



1	Changes in PM _{2.5} Peat Combustion Source Profiles with
2	Atmospheric Aging in an Oxidation Flow Reactor
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24 Abstract

Smoke from laboratory chamber burning of peat fuels from Russia, Siberia, U.S.A. (Alaska and Florida), and Malaysia representing boreal, temperate, subtropical, and tropical regions was sampled before and after passing through a potential aerosol mass-oxidation flow reactor (PAM-OFR) to simulate \sim 2- and 7-day atmospheric aging. Species abundances in PM_{2.5} between aged and fresh profiles varied by >5 orders of magnitude with two distinguishable clusters: around 0.1% for reactive and ionic species and mostly >10 % for carbon.

31 Organic carbon (OC) accounted for 58-85 % of PM2.5 mass in fresh profiles with low EC 32 abundance (0.67-4.4 %). After a 7-day aging time, degradation was 20-33 % for OC, with 33 apparent reductions (4-12 %) in low temperature OC1 and OC2 (thermally evolved at 140 and 34 280 °C), implying evaporation of higher vapor pressure semi-volatile organic compounds 35 (SVOCs). Additional losses of OC from 2- to 7-days aging is somewhat offset by the formation 36 of oxygenated organic compounds, as evidenced by the 12-19 % increase in organic mass (OM) 37 to OC ratios. However, the reduction of OM abundances in PM_{2.5} by 3-18 % after 7 days, 38 reconfirms that volatilization is the main loss mechanism of SVOCs. Although the ammonia 39 (NH_3) to PM_{2.5} ratio rapidly diminished with a 2-day aging time, it represents an intermediate 40 profile -not sufficient for completed OC evaporation, levoglucosan degradation, organic acid 41 oxidation, or secondary inorganic aerosol formation.

Week-long aging resulted in an increase to \sim 7–8 % of NH₄⁺ and NO₃⁻ abundances, but with 42 43 enhanced degradation of NH₃, low temperature OC, and levoglucosan for Siberia, Alaska, and 44 Everglasdes (FL) peats. Elevated levoglucosan was found for Russian peats, accounting for 35-45 39 % and 20–25 % of PM_{2.5} mass for fresh and aged profiles, respectively. Abundances of watersoluble organic carbon (WSOC) in PM_{2.5} was >2-fold higher in fresh Russian (37.0 \pm 2.7 %) than 46 47 Malaysian (14.6 \pm 0.9 %) peats. While Russian peat OC emissions are largely water-soluble, Malaysian peat emissions are mostly water-insoluble, with WSOC/OC ratios of 0.59-0.71 and 48 49 0.18–0.40, respectively.

50 Source profiles can change with aging during transport from source to receptor. This study 51 shows significant differences between fresh and aged peat combustion profiles among the four 52 biomes that can be used to establish speciated emission inventories for atmospheric modeling and 53 receptor model source apportionment. A sufficient aging time (~one week) is needed to allow gas-





- 54 to-particle partitioning of semi-volatilized species, gas-phase oxidation, and particle volatilization
- 55 to achieve representative source profiles for regional-scale source apportionment.

- 57 Keywords: fresh and aged source profiles, atmospheric aging, organic mass, organic carbon,
- 58 levoglucosan, oxidation flow reactor (OFR)





59 1 Introduction

60 Receptor-oriented source-apportionment models have played a major role in establishing the weight of evidence (U.S.EPA, 2007) for pollution control decisions. 61 These models, 62 particularly the different solutions (Watson et al., 2016) to the Chemical Mass Balance (CMB) 63 equations (Hidy and Friedlander, 1971), rely on patterns of chemical abundances in different 64 source types that can be separated from each other when superimposed in ambient samples of volatile organic compounds (VOC) and suspended particulate matter (PM). These patterns, termed 65 66 "source profiles," have been measured in diluted exhaust emissions and resuspended mineral dusts for a variety of representative emitters. Many of these source profiles are compiled in country-67 specific source profile data bases (CARB, 2018; Liu et al., 2017; Mo et al., 2016; Pernigotti et al., 68 69 2016; U.S.EPA, 2016) and have been widely used for source apportionment and speciated 70 emission inventories.

71 Chemical profiles measured at the source have been sufficient to identify and quantify 72 nearby, and reasonably fresh, source contributions. These source types include gasoline- and 73 diesel-engine exhaust, biomass burning, cooking, industrial processes, and fugitive dust. Ambient 74 VOC and PM concentrations have been reduced as a result of control measures applied to these 75 sources, and additional reductions have been implemented for toxic materials such as lead, nickel, 76 vanadium, arsenic, diesel particulate matter, and several organic compounds. As these fresh 77 emission contributions in neighborhood- and urban-scale environments (Chow et al., 2002) 78 decrease, regional-scale contributions that may have aged for 2- to 7-days prior to arrival at a 79 receptor gain in importance. These profiles experience augmentation and depletion of chemical 80 abundances owing to photochemical reactions among their gases and particles, as well as 81 interactions upon mixing with other source emissions.

82 Changes in source profiles have been demonstrated in large smog chambers (Pratap et al., 83 2019), wherein gas/particle mixtures are illuminated with ultraviolet (UV) light for several hours 84 and their end products are measured. Such chambers are specially constructed and limited to laboratory testing. A more recent method for simulating such aging is the oxidation flow reactor 85 (OFR), based on the early studies of Kang et al. (2007), revised and perfected by several 86 87 researchers (e.g., Jimenez, 2018; Lambe et al., 2011), and commercially available from Aerodyne 88 (2019a, b). Cao et al. (2019) evaluated the OFR (Aerodyne Research, Inc., Billerica, MA, USA) 89 for potential source emission certification in China, finding that further study and development is





90 required for this purpose. However, Cao et al. (2019) concluded that the OFR could be suitable
91 for source profile aging experiments in support of regional-scale source apportionment, and this is
92 further investigated in this paper for peat combustion.

Peatland fires produce long-lasting thick smoke that leads to adverse atmospheric, climate, ecological, and health impacts. Smoke from Indonesian and Malaysian peatlands is a major concern in the countries of southeast Asia (Wiggins et al., 2018) and elsewhere where it is transported over long distances. Aged peat smoke profiles are likely to differ from fresh emissions, as well as among the different types of peat in other parts of the world.

98 Several ground-based, aircraft, shipboard, and laboratory peat combustion experiments 99 have been carried out to better represent global peat fire emissions and estimate their 100 environmental impacts (e.g., Akagi et al., 2011; Iinuma et al., 2007; Nara et al., 2017; Stockwell 101 et al., 2014; 2016). Most peat fire studies report emission factors (EFs) for pyrogenic gases (e.g., 102 methane, carbon monoxide, and carbon dioxide) and fine particle (PM_{2.5}, particles with 103 aerodynamic diameter <2.5 microns) mass, with a few studies reporting EFs for organic and 104 elemental carbon (OC and EC) (Hu et al., 2018); no information on PM_{2.5} speciated source profiles 105 including elements, ions, and carbon is available.

106 Laboratory peat combustion EFs for gaseous carbon and nitrogen species corresponding 107 with the profiles described here, as well as PM_{2.5} mass and major chemical species (e.g., carbon 108 and ions), are reported by Watson et al. (2019). The PM2.5 speciated source profiles derive from 109 six peat fuels collected from Odintsovo, Russia; Pskov, Siberia; Northern Alaska and Florida, 110 U.S.A.; and Borneo, Malaysia, representing boreal, temperate, subtropical, and tropical climate 111 regions. Comparisons between fresh (diluted and unaged) and aged (i.e., 2- and 7-days simulated oxidation with an OFR) PM_{2.5} speciated profiles are made to highlight chemical abundance 112 113 changes with photochemical aging. Objectives are to: 1) evaluate similarities and differences 114 among the peat source profiles from four biomes; 2) examine the extent of gas-to-particle oxidation 115 and volatilization between 2- and 7-days of simulated atmospheric aging; and 3) characterize 116 carbon and nitrogen properties in peat combustion emissions.

117 2 Experiment

Peat smoke generated in a laboratory combustion chamber was diluted with clean air by factors of three to five to allow for nucleation and condensation at ambient temperatures (Watson et al., 2012). These diluted emissions were then passed through a potential aerosol mass (PAM)-





OFR (Cao et al., 2019; Watson et al., 2019) in the OFR185 mode without ozone (O₃) injection. 121 The OFR UV lamps were operated at 2 and 3.5 volts with a flow rate of 10 L min⁻¹, assuming an 122 average daily hydroxyl (OH) concentration of 1.5×10^6 molecules cm⁻³ to translate OH exposures 123 124 (OH_{exp}) of ~2.6 x 10¹¹ and 8.8 x 10¹¹ molecules-sec cm⁻³ into ~2- and 7-days of photochemical 125 aging. Cao et al. (2019) summarize published ambient OH measurements that span a range around 126 this assumed daily average, indicating that the real-world aging times can differ by factors of two or more. Nevertheless, the 1.5×10^6 molecules-sec cm⁻³ estimate for OH concentration is a de 127 128 facto standard used for OFR aging estimation. 129

Forty smoldering-dominated peat combustion tests were conducted that included three to six tests for each type of peat fuel. The following analysis uses time-integrated (~40–60 minutes) gaseous and $PM_{2.5}$ filter pack samples collected upstream and downstream of the OFR, representing fresh and aged peat combustion emissions, respectively. The sampling configuration is documented in Supplemental Fig. S1 with detailed sampling parameters reported by Watson et al. (2019).

135

2.1 PM_{2.5} mass and chemical analyses

Measured chemical abundances included PM_{2.5} precursor gases (i.e., nitric acid [HNO₃] and ammonia [NH₃]) as well as PM_{2.5} mass and major components (e.g., elements, ions, and carbon). Water-soluble organic carbon (WSOC), carbohydrates, and organic acids that are commonly used as markers in source apportionment for biomass burning were also quantified (Chow and Watson, 2013; Watson et al., 2016).

The filter pack sampling configurations for the four upstream and two downstream channels along with filter types and analytical instrument specifications are shown in Fig. 1. Multiple sampling channels accommodate different filter substrates that allow for comprehensive chemical speciation. The additional upstream Teflon-membrane and quartz-fiber filters were taken for more specific nitrogen and organic compound analyses that are not reported here. The limited flow through the OFR precludes additional downstream sampling.

147 Teflon-membrane filters (i.e., channels one and five in Fig. 1) were submitted for: 1) 148 gravimetric analysis by microbalance with $\pm 1 \mu g$ sensitivity before and after sampling to acquire 149 PM_{2.5} mass concentrations (Watson et al., 2017); 2) filter light reflectance and transmittance by 150 ultraviolet/visible (UV/Vis) spectrometer (200–900 nm) equipped with an integrating sphere that 151 measures transmitted/reflected light at 1 nm interval (Johnson, 2015); 3) 51 elements (i.e., Sodium





[Na] to uranium [U]) by energy-dispersive x-ray fluorescence (XRF) analysis (Watson et al.,
1999); and 4) organic functional groups by Fourier Transform Infrared (FTIR) Spectrometry.

154 Results from UV/Vis and FTIR spectrometry will be reported elsewhere.

155 Half of the quartz-fiber filter (i.e., channels two and six) was analyzed for: 1) four anions (i.e., chloride [CL⁻], nitrite [NO₂⁻], nitrate [NO₃⁻], and sulfate [SO₄⁼]), three cations (i.e., water-156 157 soluble sodium $[Na^+]$, potassium $[K^+]$, and ammonium $[NH_4^+]$), and nine organic acids (including 158 four mono- and five di-carboxylic acids) by ion chromatography (IC) with a conductivity detector 159 (CD) (Chow and Watson, 2017); 2) 17 carbohydrates including levoglucosan and its isomers by 160 IC with a pulsed amperometric detector (PAD); and 3) WSOC by combustion and non-dispersive infrared (NDIR) detection. A portion (0.5 cm^2) of the other quartz-fiber filter half was analyzed 161 162 for OC, EC, and brown carbon (BrC) by the IMPROVE A multiwavelength thermal/optical reflectance/transmittance method (Chen et al., 2015; Chow et al., 2007; 2015b); the IMPROVE A 163 protocol (Chow et al., 2007) reports eight operationally defined thermal fractions (i.e., OC1 to 164 165 OC4 evolved at 140, 280, 480, and 580 °C in helium atmosphere; EC1 to EC3 evolved at 580, 740, and 840 °C in helium/oxygen atmosphere; and pyrolyzed carbon [OP]) that further 166 167 characterize carbon properties under different combustion and aging conditions. Citric acid and sodium chloride impregnated cellulose-fiber filters placed behind the Teflon-membrane and 168 169 quartz-fiber filters, respectively, acquired NH_3 as NH_4^+ and HNO_3 as volatilized nitrate, 170 respectively, with analysis by IC-CD.

Detailed chemical analyses along with quality assurance/quality control (QA/QC) measures are documented in Chow and Watson (2013). For each analysis, a minimum of 10 % of the samples were submitted for replicate analysis to estimate precisions. Precisions associated with each concentration were calculated based on error propagation (Bevington, 1969) of the analytical and sampling volume precisions (Watson et al., 2001).

176 2.2 PM_{2.5} source profiles

Concentrations of two gases (i.e., NH₃ and HNO₃) and 125 chemical species acquired from each sample pair (fresh vs. aged) were normalized by the PM_{2.5} gravimetric mass to obtain source profiles with species-specific fractional abundances. The following analyses are based on the average of 24 paired profiles (shown in Table 1), grouped by upstream (fresh) and downstream (aged) samples for 2- and 7-day aging (i.e., denoted as Fresh 2 vs. Aged 2 and Fresh 7 vs. Aged 7) for each of the six peats with 25 % fuel moisture. Composite profiles are calculated based on the





average of individual abundances and the standard deviation of the average within each group (Chow et al., 2002). Although the standard deviation is termed the source profile abundance uncertainty, it is really an estimate of the profile variability for the same fuels and burning conditions, which exceeds the propagated measurement precision.

To assess changes with fuel moisture content, tests of three sets of Putnam (FL) peats at 60 % fuel moisture were conducted with resulting profiles shown in Supplemental Table S1. A few samples were voided due to filter damage or sampling abnormality, which produced five unpaired (either fresh or aged) individual profiles (Table S2). These profiles are reported as they might be useful for future source apportionment studies.

192 2.3 Equivalence measures

193 The Student *t*-test is commonly used to estimate the statistical significance of differences 194 between chemical abundances. Two additional measures are used to determine the similarities 195 and differences between profiles: 1) the correlation coefficient (r) between the source profile 196 abundances (F_{ii}, the fraction of species i in peat j) divided by the source profile variabilities (σ_{ii}) that quantifies the strength of association between profiles; and 2) the distribution of weighted 197 198 differences (residual [R]/uncertainty $[U] = [F_{i1} - F_{i2}]/[\sigma_{i1}^2 + \sigma_{i2}^2]^{0.5}$) for $< 1\sigma$, 1σ - 2σ , 2σ - 3σ , and 199 $>3\sigma$. The percent distribution of R/U ratios is used to understand how many of the chemical 200 species differ by multiples of the uncertainty of the difference. These measures are also used in 201 the effective variance-chemical mass balance (EV-CMB) receptor model solution that uses the variance (r^2) and the R/U ratio to quantify agreement between measured receptor concentrations 202 203 and those produced by the source profiles and source contribution estimates (Watson et al., 1998).

204 **3 Results and discussion**

205 3.1 Similarities and differences among peat profiles

The first comparison is made between two Florida samples from locations separated by ~485 km (i.e., Putnam County Lakebed and Everglades National Park), representing different geological areas. Table S3 shows that the two profiles have high correlations (r > 0.994), but are statistically different (P < 0.002). Over 93 % of the chemical abundance differences fall within ±3 σ . Statistical differences are not found when combining both fresh Florida profiles (e.g., all Fresh 2 vs. all Fresh 7), resulting in high correlations (r > 0.997) with over 98 % of abundance differences within ± 1 σ and P > 0.5. However, paired comparisons of other combined profiles





show statistical differences with low *P*-values (P < 0.002). These two subtropical profiles should not be combined to compare with other biomes.

215 Similarities and differences in peat profiles by biome are summarized in Table 2. 216 Comparisons are made for: 1) paired fresh vs. aged profiles (i.e., All Fresh vs. All Aged; Fresh 2 217 vs. Aged 2; Fresh 7 vs. Aged 7); 2) different experimental tests (i.e., Fresh 2 vs. Fresh 7); and 3) 218 two aging times (i.e., Aged 2 vs. Aged 7). Equivalence measures show that most of these profiles 219 are statistically different (P < 0.05) but highly correlated (r > 0.97, mostly >0.99). However, 220 statistical differences are not found between the Fresh 2 vs. Aged 2 Malaysian profiles, which may 221 be due to the low number of samples (n=2) in the comparison. Similar to the findings of combining both Florida profiles, fresh Alaskan and Malaysian profiles do not show statistical differences (P 222 223 >0.1).

Compositing profiles by averaging each of the measured abundances may disguise some useful information. For receptor model source apportionment, region-specific profiles are most accurate for estimating source contributions.

Student *t*-tests for the gravimetric $PM_{2.5}$ mass concentrations ($\mu g/m^3$) measured upstream and downstream of the OFR (Table S4) show statistically significant differences (P < 0.05) between fresh vs. aged $PM_{2.5}$ (i.e., Fresh 2 vs Aged 2 and Fresh 7 vs Aged 7). Fresh 2 and Fresh 7 $PM_{2.5}$ mass concentrations are similar, as expected from replicate tests for the same conditions. Increases in some species abundances offset decreased on other abundances, resulting in similar $PM_{2.5}$ levels for some of the fresh vs. aged comparisons.

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3.2 Sum of species to PM_{2.5} mass ratios

234 The sum of the major PM chemical abundances should be less than unity since oxygen, 235 hydrogen, and liquid water content are not measured (Chow et al., 1994; 1996). As shown in Table 236 S5, the sums of elements, ions, and carbon explain averages of \sim 70–90 % of PM_{2.5} mass for fresh 237 profiles except for Russian peat (62–64 %). The sum of species decreased by an average of 6 % 238 and 11 % after 2- and 7-days, respectively. These differences can be attributed to loss of semi-239 volatile organic compounds (SVOCs), although they are offset by formation of oxygenated 240 compounds during aging. This is true for all but Putnam (FL) peat, for which the sum of species 241 explains nearly the same fraction of PM2.5 for the fresh and aged profiles.





242 **3.3** Comparison between fresh and aged profiles

243 Fresh and aged chemical abundances are compared in Fig. 2. Species abundances vary by 244 over 5 orders of magnitude but exhibit two distinguishable clusters: around 0.1 % for reactive and secondary ionic species (e.g., NH_4^+ , NO_3^- , and SO_4^-) and >1 % (mostly >10 %) for carbon 245 246 compounds (e.g., OC fractions and WSOC). While most gaseous NH₃/PM_{2.5} ratios exceed 10 %, 247 HNO₃/PM_{2.5} ratios are well below 1 %. Reactive/ionic species and carbon components are mostly 248 above and below the 1:1 line, respectively, implying particle formation and evaporation after 249 atmospheric aging. Large variabilities are found for individual species as noted by the standard deviations associated with each average. 250

251 Figure 3 shows the ratio of averages between aged and fresh profiles with increasing ratios 252 from 2- to 7-day aging. Atmospheric aging increased oxalate, NO_3^- , NH_4^+ , and SO_4^- abundances 253 (likely due to conversion of nitrogen and sulfur gases [e.g., NO, NO₂, and SO₂] to particles), but 254 decreased NH₃, levoglucosan, and low temperature OC1 and OC2 in most cases. Large variations 255 are found among measured species as ratios (left panels in Fig. 3) range over 7 orders of magnitude 256 for mineral and ionic species. Consistent with Fig. 2 where most carbon compounds are close to 257 but below the 1:1 line, the right panels in Fig. 3 show the reduction of carbonaceous abundances 258 with ratios between 0.1 and 1 with lower ratios after 7-day aging.

Atmospheric aging should not change the abundances of mineral species (e.g., Al, Si, Ca, Ti, and Fe), except to the extent that the PM_{2.5} mass (to which all species are normalized) increases or decreases with aging. Large standard deviations associated with the ratio of averages for mineral species in the left panels of Fig. 3 illustrate variabilities among different combustion tests for the less abundant species.

264 3.4 Carbon abundances

265 3.4.1 Organic carbon and thermally-evolved carbon fractions

Total carbon (TC, sum of OC and EC) constitutes the largest fraction of $PM_{2.5}$ (Table 1), accounting for 59–87 % and 43–77 % of the $PM_{2.5}$ mass for the fresh and aged profiles, respectively. Organic carbon dominates TC with low EC abundance (0.67–4.4 %), as commonly found in smoldering-dominated biomass combustion (Chakrabarty et al., 2006; Chen et al., 2007). The largest OC fractions are high temperature OC3 (15–30 % of $PM_{2.5}$), consistent with past studies for biomass burning emissions (Chen et al., 2007; Chow et al., 2004).





272 Abundances of OC decrease with aging time. Upstream (Fresh 2 and Fresh 7) OC abundances ranged from 58-85 % and decreased by 4-12 % and 20-33 % after 2- and 7-day aging, 273 274 respectively. Part, but not all of this is due to increasing abundances of non-carbon components, particularly nitrogen-containing species. The exception is for Putnam (FL) peat, where the OC 275 abundances (67-72 %) were similar for fresh and aged profiles. OC abundance decreases after 276 277 aging may have contributed to the statistical differences found between fresh and aged PM2.5 mass 278 (Table S4). With the exception of Putnam (FL) peat, the additional 7–22% OC degradation from 279 2- to 7-days implies that much of the OC changes require about a week of aging time.

280 The Student *t*-test for fresh and aged profiles shows statistical differences (P < 0.05) for 281 TC, OC, and low temperature OC1 and OC2, but similarities for OC3 and OC4. While the OC1 282 abundance is 9–18 % for fresh profiles, it decreases to 4–10 % after aging. A similar pattern is 283 found for OC2, with 12-25 % and 9-19 % abundances for the fresh and aged profiles. The 284 exception is Putnam (FL) peats that retained a ~20 % OC2 abundance after aging. High 285 temperature OC3 and OC4 contain more polar and/or high molecular-weight organic components (Chen et al., 2007) that are less likely to photochemically degrade. Further reduction in OC 286 287 abundances (20-33 %) after 7-days is attributed to decreases of the OC1 and OC2 in the OFR as shown in the fresh vs. aged ratios of average abundances (Fig. 3). Large fractions of pyrolized 288 289 carbon (OP of 7-13%) are also found --indicative of higher molecular-weight compounds that are 290 likely to char (Chow et al., 2001; Chow et al., 2004; Chow et al., 2018).

291 3.4.2 Organic mass (OM) and OM/OC ratios

292 Reduction of the "sum of species" and OC abundances from fresh to aged profiles can be 293 offset by the formation of oxygenated organic compounds as the profiles age. Different 294 assumptions have been used to transform OC to organic mass (OM) to account for unmeasured H, 295 O, N, and S in organic compounds (Cao, 2018; Chow et al., 2015a; Riggio et al., 2018). As single 296 multipliers for OC cannot capture changes by oxidation in the OFR, OM is calculated by 297 subtracting mineral components (using the IMPROVE soil formula by Malm et al. (1994)), major 298 ions (i.e., NH4⁺, NO3⁻, and SO4⁼), and EC from PM_{2.5} mass to account for unmeasured mass in organic compounds (Chow et al., 2015a; Frank, 2006). This approach assumes that no major 299 300 chemical species are unmeasured and that the remaining mass consists of H, O, N, and S associated 301 with OC in forming OM.





- Table 3 shows that averaged OM/OC ratios are ~1.3 for fresh profiles and increases by 12– 19 % from 2- to 7-days aging. These fresh OM/OC ratios are consistent with those reported for other types of biomass burning (Chen et al., 2007; Reid et al., 2005). The increased OM/OC ratios with aging are likely due to an increase in oxygenated organics. The OM/OC ratio of 1.20 ± 0.05 for fresh Borneo, Malaysian peat is consistent with the 1.26 ± 0.04 ratio for fresh peat burning aerosol in Central Kalimantan, Indonesia (Jayarathne et al., 2018), both located on the Island of Borneo.
- 309 The highest OM/OC ratios are found for Russian peat, ranging 1.6–1.7 for fresh profiles 310 and increasing to 2.1-2.2 for aged profiles, consistent with formation of low vapor pressure oxygenated compounds in the OFR. Watson et al. (2019) report that the Russian peat fuel contains 311 312 the lowest carbon (44.20 ± 1.01 %) and highest oxygen (38.64 ± 0.78 %) contents among the six peats. The low carbon contents are consistent with the lowest "sum of species" found in Russian 313 314 peat, with 62-64 % and 50-52 % of PM_{2.5} mass for the fresh and aged profiles, respectively. After 315 7-day aging for Siberian peat, the increasing OM/OC ratios from 1.2 ± 0.14 to 1.5 ± 0.18 are 316 similar to the increase from 1.22 to 1.42 reported by Bhattarai et al. (2018).
- 317 **3.4.3**

3.4.3 Water-soluble organic carbon (WSOC)

318 WSOC abundances in PM_{2.5} was >2-fold higher in fresh Russian (37.0 \pm 2.7 %) than 319 Malaysian (14.6 \pm 0.9 %) peat. The 15–17 % WSOC in PM_{2.5} for fresh Borneo, Malaysian peat 320 (Table 1) is consistent with the 16 \pm 11 % from Central Kalimantan, Indonesia peat (Jayarathne et 321 al., 2018).

The WSOC/OC ratios also vary (Table 3), ranging from 0.18–0.64 and 0.31–0.71 for fresh and aged profiles, respectively. Russian peat OC emissions are largely water-soluble, whereas Malaysian peat emissions are mostly water-insoluble, with WSOC/OC ratios of 0.59–0.71 and 0.18–0.40, respectively. Longer aging time results in higher WSOC/OC ratios with 2–13 % and 5–19 % increases for 2- and 7-day aging.

327 3.4.4 Carbohydrates

Bates et al. (1991) found that peat from Sumatra, Indonesia consists of 18–46 % carbohydrate (mainly levoglucosan) relative to total carbon based on nuclear magnetic resonance spectroscopy. Levoglucosan and its isomers (mannosan and galactosan) are saccharide derivatives formed from incomplete combustion of cellulose and hemi-cellulose (Kuo et al., 2008; Louchouarn et al., 2009) and have been used as markers for biomass burning in receptor model





source apportionment (Bates et al., 1991; Watson et al., 2016). These carbohydrate-derived
pyrolysis products undergo heterogeneous oxidation when exposed to OH radicals in the OFR
(Hennigan et al., 2010; Kessler et al., 2010).

336 Only five of the 17 carbohydrates (Table 1) were detected with noticeable variations (e.g., 337 >2 orders of magnitude) in levoglucosan for boreal and temperate peats. Levoglucosan abundances 338 account for 35-39 % and 20-25 % of PM_{2.5} mass for fresh and aged Russian profiles, respectively. 339 On a carbon basis, Table 3 shows that levoglucosan-carbon (with an OM/OC ratio of 2.25) accounts for 43-48 % and 30-35 % of WSOC and 27-28 % and 21-24 % of OC for fresh and 340 341 aged Russian profiles, respectively. These levels are less than the 96 \pm 3.8 % levoglucosan or ~42.7 % of levoglucosan-carbon in OC reported for German and Indonesian peats (linuma et al., 342 343 2007). Elevated levoglucosan is also found for Siberian and Alaskan peats, ranging from 4-18 % 344 in PM_{2.5}. However, the levoglucosan abundances are reduced to 1-4 % for the subtropical and 345 tropical peats. The 7-day aging time resulted in an additional 1-4 % levoglucosan degradation 346 relative to 2 days with the exception of a 9 % reduction for Russian peat.

347 The extent of levoglucosan degradation depends on organic aerosol composition, OH exposure in the OFR, and vapor-wall losses (Bertrand et al., 2018a; 2018b; Pratap et al., 2019). 348 349 Figure 4 shows the presence of 11 % and 7.6 % levoglucosan-carbon for the Russian and Alaskan 350 peats after 2-day aging, in line with a chemical lifetime longer than 2 days. This is consistent with 351 the estimated 1.2-3.9 days of levoglucosan lifetimes under different environments reported by Lai 352 et al. (2014). However, other studies (Hennigan et al., 2010; May et al., 2012; Pratap et al., 2019) found that levoglucosan experiences rapid gas-phase oxidation, resulting in ~1-2 day lifetimes at 353 354 ambient temperatures.

Among the carbohydrates, Jayarathne et al. (2018) reported 4.6 ± 4.0 % of levoglucosan in OC for fresh Indonesia peat. Converting to levoglucosan-carbon in Jayarathne et al. (2018) yields a fraction of 2 %, consistent with findings for Malaysian peat (1.4–2.4 %) in this study.

While the presence of levoglucosan in peat smoke is apparent, its isomer, galactosan was not detectable. Mannosan is detectable in cold climate peats with 1-5 % in PM_{2.5} for the Russian and Alaskan peats and up to 1.3 % for Siberian peat. Apparent degradations from 3.9 to 2.5 % and from 5.0 to 2.1 % in mannosan abundances are found for Russian peat (Table 1) after 2- and 7days, respectively. A 2- to 3-fold reduction in mannosan is also shown after 7 days for the Siberian and Alaskan peats. Similar observations apply to glycerol in Russian peat, ranging 1.9–3.5 % and





1.3-1.7 % in PM_{2.5} for fresh and aged profiles, respectively. Other detectable carbohydrates are galactose and mannitol, typically present at one hundredth of one percent of the levoglucosan abundance.

367 3.4.5 Organic acids

Organic acids have been associated with a mixture of anthropogenic sources, including engine exhaust, biomass burning, meat cooking, bioaerosol, and biogenic emissions. Past studies show the presence of low molecular-weight dicarboxylic acids in biomass burning emissions (e.g., Cao et al., 2016; Falkovich et al., 2005; Veres et al., 2010).

Only four of the ten measured organic acids (Table 1) in their anion forms (i.e., proprionate, oxalate, acetate, and formate) are detectable with variable abundances (<0.02-3.9 %). The largest changes between fresh and aged profiles are found for oxalate, ranging from <0.02-0.43 % of PM_{2.5} for fresh profiles, with \sim 10- to 20-fold increase after 2 days (0.6–1.3 %), and with 1 to 2 orders of magnitude increases after 7 days (1.1–3.9 %). With the exception of Putnam, FL peat (1.1 ± 0.19 %), oxalate accounts for >2.9 % of PM_{2.5} mass after 7 days.

378 Acetate abundances are stable between fresh and aged profiles, mostly in the range of 0.2-379 0.5% except for a 6-fold increase from $0.23 \pm 0.15\%$ (Fresh 7) to $1.5 \pm 2.0\%$ (Aged 7) for Siberian 380 peat with large variability among the tests. Propionate and formate abundances are low (<0.02and <0.5 %, respectively), but increase with aging. Extending the aging time from 2- to 7-days 381 382 resulted in a notable increase in organic acid abundances, consistent with the increases in 383 WSOC/OC ratios (Table 3). By biome, the highest abundances for organic acids in $PM_{2.5}$ are 384 found for aged (Aged 7) Siberian peat, with 3.9 ± 1.4 % oxalate, 1.5 ± 2.0 % acetate, and $0.44 \pm$ 385 0.28 % formate (Table 1).

386 **3.5** Nitrogen species, sulfate, and chloride abundances

Ammonia normalized to $PM_{2.5}$ mass is high for fresh profiles, ranging 17–64 %, except for the low NH₃ content in Russian peat (6–8 %). These abundances are reduced to 3–14 % and 1–7 % after 2- and 7-day aging, respectively. As shown in Fig. 5, most of the NH₃ rapidly diminished after 2 days, with increasing particle-phase NH₄⁺ and NO₃⁻ after 7 days. The highest NH₃ to PM_{2.5} ratios are found for fresh Everglades (FL) peat profiles (51–64 %), ~2–8 fold higher than other peats. These high and low NH₃/PM_{2.5} ratios are consistent with the nitrogen contents in peat fuel: 3.93 ± 0.08 % for Everglades and 1.50 ± 0.52 % for Russian peats (Watson et al., 2019).





394	Ionic abundances are typically <0.5 %, especially in fresh profiles. Abundances of $\rm NH_4^+$
395	in PM _{2.5} are low (0.0005–0.13 %) for fresh emissions, but increase to $0.05-1.0$ % after 2 days and
396	3.4–6.7 % after 7 days, with the exception of Putnam (FL) peat (1.01 \pm 0.05 % NH ₄ ⁺). Extending
397	the aging time from 2- to 7-days results in an increase to $\sim 1-7$ % in NH ₄ ⁺ abundances, in contrast
398	to NH_3 that is largely depleted after 2 days.

399 Figure 5b shows increasing in NO_3^- abundances with aging, 0.04–0.23 % for fresh profiles, 400 increasing to 0.74–2.64 % after 2 days, and to 2.0–8.2 % after 7 days with the exception of Putnam (FL) peat (1.10 \pm 0.18 % NO₃⁻). After 7 days, NH₄⁺ and NO₃⁻ account for ~4–7 % and ~8 % of 401 402 PM_{2.5} mass, respectively, for Siberian, Alaskan, and Everglades (FL) peats. No specific trend is 403 evident for NO_2^- , mostly <0.002 % with ~0.2 % for some fresh Siberian and Alaskan peats. The 404 ratio of gaseous HNO₃ to PM_{2.5} is low, in the range of 0.2–0.5 % without much changes between 405 fresh and aged profiles. HNO₃ created through photochemistry is largely neutralized by the 406 abundant NH₃ in the emissions, resulting in the increasing NH₄⁺ and NO₃⁻ to PM_{2.5} in aged profiles. 407 The reaction of NH₃ with HNO₃ to form ammonium nitrate (NH₄NO₃) is the main pathway for inorganic aerosol formation, owing to low sulfur content in the peat fuels (Watson et al., 2019). 408 SO_4^{-} abundance is low in fresh profiles (0.13–1.4 %), but it increases by 2–3 fold after 2 days 409 except for the Alaskan (0.35–0.46 %) and Everglades (FL) (1.3–1.4 %) profiles. More apparent 410 411 changes are found for 7 days with the largest increase in $SO_4^{=}$ from 0.13 to 1.96 % for the 412 Malaysian peats – indicating formation of ammonium sulfate ([NH4]2SO4). The ion balance shows more NH₄⁺ than needed to completely neutralize NO₃⁻ and SO₄⁼ (Chow et al., 1994). Some NH₄⁺ 413 414 may be present as ammonium chloride (NH₄Cl), however, the abundance of chloride (Cl⁻) is low 415

- 415 (<0.3 %). The large increase in NO_3^- and SO_4^- after 7 days implies that a 2-day aging time is not 416 sufficient to allow the full formation of secondary NH₄NO₃ and (NH₄)₂SO₄.
- 417 3.6 Mass reconstruction

418 Mass reconstruction is applied to understand the changes in major chemical composition 419 between the fresh and aged profiles. As shown in Fig. 6, the largest component in PM_{2.5} is OM, 420 accounting for 94–99 % and 80–95 % of PM_{2.5} mass for fresh and aged profiles, respectively. 421 Although the 7-day aging time increased the OM to OC ratios (by 12–19 %), the abundances of 422 OM in PM_{2.5} are reduced (3–18 %). This indicates that volatilization becomes a significant loss 423 mechanism for SVOCs (Smith et al., 2009). The reduction of OM abundance is also partially due 424 to increased ionic species (i.e., sum of NH₄⁺, NO₃⁻, and SO₄⁼), with low abundances (0.3–1.7 %)





425 in fresh profiles, increasing to 3–16 % after aging. The sum of ionic species accounts for 11–16 426 % of PM_{2.5} mass for the Siberian, Alaskan, Everglades (FL), and Malaysian peats after 7 days, 427 mainly due to the increase in NH_4^+ and NO_3^- as shown in Fig. 5.

Elemental abundances are low (<0.0001 %), mostly below the lower quantifiable limits. Table 1 only lists 34 of the 51 elements (Na to U) detected by XRF. Using the IMPROVE soil formula (assuming metal oxides of major mineral species) yielded 0.07–2.9 % of mineral components.

This study indicates that an aging time of ~2 days represents the intermediate profile,
whereas 7 days represents the profile with adequate residence time to complete the atmospheric
process.

435 **3.7** Changes in source profiles by fuel moisture content

436 The effect of fuel moisture content on source profiles is mostly unknown. The 25 % fuel 437 moisture content selected for this study intends to better simulate the conditions of moderate to 438 severe droughts where most peat fires occur. Increasing fuel moisture content from ~ 25 to 60 % 439 for the three Putnam (FL) peat fuels yielded 12 % higher EFs for CO₂ (EF_{CO₂}), but 12–20 % lower 440 EFs for CO, NO, NO₂, and PM_{2.5} mass (Watson et al., 2019). Tests of fuel-moisture content on 441 profile changes are available for only 2-day aging. Equivalence measures (Table S6) show 442 statistical differences (P < 0.001) between 25 % and 60 % moisture profiles on either fresh or aged 443 profiles with over 93 % of species abundance fall within $\pm 3\sigma$ and high correlations (r > 0.997). While OC abundances in PM2.5 are comparable for the fresh and aged profiles (70-72 %) for 25 444 445 % fuel moisture, a reduction of 18 % OC in PM_{2.5} is found for 60 % fuel moisture (from 82 to 64 446 %) after aging (Table S1). The higher fuel moisture content also reduced WSOC by 6 % and 447 levoglucosan by 1.3 % with <1 % increases for NH_4^+ and organic acids. After aging, the NH_3 to 448 PM_{2.5} ratios reduced from 28 to 5 % and from 20 to 8 % for the 25 % and 60 % fuel moisture, 449 respectively. These results are not conclusive as most measurements are associated with high 450 variabilities.

451 **4 Summary and conclusion**

Fresh and aged peat fire emission profiles from laboratory combustion chamber and potential aerosol mass-oxidation flow reactor (PAM-OFR) for six types of peats representing boreal (Odintsovo, Russia and Pskov, Siberia), temperate (Northern Alaska, USA), subtropical (Putnam County Lakebed and Everglades National Park, Florida, USA), and tropical (Borneo,





Malaysia) biomes are compared. Analyses are focused on the average of 24 paired profiles
grouped by six peats and by fresh vs. aged profiles for 2- and 7-days of simulated atmospheric
aging.

Equivalence measures show that these profiles are highly correlated (r > 0.97, mostly >0.99) but statistically different (P < 0.05) between different biomes, suggesting that these profiles should be used independently for receptor model source apportionment studies in different climate regions.

The sum of chemical species (i.e., elements, ions, and carbon) explains an average of \sim 70– 90 % of PM_{2.5} mass for fresh profiles except for Russian peat (62–64 %), confirming that major PM_{2.5} chemical species are measured. Aging times of 2- and 7-days resulted in an average mass depletion of 6 % and 11 %, respectively. These differences are caused by: 1) loss of SVOCs with aging, as indicated by lower abundances of OC1 and OC2 (evolved at 140 and 280 °C) in the aged profiles; and 2) replacement of the lost OC mass with unmeasured oxygen associated within secondary organic aerosol formation in the OFR.

470 Species abundances in $PM_{2.5}$ between aged and fresh profiles varied by >5 orders of 471 magnitude but exhibited two distinguishable clusters, with reactive/ionic species (e.g., NH_4^+ , $SO_4^=$, 472 oxalate, and HNO₃) constituting 0.1–1 % and carbon compounds (e.g., organic carbon fractions 473 [OC1–OC4], WSOC, and OC) constituting >1 % (mostly >10 %) of PM_{2.5} mass. Most NH₃/PM_{2.5} 474 ratios are >10 % whereas HNO₃/PM_{2.5} ratios are <1 %.

475 Total carbon (TC, sum of OC and EC) is the largest component, accounting for 59–87 % 476 and 43-77 % of the PM_{2.5} mass for the fresh and aged profiles, respectively. With predominant 477 smoldering combustion, the majority of the TC is in OC, with low EC abundances (0.67-4.4 %). 478 Further degradation in OC abundances (7-22 %) from 2- to 7-days implies the incomplete 479 volatilization with short aging time. Different thermal carbon fractions are used to characterize 480 combustion and aging conditions. While most of OC evolved at high temperatures (OC3 at 480 481 °C), losses of low temperature OC1 and OC2 are found, suggesting a shift of gas-particle 482 partitioning of SVOC to gas-phase, where particle volatilization, the loss mechanism, outweighed 483 gas-to-particle conversion.

Formation of oxygenated compounds is pronounced after aging, with organic mass (OM)
to OC ratios increasing by 12–19 % from 2- to 7-days aging. The WSOC abundance in PM_{2.5}





486 varies from 14.6 ± 0.9 % to 51 ± 32 % for fresh Malaysian and Siberian peats, respectively. While 487 levoglucosan accounts for $\sim 1-4$ % of PM_{2.5} mass for fresh subtropical and tropical peats, elevated 488 levels (~10 %) are found for boreal and temperate peats. Increasing the atmospheric aging time 489 from 2- to 7-days results in additional formation of ionic species (e.g., oxalate, NO_3^- , NH_4^+ , and SO₄⁼), but enhanced losses of NH₃, levoglucosan, and low temperature OC1 and OC2. 490

491 Among the four climate regions, Russian peat with the lowest carbon (44 %) and highest 492 oxygen (39%) content, resulted in ~59-71% of WSOC in OC along with the highest levoglucosan 493 $(20-39\% \text{ of PM}_{2.5})$ and lowest NH₃/PM_{2.5} ratios (3-8%). It also yielded the highest oxygenated 494 compounds after aging with OM/OC ratios of 2.1–2.2. This contrasts with Malaysian peats that are mostly water-insoluble (WSOC/OC of 0.18-0.4) with low oxygenated compounds after aging 495 496 (OM/OC ratios of 1.3–1.5). Large increases are found for oxalate abundances from fresh (<0.02– 497 0.43 %) to 7-day aging (1-4%).

498 With the exception of Russian peats, fresh profiles contain high NH₃/PM_{2.5} ratios (17-64 499 %) with low abundances after aging (3-14 % for Aged 2 and 1-7 % for Aged 7). Extending the aging time from 2- to 7-days results in an increase to \sim 7–8 % NH₄⁺ and NO₃⁻ abundances. 500 501 Although the week-long aging time increased the OM/OC ratios, abundances of OM in PM2.5 were 502 reduced by 3-18 % with more degradation after 7 days.

503 Source profiles can change with aging during transport from source to receptor. This study 504 shows significant differences between fresh and aged peat combustion profiles between the four 505 biomes that can be used to establish speciated emission inventories for air quality modeling. A 506 sufficient aging time (~one week) is needed to allow gas-to-particle partitioning of semi-507 volatilized species, gas-phase oxidation, and volatilization to achieve representative source 508 profiles for receptor-oriented source apportionment.

- 5 **Author contribution** 509

- 510 JCC, JGW, JC, L-WAC, and XW jointly designed the study, performed the data analyses, 511 and prepared the manuscript. QW, JT, and SSHH carried out the peat combustion experiments.
- 512 TBC and SDK assembled the database and performed the similarity and difference tests between
- 513 the fresh and aged profiles.
- 514 **Competing interests** 6
- 515 The authors declare that there are no conflicts of interest.





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 0.058 ± 0.053

 0.0688 ± 0.038 0.0230 ± 0.014

 0.047 ± 0.020

 0.017 ± 0.0011 0.020 ± 0.016

 0.032 ± 0.0074

 0.022 ± 0.01 0.11 ± 0.087

 0.024 ± 0.0013

 0.018 ± 0.0015

Water-Soluble Sodium (Na⁺)

Water-Soluble Potassium

Chloride (Cl⁻)

Ę,

Ammonium (NH4⁺)

OCI (140°C) OC2 (280°C)

OC3 (480°C)

OC4 (580°C)

Nitrate (NO₃⁻)

Nitrite (NO2⁻) Sulfate (SO4["])

 0.034 ± 0.036

 6.0095 ± 0.93

 0.18 ± 0.080

na^d

na^d

 18.21 ± 3.97

 0.18 ± 0.052

 0.24 ± 0.085 4.56 ± 1.36

 0.21 ± 0.059

 0.32 ± 0.15 3.21 ± 0.78

 7.84 ± 0.31

PEAT033, PEAT034, PEAT035

na^d

na^d

 7.090 ± 5.59

 22.81 ± 5.88

 0.27 ± 0.075

 0.27 ± 0.074 8.81 ± 4.047

PEAT023, PEAT025, PEAT026

 0.39 ± 0.15

PEAT027, PEAT028, PEAT029

Aged 7

Fresh 7

Aged 2^b

Fresh 2

Aged 7

Fresh 7

Aged 2

Fresh 2

2 days

PEAT030, PEAT031, PEAT032

Peat IDs in the average^c

Aging Time

Nitric Acid (HNO₃)

Ammonia (NH₃)

7 days

2 days

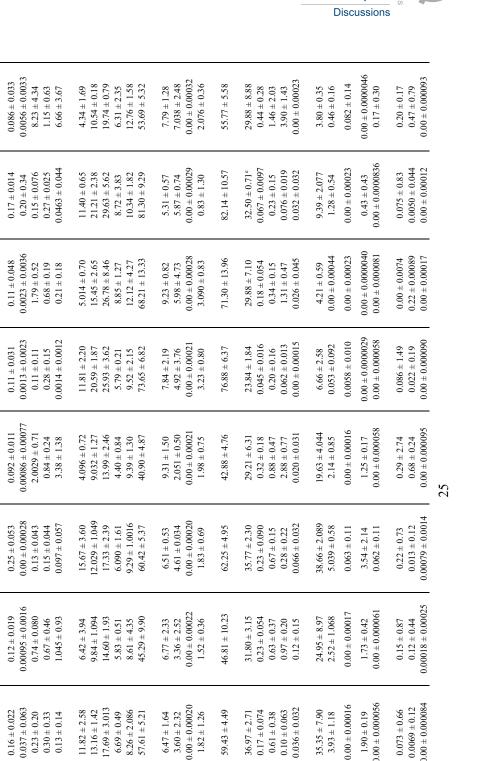
Average \pm Standard Deviation of Percent PM_{2.5} Mass^a

Boreal

7 days

Pskov, Siberia





Water-Soluble OC (WSOC)

Fotal Carbon (TC)

Elemental Carbon (EC)

EC2 (740°C) EC3 (840°C)

EC1 (580°C)

Pyrolized Carbon (OP) Organic Carbon (OC) Levoglucosan (C6H10O5)

Mannosan (C₆H₁₀O₅)

Galactose/Maltitol

(C₆H₁₂O₆/C₁₂H₂₄O₁₁) Glycerol (C₃H₈O₃)

Mannitol (C₆H₁₄O₆)

Aluminum (Al) Silicon (Si) Phosphorous (P)

Oxalate (C₂H₂O₄") Propionate (C₃H₅O₂')

Acetate (C₂H₄O₂)

Formate (CH₂O₂⁻)

Atmospheric

Techniques

Measurement

Oper

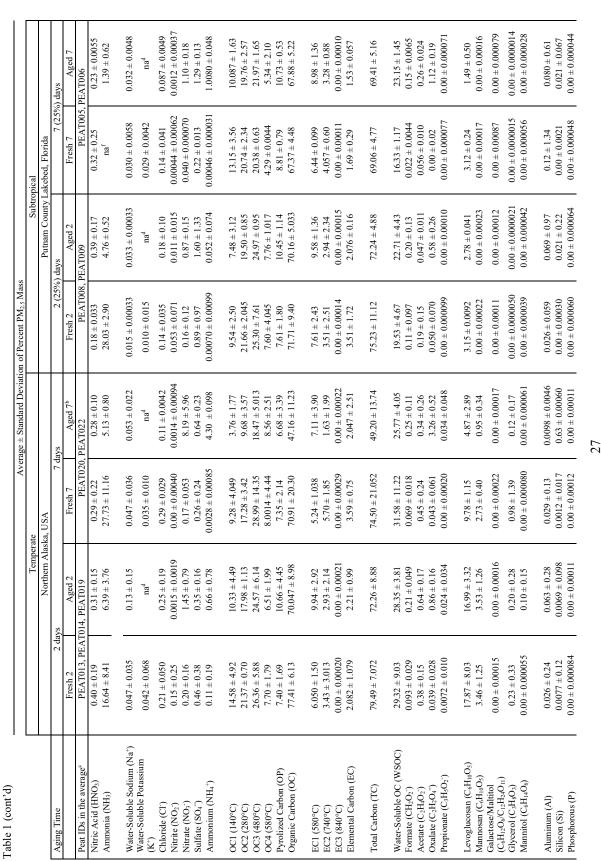
Table 1. Fresh and aged average profiles (in % of PM2.5 mass) for six types of peats Odintsovo, Russia





Table 1 (cont'd) Sulfur (S) Chlorine (Cl)	0.024 ± 0.0088 0.12 ± 0.027	$\begin{array}{c} 0.081 \pm 0.046 \\ 0.035 \pm 0.019 \end{array}$	$\begin{array}{c} 0.040 \pm 0.056 \\ 0.18 \pm 0.030 \end{array}$	$\begin{array}{c} 0.26 \pm 0.095 \\ 0.032 \pm 0.0025 \end{array}$	$\begin{array}{c} 0.081 \pm 0.030 \\ 0.11 \pm 0.015 \end{array}$	$\begin{array}{c} 0.090 \pm 0.000098 \\ 0.057 \pm 0.000068 \end{array}$	$\begin{array}{c} 0.028 \pm 0.034 \\ 0.081 \pm 0.018 \end{array}$	$\begin{array}{c} 0.31 \pm 0.0057 \\ 0.027 \pm 0.0064 \end{array}$
Potassium (K) Calcium (Ca) Scandium (Sc) Titanium (Ti) Vanadium (V)	$\begin{array}{c} 0.030\pm0.011\\ 0.018\pm0.016\\ 0.064\pm0.11\\ 0.0046\pm0.0056\\ 0.0046\pm0.00013 \end{array}$	$\begin{array}{c} 0.48 \pm 0.44 \\ 0.040 \pm 0.056 \\ 0.00 \pm 0.0021 \\ 0.00 \pm 0.00076 \\ 0.00 \pm 0.00014 \end{array}$	0.041 ± 0.018 0.031 ± 0.025 0.002 ± 0.0021 0.0055 ± 0.0049 0.002 ± 0.00014	$\begin{array}{c} 0.13 \pm 0.035 \\ 0.0034 \pm 0.0048 \\ 0.00 \pm 0.0023 \\ 0.0013 \pm 0.0018 \\ 0.00 \pm 0.00015 \end{array}$	$\begin{array}{c} 0.15 \pm 0.19 \\ 0.00 \pm 0.00 \\ 0.079 \pm 0.14 \\ 0.0079 \pm 0.014 \\ 0.00070 \pm 0.0012 \end{array}$	$\begin{array}{c} 0.096\pm0.00025\\ 0.00\pm0.00022\\ 0.00\pm0.0041\\ 0.00\pm0.00015\\ 0.00\pm0.000027\end{array}$	$\begin{array}{c} 0.11\pm0.14\\ 0.00\pm0.00065\\ 0.031\pm0.053\\ 0.00\pm0.00010\\ 0.00\pm0.00019\end{array}$	$\begin{array}{c} 0.30\pm 0.017\\ 0.028\pm 0.039\\ 0.00\pm 0.0022\\ 0.00\pm 0.00078\\ 0.00\pm 0.00015\end{array}$
Chromium (Cr) Manganese (Mn) Iron (Fe) Cobalt (Co) Nickel (Ni)	$\begin{array}{c} 0.0012 \pm 0.0020\\ 0.0014 \pm 0.0022\\ 0.038 \pm 0.021\\ 0.00032 \pm 0.00056\\ 0.00 \pm 0.000022\end{array}$	$\begin{array}{l} 0.00039\pm0.00056\\ 0.00053\pm0.00074\\ 0.091\pm0.098\\ 0.00\pm0.000094\\ 0.0026\pm0.0037\end{array}$	$\begin{array}{c} 0.00\pm0.00046\\ 0.0037\pm0.0033\\ 0.062\pm0.043\\ 0.00037\pm\\ 0.000057\pm\\ 0.000064\\ 0.000050\end{array}$	$\begin{array}{c} 0.00084\pm 0.0012\\ 0.00\pm 0.00018\\ 0.26\pm 0.32\\ 0.00049\pm 0.00069\\ 0.00\pm 0.00025\end{array}$	$\begin{array}{l} 0.00079\pm0.0014\\ 0.0018\pm0.0022\\ 0.039\pm0.035\\ 0.00018\pm0.00031\\ 0.000\pm0.00024 \end{array}$	$\begin{array}{l} 0.00\pm0.00091\\ 0.020\pm0.00032\\ 0.029\pm0.00056\\ 0.00\pm0.00018\\ 0.00\pm0.000018\\ \end{array}$	$\begin{array}{c} 0.0010\pm 0.00095\\ 0.0031\pm 0.0031\\ 0.013\pm 0.017\\ 0.00\pm 0.00013\\ 0.0014\pm 0.0017\\ \end{array}$	$\begin{array}{c} 0.00\pm 0.00049\\ 0.0051\pm 0.0072\\ 0.015\pm 0.021\\ 0.00\pm 0.00018\\ 0.0041\pm 0.00039 \end{array}$
Copper (Cu) Zinc (Zn) Arsenic (As) Selenium (Se) Bromine (Br)	$\begin{array}{c} 0.0055\pm 0.0029\\ 0.0017\pm 0.0015\\ 0.00086\pm 0.0015\\ 0.00021\pm 0.00036\\ 0.00031\end{array}$	$\begin{array}{c} 0.15\pm0.11\\ 0.054\pm0.066\\ 0.00\pm0.00038\\ 0.0026\pm0.0037\\ 0.0030\pm0.0031\end{array}$	$\begin{array}{c} 0.0052\pm0.0038\\ 0.0047\pm0.0041\\ 0.00\pm0.00037\\ 0.00067\pm0.00076\\ 0.00096\pm0.0014\\ \end{array}$	$\begin{array}{c} 0.046\pm0.054\\ 0.053\pm0.070\\ 0.00\pm0.00040\\ 0.0029\pm0.00041\\ 0.0021\pm0.0019\\ \end{array}$	$\begin{array}{c} 0.0072 \pm 0.0041 \\ 0.0055 \pm 0.0030 \\ 0.00076 \pm 0.0013 \\ 0.0018 \pm 0.0022 \\ 0.0012 \pm 0.0043 \end{array}$	$\begin{array}{c} 0.014\pm 0.00028\\ 0.0034\pm 0.00016\\ 0.0050\pm 0.000073\\ 0.0026\pm 0.00013\\ 0.0032\pm 0.000036\end{array}$	$\begin{array}{c} 0.047\pm0.052\\ 0.0058\pm0.0056\\ 0.000069\pm0.00012\\ 0.00003\pm0.00031\\ 0.00032\pm0.0006\end{array}$	$\begin{array}{c} 0.11\pm 0.067\\ 0.0019\pm 0.00081\\ 0.00013\pm 0.00019\\ 0.00029\pm 0.00041\\ 0.0066\pm 0.0014 \end{array}$
Rubidium (Sr) Strontium (Sr) Yttrium (Y) Zirconium (Zr) Niobium (Nb)	$\begin{array}{c} 0.00052\pm 0.00090\\ 0.0033\pm 0.0032\\ 0.00079\pm 0.0013\\ 0.0040\pm 0.0024\\ 0.00072\pm 0.0012 \end{array}$	$\begin{array}{c} 0.0029\pm 0.00079\\ 0.0017\pm 0.0018\\ 0.000066\pm\\ 0.000093\\ 0.00034\pm 0.0014\\ 0.0023\pm 0.0013\end{array}$	$\begin{array}{l} 0.0020\pm0.0019\\ 0.0032\pm0.0027\\ 0.0031\pm0.0035\\ 0.0013\pm0.0018\\ 0.00038\pm0.00038\\ \end{array}$	$\begin{array}{l} 0.00049\pm0.00069\\ 0.0033\pm0.0013\\ 0.00077\pm0.0011\\ 0.0017\pm0.0024\\ 0.00063\pm0.00089 \end{array}$	$\begin{array}{l} 0.00031\pm0.00054\\ 0.0027\pm0.0028\\ 0.0014\pm0.0012\\ 0.0051\pm0.0019\\ 0.00040\pm0.00069\end{array}$	$\begin{array}{l} 0.00\pm0.000045\\ 0.0039\pm0.000045\\ 0.0015\pm0.000045\\ 0.001\pm0.000017\\ 0.00\pm0.00017\\ \end{array}$	$\begin{array}{l} 0.00066\pm0.00068\\ 0.0072\pm0.0042\\ 0.0045\pm0.0045\\ 0.0060\pm0.0088\\ 0.00059\pm0.00067 \end{array}$	$\begin{array}{c} 0.0024\pm 0.0034\\ 0.0047\pm 0.0066\\ 0.0053\pm 0.0049\\ 0.0033\pm 0.0021\\ 0.00623 \end{array}$
Molybdenum (Mo) Silver (Ag) Cadmium (Cd) Indium (In) Tin (Sn)	$\begin{array}{c} 0.0020\pm 0.0035\\ 0.0010\pm 0.0015\\ 0.0034\pm 0.0059\\ 0.000\pm 0.00010\\ 0.0022\pm 0.00048 \end{array}$	$\begin{array}{c} 0.00\pm0.00000\\ 0.00\pm0.00011\\ 0.003\pm0.0053\\ 0.00\pm0.0011\\ 0.0095\pm0.013\end{array}$	$\begin{array}{c} 0.0015\pm0.0011\\ 0.00\pm0.00011\\ 0.0023\pm0.0039\\ 0.0059\pm0.0011\\ 0.0013\pm0.0022\end{array}$	$\begin{array}{c} 0.0030\pm0.0010\\ 0.00\pm0.00012\\ 0.0023\pm0.0033\\ 0.0060\pm0.0016\\ 0.0037\pm0.0053\\ \end{array}$	$\begin{array}{c} 0.0029\pm0.0051\\ 0.00\pm0.0011\\ 0.00\pm0.0016\\ 0.0065\pm0.0011\\ 0.0098\pm0.010\end{array}$	$\begin{array}{c} 0.00 \pm 0.00017\\ 0.00 \pm 0.00022\\ 0.00 \pm 0.00030\\ 0.018 \pm 0.00021\\ 0.0075 \pm 0.00038\end{array}$	$\begin{array}{c} 0.0013\pm0.0022\\ 0.0083\pm0.0074\\ 0.0024\pm0.0029\\ 0.0027\pm0.0047\\ 0.0027\pm0.014 \end{array}$	$\begin{array}{c} 0.0026\pm0.0037\\ 0.00\pm0.00012\\ 0.00\pm0.00016\\ 0.00\pm0.00011\\ 0.008\pm0.013\\ 0.0089\pm0.013 \end{array}$
Antimony (Sb) Cesium (Cs) Barium (Ba) Lanthanum (La) Wolfram (W)	$\begin{array}{c} 0.00\pm0.00028\\ 0.025\pm0.040\\ 0.014\pm0.024\\ 0.048\pm0.043\\ 0.0023\pm0.0014\end{array}$	$\begin{array}{c} 0.0086\pm0.012\\ 0.0085\pm0.012\\ 0.00\pm0.0071\\ 0.00\pm0.0012\\ 0.0073\pm0.010\end{array}$	$\begin{array}{c} 0.00\pm0.0029\\ 0.023\pm0.033\\ 0.011\pm0.020\\ 0.049\pm0.043\\ 0.0077\pm0.013 \end{array}$	$\begin{array}{c} 0.00\pm0.00032\\ 0.014\pm0.020\\ 0.00\pm0.0068\\ 0.059\pm0.083\\ 0.011\pm0.0016\end{array}$	$\begin{array}{c} 0.00\pm0.0030\\ 0.0057\pm0.0099\\ 0.023\pm0.020\\ 0.017\pm0.030\\ 0.0079\pm0.0014\\ \end{array}$	$\begin{array}{c} 0.000053\pm 0.00058\\ 0.00\pm 0.0016\\ 0.00\pm 0.0012\\ 0.00\pm 0.0024\\ 0.00\pm 0.00047\end{array}$	$\begin{array}{c} 0.00\pm0.00041\\ 0.0046\pm0.0079\\ 0.00\pm0.0086\\ 0.094\pm0.085\\ 0.0047\pm0.0082\\ \end{array}$	$\begin{array}{c} 0.00\pm 0.0031\\ 0.00\pm 0.0086\\ 0.00\pm 0.0067\\ 0.02\pm 0.028\\ 0.0048\pm 0.0054\\ \end{array}$
Gold (Au) Mercury (Hg) Lead (Pb) Uranium (U)	$\begin{array}{c} 0.0029 \pm 0.0027 \\ 0.0015 \pm 0.0014 \\ 0.0026 \pm 0.0024 \\ 0.0018 \pm 0.0031 \end{array}$	$\begin{array}{c} 0.00\pm 0.00071\\ 0.00\pm 0.00038\\ 0.0018\pm 0.0025\\ 0.0017\pm 0.0024\end{array}$	$\begin{array}{c} 0.00080\pm 0.0014\\ 0.00081\pm 0.0014\\ 0.0024\pm 0.0014\\ 0.0026\pm 0.0017\\ \end{array}$	$\begin{array}{c} 0.0024 \pm 0.0033 \\ 0.00 \pm 0.00040 \\ 0.0053 \pm 0.0074 \\ 0.0024 \pm 0.0035 \end{array}$	$\begin{array}{c} 0.00\pm 0.00071\\ 0.0013\pm 0.0023\\ 0.00\pm 0.00071\\ 0.0028\pm 0.0027\end{array}$	$\begin{array}{c} 0.012 \pm 0.00014 \\ 0.00 \pm 0.00073 \\ 0.00 \pm 0.00014 \\ 0.00 \pm 0.00025 \end{array}$	$\begin{array}{c} 0.0038\pm0.0065\\ 0.000065\pm0.00011\\ 0.0050\pm0.00088\\ 0.0025\pm0.0033\end{array}$	$\begin{array}{c} 0.0018\pm0.0025\\ 0.00\pm0.00039\\ 0.0027\pm0.0032\\ 0.0046\pm0.0066 \end{array}$





Atmospheric

Techniques

Discussions

Measurement

EGU





Table 1 ($aantable$)								
Sulfur (S) Chlorine (Cl)	0.031 ± 0.054 0.12 ± 0.068	$\begin{array}{c} 0.062 \pm 0.087 \\ 0.087 \pm 0.030 \end{array}$	0.0099 ± 0.014 0.14 ± 0.049	$\begin{array}{c} 0.34 \pm 0.00013 \\ 0.019 \pm 0.000040 \end{array}$	0.19 ± 0.056 0.12 ± 0.0064	0.37 ± 0.24 0.067 ± 0.024	0.17 ± 0.037 0.14 ± 0.022	$\begin{array}{c} 0.74 \pm 0.047 \\ 0.056 \pm 0.00047 \end{array}$
	0.017 - 0.017			0.17 - 0.00000	0.0000 - 0.010		0.0015 - 0.00011	010.010
Potassium (K)	0.046 ± 0.016	$CI.0 \pm 0.10$	0.02 ± 0.040	$0.4 / \pm 0.00022$	710.0 ± 7600.0	CCU.U ± / CU.U	0.0046 ± 0.00044	0.12 ± 0.10
Calcium (Ca)	0.032 ± 0.032	0.032 ± 0.045	0.035 ± 0.049	0.00 ± 0.00057	0.0040 ± 0.0056	0.00 ± 0.0034	0.00 ± 0.00025	0.00 ± 0.00023
Scandium (Sc)	0.00 ± 0.0020	0.00 ± 0.0025	0.00 ± 0.0029	0.00 ± 0.0026	0.00 ± 0.0014	0.00 ± 0.0015	0.022 ± 0.031	0.00 ± 0.0010
Titanium (Ti)	0.00 ± 0.00071	0.00 ± 0.000001	0.0055 ± 0.0078	0.051 ± 0.000093	0.0036 ± 0.0050	0.00 ± 0.00054	0.0086 ± 0.012	0.00 ± 0.00037
Vanadium (V)	0.00 ± 0.00013	0.00 ± 0.00017	0.00 ± 0.00019	0.00 ± 0.000017	0.00 ± 0.00004	0.00 ± 0.00010	0.00 ± 0.0000075	0.00 ± 0.0000069
Chromium (Cr)	0.00051 ± 0.00089	0.00028 ± 0.00040	0.00 ± 0.00065	0.00 ± 0.00057	0.00 ± 0.000032	0.00 ± 0.00034	0.00034 ± 0.00048	0.00 ± 0.000023
Manganese (Mn)	0.0015 ± 0.0014	0.00069 ± 0.00098	0.0016 ± 0.0023	0.0011 ± 0.00020	0.0013 ± 0.0012	0.00033 ± 0.00047	0.00057 ± 0.00080	0.0016 ± 0.0018
Iron (Fe)	0.036 ± 0.014	0.10 ± 0.095	0.049 ± 0.048	0.029 ± 0.00035	0.00 ± 0.00019	0.047 ± 0.040	0.024 ± 0.012	0.065 ± 0.0091
Cobalt (Co)	0.00 ± 0.000088	0.00 ± 0.00011	0.00 ± 0.00013	0.00013 ± 0.000011	0.00 ± 0.000063	0.00021 ± 0.00030	0.00020 ± 0.00028	0.00 ± 0.0000046
Nickel (Ni)	0.00028 ± 0.00049	0.00 ± 0.00028	0.00075 ± 0.0011	0.00 ± 0.00028	0.00045 ± 0.00064	0.00 ± 0.00017	0.00069 ± 0.00097	0.00043 ± 0.00026
Conner (Cu)	0.028 ± 0.047	0.027 ± 0.034	0.0098 ± 0.0028	0.15 ± 0.00018	0.00 ± 0.0008	0.0035 ± 0.0049	0.0019 ± 0.000053	0.069 ± 0.090
Zine (Zn)	0.076 ± 0.036	0.027 ± 0.031	0 000 0 + 9 000 0	0.011 ± 0.000007	0.0013 ± 0.0015	0.003 ± 0.003	0.00041 ± 0.00008	0.0046 ± 0.00037
	0.000 - 020.0	1000 - 1700		1/00000 = 11000	CT00:0 + CT00:0	10000 - 01000	$+ 000000 \pm 100000$	
Arsenic (As)	0.0006 ± 0.00078	0.00 ± 0.00045	0.00 ± 0.00052	0.00067 ± 0.000045	0.00 ± 0.000025	0.00 ± 0.000027	0.000087	0.00034 ± 0.00048
Selenium (Se)	0.00016 ± 0.00028	0.0064 ± 0.0017	0.0022 ± 0.0032	0.00 ± 0.00080	0.0017 ± 0.00092	0.00 ± 0.00047	0.00034 ± 0.00048	0.0034 ± 0.0017
Bromine (Br)	0.0017 ± 0.0018	0.0031 ± 0.0044	0.0079 ± 0.00064	0.0020 ± 0.000023	0.020 ± 0.0008	0.0077 ± 0.010	0.024 ± 0.0043	0.019 ± 0.0012
r.								
Rubidium (Rb)	0.00 ± 0.000022	0.0035 ± 0.0048	0.0057 ± 0.0059	0.0026 ± 0.000028	0.00011 ± 0.00016	0.00095 ± 0.0013	0.00 ± 0.00013	0.00066 ± 0.00047
Strontium (Sr)	0.0017 ± 0.00036	0.0076 ± 0.0084	0.0068 ± 0.0014	0.0028 ± 0.000028	0.0023 ± 0.00057	0.0038 ± 0.0013	0.0018 ± 0.00075	0.0046 ± 0.0025
Yttrium (Y)	0.0013 ± 0.0014	0.0037 ± 0.0013	0.0057 ± 0.0041	0.0054 ± 0.000028	0.0014 ± 0.00029	0.0012 ± 0.0018	0.00085 ± 0.000067	0.0022 ± 0.0032
Zirconium (Zr)	0.0027 ± 0.0028	0.0047 ± 0.0014	0.0025 ± 0.0027	0.011 ± 0.00011	0.0016 ± 0.0023	0.0003 ± 0.00089	0.00074 ± 0.0010	0.0013 ± 0.00079
Niobium (Nb)	0.00 ± 0.00040	0.00092 ± 0.00090	0.00027 ± 0.00039	0.00 ± 0.00051	0.0016 ± 0.0023	0.00082 ± 0.0012	0.00042 ± 0.00060	0.00 ± 0.00021
Molvhdenum (Mo)	0.0012 ± 0.0019	0.0044 ± 0.0062	0.0020 ± 0.00084	0.00 ± 0.00011	0.00 ± 0.00060	0.00063 ± 0.00089	0.0025 ± 0.00092	0.00 ± 0.00044
Silver (AG)	0.00 ± 0.00011	0.00 ± 0.00014	0.00 ± 0.0016	0.00 ± 0.0014	0.0010 ± 0.0014	0.00 ± 0.00081	0.00 ± 0.00060	0.00 ± 0.00055
Cadmium (Cd)	0.00 ± 0.00015	0.00 ± 0.00019	0.00 ± 0.00022	0.00 ± 0.0019	0.0034 ± 0.0049	0.00 ± 0.00011	0.0029 ± 0.00093	0.0020 ± 0.0029
Indium (In)	0.00082 ± 0.0013	0.0011 ± 0.0016	0.00069 ± 0.00097	0.00 ± 0.00013	0.00068 ± 0.00096	0.0025 ± 0.0036	0.0021 ± 0.0030	0.0018 ± 0.0026
Tin (Sn)	0.0045 ± 0.0078	0.014 ± 0.020	0.0067 ± 0.0025	0.00 ± 0.00024	0.0037 ± 0.00047	0.0034 ± 0.0048	0.0028 ± 0.0025	0.0074 ± 0.00049
Antimony (Sb)	0.0065 ± 0.011	0.015 ± 0.021	0.00 ± 0.00041	0.00 ± 0.0036	0.00 ± 0.00020	0.0072 ± 0.010	0.0020 ± 0.0029	0.00 ± 0.00015
Cesium (Cs)	0.0097 ± 0.0095	0.022 ± 0.031	0.010 ± 0.014	0.058 ± 0.0010	0.00 ± 0.00056	0.00 ± 0.00060	0.00 ± 0.00044	0.00 ± 0.00041
Barium (Ba)	0.00 ± 0.00059	0.00 ± 0.00077	0.00 ± 0.00086	0.00 ± 0.0089	0.00 ± 0.00042	0.00 ± 0.00046	0.00 ± 0.0034	0.00 ± 0.00031
Lanthanum (La)	0.015 ± 0.026	0.065 ± 0.025	0.055 ± 0.0026	0.00 ± 0.0015	0.042 ± 0.044	0.0053 ± 0.0075	0.019 ± 0.028	0.036 ± 0.021
Wolfram (W)	0.0034 ± 0.0059	0.0082 ± 0.0061	0.00 ± 0.00033	0.00 ± 0.00029	0.0037 ± 0.0018	0.0034 ± 0.0049	0.0019 ± 0.0028	0.00 ± 0.00012
	0.00 ± 0.00066	0.0032 ± 0.0045		0.00 ± 0.00085	88000 0 + 69000 0	0.00 ± 0.00051	0.00037 ± 0.00031	0.0012 ± 0.0017
	0.0034 ± 0.00050	0.000 ± 0.000	0.000 ± 0.00050	0.00 ± 0.00005	0.0000 ± 0.0000	10000 ± 0.000	100000 ± 770000	0.0012 ± 0.0017
Mercury (ng)	700000 ± 400000	0.0014 ± 0.0020	20000.0 ± 00.0	0.00 ± 0.00043	0.00020 ± 0.00020	0.0014 ± 0.0020	0.000 ± 0.00020	0.00024 ± 0.00030
Leau (FD)	0.000 ± 0.0000 0 0.001 0.001 0.001 0.001 0.001 0.001 0.001 0.001 0.001 0.001 0.001 0.001 0.001 0.001 0.001 0.0001 0.001	$C100.0 \pm 0100.0$	0.00 ± 0.0000	0.0035 ± 0.00015	0.0024 ± 0.0044	0.001 ± 0.00000	$0.000/0 \pm 0.0011$	100.0 ± 0.000
		1700.0 ± 0200.0	CT0007 T1000	C1000.0 ± CC00.0	++00.0 ± +000.0	$7 \leq 0 \leq $	1 CON'N T 07000	7000000 ± 0.00

> Tropical Borneo, Malaysia

Average \pm Standard Deviation of Percent PM2.5 Mass

Everglades National Park, Florida

Subtropical



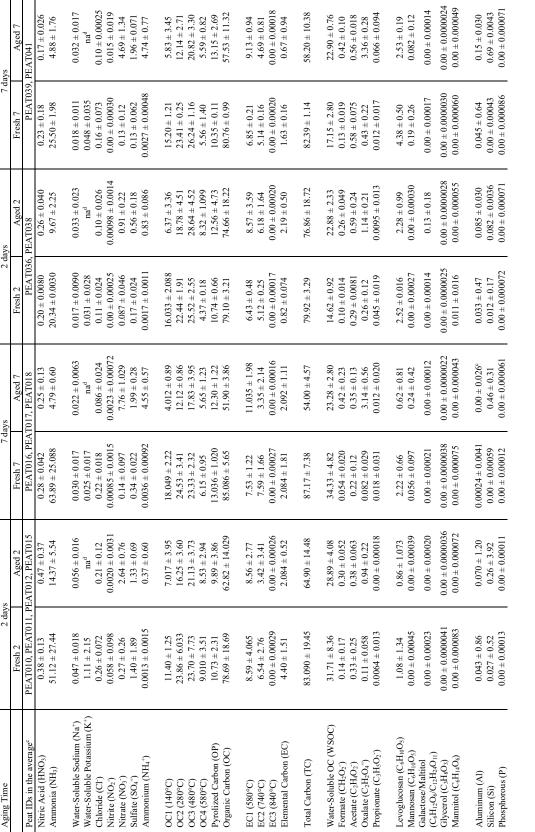


Table 1 (cont'd)

706

29

Atmospheric

Techniques

Discussions

Measurement





$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Table 1 (cont'd)								
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Sulfur (S)	0.39 ± 0.23	0.59 ± 0.27	0.42 ± 0.066	1.12 ± 0.094	0.11 ± 0.12	0.39 ± 0.00013	0.029 ± 0.0022	0.83 ± 0.00026
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Chlorine (Cl)	0.21 ± 0.088	670.0 ± C00.0	0.24 ± 0.024	0.058 ± 0.011	$0.0/4 \pm 0.0012$	$0.06/ \pm 0.00035$	0.085 ± 0.0038	$0.04 / \pm 0.000050$
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Potassium (K)	0.034 ± 0.015	0.51 ± 0.37	0.018 ± 0.014	0.22 ± 0.052	0.051 ± 0.049	0.084 ± 0.00010	0.028 ± 0.017	0.017 ± 0.00010
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Calcium (Ca)	0.00 ± 0.00067	0.0081 ± 0.016	0.00 ± 0.0061	0.010 ± 0.014	0.0058 ± 0.0082	0.00 ± 0.00037	0.00 ± 0.0046	0.023 ± 0.00038
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Scandium (Sc)	0.00 ± 0.0030	0.00 ± 0.0026	0.00 ± 0.0027	0.00 ± 0.0014	0.00 ± 0.0017	0.00 ± 0.0017	0.00 ± 0.0020	0.00 ± 0.0017
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Titanium (Ti)	0.0061 ± 0.0079	0.017 ± 0.035	0.00 ± 0.00008	0.00 ± 0.000051	0.0073 ± 0.010	0.00 ± 0.00059	0.0066 ± 0.0094	0.00 ± 0.00059
$1000 \pm 0.00053 \pm 0.0005$ 0.00053 ± 0.0001 0.00054 ± 0.0001 0.00044 ± 0.0011 0.00044 ± 0.0011 0.00044 ± 0.0001 0.00044 ± 0.0011 <th< td=""><td>Vanadium (V)</td><td>0.0010 ± 0.0020</td><td>0.00 ± 0.00017</td><td>0.00 ± 0.00018</td><td>0.0065 ± 0.0092</td><td>0.00 ± 0.00011</td><td>0.00 ± 0.00011</td><td>0.00 ± 0.00014</td><td>0.00 ± 0.00011</td></th<>	Vanadium (V)	0.0010 ± 0.0020	0.00 ± 0.00017	0.00 ± 0.00018	0.0065 ± 0.0092	0.00 ± 0.00011	0.00 ± 0.00011	0.00 ± 0.00014	0.00 ± 0.00011
$ \begin{array}{ c c c c c c c c c c c c c c c c c c c$	Chromium (Cr)	0.00 ± 0.00066	0.00056 ± 0.0011	0.00 ± 0.00061	0.00016 ± 0.00023	0.00 ± 0.00038	0.00 ± 0.00037	0.0026 ± 0.0037	0.00 ± 0.000037
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Manganese (Mn)	0.0032 ± 0.0064	0.0051 ± 0.0050	0.0017 ± 0.0015	0.0034 ± 0.0043	0.0055 ± 0.0026	0.0075 ± 0.00013	0.0088 ± 0.00010	0.0046 ± 0.00013
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Iron (Fe)	0.023 ± 0.021	0.065 ± 0.034	0.020 ± 0.016	0.091 ± 0.096	0.074 ± 0.0078	0.074 ± 0.00023	0.045 ± 0.020	0.043 ± 0.00023
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Cohalt (Co)	0.000055 ± 0.00011	0.000045 ± 0.000090	0.00024 ± 0.00041	0.00 ± 0.000064	0.00 ± 0.0000075	0.00061 ± 0.000074	0.00 ± 0.000000	0.000087 ± 0.000074
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Nickel (Ni)	0.00026 ± 0.00042	0.00 ± 0.00029	0.00 ± 0.00031	0.00038 ± 0.00054	0.00064 ± 0.00091	0.00 ± 0.00019	0.0034 ± 0.0014	0.00 ± 0.00019
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$									
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Copper (Cu)	0.010 ± 0.0080	0.21 ± 0.23	0.0033 ± 0.0036	0.021 ± 0.0024	0.0054 ± 0.0042	0.0075 ± 0.00012	0.0091 ± 0.0013	0.0017 ± 0.00012
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	ZINC (ZN) Arsenic (As)	1100.0 ± 6200.0	0.0013 ± 0.0020	0.00 ± 0.0019	0.00 ± 0.0003	0.00 ± 0.0030	0.00 ± 0.000030	0.00 ± 0.0036	0.003 ± 0.00003
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Selenium (Se)	0.0011 ± 0.0014	0.0023 ± 0.0018	0.0037 ± 0.0025	0.00016 ± 0.00023	0.0019 ± 0.0010	0.00 ± 0.000052	0.00086 ± 0.0012	0.00 ± 0.00052
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Bromine (Br)	0.030 ± 0.015	0.0090 ± 0.0049	0.022 ± 0.0072	0.0088 ± 0.0036	0.011 ± 0.0015	0.012 ± 0.000015	0.012 ± 0.0026	0.0044 ± 0.000015
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$									
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Rubidium (Rb)	0.00038 ± 0.00077	0.0015 ± 0.0014	0.0015 ± 0.0026	0.00 ± 0.00016	0.00039 ± 0.00056	0.00035 ± 0.000019	0.00 ± 0.000023	0.0017 ± 0.00019
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Strontium (Sr)	0.0031 ± 0.0012	0.0044 ± 0.0023 0.0021 ± 0.0034	0.0003 ± 0.0003	0.0055 ± 0.0022 0.00 ± 0.000016	0.0028 ± 0.00026 0.0018 ± 0.0023	0.0021 ± 0.00019 0.0032 ± 0.000019	$0.00/0 \pm 0.0009$	0.0029 ± 0.00019 0.0027 ± 0.00010
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Tirconium (T)	0.0041 ± 0.0038	0.0049 ± 0.0066	0.0040 ± 0.0069	0.0051 ± 0.0039	0.0048 ± 0.00038	0.0016 ± 0.00001	0.00052 ± 0.00074	0.00 ± 0.00071
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Niobium (Nb)	0.0016 ± 0.0022	0.00080 ± 0.0013	0.0019 ± 0.0026	0.00 ± 0.00029	0.00095 ± 0.0014	0.00 ± 0.00034	0.0021 ± 0.0030	0.00026 ± 0.000034
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Malthedamme (Ma)	0.000 ± 0.000	0.0012 ± 0.0017	0.000 ± 0.000	0.00081 ± 0.0011	0.00071 ± 0.0010	0.00 ± 0.00071	0.0014 ± 0.00018	0.0032 ± 0.000071
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Silver (AG)	0.0014 ± 0.0020	0.00 ± 0.0014	0.0012 ± 0.0015	1100.0 ± 10000.0	0.0075 ± 0.0035	0.00 ± 0.0000 1 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	0.0076 ± 0.0037	$1/0000.0 \pm 2000.0$
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Cadmium (Cd)	0.00 ± 0.00022	0.00 ± 0.0019	0.0075 ± 0.013	0.0095 ± 0.0060	0.00044 ± 0.00063	0.00 ± 0.00012	0.00 ± 0.00015	0.00 ± 0.00012
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Indium (In)	0.0069 ± 0.0049	0.0023 ± 0.0046	0.0054 ± 0.0093	0.0012 ± 0.0017	0.0048 ± 0.0067	0.0013 ± 0.000085	0.00087 ± 0.0012	0.00 ± 0.00085
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Tin (Sn)	0.0061 ± 0.0072	0.0058 ± 0.012	0.0061 ± 0.0058	0.0068 ± 0.0096	0.0022 ± 0.0031	0.013 ± 0.00016	0.0038 ± 0.0054	0.012 ± 0.00016
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Antimony (Sh)	0.00028 ± 0.00056	0.00040 ± 0.00052	0.00033 ± 0.00057	0.00050 ± 0.00071	0.00 ± 0.00024	0.0039 ± 0.00023	0.011 ± 0.0097	0.00 ± 0.00023
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Cesium (Cs)	0.000088 ± 0.00018	0.028 ± 0.037	0.037 ± 0.064	0.00 ± 0.00057	0.028 ± 0.031	0.020 ± 0.00066	0.0077 ± 0.011	0.00 ± 0.00066
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Barium (Ba)	0.00 ± 0.00088	0.00 ± 0.0085	0.00 ± 0.0081	0.00 ± 0.0044	0.00 ± 0.00050	0.00 ± 0.00050	0.00 ± 0.0060	0.00 ± 0.00050
$ \begin{array}{ c c c c c c c c c c c c c c c c c c c$	Lanthanum (La)	0.054 ± 0.039	0.033 ± 0.039	0.036 ± 0.039	0.0049 ± 0.0070	0.041 ± 0.058	0.00 ± 0.0007	0.018 ± 0.025	0.080 ± 0.00097
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Wolfram (W)	0.010 ± 0.012	0.0030 ± 0.0051	0.0080 ± 0.014	0.00 ± 0.00016	0.00 ± 0.00019	0.0058 ± 0.00019	0.00 ± 0.00023	0.00 ± 0.00019
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Gold (Au)	0.0012 ± 0.0013	0.00082 ± 0.0016	0.0046 ± 0.0045	0.00033 ± 0.00047	0.00051 ± 0.00072	0.00 ± 0.00056	0.00041 ± 0.00058	0.00 ± 0.00056
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	Mercury (H a)	0.00035 ± 0.00070	0.00091 ± 0.0015	0.00 ± 0.00049	0.00 ± 0.00025	0.00 ± 0.00030	0.00 ± 0.00030	0.00041 ± 0.00058	0.000087 ± 0.000030
$0.0027 \pm 0.0021 \pm 0.0025 \pm 0.0026 = 0.0044 \pm 0.0077 = 0.0017 \pm 0.0023 = 0.0010 = 0.0033 \pm 0.00010 = 0.0057 \pm 0.00076 = 0.0076 \pm 0.0076 = 0.00076$	Lead (Pb)	0.0017 ± 0.0035	0.0012 ± 0.0024	0.0018 ± 0.0031	0.0028 ± 0.0026	0.0031 ± 0.0044	0.00052 ± 0.000056	0.0016 ± 0.0022	0.00 ± 0.00056
	Uranium (U)	0.0027 ± 0.0031	0.0023 ± 0.0026	0.0044 ± 0.0077	0.0017 ± 0.0023	0.00 ± 0.0010	0.0033 ± 0.00010	0.0057 ± 0.00076	0.0062 ± 0.00010

-Analytical uncertainties are used for species below the minimum detection minit, mostly for carbonydrate species and elements with an at ^bOnly one sample was analyzed for elements by x-ray fluorescence with abundance and measurement uncertainty "Peat ID code, detailed operation parameters are reported in Watson et al. (2019)





⁴Data not available; water-soluble K^+ data were contaminated for aged samples due to the use of potassium iodide denuder downstream of the oxidation flow reactor ⁴WSOC measures from Peat sample ID PEAT028 was invalidated due to a crack in the test tube. Therefore, only two measurements are used to calculate the average and standard deviation. ⁴Data not available due to the invalidated citric acid impregnated filter sample

712





715 Table 2. Equivalence measures^a for comparison of PM_{2.5} peat source profiles. Highlighted P-values <0.05 indicate significant 716 differences at the 95% confidence level.

All Fresh (Profile #1) vs. A	ll Aged (Profile #2) by Biome (g	roup co	mparis	on of fresh a	nd aged sam	ples)			
					Percent Di	stribution		Correlation	
Peat region ^b	Peats Included	nl°	n2°	<1 σ	1 - 2 σ	2 - 3 σ	$> 3 \sigma$	Coefficient	P-value ^d
Boreal	Russia + Siberia	12	12	93.60%	5.60%	0.80%	0.00%	0.995	0.00012
Boreal + Temperate	Russia + Siberia + Alaska	17	17	95.20%	4.80%	0.00%	0.00%	0.996	0.00010
Temperate	Alaska	5	5	96.00%	4.00%	0.00%	0.00%	0.997	0.00008
Subtropical	Florida	11	11	92.86%	7.14%	0.00%	0.00%	0.985	0.00007
Subtropical + Temperate	Alaska + Florida	16	16	94.44%	5.56%	0.00%	0.00%	0.992	0.00004
Tropical	Malaysia	4	4	78.57%	18.25%	1.59%	1.59%	0.994	0.00195
Subtropical + Tropical	Florida + Malaysia	15	15	93.65%	6.35%	0.00%	0.00%	0.990	0.00009

Fresh 2 vs. Aged 2 by Bion	e (paired comparison for 2-day	aging)							
					Percent Di	stribution		Correlation	
Peat region ^b	Peats Included	n1°	n2°	<1 σ	1 - 2 σ	2 - 3 σ	$> 3 \sigma$	Coefficient	P-value ^d
Boreal	Russia + Siberia	6	6	94.40%	3.20%	2.40%	0.00%	0.997	0.00088
Boreal + Temperate	Russia + Siberia + Alaska	9	9	95.20%	4.00%	0.80%	0.00%	0.997	0.00237
Temperate	Alaska	3	3	86.40%	11.20%	0.80%	1.60%	0.997	0.02474
Subtropical	Florida	6	6	92.86%	6.35%	0.79%	0.00%	0.992	0.00001
Subtropical + Temperate	Alaska + Florida	9	9	96.83%	2.38%	0.00%	0.79%	0.996	0.00006
Tropical	Malaysia	2	2	80.00%	5.33%	5.33%	9.33%	0.996	0.95960
Subtropical + Tropical	Florida + Malaysia	8	8	96.83%	2.38%	0.79%	0.00%	0.995	0.00007

					Percent Di	stribution		Correlation	
Peat region ^b	Peats Included	n1°	n2°	<1 σ	1 - 2 σ	2 - 3 σ	$> 3 \sigma$	Coefficient	P-value ^d
Boreal	Russia + Siberia	6	6	76.00%	20.80%	1.60%	1.60%	0.992	0.00007
Boreal + Temperate	Russia + Siberia + Alaska	8	8	76.80%	20.00%	0.80%	2.40%	0.993	0.00003
Temperate	Alaska	2	2	64.86%	25.68%	2.70%	6.76%	0.993	0.00000
Subtropical	Florida	5	5	73.02%	23.81%	2.38%	0.79%	0.974	0.00023
Subtropical + Temperate	Alaska + Florida	7	7	75.40%	23.02%	1.59%	0.00%	0.984	0.00004
Tropical	Malaysia	2	2	41.33%	21.33%	24.00%	13.33%	0.989	0.00017
Subtropical + Tropical	Florida + Malaysia	7	7	75.40%	21.43%	2.38%	0.79%	0.983	0.00012

Fresh 2 vs. Fresh 7 by Bion	ne (comparison between differen	nt expei	riments	for unaged fr	esh profiles)			
					Percent Di	stribution		Correlation	
Peat region ^b	Peats Included	n1°	n2°	<1 σ	1 - 2 σ	2 - 3 σ	$> 3 \sigma$	Coefficient	P-value ^d
Boreal	Russia + Siberia	6	6	97.62%	2.38%	0.00%	0.00%	0.999	0.00004
Boreal + Temperate	Russia + Siberia + Alaska	9	8	100.00%	0.00%	0.00%	0.00%	0.999	0.00148
Temperate	Alaska	3	2	91.27%	6.35%	0.79%	1.59%	0.996	0.12876
Subtropical	Florida	6	5	98.41%	1.59%	0.00%	0.00%	0.997	0.52344
Subtropical + Temperate	Alaska + Florida	9	7	100.00%	0.00%	0.00%	0.00%	0.998	0.93350
Tropical	Malaysia	2	2	81.10%	10.24%	3.15%	5.51%	0.999	0.00006
Subtropical + Tropical	Florida + Malaysia	8	7	100.00%	0.00%	0.00%	0.00%	0.999	0.11445

Aged 2 vs. Aged 7 by Biom	e (comparison between differen	it experi	ments	for the 2- and	l 7-day agin	g times)			
					Percent Di	istribution		Correlation	
Peat region ^b	Peats Included	nl°	n2°	<1 σ	1 - 2 σ	2-3σ	> 3 o	Coefficient	P-value ^d
Boreal	Russia + Siberia	6	6	95.20%	3.20%	1.60%	0.00%	0.997	0.00018
Boreal + Temperate	Russia + Siberia + Alaska	9	8	94.40%	3.20%	1.60%	0.80%	0.998	0.00002
Temperate	Alaska	3	2	66.22%	27.03%	5.41%	1.35%	0.996	0.00000
Subtropical	Florida	6	5	93.65%	6.35%	0.00%	0.00%	0.998	0.00194
Subtropical + Temperate	Alaska + Florida	9	7	98.41%	1.59%	0.00%	0.00%	0.998	0.00002
Tropical	Malaysia	2	2	81.33%	13.33%	1.33%	4.00%	0.997	0.00002
Subtropical + Tropical	Florida + Malaysia	8	7	96.03%	3.97%	0.00%	0.00%	0.998	0.00026

For the t-test, a cutoff probability level of 5% is selected; if P < 0.05, there is a 95% probability that the two profiles are different. For correlations, r > 0.8 suggests similar profiles, 0.5 < r < 0.8 indicates a moderate similarity, and r < 0.5 denotes little or no similarity. The R/U ratio indicates the percentage of the >93 reported chemical abundances differ by more than an expected number of uncertainty intervals. The normal probability density function of 68%, 95.5%, and 99.7% for $\pm 1 \sigma$, $\pm 2\sigma$, and $\pm 3\sigma$, respectively, is used to evaluate the R/Uthan an expected number of uncertainty increases. The normal probability density function of 05^{20} , $5_{2,3,20}$, and $9_{2,7,70}$ for ± 10 , ± 20 , and ± 30 , respectively, is used to evaluate the ratios. The two profiles are considered to be similar, within the uncertainties of the chemical abundances when 80^{9} of the R/U ratios are within $\pm 3\sigma$, with r > 0.8 and P > 0.05. Species with R/U ratios > 3σ are further examined as these may be markers that further allow source contributions to be distinguishes by receptor measurements. They may also reflect the sampling and analysis artifacts that are not representative of the larger population of source profiles. ^bUnless otherwise noted, Boreal represents Russia and Siberia regions, Temperate represents northern Alaska region, Subtropical represents north and south Florida regions, and

Tropical represents Island of Borneo, Malaysia region.

^cn1 and n2 denote number of samples in comparison dStudent t-test P-values



Table 3. Organic carbon diagnostic ratios	bon diagnos	tic ratios for	for different peat samples.	samples.				
Peat Type	Atmospheric Aging time	$OC/TC\pm\sigma^a$	$OM^b/OC \pm \sigma^a$	WSOC%/OC $\pm \sigma^a$	$(Levoglucosan/2.25)^d/OC \pm \sigma^a$	$(Oxalate/3.75)^{e}/OC \pm \sigma^{a}$	$(Levoglucosan/2.25)^d/WSOC \pm \sigma^a$	$(Oxalate/3.75)^{e}/WSOC \pm \sigma^{a}$
	Fresh 2	0.97 ± 0.11	1.7 ± 0.15	0.64 ± 0.075	0.27 ± 0.066	0.00047 ± 0.00029	0.42 ± 0.10	0.00073 ± 0.00045
e e e	Aged 2	0.97 ± 0.30	2.1 ± 0.46	0.70 ± 0.17	0.24 ± 0.10	0.0057 ± 0.0017	0.35 ± 0.13	0.0082 ± 0.0019
Odifitsovo, Kussia	Fresh 7	0.97 ± 0.12	1.6 ± 0.14	0.59 ± 0.065	0.28 ± 0.030	0.0012 ± 0.001	0.48 ± 0.040	0.0021 ± 0.0017
	Aged 7	0.95 ± 0.16	2.2 ± 0.26	0.71 ± 0.18	0.21 ± 0.051	0.019 ± 0.0055	0.30 ± 0.089	0.026 ± 0.0090
	Fresh 2	0.96 ± 0.12	1.3 ± 0.12	0.32 ± 0.039	0.04 ± 0.016	0.00023 ± 0.000050	0.12 ± 0.049	0.00069 ± 0.00015
Dolrow Cilconio	Aged 2	0.96 ± 0.26	1.4 ± 0.27	0.44 ± 0.13	0.027 ± 0.0066	0.0051 ± 0.0021	0.063 ± 0.017	0.012 ± 0.0050
rskov, Slucila	Fresh 7	0.99 ± 0.17	1.2 ± 0.14	0.40 ± 0.046	0.051 ± 0.013	0.00025 ± 0.000067	0.13 ± 0.055	0.00063 ± 0.00015
	Aged 7	0.96 ± 0.14	1.5 ± 0.18	0.56 ± 0.17	0.031 ± 0.0043	0.019 ± 0.0073	0.057 ± 0.018	0.035 ± 0.016
	Fresh 2	0.97 ± 0.12	1.3 ± 0.10	0.38 ± 0.12	0.10 ± 0.047	0.00013 ± 0.00010	0.27 ± 0.15	0.00035 ± 0.00028
Monthous Alache IICA	Aged 2	0.97 ± 0.17	1.4 ± 0.18	0.40 ± 0.075	0.11 ± 0.025	0.0033 ± 0.00073	0.27 ± 0.063	0.0080 ± 0.0018
NOTTICITI Alaska, USA	Fresh 7	0.95 ± 0.38	1.4 ± 0.39	0.45 ± 0.20	0.061 ± 0.019	0.00016 ± 0.00023	0.14 ± 0.052	0.00037 ± 0.00053
	Aged 7	0.96 ± 0.35	1.8 ± 0.44	0.55 ± 0.16	0.046 ± 0.029	0.018 ± 0.0053	0.084 ± 0.052	0.034 ± 0.0076
	Fresh 2	0.95 ± 0.19	1.3 ± 0.18	0.27 ± 0.074	0.02 ± 0.0026	0.00019 ± 0.00026	0.072 ± 0.017	0.00068 ± 0.0010
Putnam County Lakebed,	Aged 2	0.97 ± 0.10	1.4 ± 0.10	0.32 ± 0.067	0.018 ± 0.0013	0.0022 ± 0.0010	0.054 ± 0.011	0.0068 ± 0.0033
Florida, USA	Fresh 7	0.98 ± 0.094	1.5 ± 0.10	0.24 ± 0.024	0.021 ± 0.0021	na	0.085 ± 0.009	na
	Aged 7	0.98 ± 0.10	1.4 ± 0.11	0.34 ± 0.034	0.010 ± 0.0034	0.0044 ± 0.00082	0.029 ± 0.010	0.013 ± 0.0023
	Fresh 2	0.95 ± 0.32	1.2 ± 0.28	0.40 ± 0.14	0.0061 ± 0.0077	0.00036 ± 0.00021	0.015 ± 0.019	0.00089 ± 0.00054
EE	Aged 2	0.97 ± 0.31	1.5 ± 0.33	0.46 ± 0.12	0.0061 ± 0.0077	0.0044 ± 0.00082	0.013 ± 0.017	0.0086 ± 0.0024
Evelgianes, Florina, USA	Fresh 7	0.98 ± 0.11	1.1 ± 0.079	0.40 ± 0.063	0.012 ± 0.0035	0.00026 ± 0.00092	0.029 ± 0.009	0.00064 ± 0.00024
	Aged 7	0.96 ± 0.11	1.6 ± 0.12	0.45 ± 0.063	0.0053 ± 0.007	0.016 ± 0.0031	0.012 ± 0.016	0.036 ± 0.0078
	Fresh 2	0.99 ± 0.057	1.2 ± 0.051	0.18 ± 0.014	0.014 ± 0.00058	0.00087 ± 0.00042	0.077 ± 0.005	0.0047 ± 0.0023
	Aged 2	0.97 ± 0.33	1.3 ± 0.31	0.31 ± 0.081	0.014 ± 0.0067	0.0041 ± 0.0012	0.044 ± 0.020	0.013 ± 0.0028
DUILICO, Malaysia	Fresh 7	0.98 ± 0.018	1.2 ± 0.015	0.21 ± 0.035	0.024 ± 0.0027	0.0014 ± 0.00072	0.11 ± 0.023	0.0067 ± 0.0036
	Aged 7	0.99 ± 0.26	1.5 ± 0.29	0.40 ± 0.079	0.02 ± 0.0041	0.016 ± 0.0033	0.049 ± 0.0040	0.039 ± 0.0035
^a Uncertainty associated with each ratio is calculated based	n each ratio is calc	ulated based on th	e square root of the	individual uncertaintie	on the square root of the individual uncertainties multiplied by the ratio			

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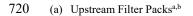
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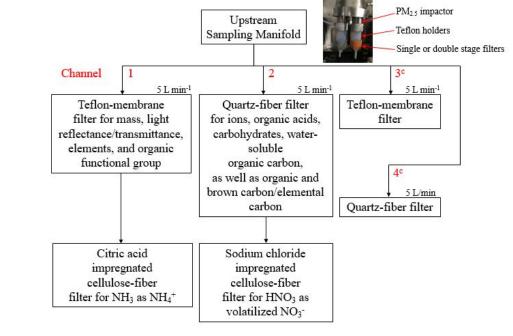




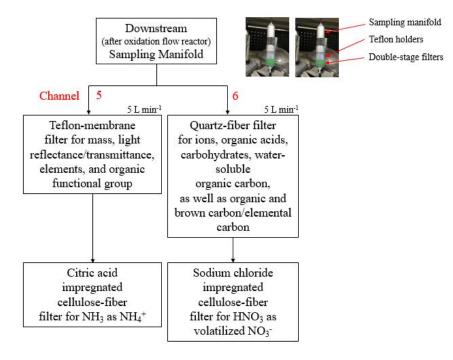








722 (b) Downstream Filter Packs^{a,b}



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^aThe filter types are: 1) Teflon-membrane filter (Teflo[©], 2 µm pore size, R2PJ047, Pall Life Sciences, Port Washington, NY, USA); 2) quartz-fiber filters (Tissuquartz, 2500 QAT-UP, Pall Life Sciences); and 3) citric acid and sodium chloride impregnated cellulose-fiber filters (31ET, Whatman Labware Products, St. Louis, MO, USA).

 $\begin{array}{c} 725\\ 726\\ 727\\ 728\\ 729\\ 730\\ 731\\ 732\\ 733\\ 734\\ 735\\ 736\\ 737\\ 738\\ 737\\ 738\\ 740\\ 741\\ 742\\ 743\\ 744\\ 745\\ 746\\ 747\\ 748\\ \end{array}$ ^bAnalyses include: 1) mass by gravimetry (Model XP6 microbalance, Mettler-Toledo, Columbus, OH, USA); 2) light reflectance/transmittance by UV/Vis spectrometry (Lambda35, Perkin Elmer, Waltham, MA, USA); 3) multiple elements by energy-dispersive x-ray fluorescence (XRF) (Epsilon 5 PANalytical, Westborough, MA, USA); 4) four anions (chloride [Cl], nitrite $[NO_2]$, nitrate $[NO_3]$, and sulfate $[SO_4^{-1}]$; three cations (water-soluble sodium $[Na^{+}]$, potassium $[K^{+}]$, and ammonium [NH4⁺]); and ten organic acids (i.e., formic acid/formate, acetic acid/acetate, lactic acid/lactate, methanesulfonic acid/methanesulfanate, oxalic acid/oxalate, propionate, succinic acid/succinate, maleic acid/maleate, malonic acid/malonate, and glutaric acid/glutarate) by ion chromatography (IC) with conductivity detector (Dionex Model ICS-5000+, Thermo Scientific, Waltham, MA, USA); 5) 17 carbohydrates (i.e., levoglucosan, mannosan, galactosan, glycerol, 2-methylerythritol, arabitol, mannitol, xylitol, erythritol, adonitol, inositol, glucose, galactose, arabinose, fructose, sucrose, and trehalose) by IC with pulsed amperometric detector (Dionex Model ICS3000, Thermo Scientific, Waltham, MA, USA); 6) water-soluble organic carbon (WSOC) by total organic carbon analyzer with non-dispersive infrared (NDIR) detector (Shimadzu Corporation, Kyoto, Japan); 7) organic functional groups by Fourier-Transform Infrared (FTIR) spectroscopy (VERTEX 70, Bruker, Billerica, MA, USA); and 8) organic, elemental, and brown carbon (OC, EC, and BrC) by multiwavelength thermal/optical carbon analyzer (DRI Model 2015, Magee Scientific, Berkeley, CA, USA).

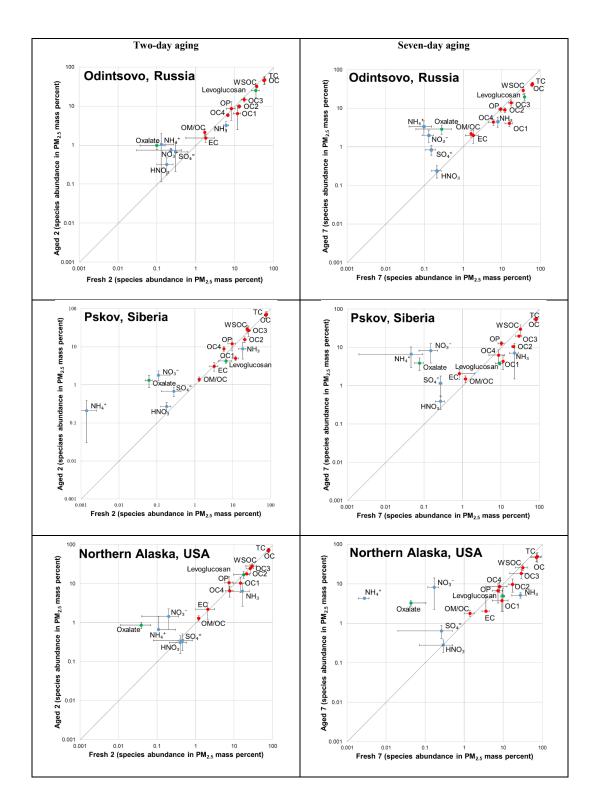
"Teflon-membrane filter samples from Channel 3 are to be analyzed for additional organic nitrogen speciation using Fourier transform-ion cyclotron resonance mass spectrometry (FT-ICR-MS) at the Michigan Technological University. Quartz-fiber filter samples from Channel 4 are to be analyzed for polar and non-polar organics at the Hong Kong Premium Services and Research Laboratory.

749 Figure 1. Filter pack sampling configurations for upstream and downstream channels of the

- 750 oxidation flow reactor.
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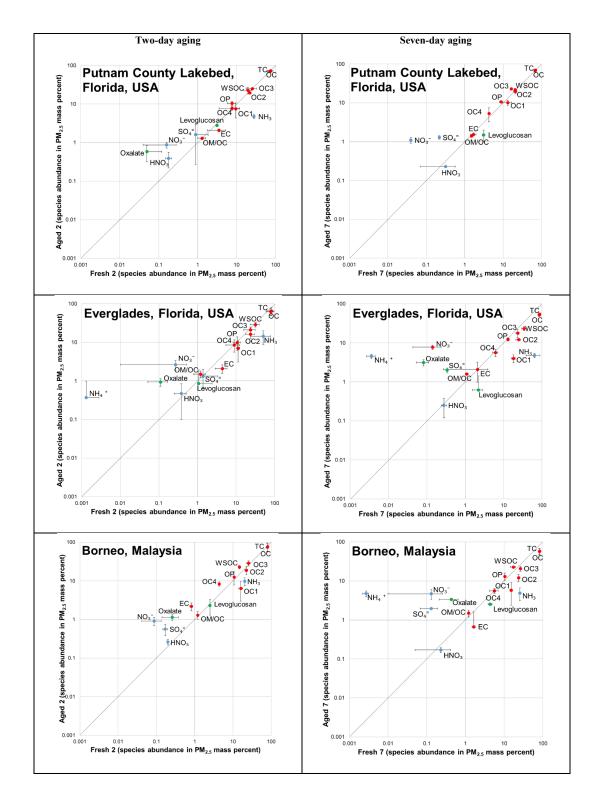












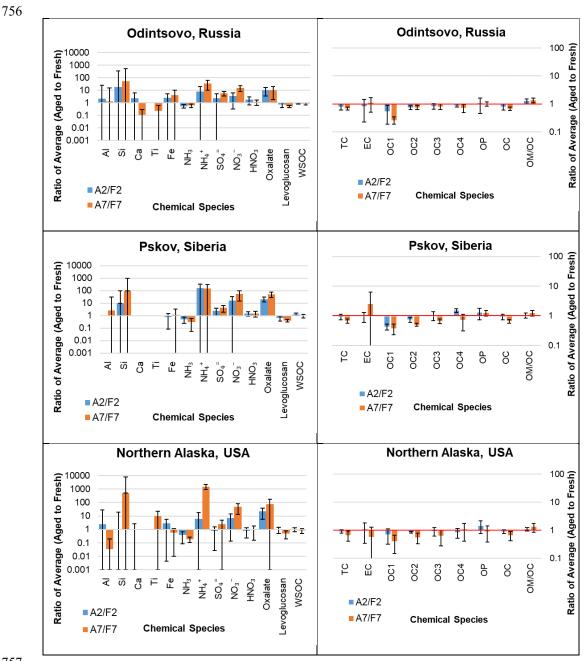




- 752 Figure 2. Comparison between fresh and aged profile chemical abundances for each of the six
- 753 types of peat with 2- and 7-day aging times. Standard deviations associated with averages in x and
- 754 755 y axes are also shown.







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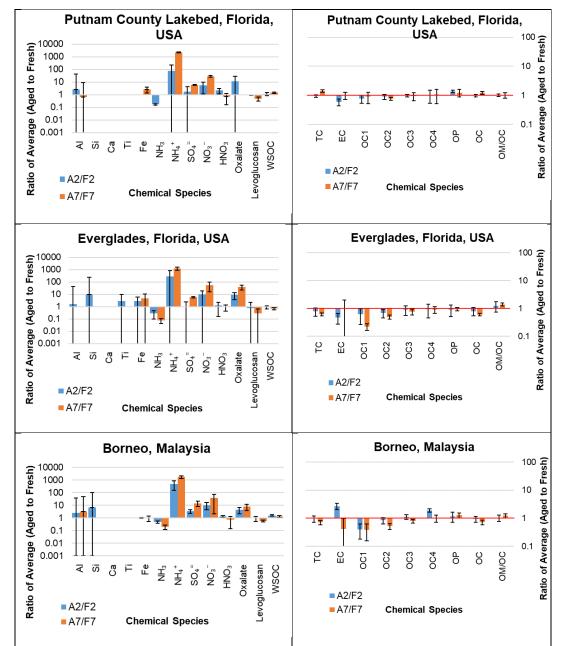
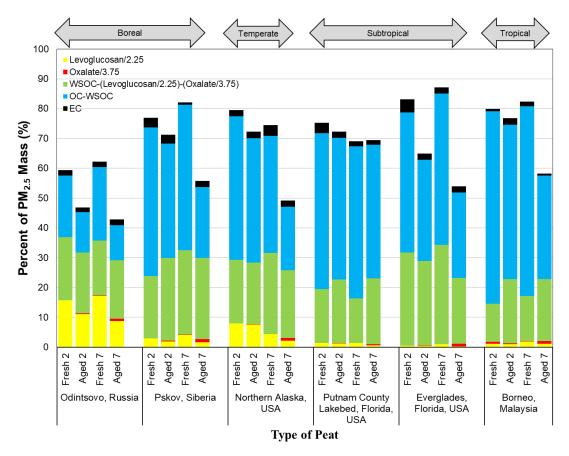


Figure 3. Ratios of average Aged (A) to Fresh (F) chemical species for 2-days (A2/F2) and 7-days
(A7/F7) of atmospheric aging of six types of peats. Vertical bars represent the standard deviations
associated with each ratio. Note that different scales were used in the two Y axes, with 0.001 to
10,000 on the left axis and 0.1 to 100 on the right axis (species abbreviations are shown in Fig. 1;
OM is organic mass).





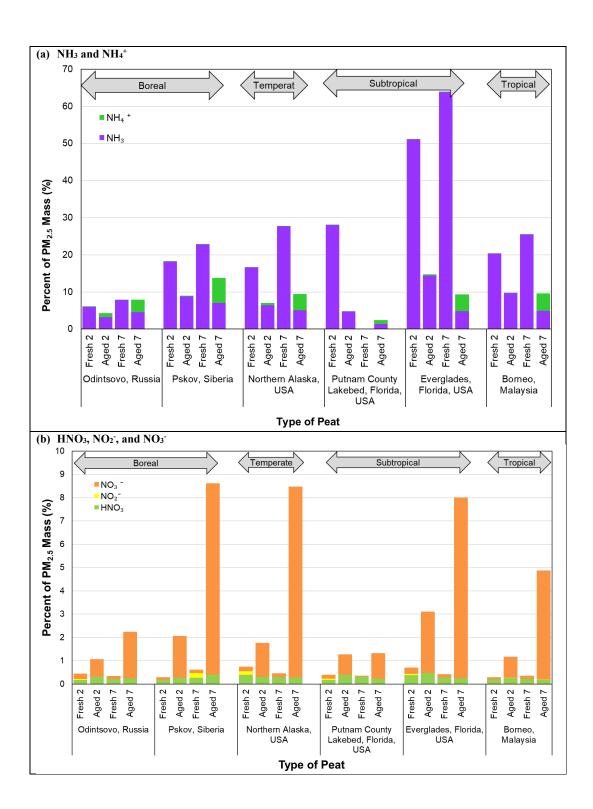


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Figure 4. Abundances of fresh and aged carbon-containing components in $PM_{2.5}$ (levoglucosan $[C_6H_{10}O_5]$ is divided by 2.25 and oxalate $[C_2H_2O_4^-]$ is divided by 3.75 to obtain the carbon content. These levels are subtracted from the water-soluble organic carbon [WSOC] to obtain the remainder, and WSOC is subtracted from organic carbon [OC] to obtain non-soluble carbon. Elemental carbon [EC] is unaltered).







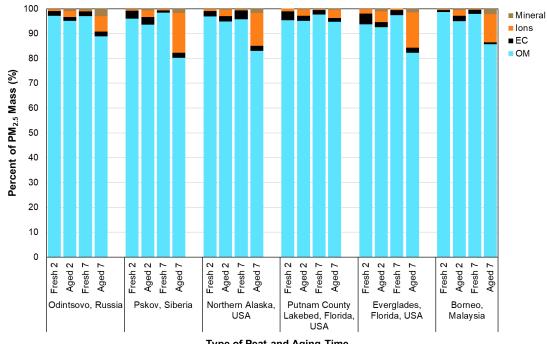




- Figure 5. Comparison of nitrogen species for: a) NH₃ and NH₄⁺; and b) HNO₃, NO₂⁻, and NO₃⁻
- between fresh and aged profiles for six types of peats.
- 773 774







775 776

Type of Peat and Aging Time

Figure 6. Reconstruction of PM_{2.5} mass with organic matter (OM, see Table 3 for OM/OC ratios), 777 elemental carbon (EC), major ions (i.e., sum of NH4⁺, NO3⁻, and SO4⁼), and mineral component 778 (=2.2 Al + 2.49 Si + 1.63 Ca + 1.94 Ti + 2.42 Fe) for six types of peat between fresh and aged 779 profiles.