PERSPECTIVE

Population connectivity: recent advances and new perspectives

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Abstract Connectivity is a vital component of metapopulation and landscape ecology, influencing fundamental processes such as population dynamics, evolution, and community responses to climate change. Here, we review ongoing developments in connectivity science, providing perspectives on recent advances in identifying, quantifying, modelling and analysing connectivity, and highlight new applications for conservation. We also address ongoing challenges

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Department of Zoology, University of Melbourne, Melbourne, VIC 3010, Australia for connectivity research, explore opportunities for addressing them and highlight potential linkages with other fields of research. Continued development of connectivity science will provide insights into key aspects of ecology and the evolution of species, and will also contribute significantly towards achieving more effective conservation outcomes.

Keywords Dispersal · Spatial ecology · Review · Tracking · Population genetics · Modeling · Conservation · Management

Introduction

Connectivity has rapidly grown into a field of great interest for scientists and conservation managers (Crooks and Sanjayan 2006; Claudet 2011; Liu et al. 2011; Luque et al. 2012; Rayfield et al. 2011). Connectivity research links a wide variety of subjects in ecology and evolution, including dispersal and migration (Baguette and Van Dyck 2007), the development of population genetic structure (Kool et al. 2011), sourcesink dynamics (Figueira and Crowder 2006) and potential responses to climate change (Munday et al. 2009; Wasserman et al. 2012). Connectivity also affects conservation decisions involving aspects of reserve network design (Cerdeira et al. 2010), restoration (Raeymaekers et al. 2008), controlling invasive species (Hulme 2009), and administration of transboundary resources (Chester 2006; Treml and Halpin in press). Over time, perspectives on connectivity have evolved considerably, departing from the view that populations are uniformly distributed and panmictic, towards a more nuanced notion of networks of patches and demes often engaged in self-replenishment, as well as dynamic and asymmetric exchanges. Yet, despite the attention connectivity has received, much work is still required in order to understand its underlying causes and consequences, and to incorporate our understanding of connectivity into operational management strategies. Although general reviews of population connectivity have appeared elsewhere (Crooks and Sanjayan 2006; Hilty et al. 2006; Cowen and Sponaugle 2009), we here focus on trends in connectivity research, highlight ongoing developments, technologies and applications, and discuss emerging challenges and opportunities.

Conceptualizing connectivity

Initially, connectivity was described in the terrestrial context as "the degree to which the landscape facilitates or impedes movement among resource patches" (Taylor et al. 1993). Over time however, different research perspectives and operational needs have led to alternative ways of defining connectivity (Panel 1). For example, Pineda et al. (2007) distinguish between transport, dispersal and connectivity in marine systems (connectivity being a function of transport, larval survival, settlement and post-larval survival), Pringle (2003) addresses the importance of hydrologic connectivity, and Lowe and Allendorf (2010) discuss aspects of demographic and genetic connectivity. A challenge in population connectivity research lies in defining what constitutes a population, subpopulation or patch, and in semi-continuous habitats, distinguishing patches may be difficult and/or counterproductive. Consequently connectivity, however defined, may vary greatly, depending not only on the abundance and density of individuals, but also on habitat characteristics and the spatial and temporal scales of interest. For example, low levels of exchange might not be significant in a short-term demographic context, but might be vital for maintaining genetic diversity (Lowe and Allendorf 2010). Furthermore, population connectivity is not limited to the movement of individuals, but can also be defined according to gene flow (Hedgecock et al. 2007), or even more abstract concepts, such as the transfer of information or behaviour (Ahmad and Teredesai 2006).

Although population connectivity can be interpreted in different ways under different circumstances, the approaches share a fundamental property in common. In all cases, connectivity corresponds to a structured set of relationships between spatially and/or temporally distinct entities, or put another way connectivity is the outcome of dependencies between populations or individuals. Exploring the nature of these dependencies and relationships, as well as the consequences of their form, is what underpins connectivity research.

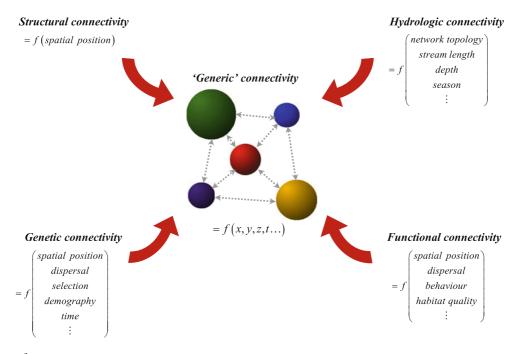
Empirically quantifying population connectivity

Direct methods

A variety of different techniques have been used to directly measure connectivity between populations (Table 1). Tracking organisms through field observation is the most basic means of evaluating population connectivity, but this can be challenging when monitoring large populations or broad spatial extents, particularly when the organisms being observed are small or cryptic. To address this difficulty, markrecapture techniques have been used extensively in the past (Webster et al. 2002) and continue to be the primary means of assessing connectivity today (Jacobson and Peres-Neto 2010), providing estimates of population size and movement patterns, often in a habitat-specific context. For organisms that are too small or fragile to carry physical tags, chemical-based analyses are often used (Rubenstein and Hobson 2004; Hobson 2008; Pauli et al. 2009; Durbec et al. 2010), particularly for aquatic populations. Researchers have been able to identify probable source populations based on chemical signatures present in otoliths and statoliths (sensory bones/stones found in fish and invertebrates respectively) (Thorrold et al. 2007; Woods et al. 2010), and more recently, artificial tagging techniques have been used to label and identify parents and progeny of marine species (Almany et al. 2007). Similar approaches have been used to study terrestrial mammal and bird populations (West et al. 2006; Newsome et al. 2007; Faaborg et al. 2010).

Panel 1: Deconstructing connectivity

Existing definitions of connectivity lie along a continuum ranging from the simple – e.g. examining aspects such as patch adjacency or straight-line relationships between objects, to the very complex – e.g. incorporating factors such as mobility, perceptual ability, and species-specific biological considerations. These definitions are not necessarily mutually exclusive and may overlap with one another (e.g. it is possible to consider functional genetic connectivity). In all cases however, the idea of connectivity being a structured set of inter-population relationships remains consistent.



f is a function linking the input variables to the output connectivity values. x,y,z,t... is a set of arbitrary variables.

Rather than pursuing a single 'ideal' operational definition of connectivity, the following questions can instead be used to distinguish between different analytical frameworks, and to clarify various study objectives:

- (1) How is landscape structure characterized? Is a simple, binary habitat/non-habitat classification used, or is continuous or semi-continuous variation in quality allowed?
- (2) How are distances between landscape elements determined? For example, are they Euclidian distances, leastcost paths, or more complex pathways (e.g. accounting for mortality, competition and selection, reproductive success etc.)?
- (3) Are distances or potential movement pathways interpreted in a species- or individual-specific manner?
- (4) Is connectivity being considered from the perspective of a patch (landscape element), between a pair of landscape elements, or as an aggregate property of the entire landscape?
- (5) Is connectivity used as a stand-alone quantity, as an explanatory variable of a statistical model, or as a part of a full population model, allowing Population Viability Analysis (PVA)-like evaluations of population survival?

Table 1	Methods for er	mpirically quar	Table 1 Methods for empirically quantifying population connectivity	ion connectivi	ity					
	Method	Extent (typical order of magnitude)	l order of	Resolution (ty magnitude)	Resolution (typical order of magnitude)	Environment	Data type	Sample scale	Constraints	Selected references
		Spatial	Temporal	Spatial	Temporal					
Direct methods	Visual observation	Hundreds of metres or less	Study duration, typically daily or aggregated over time	Metres or less	Instant	Any	Any	Individuals	Sampling effort, species visibility at observation distance. Data recording and processing effort.	Sutherland (2006)
	Mark- recapture	Conditioned on species mobility, typically kilometres	Study duration, conditioned on species mobility and life span.	Conditional on sampling effort and species behaviour	Conditional on sampling effort and species behaviour	Any	Displacement Individuals	Individuals	Tagging and sampling effort, organism must be able to carry the marker, and able to withstand handling. Isotopes require hard structures.	Sutherland (2006) Munro et al. (2009), and Williamson (2009)
	Radio tracking	Hundreds of metres to kilometres	Study duration, typically daily or aggregated over time	Metres	Seconds to Days	Typically terrestrial, some aquatic using sound	Points or routes	Individuals	Tagging and sampling effort, organism must be able to carry the transmitter, and able to withstand handling. Signal transmission is also necessary.	Millspaugh and Marzluff (2001)
	Geographic Positioning System (GPS) sensor	Global	Days to months	Metres	Seconds to Days	Terrestrial	Points or routes	Individuals	Tagging effort, organism must be capable of bearing the transmitter, and strong enough to withstand handling. Signal transmission is also necessary.	Urbano et al. (2010) and Recio et al. (2011)
	Pop-up Satellite Archival Tag (PSAT)	Global	Days to months	Metres	Instant to Days	Aquatic	Points or routes	Individuals	Tagging effort, organism must be capable of bearing the transmitter, and strong enough to withstand handling. Signal transmission is also necessary for collection.	Patterson et al. (2008a) and Musyl et al. (2011)
	Satellite/ aerial surveys	Hundreds of metres per frame to global coverage	Varies by platform and program duration	Centimetres to metres	Varies by platform	Typically terrestrial	Points or patches	Population	Organism must be visible from a vertical aerial perspective.	Turner et al. (2003) Fleming and Tracey (2008) and Wang et al. (2010)
	Radar	Kilometres	Years (station lifetime)	Sub-metre	Seconds	Terrestrial, aquatic surface	Points or patches	Population	Typically requires a radar installation, though portable radar has been developed.	Ovaskainen et al. (2008c) and Randall et al. (2011)

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	Method	Extent (typi magnitude)	cal order of	Resolution (typical order of magnitude)	cal order of	Environment Data type	Data type	Sample scale	Constraints	Selected references
		Spatial	Spatial Temporal	Spatial	Temporal					
Indirect methods	Indirect Parentage methods analysis	Any	Single generation	Parental home/ breeding range size	Single generation	Any	Displacement Individuals likelihood within population	Individuals within population	Requires comprehensive sampling of the population. Adequate genetic markers must be developed.	Jones et al. (2005) and Jones et al. (2010)
	Genetic assignment	Any	Multiple generations	Subpopulation	Multiple generations	Any	Displacement likelihood	Population	Multiple, meaningful markers must be developed. Population assignment depends on genetic models and their associated assumptions.	Berry et al. (2004) and Waples and Gaggiotti (2006)
	Genetic similarity	Any	Multiple generations	Subpopulation Multiple generati	Multiple generations	Any	Displacement as genetic distance	Population	Multiple, meaningful markers must be developed. Similarity measures depend on genetic models and associated assumptions.	Jaquiéry et al. (2011) and Foster et al. (2012)

This table summarizes and compares different methods of measuring population connectivity. Direct methods are techniques that can detect the physical movement of individuals between populations. indirect methods make strong inferences regarding movement between populations Extern The typical order of magnitude of the extent of the study area covered by the technique. In spatial terms, this corresponds to the bounding box, or areal range that the technique operates at. In temporal terms, this is the duration over which such studies typically occur Resolution: The typical order of magnitude at which the technique is able to resolve differences between objects. In spatial terms, this corresponds to the pixel size of an image, or the accuracy of determining the position of a tracking device. In temporal terms, this is the minimum time difference that is possible between successive measurements

Environment The environment for which the technique is primarily suited: terrestrial, aquatic (marine or freshwater), or any

Data type: The geometry typically obtained by this technique, point locations, line tracks, patches or displacements. Displacements only measure the differences between start and end locations with no regard to processes occurring between the two Sample scale The level at which the sample data is typically collected. Individual means that individual organisms are marked, tagged, observed. Population means that multiple individuals are monitored simultaneously Constraints Limitations associated with the use of the technique. Note that indirect methods place additional demands in terms of processing requirements (e.g. genetic analysis) and technical aspects of data analysis

Selected references Reviews or illustrative applications of the technique

Recently, there has been a dramatic expansion in the ability to remotely monitor animal movements, physiological measurements and associated environmental data (biologging-Rutz and Hays 2009). Large quantities of data are becoming available from these efforts, and are proving invaluable for understanding animal migration, behaviour and ecology for many species at a greater level of detail and at a broader range of scales than previously possible. For example, pop-up satellite archival tags (PSATs) have been used to track large numbers of pelagic predators and sea turtles over extremely large distances (Rutz and Hays 2009). Radar technology is also now providing ways of comprehensively tracking large collections of small and delicate organisms such as butterflies (Ovaskainen et al. 2008c).

Major advances are also being made towards the development of large-scale, fine-grained sensor networks for monitoring animal movement (Porter et al. 2005; Borgman et al. 2007). Cameras and environmental sensors linked to wireless communication systems provide a means of automatically detecting fine-scale movement patterns in real time (Hamilton et al. 2007; Kays et al. 2009). These data can then be filtered and queried to identify and summarize mass occurrences of movement events. These types of networks have been applied in terrestrial environments for monitoring tiger populations (Karanth et al. 2006), and are becoming increasingly prevalent in coastal and ocean systems as well (Martin Taylor 2009). Imaging systems are also being developed that are capable of capturing images of microscopic plankton (Cowen and Guigand 2008), which can then be processed using algorithms to identify species and characterize their spatial distribution within the water column (Tsechpenakis et al. 2007). Coupling these technologies with ocean sensing grids would provide an unprecedented opportunity to monitor the real-time spatial characteristics of connectivity in aquatic environments.

Connectivity studies have been greatly assisted by the extensive development of GIS and remote-sensing data, however obtaining comprehensive and simultaneous data with a high degree of resolution remains challenging, especially for features that are not highly visible. Further development of remote-sensing platforms will be necessary, as well as comprehensive field research for ground-truthing remotely sensed and modelled data. It will also be important to collect timeseries data to assess the effects of temporal changes in connectivity (e.g. successional dynamics, anthropogenic change). Understanding temporal aspects of connectivity will be key for understanding species responses (such as range expansion) to progressive habitat fragmentation and climate change (Heller and Zavaleta 2009).

Indirect methods

Direct tracking of organisms provides the most accurate information on animal movement over demographic time-scales, but over longer time scales (e.g. evolutionary) a different approach is needed. Population genetics provides a means of assessing connectivity integrated over many generations, compressing time scales that otherwise would not be observable. Rapid expansion in the availability of genetic markers (Parker et al. 1998; Broquet and Petit 2009; Francesco Ficetola and Bonin 2011) and dramatic increases in computing power have opened up new opportunities for identifying patterns of genetic connectivity (Balkenhol et al. 2009; Lowe and Allendorf 2010). Restriction fragment-length polymorphism (RFLP) and mtDNA analyses have largely given way to variable-number tandem repeat (VNTR-e.g. microsatellites) analysis, and with drastic decreases in both the cost and amount of time required to carry out genetic research, large repositories of population genetic data are becoming available for a variety of species and locations (Storfer et al. 2010). These data can be used to examine isolation by distance patterns, to back-trace migration paths and to identify potential stepping-stone populations using specialized software programs (such as MIGRATE-N-Beerli and Palczewski 2010). Assignment tests are also being used to identify barriers, spatial structuring and recent migration patterns (Excoffier and Heckel 2006; Faubet and Gaggiotti 2008). These developments have led to the expansive growth and development of the fields of landscape and seascape genetics (Manel et al. 2003; Holderegger and Wagner 2006; Selkoe et al. 2008).

Parentage analysis is increasingly being used as a means of assessing demographic connectivity over the time-scale of a single generation (Jones and Ardren 2003; Jones et al. 2005; Planes et al. 2009; Jones et al. 2010). This is typically achieved by comprehensively

sampling the population, obtaining molecular marker frequencies (e.g. microsatellites), numerically simulating progeny, and using log-likelihood scores to match the actual progeny with the most likely parent or parent pair (Saenz-Agudelo et al. 2009). Parentage analysis offers tremendous benefits in that it provides quantitative and unambiguous measures of connectivity (Harrison et al. 2012), as well as a strong means of validating other means of assessing connectivity (Berumen et al. 2010). However, the requirement that the population be comprehensively sampled (Marshall et al. 1998) makes large-scale studies difficult, or in many cases, impossible. Methods have been developed to help account for incomplete sampling (Duchesne et al. 2005; Mobley 2011), but for the time being, this approach will be generally limited to smaller, mostlyclosed populations or small groups of populations.

With major developments in next generation sequencing technology (Hudson 2008), extensive analyses of single nucleotide polymorphisms (SNPs) will become increasingly feasible and affordable. This is opening up the potential for genome-wide association studies (GWAS-Donnelly 2008), making it possible to compare differences between individuals and cohorts at the nucleotide level, the lowest possible level of genetic resolution. Reviews by Allendorf et al. (2010), Avise (2010) and Ouborg et al. (2010) all stress the ongoing shift towards the use of genomic data in conservation applications. Making effective use of the rapidly expanding sources of data will necessitate the development of not only new methods for searching and filtering genomic data for intra- and inter-population signals, but also the development of appropriate statistical tests to determine their significance. This will require moving beyond the use of simple genetic models into the extensive application of multivariate analytical techniques (Jombart et al. 2009).

Although population genetic data have the ability to reveal connectivity patterns over long time periods, they also present challenges, since a large amount of variability is introduced into the data as a result of stochastic population processes (e.g. birth, mortality, and mutation) and natural plasticity in biological parameters (e.g. life-history characteristics). The influence of contemporary landscape or seascape patterns can also be confounded by historical influences, such as demographic bottlenecks, geographic barriers or patterns of anthropogenic habitat loss (Kool et al. 2011). Furthermore, there is also a mismatch between the time scale of genetic processes and the time scales of management interest, and reconciling them will require identifying the characteristic scales of the system, as well as innovative ways of adapting our understanding/ knowledge across different scales. However, with the development of multiple genetic marker types, new opportunities will emerge for empirically examining genetic connectivity patterns over various time scales, particularly as our ability to process, analyse and compare very large data sets improves with increased computing power.

Modelling and analysing connectivity

Sampling large spatial and temporal extents with a high degree of resolution is often impossible, and consequently researchers are forced to turn to models in order to investigate these types of environments. There are many challenges associated with modelling and analysing connectivity however (Panel 2), and many different approaches have been used in both metapopulation and landscape ecology (Tables 2, 3).

Statistics and measures

The earliest and simplest means of assessing connectivity involved using buffer distances or through the use of statistics summarizing the size and arrangement of landscape patches classified in a binary manner (habitat vs. non-habitat—Dale et al. 2002). Dispersal kernels can be used as a means of scaling the effect of distance on connectivity (Moilanen and Nieminen 2002), however this approach typically assumes that the dispersal process is radially symmetric and not influenced by intervening habitat structure, which may not be true (Mitarai et al. 2008). Population geneticists have also made extensive use of isolation by distance plots, comparing physical distance (typically geographic, but see White et al. 2010) versus some measure of genetic distance (e.g. Pinsky et al. 2010). Saura and Pascual-Hortal (Pascual-Hortal and Saura 2006; Saura and Pascual-Hortal 2007) have developed indices that characterize the reachability of habitat patches. Reachability considers habitat patches themselves as spaces where connectivity occurs, taking into account resources existing within patches (intra-patch connectivity), together with those available through connections with other habitat patches (inter-patch

Table 2 Characteristics of connectivity models	connectivity models					
		Technique	Domain	Path Type	Derivation	Selected references
Estimation of pair-wise connectivity values	$\frac{1}{G_I} = \frac{\sum_{j=1,j\neq i}^n w_{ij}y_j}{\sum_{j=1,j\neq i}^n y_j}$	Neighbourhood statistics	Any	Typically aggregated	Analytical	Dale et al. (2002) and Kindlmann and Burel (2008)
	2	Dispersal kernels	Continuous	Integrated	Analytical	Fujiwara et al. (2006) and Slone (2011)
	φ	Least cost path analysis	Semi-continuous	Single	Algorithmic	Algorithmic Douglas (1994)
	4	Circuit theory	Semi-continuous	Multiple, Integrated	Algorithmic	Algorithmic McRae et al. (2008)
	S	Spatially structured diffusion	Continuous (structured)	Integrated	Analytical	Ovaskainen (2004) and Ovaskainen et al. (2008a)
	ي م	Individual-based models	Any	Multiple	Algorithmic	Algorithmic Grimm and Railsback (2005)

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Analysis

	Technique	Domain	Path Type	Derivation	Derivation Selected references
s of connectivity 7	Graph theory	Discrete, semi- continuous	Multiple	Algorithmic	Algorithmic Minor and Urban (2008) and Urban et al. (2009)
~	Matrix theory	Discrete, semi- continuous	Single*	Analytical	Analytical Caswell (2001)
6	Differential equations	Continuous	Integrated	Analytical	Holmes et al. (1994) and Cantrell and Cosner (2003)

Although matrix elements are single, it is possible for them to contain nested functions

This table summarizes and compares different connectivity modelling approaches. The methods above the blue line provide ways of estimating connectivity values between objects, whereas those beneath the blue line provide a means of analysing aspects of local to system-wide network structure and behaviour Domain Discrete environments are patches or populations with a defined boundary and homogeneous interior, and are typically represented using polygons. Semi-continuous domains are typically cell or pixel-based discretizations of landscape features (e.g. raster landscapes). Continuous domains are infinitely divisible and are typically defined using equations or a mesh framework (e.g. finite element method or triangulated irregular network) Path Type Single path methods only allow a single value to describe the connection strength between populations. Multiple path methods allow multiple channels to influence connectivity between populations. Integrated paths provide analysis integrated across a distribution of paths Derivation Analytical methods have an algebraic foundation and are solved using equations. Algorithmic methods are required when analytical solutions are complicated or impossible, and are executed out by following a block of sequential instructions, frequently in a repetitive or recursive manner. Note that many analytical equations can be characterized and solved by algorithms, and some algorithms (particularly homogeneous recursions) can be formulated as analytical equations In order to move from individual (local approaches) to summarized or integrated representations, aggregations or convolutions over some form of distance are required, which equires specifying a scale for the analysis

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Modelling/analytical	approaches	Linkage explanation
From (Table 2 row)	To (Table 2 row)	-
Neighbourhood statistics (1)	Graph theory (7)	Join count statistics can be linked to graph theory through the construction of spanning trees
	Matrix theory (8)	Statistics can be obtained through aggregating and summarizing the matrix data
	Differential equations (9)	Statistics can be obtained through integrating, differentiating, combining, or solving sets of equations
Dispersal kernels (2)	Spatially structured diffusion (5)	Spatially structured diffusion adds spatial structure to the dispersal process/ Dispersal kernels can be constructed from spatially explicit diffusion models by averaging across all points in all directions over different lag distances
	Individual-based models (6)	Dispersal kernels can be used to define movement rules for individual-based models/Dispersal kernels can be estimated from individual-based models by aggregating positions of individuals into a frequency histogram, binning on distance travelled relative to a release point
	Graph theory (7)	Dispersal kernels can be used to link weights for graph based analysis/Graph connections can be aggregated at various distance intervals to generate dispersal kernels
	Matrix theory (8)	Values for connectivity matrix elements can be obtained from dispersal kernels by using the kernel as a lookup function in conjunction with information about distances between populations
	Differential equations (9)	If parametric, dispersal kernels can be used to define differential equations/Sets of differential equations can directly define or be combined to generate a dispersal kernel
Least cost path analysis (3)	Circuit theory (4)	Circuit theory can be viewed as an extension to least cost path analysis, allowing for multiple pathways, and addressing strength of corridor use across the landscape
	Graph theory (7), Matrix theory (8)	Connectivity values between pairs of patches or populations derived from LCP analyses can be used as individual cell entries in a connectivity matrix or individual links in a graph
	Differential equations (9)	Shortest paths across surfaces defined by differential equations can be solved numerically or using Euler-Lagrange equations
Circuit Theory (4)	Individual-based models (6)	Using the values from a circuit theory layer, individual pathways can be reconstructed using a cellular automaton-type approach. The resistance values from circuit theory also correspond to the movement rules that an individual follows when traversing a given habitat type/Multiple paths derived from individual based models can be summarized using focal or block statistics to yield surfaces resembling circuit pathways
Spatially structured diffusion (5)	Individual-based models (6)	Individual paths can be generated from a diffusion process using differential equation solvers in conjunction with a random walk model/Individual-based models could also be used to generate spatial diffusion-type surfaces using focal or block statistics
	Differential equations (9)	Spatially structured diffusion directly uses partial differential equations
Individual-based models (6)	Graph theory (7), Matrix theory (8)	Matrix models arise from individual-based stochastic models where each individual moves through its life cycle independently. Probabilities or numbers of successful individuals making the transition between patches can be used as matrix cell entries or graph edge values
Graph Theory (7)	Matrix theory (8)	Graphs can be built directly using connection information stored in adjacency or distance matrices and vice versa
	Differential equations (9)	Differential equations and graph theory (and consequently matrix theory) can be linked by considering differential equations as the continuous (or limit) form of difference equations for discrete systems. Difference equations can be used to describe relationships between semi-continuous locations in space and time

Table 3 Linkages between approaches for modelling and analysing connectivity

Panel 2 - Challenges and considerations for connectivity studies

General challenges and considerations

- Different conceptual and operational definitions for functional connectivity are required.
- Species are likely to have alternative dispersal modes.
- Monitoring and modeling long-distance dispersal can be challenging.
- The effects of spatial autocorrelation on parameterization and interpretation need to be considered.
- Species interdependencies and community effects should be considered, particularly in the context of conservation planning.
- Connectivity calculations for high-dimensional problems can require significant computing resources.
- Determining connectivity for non-stationary distributions (e.g. due to climate change) may need to be considered.

Data requirements and challenges

- Minimally, information regarding location and size of habitat patches are necessary for deriving distance-based measures, such as nearest neighbour distances, buffer measures and many graph-based metrics.
- Identifying the scale(s) at which species use the landscape (dispersal capability) makes it possible to use species-specific buffer or kernel-type connectivity measures.
- Neighbourhood characteristics provide a better understanding of local landscape characteristics, and improve estimates of
 emigration and immigration rates. Unfavourable habitat surrounding a patch will tend to decrease dispersal to and from the
 patch, reducing its functional connectivity.
- Detailed information about landscape structure is needed for detailed analysis of movement including development of pathtype measures. It is assumed that different habitats have different species-specific effects on movement.
- Data regarding species behaviour, including perceptual ability and behaviour at habitat boundaries improves analyses of
 movement paths and probabilities. Direct observations, satellite tracking data or mark-recapture data are usually needed to
 parameterize these models.
- Demographic information such as population size, reproduction rates, and mortality are necessary for parameterising full
 population models that have connectivity as a component. These data can be difficult to obtain, and are typically speciesand environment-specific.
- Estimates of environmental change are needed to assess the effects of processes such as succession, climate change or anthropogenic changes on connectivity, particularly over long spatial distances or time intervals.

Habitat-specific challenges and considerations

Terrestrial environments

- Effectively 2-dimensional landscapes.
- Radially symmetric responses are frequently assumed.
- Landscape barriers, such as roads, rivers or urban areas may be important for ground-dwelling species.
- Landscape traversability (e.g., differential costs of moving through forest vs. agriculture) is an important consideration.

Riverine environments

- Typically represented using 1 or 2 dimensions.
- Strongly asymmetric flow between upstream and downstream locations is common.
- Strong physical barriers (e.g. waterfalls, dams) often exist.
- Large temporal fluctuations are common.

Marine environments

- Truly 3-dimensional.
- Fluid dynamics (e.g. advection and diffusion by winds and ocean currents) are important processes, and are computationally
 intensive to model.
- Complete dispersal barriers are less common than in terrestrial or riverine systems, and are often cryptic.
- Strong asymmetries in connectivity patterns are common.
- Species-specific transport mechanisms can be challenging to identify, and demographic rates (e.g. mortality) can be difficult to obtain.

connectivity), and consequently connectivity can be generated by large individual high-quality patches, from connections between patches, or a combination of both. Saura and Rubio (2010) also demonstrated how the probability of connection metric (PC) could be expressed in terms of the relative contribution of an individual component towards overall habitat availability in the landscape, and how that score could be partitioned into three components—intra-patch connectivity, dispersal flux through the patch, and the contribution of a component to the connectivity between other habitat patches (i.e. as a steppingstone). Additional landscape metrics have been reviewed by Kindlmann and Burel (2008).

Pathfinding

Least-cost path (LCP) analysis also provides a means of scaling distance values between patches, and continues to be influential in landscape ecology (Urban et al. 2009). With LCP analysis, connectivity values are based on the path of least resistance between any two landscape elements. Exact and approximate algorithms exist for the computation of LCPs, but computation time remains a challenge for high-dimensional landscapes (Urban et al. 2009). In addition, LCP computation requires species-specific resistance values for different habitat types, which can be difficult to parameterize. Electric circuit theory has also been used in an ecological context to investigate path-type connectivity (McRae et al. 2008), and can be considered as an extension to LCP analysis. Like LCP analysis, circuit theory operates on the basis of deriving resistance values between patches, but rather than identifying a single path, this framework allows for multiple paths between patches. This is a conceptually important development, since it becomes possible to investigate swaths as connections, as well as multiple corridor routing options (Ferreras 2001).

Spatially structured diffusion

Spatially structured diffusion provides another way of analysing animal movements in heterogeneous landscapes using mark-recapture and tracking data (Ovaskainen 2004; Ovaskainen et al. 2008a). It operates by incorporating directional biases towards particular habitats at patch boundaries using a diffusion framework. Rather than considering discrete corridors, spatially structured diffusion integrates in a continuous manner across all possible movement pathways, and allows for rigorous estimates of species observability, as well as movement rates and mortalities in different habitat types, and transition rates between different pairs of habitats (Ovaskainen et al. 2008b). Occupancy times in landscape elements, hitting probabilities of landscape elements, quasi-stationary occupancy distributions, time evolution of occupancy distribution as function of initial condition, and occupancy probability densities between two observation points can also be derived directly from the diffusion process (Ovaskainen 2008).

Individual-based simulation

For complex environments with extremely high levels of spatial and temporal variability, individual-based models (IBMs) (Grimm and Railsback 2005) are being used to generate increasingly realistic simulations based on real-world data (Paris et al. 2007; Kool et al. 2010). IBMs operate on the basis of programmatically assigning properties and behaviour to individuals and then allowing them to interact within a stochastic simulation environment (Levey et al. 2008; Kool et al. 2011). Although individual-based models are flexible in terms of their structure and dynamics, they require programming expertise, are difficult to parameterize rigorously, and cannot be manipulated, analysed and reconfigured in the manner of algebraic equations.

Graph theory

Graph theory has been extensively used to study the structure and properties of connectivity networks, as well as providing a means of displaying and visualizing them (Urban et al. 2009; Galpern et al. 2011; Luque et al. 2012). Graph theory provides a means of efficiently analysing large and complex networks, as well as their emergent properties and key structural characteristics. For example, measures of centrality (e.g., betweenness, degree, closeness) identify the position or role of a node with respect to its neighbours or the entire network (Estrada and Bodin 2008; Opsahl et al. 2010), and detecting nodes that exert a high degree of influence over the dynamics of the entire system. Network community structure can be evaluated through various clustering methods (Clauset et al. 2004; Palla et al. 2005), characterizing associations

between individuals or groups. The degree distribution of a network is the probability distribution of the number of edges a node will have across the entire network, providing an indication of resilience and communicability within the network (Minor and Urban 2008). A network with a skewed degree distribution and several large hubs would suggest resilience to random node failure, and fast spread across the network (Proulx et al. 2005). Metanetworks (networks that model the relationships between other networks) have also been proposed as a means of linking species networks with spatial networks (Luque et al. 2012; Rubio and Saura 2012). For in-depth reviews of graph and network metrics, refer to Rayfield et al. (2011). Although it is important to be mindful of some of the potential limitations of relying on a graph-theoretic approach (Moilanen 2011), recent applications in both terrestrial and marine systems (Treml et al. 2008; Minor et al. 2009; Erös et al. 2012), as well as rapidly expanding interest in the analysis of social networks (Bodin and Crona 2009; Borgatti et al. 2009) suggests that graph theory will remain an active part of connectivity research for some time.

Matrix analysis

Matrix models provide another means of analysing connectivity flows (Caswell 2001), and have recently been used to project connectivity structure over time (Kool 2009), providing a link between individualbased biophysical dispersal models and population genetic structure (Foster et al. 2012). The sensitivity and elasticity of connectivity matrices (Caswell 2001, 2007) can be used to identify connections that exert the greatest influence on the overall system, and ordering matrices through sorting (Tsafrir et al. 2005), reduction (Bode et al. 2006) or recursive partitioning (Jacobi et al. 2012) makes it possible to evaluate natural clusters of exchange. More advanced techniques, such as singular value decomposition, and matrix perturbation theory for analysing connectivity and designing optimal networks have also been explored (Aiken and Navarrete 2011; Jacobi and Jonsson 2011).

Linkages between approaches

Many similarities exist between the various connectivity measures and analyses used in landscape ecology, metapopulation ecology, and connectivity research (Table 3). Cluster analysis based on nearestneighbour distances is closely connected to the construction of minimum spanning tree type graphs. Critical distances used in graph theory are structurally the same as buffer or neighbourhood measures in metapopulation studies and statistical habitat modelling (Visconti and Elkin 2009). Pair-wise distance matrices used inside connectivity measures can be constructed based on declining-by-distance dispersal kernels or via least cost path computations (Urban et al. 2009). Graph theoretic approaches and matrices can be explicitly linked via the construction of an adjacency matrix representing the strength of connections between nodes. Caswell (2001) noted that matrix models can be linked to IBMs, and arise naturally from stochastic models where each individual moves through its life cycle independently. The various methods are in many cases closely related ways of approaching the same problem-characterizing relationships among patches, populations or demes. Rather than focusing on a particular modelling framework, it is more profitable to classify spatial studies and connectivity measures according to their structural characteristics (Panel 1).

Challenges and opportunities

Modelling and analysing connectivity presents a number of challenges. Landscape dynamics (e.g. successional changes, fragmentation) have the potential to confuse connectivity observations, leading to underestimates or even an apparent lack of connectivity effects (Hodgson et al. 2009a), and imperfect detection of species in sites has long been recognized as a problem for metapopulation studies, leading to biases in parameter estimation, including overestimation of population turnover, extinction and colonization rates, dispersal distances and connectivity as a whole (Mackenzie et al. 2003). Overestimation of connectivity can then lead to underestimation of conservation needs. One way of addressing this is through the use of stochastic state-space models (Patterson et al. 2008b). Under this framework, a process model is coupled with a separate observation model, providing a means of partitioning the sources of variability that are truly associated with the process from those associated with observation. Spatial autocorrelation is also an important consideration for connectivity studies (González-Megías et al. 2005). Autocorrelation in observations can occur due to correlation in local habitat quality, spatially correlated dynamics or synchronizing factors such as weather (Van Teeffelen and Ovaskainen 2007). If spatial autocorrelation is ignored, then events are taken as independent when in truth they are not, leading to incorrect parameter estimates and false estimates of statistical significance. Autocorrelation is particularly a problem when habitat data are represented using high-resolution grids of semi-continuously varying habitat quality (Drielsma and Ferrier 2009), and consequently individual spatial units (i.e. individual raster cells) cannot be taken as dynamically independent from their neighbourhood. This has operational significance because most spatial habitat data currently exists in raster format, and high-resolution analyses are necessary to link the data with on-theground conservation applications (Elith and Leathwick 2009).

A number of opportunities exist for moving connectivity research forward by taking advantage of advances made in other fields. Some of the challenges facing connectivity researchers correspond to problems in other disciplines, and existing solutions can be brought to bear in a biological context. For example, solvers for the knapsack problem from computer science have been applied to optimizing environmental designs (Higgins et al. 2008), and the entire framework of graph-theoretic connectivity is an import from mathematical/computational sciences (Urban and Keitt 2001). Allesina and Pascual (2009) demonstrated how an adaptation of the Google PageRank algorithm could be used to identify key species whose loss could result in cascading extinctions, and the same could be used to identify groups of co-dependent patches or demes. Stochastic control theory (Wang et al. 2008) could be used to develop management strategies that dynamically respond to changes in connectivity, and bandwidth-allocation models (Ogryczak et al. 2008) could be modified to determine how resources could be most effectively distributed to maintain existing connectivity structure. Many of these questions relating to connectivity research appear to fall under the domain of complex adaptive systems, and complexity in general (Miller and Page 2007). However, it is also important to recognize that populations are not binary switches, and ecological systems frequently exhibit non-linear and strategic behaviour. By design, many algorithms and analytical methods focus on maximal or minimal aspects of the system, but in many cases, the variability and distribution of responses are just as important, sometimes even more. Developing ways of assessing and testing how models, metrics and analyses results are affected by different forms of variability, as well as behaviour that evolves over time will be essential for moving forward with population connectivity research.

Management applications

Connectivity is a critical consideration in biodiversity conservation and management. Interactions between humans and landscapes occur through spatially defined interactions, which influence connectivity (Crooks and Sanjayan 2006). Spatial considerations were originally incorporated into conservation through the use of critical maximum dispersal distances and minimum patch size requirements, and spatial aggregation was achieved using boundary length penalties (Sarkar et al. 2006). Boundary length penalties penalize high edge-to-area ratios when carrying out optimization of reserve networks, leading to more globular delineations for individual sites and to more aggregated network solutions. This technique is still widely used in reserve network design, since structural aggregation is beneficial from both an ecological and economic perspective with respect to reserve establishment and management (Ball et al. 2009).

Presently, many different connectivity indices, both structural and functional, can be calculated using publicly available software packages such as FRAGSTATS (McGarigal et al. 2002), PATHMA-TRIX (Ray 2005), Conefor (Saura and Torné 2009), Marine Geospatial Ecology Tools (MGET-Roberts et al. 2010), or generic GIS software. These connectivity measures can be used as explanatory variables in further statistical analysis and modelling of conservation decisions. Conservation-oriented single-species spatial analysis can be carried out using empirically fitted metapopulation models (Drielsma and Ferrier 2009) or spatial population viability analyses (Naujokaitis-Lewis et al. 2009), although data demands of detailed dynamic models are generally high. Detailed mechanistic analyses of dispersal are also possible, for example, via spatially structured diffusion. Additionally, specialized software exists for advanced path-based analysis (McRae et al. 2008), and corridor building (Cushman et al. 2009).

Several software packages are publicly available for addressing connectivity in multi-species systematic conservation planning. The ResNet software package incorporates connectivity considerations into reserve network design via path-like graph-theoretic considerations (Ciarleglio et al. 2009). MARXAN and MARXAN with Zones implement patch size requirements and the boundary length penalty technique (Ball et al. 2009; Watts et al. 2009). The grid-based Zonation software implements species-specific parametric neighbourhood responses in a non-directional (terrestrial) environment (Moilanen and Wintle 2007) and for freshwater networks with strongly directed connectivity (Leathwick et al. 2010). It also implements pair-wise and many-to-one connectivity responses between species, between environments, between existing and proposed conservation areas (Lehtomäki et al. 2009), or between the present and the future in the climate change context (Carroll et al. 2010).

Despite significant progress during the past decade, many challenges remain in understanding how to best include connectivity in conservation management, when the needs of multiple species and environments, habitat quality and connectivity, direct costs and opportunity costs, short-term and long-term objectives, and multiple alternative conservation actions must be balanced (Pressey et al. 2007). Johst et al. (2011) were able to develop an analytical method for examining trade-offs between different landscape attributes, but integrating multiple forms of connectivity into the same analysis, using sparse data to effectively parameterize conservation analyses, and understanding the most robust and appropriate use of connectivity criteria in spatial conservation planning are all areas requiring further study. There are also questions regarding appropriate role of connectivity in conservation relative to strategies that primarily target habitat area or habitat quality-the two most fundamental determinants of regional carrying capacity for any species (Hodgson et al. 2009b). Using simulations, Visconti and Elkin (2009) were able to quantitatively show that connectivity metrics that take into account patch quality performed significantly better with regards to correctly ranking patches according to their contribution to overall metapopulation viability. Reinforcing connectivity for one species may add breeding habitat for another, implying potentially great benefits from strategies such as agri-environment schemes (Donald and Evans 2006). However, working with connectivity alone does not provide information regarding what is necessary or adequate for conservation. While connectivity can inform decision-makers about patterns of dispersal and colonization, alone it does not provide comprehensive information on local population dynamics, age/stage structure, or population growth and extinction.

As another general concern, connectivity is an uncertain management criterion despite it being the one most commonly proposed as a solution for conservation under climate change (Heller and Zavaleta 2009). There are numerous conceptual and operational definitions for connectivity, making discussion about connectivity prone to linguistic uncertainty in communication. Choices of connectivity metrics are also prone to human decision uncertainty about what form of connectivity measure is applied and for what species (or other biodiversity features). Further complicating use of connectivity as a management criterion is epistemic uncertainty (lack of knowledge) about the correct structure and parameterization of connectivity. Consequently, application of connectivity in multi-species conservation management needs to be implemented with care and in a manner robust to uncertainty. While our ability compute connectivity metrics improves, our understanding about the appropriate use of connectivity in conservation management does not improve at the same rate, and our linguistic and decision uncertainty have not been reduced.

Synthesis

A significant part of the value of connectivity research lies in assembling the individual pieces of a landscape or seascape together into in integrated spatially and temporally explicit whole. Studying populations an integrated manner makes it possible to test the consistency of our understanding of the system, and reveals if critical components are not being accounted for. Moreover, by examining the various components in concert, other aspects emerge. The first is the critical importance of scale. Depending on the spatial and temporal scales at which one observes a landscape or seascape, patches may change, blend into one another, or cease to effectively exist altogether. Understanding the scales at which different landscape processes operate, as well as ways of identifying those scales is essential for devising efficient monitoring strategies, as well as determining functional connections. This is no trivial task, since scales will vary not only among the different processes, but also according to how individual species perceive and use their environment. Consequently, in addition to seeking out unifying principles, it is equally important to critically evaluate the causes and consequences of variability between individuals, species and assemblages, and to address the interactions between them. Fusing the homogeneity of an integrated design with variability down to the genetic level requires reconciling topdown, holistic approaches with bottom-up, reductionist approaches. The complexity of dispersal and connectivity, augmented by the need to account for additional factors such as the role of demographic processes or to integrate with social and economic systems might seem cause for despair, but this is a challenge for which connectivity researchers are wellsuited. Teasing signal from noise, partitioning intraand inter-group variation, and developing conceptual models that explain system behaviour using the minimum amount of detail required are common practice in connectivity research. Fortunately, is it not necessary to develop methods entirely de novo. Although models designed for mechanical systems may be too simple for biological systems, they can at least serve as a basis for further development. Communication across disciplines will also be crucial when developing conservation and management strategies. Connectivity scientists need to be transparent about what their measurements and models mean, and the assumptions behind them. It is also essential to distil and simplify this knowledge into an accessible form-through the development of tools, and outreach beyond scientific publications. From managers, a clear articulation of their needs is required, as well as a transparent assessment of constraints: logistical, social and economic. Naturally, this will be an iterative and interactive process, but facilitating these connections, as well as identifying where productive new linkages could be formed will be important moving forward. Lastly, understanding the implications of long-term connectivity will involve strengthening links with population genetic theory, including aspects of speciation and biogeography. Over time, inter-population processes will be dependent on intra-population ones, such as reproductive success, carrying capacity and habitat quality. Fundamentally, an improved understanding of connectivity is needed to fully appreciate the likely development of biodiversity patterns under climate change and other human pressures.

Summary

From advances in physical tracking, to the application of new genetic techniques, as well as ongoing developments in modelling and analysis, it is clear that much work has been done, and is still going on to improve our understanding of connectivity. Methods for measuring connectivity have greatly improved in both extent and resolution, spatially and temporally. A wide range of options exist for monitoring organisms at a variety of scales in terrestrial and aquatic environments. Similarly, many techniques are available to characterizing connectivity and to represent its underlying processes. From relatively simple measures, such as summary statistics through to dynamic individual-based models and spatially structured diffusion models, researchers have many choices depending on data availability and structure, as well as how the results will ultimately be used. It is also important to give strong consideration to how connectivity data and models can be integrated into conservation and management strategies. To this end, a number of software tools have been developed, but ensuring that this information is effectively used will require careful consideration of what operational definitions of connectivity are most relevant to the problem at hand, as well as its relative importance in the decision-making process. Also, while our understanding of connectivity is improving, there will still be a strong need to gather field data on individual and species-level behaviour, habitat quality, and demography. Nevertheless, the progress to date in detecting and recognizing connectivity patterns, and understanding the processes responsible for generating them is highly encouraging, and we look forward to seeing the benefits that an improved understanding of connectivity will provide in the future.

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References

- Ahmad MA, Teredesai A (2006) Modeling spread of ideas in online social networks. In: Proceedings of the fifth Australasian conference on data mining and analystics, vol 61. Australian Computer Society, Inc., Sydney, pp 185–190
- Aiken CM, Navarrete SA (2011) Environmental fluctuations and asymmetrical dispersal: generalized stability theory for studying metapopulation persistence and marine protected areas. Mar Ecol Prog Ser 428:77–88
- Allendorf FW, Hohenlohe PA, Luikart G (2010) Genomics and the future of conservation genetics. Nat Rev Genet 11(10):697–709
- Allesina S, Pascual M (2009) Googling Food Webs: can an Eigenvector Measure Species' Importance for Coextinctions? PLoS Comput Biol 5(9):e1000494
- Almany GR, Berumen ML, Thorrold SR, Planes S, Jones GP (2007) Local Replenishment of Coral Reef Fish Populations in a Marine Reserve. Science 316(5825):742–744
- Avise JC (2010) Perspective: conservation genetics enters the genomics era. Conserv Genet 11(2):665–669
- Baguette M, Van Dyck H (2007) Landscape connectivity and animal behavior: functional grain as a key determinant for dispersal. Landscape Ecol 22(8):1117–1129
- Balkenhol N, Gugerli F, Cushman S, Waits L, Coulon A, Arntzen J, Holderegger R, Wagner H (2009) Identifying future research needs in landscape genetics: where to from here? Landscape Ecol 24(4):455–463
- Ball IR, Possingham HP, Watts M (2009) Marxan and relatives: software for spatial conservation prioritisation. In: Moilanen A, Wilson KA, Possingham HP (eds) Spatial conservation prioritisation: quantitative methods and computational tools. Oxford University Press, Oxford, pp 185–195
- Beerli P, Palczewski M (2010) Unified framework to evaluate panmixia and migration direction among multiple sampling locations. Genetics 185(1):313–326
- Berry O, Tocher MD, Sarre SD (2004) Can assignment tests measure dispersal? Mol Ecol 13(3):551–561
- Berumen M, Walsh H, Raventos N, Planes S, Jones G, Starczak V, Thorrold S (2010) Otolith geochemistry does not reflect dispersal history of clownfish larvae. Coral Reefs 29(4): 883–891

- Bode M, Bode L, Armsworth PR (2006) Larval dispersal reveals regional sources and sinks in the Great Barrier Reef. Mar Ecol Prog Ser 308:17–25
- Bodin Ö, Crona BI (2009) The role of social networks in natural resource governance: what relational patterns make a difference? Glob Environ Change 19(3):366–374
- Borgatti SP, Mehra A, Brass DJ, Labianca G (2009) Network analysis in the social sciences. Science 323(5916):892–895
- Borgman C, Wallis J, Enyedy N (2007) Little science confronts the data deluge: habitat ecology, embedded sensor networks, and digital libraries. Int J Digit Libr 7(1):17–30
- Broquet T, Petit EJ (2009) Molecular estimation of dispersal for ecology and population genetics. Ann Rev Ecol Syst 40(1):193–216
- Cantrell RS, Cosner C (2003) Spatial ecology via reaction-diffusion equations. Wiley, Chichester
- Carroll C, Dunk JR, Moilanen A (2010) Optimizing resiliency of reserve networks to climate change: multispecies conservation planning in the Pacific Northwest, USA. Glob Change Biol 16(3):891–904
- Caswell H (2001) Matrix population models: construction, analysis, and interpretation. Sinauer Associates, Sunderland
- Caswell H (2007) Sensitivity analysis of transient population dynamics. Ecol Lett 10(1):1–15
- Cerdeira JO, Pinto LS, Cabeza M, Gaston KJ (2010) Species specific connectivity in reserve-network design using graphs. Biol Conserv 143(2):408–415
- Chester CC (2006) Conservation across borders: biodiversity in an interdependent world. Island Press, Washington, DC
- Ciarleglio M, Wesley Barnes J, Sarkar S (2009) ConsNet: new software for the selection of conservation area networks with spatial and multi-criteria analyses. Ecography 32(2): 205–209
- Claudet J (2011) Marine protected areas: a multidisciplinary approach. Cambridge University Press, Cambridge
- Clauset A, Newman MEJ, Moore C (2004) Finding community structure in very large networks. Phys Rev E 70(6):066111
- Cowen RK, Guigand CM (2008) In situ ichthyoplankton imaging system (ISIIS): system design and preliminary results. Limnol Oceanogr Methods 6:126–132
- Cowen RK, Sponaugle S (2009) Larval dispersal and marine population connectivity. Annu Rev Mar Sci 1:443–466
- Crooks KR, Sanjayan M (eds) (2006) Connectivity conservation. Cambridge University Press, Cambridge
- Cushman SA, McKelvey KS, Schwartz MK (2009) Use of empirically derived source-destination models to map regional conservation corridors (Utilización de Modelos Fuente-Destino Empíricamente Derivados para Cartografiar Corredores de Conservación Regionales). Conserv Biol 23(2):368–376
- Dale MRT, Dixon P, Fortin M-J, Legendre P, Myers DE, Rosenberg MS (2002) Conceptual and mathematical relationships among methods for spatial analysis. Ecography 25(5):558–577
- Donald PF, Evans AD (2006) Habitat connectivity and matrix restoration: the wider implications of agri-environment schemes. J Appl Ecol 43(2):209–218
- Donnelly P (2008) Progress and challenges in genome-wide association studies in humans. Nature 456(7223):728–731
- Douglas DH (1994) Least-cost path in GIS using an accumulated cost surface and slopelines, Cartographica. Int J Geogr Inf Geovisual 31(3):37–51

- Drielsma M, Ferrier S (2009) Rapid evaluation of metapopulation persistence in highly variegated landscapes. Biol Conserv 142(3):529–540
- Duchesne P, Castric T, Bernatchez L (2005) pasos (parental allocation of singles in open systems): a computer program for individual parental allocation with missing parents. Mol Ecol Notes 5(3):701–704
- Durbec M, Cavalli L, Grey J, Chappaz R, Nguyen B (2010) The use of stable isotopes to trace small-scale movements by small fish species. Hydrobiologia 641(1):23–31
- Elith J, Leathwick JR (2009) Species distribution models: ecological explanation and prediction across space and time. Annu Rev Ecol Evol Syst 40(1):677–697
- Erös T, Olden J, Schick R, Schmera D, Fortin M-J (2012) Characterizing connectivity relationships in freshwaters using patch-based graphs. Landscape Ecol 27(2):303–317
- Estrada E, Bodin Ö (2008) Using network centrality measures to manage landscape connectivity. Ecol Appl 18(7): 1810–1825
- Excoffier L, Heckel G (2006) Computer programs for population genetics data analysis: a survival guide. Nat Rev Genet 7(10):745–758
- Faaborg J, Holmes RT, Anders AD, Bildstein KL, Dugger KM, Gauthreaux SA Jr, Heglund P, Hobson KA, Jahn AE, Johnson DH, Latta SC, Levey DJ, Marra PP, Merkord CL, Erica NOL, Rothstein SI, Sherry TW, Scott Sillett T, Thompson III FR, Warnock N (2010) Recent advances in understanding migration systems of New World land birds. Ecol Monogr 80(1):3–48
- Faubet P, Gaggiotti OE (2008) A new Bayesian method to identify the environmental factors that influence recent migration. Genetics 178(3):1491–1504
- Ferreras P (2001) Landscape structure and asymmetrical interpatch connectivity in a metapopulation of the endangered Iberian lynx. Biol Conserv 100(1):125–136
- Figueira W, Crowder L (2006) Defining patch contribution in source-sink metapopulations: the importance of including dispersal and its relevance to marine systems. Popul Ecol 48(3):215–224
- Fleming PJS, Tracey JP (eds) (2008) Aerial surveys of wildlife: theory and applications. CSIRO Publishing, Collingwood
- Foster NL, Paris CB, Kool JT, Baums IB, Stevens JR, Sanchez JA, Bastidas C, Agudelo C, Bush P, Day O, Ferrari R, Gonzalez P, Gore S, Guppy R, McCartney M, McCoy C, Mendes J, Srinivasan A, Steiner S, Vermeij MJA, Weil E, Mumby PJ (2012) Connectivity of Caribbean coral populations: complementary insights from empirical and modelled gene flow. Mol Ecol 21(5):1143–1157
- Francesco Ficetola G, Bonin A (2011) Conserving adaptive genetic diversity in dynamic landscapes. Mol Ecol 20(8):1569–1571
- Fujiwara M, Anderson K, Neubert M, Caswell H (2006) On the estimation of dispersal kernels from individual markrecapture data. Environ Ecol Stat 13(2):183–197
- Galpern P, Manseau M, Fall A (2011) Patch-based graphs of landscape connectivity: a guide to construction, analysis and application for conservation. Biol Conserv 144(1):44–55
- González-Megías A, Gómez JM, Sánchez-Piñero F (2005) Consequences of spatial autocorrelation for the analysis of metapopulation dynamics. Ecology 86(12):3264–3271

- Grimm V, Railsback SF (2005) Individual-based modeling and ecology. Princeton University Press, Princeton
- Hamilton MP, Graham EA, Rundel PW, Allen MF, Kaiser W, Hansen MH, Estrin DL (2007) New approaches in embedded networked sensing for terrestrial ecological observatories. Environ Eng Sci 24(2):192–204
- Harrison Hugo B, Williamson David H, Evans Richard D, Almany Glenn R, Thorrold Simon R, Russ Garry R, Feldheim Kevin A, van Herwerden L, Planes S, Srinivasan M, Berumen Michael L, Jones Geoffrey P (2012) Larval export from marine reserves and the recruitment benefit for fish and fisheries. Curr Biol 22(11):1023–1028
- Hedgecock D, Barber PH, Edmands S (2007) Genetic approaches to measuring connectivity. Oceanography 20(3): 70–79
- Heller NE, Zavaleta ES (2009) Biodiversity management in the face of climate change: a review of 22 years of recommendations. Biol Conserv 142(1):14–32
- Higgins AJ, Hajkowicz S, Bui E (2008) A multi-objective model for environmental investment decision making. Comput Oper Res 35(1):253–266
- Hilty J, Lidicker WZ Jr, Merenlender A (2006) Corridor ecology: the science and practice of linking landscapes for biodiversity conservation. Island Press, Washington, DC
- Hobson KA (2008) Applying isotopic methods to tracking animal movements. In: Keith AH, Leonard IW (eds) Terrestrial ecology. Elsevier, New York, pp 45–78
- Hodgson JA, Moilanen A, Thomas CD (2009a) Metapopulation responses to patch connectivity and quality are masked by successional habitat dynamics. Ecology 90(6):1608–1619
- Hodgson JA, Thomas CD, Wintle BA, Moilanen A (2009b) Climate change, connectivity and conservation decision making: back to basics. J Appl Ecol 46(5):964–969
- Holderegger R, Wagner H (2006) A brief guide to landscape genetics. Landscape Ecol 21(6):793–796
- Holmes EE, Lewis MA, Banks JE, Veit RR (1994) Partial differential equations in ecology: spatial interactions and population dynamics. Ecology 75(1):17–29
- Hudson ME (2008) Sequencing breakthroughs for genomic ecology and evolutionary biology. Mol Ecol Res 8(1):3–17
- Hulme PE (2009) Trade, transport and trouble: managing invasive species pathways in an era of globalization. J Appl Ecol 46(1):10–18
- Jacobi MN, Jonsson PR (2011) Optimal networks of nature reserves can be found through eigenvalue perturbation theory of the connectivity matrix. Ecol Appl 21(5): 1861–1870
- Jacobi MN, André C, Döös K, Jonsson PR (2012) Identification of subpopulations from connectivity matrices. Ecography doi:10.1111/j.1600-0587.2012.07281.x
- Jacobson B, Peres-Neto P (2010) Quantifying and disentangling dispersal in metacommunities: how close have we come? How far is there to go? Landscape Ecol 25(4):495–507
- Jaquiéry J, Broquet T, Hirzel AH, Yearsley J, Perrin N (2011) Inferring landscape effects on dispersal from genetic distances: how far can we go? Mol Ecol 20(4):692–705
- Johst K, Drechsler M, van Teeffelen AJA, Hartig F, Vos CC, Wissel S, Wätzold F, Opdam P (2011) Biodiversity conservation in dynamic landscapes: trade-offs between number, connectivity and turnover of habitat patches. J Appl Ecol 48(5):1227–1235

- Jombart T, Pontier D, Dufour AB (2009) Genetic markers in the playground of multivariate analysis. Heredity 102(4): 330–341
- Jones AG, Ardren WR (2003) Methods of parentage analysis in natural populations. Mol Ecol 12:2511–2523
- Jones GP, Planes S, Thorrold SR (2005) Coral reef fish larvae settle close to home. Curr Biol 15(14):1314–1318
- Jones AG, Small CM, Paczolt KA, Ratterman NL (2010) A practical guide to methods of parentage analysis. Mol Ecol Res 10(1):6–30
- Karanth KU, Nichols JD, Kumar NS, Hines JE (2006) Assessing tiger population dynamics using photographic capturerecapture sampling. Ecology 87(11):2925–2937
- Kays R, Kranstauber B, Jansen C, Carbone P, Rowcliffe M, Foundtain T Tilak S (2009) Camera traps as sensor networks for monitoring animal communities. In: Local computer networks, LCN 2009. IEEE 34th conference on, 2009, pp 811–818
- Kindlmann P, Burel F (2008) Connectivity measures: a review. Landscape Ecol 23(8):879–890
- Kool JT (2009) An object-oriented, individual-based approach for simulating the dynamics of genes in subdivided populations. Ecol Inform 4:136–146
- Kool JT, Paris CB, Andréfouët S, Cowen RK (2010) Complex migration and the development of genetic structure in subdivided populations: an example from Caribbean coral reef ecosystems. Ecography 33:597–606
- Kool JT, Paris CB, Barber PH, Cowen RK (2011) Connectivity and the development of population genetic structure in Indo-West Pacific coral reef communities. Glob Ecol Biogeogr 20(5):695–706
- Leathwick JR, Moilanen A, Ferrier S, Julian K (2010) Complementarity-based conservation prioritization using a community classification, and its application to riverine ecosystems. Biol Conserv 143(4):984–991
- Lehtomäki J, Tomppo E, Kuokkanen P, Hanski I, Moilanen A (2009) Applying spatial conservation prioritization software and high-resolution GIS data to a national-scale study in forest conservation. For Ecol Manage 258(11):2439–2449
- Levey DJ, Tewksbury JJ, Bolker BM (2008) Modelling longdistance seed dispersal in heterogeneous landscapes. J Ecol 96(4):599–608
- Liu J, Hull V, Morzillo AT, Wiens JA (2011) Sources. Cambridge University Press, Cambridge, MA, Sinks and Sustainability
- Lowe WH, Allendorf FW (2010) What can genetics tell us about population connectivity? Mol Ecol 19(15):3038–3051
- Luque S, Saura S, Fortin M-J (2012) Landscape connectivity analysis for conservation: insights from combining new methods with ecological and genetic data. Landscape Ecol 27:153–157
- Mackenzie DI, Nichols JD, Hines JE, Knutson MG, Franklin AB (2003) Estimating site occupancy, colonization, and local extinction when a species is detected imperfectly. Ecology 84(8):2200–2207
- Manel S, Schwartz MK, Luikart G, Taberlet P (2003) Landscape genetics: combining landscape ecology and population genetics. Trends Ecol Evol 18(4):189–197
- Marshall TC, Slate J, Kruuk LEB, Pemberton JM (1998) Statistical confidence for likelihood-based paternity inference in natural populations. Mol Ecol 7(5):639–655

- Martin Taylor S (2009) Transformative ocean science through the VENUS and NEPTUNE Canada ocean observing systems. Nucl Instrum Methods Phys Res, Sect A 602(1): 63–67
- McGarigal K, Cushman SA, Neel MC, Ene E (2002) FRAG-STATS: spatial pattern analysis program for categorical maps. University of Massachusetts, Amherst
- McRae BH, Dickson BG, Keitt TH, Shah VB (2008) Using circuit theory to model connectivity in ecology, evolution, and conservation. Ecology 89(10):2712–2724
- Miller JH, Page SE (2007) Complex adaptive systems: an introduction to computational models of social life. Princeton University Press, Princeton
- Millspaugh J, Marzluff JM (eds) (2001) Radio tracking and animal populations. Academic Press, Salt Lake City
- Minor ES, Urban DL (2008) A graph-theory framework for evaluating landscape connectivity and conservation planning. Conserv Biol 22(2):297–307
- Minor ES, Tessel SM, Engelhardt KAM, Lookingbill TR (2009) The role of landscape connectivity in assembling exotic plant communities: a network analysis. Ecology 90(7): 1802–1809
- Mitarai S, Siegel DA, Winters KB (2008) A numerical study of stochastic larval settlement in the California Current system. J Mar Syst 69(3–4):295–309
- Mobley KB (2011) Grandfathering in a new era of parentage analysis. Mol Ecol 20(6):1080–1082
- Moilanen A (2011) On the limitations of graph-theoretic connectivity in spatial ecology and conservation. J Appl Ecol 48:1543–1547
- Moilanen A, Nieminen M (2002) Simple connectivity measures in spatial ecology. Ecology 83(4):1131–1145
- Moilanen A, Wintle BA (2007) The boundary-quality penalty: a quantitative method for approximating species responses to fragmentation in reserve selection. Conserv Biol 21(2):355–364
- Munday P, Leis J, Lough J, Paris C, Kingsford M, Berumen M, Lambrechts J (2009) Climate change and coral reef connectivity. Coral Reefs 28(2):379–395
- Munro AR, Gillanders BM, Thurstan S, Crook DA, Sanger AC (2009) Transgenerational marking of freshwater fishes with enriched stable isotopes: a tool for fisheries management and research. J Fish Biol 75(3):668–684
- Musyl MK, Domeier ML, Nasby-Lucas N, Brill RW, McNaughton LM, Swimmer JY, Lutcavage MS, Wilson SG, Galuardi B, Liddle JB (2011) Performance of pop-up satellite archival tags. Mar Ecol Prog Ser 433:1–28
- Naujokaitis-Lewis IR, Curtis JMR, Arcese P, Rosenfeld J (2009) Sensitivity analyses of spatial population viability analysis models for species at risk and habitat conservation planning. Conserv Biol 23(1):225–229
- Newsome SD, Martinez del Rio C, Bearhop S, Phillips DL (2007) A niche for isotopic ecology. Front Ecol Environ 5(8):429–436
- Ogryczak W, Wierzbicki A, Milewski M (2008) A multi-criteria approach to fair and efficient bandwidth allocation. Omega 36(3):451–463
- Opsahl T, Agneessens F, Skvoretz J (2010) Node centrality in weighted networks: generalizing degree and shortest paths. Soc Netw 32(3):245–251

- Ouborg NJ, Pertoldi C, Loeschcke V, Bijlsma RK, Hedrick PW (2010) Conservation genetics in transition to conservation genomics. Trends Genet 26(4):177–187
- Ovaskainen O (2004) Habitat-specific movement parameters estimated using mark-recapture data and a diffusion model. Ecology 85(1):242–257
- Ovaskainen O (2008) Analytical and numerical tools for diffusion-based movement models. Theor Popul Biol 73(2):198–211
- Ovaskainen O, Luoto M, Ikonen I, Rekola H, Meyke E, Kuussaari M (2008a) An empirical test of a diffusion model: predicting clouded apollo movements in a novel environment. Am Nat 171(5):610–619
- Ovaskainen O, Rekola H, Meyke E, Arjas E (2008b) Bayesian methods for analyzing movements in heterogeneous landscapes from mark-recapture data. Ecology 89(2):542–554
- Ovaskainen O, Smith AD, Osborne JL, Reynolds DR, Carreck NL, Martin AP, Niitepõld K, Hanski I (2008c) Tracking butterfly movements with harmonic radar reveals an effect of population age on movement distance. PNAS 105(49):19090–19095
- Palla G, Derenyi I, Farkas I, Vicsek T (2005) Uncovering the overlapping community structure of complex networks in nature and society. Nature 435(7043):814–818
- Paris CB, Chérubin LM, Cowen RK (2007) Surfing, spinning, or diving from reef to reef: effects on population connectivity. Mar Ecol Prog Ser 347:285–300
- Parker PG, Snow AA, Schug MD, Booton GC, Fuerst PA (1998) What molecules can tell us about populations: choosing and using a molecular marker. Ecology 79(2):361–382
- 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. Landscape Ecol 21(7):959–967
- Patterson TA, Evans K, Carter TI, Gunn JS (2008a) Movement and behaviour of large southern bluefin tuna (*Thunnus* maccoyii) in the Australian region determined using popup satellite archival tags. Fish Oceanogr 17(5):352–367
- Patterson TA, Thomas L, Wilcox C, Ovaskainen O, Matthiopoulos J (2008b) State-space models of individual animal movement. Trends Ecol Evol 23(2):87–94
- Pauli JN, Ben-David M, Buskirk SW, Depue JE, Smith WP (2009) An isotopic technique to mark mid-sized vertebrates non-invasively. J Zool 278(2):141–148
- Pineda J, Hare JA, Sponaugle S (2007) Larval transport and dispersal in the coastal ocean and consequences for population connectivity. Oceanography 20(3):22–39
- Pinsky ML, Montes HR, Palumbi SR (2010) Using isolation by distance and effective density to estimate dispersal scales in anemonefish. Evolution 64:2688–2700
- Planes S, Jones GP, Thorrold SR (2009) Larval dispersal connects fish populations in a network of marine protected areas. PNAS 106:5693–5697
- Porter J, Arzberger P, Braun H-W, Bryant P, Gage S, Hansen T, Hanson P, Lin C-C, Lin F-P, Kratz T, Michener W, Shapiro S, Williams T (2005) Wireless sensor networks for ecology. Bioscience 55(7):561–572
- Pressey RL, Cabeza M, Watts ME, Cowling RM, Wilson KA (2007) Conservation planning in a changing world. Trends Ecol Evol 22(11):583–592

- Pringle C (2003) The need for a more predictive understanding of hydrologic connectivity. Aqua Conserv Mar Freshw Ecosyst 13(6):467–471
- Proulx SR, Promislow DEL, Phillips PC (2005) Network thinking in ecology and evolution. Trends Ecol Evol 20(6):345–353
- Raeymaekers JAM, Maes GE, Geldof S, Hontis I, Nackaerts K, Volckaert FAM (2008) Modeling genetic connectivity in sticklebacks as a guideline for river restoration. Evol Appl 1(3):475–488
- Randall LA, Diehl RH, Wilson BC, Barrow WC Jr, Jeske CW (2011) Potential use of weather radar to study movements of wintering waterfowl. J Wildl Manag 75(6):1324–1329
- Ray N (2005) PATHMATRIX: a geographical information system tool to compute effective distances among samples. Mol Ecol Notes 5(1):177–180
- Rayfield B, Fortin M-J, Fall A (2011) Connectivity for conservation: a framework to classify network measures. Ecology 92(4):847–858
- Recio MR, Mathieu R, Denys P, Sirguey P, Seddon PJ (2011) Lightweight GPS-tags, one giant leap for wildlife tracking? An assessment approach. PLoS ONE 6(12):e28225
- Roberts JJ, Best BD, Dunn DC, Treml EA, Halpin PN (2010) Marine geospatial ecology tools: an integrated framework for ecological geoprocessing with ArcGIS, Python, R, MAT-LAB, and C++. Environ Model Softw 25(10):1197–1207
- Rubenstein DR, Hobson KA (2004) From birds to butterflies: animal movement patterns and stable isotopes. Trends Ecol Evol 19(5):256–263
- Rubio L, Saura S (2012) Assessing the importance of individual habitat patches as irreplaceable connecting elements: an analysis of simulated and real landscape data. Ecol Complex 11:28–37
- Rutz C, Hays GC (2009) New frontiers in biologging science. Biol Lett 5(3):289–292
- Saenz-Agudelo P, Jones GP, Thorrold SR, 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(8):1765–1776
- Sarkar S, Pressey RL, Faith DP et al (2006) Biodiversity conservation planning tools: present status and challenges for the future. Annu Rev Environ Resour 31:123–159
- Saura S, Pascual-Hortal L (2007) A new habitat availability index to integrate connectivity in landscapte conservation planning: comparison with existing indices and application to a case study. Landsc Urb Plan 83:91–103
- Saura S, Rubio L (2010) A common currency for the different ways in which patches and links can contribute to habitat availability and connectivity in the landscape. Ecography 33(3):523–537
- Saura S, Torné J (2009) Conefor Sensinode 2.2: a software package for quantifying the importance of habitat patches for landscape connectivity. Environ Model Softw 24(1):135–139
- Selkoe KA, Henzler CM, Gaines SD (2008) Seascape genetics and the spatial ecology of marine populations. Fish 9(4):363–377
- Slone DH (2011) Increasing accuracy of dispersal kernels in grid-based population models. Ecol Model 222(3): 573–579

- Storfer A, Murphy MA, Spear SF, Holderegger R, Waits LP (2010) Landscape genetics: where are we now? Mol Ecol 19(17):3496–3514
- Sutherland WJ (ed) (2006) Ecological census techniques: a handbook. Cambridge University Press, Cambridge
- Taylor PD, Fahrig L, Henein K, Merriam G (1993) Connectivity is a vital element of landscape structure. Oikos 68(3): 571–573
- Thorrold SR, Zacherl DC, Levin LA (2007) Population connectivity and larval dispersal using geochemical signatures in calcified structures. Oceanography 20(3):80–89
- Treml EA, Halpin PN (in press) Marine population connectivity identifies ecological neighbors for conservation planning in the Coral Triangle. Cons Lett. doi:10.1111/j.1755-263X. 2012.00260.x
- Treml EA, Halpin PN, Urban DL, Pratson LF (2008) Modeling population connectivity by ocean currents, a graph-theoretic approach for marine conservation. Landscape Ecol 23(S1):19–36
- Tsafrir D, Tsafrir I, Ein-Dor L, Zuk O, Notterman DA, Domany E (2005) Sorting points into neighborhoods (SPIN): data analysis and visualization by ordering distance matrices. Bioinformatics 21(10):2301–2308
- Tsechpenakis G, Guigand C, Cowen RK (2007) Image analysis techniques to accompany a new in situ Ichthyoplankton imaging system. In: OCEANS 2007—Europe, pp 1–6
- Turner W, Spector S, Gardiner N, Fladeland M, Sterling E, Steininger M (2003) Remote sensing for biodiversity science and conservation. Trends Ecol Evol 18(6):306–314
- Urban D, Keitt T (2001) Landscape connectivity: a graph-theoretic perspective. Ecology 82(5):1205–1218
- Urban DL, Minor ES, Treml EA, Schick RS (2009) Graph models of habitat mosaics. Ecol Lett 12(3):260–273
- Urbano F, Cagnacci F, Calenge C, Dettki H, Cameron A, Neteler M (2010) Wildlife tracking data management: a new vision. Phil Trans R Soc Lond B 365(1550):2177–2185
- Van Teeffelen AJA, Ovaskainen O (2007) Can the cause of aggregation be inferred from species distributions? Oikos 116(1):4–16
- Visconti P, Elkin C (2009) Using connectivity metrics in conservation planning—when does habitat quality matter? Divers Distrib 15(4):602–612

- Wang A, Afshar P, Wang H (2008) Complex stochastic systems modelling and control via iterative machine learning. Neurocomputing 71(13–15):2685–2692
- Wang K, Franklin SE, Guo X, Cattet M (2010) Remote sensing of ecology, biodiversity and conservation: a review from the perspective of remote sensing specialists. Sensors 10:9647–9667
- Waples RS, Gaggiotti OE (2006) What is a population? An empirical evaluation of some genetic methods for identifying the number of gene pools and their degree of connectivity. Mol Ecol 15:1419–1439
- Wasserman TN, Cushman SA, Shirk AS, Landguth EL, Littell JS (2012) Simulating the effects of climate change on population connectivity of American marten (*Martes americana*) in the northern Rocky Mountains, USA. Landscape Ecol 27:211–225
- Watts ME, Ball IR, Stewart RS, Klein CJ, Wilson K, Steinback C, Lourival R, Kircher L, Possingham HP (2009) Marxan with zones: software for optimal conservation based landand sea-use zoning. Environ Model Softw 24(12): 1513–1521
- Webster MS, Marra PP, Haig SM, Bensch S, Holmes RT (2002) Links between worlds: unraveling migratory connectivity. Trends Ecol Evol 17(2):76–83
- West JB, Bowen GJ, Cerling TE, Ehleringer JR (2006) Stable isotopes as one of nature's ecological recorders. Trends Ecol Evol 21(7):408–414
- White C, Selkoe KA, Watson J, Siegel DA, Zacherl DC, Toonen RJ (2010) Ocean currents help explain population genetic structure. Proc R Soc B Biol Sci 277(1688):1685–1694
- Williamson David H, Jones GP, T SR, Frisch AJ (2009) Transgenerational marking of marine fish larvae: stableisotope retention, physiological effects and health issues. J Fish Biol 74(4):891–905
- Woods RJ, Macdonald JI, Crook DA, Schmidt DJ, Hughes JM (2010) Contemporary and historical patterns of connectivity among populations of an inland river fish species inferred from genetics and otolith chemistry. Can J Fish Aquat Sci 67:1098–1115