

Protecting Biodiversity (in All Its Complexity): New Models and Methods

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1 Protecting biodiversity (in all its complexity): new models and methods

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- 19
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- 23

25 Abstract

26 We are facing a biodiversity crisis at the same time as we are acquiring an unprecedented view of the 27 world's biodiversity. Vast new datasets (e.g. species distributions, traits, phylogenies, and interaction 28 networks) hold knowledge to better comprehend the depths of biodiversity change, reliably anticipate 29 these changes, and inform conservation actions. To harness this information for conservation, we need to 30 integrate the largely independent fields of biodiversity modelling and conservation. We highlight new 31 developments in each respective field, early examples of how they are being brought together, and ideas 32 for a future synthesis so conservation decisions can be made with a fuller awareness of the biodiversity at 33 stake.

34

35 Main Text

36

37 We need greater diversity in biodiversity conservation

38 Aside from a small fraction of well-known species with established conservation status, most of 39 the world's biodiversity is poorly integrated into conservation efforts. Even for iconic clades 40 such as birds and mammals, only a subset of species are considered in conservation despite a 41 continuously refined understanding of the ecology and evolution of these clades[1,2]. Lesser 42 known taxa are nearly always overlooked[3] even those most at risk of extinction[4]. First, we 43 need to expand the focus of biodiversity in conservation programs to consider more of the tree of 44 life, even taxa without names[5,6], especially as increasing evidence suggests 'invisible' 45 diversity is important to ecosystem functioning [7]. Second, we must consider the role of species in ecosystems[8]. Biodiversity is not simply the mere sum of species, but includes complex 46 47 interaction networks with fluxes of energy and nutrients[9,10]. Even conservation efforts for

48 focal charismatic species would benefit from considering the other species enabling their

49 existence. Biodiversity losses impact all facets of biodiversity (species, genetic diversity,

50 functional diversity, phylogenetic diversity, see Glossary) and spatial extents ranging from 51 local assemblages to the global pool of biodiversity. Considering these dimensions is essential to 52 prevent widespread biodiversity loss.

53

54 Conservation evaluation often falls short for under-represented biodiversity

55 Simplified metrics likely miss many types of biodiversity

56 Despite scientific recognition of the importance of biodiversity across multiple facets, most

57 planning and on-the-ground conservation still focus on a simplified representation of biodiversity

such as the amount or quality of habitat. Habitat-based metrics are used more often in

59 conservation than in ecology[11], and are often the basis of conservation offsetting, wetland

60 mitigation, conservation easements, conservation and management of listed species (alongside

61 population abundances, connectivity, and threat information when available), and are used at a

62 regional extent in coordinated reserve designs (e.g. Natura 2000,

63 https://ec.europa.eu/environment/nature/natura2000/index_en.htm). Reliance on habitat metrics

64 can overlook important biodiversity that falls outside of commonly held notions of high-quality

habitat[12], and these metrics do not necessarily represent population persistence[11]. While the

66 use of habitat metrics certainly has follow-on benefits to some diversity, its surrogacy value for

67 even well-known species is unclear[13], and unknown for many taxa and for different

68 **biodiversity facets**.

70 In cases where multiple species are considered together in applied conservation, they are most 71 often represented as species richness (the count of species in areas)[11]. The use of species 72 richness cannot address the known biases in extinction risk towards certain clades (e.g. 73 amphibians[14]) and functional types (e.g. top predators). While species richness and other 74 species-based metrics can be an effective surrogate for phylogenetic diversity[10] or functional 75 diversity[15], its surrogacy value depends on the conservation scenario[16] and tends to decrease 76 in scenarios where resources are limited (i.e. few sites available for protection)[17]. We have 77 very limited evidence for surrogacy value of different biodiversity facets outside of iconic animal 78 groups (e.g. does above ground animal richness relate to soil microbial diversity?). Finally, how 79 species interact within ecosystems is almost never considered in applied conservation despite 80 links to the important conservation goals of ecosystem functioning and resilience[9].

81

82 Independent conservation efforts fail to protect the world's biodiversity

83 Safeguarding multiple facets of biodiversity—especially for the purposes of preventing 84 extinction, protecting our biological heritage, and retaining future options for society-requires 85 an understanding of the global biodiversity pool, arguably more than for other conservation aims 86 (e.g. managing local ecosystems or protecting ecosystem services)[18]. But conservation actions 87 are typically local or regional, which can lead to unintended outcomes for global 88 biodiversity[19]. First, species are declining globally[20], but not necessarily locally[21–23], 89 leading to a homogenization of biodiversity where different areas have increasingly similar 90 biodiversity[20]. Second, conservation priorities established for local species assemblages[16] or 91 even for species at a national scale[19] differ from global biodiversity priorities[16,19]. Lastly, 92 the emphasis on protecting large patches and connected habitats (e.g. species corridor 93 initiatives), which is based on research on metapopulations and conservation genetics, risks

overlooking important biodiversity (e.g. endemic species, highly functional communities) that
occur in small and isolated habitat patches [24]. A more spatially-coordinated approach is needed
that recognizes how local conservation efforts impact the regional and global biodiversity pool.

98 The potential to bring together biodiversity modelling and conservation

99 The fields of ecology and conservation planning already consider the spatial extents necessary 100 for this coordination, but they do so in slightly different ways. In systematic conservation 101 planning (SCP), biodiversity features (e.g. species distributions) are considered in planning units. 102 Planning units are compared across the entire area of interest using the principles of spatial 103 complementarity and irreplaceability. In ecology, a similar hierarchy can be found in the 104 delineation of α -(within-site or within-ecosystem diversity), β -(differentiation of sites in a 105 region; e.g. compositional turnover), and γ -diversity (diversity of the entire region). We refer to 106 γ -diversity as the total set of biodiversity in the area of interest, which could have a regional or 107 global extent, to better align with conservation planning. Both fields are progressing 108 independently toward a multi-scale, multi-facetted view of biodiversity (Figure 1). Now is the 109 time to integrate these fields to rapidly expand the amount of biodiversity data that reaches 110 conservation practice (Figure 2).

111

112 Advances in biodiversity models

While steady progress has been made on addressing the major biodiversity shortfalls with increased data collection (e.g. citizen science[25], environmental DNA) and global initiatives to compile data on species occurrences (e.g. GEO-BON initiative; geobon.org, GBIF; gbif.org), traits (e.g. TRY;www.try-db.org), genetic and phylogenetic data (e.g. GenBank), and species 117 interaction data (e.g. GloBI; globalbioticinteractions.org), major shortfalls remain[26].

118 Biodiversity models are essential tools to rapidly overcome these shortfalls. We consider

- 119 **biodiversity models** to be any model that makes predictions for biodiversity (including species
- 120 occurrences or abundances, traits, phylogenetic placement, or species interactions). Here, we
- 121 highlight two model classes: (1) imputation models for missing biodiversity data and (2) spatial
- 122 biodiversity models for predicting taxa or other biodiversity facets for unsampled locations,
- 123 which often serve as an input for conservation planning (Figure 1).
- 124

125 Imputation models

126 Imputation models are needed to fill in the many gaps in biodiversity datasets. For example, they 127 can generate missing trait values for taxa[27], which can be estimated from phylogenies, 128 assuming traits have a general pattern of conserved evolution [28] or through multivariate 129 imputation that does not rely on phylogenies[29]. Increasingly sophisticated approaches are also 130 being used to generate phylogenies when some gene sequences are missing[30], and to combine 131 multiple datasets into aggregated phylogenies either based on the phylogenies themselves ('super 132 trees' or preferably from a combined gene matrix[31]. There is also growing interest in 133 predicting interactions between species based on their trait attributes (e.g. predator-prey trait 134 matching[32]) and spatial co-occurrence data[33]. Predicting species interactions across many 135 species or at multiple locations is difficult with severely limited data, but can be addressed by 136 combining empirical data and expert knowledge[34].

137

While imputation models are greatly expanding the coverage of biodiversity datasets, challengesremain. Data collection is systematically geographically and taxonomically biased, which could

140 bias predictions, especially for rare taxa. We are increasingly relying on the same datasets for 141 different types of analyses, so there is also the danger of circularity if those analyses are later 142 combined (e.g. using the phylogeny to predict trait values and later combining phylogeny and 143 traits in spatial models). We need more robust tests for the impact of these biases, and to ensure 144 that uncertainty is propagated from the original data sources to all subsequent analyses. A partial 145 solution is to integrate methods and create a transparent and documented pipeline of analyses 146 (Figure 2), but we still must address the underlying causes of bias and use this knowledge to 147 guide future collection.

148

149

151

150 Spatial Biodiversity models

152 The Wallacean shortfall (inadequate knowledge of geographic distributions) greatly hinders 153 progress in identifying conservation priorities[26]. This shortfall is particularly problematic at 154 local scales-it is easier to define a regional or national set of species than to pinpoint the precise 155 distributions of species at fine spatial scales[26]. Biodiversity models are essential tools to 156 address this shortfall. In many cases, they can provide continuous coverage of species 157 distributions at a resolution relevant for conservation purposes[35]. The most common examples 158 are single species distribution models (i-SDMs; Fig. I), which have been and will continue to be 159 invaluable tools for conservation applications [35]. Advanced models can additionally 160 accommodate multiple facets (e.g. traits and phylogenies) and integrate elements of α - and β -161 diversity (Box 1). A notable advance is the ability to combine a top-down approach focused on 162 an entire species assemblage with a bottom-up approach focused on single species ('sideways' 163 biodiversity models; Box 1, Box 2). One such model, Joint Species Distribution Models (j-164 SDMs, reviewed in [36]), offers potential improvements by modeling all species simultaneously

[37,38], enabling conditional predictions [39–41], and has a flexible hierarchy amenable to
adding other information such as traits[37], phylogenies[42], geographic space[43], temporal
data[44], and detection bias[45], which impact multifaceted biodiversity estimates[46]. Models
that can account for species interactions[47] are an exciting arena for future developments.

169

170 A key advantage of these advanced biodiversity models is the ability to process highly 171 multidimensional datasets (e.g. many taxa, sites, and environmental variables). Combining 172 multiple, large biodiversity datasets can quickly result in a computationally intractable problem. 173 Solutions include dimensionality reduction in a Bayesian framework[36,48] or via machine 174 learning[49]. However, what these approaches gain in flexibility may come at a cost in terms of 175 generality, and the relationships derived between explanatory variables and diversity metrics are 176 often empirical with little to no grounding in ecological theory. An important future research 177 avenue will be to rigorously test new biodiversity models to evaluate their capacity for aiding 178 ecological inference and for predicting to new situations (e.g. novel climates or altered 179 ecosystems).

180

181 Advances in conservation evaluation of multifaceted biodiversity

While the methods described above could offer a richer representation of biodiversity and improved predictive abilities, identifying conservation solutions requires more than just biodiversity predictions. These predictions must be translated to meaningful outputs that can be used to meet defined objectives (e.g. protected area targets) or provide forecasts for decision makers. In systematic conservation planning (SCP)[50], priorities are established that efficiently meet goals for representation of biodiversity under a range of spatial or other constraints (e.g. in a **spatial prioritization**). Targets can be set on diversity (e.g. 10% of each species range

189 protected[17]) or approaches without targets such as algorithms based on weighted 190 endemism[16,19]. While i-SDMs are commonly used as inputs to spatial prioritization, more 191 advanced biodiversity models are rarely used (see Box 2 for exceptions). Multiple biodiversity 192 facets are also rarely considered in conservation despite decades of recognized conservation 193 benefit of *e.g.* phylogenetic diversity[51]. However, in recent years methodological 194 advances[52-54] have enabled the use of phylogenies and functional diversity [16,53] in 195 conservation planning studies at regional[6,52] and global[16,17] extents, and initiatives such as 196 EDGE (evolutionarily distinct and globally endangered; edgeofexistence.org) are promoting the 197 conservation of multiple biodiversity facets. Species interactions are even more rarely 198 considered, although emerging examples show how interactions can be used to evaluate 199 ecological network collapse[55], and set spatial priorities[56] and management scenarios that 200 account for predators and their prey[57]

201

202 The way forward: integrating biodiversity models and conservation evaluation

203 Modeling has a critical, yet under-appreciated role in conservation planning and practice, 204 especially with respect to setting and achieving global conservation targets (Aichi targets for 205 threatened species, habitat loss, and protected areas) in the Strategic Plan for Biodiversity 2011-206 2020 and the post-2020 global biodiversity framework [58]. Models that predict the distribution 207 of species in space and time are crucial to these global initiatives, and i-SDMs are already widely 208 used in conservation[35]. We believe the more advanced biodiversity models have vast potential 209 for representing not just species but entire communities, clades, or functional roles (see early 210 examples in Box 2), but they first must be better-integrated into conservation efforts.

212 The advantages of this integration are many. First, a combined approach retains the unique 213 strengths of each individual approach. Conservation planning would benefit from having access 214 to the most advanced statistical machinery including model-calibration, model validation, and 215 the propagation of uncertainty. Such models can also incorporate more flexible hierarchical 216 structures that account for biases in sampling biodiversity, different sources of inputs (e.g. 217 museum location data with community-based eDNA sampling), different extents (α - and β -218 diversity), and different resolutions of input data. They can take advantage of new data types 219 (e.g. tracking data[59]) as they become available. Conversely, the field of biodiversity modelling 220 would benefit from tailoring predictions for their use in tangible conservation scenarios and 221 targets.

222

Second, an integrated approach reduces any redundancies that arise from using similar datasets and similar approaches in an unsynchronized manner. The potential for redundancy becomes greater the larger and the more complex the representation of biodiversity becomes. Redundancy could be reduced on two fronts—the raw data inputs and the methodological similarities (e.g. combining modelling and conservation optimization algorithms as discussed below).

228

Finally, this synthesis would make the pipeline of data-implementation more efficient and likely to result in a better representation of under-represented biodiversity in conservation. Recent calls to streamline analytical pipelines in ecology[60–62] point to increased efficiency [63] of readily available technological solutions, such as user-friendly interfaces (e.g. R Shiny, Dash).
Importantly, the integration of conservation goals, targets, and scenarios into modelling would require stakeholder input into the modelling decisions[64], which could lead to increased coordination, trust and uptake of these models, decreasing the 'black box' problem of

biodiversity models[60]. It would engage biodiversity scientists beyond academia to understand

the needs of stakeholders and provide more tailored conservation guidance.

238

239 Steps forward

We recommend three related areas of focus to speed integration of biodiversity models and
conservation: (1) adapting biodiversity models to accommodate conservation goals and
scenarios, (2) combining biodiversity models and conservation optimization algorithms, and (3)
re-evaluating model validation in light of conservation goals and scenarios. Recent examples
show this integration is indeed possible (Box 2).

245

246 In many cases, statistical methods already exist to adapt biodiversity models for use in 247 conservation. For example, management actions (or management history) could be included as 248 model predictors, which could then be used to make predictions for different future actions. 249 More complex methods of scenario-building are also possible through altering various model 250 parameters [65,66](Box 2) or even adding sub-models of ecological processes (e.g. population 251 viability). Previously collected 'prior' information (information on species threat or response to 252 management action) can be combined with newly collected data to refine model estimates[67]. 253 The challenge will be to find the best data and models for the conservation question at hand[68], 254 requiring close collaboration between modelers and conservation practitioners[65]. 255

Combining conservation optimization algorithms with biodiversity models is another important
 step (Figure 2) that will enable relevant ecological processes to be directly considered in the

258 conservation alternatives[65]. This integration will require some method developments, but will 259 benefit from the similarities in data types (Figure 1), especially when definitions are 260 synchronized (e.g. α -diversity is equivalent to diversity in planning units, β -diversity to 261 complementarity[69], γ -diversity to the total set of biodiversity). In particular, β -diversity could 262 be used in ecological models with scenarios (Box 2) or within optimization algorithms, 263 streamlining methods and opening the possibility of evaluating conservation value of 264 biodiversity not represented in a single-species approach (e.g. ecological interaction 265 networks[70]).

266

267 Finally, model evaluation—while always a critical step in biodiversity modelling—is especially 268 important with conservation scenarios that are often extrapolations to new situations (e.g. 269 different environmental conditions, alternative restoration scenarios, poorly known taxa). Model 270 evaluation will help address questions such as which type of model to use. For some 271 conservation applications, a highly flexible, accurate model is likely appropriate (e.g. predicting 272 the current distribution of a threatened species). For others, it might be preferable to have models more closely aligned with ecological theory (e.g. when predicting to novel environments such as 273 274 climate change) or when the conservation question depends on a particular variable (e.g. 275 properties of a connectivity corridor). While there are examples that evaluate the ability of 276 advanced biodiversity models to predict rare species[71] and their interactions[72], very few 277 studies have tested how well models extrapolate to new conditions with independent datasets, or 278 whether they predict realistic conservation outcomes. Much more research is needed to 279 understand how biodiversity models will perform in an applied setting. 280

282 Concluding Remarks

283 We urgently need to understand how biodiversity change impacts critical ecosystem functioning 284 and prunes the tree of life. We propose an integration of advanced biodiversity models with 285 conservation goals, targets, assessments, and practice. This integration would improve the ability 286 to rapidly evaluate biodiversity data, make predictions and recommend conservation action for 287 taxa, communities, and ecosystems. The backbone of this integration already exists with recent 288 developments in biodiversity modelling and conservation. Early examples show this integration 289 is possible, could be extended to a number of other conservation efforts and used to address 290 unresolved questions of how to best protect the world's biodiversity (see Outstanding 291 Questions). 292 293 Acknowledgements 294 LJP was supported with Marie Curie Fellowship 'Conserving the Legacy of Evolution into the 295 Future) AMD-659422-1. This work received funding from the ERA-Net BiodivERsA -296 Belmont Forum, with the national funder Agence National pour la Recherche (FutureWeb: 297 ANR-18-EBI4-0009). 298 299 300 **Glossary Box** 301 302 Alpha (α) diversity: quantifies local diversity, usually species richness or effective number of 303 species (Shannon exponent) 304

Beta (β) diversity: quantifies the dissimilarity or turnover between locations in terms of
composition. Can be pairwise between locations, global (average dissimilarity across the region)
or the distinctiveness of a location compared to all others in the region.

308

309 Biodiversity facets: categories of biodiversity that describe relevant taxonomic or ecological 310 information to support biodiversity evaluation, and that apply to and can be (relatively) easily 311 measured for all or most taxa. Examples include taxonomic diversity, phylogenetic diversity, 312 genetic diversity, trait/functional diversity and network diversity.

313

314 Biodiversity models: statistical or process-based models that are used to make inferences and 315 predictions about the effect of the environment on biodiversity, accounting for ecological processes explicitly or implicitly. Biodiversity can be represented in units ranging from 316 317 individuals to entire communities and additionally represent attributes of those taxa (e.g. 318 abundances, functional traits, phylogenetic position, threat). 319 320 Complementarity (conservation planning): quantifies the difference between locations in 321 terms of species or features represented within them (two locations are fully complementary if 322 their pairwise β -diversity is equal to 1).

323

324 Complementarity (ecology): two species are complementary if they fill different roles in an
325 ecosystem or use resources differently.

327	Functional diversity (FD): the diversity of functional forms in a species set (or community)
328	measured by a variety of metrics that use dendrograms or representations in multidimensional
329	space.
330	
331	Gamma (γ) diversity: quantifies the total diversity across locations for an entire area of interest,
332	which could be at regional or global scale depending on the study.
333	
334	Imputation model: a model that estimates values for missing data points in biodiversity datasets
335	(e.g. trait values for a particular taxa).
336	
337	Irreplaceability: the extent to which a location or species is distinct from all others (opposite of
338	redundancy). For example, a location where an endemic species occurs is irreplaceable; an
339	irreplaceable species has a unique position in its community (in terms of function or interactions,
340	i.e. distinctive) or in the phylogeny.
341	
342	Macro-ecological model : a biodiversity model that uses a top-down approach to model α - or β -
343	diversity directly rather than modeling the distributions of the component taxa.
344	
345	Phylogenetic diversity (PD): The evolutionary diversity represented by sets of taxa, with the
346	most common metric (Faith's PD) being the branch length of the minimum spanning tree
347	connecting a set of species on a phylogeny[51] (sets of species can be from a single area or
348	multiple areas combined).
349	

350	'Sideways' biodiversity models: models that predict the distribution of biodiversity with a
351	combination of bottom-up (i.e. single-taxon predictions) and top-down (i.e. models of the
352	properties of an assemblage or community of taxa) approaches.
353	
354	Spatial biodiversity models: biodiversity models that are explicitly spatial, in which inferences
355	and predictions are made for biodiversity in certain locations (e.g. planning units or grid cells).
356	
357	Spatial prioritization: A form of systematic conservation planning (SCP) that selects a set of
358	areas that maximizes conservation value given other constraints (e.g. cost, protected areas,
359	feasibility).
360	
361	Weighted endemism: the ratio between a species local range of occurrence and its total range.
362	Note: this is different than the definition of endemism which is the extent to which a species
363	range is restricted to a particular location.
364	
365	
366	Figure 1. How biodiversity facets, different scales of biodiversity and conservation concerns
367	factor into biodiversity modelling and conservation evaluation. Most modelling and evaluation
368	involve particular target species rather than other facets of diversity (e.g. evolutionary history,
369	species interactions). Most biodiversity models focus on ecological processes rather than
370	explicitly considering conservation concerns, although many models are built for threatened
371	species or species of conservation interest resulting in a mostly unidirectional flow of
372	information from models to evaluation. In the case of spatial conservation planning, both models
373	and evaluation use similar data inputs and spatial organization of biodiversity. Both have local

scale (or planning units) and a regional or global scale. In particular, the use of metrics or
algorithms to differentiate sites from one another (beta diversity in modelling, complementarity
in conservation planning) are treated similarly in modelling and planning.

377

Figure 2. The steps and information flow in a typical data-implementation pipeline (left) and a streamlined version (right) that combines models and evaluation in one step. We expect that more biodiversity information would make it through to the application stage (orange arrows) with a unified rather than disconnected method of evaluating biodiversity from the data products to their use in conservation applications, and that the increasing transparency and additional communication between biodiversity scientists and stakeholders is required by the combined approach.

385

386 Figure I. (Embedded in Box 1). The dimensions of recent biodiversity models. Models can have 387 different types of response objectives: within-site models, between-site models and models of 388 individual species. These response objectives fall along two axes: (1) top-down approaches 389 where α -diversity or other diversity metrics are the focus versus bottom-up where individual 390 species are the focus (vertical axis), and (2) whether the focus is on predicting biodiversity 391 within sites or between sites (horizontal axis). DynamicFOAM: Dynamic Framework for 392 Occurrence Allocation in Metacommunities, GDM: Generalized Dissimilarity Modelling, MEM: 393 macro-ecological model, i-SDM: individual or single-species distribution model, s-SDM: 394 stacked species distribution model, j-SDM: joint species distribution model, SESAM: Spatially 395 explicit species assemblage modelling.

396

Box 1. Modelling biodiversity from the top-down, bottom-up and sideways

398 Biodiversity models can be defined by their 'response objectives' and internal structure 399 (particularly the response variable—what is being predicted). These response objectives can be 400 represented with axes that describe whether the objective is a top-down versus bottom-up or a 401 within- versus between-site depiction of biodiversity (Fig. I). Current modelling approaches can 402 be mapped onto this space, with potentially the most information rich approaches (grey circle in 403 Fig. I) cross-linking biodiversity facets (traits and phylogenies) while jointly predicting 404 community composition (e.g. species abundance or occurrence), within-site community 405 diversity, and between-site turnover.

406

In the top-down approach, known as a **macro-ecological model** (MEM), α -diversity is itself is the response (e.g. species richness or phylogenetic diversity) and the input data is a site by diversity matrix. β -diversity models (e.g. GDM; Generalized Dissimilarity Modelling[73] or MBM; multi-facetted biodiversity models[49]) estimate turnover in species or phylogenetic diversity[74] between sites as a function of differences in environmental and geographic distances.

413

Bottom-up approaches, where individual species are the response variables (single-species distribution model, i-SDM), are by far the most common type of biodiversity model used. Note that α -diversity can also be derived from aggregating models of individual species together in a stacked species distribution model (s-SDM). This feature is important for conservation planning, because both the species pool of a set of sites and diversity metrics for entire species assemblages can be calculated (e.g. endemism-based metrics). However, resulting diversity 420 metrics are simply the sum of the parts—there is no information on species associations or421 community-level attributes.

422 'Sideways' methods are methods that combine top-down and bottom-up approaches: Joint 423 Species Distribution Models (j-SDMs), which are the multi-species version of i-SDMs that model effects of individual species and communities[39,40], and can include shared responses 424 425 between species and can include other information such as phylogeny and traits in a hierarchical 426 framework, and spatial and temporal dynamics[66]; Dynamic Framework for Occurrence 427 Allocation in Metacommunities (DynamicFOAM)[75], which balances richness and turnover to 428 predict occurrences for all species, SESAM: Spatially explicit species assemblage modelling[76], 429 which adjusts combined s-SDM predictions based on those of macro-ecological models.

430

431 Box 2. Hybrid approaches for integrating advanced biodiversity models and conservation 432 decisions

We have emphasized the need for models that join different spatial and taxonomic dimensions of
biodiversity that can be combined with conservation evaluation. Here are examples from two
classes of advanced biodiversity models (Box 1) that directly address applied conservation
questions. There is much scope for building upon and extending these early examples.

437

438 *1. Macro-ecological models*

439 For taxa for which there is no abundance or time-series data, models based on α - and β -diversity 440 (Box1) can also be used to directly design, predict, and compare conservation scenarios. For 441 example, β -diversity models can summarise the expected effects on future biodiversity of 442 changes to habitat condition, extent or management, as change in effective habitat area[77]. With 443 sophisticated models of environmental or land-use change for example, new protected areas, habitat loss[78] or degradation and climate change[79], scenarios can be planned for the most
beneficial effect on biodiversity even at global extents[77], and can also be used with

446 phylogenetic[74] or functional[49] ß-diversity.

447 2. 'Sideways' biodiversity models

448 Multi-species models that contain a hierarchy for a species-level and community (or ecosystem)-449 level (Box 1) are widely relevant to conservation applications because they can simultaneously 450 estimate ecological processes relevant for individual species and for larger-scale attributes (e.g. 451 ecosystem properties). Perhaps the best example of this approach has been for evaluating 452 management strategies in fisheries, in which models of fish species (or sets of species) are 453 combined with ecosystem level attributes[64]. In this way, important ecological and ecosystem 454 constraints can be considered (e.g. population dynamics in response to habitat or predators) 455 along with management scenarios (harvest controls)[57]. Recent versions allow for evaluation of 456 harvest impacts, survey designs, and protection scenarios for finescale habitats in a changing 457 climate[66]. While these more elaborate designs are only possible for a subset of well-studied 458 taxa, this will be increasingly possible with ever-improving sampling and imputation models. 459

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