Climate control on terrestrial biospheric carbon turnover

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Significance

Terrestrial organic-carbon reservoirs (vegetation, soils) currently consume more than a third of anthropogenic carbon emitted to the atmosphere, but the response of this “terrestrial sink” to future climate change is widely debated. Rivers export organic carbon sourced over their watersheds, offering an opportunity to assess controls on land carbon cycling on broad spatial scales. Using radiocarbon ages of biomolecular tracer compounds exported by rivers, we show that temperature and precipitation exert primary controls on biospheric-carbon turnover within river basins. These findings reveal large-scale climate control on soil carbon stocks, and they provide a framework to quantify responses of terrestrial organic-carbon reservoirs to past and future change.

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Terrarrestrial biospheric carbon residing in vegetation and soils may moderate or exacerbate ongoing buildup of atmospheric greenhouse gases on timescales that are of direct relevance to humankind (1). Much current debate surrounds the response and potential contributions of terrestrial ecosystems to climate change, with large uncertainties concerning the magnitude—and even the sign—of change in response to different environmental forcing factors such as temperature and hydrology (2, 3). Because of their large organic carbon (OC) stocks and potential to stabilize carbon on a range of timescales, soils are thought to regulate overall terrestrial ecosystem carbon storage (4, 5). Globally, soil carbon turnover time (τs) (i.e., the ratio of soil carbon stock to input flux) is estimated via remote sensing approaches (4, 5) and Earth-system models (3) that are calibrated using numerous observational and experimental studies investigating controls on soil OC turnover in a range of ecosystems and soil types (6, 7). However, findings from such studies are often relevant only to a specific experiment, plot, or environment, thus hindering extrapolation and regional validation of remote sensing and model products (8, 9).

One major reason for this limitation is our lack of constraints regarding the importance of erosion and lateral transport, despite a growing realization that these processes are pervasive on diverse landscapes (10) and link terrestrial and aquatic components...
of the carbon cycle (11). Investigations of lateral transport of terrestrial biospheric carbon have thus far focused on small spatial scales (e.g., hillslopes), which do not fully encompass heterogeneous landscape mosaics (10, 12, 13) and cannot be readily extrapolated to assess processes relevant at the ecosystem or biome scale. In particular, there is a paucity of information on how carbon turnover observed at individual plots relates to landscape- and basin-scale biospheric carbon dynamics. Depending on the mode of carbon turnover, mobilization, and transport, lateral processes can either export freshly synthesized carbon (e.g., via surface runoff) or exhume carbon stocks sequestered in deeper soils and wetlands (14, 15); the relative importance of these processes likely exerts a strong influence on carbon stocks and dynamics but is currently underrepresented in large-scale models. Given that climate change, as well as direct anthropogenic perturbations (e.g., land-use practices), may potentially modify and amplify such carbon fluxes and trajectories (3), establishing the underlying drivers and pacing of carbon cycling on appropriate spatial and temporal scales is of key importance.

Insight into the factors that control ecosystem-scale carbon turnover times can be determined using the $^{14}$C activity (reported as age in $^{14}$C years) of OC laterally exported by rivers. Rivers integrate processes within their watersheds, thus enabling investigation of biogeochemical processes at the basin scale, facilitating observational extrapolation, and linking terrestrial and marine realms. Fluvial systems form the major conduit that transfers OC from the continents to the ocean, exporting a combined $\sim 4.5 \times 10^{14}$ g of dissolved organic carbon (DOC) and particulate organic carbon (POC) annually (16, 17). The majority of this OC exported by rivers is in dissolved form, but much of this is rapidly mineralized (16). Although POC is also subject to extensive degradation (16) and riverine POC export is estimated to account for less than 0.2% of net primary production (NPP, 17), the export of terrestrial biospheric POC and its subsequent burial in marine sediments is important in modulating atmospheric $^{14}$CO$_2$ on a range of timescales (18) and provides some of the most continuous and long-term records of past climate and carbon-cycle dynamics on the continents.

Prior studies have examined the nature and magnitude of carbon transfer via rivers to the ocean (ref. 17 and references therein) and have shown that soil OC represents a dominant component of the terrestrial POC exported by many fluvial systems (19, 20). Strong contrasts in POC yield (i.e., carbon flux per unit catchment area) and composition relate to geomorphic and climatic factors influencing mobilization and retention of OC within drainage basins, as well as the proportions and fluxes of biospheric versus rock-derived (“petrogenic”) carbon inputs (14, 21). Bulk OC radiocarbon measurements in both particulate and dissolved phases reveal a wide variety of $^{14}$C ages (22); however, interpretations in terms of biospheric-carbon dynamics are confounded by diverse OC contributions (e.g., petrogenic OC; ref. 17 and references therein) and secondary overprinting (within-river autotrophy and heterotrophy) (22). Moreover, for most modern river systems, anthropogenic activities influence $^{14}$C ages through the introduction of organic contaminants that may be either relatively modern (e.g., domestic sewage) or fossil (e.g., petroleum or petrochemical contamination) in age (23).

These interferences can be obviated by determining ages of organic compounds unique to vascular land-plant biomass (24). To explore controls on the age of terrestrial biospheric carbon exported from river basins, here we compile previously reported ($n = 95$) and report additional ($n = 28$) $^{14}$C age measurements from source-specific “biomarker” tracer compounds measured on 36 fluvial systems representing diverse watersheds and collectively accounting for $\sim 42, 29,$ and $20\%$ of the global riverine water, sediment, and POC discharge, respectively (Fig. 1 and SI Appendix, Fig. S1 and Tables S1 and S2) (17). We focus on two well-established classes of terrestrial higher plant biomarkers—plant-wax lipids (25) and lignin-derived phenols (26). Because of their hydrophobic nature, plant waxes reside in the particulate phase—particularly via association with mineral surfaces (27)—and persist in soils and downstream environments (24). The abundances, distributions, and stable isotopic compositions ($^{13}$C/$^{12}$C, reported as δ$^{13}$C) of these compounds preserved in sedimentary and soil sequences carry information on past vegetation inputs, plant productivity, and environmental conditions (24). We specifically analyze n-alkanoic acids (“fatty” acids [FAs]) since n-alkanes can be influenced by contributions from bedrock- or fossil-fuel-derived sources that impact corresponding $^{14}$C ages (19). Lignin imparts structural support for the plant; phenolic monomers liberated by chemical hydrolysis of this biopolymer and its corresponding residues in soils and sediments carry information on plant and tissue type and extent of degradation (ref. 26 and references therein). Assessments of lignin stability and turnover in soils vary (7); however, like plant waxes, lignin signatures are present in fluvial sediments and deposits (15). We therefore treat measured $^{14}$C ages of these two biomarker classes
as independent yet complementary estimates of the mean age of fluvially exported biospheric OC.

**Results and Discussion**

**Relationships with Basin Properties.** We examined $^{14}$C and, in selected cases, $\delta^{13}$C variations in bulk OC, lignin phenols, and fatty acids from fluvial sediment samples (Materials and Methods and SI Appendix, Tables S1 and S2). Bulk OC $^{14}$C ages and $\delta^{13}$C values in the investigated rivers encompass much of the variability that has been observed in riverine POC worldwide (SI Appendix, Fig. S2) (22), indicating that our sample set accurately captures global trends. Bulk POC ages range from modern to almost $11,000^{14}$C y in our sample set ($n = 43$), whereas biomarker ages range from above modern (reflecting incorporation of nuclear-bomb–derived $^{14}$C) to more than $10,000^{14}$C y for fatty acids ($n = 44$) and from above modern to almost $5,000^{14}$C y for lignin phenols ($n = 16$) (SI Appendix, Tables S1 and S2). Observed age offsets between fatty acids and lignin phenols collected from the same river basins may reflect differences in their degree of mineral stabilization (28) and/or pathways of mobilization (15).

To identify key parameters governing bulk OC and biomarker $^{14}$C ages and $\delta^{13}$C values, we performed an ordinary least squares (OLS) multivariate linear regression (MLR) using a suite of climatic, geomorphic, and anthropogenic properties as control variables (Materials and Methods; SI Appendix, Table S3). This approach implicitly treats results calculated using modern values as representative of control variable values over the timescales of biomarker $^{14}$C ages (i.e., centuries to millennia). Because this assumption does not strictly hold for all control variables (e.g., anthropogenic land use), correlations calculated herein may deviate from steady-state results. Many control variables are spatially resolved and must be averaged across each river basin. However, it has been shown that fluvially exported POC and, in particular, biomarker isotope signatures exhibit a downstream bias and do not represent a uniformly integrated basin signal (29). We therefore weight spatial values by a factor—the $e$-folding distance—that decays exponentially with upstream flow distance from each sampling location (Materials and Methods). We choose as the optimal $e$-folding distance the value that maximizes the fraction of total bulk OC and biomarker isotope variance that is explained by our MLR analysis (average adjusted $r^2 = 0.72$; SI Appendix, Fig. S3). This optimal value of $\sim 500$ km agrees with previous estimates of biomarker spatial integration in large river basins (29, 30).

**Spatially resolved control variables** are thus weighted by upstream flow distance with an $e$-folding distance of $500$ km when calculating catchment averages (SI Appendix, Table S3). For all bulk and biomarker isotope measurements, MLR-predicted values show no bias (measured versus predicted slopes are always statistically identical to unity) and explain the majority of observed sample variance (measured versus predicted adjusted $r^2$ always $\geq 0.35$ and typically $\geq 0.64$; SI Appendix, Fig. S4), indicating the robustness of the chosen control variables.

Catchment-weighted geomorphic characteristics such as basin area, relief (mean basin slope), and relative floodplain extent showed no significant relationship with biomarker age (SI Appendix, Table S4). Instead, FA and lignin-phenol $^{14}$C ages are significantly correlated with climate variables—chiefly, catchment-weighted mean annual temperature (MAT) and precipitation (MAP) (Fig. 2A and SI Appendix, Table S4). Globally, biomarker $^{14}$C ages decrease (become younger) with increasing MAT and MAP; this phenomenon manifests as a relationship between biomarker $^{14}$C age and latitude because of the strong covariation of the latter with climate (Fig. 1C). A significant, albeit weaker, positive correlation is also observed between biomarker $^{14}$C ages and the fraction of catchment area that is impacted by anthropogenically perturbed land use (i.e., agriculture and urbanization; Fig. 2A and SI Appendix, Table S4), suggesting such perturbations might mobilize old OC that would otherwise be stable under natural conditions. However, directly ascribing land-use–change impacts on biomarker $^{14}$C ages is challenging since the extent of anthropogenically perturbed area additionally exhibits strong covariation with climate variables, particularly MAT and MAP. Finally, lignin-phenol and, to a lesser extent, plant-wax fatty-acid $^{14}$C ages display a significant negative correlation with runoff (Fig. 2). While other variables—for example, soil properties such as clay content (31, 32)—may contribute to these relationships, their strength implies that climate constitutes the dominant direct or indirect driver of biospheric-carbon ages. Bulk OC $^{14}$C ages display systematically weaker correlations with the set of tested control variables (e.g., MAT and MAP), as well as with latitude (SI Appendix, Fig. S5), reflecting the influence of additional OC sources (e.g., petrogenic carbon, in situ aquatic productivity) on bulk OC isotopic signatures.

To further assess underlying natural variables controlling bulk POC and biomarker isotope compositions, we additionally performed a redundancy analysis (RDA) (Materials and Methods). The results show that two orthogonal axes explain a combined
57% of the total sample variance (Fig. 2B and SI Appendix, Table S5). The first axis (RDA1) loads closely with catchment-weighted MAT, MAP, and associated highly correlated variables (e.g., permafrost cover, land use), whereas the second axis (RDA2) loads closely with precipitation seasonality (coefficient of variability). All $^{14}$C metrics, particularly fatty-acid $^{13}$C age, load almost exclusively onto RDA1, indicating that they are controlled most strongly by MAT and MAP (and/or all highly correlated variables). In contrast, $^{13}$C values of fatty acids and bulk POC, broadly reflecting biome structure within the basin (25), load almost exclusively on RDA2, revealing that they are controlled most strongly by precipitation seasonality (and/or all highly correlated variables). Specifically, lower $^{13}$C values correspond to lower seasonal variation in precipitation, as observed in other regional studies (e.g., 33).

**Relationships with $\tau_{\text{soil}}$ and Mean Age.** Plant-wax fatty-acid and, to a lesser extent, lignin-phenol $^{14}$C ages display positive relationships with catchment-weighted $\tau_{\text{soil}}$ estimated from remote-sensing-derived soil carbon stocks and NPP (Fig. 3A and C). These linear relationships imply the age of fluvially exported biospheric OC reflects soil carbon turnover at the basin scale, the latter being controlled by MAT and MAP (4, 5). Still, fatty-acid and lignin-phenol $^{14}$C ages are on average 43.6- and 16.6-fold greater than the corresponding $\tau_{\text{soil}}$. To further probe these relationships, we additionally considered recent spatially resolved estimates of soil $^{13}$C ages (34). Biomarker $^{14}$C ages display similar positive relationships with catchment-weighted soil $^{14}$C ages integrated from 0 to 100 cm depth, further demonstrating the strong imprint of soil OC aging processes on fluvial biomarker radiocarbon ages (Fig. 3B and D, Materials and Methods, and SI Appendix, Table S4). Unlike $\tau_{\text{soil}}$, however, fatty-acid and lignin-phenol $^{14}$C ages are on average 2.5- and 5.0-fold lower than corresponding soil $^{14}$C ages, suggesting that lateral transport processes do not fully capture older, deeper soil horizons. This interpretation is supported by correlations between biomarker $^{14}$C ages and catchment-weighted soil $^{14}$C ages integrated from 0 to 30 cm depth, in which biomarkers display similar or slightly older $^{14}$C ages than corresponding soils, and integrated from 30 to 100 cm depth, in which soils display significantly older $^{14}$C ages than corresponding biomarkers (SI Appendix, Fig. S6).

Nonetheless, the observed discrepancy between biomarker $^{14}$C ages and $\tau_{\text{soil}}$ contrasts with the much closer agreement between biomarker $^{14}$C ages and soil mean carbon age, regardless of soil integration depth (Fig. 3 and SI Appendix, Fig. S6). This offset between both biomass $^{14}$C ages and catchment-weighted soil $^{14}$C ages on one hand and $\tau_{\text{soil}}$ on the other hand likely reflects the fundamental principles governing organic matter degradation and aging. Natural organic matter is compositionally heterogeneous, with age heterogeneity evident even within individual compound populations (20, 35). Complex interplay between environmental properties and chemical composition results in widely variable OC degradation rates (36). We therefore attribute discrepancies between riverine biospheric-carbon age and $\tau_{\text{soil}}$ to a heavy-tailed distribution of OC degradation, as has been hypothesized previously (37). Assuming degradation rate constants follow a lognormal distribution with given variance (38), the ratio between mean age and turnover time is proportional to the natural exponential function of the variance (39). Accepting that mean ages can be approximated using $^{14}$C ages (Materials and

![Fig. 3. Relationships between $\tau_{\text{soil}}$, soil mean carbon age, and biomarker $^{14}$C age. (A and B) Plant-wax fatty-acid and (C and D) lignin-phenol $^{14}$C ages as a function of weighted-catchment $\tau_{\text{soil}}$ (Left, ref. 5), and soil mean carbon age (0 to 100 cm, Right, ref. 34). Solid and dashed black lines are reduced major-axis regression lines; reported values are the corresponding reduced major-axis regression slopes and $r^2$ values (Materials and Methods). Uncertainty ($\pm 1\sigma$) is always smaller than marker points.](https://doi.org/10.1073/pnas.2011585118)
Methods), the biomarker $^{14}$C age versus $\tau_{\text{soil}}$ slopes (Fig. 3A and C) provide estimates of the globally averaged ratio between mean age and turnover time. These ratios correspond to log-normal degradation rate-constant variances of 3.6 ± 0.7 using fatty acids ($\mu \pm 1\sigma, n = 44$) and 2.7 ± 0.8 using lignin phenols ($\mu \pm 1\sigma, n = 16$), consistent with experimentally determined plant OC degradation rate-constant variances of ~1.0 to 4.0 (38, 39).

The scaling of biomarker $^{14}$C ages (and soil mean carbon age) with ecosystem turnover time reveals the heterogenous nature of organic matter and the associated wide distribution of OC degradation rates. This is particularly relevant in the context of potential soil OC destabilization upon increases in temperature and precipitation since OC pools characterized by low decay rates (i.e., long turnover) are proportionally more vulnerable to destabilization. Specifically, the theory of temperature-dependent activation energies predicts that turnover of long-lived OC pools (characterized by low decay rates) will increase proportionally more than that of short-lived OC pools (characterized by high decay rates) for a given increase in temperature or precipitation (40). As such, the prevalence of wide OC degradation rate distributions observed here implies that some fraction of the global soil OC reservoir is characterized by low decay rates and is thus susceptible to destabilization upon increases in temperature and precipitation—a positive climate feedback.

Interpreting Exported Carbon Ages. Lignin-phenol $^{14}$C ages, fatty-acid $^{14}$C ages, and $\tau_{\text{soil}}$ all exhibit significant negative power-law relationships with MAT and the logarithm of MAP (Fig. 4). For both fatty acids and lignin phenols, this implies a ~0.4 order-of-magnitude decrease in mean age for each 10-degree increase in MAT. Interestingly, this power-law exponent is statistically identical for $\tau_{\text{soil}}$ and both biomarker classes despite differences in their absolute $^{14}$C ages and the fact that they likely reflect different biospheric-fatty acids and lignin phenols, this implies a more than that of short-lived OC pools (characterized by high decay rates) for a given increase in temperature or precipitation (40). As such, the prevalence of wide OC degradation rate distributions observed here implies that some fraction of the global soil OC reservoir is characterized by low decay rates and is thus susceptible to destabilization upon increases in temperature and precipitation—a positive climate feedback.

Environmental controls on $\tau_{\text{soil}}$ and biomarker $^{14}$C ages. Logarithmic plant-wax fatty-acid $^{14}$C ages (black circles), lignin-phenol $^{14}$C ages (white squares), and catchment $\tau_{\text{soil}}$ [gray triangles (5)] as functions of (A) MAT and (B) logarithmic MAP (SI Appendix, Tables S1 and S3). Solid black, dashed black, and gray lines are fatty-acid–, lignin-phenol–, and $\tau_{\text{soil}}$-reduced major-axis regression lines. Relationship slopes and $r^2$ values are as follows: (A) fatty-acid $^{14}$C ages: slope = −0.036 ± 0.004, $r^2 = 0.62$; lignin-phenol $^{14}$C ages: slope = −0.036 ± 0.006, $r^2 = 0.67$; and $\tau_{\text{soil}}$: slope = −0.030 ± 0.002, $r^2 = 0.83$. (B) Fatty-acid $^{14}$C ages: slope = −1.46 ± 0.17, $r^2 = 0.55$; lignin-phenol $^{14}$C ages: slope = −1.44 ± 0.30, $r^2 = 0.56$; and $\tau_{\text{soil}}$: slope = −1.13 ± 0.11, $r^2 = 0.72$. Uncertainty (±1σ) is always smaller than marker points.

Materials and Methods

Sample Selection. Rationale for selection of river systems. We focus on 36 globally distributed river systems that collectively account for one-third of the global exorheic land surface and for 42, 29, and 20% of the global riverine water, sediment, and POC discharge, respectively (SI Appendix, Table S3). The majority of these rivers have been the subject of prior in-depth biochemical studies. The 36 rivers included in this study nearly cover the full range of intrinsic basin properties (e.g., catchment area, latitude, relief, physical erosion rate, discharge, and POC flux) and are evenly distributed across the continuum of geomorphic characteristics (SI Appendix, Fig. S1). This ensures that our results are not biased toward any particular set of basin properties. Rationale for focus on POC as opposed to DOC. We focus on organic matter that is associated with fluvioloxically derived, or fluviomarine influenced sediments either transported as suspended particles or recently deposited near the terminus of the river system. The translocation and sequestration of POC in marine sediments is considered to influence atmospheric CO$_2$ over millennial and longer timescales, whereas DOC is efficiently remineralized in coastal waters (16, 18). This POC leaves a legacy of terrestrial carbon fixation that can be traced in the sediment record and can be used to reconstruct the full range of sample types (e.g., suspended sediments, river flood deposits, offshore deposioners, etc.). Our statistical analysis did not identify any obvious biases related to the heterogenous nature of the sample set. Still, below we discuss the influence of a set of individual parameters in the context of the overall observed variability and trends described.

Sample type. Our sample set includes suspended sediments, bed sediments, recently deposited bank deposits (i.e., flood deposits), and floodplain sediments, as well as fluviomarine influenced sediments collected at or close to
the river terminus. In general, biospheric OC tends to be concentrated in the finer grain-size fractions, likely reducing potential impact from sorting processes. This is not always the case, as demonstrated by the Yellow River, where sediments were presieved to concentrate finer-grained (<63 μm) material, as well as to exclude coarse-grained detritus that may have been locally introduced (e.g., from riverbank erosion) to the river. Nevertheless, given potential differences in grain size distribution and associated hydrodynamic characteristics, it is important to assess associated variability in 14C characteristics. In the Yellow River, for example, plant-wax 14C ages have been found to vary among grain-size fractions of suspended sediments (44). For systems for which more than one sample type is available (i.e., Brahmaputra, Danube, Ganges, and Mississippi), a comparison of 14C characteristics reveals no systematic offsets between sample type. In the Danube River, vertical water-column profiles of plant-wax δ13C and 14C ages showed no systematic trend with depth (45), suggesting plant waxes are relatively homogenous within the water column and rather insensitive to hydrodynamic sorting. This likely results from the association of plant-wax compounds with mineral surfaces (especially clay minerals) and the uniform distribution of fine-grain clays in the water column (46, 47). We nonetheless include sample type in the statistical analysis since it is significantly correlated with lignin 14C age (Fig. 2).

Seasonal and interannual variability. Most rivers included in this study exhibit large seasonal variations in water and sediment discharge, which has been linked to particulate 14C composition (53, 54). Our sampling generally does not capture seasonal variations in the 14C age of biomarkers. The limited number of investigations of temporal variability in rivers does not reveal marked seasonal variations in biomarker 14C age [e.g., plant waxes in the Yellow River (49); lignin phenols in the Mekong River (50)]. Furthermore, in many cases, the samples investigated in this study were collected during or shortly following maximum discharge, which accounts for much of the annual sediment and OC load. As such, we argue that overall, our sampling is not affected by a strong seasonal bias. Interannual variability of riverine sediment load and bulk POC composition has been reported for several river systems (51, 52). Our dataset includes samples collected over different years for several river systems (Brahmaputra, Danube, Ganges, Yangtze, and Yellow River). Compared to the observed overall range of variability, none of the observed interannual variations in fatty-acid 14C ages suggests that interannual variations are not a significant driver of the observed trends in biomarker 14C ages.

Sample collection date. Samples have been collected between 1976 and 2016, a period of time over which atmospheric 14C, and hence freshly produced terrestrial plant organic matter, has exhibited a marked decline in response to the redistribution of “bomb” 14C (from above-ground thermonuclear weapons testing in the late 1950s and early 1960s) in the earth’s surface carbon reservoirs (43, 53). Recently synthesized biospheric carbon then transmits this bomb 14C signal through river basins, inducing time-variable deviations from natural 14C levels. We have examined this potential influence on observed signals for selected river systems in two different ways: 1) through comparison of data from samples from the same river system but collected at different times over the last few decades; and 2) through analysis of statistically calibrated and uniformly calibrated modern-day sequences with well-defined chronologies that span a time interval encompassing the bomb spike (20, 35). Although the influence of the bomb 14C spike clearly manifests itself in plant-wax fatty acids, the induced variability is small relative to the overall 14C variability observed within the global dataset presented in this study. These muted changes in plant-wax fatty-acid 14C ages require the presence of at least two different aged populations—one reflecting rapid (i.e., within years to decades) transfer of these tracer molecules from biological source to sedimentary sink and a second, substantially preaged (i.e., hundreds to several thousand years) component that implies substantial retention prior to export and deposition. In both of the above studies, model results suggest that the majority (49 to 83%) of the plant-wax signal derives from a preaged pool that likely corresponds to a mineral-bound soil OC component. Hence, sensitivity to bomb 14C is relatively small, and the date of sampling does not impart a large bias on our results.

Anthropogenic influences. A key consideration in any study of modern rivers is the extent to which modern observations of fluvial-sediment flux and composition have been impacted by anthropogenic activity, both within the catchment area (e.g., land-use change) and on the hydrological network (e.g., damming, channelization, etc.). As a proxy for anthropogenic influences, we include in our statistical analysis the fraction of each river basin that is impacted by anthropogenic land-use change (i.e., agriculture and urbanization) (SI Appendix, Table S3). This approach implicitly assumes the following: 1) Land use is an accurate proxy for all anthropogenic disturbances, and 2) modern land-use change extent is representative of perturbations over the timescale of biomarker 14C ages (i.e., centuries to millennia). Nevertheless, our statistical correlation results indicate that biomarker 14C ages are more strongly correlated with climate as opposed to anthropogenic variables (SI Appendix, Table S4), suggesting that differences in the type and duration of anthropogenic disturbance likely impart only a small impact on resulting biomarker 14C ages.

Sample Collection and Processing. Sample collection.

Samples include suspended sediments obtained via filtration of river water, riverbed sediments sampled via a grab or bedload sampler, recently deposited river bank and floodplain sediments, and fluvially influenced coastal sediments deposited proximal to the mouth of the river. For river flank, floodplain, and coastal sediments, emphasis was placed on sampling deposits where fine-grained sediments accumulate, such as quiescent areas characterized by weak river and coastal currents. Sample processing and analysis procedures vary somewhat between field campaigns. Detailed procedures are provided in previous publications and are briefly summarized below (53, 54). Our sampled sediments generally do not capture seasonal variations in the 14C age of biomarkers. Samples were then stored, refrigerated, or frozen before freeze-drying in the laboratory. For selected samples (mostly floodplain materials), the sediment was washed sieved to <63 μm to remove large fragments that may have been directly introduced (e.g., via riverbank erosion). The <63-μm fraction is also considered to more closely resemble the riverine suspended load owing to its overall smaller grain size compared to deposited sediments (45, 56, 57).

Plant-wax lipids were extracted from freeze-dried sample aliquots using a mixture of dichloromethane and methanol. The lipid extract was subsequently treated to obtain a fatty-acid methyl esters (FAMEs). The FAMEs were further purified via chromatography and ultimately isolated by preparative gas chromatography (62). Isolated compounds were subsequently analyzed by accelerator mass spectrometry (AMS) (14C contents) and carbon chromatography (GC-IRMS) (δ13C). Resulting 14C data are corrected for any bomb 14C contribution as well as other 14C age correction factors (Table S4), and reported using the fraction modern (Fm) notation and as radiocarbon ages (63). We report abundance-weighted average 14C compositions of long-chain FA homologs in the 24 to 32 carbon number range. In some cases, sample availability and/or technical issues led to averaging over a subset of homologs within this carbon number range. For coastal sediments that may have additional sources of midchain FA homologs (e.g., ref, 35), we used corresponding δ13C values from GC-IRMS to select homologs that are exclusively derived from terrestrial vascular plants. Lignin-derived phenols were recovered by alkaline oxidative hydrolysis (CuO oxidation) of freeze-dried or solvent-extracted sediments (64). Resulting hydrolysis products were then purified via high-performance liquid chromatography (50, 64) and subsequently measured for 13C content by AMS.

Biomarker 14C data corrections. Compound-specific radiocarbon data were corrected using mass-balance equations for procedural blanks and, in the case of fatty acids, for the addition of derivative carbon during methylation with methanol with a known 14C composition. Procedural blanks varied depending on the instrumental set up used (e.g., at WHOI versus at ETH) and the measurement date, as these methods have been continuously refined over the course of this study. Overall, procedural blanks were in the range of 1 to 2 μg C, with compositions intermediate between modern and dead. Details of blank assessments and corrections can be found in Santos et al. (65), French et al. (20), Haghipour et al. (66), and Feng et al. (64). The
uncertainty related to blank corrections varies from sample to sample, primarily according to sample size and 14C age. Overall, these uncertainties remain below 0.05 fm units and are therefore small compared to the variability observed across our entire database.

**Geospatial Analysis.** All geospatial analyses were performed in ArcGIS version 10 (ESRI Corporation). River basin outlines were calculated either upstream of each sampling point (for suspended-sediment and bed-sediment samples) or at the river mouth (for shelf-deposit samples), and environmental control variables were calculated using the following data sources: 1) Elevation, slope, and floodplain fractional area (here defined as area with slope less than 1% rise) were calculated using the United States Geological Survey (USGS) GTOPO30 digital elevation model with 30-arc-second resolution [1 km at the equator (67)]. 2) Catchment-averaged MAT, temperature seasonal coefficient of variability (TCV), MAP, and precipitation seasonal coefficient of variability (PCV) were calculated from the raster data set of New et al. (68) with 10° resolution. 3) τwa and NPP were calculated from the raster data set of Bloom et al. (5) with 1° resolution. 4) Total ecosystem carbon turnover time (ecosystem τ) and soil carbon stocks were calculated from the raster data set of Carvalhais et al. (4) with 0.5° resolution. 5) Anthropogenic land-use fractional area was calculated as the sum of “croplands,” “urban and built-up,” and “cropland/natural vegetation mosaic” pixels (International Geosphere Biosphere Program classification) extracted from the USGS Global Land Cover Characterization raster data set version 2.0 with 30-arc-second resolution (69). 6) Soil carbon 14C ages (0 to 30 cm, 30 to 100 cm, and 0 to 100 cm depth integrated) were calculated from the raster data set of Shi et al. (34) with 0.5° resolution.

All other environmental control variables were compiled from the literature (15, 17, 70). When comparing biomarker 14C ages and soil turnover times, we implicitly assume that the soil turnover times determined using estimates of soil C stocks and NPP are representative of the average turnover times over the time scales referred to in the biomarker 14C ages (centuries to millennia). While short-term small variations in NPP have likely occurred over these timescales and regional-scale climate variability during the late Holocene locally impacted soil carbon stocks (71), this relatively quiescent time interval is unlikely to have seen large, globally coherent variations in soil carbon stocks. As such, temporal variations in τwa are unlikely to contribute significantly to the systematic difference observed between τwa and biomarker 14C ages (i.e., since plant-wax fatty-acid and lignin-phenol 14C ages are on average 43.6- and 16.9-fold greater, respectively, than corresponding τwa)

For spatially resolved control variables (i.e., elevation, slope, floodplain fraction, MAT, TCV, MAP, POC, τwa, NPP, ecosystem τ, and soil C stocks), weighting factors were calculated using hydrologically conditioned versions of the GTOPO30 digital elevation model (i.e., Hydro1k and HydroSHEDS (72, 73)). Hydrologic distance upstream of each sampling location was calculated using the “Flow Length” feature in ArcGIS and was weighted as an exponential decay following

\[
w = e^{-k/l}, \quad [1]
\]

where w is the weighting factor (ranging from 0 to 1), l is the upstream distance, and k is the reciprocal of the prescribed e-folding distance. All spatially resolved control variables were taken as the weighted-catchment averages.

**Statistical Analyses.**

Multiple linear regression. Regression analyses were performed using the Numpy and Scipy packages in Python version 3.5; all analysis code is provided in Dataset S1. To determine the strength of relationships between environmental control variables and biomarker and POC response variables, OLS MLR was performed following standard practices (74). In summary, all data were first “whitened” by subtracting the mean and dividing by the SD for each variable in order to facilitate comparisons between variables with differing units and ranges. The regression parameter matrix, B, was then calculated as

\[
B = (X'X)^{-1}X'Y, \quad [2]
\]

where X is the whitened matrix of environmental control variables, and Y is the whitened matrix of biomarker and POC response variables. The matrix of MLR-predicted response variable best estimates, \(\hat{Y}\), was then calculated as

\[
\hat{Y} = XB, \quad [3]
\]

and the matrix of residuals was calculated as

\[
Y = \hat{Y} - Y. \quad [4]
\]

The matrix of correlation strengths between X and Y, termed \(S_{XY}\), was then calculated following

\[
S_{XY} = \frac{1}{n}X'Y, \quad [5]
\]

where n is the number of samples in the data set, and correlation P values were calculated individually for each control and response variable pair using standard QLS methods (74). Reported MLR \(r^2\) values are the square of the diagonal elements of \(S_{XY}\). The optimal e-folding \(k\) value was calculated using an inverse approach by repeating Eqs. 1-5 and choosing the value that maximizes the average resulting \(r^2\) value (SI Appendix, Fig. 53). Because scatterplots of PCVs presented in this study generally contain uncertainty in both x and y variables (e.g., Figs. 3 and 4), predictive relationships were calculated using reduced major-axis regression (75). Note that some environmental control variables, and particularly biomarker response variables, contained missing data entries; missing data were ignored when performing statistical calculations by utilizing Numpy “masked” arrays (reference Dataset S1 for details).

Finally, we test if and how averaging all samples of a given sample type (i.e., suspended sediment, bedload sediment, or shelf/slope sediment) within a given river basin impacts the results of our statistical analysis. We repeated Eqs. 1-5 using either “catchment averaged” or “all reported samples” as the response variable dataset (i.e., SI Appendix, Tables S1 and S2, respectively). Resulting correlation coefficients exhibit only small differences and are largely independent of our choice of response variable dataset (SI Appendix, Fig. S8). We therefore use the catchment averaged dataset for all subsequent calculations as this avoids potential biases that could arise because of uneven sampling density between river basins.

**Redundancy analysis.** To extract the canonical axes and to determine the loadings of each sample, environmental control variable, and biomarker and POC response variable onto each canonical axis, RDA was performed following Legendre and Legendre (76) using the Numpy and Scipy packages in Python version 3.5. All analysis code is provided in Dataset S1. RDA is a canonical extension of MLR that transforms all control variables into orthogonal (linearly uncorrelated) axes and determines the fraction of sample variance explained by each orthogonal axis; it is ideally suited for situations with highly correlated control variables, as is the case here (76). Heuristically, RDA extracts the principal components of response variables as predicted by control variables; that is, it calculates the amount of variance within the response variables that can be explained by the set of control variables and maps all variables onto a set of underlying orthogonal axes. Briefly, RDA involves independently performing principal component analyses on Y, the matrix of MLR-predicted response variables, and on \(Y^T\), the matrix of residuals. Analogous to Eq. 5, the correlation matrix between MLR-predicted response variables is first calculated as

\[
S_{Y^T} = \frac{1}{n}Y^TY. \quad [6]
\]

Then, the following eigenvalue problem is solved:

\[
(S_{Y^T} - \lambda I)U = 0, \quad [7]
\]

where \(\lambda\) is the jth canonical eigenvalue, \(\omega_i\) is the jth canonical eigenvector, and \(I\) is the identity matrix; the response variable loadings onto the jth canonical axis (termed “species scores”) are thus the entries of \(\omega_i\). Eqs. 6 and 7 are then repeated using \(Y\) instead of \(Y^T\) to calculate the noncanonical eigenvalues and eigenvectors. The percentage of response variable variance explained by each axis is calculated as \(\lambda_i\) divided by the sum of all (canonical plus noncanonical) eigenvalues. Finally, constrained sample loadings (“constrained site scores”) onto each canonical axis, termed Z, are calculated as

\[
Z = YU, \quad [8]
\]

and control variable loadings (“constraining variable scores”) onto each canonical axis, termed C, are calculated as

\[
C = BU, \quad [9]
\]

where U is the column-wise matrix of \(\omega_i\) eigenvectors. Species scores, constrained site scores, and constraining variable scores for any two canonical axes can then be plotted as in Fig. 28; the angles between species scores and constraining variable scores thus represent the strength of their linear correlation. Reference Dataset S1 for further details and analyses.
Data Availability. All study data are included in the article and/or supporting information.

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1. P. Friedlingstein et al., Global carbon budget 2019. Earth Syst. Sci. Data 11, 1783–1838 (2019).
2. B. N. Sulman et al., Multiple models and experiments underscore large uncertainty in soil carbon dynamics. Biogeochem. 141, 109–123 (2018).
3. J. Dowling et al., Twenty-first century compatible CO2 emissions and airborne fraction simulated by CMIP5 earth system models under four representative concentration pathways. J. Clim. 26, 4398–4413 (2013).
4. N. Carvalhais et al., Global covariation of carbon turnover times with climate in terrestrial ecosystems. Nature 513, 212–217 (2014).
5. A. A. Bloom, J. F. Exbrayat, I. R. van der Velde, L. Feng, M. Williams. The decadal state of the terrestrial carbon cycle: Global retrievals of terrestrial carbon allocation, pools, and residence times. Proc. Natl. Acad. Sci. U.S.A. 113, 1285–1290 (2016).
6. N. S. Torn, S. E. Trumbore, O. A. Chadwick, P. M. Vitousek, D. M. Hendricks, Mineral control of soil organic carbon storage and turnover. Nature 389, 170–173 (1997).
7. M. W. Schmidt et al., Persistence of soil organic matter as an ecosystem property. Nature 478, 49–56 (2011).
8. S. Doetterl et al., Controls, experimental tools and models to address C cycling in dynamic landscapes. Annu. Rev. Earth Planet. Sci. 43, 123–157 (2015).
9. R. Z. Abramoff, M. S. Torn, K. Georgiou, J. Tang, W. J. Riley, Soil organic matter temperature sensitivity cannot be directly inferred from spatial gradients. Biogeochem. 321, 1455–1456 (2008).
10. R. Z. Abramoff, M. S. Torn, K. Georgiou, J. Tang, W. J. Riley, Soil organic matter temperature sensitivity cannot be directly inferred from spatial gradients. Global Biogeochem. Cycles 33, 761–776 (2019).
11. A. Behe, R. T. Barnes, J. S. Xu, E. Marin-Spiotta, Role of soil erosion in biogeochemical cycling of essential elements: Carbon, nitrogen, and phosphorus. Annu. Rev. Earth Planet. Sci. 46, 521–548 (2018).
12. T. J. Battin et al., The boundless carbon cycle. Nat. Geosci. 2, 598–600 (2009).
13. B. A. Fisher, A. K. Aufdenkampe, K. Yoo, R. E. Aalto, J. Marquard, Soil carbon redistribution and organic-mineral associations after lateral soil movement and mixing in a first-order forest watershed. Geoderma 319, 142–155 (2018).
14. S. Doetterl et al., Erosion, deposition and soil carbon: A review of process-level controls on measurement methodology and models to address C cycling in dynamic landscapes. Earth Sci. Rev. 154, 102–122 (2016).
15. R. G. Hilton, Climate regulates the erosional carbon export from the terrestrial biosphere. Geomorphology 277, 118–132 (2017).
16. K. Feng et al., Differential mobilization of terrestrial carbon pools in Eurasian Arctic river basins. Proc. Natl. Acad. Sci. U.S.A. 110, 14168–14173 (2013).
17. J. J. Hedges, R. G. Keil, R. Benner, What happens to terrestrial organic matter in the ocean? Org. Geochem. 27, 195–212 (1997).
18. N. Vally, G. P. Feuerhahn, T. I. Eglinton, Global carbon export from the terrestrial biosphere controlled by erosion. Nature 521, 204–207 (2015).
19. R. A. Benner, Burial of organic carbon and pyrite sulfur in the modern ocean: Its environmental signatures encoded in vascular plant biomarkers carried by the Brahmaputra rivers. Proc. Natl. Acad. Sci. U.S.A. 108, 16965–16970 (2011).
20. K. L. French et al., The boundless carbon cycle. Nature 517, 451 (2015).
21. J. N. Eicker, J. Bouchez, M. Lupker, J. Gaillardet, C. France-Lanord, L. Maurice, How important is it to integrate riverine suspended sediment chemical composition with depth? Clues from Amazon river depth-profiles. Geochim. Cosmochim. Acta 75, 6955–6970 (2011).
22. B. M. Voss et al., Seasonal hydrology drives rapid shifts in the flux and composition of dissolved and particulate organic carbon and major and trace ions in the Fraser River, Canada. Biogeosci. 12, 5597–5618 (2015).
23. M. Yu et al., Impacts of natural and human-induced hydrological variability on particulate organic carbon dynamics in the Yellow River. Environ. Sci. Technol. 53, 1119–1129 (2019).
24. E. G. Carvalhais et al., The boundless carbon cycle. Nat. Geosci. 6, 598–600 (2009).
25. S. Lin, T. I. Eglinton, C. France-Lanord, S. Myhre, M. Zhao, Pre-aged soil carbon as a major component of the Yellow river suspended load: Regional significance and global relevance. Earth Planet. Sci. Lett. 414, 77–86 (2015).
26. K. L. French et al., Mineralogical inventory of terrestrial organic matter deposited in the Black Fan. Earth Planet. Sci. Lett. 394, 11967–11987 (2013).
27. J. E. Blair, R. C. Aller, The fate of terrestrial organic carbon in the marine environment. Annu. Rev. Mar. Sci. 4, 401–423 (2012).
28. T. R. Marwick et al., The age of river-transported carbon: A global perspective. Global Biogeochem. Cycles 29, 613–627 (2015).
29. D. R. Griffith, R. T. Barnes, P. A. Raymond, Inputs of fossil carbon from wastewater treatment plants to U.S. rivers and oceans. Environ. Sci. Technol. 43, 5647–5651 (2009).
30. T. I. Eglinton et al., Variability in radiocarbon ages of individual organic compounds from marine sediments. Science 272, 796–799 (1997).
31. T. I. Eglinton, G. Eglinton, Molecular proxies for paleoclimatology. Earth Planet. Sci. Lett. 275, 1–16 (2008).
32. C. N. Jex et al., Lignin biogeochemistry: From modern processes to Quaternary archives. Quat. Rev. 57, 46–59 (2014).
33. T. van der Voort et al., Diverse soil carbon dynamics expressed at the molecular level. Geophys. Res. Lett. 44, 11840–11850 (2017).
34. J. D. Hemingway et al., Mineral protection regulates long-term global preservation of natural organic carbon. Nature 570, 228–231 (2019).
35. J. D. Hemingway, E. Scheufele, B. J. Dinga, H. Pryer, V. Galy, Multiple plant-wax compounds record differential sources and ecosystem structure in large river catchments. Geochim. Cosmochim. Acta 223, 389–399 (2018).
36. J. H. Whitehead et al., Pangean great paleoecology on the cusp of the end-Triassic extinction. Palaeogeogr. Palaeoclimatol. Palaeoecol. 301, 1–17 (2011).
37. A. P. McIntosh, E. A. Osborne, A. R. Gagnon, B. Fy, G. A. Jones, TIC, TOC, DIC, DOC, PIC, POOC—unique aspects in the preparation of oceanographic samples for 14CAMS. Nucl. Instrum. Methods Phys. Res. B 92, 162–165 (1994).
38. M. Christf et al., The ETH Zurich AMS facilities: Performance parameters and reference materials. Nucl. Instr. Meth. Phys. Res. Sec. B 294, 29–38 (2013).
39. C. B. McInerney et al., Online 14C and 13C gas measurements by EA-IRMS at AMS ETH Zurich. Radiocarbon 59, 893–907 (2017).
40. T. I. Eglinton, L. I. Alwihale, J. E. Bauer, E. R. M. Druffel, A. P. McNichol, Gas chromatographic isolation of individual compounds from complex matrices for radiocarbon dating. Anal. Chem. 68, 780–782 (1996).
41. M. Stuiver, H. A. Polach, Discussion: Reporting of 14C data. Radiocarbon 19, 355–367 (1977).
64. X. Feng et al., 14C and 13C characteristics of higher plant biomarkers in Washington margin surface sediments. Geochim. Cosmochim. Acta 185, 14–30 (2013).
65. G. M. Santos et al., Blank assessment for ultra-small samples: Chemical extraction and separation vs. AMS. Radiocarbon 52, 1322–1335 (2010).
66. N. Haghipour et al., Compound-specific radiocarbon analysis by elemental analyzer-accelerator mass spectrometry. Precision and limitations. Anal. Chem. 91, 2042–2049 (2019).
67. Earth Resources Observation and Science Center/U.S. Geological Survey/U.S. Department of the Interior, USGS 30 ARC-second Global Elevation Data, GTOPO30 (1997). Research Data Archive at the National Center for Atmospheric Research, Computational and Information Systems Laboratory. https://doi.org/10.5065/A1Z4-EE71. Accessed 13 March 2020.
68. M. New, D. Lister, M. Hulme, I. Makin, A high-resolution data set of surface climate over global land areas. Clim. Res. 21, 1–25 (2002).
69. A. S. Belward, ed., The IGBP-DIS global 1 km land cover data set (DISCover)-proposal and implementation plans: IGBP-DIS Working Paper No. 13, Toulouse, France, pp. 61 (1996).
70. B. Peucker-Ehrenbrink, Land2Sea database of river drainage basin sizes, annual water discharges, and suspended sediment fluxes. Geochem. Geophys. Geosyst. 10, 6 (2009).
71. R. H. Smittenberg, T. I. Eglinton, S. Schouten, J. S. Damste, Ongoing buildup of refractory organic carbon in boreal soils during the Holocene. Science 314, 1283–1286 (2006).
72. K. L. Verdin, S. K. Greenlee, “Development of continental scale digital elevation models and extraction of hydrographic features” in Proc. 3rd Int. Conf./Workshop on Integrating GIS and Environmental Modeling, Santa Fe, New Mexico, January 21-26, 1996 (National Center for Geographic Information and Analysis, Santa Barbara, California, 1996).
73. B. Lehner, K. Verdin, A. Jarvis, New global hydrography derived from spaceborne elevation data. Eos (Wash. D.C.) 89, 93–94 (2008).
74. N. R. Draper, H. Smith, Applied Statistical Analysis (John Wiley & Sons, Inc., New York City, New York, ed. 3, 1998), pp. 217–234.
75. J. M. Rayner, Linear relations in biomechanics: The statistics of scaling functions. J. Zool. 206, 415–439 (1985).
76. P. Legendre, L. Legendre, Numerical Ecology: Developments in Environmental Modelling (Elsevier B.V., Amsterdam, The Netherlands, ed. 3, 2012), pp. 424-520.