Operationalizing ecological connectivity in spatial conservation planning with Marxan Connect

- ³ *Rémi M. Daigle, Dalhousie University, Department of Oceanography, 1355 Oxford Street, Halifax, NS,
- ⁴ Canada, B3H 4R2 and Université Laval, Département de biologie, 1045, av. de la Médecine, Québec, QC,
 ⁵ Canada, G1V 0A6
- Anna Metaxas, Dalhousie University, Department of Oceanography, 1355 Oxford Street, Halifax, NS, Canada,
 7 B3H 4R2
- Arieanna Balbar, Dalhousie University, Department of Oceanography, 1355 Oxford Street, Halifax, NS,
 Canada, B3H 4R2
- Jennifer McGowan, Centre for Biodiversity and Conservation Science, School of Biological Sciences, University
 of Queensland, St. Lucia, QLD 4072, Australia.
- Eric A. Treml, School of Life and Environmental Sciences, Deakin University, Waurn Ponds, VIC 3216,
 Australia.
- Caitlin D. Kuempel, Centre for Biodiversity and Conservation Science, School of Biological Sciences, University
 of Queensland, St. Lucia, QLD 4072, Australia.
- ¹⁶ Hugh P. Possingham, The Nature Conservancy, South Brisbane, Queensland 4101, Australia.
- ¹⁷ Maria Beger, School of Biology, Faculty of Biological Sciences, University of Leeds, LS2 9JT, UK and Centre
- ¹⁸ for Biodiversity and Conservation Science, School of Biological Sciences, University of Queensland, Brisbane,
- ¹⁹ QLD 4072, Australia.

- ²⁰ * Corresponding Author: remi.daigle@dal.ca
- ²¹ Running Head (<45 characters): Operationalizing connectivity in spatial planning
- Keywords: Connectivity; Conservation; Marxan; Movement Ecology; Spatial Planning; Structural connec tivity

Operationalizing connectivity in spatial planning

²⁴ Abstract:

- Globally, protected areas are being established to protect biodiversity and to promote ecosystem
 resilience. The typical spatial conservation planning process leading to the creation of these protected
 areas focuses on representation and replication of ecological features, often using decision support
 systems such as Marxan. Unfortunately, Marxan currently requires manual input or specialised scripts
 to explicitly consider ecological connectivity, a property critical to metapopulation persistence and
 resilience.
- ³¹ 2. "Marxan Connect" is a new open source, open access Graphical User Interface (GUI) designed to assist
 ³² conservation planners in the systematic operationalization of ecological connectivity in protected area
 ³³ network planning.
- Marxan Connect is able to incorporate estimates of demographic connectivity (*e.g.* derived from tracking data, dispersal models, or genetics) or structural landscape connectivity (*e.g.* isolation by resistance). This is accomplished by calculating metapopulation-relevant connectivity metrics (*e.g.* eigenvector centrality) and treating those as conservation features, or using the connectivity data as a spatial dependency amongst sites to be included in the prioritization process.
- 4. Marxan Connect allows a wide group of users to incorporate directional ecological connectivity into
 conservation plans. The least-cost conservation solutions provided by Marxan Connect, combined with
- ecologically relevant post-hoc testing, are more likely to support persistent and resilient metapopulations
- (*e.g.* fish stocks) and provide better protection for biodiversity than if connectivity is ignored.

Operationalizing connectivity in spatial planning

Introduction 43

Connectivity, in its most general form, refers to the exchange of individuals (including genes, traits, disease, 44 etc.), energy or materials among habitat patches, populations, communities or ecosystems. Maintaining 45 connectivity can improve population resilience to perturbations, increase metapopulation viability, promote 46 genetic diversity and maintain energetic pathways among ecosystems (Palumbi 2003; Figueira & Crowder 47 2006; Lowe & Allendorf 2010). Connectivity also appears at the forefront of global international conservation 48 policy, so as Aichi Target 11, which commits 197 countries to establishing "effective, representative, and 49 well-connected" networks of reserves by 2020 (UNEP 2010). 50

- There are many metrics and methods to evaluate the connectivity of sea/landscapes and these can be used 51
- to assess networks of protected areas and influence spatial conservation planning in the future (Saura & 52
- Pascual-Hortal 2007; Beger et al. 2010a; Chollett et al. 2017; D'Aloia et al. 2017; Zeller et al. 2018). The 53
- quantity and quality of empirical data used to calculate connectivity have been growing rapidly in the last 54
- few years (Kool, Moilanen & Treml 2013; Hussey et al. 2015; Magris et al. 2018; Zeller et al. 2018). In 55 turn, methods for estimating ecological connectivity are also advancing, and new conservation planning tools 56
- are quickly emerging to capitalize on these new data and methods (Saura & Pascual-Hortal 2007; Beger et 57
- al. 2010b; White et al. 2014). Examples of connectivity data that have been incorporated in conservation 58
- applications include: gene flow (Beger et al. 2014; Marrotte et al. 2017), dynamic distributions and migratory 59 bottlenecks on migratory pathways (Iwamura et al. 2013; Runge et al. 2016), maximizing larval flow (Magris
- 60 et al. 2016; D'Aloia et al. 2017), ontogenetic shifts in habitat use (Brown et al. 2016; Weeks 2017), ensuring 61
- the movement of adult individuals pathways (Beger et al. 2015; Mazor et al. 2016; Pereira, Saura & Jordán 62
- 2017; Zeller et al. 2018), and maintaining fisheries benefits (Daigle, Monaco & Elgin 2017; Krueck et al. 63
- 2017). Despite these efforts, connectivity is not commonly being incorporated in on-the-ground decision 64
- making for planning (Beger et al. 2010a; Barnes et al. 2018; Balbar, unpublished data). This is largely 65
- because connectivity metrics are not well defined or standardized, practitioners often lack confidence in the 66
- data or the expertise to work with them, and approaches to explicitly incorporate connectivity patterns in 67
- spatial planning are rare. 68

- Spatial conservation planning is an approach that guides the allocation of conservation resources to areas 69
- identified as important for biodiversity whilst minimising the conservation impact on resource users (Margules 70
- & Pressev 2000; Moilanen, Wilson & Possingham 2009; Wilson, Cabeza & Klein 2009). The process of spatial 71
- planning demands setting broad goals, which can be turned into quantifiable objectives that lead to the 72
- conservation of biodiversity (Tear et al. 2005) and which, in turn, link back to actions, costs and feasibility 73
- (Wilson et al. 2007). Spatial planning often relies on the use of decision-support software (e.q. Marxan or74
- Zonation) to help decide the location and timing of actions (e.q. establishing protected areas) to best achieve 75
- conservation objectives. These tools are primarily used to develop representative and cost-efficient conservation 76
- plans by meeting targets for species or habitats, with the consideration of connectivity patterns primarily 77 expressed by prioritising adjacent or contiguous sites. To advance the inclusion of ecological connectivity 78
- into the spatial planning process, technical documentation, best-practice guidelines and user-friendly tools
- are needed. Knowing how to best identify, evaluate, and treat connectivity data to meet different objectives 80
- within a given spatial planning framework is important to better capture key ecological processes in planning. 81
- Here, we outline potential workflows of realising connectivity in spatial planning, including the treatment of 82 various data formats, key decision points that link back to objectives, types of data related to connectivity, 83

Operationalizing connectivity in spatial planning

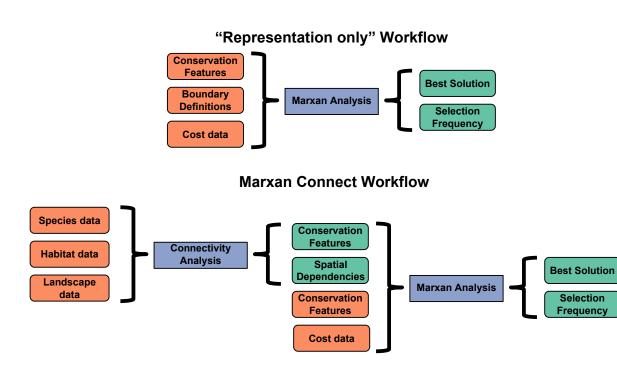


Figure 1: Comparison of workflows between the "representation only" approach to Marxan and "Marxan Connect." Marxan Connect facilitates the use of connectivity data, derived from tagging data, genetics, dispersal models, resistance models, or geographic distance, by producing connectivity metrics and connectivity strengths (*i.e.* spatial dependencies that are used in the place of boundary definitions) before running Marxan. These connectivity metrics and linkage strengths are then used as inputs (connectivity-based conservation features or spatial dependencies) in the traditional Marxan workflow. The cost data in the traditional Marxan analysis refers to the cost of protecting a planning unit (*i.e.* opportunity cost), not the cost to traverse a landscape.

evaluation and post-hoc analysis. We do so in the context of the widely used spatial planning tool Marxan, which aims to represent biodiversity whilst minimizing overall cost (Ball, Possingham & Watts 2009). We then introduce a new open source and open access tool called Marxan Connect to help users operationalize these concepts within Marxan. Our objective is to enable an overview of the selection and treatment of connectivity data to encourage its use in spatial conservation planning.

Box 1: A primer for spatial conservation planning with Marxan Marxan uses a simulated annealing algorithm to find good solutions to the "minimum set" problem. In the minimum set problem, the user specifies an amount of each conservation feature j that needs to be conserved, or conservation targets (T_j) , for each conservation feature. The basic minimum set problem is an integer linear programming problem and does not consider connectivity:

minimize
$$\sum_{i=1}^{N} c_i x_i$$
, given that we should meet all targets

Operationalizing connectivity in spatial planning

$$\sum_{i=1}^{N} x_i r_{ij} \ge T_j, \text{ for all features} j,$$

where N is the number of planning units, c_i is the cost of planning unit i, r_{ij} is the amount of 95 feature j in planning unit i, and x_i is a control variable which has the value of 1 for selected sites 96 and 0 for unselected sites. It is usually desirable to include some basic spatial properties of a 97 protected area system such as geographic proximity or adjacency information between planning 98 units to help minimize costs or maximize clumping of a protected area system. For example, if 99 the common boundary between every pair of planning units is known, then the minimum set 100 problems can be extended to include a term for the boundary length of the reserve system and an 101 effort made to minimise it: 102

minimize
$$\sum_{i=1}^{N} c_i x_i + b \sum_{i=1}^{N} \sum_{h=1}^{N} x_i (1-x_h) c v_{ih}$$
, given that we should meet all targets
 $\sum_{i=1}^{N} x_i r_{ij} \ge T_j$, for all features j ,

- where b is the boundary length modifier (BLM), and cv_{ih} represents the cost of a boundary and is typically the length of the physical boundary between sites i and h. Costs (c_i) in Marxan often pertain to socio-economic implications of protecting a site, such as management or opportunity costs. For more information see Ball et al. (2009) and Ardron et al. (2010). Key terms and definitions:
- **Planning area**: the spatial domain over which the planning process occurs. This is synonymous with terms "domain" or "extent" or "study area" in other fields. This area is subdivided into smaller "Planning Units".
 - **Planning unit**: spatial units within the entire planning area (*i.e.* domain, or study area), which can be defined using regular gridded (*e.g.* hexagonal) or using landscape features-based (*e.g.* reefs, water catchments) as in Marxan.
 - Boundary Length: the shared boundary length between adjacent planning units.
 - Boundary Length Modifier (BLM): a weighting parameter to 'tune' the influence of the boundaries. The BLM helps achieve "clumped" solutions by reducing the overall edge to area ratio. A higher BLM value results in a more 'clumped' Marxan solution.
 - **Conservation feature**: the features (*e.g.* habitats, species, processes) for which a target is set.
 - **Conservation target**: the minimum quantity or proportion of the conservation feature in the study area to be included in solutions.
 - Solution: a binary output of Marxan reflecting whether a planning unit is selected (1) or not selected (0) as part of the conservation plan.
- Selection Frequency: the summed solution output of Marxan reflecting how many times a planning unit was selected across runs

126

111

112

113

114

115

116

117

118

119

120

121

122

Operationalizing connectivity in spatial planning

¹²⁷ Understanding connectivity data

One of the challenges associated with integrating ecological connectivity in spatial planning is the wide 128 variety of entities that move (e.q. organism, gene, pollutant) and movement processes (e.q. migration route,129 larval dispersal, multi-generational gene flow, carbon flux). While there are many types of data sources, 130 connectivity data are often stored as matrices, where donor (or source) sites are rows, and the recipient (or 131 destination) sites are columns. Alternatively, connectivity data may be stored in an edge list where the first 132 column contains the donor site IDs, the second column contains the recipient site IDs, and the third column 133 contains the connectivity value. Below, we review a few of the most common data sources organized by their 134 treatment in Marxan Connect. Additional details on data format, types, mathematical representations and 135 associated assumptions can be found on the Marxan Connect tool website, marxanconnect.ca. 136

¹³⁷ Landscape-based estimates of connectivity

Some spatial planners may have access to detailed connectivity information based on demographic data (See "Demographic estimates of connectivity" section below). In these cases, Marxan Connect can generate estimates of connectivity strength (*e.g.* spatial isolation) based either on the Euclidean distance between habitats, or isolation by resistance (McRae & Nürnberger 2006). These landscape-based connectivity estimates are often more limited in their applicability than demographic data (*e.g.* self-recruitment), but require less data.

144 Linkages across a habitat matrix

The structure and spatial configuration of the land- or sea-scape (*i.e.* habitat type, size, and spacing) can 145 impede or facilitate the movement of organisms. The rate at which impediment or facilitation occurs has 146 been defined as the strength of landscape connectivity (Tischendorf & Fahrig 2000). The impediment or 147 facilitation (*i.e.* resistance or cost to traverse landscape) posed by habitat types can be estimated from 148 tracking data, genetic data, expert opinion, or habitat suitability models for species-centric approaches (Bunn, 149 Urban & Keitt 2000; Urban & Keitt 2001; Ricketts 2001; Zeller, McGarigal & Whiteley 2012). For a habitat 150 or multi-species centric approach, resistance can also be estimated from the similarity in environmental 151 variables (e.q. land cover) or that of species assemblages (Schumaker 1996). From this resistance surface, it is 152 possible to estimate the rate of movement of organisms across the landscape based on the spatial arrangement 153 of habitat patches using various methods such as least-cost path analysis, and current density approaches 154 (Fall et al. 2007; Rayfield, Fortin & Fall 2010; Koen et al. 2014). While these methods are conceptually 155 similar, they produce qualitatively different connectivity estimates and may be difficult to validate (Saura & 156 Pascual-Hortal 2007; Rayfield, Fortin & Fall 2010; Zeller et al. 2018). 157

¹⁵⁸ The conservation implications of these differences in connectivity have not been fully explored in the context

¹⁵⁹ of spatial planning. While methods based on Euclidean distance can be species non-specific, resistance-based

 $_{160}$ models necessarily focus on specific species (Ricketts 2001). However, some features of the landscape (*i.e.*

¹⁶¹ calculated using the multi-species approach) may have different, but important uses for multiple species. For

example, current density is a metric adopted from electrical circuit theory which, in movement ecology, is

¹⁶³ intended to represent the prevalence of movement of organisms across a landscape. However, some species

Operationalizing connectivity in spatial planning

(amphibians and reptiles) may use areas of high current density as movement pathways while others (fishers -

¹⁶⁵ Martes pennanti) use these areas as home ranges (Koen et al. 2014).

¹⁶⁶ Demographic estimates of connectivity

¹⁶⁷ Whether the movement of 'individual' organisms, particles of detritus or pollutants are being directly measured ¹⁶⁸ (tagging and tracking) or estimated (genetics or models), Marxan Connect treats these data as "demographic ¹⁶⁹ connectivity" (See marxanconnect.ca for more details on the mathematical representations of connectivity ¹⁷⁰ data). The strength of connectivity is measured as a probability or an absolute amount.

171 Tagging and Tracking

172

173

174

175

176

177

178

179

180

Movement ecology has seen profound advances over the last decades arising from the ability to identify and track individuals and thus understand the movement of organisms in space and time (Hussey *et al.* 2015). Traditionally, individual organisms have been identified using scarring, banding, tagging, radiotelemetry collars, passive integrated transponder devices (PIT tags), otolith/statolith microchemical signatures, or other approaches that allow an observer to track the movement of organisms through a landscape (Scott 1942; Thomas & Marburger 1964; Dunn & Gipson 1977; Twigg 1978; McNeil & Crossman 1979; Whitfield Gibbons & Andrews 2004). The advent of Global Positioning System (GPS) tags, satellite communication, and other forms of data relay provide opportunities to collect tracking data at higher spatiotemporal resolution and on a wider range of organisms, such as long-distance migratory species (Voegeli *et al.* 2001; Cagnacci *et al.* 2010).

Regardless of the approach, marking and tagging observations typically consist of a sequence of times and locations at which an individual was observed. The sequence often records numerous processes including ontogenetic movements, foraging, seasonal migrations, etc. The data can be used in spatial planning for objectives related to habitat use (*e.g.* identifying core foraging areas), movement pathways (*e.g.* finding migration routes) and species demography (*e.g.* disease spread through a population; McGowan *et al.* 2017b). Since tagging or tracking data are typically collected from relatively few individuals, they are often spatially biased and do not record multi-generational variation in movement.

¹⁸⁸ Genetic approaches for estimating ecological connectivity

Genetic approaches have long been used to estimate the degree to which populations have diverged, and 189 the degree to which gene flow via dispersal or migration influences this divergence (Palumbi 2003). As 190 molecular techniques have changed, divergence and gene flow can be now be quantified at much finer spatial 191 and temporal scales (Manel & Holderegger 2013; Hand et al. 2015). The integration of this fine-scale 192 genetic information is being proposed as central to conservation and management efforts, despite technical 193 and conceptual challenges (Beger et al. 2014). Generally, assignment methods are being used to develop 194 estimates of ecological connectivity at the scale of populations (e.g., using Structure Pritchard, Stephens & 195 Donnelly 2000), whereby individuals are 'assigned' back to the population of origin. Similarly, parent-offspring 196 assignments can be developed based on DNA fingerprinting to identify realised dispersal events over a single 197 dispersal event (e.g. Saenz-Agudelo et al. 2009). In both cases, ecological, or demographically-significant. 198 connectivity estimates between conservation planning units (and/or populations) can be estimated, if sampling 199

Operationalizing connectivity in spatial planning

is thorough and consistent with the planning unit structure. The result of these genetic approaches is often a migration matrix representing the likelihood that individuals or genotypes found at some destination populations came from the suite of sampled source populations. Unfortunately, these data are expensive to collect at the scale and scope appropriate for conservation applications, and appropriately interpreting the connectivity results is often highly context dependent. To date, the few published academic studies attempting this have struggled with significant compromises in taxonomic coverage, geographic extent, and alignment with the planning process (Harrison *et al.* 2012; *e.g.* Beger *et al.* 2014).

207 Individual-based models of movement or dispersal

Models of movement for individual organisms can be very useful in estimating connectivity. These models are 208 typically based on the physical environment (e.q. ocean currents) and behavioural (e.q. resource selection) 209 processes which influence movement. To model the movement of materials or individuals through land and 210 seascapes, advection-diffusion models are often used as they can efficiently capture the physical fluid-like 211 transport through these environments, including turbulence (Metaxas 2001), and other biological traits such 212 as behaviour and duration of the dispersal phase (Roughgarden, Gaines & Possingham 1988; Cowen et al. 213 2000; Bode, Bode & Armsworth 2006; Paris, Chérubin & Cowen 2007; Metaxas & Saunders 2009; Treml 214 et al. 2015; Daigle, Chassé & Metaxas 2016). In general, these modelling approaches are used to calculate 215 the probability of exchange of individuals between habitat patches. Particle tracking approaches link source 216 populations to destinations and vice versa allowing one to calculate the probability of linking any source to 217 any destination and vice versa. These probabilities can then be used as the connectivity strength. Behaviour 218 motivated movement requires some specific knowledge of the species being modelled. This behaviour can 219 be based on resource availability (Chetkiewicz & Boyce 2009), predator avoidance (Bracis 2015), or other 220 environmental cues (Daigle & Metaxas 2012; Zeller et al. 2018). While it is computationally feasible to scale 221 up these models, it is often difficult to appropriately validate these complex models at the broad-scale most 222 relevant to area-based management. 223

²²⁴ Using connectivity data in spatial planning

In addition to the often used "rules of thumb" for connectivity which guide sizing and spacing of marine 225 protected areas (Mora et al. 2006; Smith & Metaxas 2018), there are several different methods to directly 226 incorporate connectivity data into spatial planning tools. For Marxan specifically, these include: 1) treating 227 connectivity properties of planning units as conservation features (continuous or discrete) for which a target 228 is set; 2) including connectivity strengths among planning units as spatial dependencies within the objective 229 function; 3) treating connectivity properties of planning units as a connectivity cost; and 4) customizing the 230 objective function to incorporate connectivity performance metrics. Methods 1) and 2) are fully implemented 231 within Marxan Connect (Figure 2) while 3) is supported, but not facilitated for reasons described below. 232 Method 4) is currently an area of active research not yet implemented. We note that in the following sections, 233 our objective is to identify and discuss different treatments of connectivity data. We follow up with notes 234 on making decisions related to data or methods, and we stress the importance of post-hoc evaluations in 235 separate sections below. Our objective is not to be prescriptive about "best practices" although we do offer 236 some insights where appropriate. 237

Operationalizing connectivity in spatial planning

²³⁸ Connectivity as conservation features

A simple and accessible way to integrate connectivity data into spatial planning is to treat it as a conservation 239 feature, such that r_{ij} is the amount of connectivity feature j (e.g. reproductive outflux) in planning unit i. 240 In the classical minimum set problem (*i.e.* Marxan), a target is set for each feature, T_i , (*e.g.* 50%) and the 241 reserve system needs to contain at least that amount. This method can incorporate continuous or discrete 242 conservation features (Figure 2). Examples of conservation targets best represented as continuous data include 243 patch-level self-recruitment values (range between 0 and 1) or centrality measures (White et al. 2014; D'Aloia 244 et al. 2017; Magris et al. 2018). In this treatment, a metric that represents the process of connectivity 245 across the entire planning area receives a target. Definitions and potential conservation objectives for each 246 connectivity based conservation feature available in Marxan Connect can be found on marxanconnect.ca. 247

To increase the probability that the connectivity process is maintained, and that the spatial conservation plan is influenced by the conservation feature, a target higher than that of most other conservation features should be set. We suggest using a tunable 'constraint', a connectivity target multiplier (C), as a way to determine the higher target for connectivity-based conservation features (Figure 2 & 3). The target for the connectivity-based conservation feature, T_c , would then be:

$$T_c = T_j * C$$

with T_i being a typical conservation target for features not related to connectivity. An appropriate value for 253 C can be determined by using a cost trade-off curve, similar to calibrating the BLM, where one would test 254 the sensitivity of cost of the best solution and the total summed metric. It is worth noting that the BLM may 255 interact with C. The value of C could be chosen as the divergent point, where the greatest increase in the 256 connectivity metric is achieved for a relatively small increase in cost. However, the preferred approach would 257 be to establish conservation targets leading to a reserve network design which meets ecologically relevant 258 conservation objective(s), such as population viability or a within network metapopulation growth rate > 1259 (See "Post-hoc evaluation" section for more details). 260

In contrast, conservation targets set for discretized connectivity conservation features X_j can capture different 261 levels of importance to connectivity amongst sites (Álvarez-Romero et al. 2018; Figure 2). To create a 262 discrete feature, threshold value(s) must be chosen from the range of the continuous connectivity metric, 263 ideally after ecological assessment or sensitivity analysis. The planning units that meet the threshold(s) are 264 then discretized into unique features for which a target is set (Figure 2). Similarly, this type of threshold 265 setting is often used with species distribution models, where each planning unit is assigned a probability of 266 species occurrence, and a threshold value is used to convert these continuous distribution data to a binary 267 map (presence vs absence or suitable vs unsuitable) to represent a particular species as a conservation feature 268 (e.g. Minor et al. 2008). When appropriate, discrete and very high priority planning units could alternately 269 be "locked-in" to the final solution to guarantee their inclusion. One disadvantage of setting a connectivity 270 metric as a conservation feature is where the benefits of that connectivity conservation features are highly 271 dependent on the final set of actions. For example, this approach currently does not allow the iterative 272

²⁷³ recalculation of the metric as sites are added to the solution.

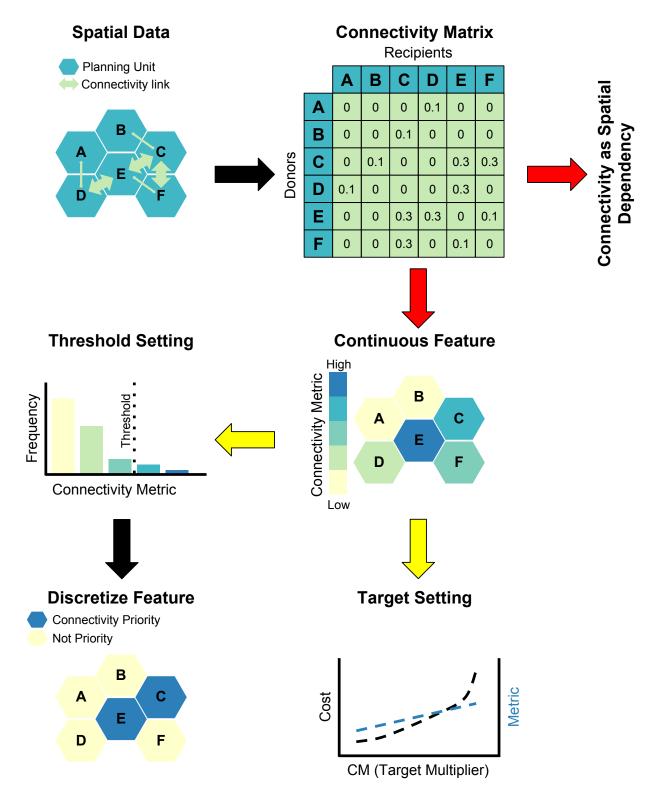


Figure 2: Data processing workflow for three possible methods for using raw connectivity data in spatial prioritization: 1) Connectivity as Spatial Dependency in the objective function (Raw Data -> Connectivity Matrix -> Connectivity as Spatial Dependency), 2) Continuous conservation features (Raw Data -> Connectivity Matrix -> Continuous Metrics -> Target Setting), or 3) Discrete conservation features (Raw Data -> Connectivity Matrix -> Continuous Metrics -> Threshold Setting -> Discrete Metrics. Coloured Arrows indicate important decision points. Red arrows indicate the conservation feature vs connectivity strength method decision point and yellow arrows indicate the continuous vs discrete conservation feature decision point. 10

Operationalizing connectivity in spatial planning

²⁷⁴ Connectivity as spatial dependencies in the objective function

Another approach to incorporate connectivity in Marxan is using the directional connectivity-based data to 275 inform the boundary cost (cv_{ih}) . This cost signifies the penalty associated with protecting one site but not 276 other sites to which it is connected (Beger et al. 2010b). Here, a directional connectivity dataset is used to 277 parameterize the boundary cost between all pairs of planning units (including non-adjacent units) and set 278 the importance of incorporating connectivity with the BLM (or b in the equation in the "A spatial planning 279 primer" section), also called "Connectivity Strength Modifier" (CSM) (Beger et al. 2010b). Increasing the 280 BLM reduces the edge to area ratio by minimizing costs associated with unprotected adjacent boundaries: 281 similarly, increasing the BLM in the connectivity context (*i.e.* CSM) will reduce the amount of leakage from 282 the network (*i.e.* amount lost from the network) by minimizing costs associated with unprotected connectivity 283

linkages, and thus maximising connectivity strengths across the entire Marxan solution.

Connectivity estimates as spatial dependencies in the objective function have been used to design marine protected area (MPA) networks in the Coral Triangle (Beger *et al.* 2010b, 2015). This approach can maximize the within-network connectivity, and may improve the metapopulation growth rate and other performance metrics. In Marxan Connect, one can combine the use of connectivity as spatial dependencies with a locked-in "Focus Area" (*e.g.* an existing protected area) to generate candidate stepping stones. However, the method will exclude isolated sites from the final solution unless these are included using other methods (*e.g.* a conservation feature for an isolated site which happens to contain a unique species).

²⁹² Connectivity as a Cost

A common approach to attributing costs in Marxan is to use inverse values as a treatment of the cost to be 293 minimized. For example, such an approach might take the distance of a planning unit to the nearest port as 294 a proxy of the cost to coastal fishing industries when establishing MPAs (Maina et al. 2015; Mazor et al. 295 2016; McGowan et al. 2017a). Here, the planning units closer to shore will be less costly than those farther 296 offshore as the distance grows. Thus, an inverse distance cost makes the planning units closer to shore more 297 expensive and less desirable for selection than those farther offshore. A more recently proposed method is to 298 use connectivity as the cost to be minimized in the Marxan objective function. For example, Weeks (2017) 299 used a "seascape connectivity cost" representing the inverse of the connectivity, expressed as the distance 300 between adult habitat to nursery habitat. A disadvantage of the "connectivity as a cost" approach is that it 301 precludes the consideration of other important socio-economic costs in the analysis, which are crucial for 302 reducing conflicts with resource users and increasing the cost-effectiveness of implementation and management 303 (Ban and Klein 2009). Further, this approach is not ideal since each planning unit's contribution to the 304 connectivity of the entire system relies on whether other sites are "in" or "out" of the reserve system. 305

³⁰⁶ Connectivity-based objective function

Where the goal of including connectivity data into the spatial planning problem is to maximise the likelihood of the species' persistence, then the most appropriate approach would be to include a persistence metric (e.q. population viability; metapopulation capacity) within the objective function. In this case, population

³¹⁰ connectivity together with fecundity, mortality and survival are included as a metapopulation model within

Operationalizing connectivity in spatial planning

the optimisation process. Currently, this is only realistic computationally for small problems (tens to hundreds 311 of planning units) because the algorithm needs to calculate the performance metric very fast for simulated 312 annealing to deliver good answers reasonably quickly. For example, Chollett *et al.* (2017) used a genetic 313 algorithm to optimize an MPA network for maximum population persistence and fisheries yield, but this 314 approach took 2 days on a high performance computing cluster which equates to ~ 300 days of single processor 315 computing time. Even with powerful computational methods, such as integer linear programming (Beyer et316 al. 2016; Hanson et al. 2017), the problem formulation would be challenging as the performance metric of 317 interest may vary non-linearly. Connectivity-based objective functions and their implementation in decision 318 support software is a research priority. 319

³²⁰ Making decisions: models, matrices and methods

Most data for the above methods focus on single species, yet many protected areas are designed to protect a 321 diversity of species. There are many strategies for combining single-species connectivity data into a single 322 multi-species connectivity matrix. Examples include taking the arithmetic or geometric means of multiple 323 connectivity matrices, or connectivity metrics derived from them (Melià et al. 2016; D'Aloia et al. 2017). 324 Others have calculated the probability of at least one species, or all species being connected (Jonsson, Nilsson 325 Jacobi & Moksnes 2016; Magris et al. 2016). However, in all these cases, generating multi-species metrics 326 or connectivity matrices resulted in some level of compromise that was suboptimal for a single species. If 327 targeting conservation features, the most efficient solutions will be achieved by using single species conservation 328 features that are representative of multiple life-stages and species with varying dispersal traits (Beger et al. 329 2015; Magris et al. 2016; Albert et al. 2017; D'Aloia et al. 2017). Alternatively, if the connectivity data 330 are used to modify the boundary definitions, then a single connectivity dataset (e.g. edge list or matrix) 331 per Marxan optimization must be used. Therefore, users are forced to calculate multi-species connectivity 332 metrics, or run Marxan once per species and potentially combine the outputs. In all cases the consequences 333 or trade-offs of the chosen strategy should be evaluated. 334

If the sole conservation objective is to maximize among-reserve connectivity, then modifying the boundary definitions and locking in existing reserves will likely produce the most efficient results. However, targeting connectivity features allows for more flexibility in objectives such as protecting areas that are important to maintain entire ecosystems (not just in reserves), avoiding areas with invasive species, or targetting areas with higher larval or adult spillover into unprotected sites. While these approaches may be slightly redundant, they are not mutually exclusive. Regardless of the data or method(s) that are chosen, post-hoc evaluations should be used to evaluate competing strategies.

The spatial and temporal scales of the connectivity data should be considered in all approaches. Ideally, 342 connectivity should be quantified at the spatial scale of the planning units because the assumptions needed 343 for rescaling can lead to erroneous results. The temporal scale of connectivity data should be aligned with 344 the conservation objectives, such as providing demographic (e.g., single generation movement) connectivity 345 or safeguarding long-term gene flow (e.q., many 100s of generations). Additionally, the planning area should 346 extend beyond jurisdictional boundaries and the focus area to avoid edge effects which are particularly 347 consequential for connectivity data (e.g. important source habitat may exist just beyond jurisdictional 348 boundaries or focus areas). 349

Operationalizing connectivity in spatial planning

350 Post-hoc evaluation

None of the above methods, aside from customising the objective function, guarantee that population viability 351 (see Tear *et al.* 2005) or metapopulation growth rate ($\lambda_M >$; Figure 3) are maximised. This is because these 352 methods are targeting the connectivity process per se but not the population outcomes explicitly. The use of 353 simulated annealing for optimising complex performance metrics (e.g. population viability) currently has 354 computational limitations. Therefore, the feasible solution to determine how well the final plan captures 355 metapopulation outcomes in the analysis, or to compare the performance between plans, is to undertake 356 post-hoc evaluations. The post-hoc analysis is structured as a sensitivity analysis, where multiple solutions 357 are generated and compared to assess their performance in achieving the chosen connectivity objective. These 358 solutions can be the result of varying Marxan parameters such as the conservation targets, the boundary 359 length modifier, connectivity metrics, and/or costs, or the post-hoc analysis can compare individual spatial 360 plans from the full ensemble created by the same input parameters and the simulated annealing process 361

³⁶² (Nicholson & Possingham 2006).

To illustrate this approach, we present a post-hoc sensitivity analysis to determine the optimal connectivity 363 target multiplier value, C, across four connectivity metrics (Figure 3). This same approach could also 364 be adapted for exploring the impact of using different methods, targets, thresholds, or data. We vary C365 in different Marxan scenarios using four different conservation features (in degree, betweenness centrality, 366 Google PageRank and avoidance area recipients) to determine the impact on cost, and metapopulation 367 growth. In this case, we used an in-reserve metapopulation growth rate greater than 1 (e.q. Figueira & 368 Crowder 2006; Hale, Treml & Swearer 2015; marxanconnect.ca) as our ecologically relevant conservation 369 objective. If the growth rate is lower than 1, the entire metapopulation would go extinct without external 370 supplementation; however, this method requires detailed biological information. In this example, using "in 371 degree" as a conservation feature increases the metapopulation growth rate linearly with cost, which has 372 limited applicability. Both "betweenness centrality" and "Google PageRank" perform quite well with the 373 latter being slightly better, likely because it considers the weight (*i.e.* strength) of the linkages. With "Google 374 PageRank" as a conservation feature and C = 3 (*i.e.* conservation target of 30%), this example species is 375 predicted to have a metapopulation growth rate > 1 and nearly the same level of growth as if the entire 376 ecosystem was protected. In this example, "avoidance area recipient" performs extremely poorly because: 377 1) there was no parameter or mechanism in the population model that represented a reason to avoid the 378 avoidance areas (e.g. impact of invasive species); and 2) the "avoidance area recipient" metric should always 379 be discretized (*i.e.* low values of "avoidance area recipient" are desirable) since using the continuous metric 380 would promote the selection of areas that receive the most propagules from the avoidance area. 381

In the second case, connectivity metrics can be calculated for individual conservation plans with varying spatial configurations that all meet the specified objective function (*i.e.* have the same input parameters). To identify the most dissimilar Marxan solutions in an analysis, a dissimilarity matrix (*i.e.* dendrogram) can be created using the Marxan cluster analysis function (Linke *et al.* 2011). The chosen metric can then be calculated for each plan to evaluate which spatial configuration best achieves the connectivity objective.

There are several different types of post-hoc assessments that can be performed, such as pattern-based assessments, evaluating metapopulation capacity, or using system models. In pattern-based assessments, for example, Krueck et al. (2017) developed metrics using local larval retention, import connectivity and export connectivity from connectivity matrices which can be calculated in a post-hoc analysis to evaluate how

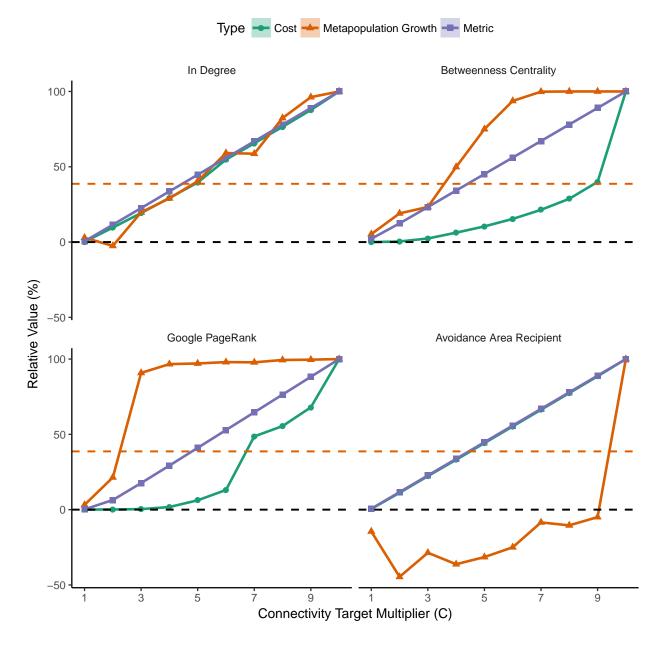


Figure 3: Connectivity target multiplier (C) sensitivity analysis, where the sum of the connectivity metric, estimated metapopulation growth rate (λ_M) , and cost of the selected network (mean \pm S.E., n = 10) are plotted as relative percentages of their maximum values. Four connectivity metrics are used as examples: In Degree, Betweenness Centrality, Google Page Rank and Avoidance Area Recipient. The black dashed line is the value of the metric, growth rate, and cost of the selected network if the connectivity based conservation feature is not included. The orange dashed line indicates where $\lambda_M = 1$, or the point at which the protected area network is self sustaining. The calculation of λ_M , included here for demonstration purposes only, is estimated using the leading eigenvalue of a theoretical connectivity matrix which includes theoretical fecundity and survival (*i.e.* flow matrix). Full details on the generation of this figure can be found on marxanconnect.ca.

Operationalizing connectivity in spatial planning

different conservation plans meet both conservation and fisheries objectives across protected and unprotected 391 sites. Graph theoretic metric(s) or those that better capture population persistence can also be used, but few 392 examples of their use in post-hoc analyses currently exist. In one example, Laita, Kotiaho, Monkkonen et al. 303 (2011) explored how network connectivity measures (*i.e.* correlation length, expected cluster size, landscape 394 coincidence probability, area-weighted flux, integral index of connectivity and probability of connectivity) 395 changed with the addition of woodland key habitats to reserve networks in Finland. However, they highlight 396 the need for a more detailed understanding of the caveats and justifications of these measures before they 397 can be used for conservation purposes. 398

If additional demographic information of species, such as survival and mortality, is known, then the suggested 399 course of action is to evaluate potential reserve networks using metapopulation models. With these models, 400 it is possible to make predictions regarding the ecological outcomes such as the probability of going extinct 401 in a certain time frame (Boyce 1992), the capacity to recover from a disturbance (Figueira & Crowder 402 2006), metapopulation lifetime (Kininmonth et al. 2010), probability of metapopulation extinction (Bode. 403 Burrage & Possingham 2008), or other possible ecologically relevant conservation objectives. While increasing 404 connectivity in reserve networks is generally desired; without models of metapopulation dynamics, connectivity 405 risks becoming a relatively meaningless objective like "percent area covered by protected area" (Tear et al. 406 2005). 407

Lastly, system models are designed to simulate one or more processes related to the conservation objectives 408 (e.q. prioritize stepping stones) or overall goals (e.q. population persistence). White et al. (2014) used 409 population models to compare the performance of Marxan solutions generated with and without the inclusion 410 of static larval connectivity information by calculating the equilibrium biomass (in and outside of protected 411 areas) and fishery yield of the different spatial configurations in California. Similarly, tools such as the 412 BESTMPA R package (Daigle, Monaco & Elgin 2017), allow users to test commercial fishery costs and 413 benefits from various spatial conservation scenarios using a spatially explicit metapopulation model that 414 interacts with fishing behaviour. 415

No matter what post-hoc analysis approach is used, selecting the most appropriate metric(s), understanding 416 the caveats of the metric, and making ad hoc assumptions on how the user expects the metric to perform 417 for the specified application is extremely important for interpreting and comparing the outcomes of these 418 different measures (Pascual-Hortal & Saura 2006; see Laita, Kotiaho & Mönkkönen 2011). In conservation 419 plans that incorporate existing protected areas, it is also important to evaluate the contribution of newly 420 selected sites to the conservation objective, which can be accomplished by performing the post-hoc analysis 421 with and without considering existing reserves (*i.e.* a gap analysis). This can reveal important information 422 on the performance of the existing reserve system and can help ensure complementarity between the existing 423 network and potential sites for protected area expansion. 424

425 Marxan Connect

Because we recognize that there is considerable investment in Marxan-based prioritization, Marxan Connect
was designed to help conservation practitioners incorporate connectivity into existing Marxan analyses. It
guides users through:

⁴²⁹ 1. Identifying and loading appropriate spatial data

Operationalizing connectivity in spatial planning

430	a. Planning units
431	b. Focus areas
432	c. Avoidance areas
433	2. Identifying and loading connectivity data
434	a. Demographic-based
435	b. Landscape-based
436	3. Calculates connectivity metrics or generates spatial dependencies
437	a. Conservation features method
438	b. Connectivity as spatial dependencies method
439	4. Optionally discretizes conservation features and exports Marxan files
440	5. Running Marxan

6. Evaluate results with basic plotting options

442 Marxan Connect allows users to export data products (e.g. connectivity metrics, Marxan files, etc.) at any of

the above steps to enable users to base their workflow in or outside Marxan Connect.

For the landscape connectivity approach, Marxan Connect calculates connectivity metrics from networks 444 based either on Euclidean distance or least-cost path between the centroid of planning units. However, other 445 software packages such as Circuitscape (McRae, Shah & Mohapatra 2009) and Conefor (Saura & Torné 2009) 446 currently provide a richer set of options and specialized methods. These software packages can be used to 447 generate custom conservation features or connectivity matrices both of which can then be used in Marxan 448 Connect. For example, one could generate a network using current density using Circuitscape and input the 449 resulting connectivity matrix into Marxan Connect to generate conservation features or spatial dependencies. 450 For a user opting to use non-Marxan spatial prioritization software such as Zonation (Lehtomäki & Moilanen 451

⁴⁵² 2013) or prioritizr (Hanson *et al.* 2017), there is a high degree of compatibility with Marxan Connect. The ⁴⁵³ approach of targeting conservation features is compatible with any spatial planning software. Certain software ⁴⁵⁴ packages such as prioritizr can read Marxan-formatted files directly; therefore, Marxan Connect could be ⁴⁵⁵ used with prioritizr to generate connectivity-related input files It also appears that modifying the boundary ⁴⁵⁶ definitions with connectivity data could be performed with prioritizr's "add_boundary_penalties" function. ⁴⁵⁷ This is in addition to prioritizr's "add_connected_constraints" function which tends to select unbroken ⁴⁵⁸ chains of physically linked planning units (Önal & Briers 2006).

459 Conclusions

The approaches for including connectivity in spatial planning are rapidly evolving and few "best practices" exist. Here, we provide some guidance on methods, data sources, and models, as well as a novel open-source tool to support these methods. However, connectivity-based conservation targets are ecologically meaningless unless placed in the context of broader ecologically relevant conservation objectives such as population viability, expected time to extinction, or metapopulation growth rate. Similarly, connectivity is usually only one criterion in planning, and will be considered alongside area-based targets, socio-economic goals, and multi-species requirements.

⁴⁶⁷ Connectivity is a complex topic with abundant terminology and a diversity of methods that require substantial
 ⁴⁶⁸ effort to understand and apply to spatial prioritization scenarios correctly (Beger et al. in prep). If

Operationalizing connectivity in spatial planning

connectivity is to widely inform protected area planning, communication channels between experts in the fields of connectivity and population dynamics and planners must be improved. The experts, in particular, should make their research outcomes more accessible to practitioners by providing openly available data and clarifying definitions, assumptions, and limitations. For example, the term "connectivity matrix", while central to the concept of connectivity, does not provide enough information to spatial planners or even to other connectivity experts to incorporate connectivity into spatial planning initiatives. With Marxan Connect, we hope to offer standardized methods and terminology to help close this research-implementation gap.

477 of protected areas, it does not guarantee that reserves will be "well connected". Only post-hoc evaluation of

the reserve design related to ecologically relevant conservation objective(s) can inform practitioners of the

⁴⁷⁹ resilience and persistence of targeted populations. However, the tools provided in Marxan Connect greatly

⁴⁸⁰ improve the likelihood that a selected reserve design will adequately meet those conservation objective(s).

481 Acknowledgements

Funding for the development of this software was provided by the Natural Sciences and Engineering Research 482 Council of Canada through the Canadian Healthy Oceans Network (NSERC NETGP 468437-14), and 483 the Discovery Grants program (NSERC DG 34851-2012), and the University of Queensland. Additional 484 contributions were provided by the University of Leeds, The Nature Conservancy, Dalhousie University, The 485 University of Melbourne, and the Australian Research Council - Centre of Excellence for Environmental 486 Decisions (CEED). This project builds on the existing Marxan (Ball, Possingham & Watts 2009) software 487 and would not be possible without the hard work of Ian Ball, Matt Watts, and Hugh Possingham. The authors also wish to thank Ryan Stanley, Marco Andrello, and Jo Clarke for constructive feedback on early 489 versions of the software or manuscript. 490

⁴⁹¹ Supplementary Material:

The source code for the software and website can be found at https://github.com/remi-daigle/MarxanConnect. There are repeated references in the tutorial and glossary section of marxanconnect.ca, the website may evolve as the software is improved. The original publication version of the website and app have been archived on Zenodo (Daigle et al. 2018 This will be archived and added to references upon acceptance of the manuscript)

496 Author's contributions

⁴⁹⁷ AM and MB conceived the application and acquired funding; RD developed the application and website;
⁴⁹⁸ RD, AM, AB, and MB worked on the initial development and early testing of the application. All authors
⁴⁹⁹ made significant contributions to the later development stages of the application and website. All authors
⁵⁰⁰ contributed critically to making improvements to the application, drafting the manuscript, and provided final
⁵⁰¹ approval for publication.

Operationalizing connectivity in spatial planning

502 **References**

- ⁵⁰³ Albert, C.H., Rayfield, B., Dumitru, M. & Gonzalez, A. (2017). Applying network theory to prioritize ⁵⁰⁴ multi-species habitat networks that are robust to climate and land-use change. *Conserv. Biol.*
- ⁵⁰⁵ Ardron, J.A., Possingham, H.P. & Klein, C.J. (2010). Marxan good practices handbook. Victoria, BC.
- ⁵⁰⁶ Álvarez-Romero, J.G., Munguía-Vega, A., Beger, M., Del Mar Mancha-Cisneros, M., Suárez-Castillo, A.N.,
- ⁵⁰⁷ Gurney, G.G., Pressey, R.L., Gerber, L.R., Morzaria-Luna, H.N., Reyes-Bonilla, H., Adams, V.M., Kolb, M.,
- ⁵⁰⁸ Graham, E.M., VanDerWal, J., Castillo-López, A., Hinojosa-Arango, G., Petatán-Ramírez, D., Moreno-Baez,
- ⁵⁰⁹ M., Godínez-Reyes, C.R. & Torre, J. (2018). Designing connected marine reserves in the face of global
- ⁵¹⁰ warming. *Glob. Chang. Biol.*, **24**, e671–e691.
- ⁵¹¹ Ball, I.R., Possingham, H.P. & Watts, M. (2009). Marxan and relatives: Software for spatial conservation
- ⁵¹² prioritisation. Spatial conservation prioritisation: quantitative methods and computational tools. Oxford
- ⁵¹³ University Press, Oxford, 185–195.
- Barnes, M.D., Glew, L., Wyborn, C. & Craigie, I.D. (2018). Prevent perverse outcomes from global protected
 area policy. *Nat Ecol Evol.*
- ⁵¹⁶ Beger, M., Grantham, H.S., Pressey, R.L., Wilson, K.A., Peterson, E.L., Dorfman, D., Mumby, P.J., Lourival,
- ⁵¹⁷ R., Brumbaugh, D.R. & Possingham, H.P. (2010a). Conservation planning for connectivity across marine,
- ⁵¹⁸ freshwater, and terrestrial realms. *Biol. Conserv.*, **143**, 565–575.
- ⁵¹⁹ Beger, M., Selkoe, K.A., Treml, E., Barber, P.H., Heyden, S. von der, Crandall, E.D., Toonen, R.J. & Riginos,
- ⁵²⁰ C. (2014). Evolving coral reef conservation with genetic information. Bull. Mar. Sci., **90**, 159–185.
- Beger, M., Linke, S., Watts, M., Game, E., Treml, E., Ball, I. & Possingham, H.P. (2010b). Incorporating asymmetric connectivity into spatial decision making for conservation: Asymmetric connectivity in conservation planning. *Conservation Letters*, **3**, 359–368.
- Beger, M., McGowan, J., Treml, E.A., Green, A.L., White, A.T., Wolff, N.H., Klein, C.J., Mumby, P.J. &
- ⁵²⁵ Possingham, H.P. (2015). Integrating regional conservation priorities for multiple objectives into national
- ⁵²⁶ policy. Nat. Commun., 6, 8208.
- ⁵²⁷ Beyer, H.L., Dujardin, Y., Watts, M.E. & Possingham, H.P. (2016). Solving conservation planning problems ⁵²⁸ with integer linear programming. *Ecol. Modell.*, **328**, 14–22.
- Bode, M., Bode, L. & Armsworth, P.R. (2006). Larval dispersal reveals regional sources and sinks in the
 great barrier reef. Mar. Ecol. Prog. Ser., 308, 17–25.
- ⁵³¹ Bode, M., Burrage, K. & Possingham, H.P. (2008). Using complex network metrics to predict the persistence ⁵³² of metapopulations with asymmetric connectivity patterns. *Ecol. Modell.*, **214**, 201–209.
- Boyce, M.S. (1992). Population viability analysis. Annu. Rev. Ecol. Syst., 23, 481–506.
- Bracis, C.I. (2015). Incorporating cognition into models of animal movement and predator-prey interaction.
 PhD thesis thesis,
- ⁵³⁶ Brown, C.J., Harborne, A.R., Paris, C.B. & Mumby, P.J. (2016). Uniting paradigms of connectivity in marine

- ⁵³⁷ ecology. *Ecology*, **97**, 2447–2457.
- ⁵³⁸ Bunn, A.G., Urban, D.L. & Keitt, T.H. (2000). Landscape connectivity: A conservation application of graph
- 539 theory. J. Environ. Manage., 59, 265–278.
- 540 Cagnacci, F., Boitani, L., Powell, R.A. & Boyce, M.S. (2010). Animal ecology meets GPS-based radiotelemetry:
- ⁵⁴¹ A perfect storm of opportunities and challenges. *Philos. Trans. R. Soc. Lond. B Biol. Sci.*, **365**, 2157–2162.
- ⁵⁴² Chetkiewicz, C.-L.B. & Boyce, M.S. (2009). Use of resource selection functions to identify conservation ⁵⁴³ corridors. J. Appl. Ecol., 46, 1036–1047.
- ⁵⁴⁴ Chollett, I., Garavelli, L., O'Farrell, S., Cherubin, L., Matthews, T.R., Mumby, P.J. & Box, S.J. (2017). A
- genuine win-win: Resolving the 'conserve or catch' conflict in marine reserve network design. *Conservation*
- ⁵⁴⁶ Letters, **10**, 555–563.
- ⁵⁴⁷ Cowen, R.K., Lwiza, K.M., Sponaugle, S., Paris, C.B. & Olson, D.B. (2000). Connectivity of marine ⁵⁴⁸ populations: Open or closed? *Science*, **287**, 857–859.
- ⁵⁴⁹ Daigle, R.M. & Metaxas, A. (2012). Modeling of the larval response of green sea urchins to thermal ⁵⁵⁰ stratification using a random walk approach. *J. Exp. Mar. Bio. Ecol.*, **438**, 14–23.
- ⁵⁵¹ Daigle, R.M., Chassé, J. & Metaxas, A. (2016). The relative effect of behaviour in larval dispersal in a low ⁵⁵² energy embayment. *Prog. Oceanogr.*, **144**, 93–117.
- ⁵⁵³ Daigle, R.M., Monaco, C.J. & Elgin, A.K. (2017). An adaptable toolkit to assess commercial fishery costs ⁵⁵⁴ and benefits related to marine protected area network design. *F1000Res.*, **4**.
- ⁵⁵⁵ Dunn, J.E. & Gipson, P.S. (1977). Analysis of radio telemetry data in studies of home range. *Biometrics*, 33,
 ⁵⁵⁶ 85–101.
- ⁵⁵⁷ D'Aloia, C.C., Daigle, R.M., Côté, I.M., Curtis, J.M.R., Guichard, F. & Fortin, M.-J. (2017). A multiple-⁵⁵⁸ species framework for integrating movement processes across life stages into the design of marine protected ⁵⁵⁹ areas. *Biol. Conserv.*, **216**, 93–100.
- Fall, A., Fortin, M.-J., Manseau, M. & O'Brien, D. (2007). Spatial graphs: Principles and applications for
 habitat connectivity. *Ecosystems*, 10, 448–461.
- Figueira, W.F. & Crowder, L.B. (2006). Defining patch contribution in source-sink metapopulations: The importance of including dispersal and its relevance to marine systems. *Popul. Ecol.*, **48**, 215–224.
- Hale, R., Treml, E.A. & Swearer, S.E. (2015). Evaluating the metapopulation consequences of ecological
 traps. *Proc. Biol. Sci.*, 282, 20142930.
- Hand, B.K., Lowe, W.H., Kovach, R.P., Muhlfeld, C.C. & Luikart, G. (2015). Landscape community genomics:
 Understanding eco-evolutionary processes in complex environments. *Trends Ecol. Evol.*, **30**, 161–168.
- Hanson, J.O., Schuster, R., Morrell, N., Strimas-Mackey, M., Watts, M.E., Arcese, P., Bennett, J. &
- ⁵⁶⁹ Possingham, H.P. (2017). Prioritizr: Systematic conservation prioritization in R.
- Harrison, H.B., Williamson, D.H., Evans, R.D., Almany, G.R., Thorrold, S.R., Russ, G.R., Feldheim, K.A.,
- ⁵⁷¹ Herwerden, L. van, Planes, S., Srinivasan, M., Berumen, M.L. & Jones, G.P. (2012). Larval export from

- marine reserves and the recruitment benefit for fish and fisheries. Curr. Biol., 22, 1023–1028.
- ⁵⁷³ Hussey, N.E., Kessel, S.T., Aarestrup, K., Cooke, S.J., Cowley, P.D., Fisk, A.T., Harcourt, R.G., Holland,
- 574 K.N., Iverson, S.J., Kocik, J.F., Mills Flemming, J.E. & Whoriskey, F.G. (2015). ECOLOGY. aquatic animal
- ⁵⁷⁵ telemetry: A panoramic window into the underwater world. *Science*, **348**, 1255642.
- ⁵⁷⁶ Iwamura, T., Possingham, H.P., Chadès, I., Minton, C., Murray, N.J., Rogers, D.I., Treml, E.A. & Fuller, R.A.
- ⁵⁷⁷ (2013). Migratory connectivity magnifies the consequences of habitat loss from sea-level rise for shorebird
- ⁵⁷⁸ populations. Proc. Biol. Sci., 280, 20130325.
- Jonsson, P.R., Nilsson Jacobi, M. & Moksnes, P.-O. (2016). How to select networks of marine protected areas
- for multiple species with different dispersal strategies (L. Beaumont, Ed.). Divers. Distrib., 22, 161–173.
- Kininmonth, S., Drechsler, M., Johst, K. & Possingham, H.P. (2010). Metapopulation mean life time within
 complex networks. *Mar. Ecol. Prog. Ser.*, 417, 139–149.
- Koen, E.L., Bowman, J., Sadowski, C. & Walpole, A.A. (2014). Landscape connectivity for wildlife:
 Development and validation of multispecies linkage maps (A. Tatem, Ed.). *Methods Ecol. Evol.*, 5, 626–633.
- Kool, J.T., Moilanen, A. & Treml, E.A. (2013). Population connectivity: Recent advances and new
 perspectives. Landsc. Ecol., 28, 165–185.
- 587 Krueck, N.C., Ahmadia, G.N., Green, A., Jones, G.P., Possingham, H.P., Riginos, C., Treml, E.A. & Mumby,
- P.J. (2017). Incorporating larval dispersal into MPA design for both conservation and fisheries. *Ecol. Appl.*,
 27, 925–941.
- Laita, A., Kotiaho, J.S. & Mönkkönen, M. (2011). Graph-theoretic connectivity measures: What do they tell us about connectivity? *Landsc. Ecol.*, **26**, 951–967.
- Lehtomäki, J. & Moilanen, A. (2013). Methods and workflow for spatial conservation prioritization using
 zonation. *Environmental Modelling & Software*, 47, 128–137.
- Linke, S., Watts, M., Stewart, R. & Possingham, H.P. (2011). Using multivariate analysis to deliver conservation planning products that align with practitioner needs. *Ecography*, **34**, 203–207.
- Lowe, W.H. & Allendorf, F.W. (2010). What can genetics tell us about population connectivity? Mol. Ecol.,
 19, 3038–3051.
- Magris, R.A., Andrello, M., Pressey, R.L., Mouillot, D., Dalongeville, A., Jacobi, M.N. & Manel, S. (2018).
- ⁵⁹⁹ Biologically representative and well-connected marine reserves enhance biodiversity persistence in conservation ⁶⁰⁰ planning. *CONSERVATION LETTERS*, **96**, e12439.
- Magris, R.A., Treml, E.A., Pressey, R.L. & Weeks, R. (2016). Integrating multiple species connectivity and habitat quality into conservation planning for coral reefs. *Ecography*, **39**, 649–664.
- Maina, J.M., Jones, K.R., Hicks, C.C., McClanahan, T.R., Watson, J.E.M., Tuda, A.O. & Andréfouët, S.
- (2015). Designing Climate-Resilient marine protected area networks by combining remotely sensed coral reef

- habitat with coastal Multi-Use maps. Remote Sensing, 7, 16571–16587.
- Manel, S. & Holderegger, R. (2013). Ten years of landscape genetics. Trends Ecol. Evol., 28, 614–621.
- ⁶⁰⁷ Margules, C.R. & Pressey, R.L. (2000). Systematic conservation planning. *Nature*, **405**, 243–253.
- Marrotte, R.R., Bowman, J., Brown, M.G.C., Cordes, C., Morris, K.Y., Prentice, M.B. & Wilson, P.J. (2017).
- ⁶⁰⁹ Multi-species genetic connectivity in a terrestrial habitat network. *Mov Ecol*, **5**, 21.
- Mazor, T., Beger, M., McGowan, J., Possingham, H.P. & Kark, S. (2016). The value of migration information
- ⁶¹¹ for conservation prioritization of sea turtles in the mediterranean: Conservation planning of migratory species.
- ⁶¹² Glob. Ecol. Biogeogr., **25**, 540–552.
- 613 McGowan, J., Smith, R.J., Di Marco, M., Clarke, R.H. & Possingham, H.P. (2017a). An evaluation of marine
- ⁶¹⁴ important bird and biodiversity areas in the context of spatial conservation prioritization. *Conservation*
- 615 Letters.
- 616 McGowan, J., Beger, M., Lewison, R.L., Harcourt, R., Campbell, H., Priest, M., Dwyer, R.G., Lin, H.-Y.,
- Lentini, P., Dudgeon, C., McMahon, C., Watts, M. & Possingham, H.P. (2017b). Integrating research using
- animal-borne telemetry with the needs of conservation management (J. du Toit, Ed.). J. Appl. Ecol., 54, 423–429.
- McNeil, F.I. & Crossman, E.J. (1979). Fin clips in the evaluation of stocking programs for muskellunge, esox masquinongy. *Trans. Am. Fish. Soc.*, **108**, 335–343.
- McRae, B.H. & Nürnberger, B. (2006). ISOLATION BY RESISTANCE. Evolution, 60, 1551–1561.
- McRae, B.H., Shah, V.B. & Mohapatra, T.K. (2009). Circuitscape user guide. The University of California,
 Santa Barbara.
- Melià, P., Schiavina, M., Rossetto, M., Gatto, M., Fraschetti, S. & Casagrandi, R. (2016). Looking for hotspots of marine metacommunity connectivity: A methodological framework. *Sci. Rep.*, **6**, 23705.
- Metaxas, A. (2001). Behaviour in flow: Perspectives on the distribution and dispersion of meroplanktonic larvae in the water column. *Can. J. Fish. Aquat. Sci.*, **58**, 86–98.
- Metaxas, A. & Saunders, M. (2009). Quantifying the 'bio-' components in biophysical models of larval
 transport in marine benthic invertebrates: Advances and pitfalls. *Biol. Bull.*, 216, 257–272.
- Minor, E.S., McDonald, R.I., Treml, E.A. & Urban, D.L. (2008). Uncertainty in spatially explicit population
 models. *Biol. Conserv.*, 141, 956–970.
- Moilanen, A., Wilson, K.A. & Possingham, H. (2009). Spatial conservation prioritization: Quantitative
 methods and computational tools. Oxford University Press.
- Mora, C., Andrèfouët, S., Costello, M.J., Kranenburg, C., Rollo, A., Veron, J., Gaston, K.J. & Myers, R.A.
- (2006). Coral reefs and the global network of marine protected areas. Science, **312**, 1750–1751.
- ⁶³⁷ Nicholson, E. & Possingham, H.P. (2006). Objectives for Multiple-Species conservation planning. *Conserv.* ⁶³⁸ *Biol.*, **20**, 871–881.
- ⁶³⁹ Önal, H. & Briers, R.A. (2006). Optimal selection of a connected reserve network. Oper. Res., 54, 379–388.
- ⁶⁴⁰ Palumbi, S.R. (2003). Population genetics, demographic connectivity, and the design of marine reserves. *Ecol.*

- ⁶⁴¹ Appl., **13**, S146–S158.
- Paris, C.B., Chérubin, L.M. & Cowen, R.K. (2007). Surfing, spinning, or diving from reef to reef: Effects on
- ⁶⁴³ population connectivity. Mar. Ecol. Prog. Ser., **347**, 285–300.
- Pascual-Hortal, L. & Saura, S. (2006). Comparison and development of new graph-based landscape connectivity
- indices: Towards the priorization of habitat patches and corridors for conservation. Landsc. Ecol., 21, 959–967.
- ⁶⁴⁶ Pereira, J., Saura, S. & Jordán, F. (2017). Single-node vs. multi-node centrality in landscape graph analysis:
- ⁶⁴⁷ Key habitat patches and their protection for 20 bird species in NE spain (F. Parrini, Ed.). Methods Ecol.
- ⁶⁴⁸ Evol., **8**, 1458–1467.
- Pritchard, J.K., Stephens, M. & Donnelly, P. (2000). Inference of population structure using multilocus
 genotype data. *Genetics*, 155, 945–959.
- Rayfield, B., Fortin, M.-J. & Fall, A. (2010). The sensitivity of least-cost habitat graphs to relative cost surface values. *Landsc. Ecol.*, **25**, 519–532.
- Ricketts, T.H. (2001). The matrix matters: Effective isolation in fragmented landscapes. Am. Nat., 158,
 87–99.
- Roughgarden, J., Gaines, S. & Possingham, H. (1988). Recruitment dynamics in complex life cycles. Science,
 241, 1460–1466.
- ⁶⁵⁷ Runge, C.A., Tulloch, A.I.T., Possingham, H.P., Tulloch, V.J.D. & Fuller, R.A. (2016). Incorporating ⁶⁵⁸ dynamic distributions into spatial prioritization (N. Roura-Pascual, Ed.). *Divers. Distrib.*, **22**, 332–343.
- Saenz-Agudelo, P., Jones, G.P., Thorrold, S.R. & Planes, S. (2009). Estimating connectivity in marine
 populations: An empirical evaluation of assignment tests and parentage analysis under different gene flow
 scenarios. *Mol. Ecol.*, 18, 1765–1776.
- ⁶⁶² Saura, S. & Pascual-Hortal, L. (2007). A new habitat availability index to integrate connectivity in landscape
- conservation planning: Comparison with existing indices and application to a case study. Landsc. Urban
 Plan., 83, 91–103.
- 665 Saura, S. & Torné, J. (2009). Conefor sensinode 2.2: A software package for quantifying the importance of
- habitat patches for landscape connectivity. Environmental Modelling & Software, 24, 135–139.
- ⁶⁶⁷ Schumaker, N.H. (1996). Using landscape indices to predict habitat connectivity. *Ecology*, 77, 1210–1225.
- 668 Scott, T.G. (1942). Ear tags on mice. J. Mammal., 23, 339.
- ⁶⁶⁹ Smith, J. & Metaxas, A. (2018). A decision tree that can address connectivity in the design of marine ⁶⁷⁰ protected area networks (MPAn). *Mar. Policy*, **88**, 269–278.
- Tear, T.H., Kareiva, P., Angermeier, P.L., Comer, P., Czech, B., Kautz, R., Landon, L., Mehlman, D.,
- ⁶⁷² Murphy, K., Ruckelshaus, M., Scott, J.M. & Wilhere, G. (2005). How much is enough? The recurrent problem
- of setting measurable objectives in conservation. *Bioscience*, **55**, 835–849.
- Thomas, J.W. & Marburger, R.G. (1964). Colored leg markers for wild turkeys. J. Wildl. Manage., 28,
 552–555.
- ⁶⁷⁶ Tischendorf, L. & Fahrig, L. (2000). On the usage and measurement of landscape connectivity. Oikos, 90,

- 677 **7**-19.
- ⁶⁷⁸ Treml, E.A., Ford, J.R., Black, K.P. & Swearer, S.E. (2015). Identifying the key biophysical drivers,
- ⁶⁷⁹ connectivity outcomes, and metapopulation consequences of larval dispersal in the sea. Mov Ecol, 3, 17.
- ⁶⁸⁰ Twigg, G.I. (1978). Marking mammals by tissue removal. Animal marking, pp. 109–118. Palgrave, London.
- ⁶⁸¹ UNEP. (2010). TARGET 11 technical rationale extended (provided in document COP/10/INF/12/Rev.1).
- 682 Convention on Biological Diversity.
- ⁶⁸³ Urban, D. & Keitt, T. (2001). LANDSCAPE CONNECTIVITY: A GRAPH-THEORETIC PERSPECTIVE.
 ⁶⁸⁴ Ecology, 82, 1205–1218.
- Voegeli, F.A., Smale, M.J., Webber, D.M., Andrade, Y. & O'Dor, R.K. (2001). Ultrasonic telemetry, tracking
- and automated monitoring technology for sharks. The behavior and sensory biology of elasmobranch fishes:
- An anthology in memory of donald richard nelson, pp. 267–282. Developments in environmental biology of
- 688 fishes. Springer, Dordrecht.
- Weeks, R. (2017). Incorporating seascape connectivity in conservation prioritisation. PLoS One, 12, e0182396.
- ⁶⁹⁰ White, J.W., Schroeger, J., Drake, P.T. & Edwards, C.A. (2014). The value of larval connectivity information ⁶⁹¹ in the static optimization of marine reserve design. *Conservation Letters*, **7**, 533–544.
- Whitfield Gibbons, J. & Andrews, K.M. (2004). PIT tagging: Simple technology at its best. *Bioscience*;
 Oxford, 54, 447–454.
- Wilson, K.A., Underwood, E.C., Morrison, S.A., Klausmeyer, K.R., Murdoch, W.W., Reyers, B., Wardell-
- Johnson, G., Marquet, P.A., Rundel, P.W., McBride, M.F., Pressey, R.L., Bode, M., Hoekstra, J.M.,
- ⁶⁹⁶ Andelman, S., Looker, M., Rondinini, C., Kareiva, P., Shaw, M.R. & Possingham, H.P. (2007). Conserving
- ⁶⁹⁷ biodiversity efficiently: What to do, where, and when. *PLoS Biol.*, **5**, e223.
- Wilson, K.A., Cabeza, M. & Klein, C.J. (2009). Fundamental concepts of spatial conservation prioritization.
 Spatial Conservation Prioritization: quantitative methods and computational tools. Oxford, UK2009.
- Zeller, K.A., Jennings, M.K., Vickers, T.W., Ernest, H.B., Cushman, S.A. & Boyce, W.M. (2018). Are all
- ⁷⁰¹ data types and connectivity models created equal? Validating common connectivity approaches with dispersal
- 702 data (J. Bolliger, Ed.). Divers. Distrib., 54, 412.
- 703 Zeller, K.A., McGarigal, K. & Whiteley, A.R. (2012). Estimating landscape resistance to movement: A
- ⁷⁰⁴ review. Landsc. Ecol., 27, 777–797.