Unveiling the Complexity of Food Webs: A Comprehensive Overview of Definitions, Scales, and Mechanisms

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Abstract

Food webs are a useful abstraction and representation of the feeding links between species in a community and are used to infer many ecosystem level processes. However, the different theories, mechanisms, and criteria that underpin how a food web is defined and, ultimately, constructed means that not all food webs are representing the same ecological process. Here we present a synthesis of the different assumptions, scales and mechanisms that are used to define different ecological networks ranging from metawebs (an inventory of all potential interactions) to fully realised networks (interactions that occur within a given community over a certain timescale). Illuminating the assumptions, scales, and mechanisms of network inference allows a formal categorisation of how to use networks to answer key ecological and conservation questions and defines guidelines to prevent unintentional misuse or misinterpretation.

At the heart of modern biodiversity science are a set of concepts and theories about biodiversity, stability and function. These relate to the abundance, distribution and services that biodiversity provides, and how biodiversity – as an interconnected set of species – responds to multiple stressors. The interaction between species is one of the fundamental building blocks of ecological communities, providing a powerful abstraction that can help quantify, conceptualise, and understand biodiversity dynamics, and ultimately, make predictions, mitigate change, and manage services [1]. Such network representations of biodiversity (including within species diversity) are increasingly argued to be an asset to predictive ecology, climate change mitigation and resource management, with the argument that characterising biodiversity in a network will afford a deeper capacity to understand and predict the abundance, distribution, dynamics and services provided by multiple species facing multiple stressors. However, there is a growing discourse around limitations to the interpretation and applied use of networks [2,3], primarily as the result of shortcomings regarding the conceptualisation of networks [4].

A ‘network’ can be defined and conceptualised in a myriad of ways, which means that different networks will be embedding different processes (or determinants) of interactions, ultimately influencing the patterns and mechanisms that are inferred [5,6]. The different ways in which a network can be represented is the result of *how* the network is constructed, which itself rests on two pillars: the data used to construct the network and the underlying theory as to what drives the interactions between species. The latter represents an expression of mechanism and process that gives rise to the patterns that emerge from collating interactions among species, and will ultimately inform which data are deemed important in the determination of interactions occurring. Each of these pillars carries with it a set of practical, semantic and conceptual constraints that not only influence progress in making network ecology more valuable and potentially predictive, but help define the spatial, temporal, and evolutionary scale of assumptions we make and the predictions we might generate from different network representations.

In this perspective we aim to provide an overview of the different **food web** representations (*a note on how there has been developments in the ‘bipartite space’ and it would be flawed to try and view them in tandem as food webs and non-trophic webs are two very different conceptualisations*), particularly how these relate to the terminology used to define a network, and how this influenced by both the processes that determine networks as well as how this relates to the way in which we construct networks. The provision of this detail ultimately leads to a set of insights and conclusions about whether, when. and under what conditions network representations of biodiversity can contribute to the advancement of ecological theory and generate value in predictive ecology. Specifically, we finish this perspective with an overview of fundamental questions in ecology that we think can benefit from network thinking and a proposal that such thinking can accelerate our capacity to predict the impact of multiple stressors on biodiverse communities.

# 1. Setting the Scene: The Not So Basics of Nodes and Edges

Defining a food web seems simple; it is the representation of the interactions (edges) between species (nodes) in the form of a network, however the definition of ‘edges’ and ‘nodes’, as well as the levels of organization at which they are aggregated can take many forms [8], which ultimately encodes a series of assumptions and criteria within a network. An awareness of variance in the way a food web can be defined is critical as a network (or its adjacency matrix) is both the ‘object’ from which inferences are made (*e.g.,* the interactions between species, or how the structure influences ecosystem level processes) as well as the ‘product’ of either the data collection [6] or prediction process [9]. One thus needs to be aware of both the criteria that is used to define nodes and edges, and what processes or mechanisms the aggregation of the two represents, as this will determine what the network can be used for.

## 1.1 How do we define a node?

Although this may seem an elementary question in the context of food webs — a node *should* represent a (taxonomic) species, the reality is that nodes can often represent an aggregation of different species - so called ‘trophic species’ [10,11] or segregation of species by life stages [12]. Practical implications of how we are aggregating the nodes is that the resolution may not always be ‘pixel perfect’, which limits the ability to make (taxonomic) species specific inferences *e.g.,* does species eat species , however there is value in having nodes that represent an aggregation of species, as the distribution of the links between them are more meaningful in terms of understanding energy flow and distribution within the system.

## 1.2 What is captured by an edge?

At its core, links within food webs can be thought of as a representation of either feeding links between species - be that realised [13] or potential [14], alternative links can represent fluxes within the system *e.g.,* energy transfer or material flow as the result of the feeding links between species [15]. Fundamentally this means that the links within a network represent different ‘currencies’ (either the feasibility of a link existing between two species or the energy that is moving through the system) and how the links within a network are specified will influence the resulting structure of the network. For example taking a food web that consists of links representing all *potential* feeding links for a community (*i.e.,* a metaweb) will be meaningless if one is interested in understanding the flow of energy through the network as the links within a metaweb do not represent environmental/energetic constraints, making them poor representations of which interactions are *realised* in a specific location [16]. In addition to the various ways of defining the links between species pairs there are also a myriad of ways in which the links themselves can be quantified. Links between species are often treated as being present or absent (*i.e.,* binary) but it is also possible to use probabilities [9,which quantifies how likely an interaction is to occur, 17] or continuous measurements [which quantifies the strength of of an interaction, 18].

## 1.3 Network representations

Broadly, networks can be thought of to fall into two different ‘types’; namely metawebs; traditionally defined as all of the *potential* interactions for a specific species pool [14], and realised networks; which is the subset of interactions in a metaweb that are *realised* for a specific community at a given time and place. The fundamental difference between these two different network representations is that a metaweb provides insight as to the viability of an interaction between two species occurring and is a means to identify links that are not ecologically plausible, *i.e.,* forbidden links [19], or provide an idea of the *complete* diet of a species [20]. In contrast realised networks are highly localised and the links between species are contingent on both the co-occurrence of species, the role of the environment, and population and community dynamics on predator choice. In the context of definitions and semantics the links that are represented by a metaweb and a realised network are different; links that are absent in a metaweb can be treated as being truly absent, however links that are absent in a realised network cannot be considered to be truly absent but rather as absent due to the broader environmental/community context. Importantly, a realised network is *not* simply the downscaling of a metaweb to a smaller scale (*e.g.,* moving from the country to the 1x1 km2 scale based on fine-scale species co-occurrence) but represents a shift towards capturing the higher level processes that determine the *realisation* of an interaction, *i.e.,* the definition of an edges shifts from being determined by interaction feasibility to that of energetic choices/consequences. Thus, different network representations are determined and constrained by different sets of assumptions as to what the processes are that determine the presence/absence of an interaction between two species as well as the resulting network structure.

# 2. From Nodes and Edges to Scale, Context, and Process

The interplay between network representation and network (node and edge) definition is primarily governed by the process(es) that determine the interaction between species, however these processes are also scale and context dependent. Here we start by introducing the five core processes that determine either the feasibility or the realisation of interactions, namely: evolutionary compatibility, co-occurrence, abundance, predator choice, and non-trophic interactions; while simultaneously contextualising them within, and linking them to, the different network representations [Figure 1](#fig-process). We can think of the different network representations to be conceptually analogous to the fundamental and realised niche, whereby the metaweb represents the ‘fundamental diet niche’ of a species and a realised network represents the ‘realised diet’ of a species. Of course these processes do not function in a vacuum and do interact with/influence one another, but it is still beneficial to present them in a categorical manner as these different processes are often the underpinning logic in the development of prediction/network models, the criteria for data collection in the field, and the scale of organisation for which they are relevant (species, population, or community).

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| Figure 1: Aligning the various processes that determine interactions with the different network representations. First we start with a ‘global metaweb’ this network which captures all possible interactions for an arbitrary collection of species, we can further refine this network by taking in to consideration the co-occurrence of these difference species - as shown here we have two regions with some species (blue) that are found in both regions and others endemic to either region one (pink) or region two (orange). These regional metawebs to capture all possible interactions, however it only considers species that co-occur. However even within a region we do not expect all interactions to be realised but rather that there are multiple configurations of the regional metaweb over both space and time. The ‘state’ of the different network realisations are ultimately influenced not just by the co-occurrence of a species pair but rather the larger community context such as the abundance of different species, maximising energy gain, or indirect/higher order interactions. |

## 2.1 The processes that determine species interactions

**Evolutionary compatibility**

There is compelling evidence that an interaction occurring between two species is the result of their shared (co)evolutionary history [21–23] which, in the more proximal sense, is manifested as the ‘trait complementarity’ between two species [24], whereby one species (the predator) has the ‘correct’ set of traits that allow it to chase, capture, kill, and consume the other species (the prey). For species pairs where this condition is not met the link is deemed to be forbidden [19]; *i.e.,* not physically possible and will always be absent within a network. A network constructed on the basis of evolutionary compatible links is most closely aligned with a metaweb, although it would not be required that the species co-occur (as shown in [Figure 1](#fig-process)), and arguably makes for a good approximation of the ‘Eltonian niche’ of species [25]. Finally, one should be aware that it is possible to represent evolutionary compatible interactions as either binary (possible vs forbidden) or as a probability [9], where the probability represents how likely the interaction between two species is to be possible.

**(Co)occurrence**

Although the outright assumption that because two species are co-occurring it must mean that they are interacting is flawed [26], it is of course impossible for two species to interact (at least in terms of feeding links) if they are not co-occurring in time and space. Thus, although co-occurrence data alone is insufficient to build an accurate and ecologically meaningful representation of *feeding links* it is still a critical process that determines the realisation of feeding links and allows us to constrain a global metaweb to only consider ‘realised’ communities [27] and an understanding of the intersection of species interactions and their co-occurrence is meaningful when one is operating in the space of trying to determine the distribution of a species [28,29], representing something of a fusion of the the Grinnellian and Eltonian niches [30].

**Abundance**

The abundance of different the species within the community is thought to influence the realisation of feeding links primarily in two ways. Firstly there is the argument that that structure of networks (and the interactions that they are composed of) are driven *only* by the abundance of the different species and that interactions are not contingent on there being any compatibility (trait matching) between them, *sensu* neutral processes [31,32]. However, a more ecologically sound assumption would be that the abundance of different prey species will influence the distribution of links in a network [33], be influencing which prey are targeted or preferred by the predator as abundance influences factors such as the likelihood of two species (individuals) meeting [9,34], or in the dynamic sense will influence the persistence of viable populations.

**Profitability (predator choice)**

Ultimately, predator choice is underpinned by the energetic cost-benefit (profitability) of trying to catch, kill, and consume prey (where a predator will optimise energy while minimising handling and search time), and is well described within both optimal foraging [35] and metabolic theory [36]. The energetic cost of feeding is itself can be deconstructed as the energy content as well as the density (abundance) of prey (as this influences search time) and how these will influence which links are realised [Figure 1](#fig-process), with an argument that body size represents a key trait that may capture and influence these processes [37,38]. Additional work on on understanding the energetic cost that the environment imposes on an individual [39] as well as the way a predator uses the landscape to search for prey [40] is bringing us closer to accounting for the energetic cost of realising feeding links.

**Non-trophic interactions**

Perhaps not as intuitive when thinking about the processes that determine feeding links (trophic interactions) is thinking about the role of the ability of non-trophic interactions to modify either the realisation or strength of trophic interactions [41,42]. Non-trophic interactions can modify interactions either ‘directly’ *e.g.,* predator *a* outcompetes predator *b* or ‘indirectly’ *e.g.,* mutualistic/facilitative interactions will alter the fine-scale distribution and abundance of species as well as their persistence [43–45]. The ‘unobservable’ nature of non-trophic interactions makes them a challenge to quantify, however their importance in network dynamics [46] as well as cascading effects [*e.g.,* 47] should not be overlooked.

## 2.2 Contextualising the processes that determine species interactions

It should be self evident that the different processes discussed above are all ultimately going to influence the realisation of interactions as well as the structure of a network, however they are acting at different scales of organisation. Both the **co-occurrence** and the **evolutionary compatibility** are valid at the scale of the species pair of interest, that is the *possibility* of an interaction being present/absent is assessed at the pairwise level and one is left with a ‘list’ of interactions that are present/absent. Although it is possible to build a network (*i.e.,* metaweb) from this information it is important to be aware that the structure of this network is not constrained by real-world dynamics or conditions, and so just because species are able to interact does not mean that they will [34]. In order to construct a network who’s structure is a closer approximation of reality (localised interactions) one needs to take into consideration the properties of the community as a whole and information about the individuals it is comprised of [48], which requires more data at the community scale, such as the abundance of species.

# 3. Network construction is nuanced

The act of constructing a ‘real world’ network will ultimately be delimited by its intended use, however the reality is that the empirical collection of interaction data is both costly and challenging to execute [19,49], especially if one wants to capture *all* aspects of the processes discussed in [Section 2](#sec-process) (owing to the different time and spatial scales they may be operating at). Thus we often turn to models to either predict networks (be that the interaction between two species, or network structure [50]), or as a means to identify missing interactions (gap fill) within an existing empirical dataset [51–53], and so for the purpose of this discussion network construction will be synonymous with using a model as a means to represent or predict a network. That is not to say that there is no need for empirical data collection but rather that using a model for food web prediction (or reconstruction) is a more feasible approach as it allows us to make inferences about interactions that are not happening in the ‘observable now’ [50], with the added benefit that one is able to build some uncertainty into the resulting network [9]. Additionally different models have different underlying philosophies that allow us to capture one or a few of the processes discussed in [Section 2](#sec-process), and although the delimits and defines what inferences can be made from the resulting network it also allows us to isolate and understand how different processes determine interactions [54,55]. Here we will introduce the three different types of network representations (metawebs, realised networks, and structural networks), how they link back to (and encode) the different processes determining interactions [Figure 1](#fig-process), and broadly discuss some of the modelling approaches that are used to construct these different network types. This is paralleled by a hypothetical case study (Box 1) where we showcase the utility/applicability of the different network representations in the context of trying to understand the feeding dynamics of a seasonal community.

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| Box 1 - Why we need to aggregate networks at different scales: A hypothetical case study |
| note I am using a figure for layout experimentation purposes  Although it might seem most prudent to be predicting, constructing, and defining networks that are the closest representation of reality there are pros and cons of constructing both realised networks as well as metawebs. Let us take for example a community that experiences a degree of species turnover between seasons. In this community we expect species to be either present or absent depending on the season (*i.e.,* changes in co-occurrence) as well as some species exhibiting seasonal shifts in their diets (be that due to changes in species occurrence or predator choice). If one were to construct a metaweb that disregards these season shifts (‘global metaweb’) these details would be lost and it would be valuable to construct either smaller metawebs for the different seasonal communities (thereby capturing the changes in community diversity), or realised networks for each season (to capture diet or ecosystem process shifts *e.g.,* [56]). However, these small-scale networks lack the context of the bigger picture that is available at the metaweb - that is it gives us a more holistic idea of the entire diet range of a specific species, which is important when one needs to make conservation-based/applied decisions (*e.g.,* conserving the entire diet of a species and not just seasonal prey items) as well as providing information on interactions that may be possible regardless of the environmental/community context (species may have the capacity to consume certain prey items but do not do so due to local conditions). With this is mind let us see how the different network aggregations can be used   |  | | --- | | Figure 2 | |

## 3.1 Models that predict metawebs (feasible interactions)

This is perhaps the most developed group of models; with a variety of approaches having been developed that typically determine the feasibility of an interaction using the trait compatibility between predator and prey (*i.e.* their evolutionary compatibility) to determine ‘feeding rules’ [57]. These feeding rules are broadly elucidated in two different ways; mechanistic feeding rules can be explicitly defined and applied to a community [*e.g.,* 58,59,60] or they are inferred from a community for which there are interaction data and the ‘rules’ are then applied to a different community [*e.g.,* 20,61–67]. The fundamental difference between these two model groups is that ‘mechanistic models’ rely on expert knowledge and make explicit assumptions on trait-feeding relationships, whereas the ‘pattern finding’ models are dependent on existing datasets from which to elucidate feeding rules. These models are useful for determining all feasible interactions for a specific community, and owing to the availability of empirical interaction datasets [*e.g.,* 68,69,70], as well as the development of model testing/benchmarking tools [71], means that these models can be validated and (with relative confidence) be used to construct first draft networks for communities for which we have no interaction data [62], and are valuable not only in data poor regions but also for predicting interactions for ‘unobservable’ communities *e.g.,* prehistoric networks [72,73] or future, novel community assemblages. Importantly metawebs are inherently ‘static’ in the sense that they are *not* able to capture dynamic processes (since the notion of feasibility is all or nothing), however they provide a bigger picture context (*e.g.,* understanding the *entire* diet breadth of a species) and often require little data to construct.

## 3.2 Models that predict realised networks (realised interactions)

In order to construct realised networks models need to incorporate *both* the feasibility of interactions (*i.e.,* determine the entire diet breadth of a species) as well as then determine which interactions are realised (*i.e.,* incorporate the ‘cost’ of interactions). As far as we are aware there is no model that explicitly accounts for both of these ‘rules’ (although see [74]) and rather *only* account for processes that determine the realisation of an interaction (*i.e.,* abundance, predator choice, or non-trophic interactions). Although the use of allometry *i.e.,* body size [*e.g.,* 75,76] may represent a first step in capturing ‘evolutionary compatibility’ alongside more energy (predator choice) driven processes we still need to account for other traits that determine feeding compatibility [*e.g.,* 77 show how incorporating prey defensive properties alongside body size improves predictions]. In terms of constructing realised networks, diet models [76,78] have been used construct networks based on both predator choice (as determined by the handling time, energy content, and predator attack rate) as well as abundance (prey density) and progress has also been made in understanding the compartmentation of energy in networks and how this influences energy acquisition [79,80]. As realised networks are are build on the concept of dynamic processes (the abundance of species will always be in flux) these networks are valuable for understanding the behaviour of networks over time or their response to change [81–83]. However, they are ‘costly’ to construct (requiring data about the entire community as it is the behaviour of the system that determines the behaviour of the part) and also lack the larger diet niche context afforded by metawebs.

## 3.3 Models that predict structure (interaction agnostic)

Although we identify mechanisms that determine species interactions in [Section 2](#sec-process) not all models that are used to predict networks explicitly operate at the ‘process’ level, but rather represent the *structure* of a network based on a series of *a priori* assumptions as to the distribution of links between species (typically trophic not taxonomic species). These models operate by parametrising an aspect of the network structure, (*e.g.,* the niche model [11] makes an assumption as to the expected connectance of the network,although see [84] for a parameter-free model) or alternatively uses structural features of an exiting *realised* network (*e.g.,* stochastic block model, [85]). Importantly these structural models do not make species specific predictions (they are usually species agnostic and treat nodes as trophic species) and so cannot be used to determine if an interaction is either possible *or* realised between two species (*i.e.,* one cannot use these models to determine if species eats species ). Although this means this suite of models are unsuitable as tools for predicting species-specific interactions, they have been shown to be sufficient tools to predict the structure of networks [86], and provide a data-light (the models often only require species richness) but assumption heavy (the resulting network structure is determined by an assumption of network structure) way to construct a network.

# 4. Making Progress with Networks

## 4.1 Further development of models and tools

There has been a suite of models that have been developed to predict feeding links, however we are lacking in tools that are explicitly taking into consideration estimating both the feasibility as well as realisation of links, *i.e.,* both interactions and structure simultaneously [50]. This could be addressed either through the development of tools that do both (predict both interactions and structure), or to develop an ensemble modelling approach [87,88] or tools that will allow for the downsampling of metawebs into realised networks [*e.g.,* 89]. Additionally although realised networks are more closely aligned with capturing interaction strength we lack models that allow us to quantify this [50,90]. In addition to the more intentional development of models we also need to consider the validation of these models, there have been developments and discussions for assessing how well a model recovers pairwise interactions [50,71], although the rate of false-negatives that may be present in the testing data still present a challenge [91], and we still lack clear set of guidelines for benchmarking the ability of models to recover structure [92].

## 4.2 At what scale should we be predicting and using networks?

We lack an understanding of which processes drive interactions at different scales [93], as well as to what the appropriate level of aggregation for a ‘network’ is [8,94]. Thus we need an understanding of not only how time and scale influence the interpretation of networks [95,96], but how this is in turn influenced by the type of networks used. Which presents a challenge both in deciding what the appropriate spatial and time scales are for constructing not only a network but also which type of network representation. Space influences both network properties [97], as well as dynamics [98,99], and time has implications when it comes to accounting for seasonal turnover in communities [100,101] as well as thinking about co-occurrence, particularly the records that are used to determine co-occurence [102]. Although multilayer networks may allow us to encode the nuances of space and time [103] we still need to understand the implications of *e.g.,* constructing networks that are not at ecologically but rather politically relevant scales [62] and what the implications of this disconnect may be.

# 5. The future value of networks

developing a dictionary of use… that helps navigate between the levels and assumptions

It should be clear that there is a high degree of interrelatedness and overlap between the way a network is constructed (modelled or predicted) and the process(es) it captures, these are encoded (embedded) within the network representation and ultimately influences how the network can and should be used [104,105], with different network representations yielding different interpretations of processes [106]. It is probably both this nuance as well as a lack of clear boundaries and guidelines as to the links between network form and function [although see 107] that has stifled the ‘productive use’ of networks beyond inventorying the interactions between species. Although, progress with using networks as a means to address questions within larger bodies of ecological theory *e.g.,* invasion biology [108] and co-existence theory [109], has been made we still need to have a discussion on what the appropriate network representation for the task at hand would be. This is highlighted in Box 1, and underscores that we need to evaluate exactly what process a specific network representation captures as well as its suitability for the question of interest.

## 5.1 How will novel communities interact?

Here we can talk about the effects of range shifts and invasions and how this will result in new/novel community assemblages. And then also the intentional changes of species compositions through rewilding.

## 5.2 How will changes in the community influence ecosystem processes?

Linking to dynamic networks and how this lets us build spatially/temporally explicit networks which can be used to infer form and function. Also bring in the discussion on the suitable aggregation (and the fact that we don’t know)

## 5.3 How do species persist/co-exist?

Specific sub points to consider here is persistence, especially persistence to perturbation. Again, dynamic networks and network/community assembly and finally extinctions [110].

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| Figure 3: An attempt to try and visualise a way to map the different scales of network representations to the way in which we can interrogate/ask questions about them? |

An informative table

| Question (broad) | Question (specific) | Network representation |
| --- | --- | --- |
| Species invasions | What species will the invading species interact with? | Regional metaweb but need to derive information from a global metaweb since these are interactions that are ‘novel’ |
| Species invasions | How does the invading species alter network dynamics and function? | Realised network (after having moved through the global metaweb to understand which interactions are feasible) |
| Range shifts and novel communities | Under global change how will novel community assemblages interact? | Global metaweb, need context of broader community |
| Extinctions | Cascading effect of the loss of a species from the network | Regional metaweb - need to account for entire diet, a realised network will exclude the entire diet but will allow to elucidate the final structure |
| Species/community persistence | Dynamics over time. Stability/resilience. How does a change in pop *A* affect pop *B*? | Realised networks - but dynamic! |
| Synthetic networks | Creating ecologically plausible communities for synthetic analyses | Structural networks - data light! |
| Practical use | What is both attainable (data constraints) but also of practical use to ‘real world’ decision making. So moving from theory to applied | ??Regional metawebs?? |

# References

1. Windsor, F.M. *et al.* (2023) [Using ecological networks to answer questions in global biogeography and ecology](https://doi.org/10.1111/jbi.14447). *Journal of Biogeography* 50, 57–69

2. Dormann, C.F. (2023) The rise, and possible fall, of network ecology. In *Defining Agroecology – A Festschrift for Teja Tscharntke*, pp. 143–159., Tredition

3. Blüthgen, N. (2010) [Why network analysis is often disconnected from community ecology: A critique and an ecologist’s guide](https://doi.org/10.1016/j.baae.2010.01.001). *Basic and Applied Ecology* 11, 185–195

4. Blüthgen, N. and Staab, M. (2024) [A Critical Evaluation of Network Approaches for Studying Species Interactions](https://doi.org/10.1146/annurev-ecolsys-102722-021904). *Annual Review of Ecology, Evolution, and Systematics* 55, 65–88

5. Proulx, S.R. *et al.* (2005) [Network thinking in ecology and evolution](https://doi.org/10.1016/j.tree.2005.04.004). *Trends in Ecology & Evolution* 20, 345–353

6. Brimacombe, C. *et al.* (2023) [Shortcomings of reusing species interaction networks created by different sets of researchers](https://doi.org/10.1371/journal.pbio.3002068). *PLOS Biology* 21, e3002068

7. Poisot, T. *et al.* (2016) [Describe, understand and predict: Why do we need networks in ecology?](https://www.jstor.org/stable/48582345) *Functional Ecology* 30, 1878–1882

8. Moulatlet, G. *et al.* (2024) [The scaling of trophic specialization in interaction networks across levels of organization](https://doi.org/10.22541/au.172977303.33335171/v1)Authorea

9. Banville, F. *et al.* (2024) [Deciphering probabilistic species interaction networks](https://doi.org/10.32942/X28G8Z)EcoEvoRxiv

10. Yodzis, P. (1982) [The Compartmentation of Real and Assembled Ecosystems](https://doi.org/10.1086/284013). *The American Naturalist* 120, 551–570

11. Williams, R.J. and Martinez, N.D. (2000) [Simple rules yield complex food webs](https://doi.org/10.1038/35004572). *Nature* 404, 180–183

12. Clegg, T. *et al.* (2018) [The impact of intraspecific variation on food web structure](https://doi.org/10.1002/ecy.2523). *Ecology* 99, 2712–2720

13. Pringle, R.M. (2020) [Untangling Food Webs](https://doi.org/10.1515/9780691195322-020). In *Unsolved Problems in Ecology*, pp. 225–238, Princeton University Press

14. Dunne, J.A. (2006) The Network Structure of Food Webs. In *Ecological networks: Linking structure and dynamics* (Dunne, J. A. and Pascual, M., eds), pp. 27–86, Oxford University Press

15. Lindeman, R.L. (1942) [The Trophic-Dynamic Aspect of Ecology](https://doi.org/10.2307/1930126). *Ecology* 23, 399–417

16. Caron, D. *et al.* (2024) [Trait-matching models predict pairwise interactions across regions, not food web properties](https://doi.org/10.1111/geb.13807). *Global Ecology and Biogeography* 33, e13807

17. Poisot, T. *et al.* (2016) [The structure of probabilistic networks](https://doi.org/10). *Methods in Ecology and Evolution* 7, 303–312

18. Berlow, E.L. *et al.* (2004) [Interaction strengths in food webs: Issues and opportunities](https://doi.org/10.1111/j.0021-8790.2004.00833.x). *Journal of Animal Ecology* 73, 585–598

19. Jordano, P. (2016) Sampling networks of ecological interactions. *Functional Ecology* DOI: [10.1111/1365-2435.12763](https://doi.org/10.1111/1365-2435.12763)

20. Strydom, T. *et al.* (2023) [Graph embedding and transfer learning can help predict potential species interaction networks despite data limitations](https://doi.org/10.1111/2041-210X.14228). *Methods in Ecology and Evolution* 14, 2917–2930

21. Segar, S.T. *et al.* (2020) [The Role of Evolution in Shaping Ecological Networks](https://doi.org/10.1016/j.tree.2020.01.004). *Trends in Ecology & Evolution* 35, 454–466

22. Gómez, J.M. *et al.* (2010) [Ecological interactions are evolutionarily conserved across the entire tree of life](https://doi.org/10.1038/nature09113). *Nature* 465, 918–921

23. Dalla Riva, G.V. and Stouffer, D.B. (2016) [Exploring the evolutionary signature of food webs’ backbones using functional traits](https://doi.org/10.1111/oik.02305). *Oikos* 125, 446–456

24. Benadi, G. *et al.* (2022) [Quantitative Prediction of Interactions in Bipartite Networks Based on Traits, Abundances, and Phylogeny](https://doi.org/10.1086/714420). *The American Naturalist* 199, 841–854

25. Soberón, J. (2007) [Grinnellian and Eltonian niches and geographic distributions of species](https://doi.org/10.1111/j.1461-0248.2007.01107.x). *Ecology Letters* 10, 1115–1123

26. Blanchet, F.G. *et al.* (2020) [Co-occurrence is not evidence of ecological interactions](https://doi.org/10.1111/ele.13525). *Ecology Letters* 23, 1050–1063

27. Dansereau, G. *et al.* (2024) [Spatially explicit predictions of food web structure from regional-level data](https://doi.org/10.1098/rstb.2023.0166). *Philosophical Transactions of the Royal Society B: Biological Sciences* 379

28. Higino, G.T. *et al.* (2023) [Mismatch between IUCN range maps and species interactions data illustrated using the Serengeti food web](https://doi.org/10.7717/peerj.14620). *PeerJ* 11, e14620

29. Pollock, L.J. *et al.* (2014) [Understanding co-occurrence by modelling species simultaneously with a Joint Species Distribution Model (JSDM)](https://doi.org/10.1111/2041-210X.12180). *Methods in Ecology and Evolution* 5, 397–406

30. Gravel, D. *et al.* (2019) [Bringing Elton and Grinnell together: A quantitative framework to represent the biogeography of ecological interaction networks](https://doi.org/10.1111/ecog.04006). *Ecography* 42, 401–415

31. Canard, E. *et al.* (2012) [Emergence of Structural Patterns in Neutral Trophic Networks](https://doi.org/10.1371/journal.pone.0038295). *PLOS ONE* 7, e38295

32. Momal, R. *et al.* (2020) [Tree-based inference of species interaction networks from abundance data](https://doi.org/10.1111/2041-210X.13380). *Methods in Ecology and Evolution* 11, 621–632

33. Vázquez, D.P. *et al.* (2009) [Uniting pattern and process in plant–animal mutualistic networks: A review](https://doi.org/10.1093/aob/mcp057). *Annals of Botany* 103, 1445–1457

34. Poisot, T. *et al.* (2015) [Beyond species: Why ecological interaction networks vary through space and time](https://doi.org/10.1111/oik.01719). *Oikos* 124, 243–251

35. Pyke, G. (1984) [Optimal Foraging Theory: A Critical Review](https://doi.org/10.1146/annurev.ecolsys.15.1.523). *Annual Review of Ecology, Evolution and Systematic* 15, 523–575

36. Brown, J.H. *et al.* (2004) [Toward a Metabolic Theory of Ecology](https://doi.org/10.1890/03-9000). *Ecology* 85, 1771–1789

37. Yodzis, P. and Innes, S. (1992) [Body Size and Consumer-Resource Dynamics](https://doi.org/10.1086/285380). *The American Naturalist* 139, 1151–1175

38. White, E.P. *et al.* (2007) [Relationships between body size and abundance in ecology](https://doi.org/10.1016/j.tree.2007.03.007). *Trends in Ecology & Evolution* 22, 323–330

39. Cherif, M. *et al.* (2024) [The environment to the rescue: Can physics help predict predator–prey interactions?](https://doi.org/10.1111/brv.13105) *Biological Reviews* n/a

40. Pawar, S. *et al.* (2012) [Dimensionality of consumer search space drives trophic interaction strengths](https://doi.org/10.1038/nature11131). *Nature* 486, 485–489

41. Golubski, A.J. and Abrams, P.A. (2011) [Modifying modifiers: What happens when interspecific interactions interact?](https://doi.org/10.1111/j.1365-2656.2011.01852.x) *Journal of Animal Ecology* 80, 1097–1108

42. Pilosof, S. *et al.* (2017) [The multilayer nature of ecological networks](https://doi.org/10.1038/s41559-017-0101). *Nature Ecology & Evolution* 1, 101

43. Kéfi, S. *et al.* (2012) [More than a meal… integrating non-feeding interactions into food webs: More than a meal …](https://doi.org/10.1111/j.1461-0248.2011.01732.x). *Ecology Letters* 15, 291–300

44. Kéfi, S. *et al.* (2015) [Network structure beyond food webs: Mapping non-trophic and trophic interactions on Chilean rocky shores](https://doi.org/10.1890/13-1424.1). *Ecology* 96, 291–303

45. Buche, L. *et al.* (2024) [Multitrophic Higher-Order Interactions Modulate Species Persistence](https://doi.org/10.1086/729222). *The American Naturalist* 203, 458–472

46. Staniczenko, P.P.A. *et al.* (2010) [Structural dynamics and robustness of food webs](https://doi.org/10.1111/j.1461-0248.2010.01485.x). *Ecology Letters* 13, 891–899

47. Kamaru, D.N. *et al.* (2024) [Disruption of an ant-plant mutualism shapes interactions between lions and their primary prey](https://doi.org/10.1126/science.adg1464). *Science* 383, 433–438

48. Quintero, E. *et al.* (2024) [Downscaling mutualistic networks from species to individuals reveals consistent interaction niches and roles within plant populations](https://doi.org/10.1101/2024.02.02.578595)bioRxiv, 2024.02.02.578595

49. Jordano, P. (2016) [Chasing Ecological Interactions](https://doi.org/10.1371/journal.pbio.1002559). *PLOS Biology* 14, e1002559

50. Strydom, T. *et al.* (2021) [A roadmap towards predicting species interaction networks (across space and time)](https://doi.org/10.1098/rstb.2021.0063). *Philosophical Transactions of the Royal Society B: Biological Sciences* 376, 20210063

51. Biton, B. *et al.* (2024) Inductive link prediction boosts data availability and enables cross-community link prediction in ecological networks

52. Stock, M. (2021) Pairwise learning for predicting pollination interactions based on traits and phylogeny. *Ecological Modelling*

53. Dallas, T. *et al.* (2017) [Predicting cryptic links in host-parasite networks](https://doi.org/10.1371/journal.pcbi.1005557). *PLOS Computational Biology* 13, e1005557

54. Stouffer, D.B. (2019) [All ecological models are wrong, but some are useful](https://doi.org/10.1111/1365-2656.12949). *Journal of Animal Ecology* 88, 192–195

55. Song, C. and Levine, J.M. (2024) [Rigorous (in)validation of ecological models](https://doi.org/10.1101/2024.09.19.613075)bioRxiv, 2024.09.19.613075

56. Schwarz, B. *et al.* (2020) [Temporal scale-dependence of plant–pollinator networks](https://doi.org/10.1111/oik.07303). *Oikos* 129, 1289–1302

57. Morales-Castilla, I. *et al.* (2015) [Inferring biotic interactions from proxies](https://doi.org/10.1016/j.tree.2015.03.014). *Trends in Ecology & Evolution* 30, 347–356

58. Shaw, J.O. *et al.* (2024) [A framework for reconstructing ancient food webs using functional trait data](https://doi.org/10.1101/2024.01.30.578036)bioRxiv, 2024.01.30.578036

59. Dunne, J.A. *et al.* (2008) [Compilation and Network Analyses of Cambrian Food Webs](https://doi.org/10.1371/journal.pbio.0060102). *PLOS Biology* 6, e102

60. Roopnarine, P.D. (2017) Ecological Modelling of Paleocommunity Food Webs. In *Conservation Paleobiology: Using the Past to Manage for the Future*, pp. 201–226, University of Chicago Press

61. Pichler, M. *et al.* (2020) [Machine learning algorithms to infer trait-matching and predict species interactions in ecological networks](https://doi.org/10.1111/2041-210X.13329). *Methods in Ecology and Evolution* 11, 281–293

62. Strydom, T. *et al.* (2022) [Food web reconstruction through phylogenetic transfer of low-rank network representation](https://doi.org/10.1111/2041-210X.13835). *Methods in Ecology and Evolution* 13, 2838–2849

63. Caron, D. *et al.* (2022) [Addressing the Eltonian shortfall with trait-based interaction models](https://doi.org/10.1111/ele.13966). *Ecology Letters* 25, 889–899

64. Llewelyn, J. *et al.* (2023) [Predicting predator–prey interactions in terrestrial endotherms using random forest](https://doi.org/10.1111/ecog.06619). *Ecography* 2023, e06619

65. Desjardins-Proulx, P. *et al.* (2017) [Ecological interactions and the Netflix problem](https://doi.org/10.7717/peerj.3644). *PeerJ* 5, e3644

66. Eklöf, A. *et al.* (2013) [Secondary extinctions in food webs: A Bayesian network approach](https://doi.org/10.1111/2041-210X.12062). *Methods in Ecology and Evolution* 4, 760–770

67. Cirtwill, A.R. *et al.* (2019) [A quantitative framework for investigating the reliability of empirical network construction](https://doi.org/10.1111/2041-210X.13180). *Methods in Ecology and Evolution* 10, 902–911

68. Poelen, J.H. *et al.* (2014) [Global biotic interactions: An open infrastructure to share and analyze species-interaction datasets](https://doi.org/10.1016/j.ecoinf.2014.08.005). *Ecological Informatics* 24, 148–159

69. Poisot, T. *et al.* (2016) [Mangal – making ecological network analysis simple](https://doi.org/10.1111/ecog.00976). *Ecography* 39, 384–390

70. Gray, C. *et al.* (2015) [Joining the dots: An automated method for constructing food webs from compendia of published interactions](https://doi.org/10.1016/j.fooweb.2015.09.001). *Food Webs* 5, 11–20

71. Poisot, T. (2023) [Guidelines for the prediction of species interactions through binary classification](https://doi.org/10.1111/2041-210X.14071). *Methods in Ecology and Evolution* 14, 1333–1345

72. Yeakel, J.D. *et al.* (2014) [Collapse of an ecological network in Ancient Egypt](https://doi.org/10.1073/pnas.1408471111). *PNAS* 111, 14472–14477

73. Fricke, E.C. *et al.* (2022) [Collapse of terrestrial mammal food webs since the Late Pleistocene](https://doi.org/10.1126/science.abn4012). *Science* 377, 1008–1011

74. Olivier, P. *et al.* (2019) [Exploring the temporal variability of a food web using long-term biomonitoring data](https://doi.org/10.1111/ecog.04461). *Ecography* 42, 2107–2121

75. Valdovinos, F.S. *et al.* (2023) [A bioenergetic framework for aboveground terrestrial food webs](https://doi.org/10.1016/j.tree.2022.11.004). *Trends in Ecology & Evolution* 38, 301–312

76. Beckerman, A.P. *et al.* (2006) [Foraging biology predicts food web complexity](https://doi.org/10.1073/pnas.0603039103). *Proceedings of the National Academy of Sciences* 103, 13745–13749

77. Van De Walle, R. *et al.* (2023) [Arthropod food webs predicted from body size ratios are improved by incorporating prey defensive properties](https://doi.org/10.1111/1365-2656.13905). *Journal of Animal Ecology* 92, 913–924

78. Petchey, O.L. *et al.* (2008) [Size, foraging, and food web structure](https://doi.org/10.1073/pnas.0710672105). *Proceedings of the National Academy of Sciences* 105, 4191–4196

79. Wootton, K.L. *et al.* (2023) [Towards a modular theory of trophic interactions](https://doi.org/10.1111/1365-2435.13954). *Functional Ecology* 37, 26–43

80. Krause, A.E. *et al.* (2003) [Compartments revealed in food-web structure](https://doi.org/10.1038/nature02115). *Nature* 426, 282–285

81. Lajaaiti, I. *et al.* (2024) [EcologicalNetworksDynamics.jl A Julia package to simulate the temporal dynamics of complex ecological networks](https://doi.org/10.1101/2024.03.20.585899)bioRxiv, 2024.03.20.585899

82. Delmas, E. *et al.* (2017) [Simulations of biomass dynamics in community food webs](https://doi.org/10.1111/2041-210X.12713). *Methods in Ecology and Evolution* 8, 881–886

83. Curtsdotter, A. *et al.* (2019) [Ecosystem function in predator–prey food webs—confronting dynamic models with empirical data](https://doi.org/10.1111/1365-2656.12892). *Journal of Animal Ecology* 88, 196–210

84. Allesina, S. and Pascual, M. (2009) [Food web models: A plea for groups](https://doi.org/10.1111/j.1461-0248.2009.01321.x). *Ecology Letters* 12, 652–662

85. Xie, J.-R. *et al.* (2017) [Completeness of Community Structure in Networks](https://doi.org/10.1038/s41598-017-05585-6). *Scientific Reports* 7, 5269

86. Williams, R.J. and Martinez, N.D. (2008) [Success and its limits among structural models of complex food webs](https://doi.org/10.1111/j.1365-2656.2008.01362.x). *Journal of Animal Ecology* 77, 512–519

87. Becker, D.J. *et al.* (2022) [Optimising predictive models to prioritise viral discovery in zoonotic reservoirs](https://doi.org/10.1016/S2666-5247(21)00245-7). *The Lancet Microbe* 3, e625–e637

88. Terry, J.C.D. and Lewis, O.T. (2020) [Finding missing links in interaction networks](https://doi.org/10.1002/ecy.3047). *Ecology* 101, e03047

89. Roopnarine, P.D. (2006) [Extinction Cascades and Catastrophe in Ancient Food Webs](https://www.jstor.org/stable/4096814). *Paleobiology* 32, 1–19

90. Wells, K. and O’Hara, R.B. (2013) [Species interactions: Estimating per-individual interaction strength and covariates before simplifying data into per-species ecological networks](https://doi.org/10.1111/j.2041-210x.2012.00249.x). *Methods in Ecology and Evolution* 4, 1–8

91. Catchen, M.D. *et al.* (2023) The missing link: Discerning true from false negatives when sampling species interaction networks

92. Allesina, S. *et al.* (2008) [A General Model for Food Web Structure](https://doi.org/10.1126/science.1156269). *Science* 320, 658–661

93. Saravia, L.A. *et al.* (2022) [Ecological network assembly: How the regional metaweb influences local food webs](https://doi.org/10.1111/1365-2656.13652). *Journal of Animal Ecology* 91, 630–642

94. Estay, S.A. *et al.* (2023) Editorial: Patterns and processes in ecological networks over space. *Frontiers in Ecology and Evolution* 11

95. Morales, J.M. and Vázquez, D.P. (2008) [The effect of space in plant–animal mutualistic networks: Insights from a simulation study](https://doi.org/10.1111/j.0030-1299.2008.16737.x). *Oikos* 117, 1362–1370

96. Blüthgen, N. and Staab, M. (2021) [Ecology: Mammals, interaction networks and the relevance of scale](https://doi.org/10.1016/j.cub.2021.05.032). *Current Biology* 31, R850–R853

97. Galiana, N. *et al.* (2018) [The spatial scaling of species interaction networks](https://doi.org/10.1038/s41559-018-0517-3). *Nature Ecology & Evolution* 2, 782–790

98. Rooney, N. *et al.* (2008) [A landscape theory for food web architecture](https://doi.org/10.1111/j.1461-0248.2008.01193.x). *Ecology Letters* 11, 867–881

99. Fortin, M.-J. *et al.* (2021) [Network ecology in dynamic landscapes](https://doi.org/10.1098/rspb.2020.1889). *Proceedings of the Royal Society B: Biological Sciences* 288, rspb.2020.1889, 20201889

100. Brimacombe, C. *et al.* (2021) [Inferred seasonal interaction rewiring of a freshwater stream fish network](https://doi.org/10.1111/ecog.05452). *Ecography* 44, 219–230

101. Laender, F.D. *et al.* (2010) [Carbon transfer in a herbivore- and microbial loop-dominated pelagic food webs in the southern Barents Sea during spring and summer](https://doi.org/10.3354/meps08335). *Marine Ecology Progress Series* 398, 93–107

102. Brimacombe, C. *et al.* (2024) [Applying a method before its proof-of-concept: A cautionary tale using inferred food webs](https://doi.org/10.13140/RG.2.2.22076.65927)

103. Hutchinson, M.C. *et al.* (2019) [Seeing the forest for the trees: Putting multilayer networks to work for community ecology](https://doi.org/10.1111/1365-2435.13237). *Functional Ecology* 33, 206–217

104. Petchey, O.L. *et al.* (2011) [Fit, efficiency, and biology: Some thoughts on judging food web models](https://doi.org/10.1016/j.jtbi.2011.03.019). *Journal of Theoretical Biology* 279, 169–171

105. Berlow, E.L. *et al.* (2008) [The “Goldilocks factor” in food webs](https://doi.org/10.1073/pnas.0800967105). *Proceedings of the National Academy of Sciences* 105, 4079–4080

106. Keyes, A.A. *et al.* (2024) [Synthesising the Relationships Between Food Web Structure and Robustness](https://doi.org/10.1111/ele.14533). *Ecology Letters* 27, e14533

107. Delmas, E. *et al.* (2019) [Analysing ecological networks of species interactions](https://doi.org/10.1111/brv.12433). *Biological Reviews* 94, 16–36

108. Hui, C. and Richardson, D.M. (2019) [How to Invade an Ecological Network](https://doi.org/10.1016/j.tree.2018.11.003). *Trends in Ecology & Evolution* 34, 121–131

109. García-Callejas, D. *et al.* (2023) [Non-random interactions within and across guilds shape the potential to coexist in multi-trophic ecological communities](https://doi.org/10.1111/ele.14206). *Ecology Letters* 26, 831–842

110. Dunhill, A.M. *et al.* (2024) [Extinction cascades, community collapse, and recovery across a Mesozoic hyperthermal event](https://doi.org/10.1038/s41467-024-53000-2). *Nature Communications* 15, 8599