



1 **Saccharide composition in atmospheric fine particulate**
2 **matter at the remote sites of Southwest China and estimates**
3 **of source contributions**

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18 **Abstract.** Based on source-specific saccharide tracers, the characteristic of
19 biomass burning (BB) and biogenic emissions to saccharides was investigated in three
20 rural sites at Lincang, where covered with 65% of forest in the southwest border of
21 China. The total saccharides accounted for $8.4\pm 2.7\%$ of OC, and $1.6\pm 0.6\%$ of $PM_{2.5}$.
22 The measured anhydrosugars accounted for 48.5% of total saccharides, among which
23 levoglucosan was the most dominant species. The high level of levoglucosan was
24 both attributed to the local BB activities and biomass combustion smoke transported
25 from the neighboring regions of Southeast Asia (Myanmar) and the northern Indian
26 Peninsula. The measured mono (di) saccharides and sugar alcohols accounted for
27 $24.9\pm 8.3\%$ and $26.6\pm 9.9\%$ of the total saccharides, respectively, were both proved to
28 be mostly emitted by direct biogenic volatilization from plant materials/surface soils,
29 rather than as byproducts of polysaccharides breakdown during BB processes. Five
30 sources of saccharides were resolved by non-negative matrix factorization (NMF)
31 analysis, including BB, soil microbiota, plant senescence, airborne pollen and plant
32 detritus with the contribution of 34.0%, 16.0%, 21.0%, 23.7% and 5.3%, respectively.
33 The results provide the information on the magnitude of levoglucosan and
34 contributions of BB, as well as the characteristic of biogenic saccharides, at the
35 remote sites of Southwest China, which can be further applied to regional source
36 apportionment models and global climate models.

37 **1 Introduction**

38 Biomass burning (BB) and biogenic aerosols are thought to play important roles on
39 air quality, human health and climate through direct or indirect effects (Jacobson et al.,
40 2000; Christner et al., 2008; Pöschl et al., 2010; Després et al., 2012; Chen et al., 2017,
41 Tang et al., 2019). Atmospheric saccharides components have been extensively
42 reported to be originate from biomass burning (natural or anthropogenic), suspended
43 soil or dust and primary biological aerosol particles (PBAPs) (e.g., fungal and fern
44 spores, pollens, algae, fungi, bacteria, and plant debris), as well as biogenic secondary
45 organic aerosol (SOA) (e.g., Rogge et al., 1993; Graham et al., 2003; Jaenicke, 2005;
46 Medeiros et al., 2006; Elbert et al., 2007; Fu et al., 2013). As one of the major classes
47 of water-soluble organic compounds, saccharides in atmospheric aerosols have been
48 detected over urban areas, forests, mountains, and remote marine regions (Pashynska
49 et al., 2002; Yttri et al., 2007; Fu et al., 2009; Burshtein et al., 2011; Jia and Fraser,
50 2011; Chen et al., 2013; Pietrogrande et al., 2014; Li et al., 2016a, b). It has been



51 reported that saccharides account for 13-26% of the total organic compound mass
52 identified in continental aerosols, and as much as 63% in oceanic aerosols (Simoneit
53 et al., 2004).

54 Levoglucosan and related anhydrosugar isomers (mannosan and galactosan),
55 produced from pyrolysis of cellulose and hemicellulose, are considered to be
56 relatively stable in the atmosphere (Schkolnik et al., 2005; Puxbaum et al., 2007), and
57 thus have been recognized as specific molecular markers for BB source emissions
58 (Simoneit et al., 1999, 2000; Fraser and Lakshmanan, 2000; Sullivan et al., 2014; Du
59 et al., 2015). However, some studies have challenged this knowledge and proved that
60 levoglucosan alone was not suitable to be a distinct marker for BB in various regions
61 and periods. Because there were evidences that levoglucosan is also emitted from
62 non-BB sources (Wu et al., 2020), such as coal burning (Rybicki et al., 2020; Yan et
63 al., 2018), open waste burning (Kalogridis et al., 2018), incense burning (Tsai et al.,
64 2010), and food cooking (Reyes-Villegas et al., 2018). It was reported that the
65 levoglucosan emission contribution of BB sources ranged from 21.3 to 95.9% (Wu et
66 al., 2021). The current studies in China have reported the value of 2.6-289.1 ng m⁻³
67 and 11.6-1803.1 ng m⁻³ respectively over Beijing and Wangdu in summer (Yan et al.,
68 2019), 2.4-1064.1 ng m⁻³ over Shanghai all year round (Xiao et al., 2018), 15.6-472.9
69 ng m⁻³ over Guangzhou (Zhang et al., 2010), 21.1-91.5 ng m⁻³ over Hongkong (Sang
70 et al., 2011), 60.2-481.9 ng m⁻³ over Xi 'an (Yang et al., 2012), 36.0-1820.9 ng m⁻³
71 over Chengdu (Yang et al., 2012) and 10.1-383.4 ng m⁻³ over the Tibetan Plateau (Li
72 et al., 2019). In north China, high concentrations of levoglucosan is a serious problem
73 due to drastic enhancement of coal and BB for house heating in winter and autumn
74 (Zhang et al., 2008; Zhu et al., 2016). The BB pollution might be exacerbated under
75 unfavorable meteorological conditions, such as in the Chengdu basin (Chen and Xie,
76 2014). In general, BB is an important source of fine particulate matter and with
77 notable contribution to OC in China (Zhang et al., 2008; Cheng et al., 2013; Chen et
78 al., 2017), controls on BB could be an effective method to reduce pollutant emissions.
79 Recently study reported that BB activities have been reduced in China, because the
80 total levoglucosan emission of China exhibited a clear decreasing trend from 2014
81 (145.7 Gg) to 2018 (80.9 Gg) (Wu et al., 2020).

82 Saccharide compounds including a variety of primary saccharides
83 (monosaccharides and disaccharides) and sugar alcohols (reduced sugars) have been
84 measured to estimate the contribution of biogenic aerosols, including fungi, viruses,



85 bacteria, pollen, and plant as well as animal debris (Simoneit et al., 2004; Jaenicke et
86 al., 2007). For instance, arabinol and mannitol have been proposed as biomarkers for
87 airborne fungal spores (Bauer et al., 2008; Zhang et al., 2010; Holden et al., 2011;
88 Liang et al., 2013a, b), because both of them can function as storage or transport
89 carbohydrates to regulate intracellular osmotic pressure (Bauer et al., 2008). Glucose
90 and sucrose are thought to originate from natural biogenic detritus, including
91 numerous microorganisms, plants, and animals (Simoneit et al., 2004; Tominaga et al.,
92 2011). As the oxidation products of isoprene, methyltetrols (including
93 2-methylthreitol and 2-methylerythritol) have been suggested as tracers of
94 isoprene-derived SOA (Claeys et al., 2004; Kleindienst et al., 2007; Ding et al., 2016).
95 In the previous study, the contributions of fungal spores to OC were estimated to be
96 $14.1 \pm 10.5\%$ and $7.3 \pm 3.3\%$ respectively at the rural and urban sites of Beijing
97 (Liang et al., 2013b). Airborne pollen and fungal spores contributed 12-22% to the
98 total OC in ambient aerosols collected in Toronto (Womiloju et al., 2003). Jaenicke
99 (2005) found that PBAPs can comprise from 20 to 30% of the total atmospheric PM
100 ($>0.2 \mu\text{m}$) from Lake Baikal (Russia) and Mainz (Germany). However, there is still a
101 limited number of studies on quantifying the abovementioned biogenic aerosol
102 contributions to ambient aerosol.

103 Lincang located in the southwest border of China is a traditional agricultural area of
104 Yunnan province, where planting a large area of tea, sugar cane, rubber, macadamia
105 nuts, etc. It is the largest production base of black tea and macadamia nut in China.
106 The forest coverage rate of Lincang reaches to 65%. It has a wide variety of plant
107 species, and has 6 nature reserves covering an area of $\sim 222,000$ hectares, accounting
108 for 8.56% of the total area. As a residential area for ethnic minorities, Lincang has
109 unique culture, humanity and living habits. A high portion of houses with wood
110 burning used for cooking in villages in proximity and a large area of Southeast Asia
111 constitute, and forest fires were frequently happened in this area especially in the dry
112 seasons (March-April). All implies that there are abundant biogenic aerosols in the
113 area, and BB pollution may be an important potential source of air pollution. However,
114 little information on the magnitude of biogenic and BB tracers in this area is available.
115 The contributions of biogenic aerosol and BB, and BB types are poorly understood.

116 In this study, the sampling were conducted from March 8 to April 8, 2019 at three
117 mountaintop sites of Lincang, where is an ideal site for investigating the BB emission
118 characteristics. BB tracers (including anhydrosugars and K^+) and biogenic aerosol



119 tracers (primary saccharides and sugar alcohols) were measured to gain the
120 information on source and contributions of BB and biogenic emissions in PM_{2.5} over
121 the rural Lincang. This study would be useful and valuable for provide reliable
122 information on sources and magnitudes of saccharides involving rural BB and
123 biological emissions in China.

124 **2 Experimental section**

125 **2.1 Aerosol sampling**

126 The PM_{2.5} samples were simultaneously collected on three mountaintop sites,
127 respectively of Datian (24.11° N, 100.13° E, 1960 m asl), Dashu (24.12° N, 100.11° E,
128 1840 m asl) and Yakoutian (24.12° N, 100.09° E, 1220 m asl), in Lincang, Yunnan
129 Province, China, which are located ~300 km west to Kunming (the capital of Yunnan
130 province) and ~120 km east from the Burma border. These sites are surrounded by
131 massive mountains and scattering villages without obvious nearby traffic or major
132 industry emissions. Each sampling was performed over a 23.5 h period every day, and
133 was collected on quartz by high-volume air samplers (Thermo) equipped with a size
134 selective inlet to sample PM_{2.5} at a flow rate of 1.13 m³ min⁻¹. Altogether, 91 samples
135 were collected during 8 March to 9 April in 2019.

136 Quartz filters (Whatman, 8 × 10 in.) were prebaked at 550 °C for 4 h in a muffle
137 furnace to remove organic material, and were then stored in pre-baked aluminum foils.
138 The samples were stored at about -20 °C in a refrigerator until analysis. Field blanks
139 were collected by mounting filters in the sampler without air flow to replicate the
140 environmental exposure. The data reported were corrected by the blanks at the
141 sampling sites.

142 **2.2 Measurements**

143 The concentrations of organic carbon (OC) and elemental carbon (EC) were
144 measured using a Multiwavelength Carbon Analyzer (DRI Model 2015; Atmoslytic
145 Inc., USA). Typically, a 0.58 cm² punch of the filter was placed on a boat inside the
146 thermal desorption chamber of the analyzer, and then stepwise heating was applied.
147 Carbon fractions were obtained following the Interagency Monitoring of Protected
148 Visual Environments (IMPROVE-A) thermal/optical reflectance (TOR) protocol
149 (Chow et al., 2007). Replicate analyses were conducted once every ten samples.
150 Blank sample was also analyzed and used to correct the sample results.

151 A punch (4.7 cm²) of each quartz filter was ultrasonically extracted with 10.0 mL of



152 de-ionized water (resistivity = 18.2 MU) for 40 min. The aqueous extracts were
153 filtrated through syringe filters (PTFE, 0.22 μm) to remove insoluble materials. Ion
154 chromatography (Metrohm, Switzerland) coupling with Metrosep C6-150 and A6-150
155 columns was used to detect water-soluble ions (Cl^- , NO_3^- , PO_4^- , SO_4^{2-} , Na^+ , NH_4^+ ,
156 K^+ , Mg^{2+} , and Ca^{2+}) with a detection limit (DL) range of 0.001-0.002 $\mu\text{g m}^{-3}$.

157 Five saccharide alcohols (glycerol, erythritol, inositol, arabitol and mannitol) and
158 five primary saccharides (fructose, glucose, mannose, sucrose and trehalose), together
159 with three anhydrosugars (levoglucosan, mannosan and galactosan) were quantified
160 by an improved high performance anion-exchange chromatography system coupled
161 with a pulsed amperometric detector (HPAEC-PAD) (Engling et al., 2006; Caseiro et
162 al., 2007; Zhang et al., 2013). This method developed by Engling et al. (2006) was
163 validated to be a powerful method for the detection of carbohydrates without
164 derivatization techniques, and has been successfully applied for the atmospheric
165 tracers (e.g., Zhang et al., 2010; Holden et al., 2011; Liang et al., 2013a, b; Li et al.,
166 2016a, b; Kalogridis et al., 2018; Yan et al., 2018). The separation of the saccharides
167 was performed on an ion chromatograph (Metrohm, Switzerland) equipped with a
168 Metrosep Carb 2-250 analytical column and a guard column. The aqueous eluent of
169 sodium hydroxide and sodium acetate was pumped by a dual pump module at a flow
170 rate of 0.4 mL min^{-1} . The low concentration of 50 mM sodium hydroxide and 10 mM
171 sodium acetate (eluent A) was applied to pump 1, while the high concentration of 250
172 mM sodium hydroxide and 50 mM sodium acetate (eluent B) was applied to pump 2.
173 The gradient generator was set as: 0-10 min, 100% of eluent A; 10-20 min, 50% of
174 eluent A and 50% of eluent B; 20-50 min, 100% of eluent B; 50-60 min, 100% of
175 eluent A for equilibration. The extraction efficiency of this analytical method was
176 determined to be better than 90% based on analysis of quartz filters spiked with
177 known amounts of mannitol. The method DL of the referred carbohydrate compounds
178 were 0.005-0.01 mg L^{-1} . All carbohydrate species were below detection limits in the
179 field blanks.

180 **2.3 Other data**

181 The meteorological parameters, including temperature (T), relative humidity (RH),
182 solar irradiation (W m^{-2}), and rainfall (mm) were obtained from the Physical Sciences
183 Laboratory of NOAA (<https://psl.noaa.gov>). The temporal changes in meteorological
184 variables over the observation sites during the sampling periods are shown in Figure



185 S1.

186 In order to characterize the origin and transport pathway of the air masses to the
187 sampling sites, 72 h back-trajectories of the aerosol were calculated using Hybrid
188 Single-Particle Lagrangian Integrated Trajectory (HYSPLIT) model developed by
189 NOAA/ARL (Draxler and Hess, 1998) via NOAA ARL READY Website
190 (<http://ready.arl.noaa.gov/HYSPLIT.php>) with an endpoint height of 1500 m. To
191 investigate the influence of BB emissions, fire pixel counts were obtained from
192 Moderate Resolution Imaging Spectroradiometer (MODIS) observations on NASA
193 satellites (<https://earthdata.nasa.gov/>).

194 **2.4 Statistical analysis**

195 A Pearson's correlation test was performed using the Statistical Product and Service
196 Solutions (SPSS) software for the dataset containing ambient concentrations of the
197 measured saccharides, inorganic ions and solar irradiation. Non-negative matrix
198 factorization (NMF) analysis was utilized to resolve potential emission source and
199 estimate their contribution to atmospheric saccharides. NMF introduced by Lee and
200 Seung (1999) was similar to positive matrix factorization (PMF). Both methods find
201 two matrices (termed the contribution matrix of W and the source profile matrix of H)
202 to reproduce the input data matrix (V) using the factorization approach ($V = WH$) as a
203 positive constraint ($W \geq 0$ and $H \geq 0$). However, PMF forces the negative factors to be
204 positive, but NMF method only retains nonnegative factors. NMF minimizes the
205 conventional least-squares error and the generalized Kullback-Leibler divergence
206 (Shang et al., 2018). Therefore, the results obtained from NMF are more responsive to
207 the original characteristics of input data set and less number of factors will be
208 extracted (Zhang et al., 2019). Half of the DL was used for the value below the
209 detection limit. In this study, galactosan, mannose and inositol were excluded because
210 their concentration was mostly below the DL. Concentrations of the other ten
211 saccharide species for the total 91 samples were subjected for NMF analysis. The
212 uncertainties in NMF analysis were estimated as 0.3 plus the analytical detection limit
213 according to the method of Xie et al. (1999). The constant 0.3 corresponding to the
214 log (Geometric Standard Deviation, GSD) was calculated from the normalized
215 concentrations for all measured species, and was used to represent the variation of
216 measurements.

217 **3 Results and Discussion**



218 **3.1 Saccharides concentration and composition**

219 The temporal variations of PM_{2.5} mass, OC, EC, and various saccharides measured
220 in all samples are shown in Figure 1. A statistical summary of all the data are listed in
221 Table S1. During the sampling periods, the PM_{2.5} mass concentrations ranged between
222 13.7-87.8 $\mu\text{g m}^{-3}$ with average values of 41.8 $\mu\text{g m}^{-3}$. The concentrations of OC and
223 EC varied between 2.5-22.4 and 0.3-4.3 $\mu\text{g m}^{-3}$ with average values of 8.4 and 1.7 μg
224 m^{-3} , respectively. OC accounted for 19.9 \pm 3.7% of total PM_{2.5} mass. The ambient
225 concentrations of the total saccharides varied between 244.5-1291.6 ng m^{-3} with
226 average values of 638.4 ng m^{-3} . The total saccharides quantified in PM_{2.5} accounted
227 for 8.4 \pm 2.7% (range: 3.8%-20.6%) of the OC, and accounted for 1.6 \pm 0.6% (range:
228 0.6%-3.0%) of the PM_{2.5}. Figure 2 presents the mean concentration levels of 12
229 measured saccharide compounds, categorized into anhydrosugar, sugar and sugar
230 alcohol, as well as the relative contribution of these saccharides for the all samples.
231 The values for each site are shown in Figure S2. The absolute concentration and
232 relative contribution of each saccharide displayed no distinct variation between the
233 three sites.

234 **3.1.1 Anhydrosugars**

235 The mean concentrations of levoglucosan and mannosan were 287.7 and 31.6 ng
236 m^{-3} , respectively with a range of 95.6-714.7 and 0-134.7 ng m^{-3} for all the 91 samples.
237 Galactosan was only detected in 6 samples, with a range of 2.5-5.5 ng m^{-3} . The
238 anhydrosugars accounted for 48.5% of total measured saccharides. Levoglucosan was
239 the most dominant specie among all the saccharides. The mean levoglucosan
240 concentration in this study was comparable to the value at urban Beijing collected in
241 spring of 2012 (above 200 ng m^{-3}) (Liang et al., 2016) and at urban Xi'an collected in
242 winter of 2015 (268.5 ng m^{-3}) (Wang et al., 2018). It was higher than the value at
243 rural Tengchong mountain site (193.8 ng m^{-3}) (Sang et al., 2013), at urban Shanghai
244 collected in spring of 2012 (66.0 ng m^{-3}) (Li et al., 2016) and at urban Hong Kong
245 collected in spring of 2004 (36.0 ng m^{-3}) (Sang et al., 2011), as well as at urban
246 Beijing collected in summer of 2013 (49.4 ng m^{-3}) (Yan et al., 2019), but was lower
247 than that at a rural site of Xi'an (0.93 mg m^{-3}) (Zhu et al., 2017) and at a rural site in
248 eastern central India (2258 ng m^{-3}) (Nirmalkar et al., 2015). During the observation
249 period, several instances of elevated levoglucosan occurred, peaked on March 8, 16,
250 23 and April 1. It was thought the ambient levoglucosan were primarily attributed to



251 domestic biomass fuel burning, the high levoglucosan emission on these peak days
252 might be from open BB events.

253 Regression analyses of levoglucosan and the other two anhydrosugars (mannosan,
254 galactosan) are shown in Figure 3a. Levoglucosan was highly correlated with
255 mannosan and galactosan, with coefficients of determination (R) of 0.81 ($P < 0.01$)
256 and 0.89 ($P < 0.01$), respectively, indicating similar combustion sources of them. The
257 ratios of levoglucosan to mannosan (L/M) and mannosan to galactosan (M/G) had
258 been employed to identify the specific types of BB, although these ratios were quite
259 variable (Fabbri et al., 2009; Sang et al., 2018). Previous studies suggested that L/M
260 ratios for burning of softwood were 3-10, hardwood were 15-25, and those from crop
261 residues were often above 40 (Cheng et al., 2013; Zhu et al., 2015; Kang et al., 2018).
262 The average L/M and M/G ratios were statistically reported as 32.6 and 1.2 for crop
263 residues combustion, 4.0 and 3.9 for softwood combustion, 21.5 and 1.5 for hardwood
264 combustion, respectively (Sang et al., 2013; Shen et al., 2018). In this study, the ratios
265 of L/M and M/G ranging from 4.7 to 16.1 (mean: 9.7, $n=91$) and from 3.9 to 6.1
266 (mean: 4.8, $n=6$), respectively, crudely indicating major contribution from softwood
267 burning. The sample collected during 31 March-1 April and during 8-10 March
268 respectively had considerably lower and higher concentrations of mannosan than
269 predicted by the levoglucosan-mannosan regression model (Figure 3a). The results
270 suggested that BB aerosols collected during 31 March-1 April ($L/M = 11.52 \pm 1.34$)
271 and during 8-10 March ($L/M = 6.57 \pm 0.53$) have originated from different types of
272 BB as compared with the remaining sampling periods ($L/M = 9.34 \pm 1.20$). Therefore,
273 the high levoglucosan emission during 31 March-1 April and during 8-10 March
274 might be from different open BB events, possibly an open agricultural waste burning
275 event or a forest fire, whilst the BB of most sampling days originated from biomass
276 fuel for domestic cooking and heating. It was worth noting that these peak days
277 neared the Qingming Festival. Another possibility of BB events was that people
278 burned ghost money to sacrifice ancestor according to Chinese tradition.

279 Anhydrosugars and water-soluble potassium (K^+) have been both widely utilized as
280 source tracers of BB emissions (e.g., Puxbaum et al., 2007; Wang et al., 2007; Zhang
281 et al., 2008; Engling et al., 2011). The daily variation on concentrations of
282 levoglucosan and K^+ are shown in Figure S3, the regression analysis of K^+ and three
283 anhydrosugars is shown in Figure 3b. K^+ was weakly correlated with levoglucosan,
284 mannosan, and galactosan, with R values of 0.33, 0.28, and 0.74, respectively. It could



285 be explained by the additional emissions of K^+ from soil and sea water. Since Lincang
286 is far from the coast, sea salt could be negligible. Because the inhomogeneity of
287 crustal K^+ associated with soil types, it was difficult to fully account for crustal K^+
288 contributions from soil (Harrison et al., 2012; Cheng et al., 2013). The ratio of
289 levoglucosan to K^+ (L/K^+) was also used to track possible sources of BB in the
290 previous studies. The ratios of L/K^+ strongly depended on BB processes, namely
291 smoldering and flaming. Studies suggested that relatively high L/K^+ ratios were
292 obtained from smoldering combustion at low temperatures compared with flaming
293 combustion (Schkolnik et al., 2005; Lee et al., 2010). Previous results showed the
294 emissions from the combustion of crop residuals such as rice straw, wheat straw and
295 corn straw exhibited comparable L/K^+ ratios, typically below 1.0. The averages of
296 L/K^+ ratios in this study was 0.48 ± 0.20 , which was higher than the ratio for corn
297 straw (0.21 ± 0.08), but was lower than the ratio for Asian rice straw (0.62 ± 0.32)
298 (Cheng et al., 2013). In this study, higher L/K^+ ratios were observed during 8-10
299 March (1.20 ± 0.19) than those during 31 March-1 April (0.40 ± 0.13), which
300 suggested that the open fire event during 8-10 March was more possibly due to
301 smoldering combustion of residues at low temperatures.

302 Figure 3c and 3d show the scatter plots and regression analyses of K^+ versus $PM_{2.5}$,
303 OC and EC, and levoglucosan versus $PM_{2.5}$, OC and EC, respectively. Linear
304 regression of K^+ on $PM_{2.5}$, OC and EC resulted in R values of 0.64, 0.63 and 0.62,
305 respectively, which were generally higher than those of levoglucosan on $PM_{2.5}$, OC
306 and EC, with R values of 0.40, 0.54 and 0.48, respectively. It showed that K^+ is more
307 highly correlated with $PM_{2.5}$, OC and EC. This can be explained by either the
308 photo-oxidative decay of levoglucosan (Hennigan et al., 2010) and/or different types
309 of BB processes (Schkolnik et al., 2005; Lee et al., 2010). Even so, the results
310 supported that the BB posed great impact on fine aerosols. The ratio of levoglucosan
311 to $PM_{2.5}$ ($L/PM_{2.5}$) is also helpful in distinguishing the contributions of different
312 levoglucosan sources (Wu et al., 2020). The ratios of $L/PM_{2.5}$ in this study was
313 0.0041-0.0162 (mean: 0.0072), indicating that levoglucosan emission in the areas
314 might mainly come from woods (0.01-0.09) and crop straws (0.001-0.008), not
315 excluding incense burning (0.001-0.007), ritual item burning (0.004-0.086), and meat
316 cooking (0.005-0.06). However, it is certainly that it was not from corncob burning
317 (0.0092-0.032), coal burning (0.0001-0.001) and waste incineration (0.0022).

318 An empirical ratio of levoglucosan to OC (8.2%), calculated from main types of



319 Chinese cereal straw (rice, wheat and corn) based on combustion chamber
320 experiments (Zhang et al., 2007), was used to estimate the BB-derived OC. The
321 average mass concentration of BB-derived OC was 3534.4 ng m^{-3} , whilst the
322 contributions of BB to OC was 41.3%, with a large range of 19.1 to 81.3%. The
323 contributions are higher than those previous reported, such as 6.5-11% in Hong Kong
324 (Sang et al., 2011), 18-38% in Beijing (Zhang et al., 2008), 18.9-45.4% over
325 southeastern Tibetan Plateau (Sang et al., 2013), 26.4-30.2% in Xi'an (Zhang et al.,
326 2014). The large range of 19.1 to 73.9% revealed that the daily contribution of BB
327 varied greatly, suggesting open BB event or forest fire happened occasionally.
328 However, the contribution apportionment of primary BB might be underestimated due
329 to the degradation of levoglucosan during atmospheric aging of BB influenced air
330 mass after long-range transport (Hennigan et al., 2010; Mochida et al., 2010; Lai et al.,
331 2014). Moreover, Wu et al (2020) have reported that the total levoglucosan emission
332 of China exhibited a clear decreasing trend and biomass burning activities have been
333 reduced in China. However, it is noteworthy that the mean concentration of
334 levoglucosan (287.7 ng m^{-3}) and the biomass burning contributions to OC (41.3%) at
335 Lincang mountain site in this study are both higher than the values of 191.8 ng m^{-3}
336 and 28.4% at Tengchong mountain site in 2004 spring (Sang et al., 2013). The result
337 suggested no significant reduction in BB emissions in Southwest Yunnan Province to
338 some extent.

339 **3.1.2 Mono (di) saccharides**

340 The total concentrations of five mono (di) saccharides, including glucose, fructose,
341 mannose, sucrose and trehalose, were $25.2\text{-}373.7 \text{ ng m}^{-3}$ (mean: 158.9 ng m^{-3}), which
342 contributed $24.9\pm 8.3\%$ of the total measured saccharides. The mean values of glucose,
343 fructose, mannose, sucrose and trehalose were 31.2, 24.6, 2.7, 86.4 and 13.8 ng m^{-3} ,
344 respectively. Sucrose was the dominant mono (di) saccharides. The results was
345 consistent with the previous studies of Yttri et al. (2007), Jia et al. (2010), and Fu et al.
346 (2012), which had found that sucrose was one of the dominate specie in spring fine
347 aerosols. The ruptured pollen may be an important source of sucrose in particular of
348 spring blossom season (Yttri et al., 2007; Fu et al., 2012; Miyazaki et al., 2012). In
349 spring and early summer, farmland tilling after the wheat harvest causes an enhanced
350 exposure of soil containing wheat roots to the air, which is beneficial to the release of
351 sucrose stored in the root (Medeiros et al., 2006), thus resulting in a sharply increased



352 sucrose.

353 It was reported that sugars, such as glucose, sucrose and fructose, could be emitted
354 from developing leaves (Graham et al., 2003). Glucose could be released from both
355 soils and plant materials (e.g., pollen, fruits and their fragments) (Graham et al., 2003;
356 Simoneit et al., 2004; Fu et al., 2012). Glucose and sucrose were rich in biologically
357 active surface soils (Rogge et al., 2007). In this study, the positive correlations were
358 found between sucrose and glucose ($R = 0.52$) (Table S2), suggesting a similar origin
359 of glucose and sucrose in this study. Glucose and fructose have also been identified as
360 a minor product of cellulose pyrolysis, because they were found to be enrich in BB
361 emission (Nolte et al., 2001), and correlated well with K^+ (Graham et al., 2002) and
362 levoglucosan (Kang et al., 2018). Herein, no significant correlation were found
363 between K^+ , levoglucosan and these mono (di) saccharides. Therefore, the detected
364 glucose, fructose and sucrose might mostly be emitted by direct volatilization from
365 plant materials/surface soils, rather than as products of polysaccharides breakdown
366 during BB processes. The high abundance of sucrose, as well as glucose and fructose
367 were responsible for biogenic aerosols associated with developing leaves and flowers,
368 and surface soil suspension.

369 Trehalose as a stress protectant of various microorganisms and plants (Medeiros et
370 al., 2006; Jia and Fraser, 2011) was found to be abundant in the fine mode soil, and
371 has been proposed as a marker compound for fugitive dust from biologically active
372 surface soils (Simoneit et al., 2004; Medeiros et al., 2006; Rogge et al., 2007; Fu et al.,
373 2012). Previous study found a positive correlation between trehalose and calcium
374 (Nishikawa et al., 2000). In this study, there was no significant correlation between
375 trehalose and calcium. Besides, mannose has been reported to be one of the major
376 monosaccharide components in phytoplankton, which is originate from marine
377 biological fragments (Tanoue and Handa, 1987). Mannose was detected in only a few
378 samples and presented in low concentrations in this study.

379 **3.1.3 Sugar alcohols**

380 Five sugar alcohol compounds, including glycerol, threitol, mannitol, arabitol and
381 inositol were detected in $PM_{2.5}$. These reduced sugars are often reported to be related
382 with the plant senescence and decay by microorganisms (Simoneit et al., 2004; Tsai et
383 al., 2013), and are produced by fungi, lichens, soil biota and algae (Elbert et al., 2007;
384 Bauer et al., 2008). The average concentration of the total sugar alcohols were 159.9



385 ng m⁻³ with a range of 53.1-254.0 ng m⁻³, which contributed 26.6±9.9% of the total
386 measured saccharides. Glycerol has been widely found in soil biota (Simoneit et al.,
387 2004). Herein, glycerol was the second most abundant saccharide with an average
388 concentration of 123.7 ng m⁻³, which comprised 5.1-44.6% (mean: 22.6%) of the total
389 measured saccharides.

390 Mannitol and arabitol have been proposed as tracers for airborne fungal spores
391 (Elbert et al., 2007; Bauer et al., 2008; Zhang et al., 2010; Burshtein et al., 2011).
392 Mannitol and arabitol were detected with a concentration range of 0.0- 38.6 ng m⁻³
393 (14.7 ng m⁻³) and 0.0-21.1 ng m⁻³ (5.8 ng m⁻³), respectively. The mean
394 concentrations of mannitol and arabitol were comparable to those (mean 11.3 and 9.1
395 ng m⁻³) reported in the Beijing spring aerosols (Liang et al., 2013b), but were lower
396 than those (mean 21.9 and 8.43 ng m⁻³) in the Mediterranean summer aerosols
397 (Burshtein et al., 2011) and (30 and 24 ng m⁻³) at Hyytiälä, Finland in summer (Yttri
398 et al., 2011). Poor correlations ($r = 0.38$) were found among mannitol and arabitol in
399 this study. Nevertheless, a positive correlations was found between trehalose and
400 mannitol ($r = 0.79$, $P < 0.05$) (Table S2).

401 In the previous studies, the total measured mannitol has been measured and used
402 for estimating the contribution of fungal spores to organic carbon (Elbert et al., 2007;
403 Bauer et al., 2008; Zhang et al., 2010). A factor of mannitol per spore (0.49 ± 0.20 pg)
404 was used to calculate the number concentrations of fungal spores (Liang et al., 2013a),
405 then the carbon content of fungal spores can be calculated using a conversion factor of
406 13 pg C per spore obtained earlier as the average carbon content of spores from nine
407 airborne fungal species, with an uncertainty of 20% (Bauer et al., 2008). The
408 diagnostic tracer ratio of mannitol to OC was calculated to be 0.0377 according to
409 these researches (Bauer et al., 2008; Liang et al., 2013a), and then used to estimate the
410 contribution of fungal spores to the OC. The average mannitol concentrations were
411 14.7 ± 11.2 ng m⁻³ during the observation period. The average spore-derived OC was
412 calculated to be 390.3 ng C m⁻³, which contributed of 4.9% of the total OC.

413 Claeys et al. (2004) firstly identified two diastereoisomeric 2-methyltetrols as
414 oxidation products of isoprene in the Amazonian rain forest aerosols. Henceforward,
415 2-methyltetrols has been used as tracers for isoprene-derived SOA (Liang et al., 2012;
416 Fu et al., 2016; Yan et al., 2019). In the previous studies, erythritol was often
417 quantified as surrogate of 2-methyltetrols (2-methylthreitol and 2-methylerythritol)
418 due to lack of standards (Claeys et al., 2004; Ding et al., 2013; Ding et al., 2016). In



419 this study, concentration ranges of erythritol were 0.4-19.8 ng m⁻³ (mean 11.1 ng m⁻³).
420 The values of inositol ranged from 0.0 to 22.8 ng m⁻³ with average values of 5.8 ng
421 m⁻³. Moreover, the sugar alcohols not only originates from biological emissions, but
422 also derives from BB (Wan and Yu, 2007; Jia et al., 2010). Different levels of glycerol,
423 arabitol, mannitol, erythritol and inositol in fine particles have been found during
424 burning of crop residues and fallen leaves as well as indoor biofuel usage for heating
425 and cooking (Graham et al., 2002; Burshtein et al., 2011; Wang et al., 2011; Yang et
426 al., 2012; Kang et al., 2018). Herein, only inositol exerted correlation with
427 levoglucosan ($r = 0.42$), suggesting inositol may be linked to biomass combustion
428 sources. Hence, the primarily source of sugar alcohols associated with fine particles
429 was biogenic aerosols at observation sites.

430 **3.2 Sources and transport**

431 Since the distinct concentration of studied compounds was caused by different
432 emission sources arising from different wind direction, the 72 h backward trajectories
433 for the samples at Dashu site (24.12° N, 100.11° E) and the spatial distribution of the
434 fire spots (March 8-April 8, 2019) were calculated to understand the source of
435 saccharides in aerosol (Figure 4). The analysis of air mass backward trajectories
436 suggested that the air mass over Lincang were almost from the westerlies during the
437 sampling periods, and could be separated into two episodes of remote western source
438 over 2000 meters and local western source below 2000 meters, as shown in wine red
439 and green lines. The air masses over 2000 meters were mainly from the regions of
440 South Asia, such as India and Myanmar. The air masses below 2000 meters were
441 mainly attributed to the local air flows in the east of Hengduan Mountains.

442 Mean concentrations of saccharide compounds, as well as the contribution of them,
443 for the episodes over and below 2000 meters are shown in Figure 5. The mean
444 concentration of levoglucosan and mannosan for the below 2000 meters samples
445 (327.4 and 35.6 ng m⁻³) were higher than those for the over 2000 meters samples
446 (250.3 and 27.3 ng m⁻³). The anhydrosugars accounted for 49.2% and 36.9% of total
447 saccharides, respectively for the below and above 2000 meters samples. It implied
448 that the levoglucosan at the observation site was both attributed to the local BB
449 activities and biomass burning smoke transported from the neighboring regions of
450 Southeast Asia (Myanmar) and the northern Indian Peninsula. These results were in
451 agreement with the fact that residents across the Southeast Asia used to utilize woods



452 as energy source to cook and generate heat.

453 While for glucose, fructose and sucrose, it was a little higher in the over 2000
454 meters samples (mean 33.5, 26.4 and 106.2 ng m⁻³) than that in the below 2000
455 meters samples (mean 29.2, 22.9 and 67.8 ng m⁻³). It implied that biogenic aerosols
456 (such as ruptured pollen) carrying sugars could pass long distance, which was
457 supported by previous study, which have observed long-range atmospheric transport
458 of fine pollen from the Asian continent to the remote island Chichi-Jima under the
459 influence of westerlies (Rousseau et al., 2008). Although the pollen are usually coarse
460 with various shapes and hard shells, which results in the relatively short retention time
461 in the atmosphere. Therefore, it could be concluded that, in addition to the local
462 pollen, the concentration of sucrose in Lincang was also influenced by the transport of
463 airborne pollen derived from South Asia areas.

464 **3.3 Source apportionment of saccharides**

465 Based on the compositional data of saccharides and key representative markers for
466 difference sources, five factors associated to the emission sources of saccharides were
467 finally resolved by NMF. As shown in Figure 6a, factor 1 was characterized by high
468 level of levoglucosan (71.8%) and mannosan (78.7%), suggesting the source of BB
469 (Simoneit et al., 1999; Nolte et al., 2001). Factor 2 was characterized by trehalose
470 (99.9%) and mannitol (100.0%), and was enriched in the other saccharides
471 components, i.e., arabitol (44.1%), glucose (29.6%), erythritol (18.2%), glycerol
472 (17.8%), levoglucosan (14.7%), and sucrose (8.6%). These saccharide compounds
473 had all been detected in the suspended soil particles and associated microbiota (e.g.,
474 fungi, bacteria and algae) (Simoneit et al., 2004; Rogge et al., 2007). Hence, this
475 factor was attributed to suspended soil dust and soil microbiota. Factor 3 has high
476 levels of glycerol (71.4%) and erythritol (58.2%), and showed loadings of glucose
477 (12.8%) and fructose (11.8%). This factor was thought as the sources from plant
478 senescence and decay by microorganisms. Factor 4 exhibited a predominance of
479 sucrose (78.7%), and showed loadings of glucose (17.2%), arabitol (11.8%). This
480 factor was regarded as the source of airborne pollen, because pollen is the
481 reproductive unit of plants and contains these saccharides and saccharide alcohols as
482 nutritional components (Bieleski, 1995; Miguel et al., 2006; Fu et al., 2012). Factor 5
483 characterized by the dominance of fructose (88.2%) was resolved, and was enriched
484 in glucose (38.2%) and arabitol (21.2%), thus it could be regarded as the source of



485 plant detritus.

486 The pie charts in Figure 6b shows the contribution of each source to total
487 saccharides. BB of factor 1 (34.0%) was found as the dominant contributor to total
488 saccharides. Factor 2-5 could all be labeled to a biogenic source accounting for a total
489 contribution of 66.0%. The sources of soil microbiota (factor 2), plant senescence
490 (factor 3), airborne pollen (factor 4) and plant detritus (factor 5) respectively
491 contributed 16.0%, 21.0%, 23.7% and 5.3% to total saccharides. During the sampling
492 periods, daily variations on proportion of the five factors are shown in Figure S4.
493 Factor 2 soil microbiota emissions could be associated to soil reclamation and
494 cultivation of farming periods, and factors 3 plant senescence and factor 5 plant
495 detritus could be associated to harvesting of vegetation or crop. During the
496 observation period of a month, along with the weather warming as sunshine enhanced,
497 human left two obvious traces of cultivated soil during 9-17 March and 27 March-8
498 April and a trace of vegetation or crop harvest during 17-30 March. The stronger
499 pollen discharge occurred in March, probably due to the flowering of certain plants.
500 The BB emissions peaked on 9, 16 March, and 1 April were more prone to be open
501 burnings.

502 Since there is still some uncertainty of the factor apportionment, the proportion of
503 sources are only relative and uncertain. It is still difficult to distinguish the
504 contributions of microorganisms and plants to biogenic aerosols. Furthermore, all the
505 above speculations about farming and harvesting periods are based on only one
506 month's observation, and long-term observations are needed to obtain more accurate
507 and effective information.

508 **4 Conclusion**

509 With the help of the various atmospheric saccharides, this study presents the
510 characteristic of BB and various biogenic emissions to ambient aerosol in the rural
511 sites of Southwest China. Levoglucosan was the most dominant specie among all the
512 saccharides with the concentration of $287.7 \mu\text{g m}^{-3}$. The ratios of levoglucosan/OC
513 were 1.9-8.9% (mean: 3.7%). BB contributed to 19.1-73.9% of OC (mean: 41.3%).
514 The results indicated that domestic biomass fuel burning, open BB events, possibly
515 open agricultural waste burning, forest fire, or sacrificial activity appeared to be
516 significant during the spring in this area. The total concentrations of five mono (di)
517 saccharides and five sugar alcohols respectively contributed $24.9\pm 8.3\%$ and $26.6\pm 9.9\%$



518 of the total measured saccharides. Based on the regression analysis, these mono (di)
519 saccharides and sugar alcohols were mostly emitted by direct biogenic volatilization
520 from plant materials/surface soils, rather than BB processes. The sampling sites
521 suffered from both local emissions and BB via long-range transport from Southeast
522 Asia (Myanmar) and the northern Indian Peninsula. Five sources of saccharides were
523 resolved by NMF analysis, including BB (34%), soil microbiota (16.0%), plant
524 senescence (21.0%), airborne pollen (23.7%) and plant detritus (5.3%) at rural
525 Lincang in spring.

526 The data herein indicated that biofuel and open BB activities in the rural Southwest
527 China and neighboring regions could have a significant impact on ambient aerosol
528 levels. In addition to the residential biofuel usage, field burnings of agricultural
529 residues and fallen leaves, as well as forest fire, were non-negligible. Some new
530 technical measures of biomass resource utilization are urgently needed to improve the
531 open burning emission scenario in rural areas, along with strict prohibition policy of
532 BB. Meanwhile, the characteristics analysis of saccharides in the region can serve as a
533 valuable reference for future studies to evaluate temporal variations of biomass
534 combustion and biogenic emission during modeling predictions and policy making.

535 **ASSOCIATED CONTENT**

536 **Supporting Information**

537 Temporal variations of RH, temperature, solar irradiation and rainfall are shown in
538 Figure S1. Mean concentrations of saccharide compounds and the contribution of
539 them for the Datian, Dashu, and Yakoutian samples are shown in Figure S2. Daily
540 variation on average concentrations of levoglucosan and K^+ (a), arabitol and mannitol
541 (b), $PM_{2.5}$, Ca^{2+} and trehalose (c) at the three sites throughout the sampling period are
542 shown in Figure S3. Figure S4 showed daily variations on proportion of the five
543 factors to the total saccharides in $PM_{2.5}$ sampled at three sites during the sampling
544 periods. Table S1 lists the concentrations of the carbonaceous components and soluble
545 inorganic ions in $PM_{2.5}$ during the sampling periods of spring 2019. Correlation
546 matrix for the dataset of the determined saccharides compounds in $PM_{2.5}$ samples is
547 shown in Table S3.

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Captions of Figure and Table

Figure 1. Temporal variations of OC, EC, PM_{2.5} and total sugars at the three sites during the sampling periods.

Figure 2. The absolute concentration (bar chart) and the relative contribution (pie chart) of various saccharide compounds during the sampling periods.

Figure 3. Regression analyses of levoglucosan versus the other two anhydrosugars (a), K⁺ versus three anhydrosugars (b), levoglucosan versus PM_{2.5}, OC and EC (c), and K⁺ versus PM_{2.5}, OC and EC (d).

Figure 4. Spatial distribution of the fire spots observed by MODIS, as well as the corresponding 72 h backward air-mass trajectory clusters arriving at 1500 m above ground level during the sampling periods for the collected samples. The backward trajectories were separated into two episodes of remote western source over 2000 meters and local western source below 2000 meters, as shown in wine red and green lines.

Figure 5. Mean concentrations and contribution of saccharide compounds for the aerosol samples separated as over and below 2000 meters.

Figure 6. Factor profile obtained by NMF analysis (a). Source contribution of the five factors to the total saccharides in PM_{2.5} samples (b).



Figure 1

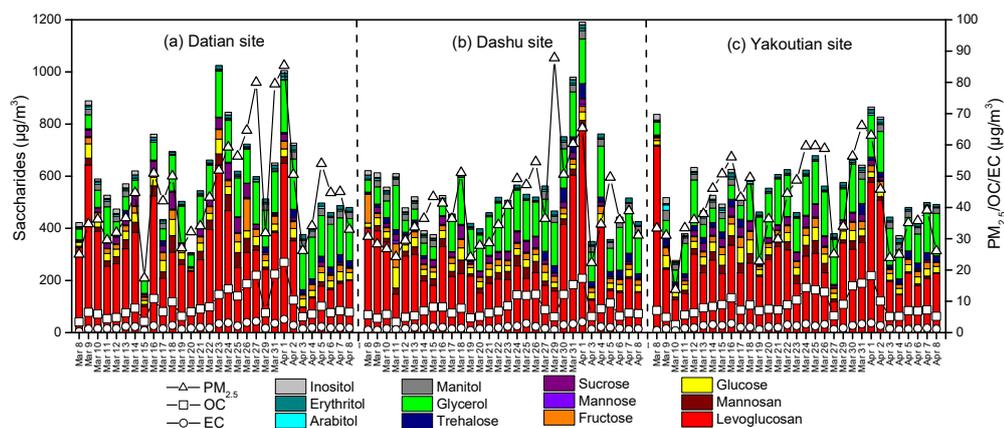




Figure 2

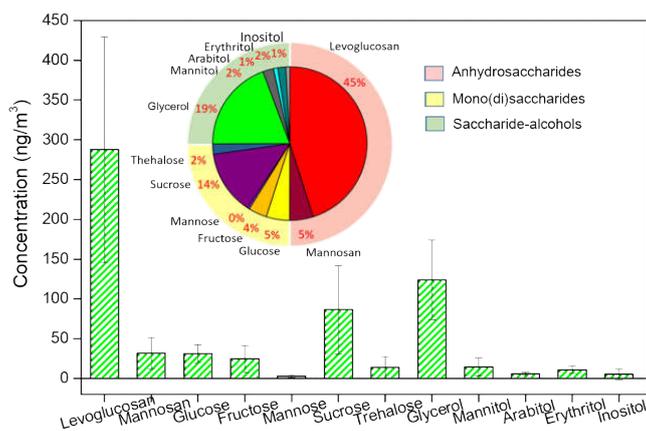




Figure 3

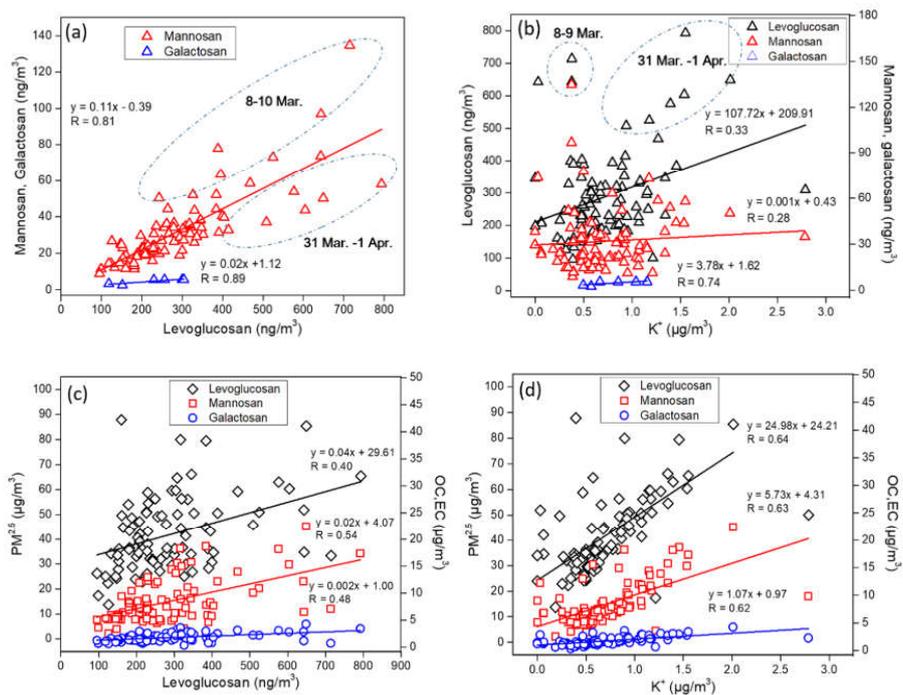




Figure 4

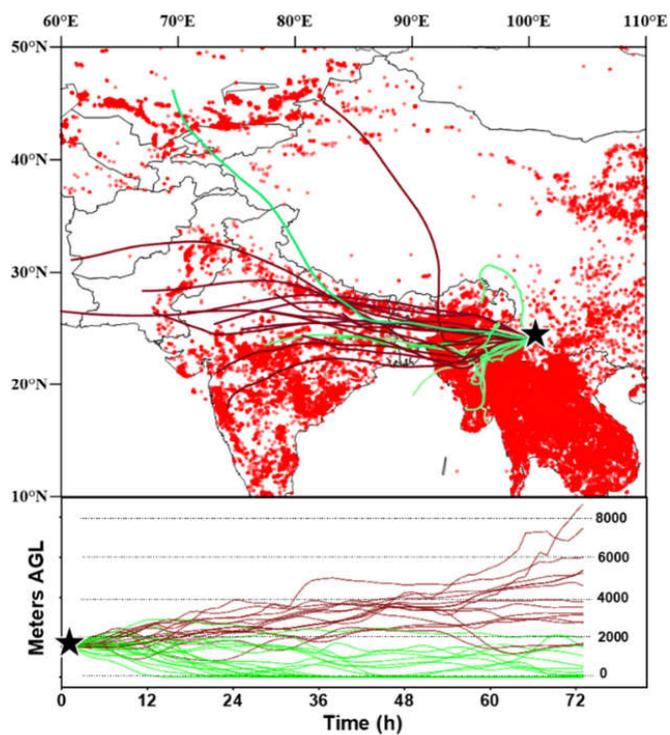




Figure 5

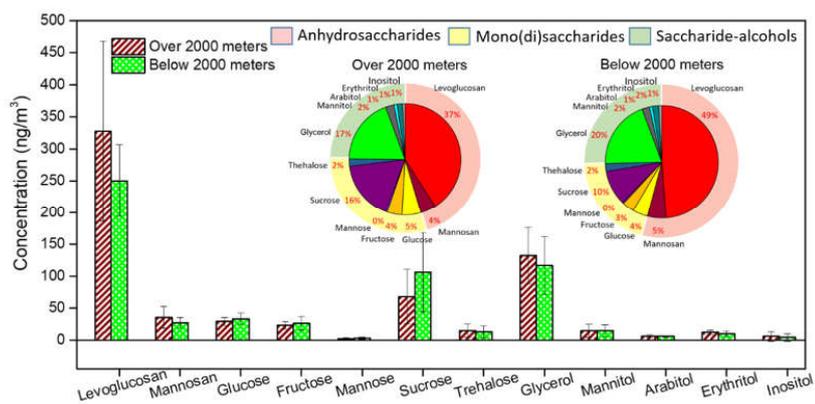




Figure 6

