

Author response on “Exploring the contributions of vegetation and dune size to early dune building using unmanned aerial vehicle (UAV)-imaging” by Marinka E. B. van Puijenbroek et al.

I would like to thank both reviewers for their positive comments and the helpful feedback, which helped us improve both clarity and impact of the MS. In this version of our manuscript, we addressed the comments of one of the reviewers (see below), thanked the reviewers in the acknowledgements and included information about the accessibility of the final dataset (in the marked-up version of the manuscript below).

Kind regards, also on behalf of all co-authors

Marinka van Puijenbroek

Comments Anne-Lise Montreuil

The new version of the manuscript is much better and clearer with the changes in the text and figures.

I have some minor remarks:

- Lines 198 and 427: it must be all in lower case letter for ‘photogrammetric’.
 - o *We thank the reviewer for noticing and changed the p to a lower-case letter.*
- Lines 430-431 are confusing and need to be split into 2 sentences.
 - o *We changed it into two sentences and rewrote the second part to clarify our results.*
- Lines 529-531: the meaning of ‘co-variation’ is not clear in this sentence.
 - o *Co-variation was not the correct word, we changed it to collinearity.*
- Lines 550-552 must be re-written.
 - o *We rewrote that sentence, to clarify our message.*

1 **Exploring the contributions of vegetation and dune size to early dune**
2 **development using unmanned aerial vehicle (UAV)-imaging**

3 Short running head: Dune size and vegetation

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Abstract

Dune development along highly dynamic land-sea boundaries is the results of interaction between vegetation and dune size with sedimentation and erosion processes. Disentangling the contribution of vegetation characteristics from that of dune size would improve predictions of nebkha dune development under a changing climate, but has proven difficult due to scarcity of spatially continuous monitoring data.

This study explored the contributions of vegetation and dune size to dune development for locations differing in shelter from the sea. We monitored a natural nebkha dune field of 8 hectares, along the coast of the island Texel, the Netherlands, for one year using an Unmanned Aerial Vehicle (UAV) with camera. After constructing a Digital Surface Model and orthomosaic we derived for each dune 1) vegetation characteristics (species composition, vegetation density, and maximum vegetation height), 2) dune size (dune volume, area, and maximum height), 3) degree of shelter (proximity to other nebkha dunes and the sheltering by the foredune). Changes in dune volume over summer and winter were related to vegetation, dune size and degree of shelter.

We found that a positive change in dune volume (dune growth) was linearly related to initial dune volume over summer but not over winter. Big dunes accumulated more sand than small dunes due to their larger surface area. Exposed dunes increased more in volume (0.81% -per dune per week) than sheltered dunes (0.2% -per dune per week) over summer, while the opposite occurred over winter.

Vegetation characteristics did not significantly affect dune growth in summer, but did significantly affect dune growth in winter. Over winter, dunes dominated by *Ammophila arenaria*, a grass species with high vegetation density throughout the year, increased more in volume than dunes dominated by

43 *Elytrigia juncea*, a grass species with lower vegetation density (0.43 vs. 0.42 (m³/m³)/week). The effect
44 of species was irrespective of dune size or distance to the sea.

45 Our results show that dune growth in summer is mainly determined by dune size, whereas in winter
46 dune growth was determined by vegetation. In our study area the growth of exposed dunes was likely
47 restricted by storm erosion, whereas growth of sheltered dunes was restricted by sand supply. Our
48 results can be used to improve models predicting coastal dune development.

49 **Key words:** Nebkha dunes, *Ammophila arenaria*, *Elytrigia juncea*, beach-dune interaction, landform
50 morphology, the Netherlands

51 **1. Introduction**

52 Coastal dunes occur along the sandy shores of most continents (Martínez and Psuty, 2008), and are
53 important to protect these coasts against flooding, provide areas for recreation, store drinking water and
54 shelter unique biodiversity (Everard et al., 2010). Coastal dunes and their services are threatened by
55 climate-induced sea-level rise (Carter, 1991; Feagin et al., 2005; Keijsers et al., 2016). However, dunes
56 also provide self-adapting systems of coastal protection, since the threat by sea-level rise can be
57 mitigated by the development of new dunes. Although the development of new dunes is well described,
58 we know little about the factors that determine the speed of early dune development. Understanding
59 these factors is essential for predicting dune development, and for safeguarding their services.

60 Dune development is the result of an interaction between vegetation and aeolian processes and
61 starts above the high-water line by the establishment of dune-building plant species (Maun, 2009). Once
62 vegetation establishes on the bare beach, it forms a roughness element that facilitates local sand
63 deposition and reduces erosion, forming a small dune within discrete clumps of vegetation (Dong et al.,
64 2008; Hesp, 2002). At the lee side of these small clumps of vegetation a shadow dune develops by sand
65 deposition, this shadow dune has a ridge parallel to the wind direction (Clemmensen, 1986; Gunatilaka
66 and Mwangi, 1989; Hesp, 1981). Vegetation and shadow dune together are known as nebkha dunes,
67 embryo dunes or incipient foredunes (Hesp, 2002; Hesp and Smyth, 2017). The further development of
68 these nebkha dunes strongly depends on the balance between summer accumulation of sand and
69 vegetation growth and winter erosion of sand and loss of vegetation (Montreuil et al., 2013). Summer
70 growth and winter erosion depend on weather conditions, such as wind speed, precipitation and storm

71 intensity (Montreuil et al., 2013; van Puijenbroek et al., 2017). As a result, net dune growth can differ
72 from year to year. Over time the smaller vegetated dunes can develop into an established foredune that
73 forms the first line of coastal defense against flooding.

74 Most research on coastal dune growth and erosion have focussed on processes and factors that
75 influence the supply of sand to the dunes and the effect of storm intensity on dune erosion (Anthony,
76 2013; Haerens et al., 2012; Houser et al., 2008; Keijsers et al., 2014; Saye et al., 2005; de Vries et al.,
77 2012). However, how coastal nebkha dune growth and erosion rates are influenced by the individual
78 dune characteristics, such as dune size, vegetation and degree of sheltering are less well studied. Dune
79 size affects the wind flow pattern, thus affecting sand deposition (Walker and Nickling, 2002) for
80 example increasing height or length of the shadow dune (Hesp, 1981; Hesp and Smyth, 2017). Dune
81 size also influences storm erosion: Claudino-Sales (2008) found that foredunes with a higher volume
82 were less sensitive to erosion. Whether the latter also applies to nebkha dunes, is unknown. Differences
83 in vegetation density between plant species are known to modify sand deposition (Arens, 1996; Hesp,
84 1983; Keijsers et al., 2014; Zarnetske et al., 2012), storm erosion (Charbonneau et al., 2017; Seabloom
85 et al., 2013), and dune morphology (Du et al., 2010; Hacker et al., 2012; Hesp, 1988). Sheltering by
86 other nebkha dunes can decrease the sand supply but can also reduce erosion by waves (Arens, 1996;
87 Lima et al., 2015; Luo et al., 2014; Montreuil et al., 2013). Although dune size, vegetation and
88 sheltering are known to be important for individual nebkha dune development, the relative contributions
89 of these factors are unknown.

90 In this study, we explored the contribution of vegetation and dune size to dune development.
91 Using an unmanned aerial vehicle (UAV) with camera we monitored a natural nebkha dune field for
92 one year. From the aerial images we constructed digital terrain models (DTM) and orthomosaics. From
93 the DTM's and orthomosaics we extracted detailed data on dune size (dune area, volume and maximum
94 height), vegetation characteristics and the degree of sheltering. We related changes in dune volume
95 (dune growth) to initial dune size, vegetation and sheltering over a summer (April - August) and winter
96 period (November - April). We expected that nebkha dune growth would be a function of vegetation
97 density, initial dune size, and shelter, with the function being modulated by season and degree of
98 shelter. We hypothesised that:

- 99 1. Nebkha dunes with high vegetation density grow faster irrespective of season or shelter.
- 100 2. In summer, growth of nebkha dunes is linearly related to initial dune size, with small
101 dunes growing at the same rate as big dunes. Exposed dunes grow faster than sheltered
102 dunes because of higher sand supply.
- 103 3. In winter dune growth is no longer linearly related to initial dune size, as small dunes are
104 more susceptible to storm erosion than big dunes. Exposed dunes grow slower than
105 sheltered dunes because of higher storm erosion.

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2. Methods

2.1 Study site

We monitored 8 hectares (200 m x 400 m) of a natural nebkha dune field with a large range of dune sizes at 'the Hors', the southern tip of the barrier island at Texel, the Netherlands, coordinates: 52°59'43.70"N, 4°43'47.53"E (Fig. 1). The Hors is a wide dissipative beach with a high degree of hydrodynamic reworking of the sand, which results in a high transport potential and opportunity for dunes to develop. In the last 5 years, between 2010 and 2015, many nebkha dunes have developed on the beach by plant species *Ammophila arenaria*, *Elytrigia juncea* or a mixture of both species. These three dunes with different species composition occur at similar distances from the sea, making this area ideal for exploring the effects of dune size and species composition on dune growth. *A. arenaria* and *E. juncea* differ in their vegetation characteristics: *A. arenaria* grows in dense patches, whereas *E. juncea* has a more sparse growth form. This difference in growth form probably also results into a different dune morphology: *A. arenaria* forms higher 'hummocky' shaped dunes, whereas *E. juncea* builds broader and lower dunes (Bakker, 1976; Hacker et al., 2012). The monitoring area is bisected by a low (maximum height of 7 m NAP, i.e. above the mean sea level near Amsterdam), continuous foredune ridge that runs parallel to the shore. The nebkha dunes that occur at the seaward side of this foredune are more exposed to the sea, while the nebkha dunes occurring at the landward side of the foredune are more sheltered from the sea, enabling us to explore whether the effects of dune size and vegetation are modified by the degree of shelter, especially since the age difference between the seaward and landward nebkha dunes is at most 5 years.

129 # Figure 1 approximately here #

130 **2.2 Weather conditions**

131 Summer conditions during our study period were similar to previous years, while winter conditions
132 were calmer than usual (Supplementary S1). The precipitation during the growing season was 276 mm,
133 and the average temperature in June and July was 16 °C. The most common wind direction was South
134 to South-West. The most common wind speed in summer was 4 - 5 m s⁻¹, and the maximum wind speed
135 was 13 m s⁻¹. In winter the wind speed was higher compared to summer, the most common wind speed
136 was 5 – 6 m s⁻¹ and the maximum wind speed was 19 m s⁻¹. We registered one storm during the study
137 period. This storm, however, could be classified as relatively weak. The highest water level was 211
138 cm NAP; compared to 248 cm NAP and 254 cm NAP from previous years. The storm, which was the
139 first of the season, occurred after the beginning of our mapping campaign.

140 **2.3 Data collection**

141 Three UAV flights in November (2015), April (2015) and August (2016) were carried out with a rotary
142 octocopter UAV system (Aerialtronics Altura Pro AT8 v1) and camera equipment of WageningenUR
143 *Unmanned Aerial Remote Sensing Facility* (Fig. 1). The octocopter was equipped with a Canon EOS
144 700D single-lens reflex camera with a 28mm f/2.8 Voigtländer Color Scopar SL-II N objective. The
145 camera sensor was modified to give a false colour output. The red channel of the camera had been
146 converted to be sensitive in the near-infrared, with centre point around 720nm. The blue channel of the
147 camera had been extended to also cover the UV region of the spectrum. The green channel was left with
148 almost original response. The false colour modification enabled the calculation of a modified

149 Normalised Difference Vegetation Index (NDVI), a commonly used measure for vitality and/or cover of
150 the vegetation (Carlson and Ripley, 1997). Aerial images were acquired by auto-piloted flights at an
151 altitude of 80 m at 4 – 5 m s⁻¹ velocity. The camera was set to take one image per second. The auto-
152 piloted flights enabled us to have the same flight paths for each of the three mapping campaigns. The
153 flight paths ensured that images had a minimum of 85% forward and 65% side-way overlap. Four
154 flights of 10 minutes were needed to cover the study area, yielding up to 900 RAW false colour images
155 per mapping campaign. Five ground control points were permanently placed in the flight area and
156 measured with a RTK-DGPS Trimble R6 Model 3 (TSC3) to calibrate our images with coordinates.
157 During our mapping campaign, a Spectralon reference panel was measured with our camera
158 immediately before take-off and after landing.

159 **2.4 Radiometric calibration**

160 In order to compare the images over the time, they were calibrated and converted from RAW to 16 bit
161 tiff format. First, we ensured that each individual pixel within an image was comparable, by converting
162 the RAW digital number into radiance units using a pixel-wise dark current and flat field calibration.
163 Second, each radiance image was calibrated to a reflectance factor image in order to correct for changes
164 in incident irradiance on different flight days. This calibration was done by using a Spectralon panel
165 with a known reflectance factor. The radiometric calibration is described in more detail by Suomalainen
166 et al. (2014).

167 The images were subsequently converted into NDVI images. Usage of the standard NDVI was
168 not possible due to lack of red channel in the false colour modified camera. Thus we used a custom

169 NDVI equation (Eq. 1), which was recommended by the company that modified the sensor. On their
170 website (MaxMax.com) this equation was shown to be just as effective for green vegetation as the
171 traditional NDVI formula ($R^2 = 0.77$) where the red band is taken as the absorption channel.

$$172 \quad 1) \quad NDVI = \frac{(NIR + G) - (2B)}{(NIR + G) + (2B)}$$

173 Where NIR, G, and B are the near-infrared, green and blue bands of the false colour image respectively.
174 For photogrammetric reconstruction, the NDVI image layer was stacked with the original green and
175 blue bands to form a three-color image.

176

177 **2.5 Photogrammetric reconstruction**

178 The large overlap between the consecutive images was necessary for photogrammetric software to
179 successfully process the aerial images into a 3D point cloud (Fig. 2). The 3D point cloud was generated
180 using Agisoft Photoscan Professional (v. 1.2.6), using the Structure-from-Motion (SfM) and Multi-
181 View Stereo (MVS) algorithms (Fonstad et al., 2013; Westoby et al., 2012). The correlated 3D points
182 are georeferenced to match the ground control points, and contain pixel intensity values of the input
183 imagery. From this 3D point cloud we interpolated a 5 cm pixel size digital surface model (DSM) and a
184 1 cm pixel size orthomosaic image. The DSM included also vegetation, which resulted in a vertical
185 error in dune height in areas where vegetation is present. We removed the vegetation from the point
186 cloud by identifying and removing the vegetation points. Vegetation points were removed by
187 distinguishing vegetation from sand using k-means clustering of the 3-D point cloud with NDVI using

188 the Hartigan and Wong (1979) algorithm in R (R Core Team, 2016). The holes in the point cloud that
189 arose by removing the vegetation were filled by using LAStools (the tool Blast2dem) (Isenburg, 2016),
190 which resulted in a Digital Terrain Model (DTM) without vegetation.

191 # Figure 2 approximately here #

192 We checked the accuracy of the **p**Photogrammetric reconstruction by measuring the vertical
193 error, the repeatability of the method and the degree in which NDVI predicted the biomass of the
194 vegetation. The vertical error of the DTM was assessed during a combined mapping and flight
195 campaign in August 2015 by measuring the elevation for 1100 points distributed over the flight area
196 with an RTK-DGPS Trimble R6 Model 3 (TSC3) and comparing the measured point measurements
197 with the DTM. The repeatability of the UAV photogrammetry was tested by repeating the same flight
198 path five times in November 2015 and comparing the similarity between the five DSMs. The NDVI
199 measurements were tested by clipping the vegetation flush with the sand surface for six *A. arenaria* and
200 seven *E. juncea* dunes and relating the biomass of the vegetation to the NDVI values.

201

202 **2.6 Defining dunes**

203 To be able to relate dune growth to characteristics of an individual nebkha dune including its shadow
204 dune, we first had to define individual dunes from the DTM. We followed a step-wise procedure for
205 each of our mapping campaigns (November, April, and August) using ArcGIS 10.3 (ESRI, 2016) that
206 resulted into different polygons in which each individual dune expanded or decreased in volume over

207 the study period. Dune volume and growth were later calculated using the same polygons for each
208 measurement campaign through time (see next section). To define the polygons we used the step-wise
209 procedure described below: 1) we constructed a baseline raster by calculating the average elevation in a
210 circle of 5m radius around each pixel in the DTM. A higher or lower radius resulted in either a too low
211 or too high baseline. 2) We then qualified pixels of the DTM as dunes, if they were 5 cm or higher
212 above a baseline raster, or had a slope of 15° or higher. The 5 cm threshold is the minimum that can be
213 accurately derived from the images and corresponds with visual estimates of nebkha dune foot; a slope
214 of 15° has been earlier identified by Baas et al (2002), as the slope for a shadow dune. From these
215 selected 'dune' pixels we created dune polygons. 3) Dune polygons of consecutive campaigns were
216 overlaid to construct the largest dune-covered area during the study period. 4) Each polygon was
217 visually checked for minimum size and presence of vegetation: dunes consisting of only one clump of
218 vegetation (0.4 m² or smaller) and dunes with no vegetation were discarded to derive conservative
219 estimates of nebkha dune volume and growth.

220

221 **2.7 Variables**

222 For each nebkha dune and for each mapping campaign we extracted dune volume (m³), max height (m)
223 and horizontal area (m²) from the dune polygons (see previous section) in the DTM. We calculated
224 changes in dune volume, i.e. absolute dune growth (m³/week) by subtracting the current dune volume
225 (V_t) from the volume of the previous mapping campaign (V_{t-1}), correcting for the number of weeks

226 between the mapping campaigns. To explore relationships irrespective of dune size, we also calculated
227 the relative dune growth ($m^3/m^3/week$).

228 We manually identified the species composition on each nebkha dune from the orthomosaic.
229 Species identification was verified in the field for a random subset of 100 dunes (23%) in May 2016. To
230 this end we created 2 transects from the southwest border to the northeast border of the area, along
231 which we determined the species on each nebkha dune. We compared the presence of species in the
232 field with the orthomosaic, and adjusted the species composition if necessary. In our dataset, dunes have
233 either *A. arenaria*, *E. juncea* vegetation, or a mixture of both species. A dune was defined as covered by
234 a mixture of both species, when it had distinct vegetation patches of both species present. For each
235 nebkha dune and mapping campaign we also extracted the vegetation density and the maximum plant
236 height. To assess vegetation density we first distinguished vegetated pixels from non-vegetated pixels
237 based on the orthomosaic using k-means classification of the NDVI using the MacQueen (1967)
238 algorithm. Hereafter, the vegetation area (m^2) and vegetation density ($NDVI/cm^2$ dune) were calculated
239 by summing the NDVI values of all vegetated pixels within the dune polygon (vegetation area) and then
240 dividing this summed NDVI by the total number of cm^2 pixels within the dune polygon (vegetation
241 density). The maximum plant height was calculated by subtracting the ~~DTM~~DSM (with vegetation)
242 from the ~~DSM~~DTM (without vegetation).

243 Sheltering can affect the sand supply and storm erosion. We used two methods to define the
244 degree of sheltering. Firstly, we distinguished whether a nebkha dune was seaward or landward from
245 the foredune. Secondly we determined how much the dune was clustered with other dunes. We

246 extracted the degree of clustering for each dune by calculating the mean height from the DTM in a 25 m
247 radius around the dune. All data extraction from the DSM, DTM and orthomosaic were done in R (R
248 Core Team, 2016).

249

250 **2.8 Statistical analysis**

251 First we explored if nebkha dune area, volume, maximum height, clustering (mean height in a 25m
252 radius around the dune), vegetation density and maximum plant height depended on species
253 composition using August 2016 data. As the number of dunes per species composition was unequal, we
254 used an ANOVA type III SS, to compensate for the unequal sample size (Fox and Weisberg, 2011) and
255 then used a Tukey HSD test (Hothorn et al., 2008) to determine significant differences between the
256 dunes with different species compositions.

257 Secondly, we tested how absolute changes in dune volume over winter (November – April) and
258 summer (April – August) periods related to the dune volume at the beginning of the period for locations
259 with different degree in sheltering with a linear regression model.

260 Thirdly, we analysed how the relative changes in dune volume over winter and summer
261 depended on dune size and vegetation characteristics in separated linear mixed models (Pinheiro et al.,
262 2016). To correct for spatial autocorrelation and species distribution we ran this analyses on a subset of
263 236 (54%) dunes. To this end we first explored the degree of spatial autocorrelation in our dataset by
264 creating a variogram. To account for the spatial autocorrelation of 25 m in our dataset we imposed a 50

265 m x 50 m grid over our study area; all dunes that were located within a grid cell (referred to as block)
266 were assumed to show spatial autocorrelation to some extent. This spatial autocorrelation was corrected
267 for in our statistical model by including block as a random intercept. We had 10 blocks seaward from
268 the foredune and 11 blocks landward from the foredune (Fig. 3), in which all species combinations
269 occurred (*A. arenaria* dunes, *E. juncea* dunes and *A. arenaria* + *E. juncea* dunes). By only including
270 dunes that were located within a block in the analysis, our selection was biased towards smaller dunes,
271 since larger dunes often fell within multiple blocks. We do expect that the effect of vegetation is more
272 apparent for these smaller dunes compared to larger dunes. To better distinguish between effects of
273 species compositions and vegetation structure we used two different models. The effect of species
274 composition was tested in a model with dune volume, maximum dune height, clustering and species,
275 whereas the effect of vegetation structure was tested in a model with dune volume, maximum dune
276 height, dune clustering, vegetation density and maximum plant height as explanatory variables. Within
277 each model we used the initial conditions for the explanatory variables, with initial conditions being the
278 values at the start of each measurement campaign. We included all two-way interactions. We selected
279 the best model by using Akaike information criterion (AIC). As we were mainly interested in the
280 importance of the explanatory variables relative to each other, we calculated the standardised estimates
281 for all the models by scaling the explanatory data.

282 The normality and homogeneity of the variance of the data was visually checked. All statistical
283 analyses were conducted in R (R Core Team, 2016). In the results we use statistic notation to show the
284 results of the ANOVA and linear regression models. We mention the F- value (ANOVA) or t-value
285 (linear regression), which indicates the difference of the explanatory variable to the variation in the data.

286 The p-value indicates the probability that the null-hypothesis is correct, we used a p-value of 0.05 as a
287 cut off to reject the null-hypothesis. The number in subscript indicates the degrees of freedom.

288

289 **3. Results**

290 **3.1 Nebkha dune characteristics**

291 Within the 8 hectare nebkha dune field we distinguished 432 polygons that were covered with nebkha
292 dunes for at least one moment during our mapping campaigns (Supplementary material S2). Half of
293 these dunes were covered by *E. juncea* vegetation (50.0%), followed by *A. arenaria* vegetation (28.2%)
294 and a mixture of both plant species (21.8%) in August 2016. Species composition of the dunes changed
295 along a gradient from sea to land. Close to the sea dunes were vegetated by *E. juncea*, while, further
296 from the sea, dunes were also vegetated by *A. arenaria* alone, or in a mix with *E. juncea* (Fig. 3).
297 Landward of the foredune dunes were also vegetated by *E. juncea*, *A. arenaria* alone, or a mix of both
298 species. The foredune bisecting our study area was mainly vegetated with *A. arenaria*.

299 # Figure 3 approximately here #

300 In August 2016 dune area, volume and maximum height differed significantly between nebkha
301 dunes differing in species composition (volume: $F_{2,426}=3.02$, $p=0.049$; max. height: $F_{2,426}=58.8$, $p <$
302 0.001), but did not differ between dunes contrasting in shelter. Dunes with a mix of *E. juncea* and *A.*
303 *arenaria* had overall the highest volume and maximum height, whereas dunes with *E. juncea* had the
304 lowest volume and height. Dunes with *A. arenaria* had the largest range in dune volume (Fig. 4A, B,

305 C). For dunes with *E. juncea* seaward from the foredune the distance between nebkha dunes was higher,
306 and thus clustering lower, than for to dunes with *A. arenaria* and dunes with both species ($F_{2,426}=51.5$,
307 $p<0.001$). The dune volume did not significantly differ between dunes seaward and landward from the
308 foredune (volume: $F_{1,426}=0.75$, $p=0.39$). In contrast, the dune height above NAP as well as the degree of
309 clustering (Fig. 4D) were significantly higher for dunes landward from the foredune (dune height:
310 $F_{1,426}=15.9$, $p<0.001$, clustering: $F_{1,426}=70.2$, $p<0.001$); we cannot exclude that part of these effects were
311 related to the slightly older age (max. 5 years) of the nebkha dunes landward of the foredune.

312 # Figure 4 approximately here #

313 For the statistical model with relative change in dune volume as response variable, we had to
314 correct for species distribution and spatial autocorrelation. We created a grid, with blocks of 50 m x 50
315 m, and we selected dunes that fell within a block. In total, we selected 236 dunes, which consisted of
316 41.95% of dunes with *E. juncea*, 36.02% of dunes with *A. arenaria*, and 22.03% of dunes with both
317 species. This subset of dunes had an overall lower dunes size compared to all the nebkha dunes in the
318 dune field, but had overall similar dune morphology and vegetation characteristics (Supplementary data
319 S3).

320 Vegetation characteristics depended on the plant species dominating the dunes and on the degree
321 of shelter. Nebkha dunes with *E. juncea* had significantly the lowest vegetation density, nebkha dunes
322 with *A. arenaria* the highest and nebkha dunes which consisted of both species had an intermediate
323 vegetation density (Fig. 4E, $F_{2,426}=48.91$, $p<0.001$). Similar to vegetation density, nebkha dunes with *E.*
324 *juncea* also had the lowest maximum plant height, whereas nebkha dunes with *A. arenaria* and

325 consisting of both species had the highest maximum plant height (Fig. 4F, $F_{2,426}=42.38$, $p<0.001$).

326 Nebkha dunes landward from the foredune had significantly higher vegetation densities compared to

327 seaward dunes ($F_{1,426}=45.49$, $p<0.001$), which is probably caused the calmer conditions landward from

328 the foredune, which benefits plant growth or the slightly older age of these nebkha dunes. There was no

329 significant difference in maximum plant height between nebkha dunes seaward and landward from the

330 foredune ($F_{1,426}=0.41$, $p=0.52$). Nebkha dunes with *E. juncea* had the smallest vegetation area

331 ($0.35\pm 0.047\text{m}^2$), nebkha dunes with mixed vegetation the largest vegetation area ($10.90\pm 3.05\text{m}^2$) and

332 nebkha dunes with *A. arenaria* have an intermediate vegetation area ($7.25\pm 4.18\text{m}^2$). The vegetation

333 area on a nebkha dune is larger landward from the foredune ($9.61\pm 3.96\text{m}^2$), compared to seaward of the

334 foredune ($2.04\pm 0.41\text{m}^2$). The vegetation area was correlated to dune volume (linear regression: $t_{430} =$

335 25.29 , $p < 0.001$), however this relationship was stronger for nebkha dunes landward from the foredune,

336 compared to nebkha dunes seaward from the foredune ($R^2 = 0.99$ vs. $R^2 = 0.69$).

337

338 **3.2 Change in nebkha dune number and volume**

339 The number of nebkha dunes within the measurement area changed over time, with nebkha dune

340 numbers declining over winter and increasing during summer. The degree of dynamics depended on

341 season, species and degree of sheltering.

342 **3.2.1 Summer**

343 Of the 434 nebkha dunes present in August 2016, 22.36% appeared over summer (April – August).
344 Most of these new dunes (65.93%) were *E. juncea* nebkha dunes, 31.87% were *A. arenaria* nebkha
345 dunes and only 2.20% were mixed dunes. Most (73.63%) new nebkha dunes developed seaward from
346 the foredune and were quite small in size with a volume of $2.72 \pm 0.29 \text{ m}^3$ (mean \pm SE). We assumed
347 that most of these dunes established over the growing season, as the orthomosaic showed a large
348 amount of wrack line material (plant material, woody debris, rope etc.) in their polygon in November
349 and April. However we cannot exclude that part of the large increase in the smaller *E. juncea* nebkha
350 dunes over summer is a result of their poor recognition in November and April.

351

352 Over summer, most nebkha dunes increased in dune volume, including the foredune which
353 increased over summer with 0.28% per week, reaching a volume of $64,444 \text{ m}^3$ in August. Only 4.16%
354 of the nebkha dunes showed a small decrease in the volume with a mean of $-0.041 \pm 0.014 \text{ m}^3/\text{week}$.
355 Changes in dune volume were positively related to the initial dune volume (Fig. 5A, $t\text{-value}_{428} = 57.11$,
356 $p < 0.001$) and were higher for nebkha dunes seaward of the foredune compared to nebkha dunes
357 landward of the foredune, resulting in a significant effect of shelter ($t\text{-value}_{428} = 2.72$, $p = 0.0069$). The
358 absolute changes in dune volume were also positively related to vegetation area, however this
359 relationship depended on the sheltering (vegetation area*sheltering by foredune: $t\text{-value}_{428} = 25.29$, $p >$
360 0.001). Nebkha dune vegetation area explained more variation in the change in dune volume for dunes
361 landward of the foredune, compared to dunes seaward of the foredune ($R^2 = 0.98$ vs. $R^2 = 0.36$).

362

363 # Figure 5 approximately here #

364 Compared to the absolute change in dune volume, the relative change in dune volume
365 ($\text{m}^3/\text{m}^3/\text{week}$) was mainly influenced by sheltering, with dunes seaward of the foredune growing faster
366 than dunes to landward of the foredune (Fig. 6A). We found no significant difference in relative change
367 in dune volume between dunes with different species composition (Fig. 6A, Table 1). In our statistical
368 model plant height had a statistically significant effect on the relative dune growth. However, when
369 tested in a single linear mixed model with block as random intercept, plant height had a R^2 of 0.0038,
370 thus hardly explaining any variation in relative dune growth (Table 2). Several dune size variables were
371 significant, but the individual variation explained by initial dune volume and dune height was very low,
372 their R^2 ranging between 0.05 – 0.0033. The significant interactions between variables were mostly
373 caused by the slight correlations between the explanatory variables. The clustering of nebkha dunes (i.e.
374 the average height within 25 m of each dune) did not significantly affect the relative dune growth. We
375 tested whether the effect of clustering was masked by the use of blocks as random intercept, since the
376 amount of clustering was different between the blocks. We re-analysed the data without the blocks as
377 random factor and again found no effect of clustering on the relative growth rate of dunes.

378 # Figure 6, Table 1 & 2 approximately here #

379 **3.2.2 Winter**

380 Over winter (November – April) 7.85% of the 344 nebkha dunes disappeared, of which 40.74% were
381 dunes with *E. juncea*, 55.56% were dunes with *A. arenaria* and 3.70% were dunes with both species.
382 These nebkha dunes disappeared both seaward (40.74%) and landward (59.26%) from the foredune and
383 were overall quite small with an average volume of $2.23 \pm 0.19 \text{ m}^3$.

384 Despite the decreasing number of nebkha dunes over winter, dunes increased in volume, the
385 large foredune even increased with 0.22% per week. However on average the change in absolute dune
386 volume was less positive than over summer, 21.30% of the dunes decreased -0.061 ± 0.015 (SE)
387 m^3/week in volume, particularly seaward of the foredune. 25.00% of these decreased dunes were
388 covered with *A. arenaria*, 50.00% with *E. juncea* and 25.00% with both species. The absolute change
389 in dune volume between November and April was positively related to the initial dune volume in
390 November (Fig. 5B, $t\text{-value}_{428}=2.12$, $p=0.034$), but was only significant for dunes landward of the
391 foredune. Dunes seaward of the foredune showed no relationship between absolute change in dune
392 volume and the dune volume in November (shelter: $t\text{-value}_{428}=-3.00$, $p=0.0029$). Similar to initial dune
393 volume, the vegetated area only explained variation in dune volume for the dunes landward from the
394 foredune (vegetated area * sheltering by foredune: $t\text{-value}_{428} = 16.17$, $p<0.001$).

395 The relative change in dune volume was influenced by species composition and degree of shelter
396 (Table 1). Nebkha dunes with *E. juncea* increased relatively less in volume than dunes with *A. arenaria*
397 (Fig. 6B); this effect was only significant for dunes seaward of the foredune. We found no significant
398 relationship between relative change in dune volume and vegetation density or maximum plant height
399 (Table 2). There was a significant interaction between vegetation density and sheltering by the
400 foredune, which could be related to the higher vegetation density at the dunes landward of the foredune.
401 Initial dune volume, and sheltering, had significant negative effects on the relative change in dune
402 volume, whereas clustering had a positive significant effect, but the relationships were very weak (R^2
403 between 0.002 – 0.05).

3.3 Net nebkha dune growth

Taken over the whole observation period November – August, the absolute nebkha dune growth (m^3/week) was higher at the seaward side of the foredune than at the sheltered landward side (slope seaward dunes: 0.37%, slope landward dunes: 0.25%, dune volume*position from foredune: $t\text{-value}_{428} = -11.7$, $p < 0.001$). Similarly, the relative dune growth ($\text{m}^3/\text{m}^3/\text{week}$) of the seaward dunes was also slightly higher than the landward dunes (seaward dunes: 0.27 ± 0.00009 (means \pm SE), landward dunes: 0.026 ± 0.0001 , $F\text{-value}_{1,230} = 18.51$, $p < 0.001$).

3.4 Accuracy of photogrammetric reconstruction

We checked the accuracy of the ~~Photogrammetric-photogrammetric~~ reconstruction by measuring the vertical error, the repeatability of the method and the degree in which NDVI predicted the biomass of the vegetation. The average vertical error was 7.3 ± 0.2 cm, with 80% of the measured points having a vertical error between -10 and 10 cm (Fig. S4.1). The vertical error increased with increasing distance from a ground control point. ~~The vertical error increased up to 20 cm for points that were 150 m at 150 m from a ground control point there was a vertical error of 20 cm~~ (Fig. S4.2). A vertical error of 10 cm could result in a deviation 3 – 6% in the dune volume, whereas the vertical error of 20 cm would result in a deviation of 5 – 12% in the dune volume (Table S4.1). The deviation depends however on the average elevation of a dune, a nebkha dune with a higher average elevation will have lower deviation of the vertical error than a nebkha dune with a low average elevation.

423 The source of error due to different conditions during consecutive mapping campaigns was
424 limited (Table S4.2). The difference between the DSMs of different flights with the same flight paths at
425 the same day was on average $3.9 \pm 3.9e^{-6}$ cm, with 80% of the raster cells of the DSM had a difference
426 between -0.07 and 0.07 cm (Fig. S4.3).

427 The degree in which NDVI represented vegetation biomass differed between species. The
428 summed NDVI of a nebkha dune with *A. arenaria* showed a trend with the biomass of *A. arenaria* ($t_4 =$
429 2.43 , $p = 0.07$, $R^2 = 0.6$), for nebkha dune consisting of *E. juncea* the summed NDVI was not
430 significantly related to the biomass of the vegetation ($t_5 = 1.43$, $p = 0.21$, $R^2 = 0.29$).

431 **4. Discussion**

432 The aim of this study was to explore the contributions of vegetation and dune size (i.e. initial dune
433 volume) to nebkha dune development expressed as change in dune volume. In addition, we were
434 interested in how- the effects of vegetation and dune size on nebkha dune development were modified
435 by the degree of shelter. Our results show that the contribution of vegetation and dune size depended on
436 season and degree of shelter. In summer dune volume change ($m^3/week$) was explained by initial dune
437 volume and to a lesser extent by dune height, while species composition, vegetation height or density
438 had no effect. In winter dune volume change was explained by vegetation and initial dune volume,
439 depending on the degree of shelter. Exposed nebkha dunes with sparsely growing *E. juncea* grew less in
440 volume than exposed nebkha dunes with densely growing *A. arenaria*. In contrast, growth of sheltered
441 nebkha dunes was a function of initial dune volume. These findings are the first to show that the effect

442 of vegetation and dune size on the nebkha dune development depends on season. These results can be
443 used to improve modelling of coastal dune development.

444 **4.1 Dune size**

445 **4.1.1 Summer growth**

446 We found a positive linear relationship between the initial dune volume and the absolute change in dune
447 volume over summer. It is known that nebkha dunes affect sedimentation by changing the wind flow
448 patterns (Dong et al., 2004; Li et al., 2008). Previous studies have found that with increased dune
449 volume the area where the wind speed is reduced increases, which result in higher sedimentation rates
450 (Hesp, 1981; Hesp and Smyth, 2017). The linear relationship between initial dune volume and dune
451 volume change found for the nebkha dunes in our study indicates that different dune sizes have similar
452 effect on the wind flow pattern per unit of area, which indicates scale invariance (Hallet, 1990). Scale
453 invariance has been used for modelling nebkha and foredune development (Baas, 2002; Durán Vinent
454 and Moore, 2013), but not yet been validated for nebkha dunes to our knowledge.

455 Our study focussed on a relatively small range in nebkha dune sizes. It is likely that the linear
456 relationship between dune volume change and dune size will saturate when dunes continue to grow and
457 processes other than wind speed reduction become important. The latter is supported by the volume
458 change of the low foredune bisecting our study area. Over summer the large foredune increased 0.28%
459 per week in volume, which is much lower than the overall increase of 0.81% per week of the dune
460 seaward of foredune. Therefore, we expect that there is a critical dune size at which the relationship
461 between dune volume and absolute dune growth is no longer linear. However, what exactly the critical

462 dune size is, is difficult to predict, it probably depends on multiple factors such as available sediment
463 supply and vegetation growth. The wind flow patterns are not only influenced by dune volume, but also
464 by maximum dune height (Walker and Nickling, 2002). In our study we found a significant, albeit weak
465 effect of the maximum dune height on the relative growth, suggesting differences in height did not have
466 a large effect on the wind flow pattern and the subsequent deposition of sand.

467 The positive linear relationship between dune volume and dune growth was modified by
468 sheltering; dunes landward of the foredune increased 0.60% per week less in volume than dunes
469 seaward of the foredune. This reduction in dune growth rate is likely the result of decreased sand supply
470 landward of the foredune; presumably a large amount of the sand was captured by the foredune as was
471 also observed for other foredunes (Arens, 1996). In our study the decrease in sand transport was less
472 sharp as observed by Arens (1996), however the difference in foredune sink strength between the
473 foredune in our study and those measured in Arens (1996) could be related to its smaller size, its
474 relatively low height and/or its sparse vegetation cover of 29% (Keijsers et al., 2015). Clustering of
475 dunes did not have any significant effect on the relative growth rate, which suggests that these smaller
476 dunes do not significantly reduce the sand supply to the landward situated dunes.

477 **4.1.2 Winter**

478 In winter initial dune size was only a good predictor for growth of the nebkha dunes occurring landward
479 of the foredune. For these sheltered dunes, increases in volume (m^3/week) again followed a linear
480 relationship with initial dune volume. The absence of a relationship between initial dune volume and
481 dune growth for the exposed dunes occurring seaward from the foredune, suggests that dune erosion is

482 less dependent on initial dune size than dune growth. Dune erosion has mainly been attributed to wave
483 run-up during storms (Haerens et al., 2012; Vellinga, 1982). Therefore, it seems reasonable to assume
484 that the degree of erosion depends on whether the dune can be reached by high energy waves. Large
485 dunes that are reached by high water levels can erode substantially, whereas small dunes can have no
486 erosion if they are protected by other dunes from the high water.

487 Interestingly, the sheltered nebkha dunes had a slightly higher dune growth in winter compared
488 to summer. This increase in dune growth for sheltered nebkha dunes can perhaps be explained by more
489 frequent and/or intensive aeolian transport events during winter resulting into higher sand supply to the
490 sheltered dunes (Davidson-Arnott and Law, 1990).

491 **4.2 Vegetation**

492 Vegetation characteristics were a poor predictor of dune volume change over the summer period, but
493 were a significant predictor for dune volume change over winter. Over summer dune growth did not
494 differ between nebkha dunes covered by different dune building plant species when corrected for dune
495 size. ~~Similarly~~Similarly, we did not find a clear effect of vegetation density and plant height on dune
496 growth. This results contrast with other studies that report a significant difference in the ability of
497 species to trap sand mediated by differences in shoot density and cover (Keijsers et al., 2015; Zarnetske
498 et al., 2012). Perhaps the discrepancy with our study can be explained by the differences in spatial scale
499 used between studies. We studied dune volume change at the scale of a nebkha dune including its
500 shadow dune, whereas the other studies focussed on the scale of the vegetation patch (Bouma et al.,
501 2007; Dong et al., 2008; Hesp, 1981, 1983; Keijsers et al., 2015; Zarnetske et al., 2012), where species

502 specifics effects are probably more pronounced than at the scale of the whole dune. Our results support
503 findings of Al-Awadhi and Al-Dousari (2013) who reported that the effects of vegetation on dune
504 growth are scale dependent for coastal nebkha dunes. They found that the linear relationship between
505 shrub vegetation characteristics and dune morphology levels off for bigger dunes. In our statistical
506 models we selected the smaller nebkha dunes, which was a consequence of only selecting dunes that
507 were located within one block. However even for these smaller nebkha dunes vegetation had no
508 significant effect on relative dune growth. The vegetated area of the nebkha dunes did have a positive
509 relationship with the change in dune volume, however this relationship could be caused by ~~eo-~~
510 variation/collinearity between the vegetated area and dune size, big dunes generally having a higher
511 vegetated area. Since initial dune volume was generally a better predictor for change in dune volume
512 than the vegetated area, our results suggest initial dune volume to be the better predictor for modelling.

513 Over winter nebkha dunes with *E. juncea* had a significantly lower relative growth rate than
514 nebkha dunes with *A. arenaria*, presumably because of their higher sensitivity to erosion. This species-
515 effect might be related to the sparser growth form of *E. juncea* in comparison to *A. arenaria* as dense
516 vegetation has been found to reduce the amount of dune erosion, by more effective wave attenuation
517 (Charbonneau et al., 2017; Koch et al., 2009; Silva et al., 2016). However, the effect of vegetation
518 density was not significant in our model suggesting that the species effect might be due to other species
519 differences, such as differences in rooting pattern. Another explanation is that the vegetation density
520 measurement did not reflect the real vegetation density, *E. juncea* was difficult to detect due to the low
521 NDVI values. The species effect was only significant for dunes situated at the exposed, seaward side of
522 the foredune where erosion by water likely occurred during the single storm covered by our study

523 period. Despite being statistically significant, the differences in relative growth rate between exposed
524 nebkha dunes with *A. arenaria* and *E. juncea* was not very large. Nevertheless the species effect might
525 become more pronounced with higher erosion pressure during more stormy winters (Charbonneau et al.,
526 2017).

527 Interestingly, our species did show differences in dunes size. On average, nebkha dunes with *A.*
528 *arenaria* were higher than nebkha dunes with *E. juncea*, that were broader (Bakker, 1976; Zarnetske et
529 al., 2012). This difference in nebkha dune morphology suggests a higher sand catching efficiency of *A.*
530 *arenaria*, as also suggested by (Zarnetske et al., 2012), this difference in sand catching efficiency might
531 have been masked by including the initial dune volume and maximum dune height as explanatory
532 variables. ~~that might be masked by using dune volume, mean height or dune area as explanatory~~
533 variables. We explored whether there is an effect of species composition on the change in maximum
534 dune height over summer, but found no consistent effect. Perhaps the difference in nebkha dune
535 morphology could be a result of differences in erosion between the nebkha dunes with different species
536 composition over winter.

537

538 **4.3 Application of UAV monitoring for nebkha dune development**

539 Measurements on the accuracy of the photogrammetric reconstruction shows that the vertical error is
540 between 0 cm – 20 cm, where most of the DTM pixels have a vertical error between 0 cm – 10 cm,
541 resulting into a deviation of dune volume between 3 – 12%. We do not expect this variation to affect
542 our results however, since the measurement error is random in nature and not systematic making

543 explanatory variables less significant rather than more significant. The vertical error increased with
544 increasing distance from the ground control markers, for future studies a maximum distance of 70 m
545 from each raster pixel to a ground control marker would be better than the 150 m we used. In our
546 statistical models for relative dune volume change ($\text{m}^3/\text{m}^3/\text{week}$) we accounted for the increasing
547 vertical error with increasing distance from the ground control marker by including blocks as a random
548 factor, since the nebkha dunes within a block have similar distances to a ground control marker.

549 The vegetation density, expressed as NDVI/cm² dune, was not significantly correlated with the
550 biomass. The poor relationship is likely a result of the low sample size (six or seven samples), in
551 combination with the high contribution of non-green parts, such as stems and dead litter, that give no or
552 weak NDVI signal. Since stems and dead litter do affect the wind flow pattern and attenuate waves, the
553 poor relationship between NDVI and biomass could explain why we did not find an effect of vegetation
554 density on dune growth and erosion. We did not measure the accuracy of the plant height, and can
555 therefore not say how well the maximum plant height represents the real plant height, however it is
556 probably an under-representation, since outliers are removed during photogrammetric processing.

557

558 **4.4 Implication for dune development**

559 **4.4.1 Net dune growth**

560 Exposed nebkha dunes had an overall higher net growth compared to sheltered nebkha dunes, indicating
561 that summer growth offset winter erosion in our study period which was characterised by an average

562 summer and calm winter. This balance might have been different if winter conditions had been more
563 severe.

564 During winter, storms determine the erosion of nebkha dunes seaward of the foredune. Multiple
565 low intensity storms can lead to more erosion than one high intensity storm (Dissanayake et al., 2015;
566 Ferreira, 2006; van Puijenbroek et al., 2017). Whether exposed dunes have a higher net dune growth
567 compared to dunes landward from the foredune depends mainly on the storm intensity and frequency. A
568 single high intensity storm can erode all the sand that exposed dunes have accumulated over a whole
569 summer, and in such case sheltered dunes could have a higher growth rate than the exposed dunes. The
570 exact relative growth rate over summer depends on the number of aeolian transport events. Linking the
571 number of aeolian transport event to the relative growth rate over summer would be a worthwhile
572 avenue for future research.

573 Sand supply and storm intensity are also affected by local conditions as beach morphology. A
574 minimum beach width is needed to reach maximum aeolian transport, the fetch length (Delgado-
575 Fernandez, 2010; Dong et al., 2004; Shao and Raupach, 1992). Our study site had a wide beach (0.9 km
576 wide), and we assume that the maximum aeolian transport was reached. The net growth of our foredune
577 was approximately 30 m³ per m foredune parallel to the sea for a period of 10 months. This growth rate
578 does also occur at other places along the Dutch coast, but is not very common (Keijsers et al., 2014).
579 Storm intensity is also influenced by beach morphology. The presence of intertidal bars and a wide
580 beach can reduce the storm intensity by wave attenuation (Anthony, 2013; Ruggiero et al., 2004).
581 Therefore, we can assume that the net dune growth we found in our study will depend on the beach

582 morphology. On smaller beaches we expect the net dune growth to be lower compared to wider
583 beaches, due to the lower sand supply by reduced fetch length and higher storm erosion of dune (van
584 Puijenbroek et al., 2017)

585 **4.4.2 Vegetation**

586 For coastal dune development vegetation is essential, however the species-composition of the
587 vegetation seems less important than we assumed: species did not seem to affect dune growth over the
588 summer, but did affect dune growth over winter.

589 We did find differences in nebkha dune morphology between the species, which suggest a causal
590 relationship. However, the difference in nebkha dune morphology between species is probably also
591 caused by differences in nebkha dune age. In Western Europe, the primary succession of coastal dunes
592 is generally assumed to start with *E. juncea*. Only after a fresh water lens has developed in the dune
593 with *E. juncea*, *A. arenaria* will establish (Westhoff et al., 1970). Over time *A. arenaria* will
594 outcompete *E. juncea*. This assumed succession pathway matches part of the spatial patterns that we
595 found in our study site and explains why nebkha dunes with only *E. juncea* are relatively small. Over
596 time these small nebkha dunes merge together after which *A. arenaria* is assumed to establish.
597 However, we found that *A. arenaria* has a large range in dune volume suggesting that, contrary to
598 current assumptions, *A. arenaria* can also establish on the bare beach without *E. juncea*, as long as the
599 soil salinity is not too high.

600 At our study site only two dune building species occur, however there are many different dune-
601 building species. It could very well be that other dune building species do have significant differences in

602 the nebkha dune growth over summer. For further research it would be interesting to study if these
603 results are similar in another nebkha dune system with different plant species.

604 **4.4.4 Application**

605 To our knowledge, we are the first to report on the relationship between initial dune volume and dune
606 growth for nebkha dunes in the field. The linear relationship that we found in our studies can be
607 incorporated in mathematical models that predict dune development. Furthermore, our research shows
608 that for predicting dune growth species identity does not matter during the summer, however it does
609 matter during the winter. This indicates that for dune building models, species identity is especially
610 important when winter survival of nebkha dunes is modelled. Furthermore, for the construction of an
611 artificial dune it appears to be crucial to plant the more storm resistant species.

612 Despite the presence of smaller nebkha dunes seaward of the foredune, the foredune showed a
613 large increase in volume compared to similar foredunes along the Dutch coast. This indicates that sand
614 supply to the foredune was not seriously hampered by the presence of the small vegetated dunes, while
615 the smaller dunes seaward of the foredune likely added to the protection of the foredune against storm
616 erosion. For coastal management it could be beneficial for foredune growth to have nebkha dunes
617 seaward of the foredune given a high sand supply.

618

619 **5. Conclusions**

620 The purpose of this study was to explore the contribution of vegetation and dune size on nebkha dune
621 development at locations differing in shelter from the sea. Our results show that 1) the contribution of
622 vegetation and dune size depend on season and degree of shelter. 2) Species composition does not affect
623 dune growth over summer, but does affect dune growth during winter, particularly at exposed sites. 3)
624 During early dune development, nebkha dune growth is linearly related to nebkha dune volume,
625 whereas dune volume does not seem to matter for nebkha dune erosion. 4) Sheltering by a foredune
626 reduces both sand supply and dune erosion; the net effect of shelter on dune growth therefore likely
627 depends on beach morphology and weather conditions. These results can be incorporated in models
628 predicting nebkha dune development and can be used by managers to determine coastal safety.

629

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635 **Competing interests** The authors declare that they have no conflict of interest.

636 **Data availability** ~~Data will be made available through the 4TU.Center for Research Data~~ Final dataset
637 used for the statistical tests and the data on the accuracy of the photogrammetric reconstruction are
638 archived in the 4TU Datacentre (<http://researchdata.4tu.nl/home/>), under
639 <https://doi.org/10.4121/uuid:8a2-30db-4328-bf04-40618bf31e4c>. The RAW images, digital surface

Gewijzigde veldcode

Met opmaak: Standaardlinea-lettertype, Lettertype:
(Standaard) Verdana, 8.5 pt, Engels (Verenigde Staten)

Met opmaak: Lettertype: (Standaard) Times New Roman,
12 pt, Tekstkleur: Auto, Engels (Verenigd Koninkrijk),
Patroon: Doorzichtig

640 [model, digital terrain model and the orthomosaic are available upon request by the corresponding](#)
641 [author.](#)

642 **Author contributions** MvP, CN and JS performed UAV flights and image calibration. MvP, CN and
643 JL analysed the data. MvP, CN, JS, AdG, MR, FB and JL provided guidance on the scope and design of
644 the project, and contributed to the writing of the manuscript.

645 **Supporting information**

646 Additional supporting information can be found in the online version of this article:

647 **Supplement S1** Weather conditions in our study site for 2013 - 2016

648 **Supplement S2** DTM, DSM and orthomosaic of each mapping campaign

649 **Supplement S3** Nebkha dune morphology of selected dunes

650 **Supplement S4** Accuracy photogrammetric reconstruction

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797

798 **Table 1.** Statistical models for the relative change in dune volume between April – August (summer)
799 and November – April (winter) for nebkha dunes. In this model we tested the effect of species, dune
800 size, and degree of sheltering. The data was analysed with a general linear mixed model with blocks as
801 random intercept. The standardized estimates and level of significance are shown for the models. Model
802 selection was performed with AIC (Akaike information criterion) as selection criteria. Marginal R² is
803 the variation explained by the fixed factors, whereas the conditional R² is the variation explained by the
804 fixed and random factors.

<i>Model with species</i>	<i>Dependent variable:</i>			
	Relative change in dune volume			
	<i>Summer</i>		<i>Winter</i>	
	Full model	Model selection	Full model	Model selection
Main effects				
Intercept	1.18** *	1.17***	0.92** *	0.94***
E. juncea	-0.02		0.005	-0.02**
Mix	0.02		0.02	-0.003
Dune volume	6.10	8.27***	-6.0*	-3.43**
Clustering	-0.22	-0.18	0.22	0.23
Max. dune height	-0.25	-0.31*	0.15	0.087
Sheltering by foredunes	0.29*	0.31**	-0.31**	-0.31**
Interaction effects				
E. juncea * Dune volume	0.90		1.90	
Mix * Dune volume	-0.11		1.41	
E. juncea * clustering	0.11		0.04	
Mix * clustering	0.01		-0.006	
E. juncea * max. dune height	-0.08		-0.09	
Mix * max. dune height	-0.02		-0.033	
E. juncea * Shel. by foredune	-0.05		0.03	
Mix * Shel. by foredune	-0.02		0.001	
Dune volume * clustering	-4.64*	-5.65**	4.44**	4.10**
Dune volume * max. dune height	-1.16	-2.01*	0.62	
Dune volume * Shel. by foredune	1.85	2.00*	-1.11	-1.31*
Clustering * max. dune height	0.31	0.34*	-0.29	-0.27*
Clustering * Shel. by foredune	-0.12	-0.17*	0.12	0.13
Max. dune height * Shel. by foredune	-0.20*	-0.18*	0.19**	0.19**
Marginal R ²	0.31	0.31	0.25	0.23
Conditional R ²	0.34	0.33	0.39	0.39
Observations	236	236	236	236

Akaike Inf. Crit.	-632.60	-685.45	-673.10	-709.11
Bayesian Inf. Crit.	-555.08	-641.04	-595.57	-661.35

Note:

*p<0.05; **p<0.01; ***p<0.001

805

806

807 **Table 2.** Statistical models for the relative change in dune volume between April – August (summer)
 808 and November – April (winter) for nebkha dunes. In this model we tested the effect of vegetation
 809 characteristics, dune size and degree of sheltering. The data was analysed with a general linear mixed
 810 model with blocks as random intercept. The standardized estimates and significance values are shown
 811 for the models. Model selection was performed with AIC as selection criteria. Marginal R² is the
 812 variation explained by the fixed factors, whereas the conditional R² is the variation explained by the
 813 fixed and random factors.

<i>Model with vegetation characteristics</i>	<i>Dependent variable:</i>			
	Relative change in dune volume			
	<i>Summer</i>		<i>Winter</i>	
	Full model	Model selection	Full model	Model selection
Main effects				
Intercept	1.24***	1.24***	0.90***	0.81***
Vegetation density	-0.003		-0.05	-0.03
Max. plant height	0.15	0.14**	0.04	
Dune volume	8.65***	6.62***	-2.72	-3.67**
Clustering	-0.21	-0.23	0.29	0.40**
Max. dune height	-0.44*	-0.41**	0.07	0.17
Sheltering by foredune	0.26*	0.29*	-0.28*	-0.25**
Veg. density * max. plant height	-0.01		0.001	
Veg. density * dune volume	0.83		0.92	
Veg. density * clustering	-0.03		0.078	0.06
Veg. density * max. dune height	0.04		-0.03	
Veg. density * Shel. by foredune	-0.005		-0.03	-0.04**
Max. plant height * dune volume	-0.58		-0.19	
Max. plant height * Clustering	0.02		-0.06	
Max. plant height * max. dune height	-0.11	-0.10**	0.04	
Max. plant height * Shel. by foredune	0.004		-0.01	
Dune volume * clustering	-6.37**	-6.30***	4.51**	4.65***
Dune volume * max. dune height	-1.54		-1.11	
Dune volume * Shel. by foredune	1.63	1.95*	-2.23*	-1.82**
Clustering * max. dune height	0.40*	0.41**	-0.32	-0.42**

Clustering * Shel. by foredune	-0.15	-0.17*	0.05	
Max. dune height * Shel. by foredune	-0.16	-0.16*	0.28**	0.31***
Marginal R ²	0.33	0.31	0.24	0.21
Conditional R ²	0.37	0.35	0.42	0.40
Observations	236	236	236	236
Akaike Inf. Crit.	-622.85	-674.05	-656.46	-704.97
Bayesian Inf. Crit.	-542.07	-626.28	-575.68	-657.20

Note:

*p<0.05; **p<0.01; ***p<0.001

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816 **Figure captions**

817 **Fig. 1** A) Overview of the Hors on Texel, the Netherlands. The white lines show the flight path for the
818 four different flights. The points show the position of the ground control markers. The white polygon is
819 the monitoring area, which is 200 m x 400 m. B) Photograph of the study site with the UAV used to
820 monitor the nebkha dunes.

821 **Fig. 2** Workflow of the methodology. The 3D point cloud from the photogrammetry was used to
822 construct a DSM, DTM and NDVI orthomosaic. The DTM and NDVI orthomosaic were used to
823 define the nebkha dunes. The explanatory variables for the statistical models were derived from the
824 DSM, DTM and NDVI orthomosaic. For a more detailed explanation see methods.

825 **Fig. 3** Overview of the monitoring area. A) The elevation is shown with the Digital Terrain Model (m
826 NAP), the green pixel indicates grass cover and the polygons indicate the nebkha dunes. B) The colour
827 indicates the species present on the nebkha dune and the squares the blocks. The foredune in the middle
828 of the monitoring area is excluded from the statistical analysis. Some dunes were cut-off by the edge of
829 the DTM, we discarded these dunes.

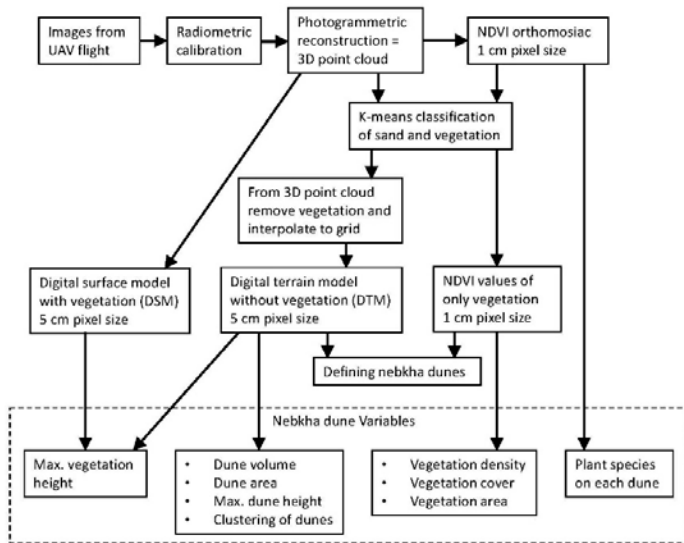
830 **Fig. 4** Different dune characteristics for nebkha dunes in August with *A. arenaria*, *E. juncea* and a mix
831 of both species separated for dunes seaward and landward of the foredune: A) Dune area, B) Maximum
832 dune height, C) Dune volume, D) Clustering: mean height around a 25m radius around the dune, E)
833 Vegetation density, F) Plant height. The letters denote the significant difference between the bars.
834 Seaward of the foredune there were 41 dunes with *A. arenaria*, 193 dunes with *E. juncea*, and 53 dunes
835 with both species, landward of the foredune there were 81 dunes with *A. arenaria*, 23 dunes with *E.*
836 *juncea*, and 41 dunes with both species. NAP refers to Amsterdam Ordnance Date, which refers to mean
837 sea level near Amsterdam

838 **Fig. 5** The relationship between initial dune volume (m^3) and the absolute change in dune volume (m^3 /
839 week) for: A) summer (April – August); B) winter (November – April). The data is shown for nebkha
840 dunes seaward and landward of the foredune. The black line shows the regression prediction, the grey
841 dashed line the 95% confidence interval. The formulas are the result of a linear regression model.

842 **Fig. 6** Relative change in dune volume (m^3/m^3)/week for nebkha dunes with *A. arenaria*, *E. juncea* and
843 a mix of both species and separated for dunes seaward and landward of the foredune for: A) summer,
844 April – August; B) winter, November – April. The letters denote the significant difference between the
845 bars. Seaward of the foredune there were 28 dunes with *A. arenaria*, 77 dunes with *E. juncea*, and 28
846 dunes with both species, landward from the foredune there were 57 dunes with *A. arenaria*, 22 dunes
847 with *E. juncea*, and 25 dunes with both species.

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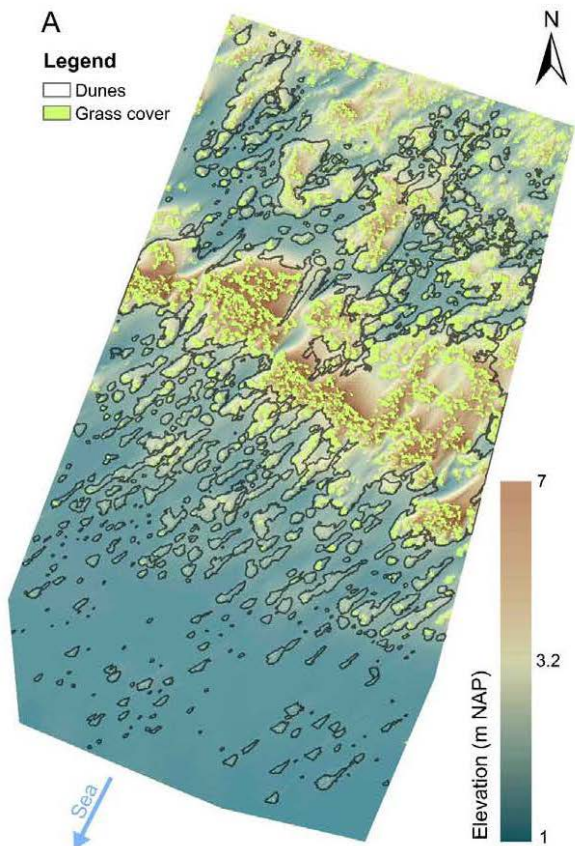




A

Legend

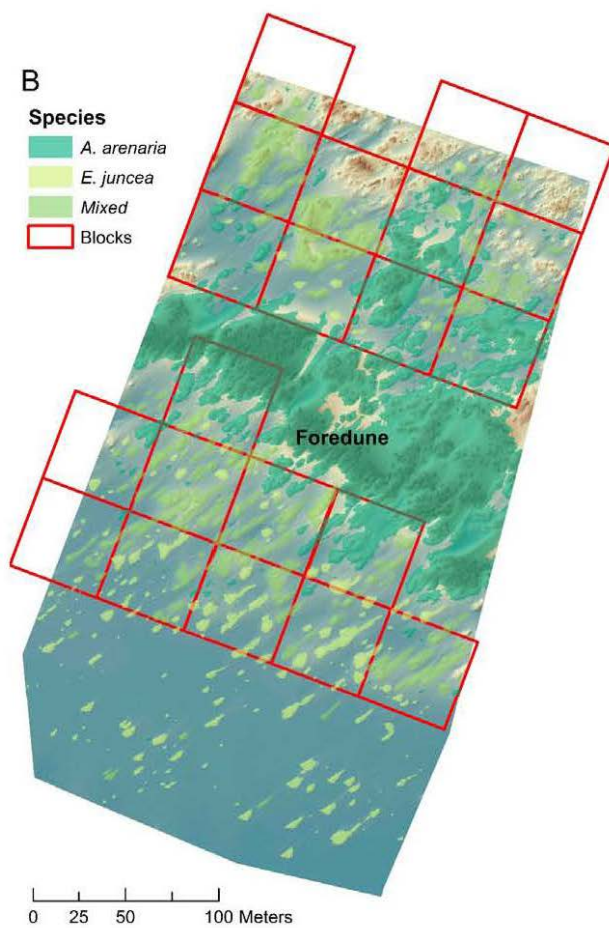
- Dunes
- Grass cover



B

Species

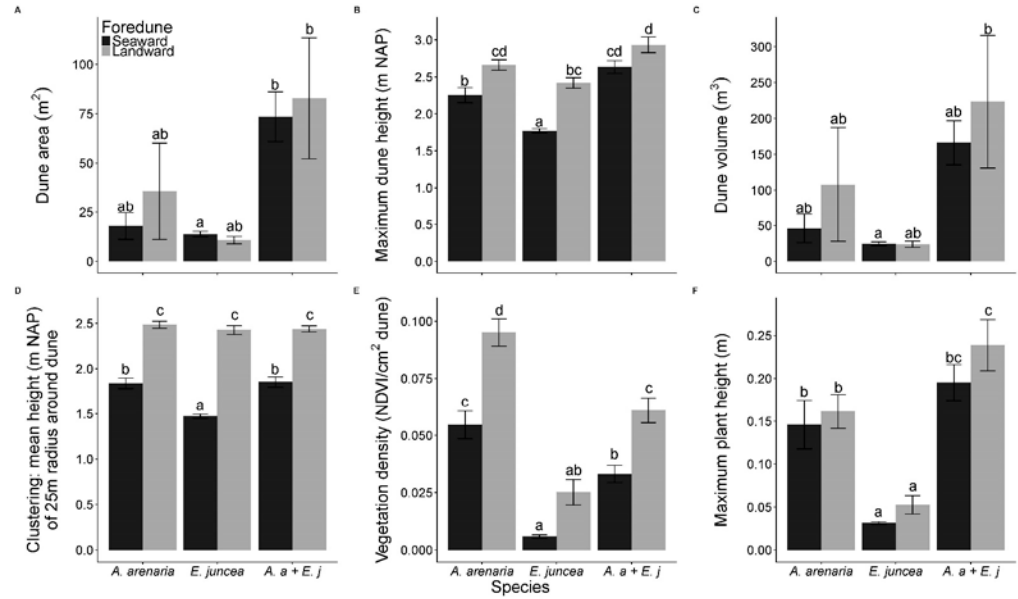
- *A. arenaria*
- *E. juncea*
- Mixed
- Blocks



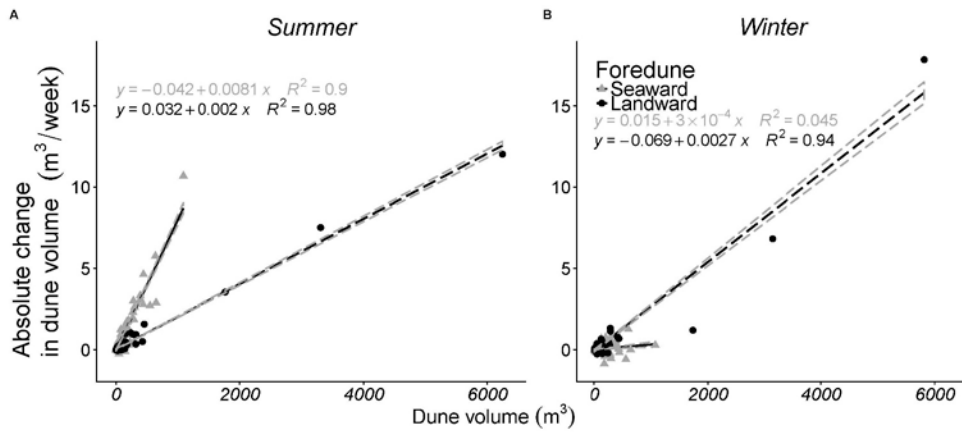
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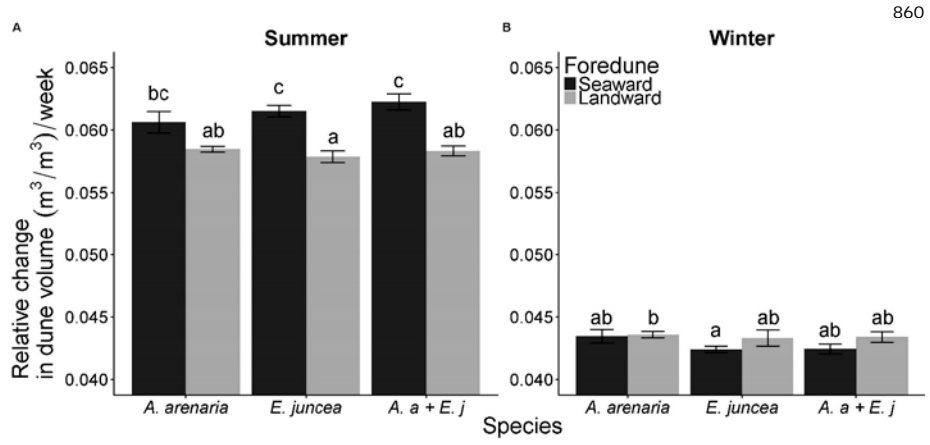


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