



Depolarization ratio of smoke and volcanic ash aerosol particles at 1565 nm using a HALO Doppler lidar

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Received: 27 April 2026 – Discussion started: 30 April 2026

Revised: 29 July 2026 – Accepted: 12 August 2026 – Published: 17 August 2026

Abstract. Particle linear depolarization ratio is a widely used parameter in lidar research to distinguish different aerosol types and the thermodynamic phase of water. It is most frequently measured at ultraviolet and visible wavelengths (355 and 532 nm), yet multi-wavelength observations suggest that this parameter can vary substantially with wavelength. In this work, we assessed particle linear depolarization ratios at 1565 nm using Halo Photonics StreamLine Doppler lidars. We examined the depolarization ratio through three case studies featuring extremely fresh and aged smoke, and volcanic ash aerosol particles in the troposphere. Both fresh and aged smoke aerosol particles induced low values. Specifically, aerosol layers dominated by extremely fresh smoke showed a depolarization ratio of 0.017 ± 0.004 , whereas aged long-range transported smoke particles exhibited marginally higher values. Volcanic aerosol layers induced high depolarization ratios with layer mean values of 0.45 ± 0.01 . For the extremely fresh smoke case, we further estimated the smoke mass concentration using the lidar observations at 1565 nm and found good agreement with the in situ observations. These results demonstrate that Halo Doppler lidars operating at 1565 nm wavelength are capable of distinguishing several key aerosol types, enabling a comprehensive characterization of atmospheric conditions by simultaneously observing aerosol properties and wind dynamics.

1 Introduction

Aerosol particles exert a profound influence on climate, yet their effects remain among the largest sources of uncertainty in quantifying anthropogenic forcing (IPCC, 2021). Despite substantial advances driven by field measurements, laboratory studies, and modeling, the magnitude of this uncertainty has remained largely unchanged over the past two decades, with estimates of aerosol effective radiative forcing being far less constrained than those of greenhouse gases (Kahn et al.,

2023). Reducing this uncertainty requires, as a fundamental first step, to reliably detect aerosol particles in the atmosphere and accurately characterize their properties. Active remote sensing techniques, particularly lidars (Ansmann et al., 1990), provide this dual capability by first identifying the vertical distribution of aerosol particles and subsequently enabling the retrieval of intrinsic aerosol properties. One such property is the ratio of cross- to co-polarized backscattered signal known as the particle linear depolarization ratio (δ), after removing the molecular contribution. This parameter

has long been recognized as a powerful indicator of particle shape, allowing discrimination between spherical and non-spherical aerosol particles (Schotland et al., 1971; Sugimoto et al., 2002; Tesche et al., 2009), thereby offering critical insights into aerosol type characterization (Nicolae et al., 2018; Papagiannopoulos et al., 2018; Floutsi et al., 2024) and their individual role in air quality, interaction with clouds, and radiative forcing (Noh et al., 2012; Donovan et al., 2015; Xian et al., 2020; Cairo et al., 2024; Papayannis et al., 2025).

Historically, aerosol measurements on δ using lidars have been conducted across a wide range of wavelengths from ultraviolet (355 nm) and visible (532 nm) to less commonly used wavelengths at near-infrared (710, 1064 nm) (Sugimoto and Lee, 2006; Ansmann et al., 2009; Freudenthaler et al., 2009; Tesche et al., 2011; Groß et al., 2013; Pappalardo et al., 2013; Burton et al., 2015). Recent developments have expanded δ observations at 458 nm (Kong et al., 2022), and at longer wavelengths, including 808 nm (González et al., 2025), 910 nm (Bedoya-Velásquez et al., 2022) and 1565 nm (Vakkari et al., 2021). As an outcome, extensive ground-based and airborne campaigns at several locations have established reference δ values for key aerosol types like mineral dust (Sugimoto and Lee, 2006; Groß et al., 2011; Filioglou et al., 2020), volcanic ash particles (Ansmann et al., 2010; Pisani et al., 2012; Shimizu et al., 2025), pollen (Bohlmann et al., 2021; Filioglou et al., 2023), marine (Haarig et al., 2017) and smoke aerosol particles (Murayama et al., 2004; Müller et al., 2005). These studies conclude that non-spherical aerosol particles such as mineral dust, volcanic ash and pollen can maintain high depolarization ratio at longer wavelengths, while smoke and marine aerosols typically induce much lower depolarization ratio, presenting similar or decreasing values with increasing wavelength.

Despite these advances, several gaps remain. Long-term, harmonized datasets at longer wavelengths (particularly 910, 1064 and 1565 nm) are only beginning to be systematically explored and the spectral dependence of δ for many natural aerosol types and mixtures of these require further investigation, particularly under varying relative humidity (RH) conditions and in under-sampled regions. Moreover, case studies demonstrate that δ and its spectral dependence is sensitive to particle size, shape, composition and mixing state. For example, Freudenthaler et al. (2009) studied Saharan desert dust aerosols at 532 and 1064 nm and concluded that δ tend to be slightly higher at 532 nm compared to the longer wavelength. In contrast, Sugimoto and Lee (2006) concluded the opposite for Asian dust particles. In line with Freudenthaler et al. (2009), Haarig et al. (2022) found that mineral dust δ values are slightly higher at 355 and 532 nm compared to the longer wavelength of 1064 nm. In turn, Vakkari et al. (2021) reported higher values at 1565 nm than at 355 and 532 nm for a mineral-dust-dominated case while for dust mixed with other aerosols they reported higher depolarization ratio at 1565 nm compared to the shorter wavelengths. For pollen aerosol particles, Filioglou et al. (2023) concluded that birch pollen par-

ticles induce higher δ at 910 and 1565 nm compared to 355 and 532 nm. The opposite is valid for pine pollen particles. They also found that δ is concentration dependent, therefore, the longer wavelengths offer an advantage in the detection of large particles such as pollen particles. For marine and polluted marine aerosols, previous studies report a decreasing trend in δ with increasing wavelength (Vakkari et al., 2021; Haarig et al., 2025). Moreover, Haarig et al. (2018) studied tropospheric smoke particles and found that they induce a less than 3 % δ at 355, 532 and 1064 nm, while higher δ values were reported for stratospheric smoke layers at shorter wavelengths (Burton et al., 2015; Haarig et al., 2018; Hu et al., 2019). To the authors' knowledge, there are no studies of δ at 1565 nm for volcanic ash and smoke aerosol particles.

Multi-wavelength lidar observations, obtained from single or multiple instruments, enhance the reliability of aerosol characterization and typing. In this context, this study estimates the δ for smoke and volcanic ash particles using a Halo Doppler lidar operating at 1565 nm. Furthermore, we extend the applicability of such lidar systems by retrieving smoke aerosol mass concentration at this wavelength. Such estimates are particularly valuable, as they provide vertically resolved, model-ready information that improves the representation of aerosol loading, plume dynamics, and radiative effects in atmospheric models (Benedetti et al., 2009; Sekiyama et al., 2010; Zhang et al., 2014). By addressing the current lack of δ observations at 1565 nm, this study supports the integration of this wavelength into operational aerosol monitoring networks, benefiting from the negligible contribution of molecular scattering and absorption, which simplifies signal processing and enhances retrieval robustness.

2 Materials and Methods

Three field campaigns were conducted over the years of 2024, 2020 and 2016 at three different locations: at a Boreal forest site at Kiviniemi, Finland (62°29'06.0'' N, 27°29'27.4'' E, 110 m above sea level), at an urban site at Kumpula (Helsinki), Finland (60°12'13.5'' N, 24°57'39.1'' E, 45 m a.s.l.) and at a grassland savanna site at Welgegund, South Africa (26°34'11.3'' S, 26°56'21.5'' E, 1480 m a.s.l.), respectively. In each location, a HALO Photonics Streamline Doppler lidar was operating. To characterize the δ_{1565} of extremely fresh and aged smoke as well as volcanic ash particles, three representative case studies, one from each field campaign, were selected and presented in Sect. 3. The extremely fresh smoke case was observed in Kiviniemi (FI), the aged smoke case in Kumpula (FI) and the volcanic ash case in Welgegund (SA). For the extremely fresh smoke case in Kiviniemi (FI), a suite of ground- and drone-based in situ instruments were employed to quantify aerosol physical and chemical properties to support the mass concentration estimation of smoke from the lidar observations. To exclude the influence of biological particles on the analysis, given that

Table 1. Specifications of Halo Doppler lidars used in this study.

Wavelength	1565 nm
Pulse repetition rate	15 kHz
Pulse energy	20 μ J
Pulse duration	0.2 μ s
Nyquist velocity	20 m s ⁻¹
Sampling frequency	50 MHz
Velocity resolution	0.038 m s ⁻¹
Points per range gate	10
Range resolution	30 m
Maximum range	9600 m
Lens diameter	8 cm
Lens divergence	33 μ rad
Telescope	monostatic optic-fibre coupled

the measurements in Kiviniemi (FI) coincided with the active pollen season, a Burkard sampler was operated to monitor airborne pollen concentrations in the near-by station in the city of Kuopio (62°53′31.5″ N, 27°38′01.0″ E).

2.1 Lidar observations

Halo Photonics StreamLine lidars are commercially available pulsed Doppler lidars that operate at 1565 nm wavelength (Pearson et al., 2009). Due to the high pulse-repetition rates and low pulse energies they are eye-safe, enabling continuous long-term operation. Here, we use Halo Doppler lidars that are equipped with a cross-polar receiver channel, which enables consecutive measurement of co- and cross-polar signals and subsequent retrieval of particle linear depolarization ratio (Vakkari et al., 2021). The cross-polar channel is implemented through a fibre-optic switch between the normal receiver path and the path with a fibre-optic polarizer. The lidars were configured with 30 m vertical resolution and focus was set to 2000 m; first 90 m were excluded due to effects from the outgoing pulse. Other technical specifications can be found in Table 1.

At Kumpula (FI) and Welgegund (SA), the Halo Doppler lidars used were capable of full hemispheric scanning (i.e., model StreamLine). At Kiviniemi (FI), the version with limited scanning (i.e. StreamLine Pro) was used. In this study, we utilize only the vertically-pointing measurements. At Kiviniemi (FI), the vertically-pointing integration time was set to 7 s per ray and 13 min out of every 15 min was vertically-pointing measurement. At Kumpula (FI), the vertically-pointing integration time was set to 3.5 s per ray and 8 min out of every 15 min was vertically-pointing measurement. At Welgegund (SA), the vertically-pointing integration time was set to 7 s per ray and 6 min out of every 15 min was vertically-pointing measurement.

For each Halo Doppler lidar the data was post-processed following Vakkari et al. (2019). The post-processing is based on hourly background checks carried out automatically by

the lidar. Quantifying the non-polynomial component in the instrument noise floor, denoted P_{amp} by Vakkari et al. (2019), requires an extended data set of background checks and internal temperature. We used background checks from the full 11-month campaign at Welgegund (SA), and 12 months centered at the case study for the two other lidars, respectively. We determined also the internal temperature dependency of P_{amp} , though it has only a very minor effect. For each measured profile, we used the aerosol- and cloud-free parts of the profile to fit the linear and parabolic components of the instrumental noise floor. These fits and the instrument and campaign specific P_{amp} were then used to process the corrected co- and cross-polar signal-to-noise ratio (SNR) and attenuated backscatter profiles following Vakkari et al. (2019).

Bleed-through, i.e. the incomplete extinction in the lidar internal polarizer where co-polarized signal is leaking into cross-receiver due to imperfect polarization separation in the instrument, was determined from liquid cloud base observations similar to Le et al. (2024). For Kiviniemi (FI), a bleed-through of 0.0165 ± 0.010 was determined. For Kumpula (FI), bleed-through was determined as 0.00715 ± 0.0028 . For Welgegund (SA), bleed-through was determined as 0.0085 ± 0.007 . These values are in good agreement with the long-term statistics in Le et al. (2024).

At 1565 nm wavelength, the molecular backscatter contribution is about 2 orders of magnitude lower than at 532 nm, being approximately $1.9 \times 10^{-8} \text{ m}^{-1} \text{ sr}^{-1}$ at standard pressure and temperature (Bucholtz, 1995) which is considered negligible. In addition, no evidence of multiple scattering was observed under the prevailing measurement conditions. Therefore, δ_{1565} as derived directly from the instrument after correcting for the bleed-through, can be regarded as a reasonable approximation of the particle linear depolarization ratio, obtained without applying a correction for the molecular scattering contribution, especially during high aerosol load conditions. The associated uncertainty, $\sigma_{\delta_{1565}}$, is estimated according to Vakkari et al. (2021) taking into consideration the instrumental noise and the uncertainty in bleed-through. In the absence of hydrometeors, the atmospheric transmittance is very close to unity therefore, the attenuated backscatter coefficient, hereafter ATB_{1565} , can be used as a proxy for the particle backscatter coefficient, hereafter β_{1565} . This assumption is valid provided that extinction within the sampled aerosol layer remains sufficiently small. Attenuation in dense aerosol layers can lead to an underestimation of the true β_{1565} . For the exceptionally fresh smoke case at Kiviniemi (FI), β_{1565} retrieved using the forward Klett method (Wiegner and Geiß, 2012) as the mass concentration parameter requires an accurate measure of backscatter intensity and could therefore be biased by substantial departures of atmospheric transmittance from unity observed during periods of high plume density. A Lidar Ratio (LR) of 51 sr was used in Klett inversion according to Mie calculations (see Sect. 2.4).

To determine the upper boundary of the smoke layer in the lidar observations at Kiviniemi, we first identified the maximum of the ATB_{1565} profile within the lowest 500 m. The layer top was then defined as the first altitude above this maximum at which the δ_{1565} exceeded 0.1. If no such threshold was reached, the layer top was instead assigned to the altitude corresponding to the maximum of the ATB_{1565} profile. For the Welgegund case, the maximum of the δ_{1565} profile was first located between 2 and 4 km and then the base and top boundaries were determined using the second derivative of ATB_{1565} . For the Kumpula case, ATB_{1565} observations with values lower than $0.2 \text{ Mm}^{-1} \text{ sr}^{-1}$ or higher than $2 \text{ Mm}^{-1} \text{ sr}^{-1}$ were excluded. The remaining observations were subsequently used to define the boundaries of the smoke layer. In all cases, cloud-free layers were considered only.

2.2 Particle mass concentration from lidar observations

Lidar-derived mass concentration (m) has been estimated using the methodology presented in Ansmann et al. (2011). The method requires the mass particle density (ρ), the extinction-to-volume conversion factor (c_v), and the particle extinction coefficient $\alpha(\lambda) = \beta(\lambda) \cdot LR(\lambda)$ for a specific aerosol type at a certain wavelength (λ) to be known according to $m = \rho \cdot c_v(\lambda) \cdot \alpha(\lambda)$. A density of $\rho = 1.35 \pm 0.05 \text{ g cm}^{-3}$ was used for smoke particles (Reid et al., 2005).

To derive the mass concentration, we first converted the β_{1565} into β_{532} using a Backscatter-related Ångström exponent (BAE) according to $\beta_{532} = \beta_{1565} \cdot (532/1565)^{-BAE}$. The BAE, as well as, the LR at both 532 and 1565 nm wavelengths were estimated through Mie calculations (see Sect. 2.4). This wavelength conversion enables us to use a literature value of $0.16 \pm 0.01 \times 10^{-12} \text{ Mm}$ for c_v at 532 nm (Ansmann et al., 2026), which has not been determined at 1565 nm wavelength, making the mass concentration estimation possible.

2.3 In situ aerosol observations

For the extremely fresh smoke event in Kiviniemi (FI), additional ground- and drone-based in situ aerosol observations were considered. The aerosol size distribution in size range of 15–710 nm was measured with a Scanning Mobility Particle Sizer (SMPS). The SMPS setup (TSI Inc., Model 3080) consists of a DMA (TSI Inc., Model 3081) and a CPC (TSI Inc., Model 3776) and temporal resolution was 3 min. Aerosol particle size distribution in the range between 0.45 and $15.5 \mu\text{m}$ was measured using an optical particle counter sensor (OPC-N2, Alphasense Ltd.; e.g., Crilley et al., 2018; Hagan and Kroll, 2020; Julaha et al., 2025) mounted onboard a DJI Matrice 600 drone. The OPC was housed within a custom-built module, similar to Brus et al. (2025), designed for uncrewed aerial vehicles. Measurements were obtained by flying the drone horizontally across the smoke plume at

multiple altitudes ranging from 50 to 110 m. There was a total of four flights, at 11:30, 12:00, 16:15 and 16:45 UTC, each lasting up to 20 min. Drone data from the first two flights are considered only, as these periods exhibited spatially consistent smoke conditions between the airborne and surface-based observations.

To crosscheck and inter-compare the measurements of OPC and SMPS, the drone platform was positioned adjacent to the measurement station. The inter-comparison was performed at 14:30–15:05 UTC and the size distributions from OPC and SMPS were averaged over this period. The intercomparison showed an approximately twofold difference in particle number size distributions, with lower concentrations measured by the OPC on the drone platform than by the SMPS at the measurement station. This discrepancy is not unexpected, given the different measurement principles of the two instruments, their detection efficiencies and the strong spatial and temporal variability within the smoke plume. Sousan et al. (2016) reported that the OPC-N2 performs well in measuring coarse particles but tends to underestimate number concentrations of submicrometer particles, with a detection efficiency of 78 % for particles smaller than $0.5 \mu\text{m}$.

To derive the complete aerosol size distribution, the SMPS and OPC size distributions (hereafter SMPS-OPC) were averaged over the time intervals during which the OPC was operating (i.e., the flight periods), and the corresponding 10th and 90th percentiles were calculated over these overlapping intervals. From the combined SMPS-OPC aerosol size distribution, the total mass of aerosol particles was estimated using a particle density of 1.35 g cm^{-3} .

To gain a more detailed understanding of the aerosol composition, and particularly the black carbon (BC) and organic aerosol (OA) content during the burning event, ground-based multi-angle absorption photometer (MAAP, model 5012, Thermo Scientific, Petzold and Schönlinner, 2004) and aerosol mass spectrometer (AMS, HR-ToF-AMS, Aerodyne Research Inc., DeCarlo et al., 2006) measurements from the container (Fig. 1c) were used to estimate the BC-to-organic aerosol ratio (BC/OA). MAAP measured dried aerosol particles from the station's main inlet at a flow rate of 7.5 LPM at 1 min time resolution. The aerosol absorption determined by MAAP was converted to BC mass concentration using the instrument manufacturer set default mass absorption cross-section (MAC) of $6.6 \text{ m}^2 \text{ g}^{-1}$, which is commonly accepted value for MAAP and broadly used in scientific literature (e.g. Ohata et al., 2021). AMS measures the mass concentration of organics, sulfate, nitrate, ammonia and chloride compounds and was operated in EI-V mode (tungsten vaporizer) at 2 min time resolution. The instrument was calibrated on-site with ammonia sulfate and ammonia nitrate aerosols to determine the ionization efficiency of nitrate particles and relative ionization efficiency (RIE) of ammonia and sulfate. AMS data analysis was performed using standard tools – SQUIRREL v1.65C and PIKA v1.25C, downloaded from

the ToF-AMS-Resources web page (<https://cires1.colorado.edu/jimenez-group/ToFAMSResources/>, last access: 14 August 2026), in Igor Pro.

SMPS, AMS and MAAP in situ aerosol observations were averaged for 15 min periods to match the temporal resolution in the lidar observations. Regarding the OPC data, only observations inside the smoke plume were considered. For each flight, we derived an average aerosol size distribution by combining measurements across all sampled altitudes.

Pollen concentrations were assessed with a Burkard sampler of Hirst design (Hirst, 1952) at Kuopio (62°53′31.5″ N 27°38′01.0″ E). The site is part of the Finnish pollen network operated by the University of Turku (<https://sites.utu.fi/siitepoly/toiminta/>, last access: 14 August 2026) and member of the European Aeroallergen Network. For the derivation of the pollen types and their respective concentrations, the recommendations of the European Aeroallergen Society (Galán et al., 2014) were followed to ensure high data quality. Pollen concentrations were utilized in the extremely fresh smoke case in Kiviniemi (FI). The distance between the two sites is about 50 km, and the temporal resolution of the observations is 2 h.

2.4 Mie calculations of smoke particles

Mie-theory predicts the optical properties of spherical particles (Mie, 1908). Here, Mie model was used to calculate the β and α at the lidar wavelengths (355, 532, 1064 and 1565 nm) in order to infer the LR and BAE from the measured in-situ aerosol particle size distributions. Mie model calculations were done using the Python-package PyMieScatt (ver 1.8.1.) (Sumlin et al., 2018a).

To perform this analysis, the particles were assumed spherical without core-shell structures. A dry aerosol was also assumed, as the surface relative humidity (RH) remained below 40 %. The assumption of spherical particles required for Mie theory is supported by the low δ in the lidar observations during the smoke event. From the in-situ observations, the average mass BC / OA-ratio during the plume was approximately 1 %.

The complex refractive indices used were $1.5 + 0.0366i$, $1.5 + 0.0064i$, $1.5 + 0.0003i$, $1.5 + 0.0001i$ at 355, 532, 1064, and 1565 nm, respectively. The imaginary part (k) was derived according to Mukherjee et al. (2025), who measured the k for fresh boreal forest surface smoldering emissions. A wavelength dependence of k was assumed using their measurements at 365 and 550 nm. The methanol soluble fraction accounted for more than 90 % of the total organic aerosol mass, and was therefore considered representative of the bulk organic aerosol. In contrast, the methanol insoluble fraction was negligible in these fresh emissions. The real part (n) was treated as constant at a value of 1.5 according to Sumlin et al. (2018b), who observed that the n was independent of the λ and was between 1.5–1.7.

The absorption by BC was taken into account by assuming an externally mixed aerosol. BC size distribution measured in a chamber for fresh boreal forest surface was used (Peltokorpi et al., 2026; Vakkari et al., 2026). The BC size distribution was scaled so that the mass of BC matched the BC / OA-ratio (about 1 %) of the mass of particles smaller than 1 μm . The BC density was assumed to be 1.80 g cm^{-3} . The mass of particles smaller than 1 μm was estimated from the derived size distribution assuming density of 1.35 g cm^{-3} . The complex refractive index for BC was defined according to Eqs. (6) and (7) in Kahnert and Kanngießer (2020).

2.5 FLEXPART air mass trajectories

Air mass history was simulated using the Lagrangian particle dispersion model FLEXPART (FLEXible PARTicle dispersion model) version 11.0 (Bakels et al., 2024). ERA5 re-analysis from European Centre for Medium-Range Weather Forecasts (ECMWF) was used as meteorological input (Hersbach et al., 2023). The ERA5 input data was retrieved at 1 h temporal, 0.25° latitude–longitude resolution and for model levels 40 to 137, which corresponds to approximately the lowest 24 km in the atmosphere. FLEXPART was run in backward simulation mode, with potential emission sensitivity (PES, s^{-1}), describing the sensitivity of the sampled air mass to emissions along its transport history, output at 1 h temporal resolution. PES output vertical resolution was 500 m up to 6 km a.g.l. and remaining PES above 6 km was integrated into one additional output layer.

For the aged smoke case in Kumpula, Finland on 22 September 2020 the FLEXPART simulation retroplume was released at height interval of 3 to 4 km above ground level (a.g.l.) and time interval of 18:00 to 19:00 UTC, respectively. FLEXPART backward simulation was computed for 16 d. For the volcanic ash particles in Welgegund, South Africa on 17 December 2016 the FLEXPART simulation retroplume was released at height interval of 2.5 to 3.2 km a.g.l. and time interval of 04:00 to 05:00 UTC, corresponding to the lidar observation of the elevated layer. The duration of the FLEXPART backward simulation for this event was 14 d.

2.6 MODIS fire radiative power

The fire radiative power (FRP) product from MODIS Aqua and Terra, collection 6.1, (Giglio et al., 2016) was downloaded through the FIRMS interface for the period 7 to 20 September 2020 covering the area of North America to examine the locations of wildfires in the path of the air mass for the Kumpula case.

3 Results

In this section, we present the observations of extremely fresh and aged smoke and volcanic ash particles. Extremely

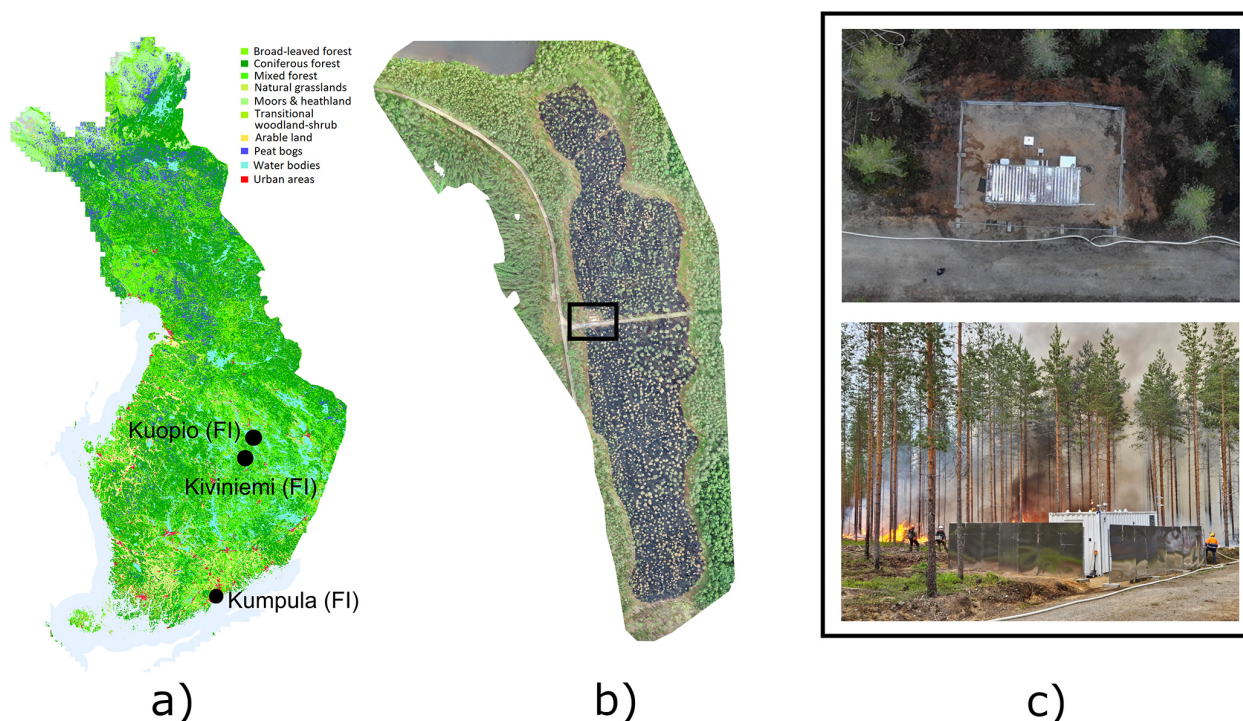


Figure 1. (a) Measurement site locations in Finland and land cover from the Copernicus Land Cover inventory CORINE (EU, 2025). (b) Aerial view of the burned area at Kiviniemi site. The black rectangle indicates the measurement station. (c) Top and side view of the measurement station in Kiviniemi.

fresh and aged smoke aerosols were observed in Finland during 2024 and 2020, and volcanic ash particles were observed in South Africa in 2016.

3.1 Case study 1: Extremely fresh smoke

A controlled prescribed burning was conducted on the 6 June 2024 in Kiviniemi ($62^{\circ}29'06.0''$ N, $27^{\circ}29'27.4''$ E) in North Savo region in Finland (Fig. 1). Approximately 8 hectares of managed boreal forest ecosystem was burned, with the objective of promoting biodiversity. The area is a 35–75 year old dry Scots pine (*Pinus sylvestris*) forest, with scattered birch (*Betula* sp.) within the stand. A commercial thinning was carried out in winter 2022, leaving logging residues unevenly distributed across the area. The understory is sparsely vegetated, with dwarf shrubs (*Vaccinium myrtillus*, *V. vitis-idaea*), mosses (*Pleurozium schreberi*), and *Cladonia* lichens dominating the forest floor. The prescribed burning was performed by the Finnish Forest Administration (Metsähallitus) and the fire was allowed to extinguish naturally – mostly within 24 h, although isolated smoldering persisted for several days after the burning event. The fire burned as a surface fire of low to moderate intensity, with spatially variable severity ranging from low to high. The prescribed burning was carried out in two events. In the first burning event, the fire started at about 11:00 UTC south from the measurement container along the horizontal road until the

whole southernmost section of the area was burned (Fig. 1b). The wind direction in this first burning was towards north, bringing the smoke plumes towards the measurement site. The second burning event started after 15:30 UTC. The fire was initiated at the easternmost area above the horizontal road and progressed northwards until the rest of the area was burned. The fire was made to progress in a direction against the wind to ensure a better control of the burning.

Figure 2 presents the atmospheric conditions over Kiviniemi, Finland on the 6 June 2024, recorded by the HALO Doppler lidar on site. Very high values in the ATB_{1565} profile are seen close to the surface starting at about 11:00 UTC, caused by high concentrations of smoke particles. In this first aerosol layer, which from now on is referred to as the smoke layer, the δ_{1565} is very low (Fig. 2b). The bottom and top boundaries of the smoke layer are also marked with red filled circles. As seen by the lidar, the smoke layer had its highest extent during the first burning event in which the direction of the wind was towards the lidar instrument and the fire was at its closest proximity to it. The maximum top height of the smoke layer reached at about 460 m and the average geometrical extent of it was 120 ± 90 m.

Figure 3 shows the relationship between ATB_{1565} and δ_{1565} for all height bins inside the smoke layer boundaries. A clear inverse relationship is observed, with increasing ATB_{1565} corresponding to decreasing δ_{1565} . This behavior is consistent with the presence of fresh smoke, which is char-

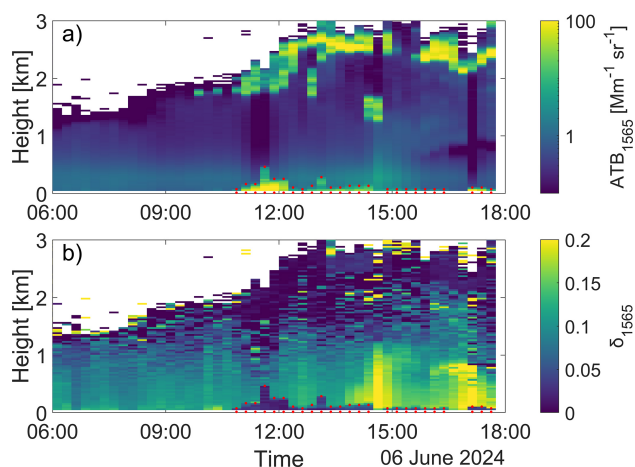


Figure 2. Time-height plot of (a) ATB_{1565} and (b) δ_{1565} measured in Kiviniemi 6 June 2024, with the HALO Doppler lidar. The red dots indicate the boundaries of the smoke layer. The temporal resolution corresponds to 15 min. The ATB_{1565} is shown on a logarithmic color scale. A co-polar SNR threshold of 0.0005 was applied to the observations.

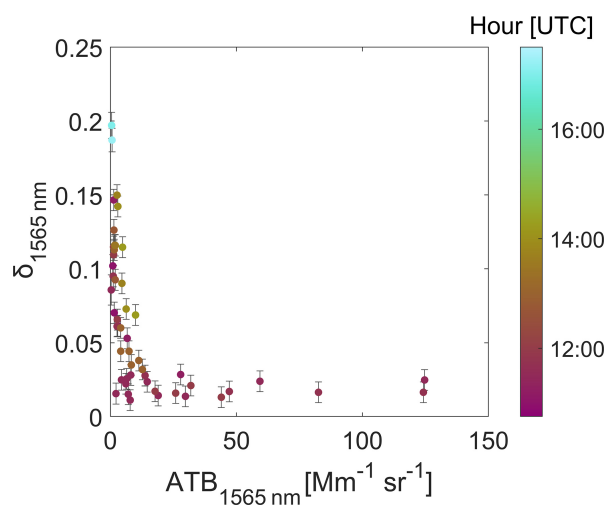


Figure 3. Scatterplot of ATB_{1565} versus δ_{1565} for all height bins within the boundaries of the smoke layer. The color indicates the time of the observations. The error bars show the absolute uncertainty of δ_{1565} .

acterized by enhanced backscatter and relatively low depolarization. For sufficiently high ATB_{1565} , δ_{1565} remains effectively constant. This behavior indicates that the signal is primarily governed by a single aerosol type, i.e. fresh smoke, with relatively uniform microphysical properties. Under these conditions, further increases in particle concentration enhance ATB_{1565} but do not significantly affect δ_{1565} , which is primarily governed by particle shape rather than concentration.

After 14:00 UTC, elevated δ_{1565} values are detected. At this time, the smoke plume was no longer aligned with the

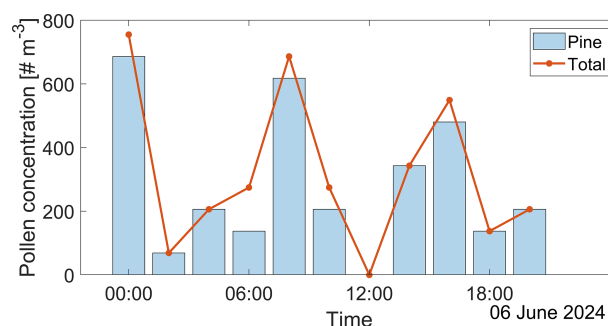


Figure 4. Evolution of the pine and total pollen concentration on the 6 June 2024 from the Burkard sampler at Kuopio site.

instrument's line of sight, as indicated by the concurrent decrease in ATB_{1565} . Nevertheless, the δ_{1565} time series reveals the presence of highly depolarizing aerosol particles. Although large fly-ash particles produced during biomass burning can contribute to increased depolarization (Martinent et al., 2025; Kalemekiewicz et al., 2018), the observed δ_{1565} enhancement extends up to approximately 1 km a.g.l., i.e. substantially higher than the fresh smoke layer observed between 11:00 and 12:00 UTC. This vertical extent suggests that the particles responsible for the elevated δ_{1565} were not associated with freshly emitted local smoke, but rather were well mixed within the boundary layer. To further examine this, Fig. 4 presents the daily evolution of pollen concentrations measured at a site in Kuopio (approximately 50 km from the lidar location). Pine pollen was the dominant type during the observation period. The elevated δ_{1565} values observed here are consistent with previous lidar observations of pine pollen (Filioglou et al., 2023). Therefore, the enhanced δ_{1565} signal is attributed to the regional presence of pine pollen particles.

A mean δ_{1565} of extremely fresh smoke aerosol particles of 0.017 ± 0.004 was estimated accounting for observations having an ATB_{1565} higher than $30 \text{ Mm}^{-1} \text{ sr}^{-1}$. Previous studies on fresh smoke particle detection report low δ due to the dominance of almost spherical morphology or weakly nonspherical aggregates shortly after emission. Typical tropospheric values of 0.01–0.03 at 355 nm, 0.02–0.06 at 532 nm and 0.01–0.02 at 1064 nm for fresh plumes have been previously reported (Pereira et al., 2014; Haarig et al., 2018; De Rosa et al., 2022). Falling well within the range of δ in shorter wavelengths, the δ_{1565} shows minimal spectral dependence.

3.1.1 Mass concentration estimation

Going beyond, the lidar-derived mass concentration was determined following the methodology outlined by Ansmann et al. (2010). Since there are no observational values on the LR and c_v at 1565 nm, parameters that are needed for the mass concentration calculation, we converted the β_{1565}

from the Doppler lidar into β_{532} using a $BAE_{532/1565}$ (See Sect. 2.2). Furthermore, in order to estimate the mass concentration at 532 nm, the $c_{v,532}$ and LR_{532} should be known. A $c_{v,532}$ of $0.16 \pm 0.01 \times 10^{-12}$ Mm for fresh smoke has been previously reported in Ansmann et al. (2026) and utilized in this study. A $BAE_{532/1565}$ and LR_{532} of 1.22 ± 0.09 and 28 ± 11 sr were estimated from Mie calculations at 532 nm (see Sect. 2.4). The Mie estimated LRs at 355, 532, 1064 and 1565 nm wavelengths for this specific aerosol mixture during the smoke event were 81 ± 13 sr, 28 ± 11 sr, 45 ± 2 sr and 51 ± 7 sr, respectively. Having the LR_{1565} , the determination of the conversion factor at 1565 nm for boreal smoke aerosol particles was then possible, yielding to a mean value of $c_{v,1565} = 0.32 \pm 0.18 \times 10^{-12}$ Mm.

Figure 5 demonstrates that the lidar-derived mass concentrations for the extremely fresh smoke event are in overall good agreement with the SMPS–OPC in situ observations. This consistency indicates that the lidar-based retrieval captures the main features of the near-source smoke plume despite the inherent differences in the sampling approaches. At the same time, both datasets reveal pronounced small-scale variability in smoke concentrations, which is expected given the close proximity of the fire source to the measurement site. This variability is particularly evident in the in situ observations. The vertical lines in Fig. 5 represent the temporal variability of the smoke concentration, expressed as the range between the 10th and 90th percentiles, highlighting strong concentration fluctuations within the plume. The lidar measurements additionally provide information on the temporal evolution and vertical structure of the smoke layer. The mass concentration is dominated by the amount of the larger aerosol particles and during the two flights moderate differences were captured. During these flights, the measurement altitude varied between approximately 50 and 100 m a.g.l., with maximum heights of 70 and 100 m during the first and second flights, respectively. During the 1st flight there is greater agreement between the lidar and in situ observation compared to the 2nd flight but at the same time the range of mass concentration is much larger. This discrepancy is expected, given the different sampling volumes and the inhomogeneity of the smoke plume and assumptions in the mass concentration estimation between these two measuring approaches. Lidar-derived mass concentration estimations are bound to 49 % uncertainty, assuming a 10 % uncertainty in the β_{1565} and the uncertainty of the rest parameters as noted above. The presence of a dense smoke layer around 12 UTC results in substantial attenuation of the lidar signal, making the forward Klett inversion inherently unstable and strongly dependent on the boundary conditions. Therefore, the retrieved β_{1565} carries considerable uncertainty and should be regarded as indicative of the plume structure only, rather than a quantitatively robust estimate. Supporting observations from surface and drone indicate that the aerosol composition during the fire event was dominated by OA, while BC contributed less than 1 % to the total mass. This

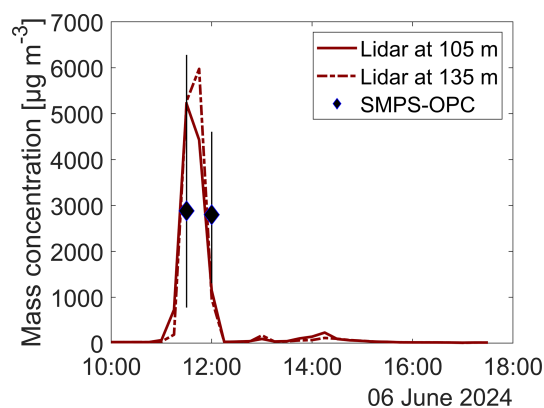


Figure 5. Mass concentration estimations from in situ SMPS–OPC observations and from lidar observations at 105 and 135 m. The SMPS–OPC observations are limited to the drone flight times between 11:30–11:40 and 12:00–12:10. The vertical lines indicate the 10th and 90th percentiles.

is consistent with the characteristics of surface fires, which burn at relatively low combustion temperatures (e.g. Cahill et al., 2008; Mukherjee et al., 2025; Vakkari et al., 2026). For crown-fire-dominated events, such as those commonly observed in North America (Rogers et al., 2015), a higher BC fraction is typical.

Related to the Mie-estimated LRs, the LR at 355 and 532 nm are consistent with previous measurements of extremely fresh smoke particles exhibiting a decreasing trend with values of 46 ± 4 sr at 355 nm and 34 ± 3 sr at 532 nm (De Rosa et al., 2025), although the Mie-estimated LR_{355} , in this study is larger than the one reported at De Rosa et al. (2025). More often, there is a neutral or slight increasing spectral dependence (Müller et al., 2005; Nicolae et al., 2013; De Rosa et al., 2022; Janicka et al., 2023) reported for fresh smoke. Although there are no literature values for the LR_{1565} , increasing LR_{1064} has been previously reported for long-range transported free-tropospheric smoke layers (Haarig et al., 2018; Su et al., 2025). This is not the case in Sayer et al. (2014) where similar LR values at 355 and 532 nm followed by a decrease towards 1064 nm were reported at boreal locations using observations from a plethora of AERONET sites. To this direction, using light-scattering simulations, Liu and Mishchenko (2020) investigated the lidar ratios at 532 and 1064 nm for a range of refractive indexes, particle sizes and shapes and found a decreasing trend with increasing wavelength for aerosol particles with an effective radius smaller than about $0.3 \mu\text{m}$ as well. In this study, the Mie estimations point to increasing LRs between 1064 and 1565 nm wavelengths but the values are smaller compared to the ones reported in Haarig et al. (2018) and Su et al. (2025), which is attributed to the aging of the smoke particles and the likely higher share of absorbing aerosols like BC in the aerosol mixture in their studies.

The lower LRs estimated from the Mie calculations at the visible and infrared wavelengths could be explained by the contribution of BrC, which absorbs more effectively at shorter wavelengths. Using a laboratory open burning setup Mukherjee et al. (2025) found that optical properties of freshly emitted organic particles are substantially diverse and dependent on the fire type (smoldering, flaming) and burning material. Among others, they used Finnish boreal peat and forest surface samples. Photochemical aging transformed the weakly absorbing particles into less absorbing while the opposite is anticipated in dark-aging conditions. In contrast, BC is more likely to become more absorbing due to the lensing effect Zhang et al. (2018). The wavelength dependence of LR in aged smoke particles (> a few hours) is also related to the composition of the smoke plume as it contains larger aggregates that become optically apparent at longer wavelengths. It is important to emphasize that the case presented here concerns smoke aerosol particles sampled within a few minutes after emission, whereas most previously reported observations of fresh smoke aerosol layers in the literature present substantially longer timescales (from hours to days). Smoke particles undergo rapid aging within hours after emission altering their size and scattering properties (Kleinman et al., 2020; Saide et al., 2022; Vakkari et al., 2018).

3.2 Case study 2: Long-range transported free tropospheric smoke

On the 22 September 2020, free-tropospheric aerosol layers were observed over Kumpula site ($60^{\circ}12'13.5''\text{N}$, $24^{\circ}57'39.1''\text{E}$) in Finland (Fig. 6). The aerosol layers which extended from 1.3 to 5.1 km in the vertical had a variable geometrical depth with a mean value of 800 m (range: 120 m–2.2 km). An example case of the hourly-averaged Doppler lidar profiles between 17:00 and 18:00 UTC is shown in Fig. 7. Elevated β_{1565} values can be seen at around 1.6 km and between 2.7 and 4 km with a mean δ_{1565} of 0.034 ± 0.01 and 0.038 ± 0.017 , respectively. For this case, the 16 d FLEXPART backward simulation showed air mass transport from California, which overlapped with very intensive fire locations according to MODIS observations (Fig. 8). In fact, long-range transport from the same wildfire episode in California (USA) was observed a few days earlier in Germany (Baars et al., 2021).

To further investigate the variation of δ_{1565} in these long-range transported smoke layers, hourly ATB_{1565} and δ_{1565} profiles were calculated to enhance the signal-to-noise ratio (SNR). Figure 9 presents the relationship between the hourly-averaged profiles of ATB_{1565} and δ_{1565} for all height bins inside the smoke layer boundaries. The smoke layer boundaries for the hourly profiles are indicated in the hourly time-height plot that can be found in the appendix (Fig. A1). Similar to the fresh smoke, lower δ_{1565} values coincide with higher ATB_{1565} . Since this is long-range transported smoke located in the free troposphere, the ATB_{1565} is lower and

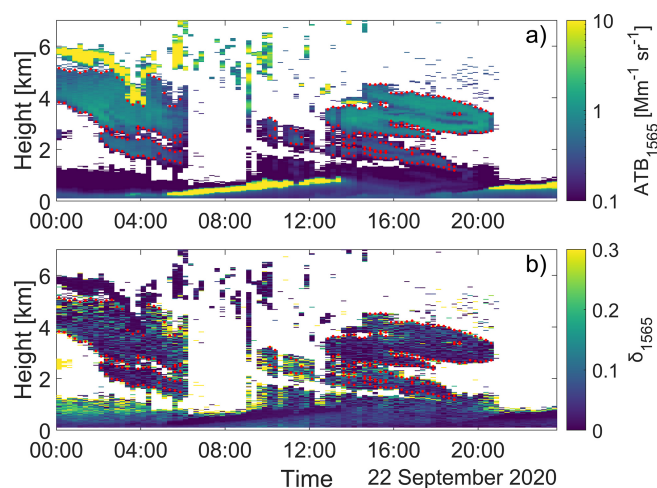


Figure 6. Time-height plot of (a) ATB_{1565} and (b) δ_{1565} measured in Kumpula 22 September 2020, with the HALO Doppler lidar. The red dots indicate the boundaries of the smoke layer. The temporal resolution corresponds to 15 min. The ATB_{1565} is shown on a logarithmic color scale. A co-polar SNR threshold of 0.0006 was applied to the observations.

the scatter of δ_{1565} is larger than in the extremely fresh smoke case. In order to retrieve the mean δ_{1565} , we have accounted both for the relative uncertainty of δ_{1565} and the intensity of ATB_{1565} in a way that observations presenting high ATB_{1565} values and low relative uncertainty in δ_{1565} are weighted more in the average. A weighted mean δ_{1565} of smoke aerosol particles of 0.04 ± 0.02 was estimated for the long-range transported smoke particles following Eq. (1). We should note here that the same conclusions are valid when estimating the mean δ_{1565} using the 15 min resolution information but the uncertainty is higher due to the poorer SNR (not shown).

Free-tropospheric smoke particles have been reported to produce depolarization ratios below 5 % across lidar wavelengths up to 1064 nm (Pereira et al., 2014; Haarig et al., 2018). The present findings are consistent with these earlier observations. In comparison to the fresh smoke, the free tropospheric smoke δ_{1565} is larger, indicating that the shape of the particles is more non-spherical. Various factors can contribute to the observed difference in δ_{1565} , including higher fraction of fractal-shaped BC in smoke emitted in fires with high FRP or the presence of soil dust in the smoke plumes. Recent findings also link organic aerosol shape changes to chemical aging (Qin et al., 2024).

$$\frac{\sum w \delta_{1565}}{\sum w} \quad \text{where} \quad w = \left(1 - \frac{\frac{\sigma_{\delta_{1565}}}{\delta_{1565}}}{\max\left(\frac{\sigma_{\delta_{1565}}}{\delta_{1565}}\right)} \right) \frac{\text{ATB}_{1565}}{\max(\text{ATB}_{1565})} \quad (1)$$

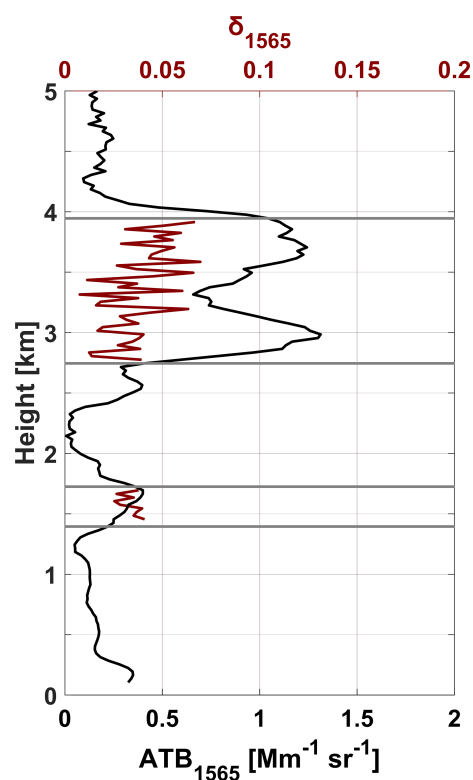


Figure 7. Mean ATB_{1565} (black) and δ_{1565} (dark red, for the smoke layers only) vertical profiles accounting for a temporal resolution of 1 h starting at 17:00 UTC. The smoke layer base and top heights are indicated by gray horizontal lines.

3.3 Case study 3: Volcanic ash in free troposphere

A free tropospheric aerosol layer with very high δ_{1565} reached the measurement site at Welgegund, South Africa ($26^{\circ}34'11.3''$ S, $26^{\circ}56'21.5''$ E) in the night from 16 to 17 December 2016. Figure 10 shows the temporal evolution of the layer's ATB_{1565} (upper panel) and δ_{1565} (lower panel) over the measurement site as monitored by the HALO Doppler lidar. The base and top heights of the layer are marked with circles. The increase of backscatter coefficient between 2.5 and 3.5 km clearly demonstrates the existence of a pronounced aerosol layer and the large values of δ_{1565} are a clear indication of non-spherical particles; a typical feature of volcanic ash (Pisani et al., 2012). Scattered clouds before and after the plume did not prevent the observation of the plume. The plume's altitude remained rather constant during the 12 h observation period as well as the layer's geometrical thickness of 850 m.

FLEXPART air mass history for this layer (Fig. 11) shows substantial contribution from South America and the direction of Copahue Volcano, which is located on the border between Chile and Argentina in the Andes Mountains ($37^{\circ}45'$ S, $71^{\circ}10.2'$ W, 2997 m a.s.l.). Historically, the volcano has exhibited intermittent mild-to-moderate explosive

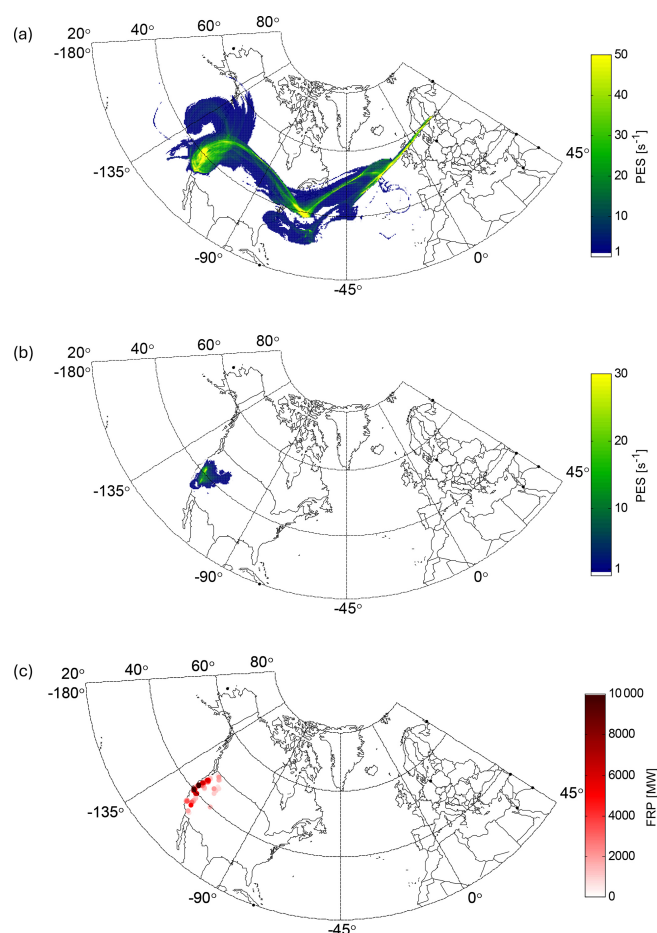


Figure 8. Air mass history for the Kumpula case on 22 September 2020 in Finland. (a) Sum of PES at all heights accounting for 16 d back in time. (b) Sum of PES for < 1 km a.g.l. for 16 d back in time. (c) MODIS observations of FRP > 500 MW for the time period 8 to 12 September 2020.

activity, with a notable increase in eruptions since the beginning of 2012. During early December 2016, diffuse gas, water vapor, and ash plumes from Copahue rose to altitudes of 3–3.3 km a.s.l. and were transported eastbound (Paez et al., 2021), which overlaps with the air mass history analysis in Fig. 11. Therefore, we attribute the pronounced high δ_{1565} layer as ash originating from the Copahue Volcano.

Figure 12 presents the hourly extracted lidar information of ATB_{1565} and δ_{1565} at the volcanic plume layer only (Fig. A1). The color indicates the relative uncertainty of δ_{1565} . In this highly depolarizing aerosol layer, we can define the δ_{1565} with a better than 10 % accuracy estimating a depolarization ratio for the volcanic ash aerosols to be 0.46 ± 0.05 at 1565 nm. A δ_{1565} of 0.45 ± 0.01 is estimated using the weighted mean method described in the previous case study. No significant changes were found between the 15 min and 1 h temporal resolutions apart from the higher uncertainty accompanying the finer temporal resolution observations.

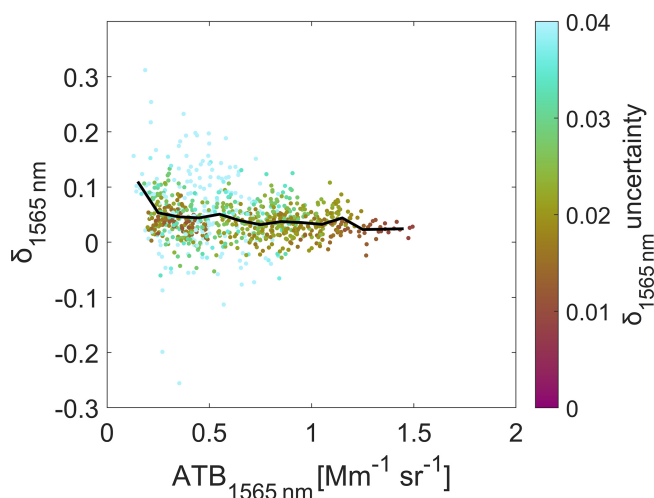


Figure 9. Scatterplot of $ATB_{1565\text{ nm}}$ versus $\delta_{1565\text{ nm}}$ at the long-range transported smoke plume layer extracted from hourly lidar data. The black line shows the mean $\delta_{1565\text{ nm}}$ binned every $0.1\text{ Mm}^{-1}\text{ sr}^{-1}$. The color indicates the absolute uncertainty of $\delta_{1565\text{ nm}}$.

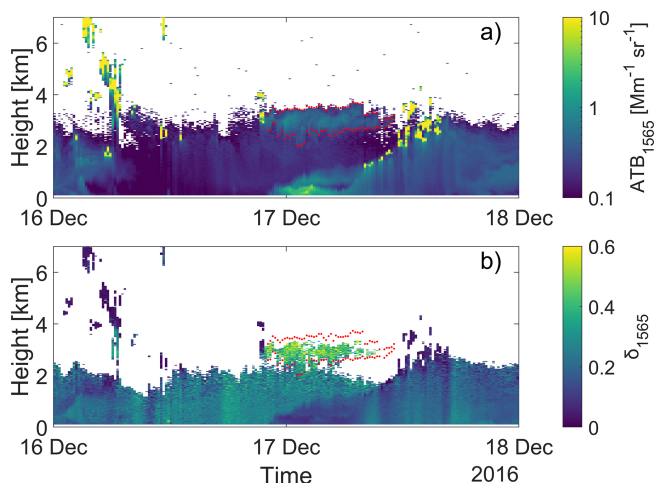


Figure 10. Time-height plot of (a) ATB_{1565} and (b) δ_{1565} measured between 16 and 17 December 2016 in Welgegund, South Africa, with the HALO Doppler lidar. The red dots indicate the boundaries of the smoke layer. The temporal resolution corresponds to 15 min. The ATB_{1565} is shown on a logarithmic color scale. A co-polar SNR threshold of 0.0005 was applied to the observations.

3.4 Depolarization ratio at 1565 nm

Here we summarize the depolarization ratios of the fresh smoke, aged smoke and volcanic ash aerosol particles from this study. We also gather δ_{1565} information for all studied aerosol particles up-to-date (Table 2). This collection of case studies at 1565 nm for various aerosol types demonstrates that lidar-based classification algorithms could benefit from the extended information at δ_{1565} , particularly for identifying large non-spherical particles. Besides non-sphericity, particle

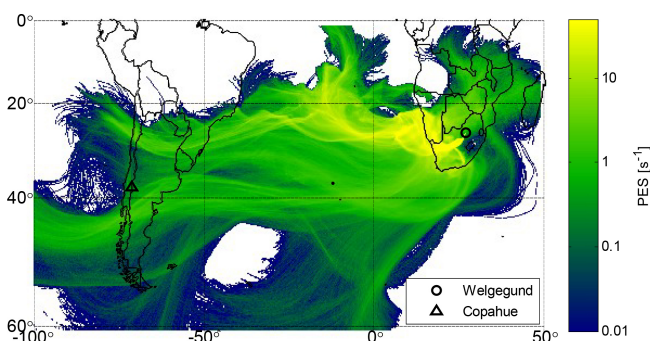


Figure 11. FLEXPART PES summed up over all heights accounting for 14 d back in time for the Welgegund case in South Africa on 17 December 2016.

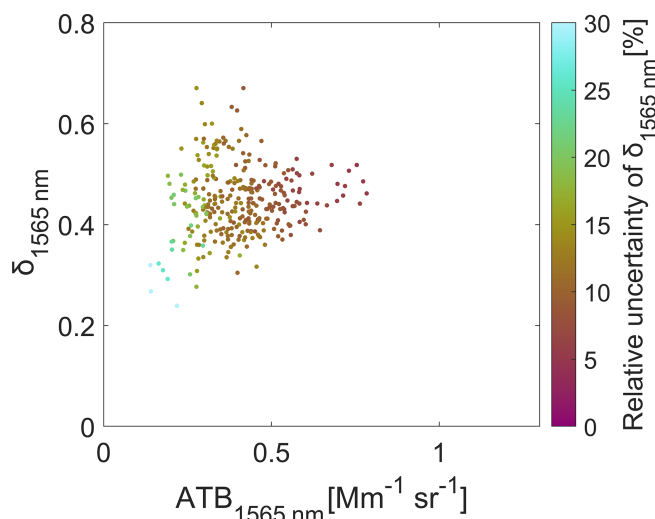


Figure 12. Scatterplot of $ATB_{1565\text{ nm}}$ versus $\delta_{1565\text{ nm}}$ at the volcanic plume layer extracted from hourly lidar data. The color indicates the relative uncertainty of $\delta_{1565\text{ nm}}$.

size should also be emphasized, because longer wavelengths are more sensitive to larger particles. The aerosol types with enhanced depolarization ratios at 1565 nm listed in Table 2, including dust, pollen, and volcanic ash, are generally large particles. In contrast, stratospheric smoke observations have shown that small particles may retain low depolarization ratios at longer wavelengths even if they are non-spherical (Haarig et al., 2018). Therefore, δ_{1565} provides complementary information related to both particle shape and size, highlighting its potential for aerosol classification and air-quality monitoring.

For the three aforementioned aerosol types δ exhibits a wavelength dependence. Specifically, for fresh smoke, depolarization ratios at shorter wavelengths (355 and 532 nm) are typically low due to the almost spherical nature of freshly emitted sub-micron particles; previous multi-wavelength lidar studies report tropospheric smoke depolarization ratios < 0.03 at 355 and 532 nm and 0.01 at 1064 nm; a consis-

Table 2. Depolarization ratio of aerosol types at 1565 nm.

Aerosol type	δ_{1565}	Reference
Polluted marine	0.009 ± 0.003	Vakkari et al. (2021)
Fresh smoke	0.017 ± 0.004	This study
Aged smoke	0.04 ± 0.02	This study
Mixture of spruce and birch pollen	0.269 ± 0.005	Vakkari et al. (2021)
Pine pollen	0.26 ± 0.03	Filioglou et al. (2023)
Birch pollen	0.33 ± 0.04	Filioglou et al. (2023)
Pollen mixture	0.23–0.30	Le et al. (2024)
Desert dust	0.30 ± 0.02	Vakkari et al. (2021)
Long-range transported desert dust	0.24 ± 0.008	Le et al. (2024)
Volcanic ash	0.45 ± 0.05	This study

tent behavior at 1565 nm was found in this study too, reflecting limited non-sphericity. In aged smoke, particle reshaping and mixing together with the smoke's physical and chemical properties and any co-lifted dust particles can enhance non-sphericity at shorter wavelengths. At longer wavelengths, however, depolarization tends to remain low because small soot aggregates scatter less efficiently, producing a characteristic spectral decline in depolarization with increasing wavelength. In contrast, volcanic ash consists of larger, non-spherical particles that produce high depolarization ratios at both UV and visible wavelengths. Previous studies at 355 and 532 nm report δ of about 0.35–0.38 for ash plumes with little spectral variation among these shorter wavelengths (Groß et al., 2012). Our findings suggest that this non-sphericity persists at 1565 nm and is further enhanced, presenting an increase of δ with increasing wavelength.

4 Conclusions

For the first time, we report particle linear depolarization ratios for extremely fresh and aged forest fire smoke as well as volcanic ash at 1565 nm. We found that (a) non-spherical large particles such as volcanic ash induce elevated particle linear depolarization ratios at 1565 nm, consistent to the previous observations at shorter wavelengths and (b) smoke aerosols, dominated by almost spherical particles, yield lower depolarization ratio, but aged smoke show some variability probably due to differences in the BC content, dust uplifting and/or processing during the plume transport. We further estimated the mass concentration of extremely fresh smoke using a combination of Mie-derived optical properties and lidar observations with reasonable agreement to in situ observations. The Mie-estimated lidar ratios increased with increasing wavelength except at 355 nm. Whether this behavior is characteristic of extremely fresh smoke particles in boreal forest environments and for this type of surface fire remains uncertain; however, previous observational and light-scattering simulations have demonstrated the high sensitivity

of lidar intensive properties such as the lidar ratio to smoke particles.

For the extremely fresh smoke event within the planetary boundary layer, we found very low particle linear depolarization ratio at 1565 nm of 0.017 ± 0.004 , which is in good agreement with previous observations at 1064 nm showing minimal wavelength dependence. For long-range transported smoke aerosols in the free troposphere, we observed marginally higher depolarization ratio of 0.04 ± 0.02 . The volcanic ash depolarization ratio induced high values of 0.45 ± 0.01 and combined with earlier lidar studies at shorter wavelengths it shows an increase with increasing wavelength. Taken together, our findings show that Halo Doppler lidars provide a valuable additional wavelength information at 1565 nm for investigating the spectral dependence of particle linear depolarization ratio, even up to the lowest 4–5 km of the atmosphere. Therefore, they build towards a more accurate aerosol typing and support the development of new aerosol climatologies, thereby enhancing network-wide utilization of aerosol measurements at this wavelength.

Aerosol observations at 1565 nm benefit from the near absence of molecular scattering and absorption, which greatly simplifies signal pre-processing. At this wavelength, Rayleigh scattering is negligible, so the lidar return is dominated by aerosol particles rather than atmospheric molecules. This provides a clear advantage over the shorter wavelengths and for real-time application of the observations. However, the trade-off is that the aerosol backscatter efficiency is much lower compared to visible wavelengths, making it harder to detect fine particles and aerosol layers with low concentrations. Furthermore, mixtures of different aerosol types could also become challenging due to the weaker scattering contrast.

Future work should further investigate aerosol optical and microphysical properties at 1565 nm, as current understanding at this wavelength emerges from limited case studies. Expanded observational studies with various aerosol mixtures would help clarify how different aerosol types behave

in the near-infrared and how these signatures vary with atmospheric conditions. Such efforts are essential for fully assessing the potential of 1565 nm measurements to complement the traditional lidar wavelengths and improve aerosol classification efforts.

Appendix A

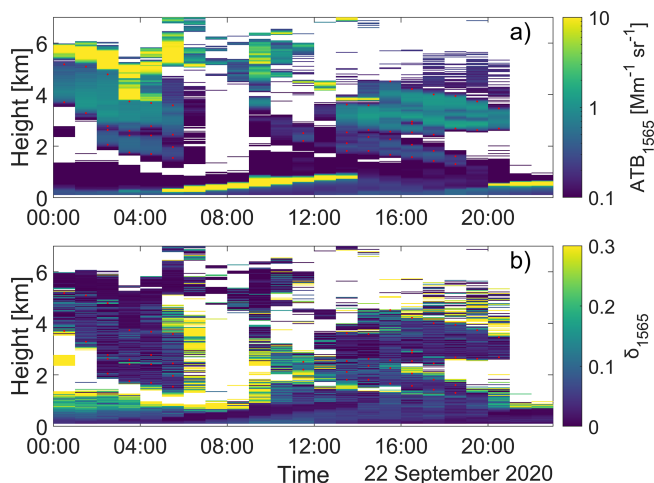


Figure A1. Similar plot to Fig. 6 for a temporal averaging of 1 h.

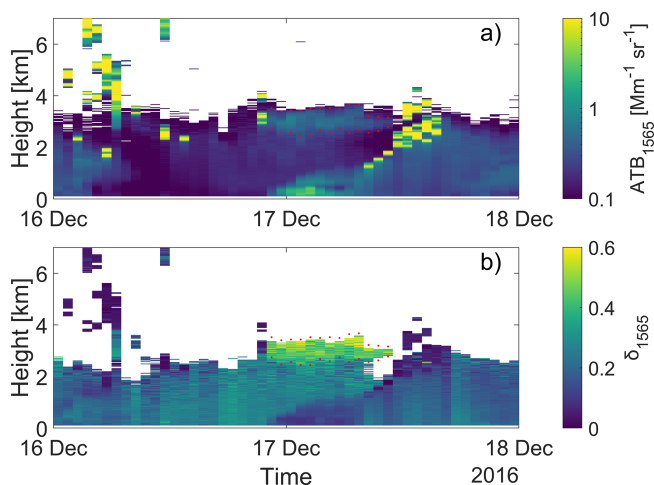


Figure A2. Similar plot to Fig. 10 for a temporal averaging of 1 h.

Data availability. HALO Doppler lidar observations are available through Cloudnet portal (<https://cloudnet.fmi.fi/file/7fc69d32-56df-48d8-ba9d-d302051bc414>, Komppula and O'Connor, 2025). Flexpart version 11.0 is available at <https://gitlab.phaidra.org/flexpart/> (last access: 14 August 2026). Fire radiative power (FRP) data are publicly available through FIRMS interface at <https://firms.modaps.eosdis.nasa.gov/map/> (last

access: 14 August 2026). In situ aerosol and pollen observations are available upon request.

Author contributions. MF conceptualized the original paper, performed the main analysis considering all data sources and wrote the manuscript. VV conceptualized the original paper and provided the Halo Doppler lidar observations, FLEXPART air mass trajectories and MODIS fire radiative power data. KL combined the in situ observations from SMPS and OPC instruments, performed and provided the Mie calculations. EA analyzed and provided the MAAP observations. EA and VL analyzed and provided the OPC observations. DB prepared the drone load and VL and KH performed the flights onsite. SK provided the SMPS observations. LH provided the AMS observations. AS and SPä provided the pollen data. All authors were involved in the editing and discussion of the manuscript.

Competing interests. At least one of the (co-)authors is a member of the editorial board of *Atmospheric Chemistry and Physics*. The peer-review process was guided by an independent editor, and the authors also have no other competing interests to declare.

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Acknowledgements. This study was supported by the Research Council of Finland (grant nos. 337552, 343359, 369600, 369601). The prescribed burning is part of the project "Climate and air quality impacts of boreal forest fires", (2023–2027) funded by the Jane and Atos Erkko Foundation. The fire area is part of the Life2Taiga project (2022–2028) funded by the European Union's LIFE Nature and Biodiversity programme, which aims to restore forests in Finland and Sweden through conservation burning. We acknowledge the use of data from NASA's Fire Information for Resource Management System (FIRMS) part of NASA's Land, Atmosphere Near real-time Capability for Earth observations (LANCE) and NASA's Earth Science Data and Information System (ESDIS).

Financial support. This research has been supported by the Research Council of Finland (grant nos. 337552, 343359, 369600, and 369601).

Review statement. This paper was edited by Matthias Tesche and reviewed by three anonymous referees.

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