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Optical characteristics of biomass burning aerosols over Southeastern Europe determined from UV-Raman lidar measurements

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Abstract

The influence of smoke on the aerosol loading in the free troposphere over Thessaloniki, Greece is examined in this paper. Ten cases during 2001–2005 were identified when very high aerosol optical depth values in the free troposphere were observed with a UV-Raman lidar. Particle dispersion modeling (FLEXPART) and satellite hot spot fire detection (ATSR) showed that these high free tropospheric aerosol optical depths are mainly attributed to the advection of smoke plumes from biomass burning regions over Thessaloniki. The biomass burning regions were found to extend across Russia in the latitudinal belt between 45° N–55° N, as well as in Eastern Europe (Baltic countries, Western Russia, Belarus, and the Ukraine). The highest frequency of agricultural fires occurred during the summer season (mainly in August). The data collected allowed the optical characterization of the smoke aerosols that arrived over Greece, where limited information has so far been available. Two-wavelength backscatter lidar measurements showed that the backscatter-related Ångström exponent ranged between 0.5 and 2.4 indicating a variety of particle sizes. UV-Raman lidar measurements showed that for smoke particles the extinction to backscatter ratios varied between 40 sr for small particles to 100 sr for large particles. Dispersion model estimations of the carbon monoxide tracer concentration profiles for smoke particles indicate that the variability of the optical parameters is a function of the age of the smoke plumes.

1 Introduction

Biomass burning is a major source of air pollution and the second largest source of anthropogenic aerosols (IPCC, 2001). The Fourth Assessment Report of the Intergovernmental Panel on Climate Change (Forster et al., 2007) reports a contribution of roughly $+0.04 \text{ W/m}^2$ of biomass burning aerosol to the global radiative forcing (RF) with a standard deviation of 0.07 W/m^2 . This estimate of the direct RF is more positive than that of the Third Assessment Report (IPCC, 2001) and it is linked with improvements in

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the models in representing the absorption properties of biomass burning aerosol and the effects of their vertical distribution. Textor et al. (2006) showed that there are still significant uncertainties in the aerosol vertical distribution in global aerosol models, information that is critical in assessing the magnitude and even the sign of the direct RF of biomass burning aerosols.

In the last decade, a large number of biomass-burning studies concentrated on the physical, chemical, and thermodynamic properties of biomass-burning particles. A considerable body of research using in situ measurement techniques has been published on the optical properties of smoke. This includes the analysis of laboratory burns (Patterson and McMahon, 1987) and intensive field campaigns involving large-scale forest fires in the North America (Hobbs et al., 1996), as well as over Brazil (Andreae et al., 1996; Kaufman et al., 1998) and over Africa (Eck et al., 2001). Finally, smoke studies using remote-sensing techniques have included ground-based sun photometry (Kaufman et al., 1992) and quantitative inversions of satellite imagery for smoke optical properties (Ferrare et al., 1990). A detailed review of biomass burning literature is presented in Reid et al. (2005a; 2005b).

Studies mentioned above are referring to laboratory, in-situ and columnar sunphotometric measurements mainly of fresh smoke. Smoke particles from biomass burning change rapidly in size and composition after being emitted into the atmosphere (Westphaal and Toon, 1991; Liousse et al., 1995; Hobbs et al., 1997; Reid et al., 1998). There are a variety of mechanical, chemical and thermodynamic processes that can alter smoke properties. Under most circumstances, measurements of the characteristics of fresh smoke plumes and laboratory burns cannot be assumed to be applicable to the aged smoke that is of greatest interest when studying its radiative effects or its impact on regional air quality (Reid et al., 2005a). Evidence suggests that most particle growth in size and mass occurs on fairly short time scales (Abel et al., 2003; Reid et al., 1998). Excluding scavenging processes, studies on smoke particle evolution and growth have focused on Brownian coagulation in dispersing plumes, and gas-to-particle conversion and exchange (e.g., Turco and Yu, 1997). These calculations indicate that

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after a plume has dispersed into a regional haze or is undergoing long-range transport, particle coagulation is the significant mechanism for particle growth over long enough timescales. If this evolution of smoke particles is not properly accounted for, large errors may result in calculations of their RF on regional and global scales (Müller et al., 2005)

For an accurate smoke characterization, first the smoke particles have to be identified in a non-mixed aerosol state and that can only be achieved by studying lofted smoke layers in the free troposphere on the basis of long-term observations by means of lidar technique. Up to date, only few lidar studies investigated the optical, radiative (Wandinger et al., 2002; Balis et al., 2003) and microphysical properties (Müller et al., 2005) of smoke particles in lofted layers. Mattis et al. (2003) reported on strongly enhanced values of particle backscattering and extinction in the free troposphere over Germany in spring/summer of the year 2003, when a large-scale event of free tropospheric transport of biomass burning particles from Siberia occurred. The above mentioned studies were based on Raman lidar observations carried out in the framework of the European Aerosol Research Lidar Network (EARLINET) (Bösenberg et al., 2003). Ground-based lidar measurements of particle optical properties with high spatial and temporal resolution give detailed information on the occurrence, extent and development of aerosol structures. The EARLINET can follow the evolution of smoke plumes from Siberia and Russia over the European continent and report on changes of the optical properties of these particles with age.

Forest fires in Russia are a major source of pollution in the Northern Hemisphere (Wotawa et al., 2001). For that reason, there has been increased interest in recent years to assess the impact of these fires on climate (e.g. Kasischke and Bruhwiler, 2003). According to an investigation of the spatial and temporal occurrence of fires in croplands based on MODIS (Moderate Resolution Imaging Spectroradiometer) active fire product (Koronzi et al., 2006), the Russian Federation was found to be the largest contributor to agricultural burning globally during the period 2001 to 2003, producing 31–36% of all agricultural fires. This globally highest concentration of agri-

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cultural fires, found to be extended across Russia in the latitudinal belt between 45° N–55° N during the spring (April–May), as well as in Eastern Europe (Baltic countries, western Russia, Belarus, and the Ukraine) during the late summer (August).

In this study we present analyses of smoke optical properties measured over South-eastern Europe based on measurements of the EARLINET lidar station of Thessaloniki during ten episodes of very high free tropospheric aerosol optical depths (AOD) caused by the advection of air masses over Greece, mainly from biomass burning agricultural regions in Russia and Ukraine. In Sect. 2 we describe the instrumentation and the dispersion modeling tools used. Section 3 presents a smoke case study as an example of the analysis followed for all the days of our smoke measurements. The variability of the optical properties is then attributed to smoke ageing by coagulation leading to variable observed mixing state and size distribution of the agricultural fire aerosol investigated. Finally, our conclusions are given in Sect. 4.

2 Instrumentation and methods

2.1 Description of the lidar system and lidar data processing

At the Laboratory of Atmospheric Physics (LAP) (50 m above sea level), Aristotle University of Thessaloniki, a 355 nm Raman-backscatter lidar is used to perform continuous measurements of suspended aerosols particles in the Planetary Boundary Layer (PBL) and the lower free troposphere (FT). It is based on the second and third harmonic frequency of a compact, pulsed Nd:YAG laser, which emits pulses of 300 and 120 mJ at 532 nm and 355 nm, respectively, with a 10 Hz repetition rate (more technical details can be found on: Balis et al., 2000; Amiridis et al., 2005). The optical receiver is a 500 mm diameter telescope. The LAP lidar was successfully intercompared with other EARLINET instruments showing on the average an agreement of better than 5% for heights above 2 km in the backscatter coefficient (Matthias et al., 2004). The lidar-data handling procedures that are routinely used at LAP for the application of Fernald-Klett

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and Raman retrieval methods were successfully tested by two algorithm intercomparisons that took place during the EARLINET project (Böckmann et al., 2004; Pappalardo et al., 2004) and include error estimates for all participating groups. According to these studies the mean deviation from the solution in the determination of the backscatter coefficient with the Klett method was 5.5% at 355 nm and 2.9% at 532 nm, while the mean deviations from the solution for extinction and backscatter calculations at 355 nm with the Raman method was around 10% and 7% respectively. For the height ranges 2000–4000 m the overall estimated error in the measurements is up to 10% for the extinction coefficient, and up to 30% for the backscattering coefficients and the lidar ratio. The incomplete overlap between the laser beam and the receiver field of view affects significantly lidar observations of particle optical properties in the near-field range. At LAP we apply an overlap correction based on a technique proposed by Wandinger and Ansmann (2002), down to the height where the overlap function is equal to 0.5. This correction allows extending the profile in most cases down to 1000–1500 m above sea level. An application of this methodology and its experimental verification is presented in Balis et al. (2002).

2.2 Particle dispersion model and satellite hotspot detection

To simulate transport processes, we employed the Lagrangian particle dispersion model FLEXPART (version 6.3) (Stohl et al., 1998, 2005). Information on FLEXPART can be obtained via the internet at <http://transport.nilu.no/flexpart>. FLEXPART was driven here by operational analysis data from the European Centre for Medium-Range Weather Forecasts (ECMWF) model. The ECMWF data had 60 model levels and 1 × 1 degree global resolution. For the emission inputs for carbon monoxide, nitrogen oxides and sulfur dioxide, the Emission Database for Global Atmospheric Research (EDGAR) version 3.2 emission inventory for the year 2000 (fast track) on a 1 × 1 degree grid was used. Backward simulations were performed for 500-m intervals along a vertically stacked array at the location of our surface station for times when particularly interesting measurements were obtained (see <http://zardozi.nilu.no/~andreas/EARLINET/>).

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Every simulation consists of 40 000 particles released in the volume of air sampled. The backward simulations were done with full turbulence and convection parameterizations. The mathematical theory of these computations is described in Seibert and Frank (2004), and an application to aircraft measurements was presented by Stohl et al. (2003). The model output is a sensitivity field of emission fluxes, which were stored on a 3-d grid with three altitude levels (0–100 m, 100–3000 m, and above) and a horizontal resolution of 1×1 degree globally (0.5×0.5 degree resolution in a nest over the area of most interest). The emission sensitivity in the 0–100 m layer is of high interest since emissions typically occur at or near the ground. By multiplying the emission sensitivity value with the emission flux, a gridded source contribution field is obtained which, after spatial integration, yields a mixing ratio at the measurement site.

For estimating the smoke emission for the fires under study, we used fire count data from the low resolution Along Track Scanning Radiometers ATSR-2 onboard the ERS-2 satellite (2000–2002) and AATSR onboard ENVISAT (2003–2005). ATSR allows the detection of active fires using its thermal infrared channel centered at 3.7 μm. ATSR data was obtained from the Ionia website of European Space Agency, ESA (<http://dup.esrin.esa.int/ionia/wfa/index.asp>). Hotspot data used in this study were calculated with the version 2 algorithm that uses a detection threshold of 308° K (Dozier, 1981).

Following Seiler and Crutzen (1980) the emissions of carbon monoxide (CO) from the fires were estimated. CO is often used as a tracer for long-range pollution transport and will be used here to identify when and at what altitudes the atmosphere above Thessaloniki was affected by fire emissions. CO emissions were calculated using the equation:

$$E = ABab \quad (1)$$

where A is the area burned, B is the biomass per area, a is the fraction of the biomass consumed by the fire, and b is the CO emission factor. Every detected fire was linked to a certain land cover type, using a global land cover classification with a resolution of 1 km (Hansen et al., 2000), for which a certain biomass per area was assumed. The

largest uncertainty in this calculation is the area burned for a detected hot spot, here taken to be 600 ha. Since this number is more or less arbitrary for the ATSR data, the resulting tracer concentrations are rather qualitative. The emission calculations (but based on MODIS hot spot data) are described in detail in Stohl et al. (2007).

3 Results and discussion

Regular aerosol extinction and backscatter measurements using a UV Raman lidar have been performed from January 2001 to December 2005 at Thessaloniki, Greece, in the framework of the EARLINET project. Dispersion model simulations showed that for ten days of our lidar measurements, smoke was advected over Thessaloniki from regions with strong fire activity in Eastern Europe and Russia. These dates of smoke presence and the main source of particles are presented in Table 1. The height ranges of smoke layers are additionally denoted in Table 1 along with the lidar derived optical parameters for the specific layers, namely the AOD, the mean lidar ratio and the mean backscatter related Ångström exponent.

Figure 1 shows an aggregate plot of all (A)ATSR fire detections for the ten days of the lidar measurements. Location of the burning regions was also verified by the MODIS Fire Product (<http://modis-fire.umd.edu/products.asp>). Symbols indicate the dominant land cover where the detection occurred based on a global land cover classification with a resolution of 300 m derived from the European Space Agency's Global Land Cover service for the production of global land-cover map for the year 2005, using ENVISAT MERIS Fine Resolution (300 m) data that replaced the GlobCover2000 product which had 1 km resolution (<http://dup.esrin.esa.int/projects/-summary68.asp>). Figure 1 strongly suggests that smoke particles discussed in this paper were produced mainly by agricultural burning activities.

To explore whether the observed aerosols in our lidar profiles have advected over Thessaloniki from the agricultural fires mentioned, the Lagrangian particle dispersion model FLEXPART was used. A case study example for the 20 August 2001

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(18:00 UTC) is presented here to describe the analysis followed for the ten days used in this work. In Fig. 2 the column-integrated emission sensitivity obtained from the backward simulation with FLEXPART for the height range 2.5–3.0 km over Thessaloniki (corresponding simulations were done for all other 500 m intervals up to 10 km altitude) is presented along with the hot spots detected by ATSR, representing the position of fires as black dots. FLEXPART was run backward in time from the smoke layers over Thessaloniki, as those detected by the lidar profiles. According to Fig. 2 for the 20 August 2001, the vertically integrated emission sensitivity shows a maximum over Bulgaria and Ukraine, where the retroplume was located approximately on days 1 and 2, backward in time. As one can see in Fig. 2, the regions with the largest emission sensitivity are regions with strong fire activity. The hot spots detected by ATSR are superimposed on the column-integrated emission sensitivity map only in grid cells where the daily column-integrated emission sensitivity on the very day of the hot spot identification is above 8 ns m /kg, indicating the significant possibility of smoke advection.

To estimate the age of the smoke air mass detected by our lidar measurements, FLEXPART simulations have been used for the estimation of a carbon monoxide (CO) tracer profile over Thessaloniki. An example is presented in Fig. 3 for the 20 August 2001 at 18:00 UTC. CO concentration profiles have been estimated with the assumption of CO emissions originating only from fires. Based on ATSR hotspot detection, CO emissions were calculated using Eq. (1). For these calculations we have assumed a fraction of the biomass consumed by the fire equal to 600 ha/hotspot, which in general is considered a low estimation. This value was subjectively determined, since no information on how much area was actually burned per detected hotspot is available. The uncertainties in CO fire emissions in our calculations are even larger due to the limited capability of ATSR to correctly detect agricultural fires. For these two reasons, the final simulated CO vertical distributions from biomass burning activities used in this study might be rather low in absolute concentration terms. However, under-estimation of CO should not affect our study, since in the following we will use only relative differences between the ten days of simulations. This under-estimation must be of common

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magnitude for all cases examined due to the linearity of Eq. (1) and changes in the constant parameters assumed would only scale CO concentrations to higher values, affecting only the absolute values of the simulated relative vertical distributions.

For the ten days of the smoke advection over Thessaloniki as indicated from dispersion calculations, we have calculated the extinction and backscatter coefficient profiles from the measured Raman lidar signal at 355 nm and the elastic-backscatter signals at 355 and 532 nm. As an example we present in Fig. 4 these profiles together with the derived lidar ratio at 355 nm and the backscatter-related Ångström exponent (calculated from 355 and 532 nm backscatter coefficients) for the 20 August 2001, between 18:24 and 19:18 UTC. The error bars indicate statistical errors corresponding to signal averaging. The backscatter maximum at both wavelengths for that case was observed within the 1.5 and 4.5 km height range. This aerosol layer showed quite a stable maximum during the measurement period. The lidar ratio was almost constant with an average value of 47.6 ± 4.6 sr. The lack of significant vertical variability of the lidar ratio suggests the presence of aerosol of the same type throughout this layer. Backscatter-related Ångström exponent for the same height range was found to be equal to 2.25 ± 0.16 . Hence the lidar profiles, the dispersion modeling results shown in Figs. 2 and 3 and additional trajectory analysis (not shown here) confirmed that air masses in the height region between 1.5 and 4.5 km originated solely from regions with biomass burning activity, suggesting the presence of smoke particles in the free troposphere over Thessaloniki.

The above presented analysis for the 20 August 2001 was repeated for the entire ten day data-set discussed in this paper, where smoke advection over Thessaloniki was strongly indicated by the synergy of lidar measurements and dispersion modeling. Vertical profiles of the extinction coefficient (355 nm), backscatter coefficient (355 and 532 nm), lidar ratio (355 nm) and backscatter-related Ångström exponent (355/532 nm) referring to smoke particles for the days discussed are presented in Fig. 5. The dates follow the same colour-code as the profiles on the figure. Smoke layers, as they have been identified by emission sensitivities calculated for different arrival heights above

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the station, are indicated with different colour for each day. Only the smoke layers are indicated with colours, while black lines correspond to the complete vertical profile measured for each day. The lidar ratio for smoke particles found to vary between 32 and 103 sr. The backscatter related Ångström exponent ranged between 0.5 and 2.4.

The variability of the lidar ratio and Ångström exponent for smoke particles cannot be attributed to different source characteristics, since for the case studies discussed here the main and common source of smoke is the agricultural fires in Russia and Ukraine. In this region, farmers burn each year the crop stubble off recently harvested fields in anticipation of the next sowing. The variability in particle size can be attributed to the fact that the fire regions were located in variable distances from Thessaloniki which affects the optical characteristics of the observed smoke aerosols due to different ageing processes, depending also to the meteorological conditions responsible for the speed of the smoke plume.

To investigate the variability of the Ångström exponent during the days of this study, we attempted to estimate the relative age of the air mass, by calculating weighted averages of the CO age over Thessaloniki from the model estimated vertical profiles based on fire emissions (shown in Fig. 3 for the example of 20 August 2001). Only the height ranges with smoke presence identified both by lidar measurements and FLEXPART simulations were considered. For the calculation of the mean age of CO from biomass burning activities, the CO concentrations were used as weights. Calculated age averages are plotted against vertical mean values of the backscatter related Ångström exponents binned at 0.5 km in Fig. 6. The correlation coefficient was found equal to -0.85 suggesting a strong anti-correlation. This suggests that smoke particles become larger with ageing. There is strong evidence from observations of biomass-burning plumes in different regions of the world that particles grow in size during the ageing of the plumes, (e.g., Fiebig et al., 2003). Processes that lead to the increase of particle size are, gas-to-particle conversion of inorganic and organic vapors (Reid and Hobbs, 1998; Reid et al., 1998), condensation of large organic molecules from their gas phase in the first few hours of ageing (Reid and Hobbs, 1998; Pósfai et al., 2004),

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particle growth due to coagulation (Westphal and Toon, 1991; Fiebig et al., 2003), and photochemical and cloud-processing mechanisms. Liousse et al. (1995) found a significant increase of particle size with age of plumes that originated from savannah fires in Ivory Coast in Africa. A decrease of the Ångström exponent (450–550 nm wavelength), which is equivalent to an increase of particle size was observed for fires in tropical forest and cerrado during SCAR-B campaign (Reid et al., 1998). In situ observations showed that Ångström exponents were on the order of 2.2 ± 0.2 for fresh smoke and 1.2 ± 0.2 for aged smoke. The values for aged smoke are very similar to the values reported here for the backscatter-related Ångström exponent (355–532 nm wavelength) calculated from lidar observations. A shift of particle size toward larger values was also found here, since Ångström exponents of the order of 0.5 were also observed. There seems to be a systematic shift of particle size toward larger values with increasing distance from the source of fires. Lidar observations of an aged biomass-burning plume that had originated from western Canada in 1998 and which was observed over Germany during the Lindenberg Aerosol Characterization Experiment (LACE) 98 showed that the Ångström exponent was as small as 0.06 which indicates well-aged smoke particles even larger than the ones found in this study (Fiebig et al., 2002; Wandinger et al., 2002). Dual-wavelength Raman lidar observations of Siberian forest fire smoke were carried out at Tokyo on 21 May 2003 (Murayama et al., 2004). Smaller particles than those observed in Leipzig on 29 May 2003 were found with an Ångström exponent on the order of 1.35. It has to be emphasized here that the extinction-related and the backscatter-related Ångström exponents can differ and a direct comparison between these parameters is not in general appropriate. However, it has been shown by Müller et al. (2005) that the lidar ratio at 355 and 532 nm for smoke particles is almost equal. No spectral variability of the lidar ratio is mathematically translated to the equivalent of the Ångström exponents for the extinction and the backscatter. Finally, we have to mention that the fires described here and the fires discussed in the cited literature, certainly have different properties. Thus, conclusions regarding the increase of particle size with ageing have to be treated with caution to each case (depending on source

characteristics) and are subject for future studies.

In Fig. 6, an anti-correlation of the same order (-0.84) is presented for the vertical mean values of the lidar ratio and the backscatter related Ångström exponent, calculated for the smoke layers under investigation. This figure suggests that the lidar ratio for smoke particles is also a function of the particle size. In general, the lidar ratio can be calculated by Raman lidars that can perform independent measurement of backscatter and extinction profiles (Ansmann et al., 1992). However, most lidars used in aerosol research are elastic-backscatter lidars and in that case, the lidar ratio is an important parameter for the determination of the particle backscatter coefficient from the elastic-backscatter signal alone. New space-borne missions like the Cloud-Aerosol Lidar and Infrared Pathfinder Satellite Observation (CALIPSO) need the lidar ratio information to invert the elastic-backscatter signals to reliable backscatter profiles. The lidar ratio for backscatter lidar inversions is usually being hypothesized based on aerosol type assumption. Usually, the aerosol type is assumed based on synergetic satellite data and modeling aerosol source estimations (for e.g. air-mass trajectories). Additional lidar parameters measured, like the depolarization ratio and the backscatter-related Ångström exponent (for multi-wavelength backscatter lidars) could be synergistically used to characterize the aerosol type. However, the choice of a representative constant lidar ratio for a specific aerosol type could be a large source of uncertainty in backscatter retrievals. As we are showing here, the lidar ratio is variable for smoke particles, depending mainly on their size. Specifically for the case of smoke particles, Reid et al. (1998) suggested that if just one optical property of the particles is measured (e.g. the Ångström exponent), then some of the other properties of the particles, such as particle size distribution parameters and single-scattering albedo, can be deduced. In this paper we apply this suggestion for the lidar ratio. For smoke particles and in the case of multi-wavelength elastic-backscatter lidar (e.g. CALIPSO), one measures the backscatter-related Ångström exponent and assumes a specific value for the lidar ratio based on the aerosol type. In this paper we demonstrated that for the smoke case the lidar ratio vary with size of the particles. The choice of the lidar ratio for the retrieval

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of backscatter profiles from multi-wavelength backscatter lidars can be improved using relationships extracted by ground-based Raman lidar measurements of a specific aerosol type like the one presented in Fig. 6. However, we have to mention that the choice of the lidar ratio is more complicated in mixed aerosol states. The state of mixing and the age of the air mass are the reasons that the lidar ratio is not constant since this parameter is dependent on the size distribution and the refractive index. Only in a well defined aerosol layer of a specific type, the lidar ratio can be assumed based on ground-based Raman lidar measurements.

4 Summary and conclusions

In this paper, Raman lidar measurements for ten selected cases during 2001–2005, when air masses were advected over Thessaloniki from regions with intense fire activity, have been presented, in terms of extinction (355 nm) and backscatter coefficient (355 and 532 nm) and the corresponding lidar ratio (355 nm) and backscatter-related Ångström exponent (355/532 nm). Lidar ratios in the range of 32–103 sr were observed for smoke particles, while the backscatter related Ångström exponent varied between 0.5 and 2.4. Knowledge of the optical properties of smoke in Southeastern Europe is very important for climatic studies, taking also into account the increasing trend of forest fires during the last decade. It was found that the optical characteristics of smoke aerosol were variable and this behaviour was mainly attributed to the fact that the burned regions are located in variable distances from the lidar station of Thessaloniki which affects the optical characteristics of the advected observed smoke aerosols due to different ageing processes. To justify this assumption we compared the backscatter-related Ångström exponent, which is mainly depended on particle size, with dispersion model estimations of the mean age of the carbon monoxide produced by the fires and advected over our site. Our results show that particle size is likely to increase with the age of the advected air mass. The increase of smoke particle size with age has fundamental implications for climate modeling and could be combined with modeling efforts

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to better constrain the radiative effect of biomass burning aerosols on regional/global scales. Finally, the correlation of the lidar ratio with particle size found in this study is considered valuable information to be used by backscatter lidar inversion algorithms in case of smoke presence (e.g. for space-borne lidars such as CALIPSO).

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Table 1. Dates of smoke presence over Thessaloniki, as indicated by the synergy of lidar measurements and dispersion modeling. The source of smoke particles is denoted along with the height range of smoke layers and the lidar derived optical parameters, namely the AOD, the mean lidar ratio and the mean backscatter related Ångström exponent.

Date	Source	Height range [km]	AOD	LR [sr]	Ångström
12 Jul 2001	Russia	2.12–3.83	0.47	86±2	1.39±0.02
16 Jul 2001	Russia	1.28–0	0.25	71±8	1.57±0.13
9 Aug 2001	Ukraine	1.67–3.47	0.74	56±1	1.71±0.26
16 Aug 2001	Ukraine	1.64–4.22	0.71	78±14	1.41±0.08
20 Aug 2001	Russia	2.42–4.73	0.31	50±1	1.80±0.06
8 Jul 2002	Russia	2.45–3.77	0.20	76±3	1.76±0.05
22 Aug 2002	Ukraine	2.63–4.10	0.20	39±12	2.20±0.21
28 Jul 2005	Russia	1.67–3.14	0.52	69±12	1.98±0.15
1 Aug 2005	Russia	2.21–3.17	0.22	94±5	0.78±0.25
12 Sep 2005	Russia	2.06–3.56	0.28	53±6	1.29±0.22

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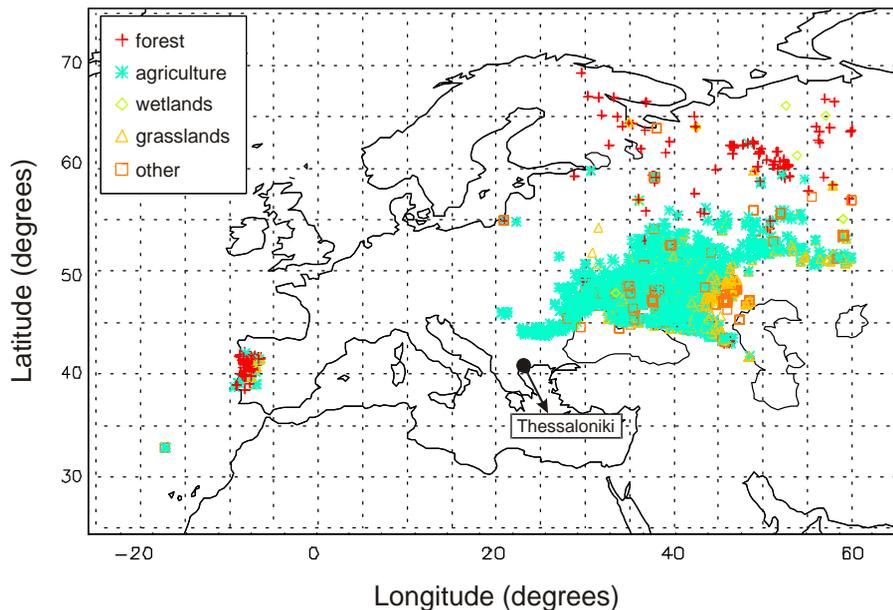


Fig. 1. Location of fires for the ten days of lidar measurements over Thessaloniki as detected by ATSR. Symbols indicate the dominant land cover where the detection occurred.

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Column-integrated emission sensitivity in global domain for Thessaloniki_010820.

Start time of sampling 20010820.180000 End time of sampling 20010820.180000

Upper release height 3000 m Lower release height 2500 m

Meteorological data used are from ECMWF

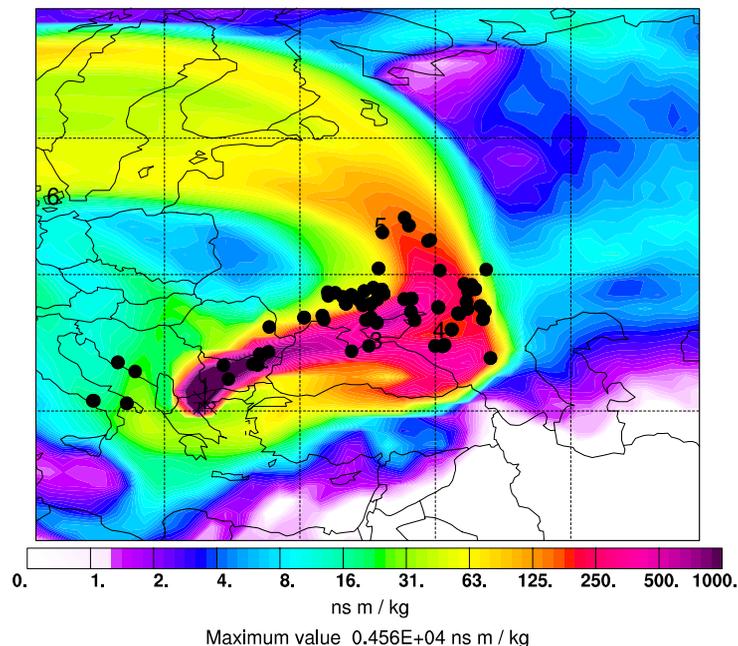


Fig. 2. Emission sensitivity obtained from the 20-day backward simulation with FLEXPART for the layer between 2500 and 3000 m on the 20 August 2001 at 18:00 UTC. The column vertical emission sensitivity is shown. Numbers represent the days backward in time as plotted at the retroplume centroid location. Hot spots detected by ATSR are superimposed (black circles) only in grid cells where the daily column-integrated emission sensitivity on the very day of the hot spot identification is above 8 ns m/kg, indicating the significant possibility of smoke advection.

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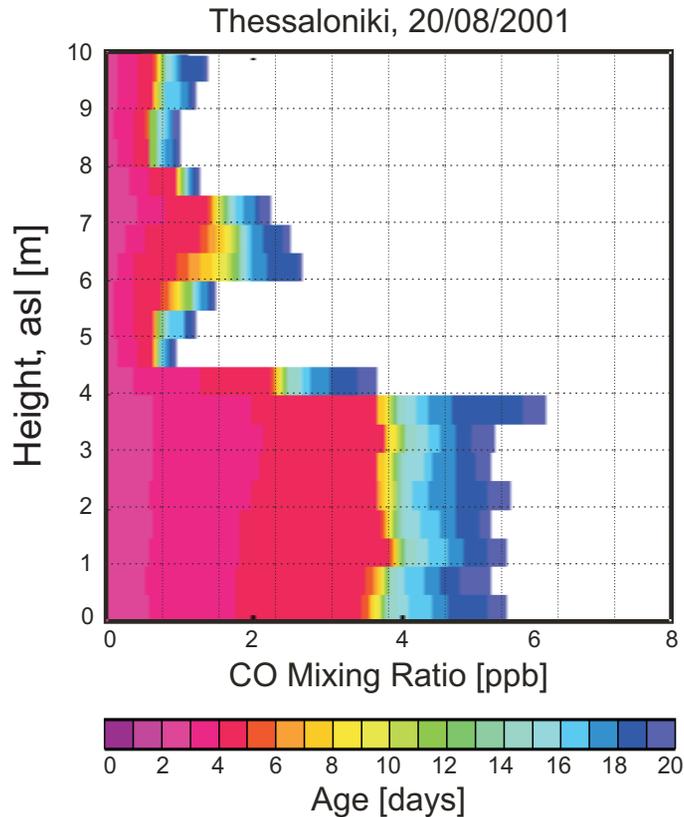


Fig. 3. Carbon monoxide concentration vertical profile over Thessaloniki for 20 August 2001 at 18:00 UTC, from FLEXPART simulations. Model calculations were performed considering CO emissions only from fire hotspots detected by ATSR. The age of CO is indicated by the color bar.

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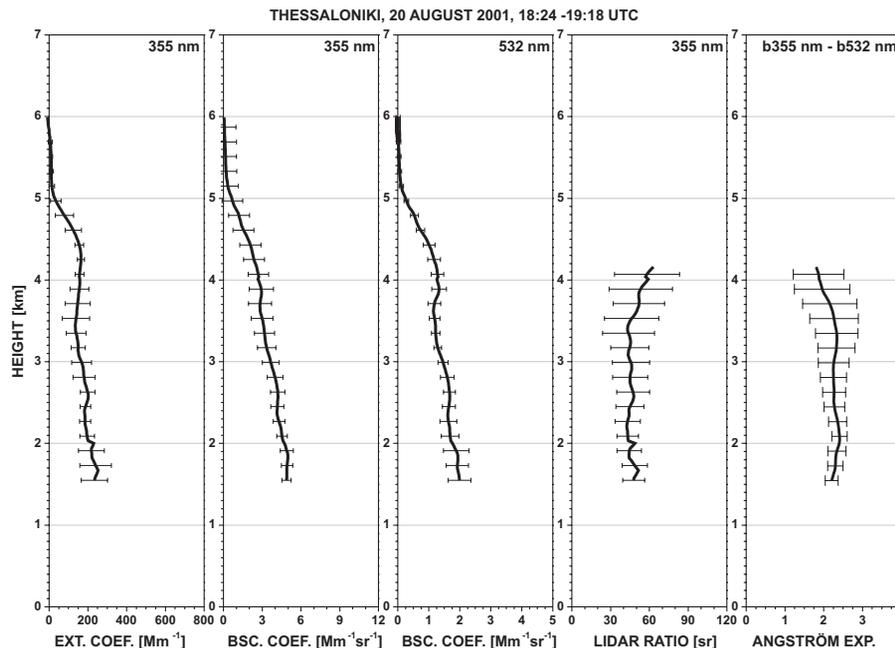


Fig. 4. Lidar derived vertical profiles of the extinction coefficient at 355 nm, the backscatter coefficients at 355 and 532 nm along with the lidar ratio at 355 nm and the backscatter-related Ångström exponent measured at Thessaloniki on 20 August 2001, between 18:24 and 19:18 UTC.

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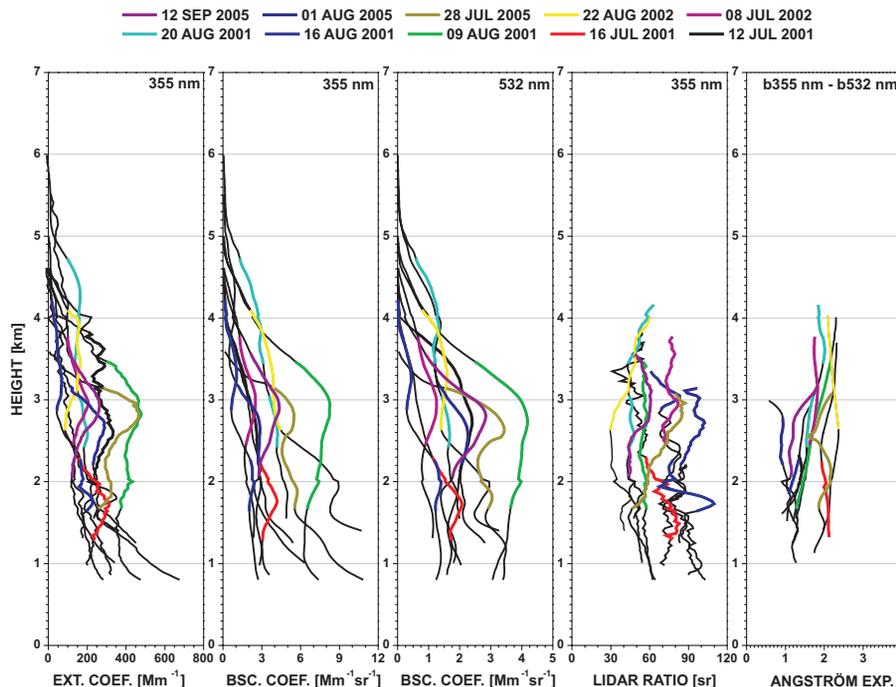


Fig. 5. Lidar derived vertical profiles of the extinction coefficient at 355 nm, the backscatter coefficients at 355 and 532 nm along with the lidar ratio at 355 nm and the backscatter-related Ångström exponent measured at Thessaloniki for ten days that smoke presence was detected. Black lines correspond to the complete vertical profile measured for each day while height ranges indicated with different colour correspond to the smoke layers.

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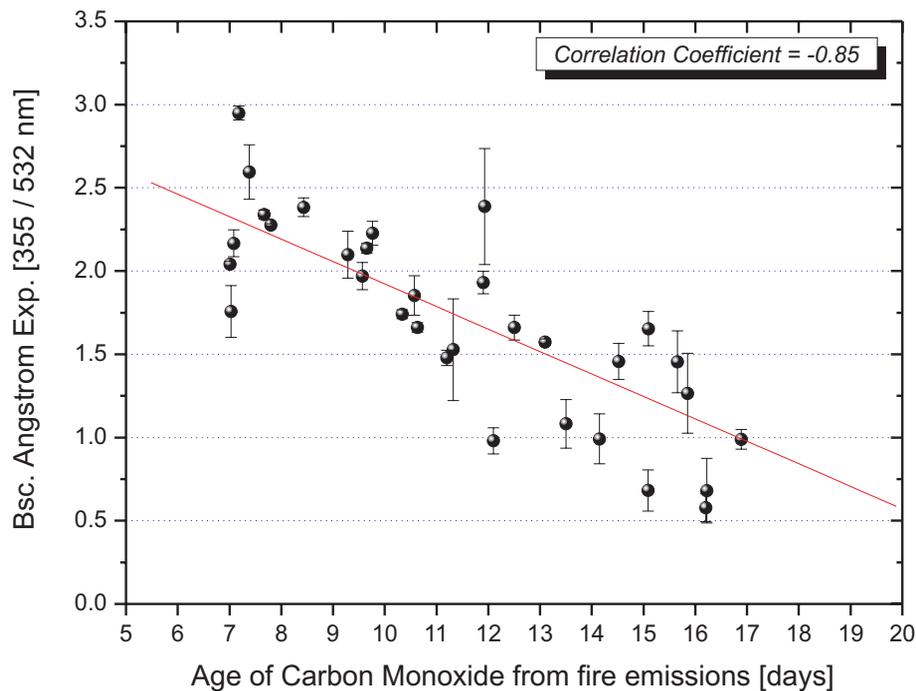


Fig. 6. Vertical mean values of the backscatter related Ångström exponent calculated from the backscatter coefficients at 355 and 532 nm versus weighted mean age of the carbon monoxide tracer, calculated for height ranges of smoke presence.

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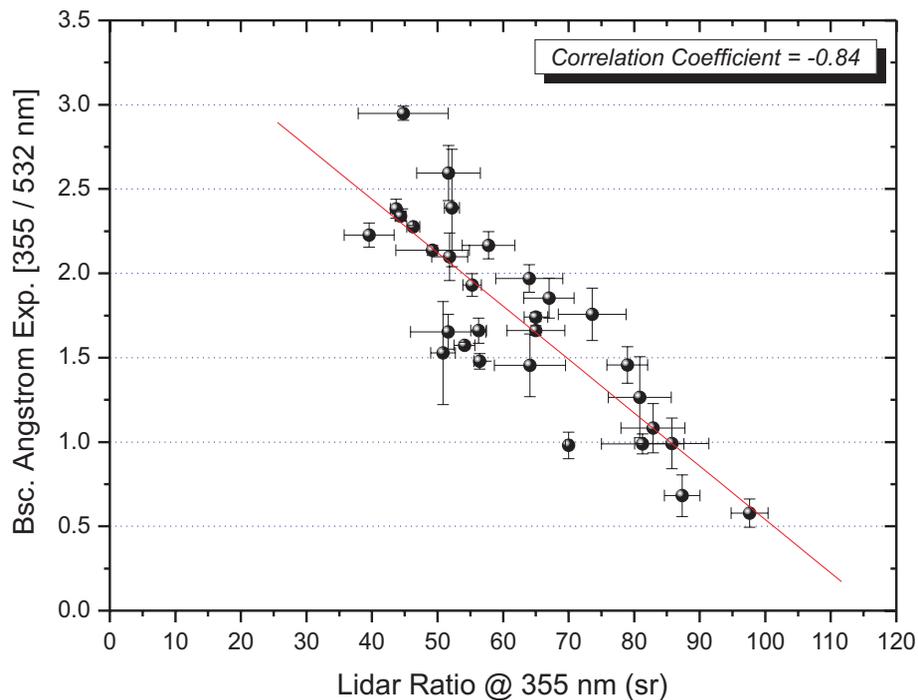


Fig. 7. Vertical mean values of the backscatter related Ångström exponent calculated from the backscatter coefficients at 355 and 532 nm versus vertical mean values of the lidar ratio at 355 nm, calculated for the same height ranges of smoke presence.

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