

2 *Submission of a perspectives piece to Integrative and Comparative Biology*

3 4 **Resolving the rules of robustness and** 5 **resilience in biology across scales**

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Abstract

Why do some biological systems and communities persist while others fail? Robustness, a system's stability, and resilience, the ability to return to a stable state, are key concepts that span multiple disciplines within and outside the biological sciences. Discovering and applying common rules that govern the robustness and resilience of biological systems is a critical step toward creating solutions for species survival in the face of climate change, as well as the for the ever-increasing need for food, health, and energy for human populations. We propose that network theory provides a framework for universal scalable mathematical models to describe robustness and resilience and the relationship between them, and hypothesize that resilience at lower organization levels contribute to robust systems. Insightful models of biological systems can be generated by quantifying the mechanisms of redundancy, diversity, and connectivity of networks, from biochemical processes to ecosystems. These models provide pathways towards understanding how evolvability can both contribute to and result from robustness and resilience under dynamic conditions. We now have an abundance of data from model and non-model systems and the technological and computational advances for studying complex systems. Several conceptual and policy advances will allow the research community to elucidate the rules of robustness and resilience. Conceptually, a common language and data structure that can be applied across levels of biological organization needs to be developed. Policy advances such as cross-disciplinary funding mechanisms, access to affordable computational capacity, and the integration of network theory and computer science within the standard biological science curriculum will provide the needed research environments. This new understanding of biological systems will allow us to derive ever more useful forecasts of biological behaviors and revolutionize the engineering of biological systems that can survive changing environments or disease, navigate the deepest oceans, or sustain life throughout the solar system.

1. The Problem

Life on Earth is shaped both by ancient and current events: no environment on Earth is fully invariant. Why particular biological systems, lineages, and communities persist while others fail is a question that spans multiple disciplines within and outside the biological sciences.

Understanding how all levels of biological organization respond to perturbation is central to decoding the rules of life. All living systems, including humans, face rapid changes in climate and landscapes that bring significant biotic (e.g., availability and phenology of prey and food items) and abiotic impacts (e.g., frequency/severity of floods, droughts, wildfires; temperature extremes) (König et al. 2020; Wintle et al. 2020). Revealing and applying common rules that govern the robustness and resilience of biological systems is an important and indispensable step toward finding solutions for preventing and curing diseases, for the ever-increasing need for food and energy, as well as for species survival. However, we lack an overarching understanding of the fundamental mechanisms that enable biological systems at various levels to appropriately respond to alterations in their environment and withstand or recover from perturbations. If researchers can decode universal rules of robustness and resilience, we can use these rules to predict how life on Earth will respond to rapidly changing conditions, to develop tools for ecosystem conservation, and to improve human conditions.

Shifting our conception of the natural world as many nested and interconnected networks (**see Fig. 1**) will transform how we view the minutiae and grandeur of biodiversity, while understanding how biological systems respond to changing conditions over time and space has a multiplicity of broader applications. How biological systems react with current, rapidly changing environmental conditions will affect every living thing on Earth (e.g., Hammerschlag et al. 2019). Outcomes of these efforts have consequences for an array of applications that will improve the quality of life for humans. The study of robustness and resilience at sub-cellular, physiological and tissue levels has medical implications; research in this area can set the stage

for advancements in disease biology and cancer treatments (Pienta et al. 2020; Rauter et al. 2020). The study of robustness and resilience can also be viewed through the lens of organismal biology and responses to environmental changes; outcomes from this area will influence conservation strategies for species in threatened ecosystems as well as providing a unique view of many potential and realized threats to biodiversity (e.g., Donelan et al. 2019). Finally, understanding resilient and robust biological systems can even facilitate improved design of energy storage/transport, urban transportation systems, and movement of resources across the globe (Tang et al. 2021, Wu et al. 2019, Ma et al. 2015).

2. Concepts of Robustness and Resilience

Processes related to robustness and resilience are studied by scientists across biological and physical disciplines, as well as social sciences, computer science and engineering (**Table 1**). At the same time, research into the responses to perturbations is often siloed at molecular, cellular, organismal, and ecological scales or within a discipline. Here we define **robustness** as the ability of a system to remain in or reach the same stable state despite diverse internal and external environments. Robustness underscores the ability of a biological system to maintain the original state even after encountering perturbations. In contrast, **resilience** (or resistance in ecological sciences) is the ability of a biological system to return to a previous state or establish a new state after significant perturbations. For example, a plant is robust and resilient if it grows normally across all different light conditions. A plant is resilient but not robust if it becomes dormant in the dark but restores growth rapidly once the desired light condition is met. A plant is robust but not resilient if it can grow under most light conditions but cannot handle switching between different light conditions. A plant is neither robust nor resilient if it only grows under one specific light condition and dies when that condition is not met. It is important to recognize in these definitions that one needs to carefully define variables into measurable characteristics or properties of a system (operationalize the system) maintaining stability, and identify what

processes or mechanisms are conferring the ability to return to the steady state in the context in which each of these terms are used (Brand and Jax 2007, Whitacre 2012, Nijhout et al 2019).

There are two overarching questions for examining robust and resilient systems: (1) how do biological systems maintain robustness and resilience in a continuously fluctuating and changing environment? (2) are there common rules that govern resilience and robustness across different scales of biological organization, from molecules to ecosystems? These questions can be addressed by examining biology as a multi-scale, nested, hierarchical system and considering how this complex system navigates changing conditions. We can then develop a holistic view of biological organization with more integrative approaches than the more discipline-specific or molecule-specific approach currently used. This approach will allow us to decode the complexity of biological systems and depict the hierarchical and network designs of biological systems more clearly. When we can deduce these rules, strategies, and mechanisms and any necessary variations, we will be better positioned to describe, model, and forecast resilience and robustness in systems across different levels. In addition, we will be able to create tools that allow us to “hack” biological systems, lending solutions to large problems involving disease, climate change, and threats to biodiversity.

Here we propose that concepts from **network theory** provide a framework for universal mathematical models to describe robustness and resilience and their relationship. First, we review properties of networks that confer robustness and resilience and provide examples of systems in which network theory has been applied (review the current state of knowledge). We then identify barriers that need to be overcome before scientific investigation can embrace network theory approaches, and describe ways a reintegration of biology and potential technological advances will allow us to overcome those barriers to advance our understanding of mechanisms underlying robustness in biological systems. Lastly, we suggest open questions

and research opportunities that remain to be addressed.

3. Review of network theory

3.1 How network theory relates to the study of robustness and resilience

The study of resilience and robustness is a transdisciplinary field that is amenable to a network science framework across different levels of biological organization. One network at a particular level of biological organization (e.g., within a cell) can become a node in a network at a different scale (e.g., across a cell population). Because networks are universal, scientists in all disciplines such as psychology, anthropology, social science, economics and engineering can benefit from a network-based, unified theory of biological robustness and resilience.

A network is defined as a collection of nodes and edges, which are abstract and universal to systems of all levels but can also be embodied with specific properties unique to individual systems. Each node in a network could be molecules, genes, cellular transduction pathways, individuals or genotypes in a population, species in a community or trophic levels in an ecosystem. For example, a biological community can be regarded as a network of interacting species within a geographic area. Within each species, different populations can have varying levels of interconnectivity, resulting in gene flow or isolation and constituting a dynamic network over time (Proulx et al. 2005; May 2006). Within each population, such as a colony of eusocial insects, individuals operate in a network to fulfill different functions of the colony (Wild et al. 2021). Within the organism, physiological regulatory networks operate to adjust functionality of multiple systems depending on environmental conditions (Cohen et al. 2012; Nijhout et al 2019). Within an embryo, different cell populations connect and operate in a developmental regulatory network to pattern the body plan of an organism (Levine and Davidson, 2005). Within each cell, functions are maintained by metabolic networks and cytoplasmic molecular networks, and in the nucleus, transcriptional networks are modulating cellular function (Gómez-Romero et al. 2020).

Borrowing from the mathematical theory of networks, we propose that key properties determining the robustness and resilience of biological systems at any organizational level are **redundancy**, **diversity**, and **connectivity** (see Fig. 2). Below we provide definitions and some examples of the relationship between these network properties and robustness and resilience.

Redundancy: Multiple nodes in a network could have the same or overlapping functions. If one or more nodes lose function, others can compensate. Similarly, there could be multiple routes of communication among nodes that confer the same functionality to a network. Redundancy is widely observed in developmental biology, where essential developmental events are often under the control of many genes that have similar or overlapping functions, and the expression of one gene compensates for the failure of another, up to a certain point. Redundancy is often used to explain how embryos tolerate developmental errors to result in the successful development of canalized body plans and morphogenesis (Lachowiec et al. 2018). Genetic knockout studies demonstrate the redundancy of many different molecular pathways (El-Brolosy and Stanier, 2017). Similarly, redundancy of neuroendocrine and genetic mechanisms regulating food intake are characteristics of a system regulating energy balance homeostasis (Schwarz et al. 2000). Lastly, food webs with overlapping ecological niches at different trophic levels are considered to confer stability to the system (Sanders et al. 2018).

Connectivity: We broadly define connectivity as the extent to which nodes communicate with each other, or specifically, the number and types of connections (edges) linking nodes in a network. Connectivity is a universal property of networks, but the specific connectivity depends on the structure of the network and mechanisms of communication and interaction. Networks can be described as distributed, decentralized, or centralized, each having

different patterns of connectivity. An important concept is the idea of ‘scale-free’ networks, describable by power law distributions of nodes with increasing connectivity degrees. Scale-free connectivity patterns are more likely to occur in biological systems than in informational or other technological systems (Broido and Clauset 2019), but the idea of universal scale-free network connectivity remains slightly contentious and requires more development (Holme, 2019). Connectivity plays a critical role in determining the robustness and resilience of a network. For example, distributed networks with high levels of edges connecting nodes confer stability, as demonstrated in the stability and persistence of metapopulations linked with migration (Hopf et al. 2019). During gastrulation, sheets of cells are robust against any “weak links” of individual cells in the population to allow for successful differentiation into germ layers. However, they are also resilient -- they can bend in response to external forces, while enabling them to still maintain cohesion and function (Davidson 2012). System feedback (i.e., negative feedback or positive feedback) is an essential part of control theory of dynamic systems. In the context of biological networks, feedback mechanisms are encoded in connectivity. Feedbacks in a network allow upstream nodes to send out signals to downstream nodes in response to signals they receive from the downstream nodes. A network with feedback connections will sense the state it is in, compare the current state to a setpoint or desired state, and then adjust its output to meet the desired state. In the scenario where the original set state cannot be met, a network with the appropriate connectivity could activate different feedbacks to break old connections, make new connections to establish a new stable state. Feedback mechanisms allow a network to correct or repair nodes and links that are perturbed or become dysfunctional under certain conditions. Common examples include negative feedbacks in predator-prey systems that result in population oscillations (Li et al. 2011), gene regulation systems that lead to constant gene expression outputs (Gjusvland et al. 2007, Hensel et al. 2012), or DNA proofreading and repair systems (Ashour & Mosammaparast 2021), and positive feedbacks

in excitable organism behaviors (O'Boyle et al. 2020) or memories in gene regulatory networks (Qiao et al. 2020, Fang et al., 2018).

Diversity: Diversity within a network can be regarded as the number, variations, and complexity of nodes of differential identities or functions. While the redundancy of nodes provides 'backups' that can compensate for potential failures in any one node, the diversity within a set of nodes provides variations in responses to heterogeneous challenges that can enable the system to function under different conditions. For example, genetic variations or differential gene expression states in microbial populations allow for the survival of resistant and persistent cells that could revive the entire population upon the termination of antibiotic treatment. High viral mutation rates create variants that escape host immune systems, resulting in robust viral infections (Drake 1993; Fitzsimmons et al. 2018). Genetic recombination and non-genetic memory (histone modifications, DNA methylation, prion-based inheritance mechanisms) are critical for adaptation to unexpected environment changes. They provide the molecular ingredients for a heritable response, fixing these changes in phenotype within a population (Payne and Wagner 2019). Animals in unpredictable or highly variable environments produce eggs of various sizes or offspring with diverse phenotypes (or genotypes) so that at least some of the offspring are suited for the environment (bet-hedging) (Olofsson et al., 2009; Morrongiello et al., 2012).

Communities with more diverse species composition and larger population sizes are more stable and resistant to invasive species than those with smaller sizes (Hopf et al. 2019).

We propose that once the redundancy, connectivity, and diversity of networks at any level of biological organization are understood, common rules of robustness and resilience will emerge. Each level of biological organization is conceptualized as a network consisting of nodes and edges, with the emergent collective behavior of the network as a node

for the network of the next higher level of organization. With this framework, we can ask interesting questions such as how robustness and resilience are related across scales; i.e., are there microscale to macroscale network dynamics that work together to facilitate robustness?

One important hypothesis that can be tested is whether resilience at lower levels of organization contributes to increasing robustness at higher spatial and biological scales. For example, ecosystem robustness may be maintained when some populations thrive while others decline during an environment change. Thus the output, e.g. survival or appropriate development of a species or an organism, may be robust to environmental insult by virtue of the resilience of underlying interaction networks (see **Box 1**).

3.2 Concept of Evolving Networks

While a network with appropriate levels of redundancy, diversity and connectivity could confer robustness and resilience under set conditions, these networks must also be able to adjust under fluctuating and changing environments and evolve over time. Here, we broadly define **evolvability** as the ability of the system to change functions in response to significant perturbations, either by maintaining the original stable state but with enhanced stability, or by moving to a new stable state with changed properties. An evolved network may have broken or established new connections, or connections that have increased/decreased in strength, or direction relative to the remaining connections. An evolvable network can provide the potential to sustain individual and/or population survival in hostile environments, such as what was shown in signaling networks (e.g., Pimpinelli & Piacentini 2020). This concept is commonly referred to as physiological acclimation, phenotypic plasticity, or evolutionary adaptation depending on the level of biological organization. For an example of how robustness, resilience and evolvability play out in metabolic networks in living cells, see **Fig. 3, Box 2**.

Evolvability can both contribute to and result from robustness and resilience under dynamic conditions. Variation in ecological niches can also promote the evolution of organismal specialization (Cordeiro et al. 2020). Organismal specialization can involve a gain or loss of a response to particular environmental conditions, depending on the dynamism of the environmental stressor (e.g., Saiz et al. 2021). The frequency, magnitude and type of environmental changes experienced by a lineage contribute to the evolution of robustness-supporting networks. The resilience of a system to environmental change is associated with the introduction of novelties into it, or the systems' adaptive capacity (Allen and Holling 2010). However, ecological, physiological or evolutionary constraints may limit a system's response during exposure to extreme conditions that are significantly different than those previously encountered (Dutta et al. 2021). Even so, there may be biological factors that contribute to a species' population robustness even in the face of rapid human-driven changes (e.g. Reid et al. 2016).

Linking the changes that promote robustness or resilience in a particular environment to a single gene or small set of genes (or a small set of organisms) may artificially limit our understanding of the nature of these emerging properties. Evolutionary history shapes responses to environmental conditions; understanding these changes in broader terms that incorporate network changes or community changes is important. It is also important to note that phenotypic plasticity within a generation that can be transmitted to the next generation via epigenetic or non-genetic changes contribute to gain or loss of robustness in an organism (Payne & Wagner 2019). Regardless of whether its origin is genetic or epigenetic, study of flexible networks that occur at different levels of organization is needed to understand generalizable strategies. These strategies can then be modeled across scales to show how robustness or resilience at one level relates to those at another. Evolutionary biologists can help us understand how stability and resilience of systems change in response to selection different pressures or how diverse

mechanisms create systems that confer stability and control.

3.5 Technological and computational advances enabling a network theory paradigm shift

Now is an opportune time to establish a framework that enables the modeling of complex systems across scales to understand biological robustness and resilience. We have access to many state-of-the-art, enabling technologies that can generate expansive molecular-level data sets, including all of the ‘omics” at the molecular levels. Population-wide and individual behaviors at the large can be recorded remotely and analyzed in near real-time, through large-scale phenomics systems or satellite images. Most importantly, we are developing better tools for data acquisition, analysis, and transfer that will allow us to bridge data from atomic to stellar scales. We now possess technologies to manipulate, observe, analyze and synthesize our understanding of model and non-model systems in controlled lab environments as well as in the field, even up to the global scale. Much is now known about the mechanisms of life, including the biochemical reactions of information and energy processing within microbial cells, programs that define the development and evolution of multicellular organisms from plants to humans, and interactions among diverse life forms that contribute to ecosystem emergence and dynamics.

At the molecular scale, we can access large quantities of genomic and transcriptomic information in near real time across phenotypes, populations, species, and lineages through NGS, single-cell sequencing and RNA-seq approaches (Estermann et al. 2020; Iacono et al. 2019). Advanced mass spectroscopic techniques provide quantitative proteomic and metabolomic analyses to address a wide range of biological questions. Cryo-electron microscopy and tomography can visualize structures of macromolecular complexes in native or near native environments with atomic resolutions. Super-resolution and single-molecule imaging push the detection of molecules and cellular structures in live cells beyond the diffraction limit of light microscopy. We also possess incredible powers in manipulating organisms through

genome editing and targeted perturbations. At the organismal level, it is feasible to build synthetic cells and grow organoids that recapitulate essential features of life, and now even sustain mammalian development in vitro (ZhuAguilera-Castrejon et al. 2021). At the population level, the most advanced tracking technologies are able to monitor the dynamics of large populations of animals and changes in ecosystems (Barnas et al. 2019). Various social media outlets offer new platforms to gather and disseminate information at the societal level. Growing computational and mathematical power, coupled with mechanistic modeling, machine learning and artificial intelligence algorithms, have the potential to describe systems and predicate outcomes at different scales, across different levels of biological organization (molecules to ecosystems), spanning broad time scales (nanoseconds, seconds, minutes, hours), or by some metric of complexity (e.g., reaction, pathway, network, hairball). We have an abundance of in-depth data not only from model systems, but also from diverse, non-lab adapted systems. If coalesced into standardized, user-accessible databases (as exemplified by Pangeo for geoscientific data (<http://pangeo.io>), these data can be used to systems and examine strategies universal to different scales. The substantial amount of historical genetic and ecological data can be integrated with current data to develop algorithms of hindcasts to forecast robustness and resilience of systems.

4. Barriers to progress: Challenges to the adoption of a network theory framework

While there are many advances that make this paradigm shift possible at this time, there are also many barriers that need to be overcome before a wide range of scientists are able to embrace applying network theory for robustness and resilience across all biological scales. As described in more detail below, engineers, computer scientists, and biologists in different research communities lack a common language for describing the meaning of robustness or resilience across different levels of biological organization, although the field of systems biology has adapted many of the ideas of network theory for some biological systems, typically focused

at the molecular, cellular, and tissue levels (e.g., Goldman et al. 2015). In addition, there are many institutional and structural barriers to be overcome. For a unified theory of robustness and resilience to emerge, meaningful incentives to promote collaborative research must be implemented, and traditional divisional barriers must be bridged.

4.1 Language: There is a lack of a common language for describing robustness or resilience across different levels of biological organization (see Table 1). Terms like “resilience” and “robustness” depend on context (molecular, cellular, multi-cellular, population) and differ depending on scientific training or field (math/systems/engineering versus molecular/cell /biology/ecology). Developing a common language across fields provides an opportunity to identify unifying threads across biological levels and across scientific fields (e.g., Davies 2018). Different fields and training have hypotheses and constructed models of “resilience” or “robustness” for certain systems, but scientists outside the field (or approach) may struggle to adopt these models to novel areas, or they may toil to adapt powerful methods of another field to test hypotheses in their own. Common terms will allow scientists to find relevant concepts and empirical data in other fields through literature searches and increase opportunities to collaborate across fields. We propose that the language of network theory (see above) could take a first step toward unifying how researchers from diverse fields conceptualize and communicate information about complex systems.

Another general problem when integrating information across subdisciplines in the biological sciences is the use of jargon, such that the same phenomena are studied independently, preventing the integration of these disciplines. For example, we have amazing tools for searching primary literature that combine sources of information across diverse scientific disciplines (e.g., Web of Science), but literature searches are restricted to the terms used. Unless this terminology is standardized, or “smart” searches that translate those terms into

others that are known to vary across fields are utilized, relevant information will be forever segregated in the minds of researchers of different fields. For example, evolutionary biologists interested in “transgenerational plasticity” may also be interested in “developmental programming” studies in the biomedical literature or “carry-over effects” in the ecological literature. As shown in Table 1, there are terms of similar meaning related to the concepts of robustness and resilience across fields, although in each case there are specific nuances, connotations or usages that differ among terms. Creating interdisciplinary educational programming will enhance this merging of language and terminology so that discipline-specific jargon will be eased.

4.2 Lack of technology and experimental testing: A process that is altered and returns to a previous state (resilient) may exhibit a robust response at a higher level of temporal, spatial, or organismal integration. Measures need to be relevant both to the physical and temporal scale of perturbation and must subsequently transmit a signal associated with this perturbation to adjacent levels. Despite access to huge sets of molecular, behavior, and population data, the current state-of-the-art techniques generally lack the ability to integrate information across length scales and time scales; how networks are defined and interactions quantified requires more development, including new technologies to measure how networks respond to perturbations across scales. It is also unclear which experimental systems best serve as case studies in which this technology can be tested and optimized.

4.3 Logistics: Even when there is a desire to collaborate across fields, finding potential colleagues with similar interests and willingness to collaborate can be challenging. Most scientific conferences are field-specific; thus, it is challenging for scientists to find opportunities to meet and discuss ideas with others in different fields. Even after finding a collaborator, there are logistical hurdles in carrying out a project such as grant administration and international

access to sensitive data. In addition, there are institutional barriers that prevent scientists from gaining access to the physical infrastructure and tools needed to study transdisciplinary robustness and resilience across scales. Often funding opportunities and financial incentives that promote the formation of novel transdisciplinary collaborations are limited. When inter- or transdisciplinary proposals are submitted to traditional funding mechanisms, the small pool of reviewers who have discipline-specific expertise but also appreciate the novelty of transdisciplinary collaborations could limit the funding of such proposals.

5. Strategies to overcome barriers to progress

5.1 Reintegration of biology: Robustness is a concept that crosses many levels of biological organization; a fuller understanding of this characteristic requires the integration of many different disciplines so that a common language emerges. A multidisciplinary team approach would eliminate the inherent scale and model bias, allowing for broader perspectives into the rules of life. We therefore need platforms for researchers who are interested in understanding robustness and resilience from biophysics, mathematics, molecular biology, physiology, population genetics, and ecosystem biology, etc. who do not otherwise interact to brainstorm ideas. This could be done in workshops resulting in new collaborations and possible research coordination networks. Funding mechanisms that promote the formation of new multidisciplinary research teams will also broaden participation of researchers from different backgrounds and institutional types (e.g., primarily teaching institutions, medical schools, and research-intensive universities). Funding agencies such as the National Science Foundation have acknowledged that they can play a major role in promoting cross-disciplinary training of a new generation of scientists by changing funding schemes, paradigms and training programs. These changes will promote cross-disciplinary training of a new generation of scientists who have the skills to discover and describe the important overarching questions of life on Earth. For example, we might harness existing big data and integrate insights from available models of community and

population dynamics that are successfully used for metabolism, viruses, microbiomes and ecosystems (Mathias et al. 2017) to construct mathematical models to elucidate common rules underlying resilience and robustness.

We can also leverage our understanding of the evolution to advance our understanding of robust and resilient systems. With large-scale, multidimensional networks, comparative analysis of network interactions over time will allow the role of evolutionary pressure to be examined in biological robustness. This analysis would move beyond our current reliance on gene or protein networks, to incorporate communications between nearest neighbors (intra- and inter-habitat) and entire communities over time. Then specific nodes or network strategies to overcome challenges and promote robustness that recur over time could then be used to re-engineer robust and scalable networks from gene to community levels.

5.2 Development of new tools: To overcome technological barriers, we need to develop suitable metrics and tools to measure robustness and resilience (or lack thereof) across space and time scales. Ideally such a tool would measure or provide a measure of the response of a system at one scale and seamlessly measure the propagation of the response across multiple scales. For example, noise in the production of RNA during the activation of gene expression can contribute to cellular heterogeneity, resulting in a robust response to perturbations across a population of cells. It is unclear how heterogeneity that is generated at the cellular level affects higher-order processes. Real-time readouts would enable us to capture events that happen throughout the life of the organism. One method of obtaining this type of data would be using optical methods, requiring the development of stable reporters that are not susceptible to bleaching or degradation biases. Optical or other readouts of behavior, neural status, and molecular reporters could then be integrated across scales to provide networks in context. Eventually, to support the development of full molecular networks in context, real-time molecular

sampling of a freely-responding (super)-organism will be necessary.

At the most ambitious level, advanced technologies would be deployed to generate and analyze network data in real time. These technologies might include real-time analysis of transcriptomes, proteomes, metabolomes, neural readouts, and behavior in an environmental context. Not all of these technologies are ready, but many are very close, enhanced by the current growth in computational power (data analytics), real-time sequencing, and computer vision. Assuming no limitations, we could have all the experimental data possible to build dynamic networks. This will require integrated hypotheses that probe networks and additional strategies to address evolutionary selection, particularly the survival of an individual and a population.

To move toward this integrative network-based analysis of robustness, in the next few years we would need to implement model test systems across multiple life scales with scientific teams to develop testable hypotheses that validated network development. understand In addition to the development of new sensing and measurement technologies, we need to develop new data analytics and computational methods to transform current data streams into multidimensional networks. Enormous, affordable computational capacity is needed in hardware for storage, fast CPU/GPU, parallel processing, and freely available open software. With these developments, we could not only test network robustness but analyze redundancy. Exploring redundancy and determining essential nodes for stability and robustness of networks at multiple levels would provide essential insight into robustness that has been inaccessible due to the lack of global monitoring systems capable of collecting data at sufficient scales. Infrastructure will also need to be created to host these databases, enable user contributions and make databases searchable and available to the public, much like NCBI databases.

5.3 Education: In order to realize a reintegration of biology and generate the workforce needed to create the technologies needed to advance network-level study of biological systems, we need to reform science and math education. Critically, science education from K-12 through the post-doctoral level should be designed to foster problem-based scientific thinking not siloed by discipline. Integration of knowledge from different scientific disciplines needs to become a common way of thinking for the next generation of scientists and innovators. In addition, curricula should include requirements that emphasize analytical reasoning and quantitative skills. Network theory and computer science courses could be included as standard biological science curricula in addition to algebra, calculus, and statistics. It is important to impress upon students how mathematical tools applied in modeling and engineering fields can be employed to derive potential solutions to important societal problems (NRC, 2009, **see Box 3**).

5.4 Reorganization of institutional funding mechanisms and infrastructure: To overcome logistical barriers to advancing research on robustness and resilience, it is important for both funding agencies and research institutions to facilitate and incentivize interdisciplinary interactions among scientists. This can be best accomplished with specialized funding mechanisms that call for such interdisciplinary teams, such as the joint National Institutes of Health and National Science Foundation Ecology and Evolution of Infectious Disease mechanism, and the newly established NSF Integrative Research in Biology (IntBIO) and the Biology Integration Institutes mechanisms. However, it is still a challenge for researchers to establish relationships with collaborators, especially biomathematicians and bioinformaticians with allied interests and expertise. Within research institutions, increasing internal funding opportunities to encourage interdisciplinary collaborations, cluster hiring around interdisciplinary research themes, and encouraging young investigators to engage in collaborative research through established (or new) institutional interdisciplinary or transdisciplinary centers could increase research in robustness and resilience.

6. Open questions and research opportunities

Studying biological systems within a unifying framework as living and interacting networks will allow us to address some of the most important biological and social questions of our time (see **Table 2**). Understanding the underlying principles of biological robustness and resilience will allow us to model and anticipate consequences of environmental changes across scales and enable controlling of biological systems for most beneficial outcomes. For example, it is desirable to destabilize the state of persistent neural seizures resulting from epilepsy or neurotoxin exposure, in which neural signals are persistently entrained. Similarly, we may want to model or forecast consequences of anthropogenic effects such as an oil spill and develop ways to return ecosystems to its healthy state. Models of robustness and resilience can inform methods to stabilize or destabilize agri- and aquaculture, improving sustainability or reducing the impact of invasive species. They could also provide insight into disease development and progression, either in natural or modified systems. In a world with a rapidly changing climate, such interventions may be essential for organismal survival and to prevent a sixth extinction but will require significant ethical restraint in their applications.

Collaboration among researchers from experimental, mathematical, computational and engineering fields will allow the application of developed models to improve the health of the ecosystem and human lives. For example, useful experimental datasets, mathematical models, and computational tools for validating and understanding behaviors of complex systems may be generated. New software incorporating improved parameter definitions and modeling techniques could facilitate the investigation and understanding of intra- and inter-level connections of complex biological systems. Synthetic datasets with standardized format could also result from this research to allow downstream applications for other multiscale studies.

A greater understanding of the theoretical mechanisms of robust or resilient networks will also help develop better computation tools and more reliable artificial intelligence (AI) algorithms. By identifying essential networks and nodes that promote robustness, we can implement them to perform complex AI-driven tasks such as self-driving vehicles, rover navigation undersea, or on Mars, or exploration of oceans and moons. Robustness and resilience theory will provide new algorithms for implementing complex tasks in constantly changing environments. Understanding the role of robustness in evolution will also enable artificial systems to learn how to rapidly navigate new and complex environmental contexts.

Finding common rules of robustness and resilience across scales in natural systems will accelerate new discoveries and progress on elucidating the rules of life on Earth, transform the way we understand biological systems and revolutionize synthetic biology. We will begin elucidating design and engineering principles of living systems and use them to deploy stable and viable synthetic systems. As biological systems of different organization levels are interconnected across scales, we may be able to forecast how changes at one organization level affect the other levels, contributing to a holistic understanding of all biological systems.

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Literature Cited

- Adger, W. N. 2000. Social and ecological resilience: are they related? *Progress in Human Geography* **24**(3):347-364. doi.org/10.1191/030913200701540465.
- Aguilera-Castrejon, A., Oldak, B., Shani, T. et al. 2021. *Ex utero* mouse embryogenesis from pre-gastrulation to late organogenesis. *Nature* doi:10.1038/s41586-021-03416-3.
- Allen, C.R., Holling, C.S. 2010. Novelty, adaptive capacity and resilience. *Ecology and Society* **15**(3): 24.
- Ashour, M.E. and Mosammaparast, N. 2021. Mechanisms of damage tolerance and repair during DNA replication. *Nucleic Acids Research*. doi: 10.1093/nar/gkab101.
- Baez, A. and J. Shiloach. 2014. Effect of elevated concentration on bacteria, yeasts and cells propagated for production of biological compounds. *Microbial Cell Factories* **13**: 181. doi.org/10.1186/s12934-014-0181-5.
- Barkai, N., Leibler, S. 1997. Robustness in simple biochemical networks. *Nature* **387**, 913–917. https://doi.org/10.1038/43199.
- Barnas, A.F., Darby, B.J., Vandeberg, G.S., Rockwell, R.F., Ellis-Felege, S.N. 2019. A comparison of drone imagery and ground-based methods for estimating the extent of habitat destruction by lesser snow geese (*Anser caerulescens caerulescens*) in La Pérouse Bay. *PLoS ONE* **14**(8): e0217049. doi: 10.1371/journal.pone.0217049.
- Basan, M., S. Hui, H. Okano, Z. Zhang, Y. Shen, J.R. Williamson and T. Hwa. 2015. Overflow metabolism in *E. coli* results from efficient proteome allocation. *Nature* **528**(7580): 99-104. doi:10.1038/nature15765.
- Brand, F. S., and K. Jax. 2007. Focusing the meaning(s) of resilience: resilience as a descriptive concept and a boundary object. *Ecology and Society* **12**(1): 23. doi:10.5751/ES-02029-120123.
- Broddrick, J. T., Rubin, B. E., Welkie, D. G., Du, N., Mih, N., Diamond, S., Lee, J. J., Golden, S. S., and Palsson, B. O. 2016. Unique attributes of cyanobacterial metabolism revealed by improved genome-scale metabolic modeling and essential gene analysis. *Proc. Natl. Acad. Sci. USA* **113**(51): E8344-E8353. doi:10.1073/pnas.1613446113.
- Broido AD, Clauset A. Scale-free networks are rare. *Nat Commun*. 2019 Mar 4;10(1):1017. doi: 10.1038/s41467-019-08746-5.
- Cannon WB. *The Wisdom of the Body*. New York: W. W. Norton & Company; 1932. pp. 177–201.
- Carlson, J.M., Doyle, J. 2002. Complexity and robustness. *PNAS* **99**: 2538-2545. doi.org/10.1073/pnas.012582499.
- Cohen, A.A., L.B. Martin, J.C. Wingfield, S.R. McWilliams, J.A. Dunne. 2012. Physiological regulatory networks: ecological roles and evolutionary constraints. *Trends in Ecology & Evolution* **27**(8): 429-435. doi: 10.1016/j.tree.2012.04.008.
- Cordeiro, J., de Oliveira, J.H.F, Schmit, H.J., Vizentin-Bugoni, J. 2020. High niche partitioning promotes highly specialized, modular and non-nested florivore-plant networks across spatial scales and reveals drivers of diversification. *Oikos* **619**-629. doi: 10.1111/oik.06866.
- Cumming, G. S., G. Barnes, S. Perz, M. Schmink, K. E. Sieving, J. Southworth, M. Binford, R. D. Holt, C. Stickler and T. Van Holt. 2005. An exploratory framework for the empirical measurement of resilience. *Ecosystems* **8**:975-987. doi.org/10.1007/s10021-005-0129-z.
- Davidson, L.A. 2012. Epithelial machines that shape the embryo. *Trends Cell Biol* **22**:82-87. doi: 10.1016/j.tcb.2011.10.005.

- Davies, KJA. 2016. Adaptive homeostasis. *Mol Aspects Med.* 49: 1–7.
doi:10.1016/j.mam.2016.04.007.
- Davies, P. 2018. *Demon in the machine: How hidden webs of information are solving the mystery of life.* University of Chicago Press, Chicago, IL.
doi: 10.7208/chicago/9780226669847.001.0001
- De Marco L, Sassera D, Epis S, Mastrantonio V, Ferrari M, Ricci I, Comandatore F, Bandi C, Porretta D, Urbanelli S. 2017. The choreography of the chemical defense response to insecticide stress: insights into the *Anopheles stephensi* transcriptome using RNA-Seq. *Sci Rep.* 7:41312. doi: 10.1038/srep41312.
- Dodds, P.S., Watts, D.J., Sabel, C.F. 2003. Information exchange and the robustness of organizational networks. *PNAS* 100 (21) 12516-12521. doi.org/10.1073/pnas.1534702100.
- Donelan, S. C., J. K. Hellman, A. M. Bell, B. Luttbeg, J. L. Orrock, M. J. Sheriff, and A. Sih. 2019. Transgenerational plasticity in human-altered environments. *Trends in Ecology and Evolution* 35(2): P115-124. doi.org/10.1016/j.tree.2019.09.003.
- Drake, J.W. 1993. Rates of spontaneous mutation among RNA viruses. *Proc. Natl. Acad. Sci. USA* 90: 4171-4175. doi:10.1073/pnas.90.9.4171.
- Dutta, A., Hartmann, F.E., Francisco, C.S., McDonald, B.A., Croll, D. 2021. Mapping the adaptive landscape of a major agricultural pathogen reveals evolutionary constraints across heterogeneous environments. *The ISME Journal*. doi: 10.1038/s41396-020-00859-w.
- Egeland, R., Carlson, E., Sroufe, L.A. 1993. Resilience as a process. *Development and Psychopathology* 5: 517-528. doi.org/10.1017/S0954579400006131.
- Elton, C. S. *The Ecology of Invasions by Animals and Plants* (Methuen, London, 1958).
- Estermann MA, Smith CA. 2020. Applying Single-Cell Analysis to Gonadogenesis and DSDs (Disorders/Differences of Sex Development). *Int J Mol Sci.* 21(18):6614. doi: 10.3390/ijms21186614.
- Falk, D.A., Watts, A.C., Thode, A.E. 2019. Scaling Ecological Resilience. *Ecol. Evol.* 7:275. doi.org/10.3389/fevo.2019.00275.
- Fitzsimmons, W.J., R.J. Woods, J.T. McCrone, A. Woodman, J.J. Arnold, M. Yennawar, R. Evans, C.E. Cameron, A.S. Luring. 2018. A speed-fidelity trade-off determines the mutation rate and virulence of an RNA virus. *PLOS Biology* 16(6): e2006459. doi: 10.1371/journal.pbio.2006459.
- Fletcher, D., Sarkar, M. 2013. Psychological resilience: A review and critique of definitions, concepts, and theory. *European Psychologist*, 18(1), 12-23. https://doi.org/10.1027/1016-9040/a000124.
- Gjuvslund, A.B., Plahte, E., Omholt, S.W. 2007. Threshold-dominant regulation hides genetic variation in gene expression networks. *BMC Systems Biology* 1:57. 0.1186/1752-0509-1-57.
- Goldman, A.W., Burmeister, Y., Cesnulevicius, K., Herbert, M., Kane, M., Lescheid, D., McCaffrey, T., Schultz, M., Seilheimer, B., Smit, A., St.Laurent, G., and Berman, B. 2015. Bioregulatory Systems medicine: an innovative approach to integrating the science of molecular networks, inflammation, and systems biology with the patient's autoregulatory capacity? *Front. Physiol.* 6:225. doi: 10.3389/fphys.2015.00225.
- Goldstone JV, Hamdoun A, Cole BJ, Howard-Ashby M, Nebert DW, Scally M, Dean M, Epel D, Hahn ME, Stegeman JJ. 2006. The chemical defense response: environmental sensing and response genes in the *Strongylocentrotus purpuratus* genome. *Dev Biol.* 300(1):366-84. doi: 10.1016/j.ydbio.2006.08.066.
- Gómez-Romero, L., López-Reyes, K., Hernández-Lemus, E. 2020. The large scale structure of

human metabolism reveals resilience via extensive signaling crosstalk. *Front. Physiol.* 11: 588012. doi: 10.3389/fphys.2020.588012.

Hammerschlag, N., O. J. Schmitz, A. S. Flecker, K. D. Lafferty, A. Sih, T. B. Atwood, A. J. Gallagher, D. J. Irschick, R. Skubel, and D. J. Cooke. 2019. Ecosystem function and services of aquatic predators in the Anthropocene. *Trends in Ecology and Evolution* 34(4): 360-383. doi.org/10.1016/j.tree.2019.01.005.

Hari, A., and Lobo, D. 2020. Fluxer: a web application to compute, analyze and visualize genome-scale metabolic flux networks. *Nucleic acids research* 48, W427-W435.

Hensel, Z., Feng, H., Han, B., Hatem, C., Xiao, J. 2012 Stochastic expression dynamics of a transcription factor revealed by single-molecule noise analysis. *Nat. Struct. Mol. Biol.* 19: 797-802. doi: 10.1038/nsmb.2336.

Holling, C. S. 1973. Resilience and stability of ecological systems. *Ann. Rev. Ecol. Sys.* 4, 1–23. doi: 10.1146/annurev.es.04.110173.000245

Holme P. 2019. Rare and everywhere: Perspectives on scale-free networks. *Nat Commun.* 10(1):1016. doi: 10.1038/s41467-019-09038-8.

Hopf, J.K., G.P. Jones, D.H. Williamson, and S.R. Connolly. 2019. Marine reserves stabilize fish populations and fisheries yields in distributed coral reef systems. *Ecological Applications* 29(5): e01905. doi.org/10.1002/eap.1905.

Iacono G, Massoni-Badosa R, Heyn H. 2019. Single-cell transcriptomics unveils gene regulatory network plasticity. *Genome Biol.* 20(1):110. doi: 10.1186/s13059-019-1713-4.

Kim, P.J., D.Y. Lee, T.Y. Kim, K.H. Lee, H. Jeong, S.Y. Lee, S. Park. 2007. Metabolite essentiality elucidates robustness of *Escherichia coli* metabolism. *Proc. Natl. Acad. Sci. USA* 104(34): 13638-13642. doi: 10.1073/pnas.0703262104.

Kitano, H. 2004. Biological robustness. *Nat Rev Genet* 5, 826–837. <https://doi.org/10.1038/nrg1471>

König, H.J., Kiffner, C., Kramer-Schadt, S., Fürst, C., Keuling, O., Ford, A.T. 2020. Human-wildlife coexistence in a changing world. *Cons. Biol.* 34(4): 786-794. doi: 10.1111/cobi.13513.

La Rosa, R., H.K. Johansen, S. Molin. 2018. Convergent metabolic specialization through distinct evolutionary paths in *Pseudomonas aeruginosa*. *mBio* 9: e00269-18. doi.org 10.1128/mBio.00269-18.

Lacohwiec, J., G.A. Mason, K. Schultz, and C. Queitsch. 2018. Redundancy, feedback and robustness in *Arabidopsis thaliana* BZR/BEH gene family. *Front. Genet.* 9: article 523. doi:10.3389/fgene.2018.00523.

Lafuente, E., Beldade, P. 2019. Genomics of developmental plasticity in animals. *Front. Genet.* <https://doi.org/10.3389/fgene.2019.00720>

Lesne, A. 2008. Robustness: confronting lessons from physics and biology. *Biological Reviews* 83(4): 509-532. doi.org/10.1111/j.1469-185X.2008.00052.x

Li, C., Wang, E., Wang, J. 2011. Potential landscape and probabilistic flux of a predator prey network. *PLoS ONE* 6(3): e17888. doi: 10.1371/ journal.pone.0017888.

Liang, F. and P. Lindblad. 2016. Effects of overexpressing photosynthetic carbon flux control enzymes in the cyanobacterium *Synechocystis* PCC 6803. *Metabolic Engineering* 38: 56-64. doi:10.1016/ j.ymben.2016.06.005.

Ma, X., Yu, H., Wang, Y., Wang, Y. 2015. Large-scale transportation network congestion revolution prediction using deep learning theory. *PLoS ONE* 10(3): 0119044. doi:

10.1371/journal.pone.0119044.

Mareya CR, Tugizimana F, Piater LA, Madala NE, Steenkamp PA, Dubery IA. 2019. Untargeted Metabolomics Reveal Defense-Related Metabolic Reprogramming in *Sorghum bicolor* against Infection by *Burkholderia andropogonis*. *Metabolites*. 9(1):8.

Mathias, M., M.D. Flavia, L.G. Rafael, P.P. Coltri, S.I. Perez, D.R. Barneche, D.Y.C. Brandt, K. Nunes, F.G. Daura-Jorge, S.R. Floeter, and P.R. Guimarães. 2017. Nestedness across biological scales. *PLOS ONE* 12(2): e0171691. doi.org/10.1371/journal.pone.0171691.

May, R.M. 2006. Network structure and the biology of populations. *Trends in Ecology and Evolution* 21(7): 394-399. doi: 10.1016/j.tree.2006.03.013.

McEwen, B.S. 2008. Central effects of stress hormones in health and disease: Understanding the protective and damaging effects of stress and stress mediators. *European Journal of Pharmacology* 583 (2008) 174–185.

Milesi P, Weill M, Lenormand T, Labbé P. Heterogeneous gene duplications can be adaptive because they permanently associate overdominant alleles. *Evol Lett*. 2017 Jul 21;1(3):169-180. doi: 10.1002/evl3.17.

Morrongiello, J. R., Bond, N. R., Crook, D. A. and Wong, B. B. 2012. Spatial variation in egg size and egg number reflects trade-offs and bet-hedging in a freshwater fish. *J Anim Ecol* 81, 806-17.

National Research Council. 2009. Vision and Change: A new biology for the 21st century. National Academies Press.

Nielsen, J. 2017. Systems biology of metabolism. *Annual Review of Biochemistry* 86: 245-275. doi 10.1146/annurev-biochem-061516-044757.

Nijhout HF, Best JA, Reed MC. Systems biology of robustness and homeostatic mechanisms. *Wiley Interdiscip Rev Syst Biol Med*. 2019 May;11(3):e1440. doi: 10.1002/wsbm.1440

O'Boyle, N., Turner, N.C.A., Roe, A.J., Connolly, J.P.R 2020. Plastic circuits: Regulatory flexibility in fine tuning pathogen circuits. *Trends in Microbiology* 28(5): 360-371. doi: 10.1016/j.tim.2020.01.002.

Olofsson, H., Ripa, J. and Jonzen, N. 2009. Bet-hedging as an evolutionary game: the trade-off between egg size and number. *Proc Biol Sci* 276, 2963-9.

Orth, J.D., I. Thiele, B.Ø. Palsson. 2010 What is flux balance analysis? *Nature Biotechnology* 28: 245-248. 10.1038/nbt.1614.

Payne, J.L. and Wagner, A. 2019. The causes of evolvability and their evolution. *Nature* 20: 24-39. doi: 10.1038/s41576-018-0069-z.

Pienta, K.J, Hammarlund, E.U., Axelrod, R., Brown, J.S., Amend, S.R. 2020. Poly-aneuploid cancer cells promote evolvability, generating lethal cancer. *Evol. Appl.* 13: 1626-1634. doi: 10.1111/eva.12929

Pimpinelli, S. and Piacentini, L. 2020. Environmental change and the evolution of genomes: Transposable elements as translators of phenotypic plasticity *Functional Ecology* 34: 428-441. doi: 10.1111/1365-2435.13497.

Proulx, S.R., Promislow, D.E.L., Phillips, P.C. 2005. Network thinking in ecology and evolution. *TREE* 20(6): 345-353. doi: 10.1016/j.tree.2005.04.004.

Qiao, Y., Yan, H., Duan, L., Miao, J. 2020. Finite-time synchronization of fractional-order gene regulatory networks with time delay. *Neural Networks* 126: 1-10. doi: 10.1016/j.neunet.2020.02.004.

Rauter, T., Burgstaller, S., Gottschalk, B., Ramandi-Muja, J., Bischof, H., Hay, J.C., Graier,

- W.F., Walli, R. 2020. ER-to-Golgi transport in HeLa cells displays high resistance to Ca²⁺ and energy stresses. *Cells* 9(10): 2311. doi: 10.1111/eva.12929.
- Reid, R.S., Nkedianye, D., Said, M.Y., Kaelo, D., Neselle, M., Makui, O., Onetu, L., Kiruswa, S., Ole Kamuaro, N., Kristjanson, P., Ogutu, J., BurnSilver, S.B., Goldman, M.J., Boone, R.B., Galvin, K.A., Dickson, N.M., Clark, W.C. 2016. Evolution of models to support community and policy action with science: Balancing pastoral livelihoods and wildlife conservation in savannas of East Africa. *PNAS* 113(17): 4579-4584. doi:10.1073/pnas.0900313106.
- Romero, L. M., Dickens, M. J. and Cyr, N. E. (2009). The reactive scope model -- A new model integrating homeostasis, allostasis, and stress. *Hormones and Behavior* 55, 375-389.
- Saiz, H., Dainese, M., Chiarucci, A., Nascimbene, J. 2021. Networks of epiphytic lichens and host trees along elevation gradients: Climate change implications in mountain ranges. *J. Ecol.* 109: 1122-1132. doi: 10.1111/1365-2745.13538.
- Sanders et al. 2018. Trophic redundancy reduces vulnerability to extinction cascades. *PNAS*. 115 (10) 2419-2424. doi: 10.1073/pnas.1716825115.
- Sauer, U. 2006. Metabolic networks in motion: ¹³C- based flux analysis. *Molecular Systems Biology* 2: 62. 10.1038/msb4100109.
- Schwartz, M. W., Woods, S. C., Porte, D., Seeley, R. J., & Baskin, D. G. (2000). Central nervous system control of food intake. *Nature.*, 404(6778), 661–671. doi: 10.1038/35007534.
- Selye, H. 1973. Homeostasis and Heterostasis. *Perspectives in Biology and Medicine*, 16, 441-445.
- Sgro, C.M, Low, A.J., Hoffman, A.A. 2011. Building evolutionary resilience for conserving biodiversity under climate change. *Evol. Appl.* 4(2): 326-337. doi: 10.1111/j.1752-4571.2010.00157.x
- Sieriebriennikov, B. and Sommer, R. J. (2018). Developmental Plasticity and Robustness of a Nematode Mouth-Form Polyphenism. *Front Genet* 9, 382.
- Stenuit, B., Agathos, S.N. 2015. Deciphering microbial community robustness through synthetic ecology and molecular systems synecology. *Curr. Opin. Biotechnology* 33: 305-317. doi.org/10.1016/j.copbio.2015.03.012.
- Sterling, P., Eyer, J., 1988. Allostasis, a new paradigm to explain arousal pathology. In: Fisher, S., Reason, J. (Eds.), *Handbook of Life Stress, Cognition and Health*. John Wiley & Sons, New York, pp. 629–649.
- Sutherland, J. P. 1990. Perturbations, resistance, and alternative views of the existence of multiple stable points in nature. *Am. Natural.* 136, 270–275. doi: 10.1086/285097.
- Tang, M., Hong, J., Guo, S., Liu, G., Shen, G.Q. 2021. A bibliometric review of urban energy metabolism: Evolutionary trends and the application of network analytical methods. *J. Cleaner Production* 279(10): 123403. doi: 10.1016/j.jclepro.2020.123403.
- Tayyrov A, Stanley CE, Azevedo S, Künzler M. 2019. Combining microfluidics and RNA-sequencing to assess the inducible defensome of a mushroom against nematodes. *BMC Genomics*. 25;20(1):243. doi: 10.1186/s12864-019-5607-3.
- Turner KM, Deshpande V, Beyter D, Koga T, Rusert J, Lee C, Li B, Arden K, Ren B, Nathanson DA, Kornblum HI, Taylor MD, Kaushal S, Cavenee WK, Wechsler-Reya R, Furnari FB, Vandenberg SR, Rao PN, Wahl GM, Bafna V, Mischel PS. 2017. Extrachromosomal oncogene amplification drives tumour evolution and genetic heterogeneity. *Nature*. 543(7643):122-125. doi: 10.1038/nature21356.
- Ukrantseva, S., Yashin, A. I. and Arbeev, K. G. (2016) Resilience versus robustness in aging. *J*

- Gerontol A Biol Sci Med Sci 71, 1533-1534.
- Varadhan, R., J.D. Walston, K. Bandeen-Roche. 2018. Can physical resilience and frailty in older adults be linked by the study of dynamical systems? *J. Am. Geriatr. Soc.* 66(8): 1455-1458. doi: 10.1111/jgs.15409.
- Wada, H. 2019. Damage-Fitness Model: the missing piece in integrative stress models. *Stress* 22, 548-562. doi.org/10.1080/10253890.2019.1614556.
- Wagner, Andreas. 2005. Robustness and evolvability in living systems. Princeton, NJ: Princeton Univ. Press.
- Walker, B.; Holling, C. S.; Carpenter, S. R. Kinzig, A. 2004. Resilience, adaptability and transformability in social–ecological systems. *Ecology and Society*. 9 (2): 5. doi:10.5751/ES-00650-090205.
- Weetman D, Djogbenou LS, Lucas E. 2018. Copy number variation (CNV) and insecticide resistance in mosquitoes: evolving knowledge or an evolving problem? *Curr Opin Insect Sci.* 27:82-88. doi: 10.1016/j.cois.2018.04.005.
- Whitacre, J. M. (2012). Biological robustness: paradigms, mechanisms, and systems principles. *Front. Genet.* 3:67. doi: 10.3389/fgene.2012.00067.
- Wild, B., Dormagen, D.M., Zachariae, A., Smith, M.L., Traynor, K.S., Brockmann, D., Couzin, I.D., Landgraf, T. 2021. Social networks predict the life and death of honey bees. *Nature Communications* 12, 1110. doi: 10.1038/s41467-021-21212-5.
- Wintle, B.A., Legge, S., Woinariski, J.C.Z. 2020. After the megafires: what next for Australian wildlife? *TREE* 35(9): 753-757. doi: 10.1016/j.tree.2020.06.009.
- Wu, D., Wang, N., Wu, N. 2019. Vulnerability analysis of global container shipping linear network based on main channel disruption. *Maritime Policy and Management* 46(4): 394-409. doi: 10.1080/03088839.2019.1571643.
- Xiuong, W., J. A. Morgan, J. Ungerer, B. Wang, P.C. Maness and J. Yu. 2015. The plasticity of cyanobacterial metabolism supports direct CO₂ conversion to ethylene. *Nature Plants* 1: 15053. 10.1038/nplants.2015.53.
- Zhang, D., Iyer, L.M., Burroughs, A.M., Aravind, L. 2014. Resilience of biochemical activity in protein domains in the face of structural divergence. *Curr. Opin. Struct. Biol.* 26: 92-103. doi: 10.1016/j.sbi.2014.05.008
- Zhu, H., Zhang, P., Zhong, Z., Xia, J., Rich, J. Mai, J., Su, X., Tian, Z., Bachman, H., Rufo, J., Gu, Y., Kang, P., Chakrabarty, K., Witelsky, T.P., Huang, T.J. 2021. Acoustohydrodynamic tweezers via spatial arrangement of streaming vortices. *Sci. Adv.* 7(2): eabc7885. doi: 10.1126/sciadv.abc7885.