

Opinion

Maximizing biological insights from instruments attached to animals

Roxanne S. Beltran ($^{1,*,@}$, A. Marm Kilpatrick (1 , Simona Picardi (2 , Briana Abrahms (3 , Gabriel M. Barrile (4 , William K. Oestreich (5 , Justine A. Smith (6 , Max F. Czapanskiy⁷, Arina B. Favilla (1,* , Ryan R. Reisinger (9 , Jessica M. Kendall-Bar (10 , Allison R. Payne (11 , Matthew S. Savoca (11 , Danial G. Palance¹, Samantha Andrzejaczek (11 , Daphne M. Shen (11 , Taiki Adachi (8 , Daniel P. Costa^{1,7}, Natalie A. Storm (11 , Conner M. Hale (11 , and Patrick W. Robinson¹

Instruments attached to animals ('biologgers') have facilitated extensive discoveries about the patterns, causes, and consequences of animal behavior. Here, we present examples of how biologging can deepen our fundamental understanding of ecosystems and our applied understanding of global change impacts by enabling tests of ecological theory. Applying the iterative process of science to biologging has enabled a diverse set of insights, including social and experiential learning in longdistance migrants, state-dependent risk aversion in foraging predators, and resource abundance driving movement across taxa. Now, biologging is poised to tackle questions and refine ecological theories at increasing levels of complexity by integrating measurements from numerous individuals, merging datasets from multiple species and their environments, and spanning disciplines, including physiology, behavior and demography.

Testing and refining ecological theories with biologgers

A holistic understanding of complex ecological processes requires creative solutions for collecting data across sites, species, and systems. Archival or transmitting instruments attached to animals (hereafter, **biologgers**) (see Glossary) have become routine tools for recording physiological, behavioral, and demographic characteristics of individuals, and for inferring their interactions with environmental and ecological features [1] (Figure 1). These animal and ecosystem insights are obtained through various levels of data processing by the researcher, ranging from minimal (sensors that measure heart rates and ambient light levels) to extensive (calculations of activity budgets and energy expenditure from sensor measurements) (Figure 1 and see Table S1 in the supplemental information online). The resulting discoveries in animal physiology, ecology, and evolution span temporal and spatial extents and resolutions that far exceed those possible from traditional observation and mark–recapture approaches. Major discoveries include understanding how navigation abilities allow animals to migrate across the globe and how physiological and anatomical adaptations allow animals to inhabit harsh regions such as polar and deep-ocean habitats.

The rapid development of biologging has motivated numerous proposals for future directions. The biologging community has been urged to integrate with other disciplines [2], use mathematics and optimality [3], re-unite big data approaches with field-based ecological processes [4], target key knowledge gaps [5], learn lessons from human mobility research [6], and reconnect tools and questions [7,8]. In recent years, biologging studies have transitioned from describing unique biological observations to adopting a more rigorous scientific

Highlights

Many animal characteristics that strongly influence population, community, and ecosystem dynamics are difficult or impossible to measure using traditional field methods.

Biologging sensors enable broad-scale, high-resolution measurements of the physiological, behavioral, demographic, social, and environmental interactions underpinning patterns in nature.

We present a framework, case studies, and outstanding questions for integrating biologging data with theoretical concepts to facilitate process explanation and prediction.

Future studies should use biologging technology to rigorously test ecological hypotheses, transitioning from pattern description to refining and advancing ecological theory.

¹Department of Ecology and Evolutionary Biology, University of California Santa Cruz, 130 McAllister Way, Santa Cruz, CA 95060, USA ²Department of Fish and Wildlife Sciences, University of Idaho, 875 Perimeter Drive MS 1136. Moscow, ID 83844, USA ³Center for Ecosystem Sentinels. Department of Biology, University of Washington, 1410 NE Campus Pkwy, Seattle, WA 98195, USA ⁴Department of Zoology and Physiology, University of Wyoming, 1000 E University Ave, Laramie, WY 82071, USA ⁵Monterey Bay Aguarium Research Institute, 7700 Sandholdt Rd, Moss Landing, CA 95039, USA

Trends in Ecology & Evolution, Month 2024, Vol. xx, No. xx https://doi.org/10.1016/j.tree.2024.09.009 1 © 2024 The Author(s). Published by Elsevier Ltd. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).



approach. Now, researchers typically formulate defined questions, ensure adequate sample sizes, and conduct in-depth analyses, shifting from simple movement descriptions to hypothesis-driven investigations. Biologging studies have also applied 'big-data' approaches to tackle behavioral and ecological mechanisms underlying animal movement and the proximate internal and external factors that constrain them [9,10]. In turn, the discoveries and natural history descriptions enabled by biologging technology are primed for testing and refining long-standing ecological theories [11], with implications for broad generalizations [12] and evidence-based conservation solutions [13].

We believe a critical next step is to use biologging to answer questions that span multiple taxa, such as whether generalizable 'rules' underpin complex movements and species interactions [14]. For many emerging questions, ecological theory exists to generate hypotheses about how animals behave, and how behavior scales up to shape species' distributions and broader ecosystem processes [15]. In contrast to controlled laboratory experiments, biologging facilitates measurements in natural settings where competing selective pressures such as predation, starvation, competition, and infection exist. For example, data from satellite trackers and accelerometers can be used to estimate foraging rates and prey densities to test hypotheses about **functional responses** (i.e., the shape of the curve that describes prey capture rates by predators as a function of prey availability) (Figure 1). We illustrate the potential of data from biologgers to integrate complex data into mechanistic evaluations, thereby offering a powerful approach to hypothesis testing and theory refinement.

Examples of development from discovery to theory

We provide three case-studies that range from single-species behavioral research to multispecies interactions and consequences for population dynamics (Figure 2). In these examples, we highlight the progression from pattern description to hypothesis testing to theory development and suggest directions for future research.

State-dependent risk-taking throughout oceanic foraging in elephant seals

Body condition is hypothesized to mediate how animals navigate **risk-reward trade-offs**; however, the difficulty of simultaneously measuring body condition, predation risk, and food rewards in the wild has limited our understanding of how intrinsic states affect risk-taking behavior [16]. Biologgers have begun to address this gap by collecting data on animal movement behavior. For instance, time-depth recorders facilitated the discovery that northern elephant seals (Mirounga angustirostris) cease swimming and passively drift during some dives [17]. The discovery of these so-called 'drift dives' led to the hypothesis that seals in inferior body condition with less body fat would sink faster [18]. Later, this hypothesis was tested with longitudinal energy gain rates [19], enabling drift dives to be used as a valuable metric for estimating body composition [20] at fine spatiotemporal scales throughout oceanic foraging trips. In addition, measurements of drifting seals were used to test hypotheses about the functions of drift dives for resting and food processing [21,22]. Finally, drift dive analysis provided a method to estimate daily body composition and time-activity budgets throughout months-long oceanic foraging trips, which enabled the testing of predictions from state-dependent risk-taking theory regarding optimal periods for animals to rest and forage [23]. This three-decade research arc, examining whether behavior is state-dependent throughout oceanic foraging trips, demonstrated that seals in superior body condition sacrifice more profitable nighttime foraging hours to sleep in the safety of darkness (Figure 2) [23]. More broadly, this research provides insights into how other wild animals may move relative to ambient light levels, particularly in marine environments where light levels are mediated by depth in addition to location and time-of-day.

⁶Department of Wildlife, Fish, and Conservation Biology, University of California Davis, 1 Shields Ave, Davis, CA 95616, USA

 ⁷Institute of Marine Sciences, University of California Santa Cruz, 115 McAllister Way, Santa Cruz, CA 95060, USA
⁸National Institute of Polar Research, 10-3 Midori-cho, Tachikawa, Tokyo 190-8518, Japan
⁹School of Ocean and Earth Science, University of Southampton, European Way, Southampton SO14 3ZH, UK
¹⁰Center for Marine Biotechnology and Biomedicine, Scripps Institution of Oceanography, UC San Diego, La Jolla, CA 92037. USA

¹¹Hopkins Marine Station, Stanford University, 120 Ocean View Blvd, Pacific Grove, CA 93950, USA

*Correspondence: roxanne@ucsc.edu (R.S. Beltran). [®]X: @roxannesbeltran



Developing new technologies and analytical approaches enables the study of state-dependent behavior in free-ranging animals. New on-board processing algorithms have been developed to estimate and transmit real-time body composition data to test state-dependent behavior in marine species that do not perform drift dives [24]. Likewise, other components of an animal's internal state (e.g., hunger, heat, stress, and exhaustion [25]) may influence behavior and could be measured with new sensors. In the future, state-of-the-art physiological biologgers that measure brain activity [26] and heart rate [27] could be used to test theories of physiological recharge, rebound, and replenishment (e.g., sleep quotas) in other species [28]. Additionally, simultaneously instrumenting predators and prey across dynamic conditions [29–31], which has been achieved in some study systems, could be used to test theoretical predictions of **dynamic landscapes of fear** or **energy landscapes** frameworks.

Social and experiential learning in whooping cranes

Understanding how animals learn long-distance migrations has implications for wildlife conservation and management, including the likely success of protected area designation and species reintroductions or translocations. The role of social versus experiential learning in migratory behavior has been tested in studies of reintroduced whooping cranes (*Grus americana*) (Figure 2). Using satellite transmitters and very high frequency (VHF) radio telemetry tags, the movements and survival of re-introduced cranes demonstrated successful migration and dispersal patterns [32] and motivated hypothesis testing about whether reintroduced cranes migrate more efficiently when flying with experienced birds [33]. A long-term satellite tracking dataset indicated an ontogenetic switch from social to experiential learning as birds age, allowing them to track resources throughout their migration at all ages [34]. This work transitioned from initial descriptive discoveries (examining whether reintroduction was a successful management strategy) to theory-testing (disentangling the roles of social versus experiential learning).

Future biologging research could provide answers to many remaining questions pertaining to animal learning and social interactions. For example, biologgers could help resolve the populationlevel consequences of interrupted information transfer (e.g., by human-made noise) in systems where social learning is a primary mechanism shaping movement strategies [35]. Likewise, many facets of information gathering, particularly social cues and eavesdropping on other species, remain poorly understood but could be answered with animal-borne microphones or video recorders [36,37].

Migration based on past and current resource availability across species

Foundational biologging discoveries about memory and **behavioral compensation** have set the stage for rich theory and emerging tools to address long-standing questions about migration [38]. One area that has seen significant theoretical progress across species and systems is the influence of variation in resource availability on migratory behavior (Figure 2). In the late 1970s, researchers observed simultaneous changes in plant phenology and migratory movements in waterfowl [39]. Later, biologgers were used to examine the link between **vegetation green-up** and ungulate migration decisions [40]. Follow-up studies tested hypotheses that mule deer (*Odocoileus hemionus*) migrate in concert with green-up, maximizing the intake of high-quality forage along migration routes [41], and that the rate and order of green-up influences the ability of animals to **green-wave surf** [42]. Simulation of zebra (*Equus burchelli antiquorum*) and mule deer movement, and empirical validation with GPS tracks, showed that previous experience (memory) of green-up patterns also plays an essential role in migration [43,44]. More recently, biologging data has been used to suggest revising movement theory to include the roles of memory [45] and social cues [37] alongside the traditional drivers of contemporaneous environmental conditions perceived by individuals.

Glossary

Behavioral compensation: animals change their behavior to overcome constraints (e.g., regulating life history timing or transit speed to reach a destination on-time or match presence with resource availability peaks). Biologgers: instruments attached to animals that record and/or transmit physiological, behavioral, and demographic characteristics of individuals, and their interactions with environmental and ecological features. Dynamic landscapes of fear: spatial and temporal variation in prey

perception of predation risk. **Energy (cost) landscapes:** spatial and temporal variation in an animal's cost of moving and foraging, irrespective of energy (gain) landscapes due to resource distribution and abundance.

Functional responses: the relationship between an animal's response to a resource and its availability.

Green-wave surfing: migratory animals track and consume high-quality vegetation during green-up in the spring. Resource pulses: sporadic, intense episodes of increased resource availability in space and time.

Resource tracking: animals increase energy gain by moving in conjunction with temporal variation in resources across space.

Risk-reward trade-offs: animals balance the need to obtain food rewards while avoiding predation risks.

State-dependent risk-taking: body condition mediates how animals navigate risk-reward trade-offs. Vegetation green-up: a burst of

vegetation productivity during spring that propagates across landscapes.



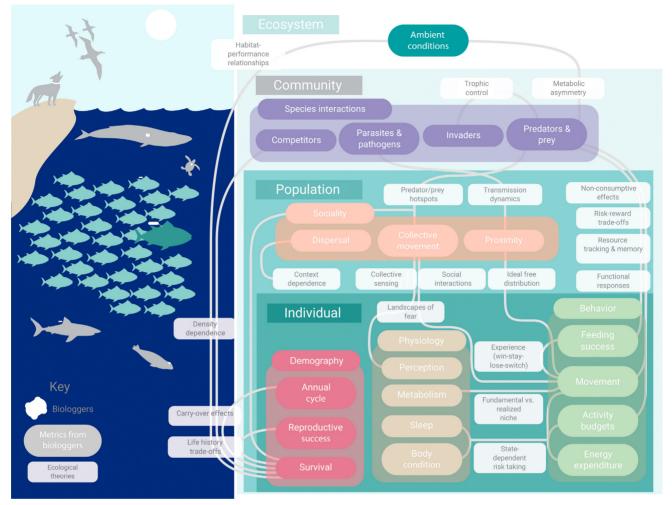


Figure 1. Data generated by archival or transmitting biologgers can be used to calculate extrinsic and intrinsic variables (colored ovals) that can be used to address key research topics (white squares) in behavior and ecology. These research topics span individual, population, community, and ecosystem scales, and cross traditional disciplinary boundaries (red, demography; gold, physiology; green, behavior; orange, sociality; purple, species interactions). Note that there are several critical steps, such as data conversions, processing, and analysis, between raw sensor-derived measurements and the variables presented here (see Table S1 in the supplemental information online). For example, species interactions between predators and prey can be measured by pairing accelerometers and acoustic loggers (attached to prey) that measure escape movement responses to predator vocalizations. Illustration by Jessica Kendall-Bar.

Biologging science could be used to address how migratory animals respond to environmental drivers. New technologies are facilitating the measurement of *in situ* resource distribution and abundance at a sufficiently fine resolution to inform theory about animal movement. For example, Chevallay *et al.* [46] used a novel animal-borne echosounder to measure prey detection and capture events so that estimates of sensory perception volume could be made. Continuing to ground these research efforts in existing theory (e.g., framing the introduction with previous work on resource patterns such as **pulses** [47] and waves [48]) can help maximize insights into how movement patterns emerge from stochastic resource patterns such as irregular rainfall or unpredictable fire events. Moreover, testing distinct theories with a common set of data [e.g., predation risk aversion (a top-down process)] and **resource tracking** (a bottom-up process)] could advance our understanding of how movement outcomes result from risk–reward trade-offs [48].



Biologging for the future

Although many ecological studies that leverage biologging tools have been descriptive, contemporary research can effectively test theory by building on past insights while focusing on generalizable patterns, testing hypotheses, and linking patterns to ecological processes (Figure 2). Many individual examples of biologging studies in both the marine and terrestrial realms test theory (Figure 3). The highlighted studies share many common approaches to their organization and focus, including (i) the introductions are often framed with generalizable patterns in addition to system-specific information, (ii) the background information often includes theory papers that

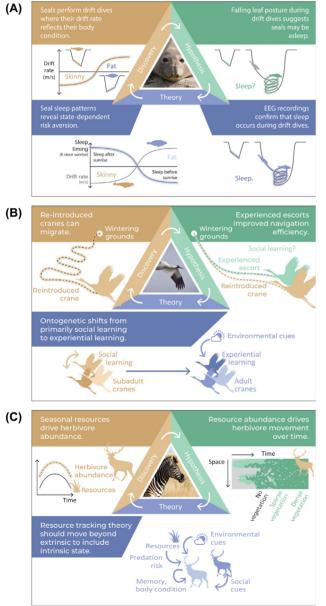
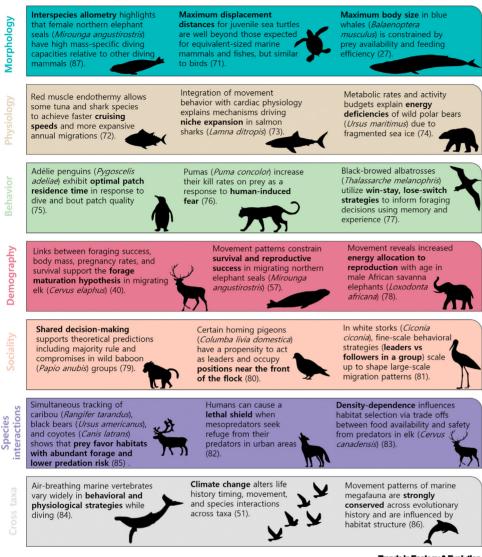


Figure 2. Illustrated examples of the development from discovery to hypothesis and testing to theory refinement for each of three case studies. (A) State-dependent risk-taking throughout migration in northern elephant seals (Mirounga angustirostris). (B) Social and experiential learning in whooping cranes (Grus americana). (C) Migration based on past and current resource availability across species. These case studies show how discoveries and theory testing can comprise a positive feedback loop, where observations lead to insights that inform hypotheses, and testing those hypotheses can lead to theory development, which can then be refined through continued observations and discoveries. Illustration by Jessica Kendall-Bar. Abbreviation: EEG, electroencephalogram.

Trends in Ecology & Evolution





Trends in Ecology & Evolution

Figure 3. Examples of biologging studies with strong conceptual/theoretical foundations. Studies are organized by discipline and cross levels of biological organization from individuals to populations and ecosystems. See [27,40,51,57,71–87].

inform testable hypotheses, and (iii) the discussions often describe how insights produced from the biologging study advance our theoretical understanding of ecological patterns and processes.

Data synthesis

Many theoretical and conceptual questions require synthesizing multiple biologging datasets. Doing so can produce more generalizable knowledge about patterns and processes in the natural world across vast spatial and temporal scales. Major synthesis efforts include the Tagging of Pacific Predators [49], Retrospective Analysis of Antarctic Tracking Data, [50], Arctic Animal Movement Archive [51], coronavirus disease 2019 (COVID-19) Bio-Logging Initiative [52], and the Wyoming Migration Initiative [53]. Many of these efforts were initially discovery-based but laid a foundation for future concept-driven research. A promising area for future conceptual



research is bridging terrestrial and marine systems. Likewise, re-analyzing and synthesizing datasets can maximize insights and minimize research impacts by reducing the need to instrument additional animals.

Integration of biologging with auxiliary data

Understanding and predicting the spatiotemporal dynamics of populations is a central goal in ecology and conservation, and requires understanding variation in, and genetic and phenotypic drivers of, demographic rates [54–56]. Integrating biologging with auxiliary datasets and complementary disciplines can enable tests of hypotheses that span traditional boundaries (Table S1). For example, integrating biologging with demographic data from mark–recapture programs could be used to quantify the fitness consequences of movement behavior [57,58]. Moreover, considering species-level traits (e.g., body size, brain size, reproductive characteristics, and diet) when analyzing tracking data [59] or conducting vulnerability risk assessments could facilitate valuable ecological-evolutionary insights. Biologging data can also facilitate insights into human–wildlife conflict theory through field-based experiments that combine camera traps, audio recorders, and biologgers to quantify movement responses to simulated human activity and conservation interventions [60,61].

Novel tools and approaches

Using existing biologging data to address new questions outside those of the initial study is sometimes problematic. In these cases, new tools may be needed to answer a given conceptual question. Here, concept-driven motivation for improved tagging technologies can drive the collaborative development of smaller and longer-lasting biologgers and more reliable sensors that overcome current limitations in measuring the covariates that matter to animals (e.g., *in situ* resources, predator abundance, multimodal sensory information perception and processing, real-time infection, and heat stress) [62].

Inclusive, equitable practices

More equitable approaches to biologging science, including data collection, processing, archiving, and reporting, will provide the greatest impact and the broadest participation by diverse individuals, institutions, and nations. For example, the research community can seek to understand and preserve the socioecological systems within which the animals exist through global and interdisciplinary collaborators [8], including Indigenous communities who should be compensated for their time and meaningfully involved in study design decisions based on their knowledge of the system [i.e., development of research question(s), instrumentation size, and handling and attachment methods] [63,64]. Likewise, a global tag registry could help the biologging community and public know whom to contact about sampling efforts and datasets to optimize the probability and efficiency of collaboration [65]. Standardized and reproducible data can fast-track theory-testing across systems by facilitating synthesis of large global biologging datasets [66]. This is especially needed for nonspatial data such as physiologging, accelerometry, and video data that do not fit the standards developed for spatial data [67]. Open-access publication of those datasets must credit those who collect and share data [68]. Finally, partnering with theoreticians will allow empiricists to identify promising areas for theory development and refinement using biologging data.

Concluding remarks

There is a wide range of novel research directions that can be pursued with biologgers, ranging across temporal and spatial scales (Figure 4; see Outstanding questions). The key to moving from anecdote to generalizable theory is to examine interdisciplinary patterns and processes across species and habitats. For example, habitat selection can be influenced by the population context (e.g., density and social dynamics), by the community context (e.g., predators and

Outstanding questions

How do system-specific findings about ecological patterns and processes generalize across systems?

How can we replicate biologging data collection processes or leverage large datasets across systems or species to test the conditions in which ecological theories are supported?

Which ecological theories are primed for testing or refinement through biologging science?

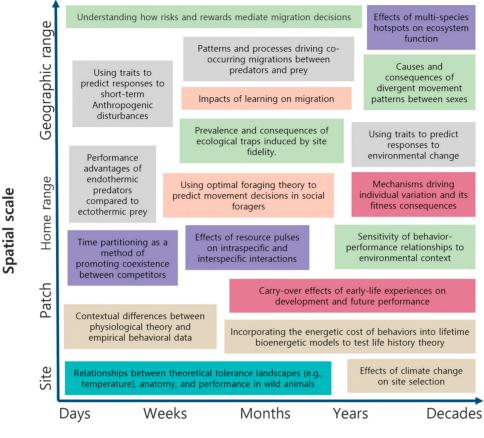
Which insights from empirical biologging studies are well poised to inform theory development?

How do we engage across disciplinary boundaries to advance, test, and apply ecological theory using biologgers? Who are key partners for removing barriers and fostering interdisciplinary collaborations in this space?

What innovations in biologging technology are needed to facilitate theoretical advancements? How can we accomplish them?

How can we widen the adoption of open science practices among biologging users to accelerate the pace of basic and applied science?





Temporal scales

Trends in Ecology & Evolution

Figure 4. Some emerging research directions that could be addressed using biologgers, across temporal and spatial scales. Each topic is colored by discipline (blue, morphology; gold, physiology; green, behavior; red, demography; orange, sociality; purple, species interactions; gray, cross taxa). Many of the featured research topics are moderately well understood from a theoretical perspective but are primed for empirical testing through finer-resolution or larger-scale biologging data. For others, extensive empirical data could challenge or refine theory by identifying deviations from optimality or highlighting discrepancies between theory and empirical data. Many topics represent significant opportunities for developing both theory and empirical evidence, especially across disciplinary boundaries. All can be used as an agenda for further research and a benchmark for future evaluations of progress in biologging science.

competitor density), and by the environmental context (e.g., drought, fire) (Figure 1). Similarly, theory can be applied to the many roles played by humans (e.g., predators, resource drivers, disease sources) to inform conservation and management actions such as reserve design and species protection [69]. Predictive models used for these decisions will be most accurate when informed by a process-based understanding of animal movements and their roles in ecological systems [70]. For example, understanding how animals move relative to human-made barriers such as fences and highways can allow predictions about how increasing urbanization and development will reduce wildlife fitness as the human population grows. Likewise, characterizing links between animal performance and environmental conditions can facilitate predictions of how climate change will trigger species range shifts and population declines.

Biologging enables efficient, fine-scale data collection that can provide the breadth and depth needed to develop, test, and refine our understanding of ecological processes in our rapidly



changing world (Figure 2). Although new theories are rare in ecology, new technological developments can allow existing theories or hypotheses to finally be tested or refined, so long as the research has a strong conceptual focus (Figure 3) [58]. Of course, theory testing and refinement are not always possible. Research is still in the description and discovery phase for species that have never been instrumented or that inhabit under-studied regions and biomes. In these cases, a closer connection between conceptual questions and biologging technology can expedite the development of new theories and contribute to the iterative process of testing and refinement [7]. Through these avenues, biologging can provide insight into how nature works and provide a roadmap for better protection of species, ecosystems, and the services they provide [68].

Acknowledgments

The authors would like to thank the many pioneers of biologging science who developed a creative vision of possibilities for studying animals, and who supported our growth and development as young scientists. Thank you to Josh London, Scott Yanco, Autumn-Lynn Harrison, and Mike Kowalski for helpful suggestions, and to Cara Munro for helping with Figure 4. R.S.B. was supported by a Packard Fellowship for Science and Engineering, a Beckman Young Investigator award, the Office of Naval Research Young Investigator Program, and a National Science Foundation grant (#2052497). Funding was also provided by a National Science Foundation grant (#1911853).

Declaration of interests

The authors declare no competing interests.

Supplementary information

Supplementary information associated with this article can be found online at https://doi.org/10.1016/j.tree.2024.09.009.

References

- Watanabe, Y.Y. and Papastamatiou, Y.P. (2023) Biologging and biotelemetry: tools for understanding the lives and environments of marine animals. *Annu. Rev. Anim. Biosci.* 11, 247–267
- Burggren, W.W. (1987) Invasive and noninvasive methodologies in ecological physiology: a plea for integration. In *New Directions in Ecological Physiology* (Feder, M.E. et al., eds), pp. 251–272, Cambridge University Press
- Nathan, R. et al. (2008) A movement ecology paradigm for unifying organismal movement research. Proc. Natl. Acad. Sci. 105, 19052–19059
- Hebblewhite, M. and Haydon, D.T. (2010) Distinguishing technology from biology: a critical review of the use of GPS telemetry data in ecology. *Philos. Trans. R. Soc. B Biol. Sci.* 365, 2303–2312
- 5. Hays, G.C. et al. (2016) Key questions in marine megafauna movement ecology. *Trends Ecol. Evol.* 31, 463–475
- Thums, M. et al. (2018) How big data fast tracked human mobility research and the lessons for animal movement ecology. Front. Mar. Sci. 5, 00021
- Börger, L. (2016) Stuck in motion? Reconnecting questions and tools in movement ecology. J. Anim. Ecol. 85, 5–10
- Williams, H. et al. (2020) Optimizing the use of biologgers for movement ecology research. J. Anim. Ecol. 89, 186–206
- 9. Kays, R. et al. (2015) Terrestrial animal tracking as an eye on life and planet. Science 348, aaa2478
- Hussey, N.E. *et al.* (2015) Aquatic animal telemetry: a panoramic window into the underwater world. *Science* 348, 1255642
- Cooke, S.J. et al. (2004) Biotelemetry: a mechanistic approach to ecology. Trends Ecol. Evol. 19, 334–343
- Moll, R.J. et al. (2007) A new 'view' of ecology and conservation through animal-borne video systems. Trends Ecol. Evol. 22, 660–668
- Wilmers, C.C. *et al.* (2015) The golden age of bio-logging: how animal-borne sensors are advancing the frontiers of ecology. *Ecology* 96, 1741–1753
- Abrahms, B. et al. (2017) Suite of simple metrics reveals common movement syndromes across vertebrate taxa. Mov. Ecol. 5, 12
- 15. Pickett, S.T. et al. (2010) Ecological Understanding: the Nature of Theory and the Theory of Nature, Elsevier
- Brown, J.S. (1992) Patch use under predation risk: I. Models and predictions. Ann Zool Fennici 29, 301–309

- Le Boeuf, B.J. et al. (1992) Swim speed in a female northern elephant seal: metabolic and foraging implications. Can. J. Zool. 70, 786–795
- Crocker, D.E. et al. (1997) Drift diving in female northern elephant seals: implications for food processing. *Can. J. Zool.* 75, 27–39
- Biuw, M. et al. (2003) Blubber and buoyancy: monitoring the body condition of free-ranging seals using simple dive characteristics. J. Exp. Biol. 206, 3405–3423
- Pirotta, E. et al. (2019) Modeling the functional link between movement, feeding activity, and condition in a marine predator. Behav. Ecol. 30, 434–445
- 21. Mitani, Y. et al. (2010) Three-dimensional resting behaviour of northem elephant seals: drifting like a falling leaf. *Biol. Lett.* 6, 163–166
- Kendall-Bar, J.M. *et al.* (2023) Brain activity of diving seals reveals short sleep cycles at depth. *Science* 380, 260–265
- Beltran, R.S. et al. (2021) Lightscapes of fear: how mesopredators balance starvation and predation in the open ocean. Sci. Adv. 7, eabd9818
- Adachi, T. et al. (2023) Body condition changes at sea: onboard calculation and telemetry of body density in diving animals. *Methods Ecol. Evol.* 14, 1457–1474
- Blecha, K.A. *et al.* (2018) Hunger mediates apex predator's risk avoidance response in wildland–urban interface. *J. Anim. Ecol.* 87, 609–622
- Libourel, P.-A. et al. (2023) Nesting chinstrap penguins accrue large quantities of sleep through seconds-long microsleeps. Science 382, 1026–1031
- Goldbogen, J.A. et al. (2019) Extreme bradycardia and tachycardia in the world's largest animal. Proc. Natl. Acad. Sci. 116, 25329–25332
- Hooten, M.B. et al. (2019) Running on empty: recharge dynamics from animal movement data. Ecol. Lett. 22, 377–389
- Palmer, M.S. et al. (2022) Dynamic landscapes of fear: understanding spatiotemporal risk. Trends Ecol. Evol. 37, 911–925
- Gallagher, A.J. et al. (2017) Energy Landscapes and the Landscape of Fear. Trends Ecol. Evol. 32, 88–96
- Papastamatiou, Y.P. et al. (2024) Dynamic energy landscapes of predators and the implications for modifying prey risk. *Funct. Ecol.* 38, 284–293

CellPress OPEN ACCESS

Trends in Ecology & Evolution

- Urbanek, R. *et al.* (2005) First cohort of migratory whooping cranes reintroduced to Eastern North America: the first year after release. *Proc. North Am. Crane Workshop* 32, Published online 2005. https://digitalcommons.unl.edu/nacwgproc/32
- Mueller, T. *et al.* (2013) Social learning of migratory performance. Science 341, 999–1002
- Abrahms, B. *et al.* (2021) Ontogenetic shifts from social to experiential learning drive avian migration timing. *Nat. Commun.* 12, 7326
- Jesmer, B.R. *et al.* (2018) Is ungulate migration culturally transmitted? Evidence of social learning from translocated animals. *Science* 361, 1023–1025
- Aikens, E.O. et al. (2022) Viewing animal migration through a social lens. Trends Ecol. Evol. 37, 985–996
- Oestreich, W.K. *et al.* (2022) The influence of social cues on timing of animal migrations. *Nat. Ecol. Evol.* 6, 1617–1625
- Ortega, A.C. *et al.* (2023) Migrating mule deer compensate en route for phenological mismatches. *Nat. Commun.* 14, 2008
- Owen, M. (1980) Wild Geese of the World: their Life History and Ecology, BT Batsford Limited
- Hebblewhite, M. et al. (2008) A multi-scale test of the forage maturation hypothesis in a partially migratory ungulate population. *Ecol. Monogr.* 78, 141–166
- Sawyer, H. and Kauffman, M.J. (2011) Stopover ecology of a migratory ungulate. J. Anim. Ecol. 80, 1078–1087
- Aikens, E.O. et al. (2017) The greenscape shapes surfing of resource waves in a large migratory herbivore. Ecol. Lett. 20, 741–750
- Bracis, C. and Mueller, T. (2017) Memory, not just perception, plays an important role in terrestrial mammalian migration. *Proc. R. Soc. B Biol. Sci.* 284, 20170449
- Merkle, J.A. et al. (2019) Spatial memory shapes migration and its benefits: evidence from a large herbivore. *Ecol. Lett.* 22, 1797–1805
 Abrahms, B. et al. (2019) Memory and resource tracking drive
- blue whale migrations. Proc. Natl. Acad. Sci. 116, 5582–5587
- 46. Chevailay, M. et al. (2023) Large sensory volumes enable Southem elephant seals to exploit sparse deep-sea prey. Proc. Natl. Acad. Sci. 120, e2307129120
- Clark-Wolf, T.J. et al. (2023) Climate presses and pulses mediate the decline of a migratory predator. Proc. Natl. Acad. Sci. 120, e2209821120
- Abrahms, B. et al. (2021) Emerging perspectives on resource tracking and animal movement ecology. Trends Ecol. Evol. 36, 308–320
- 49. Block, B.A. et al. (2011) Tracking apex marine predator movements in a dynamic ocean. *Nature* 475, 86
- Hindell, M.A. et al. (2020) Tracking of marine predators to protect Southern Ocean ecosystems. Nature 580, 87–92
- Davidson, S.C. et al. (2020) Ecological insights from three decades of animal movement tracking across a changing Arctic. Science 370, 712–715
- Rutz, C. et al. (2020) COVID-19 lockdown allows researchers to quantify the effects of human activity on wildlife. *Nat. Ecol. Evol.* 4, 1156–1159
- Kauffman, M.J. et al. (2018) Wild Migrations: Atlas of Wyoming's Ungulates, Oregon State University Press Corvallis, OR
- Bolnick, D.I. et al. (2011) Why intraspecific trait variation matters in community ecology. *Trends Ecol. Evol.* 26, 183–192
- Crossin, G.T. et al. (2014) Tracking fitness in marine vertebrates: current knowledge and opportunities for future research. Mar. Ecol. Prog. Ser. 496, 1–17
- Morales, J.M. et al. (2010) Building the bridge between animal movement and population dynamics. *Philos. Trans. R. Soc. B Biol. Sci.* 365, 2289–2301
- Beltran*, R.S. et al. (2023) Physiological tipping points in the relationship between foraging success and lifetime fitness of a longlived mammal. *Ecol. Lett.* 26, 706–716
- Schradin, C. and Hayes, L.D. (2017) A synopsis of long-term field studies of mammals: achievements, future directions, and some advice. J. Mammal. 98, 670–677
- Beltran, R. et al. (2024) Integrating animal tracking and trait data to facilitate global ecological discoveries. *EcoEvoRxiv*, Published online May 7, 2024. https://ecoevorxiv.org/repository/view/7089
- Tosa, M.I. et al. (2021) The rapid rise of next-generation natural history. Front. Ecol. Evol. 9, 698131

- Suraci, J.P. et al. (2019) Fear of humans as apex predators has landscape-scale impacts from mountain lions to mice. Ecol. Lett. 22, 1578–1586
- Couzin, I.D. and Heins, C. (2023) Emerging technologies for behavioral research in changing environments. *Trends Ecol. Evol.* 38, 346–354
- Polfus, J.L. et al. (2014) Comparing traditional ecological knowledge and western science woodland caribou habitat models J. Wildl. Manag. 78, 112–121
- Lamb, C.T. et al. (2022) Indigenous-led conservation: pathways to recovery for the nearly extirpated Klinse-Za mountain caribou. *Ecol. Appl.* 32, e2581
- Rutz, C. (2022) Register animal-tracking tags to boost conservation. Nature 609, 221
- Sequeira, A.M.M. et al. (2021) A standardisation framework for bio-logging data to advance ecological research and conservation. *Methods Ecol. Evol.* 12, 996–1007
- Czapanskiy, M.F. and Beltran, R.S. (2022) How reproducibility will accelerate discovery through collaboration in physiologging. *Front. Physiol.* 13, 917976
- Kays, R. and Wikelski, M. (2023) The Internet of Animals: what it is, what it could be. *Trends Ecol. Evol.* 38, 859–869
- Hays, G.C. *et al.* (2019) Translating marine animal tracking data into conservation policy and management. *Trends Ecol. Evol.* 34, 459–473
- Durner, G.M. *et al.* (2009) Predicting 21st-century polar bear habitat distribution from global climate models. *Ecol. Monogr.* 79, 25–58
- Hays, G.C. and Scott, R. (2013) Global patterns for upper ceilings on migration distance in sea turtles and comparisons with fish, birds and mammals. *Funct. Ecol.* 27, 748–756
- Watanabe, Y.Y. et al. (2015) Comparative analyses of animaltracking data reveal ecological significance of endothermy in fishes. Proc. Natl. Acad. Sci. 112, 6104–6109
- Weng, K.C. et al. (2005) Satellite tagging and cardiac physiology reveal niche expansion in salmon sharks. Science 310, 104–106
- Pagano, A.M. *et al.* (2018) High-energy, high-fat lifestyle challenges an Arctic apex predator, the polar bear. *Science* 359, 568–572
- Watanabe, Y.Y. et al. (2014) Testing optimal foraging theory in a penguin–krill system. Proc. R. Soc. B Biol. Sci. 281, 20132376
- Smith, J.A. et al. (2015) Top carnivores increase their kill rates on prey as a response to human-induced fear. Proc. R. Soc. B Biol. Sci. 282, 20142711
- Bonnet-Lebrun, A.-S. *et al.* (2021) A test of the win-stay-loseshift foraging strategy and its adaptive value in albatrosses. *Anim. Behav.* 182, 145–151
- Taylor, L.A. et al. (2020) Movement reveals reproductive tactics in male elephants. J. Anim. Ecol. 89, 57–67
- Strandburg-Peshkin, A. *et al.* (2015) Shared decision-making drives collective movement in wild baboons. *Science* 348, 1358–1361
- Nagy, M. et al. (2010) Hierarchical group dynamics in pigeon flocks. *Nature* 464, 890–893
- Flack, A. et al. (2018) From local collective behavior to global migratory patterns in white storks. Science 360, 911–914
- Prugh, L.R. et al. (2023) Fear of large carnivores amplifies human-caused mortality for mesopredators. *Science* 380, 754–758
- Smith, B.J. *et al.* (2023) Density-dependent habitat selection alters drivers of population distribution in northern Yellowstone elk. *Ecol. Lett.* 26, 245–256
- Favilla, A.B. and Costa, D.P. (2020) Thermoregulatory strategies of diving air-breathing marine vertebrates: a review. *Front. Ecol. Evol.* 8, 555509
- Bastille-Rousseau, G. et al. (2015) Unveiling trade-offs in resource selection of migratory caribou using a mechanistic movement model of availability. *Ecography* 38, 1049–1059
- Sequeira, A.M.M. et al. (2018) Convergence of marine megafauna movement patterns in coastal and open oceans. Proc. Natl. Acad. Sci. 115, 3072–3077
- Adachi, T. et al. (2021) Forced into an ecological corner: Roundthe-clock deep foraging on small prey by elephant seals. Sci. Adv. 7, eabg3628