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1 *Reviews for Trends in Ecology & Evolution*

2

3 **Unifying niche shift studies: insights from biological invasions**

4

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24 *A Glossary Box*

25

26

27

28 **Abstract**

29 Assessing whether the climatic niche of a species may change between different  
30 geographic areas or time periods has become increasingly important in the context of  
31 ongoing global change. However, approaches and findings have remained largely  
32 controversial so far, calling for a unification of methods. Here, we build on a review of  
33 empirical studies of invasion to formalize a unifying framework that decomposes niche  
34 change into unfilling, stability and expansion situations, taking both a pooled-range and  
35 range-specific perspective on the niche, while accounting for climatic availability and  
36 climatic analogy. This framework provides new insights into the nature of climate niche  
37 shifts and our ability to anticipate invasions, and may help guiding the design of  
38 experiments for assessing causes of niche changes.

39

40

41 **Niche shifts during invasions: setting the scene**

42 How climate determines the distribution of species is a classic question in ecology,  
43 closely tied to Hutchinson's (1957) concept of the environmental niche, and still a major  
44 research topic (Soberón & Nakamura, 2009; Peterson *et al.*, 2011). Although, for some  
45 species, it has become possible to determine the fundamental climatic niche based on  
46 physiological information and a mechanistic understanding (e.g. (Kearney & Porter,  
47 2009)), for most species only the realized climate niche can realistically be estimated  
48 through empirical studies (Guisan & Thuiller, 2005). With global change, it has become  
49 increasingly important not only to describe species' climate niches but also to  
50 understand whether these can change rapidly (niche shifts) or not (niche conservatism)  
51 between different geographic areas or time periods (Peterson, 2011) (Fig. 1). The  
52 primary approach to investigating climatic niches in space and time has been to analyse  
53 climatic conditions across a species' distributional ranges and/or over time (Pearman *et al.*, 2008a).

55 As already understood by Charles Elton (1958), biological invasions offer a unique  
56 opportunity to study how species colonize new environments (Sax *et al.*, 2007;  
57 Richardson & Pysek, 2008; Kueffer *et al.*, 2013), and whether they retain their climatic

58 niche in a new range (Pearman *et al.*, 2008a). Addressing this question has proved  
59 important in recent years as a test of our capacity to use climate matching to assess  
60 invasion risks by exotic species at transnational scales (Venette *et al.*, 2010; Guisan *et al.*,  
61 2013), in particular when using ecological niche models (ENMs), which rely heavily on  
62 climatic niche conservation between native and exotic ranges (Pearman *et al.*, 2008a;  
63 Colwell & Rangel, 2009; Peterson, 2011). Do a majority of species retain their native  
64 climatic niche when introduced elsewhere? The answer to this question is fundamental  
65 because it informs both theoretical and applied ecology, but approaches have diverged  
66 and findings have remained largely controversial so far (Table S1, supplementary  
67 material) (Pearman *et al.*, 2008a; Peterson, 2011).

68 Evidence exists both for and against climatic niche conservatism during invasions. A  
69 recent large-scale survey of 50 Holarctic terrestrial plant invaders concluded that  
70 climatic niche shifts are rare overall between the native and invaded ranges, and  
71 therefore models can usefully predict invasion in the exotic range (Petitpierre *et al.*,  
72 2012.) The same conclusions were reached for birds (Strubbe *et al.*, 2013) and other  
73 groups (see (Peterson, 2011); Table S1, suppl. mat.). But the assumption of niche  
74 conservatism was also challenged by evidence of climatic niches shifting during  
75 invasions (e.g. (Broennimann *et al.*, 2007; Fitzpatrick *et al.*, 2007; Rödder & Lötters,  
76 2009; Medley, 2010; Lauzeral *et al.*, 2011); Table S1), potentially hampering predictions  
77 in the new range. Contrasting evidence of niche dynamics during invasions, and  
78 particularly of the frequency of niche shifts (i.e. of centroid and/or limits; see Fig. 1)  
79 among various taxonomic groups, thus coexist in the literature (about 50% shifts/42%  
80 no-shifts and 8% no-conclusion in Table S1). This contrasting evidence may, however,  
81 correspond to different types of niche changes, biological and/or methodological study  
82 contexts, data types, species characteristics or methods being used (Peterson &  
83 Nakazawa, 2008; Mandle *et al.*, 2010; Peterson, 2011; Soberon & Peterson, 2011;  
84 Broennimann *et al.*, 2012) (Table S1), which confounding effects prevent sound  
85 interpretation of the possible mechanisms behind niche changes. Unification of the  
86 analytic context and metrics used, and a well-balanced comparison across different  
87 species, taxonomic groups, environmental spaces and geographic areas (Pearman *et al.*,  
88 2008a; Kueffer *et al.*, 2013), may contribute to reconcile conflicting evidence in  
89 observational studies of biological invasions requires.

90 Here, we build on a review of niche changes reported in empirical invasion studies  
91 (Table S1) to formalize a new framework that unifies the analytical context (Box 1, Figs  
92 1 and 2), clarifies the role of the niche-biotope duality (Box 2) (Colwell & Rangel, 2009;  
93 Soberón & Nakamura, 2009), and helps to identify potential factors influencing niche  
94 change between ranges. The central idea of this framework is to decompose a niche  
95 comparison between native and exotic ranges into its three basic components: niche  
96 unfilling, niche stability and niche expansion (Box 3, Fig. 2) (Petitpierre *et al.*, 2012). We  
97 present these elements and discuss them along with the importance of taking into  
98 account the available environment, distinguishing analog from non-analog climatic  
99 conditions between ranges (Box 4), and accounting for niche factors and niche dynamics  
100 at finer resolution. We conclude with recommendations on using the proposed  
101 framework for future niche change studies.

102

### 103 **Niche changes and associated metrics**

104 *Which niche is measured from field observations?*

105 The realized climatic niche quantified from field observations is determined by biotic  
106 constraints on the fundamental eco-physiological niche, population dynamics (e.g.  
107 source-sink dynamics) and dispersal limitations (i.e. accessibility; Box 2) (Pulliam, 2000;  
108 Soberon, 2007; Barve *et al.*, 2011), but it is also constrained by the availability of the  
109 environment in the areas (Box 4) at the timescale considered in the study (i.e. some  
110 conditions can be available at one time in one area, but not earlier or later) (Jackson &  
111 Overpeck, 2000; Mandle *et al.*, 2010). A change in this realized niche can thus result  
112 from adaptive evolution occurring in the colonized range (Sax *et al.*, 2007; Alexander &  
113 Edwards, 2010) or from changes in biotic interactions, dispersal limitations, or from  
114 pre-adaptation to conditions not (anymore) available in the initial range at the time of  
115 the study but available in the colonized range (Pearman *et al.*, 2008a). Hereafter, we  
116 consider a niche shift as any change of the realized niche, i.e. the niche as measured by  
117 climatic characteristics at sites of species occurrence in the field. It thus includes  
118 implicitly any potential change of the fundamental niche, although with such empirical  
119 data, a change caused by evolution of physiological tolerance cannot be differentiated  
120 from a change due to other factors (Broennimann *et al.*, 2007; Soberon & Peterson,  
121 2011).

122 *Two main approaches to quantifying niche changes*

123 Two main approaches have been used so far to compare niches between ranges, based  
124 on direct observations or on model predictions (Broennimann *et al.*, 2012) (Fig. 3, Table  
125 S1, suppl. mat.). The first approach uses observations directly and compares the  
126 difference in environmental attributes of the sites where the species occurs between the  
127 native and exotic ranges in environmental space. This comparison can be done either  
128 through univariate (e.g. (Lauzeral *et al.*, 2011)) or multivariate tests (e.g. in a reduced  
129 PCA space (Broennimann *et al.*, 2007); Fig. 3a). Such a direct approach does not rely on  
130 any underlying model that relates the occurrences to the environment. The approach  
131 can be considerably improved by calculating smooth densities of species occurrences in  
132 a gridded environmental space, as a way to avoid unrealistic 'holes' in a niche due to low  
133 sampling effort (Broennimann *et al.*, 2012) (see also (Guisan *et al.*, 2012; Webber *et al.*,  
134 2012)). The second approach relies instead on the outcomes of ecological niche models  
135 (ENMs (Peterson *et al.*, 2011); also called species distribution models, SDMs (Guisan &  
136 Thuiller, 2005)), and compares the overlap of reciprocal predictions of geographic  
137 distributions (i.e. predicting the invaded distribution with the model fitted in the native  
138 range, and vice-versa), usually comparing in the exotic range the two predictions by the  
139 models fitted in each range (Fitzpatrick *et al.*, 2007; Warren *et al.*, 2008; Warren *et al.*,  
140 2010) (Fig. 3b). Specialized software has been developed for niche comparisons based  
141 on the ENM approach (ENMTool (Warren *et al.*, 2010)). Comparative analyses with  
142 virtual species, for which distributions and niche overlap are known, showed that the  
143 first approach (ordination) quantified niche overlap overall more accurately than the  
144 second (ENM) (Broennimann *et al.*, 2012); however, the ordination approach provides a  
145 mathematically less formalized representation of the niche and is less able to optimize  
146 the weighting of the different environmental factors based on their relevance for a  
147 species' ecology. The ENM approach is particularly useful to assess ENM transferability  
148 between ranges (Randin *et al.*, 2006). Thus, although both approaches have strengths  
149 and weaknesses (Broennimann *et al.*, 2012), comparisons of niche change results  
150 between studies (meta-analyses) should include preferentially those based on  
151 ordinations, and at least make clear which approach was used (see Table S1).

152 *Different components of niche change: centroid shift, overlap, expansion and unfilling*

153 From either of these approaches, different niche change metrics can be calculated, at  
154 two levels of analyses – pooled ranges and range-specific (Box 3). The most commonly  
155 used metrics so far measure either a shift of the niche centroid, C (mean position; e.g.  
156 using Euclidean distance (Broennimann *et al.*, 2007)), or a change in the overlap, O,  
157 between the two niches (e.g. using Schoener's D (Warren *et al.*, 2008) or minimum  
158 convex polygons (Gallagher *et al.*, 2010)), and they are usually calculated in relation to  
159 the entire realized niche between two ranges (i.e. pooled; Box 3). However, a niche  
160 change detected in one of these two ways can result from multiple situations (Fig. 1):

161 (i) a change of the niche envelope (overlap  $\neq 1$ ) due to symmetric niche expansion or  
162 contraction (hereafter called 'unfilling' in the case of invasions, because it corresponds  
163 to a part of the native niche that was not filled) in climatic space, thus not shifting the  
164 niche centroid (Fig. 1a); a species may expand both to warmer and colder conditions in a  
165 way that the average temperature-related niche position remains stable as it is observed  
166 for common spotted knapweed invading North America (*Centaurea stoebe*, Fig. S1a,  
167 suppl. mat.);

168 (ii) a change of the niche centroid with displacement of the niche envelope (Fig. 1b-d)  
169 due to niche unfilling (e.g. black cherry tree invading Europe, *Prunus serotina*, Fig. S1b)  
170 and/or expansion (e.g. desert false indigo invading Europe, *Amorpha fruticosa*, Fig. S1c)  
171 in the invaded range; or

172 (iii) a change of the niche centroid only, without niche expansion or unfilling, due to a  
173 change of the density of occurrences within the same niche envelope in climatic space  
174 (Fig. 1e). The latter case can result from changes in competition, limited dispersal or  
175 availability of environmental conditions in the exotic range that reduce the density of  
176 species occurrences in some part of the niche space (Soberon & Peterson, 2011),  
177 changing the position of the centroid with only a weak impact on the niche limits, as  
178 shown for pinweed invading North America (*Erodium cicutarium*, Fig. S1d). Thus, a shift  
179 of the niche centroid between the native and the exotic range (Fig. 1b-e) can provide a  
180 first indication that a niche change occurred, but it is not sufficient to interpret its exact  
181 nature. And, reciprocally, an absence of a shift of the niche centroid does not mean that  
182 no niche shift occurred.

183 New indices were thus required to decompose niche comparisons to reveal two distinct  
184 components of niche changes: *expansion* and *unfilling* (Box 3, Fig. S1) (Petitpierre *et al.*,

185 2012). *Unfilling* (U) most commonly corresponds to the proportion of the native niche  
186 non-overlapping with the exotic niche, and *expansion* (E) refers to the proportion of the  
187 exotic niche non-overlapping with the native niche. These indices, as just defined,  
188 measure changes that are relative to one of the ranges (native or exotic), but they can  
189 also be measured with regard to the entire species distribution, where native and exotic  
190 ranges are pooled (Box 3). The pooled versions of E and U ( $E_p$  and  $U_p$  in Box 3) thus  
191 inform us about the species niche dynamic at the global scale of the study, but convey  
192 less information about our ability to predict species invasions from the native range  
193 (Box 3). E and U (and equivalently  $E_p$  and  $U_p$ ) are recently published indices (Petitpierre  
194 *et al.*, 2012) that can easily be calculated from the same two main approaches previously  
195 described (Broennimann *et al.*, 2012), but provide much more information than simple  
196 overlap or centroid changes. Studies that found overall niche conservatism for invaders  
197 relied consistently on such complete set of niche change metrics (Table S1, suppl. mat.).  
198 Later, we will refer to the whole set of niche change metrics as the COUE scheme (Box  
199 3).

#### 200 *Dealing with available and non-analog climates between ranges*

201 The availability of climatic conditions in geographic space matters when quantifying  
202 niche changes between ranges. Due to the niche-biotope duality (i.e. the correspondence  
203 between environmental and geographic spaces; Box 2), some conditions common in the  
204 exotic range may be rare in the native range (or the converse; Box 4) so that, without  
205 correction, one may detect niche shifts (measured with centroid change or overlap of  
206 percentile envelopes) only because these conditions are more or less available in one  
207 range than in the other (Soberón & Nakamura, 2009; Soberon & Peterson, 2011).  
208 Accounting for environmental availability is thus necessary and has been done so far in  
209 two ways. First, niche change metrics can be corrected by the distribution of the  
210 available environment, either by comparing the overlap between native and exotic  
211 niches with the overlap between native and exotic ranges (Mandle *et al.*, 2010), or by  
212 transforming species densities in the environmental space into species “occupancies”  
213 (i.e. the ratio of density of species to the density of available environment (Broennimann  
214 *et al.*, 2012); see also (Dormann *et al.*, 2010)). Second, niche metrics can be calculated  
215 only within the most common environments shared between native and exotic ranges  
216 (say within the shared portion of the 75<sup>th</sup> percentiles encompassing the prevailing



217 conditions in each range (Petitpierre *et al.*, 2012)). Removal of rare climates is however  
218 likely to have a strong impact on the results (with either approach) when the two ranges  
219 show important differences in climate availability. In this case we advise comparing  
220 analyses across a range of percentiles (say 75, 80, 85, 90, 95 and 100%) in order to see  
221 how the quantification of niche change can be affected by various levels of trimming (see  
222 suppl. online mat. in (Petitpierre *et al.*, 2012)) and to understand the implications  
223 (specific to each case study) for the interpretation of niche changes.

224 An extreme case of climate non-availability is when climate conditions exist only in one  
225 of the two ranges (Box 4) (Williams & Jackson, 2007). These non-analog climates  
226 represent a severe problem when calculating niche change metrics, because no insight  
227 on the biology of the species in these non-analog climates can be learned from a  
228 comparison between ranges. This is because colonization of portions of environmental  
229 space not present in the native range cannot be considered unambiguously as resulting  
230 from niche evolution in the exotic range, and the interpretation of these situations thus  
231 remains speculative (Mandle *et al.*, 2010; Soberon & Peterson, 2011). A scientifically  
232 more rigorous approach to assess niche expansion, therefore, is to restrict the analyses  
233 to the shared, analog climatic conditions between the native and exotic ranges (e.g.  
234 (Petitpierre *et al.*, 2012)), and to provide measures of expansion in non-analog  
235 situations separately (Guisan *et al.*, 2012). Studies that restricted their analyses to  
236 analog environments found niche conservatism to be dominant among invader species  
237 (Table S1, suppl. mat.). Complementary experimental approaches would then be needed  
238 to determine whether, for instance, expansions in non-analog conditions may represent  
239 a change of the fundamental niche (Pearman *et al.*, 2008a). This issue is particularly  
240 important because non-analog climates not only occur across space but will also occur  
241 over time due to climate change (Williams & Jackson, 2007). This is also the reason why  
242 projections of ecological models in non-analogous climates are considered unreliable  
243 (Fitzpatrick & Hargrove, 2009; Mandle *et al.*, 2010). Still, colonization of non-analog  
244 climates in the exotic range may represent relevant situations to consider from a  
245 management perspective, calling for separate ENM projections in both analog and non-  
246 analog climates in the invaded range (through fitting ENMs with pooled data from the  
247 native and exotic range (Broennimann & Guisan, 2008)).

248

249 **What other factors could affect the quantification of climatic niche changes?**

250 *Range unfilling in the native range*

251 Geographic range unfilling (not to be confused with niche unfilling) – i.e. when a species  
252 does not occupy all the geographic locations that have suitable conditions within its  
253 climatic niche – can occur in the native range as a result of non-equilibrium situations,  
254 such as ongoing post-glacial recolonization during the Holocene (Normand *et al.*, 2011),  
255 and can potentially affect the quantification of niche change. This problem is also known  
256 as the ‘accessible area’ issue (Barve *et al.*, 2011). But geographic range unfilling does not  
257 necessarily lead to niche unfilling in environmental space. For instance, it was shown  
258 that range unfilling can nonetheless translate into complete climatic niche filling for  
259 some tree species in Europe (Randin *et al.*, 2013). Range unfilling particularly affects  
260 niche quantification if the climates present in the unfilled geographic space are rare  
261 and/or not well represented – or even absent – in other parts of the range. However,  
262 published analyses generally calculate range filling based on a geographic projection of  
263 the realized niche at the time of the study (e.g. (Normand *et al.*, 2011)), and thus these  
264 documented cases of range unfilling cannot translate into niche unfilling. But range  
265 unfilling measured in other ways - e.g. field common garden experiments located  
266 beyond a species’ current geographic and climatic range (Alexander *et al.*, 2012;  
267 Alexander, 2013; Hargreaves *et al.*, 2014) - may reveal niche unfilling.

268 *Biased or incomplete sampling of species distributions*

269 Another issue relates to the type and quality of species distribution data. Although it is  
270 important to cover an entire species’ niche to assess niche change without bias, its  
271 complete native and exotic distribution ranges need not necessarily be considered.  
272 Because of the niche-biotope duality (Colwell & Rangel, 2009; Soberón & Nakamura,  
273 2009) (Box 2), the climatic niche of a species might well be fully captured even if only a  
274 part of its geographical distribution is sampled. However, and similarly to the issue of  
275 range unfilling, when geographic truncation leads to environmental truncation (Raes,  
276 2012), niche change studies based on both ordinations and ENMs (Fig. 3) should be  
277 considered with care, because their conclusions will only be applicable to the climate  
278 space investigated and within analog climatic combinations between the two ranges. In  
279 these situations, approaches based on ecological niche models (ENMs (Guisan &  
280 Thuiller, 2005; Peterson *et al.*, 2011); Fig. 3) may be less reliable for spatial predictions,

281 as they rely heavily on fitted species-environment response curves that could be biased  
282 (Thuiller *et al.*, 2004; Raes, 2012). In addition to environmental truncation, bias or  
283 errors in the geographic sampling of the distribution of a species may also bias  
284 measures of niche change. For instance, coarse atlas distribution data may portray a  
285 species in areas where it does not exist, while occurrence data (e.g. from herbaria) may  
286 under-represent or omit areas where the species occurs, both possibly affecting niche  
287 quantification.

### 288 *Beyond macroclimate: microclimate and non-climatic factors*

289 Climate is often seen as the main factor driving species distributions at large scales  
290 (Guisan & Thuiller, 2005), and most global-scale studies of niche changes in native  
291 (Pearman *et al.*, 2008b; Crisp *et al.*, 2009; Maiorano *et al.*, 2012) and exotic species  
292 (Broennimann *et al.*, 2007; Gallagher *et al.*, 2010; Medley, 2010; Lauzeral *et al.*, 2011;  
293 Petitpierre *et al.*, 2012) looked at changes in macroclimate (i.e. the coarse and large-  
294 scale climate that usually determines biomes). This primary role of macroclimate does  
295 not prevent finer climatic characteristics or other abiotic factors from affecting species  
296 distributions, such as the restriction to specialized habitats (e.g. mountain  
297 microclimates, stream banks or particular soil types) that must generally be  
298 characterized at a finer spatial grain (e.g. 1 km x 1 km) than that typically used in  
299 macroclimatic studies. Niche changes may be particularly observed in non-climatic  
300 components (such as soils) of a species' niche. For instance, Bertrand *et al.* (2012)  
301 showed that a shift of the climatic niche centroid can be observed when soil variables  
302 are included in the analyses. When shaping the distribution in the native range, these  
303 micro-scale factors could thus result in the detection of apparent macroclimatic niche  
304 expansion in the exotic range for two reasons: (i) part of the native macroclimate might  
305 not be occupied by the species due to spatial correlation with factors that hinder its  
306 occurrence (Bertrand *et al.*, 2012); or (ii) a species might occur under conditions in the  
307 native range that, within the coarse cells of macro-climatic maps, are scattered and  
308 marginal (and thus smoothed and hindered in niche analyses based on mean values  
309 within coarse cells), but are dominant in the exotic range and thus only revealed there in  
310 the niche quantification, causing an apparent niche shift.

311 However, these factors will only modify measures of macroclimatic niche change if: (i)  
312 their geographic distribution matches a restricted portion of the climatic niche in the

313 native or exotic range, and (ii) this restricted portion is the one that shows niche change.  
314 In this regard, studies at a finer resolution (e.g. microclimate) and/or including non-  
315 climatic factors would be useful for a more detailed understanding of niche dynamics in  
316 invaded ranges. But to be complementary to the strict macroclimatic niche studies  
317 conducted so far, findings based on macroclimate alone should be presented and  
318 compared to findings when microclimatic and non-climatic components are added (as  
319 for analog/non-analog climates), so that their relative effect can be properly assessed  
320 (e.g. (Bertrand *et al.*, 2012)).

321

### 322 **Toward a unifying framework: conclusion and remaining challenges**

323 There has recently been a great diversity of studies examining climate niche change in  
324 exotic species (Table S1, suppl. mat.), some reporting dramatic niche changes  
325 (Broennimann *et al.*, 2007; Fitzpatrick *et al.*, 2007; Gallagher *et al.*, 2010; Medley, 2010).  
326 However, how many shifts occur in analog versus in non-analog climates, and whether  
327 these only occur in specific taxonomic groups or habitats, remains to be investigated.  
328 Among 36 studies including ca. 180 species, about 50% of the species showed overall a  
329 niche shift, with a higher prevalence among plants than animals, and a majority of the  
330 studies reporting niche shifts included only one or a few species (Table S1). It might  
331 therefore be that studies reporting a shift (rather than no shift) were preferentially  
332 published, especially considering that the only two studies that concluded overall niche  
333 conservatism among a large number of invader species used an ordination approach,  
334 relied on the most complete set of niche change metrics, and accounted for  
335 environmental availability (Table S1 in supplementary material). Therefore, conclusions  
336 on niche shifts likely depend strongly on the organisms, methods and data used, and  
337 generalization about the frequency and drivers of niche shifts can only be based on a  
338 standardized and rigorous approach for quantifying niche shifts within each group. This  
339 could ultimately allow concluding if there are identifiable trends among niche shifts, or if  
340 niche changes are very idiosyncratic (i.e. species specific). In order to promote such  
341 standardization in future studies, we recommend:

342 - Using at least ordination, rather than only ENM, approaches to quantify climatic niche  
343 changes (see (Broennimann *et al.*, 2012));

344 –Using as much as possible, within a same taxonomic group, the same set of variables  
345 used in previous studies on the same group, so that proper comparisons can be ensured;  
346 this does not prevent additionally testing niche changes with other sets of variables, if  
347 thought to be more meaningful to picture species’ niches in the group considered;

348 - Disentangling all possible situations of niche change through measures of niche  
349 unfilling and expansion in complement to centriod shift and overlap metrics, at the two  
350 possible analytical levels (COUE scheme; Box 3);

351 - Correcting these niche change metrics to account for the density of occurrences and  
352 the available environment in both ranges (or time periods);

353 - Assessing whether niche metrics change when excluding rare climates along a range of  
354 percentiles, and when considering analog and non-analog environments separately; this  
355 will ensure retaining all the necessary information for further interpretation and  
356 comparison of results from different studies.

357 We suggest three important remaining challenges for studies of realized niche changes  
358 during biological invasions:

359 1) Assessing climatic niche changes at finer scales and in combination with other non-  
360 climatic factors, such as differences in soils (Bertrand *et al.*, 2012), biota, and  
361 disturbances between the native and exotic range. High-resolution data are becoming  
362 increasingly available and standardized to be comparable across large spatial areas.  
363 They constitute avenues to provide complementary answers to questions on  
364 macroclimate niche changes, and to improve our ability to predict and anticipate  
365 invasions.

366 2) Assessing invasions in non-analog environments has been poorly addressed so far. As  
367 these situation cannot be predicted from the native range with static approaches, and  
368 thus their interpretations remain speculative, they require mechanistic approaches (e.g.  
369 (Kearney & Porter, 2009)) or experiments (see below). It is however a promising field of  
370 investigation that may deliver invaluable insights on colonization processes in non-  
371 analog situations while also improving assessments of biodiversity under future climate  
372 changes (Williams & Jackson, 2007). Retrospective studies that examine the details of  
373 invasion success and failure into particular non-analog climates, relative to the native

374 climatic niche, could inform us of possible predictors of invasion into non-analog  
375 climates (e.g. for niche-based spatial predictions) (Guisan *et al.*, 2012).

376 3) Although correlative niche shift studies of exotic species may guide experimental  
377 studies (Kueffer *et al.*, 2013), a dual approach has been rare so far (but see (Hill *et al.*,  
378 2013)). Experimental studies on populations found in geographic areas where niche  
379 expansion occurred in the exotic range are needed to rigorously identify the related  
380 ecological or evolutionary causes, e.g. through rapid evolution (Sax *et al.*, 2007;  
381 Alexander, 2013), increased phenotypic plasticity (Hahn *et al.*, 2012) or biotic  
382 interactions (e.g. enemy-release) (Alexander & Edwards, 2010). Similarly, information  
383 about unfilling can help identify interesting model systems (Kueffer *et al.*, 2013) for  
384 studying why some habitats and landscapes are more resistant to invasions, e.g. due to  
385 dispersal limitations (Barve *et al.*, 2011) or abiotic or biotic resistance (Richardson,  
386 2011).

387 We expect that systematic use of this framework will substantially advance  
388 generalization about niche change, not only in invasion studies (including pests and  
389 diseases) but also in studies of niche conservatism between disjoint distributions (e.g.  
390 arctic-alpine) (Pellissier *et al.*, 2013) or across time in response to global change  
391 (Maiorano *et al.*, 2012).

392

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402

### 402 **References**

403

404 Ackerly, D.D. (2003) Community assembly, niche conservatism, and adaptive evolution  
405 in changing environments. *International Journal of Plant Sciences*, **164**, S165-  
406 S184.

- 407 Alexander, J.M. (2013) Evolution under changing climates: climatic niche stasis despite  
408 rapid evolution in a non-native plant. *Proceedings of the Royal Society B*, **280**,  
409 1446.
- 410 Alexander, J.M. & Edwards, P.J. (2010) Limits to the niche and range margins of alien  
411 species. *Oikos*, **119**, 1377-1386.
- 412 Alexander, J.M., van Kleunen, M., Ghezzi, R. & Edwards, P.J. (2012) Different genetic  
413 clines in response to temperature across the native and introduced ranges of a  
414 global plant invader. *Journal of Ecology*, **100**, 771-781.
- 415 Araujo, M.B. & Guisan, A. (2006) Five (or so) challenges for species distribution  
416 modelling. *Journal of Biogeography*, **33**, 1677-1688.
- 417 Barve, N., Barve, V., Jimenez-Valverde, A., Lira-Noriega, A., Maher, S.P., Peterson, A.T.,  
418 Soberon, J. & Villalobos, F. (2011) The crucial role of the accessible area in  
419 ecological niche modeling and species distribution modeling. *Ecological*  
420 *Modelling*, **222**, 1810-1819.
- 421 Bertrand, R., Perez, V. & Gegout, J.C. (2012) Disregarding the edaphic dimension in  
422 species distribution models leads to the omission of crucial spatial information  
423 under climate change: the case of *Quercus pubescens* in France. *Global Change*  
424 *Biology*, **18**, 2648-2660.
- 425 Broennimann, O. & Guisan, A. (2008) Predicting current and future biological invasions:  
426 both native and invaded ranges matter. *Biology Letters*, **4**, 585-589.
- 427 Broennimann, O., Treier, U.A., Muller-Scharer, H., Thuiller, W., Peterson, A.T. & Guisan, A.  
428 (2007) Evidence of climatic niche shift during biological invasion. *Ecology Letters*,  
429 **10**, 701-709.
- 430 Broennimann, O., Fitzpatrick, M.C., Pearman, P.B., Petitpierre, B., Pellissier, L., Yoccoz,  
431 N.G., Thuiller, W., Fortin, M.J., Randin, C., Zimmermann, N.E., Graham, C.H. &  
432 Guisan, A. (2012) Measuring ecological niche overlap from occurrence and spatial  
433 environmental data. *Global Ecology and Biogeography*, **21**, 481-497.
- 434 Colwell, R.K. & Rangel, T.F. (2009) Hutchinson's duality: The once and future niche.  
435 *Proceedings of the National Academy of Sciences of the United States of America*,  
436 **106**, 19651-19658.
- 437 Crisp, M.D., Arroyo, M.T.K., Cook, L.G., Gandolfo, M.A., Jordan, G.J., McGlone, M.S., Weston,  
438 P.H., Westoby, M., Wilf, P. & Linder, H.P. (2009) Phylogenetic biome conservatism  
439 on a global scale. *Nature*, **458**, 754-756.
- 440 Dormann, C.F., Gruber, B., Winter, M. & Hermann, D. (2010) Evolution of climatic niches  
441 in European mammals? *Biology Letters*, **6**, 229-232.
- 442 Elith, J., Kearney, M. & Phillips, S. (2010) The art of modelling range-shifting species.  
443 *Methods in Ecology and Evolution*, **1**, 330-342.
- 444 Elton, C.S. (1958) *The ecology of invasions by animals and plants*. Methuen, London.

- 445 Fitzpatrick, M.C. & Hargrove, W.W. (2009) The projection of species distribution models  
446 and the problem of non-analog climate. *Biodiversity and Conservation*, **18**, 2255-  
447 2261.
- 448 Fitzpatrick, M.C., Weltzin, J.F., Sanders, N.J. & Dunn, R.R. (2007) The biogeography of  
449 prediction error: why does the introduced range of the fire ant over-predict its  
450 native range? *Global Ecology and Biogeography*, **16**, 24-33.
- 451 Gallagher, R.V., Beaumont, L.J., Hughes, L. & Leishman, M.R. (2010) Evidence for climatic  
452 niche and biome shifts between native and novel ranges in plant species  
453 introduced to Australia. *Journal of Ecology*, **98**, 790-799.
- 454 Guisan, A. & Thuiller, W. (2005) Predicting species distribution: offering more than  
455 simple habitat models. *Ecology Letters*, **8**, 993-1009.
- 456 Guisan, A., Petitpierre, B., Broennimann, O., Kueffer, C., Randin, C. & Daehler, C. (2012)  
457 Response to comment on "Climatic niche shifts are rare among terrestrial plant  
458 invaders". *Science*, **338**, 193.
- 459 Guisan, A., Tingley, R., Baumgartner, J.B., Naujokaitis-Lewis, I., Sutcliffe, P.R., Tulloch,  
460 A.I.T., Regan, T.J., Brotons, L., McDonald-Madden, E., Mantyka-Pringle, C., Martin,  
461 T.G., Rhodes, J.R., Maggini, R., Setterfield, S.A., Elith, J., Schwartz, M.W., Wintle,  
462 B.A., Broennimann, O., Austin, M., Ferrier, S., Kearney, M.R., Possingham, H.P. &  
463 Buckley, Y.M. (2013) Predicting species distributions for conservation decisions.  
464 *Ecology Letters*, **16**, 1424-1435.
- 465 Hahn, M.A., van Kleunen, M. & Müller-Schärer, H. (2012) Increased phenotypic plasticity  
466 to climate may have boosted the invasion success of polyploid *Centaurea stoebe*.  
467 *PLoS ONE*, **7**, e50284.
- 468 Hargreaves, A.L., Samis, K.E. & Eckert, C.G. (2014) *The American Naturalist*, **183**, 157-  
469 173.
- 470 Hill, M.P., Chown, S.L. & Hoffmann, A.A. (2013) A predicted niche shift corresponds with  
471 increased thermal resistance in an invasive mite, *Halotydeus destructor*. *Global  
472 Ecology & Biogeography*, **22**, 942-951.
- 473 Hutchinson, G.E. (1957) Population Studies - Animal Ecology and Demography -  
474 Concluding Remarks. *Cold Spring Harbor Symposia on Quantitative Biology*, **22**,  
475 415-427.
- 476 Jackson, S.T. & Overpeck, J.T. (2000) Responses of plant populations and communities to  
477 environmental changes of the late Quaternary. *Paleobiology*, **26**, 194-220.
- 478 Kearney, M. & Porter, W. (2009) Mechanistic niche modelling: combining physiological  
479 and spatial data to predict species' ranges. *Ecology Letters*, **12**, 334-350.
- 480 Kueffer, C., Pyšek, P. & Richardson, D.M. (2013) Integrative invasion science: model  
481 systems, multi-site studies, focused meta-analysis, and invasion syndromes. *New  
482 Phytologist*, **200**, 615-633.



- 483 Lauzeral, C., Leprieur, F., Beauchard, O., Duron, Q., Oberdorff, T. & Brosse, S. (2011)  
 484 Identifying climatic niche shifts using coarse-grained occurrence data: a test with  
 485 non-native freshwater fish. *Global Ecology and Biogeography*, **20**, 407-414.
- 486 Maiorano, L., Cheddadi, R., Zimmerman, N.E., Pellissier, L., Petitpierre, B., Pottier, J.,  
 487 Laborde, H., Hurdu, B.I., Pearman, P.B., Psomas, A., Singarayer, J.S., Broennimann,  
 488 O., Vittoz, P., Dubuis, A., Edwards, M.E., Binney, H.A. & Guisan, A. (2012) Building  
 489 the niche through time: using 13,000 years of data to predict the effects of  
 490 climate change on three tree species in Europe. *Global Ecology & Biogeography*, **in**  
 491 **press**
- 492 Mandle, L., Warren, D.L., Hoffmann, M.H., Peterson, A.T., Schmitt, J. & von Wettberg, E.J.  
 493 (2010) Conclusions about Niche Expansion in Introduced *Impatiens walleriana*  
 494 Populations Depend on Method of Analysis. *Plos One*, **5**
- 495 Medley, K.A. (2010) Niche shifts during the global invasion of the Asian tiger mosquito,  
 496 *Aedes albopictus* Skuse (Culicidae), revealed by reciprocal distribution models.  
 497 *Global Ecology and Biogeography*, **19**, 122-133.
- 498 Normand, S., Ricklefs, R.E., Skov, F., Bladt, J., Tackenberg, O. & Svenning, J.C. (2011)  
 499 Postglacial migration supplements climate in determining plant species ranges in  
 500 Europe. *Proceedings of the Royal Society B-Biological Sciences*, **278**, 3644-3653.
- 501 Pearman, P.B., Guisan, A., Broennimann, O. & Randin, C.F. (2008a) Niche dynamics in  
 502 space and time. *Trends in Ecology & Evolution*, **23**, 149-158.
- 503 Pearman, P.B., Randin, C.F., Broennimann, O., Vittoz, P., van der Knaap, W.O., Engler, R.,  
 504 Le Lay, G., Zimmermann, N.E. & Guisan, A. (2008b) Prediction of plant species  
 505 distributions across six millennia. *Ecology Letters*, **11**, 357-369.
- 506 Pellissier, L., Bråthen, K.A., Vittoz, P., Yoccoz, N.G., Dubuis, A., Meier, E.S., Zimmermann,  
 507 N.E., Randin, C.F., Thuiller, W., Garraud, L., Van Es, J. & Guisan, A. (2013) Thermal  
 508 niches are more conserved at cold than warm limits in arctic-alpine plant species.  
 509 *Global Ecology and Biogeography*, **22**, 933-941.
- 510 Peterson, A.T. (2011) Ecological niche conservatism: a time-structured review of  
 511 evidence. *Journal of Biogeography*, **38**, 817-827.
- 512 Peterson, A.T. & Nakazawa, Y. (2008) Environmental data sets matter in ecological niche  
 513 modelling: an example with *Solenopsis invicta* and *Solenopsis richteri*. *Global*  
 514 *Ecology and Biogeography*, **17**, 135-144.
- 515 Peterson, A.T., Soberón, J., Pearson, R.G., Anderson, R.P., Martinez-Meyer, E., Nakamura,  
 516 M. & Araujo, M.B. (2011) *Ecological niches and geographic distributions*. Princeton  
 517 University Press, Princeton, USA.
- 518 Petitpierre, B., Kueffer, C., Broennimann, O., Randin, C., Daehler, C. & Guisan, A. (2012)  
 519 Climatic niche shifts are rare among terrestrial plant invaders. *Science*, **335**,  
 520 1344-1348.

- 521 Pulliam, H.R. (2000) On the relationship between niche and distribution. *Ecology Letters*,  
522 **3**, 349-361.
- 523 Raes, N. (2012) Partial versus Full Species Distribution Models. *Natureza & Conservação*,  
524 **10**, 127-138.
- 525 Randin, C.F., Dirnbock, T., Dullinger, S., Zimmermann, N.E., Zappa, M. & Guisan, A. (2006)  
526 Are niche-based species distribution models transferable in space? *Journal of*  
527 *Biogeography*, **33**, 1689-1703.
- 528 Randin, C.F., Paulsen, J., Vitasse, Y., Kollas, C., Wohlgemuth, T., Zimmermann, N.E. &  
529 Korner, C. (2013) Do the elevational limits of deciduous tree species match their  
530 thermal latitudinal limits? *Global Ecology and Biogeography*, **22**, 913-923.
- 531 Richardson, D.M. (ed.^eds) (2011) *Fifty years of invasion ecology: the legacy of Charles*  
532 *Elton*. Blackwell Publishing, Oxford.
- 533 Richardson, D.M. & Pysek, P. (2008) Fifty years of invasion ecology - the legacy of  
534 Charles Elton. *Diversity and Distributions*, **14**, 161-168.
- 535 Rödder, D. & Lötters, S. (2009) Niche shift versus niche conservatism? Climatic  
536 characteristics of the native and invasive ranges of the Mediterranean house  
537 gecko (*Hemidactylus turcicus*). *Global Ecology and Biogeography*, **18**, 674-687.
- 538 Sax, D.F., Early, R. & Bellemare, J. (2013) Niche syndromes, species extinction risks, and  
539 management under climate change. *Trends in Ecology & Evolution*, **28**, 517-523.
- 540 Sax, D.F., Stachowicz, J.J., Brown, J.H., Bruno, J.F., Dawson, M.N., Gaines, S.D., Grosberg,  
541 R.K., Hastings, A., Holt, R.D., Mayfield, M.M., O'Connor, M.I. & Rice, W.R. (2007)  
542 Ecological and evolutionary insights from species invasions. *Trends in Ecology &*  
543 *Evolution*, **22**, 465-471.
- 544 Soberon, J. (2007) Grinnellian and Eltonian niches and geographic distributions of  
545 species. *Ecology Letters*, **10**, 1115-1123.
- 546 Soberon, J. & Peterson, A.T. (2011) Ecological niche shifts and environmental space  
547 anisotropy: a cautionary note. *Revista Mexicana de Biodiversidad*, **82**, 1348-1355.
- 548 Soberón, J. & Nakamura, M. (2009) Niches and distributional areas: concepts, methods,  
549 and assumptions. *Proceedings of the National Academy of Sciences of the United*  
550 *States of America*, **106**, 19644-19650.
- 551 Strubbe, D., Broennimann, O., Chiron, F. & Matthysen, E. (2013) Niche conservatism in  
552 non-native birds in Europe: niche unfilling rather than niche expansion. *Global*  
553 *Ecology and Biogeography*, **22**, 962-970.
- 554 Thuiller, W., Brotons, L., Araujo, M.B. & Lavorel, S. (2004) Effects of restricting  
555 environmental range of data to project current and future species distributions.  
556 *Ecography*, **27**, 165-172.

- 557 Venette, R.C., Kriticos, D.J., Magarey, R.D., Koch, F.H., Baker, R.H.A., Worner, S.P.,  
558 Raboteaux, N.N.G., McKenney, D.W., Dobesberger, E.J., Yemshanov, D., De Barro,  
559 P.J., Hutchison, W.D., Fowler, G., Kalaris, T.M. & Pedlar, J. (2010) Pest risk maps  
560 for invasive alien species: a roadmap for improvement. *Bioscience*, **60**, 349-362.
- 561 Warren, D.L., Glor, R.E. & Turelli, M. (2008) Environmental Niche Equivalency Versus  
562 Conservatism: Quantitative Approaches to Niche Evolution. *Evolution*, **62**, 2868-  
563 2883.
- 564 Warren, D.L., Glor, R.E. & Turelli, M. (2010) ENMTools: a toolbox for comparative studies  
565 of environmental niche models. *Ecography*, **33**, 607-611.
- 566 Webber, B.L., Le Maître, D.C. & Kriticos, D.J. (2012) Comments on "Climatic niche shifts  
567 are rare among terrestrial plant invaders". *Science*, **338**, 193.
- 568 Williams, J.W. & Jackson, S.T. (2007) Novel climates, no-analog communities, and  
569 ecological surprises. *Frontiers in Ecology and the Environment*, **5**, 475-482.
- 570 Willis, K.J. & McElwain, J.C. (2002) *The evolution of plants*. Oxford University Press.  
571  
572

573

574 **Boxes**

575 **Glossary box**

576 *Analog climate*: A combination of climate factors found in one area or time period that is  
577 within the envelope of climatic conditions found in a different area or time period used  
578 for comparison (Williams & Jackson, 2007). Contrary: 'non-analog climate'.

579 *Accessible range*: The geographic locations within a given area that are accessible to a  
580 species given its current distribution and the timescale considered in the study. It is thus  
581 conditional upon spatial configuration and the species' dispersal ability (Soberon, 2007;  
582 Barve *et al.*, 2011).

583 *Available environment*: the set of environmental conditions that exist in a given area  
584 (Jackson & Overpeck, 2000) (Box 3). Synonyms: 'realized environment' (whole range,  
585 not species-specific), 'background environment'.

586 *Ecological niche model* (ENM; also called species distribution or habitat suitability  
587 models): multivariate models fitting the niche of species by relating distribution  
588 observations with environmental variables measured at the same sites, and projected  
589 over a whole study area (see (Guisan & Thuiller, 2005; Peterson *et al.*, 2011)).

590 *Exotic niche*: The niche measured based on a species' distribution in the exotic range.  
591 Synonyms: 'naturalized niche', 'adventive niche', 'invaded niche' or 'invasive niche' (for  
592 invasive species).

593 *Exotic range*: The geographic range where a species is not native. Synonyms:  
594 'naturalized range', 'adventive range', 'invaded range' (for invasive species)

595 *Exotic species*: A species present in a region where it is not native, mostly due to human  
596 actions that enabled it to overcome biogeographical barriers (Richardson, 2011).  
597 Synonyms: 'alien species', 'non-native species', 'non-indigenous species', 'introduced  
598 species'.

599 *Fundamental niche*: The envelope of environmental (abiotic) conditions allowing  
600 populations to sustain themselves in an n-dimensional environmental space. It depicts  
601 the eco-physiological requirements of species (Soberon, 2007). Synonyms:  
602 'Physiological niche'.

603 *Native niche*: The niche measured in the native range.

604 *Native range*: The complete geographic area where an exotic species is native.

605 *Niche-biotope duality*: The reciprocal correspondence between the niche conditions in  
606 multidimensional environmental space and the physical locations that a species actually  
607 occupies in geographical space (derived from (Colwell & Rangel, 2009)).

608 *Niche centroid*: the mean niche position in n-dimensional environmental space.

609 *Niche conservatism*: The tendency for species to retain their niche in space and time.  
610 Synonyms: 'niche stability'.

611 *Niche envelope*: The envelope of conditions in multivariate environmental space defining  
612 a species' niche. The boundary of the envelope can be defined in many different ways  
613 (e.g. percentiles; see (Broennimann *et al.*, 2012)).

614 *Niche expansion*: Proportion of the exotic niche non-overlapping with the native niche.

615 *Niche overlap*: the intersection of two niches in n-dimensional environmental space.

616 *Niche shift*: A change in the centroid (see above) or limits of the niche envelop in  
617 environmental space. Synonyms: niche change.

618 *Niche stability*: Proportion of the exotic niche overlapping with the native niche

619 *Niche unfilling*: Proportion of the native niche non-overlapping with the exotic niche.

620 *Non-analog climate*: See 'analog' climate.

621 *Ordination*: statistical approach used to represent the arrangement of a series of objects  
622 described by multiple descriptor variables into a reduced multidimensional space which  
623 axes represent combinations of the initial variables (see PCA).

624 *PCA*: Principal component analysis, a classical ordination approach (see above).

625 *Potential niche*: The intersection between the fundamental niche and the realized  
626 environment (see (Jackson & Overpeck, 2000; Soberón & Nakamura, 2009)).

627 *Rare climate*: Climatic conditions poorly represented overall within an area during a  
628 given time period.

629 *Realized niche*: The environmental (abiotic) niche of a species as quantified from field  
630 observations, i.e. the fundamental niche modulated by biotic exclusions, population

631 dynamics (such as source-sink dynamics) and dispersal limitations (Soberon, 2007;  
632 Colwell & Rangel, 2009). Synonyms: 'Ecological niche'.

633 *Schoener's D*: The most common measure of niche overlap (see (Warren *et al.*, 2008;  
634 Broennimann *et al.*, 2012)).

635

### 636 **Box 1: The analytical context for quantifying niche shifts**

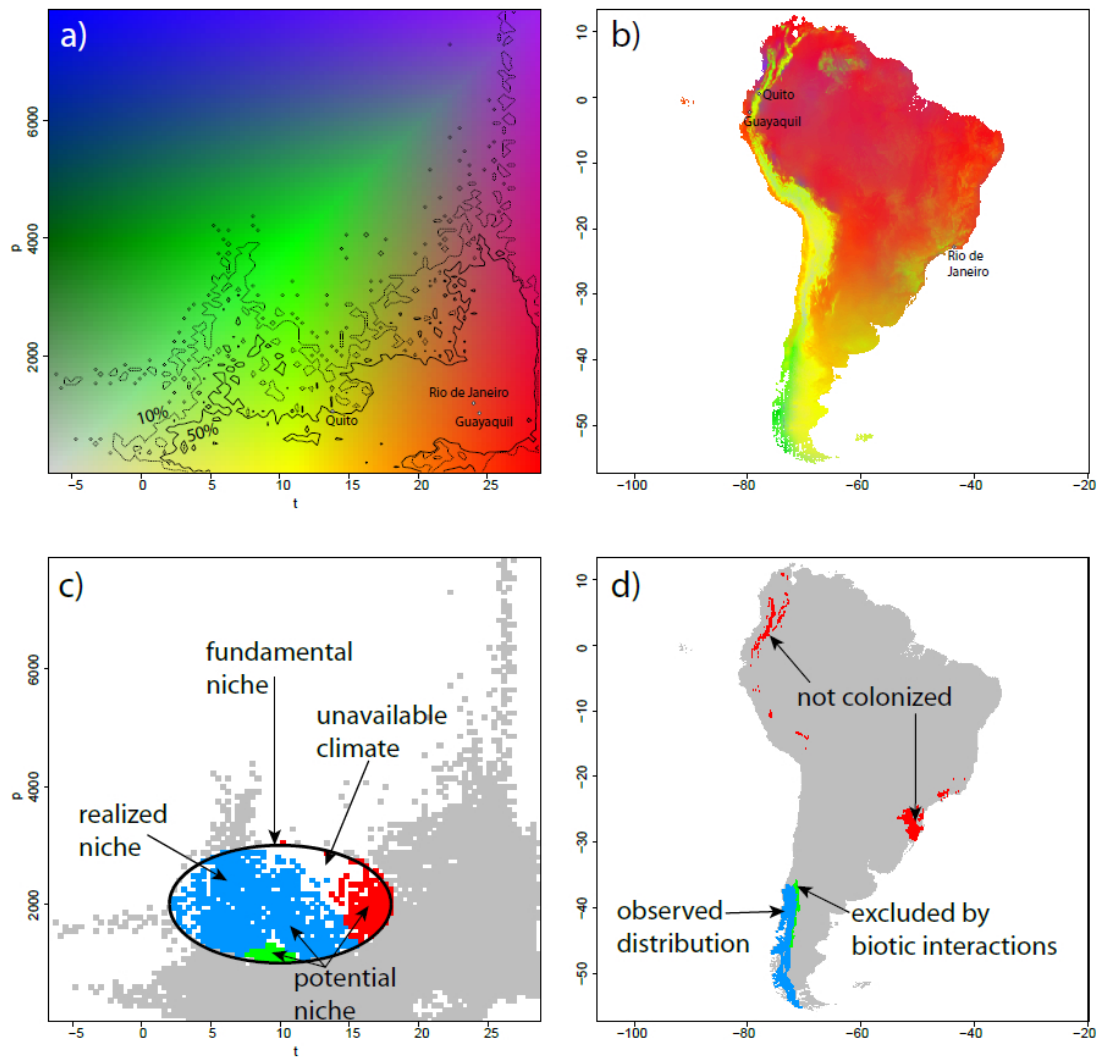
637 Assessing niche change between ranges is generally done by considering a species  
638 native in one area (its native range) and invading another (or several other)  
639 biogeographically separated area (the exotic range; e.g. (Petitpierre *et al.*, 2012)). This  
640 context could similarly apply to the same species in two (or more) time periods (e.g.  
641 (Maiorano *et al.*, 2012)). Regions large enough to include the entire (or large parts of)  
642 the native and exotic species' geographic distributions are usually considered for  
643 comparison. The choice of these areas will strongly condition the niche-biotope duality  
644 (Box 2), and thus the available environments (Fig. 2, Box 3), and ultimately the  
645 quantification of niche changes (Soberón & Nakamura, 2009; Soberon & Peterson,  
646 2011). Optimally, the studied ranges should encompass the species' complete  
647 geographic distribution in the native and introduced ranges that could potentially be  
648 reached by a species given its dispersal ability, i.e. the accessible areas (Barve *et al.*,  
649 2011). In practice we recommend defining areas with ecological relevance, such as  
650 biomes or ecoregions, and using species data (atlas or occurrences) well representing  
651 the focal species' range. The full multi-dimensional set of environmental conditions  
652 observed in one area/time period is the realized environment (Box 3; Box 2) (Jackson &  
653 Overpeck, 2000; Ackerly, 2003) and the envelop of conditions where the species is  
654 observed represents its realized environmental niche (Box 2) (Araujo & Guisan, 2006;  
655 Soberón & Nakamura, 2009).

656

### 657 **Box 2: Hutchinson's niche-biotope duality**

658 It is important to recall the niche (environmental space) versus biotope (geographic  
659 space) duality framework described by G.E. Hutchinson (see (Colwell & Rangel, 2009),  
660 Glossary). This duality means that there is no direct match between the topology of the  
661 niche space and the geographic distribution of a species (see figure below). The same  
662 combination of climate factors (colors in panel a of the figure) can occur in one or

663 several localities in geographic space (same colors in panel b), and locations close in  
664 environmental space can be far apart geographically and vice versa (Soberón &  
665 Nakamura, 2009). For instance, in South America, the cities of Quito and Guayaquil are  
666 close to each other but climatically far away, whereas Guayaquil and Rio are  
667 geographically far but climatically close (see figure below). Interpretation of niches and  
668 distributions of species thus requires careful screening of both spaces jointly (see figure  
669 a,c vs. b,d), with special attention to issues of dispersal limitations, biotic interactions  
670 and available environmental conditions (Colwell & Rangel, 2009). Blank areas in panel c  
671 of the figure below represent environments that are not available within the geographic  
672 range considered (here South America). The intersection of the available environment  
673 and the fundamental (i.e. physiological) limits of a species define its potential niche  
674 (Jackson & Overpeck, 2000; Soberón & Nakamura, 2009). Parts of this potential niche  
675 can be unoccupied by the species because of dispersal limitations (i.e. red areas in  
676 panels c and d) or exclusion by biotic interactions (i.e. green parts in panels c and d). As  
677 the potential niche rests on the hypothetical quantification of the fundamental niche  
678 (see (Sax *et al.*, 2013)), whereas we focus mainly on the realized niche here, we do not  
679 expand further on this concept. For a full theoretical development of the concepts and  
680 definitions of niches and distributional areas, with formal abbreviations, see Soberon &  
681 Nakamura (2009) and Peterson et al. (2011).



682

683 Hutchinson's duality framework, inspired by Soberon and Nakamura (2009) and  
 684 Soberon & Peterson (2011). The fundamental niche ellipse pictured in panel c) is  
 685 theoretical (artificially created) and could not be derived from field observations.

686

687

### 688 **Box 3: Metrics to quantify and decompose niche changes - The COUE scheme**

689

690 The niche space of an exotic species can be classified into three categories: niche space  
 691 occurring only in the exotic range (i.e. expansion,  $\epsilon$ ), in both exotic and native range (i.e.  
 692 stability,  $\sigma$ ) and only in the native range (i.e. unfilling,  $\upsilon$ ). Niche comparisons can then be  
 693 made at two levels: (i) relative to the entire niche of the species, pooled from the two  
 694 ranges (pooled ranges approach); or (ii) relative to the native or exotic ranges  
 695 separately (range-specific approach). The table below presents a unified terminology  
 696 (COUE, an acronym based on its main components, centroid shift, overlap, unfilling and



697 expansion, as defined below) for niche comparisons and related metrics of niche  
 698 changes.

699

Niche change component	Absolute component	Metric	
		Pooled ranges	Range-specific
Centroid shift	--	C	--
Expansion	$\varepsilon$	$E_p$	E
Stability	$\sigma$	$S_p (\approx O)$	$S_n, S_e$
Unfilling	$\upsilon$	$U_p$	U

700

701 Centroid shift measures the change in mean niche position (and thus mean intensity) in  
 702 the pooled ranges space, and thus no range-specific counterpart exists here. At the  
 703 pooled-range level, niche stability ( $S_p$ ) measures the proportion of niche assessed from  
 704 the pooled native and exotic occurrences (possibly transformed into ENM predictions or  
 705 densities in the environmental space, Fig. 3) present in both native and exotic ranges.  
 706 This is similar to the niche overlap (O) assessed through Schoener's D or Hellinger's I  
 707 (see (Warren *et al.*, 2008; Broennimann *et al.*, 2012)). The non-overlapping parts of the  
 708 two niches ( $1 - S_p$ ) can then be decomposed into global ratios of expansion ( $E_p$ ) and  
 709 unfilling ( $U_p$ ) based on the pooled ranges. Decomposing niche changes relative to the  
 710 pooled species distribution informs about the magnitude of niche changes at the global  
 711 scale (i.e. relative to the entire realized niche of the species), but may not be informative  
 712 about niche changes specific to either exotic or native ranges (as used e.g. in (Petitpierre  
 713 *et al.*, 2012)). For example, the exotic niche can be very small relative to the entire  
 714 pooled niche but entirely located in environments different from the native niche, in  
 715 which case  $E_p$  would be very small although the entire invaded niche is distinct from the  
 716 native niche and would hardly be predictable from the native range data. It is however  
 717 possible to quantify a ratio of expansion (E) and unfilling (U) relatively to the exotic or  
 718 native niches only, i.e. at the range-specific level. These inform us about the relative  
 719 importance of changes in each exotic and native niche. In turn, niche stability can be  
 720 assessed from the perspective of native or invaded niches separately, depending on  
 721 whether it complements the relative expansion or relative unfilling ratios ( $S_n = 1 - U$ ;  $S_e$   
 722  $= 1 - E$ , respectively).

723

724

725 **Box 4: The available climate and the analog/non-analog issue**

726 The available environment is a subset of all possible environmental combinations (Box  
727 2). The existence of non-available environments constrains niche shape and size  
728 (Jackson & Overpeck, 2000). For instance, places with very warm summer temperature  
729 (say >40°C) and very cold winter temperature (say <-20°C) do not currently exist on  
730 Earth (see Figure 1 in (Jackson & Overpeck, 2000)). When comparing the available  
731 environment in two areas, some habitats in one area (or time period) may be much  
732 more frequent or rare than in the other area (or time period), or some specific  
733 conditions found in one range may be totally absent from the other range. For instance,  
734 some very dry conditions of Western North America are not found in Western Europe  
735 (Broennimann *et al.*, 2007) and tropical conditions of the Tertiary in Europe are not  
736 observed anymore (Willis & McElwain, 2002). Conditions similar in two ranges or two  
737 time periods are called 'analog' and those differing 'non-analog' (Williams & Jackson,  
738 2007; Fitzpatrick & Hargrove, 2009) (or 'non-overlapping backgrounds' in (Soberon &  
739 Peterson, 2011)). Non-analog environments in an invaded range, or in the future,  
740 typically represent situations outside the range of values considered to quantify the  
741 native niche and not experienced by the species before invasion, and therefore lead to  
742 difficulty in interpreting niche shifts (Petitpierre *et al.*, 2012) and predicting species  
743 distributions (Fitzpatrick & Hargrove, 2009). Tools are available to define areas in the  
744 exotic range with climates analog to the native range. The simplest approach is to define  
745 a bounding box that encloses all the conditions present in the native range (e.g.  
746 BIOCLIM; Busby 1991). Any pixel in the exotic range outside of the bounding box range  
747 can be considered non-analogous. A more refined approach is the MESS analysis  
748 (Multivariate Environmental Similarity Surfaces)(Elith *et al.*, 2010), an index of  
749 similarity reporting the closeness of a point described by a set of climate attributes (e.g.  
750 a pixel in the exotic range) to the distribution of these attributes within a population of  
751 reference points (e.g. the native range).

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756 **Figure captions**

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758 **Figure 1:** Theoretical scenarios of realized niche changes in space (e.g. following  
759 invasions) or time (e.g. under climate change). Change of : (i) the niche envelop  
760 (expansion or contraction) without change of the niche centroid, due to symmetric niche  
761 change, i.e. in two opposite (a) or all directions in climatic space; (ii) the niche centroid  
762 with expansion (b, c) or displacement (d) of part of or the whole niche envelop , or (iii)  
763 the niche centroid only, due to a change of the density of occurrences within the same  
764 niche envelop in climatic space (e). The latter case would result in stability (no change)  
765 in figure 2. Observed changes are likely to be combinations of these cases.

766

767 **Figure 2:** Schematic 2-dimensional representation of the indices of niche change  
768 (unfilling, stability and expansion) presented in (2012) (see definitions in Box 3). Solid  
769 thin lines show the density of available environments (Box 4) in the native range (in  
770 green) and in the invaded range (in red). The gray area shows the most frequent  
771 environments common to both ranges (i.e. analog environments). The green and red  
772 thick lines show respectively the native and the invaded niches. Niche unfilling (U),  
773 stability ( $S_e$ ) and expansion (E) are shown respectively with green, blue and red hatched  
774 surfaces inside analog environments. The definition of a niche shift using the change of  
775 niche centroid only (inertia ratio, IR) is shown with a thick dotted arrow. In this context,  
776 the lower-case letters represent similar features in both graphs: a. available conditions  
777 in the native range, outside of the native niche and non-analog to the invaded range.. b.  
778 Conditions inside of the native niche but non-analog to the invaded range. c. Unfilling, i.e.  
779 conditions inside of the native niche but outside the invaded niche , possibly due to  
780 recent introduction combined with ongoing dispersal of the exotic species, which should  
781 at term fill these conditions. d. Niche stability, i.e. conditions filled in both native and  
782 invaded range. e. Niche expansion, i.e. conditions inside the invaded niche but outside  
783 the native one, due to ecological or evolutionary change in the invaded range. f.  
784 Conditions inside of the invasive niche but non-analog to the native range. g. Available

785 conditions in the invaded range but outside of the invasive niche and non-analog to the  
786 native range. h. Analog conditions between the native and invaded ranges.

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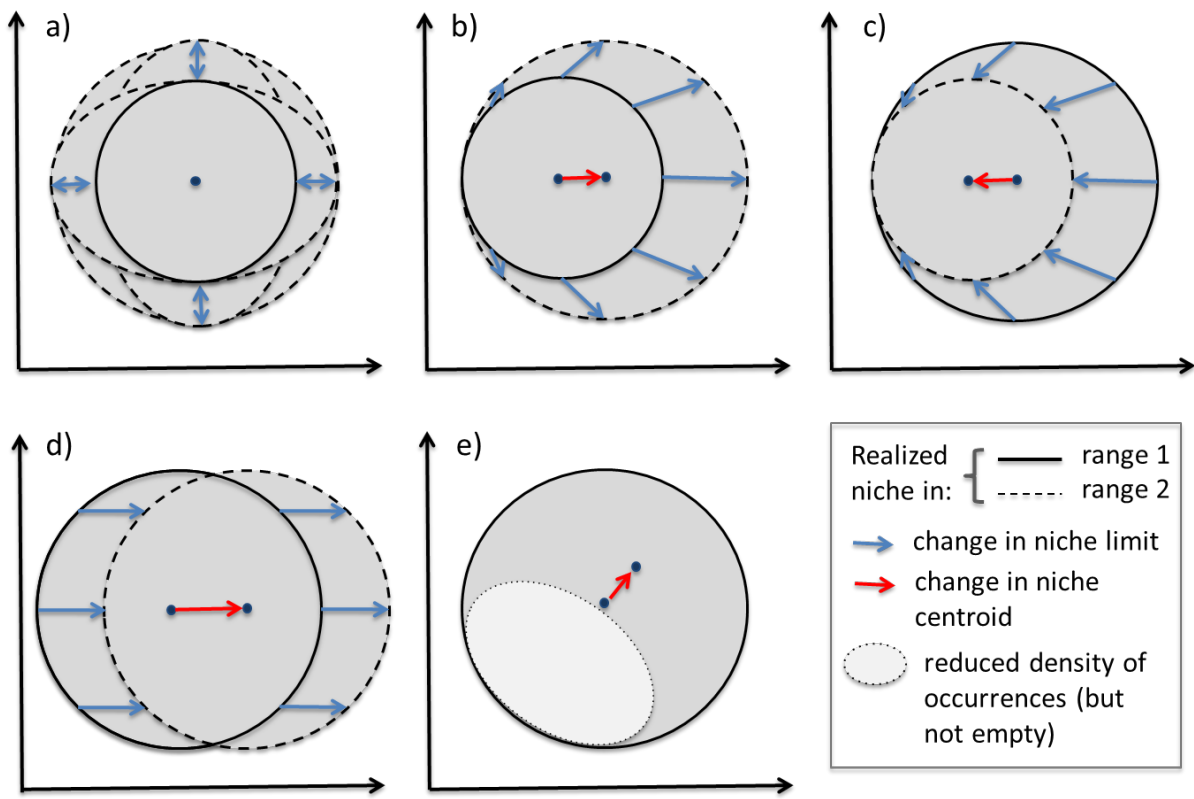
788 **Figure 3:** The two approaches commonly used to quantify niche changes between  
789 ranges (Box 1). Ordination is based on the observations directly, whereas ENM is based  
790 on predictions (see (Broennimann *et al.*, 2012), and Box 1). Steps for ordination are  
791 (square numbers): 1. Definition of the reduced multidimensional environmental space;  
792 2. Plot of the observations from each range in this space; 3. Comparison of the niche  
793 defined from observations in each range; 4. Calculation of the niche change metrics (see  
794 Box 3). Steps for ENMs are: 1. Fit of ENMs by relating field observations to  
795 environmental variables; 2. Projections of the ENMs in geographic space; 3. Compute  
796 difference in the projections; 4. Calculation of the niche change metrics (see Box 3). See  
797 main text for discussion of the respective strengths and weaknesses of the two  
798 approaches.

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802 Figure 1



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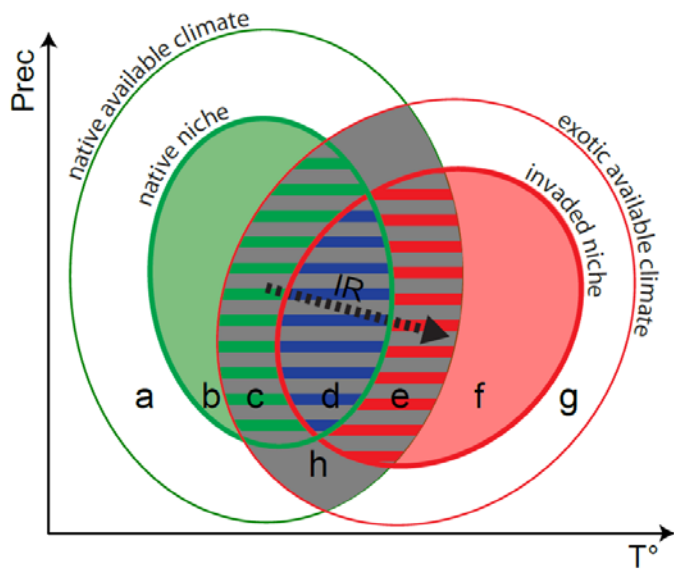
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809 Figure 2



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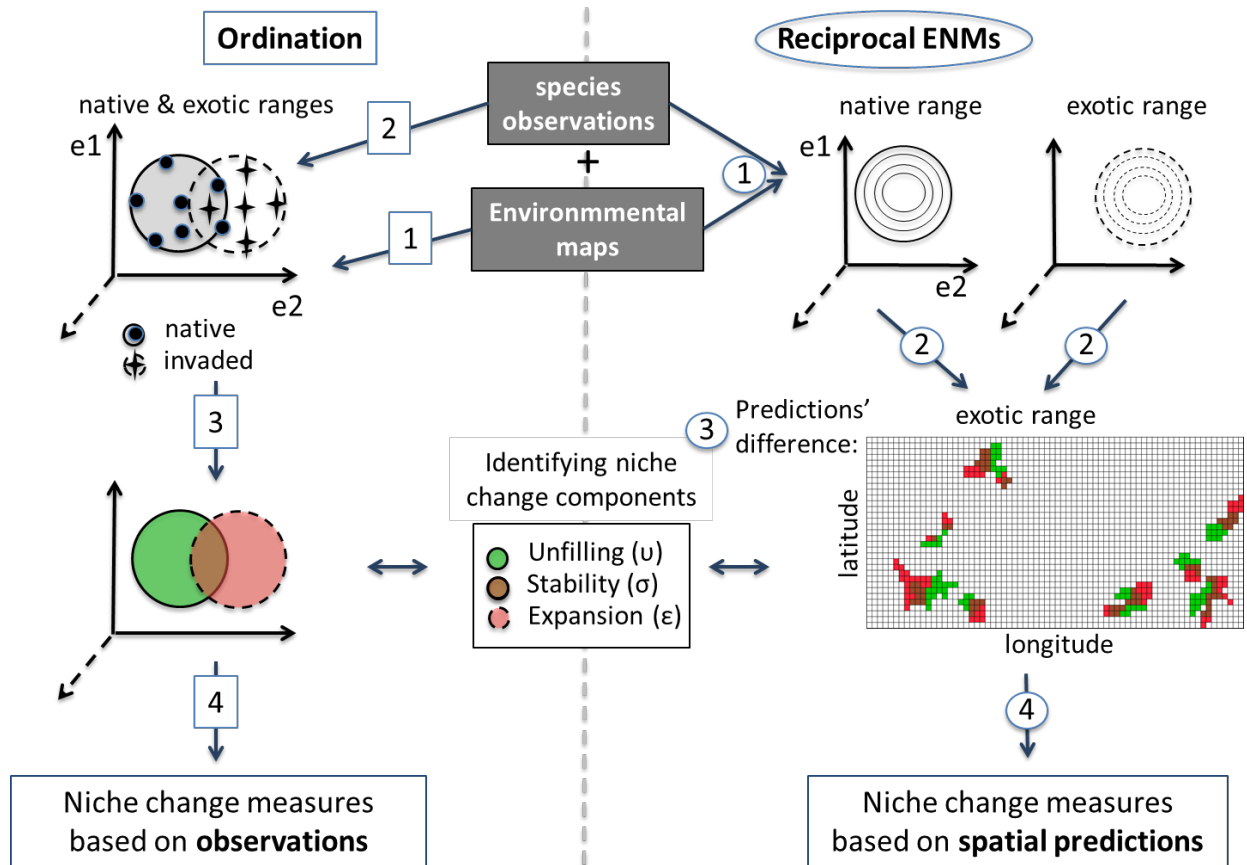
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818 Figure 3



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