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1 **The first balloon-borne sample analysis of atmospheric carbonaceous components reveals new**
2 **insights into formation processes**

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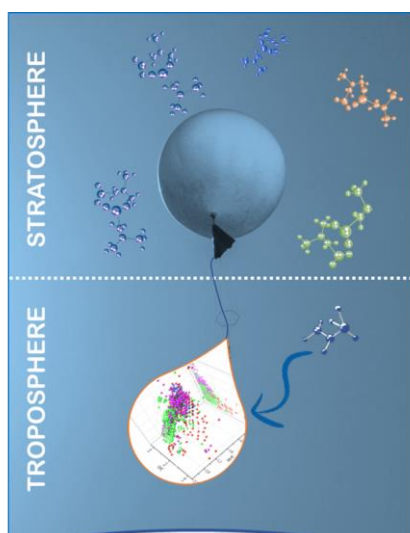
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19
20 **Abstract**

21
22 Atmospheric aerosol optical, physical, and chemical properties play a fundamental role in the Earth's climate
23 system. A better understanding of the processes involved in their formation, evolution, and interaction with
24 radiation and the water cycle is critical. We report the analysis of atmospheric molecules/particles collected
25 with a new sampling system that can fly under regular weather balloons for the first time. The flight took
26 place on 18/02/2022 from Reims (France). The samples were subsequently analyzed by high-resolution mass
27 spectrometry (Orbitrap) to specifically derive hundreds of organic components present in 4 different layers
28 from the troposphere to the stratosphere (up to 20 km). Additional measurements of O₃, CO, and aerosol
29 concentrations a few hours before this flight took place to contextualize the sampling. After separating
30 common species found on each filter that might be common to atmospheric layers or residuals for
31 contaminations, we found that each sample yields significant differences in the number and size of organic
32 species detected that must reflect the unique composition of atmospheric layers. While tropospheric samples
33 yield significantly oxidized and saturated components, with carbon numbers below 30 that might be
34 explained by complex organics chemistry from local and distant source emissions, the upper tropospheric
35 and stratospheric samples were associated with increased carbon numbers (C>30), with a significantly
36 reduced unsaturation number for the stratosphere, that might be induced by strong UV radiations. The
37 multimodal distributions of carbon numbers in chemical formulas observed between 15-20 km suggest that
38 oligomerization and growth of organic molecules may take place in aged air masses of tropical origin that
39 are known to carry organic compounds even several km above the tropopause where their lifetime
40 significantly increases. Overall, these results are consistent with the injection of fire smoke months before
41 the insitu observations and with thermodynamics inherent to conditions prevailing in the stratosphere.

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44 **Key words: DESI, Orbitrap, organic aerosols, troposphere, stratosphere**

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50 **Graphical abstract:**



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Highlights:

- A new experiment setup was designed and used to chemically characterize organic matter present in the atmosphere (troposphere to the stratosphere)
- We found that each sample collected in different atmospheric layers yields significant differences in terms of the number and size of organic species detected that must reflect the unique composition of atmospheric layers.
- The multimodal distributions of carbon numbers in chemical formulas observed between 15-20 km suggest that oligomerization and growth of organic molecules may take place in aged air masses of tropical origin that are known to carry organic compounds even several km above the tropopause.
- The present results are consistent with the injection of fire smoke months before the in situ observations and with thermodynamics inherent to conditions prevailing in the stratosphere

1. Introduction

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Particles in the atmosphere arise from natural sources, such as dust, sea spray, wildfires, and volcanoes, and from anthropogenic activities, such as the combustion of fuels. Primary aerosol particles are emitted directly into the atmosphere and secondary aerosol particles are formed by gas-to-particle conversion processes. They evolve in size and composition through condensation/evaporation of gases, by coagulating with other particles, by chemical reaction, or by activation to become nuclei for liquid or ice clouds formation

Aerosol particles play an important role in the radiative balance of the Earth, with cooling and warming effects depending on particle optical properties according to their size distribution and composition [1]. The export of aerosols and of their precursors from their source areas is associated with atmospheric chemical processes that affect air quality [2] and deplete the stratospheric ozone layer [3]. The aerosol effects depend on the altitude levels but overall are estimated to provide a net negative radiative forcing associated with a cooling of the surface [4]. However, uncertainties in the estimation of aerosol impacts are still large because of the scarcity of observations and the difficulty to characterize the complex aerosol microphysical and chemical properties with respect to altitude.

Tropospheric aerosols contain major fractions of organics (up to 90 %) [5-7]. The oxidation of volatile organic compounds results in the formation of secondary organic aerosols (SOA). A number of studies have been dedicated to the chemical characterization of tropospheric organic aerosol (OA) particles and to the investigation of the physical and chemical pathways for the formation of OA. In particular, reactions of carbonyl-containing organic species, including organosulfate formation and oligomerization reactions have been reported as potential sources of low-volatility organic products in OA [8-10]. Photochemical aging of the light-absorbing OA by photolysis and oxidation by ozone and OH is a significant sink of OA[11].

89 In the tropics, convective upwelling transports aerosol precursor gases and primary particles from the
90 lower to the upper troposphere [12] where homogeneous nucleation and particle growth processes form
91 secondary particles. Once in the upper troposphere/lower stratosphere (UTLS) region, air masses sustain
92 large-scale transport via the Brewer-Dobson (BD) circulation [13], first upward within the tropical
93 stratosphere and subsequently poleward to descend finally through the lowermost stratosphere into the
94 extratropical troposphere. In addition, direct two-way mass exchange between the upper troposphere (UT)
95 and lowermost stratosphere (LMS), through the extratropical tropopause, occurs along isentropic surfaces.
96 In the stratosphere, the dominant aerosol population consists of droplets of sulfuric acid and water, formed
97 via gas to particle conversion of mainly carbonyl sulfide (OCS) and sulfur dioxide (SO₂) [14]. OCS photo-
98 oxidizes at high altitudes, within the deep BD branch, while SO₂ oxidizes at all altitudes in the stratosphere
99 [15]. In the stratosphere particles are largely submicrometric and their sedimentation velocities are low
100 resulting in aerosol residence times of months [16]. Volcanic eruptions represent the strongest source of
101 variability in stratospheric aerosol as they can inject substantial amounts of SO₂ directly into the UTLS as
102 well as ash particles for some events [17, 18].

103 In addition to the sulphate population, aerosols in the UTLS contains fractions of other particle types, i.e.
104 organic particles, a smaller proportion of black carbon and some extraterrestrial material as derived from in
105 situ airborne observations using mass spectrometry [19, 20], single-particle soot photometry [21] or aerosol
106 sample collection/impaction with subsequent composition analysis in the laboratory [22, 23]. The increasing
107 occurrence of wildfires in the warming climate is an important source of injection of black carbons and
108 organic particles in the UTLS [24, 25]. Together with sulphates and organics, the presence of nitrates,
109 ammonium and mineral dust has been reported in the UTLS above the polluted Asian region during summer
110 monsoon by model simulations [26, 27] with possible subsequent influence on the northern hemisphere
111 aerosol content [28]. However, recent aircraft and balloon chemical composition measurements indicate the
112 dominance of secondary aerosols such as nitrate, organics and sulfate with a degree of oxidation increasing
113 with altitudes [29, 30]. It remains uncertain if these particles are in an internal or external mixture with the
114 sulfuric acid droplets though [19] suggest rather internal mixtures of sulfate and carbonaceous material.
115 During their transport, their composition of organic aerosols is transformed by oxidation, cloud processing
116 and wet/dry deposition [31-33]. However, the detailed processes involved in the formation of organic
117 aerosols, the chemical composition of the organic fraction and its potential impacts on the chemical and
118 optical properties of UTLS aerosols are still unclear. Most of the observations of OA have been obtained in
119 the troposphere (or in laboratory experiments excluding thermodynamical and chemical conditions of the
120 UTLS). Chemical composition, air mass residence times and UV flux between the lower troposphere and the
121 UTLS are too different to apply directly the results obtained for tropospheric OA. Available sampling
122 techniques in the UTLS are limited due to the restricted number of research platforms and instruments
123 capable of sampling at these altitudes. Therefore, it is particularly important to improve the knowledge about
124 the organic aerosols in the UTLS in order to identify the sources and processes leading to aerosol presence
125 and production there.

126 In this study, we analyze gas and aerosol samples collected on balloon-borne filters from the lower
127 troposphere to the stratosphere at mid-latitudes for the first time, to our knowledge, with altitude segregation
128 and subsequent information on the chemical organic composition derived from high resolution mass
129 spectrometry (Orbitrap, Q-Exactive, Thermo Scientific). The major aim of this study is to present the new
130 method and gain insight into the differences between the troposphere and the stratosphere in term of organic
131 chemicals composition. Chemical analysis may help to better understand the sources and processes
132 controlling the composition of the UTLS region.

135 **2. Experimental**

137 **2.1 Materials and methods**

139 **2.1.1 Sampler description**

140 The CHEM Filter Sampler (referred as Sampler herein below) is a system that collects particles on several
141 filters for time-resolved sampling[34]. The system was adapted for balloon flight applications using a micro-
142 controller connected via serial communication and an InterMet radiosonde that collects weather parameters
143 (Pressure, Temperature, Relative Humidity) as well as Global Position System locations (Lon, Lat, Alt). The
144

145 sampler is composed of eight filter cartridges that hold the individual filters (Fig S1-a). The eight channels
146 allow to perform time-resolved and/or altitude-based sampling. An external rotating disk with permanent
147 magnets interacts with internal valve magnets so that only one valve opens at a given time depending on
148 altitudes. The rotating disk is located downstream from the filters. The sampled flow rate is measured by a
149 laminar flow element on the inlet (Fig S1-b). The sampled flow is controlled by modulating the speed of the
150 integrated vacuum pump. The sampler is controlled based on the pressure measured by the Imet radiosonde.
151 Once started, the unit automatically exposes each filter to the sampled flow within a given atmospheric layer
152 between a lower and an upper pressure bounds. During position change, the pump is turned off. The aerosol
153 sampler was cleaned thoroughly with isopropyl alcohol before each flight.
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155 **2.1.2 Sampler preparation**

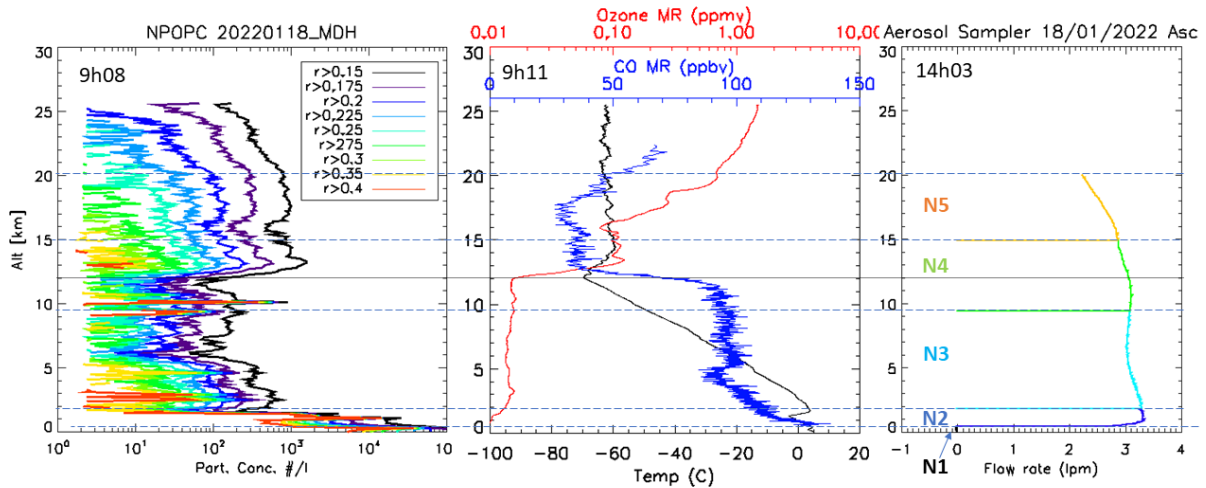
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157 Activated carbon cloth (ACC) referenced CT27 (Mast CARBON UK) prepared from a viscose precursor was
158 used as an adsorbent to collect the volatile organic compounds (VOCs) and organic aerosols (OAs) present
159 in the atmosphere[35, 36]. The nanotexture, porosity and chemical functionalities of CT27 have been fully
160 characterized in a previous work[37]. These activated carbon materials have a highly developed specific
161 surface and are used in air pollution control applications. Their effectiveness is recognized for the filtration
162 of VOCs and the trapping of molecules. In the form of fabrics, these materials are self-standing filters easy
163 to handle and present the advantage of not releasing any fine particles and dusts. The fabrics used are
164 composed of carbon fibers with an average diameter of 1 μm with a range of porosity covering several orders
165 of magnitude (micropores < 2 nm, mesopores between 2 and 50 nm, and macropores with diameter > 50nm).
166 The porosity is directly accessible all along the fibers allowing a quick diffusion of molecules within the
167 porosity followed by physical or chemical adsorption on adsorption sites and capillary condensation in the
168 micropores. These fibers were selected for their hydrophilic properties originated from a high oxygen content
169 3.7 wt % determined by temperature programmed desorption analysis; the oxygen heteroatoms present in the
170 carbon structure being arranged in acidic functionalities. Their nitrogen adsorption isotherm is of type IV
171 showing an hysteresis characteristic of a micro/mesoporous material. The specific surface area was over 1100
172 m^2/g with a total pore volume of 0.75 cm^3/g , a micropore volume (pores between 0.7 to 2 nm) of 0.43 cm^3/g
173 (N_2 adsorption) and a mesopore volume (pores from 2 to 50 nm) of 0.45 cm^3 . CO_2 adsorption showed a high
174 ultramicropore volume (pores below 0.7 nm) of 0.45 cm^3 . The pore size distribution determined by applying
175 DFT model on nitrogen adsorption isotherm shows mainly two ranges of pore sizes, around 1.2 and 3.8 nm
176 able to trap small gaseous molecules or oligomers and aerosols respectively. Based on the properties of those
177 filters, we therefore assume that the system can collect gases and aerosols.

178 Two filters, 1.3 cm in diameter, of this fabric were cut out, superimposed and loaded in each filter cartridge
179 for aerosol and gas collections along the air flow (Fig 1-c). After the flight, filters were placed in individual
180 aluminum foil and named according to their respective positions. To limit chemical transformation, the
181 sample were stored in a desiccator which was swept more than ten times its volume with nitrogen (99.995 %
182 pure) to eliminate all traces of humidity and oxygen. For the flight data presented in this study, the sampler
183 was loaded at ~13h00 and unloaded after recovery near ~19h00. The chemical analyzes were carried out
184 three days after the atmospheric sampling.
185
186

187 **2.1.3 Balloon flights (18/01/2022)**

188
189 On 18 January 2022, 3 balloon flights took place from Reims (France) as part of the Radiosondage des
190 Evenements extremes et des Aerosols de la Stratosphere (REAS) campaign (Dumelie et al., 2022, in prep).
191 The first flight was launched at 9h08 and included a Particle plus Optical Counter (POPC) for which the
192 concentration profiles are shown in Fig. 2. The second flight launched a few minutes later embarked an M20
193 radiosonde, an electrochemical Concentration Cell and an Aircore sampling system. Vertical profiles of
194 temperature, O_3 , and CO mixing ratio are shown in the middle panel of Fig. 2. Finally, a third flight took
195 place at 14h03 with the sampler. The flow rate profile (lpm) colored by position Id (N1 to N5) is shown (Fig.
196 2, right). The aerosol, trace gases and temperature profiles provide meteorological and chemical information
197 within the layers sampled during the flight. The cold point tropopause (~13km) marks a transition with a
198 positive gradient in aerosol concentration (increase by a factor 10) for all sizes as well as ozone concentration
199 while CO concentration decreases by more than 50%. The transition region between the troposphere and the
200 stratosphere also known as the UTLS region was sampled on position 4 between 9.4-14.9 km. In the free
201 troposphere where aerosol concentration for $r > 0.15 \mu\text{m}$ is lower than in the stratosphere was sampled on

202 position 3. Above the cold point tropopause near 12 km, two distinct regions of the stratosphere exist.
 203 Between 12-16 km, higher concentration of ozone is collocated with increased aerosol concentration for
 204 larger size ($>0.4\mu\text{m}$) while the layer above 16 km is marked by an increase of ozone but with smaller aerosols
 205 concentration. The region between 12-16 km corresponds to the lowermost stratosphere, an active zone of
 206 exchanges with the tropical and mid-latitude troposphere while the lower stratosphere above 16 km is
 207 generally influenced by the horizontal and downward motions of the Brewer Dobson circulation [23]. We
 208 also note that CO displays behavior changes within the two layers. The upper layer was sampled on position
 209 5 of the sampler and was likely influenced by air masses transported from the tropics.



210
 211 **Fig. 2.** (Left panel). Particle concentration for 9 size radii between 0.15 μm and 0.4 μm measured by the
 212 Particle plus Optical Counter (POPC) on a balloon flight on 18/01/2022 at 9h08 (launch time) from Reims.
 213 (middle) Corresponding temperature, ozone and CO profiles measured by an M20 radiosonde, an
 214 Electrochemical Concentration Cell (ECC) and Aircore on a second flight which took place a few minutes
 215 after. We note that the Aircore retrieved CO was measured during the descent (right). Flow rate on different
 216 filter position (N1 to N5) during a third flight with the aerosol sampler launched at 14h03.

217
 218 The goal of the sampler flight was to collect gas and aerosol samples from distinct atmospheric layers for
 219 which differences were highlighted from the POPC observations (Fig.2). The derived layers and
 220 corresponding sampling characteristics are described in Table 1. We selected the levels of each layer using
 221 an aerosol profile derived from the Compact Optical Backscatter for Aerosol Detection (COBALD) the
 222 previous night (Fig.S2). The ozone and aerosol profiles indicate that those layers, in which N4 and N5 were
 223 sampled, have likely different origins. Higher ozone values would indicate a tropical origin also associated
 224 with volcanic aerosols from La Soufriere eruption which took place in April 2021, i.e., 9 months prior to the
 225 balloon flight. The transport of the plume is apparent from the observations of the CALIPSO spaceborne
 226 lidar (Fig. S4). At lower levels (9-15 km), the second peak might be associated with residual of smoke from
 227 biomass burning which took place during the summer and fall 2021. In the troposphere, the boundary between
 228 N2 and N3 was chosen to differentiate aerosols in the planetary boundary layer with a top located near 2 km
 229 (very sharp decrease of aerosol concentration) and the free troposphere above.

230
 231 **Table 1.** Summary of the filter position number, altitude range, samples time (ST), sampled volume (SV),
 232 temperature in Kelvin, atmospheric pressure and number of moles (n) during the balloon flight on 18
 233 January 2022.

Position # (filter No.)	Altitude range (km)	Pump status	ST (s)	SV (l)	T(K)	P atm	n
P1 (N1)	Ground-0.5	Off	11714	-3	-	1	-
P2 (N2)	0.5-1.8	On	562	30	273	0.86	1.1525
P3 (N3)	1.8-9.4	On	2659	134	233	0.38	2.665
P4 (N4)	9.4-14.9	On	1629	81	208	0.08	0.38
P5 (N5)	14.9-20	On	1525	66	213	0.02	0.0756

235
 236

2.1.3 Sample analysis

Samples collected on ACCs were analyzed by high-resolution mass spectrometry (Orbitrap Q-Exactive, Thermo Scientific). The IonMax HESI II source has been modified by soldering a gold foil onto the corona needle. The filters were placed on this new gold support using mounting brackets. We used desorption electrospray ionization (DESI), a soft ionization technique, to analyze the chemicals present on the filters in their native state [38, 39]. With this mode of ionization, the ionic intensities do not directly reflect the concentrations of the chemicals present on the filter. This method of analysis is extremely sensitive (low femtograms per squared millimeter) [40] but, the present data is therefore only of a qualitative nature. Figure S3 presented in the Supplements shows the assembly used in this work.

Analyses were performed by direct instillation of toluene (flow injection analyses - FIA) for chemical ionization A Heated electrospray ionization source (HESI) was used. The FIA settings for HESI ionization were: sheath gas 20 arbitrary units (a.u.), auxiliary gas flow 0, vaporizer temperature 120°C, capillary temperature 280°C, spray voltage 3 kV, flow injection of 6µL/min recorded for 1 min for data averaging. The Orbitrap® Q-Exactive from Thermo Scientific has a mass resolution of 140,000 and mass accuracy <0.5 ppm RMS. Mass calibrations in positive and negative HESI were performed using Pierce™ calibration mixtures (Thermo Scientific). The maximum ion injection time was set to 200 ms with an automatic gain control of 10⁶. For each sample, the faces exposed to the atmosphere, were analyzed using positive and negative ionization modes. All samples were analyzed three times three mass ranges (m/z 50-750, 200-1000, and 500-2000). The use of different mass ranges aims to correct for the decrease in trapping efficiency at higher masses. A more detailed description of these technical aspects is available in a recent review[41]. Using several mass ranges allows detecting simple molecules, oligomers but also nano aerosols of a few dozen nanometers in diameter. For larger aerosols, only the parts extracted by desorption with m/z < 2000 will be detected[42]. In mass spectrometry, m/z close to 2000 Da correspond to chemicals up to 0.02 µm in size. We observed a gradual growth of the size of VOCs towards that of aerosols (POPC). The DESI sensitivity which allows the measurement of a few thousandths of a mole, is suitable for the mole concentrations observed during this flight (Tab.1).

2.2 Data Processing

High resolution mass spectrometry generates a large amount of data that is easier to interpret with visualization tools [43-46]. In this study, we used double bond equivalent (DBE), van Krevelen diagrams, carbon oxidation state (OSc), and degree of unsaturation as a function of m/z for these representations. Molecules with carbon, hydrogen, oxygen, nitrogen, and sulfur atoms were sought. Molecular formula assignments were made with a maximum mass deviation of 3 ppm and an intensity greater than 10³. Molecular formula with an intensity lower than 5.0x10³ and detected only once out of three analyzes were not retained. The DBE represents the sum of the number of unsaturation (double bond and ring) present in a chemical compound [43]. Decreasing the number of hydrogen atoms increases its value. DBE is defined as:

$$DBE = 1 + c - h/2 + n/2$$

(c= number of carbon atoms; h = number of hydrogen atoms; n= number of nitrogen atoms)

for elemental composition C_cH_hN_nO_oS_s where only divalent sulfur is considered.

The van Krevelen diagram [47, 48] shows the evolution of the H/C ratio as a function of O/C for a set of identified chemical formulas. In a complex organic mixture, this type of representation allows the identification of classes of compounds such as aliphatics, aromatics or highly oxidized compounds.

The carbon oxidation state (OSc) measures the degree of oxidation of organic species (alcohols, aldehydes, carboxylic acids, esters, ethers and ketones, but not peroxides) [49-51]. It is defined by the following equation:

$$OSc \approx 2O/C - H/C$$

The percentage of unsaturation [52] was calculated by dividing the DBE number for each chemical formula identified by the number of single bonds in a saturated hydrocarbon having the same number of carbon atoms (CN). All molecular ratios, DBE numbers, and chemical formulas presented in this article refer to neutral molecules.

294 Unsaturation % = $DBE / (3 \cdot CN + 1) \cdot 100$
295

296 3. Results and discussion

297

298 3.1 Obtained main chemical formulas and classification

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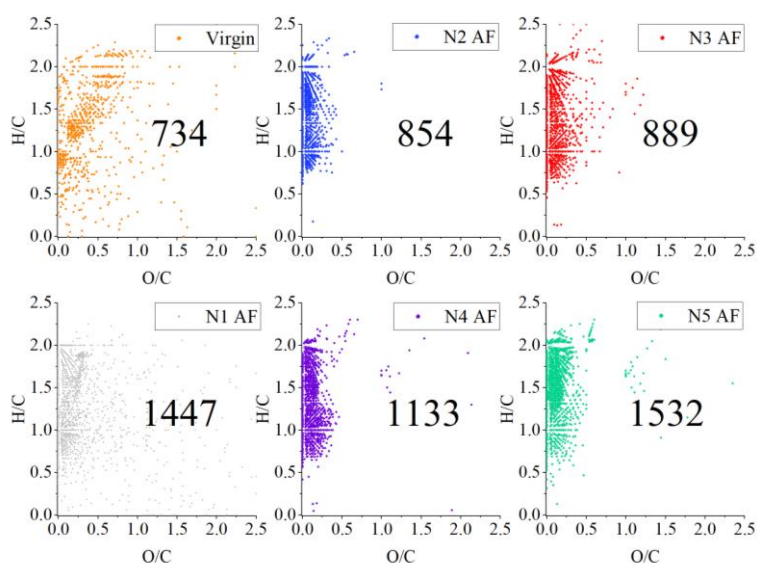
300 The data processing carried out on all six filters (Virgin + 1-5) gave six sets of chemical formulas. Given the
301 data filtering criteria presented previously, we searched for $C_cH_hN_nO_oS_s$ molecules in positive and negative
302 modes.

303 For the chemical formulas identified as containing sulfur, only the presence of H_2SO_4 could be verified on
304 the filters 1 to 5 (negative mode). This detection is expected since sulfur is a major gas emitted in the
305 troposphere from anthropogenic activities (in the form of SO_2) and that sulfuric acid droplets are ubiquitous
306 in the UTLS and the stratosphere via anthropogenic (SO_2 , DMS, OCS) and volcanic (SO_2) sources. The
307 stratosphere was likely influenced by La Soufriere eruption as seen by an increase of aerosol concentration
308 from the OPC (Fig.2) which likely led to reinforce the H_2SO_4 signature observed with the Orbitrap. For the
309 other chemical formulas, the isotopic analysis did not confirm their presence, either because of too low
310 intensity of ^{32}S (if ^{32}S is $< 10^5$ then $^{34}S < 10^3$), or by the absence of the mass containing the ^{34}S isotope. Its
311 detection on the filters is a first indication of the quality of the samples and the measurements by mass
312 spectrometry.

313 We also searched in filter 1 for potential contamination and identified a series of fluorinated compounds
314 characteristic of PolyTetraFluoroEthylene (PTFE). This contamination was also identified on filters 2 to 5
315 and coming from PTFE filters used during previous samplings. No other contamination could be identified.
316 Among all the other chemical formulas detected, we restricted the study to VOCs and OAs composed only
317 of carbon, hydrogen and oxygen which constitute the majority of VOCs and OAs present in the atmosphere,
318 i.e, hydrocarbons, oxygenated VOC, OAs, and polycyclic aromatic hydrocarbons, (PAHs)[53].

319 In the first part of this study, we compared the results obtained for the different faces of the filters for the
320 same sampling zone (face exposed to the atmosphere – AF; face facing each other– FF; and the face oriented
321 to the diaphragm pump - MF). An average reduction of 20% in intensities is observed and consequently a
322 reduction in the chemical formulas identified from AF to FF and FF to MF. Therefore, we used the data from
323 the faces exposed to the atmosphere for comparison.

324 Under these conditions, for the six filters, we identified a total of 6589 chemical formulas. In order to process
325 these large data sets, we used graphical tools previously described [43]. We used the van Krevelen
326 representation and gave for each filter the number of chemical formulas identified (Fig. 4). This allows
327 identifying families of chemical compounds [54] ; here it highlights differences in oxidation (O/C ratio) and
328 unsaturation (H/C) specific to the samples collected on each filter.
329



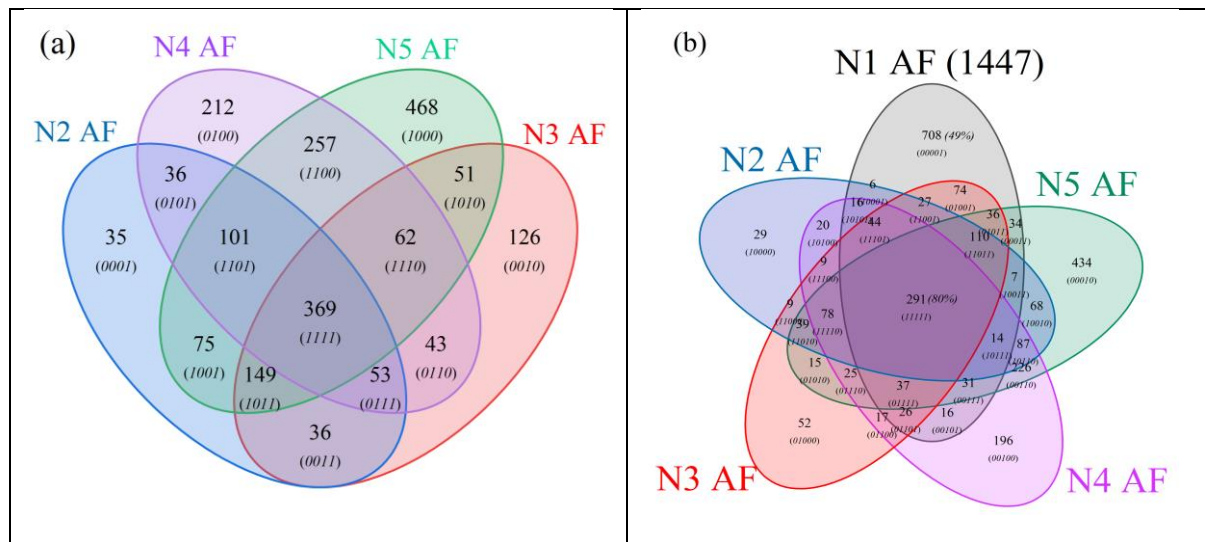
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332

Fig. 4. van Krevelen diagrams for the six filters

333 We correlated the six lists of $C_6H_8O_6$ chemical formulas identified on filters 2-5 to the physical parameters
 334 of the flight (atmospheric pressure and gas flow). It appears that the amount of gas flowing through the filters
 335 had little impact on the number of chemical formulas detected, unlike the decrease in pressure which, with
 336 altitude, enabled adsorption to be activated on the surfaces of the filters.
 337 In order to highlight the possible logical relationship between these filters, a Venn diagram was built (Fig.
 338 5).
 339



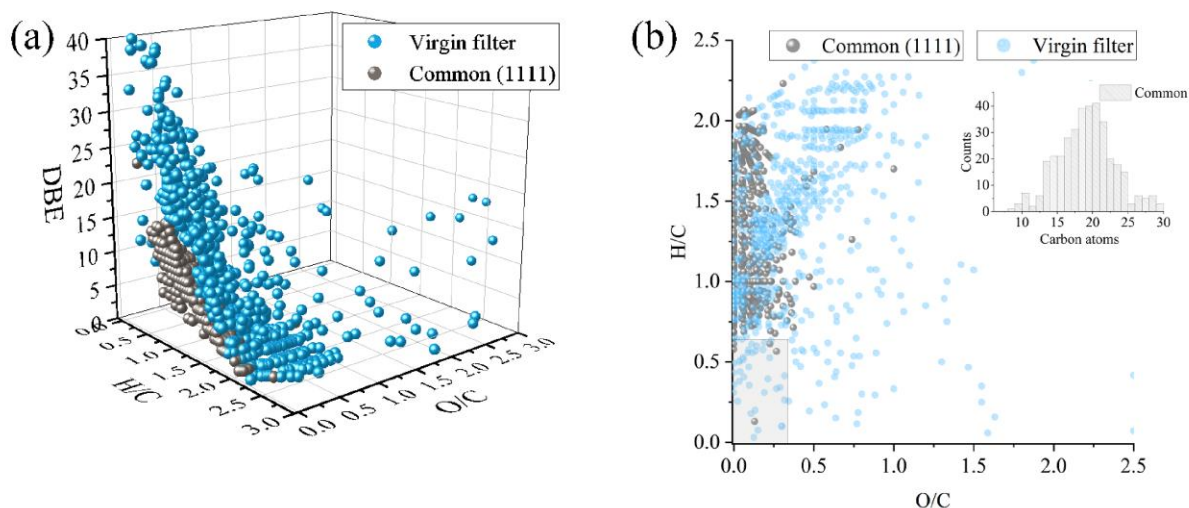
340
 341 **Fig. 5.** (a) Venn diagram showing common chemical formulas detected in samples 2-5 AF, (b) Venn
 342 diagram showing common chemical formulas detected in samples N2-N5 AF compared to N1.
 343

344 This diagram shows (Fig. 5a) 15 sets (0001-1111) which could reflect atmospheric similarities between the
 345 different sampling zones. We defined four groups of data according to the following criteria. The first group
 346 contains only the data of the set (1111) common to the four filters. These 369 chemical formulas represent
 347 30% of the total population identified for the four filters. Among the 369 common species, some might
 348 represent some levels of contamination but could also be indicative of common species throughout the
 349 atmosphere. The second group consist of sets common to two or three neighboring sampling areas. For
 350 example, these are the sets common to filters 2 and 3, or 3, 4 and 5. The third group consists of the four sets
 351 specific to a sampling zone: (0001), (0010), (0100), and (1000). Finally, the last group contains the other sets
 352 of which at least one of the sampling zones is distant, i.e., not contiguous (e.g., 2-4; 2-3-5). This last group
 353 represents only 20% of all chemical formulas. These distant groups that nevertheless share chemical formulas
 354 have not been studied. They may be the result of exchanges between the different layers. However, the
 355 absence of information on such exchanges has not allowed to confirm this hypothesis [25, 55]. The
 356 constitution of these four groups allowed us to make the new observations described in the next paragraphs.

357 358 **3.2 Origin of the group (1111) common to all four filters** 359

360 It is important to determine the origin of this group in order to know if these chemical formulas can
 361 correspond to those detected during this flight. This group can have different origins such as contamination
 362 by the media used, residual adsorption inherent to the exposure of these filters before and after the flight or
 363 chemical adsorption related to the flight. To verify the first hypothesis, we compared the data from reference
 364 filters, i.e., N1 et Virgin.

365 The N1 filter underwent the same theft as the other four filters. However, it was not subjected to a flow of
 366 gas, but simply to the pressure variations due to the theft. Adsorption was therefore reduced.
 367 For the Virgin filter, which was not flown, (no gas flow or pressure variation), the percentage of common
 368 chemical formulas with all four filters (2-5) is only 10%. Common chemical formulas set with space (1111)
 369 is reduced to 1%. In order to complete this comparison, we have represented the Virgin reference and the
 370 group (1111) using a 3D diagram including the DBE number in addition to a van Krevelen graph (Fig. 6).
 371



372
 373 **Fig. 6.** Comparison of the two groups (1111) and Virgin: (a) van Krevelen graph with DBE number, (b)
 374 without DBE number and with PAH and oxy-PAHs chemical fingerprint (grey area); inset plot: distribution
 375 of the carbon number (1111)
 376

377 This representation has the advantage (i) of avoiding the superposition of points linked to the van Krevelen
 378 representation and (ii) of providing new chemical information, i.e., unsaturation. The differences between
 379 these two groups are clearly visible in terms of the number of unsaturation and the O/C atomic ratios. We
 380 thus observe on the Virgin reference (Fig 6a), a larger number of chemical formulas with a DBE > 15
 381 compared to the group (1111). The chemical formulas with a DBE > 15 are mainly positioned in the gray
 382 zone of the Van Krevelen graph (Fig. 6b). This chemical character ($0 < H/C < 0.75$ and $O/C < 0.35$) corresponds
 383 to PAHs and oxy-PAHs. This consistent with the chemical composition of the graphite planes present in the
 384 filters used. The Virgin filter was not exposed to the pressure variations that occurred during the flight and
 385 that favored a uniform adsorption on the surfaces. Finally, the group (1111) does not present, with some
 386 exceptions, points in common with the gray area. We can then conclude that the existence of this group
 387 (1111) is indeed the result of the flight conditions, without excluding at this stage a contamination before
 388 and/or after the flight.

389 For residual adsorption inherent to the exposure of these filters before and after the flight (second hypothesis),
 390 one notes that the identified chemical formulas with dispersed low mass (86% in the range 200 to 340 m/z)
 391 and CN centered on 19 (Fig. 6b). Figure S5 shows the central position of this group (1111) among all
 392 identified chemical formulas (a) and those specific to each filter (b). Figure S5c shows the chemical formulas
 393 specific to each filter without the set (1111). Moreover, we note, that the percentage of unsaturation of the
 394 group (1111), in the range 5 to 27%, can result from a wide range of chemical reactions (addition,
 395 fragmentation, oligomerization, etc.). One notes that this set fits both in a chemical continuity (e.g. decrease
 396 of unsaturation with the increase of m/z), and on the other hand that in the absence of these data, no other
 397 chemical formula detected on another filter maintains this continuity. In conclusion, the graphs of Figure 6
 398 show a chemical evolution of the formulas indicating the presence of a common data set. It therefore seems
 399 unlikely, given these observations, that such large contaminations would fit so coherently into a remaining
 400 chemical set. Without excluding a punctual contamination, the set (1111) can be identified as a common set
 401 of chemical formulas present in the atmosphere.

402 It should be noted that apart from the sampling phases, the filters are isolated and only undergo pressure
 403 variations during the path of the balloon flight. This pressure variation induces adsorption without air flow.
 404 This passive adsorption can be observed principally on the N1 filter which has undergone the longest passive
 405 exposure, i.e., without pump generated flow. On N1, this passive adsorption is characterized by 51%
 406 similarity with the totality of the identified chemical formulas and 80% with the set (1111) present along the
 407 whole flight (Fig. 5(b)).
 408

409 3.3 Molecular formulas specific to sampling areas

410
 411 We compared the chemical formulas detected specifically in each sampling area: N2-AF (0001), N3-
 412 AF(0010), N4-AF (0100), and N5-AF(1000). The properties of these chemical formulas were plotted in a
 413 Van Krevelen diagram with an extra dimension assigned to unsaturation (DBE). In this type of representation,

414 the nature of the oxidation (formation of alcohols, carbonyls, unsaturation, etc.) is better described thanks to
 415 the inclusion of the DBE parameter (Fig.7-b). To complete these analyzes of the properties of chemical
 416 molecules, we have added for each filter the distribution of the carbon number.

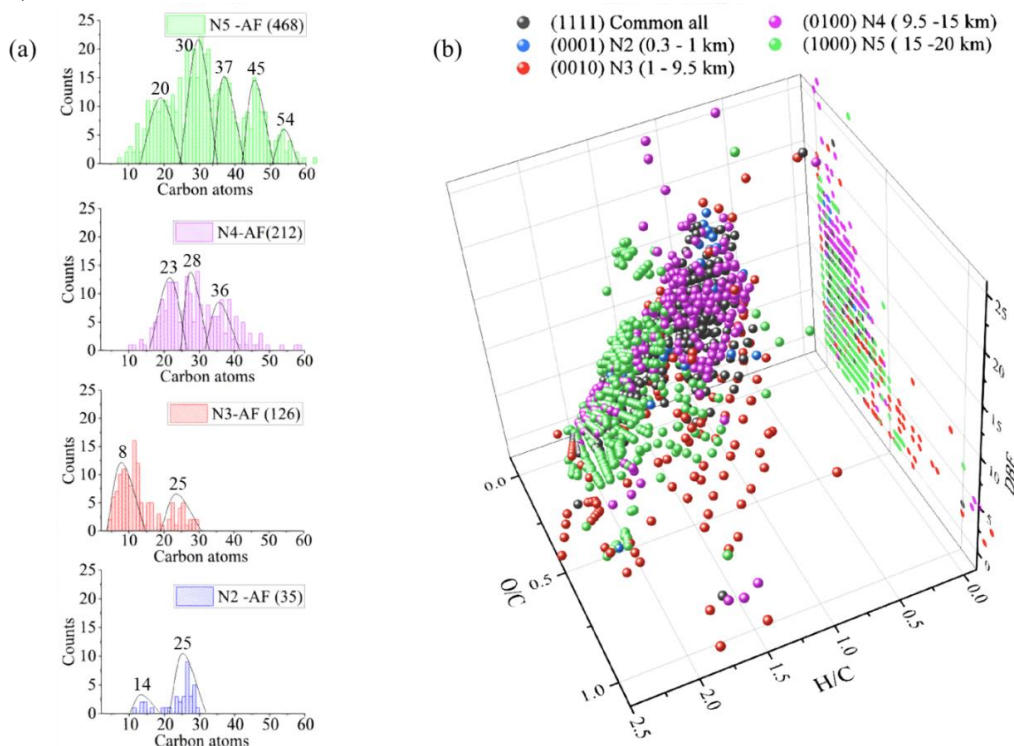
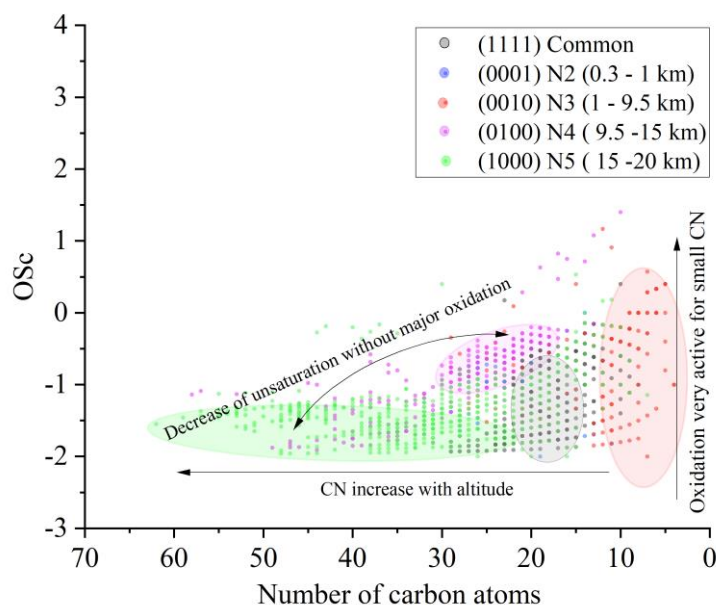


Fig. 7. (a) Carbon number distributions for filters 2 to 5; (b) Van Krevelen graph with DBE number.

417
 418
 419
 420 We observe on this graph a specific fingerprint of each sampling zone characterized by a centered dispersion
 421 of the chemical formulas, the oxidation level, and the amount of unsaturation. These results are correlated
 422 with the carbon number (Fig. 7a) and all of these data are particularly significant for filters 3, 4 and 5. For
 423 filter 2, the low number of chemical formulas specific to this filter and the dispersion of chemical formulas
 424 in this space limits the interpretation of the data. One should note that filter 2 was the one which was the less
 425 exposed to air flow between 0.5-1.8 km.
 426 For filter 3, one can observe a more important oxidation with a distribution centered on O/C = 0.5, unlike
 427 filters 4 and 5 showing a distribution centered on O/C=0.25 (Fig. 4). The distribution of carbon number is
 428 bimodal, but it is difficult to specify whether this observation is part of a process of fragmentation or growth.
 429 It is nevertheless observed that the DBE number increases with the length of the carbon number (CN = 10 to
 430 CN= 25).
 431 On filter 4, all the chemical formulas with distribution centered on O/C=0.25 and with an average DBE of
 432 12 show unsaturation not linked to the formation of ketones or aldehydes. Similar to filter 3, the growth of
 433 the molecules naturally increases this DBE number without significantly modifying the O/C ratio. This
 434 growth gives, here too, a multimodal distribution of the carbon number (CN = 23, 28 and 35) and could be
 435 assimilated to oligomerization.
 436 On filter 5, the distribution of chemical formulas is centered on O/C = 0.25, with a strong reduction of the
 437 DBE number (centered on 5), compared to filter 4. This reduction of unsaturation occurs in parallel with
 438 fragmentation /growth and a multimodal distribution of carbon number (CN = 20, 30, 37, 45, and 54). It also
 439 occurs in an environment where HO_x and O(¹D) radicals, abundant in the stratosphere, react with unsaturated
 440 species. The reduction in the DBE number could result from a breaking of double bonds after addition. At
 441 the same time, we observe a strong increase in the number of molecules with an O/C ratio close to 0.25 (2
 442 times higher than the distribution observed on filter 4), which corresponds to an increase in oxidation. (Fig
 443 S6). This decrease in unsaturation may explain this wide distribution of CN. More generally, if we observe
 444 an effect of atmospheric pressure on adsorption (i.e., the number of adsorbed chemicals), other variations,
 445 such as ozone concentration, concentration of oxygen atoms and ionizing radiation (UV) depending on
 446 altitude could affect products distribution. Thus, for filter 5, the relatively low pressure (leading to lower
 447 reaction kinetics) and the presence of UV radiations could explain the degradation of unsaturated chemicals
 448 and an increasing number of oxidized molecules. This type of oxidation would have no significant impact on

449 the O/C ratio. The OSc graph of Figure 8 offers another visualization of the results and highlights the
 450 particularities of each sample. There, we observe a clear oxidation in the low CN range of the N3 filter, much
 451 more visible than on filter 2. There is also a strong increase of the CN between the N4 and N5 filters with
 452 significant evolution of OSc. There, the group of chemical formulas (1111) is again integrated into a
 453 continuum between the different samples.

454 It is difficult to connect the evolution of chemicals' size with aerosols radii observed by POPC. However, our
 455 HRMS observations show that the size the evolution of chemicals' size follows the trend observed by POPC
 456 across the atmospheric layers.
 457



458 **Fig. 8.** OSc vs. number of carbon atoms presented for samples N2 to N5. The ellipses show the
 459 preferential zones of the different filters.
 460
 461

462 The data in general, and more particularly on filter 5, do not allow to specify if this increase in the number
 463 of carbon atoms is linked to an aging phenomenon.
 464

465 3.4 Flux between sampled neighbor areas

466 The sampling zones were defined based on aerosol concentration and backscatter properties from the
 467 troposphere to the stratosphere determined during a flight on the previous day. These boundaries are
 468 somewhat arbitrary and consecutive samples likely share chemical molecules through exchanges and
 469 transport. Table 2 shows how these exchanges evolve between the different sets. These exchanges are
 470 analyzed using the number of chemical formulas contained in the common sets. In addition to the number of
 471 common chemical formulas, we have expressed this value as a percentage with respect to each of the layers.
 472 There are many common species between these layers, greater than 50%, which nevertheless leave room for
 473 the particularities observed for each filter, i.e., (0001), (0010), (0100), and (1000).
 474
 475

476 **Table 2.** common species at the interfaces of the sampling areas
 477

Interface N2-N3	Interface N3-N4	Interface N4-N5
607 (N2-71%; N3-68%)	527 (N3-59%, N4-46%)	789 (N4-69%, N5-51%)

478
 479 **3.5 Link between distant areas sampled**
 480
 481 Fluxes of organic matter between atmospheric layers are poorly documented. Nevertheless, recent works
 482 have shown that the ejections of organic matter (e.g. during gigantic fires)[25, 56, 57] or mineral matter (e.g.

483 during volcanic eruptions) [58] could reach altitudes above 30 km. The distribution of these materials as a
 484 function of altitude and the resulting exchanges between atmospheric layers as well as the chemical evolution
 485 of carbonaceous matter along its transport, remains to be studied using chemistry-transport models. Our
 486 analyzes show common chemical formulas in non-consecutive layers (e.g., sets (0101), (1001), and (1010)).
 487 The number of these chemical formulas remains small compared to the total 2073 chemical formulas
 488 identified on the filters (2-5). The occasional presence of chemical formulas common to distant layers remain
 489 unexplained at this stage of the study, but although the number is small, it is large enough not to be considered
 490 as an experimental error.
 491

492 **4 Matching of the identified chemical formulas with the literature**

493 Stratospheric levels corresponding to filter 5 are shown to be impacted by air masses transported from the
 494 tropical lower stratosphere (Fig. S2) pointing at a tropical origin of the sources of the detected carbonaceous
 495 molecules. O/C ratios in the UTLS have been occasionally reported in the literature and have shown values
 496 of 0.18-0.2 [59][23] matching our results obtained for filters 4 and 5. This organic signature is expected to
 497 stem from organic volatiles and primary aerosols from the troposphere [23], mostly from the tropical latitudes
 498 [19, 20]. However, our method focusses on heavy molecules whereas reported studies may integrate a wider
 499 spectrum of organic species and also much large aerosols. Moreover, one should keep in mind that O/C
 500 values could depend on the location and season corresponding to various influences of overlying stratospheric
 501 layers as controlled by the BD circulation. Wildfires with subsequent injection to stratospheric levels are
 502 expected to alter the O/C ratio in this region. However, our values for UTLS levels corresponding to filter 4,
 503 possibly impacted by past fire events, appear well below 0.5-1, typically reported for young biomass burning
 504 plumes [60, 61]. The measured O/C here could be linked to chemical aging processes all along smoke
 505 transport.
 506

507 The small quantities of materials adsorbed on the filters (2-4) did not allow the use of High Performance
 508 Liquid Chromatography (HPLC) for a possible identification of the chemical compounds present.
 509 Nevertheless, in the hypothesis of an origin linked to injections from wildfires in Northern America, regularly
 510 observed in summer and fall over the past few years, we studied the similarities with chemical composition
 511 of biomass burning samples presented in the literature. This comparison remains partial, because it was
 512 carried out using chemical formulas and does not consider the phenomena of aging or dilution. Nevertheless,
 513 it constitutes a necessary step for a better understanding of the evolution of biomass burning plumes in the
 514 atmosphere. For this screening, we selected three studies dealing with biomass burning and containing
 515 chemical compounds data. Details on these studies are given in Table 3.
 516

517
 518
 519 **Table 3.** References and characteristics of biomass burning studies with percentage of similarity with
 520 identified chemical formulas in our study.
 521

Bibliographic reference	Samples	Fuels	Instrument (see supplementary)	Chemical formulae	Percentage similarity
Smith et al. [52]	Laboratory	Biofuel	LTQ-Orbitrap	73	60%
Gilman et al. [62]	Laboratory	Biofuels	GC, PTR, PIT, CIMS, FTIR	65	36%
Hatch et al. [63]	Laboratory Prescribed fires	Biofuels	GC-ToF	15	0%

522
 523 For each study, a Venn diagram was built and presented in the Supplements (Figures S7-S9). For the three
 524 studies considered, only the chemical formulas were considered. For example, the study by Hatch et al.
 525 reported 135 chemical compounds with numerous isomers which, in the end gave, 15 chemical formulas. In
 526 this case, the lack of similarity with the filter samples can be explained by this low sampling targeted on
 527 terpenes with a very low dispersion in number of carbons (mainly between 9 and 12 carbon atoms). For the
 528 other two studies, the similarities are significant and support the hypothesis that biomass burning products
 529 could have been collected on filters in the present study. The data from these two studies also indicate the
 530 existence of a common set of chemical formulas observed at different altitudes. We have 22% and 9% of the
 531 chemical formulas identical to the set (1111) for the work of Smith et al. and that of Gilman et al.,
 532 respectively. Levoglucosan, considered to be the most important and stable marker of biomass burning [64]

533 is present in the study of. Smith et al. However, due to its estimated time life of 8 days in the atmosphere, it
534 could not be observed in our samples, as could be expected.
535

536 **5 Conclusion and perspectives**

537
538 The results of several flights are presented. A range of instruments and techniques were used. A Particle plus
539 Optical Counter, an M20 radiosonde, an electrochemical Concentration Cell, an Aircore sampling system,
540 and a new sampler equipped with carbon filters were used. We collected atmospheric samples on carbon
541 filters through 4 atmospheric layers between 0.8 km to 20 km to separate different source regions. They were
542 analyzed by DESI coupled to high-resolution mass spectrometry.

543 The samples show 30% common chemical formulas. However, a non-negligible, proportion of chemical
544 formulas are only present on a specific sample (e.g. 10% for the free troposphere on N3 and 35% for the
545 stratosphere on N5) reflecting different sources, chemical processes or/and chemical evolution of the organic
546 matter from the troposphere to the lower stratosphere.
547

548 The main results of the chemical analysis show:

- 549
- 550 i) an increase of the carbon chains from the troposphere to the stratosphere
- 551 ii) In the stratosphere, the samples collected show a reduction in unsaturation compared to those
552 collected below (UTLS, sample 4)
- 553 iii) the oxidation of organic compounds measured by the O/C ratio remains stable across the layers
554 corresponding to the UTLS and the stratosphere
- 555 iv) The number of oxidized compounds increase with altitude
556

557 Above 15 km, unsaturated organic compounds are strongly influenced by UV light absorption [Ref], which
558 causes unsaturation breakdown and may also explain the multimodal carbon distribution associated with
559 molecular fragmentation/growth. Despite the increase of ozone concentration, at this altitude, the oxidation
560 of organic compounds measured by the O/C ratio does not increase unlike the number of oxidized
561 compounds, which increases. The fragmentation, possible outcome of the decrease of unsaturation, can
562 explain the observed simultaneous phenomena.
563

564 While quasi-coincident aerosol number concentration and size measurements collected with an optical
565 particle counter might be related to the chemical analysis shown here, the sizes of particle measured, a few
566 nanometers with the sampler and more than 300 nm with the optical particle counter make the comparison
567 difficult. Nevertheless, it is not unrealistic to assume that the source regions have produced specific
568 aerosols/molecules that were encountered by both instruments but the connections between both remain
569 challenging to assess.
570

571 Ozone and aerosol profiles obtained a few hours before the sampler flight shows that sample 5 was likely
572 influenced by air masses from the tropics associated with enhanced O₃ and aerosols likely influenced by La
573 Soufriere eruption which took place 9 months prior to the flights. At such levels, satellite observations point
574 at air mass origins from tropical latitudes where the higher quantities of organic material have been reported
575 in the literature [19, 20]. It is expected that carbonaceous compounds can remain for months in the
576 stratospheric aerosol particles [19].

577 The origin of the organic compounds we have detected is still to be determined especially in the UTLS and
578 the stratosphere (e.g. Asian pollution transported upward by cloud convection and then distributed to the
579 northern hemisphere, seasonal fires in South America and Africa). The way combustion products emitted
580 from anthropogenic emissions and natural wildfires evolve, sustain breakdown, results in unsaturated
581 organics and organic radicals, recombine to result in a spectrum of products under varying pressure,
582 temperature, humidity and solar flux all along their transport from the source regions is not clear.
583

584 At this stage, although efforts have been made to compute chemistry of secondary organic aerosols and
585 particulate organic matter from the surface to the stratosphere [27], no chemistry-transport model is able to
586 comprehensively simulate the various sources, content and chemical evolution of the organic matter for the
587 heavy molecules we have pointed out in our study.
588

589 An important question is whether or not our results are reproducible and correspond to a snapshot of the

590 atmosphere state or point at a steady feature. Further investigations at other seasons and latitudes, in period
591 non-impacted by volcanic and fire plumes would be necessary to address this question.

592

593 **Author statements**

594

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606 acquisition; Supervision.

607

608 **Declaration of competing interest**

609

610 The authors declare that they have no known competing financial interests or personal relationships that
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612

613

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625

626 **Acronyms:**

627

628 LTQ-Orbitrap: linear trap quadrupole Orbitrap; GC: gas chromatography; PTR: Proton-transfer-reaction;
629 PIT: Proton transfer ion; CIMS: Chemical Ionization Mass Spectroscopy; FTIR: Fourier-transform infrared
630 spectroscopy; GC-ToF: Gas chromatography coupled to time-of-flight mass spectrometry

631

632

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