

1 **Invasive Species Increase Biodiversity and, Therefore, Services:**

2 **An Argument of Equivocations**

3 **Christopher Hunter Lean**

4 **Abstract**

5 Some critics of invasion biology have argued the invasion of ecosystems by non-indigenous species
6 can create more valuable ecosystems. They consider invaded communities as more valuable because
7 they potentially produce more ecosystem services. To establish that the introduction of non-
8 indigenous species creates more valuable ecosystems they defend that value is provisioned by
9 ecosystem services. These services are derived from ecosystem productivity, the production and
10 cycling of resources. Ecosystem productivity is a result of biodiversity, which is understood as local
11 species richness. Invasive species increase local species richness and, therefore, increase the
12 conservation value of local ecosystems. These views are disseminating to the public via a series of
13 popular science books. Conservationists must respond to these views, and I outline a method of
14 rejecting such arguments against controlling invasive species. Ecological systems are valuable for
15 more than local productivity and biodiversity is not accurately described by a local species count.

16
17 **Keywords:** Invasive Species; Beta-Diversity; Biodiversity Concept; Ecosystem Services; Option
18 **Value; Invasive Species Scepticism**

19
20 **(This paper will be subject to revision and is not the penultimate copy)**

23 **Introduction**

24 It is common practice throughout the world to control invasive species populations to maintain the
25 character and composition of ecological communities. Invasive populations are controlled through the
26 reduction or elimination of their populations and preventing their movement into new areas (Kopf et
27 al. 2017). Scepticism towards the control of invasive species populations has flourished recently with
28 a series of scientists, environmental journalists, and other academics arguing there is rarely reason to
29 control invasive species (Sagoff 2005; Marris 2011; Thompson 2014; Pearce 2015; Thomas 2017).
30 This movement has been described as invasive species denialism, and while there are moments when
31 this literature tips into denialism, there are legitimate arguments that warrant serious consideration
32 (Russell & Blackburn 2017; Frank 2019). In this paper, I draw out and critique an argument that has
33 coalesced within the Invasive Species Sceptics (who I will refer to as *sceptics*) literature. This
34 argument is separate from the standard animal welfare-based arguments that motivate the
35 “compassionate conservation” movement (Wallach et al. 2018) or arguments that concepts such as
36 ‘nativeness’ or ‘invasive species’ are not well-defined or useful (Chew & Hamilton 2011). Instead,
37 the argument addressed here proposes that invasive species are, all things considered, not bad for
38 humanity. I aim to clearly represent the argument, so that scientists may directly address it, and
39 illustrate some possible responses. In my view, the argument pivots on what I consider an illegitimate
40 use of the concept ‘biodiversity’. The argument against the control of invasive species can be found
41 scattered through multiple sources and can be reconstructed as follows:

42 **Why we should not control invasive non-indigenous species:**

- 43 **1.** We should not control populations if they promote ecosystem services (more than any readily
44 available alternative).
- 45 **2.** Invasion often increases *biodiversity*.
- 46 **3.** More *biodiversity* results in more *ecosystem services*.
- 47 **4.** Invasive species often promote *ecosystem services*. (2, 3)

48 **Conclusion:** We should not control invasive species as they often promote ecosystem services (1, 4)
49 This is an extrapolation of a more moderate position, which states that invasive species can contribute
50 to ecosystem services and we should not control a population when these contributions are on sum
51 worth more than the cost of population control (Davis et al. 2011). The above argument generalises
52 the particular, stating on sum we are not warranted acting on invasive species. This implies that
53 research is required to justify preventing the movement of a population into wilderness areas or
54 eradicating a population while it has a small abundance and before it substantially impacts an area.
55 Both positions somewhat utilise the difficulty in conducting cost-benefit analyses of species impacts
56 to support inaction (Courtois et al. 2018). While some may claim this strong view is a fringe argument
57 of a small vocal minority, this is only true internally to the field of conservation science itself. Several
58 of the books that defend this view received wide media attention, particularly *The New Wild* (Pearce
59 2015). These views are disseminating through the public and it is critical to stakeholder engagement
60 for conservationists to respond to these arguments.

61 The idea that invasive species increase biodiversity, and in turn ecosystem services, is unsurprising
62 given the dominant paradigms in conservation ecology, found within the Biodiversity-Ecosystem
63 Services (BES) literature. It has only really been reapplied, with some modifications, to new
64 conclusions by the sceptics (Sagoff 2005; 2018; Pearce 2015; Thomas 2013; 2017) or accepted as an
65 implication of the BES framework (Odenbaugh 2020). This argument can also be converted into a
66 reductio against the BES conservation framework (Newman et al. 2017). In Section 2, I flesh out the
67 argument and situate it in the literature. I critique the argument for deploying impoverished
68 operationalisations of key conservation concepts, biodiversity and ecosystem services. In Section 3, I
69 discuss how ecosystem productivity fails to encompass the range of services proposed within the
70 Ecosystem Services conservation framework. In Section 4, I turn to how local species richness misses
71 many of the critical values the ‘biodiversity’ concept was designed to encompass. Finally, I conclude
72 by conceding some roles Non-Indigenous Species play in contributing to biodiversity (Section 5).

73

74 **2. Unpacking the argument**

75 **2.1. Environment as a service provider**

76 The initial premise, “*(w)e should not control populations if they promote ecosystem services*”, is a
77 corollary of the position that we should preserve species because they provide ecosystem services.
78 Following the Millennium Ecosystem Assessment (MA), which found that ecosystem degradation
79 was a major threat to current and future human wellbeing, ecosystem services have become a major
80 focus of conservation (MA 2005). Ecosystem services are, “the conditions and processes through
81 which natural ecosystems, and the species that make them up, sustain and fulfill human life” (Daily
82 1997, p. 3). More generally, they are considered goods of immediate economic utility. Varying
83 interpretations of “ecosystem services” has led to a literature in which the empirical work, ethical
84 work, and conceptual work do not always lead to the same conclusions about what is worthy of
85 conservation.

86 There is serious debate about what of nature’s value is captured by ecosystem services (Schröter et al.
87 2014). Under some interpretations, anything related to our immediate preferences for nature can be
88 labelled ecosystem services. Conservationists often raise values they believe are not contained within
89 the ecosystem services framework, only to find those within the framework replying that the value
90 raised against them are included (e.g., option value in Faith 2010; Perrings et al. 2010). Sometimes it
91 appears that ecosystem services proponents state a type of value can conceptually be part of the
92 services framework without indicating how the biological features their experiments quantify
93 represent this source of value. A crucial example of this is that many experiments examining the
94 relationship between biodiversity and ecosystem services use biomass production, or net primary
95 production, as a proxy for ecosystem services generally (Newman et al. 2017).

96 Biomass produced is not representative of the range of values people have towards the environment,
97 but it is readily measurable and represents ecosystem productivity. There is a neat conceptual
98 connection between biodiversity, functional diversity, and ecosystem productivity. The thought being
99 biodiverse assemblages will be functionally diverse, providing many ways to process resources, with

100 diverse processing and specialisation the ecosystem will be highly productive. This is appealing as
101 each component is readily quantifiable. Strong proponents of the premise that ecosystem services are
102 the sole justification for conservation can be found within the BES literature as much as within the
103 invasive species sceptic literature, some of whom recognise other types of environmental values (see
104 Marris 2011). For example, Dasgupta et al. (2013) represent biodiversity as only being valuable
105 insofar as it provides ecological functions that can then make productive ecosystems.

106 The representation of biodiversity as only being justified through its relationship to the production or
107 cycling of resources diminishes the variety of values associated with biodiversity. Sometimes
108 biodiversity is represented as either being valuable as it is a cause of services or it has *intrinsic value*,
109 which is notoriously difficult to quantify and whose existence is contested (e.g. Reyer et al. 2012).
110 This all creates the perception, whether justified or not, that biodiversity only derives value from its
111 provision of a narrow set of services, usually equated with resource production and cycling. This
112 underemphasises the cultural, regulating, and supporting services ecosystems provide. It is this narrow
113 interpretation of the relationship between biodiversity and ecosystem services, or more commonly the
114 accidental use of language which represents this relationship as narrow, which warrants the
115 conclusion we should not control invasive species.

116

117 **2.2. Invasive Species increase Biodiversity**

118 Despite many invasive species causing local extinctions, their addition to new ecosystems does not
119 necessarily lead to drastic species loss. There is strong evidence that local species richness worldwide
120 has recently either remained stable or increased (Sax & Gaines 2003; Dornelas et al. 2014). Invasive
121 species can increase the number of species locally; as Pearce (2015 p. 9) says “Rather than reducing
122 biodiversity, the novel new worlds that result [from invasives] are usually richer in species than what
123 went before”. Local species numbers generally appear to be a product of the regional pool of species
124 (Ricklefs 1987). With global connectivity increasing (the ‘*New Pangea*’ celebrated by Thomas 2017),
125 so has the ‘regional’ species pool. This has ultimately driven up local species richness.

126 Assessing species richness is not a simple process. Sometimes ecologists exclude non-indigenous
127 species from local species counts, but as Sagoff (2005 p. 229) argues excluding these populations
128 from such counts by stipulation is just dodgy accounting. But contra Sagoff and other critics of
129 invasive species science and management, any semantic argument utilizing species richness without
130 effort to address the complexities of scale will misrepresent the natural patterns of species
131 distributions. Representing species diversity at multiple scales cannot be done with any single
132 equation (Whittaker et al. 2001).

133 Local increase in species richness has been coupled with global species loss (Dirzo & Raven 2003).
134 This phenomenon has been described as ‘the biodiversity paradox’ (Vellend 2017). The explanation
135 for the paradox is evident, if you add many common non-indigenous species to an area but lose
136 endemic or rare native species there will be increasing local species counts and global species loss.
137 Australia (and the world) has lost the desert bandicoot (*Perameles eremiana*) but gained the red fox,
138 cat, black rat, and common pigeon; a triumph!

139 Ultimately, this indicates simply discussing species numbers misses much of the picture in ecological
140 systems. There must be some attempt to address the relationships between populations. Co-evolved
141 populations have interdependencies, which invasive species can disrupt causing cascading extinctions
142 (Simberloff 2013). While such losses can be recouped through introducing more species, the losses
143 are significant for community composition. The species lost are often specialists who are co-adapted
144 to other local species, the populations introduced are often generalists who can utilize a range of
145 resources and live within varied conditions (Clavel et al. 2011). This leads to the global loss of
146 functional diversity as generalist species prosper. The structure of species interactions must be
147 incorporated into any picture of conservation due to how these interdependencies both lead to species
148 loss and structure biodiversity.

149

150 **2.3. Biodiversity Yields Ecosystem Services**

151 The next step in the case against invasive species control is that the increase in local species counts,
152 due to the introduction of non-indigenous species, results in more ecosystem services. The BES
153 research program supports the case for invasive species increasing the value of ecosystems. It is
154 widely believed that biodiversity increases ecosystem functioning, which increases ecosystem
155 services (Loreau et al. 2001, Haines-Young & Potschin 2010, Mace et al. 2012). If non-indigenous
156 species increase biodiversity then they increase the ecosystems services, which facilitate nature's
157 value to humanity. Or as Mark Sagoff states, "If in any scientific (e.g., random) sample of ecosystems
158 introduced organisms generally, overwhelmingly, and typically increase species richness, and if
159 species richness supports desirable ecosystem properties, then one could argue these organisms
160 benefit those systems." (Sagoff 2005 p. 225).

161 The BES research program has predominantly considered the effects of biodiversity as measured in
162 species richness on ecosystems (Hendriks & Duarte 2008). The most studied effect variable of the
163 biodiversity and ecosystem services relationship is the extent to which ecosystems produce biomass
164 (Cardinale et al. 2011). The scales assessed in these experiments are generally local, only occurring
165 over scales up to 100m. Conservation policy likewise is conducted on the scale of hectares (Srivastava
166 & Vellend 2005). The scales considered by the science, and the policy, appear to support the sceptics'
167 conclusions that we should not control populations of invasive species as on local scales they
168 generally increase species richness and, therefore, ecosystem services.

169

170 **3. Ecosystem Services: Problems with Productivity**

171 Even granting the primacy of ecosystem services in conservation policy, these services come with
172 deceptive variations in how tangible and quantifiable they are. The Millennium Ecosystem
173 Assessment identifies four types of service: provisioning (e.g., wood), regulation (e.g., water quality),
174 cultural (e.g., recreation), and supporting (e.g., carbon cycle) (MA 2005). Despite the scope of the
175 services described, the empirical research on such services historically has narrowed its focus to
176 predominantly the relationship between species richness and biomass or net primary production (e.g.,

177 Carpenter et al. 2006; Costanza et al. 2007; Cardinale et al. 2011). Ecosystem productivity
178 undoubtedly influences the different forms of services provided, it is crucial for both the provision of
179 resources and the regulation of resource cycles. But the emphasis on resource production and cycling
180 to the exclusion of other modes by which services are provided, particularly cultural services, stack
181 the deck towards invasive species. One could counter that ecosystem services are more widely
182 measured than biomass, which is true (Costanza 2015). The issue, however, is that services have
183 historically disproportionately used biomass as a proxy (Newman et al. 2017), which allows for this
184 style of argument to be constructed. Echoes of this historical trend can be seen in the modern
185 literature, a recent metanalysis shows that while ecosystem production and ecosystem provisioning of
186 services was measured by 67% and 68% of studies, only 35% measured the cultural services
187 ecosystems provided (Boerema et al. 2017).

188 Invasive species can contribute to services and reduce services, often simultaneously doing both, and
189 empirical research is required to determine to what degree (Boltovskoy et al. 2018). But the relative
190 contribution of species to the productivity of an ecosystem is highly influenced by the sheer
191 abundance of that population (Winfree et al. 2015). This makes ecosystem productivity quite
192 antithetical to conservation's aims of preserving endemic and rare species, which are often not
193 abundant. Many rare, threatened, and endangered species are 'functionally extinct' in that they are not
194 able to have strong effects on the ecosystem they reside within. Within a BES framework, where
195 productivity and direct causal contribution is emphasised, such species lack value. Instead, it is the
196 hyper-abundant and highly productive species that contribute. The features that make invasive species
197 invasive rather than just non-indigenous is their ability to rapidly grow in abundance (Simberloff
198 2013). Their ability to produce biomass is what allows them to physically exclude local species. These
199 properties are given a new presentation by sceptics, their rapid increases in abundance and biomass
200 make them productive ecosystem services providers (Pearce 2015). Invasive species then should be
201 considered as 'super species' due to their success moving across the globe and processing biomass
202 (Hamilton 2010). It is the framing of biodiversity's value as being strongly connected to the
203 productivity of whole ecosystems that leads to these conclusions.

204 Conservationists have warned against strongly connecting conservation to ecological productivity
205 (Silvertown 2015; Faith 2018). Following his reflections on Leopold's land ethic Michael Soulé
206 warned us that justifying conservation through ecosystem processes would facilitate the conclusion
207 we should replace native species with invasives:

208 *"it is technically possible to maintain ecological processes, including a high level of economically*
209 *beneficial productivity, by replacing the hundreds of native plants, invertebrates and vertebrates with*
210 *about 15 or 20 introduced, weedy species.... WARNING! Be suspicious of "ecologists" who are*
211 *pitching ecological services (for people) and who speak of "redundant" species or "hyperdiversity."*"

212 Soulé 1996 (p. 60)

213 In the face of such warnings we now find, two decades on, significant support for the idea invasive
214 species are 'super species', which can replace natives due to their productivity (e.g. Pearce 2015).

215

216 **4. Biodiversity**

217 **4.1. Biodiversity is more than Species Richness**

218 In the case of invasive species being added to the local species pool, biodiversity is increased under
219 the assumption that biodiversity is local species richness (Pearce 2015; Thomas 2017). These critics
220 expect this increase to outpace local species extinctions. Local species count, or species richness, is
221 widely known as α (Alpha) diversity. When the local extinctions are of species endemic to that region,
222 global species counts reduce. This global inventory of species is γ (Gamma) diversity, or more
223 accurately the inventory of all the local systems being analysed. These two diversity measures take an
224 inventory of the populations or species or similar unit of biodiversity in their region. There is another
225 count, which is widely considered an essential target in conservation. This is β (Beta) diversity, which
226 is a comparative measure of diversity between regions. It considers how many new species are added
227 to the regional species pool by an area. By taking biodiversity as only α diversity, sceptics
228 significantly underplay the damage Non-Indigenous Species do through their diminishing of γ
229 diversity and β diversity.

230 β diversity is a measure of the entities which comprise biodiversity, biodiversity units; these are
231 generally counted as species but can be other entities (Sarkar 2016). For example, the entities being
232 counted could be the distinct habitat types in an area, like shrublands or deciduous tree forest, or
233 biotic 'features', which are the biotic traits possessed by populations such as their genes or their
234 'functions'. Further dimensions of biodiversity could be argued for such as diversity of biotic
235 interactions (Luna et al. 2020). These can be understood as compromising different levels of
236 biodiversity and we may have reason to count all or some (Faith 2016; Lean & Sterelny 2016). A
237 local ecosystem will have higher β diversity the more unique biodiversity units it adds to the
238 previously assessed regional pools, the 'complementary' units of diversity (See Figure 1). If there are
239 no previously assessed areas, then we are making a count of biodiversity units in an area, which is
240 equivalent to α diversity.

241 **Insert Figure 1.**

242 Adding new species to those already protected increase β diversity but species are not equally similar.
243 Many species are extremely similar (e.g., cryptic species). Complementarity has been incorporated
244 into algorithms to identify species that are themselves unique (Vane-Wright et al. 1991; Faith 1992).
245 The disparity between species can be represented through measuring phylogenetic distance or the
246 functional differentiation between populations (see Magurran & McGill 2011). There are continuing
247 debates on which measures best represent biological difference but incorporating the extent to which
248 populations themselves contribute unique features is an extension of complementarity and
249 biodiversity measurement (Lean & Maclaurin 2016; Lean 2017).

250 β diversity is generally thought of as an essential component of biodiversity preservation practice
251 (Sarkar 2012; 2016; Socolar et al. 2016). This is partially due to a conceptual claim, biodiversity as a
252 concept is designed to maximize the representation of different life forms. Regardless of the entities
253 measured as representing biodiversity, higher β diversity results in more biotic variety, therefore,
254 should be incorporated into conservation decision-making (Sarkar 2006). Complementarity already
255 has featured in the practice of conservation planning for 40 years to select areas that represent the
256 most distinct lifeforms (Kirkpatrick et al. 1980). It is both part of the practice of conservation and part

257 of the theoretical framework of biodiversity conservation. Insofar as biodiversity aims to represent
258 more than just a tally it must quantify unique entities.

259

260 **4. 2. Valuing Biodiversity beyond Species Richness**

261 The values represented through β and γ diversity are not easily captured within the α diversity focused
262 BES framework. Local α diversity is required to understand the goods local interacting populations
263 produce, but β diversity represents more abstract values. β diverse ecosystems have value over copies
264 of common ecosystem types, their uniqueness connects them to the overall range of forms found in
265 life on earth (γ diversity). Local ecosystem productivity is irrelevant to the value created by these
266 forms of diversity and vice versa. Local tallies of biological entities cannot represent the full range of
267 biological values as they ignore how the preservation of a range of unique variety is valuable.

268 Ecosystem services are not the only or original justification for preserving biodiversity. Biodiversity
269 was designed to represent the range of biological features that exist (Soulé 1985; Wilson 1992)
270 including key values overlooked in the search for productivity: heritage and option value. These
271 values are not derived from immediate use and may be difficult to represent economically (Silvertown
272 2015).

273 **Option Value:** Biodiversity is the most direct way to preserve option value. The preservation of a
274 range of biological features is a prudent bet-hedging strategy to account for future uncertainty (Faith
275 1992; Maclaurin and Sterelny 2008; Lean 2017; Owen et al. 2019; c.f. Maier 2012; Newman et al.
276 2017). The utility of diverse features of life cannot be accurately known. These values need not only
277 be in their use for commerce or medicine (future monetization). Human preferences may change in
278 their representation of what they find aesthetically appealing or culturally significant. Given that the
279 losses of biological features are irreversible, we need to guard against the risk involved in losing these
280 goods (Arrow & Fisher 1974). This is true even if in rare cases some of these goods have some
281 unexpected disutility (like zoonotic diseases), as long as the sum benefit of preserving diversity is
282 advantageous.

283 **Heritage Value:** Heritage value is commonly derived from an entity having cultural significance to a
284 group of people, usually developed over extended periods (Thompson 2000). Just as old buildings or
285 artworks have both an intellectual value, in that they are a record of history and culture, and are of
286 aesthetic value, often because they are a physical representation of the past, so too does biodiversity
287 (Russow 1981; Sober 1986). This creates a relationship between local people and the history of
288 environmental systems. While cultural significance is mentioned in the wider ecosystem service
289 framework, a focus on productivity ignores these values.

290 These values are more difficult to quantify within the ecosystem services framework but they are still
291 instrumental-anthropocentric values. A sophisticated ecosystem services framework could incorporate
292 them, but when such a framework is skewed towards ecosystem productivity and local species counts,
293 they are undervalued.

294

295 **4.3. Valuing Diversity**

296 Invasive species should be controlled because they diminish β diversity homogenizing the biological
297 world (Wright 2011). Uniqueness and diversity foster connections between local citizens and their
298 natural landscape, which can be lost through it being just like any other place in the world. This
299 grounds people's local pride in these systems and justifies their disdain for homogenisation. Heritage
300 value is created by local people interacting with their local ecological systems over time. Value is
301 created by the acknowledgement of unique experiences formed by having a relationship to a unique
302 environment. This can be described as a relational intrinsic value or as an instrumental value (Elliot
303 1992). Heritage and uniqueness increase ecosystem desirability to not just local people but also
304 tourists. There is no reason for me to travel to California to walk through Gum forests. The Gum
305 forests around Sydney provide the same aesthetic experience but also possess heritage value derived
306 from their historical relationship to this place and the other species within the Australian landscape.
307 This provides the Sydney Gum forest with a comparative advantage in its conservation value over the
308 California Gum forest. The cultural services provided by ecosystems are often recognised by

309 ecosystem services in studies (Boerema et al. 2017) but are not represented by the BES relationship
310 built from local species counts.

311 Global species richness, γ diversity, is of unique heritage value (Wilson 1992). Not only does it
312 provide local people with a unique sense of place in the world, but unique biotic forms carry
313 information about the past. Global species diversity is seen as an object of global heritage, comparable
314 to human sites like the pyramids of Giza or Stonehenge. Some are sceptical of invoking global
315 heritage, as its protection can take the form of colonialism and as such cannot be ethically enforced
316 (Sarkar 2019). While we can accept that acting on global heritage claims at times can be unethical, we
317 may still hold that such entities are of global value, and as local conservation actors, we should
318 maintain this value. Preserving global species richness is the archetypal commitment of
319 environmentalism. The founding of the International Union for Conservation of Nature and its Red
320 List was created with the goal of stopping extinctions (IUCN 2020) and The United Nations
321 Educational, Scientific, and Cultural Organization's (UNESCO) World Heritage List was created to
322 preserve sites of heritage value be they natural or man-made (UNESCO 2021). While conservationists
323 may accept that we cannot save all species, due to resource limitations, it does not imply global
324 species preservation is not a goal of conservation. Advocating for allowing 'relic' or 'loser' species to
325 become extinct stands in contrast to such aims (Pearce 2015; Thomas 2017). To claim that global
326 species loss is secondary to the primary conservation goal of resource production is to reject the
327 foundations of conservation biology.

328 The emphasis on local diversity and acceptance of global extinction, proposed by sceptics, stands as a
329 radical rejection of the principles traditionally associated with conservation. Consider the original
330 postulates of conservation described by Soulé (1985): (1) diversity should be preserved, (2) untimely
331 extinctions should be prevented, (3) ecological complexity should be maintained, (4) evolutionary
332 processes should continue, and (5) biological diversity has intrinsic value. Interpreting these
333 postulates as claims about global or local diversity results in different recommendations. By solely
334 interpreting diversity locally rather than globally, sceptics are proposing we, at the minimum, jettison

335 1, 2, and 5 as global conservation aims. They must defend such a radical change in conservation
336 values.

337 Invasive species actively diminish β diversity when they eliminate native endemic biotic variation and
338 replace them with biotic forms that are found commonly elsewhere. This not only diminishes heritage
339 value but also option value. Option value directly connects to β diversity, as unique features create
340 new options. Option value does not require large standing populations of high productivity species,
341 just preserving unique lifeforms because we may value them in unique and unpredictable ways in the
342 future.

343 Preserving diverse biotic features directly entails the preservation of unique options, it is just a
344 question of what the best way is to measure diversity to represent the unknown future uses of life on
345 earth (Lean 2017). Attempts to reduce option value to functional diversity (e.g. Mazel et al. 2018)
346 systematically underestimates the value of biotic diversity because they ignore the way human
347 preferences for the environment change over time, often in unexpected ways. While ‘swamps’ were
348 not valued highly in yesteryear, many highly value ‘wetlands’. Option value indicates we should
349 preserve the environment for changing recreational and aesthetic valuations in addition to its possible
350 immediate economic uses. There is a range of values that people, when surveyed, hold towards the
351 environment that are not captured by productivity (see the literature on Wildlife Value Orientations
352 e.g. Fulton et al. 1996). These values change between demographics and over time. Option value is
353 for preserving biodiversity so other humans can value different aspects of the environment in the
354 future.

355 There are numerous ways to describe the value that biodiversity provides. Local species richness is
356 inadequate. Adding rats, cats, and pigeons to every corner of the globe does not preserve the heritage
357 or options value of an area. Possessing unique biotic resources allows communities to bargain with
358 other communities and fosters their connection to the local environment. These values require
359 representing the range of lifeforms that exist across different ecosystems through γ biodiversity and β
360 diversity. These necessary components of biodiversity preservation are ignored when we solely focus
361 on ecosystem productivity.

362 Now one could argue that this dispute is about differing values rather than equivocation. It is, in one
363 sense. The critics of invasive species management ascent to a much narrower conception of
364 conservations goals than most conservationists have traditionally considered. Only describing
365 biodiversity as α diversity, rather than admitting the importance of β and γ diversity, and representing
366 services as being derived from high productivity and fecundity. They could argue that local species
367 richness is more significant than both heritage and option value. In partial agreement with these
368 critics, some have argued the ecosystem services paradigm justifies not preserving a large portion of
369 biodiversity (Newman 2020). But invasive species critics, however, do not provide strong arguments
370 for such a narrowing of the scope of conservation goals. Instead they use general terms (biodiversity,
371 ecosystem services) to appear to be agreeing to the more widely held views about conservations aims.
372 This appears to be a rhetorical decision to equivocate for the means of engagement with conservations
373 aims. What is required of such critics is a direct argument we should narrow the goals of conservation
374 for there to be an honest debate about values in conservation. This would then facilitate the further
375 assessment of the costs and benefits of preferring such a narrow interpretation over the wider goal's
376 conservation has traditionally held.

377

378 **5. Conclusion: Beta Diversity and Invasion**

379 Accepting that biodiversity must represent uniqueness and disparity does not imply we must always
380 control Non-Indigenous Species in wild spaces. There are a significant number of species that are
381 endangered or extinct in their native habitat but wild in an invasive habitat. Thompson (2014) frames
382 his discussion of invasive species control around the case of the Camel. Wild Camel populations no
383 longer exist in their native range, but wild Camel populations move through central Australia. If we
384 remove this population, we reduce the β diversity of this habitat and the number of wild populations
385 on earth. Accepting β diversity as a significant biodiversity measure indicates we should retain Camel
386 populations in Australia. This is, however, not without conditions. If an invasive population threatens
387 multiple endemic native populations, it will warrant the control or even eradication of this population.
388 Population control is critical for populations without consumers. Population control does not imply

389 local extinction and often the best choice is to keep the population numbers low enough so that they
390 do not impact indigenous populations.

391 The β diversity conservation framework does not necessitate invasive species control in all cases. The
392 number of species that are endangered in their native range and invasive are increasing and include
393 the wattle-necked soft-shell turtle, the Monterey pine, and the Barbary Sheep (Marchetti & Engstrom
394 2016). There will be instances where non-indigenous species have moved into a system and now
395 provide services necessary for the survival of endemic species. Chew (2009) argues Tamarisk in the
396 USA is a critical habitat for native songbirds. In such cases, consideration should be given to these
397 populations and the role they play in supporting biotic diversity and uniqueness. This does not,
398 however, warrant the rejection of invasive species control and eradication.

399 Current arguments forwarded by sceptics of invasive species control engage environmentalists on
400 their own principles rather than solely forwarding animal welfare arguments. They contest that on the
401 grounds of preserving biodiversity and promoting ecosystem services the control of invasive species
402 is not justified. Their arguments, however, require an impoverished account of biodiversity, one
403 which equates local species counts with biodiversity. This position ignores the importance of diversity
404 and the disparity of life. It ignores the value of unique biotic options, and the potential utility these
405 options could bring, and it ignores the heritage contained in life on Earth. Such values justify the
406 preservation of endemic and unique species even when they are not major contributors to local
407 productivity.

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415 **References**

- 416 Arrow, K. J., & Fisher, A. C. (1974). Environmental Preservation, Uncertainty, and
417 Irreversibility. *The Quarterly Journal of Economics*, 88(2), 312-319.
- 418 Boerema, A., Rebelo, A. J., Bodi, M. B., Esler, K. J., & Meire, P. (2017). Are ecosystem services
419 adequately quantified?. *Journal of Applied Ecology*, 54(2), 358-370.
- 420 Boltovskoy, D., Sylvester, F., & Paolucci, E. M. (2018). Invasive species denialism: sorting out facts,
421 beliefs, and definitions. *Ecology and evolution*, b8(22), 11190-11198.
- 422 Cardinale, B. J., Matulich, K. L., Hooper, D. U., Byrnes, J. E., Duffy, E., Gamfeldt, L., ... &
423 Gonzalez, A. (2011). The functional role of producer diversity in ecosystems. *American*
424 *journal of botany*, 98(3), 572-592.
- 425 Carpenter, S. R., DeFries, R., Dietz, T., Mooney, H. A., Polasky, S., Reid, W. V., & Scholes, R. J.
426 (2006). Millennium ecosystem assessment: research needs. *Science* 314: 257-258
- 427 Chew, M. K. (2009). The monsterring of tamarisk: how scientists made a plant into a problem. *Journal*
428 *of the History of Biology*, 42(2), 231-266.
- 429 Chew, M.K. and Hamilton, A.L. (2011) The rise and fall of biotic nativeness: a historical perspective.
430 In Richardson, D.M. (Eds) *Fifty Years of Invasion Ecology. The Legacy of Charles Elton*.
431 Blackwell, Oxford, 35–48.
- 432 Clavel, J., Julliard, R., & Devictor, V. (2011). Worldwide decline of specialist species: toward a
433 global functional homogenization?. *Frontiers in Ecology and the Environment*, 9(4), 222-228.
- 434 Courtois, P., Figuieres, C., Mulier, C., & Weill, J. (2018). A cost–benefit approach for prioritizing
435 invasive species. *Ecological Economics*, 146, 607-620.
- 436 Costanza, R., Fisher, B., Mulder, K., Liu, S., & Christopher, T. (2007). Biodiversity and ecosystem
437 services: A multi-scale empirical study of the relationship between species richness and net
438 primary production. *Ecological economics*, 61(2-3), 478-491.
- 439 Costanza R. (2015). Ecosystem services in theory and practice. In P Figgs, B Mackey, J Fitzsimons, J
440 Irving, P Clark (Eds) *Valuing nature: Protected areas and ecosystem services*. Australian
441 Committee for IUCN: Sydney, 6–15

442 Daily, G. C. (1997). *Nature's services*. Island Press: Washington, DC.

443 Dasgupta, P., Kinzig, A. P., & Perrings, C. (2013). The value of biodiversity. In: S. Levin, G.C. Daily
444 and R.K. Colwell et al., (Eds), *The Encyclopedia of Biodiversity*, Academic Press, San Diego,
445 CA

446 Davis, M. A., Chew, M. K., Hobbs, R. J., Lugo, A. E., Ewel, J. J., Vermeij, G. J., ... Carroll, S. P.
447 (2011). Don't judge species on their origins. *Nature*, 474(7350), 153.

448 Dirzo, R., & Raven, P. H. (2003). Global state of biodiversity and loss. *Annual review of Environment
449 and Resources*, 28.

450 Dornelas, M., Gotelli, N. J., McGill, B., Shimadzu, H., Moyes, F., Sievers, C., & Magurran, A. E.
451 (2014). Assemblage time series reveal biodiversity change but not systematic
452 loss. *Science*, 344(6181), 296-299.

453 Elliot, R. (1992). Intrinsic value, environmental obligation and naturalness. *The Monist*, 75(2), 138-
454 160.

455 Faith, D. P. (2010). Biodiversity transcends services. *Science*, 330(6012), 1744-1745.

456 Faith, D. P. (1992). Conservation evaluation and phylogenetic diversity. *Biological
457 Conservation*, 61(1), 1-10.

458 Faith, D. P. (2016). A General Model for Biodiversity and its Value. In J. Garson, A.
459 Plutynski, & S. Sarkar (Eds.), *The Routledge Handbook of Philosophy of Biodiversity*.
460 New York: Routledge, 69 – 85.

461 Faith, D. P. (2018). Avoiding paradigm drifts in IPBES: reconciling “nature’s contributions to
462 people,” biodiversity, and ecosystem services. *Ecology and Society*, 23(2).

463 Fulton, D. C., Manfredi, M. J., & Lipscomb, J. (1996). Wildlife value orientations: A conceptual and
464 measurement approach. *Human dimensions of wildlife*, 1(2), 24-47.

465 Frank, D. M. (2019). Disagreement or denialism? “Invasive species denialism” and ethical
466 disagreement in science. *Synthese*, 1-29.

467 Haines-Young, R., & Potschin, M. (2010). The links between biodiversity, ecosystem services and
468 human well-being. In D. Raffaelli & C. Frid (Eds.) *Ecosystem Ecology: a new synthesis*.
469 Cambridge, 110-139.

470 Hamilton, G. (2010). *Super Species: The Creatures That Will Dominate the Planet*. Richmond
471 Hill: Firefly.

472 Hendriks, I. E., & Duarte, C. M. (2008). Allocation of effort and imbalances in biodiversity
473 research. *Journal of Experimental Marine Biology and Ecology*, 360(1), 15-20.

474 IUCN (2020). *The IUCN Red List of Threatened Species*. Version 2020-2. URL
475 <https://www.iucnredlist.org>. Accessed 10.10.2020.

476 Kirkpatrick, J. B., Brown, M. J., & A. Moscal. (1980). *Threatened plants of the Tasmanian central*
477 *east coast*. Tasmanian Conservation Trust.

478 Kopf, R. K., Nimmo, D. G., Humphries, P., Baumgartner, L. J., Bode, M., Bond, N. R., ... & Olden, J.
479 D. (2017). Confronting the risks of large-scale invasive species control. *Nature Ecology &*
480 *Evolution*, 1(6), 1-4.

481 Lean, C., & Maclaurin, J. (2016). The Value of Phylogenetic Diversity. In R. Pellens & P.
482 Grandcolas (Eds.), *Biodiversity Conservation and Phylogenetic Systematics* Springer, 19–37.

483 Lean, C., & Sterelny, K. (2016). Ecological hierarchy and biodiversity. In J. Garson, A. Plutynski, &
484 S. Sarkar (Eds.), *The Routledge handbook of biodiversity*. London: Routledge, 110-124.

485 Lean, C. H. (2017). Biodiversity Realism: Preserving the tree of life. *Biology & Philosophy*, 32(6),
486 1083–1103.

487 Loreau, M., Naeem, S., Inchausti, P., Bengtsson, J., Grime, J. P., Hector, A., ... & Tilman, D. (2001).
488 Biodiversity and ecosystem functioning: current knowledge and future
489 challenges. *Science*, 294(5543), 804-808.

490 Luna, P., Corro, E. J., Antoniazzi, R., & Dáttilo, W. (2020). Measuring and linking the missing part of
491 biodiversity and ecosystem function: The diversity of biotic interactions. *Diversity*, 12(3), 86.

492 Mace, G. M., Norris, K., & Fitter, A. H. (2012). Biodiversity and ecosystem services: a multilayered
493 relationship. *Trends in ecology & evolution*, 27(1), 19-26.

494 Maclaurin, J., & Sterelny, K. (2008). *What is biodiversity?* Chicago: University of Chicago Press.

495 Magurran, A. E., & McGill, B. J. (2011). *Biological diversity: frontiers in measurement and*
496 *assessment*. Oxford: Oxford University Press.

497 Maier, D. S. (2012). *What's so good about biodiversity? A call for better reasoning about*

498 *nature's value*. Dordrecht: Springer.

499 Marchetti, M. P., & Engstrom, T. (2016). The conservation paradox of endangered and invasive
500 species. *Conservation Biology*, 30(2), 434-437.

501 Marris, E. (2011). *Rambunctious garden: saving nature in a post-wild world*. Bloomsbury Publishing
502 USA.

503 Mazel, F., Pennell, M. W., Cadotte, M. W., Diaz, S., Dalla Riva, G. V., Grenyer, R., ... & Pearse, W.
504 D. (2018). Prioritizing phylogenetic diversity captures functional diversity unreliably. *Nature*
505 *Communications*, 9(1), 1-9.

506 *Millennium Ecosystem Assessment*. Washington, DC, Island Press.

507 Newman, J. A., Varner, G., & Linquist, S. (2017). *Defending biodiversity: environmental science and*
508 *ethics*. Cambridge University Press.

509 Newman, J. A. (2020). Biodiversity, ecosystem functioning, and the environmentalist agenda: a reply
510 to Odenbaugh. *Biology & Philosophy*, 35(1), 1-8.

511 Odenbaugh, J. (2020). Biodiversity, ecosystem functioning, and the environmentalist agenda. *Biology*
512 *& Philosophy*, 35(1), 16.

513 Owen, N. R., Gumbs, R., Gray, C. L., & Faith, D. P. (2019). Global conservation of phylogenetic
514 diversity captures more than just functional diversity. *Nature communications*, 10(1), 1-3.

515 Pearce, F. (2015). *The new wild: Why invasive species will be nature's salvation*. Icon Books Ltd.

516 Perrings, C., Naeem, S., Ahrestani, F., Bunker, D. E., Burkill, P., Canziani, G., ... & Kawabata, Z.
517 (2010). Biodiversity Transcends Services—Response. *Science*, 330(6012), 1745-1745.

518 Sagoff, M. (2005). Do non-native species threaten the natural environment? *Journal of Agricultural*
519 *and Environmental Ethics*, 18(3), 215–236.

520 Sagoff, M. (2018). Invasive species denialism: a reply to Ricciardi and Ryan. *Biological*
521 *Invasions*, 20(10), 2723-2729.

522 Reyers, B., Polasky, S., Tallis, H., Mooney, H. A., & Larigauderie, A. (2012). Finding common
523 ground for biodiversity and ecosystem services. *BioScience*, 62(5), 503-507.

524 Ricklefs, R. E. (1987). Community diversity: relative roles of local and regional
525 processes. *Science*, 235(4785), 167-171.

- 526 Russell, J. C., & Blackburn, T. M. (2017). The rise of invasive species denialism. *Trends in Ecology*
527 *& Evolution*, 32(1), 3-6.
- 528 Russow, L. M. (1981). Why do species matter?. *Environmental Ethics*, 3(2), 101-112.
- 529 Sarkar, S. (2006). Ecological diversity and biodiversity as concepts for conservation planning:
530 comments on Ricotta. *Acta Biotheoretica*, 54(2), 133-140.
- 531 Sarkar, S. (2012). *Environmental philosophy: from theory to practice*. Sussex: John Wiley &
532 Sons.
- 533 Sarkar, S. (2016). Approaches to Biodiversity. In J. Garson, A. Plutynski, & S. Sarkar (Eds.) *The*
534 *Routledge Handbook of Philosophy of Biodiversity*. New York: Routledge, 375-399.
- 535 Sarkar, S. (2019). What Should “Biodiversity” Be? In E. Casetta, J. Marques da Silva, & D. Vecchia
536 (Eds.) *From Assessing to Conserving Biodiversity: Conceptual and Practical Challenges*.
537 Switzerland: Springer, 401–414.
- 538 Sax, D. F., & S. D. Gaines. (2003). Species diversity: From global decreases to local increases. *Trends*
539 *in Ecology & Evolution* 18:561–566
- 540 Schröter, M., Van der Zanden, E. H., van Oudenhoven, A. P., Remme, R. P., Serna-Chavez, H. M.,
541 De Groot, R. S., & Opdam, P. (2014). Ecosystem services as a contested concept: a synthesis
542 of critique and counter-arguments. *Conservation Letters*, 7(6), 514-523.
- 543 Silvertown, J. (2015). Have ecosystem services been oversold?. *Trends in ecology &*
544 *evolution*, 30(11), 641-648.
- 545 Simberloff, D. (2013). *Invasive species: what everyone needs to know*. Oxford University Press.
- 546 Sober, E. (1986). 'Philosophical problems for environmentalism'. In B. Norton (Eds.) *The*
547 *Preservation of Species*. Princeton: Princeton University Press, 173-94.
- 548 Socolar, J. B., Gilroy, J. J., Kunin, W. E., & Edwards, D. P. (2016). How should beta-diversity inform
549 biodiversity conservation?. *Trends in ecology & evolution*, 31(1), 67-80.
- 550 Soulé, M. E. (1985). What Is Conservation Biology? *BioScience*, 35(11), 727–734.
- 551 Soulé, M. E. (1996). “Are ecosystem processes enough?” *Wild Earth* 6(1): 59–60.
- 552 Srivastava, D. S., & Vellend, M. (2005). Biodiversity-ecosystem function research: is it relevant to
553 conservation?. *Annu. Rev. Ecol. Evol. Syst.*, 36, 267-294.

554 Thomas, C. D. (2013). The Anthropocene could raise biological diversity. *Nature*, 502(7469), 7-7.

555 Thomas, C. D. (2017). *Inheritors of the Earth: how nature is thriving in an age of extinction*. Hachette
556 UK.

557 Thompson, J. (2000). Environment as Heritage. *Environmental Ethics*, 22(3), 241-258.

558 Thompson, K. (2014) *Where do camels belong? the story and science of invasive species*. Profile
559 Books, London.

560 UNESCO (2021). *World Heritage Convention*. URL <https://whc.unesco.org/en/convention/>. Accessed
561 03.08.2021.

562 Vane-Wright, R. I., Humphries, C. J., & Williams, P. H. (1991). What to protect?—
563 Systematics and the agony of choice. *Biological Conservation*, 55(3), 235–254

564 Vellend, M. (2017). The biodiversity conservation paradox. *American Scientist*, 105(2), 94-101.

565 Wallach, A. D., Bekoff, M., Batavia, C., Nelson, M. P., & Ramp, D. (2018). Summoning compassion
566 to address the challenges of conservation. *Conservation Biology*, 32(6), 1255-1265.

567 Wilson, E. O. (1992). *The diversity of life*, Cambridge: Belknap Press.

568 Whittaker, R. J., Willis, K. J. & Field. R. (2001). "Scale and species richness: towards a general,
569 hierarchical theory of species diversity." *Journal of biogeography* 28.4: 453-470.

570 Winfree, R., W. Fox, J., Williams, N. M., Reilly, J. R., & Cariveau, D. P. (2015). Abundance of
571 common species, not species richness, drives delivery of a real-world ecosystem
572 service. *Ecology letters*, 18(7), 626-635.

573 Wright, S. (2011). Invasive species and the loss of beta diversity. *Ethics & the Environment*, 16(1),
574 75-97.

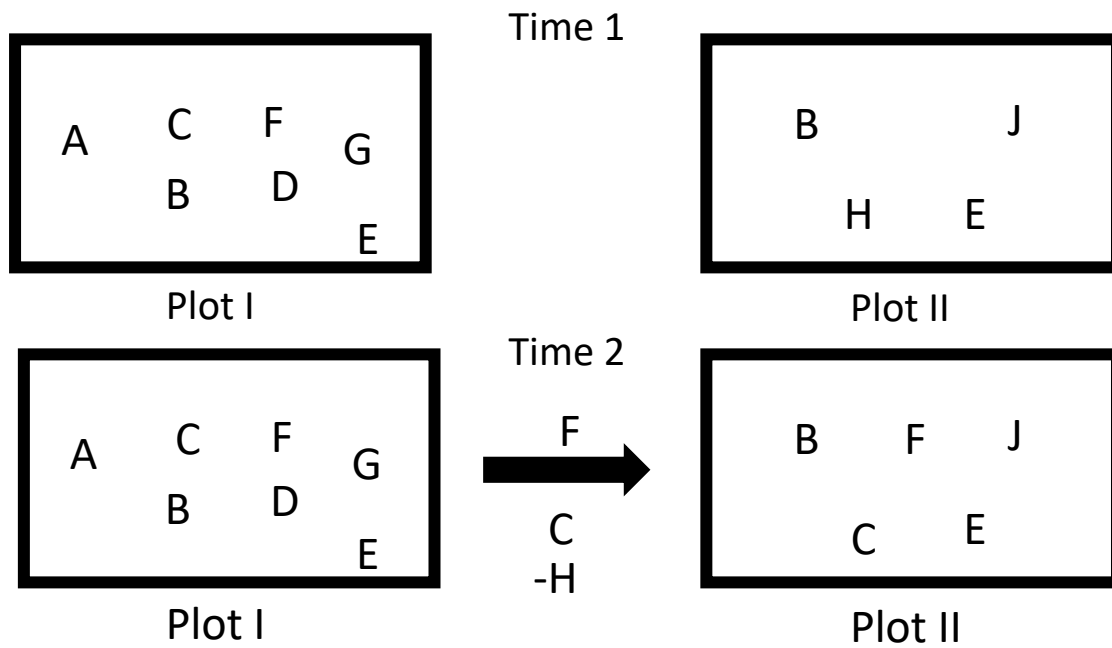


Figure 1. Plots, I, which contains 7 species, and II, which has 4 (α diversity). II adds two unique species to I (β diversity). Their combined species count is 9 (γ diversity). If through introduction, two of I's species (F,C) invade II, and one of II's unique species (H) is eradicated then II increases its α diversity by 1 but its β diversity is reduced by 1.