

# A framework to quantify the vulnerability of insular biota to global change

Céline Bellard, Clara Marino, Nathalie Butt, José María Fernández-Palacios, François Rigal, Marine Robuchon, Jonathan Lenoir, Severin Irl, Ana Benítez-López, Pol Capdevila, et al.

#### ▶ To cite this version:

Céline Bellard, Clara Marino, Nathalie Butt, José María Fernández-Palacios, François Rigal, et al.. A framework to quantify the vulnerability of insular biota to global change. 2024. hal-04550966v1

# HAL Id: hal-04550966 https://hal.science/hal-04550966v1

Preprint submitted on 18 Apr 2024 (v1), last revised 14 Oct 2024 (v2)

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

#### A framework to quantify the vulnerability of insular biota to global change 1

- C. Bellard<sup>1\*</sup>, C. Marino<sup>1,2</sup>, N. Butt<sup>3</sup>, José M. Fernández-Palacios<sup>4</sup>, F. Rigal<sup>5</sup>, M. Robuchon<sup>6</sup>, J. 2
- 3
- Lenoir<sup>7</sup>, Severin D. H. Irl<sup>8</sup>, A. Benítez-López<sup>9</sup>, P. Capdevila<sup>10,11</sup>, G. Zhu<sup>12</sup>, Gabriel H. O. Caetano<sup>1</sup>, P. Denelle<sup>13</sup>, M. Philippe-Lesaffre<sup>1</sup>, A.M. Schipper<sup>14</sup>, W.B. Foden<sup>15,16</sup>, W. Daniel Kissling<sup>17</sup>, C. Leclerc<sup>18</sup> 4
- 5
- \*Email: celine.bellard@universite-paris-saclay.fr 6

7

- <sup>1</sup> Université Paris-Saclay, CNRS, AgroParisTech, Ecologie Systématique et Evolution, 91190, 8
- 9 Gif-sur-Yvette, France
- <sup>2</sup> FRB-Cesab, 5 rue de l'école de médecine, 34000 Montpellier, France 10
- <sup>3</sup> School of the Environment, The University of Queensland, St. Lucia, 4072, Queensland, 11
- 12 Australia.
- <sup>4</sup> Island Ecology and Biogeography Research Group, Instituto Universitario de Enfermedades 13
- Tropicales y Salud Pública de Canarias (IUETSPC), Universidad de La Laguna, Tenerife, 14
- Canary Islands, Spain 15
- <sup>5</sup> Universite de Pau et des Pays de l'Adour, E2S UPPA, CNRS, IPREM, 64000 Pau, France 16
- <sup>6</sup> Joint Research Centre (JRC) of the European Commission, Directorate for Sustainable 17
- 18 Resources, Ispra, VA, 21027 Italy
- <sup>7</sup> UMR CNRS 7058 "Ecologie et Dynamique des Systèmes Anthropisés" (EDYSAN), 19
- Université de Picardie Jules Verne, Amiens, France 20
- <sup>8</sup> Biogeography and Biodiversity Lab, Institute of Physical Geography, Goethe-University 21
- Frankfurt, 60438 Frankfurt am Main, Germany 22
- <sup>9</sup> Department of Biogeography and Global Change, Museo Nacional de Ciencias Naturales 23
- (BGC-MNCN-CSIC), E-28006 Madrid, Spain 24
- <sup>10</sup> Departament de Biologia Evolutiva, Ecologia i Ciències Ambientals, Facultat de Biologia, 25
- Universitat de Barcelona (UB), Spain 26
- <sup>11</sup> Institut de Recerca de la Biodiversitat (IRBio), Universitat de Barcelona (UB), Barcelona, 27
- 28 Spain.
- <sup>12</sup> Department of Entomology, Washington State University, Pullman, Washington State, 29
- USA 99163 30

31

- <sup>13</sup> University of Göttingen, Department of Biodiversity, Macroecology & Biogeography, 32
- Büsgenweg 1, 37077 Göttingen, Germany 33

- <sup>14</sup> Radboud University, Radboud Institute for Biological and Environmental Sciences, 35
- Heyendaalseweg 135, 6525 AJ, Nijmegen, The Netherlands 36
- <sup>15</sup> Cape Research Centre, South African National Parks, 643 Levds St, Muckleneuk, Pretoria, 37
- South Africa. 38
- <sup>16</sup> School for Climate Studies, Stellenbosch University, Private Bag X1 Matieland, 7602. 39
- South Africa. 40

| 41<br>42 | <sup>17</sup> Institute for Biodiversity and Ecosystem Dynamics (IBED), University of Amsterdam, P.O. Box 94240, 1090 GE Amsterdam, The Netherlands |
|----------|---|
| 43<br>44 | <sup>18</sup> INRAE, Aix Marseille Univ., RECOVER, Aix-en-Provence, France  |
| 45       | Running head : Vulnerability of islands   |
| 46       | Keywords (6-10): islands, biodiversity, anthropogenic threats, risk, adaptive capacity,   |
| 47       | exposure, sensitivity   |
| 48       |   |
| 49       |   |
| 50       | Words: 3388   |
| 51       | Figures: 2  |
| 52       | Table: 1  |
| 53       | Box: 2  |
| 54       |   |
| 55       |   |
| 56       |   |
| 57       |   |
| 58       |   |
| 59       |   |
| 60       |   |
| 61       |   |
|          |   |

Abstract:

The majority of vulnerability assessments of biodiversity to global changes have so far focused on mainland systems, ignoring islands. However, islands harbour unique biodiversity and are epicentres of ongoing extinctions. We thus introduce a new framework for quantifying the vulnerability of terrestrial insular biota to multiple threats. This framework uses markers of exposure, sensitivity, and adaptive capacity that reflect the unique characteristics of island biodiversity, from population to island levels. Our framework involves four steps: (1) identifying the scope of the vulnerability assessment, (2) selecting the most appropriate markers, (3) computing the vulnerability, and (4) provide recommendations for conservation. We discuss the need and urgency to deploy this framework to guide evidence-based decisions for the conservation of insular biodiversity and for an improved attention of insular biota at the science-policy interface. 

#### **Introduction:**

83

84

85

86

87

88

89

90

91

92

93

94

95

96

97

98

99

100

101

102

103

104

105

106

107

Islands are among the most vulnerable ecosystems affected by global change (Fernández-Palacios et al., 2021). Approximately ca. 75% of extinctions have occurred on islands, and over half of all terrestrial species that face imminent extinction are island-dwellers, with invasive alien species and land-use change, as leading drivers of species' declines. Climate change emerges as a growing threat for island biota and it may interact in unexpected ways with invasive alien species and land-use change (Capdevila et al., 2021; Mantyka-Pringle et al., 2011). As a result, islands are commonly considered the epicentres of past, imminent, and potential future species extinctions (Supplementary appendix S1). At least three reasons can explain the disproportionate vulnerability of island ecosystems. First, insular species are usually intrinsically more sensitive to any given threat due to specific traits deriving from the so-called 'island syndrome' (Benítez-López et al., 2021; Biddick et al., 2019; Lomolino, 1985). Second, insular species are less likely to adapt to new threats due to their inherent demographic features (e.g., small population sizes, naturally fragmented distribution ranges) (Fernández-Palacios et al., 2021). Lastly, the physiography of islands, specifically in the case of isolated, small-sized ones, renders their biota more exposed to threats and also less able to escape compared to their continental counterparts (Harter et al., 2015, p. 201). Despite the urgency needed to protect the unique and highly vulnerable island biodiversity from ongoing global change, insular biota is only briefly mentioned in international biodiversity policy frameworks. In December 2022 the 196 parties of the Convention on Biological Diversity (CBD) adopted the Kunming-Montreal Global Biodiversity Framework (GBF), involving 23 action-oriented global targets. Among them, only one target explicitly mentions islands as priority areas for conservation (though not explicitly mentioning insular biota) and only two other targets require strong actions on islands. This lack of attention

reflects the scientific bias towards mainland and charismatic species (Albert et al., 2018; Rodrigues et al., 2010), with remote islands being marginally considered in biodiversity assessments, which hinders the development of effective conservation plans on insular biota (Troudet et al., 2017). Here, we introduce a novel framework specifically designed to assess the vulnerability of insular biota to global change, across multiple biodiversity dimensions, and based on insular biota's exposure, sensitivity, and adaptive capacity to multiple threats. We first describe the characteristics that contribute to the vulnerability of island biota to global change, which need to be considered in insular biodiversity vulnerability assessments. Then, we describe how our framework can address some of the ongoing questions on island biota vulnerability.

#### Challenges

The uniqueness of insular biota enhances their vulnerability to threatening processes. The inherent uniqueness of insular biota (e.g., endemic species, distinct traits, unique lineages) increases their overall vulnerability to global change drivers (Fernández-Palacios et al., 2021). Their isolated nature and complex topography, prone to extreme events, make them more exposed to threats (Russell & Kueffer, 2019) (Figure 1A) because islands per se harbour fewer potential refugia (compared with continental landmasses). Islands are also known to host a large number of endemic species: up to 90% of endemic non-vagile taxa (i.e., non-flying vertebrates, seed plants, molluscs or arthropods) occur in oceanic islands and continental fragments (e.g., Madagascar, Antonelli et al., 2022). Endemics have small population sizes and restricted geographical ranges (sometimes limited to an archipelago, an island or even a single peak, volcano or cliff), increasing their extinction probability (Manes et al., 2021). Iconic examples are the radiation of the Hawaiian silversword alliance, which

gave rise to the radiation of Hawaiian Honeycreepers that are particularly vulnerable to threats (Box 1).

## A Why are island biota particularly vulnerable to global change?

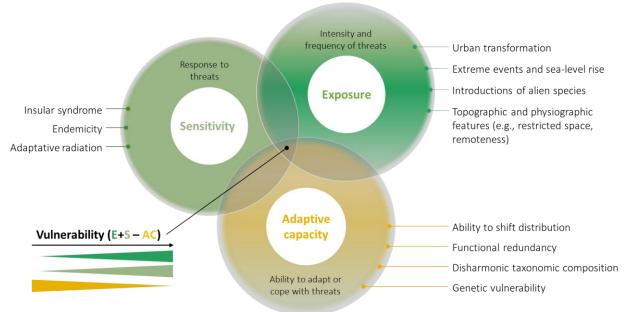




Figure 1: A Conceptual figure of the vulnerability of insular biota to global change with a non-exhaustive list of the characteristics contributing to each vulnerability component (i.e., exposure, sensitivity, and adaptive capacity). For example, insular syndrome traits, endemicity, and adaptive radiation are characteristics that increase the sensitivity of insular biota to global change compared to mainland biota, whereas lack of functional redundancy, disharmonic taxonomic composition, and genetic vulnerability are characteristics that decrease adaptive capacity. B Bias in current climate change vulnerability assessments in terms of island biota representation, taxonomic representation, and vulnerability components, numbers calculated from (de los Ríos et al., 2018).

Insular species have also evolved distinct and predictable ecological, physiological,

behavioural, morphological, and life-history traits, a phenomenon commonly referred to as

the 'island syndrome' (Baeckens & Van Damme, 2020), worsening their sensitivity to current and future anthropogenic threats, including overharvesting, habitat loss, increased drought and alien invasive species (Rozzi et al., 2023; Zizka et al., 2022), (Figure 1A). This phenomenon is for example associated with dioecism in plants, which makes them more likely to disappear if their mutualists disappear, presumably causing higher vulnerability to human induced perturbations (Schrader et al., 2021). The reduced herbivory and predation pressure on islands leads to low levels of spinescence compared to mainland, which makes island species more likely to be affected by the introduction of exotic herbivores and predators (K. Burns, 2016; Clavero et al., 2009, p. 200). The loss of flight capacities also makes insular birds unable to escape from anthropogenic threats (Sayol et al., 2021) and more likely to be threatened by extreme events, such as hurricanes, or gradual environmental change (e.g., progressive directional changes such as climate change) (Burns, 2019; Roff, 1990). Additionally, the intricate and often rugged topography of oceanic islands frequently leads to isolations of populations, which may result in genetic drift and then inbreeding when the population sizes are small. Subsequent island colonization within archipelagos and isolation by distance could too led to high population structure between island (Parmakelis et al., 2015; White & Searle, 2007). These populations are thus more prone to genetic diversity loss, which may lead to fewer opportunities to adapt to changes. Although this does not necessarily increase their vulnerability per se, the loss of genetic diversity due to disturbances, especially when exposed to disturbances over a short time period (Inamine et al., 2022), can cause a demographic or genetic collapse. Thus, losing an insular population can cause demographic and genetic loss, whereas it causes only demographic loss on mainland. Because of that, the effective conservation units on islands should be at the population level, rather than species level (e.g., (Melo & O'Ryan, 2007)).

147

148

149

150

151

152

153

154

155

156

157

158

159

160

161

162

163

164

165

166

167

168

169

Besides population- and species-level characteristics, the composition of insular species' assemblages harbours features that make them more vulnerable. An example is their disharmonic taxonomic composition, which refers to the systematic over- or under-representation of certain taxonomic groups compared to continental assemblages. Notable examples include the absence of several families of non-volant mammals on many isolated islands (Brace et al., 2015), and the over-representation of pteridophytes in plant communities (Kreft et al., 2010). The absence of certain functional groups, although partially compensated by in-situ diversification and differentiation, allows the establishment of new exotic species (Vitousek et al., 1997). These exotic species can exert a new role (predation, competition) unknown to the native community, which is evolutionarily and functionally unequipped to withstand these novel pressures. Additionally, species traits in island communities are often complementary rather than redundant, leading to communities with low functional redundancy (Harter et al., 2015; Whittaker et al., 2014), making them more sensitive to threats (McConkey & Drake, 2015) and with no 'ecological insurance' to replace the missing functions (Loreau et al., 2021).

State of the art of vulnerability assessments on insular biota

Despite the inherent vulnerability of insular biota to global changes, the majority of vulnerability assessments of biodiversity have so far focused on mainland systems, ignoring islands. Climate change vulnerability assessments emerged in the 1990s to pre-empt impacts and prepare appropriate responses (Foden et al., 2019; Solomon et al., 2007). Most definitions concur that a species' vulnerability to a threat is determined by three factors: **exposure**, **sensitivity**, and **adaptive capacity** (Butt et al., 2022; Foden et al., 2013, 2019). Using this definition, a recent review showed that among the 741 studies assessing climate change vulnerability, the large majority focused on mainland systems, and less than one third (n=231) included islands (de los Ríos et al., 2018) (see Table S2 for examples) and with only 136

studies associated with a specific insular country. Those island vulnerability assessments were often geographically or taxonomically restricted towards high income countries (e.g., Australia, the UK, and the USA account for 60% of studies on insular biota), and plants (49% of studies) (Figure 1B). In addition, the large majority of vulnerability assessments do not consider the influence of multiple threats (but see Santos et al., 2021; Sousa et al., 2021; Ureta et al., 2022), which are likely to act together and be exacerbated by climate change (e.g., habitat loss, overexploitation, biological invasions). Making the assumption that exposure is a suitable proxy for the impact of a particular threat, most vulnerability assessment studies focused on exposure alone, and less than 10% relied on the three specified components of vulnerability (Butt et al., 2016; de los Ríos et al., 2018). The implicit associated assumption is that all species will have equal responses to a threat, which is highly unlikely. In fact, species' responses to threats, either on mainland or islands, are modulated by their traits (Fromm & Meiri, 2021; Leclerc, Villéger, et al., 2020; Marino et al., 2022; Soares et al., 2022). We argue here that, applied to a range of taxa and threats, traitbased vulnerability assessments can provide a useful approach for (i) developing a more comprehensive index of vulnerability to threats, and (ii) informing effective management actions for conservation (Gallagher et al., 2021).

#### **Perspective**

196

197

198

199

200

201

202

203

204

205

206

207

208

209

210

211

212

213

214

215

216

217

218

219

220

We present a conceptual framework divided in four steps that provides a roadmap for vulnerability assessments in insular systems, considering the three components of vulnerability (i.e., exposure, sensitivity, and adaptive capacity; Figure 2). We built upon previous works focused on taxonomic diversity in mainland ecosystems (Foden et al., 2013; Parravicini et al., 2014) to include in our framework various markers for each vulnerability component related to the inherent characteristics of island biota, as well as multiple threats, taxa, and dimensions of diversity (i.e., taxonomic, functional and phylogenetic diversity; TD,

FD and PD). The development of this framework is part of a larger initiative to meet 221 international policy targets that better integrate biodiversity threats and dimensions (Box 2). 222 223 Step 1: Identify the scope and aim of the vulnerability of insular biota The first step is to define the scope of the vulnerability assessment in terms of spatial and 224 225 temporal extent, relevant threats, and studied biota (taxonomic group, biological level, 226 species biogeographical origin). This challenging task is pivotal for the assessment design, which in turn affects the final step of informing conservation actions (Step 4). For example, 227 broad-scale assessments (e.g., at the global extent or among several archipelagos) contribute 228 229 to strategic planning and to establish a common baseline of vulnerability information (e.g., IPBES assessments), while local-scale assessments (e.g., within archipelagos or group of 230 islands) are appropriate for informing site-level management decisions. Also at the global 231

scale, special attention should be paid to endemic species or insular populations, since island

biodiversity conservation operates at the population level.

232

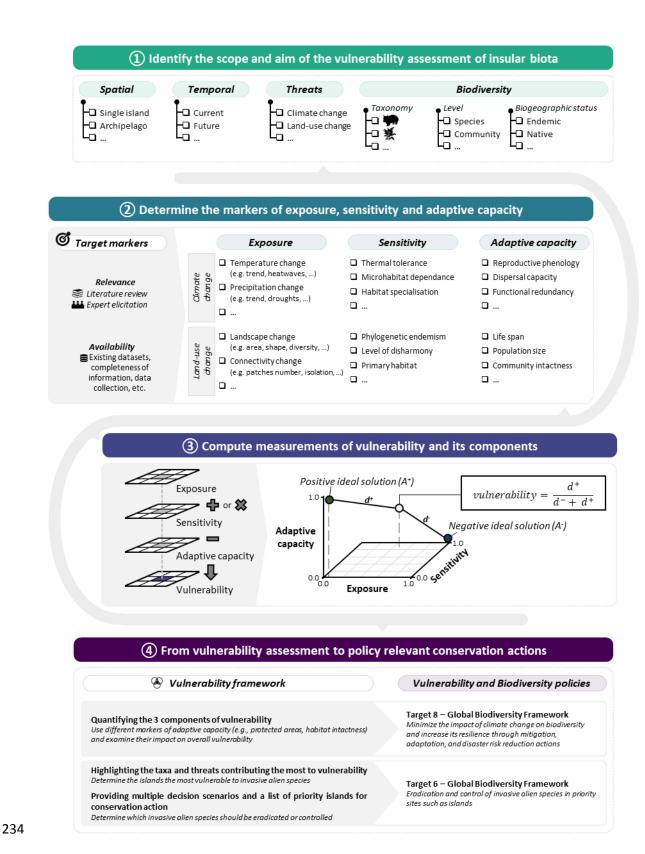


Figure 2: Conceptual framework for assessing island biodiversity vulnerability to global change.

239

There are many different relevant markers associated with species' vulnerability to threats. 240 241 Those markers may vary in type (e.g., ecological or demographic) and organisation level (e.g., 242 population, species, community, or ecosystem). One can review the scientific literature and elicit expert opinion to identify and collect relevant markers for each component of 243 244 vulnerability based on the scope and purpose of the assessment. For instance, assessing the vulnerability of terrestrial species to land use change and climate change requires markers of 245 246 exposure to climate change (e.g., change in precipitation regimes, sea-level rise) and land use change (e.g., urbanization rates, forest conversion into agriculture, infrastructure 247 development, shift from agriculture to tourism). Moreover, a list of biological traits to infer 248 249 the sensitivity and adaptive capacity of the studied insular species to those threats are needed 250 (Figure 2 and Table S1 for examples of exposure, sensitivity, and adaptive capacity markers). 251 Markers for exposure can be linked to the focal threat(s) or general to global changes, and multiple markers should be used to capture the multiple dimensions of changes (e.g., 252 temperature change, heat waves, droughts). Species' sensitivity is determined by their 253 254 intrinsic traits (e.g., climatic niche breadth, habitat specialization, diet breadth, body size, etc.). Some sensitivity traits might be specific to a given threat (e.g., temperature tolerance), 255 while others are general and encompass the sensitivity to different threats (e.g., body size, 256 257 dependence on interspecific interactions). Note that for both sensitivity and adaptive capacity, a long list of markers is applicable at either the population or species level (see (Thurman et 258 259 al., 2020) for a list and Table S1). However, markers applicable at the community level such 260 as functional or phylogenetic diversity could also be included (Table 1 for example). 261 Adaptive capacity refers to the ability of insular biota to respond to a stressor by persisting in 262 situ in the face of threats or novel conditions, or by shifting in space or time (following (Thurman et al., 2020)). Acclimation, behavioural change, phenotypic plasticity and 263

evolutionary adaptation may all contribute to adaptative capacity (Foden et al., 2018; Royer-Tardif et al., 2021). For mobile animals, especially the most vagile organisms, markers of adaptive capacity can incorporate traits linked to movement or mobility (e.g., migration frequency and distance, site fidelity) (Butt et al., 2022; Thurman et al., 2020). Note that for plants and sessile organisms in general, adaptive capacity mostly refers to their capacity to persist *in situ* with mating system and fecundity as strong markers. Finally, adaptive capacity of insular biota can also be modulated by extrinsic factors, such as habitat quality, area connectivity and level of protection, factors which could therefore be included in the measure of adaptive capacity at the system level (i.e., island, archipelago).

| Functional-based markers | Phylogeny-based markers |
|--------------------------|-------------------------|
|--------------------------|-------------------------|

Table 1: Proposed functional-based and phylogeny-based metrics that could be used as markers at the community level in vulnerability frameworks.

| g                 | - · · · ·  | 77.1   |
|-------------------|--|--|
| Sensitivity       | Functional rarity:   | Phylogenetic endemism:   |
|                   | Definition - Functional rarity, expressed at the species level, is the combination   | Definition - Phylogenetic endemism is the spatial restriction of the phylogenetic  |
|                   | of the functional uniqueness, the  | diversity of a community (ROSAUER et   |
|                   | scarcity and the geographic  | al., 2009). In other words, this is a relative   |
|                   | restrictedness of a species (Violle et al.,  | measure of endemism that represents the  |
|                   | 2017).   | degree to which lineages or branches of  |
|                   |  | the tree of life are spatially restricted.   |
|                   |  |  |
|                   | Rationale - Given that functionally rare   | Rationale - Areas with high phylogenetic   |
|                   | species can play a critical role in  | endemism are characterized by the  |
|                   | ecosystem functioning, and that they   | presence of species that have diversified  |
|                   | rarity make them heightened sensitivity  | and evolved in response to specific  |
|                   | to anthropogenic pressures (Davies et  | environmental conditions within a  |
|                   | al., 2004; Loiseau et al., 2020), areas with a high proportion of functionally   | particular location. Among the drivers shaping phylogenetic endemism, climate  |
|                   | rare species are particularly susceptible  | plays a significant role (Guo et al., 2023),   |
|                   | to threats?  | implying that changes in climate   |
|                   |  | conditions may disproportionately affect   |
|                   |  | these species, potentially leading to their  |
|                   |  | decline or extinction.   |
|                   |  |  |
|                   |  |  |
| Adaptive          | Functional redundancy:   | Phylogenetic distinctiveness:  |
| Adaptive capacity | Functional redundancy:  Definition - Functional redundancy of a  | Phylogenetic distinctiveness:  Definition - Phylogenetic distinctiveness   |
|                   | Definition - Functional redundancy of a given community reflects the tendency  | Definition - Phylogenetic distinctiveness reflects the degree of isolation of a species  |
|                   | Definition - Functional redundancy of a given community reflects the tendency for the constituent species to perform   | Definition - Phylogenetic distinctiveness reflects the degree of isolation of a species or a group of species within a   |
|                   | Definition - Functional redundancy of a given community reflects the tendency  | Definition - Phylogenetic distinctiveness reflects the degree of isolation of a species or a group of species within a phylogenetic tree (Pavoine & Ricotta,   |
|                   | Definition - Functional redundancy of a given community reflects the tendency for the constituent species to perform similar functions (Mouillot et al., 2014).  | Definition - Phylogenetic distinctiveness reflects the degree of isolation of a species or a group of species within a phylogenetic tree (Pavoine & Ricotta, 2021).  |
|                   | Definition - Functional redundancy of a given community reflects the tendency for the constituent species to perform similar functions (Mouillot et al., 2014).  Rationale - Functional redundancy has   | Definition - Phylogenetic distinctiveness reflects the degree of isolation of a species or a group of species within a phylogenetic tree (Pavoine & Ricotta, 2021).  Rationale - Evolutionarily distinct species   |
|                   | Definition - Functional redundancy of a given community reflects the tendency for the constituent species to perform similar functions (Mouillot et al., 2014).  Rationale - Functional redundancy has been theoretically and empirically linked   | Definition - Phylogenetic distinctiveness reflects the degree of isolation of a species or a group of species within a phylogenetic tree (Pavoine & Ricotta, 2021).  Rationale - Evolutionarily distinct species or group of species represent uniquely  |
|                   | Definition - Functional redundancy of a given community reflects the tendency for the constituent species to perform similar functions (Mouillot et al., 2014).  Rationale - Functional redundancy has been theoretically and empirically linked to the concepts of resistance and   | Definition - Phylogenetic distinctiveness reflects the degree of isolation of a species or a group of species within a phylogenetic tree (Pavoine & Ricotta, 2021).  Rationale - Evolutionarily distinct species or group of species represent uniquely divergent genomes (Warren et al., 2008).   |
|                   | Definition - Functional redundancy of a given community reflects the tendency for the constituent species to perform similar functions (Mouillot et al., 2014).  Rationale - Functional redundancy has been theoretically and empirically linked   | Definition - Phylogenetic distinctiveness reflects the degree of isolation of a species or a group of species within a phylogenetic tree (Pavoine & Ricotta, 2021).  Rationale - Evolutionarily distinct species or group of species represent uniquely divergent genomes (Warren et al., 2008). Consequently, sets of evolutionarily distinct species are expected to encompass   |
|                   | Definition - Functional redundancy of a given community reflects the tendency for the constituent species to perform similar functions (Mouillot et al., 2014).  Rationale - Functional redundancy has been theoretically and empirically linked to the concepts of resistance and resilience of ecosystem functioning to species loss (Biggs et al., 2020) via the insurance hypothesis (McCann, 2000).   | Definition - Phylogenetic distinctiveness reflects the degree of isolation of a species or a group of species within a phylogenetic tree (Pavoine & Ricotta, 2021).  Rationale - Evolutionarily distinct species or group of species represent uniquely divergent genomes (Warren et al., 2008). Consequently, sets of evolutionarily distinct species are expected to encompass a large proportion of the parental clade's  |
|                   | Definition - Functional redundancy of a given community reflects the tendency for the constituent species to perform similar functions (Mouillot et al., 2014).  Rationale - Functional redundancy has been theoretically and empirically linked to the concepts of resistance and resilience of ecosystem functioning to species loss (Biggs et al., 2020) via the insurance hypothesis (McCann, 2000). Hence, more functionally redundant  | Definition - Phylogenetic distinctiveness reflects the degree of isolation of a species or a group of species within a phylogenetic tree (Pavoine & Ricotta, 2021).  Rationale - Evolutionarily distinct species or group of species represent uniquely divergent genomes (Warren et al., 2008). Consequently, sets of evolutionarily distinct species are expected to encompass a large proportion of the parental clade's total phylogenetic diversity, which may  |
|                   | Definition - Functional redundancy of a given community reflects the tendency for the constituent species to perform similar functions (Mouillot et al., 2014).  Rationale - Functional redundancy has been theoretically and empirically linked to the concepts of resistance and resilience of ecosystem functioning to species loss (Biggs et al., 2020) via the insurance hypothesis (McCann, 2000). Hence, more functionally redundant systems should show greater resilience   | Definition - Phylogenetic distinctiveness reflects the degree of isolation of a species or a group of species within a phylogenetic tree (Pavoine & Ricotta, 2021).  Rationale - Evolutionarily distinct species or group of species represent uniquely divergent genomes (Warren et al., 2008). Consequently, sets of evolutionarily distinct species are expected to encompass a large proportion of the parental clade's total phylogenetic diversity, which may play a crucial role in ensuring long-term  |
|                   | Definition - Functional redundancy of a given community reflects the tendency for the constituent species to perform similar functions (Mouillot et al., 2014).  Rationale - Functional redundancy has been theoretically and empirically linked to the concepts of resistance and resilience of ecosystem functioning to species loss (Biggs et al., 2020) via the insurance hypothesis (McCann, 2000). Hence, more functionally redundant systems should show greater resilience to perturbation (Mouillot et al., 2014)   | Definition - Phylogenetic distinctiveness reflects the degree of isolation of a species or a group of species within a phylogenetic tree (Pavoine & Ricotta, 2021).  Rationale - Evolutionarily distinct species or group of species represent uniquely divergent genomes (Warren et al., 2008). Consequently, sets of evolutionarily distinct species are expected to encompass a large proportion of the parental clade's total phylogenetic diversity, which may play a crucial role in ensuring long-term stability and resilience (Cadotte et al.,  |
|                   | Definition - Functional redundancy of a given community reflects the tendency for the constituent species to perform similar functions (Mouillot et al., 2014).  Rationale - Functional redundancy has been theoretically and empirically linked to the concepts of resistance and resilience of ecosystem functioning to species loss (Biggs et al., 2020) via the insurance hypothesis (McCann, 2000). Hence, more functionally redundant systems should show greater resilience   | Definition - Phylogenetic distinctiveness reflects the degree of isolation of a species or a group of species within a phylogenetic tree (Pavoine & Ricotta, 2021).  Rationale - Evolutionarily distinct species or group of species represent uniquely divergent genomes (Warren et al., 2008). Consequently, sets of evolutionarily distinct species are expected to encompass a large proportion of the parental clade's total phylogenetic diversity, which may play a crucial role in ensuring long-term  |
|                   | Definition - Functional redundancy of a given community reflects the tendency for the constituent species to perform similar functions (Mouillot et al., 2014).  Rationale - Functional redundancy has been theoretically and empirically linked to the concepts of resistance and resilience of ecosystem functioning to species loss (Biggs et al., 2020) via the insurance hypothesis (McCann, 2000). Hence, more functionally redundant systems should show greater resilience to perturbation (Mouillot et al., 2014) since when a species becomes extinct, its role can be fulfilled by functionally close species. This indirectly reflects the                                   | Definition - Phylogenetic distinctiveness reflects the degree of isolation of a species or a group of species within a phylogenetic tree (Pavoine & Ricotta, 2021).  Rationale - Evolutionarily distinct species or group of species represent uniquely divergent genomes (Warren et al., 2008). Consequently, sets of evolutionarily distinct species are expected to encompass a large proportion of the parental clade's total phylogenetic diversity, which may play a crucial role in ensuring long-term stability and resilience (Cadotte et al., 2012). This implies that communities exhibiting higher phylogenetic distinctiveness are more likely to harbor  |
|                   | Definition - Functional redundancy of a given community reflects the tendency for the constituent species to perform similar functions (Mouillot et al., 2014).  Rationale - Functional redundancy has been theoretically and empirically linked to the concepts of resistance and resilience of ecosystem functioning to species loss (Biggs et al., 2020) via the insurance hypothesis (McCann, 2000). Hence, more functionally redundant systems should show greater resilience to perturbation (Mouillot et al., 2014) since when a species becomes extinct, its role can be fulfilled by functionally close species. This indirectly reflects the ability of the system to adapt to | Definition - Phylogenetic distinctiveness reflects the degree of isolation of a species or a group of species within a phylogenetic tree (Pavoine & Ricotta, 2021).  Rationale - Evolutionarily distinct species or group of species represent uniquely divergent genomes (Warren et al., 2008). Consequently, sets of evolutionarily distinct species are expected to encompass a large proportion of the parental clade's total phylogenetic diversity, which may play a crucial role in ensuring long-term stability and resilience (Cadotte et al., 2012). This implies that communities exhibiting higher phylogenetic distinctiveness are more likely to harbor increased evolutionary potential, enabling |
|                   | Definition - Functional redundancy of a given community reflects the tendency for the constituent species to perform similar functions (Mouillot et al., 2014).  Rationale - Functional redundancy has been theoretically and empirically linked to the concepts of resistance and resilience of ecosystem functioning to species loss (Biggs et al., 2020) via the insurance hypothesis (McCann, 2000). Hence, more functionally redundant systems should show greater resilience to perturbation (Mouillot et al., 2014) since when a species becomes extinct, its role can be fulfilled by functionally close species. This indirectly reflects the                                   | Definition - Phylogenetic distinctiveness reflects the degree of isolation of a species or a group of species within a phylogenetic tree (Pavoine & Ricotta, 2021).  Rationale - Evolutionarily distinct species or group of species represent uniquely divergent genomes (Warren et al., 2008). Consequently, sets of evolutionarily distinct species are expected to encompass a large proportion of the parental clade's total phylogenetic diversity, which may play a crucial role in ensuring long-term stability and resilience (Cadotte et al., 2012). This implies that communities exhibiting higher phylogenetic distinctiveness are more likely to harbor  |

Step 3: Compute measures of vulnerability and its components

Once all the markers are collected, the next step is to combine those markers to determine a composite vulnerability measure. Of the myriad of vulnerability assessments that already exist (e.g., correlative, trait-based, mechanistic), not all are equivalent in terms of data requirements and objectives (see Foden et al., 2019 for an extensive review on the topic), with the correlative approaches being the most frequently used. Despite the increasing availability of traits for birds, mammals, reptiles, and plants (Díaz et al., 2022; Faurby et al., 2018; Soria et al., 2021; Tobias et al., 2021), trait-based approaches have been barely applied to date. Among the trait-based approaches, most are criteria-based, in which biota are classified into categories of vulnerability from low to high when summing up the different components of vulnerability (hereby referred to as qualitative or semi-quantitative frameworks). To prevent the use of arbitrary thresholds, we propose a continuous quantitative framework where a multicriteria decision analysis can be applied (Leclerc, Courchamp, et al., 2020; Parravicini et al., 2014) and provide continuous values and a ranking of islands or archipelagos based on their vulnerability scores. This technique ranks alternatives according to their distance to positive (i.e., no vulnerability, where exposure and sensitivity are minimized and adaptive capacity is maximized) and negative ideal solutions (see (Leclerc, Courchamp, et al., 2020) for details). Note that alternative approaches exist for combining the three vulnerability components such as additive effect between components (Nyboer et al., 2021) or interacting effect between exposure and sensitivity for instance (Silva Rocha et al., 2024).

310

311

312

313

314

315

291

292

293

294

295

296

297

298

299

300

301

302

303

304

305

306

307

308

309

Step 4: From vulnerability assessment to policy relevant conservation actions

The vulnerability framework developed could be used to increase the fundamental knowledge on vulnerability (Step 1-3) but has also the potential to help implementing biodiversity policies (Step 4, Box 2). For instance, one of the targets of the Global Biodiversity Framework is to minimize the impact of climate change on biodiversity through (among

others) mitigation, adaptation and risk disaster reduction actions. Such aims require to use different markers of adaptive capacity through multiple plausible scenarios (e.g., protected areas, habitat intactness) to inform on how different management actions can impact the overall vulnerability. Vulnerability assessments can help targeting the control and eradication of invasive alien species in conservation priority sites such as islands (Target 6), but also identifying the sites that need to be under protection (Target 3) to conserve biodiversity. Effective conservation practice relies on understanding biodiversity's vulnerability through multiple lenses. Where the exposure type is common across species (e.g., inundation, invasive species), the threat itself may be managed through dedicated programs for, like mangrove rehabilitation, or invasive species control (Jones et al., 2016). Finally, beyond its use in current policies, our vulnerability framework could be useful in drafting new policies. Indeed, to set relevant targets in the new policies being drafted (e.g., by [year], reduce by [number] % the number of species threatened by [threat]), policy makers need to know (i) what is the current situation and (ii) what could be the future situation under different scenarios. Our vulnerability framework may provide both, and can contribute to the setting of policy targets that are simultaneously ambitious and reachable for insular biodiversity.

332

333

334

335

336

337

338

339

340

316

317

318

319

320

321

322

323

324

325

326

327

328

329

330

331

#### **Concluding remarks**

Given the proliferation of threats that islands are facing, efforts must be made to study island biota in the light of global change, and thus to embrace the whole of biodiversity, mainland and insular, in current vulnerability assessments in international instance such as IPCC or IPBES. Our comprehensive and detailed framework lays the foundations to understand and predict island biodiversity vulnerability to global change. Yet, the outcomes of vulnerability assessments will be challenged by missing data and, in the case of future assessments, by uncertainties about the future trajectories of anthropogenic threats. In this context, it is crucial

| 341 | to estimate the existing uncertainty as wen as to communicate this uncertainty (e.g., (Hossain   |
|-----|--|
| 342 | et al., 2019; Rocchini et al., 2011; Tessarolo et al., 2021; Thuiller et al., 2019)), which will |
| 343 | ultimately help to identify future research priorities.  |
| 344 |  |
| 345 |  |
| 346 |  |
| 347 |  |
| 348 |  |
| 349 |  |
| 350 |  |
| 351 |  |
| 352 |  |
| 353 |  |
| 354 |  |
| 355 |  |
| 356 |  |
| 357 |  |
| 358 |  |
| 359 |  |
| 360 | Glossary:  |

| 361<br>362<br>363<br>364<br>365 | Adaptive capacity: the ability to adapt to changing conditions for an individual, population or a species; this may be via ecological (i.e., physiological and/or behavioural plasticity) or evolutionary adaptation (i.e., through natural selection acting on quantitative traits).  Adaptive radiation: the rapid diversification of a single evolutionary lineage into multiple |
|---------------------------------|---|
| 366                             | ecologically or morphologically distinct species.   |
| 367<br>368<br>369               | <b>Exposure</b> : the extent to which each species' physical environment changes due to global threats. It includes the intensity, magnitude and frequency of the threat(s).  |
| 370<br>371                      | <b>Sensitivity</b> : the intrinsic capacity of species to cope with threats, based on their life-history, ecology, morphology, or behavior.   |
| 372                             |   |
| 373<br>374                      | <b>Threats</b> : external factors that have the potential to impact the viability, abundance, distribution, or behavior of an individual, species, community or an ecosystem.   |
| 375<br>376                      | <b>Vulnerability</b> : susceptibility of a system/species to a negative impact following a threat   |
| 377                             | value and the state of a system species to a negative impact ions wing a uncar  |
| 378                             |   |
| 379                             |   |
| 380                             |   |
| 381                             |   |
| 382                             |   |
| 383                             |   |
| 384                             |   |
| 385                             |   |
| 386                             |   |
| 387                             |   |
| 388                             |   |
| 389                             |   |
| 390                             |   |
| 391                             |   |
| 392                             |   |
| 393                             |   |
| 394                             |   |
| 395                             |   |
| 396                             |   |
| 397                             |   |
| 398                             |   |
| 399                             |   |
| 400                             |   |
| 401                             |   |
| 402                             |   |
| 403<br>404                      |   |
| 404<br>405                      |   |
| 405<br>406                      |   |
| 407                             |   |
| 407                             | Box 1   The case of the Hawaiian honeycreepers  |
| 400                             | DOX 1   THE CASE OF THE FIAWAHAN HONEYCI CEPELS   |

The Hawaiian Islands constitute an archipelago of eight larger and numerous smaller volcanic

islands in the North Pacific Ocean, in total about 16,644 km² in size, and about 3,200 km from

the nearest mainland. The islands in the northwest are older and typically smaller, due to

progressive erosion, while the islands in the southeast are larger and still volcanically active.

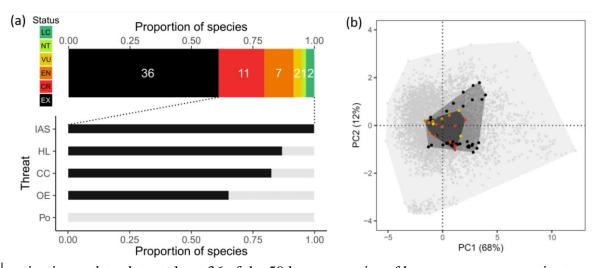
Thanks to the archipelago's extreme isolation and the islands' variation in size and

|  | environmental conditions | , Hawaii has been | n a hotspot of s | speciation and | adaptive radiation |
|--|--------------------------|-------------------|------------------|----------------|--------------------|
|--|--------------------------|-------------------|------------------|----------------|--------------------|

(Baldwin & Sanderson, 1998; Lerner et al., 2011; Price, 2004). The Hawaiian honeycreepers

(Fringillidae: Drepanidinae) represent one of the most iconic examples of adaptive radiation,

which has resulted in a striking variation in bill morphology (Lerner et al., 2011). This strong adaptive radiation, however, has also led to species with highly restricted ranges and naturally small population sizes. Hence, little is needed to push these species over the brink of



extinction and, to date, at least 36 of the 59 known species of honeycreepers are extinct (Figure 3a). Extinction has been non-random with respect to the functional trait space of the species, with extinct species generally larger in size (Figure 3b). The main drivers of past extinction, as well as current threats to the species, are the introduction of alien species and the loss of habitat due to conversion to agricultural land. Introduced species include predators of birds and eggs (rats, cats, dogs), herbivores that modify the habitat (for example, the extinction of Laysan Honeycreeper *Himatione fraithii* is ascribed to the introduction of rabbits, which eliminated virtually all vegetation cover from the Laysan islands), and vector species of infectious diseases, such as avian malaria (Benning et al., 2002). Habitat loss began with Polynesian colonists, who cleared much of the low-elevation and seasonally dry forest for agricultural purposes, and was continued by later European colonists, who additionally converted high-elevation forests for pasturage (Riper & Scott, 2017). Climatic factors contributed little to past extinctions but are a progressive threat to extant species, especially because it may lead to an upslope shift of infectious diseases (notably avian malaria) (Benning et al., 2002).

Figure 3: Conservation status, threats, and morphospace of Hawaiian Honeycreepers. (a) IUCN Red List threat status of 59 known species of honeycreeper (LC: least concern, NT: near threatened, VU: vulnerable, EN: endangered, CR: critically endangered, EX: extinct) and prevalence of threats among the 23 extant species (IAS: invasive alien species, HL: habitat loss, CC: climate change and extreme events, OE: over-exploitation, Po: pollution) (b) The morphospace compared between 19 extant honeycreepers (dark grey polygon), 54 extant and extinct honeycreepers (medium grey polygon), and 5,974 extant passerine birds globally (light grey polygon). The morphospaces for the extant species were obtained through a principal component analysis (PCA) based on the eight morphological traits available in AVONET (Tobias et al., 2021), namely bill length from tip to skull along the culmen, bill length to nostrils, bill width and depth to nostrils, tarsal, wing and tail length, and Kipp's distance. The same eight traits for the extinct honeycreepers were taken from (Matthews et al., 2023). The first two principal components explain 80% of the total morphological variance and are primarily associated with body size (PC1) and tarsus length versus Kipp's distance (PC2), respectively. The list of species was obtained from (Matthews et al., 2023; Ricklefs, 2017) and the IUCN Red List of Threatened Species v2023-1. Species names were harmonized and duplicates removed based on the taxonomic backbone used by the IUCN.

435 436

437

438

439 440

441

442 443

444

445

446 447

448

449 450

451452453

### Box 2 A novel vulnerability framework fit for multiple policies

455

456

457

458

459

460

461

462

463

464

465

466

467

468

469

470

471

472

473

474

475

476

477

478

This novel vulnerability framework proposed for insular biodiversity is particularly relevant to inform various vulnerability policies (in orange) and biodiversity policies (in green) at both international and supranational levels such as the European Union (EU). First, by quantifying the three components of island vulnerability, our framework can directly feed the currently missing biodiversity component of the Multidimensional Vulnerability Index (MVI). The MVI is developed by the Small Island Developing States (SIDS; https://www.un.org/ohrlls/mvi/) with the United Nations to characterise the financing help for sustainable development needed in the face of global change (1). The SIDS is a political coalition of 39 low-lying islands that are united by the threat posed by climate change to their survival. In climate change negotiations, they are a loud and powerful voice for upscaling climate action since they are disproportionally affected by climate change consequences. To understand and respond, several assessments of these islands' climate change vulnerability have been carried out (e.g. (UN High Level Panel, 2022; UNFCCC, 2007)), but none includes more than a broad mention of biodiversity impacts. A scientifically rigorous assessment of climate change vulnerability of island biodiversity is, therefore, both extremely important and critically urgent. Our framework can also help EU Member States, which are required to report to the European Commission on their disaster risk management activities, to identify which components of vulnerability they can act on (2). The risk of biodiversity loss has been recently included in the Recommendations for National Risk Assessment for Disaster Risk Management in the EU (Poljansek et al., 2021), but it does not cover specifically insular biodiversity. In that context, our framework can provide a real additional value for the EU Member States with insular territories. Our framework can also further help identifying whether mitigation strategies for islands to reach Target 8 of the Kunming-Montreal Global

Biodiversity Framework (hereafter, GBF) should focus on limiting territories' exposure or promoting the adaptive capacity of insular biota (3). Second, our framework can help EU Member States with insular territories to identify and prioritise which species, sites and threats they should use to best mitigate their vulnerability in the context of their National Risk Assessment (4). Such actions can also contribute to (i) identifying the species, sites and to focus the management measures to reach Target 4 of the GBF (5), (ii) determining the most vulnerable islands to invasive alien species to reach Target 6 of the GBF (6), and helping EU Member States with insular territories to identify species threatened by invasive alien species on which to focus conservation actions to reach Target 12 of the EU Biodiversity Strategy for 2030 (EU BDS, 7). Finally, by providing multiple scenarios of threats trajectories and a list of priority islands for conservation action, our framework can contribute to document the risk of biodiversity loss under different conservation scenarios to EU Member States with insular territories (8). The vulnerability framework could contribute to identifying priority islands for restoration (target 2 of the GBF and target 4 of the EU BDS (9)) and for protection (target 3 of the GBF and target 1 of the EU BDS (10)), as well as in determining which invasive alien species should be eradicated or controlled (target 6 of the GBF (11)).

479

480

481

482

483

484

485

486

487

488

489

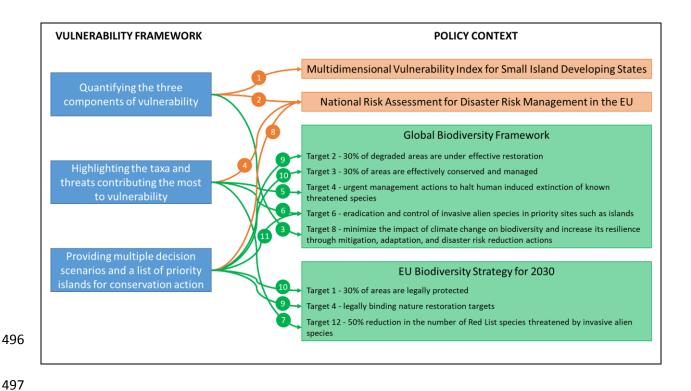
490

491

492

493

494



**Funding :** This research is product of the RIVAGE group funded by the synthesis center CESAB of the French Founda $\Theta$ on for Research on Biodiversity

(FRB; www.fondationbiodiversite.fr)

| 528 | Albert, C., Luque, G. M., & Courchamp, F. (2018). The twenty most charismatic species. <i>PLOS ONE</i> ,        |
|-----|---|
| 529 | 13(7), e0199149. https://doi.org/10.1371/journal.pone.0199149   |
| 530 | Antonelli, A., Smith, R. J., Perrigo, A. L., Crottini, A., Hackel, J., Testo, W., Farooq, H., Torres Jiménez,   |
| 531 | M. F., Andela, N., Andermann, T., Andriamanohera, A. M., Andriambololonera, S., Bachman,                        |
| 532 | S. P., Bacon, C. D., Baker, W. J., Belluardo, F., Birkinshaw, C., Borrell, J. S., Cable, S.,                    |
| 533 | Ralimanana, H. (2022). Madagascar's extraordinary biodiversity: Evolution, distribution, and                    |
| 534 | use. Science, 378(6623), eabf0869. https://doi.org/10.1126/science.abf0869                                      |
| 535 | Baeckens, S., & Van Damme, R. (2020). The island syndrome. <i>Current Biology</i> , 30(8), R338–R339.           |
| 536 | https://doi.org/10.1016/j.cub.2020.03.029   |
| 537 | Baldwin, B. G., & Sanderson, M. J. (1998). Age and rate of diversification of the Hawaiian silversword          |
| 538 | alliance (Compositae). Proceedings of the National Academy of Sciences, 95(16), 9402–9406.                      |
| 539 | https://doi.org/10.1073/pnas.95.16.9402   |
| 540 | Benítez-López, A., Santini, L., Gallego-Zamorano, J., Milá, B., Walkden, P., Huijbregts, M. A. J., &            |
| 541 | Tobias, J. A. (2021). The island rule explains consistent patterns of body size evolution in                    |
| 542 | terrestrial vertebrates. Nature Ecology & Evolution, 5(6), Article 6.   |
| 543 | https://doi.org/10.1038/s41559-021-01426-y  |
| 544 | Benning, T. L., LaPointe, D., Atkinson, C. T., & Vitousek, P. M. (2002). Interactions of climate change         |
| 545 | with biological invasions and land use in the Hawaiian Islands: Modeling the fate of endemic                    |
| 546 | birds using a geographic information system. Proceedings of the National Academy of                             |
| 547 | Sciences, 99(22), 14246–14249. https://doi.org/10.1073/pnas.162372399   |
| 548 | Biddick, M., Hendriks, A., & Burns, K. C. (2019). Plants obey (and disobey) the island rule. <i>Proceedings</i> |
| 549 | of the National Academy of Sciences, 116(36), 17632–17634.  |
| 550 | https://doi.org/10.1073/pnas.1907424116   |
| 551 | Biggs, C. R., Yeager, L. A., Bolser, D. G., Bonsell, C., Dichiera, A. M., Hou, Z., Keyser, S. R., Khursigara,   |
| 552 | A. J., Lu, K., Muth, A. F., Negrete Jr., B., & Erisman, B. E. (2020). Does functional redundancy                |

| 553 | affect ecological stability and resilience? A review and meta-analysis. Ecosphere, 11(7),                     |
|-----|---|
| 554 | e03184. https://doi.org/10.1002/ecs2.3184   |
| 555 | Brace, S., Turvey, S. T., Weksler, M., Hoogland, M. L. P., & Barnes, I. (2015). Unexpected evolutionary       |
| 556 | diversity in a recently extinct Caribbean mammal radiation. Proceedings of the Royal Society                  |
| 557 | B: Biological Sciences, 282(1807), 20142371. https://doi.org/10.1098/rspb.2014.2371                           |
| 558 | Burns, K. (2016). Spinescence in the New Zealand flora: Parallels with Australia. New Zealand Journal         |
| 559 | of Botany, 54(2), 273-289. https://doi.org/10.1080/0028825X.2015.1130727                                      |
| 560 | Burns, K. C. (2019). Evolution in Isolation: The Search for an Island Syndrome in Plants. Cambridge           |
| 561 | University Press.   |
| 562 | Butt, N., Halpern, B. S., O'Hara, C. C., Allcock, A. L., Polidoro, B., Sherman, S., Byrne, M., Birkeland, C., |
| 563 | Dwyer, R. G., Frazier, M., Woodworth, B. K., Arango, C. P., Kingsford, M. J., Udyawer, V.,                    |
| 564 | Hutchings, P., Scanes, E., McClaren, E. J., Maxwell, S. M., Diaz-Pulido, G., Klein, C. J. (2022).             |
| 565 | A trait-based framework for assessing the vulnerability of marine species to human impacts.                   |
| 566 | Ecosphere, 13(2), e3919. https://doi.org/10.1002/ecs2.3919  |
| 567 | Butt, N., Possingham, H. P., De Los Rios, C., Maggini, R., Fuller, R. A., Maxwell, S. L., & Watson, J. E. M.  |
| 568 | (2016). Challenges in assessing the vulnerability of species to climate change to inform                      |
| 569 | conservation actions. <i>Biological Conservation</i> , 199, 10–15.  |
| 570 | https://doi.org/10.1016/j.biocon.2016.04.020  |
| 571 | Cadotte, M. W., Dinnage, R., & Tilman, D. (2012). Phylogenetic diversity promotes ecosystem                   |
| 572 | stability. <i>Ecology</i> , 93(sp8), S223–S233. https://doi.org/10.1890/11-0426.1                             |
| 573 | Capdevila, P., Noviello, N., Mcrae, L., Freeman, R., & Clements, C. F. (2021). Global patterns of             |
| 574 | resilience decline in vertebrate populations. <i>Ecology Letters, October</i> , 1–12.                         |
| 575 | https://doi.org/10.1111/ele.13927   |
| 576 | Clavero, M., Brotons, L., Pons, P., & Sol, D. (2009). Prominent role of invasive species in avian             |
| 577 | biodiversity loss. <i>Biological Conservation</i> , 142(10), 2043–2049.                                       |
| 578 | https://doi.org/10.1016/j.biocon.2009.03.034  |

| 579 | Davies, K. F., Margules, C. R., & Lawrence, J. F. (2004). A SYNERGISTIC EFFECT PUTS RARE,                          |
|-----|--|
| 580 | SPECIALIZED SPECIES AT GREATER RISK OF EXTINCTION. Ecology, 85(1), 265–271.  |
| 581 | https://doi.org/10.1890/03-0110  |
| 582 | de los Ríos, C., Watson, J. E. M., & Butt, N. (2018). Persistence of methodological, taxonomical, and              |
| 583 | geographical bias in assessments of species' vulnerability to climate change: A review. Global                     |
| 584 | Ecology and Conservation, 15, e00412. https://doi.org/10.1016/j.gecco.2018.e00412                                  |
| 585 | Díaz, S., Kattge, J., Cornelissen, J. H. C., Wright, I. J., Lavorel, S., Dray, S., Reu, B., Kleyer, M., Wirth, C., |
| 586 | Prentice, I. C., Garnier, E., Bönisch, G., Westoby, M., Poorter, H., Reich, P. B., Moles, A. T.,                   |
| 587 | Dickie, J., Zanne, A. E., Chave, J., Zotz, G. (2022). The global spectrum of plant form and                        |
| 588 | function: Enhanced species-level trait dataset. Scientific Data, 9(1), 755.  |
| 589 | https://doi.org/10.1038/s41597-022-01774-9   |
| 590 | Faurby, S., Davis, M., Pedersen, R. Ø., Schowanek, S. D., Antonelli1, A., & Svenning, JC. (2018).                  |
| 591 | PHYLACINE 1.2: The Phylogenetic Atlas of Mammal Macroecology. <i>Ecology</i> , 99(11), 2626.                       |
| 592 | https://doi.org/10.1002/ecy.2443   |
| 593 | Fernández-Palacios, J. M., Kreft, H., Irl, S. D. H., Norder, S., Ah-Peng, C., Borges, P. A. V., Burns, K. C.,      |
| 594 | de Nascimento, L., Meyer, JY., Montes, E., & Drake, D. R. (2021). Scientists' warning – The                        |
| 595 | outstanding biodiversity of islands is in peril. Global Ecology and Conservation, 31, e01847.                      |
| 596 | https://doi.org/10.1016/j.gecco.2021.e01847  |
| 597 | Foden, W. B., Butchart, S. H. M., Stuart, S. N., Vié, JC., Akçakaya, H. R., Angulo, A., DeVantier, L. M.,          |
| 598 | Gutsche, A., Turak, E., Cao, L., Donner, S. D., Katariya, V., Bernard, R., Holland, R. A., Hughes,                 |
| 599 | A. F., O'Hanlon, S. E., Garnett, S. T., Sekercioğlu, C. H., & Mace, G. M. (2013). Identifying the                  |
| 600 | world's most climate change vulnerable species: A systematic trait-based assessment of all                         |
| 601 | birds, amphibians and corals. PloS One, 8(6), e65427.  |
| 602 | https://doi.org/10.1371/journal.pone.0065427   |
| 603 | Foden, W. B., Young, B. E., Akçakaya, H. R., Garcia, R. A., Hoffmann, A. A., Stein, B. A., Thomas, C. D.,          |
| 604 | Wheatley, C. J., Bickford, D., Carr, J. A., Hole, D. G., Martin, T. G., Pacifici, M., Pearce-Higgins,              |

| 605 | J. W., Platts, P. J., Visconti, P., Watson, J. E. M., & Huntley, B. (2018). Climate change                       |
|-----|--|
| 606 | vulnerability assessment of species. Wiley Interdisciplinary Reviews: Climate Change, July,                      |
| 607 | e551. https://doi.org/10.1002/wcc.551  |
| 608 | Foden, W. B., Young, B. E., Akçakaya, H. R., Garcia, R. A., Hoffmann, A. A., Stein, B. A., Thomas, C. D.,        |
| 609 | Wheatley, C. J., Bickford, D., Carr, J. A., Hole, D. G., Martin, T. G., Pacifici, M., Pearce-Higgins,            |
| 610 | J. W., Platts, P. J., Visconti, P., Watson, J. E. M., & Huntley, B. (2019). Climate change                       |
| 611 | vulnerability assessment of species. WIREs Climate Change, 10(1), e551.  |
| 612 | https://doi.org/10.1002/wcc.551  |
| 613 | Fromm, A., & Meiri, S. (2021). Big, flightless, insular and dead: Characterising the extinct birds of the        |
| 614 | Quaternary. Journal of Biogeography, 48(9), 2350–2359. https://doi.org/10.1111/jbi.14206                         |
| 615 | Gallagher, R. V., Butt, N., Carthey, A. J. R., Tulloch, A., Bland, L., Clulow, S., Newsome, T., Dudaniec, R.     |
| 616 | Y., & Adams, V. M. (2021). A guide to using species trait data in conservation. One Earth, 4(7)                  |
| 617 | 927–936. https://doi.org/10.1016/j.oneear.2021.06.013  |
| 618 | Guo, WY., Serra-Diaz, J. M., Eiserhardt, W. L., Maitner, B. S., Merow, C., Violle, C., Pound, M. J., Sun,        |
| 619 | M., Slik, F., Blach-Overgaard, A., Enquist, B. J., & Svenning, JC. (2023). Climate change and                    |
| 620 | land use threaten global hotspots of phylogenetic endemism for trees. Nature                                     |
| 621 | Communications, 14(1), 6950. https://doi.org/10.1038/s41467-023-42671-y  |
| 622 | Harter, D. E. V., Irl, S. D. H., Seo, B., Steinbauer, M. J., Gillespie, R., Triantis, K. A., Fernández-Palacios, |
| 623 | JM., & Beierkuhnlein, C. (2015). Impacts of global climate change on the floras of oceanic                       |
| 624 | islands – Projections, implications and current knowledge. Perspectives in Plant Ecology,                        |
| 625 | Evolution and Systematics, 17(2), 160–183. https://doi.org/10.1016/j.ppees.2015.01.003                           |
| 626 | Hossain, M. A., Kujala, H., Bland, L. M., Burgman, M., & Lahoz-Monfort, J. J. (2019). Assessing the              |
| 627 | impacts of uncertainty in climate-change vulnerability assessments. Diversity and                                |
| 628 | Distributions, 25(8), 1234–1245. https://doi.org/10.1111/ddi.12936   |

| 629 | Inamine, H., Miller, A., Roxburgh, S., Buckling, A., & Shea, K. (2022). Pulse and Press Disturbances       |
|-----|--|
| 630 | Have Different Effects on Transient Community Dynamics. The American Naturalist, 200(4),                   |
| 631 | 571–583. https://doi.org/10.1086/720618  |
| 632 | Jones, H. P., Holmes, N. D., Butchart, S. H. M., Tershy, B. R., Kappes, P. J., Corkery, I., Aguirre-muñoz, |
| 633 | A., Armstrong, D. P., Bonnaud, E., Burbidge, A. A., Campbell, K., Courchamp, F., Cowan, P. E.,             |
| 634 | Cuthbert, R. J., Ebbert, S., Genovesi, P., Howald, G., Keitt, B. S., Kress, S. W., Croll, D. A.            |
| 635 | (2016). Invasive mammal eradication on islands results in substantial conservation gains.                  |
| 636 | Proceedings of the National Academy of Sciences, 1–6.  |
| 637 | https://doi.org/10.1073/pnas.1521179113  |
| 638 | Kreft, H., Jetz, W., Mutke, J., & Barthlott, W. (2010). Contrasting environmental and regional effects     |
| 639 | on global pteridophyte and seed plant diversity. Ecography, 33(2), 408-419.                                |
| 640 | https://doi.org/10.1111/j.1600-0587.2010.06434.x   |
| 641 | Leclerc, C., Courchamp, F., & Bellard, C. (2020). Future climate change vulnerability of endemic island    |
| 642 | mammals. Nature Communications, 11(1), 4943. https://doi.org/10.1038/s41467-020-18740-                     |
| 643 | x  |
| 644 | Leclerc, C., Villéger, S., Marino, C., & Bellard, C. (2020). Global changes threaten functional and        |
| 645 | taxonomic diversity of insular species worldwide. Diversity and Distributions, 26(4), 402–414.             |
| 646 | https://doi.org/10.1111/ddi.13024  |
| 647 | Lerner, H. R. L., Meyer, M., James, H. F., Hofreiter, M., & Fleischer, R. C. (2011). Multilocus Resolution |
| 648 | of Phylogeny and Timescale in the Extant Adaptive Radiation of Hawaiian Honeycreepers.                     |
| 649 | Current Biology, 21(21), 1838–1844. https://doi.org/10.1016/j.cub.2011.09.039                              |
| 650 | Loiseau, N., Mouquet, N., Casajus, N., Grenié, M., Guéguen, M., Maitner, B., Mouillot, D., Ostling, A.,    |
| 651 | Renaud, J., Tucker, C., Velez, L., Thuiller, W., & Violle, C. (2020). Global distribution and              |
| 652 | conservation status of ecologically rare mammal and bird species. Nature Communications,                   |
| 653 | 11(1), 5071. https://doi.org/10.1038/s41467-020-18779-w  |

| 654 | Lomolino, M. V. (1985). Body Size of Mammals on Islands: The Island Rule Reexamined. <i>The American</i>    |
|-----|---|
| 655 | Naturalist, 125(2), 310–316.  |
| 656 | Loreau, M., Barbier, M., Filotas, E., Gravel, D., Isbell, F., Miller, S. J., Montoya, J. M., Wang, S.,      |
| 657 | Aussenac, R., Germain, R., Thompson, P. L., Gonzalez, A., & Dee, L. E. (2021). Biodiversity as              |
| 658 | insurance: From concept to measurement and application. Biological Reviews, 96(5), 2333-                    |
| 659 | 2354. https://doi.org/10.1111/brv.12756   |
| 660 | Manes, S., Costello, M. J., Beckett, H., Debnath, A., Devenish-Nelson, E., Grey, KA., Jenkins, R., Khan,    |
| 661 | T. M., Kiessling, W., Krause, C., Maharaj, S. S., Midgley, G. F., Price, J., Talukdar, G., & Vale, M.       |
| 662 | M. (2021). Endemism increases species' climate change risk in areas of global biodiversity                  |
| 663 | importance. Biological Conservation, 257, 109070.   |
| 664 | https://doi.org/10.1016/j.biocon.2021.109070  |
| 665 | Mantyka-Pringle, C. S., Martin, T. G., & Rhodes, J. R. (2011). Interactions between climate and habitat     |
| 666 | loss effects on biodiversity: A systematic review and meta-analysis. Global Change Biology,                 |
| 667 | n/a-n/a. https://doi.org/10.1111/j.1365-2486.2011.02593.x   |
| 668 | Marino, C., Leclerc, C., & Bellard, C. (2022). Profiling insular vertebrates prone to biological invasions: |
| 669 | What makes them vulnerable? Global Change Biology, 28(3), 1077–1090.  |
| 670 | https://doi.org/10.1111/gcb.15941   |
| 671 | Matthews, T. J., Wayman, J. P., Whittaker, R. J., Cardoso, P., Hume, J. P., Sayol, F., Proios, K., Martin,  |
| 672 | T. E., Baiser, B., Borges, P. A. V., Kubota, Y., dos Anjos, L., Tobias, J. A., Soares, F. C., Si, X.,       |
| 673 | Ding, P., Mendenhall, C. D., Sin, Y. C. K., Rheindt, F. E., Rigal, F. (2023). A global analysis of          |
| 674 | avian island diversity—area relationships in the Anthropocene. Ecology Letters, 26(6), 965—                 |
| 675 | 982. https://doi.org/10.1111/ele.14203  |
| 676 | McCann, K. S. (2000). The diversity-stability debate. <i>Nature</i> , 405(6783), 228–233.                   |
| 677 | https://doi.org/10.1038/35012234  |
| 678 | McConkey, K. R., & Drake, D. R. (2015). Low redundancy in seed dispersal within an island frugivore         |
| 679 | community. AoB PLANTS, 7, plv088. https://doi.org/10.1093/aobpla/plv088                                     |

| 680 | Melo, M., & O'ryan, C. (2007). Genetic differentiation between Príncipe Island and mainland                 |
|-----|---|
| 681 | populations of the grey parrot (Psittacus erithacus), and implications for conservation.                    |
| 682 | Molecular Ecology, 16(8), 1673–1685. https://doi.org/10.1111/j.1365-294X.2006.03128.x                       |
| 683 | Mouillot, D., Villéger, S., Parravicini, V., Kulbicki, M., Arias-González, J. E., Bender, M., Chabanet, P., |
| 684 | Floeter, S. R., Friedlander, A., Vigliola, L., & Bellwood, D. R. (2014). Functional over-                   |
| 685 | redundancy and high functional vulnerability in global fish faunas on tropical reefs.                       |
| 686 | Proceedings of the National Academy of Sciences of the United States of America, 111(38),                   |
| 687 | 13757–13762. https://doi.org/10.1073/pnas.1317625111  |
| 688 | Nyboer, E. A., Lin, HY., Bennett, J. R., Gabriel, J., Twardek, W., Chhor, A. D., Daly, L., Dolson, S.,      |
| 689 | Guitard, E., Holder, P., Mozzon, C. M., Trahan, A., Zimmermann, D., Kesner-Reyes, K., Garilao               |
| 690 | C., Kaschner, K., & Cooke, S. J. (2021). Global assessment of marine and freshwater                         |
| 691 | recreational fish reveals mismatch in climate change vulnerability and conservation effort.                 |
| 692 | Global Change Biology, 27(19), 4799–4824. https://doi.org/10.1111/gcb.15768                                 |
| 693 | Parmakelis, A., Rigal, F., Mourikis, T. P., Balanika, K., Terzopoulou, S., Rego, C., Amorim, I. R., Crespo, |
| 694 | L. C. F., Pereira, F. L., Triantis, K. A., Whittaker, R. J., & Borges, P. A. V. (2015). Comparative         |
| 695 | phylogeography of endemic Azorean arthropods. BMC Evolutionary Biology, 15.                                 |
| 696 | https://api.semanticscholar.org/CorpusID:17414518   |
| 697 | Parravicini, V., Villéger, S., Mcclanahan, T. R., Arias-González, J. E., Bellwood, D. R., Belmaker, J.,     |
| 698 | Chabanet, P., Floeter, S. R., Friedlander, A. M., Guilhaumon, F., Vigliola, L., Kulbicki, M., &             |
| 699 | Mouillot, D. (2014). Global mismatch between species richness and vulnerability of reef fish                |
| 700 | assemblages. Ecology Letters, 17(9), 1101–1110. https://doi.org/10.1111/ele.12316                           |
| 701 | Pavoine, S., & Ricotta, C. (2021). On the relationships between rarity, uniqueness, distinctiveness,        |
| 702 | originality and functional/phylogenetic diversity. Biological Conservation, 263, 109356.                    |
| 703 | https://doi.org/10.1016/j.biocon.2021.109356  |
| 704 | Poljansek, K., Casajus, V. A., Marin, F. M., Artes, V. T., Boca, R., Bonadonna, C., Branco, A.,             |
| 705 | Campanharo, W., De, J. A., De, R. D., Dottori, F., Durrant, T., Estreguil, C., Ferrari, D.,                 |

| 706 | Frischknecht, C., Galbusera, L., Garcia, P. B., Giannopoulos, G., Girgin, S., Wood, M. (2021,                  |
|-----|--|
| 707 | February 23). Recommendations for National Risk Assessment for Disaster Risk Management                        |
| 708 | in EU. JRC Publications Repository. https://doi.org/10.2760/80545  |
| 709 | Price, J. P. (2004). Floristic biogeography of the Hawaiian Islands: Influences of area, environment           |
| 710 | and paleogeography. Journal of Biogeography, 31(3), 487–500.   |
| 711 | https://doi.org/10.1046/j.0305-0270.2003.00990.x   |
| 712 | Ricklefs, R. E. (2017). Historical Biogeography and Extinction in the Hawaiian Honeycreepers. <i>The</i>       |
| 713 | American Naturalist, 190(4), E106–E111. https://doi.org/10.1086/693346   |
| 714 | Riper, C. G. V., & Scott, J. M. (2017). Limiting factors affecting hawaiian native birds.                      |
| 715 | https://api.semanticscholar.org/CorpusID:244917512   |
| 716 | Rocchini, D., Hortal, J., Lengyel, S., Lobo, J. M., Jiménez-Valverde, A., Ricotta, C., Bacaro, G., &           |
| 717 | Chiarucci, A. (2011). Accounting for uncertainty when mapping species distributions: The                       |
| 718 | need for maps of ignorance. Progress in Physical Geography: Earth and Environment, 35(2),                      |
| 719 | 211–226. https://doi.org/10.1177/0309133311399491  |
| 720 | Rodrigues, A. S. L., Gray, C. L., Crowter, B. J., Ewers, R. M., Stuart, S. N., Whitten, T., & Manica, A.       |
| 721 | (2010). A Global Assessment of Amphibian Taxonomic Effort and Expertise. BioScience,                           |
| 722 | 60(10), 798-806. https://doi.org/10.1525/bio.2010.60.10.6  |
| 723 | Roff, D. A. (1990). The Evolution of Flightlessness in Insects. <i>Ecological Monographs</i> , 60(4), 389–421. |
| 724 | https://doi.org/10.2307/1943013  |
| 725 | Rosauer, D., Laffan, S. W., Crisp, M. D., Donnellan, S. C., & Cook, L. G. (2009). Phylogenetic endemism:       |
| 726 | A new approach for identifying geographical concentrations of evolutionary history.                            |
| 727 | Molecular Ecology, 18(19), 4061–4072. https://doi.org/10.1111/j.1365-294X.2009.04311.x                         |
| 728 | Royer-Tardif, S., Boisvert-Marsh, L., Godbout, J., Isabel, N., & Aubin, I. (2021). Finding common              |
| 729 | ground: Toward comparable indicators of adaptive capacity of tree species to a changing                        |
| 730 | climate. Ecology and Evolution, 11(19), 13081–13100. https://doi.org/10.1002/ece3.8024                         |

| /31 | ROZZI, R., LOMOIINO, M. V., Van der Geer, A. A. E., Silvestro, D., Lyons, S. K., Bover, P., Alcover, J. A., |
|-----|---|
| 732 | Benítez-López, A., Tsai, CH., Fujita, M., Kubo, M. O., Ochoa, J., Scarborough, M. E., Turvey, S.            |
| 733 | T., Zizka, A., & Chase, J. M. (2023). Dwarfism and gigantism drive human-mediated                           |
| 734 | extinctions on islands. <i>Science</i> , <i>379</i> (6636), 1054–1059.                                      |
| 735 | https://doi.org/10.1126/science.add8606   |
| 736 | Russell, J. C., & Kueffer, C. (2019). Island Biodiversity in the Anthropocene. <i>Annual Review of</i>      |
| 737 | Environment and Resources, 44(1), 31–60. https://doi.org/10.1146/annurev-environ-101718-                    |
| 738 | 033245  |
| 739 | Santos, M. J., Smith, A. B., Dekker, S. C., Eppinga, M. B., Leitão, P. J., Moreno-Mateos, D., Morueta-      |
| 740 | Holme, N., & Ruggeri, M. (2021). The role of land use and land cover change in climate                      |
| 741 | change vulnerability assessments of biodiversity: A systematic review. Landscape Ecology,                   |
| 742 | 36(12), 3367–3382. https://doi.org/10.1007/s10980-021-01276-w   |
| 743 | Sayol, F., Steinbauer, J., Blackburn, T., Antonelli, A., & Faurby, S. (2021). Anthropogenic extinctions     |
| 744 | conceal widespread evolution of flightlessness in birds. Science Advances, 6(49), eabb6095.                 |
| 745 | https://doi.org/10.1126/sciadv.abb6095  |
| 746 | Schrader, J., Wright, I. J., Kreft, H., & Westoby, M. (2021). A roadmap to plant functional island          |
| 747 | biogeography. Biological Reviews, 96(6), 2851–2870. https://doi.org/10.1111/brv.12782                       |
| 748 | Silva Rocha, B., Jamoneau, A., Logez, M., Laplace-Treyture, C., Reynaud, N., & Argillier, C. (2024).        |
| 749 | Measuring biodiversity vulnerability in French lakes – The IVCLA index. Science of The Total                |
| 750 | Environment, 908, 168205. https://doi.org/10.1016/j.scitotenv.2023.168205                                   |
| 751 | Soares, F. C., de Lima, R. F., Palmeirim, J. M., Cardoso, P., & Rodrigues, A. S. L. (2022). Combined        |
| 752 | effects of bird extinctions and introductions in oceanic islands: Decreased functional diversity            |
| 753 | despite increased species richness. Global Ecology and Biogeography, 31(6), 1172–1183.                      |
| 754 | https://doi.org/10.1111/geb.13494   |

- 755 Solomon, S., Qin, D., M. Manning, Z. Chen, M. Marquis, K.B. Averyt, M. Tignor, & H.L. Miller. (2007). 756 Contribution of Working Group I to the Fourth Assessment Report of the Intergovernmental 757 Panel on Climate Change (p. 996). IPCC. Soria, C. D., Pacifici, M., Di Marco, M., Stephen, S. M., & Rondinini, C. (2021). COMBINE: a coalesced 758 759 mammal database of intrinsic and extrinsic traits. Ecology, 102(6), e03344. https://doi.org/10.1002/ecy.3344 760 761 Sousa, A., Alves, F., Arranz, P., Dinis, A., Fernandez, M., González García, L., Morales, M., Lettrich, M., 762 Encarnação Coelho, R., Costa, H., Capela Lourenço, T., Azevedo, N. M. J., & Frazão Santos, C. 763 (2021). Climate change vulnerability of cetaceans in Macaronesia: Insights from a trait-based 764 assessment. Science of The Total Environment, 795, 148652. 765 https://doi.org/10.1016/j.scitotenv.2021.148652 766 Tessarolo, G., Ladle, R. J., Lobo, J. M., Rangel, T. F., & Hortal, J. (2021). Using maps of biogeographical ignorance to reveal the uncertainty in distributional data hidden in species distribution 767 768 models. *Ecography*, 44(12), 1743–1755. https://doi.org/10.1111/ecog.05793 Thuiller, W., Guéguen, M., Renaud, J., Karger, D. N., & Zimmermann, N. E. (2019). Uncertainty in 769 770 ensembles of global biodiversity scenarios. Nature Communications, 10(1), Article 1. 771 https://doi.org/10.1038/s41467-019-09519-w 772 Thurman, L. L., Stein, B. A., Beever, E. A., Foden, W., Geange, S. R., Green, N., Gross, J. E., Lawrence, D. J., LeDee, O., Olden, J. D., Thompson, L. M., & Young, B. E. (2020). Persist in place or shift 773 in space? Evaluating the adaptive capacity of species to climate change. Frontiers in Ecology 774 775 and the Environment, 18(9), 520-528. https://doi.org/10.1002/fee.2253
- Tobias, J. A., Sheard, C., Pigot, A. L., Devenish, A. J. M., Yang, J., Neate-Clegg, M. H. C., Alioravainen,
   N., Weeks, T. L., Barber, R. A., Walkden, P. A., MacGregor, H. E. A., Jones, S. E. I., Vincent, C.,
   Phillips, A. G., Marples, N. M., Montaño-Centellas, F., Leandro-Silva, V., Claramunt, S., Darski,
   B., ... Schleunning, M. (2021). AVONET: morphological, ecological and geographical data for
   all birds. *Ecology Letters*, 25(March), 581–597. https://doi.org/10.1111/ele.13898

| 781 | Troudet, J., Grandcolas, P., Blin, A., Vignes-Lebbe, R., & Legendre, F. (2017). Taxonomic bias in        |
|-----|--|
| 782 | biodiversity data and societal preferences. Scientific Reports, 7(1), 9132.                              |
| 783 | https://doi.org/10.1038/s41598-017-09084-6   |
| 784 | UN High Level Panel. (2022). United Nations High Level Panel on the Development of a                     |
| 785 | Multidimensional Vulnerability Index Interim Report. https://www.oecd.org/dac/financing-                 |
| 786 | sustainable-   |
| 787 | UNFCCC. (2007). VULNERABILITY AND ADAPTATION TO CLIMATE CHANGE IN SMALL ISLAND                           |
| 788 | DEVELOPING STATES. the secretariat of the United Nations Framework Convention on                         |
| 789 | Climate Change.  |
| 790 | https://unfccc.int/files/adaptation/adverse_effects_and_response_measures_art_48/application             |
| 791 | tion/pdf/200702_sids_adaptation_bg.pdf   |
| 792 | Ureta, C., Ramírez-Barrón, M., Sánchez-García, E. A., Cuervo-Robayo, A. P., Munguía-Carrara, M.,         |
| 793 | Mendoza-Ponce, A., Gay, C., & Sánchez-Cordero, V. (2022). Species, taxonomic, and                        |
| 794 | functional group diversities of terrestrial mammals at risk under climate change and land-               |
| 795 | use/cover change scenarios in Mexico. Global Change Biology, 28(23), 6992–7008.                          |
| 796 | https://doi.org/10.1111/gcb.16411  |
| 797 | Violle, C., Thuiller, W., Mouquet, N., Munoz, F., Kraft, N. J. B., Cadotte, M. W., Livingstone, S. W., & |
| 798 | Mouillot, D. (2017). Functional Rarity: The Ecology of Outliers. Trends in Ecology & Evolution,          |
| 799 | 32(5), 356–367. https://doi.org/10.1016/j.tree.2017.02.002   |
| 800 | Vitousek, P. M., D'Antonio, C. M., Loope, L. L., Rejmanek, M., & Westbrooks, R. (1997). Introduced       |
| 801 | species: A significant component of Human-caused global environmental change The scope                   |
| 802 | and distribution of invasions. Journal of Ecology, 21, 1–16.   |
| 803 | Warren, D. L., Glor, R. E., & Turelli, M. (2008). Environmental niche equivalency versus conservatism:   |
| 804 | Quantitative approaches to niche evolution. Evolution; International Journal of Organic                  |
| 805 | Evolution, 62(11), 2868–2883. https://doi.org/10.1111/j.1558-5646.2008.00482.x                           |

| 806                      | White, T. A., & Searle, J. B. (2007). Genetic diversity and population size: Island populations of the      |
|--------------------------|---|
| 807                      | common shrew, Sorex araneus. Mol Ecol, 16(10), 2005–2016. PubMed.   |
| 808                      | https://doi.org/10.1111/j.1365-294x.2007.03296.x  |
| 809                      | Whittaker, R. J., Rigal, F., Borges, P. A. V., Cardoso, P., Terzopoulou, S., Casanoves, F., Pla, L.,        |
| 810                      | Guilhaumon, F., Ladle, R. J., & Triantis, K. A. (2014). Functional biogeography of oceanic                  |
| 811                      | islands and the scaling of functional diversity in the Azores. Proceedings of the National                  |
| 812                      | Academy of Sciences, 111(38), 13709 LP — 13714.   |
| 813                      | Zizka, A., Onstein, R. E., Rozzi, R., Weigelt, P., Kreft, H., Steinbauer, M. J., Bruelheide, H., & Lens, F. |
| 814                      | (2022). The evolution of insular woodiness. Proceedings of the National Academy of Sciences,                |
| 815                      | 119(37), e2208629119. https://doi.org/10.1073/pnas.2208629119   |
| 816<br>817<br>818<br>819 |   |
| 820                      |   |
| 821                      |   |
| 822                      |   |
| 823                      |   |
| 824                      |   |
| 825                      |   |
| 826                      |   |
| 827                      |   |
| 828                      |   |
| 829                      |   |
| 830                      |   |

## **Supplementary material:**

831 832

833

834

835

836

837

838

839

840

841

842

843

844

845

846

847

848

849

850

851

852

853

854

## **Supplementary appendix 1:**

Islands as past, current, and future epicentres of extinctions: The IUCN Red List documented nearly 1,000 species as globally extinct or extinct in the wild (IUCN Red list access 04/07/2023). Among the documented extinctions, the overwhelming majority occurred on oceanic or continental islands. Extinction hotspots are located in the Oceanic realm, followed by the Afrotropical realm (e.g., Madagascar) and the Nearctic. Among reported extinctions in the IUCN Red List, the large majority are represented by animal species (>80%), followed by plants. Most of these species have gone extinct because of invasive alien species, but a large number have also concurrently been harmed by wildlife exploitation and/or land use change due to cultivation (Leclerc et al., 2018; Maxwell et al., 2016). In fact, extinct species within insular regions have faced, on average, four threats (i.e., biological invasions, wildlife exploitation, cultivation, and habitat modifications), and the exposure to threats for birds, mammals, amphibians, reptiles, freshwater fish, plants, arthropods, and gastropods continue to increase, with an average of ten threats being faced by threatened species (Leclerc et al., 2018). Moreover, because insular biota makes a disproportionate contribution to global biodiversity in relation to their surface area (Fernández-Palacios et al., 2021), the risk for future extinctions is high. The first extinction due to climate change already occurred on an island near Papua New Guinea (Fulton, 2017). In 2016, Melomys rubicola (also called the Bramble Cay melomys) became the very first documented extinction due to climate change (Waller et al., 2017). This small rodent was endemic to a low-island of Bramble Cay in Australia and was periodically recorded from 1978 to 2009. Seven years after its last observation in the field and numerous efforts to trap the species the years after, the species was officially declared extinct. Although only symbolic, compared to a thousand of species already extinct since 1500, this is one of the first documented victims of climate change as a major factor of near-future extinctions in our contemporary history. Unfortunately, under future climate change, we can expect that islands will continue to contribute disproportionately to the loss of biodiversity, given that the overwhelming majority of critically endangered species are endemic to island systems, and their inherent characteristics make them extremely vulnerable to global change drivers

**Table S1:** Traits and characteristics from population to the community that may be included in the island biota vulnerability framework (IBVF) with examples of sources where to calculate or when directly available.

|                            | Traits  | Level      | Availability   |  |
|----------------------------|---|------------|--|--|
| Population dynamic         | Growth rate   | population | Living planet database<br>(vertebrates), FishGlob<br>(fishes), German vegetation<br>(plants) |  |
|                            | Population size   | population | CoralTraits (corals), BIOTime (multiple taxa)  |  |
|                            | Reproductive strategies                                 | species    | GIFT (plants),<br>COMPADRE, GARD<br>(reptiles)   |  |
|                            | Mating system   | species    |  |  |
| Reproduction               | Clutch size species                                     |            | GARD (reptiles),<br>AmphiBio (amphibians),<br>COMBINE (mammals),<br>AVONET (birds)           |  |
|                            | Fecundity   | species    | GARD (reptiles),<br>AmphiBio (amphibians)  |  |
|                            | Temperature-dependent sex determination                 | species    |  |  |
|                            | Generation time   | species    | Amniote (vertebrates),<br>CoralTraits (corals),<br>COMPADRE (plants)                         |  |
|                            | Parental care (time)                                    | species    | Amniote (vertebrates)  |  |
|                            | Life-cycle  | species    | TRY (plants), GIFT (plants)  |  |
| Life-history               | Life stage  | population | FishBase (fishes),<br>CoralTraits (corals),<br>Amphibio (amphibians)                         |  |
| Life-mistory               | Dependence on intraspecific interactions                | species    | CoralTraits (corals)   |  |
|                            | Body dimension (mass/size?)                             | species    | Amniote (vertebrates), TRY (Plants), GIFT (plants), COMBINE (mammals), AVONET (birds).       |  |
|                            | Clonality   | species    | GIFT (plants),<br>COMPADRE (plants)  |  |
|                            | Phenotypic plasticity                                   | population | Noble et al 2018 (reptiles)  |  |
| Acclimatation or evolution | Genetic diversity                                       | population | Miraldo et al. 2016<br>(mammals)   |  |
|                            | Life span   | species    | AmphiBio (amphibians),   |  |
|                            | Climatic niche breadth                                  | species    | GARD (reptiles), GBIF (plants), BIEN (plants)  |  |
|                            | Endemicity  | species    | Global Species Database (vertebrates, invertebrates, plants)                                 |  |
| Ecological                 | Habitat specialization                                  | species    | IUCN   |  |
| characteristics            | Habitat condition                                       | species    | IUCN   |  |
|                            | Diet breadth  | species    | GARD (reptiles),<br>AmphiBio (amphibians),   |  |
|                            | Nocturnality  | species    | GARD (reptiles), Amphibio (amphibians)   |  |
|                            | Dependence on water for reproduction, foraging, shelter | species    | Amphibio (amphibians)  |  |

|                             | Competitive ability                     | species      |   |
|-----------------------------|---|--------------|---|
|                             | Dispersal capacity                      | species      | AVONET (birds), TRY (plants), GIFT (plants)   |
| 3.5                         | Site fidelity                           | species      |   |
| Movement and mobility       | Migration frequency and distance        | species      | AVONET (birds)  |
|                             | Flight efficiency                       | species      | AVONET (birds)  |
|                             | Number of insular populations           | population   |   |
|                             | Home range                              | population   | HomeRange (mammals)   |
| D: 4 11 41                  | Geographic rarity                       | population   |   |
| Distribution                | Low local abundance                     | island level |   |
|                             | Range size                              | population   | IUCN  |
|                             | Level of disharmony                     | community    |   |
|                             | Functional redundancy                   | community    | TRY and GIFT(plants)  |
|                             | Phylogenetic diversity                  | community    | GIFT, WCVP (plants)   |
| <b>Community properties</b> | Richness                                | community    | GIFT, WCVP (plants)   |
|                             | Intactness                              | ecosystem    |   |
|                             | Phylogenetic endemism                   | community    | Vertlife (vertebrates),<br>PhylomeDB (plants), GIFT,<br>WCVP (plants)   |
|                             | Behaviour (aggressiveness)              | species      |   |
|                             | Dwarfism-gigantism                      | species      | Meiri et al 2008<br>(mammals), Meiri 2007<br>(reptiles), Benítez-López et<br>al. 2021, Rozzi et al. 2023<br>GIFT (plants) |
| Inherent vulnerability      | Vulnerability to diseases               | species      | GABiP (amphibians)  |
|                             |   |              | try-db.org (plants), Le   |
|                             | Tolerance to drought                    |              | Galliard et al. 2021 (reptiles)   |
|                             | Tolerance to drought  Tolerance to fire | species      |   |

900

901 902

## **References:**

- Benítez-López, A., Santini, L., Gallego-Zamorano, J. *et al.* The island rule explains consistent patterns of body size evolution in terrestrial vertebrates. *Nat Ecol Evol* **5**, 768–786 (2021).
- 905 https://doi.org/10.1038/s41559-021-01426-y
- Broekman, M. J. E., Hoeks, S., Freriks, R., Langendoen, M. M., Runge, K. M., Savenco, E.,
- 907 ... & Tucker, M. A. (2023). HomeRange: A global database of mammalian home
- 908 ranges. Global Ecology and Biogeography, 32(2), 198-205.
- Brown, M.J.M., Walker, B.E., Black, N., Govaerts, R.H.A., Ondo, I., Turner, R. and Nic
- 910 Lughadha, E. (2023), rWCVP: a companion R package for the World Checklist of Vascular
- 911 Plants. New Phytol, 240: 1355-1365. <a href="https://doi.org/10.1111/nph.1891">https://doi.org/10.1111/nph.1891</a>9

912

- Denelle, P., Weigelt, P., & Kreft, H. (2023). GIFT—An R package to access the Global
- 914 Inventory of Floras and Traits. Methods in Ecology and Evolution, 14, 2738–2748.
- 915 https://doi.org/10.1111/2041-210X.14213

916

- Dornelas M, Antão LH, Moyes F, Bates, AE, Magurran, AE, et al. BioTIME: A database of
- 918 biodiversity time series for the Anthropocene. *Global Ecol Biogeogr.* 2018; 27:760 786.
- 919 https://doi.org/10.1111/geb.12729
- 920 Le Galliard, Jean- François, et al. "A worldwide and annotated database of evaporative water
- loss rates in squamate reptiles." Global Ecology and Biogeography 30.10 (2021): 1938-1950.

922

- 923 Maitner BS, Boyle B, Casler N, et al. The bien r package: A tool to access the Botanical
- 924 Information and Ecology Network (BIEN) Database. *Methods Ecol Evol.* 2018; 9: 373–379.
- 925 https://doi.org/10.1111/2041-210X.12861

926

- 927 Salguero-Gómez, R., Jones, O.R., Archer, C.R., Buckley, Y.M., Che-Castaldo, J., Caswell,
- 928 H., Hodgson, D., Scheuerlein, A., Conde, D.A., Brinks, E., de Buhr, H., Farack, C.,
- 929 Gottschalk, F., Hartmann, A., Henning, A., Hoppe, G., Römer, G., Runge, J., Ruoff, T., Wille,
- J., Zeh, S., Davison, R., Vieregg, D., Baudisch, A., Altwegg, R., Colchero, F., Dong, M., de
- 931 Kroon, H., Lebreton, J.-D., Metcalf, C.J.E., Neel, M.M., Parker, I.M., Takada, T., Valverde,
- T., Vélez-Espino, L.A., Wardle, G.M., Franco, M. and Vaupel, J.W. (2015), The compadre
- Plant Matrix Database: an open online repository for plant demography. J Ecol, 103: 202-218.
- 934 https://doi.org/10.1111/1365-2745.12334

935

- 936 Soria, C. D., M. Pacifici, M. Di Marco, S. M. Stephen, and C. Rondinini. 2021. COMBINE: a
- 937 coalesced mammal database of intrinsic and extrinsic traits. *Ecology*
- 938 102(6):e03344. 10.1002/ecy.3344
- 839 Kattge J, Bönisch G, Díaz S, et al. TRY plant trait database enhanced coverage and open
- 940 access. Glob Change Biol. 2020; 26: 119–188. https://doi.org/10.1111/gcb.14904

941

- Noble, Daniel WA, et al. "A comprehensive database of thermal developmental plasticity in
- 943 reptiles." Scientific Data 5.1 (2018): 1-7.

944

- Meiri, Shai, Natalie Cooper, and Andy Purvis. "The island rule: made to be broken?."
- Proceedings of the Royal Society B: Biological Sciences 275.1631 (2008): 141-148.

947

- 948 Meiri, Shai. "Size evolution in island lizards." Global Ecology and Biogeography 16.6 (2007):
- 949 702-708.
- Position Roberto Rozzi et al, Dwarfism and gigantism drive human-mediated extinctions on
- 951 islands.Science **379**, 1054-1059(2023).DOI: 10.1126/science.add8606
- Tobias, J.A., Sheard, C., Pigot, A.L., Devenish, A.J.M., Yang, J., Sayol, F., et al. (2022)
- 953 AVONET: morphological, ecological and geographical data for all birds. *Ecology Letters*, 25,
- 954 581–597. https://doi.org/10.1111/ele.13898

955

956

**Table S2:** An overview of studies assessing the vulnerability of species to climate change. This is a non-exhaustive summary that focuses on studies that have been conducted recently, use trait-based approaches, and include a full assessment of vulnerability with exposure, sensitivity, and adaptive capacity markers.

| Study                           | Ecosystems               | Spatial coverage        | Taxa   | n      | Main findings  |
|---------------------------------|--------------------------|-------------------------|--|--------|--|
| (Barry et al., 2023)            | Freshwater               | Ireland                 | Fish   | 32     | All species were vulnerable to some effect of climate change with cold water species more vulnerable to climate change than warm water species.  |
| (Boyce et al., 2022)            | Marine                   | Global                  | Multiple groups<br>(animals, plants,<br>chromists,<br>protozoans, and<br>bacteria) | 24,975 | Almost 90% of all species are at high or critical risk under high emissions, with exploited species in low-income countries with heavy dependence on fisheries at greatest risk.   |
| (Nyboer et al., 2021)           | Freshwater /<br>Marine   | Global                  | Fish   | 415    | Over 20% of recreationally fishes are vulnerable under a high emission scenario, with 72% of vulnerable freshwater fish and 33% of vulnerable diadromous fish being without conservation effort, compared to only 19% for vulnerable marine species.       |
| (Vaz-Canosa et al., 2023)       | Terrestrial              | Uruguay                 | Amphibians<br>Reptiles   | 112    | 14.6% of amphibians and 10.9% of reptiles were identified as highly vulnerable to climate change.  |
| (Bueno-Pardo et al., 2021)      | Marine                   | Portugal                | Fish<br>Invertebrates  | 74     | Under the RCP 8.5 scenario, only two species were classified as at very high vulnerability. Overall vulnerability scores were low, likely due to the high adaptive capacity of species from temperate ecosystems.  |
| (Leclerc et al., 2020)          | Terrestrial –<br>insular | Global                  | Mammals  | 873    | All islands have some degree of vulnerability to future climate change, especially those in the Pacific Ocean. Among endemic mammals, those with long generation times and high food specialization are predicted to be most vulnerable to climate change. |
| (Ramírez-Bautista et al., 2020) | Terrestrial              | Oaxaca State,<br>Mexico | Rodents  | 55     | Under the higher impact (MPI-RCP 8.5) climate scenarios, some level of threat was predicted for all species assessed, with 4 species predicted to be highly vulnerable.  |

## **References:**

Barry, J., Radinger, J., Connor, L., Coyne, J., Kelly, S., & Kelly, F. L. (2023). The vulnerability of
 Ireland's freshwater fish to climate change. *Fisheries Management and Ecology*, 30(2), 142–
 https://doi.org/10.1111/fme.12606

- Boyce, D. G., Tittensor, D. P., Garilao, C., Henson, S., Kaschner, K., Kesner-Reyes, K., Pigot, A.,
- 970 Reyes, R. B., Reygondeau, G., Schleit, K. E., Shackell, N. L., Sorongon-Yap, P., & Worm, B.
- 971 (2022). A climate risk index for marine life. *Nature Climate Change*, 12(9), 854–862.
- 972 https://doi.org/10.1038/s41558-022-01437-y
- 973 Bueno-Pardo, J., Nobre, D., Monteiro, J. N., Sousa, P. M., Costa, E. F. S., Baptista, V., Ovelheiro, A.,
- Vieira, V. M. N. C. S., Chícharo, L., Gaspar, M., Erzini, K., Kay, S., Queiroga, H., Teodósio,
- 975 M. A., & Leitão, F. (2021). Climate change vulnerability assessment of the main marine
- ommercial fish and invertebrates of Portugal. *Scientific Reports*, 11(1), 2958.
- 977 https://doi.org/10.1038/s41598-021-82595-5
- 978 Fernández-Palacios, J. M., Kreft, H., Irl, S. D. H., Norder, S., Ah-Peng, C., Borges, P. A. V., Burns, K.
- 979 C., de Nascimento, L., Meyer, J.-Y., Montes, E., & Drake, D. R. (2021). Scientists' warning –
- The outstanding biodiversity of islands is in peril. *Global Ecology and Conservation*, 31,
- 981 e01847. https://doi.org/10.1016/j.gecco.2021.e01847
- 982 Fulton, G. R. (2017). The Bramble Cay melomys: The first mammalian extinction due to human-
- 983 induced climate change. *Pacific Conservation Biology*, 23(1), 1–3.
- Leclerc, C., Courchamp, F., & Bellard, C. (2018). Insular threat associations within taxa worldwide.
- 985 *Scientific Reports*, 8(8), 6393. https://doi.org/10.1038/s41598-018-24733-0
- 986 Leclerc, C., Courchamp, F., & Bellard, C. (2020). Future climate change vulnerability of endemic
- 987 island mammals. *Nature Communications*, 11(1), 4943. https://doi.org/10.1038/s41467-020-
- 988 18740-x
- 989 Maxwell, S. L., Fuller, R. A., Brooks, T. M., & Watson, J. E. M. (2016). Biodiversity: The ravages of
- 990 guns, nets and bulldozers. *Nature*, 536, 143–145. https://doi.org/10.1038/536143a.
- 991 Nyboer, E. A., Lin, H.-Y., Bennett, J. R., Gabriel, J., Twardek, W., Chhor, A. D., Daly, L., Dolson, S.,
- Guitard, E., Holder, P., Mozzon, C. M., Trahan, A., Zimmermann, D., Kesner-Reyes, K.,
- 993 Garilao, C., Kaschner, K., & Cooke, S. J. (2021). Global assessment of marine and freshwater
- recreational fish reveals mismatch in climate change vulnerability and conservation effort.
- 995 *Global Change Biology*, 27(19), 4799–4824. https://doi.org/10.1111/gcb.15768

| 996  | Ramirez-Bautista, A., Thorne, J. H., Schwartz, M. W., & Williams, J. N. (2020). Trait-based climate    |
|------|--|
| 997  | vulnerability of native rodents in southwestern Mexico. Ecology and Evolution, 10(12), 5864            |
| 998  | 5876. https://doi.org/10.1002/ece3.6323  |
| 999  | Vaz-Canosa, P., Laufer, G., Borteiro, C., Baldo, D., Prigioni, C., & Soutullo, A. (2023). Expert-based |
| 1000 | assessment of the climate change vulnerability of amphibians and reptiles of Uruguay.                  |
| 1001 | Environmental Conservation, 50(1), 12–21. https://doi.org/DOI:   |
| 1002 | 10.1017/S0376892922000418  |
| 1003 | Waller, N. L., Gynther, I. C., Freeman, A. B., Lavery, T. H., & Leung, L. KP. (2017). The Bramble      |
| 1004 | Cay melomys Melomys rubicola (Rodentia: Muridae): A first mammalian extinction caused                  |
| 1005 | by human-induced climate change? Wildlife Research, 44(1), 9–21.                                       |
| 1006 |  |
| 1007 |  |
| 1008 |  |
| 1009 |  |
| 1010 |  |
| 1011 |  |
| 1012 |  |
| 1013 |  |
| 1014 |  |
| 1015 |  |
| 1016 |  |
| 1017 |  |
| 1018 |  |
| 1019 |  |
| 1020 |  |
| 1021 |  |
| 1022 |  |
| 1023 |  |
| 1024 |  |
| 1025 |  |
| 1026 |  |