

African volcanic emissions influencing atmospheric aerosols over the Amazon rain forest

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A  **bstract.**

The long-range transport (LRT) of trace gases and aerosol particles plays an important role for the composition of the Amazonian rain forest atmosphere. Sulfate aerosols originate to a substantial extent
30 from LRT sources and play an important role in the Amazonian atmosphere as strongly light-scattering particles and effective cloud condensation nuclei. The transatlantic transport of volcanic sulfur emissions from Africa has been considered as a source of particulate sulfate in the Amazon, however, direct observations have been lacking so far. This study provides expermental evidence for the influence of volcanic emissions on the Amazonian aerosol properties and atmospheric composition
35 during September 2014, based on comprehensive ground-based and airborne aerosol measurements

together with satellite observations. Under the volcanic influence, hourly mean sulfate mass concentrations in the sub-micron size range reached up to $3.6 \mu\text{g m}^{-3}$ at the Amazon Tall Tower Observatory, the highest value ever reported in the Amazon region. The substantial sulfate injection increased the aerosol hygroscopicity with κ values up to 0.36, thus, altering aerosol-cloud interactions over the rain forest. Airborne measurements and satellite data indicate that the transatlantic transport of the volcanogenic aerosols occurred in two major volcanic plumes with a sulfate-enhanced layer between 4 and 5 km altitude. This study demonstrates how remote aerosol sources in Africa, such as the volcanic sulfur emissions, can substantially affect the aerosol cycling and atmospheric processes over the Amazon rain forest.

45 **1 Introduction**

Sulfate aerosol particles are produced in the atmosphere by oxidation of sulfur dioxide (SO_2) emitted by fossil fuel (FF) combustion, volcanic emissions, and by oxidation of reduced sulfur compounds, such as DMS (Andreae and Rosenfeld, 2008). These particles scatter solar radiation and act as efficient cloud condensation nuclei (CCN) (Stevens and Feingold, 2009). Anthropogenic SO_2 emissions have increased over the 20th century to a maximum around the year 1980 and declined somewhat thereafter to around 100 Tg SO_2 per year, but they are still the most important source of sulfur to the atmosphere (Boucher et al., 2013; <http://edgar.jrc.ec.europa.eu/overview.php?v=431>, last access: 6 Sep 2017). Volcanic emissions are the predominant natural source of SO_2 and account for about 5 % of total annual SO_2 emissions (Textor et al., 2004; Yang et al., 2017). The emitted sulfur dioxide is oxidized in the atmosphere to gaseous sulfuric acid, which is quickly converted to sulfate aerosol particles (Chin et al., 1996; Reiner and Arnold, 1994). According to Chin and Jacob (1996), volcanic sulfur emissions can account for 20 – 40 % of sulfate aerosol particle mass concentrations in the troposphere and might therefore be able to substantially alter the atmospheric composition. Experimental evidence of drastic changes hundreds of kilometers downwind of volcanic eruptions affecting physical and chemical aerosol properties can be found in Mather et al. (2003) and Robock (2000, 2007). Two prominent examples are the Pinatubo eruption in 1991 that reached the stratosphere (> 10 km) (Kirchner et al.,

1999) and the 2014 – 2015 tropospheric effusive eruption of the Holuhraun volcano in Iceland (Ilyinskaya et al., 2017), where the emissions affected the cloud-drop effective radius (r_{eff}) while other cloud properties, like the cloud optical depth and the cloud liquid water path, remained unaffected (Malavelle et al., 2017; Yuan et al., 2011). Moreover, a connection between tropical volcanic explosive eruptions and El Niño-like events has been described recently (Khodri et al., 2017). Besides explosive-effusive eruptions, small eruptions and passive degassing account for relatively stable SO_2 fluxes ($23.0 \pm 2.3 \text{ Tg yr}^{-1}$, 2005–2015), and on average emit approximately one order of magnitude more SO_2 compared to explosive eruptions (Carn et al., 2017). To what extent volcanic passive sulfur emissions can affect cloud properties is still debated (Ebmeier et al., 2014; Malavelle et al., 2017).

There are several active volcanoes in Africa. The Nyamuragira-Nyiragongo neighboring volcanoes in the Democratic Republic of the Congo (DRC) were among the most persistent passively degassing volcanoes worldwide between 2004 and 2014, with around 150 days with satellite-detected degassing in 2014 and the highest average SO_2 index observed in the period 2004 – 2014 (Carn et al., 2016). The strong passive degassing activity of Nyamuragira starting in 2011 and culminating in the formation of a lava lake in late 2014 (Campion, 2014) lead to two orders of magnitude higher SO_2 mass emissions from the Nyiragongo-Nyamuragira complex (Barrière et al., 2017; Bobrowski et al., 2017). The emissions from Nyamuragira in 2014 have been estimated at ca. 2.5 Tg SO_2 (Fioletov et al., 2016).

In the Amazon rain forest, biogenic sulfate aerosol is sustained by oceanic DMS emission, and to a lesser degree by hydrogen sulfide (H_2S), methanethiol (MeSH), and DMS emissions from soils and vegetation (Andreae and Andreae, 1988; Jardine et al., 2015; Martin et al., 2010). Occasional anthropogenic sulfur injections have been attributed to open biomass burning and fossil fuel combustion emissions, either from Brazil or, via LRT, from Africa (Andreae et al., 1990; Talbot et al., 1988). Fossil fuel sources (e.g., ship traffic, power plants) do not often influence aerosol particles measured in the remote Amazon forest (Saturno et al., 2017) but can be important downwind of populated areas like Manaus, Brazil (Kuhn et al., 2010; Martin et al., 2016). Recent measurements during the South American Biomass Burning Analysis (SAMBBA) campaign, which focused on biomass burning (BB) emissions, found no correlation between sulfate aerosol and various kinds of BB aerosol particles (Brito et al., 2014). On the other hand, aircraft observations of haze layers at 2 – 4 km altitude over the

90 Amazon rain forest have shown high sulfate enrichment in comparison to the boundary layer and the free troposphere concentrations and indicated these haze layers to be linked to LRT of aerosol particles from Africa (Andreae et al., 1988; Holanda et al., 2018). A modeled global sulfate source attribution study showed that southern Africa peak sulfate concentrations occur between June and August. For this three-month period, estimated emissions were 0.81 Tg S and decreased in the following three months to 95 0.66 Tg S (Yang et al., 2017). Observations in the Amazon rain forest have shown that sulfate and organic aerosol (OA) mass concentrations can increase up to ten times from the wet to the dry season with rather stable sulfate-to-OA mass ratio throughout the year (Andreae et al., 2015; Fuzzi et al., 2007; Martin et al., 2010).

Northeasterly and southeasterly trade winds [north and south of the inter-tropical convergence zone (ITCZ)] are able to transport aerosols over large distances, given the typically weak wet deposition in this latitude band (Wang et al., 2016). During the Amazonian dry season (August – November), the transport of African smoke from southern Africa savanna and shrubland fires is an important source of aerosol in addition to the regional emissions (Andreae et al., 1994). Even though the potential impact of transatlantic transport of volcanic sulfur emission has been suggested before (Yang et al., 2017), no 105 ground-based evidence has been reported previously in the literature concerning the impact of African volcanic sources.

To analyze a major degassing period of the Nyamuragira in September 2014 and the subsequent transatlantic transport of the volcanic plume, this paper combines satellite and backward trajectory data with measurements at the Amazon Tall Tower Observatory as well as during the ACRIDICON-CHUVA 110 aircraft campaign in the Amazon Basin. This comprehensive data set indicates that the African volcanic SO₂ emissions were oxidized to particulate sulfate, which reached the central Amazon rain forest and influenced aerosol physical and chemical properties significantly.

2 Data and methods

2.1 Ground-based instrumentation

115 The ground-based aerosol data presented here have been collected at the Amazon Tall Tower
Observatory (ATTO) site [called *T0a* in the GoAmazon2014/5 experiment (Martin et al., 2016)], located
in the Uatumã Sustainable Development Reserve, Amazonas, Brazil. Details about the ATTO site
infrastructure, instrumentation and an overview of running measurements can be found in Andreae et al.
(2015). Figure 1 shows the ATTO site location and the location of the Nyamuragira volcano in the DRC,
120 located at 1.41° S, 29.2° E, 3058 m a.s.l. The long-term measurements at ATTO started in 2012. A
systematic backward trajectory (BT) analysis of air masses arriving at ATTO can be found in (C.
Pöhlker et al., 2018). The ATTO aerosol measurements were taken on a triangular mast (02° 08.602' S,
59° 00.033' W, 130 m above sea level, a.s.l.) using a 25 mm diameter, 60 m high stainless steel tube
with a laminar sampling flow rate of 30 L min⁻¹. The instruments were installed inside an
125 air-conditioned container where the temperature was kept between 29 and 31 °C. All aerosol data were
aggregated to hourly mean values. More details on the aerosol sampling setup can be found elsewhere
(Moran-Zuloaga et al., 2017; M. L. Pöhlker et al., 2016; Saturno et al., 2017).

Equivalent black carbon (BC_e) mass concentrations, M_{BC_e} , were calculated from absorption
measurements by a multi-angle absorption photometer, MAAP (Model 5012, Thermo Electron Group,
130 Waltham, USA). The details of the instrument are described elsewhere (Petzold and Schönlinner, 2004).
The BC mass absorption cross-section (MAC) was retrieved from fitting MAAP absorption coefficients
at 637 nm wavelength and refractory black carbon (rBC) mass concentrations measured by using a
single-particle soot photometer (SP2) revision D (Droplet Measurement Technologies, Longmont,
USA). Details of the technique can be found in Stephens et al. (2003). The MAC calculations are
135 described in Saturno et al. (2017). Light scattering coefficients were measured using a three-wavelength
integrating nephelometer (Aurora 3000, Ecotech Pty Ltd., Knoxfield, Australia). For details of the
instrument, see Müller et al. (2011). Absorption and interpolated scattering measurements at 637 nm
wavelength were used to calculate the single scattering albedo of dry aerosol particles, ω_0 , at this
wavelength, which is defined as the ratio of scattering to extinction coefficients (extinction = scattering

140 + absorption). Strong BB and FF influenced observations were defined as periods when M_{BCe} exceeded the 75th percentile ($0.27 \mu\text{g m}^{-3}$) and the absorption wavelength dependence was above or within the BC-only regime, respectively. A detailed discussion on the absorption wavelength dependence can be found in Saturno et al. (2017).

An aerosol chemical speciation monitor (ACSM) (Aerodyne Research Inc., Billerica, USA) was used to
145 measure online non-refractory aerosol chemical composition (Carbone et al., 2018). These measurements started in February 2014 at the ATTO site. The technique resolves the sub-micron aerosol chemical species in the following categories: Organics, sulfate, nitrate, ammonium and chloride (Ng et al., 2011). In this paper, we only use organics and sulfate mass concentration data, M_{org} and M_{sulfate} , respectively.

150 Cloud condensation nuclei (CCN) number concentrations, N_{CCN} , were measured with a CCN counter (CCNC, model CCN-100, Droplet Measurement Technologies, Longmont, USA), which was deployed at the ATTO site starting in March 2014. The instrument scanned over a range of different supersaturations and particle diameters; more details can be found elsewhere (M. L. Pöhlker et al., 2016). The hygroscopicity parameter, κ , retrieved for a CCN activation ratio of 50 % is used in this
155 study. Condensation nuclei number concentrations ($> 5 \text{ nm}$), N_{CN} , were measured with a condensation particle counter (CPC, model 5412, Grimm Aerosol Technik, Germany).

2.2 Airborne *in-situ* measurements

Chemical species of sub-micron aerosol particles were measured using a compact time-of-flight aerosol mass spectrometer (C-ToF-AMS) installed on board of the German High-Altitude and Long Range
160 Research Aircraft (HALO, <http://www.halo.dlr.de>, last access: 13 September 2017), a modified business jet G550 (Gulfstream, Savannah, USA). The C-ToF-MS details are presented in Drewnick et al. (2005) and Schulz et al. (2018). A single-particle soot photometer (SP2, Droplet Measurement Technologies, Longmont, USA) was used to measure rBC mass concentration. The measurements took place between 6 September and 1 October 2014, during the “Aerosol, Cloud, Precipitation, and Radiation Interactions
165 and Dynamics of Convective Cloud Systems” (ACRIDICON) - “Cloud Processes of the Main

Precipitation Systems in Brazil: A Contribution to Cloud Resolving Modeling and to the GPM (Global Precipitation Measurement)” (CHUVA) campaign over the Amazon rain forest (Machado et al., 2014). More details on the flight trajectories and instrumentation can be found in Wendisch et al. (2016). In this study, only data up to 7 km altitude have been used.

170 **2.3 Air mass trajectories**

To investigate the probability of the volcanic sulfate plume reaching the ATTO site, trajectories were calculated using the National Oceanic and Atmospheric Administration (NOAA) hybrid single-particle Lagrangian integrated trajectory HYSPLIT model (Draxler and Hess, 1997, 1998; Stein et al., 2015). NOAA Global Data Assimilation System (GDAS) (Kleist et al., 2009) data at $1^\circ \times 1^\circ$ resolution were
175 used as meteorological input for HYSPLIT.

2.4 Satellite SO₂ vertical column density data

As one of the most abundant gases in a volcanic plume, SO₂ is often used as a tracer for volcanic emissions by a variety of spectroscopic remote sensing techniques. The strong characteristic absorption features in the UV spectral range allow the quantification of SO₂ using differential optical absorption
180 spectroscopy (DOAS, see e.g., Platt and Stutz, 2008 and Richter and Wagner, 2011), both from the ground (e.g., Bobrowski and Platt, 2007; Galle et al., 2003) and from space (e.g., Eisinger and Burrows 1998, Khokhar et al. 2005 and Krueger, 1985).

The ozone monitoring instrument (OMI) on board of the National Aeronautics and Space Administration (NASA) Aura satellite, launched in 2004, detects backscattered solar radiation in the
185 UV-visible range (Levelt et al., 2006). The polar-orbiting instrument crosses the equator at 13:30 local time. DOAS analysis of OMI spectra yields column densities of trace gases such as NO₂, SO₂, and HCHO with a spatial resolution of about 13×24 km² away from the swath edges. OMI's wide swath of 2600 km allowed daily global coverage until the first occurrence of the so-called row anomaly in June 2007, an instrumental problem that causes grievous radiance errors in up to half of OMI's ground
190 pixels (Van Hoek and Claas, 2010). The row anomaly strongly affects the reliability of observations; therefore all affected pixels were removed from the data set prior to analysis.

The OMI SO₂ vertical column density (VCD) data presented in this paper were retrieved using NASA's principal-component based algorithm with an a-priori vertical profile representative of a volcanic plume in the middle troposphere (TRM, Li et al., 2013, 2017). The assumption that the volcanic plume is located in the mid-troposphere is justified by the elevation of the volcano (3058 m), the strength of the eruption, and, particularly, the HYSPLIT trajectory analysis presented later in this paper. It is, however, important to note that the sensitivity of the satellite measurements depends systematically on plume altitude. Thus, the absolute values of the SO₂ VCD derived from the satellite observations over- or underestimate the true values if the plume is located at a higher or lower altitude, respectively. Fortunately, this does not influence our study, as the focus of this paper is on the spatial pattern of the SO₂ plumes, and not on the SO₂ amount. The level-2 data were downloaded from: <https://mirador.gsfc.nasa.gov/> (last access: 27 October 2017) and gridded to a regular, 0.1° × 0.1° grid for easily handling.

3 Results and discussion

The Amazonian dry season (August – November) M_{sulfate} median over 3 years of measurements at the ATTO site was 0.6 $\mu\text{g m}^{-3}$ (0.41 – 0.79, inter-quartile range, IQR) as shown in Fig. 2a. This value slightly increased under the influence of BB (median: 0.83 $\mu\text{g m}^{-3}$) and did not change significantly during FF combustion influence (median: 0.56 $\mu\text{g m}^{-3}$). During African dust advection periods, when mineral dust particles are usually mixed with BB emissions and sea-salt aerosol particles, Moran-Zuloaga et al. (2017) measured sulfate mass concentrations of $\sim 0.44 \mu\text{g m}^{-3}$ using energy-dispersive X-ray analysis and M. L. Pöhlker et al. (2017) reported a M_{sulfate} mean of $0.25 \pm 0.19 \mu\text{g m}^{-3}$ measured by ACSM, both studies at the ATTO site. All these values and also measurements over the South Atlantic Ocean (Huang et al., 2018) were well below the M_{sulfate} median values measured at ATTO between 21 and 30 September 2014 (1.60 $\mu\text{g m}^{-3}$, see Fig. 2a). Regarding the single scattering albedo, $\omega_{0,637}$, shown in Fig. 2b, the lowest average was observed during FF influenced periods, indicating the presence of dark aerosol particles, rich in BC, which contrasts with the higher $\omega_{0,637}$ observed in September 2014. As can be seen in Fig. 2, the ATTO observations between 21 and 30 September 2014

are remarkably different from strong BB and FF combustion influence periods in terms of M_{sulfate} and $\omega_{0,637}$. Given this, the elevated M_{sulfate} observed in September 2014 could not be explained by combustion sources. Therefore, the possibility of an additional sulfate source, like volcanogenic sulfate aerosol, was considered. In the following section, satellite data is used to study the possibility of a volcanic plume reaching the Amazon rain forest during the period of interest. The last two sections are dedicated to the discussion of the aerosol physicochemical properties measured by aircraft- and ground-based instruments.

225 3.1 Satellite measurements and trajectory analysis of the volcanic plume

Given that the Nyamuragira and Nyiragongo volcanoes are so close to each other (within ~15 km), their emissions detected by remote sensors are often treated as a paired source (Carn et al., 2017). Hereafter, the term “Nyamuragira” refers to the couple Nyamuragira-Nyiragongo in this text. Time series of area-averaged OMI SO₂ observations are shown in Fig. 3a. The area where the average was calculated (Fig. 3b) corresponds to approximately 200,000 km² around the volcanoes. Nyamuragira produced frequent intensive SO₂ emission events in 2014 with a mean emission of 14.4×10^6 kg SO₂ day⁻¹ (Barrière et al., 2017). Multi-year OMI SO₂ VCD daily averages from 11° S to 17° N are summarized in Fig. S1 as a function of time and longitude. The figure shows observations during the ATTO measurement period (March 2012 to July 2017) and a snapshot of September 2014. According to Barrière et al. (2017), the emissions from June to October 2014 were mostly due to lava fountaining activity in Nyamuragira, characterized by strong tremors. The high emission activity of Nyamuragira at the end of 2014 was also observed by in-situ measurements from November 2014 to October 2015, when SO₂ emission fluxes reached up to 12.7×10^6 kg day⁻¹ from Nyamuragira (Bobrowski et al., 2017). These emissions were often transported westward, as can be observed in HYSPLIT forward trajectories calculated for 2014 (Fig. S2).

Fortunately, during the high emission period mentioned above, specifically in September – October 2014, ground and airborne measurement campaigns took place in the Amazon Basin (Andreae et al., 2015; Wendisch et al., 2016; Martin et al., 2017). Satellite SO₂ VCD observations over central Africa and the Atlantic Ocean were examined during this period in order to precisely identify the plume

245 trajectory. A map of gridded OMI SO₂ TRM VCD observations from 7 to 17 September 2014, is available as supplementary material (Fig. S3). Two important emission events were observed at the Nyamuragira location, one on 7 September and another on 12 September. The latter exhibits a clear westward transport of the SO₂ plume starting from 13 to 17 September. Fig. 4 shows SO₂ VCD observations during this period within the region between 20 W – 30 E, and 15 S – 5 N with SO₂ VCD larger than 2.5×10^{16} molecules cm⁻². Several sets of trajectory calculations were performed. First, three to seven starting locations were selected within the SO₂ plumes detected by OMI on 12–17 September 2014. At each location, 15-day (360 hours) forward trajectories were started at the time of the satellite overpass (11 – 14 UTC) at seven altitudes spaced equally between 1 and 7 km. The resulting trajectories initialized at 4 km altitude on 13 September are in best agreement with the satellite data and are shown in Fig. 4. All starting parameters were systematically varied and very consistent patterns were found (see Fig. 4). The trajectories are superimposed on the map presenting all SO₂ plumes detected by OMI between 12 and 17 September. Trajectories started within the leading edge of the plume are in good agreement with the OMI data, as after 24 hours (second marker) both trajectories are located within the plume detected on 14 September, and after 48 hours (third marker) both trajectories are located within the plume detected on 15 September.

This approach of using the OMI data to evaluate the trajectories was used also with the purpose of minimizing uncertainties by calculating shorter trajectories instead of initializing them from the volcano location. It should be noted here that the trajectory calculations by HYSPLIT yield a line, which can be understood as the center line of a propagating plume that widens both by stochastic uncertainty in the model calculations and by dispersive processes in the atmosphere. Consequently, the location of the plume becomes more uncertain the longer the model is run, and the physical size of the plume increases as well. Given the 9000-km distance between the volcanoes and ATTO, the uncertainty in the calculated plume trajectory position would become very large. To mitigate this problem, we use a multi-step approach, where we follow the emissions using the satellite-observed locations of the SO₂ plume for the initial days, and then calculate forward trajectories from the observed location of the plume.

The two southernmost trajectories in Fig. 4 make a sharp turn after 15 September, which is in agreement with the observed pattern, although there is no longer an exact match with the respective OMI

observations (in red and maroon). This discrepancy may be due to inaccuracy of the individual trajectories, or the SO₂ plumes might have been below OMI's detection limit. The southernmost trajectories reach South America and come within several hundreds of kilometers of ATTO within 15 days, which is well within the HYSPLIT uncertainty, estimated at 15-30 % of the trajectory length (<https://www.arl.noaa.gov/hysplit/hysplit-frequently-asked-questions-faqs/faq-hg11/>, last access: 19 March 2018). One of them reaches ATTO on 25 September at 1.8 km altitude, whereas the other one passed at an altitude of 1.5 km at the point nearest to ATTO on 24 September. As discussed above, the locations along the trajectory should be interpreted as the centers of a plume, which had dispersed both vertically and horizontally. This dispersion is not considered in the trajectory mode of HYSPLIT (Stein et al., 2015), but of course takes place in reality by turbulent and convective processes. Consequently, a trajectory “parcel” moving at 1.8 km altitude will certainly be dispersed to the surface in the highly convective Amazonian lower troposphere.

In addition to the plume forward trajectory analysis, backward trajectories initiated at the ATTO site at an altitude of 300 m (approximately 170 m above ground) were calculated for 360 hours. These trajectories were initiated every hour from 20 September 0:00 UTC up to 30 September 23:00 UTC. The results, presented as a trajectory density plot in Fig. 5, indicate that although during this time period essentially all air masses come from southern Africa, they usually come from further south. Nevertheless, a significant number of trajectories originates close to the volcano and its plume.

3.2 Airborne aerosol observations

Enhanced sulfate aerosol mass concentrations were observed above 3 km height over the Amazon Basin during the ACRIDICON-CHUVA campaign, which were substantially greater than the concentrations measured at lower altitudes for some flights. A map including all airborne observations on the different flights can be found in the supplementary material (Fig. S5). However, given the multiple sulfate aerosol sources, sulfate itself cannot be used as a tracer of volcanic emissions. In order to distinguish the volcanogenic sulfate from other aerosol sources like BB, which is important during this time of the year, we examined the M_{sulfate} vertical profiles together with their sulfate-to-OA mass ratio ($M_{\text{sulfate}} / M_{\text{org}}$). A

list of the ACRIDICON-CHUVA flights and M_{sulfate} vertical profiles are presented as supplementary
300 information in Table S1 and Fig. S6, respectively.

From the different airborne observations, the elevated plumes observed on flights AC14 and AC17
exhibited median sulfate-to-OA mass ratios larger than 1.0, indicating strong volcanogenic influence.
The M_{sulfate} vertical profile measured on 21 September 2014 (AC14) is presented in Fig. 6. The
observations show a sulfate-enhanced layer between 4 and 5 km height, which was also observed during
305 flight AC17 on 27 September 2014. The average M_{sulfate} observed during flight AC14 was
 $1.1 \pm 0.5 \mu\text{g m}^{-3}$ between 3 and 6 km height. This sulfate-enhanced layer exhibits a sulfate-to-OA ratio
generally larger than 1. It can be distinguished from lower layers, below 3 km height, which are
characteristically rich in OA due to BB and biogenic emissions. Usually, BB aerosol particles have been
shown to have higher OA mass concentrations than other aerosol particles (McNaughton et al., 2011;
310 Saturno et al., 2017). The concentrations of rBC in the region between 4 and 5 km were also very low (9
 ng m^{-3}), compared to the values below 3 km altitude (270ng m^{-3}), ruling out a combustion source of the
sulfate. In the light of all the observations (i.e., enhanced sulfate layer above 4 km height, high sulfate-
to-OA, very low rBC) there is no other plausible explanation for the source of this sulfate plume than
the LRT of volcanogenic aerosols.

315 Backward trajectories were calculated from several points along the flight paths. Figure 7 shows
backward trajectories started at nine points along the AC14 flight track, where sulfate-to-OA ratios
larger (colored lines) or smaller than 1 (gray lines) were detected; the flight and aerosol data measured
at each point are presented in Table 1. Figure 7 clearly shows that the colored trajectories, with one
exception, initialized within the sulfate plume originate from central Africa, whereas the gray
320 trajectories, started outside of the sulfate plume, appear to originate from South America or from more
southern regions over or across the Atlantic Ocean. The air mass trajectory analysis supports our
hypothesis that the AC14 observations were likely the result of probing the volcanic plume emitted on 7
September, the first one detected by OMI (see Fig. S3). For flight AC17 a similar pattern is observed,
with three out of four (colored) trajectories started within the sulfate plume originating from central
325 Africa and half of the other trajectories clearly coming from regions more to the South (see supplement,
Fig. S7 and Table S2).

Figures 4, 5, and 7 show that the trajectories agree well, but not perfectly with the ground-based, airborne, and satellite measurements, which is mainly caused by the uncertainty of such long trajectories. Nevertheless, the fact that forward and backward trajectories calculated from various starting points and times agree on the sulfate source is a further indication that the sulfate plumes observed at and near ATTO originate from the Nyamuragira volcano. Combined with the westward transport pattern derived from SO₂ satellite data and the lack of an alternative strong sulfate source makes this a quite convincing case.

3.3 Ground-based aerosol observations

The arrival of the African volcanic emissions over the Amazon rain forest is reflected in the aerosol physical and chemical properties measured at the ATTO site. The most evident effect was the significant increase in M_{sulfate} . The 90th percentile of M_{sulfate} measured at the ATTO site during the dry season 2014 was used as a threshold to define the volcanic influence event (Nya2014) as the time when this threshold was exceeded. By this criterion, the Nya2014 event spanned from 21 September 2014 at 02:00 UTC to 1 October 2014 at 01:00 UTC. Figure 8 shows different aerosol parameters measured before, during and after the Nya2014 event. The N_{CN} , shown in Fig. 8a, did not vary greatly from the values typical of the season (M. L. Pöhlker et al., 2016). The average N_{CN} during the dry season in 2014 was $(1.3 \pm 0.6) \times 10^3$ particles cm⁻³, whereas during the Nya2014 event, there were three peaks lasting for few hours with particle number concentrations higher than 3×10^3 particles cm⁻³ on 27, 29, and 30 September 2014, likely related to biomass burning (M. L. Pöhlker et al., 2017).

On the other hand, during the Nya2014 period, M_{sulfate} averaged $1.7 \pm 0.6 \mu\text{g m}^{-3}$, which was significantly above the dry season 2014 average of $0.7 \pm 0.3 \mu\text{g m}^{-3}$, see Fig. 8b. The highest M_{sulfate} value observed at the ATTO site was $3.6 \mu\text{g m}^{-3}$ (hourly mean) on 26 September 2014. For comparison, during the SAMBBA campaign in southern Amazonia, M_{sulfate} barely exceeded $1.0 \mu\text{g m}^{-3}$, despite organics nearly reaching $100 \mu\text{g m}^{-3}$, M_{BCe} of $5 \mu\text{g m}^{-3}$ and N_{CN} above 25×10^3 particles cm⁻³ during the peak of biomass burning (Brito et al., 2014). A long-term measurement study, also conducted in southern Amazonia, reported M_{sulfate} of $1.1 \pm 0.7 \mu\text{g m}^{-3}$ during the dry season, with a maximum of $3.3 \mu\text{g m}^{-3}$ for aerosol particles with mobility diameters smaller than $2 \mu\text{m}$ (Artaxo et al., 2002). It is

important to note that the observations by Artaxo et al. (2002) were done under strong BB influence
355 with average elemental carbon (EC) mass concentrations, M_{EC} , of $3.8 \pm 4.2 \mu\text{g m}^{-3}$, with a maximum of
 $25 \mu\text{g m}^{-3}$. In contrast, the BC_e measurements at the ATTO site during the Nya2014 event had an
average of $0.4 \pm 0.1 \mu\text{g m}^{-3}$, with a maximum of $0.8 \mu\text{g m}^{-3}$, indicating that the BB influence was
relatively weak during the period of interest, with some short (few hours) spikes due to the influence of
near-by fire events, see Fig. 8b. At a sampling site impacted by Manaus emissions, the sub-micron
360 M_{sulfate} was about $0.2 \mu\text{g m}^{-3}$ during the wet season, rarely exceeding $0.6 \mu\text{g m}^{-3}$ (de Sá et al., 2017).
Therefore, even considering a range of pollution sources, our measurements at ATTO during the
Nya2014 event are the highest sub-micron sulfate concentration ever reported in the Amazon Basin; see
Martin et al. (2010) for a summary of wet and dry season aerosol observations. For comparison, the
ACRIDICON-CHUVA airborne measurements are also included in Fig. 8b. The M_{sulfate} measured on
365 flight AC14 was significantly enhanced between 3 to 6 km altitude, reaching a median of $1.0 \mu\text{g m}^{-3}$ and
a 75th percentile of $1.6 \mu\text{g m}^{-3}$. Previous aircraft measurements during the SAMBBA campaign reported
a M_{sulfate} flight average of $0.48 \mu\text{g m}^{-3}$ (Allan et al., 2014).

The increased M_{sulfate} period was accompanied by an enhanced sulfate-to-OA mass ratio, according to
the ATTO observations (Fig. 8c). The Nya2014 sulfate-to-OA average over about 10 days was
370 0.38 ± 0.09 , significantly higher than the dry season average of 0.18 ± 0.15 . During some BB pulses,
decreased sulfate-to-OA ratios were observed, but the whole Nya2014 period was exceptionally high
compared to the typical dry season conditions. The sulfate-to-OA values measured at ground level were
usually lower than the airborne values observed between 3 and 6 km height because the OA sources
(BB and biogenic emissions) are located at ground level and the LRT sulfate that arrives at higher
375 altitudes is diluted upon vertical mixing. The possibility of FF burning was ruled out as an important
sulfur source during the event discussed here because of the particularly high dry-aerosol ω_0 measured
during the event (0.89 ± 0.04), as can be observed in the color code data in Fig. 8c. Usually FF
emissions, rich in BC, present characteristically low ω_0 (0.2 – 0.3) (Bond and Bergstrom, 2006) and its
addition would have lowered the value of ω_0 . Instead, an increase in ω_0 was observed to values higher
380 than 0.90 during the period of maximum M_{sulfate} (26 – 27 September 2014).

The effects of the volcanic sulfur plume on the aerosol hygroscopicity was explored by analyzing the κ values measured at different supersaturations. Higher κ values were measured for the accumulation mode aerosol (particles with diameter greater than 100 nm) (Fig. 8d; note the color-coded particle activation diameter, D_a). During the Nya2014 event, the κ values increased significantly, especially
385 when the maximum M_{sulfate} was observed. For example, the average κ for a supersaturation of 0.10 % ($D_a = 167 - 179$ nm, 25th and 75th percentile, respectively) was 0.26 ± 0.04 during the Nya2014 event, with a maximum of 0.36. A similar increase was observed at Jungfrauoch (Switzerland) under the influence of the Eyjafjallajökull volcanic emissions in 2010 (Bukowiecki et al., 2011). In that case the Aitken mode particles (10 to 50 nm diameter) were more strongly affected than the accumulation mode
390 particles because of the shorter ageing period of the sulfate particles. The Nya2014 κ average was slightly higher than the 2014 dry season average of 0.21 ± 0.03 for 0.10 % supersaturation (excluding the volcanic event), and significantly higher than a strong BB event average of 0.18 ± 0.01 for 0.10 % supersaturation, whose high OA content (sulfate-to-OA < 0.04) caused a significant drop in κ (M. L. Pöhlker et al., 2017).

395 **Summary and conclusions**

Satellite SO₂ observations showed two explosive events at the Nyamuragira volcano on 7 and 12 September 2014. These emissions were observed to be transported over the South Atlantic Ocean. Modelled air mass trajectories, starting at the satellite-observed plume locations, showed that the plume was transported towards South America, eventually reaching the region of the Amazon Tall Tower
400 Observatory in central Amazonia. Airborne observations during the ACRIDICON-CHUVA campaign probed a sulfate-enhanced layer between 4 and 5 km height on 21 September 2014 (flight 4). Additionally, this layer exhibited very low rBC concentrations and an increased sulfate-to-OA mass ratio with medians higher than 1 for measurements between 3 and 6 km height.

The ground-based M_{sulfate} measured at the ATTO site reached hourly mean levels up to $3.6 \mu\text{g m}^{-3}$ on
405 26 September 2014, the highest values ever reported in the Amazon Basin. The sulfate-to-OA mass ratio increased from a dry-season average of 0.18 ± 0.15 to an average of 0.38 ± 0.09 upon volcanic

influence, which spanned over about 10 days. Increased sulfate-to-OA and single scattering albedo (ω_0) suggest a rather low influence of BB and FF sources. In terms of aerosol hygroscopicity, the values of κ (for 0.10 % supersaturation) measured during the volcanic event reached an average of 0.26 ± 0.04 (and
410 a maximum of 0.36), the highest levels observed under dry season conditions (average of 0.21 ± 0.03).

The evidence presented here shows one particular event of volcanic SO_2 emissions in Africa influencing the aerosol particles' chemical composition, hygroscopicity, and optical properties in the Amazon Basin. Therefore, our study indicates that these emissions and their transatlantic transport could potentially affect the Amazonian cloud microphysical properties. However, the extent and relevance of the episodic
415 volcanic influence on the Amazonian atmosphere would require more extensive studies. Beyond the effects and implications of this particular event, the results of our study represent a reference case of the dynamics and conditions of transatlantic aerosol transport from southern Africa to South America. This could help to understand the inter-continental advection of other aerosol species, such as combustion aerosol particles that are more difficult to trace.

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Data availability. The data presented in this paper can be accessed via e-mail request to Jorge Saturno (j.saturno@mpic.de) or Christopher Pöhlker (c.pohlker@mpic.de). OMI data are available online at <https://disc.gsfc.nasa.gov/>.

425 *Competing interests.* The authors declare that they have no conflict of interest.

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Table 1. Measurements at points along track of the flight AC14 (21 September 2014) selected as starting points for backward trajectories presented in Fig. 7. Data points with sulfate-to-OA > 1 are emphasized by bold font.

Time (UTC)	Latitude [°N]	Longitude [°E]	Altitude [km]	M_{sulfate} [$\mu\text{g m}^{-3}$]	Sulfate-to-OA	Color in Fig. 7
15:14	-2.75	-60.34	1.61	1.2	0.5	Gray
16:19	-3.19	-60.21	4.50	1.0	1.1	Blue
16:36	-4.00	-59.50	4.50	3.0	2.9	Green
16:54	-5.20	-59.25	4.49	3.1	7.8	Red
17:11	-6.45	-58.98	6.45	0.6	0.8	Gray
17:46	-4.52	-59.40	6.43	0.5	0.8	Gray
19:40	-4.44	-59.41	7.68	1.2	1.0	Gray
21:41	-3.79	-59.55	4.79	1.8	2.2	Light blue
21:59	-3.04	-60.20	0.90	1.2	0.3	Gray

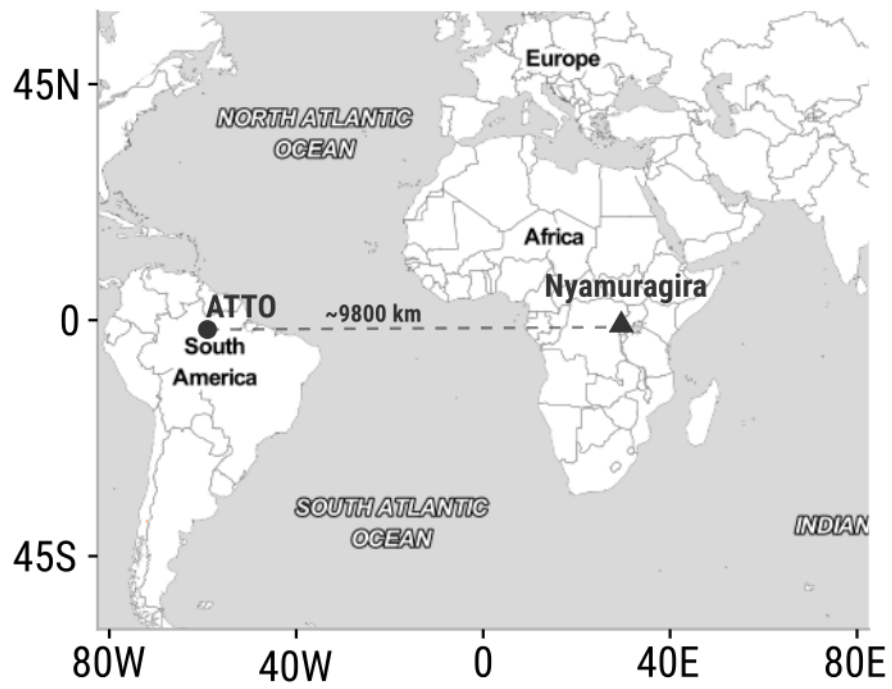


Figure 1. ATTO site and Nyamuragira volcano locations.

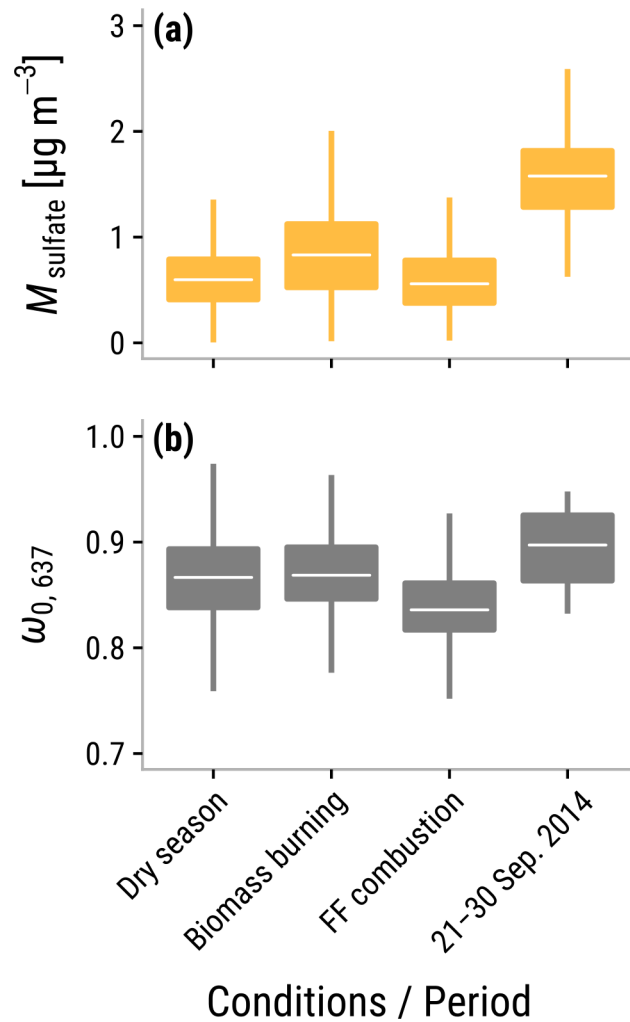


Figure 2. Box and whisker plots of hourly-mean (a) sulfate mass concentration, M_{sulfate} , and (b) single scattering albedo at 637 nm, $\omega_{0,637}$, during different periods and conditions including dry season average 2014 – 2016, biomass burning and fossil-fuel combustion influenced conditions and the period of interest from 21 to 30 September 2014. The white segment inside the box represents the median. Lower and upper box edges represent the first and the third quartiles, respectively. The whiskers represent the lowest and highest observations within the 99.3 % confidence interval.

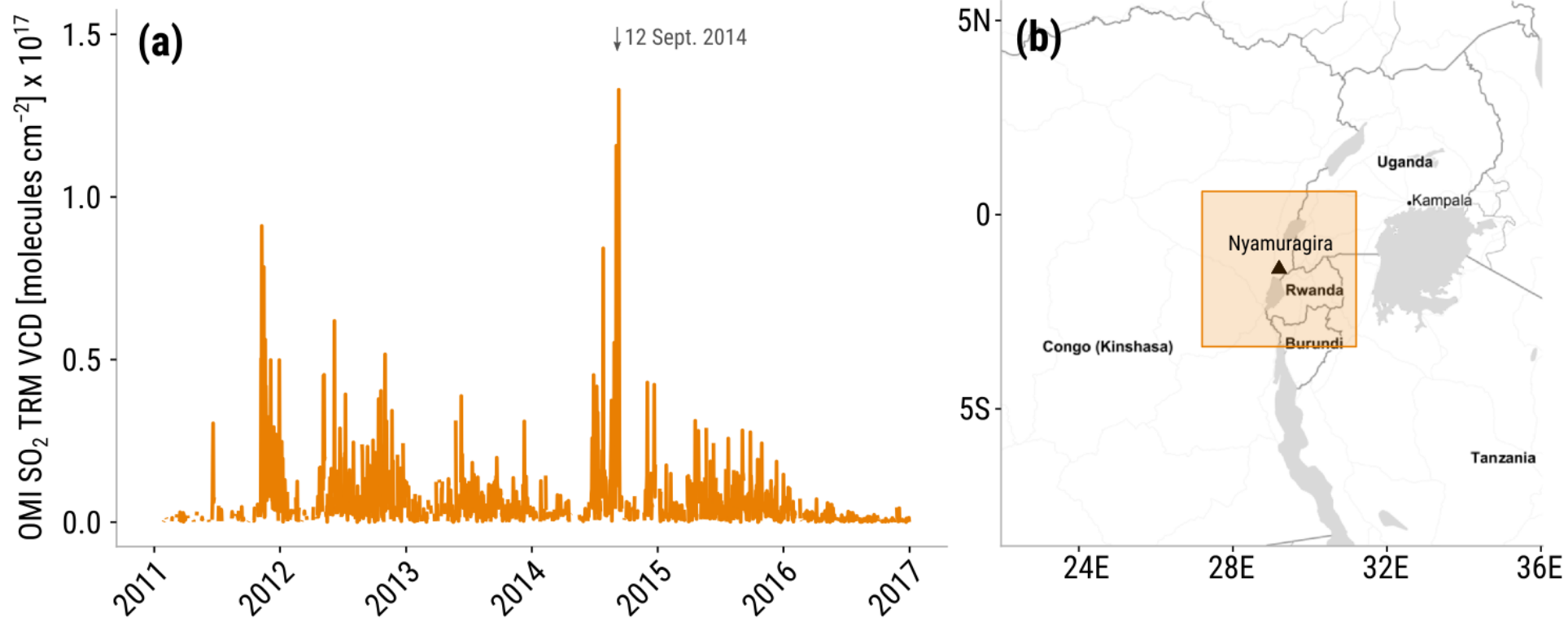


Figure 3. (a) Time series of daily-averaged OMI SO₂ TRM VCD observations corresponding to the averages over the area delimited by 27.2° E, 3.4° S, 31.2° E, and 0.6° N (b). Map of eastern Africa showing the averaging area (orange square). The location of Nyamuragira is represented by a black triangle.

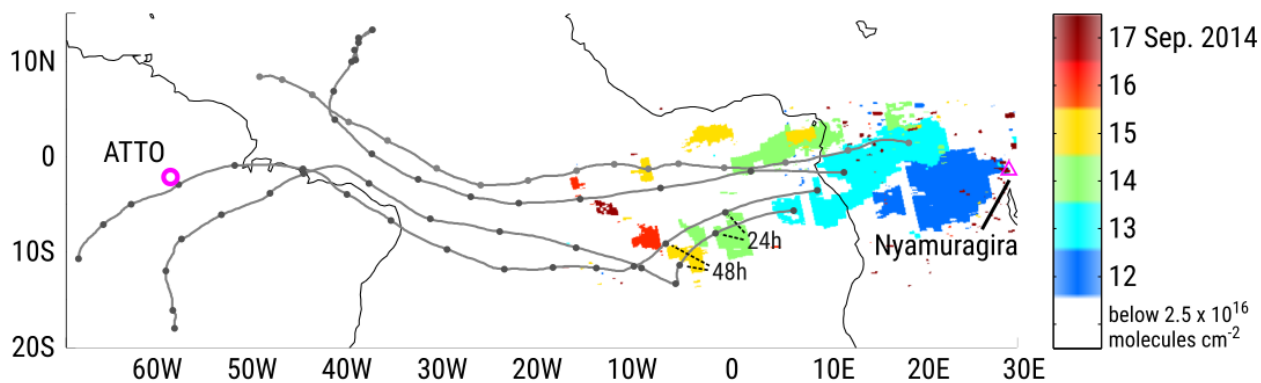


Figure 4. Map of SO₂ plumes with VCD $> 2.5 \times 10^{16}$ molecules cm⁻² color-coded by date of observation. 15-day forward trajectories started at 4 km (above mean sea level, a.m.s.l.) at four locations within the plume detected on 13 September 2014 (light blue) are indicated by black lines with markers at 24-hour intervals.

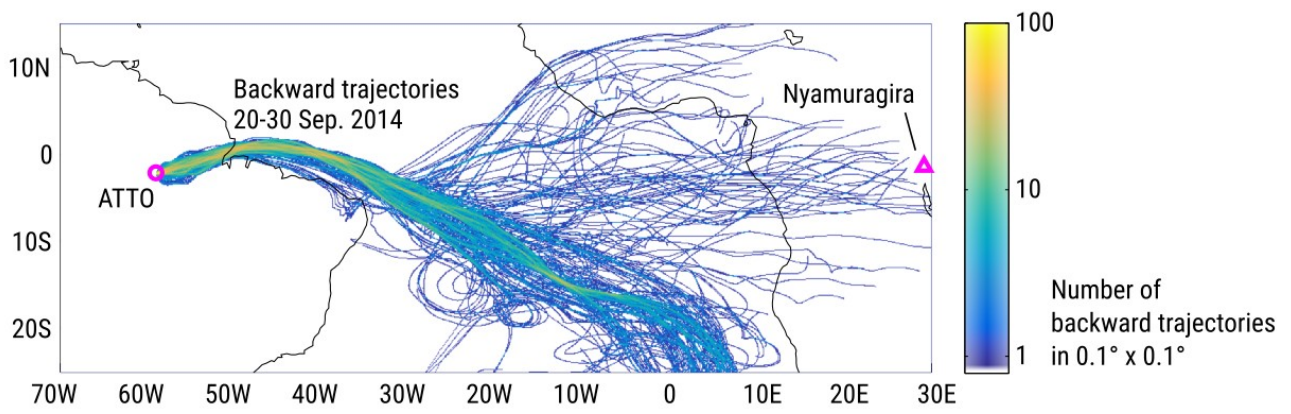


Figure 5. Density of 15-day backward trajectories started from ATTO at an altitude of 300 m a.m.s.l. on every hour starting at 0:00 UTC on 20 September 2014 up to 23:00 UTC on 30 September 2014.

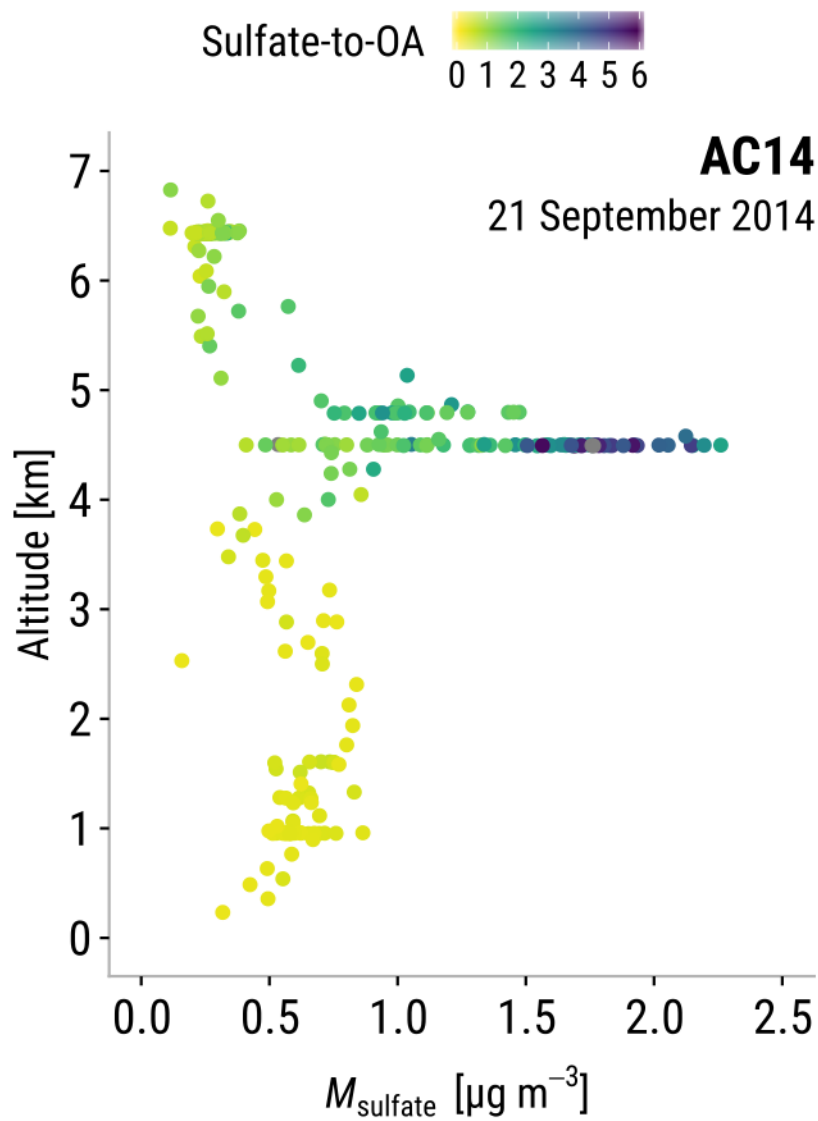


Figure 6. Sulfate layer observations over the Amazon rain forest. The figure shows the M_{sulfate} vertical profile observed during flight AC14 (21 September 2014). Color-coded sulfate-to-OA values are truncated at a maximum of 6.

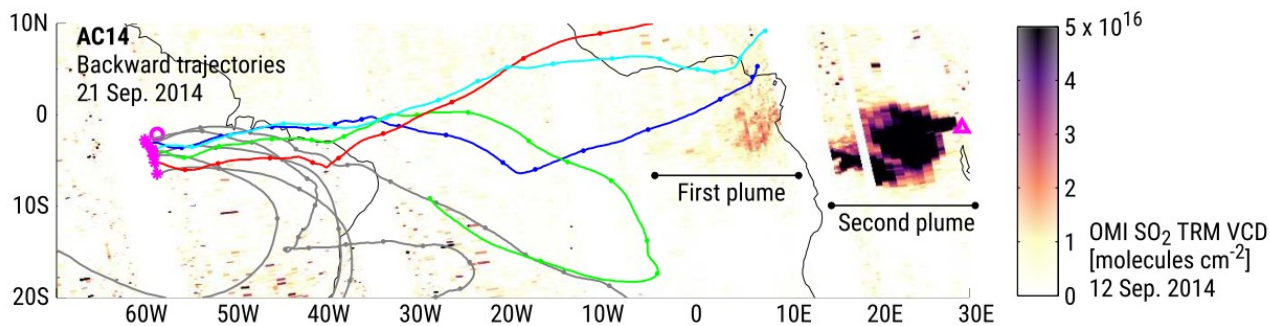


Figure 7. Map of gridded OMI SO₂ VCD, observed on 12 September 2014. Two plumes are shown on the map; the first one emitted on 7 September 2014, and the second one emitted on 12 September 2014. 15-day backward trajectories initialized at several points along track of the flight AC14 (21 September 2014) at flight altitude are shown as continuous lines with dots. Trajectories starting at points where sulfate-to-OA > 1 are shown in color (see Table 1 for details), all other trajectories are shown in gray; dots are placed at 24-hour intervals. The path of flight AC14 is marked in pink, with stars denoting the starting points of the backward trajectories. The locations of the ATTO site and Nyamuragira are marked with a pink circle and a triangle, respectively.

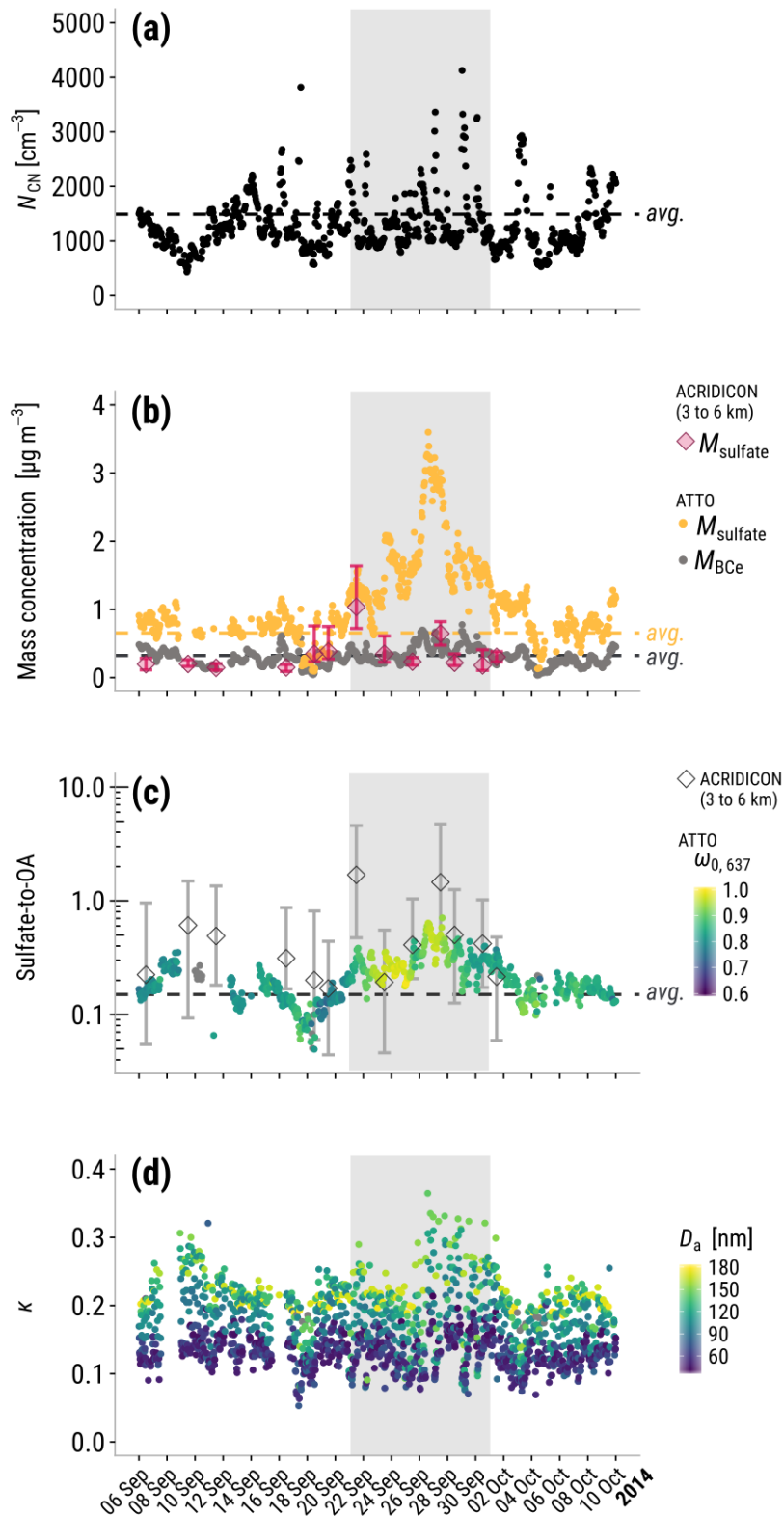


Figure 8. Different aerosol properties measured during the Nya2014 event (gray shaded area), These time series include ATTO hourly means of (a) aerosol particle number concentration, N_{CN} , (b) sulfate and BC_e mass concentration, (c) sulfate-to-OA mass ratio with color-coded aerosol particle single scattering albedo at 637 nm wavelength, and (d) hygroscopicity parameter, κ , original time resolution, with color-coded particle activation diameter. Dry season averages are shown as dashed lines. ACRIDICON-CHUVA flight medians and inter-quartile ranges from 3 to 6 km altitude are shown in (b) and (c).