Title: Ecological networks: Pursuing the shortest path, however narrow and crooked

Running title: *Shortest paths in ecological networks*

Andrea Costa[1,2]*, Ana M. Martín González[3]*, Katell Guizien[4], Andrea M. Doglioli[5], José María Gómez[6],

Anne A. Petrenko[5], Stefano Allesina[7,8]

* shared first authorship

1- Center for Climate Physics, Institute for Basic Science, Busan, South Korea, 46241

2- Department of Atmospheric Sciences, Pusan National University, Busan, South Korea, 46241

3- Center for Macroecology, Evolution and Climate, Natural History Museum of Denmark, University of

Copenhagen, Copenhagen, Denmark

4- CNRS, Sorbonne Universités (UPMC Univ. Paris 06), Laboratoire d'Ecogéochimie des Environ-

nements Benthiques (LECOB), 1 avenue Pierre Fabre Observatoire Océanologique, Banyuls/Mer,

France

5- Aix Marseille Université, CNRS, Université de Toulon, IRD, OSU Pythèas, Mediterranean Institute of

Oceanography (MIO), UM 110, 13288, Marseille, France

6- Departamento de Ecología Funcional y Evolutiva, Estación Experimental de Zonas Áridas (EEZA-

CSIC), Almería, Spain

7- Department of Ecology and Evolution, University of Chicago, Chicago, IL, USA

8- Northwestern Institute on Complex Systems, Northwestern University, Evanston, IL, USA

Corresponding authors: Andrea Costa (IBS Center for Climate Physics, Tonghapgigyegwan Bldg.

Busandaehak-ro 63beon-gil 2, Jangjeon-dong, Geumjeong-gu, Busan 46241, Republic of Korea,

+82 515107641) and **Ana M. Martín González** (Center for Macroecology, Evolution and Climate, Nat-

ural History Museum of Denmark, University of Copenhagen. Universitetsparken 15, Building 3. 2100

Copenhagen V. +45 35333734).

ABSTRACT

1. Representing data as networks cuts across all sub-disciplines in ecology and evolutionary biology. Besides providing a compact representation of the interconnections between agents, network analysis allows the identification of especially important nodes, according to various metrics of network centrality. These centrality measures often rely on the calculation of the shortest path connecting any two nodes, and while the interpretation of a shortest paths is straightforward in binary, unweighted networks, whenever weights are accounted for, inconsistency between weight definition and shortest path interpretation could yield unexpected results.

2. Here we review 129 studies of ecological networks published in the last decade and making use of shortest paths.

3. We evidenced a methodological inaccuracy related to the edge weights used to calculate shortest paths (and related centrality measures) in ecological studies, particularly in interaction networks. Specifically, we found numerous studies in which the edge weights were not transformed prior to calculating shortest paths when the edge weights were proportional to the information flow between the nodes of the network at study. 49% of the studies do not report sufficient information on the calculation to allow their replication, and 61% of the studies on weighted networks may contain errors in how shortest paths are calculated. Using toy models and empirical ecological data, we show how to transform the data prior to calculation and illustrate the pitfalls that need to be avoided.

4. We conclude by proposing a five-point check-list to foster best-practices in the calculation and reporting of centrality measures in ecology and evolution studies.

Keywords: Betweenness centrality, Closeness centrality, graph theory, interaction networks, keystone species, landscape networks, shortest path, stepping stones.

50   *Pursue some path, however narrow and crooked, in*

51   *which you can walk with love and reverence.*

52   *– Henry David Thoreau*

INTRODUCTION

The last two decades have witnessed an exponential increase in the use of graph analysis in

ecological and conservation studies (see Dale, 2017 and Delmas et al. 2018 for recent introductions to

network theory in ecology and evolution). Networks (graphs) represent agents as nodes linked by

edges representing pairwise relationships. For instance, a food web can be represented as a network

of species (nodes) and their feeding relationships (edges) (e.g., Brose et al., 2005). Similarly, the

spatial dynamics of a metapopulation can be analyzed by connecting the patches of suitable habitat

(nodes) with edges measuring dispersal between patches (e.g., Altermatt, Seymour & Martinez,

2013). Data might either simply report the presence/absence of an edge (binary, unweighted

networks), or provide a strength for each edge (weighted networks). In turn, these weights can

represent a variety of ecologically-relevant quantities, depending on the system being described. For

instance, edge weights can quantify interaction frequency (e.g., visitation networks, Martín González

et al., 2015), interaction strength (e.g., per-capita effect of one species on the growth rate of another,

Brose et al., 2005), carbon-flow between trophic levels (Jordán, Liu, & Davis, 2006), genetic similarity

(Rozenfeld et al., 2008), niche overlap (e.g., number of shared resources between two species, Pires,

Marquitti, & Guimarães, 2017), affinity (e.g. Luthe & Wyss, 2016), dispersal probabilities (e.g., the rate

at which individuals of a population move between patches, Zamborain-Mason, Russ, Abesamis,

Bucol, & Connolly, 2017), cost of dispersal between patches (e.g., resistance, Girardet, Conruyt-

Rogeon, & Foltête, 2015), etc.

Despite such large variety of ecological network representations, a common task is the identification of

nodes of high importance, such as keystone species in a food web, patches acting as stepping stones

in a dispersal network, or genes with pleiotropic effects. The identification of important nodes is

typically accomplished through centrality measures (e.g., Treml, Halpin, Urban, & Pratson, 2008;

Martín González, Dalsgaard, & Olesen, 2010). A large number of centrality measures has been

proposed, each probing complementary aspects of node-to-node relationships (e.g., Kool, Moilanen, &

Treml, 2013). For instance, Closeness centrality (Bavelas, 1950; Freeman, 1979) highlights nodes that

are "near" to all other nodes in the network in terms of average distance from all other nodes.

Whenever the effects of a node on another weaken along the path (Burt, 2000), then central nodes

are those having the largest capacity to influence the others. Consider however highly modular

networks, in which tightly knit communities of nodes are loosely connected to one another; then, one

may be interested in identifying nodes that act as bridges connecting the different communities,

allowing for the spread of perturbations across the entire network. Stress centrality (Shimbel, 1953),

and Betweenness centrality (Freeman, 1979) serve this purpose. The choice of a centrality measure

thus depends on the research question at hand, and on the characteristics of the data being analyzed.

Different centrality measures have been used to identify keystone species in networks of biotic

interactions (Jordán, 2009; Martín González, Dalsgaard, & Olesen, 2010), to explore the robustness of

metapopulations (Thompson, Rayfield, & Gonzalez, 2015), to describe connectivity patterns across

fragmented habitats (Carroll, McRae, & Brookes, 2012), to explore social behavior and pathogen

spread within populations (Aplin et al., 2013), and to provide a theoretical background to support

decision-making in conservation planning and urban management (Girardet, Conruyt-Rogeon, &

Foltête, 2015; Poodat, Arrowsmith, Fraser, & Gordon, 2015; see Supporting Information for a complete

list). Given the wide array of available techniques and the span of ecological applications, confusion

may arise when performing and reporting centrality analysis. An understanding of how the calculations

are performed, as well as a clear and sufficient reporting of the details of the analysis, are necessary

in order to avoid misinterpretation of the results and ensure the reproducibility of published studies.

In particular, edge weights exert a substantial influence on all measures of centrality. Many centrality

metrics rely on calculating the shortest path connecting any two nodes, and for weighted edges this

translates into finding the path with the smallest sum of weights. Edge weights definition is crucial in

all measures of centrality. When edge weights represent cost, resistance, or in general scale inversely

to the strength of the relationship between two nodes, then the definition of "shortest" paths retains its

simple interpretation. However, whenever edge weights are proportional to the strength of the

relationship between two nodes (e.g., probability of dispersal, interaction frequency, contact rate, etc.),

then minimizing the sum of edges along the path makes no sense: the data need to be transformed

prior to the analysis, or one has to choose an appropriate method able to deal with this situation. This

issue has been raised before (e.g., Foltête, Clauzel, Vuidel, & Tournant, 2012a; Costa, Petrenko,

Guizien, & Doglioli, 2017) and is widely acknowledged among studies describing community structure,

where measures of network structure have been specifically developed to account for weighted edges

(e.g., Newman, 2004; Almeida-Neto, & Ulrich, 2011; Dormann, & Strauss, 2014). However, our review

suggests that this issue is not fully resolved, and that incorrect interpretations of centrality measures

linger on, particularly among weighted networks.

Furthermore, most published studies do not report with sufficient detail the calculation of the node-to-

node distance definition used in conjunction with the calculated centrality measures. As a

consequence, it is often impossible to evaluate the correctness of the calculations. We have

performed a systematic review of the ecological network literature, selecting ecological studies from

the Web of Science that use a network approach. We searched on topics TS=(network AND ecolog*),

limiting our search to Articles written in English in the science-related citation indexes (SCI-

EXPANDED, CPCI-S, BKCI-S, ESCI) since 2006 and added 22 records obtained from other sources.

From the resulting articles, we refined our search to those mentioning "centrality" (n = 210). We further

refined our set by choosing studies that studied ecological communities using centrality metrics

requiring the calculation of shortest paths and discarding purely methodological studies. Finally, armed

with a list of 129 articles, we checked whether the analysis was reported with sufficient detail to

determine whether the calculations were appropriate. In Figure 1 we summarize this information with a

frequency chart and in the Supporting Information we provide the full list of articles. 63 articles (49%)

did not report sufficient information on the calculation of centrality to allow their replication, six of which

were unclear even whether edges were binary or weighted. Moreover, 61% of the studies using

weighted edges may contain errors in how shortest path centralities are calculated, a figure that grows

to 89% if we limit the analysis to the case of weighted interaction networks. Noticeably, 88% of the

studies that correctly accounted for weighted edges in the calculation of shortest paths, considered

networks in which weights are inversely proportional to the strength of association – thus not requiring

transformation. Furthermore, nine studies (eight of which examine interaction networks) calculated

centrality using the binary version of the weighted data, without providing any justification for this

methodological choice.

The final goal of this review is to offer a precise and detailed protocol for the calculation of shortest

path centralities in ecological networks. Given that the correct way to calculate shortest path-based

centrality measures depends on the type of network considered (binary or weighted) and on the

meaning of edge weights, we use simple examples as well as real data to show how different

approaches can result in unreliable estimates of centrality. Finally, since most studies omit to report

key aspects of the definition of the edge weights, we propose a simple checklist to foster best-

practices in the calculation and reporting of shortest path-related centrality analysis.

THE SHORTEST PATH IS FULL OF PITFALLS

In network analysis, the interaction between nodes can be thought of as a flow of information between

the nodes that are linked by edges. The sequence of edges that information must cross in order to

reach a specific node is called a path. It is generally assumed that the bulk of information between any

two given nodes (among all the possible paths between these two nodes) passes through the shortest

path connecting them (i.e., the one with "lowest weight"). However, it should be emphasized that while

the concept of information flow is general, its immanence can differ dramatically from case to case,

depending on which network feature weights quantify. For reference, in Table 1 we list the principal

types of networks and the proportionality of the edges to the information flow between nodes found in

the ecological literature.

The interpretation of a shortest path as the path that funnels the bulk of information flow relies on it

being the least weight path (i.e., the path of least resistance) between two nodes. Indeed, all the

shortest path algorithms currently available (e.g., Djikstra, 1959; Brandes, 2008) and generally

implemented in graph theory software (Table 2) seek to minimize the value of the path between two

nodes calculated as the sum of the edge weights. The reason being that a minimization problem

converges, while maximization can fail (see Costa, Petrenko, Guizien, & Doglioli, 2017 for a detailed

explanation). Nevertheless, the identification of the shortest paths is far from trivial, as one has to pay

attention to what edge weights represent. That is, one has to ensure that the edge weight is inversely

proportional to the flow of information between the nodes. This condition is automatically fulfilled if the

natural weight suggested by the network at study is already inversely proportional to the information

flow (e.g., resistance distance, dispersal time). However, when the weight is directly proportional to the

information flow (e.g., interaction frequency, individual transfer, dispersal probability, pathogen

transmission), it is necessary to transform the edge weight in order to calculate the shortest paths, and

the centrality measures that rely on them (Table 3). In particular, this is important when using user-

friendly software packages that automate the calculation of centrality measures (Table 2).

There is a wide range of functions that can accomplish this transformation. For example, if a_{ij}

measures the flow of information between nodes *i* and *j*, the functions $1-a_{ij}$, $\exp(-a_{ij})$, $1/a_{ij}$, $\log(1/a_{ij})$ and

$\log(a_{ij}/(1-a_{ij}))$ are all found in the graph-theoretical literature (e.g., Augustin, Cummins, & French, 2001;

Estrada & Bodin, 2008; Brandes, 2008; Opsahl, Agneessens, & Skvoretz, 2010; Crandal et al., 2014).

Note that one must pay attention to the range of values that the edge weights can span, and to the

values induced by the transformation – for example, one must avoid the use of negative edges (e.g.,

log transformation of values between 0 and 1), as these can greatly hamper the interpretation of the

results. Finally, if the edge weights represent probabilities, one should account for the independence

(or lack of independence) of the different edges.

AVOIDING THE PITFALLS

When computing any shortest path-related measure, various decisions need to be made, and the

wrong decision could lead to unexpected results, as we will illustrate in the following examples. We

focus on the effects on Betweenness (BC) and Closeness centralities (CC) as these are the most

commonly used centrality measures. In Figure 2 we summarize this decision process.

Binary or weighted?

The first methodological choice when computing shortest paths is whether to consider edge weights.

Binary data can be highly informative: for example, it has been used to identify species fundamental

niches (e.g., Fründ et al., 2016), and key species in pollination networks (Martín González et al.,

2012). Nonetheless, studies must clearly state whether the analysis is performed on weighted or

unweighted networks, and provide an ecological justification supporting either choice (e.g., Mello et

al., 2015). Seminal studies (e.g., Minor & Urban, 2007) as well as recent ones (e.g., Zamborain-

Mason, Russ, Abesamis, Bucol, & Connolly, 2017) analyzed unweighted versions of their data arguing

that paths with fewer edges would inherently be stronger than those composed of multiple ones. In our

review of the literature, we have found nine studies that, despite using weighted data for some

calculations, revert to binary versions of the network for the calculation of centrality measures without

providing a justification (Figure 1). Indeed, note that unipartite projections of bipartite networks result

in weighted networks even when the original network is binary. Seven articles out of nine use binary

unipartite projections without any justification. Although calculations in these are technically correct,

one must be aware that the calculation of shortest paths and related measures in binary and weighted

versions of the same network can lead to dramatically different conclusions.

To see how discarding the edge weights can significantly change the results of network analysis, let us

start with a highly idealized example. In Figure 3a we show a network such that there are two possible

paths between from Busan (South Korea) to Almería (Spain): one containing two edges (Busan-

Chicago-Almería), and the other one three (Busan-Copenhagen-Marseille-Almería). If one considers

only the binary data (implicitly assuming that the distance between any cities is the same), the

shortest path from Busan to Almería would be crossing Chicago (one stopover vs. the two stopovers

of the other possible path). However, if the distance between cities is considered, it can be easily seen

that the path Busan-Copenhagen-Marseille-Almería is much shorter (~11000 vs. ~18000 Km).

For an ecologically-relevant example, we constructed a connectivity matrix for a hypothetical bird

species living between 500 and 2000 m above sea-level, and with a typical habitat size of about 15

km^2. For this purpose, the Global Relief Database ETOPO1 (Amante & Eakins, 2009) data in the

region we considered were coarse-grained to 15 km^2 horizontal resolution, resulting in 787 habitat

patches (Figure 4a). Dispersal probability between patches was calculated as $p_{ij} = \exp(-\alpha\, d_{ij})$ following

Minor & Urban (2007), where d_{ij} is the geographical distance between the patches boundaries, and α

is a parameter chosen to be 0.03 in order to have a (hypothetical) median dispersal distance of 100

km. This probability can be stored in a connectivity matrix (see Supporting Information, Figure S1) and

graph theory can be used to identify the habitat patches that have high Betweenness and Closeness

centrality scores. Calculating BC and CC on binary and weighted versions of this dataset resulted in

markedly different outcomes. None of the 20 habitat patches with highest BC and CC in the binary

network (Figure 4b-c) match the ones obtained from the analysis of the weighted network (Figure 4d-

e). Note that to calculate Betweenness and Closeness centralities in weighted data we first inverted

the edge weights using $\log(1/p_{ij})$ (see next section for a detailed explanation).

Modifying edge weights

The next question one should answer when computing shortest paths is whether edge weights are

inversely proportional to the information flow between the nodes in the network (Figure 2). If this is the

case, the shortest path between nodes can be calculated directly using the edge weights. If, on the

other hand, edge weights are proportional to the flow of information, one must transform them before

using shortest path algorithms. If one does not modify the edge weights, the shortest path algorithms

will either fail (and identify the longest, rather than shortest path), or will be unable to identify a

shortest path at all. We use the transformations $1/a_{ij}$ or $\log(1/a_{ij})$ for the edges (although for this

operation there are several alternative options reviewed below).

For example, let us consider a small network of four famous primates sharing a certain number of

parasites (Figure 5a). In order to detect the primate mediating the transmission of infectious diseases

in this network, one could identify the primate displaying the largest number of parasites common to

other primates – indicated by stronger edges. In this simple network it is easy to verify that most of the

paths with the highest edge weight (largest number of shared parasites) pass through Rafiki.

However, if BC and CC were calculated directly on unmodified edge weights, we would conclude that

Abu is the key primate in this network (Figure 5a). If, on the other hand, we transform edge weights

using the function $1/a_{ij}$, we correctly identify Rafiki as the primate with the highest BC and CC (Figure

5b).

Using a real-world case, the impact of not reversing edge weights can be illustrated using the Global

Mammal Parasites Database (GMPD, https://parasites.nunn.lab.org) containing data on 542 primate

species and their 750 parasites (Stephens et al., 2017 and references therein). To identify the species

mediating the transmission of parasites (similarly to Gómez, Nunn & Verdú , 2003), a connectivity

matrix linking primates that have been found to host the same parasite was built (see Supporting

Information, Figure S2). Edge weights were defined as the number of shared parasites between any

two species. Therefore, edge weight is directly proportional to the relationship strength between two

species and, as in the previous example, the quantity should be transformed before calculating the

shortest paths. In Figure 6a we show the lack of correlation between species rankings based on the

centrality scores calculated on modified and unmodified edge weights. For this example, we use the

inverse of the edge weights, e.g., $1/a_{ij}$. Interestingly, the BC scores calculated using the modified edge

weights highlight only few species, one of which has by far the highest BC score. Instead, if we

directly use the unmodified values, several more species have comparable BC scores. This is not

surprising if we consider that, when using the raw weights, shortest paths pass through weak

connections, which are likely to be numerous. Differences in ranks are substantial. For instance,

among the top ten high-Betweenness species identified using modified weights, only one is also

among the high-Betweenness species identified using raw weights (see Supporting Information, Table

S1).

Likewise, the results from the Closeness-based rankings (Figure 6b) show that species rankings

based on CC also differ significantly between modified and unmodified edge weights. The CC scores

calculated on modified edge weights also support the importance of a handful of species (see

Supporting Information, Table S1). Unlike with BC, nine of the top 10 high-Closeness species are the

same ones when using the unmodified weights (also in Supporting Information) but the exact ranking

differs between the two cases.

Other modifying functions

Adding constants

When edges are directly proportional to the information flow, it is frequent practice to make them

inversely proportional by subtracting their value from a theoretical maximum or some other meaningful

constant (Brandes, 2008; Ophsal, 2010). For example, in the case of dispersal probabilities, one could

choose to subtract the edge weights a_{ij} from 1. However, the new edge weight $1-a_{ij}$ biases the

calculation of the shortest paths towards the path with the lowest number of edges (because

probabilities sum to one, nodes with many edges tend to have lower values).

Again, we will use the simple toy matrix presented in Figure 3, where we show a network connecting

different cities. We then consider three different quantities to weight the edges that represent the

movement of researchers between these cities. In the first case (Figure 3b), edge weights quantify the

number of researchers who moved from one site to another. In this case, the path sustaining the

largest "flow of researchers" between Busan and Almería is the three-steps path Busan-Copenhagen-

Marseille-Almería (30 vs 10). However, applying a shortest path algorithm directly to this network

would identify the two-step path as more important. One possible way to reverse the edge weight is to

subtract the edge weights from a large (and in most instances arbitrary) constant C – de facto adding

a constant to all edges. For example, if one chooses C = 100, now the largest edge weights (those

representing largest flows) are the smallest and would hence be identified by the shortest-path

centrality algorithm as more central. However, one can easily verify that such a transformation did not

change the fact that the two-step path is the shortest between Busan and Almería (190 vs 270). The

reason is that adding a constant to all the edge weights biases the shortest paths algorithm towards

paths with fewer edges.

In Figure 3c we see that subtracting the edge weights from a theoretical maximum C (in the case of

transfer probabilities of interaction frequencies, C = 1) makes the three-step path with higher

information flow the shortest path (3x(1-0.9) vs 2x(1-0.1)). However, this transformation does not work

in all cases. In fact, if the edge weights, as frequently occurs in ecological studies, span different

orders of magnitude (e.g., Figure 3d), the three-step path will not be the shortest path anymore (3x(1-

10^{-3}) vs 2x(1-10^{-5})), as in Figure 3b. Furthermore, we note that this type of transformation, even when

it is likely to work, cannot be used for all the edge weights. For example, it cannot be used for

probabilities, as the values of log(1-a_{ij}) are negative and, consequently, cannot be used to find shortest

paths (see next section).

A real-world example of the effect of adding constants to the edge weights is provided in the

Supporting Information (Figures S3, S4 and S5).

Negative weights and loops

Another way to reverse the edge weights is to reverse the sign of the weights (i.e., using -a_{ij}).

However, as shortest path algorithms seek to minimize the value of a path, they would keep looping

closed paths (cycles) ad infinitum, without ever converging. It must be noted that there are alternative

algorithms that can handle negative edges values (e.g., the Bellman-Ford-Moore algorithm, Bang-

Jensen & Gutin, 2000) cannot handle cycles. As cycles are essentially ubiquitous in ecological

applications, edge weight transformations that result in negative values should therefore be avoided.

As an example, consider a simple toy network depicting the carbon flow between different layers of

the food chain (Figure 7a). In this case, given that the flow of carbon is directly proportional to the

strength of the connection between two layers of the food chain, we need to transform the weights.

However, if one uses $-a_{ij}$ (Figure 7b), the shortest path algorithms would never converge, and would

keep circling the loop. On the other hand, using another weight reversing function, such as $1/a_{ij}$, would

correctly identify Merlin and Wart as key species pivoting the carbon flow in the moat's food chain

network (Figure 7c).

Independence of probabilities

Finally, another important aspect to consider when modifying the edge weight is whether edges along

a path are independent from each other. For instance, in the case of genetic diversity, one can

reasonably assume that genetic differences between populations are independent from one another.

However, when the edges a_{ij} represent probabilities (e.g., dispersal probabilities), the value of a path

from node A to node C passing through node B must consider the dependence of the edges AB and

BC. This can be done by defining the edge weights as $\log(1/a_{ij})$ so that the shortest path algorithms

will calculate the path values as the product of the a_{ij} values (see Costa, Petrenko, Guizien & Doglioli,

2017 for a detailed explanation). In a real-world case, we show the consequences of not accounting

for the independence of the edges' values using the ETOPO1 dataset (see Supporting Information,

Figure S6).

One or all shortest paths?

In most networks, whether binary or weighted, there may be more than one shortest path connecting

any two nodes. To account for this fact, Newman (2005) introduced an alternative definition of

Betweenness centrality based on random walks and Opsahl, Agneessens, & Skvoretz (2010) a

generalization of node centrality that considers both edge weight and number when calculating

centrality measures. Although the discussion of the pros and cons of the different formulations falls

beyond the scope of this review, we encourage the reader to be aware that considering a single or all

shortest paths may also introduce differences in the resulting centrality values, and the decision

should hence also be reported.

WIDENING THE PATH

Network analysis has been developing quite independently in different branches of ecology. However,

dissemination between ecological disciplines and reproduction of published studies are being

hampered, at least partially, by the lack of transparency when describing the methodologies used.

Establishing a protocol for the analysis and reporting of calculations would ease these obstacles. For

this reason, we urge all the researchers applying graph theory to ecological data to pay special

attention when reporting their calculations, and, in particular, to provide a description of the network

and edge weight they used.

Here is a checklist of crucial methodological information that should always be reported (Figure 2):

1) A clear definition of what nodes and edges represent.

2) Are edges binary or weighted? If weighted, report the proportionality of the edge weight to the

information flow between the nodes in order to evaluate whether edges need to be modified.

3) Report the eventual transformation applied to edge weight before the calculation of centrality

measures. Moreover, carefully justify any conceptual reason to not transform edge weights, or to use

the unweighted versions of weighted data. For example, in a species-interaction network (where

species are typically considered closer if they interact with higher frequencies), one could purposely

choose to calculate shortest path-centrality measures without transforming the weights in order to

study the effect of weak interactions across the network.

4) Report the formula used for the calculation of the centrality measure and whether it considers all

shortest paths or a single one.

5) Report the full version of the software or package used for the calculation of centrality.

CONCLUSIONS

Graph theory enables to achieve precious insights on ecological networks. For this reason, it has gained popularity in ecology and has developed quite independently in different disciplines, becoming a routine analysis in ecological studies. Our review of the literature evidenced that this familiarity is however associated to a lack of methodological rigor in the published studies. Indeed, by reading the methodological sections of a large portion of the published studies, we were not able to clearly ascertain what edges represented when centrality measures calculations were carried out.

Using both theoretical and real-world case studies we showed that such an oversight can lead to radically different results. We provide an overview of different methods to meaningfully calculate shortest paths and related centrality measures in ecological systems, and a checklist to ensure clear and sufficient reporting of such calculations. We hope that following the protocol we suggest will further increase the popularity of centrality measures in ecology, and, at the same time, guarantee the reproducibility of these studies.

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AUTHOR'S CONTRIBUTIONS

AC, AMMG, SA, AAP, KG & AMD designed the study; AC conducted the data analysis, AMMG performed the literature search and AC & AMMG performed the review and wrote the first draft of the manuscript. All authors contributed substantially to the drafts and gave final approval for publication.

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Tables

Table 1. Summary of different network types, describing the type of information flowing along edges and the consequences for weight transformation. Landscape networks are spatially-explicit, while interaction networks depict relationships among entities. Social networks are here included as interaction networks, where shared traits include shared interacting organisms, foraging time, etc. depicting social relationships or common characteristics.

Network type	Edge weight	Proportionality to information flow	Requires transformation	Example references
Landscape	Cost-distance	Inverse	No	Carroll, McRae & Brookes, 2012; Girardet, Conruyt-Rogeon & Foltête, 2015.
	Dispersal probability	Direct	Yes	Magris, Treml, Pressey, & Weeks, 2016; Zamborain-Mason, Russ, Abesamis, Bucol, & Connolly, 2017
	Dispersal time	Inverse	No	Treml, Halpin, Urban, & Pratson, 2008
	Exchange of individuals	Direct	Yes	Emer et al., 2018
	Genetic similarity	Direct	Yes	Herrera-Arroyo et al., 2013; Naujokaitis-Lewis, Rico, Lovell, Fortin, & Murphy 2013
Interaction	Frequencies	Direct	Yes	Ruggera, Blendinger, Gomez, & Marshak, 2015; Lozano, Mateos, & Rodríguez, 2016
	Shared traits, affinity	Direct	Yes	Reino et al., 2017; Blaszczyk 2018
	Trophic flow	Direct	Yes	Livi, Jordán, Lecca, & Okey, 2011; Lai, Liu, & Jordán, 2012

551　Table 2. Analytical packages commonly used for the calculation of centrality in ecological studies and

552　references for each package. Only *sna* and *Ucinet* packages caution in their documentation about

553　edge transformations when calculating shortest paths.

Package	References
igraph	Csardi & Nepusz (2006) http://igraph.org/
sna	Butts (2016) https://cran.r-project.org/web/packages/sna/index.html
tnet	Opsahl (2015) https://cran.r-project.org/web/packages/tnet/index.html
Pajek	Mrvar & Batagelj (2016) http://mrvar.fdv.uni-lj.si/pajek/
Ucinet	Borgatti, Everrett & Freeman (2002) https://sites.google.com/site/ucinetsoftware/home
CONEFOR	Saura & Torné (2009) http://www.conefor.org/
Graphab	Foltête, Clauzel & Vuidel (2012b) https://sourcesup.renater.fr/graphab/en/home.html

554　Table 3. Measures of shortest path-related centrality measures commonly used in ecological network analysis. Notice that other common

555　centrality measures are based on eigenvectors or dissimilarity scores instead of on the identification of shortest paths and are hence not

556　reviewed in this work.

Centrality Measure	Definition	Formula	Intended Network type	Reference
Betweenness, BC	Quantifies the proportion of shortest paths g between any two nodes i, j, that pass through a focal node v.	$$BC(v) = \sum_{i \neq v \neq j} \frac{g_{ij}(v))}{g_{ij}}$$	All types	Freeman, 1979
Stress, SC	Measures the number of shortest paths g between any two nodes i, j, that pass through a focal node v.	$$SC(v) = \sum_{i \neq v \neq j} g_{ij}(v)$$	All types	Shimbel, 1953
Closeness, CC	Measures the average length of the shortest paths from a node v to all the other nodes in the network.	$$CC(v) = \frac{N-1}{\sum_j g_{vj}}$$	All types	Freeman, 1979
Integral Index of Connectivity, IIC	Measures the degree of connectivity of the entire landscape (of total area A_L) through the calculation of the number of edges in the shortest path nl_{ij} between patches with area a_i and a_j.	$$IIC(v) = \frac{\sum_{i=1}^{n} \sum_{j=1}^{n} a_i a_j / (1 + nl_{ij})}{A_L^2}$$	Binary landscape networks	Pascual-Hortal & Saura, 2006
Probability of Connectivity Index, PC	Quantifies the probability that two species randomly placed across a patchy landscape (of total area A_L) fall into habitat patches a_i and a_j that are reachable from each other with a maximum connectivity probability p_{ij}, defined as the maximum product probability of all possible paths between patches i and j (including single-step paths).	$$PC(v) = \frac{\sum_{i=1}^{n} \sum_{j=1}^{n} a_i a_j p_{ij}^*}{A_L^2}$$	Landscape networks	Saura & Pascual-Hortal 2007

Figure 1. Summary of our literature review, detailing the number of studies by (a) edge weights, and

(b) whether the information provided was sufficient or insufficient, and network type (as "landscape",

"interaction", and "others", which include social, co-occurrence, etc. networks). We only categorize as

"correct" or as "wrong" studies on which we had enough information to support such a claim, and as

"seems correct" and "seems wrong" studies on which the insufficient information available suggest that

calculations are correct or wrong, respectively. Notice that (1) most of the correct or probably correct

studies use unweighted edges (n = 52, 66%); (1) for all network types half of the studies do not report

enough information to validate whether calculations were correct (n = 63, 49%); and that (3) numerous

studies report unclear calculations (n = 19). Differences in the number of oversights in centrality

calculations between weighted and landscape and interaction networks are probably due to the fact

that in interaction networks weighted edges typically require transformation (see Table 1), whereas in

landscape networks edge weights tend to be inversely proportional, requiring no transformation.

Figure 2. Scheme illustrating the step-by-step decision process for the calculation of shortest path-

centrality measures in ecological networks and the 5-point guide of information require for good-

practices.

Figure 3. Toy networks depicting two possible paths to arrive from the city of Busan to the city of

Almería. Edges connecting the different cities quantify distance in Km (a), or using different

measurements of the movement of researchers between these cities (b-d; see text for explanation).

Figure 4. Land topography and ocean bathymetry (m) of the Italian region (data from the ETOPO1

Global Relief database; doi:10.7289/V5C8276M). A hypothetical bird species lives between 500 and

2000 m a.s.l., with a typical habitat size of about 15 km^2 and a dispersal distance of 100km. a) Pixels

in lighter tones denote patches of suitable habitat (colored in black in the following panels). After

computing Betweenness (BC) and Closeness centrality (CC), we highlighted in red the 20 pixels with

highest BC in the binary (b) and weighted network (c), and the 20 pixels with highest CC in the binary

(d) and weighted network (e). Note the differences in the identification of the pixels between weighted

and binary versions of the data.

Figure 5. Toy network describing the social affinity between four famous characters. Edges quantify

proportion of shared parasites (as in Gómez, Nunn & Verdú, 2003).

Figure 6. Scatter plot showing species ranks based on Betweenness (a) and Closeness centrality (b)

values calculated on untransformed and inversed edge weights.

Figure 7. Toy network describing the carbon flow through a food chain in a famous British moat.

Supporting Information

Supporting Information 1. Details and references of the literature review.

Supporting Information 2. Review of other potential pitfalls on shortest-path centrality calculations and additional figures.

Figure 1.

a)

b)

Legend:
- Correct (dark green)
- Seem correct (light green)
- Unclear (gray)
- Seem wrong (salmon/red)

Figure 2.

Figure 3.

Figure 4.

Figure 5.

a)

b)

Figure 6.

a) Betweenness Centrality

b) Closeness Centrality

Untransformed edge weight (a_{ij})

Transformed edge weight ($1/a_{ij}$)

Figure 7.