

Coevolution is detected regionally but not locally in ecological communities

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1 Coevolutionary dynamics acting on both species and their interactions are a
2 key driving force behind the structure of ecological communities. The Geo-
3 graphic Mosaic Theory of Coevolution (GMTC) provides a spatial perspective
4 to these dynamics by proposing the existence of feedbacks between local and re-
5 gional scales. It remains unclear, however, how the structure of communities
6 at larger spatial scales either influences or is influenced by local coevolutionary
7 processes. Despite this, ecological networks are known to have an ecological
8 structure, which suggest that coevolution may play a role in their evolution-
9 ary dynamics. Here we show that the coevolutionary significance of individual
10 interactions is maintained when moving between the local and regional scale.
11 Importantly, this occurs despite the fact that community variation at the local
12 scale tends to weaken or remove community-wide coevolutionary signal. This
13 apparent mismatch between our interaction-level and community-level results
14 strongly suggests that interactions provide the correct scale to study coevolu-
15 tion at small spatial scales while communities are the relevant scale only at larger
16 spatial extents. We provide a new perspective on the interplay between coevolu-

1 tionary theory and community ecology, by establishing the organisational scales
2 at which the different theories have relevance. Although it has been tempting
3 so far to understand how coevolution relates to network structure, our results
4 suggest that the way forward is to understand how network structure may affect
5 coevolution over space instead.

6 Ecological interactions introduce selective pressures on the species involved. At large
7 organisational scales, this results in taxonomic boundaries delineating groups of in-
8 teracting species ¹, invariant structures in some ecological communities ², and con-
9 servatism of both the distribution of community modules ⁴ and the role of species
10 occupy within them ⁵. Although the evolutionary dynamics for a pair of interact-
11 ing species has been well described ⁶, attempts to understand how these mechanisms
12 cascade up to generate species diversity observed in large ecological networks have
13 been inconclusive ^{7,8}, despite the well known effect of antagonistic coevolution on
14 genomic diversification ^{9,10}. The scales at which these diversities happen are hard
15 to reconcile: coevolution is expressed within patches connected by gene-flow ¹¹⁻¹³,
16 whereas the species diversity of complex networks is typically observed at spatial
17 scales matching the species distribution ^{14,15}. Because these scales differ by orders of
18 magnitude, one must question the relevance of previous calls to scale the theory on
19 coevolution up to multi-species systems covering large spatial extents ¹⁶.

20 Network-based approaches ^{17,18}, on the other hand, were designed to study and de-
21 scribe species-rich systems. Previous empirical findings revealed the impact of evo-
22 lutionary dynamics on overall network structure in food webs ^{1,19}, pollination net-
23 works ^{20,21}, and host-parasite networks ^{22,23}. Both micro ^{3,24} and macro ^{25,26} evolu-
24 tionary dynamics have been measured or modelled in species-rich communities, but

1 there is no understanding of how, or even of whether, local/micro-evolutionary and
2 regional/macro-evolutionary feedback into one another. Another layer of complex-
3 ity is that ecological networks are known to vary in their structure over time and
4 space ²⁷: the same two species will not interact in a consistent way locally, either
5 because of local environmental contingencies, by chance, or because their phenolo-
6 gies do not positively covary in space ²⁸. As a consequence, *locally*, the evolutionary
7 signal on network structure is expected to be buried under much ecological noise,
8 and the effect of coevolution can only be inferred *regionally*.

9 In multi-species systems that typically span a large taxonomic range, coevolution is
10 often measured as the matching between the phylogenies of two sets of interacting
11 organisms ^{22,29}. This build on the century-old ideas that extant species interact in a
12 way similar to the way their ancestors did ³⁰. “Coevolved” systems should (i) have
13 approximately similar phylogenetic trees and (ii) species at matching positions in
14 either trees should interact. It is not clear, however, how this idea relates to dynamics
15 occurring at smaller scales ³¹: many ecological and evolutionary processes that occur
16 locally, or over small spatial scales, can disturb this expected structure. Notably, it
17 has been shown that species interactions are not consistent through space ^{27,32}. Local
18 loss of both interactions and species from the regional pool is most likely to result in
19 observed communities that do not appear to have been shaped by coevolution.

20 We use data on ectoparasites of rodents from Western to Eastern Europe ³³ to test
21 the following four hypotheses. First, local species assemblages do not show evidence
22 of coevolution even though the system as a whole does. Second, interaction-level
23 coevolutionary signal is conserved. Third, interaction-level coevolutionary signal
24 does not predict the spatial consistency of interactions. We do so by coupling two

1 novel methods: the *PACO* algorithm for detection of phylogenetic congruence ³⁴,
2 and a general framework for the variation of ecological networks ³².

3 Local observations on the 51 localities (*Supp. Mat. 1*) are aggregated into a regional
4 metanetwork ³², and the two phylogenetic trees have been rendered ultrametric (see
5 Suppl. Methods). We use *PACO* ³⁴ to measure the congruence between trees know-
6 ing the interactions. *PACO* yields a network-level significance value for the likeli-
7 hood that hosts and parasites have coevolved. For each local network, we measure
8 the strength of coevolution using (i) only local observations and (ii) all possible in-
9 teractions between local species (as known from the regional aggregation of all local
10 networks). This allows us to separate the effect of species sorting (regional interac-
11 tions) and interaction sorting (local interactions). At the regional scale, coevolution-
12 ary signal is extremely strong ($p \leq 10^{-4}$), as established by previous analysis of this
13 system ³⁵. Most local networks, on the other hand, show very little evidence of phy-
14 logenetic congruence. Out of 51 local networks, 35 show no signal of coevolution,
15 11 show coevolution when using the regional interactions, and 12 show coevolution
16 using the local interactions (see *Supp. Mat. 1* for network-level significance values).

17 **Figure?.**

18 This suggests that macro-evolutionary processes (such as co-diversification) have con-
19 sequences at the macro-ecological level ³⁶, but may not be detected at finer spatial
20 scales due to a stronger effect of ecological processes locally. *PACO* permits the anal-
21 ysis of *how strongly* each interaction contribute to coevolution, in a way that is as
22 independent as possible from other interactions. As interactions vary only insofar
23 that there are some locations in which they *do not* happen, we expect that the over-
24 all contribution of interactions will be the same in the local and regional networks.

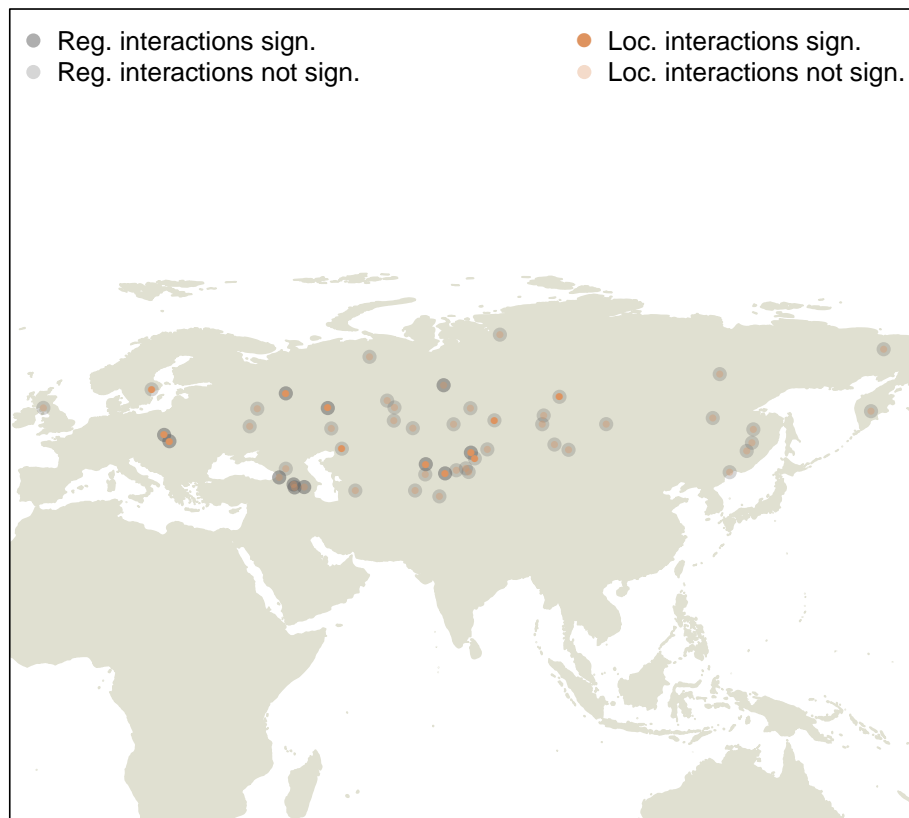


Figure 1: figure1

1 For the 5 networks that show evidences of coevolution accounting both for species
2 and interactions sorting, we measured the contribution of each interaction locally,
3 and compared it to its contribution to the regional network. Results are presented
4 in **FIG. one-sentence summary**. This is a key result, as it establishes that although
5 coevolution does not leave an imprint on local networks, it is still detectable in *in-*
6 *teractions*. This is in line with recent results that established that, although networks
7 are composed of interactions, both objects seem to have uncoupled behaviors ³⁷.

8 Species interactions vary in a way that is independent from species distribution ³².
9 One possible explanation is that species that have a strong (coevolutionary) rela-
10 tionship would either co-distribute more, or interact more frequently when they
11 co-occur. Should it be true, we would expect that *coevolved species pairs*, or in other
12 words, species involved in an interaction contributing strongly to the community-
13 wide coevolution, should be interacting frequently ³⁸. This would result in a positive
14 association between the frequency of the interaction (the number of observations
15 of a particular interaction divided by the number of observations of the tow species
16 together), and its overall importance for coevolution (here measured in the *regional*
17 network). As we report in **FIGURE**, we do not find this relationship – how strongly
18 an interaction contributes to overall coevolution does not predict how frequently it
19 will be realized when the two species are put together.

20 Our results, that (i) local networks show no signal of coevolution and (ii) the strength
21 of coevolution between two species does not predict how frequently they interact,
22 fall when in line with recent conclusions about the spatial dynamics of species inter-
23 actions. Species interactions vary according to ecological mechanisms ²⁷: local popu-
24 lation abundance ³⁹, local mis-matches of phenologies ⁴⁰, local micro-environmental

1 conditions ⁴¹. And even though network composition varies, the overall network
 2 *structure* remains constant over time ⁴², suggesting either (i) higher-order constraints
 3 or (ii) replacement of species by functionally equivalents from the regional pool.
 4 These result show that our current understanding of coevolution in multi-species in-
 5 teractions does not scale well to ecological questions – although phylogenetic struc-
 6 ture and interaction show a strong agreement at the regional scale, the structure of
 7 local communities remains largely driven by ecological constraints. The analysis
 8 of ecological networks has often focused on emerging properties ⁴³ rather than on
 9 the building blocks of the networks, that is species and interactions. Contrary to
 10 the often-argued point that coevolution should explain the local structure of interac-
 11 tions ⁴⁴, our result suggests that given the high variance in local interactions, coupled
 12 with the lack of relationship between coevolution and interaction frequency, local
 13 network structure is more likely to affect coevolution than the other way around.

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