

~~Flexibility of interactions depends on habitat suitability~~

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16 Abstract

17 Evidence that the environment influences the interaction between species
18 is rapidly accumulating. However, how it happens is currently unclear as
19 environmental gradients appear to have contrasting or non-linear effects on
20 the species' trophic niche breadth depending on the environmental variable.
21 Here, we explore the relationship between the stresses imposed by the envi-
22 ronment, instead of environmental gradients directly, and niche breadth using
23 a global dataset of plant-pollinator interactions. We found that environmental
24 stress plays a significant role in determining the number of partners a species
25 interacts with, but this role is highly variable across species. In particular,
26 when faced with environmental stress, species that have a large number of
27 interactions are more likely to focus on a smaller number of, presumably
28 higher-quality, interactions. Contrastingly, the specialists that can cope
29 with increased stress are more likely to broaden their niche and engage in
30 opportunistic interactions, effectively behaving as facultative generalists.

→ define this

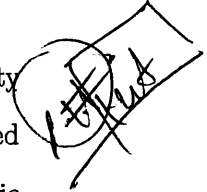
31 Introduction

32 Species interactions are known to vary widely across space and time. There
33 are multiple examples of species that interact tightly in an ecological com-
34 munity or a particular season but not in another. It is crucial to understand
35 why and how this turnover of interactions occur because the network of
36 interactions of an ecological community is known to underpin ecosystem
37 function and stability. Despite much progress in the last decades, we are
38 still unable to predict species interactions, partially because we still have not
39 accurately identified the factors that drive species to interact under certain
40 conditions and not under another.

41 There are two main processes involved in whether ~~two~~ species in a community
42 interact or not. The first is an "Eltonian" niche process that is underpinned
43 by the traits of the interacting species. Compared to other kinds of trophic
44 interactions, mutualistic interactions, and plant-pollination interactions,
45 in particular, are relatively generalised. While a large proportion of the
46 interactions appear to be opportunistic, a fraction of interactions shows a
47 strong signature of deep co-evolutionary history (Hutchinson, Cagua, and
48 Stouffer 2017). In contrast to opportunistic interactions with low trait-
49 matching, the more co-evolved interactions with optimally matched traits are
50 more persistent and presumably lead to a more substantial per-capita benefit
51 for the interacting species. The second is a neutral process underpinned by
52 the encounter rate of the interacting species. All else being equal, when the
53 encounter rate of interacting species is high, there is a higher probability to
54 interact than if the rate were lower, for example, due to species rarity.

55 Perhaps, because it operates at scales larger than those of the Eltonian
56 niche concept, the abiotic environment is one of the factors that is commonly
57 overlooked by community ecologists when explaining interaction turnover (but
58 see Gravel et al. 2018). The most evident—and best-studied—way through

→ define what is



in time / space?

need to clarify
this, because there
are tons of
studies looking
at the eff. of
e.g. habitat types
on interactions

identity

59 which the environment can affect the structure of ecological networks is by

not clear

60 way of patterns of community composition. At the heart of biogeography theory?

61 is the idea that the abiotic environment can determine the "Grinnellian"

62 niche of a species, this is whether the species can have a positive growth-

63 rate when rare in a particular location. For two species to interact, they

64 first need to co-occur in the same community, which can only happen if

65 the environmental conditions of the ^{location} community fall within the Grinnellian

66 niche of both species. Although less understood, the environment can also

67 affect the interactions themselves, after community composition has been

68 established (Tylianakis and Morris 2017). For instance, some studies show

69 that trophic interactions, like predation (McKinnon et al. 2010; Vucic-Pestic

70 et al. 2011) and herbivory (Baskett and Schemske 2018), can increase with

71 temperature but might decrease with precipitation (Pires et al. 2016), and

72 both plant-plant (Bertness and Callaway 1994) and plant-pollinator (Tur

73 et al. 2016) interactions can switch from facilitative to competitive along

74 an elevation/temperature gradient. Contrastingly, other studies have shown

75 either no effect or non-linear effects of temperature or precipitation on

76 plant-pollinator interactions (Devoto, Medan, and Montaldo 2005; Gravel

77 et al. 2018). One way to summarise the seemingly contradictory evidence

78 of whether the environment can influence whether two species interact or

79 not is by moving from gradients of temperature or precipitation to gradients

80 of environmental stress. Previous research is so far equivocal and, taken

81 together, suggest that environmental stress can have two contrasting effects

82 on species' Eltonian-niche (Tylianakis and Morris 2017). On the one hand,

83 it is possible that when species are under environmental stress, they might

84 be "pressured" to focus on partners with which they are best adapted to

85 interact. For instance, Hoiss et al. (2012) found increased phylogenetic

86 clustering between plants and pollinators at higher altitudes; while Peralta et

87 al. (2015) found that parasitoids in plantation forest, where environmental

88 stress was higher than in native forests, were constrained to interact with

what do you mean by this?

what freq?

refs cover

here define it as it is the first time you mention this term

what do you mean by this?

What about focusing only on plant-poll int throughout the intro? I say this because parasitoids are generally more specialized than other groups, so it could be argued that even specialized groups could increase their specialization (contrary to what your hypothesis is).

89 hosts they were best adapted to attack. Similarly, Lavandero and Tylianakis
90 (2013) found that environmental stress due to higher temperature reduced
91 the breadth of the Eltonian niche of parasitoids. On the other, it is also
92 possible that when species are under environmental stress, they are forced to
93 be more flexible in their interactions as higher environmental stress is likely
94 to be reflected in greater energetic or reproductive costs and therefore they
95 might not be able to sustain encounter rates with their preferred partners
96 at sufficient levels. In line with this hypothesis, Hoiss, Krauss, and Steffan-
97 Dewenter (2015) found that the specialisation of plant-pollinator networks
98 decreased both with elevation and after extreme drought events. Likewise,
99 Pellissier et al. (2010) found a positive relationship between niche breadth
100 and environmental stress: disk- or bowl-shaped blossoms (which allow a large
101 number of potential pollinator species to access pollen and nectar rewards)
102 dominated at high altitude flower communities.

103 We analyse data on plant-pollinator interactions globally and calculate the
104 environmental suitability of all species present at two or more communities
105 to. Our main aim is to leverage that information to test the two competing
106 hypotheses that relate environmental stress and Eltonian niche breadth
107 and whether there is a pattern within and across species. Specialist species
108 can become "facultative" generalists to reduce their vulnerability to the
109 absence of preferred partners (for example, when variations in climate de-
110 couple phenologies; Benadi et al. 2014). We therefore also expect that
111 as environmental stress increases species with a relatively small number of
112 partners are more likely to broaden their Eltonian niche. Species with a
113 large number of partners, on the other hand, should have a larger pool of
114 available partners and might, therefore, be more likely to narrow their niche
115 under environmental stress. Importantly, when testing these hypotheses, we
116 control for the potential effects of the environment in community composi-
117 tion and the size of the species fundamental niche, both from an Eltonian

and this
would be
env. stress?

for what?

any pattern?

Separar
OK!
revis

not clear
Become more
generalists
you mean?

Ass repetição
la idea de
esta redución
de nichos sería
pl conectarlos en
más beneficios.

Does this mean that you
control for the fact that the
hosts (or interacting partners) are present or not?
And their abundances??

(interactions) and Grinnellian (environment) perspective.

Methods

We retrieved plant-pollinator networks from the Web of Life database (Fortuna, Ortega, and Bascompte 2014). This database contains datasets originating from 57 studies published in the primary literature between 106 and 2016 (Kaiser-Bunbury et al. 2010; Bartomeus, Vilà, and Santamaría 2008; Arroyo, Primack, and Armesto 1982; Hattersley-Smith 1985; Barrett and Helenurm 1987; Dicks, Corbet, and Pywell 2002; Dupont, Hansen, and Olesen 2003; Elberling and Olesen 1999; Herrera 1988; Hocking 1968; Inouye and Pyke 1988; Kato et al. 1990; Medan et al. 2002; Memmott 1999; Motten 1986; Olesen, Eskildsen, and Venkatasamy 2002; Ollerton 2003; Petanidou and Vokou 1993; McMullen 106AD; Mosquin 1967; Clements and Long 1923; Kevan 1970; Small 1976; Montero 2005; Stald, Valido, and Olesen 2003; Ingversen 2006; Bundgaard 2003; Bek 2006; KATO 2000; Dupont and Olesen 2009; Lundgren and Olesen 2005; Percival 1974; Philipp et al. 2006; Primack 1983; Ramirez 1989; Ramirez and Brito 1992; Schemske et al. 1978; Smith-Ramírez et al. 2005; Vázquez 2002; Robertson 1929; Ferreira Canela 2006; Kohler 2011; Lasprilla and others 2003; Sabatino 2010; Lara 2006; Las-Casas, Azevedo Júnior, and Dias Filho 2012; del Coro Arizmendi and Ornelas 1990; Abreu and Vieira 2004; Vizentin-Bugoni et al. 2016; Kaiser-Bunbury et al. 2014; Bezerra, Machado, and Mello 2009; INouE et al. 1990; KATO, Matsumoto, and Kato 1993; Yamazaki and Kato 2003; Gutierrez Z, Rojas-Nossa, and Stiles 2004; Kato and Miura 1993; Kakutani et al. 1990). Calculating the environmental stress of species in their community and their Eltonian niche breadth requires that the scientific names used to identify them are valid and unambiguous. This is because valid scientific names are necessary to both accurately match species across studies and to obtain other information from biological databases. However, a significant proportion