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Atmospheric Measurement Techniques Discussions

LOAC: a small aerosol optical counter/sizer for ground-based and balloon measurements of the size distribution and nature of atmospheric particles – Part 1: Principle of measurements and instrument evaluation

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AMTD
8, 1203–1259, 2015

Abstract

The study of aerosols in the troposphere and in the stratosphere is of major importance both for climate and air quality studies. Among the numerous instruments available, aerosol particles counters provide the size distribution in diameter range from few hundreds of nm to few tens of µm. Most of them are very sensitive to the nature of aerosols, and this can result in significant biases in the retrieved size distribution. We describe here a new versatile optical particle/sizer counter (OPC) named LOAC (Light Optical Aerosols Counter), which is light and compact enough to perform measurements not only at the surface but under all kinds of balloons in the troposphere and in the stratosphere. LOAC is an original OPC performing observations at two scattering angles. The first one is around 12°, and is almost insensitive to the nature of the particles; the second one is around 60° and is strongly sensitive to the refractive index of the particles. By combining measurement at the two angles, it is possible to retrieve accurately the size distribution and to estimate the nature of the dominant particles (droplets, carbonaceous, salts and mineral particles) in several size classes. This speciation is based on calibration charts obtained in the laboratory. Several campaigns of cross-comparison of LOAC with other particle counting instruments and remote sensing photometers have been conducted to validate both the size distribution derived by LOAC and the retrieved particle number density. The speciation of the aerosols has been validated in well-defined conditions including urban pollution, desert dust episodes, fog, and cloud. Comparison with reference aerosol mass monitoring instruments also shows that the LOAC measurements can be successfully converted to mass concentrations. All these tests indicate that no bias is present in the LOAC measurements and in the corresponding data processing.

AMTD
8, 1203–1259, 2015

1 Introduction

The importance of measuring the concentration and size distribution of aerosols in the lower atmosphere has been highlighted by various studies. For instance, their presence in ambient air can have direct effects on human health (e.g. Zemp et al., 1999; Brunekreef and Holgate, 2002), and their interaction with solar radiation and clouds are affectingregional and global climate (Ramanathan et al., 2001; Diner et al., 2004; Kanakidou et al., 2005; Quaas et al., 2008). When very high concentrations of ashes after volcanic eruptions are present at cruise altitude, they can affect air traffic (e.g. Chazette et al., 2012). In the middle atmosphere, aerosols play a significant role in stratospheric chemistry through heterogeneous reactions with nitrogen and halogen species (e.g. Hanson et al., 1994, 1996), and they can affect climate through their role in the global radiative balance of the Earth (e.g. Hansen et al., 1992; Ammann et al., 2003). The concentration and size of the particles are highly variable due to the large variety of aerosol sources and properties, both of natural and man-made origin, and because of their relatively short residence time in the atmosphere. To understand and predict aerosol impacts, it is important to develop observation and monitoring systems allowing for their full characterization.

Instruments have been developed for routine measurements or for dedicated campaigns. Observations can be conducted from the ground, from unmanned aerial vehicles (UAV), from aircrafts, from balloons, and from satellites. To retrieve the physical properties of the aerosols, it is necessary to combine the information obtained with different instruments. In situ mass-spectrometers (Murphy et al., 2007) and aerosol collecting instruments (Brownlee, 1985; Blake and Kato, 1995; Allan et al., 2003; Bahreini et al., 2003; Ciucci et al., 2011) provide their composition. Optical instruments performing remote sensing measurements from the ground or from space with photometric, lidar, and extinction techniques (Shaw et al., 1973; Dubovik and King, 2000; Bitar et al., 2010; Winker et al., 2010; Salazar et al., 2013) provide indications on the size distribution and on the nature of the particles, generally assuming a priori hypotheses in

AMTD
8, 1203–1259, 2015

the retrieval process. Complementarily, in situ optical measurements with optical particle counters can provide more accurate information on the size distributions of the particles.

The present study deals with optical aerosol particles counters (OPCs). The corresponding measurement principle relies on the properties of light scattered by particles injected in an optical chamber and crossing a light beam (e.g. Grimm and Eatough, 2009). The measurements are usually conducted at "large" scattering angles, typically around 90° with collecting angle of a few tens of degrees. At such angles, the light scattered is depending both on the size of the particles and on their refractive index. Conventional counters are calibrated using latex and glass beads and are post-calibrated using Mie calculations (Mie, 1908) for liquid aerosols (the refractive index of latex beads and liquid aerosols is well known, assuming no imaginary part of the index i.e. non-absorbing aerosols). Some instruments can be also be post-calibrated for the observation of specific particles, as desert dust or urban pollutants, assuming a given value of their refractive index.

The refractive index dependence can be partially determined by performing measurements at different scattering angles. Since the variation of the scattered intensity with scattering angles is strongly dependent on the refractive index of the particles (Volten et al., 2006; Francis et al., 2011). Thus, performing simultaneous measurements at different angles can provide an indication of the nature of the particles. Such an approach was used by Eidhammer et al. (2008) at angles of 40° and 74° mainly for the identification of mineral particles, and by Gayet et al. (1997) with a ring of detectors covering the whole scattering angle range for the identification of cloud droplets and icy particles.

Another approach was proposed by Renard et al. (2010a); in this case, measurements are conducted at small scattering angles, below 20°, where the light scattered is less sensitive to the refractive index of the particles. In this angular region, the scattered light is dominated by diffraction (which is not sensitive to the refractive index), at least for irregular grains as those found in the atmosphere. Such non-dependence of

AMTD
8, 1203–1259, 2015

the refractive index was confirmed by measurements conducted at a scattering angle around 15° for different types of irregular grains (Lurton et al., 2014). In this case, the light scattered is dependent only on the size of the particles, allowing a better determination of the corresponding size distribution. However, the main problem of measurement at small angles is stray-light contamination. Thus a real-time correction of this signal offset due to the stray-light, which can vary with time, must be developed.

Aerosol particles counters are often used on the ground; some of them are used in the free atmosphere on-board aircraft or large balloons during dedicated campaigns, for example for the studies of desert dust events or volcanic aerosols (Bukowiecki et al., 2011; Jégou et al., 2013; Ryder et al., 2013) or for stratospheric studies (Rosen, 1964; Ovarlez and Ovarlez, 1995; Deshler et al., 2003; Renard et al., 2008, 2010b). We propose here a new optical particle counter concept, called LOAC (Light Optical Aerosols Counter) that is light and compact enough to perform measurements on the ground and under all kinds of balloons in the troposphere and in the stratosphere, including meteorological balloons. LOAC uses a new approach combining measurements at two scattering angles. The first one is around 12°, an angle for which scattering is weakly sensitive to the nature of the aerosols, allowing the retrieval of the particle size distribution. The second one is around 60° where the light scattered is strongly sensitive to the refractive index of the particles, and this can be used to determine their nature.

In this first paper, we will present the principle of measurements and calibration, and cross-comparison exercises with different instruments that detect atmospheric aerosols. In a companion paper, we illustrate first scientific results from airborne observations on-board balloons and unmanned aircraft.

AMTD
8, 1203–1259, 2015

2 Principle of measurements

2.1 Instrument concept

LOAC is a modular instrument, for which some parts can be changed depending on the measurements conditions. For measurements under balloon or on the ground in low wind conditions, the aerosols are collected by a metal profiled inlet designed to optimize the sampling conditions when oriented in the wind direction. The particles are drawn up to the optical chamber through an isostatic tube by a small pump (having a life-time of 3 weeks in continuous operation) working at $\sim 2\,\mathrm{L\,min^{-1}}$. The pump is connected to the exit of the optical chamber by a flexible plastic tube. For measurements in windy and rainy conditions, the inlet can be replaced by a total suspended particulate or TSP inlet rejecting rain droplets and particles greater than 100 µm. For long-duration measurements, the small pump can be replaced by a robust pump; to maintain the aerosol detection efficiency, the pump flux must be in the range 1.3–2.7 $\mathrm{L\,min^{-1}}$.

The sampled air crosses a laser beam of 25 mW working at the wavelength 650 nm. The scattered light is recorded by two photodiodes at scattering angles of respectively ~ 12 and $\sim 60^{\circ}$ (Fig. 1). Instead of using lenses to collect the light, the photons travel directly to the photodiodes through pipes, providing fields of view with a few degrees. The collecting area of the photodiodes is larger than the diameter of the pipes. This system prevents optical misalignment problems in case of vibrations and strong temperature variations like those encountered during atmospheric balloon flights. Such a concept of scattering measurements without collecting lenses was tested and validated by Daugeron et al. (2007).

The electronic sampling is at 40 kHz and the transit time of particles inside the laser beam is of ~ 500 µs. As said before, stray light contamination is high at small scattering angles and needs real-time correction. The stray light correction method presented in Renard et al. (2010a) was applied to the LOAC measurements. The stray light acts as a flux continuum, which can slightly vary over time due to changes in the temperature and pressure conditions and possible dust contamination in the optical chamber. The

AMTD
8, 1203–1259, 2015

light scattered by the particles is superimposed on this continuum, which can be assumed as a continuous base-line over a short time interval. This baseline is determined before and after the intensity pulse produced by the particles that cross the laser beam.

The maximum of the intensity pulse is obtained after subtracting the stray-light contamination. Figure 2 presents an example of real measurements of the time evolution of the flux scattered by a 5 µm particle and by few submicronic particles. The pulse is slightly asymmetric, which corresponds to the deceleration of the particle while crossing the laser beam. Some secondary flux maxima may be present in the pulse and are due to the rotation of particles having an irregular shape. The search for a new intensity peak is inhibited until the flux decreases to a given threshold, represented in Fig. 2 by the red line. This procedure prevents multiple counting of the same particle (of irregular shape) that exhibits secondary flux maxima.

To minimize its weight, the optical chamber is in plastic Delrin®. The weight, including the pump, is of 350 g. The electric consumption is of 340 mA under 8 V (which corresponds to a power of 3 W).

2.2 Calibration

The calibration of an optical counter is not an easy task, especially for the detection of irregular particles (Whitby and Vomela, 1967; Gebhart, 1991; Hering and McMurry, 1991; Belosi et al., 2013). A first presentation of the calibration procedure for measurements at small scattering angles using a LOAC optical chamber can be found in Lurton et al. (2014).

Latex beads, which are perfect transparent spheres, have been used for diameter calibration below 2 µm. In fact, Mie calculations show that the scattered flux encounters strong oscillations linked to small changes both in diameter and in scattering angle. Conventional aerosol counters use large field of view, typically a few tens of degrees, to average these oscillations. On the opposite, the LOAC 12 and 60° measurement channels have a field of view only of few degrees and use no lens. The detected flux is then very sensitive to the position of the individual bead inside the laser beam, and thus

AMTD
8, 1203–1259, 2015

to its scattering angle. Fortunately, real atmospheric particles are not perfect spheres and will not produce Mie oscillations. Also, such grains rotate when crossing the laser beam, producing scattered flux variations independent of the grain position inside the beam. Thus, we considered here the highest flux scattered by the beads.

For the calibration in the 5–45 µm size range, different natures of irregular grains have been used: glass beads (quite irregular), carbon particles, dust sand of various types, ashes and salts. The size selection was obtained using sieves. For diameters at ∼ 90 µm, calibrated silicon carbide grains were used, the size being characterised by the provider. As expected, no significant dependence with the nature of these particles was found for the variation of the (peak) scattered flux with their diameter, as shown in Fig. 8 of Renard et al. (2010a) and in Fig. 5 of Lurton et al. (2014).

Figure 3 presents the calibration curve; where the scattered flux is given in mV, which corresponds to the photodiode output voltage (updated from Lurton et al., 2014). The diameter presented here corresponds to an equivalent (or optical) diameter, which can differ significantly from the aerodynamic diameter or from the electric mobility diameter used by non-optical instruments for ambient air measurements. The electronic noise is taken into account, and acts as an offset in the output voltage. The calibration captures well the large-amplitude Mie oscillations calculated by integrating the scattered fluxes over the whole LOAC field of view. In particular, the amplitude of the $\sim 2\lambda_0$, and $\sim 3\lambda_0$ enhancements (λ_0 being the laser diode wavelength) is well reproduced. For the larger sizes, the evolution of the scattered flux with size is lower than the one expected from the Mie calculation. This is due to both the small aperture of the field of view and to the roughness of particles, the recorded flux being dominated by diffraction (Lurton et al., 2014). Taking into account this advantage, the LOAC detection size range is between 0.2 and ∼ 100 µm; the upper limit can be lower, however, depending on the sampling collection cut-off of the inlet.

Particles found in ambient air are not perfectly spherical and have some irregularity on their surface, even for the sub-micrometre (sub-µm) sizes (e.g. Xiong and Friedlander, 2001; McDonald and Biswas, 2004). Thus the large-amplitude Mie oscillations are

AMTD
8, 1203–1259, 2015

disappearing. It is then realistic to fit the calibration curve using a power law in $D^{-1.0}$, where D is the particle diameter, as shown in Fig. 3. Such an approach is validated by performing measurements with carbon particles in urban air, as shown in Fig. 4. LOAC captures well the continuous decrease of concentration with increasing size, as previously established by various kinds of instruments, such as electrostatic low pressure impactors (e.g. Shi et al., 1999), while a calibration error would produce oscillations in the size distribution. On the other hand, we would expect that the LOAC calibration could be inappropriate for droplets, which are supposed to be perfectly spherical. In fact, droplets are slightly distorted when entering the optical chamber due to changes in the air flow speed. Thus, no bias on the size distribution nor oscillation correlated with the Mie oscillations were detected for the measurements inside fogs and clouds.

LOAC, with its present calibration procedure, is operated to the detection of irregular grains and droplets, but not to perfect spherical solid grains, such as latex or metal beads for which uncertainties arise from the smoothing of Mie oscillations by the calibration curve.

Overall, a total of 19 size classes are defined for diameters between 0.2 and 100 µm (Table 1). The size classes are chosen as a good compromise between the instrument sensitivity and the expected size distribution of ambient air aerosols.

2.3 Concentration measurements

Counting is conducted while the particles cross the laser beam one by one, and are classified in size classes corresponding to the scattered flux. The measurements are integrated during 10 s and are converted to number densities or particles cm^{-3}. The detectors of the two channels (12 and 60°) work asynchronously.

The optical and electronic response of the system has been modelled by a numerical Monte-Carlo method, taking into account the shape of the laser beam, the speed of the particles inside the laser beam and the instrument noise. To ensure a good statistical approach, 10^4 particles were randomly injected for each size class. The ratio

AMTD
8, 1203–1259, 2015

of the number of detected particles over the number of injected particles provides the detection efficiency. For the smallest particles, only the brightest part of the peak of the pulse of the scattered flux is observable and the apparent transit time in the laser beam is a few tens of µs (in Figs. 2 and 4 such small peaks are present). Thus, some particles cannot be detected. The detection efficiency increases as the diameter of the particles increases, and is reaching 100 % for particles larger than 1 µm. The concentrations of submicron size particles are then corrected using these detection efficiency coefficients.

LOAC can count up to ~ 3000 particles smaller than 1 µm cm^{-3} thanks to the short apparent transit time for the smallest particles. For particles larger than 1 µm, the observed transit time in the laser beam is at its maximum (~ 500 µs) and the expected maximum concentration is of about 20 particles cm^{-3}. In fact, even higher concentrations can be determined using a statistical approach when several particles cross the laser beam almost simultaneously. The higher are the concentration, the lower is the probability that the scattered intensity peaks decreases below the threshold to start a new counting. In this extreme case, the real concentrations are inversely proportional to the detected concentrations. Another Monte-Carlo numerical modelling was conducted to establish the relationship between the number of particles > 1 µm detected and the number of particles injected in the laser beam (Fig. 5). In the simulations, particles were randomly injected (in time), with concentrations increasing from 0 to 500 particles cm^{-3} by step of 1 particles cm^{-3}. The response is almost linear up to 10 particles cm^{-3}, reaching a kind of saturation value at around 15 particles cm^{-3}, and decreases for larger concentrations. It is obvious that such a corrective procedure must be used only in dense aerosol media (more than 10 particles greater than 1 µm cm^{-3}), as fog or clouds, i.e. in conditions which must be confirmed by independent measurements. At present, this procedure is applied only when large droplets are detected by LOAC using the speciation procedure presented below. In this case, up to 200 large particles cm^{-3} can be detected.

AMTD
8, 1203–1259, 2015

For the LOAC integration time of 10 s, the counting uncertainty can be derived from the Poisson counting statistics. This uncertainty, defined as the relative SD, is 60 % for aerosol concentrations of 10^{-2} cm^{-3}, 20 % for 10^{-1} cm^{-3}, and 6 % for concentrations higher than 1 cm^{-3}. Such uncertainties can be reduced by averaging the concentration measurements for each size classes or by increasing the integration time. Nevertheless, such calculation does not take into account the real instrumental uncertainties dominated by the electronic noise and the inlet sampling efficiency.

LOAC is designed to be used in various atmospheric conditions. The temperature can dramatically change, in particular during balloon flights up to the middle stratosphere. The electronic offset can change with time because of the sensitivity of the electronic components to atmospheric temperature variations. The instrument performs a check of its noise level after 10 min of measurements. If the noise differ by more than 50 % from the previous check, an electronic re-calibration is automatically performed to estimate the offset variation and to adjust the calibration. A processing software is applied after the experiment to check the offset time-evolution during the 10 min periods and to then correct the raw measurements.

2.4 Speciation

The scattered flux recorded at 60° is very sensitive to the refractive index of the particles and thus to their nature (as said before this phenomenon appears at scattering angles greater than ∼ 20°). The more absorbing they are, the lower the recorded fluxes. Thus, we use the "60° channel" as a diagnostic for the effect of the refractive index on the scattered fluxes. This channel uses the same calibration threshold (in mV) as the 12° channel, in order to perform a direct comparison of the fluxes recorded by two channels. For a given size class and for a given particle concentration recorded in the 12° channel, the concentration detected by the 60° channel decreases when the imaginary part of the refractive index increases. This increase of the imaginary part leads to an underestimation of the real size of the particles, and thus produces a diameter bias in the size distribution (diameter vs. concentration) for the 60° channel with respect to the

AMTD
8, 1203–1259, 2015

12° channel. An example of the procedure used to determine this effect is presented in Fig. 6, where the size distributions of the two channels are presented. For a given particle size of the 12° channel (noted D1), we consider the concentration value of the 60° channel. Then we search for the same concentration value on the 12° channel (a linear interpolation is used if needed). The corresponding diameter is then determined (D2). Finally, we define a so-called "speciation index" as the ratio D2/D1. The more absorbing the particles are, the higher this ratio. This procedure is conducted for each size class.

This procedure works for a large enough number of detected particles per size class, because of the irregular shape of the particles. In its nominal operating mode, LOAC provides the speciation index every 1 min. For the analysis of continuous ground-based measurements presented below, we have conducted the speciation with an integration time of 15 min (assuming that the aerosols are stationary).

Different types of particles have been tested in the laboratory to assess the amplitude of the speciation index throughout the measurement size range: organic carbon, black carbon, desert dust or sand from different origins (excluding black sand), volcanic ashes, plaster, salt (NaCl), water droplets, droplets of mixture of water and sulphuric acid. They can be classified in 4 families: carbonaceous particles, minerals, salts and liquid droplets. Then, "speciation zones" charts (speciation index vs. real diameter) are defined by the minimum and maximum speciation index values reached by each family, taking into account the measurement uncertainties. Among solid particles, carbononaceous particles produce the higher speciation index and salt the lower, mineral particles being in between. Detailed analysis has shown that most of the carbon particles are in the lower part of the carbon speciation zone while some strongly absorbing particles, perhaps black carbon having fractal shape, are in the middle and upper part of the carbon speciation zone. For all solid particles, the global trend is a decrease of the speciation index with increasing size. On the contrary, the liquid droplets speciation index exhibits an increase with increasing diameter.

AMTD
8, 1203–1259, 2015

The case presented in Fig. 6 has $D1 = 0.35\,\mu m$ and $D2 = 0.51\,\mu m$, leading to a ratio of 1.46, which is in the carbon speciation zone.

The speciation indices obtained from LOAC observations in the atmosphere are compared to the reference charts obtained in the laboratory. The position of the data points in the various speciation zones provides the main nature of the particles. In principle, this procedure can be conducted for each size class. In fact, due to the statistical dispersion of the results, it is better to consider several consecutive suze classes to better conduct the identification. This is in particular necessary for the identification of droplets, whose speciation zone crosses all the speciation zones of the solid particles.

It is obvious that the identification of the nature of the particles works well in case of an homogenous medium, when the speciation indices are not scattered through the various speciation zones.

At present, the speciation zones are established for particles expected to be found in the troposphere and stratosphere, but it is an evolving data base. Additional laboratory measurements can be conducted to retrieve the speciation zone for specific particles in case of measurements in new specific environments.

2.5 Reproducibility

The instrument is industrially produced by Environnement-SA (Environnement-SA, http://www.environnement-sa.fr; more than 80 copies were produced up to mid-2014). Tests have been conducted for the different parts of the instrument: diode, pump, photodiode and electronics, to assess their reproducibility and thus to establish the measurements uncertainty.

The variation of the laser flux from one copy to another is less than 5 %, which has no significant effect on the flux scattered by the particles. The variability of the pump flow was less than ±0.2 L from one pump to another. The flow can be controlled manually by a flow-metre before a balloon flight or during ground based measurements; it can used in the post-processing for the concentration retrieval. The stability of the pump flow over one hour is of about ±5 %, which induces a ±5 % concentration uncer-

AMTD
8, 1203–1259, 2015

tainty. The pump was tested at low temperature and low pressure in balloon flights in the stratosphere and no obvious instability nor loss of performance has been detected. Finally, optical tests have been conducted with a rotating wheel inserted between the laser and the photodiodes. The wheel had pinholes of different diameters to produce different flux levels. Overall, the photodiodes and the electronic contribute to a reproducibility uncertainty of less than 5 %.

Taking into account all these uncertainties, we can expect a total uncertainty of about ±15 %. To confirm this estimate, tests have been conducted with 8 LOAC in a "pollution test room" at LPC2E laboratory (Orléans, France). Various types of solid particles, following mainly a power law size distribution, have been injected in the chamber. A SD of ±15 % (1σ) has been obtained between the different instruments for particles smaller than 10 µm from measurements of these 8 LOACs using the two channels. The SD increases up to ±30 % for particles larger than 10 µm. This increase is due to the low concentrations of large particles, as resulting from the Poisson counting statistics.

2.6 Inlet sampling efficiency

LOAC will be used in different conditions, mainly on the ground and under balloons. Depending on the chosen inlet and the relative speed between the inlet and the wind, the isokinetic sampling is respected or not, and the efficiency of collecting the largest particles can change.

On the ground, a total suspended particulate (TSP) inlet can be used, ensuring efficiency close to 100 % for collecting all the particles up to a few tens of µm. For some specific studies where very large particles dominate, as measurements inside fog or clouds, or because of mechanical constraint if a TSP inlet cannot be mounted, the particles can be collected by a tube having a bevelled metal inlet and downward oriented. In this case, the largest particles are generally under-sampled and a corrective coefficient must be applied, taking into account the direction and the speed of the wind.

For measurements under balloons floating at constant altitude, the relative speed between ambient air and the inlet is close to zero. The sampling efficiency assessed

AMTD
8, 1203–1259, 2015

using the Agarwal and Liu (1980) criterion for an upward-facing inlet shows that the sampling is unbiased for particle with diameter below 20 µm.

The sampling line used during the flights is composed of a thin wall metallic probe and antistatic tubing. The thin wall aerosol probe has an inlet diameter equal to 5.4 mm and is connected to a tube of about 20 cm long and 6.7 mm internal diameter. The sampling line is connected vertically to the LOAC. Nevertheless, due to the tube stiffness, the line can be inclined with a maximum sampling angle of 30° from vertical. The sampling efficiency of the line was assessed using modelling calculations in order to account for changes in atmospheric pressure, temperature and possible changes of the probe orientation during these flights. For that purpose, the values of pressure and temperature as a function of altitude are taken from the international standard atmosphere. Sampling efficiency calculations have been made by considering a balloon ascending velocity of 5 m s^{-1}, a LOAC sampling flow rate equal to 1.7 L min^{-1} and two angles of the sampling line from the vertical (0 and 30°). According to these parameters, the inlet aspiration velocity of the probe is equal to 1.24 m s^{-1} (sub-isokinetic) and the flow is laminar in the tubing for all altitudes.

The mechanisms considered to calculate the sampling efficiency are the inlet efficiency of the probe in isoaxial and isokinetic sampling conditions (Belyaev and Levin, 1974; Hangal and Willeke, 1990) and particle losses in the tubing due to gravitational settling when the line is not perfectly vertical (Heyder and Gehbart, 1977). Calculations have been conducted for particles with diameter ranging from 0 to 20 µm, and from the ground to an altitude of 30 km. Figure 7 presents the sampling efficiency for a 0° deviation (isoaxial) and for a 30° deviation of the sampling line with respect to the vertical. Data are plotted according to the particle aerodynamic diameter which describes particle settling and inertia phenomenon.

In isoaxial conditions, results show for all altitudes an increase of sampling efficiency with the particle diameter, up to a factor > 3 for the largest particles. In this case, there is no particle deposition in the sampling line and the sampling is dominated by sub-isokinetic conditions (apparent wind velocity higher than inlet probe velocity). A sam-

AMTD
8, 1203–1259, 2015

pling efficiency higher than unity is explained by the particle inertial effect resulting from the divergence of the flow field at the inlet of the probe. The increase in sampling efficiency with altitude is due to changes in air viscosity and gas mean free path with temperature and pressure.

When the tube is inclined by 30° from the vertical, the sampling efficiency is between 1 and 2. The sampling efficiency is lower than for the 0° isoaxial conditions. Firstly, the sub-isokinetic effect is reduced by the orientation of the tube, and secondly, deposition can occur in the tubing due to particle settling.

Since the tube always has a deviation of about 30° during the balloon flights, we consider only the results at 30° from the vertical. The over-sampling effect is negligible for particles smaller than 5 µm up to the lower stratosphere and for particles smaller than 2 µm in the middle stratosphere. Thus, this effect will just affect the retrieved concentrations of the largest particles.

The results of these theoretical calculations are not yet fully validated by an experimental approach with LOAC itself. Thus, all balloon measurements in the stratosphere will not be corrected from this aerodynamic effect. It could be taken into account in future work involving large particles, for example when converting concentrations to extinction by comparison with remote sensing instruments, or to estimate the real concentration of the interplanetary dust in the middle atmosphere.

3 Cross-comparison with other instruments

Various cross-comparisons have been conducted in ambient air at ground and during balloon flights for concentrations and speciation. LOAC concentrations are compared to other commercial particle counter instruments and photometer measurements, although there is no absolute reference, many of them are using different technical approaches and calibration procedures. The LOAC speciations are validated during well-identified atmospheric events of liquid and solid particles. Finally, the LOAC particle concentrations are converted to mass concentrations to be compared to commercial

AMTD
8, 1203–1259, 2015

microbalance mass instruments used as reference instruments in air quality monitoring. Table 2 summarizes the conditions of measurements.

3.1 Concentrations and size distribution

Continuous measurements have been conducted in ambient air at the SIRTA observatory (Site Instrumental de Recherche par Télédétection Atmosphérique, http://sirta.ipsl.fr/) at Palaiseau, South of Paris, France (48.713° N, 2.208° E), during the ParisFog campaign, http://parisfog.sirta.fr/, from November 2012 to April 2013. During this period, the total concentrations of aerosols have been monitored by a WELAS aerosols counter, a Fog Monitor (counter for large droplets) and a SMPS (Scanning Mobility Particle Sizer) in common size range domains with LOAC. SMPS measurements are based on the electric mobility diameter of the particles, while the other instruments determine optical diameters. These two diameters could differ depending on the nature of the particles; thus the direct comparison of the measurements (even in the same air mass) could be sometimes biased.

Figure 8 presents the cross-comparison of the instruments in January 2013. Most of the measurements were conducted in background aerosol conditions, although some small fog events were detected and can be identified by concentration peaks in the Fog Monitor measurements. Roughly speaking, the order of magnitude of the concentrations is similar, although some significant discrepancies are present. To investigate their possible origin, the size distribution obtained in different conditions of aerosol content can be compared. Figure 9 (upper panel) presents an example where the agreement in total concentration during background aerosol conditions is very good between LOAC and SMPS. On the other hand, the shape of the size distribution of the WELAS instrument is unusual with a decrease of the sub-µm aerosol concentrations with decreasing size (the opposite trend is expected for background aerosol conditions). This could be due to calibration problem of the latter instrument; indeed, Heim et al. (2008) report a strong decrease in counting efficiency with decreasing particle diameter for submicron particles for the WELAS-2100 OPC. Finally, as expected, the Fog Monitor

AMTD
8, 1203–1259, 2015

measurements are not useable (no fog at this time and only noise is present). Figure 9 (lower panel) presents the same cross-comparison in case of significant disagreements between all the instruments. LOAC seems to underestimate the concentrations of the smallest particles, but this can be due to the difficulty of reconciling the retrieved diameters for the different measurement techniques in case of different types of particles.

Strong fog events were observed in November 2012. LOAC, WELAS and Fog Monitor are in very good agreement during these events (Fig. 10). This result is validating the correction procedure applied to the LOAC measurements in case of dense medium of liquid particles. Between fog events, LOAC and WELASwere sometimes in disagreement. This was due to the difference in the concentration values obtained by the two instruments for the particles smaller than $\sim 0.5\,\mu m$, which may be attributed to the WELAS undercounting. Figure 11 presents the size distribution at the beginning of a fog event, with the typical enhancement around a diameter of $10\,\mu m$ (e. g. Singh et al., 2011), and at the end of the event. Both LOAC and WELAS found a bimodal size distribution although the WELAS shows a doubtful sharp decrease in concentration for the smallest sizes., but disagree for the size of the second mode at $\sim 10\,\mu m$. On the opposite, LOAC and Fog Monitor were in good agreement for the position of the second mode, although the population of the first size class of Fog Monitor was obviously underestimated. Finally, for the largest sizes, LOAC concentrations are in-between those of the WELAS and Fog Monitor.

A ground-based measurement session was conducted from Minorca (Spain) during the ChArMEx campaign (Chemistry Aerosol Mediterranean Experiment, https://charmex.lsce.ipsl.fr/).in parallel with measurements of an HHPC-6 aerosol counter in the period 12 June–2 July 2013. The orders of magnitude for the different size classes were in good agreement. In particular, both instruments captured an aerosol enhancement of large solid particles between 18 and 21 June 2013, as shown in Fig. 12 for the size distribution.

The last cross-comparison exercise was conducted during an ambient air campaign at SIRTA observatory, site#5 near Gif-sur-Yvette, South of Paris, France (48.709° N,

AMTD
8, 1203–1259, 2015

2.149° E), in the beginning of 2014. LOAC performed measurements from 3 to 13 February 2014 in parallel with a SMPS, a Grimm aerosol counter and a HHPC-6 aerosol counter. Due to the sampling conditions that vary from one instrument to another (direct sampling, TSP inlet, dryer, direct or curved tubes), the analysis is limited to the smallest particles (diameter < 1 µm) which are almost insensitive to the sampling techniques. Figure 13 presents the temporal cross-comparison for 4 size-classes: 0.2–0.3, 0.3–0.5, 0.5–0.7 and 0.7–1.0 µm. In fact, the size classes of the 4 instruments are not always the same, thus the closest ones have been considered for the comparison.

Globally, all the instruments give similar concentrations for all size classes, the better agreement being for the 0.5–0.7 µm diameter range. Some discrepancies appear for some time periods, but several reasons can explain them. First, the SMPS instrument determines the electric mobility diameter that can depend on the nature of the aerosols, whereas the other instruments determine optical diameters. This could explain why LOAC has missed some concentration peaks detected by SMPS. Secondly, the particles size distribution of sub-µm particles exhibits a strong decreasing when the diameter increases. Thus uncertainty in the size calibration of a few hundredth of µm could induce concentration differences of at least a factor 2. This is presented in Fig. 13 for the 0.7–1 µm comparison with the Grimm instrument for which both 0.65–1 and 0.8–1 µm concentrations are plotted. Finally, the Grimm and HHPC-6 instruments are sensitive to the nature of the particles, and changes in the type of aerosol (for example mineral or carbon particles) could partially affect their size determination.

An indirect evaluation of the LOAC size calibration has been conducted during the ChArMEx campaign on the Balearic island of Minorca, Spain. A total of 9 flights of LOAC have been performed under a meteorological sounding balloon launched from Sant Lluís airfield (39.865° N, 4.254° E) in the 15–19 June 2013 period during a desert dust transport event. The aerosol concentration has been integrated for all size classes from the ground to the highest altitude reached by the balloon, i.e. an altitude of about 30 km, to be compared to ground-based remote sensing measurements provided by the AERONET photometer network (http://aeronet.gsfc.nasa.gov/) station of Cap d'En

AMTD
8, 1203–1259, 2015

Font (39.826° N, 4.208° E), which performed measurements close the trajectory of the LOAC balloon measurements. AERONET provides the vertically integrated volume concentration of aerosols (in $dV/d\ln(r)$, where r is the radius of the particles) in the 0.13–30 µm radius range (Dubovik et al., 2000).

The LOAC integrated concentrations are converted to volume concentrations by using the mean volumetric diameter D_v calculated for each size class by the formula:

$$D_v = 0.5 \times \left[\left(D_{min}^3 + D_{max}^3\right)\right]^{(1/3)}$$

where D_{min} and D_{max} are the lower and upper diameter of a given size class, respectively. With such a formula, the mean volumetric diameter is at about 60 % of the size class width instead of 50 % for the mean geometric diameter. For each size class, the volume of the particles is calculated assuming sphericity. To be consistent with the AERONET data, the LOAC results are presented in radius instead of diameter.

Figure 14 presents two examples of comparison between LOAC and AERONET size distributions for two different amounts of sand particles in the troposphere (the contribution of the stratospheric particles is negligible). The bi-modal distribution is typical for a desert dust or sand plume event. The two instruments are in excellent agreement, both in size distribution and volume concentrations It is worth noting that the volume concentrations are proportional to the cube of the size of the particles, an error in the LOAC calibration would lead to strong discrepancies, which is not the case.

All these cross-comparison exercises have shown that the LOAC measurements are consistent with those of the other instruments considered here, accounting for the errors and the limitation of the various techniques. This confirms that no systematic bias are present in the LOAC calibration and in the data analysis.

3.2 Speciation

The speciation zones, obtained from laboratory measurements, must be validated in real atmospheric conditions.

AMTD
8, 1203–1259, 2015

Urban ambient air measurements are proper for the detection of carbon particles (black and organic carbon), especially during well-identified pollution events. Permanent LOAC measurements have been conducted at "Observatoire Atmosphérique Generali" (OAG) in the South-West of Paris since May 2013 (48.841° N, 2.274° E). This observatory is a recreational tethered balloon operated in a public park; the LOAC measurements nominal maximum altitude is 120 m but some flights could be conducted up to an altitude of 270 m. The measurements can be sorted out between measurements with the balloon at ground level and measurements during flight. Figure 15 presents an example of carbon particles detected at the OAG on 29 December 2013 around 07:30 UT. In this example, the "speciation index" is well inside the carbon speciation zone in the whole size range up to ~ 10 µm.

In addition to sounding balloons mentioned above, measurements under drifting balloons launched from Sant Lluís on Minorca Island were also conducted during several well-identified desert dust events above the Mediterranean Sea during the summer ChArMEx campaign. Figure 16 presents an example on 17 June 2013, around 14:30 UT (approximative balloon position: 41.9° N, 4.1° E) at an altitude of 2050 m under a low altitude pressurized drifting balloon. The speciation curve is well inside the mineral dust zone, showing that LOAC has indeed detected the desert dust event.

Measurements in the marine atmospheric boundary layer were also conducted with a low altitude balloon on 22 July 2013 drifting in an altitude range of 250–400 m, launched from the French Levant Island off Hyères on the Mediterranean French coast (43.021° N, 6.461° E). Figure 17 presents the measurements at 21:25 UT (approx. balloon position: 43.0° N, 6.55° E, alt. ~ 275 m), and the speciation is mainly in the "salt zone", as expected for a measurement close to the sea surface.

Droplet speciation was validated in fog events during the ParisFog campaign; but also during cloud measurements conducted in May 2013 at the Puy de Dôme observatory (45.772° N, 2.964° E, alt. 1465 m). Figure 18 presents an example of measurements inside a cloud on 15 May 2013 at 10:30 UT. Globally, the speciation is inside the droplets zone, which indicates that all of the particles were indeed liquid. In addition,

AMTD
8, 1203–1259, 2015

measurements were conducted inside haze or thin cloud at an altitude of 1.2 km during a flight under a meteorological balloon launched from Reykjavik, Iceland (64.127° N, 21.904° W), on 7 November 2013 at 12:30 UT in the frame of the VOLTAIRE-LOAC campaign for the study of the stratospheric aerosol trend. The presence of the droplets was confirmed by the on-board humidity sensor, with a hygrometry of 90 %. The speciation in Fig. 19 is well inside the droplets zone.

Finally, most of the measurements under meteorological balloons in the middle atmosphere show that (pure) liquid water and sulphuric acid droplets largely present in the stratosphere are close to the lower part of the droplets zone, and sometimes slightly below. Vertical profiles of LOAC concentration and speciation measurements are presented in paper 2.

These examples show that the speciation determination works well in case of homogeneous aerosol media. Nevertheless, there are two limitations of this process. First, the analysis of measurements conducted in heterogeneous media could be difficult or even inaccurate, in particular when different size modes are present. In this case, the speciation curve exhibits unusual oscillations that match none of the speciation zones. Secondly, some high porosity aerosols can exhibits high values for the "speciation index", even if they are not black (as fluffy silica). Then, the speciation determination is providing most of the time the main nature of the particles, but one has to be cautious in the analysis when the speciation curves are non-typical.

3.3 Mass concentrations

Our final test to evaluate both the calibration of LOAC and the retrieval of concentrations in all size classes (but especially large particles) is to convert the number size distribution measurements to mass concentrations and to compare the results to reference mass measurements. This is the most sensitive test to evaluate LOAC, since mass concentrations are proportional to the cubic diameter of the particles (and to their density). The speciation helps to determine the type of aerosols, from which a density

AMTD
8, 1203–1259, 2015

can be deduced. The density determination is necessary for the conversion of number concentrations (in cm^{-3}) to mass concentrations (in $\mu g\,m^{-3}$).

Measurements were conducted first in indoor air (in the "pollution room" at the LPC2E laboratory) in autumn 2013, by injecting in the air of the room different kinds of carbonaceous and mineral particles (smaller than 20 µm) in various concentrations to produce a large range of mass concentrations. The reference mass measurements were achieved with a calibrated TEOM microbalance. An air flow system was used (when needed) to prevent sedimentation of the particles in the room. Also, some measurements have been conducted without injecting particles, to detect only the smallest particles present in the ambient air, in particular during the night without convection in the room.

The volume concentration is calculated for each size class, using the mean volumetric diameter, assuming spherical particles, and is multiplied by the corresponding concentrations. The mass concentration is obtained by multiplying these results by the particle density. The mass densities were determined for each size class by identifying the nature of the particles though their speciation index. The mass densities chosen here are:

- $2.2\,g\,cm^{-3}$ for salt; a value corresponding to NaCl particles;
- $2.2\,g\,cm^{-3}$ for mineral particles; this value is a compromise for common mineral particles present in ambient air: compact sand ($2.1\,g\,cm^{-3}$), quartz ($2.7\,g\,cm^{-3}$), limestone ($2.5\,g\,cm^{-3}$) and silicon ($2.3\,g\,cm^{-3}$);
- $1.4\,g\,cm^{-3}$ for carbonaceous particles. This value was derived after detailed tests during the comparison between LOAC and microbalance measurements in the laboratory. It lies well inside the range of values proposed in the literature for such particles (e.g. Chen et al., 2010; Virtanen et al., 2006; Spencer et al., 2007). Sensitivity tests have shown that a 10 % variation of this value will not induce strong changes in the results presented below.

AMTD
8, 1203–1259, 2015

– A value of 0.0 g cm^{-3} was used for water droplets, for comparing LOAC measurements to those of the TEOM instrument, which tends to evaporate condensed water.

The duration of the sessions was from several hours to several days. Figure 20 presents the mass measurements for particles smaller than 20 µm, averaged on 24 h for the two instruments. The variability of the concentrations is related to the amount of particles injected into the room. The lowest values correspond to measurements without injection. In this case, LOAC indicates that only particles smaller than 2 µm were present in the air. The LOAC and TEOM measurements are in very good agreements, with a correlation of 0.97. The correlation curve has the slope of 0.98, with an offset at the origin of 2.2 µg m^{-3}, and a mean error of 4.8 µg m^{-3}.

Sessions of ambient air measurements were conducted in Paris and in its suburbs, to test the retrieval of $PM_{2.5}$ and PM_{10} mass concentrations, with pumps working at 2.7 L min^{-1}. The first location of measurements is at the "Observatoire Atmosphérique Generali (OAG)" in Paris (latitude 48.8417° N, longitude 2.2736° E). The LOAC measurements were conducted using a TSP inlet. The second location is at SIRTA observatory at Palaiseau (48.7180° N, 2.2075° E) during the ParisFog campaign. The LOAC measurements were conducted with the metal inlet directed towards the ground. The OAG and SIRTA measurements considered here were conducted in the periods September 2013–April 2014 and September–December 2013, respectively. The $PM_{2.5}$ and PM_{10} LOAC mass concentrations were retrieved by combining the results for particles smaller than 3 µm an smaller than 10 µm, respectively, taking into account the sampling efficiency of the $PM_{2.5}$ and PM_{10} inlets currently used by the air quality networks (cut-off at 2.5 µm for $PM_{2.5}$ inlet and cut-off at 10 µm for PM_{10} inlet).

Reference mass concentrations data of urban ambient air in the Paris region are provided by the AirParif network (http://www.airparif.asso.fr/), operating TEOM microbalance instruments. Unfortunately, there is no AirParif station very close to the OAG site nor to the SIRTA site at the time of the measurements. Therefore, we decided to use data recorded at 3 stations that have environmental conditions close to those at

AMTD
8, 1203–1259, 2015

OAG and SIRTA: Paris Centre (latitude 48. 8528° N, longitude 2.3600° E), Vitry (latitude 48.7820° N, longitude 2.3992° E) in the south-eastern suburb area of Paris, and "Rural South" at Bois-Herpin (latitude 48.3725° N, longitude 2.2258° E) in the South of the Paris region; the last station provides background conditions measurements.

Figures 21 and 22 present the comparison of $PM_{2.5}$ and PM_{10} concentrations, for the 2013 and 2014 period, respectively. The LOAC measurements are most of the time between the background and the urban conditions, the small discrepancies with the reference mass concentrations are probably due to a difference in the wind direction and to the regional-scale transport of the particles. It is worth noting that LOAC did capture well the 10–15 December 2013 and the 11–14 March 2014 pollution peaks.

These measurement sessions have been conducted with different kinds of pumps and of inlet systems. The agreement with reference mass concentration measurements is very good. This confirms that no obvious bias is present in LOAC observations for the sizes of particles considered here (~ 0.2–20 µm), and that the speciation procedure is providing useful information to convert the LOAC concentrations for the 19 size classes to mass concentrations.

4 Conclusions

LOAC is a modular optical particle counter/sizer, of which the pump and the air inlet can be changed, depending on the conditions of measurements. Extensive tests performed in different atmospheric conditions have shown that LOAC can be used to retrieve the size distribution of irregular-shaped or liquid aerosols with a satisfactory accuracy at ground level and from all kinds of balloons. The uncertainty (at 1σ) is of about ±15 % for concentrations greater than 10^{-1} particles cm^{-3} and of about ±30 % for lower concentrations in an integration time between 2 and 15 min depending on the concentration and flow rate. LOAC can also provide an estimate of the nature of the particles in case of a relatively homogeneous media. Finally, LOAC can be used for monitoring the mass concentration of $PM_{2.5}$ and PM_{10} (and of course of largest parti-

AMTD
8, 1203–1259, 2015

cle) in ambient air. in ambient air with reasonable accuracy. The companion paper will present first scientific results from balloons and an unmanned aerial vehicle (Renard et al., 2015).

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AMTD

8, 1203–1259, 2015

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AMTD
8, 1203–1259, 2015

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AMTD
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AMTD
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AMTD
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AMTD
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Table 1. The 19 size classes of LOAC for concentration measurements.

Diameter range (µm)
0.2–0.3
0.3–0.4
0.4–0.5
0.5–0.6
0.6–0.7
0.7–0.9
0.9–1.1
1.1–3.0
3.0–5.0
5.0–7.5
7.5–10.0
10.0–12.5
12.5–15.0
15.0–17.5
17.5–20.0
20.0–22.0
20.0–30.0
30.0–40.0
40.0 – up to 100.0

AMTD
8, 1203–1259, 2015

Table 2. Conditions of measurements for evaluation exercises.

Campaign	Location	Date	Installation	Instruments for validation
ParisFog	SIRTA Observatory, Palaiseau (France)	Nov 2012–Apr 2013 Sep 2013–Jan 2014	Continuous ground measurements	– WELAS counter – Fog monitor counter – Scanning Mobility Particle Sizer (SMPS)
Cloud measurements	Puy de Dôme (France)	May 2013	Continuous ground measurements	Well-known atmospheric conditions for the speciation
ChArMEx	Minorca (Spain)	17 Jun 2013	Tropospheric pressurized balloon	Well-known atmospheric conditions for the speciation
ChArMEx	Ile du Levant (France)	22 Jul 2013	Tropospheric pressurized balloon	Well-known atmospheric conditions for the speciation
ChArMEx	Minorca (Spain)	15 Jun–2 Jul 2013	Continuous ground measurements	HHPC-6 counter
ChArMEx	Minorca (Spain)	16 and 17 Jun 2013	Meteorological sounding balloon flights	Well-known atmospheric conditions for the speciation
QAIDOMUS	Orléans (France)	Sep–Nov 2013	Indoor air	TEOM microbalance
VOLTAIRE-LOAC	Reykjavik (Iceland)	7 Nov 2013	Meteorological balloon flight	Well-known atmospheric conditions for the speciation
Observatoire Atmosphérique Generali	Paris (France)	Jan – Apr 2014	Permanent measurements on tethered balloon (at ground and up to an altitude of 270 m)	– TEOM microbalances (AirParif air quality network) – Well-known atmospheric conditions for the speciation
SIRTA5 campaign	Gif-sur-Yvette (France)	3–13 Feb 2014	Continuous ground measurements at SIRTA	– Grimm counter – HHPC-6 counter – SMPS

AMTD
8, 1203–1259, 2015

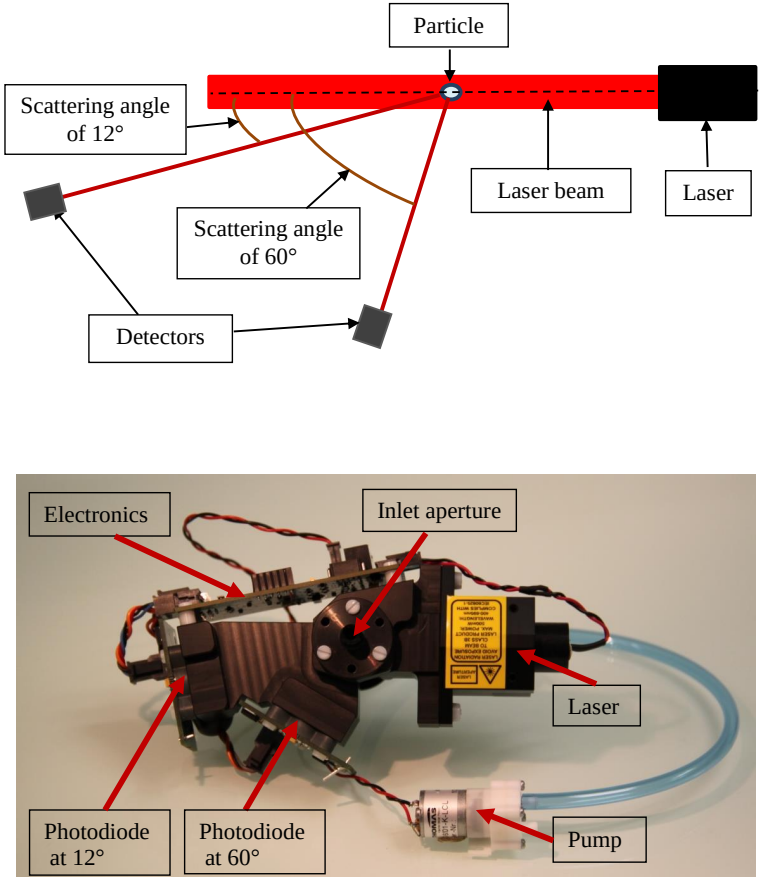

Figure 1. The LOAC instrument; upper panel: principle of measurement; lower panel: picture of the instrument (the inlet tube is not presented here).

AMTD
8, 1203–1259, 2015

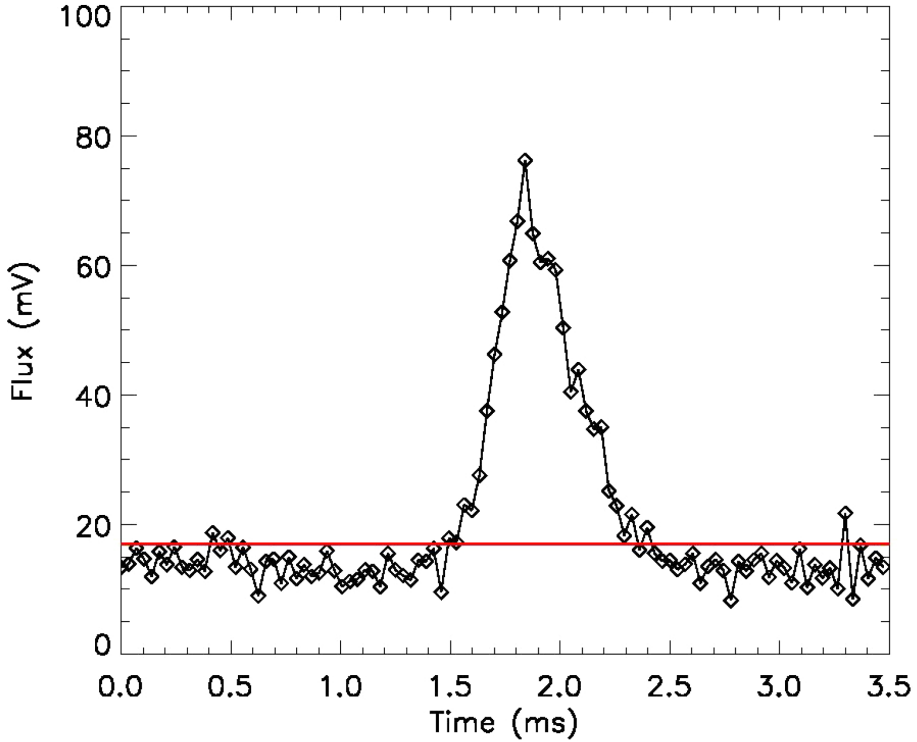

Figure 2. Example of the flux scattered by particles while crossing the laser beam. The red line corresponds to the threshold for the peak detection. When a particle is detected, the signal must return back below the threshold to allow the detection of the next one. In this example, the small particle causing the small secondary peak at 2.1 ms is not counted.

AMTD
8, 1203–1259, 2015

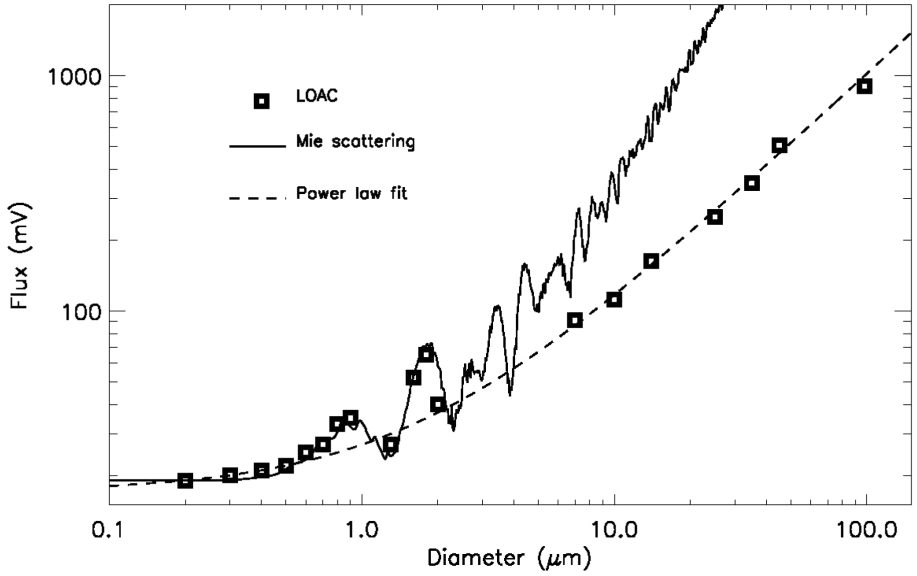

Figure 3. Calibration curve of the scattered flux at 12° as a function of particle diameter; the difference between the Mie scattering calculations and LOAC measurements for diameters greater than 5 µm is due to the small aperture of the field of view coupled with the roughness of the particle shapes; the measurement curve is fitted by a power law.

AMTD
8, 1203–1259, 2015

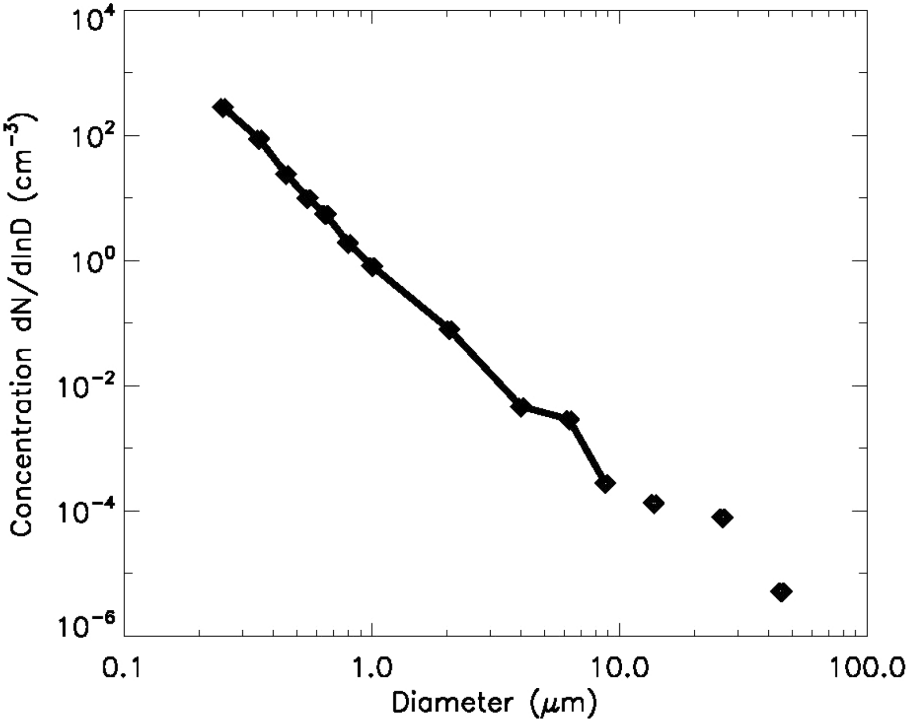

Figure 4. Typical size distribution in a suburban ambient air with carbon particles (Palaiseau, South of Paris) on 14 October 2013 during ParisFog campaign; the data are integrated during 15 min; the last points are not related because of zero concentration measured between them.

AMTD
8, 1203–1259, 2015

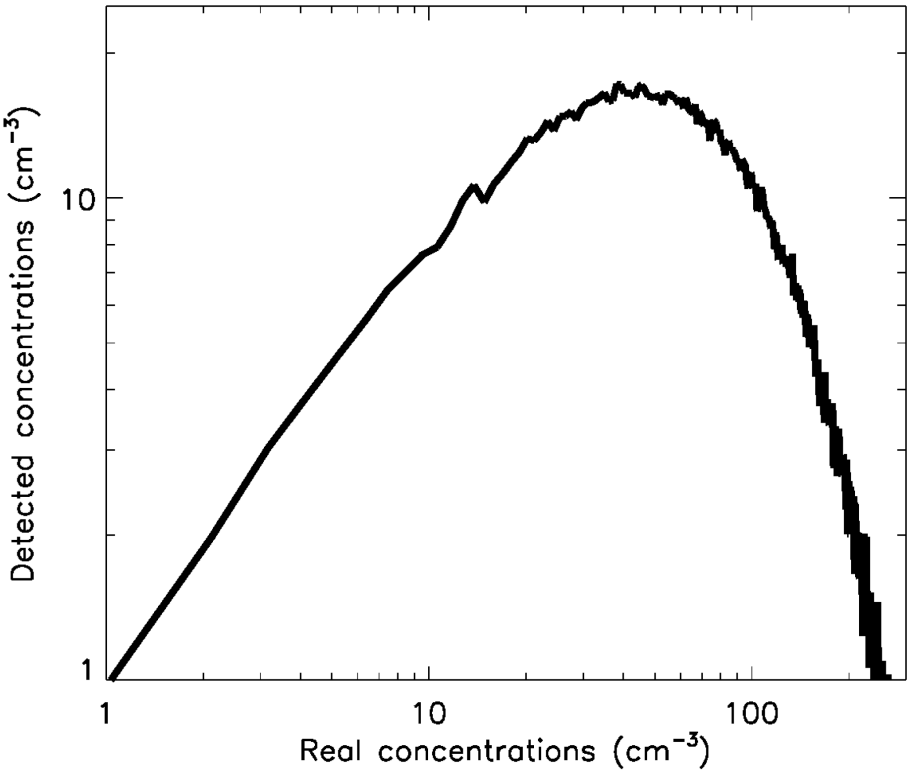

Figure 5. Monte-Carlo modelling for the response of the counting system for particles larger than 1 µm. The response is almost linear up to 10 particles cm^{-3}, and decreases for large concentrations.

AMTD
8, 1203–1259, 2015

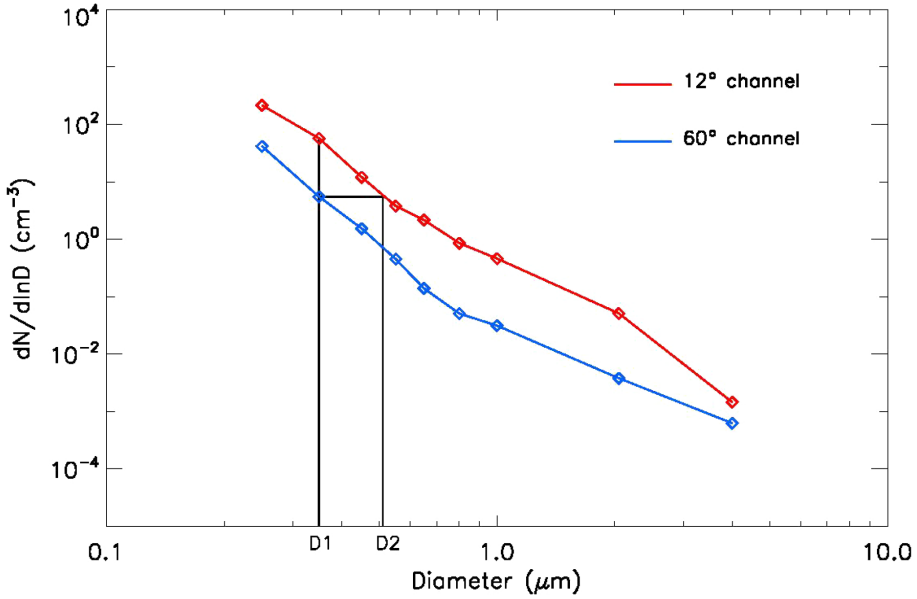

Figure 6. Principle of the determination of the "speciation index" D2/D1 (the example presented here uses real measurements).

AMTD
8, 1203–1259, 2015

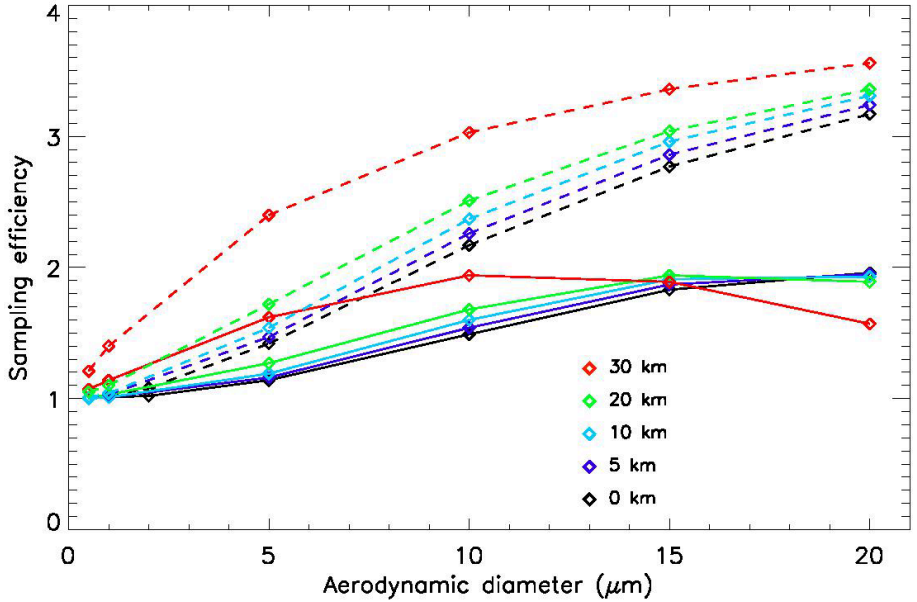

Figure 7. Efficiency of the sampling line at different altitudes from the surface up to 30 km; dashed lines: isoaxial conditions; full lines: 30° deviation from isoaxial conditions.

AMTD
8, 1203–1259, 2015

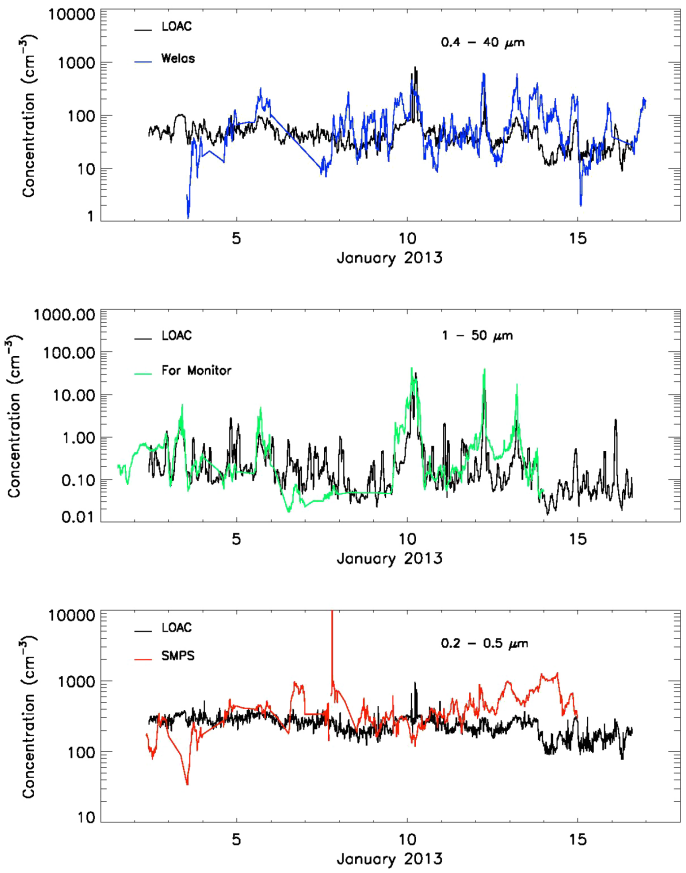

Figure 8. Cross-comparison of LOAC with 3 other instruments (WELAS, Fog Monitor and SMPS) for the total concentrations of aerosols in the size range domain in common, during the ParisFog campaign south of Paris. The LOAC uncertainties are of ±15 %.

AMTD
8, 1203–1259, 2015

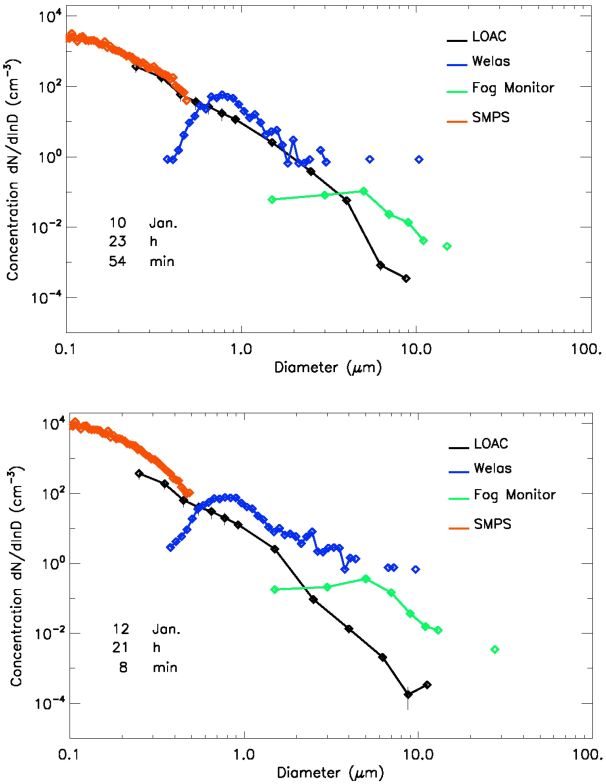

Figure 9. Cross-comparison of the 4 instruments during background conditions, in case of good agreement for the total concentrations measurements, during the ParisFog campaign. Upper panel: 10 January 2013, good agreement between the instruments; lower panel: 12 January 2013, poor agreement. The LOAC uncertainties are of ±15 %. The WELAS probably underestimates sub-µm particles (Heim et al., 2008).

AMTD
8, 1203–1259, 2015

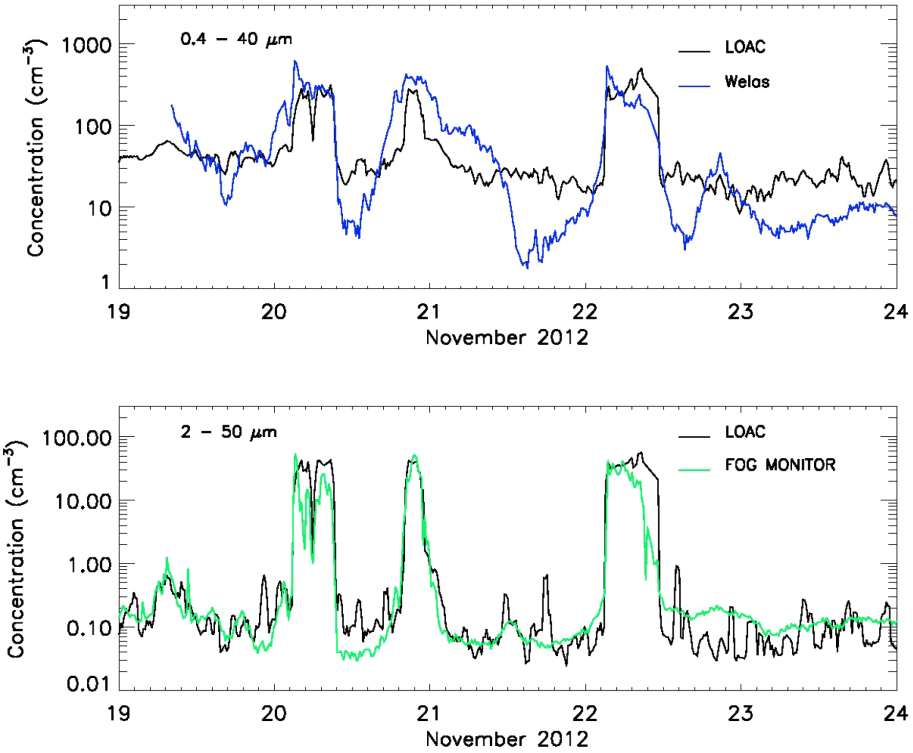

Figure 10. Cross-comparison of LOAC with 2 other instruments (WELAS and Fog Monitor) for the total concentrations of aerosols in the size range domain in common, during the ParisFog campaign. The LOAC uncertainties are of ±15 %.

AMTD
8, 1203–1259, 2015

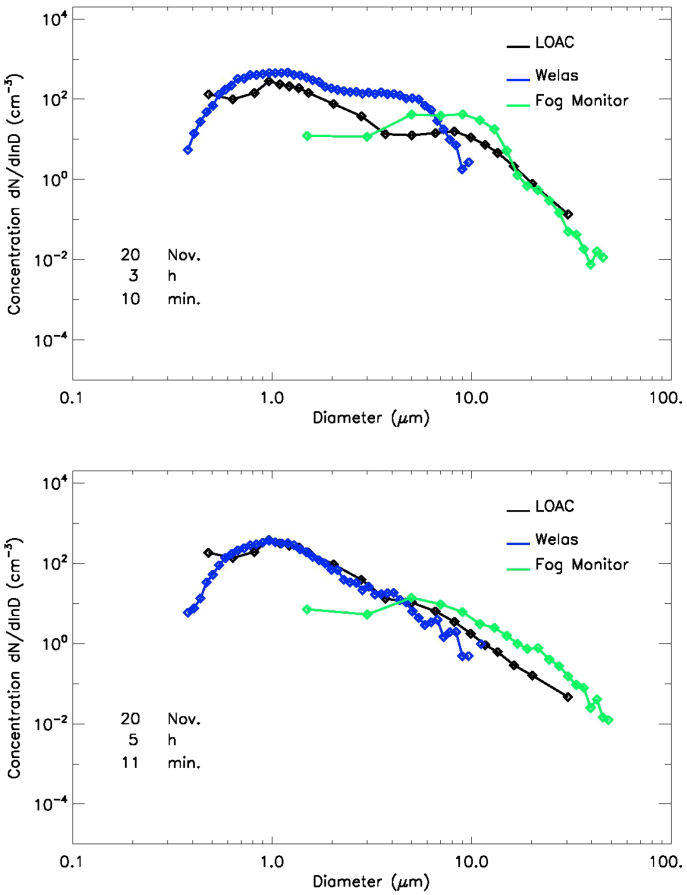

Figure 11. Cross-comparison of the 3 instruments at the beginning of the fog event (top) and at the end (bottom), during the ParisFog campaign on 20 November 2012 during a fog event. The LOAC uncertainties are of ±15 %. The WELAS probably underestimates sub-µm particles.

AMTD
8, 1203–1259, 2015

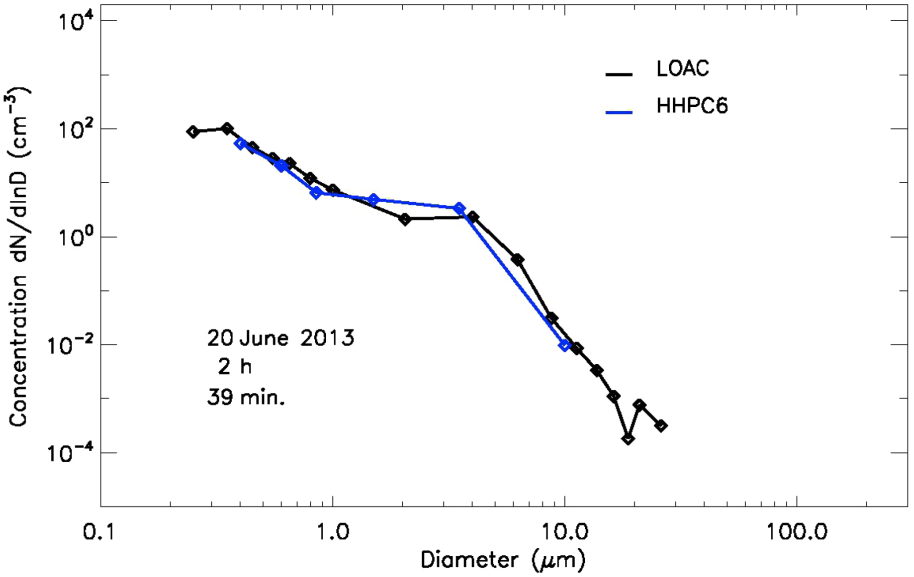

Figure 12. Example of size distribution for LOAC and HHPC-6 during an event of solid particles during the ChArMEx campaign at Minorca on 20 June 2013.

AMTD
8, 1203–1259, 2015

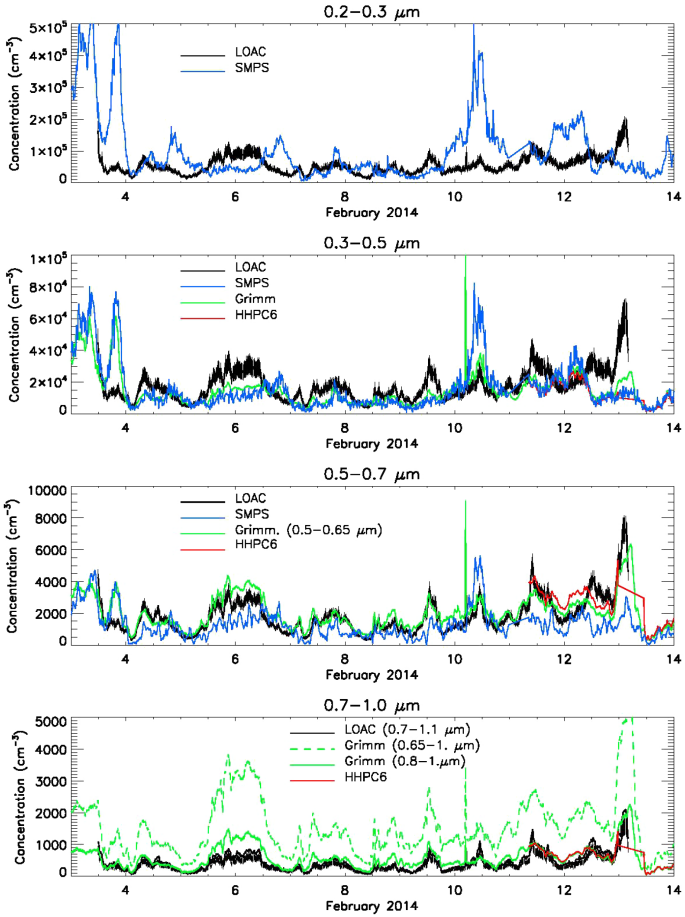

Figure 13. Comparison (in linear scale) between the ambient air measurements obtained during the campaign at the SIRTA-5 station South of Paris.

AMTD
8, 1203–1259, 2015

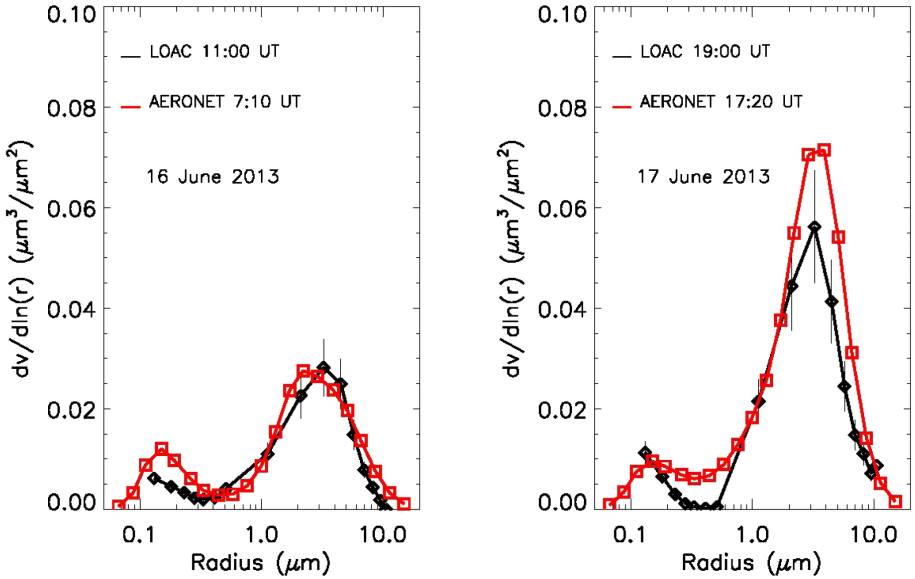

Figure 14. Comparison between integrated LOAC measurements from vertical profiles obtained under meteorological balloons and AERONET measurements during an African dust transport event during the ChArMEx 2013 campaign (note that the LOAC data are given in radius to match the AERONET format).

AMTD
8, 1203–1259, 2015

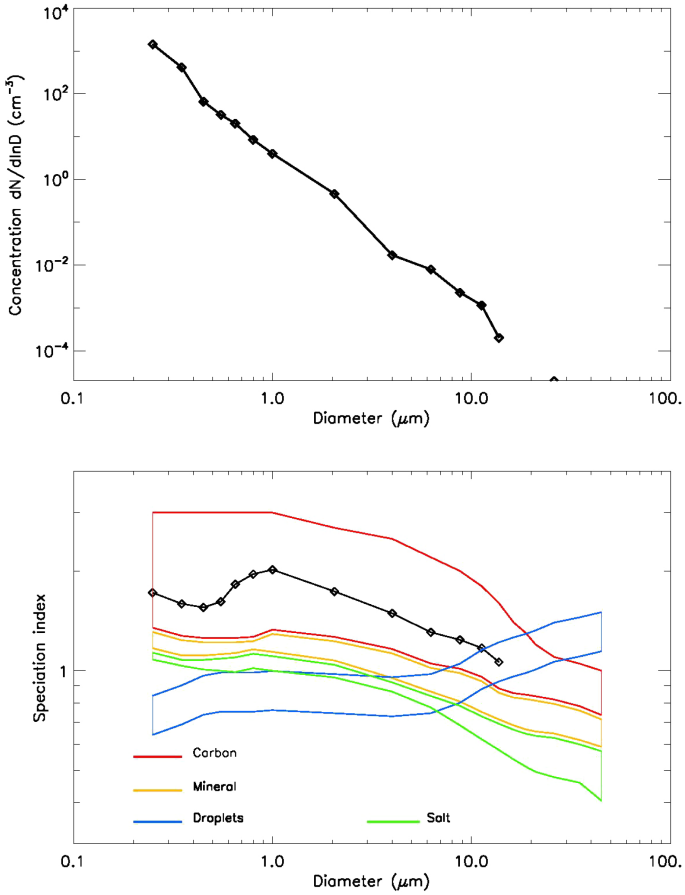

Figure 15. Example of the detection of carbon particles in urban air, in South-West of Paris on 29 December 2013 around 07:30 UT, at the "Observatoire Atmosphérique Generali"; upper panel: size distribution; lower panel: speciation.

AMTD
8, 1203–1259, 2015

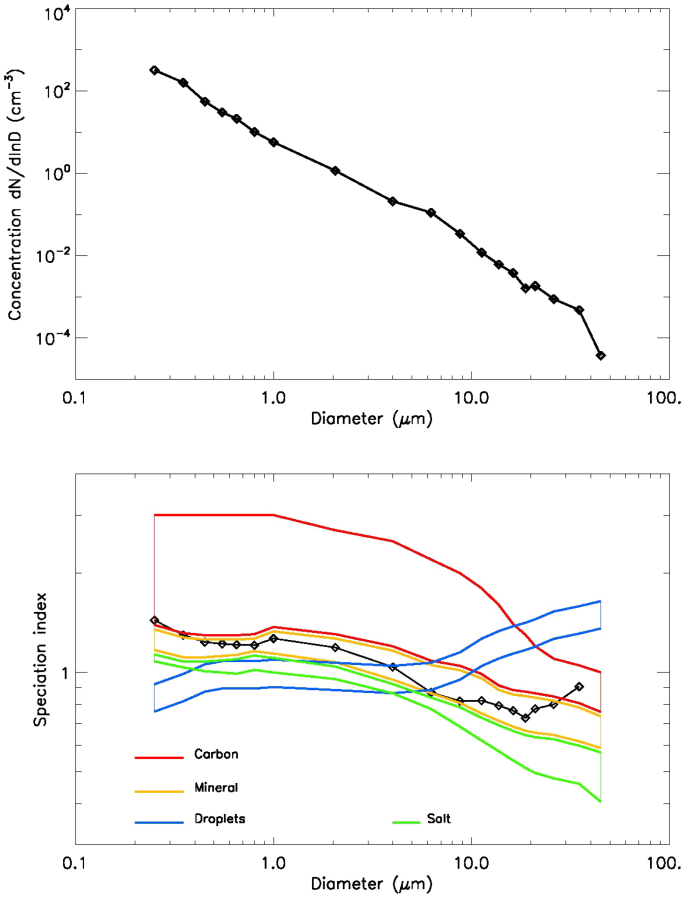

Figure 16. Example of the detection of sand particles above Mediterranean Sea (close to Minorca) from a drifting pressurized tropospheric balloon on 17 June 2013 around 14:30 UT at an altitude of 2050 m, during the ChArMEx campaign; upper panel: size distribution; lower panel: speciation.

AMTD
8, 1203–1259, 2015

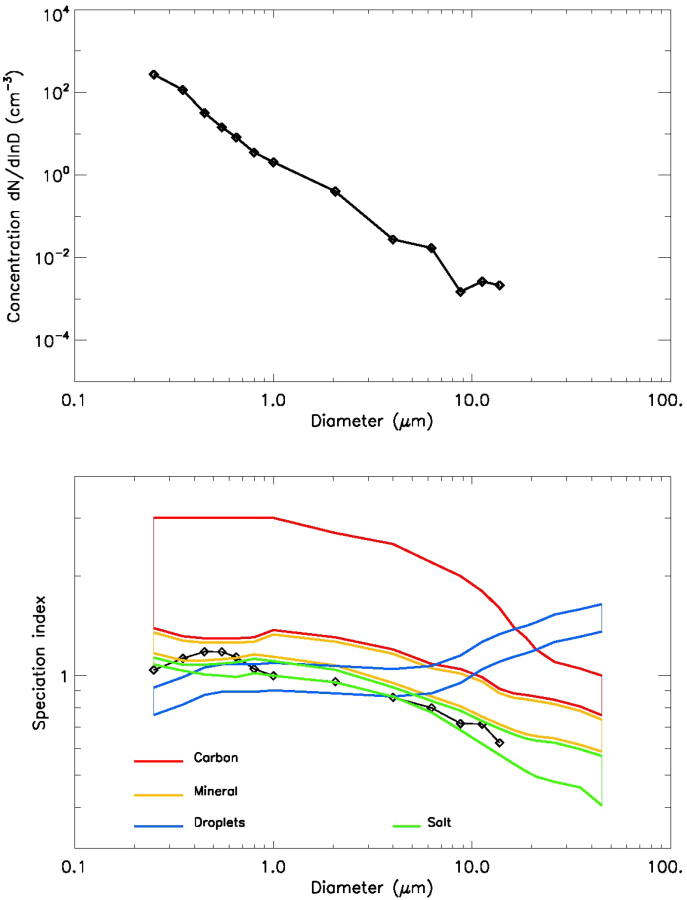

Figure 17. Example of the detection of salt particles above Mediterranean Sea (close to Minorca, Spain) from balloon on 22 July 2013 at 21:25 UT at an altitude of 275 m during the ChArMEx campaign; upper panel: size distribution; lower panel: speciation.

AMTD
8, 1203–1259, 2015

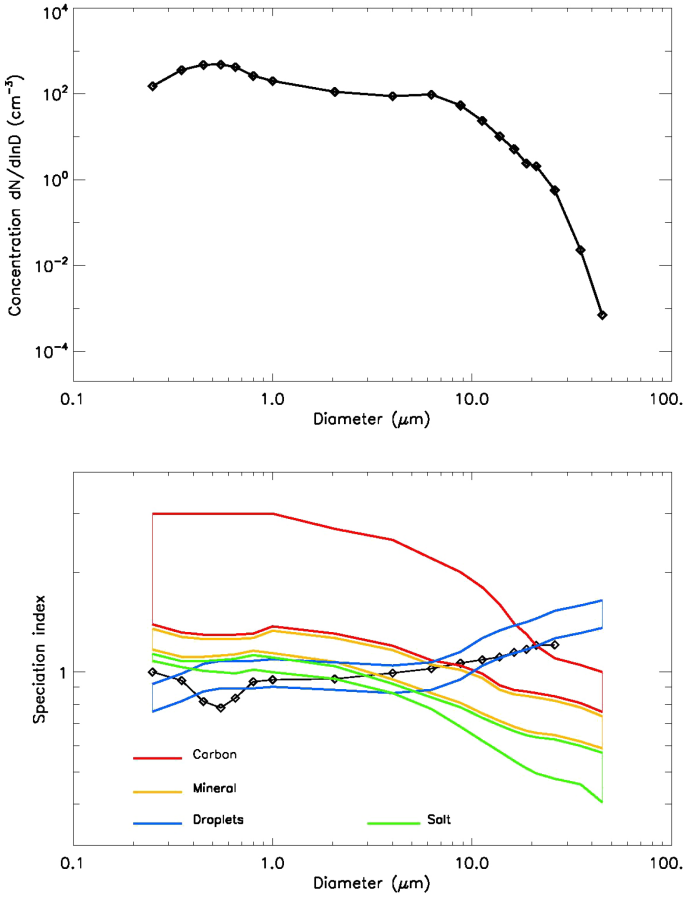

Figure 18. Example of measurements inside a cloud at Puy de Dôme observatory (France) on 15 May 2013 at 10:30 UT; upper panel: size distribution; lower panel: speciation.

AMTD
8, 1203–1259, 2015

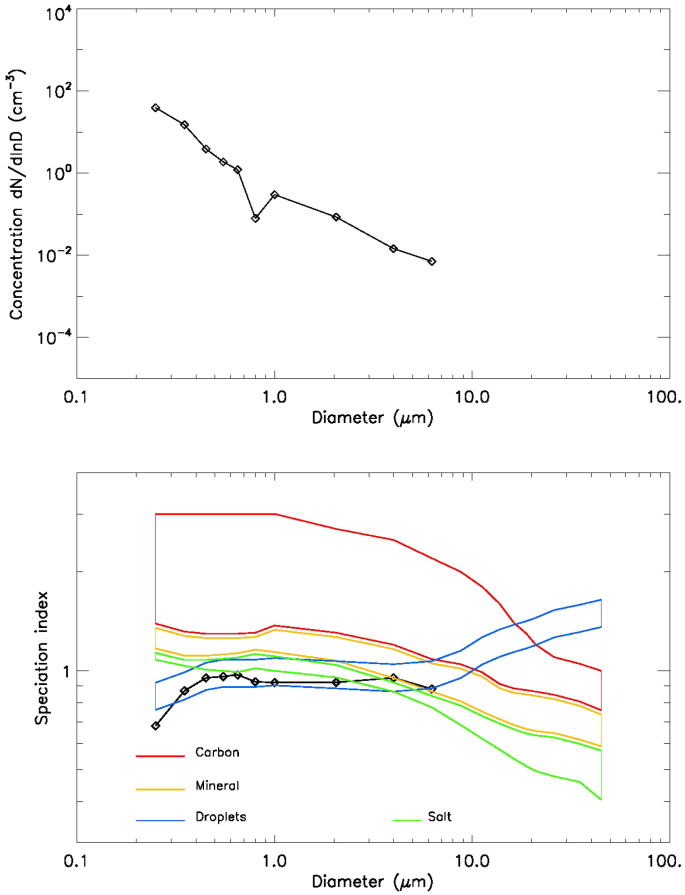

Figure 19. Example of measurements inside a haze or thin cloud at an altitude of 1.2 km during a flight under meteorological balloon from Reykjavik (Iceland) on 7 November 2013 at 12:30 UT; upper panel: size distribution; lower panel: speciation.

AMTD
8, 1203–1259, 2015

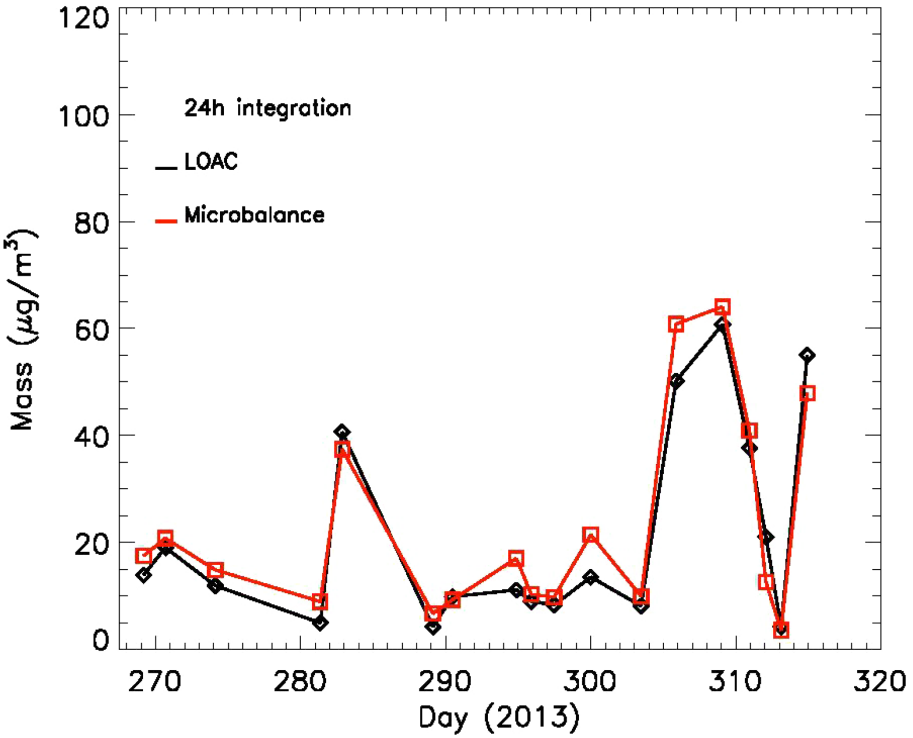

Figure 20. Comparison of coincident LOAC and TEOM microbalance measurement in indoor air (averaged over 24 h); particles have been injected with various concentrations to document a large range of mass concentration.

AMTD
8, 1203–1259, 2015

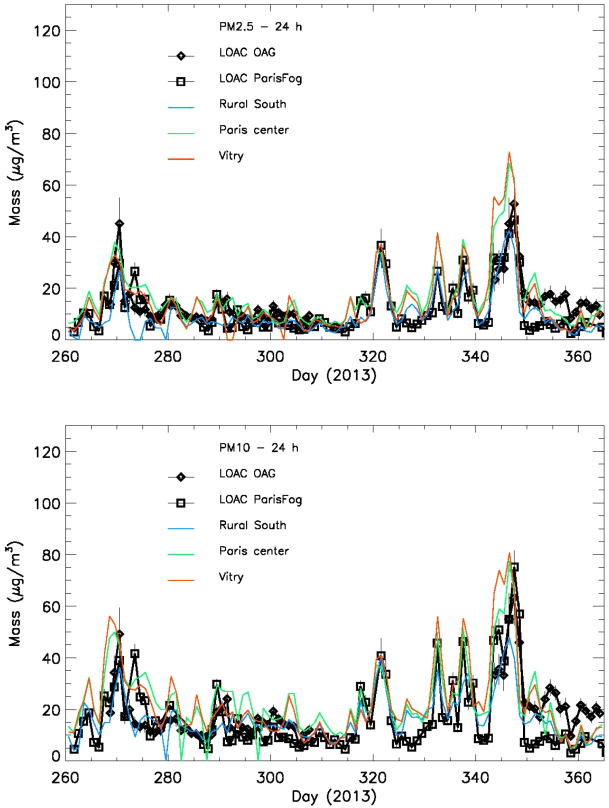

Figure 21. $PM_{2.5}$ (upper panel) and PM_{10} (lower panel) LOAC mass concentrations measurements in 2013 during the ParisFog campaign at SIRTA Observatory in Palaiseau, South of Paris, and at the Observatoire Atmosphérique Generali (OAG) in the South-West of Paris, and comparison with reference TEOM data from the AirParif air quality monitoring network.

AMTD
8, 1203–1259, 2015

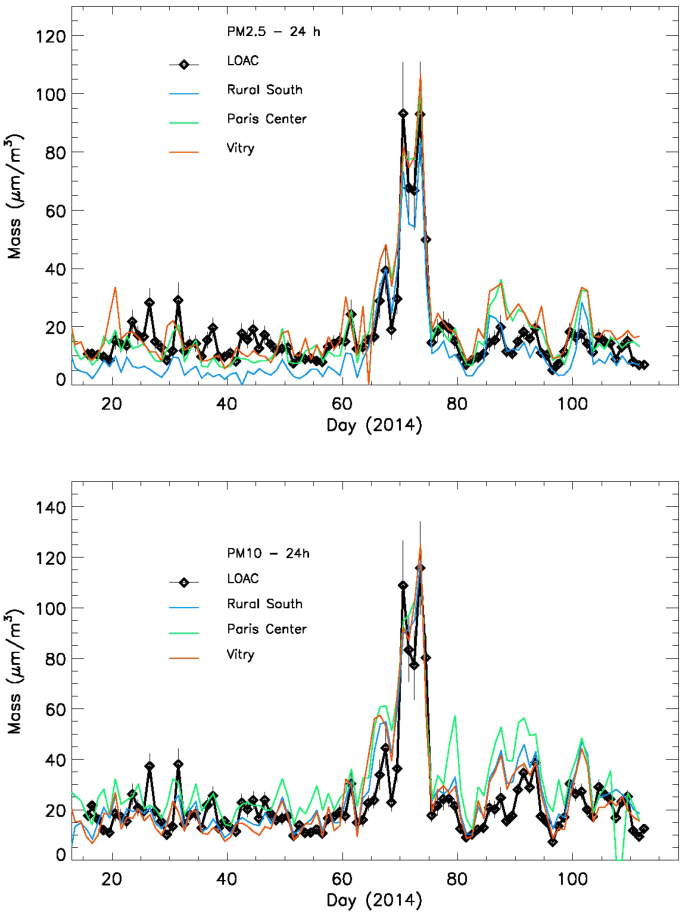

Figure 22. $PM_{2.5}$ (upper pannel) and PM_{10} (lower panel) LOAC mass concentrations measurements in 2014 at the "Observatoire Atmosphérique Generali" (South-West of Paris) and comparison with reference TEOM data from the AirParif air quality monitoring network.

AMTD
8, 1203–1259, 2015