

The complex network of trophic interactions in a subAntarctic Marine Protected Area

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Abstract

Globally, the total area of the world ocean designated under marine protection has increased in the recent decades. The majority of these Marine Protected Areas (MPAs) focus on the presence of particularly vulnerable, keystone, or charismatic species, the presence of large numbers of endemic species, and/or the presence of high biodiversity across taxonomic levels. In the sub-Antarctic region, the level of ocean protection is mainly associated to oceanic islands, except for the MPAs Namuncurá - Banco Burdwood I and II (MPA N-BB, 53°–55°S and 56°–62°W), which are associated to a submarine plateau and its southern adjacent deep slope region, respectively. Here we present the first detailed analysis of the network of predator-prey interactions (food web) for the MPA N-BB ecosystem. We applied a network approach with the objective of characterizing the food web in terms of complexity and structure, and identifying the species' role in such framework. In terms of complexity, the MPA N-BB food web consisted of 1778 predator-prey interactions and 379 species, with a link density of 4.69 and a connectance of 0.01. In terms of structure, almost half of the consumers were omnivores (0.48), and the network displayed a small-world pattern. All this suggest that the ecosystem might be fragile to external perturbations targeting highly connected species, although structural properties might provide resilience and resistance with the final outcome of a rearrange structure maintaining their functions. On the other hand, we identified several species as important in terms of different aspects of trophic structure and functioning, and response to perturbations. We suggest that generalist species, mostly fishes, play a crucial role in the ecosystem's benthic-pelagic coupling process. In addition, we propose that other species rather than the longtail southern cod *Patagonotothen ramsayi* and the Fuegian sprat *Sprattus fuegensis* should be considered as relevant energy transfers for the ecosystem. Finally, we argue that it is the diversity of species, representing the benthic and pelagic habitats, what maintains the connectivity of the food web against perturbations, therefore contributing to the structure and stability of the ecosystem.

Keywords: Food web, Complexity, Structure, Marine Protected Area, Southwest Atlantic

1. Introduction

The evidence of benefits provided by Marine Protected Areas (MPAs) as well as the urgent need for ocean protection have driven an unprecedented increase in the number of MPAs worldwide in recent years (Roberts et al., 2017; Sala et al., 2018). Globally, the total area of the world ocean designated under marine protection adds up to nearly 29,600,000 km², distributed across nearly 18,444 MPAs and covering 8.16% of the ocean's surface (IUCN & UNEP-WCMC, 2023), and therefore approaching the 10% goal of the Convention of Biological Diversity (CBD, 2004). Despite this progress, recent reports have shown that actual protection has been overestimated because it includes areas that are not yet effectively protected (only declared) as well as areas that allow significant extractive activities (Sala et al., 2018).

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In the sub-Antarctic region, the level of ocean protection is mainly associated to oceanic islands, such as the South Georgia and South Sandwich, Bouvet, Prince Edward, and Macquarie islands (IUCN & UNEP-WCMC, 2023). Interestingly, the case of the MPAs Namuncurá - Banco Burdwood I and II (MPA N-BB), which is the focus of this work, is unique since these MPAs are associated to a submarine plateau and its southern adjacent deep slope region, respectively (Falabella, 2017). Added to this, such MPAs are part of a network of protected areas in the sub-Antarctic (jointly with MPA Yaganes), that aims to protect and strengthen the functioning of this southern region to maintain global ocean health.

Many of these MPAs focus on the presence of particularly vulnerable, keystone, or charismatic species, the presence of large numbers (or proportions) of endemic species, and/or the presence of high biodiversity across taxonomic levels (Hogg et al., 2016). Indeed, the MPA N-BB was created to protect a sensitive and biodiverse benthic habitat that was barely known (Falabella, 2017; L. Schejter et al., 2016). The benthic community is featured by a high biomass of vulnerable and fragile species (mainly Porifera and Cnidaria) that meet the characteristics of vulnerable marine ecosystems (Schejter & Albano, 2021), here defined as sites that present taxa with a minimum biomass of 10 kg per 1200 m² (Commission for the Conservation of Antarctic Marine Living Resources, 2009). Also, it provides habitat to several small-sized species (López-Gappa et al., 2018; L. Schejter & Bremec, 2019; Sirito, 2019), and has an important role in the life history of fishes as a food source, refuge and nursery area (Delpiani et al., 2020; García Alonso et al., 2018; Troccoli et al., 2020). The maintenance of this singular community is related to local and regional oceanographic processes, including the circulation of the rich Malvinas (Falkland) current in the area (Guerrero et al., 1999; A. R. Piola & Gordon, 1989) and the upwelling and mixing phenomena (Matano et al., 2019). The input of nutrients coming from the Malvinas (Falkland) current also supports a diverse plankton community (Guinder et al., 2020). Overall, 811 benthic and plankton species have been identified for the MPA N-BB ecosystem, where 349 were reported for the first time for the area (APN, 2022). However, identifying the main species for maintenance of ecosystem services and health as well as for management and conservation is essential.

Recently, the structure of the southwestern South Atlantic Ocean has been proposed to be under a ‘wasp-waist’ control, meaning that the structure and dynamics of the ecosystem is regulated primarily by mid-trophic level species (e.g., fishes, crustaceans) (Padovani et al., 2012; Riccialdelli et al., 2020; Saporiti et al., 2015). In particular, the ecosystem of the MPA N-BB shows a more pronounced ‘wasp-waist’ structure than other sub-Antarctic areas, such as the Atlantic coast of Tierra del Fuego. The Fuegian sprat *Sprattus fuegensis* and longtail southern cod *Patagonotothen ramsayi* are considered the most plausible ‘wasp-waist’ species (Riccialdelli et al., 2020).

High-latitude marine ecosystems, such as the MPA N-BB, are characterized by an unexpected complexity in terms of biodiversity and ecological interactions (Cordone et al., 2020; Day et al., 2013; Kortsch et al., 2019; Trathan et al., 2021). Although there is a robust knowledge about the complexity considering the richness of the benthic and plankton communities in the MPA N-BB ecosystem (APN, 2022; Guinder et al., 2020; L. Schejter et al., 2016, 2020), there is a need for a better understanding of species interactions’ complexity and structure. This aspect can be tackled by analyzing one of the most-frequent relationships occurring between species: the predator-prey interaction (Bascompte, 2009). The sum of predator-prey or trophic interactions of a particular region is referred to as a food web, which represents the roadmap for matter and energy flow in an ecosystem. In recent years, network approaches have been successfully applied to study complex high-latitude marine ecosystems, improving our knowledge on structure, functioning, and response to environmental/anthropogenic changes (Cordone et al., 2018; Funes et al., 2022; Kortsch et al., 2015; Marina et al., 2023). Among anthropogenic threats, it is worth mentioning that contaminants like mercury and microplastics have been recently reported as important threats for the MPA N-BB region (Cossi et al., 2021; Di Mauro et al., 2022; Fioramonti et al., 2022).

In the present work, we present the first, detailed analysis of the network of predator-prey interactions, hereafter food web, for the MPA N-BB ecosystem. For this, we applied a network approach to a highly resolved food web. The objective was twofold: characterize the food web in terms of complexity and structure, and identify the species’ role in such network framework.

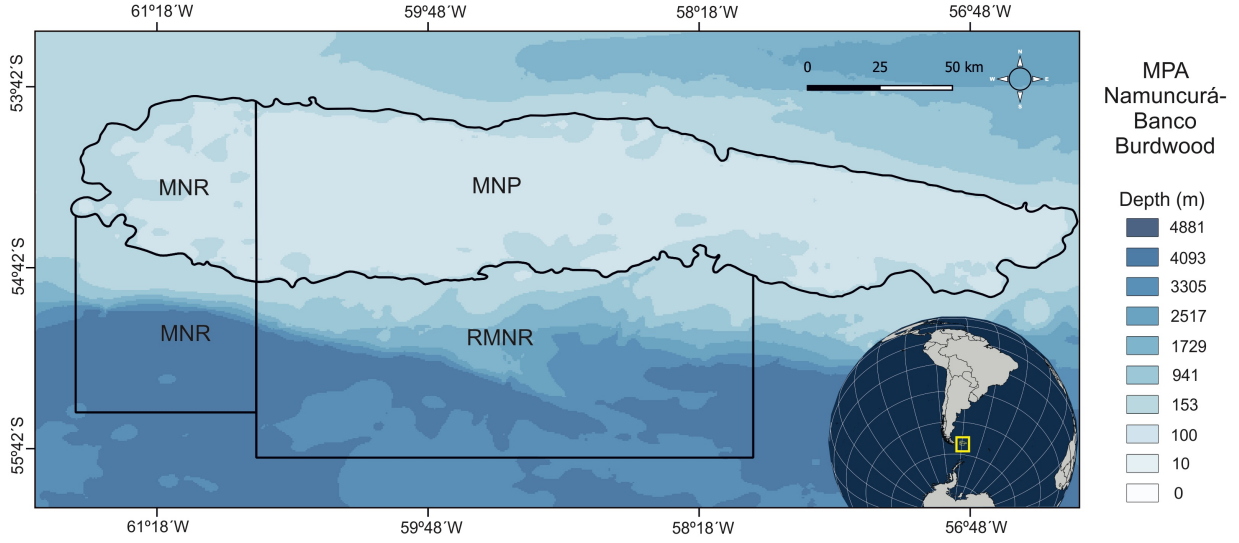


Figure 1: Marine Protected Areas Namuncurá - Banco Burdwood I (MNR and MNP, northern section) and II (MNR and SMNR, southern section). Acronyms indicate categories according to the management plan: MNR - Marine National Reserve, MNP - Marine National Park and RMNR - Restricted Marine National Reserve.

2. Methodology

2.1. Study area

The MPAs Namuncurá - Banco Burdwood I and II, created by National Laws 26.875 and 27.490, comprise a shallow submarine plateau called Burdwood Bank (BB) and a deep slope that reaches 4000 m in depth, N-BB I and N-BB II, respectively (APN, 2023). They are located 150 km east of isla de los Estados and 200 km south from Malvinas/Falkland Islands (Figure 1). The MPA N-BB I comprises nearly 34,000 km² circumscribed by the 200 m isobath, between 54°–55°S and 56°–62°W, with a slight slope extended nearly 370 km east–west. Physical features in the BB are fairly stable, with salinity averaging 34 all year round and temperature ranging between 4 and 8°C (Acha et al., 2004; Guerrero et al., 1999; A. Piola & Falabella, 2009). The BB is surrounded by steep flanks of up to 4000 m depth through which strong currents circulate (Matano et al., 2019; A. R. Piola & Gordon, 1989; Reta, 2014). Such deep slope form the basis of N-BB II that protects about 32,000 km². Intense upwelling and mixing occur over it, entraining deep nutrient-rich waters into the photic layer (Matano et al., 2019; A. Piola & Falabella, 2009), and resulting in a fairly homogeneous water column both spatially and temporally (Glorioso & Flather, 1995; Guerrero et al., 1999; Matano et al., 2019).

Given the evidence collected during several research cruises about the oceanographic and ecological processes connecting MPAs N-BB I and II (references in APN, 2022), a joint management plan was recently proposed (APN, 2022).

2.2. Network construction

In order to build the network of predator-prey interactions we reviewed more than 150 references considering published articles, databases and doctoral theses. Furthermore, we took into account personal communications from experts belonging to the working group of the study area (<https://www.pampazul.gob.ar/tag/banco-burdwood/>). The diversity of the expertise of the authors contributing to the present study was a key factor in enhancing the quality of the network, and inherently improved the network representation. A list of the references used to build the network is presented in Supplementary Material (Table S1).

Due to a lack of trophic data resolution for some species inhabiting the study area, we followed the concept of trophic species, here defined as aggregated groups of taxa. In most cases, we followed it when

specific data on species, in the taxonomic sense, were not available. In some cases, we collapsed species when taxa shared the same set of predators and prey (trophic similarity, Martinez (1991)), one of the aggregation methods that better preserves food web functional properties (Gauzens et al., 2013). Details about this can be found in Supplementary Material (Table S2).

With the gathered trophic data we constructed a matrix of pairwise interactions; a value of 1 or 0 was assigned to each element a_{ij} of the matrix depending on whether the j -species preyed or not on the i -species. Then we transformed such matrix into an oriented graph with L trophic interactions between S nodes or species. The orientation or direction of the graph follows the flow of energy and matter in the network, from prey to predator.

2.3. Network analysis

We analyzed the MPA N-BB network of trophic interactions, or food web, at two levels: A) network, considering species and interactions of the whole network; and B) species, considering interactions and species related to a particular species (Table 1).

The network-level analysis aims to characterize the food web in terms of complexity and structure. For this we calculated several network properties commonly used to describe empirical food webs (Pascual & Dunne, 2005): (1) number of species S ; (2) number of interactions or links L ; (3) link density L/S ; (4) connectance L/S^2 ; (5) omnivory Omn ; and (6) small-world pattern. In order to explore the small-world phenomenon we analyzed the characteristic path length (CPL) and the clustering coefficient (CC). The CPL is defined as the average shortest path length between all pairs of nodes (Watts & Strogatz, 1998). Here, CPL was calculated as the average number of nodes in the shortest path $CPL_{Min}(i, j)$ between all pairs of nodes $S(i, j)$ in a network averaged over $S(S - 1)/2$ nodes:

$$CPL = \frac{2}{S(S - 1)} \sum_{i=1}^S \sum_{j=1}^S CPL_{Min}(i, j)$$

The CC quantifies the local interconnectedness of the network and it is defined as the fraction of the number of existing links between neighbours of node i among all possible links between these neighbours. In this study, the CC was determined as the average of the individual clustering coefficients CC_i of all the nodes in the network. Individual CC_i were determined as follows:

$$CC_i = \frac{2E_i}{K_i(K_i - 1)}$$

where E_i is the effective number of interactions between K_i nearest-neighbor nodes of node i and the maximal possible number of such interactions (Newman, 2003). With the aim of testing whether the food web presented the small-world pattern, we compared the empirical values of CPL and CC with those resulting from 1000 randomly generated networks with the same size (S) and number of interactions (L), following the method proposed by Marina et al. (2018).

Also, we estimated the (7) degree distributions for the food web, for prey and predators, and for each functional group (e.g., Amphipoda, Ascidiacea, Bivalvia, Fish, Marine mammals, Sea birds, etc.). The prey and predator distributions indicate the frequency of prey among predators, and viceversa; the functional group's degree shows the distribution of interactions within groups.

The species-level analysis aims to describe the species' role in the food web. For this, we considered the following properties: (1) betweenness Btw ; (2) closeness Cl ; (3) trophic similarity TS ; (4) topological role TR ; and (5) trophic level TL (Table 1). Topological roles refers to the fact that food webs tend to naturally organize in non-random, modular patterns, where modules are defined as a group of species that interact more frequently among themselves than with species that are not members of the module (Guimerà & Nunes Amaral, 2005). Species can play different roles in this respect, according to the pattern of interactions they have within their own module and/or across modules. We computed the topological role for each species, classified as: module hub, species with a relatively high number of interactions, but most within its own module; module specialist, species with relatively few interactions and most within its own module; module

connector, species with relatively few interactions mainly between modules; and network connector, species with high connectivity between and within modules (Guimerà & Nunes Amaral, 2005).

We also studied the relationship between species TL and the other species properties by performing linear regression analyses. Thus, we considered the TL as the dependent variable and the given property (i.e. betweenness, closeness, trophic similarity) as the independent variable, and obtained the coefficients (slope and intercept) for the linear model. Models were fitted using the least squares approach. We also explored the topological role categories with the species TL. These species-level properties provide a general appropriate description of species' role in empirical complex food webs (Cirtwill et al., 2018).

All network analyses and graphs were performed in R version 4.2.2 (Team, 2022), mainly using 'igraph' (Csardi & Nepusz, 2006) and 'multiweb' (Saravia, 2022) packages. The source code and data are available at <https://github.com/TomasMarina/Banco-Burdwood>.

Table 1: List of network and species-level properties analysed, definitions, and relevant ecological implications related to food web complexity and structure.

Name	Definition	Implications	Reference
Number of species	Number of trophic species in a food web.	It represents the species diversity and has implications for the persistence of the ecosystem.	May 1973, Tilman 1996
Number of interactions	Total number of trophic interactions in a food web.	It represents the number of pathways along which matter and energy can flow.	Dunne et al. 2002
Link density	Ratio of interactions to species in a food web	It represents the average number of interactions per species; informs about how connected species are in the food-web.	Dunne et al. 2002
Connectance	Proportion of potential links among species that are actually realized. Range = 0 - 1.	It measures the probability of interactions and is a fundamental measure of network complexity. Connectance can be negatively or positively associated with food web robustness, depending on the network structure (random vs non-random) or how the strength of the interactions are distributed.	Martinez 1992
Degree distribution	Frequency of trophic species that have k or more interactions.	It suggests on the vulnerability of complex food webs against random failures and intentional attacks (i.e. species extinctions).	Albert & Barabási 2002

Name	Definition	Implications	Reference
Omnivory	Species feeding on prey from more than one trophic level.	It influences food web's stability; intermediate levels of omnivory may stabilize it and may diffuse top-down effects thus reduce the probability of trophic cascades.	McCann & Hastings 1997
Small-world pattern	A network with short path length (distance between nodes) and high clustering coefficient (formation of compartments) compared to random networks.	Consequences of this structural pattern in food webs are of great importance in recognizing evolutionary paths and the vulnerability to perturbations.	Watts & Strogatz 1998, Montoya & Solé 2002
Betweenness	Number of shortest paths going through a species.	Species with high betweenness act as "bridges"; if removed, would have rapidly spreading effects in the food web.	Freeman 1978, Lai et al. 2012
Closeness	Number of steps required to reach every other species from a given species.	The removal of a species with high closeness will affect the most other species in the food web.	Freeman 1978, Lai et al. 2012
Trophic similarity	Trophic overlap based on shared and unique resources (prey) and consumers (predators).	It measures one of the most important aspects of species' niches, the trophic niche, and functional aspects of biodiversity.	Martinez 1992
Topological role	Species role according to interactions within and across modules (subgroups of species).	Four roles are defined: module hub, module specialist, module connector and network connector. Network connector and module connector roles maintain the connectivity of the food web.	Guimera & Nunes Amaral 2005

3. Results

3.1. Network-level properties

In terms of complexity, the MPA Namuncurá - Banco Burdwood food web (N-BB I and II) consisted of 1778 predator-prey interactions and 379 species, where 93% of them were defined at the species taxonomical level (Figure 2, Table S2). The food web presented a link density (e.g., the average number of interactions per species) of 4.69, and a connectance of 0.01. Almost half of the consumers were omnivores (0.48), feeding on sources at different trophic levels. The food web displayed a small-world pattern, meaning that the path length was lower and the clustering coefficient higher than the random networks (Table 2).

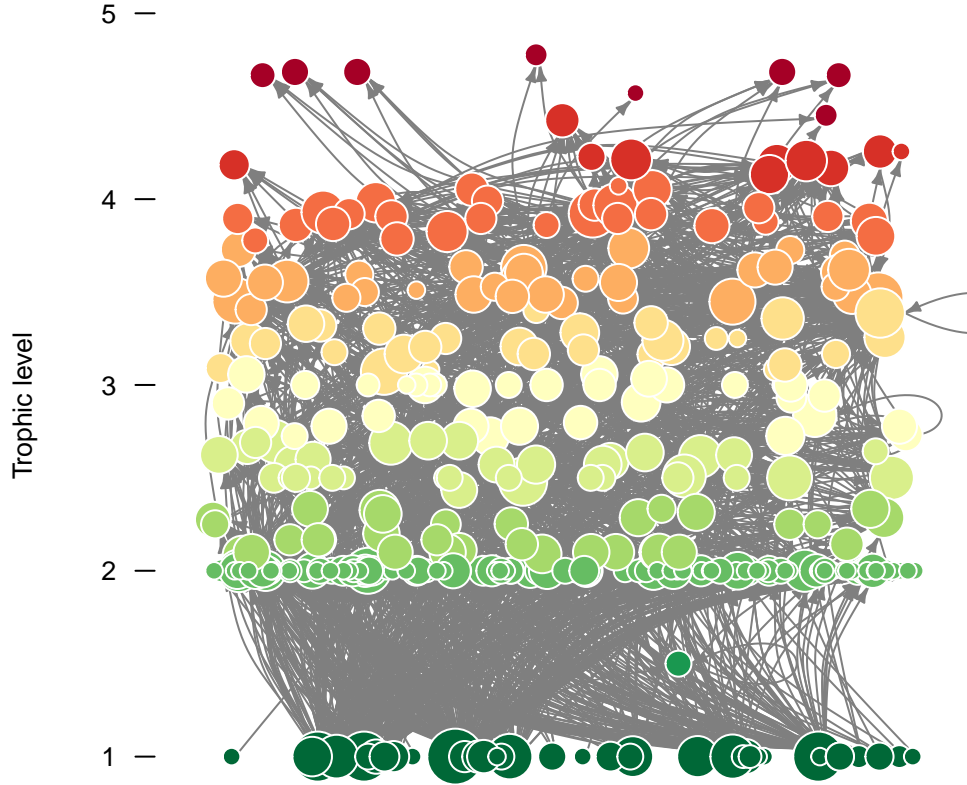


Figure 2: Graph of the food web for the MPA Namuncurá - Banco Burdwood. Circles represent species and arrows trophic interactions. Circle diameter is relative to the number of interactions. Colour gradient indicates the trophic level.

Table 2: Network-level properties of the MPA Namuncurá - Banco Burdwood food web. CPL: Characteristic Path Length; CC: Clustering Coefficient; SW: Small-World pattern. See table 1 for definitions and ecological relevance.

Species	Interactions	Density	Connectance	Omnivory	CPL	CC	SW
379	1778	4.69	0.01	0.48	2.99	0.08	True

The degree distribution of the food web showed an asymmetric frequency in the number of interactions, where most of the species had a relatively low number of interactions and few species concentrated the majority of them (Figure 3A). The distribution of prey among predators showed that most of the consumers fed on a low number of prey whereas few of them had multiple preys (Figure 3B). These were the top-five predators in number of prey: *Patagonotothen guntheri* (Notothenioid fish, 52 prey), *Patagonotothen ramsayi* (Notothenioid fish, 50 prey), *Dissostichus eleginoides* (Notothenioid fish, 30 prey), *Bathyraja brachyurops* (Chondrichthyan, 30 prey), and *Bathyraja griseocauda* (Chondrichthyan, 28 prey). Following the same distribution pattern, few prey presented multiple predators (Figure 3C). These were the top-five prey in number of predators: Detritus (Non-living, 153 predators), the three categories of Diatoms considered (benthic, centric and pennate, 72 predators on average), and species of the genus *Euphausia* (Zooplankton, 46 predators). Finally, taking into account the interactions within each functional group, again the majority of the interactions were concentrated in few species (Figure 3D). The most evident species were: *Themisto gaudichaudii* (Amphipoda), *Zygochlamys patagonica* (Bivalvia), *Aspidostoma giganteum* (Bryozoa), *Munida gregaria* (Decapoda), *Patagonotothen ramsayi* and *Patagonotothen guntheri* (benthic-pelagic fish), *Psychrolutes marmoratus* (demersal fish), and species of *Euphausia* (Zooplankton). Overall, there is an evident asymmetry in the distribution of interactions among species at different levels in the MPA Namuncurá - Banco Burdwood food web.

A list of the distribution of interactions per species is presented in Supplementary Material (Table S3).

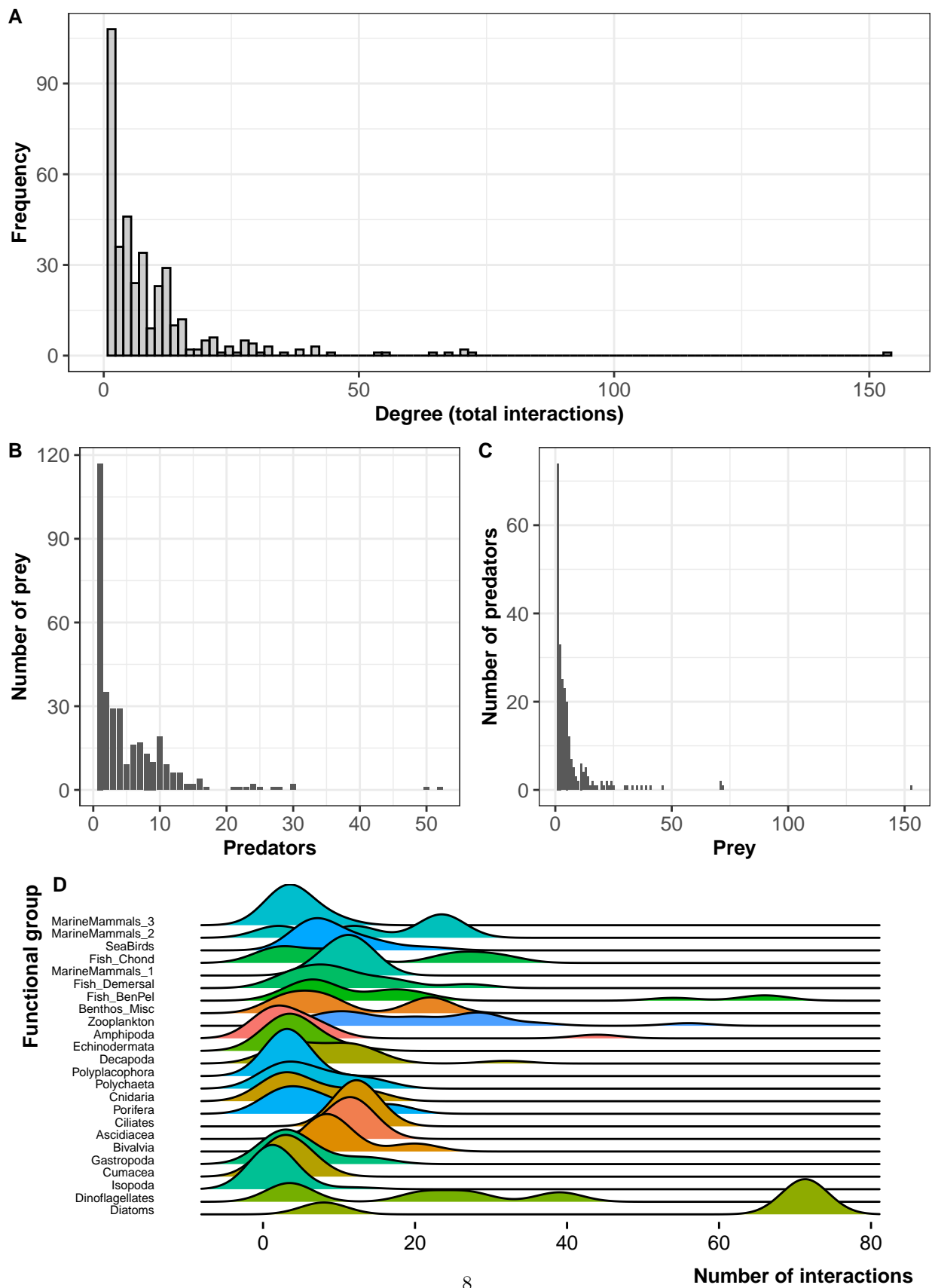


Figure 3: Degree distributions for the (A) food web, for (B) prey among predators, (C) predators among prey, and (D) for each functional group. Groups are vertically ordered by increasing trophic level; groups with less than 3 species were not plotted (e.g., pelagic fish).

3.2. Species-level properties

The majority of the species in the food web were consumers, 336 out of 379; the rest were primary producers, such as diatoms (phytoplankton), and non-living food sources like detritus and necromass.

We found different relationships between the species trophic level (TL) and the rest of the analysed species-level properties (Figure 4A-D). The most evident significant relationship was with trophic similarity, i.e. the higher the species' TL, the lower the trophic similarity or the higher the uniqueness in terms of trophic role (Figure 4C). Here it is noteworthy to highlight those high-trophic level species with low values of trophic similarity: *Bathyraxia macloviana* and *Squalus acanthias* (Chondrichthyans), *Diplopteraster clarki* and *Pteraster sp* (echinoderms), *Phalacrocorax atriceps* and *Eudyptes chrysocome* (sea birds), and *Lagenorhynchus cruciger* and *Mesoplodon bowdoini* (marine mammals) (Table S3).

We also found a negative significant relationship between TL and closeness, however less evident, meaning that low-TL species are relatively closer to any other species in the food web (Figure 4B). Species of genera *Calanus* and *Euphausia*, and species of Brachiopoda, all of them with $TL < 3$, registered the highest values of closeness (Table S3).

It is noteworthy that the highest values of betweenness were shown by species of mid-TLs (3-4), meaning that those species participated in the highest number of shortest paths between species (Figure 4A). The following are the species with the highest values: *Patagonotothen ramsayi*, *Dissostichus eleginoides*, *Salilota australis* (fishes), *Doryteuthis gahi* (Cephalopoda), and *Patagonotothen guntheri* (Notothenioid fish) (Table S3).

Considering the topological role, 'module specialist' species were the most frequent and presented a wide TL range (1 - 4.77); 'module hub' was constrained to mid-TL species (2.48 - 3.92); 'module connector' from low to mid-TLs (1 - 3.86); and 'network connector', the least frequent, had all of its species in $TL = 1$, except for one with $TL = 3.47$ (Figure 4D, see Figure S2 for species' topological roles in a food web graph framework). Here it is important to highlight the two latter topological roles, because they are responsible for linking modules and maintaining the connectivity of the food web: 40 species (5 network connectors + 35 module connectors) from 20 different functional groups with a TL range = 1 - 3.86. The following were the five species with a network connector role: Detritus, Diatoms (benthic, centric and pennate), and *Patagonotothen ramsayi*. The 35 species with a module connector role represent these functional groups: Amphipoda, Bivalvia, Brachiopoda, Bryozoa, Cnidaria, Cumacea, Decapoda, Detritus, Diatoms, Dinoflagellates, Echinodermata, Fish (benthic-pelagic, demersal, chondrichthyans), Foraminifera, Polychaeta, Porifera, Pycnogonida, Sea birds, and Zooplankton (see Supplementary Material Table S3 for the identity of the species).

An exhaustive list of the species-level properties is presented in Supplementary Material (Table S3).

4. Discussion

4.1. The food web of the MPA Namuncurá - Banco Burdwood ecosystem

The food web of the MPA Namuncurá - Banco Burdwood ecosystem analysed in this study is one of the most highly-resolved networks of trophic interactions ever studied not only for a high-latitude marine ecosystem but also for a marine protected area worldwide. It is of paramount importance to consider the complexity of species interactions in order to gain insights into the structure and functioning of the ecosystem, since the aggregation of species might mask food web properties and/or produce type II errors (false positive) (Gauzens et al., 2013; Martinez, 1993).

Food web connectance is a feature that resumes the complexity of the network, but more importantly, it is an emergent property of pairwise species interactions (Poisot & Gravel, 2014). It contains information regarding how interactions within an ecological network are distributed, and predicts reasonably well key dynamical properties of ecological networks (Dunne et al., 2002a). Complex marine food webs (i.e. with more than 25 trophic species) show connectance values ranging from 0.01 - 0.27 (Marina et al., 2018). In particular, food webs from high-latitude regions tend to exhibit a connectance closer to the minimum (between 0.01 and 0.05) (Kortsch et al., 2015; Rodriguez et al., 2022; Santana et al., 2013). Whether food webs display a low or a high connectance helps to better comprehend ecosystem's synthetic properties like robustness. In this sense, empirical analyses support the notion that highly-connected ecological networks are robust against

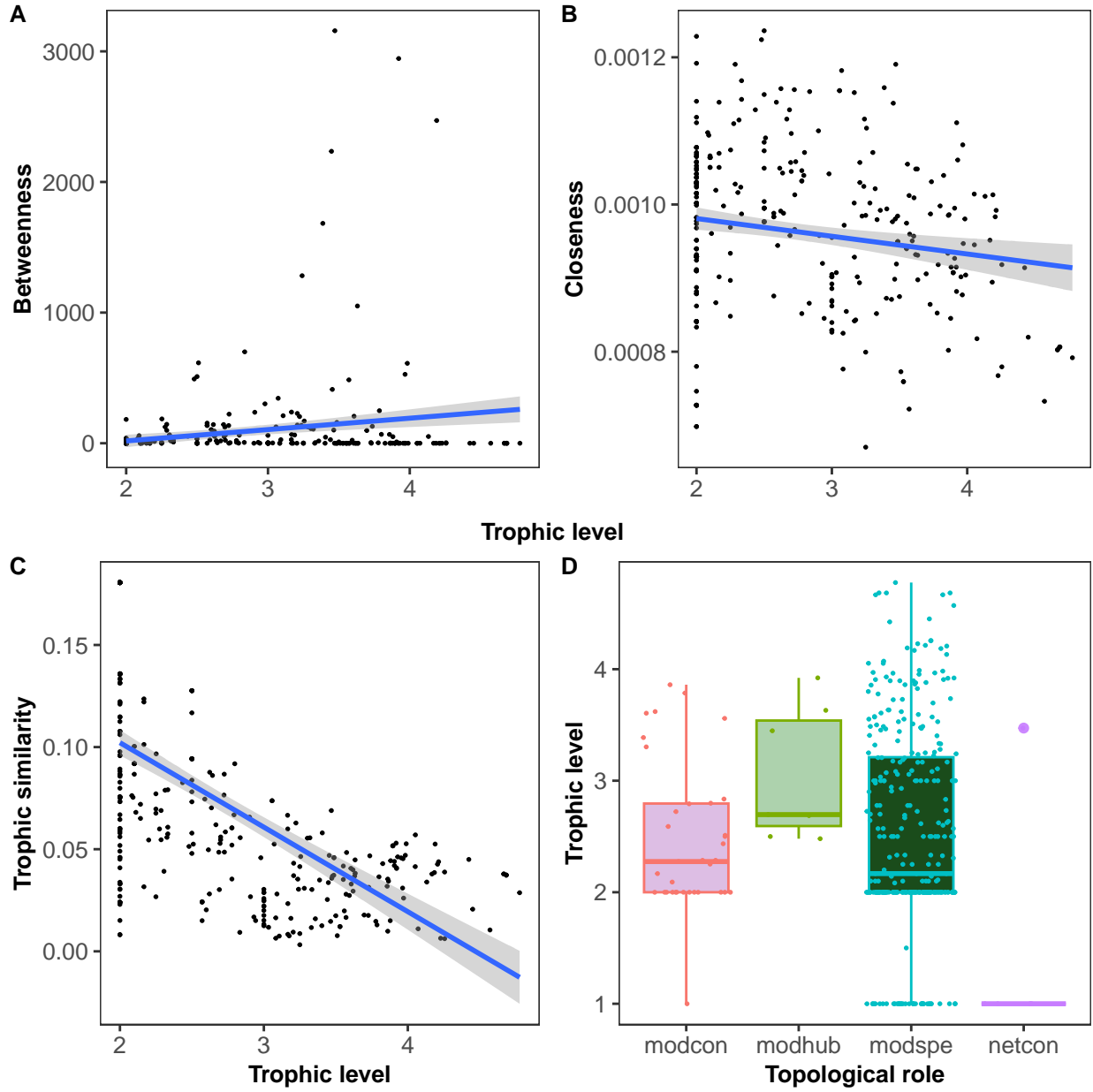


Figure 4: Species-level properties by trophic level: (A) betweenness, (B) closeness, (C) trophic similarity, and (D) topological role. Each point represents a species. Linear regressions for betweenness ($y = 72.48x - 111.98$, $R^2 = 0.04$, $p\text{-value} < 3.38e-05$), closeness ($y = 5.78e-06x - 9.37e-04$, $R^2 = -0.0005$, $p\text{-value} = 0.37$) and trophic similarity ($y = -0.01x - 0.11$, $R^2 = 0.07$, $p\text{-value} = 6.76e-08$). Note that for A, B and C panels only species with TLs equal or greater than 2 were considered.

external perturbations such as the introduction of new (e.g., invasive) species (Smith-Ramesh et al., 2017) as well as species removal (e.g., local extinction) (Dunne et al., 2002b; Montoya & Solé, 2003). The connectance of the food web of the MPA Namuncurá - Banco Burdwood (0.01) is one of the lowest reported so far for these regions; in particular, it appears to be five times lower than that of Beagle Channel, an adjacent area (Rodríguez et al., 2022).

The degree distribution or the distribution of number of interactions per species is the core for the structure of species interactions, which influences the opportunities for multiple species to persist in the long term and, therefore, their coexistence (Godoy et al., 2018). The food web for the MPA N-BB presents an asymmetric degree distribution. This pattern was identified at different levels of analysis: food web, predator, prey, and functional group. Such asymmetry is a well-known feature in empirical complex food webs in particular (Dunne et al., 2002a; Montoya & Solé, 2003; Stouffer et al., 2005), and has received great attention in complex networks in general (Albert & Barabási, 2002; Newman, 2003). The degree distribution affects the resilience of complex food webs against random failures and intentional attacks: food webs showing right-skewed distributions, like the one described in this study, are more vulnerable to the removal of the most connected species or hubs, with the potential of producing secondary extinctions and a catastrophic fragmentation of the network (Albert et al., 2000; Dunne et al., 2002b; Eklöf & Ebenman, 2006).

It is suggested that the small-world pattern, i.e., a network with short path length and high clustering coefficient is not frequent in complex marine food webs, mainly due to a low clustering coefficient compared to random networks (Dunne et al., 2002c; Marina et al., 2018). However, the food web of the MPA N-BB does display a small-world pattern. Consequences of this could be of great importance in recognizing evolutionary paths and the vulnerability to perturbations (Montoya & Solé, 2002). On the one hand, a short path length implies a rapid spread of an impact (e.g., contaminant, population fluctuation, local extinction) throughout the network and, at the same time, a more potentially adaptive dynamics in face of external perturbations (Montoya & Solé, 2002; Williams et al., 2002). On the other hand, a high clustering coefficient indicates the formation of subnetworks composed only by the neighbours of particular species. This translates into a greater resistance of the network due to the confinement of perturbations mainly within subnetworks and not spreading between them (Heer et al., 2020; Kortsch et al., 2019). Overall, a small-world topology provides ecological networks with a greater resilience and resistance (Bornatowski et al., 2017; Dormann et al., 2017).

Omnivory acts as a buffer to changes as the ecosystem presents alternative energy pathways in the face of perturbations; i.e., reducing the risk of cascading extinctions following primary loss of species (Borrvall et al., 2000). This is based on the fact that omnivores are species able to adapt faster and to a wider range of environmental conditions by changing their foraging habits to feed on the most abundant prey (Fagan, 1997). Furthermore, omnivory can be analysed from the interaction point of view: theoretical studies have identified omnivorous interactions as a possible candidate for a keystone interaction, sensu Kadoya et al. (2018), highlighting the importance of omnivory in stabilizing food web dynamics (McCann & Hastings, 1997; Neutel et al., 2002). The high proportion of omnivory in the food web of the MPA N-BB suggests that the network might be robust to variations in prey abundances, which could increase food web's persistence and stability (Stouffer & Bascompte, 2010).

To resume, the food web of the MPA Namuncurá - Banco Burdwood presents a combination of network properties that make it unique in terms of network resolution, complexity, and structural pattern. All this suggest that the ecosystem might be fragile to external perturbations targeting highly connected species. However, structural properties might provide resilience and resistance with the final outcome of a rearrange structure maintaining their functions.

4.2. Dominant consumers and food sources

The degree distribution allows to identify important species, such as potential keystone species (Dunne et al., 2002b; Solé & Montoya, 2001), generalist/specialist species, and dominant food sources (Kondoh et al., 2010).

We have identified that the majority of the consumers in the food web of the MPA N-BB either has a narrow diet or are specialists, while few present a broad diet. The most evident generalist species are *Patagonotothen guntheri*, *P. ramsayi*, *Dissostichus eleginoides*, *Bathyraja brachyurops*, and *B. griseocauda*,

with more than 25 potential prey. Since these species present mid-trophic positions in the food web (3-4), acting as predator and prey, they might be important links between lower and higher trophic levels. This is in agreement with the sole analysis, using stable isotopes, that exists so far for the trophic structure of the MPA N-BB (Ricciardelli et al., 2020), and resembles other high-latitude marine systems of the Southwest Atlantic and Arctic regions (Arkhipkin & Laptikhovsky, 2013; Kortsch et al., 2019). The importance of these particular generalist species also arises since they feed on the benthic and pelagic habitats (Covatti Ale et al., 2022; Fischer et al., 2022; Troccoli et al., 2020), linking these realms and contributing to the vertical carbon flow.

On the other hand, few prey are consumed by many predators in the food web of the MPA N-BB. This suggests that there are dominant food sources from which most consumers depend on, and from where the ecosystem energy is being transferred to the upper trophic levels. The most demanded source we identified in this study (i.e. detritus) supports the abundant benthic community of filter-feeders (L. Schejter et al., 2016). Furthermore, we found that the second and third-most consumed prey were diatoms and species of *Euphausia*, respectively, which are essential sources for the diverse zooplankton community (Spinelli et al., 2020), mid-TL consumers like the Fuegian sprat *Sprattus fuegensis* (Padovani et al., 2021) and *Patagonotothen ramsayi* (Fischer et al., 2022), and top predators (Raya Rey, 2022; Ricciardelli, 2022).

4.3. Species' role related to their trophic level

Describing species' roles in food webs provides a toolbox with which to assess the significance of species in terms of community's functioning and overall stability (Cirtwill et al., 2018; Thébault & Fontaine, 2010). We used a range of descriptors to characterise the dynamic and multifaceted nature of the species forming the MPA N-BB food web.

Closeness and betweenness are defined as “meso-scale” properties because they consider direct and indirect interactions, therefore describing the focal species' ability to influence the rest of the species of the food web (Lai et al., 2012). Closeness quantifies how many steps away species *i* is from all other species in the food web, and is proportional to how rapidly the indirect effects of the focal species can spread to other species in the network (Scotti & Jordán, 2010). In the food web of the MPA N-BB, low-TL consumers arise as important in this regard: species of the zooplankton community, *Calanus* and *Euphausia*, and Brachiopoda. Any perturbation affecting these species, such as the recently confirmed contaminants mercury (Fioramonti et al., 2022) and microplastics (Cossi et al., 2021; Di Mauro et al., 2022) for the area, should be of concern since it might reach many other species in the food web. Otherwise, betweenness measures the number of shortest paths between species, providing information on the importance of species as “bridges” for energy transfer: a species with high betweenness takes part in more food chains and therefore affects more energy flows (Scotti & Jordán, 2010). We have identified the longtail southern cod *Patagonotothen ramsayi* as the most important species in this sense. Moreover, in light of our analysis, other species like the Patagonian toothfish *Dissostichus eleginoides*, the Patagonian cod *Salilota australis*, the yellowfin notothen *Patagonotothen guntheri*, and the Patagonian longfin squid *Doryteuthis gahi* should be considered as relevant in the energy transfer in the ecosystem. All these species have a mid-trophic position in the food web, supporting the ‘wasp-waist’ control hypothesis for the MPA N-BB (Ricciardelli et al., 2020).

Ecosystems with a pronounced ‘wasp-waist’ structure are suggested to present a high trophic redundancy, since a large proportion of species would show similar trophic habits (Cury et al., 2000). The significant negative relationship we found between trophic similarity and trophic level enhances the hypothesis of functional redundancy at low and mid-TL species compared to higher TL species for the MPA N-BB food web (Ricciardelli et al., 2020). At the same time, our results highlight the uniqueness in terms of trophic role for high-TL predators. Here, not only the expected pelagic animals such as marine mammals and sea birds arise as relevant, but also benthic vertebrate (chondrichthyans *Bathyraja macloviana* and *Squalus acanthias*) and invertebrate species (echinoderms *Diplopteraster clarki* and *Pteraster sp*) are noteworthy. The role that such species play in the MPA N-BB ecosystem is unique and perturbations on them might result in unprecedented changes at the trophic structure and functioning level.

Species' role can be also assessed in a module-based context. Among the varying numbers of topological roles in which species can be divided, two are remarkable: ‘module connector’ and ‘network connector’. Here again the longtail southern cod *Patagonotothen ramsayi* is emphasized as an influential species for

the connectivity of the food web, which agrees with the suggested ‘wasp-waist’ control for the MPA N-BB ecosystem (Ricciardelli et al., 2020). But most importantly, our results broadens this issue by pointing out that there are other species, belonging to a wide range of trophic positions (1 to 3.86) and representing 20 different functional groups, that should be considered. Thus, we propose that it is the diversity of species (benthic and pelagic) what maintains the connectivity of the food web, therefore contributing to the trophic structure and ecosystem’s stability.

4.4. Caveats and future perspectives

In construction. . .

Caveats:

Seasonal variation in predators’ diet due to food source availability. Brose et al. (2019)

Spatial variation of study area: Banco Burdwood west-east gradient. In the oceanic BB area, complex oceanographic conditions generate an internal spatial heterogeneity, mainly along its longitudinal axis (Matano et al., 2019), and consequently plankton communities respond with different spatial distributions (Spinelli et al., 2020). Kortsch et al. (2019)

Perspectives:

Incorporate habitat heterogeneity, since benthic community in the BB and deep flanks is different (Schejter & Albano, 2021). This might lead to distinct food web properties, hence differences in ecosystem functioning and response to anthropogenic and environmental perturbations. Cordone et al. (2020)

Simulate perturbations on food web and target species: mercury transfer, microplastic pollution, fisheries. Cossi et al. (2021); Di Mauro et al. (2022); Fioramonti et al. (2022)

Estimation of interaction strength considering species and interaction traits (i.e. body size, body mass, interaction dimensionality) to gain insights into food web stability and response to anthropogenic and environmental perturbations. Nilsson & McCann (2016); Cirtwill et al. (2018); Kortsch et al. (2021)

5. Conclusion

We compiled information on the species and trophic diversity of the Marine Protected Area Namuncurá - Banco Burdwood, generating an unprecedented, well-resolved network of trophic interactions for a sub-Antarctic ecosystem, identifying the complexity and structure of the system, and the main species role in a network framework. Particular properties at the network level allowed us to identify ecosystem’s vulnerability and potential response to perturbations in the presence of highly-connected species, with a rearrange structure maintaining their functions due to its potential resilience and resistance.

We identified several species as important in terms of different aspects of trophic structure and functioning, and response to perturbations (i.e. environmental/anthropogenic changes). On one hand, we suggest that generalist species, mostly fishes, play a crucial role in the ecosystem’s benthic-pelagic coupling process. At the same time, we propose that other species rather than the longtail southern cod *Patagonotothen ramsayi* and the Fuegian sprat *Sprattus fuegensis* should be considered as relevant energy transfers for the ecosystem. Finally, we argue that it is the diversity of species, representing the benthic and pelagic habitats, what maintains the connectivity of the food web against perturbations, therefore contributing to the structure and stability of the ecosystem.

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References

- Acha, E. M., Mianzan, H. W., Guerrero, R. A., Favero, M., & Bava, J. (2004). Marine fronts at the continental shelves of austral South America: Physical and ecological processes. *Journal of Marine Systems*, 44(1), 83–105. <https://doi.org/10.1016/j.jmarsys.2003.09.005>
- Albert, R., & Barabási, A.-L. (2002). Statistical mechanics of complex networks. *Reviews of Modern Physics*, 74(1), 47–97. <https://doi.org/10.1103/RevModPhys.74.47>
- Albert, R., Jeong, H., & Barabási, A.-L. (2000). Error and attack tolerance of complex networks. *Nature*, 406(6794), 378–382. <https://doi.org/10.1038/35019019>
- APN. (2022). *Plan de gestión AMP Namuncurá Banco Burdwood*.
- APN. (2023). *Área Marina Protegida Namuncurá Banco Burdwood*.
- Arkhipkin, A., & Laptikhovsky, V. (2013). From gelatinous to muscle food chain: Rock cod Patagonotothen ramsayi recycles coelenterate and tunicate resources on the Patagonian Shelf. *Journal of Fish Biology*, 83(5), 1210–1220. <https://doi.org/10.1111/jfb.12217>
- Bascompte, J. (2009). Disentangling the Web of Life. *Science*, 325(5939), 416–419. <https://doi.org/10.1126/science.1170749>
- Bergagna, L. (2019). *Descripción de los ensambles bentónicos del AMP Namuncurá-Banco Burdwood y estimación del contenido de carbono -orgánico e inorgánico- allí presente*. [Bachelor Thesis]. Universidad Nacional de Córdoba.
- Bergagna, L., Chiesa, I., Álvarez, Y. P., Ceballos, S. G., Boy, C. C., Riccialdelli, L., & Lovrich, G. A. (2020). En el fondo el Banco es bueno. *La Lupa. Colección Fueguina de Divulgación Científica*, 16, 20–24.
- Bornatowski, H., Barreto, R., Navia, A. F., & de Amorim, A. F. (2017). Topological redundancy and “small-world” patterns in a food web in a subtropical ecosystem of Brazil. *Marine Ecology*, 38(2), e12407. <https://doi.org/10.1111/maec.12407>
- Borrelli, J. J., & Ginzburg, L. R. (2014). Why there are so few trophic levels: Selection against instability explains the pattern. *Food Webs*, 1(1), 10–17. <https://doi.org/10.1016/j.fooweb.2014.11.002>
- Borrvall, C., Ebenman, B., & Tomas Jonsson, T. J. (2000). Biodiversity lessens the risk of cascading extinction in model food webs. *Ecology Letters*, 3(2), 131–136. <https://doi.org/10.1046/j.1461-0248.2000.00130.x>
- Brickle, P., Arkhipkin, A. I., Laptikhovsky, V., Stocks, A., & Taylor, A. (2009). Resource partitioning by two large planktivorous fishes *Micromesistius australis* and *Macrurus magellanicus* in the Southwest Atlantic. *Estuarine, Coastal and Shelf Science*, 84(1), 91–98. <https://doi.org/10.1016/j.ecss.2009.06.007>
- Brose, U., Archambault, P., Barnes, A. D., Bersier, L.-F., Boy, T., Canning-Clode, J., Conti, E., Dias, M., Digel, C., Dissanayake, A., Flores, A. A. V., Fussmann, K., Gauzens, B., Gray, C., Häussler, J., Hirt, M. R., Jacob, U., Jochum, M., Kéfi, S., ... Iles, A. C. (2019). Predator traits determine food-web architecture across ecosystems. *Nature Ecology & Evolution*, 3(6), 919–927. <https://doi.org/10.1038/s41559-019-0899-x>
- CBD. (2004). *Secretariat of the convention on biological diversity: Technical advice on the establishment and management of a national system of marine and coastal protected areas, Technical Series number 13*.
- Cirtwill, A. R., Dalla Riva, G. V., Gaiarsa, M. P., Bimler, M. D., Cagua, E. F., Coux, C., & Dehling, D. M. (2018). A review of species role concepts in food webs. *Food Webs*, 16, e00093. <https://doi.org/10.1016/j.fooweb.2018.e00093>
- Commission for the Conservation of Antarctic Marine Living Resources. (2009). *Vulnerable Marine Ecosystem taxa identification guide*.
- Cordone, G., Marina, T. I., Salinas, V., Doyle, S. R., Saravia, L. A., & Momo, F. R. (2018). Effects of macroalgae loss in an Antarctic marine food web: Applying extinction thresholds to food web studies. *PeerJ*, 6, e5531. <https://doi.org/10.7717/peerj.5531>
- Cordone, G., Salinas, V., Marina, T. I., Doyle, S. R., Pasotti, F., Saravia, L. A., & Momo, F. R. (2020). Green vs brown food web: Effects of habitat type on multidimensional stability proxies for a highly-resolved Antarctic food web. *Food Webs*, 25, e00166. <https://doi.org/10.1016/j.fooweb.2020.e00166>
- Cossi, P. F., Ojeda, M., Chiesa, I. L., Rimondino, G. N., Frayssé, C., Calcagno, J., & Pérez, A. F. (2021). First evidence of microplastics in the Marine Protected Area Namuncurá at Burdwood Bank, Argentina:

- A study on *Henricia obesa* and *Odontaster penicillatus* (Echinodermata: Asteroidea). *Polar Biology*, 44(12), 2277–2287. <https://doi.org/10.1007/s00300-021-02959-5>
- Covatti Ale, M., Fischer, L., Deli Antoni, M., Díaz de Astarloa, J. M., & Delpiani, G. (2022). Trophic ecology of the yellowfin notothen, *Patagonotothen guntheri* (Norman, 1937) at the Marine Protected Area Namuncurá-Burdwood Bank, Argentina. *Polar Biology*, 45(4), 549–558. <https://doi.org/10.1007/s00300-022-03011-w>
- Csardi, & Nepusz. (2006). *The igraph software package for complex network research*.
- Cury, P., Bakun, A., Crawford, R. J. M., Jarre, A., Quiñones, R. A., Shannon, L. J., & Verheye, H. M. (2000). Small pelagics in upwelling systems: Patterns of interaction and structural changes in “wasp-waist” ecosystems. *ICES Journal of Marine Science*, 57(3), 603–618. <https://doi.org/10.1006/jmsc.2000.0712>
- Day, R. H., Weingartner, T. J., Hopcroft, R. R., Aerts, L. A. M., Blanchard, A. L., Gall, A. E., Gallaway, B. J., Hannay, D. E., Holladay, B. A., Mathis, J. T., Norcross, B. L., Questel, J. M., & Wisdom, S. S. (2013). The offshore northeastern Chukchi Sea, Alaska: A complex high-latitude ecosystem. *Continental Shelf Research*, 67, 147–165. <https://doi.org/10.1016/j.csr.2013.02.002>
- Delpiani, S. M., Bruno, D. O., Vazquez, D. M., Llompарт, F., Delpiani, G. E., Fernández, D. A., Rosso, J. J., Mabragaña, E., & Díaz de Astarloa, J. M. (2020). Structure and distribution of fish assemblages at Burdwood Bank, the first Sub-Antarctic Marine Protected Area “Namuncurá” in Argentina (Southwestern Atlantic Ocean). *Polar Biology*, 43(11), 1783–1793. <https://doi.org/10.1007/s00300-020-02744-w>
- Di Mauro, R., Castillo, S., Pérez, A., Iachetti, C. M., Silva, L., Tomba, J. P., & Chiesa, I. L. (2022). Anthropogenic microfibers are highly abundant at the Burdwood Bank seamount, a protected sub-Antarctic environment in the Southwestern Atlantic Ocean. *Environmental Pollution*, 306, 119364. <https://doi.org/10.1016/j.envpol.2022.119364>
- Dormann, C. F., Fründ, J., & Schaefer, H. M. (2017). Identifying Causes of Patterns in Ecological Networks: Opportunities and Limitations. *Annual Review of Ecology, Evolution, and Systematics*, 48, 559–584. <https://doi.org/10.1146/annurev-ecolsys-110316-022928>
- Dunne, J. A., Williams, R. J., & Martinez, N. D. (2002a). Food-web structure and network theory: The role of connectance and size. *Proceedings of the National Academy of Sciences*, 99(20), 12917–12922. <https://doi.org/10.1073/pnas.192407699>
- Dunne, J. A., Williams, R. J., & Martinez, N. D. (2002b). Network structure and biodiversity loss in food webs: Robustness increases with connectance. *Ecology Letters*, 5(4), 558–567. <https://doi.org/10.1046/j.1461-0248.2002.00354.x>
- Dunne, J. A., Williams, R. J., & Martinez, N. D. (2002c). *Small Networks but not Small Worlds: Unique Aspects of Food Web Structure*. Santa Fe Institute.
- Eklöf, A., & Ebenman, B. (2006). Species loss and secondary extinctions in simple and complex model communities. *Journal of Animal Ecology*, 75(1), 239–246. <https://doi.org/10.1111/j.1365-2656.2006.01041.x>
- Estrada, E. (2007). Food webs robustness to biodiversity loss: The roles of connectance, expansibility and degree distribution. *Journal of Theoretical Biology*, 244(2), 296–307. <https://doi.org/10.1016/j.jtbi.2006.08.002>
- Fagan, W. F. (1997). Omnivory as a Stabilizing Feature of Natural Communities. *The American Naturalist*, 150(5), 554–567. <https://doi.org/10.1086/286081>
- Falabella, V. (2017). *Área Marina Protegida Namuncurá-Banco Burdwood. Contribuciones para la línea de base y el plan de manejo*.
- Fernández, M., Rodríguez-Ruiz, M., Gelcich, S., Hiriart-Bertrand, L., & Castilla, J. C. (2021). Advances and challenges in marine conservation in Chile: A regional and global comparison. *Aquatic Conservation: Marine and Freshwater Ecosystems*, 31(7), 1760–1771. <https://doi.org/10.1002/aqc.3570>
- Fioramonti, N. E., Ribeiro Guevara, S., Becker, Y. A., & Riccialdelli, L. (2022). Mercury transfer in coastal and oceanic food webs from the Southwest Atlantic Ocean. *Marine Pollution Bulletin*, 175, 113365. <https://doi.org/10.1016/j.marpolbul.2022.113365>
- Fischer, L., Covatti Ale, M., Deli Antoni, M., Díaz de Astarloa, J. M., & Delpiani, G. (2022). Feeding ecology of the longtail southern cod, *Patagonotothen ramsayi* (Regan, 1913) (Notothenioidei) in the

- Marine Protected Area Namuncurá-Burdwood Bank, Argentina. *Polar Biology*, 45(9), 1483–1494. <https://doi.org/10.1007/s00300-022-03082-9>
- Franco, B. C., Defeo, O., Piola, A. R., Barreiro, M., Yang, H., Ortega, L., Gianelli, I., Castello, J. P., Vera, C., Buratti, C., Pájaro, M., Pezzi, L. P., & Möller, O. O. (2020). Climate change impacts on the atmospheric circulation, ocean, and fisheries in the southwest South Atlantic Ocean: A review. *Climatic Change*, 162(4), 2359–2377. <https://doi.org/10.1007/s10584-020-02783-6>
- Freeman, L. C. (1978). Centrality in social networks conceptual clarification. *Social Networks*, 1(3), 215–239. [https://doi.org/10.1016/0378-8733\(78\)90021-7](https://doi.org/10.1016/0378-8733(78)90021-7)
- Funes, M., Saravia, L. A., Cordone, G., Iribarne, O. O., & Galván, D. E. (2022). Network analysis suggests changes in food web stability produced by bottom trawl fishery in Patagonia. *Scientific Reports*, 12(1), 10876. <https://doi.org/10.1038/s41598-022-14363-y>
- García Alonso, V. A., Brown, D., Martín, J., Pájaro, M., & Capitano, F. L. (2018). Seasonal patterns of Patagonian sprat *Sprattus fuegensis* early life stages in an open sea Sub-Antarctic Marine Protected Area. *Polar Biology*, 41(11), 2167–2179. <https://doi.org/10.1007/s00300-018-2352-z>
- Gauzens, B., Legendre, S., Lazzaro, X., & Lacroix, G. (2013). Food-web aggregation, methodological and functional issues. *Oikos*, 122(11), 1606–1615. <https://doi.org/10.1111/j.1600-0706.2013.00266.x>
- Glorioso, P. D., & Flather, R. A. (1995). A barotropic model of the currents off SE South America. *Journal of Geophysical Research: Oceans*, 100(C7), 13427–13440. <https://doi.org/10.1029/95JC00942>
- Godoy, O., Bartomeus, I., Rohr, R. P., & Saavedra, S. (2018). Towards the Integration of Niche and Network Theories. *Trends in Ecology & Evolution*, 33(4), 287–300. <https://doi.org/10.1016/j.tree.2018.01.007>
- Grilli, J., Rogers, T., & Allesina, S. (2016). Modularity and stability in ecological communities. *Nature Communications*, 7(1), 12031. <https://doi.org/10.1038/ncomms12031>
- Guerrero, R. A., Baldoni, A. G., & Benavides, H. R. (1999). *Oceanographic conditions at the southern end of the argentine continental slope*. <https://doi.org/http://10.0.64.26/handle/inidep/247>
- Guimerà, R., & Nunes Amaral, L. A. (2005). Functional cartography of complex metabolic networks. *Nature*, 433(7028), 895–900. <https://doi.org/10.1038/nature03288>
- Guinder, V. A., Malits, A., Ferronato, C., Krock, B., Garzón-Cardona, J., & Martínez, A. (2020). Microbial plankton configuration in the epipelagic realm from the Beagle Channel to the Burdwood Bank, a Marine Protected Area in Sub-Antarctic waters. *PLOS ONE*, 15(5), e0233156. <https://doi.org/10.1371/journal.pone.0233156>
- Heer, H., Streib, L., Schäfer, R. B., & Ruzika, S. (2020). Maximising the clustering coefficient of networks and the effects on habitat network robustness. *PLOS ONE*, 15(10), e0240940. <https://doi.org/10.1371/journal.pone.0240940>
- Hogg, O. T., Huvenne, V. A. I., Griffiths, H. J., Dorschel, B., & Linse, K. (2016). Landscape mapping at sub-Antarctic South Georgia provides a protocol for underpinning large-scale marine protected areas. *Scientific Reports*, 6(1), 33163. <https://doi.org/10.1038/srep33163>
- IUCN, & UNEP-WCMC. (2023). *The world database on protected areas (WDPA)*.
- Kadoya, T., Gellner, G., & McCann, K. S. (2018). Potential oscillators and keystone modules in food webs. *Ecology Letters*, 21(9), 1330–1340. <https://doi.org/10.1111/ele.13099>
- Kondoh, M., Kato, S., & Sakato, Y. (2010). Food webs are built up with nested subwebs. *Ecology*, 91(11), 3123–3130. <https://doi.org/10.1890/09-2219.1>
- Kortsch, S., Frelat, R., Pecuchet, L., Olivier, P., Putnis, I., Bonsdorff, E., Ojaveer, H., Jurgensone, I., Sträke, S., Rubene, G., Krüze, E., & Nordström, M. C. (2021). Disentangling temporal food web dynamics facilitates understanding of ecosystem functioning. *Journal of Animal Ecology*, 90(5), 1205–1216. <https://doi.org/10.1111/1365-2656.13447>
- Kortsch, S., Primicerio, R., Aschan, M., Lind, S., Dolgov, A. V., & Planque, B. (2019). Food-web structure varies along environmental gradients in a high-latitude marine ecosystem. *Ecography*, 42(2), 295–308. <https://doi.org/10.1111/ecog.03443>
- Kortsch, S., Primicerio, R., Fossheim, M., Dolgov, A. V., & Aschan, M. (2015). Climate change alters the structure of arctic marine food webs due to poleward shifts of boreal generalists. *Proceedings of the Royal Society B: Biological Sciences*, 282(1814), 20151546. <https://doi.org/10.1098/rspb.2015.1546>

- Lai, S.-M., Liu, W.-C., & Jordán, F. (2012). On the centrality and uniqueness of species from the network perspective. *Biology Letters*, 8(4), 570–573. <https://doi.org/10.1098/rsbl.2011.1167>
- López-Gappa, J., Liuzzi, M. G., & Zelaya, D. G. (2018). A new genus and species of cheilostome bryozoan associated with hermit crabs in the subantarctic Southwest Atlantic. *Polar Biology*, 41(4), 733–741. <https://doi.org/10.1007/s00300-017-2234-9>
- Marina, T. I., Saravia, L. A., Cordone, G., Salinas, V., Doyle, S. R., & Momo, F. R. (2018). Architecture of marine food webs: To be or not be a “small-world.” *PLOS ONE*, 13(5), e0198217. <https://doi.org/10.1371/journal.pone.0198217>
- Marina, T. I., Saravia, L. A., & Kortsch, S. (2023). *New insights into the Weddell Sea ecosystem applying a quantitative network approach*. EGUsphere preprint. <https://doi.org/10.5194/egusphere-2022-1518>
- Martinez, N. D. (1992). Constant Connectance in Community Food Webs. *The American Naturalist*, 139(6), 1208–1218. <https://doi.org/10.1086/285382>
- Martinez, N. D. (1991). Artifacts or Attributes? Effects of Resolution on the Little Rock Lake Food Web. *Ecological Monographs*, 61(4), 367–392. <https://doi.org/10.2307/2937047>
- Martinez, N. D. (1993). Effects of Resolution on Food Web Structure. *Oikos*, 66(3), 403–412. <https://doi.org/10.2307/3544934>
- Matano, R. P., Palma, E. D., & Combes, V. (2019). The Burdwood Bank Circulation. *Journal of Geophysical Research: Oceans*, 124(10), 6904–6926. <https://doi.org/10.1029/2019JC015001>
- Matusevich. (2022). *Chondrichthyan fauna from the Marine Protected Area Namuncurá at Burdwood Bank: Exploring egg nursery grounds*. <https://www.researchsquare.com>. <https://doi.org/10.21203/rs.3.rs-2247873/v1>
- May, R. (1973). *Stability and complexity in model ecosystems*. Princeton University Press.
- McCann, K., & Hastings, A. (1997). Re-evaluating the omnivory–stability relationship in food webs. *Proceedings of the Royal Society of London. Series B: Biological Sciences*, 264(1385), 1249–1254. <https://doi.org/10.1098/rspb.1997.0172>
- Montoya, J. M., & Solé, R. V. (2002). Small World Patterns in Food Webs. *Journal of Theoretical Biology*, 214(3), 405–412. <https://doi.org/10.1006/jtbi.2001.2460>
- Montoya, J. M., & Solé, R. V. (2003). Topological properties of food webs: From real data to community assembly models. *Oikos*, 102(3), 614–622. <https://doi.org/10.1034/j.1600-0706.2003.12031.x>
- Neutel, A.-M., Heesterbeek, J. A. P., & de Ruiter, P. C. (2002). Stability in Real Food Webs: Weak Links in Long Loops. *Science*, 296(5570), 1120–1123. <https://doi.org/10.1126/science.1068326>
- Newman, M. E. J. (2003). The Structure and Function of Complex Networks. *SIAM Review*, 45(2), 167–256. <https://doi.org/10.1137/S003614450342480>
- Nilsson, K. A., & McCann, K. S. (2016). Interaction strength revisited—clarifying the role of energy flux for food web stability. *Theoretical Ecology*, 9(1), 59–71. <https://doi.org/10.1007/s12080-015-0282-8>
- Padovani, L. N., Álvarez, N., & Farías, A. (2021). *Alimentación de la sardina fueguina (Sprattus fuegensis) en la región patagónica austral durante la época reproductiva*. INIDEP.
- Padovani, L. N., Viñas, M. D., Sánchez, F., & Mianzan, H. (2012). Amphipod-supported food web: Themisto gaudichaudii, a key food resource for fishes in the southern Patagonian Shelf. *Journal of Sea Research*, 67(1), 85–90. <https://doi.org/10.1016/j.seares.2011.10.007>
- Pascual, M., & Dunne, J. A. (2005). *Ecological Networks: Linking Structure to Dynamics in Food Webs*. Oxford University Press.
- Pastorino, G. (2019). A new deep water gastropod of the genus Parabuccinum (Neogastropoda: Buccinulidae) from southwestern Atlantic waters with new data on the distribution of all species. *Marine Biodiversity*, 49(2), 913–922. <https://doi.org/10.1007/s12526-018-0876-7>
- Pawar, S., Dell, A. I., & Van M. Savage. (2012). Dimensionality of consumer search space drives trophic interaction strengths. *Nature*, 486, 485. <https://doi.org/10.1038/nature11131>
- Piola, A. R., & Gordon, A. L. (1989). Intermediate waters in the southwest South Atlantic. *Deep Sea Research Part A. Oceanographic Research Papers*, 36(1), 1–16. [https://doi.org/10.1016/0198-0149\(89\)90015-0](https://doi.org/10.1016/0198-0149(89)90015-0)
- Piola, A., & Falabella, V. (2009). *El mar Patagónico*. Wildlife Conservation Society y Birdlife Internacional.
- Poisot, T., & Gravel, D. (2014). When is an ecological network complex? Connectance drives degree

- distribution and emerging network properties. *PeerJ*, 2, e251. <https://doi.org/10.7717/peerj.251>
- Raya Rey, A. (2022). *Food items of sea birds of the Marine Protected Area Namuncurá - Banco Burdwood*.
- Reta, R. (2014). *Oceanografía del Banco Burdwood: Estado Actual del Conocimiento y Perspectivas*. INIDEP.
- Riccialdelli, L. (2022). *Feeding items of marine mammals of the Marine Protected Area Namuncurá - Banco Burdwood*.
- Riccialdelli, L., Becker, Y. A., Fioramonti, N. E., Torres, M., Bruno, D. O., Rey, A. R., & Fernández, D. A. (2020). Trophic structure of southern marine ecosystems: A comparative isotopic analysis from the Beagle Channel to the oceanic Burdwood Bank area under a wasp-waist assumption. *Marine Ecology Progress Series*, 655, 1–27. <https://doi.org/10.3354/meps13524>
- Roberts, C. M., O’Leary, B. C., McCauley, D. J., Cury, P. M., Duarte, C. M., Lubchenco, J., Pauly, D., Sáenz-Arroyo, A., Sumaila, U. R., Wilson, R. W., Worm, B., & Castilla, J. C. (2017). Marine reserves can mitigate and promote adaptation to climate change. *Proceedings of the National Academy of Sciences*, 114(24), 6167–6175. <https://doi.org/10.1073/pnas.1701262114>
- Rodriguez, I. D., Marina, T. I., Schloss, I. R., & Saravia, L. A. (2022). Marine food webs are more complex but less stable in sub-Antarctic (Beagle Channel, Argentina) than in Antarctic (Potter Cove, Antarctic Peninsula) regions. *Marine Environmental Research*, 174, 105561. <https://doi.org/10.1016/j.marenvres.2022.105561>
- Sala, E., Lubchenco, J., Grorud-Colvert, K., Novelli, C., Roberts, C., & Sumaila, U. R. (2018). Assessing real progress towards effective ocean protection. *Marine Policy*, 91, 11–13. <https://doi.org/10.1016/j.marpol.2018.02.004>
- Santana, C. N. de, Rozenfeld, A. F., Marquet, P. A., & Duarte, C. M. (2013). Topological properties of polar food webs. *Marine Ecology Progress Series*, 474, 15–26. <https://doi.org/10.3354/meps10073>
- Saporiti, F., Bearhop, S., Vales, D. G., Silva, L., Zenteno, L., Tavares, M., Crespo, E. A., & Cardona, L. (2015). Latitudinal changes in the structure of marine food webs in the Southwestern Atlantic Ocean. *Marine Ecology Progress Series*, 538, 23–34. <https://doi.org/10.3354/meps11464>
- Saravia, L. A. (2022). *Multiweb: Ecological network analyses including multiplex networks*.
- Schejter, L., & Bremec, C. S. (2019). Stony corals (Anthozoa: Scleractinia) of Burdwood Bank and neighbouring areas, SW Atlantic Ocean. *Scientia Marina*, 83(3), 247–260. <https://doi.org/10.3989/scimar.04863.10A>
- Schejter, L., Genzano, G., Gaitán, E., Perez, C. D., & Bremec, C. S. (2020). Benthic communities in the Southwest Atlantic Ocean: Conservation value of animal forests at the Burdwood Bank slope. *Aquatic Conservation: Marine and Freshwater Ecosystems*, 30(3), 426–439. <https://doi.org/10.1002/aqc.3265>
- Schejter, L., Rimondino, C., Chiesa, I., Díaz de Astarloa, J. M., Doti, B., Elías, R., Escolar, M., Genzano, G., López-Gappa, J., Tatián, M., Zelaya, D. G., Cristobo, J., Perez, C. D., Cordeiro, R. T., & Bremec, C. S. (2016). Namuncurá Marine Protected Area: An oceanic hot spot of benthic biodiversity at Burdwood Bank, Argentina. *Polar Biology*, 39(12), 2373–2386. <https://doi.org/10.1007/s00300-016-1913-2>
- Schejter, & Albano. (2021). Benthic communities at the marine protected area Namuncurá/Burdwood bank, SW Atlantic Ocean: Detection of vulnerable marine ecosystems and contributions to the assessment of the rezoning process. *Polar Biology*, 44(10), 2023–2037. <https://doi.org/10.1007/s00300-021-02936-y>
- Scotti, M., & Jordán, F. (2010). Relationships between centrality indices and trophic levels in food webs. *Community Ecology*, 11(1), 59–67. <https://doi.org/10.1556/ComEc.11.2010.1.9>
- Sirito, M. (2019). *Fauna asociada a corales (Octocorallia) e hidroides (Hydrozoa) del Área Marina Protegida “Namuncurá” (Banco Burdwood) y zonas profundas adyacentes* [PhD thesis]. Universidad Nacional de Mar del Plata.
- Smith-Ramesh, L. M., Moore, A. C., & Schmitz, O. J. (2017). Global synthesis suggests that food web connectance correlates to invasion resistance. *Global Change Biology*, 23(2), 465–473. <https://doi.org/10.1111/gcb.13460>
- Solé, R. V., & Montoya, M. (2001). Complexity and fragility in ecological networks. *Proceedings of the Royal Society of London. Series B: Biological Sciences*, 268(1480), 2039–2045. <https://doi.org/10.1098/rspb.2001.1767>
- Spinelli, M. L., Malits, A., García Alonso, V. A., Martín, J., & Capitano, F. L. (2020). Spatial gradients

- of spring zooplankton assemblages at the open ocean sub-Antarctic Namuncurá Marine Protected Area/Burdwood Bank, SW Atlantic Ocean. *Journal of Marine Systems*, 210, 103398. <https://doi.org/10.1016/j.jmarsys.2020.103398>
- Stouffer, D. B., & Bascompte, J. (2010). Understanding food-web persistence from local to global scales. *Ecology Letters*, 13(2), 154–161. <https://doi.org/10.1111/j.1461-0248.2009.01407.x>
- Stouffer, D. B., Camacho, J., Guimerà, R., Ng, C. A., & Nunes Amaral, L. A. (2005). Quantitative Patterns in the Structure of Model and Empirical Food Webs. *Ecology*, 86(5), 1301–1311. <https://doi.org/10.1890/04-0957>
- Strogatz, S. H. (2001). Exploring complex networks. *Nature*, 410(6825), 268–276. <https://doi.org/10.1038/35065725>
- Team, R. C. (2022). *R: A Language and Environment for Statistical Computing*. R Foundation for Statistical Computing.
- Thébault, E., & Fontaine, C. (2010). Stability of Ecological Communities and the Architecture of Mutualistic and Trophic Networks. *Science*, 329(5993), 853–856. <https://doi.org/10.1126/science.1188321>
- Tilman, D. (1996). Biodiversity: Population Versus Ecosystem Stability. *Ecology*, 77(2), 350–363. <https://doi.org/10.2307/2265614>
- Trathan, P. N., Fielding, S., Hollyman, P. R., Murphy, E. J., Warwick-Evans, V., & Collins, M. A. (2021). Enhancing the ecosystem approach for the fishery for Antarctic krill within the complex, variable, and changing ecosystem at South Georgia. *ICES Journal of Marine Science*, 78(6), 2065–2081. <https://doi.org/10.1093/icesjms/fsab092>
- Troccoli, G. H., Aguilar, E., Martínez, P. A., & Belleggia, M. (2020). The diet of the Patagonian toothfish *Dissostichus eleginoides*, a deep-sea top predator off Southwest Atlantic Ocean. *Polar Biology*, 43(10), 1595–1604. <https://doi.org/10.1007/s00300-020-02730-2>
- Watts, D. J., & Strogatz, S. H. (1998). Collective dynamics of “small-world” networks. *Nature*, 393(6684), 440–442. <https://doi.org/10.1038/30918>
- Williams, R. J., Berlow, E. L., Dunne, J. A., Barabási, A.-L., & Martinez, N. D. (2002). Two degrees of separation in complex food webs. *Proceedings of the National Academy of Sciences*, 99(20), 12913–12916. <https://doi.org/10.1073/pnas.192448799>
- Woodward, G., Ebenman, B., Emmerson, M., Montoya, J. M., Olesen, J. M., Valido, A., & Warren, P. H. (2005). Body size in ecological networks. *Trends in Ecology & Evolution*, 20(7), 402–409. <https://doi.org/10.1016/j.tree.2005.04.005>