

Does the Asian Summer Monsoon Play a Role in the Stratospheric Aerosol Budget of the Arctic?

Sandra Graßl^{1,2}, Christoph Ritter¹, Ines Tritscher³, and Bärbel Vogel³

¹Alfred Wegener Institute, Helmholtz Centre for Polar and Marine Research, 14473 Potsdam, Germany

²Institute of Physics and Astronomy, University of Potsdam, Karl-Liebknecht 24/25, 14476 Potsdam, Germany

³Forschungszentrum Jülich, Institute of Energy and Climate Research – stratosphere (IEK-7), Jülich, Germany

Correspondence: Sandra Graßl (Sandra.Grassl@awi.de)

Abstract. The Asian summer monsoon has a strong convectonal component, with which aerosols are able to be lifted up into the lower stratosphere. Due to usually long lifetimes and long-range transport aerosols remain there much longer than in the troposphere and are also able to be advected around the globe. Our aim of this study is a synergy between modelled tropical aerosol tracers by Chemical Lagrangian Model of the stratosphere (CLaMS) and KARL (Koldewey Aerosol Raman Lidar) at AWIPEV, Ny-Ålesund in the Arctic, by comparing back- and forward trajectories with exemplary days of Lidar measurements as well as analyse the stratospheric aerosol background. We use global 3-dimensional Lagrangian transport simulations including surface origin tracers as well as back-trajectories to identify source regions of the aerosol particles measured over Ny-Ålesund. We analysed Lidar data for the year 2021 and found the stratosphere generally clear, without obvious aerosol layers from volcanic eruptions or biomass burnings. Still an obvious annual cycle of the backscatter coefficient with higher values in late summer to autumn and lower values in late winter have been found. Results from CLaMS model simulations indicate that from late summer to early autumn filaments with high fractions of air which originate in South Asia – one of the most polluted regions in the world – reach the Arctic in altitudes between 360 K and 380 K potential temperature. We found a coinciding measurement between the overpass of such a filament and Lidar observations, we estimated that backscatter and depolarisation increased by roughly 15% during this event compared to the background aerosol concentration. Hence we demonstrate that the Asian summer monsoon is a weak but measurable source for Arctic stratospheric aerosol in late summer to early autumn.

1 Introduction

Already in 1960 Junge et al. (1961a) characterised some basic properties of stratospheric aerosols like size distribution, physical structure and chemical composition up to 30 km altitude. Stratospheric aerosol can either be long-range transported, injected from the troposphere or even directly created in the stratosphere and has a high sulfuric content (Junge et al., 1961a; Kremser et al., 2016), they have a big impact on the local ozone layer but also on the radiative budget with a cooling factor due to additional scattering of direct solar radiation. Already Junge et al. (1961a) pointed out different mechanisms acting on stratospheric aerosols, like horizontal mixing due to large-scale exchange by intrusions through the tropopause, meridional, hemispheric circulations or gravitational settling of particles determined by Stokes' law. Junge et al. (1961a) as well as Murphy

25 et al. (2021) found in stratospheric in-situ measurements a remarkable constant layer of particles in the accumulation mode
(from 0.1 μm to 1 μm). Particles larger than that are created by oxidation of H_2S and SO_2 in the layer they are found. Strong
zonal winds in the stratosphere lead to a fast homogenisation of aerosols, while vertical and meridional transport is controlled
by the Brewer-Dobson-Circulation (BDC) (Holton et al., 1995; Butchart, 2014). The BDC is characterised by a meridional
transport from rising air at the tropics towards high latitudes and a quasi-horizontal mixing over the extra-tropics (McIntyre
30 and Palmer, 1983). This circulation and mixing controls not only long-range transport (Shepherd, 2002) but also ageing of
aerosols (Vaughan and Hall, 2002).

As Junge et al. (1961a) found out, life-times of aerosols in the lower stratosphere are in the order of magnitude of several
hundred days and can easily be advected to higher latitudes and distributed over the entire globe. Due to the absence of water
vapour in the stratosphere aerosol removal by precipitation does not take place. Therefore they are usually removed by sedi-
35 mentation.

During the first four years of CALIPSO measurements (Cloud-Aerosol Lidar and Infrared Pathfinder Satellite Observations)
an Asian Tropopause Aerosol Layer (ATAL) was found during the Asian summer monsoon season in altitudes of 13 to 18 km
equivalent to 360 K to 420 K potential temperature, while the Junge layer is usually at around 20 km (Junge et al., 1961b). With
usually occurring depolarisation values of up to 5% and backscattering ratios of around 1.10 to 1.15, this layer is driven by an
40 enhancement of background aerosol (Vernier et al., 2011). While overall the amount of non-sulfate aerosol or anthropogenic
sulfur dioxide sources are difficult to assess (Kremser et al., 2016), the ATAL might be responsible for about 15% of the aerosol
content in the Northern Hemisphere. This is comparable with all volcanic eruptions from 2000 to 2015 (Yu et al., 2017) and
within the order of magnitude of the estimated amount of ejected SO_2 during the Pinatubo eruption in 1991 (Andersson et al.,
2015; Kremser et al., 2016; Quaglia et al., 2022). Chemical in-situ measurements revealed that the ATAL mainly consists of
45 secondary substances containing nitrate, ammonium, sulfate and organics and is therefore more diverse in its chemical compo-
sition than the Junge layer (Appel et al., 2022).

The Asian summer monsoon has very strong deep convection, where updrafts of 4 m/s are common (Wang, 2004), into
altitudes of 400 K (Randel et al., 2010; Bian et al., 2020) which allow both gas-phase aerosol precursors and aerosol particles
50 from surface sources to reach the stratosphere (Vogel et al., 2019; Brunamonti et al., 2018; Hanumanthu et al., 2020). Several
studies confirm that the horizontal transport out of the Asian monsoon anticyclone has an impact on the lower extra-tropical
stratosphere of the Northern Hemisphere (Vogel et al., 2016; Yu et al., 2017; Yan et al., 2024, e.g.). Beside of volcanic ac-
tivities, anthropogenic emissions make a large contribution to the total concentration of aerosol with sulfuric components. As
Notholt et al. (2005); Klimont et al. (2013) show, the sulfur emission of Europe, North America and the former Soviet Union
55 are constantly decreasing, while the emission drastically increases in Southeast Asia, where also the very effective uplift of
pollution, gas-phase aerosol precursors and aerosol particles in the stratosphere by the Asian summer monsoon takes place.
Some observational evidence of an increasing of stratospheric aerosols globally was found by Vernier et al. (2011) who used
observations by Cloud-Aerosol Lidar and Infrared Pathfinder Satellite Observations (CALIPSO) that revealed the ATAL near
16 km in June to September is associated with the Asian monsoon anticyclone.

60 Using the CLaMS model (Chemical Lagrangian Model of the stratosphere) and aircraft measurements over Europe and the northern Atlantic, horizontal pathways out of the Asian monsoon anticyclone were found exporting young polluted air masses eastward into the extra-tropical northern lower stratosphere within several weeks (e.g. Vogel et al., 2016; Wetzel et al., 2021; Lauther et al., 2022). In particular source regions of anthropogenic short-lived species such as CH_2Cl_2 emission are mainly located in Asia, therefore CH_2Cl_2 is a very good marker for transport out of the Asian monsoon anticyclone into the northern
65 extratropical stratosphere (Lauther et al., 2022). The good linkage between CLaMS surface origin tracers and measured pollution tracers such as CH_2Cl_2 and PAN (Wetzel et al., 2021; Lauther et al., 2022) show that CLaMS is very well suited for analysing aerosol transport out of ATAL.

Furthermore, aerosol originating from extended boreal biomass burnings especially in Russia and North America have been
70 seen in the Arctic summer stratosphere at different sights and years in the Arctic (Zielinski et al., 2020; Ohneiser et al., 2021; Cheremisin et al., 2022). According to satellite-based studies of global extinction results wild fires can only be found in the lowermost part of the stratosphere with a relatively short lifetime (Thomason and Knepp, 2023). Therefore, the annual cycle with its natural variability and the origin of the Arctic stratospheric aerosol layer is important.

75 Currently stratospheric aerosol represents an interesting research topic as already several studies analyse a potential impact of stratospheric aerosol injections on geoengineering (e.g. Irvine and Keith, 2020; Cheng et al., 2022), to counteract the global rise of temperature. As Sun et al. (2020) point out, the impact of an injection of sulfuric aerosols into the stratosphere will change globally the incoming radiation but also the precipitation. Additionally, the effect will be different depending on the location of injection as well as the amount.

80 However, clearly all potential impacts on the climate system need to be understood prior to such an execution of costly and risky environmental endeavour. To minimise impacts of precipitation suppression some authors favour aerosol injection into the Arctic summertime stratosphere (Lee et al., 2021). It is therefore important to understand the existing contribution of the un-modified stratospheric aerosol content in the Arctic, which may also originate from the Asian summer monsoon. As a first step we look into the seasonal variability of the Arctic stratospheric background aerosol impacted by the Asian summer mon-
85 soon.

Therefore we use model simulations to identify the contribution of the Asian summer monsoon to air masses detected by Lidar measurements in the Arctic. For this study we focus on the year 2021, since it was a year with a statistically average amount of biomass burnings on the northern hemisphere and we have a qualitatively good Lidar data set for every month of
90 the year. In Section 2.1 we describe the instrument sight, followed by a short overview of the Lidar KARL (see Section 3.3), radiosondes in Section 2.3, the model CLaMS (see Section 2.4) and the satellite MODIS in Section 2.5. We present an overview of basic stratospheric aerosol properties above our Arctic site in Section 2.2 which shows a weak but clear annual cycle. Further we show the output of a Lagrangian chemical transport model in Section 3.5 which shows the occurrences of air which

originated in the Asian Summer monsoon region over the Arctic. The discussion of the results will be presented in Section 4 and the conclusion will follow in Section 5.

2 Measurement Site, Instrument and Model

2.1 Arctic Measurement Site: Ny-Ålesund

The observation site in Ny-Ålesund is located in the European Arctic (78.923°N , 11.928°E) on the northwest coast of the archipelago of Svalbard along the shore of Kongsfjord, which is orientated in a SE to NW direction on the west coast of Svalbard (Figure 1).

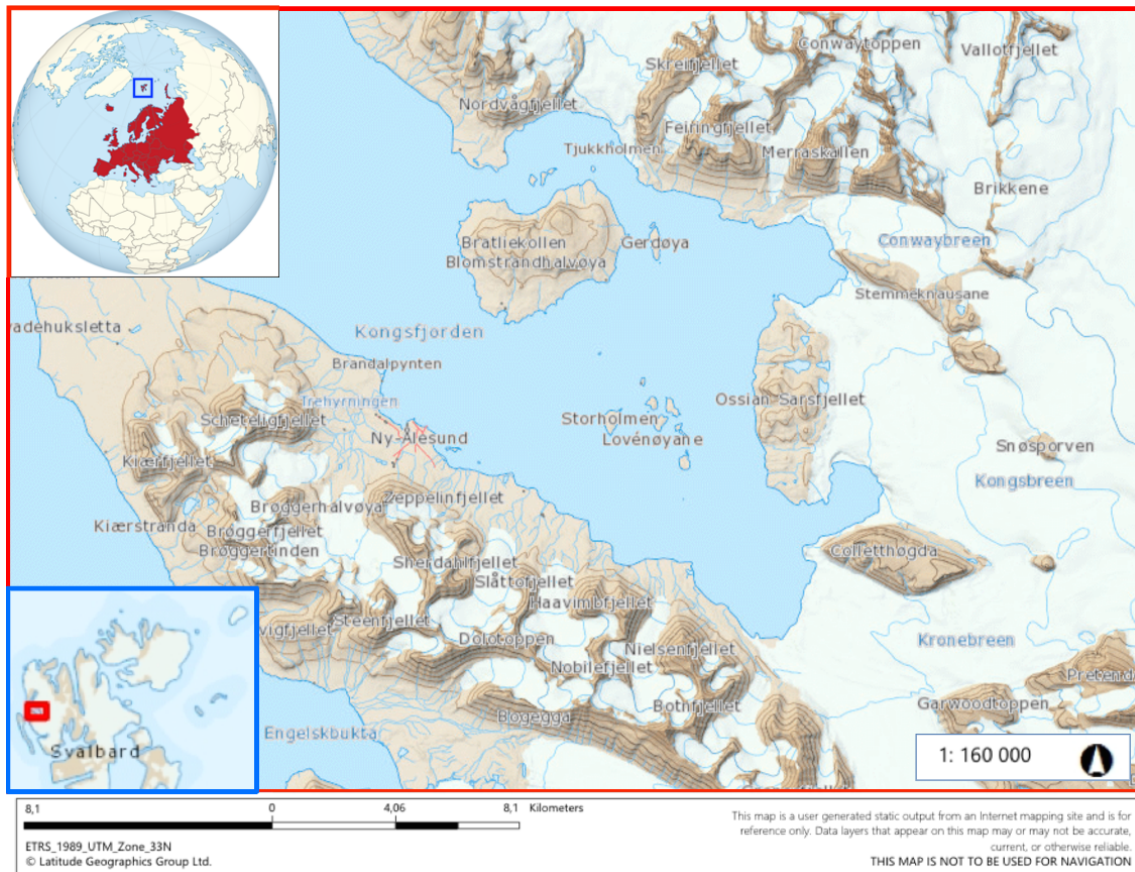


Figure 1. Map of the vicinity of Ny-Ålesund, Kongsfjord and its location in Europe. Sources: <https://geokart.npolar.no/Html5Viewer/index.html?viewer=Svalbardkartet> (accessed on April 23, 2024), courtesy of Norsk Polarinstitut and [https://de.wikipedia.org/wiki/Datei:Europe_on_the_globe_\(red\).svg](https://de.wikipedia.org/wiki/Datei:Europe_on_the_globe_(red).svg) (accessed on April 23, 2024)

2.2 KARL – Koldewey Aerosol Raman Lidar

2.2.1 Description of KARL

The Lidar measurements were performed at the German-French AWIPEV research base in Ny-Ålesund, Svalbard by the "Koldewey Aerosol Raman Lidar" (KARL). This system consists of a Spectra 290/50 Nd:YAG laser, which emits a laser beam at 355 nm, 532 nm and 1064 nm, with 50 Hz and 200 mJ per pulse and color vertically into the atmosphere. The backscattered photons are collected by a 70 cm telescope with a field of view of about 2 mrad in the elastic colours as well as in the Raman-scattered signals at N₂ at 387 nm and 607 nm. Data recording is done via Hamamatsu photomultipliers and Licel transient recorders with 7.5 m resolution. Full overlap is reached at about 700 m altitude. A more detailed description of the instrument is given by Hoffmann (2011). The aerosol backscatter coefficient $\beta^{aer}(\lambda)$ [(m · sr)⁻¹] and the aerosol extinction coefficient $\alpha^{aer}(\lambda)$ [m⁻¹] were calculated following the principle of Ansmann et al. (1992). For better readability and simplicity we use $\beta(\lambda) := \beta^{aer}(\lambda)$ and $\alpha(\lambda) := \alpha^{aer}(\lambda)$ in the following, if not stated otherwise. The Lidar ratios, $LR = \frac{\alpha^{aer}(\lambda)}{\beta^{aer}(\lambda)}$ [sr], for the other wavelengths were chosen $LR_{355} = 50$ sr for the troposphere and $LR_{355} = 70$ sr in the stratosphere, for $\lambda = 532$ nm $LR_{532} = 36$ sr and $LR_{532} = 42$ sr in the troposphere and stratosphere, respectively, if not stated otherwise. High values of LR indicate highly absorbing particles and is therefore mostly related to the refractive index but also to shape and size. The fact that we found $LR_{532} < LR_{355}$ in the stratosphere was derived by our Lidar data via two conditions: same backscatter at different wavelengths for (tropospheric) cirrus clouds and the same gradients for the aerosol backscatter in clear layers of the troposphere. Since we do not expect rapid changes in the physical and chemical composition of aerosols in the lower stratosphere and to reduce the uncertainty due to noise, especially during polar day, we chose a temporal and spacial resolution of 60 min and 150 m for the elastic profiles at 355 nm and 532 nm. Both signals can always be evaluated up to almost 30 km height.

2.2.2 Parameters and Fundamental Equations

It is well known from Klett (1981, 1985) that the solution of the elastic Lidar equation is:

$$P(z) = Const \cdot \frac{\beta^{tot}(z)}{z^2} \cdot \exp\left(-2 \int_{z_0}^z \alpha^{tot}(\hat{z}) d\hat{z}\right) \quad (1)$$

where $P(z)$ is the signal power at a given height z , $Const$ a constant determined by the instrument itself and its setup, $\beta^{tot}(z) = \beta^{aer} + \beta^{Ray}$ the total backscatter signal and $\alpha^{tot}(z) = \alpha^{aer} + \alpha^{Ray} + \alpha^{abs}$ the total extinction, depends on the choice of a Lidar ratio and on the choice of a boundary condition. The extensions aer , Ray and abs are the contributions of aerosols and Rayleigh scattering as well as the absorption by trace gases. β^{aer} needs to be prescribed at a certain altitude. Klett (1981, 1985) pointed out that the solution is stable and less critical dependent on the assumed boundary condition, when the integration is done "backwards": from large distances back to the Lidar.

We set the boundary condition, ϵ , in the altitude interval from 27 km to 30 km for March to November. In winter times, Polar Stratospheric Clouds (PSC) might occur in this altitude (Tritscher et al., 2021), so we lowered the fitting range to 24 km to 27 km. No PSCs were found in our data set after this adjustment. In this altitude range we assume, the average total backscatter coefficient in that interval is $\epsilon_{532} = 1.10 \cdot \beta^{Ray}$ and $\epsilon_{355} = 1.05 \cdot \beta^{Ray}$ for parallel and perpendicular. The signal-to-noise ratio, SNR, is calculated following Equation 2:

$$SNR(z) = \frac{P(z)}{P_{err}(z)} = \frac{P(z)}{\sqrt{P(z) + P_{bgd}}} \quad (2)$$

Where P_{err} is the estimated noise profile and P_{bgd} the averaged signal in an altitude range, which is dominated by electronic noise and background light in an altitude range of 52 km to 60 km. The Lidar signal $P(z)$ is given in the units of "counts" like in the photo-counting channels. This altitude range has a sufficiently small median signal-to-noise ratio. Therefore it is valid to assume there is just noise in the signal. The SNR is shown in Table 1 at an altitude of 20 km for the different wavelengths and directions of polarisation.

Table 1. Mean of the signal-to-noise ratio for all emitted wavelengths and directions of polarisation at 20 km altitude

λ [nm]	355	355 _⊥	532	532 _⊥	1064
SNR	52.7	7.5	15.6	10.6	4.06

With these high values of signal-to-noise ratio we conclude, that the data set is sufficiently good for a further evaluation of observations of physical features in the lower stratosphere. Since the depolarisation is very low (see Section 3.3.2), the signal is much weaker than for the parallel polarised light and therefore the SNR is generally smaller. Due to the small SNR at 20 km the signal of $\lambda = 1064$ nm is neglected for this study.

From the Lidar data we derive the following quantities: The Lidar ratio LR , aerosol depolarisation, $\delta^{aer}(\lambda)$ and the backscatter color ratio, CR , were calculated following Equation 3:

$$LR(\lambda) = \frac{\alpha^{aer}(\lambda)}{\beta^{aer}(\lambda)} \quad \delta^{aer}(\lambda) = \frac{\beta_{\perp}^{aer}(\lambda)}{\beta_{\parallel}^{aer}(\lambda)} \quad CR(\lambda_1, \lambda_2) = \frac{\beta^{aer}(\lambda_1)}{\beta^{aer}(\lambda_2)} \quad (3)$$

Here, β_{\perp}^{aer} and β_{\parallel}^{aer} are the backscatter coefficients with respect to the perpendicular and parallel laser polarisation, respectively, which are measured separately in the elastic 355 nm and 532 nm channels. For β^{aer} and δ^{aer} only 532 nm and 355 nm signals were used. The color ratio is defined with $\lambda_1 < \lambda_2$.

In contrast to the backscatter coefficient, β , the backscatter ratio, R , is dimensionless and does not decrease with increasing height. It is defined as in Equation 4:

$$R(\lambda) = \frac{\beta^{tot}(\lambda)}{\beta^{Ray}(\lambda)} = 1 + \frac{\beta^{aer}(\lambda)}{\beta^{Ray}(\lambda)} \quad (4)$$

An ideal, aerosol-free atmosphere would have $R = 1$ and increases with stronger pollution. The parameter R is therefore a good indication for aerosol existence.

160 With the ERA5 reanalysis data by Hoffmann and Spang (2021) the mean height of the tropopause was found in an average height of 9137 m with a standard deviation of 775 m at the study site of Ny-Ålesund during the Lidar measurement time and will be further discussed in Section 3.2. To investigate the properties of stratospheric aerosol, we focus on the altitude range between 10 km to 15 km; most of the troposphere is neglected and not shown in the plots.

165 2.2.3 Data Quality Management of Lidar Data

For this study, we only used automatically flagged data, which fulfilled the following criteria. Each wavelength and profile is tested and eventually removed individually:

1. Elimination of negative or too many (> 20) too low values ($R \leq 1.003$)
- 170 2. Elimination of artificial too high values ($\beta^{aer}(\lambda) \geq 5 \cdot 10^{-8}(\text{m sr})^{-1}$), when the error and parameter variables are equally large, more than $2/3$ of the data points are not-a-number or, if in more than $2/3$ of the cases values of $R \geq 2.0$ is reached
3. Clouds were detected in the troposphere with the criteria of two adjacent data points reach the limit of $\Delta\beta = 1 \cdot 10^{-4}(\text{m sr})^{-1}$

175 A qualitatively good profile has to fulfil all of these criteria at the same time. If one is not satisfied, the profile is eliminated out of the further processed data set. These criteria were chosen by manually looking at the data of different times of the year and atmospheric conditions with the aim of finding a good balance between eliminating as many disturbed profiles as necessary and keeping the good ones.

2.3 RS41 Radiosondes

180 At least once a day a radiosonde of the type RS41 by Vaisala is launched from AWIPEV at 11 UT. The data is processed according to the standards by the Global Climate Observing System (GCOS) Reference Upper-Air Network (GRUAN). A more detailed description of the radiosondes is given by Maturilli and Kayser (2017). With the quality checked and homogenised data set the Rayleigh atmosphere for the correction of Lidar profiles is calculated as well as the tropopause height.

2.4 CLaMS – Chemical Lagrangian Model of the stratosphere

185 To analyse the origin of air masses over Ny-Ålesund in 2021, we conducted 3-dimensional global simulations with the Chemical Lagrangian Model of the stratosphere (CLaMS; McKenna et al., 2002b, a; Pommrich et al., 2014, and references therein) as well as CLaMS back-trajectory calculations starting along the Lidar measurements. The global model simulations are driven by

horizontal winds from ERA5 reanalysis (Hersbach et al., 2020), provided by the European Centre for Medium-Range Weather Forecasts (ECMWF).

We use ERA5 data on 137 vertical model levels up to 0.01 hPa (80km), but in lower horizontal and time resolution referred to as ERA5 ($1^\circ \times 1^\circ$) (similar to Ploeger et al., 2021; Konopka et al., 2022; Clemens et al., 2024). Here, ERA5 data are truncated to a $1^\circ \times 1^\circ$ horizontal grid and a 6-hourly time resolution, whereby the vertical resolution is the same as in the original ERA5 reanalysis (about 0.3 to 0.4 km up to 20 km). ERA5 $1^\circ \times 1^\circ$ data are a computing-time-saving alternative to the full resolution ERA5 data and are used for 3-dimensional global CLaMS simulations. Vertical velocities are calculated following the diabatic approach by Ploeger et al. (2010). CLaMS uses a hybrid vertical coordinate (ζ). At pressure levels lower than 300 hPa, ζ can be interpreted as potential temperature (θ). Towards higher pressure levels, ζ transforms from an isentropic to a pressure-based orography-following hybrid coordinate. Global 3-dimensional CLaMS simulations are based on 3-dimensional forward trajectories and a parametrization of small-scale mixing depending on the shear in the large-scale wind flow (e.g. Pommrich et al. (2014)).

A 3-dimensional CLaMS simulation starting from 1st December 2020 until the end of 2021 to cover the entire time period of the year 2021 was performed including artificial surface origin tracers (Vogel et al., 2015, 2016, 2019). The surface origin tracers used in this study cover the entire Earth's surface and are associated to 32 defined regions in the model boundary layer that is about 2 km to 3 km above surface considering orography. They are released at the model boundary layer every 24 hours and are further transported (advected and mixed) to the free atmosphere. Here in this study was found, that regions in South Asia and in the tropical Pacific region (see Figure 2) have the major impact on air masses over Ny-Ålesund. The impact of all other regions of the world are minor and are summarised as 'residual' and 'residual ocean'. Air masses from South Asia are during summer associated with the Asian monsoon circulation and are here used as a marker for Asian monsoon air. Besides of continental regions in South Asia in addition parts of the Indian Ocean and the Bay of Bengal are included into the surface origin tracer for South Asia.

Starting with the onset of the Indian monsoon at the beginning of June, air masses from the Indian Ocean are transported long-range to the North of the Indian subcontinent where they are uplifted into the upper troposphere and lower stratosphere (e.g. Fadnavis et al., 2013; Lau et al., 2018; Nomura et al., 2021; Vogel et al., 2024). These air masses can uptake polluted air while passing over the Indo-Gangetic Plain where anthropogenic emission are higher compared to other regions in India caused by the dense concentration of industries and the high population (e.g. Fadnavis et al., 2016). Therefore, the tracer for the Indian Ocean and the Bay of Bengal are included into the marker for the Asian monsoon air.

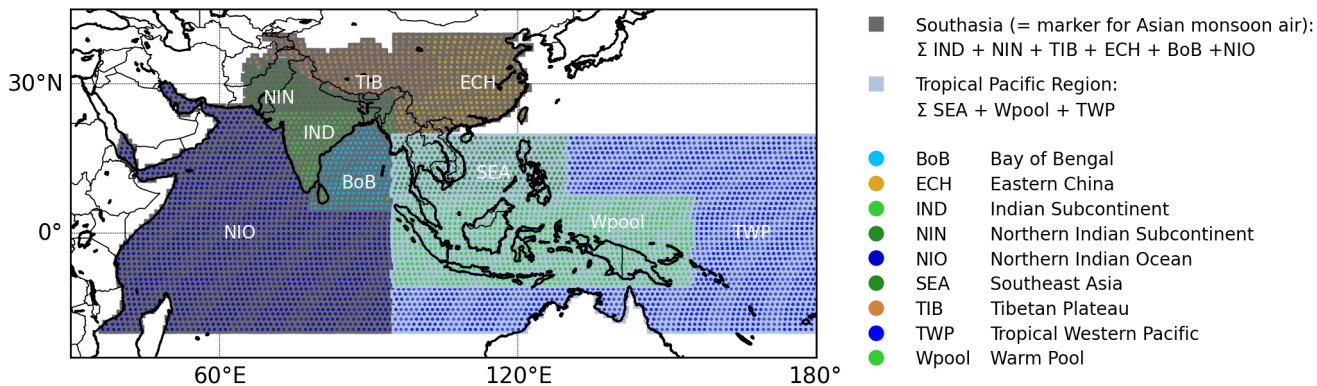


Figure 2. Geographical map showing the location of CLaMS's surface origin tracers in South Asia and the tropical Pacific region. Regions that mostly impacting the Lidar measurements in Ny-Ålesund are combined into two tracers referred to as 'South Asia' and 'tropical Pacific region'. The names of the surface origin tracers and their corresponding abbreviations are listed right beside the map

2.5 Moderate Resolution Imaging Spectroradiometer – MODIS

Moderate Resolution Imaging Spectroradiometer (MODIS) is a NASA satellite-based radiometer for Earth observations in 36 different spectral bands reaching from $0.4 \mu\text{m}$ to $14.4 \mu\text{m}$. It has, depending on the selection of bands, a spatial resolution of $220 \times 0.25^\circ \times 0.25^\circ$ and a temporal resolution of about two days. Wild fires are detected by the $4 \mu\text{m}$ and $11 \mu\text{m}$ bands by observing temperature anomalies relative to the background and absolute. In this study only the parameter Fire Radiative Power (FRP) from Aqua and Terra with a gridded spacial resolution of 1 km were used to characterise wild fires. More information can be found at MODIS.

3 Results

225 3.1 Wild Fires in North America and Russia

Wild fires occur regularly in the Northern Hemisphere and are a big source for aerosol, which can be lifted by several processes into the stratosphere (Ansmann et al., 2018; Ohneiser et al., 2021). Since air masses coming from Southeast Asia pass Russia and potentially even Canada the contribution of wild fires has to be considered, if the observed increased backscatter is influenced by these or is actually a direct signal from the monsoon region. The area "Russia" was defined within the rectangular defined by the geographical coordinates $77^\circ\text{N } 31^\circ\text{E}$ and $48^\circ\text{N } 180^\circ\text{E}$. "Canada" is defined by the corners at $71^\circ\text{N } 170^\circ\text{W}$ and $48^\circ\text{N } 52^\circ\text{W}$. Active wild fires are combined to monthly overviews. Figure 3 gives an overview about the median and mean Fire Radiative Power (FRP) measured by MODIS and averaged over all grid cells in the defined regions "Canada" and "Russia" from 2001 to 2022.

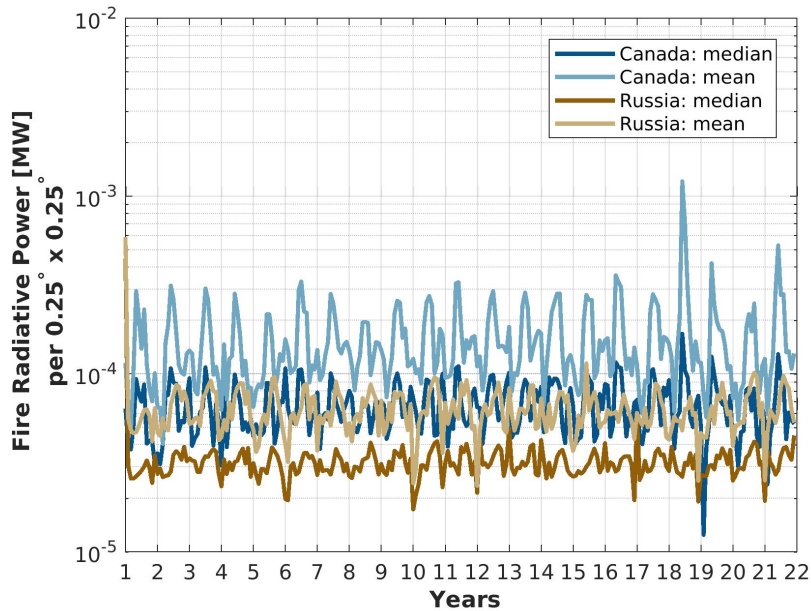


Figure 3. Overview of monthly averaged over the entire domain of either "Canada" or "Russia" fire radiative power (FRP) from 2001 – 2022

235 The wild fire season is in general more pronounced in Canada than in Russia throughout a year after averaging over the entire domain, since the variability from year-to-year is larger. It can be seen that the observed fire radiative power per grid cell does not increase during the observed 21 years, neither for Russia nor for Canada. The year 2021, which is selected for this study, shows a slightly increased higher FRP than other years in "Canada", but not in "Russia". Generally, strong biomass burnings produce clearly visible aerosol layers in Lidar observations, which are visible at least for several weeks (e.g. Zielinski et al., 2020; Ohneiser et al., 2021; Cheremisin et al., 2022). Since the aerosol layers from biomass burnings can reach altitudes
 240 of 20 km and more (Ohneiser et al., 2023), they would also influence our Lidar observations. We have not found such layers in our data set of 2021. Nevertheless the possibility must be considered that aerosol from biomass burning events lead to a higher stratospheric background in the Arctic in summer and autumn. We therefore conclude, that biomass burnings did not play a major role in the stratospheric aerosol budget in 2021.

3.2 Tropopause Calculations

245 The tropopause is a transport barrier between the troposphere and the stratosphere. We used the daily launched radiosonde from AWIPEV as well as ERA5-reanalysis data with a hourly global resolution of $0.3^\circ \times 0.3^\circ$ to determine the tropopause height for every Lidar profile. A full description of the two data sets are given by Maturilli and Kayser (2017) and Hoffmann and Spang (2021), respectively. As the definition of the tropopause the lapse rate definition, $\frac{dT}{dz} \leq 2 \frac{K}{km}$, of the World Meteorology Organisation (WMO) (Organization, 1957) is used for both data sets.

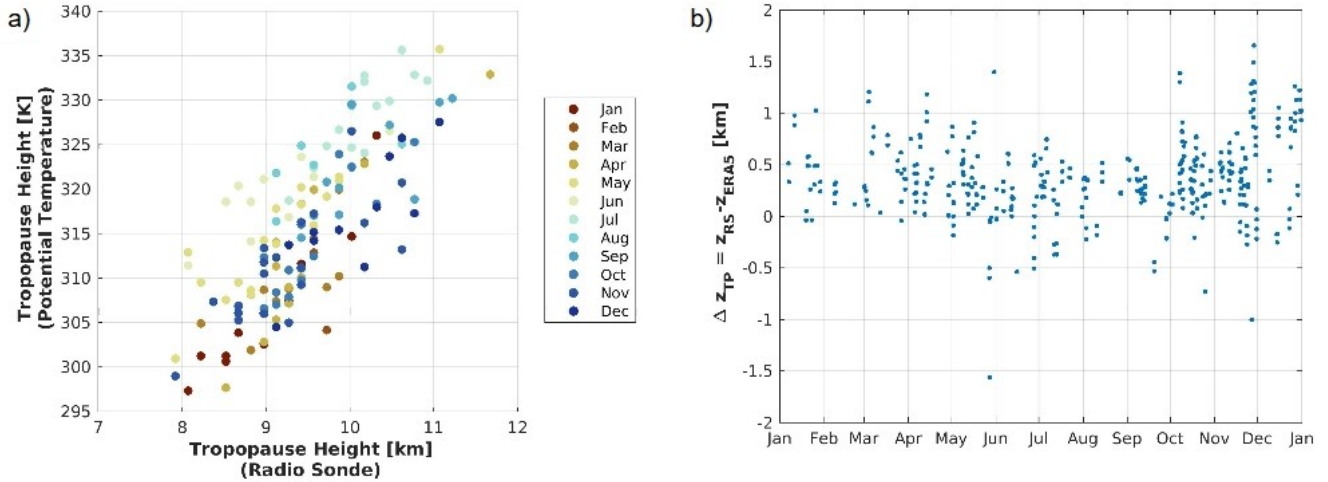


Figure 4. Annual cycle of Tropopause (TP) height measured by radiosondes in relation to the potential height of the tropopause derived from radiosondes (Figure 4a). The deviation between radiosonde tropopause height and ERA5 data is given in [km] (Figure 4b)

250 The tropopause height is shown in Figure 4. Since the measurements of radiosonde and Lidar are based on the geometric height, we transformed the altitude into potential temperature for better comparison with CLaMS results that is based on isentropic coordinates in the stratosphere. Figure 4a shows the geometric height in [km] and compares it with the corresponding potential temperature at this point, θ :

$$\theta = T \left(\frac{p_0}{p} \right)^{R/c_p} \quad (5)$$

255 Where T and p are the current temperature [K] and pressure [hPa] of the air parcel respectively, $p_0 = 1013$ hPa the ground reference pressure, R the gas constant of air and c_p the specific heat capacity. We set $R/c_p = 0.286$.

Every data point in Figure 4a corresponds to a measurement day. For comparison we also calculated the geometric tropopause with ERA5 data and compared it with radiosonde data throughout 2021 (Figure 4b). Since there is only one radiosonde launch
 260 a day, the Lidar measurement time and the launch does not always match, but the deviation between both measurements is maximum 12 h. On the other hand the data by Hoffmann and Spang (2021) has an hourly resolution corresponding to the chosen Lidar resolution. The median tropopause height in the radiosonde data is at 314.5 K (9011 m) and on average 329 m above the results of the ERA5-reanalysis model.

It can be seen, that in general the tropopause has a higher potential temperature in summer than in winter, but there is also a
 265 variability within a month. We conclude nevertheless that the tropopause is always at $\theta < 340$ K and aerosol above it is always in the stratosphere with increasing geometric height during summer months. Generally, the annual cycle of the tropopause observed by radiosondes is well represented in the ERA5 data.

3.3 Observations by KARL

In the following study we concentrate on the year 2021, since we have qualitatively good Lidar measurements throughout the entire year as well as every month with 481 h in 532 nm and 474 h for 355 nm. Due to additional campaign activity, especially in November and December, and due to high cloud cover fraction especially in summer (Graßl et al., 2022), the absolute number of measurement time varies a lot between months and seasons. February has the smallest amount of good data with 3 h, while in November 103 h of measurements are available. An overview of the monthly measurement time [h] is given in Table 2. Since the quality checks are performed for each emitted wavelength independently, a deviation in the available amount of measurement time might occur.

Table 2. Overview over the monthly available measurement time [h] for 355 nm and 532 nm after quality check

	Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sep	Oct	Nov	Dec	Total
355 nm	22	3	22	32	35	43	44	16	44	87	102	24	474
532 nm	21	3	22	34	42	44	44	17	44	85	102	23	481

Figure 5 shows the monthly medians of two aerosol backscatter coefficients β_{532} and β_{355} for the selected four heights of the lower stratosphere with a thickness of 20 K, which corresponds to about nine vertical measurement points with the given resolution.

280

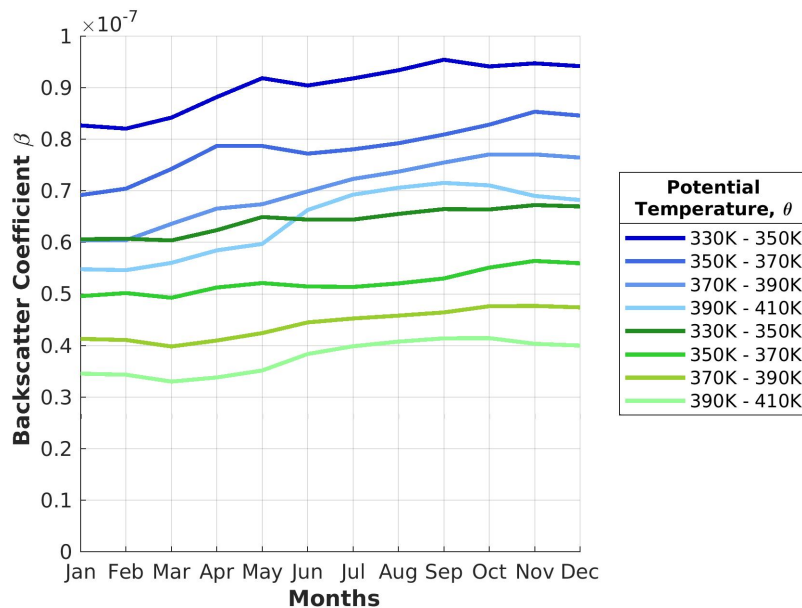


Figure 5. Monthly median backscatter values for β_{532} (green colors) and β_{355} (blue colors) for the lower stratosphere (330 K to 410 K potential temperature)

Generally, a weak annual cycle with minimal values of backscatter in spring and highest values in autumn can be seen. However, the curves are quite smooth, especially for 532 nm. At 400 K the amplitude of the annual variability is around 27% and 25% for 355 nm and 532 nm, respectively.

For all four selected layers β_{355} is always larger than β_{532} . Further, the backscatter decreases noticeably with increasing potential temperature. While the annual cycle of the backscatter in the lower two altitude ranges seems to be more variable and "disturbed", with a secondary maximum already in April or May. Between 370 K and 410 K, an increase of both backscatter coefficients is found from June to October 2021. As the backscatter and the amplitude of the annual cycle is larger in the UV we conclude that these changes refer to smaller particles, since the scattering efficiency strongly depends on the particle radius (Mie, 1908). The estimation of the effective radius of stratospheric aerosol by Mie calculations will follow in Section 3.4. Exemplary days will be later discussed and linked to CLaMS simulations in Section 3.5.

3.3.1 Color Ratio

The color ratio, CR , is a crucial parameter to determine the size of aerosols and is defined according to Equation 3. An overview of the CR in 2021 is shown in Figure 6. As a ratio of two small quantities, CR critically depends on noise and numerical assumptions in the Lidar evaluation (Foken, 2021). Hence, some noise is noticeable in form of rapid changing coloured dots in Figure 6.

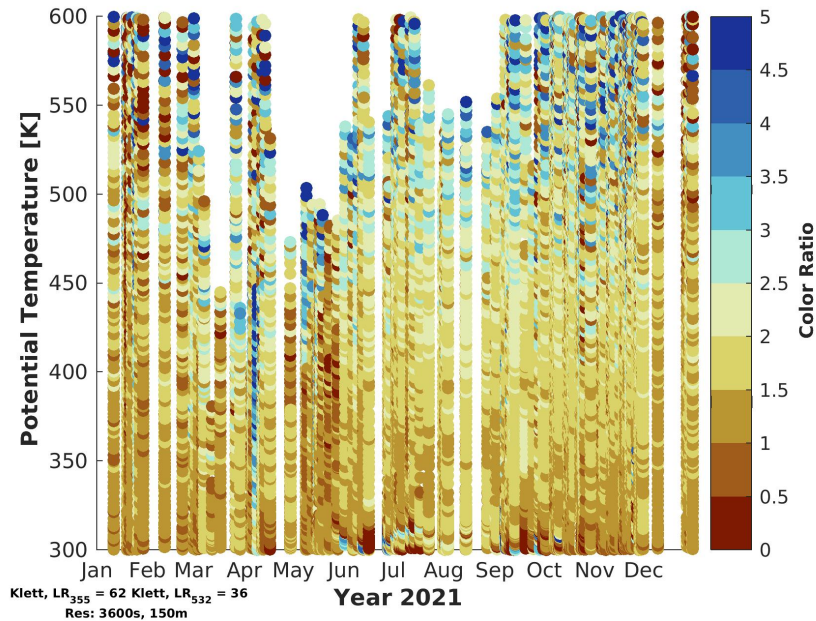


Figure 6. Overview of the color ratio, CR , throughout 2021

Typically, CR -values around 2 have been found. Remarkable is the fact that the second half of the year becomes more homogeneously with slightly larger values than the first half. From June to October a more homogeneous distribution of a $CR \sim 2$ is detected between 320 K and 440 K potential temperature. Since CR is a function of particle radius, the particle size decreases towards the end of the year and additionally with increasing height as it was also seen by Junge et al. (1961a). According to 300 Murphy et al. (2021) the generally larger particles (lower CR) in the first half of the year may be an indication of reduced tropospheric advection in the Arctic lower stratosphere in this time of year.

The relationship between β_{532} and CR is depicted for the previously defined four layers of the lower stratosphere in Figure 7. With this dependency a better understanding of the particle size and concentration can be made.

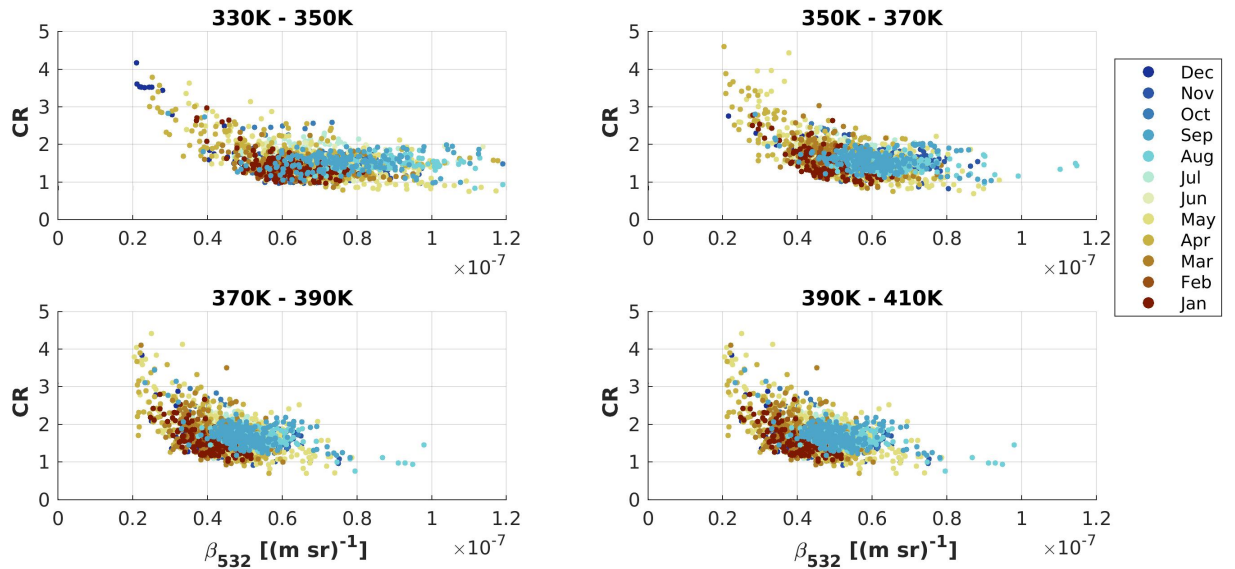


Figure 7. Dependency of CR and β_{532} for four selected height intervals of the lower stratosphere

305 While the spread in β_{532} decreases with height, CR remains in this same range of about $CR \in [1, 4]$. In altitudes of 330 K – 350 K a small influence of the troposphere might be seen in summer months, since the tropopause can reach these altitudes exceptionally. In the range of $CR \in [1, 2]$ all altitude layers have a backscatter coefficient larger at the end of the year than in the beginning. Under pristine conditions, $\beta_{532} \leq 5 \times 10^{-8} (\text{m sr})^{-1}$ we see always a clear dependence between CR and β : The smaller β (the clearer the atmosphere) becomes the larger CR becomes, because small particles show only a small scattering

310 efficiency. Hence, below $\beta_{532} = 5 \times 10^{-8} (\text{m sr})^{-1}$ the stratospheric β is mainly driven by the size of aerosols. This effect levels to above the limit of $\beta_{532} = 5 \times 10^{-8} (\text{m sr})^{-1}$. Larger than a certain limit the stratospheric aerosol does not grow apparently. This is in agreement to the annual cycle already shown in Figure 5. Since the color ratio does not change much in the four discussed intervals it also means the aerosol size is very similar, but only the concentration becomes lower the higher up the layer is located.

315

3.3.2 Aerosol Depolarisation

Another parameter to determine physical properties of aerosol by remote sensing is using the so-called Aerosol depolarisation, $\delta^{aer}(\lambda)$. It is defined according to Equation 3. The laser of KARL is linearly polarised and is always the reference direction of the scattered light. With Mie theory it can be shown, that spherical particles do not change the polarisation of the laser light,

320 while non-spherical ones do. Due to surface tension liquid droplets are in general spherical (Foken, 2021).

In general stratospheric aerosols have the property of $\beta_{\perp}^{aer} \ll \beta_{\parallel}^{aer}$. Therefore, it is expected that δ^{aer} becomes quite noisy for a standard aerosol measurement in higher altitudes. Due to generally low values of the depolarisation we only show and

discuss δ_{532} in this study.

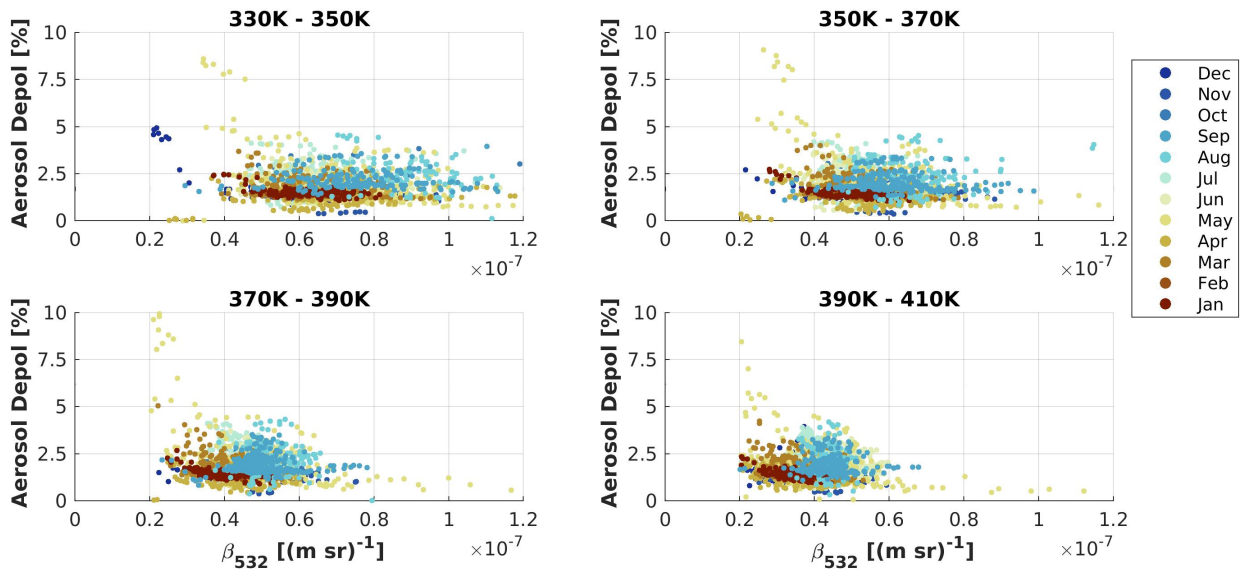


Figure 8. Dependency of β_{532} and δ_{532} for the four selected altitudes in the lower stratosphere

325 The overview of the aerosol depolarisation, δ_{532} , is given in Figure 8 for the different layers of the lower stratosphere color coded for every month. Due to case studies in Section 3.5 September is highlighted already here. The depolarisation is low ($< 10\%$) for all corresponding values of β_{532} . All four height intervals a common depolarisation is between $\delta_{532} \geq 1\%$ and $\delta_{532} \leq 4\%$. Additionally May is the most divers month with single cases of $\delta_{532} \rightarrow 10\%$, while January shows the smallest depolarisation. In general, summer has a higher depolarisation than winter months.

330 We define a "weak" depolarisation with $\delta_{532} = 2\% - 5\%$ (see Table 3) a "medium" depolarisation with $\delta_{532} = 5\% - 10\%$ (see Table 4). A "strong" depolarisation with $\delta_{532} \geq 10\%$, but was not found in the lower stratosphere of 2021.

Table 3. Relative occurrence of at least a "weak" polarisation ($\delta_{532} = 2\% - 5\%$) in different altitude layers of the lower stratosphere. All numbers are given in [%]. The low numbers in February can also be explained by the low number of measurement points (see Table 2)

	Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sep	Oct	Nov	Dec
330 – 350 K	3.81	0	21.43	7.29	13.94	13.96	25.76	49.33	17.50	3.96	0.39	4.89
350 – 370 K	3.81	0	22.92	5.83	14.83	14.02	31.82	50.56	14.17	1.92	0	4.80
370 – 390 K	2.91	0	18.67	6.85	17.18	15.28	27.53	34.67	12.36	1.68	0	5.78
390 – 410 K	3.89	0	24.29	3.91	18.17	16.90	24.01	32.50	9.58	0.94	0	18.40

335 A big variability between the months can be observed in Table 3 for depolarisation of $\delta_{532} = 2\% - 5\%$. While the depolarisation is in general lower in the winter months, it reaches its maximum in August and has a clear annual cycle. The depolarisation never reached values $> 2\%$. This indicates that the stratospheric aerosol observed above Ny-Ålesund is not constant throughout the year but changes its chemical composition and / or its shape (Foken, 2021). Furthermore, a more frequent depolarisation is found in lower altitudes than in higher ones. Our Lidar observations fit very well to the physical properties of aerosols in the Asian Tropopause Aerosol Layer (ATAL), which consist of organics, nitrate, sulfate and ammonium, especially in combination to Figure 8, where it can be seen, that the aerosol is also slightly non-spherical (Appel et al., 2022). The altitude, in which this aerosol layer in the Arctic was found also matches very well to the altitude of the ATAL.

Table 4. Relative occurrence of at least a "medium" polarisation ($\delta_{532} = 5\% - 10\%$) in the lower stratosphere from 330 K – 410 K. The numbers are given in [%]

	Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sep	Oct	Nov	Dec
330 – 410 K	0	0	0.10	0	2.08	0	0	0	0.15	0	0	1.16

340 Table 4 shows on the other hand the relative occurrence of $\delta_{532} = 5\% - 10\%$ for the entire layer of 330 K – 410 K. Only in a few months a "medium" depolarisation were observed. Even though the depolarisation has a annual cycle all values are very small compared to lower latitudes (Foken, 2021). No young extreme pollution events, like wild fire aerosol, dust intrusion or volcanic ashes with typical values of $\delta_{532} \geq 16\%$ (Foken, 2021) were found in the Lidar measurements over Ny-Ålesund in 2021. Therefore we conclude that our data set indeed mainly represents stratospheric background conditions even with the contribution of the ATAL. Therefore we conclude that our data set represents stratospheric conditions free from local events, however there is evidence of a seasonal impact of ATAL particles to the lower stratosphere during summer.

Figure 9 shows for two months with the lowest (February) and highest median depolarisation (August) in comparison for the lower stratosphere and further investigate the difference between the relative occurrence of depolarisation.

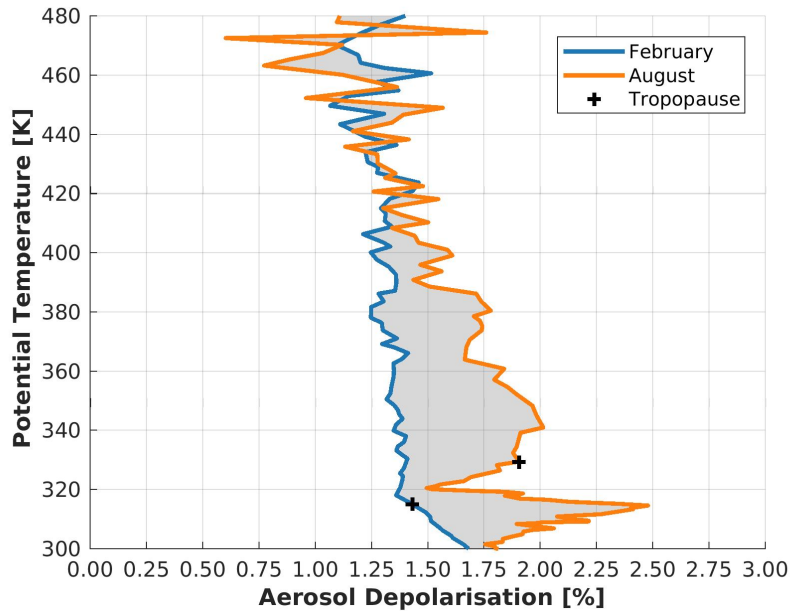


Figure 9. Mean aerosol depolarisation, δ_{532} , for the two exemplary months February and August. The grey shaded area shows the difference between both months

350 While February has a more or less constant depolarisation at $\delta_{532} = 1.35\%$ in the atmosphere between 320 K and 480 K, August on the other hand has in general higher depolarisation values, especially in the lower parts of the atmosphere (until about 420 K). The grey shaded area highlights the difference between both months. Especially in the troposphere and lower stratosphere a significant deviation can be found and shows the impact of the ATAL. Additionally, the summer month also has a maximum around 315 K, where it reaches $\delta_{532} = 2.5\%$, followed by a second, local maximum at around 340 K with

355 $\delta_{532} = 2.0\%$. From that point on it decreases to about $\delta_{532} = 1.25\%$ and converges to the same value as in February. Due to solar radiation in August, the data becomes noisy quicker than during polar night. At an altitude of 400 K the depolarisation in August is still significantly higher than in February. The median tropopause heights according to radiosonde data is in February at 307 K height, while it is at 324 K in August. This plot also shows that even if no distinct biomass burning layer has been

360 seen in our Lidar data during summer, the slightly less spherical shape of the particles indicates a non-sulfuric component that is an evidence for an impact of ATAL particles on the Arctic background aerosol up to ~ 450 K in August 2021, even though aged biomass burning aerosol may still contribute to the stratospheric aerosol background, which were even found in altitudes of up to 27 km (Ohneiser et al., 2023).

3.3.3 Seasonal variability of stratospheric Arctic background Aerosol

365 Our measurements indicate evidence for the impact of ATAL particles on the Arctic stratosphere during summer, therefore we now present the annual cycle of β_{355} and β_{532} of the background aerosol in Figures 10 and 11, respectively, for two different heights of 360 K and 380 K. Data from ± 10 K around the chosen level are taken into account for the following study.

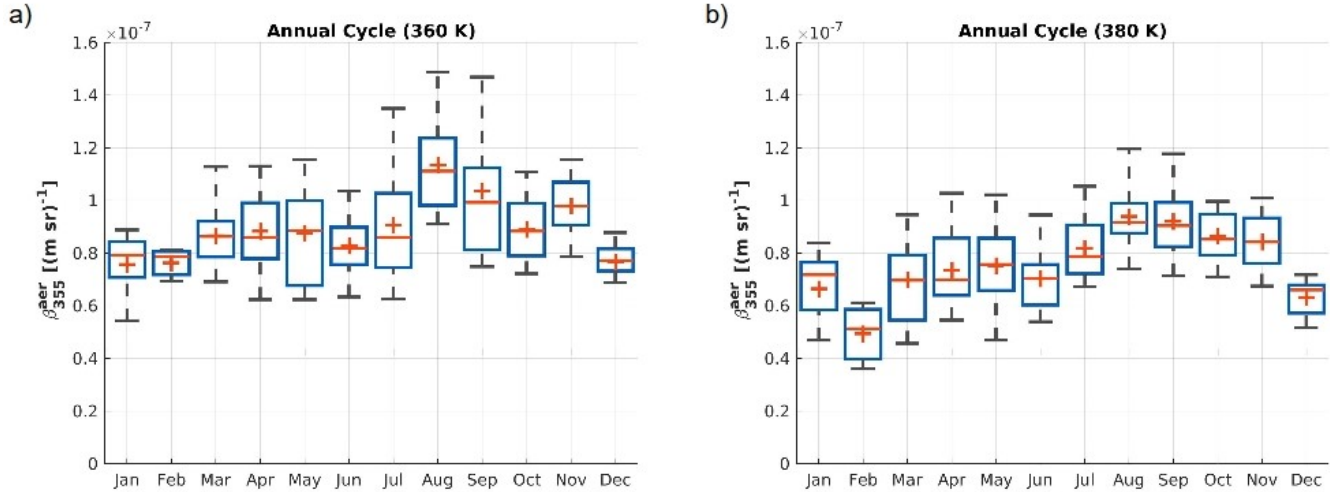


Figure 10. Annual cycle of stratospheric aerosol backscatter coefficient β_{355} in 360 K and 380 K height. The red symbols, $-$ and $+$, show median and average, respectively. The blue box symbolise the 25th and 75th percentiles, respectively. The black lines show accordingly the 9th and 91th percentiles

The annual cycle of β_{355} in 360 K altitude (Figure 10a) shows a small seasonal dependency: While the first half of the year is characterised by lower values of the backscatter coefficient, the mean values of August, September and November are $> 10^{-7}(\text{m sr})^{-1}$. While winter (DJF) is the clearest season, May, July and September are the months with the broadest distribution. Overall the median and mean values are for all months very similar, indicating that β_{355} follows almost a Gaussian distribution.

Likewise there is a similar annual cycle in the data at the 380 K level (Figure 10b) with larger values from June to November. Also here winter (DJF) is the clearest season. While the absolute values of β_{355} are consistently lower than at 360 K height, the relative spreading of β_{355} is also smaller. Typical values are for 360 K altitude are $\beta_{355} \in [7, 11] \times 10^{-8}(\text{m sr})^{-1}$, they are smaller for the 380 K-level, $\beta_{355} \in [6, 10] \times 10^{-8}(\text{m sr})^{-1}$. In summary, a clear annual cycle in the stratospheric backscatter coefficient has been found.

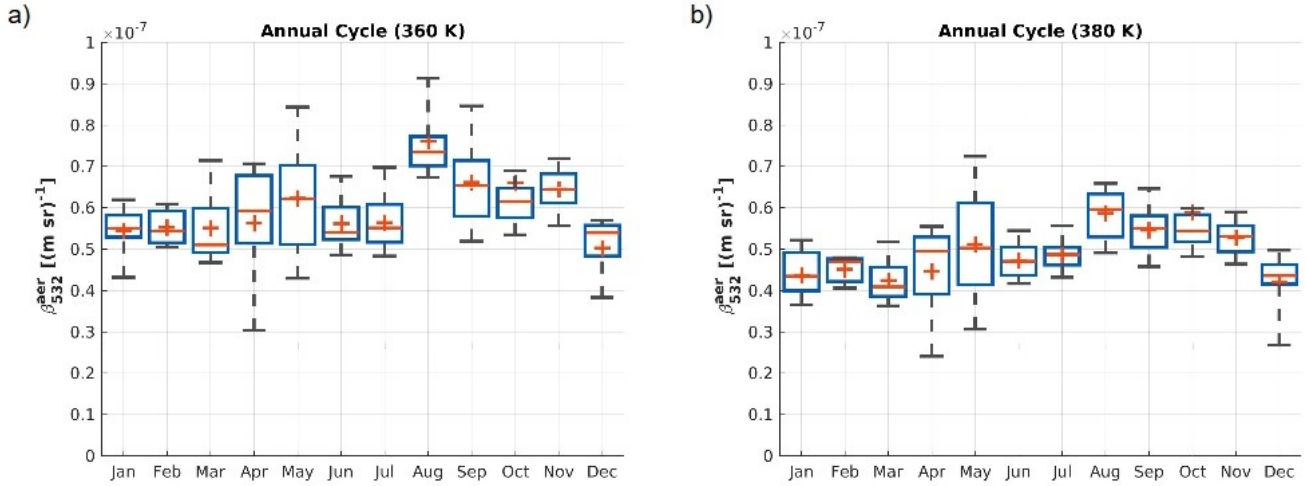


Figure 11. Annual cycle of stratospheric aerosol backscatter coefficient β_{532} in 360 K and 380 K height. The red symbols, $-$ and $+$, show median and average, respectively. The blue box symbolise the 25th and 75th percentiles, respectively. The black lines show accordingly the 9th and 91th percentiles

While the values of β_{532} are by about a factor of 1.5 to 2 smaller than for β_{355} a similar pattern is shown in Figure 11a compared to β_{355} with some differences: The variability in the first half of the year is more pronounced and the increased backscatter coefficient peaks in August. On the other hand β_{532} is less homogeneously distributed than β_{355} throughout the entire year. The 380 K-level (Figure 11b) shows the same trend throughout the year with a higher month-to-month variability in April and May. Median and mean values of β_{532} spread more than for β_{355} , indicating a more asymmetrical distribution and more variability within a month. Typical values are for 360 K altitude $\beta_{532} \in [5, 7] \times 10^{-8}(\text{m sr})^{-1}$ and $\beta_{532} \in [1, 6] \times 10^{-8}(\text{m sr})^{-1}$ for the 380 K-level.

385

3.4 Mie Calculus for Particle Size Estimations

Absorption and scattering of a single spherical particle surrounded by a non-absorbing medium is described by Mie scattering (Mie, 1908). Particles can have a complex refractive index, m , in general:

$$m = n + ik \tag{6}$$

390 where $\Re(m) = n$ describes the scattering and $\Im(m) = k$ the absorption of the light by aerosols. The scattering efficiency factor is a parameter, which is defined as the ratio of the scattering cross section, σ_{eff} (Equation 7), to the geometrical cross section. Therefore, it is a good measure to determine the effective radius, r_{eff} , for a mono-modal log-normal aerosol size distribution:

$$\sigma_{eff} = \pi \int_0^{\infty} Q_{sca}^{FF}(r_{eff}) r^2 f_1(r_{eff}) dr_{eff} \quad (7)$$

395 The scattering efficiency $Q_{sca}^{FF}(r_{eff})$ for the far field, FF , and f_1 the mono-modal log-normal aerosol size distribution following the calculations according to Bohren et al. (2023).

We expect small or already aged aerosols in the stratosphere (Turco et al., 1982; Kremser et al., 2016) and are therefore able to estimate the effective radius of these particles by the Library for Radiative Transfer – libRadtran (Mayer and Kylling, 400 2005; Emde et al., 2016). As Kremser et al. (2016) pointed out, sulfur chemistry is a very common and frequent process in the stratosphere. Since sulfate containing particles are not the only aerosol type in the ATAL, we also chose biomass burning particles as the second example as a highly absorbing particle type.

We roughly estimate an effective radius for the observed aerosol, which can have a lifetime of up to several weeks (Openheimer et al., 1998). Since the refractive index is wavelength-dependent two different indices were chosen for sulfate: 405 $m_{532}^{Sulfate} = 1.44 + 10^{-4}i$ (Russell and Hamill, 1984; Yue et al., 1994) and $m_{355}^{Sulfate} = 1.41 + 0i$. The imaginary part of Equation 6 can be neglected for $m_{355}^{Sulfate}$, since sulfate does not absorb in UV (Beyer and Ebeling, 1998; Washenfelder et al., 2013). As a comparison we also calculated the effective radii for a strongly absorbing aerosol type, like from wild fires and chose a refractive index of biomass burning aerosols (BB) of $m^{BB} = 1.52 + 10^{-2}i$ for 355 nm and 532 nm (Sumlin et al., 2018).

With the above mentioned assumptions we used the two refractive indices for 355 nm and 532 nm, a log-normal distribution 410 of particles with a standard deviation of $\sigma = 1.5$ up to a maximum size of 600 nm and a bin width of the distribution of 10 nm.

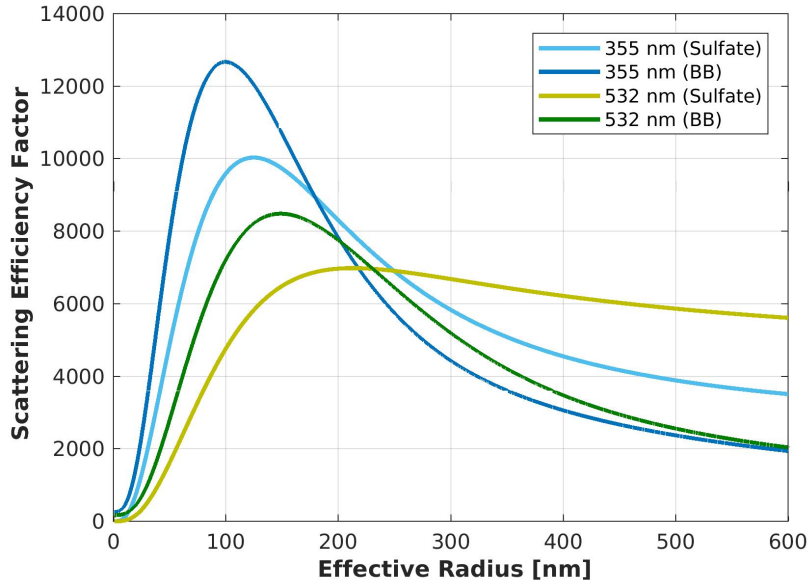


Figure 12. Scattering efficiency factor, Q_{sca}^{FF} , in arbitrary units for the emitted 355 nm and 532 nm laser beam for a log-normal distribution of purely sulfate particles and strongly absorbing BB aerosols

As Figure 12 shows, the larger the real part of the refractive index the quicker the scattering efficiency drops for larger effective radii. Simultaneously the maximum of scattering efficiency moves towards larger r_{eff} for larger real parts and wavelengths. On the other hand, there is not a big difference in scattering efficiency for large particles especially for biomass burning aerosol observed in the wavelengths 355 nm and 532 nm. Even though the efficiency is much lower for smaller particles in
415 532 nm. An overview of the different effective radii correlated to typical color ratios (see Section 3.3.1) are given in Table 5.

Table 5. Calculated effective radii, $r_{eff}[nm]$ corresponding to typical Color Ratios with $m^{BB} = 1.52 + 10^{-2}i$, $m_{532}^{Sulfate} = 1.44 + 10^{-4}i$ and $m^{Sulfate}_{355} = 1.41 + 0i$

r_{eff}	CR			
[nm]	2.00	2.25	2.50	2.75
Sulfate	97	82	70	59
BB	84	72	61	52

It is expected to find a small effective radius for stratospheric aerosols distributions, since the particles form in chemical reactions in Aitken mode on salt solutions or salt embryos. The reaction takes also place at stratospheric temperatures of $T \leq -55^{\circ}C$ (Friend et al., 1973). With these small effective radii corresponding to an height dependent value of CR (see Figure 6) we conclude, that in general the effective radius of aerosol in the lower stratosphere is relatively small (Murphy

420 et al., 2021) and the particles are sorted according to their size by for example sedimentation processes with an accumulation of largest particles close to the tropopause.

3.5 Origin of air masses inferred from CLaMS simulations

Backscatter coefficients, color ratio and depolarisation of the aerosol particles detected by KARL show evidence that the ATAL has an impact on the Arctic lower stratosphere. To confirm this assumption we link KARL measurements to Lagrangian transport simulations. To identify the source regions of the aerosols detected by KARL, global 3-dimensional CLaMS simulations including surface origin tracers were analysed and discussed for two exemplary days in September 2021 (13th and 28th). The identification of possible source regions of the aerosol particles found above Ny-Ålesund (Section 3.5.1) are investigated by global 3-dimensional CLaMS simulations including surface origin tracers (Section 3.5.2) as well as CLaMS back-trajectories (Section 3.5.3).

430 3.5.1 Seasonal variability of air mass origins over Ny-Ålesund

The contributions of young air masses from South Asia and the tropical Pacific over the course of the year 2021 to the air over Ny-Ålesund is inferred for the heights (360 ± 5) K (Figure 13a) and (380 ± 5) K (Figure 13b), respectively. Mean values are calculated for the surface origin tracers for each day.

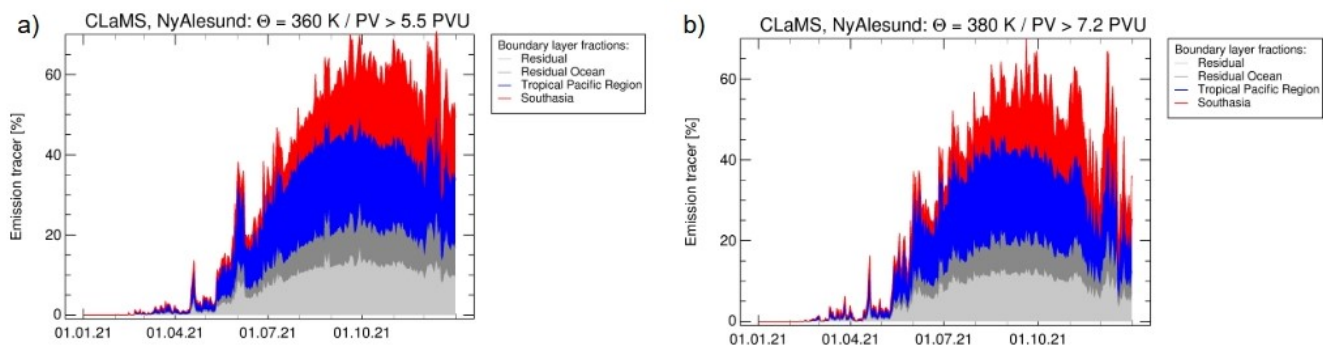


Figure 13. Contribution of different surface emission tracers from South Asia, Tropical Western Pacific, Residual and Residual Ocean to the lower stratosphere above Ny-Ålesund at 360 K (Figure 13a) and 380 K (Figure 13b) potential temperature for the year 2021. The 5.5 PVU and 7.2 PVU surface takes the climatological isentropic transport barrier at 360 K and 380 K, respectively, into account (Kunz et al., 2015)

Artificial tracers released in the model simulations since 1st December 2020 in Figure 13 do not reach Ny-Ålesund at levels of potential temperature of 360 K and 380 K before May 2021. During summer and autumn, the highest fractions of tropospheric air over Ny-Ålesund are from South Asia and the tropical Pacific region. Especially in September and October almost 20% of the air is from South Asia. These model results support the evidence that the Arctic lower stratosphere is impacted by aerosol particles from the ATAL (Yu et al., 2017; Bian et al., 2020). All other regions combined are of minor importance. With few exceptions the "Tropical Pacific" contributes constantly more, sudden changes in its impact are rare.

440 The accumulation of young air masses in the lower stratosphere over Ny-Ålesund is lower 100% during the year 2021. The remaining fraction consists of aged air, that is older than 1st December 2020 when the CLaMS simulation was initialised.

3.5.2 Three-dimensional CLaMS simulation

Figure 14 gives for two exemplary days in September (13th and 28th) the distribution of the surface origin tracer for South Asia (a marker for air from the Asian monsoon region) over northern Europe at 360 K and 380 K potential temperature. A high
445 fraction from South Asia is found over the Arabian Peninsula marking the western outflow of the Asian monsoon anticyclone (e.g. Vogel et al., 2016). Air masses contributing to the ATAL can be transported eastwards along the subtropical jet and enter the lower stratosphere by quasi-horizontal transport. As a result, thin filaments with enhanced signatures of air mass tracers origin from South Asia were found over the northern Atlantic Ocean and Europe (e.g. Vogel et al., 2016; Wetzel et al., 2021; Lauther et al., 2022). On 28 September a filament located over Ny-Ålesund with contribution from South Asia up to 40% is
450 simulated. This filament was not found on 13 September.

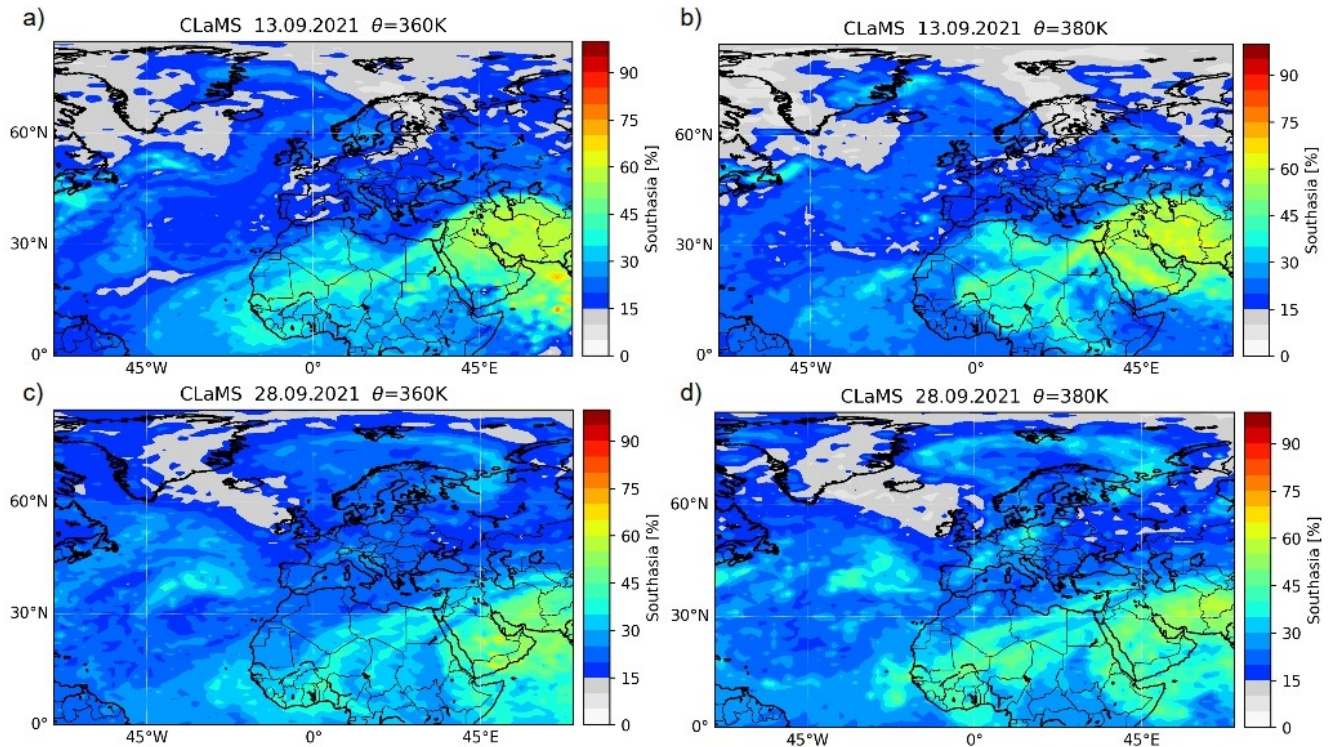


Figure 14. Horizontal distribution of the fraction of air originating in South Asia (details of the region can be found in Figure 2) at 360 K (left column) and 380 K (right column) potential temperature on 13 (top), and 28 (bottom) September 2021

Figure 14 shows in the stratosphere from Southeast Asia for the two different heights of 360 K and 380 K. The contribution is quite low on 13 September in 360 K in 380 K altitude (Figures 14a and 14b). As it can be seen in both figures, the contribution is constant over entire Europe, but increases with increasing height and changes on time scales of less than a day. As it
455 can be seen in both altitudes the impact is already very low over Svalbard ($\approx 10\%$). On 28 September a filament was seen faint in 360 K altitude (Figure 14c) but very pronounced in 380 K (Figure 14d). The presented values are strong for Ny-Ålesund up to 20% are comparable small compared to other sites at lower latitudes.

When comparing Figures 13 and 14 with each other the contributions of the two main sources of tracers can be identified. The region 'Tropical Pacific' is responsible more for the background amount, which can be observed in Figure 14. Contrary
460 to this, 'Asia' has a smaller contribution but shows also a very large day-to-day variability. These spikes can be nicely seen as filaments in the three dimensional simulations of tracers of Figure 14.

The following preliminary conclusion can be made: There is a large day-to-day variability in the distribution of young air masses from Asia contributing the Arctic lower stratosphere. This variability strongly depends on the dynamics of the Asian
465 monsoon anticyclone itself and the transport of air masses from Asia to the extra-tropical lower stratosphere.

3.5.3 Case studies using back-trajectory calculations

Backward trajectories are well suited to analyse the detailed transport pathway of an air parcel and therefore complement the three-dimensional CLaMS simulations (e.g. Vogel et al., 2014; Wetzel et al., 2021; Lauther et al., 2022). Within this study, we calculated an ensemble of backward trajectories for the two exemplary days (13 and 28 September 2021). Within $\pm 2^\circ$ in
470 latitudinal and longitudinal direction around Ny-Ålesund and in a height range of 360 K to 380 K, 891 trajectories have been calculated in total for each day. Markers in Figure 15 show the location, where the trajectory first reaches the boundary layer. The color indicates the transport time.

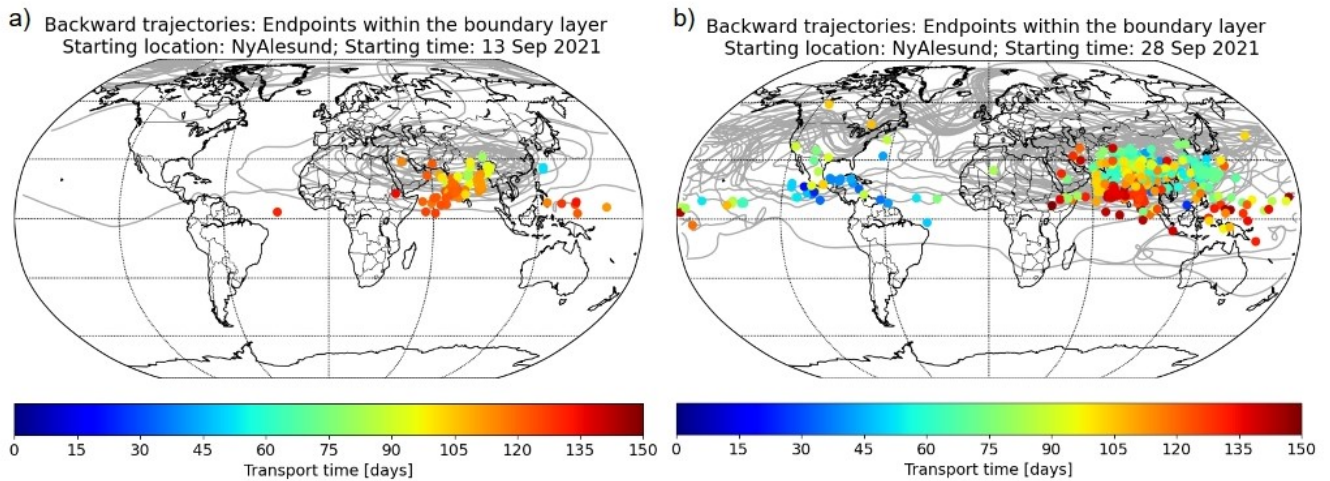


Figure 15. Location of trajectory end points where they reach the model boundary layer (BL). Trajectories are initialised on 13 and 28 September within 360 K and 380 K in altitude and $\pm 2^\circ$ horizontally around Ny-Ålesund. The end points are colour-coded by the transport time between reaching the Boundary Layer the first time and the Lidar measurement at Ny-Ålesund

As it can be seen in Figure 15a on 13 September in total 41 trajectories (4.6%) reached the boundary layer in Southeast Asia after travelling for about three to four months, 6 other trajectories (0.7%) are outside of the monsoon region. 850 trajectories
475 (89.6%) did not touch the boundary layer at all within the model run time.

The situation is a bit different for 28 September 2021 (Figure 15b), where the fraction of contacts outside of the monsoon region is larger (7.6%), but also the fraction of end points in Southeast Asia (37.9%). On this day only 54.4% of the 891 trajectories did not reach the model boundary layer within the computational time. It took the air parcel on average 87 days to reach Ny-Ålesund on 28 September. As it can be seen on both days, back-trajectories also end in different parts of the world,
480 like the equatorial pacific ocean but also central America. This result agrees very well with the forward trajectories of aerosol tracers of Section 3.5.2. The average transportation time from the Asian summer region to Ny-Ålesund is about one month longer on 13 September (111 days) than on the 28 September (87 days).

The presented numbers are also given in Table 6 for a better overview with the absolute and relative numbers of back-trajectories touching the boundary layer inside and outside of the monsoon region in South Asia.

Table 6. Contact points of all calculated back-trajectories, which are presented in Figure 15 with their relative and absolute frequencies as well as the mean propagation time between the Asian summer monsoon region and Ny-Ålesund

	13 Sep 2021	28 Sep 2021
Boundary Layer	41 (4.6%)	213 (23.9%)
Asia	35 (3.9%)	159 (17.8%)
Residual	6 (0.7%)	54 (6.1%)
no contact	850 (95.4%)	678 (76.1%)
Sum	891 (100%)	891 (100%)
Mean transport time	111 days	87 days

485 Additionally one trajectory ensemble is given for each day in Figure 15. As it can be seen in both cases, the artificial tracer circulates from Ny-Ålesund first around the pole, before it moves to lower latitudes until it reaches South Asia, where it also circles, until it reaches the boundary layer. Since both trajectories have similar pathways, we conclude that the intake of aerosols into the lower stratosphere depends on the local conditions in South Asia and the effectivity for aerosols being lifted into the ATAL.

490 **3.6 Detailed Analysis with KARL**

In the following these two exemplary days will be discussed in detail. On one of them CLaMS observed a filament in different heights. The other day is an example with a comparable clear stratosphere. As a reference day a profile of February was additionally plotted to emphasise the difference between seasons and days. Here the backscatter ratio, defined according to Equation 4, is used as it does not decrease with increasing height and layers are easier to detect.

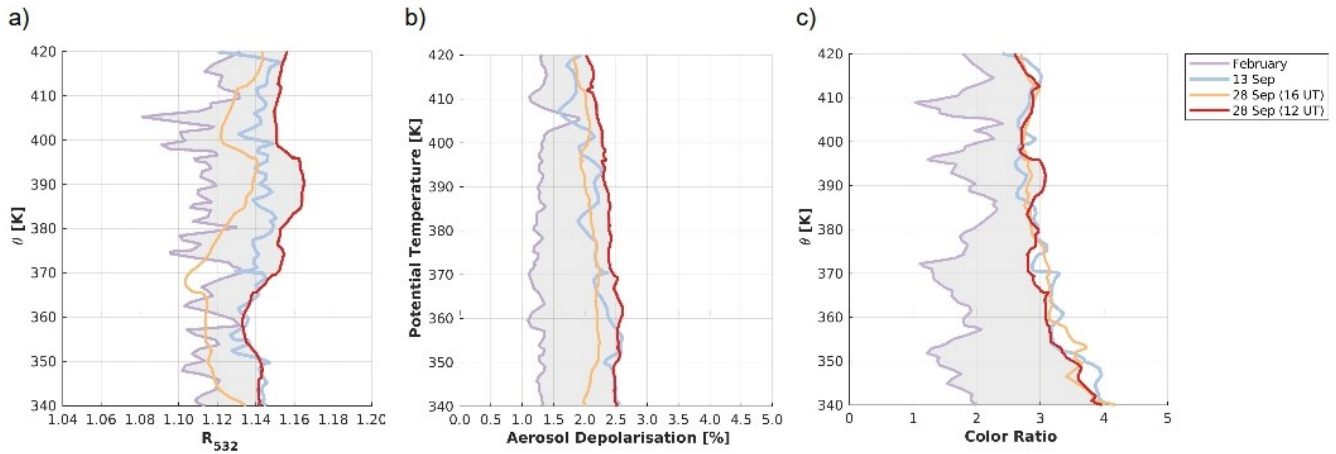


Figure 16. Profiles of different parameters, R_{532} (Figure 16a), δ_{532} (Figure 16b) and CR (Figure 16c), for the two exemplary days. 28 September is shown here twice, since the Lidar measurement simultaneously took place with the model output at 12 UT. As a comparison the Lidar results are also shown at 16 UT for 28 September. The shaded area highlights the difference between February and the filament on 28 September

495 Figure 16 shows for the two exemplary days in September with and without a detected filament by CLaMS, R_{532} (Figure 16a), δ_{532} (Figure 16b) and CR (Figure 16c). The shaded area highlights the difference between a typical profile of February, which is also not affected by the ATAL, and the detection of the filament on 28 September at 12 UT. Since the model output of CLaMS is shown in Figure 14 for 12 UT and the size and movement of the filaments indicate only a short temporal increasing of the backscatter ratio, the measurements by KARL on 28 September are shown twice: for 12 UT and 16 UT. Due to the horizontal transports in the stratosphere the filament of this day has already moved out of the line-of-sight of the Lidar for the later time. According to the radiosonde data, the tropopause is well defined on the 28 September 2021 at 332.9 K. The reanalysis data of tropopause heights from the data repository Jülich Data (Hoffmann and Spang, 2021) reveals a tropopause height of 329.1 K to 335.1 K over Ny-Ålesund for the two Lidar profiles. For this case the air masses of troposphere and stratosphere are well separated by each other and an exchange is not possible.

505

Generally, it can be seen, that all parameter, backscatter ratio, aerosol depolarisation and color ratio, are larger and less variable for all days in September than in February as it was already shown in Figures 10 and 11. This implies that the stratospheric aerosol is already aged and well-mixed.

The backscatter ratio reveals a layer on the 28 September at 12 UT between 360 K and 400 K. This one is also seen in CLaMS as filaments (Figures 14c and 14d). KARL and CLaMS also agree that the layer is more pronounced at 380 K than at 360 K. According to KARL the layer reaches until 400 K altitude. Compared to 12 UT the layer at 16 UT has already a significantly decreased backscatter ratio, which is almost on the background level (defined as the mean backscatter ratio of the measurements not affected by the filament). From Figure 16a it can be seen that the strongest impact of the filament occurred around

510

390 K. R_{532} rises from about $R_{532} = 1.14$ to $R_{532} = 1.165$. This corresponds to an increase of aerosol backscatter by about
515 15%.

The depolarisation is in general quite small with values around 2.5% at 340 K, a maximum of all profiles at around 360 K. δ decreases with increasing heights to values around 2% at 420 K. These small values indicate nearly spherical particles throughout the lower stratosphere. A slight increase in depolarisation throughout the entire lower stratosphere is found for 12 UT of 28 September, when the filament passes above the station.

520 The same result of small particles can also be found in the color ratio in Figure 16c. While CR reaches values around 4 at 340 K, it decreases to $CR \approx 2.9$ in the altitude range of 370 K to 410 K. This means, aerosols are smallest and reach a constant size distribution. The filament of 28 September changes the overall color ratio only in the altitude range of 385 K to 395 K by $CR \approx 0.2$, meaning that the particle size is slightly smaller in times with a filament. February reveals a comparable low color ratio (larger particles), but also a lower backscatter ratio. We therefore conclude, aerosols are in general larger but less
525 abundant in February than during September. The fluctuations are also larger, which indicates a less effective vertical mixing of the lower polar stratospheric air.

Observations by KARL and modelling studies by CLaMS reveal a good agreement between the results. Filaments, which were forecasted by CLaMS were found in KARL data. Physical properties of the stratospheric aerosol in the Arctic in September reveal very similar properties as in the ATAL. We can therefore conclude, the filaments above Ny-Ålesund originate from
530 the ATAL. Stratospheric aerosol in other seasons, like February, has different physical properties and can clearly be distinguished from aerosols arriving in September.

4 Discussion

In MODIS data we did not find any hint, that 2021 was exceptionally in regards to the wild fire season on the northern hemisphere. The detailed studies of Canadian wild fires by Haarig et al. (2018) and Baars et al. (2019) reveal a aerosol depolarisation
535 of up to $\delta_{532} \leq 20\%$ in the stratosphere. An explanation for the deviation of the aerosol depolarisation is the very low relative humidity in the stratosphere (Maturilli and Kayser, 2017). Since the aerosol remained there for already some days, it dried out and retrieved a spherical shape, with which a depolarisation of $\delta_{532} \approx 2\%$ is expected and was found in our data. A similar strong exemplary wild fire event, like it was analysed by Ansmann et al. (2018), was not found in our data.

540

In this work we showed a clear annual cycle of the stratospheric aerosol for the Arctic in the year 2021. In summer and autumn the aerosol backscatter is larger (by about 20%) than in winter and spring, especially in the UV. At ALOMAR (Arctic Lidar Observatory for Middle Atmosphere Research, northern Norway) a study about stratospheric aerosols was performed by Langenbach et al. (2019) for the years 2014 – 2017. The authors found a significantly increased backscatter ratio in the lower
545 stratosphere (12 km to 18 km) in the months July and August for an emitted wavelength of $\lambda = 1064$ nm. Hence, the Arctic or sub-Arctic stratosphere is not pristine, especially aerosol contamination in summer may be typical. Compared to the strato-

spheric background aerosol at ALOMAR, we observe the aerosol layer in lower altitudes (in km) and the strongest appearance in August and September. Deviations between these two measurement stations are probably due to the geographical location and the in general lower tropopause height in the Arctic than in lower latitudes.

550

After aerosol particles or their gas-phase precursors were uplifted into the lower stratosphere by the monsoon system (e.g. Brunamonti et al., 2018; Hanumanthu et al., 2020; Weigel et al., 2021a, b; Appel et al., 2022), the air mass are confined within the Asia monsoon anticyclone from which eddies and filaments can be shed to the east. These polluted air masses are further transported into the stratosphere of the Northern hemisphere by the breaking of Rossby waves (e.g. Waugh, 1996; Vogel et al., 555 2016), therefore it is expected to find filaments with an enhanced number of aerosol particles in higher latitudes. Aerosols contributing to the ATAL can circulate a few times around the Asian monsoon anticyclone before leaving Asia. In addition they can be transported along the subtropical jet a few time around the globe until they reach the Arctic. There it can be then further investigated by the Lidar KARL by assuming similar propagation velocities as Jumelet et al. (2020). Afterwards the aerosol load decreases slowly by sedimentation and other removal processes happening in the stratosphere described in general 560 by Junge et al. (1961a). From December to July the aerosol load is low and about constant. This is in good agreement with (e.g. Junge et al., 1961a; Kremser et al., 2016), who observed both an equilibrium state of new particle formation out of the gas phase and removal processes from the stratosphere into the troposphere. The physical properties of the filaments in the Arctic stratosphere reveal that the particles originate from the ATAL.

565 Since various types of aerosols are presented in the ATAL, we estimated the effective radii of two exemplary types: One with purely sulfate-containing particles, the other highly absorbing one from biomass burnings. Mie calculus revealed an effective radius of 52 nm to 97 nm as two more extreme examples of the aerosol classes. Since the depolarisation is comparably small, we also conclude that the predominant aerosol type has a spherical shape, which agrees to the in-situ measurements of Junge et al. (1961a); Hofmann et al. (1975); Murphy et al. (2021) in lower latitudes or of aged aerosols emitted by wild fire events 570 agree with the calculated effective radius of the in the Arctic observed aerosols (120 nm to 160 nm) and additionally enhances the assumption of a log-normal particle size distribution (Dahlkötter et al., 2014; Baars et al., 2019).

Taking now the height-dependent calculations of Junge et al. (1961a) into account sedimentation velocities of about $v_{min} = 3 \times 10^{-3} \frac{\text{cm}}{\text{s}}$ to $v_{max} = 2 \times 10^{-2} \frac{\text{cm}}{\text{s}}$ can be found. Rough estimations about the altitude loss of aerosols due to sedimentation are presented in Table 7, with travelling times of $t_{min} = 60$ d to $t_{max} = 120$ d.

Table 7. Height loss of stratospheric aerosol due to sedimentation while travelling two to four months from the Asian summer monsoon region to Ny-Ålesund

	t_{min}	t_{max}
v_{min}	155 m	311 m
v_{max}	1037 m	2074 m

575 Since the fall velocities are such small for these particles, it is probable that they remain long enough in the stratosphere to be transported to the European Arctic. Additionally, due to the slow sedimentation velocity as well as their small size (see Section 3.4), aerosol from the monsoon region can reach Ny-Ålesund.

According to CLaMS back-trajectory calculations for the two exemplary days (13 and 28 September 2021) air parcels from
580 surface sources in Asia have transport times from about two to four months until they reach the lower stratosphere over Ny-Ålesund. Therefore, enhanced stratospheric backscatter coefficients measured from April to May 2021 are not caused by the Asian summer monsoon that usually starts at the beginning of June. Hence, the monsoon contributes to the aerosol load but is overlaid by other aerosol sources and the beginning of the increased backscatter coefficient is based on other contributions not only by the monsoon. The Lidar data of 28 September in comparison with CLaMS agree very well that the monsoon signal
585 can be clearly found in the Arctic as filaments by a noticeable increased backscatter ratio of about 15%.
Furthermore, the spreading between more and less polluted days is between August to October at the same level as in the other months (Figures 10 and 11). Therefore, we conclude that the increased backscatter coefficient in summer is caused by advection of aerosol from the Asian summer monsoon region and the advected particles from the ATAL. We clearly captured one event where a filament of “monsoon air” increased the particle backscatter by about 15% at 380 K to 400 K (28 September,
590 Figure 16a). The increased backscatter also happens about two months shown in Figures 10 and 11 after the monsoon in South Asia started. As it can be seen in Figure 15 it is possible for air parcels to travel within these two months from South Asia to Ny-Ålesund. Therefore we conclude, we can observe the signal of the Asian summer monsoon in the Arctic stratosphere.

Aerosol model simulations by Yu et al. (2017) proposed that aerosol particles from the ATAL can spread throughout the
595 entire extra-tropical northern lower stratosphere and contribute significantly ($\approx 15\%$) to the Northern hemisphere stratospheric column aerosol surface area on an annual basis. Eastward air mass transport of aerosol particles from the ATAL was already detected over Japan using Lidar measurements (Fujiwara et al., 2021). Khaykin et al. (2017) used long-term Lidar and satellite data over France and concluded that the ATAL is the main source for the increase of stratospheric aerosol in years without major volcanic eruptions after the maximum of aerosol load in summer. Our findings, show that the ATAL during summer and
600 autumn even has an impact on the Arctic lower stratosphere.

Stratospheric aerosol pollution from the lower latitudes has already been observed over Antarctica as well. In the study of Jumelet et al. (2020) aerosols by Australian wild fires reached the stratosphere by strong convection in Pyro-Cumulonimbus (PyroCb). They were traced and measured by the satellite-based Lidar on CALIOP as well as by the Lidar operated at the French Antarctic station Dumont d’Urville and cycled around the pole within about 6 weeks following the dominant wind
605 fields. In slightly higher altitudes (400 K to 450 K altitude) smoke layers in the shape of filaments were detected over Antarctica, which passed by within days. The observations of Jumelet et al. (2020) perfectly agree with the observations by KARL and the model output by CLaMS.

5 Conclusions and Outlook

610 In this work an annual record of Lidar observations at AWIPEV in Ny-Ålesund has been presented which is free from obvious layers like polar stratospheric clouds (PSC), volcanic eruptions or biomass burnings. The Lidar measurements have been linked to Lagrangian transport simulations with the CLaMS model using artificial surface origin tracers as well as back-trajectory calculations. The main findings of this work can be summarised:

- 615 – The low stratosphere reveals an annual cycle with lower backscatter in winter until early spring and higher values in summer to autumn
- Between 380 K and 400 K altitude the backscatter coefficient is by 25% larger in summer compared to autumn for 532 nm. The annual cycle is even slightly more pronounced for 355 nm
- 620 – The increased backscatter coefficient in June to early August can be explained by the Asian monsoon due to the time, the air is advected. The observed increasing backscatter coefficient during summer and autumn correlates with the advection time and the beginning of the Asian summer monsoon. Furthermore, the depolarisation shows a maximum during summer, even though it is generally low in our data set. Hence, probably biomass burning events contribute to stratospheric background aerosol even if no clear distinct layer can be seen by the Lidar
- 625 – Stratospheric aerosol generally shows a very low depolarisation, indicating almost spherical particles. Similar to the backscatter we found an annual cycle with slightly larger depolarisation in summer. The measured depolarisation as well as the back-scatter coefficient fit very well to the physical properties of the ATAL (Vernier et al., 2011).
- 630 – Lagrangian transport simulation with CLaMS show that air masses found between 360 K to 380 K air over Ny-Ålesund were transported from surface sources in Asia into the Arctic lower stratosphere. Case studies using back-trajectory calculations for two different days are presented. Our findings show, that the increased backscatter coefficient during summer can be linked to transport of aerosol particles from the Asian Tropopause Aerosol Layer (ATAL) into the Arctic lower stratosphere. Maximum backscatter coefficients were found during the peak season of the Asian monsoon in August. We demonstrate that the ATAL and thus the Asian monsoon has an impact on enhanced stratospheric aerosol backscatter coefficients found from June to October in the Arctic
- 635 – When assuming a mixed aerosol composition, which is typical of the ATAL, we chose optical properties of sulfate and biomass burning aerosol to calculate the aerosol effective radii using Mie calculation. In the first half of the year we obtained radii of 80 nm to 100 nm and 60 nm to 70 nm in the second half of the year

Our findings show that transport of aerosol particles from ATAL have an impact on the Arctic lower stratosphere. In future studies regional radiative forcing due to the ATAL can be compared with the global aerosol forcing caused by moderate volcanic eruptions since 2000 (Yu et al., 2017). We argue that further increasing industrial emissions in Asia will lead to a wider and thicker ATAL. An enhanced ATAL most likely yield increased transport of aerosol particles to the extra-tropical northern

640 stratosphere that could enhance the climate impact of aerosol particles in this region in the future.

Since stratospheric aerosol has a very important impact on the climate, further investigations of the natural aerosol load are important. It would increase our knowledge of its contribution, its sources and ageing processes by adding more observation sites worldwide to the study. Stations at remote places and high altitudes would be preferred to minimise the tropospheric
645 influence to the signal. Especially the upcoming EarthCARE satellite may help to track faint stratospheric aerosol layers from the Asian monsoon region also over the Southern hemisphere. Furthermore, in-situ measurements by plane and balloons at different locations and times within the lower stratosphere would also quantify the properties of aerosol better.

Data availability. The data for the tropopause heights for 2021 can be found on the data repository Jülich Data including further information
650 (Hoffmann and Spang, 2021, 2022). The radiosonde data is available at the PANGAEA repository (Maturilli, 2020). MODIS data about biomass burnings was taken from EarthData Repository. The Lidar evaluation software is written in MATLAB and can be obtained from the authors

Author contributions. Lidar, MODIS, Radiosonde and ERA5 tropopause data have been evaluated by S.G. C.R. participated in Lidar evaluation. The CLaMS simulations and analysis were performed by I.T. and B.V. All authors participated in the conceptualisation. The draft has
655 mainly been written by S.G. All authors have read and agreed to the published version of the manuscript

Competing interests. The authors declare no conflict of interest

Acknowledgements. The Lidar KARL was serviced and operated by several Observatory Engineers and impres GmbH at AWIPEV, Ny-Ålesund. We also thank Roland Neuber and Marion Maturilli as scientific coordinators of AWIPEV, who supported the Lidar project over all the years. The presented work includes contributions to the NSFC–DFG 2020 project ATAL-track (VO 1276/6-1). The research based on
660 KARL data did not receive any external funding

References

- Andersson, S. M., Martinsson, B. G., Vernier, J.-P., Friberg, J., Brenninkmeijer, C. A., Hermann, M., Van Velthoven, P. F., and Zahn, A.: Significant radiative impact of volcanic aerosol in the lowermost stratosphere, *Nature communications*, 6, 7692, <https://doi.org/10.1038/ncomms8692>, 2015.
- 665 Ansmann, A., Wandinger, U., Riebesell, M., Weitkamp, C., and Michaelis, W.: Independent measurement of extinction and backscatter profiles in cirrus clouds by using a combined Raman elastic-backscatter lidar, *Applied optics*, 31, 7113–7131, <https://doi.org/10.1364/AO.31.007113>, 1992.
- Ansmann, A., Baars, H., Chudnovsky, A., Mattis, I., Veselovskii, I., Haarig, M., Seifert, P., Engelmann, R., and Wandinger, U.: Extreme levels of Canadian wildfire smoke in the stratosphere over central Europe on 21–22 August 2017, *Atmospheric Chemistry and Physics*, 670 18, 11 831–11 845, <https://doi.org/10.34657/10755>, 2018.
- Appel, O., Köllner, F., Dragoneas, A., Hünig, A., Molleker, S., Schlager, H., Mahnke, C., Weigel, R., Port, M., Schulz, C., et al.: Chemical analysis of the Asian tropopause aerosol layer (ATAL) with emphasis on secondary aerosol particles using aircraft-based in situ aerosol mass spectrometry, *Atmospheric Chemistry and Physics*, 22, 13 607–13 630, <https://doi.org/10.5194/acp-22-13607-2022>, 2022.
- Baars, H., Ansmann, A., Ohneiser, K., Haarig, M., Engelmann, R., Althausen, D., Hanssen, I., Gausa, M., Pietruczuk, A., Szkop, A., et al.: 675 The unprecedented 2017–2018 stratospheric smoke event: decay phase and aerosol properties observed with the EARLINET, *Atmospheric chemistry and physics*, 19, 15 183–15 198, <https://doi.org/10.5194/acp-19-15183-2019>, 2019.
- Beyer, K. D. and Ebeling, D. D.: UV refractive indices of aqueous ammonium sulfate solutions, *Geophysical research letters*, 25, 3147–3150, <https://doi.org/10.1029/98GL02333>, 1998.
- Bian, J., Li, D., Bai, Z., Li, Q., Lyu, D., and Zhou, X.: Transport of Asian surface pollutants to the global stratosphere from the Tibetan 680 Plateau region during the Asian summer monsoon, *National science review*, 7, 516–533, <https://doi.org/10.1093/nsr/nwaa005>, 2020.
- Bohren, C., Huffman, D. R., and Clothiaux, E. E.: *Absorption and Scattering of Light by small particles*, Wiley-VCH, 2023.
- Brunamonti, S., Jorge, T., Oelsner, P., Hanumanthu, S., Singh, B. B., Kumar, K. R., Sonbawne, S., Meier, S., Singh, D., Wienhold, F. G., Luo, B. P., Boettcher, M., Poltera, Y., Jauhiainen, H., Kayastha, R., Karmacharya, J., Dirksen, R., Naja, M., Rex, M., Fadnavis, S., and Peter, T.: 685 Balloon-borne measurements of temperature, water vapor, ozone and aerosol backscatter on the southern slopes of the Himalayas during StratoClim 2016–2017, *acp*, 18, 15 937–15 957, <https://doi.org/10.5194/acp-18-15937-2018>, 2018.
- Butchart, N.: The Brewer-Dobson circulation, *Reviews of geophysics*, 52, 157–184, <https://doi.org/10.1002/2013RG000448>, 2014.
- Cheng, W., MacMartin, D. G., Kravitz, B., Visoni, D., Bednarz, E. M., Xu, Y., Luo, Y., Huang, L., Hu, Y., Staten, P. W., et al.: Changes in Hadley circulation and intertropical convergence zone under strategic stratospheric aerosol geoengineering, *npj Climate and Atmospheric Science*, 5, 32, <https://doi.org/10.1038/s41612-022-00254-6>, 2022.
- 690 Cheremisin, A., Marichev, V., Bochkovskii, D., and et al.: Stratospheric Aerosol of Siberian Forest Fires According to Lidar Observations in Tomsk in August 2019., *Atmos Ocean Opt*, 35, 57–64, <https://doi.org/https://doi.org/10.1134/S1024856022010043>, 2022.
- Clemens, J., Vogel, B., Hoffmann, L., Griessbach, S., Thomas, N., Fadnavis, S., Müller, R., Peter, T., and Ploeger, F.: A multi-scenario Lagrangian trajectory analysis to identify source regions of the Asian tropopause aerosol layer on the Indian subcontinent in August 2016, *Atmospheric Chemistry and Physics*, 24, 763–787, <https://doi.org/10.5194/acp-24-763-2024>, 2024.
- 695 Dahlkötter, F., Gysel, M., Sauer, D., Minikin, A., Baumann, R., Seifert, P., Ansmann, A., Fromm, M., Voigt, C., and Weinzierl, B.: The Pagami Creek smoke plume after long-range transport to the upper troposphere over Europe–aerosol properties and black carbon mixing state, *Atmospheric Chemistry and Physics*, 14, 6111–6137, <https://doi.org/10.5194/acp-14-6111-2014>, 2014.

- Emde, C., Buras-Schnell, R., Kylling, A., Mayer, B., Gasteiger, J., Hamann, U., Kylling, J., Richter, B., Pause, C., Dowling, T., et al.: The libRadtran software package for radiative transfer calculations (version 2.0. 1), *Geoscientific Model Development*, 9, 1647–1672, <https://doi.org/10.5194/gmd-9-1647-2016>, 2016.
- 700 Fadnavis, S., Semeniuk, K., Pozzoli, L., Schultz, M. G., Ghude, S. D., Das, S., and Kakatkar, R.: Transport of aerosols into the UTLS and their impact on the Asian monsoon region as seen in a global model simulation, *acp*, 13, 8771–8786, <https://doi.org/10.5194/acp-13-8771-2013>, 2013.
- Fadnavis, S., Ravi Kumar, K., Tiwari, Y. K., and Pozzoli, L.: Atmospheric CO₂ source and sink patterns over the Indian region, *Annales Geophysicae*, 34, 279–291, <https://doi.org/10.5194/angeo-34-279-2016>, 2016.
- 705 Foken, T.: Springer handbook of atmospheric measurements, Springer, 2021.
- Friend, J. P., Leifer, R., and Trichon, M.: On the formation of stratospheric aerosols, *Journal of Atmospheric Sciences*, 30, 465–479, [https://doi.org/10.1175/1520-0469\(1973\)030<0465:OTFOSA>2.0.CO;2](https://doi.org/10.1175/1520-0469(1973)030<0465:OTFOSA>2.0.CO;2), 1973.
- Fujiwara, M., Sakai, T., Nagai, T., Shiraiishi, K., Inai, Y., Khaykin, S., Xi, H., Shibata, T., Shiotani, M., and Pan, L. L.: Lower-stratospheric aerosol measurements in eastward-shedding vortices over Japan from the Asian summer monsoon anticyclone during the summer of 2018, *Atmospheric Chemistry and Physics*, 21, 3073–3090, <https://doi.org/10.5194/acp-21-3073-2021>, 2021.
- 710 Graßl, S., Ritter, C., and Schulz, A.: The Nature of the Ny-Ålesund Wind Field Analysed by High-Resolution Windlidar Data, *Remote Sensing*, 14, 3771, <https://doi.org/10.3390/rs14153771>, 2022.
- Haarig, M., Ansmann, A., Baars, H., Jimenez, C., Veselovskii, I., Engelmann, R., and Althausen, D.: Extreme levels of Canadian wildfire smoke in the stratosphere over central Europe–Part 2: Lidar study of depolarization and lidar ratios at 355, 532, and 1064 nm and of microphysical properties, *Atmospheric Chemistry and Physics*, <https://doi.org/10.5194/acp-18-11847-2018>, 2018.
- 715 Hanumanthu, S., Vogel, B., Müller, R., Brunamonti, S., Fadnavis, S., Li, D., Ölsner, P., Naja, M., Singh, B. B., Kumar, K. R., et al.: Strong day-to-day variability of the Asian Tropopause Aerosol Layer (ATAL) in August 2016 at the Himalayan foothills, *Atmospheric chemistry and physics*, 20, 14 273–14 302, <https://doi.org/10.5194/acp-20-14273-2020>, 2020.
- 720 Hersbach, H., Bell, B., Berrisford, P., Hirahara, S., Horányi, A., Muñoz-Sabater, J., Nicolas, J., Peubey, C., Radu, R., Schepers, D., Simmons, A., Soci, C., Abdalla, S., Abellan, X., Balsamo, G., Bechtold, P., Biavati, G., Bidlot, J., Bonavita, M., De Chiara, G., Dahlgren, P., Dee, D., Diamantakis, M., Dragani, R., Flemming, J., Forbes, R., Fuentes, M., Geer, A., Haimberger, L., Healy, S., Hogan, R. J., Hólm, E., Janisková, M., Keeley, S., Laloyaux, P., Lopez, P., Lupu, C., Radnoti, G., de Rosnay, P., Rozum, I., Vamborg, F., Villaume, S., and Thépaut, J.-N.: The ERA5 global reanalysis, *Quarterly Journal of the Royal Meteorological Society*, 146, 1999–2049, <https://doi.org/https://doi.org/10.1002/qj.3803>, 2020.
- 725 Hoffmann, A.: Comparative aerosol studies based on multi-wavelength Raman LIDAR at Ny-Ålesund, Spitsbergen, Ph.D. thesis, Universität Potsdam, 2011.
- Hoffmann, L. and Spang, R.: Reanalysis Tropopause Data Repository, <https://doi.org/10.26165/JUELICH-DATA/UBNGI2>, 2021.
- Hoffmann, L. and Spang, R.: An assessment of tropopause characteristics of the ERA5 and ERA-Interim meteorological reanalyses, *Atmospheric Chemistry and Physics*, 22, 4019–4046, <https://doi.org/10.5194/acp-22-4019-2022>, 2022.
- 730 Hofmann, D., Rosen, J., Pepin, T., and Pinnick, R.: Stratospheric aerosol measurements. I- Time variations at northern midlatitudes, *Journal of the Atmospheric Sciences*, 32, 1446–1456, [https://doi.org/10.1175/1520-0469\(1975\)032<1446:SAMITV>2.0.CO;2](https://doi.org/10.1175/1520-0469(1975)032<1446:SAMITV>2.0.CO;2), 1975.
- Holton, J. R., Haynes, P. H., McIntyre, M. E., Douglass, A. R., Rood, R. B., and Pfister, L.: Stratosphere-troposphere exchange, *Reviews of geophysics*, 33, 403–439, <https://doi.org/10.1029/95RG02097>, 1995.

- 735 Irvine, P. J. and Keith, D. W.: Halving warming with stratospheric aerosol geoengineering moderates policy-relevant climate hazards, *Environmental Research Letters*, 15, 044 011, <https://doi.org/10.1088/1748-9326/ab76de>, 2020.
- Jumelet, J., Klekociuk, A., Alexander, S., Bekki, S., Hauchecorne, A., Vernier, J.-P., Fromm, M., and Keckhut, P.: Detection of aerosols in Antarctica from long-range transport of the 2009 Australian wildfires, *Journal of Geophysical Research: Atmospheres*, 125, e2020JD032 542, <https://doi.org/10.1029/2020JD032542>, 2020.
- 740 Junge, C. E., Chagnon, C. W., and Manson, J. E.: Stratospheric aerosols, *Journal of Atmospheric Sciences*, 18, 81–108, [https://doi.org/10.1175/1520-0469\(1961\)018<0081:SA>2.0.CO;2](https://doi.org/10.1175/1520-0469(1961)018<0081:SA>2.0.CO;2), 1961a.
- Junge, C. E., Chagnon, C. W., and Manson, J. E.: A world-wide stratospheric aerosol layer, *Science*, 133, 1478–1479, <https://doi.org/10.1126/science.133.3463.1478.b>, 1961b.
- Khaykin, S. M., Godin-Beekmann, S., Keckhut, P., Hauchecorne, A., Jumelet, J., Vernier, J.-P., Bourassa, A., Degenstein, D. A., Rieger, L. A., Bingen, C., et al.: Variability and evolution of the midlatitude stratospheric aerosol budget from 22 years of ground-based lidar and satellite observations, *Atmospheric Chemistry and Physics*, 17, 1829–1845, <https://doi.org/10.5194/acp-17-1829-2017>, 2017.
- 745 Klett, J. D.: Stable analytical inversion solution for processing lidar returns, *Appl. Opt.*, 20, 211–220, <https://doi.org/10.1364/AO.20.000211>, 1981.
- Klett, J. D.: Lidar inversion with variable backscatter/extinction ratios, *Applied optics*, 24, 1638–1643, <https://doi.org/10.1364/AO.24.001638>, 1985.
- 750 Klimont, Z., Smith, S. J., and Cofala, J.: The last decade of global anthropogenic sulfur dioxide: 2000–2011 emissions, *Environmental Research Letters*, 8, 014 003, <https://doi.org/10.1088/1748-9326/8/1/014003>, 2013.
- Konopka, P., Tao, M., von Hobe, M., Hoffmann, L., Kloss, C., Ravegnani, F., Volk, C. M., Lauther, V., Zahn, A., Hoor, P., and Ploeger, F.: Tropospheric transport and unresolved convection: numerical experiments with CLaMS-2.0/MESSEY, *Geoscientific Model Development*, 15, 7471–7487, <https://doi.org/10.5194/gmd-15-7471-2022>, 2022.
- 755 Kremser, S., Thomason, L. W., von Hobe, M., Hermann, M., Deshler, T., Timmreck, C., Toohey, M., Stenke, A., Schwarz, J. P., Weigel, R., et al.: Stratospheric aerosol—Observations, processes, and impact on climate, *Reviews of Geophysics*, 54, 278–335, <https://doi.org/10.1002/2015RG000511>, 2016.
- Kunz, A., Sprenger, M., and Wernli, H.: Climatology of potential vorticity streamers and associated isentropic transport pathways across PV gradient barriers, *Journal of Geophysical Research: Atmospheres*, 120, 3802–3821, <https://doi.org/10.1002/2014JD022615>, 2015.
- 760 Langenbach, A., Baumgarten, G., Fiedler, J., Lübken, F.-J., von Savigny, C., and Zalach, J.: Year-round stratospheric aerosol backscatter ratios calculated from lidar measurements above northern Norway, *Atmospheric Measurement Techniques*, 12, 4065–4076, <https://doi.org/10.5194/amt-12-4065-2019>, 2019.
- Lau, W. K., Yuan, C., and Li, Z.: Origin, maintenance and variability of the Asian Tropopause Aerosol Layer (ATAL): the roles of monsoon dynamics, *Scientific Reports*, 8, 3960, <https://doi.org/10.1038/s41598-018-22267-z>, 2018.
- 765 Lauther, V., Vogel, B., Wintel, J., Rau, A., Hoor, P., Bense, V., Müller, R., and Volk, C. M.: In situ observations of CH₂Cl₂ and CHCl₃ show efficient transport pathways for very short-lived species into the lower stratosphere via the Asian and the North American summer monsoon, *Atmospheric chemistry and physics*, 22, 2049–2077, <https://doi.org/10.5194/acp-22-2049-2022>, 2022.
- Lee, W. R., MacMartin, D. G., Vioni, D., and Kravitz, B.: High-Latitude Stratospheric Aerosol Geoengineering Can Be More Effective if Injection Is Limited to Spring, *Geophysical Research Letters*, 48, e2021GL092 696, <https://doi.org/https://doi.org/10.1029/2021GL092696>, e2021GL092696 2021GL092696, 2021.
- 770

- Maturilli, M.: High resolution radiosonde measurements from station Ny-Ålesund (2017-04 et seq), <https://doi.org/10.1594/PANGAEA.914973>, 2020.
- 775 Maturilli, M. and Kayser, M.: Arctic warming, moisture increase and circulation changes observed in the Ny-Ålesund homogenized radiosonde record, *Theoretical and Applied Climatology*, 130, 1–17, <https://doi.org/10.1007/s00704-016-1864-0>, 2017.
- Mayer, B. and Kylling, A.: The libRadtran software package for radiative transfer calculations-description and examples of use, *Atmospheric Chemistry and Physics*, 5, 1855–1877, <https://doi.org/10.5194/acp-5-1855-2005>, 2005.
- McIntyre, M. E. and Palmer, T.: Breaking planetary waves in the stratosphere, *Nature*, 305, 593–600, <https://doi.org/10.1038/305593a0>, 1983.
- 780 McKenna, D. S., Groöß, J.-U., Günther, G., Konopka, P., Müller, R., Carver, G., and Sasano, Y.: A new Chemical Lagrangian Model of the Stratosphere (CLaMS) 2. Formulation of chemistry scheme and initialization, *Journal of Geophysical Research: Atmospheres*, 107, 4256, <https://doi.org/10.1029/2000JD000113>, 2002a.
- McKenna, D. S., Konopka, P., Groöß, J.-U., Günther, G., Müller, R., Spang, R., Offermann, D., and Orsolini, Y.: A new Chemical Lagrangian Model of the Stratosphere (CLaMS) - 1. Formulation of advection and mixing, *Journal of Geophysical Research: Atmospheres*, 107, 4309, <https://doi.org/10.1029/2000JD000114>, 2002b.
- 785 Mie, G.: Beiträge zur Optik trüber Medien, speziell kolloidaler Metallösungen, *Annalen der physik*, 330, 377–445, <https://doi.org/10.1002/andp.19083300302>, 1908.
- Murphy, D. M., Froyd, K. D., Bourgeois, I., Brock, C. A., Kupc, A., Peischl, J., Schill, G. P., Thompson, C. R., Williamson, C. J., and Yu, P.: Radiative and chemical implications of the size and composition of aerosol particles in the existing or modified global stratosphere, *Atmospheric Chemistry and Physics*, 21, 8915–8932, <https://doi.org/10.5194/acp-21-8915-2021>, 2021.
- 790 Nomura, S., Naja, M., Ahmed, M. K., Mukai, H., Terao, Y., Machida, T., Sasakawa, M., and Patra, P. K.: Measurement report: Regional characteristics of seasonal and long-term variations in greenhouse gases at Nainital, India, and Comilla, Bangladesh, *acp*, 21, 16427–16452, <https://doi.org/10.5194/acp-21-16427-2021>, 2021.
- Notholt, J., Luo, B., Fueglistaler, S., Weisenstein, D., Rex, M., Lawrence, M., Bingemer, H., Wohltmann, I., Corti, T., Warneke, T., et al.: Influence of tropospheric SO₂ emissions on particle formation and the stratospheric humidity, *Geophysical Research Letters*, 32, <https://doi.org/10.1029/2004GL022159>, 2005.
- 795 Ohneiser, K., Ansmann, A., Chudnovsky, A., Engelmann, R., Ritter, C., Veselovskii, I., Baars, H., Gebauer, H., Griesche, H., Radenz, M., et al.: The unexpected smoke layer in the High Arctic winter stratosphere during MOSAiC 2019–2020, *Atmospheric Chemistry and Physics*, 21, 15783–15808, <https://doi.org/10.5194/acp-21-15783-2021>, 2021.
- 800 Ohneiser, K., Ansmann, A., Witthuhn, J., Deneke, H., Chudnovsky, A., Walter, G., and Senf, F.: Self-lofting of wildfire smoke in the troposphere and stratosphere: simulations and space lidar observations, *Atmospheric Chemistry and Physics*, 23, 2901–2925, <https://doi.org/10.5194/acp-23-2901-2023>, 2023.
- Oppenheimer, C., Francis, P., and Stix, J.: Depletion rates of sulfur dioxide in tropospheric volcanic plumes, *Geophysical Research Letters*, 25, 2671–2674, <https://doi.org/10.1029/98GL01988>, 1998.
- 805 Organization, W. M.: WMO Bulletin, vol. VI, World Meteorological Organization, 1957.
- Ploeger, F., Konopka, P., Günther, G., Groöß, J.-U., and Müller, R.: Impact of the vertical velocity scheme on modeling transport in the tropical tropopause layer, *Journal of Geophysical Research: Atmospheres*, 115, D03301, <https://doi.org/10.1029/2009JD012023>, 2010.

- Ploeger, F., Diallo, M., Charlesworth, E., Konopka, P., Legras, B., Laube, J. C., Grooß, J.-U., Günther, G., Engel, A., and Riese, M.: The stratospheric Brewer–Dobson circulation inferred from age of air in the ERA5 reanalysis, *Atmospheric Chemistry and Physics*, 21, 8393–8412, <https://doi.org/10.5194/acp-21-8393-2021>, 2021.
- Pommrich, R., Müller, R., Grooß, J.-U., Konopka, P., Ploeger, F., Vogel, B., Tao, M., Hoppe, C. M., Günther, G., Spelten, N., Hoffmann, L., Pumphrey, H.-C., Viciani, S., D’Amato, F., Volk, C. M., Hoor, P., Schlager, H., and Riese, M.: Tropical troposphere to stratosphere transport of carbon monoxide and long-lived trace species in the Chemical Lagrangian Model of the Stratosphere (CLaMS), *Geoscientific Model Development*, 7, 2895–2916, <https://doi.org/10.5194/gmd-7-2895-2014>, 2014.
- Quaglia, I., Timmreck, C., Niemeier, U., Visioni, D., Pitari, G., Brühl, C., Dhomse, S., Franke, H., Laakso, A., Mann, G., et al.: Interactive Stratospheric Aerosol models response to different amount and altitude of SO₂ injections during the 1991 Pinatubo eruption, *Atmospheric chemistry and physics discussions*, 2022, 1–35, <https://doi.org/10.5194/acp-23-921-2023>, 2022.
- Randel, W. J., Park, M., Emmons, L., Kinnison, D., Bernath, P., Walker, K. A., Boone, C., and Pumphrey, H.: Asian monsoon transport of pollution to the stratosphere, *Science*, 328, 611–613, <https://doi.org/10.1126/science.1182274>, 2010.
- Russell, P. B. and Hamill, P.: Spatial variation of stratospheric aerosol acidity and model refractive index: Implications of recent results, *Journal of Atmospheric Sciences*, 41, 1781–1790, [https://doi.org/10.1175/1520-0469\(1984\)041<1781:SVOSAA>2.0.CO;2](https://doi.org/10.1175/1520-0469(1984)041<1781:SVOSAA>2.0.CO;2), 1984.
- Shepherd, T. G.: Issues in stratosphere-troposphere coupling, *Journal of the Meteorological Society of Japan. Ser. II*, 80, 769–792, <https://doi.org/10.2151/jmsj.80.769>, 2002.
- Sumlin, B. J., Heinson, Y. W., Shetty, N., Pandey, A., Pattison, R. S., Baker, S., Hao, W. M., and Chakrabarty, R. K.: UV–Vis–IR spectral complex refractive indices and optical properties of brown carbon aerosol from biomass burning, *Journal of Quantitative Spectroscopy and Radiative Transfer*, 206, 392–398, <https://doi.org/10.1016/j.jqsrt.2017.12.009>, 2018.
- Sun, W., Wang, B., Chen, D., Gao, C., Lu, G., and Liu, J.: Global monsoon response to tropical and Arctic stratospheric aerosol injection, *climate Dynamics*, 55, 2107–2121, <https://doi.org/10.1007/s00382-020-05371-7>, 2020.
- Thomason, L. W. and Knepp, T.: Quantifying SAGE II (1984–2005) and SAGE III/ISS (2017–2022) observations of smoke in the stratosphere, *Atmospheric Chemistry and Physics*, 23, 10361–10381, <https://doi.org/10.5194/acp-23-10361-2023>, 2023.
- Tritscher, I., Pitts, M. C., Poole, L. R., Alexander, S. P., Cairo, F., Chipperfield, M. P., Grooß, J.-U., Höpfner, M., Lambert, A., Luo, B., et al.: Polar stratospheric clouds: Satellite observations, processes, and role in ozone depletion, *Reviews of geophysics*, 59, e2020RG000702, <https://doi.org/10.1029/2020RG000702>, 2021.
- Turco, R., Whitten, R., and Toon, O.: Stratospheric aerosols: Observation and theory, *Reviews of Geophysics*, 20, 233–279, <https://doi.org/10.1029/RG020i002p00233>, 1982.
- Vernier, J.-P., Thomason, L., and Kar, J.: CALIPSO detection of an Asian tropopause aerosol layer, *Geophysical Research Letters*, 38, <https://doi.org/10.1029/2010GL046614>, 2011.
- Vogel, B., Günther, G., Müller, R., Grooß, J.-U., Hoor, P., Krämer, M., Müller, S., Zahn, A., and Riese, M.: Fast transport from Southeast Asia boundary layer sources to northern Europe: rapid uplift in typhoons and eastward eddy shedding of the Asian monsoon anticyclone, *Atmospheric Chemistry and Physics*, 14, 12745–12762, <https://doi.org/10.5194/acp-14-12745-2014>, 2014.
- Vogel, B., Günther, G., Müller, R., Grooß, J.-U., and Riese, M.: Impact of different Asian source regions on the composition of the Asian monsoon anticyclone and of the extratropical lowermost stratosphere, *Atmospheric chemistry and physics*, 15, 13699–13716, <https://doi.org/10.5194/acp-15-13699-2015>, 2015.

- Vogel, B., Günther, G., Müller, R., Groß, J.-U., Afchine, A., Bozem, H., Hoor, P., Krämer, M., Müller, S., Riese, M., et al.: Long-range
845 transport pathways of tropospheric source gases originating in Asia into the northern lower stratosphere during the Asian monsoon season
2012, *Atmospheric Chemistry and Physics*, 16, 15 301–15 325, <https://doi.org/10.5194/acp-16-15301-2016>, 2016.
- Vogel, B., Müller, R., Günther, G., Spang, R., Hanumanthu, S., Li, D., Riese, M., and Stiller, G. P.: Lagrangian simulations of the transport
of young air masses to the top of the Asian monsoon anticyclone and into the tropical pipe, *Atmospheric Chemistry and Physics*, 19,
6007–6034, <https://doi.org/10.5194/acp-19-6007-2019>, 2019.
- 850 Vogel, B., Volk, C. M., Wintel, J., Lauther, V., Clemens, J., Groß, J.-U., Günther, G., Hoffmann, L., Laube, J. C., Müller, R., et al.: Evaluation
of vertical transport in ERA5 and ERA-Interim reanalysis using high-altitude aircraft measurements in the Asian summer monsoon 2017,
Atmospheric Chemistry and Physics, 24, 317–343, <https://doi.org/10.5194/acp-24-317-2024>, 2024.
- Wang, J.-J.: Evolution and structure of the mesoscale convection and its environment: A case study during the early
onset of the Southeast Asian summer monsoon, *Monthly weather review*, 132, 1104–1120, [https://doi.org/10.1175/1520-0493\(2004\)132<1104:EASOTM>2.0.CO;2](https://doi.org/10.1175/1520-0493(2004)132<1104:EASOTM>2.0.CO;2), 2004.
- 855 Washenfelder, R., Flores, J., Brock, C., Brown, S., and Rudich, Y.: Broadband measurements of aerosol extinction in the ultraviolet spectral
region, *Atmospheric Measurement Techniques*, 6, 861–877, <https://doi.org/10.5194/amt-6-861-2013>, 2013.
- Waugh, D. and Hall, T.: Age of stratospheric air: Theory, observations, and models, *Reviews of Geophysics*, 40, 1–1,
<https://doi.org/10.1029/2000RG000101>, 2002.
- 860 Waugh, D. W.: Seasonal variation of isentropic transport out of the tropical stratosphere, *Journal of Geophysical Research: Atmospheres*,
101, 4007–4023, <https://doi.org/10.1029/95JD03160>, 1996.
- Weigel, R., Mahnke, C., Baumgartner, M., Dragoneas, A., Vogel, B., Ploeger, F., Viciani, S., D’Amato, F., Bucci, S., Legras, B., Luo, B., and
Borrmann, S.: In situ observation of new particle formation (NPF) in the tropical tropopause layer of the 2017 Asian monsoon anticyclone
– Part 1: Summary of StratoClim results, *acp*, 21, 11 689–11 722, <https://doi.org/10.5194/acp-21-11689-2021>, 2021a.
- 865 Weigel, R., Mahnke, C., Baumgartner, M., Krämer, M., Spichtinger, P., Spelten, N., Afchine, A., Rolf, C., Viciani, S., D’Amato, F., Tost,
H., and Borrmann, S.: In situ observation of new particle formation (NPF) in the tropical tropopause layer of the 2017 Asian monsoon
anticyclone – Part 2: NPF inside ice clouds, *acp*, 21, 13 455–13 481, <https://doi.org/10.5194/acp-21-13455-2021>, 2021b.
- Wetzel, G., Friedl-Vallon, F., Glatthor, N., Groß, J.-U., Gulde, T., Höpfner, M., Johansson, S., Khosrawi, F., Kirner, O., Kleinert, A., et al.:
Pollution trace gases C₂H₆, C₂H₂, HCOOH, and PAN in the North Atlantic UTLS: observations and simulations, *Atmospheric chemistry
870 and physics*, 21, 8213–8232, <https://doi.org/10.5194/acp-21-8213-2021>, 2021.
- Yan, X., Konopka, P., Ploeger, F., and Podglajen, A.: Transport into the polar stratosphere from the Asian monsoon region, *EGU sphere*, 2024,
1–22, <https://doi.org/10.5194/egusphere-2024-782>, 2024.
- Yu, P., Rosenlof, K. H., Liu, S., Telg, H., Thornberry, T. D., Rollins, A. W., Portmann, R. W., Bai, Z., Ray, E. A., Duan, Y., et al.: Efficient
transport of tropospheric aerosol into the stratosphere via the Asian summer monsoon anticyclone, *Proceedings of the National Academy
875 of Sciences*, 114, 6972–6977, <https://doi.org/10.1073/pnas.1701170114>, 2017.
- Yue, G., Poole, L., Wang, P.-H., and Chiou, E.: Stratospheric aerosol acidity, density, and refractive index deduced from SAGE II and NMC
temperature data, *Journal of Geophysical Research: Atmospheres*, 99, 3727–3738, <https://doi.org/10.1029/93JD02989>, 1994.
- Zielinski, T., Bolzacchini, E., Cataldi, M., Ferrero, L., Graßl, S., Hansen, G., Mateos, D., Mazzola, M., Neuber, R., Pakszys, P.,
Posyniak, M., Ritter, C., Severi, M., Sobolewski, P., Traversi, R., and Velasco-Merino, C.: Study of Chemical and Optical Prop-
880 erties of Biomass Burning Aerosols during Long-Range Transport Events toward the Arctic in Summer 2017, *Atmosphere*, 11,
<https://doi.org/10.3390/atmos11010084>, 2020.