

1 **A Community-Trait-Phylogenetic Framework: Ecological and Evolutionary Integration for**
2 **Soil Microarthropod Assembly**

3

4 **Ting-Wen Chen***

5 J.F. Blumenbach Institute of Zoology and Anthropology, University of Göttingen, Göttingen,
6 Germany

7 Biology Centre of the Czech Academy of Sciences, Institute of Soil Biology and Biogeochemistry,
8 České Budějovice, Czech Republic

9 Department of Life Sciences, National Chung Hsing University, Taichung, Taiwan

10 *Correspondence: tchen2@gwdg.de

11 **Abstract**

12 Why does a single square meter of forest soil harbour thousands of animal species? Fifty years
13 after Jonathan M. Anderson raised this question, soil ecology still struggles with a fragmented
14 view on the coexistence of species. Researchers often study taxonomy, functional traits, and
15 phylogeny in isolation. Each approach adds insight but leaves gaps in the picture of soil
16 biodiversity.

17 In this paper, I propose a Community-Trait-Phylogenetic Ecology framework that integrates
18 evolutionary and ecological perspectives to explain how soil animal communities form and
19 persist. The framework combines three research fields:

- 20 - Biogeography – describes species composition across local, regional, and global scales.
- 21 - Functional traits – divided into α -niche traits (resource use) and β -niche traits (environmental
22 tolerance), showing whether resource partitioning or filtering by environment drives community
23 assembly.
- 24 - Phylogeny – shapes trait expression and defines the pool of species.

25 Evidence from the dominant soil microarthropods, springtails (Collembola, Hexapoda) and
26 oribatid mites (Oribatida, Acari), shows the value of this framework. Global data synthesis
27 reveals a mismatch between density and diversity, which challenges traditional biogeographic
28 predictions. Trait analyses show that environmental filtering occurs at global scales. At regional
29 and local scales, cryptic species that diverged millions of years ago coexist with distinct habitat
30 preferences. In addition, ancient and recent lineages coexist across elevations. Morphological
31 and physiological traits usually follow phylogenetic constraints. In contrast, trophic traits show
32 high flexibility, which allows closely related species to coexist.

33 This integrative view shifts soil animal ecology from describing patterns to understanding the
34 mechanisms responsible for them. It also supports predictions of community responses to
35 climate change and land-use change. Finally, it can guide conservation strategies for soil
36 habitats that protect species, functional, and evolutionary diversity of soil biota.

37
38 **Keywords:** coexistence mechanisms; community assembly; environmental filtering; evolution;
39 ecology; functional traits; niche; phylogenetic comparative methods; resource partitioning; soil
40 arthropods; soil biodiversity; soil macroecology; species pool

41 Fifty years ago, British ecologist Jonathan M. Anderson posed a deceptively simple yet
42 profoundly difficult question: *Why can a single square meter of forest soil harbour thousands of*
43 *animal species comprising millions of individuals* (Anderson, 1975)? This question challenges
44 our understanding of biodiversity and species coexistence mechanisms, revealing the complex
45 and delicate balance of natural forces.

46

47 **The Astonishing Complexity of Soil Life**

48 In a temperate deciduous forest just one square meter of litter and soil contains 10,000 to
49 200,000 tiny animals, often overlooked by the naked eye. These animal communities comprise
50 roughly 60-200 mesofauna species, each measuring about 0.2mm to 2mm in width (Petersen
51 and Luxton, 1982). Unlike microfauna such as nematodes that inhabit water films, mesofauna
52 like microarthropods occupy the air-filled pore space of the soil matrix. This distinct porosphere
53 subjects them to unique selection pressures on morphology and environmental tolerance. Within
54 soil pores and litter, springtails (Collembola, Hexapoda) and oribatid mites (Oribatida, Acari)
55 stand out for their remarkable abundance and ecological importance. These two taxa account
56 for approximately 95% of global soil arthropod abundance (Rosenberg et al., 2023) and rank
57 among the most diverse soil animal taxa with over 9000 and 11,000 described species,
58 respectively (Potapov et al., 2020; Subías, 2022). Despite sharing the same habitat and feeding
59 on similar resources, hundreds of these species coexist without apparent conflict. This is a
60 phenomenon that traditional soil ecology cannot fully explain. How is it possible? The answer
61 requires understanding of soil biodiversity from three complementary perspectives.

62

63 **Limitations and Breakthroughs of Traditional Approaches**

64 **Community ecology and biogeography** take species as the units. They focus on how
65 environmental factors, such as temperature, moisture, and soil pH, correlate with species
66 occurrence, and how species interactions and dispersal limitations shape local community
67 composition (Potapov et al., 2023). Although precise, this taxonomy-based approach is labour-
68 intensive and time-consuming. Species determination requires considerable training. It is also
69 defective because, though it indicates which species adapt to which environments (i.e. habitat

70 niches), it struggles to explain how they adapt to the environment and how species achieve
71 coexistence. Thus, the traditional species-centred perspective describes the patterns of
72 coexistence but not the mechanisms responsible for coexistence.

73 **Functional trait ecology** brought a breakthrough. Rather than treating each species as an
74 independent entity, it identifies niche dimensions based on specific attributes of the organisms
75 (i.e. traits) (Winemiller et al., 2015). Different species differ in multiple traits, such as morphology
76 (body size, colouration), life history (reproductive strategy, development rate), and resource
77 utilization (diet). These traits are assumed to be functional and to affect their fitness (but see
78 Gould and Lewontin, 1979). Through analysing functional traits, it is possible to predict which
79 species are able to coexist and how communities change along environmental gradients
80 (Brousseau et al., 2018). However, even this perspective has shortcomings, as it typically
81 overlooks the influence of evolutionary legacy on traits. Traits do not appear from nowhere but
82 are the result of millions of years of evolution (van Straalen, 2021).

83 **Phylogenetic and comparative methods** reveal the deep shared evolutionary history of
84 species. The evolutionary past of species influences which traits they possess and how
85 ecological roles are constrained. Related species tend to resemble each other, and such a
86 pattern is called "phylogenetic signal" (Silvertown et al., 2005). However, until today evolutionary
87 perspectives are little considered by soil ecologists as it requires specific conceptual and
88 practical skills. As a consequence, soil animal research typically treats ecology and evolution as
89 separate domains (but see Ponge, 2020 and van Straalen, 2021).

90

91 **An Integrative Perspective: The Community-Trait-Phylogenetic Ecology** 92 **Framework**

93 To overcome the limitations of the three traditional approaches and to better understand soil
94 animal communities, I propose a "Community-Trait-Phylogenetic Ecology" (CTPE) framework
95 (**Figure 1**). This framework attempts a more holistic understanding of the complex mechanisms
96 driving the composition of soil animal communities.

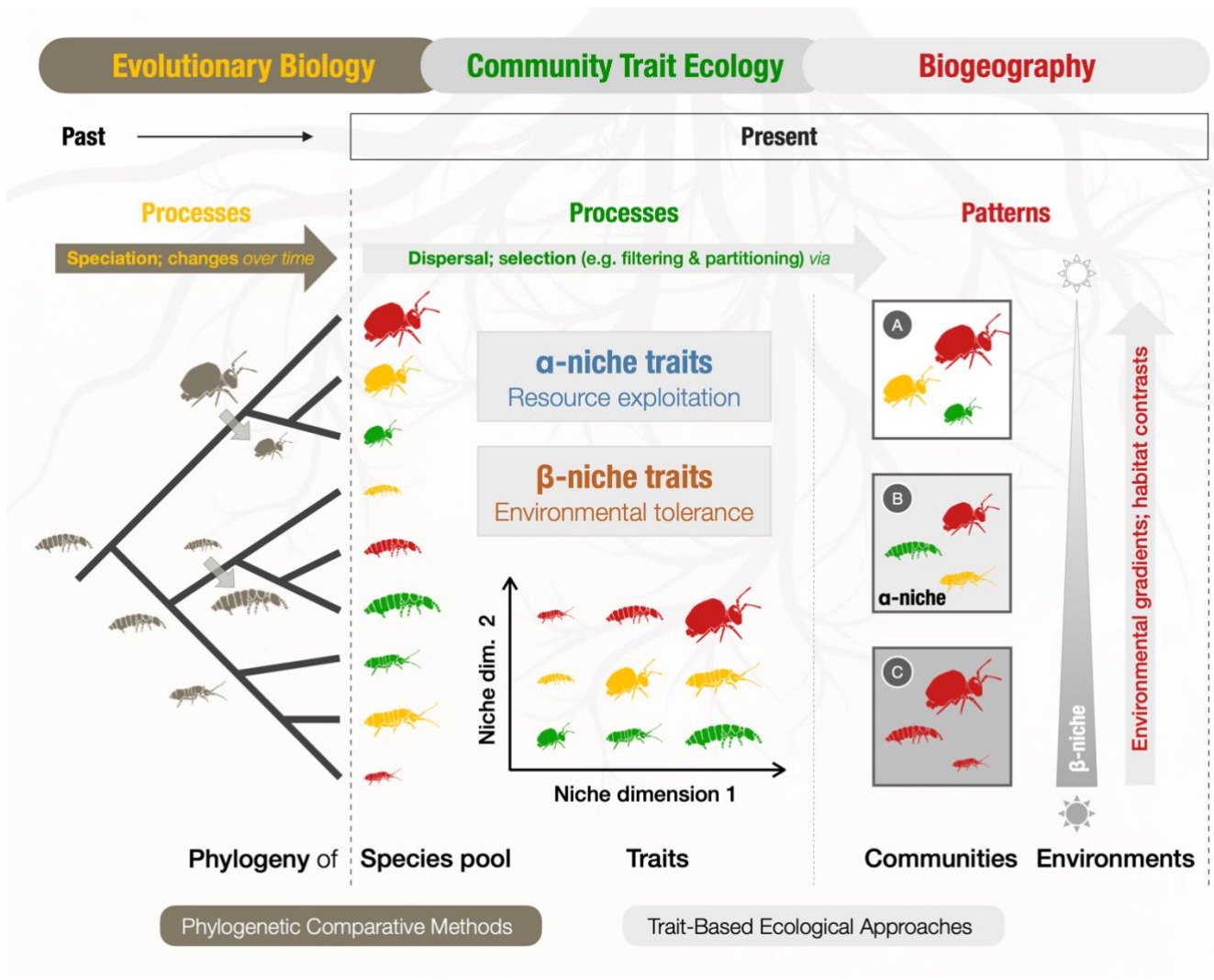


Figure 1. The "Community-Trait-Phylogenetic Ecology" (CTPE) Framework: Ecological and Evolutionary integration for Soil Animal Diversity. The CTPE framework integrates ecological and evolutionary processes to understand mechanisms driving soil biodiversity across spatial scales. The spatial distribution of soil communities is the central focus of biogeography (right, red panel). Even when species richness remains constant, species composition often changes along environmental gradients, leading to local communities that differ in their trait composition (for example, body size, colour and shape). Functional traits reflect multidimensional ecological niches and help explain species occurrence in particular habitats. For examples, community A occurs at high elevation where solar radiation is strong, while community C is located at low-elevation forest cover with dense canopy cover. By examining functional traits, defined as α -niche traits (associated with resource use and partitioning) and β -niche traits (related to environmental tolerance), this framework can infer the processes shaping community patterns and predict community responses to environmental changes (middle, green panel). In community A, springtail species share a similar body shape

but differ in body size and colour, whereas in community C, species are similar in colour but differ in the other two traits. Patterns in trait distributions, when compared to null expectations, reveal the underlying deterministic processes of community assembly. With phylogenetic comparative methods, the CTPE framework also infers the evolutionary processes that have generated present-day trait variation among species, offering a historical context for contemporary ecological patterns (left, yellow panel). Over evolutionary time, species have diversified from common ancestors by shifting traits (for example, from large to small body size, or *vice versa*). Speciation and trait evolution have thus shaped the regional species pool from which local communities assemble under the combined influence of environmental filtering and niche partitioning.

This framework comprises three elements.

The first element is the **species level - multiscale variation in community composition**. Variations in soil animal community is reflected not only in differences in species richness but also in species composition and their distribution across space and time (biogeographic patterns; e.g. Gao et al., 2014; Potapov et al., 2023; Junggebauer et al., 2024). To understand changes in soil animal diversity, it is necessary to ask what patterns characterize communities at local, regional, and global scales.

The second element is the **functional level - two niche processes reflected by traits**. Functional traits of species may be divided into two categories, α - and β -niche traits (Ackerly and Cornwell, 2007). Each category refers to different yet complementary community assembly processes (Chen et al., 2017; Noske et al., 2024). The β -niche traits reflect species environmental tolerances, such as body surface pigmentation (affecting thermoregulation and UV resistance; Xie et al., 2022) and temperature-moisture tolerance (Janion-Scheepers et al., 2018). Coexisting species typically resemble each other in these traits, indicating that they tolerate similar environmental conditions. By contrast, the α -niche traits reflect resource utilization strategies of species. Despite sharing the same habitat in soil, springtails and oribatid mites may exploit different food resources, including plant roots and exudates, organic matter, bacteria, fungi, lichens, mosses, algae, and even other soil animals (Potapov et al., 2022). Differences in the use of these resources are likely to reduce competition, thereby allowing to coexist locally. Therefore, the distinction between β - and α -niche traits helps predict which species are able to coexist and how communities change along environmental gradients.

141 Third is the **phylogenetic level – evolutionary constraints and possibilities**. Evolutionary
142 history not only provides context for trait variation but also constrains ecological possibilities.
143 Combining molecular techniques (such as mitochondrial genome sequencing; Xie et al., 2022)
144 with comparative methods (such as phylogenetic signal testing; Revell, 2024) makes it possible
145 to determine the nature of traits. These methods allow to identify which traits are of ancient
146 evolutionary origin and which evolved recently in response to contemporary selection processes,
147 or which traits are novel or have changed their original functions so that species exploit novel
148 niches. Morphological and physiological traits typically show strong phylogenetic signal;
149 conversely, trophic traits (resource utilization) often lack phylogenetic signal (Chen et al., 2017;
150 Gong et al., 2018; Xie et al., 2022; Noske et al. 2024). These patterns suggest that environmental
151 tolerance is evolutionarily constrained and that resource utilization strategies have evolved
152 independently multiple times, allowing closely related species to partition resources and coexist.

153

154 **Evidence from Springtails and Oribatid Mites: from Global to Local**

155 The international collaboration, the Global Collembola Initiative (#GlobalCollembola; Potapov et
156 al., 2020), compiled nearly 3000 community composition records. These records cover eight
157 biogeographic regions and 10 biomes from the tropics to the poles (Potapov et al., 2024). This
158 unprecedented dataset reveals a striking mismatch between density and diversity. Polar regions
159 show high density but only moderate species richness, temperate forests are characterized by
160 moderate density yet highest richness, tropical regions show lowest density but highest richness,
161 and arid systems show both low density and low richness (Potapov et al., 2023). This "density-
162 diversity mismatch" challenges traditional perspectives.

163 Recent compilation of data by the Global Collembola Initiative on 10 different traits from over
164 7000 springtail species and the ongoing global-scale trait distribution analyses, provide evidence
165 for the effect of environment on traits. The compilation indicates that springtail body pigmentation,
166 colour patterns, ommatidia number, furca development, and body size are differentially
167 influenced by habitat type, latitude, biome, and local density (Chen, T.-W. et al., unpubl. data).
168 This suggests that environmental filtering operates at global scales, selecting particular trait
169 combinations.

170 At the regional level, recent biogeographic research on soil animals of Changbai Mountain in
171 Northeast Asia tracked variation in isotomid springtail communities across 1400m of elevation
172 (Xie et al., 2022). This study found that soil nitrogen content, changing with elevation, acts as a
173 key environmental filter selecting particular pigmentation patterns and trait combinations. This
174 mountain study also raises an evolutionary puzzle: are mountains "cradles" of speciation? Or
175 are they "museums" preserving ancient lineages? Linking phylogeny with geological events
176 reveals that both scenarios apply to isotomid springtails. Ancient lineages persist while new
177 species have emerged alongside mountain uplift. These patterns provide the basis for inferring
178 historical speciation and diversification processes that shaped the current species pools from
179 which local communities are assembled (Vasconcelos et al., 2022).

180 At local scales, ongoing research on the winter springtail communities in the wetlands,
181 secondary forests, farmlands, and plantations of northeastern China, revealed different
182 dominant community assembly processes across habitats (Chen, T.-W., unpubl. data). Even
183 within a single community, species are simultaneously influenced by both filtering and
184 partitioning processes, depending on the traits considered. For example, compared to forests,
185 farmland springtails show similarity in furca and eye traits presumably due to environmental
186 filtering, while body size, pigmentation, and colour patterns reveal niche partitioning among
187 coexisting species. This suggests how trait-based perspectives on two niche processes can
188 operate simultaneously within the same community.

189

190 **The Multidimensional View of Trophic Ecology: Soil Animal Diets are** 191 **Complex**

192 Even though morphological traits can reflect partitioning processes, traits more directly related
193 to the use of food resources (α -niche) are usually derived from food-web methods that
194 characterize soil animal diet or food resources (i.e. trophic niches; Potapov et al., 2021). Gut
195 contents, digestive enzymes, neutral lipid fatty acids, and stable isotopes (^{15}N and ^{13}C), each
196 provide complementary information on the diet of consumers. Most pairwise correlations of
197 trophic niche parameters deduced from these methods are weak, indicating that each method
198 captures different dimensions of the trophic niche (Potapov et al., 2021). Stable isotope values

and gut microbiota of winter-active springtails show that they actually feed on resources on snow cover (such as cyanobacteria) rather than from litter (Hao et al., 2020). These results indicate that by combining multiple methods can we truly understand soil animal diets and reveal their realized trophic niches.

Furthermore, microbiota associated with soil animals can be viewed as a special type of trait (Gong et al., 2018, 2022; Hao et al., 2025). The study on bacteria and fungi associated with oribatid mites reveal subtle differences in how evolutionary history and trophic ecology of oribatid mites influence their microbial communities (Gong et al., 2018). Variations in fungal communities are better explained by stable isotope differences than by phylogenetic distance between mite species, suggesting fungi as food resources. Bacterial communities, in contrast, are more strongly related to host phylogeny than to trophic niche differences, indicating that closely related mites harbour similar bacteria, presumably shaped by host physiology. This suggests that mite-bacteria associations evolved interdependently. The ancient coevolution between animal hosts and gut symbionts results in a phylogenetic signal of bacterial communities across soil oribatid mite species.

Neutral lipid fatty acids also reveal the distinction between trait evolutionary dependence and resource flexibility (Chen et al., 2017). In this study we measured neutral lipid fatty acid composition in springtails and found closely related species to share similar fatty acid profiles. Long-chain polyunsaturated fatty acids, which are related to physiological function (β -niche), show a strong phylogenetic signal, while fatty acid biomarkers representing food resources (bacteria, fungi, and plants; α -niche) show almost no signal. This means that β -niche physiological functions are evolutionarily constrained, whereas α -niche resource utilization is highly flexible, indicating closely related species evolve different feeding strategies.

From Deep Time to Present: Multiscale Coupling of Evolution and Ecology

The CTPE framework reveals how evolution and ecology interact across different temporal scales. Springtails and oribatid mites have persisted since the Paleozoic (Schaefer et al., 2010; Yu et al., 2024). At hundred-million-year scales, they diversified alongside plant evolution and geological events (such as continental drift and mountain building), forming regional species

pools (Schaefer and Caruso, 2019; Xie et al., 2022). At million-year scales, even cryptic species indistinguishable morphologically have long diverged at the genetic level (Heethoff et al., 2007; Zhang et al., 2018). For example, three lineages of the springtail *Lepidocyrtus lanuginosus* diverged about 15.9 to 9.7 million years ago (Miocene), yet still coexist in the same region today, each preferring different habitats (forest, grassland, or farmland; Zhang et al., 2018). Different lineages of the oribatid mite *Oppiella nova* likewise occupy distinct ecotones, having diverged about 16 to 6 million years ago (von Saltzwedel et al., 2014). These studies demonstrate that both persistence of old lineages and environmental filtering are at work. At contemporary ecological timescales, in addition to long evolutionary paths, present community composition remains highly determined by current environmental conditions and resource availability. Evolution provides the "toolbox" (traits), while contemporary ecological conditions determine which "tools are selected" (species coexistence). This phenomenon indicates a close evolutionary-ecological coupling in soil animals, where species that diverged in deep time now coexist through particular mechanisms. Trait evolution enables environmental (β -niche) differentiation at large spatial and long evolutionary time scales but resource (α -niche) partitioning at both small spatial and temporal scales.

Insights and Significance of the CTPE Framework

This integrative framework brings several novel insights to the explanation of soil biodiversity. First, traits reveal ecological processes. Measuring functional traits is not merely for describing patterns but reveal which ecological processes dominate in communities. Environmental filtering is usually revealed by similar β -niche traits, while partitioning processes can be shown in different resources (α -niche) used by coexisting species. Second, evolutionary history reveals constraints and opportunities. Physiological functions are usually evolutionarily constrained and exhibit a phylogenetic signal. In contrast, resource utilization strategies are less evolutionarily constrained. They usually display flexibility between species (e.g. convergent evolution; Schaefer and Caruso, 2019) and even within species (e.g. trophic plasticity; Krause et al., 2019; Yang et al., 2025). In combination, contemporary ecological processes reshape soil microarthropod communities via existing traits and coexistence arises from multiple overlapping mechanisms. Evolution adds trait diversity, the environment selects particular traits, and resource differentiation enables local

diversity. Importantly, this integrative framework can extend beyond soil microarthropods to explain diversity patterns across multiple scales, systems, and taxa (Junker et al., 2022; Luza et al., 2023). Soil animals are far more complex than previously recognized. They occupy diverse niches, display remarkable adaptability, and utilize various resources. They are not merely microscopic detritivores but multitaskers with specific effect traits (Bonfanti et al., 2025), ensuring ecosystem functioning through redundancy and complementarity.

Future Applications: from Understanding to Prediction and Conservation

By integrating species distribution and community patterns, multiple trait measurements, and phylogenetic analyses, we are making progress toward understanding the enigma Jonathan M. Anderson posed fifty years ago (Anderson, 1975). This understanding is essential for predicting consequences of global change: Climate change alters temperature and moisture regimes, thereby changing directions and strengths of environmental filtering and selecting different trait combinations (Ferrín et al., 2023). Land-use change may fragment communities and limit connectivity (Susanti et al., 2021). Invasive species may disrupt the existing functional and phylogenetic structures of communities (Janion-Scheepers et al., 2018). Understanding which traits confer ecosystem resistance and resilience (Bonfanti et al., 2022), and which are shaped by evolution (Noske et al., 2024), enables the prediction of structural and functional changes of soil animal communities under these scenarios. This allows to identify key species and vulnerable taxa, thereby providing the basis for optimizing conservation strategies that maintain multidimensional diversity. The CTPE framework, thus, also emphasizes that protecting soil functioning is not just a matter of species numbers but requires maintaining multiple levels of diversity including trait diversity, functional diversity, and evolutionary diversity (Véron et al., 2019). Insights gained from the CTPE framework are useful for global soil conservation policy.

Why This Matters

Soil animals differ in their sensitivity to environmental stress, making them valuable bioindicators of ecosystem health (Shimano, 2011; Yin et al., 2020). They contribute to essential ecosystem functions by regulating nutrient cycling, supporting plant production, and mediating carbon

sequestration (Bardgett and van der Putten, 2014; Bonfanti et al., 2025). Fifty years after Anderson's original formulation, researchers now have new tools, new theories, and new frameworks to solve this enigma. Advances in high-throughput sequencing, stable isotope analysis, fatty acid and amino acid profiling, and computational methods now enable unprecedented data integration and synthesis. The coming decade of soil biodiversity research will elucidate the mechanisms that generate and maintain soil biodiversity and help predict how it responds to climate change, land-use alteration, and other global pressures. Now we can integrate ecological processes, functional traits, and evolutionary history to better understand and predict soil biodiversity. Soil will no longer be a black box but a kaleidoscope revealing the hidden wealth of life beneath our feet (Andr  n and Balandreau, 1999; van Straalen, 2023).

Concluding Remarks

- The integrated "community-trait-phylogenetic" framework depicts soil biodiversity as a multidimensional pattern shaped by ancient evolutionary processes and maintained by contemporary ecological processes. Community coexistence depends on multiple mechanisms operating in parallel: environmental filtering, resource differentiation, and evolutionary constraints and novelty.
- Past environmental conditions shaped ancient lineages with specific traits and their descendants now display unique preferences in contemporary habitats.
- β -niche traits (such as morphological and physiological traits which reflect environmental tolerance) often show a strong phylogenetic signal, indicating evolutionary constraints. Coexisting species in communities typically are characterized by trait similarity, reflecting contemporary environmental filtering.
- α -niche traits (such as multidimensional trophic niche parameters which reflect resource utilization) can be revealed through complementary diet methods. These traits typically show no, or a weak, phylogenetic signal, highlighting ecological opportunity.
- Microbiota associated with soil animals can be viewed as functional traits. Bacteria (especially symbionts) are closely tied to host evolutionary history, while fungi serve primarily as food

resources, reflecting the trophic niches of the animals.

- Integrating knowledge of community ecology, functional traits, and evolutionary history helps predict soil biodiversity responses to global change and their effects on ecosystem functioning.

Acknowledgments

I thank Stefan Scheu, Mark Maraun, Ina Schaefer, Anton M. Potapov, Donghui Wu, Meixiang Gao, Johanna Elisabeth Noske, Md Ekramul Haque, Zhijing Xie, Isma Dwi Kurniawan, Xin Gong, Cao Hao, Yan Zhang, and Jo-Fan Chao for their inspiration, discussions, and comments, Svenja Meyer for the springtail silhouettes and Andrew J. Davis (English Experience Language Services, Göttingen, Germany) for improving the English of the manuscript.

Funding

The idea of the paper was developed with support from BEST (Bioenergieregionen stärken) Project, Federal Ministry of Education, Science and Technology (BMBF, Bundesministerium für Bildung und Forschung), Germany; Government Scholarship to Study Abroad, Ministry of Education, Taiwan; CAS-DAAD mobility project (DAAD-19-10/57448388) co-funded by Czech Academy of Sciences (CAS) and German Academic Exchange Service (DAAD, Deutscher Akademischer Austauschdienst); MSM project for research and mobility of starting researchers (MSM200962001) from Czech Academy of Sciences; Short-Term Scientific Mission (STSM) under the European Cooperation in Science & Technology (COST) Action “European Soil-Biology Data Warehouse for Soil Protection” (CA18237); the collaborative German-Indonesian research project CRC990/EFForTS (192626868–SFB 990), German Research Foundation (DFG, Deutsche Forschungsgemeinschaft); and DFG Priority Program 1374 “Biodiversity-Exploratories” (SCHE 376/38-2).

References

Ackerly, D.D., Cornwell, W.K., 2007. A trait-based approach to community assembly: partitioning

of species trait values into within- and among-community components. *Ecology Letters* 10,
135–145. doi:10.1111/j.1461-0248.2006.01006.x

Anderson, J.M., 1975. The enigma of soil animal species diversity, in: Vaněk, J. (Ed.), *Progress in Soil Zoology: Proceedings of the 5th International Colloquium on Soil Zoology*. Springer Netherlands, Dordrecht, pp. 51–58. doi:10.1007/978-94-010-1933-0_5

Andrén, O., Balandreau, J., 1999. Biodiversity and soil functioning - From black box to can of worms? *Applied Soil Ecology* 13, 105–108. doi:10.1016/S0929-1393(99)00025-6

Bardgett, R.D., van der Putten, W.H., 2014. Belowground biodiversity and ecosystem functioning. *Nature* 515, 505–511. doi:10.1038/nature13855

Bonfanti, J., Hedde, M., Cortet, J., Krogh, P.H., Larsen, K.S., Holmstrup, M., 2022. Communities of Collembola show functional resilience in a long-term field experiment simulating climate change. *Pedobiologia* 90, 150789. doi:10.1016/j.pedobi.2022.150789

Bonfanti, J., Potapov, A.M., Angst, G., Ganault, P., Briones, M.J.I., Calderón-Sanou, I., Chen, T.-W., Conti, E., Degrune, F., Eisenhauer, N., Ferlian, O., Hackenberger, D., Hauer, A., Hedde, M., Hohberg, K., Krogh, P.H., Mulder, C., Perez-Roig, C., Russell, D., Shelef, O., Zhou, Z., Zuev, A.G., Berg, M.P., 2025. Linking effect traits of soil fauna to processes of organic matter transformation. *Functional Ecology* 39, 446–461. doi:10.1111/1365-2435.14720

Brousseau, P.M., Gravel, D., Handa, I.T., 2018. On the development of a predictive functional trait approach for studying terrestrial arthropods. *Journal of Animal Ecology* 87, 1209–1220. doi:10.1111/1365-2656.12834

Chen, T.-W., Sandmann, P., Schaefer, I., Scheu, S., 2017. Neutral lipid fatty acid composition as trait and constraint in Collembola evolution. *Ecology and Evolution* 7, 9624–9638. doi:10.1002/ece3.3472

Ferrín, M., Márquez, L., Petersen, H., Salmon, S., Ponge, J.-F., Arnedo, M., Emmett, B., Beier, C., Schmidt, I.K., Tietema, A., de Angelis, P., Liberati, D., Kovács-Láng, E., Kröel-Dulay, G., Estiarte, M., Bartrons, M., Peñuelas, J., Peguero, G., 2023. Trait-mediated responses to aridity and experimental drought by springtail communities across Europe. *Functional*

- Ecology 37, 44–56. doi:10.1111/1365-2435.14036
- Gao, M., He, P., Zhang, X., Liu, D., Wu, D., 2014. Relative roles of spatial factors, environmental filtering and biotic interactions in fine-scale structuring of a soil mite community. *Soil Biology and Biochemistry* 79, 68–77. doi:10.1016/j.soilbio.2014.09.003
- Gong, X., Chen, T.W., Zhang, L., Pižl, V., Tajovský, K., Devetter, M., 2022. Gut microbiome reflect adaptation of earthworms to cave and surface environments. *Animal Microbiome* 4, 47. doi:10.1186/s42523-022-00200-0
- Gong, X., Chen, T.-W., Zieger, S.L., Bluhm, C., Heidemann, K., Schaefer, I., Maraun, M., Liu, M., Scheu, S., 2018. Phylogenetic and trophic determinants of gut microbiota in soil oribatid mites. *Soil Biology and Biochemistry* 123, 155–164. doi:10.1016/j.soilbio.2018.05.011
- Gould, S.J., Lewontin, R.C., 1979. The spandrels of San Marco and the Panglossian paradigm: a critique of the adaptationist programme. *Proceedings of the Royal Society of London. Series B. Biological Sciences* 205, 581–598. doi:10.1098/rspb.1979.0086
- Hao, C., Chen, T.-W., Wu, Y., Chang, L., Wu, D., 2020. Snow microhabitats provide food resources for winter-active Collembola. *Soil Biology and Biochemistry* 143, 107731. doi:10.1016/j.soilbio.2020.107731
- Hao, C., Zhang, B., Guan, P., Xie, Z., Xu, G., Wu, D., Chen, T.-W., 2025. Microbiota as potential functional traits facilitating springtail activity in winter. *Ecology and Evolution* 15, e71448. doi:doi.org/10.1002/ece3.71448
- Heethoff, M., Domes, K., Laumann, M., Maraun, M., Norton, R.A., Scheu, S., 2007. High genetic divergences indicate ancient separation of parthenogenetic lineages of the oribatid mite *Platynothrus peltifer* (Acari, Oribatida). *Journal of Evolutionary Biology* 20, 392–402. doi:10.1111/j.1420-9101.2006.01183.x
- Janion-Scheepers, C., Phillips, L., Sgrò, C.M., Duffy, G.A., Hallas, R., Chown, S.L., 2018. Basal resistance enhances warming tolerance of alien over indigenous species across latitude. *Proceedings of the National Academy of Sciences* 115, 145–150. doi:10.1073/pnas.1715598115
- Junggebauer, A., Bluhm, C., Erdmann, G., Bluhm, S.L., Pollierer, M.M., Stefan, S., 2024.

398 Temporal variation of soil microarthropods in different forest types and regions of central
399 Europe. *Oikos* 2024, e10513. doi:doi.org/10.1111/oik.10513

400 Junker, R.R., Albrecht, J., Becker, M., Keuth, R., Farwig, N., Schleuning, M., 2022. Towards an
401 animal economics spectrum for ecosystem research. *Functional Ecology* 37, 57–72.
402 doi:10.1111/1365-2435.14051

403 Krause, A., Sandmann, D., Bluhm, S.L., Ermilov, S., Widyastuti, R., Haneda, N.F., Scheu, S.,
404 Maraun, M., 2019. Shift in trophic niches of soil microarthropods with conversion of tropical
405 rainforest into plantations as indicated by stable isotopes (15N, 13C). *PLOS ONE* 14,
406 e0224520. doi:10.1371/journal.pone.0224520

407 Luza, A.L., Barneche, D.R., Cordeiro, C.A.M.M., Dambros, C.S., Ferreira, C.E.L., Floeter, S.R.,
408 Giglio, V.J., Luiz, O.J., Mendes, T.C., Picoletto, V.A.P., Quimbayo, J.P., Silva, F.C.,
409 Waechter, L., Longo, G.O., Bender, M.G., 2023. Going across taxa in functional ecology:
410 Review and perspectives of an emerging field. *Functional Ecology* 37, 3091–3110.
411 doi:10.1111/1365-2435.14442

412 Noske, J.E., Lu, J.-Z., Schaefer, I., Maraun, M., Scheu, S., Chen, T.-W., 2024. Niche dimensions
413 in soil oribatid mite community assembly under native and introduced tree species. *Ecology*
414 and Evolution 14, e11431. doi:10.1002/ece3.11431

415 Petersen, H., Luxton, M., 1982. A comparative analysis of soil fauna populations and their role
416 in decomposition processes. *Oikos* 39, 288–388. doi:10.2307/3544689

417 Ponge, J.-F., 2020. Move or change, an eco-evolutionary dilemma: The case of Collembola.
418 *Pedobiologia* 79, 150625. doi:10.1016/j.pedobi.2020.150625

419 Potapov, A., Bellini, B., Chown, S., Deharveng, L., Janssens, F., Kováč, L., Kuznetsova, N.,
420 Ponge, J.-F., Potapov, M., Querner, P., Russell, D., Sun, X., Zhang, F., Berg, M., 2020.
421 Towards a global synthesis of Collembola knowledge: challenges and potential solutions.
422 *Soil Organisms* 92, 161–188. doi:10.25674/so92iss3pp161

423 Potapov, A.M., Beaulieu, F., Birkhofer, K., Bluhm, S.L., Degtyarev, M.I., Devetter, M., Goncharov,
424 A.A., Gongalsky, K.B., Klärner, B., Korobushkin, D.I., Liebke, D.F., Maraun, M., Mc Donnell,
425 R.J., Pollierer, M.M., Schaefer, I., Shrubovych, J., Semenyuk, I.I., Sendra, A., Tuma, J.,

Tůmová, M., Vassilieva, A.B., Chen, T.-W., Geisen, S., Schmidt, O., Tiunov, A. V, Scheu, S., 2022. Feeding habits and multifunctional classification of soil-associated consumers from protists to vertebrates. *Biological Reviews* 97, 1057–1117. doi:doi.org/10.1111/brv.12832

Potapov, A.M., Chen, T.-W., Striuchkova, A. V, Alatalo, J.M., Alexandre, D., Arbea, J., Ashton, T., Ashwood, F., Babenko, A.B., Bandyopadhyaya, I., Baretta, C.R.D.M., Baretta, D., Barnes, A.D., Bellini, B.C., Bendjaballah, M., Berg, M.P., Bernava, V., Bokhorst, S., Bokova, A.I., Bolger, T., Bouchard, M., Brito, R.A., Buchori, D., Castaño-Meneses, G., Chauvat, M., Chomel, M., Chow, Y., Chown, S.L., Classen, A.T., Cortet, J., Čuchta, P., de la Pedrosa, A.M., De Lima, E.C.A., Deharveng, L.E., Doblas Miranda, E., Drescher, J., Eisenhauer, N., Ellers, J., Ferlian, O., Ferreira, S.S.D., Ferreira, A.S., Fiera, C., Filser, J., Franken, O., Fujii, S., Koudji, E.G., Gao, M., Gendreau-Berthiaume, B., Gers, C., Greve, M., Hamra-Kroua, S., Handa, I.T., Hasegawa, M., Heiniger, C., Hishi, T., Holmstrup, M., Homet, P., Høye, T.T., Ivask, M., Jacques, B., Janion-Scheepers, C., Jochum, M., Joimel, S., Jorge, B.C.S., Juceviča, E., Kapinga, E.M., Kováč,L., Krab, E.J., Krogh, P.H., Kuu, A., Kuznetsova, N., Lam, W.N., Lin, D., Lindo, Z., Liu, A.W.P., Lu, J.-Z., Luciáñez, M.J., Marx, M.T., Mawan, A., McCary, M.A., Minor, M.A., Mitchell, G.I., Moreno, D., Nakamori, T., Negri, I., Nielsen, U.N., Ochoa-Hueso, R., Oliveira Filho, L.C.I., Palacios-Vargas, J.G., Pollierer, M.M., Ponge, J.-F., Potapov, M.B., Querner, P., Rai, B., Raschmanová, N., Rashid, M.I., Raymond-Léonard, L.J., Reis, A.S., Ross, G.M., Rousseau, L., Russell, D.J., Saifutdinov, R.A., Salmon, S., Santonja, M., Saraeva, A.K., Sayer, E.J., Scheunemann, N., Scholz, C., Seeber, J., Shaw, P., Shveenkov, Y.B., Slade, E.M., Stebaeva, S., Sterzynska, M., Sun, X., Susanti, W.I., Taskaeva, A.A., Tay, L.S., Thakur, M.P., Treasure, A.M., Tsiafouli, M., Twala, M.N., Uvarov, A. V, Venier, L.A., Widenfalk, L.A., Widyastuti, R., Winck, B., Winkler, D., Wu, D., Xie, Z., Yin, R., Zampaulo, R.A., Zeppelini, D., Zhang, B., Zoughailech, A., Ashford, O., Klauberg-Filho, O., Scheu, S., 2024. Global fine-resolution data on springtail abundance and community structure. *Scientific Data* 11, 22. doi:10.1038/s41597-023-02784-x

Potapov, A.M., Guerra, C.A., van den Hoogen, J., Babenko, A., Bellini, B.C., Berg, M.P., Chown, S.L., Deharveng, L., Kováč,L., Kuznetsova, N.A., Ponge, J.-F., Potapov, M.B., Russell, D.J.,

Alexandre, D., Alatalo, J.M., Arbea, J.I., Bandyopadhyaya, I., Bernava, V., Bokhorst, S.,
 Bolger, T., Castaño-Meneses, G., Chauvat, M., Chen, T.-W., Chomel, M., Classen, A.T.,
 Cortet, J., Čuchta, P., Manuela de la Pedrosa, A., Ferreira, S.S.D., Fiera, C., Filser, J.,
 Franken, O., Fujii, S., Koudji, E.G., Gao, M., Gendreau-Berthiaume, B., Gomez-Pamies,
 D.F., Greve, M., Tanya Handa, I., Heiniger, C., Holmstrup, M., Homet, P., Ivask, M., Janion-
 Scheepers, C., Jochum, M., Joimel, S., Claudia S. Jorge, B., Jucevica, E., Ferlian, O., Iuñes
 de Oliveira Filho, L.C., Klauberg-Filho, O., Baretta, D., Krab, E.J., Kuu, A., de Lima, E.C.A.,
 Lin, D., Lindo, Z., Liu, A., Lu, J.-Z., Luciañez, M.J., Marx, M.T., McCary, M.A., Minor, M.A.,
 Nakamori, T., Negri, I., Ochoa-Hueso, R., Palacios-Vargas, J.G., Pollierer, M.M., Querner,
 P., Raschmanová, N., Rashid, M.I., Raymond-Léonard, L.J., Rousseau, L., Saifutdinov,
 R.A., Salmon, S., Sayer, E.J., Scheunemann, N., Scholz, C., Seeber, J., Shveenkova, Y.B.,
 Stebaeva, S.K., Sterzynska, M., Sun, X., Susanti, W.I., Taskaeva, A.A., Thakur, M.P.,
 Tsiafouli, M.A., Turnbull, M.S., Twala, M.N., Uvarov, A. V., Venier, L.A., Widenfalk, L.A.,
 Winck, B.R., Winkler, D., Wu, D., Xie, Z., Yin, R., Zeppelini, D., Crowther, T.W., Eisenhauer,
 N., Scheu, S., 2023. Globally invariant metabolism but density-diversity mismatch in
 springtails. *Nature Communications* 14, 674. doi:10.1038/s41467-023-36216-6

Potapov, A.M., Pollierer, M.M., Salmon, S., Šustr, V., Chen, T.-W., 2021. Multidimensional
 trophic niche revealed by complementary approaches: Gut content, digestive enzymes, fatty
 acids and stable isotopes in Collembola. *Journal of Animal Ecology* 90, 1919–1933.
 doi:10.1111/1365-2656.13511

Revell, L.J., 2024. phytools 2.0: an updated R ecosystem for phylogenetic comparative methods
 (and other things). *PeerJ* 12, e16505. doi:10.7717/peerj.16505

Rosenberg, Y., Bar-On, Y.M., Fromm, A., Ostikar, M., Shoshany, A., Giz, O., Milo, R., 2023. The
 global biomass and number of terrestrial arthropods. *Science Advances* 9, eabq4049.
 doi:10.1126/sciadv.abq4049

Schaefer, I., Caruso, T., 2019. Oribatid mites show that soil food web complexity and close
 aboveground-belowground linkages emerged in the early Paleozoic. *Communications
 Biology* 2, 387. doi:10.1038/s42003-019-0628-7

Schaefer, I., Norton, R. A., Scheu, S., Maraun, M., 2010. Arthropod colonization of land – Linking

485 molecules and fossils in oribatid mites (Acari, Oribatida). *Molecular Phylogenetics and*
486 *Evolution*, 57, 113–121.

487 Shimano, S., 2011. Aoki's oribatid-based bioindicator systems*. *Zoosymposia* 6, 200–209.
488 doi:10.11646/zoosymposia.6.1.30

489 Silvertown, J., McConway, K., Gowing, D., Dodd, M., Fay, M.F., Joseph, J.A., Dolphin, K., 2005.
490 Absence of phylogenetic signal in the niche structure of meadow plant communities.
491 *Proceedings of the Royal Society B* 273, 39–44. doi:10.1098/rspb.2005.3288

492 Subías, L. S., 2022. Listado sistematico, sinonimico y biogeografico de los acaros oribatidos
493 (Acariformes: Oribatida) del Mundo (except fosiles). *Monografías electrónicas S.E.A.*, no.
494 12.

495 Susanti, W.I., Bartels, T., Krashevskaya, V., Widyastuti, R., Deharveng, L., Scheu, S., Potapov, A.,
496 2021. Conversion of rainforest into oil palm and rubber plantations affects the functional
497 composition of litter and soil Collembola. *Ecology and Evolution* 11, 10686–10708.
498 doi:10.1002/ece3.7881

499 van Straalen, N.M., 2023. *Soil Invertebrates: Kaleidoscope of Adaptations*. CRC Press.

500 van Straalen, N.M., 2021. Evolutionary terrestrialization scenarios for soil invertebrates.
501 *Pedobiologia* 87–88, 150753. doi:10.1016/j.pedobi.2021.150753

502 Vasconcelos, T., O'meara, B.C., Beaulieu, J.M., 2022. Retiring “cradles” and “museums” of
503 biodiversity. *American Naturalist* 199, 194–205. doi:10.1086/717412

504 Véron, S., Saito, V., Padilla-García, N., Forest, F., Bertheau, Y., 2019. The use of phylogenetic
505 diversity in conservation biology and community ecology: A common base but different
506 approaches. *The Quarterly Review of Biology* 94, 123–148. doi:10.1086/703580

507 von Saltzwedel, H., Maraun, M., Scheu, S., Schaefer, I., 2014. Evidence for frozen-niche
508 variation in a cosmopolitan parthenogenetic soil mite species (Acari, Oribatida). *PLoS ONE*
509 9, e113268. doi:10.1371/journal.pone.0113268

510 Winemiller, K.O., Fitzgerald, D.B., Bower, L.M., Pianka, E.R., 2015. Functional traits, convergent
511 evolution, and periodic tables of niches. *Ecology Letters* 18, 737–751. doi:10.1111/ele.12462

- 512 Xie, Z., Chen, T.-W., Potapov, M., Zhang, F., Wu, D., Scheu, S., Sun, X., 2022. Ecological and
513 evolutionary processes shape below-ground springtail communities along an elevational
514 gradient. *Journal of Biogeography* 49, 469–482. doi:<https://doi.org/10.1111/jbi.14317>
- 515 Yang, J., Pu, G., Jüds, M., Erktan, A., Scheu, S., Lu, J.Z., 2025. Microhabitat more than
516 ecosystem type determines the trophic position of springtail species. *Soil Biology and*
517 *Biochemistry* 209, 109912. doi:10.1016/j.soilbio.2025.109912
- 518 Yin, R., Kardol, P., Thakur, M.P., Gruss, I., Wu, G.L., Eisenhauer, N., Schädler, M., 2020. Soil
519 functional biodiversity and biological quality under threat: Intensive land use outweighs
520 climate change. *Soil Biology and Biochemistry* 147, 107847.
521 doi:10.1016/j.soilbio.2020.107847
- 522 Yu, D., Du, S., Wei, X., Zhu, J., Ding, Y., Hu, F., Liu, M., Zhang, F., 2024. Whole-genome-based
523 phylogenetic analyses provide new insights into the evolution of springtails (Hexapoda:
524 Collembola). *Molecular Phylogenetics and Evolution* 200, 108169.
525 doi:doi.org/10.1016/j.ympev.2024.108169
- 526 Zhang, B., Chen, T.-W., Mateos, E., Scheu, S., Schaefer, I., 2018. Cryptic species in
527 *Lepidocyrtus lanuginosus* (Collembola: Entomobryidae) are sorted by habitat type.
528 *Pedobiologia* 68, 12–19. doi:10.1016/j.pedobi.2018.03.001