

1 **Microbial necromass contribution to topsoil organic carbon storage of natural**
2 **and agricultural ecosystems,**

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Abstract

Microbial necromass is an important component of soil carbon (C). Yet, the relative contribution of microbial necromass in shaping the global C stocks in agricultural and natural ecosystems worldwide remains virtually unknown. Here, we compiled data on fungal and bacterial necromass along with soil organic carbon (SOC) from the 0–20 cm soil layer across 486 study sites (145 agricultural and 341 natural ecosystems) to evaluate the relative contribution of fungal necromass C (FNC) and bacterial necromass C (BNC) to SOC. Our results indicated that, on average, FNC is two times more important than BNC in explaining SOC in both agricultural and natural ecosystems. The contributions of FNC and BNC to SOC were markedly higher in agricultural ecosystems compared with natural ecosystems, with a contrasting trend in the FNC/BNC ratio. Soil physicochemical properties (soil C/N ratio and clay content) were the most important predictors of the contributions of FNC and BNC to SOC in both ecosystems, while geographical factor (elevation) was the most important predictor of the FNC/BNC ratio. Our study enhances the current level of understanding regarding microbially mediated biogeochemical cycling and SOC dynamics, underscoring the critical role of microbial necromass in the global C cycle.

Keywords: agricultural ecosystems, bacterial necromass carbon, fungal necromass carbon, microbial necromass carbon, natural ecosystems

1 Introduction

As the largest carbon (C) pool in the terrestrial biosphere, soil organic carbon (SOC) plays a pivotal role in shaping the global C cycle and climate system (Bellamy et al., 2005; Crowther et al., 2015). In brief, plant inputs provide the primary carbon source to soils, and microbial processing transforms these inputs into microbial necromass that can persist over long turnover times (Angst et al., 2021; Cotrufo et al., 2013). Although the living soil microbial biomass typically constitutes only about 2% of SOC (a ratio referred to as the microbial quotient; Anderson & Domsch, 1989; Liu et al., 2023), microbial necromass has been shown to contribute more than half and up to approximately 80% of SOC, depending on soil type and analytical methods (Liang & Balser, 2011; Kallenbach et al., 2016; Liang et al., 2019). In other words, microbial necromass C (MNC) constitutes a substantial and critical component of stable SOC (Ma et al., 2018), and its dynamics are increasingly recognized for their role in regulating the terrestrial carbon cycle and climate feedbacks (Zhao et al., 2023). As such, there is growing scientific attention on the forces driving the accumulation of MNC and its contribution to SOC (Liang et al., 2017; Ni et al., 2020; Luo et al., 2022; Zhou et al., 2023). To gain a comprehensive understanding of MNC in the global C cycle, recent research has highlighted the distinct roles of fungal and bacterial necromass, revealing their contrasting responses to environmental and anthropogenic drivers. For instance, studies have shown that the accumulation and contribution of MNC are sensitive to factors such as aridity, primary productivity, agricultural management practices like tillage and fertilization, as well as key soil properties including pH and clay content (Zhang et al., 2021; Zhou et al., 2023; Xu et al., 2024).

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删除了: The traditional paradigm is that SOC storage is primarily formed directly based on plant material (Zhu and Miller, 2003; von Lützow et al., 2008). However, emerging evidence demonstrates that the stable SOC pool is predominantly composed of microbially derived organic compounds rather than plant-derived residues, indicating that microbial necromass may in fact be a primary source of stable SOC (Kallenbach et al., 2016; Liang et al., 2017). Microbes generate biomass by degrading plant-derived C, which is subsequently stabilized as microbial necromass through interactions with minerals and various processes within soil aggregates (Liang et al., 2017)...

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删除了: Although the living soil microbial biomass constitutes only about 2% of SOC, microbial necromass carbon (MNC) can contribute up to 50%–80% of SOC (Liang and Balser, 2011; Kallenbach et al., 2016; Liang et al.,

106 Despite these advances, it remains unclear whether these organism-specific
107 mechanisms translate into systematic differences in necromass contributions between
108 ecosystems under varying degrees of human interference, such as agricultural versus
109 natural systems.

110 With the distinct roles of fungi and bacteria in decomposing organic matter and
111 stabilizing organic carbon in soil, the relative contribution to SOC of fungal and
112 bacterial necromass C could be used to track the dynamics of SOC storage (Malik et
113 al., 2016). The cell walls of fungi primarily consist of chitin (a nitrogen-containing
114 polysaccharide) and β -glucans, whereas bacterial cell walls are mainly composed of
115 peptidoglycan—a complex of sugars and amino acids (Lenardon et al., 2007). As
116 bacterial amino sugars is degradable rather than fungal chitin or β -glucans, fungal
117 necromass existed in soil generally with longer turnover time than bacterial necromass,
118 (Xu et al., 2022). Wang et al. (2021a) reported that the contribution of fungal
119 necromass carbon (FNC) to SOC exceeded 65%, considerably higher than that of
120 bacterial necromass carbon (BNC, 32–36%). This pattern is likely attributed to the
121 slower decomposition rate and stronger mineral-associative capacity of fungal
122 necromass. Furthermore, greater fungal biomass and higher turnover rates may
123 enhance the input flux of fungal necromass (Klink et al., 2022). The contributions of
124 FNC and BNC to SOC depended on the type of ecosystems (Wang et al., 2021a; Cao
125 et al., 2023; Xu et al., 2024). However, few studies on fungal and bacterial necromass
126 carbon and their contribution to SOC has been reported for ecosystems under human
127 interference (Zhou et al., 2023).

128 Terrestrial ecosystems can be broadly categorized into managed (agricultural) and
129 natural ecosystems (Hobbs et al., 2011; Keith et al., 2022). The agricultural
130 ecosystems are typical of plant litter derived from single crops under intensive human
131 management (Bohan et al., 2013), a context that typically leads to bacterial-dominated
132 soil communities (van Der Heijden et al., 2008). In contrast, natural ecosystems
133 display greater diversity in plant litter and root deposits (Wu et al., 2019). Within
134 these natural systems, minimal human interference enhances fungal mycelial
135 networks and stable soil aggregates, leading to higher FNC sequestration and
136 contributions to SOC (Sanaullah et al., 2020; Sae-Tun et al., 2022). While bacteria are
137 undoubtedly vital decomposers, fungi play a distinct and often dominant role in the
138 initial breakdown of complex plant polymers such as cellulose and lignin. This
139 functional prominence stems from their potent enzymatic machinery and hyphal
140 growth form, which enable physical penetration and decay of solid organic matter (de
141 Boer et al., 2005). As key decomposers, fungi are thus critical in processing cellulose
142 and other complex organic compounds (Hättenschwiler et al., 2005). Accordingly, as
143 demonstrated by Choi et al. (2018), soil cellulose-degrading genes are frequently
144 linked to fungal activity and abundance. Rather than implying higher cellulose
145 concentration per se, diverse plant inputs increase the chemical heterogeneity of
146 plant-derived polymers (e.g., cellulose, hemicelluloses, and lignin), which broadens
147 decomposer niches and often favors fungal communities in litter horizons
148 (Hättenschwiler et al., 2005; Štursová et al., 2012). In contrast, agricultural
149 monocultures tend to reduce fungal diversity unless mitigated by management

删除了: In order to gain a comprehensive and in-depth understanding of the role of MNC in the global C cycle, many studies have focused on the relative contributions of fungal (FNC) versus bacterial necromass (BNC; Zhang et al., 2021; Zhou et al., 2023; Xu et al., 2024)....

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删除了: To be more specific, the chemical composition and functional characteristics of fungi and bacteria differ considerably, which may also contribute to different mechanisms of SOC formation. ...

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215 practices (Chen et al., 2020). Reflecting this context dependence, cellulose-rich inputs
216 can enrich saprotrophic fungi in arable soils (Clocchiatti et al., 2021), whereas
217 bacteria may contribute substantially in mineral soils or under specific microhabitat
218 and land management conditions (Štursová et al., 2012; Choi et al., 2018). Due to
219 distinct chemical properties and organo-mineral stabilization pathways, fungal and
220 bacterial necromass exhibit differing turnover times, making the FNC/BNC ratio a
221 mechanistic tracer of SOC formation (Angst et al., 2021; Kleber et al., 2021).
222 Therefore, elucidating the global distribution and drivers of FNC, BNC, and their ratio
223 across agricultural and natural ecosystems is essential for predicting
224 management-induced shifts in SOC under varying climatic and soil conditions (Zhang
225 et al., 2021; Zhou et al., 2023; Xu et al., 2024).

226 In order to explore the global patterns and drivers of FNC, BNC and the
227 FNC/BNC ratio in agricultural and natural ecosystems, we compiled data from 486
228 study sites worldwide. The aims of this study were: (1) to quantify the contributions
229 of FNC and BNC to SOC and the FNC/BNC ratio in agricultural and natural
230 ecosystems; and (2) to investigate the primary driving factors influencing the
231 contributions of FNC and BNC to SOC and the FNC/BNC ratio.

233 2 Materials and methods

234 2.1 Data collection

235 We compiled a comprehensive dataset following the stepwise workflow. (1) We
236 collected peer-reviewed papers published from 1996 to 31 December 2022 from Web
237 of Science (<http://apps.webofknowledge.com>), Google Scholar
238 (<http://scholar.google.com>), and the China National Knowledge Infrastructure
239 (<http://cnki.net>), using the keywords: ‘amino sugars’, ‘microbial necromass’,
240 ‘microbial residue’, ‘fungal residue’, and ‘bacterial residue’. Records from different
241 databases were merged and deduplicated to form an initial compilation. (2) We then
242 filtered the compiled studies to include only those focusing on topsoil, defined as the
243 0–20 cm layer. Studies reporting deeper or unspecified sampling depths (e.g., 0–30
244 cm) were excluded to ensure spatial comparability. (3) Full texts were assessed to
245 confirm the presence of paired fungal and bacterial residue data from the same
246 sample—specifically, glucosamine (GluN) and muramic acid (MurA), or directly
247 reported FNC and BNC values—to enable consistent cross-study calculation of the
248 FNC/BNC ratio. Studies lacking either biomarker were excluded from ratio analyses,
249 though those directly reporting the FNC/BNC ratio were retained. (4) Eligible
250 observations were classified into agricultural ecosystems (including dry land, irrigated
251 cropland, and submerged paddy) or natural ecosystems (forest and grassland) based
252 on study metadata. (5) For natural ecosystems, data from fertilized, polluted,
253 experimentally treated, or otherwise anthropogenically disturbed sites were excluded.

254 In total, the final dataset consisted of 2094 observations from 486 sites worldwide
255 (145 agricultural and 341 natural sites; Figure 1) reported in 164 peer-reviewed papers.
256 Of these observations, 1001 were from agricultural ecosystems, and 1093 from natural
257 ecosystems. Among the 341 natural sites, 195 were forests and 146 were grasslands.
258 For agricultural sites, we used Google Earth Engine with the LGRIP30 V1 dataset to

删除了: This suggests cellulose-rich soils may support higher fungal diversity and abundance. The low cellulose and chitin abundance in agricultural ecosystems might result from plant monoculture, while diverse plant inputs in natural ecosystems lead to rich soil cellulose content and higher fungal abundance and FNC (Song et al., 2022). Due to di

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删除了: e compiled a comprehensive dataset from a range of peer-reviewed papers published before 2022 through the Web of Science (<http://apps.webofknowledge.com>), Google Scholar (<http://scholar.google.com>), and the China National Knowledge Infrastructure (<http://cnki.net>), using the keywords ‘amino sugars’, ‘microbial necromass’, ‘microbial residue’, ‘fungal residue’, and ‘bacterial residue’. Data compilation was conducted following four stringent criteria to reduce bias from selected publications: (1) fungal and bacterial necromass (or glucosamine [GluN] and muramic acid [MurA]) had to be reported simultaneously for the same samples; (2) only data from agricultural and natural ecosystems (i.e., grasslands and forests) were used; (3) for natural ecosystems, we excluded the data from fertilized, polluted, treated, or anthropogenically disturbed sites; and (4) we only included data from the top 0–20 cm of the soil profile reported in the publications, with other depths or unspecified soil depths excluded from our dataset...

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298 classify agricultural ecosystem into dry land and irrigated cropland, and we overlaid
299 the JRC surface-water seasonality layer to extract submerged paddy from the irrigated
300 class (LGRIP30 irrigated value = 2 and JRC seasonality > 1). We ultimately classified
301 145 samples of agricultural ecosystems into 32 dry land, 72 irrigated, and 41
302 submerged paddy sites.

303 We calculated the FNC and BNC based on amino sugar concentrations following
304 widely used conversion factors, correcting total GluN for its bacterial share using
305 MurA;

$$306 \quad FNC = \left(\frac{GluN}{179.17} - 2 \times \frac{MurA}{251.23} \right) \times 179.17 \times 9 \quad \text{Equation 1}$$

307 where 9 (unitless) is the conversion factor from GluN to FNC. To estimate
308 fungus-derived GluN, we subtracted the bacterial share of GluN assuming an
309 empirical GluN:MurA molar ratio of 2:1 for bacterial residues. 179.17 and 251.23 are
310 the molecular weights of GluN and MurA, respectively. And their units are all g/mol.
311 The unit of FNC is mg/kg.

$$312 \quad \frac{BNC}{45} = MurA \times 45 \quad \text{Equation 2}$$

313 where 45 (unitless) is the conversion factor from MurA to BNC. The unit of BNC
314 is mg/kg.

315 Additional information including site geographic location (latitude and longitude),
316 topographical condition (elevation), climatic factors (MAT) and mean annual
317 precipitation [MAP]), soil physicochemical properties (pH, SOC, TN, clay content,
318 and soil temperature), and biotic (microbial and plant) factors were recorded.
319 Specifically, biotic factors included microbial biomass carbon (MBC), microbial
320 biomass nitrogen (MBN), MBC/MBN, net primary production (NPP), and
321 belowground biomass C density (BGBC). The data of topographical condition
322 (elevation) was classified as geographical factor in this study. When MAT and MAP
323 were unavailable in the original articles, we extracted them from the global climate
324 layers of WorldClim (<http://www.worldclim.org/>) with a grid precision of 30 × 30 arc
325 sec according to geographic location. Missing elevation data were extracted using the
326 *elevatr* package v.0.4.2 (Hollister, 2021) in the R environment. We acquired the data
327 on annual mean soil temperature from the study of Lembrechts et al. (2022), while
328 other absent soil physicochemical data were extracted from the Harmonized World
329 Soil Database

330 (<https://www.fao.org/soils-portal/data-hub/soil-maps-and-databases/harmonized-world-soil-database-v12/en/>)
331 and SoilGrids 2.0 (Poggio et al., 2021; <https://www.soilgrids.org/>) using ArcGIS 10.3. We supplemented missing climatic and
332 soil variables using high-resolution, globally interpolated datasets to ensure consistent
333 spatial coverage across all sites. After retrieving missing value from gridded data, we
334 typically calibrate them against field-reported values via a field-anchored bias
335 correction (i.e., a site- or region-specific “delta” adjustment) to minimize errors
336 introduced by gridded data. While the use of such data introduces inherent
337 uncertainties, these databases are widely adopted in global-scale ecological analyses
338 and provide the most feasible approach for a unified assessment. In addition, the data
339 on NPP and BGBC were acquired from the studies of Zhao and Running (2010) and
340

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352 Spawn et al. (2020), respectively. Missing MBC and MBN data were acquired using a
353 global database with a high resolution of 30×30 arc sec (Wang et al., 2022).

354 355 2.2 Statistical analysis

356 All the statistical analyses were performed using R v4.1.3 (R Core Team, 2021).
357 Initially, the Shapiro–Wilk test was employed to assess the normality of our data,
358 followed by the application of Levene's test to evaluate the homogeneity of variances
359 across different groups. To detect the significant differences in the contributions of
360 FNC and BNC to SOC, and the FNC/BNC ratio between agricultural and natural
361 ecosystems, as well as between forest and grassland ecosystems, the Wilcoxon rank
362 sum test was conducted. [Kruskal–Wallis and Dunn's post hoc tests were used to](#)
363 [assess the significant differences of the contributions of FNC and BNC to SOC, and](#)
364 [the FNC/BNC ratio among dry land, irrigated cropland, and submerged paddy.](#) We
365 used Spearman's rank correlation coefficient to explore the connections between the
366 16 variables considered in this study, including geographical and climatic factors, soil
367 physicochemical properties, and biotic factors. Since there was a strong positive
368 correlation between MAT and soil temperature (Figure S1), soil temperature was
369 excluded from our subsequent analyses. Linear regressions between different factors
370 and the contributions of FNC and BNC to SOC and the FNC/BNC ratio were
371 performed. Dots and smoothing curves were drawn using the *geom_point* and
372 *geom_smooth* functions, respectively, in the *ggplot2* package v.3.4.0 (Wickham,
373 2016).

374 Variation partitioning analysis was conducted using the *vegan* package v.2.5.7
375 (Oksanen et al., 2020) to evaluate the effects of four types of factors on the
376 contributions of FNC and BNC to SOC and the FNC/BNC ratio in agricultural and
377 natural ecosystems at global scale. We used a variance inflation factor threshold of 3.3
378 to eliminate those variables that were strongly correlated and avoid multicollinearity
379 (Figure S2; Kock, 2015; Fanin et al., 2020). Following factor selection, boosted
380 regression trees (BRTs) were used to partition independent influences of geographical
381 (elevation) and climatic (MAT and MAP) factors, soil physicochemical properties (pH,
382 clay, C/N, and SOC), and biotic factors (NPP, BGBC, MBC, and MBC/MBN) on the
383 contributions of FNC and BNC to SOC and the FNC/BNC ratio with the *gbm* package
384 v.2.1.8.1 (Greenwell et al., 2022).

385 Utilizing the selected factors, we performed structural equation models (SEMs)
386 to quantify the effects (direct, indirect and both) of four types of factors on the
387 contributions of FNC and BNC to SOC and the FNC/BNC ratio using *lavaan* package
388 v.0.6.19 (Rosseel, 2012). According to the previously reported potential causal
389 relationships between explanatory and response variables (Wang et al., 2021a, 2021b;
390 Li et al., 2024), we established the *priori* structural equation models for agricultural
391 and natural ecosystems, respectively (Figure S3). The SEMs were fitted via maximum
392 likelihood estimation, with non-significant paths iteratively pruned through stepwise
393 exclusion, followed by model evaluation using modification indices and
394 goodness-of-fit criteria. The fit indices included degrees of freedom (df), chi-square
395 (χ^2 , $0 \leq \chi^2/df \leq 2$), comparative fit index (CFI > 0.9), and root mean square error of

396 approximation (RMSEA < 0.08), which were used to assess the adequacy of the SEM.
397 Map, box, bar, bubble, and lollipop charts were plotted with the *ggplot2* package
398 v.3.4.0 (Wickham, 2016). To enhance map visualization, the *ggnewscale* package
399 v.0.4.8 (Campitelli, 2022) was necessary alongside the *ggplot2* package v.3.4.0
400 (Wickham, 2016). Similarly, the *ggpubr* package v.0.5.0 (Kassambara, 2022) was an
401 additional necessity when creating lollipop charts.

402 3 Results

403 3.1 Fungal and bacterial necromass contribution to SOC in agricultural and natural 404 ecosystems

405 Our analysis revealed statistically significant disparities in the contributions of FNC
406 and BNC to SOC in agricultural and natural ecosystems at the global scale ($P < 0.05$;
407 Figure 2a, b). Notably, the average contributions of FNC and BNC to SOC were
408 substantially higher in agricultural ecosystems than in natural ecosystems ($P < 0.001$;
409 Figures 2a, b). For FNC, the average contribution was 34.39% in agricultural
410 ecosystems, versus 29.24% in natural ecosystems. BNC contributed an average of
411 15.65% to SOC in agricultural ecosystems, compared to 14.02% in natural
412 ecosystems (Table 1). Our results also indicated that the contributions of FNC to SOC
413 were approximately twice those of BNC in agricultural and natural ecosystems (Table
414 1).

415 The contributions of FNC and BNC to SOC indicated no significant difference
416 between dry land and irrigated cropland ($P > 0.05$), whereas both differed
417 significantly from submerged paddy ($P < 0.05$; Figure S4a, b). In detail, across dry
418 land, irrigated cropland, and submerged paddy, the mean contributions of FNC to
419 SOC were 37.77%, 35.35%, and 22.82%, respectively, whereas those of BNC were
420 17.34%, 15.95%, and 10.55% (Table 1). Moreover, there were no significant
421 differences in the contributions of FNC and BNC to SOC between forest and
422 grassland ecosystems ($P > 0.05$; Figure S5). Specifically, FNC contributed, on
423 average, 29.11% to SOC in forests and 26.75% in grasslands, while BNC contributed
424 13.48% in forests and 14.34% in grasslands (Table 1).

425 3.2 Ratios of fungal and bacterial necromass in agricultural and natural ecosystems,

426 Our results indicated that, at the global scale, the soil FNC/BNC ratio differs
427 significantly between agricultural and natural ecosystems ($P < 0.05$; Fig. 2c), with a
428 higher ratio in natural ecosystems (3.22) than in agricultural ecosystems (2.61; Table
429 1). The FNC/BNC ratio did not differ significantly among dry land, irrigated cropland,
430 and submerged paddy ($P > 0.05$; Figure S4c), with average FNC/BNC ratios of 2.87,
431 2.51, and 2.62, respectively (Table 1). Similarly, there was no significant difference in
432 the FNC/BNC ratio between forest and grassland ecosystems ($P > 0.05$; Figure S5),
433 and the average FNC/BNC ratios for forests and grasslands were 2.80 and 3.58 (Table
434 1), respectively.

435 3.3 Associations of abiotic and biotic factors with microbial necromass parameters

436 Soil physicochemical factors were the most important influence on the contributions

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These findings were further substantiated by the results obtained from the BRTs. These results indicated that soil physicochemical factors accounted for a substantial portion of the variance in the contributions of FNC and BNC to SOC in agricultural and natural ecosystems (Figures 4a–d), and geographical factors played a similar role in explaining the FNC/BNC ratio (Figures 4e, f). More precisely, soil physicochemical factors were identified as the primary

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583 of FNC and BNC to SOC across both ecosystem types (Figures 3a–d, 4a–d).
584 Specifically, they explained 16% and 17% of the variance in the contributions of FNC
585 and BNC to SOC in agricultural ecosystems, respectively (Figures 3a, c), and 20%
586 and 24% in natural ecosystems (Figures 3b, d). BRTs corroborated this pattern, with
587 soil physicochemical factors showing the highest relative influence (51% for FNC,
588 and 44% for BNC) in agricultural systems and 44% in natural systems (Figures 4a–d).
589 All BRT models were significant ($P < 0.001$), with explained variance 36–66%. While
590 soil factors dominated overall, responses to individual variables differed between
591 ecosystems. In detail, in agricultural systems, the C/N ratio ranked third for FNC after
592 clay and SOC (Figure 4a), whereas C/N was the top predictor for FNC in natural
593 systems and for BNC in both ecosystems (Figures 4b–d). Consistently, linear models
594 showed declines in the contributions of FNC and BNC with increasing C/N in both
595 ecosystems (Figures S6g, S7g). SEMs yielded convergent results, indicating both
596 direct and indirect pathways (Figures 5a–d, 6a–d). Notably, the direct and total effects
597 of soil physicochemical factors on FNC were negative in agricultural but positive in
598 natural ecosystems (Figures 5a, b, 6a, b), whereas the effects on BNC were negative
599 in both ecosystem types (Figures 5c, d, 6c, d).

600 Our results indicated that geographical factors were the most important
601 contributors to explain the FNC/BNC ratio in both agricultural and natural ecosystems,
602 accounting for 21% and 10% of the explained variance in the FNC/BNC ratio,
603 respectively (Figures 3e, f). The results of the BRTs suggested that geographical
604 factors played a similar role in explaining the FNC/BNC ratio (Figures 4e, f). In the
605 BRT models, geographical factors emerged as the primary influencers of the
606 FNC/BNC ratio in agricultural and natural ecosystems, accounting for 32% and 44%
607 of the variance in each case, respectively (Figures 4e, f). To be more specific,
608 elevation was the most significant geographical factors influencing the FNC/BNC
609 ratio in both ecosystems (Figures 4e, f). Moreover, the FNC/BNC ratio in agricultural
610 and natural ecosystems show significantly increased with an increase elevation
611 (Figure S8a). The results of SEMs also indicated that geographical factors were the
612 most influential factors for the FNC/BNC ratio in agricultural and natural ecosystems,
613 exerting both direct and indirect effects on this ratio (Figures 5e, 6e), with the
614 standardized total effect being positive (Figures 5f, 6f).

616 4 Discussion

617 4.1 Fungal necromass contributes two times more to SOC than bacterial necromass.
618 Our results show that in agricultural ecosystems, FNC/SOC ranged from 0.09% to
619 97.53% (mean \pm SE: $34.39 \pm 0.67\%$), and BNC/SOC ranged from 0.81% to 65%
620 ($15.65 \pm 0.33\%$). In natural ecosystems, FNC/SOC ranged from 0.92% to 96.29%
621 ($29.24 \pm 0.51\%$), and BNC/SOC ranged from 0.25% to 89.45% ($14.02 \pm 0.36\%$)
622 (Table 1). The FNC/BNC ratio ranged from 0.02 to 12.74 (2.61 ± 0.06) in agricultural
623 ecosystems and from 0.12 to 44.24 (3.22 ± 0.11) in natural ecosystems (Table 1).
624 Despite substantial variability at the individual sample level, the mean contribution of
625 FNC was approximately twice that of BNC in both ecosystem types. Moreover, the
626 mean FNC/BNC ratio was significantly higher in natural ecosystems than in

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MNC is an important component of SOC (Ma et al., 2018), and its variations can influence the feedback effects on the C cycle and global climate change (Zhao et al., 2023). Our study indicated that the contributions of FNC to SOC were approximately twice those of BNC in agricultural and natural ecosystems. Although the contributions of FNC and BNC to SOC were significantly higher in agricultural ecosystems than in natural ecosystems, the FNC/BNC ratio was significantly higher in the latter. In addition, soil physicochemical properties and geographical factors were the most important predictors of the contributions of MNC (FNC and BNC) to SOC and the FNC/BNC ratio in the two ecosystems, respectively. These findings enhance our understanding of microbially mediated biogeochemical cycling processes under current and future climate scenarios.

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删除了: Variation in the contributions of FNC and BNC to SOC and the FNC/BNC ratio in agricultural and natural ecosystems... With growing appreciation for the critical role of microbial necromass constituents in forming SOC, it is critical to understand the different drivers of this process across the globe. Our results show that in agricultural ecosystems,

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655 agricultural ecosystems ($P < 0.05$; Figure 2). Although this general pattern has been
656 reported in previous studies (Liang et al., 2019; Wang et al., 2021a; Zhang et al., 2023;
657 Ding et al., 2024), the systematic differences in the magnitude of these contributions
658 between agricultural and natural ecosystems—and their underlying drivers—have
659 remained poorly understood. Our study not only confirms the broad pattern but also
660 elucidates these ecosystem-level disparities and their environmental determinants.
661 Consistent with our finding that the contribution of fungal necromass carbon (FNC) to
662 SOC exceeded that of bacterial necromass carbon (BNC) in both ecosystem types
663 (Table 1), the predominance of fungal necromass may be attributed to its more
664 recalcitrant cell wall composition (e.g., chitin) and slower decomposition rate (Wang
665 et al., 2021a). Our BRT and SEM analyses further identified soil clay content and C/N
666 ratio as key drivers of FNC accumulation (Figs. 4a, 5a), reinforcing the importance of
667 organo-mineral associations in the stabilization of fungal-derived carbon.
668 Additionally, our study reveals previously unreported disparities between
669 ecosystem types, for example, the contributions of both fungal and bacterial
670 necromass carbon (FNC and BNC) to SOC were significantly higher in agricultural
671 ecosystems, while the FNC/BNC ratio was substantially elevated in natural
672 ecosystems. The higher contributions of FNC and BNC to SOC in agricultural
673 ecosystems may be attributable to two main factors. First, natural ecosystems
674 typically receive larger and more heterogeneous plant-derived carbon inputs than
675 agricultural systems. These inputs expand the plant-derived SOC pool and can dilute
676 the relative contribution of microbial necromass to SOC, thereby resulting in a lower
677 perceived contribution of microbial necromass in natural ecosystems (Angst et al.,
678 2021; Kleber et al., 2021). Second, the significantly lower soil C/N ratio in
679 agricultural ecosystems (10.78) compared to natural ecosystems (27.44) reflects
680 relative nitrogen enrichment, largely resulting from anthropogenic fertilization
681 (Castellano et al., 2015; Chen et al., 2020). This nitrogen-rich environment can
682 enhance microbial carbon use efficiency and alleviate nutrient limitation, thereby
683 promoting the production and accumulation of microbial necromass (Liang et al.,
684 2017). Supporting this mechanism, we found that the contributions of both FNC and
685 BNC to SOC decreased significantly with increasing soil C/N ratio in both
686 agricultural ecosystems (FNC/SOC: $R = -0.27$, $P < 0.001$; BNC/SOC: $R = -0.29$, $P <$
687 0.001) and natural ecosystems (FNC/SOC: $R = -0.17$, $P < 0.001$; BNC/SOC: $R =$
688 -0.35 , $P < 0.001$; Figures S6g, S7g). These results further underscore that a lower soil
689 C/N ratio—often indicative of higher nitrogen availability—is a key driver of
690 microbial necromass accumulation. It should be noted that although in situ plant
691 residues in agricultural systems (e.g., cereal straw) may have high C/N ratios, the
692 overall soil C/N ratio is reduced by management practices such as mineral fertilization
693 and the incorporation of low C/N organic amendments.
694 Furthermore, nutrient-rich conditions prevalent in agricultural systems (e.g., due
695 to fertilization) often select for bacterial-dominated communities, as many bacteria
696 exhibit r -strategist traits that support rapid growth under high resource availability. In
697 contrast, natural ecosystems—characterized by lower nutrient availability and greater
698 resource heterogeneity—tend to favor fungal dominance, since fungi often function as

删除了: In this study, we found that the contribution of FNC to SOC significantly exceeded that of BNC, with the former contributing approximately twice as much as BNC to SOC in both agricultural and natural ecosystems (including forests and grasslands; Table 1). ...

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删除了: These findings were consistent with previous studies (Liang et al., 2019; Wang et al., 2021a; Zhang et al., 2023; Ding et al., 2024). On the one hand, this might be attributable to the various nutrient utilization strategies of the different microbial groups. Labile compounds in bacterial cell walls are preferentially utilized by microbes, resulting in a lower content of bacterial residue in soil (Hu et al., 2020). In contrast, fungi, with their more recalcitrant compounds (e.g., chitin), decompose slowly, allowing them to persist longer in soil. Over time, fungal residue contributes more significantly to SOC pools (Wang et al., 2021a). On the other hand, the thicker cell walls of fungi result in a lower surface area-to-volume ratio of their residual fragments, facilitating the easy formation of complex macromolecular aggregates (Zhao et al., 2023). This contributes significantly to the stability of fungal necromass and plays a crucial role in the accumulation of SOC. Additionally, the comparatively larger biomass of fungi may be a contributing factor to the predominance of FNC over BNC in terms of contributing to SOC in agricultural and natural ecosystems (He et al., 2020). In the present study, their contributions were found to be consistently higher in agricultural ecosystems as compared to natural ecosystems

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删除了: First, owing to the diverse range of plant communities and minimal human interference in natural ecosystems

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776 K-strategists with higher efficiency in decomposing complex organic matter under
777 resource-limited conditions (Strickland & Rousk, 2010; Yu et al., 2022). This shift in
778 microbial community composition is observed in our results, which show a
779 significantly higher FNC/BNC ratio in natural ecosystems across our global dataset
780 (Figure 2c, Table 1). A high FNC/BNC ratio signifies a fungal-dominated
781 decomposition pathway. Fungal necromass—rich in recalcitrant compounds such as
782 chitin—is more resistant to decay, and fungal hyphae play a key role in the formation
783 of stable soil aggregates that physically protect organic matter from degradation
784 (Lenardon et al., 2007). This pathway promotes the formation of stable, long-turnover
785 SOC pools essential for long-term carbon sequestration (Six et al., 2006; Lehmann et
786 al., 2020). Furthermore, fungi generally exhibit higher carbon use efficiency than
787 bacteria, meaning a larger proportion of assimilated carbon is allocated to biomass
788 production (and subsequently necromass) rather than being respired as CO₂ (Wang &
789 Kuzyakov, 2024). Thus, the fungal-driven pathway characteristic of natural
790 ecosystems represents a highly efficient conversion of plant litter into persistent soil
791 organic matter (Kallenbach et al., 2016; Malik et al., 2016). Conversely, the lower
792 FNC/BNC ratio observed in agricultural ecosystems reflects a bacterial-dominated
793 pathway, accelerated by practices such as tillage and nutrient amendments. This
794 pathway is associated with faster carbon cycling and greater carbon loss through
795 respiration. Although microbial necromass can accumulate under these
796 conditions—sometimes contributing more significantly to a reduced total SOC
797 pool—the resulting carbon is often less stabilized (Zhou et al., 2023). Therefore, the
798 FNC/BNC ratio serves not merely as a descriptive metric, but as a functional
799 biomarker that elucidates fundamental differences in the stability and persistence of
800 SOM between managed agricultural systems and natural ecosystems.

801 Notably, as major components of agricultural ecosystems, both dryland and
802 irrigated croplands exhibited significantly greater contributions of FNC and BNC to
803 SOC than submerged paddy soils, although the FNC/BNC ratio did not differ
804 significantly among these three systems (Figure S4). This pattern may reflect similar
805 aeration regimes in dryland and irrigated systems (predominantly oxygenated),
806 leading to comparable decomposition–transformation–mineral association pathways
807 and, thus, similar net contributions of fungal and bacterial residues to SOC
808 (Ghezzehei et al., 2019). By contrast, persistent or periodic flooding in paddy soils
809 induces anoxia, suppresses aerobic decomposition, and shifts metabolic pathways
810 (e.g., denitrification and methanogenesis), potentially suppressing fungal activity or
811 dominance and altering the relative accumulation and turnover of fungal and bacterial
812 necromass (Qiu et al., 2017), resulting in contributions that differ significantly
813 from—and are lower than—those in the other two systems. Flooding can suppress
814 fungi yet also enhance the joint retention of both fungal and bacterial necromass via
815 slower decomposition and mineral protection, yielding unchanged ratios but altered
816 totals or compositional pathways (Chen et al., 2021; Gao et al., 2024).

817 4.2 Driving factors of the change in fungal and bacterial necromass contribution to 818 SOC and their ratio.

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删除了: Important driving factors for the contributions of FNC and BNC to SOC and the FNC/BNC ratio...Our results indicated that soil physicochemical factors were key in determining the contributions of FNC and BNC to SOC (Figures 3a–d, 4a–d, 5a–d). ...

827 Deng and Liang (2022) suggested that the potential contribution of microbial
828 necromass to the SOC pool was governed by the C/N ratio. This finding was
829 confirmed by our results (Figures 4b–d). As elaborated in Section 4.1, high N
830 availability (i.e., low soil C/N ratio) promotes the production and accumulation of
831 microbial necromass (Wu et al., 2025). Consequently, the contributions of both FNC
832 and BNC to SOC decreased with increasing soil C/N ratio (Figures S6g, S7g). In
833 agricultural ecosystems, high soil N levels primarily result from fertilization (Chen
834 et al., 2020). In contrast, natural ecosystems experience minimal anthropogenic
835 disturbance, N often acts as the key limiting factor for microbial activity (Elser et al.,
836 2007). Under N-limited conditions, microbes (both fungi and bacteria) allocate more
837 energy and C resources to the synthesis of N-acquiring enzymes (e.g., proteases and
838 chitinases). This shift in metabolic strategy reduces the C allocated to biomass
839 synthesis, thereby diminishing the amount of C ultimately converted into microbial
840 necromass (Mooshammer et al., 2014; Liu et al., 2024). Thus, although microbial
841 community composition differs between natural and agricultural ecosystems, the
842 regulatory role of soil C/N ratio in shaping their structure and function remains
843 consistent (Han et al., 2024). In our study, soil clay content was identified as the
844 predominant factor governing the contribution of FNC to SOC in agricultural
845 ecosystems (Figure 4a), with this contribution increasing concomitantly with clay
846 content (Figure S6d). This suggests that soils with higher clay and silt contents
847 generally accumulate greater amounts of microbial residues, particularly those derived
848 from fungi, which can be attributed to the promotion of stable organo-mineral
849 complex formation by abundant fine soil particles (Six et al., 2006 and Liang et al.,
850 2017). Furthermore, although agricultural management practices often disturb soil
851 structure, they simultaneously enhance clay enrichment and aggregate formation,
852 thereby providing effective physical protection for the long-term stabilization of
853 fungal-derived C (Chen et al., 2020; Mou et al., 2021; Zhou et al., 2023).

854 On the contrary, geographical factor (elevation) was identified as the most
855 influential predictor of the FNC/BNC ratio in both agricultural and natural ecosystems
856 (Figures 4e, f, 5f, 6f), with the ratio increasing significantly with elevation (Figure
857 S8a). Increasing elevation typically leads to decreased temperature, and increased
858 precipitation (Körner, 2007), conditions favoring fungi over bacteria due to higher
859 enzymatic capabilities and resource-use efficiency of fungi under the environments
860 (Chen et al., 2020; Yu et al., 2022; Zhang et al., 2025). High elevation also results in
861 slower soil development, which can reduce the availability of soil nutrients
862 (Guerrero-Ramírez et al., 2020). This in turn increases environmental stress and
863 restricts bacterial activity, thereby favoring the accumulation and conversion of fungal
864 residues into necromass (Li et al., 2024). Our study further demonstrated that although
865 elevation had a direct effect on the FNC/BNC ratio, it also indirectly influenced the
866 ratio by modulating climatic factors, soil physicochemical properties, and biological
867 factors (Figures 5e, 6e). This may explain why elevation is always integrate other
868 environmental factor effects in the studies of MNC (Cui et al., 2023; Zhang et al.,
869 2025).

870

删除了: Our results indicated that soil physicochemical key in determining the contributions of FNC and BNC to SOC (Figures 3a–d, 4a–d, 5a–d). ...

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删除了: Several factors may account for the results. On the one hand, the soil C/N ratio regulates microbial community composition, thereby influencing the accumulation of FNC and BNC (Han et al., 2023). Specifically, a high C/N ratio often indicates a relative surplus of C and a limitation of N, which constrains microbial metabolism, reduces microbial turnover and biomass input, and consequently decreases the accumulation of FNC and BNC (Wu et al., 2025). On the other hand, under high C/N conditions, microbes tend to allocate more C to maintenance respiration or CO₂ release rather than biomass synthesis, leading to a lower efficiency of necromass production and diminished contribution to SOC (Liu et al., 2024). In addition, a high C/N ratio may also impair the binding capacity of microbial residues to soil minerals, thereby reducing their long-term stabilization and further weakening their contribution to SOC (Li et al., 2024). In contrast, soil clay content was the most predominant factor influencing the contribution of FNC to SOC in agricultural ecosystems (Figure 4a), and this contribution increased with increasing clay content (Figure S5g). Consistent findings have been reported by Six et al. (2006) and Liang et al. (2017), demonstrating that soils with higher clay and silt contents generally harbor greater quantities of microbial residues, particularly fungal-derived residues, due to the enhanced formation of stable organo-mineral complexes promoted by abundant fine soil particles. Moreover, agricultural management practices frequently disturb soil structure but simultaneously enhance clay enrichment and aggregate formation, thereby providing effective physical

删除了: On the contrary, geographical factor (elevation) was identified as the most influential predictors of the FNC/BNC ratio in both agricultural and natural ecosystems (Figures 4e, f, 5f, 6f), and the FNC/BNC ratio increased with increasing elevation (Figure S7a). ...

944 4.3 Limitations and uncertainties

945 Although the present study provides important insights on global patterns and drivers
946 of soil microbial necromass in agricultural and natural ecosystems, we must clarify
947 two limitations. First, the limited data available on microbial characteristics, such as
948 microbial community composition, enzymatic activities, and the content of soil
949 aggregates and minerals hinder exploration of the drivers of soil microbial necromass.
950 Second, it is undeniable that our dataset is unevenly distributed, primarily
951 concentrating on the Northern Hemisphere, with sparse or nearly no data from other
952 regions such as Africa, South America, and Australia (Figure 1). Additionally, the
953 natural ecosystems in this study were limited to forests and grasslands, excluding
954 other natural habitats such as wetlands and deserts. The uneven distribution of data
955 may reduce the universality of MNC as a key driver of soil carbon pools in global
956 terrestrial ecosystems. Furthermore, the compiled studies employed varied
957 methodologies regarding sampling time, depth, and laboratory protocols. While such
958 heterogeneity is an inherent challenge in global meta-analyses, it likely introduces
959 additional variability and may constrain the direct comparability of certain data points.
960 Therefore, more standardized data from these important areas and biomes are clearly
961 required, and further investigation is warranted to fill the data gaps regarding the
962 contribution of MNC to SOC in terrestrial ecosystems.

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963 **5 Data availability**

964 The data and R code for this manuscript are available at
965 <https://doi.org/10.6084/m9.figshare.28827383> (Lu, 2025).

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967 **6 Conclusions**

968 Our results indicate that, on average, fungal necromass carbon (FNC) contributes
969 approximately twice as much to soil organic carbon (SOC) as bacterial necromass
970 carbon (BNC) in both agricultural and natural ecosystems. The relative contributions
971 of FNC and BNC to SOC were found to be higher in agricultural ecosystems—an
972 effect that is mediated by differences in soil physicochemical factors. The FNC/BNC
973 ratio was significantly higher in natural ecosystems than in agricultural ecosystems,
974 albeit with a modest effect size, and was primarily driven by geographical
975 factors—particularly elevation. Our findings demonstrate that, despite considerable
976 variability among individual sampling sites, statistically significant differences exist
977 between agricultural and natural ecosystems in the contributions of fungal and
978 bacterial necromass carbon (FNC and BNC) to soil organic carbon (SOC), as well as
979 in the FNC/BNC ratio, at a global scale. These results underscore a potential
980 fundamental divergence in the pathways and mechanisms of carbon turnover and
981 stabilization between these two broad ecosystem types. These insights provide novel
982 evidence that ecosystem management type (agricultural versus natural) is a key
983 determinant of the pathways through which microbial necromass contributes to the
984 global soil organic carbon (SOC) pool. Future studies that integrate microbial
985 community composition with necromass dynamics across a broader range of biomes
986 will be essential to predict ecosystem-specific responses of this critical carbon pool to
987

删除了: Our results suggest that the contribution of FNC to SOC is significantly greater than that of BNC in both agricultural and natural ecosystems. The relative

删除了: The FNC/BNC ratio was significantly higher in natural ecosystems compared to agricultural ecosystems, primarily driven by geographical factors (elevation)...

删除了: . Our results reveal consistent trends in the contributions of FNC and BNC to SOC, and the FNC/BNC ratio, which are indicative of ecosystem-level differences in C turnover and stabilization mechanisms across the globe.

1001 [global change](#),

1002

1003 **Author contributions**

1004 JL performed the data analysis and prepared the original draft. TC and MDB
1005 contributed to manuscript review and editing. WL and HS contributed to data
1006 collection. YJ contributed to data analysis. ZW supervised the project and contributed
1007 to the original draft.

1008

1009 **Competing interests**

1010 The contact author has declared that none of the authors has any competing interests.

1011

1012 **Disclaimer**

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1026

1027 **References**

1028 [Anderson, T. H., and Domsch, K. H.: Ratios of microbial biomass carbon to total
1029 organic carbon in arable soils, *Soil Biol. Biochem.*, 21, 471–479,
1030 \[https://doi.org/10.1016/0038-0717\\(89\\)90117-X\]\(https://doi.org/10.1016/0038-0717\(89\)90117-X\), 1989.](#)

1031 [Angst, G., Mueller, K. E., Nierop, K. G., and Simpson, M. J.: Plant-or
1032 microbial-derived? A review on the molecular composition of stabilized soil
1033 organic matter, *Soil Biol. Biochem.*, 156, 108189,
1034 <https://doi.org/10.1016/j.soilbio.2021.108189>, 2021.](#)

1035 [Bellamy, P. H., Loveland, P. J., Bradley, R. I., Lark, R. M., and Kirk, G. J.: Carbon
1036 losses from all soils across England and Wales 1978–2003, *Nature*, 437, 245–248,
1037 <https://doi.org/10.1038/nature04038>, 2005.](#)

1038 [Bohan, D. A., Raybould, A., Mulder, C., Woodward, G., Tamaddoni-Nezhad, A.,
1039 Bluthgen, N., Pocock, M. J. O., Muggleton, S., Evans, D. M., Astegiano, J.,
1040 Massol, F., Loeuille, N., Petit, S., and Macfadyen, S.: Networking agroecology:
1041 integrating the diversity of agroecosystem interactions, *Adv. Ecol. Res.*, 49, 1–67,
1042 <https://doi.org/10.1016/B978-0-12-420002-9.00001-9>, 2013.](#)

1043 [Campitelli, E.: *ggnewscale*: Multiple Fill and Colour Scales in 'ggplot2', R package
1044 version 0.4.8., <https://CRAN.R-project.org/package=ggnewscale>, 2022.](#)

删除了: These insights are conducive to a better mechanistic understanding of the role of microbes in the C cycle in terrestrial ecosystems, and could inform strategies for C sequestration and sustainable land management strategies in the context of global change.

1050 [Cao, Y., Ding, J., Li, J., Xin, Z., Ren, S., and Wang, T.: Necromass-derived soil](#)
1051 [organic carbon and its drivers at the global scale, *Soil Biol. Biochem.*, 181,](#)
1052 [109025, <https://doi.org/10.1016/j.soilbio.2023.109025>, 2023.](#)

1053 [Castellano, M. J., Mueller, K. E., Olk, D. C., Sawyer, J. E., and Six, J.: Integrating](#)
1054 [plant litter quality, soil organic matter stabilization, and the carbon saturation](#)
1055 [concept, *Global Change Biol.*, 21, 3200–3209, <https://doi.org/10.1111/gcb.12982>,](#)
1056 [2015.](#)

1057 [Chen, G., Ma, S., Tian, D., Xiao, W., Jiang, L., Xing, A., Zou, A., Zhou, L., Shen, H.,](#)
1058 [Zheng, C., Ji, C., He, H., Zhu, B., Liu, L., and Fang, J.: Patterns and](#)
1059 [determinants of soil microbial residues from tropical to boreal forests, *Soil Biol.*](#)
1060 [1061 \[Biochem.\]\(#\), 151, 108059, <https://doi.org/10.1016/j.soilbio.2020.108059>, 2020.](#)

1061 [Chen, X., Hu, Y., Xia, Y., Zheng, S., Ma, C., Rui, Y., He, H., Huang, D., Zhang, Z., Ge,](#)
1062 [T., Wu, J., Guggenberger, G., Kuzyakov, Y., and Su, Y.: Contrasting pathways of](#)
1063 [carbon sequestration in paddy and upland soils, *Global Change Biol.*, 27, 2478–](#)
1064 [2490, <https://doi.org/10.1111/gcb.15595>, 2021.](#)

1065 [Choi, J., Bach, E., Lee, J., Flater, J., Dooley, S., Howe, A., and Hofmockel, K. S.:](#)
1066 [Spatial structuring of cellulase gene abundance and activity in soil, *Front.*](#)
1067 [1068 \[Environ. Sci.\]\(#\), 6, 107, <https://doi.org/10.3389/fenvs.2018.00107>, 2018.](#)

1068 [Clocchiatti, A., Hannula, S. E., Hundscheid, M. P., Klein Gunnewiek, P. J., and de](#)
1069 [Boer, W.: Stimulated saprotrophic fungi in arable soil extend their activity to the](#)
1070 [rhizosphere and root microbiomes of crop seedlings, *Environ. Microbiol.*, 23,](#)
1071 [6056–6073, <https://doi.org/10.1111/1462-2920.15563>, 2021.](#)

1072 [Cotrufo, M. F., Wallenstein, M. D., Boot, C. M., Denef, K., and Paul, E.: The](#)
1073 [Microbial Efficiency-Matrix Stabilization \(MEMS\) framework integrates plant](#)
1074 [litter decomposition with soil organic matter stabilization: do labile plant inputs](#)
1075 [form stable soil organic matter?, *Global Change Biol.*, 19, 988–995,](#)
1076 [1077 \[https://doi.org/10.1111/gcb.12113\]\(#\), 2013.](#)

1077 [Crowther, T. W., Sokol, N. W., Oldfield, E. E., Maynard, D. S., Thomas, S. M., and](#)
1078 [Bradford, M. A.: Environmental stress response limits microbial necromass](#)
1079 [contributions to soil organic carbon, *Soil Biol. Biochem.*, 85, 153–161,](#)
1080 [1081 \[https://doi.org/10.1016/j.soilbio.2015.03.002\]\(#\), 2015.](#)

1081 [Cui, W., Li, R., Fan, Z., Wu, L., Zhao, X., Wei, G., and Shu, D.: Weak environmental](#)
1082 [adaptation of rare phylotypes sustaining soil multi-element cycles in response to](#)
1083 [decades-long fertilization, *STOTEN.*, 871, 162063,](#)
1084 [1085 \[https://doi.org/10.1016/j.scitotenv.2023.162063\]\(#\), 2023.](#)

1085 [de Boer, W. D., Folman, L. B., Summerbell, R. C., and Boddy, L.: Living in a fungal](#)
1086 [world: impact of fungi on soil bacterial niche development, *FEMS Microbiol.*](#)
1087 [1088 \[Rev.\]\(#\), 29, 795–811, <https://doi.org/10.1016/j.femsre.2004.11.005>, 2005.](#)

1088 [Deng, F., and Liang, C.: Revisiting the quantitative contribution of microbial](#)
1089 [necromass to soil carbon pool: stoichiometric control by microbes and soil, *Soil*](#)
1090 [1091 \[Biol. Biochem.\]\(#\), 165, 108486, <https://doi.org/10.1016/j.soilbio.2021.108486>,](#)
1092 [2022.](#)

1092 [Ding, Z., Mou, Z., Li, Y., Liang, C., Xie, Z., Wang, J., Hui, D., Lambers, H., Sardans,](#)
1093 [J., Peñuelas, J., Xu, H., and Liu, Z.: Spatial variation and controls of soil](#)

1094 [microbial necromass carbon in a tropical montane rainforest, STOTEN., 921,](#)
1095 [170986, https://doi.org/10.1016/j.scitotenv.2024.170986, 2024.](#)

1096 [Elser, J. J., Bracken, M. E., Cleland, E. E., Gruner, D. S., Harpole, W. S., Hillebrand,](#)
1097 [H., Ngai, J. T., Seabloom, E. W., Shurin, J. B., and Smith, J. E.: Global analysis](#)
1098 [of nitrogen and phosphorus limitation of primary producers in freshwater, marine](#)
1099 [and terrestrial ecosystems, Ecol. Lett., 10, 1135–1142,](#)
1100 [https://doi.org/10.1111/j.1461-0248.2007.01113.x](#), 2007.

1101 [Fanin, N., Bezaud, S., Sarneel, J. M., Cecchini, S., Nicolas, M., and Augusto, L.:](#)
1102 [Relative importance of climate, soil and plant functional traits during the early](#)
1103 [decomposition stage of standardized litter, Ecosystems, 23, 1004–1018,](#)
1104 [https://doi.org/10.1007/s10021-019-00452-z](#), 2020.

1105 [Gao, W., Duan, X., Chen, X., Wei, L., Wang, S., Wu, J., and Zhu, Z.: Iron-carbon](#)
1106 [complex types and bonding forms jointly control organic carbon mineralization](#)
1107 [in paddy soils, STOTEN., 953, 176117,](#)
1108 [https://doi.org/10.1016/j.scitotenv.2024.176117](#), 2024.

1109 [Ghezzehei, T. A., Sulman, B., Arnold, C. L., Bogie, N. A., and Berhe, A. A.: On the](#)
1110 [role of soil water retention characteristic on aerobic microbial respiration,](#)
1111 [BIOGEOSCIENCES., 16, 1187–1209, https://doi.org/10.5194/bg-16-1187-2019,](#)
1112 [2019.](#)

1113 [Greenwell, B., Boehmke, B., Cunningham, J., and Developers, G. B. M.: *gbm*:](#)
1114 [Generalized Boosted Regression Models, R package version 2.1.8.1.,](#)
1115 [https://CRAN.R-project.org/package=gbm](#), 2022.

1116 [Han, B., Yao, Y., Wang, Y., Su, X., Ma, L., Chen, X., and Li, Z.: Microbial traits](#)
1117 [dictate soil necromass accumulation coefficient: A global synthesis, Global Ecol.](#)
1118 [Biogeogr., 33, 151–161, https://doi.org/10.1111/geb.13776, 2024.](#)

1119 [Hao, Z., Zhao, Y., Wang, X., Wu, J., Jiang, S., Xiao, J., Wang, K., Zhou, X., Liu, H., Li,](#)
1120 [J., and Sun, Y.: Thresholds in aridity and soil carbon-to-nitrogen ratio govern the](#)
1121 [accumulation of soil microbial residues, Commun. Earth Environ., 2, 236,](#)
1122 [https://doi.org/10.1038/s43247-021-00306-4](#), 2021.

1123 [Hättenschwiler, S., Tiunov, A. V., and Scheu, S.: Biodiversity and litter decomposition](#)
1124 [in terrestrial ecosystems, Annu. Rev. Ecol. Evol. Syst., 36, 191–218,](#)
1125 [https://doi.org/10.1146/annurev.ecolsys.36.112904.151932](#), 2005.

1126 [Hobbs, R. J., Hallett, L. M., Ehrlich, P. R., and Mooney, H. A.: Intervention ecology:](#)
1127 [applying ecological science in the twenty-first century, BioScience, 61, 442–450,](#)
1128 [https://doi.org/10.1525/bio.2011.61.6.6](#), 2011.

1129 [Hollister, J. W.: *elevatr*: Access Elevation Data from Various APIs, R package version](#)
1130 [0.4.2., https://CRAN.R-project.org/package=elevatr/, 2021.](#)

1131 [Kallenbach, C. M., Frey, S. D., and Grandy, A. S.: Direct evidence for](#)
1132 [microbial-derived soil organic matter formation and its ecophysiological controls,](#)
1133 [Nat. Commun., 7, 13630, https://doi.org/10.1038/ncomms13630, 2016.](#)

1134 [Kassambara, A.: *ggpubr*: 'ggplot2' Based Publication Ready Plots, R package version](#)
1135 [0.5.0., https://CRAN.R-project.org/package=ggpubr, 2022.](#)

1136 [Keith, D. A., Ferrer-Paris, J. R., Nicholson, E., Bishop, M. J., Polidoro, B. A.,](#)
1137 [Ramirez-Llodra, E., Tozer, M. G., Nel, J. L., Nally, R. M., Gregr, E. J.,](#)

1138 [Watermeyer, K. E., Essl, F., Faber-Langendoen, D., Franklin, J., Lehmann, C. E.](#)
 1139 [R., Etter, A., Roux, D. J., Stark, J. S., Rowland, J. A., Brummitt, N. A.,](#)
 1140 [Fernandez-Arcaya, U. C., Suthers, I. M., Wisser, S. K., Donohue, I., Jackson, L. J.,](#)
 1141 [Pennington, R. T., Iliffe, T. M., Gerovasileiou, V., Giller, P., Robson, B. J.,](#)
 1142 [Pettorelli, N., Andrade, A., Lindgaard, A., Tahvanainen, T., Terauds, A.,](#)
 1143 [Chadwick, M. A., Murray, N. J., Moat, J., Pliscoff, P., Zager, I., and Kingsford, R.](#)
 1144 [T.: A function-based typology for Earth's ecosystems, *Nature*, 610, 513–518,](#)
 1145 <https://doi.org/10.1038/s41586-022-05318-4>, 2022.

1146 [Kleber, M., Bourg, I. C., Coward, E. K., Hansel, C. M., Myneni, S. C., and Nunan, N.:](#)
 1147 [Dynamic interactions at the mineral–organic matter interface, *NAT REV EARTH*](#)
 1148 [ENV.](#), 2, 402–421, <https://doi.org/10.1038/s43017-021-00162-y>, 2021.

1149 [Klink, S., Keller, A. B., Wild, A. J., Baumert, V. L., Gube, M., Lehndorff, E., Meyer,](#)
 1150 [N., Mueller, C. W., Phillips, R. P., and Pausch, J.: Stable isotopes reveal that](#)
 1151 [fungal residues contribute more to mineral-associated organic matter pools than](#)
 1152 [plant residues, *Soil Biol. Biochem.*, 168, 108634,](#)
 1153 <https://doi.org/10.1016/j.soilbio.2022.108634>, 2022.

1154 [Kock, N.: Common method bias in PLS-SEM: A full collinearity assessment approach,](#)
 1155 [International Journal of e-Collaboration \(IJeC\), 11, 1–10,](#)
 1156 <https://doi.org/10.4018/ijec.2015100101>, 2015.

1157 [Körner, C.: The use of 'altitude' in ecological research, *Trends Ecol. Evol.*, 22, 569–](#)
 1158 [574, <https://doi.org/10.1016/j.tree.2007.09.006>, 2007.](#)

1159 [Lembrechts, J. J., van den Hoogen, J., Aalto, J., Ashcroft, M. B., De Frenne, P.,](#)
 1160 [Kempinen, J., Kopecký, M., Luoto, Maclean, M. I. M. D., Crowther, T. W.,](#)
 1161 [Bailey, J. J., Haesen, S., Klinges, D. H., Niittynen, P., Scheffers, B. R., Van](#)
 1162 [Meerbeek, K., Aartsma, P., Abdalaze, O., Abedi, M., Aerts, R., Ahmadian, N.,](#)
 1163 [Ahrends, A., Alatalo, J. M., Alexander, J. M., Allonsius, C. N., Altman, J.,](#)
 1164 [Ammann, C., Andres, C., Andrews, C., Ardö, J., Arriga, N., Arzac, A., Aschero,](#)
 1165 [V., Assis, R. L., Assmann, J. J., Bader, M. Y., Bahalkeh, K., Barančok, P., Barrio,](#)
 1166 [I. C., Barros, A., Barthe, M., Basham, E. W., Bauters, M., Bazzichetto, M.,](#)
 1167 [Marchesini, L. B., Bell, M. C., Benavides, J. C., Alonso, J. L. B., Berauer, B. J.,](#)
 1168 [Bjerke, J. W., Björk, R. G., Björkman, M. P., Björnsdóttir, K., Blonder, B.,](#)
 1169 [Boeckx, P., Boike, J., Bokhorst, S., Brum, B. N. S., Brûna, J., Buchmann, N.,](#)
 1170 [Buysse, P., Camargo, J. L., Campoe, O. C., Candan, O., Canessa, R., Cannone,](#)
 1171 [N., and Hik, D. S.: Global maps of soil temperature, *Global Change Biol.*, 28,](#)
 1172 [3110–3144, <https://doi.org/10.1111/gcb.16060>, 2022.](#)

1173 [Lehmann, J., Hansel, C. M., Kaiser, C., Kleber, M., Maher, K., Manzoni, S., Nunan,](#)
 1174 [N., Reichstein, M., Schimel, J. P., Torn, M. S., Wieder, W. R., and](#)
 1175 [Kögel-Knabner, I.: Persistence of soil organic carbon caused by functional](#)
 1176 [complexity, *Nat. Geosci.*, 13, 529–534,](#)
 1177 <https://doi.org/10.1038/s41561-020-0612-3>, 2020.

1178 [Lenardon, M. D., Whitton, R. K., Munro, C. A., Marshall, D., and Gow, N. A. R.:](#)
 1179 [Individual chitin synthase enzymes synthesize microfibrils of differing structure](#)
 1180 [at specific locations in the *Candida albicans* cell wall, *Mol. Microbiol.*, 66,](#)
 1181 [1164–1173, <https://doi.org/10.1111/j.1365-2958.2007.05990.x>, 2007.](#)

1182 [Liang, C., Amelung, W., Lehmann, J., and Kästner, M.: Quantitative assessment of](#)
1183 [microbial necromass contribution to soil organic matter, *Global Change Biol.*, 25,](#)
1184 [3578–3590, <https://doi.org/10.1111/gcb.14781>, 2019.](#)

1185 [Liang, C., and Balser, T. C.: Microbial production of recalcitrant organic matter in](#)
1186 [global soils: implications for productivity and climate policy, *Nat. Rev.*](#)
1187 [Microbiol., 9, 75–75, <https://doi.org/10.1038/nrmicro2386-c1>, 2011.](#)

1188 [Liang, C., Schimel, J. P., and Jastrow, J. D.: The importance of anabolism in microbial](#)
1189 [control over soil carbon storage, *Nat. Microbiol.*, 2, 17105,](#)
1190 [<https://doi.org/10.1038/nmicrobiol.2017.105>, 2017.](#)

1191 [Liu, C., Tian, J., Cheng, K., Xu, X., Wang, Y., Liu, X., Liu, Z., Bian, R., Zhang, X.,](#)
1192 [Xia, S., Zheng, J., Li, L., and Pan, G.: Topsoil microbial biomass carbon pool](#)
1193 [and the microbial quotient under distinct land-use types across China: A data](#)
1194 [synthesis, *SSE.*, 2, 5, <https://doi.org/10.48130/SSE-2023-0005>, 2023.](#)

1195 [Liu, X., Tian, Y., Heinze, J., Salas, E., Kwatcho-Kengdo, S., Borken, W.,](#)
1196 [Schindlbacher, A., and Wanek, W.: Long-term soil warming decreases soil](#)
1197 [microbial necromass carbon by adversely affecting its production and](#)
1198 [decomposition, *Global Change Biol.*, 30, e17379,](#)
1199 [<https://doi.org/10.1111/gcb.17379>, 2024.](#)

1200 [Li, Y., Wang, S., Yang, Y., Ren, L., Wang, Z., Liao, Y., and Yong, T.: Global synthesis](#)
1201 [on the response of soil microbial necromass carbon to climate-smart agriculture,](#)
1202 [*Global Change Biol.*, 30\(5\), e17302, <https://doi.org/10.1111/gcb.17302>, 2024.](#)

1203 [Lu, J.: Microbial necromass contribution to topsoil organic carbon storage of natural](#)
1204 [and agricultural ecosystems, *figshare* \[data set\],](#)
1205 [<https://doi.org/10.6084/m9.figshare.28827383>, 2025.](#)

1206 [Luo, R., Kuzyakov, Y., Zhu, B., Qiang, W., Zhang, Y., and Pang, X.: Phosphorus](#)
1207 [addition decreases plant lignin but increases microbial necromass contribution to](#)
1208 [soil organic carbon in a subalpine forest, *Global Change Biol.*, 28, 4194–4210,](#)
1209 [<https://doi.org/10.1111/gcb.16205>, 2022.](#)

1210 [Malik, A. A., Chowdhury, S., Schlager, V., Oliver, A., Puissant, J., Vazquez, P. G.,](#)
1211 [Jehmlich, N., von Bergen, M., Griffiths, R., and Gleixner, G.: Soil fungal:](#)
1212 [bacterial ratios are linked to altered carbon cycling, *Front. Microbiol.*, 7, 1247,](#)
1213 [<https://doi.org/10.3389/fmicb.2016.01247>, 2016.](#)

1214 [Ma, T., Zhu, S., Wang, Z., Chen, D., Dai, G., Feng, B., Su, X., Hu, H., Li, K., Han, W.,](#)
1215 [Liang, C., Bai, Y., and Feng, X.: Divergent accumulation of microbial necromass](#)
1216 [and plant lignin components in grassland soils, *Nat. Commun.*, 9, 3480,](#)
1217 [<https://doi.org/10.1038/s41467-018-05891-1>, 2018.](#)

1218 [Mooshammer, M., Wanek, W., Zechmeister-Boltenstern, S., and Richter, A.:](#)
1219 [Stoichiometric imbalances between terrestrial decomposer communities and their](#)
1220 [resources: mechanisms and implications of microbial adaptations to their](#)
1221 [resources, *Front. Microbiol.*, 5, 22, <https://doi.org/10.3389/fmicb.2014.00022>,](#)
1222 [2014.](#)

1223 [Mou, Z., Kuang, L., He, L., Zhang, J., Zhang, X., Hui, D., Li, Y., Wu, W., Mei, Q., He,](#)
1224 [X., Kuang, Y., Wang, J., Wang, Y., Lambers, H., Sardans, J., Peñuelas, J., and Liu,](#)
1225 [Z.: Climatic and edaphic controls over the elevational pattern of microbial](#)

1226 [necromass in subtropical forests, *Catena*, 207, 105707,](#)
1227 <https://doi.org/10.1016/j.catena.2021.105707>, 2021.

1228 [Ni, X., Liao, S., Tan, S., Peng, Y., Wang, D., Yue, K., Wu, F., and Yang, Y.: The](#)
1229 [vertical distribution and control of microbial necromass carbon in forest soils,](#)
1230 [Global Ecol. Biogeogr., 29, 1829–1839, https://doi.org/10.1111/geb.13159, 2020.](#)

1231 [Oksanen, J., Blanchet, F. G., Friendly, M., Kindt, R., Legendre, P., McGlenn, D., and](#)
1232 [Wagner, H.: *vegan*: Community Ecology Package, R package version 2.5.7.,](#)
1233 <https://CRAN.R-project.org/package=vegan>, 2020.

1234 [Poggio, L., De Sousa, L. M., Batjes, N. H., Heuvelink, G., Kempen, B., Ribeiro, E.,](#)
1235 [and Rossiter, D.: SoilGrids 2.0: producing soil information for the globe with](#)
1236 [quantified spatial uncertainty, *Soil*, 7, 217–240,](#)
1237 <https://doi.org/10.5194/soil-7-217-2021>, 2021.

1238 [Qiu, H., Zheng, X., Ge, T., Dorodnikov, M., Chen, X., Hu, Y., Kuzyakov, Y., Wu, J.,](#)
1239 [Su, Y., and Zhang, Z.: Weaker priming and mineralisation of low molecular](#)
1240 [weight organic substances in paddy than in upland soil, *Eur. J. Soil Biol.*, 83, 9–](#)
1241 [17, https://doi.org/10.1016/j.ejsobi.2017.09.008, 2017.](#)

1242 [R Core Team: R: A language and environment for statistical computing, R Foundation](#)
1243 [for Statistical Computing, Vienna, Austria, https://www.R-project.org, 2021.](#)

1244 [Rosseel, Y: *lavaan*: An R package for structural equation modeling, *J. Stat. Softw.*, 48,](#)
1245 [1–36, https://doi.org/10.18637/jss.v048.i02, 2012.](#)

1246 [Sae-Tun, O., Bodner, G., Rosinger, C., Zechmeister-Boltenstern, S., Mentler, A., and](#)
1247 [Keiblinger, K.: Fungal biomass and microbial necromass facilitate soil carbon](#)
1248 [sequestration and aggregate stability under different soil tillage intensities, *Appl.*](#)
1249 [Soil Ecol., 179, 104599, https://doi.org/10.1016/j.apsoil.2022.104599, 2022.](#)

1250 [Sanauallah, M., Usman, M., Wakeel, A., Cheema, S. A., Ashraf, I., and Farooq, M.:](#)
1251 [Terrestrial ecosystem functioning affected by agricultural management systems:](#)
1252 [A review, *Soil Tillage Res.*, 196, 104464,](#)
1253 <https://doi.org/10.1016/j.still.2019.104464>, 2020.

1254 [Six, J., Frey, S. D., Thiet, R. K., and Batten, K. M.: Bacterial and fungal contributions](#)
1255 [to carbon sequestration in agroecosystems, *Soil Sci. Soc. Am. J.*, 70, 555–569,](#)
1256 <https://doi.org/10.2136/sssaj2004.0347>, 2006.

1257 [Spawn, S. A., Sullivan, C. C., Lark, T. J., and Gibbs, H. K.: Harmonized global maps](#)
1258 [of above and belowground biomass carbon density in the year 2010, *Sci. Data*, 7,](#)
1259 [112, https://doi.org/10.1038/s41597-020-0444-4, 2020.](#)

1260 [Strickland, M. S., and Rousk, J.: Considering fungal: bacterial dominance in soils–](#)
1261 [methods, controls, and ecosystem implications, *Soil Biol. Biochem.*, 42, 1385–](#)
1262 [1395, https://doi.org/10.1016/j.soilbio.2010.05.007, 2010.](#)

1263 [Štursová, M., Žifčáková, L., Leigh, M. B., Burgess, R., and Baldrian, P.: Cellulose](#)
1264 [utilization in forest litter and soil: identification of bacterial and fungal](#)
1265 [decomposers, *FEMS Microbiol. Ecol.*, 80, 735–746,](#)
1266 <https://doi.org/10.1111/j.1574-6941.2012.01343.x>, 2012.

1267 [van Der Heijden, M. G., Bardgett, R. D., and van Straalen, N. M.: The unseen](#)
1268 [majority: soil microbes as drivers of plant diversity and productivity in terrestrial](#)

1269 ecosystems, Ecol. Lett., 11, 296–310,
1270 <https://doi.org/10.1111/j.1461-0248.2007.01139.x>, 2008.

1271 Wang, B., An, S., Liang, C., Liu, Y., and Kuzyakov, Y.: Microbial necromass as the
1272 source of soil organic carbon in global ecosystems, *Soil Biol. Biochem.*, 162,
1273 108422, <https://doi.org/10.1016/j.soilbio.2021.108422>, 2021a.

1274 Wang, B., Liang, C., Yao, H., Yang, E., and An, S.: The accumulation of microbial
1275 necromass carbon from litter to mineral soil and its contribution to soil organic
1276 carbon sequestration, *Catena*, 207, 105622,
1277 <https://doi.org/10.1016/j.catena.2021.105622>, 2021b.

1278 Wang, C., and Kuzyakov, Y.: Mechanisms and implications of bacterial–fungal
1279 competition for soil resources, *ISME J.*, 18, wrae073,
1280 <https://doi.org/10.1093/ismejo/wrae073>, 2024.

1281 Wang, Z., Zhao, M., Yan, Z., Yang, Y., Niklas, K. J., Huang, H., Mipam, T. D., He, X.,
1282 Hu, H., and Wright, S. J.: Global patterns and predictors of soil microbial
1283 biomass carbon, nitrogen, and phosphorus in terrestrial ecosystems, *Catena*, 211,
1284 106037, <https://doi.org/10.1016/j.catena.2022.106037>, 2022.

1285 Wickham, H.: *ggplot2: elegant graphics for data analysis*. Springer-Verlag New York,
1286 <https://ggplot2.tidyverse.org>, 2016.

1287 Wu, H., Xiang, W., Ouyang, S., Forrester, D. I., Zhou, B., Chen, L., Ge, T., Lei, P.,
1288 Chen, L., Zeng, Y., Song, X., Peñuelas, J., and Peng, C.: Linkage between tree
1289 species richness and soil microbial diversity improves phosphorus bioavailability,
1290 *Funct. Ecol.*, 33, 1549–1560, <https://doi.org/10.1111/1365-2435.13355>, 2019.

1291 Wu, W., Feng, J., Wang, X., Xiao, J., Qin, W., and Zhu, B.: The response of soil
1292 microbial necromass carbon to global change: A global meta-analysis, *Catena*,
1293 249, 108693, <https://doi.org/10.1016/j.catena.2024.108693>, 2025.

1294 Xu, S., Song, X., Zeng, H., and Wang, J.: Soil microbial necromass carbon in forests:
1295 A global synthesis of patterns and controlling factors, *Soil Ecol. Lett.*, 6(4),
1296 240237, <https://doi.org/10.1007/s42832-024-0237-3>, 2024.

1297 Xu, Y., Sun, L., Gao, X., and Wang, J.: Contrasting response of fungal versus bacterial
1298 residue accumulation within soil aggregates to long-term fertilization, *Sci. Rep.*,
1299 12, 17834, <https://doi.org/10.1038/s41598-022-22064-9>, 2022.

1300 Yu, K., van den Hoogen, J., Wang, Z., Averill, C., Routh, D., Smith, G. R., Drenovsky,
1301 R. E., Scow, K. M., Mo, F., Waldrop, M. P., Yang, Y., Tang, W., Vries, F. T. D.,
1302 Bardgett, R. D., Manning, P., Bastida, F., Baer, S. G., Bach, E. M., García, C.,
1303 Wang, Q., Ma, L., Chen, B., He, X., Teurlinx, S., Heijboer, A., Bradley, J. A.,
1304 and Crowther, T. W.: The biogeography of relative abundance of soil fungi
1305 versus bacteria in surface topsoil, *Earth Syst. Sci. Data*, 14, 4339–4350,
1306 <https://doi.org/10.5194/essd-14-4339-2022>, 2022.

1307 Zhang, B., Zhu, S., Guo, L., Chen, G., Zhang, G., and Li, J.: Elevation-dependent
1308 distribution of soil microbial necromass carbon in *Pinus densata* Mast. Forests,
1309 *Appl. Soil Ecol.*, 209, 106049, <https://doi.org/10.1016/j.apsoil.2025.106049>,
1310 2025.

1311 Zhang, Q., Li, X., Liu, J., Liu, J., Han, L., Wang, X., Liu, H., Xu, M., Yang, G., Ren,
1312 C., and Han, X.: The contribution of microbial necromass carbon to soil organic

1313 carbon in soil aggregates, *Appl. Soil Ecol.*, 190, 104985,
 1314 <https://doi.org/10.1016/j.apsoil.2023.104985>, 2023.
 1315 Zhang, X., Jia, J., Chen, L., Chu, H., He, J. S., Zhang, Y., and Feng, X.: Aridity and
 1316 NPP constrain contribution of microbial necromass to soil organic carbon in the
 1317 Qinghai-Tibet alpine grasslands, *Soil Biol. Biochem.*, 156, 108213,
 1318 <https://doi.org/10.1016/j.soilbio.2021.108213>, 2021.
 1319 Zhao, M., and Running, S. W.: Drought-induced reduction in global terrestrial net
 1320 primary production from 2000 through 2009, *Science*, 329, 940–943,
 1321 <https://doi.org/10.1126/science.1192666>, 2010.
 1322 Zhao, X., Tian, P., Liu, S., Yin, P., Sun, Z., and Wang, Q.: Mean annual temperature
 1323 and carbon availability respectively controlled the contributions of bacterial and
 1324 fungus necromass to organic carbon accumulation in topsoil across China's
 1325 forests, *Global Ecol. Biogeogr.*, 32, 120–131, <https://doi.org/10.1111/gcb.13605>,
 1326 2023.
 1327 Zhou, R., Liu, Y., Dungait, J. A., Kumar, A., Wang, J., Tiemann, L. K., Zhang, F.,
 1328 Kuzyakov, Y., and Tian, J.: Microbial necromass in cropland soils: A global
 1329 meta-analysis of management effects, *Global Change Biol.*, 29, 1998–2014,
 1330 <https://doi.org/10.1111/gcb.16613>, 2023.
 1331

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上移了 [1]: Zhang, X., Jia, J., Chen, L., Chu, H., He, J.S., Zhang, Y., Feng, X., 2021. Aridity and NPP constrain contribution of microbial necromass to soil organic carbon in the Qinghai-Tibet alpine grasslands. *Soil Biol. Biochem.* 156, 108213. <https://doi.org/10.1016/j.soilbio.2021.108213>

删除了: Angst, G., Mueller, K.E., Nierop, K.G., Simpson, M.J., 2021. Plant-or microbial-derived? A review on the molecular composition of stabilized soil organic matter. *Soil Biol. Biochem.* 156, 108189.

<https://doi.org/10.1016/j.soilbio.2021.108189>

Bellamy, P.H., Loveland, P.J., Bradley, R.I., Lark, R.M., Kirk, G.J., 2005. Carbon losses from all soils across England and Wales 1978–2003. *Nature* 437, 245–248.

<https://doi.org/10.1038/nature04038>

Bohan, D.A., Raybould, A., Mulder, C., Woodward, G., Tamaddoni-Nezhad, A., Bluthgen, N., Pockock, M.J.O., Muggleton, S., Evans, D.M., Astegiano, J., Massol, F., Loeuille, N., Petit, S., Macfadyen, S., 2013. Networking agroecology: integrating the diversity of agroecosystem interactions. *Adv. Ecol. Res.* 49, 1–67.

<https://doi.org/10.1016/B978-0-12-420002-9.00001-9>

Campitelli, E., 2022. *ggnewscale*: Multiple Fill and Colour Scales in 'ggplot2'. R package version 0.4.8.

<https://CRAN.R-project.org/package=ggnewscale>

Cao, Y., Ding, J., Li, J., Xin, Z., Ren, S., Wang, T., 2023. Necromass-derived soil organic carbon and its drivers at the global scale. *Soil Biol. Biochem.* 181, 109025.

<https://doi.org/10.1016/j.soilbio.2023.109025>

Castellano, M.J., Mueller, K.E., Olk, D.C., Sawyer, J.E., Six, J., 2015. Integrating plant litter quality, soil organic matter stabilization, and the carbon saturation concept. *Global Change Biol.* 21, 3200–3209.

<https://doi.org/10.1111/gcb.12982>

Chen, G., Ma, S., Tian, D., Xiao, W., Jiang, L., Xing, A., Zou, A., Zhou, L., Shen, H., Zheng, C., Ji, C., He, H., Zhu, B., Liu, L., Fang, J., 2020. Patterns and determinants of soil microbial residues from tropical to boreal forests. *Soil Biol. Biochem.* 181, 109025.

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删除了: Zhu, Y.G., Miller, R.M., 2003. Carbon cycling by arbuscular mycorrhizal fungi in soil-plant systems. *Trends Plant Sci.* 8, 407–409.

[https://doi.org/10.1016/S1360-1385\(03\)00184-5](https://doi.org/10.1016/S1360-1385(03)00184-5)

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1426 **Table 1. Summary of the contributions of fungal necromass carbon (FNC) and**
 1427 **bacterial necromass carbon (BNC) to SOC and the FNC/BNC ratio in**
 1428 **agricultural and natural ecosystems at the global scale investigated in this study.**

Ecosystem	FNC/SOC (%)		BNC/SOC (%)		FNC/BNC	
	Range	Mean± SE	Range	Mean± SE	Range	Mean± SE
Natural ecosystem^{&} (N = 341)	0.92– 96.29	29.24 ± 0.51 b*	0.25– 89.45	14.02 ± 0.36 b	0.12– 44.24	3.22 ± 0.11 a
<i>Forest</i> (N = 195)	0.92– 96.29	29.11 ± 0.63 A [#]	0.94– 96.47	13.48 ± 0.43 A	0.22– 11.56	2.80 ± 0.07 A
<i>Grassland</i> (N = 146)	0.96– 93.89	26.75 ± 0.74 A	0.25– 89.45	14.34 ± 0.60 A	0.05– 44.24	3.58 ± 0.22 A
Agricultural ecosystem[%] (N = 145)	0.09– 97.53	34.39 ± 0.67 a	0.81– 65.00	15.65 ± 0.33 a	0.02– 12.74	2.61 ± 0.06 b
<i>Dry land</i> (N = 32)	<u>3.01–</u> <u>96.81</u>	<u>37.77 ±</u> <u>1.15 A[#]</u>	<u>0.81–</u> <u>65.00</u>	<u>17.34 ±</u> <u>0.65 A</u>	<u>0.13–</u> <u>9.12</u>	<u>2.87 ±</u> <u>0.12 A</u>
<i>Irrigated cropland</i> (N = 72)	<u>0.09–</u> <u>97.25</u>	<u>35.35 ±</u> <u>0.73 A</u>	<u>1.18–</u> <u>62.47</u>	<u>15.95 ±</u> <u>0.38 A</u>	<u>0.02–</u> <u>12.74</u>	<u>2.51 ±</u> <u>0.06 A</u>
<i>Submerged paddy</i> (N = 41)	<u>4.96–</u> <u>97.53</u>	<u>22.82 ±</u> <u>1.55 B</u>	<u>1.48–</u> <u>30.97</u>	<u>10.55 ±</u> <u>0.66 B</u>	<u>0.31–</u> <u>10.40</u>	<u>2.62 ±</u> <u>0.16 A</u>

1429 Note: N refers to the number of study sites;
 1430 [&]Natural ecosystem includes forest and grassland;
 1431 [%]Agricultural ecosystem includes dry land, irrigated cropland, and submerged paddy;
 1432 *Within the same column, values with different lowercase letters indicate a significant
 1433 difference in the same variable between agricultural and natural ecosystems
 1434 (Wilcoxon rank sum test; P < 0.05);
 1435 [#]Within the same column, values with different capital letters indicate a significant
 1436 difference in the same variable between forests and grasslands (Wilcoxon rank sum
 1437 test), as well as between dry land, irrigated cropland, and submerged paddy (Kruskal–
 1438 Wallis and Dunn's post hoc tests; P < 0.05).

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1442 **Figure legends**

1443 **Figure 1. Global distribution of the sites used in this study.** Ecosystem types are
1444 distinguished by distinct shapes and colors, with the numbers in parentheses
1445 indicating the number of study sites for each ecosystem type.

1446 **Figure 2. Comparison of the contributions of MNC to SOC and their ratio in**
1447 **agricultural and natural ecosystems.** Colors indicate different ecosystems types.
1448 Significance levels: *** $P < 0.001$ and * $P < 0.05$.

1449 **Figure 3. Variations in the contributions of MNC to SOC and their ratio**
1450 **explained by four types of factors in agricultural and natural ecosystems.** Colors
1451 indicate different types of factors.

1452 **Figure 4. Relative influence of different factors on the contributions of MNC to**
1453 **SOC and their ratio in agricultural and natural ecosystems.** MAT, mean annual
1454 temperature; MAP, mean annual precipitation; MBC, microbial biomass carbon; SOC,
1455 soil organic carbon; C/N, the ratio of SOC to total nitrogen (TN); MBC/MBN, the
1456 ratio of MBC to microbial biomass nitrogen (MBN); NPP, net primary production;
1457 BGBC, belowground biomass carbon density. Colors indicate different types of
1458 factors.

1459 **Figure 5. The influence pathways of four types of factors on the contributions of**
1460 **MNC to SOC and their ratio in agricultural ecosystems.** Direct and indirect effects
1461 (a, c, e) and the standardized total effects (b, d, f) of different factors on the
1462 contributions of MNC to SOC and their ratio of agricultural ecosystems are shown.
1463 Standardized path coefficients representing the effect sizes of potential causal factors
1464 are indicated by numbers adjacent to arrows. The width of arrows is proportional to
1465 the potential causal effect between variables. The red arrows indicate positive effects,
1466 and the blue arrows indicate negative effects. The numbers adjacent to boxes of
1467 response variables denote the explained variance (R^2). Right-angled rectangles denote
1468 single variables, whereas rounded rectangles represent composite variables. Colors
1469 indicate different types of factors. Significance levels: *** $P < 0.001$, ** $P < 0.01$ and
1470 * $P < 0.05$. The *priori* models are shown in Figure [S3](#).

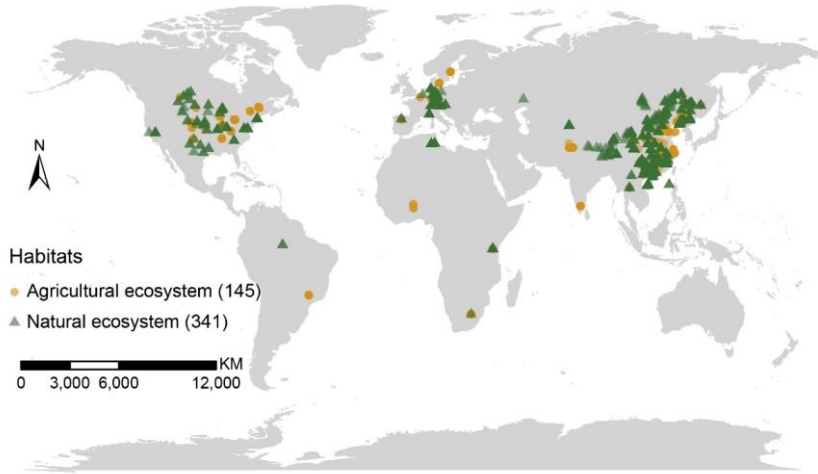
1471 **Figure 6. The influence pathways of four types of factors on the contributions of**
1472 **MNC to SOC and their ratio in natural ecosystems.** Direct and indirect effects (a, c,
1473 e) and the standardized total effects (b, d, f) of different factors on the contributions of
1474 MNC to SOC and their ratio of natural ecosystems are shown. Standardized path
1475 coefficients representing the effect sizes of potential causal factors are indicated by
1476 numbers adjacent to arrows. The width of arrows is proportional to the potential
1477 causal effect between variables. The red arrows indicate positive effects, and the blue
1478 arrows indicate negative effects. The numbers adjacent to boxes of response variables
1479 denote the explained variance (R^2). Right-angled rectangles denote single variables,
1480 whereas rounded rectangles represent composite variables. Colors indicate different
1481 types of factors. Significance levels: *** $P < 0.001$, ** $P < 0.01$ and * $P < 0.05$. The
1482 *priori* models are shown in Figure [S3](#).

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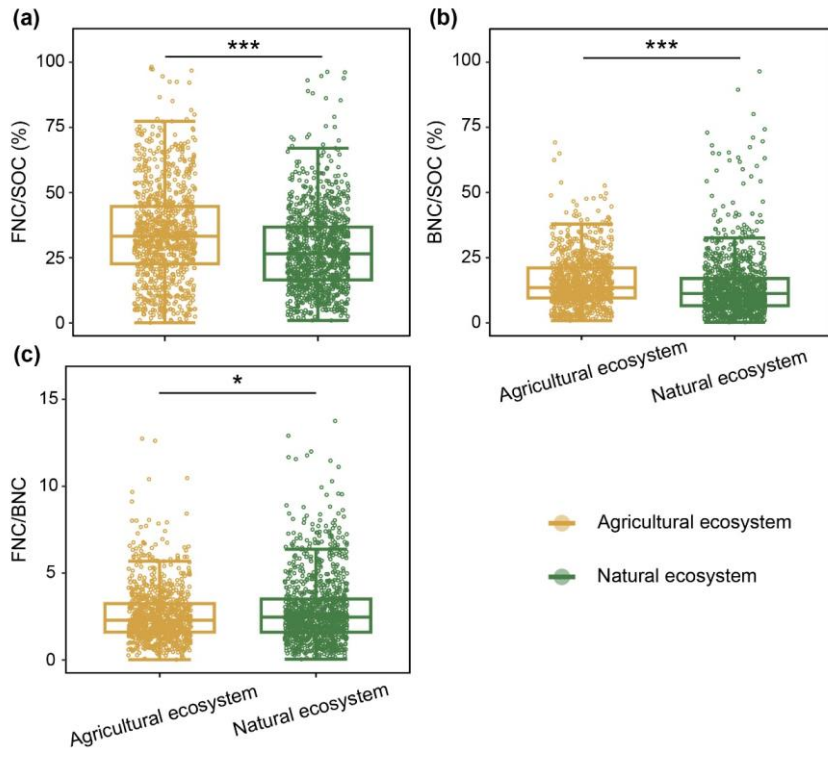
Figure 1.



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Figure 2.



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Figure 3.

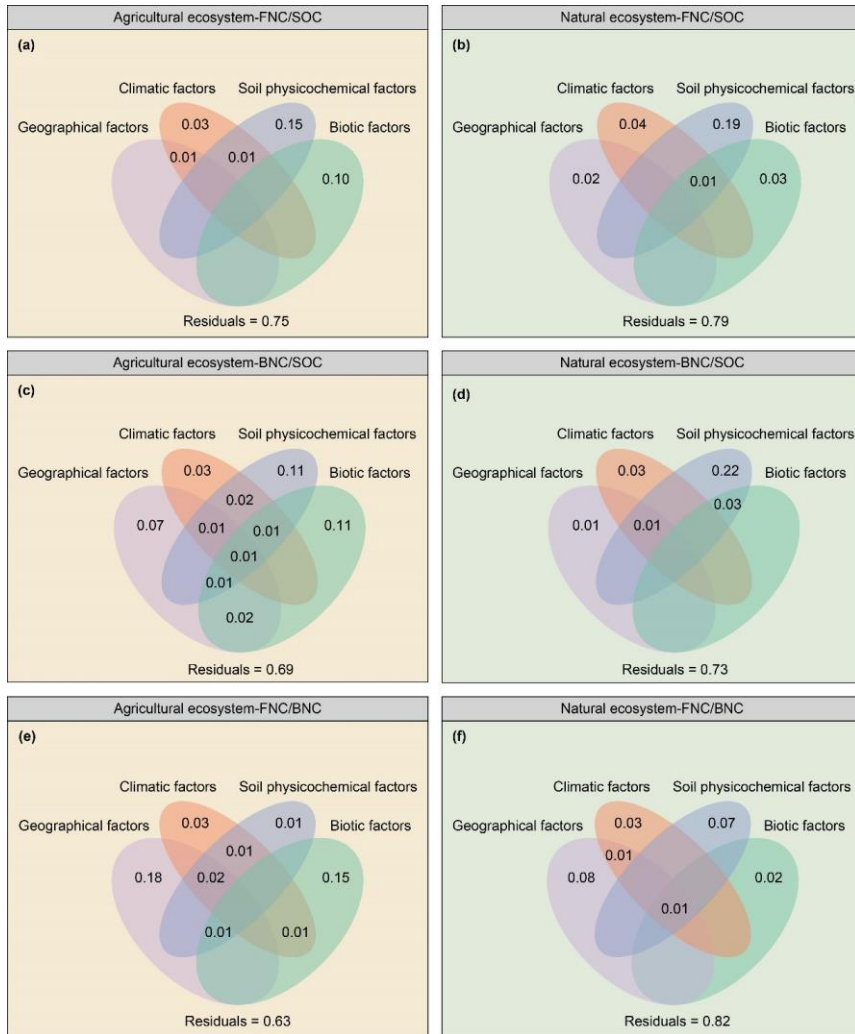


Figure 4.

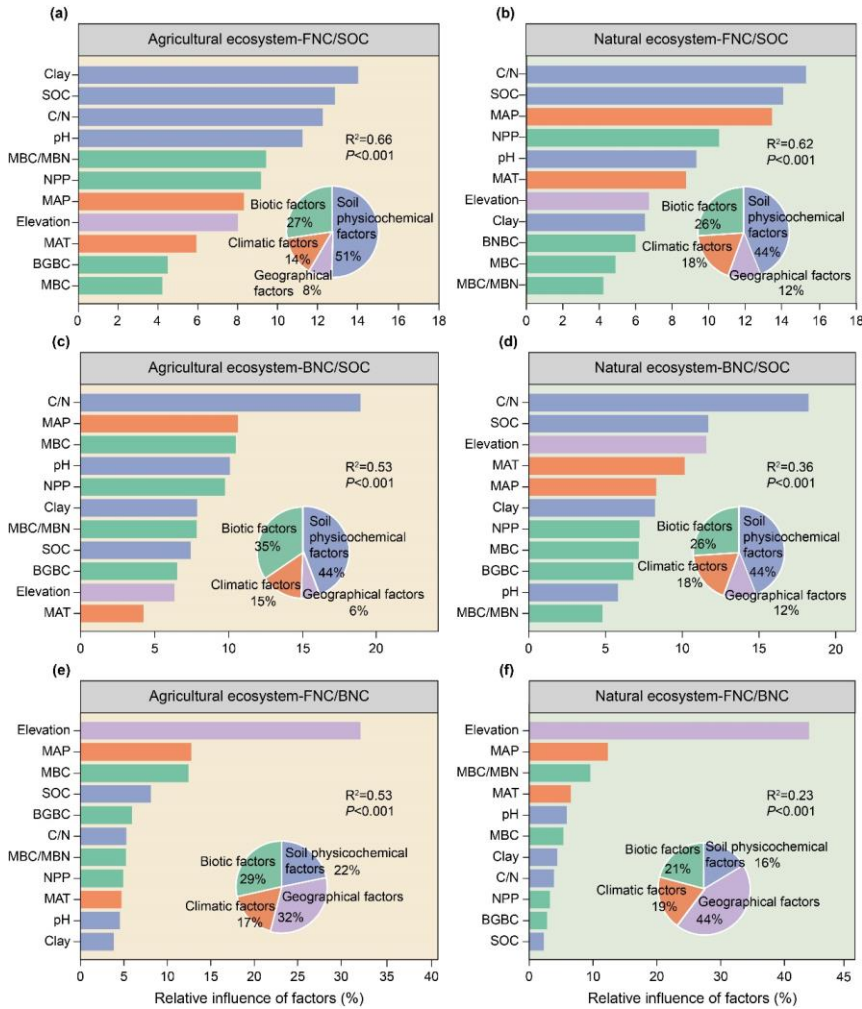


Figure 5.

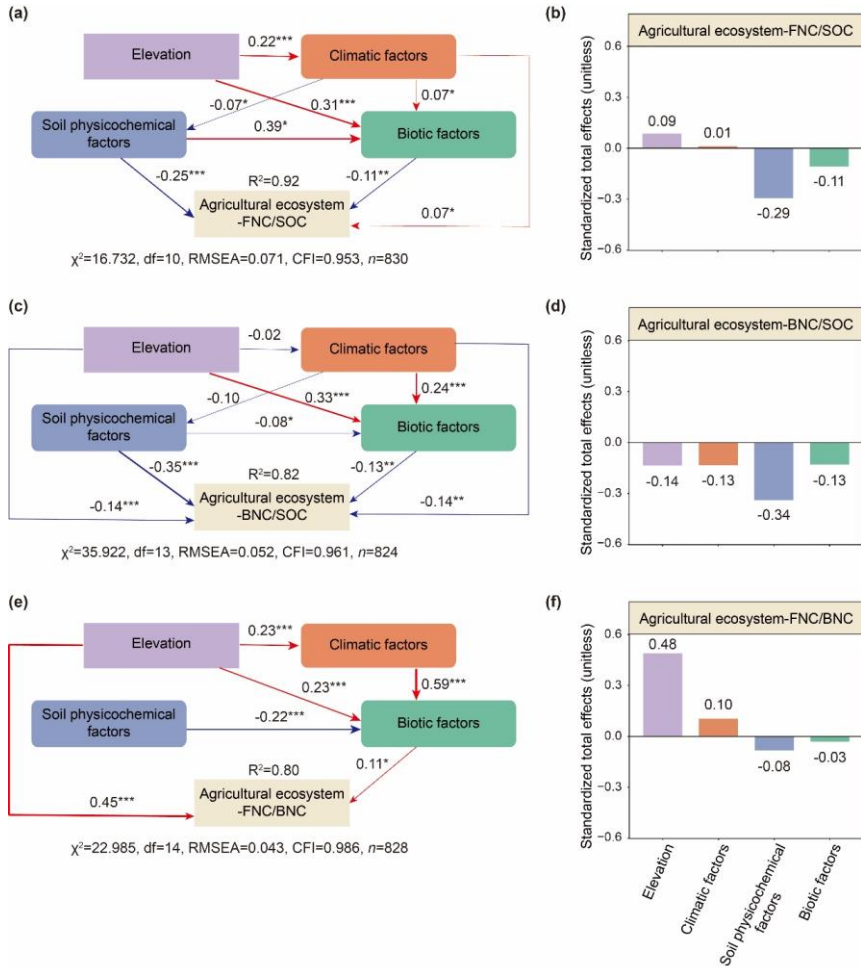


Figure 6.

