2019) Kin selection and altruism. Current Biology **29**, R438–
581 R442.

582 Lepik A., Abakumova M., Zobel K. & Semchenko M. (2012) Kin recognition is density-
583 dependent and uncommon among temperate grassland plants. Functional Ecology **26**, 1214–
584 1220.

585 Li J., Xu X. & Feng R. (2018a) Soil fertility and heavy metal pollution (Pb and Cd) alter kin
586 interaction of *Sorghum vulgare*. Environmental and Experimental Botany **155**, 368–377.

587 Li J., Xu X.-L. & Liu Y.-R. (2018) Kin recognition in plants with distinct lifestyles:
588 implications of biomass and nutrient niches. *Plant Growth Regulation* **84**, 333–339.

589 Loiselle B.A., Sork V.L., Nason J. & Graham C. (1995) Spatial genetic structure of a tropical
590 understory shrub, *Psychotria officinalis* (Rubiaceae). *American Journal of Botany* **82**, 1420–
591 1425.

592 Maina G.G., Brown J.S. & Gersani M. (2002) Intra-plant versus inter-plant root competition in
593 beans: avoidance, resource matching or tragedy of the commons. *Plant Ecology* **160**, 235–247.

594 Masclaux F., Hammond R.L., Meunier J., Gouhier-Darimont C., Keller L. & Reymond P.
595 (2010) Competitive ability not kinship affects growth of *Arabidopsis thaliana* accessions. *New*
596 *Phytologist* **185**, 322–331.

597 Maynard-Smith J. (1982) *Evolution and the theory of games*. Cambridge University Press,
598 Cambridge.

599 Mazal L., Corenblit D., Barsoum N., Steiger J., Skot L., Fumanal B. & Till-Bottraud I. (2021)
600 Fine-scale spatial genetic structure and intra-specific interactions of *Populus nigra* within a
601 natural river corridor along the lower Allier River (France). *Flora* **275**, 151763.

602 Mazal L., Corenblit D., Fumanal B. & Till-Bottraud I. (2022) Black poplar establishment on
603 alluvial bars: seed rain homogeneity over a few kilometres. *Dendrobiology* **87**, 69–78.

604 McConnaughay K.D.M., & Bazzaz F.A. (1991) Is physical space a soil resource? *Ecology* **72**,
605 94–103.

606 McIntire E.J.B. & Fajardo A. (2011) Facilitation within Species: A Possible Origin of Group-
607 Selected Superorganisms. *The American Naturalist* **178**, 88–97.

608 McIntire E.J.B. & Fajardo A. (2014) Facilitation as a ubiquitous driver of biodiversity. *New*
609 *Phytologist* **201**, 403–416.

610 Milla R., Forero D.M., Escudero A. & Iriando J.M. (2009) Growing with siblings: a common
611 ground for cooperation or for fiercer competition among plants? *Proceedings of the Royal*
612 *Society B: Biological Sciences* **276**, 2531–2540.

613 Murphy G.P. & Dudley, S. A. (2009). Kin Recognition: Competition and Cooperation in
614 *Impatiens* (Balsaminaceae). *American Journal of Botany*, **96(11)**, 1990–1996.
615 <http://www.jstor.org/stable/20621978>

616 Murphy G.P., Swanton C.J., Van Acker R.C. & Dudley S.A. (2017) Kin recognition, multilevel
617 selection and altruism in crop sustainability. *Journal of Ecology* **105**, 930–934

618 O’Brien E.E., Gersani M. & Brown J.S. (2005) Root proliferation and seed yield in response to
619 spatial heterogeneity of below-ground competition. *New Phytologist* **168**, 401–412.

620 Penn D.J. & Frommen J.G. (2010) Kin recognition: an overview of conceptual issues,
621 mechanisms and evolutionary theory. In *Animal Behaviour: Evolution and Mechanisms*. (ed P.
622 Kappeler), pp. 55–85. Springer Berlin Heidelberg, Berlin, Heidelberg.

623 Pezzola E., Pandolfi C. & Mancuso S. (2020) Resource availability affects kin selection in two
624 cultivars of *Pisum sativum*. *Plant Growth Regulation* **90**, 321–329.

625 Pickles B.J., Wilhelm R., Asay A.K., Hahn A.S., Simard S.W. & Mohn W.W. (2017) Transfer
626 of 13 C between paired Douglas-fir seedlings reveals plant kinship effects and uptake of
627 exudates by ectomycorrhizas. *New Phytologist* **214**, 400–411.

628 Proença B., Nez T., Poli A., Ciutat A., Devaux L., Sottolichio A., ... Michalet R. (2019)
629 Intraspecific facilitation explains the spread of the invasive engineer *Spartina anglica* in
630 Atlantic salt marshes. *Journal of Vegetation Science* **30**, 212–223.

631 Reijers V.C., van den Akker M., Cruijssen P.M.J.M., Lamers L.P.M. & van der Heide T. (2019)
632 Intraspecific facilitation explains the persistence of *Phragmites australis* in modified coastal
633 wetlands. *Ecosphere* **10**, e02842.

634 Rousset F. & Roze D. (2007) Constraints on the origin and maintenance of genetic kin
635 recognition. *Evolution* **61**, 2320–2330.

636 Segovia N.I., Vásquez J.A., Faugeron S. & Haye P.A. (2015) On the advantage of sharing a
637 holdfast: effects of density and occurrence of kin aggregation in the kelp *Lessonia berteroana*.
638 *Marine Ecology* **36**, 1107–1117.

639 Semchenko, M., Saar, S. & Lepik, A. (2014) Plant root exudates mediate neighbour recognition
640 and trigger complex behavioural changes. *New Phytologist*, **204**(3), 631–637.
641 <https://doi.org/10.1111/nph.12930>

642
643 Semchenko M., Saar S. & Lepik A. (2017) Intraspecific genetic diversity modulates plant-soil
644 feedback and nutrient cycling. *New Phytologist* **216**, 90–98.

645 Siefert A., Violle C., Chalmandrier L., Albert C.H., Taudiere A., Fajardo A., ... Wardle D.A.
646 (2015) A global meta-analysis of the relative extent of intraspecific trait variation in plant
647 communities. *Ecology Letters* **18**, 1406–1419.

648 Simonsen A.K., Chow T. & Stinchcombe J.R. (2014) Reduced plant competition among kin
649 can be explained by Jensen’s inequality. *Ecology and Evolution* **4**, 4454–4466.

650 Stoll P. & Weiner J. (2000) A Neighborhood View of Interactions among Individual Plants. In
651 *The Geometry of Ecological Interactions*, 1st ed. (eds U. Dieckmann, R. Law & J.A.J. Metz),
652 pp. 11–27. Cambridge University Press.

653 Subrahmaniam H.J., Libourel C., Journet E.-P., Morel J.-B., Muños S., Niebel A., ... Roux F.
654 (2018) The genetics underlying natural variation of plant-plant interactions, a beloved but
655 forgotten member of the family of biotic interactions. *The Plant Journal* **93**, 747–770.

656 Subrahmaniam H.J., Roby D. & Roux F. (2021) Toward Unifying Evolutionary Ecology and
657 Genomics to Understand Positive Plant–Plant Interactions Within Wild Species. *Frontiers in*
658 *Plant Science* **12**, 683373.

659 Takigahira H. & Yamawo A. (2019) Competitive responses based on kin-discrimination
660 underlie variations in leaf functional traits in Japanese beech (*Fagus crenata*) seedlings.
661 *Evolutionary Ecology* **33**, 521–531.

662 Tarroux E. & DesRochers A. (2011) Effect of natural root grafting on growth response of jack
663 pine (*Pinus banksiana*; Pinaceae). *American Journal of Botany* **98**, 967–974.

664 Tarroux E., DesRochers A. & Tremblay F. (2014) Molecular analysis of natural root grafting
665 in jack pine (*Pinus banksiana*) trees: how does genetic proximity influence anastomosis
666 occurrence? *Tree Genetics & Genomes* **10**, 667–677.

667 Till-Bottraud I., Fajardo A. & Rioux D. (2012) Multi-stemmed trees of *Nothofagus pumilio*
668 second-growth forest in Patagonia are formed by highly related individuals. *Annals of Botany*
669 **110**, 905–913.

670 Till-Bottraud I. & de Villemereuil P. (2016) Kin recognition or phenotype matching? *New*
671 *Phytologist* **209**, 13–14.

672 Torices R., DeSoto L., Narbona E., Gómez J.M. & Pannell J.R. (2021) Effects of the
673 Relatedness of Neighbours on Floral Colour. *Frontiers in Ecology and Evolution* **9**, 589781.
674 doi: 10.3389/fevo.2021.589781

675 Torices R., Gómez J.M. & Pannell J.R. (2018) Kin discrimination allows plants to modify
676 investment towards pollinator attraction. *Nature Communications* **9**.
677 <https://doi.org/10.1038/s41467-018-04378-3>

678 de Villemereuil P., Gaggiotti O.E., M Mouterde M. & Till-Bottraud I. (2016) Common garden
679 experiments in the genomic era: new perspectives and opportunities. *Heredity* **116**, 249–254.
680 [doi:10.1038/hdy.2015.93](https://doi.org/10.1038/hdy.2015.93).

681 de Villemereuil, P., Mouterde, M., Gaggiotti, O.E. & Till - Bottraud, I. (2018) Patterns of
682 phenotypic plasticity and local adaptation in the wide elevation range of the alpine plant *Arabis*
683 *alpina*. *Journal of Ecology*, **106**, 1952-1971.

684 Waldman B. (1988) The ecology of kin recognition. *Annual Review of Ecology and*
685 *Systematics* **19**, 543–571.

686 West, S.A., Griffin, A.S. & Gardner, A. (2007) Social semantics: altruism, cooperation,
687 mutualism, strong reciprocity and group selection. *Journal of Evolutionary Biology*, **20**(2),
688 415-432. <https://doi.org/10.1111/j.1420-9101.2006.01258.x>

689 Westerband A.C., Funk J.L. & Barton K.E. (2021) Intraspecific trait variation in plants: a
690 renewed focus on its role in ecological processes. *Annals of Botany* **127**, 397–410.

691 Xu, Y., Cheng, H.F., Kong, C.H., & Meiners, S.J. (2021) Intra-specific kin recognition
692 contributes to inter-specific allelopathy: A case study of allelopathic rice interference with
693 paddy weeds. *Plant, Cell & Environment* **44**, 3709– 3721. <https://doi.org/10.1111/pce.14083>

694 Yamawo A., Sato M. & Mukai H. (2017) Experimental evidence for benefit of self-
695 discrimination in roots of a clonal plant. *AoB PLANTS* **9**.

696 Yang X.-F., Li L.-L., Xu Y. & Kong C.-H. (2018) Kin recognition in rice (*Oryza sativa*) lines.
697 *New Phytologist* **220**, 567–578.

698 Young J.P.W. (1981) Sib competition can favour sex in two ways. **88**, 755–756.

699 Yoda, K., Kira, T., Ogawa, H. & Hozumi, K. (1963) Self - thinning in overcrowded pure stands
700 under cultivated and natural conditions. *Journal of Biology*, **14**, 107-129.

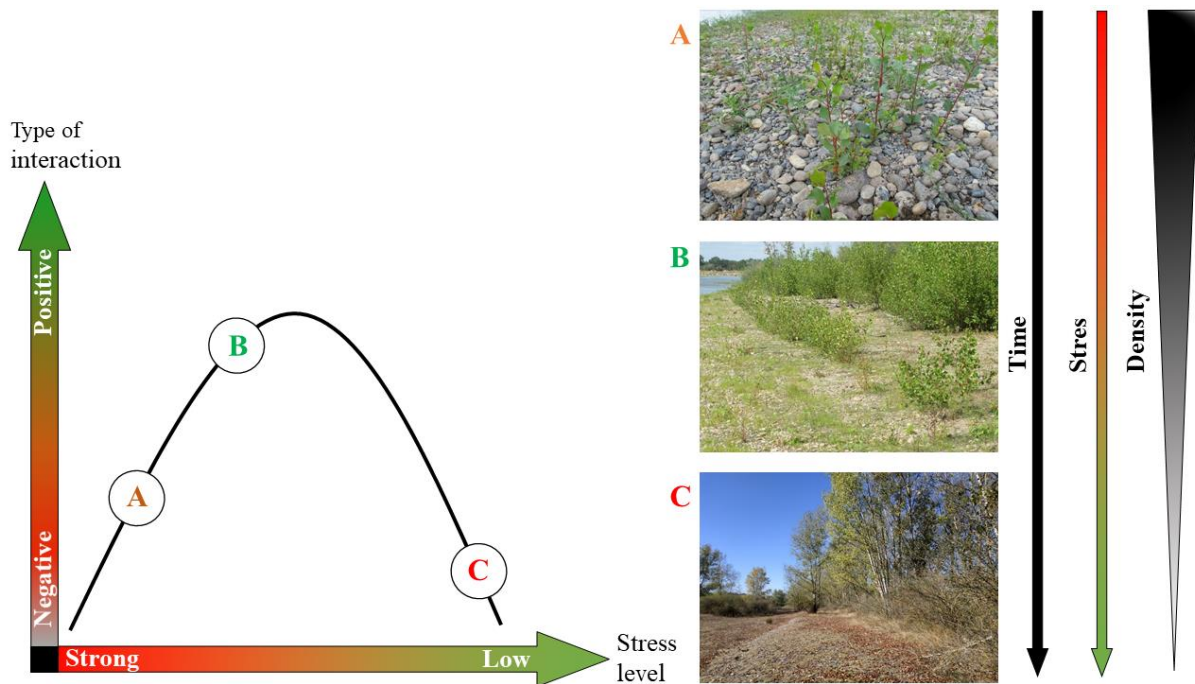
701 Younginger B.S., Sirová D., Cruzan M.B. & Ballhorn D.J. (2017) Is biomass a reliable estimate
702 of plant fitness? *Applications in Plant Sciences* **5**, 1600094.

703 Zhang L., Liu Q., Tian, Y., Xu Xingliang. & Ouyang Hua. (2016) Kin selection or resource
704 partitioning for growing with siblings: implications from measurements of nitrogen uptake.
705 Plant Soil **398**, 79–86. <https://doi.org/10.1007/s11104-015-2641-z>

706 Zhang Z., Zhou F., Pan X., Kleunen M., Liu M. & Li B. (2019) Evolution of increased
707 intraspecific competitive ability following introduction: The importance of relatedness among
708 genotypes. Journal of Ecology **107**, 387–395.

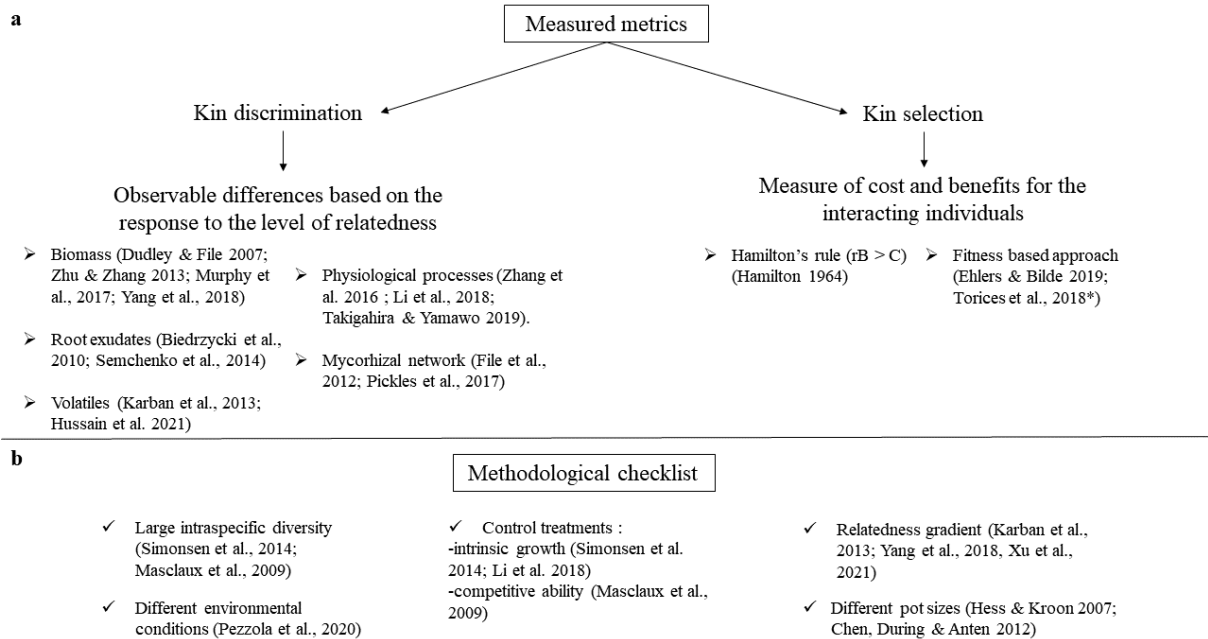
709

710 Figures



711

712 **Figure 1.** The stress gradient hypothesis (SGH, Bertness & Callaway, 1994) applied to the case
713 of *Populus nigra*. A: Seedlings establish in densities on bare surfaces along rivers and
714 experience harsh conditions (mechanical and hydric stresses). Facilitation occurs due to the
715 high densities of individuals and reduces the stresses for each individual. At this stage no spatial
716 genetic structure (SGS) is found and the main direct interactions between individuals are neutral
717 (commensalism). B: Patches of young individuals that survived now exhibit SGS. The effect of
718 *P. nigra* on the environment enhance the local conditions (soil moisture and nutrients).
719 Facilitation still occurs because of the protective effect of individuals located at the upstream
720 part of the stand (Mazal et al., 2021). Main direct interactions shift to positive (potentially
721 cooperation) and could be directed towards related individuals (as SGS is significant at this
722 stage). C: At older stages, individuals no longer benefit from the presence of their neighbours
723 and the local conditions are no longer stressful. Stem density is highly reduced through self -
724 thinning processes (i.e., competition) (Yoda et al., 1963). The main interactions between
725 individual are either neutral (commensalism) or negative (competition).



726
727

728 Figure 2. Suggested guidelines for the design of experiments on kinship studies in plants. (a)
729 Measured metrics commonly used in the conceptual frameworks studied: kin recognition
730 discrimination or kin selection. (b) Checklist for the design of future studies outlining key
731 points to be controlled. For both (a) and (b), we have mentioned the studies that in our opinion
732 have taken some of the points above into account and that could be used as a reference. *As
733 discussed in the text, although this study did not measure the fitness of individuals, it is the
734 closest evidence to date for kin selection in plants.
735