

# Coevolution leaves a stronger imprint on interactions than on community structure

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November 28, 2014

1 Coevolutionary dynamics act on both species and their interactions  
2 in ways that shape ecological communities. It remains unclear, how-  
3 ever, how the structure of communities at larger spatial scales either  
4 influences or is influenced by local coevolutionary processes, and how  
5 mechanisms acting at these different scales feedback onto one another.  
6 Here we show that, though species interactions vary substantially over  
7 a continental gradient, the coevolutionary significance of individual in-  
8 teractions is maintained across different scales. Notably, this occurs  
9 despite the fact that observed community variation at the local scale  
10 frequently tends to weaken or remove community-wide coevolutionary  
11 signal. When considered in terms of the interplay between community  
12 ecology and coevolutionary theory, our results demonstrate that indi-  
13 vidual interactions are capable and likely to show a consistent signature  
14 of past coevolution even when woven into communities that do not.

15 Ecological interactions often exert important selective pressures on the species  
16 involved. For example, the phenologies of lodgepole pines and red crossbills re-

1 spond spatially to the presence of squirrels <sup>1</sup> and palm species undergo changes  
2 in seed morphology in response to the extinction of bird dispersing their seeds  
3 <sup>2</sup>. Given that interactions are distributed in similar ways across communities, at  
4 both the large <sup>3</sup> or small <sup>4</sup> scale, it can be argued that much ecological structure  
5 is the end result of evolutionary or coevolutionary dynamics between species <sup>5</sup>.  
6 Unfortunately, while the coevolutionary dynamic of pairs of interacting species  
7 has been well described at macro <sup>7</sup> and micro <sup>8</sup> evolutionary timescales, most  
8 attempts to understand how they cascade up to the levels of diversity of both  
9 species and interactions found within empirical communities have been inconclu-  
10 sive <sup>9</sup>. Moreover, because coevolutionary dynamics are often presented as a key  
11 driving force behind ecological structure across both time and space <sup>10</sup>, it is crucial  
12 to determine the scale at which they are both relevant and quantifiable.

13 Historically, the evidence for coevolution in taxonomically diverse communities  
14 is quantified as the degree of matching between the phylogenies of two sets of  
15 interacting organisms <sup>11</sup>. This notion builds on the century-old idea that extant  
16 species interact in a way similar to the way their ancestors did <sup>12</sup>. Yet it is consid-  
17 erably more restrictive than just phylogenetic conservation of species' interactions  
18 <sup>13</sup>, as it accounts for higher-order constraints. More explicitly, it is thought that  
19 communities that have assembled by successive divergence events should display  
20 phylogenetic congruence, that is (i) have similar phylogenetic trees and (ii) have  
21 species at matching positions in the trees that tend to interact <sup>???,14</sup>. On the  
22 other hand, many ecological and evolutionary processes that occur locally are  
23 expected to blur community-wide coevolutionary signal <sup>15</sup>. One possible explana-  
24 tion is that interactions can display substantial turnover at ecologically relevant  
25 temporal and spatial scales <sup>16</sup>: the same two species can interact in different ways

1 under the effect of local environmental contingencies, spatial mismatch in species  
2 phenologies, variations in population abundances, and chance events <sup>17</sup>. It is un-  
3 clear, however, whether these mechanisms influence how the coevolutionary signal  
4 within individual interactions should vary across spatial scales.

5 To answer these questions, we study a dataset of interactions between rodents  
6 and their ectoparasites from Western to Eastern Europe <sup>18</sup> (Methods Summary).  
7 This dataset is uniquely suited for this task as it represents a paradigmatic system  
8 in which species-species interactions are thought to be driven by macro-evolution  
9 and co-speciation events <sup>19</sup>, and coevolutionary signal is indeed significant at the  
10 continental level <sup>20</sup> ( $p \leq 10^{-4}$ ; Methods Summary). Importantly, it also provides  
11 spatial replication and variability <sup>21</sup> at a scale large enough to capture macro-  
12 ecological processes.

13 As host-macroparasites interactions are hypothesized to be both ecologically con-  
14 strained and evolutionary conserved <sup>22</sup>, the phylogenetic congruence observed at  
15 the continental level provides the baseline for expectations in local communities.  
16 Of course, if ecological mechanisms reduce coevolutionary signal, we should de-  
17 tect coevolution at the continental scale but not locally. Noting that variation  
18 of interactions can decrease congruence, we analyse the local data at two differ-  
19 ent levels to test these hypotheses; we first use *regional* interaction data, which  
20 accounts for different species composition across sites, then the *local* interaction  
21 data, which also accounts for variation in the interactions between observed these  
22 species (Methods Summary). Out of 51 local networks, 35 show no signal of  
23 coevolution, 11 show significant coevolutionary signal when using the regional  
24 interactions, and 12 show significant coevolutionary signal using the local inter-

1 actions (see *Supp. Mat. 1* for network-level significance values). These results  
2 would appear to support the idea that macro-evolutionary processes such as co-  
3 diversification can have consequences at the macro-ecological level <sup>23</sup> but may not  
4 in fact be detectable at finer spatial scales.

5 This is particularly apparent when comparing the distribution of network-level  
6 coevolutionary signal at the local and regional scales, relative to the continental  
7 scale (Fig. 2A). Yet this system-level difference says little about how individual  
8 interactions, on which most of the coevolutionary mechanisms purportedly act <sup>24</sup>,  
9 behaves. We show that regardless of the scale at which the system is observed,  
10 the distribution of the contribution of individual interactions to coevolution is  
11 conserved (Fig. 2B). This allows to understand why networks differ from the  
12 continental-level expectation: networks with interactions that are important for  
13 coevolution at the continental scale have more coevolutionary signal at the local  
14 and regional scale alike (Fig. 2C). This result shows that network-level property  
15 emerge from the properties of their interactions: because interactions differ in their  
16 total contribution to coevolution, their distribution across different networks will  
17 drive differences in overall coevolutionary signal.

18 Interactions also ultimately differ in how frequently they vary when the species  
19 involved co-occur <sup>???,25</sup>. Once more, the literature on host-parasite interactions  
20 usually assumes that the reason why some interactions are more frequent is be-  
21 cause they reflect a significant past history of coevolution <sup>26</sup>. If this were true,  
22 we should observe a significant, positive correlation between the probability of  
23 observing an interaction and the importance of that interaction for coevolution  
24 at the continental scale (Methods Summary). Surprisingly, we find that neither

1 is true here since interactions that are important for coevolution are not more  
2 conserved (Fig. 3).

3 Nonetheless, interactions that *are* observed at the local scale should retain their  
4 signal of contribution to coevolution because they involve species at matching po-  
5 sitions in the phylogenetic trees. Indeed, we find that interactions that contribute  
6 strongly to coevolutionary signal at the continental scale *also* show a significant  
7 tendency to contribute strongly at the local scale (Fig. 4). Remarkably, this result  
8 implies that the remnants of coevolution are still locally detectable in *individual*  
9 *interactions* even though it does not leave its imprint on most local networks.

10 Overall, the results of our analyses demonstrate that there is a sizeable gap be-  
11 tween our current understanding of coevolution as the basis of multi-species in-  
12 teractions and its applicability to ecological questions. Local networks show little  
13 to no signal of coevolution and the strength of coevolution between two species  
14 is a surprisingly poor predictor of how frequently they interact. In contrast to  
15 the frequent assumption that phylogenetic structure is a key driver of community  
16 structure<sup>27,28</sup>, these data reveal that this impact is actually minimal at ecolog-  
17 ically relevant spatial scales. Despite all the above, individual interactions are  
18 able to maintain their coevolutionary signal even when the community they are  
19 woven into does not. Thinking more broadly, these discrepancies provide a clear  
20 roadmap for bridging the aforementioned gap between our appreciation of the role  
21 of coevolution and its empirically measurable outcomes. Network structure is the  
22 most parsimonious *mechanism* by which coevolution proceeds, not the imprint  
23 coevolution leaves on ecological communities.

## 1 Methods Summary

2 We use data on observations of interactions between 121 species of rodents and  
3 205 species of parasitic fleas in 51 locations across Europe <sup>18</sup> to build 51 species-  
4 species interaction networks. Interactions were measured by combing rodents for  
5 fleas, a method that gives high quality data as it has a high power of detection. To  
6 account for differential sampling effort and across site variations in abundance, we  
7 only study the networks' incidence matrices (presence and absence of interactions).

8 In our study, we define three scales for the network data and analysis— conti-  
9 nental, regional, and local. The continental scale is the aggregated “metanetwork”  
10 which includes all potential interactions between co-occurring species <sup>16</sup> (*i.e.*, all  
11 species and all their interactions across the 51 networks). Within each site, the  
12 regional scale is given by the list of observed species and all their possible interac-  
13 tions. Hence the regional networks are always a perfect subset of the continental  
14 network. The local scale includes only the interactions that were actually observed  
15 in the field at a given site. Therefore, the local and regional networks always in-  
16 clude the same species, but the local network has only a subset (or, at most, an  
17 exact match) of the interactions in the regional network. The spatial consistency  
18 of every individual interaction is measured as the number of sites in which the  
19 two species involved co-occur.

20 We quantified the coevolutionary signal in terms of the degree of matching be-  
21 tween host and parasite phylogenies given knowledge of species interactions using  
22 the *PACO* method <sup>29</sup>, which is robust to variations in number of species. *PACO*  
23 provides measures of both the network-level congruence (*i.e.*, is the network co-  
24 evolved?) and the interaction-level signal (*i.e.*, what is the contribution of each

1 interaction to the overall coevolutionary signal?). We measured the phylogenetic  
2 dissimilarity between two sites for hosts and parasites using PCD<sup>30</sup>, a measure  
3 that accounts for the dissimilarity of species, corrected for the phylogenetic dis-  
4 tance between all species in the dataset. Since it is a requirement of the methods  
5 we use here, the phylogenetic trees for hosts and parasites were rendered ultra-  
6 metric (i.e., all species are at the same distance from the root).

## 7 References

8 **Acknowledgments.** We thank Juan Antonio Balbuena for discussions about the  
9 *PACo* method, and members of the Stouffer and Tylianakis groups for comments  
10 on an early draft of this manuscript. Funding to TP and DBS was provided  
11 by a Marsden Fund Fast-Start grant (UOC-1101) and to DBS by a Rutherford  
12 Discovery Fellowship, both administered by the Royal Society of New Zealand.

13 1.Benkman, C. W., Parchman, T. L., Favis, A. & Siepielski, A. M. *et al.* Reciprocal  
14 selection causes a coevolutionary arms race between crossbills and lodgepole pine.  
15 *Am. Nat.* **162**, 182–194 (2003).

16 2.Galetti, M. *et al.* Functional Extinction of Birds Drives Rapid Evolutionary  
17 Changes in Seed Size. *Science* **340**, 1086–1090 (2013).

18 3.Jordano, P., Bascompte, J. & Olesen, J. M. *et al.* Invariant properties in coevo-  
19 lutionary networks of plant-animal interactions. *Ecol. Lett.* **6**, 69–81 (2003).

20 4.Stouffer, D. B., Camacho, J., Jiang, W. & Amaral, L. A. N. *et al.* Evidence for  
21 the existence of a robust pattern of prey selection in food webs. *Proc. R. Soc. B*  
22 *Biol. Sci.* **274**, 1931–40 (2007).

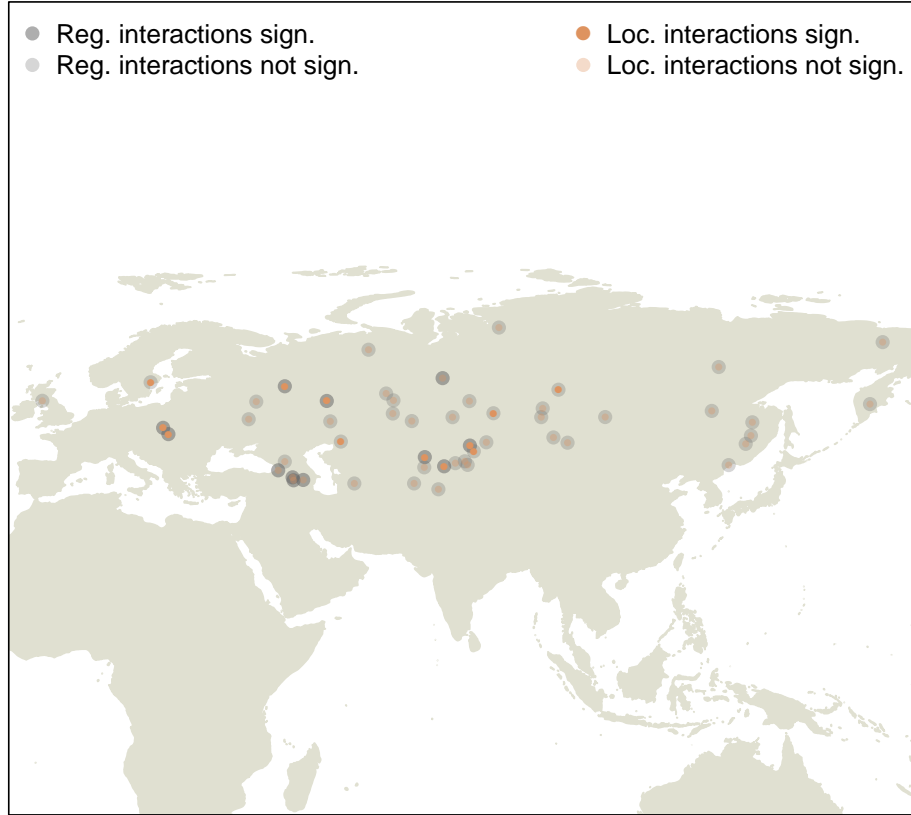


Figure 1: Spatial distribution of coevolutionary signal across the 51 sites. For each location, we indicate whether or nor the structure of regional and local interaction networks is consistent with phylogenetic congruence. The color of the circle corresponds to regionally significant or nonsignificant (foo and bar, respectively) while the symbol within corresponds to locally significant or nonsignificant (Zoroaster’s barsom and varza, respectively).



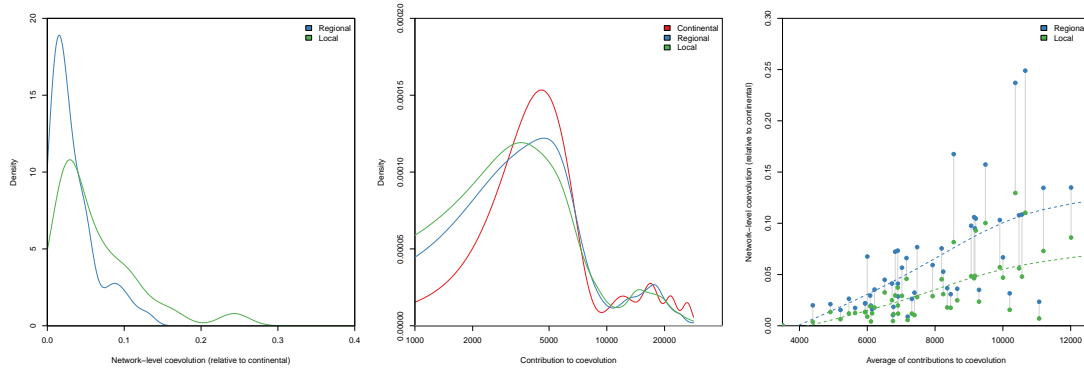


Figure 2: Distribution of coevolutionary signal at the network and interaction level. **A** Networks at the local and regional scale show less coevolutionary signal than the network at the continental scale. **B** Interactions in the local, regional, and continental scale have equal contributions to coevolution. **C** Networks that have lower coevolutionary signal are made of interactions that contribute little to coevolution at the continental scale.

- 1 5.Eklof, A., Helmus, M. R., Moore, M., Allesina, S. & Eklöf, A.*et al.* Relevance
- 2 of evolutionary history for food web structure. *Proc. R. Soc. B Biol. Sci.* **279**,
- 3 1588–1596 (2011).
- 4 6.Stouffer, D. B., Sales-Pardo, M., Sirer, M. I. & Bascompte, J.*et al.* Evolutionary
- 5 Conservation of Species' Roles in Food Webs. *Science* **335**, 1489–1492 (2012).
- 6 7.Van Valen, L. *et al.* A new evolutionary law. *Evol. Theory* **1**, 1–30 (1973).
- 7 8.Gandon, S., Buckling, A., Decaestecker, E. & Day, T.*et al.* Host-parasite co-
- 8 evolution and patterns of adaptation across time and space. *J. Evol. Biol.* **21**,
- 9 1861–1866 (2008).
- 10 9.Hembry, D. H., Yoder, J. B. & Goodman, K. R.*et al.* Coevolution and the
- 11 Diversification of Life. *The American Naturalist* **184**, 425–438 (2014).

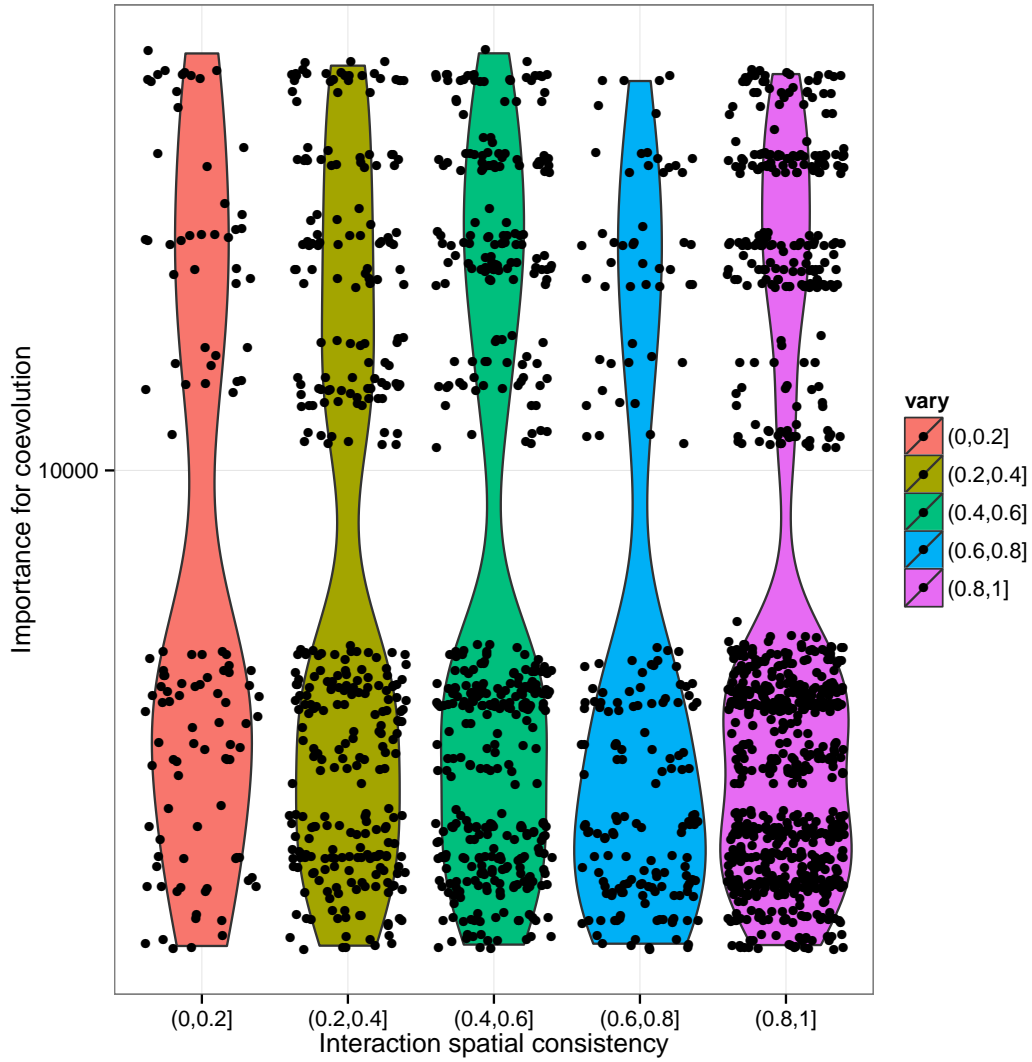


Figure 3: Lack of a relationship between the spatial consistency of an interaction and its importance for coevolution in the continental network. Spatial consistency is defined as the probability of observing an interaction between two species given that they were observed to co-occur.

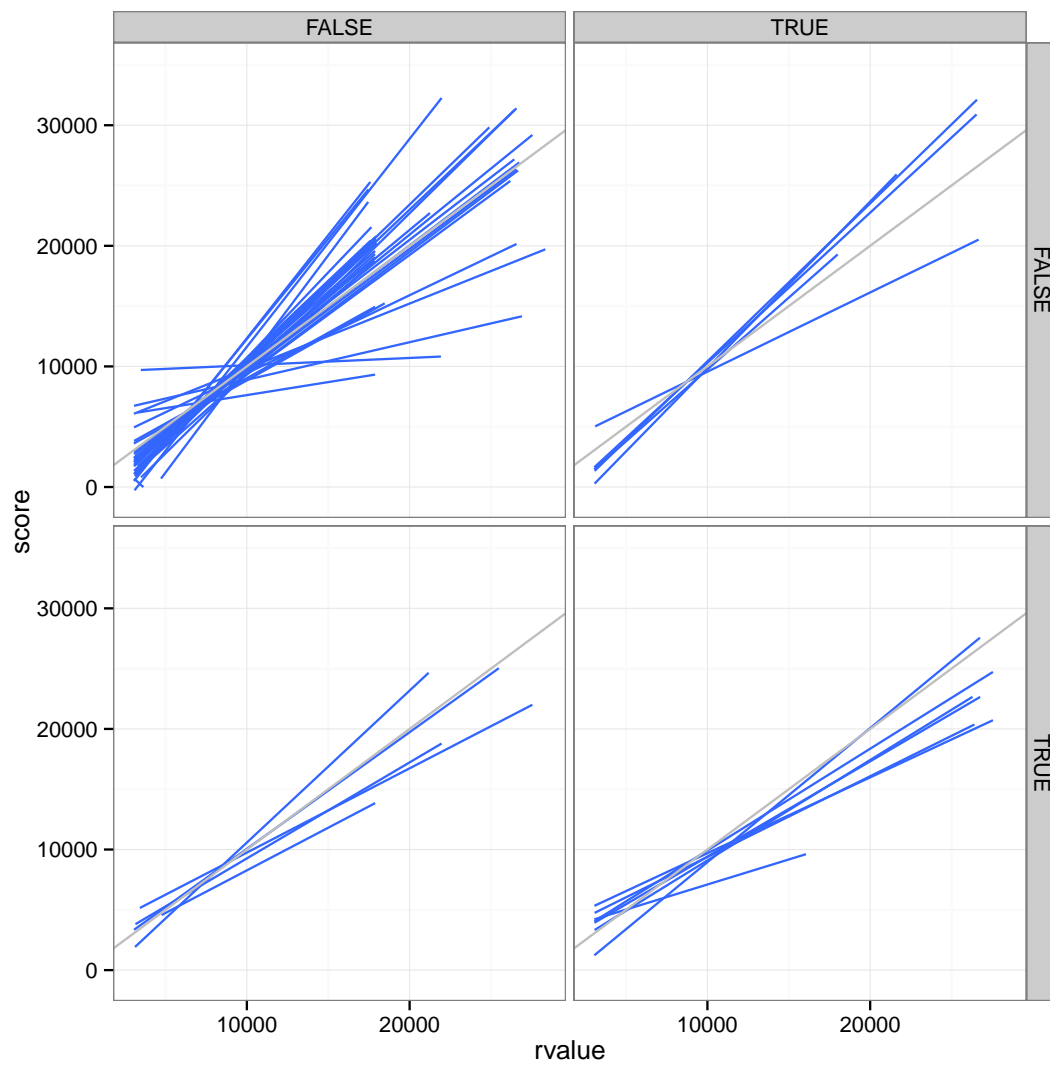


Figure 4: Relationship between continental and local contribution to coevolution. Each line corresponds to a location.

- 1 10. Thompson, J. N. *et al.* *The Geographic Mosaic of Coevolution*. (University Of  
2 Chicago Press, 2005).
- 3 11. Legendre, P., Desdevises, Y. & Bazin, E. *et al.* A statistical test for host-  
4 parasite coevolution. *Syst. Biol.* **51**, 217–234 (2002).
- 5 12. Fahrenholz, H. *et al.* Ectoparasiten und abstammungslehre. *Zool. Anz.* **41**,  
6 371–374 (1913).
- 7 13. Rezende, E. L., Lavabre, J. E., Guimarães, P. R., Jordano, P. & Bascompte,  
8 J. *et al.* Non-random coextinctions in phylogenetically structured mutualistic net-  
9 works. *Nature* **448**, 925–8 (2007).
- 10 14. Page, R. D. M. *et al.* *Tangled trees: Phylogeny, cospeciation, and coevolution*.  
11 (University of Chicago Press, 2003).
- 12 15. Poisot, T. *et al.* in *Evolutionary Ecology of Host-Parasite Systems* (eds. Morand,  
13 S., Littlewood, D. T. & Poulin, R.) (Cambridge University Press, 2015).
- 14 16. Poisot, T., Canard, E., Mouillot, D., Mouquet, N. & Gravel, D. *et al.* The  
15 dissimilarity of species interaction networks. *Ecol Lett* **15**, 1353–1361 (2012).
- 16 17. Poisot, T., Stouffer, D. B. & Gravel, D. *et al.* Beyond species: why ecological  
17 interaction networks vary through space and time. *Oikos* n/a–n/a (2014).
- 18 18. Data from: Phylogenetic signal in module composition and species connectivity  
19 in compartmentalized host-parasite networks. (2012).
- 20 19. Verneau, O., Du Preez, L. & Badets, M. *et al.* Lessons from parasitic flatworms  
21 about evolution and historical biogeography of their vertebrate hosts. *C. R. Biol.*  
22 **332**, 149–158 (2009).
- 23 20. Krasnov, B. R. *et al.* Phylogenetic Signal in Module Composition and Species

- 1 Connectivity in Compartmentalized Host-Parasite Networks. *Am. Nat.* **179**,  
2 501–511 (2012).
- 3 21.Krasnov, B. R., Mouillot, D., Shenbrot, G. I., Khokhlova, I. S. & Poulin, R.*et*  
4 *al.* Beta-specificity: the turnover of host species in space and another way to  
5 measure host specificity. *Int. J. Parasitol.* **41**, 33–41 (2011).
- 6 22.Combes, C. *et al.* *Parasitism - The Ecology and Evolution of Intimate Inter-*  
7 *actions.* (University Of Chicago Press, 2001).
- 8 23.Price, P. W. *et al.* *Macroevolutionary Theory on Macroecological Patterns.*  
9 (Cambridge University Press, 2003).
- 10 24.Thompson, J. N. *et al.* The raw material for coevolution. *Oikos* **84**, 5–16  
11 (1999).
- 12 25.Olito, C. & Fox, J. W. *et al.* Species traits and abundances predict metrics of  
13 plant–pollinator network structure, but not pairwise interactions. *Oikos* n/a–n/a  
14 (2014).
- 15 26.*Biogeography of host-parasite interactions.* (Oxford University Press, 2010).
- 16 27.Cavender-Bares, J., Kozak, K. H., Fine, P. V. A. & Kembel, S. W.*et al.* The  
17 merging of community ecology and phylogenetic biology. *Ecol. Lett.* **12**, 693–715  
18 (2009).
- 19 28.Mouquet, N. *et al.* Ecophylogenetics: advances and perspectives. *Biol. Rev.*  
20 **87**, 769–785 (2012).
- 21 29.Balbuena, J. A., Míguez-Lozano, R. & Blasco-Costa, I.*et al.* PACo: A Novel  
22 Procrustes Application to Cophylogenetic Analysis. *PLoS ONE* **8**, e61048 (2013).

- 1 30. Ives, A. R. & Helmus, M. R. *et al.* Phylogenetic Metrics of Community Simi-  
2 larity. *The American Naturalist* **176**, E128–E142 (2010).