

Beyond the tangled bank: An evolving meta-ecosystems framework to integrate ecological and evolutionary perspectives

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28 “It is interesting to contemplate a tangled bank, clothed with many plants of many kinds, with
29 birds singing on the bushes, with various insects flitting about, and with worms crawling through
30 the damp earth, and to reflect that these elaborately constructed forms, so different from each
31 other, and dependent upon each other in so complex a manner, have all been produced by laws
32 acting around us.”

33 Charles Darwin, *On the Origin of Species*

34 Darwin’s tangled bank (Darwin 1859) describes life as a network of interdependent organisms
35 shaped by eco-evolutionary processes. This metaphor has gained renewed relevance in the 21st
36 century, as rapid environmental and land-use changes drive ecological reorganization at multiple
37 biological and spatial scales. Yet Darwin’s metaphor was largely local – a single bank. While
38 species interactions and selection pressures can take place at relatively local spatial scales,
39 energy, resources, and organisms move across atmospheric, terrestrial, and aquatic boundaries,
40 connecting ecosystems into larger interacting meta-ecosystem networks (Loreau et al. 2003).
41 The meta-ecosystem framework combines concepts from meta-population and metacommunity
42 theory – which describe spatially structured populations of one or more species connected by
43 the flow of individuals and their interactions – with those from ecosystem ecology – through
44 the tracking of the sources, sinks, and fluxes of energy and materials. Together, it extends
45 beyond the study of individual systems to describe the dynamics of *multiple* ecosystems
46 connected by the “flows of energy, materials, and organisms across ecosystem boundaries”
47 (Loreau et al. 2003). This conceptual foundation has provided a powerful framework to
48 understand emergent properties of these networks that arise from spatiotemporal coupling
49 within and across constituent components. For instance, Massol et al. (2011) theoretically
50 demonstrated that interactions between flows of materials and organisms can affect population
51 persistence and the stability of connected systems in ways that are not apparent when
52 community and ecosystem processes are considered in isolation. Indeed, subsequent empirical
53 studies have found that fluxes across system boundaries can fundamentally change community
54 structure, population dynamics, and ecosystem properties and functions (Gounand et al. 2018).
55 Yet, despite its strengths, meta-ecosystem theory has remained predominantly focused on
56 ecological dynamics, largely ignoring intraspecific genetic variation that provides an additional
57 subsidy - the substrate for evolution by natural selection. Therefore, the very processes implicit
58 in Darwin’s *On the Origin of Species* and his tangled bank metaphor are rarely considered within
59 the meta-ecosystem framework.

60 **The evolving meta-ecosystem framework**

61 Over the last several decades, there has been increasing recognition that adaptive evolution can
62 take place over a few generations, on similar timescales as ecological change, and over fine,
63 microgeographic scales (Carroll et al. 2007; Richardson et al. 2014). Furthermore, the traits that
64 respond to natural selection – such as phenology, growth, stress tolerance, resource allocation,
65 and reproduction – can influence ecosystem processes, such as productivity, decomposition,

66 carbon sequestration, and even landscape morphology and geophysical properties (Dong et al.
67 2024).

68 Despite the exclusion of evolutionary dynamics within theoretical meta-ecosystem models,
69 there is ample evidence that evolutionary change can cause ecological change across ecosystem
70 boundaries. For example, black cottonwood (*Populus trichocarpa*) grows in riparian zones and
71 delivers annual pulses of leaf litter into streams and rivers, providing a major subsidy that
72 supports aquatic food webs. Common gardens have shown that genetic variation in leaf
73 senescence alters decomposition rates and aquatic invertebrate richness (Rodriguez-Cabal et al.
74 2017). Further, studies in other *Populus* species have shown that genetic variation in leaf
75 chemistry also influences decomposition, invertebrate assemblages, and soil net nitrogen
76 (Schweitzer et al. 2008). These and other examples – such as recent body size declines in Pacific
77 salmon and subsequent decreases in marine-derived nutrient transport to freshwater systems
78 (Oke et al. 2020) – illustrate that variation and contemporary changes in key traits can alter
79 meta-ecosystem dynamics through the flow of material, energy, and organisms across
80 ecosystem boundaries.

81 In the face of global change and growing recognition of the potential for evolutionary influence
82 on meta-ecosystem coupling, we must ask: How does adaptive evolution shape mechanistic
83 responses to environmental change at multiple levels of biological organization – from genes to
84 ecosystems? How do these interacting processes determine the resilience of meta-ecosystem
85 functions? To address these questions, meta-ecosystems cannot be treated simply as networks
86 of habitats connected by flows, but instead as dynamic systems with connections that are
87 mediated by organismal evolution and ecological feedbacks – an *evolving* meta-ecosystem (Figure
88 1).

89 **Experimental designs and the necessary data to understand evolving meta-** 90 **ecosystem dynamics**

91 Determining whether evolutionary responses to environmental change have consequential
92 impacts on spatiotemporal meta-ecosystem dynamics requires quantifying and linking: (1)
93 underlying relationships between environmental change and the evolutionary responses of
94 heritable, genetically-based traits (including heritable plasticity); (2) the extent to which those
95 traits directly or indirectly impact cross-boundary flows; and (3) how changes in the properties
96 of these flows (e.g., the direction, magnitude, and spatiotemporal variation and synchrony)
97 caused by evolutionary responses impact meta-ecosystem function and stability – for example,
98 in carbon or nutrient balance, trophic subsidies, connectivity, feedback dynamics, or the
99 resilience of coupled meta-ecosystem functions to disturbance or environmental change.

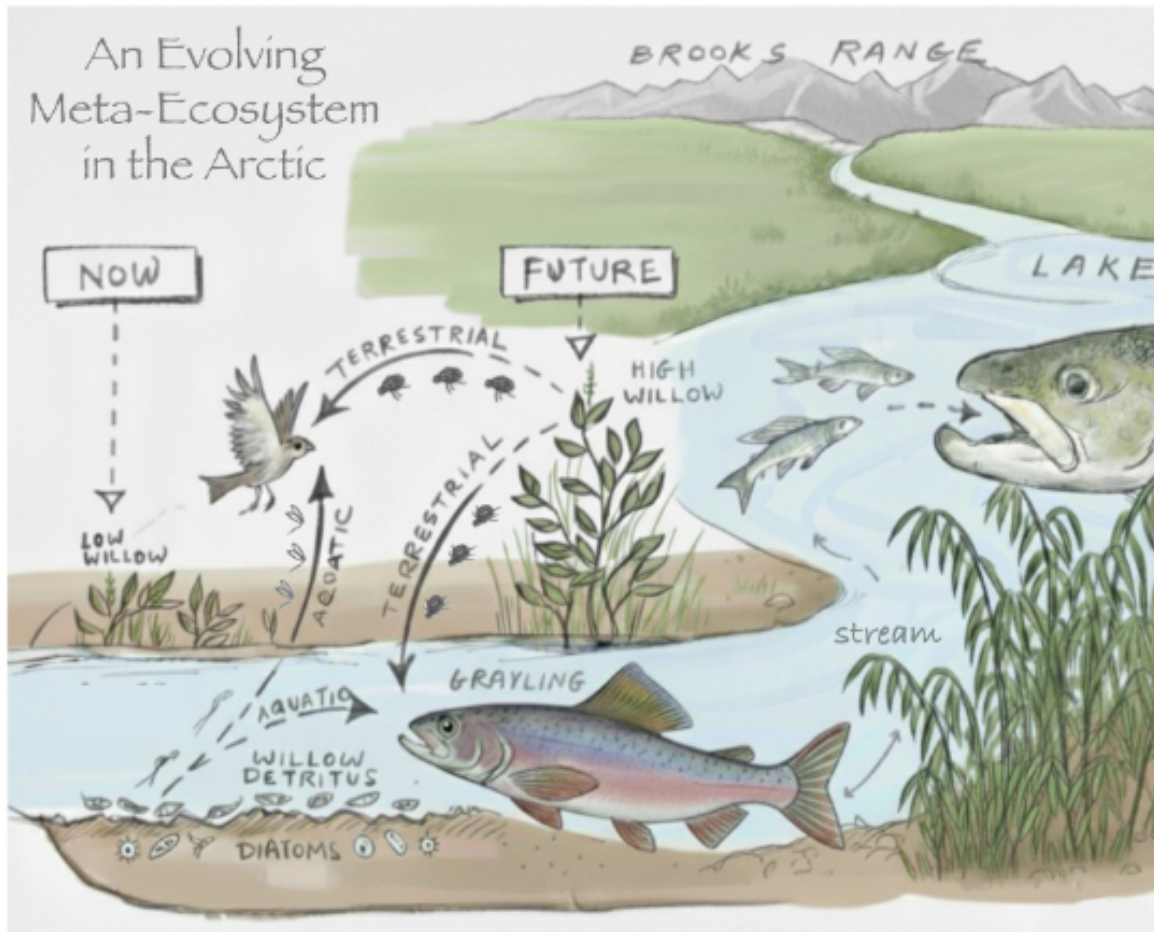


Figure 1 Evolutionary processes impact cross-boundary flows of energy, materials, and organisms in an evolving meta-ecosystem. This figure depicts a tangled bank within an Arctic stream-riparian meta-ecosystem network, where contemporary climate has driven expansion and increased growth of a dominant willow species (*Salix alaxensis*). These evolutionary responses by *S. alaxensis* and other Arctic shrubs are expected to continue and become more prevalent in the future, with the potential to dramatically alter ecological and evolutionary dynamics within and across terrestrial and aquatic ecosystems. In this evolving meta-ecosystem, cross-boundary flows – including willow leaf litter, emerging insects, and movement of Arctic grayling (*Thymallus arcticus*) and white-crowned sparrow (*Zonotrichia leucophrys*) – influence, interact with, and feedback into evolutionary responses of plant phenology and growth, the population dynamics of insect, bird, and fish species, species interactions within and across ecosystem boundaries, and nutrient cycling regimes. Likewise, each of these aspects can influence, interact with, and feedback into other properties within and across boundaries of the meta-ecosystem. Here, transplant experiments and genomic studies can establish signals of local adaptation and contemporary adaptive evolution (e.g., selective sweeps) in these species. The impact of evolutionary responses from increased shrub growth on trophic subsidies across ecosystem boundaries could be quantified through leaf litter measurements (including chemical analysis), the resulting dynamics of aquatic and terrestrial insect abundance, and tracking leaf and insect subsidies to birds and fish through isotopic measures of gut and fecal samples. *In situ* field manipulations such as litter and insect-emergence experiments, controlled dietary trials in grayling, long-term population surveys (through satellite imagery, mark-recapture, nest monitoring), and additional genomic datasets can reveal further aspects of how shrub evolution reshapes the flow of energy, materials, and organisms in this strongly coupled evolving meta-ecosystem.

Temporal and contemporary approaches offer complementary perspectives for understanding evolving meta-ecosystems. For instance, retrospective or long-term observations and experiments can each be used to demonstrate evolutionary change. Moreover, evidence of evolutionary change that coincides with environmental change can provide evidence for an evolutionary response. Resurrection studies of seeds collected from natural or human-mediated seed banks, for example, can reveal how ecologically relevant traits have shifted over periods of changing temperature, hydrology, or nutrient regimes (Christie et al. 2023). Herbarium and museum collections can also provide material for time-series genomics data, which can be used to quantify signals of adaptive responses at the genetic level. Similarly, long-term ecological research and monitoring networks can capture both ecological and evolutionary change (Cocciardi et al. 2024).

Contemporary organismal approaches complement these temporal perspectives. Common gardens or long-term transplant experiments along ecological gradients can quantify the heritable and environmental components of phenotypic variation – including genetic correlations (e.g., trade-offs) – for traits that underlie fitness, mediate meta-ecosystem flows, or both (Blum et al. 2021). These experiments can also disentangle heritable plasticity from non-heritable plasticity, an important factor to consider for species persistence (Ashander et al. 2016). Additionally, population genomic tools (e.g., genome-wide association studies and scans for selective sweeps) and landscape genomic approaches (e.g., estimating demographic history, spatial genetic structure, and relationships between environmental and genetic variation) can identify the genetic components underlying these traits while estimating the strength, timing, and spatial pattern of selection and evolution (Bernatchez et al. 2023). Similarly, comparative genomics can reveal patterns of gene family evolution, structural variation, and selective signals that may underlie adaptive strategies within lineages.

These temporal and contemporary methods quantifying organismal evolution, alongside ecosystem-level measurements or experimental manipulations – on properties such as nutrient supply and stoichiometry, productivity, matter transport rates, dissolved gases, hydrologic discharge, organismal emergence rates, movement, and migration, or trophic subsidy fluxes – allow for explicit analyses of whether genetically based trait variation alters cross-ecosystem fluxes. Establishing these links therefore requires evidence for an evolutionary response, and then tracing and quantifying its direct or indirect impact on population, community, and ecosystem dynamics throughout the meta-ecosystem - an approach outlined below for stream-riparian meta-ecosystems in the Arctic (Figure 1).

The Arctic as a tractable system to observe evolving meta-ecosystem dynamics

Ecosystems characterized by relatively low species diversity and strong, well-defined coupling between ecosystems present tractable settings for developing the evolving meta-ecosystem framework. Identifying focal study organisms is critical, but even within simplified ecosystems,

160 this can be difficult. Thus, dominant or foundation species – including those that occupy or
161 move across ecosystem boundaries – that mediate the abundance and flow of energy, material,
162 and organisms offer a practical starting point (Whitham et al. 2003). Systems experiencing
163 ongoing and rapid environmental change offer further opportunities to observe relationships
164 between evolution and evolving meta-ecosystem dynamics.

165 The Arctic is well-suited for observing and understanding such dynamics. For example, the rate
166 of Arctic warming is four times faster than global averages (Rantanen et al. 2022), leading to
167 spatially structured selective pressures and large-scale shifts in vegetation, permafrost,
168 hydrology, and biogeochemical cycles (Frost et al. 2025) that provide opportunities to observe
169 evolutionary and ecological change in real time. The Arctic has a history of extensive long-term
170 ecological research and observatory networks that provide the temporal and spatial resolution
171 necessary to investigate multi-level dynamics. The Arctic’s tundra and stream-riparian meta-
172 ecosystems are each connected by strong coupling between biotic and abiotic processes and are
173 composed of a few key species that have disproportionately large ecosystem impacts, where
174 even subtle shifts in community or functional composition can trigger cascading effects at the
175 ecosystem level (Wookey et al. 2009). The expansion and increased growth of deciduous
176 shrubs, such as willows, birches, and alders along stream networks is one of the clearest
177 biological signals of Arctic warming (Myers-Smith et al. 2020; Tape et al. 2012) and underscores
178 the potential for rapid change to affect multiple dimensions of meta-ecosystem coupling.

179 As a dominant species at the boundary of stream-riparian zones, the widespread fletleaf willow
180 (*Salix alaxensis*) has an outsized influence on the flow of energy, materials, and organisms across
181 ecosystem boundaries. For example, taller, denser Arctic willows increase habitat and resources
182 for both terrestrial insects and birds (Boelman et al. 2014) and influence levels of organic matter
183 entering streams. Impacts to aquatic and terrestrial insect abundance may in turn affect trophic
184 subsidies for freshwater fish such as Arctic grayling (*Thymallus arcticus*) and migratory birds such
185 as white-crowned sparrows (*Zonotrichia leucophrys*), potentially altering the flow of energy,
186 materials, and organisms between terrestrial and aquatic ecosystems at both local and
187 continental scales. Fletleaf willow therefore provides an excellent opportunity to understand
188 how evolutionary changes in traits related to growth, phenology, and resource allocation could
189 have substantial cross-system impacts (Figure 1). By revealing how evolutionary responses in
190 dominant species can cascade across meta-ecosystem boundaries – affecting temperature
191 regimes, nutrient fluxes, organismal abundance and diversity – Arctic stream-riparian networks
192 provide a tractable model with principles that extend beyond polar environments, offering a
193 translational framework for studying evolving meta-ecosystem dynamics.

194 **Summary**

195 Darwin’s tangled bank describes a local network of interconnected organisms resulting from
196 ecological and evolutionary processes. An evolving meta-ecosystem perspective extends that
197 metaphor to spatially dynamic interactions with consequences and feedbacks at the landscape

198 scale, mediated by evolutionary and ecological dynamics in real time. This integrative and cross-
199 disciplinary framework therefore offers a generalizable approach to studying eco-evolutionary
200 dynamics in a rapidly changing world.

201 **Author Contributions**

202 B.M.L. drafted and revised the manuscript with input and assistance from I.N.C.S., L.A.D.,
203 M.H., C.N., M.C.U., J.L.W., and P.A.H.W. Authors L.A.D., M.H., C.N., M.C.U., J.L.W., and
204 P.A.H.W. led the conceptualization of the EVOME framework and funding acquisition.

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