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1 Vulnerability of mineral-associated soil organic carbon to climate across

2 global drylands

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Mineral-associated organic carbon (MAOC) constitutes a major fraction of global soil 154 carbon (C), and is assumed less sensitive to climate than particulate organic C (POC) due 155 to protection by minerals. Despite its importance for long-term C storage, the response of 156 MAOC to changing climates in drylands, which cover more than 40% of the global land 157 area, remains unexplored. Here we assess topsoil organic C fractions across global 158 159 drylands using a standardized field survey in 326 plots from 25 countries and six continents. We find that soil biogeochemistry explained the majority of variation in both 160 MAOC and POC. Both C fractions decreased with increases in mean annual temperature 161 162 and reductions in precipitation, with MAOC responding similarly to POC. Therefore, our results suggest that ongoing climate warming and aridification may result in unforeseen C 163 losses across global drylands, and that the protective role of minerals may not dampen 164 165 these effects.

166

Soils in drylands—the largest set of biomes of the planet —store 646 Pg of organic C, more than 167 all living vegetation on Earth ^{1,2}. This vast soil organic C pool supports essential ecosystem 168 169 services, including food provision and water and climate regulation for more than 2.5 billion people ^{3,4}. Yet, temperature increases and precipitation reductions forecasted for many dryland 170 171 regions are expected to disrupt the balance of soil organic C, accelerating microbial 172 decomposition, reducing plant C inputs into the soil, and resulting in more CO₂ emissions to the atmosphere 5,6 . 173 The sensitivity of organic C in soils (sensu ref. 7) to temperature and precipitation at 174

timescales relevant to climate change mitigation is thought to be controlled largely by

interactions with soil minerals, which restrict the accessibility of microbial decomposers by

encapsulating and adsorbing organic matter ^{8–10}. Plant-derived materials at early stages of 177 decomposition are the main constituents of the mineral-unprotected, particulate organic C (POC) 178 fraction of soil organic matter ⁹. The POC fraction is thus directly affected by changes in plant C 179 180 inputs into the soil and is more exposed to microbial decomposition than the organic component of the mineral-associated organic C (MAOC) fraction, which has, therefore, a lower turnover rate 181 ^{11,12}. As a result, large scale meta-analyses and observational studies suggest that POC is more 182 sensitive to changes in climate, and particularly to warming, than MAOC ^{7,13–16}. Because of the 183 184 typically large ratio of soil minerals to organic matter in drylands, MAOC is expected to 185 dominate over POC, potentially driving a high persistence of soil organic C in these ecosystems ^{7,10,17}. However, no studies to date have examined the relationship of POC and MAOC with 186 187 climate across the diverse environmental gradients that characterize global drylands. 188 Investigating this relationship is particularly timely and relevant, as it would significantly reduce 189 the uncertainty surrounding the land carbon-climate feedback. Additionally, it would provide valuable insights for adapting soil carbon-related ecosystem services to ongoing climate change. 190 Here we evaluated how mean annual temperature and precipitation relates to POC and 191 MAOC contents across global drylands after accounting for major biotic (net primary 192 193 productivity, vegetation type, woody cover, plant and herbivore richness, and grazing pressure) 194 and soil biogeochemistry (clay and silt contents, pH, chemical index of alteration, exchangeable Ca, non-crystalline Al and Fe, available N and P, and microbial biomass C) factors known to 195 potentially affect soil organic C content by regulating C inputs and stabilization processes ^{5,18}. To 196 do so, we surveyed in situ 326 plots from 98 dryland ecosystems located in 25 countries from six 197 198 continents (Extended Data Fig. 1). Our survey spans the broad gradients of temperature, 199 precipitation, aridity, soil properties, vegetation types, and grazing pressures that can be found

across drylands worldwide (Extended Data Tables 1 and 2)^{19,20}. At each site, we collected 200 topsoil samples (0-7.5 cm) from areas both covered (322) and not covered (326) by perennial 201 vegetation from two to four plots located across a local gradient of extensive grazing pressure 202 203 (648 samples in total, see Methods). We subjected all samples to a size fractionation procedure to separate and quantify C content in POC and MAOC pools ^{9,21}. Using these data, we tested the 204 hypothesis that MAOC, being protected by minerals, is less sensitive than POC to increases in 205 temperature and decreases in precipitation ^{7,10,16,22}. We also hypothesize that the presence of 206 vegetation mitigates declines in soil C, particularly POC, by increasing soil C inputs. 207

208

209 MAOC dominates soil organic C and is sensitive to climate

Our results show that MAOC was the dominant soil organic C fraction in drylands globally (Fig. 1a). In particular, median MAOC content was 5.2 g C kg⁻¹ soil, equivalent to 66% of the total soil organic C content, whereas median POC content was 2.3 g C kg⁻¹ soil. This quantification falls within the range of soil organic C content (MAOC and POC) commonly found in drylands, and is relevant to improve the performance of emerging models of soil organic C formation and persistence using POC and MAOC frameworks $^{2, 23-25}$.

Contrary to our hypothesis, we found that MAOC and POC were equally sensitive to differences in climate across global drylands. In particular, both MAOC and POC were negatively associated with increasing temperature and decreasing precipitation to a similar extent, as indicated by the similar slopes of the associations (Fig. 1bc). These results were supported by the lack of a significant interaction between the effects of temperature and precipitation and the type of fraction (MAOC versus POC) tested by a linear mixed-effects model (Fig. 1d, see Methods). Based on the results from this model, we estimated that POC and

MAOC contents significantly declined with temperature at an average rate of 3.2% per °C (95% confidence interval (CI): 1.8, 4.6) and increased with precipitation at an average rate of 6.6% per 100 mm (95% CI: 0.6, 12.6).

226 Warming accelerates the microbial decomposition of soil organic matter, and precipitation reduction constrains plant production and organic matter inputs into the soil ^{5,26}. Our results are, 227 228 therefore, consistent with previously reported reductions in soil organic C content with increasing temperature and reducing precipitation across terrestrial ecosystems ^{27–29}. However, 229 and contrary to expectations of smaller sensitivity of MAOC versus POC to changes in climate 230 observed in more mesic systems ^{14,15}, our findings based on a space-fir-time substitution 231 highlight that the MAOC and POC fractions may decrease at similar rates in response to climate 232 warming and precipitation reduction across global drylands. Therefore, they suggest that the 233 234 current paradigm of mineral protection may not determine soil C persistence in dryland ecosystems ^{8,30–32}. The apparent lack of protection by minerals, which contrasts with what was 235 236 observed in mesic systems richer in organic matter, was consistent across the range of soil organic C content found in drylands (Extended Data Fig. 2). There is recent evidence that 237 MAOC is controlled not only by C stabilization in soil organo-mineral complexes, but also by 238 changes in C inputs driven by climate¹⁵. In drylands, not only precipitation reduction but also 239 warming may increase water deficit, which may decrease plant productivity⁵, C inputs into the 240 soil and C accumulation into the MAOC fraction. These is also evidence that dryland soils 241 242 maintain a high oxidative potential during dry periods, mainly through the stabilization of enzymes, which result in a rapid organic matter decomposition in wet periods ^{28,29} and may 243 244 further limit C inputs to the MAOC fraction.

245

246 Vegetation buffers soil C declines with warming

Both POC and MAOC contents were higher in soil beneath perennial vegetation (Fig. 2). We 247 248 further observed that as mean annual temperature increased, POC and MAOC contents 249 decreased, but to a lesser extent, beneath vegetation. Conversely, as mean annual precipitation increased, both contents increased in a similar manner in open areas and in areas under the 250 251 canopy of perennial vegetation (Fig. 2). These results are important because they suggest that the 252 presence of vegetation buffers, but does not fully compensate for, the negative effects of higher 253 temperature on soil C fractions. While the buffering effect of vegetation did not completely 254 counteract the vulnerability of organic C pools to increasing temperatures, our findings indicate that management practices aimed at protecting vegetation in drylands may help to maintain soil 255 256 organic C stocks in global drylands and reducing their losses in response to a changing climate.

257

258 Coupling of POC and MAOC in drylands

We found that POC and MAOC contents were strongly correlated across global drylands (r =
0.83, n = 326, P < 0.001; Fig. 3a). These results strongly suggest that both fractions remain
highly coupled in drylands despite their different levels of putative protection against
decomposition by microorganisms.

Variance partitioning of linear mixed-effects models and random forest analysis showed that the order of importance of the group of factors that explained most of the variation of POC and MAOC across global drylands was essentially the same for both organic C fractions (Fig. 3b, Extended Data Fig. 3). Soil biogeochemistry, above climate and biotic factors, was the most important predictor of both POC and MAOC contents. Both C fractions were negatively associated with soil pH and positively associated to exchangeable Ca, available N and P, and

269 microbial biomass C contents; additionally, MAOC was associated positively with clay and silt and non-crystalline Al and Fe contents (Extended Data Fig. 4). Slightly-acidic-to-neutral soils 270 generally feature higher nutrient availability and more fertility than alkaline soils ³³, which may 271 272 thus favor soil organic C accumulation in drylands through increased plant-derived C inputs and microbial activity. The prevalent role of soil fine texture and non-crystalline Al and Fe in MAOC 273 formation has been widely documented in the literature ³¹. Sorption of organic matter to mineral 274 surfaces is known to be promoted by the relatively high specific surface area and charge of clay 275 and silt, while non-crystalline Fe and Al phases are also known to form strong associations with 276 organic matter ³¹. 277

The coupling of POC and MAOC observed here for drylands may be, however, disrupted in more productive terrestrial ecosystems, where higher plant inputs may result in larger POC contents ^{13–15}. In contrast to experimental manipulation studies ¹⁴, our work addresses the vulnerability of soil C fractions using a space-for-time substitution. Further research into the pace of the climate-induced changes and the causality of the associations found in our study is thus warranted.

284

285 Concluding remarks

By using a global standardized field study and by focusing exclusively on dryland ecosystems, our work expands previous efforts to understand abiotic and biotic drivers of POC and MAOC along large geographical gradients, which have either been based on literature syntheses, which use datasets that are inherently heterogenous, or have focused on ecosystems other than drylands ¹⁶. Our study generated highly standardized field data on the POC and MAOC fractions of

dryland soils worldwide, along with their major predictors. These data significantly expandexisting global databases and can be used to refine current soil organic C models.

293 Our findings suggest that ongoing changes in climate, particularly warming, may adversely affect both unprotected and mineral-protected soil C content in drylands to a similar extent. The 294 results obtained also indicate that maintaining vegetation cover can mitigate, but not fully 295 296 counteract, the negative impacts of rising temperatures on soil organic C fractions. Our study 297 enhances our understanding of how POC and MAOC contents in soil respond to key abiotic and 298 biotic drivers, revealing that mineral protection has limited potential to sustain organic C storage 299 in dryland soils in the face of ongoing global warming. The novel insights about dryland soil C pools and their sensitivity provided here could facilitate much-needed advances in our model 300 representation of dryland ecosystems and their response to climate change. 301

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331 Author contributions

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344	analysis, after discussion, suggestions, and contributions from F.T.M., E.M.J., M.D.B., N.G.,
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346	manuscript draft, with contributions from F.T.M., E.M.J., and M.D.B. All authors discussed the
347	results and contributed to editing the manuscript.
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351	
352	Figure captions
353	
354	Fig. 1 Distribution of soil organic carbon (C) contents in particulate organic C (POC) and
355	mineral-associated organic C (MAOC) fractions and their relationships with climate in
356	global drylands. a, Boxplot of POC and MAOC contents. Box, 1 st , and 3 rd quartiles; central
357	horizontal line, median; upper vertical line end, largest value smaller than 1.5 times the
358	interquartile range; lower vertical line, smallest value larger than 1.5 times the interquartile range
359	(n = 326 plots). b-c , Relationships between POC and MAOC contents and mean annual
360	temperature (MAT, b) and precipitation (MAP, c). Lines and shading represent linear regressions
361	and 95% confidence intervals. d, Summary of a linear mixed-effects model, controlling for biotic
362	factors and soil biogeochemistry (see Methods). The panel shows coefficients (circles) and 95%

363 confidence intervals (CI, bars) for main and interaction effects of C fraction type (binary

variable, either POC or MAOC) and climate (MAT and MAP) on POC and MAOC contents.

365 The variance explained (R^2) by the fixed and random effects relative to the total variance was

366 77% and 12%, respectively (n = 634 POC and MAOC observations). Carbon fraction contents

367 were natural-logarithm transformed, and all the predictors were standardized. The positive

368 coefficient of C fraction type (MAOC vs. POC) indicate that MAOC contents are significantly

369 greater than POC contents (P < 0.001). For the observed negative association of MAT and

positive association of MAP with C content ($P \le 0.001$ and P = 0.039 respectively), negative

371 coefficients for the interaction of C fraction type with MAT and MAP indicate that increasing

MAT has a stronger negative effect on MAOC than on POC contents (P = 0.053), while

decreasing MAP has a stronger negative effect on POC than on MAOC (P = 0.181).

Fig. 2 | Relationships between climate and particulate organic C (POC) and mineral-

associated organic C (MAOC) contents in soils under the canopy of the dominant perennial

vegetation (V) and in open areas (O) across global drylands. a-d, Relationships between POC

and mean annual temperature (MAT, **a**) and precipitation (MAP, **c**), and between MAOC and

378 MAT (**b**) and MAP (**d**) in both O and V microsites. Lines and shading represent linear

regressions and 95% confidence intervals (n = 326 and 322 for O and V, respectively). e,

380 Coefficients (dots) and 95% confidence intervals (bars) of linear mixed-effects model illustrating

the fixed main and interaction effects of MAT, MAP, and the presence of vegetation cover (V vs.

382 O) on POC and MAOC contents (n = 648 V and O areas). The variance explained (R^2) by the

fixed and random effects relative to the total variance was 30% and 55%, respectively, for POC,

and 32% and 61%, respectively, for MAOC.

Fig. 3 | Coupling and drivers of particulate organic C (POC) and mineral-associated

organic C (MAOC) in global drylands. a, Relationship between POC and MAOC contents.

387 Dots represent individual dryland plots, with the colors of the dots illustrating their aridity (1 -

annual precipitation/potential evapotranspiration) values. The line and shading represent the

fitted linear regression and 95% confidence interval, respectively. **b**, Variance explained (R^2) by

- 390 linear mixed-effects models for POC and MAOC contents partitioned into the fraction
- attributable to unique and shared among groups of drivers (climate: mean annual temperature
- and mean annual precipitation; biotic factors: net primary productivity, type of vegetation,
- 393 woody cover, plant richness, grazing pressure, and herbivore richness; and soil biogeochemistry:

394	clay	y and silt, pH, chemical index of alteration, exchangeable Ca, non-crystalline Al and Fe,			
395	available N and P, and microbial biomass carbon). The variance explained (R^2) by the fixed and				
396	random effects relative to the total variance was 69% and 20% for POC ($n = 317$) and 84% and				
397	119	% for MAOC ($n = 317$), respectively.			
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478 Methods

Global field survey and soil sampling. Fieldwork was conducted from January 2016 to
September 2019. A total of 326 plots distributed across 98 study sites in 25 countries from all
continents except Antarctica (Algeria, Argentina, Australia, Botswana, Brazil, Canada, Chile,
China, Ecuador, Hungary, Iran, Israel, Kazakhstan, Kenya, Mexico, Mongolia, Namibia, Niger,
Palestine, Peru, Portugal, South Africa, Spain, Tunisia, and USA) and encompassing the wide
range of vegetation, soil, climate, and grazing pressure levels found in drylands worldwide were
surveyed using a common and standardized protocol ^{19,20}.

486 At each site, we gathered field data within multiple 45 m x 45 m plots situated along a gradient of grazing pressure, encompassing high (n = 98), medium (n = 97), and low (n = 88)487 pressure levels, as well as ungrazed areas (n = 43). To establish the grazing gradients, in 90 out 488 of the 98 sites surveyed, we strategically positioned these plots at varying distances from 489 artificial watering points, which are usually created in drylands to supply introduced livestock 490 with permanent water sources ³⁴. The closer the plot to the permanent water source, the more 491 intense the grazing ^{34,35}. In the remaining eight sites, local variations in grazing pressure 492 493 gradients were ascertained by observing different paddocks featuring varying grazing intensities. See ref.²⁰ for additional details on the characterization and validation of the local grazing 494 pressure gradients established. 495

A portable Global Positioning System was used to record the coordinates and elevation of
each plot, which were standardized to the WGS84 ellipsoid for visualization and analyses.
During the dry season at each site, four soil cores (145 cm³) from 0 to 7.5-cm depth (topsoil)
were collected from five 50 × 50-cm quadrats randomly placed in areas under the canopy of the
dominant perennial vegetation and five placed in open areas not covered by perennial vegetation.

The soil cores were homogenized and composited to form a sample representative of the soil under the dominant vegetation and a sample representative of the soil in open areas within each plot. The soil samples were passed through a 2-mm sieve. A portion of each soil sample was airdried and used for organic matter fractionation and texture and pH analysis, and another portion was stored at -20 °C and used for microbial biomass C analysis. A portion of the air-dried soil samples was ground with a ball mill for additional chemical analysis.

Soil organic carbon fractionation and quantification. All the soil samples, a total of 648 (326 507 from open areas and 322 from under the canopy of the dominant vegetation), were subjected to a 508 size fractionation method ^{21,36} to separate the POC (not protected by minerals from microbial 509 decomposition) and MAOC (protected by minerals) fractions. Aggregates were dispersed by 510 adding 30 mL of sodium hexametaphosphate (5 g L^{-1}) to 10 g of soil and shaking with an 511 512 overhead shaker for 18 h. After dispersion, the mixture was thoroughly rinsed through a 53-um sieve, to separate the POC (> 53 μ m) and MAOC (< 53 μ m) fractions using an automated wet 513 sieving system. The isolated fractions were oven-dried at 60 °C, weighed, and ground with a ball 514 mill. The whole soil samples and the POC and MAOC fractions were analyzed for organic C 515 516 contents by dry combustion and gas chromatography using a ThermoFlash 2000 NC Soil Analyzer (Thermo Fisher Scientific, MA) after removing carbonates by acid fumigation ³⁷. 517 **Climate data.** Mean annual temperature and mean annual precipitation data were obtained from 518 WorldClim 2.0 38 a high resolution (30 arc seconds or ~ 1 km at the equator) database based on a 519 520 large number of climate observations and topographical data for the 1970-2000 period. Aridity index (ratio of average annual precipitation to potential evapotranspiration) data were obtained 521 from the Global Aridity Index and Potential Evapotranspiration Climate Database v3³⁹. Aridity 522 523 was calculated as 1 - aridity index.

524 Vegetation and herbivore richness survey. Each plot was classified as grassland, shrubland, or forest by identifying the dominant type of vegetation. Net primary productivity (NPP) was 525 estimated using the mean annual Normalized Difference Vegetation Index (NDVI) averaged 526 527 monthly values between 1999 and 2019 at a resolution of 30 m from Landsat 7 Enhanced Thematic Mapper Plus (ETM+)⁴⁰. The cover of perennial vascular plants (plant cover) was 528 measured along four parallel 45-m transects separated by 10 m and oriented downslope during 529 the peak of the growing season using the line-intercept method ^{19,41,42}. Woody cover was 530 measured in 25 contiguous quadrats $(1.5 \text{ m} \times 1.5 \text{ m})$ placed in each transect (100 quadrats per 531 532 plot). Plant richness was the total number of unique perennial species found along the quadrats and transects surveyed. The richness of herbivores was quantified at each plot using dung data 533 collected systematically in situ along the four 45-m transects established as described in ref.²⁰. 534 535 **Soil analyses.** All the bulk soil samples were analyzed as follows. Clay and silt contents were determined by sieving and sedimentation ⁴³. Soil pH was measured in a water suspension at a 536 soil-to-water ratio of 1:2.5⁴⁴. The chemical index of alteration, which is an indicator of the 537 degree of weathering, was calculated as the molecular proportion of Al₂O₃ versus Al₂O₃ + CaO + 538 $Na_2O + K_2O^{45}$, using total Al, Ca, Na, and K contents and after correcting Ca for soils with 539 carbonates ¹⁸; total Al, Ca, Na and K contents were determined by inductively coupled plasma 540 atomic emission spectroscopy (ICP-AES) after digestion in nitric and perchloric acids ^{44,46}. 541 Exchangeable Ca content was determined by ICP-AES after extraction with ammonium acetate 542 at pH 7.0^{44,47}. Non-crystalline Fe and Al contents were determined by ICP-AES after extraction 543 with acid ammonium oxalate ⁴⁸. Available N (ammonium and nitrate) content was determined by 544 extraction with 0.5 M K₂SO₄ and the indophenol blue method using a microplate reader ⁴⁹. 545

Available P content was determined by the Olsen method ⁵⁰. Microbial biomass C was 546 determined by substrate-induced respiration ⁵¹ using an automated microrespirometer ⁵². 547 Statistical analyses. We compared the content of MAOC with that of POC in global dryland 548 549 soils controlling for confounding factors, and tested the hypothesis that the effects of climate (mean annual temperature and precipitation) on POC and MAOC contents depends on (interacts 550 551 with) the C fraction type. For these analyses, we aggregated soil data for open and vegetation covered areas by plot using plant cover area as a weighting factor, and fitted a linear mixed-552 553 effects model on the response of C content with C fraction type as a binary categorical predictor 554 (either MAOC or POC). In the fixed-effects term of the model, we also included mean annual temperature, mean annual precipitation, and the interactions of mean annual temperature and 555 mean annual precipitation with C fraction type, as well as key biotic (net primary productivity, 556 557 type of vegetation, woody cover, plant richness, grazing pressure, and herbivore richness) and soil biogeochemical (clay and silt, pH, chemical index of alteration, exchangeable Ca, non-558 559 crystalline Al and Fe, available N and P, and microbial biomass C) covariates to control for confounding factors. In the random term of the model, we incorporated an intercept structure 560 with plot nested within site as a categorical variable to account for the lack of independence in 561 562 the residuals due to the paired POC and MAOC separation and the plot sampling design. We checked whether the fit of this linear mixed-effects model improved by including quadratic terms 563 564 of mean annual temperature, mean annual precipitation, and both mean annual temperature and 565 precipitation, using the Akaike information criterion (AIC) and likelihood ratio tests. None of the quadratic models tested was a significantly better fit to the data (χ^2 (1) < 1.0, P > 0.3) than the 566 567 linear model (lowest AIC).

568 To examine separately the variance of POC and MAOC contents explained by the groups of predictors (climate: mean annual temperature and mean annual precipitation; biotic factors: net 569 primary productivity, type of vegetation, woody cover, plant richness, grazing pressure, and 570 herbivore richness; soil biogeochemistry: clay and silt, pH, chemical index of alteration, 571 exchangeable Ca, non-crystalline Al and Fe, available N and P, and microbial biomass C), we 572 573 built two linear mixed-effects models (one for POC and another one for MAOC) with site as a random categorical variable. These two separate models were used to assess the importance of 574 the different groups of predictors in explaining either POC or MAOC, and not to test statistically 575 576 for differences in the size of the effects of the predictors between POC and MAOC. To support the linear mixed-effects models, we tested the importance of the same groups of predictors of 577 POC and MAOC using random forest regression modeling ⁵³. In particular, we built two random 578 579 forest models, one for POC and one for MAOC, combining 500 trees, and quantified the importance of each predictor by computing the increase in mean squared error across trees when 580 the predictor was permuted. 581

We tested whether the presence of vegetation cover interacted with the effects of temperature and precipitation also by linear mixed-effects modeling. For this purpose, we built two linear mixed-effects models, one for POC content and another one for MAOC content in areas under the canopy of the dominant perennial vegetation and open areas, with vegetation cover as a binary predictor and plot nested within site in the random term.

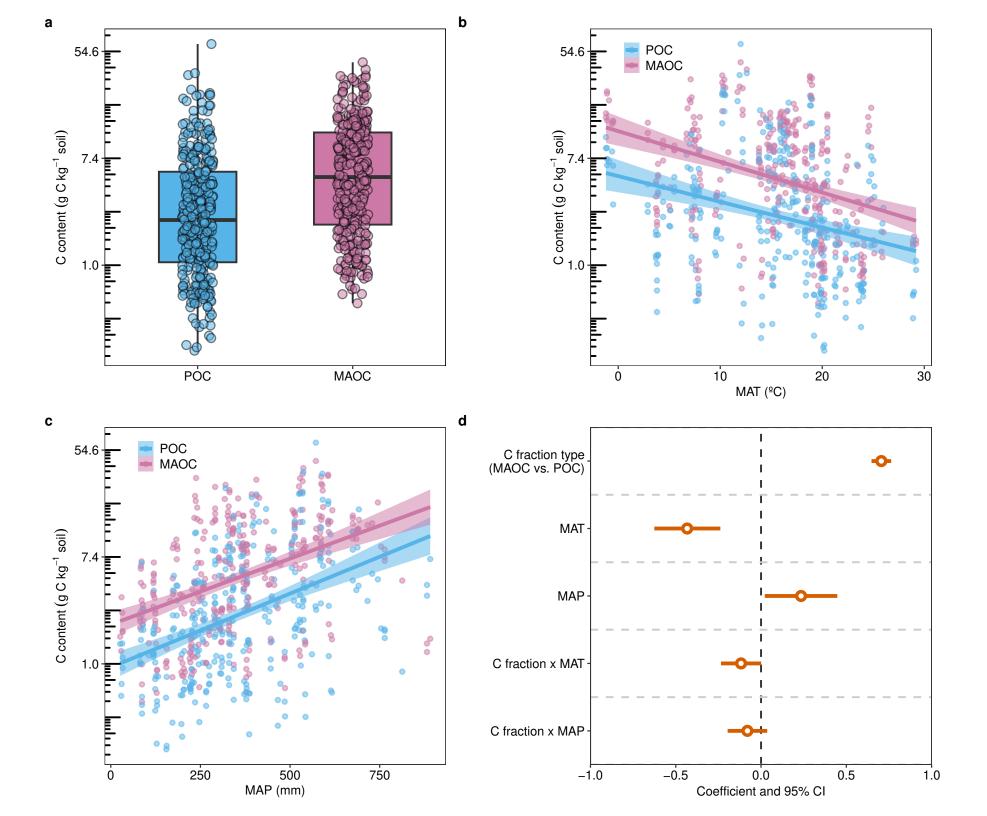
587 For all the linear mixed-effects models, POC, MAOC, exchangeable Ca, non-crystalline Al 588 and Fe, available N and P, and microbial biomass C were natural-logarithm transformed to 589 reduce the skewness of the data. To compare effect sizes, all the numeric predictors were 590 standardized by subtracting the mean and dividing by two standard deviations, and the binary

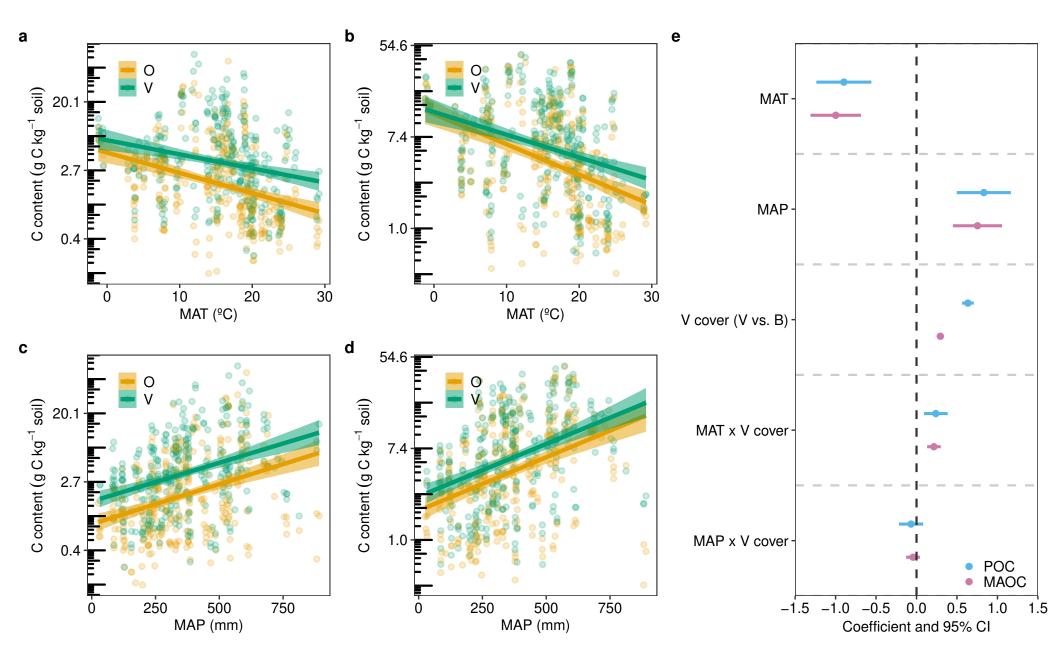
591	variables (C fraction type and vegetated vs. open areas) were rescaled to -0.5 and 0.5 54 . The				
592	coefficients of the models were estimated by the restricted maximum likelihood approach, 95%				
593	confidence intervals were calculated, and P-values were computed based on Satterthwaite				
594	approximation ⁵⁵ . The validity of the assumptions of normality, homoscedasticity and linearity				
595	were examined using residual plots. The generalized variance inflation factors (GVIFs) were				
596	computed to check for multicollinearity among predictors (GVIF values were less than 3 in all				
597	cases, suggesting that multicollinearity was low ⁵⁶). All statistical analyses were performed using				
598	R ⁵⁷ and the R packages arm ⁵⁸ , ggplot2 ⁵⁹ , lme4 ⁶⁰ , lmerTest ⁵⁵ , partR2 ⁶¹ , patchwork ⁶² ,				
599	rnaturalearth ⁶³ , randomForest ⁶⁴ , sf ⁶⁵ , terra ⁶⁶ , and viridis ⁶⁷ .				
600					
601	Data availability				
602	The data associated with this study are publicly available in				
603	https://figshare.com/s/8aeac2300650181f2c86 (https://doi.org/10.6084/m9.figshare.24678891) ⁶⁸ .				
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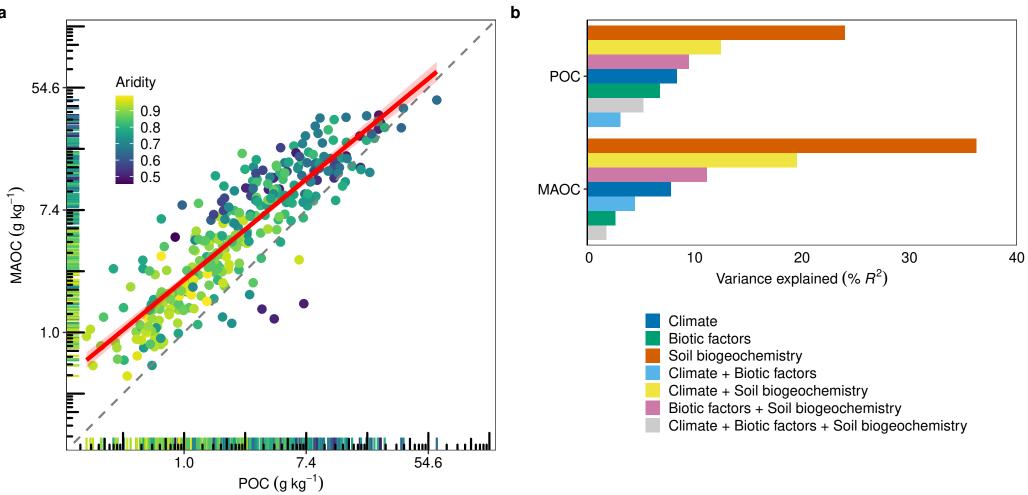
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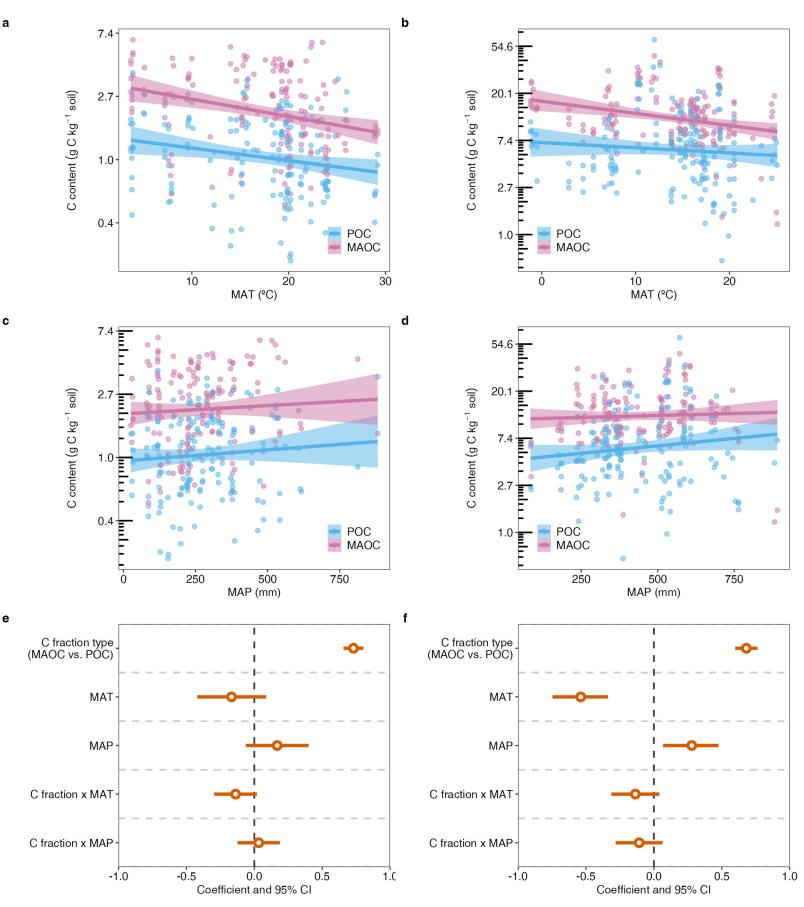
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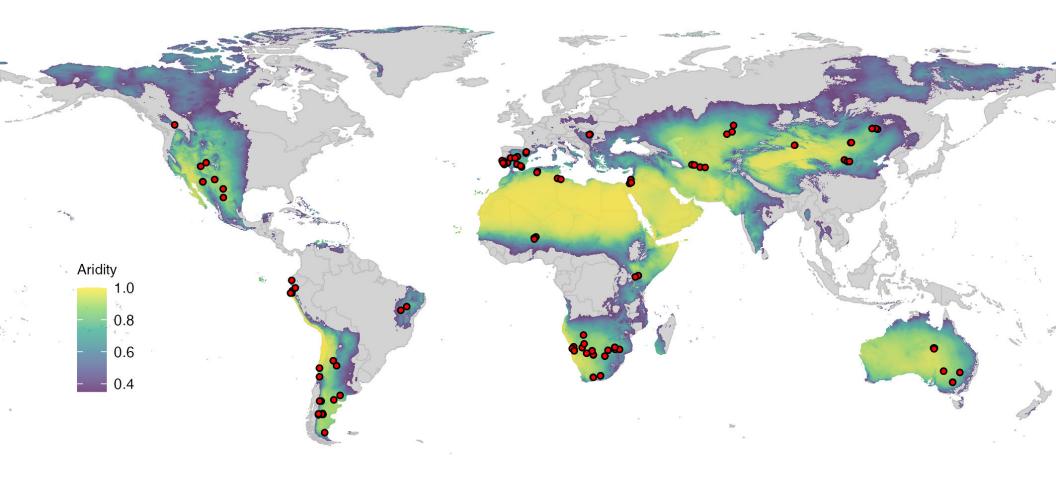
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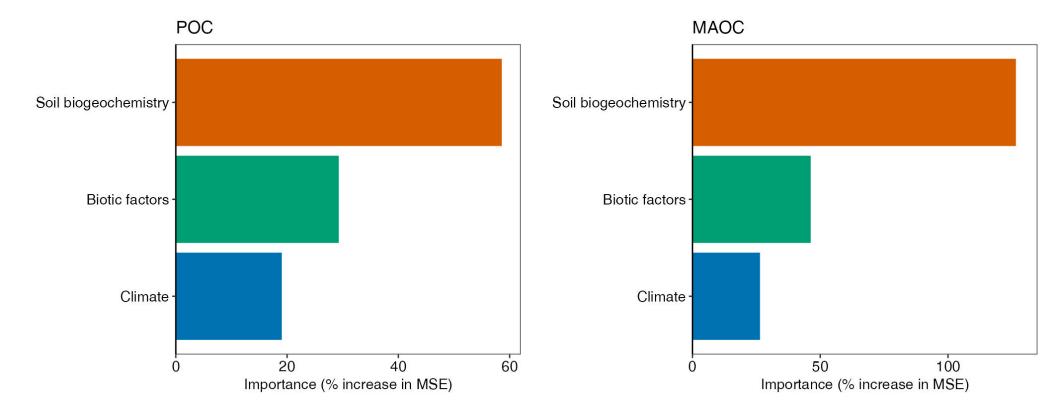


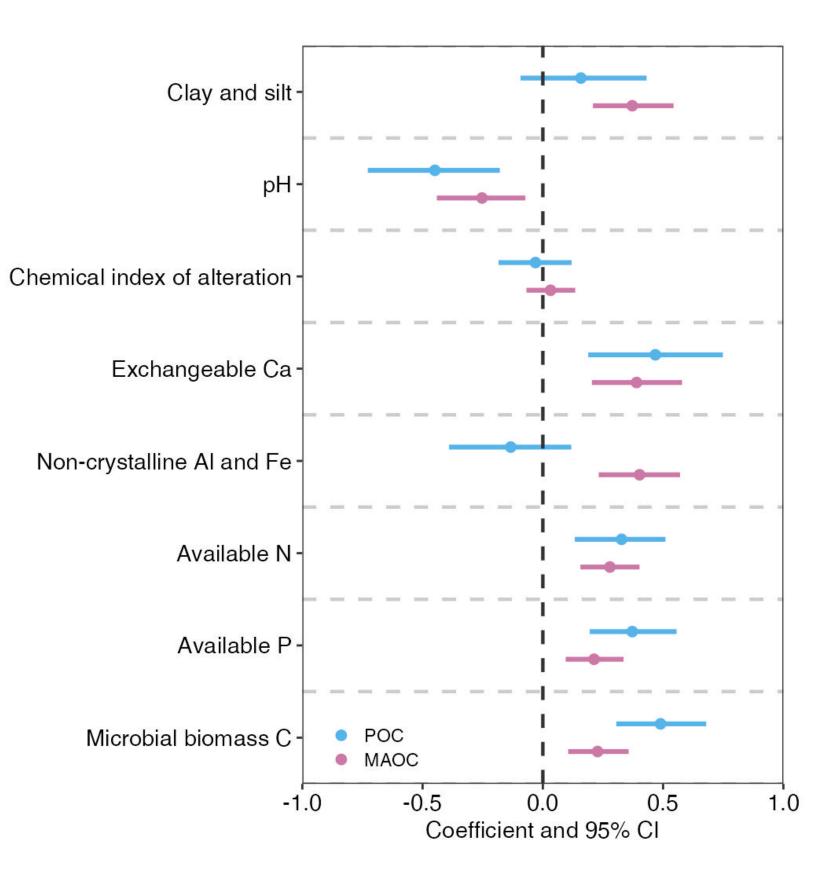












Variable	n	Min	Q1	Median	Mean	Q3	Max
MAT (°C)	326	-1.2	10.4	16.6	15.5	19.9	29.2
MAP (mm)	326	26	233	332	357	505	891
Net primary productivity (NDVI, unitless)	326	0.06	0.13	0.17	0.19	0.25	0.43
Woody cover (%)	326	0	15	46	48	83	100
Plant richness (number of species)	326	0	8	16	19	26	57
Herbivore richness (number of species)	326	0	1	2	2	3	6
Clay and silt (g kg ⁻¹)	326	10	120	271	325	512	870
рН	326	4.5	6.1	7.0	6.9	7.8	9.9
Chemical index of alteration (%)	326	42	74	81	79	87	97
Exchangeable Ca (mg kg ⁻¹)	321	39	843	1730	3394	3443	42446
Non-crystalline AI and Fe (mg kg ⁻¹)	326	28	475	932	1357	1620	9889
Available N (mg kg ⁻¹)	326	1	8	14	21	26	143
Available P (mg kg ⁻¹)	323	0.1	5.5	11.5	13.6	17.8	87.6
Microbial biomass C (mg kg ⁻¹)	326	16	101	186	245	331	1065

Variable	Category	Number of observations
Vegetation type	Grassland	94
	Shrubland	160
	Forest	72
Grazing pressure	Zero	43
	Low	88
	Medium	97
	High	98