

Forest Diversity and Environmental Factors Shape Contrasting Soil-Litter BVOC and Methane Fluxes in Three Central Amazonian Ecosystems

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Abstract

Biogenic volatile organic compounds (BVOCs) play a crucial role in biosphere-atmosphere interactions and the global carbon cycle. While vegetation is recognized as the primary source of BVOC fluxes in forest ecosystems, recent studies suggest that the carbon-rich soil-litter compartment plays a significant role in gas fluxes. However, the drivers, variability, and magnitude of these fluxes across different forest types remain poorly understood. This is particularly notable in the Amazon rainforest, the world's largest source of BVOCs, where measurements remain scarce. In this study, we investigated the net soil-litter gas exchange of BVOCs and methane, along with their potential drivers - soil and litter nutrient content, soil and litter microbial biomass, soil temperature, and soil moisture - across three forest types in central Amazonia: White Sand Forest (WS), Upland Forest (Up), and Ancient River Terrace Forest

(AR). Our results showed distinct soil-litter gas exchange patterns across the forest types. WS exhibited both high emissions and consumption of gases, notably high acetaldehyde and methane emissions, along with a uptake of monoterpenes. Up showed lower overall fluxes, with moderate emissions and consumption of dimethyl sulfide (DMS), isoprene, and acetaldehyde. In contrast, AR presented no significant fluxes. Among the variables tested, models indicated that soil moisture and temperature were the strongest drivers of fluxes in WS, whereas microbial biomass was the main driver in Up. Our measurements suggest that, despite covering a relatively small area in the Amazon basin, WS can be a significant ecosystem for BVOC and methane fluxes, where these fluxes are influenced by soil moisture and temperature. Our findings underscore the need to account for forest-type-specific fluxes when modeling BVOC and methane emissions in the Amazon, particularly under changing climate conditions.

Key words

Amazon rainforest; Biogenic Volatile Organic Compounds (BVOC); Methane (CH₄); rain-induced emissions; Soil-litter fluxes; Forest heterogeneity; Soil-litter microorganisms

1. Introduction

Biogenic Volatile Organic Compounds (BVOCs) play critical roles across scales, from cellular processes to global climate regulation. While primarily emitted by plants, BVOCs can also be produced and consumed by soils, litter and microorganisms. Once released into the atmosphere, they actively participate in atmospheric chemistry and physics, influencing climate dynamics. BVOCs react with key atmospheric oxidants - including hydroxyl radicals (OH), ozone (O₃), and nitrate radicals (NO₃) - to form secondary organic aerosols (SOAs) (Artaxo et al., 2022; Yáñez-Serrano et al., 2020). SOAs, in turn, have a major influence on cloud properties, enhancing cloud condensation nuclei (CCN) concentrations, which impacts precipitation patterns and alters cloud lifecycles (Liu and Matsui, 2022). Depending on their chemical composition, SOAs can also influence the Earth's radiation budget by scattering incoming solar radiation (resulting in a cooling effect) or absorbing outgoing longwave radiation. Additionally, depending on the concentration of NO_x, BVOCs contribute to the formation of tropospheric ozone - an important greenhouse gas and a major air pollutant (Vella et al., 2025). Given these large-scale impacts, accurately quantifying BVOC fluxes in terrestrial ecosystems is essential for advancing

our understanding of forest–atmosphere interactions and for improving Earth system models, thereby improving climate predictions.

Global emissions of BVOCs from terrestrial vegetation are estimated at approximately 760 Tg C yr⁻¹, with isoprene (C₅H₈) and monoterpenes (C₁₀H₁₆) accounting for around 70% and 11% of these emissions, respectively (Guenther et al., 2012, Sindelarova et al., 2014). Isoprene is a simple building block compound emitted in large quantities, particularly by tropical forests. Monoterpenes (e.g., α -pinene, β -pinene, limonene) are structurally more complex (Guenther et al., 2012; Alves et al., 2016) with (-)- α -pinene being the second most emitted compound (Zannoni et al., 2020; Yanez-Serrano et al., 2018). The Amazon rainforest alone contributes about 40% of global BVOC emissions, playing a critical role in the global carbon cycle (Guenther et al., 2012; Wang et al., 2024; Tripathi et al., 2025). However, these global estimates primarily consider emissions from plants, neglecting potential contributions from soil and litter, which might also include a large variety of BVOC chemical species. This gap is particularly significant given recent evidence that the soil–litter together is a compartment that can also play a crucial role in BVOC emissions (Bourtsoukidis et al., 2018; Drewer et al., 2021; Fan et al., 2020, 2024; Peñuelas et al., 2014; Tang et al., 2019). Within this compartment, multiple biological and physical processes influence BVOC dynamics. These include plant-related processes such as intra- and inter-organism communication, herbivore defense, and symbiotic interactions (Gfeller et al., 2013; Lin et al., 2007; Rasheed et al., 2021; Steeghs et al., 2004; Tang et al., 2019; Trowbridge et al., 2020). Additionally, soil microorganisms produce and consume BVOCs for communication and ecological interactions (e.g., defense and competition), with these compounds also being released as residual metabolic products (Isidorov & Jdanova, 2002; Leff & Fierer, 2008; Liu et al., 2024; Monard et al., 2021).

Greenhouse gases (GHGs), such as methane (CH₄), carbon dioxide (CO₂) and nitrous oxide (N₂O), are also produced and consumed by soil microorganisms through key metabolic processes, including methanogenesis, methanotrophy, and respiration (Conrad, 2009,2020; Hofmann et al., 2016). While CO₂ and methane are not classified as BVOCs, they play a crucial role in overall gas exchange processes and provide a broader perspective of soil-litter gas (carbon) fluxes. Moreover, environmental factors such as soil moisture, temperature, and nutrient availability influence both BVOC and GHG fluxes, albeit through distinct-but

interconnected-biological and physical mechanisms (Greenberg et al., 2012; Tang et al., 2019; Asensio et al., 2007). These interconnected processes drive net ecosystem gas exchange between the soil-litter compartment and the atmosphere, making methane and CO₂ key components for understanding processes driving BVOC flux dynamics.

Fluxes of GHGs and BVOCs can also be linked to litter decomposition. This process is influenced by various physical factors, including soil moisture, temperature, and nutrient availability, which collectively can enhance microbial activity—one of the main drivers of these fluxes (Greenberg et al., 2012; Tang et al., 2019; Mäki et al., 2017; Asensio et al., 2007). Specifically, microbial processes such as nitrification and denitrification can result in the production and consumption of N₂O in soils (Butterbach-Bahl et al., 2013; Snyder et al., 2009). Together, these processes drive the net ecosystem exchange of BVOCs and GHGs between the soil-litter compartment and the atmosphere, and the magnitude and direction of this exchange may vary across different ecosystem types.

The Amazon Basin is a mosaic of diverse forest types (Oliveira-Filho et al., 2021), each with distinct plant species composition (Ter Steege et al., 2013), shaped by the region's highly variable soil properties (Quesada et al., 2011; Quesada et al., 2012). Although Amazonian heterogeneity is known to play a critical role in regulating biogeochemical cycles, comparative studies across forest types, especially at the soil–litter interface, are still scarce. Distinct interactions between vegetation and soil can lead to highly variable patterns of BVOC and GHG exchange. This lack of representation underscores the urgent need for studies that account for the region's ecological diversity to better capture the unique contributions of each forest type to biogeochemical processes. Quantifying this variability is key to improving both regional and global models, as gas fluxes are unlikely to be uniform within the Amazon.

To address these gaps, we investigated soil-litter fluxes of BVOCs (acetaldehyde, methanol, m/z 42, dimethyl sulfide, isoprene and monoterpenes) and GHGs (CH₄ and CO₂), soil and litter nutrient content and microbial biomass, and soil temperature and moisture from three forest types in central Amazonia: (i) Ancient River Terrace Forest - a forest that was flooded in the past and is no longer flooded due to changes in the river course (paleoigapó); (ii) White Sand Forest (locally called campinarana) - a less common forest type that occupies about 5% of the

Amazon basin (Adeney et al., 2016); and (iii) Upland Forest (locally called terra-firme) - the most common forest in Amazonia, with the highest plant species richness (Emidio et al., 2016; Luize et al., 2018). We aimed to answer the following questions: (i) what is the emission/consumption of BVOCs, CO₂, and CH₄ in magnitude and chemical diversity, and; (ii) what are the main drivers of soil-litter gas exchanges across these three forest types in central Amazonia (specifically, soil moisture, soil temperature, nutrient content and microbial biomass from soil and litter)?

2. Material and Methods

2.1 Site Description

This study was conducted in the MAUA–PELD experimental plots (PELD is the abbreviation in Portuguese for long-term ecological research) (Fig. 1) at the Amazon Tall Tower Observatory (ATTO) experimental site. This site is located 150 km northeast of Manaus and is part of the Uatumã Sustainable Reserve (USDR), which covers an area of 424,430 hectares (Andreae et al., 2015). The climate is tropical humid, with average annual rainfall of 2,376 mm and a temperature of 28°C (Botía et al., 2022). There are two distinct seasons, the wet season from December to May and the dry season from July to October, with transition seasons in between. The ATTO site contains three dominant non-flooded ecosystems: a dense Upland Forest (Up) on the plateau, with an elevation close to 100 m (*terra-firme*); a White Sand Forest (WS) (*campinarana*); and another type of *terra-firme* vegetation that developed on the lower-laying ancient river terraces (Ancient River Terrace Forests (AR)) (Fig. 1) (Andreae et al., 2015).

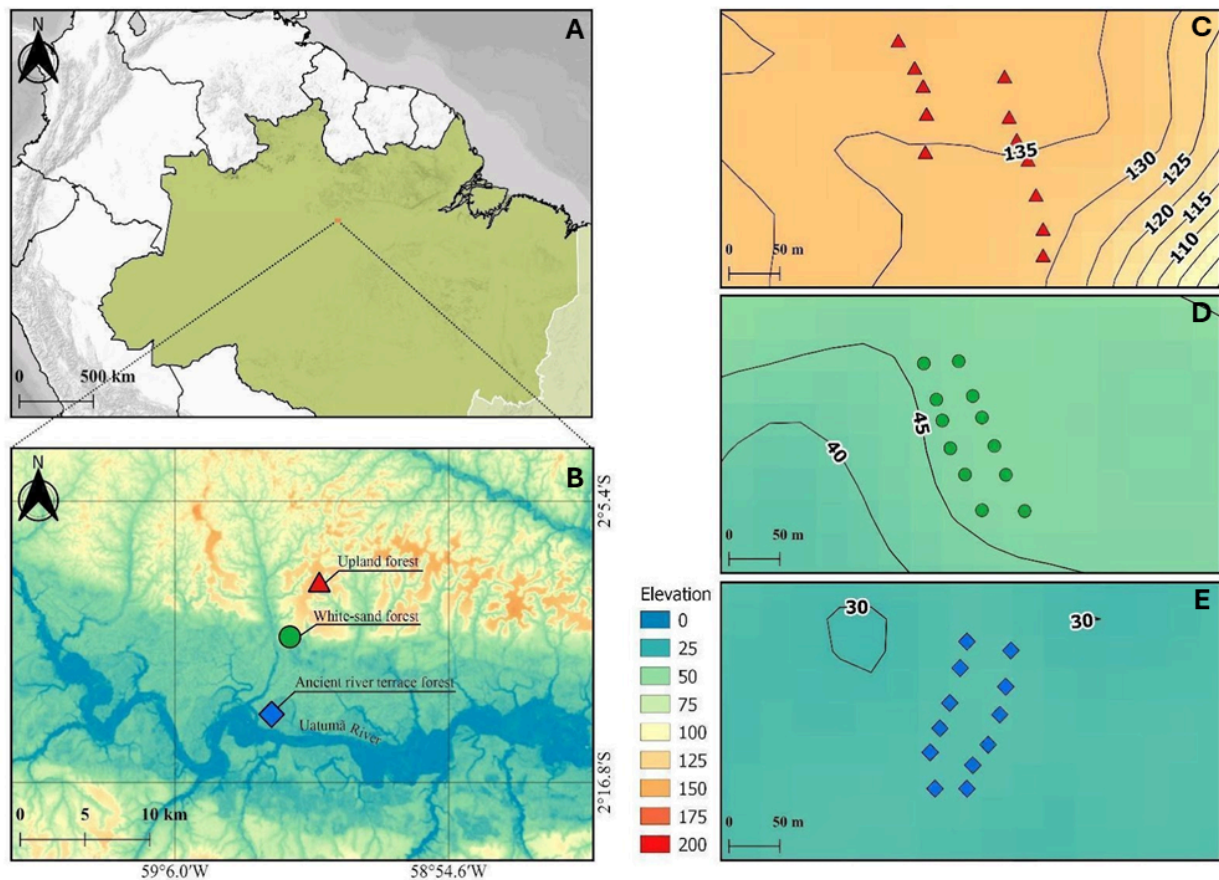


Figure 1. (A) Location of the ATTO site. (B) A map illustrating the locations of the different forest types evaluated in this study: Upland Forest, White Sand Forest, and Ancient River Terrace Forest; and showing the Uatumã River, a tributary of the Amazon River. (C), (D) and (E): The distribution of sampling points along the Transects in each forest type (Upland Forest - top, White Sand Forest - middle, and Ancient River Terrace Forest - bottom); black lines and numbers indicate the elevation (above sea level – a.s.l.).

Topography is critical to soil formation in the central Amazon region (Quesada et al., 2009). At the ATTO site, a clear topographic gradient is associated with different soil characteristics (Fig. 1). In the Ancient River Terrace Forest, soil contains more silt and clay (39% sand, 37% silt, 23% clay) in comparison to the adjacent sandy White Sand Forest soils (57% sand, 40% silt, 1.50% clay). Upland Forest soils are more clayey and contain very little sand (13% total sand, 34% silt, 52% clay) (data from this study; supplementary material, Section 1; Table S1). Upland Forest soils, which are predominantly ferrasols, are known to hold more water

than other tropical soils, benefiting forest activity during the dry season (Quesada et al., 2009). Ancient River Terrace Forest soils are typically allisols, younger and richer in nutrients compared to upland ferralsols (Andreae et al., 2015). White Sand Forest soils are arenosols, characterized by high permeability and low water retention, with low specific heat capacity and often nutrient-poor organic layers (Quesada et al., 2011). The study area in the White Sand Forest has high water table variability, with a hard subsoil layer that restricts drainage and can flood the root system during the wet season (Demarchi et al., 2022).

2.2 Sampling Design

For each forest type, a PELD-MAUA plot (~1 hectare) (<https://peld-maui.inpa.gov.br>) was selected. Within each plot, two 150 m Transects were established (Fig. 1) in homogeneous areas characterized by consistent vegetation structure, soil characteristics, and topography, to minimize spatial variability and avoid pseudo replication. Along each Transect, six sampling points were marked at ~30 m intervals, resulting in a total of 36 soil chamber measurements conducted on consecutive days; although this design was necessary for logistical reasons, it also allowed us to examine the influence of external factors beyond forest-type differences. Chamber-based methods (Section 2.3; Fig. 1) were employed for the in-situ quantification of CO₂, CH₄, and BVOC fluxes from the soil–litter compartment. These chambers were installed directly in the field with minimal disturbance to the surrounding environment. To account for potential background signals and chamber interferences, three blank chambers, each featuring completely bottom-sealed collars, were deployed per Transect and measured simultaneously alongside the sample chambers (Fig. 2b).

Tedlar bags (CEL Scientific, Cerritos, CA, USA) were used to collect gas samples directly from the outlet of the pump connected to the chambers, capturing the air for subsequent analysis of BVOCs, CO₂ and CH₄. After each gas flux measurement, soil temperature (T, °C) (TP-101, Delhi, India) and soil volumetric water content (VWC, %) were measured around the collar five times using a probe (AT SMT150, Cambridge, UK), and the averages were calculated. Surface soil samples were collected from the organic layer, approximately within the upper 5 cm of the soil. Samples from the litter and surface soil layers were collected inside the chamber and stored for analysis of chemical and physical characteristics and microbial biomass. Due to

expected low variation and limited possibility for laboratory analyses, nutrient samples from soil and litter (excluding carbon and nitrogen) and granulometry were collected as mixed samples pooled and homogenized from two collars. To minimize diurnal variation, each Transect was measured between 8:00 and 10:00 (local time), after which collected samples were processed and analyzed for BVOC and GHG concentrations. During the measurements, no precipitation was observed, but one large rain event occurred just before the measurement of Transect 2 of the White Sand Forest.

2.3 Flux Chamber Measurements

The flux chambers used in this study were produced by the Max Planck Institute for Biogeochemistry. The soil chamber, consisting of a lid and a soil collar (Fig. 2a), was made entirely of 100% stainless steel, with a total volume of 21 L and a surface area of 855 cm² (0.0855 m²). Two Teflon inlets were connected to the top of the chamber, and inside the chamber was a fan that mixed the gases in the chamber headspace. A PTFE-coated Viton O-ring was positioned at the edge of the collar over which the chamber was placed, ensuring a tight seal between the chamber and the collar.

Before gas sampling, each collar was carefully installed in a non-invasive manner by gently pressing its edge into the litter and surface soil to minimize disturbance to plant shoots and roots. To further ensure a tight seal preventing any potential leakage, the surrounding soil was carefully pressed against the outer edges of the collar (Aaltonen et al., 2011). This method ensured that the chamber system was effectively isolated from external gas exchange. The chamber and collar were sealed together with multiple clamps to prevent outside air from entering the chamber during measurements. The collars were installed approximately 24 hours prior to sampling to allow the surrounding environment to stabilize.

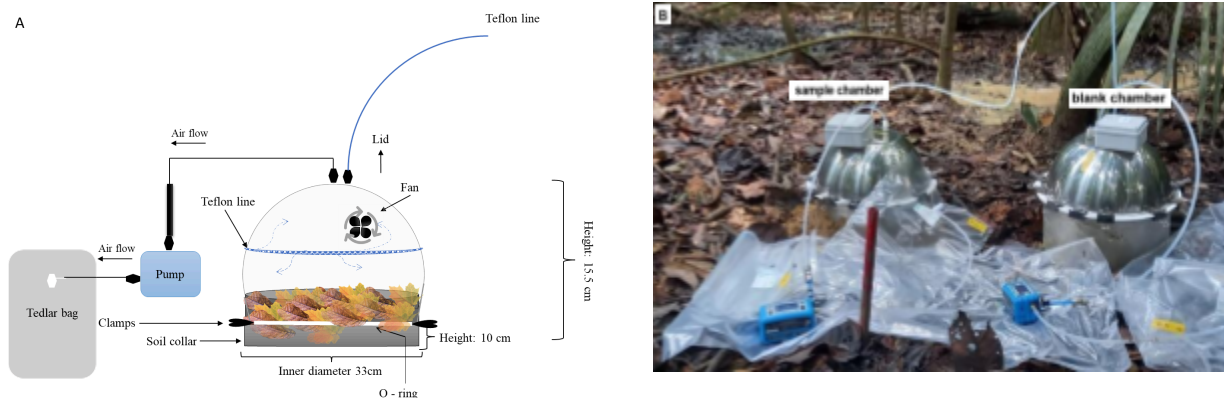


Figure 2. (A) Schematic of the flux chamber. (B) Photo of the measurement setup of the sample and the blank chamber.

Gas collection took place in December 2021, during the dry-to-wet season transition. Before placing the lid on the collar, the chamber was manually ventilated to minimize gas accumulation caused by the collar. Once the chamber was closed and sealed with clamps, the internal fan was turned on to ensure mixing. An air sampling pump (GilAir® Plus, Levitt Safety, Ottawa, ON), operating at a flow rate of 500 sccm, initiated continuous flow from the chamber outlet immediately after closure to maintain constant conditions throughout the measurement.

After 20 minutes of chamber closure with continuous air circulation, a Tedlar bag was connected to the outlet of the Teflon pump, and a 5 L air sample was collected over the next 10 minutes. By the end of the 30-minute measurement process, a total of 15 L of air had flowed through the chamber, and the last 5 L were used for subsequent analyses. The sample volumes collected were determined by the specific requirements of each analytical instrument. Measurements of BVOCs were performed using proton-transfer-reaction quadrupole mass spectrometry (PTR-QMS; IONICON Analytik, Innsbruck, Austria), prioritizing sufficient ion counts per second (CPS) and integration cycles for reliable detection of BVOC concentrations with the required precision, especially considering their typically trace levels. Greenhouse gases (GHGs) were measured using a Los Gatos Ultra-Portable Gas Analyzer (California, USA), hereafter referred to as the Los Gatos analyzer. Finally, for the offline measurements with cartridges and for specific compound identification and qualitative analysis with thermal-desorption gas chromatography time-of-flight mass spectrometry (TD-GC-TOF-MS;

Bench ToF Tandem Ionisation, Markes International, Bridgend, UK), at least 2 L of sample air was required to effectively load the adsorbent cartridges. These analyses are further detailed in the Supplementary Materials, Section 3.2.

The same procedure was followed for a blank chamber, which was measured under identical conditions to account for gas contributions unrelated to soil and litter processes. For logistical reasons, measurements were conducted with three chambers simultaneously, pairing two sample chambers with one blank chamber, followed by two additional sets, resulting in the measurement of six samples and three blank chambers per day. Because air was continuously extracted from the chamber headspace by the pump, ambient air entered the chamber through one additional inlet, which consisted of a 2 m long open Teflon tube, fixed approximately 2 m above the ground and positioned at the same location for both sample and blank chambers. The setup ensured that both chambers (sample and blank) were diluted or affected by ambient air to the same degree, minimizing potential biases.

After sampling, bags were handled carefully to prevent leakage. Potential compound losses due to adsorption onto the inner walls or diffusion through the bag material were minimized by storing all samples in a dark, stainless-steel box to avoid light exposure, and keeping them in air-conditioned lab containers at low temperatures until analysis. All samples were analyzed on the same day, within a maximum of 8 hours post-collection, following the recommendations of Beauchamp et al. (2008). Gas analysis began with the quantification of BVOCs using a PTR-QMS. Subsequent analyses of CO₂ and CH₄ concentrations were conducted using a Los Gatos analyzer. Each sample bag was then used to fill stainless steel cartridges (containing Tenax TA and Carbograph 5TD adsorbents), which were later analyzed via TD-GC-TOF-MS. Detailed descriptions of the analytical procedures and results are in Sections 3 and 3.1. of the Supplementary Material.

2.4 PTR-QMS measurements and Los Gatos analyzer measurements

Tedlar bags were connected to the PTR-QMS for analysis of BVOCs (Lindinger et al., 1998). The PTR-QMS H₃O⁺ mode was used for chemical ionization, which is extremely sensitive to all BVOCs that have a higher proton affinity than water, covering most volatile organic compounds (Edtbauer et al., 2021). Six compounds were analyzed (Table 1). The

PTR-QMS was operated under standard conditions at 2.3 mbar, and E/N 120, with 60°C, with a drift tube voltage of 600 V. During each PTR-QMS measurement cycle, the following specific protonated mass-to-charge ratios (m/z) were measured, 21 (H_3O^{18+}), 32 (O_2^+), and 37 ($\text{H}_2\text{O}-\text{H}_3\text{O}^+$), with a dwell time of 500 ms each; and Methanol (33), compound not identified (m/z 42), Acetaldehyde (45), Dimethyl sulfide - DMS (63), Isoprene (69) and Monoterpenes (137), with a dwell time of 1 second. We measured approximately 17 cycles for each bag. Mass identifications were based on the available literature (Warneke et al., 2015), and were consistent with a PTR-MS mass library database - GLOVOC (Yañez-Serrano et al., 2021) and gas calibration with certified standards.

Table 1. Compounds analyzed by the PTR-QMS

BVOC	Chemical formula (H ⁺)	m/z	Group
Methanol	CH_4O^+	33	Alcohol
not identified		42	N-compound
Acetaldehyde	$\text{C}_2\text{H}_4\text{O}^+$	45	Aldehyde
Isoprene	C_5H_8^+	69	Alkenes
Dimethyl sulfide (DMS)	$\text{C}_2\text{H}_6\text{S}^+$	63	Organosulfides
Monoterpenes	$\text{C}_{15}\text{H}_{16}^+$	137	Alkenes

Calibrations were performed before the experiment using a multi-component calibration mix containing various gases of known concentrations (Apel-Riemer Environmental, Inc.) (supplementary material; Section 2; Table S2). Four-point calibration curves were generated by diluting the multicomponent with synthetic air, humidifying the air stream with a water bubbler filled with distilled water, and controlling the flow with mass flow controllers (supplementary material; Section 2; Fig. S1). Curves were calculated considering the normalized counts per second as a function of the mixing ratio. Previously, some compounds important for soil-litter processes (Peñuelas et al., 2014) - such as acetone, ethanol, and formaldehyde - were considered for this study, but as they did not show a good fit, they were excluded from this work. The error of PTR-QMS concentration measurements of the six presented compounds is expected to consist of a systematic part and a statistical part. The systematic error consists of the uncertainty of the calibration gas standard (+5%), the error of the flow measurements (+5%), and the error of the calibration slope (14.8%, 52.4%, 12.5%, 18.4%, 18.2%, 20.4% for methanol, m/z 42,

acetaldehyde, DMS, monoterpenes and sesquiterpenes, respectively). The statistical error is based on the repeatability of the concentration measurement during the calibration routine, and was found to be 14.4%, 26.2%, 14.8%, 13.7%, 6.0%, 8.5% for methanol, m/z42, acetaldehyde, DMS, monoterpenes and sesquiterpenes, respectively. The systematic error affects both bags in the same direction, whereas the statistical error can differ between the two bags in a pair. Therefore, to evaluate the uncertainty of the fluxes, we focused on the propagated statistical uncertainty, as described in Section 2.5.

After PTR-QMS analysis, the bags were connected to a Los Gatos analyzer to measure the mixing ratios of methane and CO₂. The Los Gatos analyzer is an instrument based on laser absorption spectroscopy specifically Off-axis Integrated Cavity Output Spectroscopy (OA-ICOS), enabling ultra-sensitive, precise, and real-time measurements of trace gases in gas samples (Pohlman et al., 2021; van Asperen et al., 2024). The instrument operates at a relatively low sample flow rate (~0.1 L min⁻¹), and a minimum gas volume of 0.3 L was used as a precaution to ensure measurement stability and accurate determination of CH₄ and CO₂ concentrations; air from the sample bags was analyzed for 3 minutes, and mean concentrations were calculated from the final 2 minutes of measurement.

2.5 BVOC & GHG flux calculation

To calculate BVOC and GHG fluxes, the Volumetric Mixing Ratios of the blank chamber bags (VMR_b) were subtracted from the sample chamber bags (VMR):

$$dVMR = VMR - VMR_b \quad (1)$$

in which VMR is expressed in pptv or ppbv. By subtracting the mixing ratios of a blank chamber, dVMR represents the concentration difference attributable solely to soil and litter fluxes, corrected for potential chamber effects or the influence of ambient air entering the system. To ensure data reliability, bag pairs for which the concentration difference (dVMR) was less than or equal to the combined statistical uncertainty (calculated using the Root-Sum-Square method from the individual bag uncertainties) were assigned a value of zero. This approach ensures that only reliably detected fluxes are considered, while retaining the full sample size for modeling purposes. To convert dVMR to fluxes, we used:

$$F = dVMR * N * (V / A) * (1/T) \quad (2)$$

where N is the value of fixed molar volume at 25 °C (24.8 L mol^{-1} ; 40.3 mol m^{-3}), V is the chamber volume (0.021 m^3), A is the chamber area (0.0855 m^2), and T is the average sampling time (25 min). Flux values in $\text{nmol m}^{-2} \text{ min}^{-1}$ were converted to $\text{ng m}^{-2} \text{ h}^{-1}$.

2.6 Soil and Litter Analyses

The Thematic Laboratory of Soils and Plants (LTSP, at the National Institute for Amazonian Research - INPA) analyzed soil and litter nutrient content according to adapted protocols (EMBRAPA, 1999). The nutrients - iron (Fe^{+2}), calcium (Ca^{+2}), magnesium (Mg^{+2}), zinc (Zn^{+2}), potassium (K^+), manganese (Mn^+), phosphorus (P), and aluminum (Al) - were determined by digestion with a nitro-perchloric acid solution (Malavolta et al., 1989). Total phosphorus (P) was quantified using colorimetry (Murphy & Riley, 1962; Olsen & Sommers, 1982) and measured using a UV spectrophotometer (Model 1240, Shimadzu, Kyoto, Japan). Potassium (K), calcium (Ca), and magnesium (Mg) concentrations were determined by atomic absorption spectrophotometry (AAS, 1100 B, 250 Perkin Elmer, Ueberlingen, Germany), as described by Anderson and Ingram (1993). Soil carbon and nitrogen content was determined by the Routine Measurements & Analyses Lab (RoMA, MPI-BGC) with the elemental analyzer "varioEL" (Elementar Analysensysteme GmbH, Elementar-Straße 1, D-63505 Langenselbold, Germany). Soil porosity was analyzed using the pycnometer method described in Flint & Flint (2002). The amount of water was corrected for soil density.

For analysis of soil and litter microbial Carbon, Nitrogen, and Phosphorus (C, N, and P) contents, 2g of fresh litter and 5g of fresh soil were used from each sample chamber. These were separated into fumigated and non-fumigated samples. The fumigated samples were left with chloroform for 24 hours and then divided into two sub-samples. For first, 50 mL of KCl (Potassium Chloride) was added, and total C and N were extracted, and for the second, 50 mL of NaHCO_3 (Sodium Bicarbonate) was added for total P extraction. Following the same extraction protocol, the non-fumigated samples were prepared for direct extraction without going through the 24-hour fumigation period. Microbial C, N, and P content was estimated in fumigated and non-fumigated extracts from the difference in organic C, N, and total P measured by a TOC/TN analyzer (Jenkinson et al., 2004). The extraction of the microbial biomass was performed at INPA, and the analyses were done by the Routine Measurements & Analyses Lab (RoMA, MPI-BGC).

2.7 Statistical analyses

A total of 36 samples were evaluated ($n = 12$ per forest type). Gas fluxes were first correlated with potential predictors (soil and litter characteristics, Table 2), revealing variations between forest types. Separate regression models were built for each forest type to maximize predictive ability, with variable selection based on the following criteria: 1) given the statistical power limitation of models ($n = 12$), the maximum number of independent variables possible to include was two; thus, 2) we tested all models with one or two independent variable combinations; finally, 3) we chose the models which showed no multicollinearity and had the highest adjusted R-squared and lowest Akaike's information criterion (AIC). The "ols_step_all_possible" function from the "olsrr" package (Hebbali, 2024) was used, and multicollinearity was assessed via VIF (<2.5 ; Hair et al., 2009). Principal Component Analysis (PCA) and Pearson's correlation (Hmisc package; Harrell, 2018) were performed to explore variable interactions. Variations within forest types (e.g., between Transects) were analyzed using t-tests for normal data and Kruskal-Wallis tests for non-normal data, with a significance level of 0.05. All analyses were conducted in R (v4.3.0; R Core Team, 2023).

Table 2. Variables, their respective codes, and units.

Variable	Code	Unit
Soil carbon	c_soil	%
Soil nitrogen	n_soil	%
Soil phosphorus	p_soil	P mg/kg
Soil potassium	k_soil	K ⁺ mg/kg
Soil calcium	ca_soil	Ca ⁺² mg/kg
Soil magnesium	mg_soil	Mg ⁺² mg/kg
Soil aluminum	al_soil	Al ⁺³ mg/kg
Soil iron	fe_soil	Fe ⁺² mg/kg
Soil zinc	zn_soil	Zn ⁺² mg/kg
Soil manganese	mn_soil	Mn ⁺² mg/kg
Soil ph	ph_soil	pH
Soil temperature	soil_temp	Celsius
Soil moisture	soil_moisture	%
Litter carbon	c_litter	%
Litter nitrogen	n_litter	%

Litter calcium	ca_litter	Ca ⁺² mg/kg
Litter magnesium	mg_litter	Mg ⁺² mg/kg
Litter potassium	k_litter	K ⁺ mg/kg
Litter iron	fe_litter	Fe ⁺² mg/kg
Litter zinc	zn_litter	Zn ⁺² mg/kg
Litter manganese	mn_litter	Mn ⁺² mg/kg
Microbial biomass soil carbon	c_mic_soil	g/kg
Microbial biomass soil nitrogen	n_mic_soil	g/kg
Microbial biomass soil phosphorus	p_mic_soil	g/kg
Microbial biomass litter carbon	c_mic_litter	g/kg
Microbial biomass litter nitrogen	n_mic_litter	g/kg
Microbial biomass soil phosphorus	p_mic_litter	g/kg

3. Results

3.1 Comparison between forest types

The three forest types showed very different gas fluxes for BVOCs and GHGs (Fig. 3), with the highest fluxes observed in the White Sand Forest. Fluxes were very low in the Upland Forest, and almost no gas fluxes were observed in the Ancient River Terrace Forest. Acetaldehyde emissions showed the most significant differences between forest types, with flux averages of $35.87 \pm 46.86 \text{ mg m}^{-2} \text{ h}^{-1}$ (mean \pm SD) for the White Sand Forest, -0.09 ± 0.02 (mean \pm SD) $\text{mg m}^{-2} \text{ h}^{-1}$ for the Upland Forest, and -0.02 ± 0.008 (mean \pm SD) $\text{mg m}^{-2} \text{ h}^{-1}$ for the Ancient River Terrace Forest. Isoprenoid (isoprene and monoterpenes) fluxes were also high in the White Sand Forest, and clear differences were found between forest types concerning the speciation of monoterpenes (supplementary material; Fig. S2).

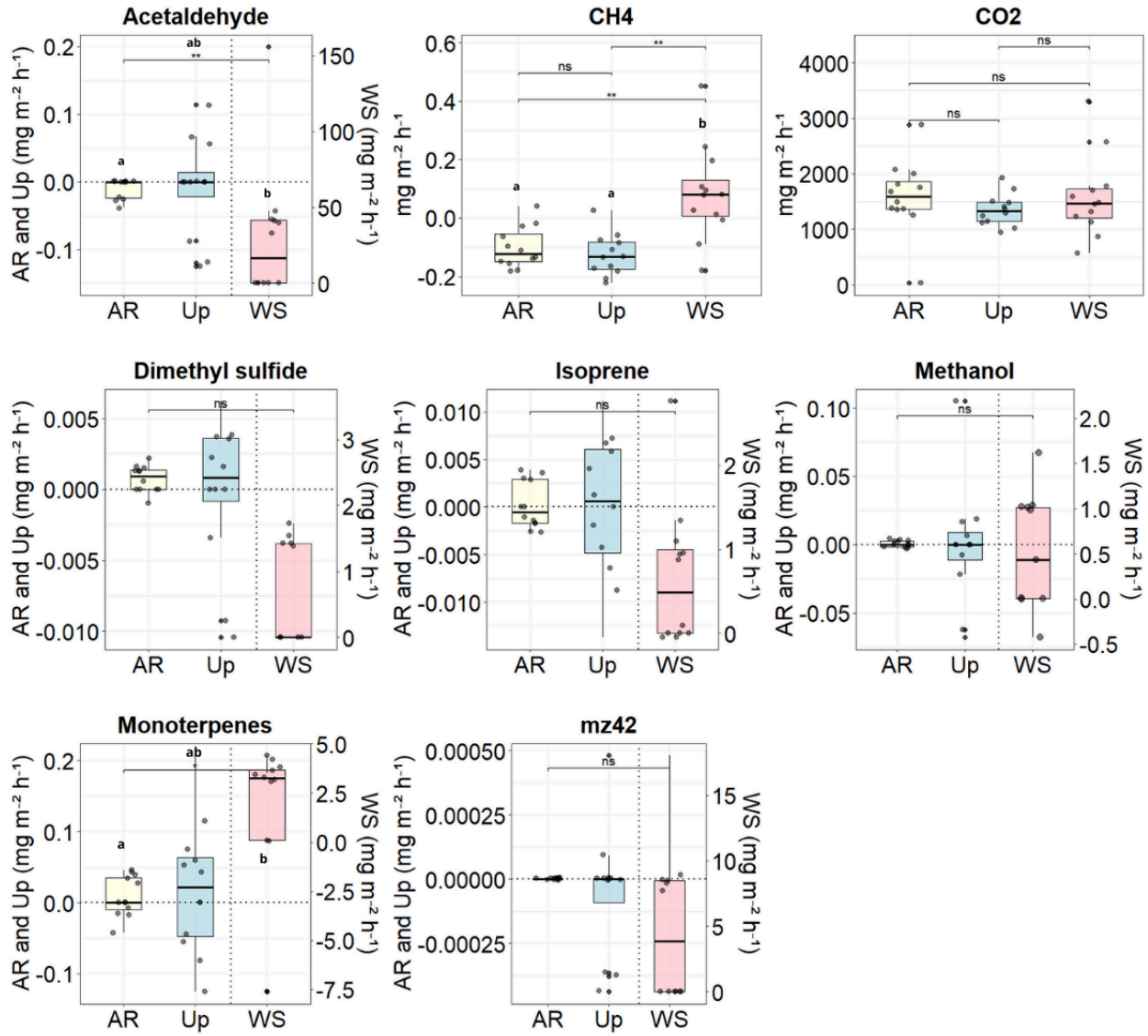


Figure 3. Biogenic Volatile Organic Compounds (BVOC) and Greenhouse Gas (GHG) fluxes from soil and litter across three forest types: Ancient River Terrace Forest (AR), White Sand Forest (WS), and Upland Forest (Up). Statistically significant differences between forest types are indicated by lowercase letters (* $p < 0.05$ *, $N=36$), “ns” is not significant. Differences were assessed using the Kruskal-Wallis test for non-normal data and ANOVA for normally distributed data, followed by Dunn’s post hoc test with Bonferroni. The left axis represents BVOC and GHG fluxes ($\text{mg m}^{-2} \text{h}^{-1}$) for AR and Up, while the right axis represents flux values for WS ($\text{mg m}^{-2} \text{h}^{-1}$). Boxes represent the median (horizontal line) and the interquartile range (IQR, first and third

quartiles). Whiskers extend up to 1.5 times the IQR, and points beyond whiskers represent statistical outliers.

In the White Sand Forest, high isoprenoid emissions were observed, with isoprene fluxes averaging $0.66 \pm 0.84 \text{ mg m}^{-2} \text{ h}^{-1}$, and monoterpene fluxes averaging $1.12 \pm 4.31 \text{ mg m}^{-2} \text{ h}^{-1}$, details on the monoterpene and sesquiterpene speciation can be found in Supplementary Material, Section 3.2. Conversely, very low (negative) values were recorded in the Upland Forest (Isoprene, $-0.005 \pm 0.003 \text{ mg m}^{-2} \text{ h}^{-1}$ and monoterpenes $-0.02 \pm 0.08 \text{ mg m}^{-2} \text{ h}^{-1}$) and Ancient River Terrace Forest (Isoprene, $-0.002 \pm -0.001 \text{ mg m}^{-2} \text{ h}^{-1}$ and monoterpenes $-0.011 \pm 0.015 \text{ mg m}^{-2} \text{ h}^{-1}$). In the WS we also observed the consumption of monoterpenes (with $-7.63 \text{ mg m}^{-2} \text{ h}^{-1}$, highlighted as an outlier in Fig. 3) and high emissions of DMS ($1.10 \pm 1.14 \text{ mg m}^{-2} \text{ h}^{-1}$), whereas a consumption of the latter was observed in the Upland Forest ($-0.001 \pm 0.005 \text{ mg m}^{-2} \text{ h}^{-1}$) and Ancient River Terrace Forest ($-0.0003 \pm 0.001 \text{ mg m}^{-2} \text{ h}^{-1}$). CH_4 fluxes showed substantial variation in the White Sand Forest, with both uptake ($-0.09 \pm 0.16 \text{ mg m}^{-2} \text{ h}^{-1}$) and emission ($0.08 \pm 0.07 \text{ mg m}^{-2} \text{ h}^{-1}$). In contrast, the Upland Forest and Ancient River Terrace Forest showed primarily methane uptake, with average fluxes of $-0.14 \pm 0.05 \text{ mg m}^{-2} \text{ h}^{-1}$ and $-0.13 \pm 0.03 \text{ mg m}^{-2} \text{ h}^{-1}$, respectively.

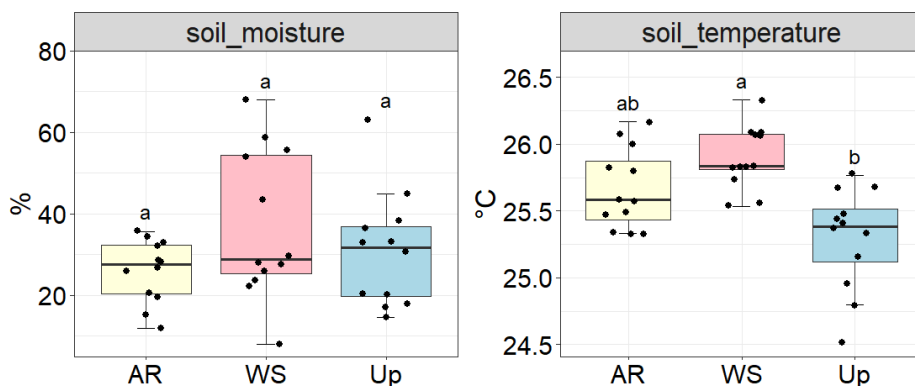


Figure 4. Soil moisture and soil temperature (expressed % for moisture °C for temperature) across the three forest types: Ancient River Terrace Forest (AR), White Sand Forest (WS), and Upland Forest (Up). Letters indicate statistically significant differences in fluxes between forest types at $p < 0.05$, $N=36$ (ANOVA test for normal distribution). Boxes show median and first and third quartiles, with whiskers and points distinguished at 1.5 times the interquartile range.

There were no statistically significant differences in soil moisture across forest types (Fig. 4); however, the White Sand Forest showed the highest and the lowest soil moisture values.

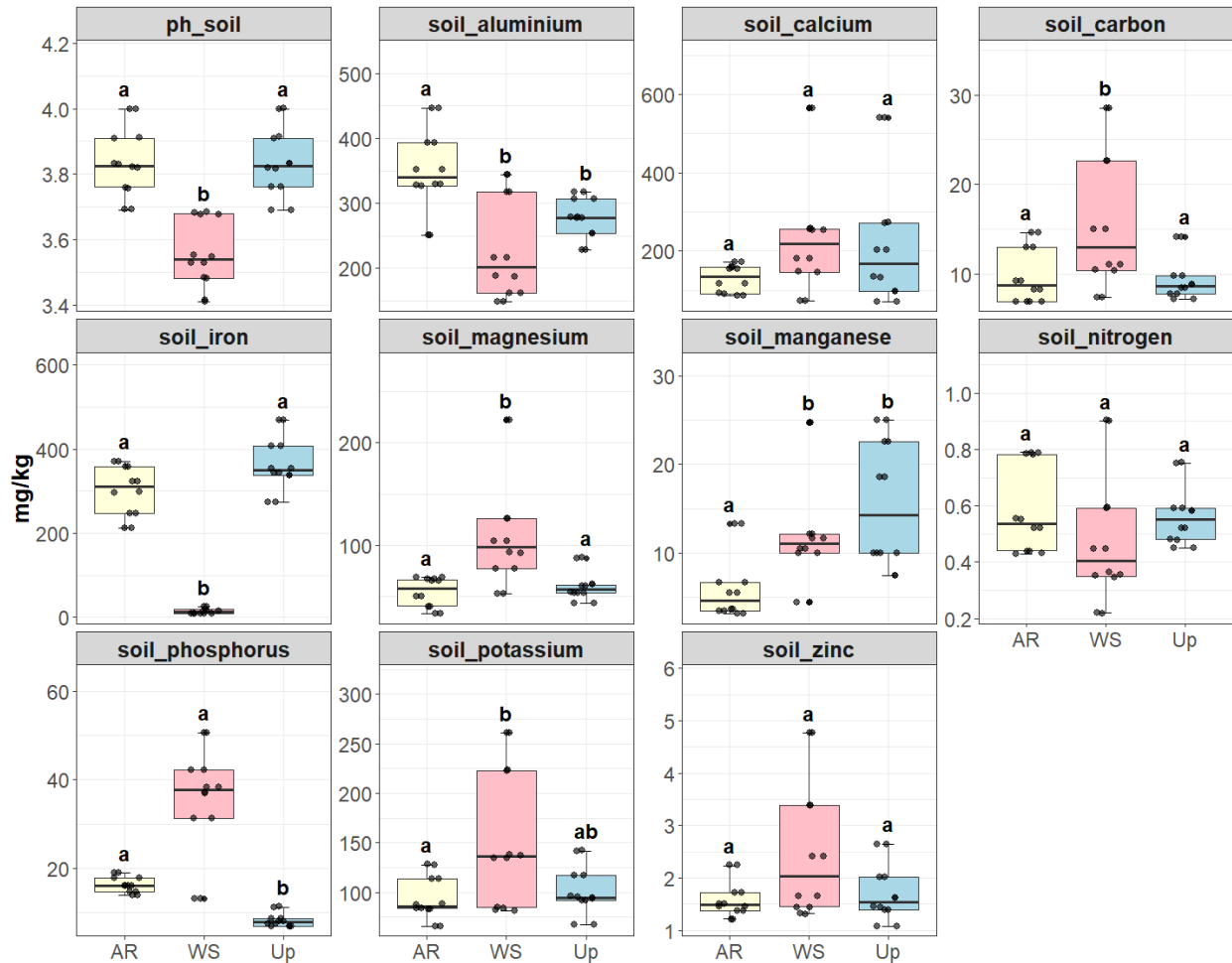


Figure 5. Concentrations of soil micro- and macronutrients in the three forest types: Ancient River Terrace Forest (AR), White Sand Forest (WS), and Upland Forest (Up). Letters indicate statistically significant differences in nutrients between forest types at $p < 0.05$, $N=36$. (ANOVA test for normal data (aluminum) and Kruskal-Wallis test for non-normal data (carbon, calcium, iron, potassium, magnesium, manganese, nitrogen, phosphorus, pH, and zinc)). Boxes show median and first and third quartiles, with whiskers and points distinguished at 1.5 times the interquartile range.

Soil macro- and micronutrients varied considerably between the forest types, with statistically significant differences in carbon, magnesium, phosphorus, and iron for the White Sand Forest. Phosphorus content was the highest in the White Sand Forest compared to other forest types (Fig. 5). All litter nutrients exhibited significant differences between forest types: Upland Forest showed the highest average concentrations of calcium, iron, manganese, and zinc, while the Ancient River Terrace Forest had the highest nitrogen, potassium, and phosphorus concentrations, and the White Sand Forest had slightly higher carbon concentrations (Fig. 6).

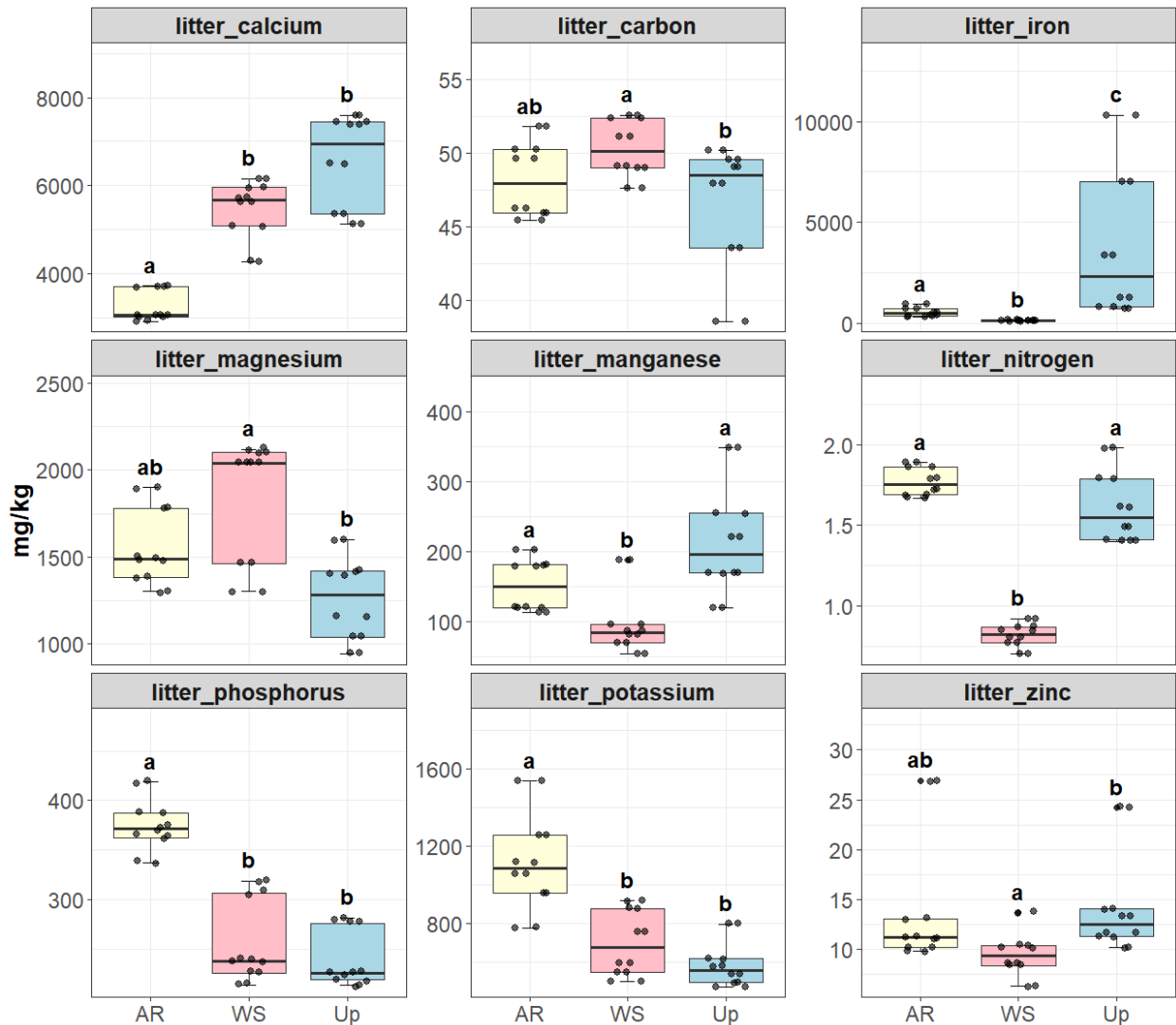


Figure 6. Concentrations of litter micro- and macronutrients in the three forest types: Ancient River Terrace Forest (AR), White Sand Forest (WS), and Upland Forest (Up). Letters indicate statistically significant differences in nutrients between forest types at $p < 0.05$, $N=36$. (ANOVA

test for normal data - potassium and nitrogen, and Kruskal-Wallis test for non-normal data - carbon, calcium, iron, magnesium, manganese, phosphorus, and zinc). Boxes show median and first and third quartiles, with whiskers and points distinguished at 1.5 times the interquartile range.

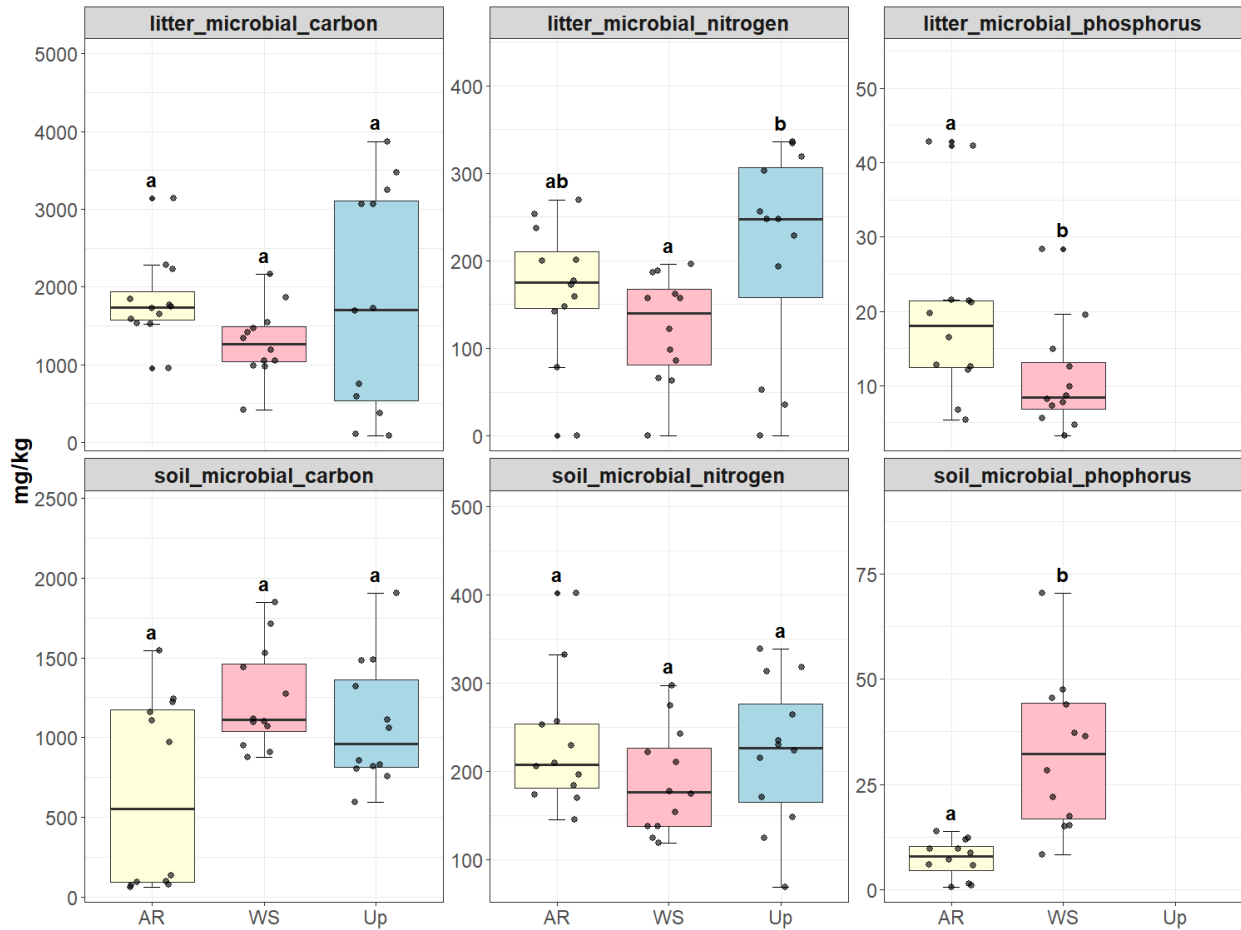


Figure 7. Concentrations of C, N, and P microbial biomass ($\mu\text{g g}^{-1}$ dry mass) of soil and litter in the three forest types: the Ancient River Terrace Forest (AR), the White Sand Forest (WS), and the Upland Forest (Up). Letters indicate statistically significant differences in microbial biomass between forest types at $p < 0.05$, $N=36$. ANOVA test for normal data (litter microbial carbon, soil microbial phosphorus, and nitrogen) and Kruskal-Wallis test for non-normal data (soil microbial carbon, litter microbial nitrogen, and litter microbial phosphorus). Boxes show median and first and third quartiles, with whiskers and points distinguished at 1.5 times the interquartile range.

Microbial biomass (soil and litter), used here as a proxy for microbial activity, showed significant differences between forest types only for soil phosphorus, which was higher in the White Sand Forest compared to the Ancient River Terrace Forest (no data available for the Upland Forest). For carbon and nitrogen, significant differences were observed only in litter microbial biomass, with the upland forest exhibiting the highest values and the white sand forest the lowest. However, for most microbial biomass parameters measured across soil and litter, differences between forest types were not statistically significant (Fig. 7).

3.2 Identification of drivers of BVOC and GHG fluxes

3.2.1 Principal Component Analysis

A PCA of soil and litter characteristics and microbial biomass, and gas fluxes (BVOC and GHG) indicated that PC1 and PC2 axes accounted for 48.5% of the data variation (Fig. 8). The first axis explained 31.6% and the second 12.6% (Table 3). The PCA grouped forest types into two distinct groups: Ancient River Terrace and Upland Forests showed considerable overlap, with lower fluxes linked to litter characteristics, soil and litter microbial biomass, CO₂, and soil pH. In contrast, the White Sand Forest formed a separate group with higher fluxes associated with soil temperature, moisture, and elevated levels of phosphorus, magnesium, and potassium.

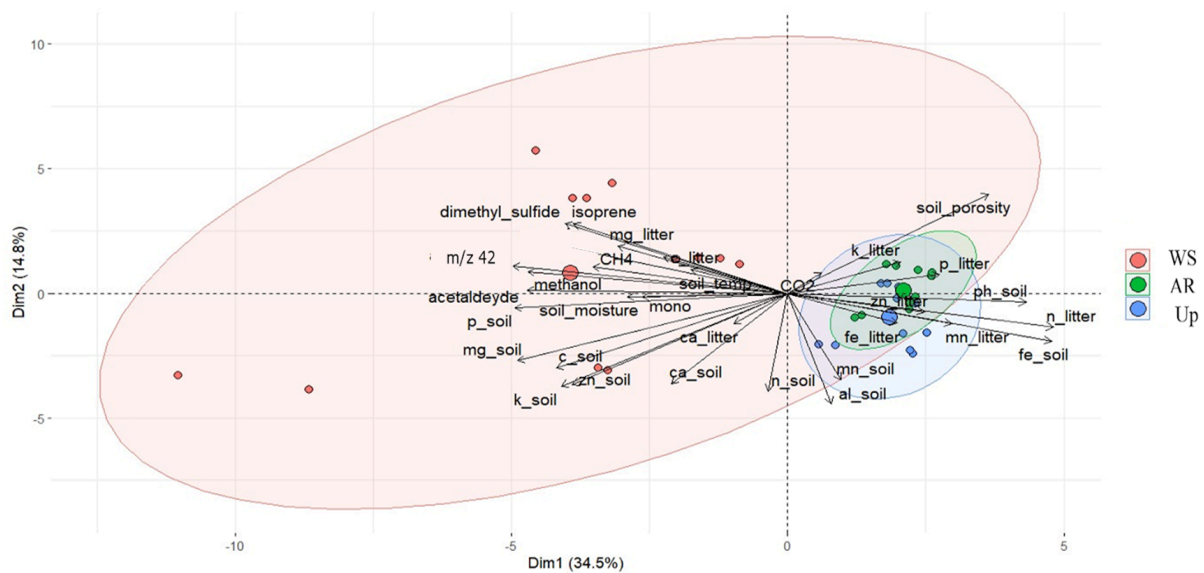


Figure 8. PCA wherein the vectors reflect their correlation with the variables, and the colored circles represent the average PCA score related to each ambient. The analyzed variables are BVOCs (methanol, m/z 42, acetaldehyde, DMS, isoprene and monoterpenes), greenhouse gases (methane and CO₂), soil characteristics (carbon, nitrogen, phosphorus, potassium, calcium, magnesium, aluminum, iron, zinc, manganese, pH, temperature, and soil moisture), litter characteristics (carbon, nitrogen, calcium, magnesium, potassium, iron, zinc, manganese), and soil and litter microorganism dynamics (soil microbial nitrogen, soil microbial phosphorus, soil microbial carbon, litter microbial nitrogen, litter microbial phosphorus and litter microbial carbon). AR = Ancient River Terrace Forest, Up = Upland Forest, WS = White Sand Forest

Table 3. Percentage correlation values extracted from PCA; Fig. 8.

	PC1		PC2
Soil iron	83.480	Isoprene	52.345
Litter nitrogen	83.440	Dimethyl sulfide	51.718
Litter manganese	52.649	Litter Iron	21.662
Litter phosphorus	46.040	Soil moisture	20.658
Soil magnesium	-79.524	Soil carbon	-63.818
Methanol	-80.097	Soil zinc	-64.436
Soil phosphorus	-83.184	Soil aluminum	-71.054
m/z 42	-84.917	Soil nitrogen	-71.553

3.2.2 Linear regression models for different forest types

We used linear regression models (referred to as linear models) to better understand the relationships between predictor variables and fluxes, as identified by the PCA analyses. Flux predictors showed substantial variation between the forest types (Fig. S3a, b, S4a, b, and S5a, b, in Section 4; supplementary material). Model comparisons for each forest type revealed similarities between Ancient River Terrace Forest and Upland Forests (Table 4). In contrast, the White Sand Forest was distinct (Table 4), as also shown by the PCA analysis.

In the Ancient River Terrace Forest, linear models for gas fluxes and predictor variables showed coefficients of determination (R^2) above 0.8 for methanol, acetaldehyde, isoprene, and monoterpenes (Table 4). The most important nutrients for predicting gas fluxes from this forest type were potassium, manganese, magnesium, iron, carbon, and phosphorus. The linear models

for monoterpenes had soil microbial biomass carbon and litter potassium as predictors. The linear models for GHG had soil temperature, soil moisture, and litter nutrients as predictors.

Table 4. Multiple linear regression models with soil and litter characteristics and microbial biomass as predictors of gas fluxes in the three forest types – Ancient River Terrace Forest (AR), Upland Forest (Up), and White Sand Forest (WS). B = unstandardized coefficients. CI = confidence interval. f^2 = Cohen’s f^2 effect size. R^2 = R-squared value. R^2_{adj} = Adjusted R-squared value. N = 36.

Forest Type	Gas	R^2/R^2_{adj}	Predictor Variable	B	95% CI	P	f^2
AR	Methanol	0.839/0.803	Soil potassium	0.03	0.02; 0.04	< .001	1.92
			Litter nitrogen	0.02	0.01; 0.03	< .001	2.31
	Acetaldehyde	0.629/0.547	Soil iron	0.00	-0.00;0.00	0.059	0.12
			Soil manganese	0.004	0.00;0.00	0.004	1.58
	Dimethyl sulfide	0.574/0.479	Soil magnesium	0.006	0.00;0.00	0.2	0.059
			Litter magnesium	-0.004	-0.01;0.00	0.013	1.06
	Isoprene	0.960/0.951	Soil Phosphorus	0.00	0.00; 0.00	< .001	15.03
			Litter Phosphorus	0.06	0.04;0.07	< .001	9.38
	Monoterpenes	0.811/0.769	Soil microbial carbon	0.00	0.00; 0.00	0.084	4.26
			Soil Moisture	0.03	0.000; 0.05	< .001	0.05
	Methane	0.276/0.203	Soil moisture	0.00	0.00; 0.00	0.0824	0.5
			Litter carbon	0.00	0.00;0.00	0.024	0.32
	CO ₂	0.685/0.615	Soil temperature	9.054	3.10; 15.00	0.007	1.232

Up

		Litter phosphorus	-87.94	-156.2; -19.6	0.017	0.943
Methanol	0.617/0.532	Soil potassium	0.61	0.18; 1.0	0.010	0.17
		Soil microbial nitrogen	0.00	0.00; 0.00	0.006	1.44
m/z 42	0.689/0.602	Soil potassium	0.00	0.00; 0.00	0.002	0.73
		Soil microbial nitrogen	0.00	0.00; 0.00	0.006	1.50
Acetaldehyde	0.608/0.521	Soil iron	0.00	0.00; 0.00	0.14	0.17
		Litter microbial carbon	0.02	0.00; 0.00	0.006	1.38
Dimethyl sulfide	0.747/0.690	Litter microbial carbon	0.00	0.00; 0.00	<0.001	1.10
		Litter microbial nitrogen	0.00	0.00; 0.00	0.003	1.85
Isoprene	0.774/0.724	Soil carbon	0.00	0.00; 0.00	0.007	1.25e-03
		Liter magnesium	0.00	0.00; 0.00	< .001	5.94
Monoterpenes	0.792/0.746	Soil potassium	1.6	0.94;2.3	< .001	0.96
		Litter microbial nitrogen	0.00	0.00;0.00	< .001	2.86
Methane	0.888/0.863	Soil carbon	0.231	0.00; 0.00	0.043	0.06
		Soil moisture	0.00	0.00; 0.00	< .001	7.91
CO ₂	0.626/0.543	Litter microbial nitrogen	0.00	0.01; 0.06	0.025	0.25
		Litter microbial carbon	0.00	0.00;0.00	0.006	1.43

WS	Methanol	0.828/0.790	Soil temperature	-3.0	-4.4, -1.5	< 0.001	2.60
			Litter phosphorus	-8.4	-16,-0.37	< 0.042	0.62
	m/z 42	0.866/0.837	Soil moisture	0.187	0.09; 0.27	< .001	2.938
			Litter nitrogen	-54.19	-75.90; -32.49	< .001	3.545
	Acetaldehyde	0.653/0.576	Soil moisture	1.368	0.28; 2.45	0.019	1.022
			Litter nitrogen	-327.4	-593.33; -61.59	0.021	0.863
	Dimethyl sulfide	0.763/0.711	Soil temperature	-0.06	-0.09; -0.03	0.003	1.87
			Soil phosphorus	-0.06	-0.09,-0.0 3	0.003	1.36
	Isoprene	0.764/0.712	Soil temperature	-2.3	-3.6; -0.96	0.004	1.70
			Soil phosphorus	-0.05	-0.07; -0.02	0.003	1.54
	Monoterpene	0.857/0.825	Soil moisture	0.13	0.06; 0.20	0.003	0.36
			Litter nitrogen	2.1	1.5; 2.8	< 0.001	5.66
	Methane	0.508/0.399	Soil moisture	0.00	0.0; 0.0	0.027	0.35
			Litter zinc	0.00	0.0; 0.0	0.035	0.69
CO ₂	0.742/0.685	Soil microbial carbon	0.02	0.0; 0.03	0.029	1.07	
		Litter zinc	3.5	1.6; 5,5	0.003	1.81	

For the Upland Forest, linear models for gas fluxes showed R² higher than 0.8 for isoprene and methane (Table 4), and key nutrients for predicting gas fluxes included potassium,

iron, manganese, and carbon. Microbial biomass was significant in predicting gases like methanol and DMS. Acetaldehyde and isoprene shared soil iron and manganese as predictors, while DMS and CO₂ were linked to litter carbon and microbial nitrogen.

In the White Sand Forest, linear models showed R² higher than 0.8 for methanol, m/z 42 and monoterpenes (Table 4). Key nutrient predictors included phosphorus, nitrogen, and zinc. All emitted gases (except CO₂) were influenced by soil temperature or moisture. Soil temperature was inversely related to fluxes of methanol, DMS, and isoprene, while emissions of m/z 42, acetaldehyde, monoterpenes and methane increased with soil moisture.

3.3 Spatial variability within forest types

Figure 9 shows BVOC and GHG fluxes of each Transect, Figure 10 shows soil temperature and soil moisture of each Transect, and Figure 11 illustrates the spatial variability within and between Transects for isoprene and monoterpenes (see supplementary material, Section 8; Fig. S8, S9, and S10 for other gases). In the Ancient River Terrace Forest, BVOC fluxes were generally lower in Transect 1, while GHG fluxes were similar between Transects (Fig. 9); soil temperature was higher in Transect 1, while Transect 2 was slightly wetter (although not statistically significant) (Fig.10). White Sand Forest exhibited the greatest variation between Transects, with the highest BVOC emissions in Transect 2, and significant variations in acetaldehyde, m/z 42, DMS, isoprene, and methanol; monoterpene fluxes showed high variation in emissions and consumption in Transect 1, while Transect 2 had low variation and high emissions; and methanol was emitted in Transect 1 and consumed in Transect 2. In the Upland Forest, significant differences between Transects were noted for acetaldehyde, m/z 42, DMS, and isoprene.

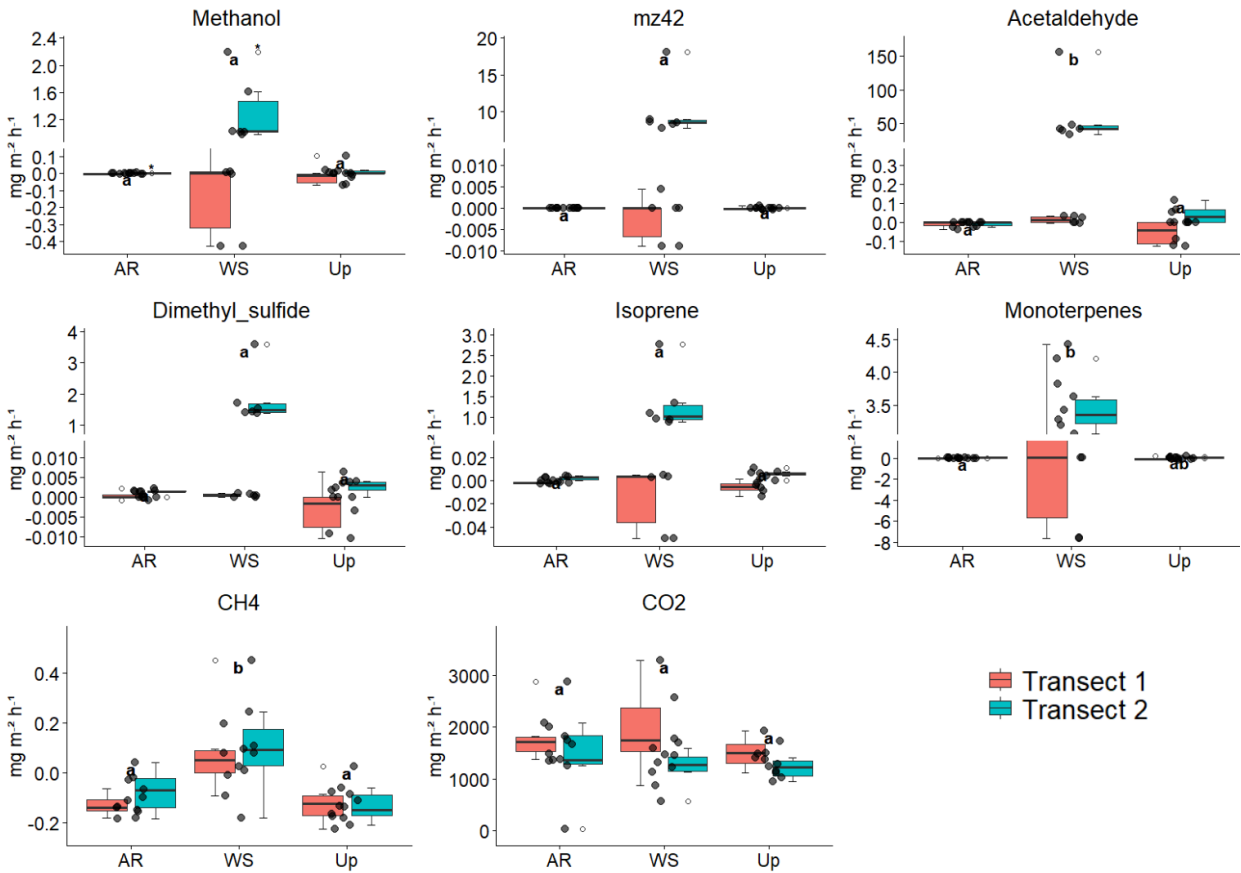


Figure 9. Soil and litter BVOC and GHG fluxes in each forest type - Ancient River Terrace Forest (AR), White Sand Forest (WS), Upland Forest (Up), and Transects within forest types. Letters indicate statistically significant differences in fluxes between the forest types at $p < 0.05$, $N=36$ (Kruskal-Wallis test for non-normal data - BVOC and GHG). Boxes show median and first and third quartiles, with whiskers and points distinguished at 1.5 times the interquartile range. Axes are broken to enhance the visibility of data variation.

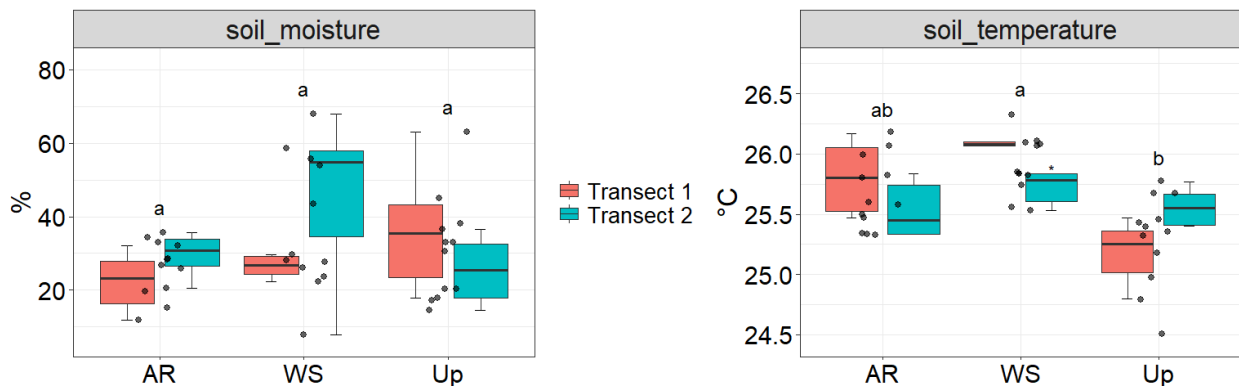


Figure 10. Soil moisture and soil temperature in each forest type - Ancient River Terrace Forest (AR), White Sand Forest (WS), Upland Forest (Up), and Transects within forest types. Letters indicate statistically significant differences in fluxes between the forest types at $p < 0.05$, $N=36$ (ANOVA test for normal distribution). Boxes show median and first and third quartiles, with whiskers and points distinguished at 1.5 times the interquartile range. Axes are broken to enhance the visibility of data variation.

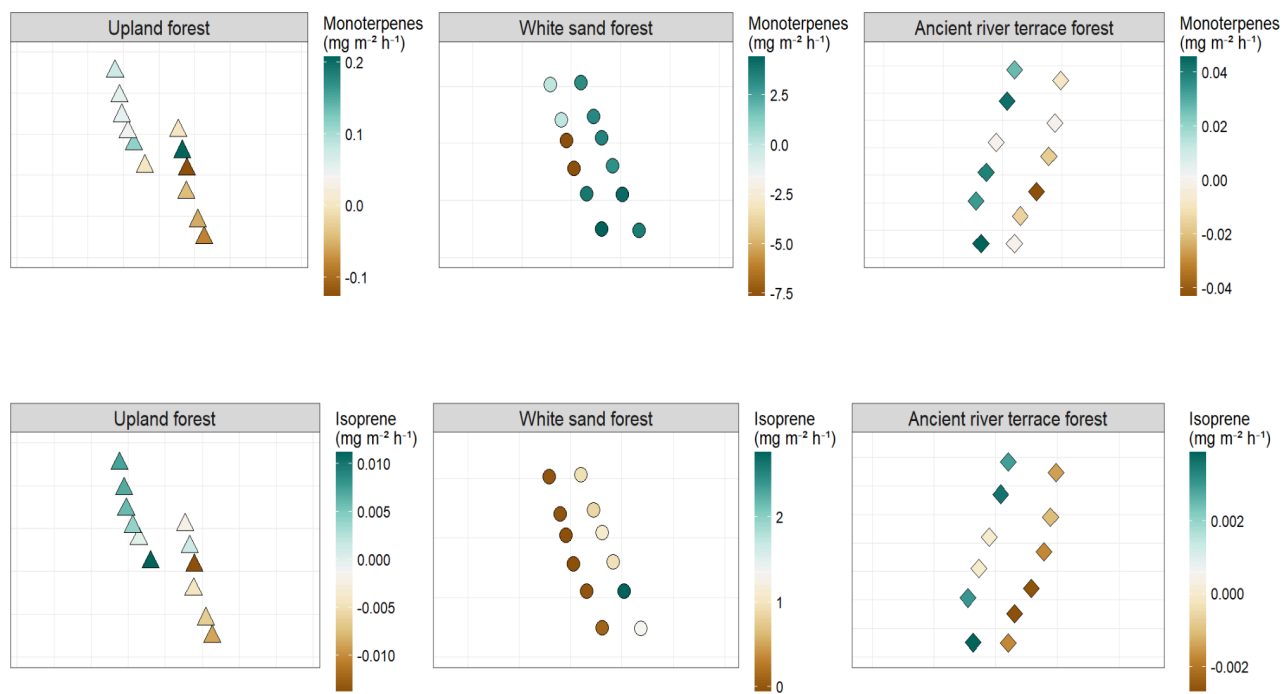


Figure 11. Map of the sampling points visualizing the spatial heterogeneity of BVOC fluxes in each of the three forest types. Each Transect, sampled on different but subsequent days, contains six sampling points, totaling 12 measurement points per forest type. Left - Transect 1; Right - Transect 2. Mono = Monoterpenes. The gas flux data are expressed in $\text{mg m}^{-2} \text{h}^{-1}$.

4. Discussion

Previous studies investigating tropical soil BVOC fluxes using in-situ measurements and incubation (Bourtsoukidis et al., 2018) and fertilization experiments (Llusià et al., 2022) showed that fluxes were higher than previously anticipated. Although these pioneer studies were very

important to understand soil BVOC fluxes from the tropics, our study took one step further by investigating soil-litter BVOC and GHG fluxes across Amazonian Forest types. We found that soil-litter BVOC and GHG fluxes changed across forest types and were influenced by differences in nutrient content, soil moisture and temperature, and microbial biomass. Given the extensive number of measured variables, we chose to focus our discussion on the key observations related to BVOC and methane fluxes and their drivers, rather than covering all variables and fluxes. However, since these variables may still be of interest to the reader, detailed analyses are provided in the supplementary material (Section 2). In the following sections, we first compare soil–litter properties among forest types, followed by a comparison of observed BVOC and GHG fluxes both between forest types and with other tropical studies. We then discuss their potential drivers and the broader significance of our findings.

4.1 Differences in soil and litter nutrient contents across forest types

Variations in soil and litter properties were observed among forest types, particularly concerning their total nutrient contents. These total nutrient pools reflect long-term nutrient availability and reservoirs within each ecosystem, shaped by distinct biogeochemical processes. In the Ancient River Terrace Forest, high concentrations of potassium and phosphorus were found in the litter, suggesting strong nutrient retention and efficient cycling within this forest type. This may result from its history of periodic flooding and the presence of allisols—relatively young, nutrient-rich soils (Andreae et al., 2015).

In the Upland Forest, the dominance of soil iron total content is likely related to the intense leaching common in its ferrasols (oxisols), resulting in iron enrichment due to the removal of other nutrients (Mosquera et al., 2024). In addition, the formation of iron oxides can reduce the mineralization of organic matter, promoting iron accumulation in the leaf litter (Li et al., 2023). The White Sand Forest showed differences in soil properties when compared to the other forest types analyzed. Despite its well-documented low fertility (Mendonça et al., 2015; Demarchi et al., 2022) and arenosol characteristics, this forest type displayed unexpectedly high soil phosphorus and carbon total concentrations. Phosphorus total concentrations, for instance, were up to four times higher than in Upland and Ancient River Terrace Forests. This observation

may possibly reflect the role of dissolved organic nutrients in mitigating nutrient limitations (Lange et al., 2024), or the efficient capture and retention of nutrients within the forest's extensive root mats, which can enhance carbon storage and nutrient cycling in structurally analogous ecosystems (Draper et al., 2014). Furthermore, iron total concentrations in the soil of this forest were lower than expected (Cornu et al., 1997), possibly attributed to spatial variability and seasonal dynamics. During the dry season, the low water-retention capacity of sandy soils leads to drought stress, whereas in the wet season, leaching redistributes iron, aluminum, and magnesium (García-Villacorta et al., 2016) – a process that can form cemented horizons, impeding drainage and elevating water tables (Franco & Dezzeo, 1994; Demarchi et al., 2022). Variable iron concentrations, together with wet-season leaching and elemental redistribution, shape the White Sand Forest into a highly distinct environment. In addition, tree species composition could play a role in influencing nutrient stocks in this forest type (García-Villacorta et al., 2016; Gomes Alves et al., 2022).

4.2 Differences in Gas Fluxes across Forest Types

Here, by comparing forest type fluxes, it is important to recognize that our chamber measurements represent the combined (net) flux from both soil and litter. Observed differences between sites may therefore reflect variations in the relative contributions of soil and litter, for example, due to different amounts of litter. Because our measurements do not allow us to separate these sources, we cannot determine to what extent the observed differences are attributable to each component. Consequently, we treat soil and litter together as a single compartment in our analysis, and discuss the differences in soil-litter gas fluxes across forest types.

The White Sand Forest exhibited the highest emissions and consumption of BVOCs and GHGs, accompanied by the greatest chemical diversity in gas fluxes. Plant species endemic to this ecosystem may influence BVOC emission patterns and their chemical speciation: Fine et al. (2004, 2006) showed that tree species adapted to very nutrient-poor sandy soils highly invest in secondary metabolite compounds in defense against herbivory, since leaves are very energetically costly for the plant. This large quantity of secondary compounds can directly

influence litter decomposition rate (Chomel et al., 2016) and probably release gases and various compounds into the soil and water (Caetano, 2022).

Isoprenoids were emitted in considerable amounts in the White Sand Forest. As isoprenoids are not expected to be emitted from soil (Bach & Rohmer, 2013; Asensio et al., 2008), the observed high emissions might indicate contributions from the activity of microorganisms living in the soil and litter (Carruthers & Lee, 2021; Hernandez-Arranz et al., 2019). In addition, it is important to note that, although emissions in this study are expected to come from soil and litter, the contribution of root emissions cannot be ruled out, as the main source of isoprenoids is expected to be the plant metabolism (Pulido et al., 2012; Thulasiram et al., 2007).

A previous study on experimental rainforest soils - similar to Upland Forest soils - showed BVOC soil uptake (under wet conditions) primarily for isoprenoids, carbonyls, and alcohols, as well as soil emissions of DMS and carbonyl compounds such as acetaldehyde and acetone (Pugliese et al., 2023). Our Upland Forest isoprene fluxes exhibited lower soil uptake ($-0.005 \text{ mg m}^{-2} \text{ h}^{-1}$) compared to the increased uptake fluxes under drier conditions ($\sim -2.38 \text{ mg m}^{-2} \text{ h}^{-1}$) observed by Pugliese et al. (2023). This lower isoprene uptake by the soil observed in the Upland Forest likely reflects the combined effects of the amount of plant-emitted isoprene diffusing into the soil and the subsequent microbial assimilation and soil physical adsorption. These processes are strongly influenced by forest structural attributes and underlying soil properties, which are likely different between this study and Pugliese et al. (2023). Additionally, the higher atmospheric isoprene concentrations reported by Pugliese et al. (2023) compared with those in this study may help explain the greater soil isoprene uptake observed in their work.

In general, our Upland and Ancient River Terrace Forests showed lower average emissions and uptake than those reported by Pugliese et al. (2023). A study focusing on methanol fluxes in cropland soils observed values ranging from 0.53 to $2.93 \text{ mg m}^{-2} \text{ h}^{-1}$ (Liu et al., 2024), which are higher than those observed in our study in Upland and Ancient River Terrace Forests but comparable to the White Sand Forest fluxes ($0.61 \pm 0.81 \text{ mg m}^{-2} \text{ h}^{-1}$). These higher emissions in crop soils can likely be attributed to factors such as crop species, tillage, fertilization, and irrigation, which can all influence BVOC emission rates; whereas the high

methanol emission observed in our study could be related to the root growth of White Sand Forest's extensive root mats, although future studies are necessary to confirm this hypothesis.

The highest DMS emissions were observed in the White Sand Forest ($\sim 1.10 \pm 1.14 \text{ mg m}^{-2} \text{ h}^{-1}$), which were higher than the DMS emission of $5.76 \mu\text{g m}^{-2} \text{ h}^{-1}$ reported by Jardine et al. (2015) for Amazon soils. However, while an interesting observation, it is important to note that the high magnitude of DMS fluxes presented here might partly be caused by a potential agglomerate of acetaldehyde (mass 45) with water, resulting in the same mass as DMS (63), suggesting that future studies should make use of techniques that differentiate these compounds.

A compound with a mass-to-charge ratio (m/z) of 42 was observed in the White Sand Forest, but its identity could not be confirmed due to technical limitations (Dunne et al., 2012). This m/z 42 is frequently attributed to acetonitrile, a known biomass burning marker primarily associated with anthropogenic sources (Huangfu et al., 2021). However, since it can also be emitted by microorganisms (Raio et al., 2020), it is possible that the microbial communities of the White Sand Forest contributed to potential acetonitrile (m/z 42) emissions.

Methane uptake was observed in the Upland ($-0.12 \text{ mg m}^{-2} \text{ h}^{-1}$) and Ancient River Terrace Forests ($-0.10 \text{ mg m}^{-2} \text{ h}^{-1}$), whereas emissions were observed in the White Sand Forest ($0.12 \text{ mg m}^{-2} \text{ h}^{-1}$). In another central Amazonia site, Upland Forest methane fluxes of similar magnitude were observed (-0.02 to $-0.09 \text{ mg m}^{-2} \text{ h}^{-1}$) (van Asperen et al., 2020). However, the discrepancy in White Sand Forest fluxes, with uptake reported in their study (-0.38 to $-0.25 \text{ mg m}^{-2} \text{ h}^{-1}$) and emissions observed here, can likely be attributed to the high spatial variability characteristic of white sand forest ecosystems.

4.3 Drivers of Soil and Litter Gas Fluxes

4.3.1 Soil moisture and soil temperature as drivers of soil-litter gas fluxes

A PCA was initially conducted to identify variables that might differentiate forest types and their gas fluxes. However, due to the overlapping ellipses and limited discrimination capacity, linear models were applied to further explore the findings. These models highlighted

soil temperature and moisture as key physical drivers across all three forest types, aligning with observations in other ecosystems (Trowbridge et al., 2020; Pugliese et al., 2023; Liu et al., 2024).

Nevertheless, before evaluating these key drivers, it is important to recognize that external factors inevitably influenced soil moisture and temperature. Although transects were consistently measured at the same time (08:00–10:00 local time), they were conducted on consecutive days under varying weather conditions. In addition, while radiation was not quantified during the campaign, its effects should be considered. Despite the use of opaque chambers, the chamber surrounding environment - and thus the prevailing BVOC flux dynamics - may still be modulated by incoming radiation. These factors are particularly important in the White Sand Forest, where the open canopy, short stature, and shallow water table (Adeney et al., 2016; Rossetti et al., 2019) promote highly dynamic conditions, reflected in the strong variability of soil temperature and moisture between consecutive transects. While on one hand these factors represent a limitation, it also allowed us to explore the influence of external factors on gas fluxes within the same forest type, such as the rain event which will be discussed hereafter.

In addition to the complicating external factors, we should also note that high soil moisture values were generally accompanied by lower soil temperatures, thereby complicating separating individual drivers. A similar pattern was observed by van Asperen et al., (2024) who showed that the direct effects of temperature and moisture are difficult to differentiate under tropical field conditions, where temperature variation is usually small. Furthermore, it should be noted that this study is additionally limited by a small temporal dataset. Thus, although some BVOC fluxes showed a positive association with soil moisture and a negative association with soil temperature, distinguishing the primary driver remains difficult. Nevertheless, despite the limitations described above, we discuss the expected effects of soil moisture and temperature on our fluxes below.

We observed a negative relationship between temperature and certain BVOC fluxes, but we interpret this as an indirect effect mediated by the previously described link to soil moisture. In general, elevated temperatures are expected to enhance both BVOC emissions and biological uptake, as described by Baggesen et al. (2022). If the temperature sensitivity of uptake exceeds

that of emission, this may result in reduced net emissions or even a net sink for BVOCs (Asensio et al., 2007; Peñuelas et al., 2014; Jiao et al., 2023).

For soil moisture, we found a clear relation with most of BVOC fluxes, especially for the White Sand Forest. The effects of moisture on BVOC fluxes were also described by Pugliese et al. (2023) who reported that rainforest soils acted as net BVOC sinks under moist conditions and as net BVOC sources under dry conditions. In the Upland Forest, we observed a similar pattern as Pugliese et al (2023), with the wetter Transect showing BVOC consumption while the drier Transect showed emissions. The White Sand Forest showed even stronger inter-Transect differences, with high BVOC emissions observed in the wetter Transect, and low emissions and uptake observed in the drier Transect. The pronounced differences between these Transects suggest that the rainfall event preceding Transect 2 was an important driver of the observed variation. A similar hypothesis was posed by Bourtsoukidis et al. (2018), who also observed that sesquiterpene emissions from Upland Forest soils in the dry season (after a rain event) were comparable to those from vegetation. In addition to increasing soil moisture, the physical impact of rainfall may trigger short-term emission bursts (Miyama et al., 2020). As we observed substantially high isoprene, monoterpenes, and acetaldehyde emissions in Transect 2 of the White Sand Forest, we argue that these observed BVOC emissions represent a burst induced by the preceding rainfall event. Similar observations have been reported by Greenberg et al.,(2012), who observed increased BVOC emissions during and immediately after rainfall in a Ponderosa pine plantation, and by Jardine et al. (2015), who reported a peak in soil DMS emissions following rainfall.

Overall, we hypothesize that the rainfall event induced an emission burst, mostly explaining the elevated BVOC fluxes observed in that transect. Nevertheless, interactions between soil temperature, soil moisture, and BVOC fluxes remain highly complex across heterogeneous forest types, with external drivers contributing to strong spatial and temporal variability and making their combined effects difficult to separate under natural field conditions.

4.3.2 Forest Type-specific Drivers of Soil-litter Gas Fluxes

We observed that drivers of soil and litter gas fluxes varied across forest types, reflecting their unique environmental conditions and nutrient dynamics. In general, Ancient River Terrace and Upland Forests showed many similarities in the predictors of certain gases, while other drivers were found for the White Sand Forest. Here, we focus on the key factors influencing gas fluxes: soil nutrients, microbial biomass, and their interactions with environmental conditions.

Soil potassium was found to be a significant factor influencing Ancient River Terrace and Upland Forest fluxes, being identified as a predictor of methanol and monoterpenes. In addition, it was also identified as a predictor of m/z 42 fluxes in the Upland Forest. Although we did not find studies directly relating BVOCs and GHGs fluxes to soil potassium content, potassium is an essential macronutrient for plant growth and metabolism. Its availability is known to affect plant physiological processes (Wang et al., 2013), and its cycling within the soil environment, often mediated by microbial activity, influences potassium's uptake by plants (Mazahar & Umar, 2022). These plant- and soil-mediated processes can, in turn, indirectly influence the BVOC production and release observed within the soil-litter compartment of our study, by affecting the overall ecosystem health and the quality of organic matter available for decomposition.

In the Upland Forest, our methane consumption fluxes correlated well with soil carbon (in conjunction with soil moisture, as mentioned previously). Soil organic carbon is known to play an important role in supporting methanotrophic bacteria, which are responsible for methane oxidation (Lee et al., 2023); therefore, we suggest that the total soil carbon observed in our study might affect methane uptake through a similar process. Phosphorus, like carbon, is a key nutrient in the soil and significantly affected BVOC soil-litter fluxes, especially for methanol in the White Sand Forest. The relationship between phosphorus and BVOC emissions is well documented for plants since the availability of phosphorus can influence the production and emission of BVOCs (Ndah et al., 2022). However, some fertilization studies have also shown that increasing soil nutrient status (nitrogen, phosphorus, and potassium) can modify pH levels, affecting microorganisms and their health state (Stotzky et al., 1976), which directly or indirectly promotes or inhibits soil BVOC fluxes (Liu et al., 2024; Raza et al., 2017). Our findings with bi-directional soil-litter fluxes in the White Sand Forest are consistent with these previous studies showing increase or inhibition of soil BVOC fluxes with soil nutrient content.

Interestingly, our results suggested that lower phosphorus levels were associated with higher isoprene emissions. The mechanisms behind this relationship remain unclear. However, studies on soil fertilization in tropical forests by Llusà et al. (2022) found that phosphorus fertilization is less efficient than nitrogen fertilization in increasing monoterpene and sesquiterpene emissions (they did not find isoprene emissions). They observed that emissions increased when the soil was fertilized only with nitrogen - consistent with a phosphorus-limited system - because excess nitrogen stimulates the enzymes responsible for producing monoterpenes and sesquiterpenes. Conversely, the addition of phosphorus likely redirected this nutrient toward plant growth, resulting in lower emissions of monoterpenes and sesquiterpenes in the phosphorus-fertilized plots compared to those fertilized with nitrogen. As in this study there was no fertilization or a controlled environment, we cannot draw similar conclusions. However, our findings provide valuable insights into the possible interactions between phosphorus, nitrogen, and soil-litter BVOC fluxes in tropical ecosystems. These observations align with previous studies on the influence of soil nutrients (Liu et al., 2024; Llusà et al., 2022) and we suggest future soil fertilization studies to explore these relationships across soil and forest types in Amazonia.

For the Upland Forest, it was found that microbial biomass was a significant driver for almost all soil-litter fluxes, except for isoprene and methane. This aligns with previous studies that have identified microbial biomass as an important driver for soil-litter gas fluxes (Leff & Fierer, 2008; Mancuso et al., 2015; Tang et al., 2019). For example, research on soil organic matter degradation by Lehnert et al. (2024) demonstrated that it is an important source of DMS emissions, highlighting the role of microorganisms associated with decomposition. Jardine et al., (2015) point out that DMS emissions in Amazonian soils are related to microbial processes, a relationship also reported in litter studies by Kesselmeier and Hubert (2002). DMS can be produced in anaerobic environments, such as saturated soils or lakes (Lehnert et al., 2024). This may explain the high emissions observed in transect 2 (wetter and more saturated) of the White Sand Forest, where conditions favorable to anaerobic processes are common and frequently linked to the production of sulfur compounds such as DMS. In contrast, in the drier transect 1 of the upland forest, DMS consumption was observed, suggesting the occurrence of microbial uptake processes. In fact, another study has shown that bacteria can consume carbon from DMS as an energy source in soil and lake sediments (Eyice et al., 2015). Therefore, the observed

uptake may be the result of microorganisms utilizing the carbon present in DMS as an energy source, leading to uptake rather than production. This dual role of microorganisms - as both producers and consumers of DMS - highlights the complexity of sulfur cycling in terrestrial ecosystems.

Based on the limited number of studies investigating the relationship between microorganisms and BVOC dynamics, it has been shown that some Proteobacteria, Actinobacteria, and Firmicutes can produce isoprene (Kuzma et al., 1995; McGenity et al., 2018). *Bacillus subtilis* can produce isoprene in response to stress; however, the mechanism is still not clear (McGenity et al., 2018). Some studies have shown that reduced microbial diversity, whether in soil (Abis et al., 2020; Sillo et al., 2024) or associated with plant surfaces (Saunier et al., 2020), can increase BVOC fluxes and alter the chemical composition of emitted compounds. Although microbial community data were unavailable in this study, we suggest that potential differences in microbial diversity have influenced emission and consumption patterns. Therefore, we strongly recommend that future studies investigate gas flux measurements with microbial community analyses to better understand these dynamics.

4.4 The relevance of White Sand Forest ecosystems

This study showed large variability across forest types and unexpectedly high BVOC emissions from the White Sand Forest. In general, relatively few studies have been performed on White Sand Forests, which is partly explained by the challenging conditions of this forest type, such as flooding and extreme temperatures (Adeney et al., 2016). In addition, the scattered patches of differentiated vegetation distributed within extensive Upland Forests (Demarchi et al., 2022) can make access to these sites even more difficult and require specific infrastructure for data collection. While our observations constitute one of the first characterizations of BVOCs in this unique forest type, their capacity to capture the full spectrum of extreme conditions is inevitably limited by the short temporal coverage of the dataset. To the best of our knowledge, only one study has provided data on BVOC fluxes with soil incubation lab measurements (Bourtsoukidis et al., 2018), and another measuring GHGs in-situ (van Asperen et al., 2020) in White Sand Forests. Despite representing only 5% of the Amazon basin area (Adeney et al., 2016) and 8% of the Reserve of this study (Demarchi et al., 2022), White Sand Forests are

extremely important environments. Their sandy, nutrient-poor soil type has created a challenging ecosystem for plant growth (Fine & Baraloto, 2016), and this unique condition has selected specialized flora and fauna adapted to thrive in these ecosystems (Adeney et al., 2016; Demarchi et al., 2022). This high level of endemism contributes significantly to the overall biodiversity of the Amazon Basin (García-Villacorta et al. 2016). Moreover, White Sand Forests have been shown to play a crucial role in the chemistry of dissolved organic matter (DOM) in Amazonian blackwater rivers, linking terrestrial ecosystem processes to aquatic biogeochemistry (Simon et al., 2021). Our results suggest a stronger link between White Sand Forest gas fluxes and physical factors (more than other forest types), which indicates a possible sensitivity to upcoming climate extremes. Although Costa et al. (2023) did not focus specifically on the White Sand Forest, they showed that regions of the Amazon with shallow water tables - such as the White Sand Forests - can act as hydrological refuges during droughts. In these areas, higher productivity under dry conditions may maintain relatively stable carbon dynamics, presenting a contrasting response to the substantial carbon losses typically observed in deep water table Upland Forests, during drought. In addition, a recent study reported high atmospheric isoprene concentrations in the northwestern Amazon throughout most of the year (Wells et al., 2022) - a region characterized by extensive and continuous White Sand Forest cover (Borges et al., 2014). In this context, as White Sand Forests play a critical role in regulating the carbon cycle and maintaining Amazonian biodiversity (Rossetti et al., 2019), our study suggest that White Sand Forests may represent significant source and sink of BVOCs, together, this highlights the need to better integrate White Sand Forests into future BVOC and GHG flux studies and Earth System modeling.

5. Summary and future directions

Multiple interconnected factors influence BVOC and GHG soil-litter fluxes in the central Amazon. This study highlights the significant roles of soil and litter properties, as well as microbial biomass, in driving these fluxes, with distinct patterns observed across forest types. Given the complexity of the mechanisms influencing BVOC and GHG fluxes, future studies should prioritize microbial activity and diversity, along with diurnal and seasonal cycles, to better identify the key drivers of emission and consumption in these diverse forest ecosystems. In addition, it is important to note that this research serves as a pilot study aimed at scoping out general trends, and many sampling issues can be addressed in future work. For instance, utilizing

a PTR-ToF-MS could alleviate the challenges associated with measuring acetaldehyde, DMS and m/z 42. Longer sampling periods, ideally continuous, would allow for capturing daily variations in emissions.

Surprisingly, despite being the least fertile and diverse forest type, the White Sand Forest exhibited the highest uptake and emission fluxes. This is likely due to intrinsic environmental factors, such as soil temperature and moisture, influencing microbial activity and gas fluxes, as well as the unique vegetation composition of this ecosystem. Furthermore, external factors, such as the preceding rainfall event, may have contributed to very high emissions, potentially reflecting short-term post-rainfall pulses that would have a low or moderate effect when averaged over longer periods that capture the full range of environmental conditions in these ecosystems. Therefore, future studies extending the measurement duration would provide a clearer understanding of how rainfall events influence average soil BVOC emissions. Finally, despite their limited spatial extent in Amazonia, White Sand Forests warrant further research to elucidate their ecological processes and their influence on atmospheric dynamics. Their high BVOC fluxes may substantially affect key physical and chemical processes in the atmosphere, with potential implications for the climate system.

Code/Data availability

All data supporting the findings of this study will be made available in a public repository upon publication.

Authors' contributions

Débora Pinheiro Oliveira, Hella van Asperen, and Eliane Gomes Alves contributed to the development and design of the study, as well as the collection, processing, and statistical analysis of the datasets. Murielli Garcia Caetano and Michelle Robin contributed to field data collection and data analysis. Achim Edtbauer helped design the methodology used in the PTR-QMS and contributed to its calibration improvement. Nora Zannoni, Joseph Byron, Jonathan Williams, Sergio Duvoisin-Junior, and Carla Batista contributed to the chemical analysis of BVOC samples with the GC-TOF-MS and GC-MS. Layon Demarchi and Maria T. F. Piedade contributed to the data analysis of the White Sand Forest. Maria T. F. Piedade, Jochen Schöngart, and Florian Wittmann contributed to the dataset for the initial selection of the points in the PELD-MAUA

project plots where the soil chambers were installed. Rodrigo Augusto Ferreira de Souza contributed to the development of the study. All authors contributed to the writing of the manuscript.

Competing interests

The authors declare that they have no conflict of interest.

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