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Article

Impact of Desert Dust on Air Quality: What is the Meaningfulness of Daily PM Standards in Regions Close to the Sources? The Example of Southern Tunisia

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Abstract: Desert dust is now recognized as a major health hazard. However, there still exists a lack of measurements of desert dust atmospheric loads in regions located in the vicinity of the major desert areas, where a growing part of the world population is living. Dust emission is a sporadic and intense phenomenon so that the classical air quality standards should not be appropriate to reflect the real population exposure to desert dust. In order to give some insight to answer this question, PM₁₀ concentrations were continuously measured at a five-minute time step in southern Tunisia from February 2014 to February 2019. The daily and annual PM₁₀ concentrations were first discussed according to the Tunisian air quality standards: In this relatively remote area, close to dust source, these standards were respected at the annual, but no longer at the daily scale. Measurements performed at a high temporal resolution (five minutes) allowed to discriminate the different situations that led to exceed Tunisian daily standards in southern Tunisia. In particular, intense (five-minute PM₁₀ concentrations up to more than 1500 µg m⁻³) and short-lived (a few hours) dust outbreaks were regularly observed. This result raises the question of the necessity of defining specific air quality standards at the sub-daily scale in countries affected by intense desert dust outbreaks.

Keywords: desert dust; air quality standards; PM₁₀

1. Introduction

During recent decades, more and more attention has been paid to the atmospheric level of particulate matter (PM), especially for their potential impact on human health. Indeed, the fine particles can be inhaled and, according to their diameter, can enter the respiratory system up to the alveoli (e.g., [1–3]) so that they are now considered as a major health hazard (e.g., [4–7]).

Among them, mineral dust, also commonly referred to as desert dust because of their geographical origins, is of crucial interest as it represents about 30% to 35% of the total mass of aerosols emitted into the atmosphere every year [8]. Most of the dust emissions come from the Sahara Desert, followed by Chinese and Central Asian deserts, the Arabian Peninsula, and the Australian deserts [9]. The population living in the vicinity of these regions can be exposed to very high levels of mineral dust concentrations. Considering their physical, chemical, and biological properties, the impacts of desert

dust on human health have been widely studied: Since 2010, about ten literature reviews dealing with the impact of desert dust on human health were published [9–17]. Indeed, since mineral dust covers a large size spectrum, ranging from some tenths of μm to 40 or 50 μm in diameter, these particles induce various impacts on human health. Particles with diameters larger than 10 μm are not inhaled, but are mainly responsible for skin and eye irritations or infections [11,15,17]. Breathable particles, those smaller than 10 μm in diameter (PM_{10}), are frequently trapped inside the upper respiratory system, while the smallest ones are associated with respiratory disorders such as asthma, tracheitis, pneumonia, allergic rhinitis, and silicosis [11,15]. Death and injuries are also associated with transport accidents due to a reduction of the visibility during dust storms (e.g., [18]). Giannadaki et al. [19], based on a modelling study of the PM smaller than 2.5 μm in diameter ($\text{PM}_{2.5}$), suggested that desert dust are responsible for a large number of premature deaths by cardiopulmonary diseases and for a significant number of deaths by lung cancer, mostly in the “dust belt” region (which is defined as an area of strong desert dust sources that extends from the west coast of North Africa, through the Middle East, South and Central Asia, to eastern China). These results were supported by the previous findings by Evans et al. [20] who used, for the first time, satellite imagery to assess the expected number of deaths due to chronic $\text{PM}_{2.5}$ exposure. These authors found that the $\text{PM}_{2.5}$ premature mortality in the Mediterranean region (several countries in North Africa and in the Middle East) was mainly related to the non-anthropogenic component (dust) of total $\text{PM}_{2.5}$. Giannadaki et al. [19] also estimated 3.56 million as the annual years of human life lost per year associated to desert dust $\text{PM}_{2.5}$ exposure.

Even if countries located in the “dust belt” region are the most impacted by desert dust, de Longueville et al. [10,21] pointed out the fact that a few epidemiological studies were conducted in this area. In particular, none of the studies listed by these authors were conducted in Africa, and only one in the Middle East. After having conducted the same type of bibliographic investigation as de Longueville et al. [10], we found that: (i) For the period October 2011–August 2017, the number of epidemiological studies focusing on the impact of desert dust on human health has significantly increased (50 studies for the period January 1999–September 2011 [10] compared to 111 for the period October 2011–August 2017); and (ii) such epidemiological studies are now conducted in the countries located in the “dust belt”—the majority of these studies concerns the Middle East (including Iran, Israel, Kuwait and Saudi Arabia), but a few of them concerned African countries (zero in the Maghreb). de Longueville et al. [10] identified two main causes to this lack of information on the effect of dust on the population living in the countries bordering the Sahara: (i) The absence of systematic particulate air pollution data (e.g., [22]); and (ii) poor-quality public health data that could be used to adequately support impact studies (e.g., [23]). The absence of systematic particulate air pollution data is still a reality when focusing on Africa: In the 2016 update of the WHO Global Urban Ambient Air Pollution Database (http://www.who.int/phe/health_topics/outdoorair/databases/cities/en/), which reports annual means of $\text{PM}_{2.5}$ and PM_{10} concentrations measured throughout the world (2975 locations), measurements for only 52 cities located in 13 African countries are reported, including seven cities in Morocco and four cities in Tunisia.

The south of Tunisia is a region located at the northern border of the Sahara Desert. It is composed of five Governorates (Tataouine, Médenine, Gabès, Kébili, and Tozeur) representing about one million of inhabitants (i.e., 9% of the whole population of Tunisia) [24]. It is a region where soils are sandy, the precipitation scarce, and the vegetation sparse, so that wind erosion, the mechanism that produces mineral dust, is very active. Moreover, the development of mechanized agriculture has increased wind erosion during the last decades (e.g., [25]). Finally, this region is known to be one of the main pathways for air masses coming from the Sahara to the central Mediterranean Basin [26–33]. For all these reasons, dust haze is frequently observed in southern Tunisia: For the period 1965–2011, the annual mean number of days of “dusty wind” was 74 in El Borma (Governorate of Tataouine), 63 in Tozeur, 45 in Remada (Governorate of Tataouine) and in Kébili, 23 in Médenine, 20 in Gabès, and 18 in Djerba [34].

In 2007, Tunisia adopted air quality standards to ensure every citizen the right to a healthy environment and sustainable development (Law No. 2007-34 of 4 June 2007 on air quality). Standards for

airborne particles (PM₁₀ [35,36]) were defined with daily and annual limit values for health and for well-being (Table 1). Tunisian air quality standards fix an annual limit value relative to health of 80 µg m⁻³, which is twice higher than the European air quality standards for PM₁₀ (40 µg m⁻³) and four times higher than the WHO air quality guidelines for PM₁₀ (20 µg m⁻³; [37]). Considering the second air quality standard defined on the daily level, the limit value of concentration relative to health in Tunisia cannot exceed 260 µg m⁻³ more than once a year while in Europe, a daily mean PM₁₀ concentration of 50 µg m⁻³ (as defined by the WHO and which is about 1/5 of the Tunisian daily limit) cannot be exceeded more than 35 times per year.

Table 1. Tunisian ambient air standards NT 106 04 for airborne particles.

Type of Average	Authorization of Excess	Limit Value Relative to Health (µg m ⁻³)	Limit Value Relative to Well-Being (µg m ⁻³)
Annual	-	80	40–60
Daily	once a year	260	120

In this framework, a national network of air quality monitoring stations (Réseau National de Surveillance de la Qualité de l’Air (RNSQA); www.anpe.nat.tn) has been deployed. In 2011, 30 stations were already installed in the main urban and industrial sites of northern Tunisia, and seven allowed the measurement of PM₁₀ concentrations by means of automatic beta radiation attenuation monitors [35,36]. However, none of these stations are located in the south of the country; therefore, population exposure to high desert dust atmospheric loads is not characterized. To do so, as a first step, PM₁₀ concentration has to be monitored in a remote place, closer to dust sources and away from other sources of pollutants, especially from anthropogenic activities. This is why, in February 2014, a ground-based station dedicated to the monitoring of PM₁₀ concentration was installed in the campus of the Institut des Régions Arides (IRA) of Médenine. At the monitoring station of El Fjé, measurements are performed with a short time step (five minutes) in order to assess, as well as possible, the high concentration periods.

In this paper, PM₁₀ concentrations measured during the period February 2014–February 2019 at the station of El Fjé in southern Tunisia are discussed first according to the Tunisian air quality standards, but also according to the way these standards were exceeded. This allows one to conclude on the relevance of actual air quality standards used in regions downwind of active desert dust sources.

2. Materials and Methods

2.1. Location of the Station

The dust station is installed inside the campus of the IRA Médenine in El Fjé (33.49963N; 10.64255E), as identified by the red dot in the satellite image in Figure 1. This place is located away from any other pollutant sources, especially from anthropogenic origin (biomass burning, road traffic, etc.) and is representative of the atmospheric background of the region. Indeed, the surroundings of the station are an agricultural area, mainly dominated by olive tree plantations, the most important town, Médenine (about 70,000 inhabitants [24]), being at more than 20 km south west. The Mediterranean Sea is about 5 km to the east, and 20 km to the north, while the Matmata Mounts are about 50 km to the west. From a climatological point of view, the monitoring station is located in the north east of the Jeffara plain, in the arid part of Tunisia where mean annual rainfalls ranging between 135 and 215 mm yr⁻¹ were recorded during the 1949–2001 period in 8 meteorological stations [38].



Figure 1. Location of the dust monitoring station (red dot), of the stations of the Réseau National de Surveillance de la Qualité de l’Air (RNSQA) mentioned in this paper (yellow dots), of the stations of the Tunisian National Institute of Meteorology mentioned in this paper (green dots), and of some the main geomorphological features of the region (terms in italics) (adapted from a Google Earth image).

2.2. Instruments and Method

The monitoring station of El Fjé was set up following the model of the stations of the Sahelian Dust Transect deployed in the Sahel since 2006 ([39]; <http://www.lisa.u-pec.fr/SDT/index.php?p=1>), now part of the International Network to Study Deposition and Atmospheric chemistry in Africa (INDAAF; <https://indaaf.obs-mip.fr/?rubrique38>). The same kind of instruments were chosen for their capability to resist severe dusty and meteorological conditions. All these instruments were installed at 2 m height from the flat roof of the higher building of the campus (i.e., about 14.5 m above the ground level).

2.2.1. PM₁₀ Concentration

PM₁₀ concentrations were measured using a Tapered Element Oscillating Microbalance (TEOM™ 1405 from Thermo Scientific™, Franklin (MA), USA) equipped with a PM₁₀ inlet (Thermo Scientific™). This instrument, widely used in air quality networks, allows one to measure atmospheric particulate concentrations ranging from a few micrograms up to one gram per cubic meter.

The microbalance was installed in an air-conditioned room to maintain ideal conditions for the TEOM™ sensor and control units. It was positioned about 5 m vertically below the PM₁₀ inlet. The flow rate is 3 L min⁻¹ and the averaging time is 5 min in these operational conditions, the nominal precision of the instrument is ± 2.0 µg m⁻³ for 1 h averaging, and ± 1.0 µg m⁻³ for 24 h averaging.

The lower part of the sampling tube (60 cm length) and the measurement chamber of the TEOM™ were heated at a temperature of 50 °C in order to allow a thermal stability of the particles and the filter and to minimize water condensation interferences. This heating may induce a loss of a part of the most volatile particulate compounds, such as ammonium nitrates or condensate organic compounds [40,41]. Thus, the contribution of carbonaceous and/or anthropogenic aerosols to the PM₁₀ concentration is

probably slightly underestimated when using this type of measurement. Moreover, the heating of the sampled air can lead to negative values of PM₁₀ concentrations when relative humidity is high, and PM₁₀ concentrations especially low. This can be observed during intrusions of air masses coming from the sea and lightly loaded in PM.

2.2.2. Sunphotometric Measurements

A Cimel sunphotometer from the AERONET (AERosol RObotic NETwork)/PHOTONS (Photométrie pour le Traitement Opérationnel de Normalisation Satellitaire) network ([42]; <http://aeronet.gsfc.nasa.gov/>) was installed in July 2014 at the monitoring station. This instrument allows one to retrieve the Aerosol Optical Depth (AOD) and the corresponding Angström exponent (α) from the measurements of the direct solar radiation at 4 wavelengths (1020, 870, 675, and 440 nm) during daytime [43,44]. In this study, data of level 1.5 (i.e., quality checked data) were used because data of level 2.0 (i.e., quality checked and cloud screened data) was only available until 27 September 2018.

2.2.3. Data Selection

The monthly recovery rate ranged between 65 and nearly 100% for PM₁₀ concentrations (see Table A1 in Appendix A), with the exception of October 2015 when the main electronic card of the TEOM™ broke down. Most of the time, missing data corresponded to particular events such as operations of maintenance on the instruments (filters changing, cleaning, control, etc.), electricity failures, or break down of the computer connections.

In order to compare to the Tunisian air quality standards, daily mean PM₁₀ concentrations were computed. Firstly, the hourly mean PM₁₀ concentration was computed from the 5 min measurements. Secondly, the daily mean PM₁₀ concentration was computed from the hourly mean PM₁₀ concentration (without accounting for the hourly negative values when there were). Considering a one-month period, 9% at a maximum (in June/July) of the hourly mean PM₁₀ concentration were removed for the computation of the daily mean PM₁₀ concentration because it was negative. For the rest of the year, negative values of the hourly mean PM₁₀ concentration never accounted for more than 5% of the total values for one month.

As it was mentioned in Section 2.2.1., the cases corresponding to negative values of the hourly mean PM₁₀ concentration were mainly cases during which PM₁₀ concentration was low. Thus, eliminating these negative values for the computation of the daily mean PM₁₀ concentration could generate a bias (i.e., could lead to overestimation of daily means). In a context of air quality, this could produce “false” excesses of air quality standards, and the impact of PM₁₀ could be overestimated. In order to test the impact of the method used to compute the daily mean PM₁₀ concentration, we replaced the negative hourly values (instead of removing them) by the nominal precision of the instrument for 1 h averaging (i.e., 2.0 $\mu\text{g m}^{-3}$). In these conditions, the coefficient of determination R^2 between the daily mean PM₁₀ concentrations computed using these two methods was 0.9995 with a slope of 0.999 for the regression line ($N = 1790$). In terms of air quality, a difference of 1 day over the 5-year period was observed between the two methods when estimating the number of times the limit values relative to well-being and relative to health were exceeded. As a conclusion, we can consider that removing the negative values of the hourly mean PM₁₀ concentration for the computation of the daily mean PM₁₀ concentration does not significantly affect the results that are presented and discussed in the present paper.

3. Results and Discussion

3.1. Tunisian Air Quality Standards

3.1.1. Annual Mean PM₁₀ Concentrations

The annual mean concentrations of PM₁₀ in El Fjé range between 32 and 56 µg m⁻³ from 2014 to 2018 (Table 2). These values are lower than 60 µg m⁻³ for all years respecting both the limit values relative to health and to well-being defined by Tunisian air quality standards for airborne particles (Table 1). On the other hand, for the five years, the WHO air quality guidelines for PM₁₀ were exceeded by, at least, a factor 1.5 (Table 2).

Table 2. Annual mean PM₁₀ concentrations (in µg m⁻³) recorded in El Fjé from 2014 to 2018. The asterisk highlights the years for which a month of data is missing.

Year.	Annual Mean PM ₁₀ Concentrations (µg m ⁻³)
2014 *	56
2015 *	44
2016	51
2017	32
2018	42

Figure 2 presents an inventory of the most recent available measurements of annual mean PM₁₀ concentrations in the countries downwind the Sahara and frequently impacted by Saharan dust outbreaks (i.e., North Africa, South Europe, and West Africa). Firstly, it can be noted that most of the available measurements are located in urban and suburban areas (63%): Only 20 stations (upon 116 in total) are classified as rural or regional background stations (see Table A2 in Appendix B). El Fjé appears as the unique rural station in Tunisia and, to a regional extent, in North Africa. The annual mean PM₁₀ concentrations measured in El Fjé are low (about three times lower for 2017) compared to those measured in the northern Tunisian urban centers (except Sousse—see measurements reported for Bizerte, Sfax, and Tunis in Figure 2). On the other hand, the annual mean PM₁₀ concentrations measured in El Fjé are 1.6 to 5 times higher than the values reported by Pey et al. [45] for south European urban centers located farther from the dust sources (Figure 2). Finally, the annual mean PM₁₀ concentrations measured in El Fjé are 2.5 to 3.5 times lower than what can be measured at the rural stations in the Sahel (see measurements reported for Cinzana and Banizoumbou by Kaly et al. [46] in Figure 2), which suggests that southern Tunisia is not a dust source or a pathway for Saharan dust transport as important as the Sahel is.

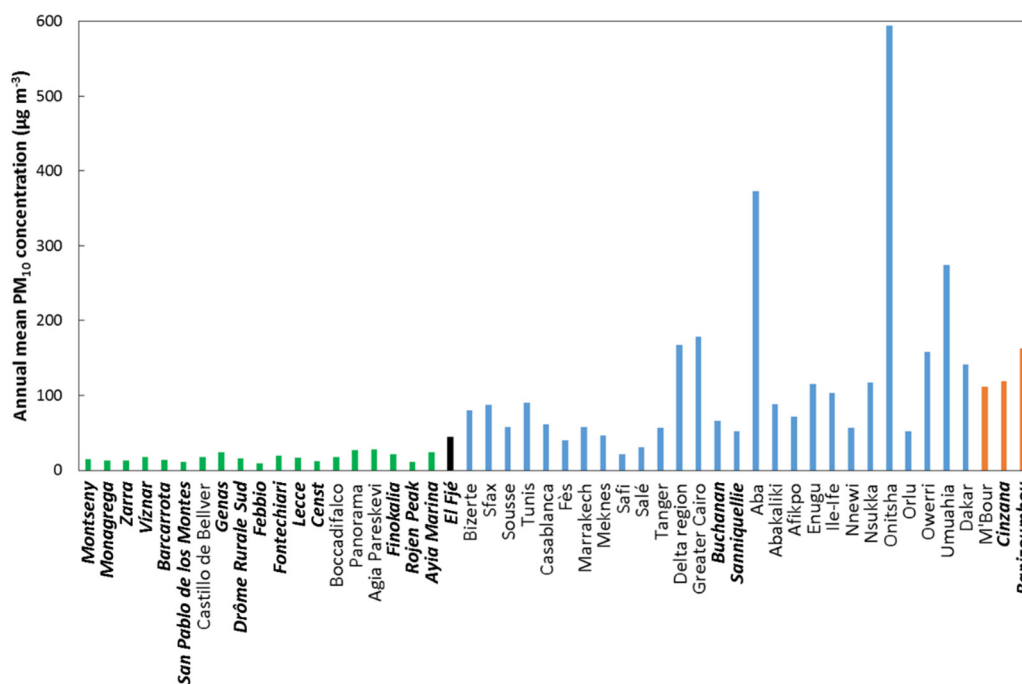


Figure 2. Annual mean PM₁₀ concentrations (in $\mu\text{g m}^{-3}$) recorded in the countries downwind the Sahara. Colors indicate data source: Green = Pey et al. [45]; black = this study; blue = the 2016 WHO GUAAP Database; orange = Kaly et al. [46]. Details on the location, type, and data coverage is given for each station in Table A2 (Appendix B). Rural and regional background stations are identified using bold italics font.

3.1.2. Daily Mean PM₁₀ Concentrations

The Tunisian air quality standards were no longer respected at the daily time scale in El Fjé: during the period 12 February 2014–28 February 2019, the daily limit value relative to health was exceeded 40 times (i.e., 2% of all the measured PM₁₀ concentrations), while the daily limit value relative to well-being was exceeded 107 times (i.e., 6% of all the measured PM₁₀ concentrations) (see Table A3 in Appendix C).

Figure 3 presents the monthly frequency of occurrence of days with daily mean PM₁₀ concentrations greater than 120 and 260 $\mu\text{g m}^{-3}$. Daily air quality standards were more frequently exceeded during winter (DJF) and spring (MAM), than in autumn (SON) and summer (JJA), during which it is frequently null. This seasonal cycle follows the seasonal cycle of desert dust outbreaks observed in this region, strongly suggesting that desert dust aerosols were responsible for daily air quality standards excesses. Indeed, Moulin et al. [31] reported that April was a period of intense dust uplift over North Africa because of the Sharav cyclones [47], and that the major source regions for the Mediterranean were: (i) South Tunisia around 30° N for the western central basin [26], and Libyan desert just below 30° N for the eastern central basin [48]. More recently, Fiedler et al. [49] showed that above 20° N, up to 80% of desert dust emissions was due to atmospheric depressions whose frequency over the north of Africa is the highest between March and May, and then between December and February.

Despite this obvious mean annual cycle, daily mean PM₁₀ concentration highly varied throughout the year (Figure 4): High daily mean PM₁₀ concentrations were frequently observed, sometimes recorded during several consecutive days, and can be greater than 1500 $\mu\text{g m}^{-3}$ (up to 1642 $\mu\text{g m}^{-3}$ on 22 March 2016). As a comparison, in the Sahel, where dust also dominate the background concentrations, the maximum daily PM₁₀ concentrations were 4020 $\mu\text{g m}^{-3}$ in Banizoumbou, 2503 $\mu\text{g m}^{-3}$ in Cinzana, and 2250 $\mu\text{g m}^{-3}$ in M'Bour over the period 2006–2008, and daily mean PM₁₀ concentrations greater than 100 $\mu\text{g m}^{-3}$ represented about 40% of all the measured daily mean PM₁₀ concentrations [39].

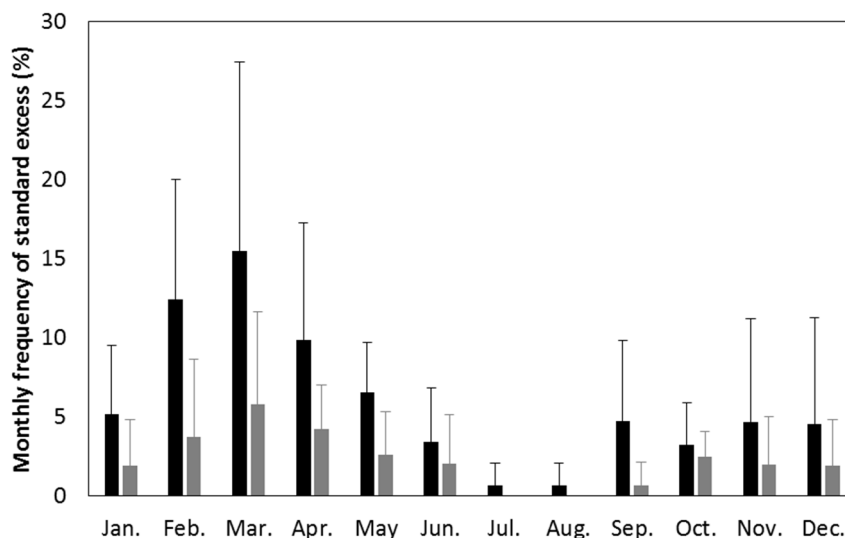


Figure 3. Percentage of days in the month with daily mean PM₁₀ concentrations greater than 120 µg m⁻³ (grey bars) and 260 µg m⁻³ (black bars) averaged for the period February 2014–February 2019.

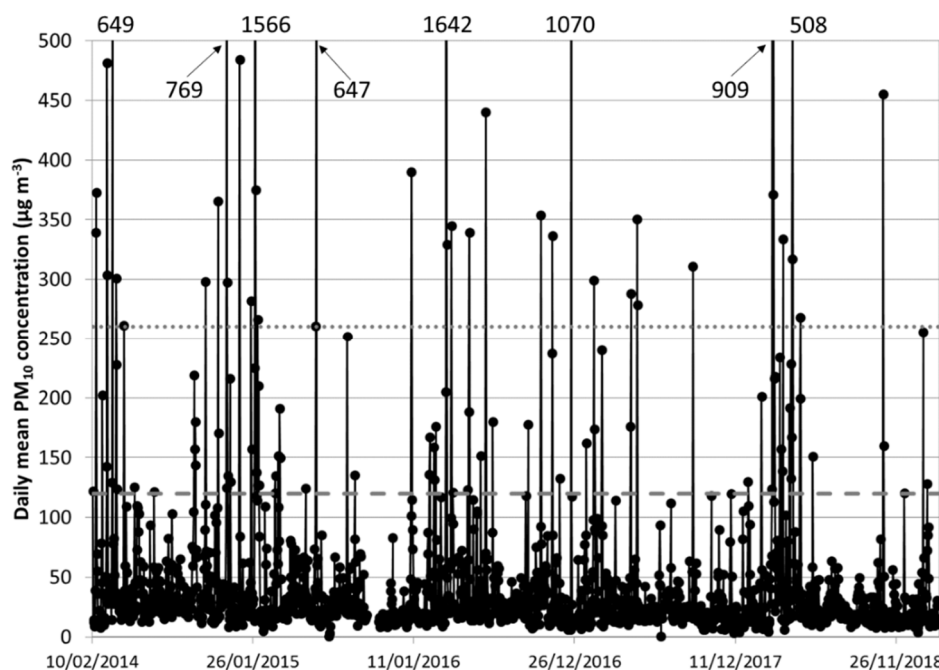


Figure 4. Daily mean of PM₁₀ concentration (in µg m⁻³; black line with dots) from 10 February 2014 to 28 February 2019. Numbers in and above the graph give the highest measured values. Grey dashed and dotted lines indicate the limit value relative to well-being (120 µg m⁻³) and health (260 µg m⁻³) according to Tunisian standards, respectively.

3.2. The Situations of the Excess of the Tunisian Daily Air Quality Standards

3.2.1. Nature of the Aerosol

In order to identify the type(s) of particles responsible for the 107 excesses of the Tunisian daily limit value relative to well-being, we used as an indicator the Angström exponent (440–870 nm), $\alpha_{440-870}$. Indeed, Holben et al. [50] considered that $\alpha_{440-870}$ values less than 0.4 are typical of situations dominated by coarse aerosols (sea salt, mineral dust) whereas larger $\alpha_{440-870}$ values denote the presence of submicron aerosols such as those produced by biomass burning or other combustion processes. Figure 5 reports the daily minimum $\alpha_{440-870}$ as a function of the daily mean PM₁₀ concentration for

the whole period. It can be seen that, for all days (except one) with a daily mean PM_{10} concentration greater than $120 \mu\text{g m}^{-3}$, the daily minimum $\alpha_{440-870}$ is systematically lower than 0.4, which means that AOD was strongly controlled by coarse particles. Kchih et al. [51] analyzed the composition of PM_{10} collected in the southern Mediterranean coast of Tunis. These authors found that in all sites where PM_{10} were collected, these particles were mainly composed, in mass, of crustal matter (45%–50%) and poorly of sea salt (3%–4%), which showed that aerosol chemical composition was slightly impacted by the nearby Mediterranean Sea. These results confirm that the recorded PM_{10} maximums in El Fjé were due to mineral dust.

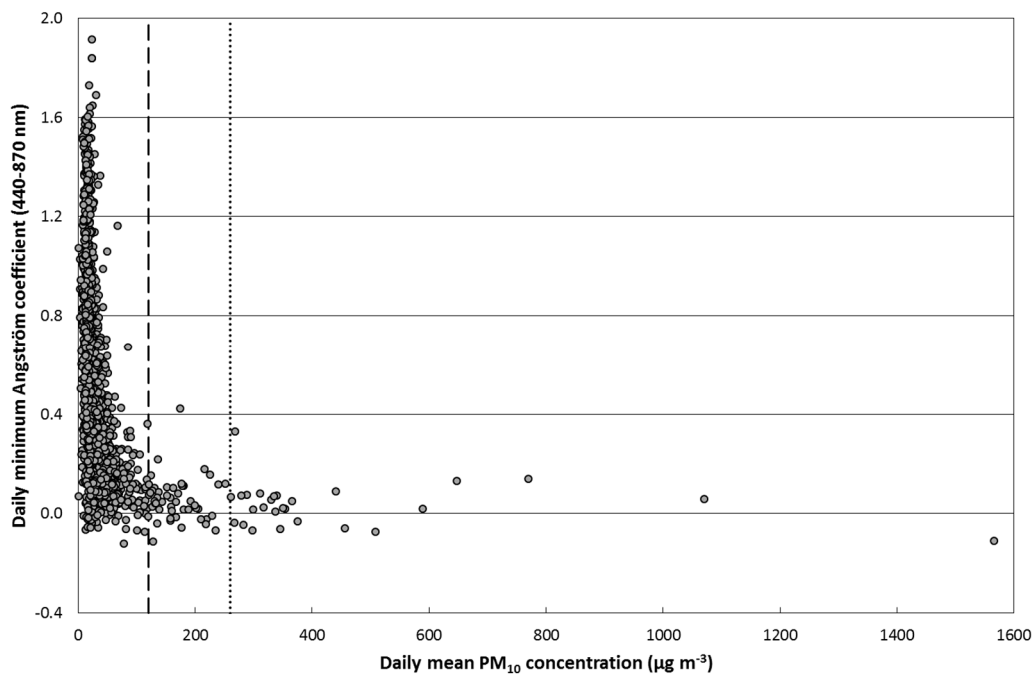


Figure 5. Daily minimum $\alpha_{440-870}$ as a function of daily mean PM_{10} concentration (in $\mu\text{g m}^{-3}$) from 24 July 2014 to 28 February 2019. Dashed and dotted lines indicate the limit value relative to well-being ($120 \mu\text{g m}^{-3}$) and health ($260 \mu\text{g m}^{-3}$) according to Tunisian standards, respectively.

3.2.2. Sub-Daily Variability

Considering the impact of desert dust on human health, we can wonder what differences exist between a long-term exposure to high levels of PM_{10} concentration and a short-term exposure to very high PM_{10} concentration. Indeed, in extreme cases, people and animals caught in a severe dust storm may die due to suffocation [9]. For instance, a major dust storm in April 1895 in the US state of Colorado was reported to have caused the loss of 20% of the cattle in the east of the state [52]. When examining all the days with a daily mean PM_{10} concentration greater than $120 \mu\text{g m}^{-3}$, we noticed that the daily limit value can be exceeded in different ways. In order to investigate it, we searched for two days for which the daily mean PM_{10} concentration was greater than $120 \mu\text{g m}^{-3}$ and in the same order of magnitude, but during which the five-minute PM_{10} concentration dynamics was completely different. Because the number of days exhibiting a daily mean PM_{10} concentration below $260 \mu\text{g m}^{-3}$ represents more than a half of the days for which the daily limit value relative to well-being was exceeded, we selected, as an example, two days for which a daily mean PM_{10} concentration of $\sim 125 \mu\text{g m}^{-3}$ was recorded—12 February 2014 and 13 May 2014.

Figure 6 presents the temporal evolution of the five-minute PM_{10} concentrations during these two days. In the first case, on 12 February 2014, five-minute PM_{10} concentrations were higher than $100 \mu\text{g m}^{-3}$ during more than half of the day, but never exceeding $400 \mu\text{g m}^{-3}$, while, in the second case, on 13 May 2014, the daily limit value relative to well-being was exceeded because of the occurrence of a peak of very high PM_{10} concentration (up to $3398 \mu\text{g m}^{-3}$ on five minutes) over a short time period

(less than 1 h). Such differences in the five-minute PM_{10} concentration dynamics were also observed during days with higher daily mean PM_{10} concentrations (not shown).

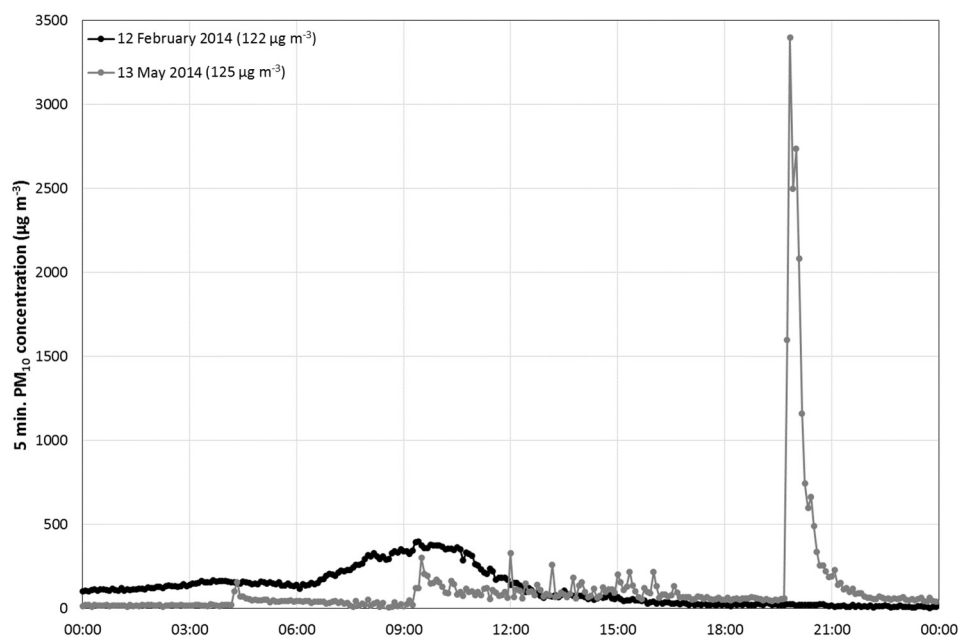


Figure 6. Five-minute PM_{10} concentrations (in $\mu\text{g m}^{-3}$) on 12 February 2014 (black), and 13 May 2014 (grey). For each date, daily mean PM_{10} concentration is given in parenthesis.

The differences observed in the temporal evolution of PM_{10} concentrations between the two presented days can be explained by the meteorological situation (see the detailed analysis presented in Appendix D): On 12 February 2014, high PM_{10} concentrations measured at the station of El Fjé resulted from dust transport from the region of the Chotts in the western part of Tunisia, while, on 13 May 2014, the same PM_{10} level was the consequence of local wind erosion in the Jeffara plain.

These observations raise the question of the possible need to define sub-daily limits to warn populations in case of very intense dust events. Middleton [9] reported on the public information system established by the Korea Meteorological Administration (Asian dust warning system; <http://web.kma.go.kr/eng/weather/asiandust/intro.jsp>) to warn the population of possible health effects in case of intrusion of desert dust coming from long-range transport from China and Mongolia across the Korean peninsula. This warning system includes two levels based on the value of the hourly mean PM_{10} concentration: Advisory (respectively warning) when the hourly mean PM_{10} concentration is expected to exceed $400 \mu\text{g m}^{-3}$ (respectively $800 \mu\text{g m}^{-3}$) for over 2 h.

During the period February 2014–February 2019, in El Fjé, the hourly mean PM_{10} concentration was greater than $400 \mu\text{g m}^{-3}$ 472 times, and greater than $800 \mu\text{g m}^{-3}$ 170 times, so that the advisory level would have been triggered 360 times while the warning level would have been triggered 115 times. At a maximum, the hourly mean PM_{10} concentration was greater than $400 \mu\text{g m}^{-3}$ during 27 consecutive hours from 30/01/2015 22:00UTC to 01/02/2015 00:00UTC; during this period, the hourly mean PM_{10} concentration was greater than $800 \mu\text{g m}^{-3}$ during 20 h. Even if such situations are exceptional (at the hourly timescale, PM_{10} concentrations were, most of the time (77%), lower than $40 \mu\text{g m}^{-3}$, Figure 7), they clearly suggest that people living in this region (and more generally, those living close to dust sources) are regularly exposed during several hours to very high PM_{10} concentrations, exceeding by far all the air quality standards. The health consequences of such intense but relatively short exposure of the population to dust events remain to be documented and the definition of air quality standards accounting for the intermittency of dust emission should be encouraged in regions where this phenomenon strongly controls air quality.

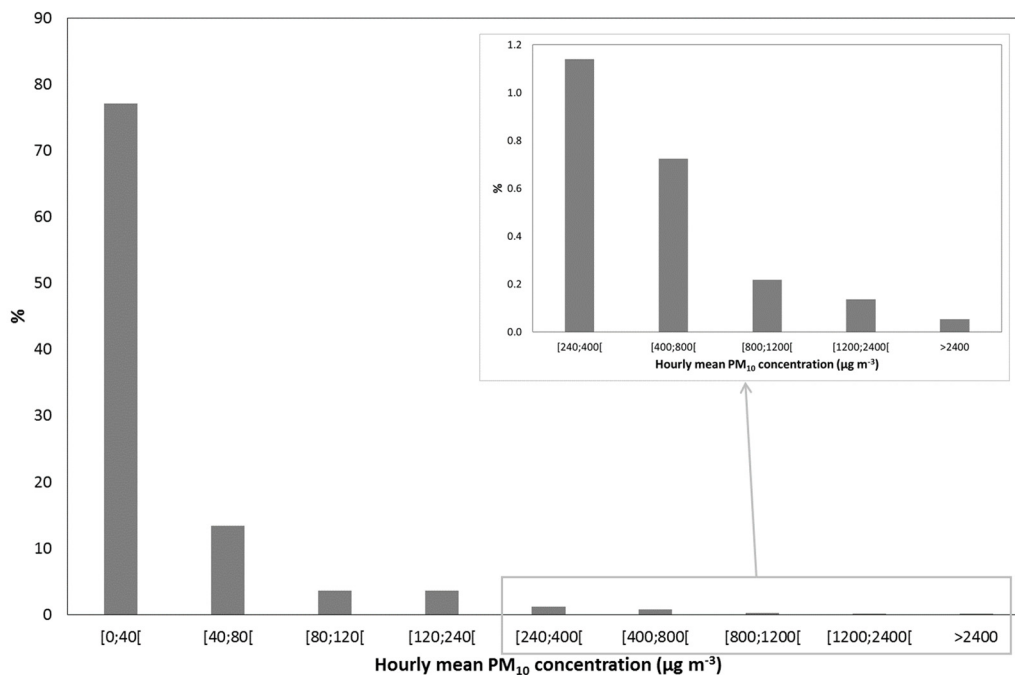


Figure 7. Percentage of time during which the hourly mean PM₁₀ concentration was between [0;40[, [40;80[, [80;120[, [120;240[, [240;400[, [400;800[, [800;1200[, [1200;2400[and greater than 2400 µg m⁻³ in El Fjé during the period February 2014–February 2019.

4. Conclusions

In order to investigate the role of mineral dust on air quality in a region where this aerosol species is dominant, PM₁₀ concentrations were measured at a five-minute time step from February 2014 to February 2019 in a remote place of southern Tunisia (El Fjé).

Simultaneous measurements of the AOD and the Angström exponent with a sunphotometer were also performed in the station of El Fjé. These complementary measurements allowed to identify the nature of the aerosols responsible for the excess of the daily Tunisian air quality standards—mineral dust.

The results showed that, on an annual basis, the Tunisian air quality standards were respected in this part of the country contrary to what was measured for the northernmost urban and industrial stations of Tunisia: Annual means of PM₁₀ concentration recorded in El Fjé were significantly lower than those recorded in the main coastal urban centers of northern Tunisia. On the other hand, they appeared high (up to twice higher) for a rural area when compared to stations located far from dust sources (southern Europe), but low when compared to Sahelian stations. In the latter case, this suggests that southern Tunisia cannot be considered as a continuous active source of mineral dust, nor as a regular pathway for mineral dust emitted in the Sahara. On a daily basis, Tunisian air quality standards were no longer respected.

The five-minute time step of PM₁₀ concentration measurements allowed to identify the manner these standards were exceeded. Two contrasted cases were identified: The high daily mean PM₁₀ concentration that was measured during these days was either due to high PM₁₀ concentrations (some thousands of µg m⁻³) lasting several hours or due to a brief, but very high peak of PM₁₀ concentration (> 1000 µg m⁻³) over a short time period (less than 2 h). Using concomitant measurements of mean and maximum wind speed at the same time step as PM₁₀ concentrations should give some insight on the local or regional origin of the dust responsible for the excess of air quality standards. Nevertheless, the use of back-trajectory analysis and space-borne measurements in complement would be necessary to confirm the obtained results.

The different ways daily standards were exceeded at the sub-daily scale question the fact that daily mean PM₁₀ concentration is not the most relevant indicator to quantify population exposure to

desert dust. Indeed, during the period February 2014–February 2019, in El Fjé, the hourly mean PM₁₀ concentration was greater than 400 $\mu\text{g m}^{-3}$ 472 times, and greater than 800 $\mu\text{g m}^{-3}$ 170 times, while the daily limit value relative to health (daily mean PM₁₀ concentration greater than 260 $\mu\text{g m}^{-3}$) was only exceeded 40 times. As a conclusion, the definition of sub-daily indicators should be then considered at the same time as the impact of the high peaks of PM₁₀ concentration on human health should be investigated. This would require epidemiological studies to be conducted on the association of high dust concentration at the sub-daily scale with health effect (such as irritation symptoms, hospital admission respiratory disease, hospital admission cardiovascular disease, respiratory mortality, cardiovascular mortality, or total mortality) in regions located downwind desert areas, such as the Sahara. As a first step, it should be useful to compare the respiratory and/or cardiovascular hospitalization rates in these areas with those in other urban regions in north Tunisia. Nevertheless, in the inhabited areas located downwind of deserts, atmospheric pollution due to mineral dust cannot be reduced, such as pollution due to road traffic or industries. Before precisely assessing the role of short bursts of mineral dust on public health, a simple preventive step to protect the local populations should be to set up warning systems based on existing weather and atmospheric dust load forecasts, such as the one already running in Korea. This would allow the local population to be protected during intense dust events following simple rules such as avoiding outdoor activities, wearing anti-pollution masks, etc.

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Appendix A

Table A1. Percentage of available data on a five-minute time step for the TEOM™, and number of days in the month where Aerosol Optical Depth (AOD) measurement is available. ND stands for “no data”.

	TEOM™ (%)	Photometer (Number of Days)
February 2014	99	ND
March 2014	100	ND
April 2014	100	ND
May 2014	100	ND
June 2014	100	ND
July 2014	99	4
August 2014	77	31
September 2014	100	29

Table A1. Cont.

	TEOM™ (%)	Photometer (Number of Days)
October 2014	94	30
November 2014	100	27
December 2014	98	9
January 2015	100	24
February 2015	100	20
March 2015	100	ND
April 2015	99	ND
May 2015	89	26
June 2015	94	30
July 2015	99	31
August 2015	100	28
September 2015	88	21
October 2015	17	26
November 2015	100	25
December 2015	99	30
January 2016	98	23
February 2016	99	25
March 2016	99	29
April 2016	65	29
May 2016	99	26
June 2016	99	16
July 2016	99	31
August 2016	99	31
September 2016	97	29
October 2016	99	31
November 2016	99	22
December 2016	100	27
January 2017	100	29
February 2017	100	28
March 2017	99	27
April 2017	99	23
May 2017	99	30
June 2017	97	29
July 2017	82	30
August 2017	99	31
September 2017	99	29
October 2017	100	28
November 2017	89	12
December 2017	99	26
January 2018	100	27
February 2018	99	20
March 2018	100	28
April 2018	99	28
May 2018	99	29
June 2018	99	29
July 2018	90	31
August 2018	100	30
September 2018	99	26
October 2018	99	16
November 2018	99	27
December 2018	100	31
January 2019	99	27
February 2019	100	18

Appendix B

Table A2. Location, type, data coverage, and source of the annual mean PM₁₀ concentrations (in $\mu\text{g m}^{-3}$) for each station presented in Figure 2. NA means that the type of station was not given in the original database.

Region	Country	City	Type of Station	Year	Source
South Europe	Spain	Montseny	Regional Background	2002–2011 mean value	Pey et al. [45]
South Europe	Spain	Monagrega	Regional Background	2001–2011 mean value	Pey et al. [45]
South Europe	Spain	Zarra	Regional Background	2001–2011 mean value	Pey et al. [45]
South Europe	Spain	Víznar	Regional Background	2001–2011 mean value	Pey et al. [45]
South Europe	Spain	Barcarrota	Regional Background	2001–2011 mean value	Pey et al. [45]
South Europe	Spain	San Pablo de los Montes	Regional Background	2001–2011 mean value	Pey et al. [45]
South Europe	Spain	Castillo de Bellver	Sub-urban regional	2001–2011 mean value	Pey et al. [45]
South Europe	France	Genas	Regional Background	2001–2010 mean value	Pey et al. [45]
South Europe	France	Drôme Rurale Sud	Regional Background	2004–2010 mean value	Pey et al. [45]
South Europe	Italy	Febbio	Regional Background	2005–2010 mean value	Pey et al. [45]
South Europe	Italy	Fontechiari	Regional Background	2001–2010 mean value	Pey et al. [45]
South Europe	Italy	Lecce	Regional Background	2009–2010 mean value	Pey et al. [45]
South Europe	Italy	Censt	Regional Background	2005–2010 mean value	Pey et al. [45]
South Europe	Italy	Bocadifalco	Sub-urban regional	2001–2010 mean value	Pey et al. [45]
South Europe	Greece	Panorama	Sub-urban regional	2001–2010 mean value	Pey et al. [45]
South Europe	Greece	Agia Pareskevi	Sub-urban regional	2001–2010 mean value	Pey et al. [45]
South Europe	Greece	Finokalia	Regional Background	2004–2010 mean value	Pey et al. [45]
South Europe	Bulgaria	Rojen Peak	Regional Background	2005–2010 mean value	Pey et al. [45]
South Europe	Cyprus	Ayia Marina	Regional Background	2003–2010 mean value	Pey et al. [45]
North Africa	Tunisia	El Fjé	Rural	2014–2018 mean value	This work
North Africa	Tunisia	Bizerte	Urban residential/traffic/commercial	2010	WHO GUAAP Database (2016) *
North Africa	Tunisia	Sfax	Urban	2010	WHO GUAAP Database (2016)
North Africa	Tunisia	Sousse	Urban residential/traffic/commercial	2010	WHO GUAAP Database (2016)
North Africa	Tunisia	Tunis	Urban	2010	WHO GUAAP Database (2016)
North Africa	Morocco	Casablanca	4 stations, urban background station	2013	WHO GUAAP Database (2016)
North Africa	Morocco	Fès	Urban background station	2013	WHO GUAAP Database (2016)
North Africa	Morocco	Marrakech	2 stations, urban background station	2012	WHO GUAAP Database (2016)
North Africa	Morocco	Meknes	Urban	2008	WHO GUAAP Database (2016)
North Africa	Morocco	Safi	Urban background station	2011	WHO GUAAP Database (2016)
North Africa	Morocco	Salé	Urban background station	2011	WHO GUAAP Database (2016)
North Africa	Morocco	Tanger	Urban background station	2013	WHO GUAAP Database (2016)
North Africa	Egypt	Delta region	14 stations	2013	WHO GUAAP Database (2016)

Table A2. Cont.

Region	Country	City	Type of Station	Year	Source
North Africa	Egypt	Greater Cairo	47 stations	2013	WHO GUAAP Database (2016)
West Africa	Liberia	Buchanan	Rural, 500m away from community, Osris	2009	WHO GUAAP Database (2016)
West Africa	Liberia	Sanniquellie	Rural, within the community, Osiris monitors	2009	WHO GUAAP Database (2016)
West Africa	Nigeria	Aba	NA	2009	WHO GUAAP Database (2016)
West Africa	Nigeria	Abakaliki	NA	2009	WHO GUAAP Database (2016)
West Africa	Nigeria	Afikpo	NA	2009	WHO GUAAP Database (2016)
West Africa	Nigeria	Enugu	NA	2009	WHO GUAAP Database (2016)
West Africa	Nigeria	Ile-Ife	University campus	2010	WHO GUAAP Database (2016)
West Africa	Nigeria	Nnewi	NA	2009	WHO GUAAP Database (2016)
West Africa	Nigeria	Nsukka	NA	2009	WHO GUAAP Database (2016)
West Africa	Nigeria	Onitsha	NA	2009	WHO GUAAP Database (2016)
West Africa	Nigeria	Orlu	NA	2009	WHO GUAAP Database (2016)
West Africa	Nigeria	Owerri	NA	2009	WHO GUAAP Database (2016)
West Africa	Nigeria	Umuahia	NA	2009	WHO GUAAP Database (2016)
West Africa	Senegal	Dakar	4 stations, traffic, urban, industrial	2013	WHO GUAAP Database (2016)
West Africa	Senegal	M'Bour	Coastal/urban	2006–2010 mean value	Kaly et al. [46]
West Africa	Mali	Cinzana	Rural	2006–2010 mean value	Kaly et al. [46]
West Africa	Niger	Banizoumbou	Rural	2006–2010 mean value	Kaly et al. [46]

* 2016 update of the WHO Global Urban Ambient Air Pollution (GUAAP) Database (http://www.who.int/phe/health_topics/outdoorair/databases/cities/en/).

Appendix C

Table A3. Number of days by month with PM₁₀ measurements ($N_{\text{obs.}}$), and with daily mean PM₁₀ concentrations greater than 120 $\mu\text{g m}^{-3}$ and greater than 260 $\mu\text{g m}^{-3}$ for the period February 2014–February 2019.

Month	$N_{\text{obs.}}$	$[\text{PM}_{10}]_{\text{daily}} > 120 \mu\text{g m}^{-3}$	$[\text{PM}_{10}]_{\text{daily}} > 260 \mu\text{g m}^{-3}$
February 2014	17	3	2
March 2014	31	6	3
April 2014	30	4	2
May 2014	31	1	0
June 2014	29	1	0
July 2014	31	0	0
August 2014	25	0	0
September 2014	30	4	0
October 2014	30	1	1
November 2014	30	4	2
December 2014	31	5	2
January 2015	31	4	2
February 2015	28	6	2
March 2015	31	5	0
April 2015	30	0	0
May 2015	29	1	0
June 2015	29	2	2
July 2015	31	0	0
August 2015	31	1	0
September 2015	27	1	0
October 2015	5	0	0
November 2015	30	0	0
December 2015	31	0	0
January 2016	31	1	1
February 2016	29	5	0
March 2016	31	3	2
April 2016	22	2	1
May 2016	31	3	1
June 2016	30	2	1
July 2016	31	1	0
August 2016	31	0	0
September 2016	30	1	0
October 2016	31	1	1
November 2016	30	3	1
December 2016	31	1	1
January 2017	31	1	0
February 2017	28	3	1
March 2017	31	0	0
April 2017	30	2	1
May 2017	31	2	2
June 2017	30	0	0
July 2017	28	0	0
August 2017	31	0	0
September 2017	30	1	1
October 2017	31	0	0
November 2017	30	0	0
December 2017	31	0	0
January 2018	31	1	0
February 2018	28	1	0
March 2018	31	10	4
April 2018	30	6	2
May 2018	31	3	1
June 2018	30	0	0
July 2018	29	0	0
August 2018	31	0	0
September 2018	30	0	0
October 2018	31	2	1
November 2018	30	0	0
December 2018	31	1	0
January 2019	31	1	0
February 2019	28	1	0

Appendix D Detailed Analysis of the 12 February and 13 May 2014 Dust Events

At the station of El Fjé, a Gill 2D-type Windsonic anemometer (Campbell® Scientific Instruments) is also installed to measure wind speed and direction at a five-minute time step. As measurement frequency is 10 s, maximum and mean wind speed values measured during the five-minute time period are stored on a CR200 data logger (Campbell® Scientific Instruments, Shepshed, Loughborough, UK).

In order to determine the differences observed in the sub-daily variations of PM₁₀ concentrations observed on 12 February and 13 May 2014, as a first step, the temporal evolution of the five-minute PM₁₀ concentration and five-minute mean and maximum wind speed were analyzed (Figure A1). It can be seen that, on 12 February 2014, the increase in PM₁₀ concentration occurred when wind speed was relatively low (below 4 m s⁻¹ on average), which shows that dust was not produced locally. On the opposite, on 13 May 2014, the sudden increase in PM₁₀ concentration occurred almost at the same time as wind speed increases (up to ~12 m s⁻¹ for five-minute mean wind speed and up to ~17 m s⁻¹ for five-minute maximum wind speed), which suggests that dust was at least partially due to local wind erosion.

In order to confirm the regional or local origin of dust during these two days, we combined back-trajectory analysis and space-borne measurements. The origin of the air-masses was determined using the online version of the HYbrid Single Particle Lagrangian Integrated Trajectory model (HYSPLIT; [53,54]). This method is indeed widely used to determine the origin of African dust contributions to the total aerosol load (e.g., [55–57]) or the origin of mineral dust deposits (e.g., [33]). 72 h backward trajectories starting at 12:00UTC on both days were computed, ending at the ground level (0 m) at El Fjé, using the ensemble model (i.e., 27 backward trajectories were computed for all identified days) forced by the GDAS (Global Data Assimilation System) meteorological database at 0.5°.

In complement to the air mass trajectories, we used the AOD at 550 nm derived from the measurements of the Moderate Resolution Imaging Spectroradiometer (MODIS) on-board the Aqua satellite: 1° × 1° daily maps of the merged Dark Target/Deep Blue product available in the collection 6 MODIS aerosol products over land and ocean [58] were used to determine the most probable origin of mineral dust observed during both days. To do so, AOD maps were produced for each of the days composing the 72 h backward trajectories computed for 12 February and 13 May 2014, and the most probable origin of mineral dust corresponds to the location where the air-mass trajectory crosses near the surface a region with a high AOD.

For 12 February 2014 (Figure A2), back-trajectories indicate that the air-mass travelled close to the surface from the regions of the Chotts, located at the border between Algeria and Tunisia in the West, where AOD was high (between 0.6 and 0.8) on 11 February 2014. This analysis confirms that dust arriving at El Fjé on 12 February 2014 was not produced locally. For 13 May 2014 (Figure A3), back-trajectories arrived from the South, where AOD was low (between 0.1 and 0.2). On the other hand, AOD on 13 May 2014 was between 0.3 and 0.4 in the Jeffara plain, where the station of El Fjé is located, which confirms the local origin of the dust responsible for the high PM₁₀ concentrations recorded during this day.

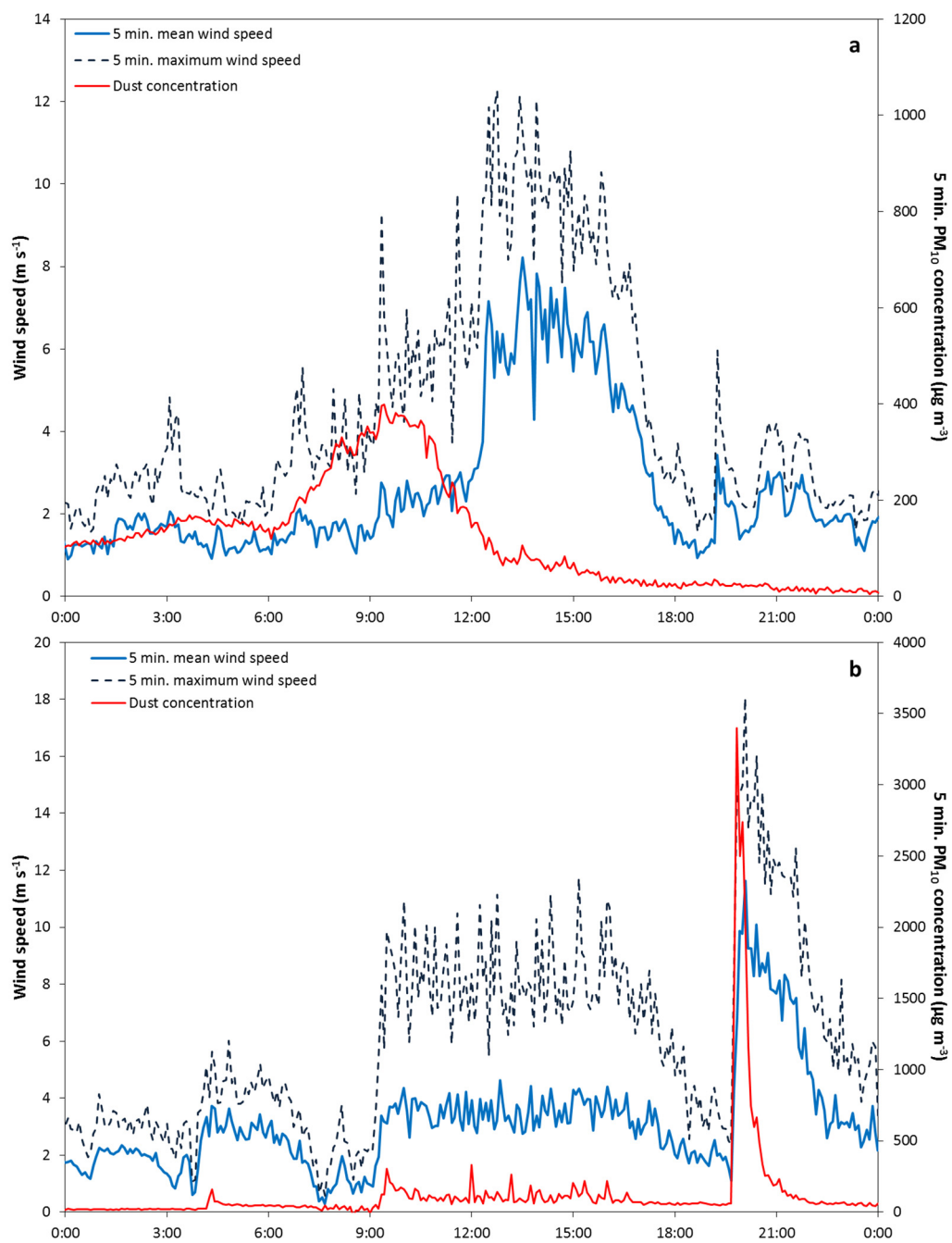


Figure A1. Temporal evolution of the five-minute PM₁₀ concentration (in µg m⁻³—red line) and five-minute mean (light blue line) and maximum (dark blue dashed line) wind speed (in m s⁻¹) for (a) 12 February 2014 and (b) 13 May 2014.

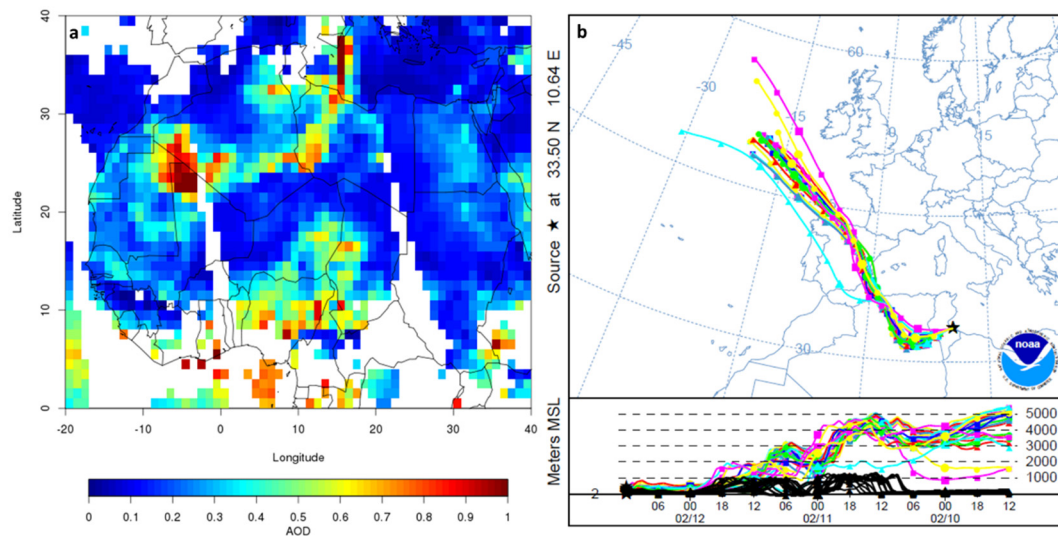


Figure A2. (a) Moderate Resolution Imaging Spectroradiometer (MODIS) (Aqua) merged Dark Target/Deep Blue AOD product at 550 nm on 11 February 2014; (b) 72 h backward trajectories ending at 1200 UTC 12 February 2014, computed using the online version of the Hybrid Single Particle Lagrangian Integrated Trajectory (HYSPLIT) model from El Fjé (black star—33.50N; 10.64E), and corresponding altitudes (in m) of the air-mass along its trajectory.

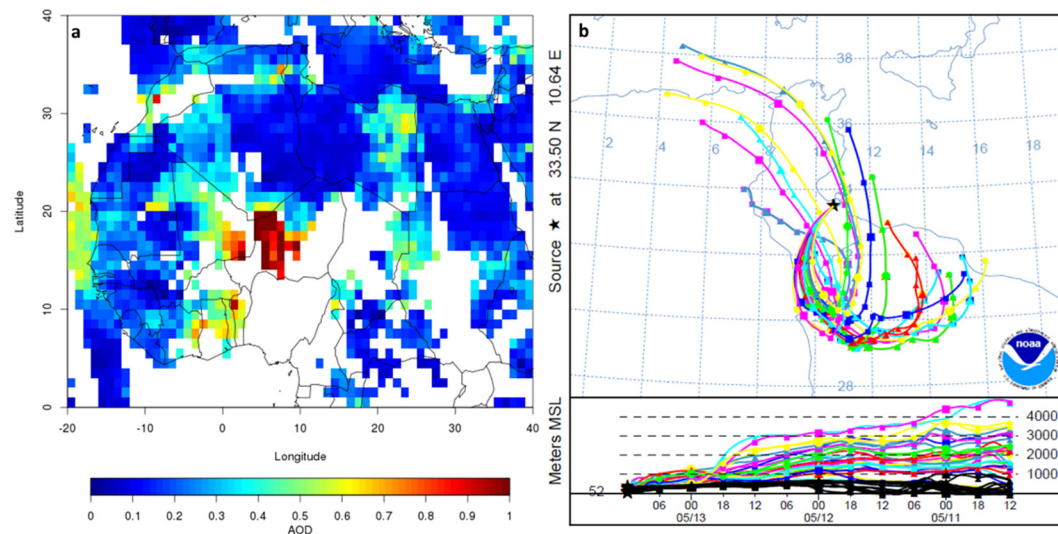


Figure A3. (a) MODIS (Aqua) merged Dark Target/Deep Blue AOD product at 550 nm on 13 May 2014; (b) 72 h-backward trajectories ending at 1200 UTC 13 May 2014 computed using the online version of the HYSPLIT model from El Fjé (black star—33.50N; 10.64E), and corresponding altitudes (in m) of the air-mass along its trajectory.

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