



Supplement of

Variability in black carbon mass concentration in surface snow at Svalbard

Michele Bertò et al.

Correspondence to: Andrea Spolaor (andrea.spolaor@cnr.it) and Michele Bertò (michele.bereto@gmail.it)

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1 **1. Calibration of the SP2**

2 The empirical calibration of the SP2, performed using size selected fullerene soot particles is linear up
3 to 500 nm, and the assumption is that it is also linear beyond that size (and the corresponding mass).
4 However, when a massive particle enters the laser beam the incandescence signal might saturate the
5 detector; therefore, in this analysis only the particles below 700 nm were considered. The evaluation
6 of the BC mass for the samples showing a BC geometric mean mass equivalent diameter above
7 300/400 nm might therefore be more influenced resulting in underestimation of the mass. However,
8 the evaluation of the missing mass is beyond the scope of this manuscript and require further analyses.
9 The losses of mass due to the presence of undetected small particles, below 80/70 nm of MED, is not
10 significant given that the geometric mean of the MED of the mass size distributions is always above
11 150 nm.

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13 **2. Possible interferences during the SP2 rBC mass concentration in Arctic snow**

14 The surface snow of the Svalbard archipelago is normally characterized by high content of sea salt
15 particles, for example those containing Na, due to its geographical position surrounded by the ocean.
16 The Na concentration in the samples analyzed in this work is, on average, about 500 ng g⁻¹. We
17 cannot exclude that the rBC mass concentration variability for the 85-days experiment presented in
18 this work might have been slightly influenced by the Na content (tracer of sea salt deposition),
19 however no clear relation appeared from the comparison between the rBC and the Na mass
20 concentrations during the “85 days” experiment (Fig. SI 3). As a precaution, all the samples from the
21 3-days experiments were diluted five times with Milli-Q water prior to the SP2 analyses. The SP2
22 instrumental performances during the analyses, in terms of laser power and incandescence signal
23 quality, were monitored and constant. At the best of our knowledge, no study focusing on the possible
24 effects of the Na particles on the rBC mass concentration retrieved from the SP2 has been published
25 in the literature. Further analyses of snow samples collected in areas characterized by a high influence
26 of marine-like aerosols could shed light on the effects of salt particles on the rBC mass and number
27 concentration via SP2 measurements, as well as establishing a shared procedure to avoid measuring
28 artifacts.

29 Mineral dust particles might have an influence on the SP2 measurements, depending on their chemical
30 and physical properties. Currently, the literature lacks investigation on these specific SP2-mineral dust
31 particles interactions. However, a few studies on snow and ice samples measured with an SP2 take the
32 assumptions that mineral dust particles are not detected by the SP2 as incandescence signals, but only
33 as scattering signals (Kaspari et al., 2011). Recent studies, based on atmospheric measurements
34 suggest that the SP2 rBC mass concentration measurement can potentially be interfered by the
35 presence of metals and metals-oxide (Moteki et al., 2017), of volcanic ashes or of dust (Kupiszewski
36 et al., 2016).

37 **3. Ancillary measurements for the 3-days samples: Na, Mn, EC, OC**

38 Ancillary data were measured/gathered for the 3-days experiment in order to strengthening
39 the interpretations even though they were not considered in the statistical exercise. The Mn
40 concentrations are considered to be a good proxy for mineral dust aerosols (Baker et al., 2006). The
41 Mn concentration profile was compared with that of the coarse mode particles concentration showing
42 a very similar behavior and suggesting a common source. A similar pattern is also clearly visible for
43 all the various chemical species.

44 Every six hours a surface snow sample was collected in parallel to that of the hourly
45 sampling. The melt snow volume of these samples were of about $1618 \pm 290 \text{ mL cm}^3$ and they were
46 used for TC (Total Carbon), EC (Elemental Carbon) and OC (Organic Carbon) measurements. Results
47 are shown in Fig. SI 4. A different profile is shown for the three compounds compared to that of the
48 rBC mass concentration. As reported above, the two measuring techniques are different and, in
49 particular, the size range of particles detected by the two instruments is different, from 80 to 600 nm
50 for the SP2 whereas a much broader dimensional spectrum for the Sunset, potentially explaining part
51 of the observed difference. Interestingly, the EC daily values increased of one order of magnitude,
52 from 1 to $10 \mu\text{g l}^{-1}$, during the sampling period, showing a maximum during the precipitation episode.
53 For more info and results about the comparison between rBC and EC snow/ice measurement see Sigl
54 et al. (2018). On the contrary, the OC atmospheric concentration showed a decreasing trend with the
55 highest values at the beginning of the sampling period and the lowest at the end, similarly to the
56 atmospheric eBC behavior. Remarkably, the highest OC concentration was found in the same sample
57 where the highest concentration of all the other measured compounds was found (at the very beginning
58 of the snow episode) suggesting a common atmospheric scavenging process, although not above the
59 average for the rBC mass concentration.

60 **4. Sodium/manganese concentrations measurements**

61 Concentrations of sodium (Na) and manganese (Mn) were also determined as tracer of sea
62 spray emission and dust deposition by Inductively Coupled Plasma Sector Field Mass Spectrometry
63 (ICP-SFMS; Element2, ThermoFischer, Bremen, Germany) equipped with a cyclonic Peltier-cooled
64 spray chamber (ESI, Omaha, USA). The sample flow was maintained at 0.4 mL min^{-1} . Detection
65 limits, calculated as three times the standard deviation of the blank, were 0.5 ng g^{-1} for ^{23}Na and 0.3
66 ng g^{-1} for Mn. The residual standard deviation (RSD) for Na and Mn ranged between 2–5%.

67 **5. Back trajectories calculation and Potential Source Contribution Function analysis**

68 Air mass back-trajectories (BT) were calculated using the NOAA ARL HYSPLIT 4 rev. 513
69 transport model (Stein et al., 2015). Global Data Assimilation System (GDAS) meteorological input
70 fields with 0.5×0.5 degree resolution and a propagation time of 240 hours was employed. The
71 trajectories were calculated every hour for an endpoint of 500 m above ground level in Ny-Ålesund. A
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73 potential source contribution function (PSCF) analysis has been applied to the BTs exploiting a
74 specifically developed FORTRAN computer code (Petroselli et al., 2018). That analysis considered
75 BC concentration measured in the air by both AE31 and PSAP. Briefly, the method calculates the
76 probability of finding a source of a particular pollutant on a certain region by superimposing grid cells
77 to it and estimating the fraction of the total time spent on each cell by trajectories associated with a
78 high concentration measured at the receptor site. The 90th percentile was used to define the high
79 concentration limit and cells of 3 x 3 degrees (lat-long) were exploited in the calculation of
80 probabilities. Details of the PSCF methodology employed here are described in Petroselli et al.,
81 (2018). The data of the active fires, covering the last 12 days before the sampling day, are from the
82 MODIS active fire products (<https://firms.modaps.eosdis.nasa.gov/firemap/>), offered by NASA
83 LANCE.

84 In order to evaluate the impact of the Eurasian fires on the measured atmospheric eBC
85 concentrations, a thorough back-trajectories analysis was performed for both the snow-sampling
86 periods. Results of PSCF analysis on eBC (Figures SI 1a, SI 1b and SI 1c; open-fire episodes are
87 reported in red on the map) show a clear maximum of probability over the Central Siberia, which
88 appears to be the major source area of eBC in this period over Ny-Ålesund. Some false positive
89 source areas are located in Greenland, the Queen Elisabeth Islands region and the Arctic Ocean, even
90 if associated to a lower probability. These artifacts are due to the persistent circulation of BTs in the
91 Arctic vortex. An example of BTs generating the above salient features in the PSCF plot is reported in
92 Figure SI 1b. Here BTs are shown to loop for few days around the Arctic at high altitudes and
93 afterwards to descend at lower altitudes over Siberia, just four days before reaching Ny-Ålesund on
94 April 22, when a clear maximum in the eBC trend has been recorded. Back trajectory analysis
95 supports the idea that the peaks of eBC in the atmosphere in early spring are directly correlated with
96 long-range transport from Eurasia, whereas the peaks in late May and June are much lower in
97 intensity and they seem to be more related to a Western circulation pattern.

98 The results of BT analysis for the 3-days experiment are reported in Figure SI 2, suggesting
99 that the air masses were persistently circulating in the polar vortex causing a similar weather condition
100 within the three days of the experiment.

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107 **Table S1.** Abakus Klotz selected dimensional bins.

Channel	Size (μm)				
1	0.8	11	2.2	22	6.4
2	0.9	12	2.4	23	7.1
3	1	13	2.6	24	7.8
4	1.1	14	2.9	25	8.6
5	1.2	15	3.2	26	9.5
6	1.3	16	3.5	27	10.5
7	1.4	17	3.9	28	11.6
8	1.6	18	4.3	29	12.8
9	1.8	19	4.8	30	14.1
10	2	20	5.3	31	15.5
		21	5.8	32	80

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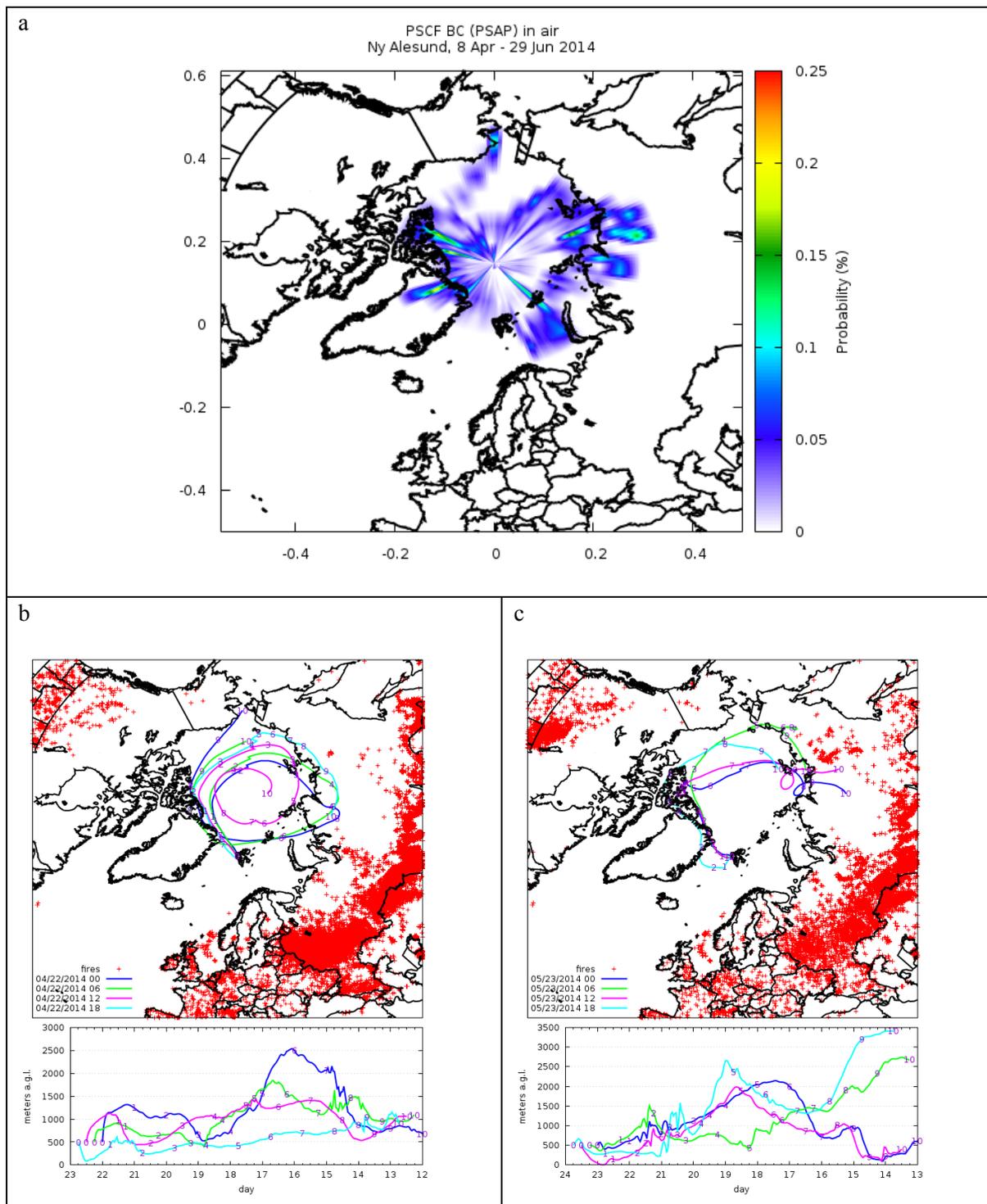
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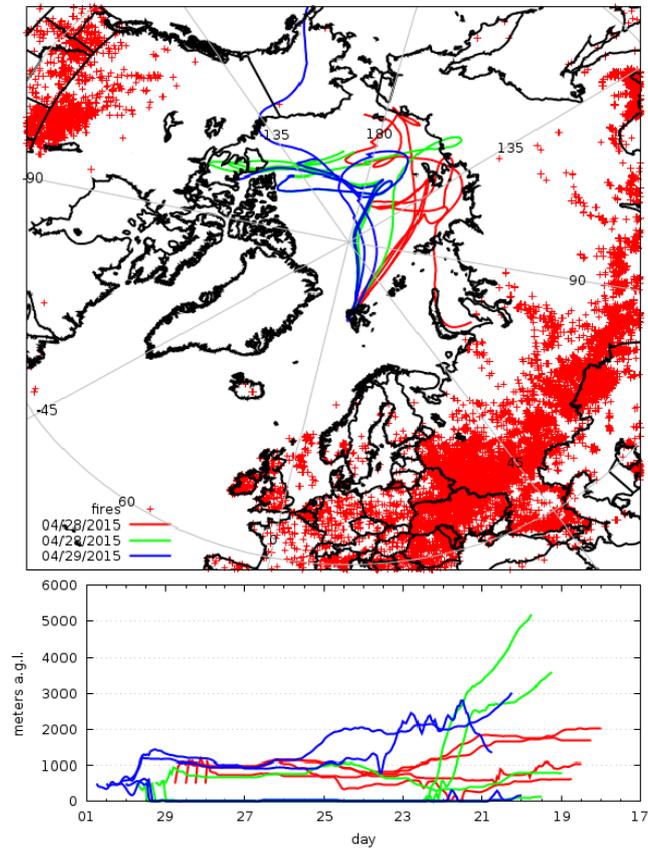
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126 **Figure S1.** a) Potential source contribution function analysis (PSCF) of eBC recorded in the 85-days
 127 experiment (8 April-29 June). 10 days back-trajectories for two selected days: b) 22 April and c) 23
 128 May. Four BTs were calculated for the two selected days, with a 6 hours interval. The red crosses
 129 represent the fires taking place during the last 10 days (data from the MODIS active fire products
 130 (<https://firms.modaps.eosdis.nasa.gov/firemap>)).



132 **Figure S2.** 10 days back-trajectory results for the 3-days experiment. The red crosses represent the
133 fires taking place during the last 10 days (data from the MODIS active fire products ([https://firms.
134 modaps.eosdis.nasa.gov/firemap/](https://firms.modaps.eosdis.nasa.gov/firemap/))).



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145 **Figure S3.** Results for the 85-days experiment from ancillary measurements. Upper panel: rBC mass
146 concentration (gray line), Na concentration (red line) and conductivity (green line). Lower panel:
147 atmospheric eBC mass concentration (black line) and ammonia as measured at the Zeppelin station,
148 Svalbard (gray bars).

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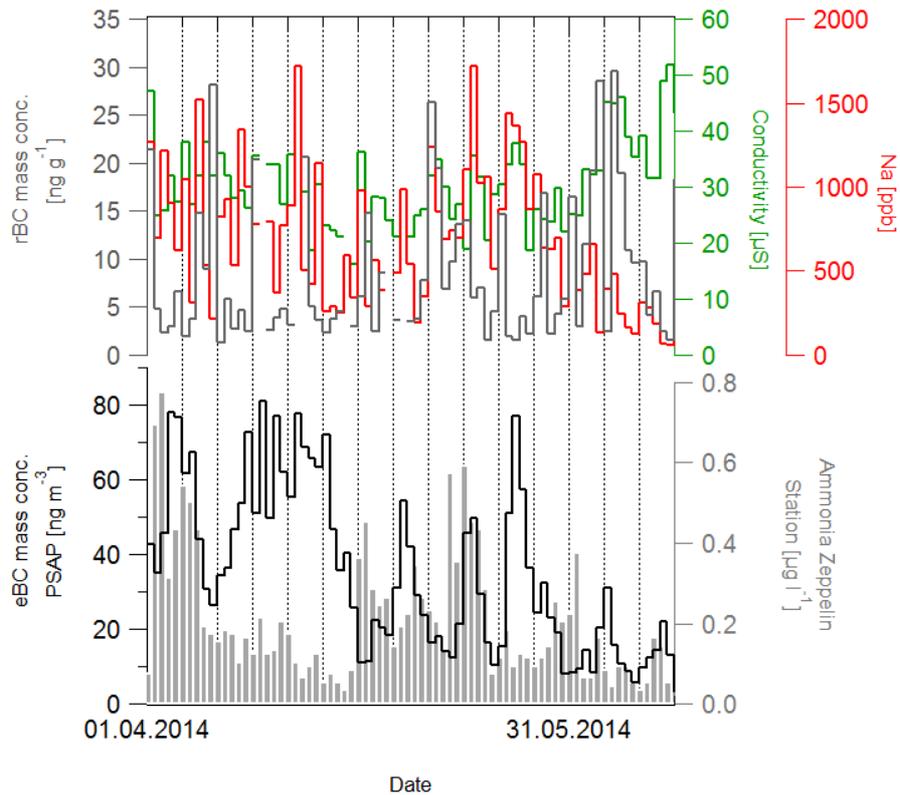
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170 **Figure S4.** Ancillary measurements for the 3-days experiments. Uppermost panel: manganese (Mn, 171 dark-yellow line) mass concentration and the coarse mode particles number (blue line). Second 172 uppermost panel: rBC mass concentration (gray line) with sodium (Na) concentration (red line) and 173 conductivity (green line). Second lowermost panel: atmospheric eBC mass concentration (black line), 174 snow OC concentration (blue bars) and daily average of atmospheric ammonia as measured at the 175 Zeppelin station (gray bars). Lowermost panel: TC (red bars), OC (blue bars) and EC (green bars) and 176 EC daily average (black line).

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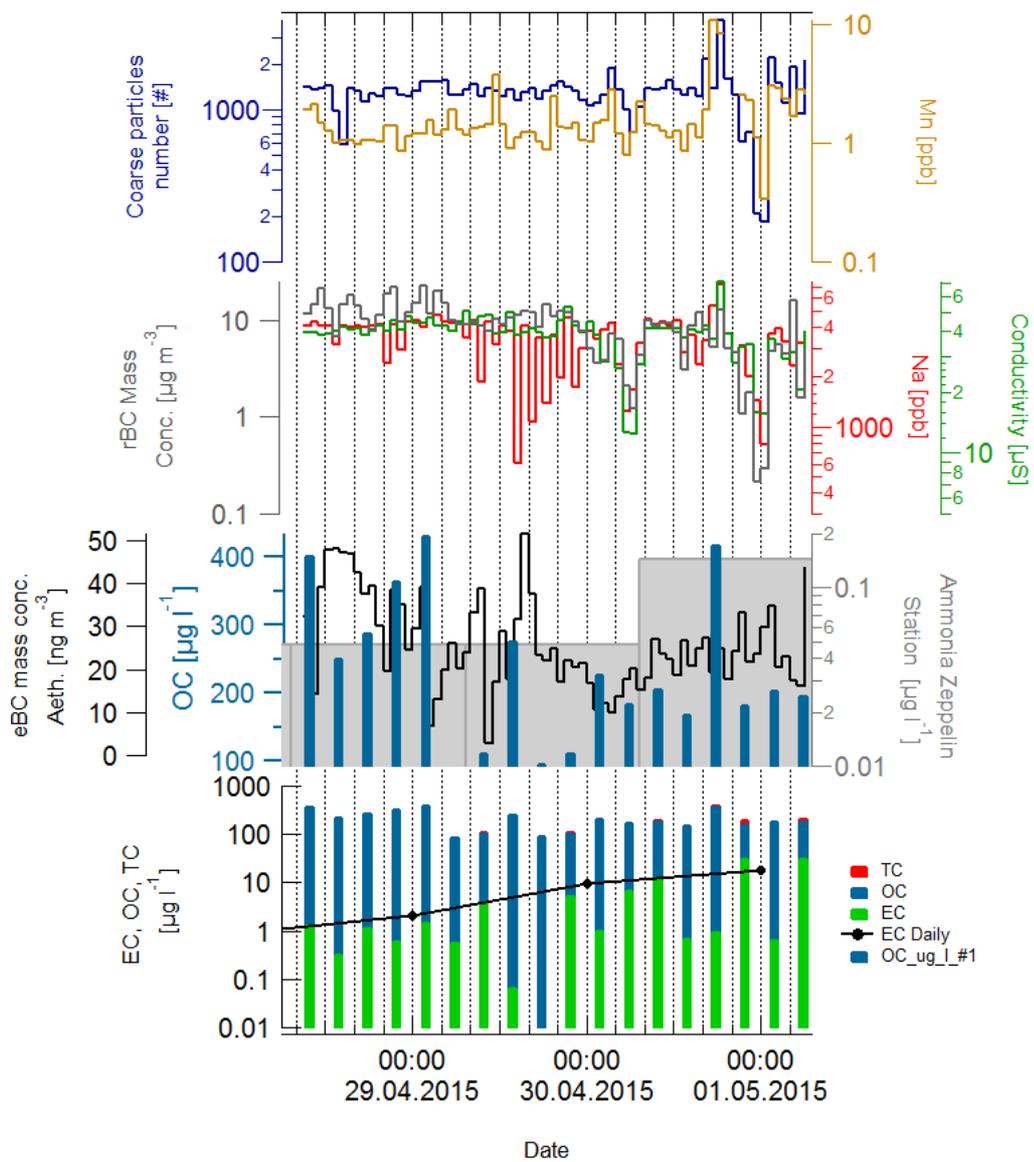
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196 **References:**

- 197 Kaspari, S. D., Schwikowski, M., Gysel, M., Flanner, M. G., Kang, S., Hou, S. and Mayewski, P. A.:
198 Recent increase in black carbon concentrations from a Mt. Everest ice core spanning 1860–2000
199 AD, *Geophysical Research Letters*, 38(4), doi:10.1029/2010GL046096, 2011.
- 200 Kupiszewski, P., Zanatta, M., Mertes, S., Vochezer, P., Lloyd, G., Schneider, J., Schenk, L.,
201 Schnaiter, M., Baltensperger, U., Weingartner, E. and Gysel, M.: Ice residual properties in mixed-
202 phase clouds at the high-alpine Jungfrauoch site, *Journal of Geophysical Research: Atmospheres*,
203 121(20), 12,343–12,362, doi:10.1002/2016JD024894, 2016.
- 204 Moteki, N., Adachi, K., Ohata, S., Yoshida, A., Harigaya, T., Koike, M. and Kondo, Y.:
205 Anthropogenic iron oxide aerosols enhance atmospheric heating, *Nature Communications*, 8, 15329,
206 doi:10.1038/ncomms15329, 2017.
- 207 Sigl, M., Abram, N. J., Gabrieli, J., Jenk, T. M., Osmont, D. and Schwikowski, M.: 19th century
208 glacier retreat in the Alps preceded the emergence of industrial black carbon deposition on high-
209 alpine glaciers, *The Cryosphere*, 12(10), 3311–3331, doi:https://doi.org/10.5194/tc-12-3311-2018,
210 2018.
- 211 Petroselli, C., Crocchianti, S., Moroni, B., Castellini, S., Selvaggi, R., Nava, S., Calzolari, G.,
212 Lucarelli, F. and Cappelletti, D.: Disentangling the major source areas for an intense aerosol
213 advection in the Central Mediterranean on the basis of Potential Source Contribution Function
214 modeling of chemical and size distribution measurements, *Atmospheric Res.*, 204, 67–77,
215 doi:10.1016/j.atmosres.2018.01.011, 2018.
- 216 Stein, A. F., Draxler, R. R., Rolph, G. D., Stunder, B. J. B., Cohen, M. D. and Ngan, F.: NOAA’s
217 HYSPLIT Atmospheric Transport and Dispersion Modeling System, *Bull. Am. Meteorol. Soc.*,
218 96(12), 2059–2077, doi:10.1175/BAMS-D-14-00110.1, 2015.