



1 Vertical observations of aerosol properties in the Arctic, Antarctic and Alpine 2 atmospheric boundary layers with a tethered balloon

3 Julia Schmale¹, Roman Pohorsky¹, Yolanda Temel¹, Michael Lonardi¹, Joanna Alden^{1,*}, Radiance
4 Calmer¹, Evan Thomas², Andrea Baccarini^{1,+}, Nora Bergner¹, Berkay Dönmez¹, Lionel Favre¹, Henrik
5 Helmig¹, Benjamin Heutte¹, Marie Lou Hirschy¹, Romain Phan¹, Orpheas Solidakis¹, Aleksandrina
6 Streltsova¹, Mihnea Surdu¹, Arthur Tabary¹

7 ¹Extreme Environments Research Laboratory, École Polytechnique fédérale de Lausanne, Sion, 1951, Switzerland

8 ²École Polytechnique fédérale de Lausanne, EPFL Valais Wallis, Sion, 1951, Switzerland

9 * now at Centre Alpin de Recherche sur les Réseaux Trophiques et Écosystèmes Limniques, Université Savoie Mont Blanc:
10 Chambéry, Auvergne-Rhône-Alpes, FR

11 + now at Aerospec, Lausanne, Switzerland

12 *Correspondence to:* Julia Schmale (Julia.schmale@epfl.ch)

13 **Abstract.**

14 Information on vertical profiles of aerosols in the lowermost kilometer of the atmosphere is still scarce in general, and
15 particularly in polar regions and complex terrain due to observational limitations. However, the vertical distribution of various
16 aerosol properties, such as the particle number size distributions, are essential climate variables, because aerosols influence
17 Earth's radiation budget and their vertical position in the atmosphere matters. Specifically for aerosol-cloud interactions, the
18 presence and properties of particles able to act as cloud condensation nuclei and ice nucleating particles at cloud level, is
19 critical information. With this data descriptor we aim to start filling the observational gaps systematically, and introduce
20 datasets of vertical aerosol properties from nearly 300 tethered balloon flights between 2022 and 2025 in the Arctic, Antarctica
21 and alpine terrain in Europe. Flights up to 900 m above ground describe various boundary layer conditions throughout all
22 seasons. We describe the processing from raw to level 2 data, provide processing and quick look codes, and give examples of
23 how data may be used and analyzed by the community. All data and code are openly accessible.

24 **1 Introduction**

25 Aerosol-radiation interactions (ARI) and aerosol-cloud interactions (ACI) continue to add uncertainty to anthropogenic
26 radiative forcing, and specifically ACI is the largest single contributor to the overall uncertainty (e.g., Bellouin et al., 2020;
27 Gordon et al., 2023). In part this challenge is due to a lack of observations of the vertical distribution of aerosols and their
28 properties. For example, the efficacy of the climatic effect of black carbon at the surface depends strongly on the height above
29 ground, where the absorbing particles reside, and in which geographical region (Flanner, 2013; Samset et al., 2013). In the
30 Arctic, black carbon has the strongest effect in the lowermost part of the atmosphere, while presence at high altitudes is much
31 less effective (Flanner, 2013). With reference to ACI, the number and size of aerosol particles at the height, where clouds form



32 and act as cloud condensation nuclei (CCN) and ice nucleating particles (INP), will influence the cloud droplet number
33 concentrations and size distributions as well as the presence of ice crystals (Carslaw et al., 2013; Kanji et al., 2017). The
34 resulting cloud microphysics determine cloud radiative properties (e.g., Zelinka et al., 2014) and precipitation processes (e.g.,
35 Rosenfeld et al., 2014).

36 To capture the potential effects of ARI and ACI, surface-based observations alone are not sufficient, because aerosol
37 concentrations and properties near the ground are not necessarily representative of those aloft due to the structure of the
38 troposphere, and vertical transport and atmospheric chemical processes (Clarke and Kapustin, 2010). Aerosol properties in the
39 free troposphere are typically different from the boundary layer (e.g., Brock et al., 2019; Collaud Coen et al., 2011; Engelmann
40 et al., 2021), and even within the remote and urban boundary layer (BL) they might vary substantially (Lei et al., 2021; Pilz et
41 al., 2024; Pohorsky et al., 2025b; Rosati et al., 2015). This variation owes to the thermodynamic structure, particularly
42 inversions that can occur at the surface (surface-based inversion) or at elevated heights (elevated inversions) within the BL
43 (Ferrero et al., 2016; Pace et al., 2015; Prasad et al., 2022; Thomas et al., 2019). Those typically inhibit the up-mixing of
44 surface emissions and the down-mixing of elevated aerosol sources.

45 Observations aiming to elucidate the climatic effects of ARI and ACI are however often confined to the surface, e.g., in
46 established monitoring networks such as Global Atmosphere Watch (GAW) or the European Aerosols, Clouds and Trace
47 Gases research infrastructure ACTRIS (e.g., Asmi et al., 2013), because regular vertical profiling for in situ measurements is
48 challenging. The challenge results e.g., from the weight and complexity of the instruments such that operation typically requires
49 research aircraft with operators or helicopters towing complex flying laboratories (Pätzold et al., 2023; Siebert et al., 2021).
50 This means that both ARI and ACI processes as well as essential climate variables (ECVs) of aerosols and clouds are not easy
51 to observe along vertical profiles. Vertical ECVs of aerosols include, amongst others, particle number size distribution, number
52 of CCN, and optical properties such as extinction coefficients and single scattering albedo (The 2022 GCOS Implementation
53 Plan (GCOS-244), 2025). ECVs of clouds are, amongst others, cloud droplet effective radius and cloud liquid and ice water
54 contents. In addition to ECVs, detailed process observations are necessary to gain insights into unresolved questions, for
55 example: in which thermodynamic conditions are surface-based observations representative of aerosol properties at cloud
56 level; whether clouds are seeded by aerosols from below, aloft or both in varying thermodynamic conditions; or whether the
57 number of INP can explain the number of ice crystals or if secondary ice processes are at work.

58 Remote sensing, and specifically ground-based remote sensing, can deliver information on cloud properties and to a certain
59 degree also on aerosol properties along the vertical profile (Ansmann et al., 2023; Engelmann et al., 2021; Radenz et al., 2024).
60 However, unless the suite of remote sensing instrumentation is highly sophisticated, uncertainties can remain large and certain
61 proxies, e.g., for aerosol and cloud droplet number size distributions, cannot be retrieved (Gast et al., 2025; Schlosser et al.,
62 2022). In recent years, there have been significant advances in vertical observations of the aerosol and cloud ECVs mentioned
63 above, mainly through development and deployment of uncrewed aerial vehicles (Altstädter et al., 2015; de Boer et al., 2022;



64 Lappin et al., 2024; Pätzold et al., 2023) and tethered balloons (Creamean et al., 2021; Ferrero et al., 2016; Pasquier et al.,
65 2020; Pilz et al., 2022; Pohorsky et al., 2024; Porter et al., 2020), as well as due to progress in miniaturized high-quality
66 instrumentation (Liu et al., 2021b; Pilz et al., 2022; Pohorsky et al., 2024). Nevertheless, in situ vertical measurements of
67 aerosols and clouds remain sparse. Consequently, there remains a gap in information on ECVs, other variables and in process
68 understanding (Creamean et al., 2021; Pilz et al., 2024; Schmale et al., 2021). Filling this gap is important to constrain satellite-
69 based observations that have a larger temporal and geographical coverage than in situ measurements, as well as numerical
70 simulations that ultimately determine the climatic effects of ARI and ACI.

71 Here, we present quality assessed data on aerosol and cloud droplet properties from tethered balloon (Helikite, Allsopp, UK)
72 observations in the BLs and lower free tropospheres of the Arctic, Antarctica and mountainous terrain in Europe. The overall
73 number of flights is 276 up to an altitude of 990 m above ground at the time of writing this manuscript. We focus on polar and
74 mountain BLs because they often feature multiple layers of aerosols due conditions with reduced vertical mixing (e.g., Jozef
75 et al., 2023; Rotach and Zardi, 2007), and specifically in polar regions, low level clouds within or at the top of the BL exert a
76 strong effect on the surface energy budget (Shupe and Intrieri, 2004). This data descriptor provides information on how data
77 were processed, which uncertainties to expect, and provides guidance on how the data can be used for scientific research.

78 **2 Overview on campaigns**

79 Campaigns were carried out between 2021 and 2025 in nine locations, thereof five in the Arctic, one in coastal Antarctica, two
80 in the Swiss Alps, and one in the Greek Peloponnese mountains (see details in Table 1 and map in Fig. 1). An online overview
81 map (<https://www.epfl.ch/labs/eerl/eerl-home-page/research/field-work-overview/>) provides interactive information on
82 locations, associated publications and collaboration partners. Here, we briefly present the purpose and larger context for each
83 campaign.

84 **Arctic campaigns:** The common denominator for these campaigns is the characterization of aerosol layers throughout the
85 often-stratified Arctic BL and the investigation of in how far surface observations can be representative of aerosol above the
86 surface mixed layer, which can be as shallow as 20 m (Pohorsky et al., 2025b). Specifically, the Alaskan Layered Pollution
87 And Chemistry Analysis (ALPACA) project focused on extreme wintertime urban pollution and BL stratification (Simpson et
88 al., 2024) as well as the up-mixing of surface and down-mixing of power plant emissions (Pohorsky et al., 2025b). Observations
89 at Villum Research Station with the CLeancloud Arctic Villum ExpeRiment (CLAVIER) focused on the transition period
90 between winter and spring, within a highly stratified BL including haze from long-range transport, new particle formation
91 events, and a warm air-mass intrusion. The GreenFjord campaign aimed to investigate the vertical footprint of climate-relevant
92 aerosol properties of local urban emissions in an environment which is otherwise influenced by marine emissions and
93 occasional long-range transported pollution (Alden et al., 2025b). The Pallas Cloud Experiment (PaCE) was geared towards
94 understanding the vertical extent of new particle formation and aerosol-cloud interactions in a boreal environment (Brus et al.,



95 2025). And finally, the ARTofMELT (AOM) expedition focused on the melt onset in the marginal ice zone and the role of
 96 aerosol-cloud interactions for the surface energy budget.

97 **Antarctic campaign:** The project ‘ORigin of Antarctic CLoud particles and their Effects on the Surface radiation budget’
 98 (ORACLES) investigates the vertical profiles of aerosol properties and ACI as a function of marine and Antarctic continental
 99 air masses as well as the BL structure at Neumayer III Station operated by the Alfred Wegener Institute.

100 **Alpine campaigns:** Both Brigerbad and Turtmann are located in the narrow Rhone valley of the Swiss Alps, where dynamic
 101 air mass movement is induced by down and up-valley winds in the morning and afternoon, respectively, as well as by the
 102 appearance of sunlight on the slopes and valley floor. Focus was on the dispersion of aerosols during transition seasons, i.e.,
 103 between summer and winter in Brigerbad, and winter to summer in Turtmann. In Greece, observations were carried out half-
 104 way up Mt. Helmos with the Cleancloud Helmos OrograPhic sIte experimeNt (CHOPIN) with the aim to measure aerosol
 105 properties throughout the evolution of the mountain BL and aerosol-cloud interactions under local and long-range transported
 106 aerosol pollution.

107 **Table 1: Overview information on flights per campaign. SBI means surface-based inversion, EI means elevated inversion.**

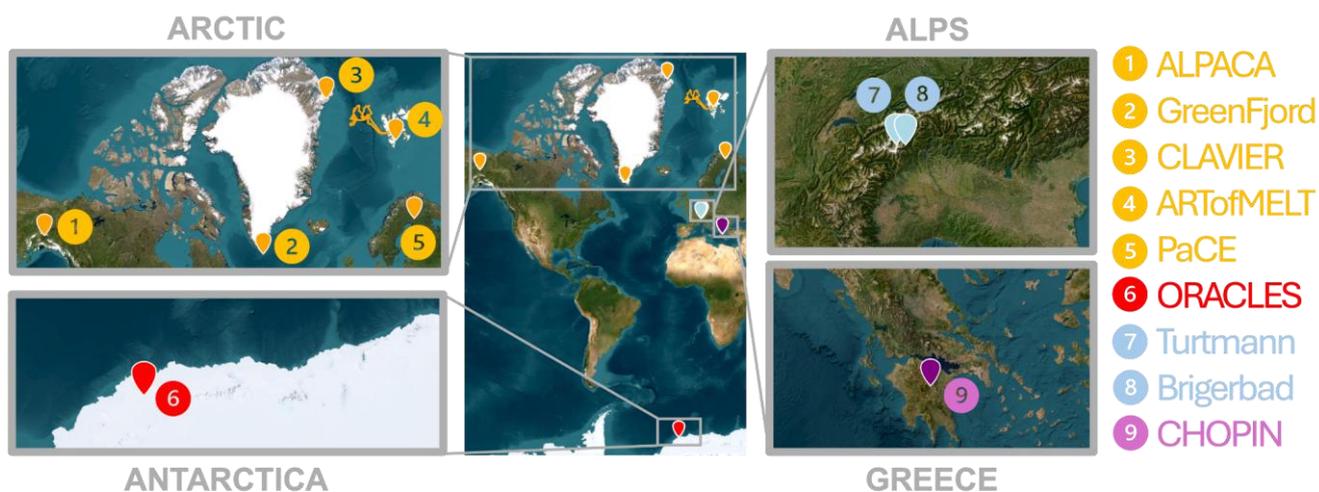
Campaign	Environment	Dates	Coordinates, elevation above ground level (a.s.l.)	Helikite model	No. of flights	Maximum flight altitude	No. of cloud cases	No. of inversions
ALPACA	Suburb of Fairbanks, Alaska, USA	2022-01-26 to 2022-02-25	64.853 N, 147.859 W, 138 m	45 m ³	24	345 m	0	14 (SBI) 4 (EI)
ARTofMELT	Arctic Ocean, marginal ice zone	2023-05-16 to 2023-06-10	80.482 to 78.842 N, 3.778 W to 6.267 E, 0 m	45 m ³	23	645 m	10	4 (SBI) 10 (EI)
Brigerbad	Narrow alpine valley surrounded by high mountains	2021-09-22 to 2021-10-14	46.300 N, 7.921 E, 652 m	45 m ³	26	760 m	0	9 (SBI) 0 (EI)
CHOPIN	Peloponnese (Greece), mountain ski resort	2024-10-23 to 2024-11-05	38.005 N, 22.198 E, 1700 m	45 m ³	23	527 m	8	2 (SBI) 8 (EI)
CLAVIER	Northern	2024-03-	81.581 N,	45 m ³	57	750 m	2	48 (SBI)



	Greenland, flat terrain near coast	23 to 2024-05-02	16.641 E				(fog)	4 (EI)
			30 m					
GreenFjord	Southern Greenlandic fjord system	2023-06-27 to 2023-08-02	60.916 N, 46.053 W, 17 m	64 m ³	24	780 m	7	7 (SBI), 14 (EI)
ORACLES	Coastal Antarctic, flat terrain, 6 km from coast	2024-12-22 to 2025-02-15	70.662 S, 8.286 W 40 m	24 m ³ and 64 m ³	68	992 m	21	11 (SBI) 23 (EI)
PaCE	Boreal forest	2022-09-28 to 2022-10-13	68.000 N, 24.239 E 334 m	45 m ³	19	539 m	12	1 (SBI), 10 (EI)
Turtmann	Narrow alpine valley surrounded by high mountains	2024-02-07 to 2024-02-28	46.305 N, 7.691 E, 640 m	64 m ³	12	700 m	0	5 (SBI) 4 (EI)

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Figure 1: Map of campaign locations. Basemap tiles: Esri | Powered by Esri, DigitalGlobe, GeoEye, i-cubed, USDA FSA, USGS, AEX, Getmapping, Aerogrid, IGN, IGP, swisstopo, and the GIS User Community.

113



114 3 Instrumentation

115 The EERL Helikite payload is called MoMuCAMS (Fig. 2) for ‘Modular Multiplatform Compatible Atmospheric
116 Measurement System’. The instruments have previously been described and characterized in Pohorsky et al. (2024 and
117 references therein). Since that publication, more instruments have been included for regular operations, specifically, a handheld
118 condensation particle counter (CPC model 3007, TSI), a Partector 2 (Naneos), a microAethalometer (microAeth, Aethlabs), a
119 miniaturized Cloud Droplet Analyzer (miniCDA, Palas) and a light optical aerosols counter (LOAC, LPC2E – CNRS). Table
120 2 provides a summary of all instruments operated per campaign.

121



122

123 **Figure 2: The Extreme Environments Research Laboratory (EERL) Helikite with payload MoMuCAMS and SmartTether during**
124 **deployment at Neumayer Station III, Antarctica, 2025. Picture credit: M. Lonardi.**

125



126 **Particle counters and particle number size distributions.** The aerosol number size distribution is obtained from the
127 miniaturized Scanning Electrical Mobility Sizer (mSEMS model 9404, Brechtel Manufacturing Inc.) and the Portable Optical
128 Particle Spectrometer (POPS, Handix Scientific). The mSEMS measures the electrical mobility diameter of particles ranging
129 from 8 to a maximum of 300 nm. The time resolution depends on the selected size resolution (number of bins) and the time
130 spent measuring in each size bin. These parameters are typically defined according to the local environment and ambient
131 aerosol concentration. For an urban environment, a 60-bin resolution with a 1 s bin time was selected, yielding a 1-min scan
132 time (Pohorsky et al., 2025b). In pristine areas, where concentrations can fall below 10 cm^{-3} , the bin time needs to be increased
133 to allow for sufficient counting statistics (Alden et al., 2025b). To limit the overall scan time, the number of bins is typically
134 reduced (e.g., 40 bins).

135 The POPS measures the optical diameter of particles ranging from $\sim 150 \text{ nm}$ to $> 3 \mu\text{m}$. The optical diameter is obtained from
136 the scattering signal of particles sampled through a laser beam and converted to a size, assuming a spherical diameter with a
137 refractive index of $1.615 + 0.001i$ at 405 nm (i.e., refractive index of polystyrene latex spheres). Note that these assumptions
138 can lead to sizing errors in environments with highly absorbing and/or non-spherical particles. The POPS records data at a 1
139 Hz resolution and classifies particles into 16 log-spaced size bins. For both instruments, the concentrations in each bin are
140 normalized (i.e., expressed as $dN/d\log Dp$) to account for the logarithmic bin spacing.

141 For the total aerosol number concentration, the advanced Mixing Condensation Particles Counter (aMCPC model 9403,
142 Brechtel Manufacturing Inc.) or the handheld Condensation Particle Counter (CPC3007, TSI) are used. The size detection cut-
143 offs (d_{50}) are approximately 7 and 10 nm, respectively, and data are recorded at a 1 Hz time resolution and expressed in particles
144 per cubic centimeter (cm^{-3}). The handheld CPC is a well characterized concentration particle counter, which has also been
145 used in polar regions (Pilz et al., 2023).

146 The Partector 2 is a nanoparticle detector (Fierz et al., 2014, 2015) occasionally deployed as a backup particle counter on the
147 MoMuCAMS due to its low weight (415 g) and ease of use. The instrument uses an open-path electrometer followed by an
148 electrostatic deposition stage and another open-path electrometer in an in-series configuration, operating with a flow rate of
149 0.5 l min^{-1} (Bezantakos et al., 2024). The Partector 2 detects particles in the size range of 10 to 300 nm, covering concentration
150 ranges between 0 and $10^6 \text{ particles cm}^{-3}$. The measurements are obtained with 1 Hz temporal resolution, additionally recording
151 ambient pressure for subsequent altitude calibration. The calibrated data, has a 30% uncertainty in the accuracy
152 (<https://www.naneos.ch/partector2.html>). Poor signal to noise ratio can be overcome by averaging over time.

153
154 **Droplet number size distribution.** The miniaturized cloud droplet analyzer (miniCDA, Palas GmbH) is a new, miniaturized
155 optical particle counter for cloud droplets (Solidakis, 2024). It has a specific inlet oriented vertically and protected against
156 precipitation, providing a flow rate of 2.8 l min^{-1} . The measurements can be used to derive ECVs such as droplet effective
157 radius and liquid water content. The instrument detects hydrometeor and particle size distributions in the range of 0.24 to 43
158 μm using 256 manufacturer-calibrated size bins, covering concentration ranges between 0 and 600 droplets cm^{-3} . The
159 measurements are obtained with a 1.5 s temporal resolution, complemented by pressure measurements (every 15 s), which are



160 used for data synchronization with the MoMuCAMS flight computer. There are no peer-reviewed publications yet on this
161 instrument.

162 The LOAC has been used during the ARTofMELT campaign to measure the size distribution of water droplets in (Renard et
163 al., 2016). The LOAC measures the aerosol and droplet size distribution from 0.2 to 50 μm , at a time resolution of 30 s. Two
164 detectors measure the scattering, at 12° and 60° , produced by the sampled particles traveling through a laser beam. The intensity
165 of the forward scattering signal is used to measure the size of the particles and the difference between the two angles provides
166 additional information to classify particles types (e.g., carbonaceous, minerals, liquid droplets).

167 The miniCDA is deployed with larger balloons, while the LOAC is routinely deployed on balloons smaller than 24 m^3 .

168

169 **Optical properties.** For light absorbing particles, filter-based absorption detection instruments are used. The single Channel
170 Tricolor Absorption Photometer (STAP model 9406, Brechtel Manufacturing Inc) provides the aerosol light absorption
171 coefficient at three wavelengths (450, 525 and 624 nm, e.g., Pilz et al., 2022). The absorption data are initially recorded at a 1
172 Hz resolution and then averaged over a longer integration time. For a 2-min integration, the detection limit of the instrument
173 is 0.2 Mm^{-1} , which corresponds to an equivalent black carbon (eBC) mass concentration of about 20 ng m^{-3} (Bates et al.,
174 2013).

175 The MicroAeth (MA200, AethLabs) is also a filter-based absorption photometer measuring aerosol light attenuation at five
176 wavelengths (375, 470, 528, 625 and 880 nm, Liu et al., 2021a). Aerosols are continuously deposited onto a
177 Polytetrafluoroethylene (PTFE) filter band, and the attenuation through the filter is then related to an eBC mass concentration.
178 On MoMuCAMS, measurements were recorded at 60 s time resolution in SingleSpot™ mode for a detection limit of 30 ng m^{-3}
179 eBC for a 5-min average at a 150 ml min^{-1} flow rate.

180 The choice between STAP and MicroAeth depends on instrument weight and expected ambient eBC concentrations.

181

182 **Particle sampling.** An 8-channel filter sampler (FILT Model 9401, Brechtel Manufacturing Inc., USA) enables time-resolved
183 aerosol collection on various filter media. The FILT can operate at a flow rate between 0.5 and 3.3 lpm with an accuracy of
184 $\pm 5\%$. The sampling channel can be switched remotely, allowing up to 8 samples to be collected during a single flight.
185 Housekeeping data of the FILT, including channel position and flow rate are recorded at a 1 Hz time resolution and used
186 afterwards to calculate the total sampled volume.

187

188 **Trace gases.** CO is measured with the Mira Pico (Aeris Technologies). The Mira Pico is based on non-dispersive infrared
189 (NDIR) spectroscopy. For ground-based operations, the instrument is operated with a CO scrubber, which allows for
190 continuous corrections of the instrument's baseline (zero measurement) and a ppb level accuracy according to the
191 manufacturer. This mode of operation reduces the time resolution. For vertical profiling, the instrument is operated in its
192 "manual" mode (without live baseline corrections). The time resolution is 1 Hz and the uncertainty reaches a maximum of 17
193 ppb due to potential fluctuations of the baseline during the flight (Pohorsky et al., 2024).



194 CO₂ is measured with an infrared sensor (GMP343, Vaisala). The instrument measures the absorption of a NDIR beam by CO₂
195 molecules through an open path cavity. A sensor switches every second between the absorbing band of CO₂ and a reference
196 wavelength, yielding a measurement time resolution of 2 s with an accuracy of 3 ppm +/- 1% according to the manufacturer.
197 O₃ is measured with the Dual Beam Ozone monitor (model 205, 2BTech). The instrument measures the intensity reduction of
198 a UV beam from O₃ molecules. A reference detection cell with an O₃ scrubber is used in parallel to improve its accuracy. The
199 time resolution of the O₃ monitor is 2 s with an accuracy of 1 ppb or 2% of the readings for 10-s averaged data.

200

201 **Meteorological variables.** On the MoMuCAMS payload, meteorological variables are obtained from various sensors placed
202 on the outside of the instrumental platform and complement the information of aerosol and trace gas measurements.
203 Temperature and relative humidity (both with a pair of sensors SHT85, Sensirion), barometric pressure (BME280, Bosch),
204 wind speed and wind direction (LI-550 TriSonica Mini, LiCOR) are typically recorded at a 1 Hz time resolution. In addition,
205 a SmartTether Weather Sonde (ST, Anasphere, Inc.) measures the latitude and longitude, temperature, relative humidity,
206 pressure, as well as wind speed and wind direction. The SmartTether operates at a 1 s time resolution with accuracies of ±0.5
207 mbar for pressure, ±0.5°C for temperature, ±1.7% for RH, ±0.1 m/s or 5% (whichever is greater) for wind speed, and ±2
208 degrees for wind direction.

209

210 **GPS.** GPS coordinates and attitude angles (pitch, roll, and yaw angles) are collected during flights with a VN200 GNSS/INS
211 (VectorNav).

212

213



214

215 **Table 2: Overview of instruments operated on each campaign. An ‘x’ denotes that the instrument was operated in general, however**
 216 **not necessarily on every flight.**

Instrument	Thermo-dynamics		particle number concentration and size distribution					Droplet number concentration		Aerosol absorption properties		Samp-ling	Trace gases		
	T and RH	SmartTether	m-SEMS	a-MCPC	CPC-3007	Parator2	POPS	LOAC	miniCDA	STAP	microAeth	FILT	CO ₂	Pico CO	Ozone
ALPACA	x		x				x			x		x	x		x
ARTofMELT	x	x	x				x	x		x		x			
Brigerbad		x	x	x			x			x		x	x	x	x
CHOPIN	x	x	x	x			x		x	x	x	x			
CLAVIER	x	x	x				x		x	x		x			
GreenFjord	x	x	x	x			x			x		x	x		x
ORACLES	x	x	x	x	x		x		x			x			
PaCE	x	x	x	x	x	x	x			x		x	x		
Turtmann	x	x	x				x			x		x			

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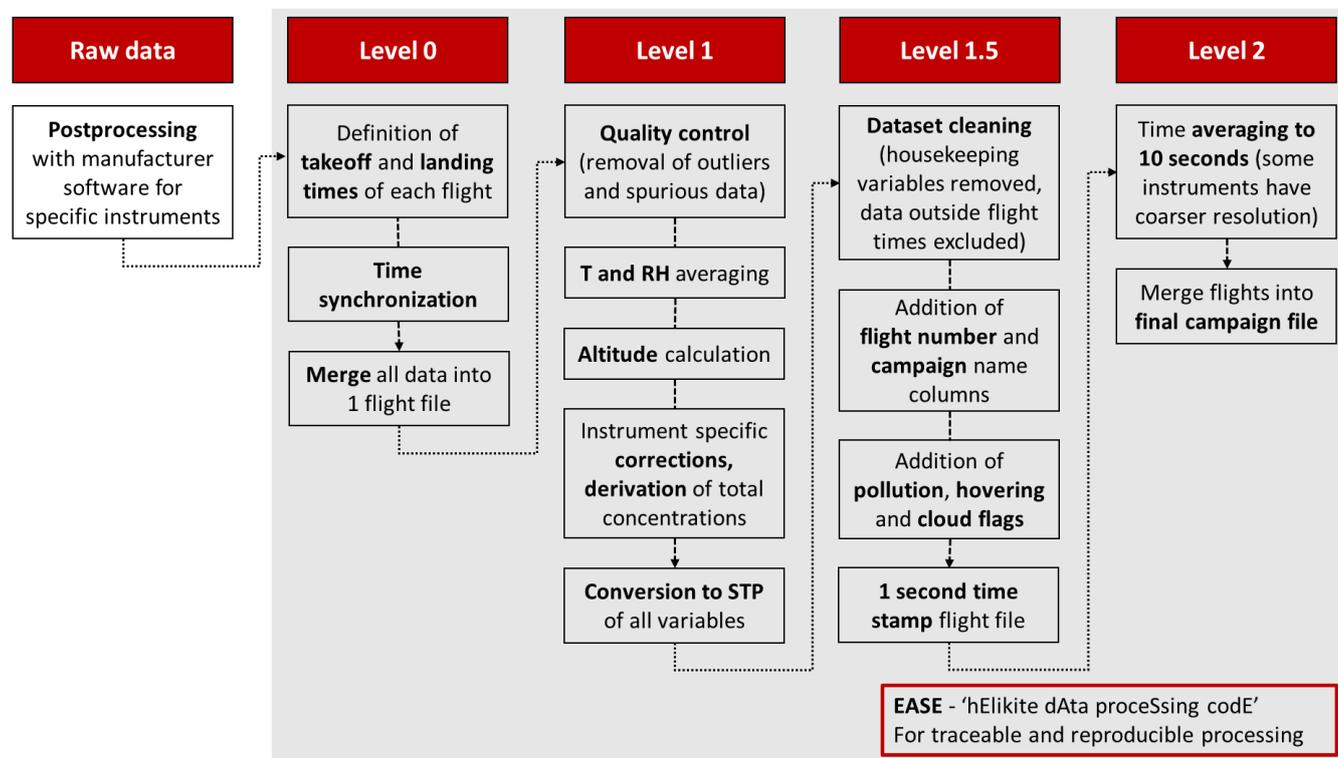


219 4 Data Processing and example analyses

220 4.1 Work flow and data levels

221 The MoMuCAMS is operated with a custom-made flight computer (FC), which collects data from most instruments and
222 provides a reference time stamp and one merged raw data file. Data from instruments, which are not connected to the flight
223 computer, are downloaded separately. To handle the various input files, a ‘hElkite dAta proceSsing codE’ (EASE) was
224 developed in Python. The code is structured such that future additional instrument files can be integrated. The objective of the
225 processing code is to generate quality controlled and assured data in a standard and traceable way before conducting scientific
226 analyses. It also ensures a homogenization of data processing steps and data products for different measurement campaigns.
227 Figure 3 summarizes the workflow. The following description provides details on the processing of data from raw to the final
228 level 2 with Table 3 providing a summary of all steps. The definition of levels follows in principle the guidelines by EBAS
229 (<https://ebas-submit.nilu.no/>), the data base of the ACTRIS network.

230



231

232 **Figure 3: Workflow diagram for EERL helikite data processing with the ‘hElkite dAta proceSsing codE’ (EASE).**



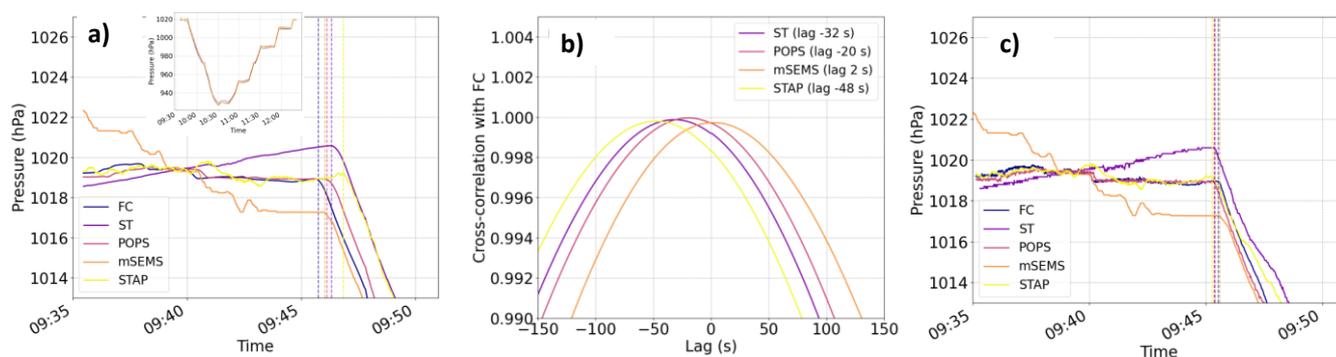
233 4.1.1 Raw data

234 Files directly generated from individual instruments are considered raw data and are imputed to EASE. There are exceptions
235 for two instruments, the STAP and mSEMS, because they require specific processing to obtain the desired variables. Codes
236 from the manufacturer to be run in Igor Wave Metrics are executed manually first and the output files are subsequently used
237 in EASE. For the STAP, the absorption coefficients are calculated based on a methodology developed by Ogren (2010). For
238 the mSEMS, an inversion routine as described in Wiedensohler et al. (2012), is applied to the raw particle counts to calculate
239 the particle number size distributions from the measured electrical mobility distributions. After these steps, all data are treated
240 to create level 0 (L0).

241 4.1.2 Level 0 data processing

242 At the end of the L0 processing step, all instrumental data are time synchronized, the start and end of each flight are defined
243 as takeoff and landing, and all variables are merged into one data file per flight.

244 While all internal clocks of instruments are synchronized at the beginning of each measurement campaign, time drifts of up to
245 a few seconds can occur. These drifts lead to shifted time series and vertical misalignments between the different variables,
246 which are not directly saved by the central flight computer. Because of the vertical motions of the Helikite, the variations in
247 the high-resolution pressure readings of each instruments allow to identify these time shifts. Using a cross-correlation function
248 for two independent variables, we use the pressure readings of the instruments to identify shifts between each instrument and
249 a reference instrument, whose time is known to be correct (i.e., flight computer). To find the time lag (τ) between instruments'
250 clocks, the pressure time series of the unsynchronized instrument is shifted by 1-s steps within a time window of ± 180 s with
251 reference to the flight computer's time. A Pearson correlation between the two pressure time series is then computed at each
252 τ . The τ with the highest Pearson correlation is then defined as the instrument's time shift and used to time synchronize the
253 measurements of the latter. Figure 4 illustrates the process.



254

255 **Figure 4: Example of time synchronization based on pressure data. The legend refers to individual instruments, where FC is flight**
256 **computer, and ST is SmartTether. a) Time series before synchronization, the insert shows pressure time series for the entire flight,**



257 **b) cross-correlation results and the associated lag per instrument, c) time series after synchronization. The vertical dashed bars**
258 **indicate the original and final time stamps.**

259

260 Adapted procedures are used for instruments that record at a time resolution other than 1 s. Pressure readings from the mSEMS
261 are obtained at 1 s resolution. However, the first step of processing with the manufacturer software provides the aerosol number
262 size distribution at a coarser resolution. The time resolution is between 1 and 3 minutes and depends on the number of bins
263 and the measurement time per bin, which can vary between campaigns based on the ambient aerosol concentration and the
264 trade-off between scanning time and counting statistics. We calculate the time lag from the raw mSEMS file and then apply it
265 to the coarser resolution data.

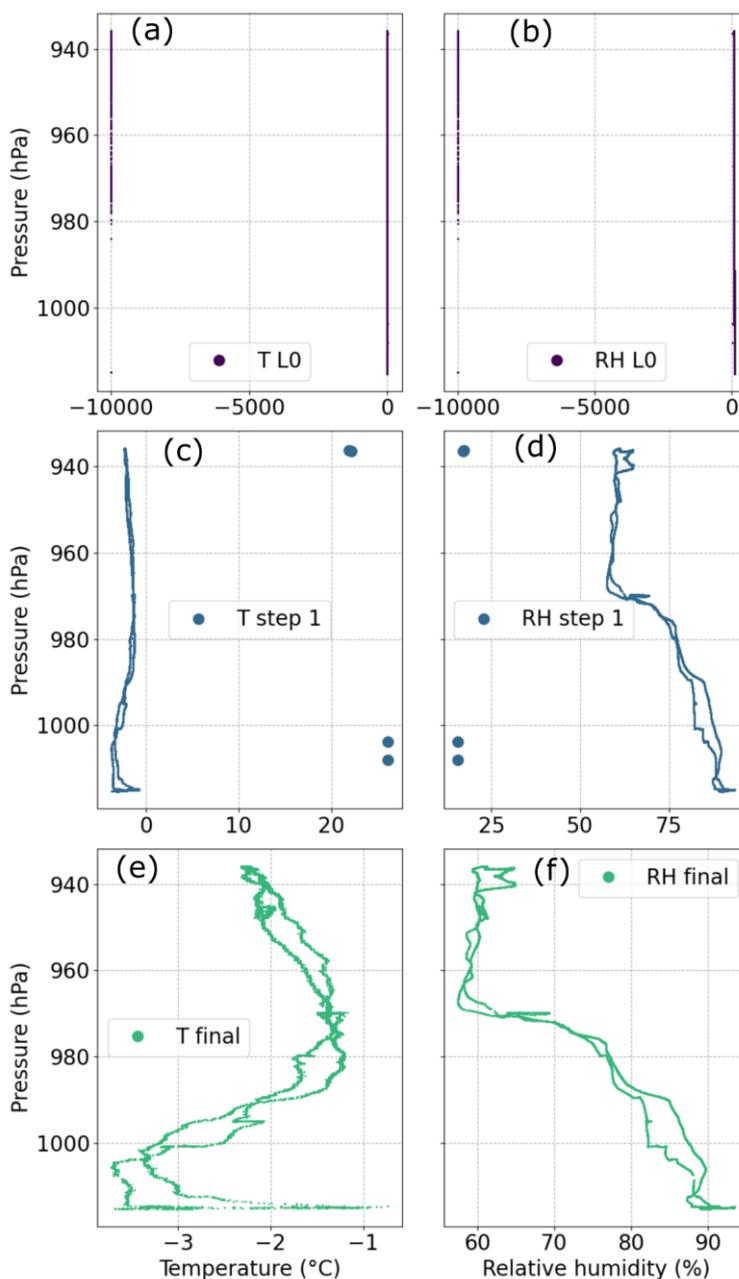
266 For instruments that do not record pressure data, regardless of their time resolution, such as the CPC3007 (1 s resolution), the
267 microAeth (1 min resolution) or the CO and O₃ sensors (1 and 2 s resolution, respectively), the internal clocks are regularly
268 synchronized with the flight computer to ensure accurate alignment based on timestamps.

269 After all data are time synchronized, the local time might be converted to UTC, depending on the campaign, and the data from
270 all instruments are merged into a single flight file.

271 **4.1.3 Level 1 data processing**

272 The objective of this step is to remove outliers, spurious data from instrument malfunction, and to apply all available data
273 corrections from e.g., instrument intercomparisons and calibrations. Flight altitude is calculated with a barometric equation.
274 The final step at this stage is the conversion of all data into standard temperature and pressure (STP) conditions. Hereafter, the
275 different processing steps for each variable are described.

276



277

278 **Figure 5: Example of outlier removal for one flight profile during ARTofMELT on 3 June 2023. a, b) L0 data for T and RH; c, d)**
279 **data after removing obvious outliers outside reasonable range (here negative values); e, f) final L1 data after despiking and visual**
280 **inspection.**

281 **Temperature, relative humidity, pressure and altitude.** Obvious outliers are data points outside of the realistic range of
282 values for a given variable (e.g., negative relative humidity or temperature above 100 °C). Those data points are removed first
283 based on thresholds of realistic values. Less obvious outliers could be within the conceivable measurement range, but are only



284 few data points with values different from the adjacent data points. They are identified by applying a despiking function, and
285 by subsequent visual inspection by the data processor. The despiking function is based on a comparison between the raw
286 measurements and the exponentially weighted moving average, with a 10 s window, which could be adjusted depending on
287 the dataset (Patel and Divecha, 2010) to exclude differences higher than a defined threshold. Figure 5 illustrates the stepwise
288 removal process of erroneous data. Panels a) and b) show the temperature and RH, respectively, with large negative values
289 (-9999), which correspond to sensor or reading issues during sampling. After removal of these data points, the vertical structure
290 of the temperature and RH profiles emerges, but a few remaining outliers can be easily identified (Fig. 5c and d). These outliers
291 are removed with the despiking function (Fig. 5e and f).

292 Once the temperature, pressure, and relative humidity data are processed, the flight altitude H can be calculated based on:

$$293 \quad H = -\frac{P_0}{\rho_0 g} \times \log\left(\frac{P}{P_0}\right) \quad \text{Eq. 1}$$

294 P_0 is the ground pressure, P is the observed pressure, g is the standard acceleration of gravity $=9.807 \text{ m s}^{-2}$, and ρ_0 is the moist
295 air density (WMO, 2025). An important point to consider is the evolution of ground pressure during a flight. If ground
296 measurements of pressure are available, they are directly implemented in Eq.1 as P_0 . If no ground pressure measurements were
297 available for the duration of the flight, the average pressures before takeoff and after landing are considered as the ground
298 reference, and a linear detrending function was applied for the EERL Helikite altitude to match before and after the flight on
299 the ground.

300
301 **Wind speed and direction.** For the campaigns listed in Table 1, the SmartTether was deployed roughly 2 m below the
302 MoMuCAMS and attached directly to the tether (Fig. 1). It provides direct measurements of wind speed and wind direction in
303 the Earth's coordinate system. Outliers are identified and removed as for e.g., temperature data as described above.
304 More recently, a sonic anemometer was installed on MoMuCAMS and provided additional wind measurement. While it is not
305 yet reported in any of the datasets described here, we provide details on the data processing for the future. The sonic
306 anemometer provides wind measurements in the Helikite coordinate system, i.e., facing the wind direction. To retrieve the
307 meteorological wind in the North-East coordinate system, positions and attitude angles of the MoMuCAMS were recorded by
308 a VectorNav VN-200 Inertial Navigation System. The equations from Lenschow and Spyers-Duran (1989) were applied to get
309 the wind speed and direction in the Earth's coordinate system.

310 **Aerosol number size distribution from the mSEMS.** Data from the mSEMS are reported as number size distribution
311 ($dN/d\log D_p$). From there the integrated particle number concentration is calculated and provided as total particle number
312 concentration (lower and upper diameters can vary between campaigns). Data showing signs of instrument malfunction are
313 identified. These include indications of i) overheating of the instrument (too high concentration for the small diameter bins
314 only), ii) flooding of the optical chamber with butanol due to strong motions of MoMuCAMS (too large concentration in all



315 size bins), or iii) low butanol levels in the condensation chamber (concentrations dropping towards zero). The corresponding
316 data are excluded from the datasets. Potential size-shift and counting efficiency corrections, based on calibration data
317 associated with a specific campaign, can also be carried out at this step if necessary.

318 **Total particle number concentration from particle counters.** For the aMCPC the same checks as for the mSEMS are carried
319 out. Mostly it is part of the mSEMS set up. For the CPC3007 and the Partector 2, outliers corresponding to spurious values
320 inconsistent with the adjacent data, such as unrealistically high concentrations, are identified and removed through visual
321 inspection by the data processor (similarly to the example in Fig. 5). Counting efficiency corrections are carried out at this step
322 if necessary and possible.

323 **Aerosol number size distribution from the POPS.** L0 POPS data is in units of particle counts per second and per bin. From
324 there, the concentration in cm^{-3} is obtained by dividing the particle counts by the mean flow rate over the duration of the flight.
325 Bins 1 to 3 corresponding to mean diameters of 149, 162, and 178 nm are excluded due to overcounting. A detailed explanation
326 can be found in Pohorsky et al. (2024). The total particle number concentration for aerosol diameters between 186 nm and
327 3370 nm is determined as the sum of the remaining bins. The aerosol number size distribution in $dN/d\log D_p$ is derived by
328 dividing the bin concentrations by $d\log D_p$ (with D_p as the midpoint diameter of each bin). Data showing signs of instrument
329 malfunction are identified and excluded. These include indications of i) temporary decrease in the pump flow (concentrations
330 dropping towards zero) or ii) spurious values inconsistent with the adjacent measurements (unrealistically high or low
331 concentrations in all size bins). Potential diameter size-shift and counting efficiency corrections, based on calibration data
332 associated with a specific campaign, can also be carried out at this step if necessary.

333 **Cloud droplet number size distribution from the miniCDA.** The cloud droplet number concentration is obtained by dividing
334 the bin counts that are provided at L0 by the mean flow rate over the duration of a flight. Bins 1 to 71, corresponding to
335 midpoint diameters between 0.24 and 0.66 μm , are excluded as they fall below the instrument's detection limit. The total
336 droplet concentration is then obtained by summing the concentrations from bins 72 to 256. Similarly to aerosol concentrations,
337 the counts are converted into a $dN/d\log D_p$ size distribution. Counting efficiency and size corrections could be implemented at
338 this step, if intercomparison data with standard instruments are available.

339 **CO₂.** Raw measurements are directly corrected for temperature variations with an internal temperature sensor but need to be
340 corrected for pressure variations. The conversion to standard pressure is described at the end of this section. After this
341 correction, the mixing ratio is further converted to a dry mole fraction, using the ambient relative humidity measurements. A
342 calibration factor also needs to be applied. The calibration factor is obtained from lab experiments with specific CO₂ mixing
343 ratios using a linear regression between the observed and calibration gas CO₂ mixing ratios. Finally, data are visually inspected
344 for outliers (i.e., unrealistic values), which are flagged and removed.



345 **CO.** The CO data at L0 are provided as dry mixing ratios in standard temperature and pressure (STP). They are corrected by
346 removing the instrument's measured baseline. The baseline is obtained through operating the CO scrubber 30 min before and
347 after the flight. A linear interpolation of this baseline is applied to account for changes between the beginning and the end of
348 each flight. It is then subtracted from the L0 measurements of CO. Data processing includes a visual quality inspection and
349 removal of outliers similarly to the other data products.

350 **O₃.** L0 O₃ measurements are reported in STP and dry mixing ratios. Using an ozone scrubber, a zero calibration is carried out
351 in each campaign and the dry mixing ratios may be corrected for an offset. The resulting ozone data are visually inspected for
352 outliers that are removed like for the other data products.

353 **Aerosol absorption properties from the STAP and microAeth.** For the STAP measurements, a 2-min moving average has
354 already been applied during the pre-processing step described in Sect. 4.1.1. The absorption coefficients of L0 are inspected
355 for outliers, such as spurious values inconsistent with the adjacent data, which are removed. In certain situations, when
356 coefficient values are low and a rapid change of ambient relative humidity occurs, data become very noisy or can even display
357 negative values. These data are considered faulty and are removed.

358 The microAeth provides eBC concentration data. Values below detection limit and positive outliers are removed. In addition,
359 absorption coefficients at each wavelength can be derived from the eBC concentrations using the wavelength-dependent mass
360 absorption cross sections (MACs) provided by AethLabs (7.78, 10.84, 13.10, 14.67, and 18.52 m² g⁻¹ at 880, 625, 528, 470
361 and 375 nm, respectively). More details of the instrument and calculation approach are provided in Alden et al. (2025b).

362 **Standard Temperature and Pressure correction.** After the above processing steps, all data, which were not reported in STP,
363 are converted to standard conditions. The following equation is used:

364
$$x_{STP} = x \times \frac{T}{T_0} \times \frac{P_0}{P} \quad \text{Eq. 2}$$

365 With x the measurements to correct, and x_{STP} is the STP corrected values. T and P are the temperatures in K and the pressures
366 in hPa observed for every x during the flight. T_0 is 273.15 K and P_0 equals 1013.25 hPa.

367 **4.1.4 Level 1.5 data processing**

368 At this stage, only the scientifically relevant variables are retained for the final dataset (i.e., housekeeping variables e.g., internal
369 temperature, flow rate, or laser power variables are removed). The column names are standardized to ensure consistency across
370 different campaigns. Data recorded before takeoff and after landing are also excluded, keeping only the measurements relevant
371 to the flight. In addition, three flags are added to the datasets to facilitate the scientific analysis of the EERL Helikite
372 measurements, and information on the flight number and campaign name is included for reference.



373 The “flag_pollution” identifies periods when the data are affected by local pollution, e.g., fossil fuel emissions from a ship
374 stack during ARTofMELT or local activities at the observatories during CLAVIER and ORACLES. When pollution is part of
375 the measured environment and aligned with the campaign objectives, it is not flagged, as it represents a relevant component
376 of the observations as e.g., in ALPACA. For ARTofMELT, pollution was detected near the stack height by the POPS, the
377 STAP, and the mSEMS, which showed a pronounced Aitken mode. During CLAVIER, local pollution was identified using
378 the POPS and mSEMS, again showing a distinct Aitken mode. For ORACLES, local pollution was visible in the CPC3007
379 data and confirmed using the flight notes. In contrast, during ALPACA and CHOPIN, local pollution was part of the study
380 objectives, and no specific pollution flag was applied in that dataset. When “flag_pollution” is set to 1, pollution is present.

381 The second flag, “flag_hovering” is applied when the EERL Helikite is stationary, i.e., remaining at constant altitude (within
382 ± 10 -20 m, depending on the flight) for a period exceeding 2 minutes. If “flag_hovering” is set to 1, hovering occurs. On the
383 one hand, hovering information is important when performing data averaging, temporally or spatially, because the number of
384 data points per altitude is strongly impacted. On the other hand, hovering is necessary for certain types of observations, e.g.,
385 for filter sampling, to increase the counting statistics from low time resolution instruments such as the mSEMS, and if a
386 scientifically interesting feature is observed (e.g., cloud, pollution plume) or for technical reasons. To accommodate the coarser
387 time resolution for particle number size distributions from the mSEMS, flight patterns typically include a step-wise ascent or
388 descent to improve counting statistics at different altitudes. This flag can hence be used to more easily select data to refine the
389 analysis of data with a coarser resolution for instance.

390 The third flag, “flag_cloud” informs about cloud conditions. “flag_cloud” is set to 1, when measurements are conducted in
391 cloud or fog and 0 otherwise. For example, in the ARTofMELT campaign, clouds were identified based on aerosol data, high
392 relative humidity, videos and flight notes, as well as ceilometer and cloud radar data providing cloud base height and
393 reflectivity information. During CLAVIER, cloud periods were flagged using aerosol data and RH. For ORACLES, clouds
394 were primarily identified from the miniCDA data and when unavailable, aerosol data and RH were used together with the
395 camera footage. In CHOPIN, the flag was based on flight notes, camera footage, and miniCDA data, while for PaCE, cloud
396 presence was determined from remote sensing and RH data. Note that defining the presence of a cloud is inherently difficult
397 due to rapidly changing conditions, patchy clouds or multiplayer clouds. In addition, the same measurements are not always
398 available. The cloud flag should therefore be considered with caution.

399 At the end of Level 1.5, a 1 s resolution file is produced for each flight.

400



401 **4.1.5 Level 2 data processing**

402 Data are averaged to 10 seconds. This averaging period is selected to keep the high-resolution information, while removing
 403 noisy signals resulting from low concentrations, the platform motions or the sensitivity of the instruments. Once all flights
 404 have been processed and averaged to 10 second resolution, the individual flight files are merged into a single final campaign
 405 dataset.

406 **Table 3: Variables provided in the data products and key processing steps at all levels**

Variable name (column header in data product)	Unit	Raw data and Level 0 comment	Level 1 comment	Level 2 comment
Altitude	m	n.a.	Calculated using barometric equation (Eq. 2) 1 s resolution Quality control (QC)	10 s averaged
Latitude (Lat)	deg	Time synchronization with cross correlation (Eq. 1) Then all variables merged directly into L0 file with 1 s resolution	QC	10 s averaged
Longitude (Long)	deg			
Pressure (P)	hPa			
Temperature (Temperature)	°C		QC Average from 2 sensors for each variable	10 s averaged
Relative humidity (RH)	%		QC	10 s averaged
Wind speed (WindSpeed)	m s ⁻¹			
Wind direction (WindDir)	deg		corrected + STP + QC	10 s averaged
CO ₂ mixing ratio (CO ₂)	ppm			
Total particle number concentration $N_{186-3370}$ (POPS_total_N)	cm ⁻³		Apply correction factors + QC + STP	10 s averaged
POPS bin concentration (POPS_bX)	cm ⁻³ (dN/dlogDp)			
Total droplet number concentration $N_{0.66-43}$ (mCDA_total_N)	cm ⁻³		QC + STP	10 s averaged
miniCDA bin concentration (mCDA_dataBX)	cm ⁻³ (dN/dlogDp)			
Filter sampling (Filter_position)	-		n.a.	10 s averaged
Filter flow (Filter_flow)	lpm			



Total particle number concentration $N_{10-1000}$ (CPC_total_N)	cm^{-3}	No pressure reading. Data merging based on manual time synchronization (pre-flight clock synchronization) into L0 file with 1 s resolution	Apply correction factors + QC + STP	10 s averaged
Total particle number concentration N_{8-250} (mSEMS_total_N)	cm^{-3}	Time synchronization with cross correlation (Eq. 1) at raw data level	Apply correction factors + QC + STP	3 min resolution
mSEMS bin concentration (mSEMS_Bin_ConcX)	cm^{-3} (dN/dlogDp)	Treatment of raw data with manufacturer software Then variables merged directly into L0 file with x min resolution		
Aerosol light absorption coeff. 450 nm (STAP_sigmasb_smth)	Mm^{-1}	Time synchronization with cross correlation (Eq. 1) at raw data level	QC + STP	10 s averaged
Aerosol light absorption coeff. 525 nm (STAP_sigmag_smth)	Mm^{-1}	Treatment of raw data with manufacturer software		
Aerosol light absorption coeff. 624 nm (STAP_sigmar_smth)	Mm^{-1}	Then variables merged directly into L0 file with 1 s resolution		
MicroAeth eBC mass value for 375, 470, 528, 625, 880 nm (MA_eBC_X)	ng/m^3	No pressure reading. Data merging based on manual time synchronization (pre-flight clock synchronization) into L0 file with 1 min resolution	QC + STP	10 s averaged
MicroAeth light absorption coeff. 375, 470, 528, 625, 880 nm (MA_Abs_Coeff_X)	Mm^{-1}			
O ₃ mixing ratio (O3)	ppb	Time synchronization with cross correlation (Eq. 1) Then variable merged directly into L0 file with 2 s resolution	Apply correction factors + QC + STP (automatic)	10 s averaged
CO mixing ratio (CO)	ppb	No pressure reading. Data merging based on manual time synchronization (pre-flight clock synchronization) into L0 file with 1 s resolution	Apply correction factors + QC + STP (automatic)	10 s averaged

407

408

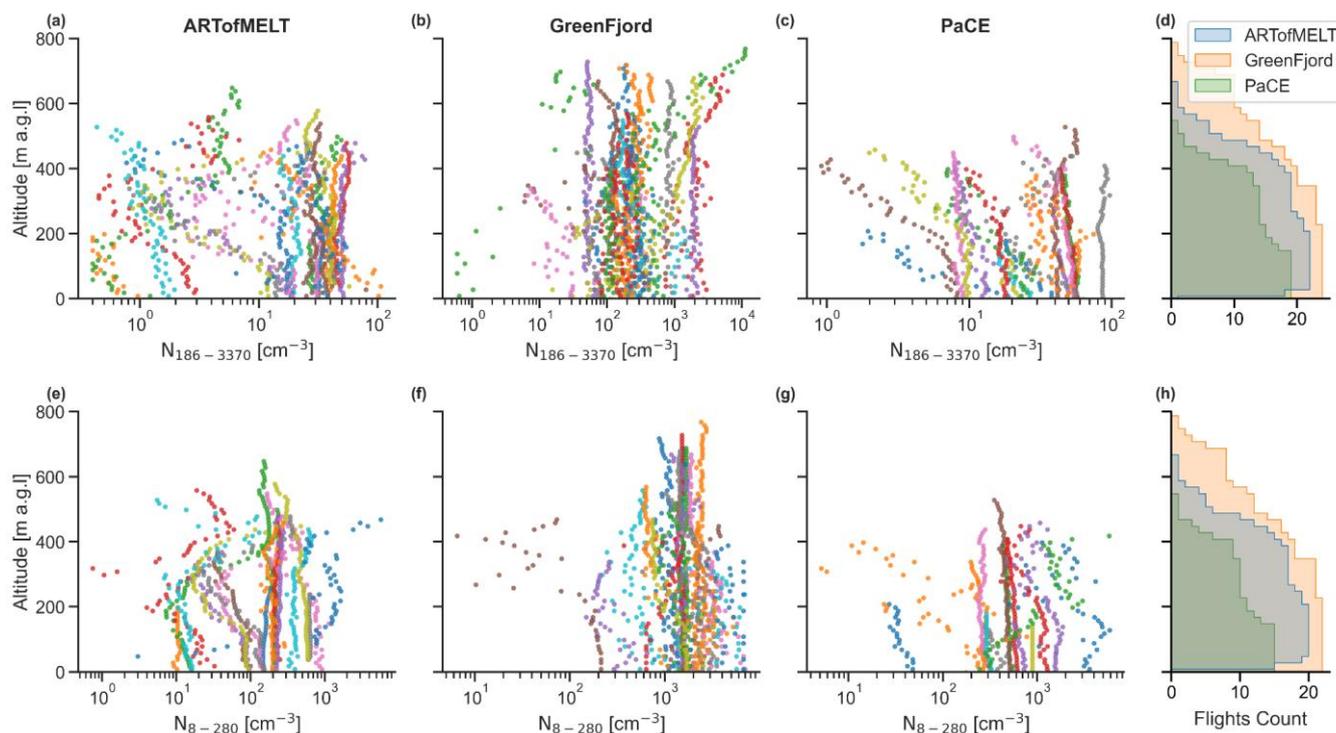


409 **4.2 Example analyses**

410 The purpose of this section is to provide examples of how data might be visualized and analyzed by the community. We do
411 not provide extensive scientific interpretation, which can be found in campaign specific publications.

412 **4.2.1 Vertical profiles of particle number concentrations**

413 We present example vertical profiles of particle number concentrations between 186 and 3370 nm, measured with a POPS,
414 and between 8 and 280 nm, measured with an mSEMS, from three different campaigns (Fig. 6). During the ARTofMELT
415 expedition, 23 flights were carried out over the Arctic sea ice during the melt onset in May-June 2023. In a southern
416 Greenlandic fjord system, during GreenFjord, 24 flights were performed between June and July 2023 (Alden et al., 2025).
417 Finally, 19 flights were conducted during the Pallas Cloud Experiment (PaCE) in September-October 2022 in the boreal forest
418 of Finland. Focusing on the larger accumulation and coarse mode, number concentrations vary between 10 and 100 cm⁻³ over
419 the Arctic sea ice (Fig. 6a) and in the boreal forest (Fig. 6c), and occasionally drop below 10 cm⁻³ in the central Arctic. The
420 individual profiles show situations with vertically homogeneously distributed aerosol concentrations and also stratified
421 situations, where concentrations either decrease or increase with altitude. Observations in southern Greenland in summer (Fig.
422 6b) are, however, different. They are generally higher compared to the other two campaigns, ranging from 100 to 3000 cm⁻³,
423 most likely because of the proximity to the open ocean and sea spray generation. On specific occasions, concentrations
424 increased with altitudes owing to three long-range transport events of biomass burning emissions from Canada at elevated
425 altitudes (Alden et al., 2025b) as can be seen from the markers of individual flights. The fine mode particle vertical profiles
426 (Fig. 6e-g) generally resemble the coarse mode profiles at each site. Surface concentrations are however higher and range from
427 roughly 10 to almost 1000 cm⁻³ in the central Arctic (ARTofMELT). Over the boreal forest (PaCE), concentrations are similar
428 but occasionally exceed 1000 cm⁻³, while in southern Greenland they often exceed 1000 cm⁻³.



429

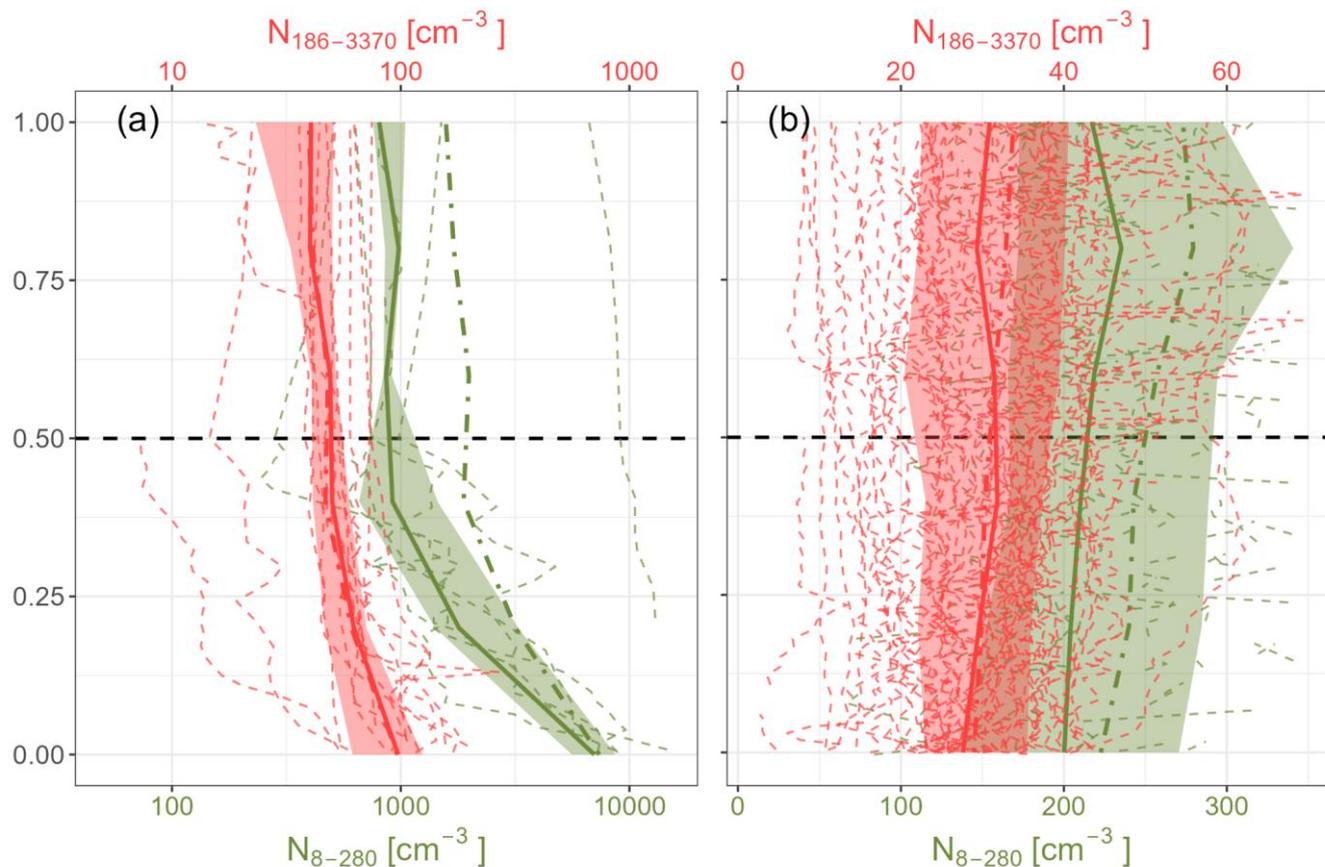
430 **Figure 6:** Vertical profiles of particle number concentrations from the ARTofMELT (a, e), GreenFjord (b, f) and PaCE (c, g)
431 campaigns. The top row shows particle number concentrations between 186 and 3370 nm measured by a POPS. The bottom row
432 displays the number concentration of particles between 8 and 280 nm measured by an mSEMS. The colored dots represent individual
433 flights in each campaign and data are averaged per 10 m altitude. Panels d and h provide the number of flights for which data were
434 recorded at specific altitudes.

435 4.2.2 Particle number concentrations relative to inversion heights

436 The recorded temperature profiles allow for the determination of inversions and investigation on how aerosols are transported
437 across inversions. We provide examples on the role of surface-based inversions in the presence (ALPACA) and absence
438 (CLAVIER) of surface aerosol sources (Fig. 7). In this example, surface-based inversions were determined by identifying the
439 height where $dT/dz = 0$ as in Kahl (1990). The height of the top of the surface-based inversion was defined as 0.5 and data
440 from each flight at altitudes below were normalized between 0 and 0.5, and between 0.5 and 1 for observations above the
441 inversion top. In the urban winter environment of Fairbanks, Alaska (panel a, ALPACA), the aerosol number concentration
442 below the surface-based inversion top (dashed horizontal line) is significantly higher than above. Below the inversion the
443 vertical profiles exhibit a decrease from the surface upward which is expected since the surface emissions can only slowly be
444 mixed upward. Above the inversion, aerosol number concentrations reach their minimum values reflecting a type of
445 background concentration with little influence from surface emissions (Pohorsky et al., 2025b). The observations during
446 CLAVIER in northern Greenland are different. Below the inversion the particle number concentrations are lower due to the
447 absence of surface sources, while above the inversion, concentrations tend to be higher due to long-range transport of mid-



448 latitude pollution during the Arctic haze season. This feature is however less evident since long-range transport events are
449 more sporadic as they depend on large-scale circulation and synoptic situations. Fig. 7b shows therefore more variability
450 between each individual profile.



451

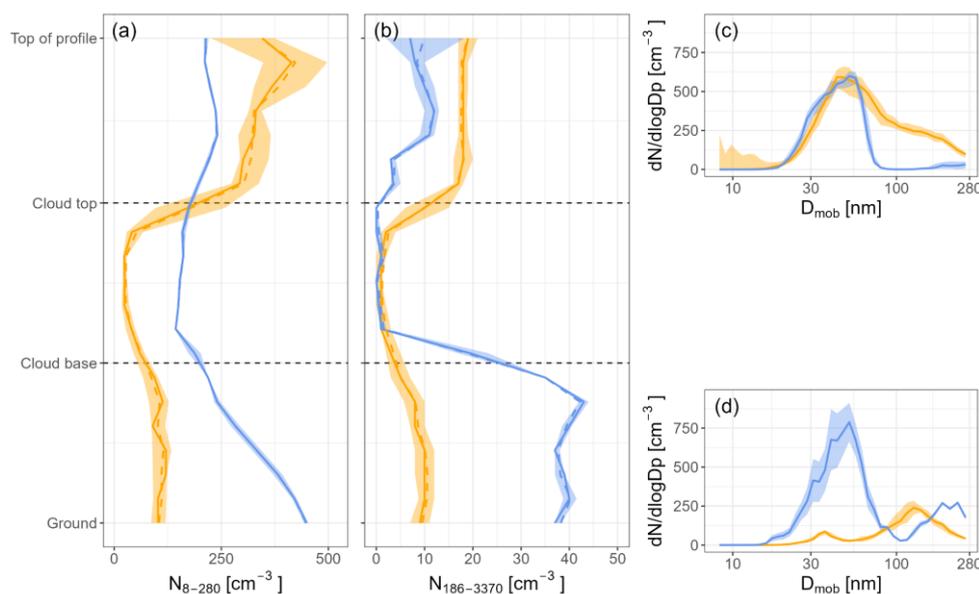
452 **Figure 7: Vertical profiles of particle number concentrations normalized to the surface-based inversion height at 0.5 for a) the urban**
453 **winter atmosphere of Fairbanks (ALPACA, n=10), and b) a remote high Arctic site during spring (CLAVIER, n=51). For both**
454 **panels, the full line represents the median of all individual profiles, the shading indicates the interquartile range and the dashed lines**
455 **represent the mean.**

456 4.2.3 Aerosol properties below, in and above clouds

457 In Fig. 8 we illustrate one possible approach to using the cloud flag described in Sect. 4.1.4 by normalizing the vertical profiles
458 of particle number concentrations to the cloud base height and cloud top. During the ARTofMELT expedition there were two
459 distinct types of aerosol vertical profiles when clouds were present and the EERL Helikite was also able to conduct
460 measurements above the cloud. One type (orange profile, flight on June 7th, 2023) featured very low particle number
461 concentrations below cloud, i.e., around a 100 cm^{-3} for the fine mode, and around 10 cm^{-3} for the coarse mode, while above



462 cloud, the concentrations increased by a factor of 3 and 1.5, respectively. The higher particle number concentrations aloft were
463 typically observed in conjunction with Aitken mode particles (Fig. 8c). Whereas near the surface (Fig. 8d), particle number
464 size distributions with a dominant accumulation mode and Hoppel minimum occurred, typical for cloud processed particles.
465 Note that the near zero concentrations inside the cloud are related to the inlet type, which is near-interstitial and does not
466 sample droplets unless they are very small. The other cloud vertical profile type (blue profile, flight on June 10th, 2023) features
467 higher concentrations of both particle populations below the cloud with $> 300 \text{ cm}^{-3}$ for N_{8-280} and around 40 cm^{-3} for $N_{186-3370}$
468 compared to above the cloud. Concentrations decrease steadily with height for the fine mode, but remain constant until cloud
469 base for the coarser particles. Above the cloud, concentrations for both modes are lower than below the cloud. Fig. 8c) and d)
470 indicate an Aitken mode-dominated size distribution above and below the cloud, with an additional accumulation mode below
471 the cloud, indicative of cloud-processed aerosols.



472

473 **Figure 8: Comparison of two vertical profiles normalized to cloud base height during the ARTofMELT campaign on June 7th, 2023**
474 **(orange) and June 10th, 2023 (blue). a) Total aerosol concentration between 8 and 280 nm measured with the mSEMS. b) Total**
475 **aerosol concentration between 186 and 3370 nm measured with the POPS. Normalization means that the vertical extent of each**
476 **observation category (i.e., below, within and above cloud) is represented as one third of the profile. c) Particle number size**
477 **distributions above the cloud. d) Particle number size distributions below the cloud. For all panels, the full line represents the median**
478 **of all individual profiles, the shading indicates the interquartile range and the dashed lines represent the mean.**

479 5 User guidance

480 Here we provide considerations for users of these, and future, EERL Helikite datasets. Tethered-balloon data offer the
481 possibility to visualize the vertical distribution of the different variables as illustrated above. Fast profiles provide a quasi-
482 snapshot of the state of the vertical column. However, flights typically include “hovering” periods where the EERL Helikite



483 is maintained at a constant altitude. Under these conditions, more data points are collected at a given altitude but the
484 measurements are subject to local and temporal variability (e.g., from eddies or horizontal advection). Additionally, the
485 advection of sporadic features such as a small cumulus cloud or a pollution plume can sometimes be captured very briefly
486 while profiling. Because of the very sporadic nature of these features, their full vertical extent may not always be fully captured.
487 For an analysis of the vertical structure of the atmosphere, it is therefore recommended to first observe the flight pattern (i.e.,
488 altitude versus time) to identify fast profiles and hovering periods to select flight segments appropriate for the intended analysis,
489 and to spatially average the data (e.g., in vertical layers of 10 m as in Fig. 4, or levels corresponding to a model output).
490 Hovering periods, defined by the “flag_hovering” in the datasets, are useful to obtain better counting statistics from instruments
491 with coarser time resolution (e.g., mSEMS) to evaluate variations between different atmospheric layers or to compare the
492 temporal evolution of a specific variable at a specific altitude. Information on local pollution that might affect the aerosol
493 measurements is also included in the datasets with the “flag_pollution” and should be considered and removed if not desired.
494 In addition, it is recommended to make use of ancillary data such as from ground-based remote sensing that is frequently
495 available (e.g., ceilometer observations during Greenland or CloudNet during ORACLES).

496 The aerosol inlet of MoMuCAMS can be considered as interstitial inlet due to its horizontal orientation and generally low flow
497 rate between 0.54 and 3 lpm (compared to standard ground-based monitoring stations) with a 7.5-mm inner diameter. In clouds,
498 water droplets are not captured by the inlet. Consequently, measurements usually show an apparent depletion of aerosols inside
499 clouds, and only interstitial particles remain visible. It is possible that very small droplets are still captured by the inlet and
500 measured. However, since the inlet is heated, those might evaporate in the inlet. The transmission of the very small droplets
501 depends hence on the flow rate of the system and can therefore change from one flight to another.

502 Relative humidity measurements typically show higher uncertainties at elevated RH (> 90 %). We also observe that RH does
503 not always reach 100 % in clouds. Both temperature and RH measurements are subject to ground effects (e.g., from excessive
504 heating of the surface from solar radiation). Despite careful data cleaning, data points near the ground should always be treated
505 with care, especially if a spatial data averaging is applied.

506 **6 User resources and code availability**

507 All data were processed in Python with the EASE code. The code package is available on Github: [https://github.com/EERL-](https://github.com/EERL-EPFL/helikite-data-processing)
508 [EPFL/helikite-data-processing](https://github.com/EERL-EPFL/helikite-data-processing).

509 A Python Jupyter Notebook script is available:
510 https://github.com/EERL-EPFL/helikite-data-processing/blob/main/notebooks/ESSD_Plot_Helikite_level2.ipynb to support
511 the initial handling of the datasets and to produce quick-looks of the EERL Helikite flights. The software provides different
512 visualization functions for each of the datasets:



- 513 a) Vertical profiles for each flight for each variable. Each flight is subdivided into ascent and descent based on the
514 maximum altitude reached during the flight. Flags are included as shaded areas.
- 515 b) Time series for each flight for each variable. A dotted line marks the transition between ascent and descent. Flags
516 for cloud presence, pollution, and stationary flight, are displayed as well.
- 517 c) Average profile: for each variable, an average profile is obtained from all the profiles of the campaign.
- 518 d) Overview of the campaign: all the profiles of the campaign are plotted and color coded based on the flight
519 number.

520 See Appendix A for examples.

521 **7 Summary**

522 In this data descriptor we provide information on the processing from raw to level 2 data, following recommendations from
523 EBAS for data levels for 15 different instruments typically deployed on the EERL Helikite. The specifically developed Python
524 code EASE is used for this purpose and available on Github. We further provide a quick look plotting tool that users can apply
525 to explore the EERL Helikite data. To guide the exploration of the data, examples are given on how data can be visualized and
526 analyzed for specific purposes such as aerosol dispersion across temperature inversions or aerosol-cloud interactions. The data
527 can be used e.g., for model and remote sensing validation or as contextual information of vertical aerosol profiles at the given
528 locations and seasons for future measurements there that do not dispose of vertical observational facilities. For most of the
529 campaigns mentioned in this data descriptor, specific manuscripts have been published or are in preparation. Nevertheless, the
530 datasets are rich in information and it requires the wider research community to fully take advantage of the vertical aerosol
531 information in the lower atmospheric boundary layer in polar and alpine regions. More data will be added from future
532 campaigns and updates on the availability of new data sets can be obtained here: [https://www.epfl.ch/labs/eerl/eerl-home-
533 page/research/field-work-overview/](https://www.epfl.ch/labs/eerl/eerl-home-page/research/field-work-overview/).

534

535



536 **Data availability**

537 All data described above are available here:

- 538 1. ARTofMELT: Pohorsky et al. (2025), <https://bolin.su.se/data/oden-artofmelt-2023-aerosols-vertical>
- 539 2. ALPACA: Pohorsky et al. (2025a), <https://arcticdata.io/catalog/view/doi%3A10.18739%2FA2MS3K363>
- 540 3. Brigerbad: Schmale et al. (2026), <https://zenodo.org/records/18327462>
- 541 4. CHOPIN: Alden et al. (2026), will be available on the ACTRIS homeless data portal: <https://dc.actris.nilu.no/search>
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548 **Author contributions**

549 JS, RP, YT, ML, JA, RC wrote the manuscript. RP, JA, RC, YT, ML, OS, ET, AS wrote data processing code. RP, JA, RC,
550 YT, ML, NB, BH, AB, MLH, RP, OS, AT, HH processed and curated data. All authors contributed to creating the final data
551 sets and commented on the manuscript.

552 **Competing Interests**

553 The authors declare that they have no conflict of interest.

554 **Special issue statement**

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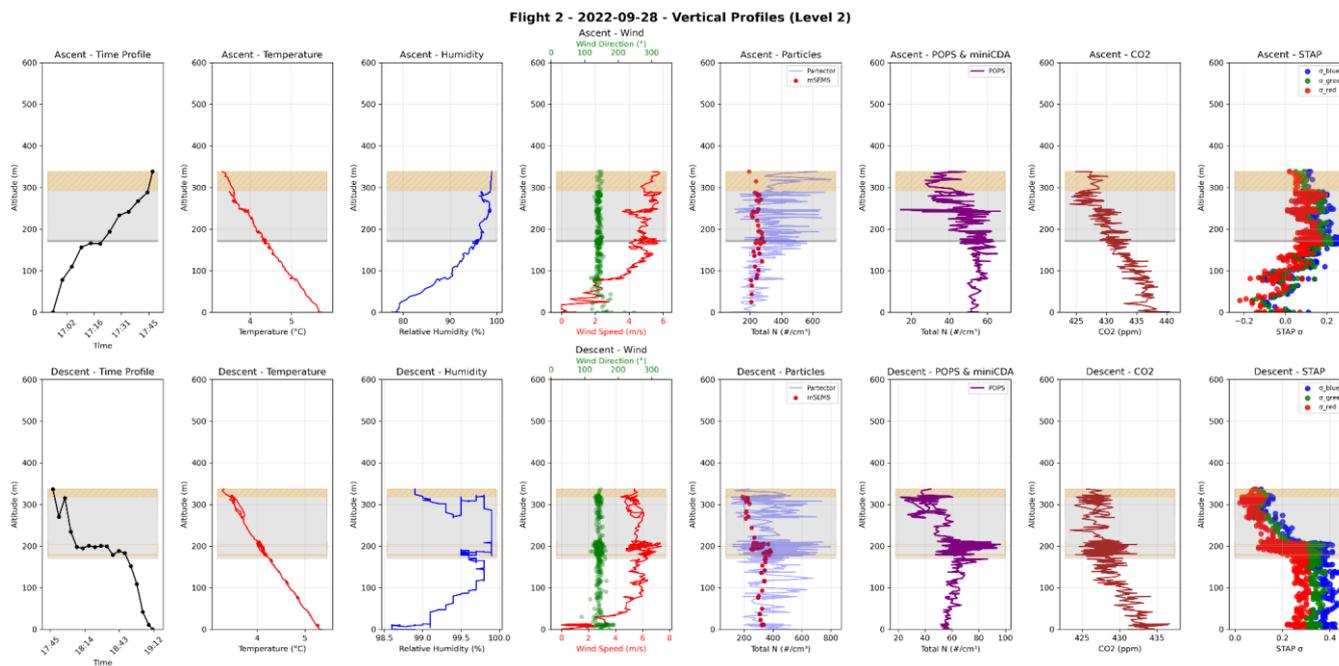
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578 **Appendix A**

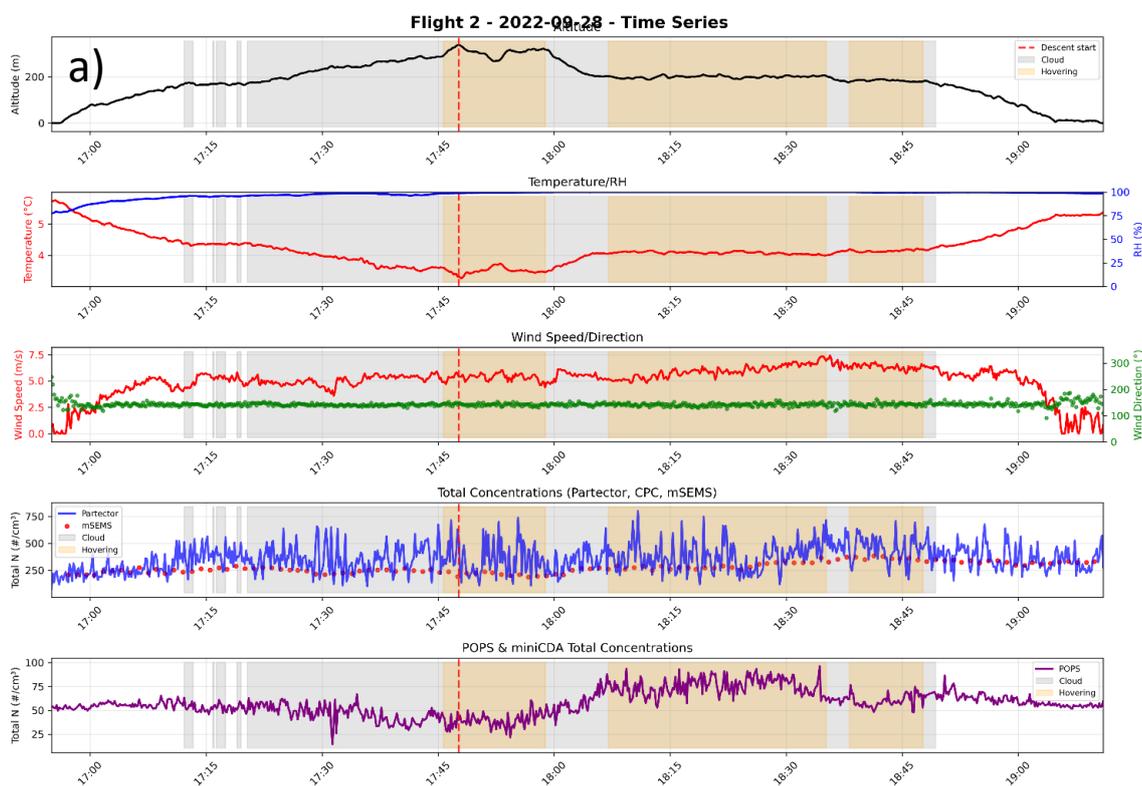


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580 **Figure A1.** Vertical profiles of all variables for one example flight (flight 2 of PaCE). The upper panels display the ascent profile,
 581 and the bottom panels the descent profile. In each panel, the presence of a cloud is indicated by a gray shading based on the cloud
 582 flag. A yellow shading indicates a stationary period lasting more than 2 minutes, based on the hovering flag. The first panel indicates
 583 the altitude of the balloon resampled every 5 minutes to ease the reading. The second and third panels indicate temperature and
 584 relative humidity. The fourth panel combines wind speed (red, bottom x-axis) and the wind direction (green, upper x-axis). The fifth
 585 panel indicates the total concentrations of sub-micron particles, displaying measurements by mSEMS (red), Partector (blue), CPC
 586 (green, not flown). The next panel showcases the number concentrations of larger particles and droplets with measurements from
 587 the POPS (purple) and miniCDA (yellow, not flown). The seventh panel indicates the mixing ratio of CO₂. The last panel indicates
 588 the absorption coefficients of the STAP or microAeth, data are color-coded based on the wavelength.

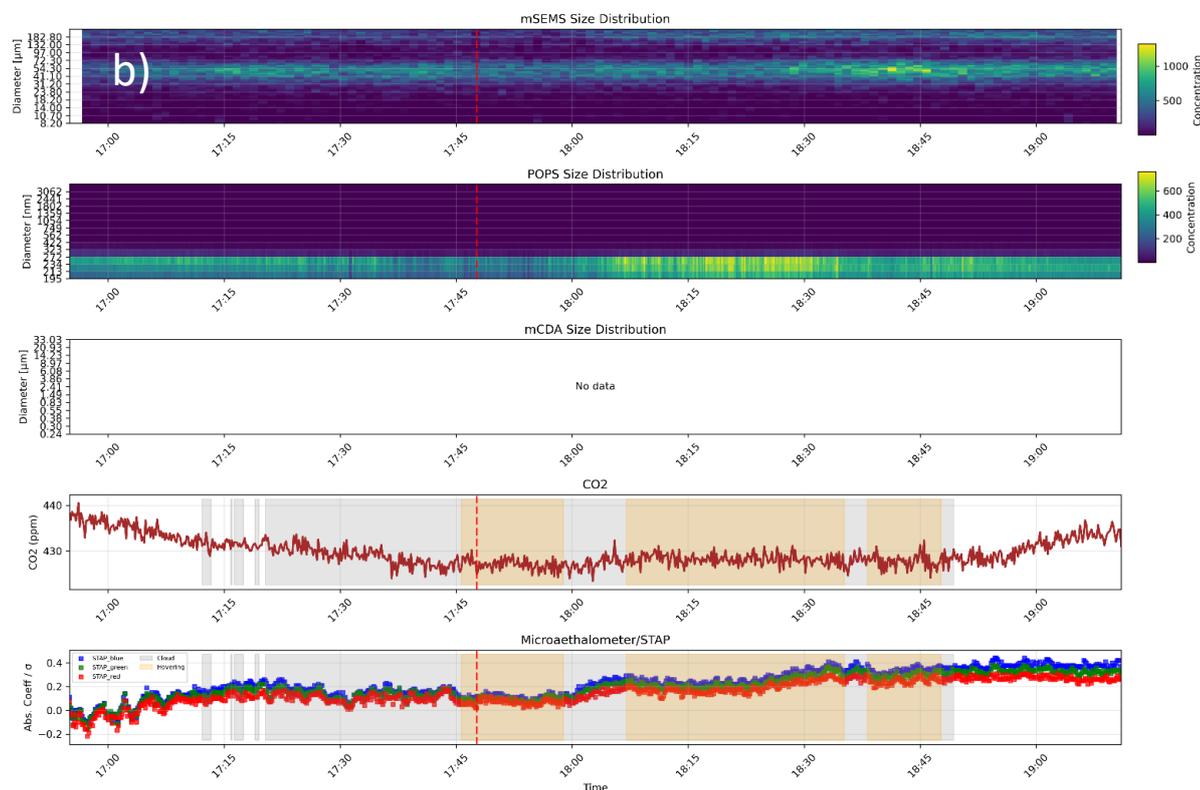


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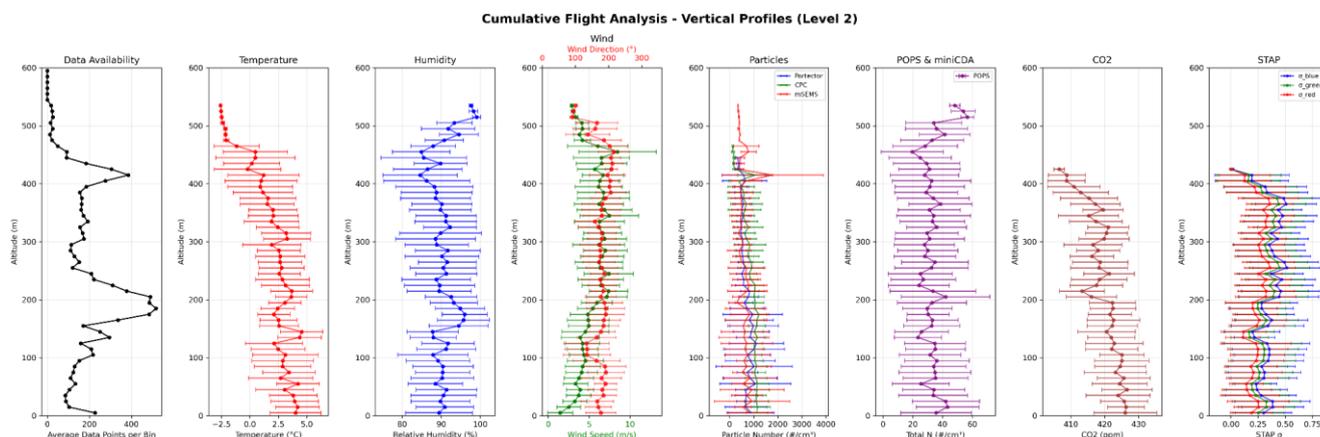


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593 **Figure A2.** Time series from flight 2 of PaCE. The temporal evolution of selected variables is displayed in 10 stacked panels (here
594 separated into panels a and b) following the same time axis. A red dashed line separates the ascent from the descent. Flags for clouds,
595 stationary flight, are displayed as shaded areas. The first panel indicates the altitude of the profile. The second panel combines
596 temperature (red, left y-axis) and relative humidity (blue, right y-axis). Wind speed (red, left y-axis) and wind direction (green, right
597 y-axis) are shown in the third panel. Panel 4 displays the total concentration of sub-micron particles, with data from mSEMS (red),
598 Partector (blue), and CPC (green, not flown). Similarly, the fifth panel features the total concentration of larger particles by the
599 POPS (purple) and droplets by the miniCDA (yellow, not flown). Panels 6 to 8 display heatmaps of the size distributions by mSEMS,
600 POPS, and miniCDA. The left axes indicate the mid-diameter of some of the bins, while individual colorbars are provided on the
601 right side of the panels. Panel 9 indicates the CO₂ concentration. The last panel indicates the absorption coefficient of STAP of
602 microAeth, with color-coding based on the wavelength.



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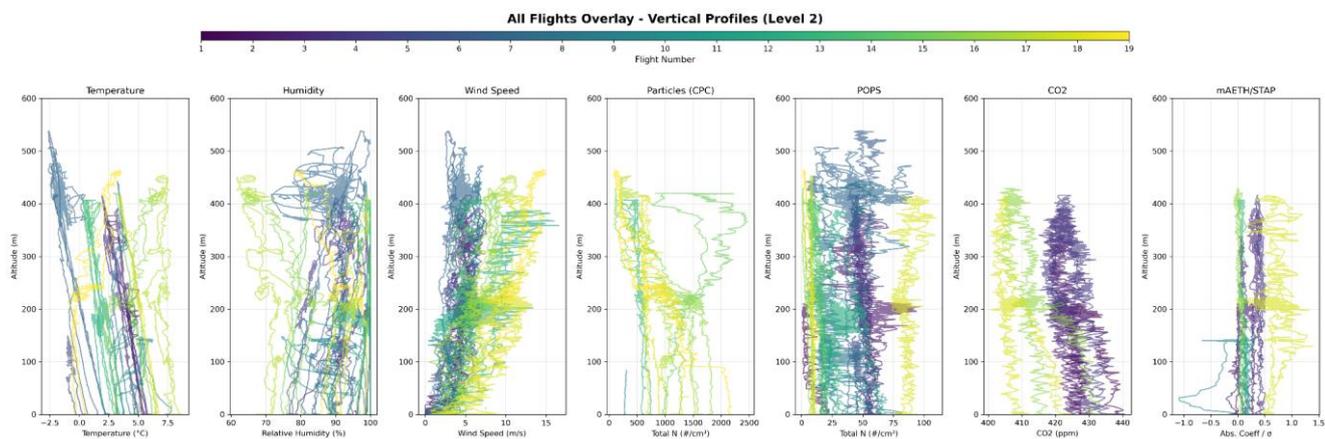
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Figure A3. Vertical profiles of averaging all helikite data from the PaCE campaign. This figure is obtained by combining the ascents and descents of all the flights, binning data every 10 m of altitude. Panel 1 indicates the average number of points per bin. All other panels 2-8 are structured as in figure A1. a solid line indicates the average, horizontal bars indicate the standard deviation.



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Figure A4. Overview of all vertical profiles from the PaCE campaign. The figure is obtained by combining all the vertical profiles, and color coding them based on the flight number. The color variation represents the – non-normalized - temporal evolution of the campaign.



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