

1Can active sands generate dust particles by wind-induced processes?

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17**Abstract**

18Mineral dust emission is a major process in determining the global dust cycle. Surfaces
19composed of sand grains (dunes, sand sheets) cover more than 10 % of the Earth land
20surfaces, but also common on Mars. Active (dune) sands have been identified recently as
21dust sources in northern Africa, China, and elsewhere. Previous studies on dust emission
22from active sands suggested that dust can be generated by different aeolian mechanisms
23that are related to (i) re-emission of settled dust particles, (ii) clay coating removal, and (iii)
24abrasion of the sand grains. However, little empirical evidence of dust emission from active
25sands under natural conditions of wind (aeolian) transport has yet been reported. This study
26integrates wind tunnel experiments and high resolution laboratory sand analyses to explore
27aeolian dust emission from active sands with conditions simulating the natural processes of
28saltation. Sand samples from three sites with different characteristics of grain size, dust

29content, morphology, and mineralogy were used in the experiments. The aeolian
30experiments were conducted under various wind velocities. No dust emission was recorded
31for shear velocities below the saltation threshold. Increasing the wind velocity above the
32saltation fluid threshold caused an increase in atmospheric PM₁₀ concentrations, the
33magnitude of which depended on the specific shear velocity and the saltation flux. The initial
34content of dust-sized particles in the sand sample was found to influence PM₁₀ emission.
35Higher PM₁₀ concentrations were recorded from sand samples initially containing more than
362 % of dust-sized particles. The experiments identify clay coatings removal as the dominant
37mechanism over time of dust emission in typical active sand dunes (< 2 % dust content) with
38an addition of re-emission of existing dust-sized particles (< 63 μm). The rate of such re-
39emission is determined by the initial amount of dust-sized particles in the sand bed. The dust
40emission observed in this study indicates that, in addition to the classic dust sources of non-
41sandy soils, sand bodies should also be taken into consideration in determining global dust
42emission.

43**Keywords:** Aeolian processes; Dust sources; PM₁₀; Wind tunnel; Saltation

441. Introduction

45Aeolian (wind-driven) dust emission has a major impact on a variety of environmental and
46socioeconomic issues. Airborne dust particles can affect climate (Nenes et al., 2014; Kok et
47al., 2017), biogeochemical cycles (Jickells et al., 2005), and soil ecology (Okin et al., 2004;
48Field et al., 2010). Substantial loss of nutrients and clays by dust emission reduces the soil
49fertility, leading to soil loss and degradation (Katra et al., 2016a). Dust events significantly
50increase air pollution (Katra et al., 2014a; Krasnov et al., 2016) and thus can impact human
51health (Vodonos et al., 2015). Models estimate that the global dust emission rate is between
52~ 500 Tg yr⁻¹ and ~ 4000 Tg yr⁻¹ (Evan et al., 2015; Huneeus et al., 2010; Kok et al., 2014a,
532017; Shao et al., 2011). Comparisons of model results against dust measurements still show

54large discrepancies (Evan et al., 2014; Huneus et al., 2010; Kok et al., 2014b) due to a
55number of major gaps in our understanding of dust source dynamics and mechanisms of dust
56emission. It is commonly assumed that dust sources consist of soils rich in clay and silt sized
57particles (< 63 μm in diameter). These fine particles are subjected to cohesive inter-particle
58forces and therefore rarely occur as loose particles in soil but as part of aggregates.
59Therefore, impacts by saltating particles (sandblasting) have been found to play a major role
60in dust emission from soil aggregates (Alfaro et al., 1997; Kok et al., 2012, 2014a; Shao et al.,
611993; Shao, 2008; Swet and Kutra, 2016).

62 Little attention has been paid to the contribution of active sand dunes as dust sources.
63The possibility to generate dust, i.e., clay (< 2 μm in diameter) and silt (between 2 and 63 μm
64in diameter) sized particles from active sands has been suggested over the years. Active sand
65refers to un-stabilized (loose) sand-sized particles that are available for wind transport. Most
66studies dealing specific with active sand have proposed aeolian abrasion of the grains as the
67mechanism for dust generation (Bhattachan et al., 2012; Bullard et al., 2004, 2007; Crouvi et
68al., 2012; Sweeney et al., 2016; Wright et al., 1998). Aeolian abrasion refers to the reduction
69in the physical size and angularity of parent sands due to the impact of saltators at the sand
70bed or by particle collisions in the air (Bagnold, 1937; Jerolmack and Brzinski, 2010;
71Jerolmack et al., 2011; Kuenen, 1960). However, dust that is apparently generated by active
72sands may also be produced through other mechanisms: re-emission of dust previously
73trapped in dunes from exogenous sources (Muhs et al., 2008), and/or by the detachment of
74clay-rich coatings present on the surfaces of sand grains (Bullard and White, 2005). Studies
75have shown that many of the sand bodies worldwide consist of clays and iron oxides coatings
76above the sand grains (Walden and White, 1997).

77 A recent remote sensing study identified that over 40 % of dust storms in Northern Africa
78originate from areas covered by sand dunes (Crouvi et al., 2012). The occurrence of fine
79particle production from sand has also been deduced from field (Crouvi et al., 2008, 2012;

80Jerolmack and Brzinski, 2010; Jerolmack et al., 2011; Sweeney et al., 2016) and experimental
81(Bullard et al., 2004, 2007; Bullard and White, 2005; Kuenen, 1960; Smalley and Vita-Finzi,
821968; Whalley et al., 1982; Wright, 2001) studies. Field studies proposing aeolian abrasion as
83the primary generator of dust particles are based on identification of downwind fining of
84aeolian sediment. However, the observed spatial fining trends may also result from sorting or
85fractionation caused by differences in transportability of different grain sizes (Roskin et al.,
862014). In addition, the few existing studies on dust generation from active sand were
87performed under conditions that do not directly reproduce the natural processes of
88saltation. Thus, our understanding of aeolian dust emission from sands remains limited.

89 Sand dunes cover around 20 % of arid areas worldwide, and about half of them are
90considered as active sand dunes (Ashkenazy et al., 2012; Pye and Tsoar, 2009). Sand dunes
91are also a dominant formation covering wide areas of other planets as Mars and Venus
92(Claudin et al., 2006; Runyon et al., 2017). Typical active sand dunes are characterized by
93more than 98 % of sand-sized grains (63-2000 μm) with a size distribution mode of 200-300
94 μm (Ahlbrandt, 1979). In addition to sand dunes, there are other forms of active sand with
95different particle composition. Sandy soils contain relatively high percentages of clay-silt
96particles (up to ~ 10 %). Many of these arid soils are located in close proximity to dust
97sources and are subjected to aeolian deposition of airborne dust. Another sand form is
98mega-ripple fields composed of fine sand and very coarse sand with a mode of up to 2000
99 μm (Yizhaq and Katra, 2015). It can be hypothesized that different active sand compositions
100will respond differently to aeolian processes and produce different rates and types of dust
101emission over time.

102 Understanding the role of active sand as a dust source can provide a more accurate
103estimation of quantities and particle characteristics of global dust loading to the atmosphere,
104thereby reducing uncertainties in chemical transport and global climate models. It can also
105contribute to our understanding of sand transport and landscape development on both Earth

106and Mars. The aim of this study is to quantify dust emission from active sands under
107different conditions simulating the natural processes of saltation. The study integrates
108targeted laboratory experiments and sand analyses to fill this apparent research gap.

1092. Materials and Methods

1102.1. Sand samples

111Three samples of active sand were utilized to represent different sand particle compositions.
112Sand was collected from two dunefields in the northwestern Negev (N_1 and N_2), Israel, and
113from Oceano Dunes, California (C_1). In both sites there is an ongoing in-situ study of dust
114emission. The Negev dunefield is located in the eastern part of the Sinai-Negev Erg (Fig. S1).
115Currently some dunes are partially stabilized by biological crusts, but their crests are still
116active (Tsoar et al., 2008; Zaady et al., 2014). The Negev dune sand has a typical size of sand
117for active dunes (mode at $\sim 250 \mu\text{m}$; Roskin et al., 2014). The N_1 sample was taken from an
118active linear sand dune, and contains less than 2 % (by volume) of clay and silt-sized
119particles. N_2 was sampled in sand at the northernmost edge of the Negev dunefield. The
120sand of N_2 is composed of active sand with relatively high percentages of silt and clay sized
121particles ($< 63 \mu\text{m}$) of up to 10 %. The higher amount of dust in N_2 compared with N_1 is due
122to the proximity of N_2 to the Negev loess plane. Nevertheless, this region is associated with
123particle-size fractionation of aeolian sand transport along the Sinai-Negev erg (Roskin et al.,
1242014).

125 The Oceano dunefield on the Central Coast of California (Fig S1) was formed by strong
126onshore sea breezes transporting sand derived from fluvial deposits (Cooper, 1967), and thus
127contains a mixture of quartz, feldspar, and other minerals (Huang et al., 2018; Bedrossian
128and Schlosser, 2007). The sample from C_1 is composed of relatively coarse sand particles
129(mode $> 400 \mu\text{m}$) with low amount ($< 1 \%$) of dust sized particles (Huang et al., 2018; Martin
130et al., 2018). Sand samples from each site were taken from the upper 2-cm layer of the dunes
131for wind tunnel experiments and laboratory analyses.

1322.2. Aeolian experiments

133 Laboratory wind tunnel experiments were performed to quantify dust emission from the
134 sand samples. The experiments were conducted under various wind velocities, above and
135 below the saltation threshold, to examine two components of dust emission: re-emission of
136 loose dust particles in the sand samples by direct aerodynamic lifting (no saltation), and dust
137 emission caused by saltation impacts onto the sand surface. For each wind velocity and sand
138 sample, the wind profile was measured at different heights (cm) above the tunnel bed: 2,
139 3.5, 5, 7.5, 10, 15, 20, 25, 30, 35, 40, and 45 (Fig. S2). These wind profiles were used for
140 determining shear velocities (u_* , m s^{-1}) following the logarithmic *law of the wall*.

141 The aeolian experiments were performed using the boundary-layer wind tunnel of Ben-
142 Gurion University (BGU) described in Katra et al. (2014b). The BGU wind-tunnel is an open
143 circuit tunnel consisting of three parts: an entrance cone, a test section, and a diffuser (Fig.
144 S2). Air is sucked in through the bell-shaped entrance by a fan located at the end of the
145 diffuser. The cross sectional area of the tunnel is $\sim 0.7 \times 0.7$ m and the working length is 7 m
146 for measurements in the test section. The boundary layer in the wind tunnel is ~ 12 cm
147 above the tunnel bed (Fig S2). For each experimental run at a specific shear velocity, the
148 saltation flux remains constant ($\pm 10\%$) and does not fade or intensify over time (Katra et al.,
149 2014b; Schmerler et al., 2016). Instruments installed in the wind tunnel enable the
150 determination of the following parameters (Fig. S2B): (i) wind velocity in vertical and
151 horizontal cross sections by micro-vane probes (www.kimo.com) for calculation of shear
152 velocity (u_*); (ii) collection of saltating sand grains by an array of traps oriented along the
153 wind direction for calculating average saltation mass flux ($\text{kg m}^{-1} \text{s}^{-1}$) over time. The traps
154 were placed at heights of 2.5, 4.5, 6.5, 8.5 and 10.5 cm above ground, and each trap had a
155 cross-sections of 2×1 cm; (iii) dust concentrations ($\mu\text{g m}^{-3}$) of particles that are less than 10
156 μm in aerodynamic diameter (PM_{10}) recorded by a light-scattering device, DustTrak DRX 8534
157 (www.tsi.com), in the range of 0.001–150 mg m^{-3} ($\pm 0.1\%$ of reading) at 1-second intervals

158 placed at 25 cm above the tunnel bed; (iv) collection of suspended dust by active (isokinetic
159 filter) gravimetric samplers that include a pump to maintain a constant flow and an inertial
160 Anderson impactor (Andersen Instruments Inc., USA) for dust characteristic analyses.

161 In each experiment, the sand was placed in a ~ 3-cm thick layer on the full length of the
162 wind tunnel bed. The first test was conducted under a free stream wind speed of 4 m s^{-1} ,
163 corresponding to a shear velocity of 0.28 m s^{-1} , below the saltation threshold for each
164 sample. The test was run for a relatively short time of 900 seconds. The second test was run
165 under higher wind shear velocities and above the saltation threshold of the different
166 samples, at $u_* = 0.30\text{--}0.36 \text{ m s}^{-1}$ ~ 5 m s^{-1} to ~ 8.5 m s^{-1} , measured at 25 cm above the tunnel
167 bed). In this case, dust emission can be a result of sand abrasion and/or removal of coatings,
168 but also by aerodynamic lifting of loose particles that are held between the coarser sand
169 grains and may be released upon their movement or impacts during the saltation transport.
170 The time duration of each experiment (shear velocity) was up to 9000 seconds (150 min),
171 which is much longer than a single wind shear velocity would typically be sustained in the
172 field. Wind events can last for hours, but the cumulative time of specific shear velocity at a
173 specific direction will be significantly shorter.

174 Before each experiment, the PM_{10} background levels were measured inside the tunnel to
175 account for noise in the measured PM_{10} signal. The measured background levels (~ $0.30 \mu\text{g}$
176 m^{-3}) were subtracted from the data recorded during the experiment. In order to optimize the
177 measurement procedure, the sand was manually recycled in the tunnel during these long
178 tests to allow a sufficient sand supply and ensure a saturated airstream and steady-state
179 saltation. Each test was repeated 3 times to determine the mean values of saltation and dust

180 emission. The recorded PM_{10} concentrations were converted into mass flux (F_{PM})
181 emitted from the soil surface ($\text{kg m}^{-2} \text{ s}^{-1}$) based on the wind tunnel dimensions and area of
182 the sand bed:

183

$$(1) \quad F_{PM} = C_{PM} V_t / (A_p t)$$

184Where C_{PM} is the recorded PM concentrations ($\mu\text{g m}^3$), V_t is the volume air in the
185wind tunnel (3.43 m^3), A_p is the area of the experimental plot (4.9 m^2), and t is time
186(in Seconds) see Katra et al., 2016b. The PM_{10} ($\text{kg m}^{-2} \text{ s}^{-1}$) was used to calculate the
187sandblasting efficiency a (m^{-1}):

188

$$(2) \quad a = F_{PM} / Q$$

189Where Q ($\text{kg m}^{-1} \text{ s}^{-1}$) is the total horizontal sand flux integrated over all sand grain sizes (see
190Kok et al., 2014a).

191 All of the above procedures were performed also on dust-free 'clean' sand to separate
192between the mechanisms of dust emission. The raw sand (bulk samples) underwent a series
193of gentle rinsing and washing to remove the loose dust-sized particles. Following the results
194obtained for the bulk samples (see section 3; Fig. 4), in which the dust emission of C_1 sample
195stopped after a period of time (reduced to the background values), and following a
196preliminary experiment on 'clean' sand from C_1 sample, in which no dust emissions were
197detected, the wind tunnel experiments on 'clean' sand were conducted only for N_1 and N_2
198samples. 2.3. Particle analyses

199Physical and chemical properties of the sand (from the tunnel bed before the aeolian
200experiments and from the sand traps during the experiments) and of the dust (collected
201during the experiments) were analyzed in the laboratory.

202 The Particle Size Distribution (PSD) was analyzed using an ANALYSETTE 22 MicroTec Plus
203(Fritsch) laser diffractometer, which measures particles in the size range of $0.08\text{--}2000 \mu\text{m}$.
204PSD data were calculated using the Fraunhofer diffraction model with a size resolution of 1
205 μm using MasControl software. The software was employed to determine the mean

206 diameters, median diameters, modes of multi-modal distributions, sorting values, and size
207 fraction weights. Mineralogical composition was analyzed using the X-ray power diffraction
208 (XRPD) method (Philips 1050/70 power diffractometer). A Panalytical Empyrean Powder
209 Diffractometer equipped with position sensitive detector X'Celerator was used. Data were
210 collected in the $\theta/2\theta$ geometry using Cu K_{α} radiation ($\lambda=1.54178 \text{ \AA}$) at 40 kV and 30 mA.
211 Scans were run during ~15 min in a 2θ range of $4-60^{\circ}$ with step equal to $\sim 0.033^{\circ}$. Elemental
212 composition analyses were performed by the X-Ray Fluorescence (XRF) method using an XRF
213 spectrometer PANalytical Co., model Axios (wavelength dispersive -WDXRF, 1kW). The
214 Omnia software was used for the quantitative analysis. Morphological and chemical
215 characteristics of the particles were examined using a Scanning Electron Microscope (SEM)
216 (Quanta 200, FEI). The high magnification ($6 \times$ to $> 1,000,000 \times$) enabled the analysis of the
217 smallest dust particles ($< 2 \mu\text{m}$). Chemical analysis in this device was performed using the
218 Energy Dispersive X-ray Spectroscopy (EDS). Sand-grain roundness was assessed for each
219 SEM image using the grain roundness chart of Powers (1953).

2203. Results

221 The PSDs of the three bulk samples used in the aeolian experiments are presented in Fig. 1.
222 All the samples are characterized by a distribution with a single mode in the range of sand-
223 sized particles. However, there are significant differences ($P \leq 0.05$) in the size mode and in
224 the initial dust content between the samples. N_1 contains a relatively high percentage (58.7
225 %) of medium-sized sand (250-500 μm), whereas N_2 is characterized by a relatively large
226 amount (64.4 %) of fine sand (63-250 μm) compared to the N_1 dune (23.5 %). C_1 has a much
227 coarser composition with 44.7 % of sand larger than 500 μm . All the samples contain dust-
228 sized particles ($< 63 \mu\text{m}$) that can be found between or attached to the sand grains. N_2 dune
229 can be considered as a "dusty" sand sample with 8.00 % content of dust-sized particles as
230 opposed to only 1.81 % in N_1 and 0.95 % in C_1 (Fig. 1). In all the samples, over 60 % of the

231 dust sized fraction is fine particles ($< 20 \mu\text{m}$), which are subject to long-term suspension (Kok
232 et al., 2017). The PM_{10} part out of the dust content is 64 % in N_2 and ~ 40 % in N_1 and C_1
233 samples.

234 Mineralogical analyses (XRPD) of the samples show that N_1 and N_2 consist of over 90 %
235 quartz sand grains, while the C_1 sample is a mixture of quartz (45 %) and feldspar (K-silicate
236 30 % and Na-silicate 22 %) grains. From the SEM images it seems that N_1 and N_2 sand grains
237 are characterized as sub-rounded grains with a relatively smooth surface (Fig. 2A, B). C_1 is
238 composed of mostly sub-angular and angular sand grains (Fig. 2C). The feldspar sand grains
239 look more angular and their surfaces are more abraded compared with the surfaces of the
240 quartz sand grains (Fig. 2C). Clay and iron-rich coatings are found on top of the sand grains in
241 all of the tested samples (Fig. 2D, E, F). Clay minerals were found also as part of the loose
242 dust-sized particles ($< 63 \mu\text{m}$) within the sand samples (Fig. 2A).

243 Subjecting the bulk N_1 , N_2 , and C_1 samples to a range of wind velocities in the boundary
244 layer wind tunnel (Fig. 3) revealed a distinct pattern in the measured atmospheric PM_{10}
245 concentrations ($\mu\text{g m}^{-3}$), depending on initial dust content in the sand sample, shear velocity,
246 and saltation flux (Fig. 1; Table 1). At low wind shear velocities below the saltation threshold
247 of all samples ($< 0.29 \text{ m s}^{-1}$), no PM_{10} emissions were recorded (Figs. 3A, C, E). The threshold
248 shear velocities were measured by a careful and gradual increase of the wind velocity in the
249 tunnel to the moment of which the sand grains entered saltation transport. The recorded
250 thresholds were 0.29 m s^{-1} (N_2), 0.30 m s^{-1} (N_1), and 0.33 m s^{-1} (C_1). Notably, the wind-tunnel
251 observed threshold at C_1 is similar to the 0.32 m s^{-1} fluid threshold shear velocity calculated
252 independently from field measurements by Martin and Kok (2018). At a wind shear velocity
253 of 0.30 m s^{-1} , PM_{10} emission was recorded only in the N_2 sand (Fig. 3C) as a response to the
254 initiation of saltation transport (Table 1). In the N_1 and C_1 samples, this wind was not
255 sufficient for dust emission (Fig. 4A, E). In the N_1 sample, only a small amount of sand grains
256 were ejected into saltation, while no sand transport was observed in the C_1 sample (Table 1).-

257 Increasing the wind shear above the saltation threshold ($\geq 0.33 \text{ m s}^{-1}$) resulted in dust
258 emission and enhanced PM_{10} concentrations for all sand samples. For each constant shear
259 velocity experimental run, the dust emission over time was characterized by a distinct
260 pattern of an initial sharp rise in PM_{10} concentrations, followed by a gradual decline until
261 stabilizing at low values (Fig. 3B, D, F). However, clear differences in PM_{10} concentrations can
262 be detected between the sand samples (Fig. 3B, D, F). The average PM_{10} concentration
263 produced by the N_2 sample was ~ 8 times higher than by N_1 , although both sand samples
264 produced very similar saltation fluxes (Table 1). The saltation flux (Table 1) of the coarser
265 saltating particles of C_1 ($418 \text{ } \mu\text{m}$; Fig. 4) was found to be greater than in
266 N_1 and N_2 samples under shear velocity of 0.36 m s^{-1} , although the amount of particles
267 entering transport is expected to be lower than in N_1 and N_2 samples. However, the
268 calculated sandblasting efficiency (m^{-1}), which is the ratio of the dust emission flux ($\text{kg m}^{-2} \text{ s}^{-1}$)
269 to the sand saltation flux ($\text{kg m}^{-1} \text{ s}^{-1}$), is substantially smaller for C_1 than for the samples from
270 the other sites under all shear velocities. In all samples there was an increase in sandblasting
271 efficiency with shear velocity (Table 1). The highest efficiency obtained was for the N_2
272 sample, although associated saltation fluxes were similar to those from the N_1 sample. The
273 efficiency recorded for C_1 sample is considered as relatively low but with a close proximity to
274 those found in a field experiment in Oceano dunes (10^{-6} m^{-1} ; Huang et al., 2018). The
275 sandblasting efficiency reduces in all sand samples as the PM_{10} emission decreases over time,
276 while the saltation flux remains constant. The efficiency results obtained for all of the sand
277 samples (10^{-7} to 10^{-4} m^{-1}) were found as smaller than typical non-sandy soils (10^{-4} to 10^{-2} m^{-1})
278 (Kok et al., 2012).

279 Following the results of the bulk sand samples (Fig. 3), only N_1 and N_2 samples were
280 washed of loose dust particles to examine the emission mechanisms. The cleaning of the
281 sand samples did not have any mineralogical, chemical, or physical effect on the sand grains
282 or on the coatings on the grain surfaces (Fig. 5). The cleaning of the sand only reduced the

283 amount of dust-sized particles in the sand to a minimum of no more than 0.6 % in both N₁
284 and N₂ samples (Table. S1). The PM₁₀ concentrations produced from the 'clean' sand were
285 lower than those from the bulk samples (Fig. 6), while the saltation fluxes did not change
286 ($2.89 \times 10^{-3} \text{ kg m}^{-1} \text{ s}^{-1}$ for 'clean' N₁ and $2.98 \times 10^{-3} \text{ kg m}^{-1} \text{ s}^{-1}$ for 'clean' N₂). The resulted dust
287 emission from N₁ and N₂ 'clean' sand samples show similar PM₁₀ concentrations (red line, Fig.
288 6) with $56.5 \mu\text{g m}^{-3}$ and $60.5 \mu\text{g m}^{-3}$, respectively.

289 The dust emitted during the aeolian experiments was collected for laboratory analysis
290 (Fig. 7). The SEM images indicate that the emitted dust from the N₁ and N₂ bulk samples are
291 composed mostly of clay minerals (Fig. 7A, B). The chemical and mineralogical composition
292 of the emitted dust of the bulk samples was similar to that of the loose dust-sized particles
293 found between sand grains and to the coatings on top of the grain surfaces (Fig. 2). Only a
294 few isolated quartz fragments were found among the dust particles. In the C₁ sample, the
295 emitted dust consists of a mixture of clays, feldspar, and quartz particles, in comparable
296 quantities (Fig. 7C). The quartz fragments were relatively coarser (30-40 μm) than the
297 feldspar and the clay particles (< 20 μm). The analysis of the emitted dust from the 'clean'
298 sand samples (N₁ and N₂) show similar composition to those of the bulk samples, with mostly
299 clay minerals and only some single coarser quartz fragments (> 40 μm) (Fig. 7D, E).

3004. Discussion

301 By subjecting three distinctive natural sand samples to a range of wind strengths in a
302 laboratory wind tunnel, we were able to simulate the process of dust emission from active
303 sands during aeolian saltation. Throughout the experiments, it was found that dust emission
304 from the sand samples was directly associated with the occurrence of saltation transport,
305 where PM₁₀ emission occurs only in the presence of saltation (Fig. 3; Table 1). As such, direct
306 aerodynamic entrainment of dust was not detectable. Dust emission from active sand in our
307 experiments thus requires that wind strength exceeds the threshold shear velocity, which in

308turn depends on the surface PSD (Bagnold, 1937; Kok et al., 2012; Schmerler et al., 2016). In
309the N₂ sample, the PSD (mode of 251 μm) is finer than for the N₁ and C₁ samples (modes at
310342 μm and 461 μm, respectively), and therefore its threshold shear velocity is lower (0.29 m
311s⁻¹).

312 For a specific shear velocity and saltation flux, the dust emission flux appears to be
313primarily controlled by the dust-sized particle content of the sand surface. The results show
314that when the wind shear was strong enough ($\geq 0.33 \text{ m s}^{-1}$) the saltation flux of N₁ and N₂
315samples was similar (Table 1). However, the recorded PM₁₀ and therefore the calculated
316sandblasting efficiency were much higher for N₂ than for N₁ (Table 1). The reason for these
317differences can be explained by the higher initial content of dust in the N₂ sand sample. N₂
318contains relatively high amounts of dust-sized particles, especially PM₁₀ particles (Fig. 1). It is
319hypothesized that the dust flux emitted per unit horizontal saltation flux increases sharply
320with the content of fine particles (Kok et al., 2014a; Marticorena and Bergametti, 1995). In
321the C₁ sample, for which the highest saltation fluxes were recorded, the PM₁₀ concentration
322(and thus also sandblasting efficiency) was much lower. C₁ is composed of coarser sand with
323a mode of 417 μm (Fig. 1), and therefore the sand grains will enter into saltation transport
324only at higher wind velocities (Table 1), and thus dust emission will also be confined to higher
325wind velocities (Fig. 3), although the number of saltating particles for the C₁ sample can be
326much smaller than in N₁ and N₂ samples for a specific wind shear velocity. Therefore, the
327relatively low sandblasting efficiency of C₁, which is consistent with field measurements at
328the collection site (Huang et al., 2018), is likely related to the low initial PM₁₀ content (0.41
329%).

330 The PM₁₀ emission patterns observed in the wind tunnel experiments (Fig. 3) provide
331evidence for the relative importance of three possible dust emission mechanisms for sandy
332surfaces: (i) re-emission of previously settled dust particles in the sand (Muhs et al., 2008),
333(ii) clay coating removal from sand grains (Bullard and White, 2005), and (iii) abrasion of the

334 sand grains (Bhattachan et al., 2012; Bullard et al., 2007; Sweeney et al., 2016; Wright et al.,
335 1998). The sharp increase in dust concentrations obtained at the beginning of saltation (Fig.
336 B, D, F) can be generated from one or all of mechanisms listed above. However, the
337 subsequent gradual decrease in the PM₁₀ concentrations may indicate gradual exhaustion of
338 the limited supply of loose dust particles for direct re-emission as the saltation flux remains
339 the same over time (Zhang et al., 2016). From the results it seems that N₁ and N₂ samples
340 have comparable sand characteristics of mineralogy, grain roundness, and saltator PSD, in
341 addition to the similar saltation fluxes (Fig. 2; Fig. 4; Table 1); thus, no difference is expected
342 in the mechanism generating the dust emission. Therefore the differences observed in the
343 sandblasting efficiency and thus in the PM₁₀ emission (Table 1) can thus be related to the
344 higher initial content of loose dust-sized particles in N₂ (Fig. 1).

345 Comparing the PM₁₀ emission patterns of the bulk samples to those of the 'clean' sand
346 samples can provide further evidence for the relative importance of the different dust
347 emission mechanisms (Fig. 6). Both 'clean' sand samples of N₁ and N₂ emitted very similar
348 and relatively low amounts of PM₁₀ over time ($u=0.36 \text{ m s}^{-1}$), while the bulk samples showed
349 significant differences in PM₁₀ concentration in the beginning of each experiment (Fig. 6).
350 After a period of time when the loose dust is emitted, the dust emission from the bulk
351 samples reaches the minimum value of the 'clean' sand emission of $\sim 0.06\text{-}0.1 \mu\text{g m}^{-3}$ (N₁
352 after ~ 300 seconds; N₂ after ~ 7000 seconds, Fig. 3). The differences in PM₁₀ concentrations
353 found between the bulk samples (Fig. 3) can be related to the initial amount of loose dust-
354 sized particles in the sand (Fig. 1). Therefore it can be assumed that in typical dune sands like
355 the N₁ sample, which contains $< 2 \%$ of dust-sized particles, the re-emission of loose dust is
356 relatively minor (Fig. 6A) and the continuous PM₁₀ emission over time (Fig. 3B, D) is
357 controlled by clay coating removal and/or abrasion.

358 The analysis of the emitted dust particles collected during the aeolian experiments
359 provides further evidence for the relative contributions of the different dust emission

360mechanisms. The dust emitted from the N₁ and N₂ bulk samples consisted mostly of very fine
361particles of clay minerals (Fig. 7A, B), indicating similar primary dust sources from loose dust
362particles contained in the pore spaces among sand bed grains and from the coatings on these
363sand grains (Fig. 2). The fact that the dust emitted from the 'clean' sand of both N₁ and N₂
364samples had barely any PM₁₀ quartz particles (Fig. 7D, E), and that the clay dust particles are
365similar to the coatings found on top of the 'clean' sand grains (Fig. 5), [indicate](#) the dominance
366of the clay coating removal mechanism in these samples.

367 The kinetic energy reached by coarse grains (C₁) during saltation is higher than for finer
368grains (i.e., sand in the N samples) (Kok et al., 2012), thereby enhancing their potential for
369aeolian abrasion. In addition, the relatively sharp-edged grains of C₁ have greater potential to
370break during saltation to produce coarse dust particles. Saltation of rounded sand like N₁ and
371N₂ was found to be less efficient than saltation of angular sand at generating dust in abrasion
372(Bullard et al., 2004; Kuenen, 1960; Whalley et al., 1982; Wright et al., 1998). In typical active
373desert sand dunes, where quartz sand grains (N₁, N₂) tend to be smaller and more rounded
374(compared to coastal sites as C₁), aeolian abrasion is therefore suggested to play a very minor
375role as a dust generator. In addition, the relatively large-sized quartz dust particles (20-63
376μm) that may be released by abrasion will not suspend for long distances in a wind event as
377the fine dust (< 20 μm) (Kok et al., 2017; Mahowald et al., 2014; Nenes et al., 2014).
378Consequently, dust emission by aeolian abrasion is likely to play a relatively small role in
379global dust emissions.

380 The dust emission flux (Table 1) recorded from all of our sand samples are considered as
381very low compared to those produced by many other global dust sources. For example, the
382results obtained for N₂ sample were 10 times lower than those received during aeolian wind
383tunnel experiments in natural (undisturbed) Loess soils (northern Negev-Israel), which
384contain more than 40 % dust-sized particles under similar wind velocity of ~ 7 m s⁻¹ (Swet
385and Katra, 2016; Tanner et al., 2016). However, even the lower PM₁₀ concentrations from

386active sands can be significant when considering the wide extent of dune fields around the
387globe. A quantitative assessment of the potential of dust emission from global active sand
388dunes is thus needed to establish its contribution to the global dust cycle.

389 It should be noted that the aeolian saltation and dust emission in our experiments differ
390from natural settings in two key ways. First, whereas our experiments sustained a constant
391wind velocity and direction over a long duration to utilize the full emission potential of the
392sand bed, typical wind gust events that enable dust emission are significantly shorter in time.
393Second, whereas dust was only emitted from the wind tunnel during any particular
394experimental run, surface dust supply in natural sand dunes can be renewed by deposition of
395dust originating from nearby source areas. Thus, the depletion of dust under sustained wind
396and non-renewing conditions may have led to lower dust emission rates in our experiments
397than in similar natural settings.

3985. Conclusions

399Large discrepancies in global dust emission models arise from a number of major gaps in our
400understanding of the dust emission mechanisms from different source areas. This study
401utilized aeolian experiments to explore the potential for dust emission from sands containing
402different sample compositions, and to distinguish the different mechanisms of dust
403generation from sand. We provided empirical evidence that dust can be emitted from active
404sands under natural conditions of saltation, where significantly higher PM₁₀ concentrations
405were generated from sands that initially contained more than 2 % dust.

406 The results obtained in this study provide insight into the dust generation mechanisms
407from active sand dunes. Our results indicate that the dominant dust emission mechanism
408over time for typical active sand dunes (< 2 % dust content) is clay coatings removal, with a
409relatively small contribution from re-emission of loose-settled dust. In sands containing

410higher amounts of dust-sized particles, the relative contribution of the re-emission
411mechanism increases drastically.

412 Despite the commonly accepted hypothesis for dust emission from active sands by the
413aeolian abrasion mechanism, this study suggests, based on analyses of emitted dust
414particles, that abrasion has only a minor contribution to dust generation from active sands,
415and largely produces coarse dust particles ($> 30 \mu\text{m}$). Although the dust emission rates from
416sand recorded in this study are lower in comparison to emission rates from classic dust
417sources of non-sandy soils, the spatial extent of sand bodies is substantial, such that they
418should be taken into consideration in determining global dust emissions. Further analyses of
419the characteristics of dust emitted from sand dunes, such as chemical composition and size
420distribution, are needed for better representation of dust in climate models.

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601 **Tables**

	0.28	0.30	0.33	0.36
N ₁ saltation	0.00	1.46×10 ⁻⁴	1.27×10 ⁻³	2.13×10 ⁻³
N ₂ saltation	0.00	1.22×10 ⁻³	1.19×10 ⁻³	2.29×10 ⁻³
C ₁ saltation	0.00	0.00	8.53×10 ⁻⁴	5.68×10 ⁻³
N ₁ PM ₁₀	0.00	0.01 (1.66×10 ⁻¹⁰)	13.64 (9.17×10 ⁻⁹)	66.08 (4.52×10 ⁻⁸)
N ₂ PM ₁₀	0.00	28.65 (5.66×10 ⁻⁹)	248.34 (8.92×10 ⁻⁸)	1065.86 (3.24×10 ⁻⁷)
C ₁ PM ₁₀	0.00	0.00	4.00 (6.67×10 ⁻¹⁰)	23.36 (2.03×10 ⁻⁸)
N ₁ efficiency	N/A	1.73×10 ⁻⁷	4.64×10 ⁻⁶	1.37×10 ⁻⁵
N ₂ efficiency	N/A	3.88×10 ⁻⁵	7.03×10 ⁻⁵	1.52×10 ⁻⁴
C ₁ efficiency	N/A	N/A	7.81×10 ⁻⁷	3.58×10 ⁻⁶

602 **Table 1.** Saltation flux by mass (kg m⁻¹ s⁻¹) of the bulk and the ‘clean’ sand; average
603 atmospheric PM₁₀ concentration (µg m⁻³) due to emitted PM₁₀ flux (kg m⁻² s⁻¹) from the bed
604 (average PM₁₀ flux in brackets); and sandblasting efficiency (m⁻¹) during the aeolian

605 experiments under the different shear velocities ($0.28-0.36 \text{ m s}^{-1}$). The background values
606 were subtracted from the PM_{10} concentrations and fluxes. Note the different duration of the
607 aeolian experiments, 900 seconds for $0.28-0.30 \text{ m s}^{-1}$ and 9000 seconds for $0.33-0.36 \text{ m s}^{-1}$. In
608 all the experiments, the saltation rate remained approximately constant ($\pm 10 \%$) while the
609 PM_{10} concentration and flux reduced over time.

610 **Figures caption**

611 **Fig. 1.** Average particle size distribution (PSD) of sand from N_1 , N_2 and C_1 sites obtained by the
612 laser diffraction technique. On the right side are statistical parameters of the distributions.
613 The sample PSDs are significantly different with $P < 0.05$.

614 **Fig. 2.** Scanning electron microscope (SEM) images of sand particles collected from N_1 (A), N_2
615 (B) and C_1 (C) sand samples. The red arrows in A point to dust-sized particles ($< 63 \mu\text{m}$) found
616 between the sand grains. In C the yellow arrows point to quartz grains, while the orange
617 arrows point to feldspar sand grains. D-F are close-up images of the coatings attached to
618 sand particles from N_1 , N_2 and C_1 , respectively. In the yellow box is a chemical composition
619 analysis (%) using SEM-EDS. The location of the EDS analysis is marked by an asterisk.

620 **Fig. 3.** PM_{10} concentrations [$\mu\text{g m}^{-3}$] following dust emission in the wind tunnel under various
621 shear velocities in N_1 (top), N_2 (middle), and C_1 (bottom). (A), (C) and (E) show results of the
622 experiments at the lower shear velocities (u^* of 0.28 and 0.30 m s^{-1}), for convenient display of
623 the results, the background values were not reduced from the lower shear velocities; (B), (D)
624 and (F) show dust emission over time (9000 seconds) at the higher shear velocities of 0.33 m
625 s^{-1} and 0.36 m s^{-1} . Note the different Y axis scales.

626 **Fig. 4.** Average particle size distribution (PSD) of the sand collected from the **saltation traps**
627 after the wind tunnel experiments ($u^* = 0.33 \text{ m s}^{-1}$) for N_1 , N_2 and C_1 dune samples. On the
628 right are statistical parameters of the distributions ($P \leq 0.05$).

629 **Fig. 5.** Scanning electron microscope (SEM) images of the clean sand N_1 (A) and N_2 (C). B and
630 D are close-up images of the coatings attached to a sand particle. In the yellow box is a
631 chemical composition analysis (%) using SEM-EDS. The location of the EDS analysis is marked
632 by the asterisk.

633**Fig. 6.** PM₁₀ concentration [$\mu\text{g m}^{-3}$] before (black) and after (red) loose dust removal by
634washing of the N₁ (A) and N₂ (B) samples under shear velocity (u_*) of 0.36 m s⁻¹. The
635background levels were subtracted from all measured PM₁₀ concentration levels.

636**Fig. 7.** Scanning electron microscope (SEM) images of the emitted dust collected during the
637aeolian experiments for shear velocity of 0.33 m s⁻¹ from N₁ (A), N₂ (B) and C₁ (C) samples. D
638and E are images of the emitted dust from the 'clean' sand of N₁ and N₂, respectively. The
639yellow arrows point to quartz fragments, while the orange arrows in C₁ point to feldspar dust
640size particles (< 63 μm). All the remaining particles are composed of clay minerals with some
641carbonates and metallic materials. F is a close-up of different types of dust particles in the
642samples.