



# Retrieval and evaluation of tropospheric-aerosol extinction profiles using multi-axis differential optical absorption spectroscopy (MAX-DOAS) measurements over Athens, Greece

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**Abstract.** In this study, we report on the retrieval of aerosol extinction profiles from ground-based scattered sunlight multi-axis differential optical absorption spectroscopy (MAX-DOAS) measurements, carried out at Athens, Greece. It is the first time that aerosol profiles are retrieved from MAX-DOAS measurements in Athens. The reported aerosol vertical distributions at 477 nm are derived from the oxygen dimer (O<sub>4</sub>) differential-slant-column-density observations at different elevation angles by applying the Bremen Optimal estimation RETrieval for Aerosol and trace gasS (BOREAS) retrieval algorithm. Four case studies have been selected for validation purposes; the retrieved aerosol profiles and the corresponding aerosol optical depths (AODs) from the MAX-DOAS are compared with lidar extinction profiles and with sun-photometric measurements (Aerosol Robotic Network, AERONET, observations), respectively. Despite the different approach of each method regarding the retrieval of the aerosol information, the comparison with the lidar measurements at 532 nm reveals a very good agreement in terms of vertical distribution, with  $r > 0.90$  in all cases. The AODs from the MAX-DOAS and the sun pho-

tometer (the latter at 500 nm) show a satisfactory correlation (with  $0.45 < r < 0.7$  in three out of the four cases). The comparison indicates that the MAX-DOAS systematically underestimates the AOD in the cases of large particles (small Ångström exponent) and for measurements at small relative azimuthal angles between the viewing direction and the sun. Better agreement is achieved in the morning, at large relative azimuthal angles. Overall, the aerosol profiles retrieved from MAX-DOAS measurements are of good quality; thus, new perspectives are opened up for assessing urban aerosol pollution on a long-term basis in Athens from continuous and uninterrupted MAX-DOAS measurements.

## 1 Introduction

Tropospheric aerosols originate from both natural and anthropogenic sources. The lifetime of aerosols in the troposphere ranges from a few days to a few weeks, depending on their size and meteorology (e.g. Pandis et al., 1995). They

take part in atmospheric processes through (i) nucleation and interaction with clouds (e.g. Twomey et al., 1977; Rosenfeld et al., 2014); (ii) participation in chemical and photochemical reactions by providing the required surface for heterogeneous reactions to take place (Andreae and Crutzen, 1997); and (iii) absorption and scattering of incoming solar and earth's IR radiation, affecting atmospheric dynamics and stability (e.g. Dubovik et al., 2002) and the earth's climate (IPCC, 2001). Significant decrease in UV-vis irradiance reaching the ground due to urban aerosol pollution has been reported in various cases (e.g. Zerefos et al., 2009; Chubarova et al., 2011).

According to a survey conducted in 25 large European cities, Athens occupies the third position on a European level in exceedances of particle pollution regulations (Pascal et al., 2013). Saharan dust transported from the African continent is the main natural source of tropospheric aerosols in Athens (e.g. Kanakidou et al., 2007; Gerasopoulos et al., 2011; Raptis et al., 2020), while common anthropogenic sources are traffic emission and domestic heating (Markakis et al., 2010; Gratsea et al., 2017). Wildfires also contribute to the aerosol mixture in the area occasionally, either from local events (Amiridis et al., 2012) or by long-range transport (Papayannis et al., 2009, Amiridis et al., 2011; Mona et al., 2012). Whereas emissions of most air pollutants, such as SO<sub>2</sub>, are expected to decrease by more than 80 % by the end of the 21st century, the decrease in aerosol emissions is projected to be small (IPCC, 2007), and thus aerosols may play an even more critical role in air quality in the future. Therefore, long-term continuous measurements, providing information on the spatial and temporal distribution of aerosols, are of great importance to urban air pollution assessment and to the understanding of the aerosol contribution to earth's climate. The knowledge of the vertical distribution of aerosols is necessary for understanding the mechanisms underlying the formation and development of urban smog.

Satellite, airborne and ground-based measurements are widely used to derive aerosol vertical profiles (e.g. Papayannis et al., 2005; Schmid et al., 2006; DeCarlo et al., 2008; Solanki and Singh, 2014); satellite measurements sometimes fail to be accurate in the lower atmosphere, while airborne measurements, although accurate in the lower atmosphere, are temporally restricted. In contrast, ground-based measurements can provide both a very good record of the lower troposphere and a satisfactory temporal resolution. However, since the ground-based profile measurements are mainly relying on lidar systems (e.g. the European Aerosol Research Lidar Network – EARLINET – within the European Research Infrastructure for the observation of Aerosol, Clouds and Trace Gases Research Infrastructure – ACTRIS), they are costly in terms of set-up and operation. An additional option for ground-based observations is the multi-axis differential optical absorption spectroscopy (MAX-DOAS) technique, which has been gaining ground over the last years (e.g. Wittrock et al., 2004; Heckel et al., 2005; Ma et al., 2013;

Schreier et al., 2020) since it can provide low-cost, continuous and uninterrupted measurements without the need for absolute radiometric calibration. The MAX-DOAS technique has also been shown to be very promising for the retrieval of aerosols' vertical distribution (e.g. Sinreich et al., 2005; Lee et al., 2009; Cl  mer et al., 2010; Wagner et al., 2011). However, its sensitivity at higher altitudes is low, and compared to the lidar technique, it provides profiles with much coarser vertical resolution. It also performs only daylight measurements, which can be considered to be a limitation of this technique. In some studies, the retrieved aerosol extinction profiles from MAX-DOAS measurements are compared to the corresponding profiles derived from lidar (e.g. Irie et al., 2008; Zieger et al., 2011; B  sch et al., 2018) or Aerosol Robotic Network (AERONET)-based measurements (e.g. Wang et al., 2016). For the Athens area, although several studies have been published on aerosol extinction profiles from lidar measurements (e.g. Papayannis et al., 1998, 2005; Matthias et al., 2004), vertical trace gas and aerosol profile retrievals from MAX-DOAS have not been published so far.

In the scope of this paper, a retrieval algorithm, recently developed by the Institute of Environmental Physics and Remote Sensing of University of Bremen (B  sch et al., 2018), is employed in order to obtain vertical distributions of aerosol extinction from O<sub>4</sub> MAX-DOAS measurements over the urban environment of Athens. O<sub>4</sub> is an atmospheric absorber with a known concentration profile; therefore measurements of the O<sub>4</sub> column can be used to retrieve the aerosol-induced light path changes (Wagner et al., 2004).

For validation purposes the outcomes of our calculations are compared to established techniques; the retrieved profiles are compared to profiles from ground-based lidar measurements (EARLINET station) and the aerosol optical depth (AOD) to sun photometer measurements (AERONET station).

A description of the instruments used in this study (location, instrumentation and data retrieval) along with a brief description of the profile retrieval algorithm is given in Sect. 2. In Sect. 3, we present the derived aerosol vertical distributions for four selected case studies, and we compare the MAX-DOAS aerosol extinction coefficient profiles and the AOD with lidar and sun-photometric measurements, respectively. The findings are summarised in Sect. 4, where the conclusions of this study are also provided.

## 2 Methodology

### 2.1 Location

Four mountains surround the city of Athens, forming a basin that is open to the south and south-west. This special topography plays an essential role in the accumulation of atmospheric pollutants over the city under certain meteorological

conditions (Kassomenos et al., 1995). Moreover, dust transport episodes from North Africa also contribute to the aerosol load of the city (e.g. Gerasopoulos et al., 2009; Kosmopoulos et al., 2017). In general, the Athens area can be considered to be an example of various aerosol types such as dust, local pollution, marine, biomass combustion and their mixtures (Souppiona et al., 2019).

Figure 1 shows the greater Athens area and the location of each instrument used in this study. The MAX-DOAS instrument is located at the premises of the National Observatory of Athens (NOA; 38.05° N, 23.86° E; 527 m a.s.l.) to the north of the city. No strong emission sources are present around the measurement area, which is considered to be suburban background. The lidar system performs measurements at the National and Technical University of Athens (NTUA; 37.97° N, 23.79° E; 212 m a.s.l.), and the site is considered to be suburban background. The CIMEL sun photometer is installed at the premises of NOA at Thissio hill (37°58' N, 23°43' E; 150 m a.s.l.), which, despite being located in the city centre, is considered to be urban background (Paraskevopoulou et al., 2015). Information about the instruments is provided in Table 1.

## 2.2 Instrumentation and data retrieval

### 2.2.1 MAX-DOAS

The MAX-DOAS instrument employed in this study is part of the BREDOM network (Bremen DOAS Network for Atmospheric Measurements; [http://www.iup.uni-bremen.de/DOAS/groundbased\\_data.htm](http://www.iup.uni-bremen.de/DOAS/groundbased_data.htm), last access: 14 February 2020) and has been operating continuously since October 2012. It comprises a grating spectrometer (LOT 260S, 600 L mm<sup>-1</sup> ruled grating) connected via an optical-fibre bundle to a computer-controlled telescope unit. The spectrometer covers a spectral range from 330 to 500 nm with a spectral resolution of approximately 0.7 nm. The detector used is a CCD (charge-coupled device) by Andor Technology, with 2048 pixel × 512 pixel resolution, cooled to -40 °C.

The telescope performs intensity measurements at eight elevation angles (-1, 0, 1, 2, 4, 8, 15, 30°) as well as to the zenith. However, the current retrieval algorithm only considers upward viewing directions, excluding -1 and 0°. With this choice, little information is available for the profile retrieval below the station altitude; therefore profiles are retrieved and presented only for altitudes above 500 m a.s.l. Measurements in eight azimuthal directions are performed, but in this study, only the *S* direction – pointing at 52.5° (with respect to south) and associated with the urban atmospheric conditions of the city (Gratsea et al., 2016) – is considered (Fig. 1). The *S* direction also covers the sun photometer's location and points close to the lidar's measurement site. The duration of one full scanning cycle (azimuthal and elevation scanning) is about 15 min; thus about 30 measurement cycles per day are available in winter and 45 in summer.

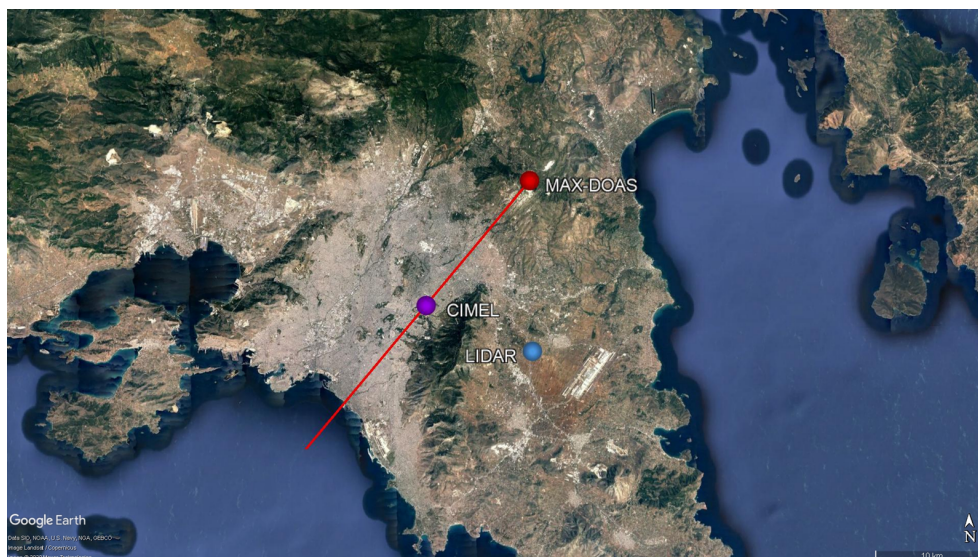
The spectral measurements are analysed using the DOAS technique; the Beer–Lambert law is considered to be the solution of the radiative transfer equation (Platt and Stutz, 2008), and the absorption spectrum is separated into broad and narrow spectral features that show low- and high-frequency variations, respectively, as a function of wavelength. The narrow spectral features correspond to the unique narrow-band absorption structures of the trace gases, while the broad ones represent the attenuation of solar radiation by scattering processes in the atmosphere as well as the continuum absorption by trace gases and the instrument. For the derivation of the slant column density (SCD; defined as the concentration of the absorber integrated along the light path), a polynomial accounting for the broad spectral features and the laboratory cross sections of the retrieved species are fitted to the measured optical depth. To determine the optical depth, the logarithm of the ratio of the current horizon measurement (*I*) and the reference intensity (*I*<sub>0</sub>) is taken.

The SCD of the oxygen dimer (O<sub>4</sub>), i.e. the slant optical thickness of the absorber divided by the absorption cross section, measured at different elevations is used as input to the retrieval algorithm for the calculation of the aerosol distribution. The slant column of the O<sub>4</sub>, a weak molecular absorber with a well-known vertical profile (the O<sub>4</sub> concentration is proportional to the square of the O<sub>2</sub>), is almost linearly dependent on the average photon path lengths (Pfeilsticker et al., 1997) and thus can be used as an indicator of the presence of clouds or aerosols in the atmosphere. The SCD<sub>O<sub>4</sub></sub> is calculated by fitting to the measured optical depth the laboratory spectrum of O<sub>4</sub> (Hermans et al., 2003), NO<sub>2</sub> (Vandaele et al., 1998) and of O<sub>3</sub> (Bogumil et al., 2000) and a polynomial of degree 4 which accounts for the broad spectral features. The fitting spectral window used is 425–490 nm. In order to retrieve the tropospheric SCD<sub>O<sub>4</sub></sub>, the zenith observation, corresponding to each measurement cycle, is used as the reference measurement *I*<sub>0</sub>, cancelling in this way the Fraunhofer lines in the solar spectrum and the stratospheric contributions to the SCD.

Tropospheric vertical column densities (VCDs) of NO<sub>2</sub>, shown in Sect. 3.1, can be derived by using air mass factors (AMFs) calculated with the SCIATRAN radiative transfer model (Rozanov et al., 2000). To convert the differential tropospheric SCD to the corresponding tropospheric VCD, the differential AMF (AMF<sub>α</sub> - AMF<sub>90°</sub>) is required, namely the difference between the AMF at the same elevation *α* as the SCD measurement and the AMF at the zenith (Eq. 1).

$$\text{VCD} = \frac{\text{SCD}_\alpha - \text{SCD}_{90^\circ}}{(\text{AMF}_\alpha - \text{AMF}_{90^\circ})} \quad (1)$$

The AMF describes the weighting of the absorption as a function of the relative azimuth and the solar zenith angle (SZA) for a given atmospheric profile and at a specific wavelength.



**Figure 1.** Measurement sites and MAX-DOAS viewing direction ( $S$ ). The distances between instruments are 16 km between MAX-DOAS and the sun photometer (CIMEL) and 13 km between MAX-DOAS and the lidar. © Google Earth

**Table 1.** Instruments and data products used in the present study.

Instrument	Location	Institute	Products
MAX-DOAS	Penteli National Observatory of Athens (38.05° N, 23.86° E; 527 m a.s.l.)	BREDOM network, Institute of Environmental Physics and Remote Sensing, University of Bremen	SCD <sub>NO<sub>2</sub></sub> , VCD <sub>NO<sub>2</sub></sub> , aerosol extinction profile, AOD
EOLE lidar	Zografou (37.97° N, 23.79° E; 212 m a.s.l.)	National Technical University of Athens, Laser Remote Sensing Laboratory (NTUA-LRSU)	Aerosol backscatter profile, aerosol extinction profile, columnar AOD
CIMEL sun–sky radiometer	Thisio 2(37.96° N, 3.72° E; 150 m a.s.l.)	National Observatory of Athens, Institute for Astronomy, Astrophysics Space Application and Remote Sensing (NOA-IAASARS)	AOD, inversion data products (ssa, asymmetry factor, refractive index, phase function, size distribution)

## 2.2.2 EOLE lidar system

The six-wavelength Raman-backscatter lidar system (EOLE) has been operating in Athens since February 2000 as part of the EARLINET network (Pappalardo et al., 2014). The system is designed following the optical set-up of a typical member station (Kokkalis, 2017), meeting all the quality assurance requirements of the network. The emission unit is based on a pulsed Nd:YAG laser, emitting high-energy pulses at 355, 532 and 1064 nm with a repetition rate of 10 Hz. The optical receiver is based on a Cassegrainian telescope (600 mm focal length and a clear aperture diameter of 300 mm), directly coupled with an optical fibre to the wavelength separation unit, detecting signals at 355, 387 (N<sub>2</sub> Raman line of 355 nm), 407 (H<sub>2</sub>O Raman line of 355 nm), 532,

607 (N<sub>2</sub> Raman line of the 532 nm) and 1064 nm. For every measuring cycle 1000 lidar signal returns are stored (every  $\sim 1.66$ ). For each case presented in this study, we used hourly averaged profiles, which correspond to approximately 34 individual signal acquisitions (Kokkalis et al., 2012).

During daytime operation, the system is capable of providing aerosol backscatter profiles ( $\beta_{\text{aer}}$ ) at 355, 532 and 1064 nm, based on the standard backscatter lidar technique and employing the Klett inversion method (Klett, 1981). This technique assumes the existence of an aerosol-free region (e.g. upper troposphere) and requires an a priori assumption of the lidar ratio value (the ratio of the extinction to backscatter coefficient,  $S_{\text{aer}}$ ). A variety of studies revealed a wide range for the lidar ratios, covering values from 20

to 100 sr (Ackermann, 1998; Mattis et al., 2004; Amiridis et al., 2005; Müller et al., 2007; Papayannis et al., 2008; Groß et al., 2011; Giannakaki et al., 2015). When the elastic backscatter lidar technique is used, the assumption of a constant lidar ratio value throughout the laser sounding range becomes very critical when solving the lidar equation; in this case, the overall uncertainty, including both statistical and systematic errors, on the retrieved  $\beta_{\text{aer}}$  values, is of the order of 20 %–30 % (e.g. Rocadenbosch et al., 2010). In this study, the aerosol extinction profiles have been retrieved under the assumption of three lidar ratio values: 30, 50 and 70 (i.e.  $50 \pm 20$  sr). This range is realistic for pollution and dust cases presented herein (Groß et al., 2013), and it is also in accordance with columnar lidar ratio values (interpolated to 532 nm) obtained by AERONET for the cases of this study, which vary from  $48.8 \pm 7.5$  to  $59.9 \pm 12.1$  sr. As a result of this variability (i.e.  $50 \pm 20$  sr), the uncertainties introduced to the aerosol extinction profiles vary from 10 %–40 %; the higher uncertainties appear at the upper atmospheric layers, where the signal-to-noise ratio of the system decreases. The corresponding uncertainties of the lidar-derived AOD values due to this assumption were estimated to be up to 11 %. All the lidar profiles were obtained with the Single Calculus Chain (SCC) processing platform (D'Amico et al., 2016; Mattis et al., 2016), which is developed in the framework of EARLINET to ensure the high-quality products of the network, by implementing quality checks on both raw lidar data and final optical products.

One of the lidar's main limitations is the distance of full overlap between the laser beam and the receiver's field of view, which makes it difficult for the instrument to obtain useful and accurate aerosol-related information below that height. Wandinger and Ansmann (2002) demonstrated that when not applying overlap correction in lidar signals, the retrieved aerosol extinction coefficient may take even non-physical negative values for heights up to the full overlap. The incomplete overlap effect can be solved by using Raman measurements under nighttime conditions. In this study, only daytime measurements are used, and therefore no overlap correction is applied on the signals. The geometrical configuration of EOLE results in full overlap distance of 500–800 m a.g.l. (Kokkalis, 2017). The aerosol extinction values below the 1000 m a.s.l. height are considered to be inside the overlap region and therefore were omitted from the extinction profile comparison. Nevertheless, in order to calculate the AOD from the lidar profiles, the lowermost trustworthy value of the extinction coefficient was assumed constant (height-independent). During daytime, the upper limit of the planetary boundary layer over Athens ranges between 1500 and 2100 m a.s.l. (Kokkalis et al., 2020); thus the minimum height of lidar profiles at 1000 m a.s.l. is well within the planetary boundary layer (PBL). Our assumption of a well-mixed atmosphere below 1000 m a.s.l. – which means that a constant lidar ratio value is considered for this part of the atmosphere (Wandinger and Ansmann, 2002) – may lead to an

underestimation of the AOD at the lowest troposphere since the city is most probably an additional source of particles. This underestimation cannot be estimated because of the lidar overlap issue.

### 2.2.3 CIMEL sun photometer

The reported columnar aerosol optical properties have been retrieved by a CIMEL sun photometer (Holben et al., 1998). The instrument is part of NASA's global sun-photometric network, AERONET, and performs automatic measurements of the direct solar radiance at the common wavelengths of 340, 380, 440, 500, 675, 870, 940 and 1020 nm every 15 min and diffuse sky radiance at 440, 675, 870 and 1020 nm. These measurements are further used to provide both optical and microphysical aerosol properties in the atmospheric column (Dubovik et al., 2006). The CIMEL data used in this study are the cloud-screened and quality-assured level 2.0 data products, providing information about the columnar AOD and the Ångström exponent. The AOD uncertainty is  $< \pm 0.02$  for UV wavelengths and  $< \pm 0.01$  for wavelengths larger than 440 nm (Eck et al., 1999).

### 2.3 BOREAS profile retrieval algorithm

The BRemen Optimal estimation REtrieval for Aerosol and trace gasS (BOREAS) is an optimal estimation-based profile retrieval algorithm developed at the Institute of Environmental Physics, University of Bremen (Bösch et al., 2018). It applies the optimal estimation technique for the retrieval of trace gas concentration profiles, while for our case – the aerosol retrievals – it uses an iterative Tikhonov regularisation approach. The main concept of the algorithm for the aerosol retrieval is to minimise the difference between modelled and measured  $O_4$  slant optical depths by applying the iterative Tikhonov technique to varied aerosol extinction profiles. This method uses the difference in the slant optical depth from an a priori state to obtain information on the aerosol concentration that caused this difference through multiple iterations. Slant column densities of trace gases and  $O_4$  from MAX-DOAS measurements at different line-of-sight (LOS) directions as well as climatology profile files are used as inputs. The BOREAS algorithm is based on the SCIATRAN radiative transfer model (Rozanov et al., 2005), which is used to calculate box air mass factors (BAMFs) and weighting functions needed for the profile inversion. The BAMF – in contrast to the total AMF – is a function of altitude describing the sensitivity of measurements to the profile at different atmospheric height layers. The aerosol weighting function matrices express the sensitivity of the  $O_4$  measurements to changes in the aerosol extinction coefficient profile. For the radiative transfer model (RTM) calculations, scattered light in a spherical atmosphere (multiple-scattering) and atmospheric profiles of pressure and temperature for Athens from the At-

mospheric Science Radiosonde Archive of the University of Wyoming (<http://weather.uwyo.edu/upperair/bufr/aob.shtml>, last access: 21 July 2020) are considered. The instrument was set to the station's altitude, and the surface was set at sea level. The aerosol inversion problem is expressed through the minimisation of Eq. (2):

$$\|\Delta\tau(\lambda, \Omega) - \Delta\tilde{\tau}(\lambda, \Omega, N_\alpha(z)) - P(\lambda, \Omega)\|^2 \rightarrow \min, \quad (2)$$

where  $\Delta\tau$  denotes the measured  $O_4$  differential slant optical thickness;  $\Delta\tilde{\tau}$  the simulated differential slant optical thickness;  $\Omega$  the measurement geometry (LOS, SZA, relative azimuth);  $N_\alpha(z)$  the a priori aerosol number concentration profile, which is used as a starting point for the iterations; and  $P$  a polynomial of lower order, which accounts for the attenuation due to scattering processes. Since the relationship between the concentration profile and the  $O_4$  differential slant optical depth is not linear, the iterative Tikhonov regularisation technique, along with weighting function matrices, is used for the solution of the minimisation problem (Bösch et al., 2018).

The uncertainty associated with each retrieved profile is computed by the algorithm. It is the sum of the noise and smoothing errors, which represent the impact of the measurements and of the a priori profile on the retrieved profile, respectively. These two errors have been calculated for each of our case studies separately and are presented in Sect. 3.2.

The temporal resolution of the measurements is about 15 min, which corresponds to the duration of one full scanning cycle through all directions over the city. The vertical sampling of the retrieved profile is 0.05 km, with the bottom layer considered at the sea level and the top layer at 4 km a.s.l. The AOD is calculated by integrating the BOREAS-retrieved aerosol extinction coefficient vertically. More details about the values assigned to each parameter are given in Sect. 3.2.

### 3 Results and discussion

#### 3.1 Selected case studies

The main objective of this study is to assess the retrieved aerosol profiles from MAX-DOAS measurements by comparing them with well-established sun-photometric measurements (CIMEL) and lidar retrievals. Therefore, certain cases had to be selected with available and valid data from all three instruments. Additionally, the selected cases had to coincide with cloud-free days as all of the used measurement techniques have more substantial uncertainties in the presence of clouds. During the period from January 2015 to June 2016, four cases were found to meet the above conditions, covering winter, summer and spring: (i) 5 February 2015 under the influence of a weak dust event, (ii) 9 July 2015 with enhanced morning levels of  $NO_2$  for this season (Gratsea et al., 2016), (iii) 10 July 2015 with typical levels of pollution and

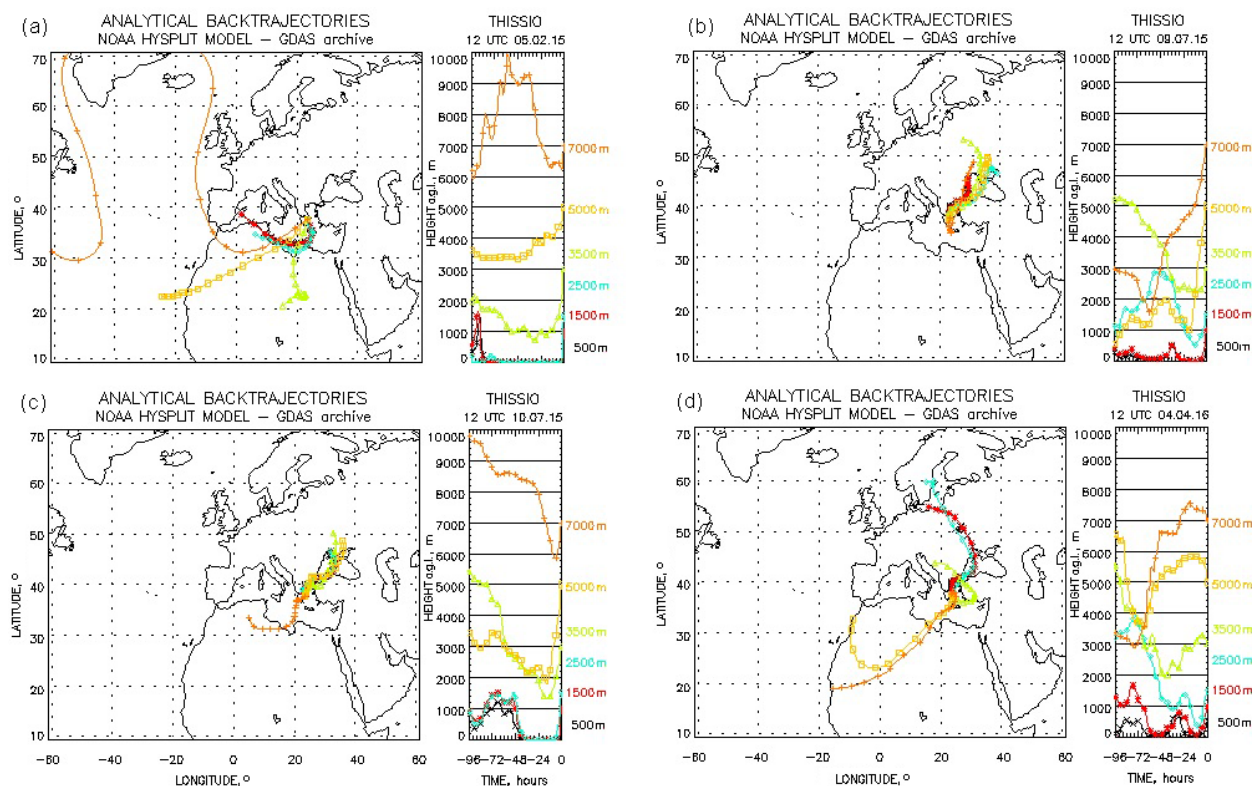
(iv) 4 April 2016 with enhanced levels of  $NO_2$ . In order to identify the sources of air masses reaching Athens on the specific dates, 4 d air mass back trajectories at different altitudes, calculated using the NOAA HYSPLIT (Hybrid Single-Particle Lagrangian-Integrated Trajectory) model (Draxler and Hess, 1997), were used. Potential for Saharan-dust transport below 4 km, which is the highest point of our retrievals, was identified only for case (i) (Fig. 2). In the rest of the cases, the air masses below 4 km originate from NNE directions and are thus not associated with dust aerosols. The  $NO_2$  levels, measured by MAX-DOAS and presented in Fig. 3, are used as an indicator for the pollution levels over the city. The mean diurnal course of  $NO_2$  differential SCs for winter and summer months, as reported by Gratsea et al. (2016), range from  $6 \times 10^{16}$  to  $9 \times 10^{16}$  and  $5 \times 10^{16}$  to  $11 \times 10^{16}$  molec  $cm^{-2}$ , respectively. Thus, enhanced pollution levels are observed during the morning hours in cases (ii) and (iv). The absence of clouds is established using in situ empirical meteorological observations from the monitoring station of the National Observatory of Athens at the centre of the city and is also verified by the MAX-DOAS-retrieved  $O_4$  slant columns throughout the day. The above-mentioned cases are henceforth referred to as cases (i), (ii), (iii) and (iv), respectively, and information about each case is summarised in Table 2.

#### 3.2 Aerosol extinction vertical profile retrievals

MAX-DOAS measurements and the BOREAS retrieval algorithm were used for the calculation of the diurnal aerosol extinction vertical distribution over the urban ( $S$ ) area (Fig. 4) for the selected case studies and for altitudes 0.5–4 km a.s.l. Single scattering albedo (SSA) and phase functions are not retrieved in BOREAS and have to be prescribed. Therefore AERONET measurements are used for specifying SSA ( $\omega$ ) and asymmetry factor ( $g$ ) values. However,  $\omega$  and  $g$  were not available in AERONET data for case (iv); therefore in this case the algorithm was run using the Henyey–Greenstein phase function with the monthly mean of SSA ( $\omega = 0.91$ ) and asymmetry factor ( $g = 0.68$ ) from the following year, as derived from the AERONET data (Table 3). Specifically for this case, sensitivity tests with varying  $\omega$  and  $g$  were carried out. It was found that the variability due to asymmetry factor is small and the impact of SSA negligible. A fixed surface albedo ( $\alpha = 0.15$ ), based on a previous study for Athens (Psiloglou et al., 2009), was used in all cases. Table 3 summarises the parameter settings used for the BOREAS retrieval.

The results for case (i) reveal a significant variation in the aerosol distribution in the vertical direction. Although the maximum retrieved extinction values in this case reach almost  $0.2 \text{ km}^{-1}$  at 1.5 km height in the afternoon, persistent high values are displayed until around local noon (Fig. 4). The temporal variation in the vertical distribution can be attributed to changes in the prevailing wind speed and direction





**Figure 2.** Analytical back trajectories for Athens as derived from the NOAA HYSPLIT model for the case studies (i) 5 February 2015 (a), (ii) 9 July 2015 (b), (iii) 10 July 2015 (c) and (iv) 4 April 2016 (d).

**Table 2.** Information about the selected case studies.

	Case (i)	Case (ii)	Case (iii)	Case (iv)
Date	5 February 2015	9 July 2015	10 July 2015	4 April 2016
Atmospheric conditions	Weak dust event, low pollution levels	High pollution levels in the morning	Typical pollution levels	High pollution levels
Air masses origin below 4 km	SSW	NNE	NNE	NNE

throughout the day; as recorded by the NOA's meteorological monitoring station at Thissio, the prevailing wind direction from 07:00 UTC until 10:00 UTC (LT = UTC+2 winter time and UTC+3 summer time) was from the south with speed from 1 to 4  $\text{m s}^{-1}$ , while easterly winds with speed reaching 10  $\text{m s}^{-1}$  started blowing at 11:00 UTC, efficiently ventilating the Athens basin and removing the dust and atmospheric pollutants. As shown in previous works conducted in the area (e.g. Fourtziou et al., 2017), wind speed below 3  $\text{m s}^{-1}$  favours the accumulation of pollutants.

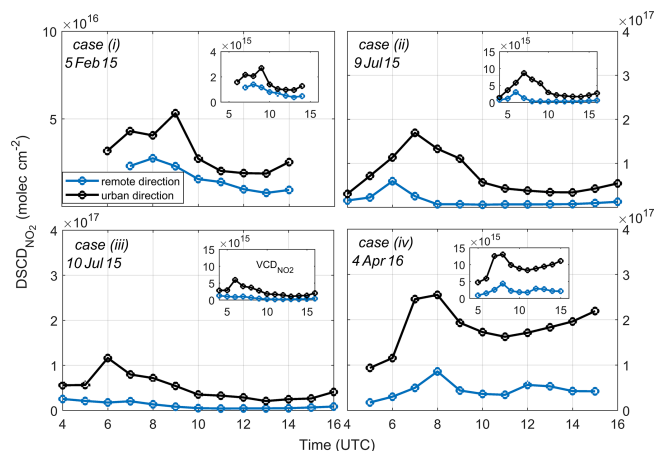
The two cases, (ii) and (iii), present an elevated aerosol layer extending up to 3 km between 10:00 and 14:00 UTC. Lidar retrievals also show an elevated extinction layer in both cases, as discussed in Sect. 3.3. However, the separation of the two layers could be an artefact which arises from the fact that the MAX-DOAS retrieval's response to a box-like dis-

tribution (e.g. a well-developed planetary boundary layer – PBL) leads to slight oscillations around this box due to the a priori smoothing. Both cases are related to weak prevailing winds ( $< 4 \text{ m s}^{-1}$ ), which favour the development of a vertically extended aerosol layer. The higher aerosol load in case (iii) is also corroborated by sun-photometric measurements, which are presented and discussed in Sect. 3.4.

Low levels of aerosol extinction (less than 0.1  $\text{km}^{-1}$ ) are present over the urban area throughout the whole day in case (iv). The highest values of the day (almost 0.14  $\text{km}^{-1}$ ) are displayed up to 800 m a.s.l. Given that the  $\text{NO}_2$  level, characteristic of anthropogenic pollution, is high during this day (Fig. 3), higher particle pollution levels would be expected.

**Table 3.** Settings used for the BOREAS retrieval. The mean daily value of each parameter ( $\omega$  and  $g$  retrieved from AERONET) is mentioned for cases (i), (ii) and (iii). The mean monthly values of  $\omega$  and  $g$  (provided from AERONET for April 2017) were used for case (iv) due to unavailable AERONET daily data around this date.

	Case (i)	Case (ii)	Case (iii)	Case (iv)
Surface albedo	0.15	0.15	0.15	0.15
Single scattering albedo ( $\omega$ )	0.92	0.96	0.93	0.91
Asymmetry factor ( $g$ )	0.78	0.65	0.68	0.68
Tikhonov parameter	20	20	20	20



**Figure 3.** Tropospheric retrievals of diurnal  $\text{SCD}_{\text{NO}_2}$  (elevation angle  $+1^\circ$ ) from MAX-DOAS measurements for the four selected case studies. The blue and the black curves correspond to the remote (W) and the urban (S) viewing direction, respectively. In the internal panels the corresponding tropospheric  $\text{VCD}_{\text{NO}_2}$  is also shown. Please consider the different scale used in case (i).

### 3.3 MAX-DOAS aerosol extinction profile evaluation

The BOREAS-retrieved aerosol extinction profiles from the MAX-DOAS measurements at 477 nm, between 0.5 km (station's elevation) and 4 km height, are compared with the lidar aerosol extinction coefficient measurements at 532 nm, between 1 and 4 km height, for the selected case studies (Fig. 6). Representative morning and afternoon snapshots during each day have been chosen to be presented and discussed. The lack of morning profiles for some days is due to the absence of lidar data; thus, both morning (mor) and evening (aft) data are available only for cases (i) and (ii). The lidar profile presented in each figure is the result of the mean lidar signal, averaged between the starting and the ending time of the corresponding MAX-DOAS profiles. The uncertainty in the lidar extinction profiles increases substantially for altitudes below 1000 m a.s.l. due to the loss of overlap between the telescope field of view and the laser beam (Wandinger and Ansmann, 2002; Kim et al., 2008; Papayannis et al., 2008); hence the lidar data for altitudes below 1000 m a.s.l. are not presented, and only measurements

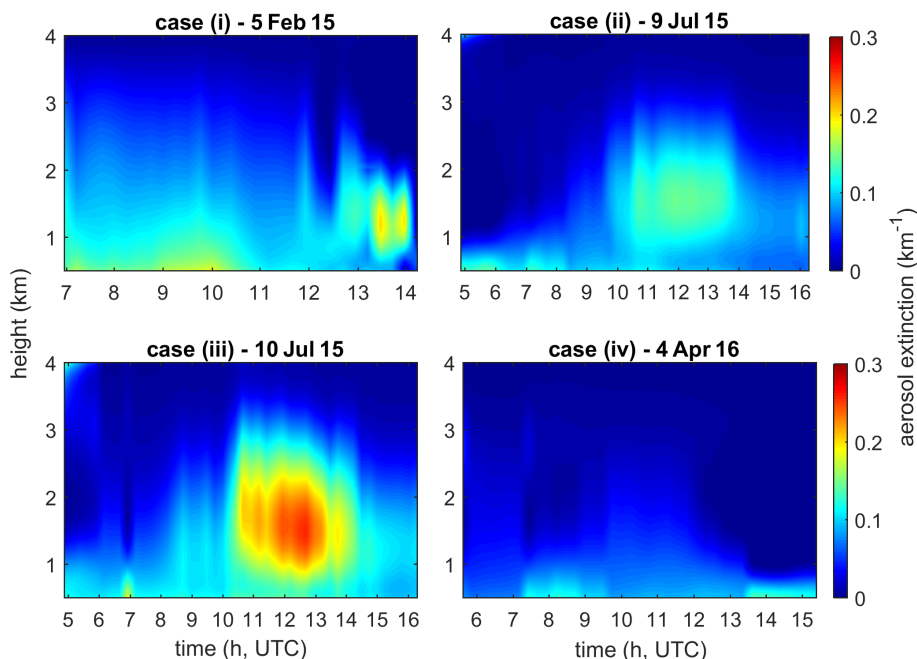
above 1000 m a.s.l. are considered for the calculation of the correlation between the two instruments. Another point that has to be considered when comparing the results from the two instruments is that the lidar profiles are characterised by high vertical and temporal resolution, and degradation to the sensitivity of the MAX-DOAS profiles is necessary in order to have a meaningful comparison to the MAX-DOAS data. According to the method described by Rodgers and Connor (2003), the degraded lidar profile  $x_f$  can be estimated by applying the equation

$$x_f = x_a + \text{AK} \cdot (x - x_a), \quad (3)$$

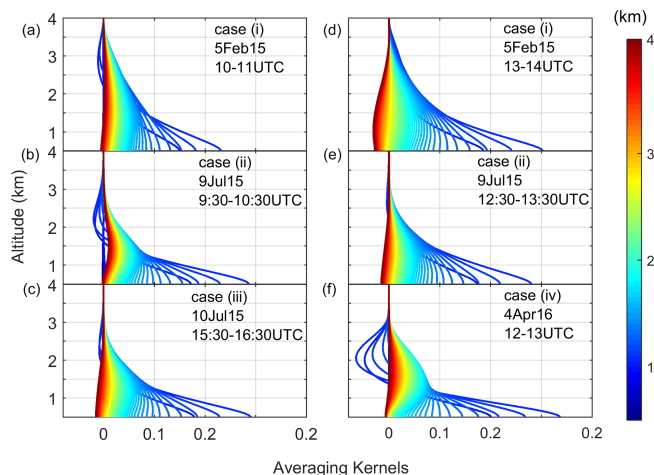
with  $x_a$  being the a priori profile used in the algorithm calculations,  $x$  the initial lidar profile and AK the averaging kernel from the BOREAS retrieval. The averaging kernel (Fig. 5) denotes the sensitivity of the retrieved profile to the true atmospheric profile for each layer, and in fact it represents the smoothing of the true profile in the retrieval. The lidar profile, degraded to 50 m vertical resolution, represents the MAX-DOAS profile that would have been retrieved if the true extinction profile were  $x$ . Last but not least, the horizontal distance (13 km) between the two measurement sites and the different operation principles of the two instruments should be noted. The lidar system retrieves information from the air mass right above the measurement site, while MAX-DOAS probes air masses along the line of sight of the telescope pointing from the top of a hill towards the city centre; hence the retrieved aerosol profiles from the two instruments correspond to different air masses and are not expected to fully agree, especially when the aerosol pollution is not horizontally homogeneous over the Athens basin. Thus, the comparison is mainly focused on a qualitative basis.

Each case is examined separately. Comparison information is given in the form of performance statistics – correlation coefficient ( $r$ ), median lidar / MAX-DOAS ratio, root mean square error (RMSE) and fractional gross error (FGE) – and is shown in Table 4. This set of statistics has been chosen as suitable to provide a detailed view of the algorithm performance; it has been proposed (Morris et al., 2005) that an FGE less than or equal to 0.75 is a criterion to evaluate good performance of an algorithm; therefore, any  $\text{FGE} > 0.75$  is used as an indicator of a relatively poor performance in this study. In order to perform the statistical cal-





**Figure 4.** MAX-DOAS-retrieved aerosol extinction vertical distributions (from instrument's height up to 4 km a.s.l.) for the four case studies over the urban area ( $S$ ). The spatial and vertical resolution of the retrievals is 50 m and 15 min, respectively.



**Figure 5.** Averaging kernels of the aerosol retrievals for the four case studies. For cases (i) and (ii), the left and right panels correspond to the morning and afternoon profiles, respectively. The colour bar represents the height of the atmospheric layers.

culations we averaged the four MAX-DOAS profiles comprising each case. Thus, all performance statistics have been calculated using the temporally averaged MAX-DOAS profile for each case and the corresponding degraded lidar profile so that both profiles are of the same temporal and vertical resolution. In all cases, 61 data points are used for the derivation of the statistics. The average smoothing and noise errors for the MAX-DOAS retrievals are given for each case study

separately in Table 5. In all cases, the noise error ranges between about 1 % and 5 %. In cases (i) (mor), (ii) and (iii), the smoothing error is about 15 %. The large smoothing errors in cases (i) (aft) and (iv) are due to the very small extinction values at higher altitudes.

### 3.3.1 Case study (i) – 5 February 2015

In case (i), the two instruments seem to be in excellent agreement in terms of correlation, with a very high correlation coefficient ( $r > 0.95$ ). In the afternoon, a peak in aerosol extinction ( $\sim 0.15 \text{ km}^{-1}$ ) between 1 and 1.5 km is captured by both instruments. The large discrepancy between the original and the degraded lidar profile is attributed to the fact that the AKs of the afternoon retrievals illustrate low sensitivity of the retrieved profile to the true atmospheric profile for altitudes up to 2.5 km (Fig. 5).

It should be mentioned that this is the only case in the present study where high aerosol load is found in the upper levels (free troposphere) in the original lidar profiles due to transboundary transport of aerosols at higher altitudes. The fact that, at these altitudes, the MAX-DOAS only agrees with the degraded lidar profiles (which means after including the AK information) suggests more significant errors in the a priori aerosol profiles and the reduced capacity of the MAX-DOAS to capture the characteristic inhomogeneity at higher atmospheric layers during aerosol transport episodes. Nevertheless, an overall satisfactory performance of the algorithm is indicated for the morning measurements by the FGE (0.31).

**Table 4.** Quantitative performance statistics of MAX-DOAS aerosol extinction calculations (BOREAS algorithm) compared to lidar measurements.

Performance measure	Case (i): mor	Case (i): aft	Case (ii): mor	Case (ii): aft	Case (iii)	Case (iv)
$r$	0.97	0.96	0.92	0.95	0.97	0.90
Median ratio (lidar / MAX-DOAS)	1.37	1.11	0.91	0.60	0.99	1.58
RMSE ( $\text{km}^{-1}$ )	0.03	0.03	0.02	0.04	0.01	0.02
FGE	0.31	0.80	0.37	0.54	0.20	0.61

**Table 5.** The MAX-DOAS average smoothing and noise errors (%) for each case study.

Uncertainties (%)	Case (i): mor	Case (i): aft	Case (ii): mor	Case (ii): aft	Case (iii)	Case (iv)
Smoothing error	15.59	90.52	16.69	13.61	17.46	53.65
Noise error	3.94	2.03	2.69	1.93	2.25	5.53

### 3.3.2 Case study (ii) – 9 July 2015

The retrieved MAX-DOAS profiles agree quite well with the degraded lidar profiles; they both show an aerosol layer extending up to about 2.5 km, and the correlation coefficient is very high ( $r \approx 0.95$ ) during both morning and afternoon measurements. In the afternoon, however, the MAX-DOAS measurements result in higher extinction levels by almost 65 % compared to the degraded lidar profile. As shown in Fig. 6 (middle row panels), in this case MAX-DOAS tends to overestimate the lidar extinction levels mainly at higher altitudes, a fact that can be attributed to the smoothing effect of the retrieval procedure on the true profile; given that a MAX-DOAS profile algorithm cannot retrieve sharp edges, the underlying narrow high-altitude enhancement in the afternoon propagates through the retrieval into a smoother and broader aerosol peak. The FGE, ranging from 0.35 to 0.55, indicates a good performance of the algorithm.

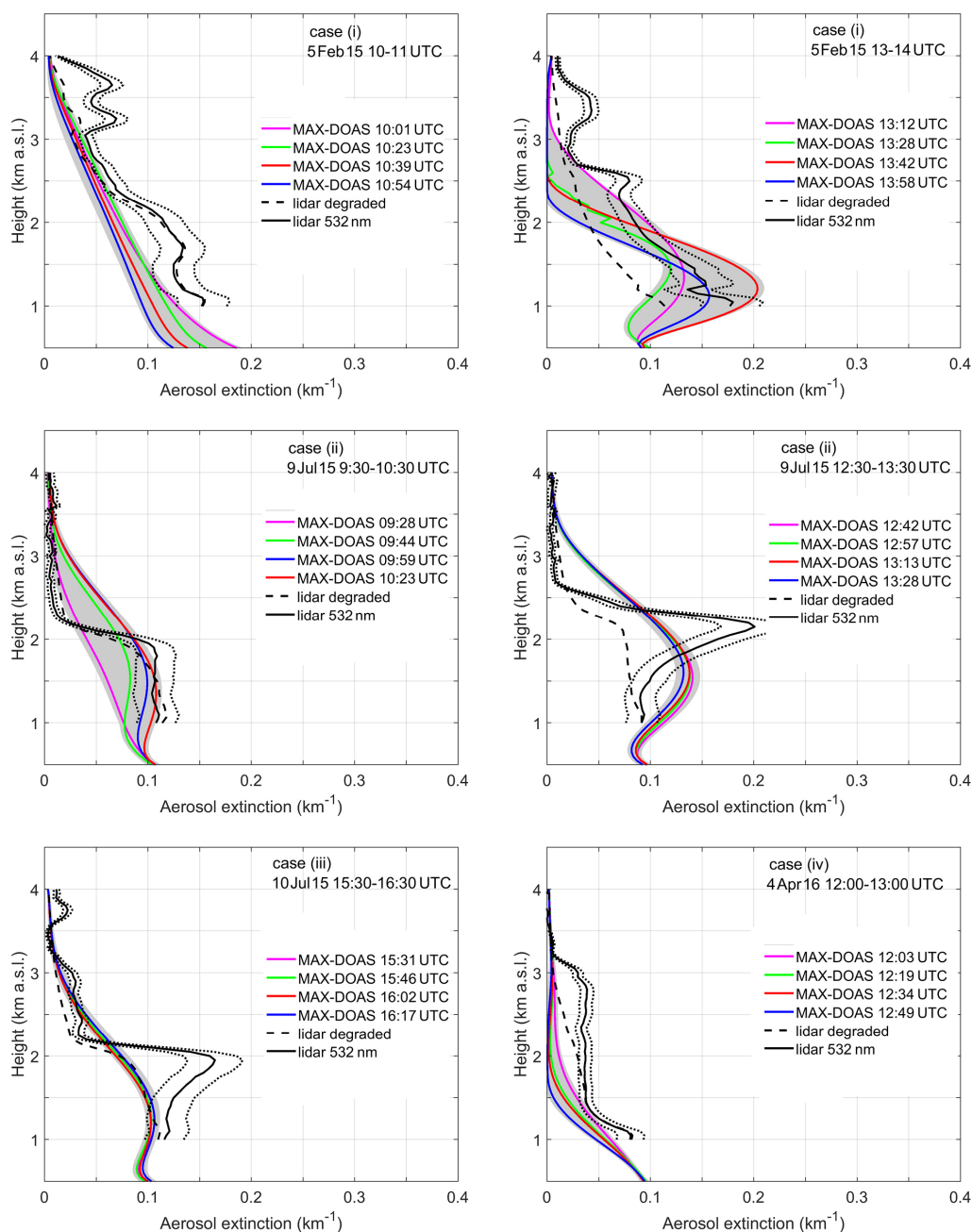
### 3.3.3 Case study (iii) – 10 July 2015

The two instruments seem to correlate very well ( $r = 0.97$ ). The MAX-DOAS coincides well with the aerosol extinction levels from the degraded lidar profile; the lidar-to-MAX-DOAS ratio is equal to 0.99. Nevertheless, when the original lidar profile is considered, a clear discrepancy in the extinction levels is present; the lidar peak value ( $0.16 \text{ km}^{-1}$ ) is enhanced by a factor of 2. The discrepancy between the original and the degraded lidar profile results from the low sensitivity of the averaging kernels for heights up to about 2 km (Fig. 5, case iii), which plays significant role in the degradation (smoothing) of the lidar retrieval. The RMSE is small ( $0.01 \text{ km}^{-1}$ ), and the low FGE (0.20) indicates good performance of the algorithm.

### 3.3.4 Case study (iv) – 4 April 2016

The profiles resulting from both instruments display an aerosol layer extending from the lower atmospheric layers up to 1.5 km height. The MAX-DOAS and degraded lidar profile shapes are very similar and highly correlated ( $r = 0.90$ ). However, the MAX-DOAS underestimates by almost 30 % the lidar aerosol extinction (median lidar / MAX-DOAS ratio = 1.58). Although the correlation is high, and the RMSE is small (0.02), the FGE (0.61) indicates a moderate performance of the algorithm for the specific case. This FGE value, however, results from the high median ratio of the two profiles, which in turn results from the low extinction levels since the absolute difference between the two profiles is not that large.

Overall, the correlation between lidar and MAX-DOAS measurements is very good ( $0.90 < r < 0.97$ ) in all cases, and a good agreement in the profile shape and altitude of the peak extinction level is also observed. The failure of the MAX-DOAS to capture clearly distinguished aerosol layers is attributed to the smoothing effect due to the presence of a priori constraints during the retrieval procedure. The RMSE ranges from 0.01 to  $0.04 \text{ km}^{-1}$  in all cases. A high FGE (0.80) has been calculated only in the case of measurements at small relative azimuthal angles between the viewing direction and the sun and in parallel presence of large particles. Due to the different operation principles of each instrument (active and passive remote sensing), the different wavelengths and the different air masses probed by each instrument, a full agreement in the derived profiles would not be expected. In particular, the lidar profiles represent the aerosols which are directly over the measurement site, whereas the MAX-DOAS profiles are representative of the atmosphere at a distance of several kilometres along the line of sight of the instrument. Another conclusion arising from these four cases is that the MAX-DOAS fails to detect part of the urban aerosol pollution when the pollution levels are low (e.g. case iv) and also



**Figure 6.** Comparison of retrieved MAX-DOAS aerosol extinction profiles at 477 nm (multicoloured curves), lidar aerosol extinction coefficient vertical profile at 532 nm (black curve) and the corresponding degraded lidar profile (dashed black curve) for the selected case studies. The lidar profile used in each case is the average profile retrieved between the starting and the ending time of the MAX-DOAS retrievals, and the light dashed black curves are the lidar-derived aerosol extinction uncertainty obtained by the lidar assumption of  $50 \pm 20$  sr. The grey shaded area represents the corresponding MAX-DOAS uncertainty.

fails to capture the inhomogeneity at higher altitudes in case of aerosol transport episodes.

### 3.4 AOD evaluation

In the previous section, lidar measurements were used for the evaluation of the aerosol extinction profiles obtained from

MAX-DOAS. However, a conclusive evaluation of MAX-DOAS aerosol optical depth (AOD) cannot be done strictly with lidar-derived AOD values mainly due to the lidar blind range (overlap height), resulting in AOD underestimation. Therefore, in this section, we focus on the comparison between the retrieved AOD from MAX-DOAS measurements

at 477 nm and from CIMEL measurements at two wavelengths (440 and 500 nm) during the aforementioned case studies (Fig. 7). For the MAX-DOAS AOD calculation, the missing values in the extinction coefficient profiles below 500 m are set to a constant value (equal to the retrieved value at 500 m). This assumes that the atmosphere is well mixed below 500 m, which probably results in an underestimation of the calculated AOD in case of enhanced surface aerosol layer. When looking at the figures, one should consider that the CIMEL AOD uncertainty is estimated to be approximately 0.01 for wavelengths  $> 400$  nm (Eck et al., 1999). The MAX-DOAS AOD uncertainties are shown in the figures. The Ångström exponent, derived from the CIMEL measurements (400–870 nm), is also taken into account as a qualitative indicator of aerosol particle size in order to investigate the origin of the aerosols (natural-dust or anthropogenic sources) and the performance of the MAX-DOAS retrievals for different aerosol types and sizes. An overview of the comparison statistics (described in Sect. 3.3), representative of the degree of agreement between MAX-DOAS and CIMEL measurements at 500 nm, is presented in Table 6. The calculations were made on an hourly basis to achieve uniform results regarding the air masses. Although this section is focused on the comparison with the AOD from CIMEL, the AOD from lidar measurements (calculated by integrating the aerosol extinction coefficient from ground up to the identified reference height of 4 km a.s.l.) is also presented indicatively. Nevertheless, as a complementary analysis, the comparison between lidar and MAX-DOAS AOD for the common altitude (1–4 km) – along with the corresponding uncertainties – is presented in Table 7.

### 3.4.1 Case study (i) – 5 February 2015

The very small Ångström exponent, ranging between 0.05 and 0.13 throughout the day, indicates the dominance of coarse particles in the aerosol distribution. Given the cloud-free sky conditions and the potential for dust transport found for this day by the NOAA HYSPLIT (Fig. 2), these particles are probably associated with the presence of dust in the atmosphere. The correlation between the two instruments is moderate ( $r = 0.47$ ), and the calculated AOD levels from the MAX-DOAS measurements underestimate by about 20 % the CIMEL measurements (median ratio CIMEL / MAX-DOAS = 1.22). The daily averaged AOD values are 0.33 ( $\pm 0.02$ ) and 0.39 ( $\pm 0.03$ ) for MAX-DOAS and CIMEL, respectively, and are much higher than the climatological monthly average value ( $0.27 \pm 0.03$ ) for February in Athens, as reported in Gerasopoulos et al. (2011). The RMSE is 0.07; however the FGE, which is small (0.17), implies excellent performance of the algorithm. The moderate correlation results may arise from the fact that CIMEL performs direct sun measurements, whereas MAX-DOAS measurements – and the subsequent AOD retrieval – are performed at a fixed azimuthal direction. Thus, the CIMEL measurements

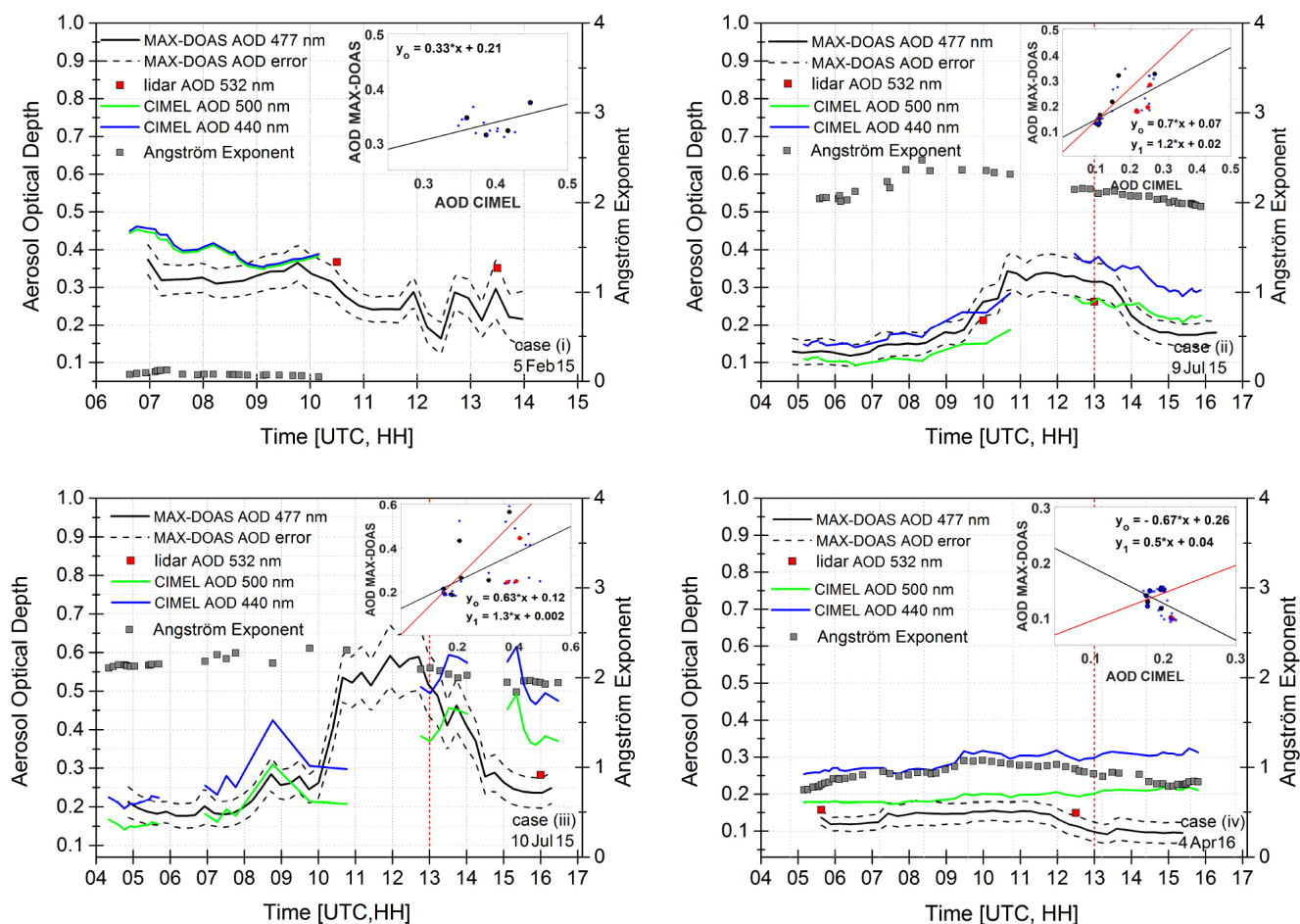
are highly affected by variations in the temporal and spatial distribution of the aerosols. The AOD from the lidar measurements at 10:00 and 13:00 UTC (0.37 and 0.35, respectively) coincides well both with CIMEL and MAX-DOAS measurements.

### 3.4.2 Case study (ii) – 9 July 2015

In this case, the large values of the Ångström exponent ( $\alpha \geq 2$ ) are indicative of the presence of fine-mode aerosols that are associated with urban pollution (Westphal and Toon, 1991; Eck et al., 1999; Gerasopoulos et al., 2011). The considerable levels of  $\text{NO}_2$  measured during this day (Fig. 3) indicate the presence of anthropogenic pollution. The two instruments are again moderately correlated ( $r = 0.67$ ); however if the afternoon measurements – after 16:00 LT – are excluded, the correlation becomes very good ( $r = 0.87$ ). The median ratio (0.85) indicates that the MAX-DOAS overestimates the AOD levels measured by the CIMEL. However, the overestimation is more profound during the morning, while in the afternoon the MAX-DOAS slightly underestimates the measured AOD by about 20 %, a fact that can be attributed to inaccuracies in the radiative transfer calculation for the forward-scattering geometry. The daily averaged AOD values are 0.20 and 0.19 for MAX-DOAS and CIMEL, respectively. The small values of both the RMSE (0.07) and the FGE (0.26) are indicators of very good performance of the algorithm. The gaps in the CIMEL data in this case, as well as in case (iii), are probably due to saturation of the instrument. The lidar AOD in this case (0.21 at 10:00 UTC and 0.26 at 13:00 UTC) agrees well with the CIMEL measurements at 500 nm but is lower than the AOD from the MAX-DOAS; the difference is more considerable in the afternoon.

### 3.4.3 Case study (iii) – 10 July 2015

In the third case study, the measurements from the two instruments seem to be in better agreement during morning hours. Overall, the MAX-DOAS AOD levels coincide well with CIMEL (median CIMEL / MAX-DOAS ratio is 0.95); however the underestimation due to light-scattering geometry after 13:00 UTC is about 35 %. The moderate results with respect to correlation ( $r = 0.53$ ) are due to the large discrepancy between the two instruments during the afternoon. If only the measurements until 13:00 UTC are considered, the correlation is considerably improved ( $r = 0.75$ ). Despite the non-satisfactory correlation, the calculated FGE (0.28) indicates a very good performance of the algorithm. The lidar-derived AOD in the afternoon is higher than the MAX-DOAS measurement. Unfortunately, no CIMEL or lidar data are available around noon in order to validate the aerosol plume captured by MAX-DOAS. It should be noted, though, that case studies (ii) and (iii) (both summer days in July) exhibit the same diurnal pattern: lower values in the morning, steadily increasing throughout the day and then



**Figure 7.** AOD as derived from MAX-DOAS (black curve) and CIMEL at 440 nm and 500 nm (green and blue curve, respectively). The grey and the red square markers represent the Ångström exponent derived from 400 and 870 nm and the lidar-derived AOD, respectively. The dashed black curves represent the MAX-DOAS AOD uncertainties. The scatterplots between hourly AOD calculated from MAX-DOAS measurements and hourly AOD from CIMEL at 500 nm are shown in the internal panels; the red points correspond to measurements after 13:00 UTC. Accordingly,  $y_0$  is the linear-regression equation with all the data points included, and  $y_1$  is the linear-regression equation when the data points after 13:00 UTC have been excluded. The smaller blue points are the raw data points. The vertical dashed red line separates the measurement data before and after 13:00 UTC.

**Table 6.** Quantitative performance statistics of MAX-DOAS AOD calculations (BOREAS algorithm) at 477 nm compared to CIMEL measurements at 500 nm.

Performance measure	Case (i)	Case (ii)	Case (iii)	Case (iv)
$r$	0.47	0.67	0.53	-0.42
Median ratio (CIMEL / MAX-DOAS)	1.22	0.85	0.95	1.37
RMSE	0.07	0.06	0.11	0.07
FGE	0.17	0.26	0.28	0.40

slightly declining in the afternoon. A similar diurnal AOD pattern was found for summer in Athens by Gerasopoulos et al. (2011), and this pattern has been associated with local urban or industrial sources (Smirnov et al., 2002).

### 3.4.4 Case study (iv) – 4 April 2016

The Ångström exponent in this case ( $\alpha \approx 1$ ) indicates the presence of coarse aerosols (radii  $\geq 0.5 \mu\text{m}$ ) in the atmosphere (Westphal and Toon, 1991; Eck et al., 1999). The NOAA HYSPLIT back trajectories show the potential for African dust transport to Athens, however, at higher altitudes



of up to 5 km. Nevertheless, despite the presence of coarse particles, the AOD levels are low; the daily averaged AOD values are 0.13 and 0.19 for MAX-DOAS and CIMEL, respectively. The MAX-DOAS underestimates the AOD with respect to CIMEL by about 25 % (and by 50 % if only the afternoon measurements are considered), the RMSE is 0.07, and the FGE is 0.40, yet it seems that the two measurement techniques are not correlated ( $r = -0.42$ ). However, the comparison in terms of correlation results in better outputs if only the morning measurements are considered; in this case the correlation coefficient is 0.75. It seems that MAX-DOAS, in this case, fails to detect the accumulation of coarse particles. Nevertheless, the AOD lidar measurements agree very well with the MAX-DOAS measurements (Table 7).

Overall, a systematic underestimation of the AOD, by 20 % to 35 %, by the MAX-DOAS is observed in the afternoon measurements, when the relative azimuthal angle between the MAX-DOAS viewing direction and the sun is small. Better agreement is achieved at large relative azimuthal angles in the morning. This finding has also been reported by Frieß et al. (2016) when comparing different retrieval algorithms with sun photometer measurements. Considering that (i) the sun photometer is located downtown (150 m a.s.l.), at lower altitude than the MAX-DOAS (527 m a.s.l.) and thus more sensitive to aerosols in the lower troposphere, and taking into account (ii) the absence of real measurements from MAX-DOAS for altitudes below 500 m a.s.l., an underestimation of the contribution of the urban pollution to the AOD retrieved by MAX-DOAS would be expected. Nevertheless, the MAX-DOAS seems to detect well the typical urban aerosols in the boundary layer; the mean AOD difference (CIMEL minus MAX-DOAS) of all the measurements is 0.03 with SD 0.08. Furthermore, CIMEL is a direct sun photometer, which means that in each measurement different air masses are detected, while the MAX-DOAS always points at the same direction; this operational difference is reflected in the non-satisfactory correlation. When fixed values of SSA and asymmetry factor (instead of AERONET data) are used by BOREAS, it seems that MAX-DOAS fails to detect accumulated coarse particles (e.g. case iv), leading to underestimation in case of small Ångström exponent values ( $< 1$ ). Frieß et al. (2016) have also come to this conclusion during the CINDI-2 campaign. However, the underestimation could also be attributed to the high-altitude aerosol layer detected by the lidar (Fig. 6); the MAX-DOAS' sensitivity at higher altitudes is low, and the calculated AOD is limited up to 4 km, while the AOD from CIMEL refers to the total atmospheric column. In addition, the presence of aerosol layers above 4 km that could contribute to the AOD was examined by using the lidar signals, and no significant aerosol load was observed above this height. It should also be noted that the standard AERONET version 2 algorithm uses an NO<sub>2</sub> climatology with a spatial resolution much coarser than the area of the city of Athens (Giles et al., 2019); hence in certain cases the difference be-

tween the MAX-DOAS and the higher CIMEL AOD levels at 440 nm could result from additional NO<sub>2</sub> content in the atmosphere, which is the case of highly polluted days. The lidar-derived AOD levels coincide well with the MAX-DOAS measurements.

#### 4 Summary and conclusions

An assessment of the retrieval of aerosol extinction profiles and AOD from MAX-DOAS measurements is presented for the first time for the urban environment of Athens. The profiling results are compared to lidar extinction profiles and to AODs obtained from sun-photometric measurements. The intercomparison results are very promising, showing that the MAX-DOAS measurements provide a good estimation of the aerosol vertical profile over Athens. Although this intercomparison is of great importance for the validation of the MAX-DOAS retrieval, the different operation, characteristics and measurement principles of each instrument, in addition to some comparison restrictions, have to be considered.

Regarding the spatial characteristics, (i) the measurements with the MAX-DOAS technique represent an area that includes the AERONET and the lidar locations, but it is not limited to them. Regarding the vertical aerosol information, (ii) the MAX-DOAS retrievals are representative of 500–4000 m a.s.l., while the lidar profiles are valid for altitudes higher than 1000 m above the station, and finally (iii) the AOD AERONET measurements describe the columnar aerosol properties representative of an area ranging from a radius of a few kilometres up to 10 km above the Athens area, depending on solar elevation. Also, (iv) the sun photometer AOD observations probe the extinction in the full atmospheric column, while MAX-DOAS retrievals are sensitive only to the lowest kilometres, leading to differences in the presence of aerosol layers at altitudes above 4 km. Nevertheless, despite the comparison restrictions and the differences in the three instruments, the comparison of the retrieved profiles and the AODs shows that the MAX-DOAS measurements bode well for the future of aerosol measurements, and they are able to provide a good estimation of the aerosol vertical distribution over Athens.

The vertical profiles retrieved by the BOREAS profiling algorithm applied to the MAX-DOAS measurements are qualitatively in good agreement with the lidar profiles smoothed with the MAX-DOAS averaging kernels; there is good agreement in aerosol layer shape and aerosol extinction levels, except in cases of inhomogeneity at higher altitudes, characteristic of aerosol dust transport episodes. Very good correlation ( $r > 0.90$ ) was found in all cases. A satisfactory fractional gross error ( $0.20 < \text{FGE} < 0.54$ ) has been calculated in all cases with fine aerosol particles (urban pollution), indicating a good performance of the BOREAS profiling algorithm in these cases. In some cases, the observed underestimation of the aerosol extinction (by 20 % to 35 %)

**Table 7.** MAX-DOAS (477 nm) and lidar (532 nm) AOD calculations for the atmospheric layer 1–4 km.

AOD (1–4 km)	Case (i): mor	Case (i): aft	Case (ii): mor	Case (ii): aft	Case (iii)	Case (iv)
Lidar	$0.24 \pm 0.04$	$0.21 \pm 0.03$	$0.13 \pm 0.03$	$0.19 \pm 0.03$	$0.19 \pm 0.03$	$0.09 \pm 0.01$
MAX-DOAS	$0.16 \pm 0.03$	$0.15 \pm 0.06$	$0.18 \pm 0.04$	$0.27 \pm 0.05$	$0.19 \pm 0.04$	$0.07 \pm 0.03$

by the MAX-DOAS at small relative azimuth angles can be attributed to the geometry of Mie scattering in relation to the location and viewing geometry of MAX-DOAS, resulting in MAX-DOAS' failure to detect part of the urban aerosol pollution. Overall, the agreement between the two instruments is encouraging, especially when considering the different nature of each technique and the different instrument locations, suggesting that the MAX-DOAS can accurately enough represent the aerosol vertical distribution.

The MAX-DOAS-retrieved AODs show satisfactory agreement with the sun-photometric measurements in terms of AOD levels. The MAX-DOAS underestimates the AOD in the presence of coarse particles; CIMEL / MAX-DOAS ratio > 1 coincides with Ångström exponent values < 1. A systematic underestimation by MAX-DOAS is observed in the afternoon measurements due to MAX-DOAS' viewing geometry. Overall, the MAX-DOAS can be considered to be an effective means for measuring the aerosol levels in Athens; the average AOD difference in all measurements between the two instruments is 0.03. It is important to note that in Athens, a highly populated and polluted area, horizontal gradients, especially in anthropogenic aerosols, are very likely to occur, resulting in different air masses detected by each instrument and subsequently in discrepancies between MAX-DOAS and CIMEL measurements.

This intercomparison is of great importance for the validation of the MAX-DOAS retrieval. Despite the already-mentioned limitations due to different operation, characteristics and measurement principles of each instrument, this work demonstrates that the MAX-DOAS measurements in Athens and the BOREAS algorithm can provide a good estimation of the aerosol vertical structure of the urban atmosphere on a continuous and long-term basis, offering a reliable data set for scientific studies. There is certainly more work to be conducted in future studies in order to understand the sensitivity of the MAX-DOAS aerosol measurements based on different aspects of urban-pollution evolution and long-range-transported aerosols.

*Data availability.* All data sets used and produced for the purposes of this work are freely available and can be requested from the corresponding author.

*Author contributions.* MG, PK and AR conceived the presented idea. MG performed the analysis and prepared the manuscript. TB

developed the profile retrieval algorithm and provided guidance to MG on the algorithm calculations and parameterisation. AR provided guidance to MG on MAX-DOAS data retrieval and profile calculations. MG designed the figures with support from PK and TB. AR and SK advised MG on the results interpretation. PK, AP and MM provided the lidar data. VA provided the sun photometer measurements. AT made comments on the comparison between the sun photometer and MAX-DOAS. AR, EG, SK, NM and MV provided critical feedback. All authors provided comments that helped shape the manuscript.

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