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Long-term patterns of peat accumulation and organic matter decomposition in Costa Rican peatlands

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Abstract. Tropical peatlands, compared to their boreal counterparts, are vastly understudied despite these ecosystems acting as a significant terrestrial carbon sink, sequestering 100-300 gigatons of carbon. In particular, the low number of field-based studies from Latin America and the Caribbean limits our knowledge of these important wetland ecosystems: across the tropical Panamerican region, peatland location, soil characteristics, inception ages, and carbon accumulation histories remain largely unknown. These datasets are needed to inform a mechanistic understanding of why peat develops in certain areas but not in others, both in terms of peat initiation conditions as well as the factors that enable peat to subsist over centuries and millennia. Here we present extensive, high-resolution laboratory datasets from 11 peat cores from four peatland types from Costa Rica (high-elevation, riverine, coastal palm swamp, and mangrove). A multi-proxy palaeoecological approach was employed to shed light on the successional pathways and past conditions that have allowed these peatlands to form, as well as to provide a first estimate of their carbon stock. The core characterization includes radiocarbon dating, loss-on-ignition, carbon and nitrogen content, and plant macrofossils. Fourier transform infrared spectroscopy (FTIR) was also used to assess changes in organic matter quality across sites and over time, ultimately to clarify the role of recalcitrant material in tropical peat accumulation. Overall, this research provides a basis for understanding long-term carbon accumulation within Caribbean tropical peatlands and is critical to advancing knowledge of the structure of tropical peatland systems.

1 Introduction

Tropical peatlands are estimated to occupy around 14%, and potentially up to 30%, of the global peatland area (Gumbricht et al., 2017; UNEP, 2022; Xu et al., 2018). Carbon stock estimates for tropical peatlands similarly vary by a factor of about two, from ~120-130 gigatonnes (Dargie et al., 2017; Leifeld and Menichetti, 2018) to 288 gigatonnes (Ribeiro et al., 2021). The reason for these large ranges is multifold and mainly attributable to the scarcity of field-based datasets pertaining to (1) peatland distribution / surface area, (2) peat depth, and (3) peat carbon density (Gumbricht et al., 2017; Melton et al., 2022; Page et al., 2011). At times, the tropical peatland carbon pool has been estimated on the basis of peat carbon density datasets from extra-tropical regions (Yu et al., 2010), adding unquantifiable uncertainties. Similarly, averaged bulk density and carbon concentration values from Southeast Asian peatlands have been extrapolated to the Panamerican region due to a lack of data (Page et al., 2011). These data and knowledge gaps undermine our ability to quantify the tropical peatland soil carbon pool and assess the past, present, and future roles of these ecosystems in the global carbon cycle.

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In the tropics, the existence of peat has often been referred to as 'surprising'. This idea stems from the notion that tropical terrestrial ecosystems tend to be characterized by rapid rates of microbial decomposition due to consistently hot and humid conditions (Knorr et al., 2005; Meentemeyer, 1978). The accumulation of peat in the tropics therefore requires special conditions that either dampen the intensified decomposition process near the soil surface and/or a carbon-soil storing mechanism that bypasses the aerobic decay zone. Previous studies have proposed the following conditions and mechanisms, which are not mutually exclusive: (1) peat soils must be flooded (or waterlogged all the way up to the near surface) for long enough throughout the year to induce anoxia, which limits organic matter decay and allows for peat buildup (Chimner and Ewel, 2005; Gillman et al., 2015); (2) plant litter inputs to the peat must be very high to maintain a positive balance between litter production and peat decomposition; (3) the peat litter is composed of recalcitrant compounds and/or abundant stems, branches, and wood, all of which inhibit decomposition (Couwenberg et al., 2010; Dommain et al., 2015; Hodgkins et al., 2018; Hoyos-Santillan et al., 2015; Phillips et al., 1997; Verbeke et al., 2022; Wright et al., 2013); and (4) root biomass inputs (down into the constantly anoxic zone) circumvent the intensive near-surface peat decomposition processes and lead to peat buildup (Chimner & Ewel, 2005).

The bulk of the literature on tropical peatland carbon storage dynamics stems from Southeast Asia (Cobb et al., 2024; Dadap et al., 2022; Dommain et al., 2015; Hergoualc'h and Verchot, 2011; Hoyt et al., 2020; Ruwaimana et al., 2020; Sasmito et al., 2025) and, to a lesser extent, the Congo Basin (Dargie et al., 2017; Crezee et al., 2022; Garcin et al., 2022; Young et al., 2023). In the lowlands of Latin America and the Caribbean (LAC), a region that may encompass the largest tropical peatland complexes in the world (Gumbricht et al., 2017), regional knowledge on peatland distribution, extent, and dynamics remains sparse. Across this region, the most extensive work has been performed in Western and Central Amazonia (Lähteenoja et al., 2009, 2012, 2013; Lähteenoja and Page, 2011; Draper et al., 2014; Dargie et al., 2024; Winton et al., 2025; Roucoux et al., 2013; Lawson et al., 2014, 2026; Hastie et al., 2022), leaving Central America and the Caribbean particularly understudied. One exception is the Changuinola peat deposit in Bocas del Toro, Panama, where detailed paleoecological information reveals the historical peatland developmental pathways (Cohen et al., 1989; Phillips, 1995; Phillips and Bustin, 1996; Phillips et al., 1997; Swindles et al., 2024). A few studies have also looked at carbon and water dynamics in the Changuinola peatlands (Baird et al., 2017; Girkin et al., 2025; Hedgpeth et al., 2025; Hoyos-Santillan et al., 2016; Sjögersten et al., 2020). But beyond Changuinola, there is little knowledge on where peat is found, how deep, old, and carbon-rich it might be, and what factors control long-term peat accumulation processes (Rabel and Loisel, 2024).

In this study, we present high-resolution, multi-proxy analyses from a series of peat cores that were collected across 9 sites that span four distinct hydroecoclimatic and geomorphic settings. The main goal of the study is to characterize peat accumulation dynamics across these diverse sites and decipher spatial and temporal differences across those peatlands, if any. First, we provide the paleoecological history from four of our study sites to document changes in the major peat-forming plant macrofossil types and humification to identify successional pathways. We also discuss the differences that were found across our sites. Second, we present peat basal ages for our 9 study sites to assess the timing of peat inception in Costa Rica. Third, we provide a suite of geochemical measurements (organic matter content, dry bulk density, carbon and nitrogen content, C/N ratio) that can be used to estimate carbon storage. Lastly, through the use of infrared spectroscopy, we endeavor to improve our understanding of the peat formation process itself. By correlating our plant macrofossil and von Post humification analyses with Fourier-Transform infrared spectroscopy (FTIR) data, we identify shifts in organic matter quality that coincide with successional changes and/or humification.

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For instance, the accumulation of recalcitrant compounds (e.g., aromatics) is expected in the deeper portions of the peat, alongside high levels of humification. An alternative scenario combines high root abundance with carbohydrate compounds at depth, which could indicate that peat buildup relies, at least in part, on root accumulation that bypasses the oxic zone. Overall, the findings of this study shed light on biogeochemical processes that underpin the formation, development, and resilience of tropical peatland ecosystems, thus providing knowledge to scientists, landowners, managers, and policymakers.

2 Methods

2.1 Study area

While Costa Rica is rarely mentioned as a peat-rich country, a recent probabilistic map suggests that it may harbor 1456 km^2, equivalent to 3% of its land area (Rabel et al. 2025; Fig. 1). With that said, it is important to note that Costa Rica does not yet possess an official peatland inventory. In July and August 2023, our team traveled across Costa Rica and visited over 15 sites (Loisel et al., 2024) that had been identified on the basis of published wetland and vegetation maps as well as few existing peat depth point data (SINAC, 2018; Villegas-Mejía, 2018). We corroborated those data with global-scale peatland probability maps (Gumbricht et al., 2017; Melton et al., 2022) and a preliminary map that was produced by Peters and Tegetmeyer (2019). In most places that were visited, we found peaty soils (i.e., organic-rich soil layers in excess of 30 cm in thickness, *sensu* Lourenco et al., 2023). Our site selection was ultimately based on site accessibility as well as regional representativity, such that we made sure to sample the following peatland types: coastal peat swamps, mangroves, riverine complexes, and montane peatbogs (Jiménez, 2016).

Costa Rican peatlands are found under a broad range of mean annual precipitation and precipitation seasonality (Rabel et al., 2025; Fig. 1), and different peatland types are delineated within this climate space. As such, the main peatland types mentioned above are also distinct in terms of their climatology: the Caribbean coastal sites (palm swamps and mangroves) receive high annual precipitation amounts with low precipitation seasonality (and annual precipitation increasing along a north-to-south latitudinal gradient along the coast), whereas the (inland) riverine sites are intermediate in both annual precipitation and its seasonality, which contrast with the montane sites that are characterized by lower annual precipitation with higher precipitation seasonality.

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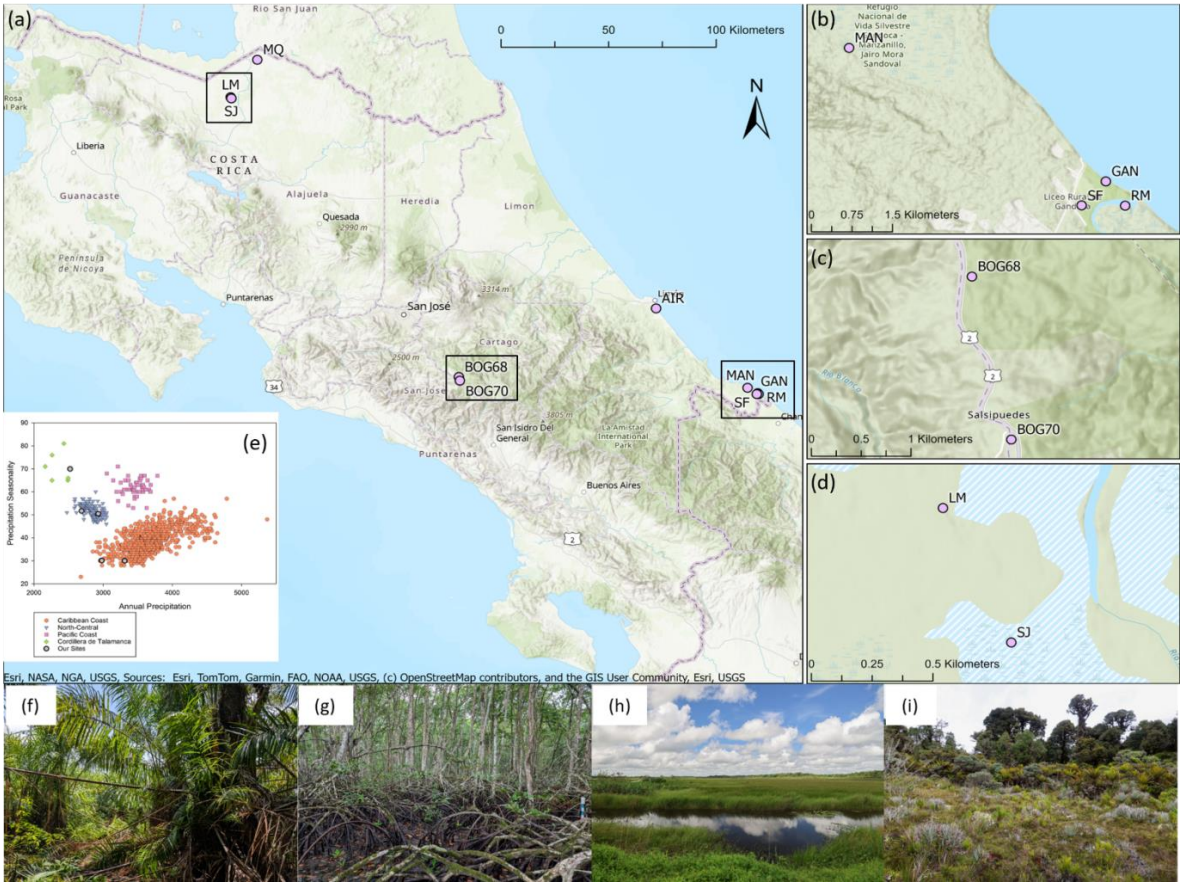

Figure 1: Locations and photos of sites from this study. (a) The country-wide Costa Rica map, (b) Inset map of the MAN, RM, and GAN sites, (c) Inset map of the BOG70 and BOG68 sites, and (d) Inset map of the LM and SJ sites. (e) Climate envelope for Costa Rican peatlands, adapted from Rabel et al., 2025. Each circle represents a pixel that has been classified as peat. Photos of the sites include: (f) The coastal palm swamp "Gandoca", (g) The "Red Mangrove" site, (h) The riverine wetland complex "Medio Queso", and (i) the high-elevation "Bog 70" peatland. Photo credits: Hannah Mitchell and Patrick Campbell. Source for panels a-d: Esri, TomTom, FAO, USGS | Powered by Esri.

2.2 Study sites

The sections below describe the four main peatland types that were sampled and provide information for each coring site. Table 1 offers a summary of the most relevant information.

Peatland region	Site code	Site name	Latitude	Longitude	Elevation (m)	Peat depth (cm)	Full cores (n)	Bottom cores (n)
High elevation	BOG70	Bog 70	9.64768	-83.84597	2755.5	100	1	0
	BOG68	Bog 68	9.66292	-83.84952	2663.2	80	0	1

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Palm swamp	GAN	Gandoca	9.59253	-82.60241	10.8	130	1	1
	AIR	Airport	9.96024	-83.02612	6.1	140	0	1
	MAN	Manzanillo	9.61640	-82.64393	14.3	239	0	1
Mangrove	RM	Red Mangrove	9.58899	-82.59794	14.4	187	1	0
Riverine	MQ	Medio Queso	11.03382	-84.68874	32.2	230	1	1
	LM	Laguna Martin San Jeronimo	10.86676	-84.80004	35.7	96	0	1
	SJ		10.86676	-84.80004	35.7	208	0	1

Table 1: Summary information for the Costa Rican peatland study sites.

(1) Coastal palm swamps (n = 3) Those sites are located along the southern Caribbean coast of Costa Rica, between the towns of Gandoca and Limón. All three sites are dominated by palm swamp species *Raphia taedigera* and harbor other plants, shrubs, and tree species, such as *Campnosperma panamensis* (Cortés, 1998; Peters and Tegetmeyer, 2019). These coastal sites have developed in back-barrier topographic lows, in close proximity to the coastline (Fig. 1). The Gandoca site (GAN) is located in the southern part of the country, close to the Panama border; two peat cores were collected along a 1km-long transect that crosses the site (GAN-4; GAN-SF). The Manzanillo site (MAN) is located approximately 10 km north of GAN and it is part of the Gandoca-Manzanillo Wildlife Refuge and Ramsar site #783 (Ramsar, 2026). Lastly, the Airport site (AIR) is located just south of Limón, in Moín. Those three sites were either accessed by boat (via canals) or on foot. To our knowledge, these ecosystems remain unstudied, with the exception of prospective work in the 1980s and 1990s that was aimed at assessing the potential economic development of these peat deposits as fuel resources (Obando et al., 1995; Thayer et al., 1995).

(2) Mangroves (n = 1) The Red Mangrove site (RM) is located a few meters away from the Gandoca site (Fig. 1). These two ecosystems are separated by Laguna Gandoca, a meandering stream that reaches the Atlantic Ocean. The northern side of the stream supports peat-forming *Raphia taedigera* palm swamps, which are probably also aided by the back-barrier geomorphology that limits saltwater intrusion, whereas the southern shore is instead characterized by brackish water, tides, and *Rhizophora mangle* communities. As only 1% of Costa Rica's mangrove forests are located along the Caribbean coast, our RM site presents a rare opportunity to study one of the largest (and preserved) mangrove stands along the Caribbean coast (Álvarez-Sánchez and Piedra-Castro, 2020; Cortés, 1998, 2016). There are local studies on the floristic composition of those mangroves (Álvarez-Sánchez and Piedra-Castro, 2020) as well as the physico-chemical characterization of the Gangoca Lagoon (Coll et al., 2004), but otherwise there is little information on mangrove soil properties from this region with the exception of a study pointing to high soil organic matter content in the Gandoca mangroves (Rovai et al., 2018).

(3) Riverine sites (n = 3) Those sites are located in the north-central portion of Costa Rica, near the Nicaraguan border (Fig. 1). The riverine sites all constitute peatland/wetland complexes that experience relatively large water table level fluctuations on an annual basis, with surface flooding that can reach 2 m during the wet season (Moreno-Casasola and Warner, 2009; Pérez-Castillo

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et al., 2024). The 'Medio Queso' site (MQ) is located approximately 1 km away from the town of Los Chiles and is found within the watershed of the Rio Medio Queso. This wetland complex is used for pasture and it is frequently burned to maintain land clearing (Camacho-Navarro et al., 2017; Pérez-Castillo et al., 2024). A recent study at the site evaluated the impacts of fires on vegetation and surface peat chemistry (Camacho-Navarro et al., 2017; Pérez-Castillo et al., 2024). The MQ study site is primarily colonized by rushes (*Eleocharis interstincta*), though many sedge and small bush species are also found (Obando and Malavassi, 1993), along with small 'islands' of Cuba palm (*Acoelorraphe wrightii*). At the time of sampling (August 2023), the wet season had begun, but we were nevertheless able to walk onto the peatland to retrieve peat cores. Two peat cores (MQ-9; MQ-10) were collected along a 500 m-long transect that runs from the western edge of the site towards Rio Medio Queso. Our other two riverine study sites are located within a lagoon complex that is primarily fed by Ríos Frío and Mónico. The lagoon is part of the Caño Negro Wildlife Refuge and Ramsar site #541 (Ramsar, 2026), which is classified as a mixed-use protected wild area that is home to local communities who make sustainable use of the ecosystem through subsistence and cultural connections (Jiménez, 2018). The two study sites (San Jeronimo (SJ) and Laguna Martin (LM) are located within 1 km of one another; the vegetation communities are dominated by *Eleocharis equisetoides* and *Scleria microcarpa* (Jiménez, 2018). The SJ and LM sites were accessed by boat and the peat cores were extracted from the boat platform, as the coring sites were flooded by approximately 1m of water at the time of core collection (August 2023). We did not find any published soil information about Caño Negro.

(4) Montane peatbogs (n = 2) These sites are located in the Cordillera de Talamanca, Ramsar site #1286 (Ramsar, 2026), in central Costa Rica. In this high-elevation area (~ 3000 m.a.s.l.), peatlands tend to be very small in size (often < 0.05 km^2; Rabel et al. 2025). Their existence has been linked to the mountain range's Late Pleistocene glacial history, with montane peatlands forming in small, poorly drained depressions following the last deglaciation around 10,000 years ago (Horn, 1990). Today, the area is considered part of a subalpine paramo ecosystem, itself located within a cloud forest, with a cool and moist climate, and the small peatland-bearing depressions are prone to seasonal flooding (Jiménez, 2016). The two sites that we sampled, Bog 70 and Bog 68, were named after the km markers on the side of the road (Rodgers and Horn, 1996). Those sites are dominated by a unique mixture of peat moss (*Sphagnum* sp.), reindeer lichen (*Cladonia rangiferina*), and tropical plants such as tree ferns (*Blechnum auratum*), bromeliads (*Puya dasylirioides*), and numerous herbaceous species (see Rodgers and Horn (1996) for details). Sediment cores have been retrieved from these sites before and used for pollen-based paleoenvironmental reconstructions (Hooghiemstra et al., 1992; Islebe et al., 1995, 1996), but studies of the peat characteristics themselves were not conducted to our knowledge.

2.3 Field sampling strategy and approach

The work herein stems from an exploratory trip; we opted for visiting a larger number of sites while obtaining fewer data points for each site. The general approach that was used to sample sites included: (1) trace a transect line from the edge of a potential peatland towards its center, (2) walk (or boat) along said transects, and (3) probe the soil along the transects to confirm the presence (or absence) of peat. The length and orientation of those transects varied for each site and mainly depended on site conditions and time constraints. Peat depth and coordinates were recorded for each probing.

A 'full core' was retrieved from the deepest portion of four peatlands using a 50 cm long, 5 cm diameter Russian-style peat borer. A full core consists of the entire peat column, from the peatland surface to the peat-to-mineral interface. The multi-proxy analyses presented in this study were performed on those four cores. In addition to these full cores, several 'bottom cores' (n = 7) were

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collected along the transects from all 9 peatlands (Table 1). Those bottom cores consist of the deepest 50 cm long drive that was collected at the site and they include the peat-to-mineral interface. Those bottom sediment cores are primarily used to analyze patterns in peat basal ages along transects, which may help understand the history of peatland expansion across individual basins. In the field, all freshly collected cores were placed in PVC pipes and wrapped in plastic film and aluminum foil. The samples were kept at 4° C until laboratory analysis.

2.4 Laboratory sample preparation and analysis

In the laboratory, all peat cores were photographed and sliced into contiguous, 1 cm thick slices, which were then placed into labeled plastic bags and stored at 4° C. The following sections describe the methodological approach that was followed to prepare our samples for different proxy analyses. Although not every analysis was conducted at the same resolution (e.g., every 2 cm for LOI vs. every 8 cm for FTIR), most measurements were made on the same peat slices to allow for direct comparison across methods (see Table A1 for a list of proxies and analytical resolution).

2.4.1 Peat geochemistry (loss-on-ignition and elemental analysis)

The loss-on-ignition method was followed (Dean, 1974) to determine peat water content, dry bulk density, and organic matter content. Every 2 cm, a sub-sample (1 cm^3) was extracted from the peat slice, weighed, and sequentially oven dried to determine water content and dry bulk density (40° C, 48 h) as well as organic matter content (550° C, 4 h). These measurements were also used to calculate organic matter (Loisel et al., 2014).

For carbon and nitrogen content determinations, 1 cm^3 sub-samples were collected every 8 cm. Samples were oven dried (40°C) and ground into a fine powder using a Retsch MM 300 Vibration Mill Mixer (Verder Company, Germany). The homogenized samples were then placed into tin capsules. C and N content were measured using a Costech ECS 4010 Elemental Analyzer (Costech Analytical Technologies, Inc., Valencia, CA USA). Different sample weights were used on the basis of organic matter content, with smaller sample weights for organic-rich samples and vice versa. The carbon-to-nitrogen ratio (C/N) is presented as a mass ratio. The C/N ratio of peat can be used as a proxy for decomposition and humification (Biester et al., 2014; Kuhry and Vitt, 1996; Malmer and Holm, 1984). Organic carbon density was subsequently calculated for each sample by multiplying the organic matter density (from LOI data) with the measured C content. Finally, the regression between peat C% and peat OM% is presented as an estimate for organic carbon content that can be used in future studies that may lack elemental analysis.

2.4.2 Peat organic matter quality (FTIR spectroscopy)

FTIR spectroscopy was used to analyze peat organic matter quality (Artz et al., 2008; Holmgren and Nordén, 1988; Krumins et al., 2012; Uhelski et al., 2022). FTIR spectroscopy has been shown to be a useful proxy for humification index in a peat column (Biester et al., 2014; Broder et al., 2012; Tfaily et al., 2014; Upton et al., 2018). Sub-samples (1 cm^3) were collected every 8 cm along the four full cores. Samples were oven dried (40° C) and ground into a fine, homogenized powder using a Retsch MM 300 Vibration Mill Mixer. Spectra were acquired on a Thermo Nicolet 380 FTIR spectrometer (Thermo Electron Scientific Instruments Corporation, Madison, WI USA) by averaging 100 scans at 8 m^{-1} resolution over a range of 4000-550 cm^{-1}. Samples were placed directly on an attenuated total reflectance (ATR) crystal, and light force was applied so the samples came into direct contact with the crystal. ATR correction and atmospheric suppression took place to account for variability in background CO_2 and water vapor.

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A background scan was collected before every sample. The spectra were then manually baseline-corrected between 4000-3750 and 2600-1800 cm^{-1}. A custom R script (R Core Team, 2024) was used to determine the peak heights and relative abundances of carbohydrates, aromatics, acids, and aliphatics in predetermined regions of the spectra (Hodgkins et al., 2018; Fig. A1). The ratios of carbohydrates/aromatics, carbohydrates/aliphatics, and aromatics/aliphatics were also calculated.

2.4.3 Peat composition (plant macrofossils and von Post humification index)

Plant macrofossils were analyzed following standard procedures (Loisel et al., 2009; Mauquoy et al., 2002). Sub-samples (1 cm^3) were analyzed every 8 cm (or at higher resolution near stratigraphic boundaries) along each full core. Samples were mixed with deionized water and suspended in a petri dish, then scanned under a stereomicroscope. The first step involved the visual estimation of the percentage of mosses, herbaceous, ligneous, mineral, and unidentified plant material (UOM) (Mauquoy et al., 2010). A peat type category (bryophyte, herbaceous, ligneous) was assigned to each sample on the basis of the plant macrofossil analysis, where the dominant plant type found in each sample was used to infer the peat type (Loisel et al., 2014). We note that the 'ligneous' peat type was split into two sub-categories: ligneous-general vs. ligneous-mangrove to better represent our peat material. Lastly, we specified the presence of 'root material' both herbaceous and ligneous material when possible. We note that this differentiation is often challenging, given structural similarities between aerial vs. subterranean plant tissue. But in some cases, such as for mangrove roots, the distinction was safe to make.

Peat humification was determined at 8cm increments using the von Post humification scale (ASTM, 1987), a simple, widely used method to estimate the level of peat decomposition by pressing and manipulating a small sample of peat (~1 cm^3) in one's hands (Kaila, 1956; Rydin and Jeglum, 2013; Stanek and Silc, 1977).

2.4.4 Peat basal ages and chronologies (radiocarbon dates and age-depth models)

Peat inception ages were determined using radiocarbon (^{14}C) dating (Table 2). Along each core, the deepest (and presumably oldest) peat sample was identified using LOI results (see below). A 30% organic matter threshold was used to identify the peat-to-mineral boundary along our peat cores (Joosten and Clark, 2002; Lourenco et al., 2023). Given the humic nature of the peat, dated samples consisted of bulk, root-free samples (63-125 µm). Samples were wet-sieved using deionized water, oven dried at 40° C, and packaged in aluminum foil. Additional samples were extracted along the four full cores and similarly prepared for ^{14}C dating, to generate peat accumulation histories. Samples were shipped to Lawrence Livermore National Laboratory's Center for Accelerator Mass Spectrometry (CAMS), where they underwent acid-base-acid (ABA) treatment (Gillespie and Hedges, 1983; Hedgpeth et al., 2025; Norris et al., 2020). Dates were first individually calibrated to calendar age (cal. yr BP) using the Intcal20 calibration curve (Reimer et al., 2020) in CALIB v8.2 (Stuiver and Reimer, 1993). As for constraining peat chronologies of the full cores using multiple ^{14}C ages, a Bayesian age-depth modeling approach was used (package *rbacon*) v.3.5.2 in R (Blaauw and Christen, 2011).

Core name	Sample Depth (cm)	^{14}C Age (years BP)	Error (±)	Median Age (cal. yr BP)	Error (2 sigma) (±)	Laboratory ID (CAMS)
BOG70	21.5	>Modern*		-59	2	195269

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	100.5	8765	35	9759	173	193837
BOG68	80.5	9640	90	10967	248	193857
GAN-4	45.5	220	25	173	77	193839
	90.5	350	25	435	77.5	193838
	130.5	725	25	668	72.5	193836
GAN-SF	170.5	990	35	856	84.5	195199
AIR	140.5	450	25	507	23.5	193832
MAN	223.5	2380	30	2403	161.5	193858
	239.5	2595	25	2740	18	193835
RM	47.5	315	25	389	75	193842
	83.5	220	30	447	57.5	195272
	94.5	495	30	524	23.5	193859
	141.5	895	25	787	86.5	193841
	187.5	1135	25	1021	108.5	193840
MQ-S9	35.5	860	30	802	141.5	195271
	45.5	1785	25	1690	156.5	193845
	59.5	2700	35	2842	147.5	195200
	69.5	3625	35	3998	359	195201
	75.5	4555	35	5233	196	195202
	80.5	5065	30	5806	139	195271
	90.5	5305	25	6082	95.5	193844
	135.5	5885	25	6706	65	193843
	180.5	6855	35	7683	83.5	193860
MQ-S10	206.5	5690	25	6465	74.5	193829
	230.5	7535	30	8361	96	193830
LM	96.5	1840	25	1741	89	193833
SJ	208.5	1085	25	988	64	193831

Table 2: AMS radiocarbon dating results from the new peat cores in Costa Rica. *The modern age for sample CAMS-195269 returned a fraction modern of 1.0520 ± 0.0041.

2.5 Statistical Analysis

First, an analysis of variance (ANOVA) was used to test the effect of 'peat type' on the peat geochemical properties (Loisel et al. 2014). Second, a correlation matrix was completed in R with the packages *energy, corrplot, dplyr,* and *ggplot2* to compare the relationship between peat proxies (Rizzo and Szekely, 2024; Wei and Simko, 2024; Wickham, 2016; Wickham et al., 2023). To account for possible non-linear trends in the data, we opted for a Spearman correlation (Zar, 2005). Third, a Principal Coordinates Analysis (PCoA) based on the Bray-Curtis distance was used to capture variations in the peat properties across peat types and sites, and over time (Tian et al., 2019). The analysis was performed in R with the *vegan* package (Oksanen et al., 2025). Bray-Curtis distances were chosen to understand differences in ecological compositions while also handling the differences in relative abundances gathered from the FTIR analysis (Kellerman et al., 2015; Wang et al., 2022). Lastly, PERMANOVA and Distance-based redundancy (dbRDA) analyses were performed in conjunction with the PCoA to statistically test differences between sites (Anderson, 2001; Legendre and Anderson, 1999; Tian et al., 2019).

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3 Results

3.1 Peat accumulation histories across four Costa Rican sites

3.1.1 Coastal palm swamp site (Gandoca - GAN)

The GAN core is characterized by 130 cm of peat that is underlain by fine mineral particles. Peatland initiation began at 668 cal. yr BP and the plant macrofossil record, which is dominated by ligneous palm remnants, suggests that a palm swamp community has likely occupied this site since peat inception (Fig. 2). The peat humification index was consistently high, with slightly more humified peat towards the base of the peat column. The apparent rate of peat accumulation is steady and relatively rapid (0.2 cm/yr), and the age-depth relationship follows a linear trend. The peat stratigraphy is high in organic matter throughout the profile (mean = 70.2%). Expectedly, bulk density tends to be relatively low (mean = 0.164 g/cm^3), with a sharp peak that corresponds to the abovementioned mineral interval. As for peat quality, the GAN core is characterized by low variability in its abundance of carbohydrate, aromatic, acid, and aliphatic compounds (Fig. 2 and A2), with the exception of near-surface samples (16 and 24 cm) and the mineral transition at the end of the peat (136 cm), where peaks associated with greater amounts of carbohydrates (in combination with lower aromatics and aliphatics) were identified.

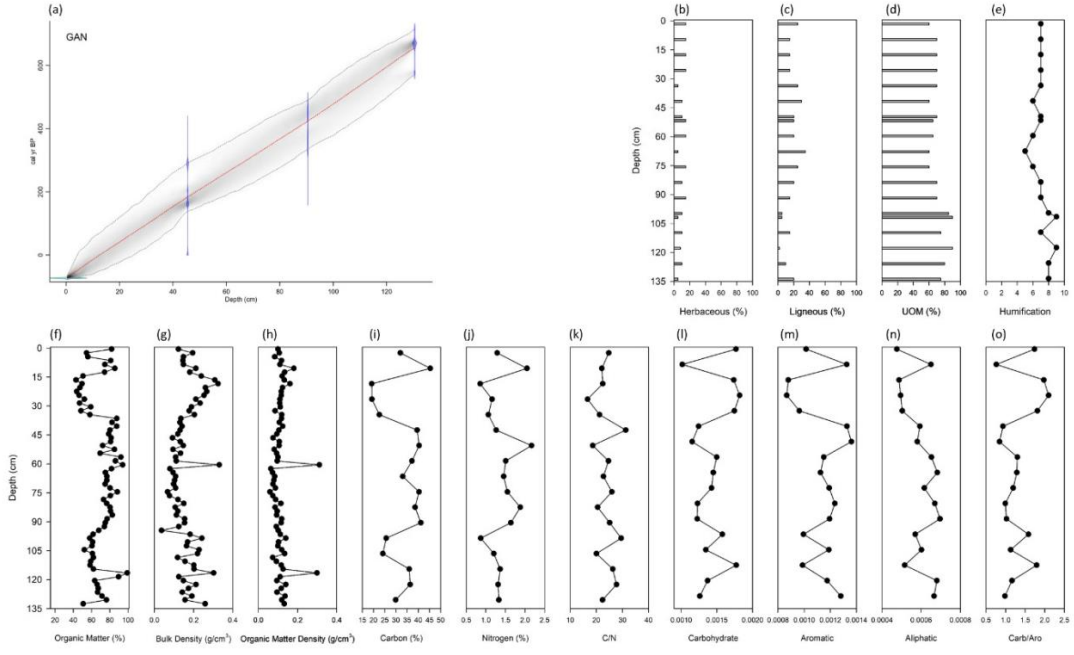

Figure 2. Accumulation history, macrofossil composition, geochemical, and FTIR results for the coastal palm swamp (GAN) site. (a) Age-depth model, (b) herbaceous macrofossil composition (%), (c) ligneous macrofossil composition (%), (d) unidentified organic matter

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(UOM) composition (%), (e) von Post humification, (f) organic matter content (%), (g) bulk density (g/cm^3), (h) organic matter density (g/cm^3), (i) carbon content (%), (j) nitrogen content (%), (k) carbon/nitrogen ratio, (l) carbohydrate abundance, (m) aromatic abundance, (n) aliphatic abundance, and (o) carbohydrate/aromatic ratio.

3.1.2 Mangrove site (Red mangrove - RM)

The RM core is 187 cm-long and primarily composed of organic-rich material, which is why this site is included in our peat study. As a point of reference, other mangrove studies in the region have described these ecosystems as mineral-rich, with 5-8% OM (Rodgers and Horn, 1996). In contrast, our RM site harbors OM% in the order of 25-71%. In addition, the record begins as a palm swamp that initiated around 1020 cal. yr BP and persisted until 465 cal. yr BP (Fig. 3). The underlying sediment consists of sandy material, and the stratigraphy is interrupted by one mineral-rich layer around ~740-620 cal. yr BP (110-134 cm). Around 465 cal. yr BP (83 cm), the stratigraphy abruptly shifts to a dominance of *Rhizophora* plant remnants (mangrove roots). We caution that, while the shift from palm swamp to mangrove seems to take place at 465 cal. yr BP according to our Bayesian age-depth model, said model also rejected a ^{14}C date from a sample that was picked at the ecological transition (83.5cm) but that was deemed 'too young' to fit the rest of the curve (^{14}C age: 220 ± 30, which yields a median age of 189 cal. yr BP with a 2-sigma range of 0-310 cal. yr BP; Fig. 3). This age reversal, if representative of the sediment history, would suggest that the 'mangrove root deposit' is in fact much more recent than indicated by the age-depth relationship, possibly as young as a few decades only.

The peat humification index is generally mesic to humic, with the exception of relatively undecomposed peat following the transition from palm swamp to mangrove. The apparent rate of peat/sediment accumulation is steady and relatively rapid (0.2 cm/yr; similar to the nearby GAN core), and the age-depth relationship follows a linear trend - with the caveat of a possible hiatus at the transition from palm swamp to mangrove (83 cm). The peat stratigraphy is characterized by moderate organic matter content (mean = 43.83%), the lowest average of all our sites. The RM results are characterized by wide changes between OM-rich (mean = 47.45%) and mineral-rich (mean = 18.80%) intervals, generally in a reverse relationship with bulk density. Those changes in organic matter content and bulk density follow a ~ 200-yr cycle (Fig. 3). As for peat quality, the spectroscopic dataset is characterized by very high peaks within the carbohydrate region of the spectra contrasted by low aromatics, acids, and aliphatics throughout the core (Fig. 3 and A3). The mineral-rich layer is expressed as flattened spectra (112-128 cm). We note that the aromatic and aliphatic abundances tend to correlate with %C and %OM, with greater compound abundances corresponding to OM- and C-rich material. However, some of the carbohydrate peaks correspond to intervals with high bulk density, which is in opposition with the hypothesized negative relationship between those two proxies. Indeed, the expectation is to see high carbohydrates correlating with low bulk density, both representing a low degree of decomposition. We offer an explanation of this incongruent result in sect. 3.1.3 and 3.2.1.

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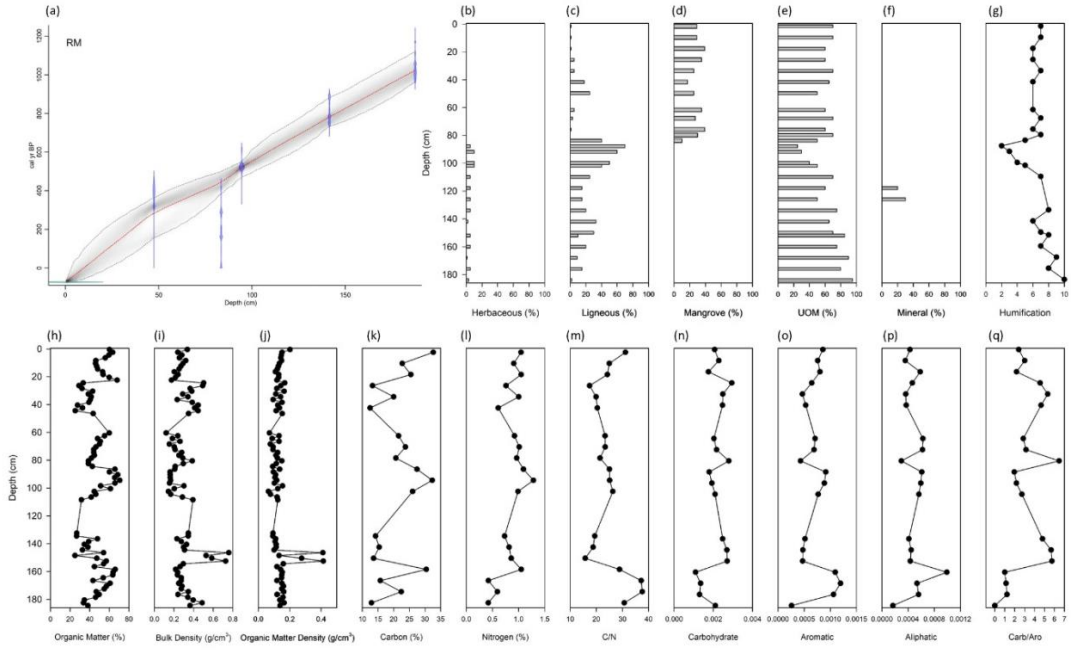

Figure 3. Accumulation history, macrofossil composition, geochemical, and FTIR results for the mangrove (RM) site. (a) Age-depth model, (b) herbaceous macrofossil composition (%), (c) ligneous macrofossil composition(%), (d) ligneous mangrove macrofossil composition (%), (e) unidentified organic matter (UOM) composition (%), (f) mineral composition (%), (g) von Post humification, (h) organic matter content (%), (i) bulk density (g/cm^3), (j) organic matter density (g/cm^3), (k) carbon content (%), (l) nitrogen content (%), (m) carbon/nitrogen ratio, (n) carbohydrate abundance, (o) aromatic abundance, (p) aliphatic abundance, and (q) carbohydrate/aromatic ratio.

3.1.3 Riverine site (Medio Queso - MQ)

The MQ core presents 180 cm of peat material with an inception age of 7683 cal. yr BP. Initially, the site was likely a palm swamp (Fig. 4), as shown by the presence of ligneous palm fragments akin to those found at our coastal (GAN) and mangrove (RM) sites. The palm swamp ecosystem persisted for at least two millennia (until ~ 5500 cal. yr BP), but perhaps for a longer time interval. A hiatus from ~ 5500 to ~ 2500 cal. yr BP (75-55 cm) challenges our ability to determine the timing of the transition into the modern-day herbaceous peatland. The apparent long-term rate of peat accumulation for the palm swamp (7683-5500 cal. yr BP) was 0.05 cm/yr, which is an order of magnitude slower than our observations at GAN and RM. The peat swamp portion of the profile is characterized by relatively high OM% (> 50%) and low BD (< 0.3 g/cm^3), with the exception of two layers with ~ 30% OM and densities of ~0.5 g/cm^3. Those two layers are also associated with very high peaks within the carbohydrate region, which we posit are overprinted by silicated minerals (Fig. A4). Indeed, the MQ core exhibits a split peak in the carbohydrate region of the spectra, most likely attributed to both organic matter carbohydrates (1030 cm^{-1}) and silicate minerals (1005 cm^{-1}) (Reig et al., 2002). This interpretation is corroborated by the presence of bands at 912, 3620, 3695 cm^{-1} that are likely linked to the presence of silicates

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and kaolinite clays (Müller et al., 2014; Seaton et al., 2024; Sengyang et al., 2015). Repeated flooding may have increased clay content, confounding the carbohydrate signal, thereby also explaining why our high carbohydrate abundances are synchronous with high bulk density values, which is also a phenomenon that we observed at our mangrove site (RM; see sect. 3.1.2).

There is then a long-time interval (from 5500 to ~ 2500 cal. yr BP; from 75 to 55 cm) during which the record may have been altered, with a two-fold slowdown in apparent rate of peat accumulation (0.007 cm/yr). This interval coincides with high bulk density values (> 0.4 g/cm^3), humic peat (10 on the von Post scale), and peaks in aliphatics that suggest the presence of herbaceous peat. Our field notes also indicate the presence of thin mineral layers throughout this part of the profile. The uppermost peat section (top 55 cm) corresponds to the past 2500 years and is composed of herbaceous material with an averaged apparent rate of peat accumulation of 0.02 cm/yr. This herbaceous section is characterized by noticeably lower C/N values and the presence of herbaceous plant remnants (up to 20%). We note that the uppermost section of the record was compressed during the coring process due to the presence of surface water (0-18cm), making it difficult to assess changes in peat properties in the near-surface samples. Nevertheless, OM% is highest in the uppermost peat layers (60-95%) and corresponds to low bulk density values (< 0.2 g/cm^3), despite the peat being already very humified near the top of the core (von Post = 8).

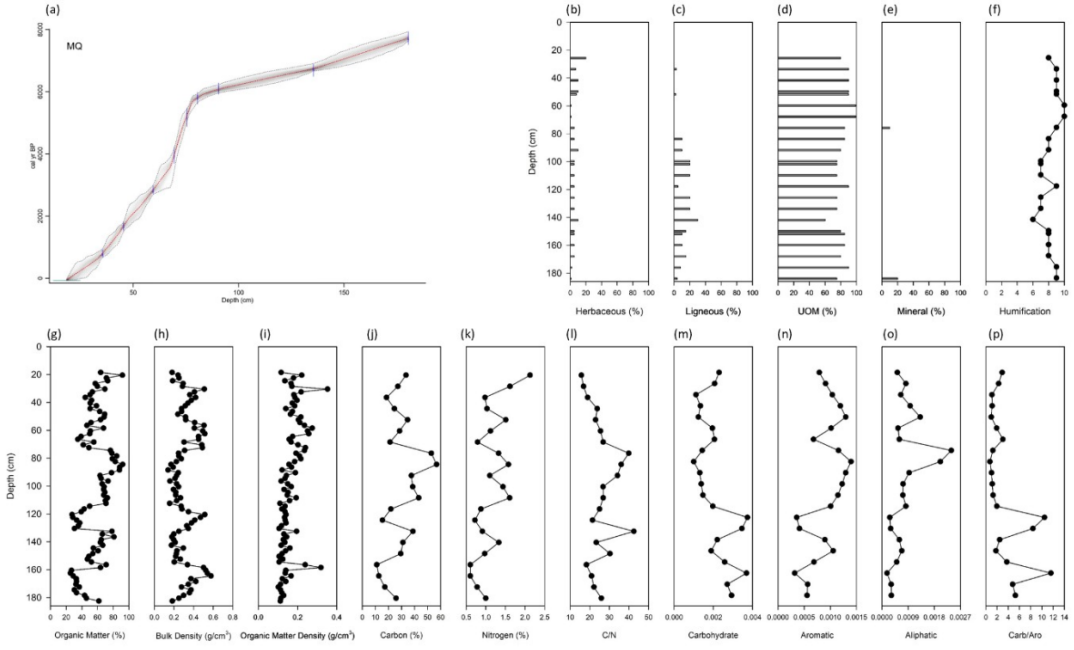

Figure 4. Accumulation history, macrofossil composition, geochemical, and FTIR results for the riverine (MQ) site. (a) Age-depth model, (b) herbaceous macrofossil composition (%), (c) ligneous macrofossil composition (%), (d) unidentified organic matter (UOM) composition (%), (e) mineral composition (%), (f) von Post humification, (g) organic matter content (%), (h) bulk density (g/cm^3), (i)

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organic matter density (g/cm^3), (j) carbon content (%), (k) nitrogen content (%), (l) carbon/nitrogen ratio, (m) carbohydrate abundance, (n) aromatic abundance, (o) aliphatic abundance, and (p) carbohydrate/aromatic ratio.

3.1.4 Montane site (BOG70)

The BOG70 core is our shortest but longest record, at 100 cm and with an initiation age of 9759 cal. yr BP. The ecosystem appears to have started as a herbaceous-dominated peatland (Fig. 5), atop clayey sediment, though the peat is highly humified, which limits the use of plant macrofossil analysis. Ligneous fragments were found starting around 1823 cal. yr BP (42 cm). *Sphagnum* and Amblystegiaceae remnants appear in the record at -65 cal. yr BP (18.5 cm) and form the modern-day moss carpet. The apparent long-term rate of peat accumulation was the slowest of all 4 study sites at 0.01 cm/yr, which is 15-20 times lower than the coastal palm sites. Peat humification is high along the entire record though it trends towards lower humification towards the top of the profile. Organic matter content is high throughout the majority of the core (mean = 86.25%); this montane site in fact presents the highest organic matter content of all 4 study sites. Those results are corroborated by a prior study that reported high organic matter content (60-96%) for these montane sites (Rodgers and Horn, 1996). The FTIR data show that the record is characterized by low carbohydrates relative to aromatic and aliphatic components (Fig. 5 and A5). Notably, high aliphatic abundance appears contemporaneous to the herbaceous peat. There is also a unique signature at the core surface, where the two aliphatic C-H bands are less resolved than in all the other samples (Fig. A5, top sample). This feature is a hallmark of fresh *Sphagnum* (Heller et al., 2015). At the base of the peat column (105 cm), an increase in clay content is inferred from spectral features at 3620 and 3695 cm^{-1}.

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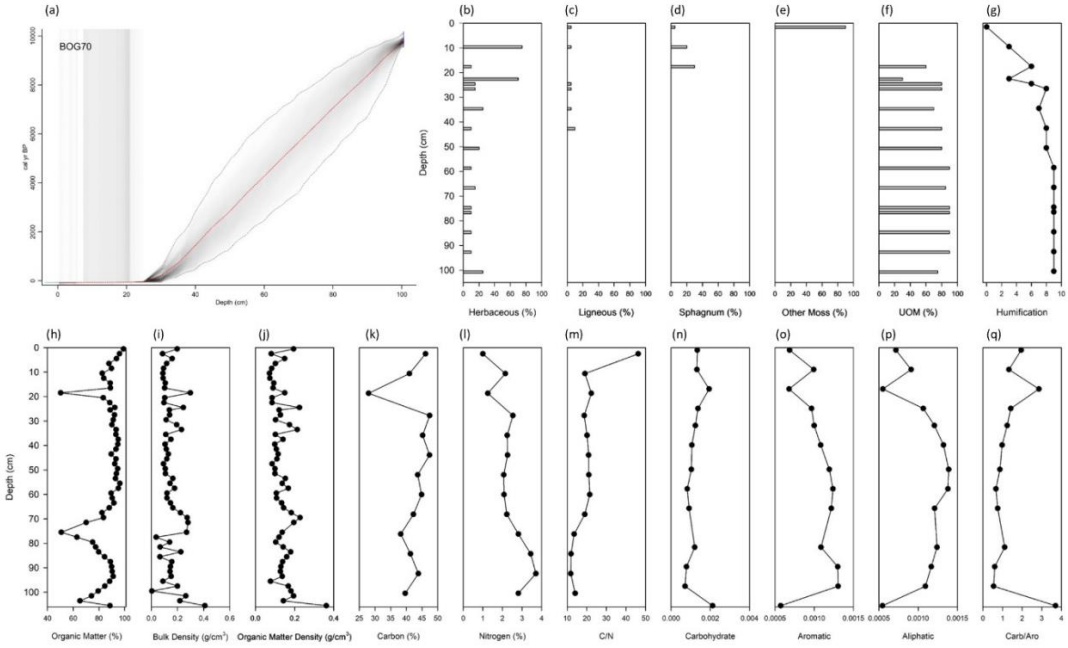

Figure 5. Accumulation history, macrofossil composition, geochemical, and FTIR results for the montane (BOG70) site. (a) Age-depth model, (b) herbaceous macrofossil composition (%), (c) ligneous macrofossil composition (%), (d) *Sphagnum* macrofossil composition (%), (e) other (non-*Sphagnum*) moss macrofossil composition(%), (f) unidentified organic matter (UOM) composition(%), (g) von Post humification, (h) organic matter content (%), (i) bulk density (g/cm^3), (j) organic matter density (g/cm^3), (k) carbon content (%), (l) nitrogen content (%), (m) carbon/nitrogen ratio, (n) carbohydrate abundance, (o) aromatic abundance, (p) aliphatic abundance, and (q) carbohydrate/aromatic ratio.

3.2 Spatiotemporal patterns in peat properties

3.2.1 Peat organic matter quality (FTIR data)

Correlations between functional groups were found across all sites (Fig. 6 and A6), with carbohydrates negatively correlated with aromatics (Spearman = -0.881, $p < 0.0001$), aliphatics (Spearman = -0.794, $p < 0.0001$), and acids (Spearman = -0.679, $p < 0.0001$). We also note that aromatics, and especially aliphatics, correspond with %C and %OM, confirming these spectral features as proxies for stable organics (Fig. A6). All those associations are stronger at the site level (Fig. 6), though downcore trends are complex. For instance, the coastal swamp site (GAN) and the montane site (BOG70) both show progressive decreases in carbohydrates with depth inverse to the increases in aromatic and aliphatic abundances. This pattern is expected, as carbohydrates represent comparatively labile organic matter, whereas aromatic and aliphatic moieties tend to show greater long-term stability and are thought to 'accumulate' in older, more refractory peat (Leri et al., 2025; Verbeke et al., 2022). We note that aliphatic abundances are greater at the montane, herbaceous-dominated site (BOG70) than in the lowland, ligneous-dominated swamp (GAN). As for

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the riverine (MQ) and mangrove (RM) sites, large variations along the core do not follow the expected increase in recalcitrant fractions with depth.

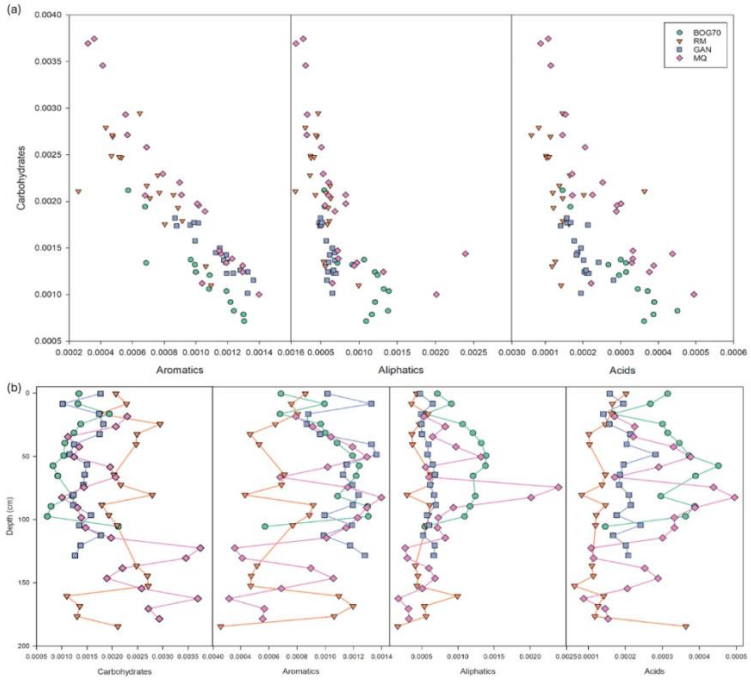

Figure 6. FTIR data from our four Costa Rican peatlands. (a) the relative abundances of carbohydrates are plotted against aromatics, aliphatics, and acids, (b) changes in carbohydrates, aromatics, aliphatics, and acids are shown as time series (i.e., along peat core depth).

Average spectra from all of the depths in each core were compared (Fig. 7) in an attempt to detect site-specific signatures. The presence of distinct dual aliphatic C-H features (2850 and 2920 cm^{-1}) across all sites indicate high organic matter content and the presence of peat (Martínez Cortizas et al., 2021). As mentioned above, the coastal peat swamp (GAN) and the montane site (BOG70) exhibit similarities in terms of their averaged carbohydrate and aromatic content, though the herbaceous site (BOG70) is characterized by greater aliphatic content. Conversely, the mangrove (RM) and riverine (MQ) sites seem characterized by very high carbohydrate peaks (1030 cm^{-1}), but a closer look at the wavenumbers reveal a probable influence by silicate compounds (1005 and 912 cm^{-1}), which is confirmed by the presence of kaolinite peaks (3620 and 3695 cm^{-1}). Given that the mangrove and riverine sites are prone to flooding, it is perhaps not surprising to see this overprint by mineral compounds, but those peaks obscure the carbohydrate signature (Verbeke et al., 2022). The mangrove site also presented low aromatic (1510 and 1630 cm^{-1}) and aliphatic peaks, which can be interpreted as indicating a lower degree of humification when compared to the coastal swamp and the montane peatbog.

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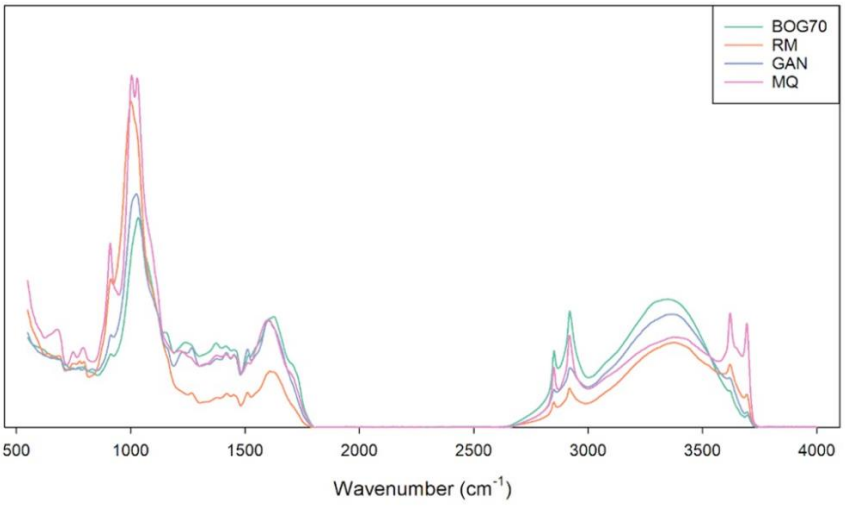

Figure 7. Average FTIR spectrum from our four full cores from Costa Rica.

3.2.2 Differences in peat composition and properties across sites

Peat composition and properties differ among the four sites (Fig. 8 and A7; Table A2), likely due to differences in organic matter quality. The first PCoA axis, explaining 79.8% of the observed variation, places %C as well as the aromatics, aliphatics, and acids on the left side of the ordination vs. the carbohydrates and BD on the right side of the ordination, as expected. The second axis (12%) was not strongly linked to any environmental variables. The PCoA plot showed that the peat properties from most samples from the montane peatbog (BOG70) and the coastal peat swamp (GAN) sites can be distinguished from those of the other two sites. In contrast, the mangrove (RM) and riverine (MQ) sites, regardless of peat type, depth, or age (Fig. A7), were scattered across the horizontal axis. Lastly, the PCoA plot that has the samples color-coded by depth (Fig. A7) shows a cluster of the shallower samples towards the left-hand side of the statistical space, suggesting the role of age on peat properties.

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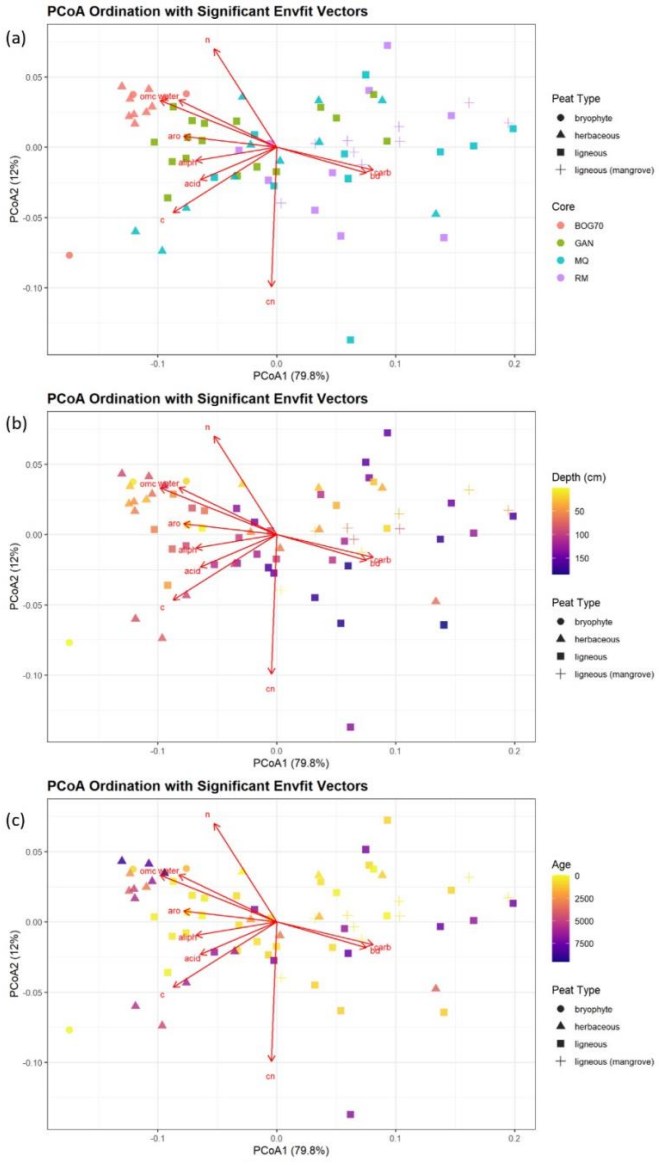

Figure 8. Principal Coordinates Analysis (PCoA) using Bray-Curtis distances overlaid with envfit vectors. The data in the ordination are visualized by (a) peat type and core, (b) peat type and depth, and (c) peat type and age.

We note that bulk density and carbohydrates seem to covary (Fig. 8, on the right-hand side of axis 1). The expectation is that carbohydrate relative abundance would be negatively correlated with bulk density, but the ordination instead suggests a strong positive relationship between those two variables. We posit that this unexpected result may reflect the contribution of Si-O bands from clay minerals in the C-O region at ~1030 cm^{-1}, confounding carbohydrates with silicates in the spectra (Fig. 7; Fig. A3 and A4). With mineral contributions enhancing the signals of both carbohydrate relative abundance and bulk density, those parameters may co-vary in mineral-rich samples.

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Subsequent PERMANOVA analysis (Table A3) revealed differences between sites (R^2 = 0.434, $F_{3,65}$ = 16.64, $p \leq 0.001$) and among peat types (R^2 = 0.282, $F_{3,65}$ = 8.52, $p \leq 0.001$), explaining 43.4 and 28.2% of the variation in the Bray-Curtis ordination, respectively. When tested together, site and peat type explained 50.1% of the variation (R^2 = 0.501, $F_{6,62}$ = 10.38, $p \leq 0.001$). The homogeneity of dispersion (betadisper), which shows the internal variation within each group, was determined for both sites ($F_{3,65}$ = 5.21, p = 0.003) and peat types ($F_{3,65}$ = 1.63, p = 0.19, Fig. A8). Dispersion was shown to be significantly different across sites, with the mangrove (RM) and the riverine (MQ) sites displaying greater heterogeneity, but not significantly different among peat types.

Lastly, a distance-based redundancy analysis (dbRDA) constrained by site and peat type was run to constrain the community variables (Table A3). Site and peat type together ($F_{6,62}$ = 8.253, $p \leq 0.001$) explained 44.4% of total dispersion in the Bray-Curtis ordination. The majority of the variation was explained by the sites themselves ($F_{3,62}$ = 14.02, $p \leq 0.001$), with 'peat types' adding explained variance ($F_{3,62}$ = 2.49, p = 0.024). A final pairwise PERMANOVA was run to determine which sites (Table A4) and peat types (Table A5) differed specifically from one another. Among the sites, all pairwise comparisons were statistically different from one another, except for MQ and RM (R^2 = 0.071, $F_{1,38}$ = 2.912, p = 0.066). The strongest difference was observed between BOG70 and RM (R^2 = 0.667, $F_{1,29}$ = 58.764, p = 0.0015). Among peat types, all comparisons were statistically different, except for bryophyte and herbaceous peats (R^2 = 0.103, $F_{1,21}$ = 2.402, p = 0.092). Ligneous and ligneous (mangrove) showed strong differentiation from bryophyte (Ligneous: R^2 = 0.153, $F_{1,38}$ = 6.908, p = 0.004; Ligneous (mangrove): R^2 = 0.687, $F_{1,10}$ = 21.976, p = 0.009) and herbaceous peats (Ligneous: R^2 = 0.144, $F_{1,55}$ = 9.275, p = 0.004; Ligneous (mangrove): R^2 = 0.412, $F_{1,27}$ = 18.924, p = 0.004). Ligneous and ligneous (mangrove) did display significant differences from one another (R^2 = 0.090, $F_{1,44}$ = 4.343, p = 0.025), but albeit at a lesser significance than bryophyte and herbaceous.

4 Discussion

4.1 Peat inception ages and age-depth relationships of Costa Rican peatlands

The timing of peatland inception has been used in many studies worldwide to help constrain changes in hydroclimatic conditions (Dommain et al., 2014; Thomas et al., 2025; Yu et al., 2016). In the case of Costa Rica, the dataset shows a wide range of peat inception age, with no particular time period during which peats primarily initiated. Peat inception ages range from 507 to 10,967 cal. yr BP (Fig. 9) and tend to cluster on the basis of peatland type / geomorphological setting. The montane peatlands exhibited the oldest ages, with BOG68 dating back to 10,967 ± 248 cal. yr BP. Conversely, the youngest peatlands were observed along the Caribbean coast, with the coastal palm swamp near Limon (AIR site) only dating back to 507 ± 24 cal. yr BP. The inland riverine sites encompass a wider age range, between 988 ± 64 and 8361 ± 96 cal. yr BP. A histogram combining all peat basal ages from this study suggests that most peatlands in Costa Rica are very young (less than 2000 years old), but this finding is partly due to a sampling artefact, as most of the sites are located in the coastal area. We have not found peats of pre-Holocene age in Costa Rica, a common theme across Latin America and the Caribbean (e.g., Lawson et al., 2026). Overall, the spatiotemporal pattern of peat inception primarily relates to elevation (high-elevation sites are the oldest) and proximity to the coast (coastal sites are the youngest).

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While the age of the montane peatlands was likely constrained by deglacial processes (Horn, 1990), the relatively recent inception and expansion of peatlands along the Caribbean coast (~1000 cal. yr BP) was probably linked to increased stability in base level, which combines stable sea level conditions as well as lowered variability in freshwater and sediment inputs from the inland rivers that feed the coastal lowlands. Regional sea-level stabilization is thought to have taken place around 6000 cal. yrs BP (Castañeda-Posadas et al., 2022), suggesting that other local factors might have prevented peatland establishment or development along the coast for many thousands of years, such as dynamic floodplains, earthquakes, erosion caused by storms, etc. As for the riverine sites, our limited sampling does not allow us to generalize, but it seems that our three sites (MQ, SJ, LM) lay within a lowland area that used to be flooded by Lago Nicaragua (also known as Lago Cocibolca) (Fig. 3; Bergoeing and Protti, 2006). Holocene fluctuations in the lake level, which could have been caused by a changing climate as well as tectonic activity (Kutterolf et al., 2023; Slate et al., 2013), have likely caused large shifts in the hydrological regime of the studied wetland complexes that span hundreds of km^2 in the area. Alas, the general lack of paleodata from the region makes it difficult to further address the cause and timing for peatland inception, though a 5700-year-old record from Lago Nicaragua reveals a potential impact of pre-colombian agriculture (5400 cal. yr BP) on the region as well as a general drying trend throughout the mid- to late-Holocene (Slate et al., 2013).

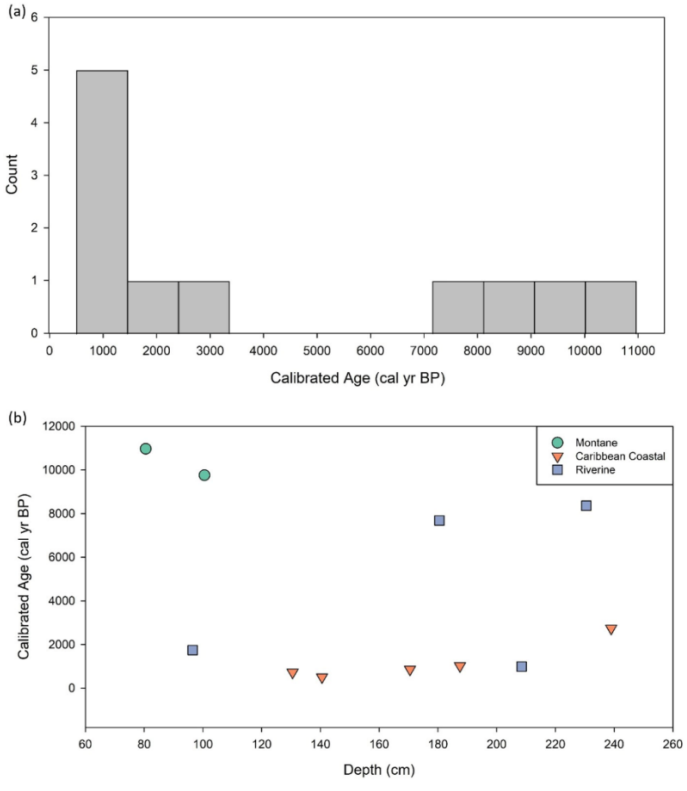

Figure 9. Peat inception ages from Costa Rican peatlands. (a) Peat initiation frequency distribution, presented in 1000-yr bins, (b) peat basal age vs. depth. The ecosystem types are displayed using different symbols and colors.

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Two of our study sites combine a full core and a bottom core, allowing us to provide some preliminary information on the spatio-temporal pattern of peatland expansion in Costa Rican peat basins. In the coastal site Gandoca (GAN), two cores were retrieved towards opposite ends of a 1 km long transect that runs perpendicular to the ocean. The full core (GAN) is located close to the beach and is characterized by 130 cm of peat that is aged at 668 cal. yr BP. Near the other end of the transect is core GAN-SF, with 170 cm of peat that dates back to 863 cal. yr BP. Those results point to a somewhat homogeneous back-barrier depression that could have filled up more or less synchronously. This preliminary finding is supported by our probe peat depth data points along the GAN transect (n = 10), which are all between 130 and 170 cm. Interestingly, we found a similar depth pattern at the AIR site, with peat depth values around 140 cm (n = 7).

In riverine wetland complex Medio Queso (MQ), 2 cores were also collected along a 600 m long, west-east transect from the edge of the site towards Rio Medio Queso. The full core (MQ; depth = 180 cm, age = 7683 cal. yr BP) and the bottom core (MQ-10; depth = 230 cm; age = 8361 cal. yr BP) are located approximately 100 m apart, with MQ-10 closer to the river. At this site, deeper (and likely older) peats are found along the transect, as corroborated by the probing depths (n = 10), which start at 40 cm near the edge of the site and reach 230 cm near the river, suggesting a progressive infilling of a local depression bordering the river.

Long-term apparent rates of peat accumulation also vary across peatland types and geomorphological settings (see age-depth models on Fig. 2-5). The nearly 10,000-year-old, but shallower (100 cm deep) montane site (BOG70) was characterized by an average apparent rate of peat accumulation (0.01 cm/yr) that is an order of magnitude slower than its coastal lowland counterparts. Indeed, the coastal peat swamp site (GAN) accumulated 130 cm of peat in only about 700 years; similarly, the mangrove site (RM) accreted 185 cm of peat in about 1000 years. The nature of the peat deposits differs across these sites: herbaceous at BOG70 vs. ligneous at GAN and RM, which might factor in the variations reported above. The differences in climatic regimes between the cool, high-altitude montane sites vs. the warm, lowland sites could also explain the reported differences in peat accumulation rates, with the cooler conditions acting as a limit to plant productivity. A third, and non-exclusive, process at play is that of long-term peat accumulation itself: the oldest peat (montane site) has had much more time to decompose, albeit slowly, than the material found in at the younger sites. With all that said and despite these large differences in accumulation rate, the organic carbon density was similar across all sites (mean = 0.04 gC/cm^3). The stratigraphies were also all characterized by large amounts of recalcitrant compounds (aromatics and aliphatics), suggesting that the peat carbon store is stable and thus somewhat protected from further mineralization, regardless of age and peat type.

4.2 Geochemical characterization of Costa Rican peatlands

Combining all the loss-on-ignition measurements (n = 279) and elemental analysis data (n = 70) across our four cores provides an overview of peat water content, dry bulk density, organic matter content, organic matter density, carbon and nitrogen content, and C/N ratio for Costa Rican peatlands. Table 3 and Fig. 10 present the data ranges as well as the mean values and standard deviation for all those variables. One-way analyses of variance (ANOVA) revealed that all peat types were statistically different from each other in the case of water content ($F_{2,276} = 11.94$, $p < 0.0001$) and organic matter content ($F_{2,276} = 44.66$, $p < 0.0001$). In the case of dry bulk density, the ANOVA revealed an overall effect of peat type ($F_{2,276} = 4.95$, $p < 0.01$), with the bryophyte type significantly different (Tukey's LSD: $p < 0.001$) from herbaceous and ligneous peats. Lastly, for organic matter density, we also report an

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overall effect of peat type ($F_{2,276}$ = 18.38, $p < 0.0001$), with the herbaceous peat type significantly different (Tukey's LSD: $p < 0.001$) from the bryophyte and ligneous peat types. Statistical tests were not applied to %C, %N, and C/N, given the lower number of samples available. We also note that the correlation between %C and %OM in the Costa Rican samples was strong and statistically significant (Spearman correlation = 0.847; $p < 0.0001$, Fig. 11). The slope (0.51) of the linear regression indicates that %C is about half the OM content, similar as the relationship found in the northern peatland database for non-*Sphagnum* peat (0.514; Loisel et al. 2014), supporting the idea that OM content and %C are closely related in peatlands across biomes, despite the large differences in their relative geochemical characteristics.

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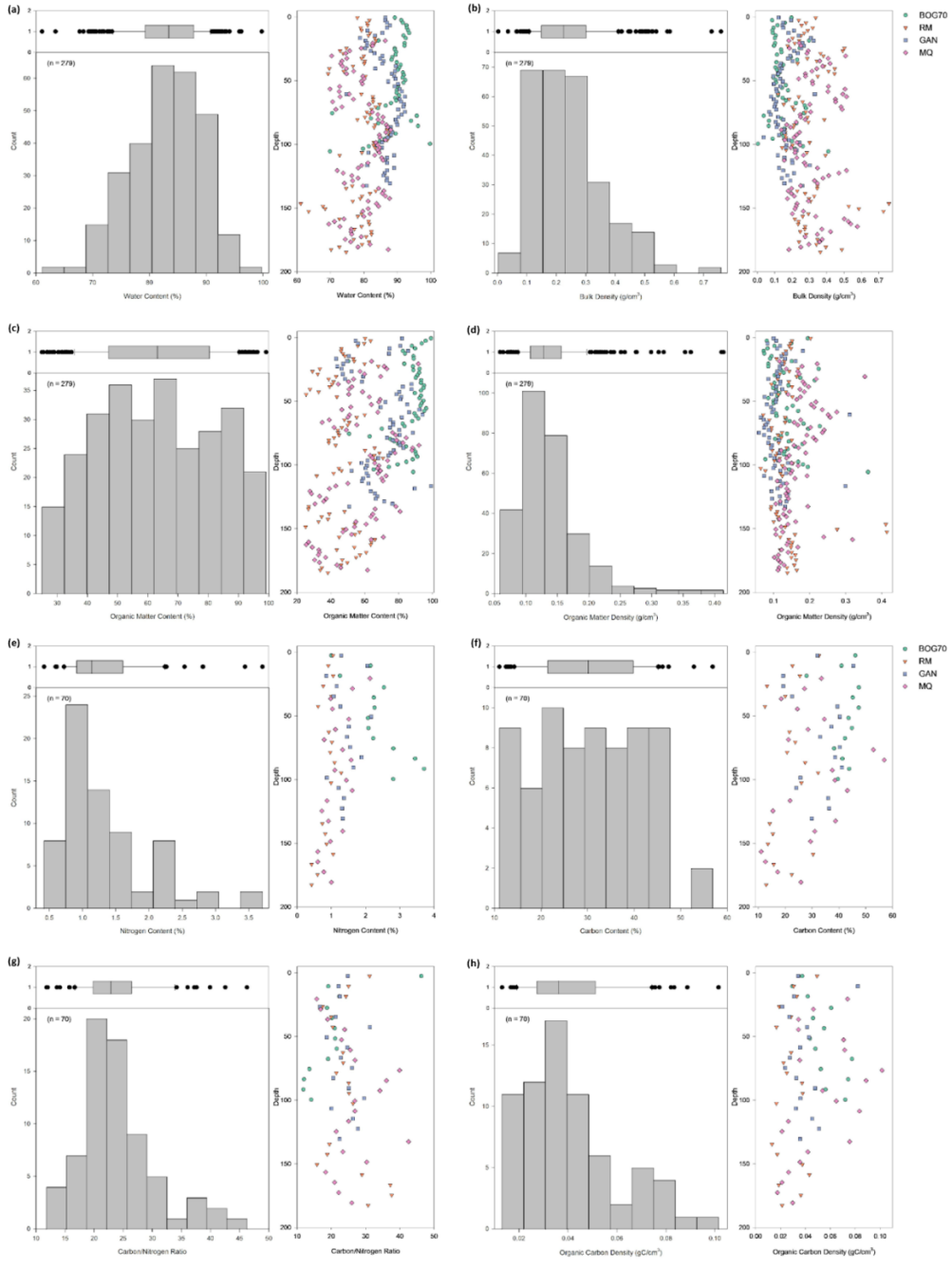

Figure 10: Frequency distribution histograms and downcore profiles of peat properties in Costa Rican peatlands. (a) Water content (%), (b) bulk density (g/cm^3), (c) organic matter content (%), (d) organic matter density (g/cm^3), (e) nitrogen content (%), (f) carbon content (%), (g) carbon/nitrogen ratio, and (h) organic carbon density (gC/cm^3).

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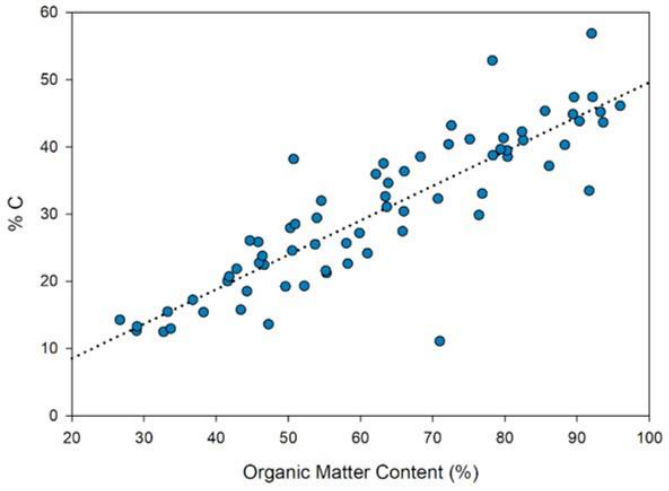

Figure 11: Relationship between organic matter content and carbon content in Costa Rican peat.

	Mean	Variance (σ^2)	Minimum	Maximum
Water Content (%)	82.90	41.74	61.07	99.77
Bulk Density (g/cm^3)	0.242	0.015	0.003	0.759
Organic Matter Content (%)	63.10	383.05	24.99	99.14
Organic Matter Density (g/cm^3)	0.139	0.003	0.059	0.414
Nitrogen Content (%)	1.358	0.448	0.421	3.705
Carbon Content (%)	30.463	125.145	11.065	56.870
Carbon/Nitrogen Ratio	23.983	44.968	11.830	46.466
Organic Carbon Density (gC/cm^3)	0.0418	0.0004	0.0131	0.1014

Table 3: Descriptive statistics for Costa Rican peat geochemical properties.

Combining the geochemical data together by peat type (bryophyte, ligneous, herbaceous) provides the first dataset of its kind for Costa Rica (and the Caribbean more broadly), with direct usability for regional carbon analysis and modeling (Fig. 12). Bryophyte peat (n = 10, only coming from the montane site) is characterized by high organic matter content (86 ± 14%), the highest of all peat types. This result is approximately 10% lower than, though still in line with, results from the northern hemisphere peatland database (94 ± 9%; Loisel et al. 2014). As for bulk density and organic matter density, the Costa Rican bryophyte peats are about 60% more dense than their northern counterparts (BD: 0.13 ± 0.07 and OMD 0.11 ± 0.04 g/cm^3 in Costa Rica vs. 0.08 ± 0.04 and

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0.07 ± 0.03 g/cm^3). Carbon and nitrogen content also differed between these populations: %C was lower (38 ± 9%) than in the northern peats (46 ± 4%) whereas %N was about twice as high (1.5 ± 0.6% vs. 0.7 ± 0.3%). Herbaceous peat (n = 84, with data coming from our montane and riverine sites) yielded moderately high organic matter content (75 ± 17%). Compared to northern herbaceous peats, Costa Rica is approximately 10% lower (86 ± 15%). Costa Rican peats are about 50% more dense than their northern counterparts, in terms of both bulk density and organic matter density (BD: 0.23 ± 12 and OMD: 0.15 ± 0.05 g/cm^3 in Costa Rica vs. 0.12 ± 0.08 and 0.09 ± 0.04). %C in herbaceous Costa Rican peats were lower (38.5 ± 10.3%) than in northern peats (50.5 ± 4.9%); %N was only slightly higher in Costa Rica (2 ± 0.8%) than in the north (1.7 ± 0.6). Ligneous peat (n = 185, with data from the riverine, coastal swamp, and mangrove) yielded around 30% lower organic matter content (56 ± 17%) than northern ligneous peats (92 ± 14%). Bulk density and organic matter density from ligneous peats was higher in Costa Rica than in the north (BD: 0.25 ± 12 and OMD: 0.13 ± 0.05 g/cm^3 in Costa Rica vs. 0.11 ± 0.05 and 0.1 ± 0.03). %C was almost double in northern peatlands (50.9 ± 4.0%) than in Costa Rica (26.7 ± 9.8%). %N was also higher in ligneous northern peats (1.3 ± 0.5%) than in Costa Rica (1.1 ± 0.4%).

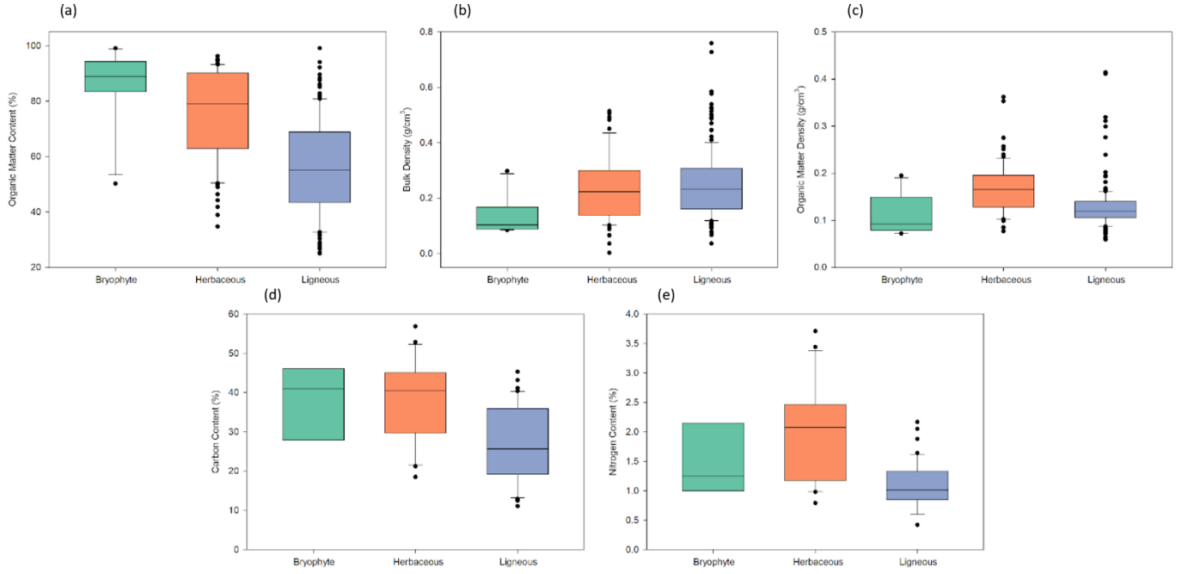

Figure 12: Box plots of geochemical data by peat type. (a) Organic matter content (%), (b) bulk density (g/cm^3), (c) organic matter density (g/cm^3), (d) carbon content (%), and (e) nitrogen content (%).

4.3 Costa Rican peatland carbon stock

To estimate peat carbon storage, previous studies have often reverted to a simple formula that combines peat extent, average peat depth, average bulk density, and an assumed carbon content of 50 or 51% (Gorham, 1991; Loisel and Yu, 2013; Yu et al., 2010). Here, we used a peatland extent of 1,456 km^2 (Rabel et al., 2025), an average peat depth of 150 cm (which is the average depth for the 9 sites presented in this study), an average bulk density of 0.24 gC/cm^3 and an average carbon content of 30%. These values yielded a peat carbon store of 1.573 x 10^{14} g C, or 0.1573 Gt C.

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Of course, every one of the values used above carries uncertainties, starting with the peatland extent, which was performed using a probabilistic approach (Rabel et al. 2025) that may under- or over-estimate the extent. Illustrating this uncertainty is the range of peat extent for Costa Rica that was provided in previous (global) peat mapping efforts, from 577 to 2670 km^2 (Melton et al., 2022; UNEP, 2022). Using these peatland extents results in carbon stores ranging between 0.0623 and 0.2883 Gt C for Costa Rican peatlands. An important aspect to highlight is that previous peat-carbon store estimates often use 0.1 g/cm^3 as their bulk density in combination with an assumed 50% carbon content, yielding an organic carbon density of 0.05 gC/cm^3. The results for Costa Rica suggest an organic carbon density of 0.072 gC/cm^3 (0.24 g/cm^3 * 30%), about 30% greater than the previous value. While this result should be further confirmed through additional measurements, it cautions against the broad application of well-established knowledge from extra-tropical peatlands to discuss peat-carbon characteristics in tropical peatlands. Estimates created out of those values would greatly underestimate the tropical carbon store.

4.4 Peatland successional pathways in Costa Rica

Ecological succession can be inferred from the plant macrofossil results. It is noteworthy that three out of our four study sites started as palm swamps (GAN, RM, MQ). While GAN has remained a palm swamp throughout its development (800 years), the mangrove site (RM) switched from palm to mangrove around 540 years ago. While ecological succession in wetlands and peatlands tends to go from wetter to drier environments, there exist many examples of opposite trends (from drier to wetter) that can be caused by changes in local hydrology that might have been induced by a lowering of the base level (land subsidence, sometimes caused by earthquakes or large erosional storms such as hurricanes) or an increase in sea level (causing flooding and saltwater intrusion) (McCloskey and Liu, 2012; Phillips and Bustin, 1996; Urquhart, 2009; Urrego et al., 2019). The coastal area where our mangrove site is located is both geomorphologically and tectonically active, with an estuary in close proximity and a history of paleo-earthquakes (Denyer, 1998). As for the riverine site (MQ), the switch from palm swamp to herbaceous peatland took place around 5500 years ago and suggests a transition into a drier (though still wet) environment. The river that transects this peatland complex (Rio Medio Queso) is a tributary of Rio San Juan, itself a major outlet of Lago Nicaragua (Fig. 3). Mid-Holocene fluctuations in the lake level would have likely caused large shifts in the hydrological regime and botanical nature of the regional wetland complexes that span hundreds of km^2 in the area. Unfortunately, there is a lack of paleodata from the lake to corroborate with the peat core.

The montane peatland presented a different story. The BOG70 core indicates a recent switch from herbaceous to bryophyte peat that might have taken place within the past few decades (Fig. 5a). It is important to note that the modern ecological assemblage of the peatland is unusual for Costa Rica; *Sphagnum* and lichen species are typically found in cool environments of the mid- and high-latitude regions of the world. Atop the Cordillera de Talamanca, where the site is found, temperatures are cool and much of the moisture is brought in the form of fog (cloud forest). The little evidence of Holocene-scale changes in paleoclimate in this area comes from a lake study (Lago Morrenas) that shows a mesic early Holocene, followed by wetter conditions during the middle Holocene, and high variability throughout the late Holocene (Kerr et al., 2024). It is possible that regional changes in hydroclimate have allowed for *Sphagnum* peat to develop here. However, the lake-based palynological record available for the region does not indicate clear shifts in vegetation in the late Holocene (Horn, 1993).

4.5 Peat type as a driver of organic matter quality in Costa Rica

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It has been shown that tropical lowland peats typically contain high levels of aromatics and low levels of carbohydrate compounds, indicative of humified, but stable, conditions that allow for long-term carbon accumulation and storage (Hodgkins et al., 2018; Verbeke et al., 2022). Our results generally corroborate the finding that tropical peat deposits contain substantial relative abundance of aromatic (and aliphatic) components.

The effect of peat type on the nature of the peat carbon compounds is worth discussing. But first, we note that peat was generally very humified throughout all the profiles (von Post and C/N), except for the near surface of BOG70 and one thin layer in RM, making correlations with our FTIR data challenging. As for peat types, our analysis indicates a much greater abundance of aliphatic compounds in the herbaceous peat when compared to the other types, suggesting a potential signature for herbaceous plants. The relationships between OM content, aliphatics, and aromatics were firmly positive, confirming that stable components are controlling organic matter accumulation (e.g., Hodgkins et al., 2018; Leri et al., 2025; Verbeke et al., 2022).

The anticipated effect of depth (and age) on the carbon compounds was not straightforward. The relative abundance of carbohydrates was expected to decrease downcore (representing a progressive decay of those labile compounds) and in turn induce relative increases in the more stable components of organic matter (aromatics and aliphatics) (Broder et al., 2012; Tfaily et al., 2014; Upton et al., 2018). We identified those trends at the montane (BOG70) and the coastal peat swamp (GAN) sites. However, in the case of the riverine (MQ) and mangrove (RM) sites, no such trends along-core were discerned. While it is possible that the nature of some tropical peats does not lend itself to those trends, as the organic matter mineralizes very quickly (i.e., near the top of the soil profile), the interference of silica in the carbohydrate signature could have also further hindered the visibility of those expected relationships with depth by creating anomalously high carbohydrate peaks where, in reality, little carbohydrates were to be found.

Lastly, it is worth mentioning the potential importance of root biomass in the process of carbon accumulation at our study sites. While we did not quantify root contribution to the peat mass (those analyses will be underway shortly), it is clear from the plant macrofossil analysis that the peat itself is often almost entirely composed of fine roots, as observed in other tropical settings (Chimner and Ewel, 2005). These fine roots have been previously shown to be the driving contributor of organic matter accumulation in some mangrove systems (Liu et al., 2017; McKee et al., 2007; Middleton and McKee, 2001; Xiong et al., 2017) and tropical riverine swamps (Sciumbata et al., 2023). A high ratio of root contribution to the peat mass would support the hypothesis that tropical peatlands primarily rely on belowground organic matter inputs, in contrast with the more classic acrotelm / surface litter contribution to a growing mound where the water table rises and engulfs the dead plant matter (Clymo, 1984).

5 Conclusions and future work

This study offers the first systematic analysis of peat sediments for Costa Rica. The lab-based work presents important new datasets on peat geochemical properties (bulk density, organic matter content, carbon and nitrogen content). A key finding is that bulk density values tend to be greater in tropical lowland peatlands than what is typically reported for extra-tropical peatlands. The infrared spectroscopy confirms the mineralized, stable nature of the tropical peats, both in the lowlands and at high-elevation sites. Importantly, this observation applies across our four (diverse) sites, regardless of peatland type, age, or geomorphological setting, suggesting that rapid and intensive mineralization within the uppermost portion of the peat profile may lead to long-term carbon

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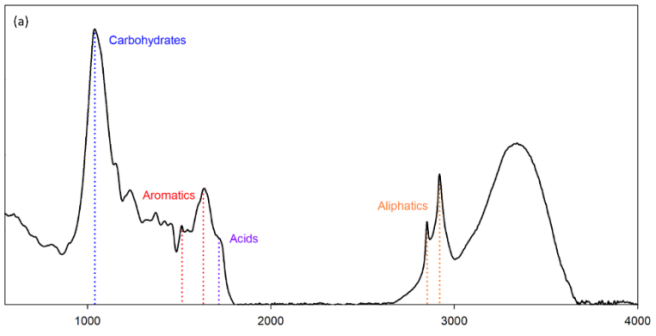
storage. Aliphatic compounds were found in much greater abundances in herbaceous-dominated peatlands than the other types, suggesting that their presence could potentially be used as a signature for this herbaceous peat. In terms of peatland dynamics, a broad range of peatland inception ages were found across Costa Rica. Spatiotemporal patterns are thus far related to peatland type, with older peats found in the montane region, intermediate and variable peat ages found along inland, river floodplains, and young peats found along the coast. Averaged apparent rates of peat accumulation in the lowlands have been rapid and steady over time, with age-depth models generally displaying linear trends, likely due to the humified nature of peat. High-elevation peat has accumulated at a markedly lower rate than the lowland sites, most likely because of the temperature limitation on plant production as well as the older age of those peat deposits. Lastly, the total peat carbon store in Costa Rica is estimated at 0.1573 gigatonnes, on the basis of our new analyses combined with a recent peatland probability map. Overall, this study provides critical, first of its kind information on the characteristics of Costa Rican peats, deepening our overall knowledge on tropical peatlands.

Appendix A



(b)

Compound	Wavelength
Carbohydrates	1030
Aromatics	1510 1630
Acids	1720
Aliphatics	2850 2920

Figure A1: Example of an FTIR spectra and peak finding for a peat sample. (a) The major functional compounds are identified after baseline correcting and normalization within the custom R script, and (b) wavenumbers that relate to carbon compounds of interest.

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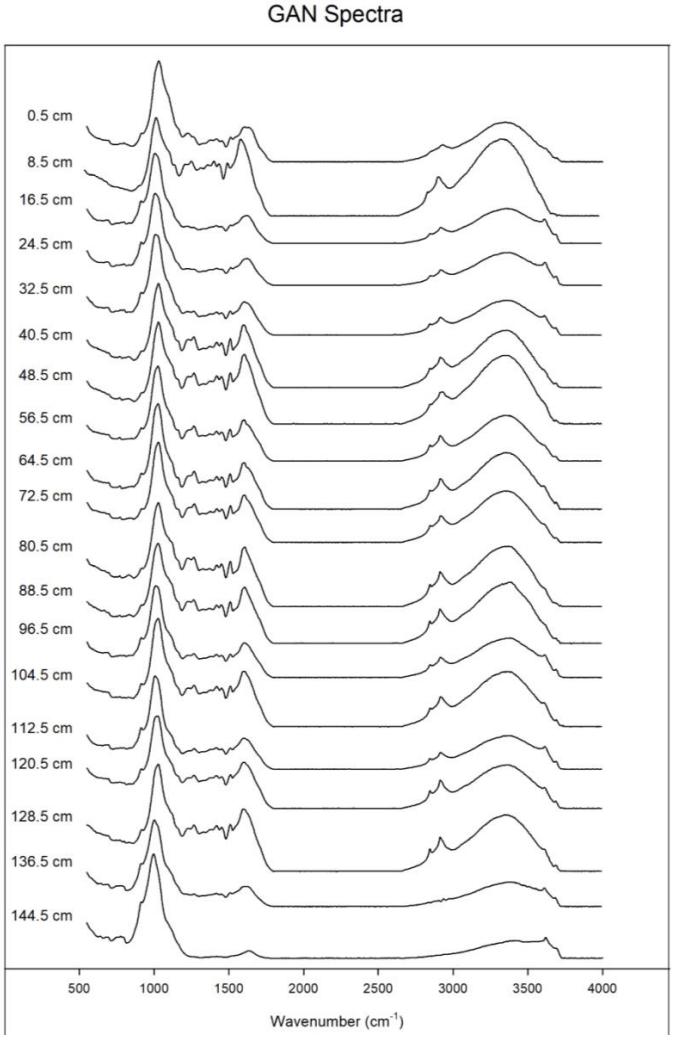

Figure A2: Stacked overlay of the FTIR spectra, GAN core.

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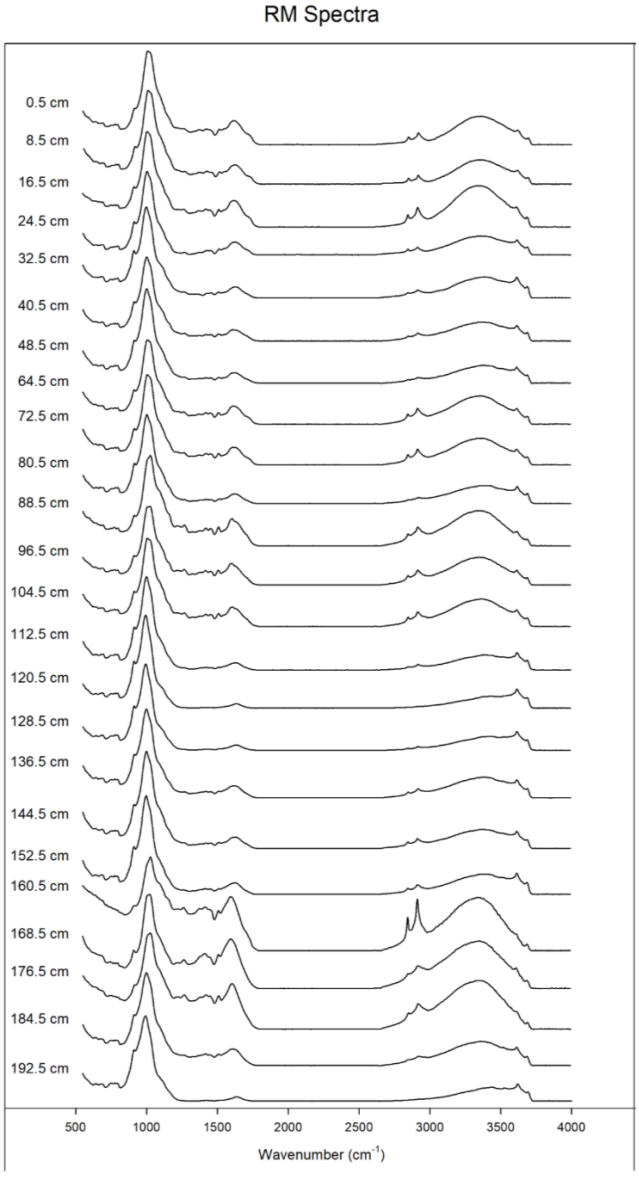

Figure A3: Stacked overlay of the FTIR spectra, RM core.

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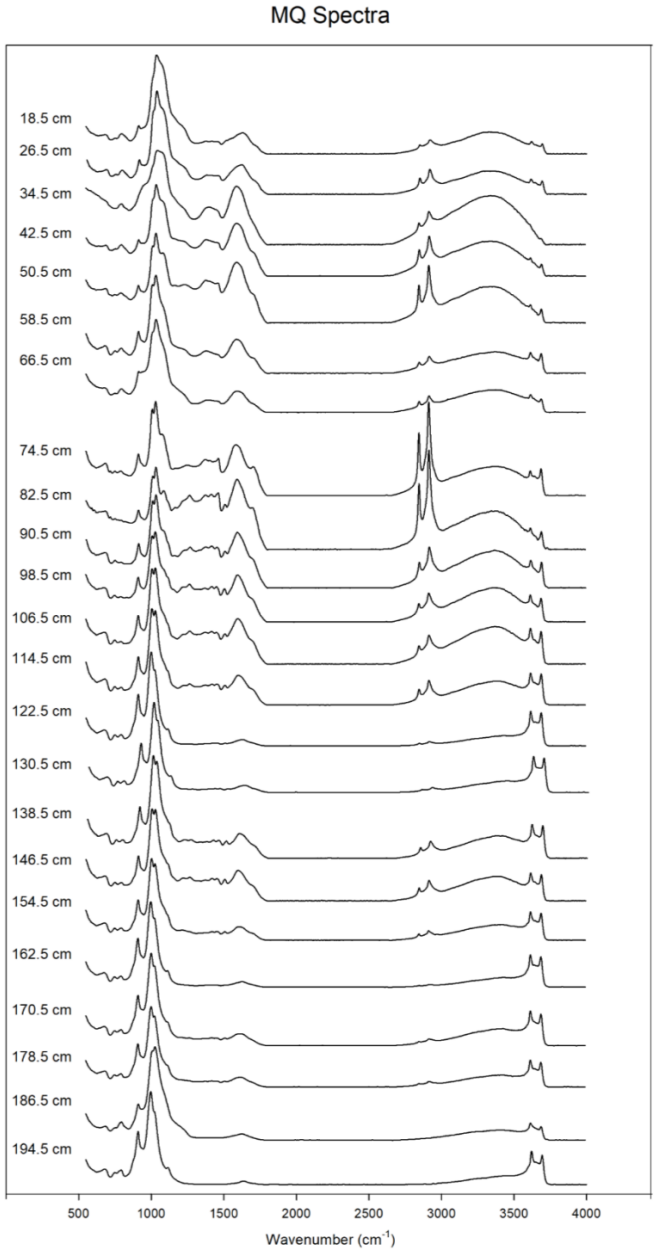

Figure A4: Stacked overlay of the FTIR spectra, MQ core.

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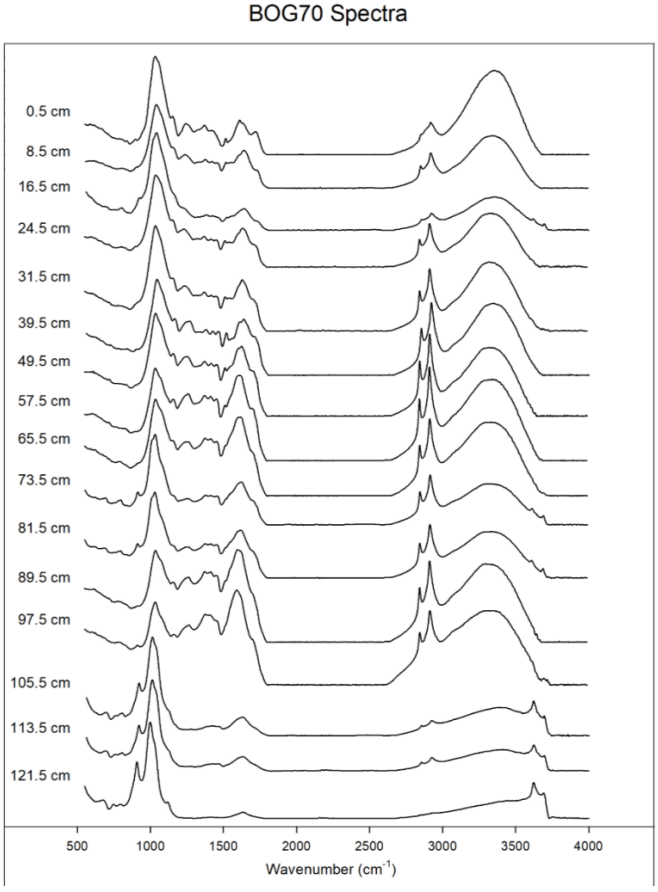

Figure A5: Stacked overlay of the FTIR spectra, BOG70 core.

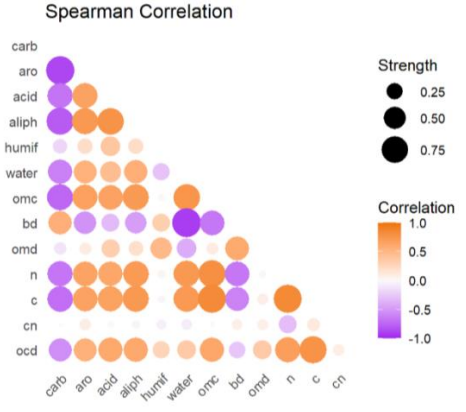

Figure A6: Spearman Correlation Matrix. Carb: carbohydrate; Aro: aromatic; Acid: acids; Aliph: aliphatics; Humif: humification; Water: water; Bd: bulk density; Omc: organic matter content; N: nitrogen; C: carbon; CN: Carbon/Nitrogen

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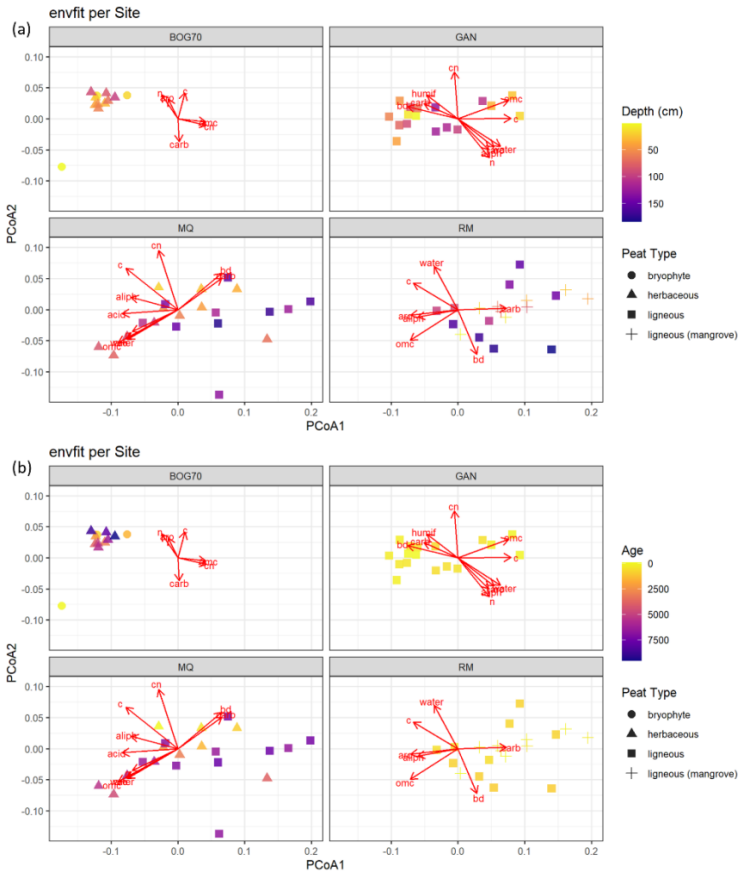

Figure A7: Principal Coordinates Analysis (PCoA) using Bray-Curtis Distances overlaid by envfit vectors separated by core. The data in the ordination is additionally visualized by (a) depth and peat type and (b) age and peat type.

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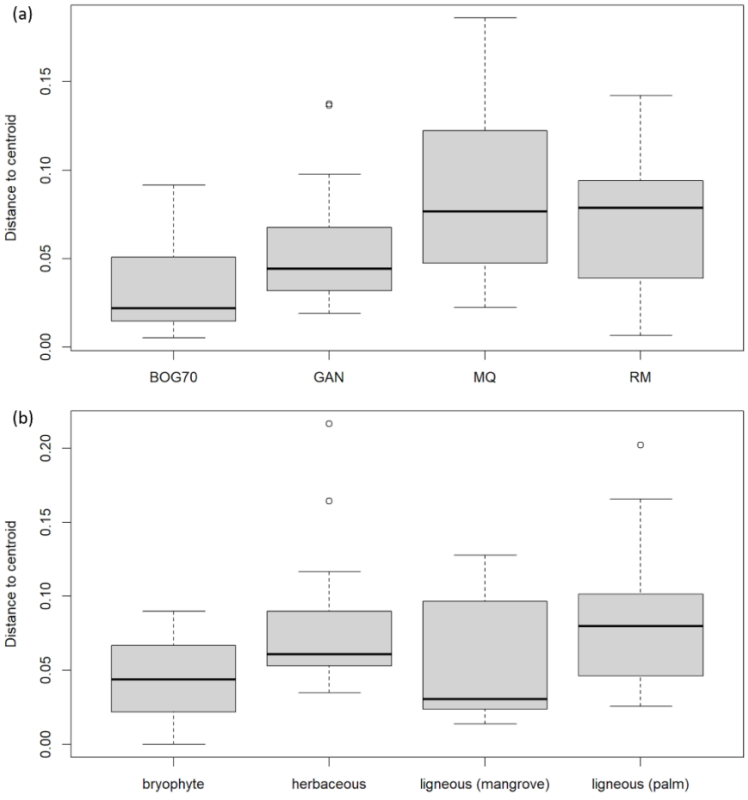

Figure A8: (a) Betadisper box plot among cores, and (b) betadisper box plot among peat types

Proxy and Resolution	Cores
Loss-on-ignition (every 2 cm)	GAN-4 GAN-SF AIR MAN RM MQ-S9 MQ-S10 LM SJ BOG70 BOG68
Carbon & Nitrogen Content (every 8 cm)	GAN-4 RM MQ-S9 BOG70
von Post Humification Index (every 8 cm)	GAN-4 RM MQ-S9 BOG70
Macrofossil Composition (every 8 cm)	GAN-4

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	RM MQ-S9 BOG70
Fourier-Transform Infrared Spectroscopy (every 8 cm)	GAN-4 RM MQ-S9 BOG70
Radiocarbon (^{14}C) Dating	GAN-4 GAN-SF AIR MAN RM MQ-S9 MQ-S10 LM SJ BOG70 BOG68

Table A1: Summary of the proxies and analytical resolution used to analyze the Costa Rican peat cores.

Variable	R^2	p value	Significance
carb	0.607	≤ 0.001	***
aro	0.549	≤ 0.001	***
acid	0.415	≤ 0.001	***
aliph	0.417	≤ 0.001	***
humif	0.034	0.334	
water	0.704	≤ 0.001	***
bd	0.533	≤ 0.001	***
omc	0.952	≤ 0.001	***
n	0.682	≤ 0.001	***
c	0.866	≤ 0.001	***
cn	0.880	≤ 0.001	***

Table A2: Statistical summary for the Environmental Fit (envfit) variables. Carb: carbohydrate; Aro: aromatic; Acid: acids; Aliph: aliphatics; Humif: humification; Water: water; Bd: bulk density; Omc: organic matter content; N: nitrogen; C: carbon; CN: Carbon/Nitrogen

Test	Factor(s)	Df	Sum of Squares	R^2	F	p-value	Significance
PERMANOVA	Core	3	0.31453	0.434	16.64	≤ 0.001	***

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	Peat type	3	0.20441	0.282	8.52	≤ 0.001	***
	Core + Peat type	6	0.36289	0.501	10.38	≤ 0.001	***
Betadisper (dispersion)	Core	3	0.02396	–	5.21	0.003	**
	Peat type	3	0.00962	–	1.63	0.192	
dbRDA	Core + Peat type	6	0.378	0.444	8.25	≤ 0.001	***
	Core	3	0.321	–	14.02	≤ 0.001	***
	Peat type	3	0.057	–	2.49	0.024	*

Table A3: Statistical summary for PERMANOVA, Betadisper, and dbRDA analyses. The significance levels are reported as follows: * = 0.05, ** = 0.01, * = 0.001**

Core 1	Core 2	R^2	F	p-value	Significance
BOG70	GAN	0.365963	15.58425	0.0015	**
BOG70	MQ	0.444373	24.79284	0.0015	**
BOG70	RM	0.669569	58.76417	0.0015	**
GAN	MQ	0.129964	5.377604	0.0168	*
GAN	RM	0.331023	16.82385	0.0015	**
MQ	RM	0.071174	2.911872	0.066	

Table A4: Statistical summary for pairwise PERMANOVA among cores.

Peat type 1	Peat type 2	R^2	F	p-value	Significance
bryophyte	herbaceous	0.102651	2.402275	0.092	
bryophyte	ligneous	0.15383	6.908256	0.004	**
bryophyte	ligneous (mangrove)	0.68727	21.97643	0.009	**

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herbaceous	ligneous	0.1443	9.274842	0.004	**
herbaceous	ligneous (mangrove)	0.412068	18.92369	0.004	**
ligneous	ligneous (mangrove)	0.089831	4.342661	0.0252	*

Table A5: Statistical summary for pairwise PERMANOVA among peat types.

Data availability

The data will be made publicly available through Zenodo upon publication of this study.

Author contributions

JL conceptualized the study; HM, JL, and ACL performed the data analysis; HM and MA performed the labwork; JL, HM, MA, and JRW completed the fieldwork and acquired the samples; JL administered the project; HM and MA completed the study visualizations (figures); HM wrote the original draft; HM, JL, JRW, ACL, and MA contributed to manuscript editing and review.

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