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Seasonal distribution of aerosol properties over Europe

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Seasonal distribution of aerosol properties over Europe and their impact on UV irradiance

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Abstract

Using the aerosol optical thickness at 550 nm (τ_{550}) from MODIS (collection 5) combined with the aerosol products from the ground-based AERONET network, key aerosol parameters have been obtained with 1 degree resolution over Europe. Additional tests have revealed a satisfactory quality of the MODIS data, except in a few cases. Quality assured AERONET data are used for evaluating the Angstrom exponent, single scattering albedo and asymmetry factor, and for validating the final aerosol optical thickness in the UV spectral region. A method for extrapolating the aerosol parameters into the UV spectral region is discussed. The aerosol optical thickness distributions are considered together with meteorological fields from NOAA_NCEP_CPC_CAMS_OPI climatology. The τ_{340} is shown to vary significantly from approximately 0.01 to 0.9 depending on the season and location. Permanent elevated aerosol loading over several industrial areas is observed, which agrees with the output of chemical transport models. Using radiative transfer modeling, monthly mean UV loss due to aerosol was estimated. The absolute decrease in UV indices varies from less than 0.1 to 1.5. The relative UV attenuation has large spatial and temporal variations (from -1% to -17%) with a minimum towards the northwest and maxima over several southern local areas (Northern Italy, etc.) during the warm period.

1 Introduction

Atmospheric aerosols are one of the important factors influencing solar UV irradiance, which play an important role in the biosphere and have a strong impact on human health (Van der Leun et al., 1998). Therefore, it is necessary to have reliable spatial and temporal distribution of aerosol parameters to get proper UV estimates. The information on aerosol is also necessary for correcting satellite UV retrievals, which can overestimate the measurements up to 40% (McKenzie, et al., 2002; Chubarova et al., 2005). The objective of this study is to develop the aerosol climatology over Europe

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and to characterize the aerosol effects on erythemally-weighted UV irradiance (Q_e) reaching the ground.

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Aerosol properties of the atmosphere can be obtained from both space and ground. At present several aerosol products are available from different satellite sensors (i.e. 5 AVHRR, OMI, MODIS, CERES, AATSR, MERIS, GLAS, SeaWiFs, MISR) as well as from ground-based aerosol networks (AERONET/PHOTONS, SKYNET, AEROSIB-NET, GLOBE, etc.) (IPCC, 2007). However, it is necessary to emphasize that the development of the aerosol climatology requires long-term period of observations and should have good spatial resolution. Unfortunately, only several aerosol datasets at 10 present can satisfy these conditions. In addition, since most of aerosol measurements are being carried out in visible spectral range, another problem can arise, how to extrapolate the aerosol parameters to UV spectral region. It should be also noted that recently a new aerosol dataset from OMI satellite instrument has been developed providing the estimation of aerosol parameters directly in the UV spectral region (Torres et al., 2002). However, not very long period of observations (since 2004) and current 15 problems with cloud screening procedures over Europe with prevailing cloudy conditions do not allow us to apply these records in our study.

The approach used in this study is based on the application of the data from satellite measurements of τ_λ at 550 nm (τ_{550}) from moderate resolution imaging spectrometer (MODIS) (collection 5) combined with the aerosol products from ground-based aerosol network AERONET/PHOTONS over Europe. According to (Remer et al., 20 2008), the latest MODIS dataset (collection 5) has a good quality over both land and ocean, except bright surfaces (snow and desert) and is available since 2000. The AERONET/PHOTONS network (<http://aeronet.gsfc.nasa.gov/>) (Holben et al., 1998) has been in operation since the middle of 1990s and has quite satisfactory spatial 25 coverage over Europe. Accurate multi-channel measurements by CIMEL sun/sky photometer through UV to near-infrared spectral region provide the data for evaluating a spectral dependence of aerosol optical thickness as well as many other inversion products including single scattering albedo and asymmetry factor (Dubovik and King,

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2000). In addition, AERONET/PHOTONS measurements of aerosol optical thickness at 340 nm can be used for direct validation of the final aerosol product in UV spectral region.

Since both satellite and ground-based measurements refer to the middle 1990s–beginning of 2000s period, the influence of Pinatubo volcanic eruption on the increase of total aerosol loading can be neglected. At the same time, we should emphasize the existence of negative trends in aerosol optical thickness of about $-0.04\text{--}0.05$ per decade (at 500 nm), which have been obtained over some areas in Europe (Ruckstuhl et al., 2008), and over the former Soviet Union since the 1980s (Gorbarenko et al., 2006; Makhotkina et al., 2005). Their nature is determined by both anthropogenic and natural processes and will not be discussed in this paper. However, one should take these trends into account while applying the results to another time period.

2 Data and method description

For estimating the aerosol impact on UV irradiance it is necessary to know several key aerosol characteristics: the aerosol optical thickness (τ_λ), the single scattering albedo (ω_λ), which characterizes the ratio of scattering to scattering plus absorption, and asymmetry factor (g_λ), which can be written in the form:

$$g_\lambda = \frac{\int_{-1}^1 \cos \theta \cdot P(\theta) d(\cos \theta)}{\int_{-1}^1 P(\theta) d(\cos \theta)} \quad (1)$$

where θ is the scattering angle, $P(\theta)$ is the phase function.

Spectral dependence of aerosol optical thickness is usually defined by Angstrom parameter α , which is estimated as follows:

$$\alpha = -d \ln(\tau_\lambda) / d \ln \lambda, \quad (2)$$

The monthly mean MODIS τ_{550} have been taken from <http://disc.sci.gsfc.nasa.gov/giovanni/> for the 2000–2008 period with 1° spatial resolution over the area of $30\text{--}80^\circ \text{N}$,

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28° W–40° E, which covers Europe and nearby territories. The measurements from both TERRA and AQUA satellite platforms were used to increase the statistics and to account possible aerosol changes during daytime.

The ground-based AERONET/PHOTONS quality assured measurements at level 2 (version 2) were used over the 1994–2006 period for evaluating the Angstrom parameter α to account spectral τ_λ dependence and for estimating the retrievals of single scattering albedo and asymmetry factor (Dubovik and King, 2000; Dubovik et al., 2002). Figure 1 presents the spatial distribution of Cimel sun/sky photometers, which data were used in the analysis. To exclude the systematical changes of aerosol parameters with surface elevation, only the data from the sites located lower than 1000 m were processed. Since overcast cloudy conditions prevent direct sun measurements, the total number of stations with available datasets varies from 73 in summer to 60 in winter. It should be mentioned that the number of Cimel sun/sky photometers with the UV channels is much smaller: the τ_{340} measurements are available from 39 sites in June–July and only from 31 in December (see Fig. 1).

Although the quality of MODIS data may be considered as satisfactory, several additional tests have been fulfilled. Figure 2 presents the difference in τ_{550} and in τ_{340} between satellite and ground-based datasets. The average uncertainty of monthly mean satellite τ_{550} retrievals lies within ± 0.02 – 0.03 with small but statistically significant difference in the fall. However, the difference is not large and is much less than τ_{550} seasonal changes over all European regions. To get τ_λ estimates in the UV spectral range two approaches were used: by accounting for spectrally-dependent Angstrom parameter, which can be observed in some conditions (Eck et al., 1999; Dubovik et al., 2002), and with the application of a constant α value retrieved from 440–870 nm spectral range. The comparisons between the τ_{340} estimated from MODIS data using both methods and direct ground τ_{340} CIMEL measurements have revealed the same character of the difference, though with higher deviation (± 0.05) for the second one. The difference between the two methods is small and lies within 0.01–0.03 that is close to the uncertainty of τ_{340} estimates. At the same time, due to the much smaller number

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of CIMEL instruments with UV channels (see Fig. 1) the advantages of accounting for spectral Angstrom curvature have been outweighed by the much lower statistics and, hence, larger uncertainty in the resulting Angstrom parameter spatial distribution. Finally, the application of the constant $\alpha_{440-870}$ values was chosen.

5 The additional analysis has revealed the presence of unrealistically high τ_{550} values (up to ~ 0.7 –0.8) observed over some regions in the northern Arctic regions, probably, due to the existence of small spots of snow or ice within the pixel. Note, that according to the measurements at AERONET/PHOTONS sites located at high latitudes ($>58^\circ$ N), the average τ_{550} values are about 0.1 with the maximum monthly mean value, not exceeding 0.2. This threshold ($\tau_{550}=0.2$) has been chosen as an additional filter, and has been applied to the MODIS data over the northern regions $>60^\circ$ N to remove unrealistically high τ_{550} .

As a result, the τ_{550} climatology over Europe has been obtained (Fig. 3), which is used as a base for developing the UV aerosol climatology. Spatial distributions of 15 Angstrom parameter $\alpha_{440-870}$ have been estimated using geostatistical ordinary Kriging method with a linear type of variogram model. The results are shown in Fig. 4. Using these distributions the final UV aerosol (τ_{340}) climatology has been generated (Fig. 5).

The final distributions of aerosol parameters over Europe are analyzed together 20 with the spatial distribution of meteorological parameters. Near-surface wind speed is known to determine the source strength for several aerosols types (for example, sea salt, dust, primary organic particles) (IPCC, 2007). Wet removal of atmospheric aerosols is affected by precipitation, which can also change the soil moisture and vegetation, and, hence, generation of dust particles and biogenic emission. In order to qualify 25 the possible reasons of aerosol optical thickness changes the data from IRI/LDEO Climate Data Library (<http://iridl.ideo.columbia.edu/>) were used to produce maps over Europe with precipitation and wind fields at 925 hPa for the period 1961–1990 on the base of NOAA NCEP CPC CAMS_OPI climatology (Janowiak and Xie, 1999; Kalnay et al., 1996). The monthly maps with meteorological fields are presented in Fig. 6.

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3 Results

3.1 Distribution of aerosol optical thickness and Angstrom parameter over Europe

The monthly mean distributions of aerosol optical thickness and Angstrom parameter shown in Figs. 3–5 have some typical features in their seasonal and temporal variations. The changes in aerosol optical thickness have a tendency to increase towards the south-east, inward the continent, though with different gradients over the year. The τ_{340} values vary from ~ 0.01 over northern and north-western areas during cold period to $\tau_{340} \sim 0.9$ over the south, south-eastern regions during spring and summer. One should note very pronounced seasonal τ_λ changes, which are sometimes higher than their difference in space. The Angstrom parameter also increases towards the south-east and has noticeable seasonal changes with maximum in summer over continental area at the south-east and minimum over Atlantic ocean in winter. Figure 7 presents the examples of different seasonal τ_{340} distributions from both direct AERONET/PHOTONS measurements and the developed MODIS/AERONET climatology in the arctic (1), temperate (2), and subtropical (3) climate zones and several of their subtypes. The lowest τ_λ are observed in the arctic climate zone with the spring maximum which, however, does not exceed $\tau_{340} \sim 0.23$. Within the temperate climate zone two subtypes can be found over Europe: the area with uniform humidification with one summer τ_λ maximum (the Oostende site) and the area transitional to continental type, located approximately eastward of $\sim 15^\circ$ E and northward of $\sim 45^\circ$ N with two τ_λ maxima in spring and summer like at Gotland, Toravere and Moscow sites. One can see much higher τ_{340} variations there with maxima reaching ~ 0.4 – 0.5 . In the subtropical zone there is a distinct τ_λ increase towards inner continental areas from Spain to Greece with maxima observed during warm period. At the same time the characteristics of the changes in τ_λ are much more irregular over subtropics, which can be attributed to the effects of the mineral dust aerosol advection to different Mediterranean regions from the surrounding deserts. It is necessary to note, that at some sites (for

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example, in Thessaloniki) the long-term τ_λ dataset obtained from Brewer measurements provides much better agreement with MODIS data than that obtained between MODIS and AERONET dataset, possibly, due to larger statistics and the application of additional clear-sky filter (Kazadzis, 2007; Koukouli et al., 2007)¹.

5 The analysis of monthly changes in Angstrom parameter over Europe, shown in Fig. 4, has revealed its substantial growth towards the inner continental regions during warm period with the high values ($\alpha > 1.4$) in June–July. This can be explained by relative decrease of sea salt aerosols and additional generation of fine mode particulate matter from different biogenic sources in conditions of high photolysis rates, elevated
10 temperatures and increase of aerosol VOC precursors. During spring and summer one can see a substantial increase of α values even over the Atlantic area that can be explained, to some extent, by the formation of fine mode sulphate aerosol from photochemical reactions with dimethyl sulfide (DMS) from marine plankton (Andreae and Crutzen, 1997; Denman et al., 2007). Another source of fine-mode aerosol can be
15 the advection of continental aerosol from the eastern coast of North America, which enhances both aerosol optical thickness and Angstrom parameter over Atlantic ocean. The back trajectory analysis described in (Smirnov et al., 2000) also confirmed that air originating above the continent was responsible for the increase in aerosol optical thickness and Angstrom parameter in remote Atlantic area during the TARFOX. In (Smirnov
20 et al., 2002) it was also mentioned the noticeable effects of different kinds of fine mode continental aerosol in Atlantic ocean. The increase of these aerosol parameters near the European coast can be also explained by the diffusion of polluted air directly from the European continent. All these assumptions should be studied further. During cold period the Angstrom parameter has a distinct decrease over almost the whole Europe

¹ It should be noted that the data from the developed MODIS/AERONET climatology have been interpolated against the coordinates of the AERONET sites with one degree resolution dataset. Therefore, this picture should be considered not as the example of the validation but as a presentation of typical τ_{340} seasonal features in different areas of Europe obtained both from satellite and ground.

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due to significant enhancing of westerlies and precipitation (see Fig. 6), which result in the growth of the coarse mode sea salt aerosol concentration. However, even in cold period there is a tendency of increasing the fine mode aerosol towards inner continental regions at the south. A local α maximum at the south-eastern part of Europe over Balkan peninsula throughout the year can be explained by the joint effect of meteorological factors such as the lack of precipitation and attenuation in wet removal of aerosol, some specific features in air advection (see the discussion below).

Figure 5 demonstrates a permanent elevated aerosol loading due to pollution effects over several local areas: Northern Italy, the coast of Netherlands, the southern regions of Poland, and, especially in summer, over the lowlands of the mid and low Danube areas. The qualitative analysis of distribution of fine mode particles $PM_{2.5}$ obtained from chemical transport model LOTOS-EUROS (Schaap et al., 2008) for 2000 (see <http://home.planet.nl/~scha1378/tno/promote/>) has also confirmed the existence of local maxima approximately over the same regions. The maximum monthly mean value can reach $\tau_{340} \sim 0.9$ ($\tau_{550} \sim 0.45$) over Northern Italy during spring-summer months, while τ_{340} is about 0.5 ($\tau_{550} \sim 0.25$) over nearby unpolluted area. In other polluted regions the differences are not so large and are about $\Delta\tau_{340} \sim 0.05\text{--}0.2$ ($\Delta\tau_{550} \sim 0.03\text{--}0.1$). Direct comparisons of simultaneous measurements obtained by two collocated CIMEL sun/sky photometers in Moscow region (Meteorological Observatory of Moscow State University and Zvenigorod sites) have revealed an additional aerosol loading over Moscow megalopolis of about $\Delta\tau_{340} \sim 0.02\text{--}0.12$ ($\Delta\tau_{500} \sim 0.01\text{--}0.06$) (Chubarova and Sviridenkov, 2008), which is in agreement with the MODIS estimates over the Moscow region. This analysis has also confirmed the ability of MODIS retrieval algorithm to produce high quality data and to reveal the areas with high local pollution.

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3.2 Month-to-month variability of aerosol optical thickness and Angstrom parameter

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ates favorable conditions for aerosol accumulation, and, hence, for the increase of the aerosol loading over this area, compared with that over Western Europe and, especially, over the Atlantic coast. However, one can see a slight reduce of aerosol loading over the regions located to the east of 30° E due to the presence of snow cover there.

5 The local τ_λ maxima over the polluted areas described above are getting much more pronounced with $\tau_{340} \sim 0.6\text{--}0.8$ ($\tau_{550} \sim 0.3\text{--}0.4$). In *April* there is still the same type of atmospheric circulation over Eastern Europe with the domination of south-eastern winds from the steppe and desert areas, which is characterized by the lack of precipitation. The beginning of agricultural activity, accompanied usually by the prescribed fires in
10 the Eastern Europe, also creates additional source of fine-mode aerosol. Altogether these processes generate a distinct seasonal τ_λ maximum over Eastern Europe. One can also see an increase of the square with high Angstrom parameter ($\alpha > 1.4$) over the continental areas in Eastern and Central Europe. Low precipitation and the advection from Central European areas create favorable conditions to the accumulation of aerosol
15 over Northern Europe and nearby North Sea area. The snow cover, which is still existed over this area, also reduces the uptake of aerosol by ground and contributes to the increase of the aerosol lifetime in the atmosphere, thus creating a seasonal maximum in aerosol optical thickness over Northern Europe. Western Europe is under the influence of the nearby Atlantic ocean, which is characterized by higher precipitation
20 leading to active wet removal of aerosol. The location of the area with large precipitation over Western Europe is in a good agreement with the lower aerosol level with the approximate border at 10° E. In *May* there are pronounced changes in the circulation processes over Europe with amplifying the Azore anticyclone and spreading its wedges over the Mediterranean area. This is accompanied by the decrease in precipitation
25 over this area and, hence, by the accumulation of the aerosol particles. Over the northern areas one can see high aerosol loading due to the lack of precipitation and still snow surface. This April-May maximum in aerosol optical thickness agrees well with the results of long-term measurements of air transparency over Russian Arctic regions (Makhotkina et al., 2005). At the same time, the aerosol loading over Central and

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Eastern Europe reduces due to the increase in precipitation, the domination of northern air advection from Scandinavian regions and more intensive uptake of aerosol by grass and leaves. In *June* further increase of precipitation over the northern Eastern Europe and ceasing the air advection from the south are responsible for distinct local minimum in aerosol optical thickness over the vast territory to the north of 45° N and to the east of 15° E. In addition, this local τ_{λ} minimum can be attributed to comparatively high water store in soil and vegetation, which can also prevent active mineral dust aerosol formation. At the same time, at the south and, especially, at the south-eastern areas (the eastern Mediterranean and western region of the Black sea) due to the lack of precipitation the processes of aerosol accumulation are observed. In addition, the air advection from the east (from Asia Minor) provides an additional mineral dust aerosol (see Fig. 6), that was also described in (Kazadsis et al., 2007). In addition, elevated temperatures and high photolysis rates in the presence of volatile organic precursors create favorable conditions for active generating there different kinds of fine mode biogenic and anthropogenic aerosols. We can see the increase in the area with fine mode particles over the whole Europe, except its western and the shores of the Mediterranean Sea, where the influence of the coarse mode sea-salt and mineral dust aerosols from deserts are dominating. As a result, the yearly maximum of $\tau_{340} \sim 0.9$ ($\tau_{550} \sim 0.4$), which is the highest value recorded over Europe, is observed over Northern Italy in June. In *July* aerosol optical thickness is very high over the Mediterranean area and nearby territories, especially over Balkan area: Northern Greece, Bulgaria, Romania (the lowlands of the mid and low Danube areas) which can be characterized by additional generation of aerosol particles from anthropogenic emissions in conditions of high photolysis rates and elevated temperatures in the absence of wet removal of aerosol from the atmosphere. The local aerosol maxima over the polluted areas mentioned above are also observed. Like in June, one can see a dominating role of fine mode particles over the whole Europe, except areas nearby the Atlantic Ocean and the Mediterranean Sea. In *August* the high aerosol loading over southern Europe is still observed due to the influence of the Azore anticyclone, which is characterized by

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hot and dry weather. At the same time, over the northern areas (higher 60° N) one can see gradual decrease of aerosol optical thickness and Angstrom parameter. In summer, aerosol optical thickness undergoes the most strong spatial variability over Europe from the very low values over Skandinavia and Arctic ocean ($\tau_{340} \sim 0.1$, $\tau_{550} \sim 0.05$) to the very high ones over the Mediterranean south-eastern area of Europe ($\tau_{340} \sim 0.7$ –0.9, $\tau_{550} \sim 0.3$ –0.4).

In *September* lower temperatures and photolysis rates as well as the intensification of westerlies decrease the aerosol loading over the whole Europe area, except its south eastern part. The local maximum, which is observed over the central regions of Eastern Europe (Russia, Belarus, Poland, Moldova), has been generated due to the strong impact of extreme forest and peatbog fires observed in 2002 near Moscow (Chubarova et al., 2009). However, there is a high probability of such conditions due to the drainage of peatbogs and overall air temperature increase. In *October* further cleaning of the atmosphere is observed over the whole territory of Europe. This process is consistent with attenuation of the α values over the vast area with a distinct tendency to decreasing towards the Atlantic Ocean. There are still the centers over the south-eastern part of Europe with the high Angstrom parameter and the aerosol optical thickness. The location of the area with higher aerosol loading over central Europe (Czech Republic, Austria and nearby countries) corresponds well with the area with less precipitation. In *November* the aerosol optical thickness is quite low ($\tau_{340} \sim <0.2$, $\tau_{550} < 0.12$) over the whole European territory, except several local areas. The Angstrom parameter continues to decrease, that means further reduction of the square with the domination of the fine mode particles. Relatively high aerosol loading is observed over the Mediterranean area and, especially, over Northern Italy and the Balkan peninsula. The local area with high Angstrom parameter values ($\alpha > 1.4$) over the Balkan peninsula might be attributed to the joint effect of favorable conditions for generating the biogenic as well as anthropogenic aerosols from both natural and industrial emissions. In *December* the aerosol loading is the lowest in seasonal cycle due to the active removal of aerosol and the absence of its active generation in the atmosphere. However, the ele-

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vated aerosol optical thickness can be observed over several areas: the eastern part of the Mediterranean, which is under the influence of mineral dust advection from the deserts, over the lowlands of the mid and low Danube areas, over the industrial zones in Northern Italy, etc. The α values are also the lowest ones, and they are characterized by a pronounced increase towards the southeast. Unfortunately, it is impossible to retrieve the aerosol optical thickness over the large northern area due to bright snow conditions, where MODIS algorithm does not work properly. However, the data available from ground-based AERONET/PHOTONS network confirm the existence of low aerosol optical thickness over snow cover areas at the north.

Table 1 presents monthly mean and yearly mean τ_{340} over Europe averaged with $10 \times 10^\circ$ grid over 10° W– 40° E 30° – 80° N region. The data over Africa and Asia continents were excluded from the statistics. It summarizes the main features of seasonal behavior of aerosol optical thickness showed above.

3.3 Aerosol single scattering albedo and factor of asymmetry

Single scattering albedo and asymmetry factors are the important factors which is necessary to know for reliable UV estimates in aerosol atmosphere. In our approach they were retrieved through the special inversion algorithm (Dubovik et al., 2000) developed for sun/sky CIMEL photometers AERONET network. However, the method has some constraints. The most important ones are the existence of clear sky conditions during a series of observations and high aerosol loading ($\tau_{440} > 0.4$). As a result, during cold period over for the most of European territory there are no estimates of single scattering albedo due to low aerosol optical thickness and dominating of cloudy conditions. The ω_λ estimates are available only from few sites over southern regions (for example, only from 4 sites in December). Even in summer the statistics is quite poor: the data are available from approximately 40–50 sites, which monthly average values can be based sometimes only on 1–2 observations per month. The lack of statistics and the uncertainty of the ω_λ retrievals ($\Delta\omega = \pm 0.03$) do not allow us to generate spatial distributions of these parameters. Instead, a study of their month-to-month variability has

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been conducted. The month-to-month variations of both single scattering albedo and factor of asymmetry at 440 nm retrieved, using a standard AERONET algorithm, are shown in Fig. 8. No seasonal change in these characteristics could be detected at the 95% confidence level, except some small difference (about 0.01) in asymmetry factor values, which are higher in February, March and October. However, the difference is rather small and can be of random character that should be studied further. It should be emphasized that the absence of seasonal variations in single scattering albedo, which has been obtained, corresponds to the situations of high aerosol loading, that is the constraint for the ω retrieval procedure. At the same time, it is high aerosol loading that influences significantly the solar irradiance reaching the surface, while solar irradiance is not so sensitive to the changes in ω at small aerosol optical thickness. Therefore, these retrievals can be chosen in the first approach for estimating UV irradiance in Europe. As a result, yearly mean single scattering albedo at 440 nm over European region is about $\omega_{440}=0.93$ with standard deviation of 0.03 varying from 0.77 to 0.99. Asymmetry factor comprises $g_{440}=0.71$ with standard deviation of 0.02 varying from 0.58 to 0.79.

Then, it was necessary to make spectral adjustments and to re-evaluate single scattering albedo and asymmetry factor in UV range. This has been made with the application of Mie theory with accounting for the characteristics of volume particle size distribution and refractive indices at 440 nm, which in turn are the standard products of AERONET CIMEL sun/sky photometer network. The assumption of the same values of refractive indices in visible and UV spectra will not work properly in some cases, especially in smoke aerosol conditions (Chubarova et al., 2009). However, we can use this approximate approach due to the large statistics and the low frequency of fire smoke conditions in the overall dataset. As a result, the adjustment factors of $\Delta\omega\sim0.01$ and of $\Delta g\sim0.04$ have been obtained to provide the average estimates of $\omega\sim0.94$ and of $g\sim0.75$ in the UV region of the spectrum. These values were taken as constants within the whole UV range. In order to obtain more accurate retrievals of these aerosol parameters one should use additional spectral or UV-MFRSR measurements and follow

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more sophisticated methods described, for example, in (Bais et al., 2005; Krotkov et al., 2005). Unfortunately, at present there is no spread network of these measurements to give good spatial distribution of these parameters.

3.4 Validation against measurements

5 The examination of the proposed method for evaluation the UV single scattering albedo and factor of asymmetry has been made by comparing model UV estimates with the experimental data in clear sky conditions with different aerosol properties in Moscow. For this purpose a Troposphere Ultraviolet and Visible (TUV) radiative transfer model version 4.1 has been used (Madronich and Flocke, 1998) with 8 stream DISORT solver and pseudo spherical corrections.

10 The measurements of erythemally weighted irradiance (Q_e) in Moscow State University Meteorological Observatory (MO MSU) are being carried out by broadband YES UVB-1 pyranometer (Chubarova 2008). The instrument is regularly tested against the reference instrument, which in turn is regularly calibrated in erythemally-weighted units
15 against the ultraviolet spectroradiometer Bentham DTM-300 of the Medical University of Innsbruck in 1999, 2005, 2008 (courtesy of M. Blumthaler). The additional corrections on solar zenith angle and ozone content are applied following the WMO recommendations (Seckmeyer et al., 2006) and cosine characteristics of the instrument have been also taken into account. The clear sky conditions were chosen from hourly visual
20 observations. In addition, the account of typical NO_2 gas absorption, which is the most important absorber in Moscow conditions, has been considered in model calculations according to the available surface NO_2 measurements combined with its model profile.

25 The comparisons of model and measured Q_e at low aerosol optical thickness ($\tau_{340} < 0.1$), where the effects of uncertainty in single scattering albedo and factor of asymmetry are negligible, has revealed the existence of some model overestimation of about 6%. After taking this shift into consideration, we estimated the Q_e loss due to aerosol $L = (Q_{e_aerosol} - Q_{e_no_aerosol})/Q_{e_no_aerosol}$, where the $Q_{e_aerosol}$ values were taken both from the results of modelling and observations. The comparison between

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observed and modelled L values is shown in Fig. 9a for a wide range of aerosol optical thickness ($\tau_{340}=0.53\text{--}1.7$), and single scattering albedo ($\omega=0.8\text{--}0.97$) obtained from AERONET measurements according to the approach described above. One can see quite satisfactory agreement between modelled and measured Q_e aerosol loss with correlation coefficient $r=0.91$. Even in typical, non-smoke conditions the attenuation by aerosol can be very significant and exceed 50% due to comparatively small ω values and the episodes with high AOT loading.

In addition, using model calculations a simple equation can be obtained to quantify the dependence of relative Q_e loss on aerosol optical thickness (for $\tau_{340}<0.8$), single scattering albedo (for $\omega\geq0.8$), and airmass (for $m\leq2$):

$$\frac{\Delta Q_e}{Q_e} = \tau_{340} [(0.42m + 0.93)\omega + (-0.49m - 0.97)] \quad R^2 = 0.99 \quad (3)$$

The equation in the brackets is the attenuation coefficient, which characterizes the attenuation at $\tau_{340}=1$ and is a function of airmass m and single scattering albedo ω . For typical European summer conditions with $\omega\sim0.94$ and at $m=1.2$, and $\tau_{340}=0.4$ this attenuation coefficient is about -0.08 . The uncertainty of this equation is less than 1%, except the cases with high aerosol loading and low single scattering albedo, when the uncertainty can reach 2–3%.

The Fig. 9b presents the dependence of relative Q_e loss on aerosol optical thickness. The data were combined into two groups with different ω values in order to test the ability of the proposed method to reveal the effects of single scattering albedo. One can see quite pronounced distinction due to different ω in both observed and modelled L values and good agreement between modelled and measured Q_e loss. In conditions with relatively high single scattering albedo ($\omega\sim0.94$) the Q_e aerosol loss is typically less than 20%, while at $\omega\sim0.87$ it comprises about 25–30% at the same aerosol optical thickness. Thus, the AERONET inversion products with some adjustments can be used for the assessment of aerosol properties in UV range.

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3.5 UV changes due to aerosol

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The TUV model has been also applied for estimating the Q_e aerosol loss over Europe. Model simulations were fulfilled for noon conditions (minimum solar zenith angle) in aerosol and in aerosol free situations. In addition, seasonal and spatial changes in 5 total ozone content retrieved from TOMS ozone climatology dataset (<http://jwocky.gsfc.nasa.gov/>) have been also accounted.

Both relative and absolute changes of UV irradiance due to aerosol have been evaluated for central months of the seasons. The absolute changes due to aerosol are shown in UV indices (UVI), which are defined by simple multiplying the Q_e (W/m^2) by 10 40 (Vanicek et al., 2000). Figure 10a presents a significant variation of monthly mean Q_e relative losses over Europe due to aerosol loading, which lie within $\sim(-1)\text{--}(-17)\%$. Here, the Q_e attenuation is mainly regulated by aerosol optical thickness and has a 15 small negative dependence on airmass m (see the Eq. 3). As a result, the most significant Q_e loss of about $(-15)\text{--}(-17)\%$ was estimated over Northern Italy in spring and in autumn due to high aerosol loading and lower solar elevations compared with 20 summer conditions. The minimum Q_e attenuation ($>-3\%$) is observed over the North Atlantic ocean, Scandinavia in autumn and in winter, where the aerosol loading is small ($\tau_{340}\sim 0.1$) and over some other clean regions like, for example, western coast of Spain in January. On average, there is a tendency of increasing the Q_e loss towards inner continental regions, especially during warm period.

In winter, in spite of relatively small aerosol optical thickness, the Q_e attenuation over Europe is quite pronounced varying within $(-3)\text{--}(-10)\%$ and increases on 3–5% over urban polluted areas (i.e. the coast of Netherlands, south of Poland) and towards the south. It is necessary to note that due to relatively higher solar elevation 25 over the Mediterranean area, compared with North Europe, the loss of Q_e over the polluted Northern Italy is not so large. In spring, due to the increase in aerosol loading one can see the enhancement of Q_e loss over the whole Europe. The attenuation of Q_e varies within $(-5)\text{--}(-15)\%$ with a pronounced drop up to -15% over Central,

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Eastern and North-Eastern Europe. The distinct shifting of high Q_e loss to the north in spring, where the aerosol optical thickness is not so high, is also attributed to smaller solar elevations there. In summer, the maximum loss in Q_e reaching $(-10)\text{--}(-15)\%$ is observed at the southern and south-eastern European areas including several local 5 polluted regions like over the lowlands of the mid and low Danube areas, Northern Italy, south of Poland, etc. In autumn, there is a pronounced decrease of Q_e loss over the whole European territory, and, especially over the north Atlantic area, except several polluted areas and, especially, Northern Italy.

Figure 10b represents the spatial distribution of absolute attenuation in monthly 10 mean UVI due to aerosol over Europe. Since absolute UVI loss has a positive correlation with both aerosol optical thickness and solar elevation, one can see a different picture. Over the southern regions with high aerosol loading the absolute decrease of UVI is enhanced, while over northern areas, especially in winter, it is quite negligible in spite of their relative significant loss. As a result, in winter a typical absolute decrease 15 of UV indices due to aerosol varies from less than $\Delta\text{UVI}\sim-0.1$ over northern and central European territory to $\Delta\text{UVI}\sim-0.3$ over southern regions. However, significant changes in aerosol loading can provide large variations in UVI. For example, at the same latitude ($\sim 40^\circ\text{N}$) in spring the absolute UVI difference varies within the order of magnitude (from -0.1 to -1.0). In summer the absolute UVI loss can exceed 20 -1.0 over the south-eastern areas, where the aerosol loading is extremely high. The maximum effect of aerosol is observed over Northern Italy, where the difference in UV indices due to aerosol loading can reach 1.5. At high latitudes, where the absolute level of UV irradiance is not so high, the absolute changes in UVI also do not exceed 0.5. In autumn one can see typical decrease of less than 0.5 over the whole Europe, 25 except Northern Italy, where aerosol optical thickness is still high.

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4 Discussions and conclusions

In the paper we have shown that the combined MODIS/AERONET datasets can be used for the development of the UV aerosol climatology over Europe. The comparisons with direct ground-based CIMEL measurements at 340 nm have revealed the uncertainty of $\Delta\tau_{340}=\pm 0.05$, which is much less than the seasonal τ_{340} variations over the most territory of Europe².

The analysis of aerosol optical thickness and Angstrom parameter spatial distribution over Europe has been carried out together with the analysis of meteorological fields, which have revealed several interesting features in seasonal and spatial distribution:

1. Increasing of the aerosol optical thickness from the north-west towards south-east is observed with τ_{λ} maxima during warm period. The existence of spring aerosol maximum over northern arctic areas has been confirmed. Summer τ_{λ} maximum is typical for the western coastal zone of Europe as well as over the south and south-east European areas which, however, is much stronger. The bimodal seasonal distribution in aerosol optical thickness with spring and summer maxima and the pronounced drop of τ_{λ} in June refers to the large continental area to the north of $\sim 45^{\circ}$ N and to the east of $\sim 15^{\circ}$ E. Most clean areas with τ_{340} less than 0.1 are observed during cold period over north-western part of the Atlantic ocean, over Scandinavia, and some other local areas. The highest monthly mean aerosol optical thickness values (up to $\tau_{340} \sim 0.9$) have been obtained during warm period over several local areas mostly at the southern, south-eastern part of Europe.
2. There are noticeable seasonal changes in Angstrom parameter, which are characterized by its increase in summer, even over the Atlantic Ocean with a permanent local maximum over south-eastern part of Europe. These features can

²It should be noted that the same approach has been used for the evaluating of aerosol optical thickness climatology at 308 nm over Europe within the frame of COST726 project. The dataset can be found at the site <http://www.cost726.org>.

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be explained by joint effects of air advection, lack of precipitation and vegetation activity, which determines the photochemical generation of different types of fine mode aerosol.

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5 3. There have been revealed several local polluted centers with elevated aerosol optical thickness. The most significant ones are located in Northern Italy, over the coast of Netherlands, the southern regions of Poland (carboniferous basin, and different industries), and, in summer, over the lowlands of the mid and low Danube areas. These areas are consistent with the areas with high concentration of $PM_{2.5}$ particles obtained from chemical transport model LOTOS-EUROS.

10 10 The average estimates of single scattering albedo and asymmetry factor ($\omega \sim 0.94$, $g \sim 0.75$) have been obtained in UV spectral region on the base on AERONET/PHOTONS retrievals. No seasonal dependence has been revealed for these parameters. However, it should take into account the constraints of the inversion method, which can be only applied for large τ_λ .

15 15 Satisfactory agreement between UV measurements and model estimates in conditions with different aerosol properties has confirmed a possibility of the retrievals of single scattering albedo and asymmetry factor from AERONET inversion products and its extrapolation to UV range using the Mie theory in the absence of fire conditions.

20 20 The estimates of UV irradiance attenuation due to aerosol have been obtained for different seasons. It has been shown that the absolute loss of UV indices due to a positive correlation with both aerosol optical thickness and solar elevation is most pronounced over southern areas. The UVI varies from less than 0.1 over the northern areas in winter up to 1.5 over polluted areas at the south (Northern Italy). However, significant variations in UVI (within the order of magnitude) can be observed only due 25 to τ_λ variations, especially at the south of Europe.

The analysis of relative changes in Q_e due to aerosol has revealed large spatial and temporal variations within $\sim(-1)\text{--}(-17)\%$ with minimum Q_e attenuation ($>-3\%$) during cold period over the North Atlantic area, Scandinavia, etc. The most significant Q_e

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loss ((−15)–(−17)%)) was estimated over Northern Italy in spring and in autumn due to high aerosol loading and lower solar elevations compared with summer conditions. In winter the Q_e attenuation over Europe varies within 10% and increases on 3–5% over the polluted centers. In spring and summer Q_e attenuation can reach (−10)–(−15)% over the Central, Eastern and North-Eastern Europe. In autumn, there is a pronounced decrease in Q_e loss over the whole European territory, except Northern Italy.

5 Due to the existence of different dependence on solar elevation one can see a pronounced shift to the north for the relative Q_e attenuation and to the south for the absolute Q_e attenuation.

10 The effects of aerosol loading on UV irradiance can be compared with total ozone influence. The same aerosol loading at high latitudes can provide much stronger relative changes in Q_e while there is vice versa situation with total ozone, which has less effects on Q_e at low elevations (high latitudes) than at low latitudes (see, for example, Fioletov et al., 2001). In other words, in spring at high latitudes the potential UVI growth
15 due to the total ozone decrease can be much more effectively cancelled by the aerosol loading. In addition, at high latitudes the Q_e variations due to aerosol are much more pronounced and can significantly influence the Q_e year-to-year variability, that should be taken into account while revealing the ozone component in Q_e long-term trend.

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25 in this investigation. MODIS data were produced with the Giovanni online data system, developed and maintained by the NASA Goddard Earth Sciences (GES) Data and Information Services Center (DISC). The work has been also partially supported by the RSFBR grants #07-05-00860, and #09-05-00582.

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Table 1. Monthly mean and yearly mean τ_{340} over Europe according to the developed MODIS/AERONET climatology averaged with 10×10 grid.

Longitude, $^{\circ}$	Latitude, $^{\circ}$	1	2	3	4	5	6	7	8	9	10	11	12	Year
-5	35	0.14	0.22	0.22	0.25	0.28	0.35	0.34	0.34	0.3	0.18	0.14	0.13	0.24
-5	45	0.14	0.21	0.28	0.28	0.31	0.31	0.32	0.3	0.23	0.19	0.13	0.14	0.24
-5	55	0.12	0.17	0.31	0.3	0.33	0.27	0.28	0.25	0.19	0.13	0.12	0.06	0.25
-5	65	0.18	0.22	0.3	0.31	0.27	0.25	0.21	0.18	0.1				
-5	75		0.18	0.26	0.28	0.22	0.16	0.14	0.1					
5	35	0.17	0.26	0.28	0.38	0.37	0.43	0.47	0.43	0.37	0.28	0.2	0.18	0.32
5	45	0.16	0.25	0.32	0.34	0.35	0.41	0.38	0.33	0.24	0.21	0.13	0.13	0.27
5	55	0.16	0.21	0.32	0.37	0.35	0.36	0.36	0.35	0.27	0.18	0.15		
5	65	0.12	0.2	0.29	0.28	0.23	0.23	0.21	0.16	0.1				
5	75		0.2	0.26	0.3	0.24	0.17	0.14	0.11					
15	35	0.2	0.29	0.34	0.41	0.39	0.41	0.45	0.47	0.39	0.32	0.24	0.2	0.34
15	45	0.17	0.28	0.38	0.45	0.4	0.46	0.45	0.43	0.31	0.25	0.19	0.17	0.34
15	55	0.18	0.18	0.33	0.39	0.35	0.35	0.38	0.37	0.27	0.2	0.14		
15	65	0.09	0.18	0.24	0.23	0.17	0.17	0.15	0.11	0.09				
15	75		0.22	0.25	0.27	0.23	0.18	0.15	0.12					
25	35	0.2	0.31	0.39	0.45	0.38	0.4	0.47	0.46	0.4	0.34	0.26	0.21	0.36
25	45	0.18	0.28	0.37	0.42	0.34	0.43	0.53	0.51	0.36	0.21	0.18	0.16	0.33
25	55	0.24	0.24	0.37	0.43	0.35	0.24	0.37	0.37	0.33	0.18	0.14		
25	65	0.1	0.22	0.33	0.22	0.16	0.2	0.17	0.13	0.1				
25	75		0.24	0.28	0.33	0.21	0.2	0.16	0.12					
35	45	0.19	0.27	0.3	0.41	0.31	0.37	0.47	0.47	0.32	0.24	0.2	0.16	0.31
35	55	0.2	0.31	0.31	0.43	0.33	0.24	0.37	0.37	0.35	0.18	0.13		
35	65	0.04	0.21	0.35	0.27	0.17	0.25	0.21	0.16	0.16				
35	75		0.28	0.32	0.34	0.24	0.3	0.19	0.13					

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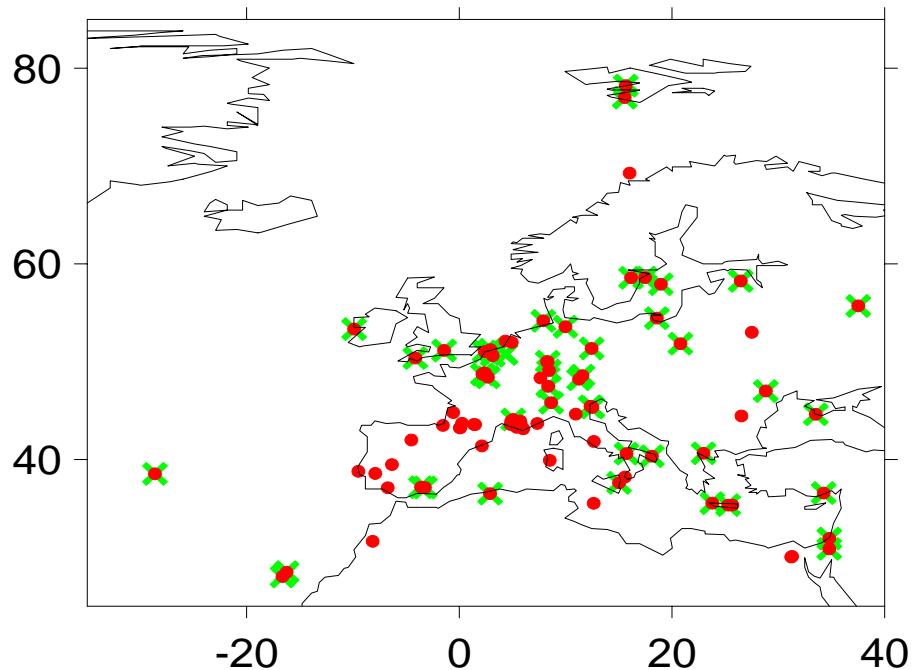


Fig. 1. Distribution of CIMEL sun/sky photometers in Europe and nearby areas. The sites, where the instrument has UV channels (340 and 380 nm), are marked by green crosses.

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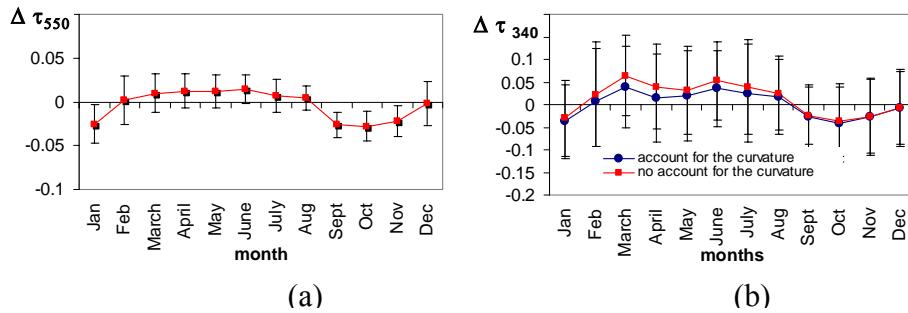


Fig. 2. The difference between MODIS and AERONET/PHOTONS aerosol optical thickness over Europe as a function of month at 550 nm **(a)** and at 340 nm **(b)**. The confidence intervals were obtained at 95% significant level. The difference with AERONET data at 340 nm is shown with the account and without account for spectral dependence of Angstrom parameter. See the details in the text.

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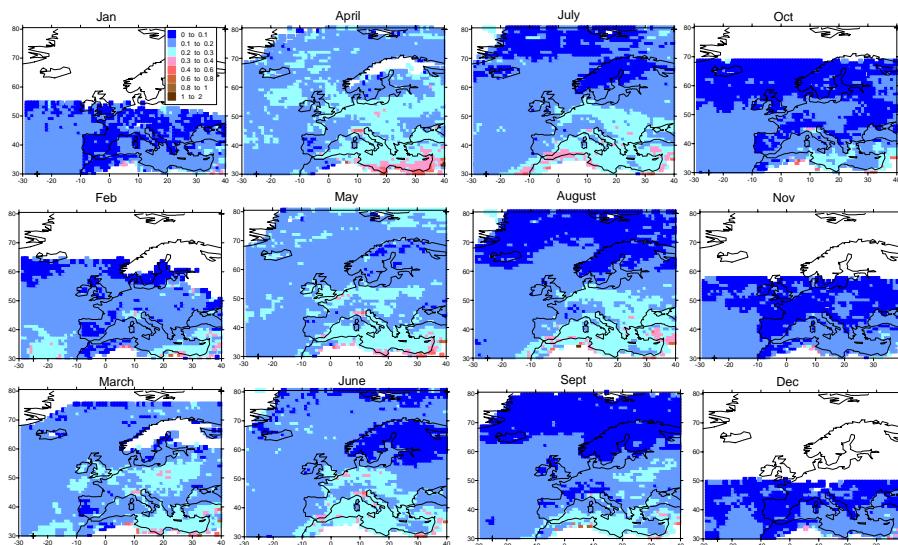


Fig. 3. Monthly spatial distribution of τ_{550} over Europe according to MODIS data with account of some corrections in high latitudes according to AERONET/PHOTONS dataset.

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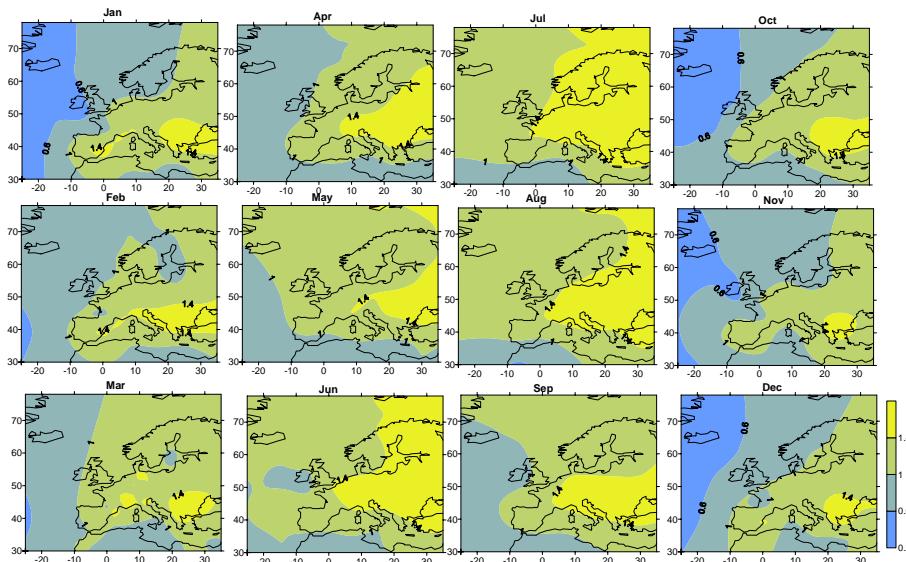


Fig. 4. Monthly spatial distribution of Angstrom parameter according to AERONET/PHOTONS dataset over Europe.

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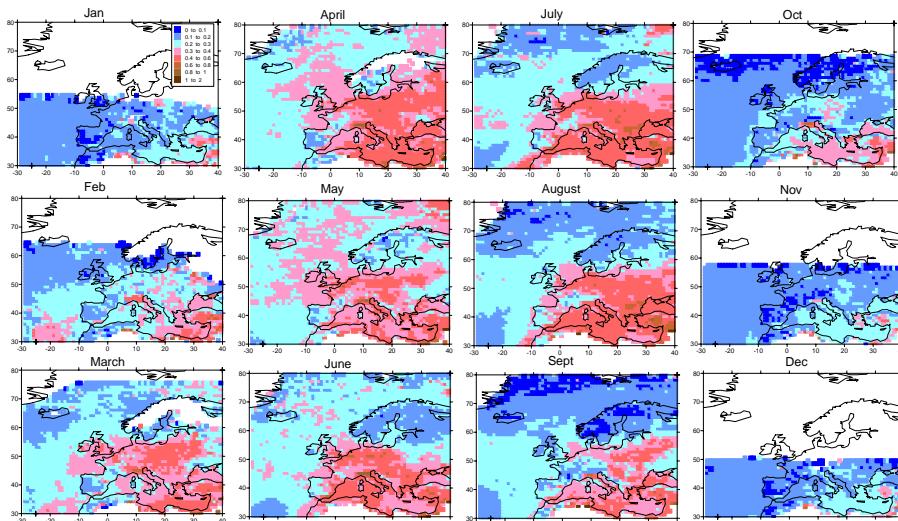


Fig. 5. Monthly spatial distribution of τ_{340} according to MODIS/AERONET data.

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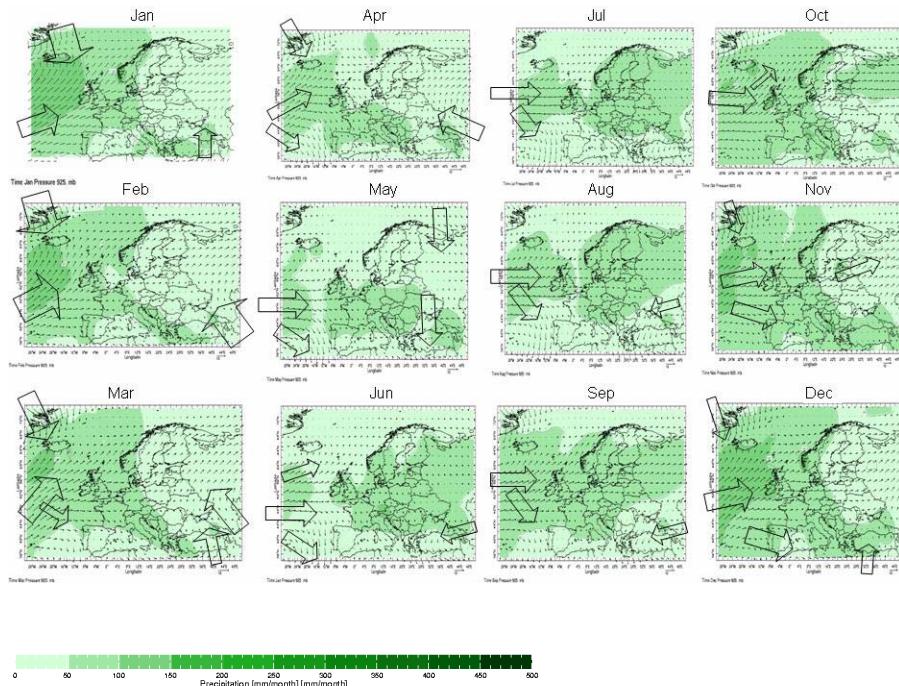


Fig. 6. Monthly spatial distribution of climate meteorological parameters (wind characteristics and precipitation) according to NCEP CPC CAMS_OPI data from the IRI/LDEO Climate Data Library 1961–1990.

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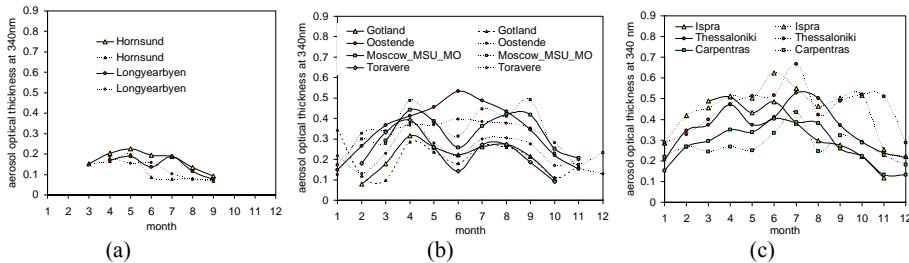


Fig. 7. Seasonal differences τ_{340} in the arctic **(a)**, temperate **(b)** and subtropical **(c)** climate zones. The MODIS τ_{340} retrievals are shown by solid lines and AERONET/PHOTONS data – by dotted lines. The coordinates of the sites: Hornsund (15.55° E, 77° N), Longyearbyen (15.649° E 78.2° N), Gotland (19.0° E 57.9° N), Moscow_MSU_MO (37.5° E, 55.7° N), Oostende (2.9° E 51.2° N), Toravere (26.5° E, 58.3° N), Ispra (8.6° E, 45.8° N), Carpentras (5.1° E, 44.1° N), Thessaloniki (23.0° N, 40.6° E).

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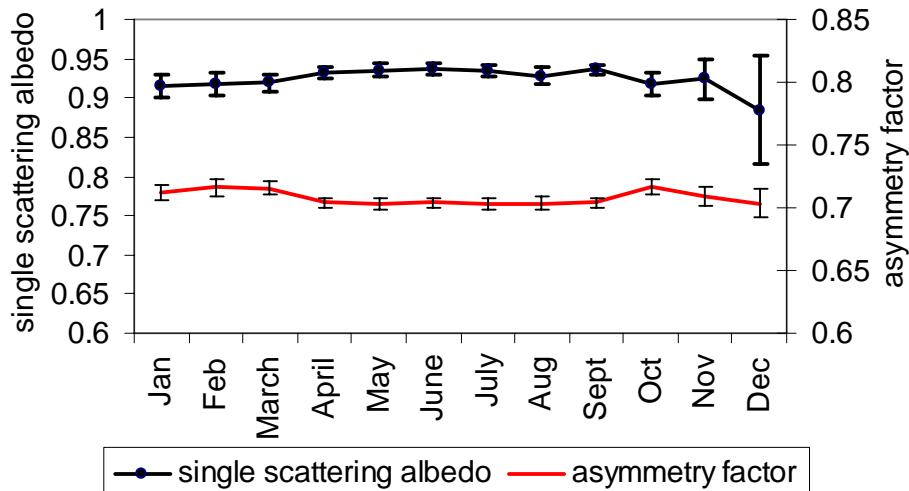


Fig. 8. Month-to-month variations of single scattering albedo (ω_{440}) and asymmetry factor (g_{440}) according to CIMEL sun/sky photometer measurements over Europe. The confidence intervals are shown at 95% level.

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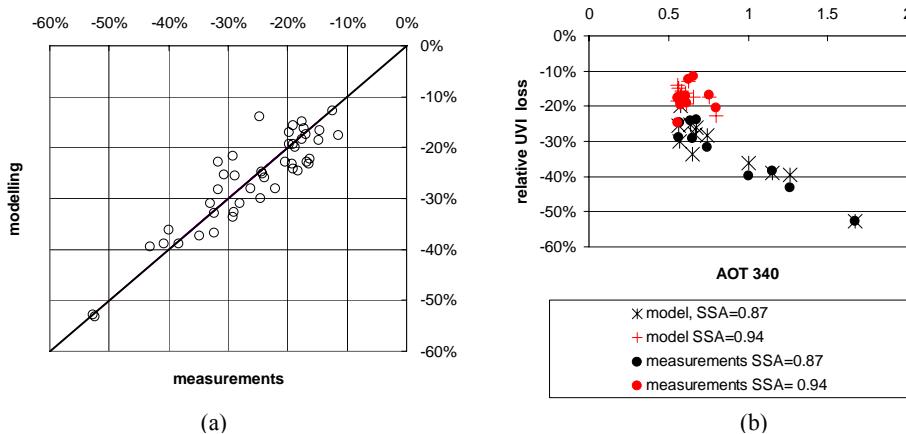


Fig. 9. The comparison between model and measured erythemally-weighted irradiance losses in clear sky conditions with single scattering albedo and factor of asymmetry parameters derived from the AERONET dataset **(a)** and relative UV loss as a function of aerosol optical thickness at 340 nm **(b)**. Moscow. Snow-free conditions.

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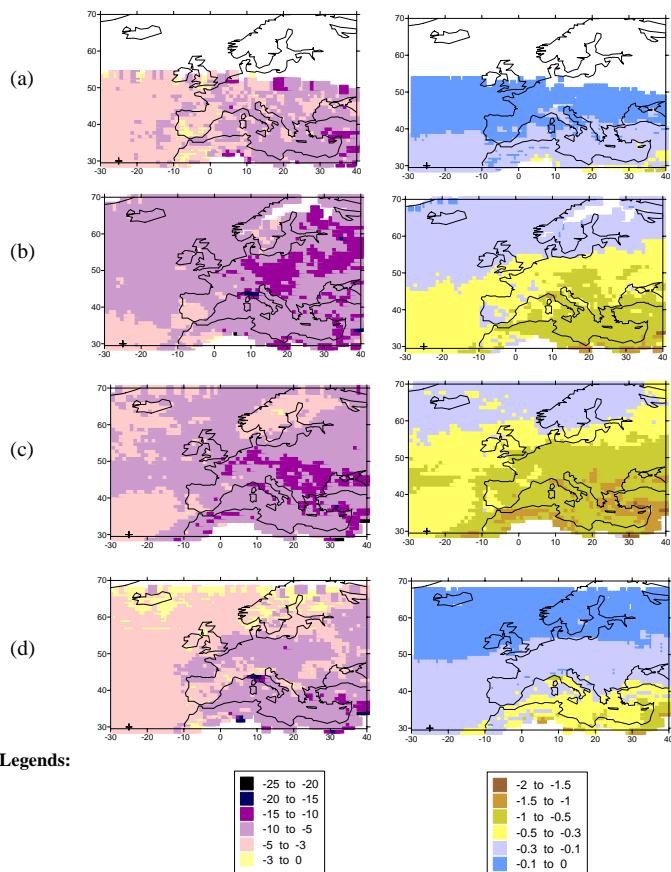


Fig. 10. Relative (left) (in percents) and absolute (right) (dimensionless) attenuation of UV indices due to aerosol in January **(a)**, April **(b)**, July **(c)** and October **(d)** over Europe. Noon conditions.