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1 **Erosion-initiated stromatolite and thrombolite formation in a present-day**  
2 **coastal sabkha setting**

3

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14 **ABSTRACT**

15 Laminated microbial mats and microbialites have been documented from a variety of coastal  
16 marine environments. This study aims to provide the first detailed descriptions of intertidal  
17 pools, along with their hosted thrombolite and stromatolite structures, from Abu Dhabi, and  
18 to propose a model for their formation and evolution. We propose that the development of  
19 pools within the upper intertidal zone was initiated by localised erosion of the laminated  
20 microbial mats during high energy events. The removal of the protective mats permitted  
21 erosion of the underlying unconsolidated sediment to produce erosional scours that continued  
22 to develop to produce the pools observed today. The margins of the newly-created submerged  
23 environment were colonised by a cyanobacteria dominated microbial community. The  
24 precipitation of aragonite cement, associated with the cyanobacteria, stabilised the pool walls  
25 and cemented the microbial communities to form stromatolitic and thrombolitic fabrics.  
26 Syndepositional cementation was further enhanced by the precipitation of marine cements as  
27 a result of evaporation-driven Ca<sup>2+</sup> and Mg<sup>2+</sup> supersaturation. Erosion behind and below the  
28 cemented pool wall eventually resulted in rim-collapse and the formation of the observed  
29 pool margin parallel thrombolite bands. Successive generations of lithification and erosion  
30 increased the area of the pool with the earliest thrombolites eroding and becoming  
31 increasingly isolated. In summary, the resultant microbialites developed through the complex  
32 interplay of erosion, abiotic early lithification and microbially-mediated processes and  
33 represent a continuum between unlithified laminated microbial mats and domal microbialites.  
34 These features are most likely produced during a sea level scenario of stillstand or  
35 transgression, and, as such, may be useful as a diagnostic tool to elucidate the onset of  
36 transgression. The newly proposed model for stromatolite formation has significant  
37 implications for the recognition and interpretation of similar structures observed in the fossil  
38 record.

39



40 **KEYWORDS**

41 sabkha, erosion, stromatolite, thrombolite, microbialite, lithification

42

## 43 INTRODUCTION

44 Stromatolites are laminated benthic microbial deposits (Riding, 1999) formed through  
45 lithification processes mediated or controlled by microbial communities of archaea, bacteria  
46 and/or diatoms (Dupraz et al., 2011). Stromatolites appear throughout much of the  
47 stratigraphic record, occurring from the Early Archean (Allwood et al., 2006) to the Recent  
48 with different organisms taking on the role of bioconstructors at different times. Extant  
49 stromatolites have been documented from a variety of coastal marine settings, the best well-  
50 known being the structures observed at Hamlin Pool, Shark Bay, Western Australia (Golubic,  
51 1985; Jahnert and Collins, 2013; Logan, 1961; Reid et al., 2003) and from the Bahamas  
52 (Andres and Reid, 2006; Bowlin et al., 2012; Monty, 1972; Stolz et al., 2009). Microbial  
53 mats, thrombolites and stromatolites have also been described from numerous other coastal  
54 settings, including Lagoa Vermelha, Brazil (Vasconcelos et al., 2006), Cayo Coco lagoon,  
55 Cuba (Bouton et al., 2016), the Caicos Platform (Trembath-Reichert et al., 2016) and the  
56 coastal sabkhas of the Persian Gulf (Brauchli et al., 2016; Court et al., 2017; Evans, 1966;  
57 Kendall and Skipwith, 1968; Lokier et al., 2017).

58 Despite the long stratigraphic history and significant geographical range of these  
59 biologically-generated sedimentary structures, models for their mode of formation have been  
60 surprisingly limited to studies of Recent examples from only a limited number of depositional  
61 settings. Such a paucity of studies is even more surprising when one considers that in many  
62 'stressed' environments the microbial communities associated with stromatolite and  
63 thrombolite formation were the dominant and, often, only biota interacting with sediments at  
64 the time of deposition. Succinctly, while stromatolites and thrombolites have a significant  
65 potential as a tool for paleoenvironmental interpretation, such utility requires that their mode  
66 of formation is considered from a range of depositional settings and potential distinguishing  
67 characteristics are identified.

68 The stromatolite communities living at the southern shoreline of the Persian/Arabian Gulf  
69 (hereafter referred to as the Gulf) provide an ideal site to study stromatolite formation in a  
70 previously-undescribed coastal environment. The presence of thrombolitic, hemispheroid  
71 microbial mats and stromatolite-like features from the coastal sabkha of Abu Dhabi has been  
72 noted (Kendall and Skipwith, 1968), yet, to date, no detailed description and interpretation of  
73 these features has been made. This study aims to provide the first detailed descriptions of the  
74 thrombolite and stromatolite structures at the seaward margin of the coastal sabkha complex  
75 of Abu Dhabi. The study does not aim to provide a detailed analysis of the composition of the  
76 microbial communities forming the stromatolites. Rather, the focus of the study is to describe  
77 and explain the morphology of the resultant stromatolite structures within the context of the  
78 environmental factors observed in the depositional setting and to consider their mode-of  
79 formation in the context of how these observations may be employed to constrain  
80 paleoenvironmental interpretations of similar features from the depositional record.

81

## 82 STUDY AREA

83 The study area is located at the southern shore of the Gulf, approximately 40 km to the  
84 southwest of Abu Dhabi City (Fig. 1A, B). The Gulf is a shallow microtidal sea with an  
85 average water depth of 35 m and a tidal range of 1.5 m in open-marine areas and less than 1  
86 m in lagoons. At the Abu Dhabi coastline, temperatures range between 7 °C at night during

87 the winter and exceed 50 °C during summer months (Lokier, 2012). Average annual rainfall  
88 is 72 mm, predominantly occurring as brief, heavy, downpours during winter months (Raafat,  
89 2007). Arid conditions result in evaporation (2.75 m per year) far exceeding precipitation  
90 (Bottomley, 1996).

91 The southeast region of the Gulf comprises a low-angle, north-sloping, ramp geometry that  
92 was rapidly flooded as post-glacial eustatic sea level rise transgressed an aeolian dune system  
93 resulting in a transition from aeolian-siliciclastic to marine-carbonate dominated deposition  
94 (Evans et al., 1969; Lambeck, 1996; Lokier and Steuber, 2008). A short-lived still stand,  
95 between 7,100 - 6,890 cal yrs BP (Lokier et al., 2015), facilitated the development of coastal  
96 sabkhas prior to renewed transgression with the onset of the Middle Holocene Atlantic stage.  
97 The subsequent relative sea level fall and stillstand resulted in the development of a strongly  
98 progradational geometry throughout the Middle to Late Holocene (Lokier and Steuber, 2008;  
99 Lokier et al., 2015).

100 From seaward to landward, the surface facies of the modern sabkha (Fig. 1C) are  
101 characterised by peloidal carbonate sands in the lower and middle intertidal zone (lagoon), a  
102 coast-parallel microbial mat belt in the upper intertidal zone, and evaporite precipitates in the  
103 supratidal zone (Evans et al., 1969; Kendall and Skipwith, 1969). The spatial occurrence of  
104 these facies belts is controlled by the local angle of slope which governs the duration of tidal  
105 flooding (Court et al., 2017). The microbial mat belt varies in width between 150 to 800 m  
106 and is classified (Court et al., 2017) based on surface morphologies resulting from the  
107 duration of inundation.

108

## 109 **METHODOLOGY**

110 Google Earth satellite imagery was used to identify areas of the intertidal zone where  
111 channels and pools were present. A field reconnaissance campaign, in January 2016,  
112 identified an area containing an ephemeral ponds network, tidal channels and the intertidal  
113 pool that is the focus of this study. The morphology of the pool was mapped using a Garmin  
114 GPSmap 78 device; the data was subsequently imported and processed in Quantum GIS 2.18.  
115 Microbial, including stromatolite and, thrombolites features, and other sedimentary and  
116 biological characteristics of the pool were documented in detail. Monitoring and sampling  
117 visits were regularly made between January 2016 and May 2017 with any changes to the  
118 morphology of the pool or the associated features being documented in detail. Three  
119 specimens of microbialite from the pool margin and one specimen of thrombolite from the  
120 pool centre were recovered and subsequently stored in seawater in plastic containers.

121 A weather station (WS-GP1, Delta-T Devices), 3 km landwards from the study site (Fig. 1C)  
122 recorded air temperature, air humidity, wind direction, wind speed, and precipitation at a  
123 height of 1.2 m between April 2016 and July 2017. Battery failure resulted in data loss  
124 between February and March 2017. Barometric pressure and temperature data were recorded  
125 at 30 minute intervals between February 2016 and June 2017 using a Baro-Diver barometric  
126 logger (Van Essen Instruments) sited approximately 2 km to the east of the weather station  
127 (Fig. 1C). Data analysis and visualisation were conducted in the statistical computing  
128 language R, in the programming language Python version 2.7.10 and in PAST version 3.15.

129 The physico-chemical characteristics of the water in the studied pool were measured using an  
130 Ultrameter II 6PFC device (Myron L Company) to record temperature, conductivity, total  
131 dissolved solids (TDS), oxidation-reduction potential (ORP), resistivity and pH. Very high  
132 dissolved solid load resulted in resistivity measurements that exceeded instrument parameters  
133 and, thus, were not used for interpretations. Salinity was measured using a Brix-type  
134 refractometer with automatic temperature compensation. The water level in the studied pond  
135 was monitored between February to October 2016 using a CTD-diver water level logger (Van  
136 Essen Instruments) installed 5 cm above the pool base. Water level was recorded as pressure  
137 in mbar, with a barometric conversion applied by subtracting barometric pressure from the  
138 water level pressure, under the assumption that 1 mbar = 1 cm of H<sub>2</sub>O.

139 The microbialite structures were investigated at the micro-scale using a Quanta 200 (FEI)  
140 scanning electron microscope (SEM), located at The Petroleum Institute of Khalifa  
141 University of Science & Technology. One sample was selected from which representative  
142 sections were broken off, cleaned with compressed air, mounted on metal stubs, and coated  
143 with a palladium/gold mixture. The energy-dispersive X-ray spectroscopy mode (EDX) of  
144 the SEM was used to semi-quantitatively characterise element compositions of specific  
145 features. Samples were also prepared for analysis under the low-vacuum conditions of the  
146 environmental SEM mode. These samples rapidly dehydrated and became charged, it was  
147 therefore decided to focus on careful sample preparation and application of the conventional  
148 SEM mode.

149 Petrographic thin sections were prepared from three microbialite specimens, including the  
150 specimen that was investigated in the SEM. The samples were cut using a diamond rock saw  
151 and subsequently left to dry in the laboratory for 24 hours. Each sample was then  
152 impregnated with blue resin to enhance the visibility of pore space under the optical  
153 microscope.

154 DNA was extracted from samples of polygonal microbial mat, using a FastDNA SPIN kit for  
155 soil (Q-BIOgene, Cambridge, UK), according to the manufacturer's instructions. DNA  
156 sequencing was performed on a MiSeq Personal Sequencer (Illumina) at the NUomics  
157 Facility (Northumbria University, Newcastle, UK). Gene amplicon libraries (16S rRNA)  
158 were prepared, including PCR amplification, purification of amplicons and quantification  
159 procedures, according to the procedures of Kozich et al., 2013. Aliquots of purified  
160 amplicons were sequenced to produce 2 x 250 bp paired-end reads with v2 chemistry. Raw  
161 sequencing reads were analysed using QIIME2 (Bolyen et al., 2019), with Amplicon  
162 Sequence Variants (ASVs) determined using Divisive Amplicon Denoising Algorithm  
163 (DADA2, Callahan et al., 2016). Taxonomic classification of representative ASV sequences  
164 was performed using the Naïve Bayesian classifier (Bokulich et al., 2018). The relative  
165 abundance of ASVs was calculated by taking the sum of the reads for individual ASV  
166 abundances and dividing by the total number of reads for all ASVs within a sample and  
167 multiplying by 100.

168

## 169 **RESULTS**

### 170 **Environmental data**

171 Air temperatures recorded by the weather station ranged between 8.4 °C and 48.9 °C; air  
172 temperatures recorded by the barometric logger ranged between 8.9 °C and 53.7 °C (Fig.  
173 2A). The highest temperatures were recorded during July and August, while the lowest  
174 temperatures were recorded during February. Total precipitation over the measurement  
175 period amounted to 7.6 mm, mainly resulting from torrential rainfall events in January and  
176 February 2017 (Fig. 2A). March 2017 experienced heavy rainfall, but, as previously  
177 mentioned, this data was not recorded by the weather station due to a battery failure.  
178 Therefore, the rainfall data represent a minimum value for the measurement interval. The  
179 primary wind direction throughout the recording period was from the north-west, with  
180 subordinate winds from the south (Fig. 2C). Wind speed varied between 0.3 - 15.4 m/s with a  
181 mean of 3.7 m/s (Fig. 2C, D).

182 Water temperatures in the pool ranged between 11.7 °C and 46.8 °C during the measurement  
183 period (Fig. 2B). The lowest water temperatures were recorded in February while the highest  
184 water temperatures were recorded in August. Salinity ranged between 75.0 ‰ to 93.0 ‰,  
185 accompanied by pH values between 7.3 to 8.1. Of the remaining physico-chemical  
186 parameters, conductivity ranged between 100.0 - 118.7 mS/cm, TDS ranged between 75.0 -  
187 91.26 parts per thousand (ppt), and ORP values ranged from 79 mV to 100 mV. The tidal  
188 regime in the studied pool was semi-diurnal micro-tidal, with a water depth ranging from a  
189 minimum of 19 cm when the pool was emergent up to a maximum of 109 cm during an  
190 unusually high tide (Fig. 2B).

191

## 192 **Pool morphology and hydrological regime**

193 The studied intertidal pool is located at the seaward margin of the polygonal microbial mat  
194 zone (Fig. 1C). The pool is broadly U-shaped and hydrologically open at its seaward and  
195 landward sides (Fig. 3A, B). The seaward opening of the pool corresponds to the landward  
196 termination of a tidal channel that extends into the lower intertidal and subtidal zones. The  
197 landward opening to the pool is defined by a narrow, well-lithified, sill that provides a  
198 connection to an elevated landward ephemeral pond system (Fig. 4). Ebb tide water continues  
199 to flow into the pool over this sill even whilst the next flood tide is recharging the pool via  
200 the tidal channel. The pool exhibits a perimeter of approximately 80 m and an area of  
201 approximately 250 m<sup>2</sup>. The height difference between the cemented floor of the pool and its  
202 upper rim ranges between 20 to 30 cm. The floor of the pool is always submerged, this is in  
203 stark contrast to the elevated surrounding areas of microbial mat that experience daily cycles  
204 of exposure and inundation.

205

## 206 **Macro-scale sedimentary and biological features**

207 The general stratigraphy in the area of the pool (Fig. 3C) consists of Pleistocene to Early  
208 Holocene aeolian mixed siliciclastic-carbonate sands, overlain by a shallow-marine  
209 transgressive carbonate hardground of mid-Holocene age (Ge et al.; Lokier and Steuber,  
210 2009; Paul and Lokier, 2017), followed by an overlying unconsolidated, to lightly-cemented,  
211 bioclastic rudstone. This sequence is terminated by laminated microbial mats with a

212 polygonal surface architecture. At the pool-floor, the hardground is locally exposed or is  
213 covered by an organic ooze that increases in thickness (up to 7 cm) landward (Figs 3 and 5).  
214 Accumulations of spongy, unlithified, mobile, gravel-sized grains are observed throughout  
215 the pool. These grains vary in colour and typically accumulate in small troughs and in the lee  
216 of thrombolite patches and bands. The grains vary in diameter between 5-10 mm and are  
217 irregularly shaped (Fig. 6A). The grains contain sub-mm scale inclusions of bioclastic grains  
218 including benthic foraminifera (Fig. 6B-D).

219 Clotted microbial (thrombolite) fabrics are distributed within the pool as isolated domes or  
220 discontinuous dm to m-long bands that are typically oriented parallel to the pool margin (Fig.  
221 7A). The thrombolite domes typically measure between 5-20 cm in width, and exhibit relief  
222 above the hardground of between 10-25 cm. The structures are coloured brown to dark  
223 brown, and are characterised by a spongy consistency that reflects a dominantly  
224 cyanobacteria composition. Cyanobacteria were detected throughout polygonal mats in the  
225 area ( $\Sigma$  relative abundance;  $8.5\% \pm 2.1$ ), with different morphologies including coccoid  
226 (*Halothece*, *Gleocapsa*) and filamentous (*Coleofasciculus*, *Lyngbya*, *Phormidium*) forms. A  
227 mm-thick cover of bioclastic grains is locally present at the upper surface of the thrombolites  
228 (Fig. 7B). Marine brown and green algae grow at the outer margins of individual  
229 thrombolites, defining the minimum water depth within the pool. Internally, the domal  
230 structures comprise two distinct intervals (Fig. 3C). The upper interval is a thrombolite,  
231 exhibiting crude laminations corresponding to different colours of microbial communities  
232 interlayered with bioclasts (Fig. 8A). The lower layer is a ~10 cm thick laminated  
233 stromatolite that is attached to the underlying hardground (Fig. 8B). The margins of the pool  
234 exhibit the same vertical facies distribution as the thrombolite domes and bands but are  
235 locally undercut by up to 10 cm (Figs 3 and 9).

236 Moving away from the pool, into the adjacent microbial mats, the lithified cyanobacterial  
237 communities of the pool margin transition over decimetres into an unlithified finely-  
238 laminated and polygonal microbial mat dominated by filamentous cyanobacteria (Fig. 10).  
239 Elongate scours in excess of 1 m long are distributed throughout the polygonal microbial mat  
240 zone (Fig. 11). The scours are oriented perpendicular to the shoreline, are shallowest at their  
241 seaward end and gradually deepen landward. The scours host a variety of microbial  
242 communities that closely resemble the organic ooze observed in the pool.

243

## 244 **Micro-facies and cement fabrics**

245 The pool margin microbialites contain abundant lithoclasts, bioclasts, peloids and coated  
246 grains (Fig. 12A) surrounded by pervasive microbial extracellular polymeric substance (EPS)  
247 (Figs 13A, B). The skeletal assemblage observed within the microbialites is dominated by  
248 benthic foraminifera (primarily peneropolids) and ostracods with subordinate bivalves.  
249 Skeletal grains are typically whole with little evidence of abrasion or breakage (Fig. 12A).  
250 SEM observations reveal the presence of cyanobacteria tubes and filaments within the EPS  
251 (Fig. 13 B, E). Primary microbial laminae are difficult to discern in thin section, occurring as  
252 discontinuous degraded organic seams that could be easily confused with dissolution seams  
253 (Fig. 12E, F).

254 Grains are coated by an isopachous fringing acicular (Sandberg et al., 1985) aragonite cement  
255 with fibrous needles oriented perpendicular to grain surfaces (Figs 12B-D). The aragonite

256 needles exhibit pointed crystal terminations, have a length of 16-185  $\mu\text{m}$  with a width of 3-5  
257  $\mu\text{m}$  (axial ratio typically  $>22$ ). Rare dolomite cements occur as isolated rhombs between 3-6  
258  $\mu\text{m}$  and more-massive clusters of slightly coarser (up to 8  $\mu\text{m}$ ) equant euhedral crystals (Fig.  
259 13C, D). These dolomite crystals are enclosed in microbial EPS and acicular aragonite. Semi-  
260 quantitative EDX elemental analysis of the dolomite crystals yield an Mg content of between  
261 approximately 20-40% compared to significantly lower values ( $\sim 14\%$ ) in the surrounding  
262 matrix (Fig. 13 C, D).

263 Primary interparticle porosity has been largely occluded by the pervasive precipitation of the  
264 aragonite cement phase that has cemented grains and significantly restricted pore throats (Fig.  
265 12C, D). Secondary, yet syndepositional, mouldic porosity after bioclasts (Fig. 12B), has also  
266 been partially to completely occluded by the precipitation of the aragonite cement phase.

267

## 268 **INTERPRETATION AND DISCUSSION**

269 On the basis of field and laboratory observations, it is here proposed that the formation of the  
270 intertidal pools and the subsequent genesis of the hosted stromatolite and thrombolite  
271 microbial fabrics are genetically related.

272

### 273 **Formation of intertidal pools and associated microbialites**

274 As has been previously documented, the development of significant modern-day microbial  
275 mat communities requires stressed environmental conditions that prevent competition from  
276 macroalgae and other eukaryotic organisms (Bouton et al., 2016; Bowlin et al., 2012;  
277 Suosaari et al., 2016b). In the case of the Abu Dhabi sabkha, these conditions are met in the  
278 uppermost intertidal zone where diurnal flooding is sufficient to maintain microbial growth,  
279 yet exposure inhibits competition. Under such conditions, laminated, unlithified, microbial  
280 communities are able to thrive.

281 Pool formation is initiated where a high energy event, such as a storm surge or high tide,  
282 removes the protective binding microbial mat and exposes the underlying unconsolidated  
283 sediment (Fig. 11A). Under the prevailing relatively low-energy conditions of the Abu Dhabi  
284 sabkha, such events are relatively rare and of limited duration, thus normally allowing  
285 recolonisation of the sediment surface by a new generation of microbial mat and the effective  
286 'healing' of the protective binding layer. However, prolonged episodes of high energy, or  
287 multiple high energy events over a short time interval, will erode the unprotected sediment to  
288 produce an erosional scour that may extend to the depth of the underlying lithified  
289 hardground (Figs 11B, 14B). Initially, the unprotected vertical walls of the pond are  
290 susceptible to continued erosion, thereby facilitating lateral growth of the pool.

291 With a return to quiescent conditions, the newly formed pool walls (Fig. 11B) provide a  
292 submerged substrate for the development of cyanobacteria dominated microbial communities  
293 that overgrow the exposed truncated surface of the laminated microbial mats (Figs 9, 10) and  
294 promote aragonite precipitation (Figs 12, 13). Lithification stabilises both the underlying  
295 laminated microbial mat substrate, producing stromatolite fabrics, and the clotted textures to

296 form a thrombolitic fabric (Fig. 14C). Increasing lithification stabilises the pool walls and  
297 inhibits erosion during subsequent high energy events.

298 Microtidal flooding cycles ensure regular marine inundation of both the pool and surrounding  
299 microbial mats by seawater that is oversaturated with respect to  $\text{Ca}^{2+}$  and  $\text{Mg}^{2+}$  (Paul and  
300 Lokier, 2017; Shinn, 1969; Wood and Sanford, 2002). Ebb tide surface flow is concentrated  
301 into tidal channels that rapidly drain much of the upper intertidal zone. High evaporation  
302 rates and shallow sub-surface bacterial sulphate reduction (Lokier and Steuber, 2008) further  
303 increase the concentrations of major ions in ponded surface and shallow subsurface waters.  
304 Localised dissolution of aragonite allochems (Fig. 12B) occurs where the pH of the pore-  
305 water is reduced by the oxidisation of hydrogen sulphide (Jordan et al., 2015). These highly  
306 saline,  $\text{Ca}^{2+}$  and  $\text{Mg}^{2+}$  oversaturated pore-waters flow seawards in the shallow subsurface  
307 until they intercept the pool margin and discharge into the pool, further promoting the  
308 precipitation of aragonite cements (Fig. 14C).

309 Over time, the pool walls become increasingly lithified through the continued precipitation of  
310 aragonite cements. During high tides, waves breaking at the pool 'shore' will erode the  
311 unlithified microbial mats immediately behind the wave-resistant cemented pool margin  
312 (Figs 3, 14C). Simultaneously, currents will erode the unlithified basal rudstone to undercut  
313 the pool wall (Fig. 9) and further destabilise the edge of the pool. This process will eventually  
314 result in the detachment of a section of the cemented microbial thrombolites and stromatolites  
315 which will founder onto the exposed hardground at the pool floor to create detached sub-  
316 aqueous margin-parallel thrombolite bands (Figs 3, 7A, 14D). The submerged thrombolite  
317 bands will continue to grow and precipitate aragonite cements within their clotted fabric (Fig.  
318 13). High energy events, such as storm surges, will detach and remove sections of the  
319 thrombolite bands. Erosion will decrease over time as the precipitation of early marine  
320 aragonite (Lokier and Steuber, 2008; Paul and Lokier, 2017) cements the thrombolite to the  
321 underlying hardground and stabilises the structure (Fig. 14E).

322 The freshly-exposed pool margin is now subject to renewed microbial colonisation and  
323 associated aragonite precipitation and the process is repeated with the genesis of a new  
324 generation of margin-parallel thrombolite band (Fig. 14F). With repeated cycles, this process  
325 will result in significant lateral expansion of the pool, possibly resulting in coalescence with  
326 any adjacent pools. Over time, the earlier thrombolites will become increasingly isolated  
327 (Fig. 3) and may be subjected to undercutting to produce the distinctive dome-shaped  
328 morphology with clotted thrombolitic fabrics sitting atop laminated stromatolites (Figs 7B, 8,  
329 14F). These thrombolite domes superficially resemble the 'isolated patch' macrofabrics  
330 described from the Cayo Coco Lagoon, Cuba (Bouton et al., 2016). However, the Cayo Coco  
331 structures appear to be entirely thrombolitic in nature, lacking either a laminated stromatolite  
332 component or a relationship with an underlying hardground.

333 The accumulations of unlithified, spongy, microbial gravel-grade grains at the pool floor  
334 (Figs 6, 7A) are comprised of cyanobacteria and EPS forming a clotted microbial fabric akin  
335 to that of the thrombolites. These soft grains are inferred to have been sourced from the  
336 thrombolite domes or pool margin thrombolite bands either through erosion or via exfoliation  
337 due to high rates of microbial production. Once disarticulated, the low-density grains are  
338 particularly susceptible to reworking and concentration in the lee of thrombolites and in  
339 bathymetric lows. Where a significant amount of carbonate material is bound within these  
340 grains, it is possible that they may be preserved in, and recognised from, the geologic record  
341 as large aggregate grains with a clotted matrix (Gerdes et al., 1994). In more-open areas of



342 the pond, agitation of the microbial grains, by wave and current action, will result in abrasion  
343 and disarticulation to generate the microbial ooze that covers much of the pool floor.

344 The rare small, EPS-hosted, dolomite crystals observed in the pool-margin stromatolites are  
345 consistent with a microbially-mediated origin as proposed previously (Bontognali et al.,  
346 2010). Under this model, the fluctuating hypersaline conditions of the sabkha environment  
347 result in the EPS generating carboxyl functional groups that overcome the kinetic barrier to  
348 the low temperature incorporation of Mg into carbonate minerals (Bontognali et al., 2014;  
349 Petrash et al., 2017). These clusters of dolomite crystals bare a remarkably similarity, in  
350 terms of size, distribution and association with biofilms, to those previously documented  
351 from the Lower Messinian of Sicily (Oliveri et al., 2010).

352 The well-lithified nature of the sill separating the pool from the landward ephemeral ponds is  
353 inferred to result from a combination of microbially-mediated aragonite precipitation  
354 enhanced by degassing-related precipitation as the solubility product of the outflowing water  
355 is lowered as it cascades over the sill margin. This process is similar to the precipitation of  
356 carbonates at travertine terraces and tufas (Fig. 15; Gandin and Capezzuoli, 2014; Özkul et  
357 al., 2014).

358

### 359 **Microbialite formation in a sea level context**

360 The formation of the pools and their associated microbial communities is inferred to  
361 preferentially occur under prevailing conditions of relative sea level stillstand or, more likely,  
362 under the increased hydrodynamic conditions associated with relative sea level rise. The  
363 present-day rate of eustatic sea level rise ( $3.2 \text{ mm yr}^{-1}$ ) (Church et al., 2013) has been  
364 previously documented as resulting in significant shoreline retreat (up to  $29 \text{ m yr}^{-1}$ ) and  
365 associated hydrodynamic energy increase at the Abu Dhabi coastline (Lokier et al., 2018).  
366 Under such a transgressive (flooding) scenario it is expected that we shall observe an increase  
367 in the frequency and amount of erosion of the unlithified upper intertidal microbial mats and,  
368 consequently, the increased initiation and development of pools with their associated  
369 stromatolite and thrombolite fabrics. However, under conditions where the rate of relative sea  
370 level rise is 'too rapid', the associated increase in energy at the shoreline may result in  
371 dynamic coastline retreat at a rate that exceeds the ability of the microbial systems to re-  
372 establish and 'track' retrogradation.

373 From an environmental perspective, as the rate of present-day eustatic sea level rise continues  
374 to increase (Church et al., 2013), coastal microbial communities will be placed increasingly  
375 under threat and should thus be added to the growing list of coastal and shallow-marine  
376 ecosystems (Lovelock et al., 2015; Perry et al., 2018; Thorner et al., 2014) that are at risk due  
377 to anthropogenic-driven global warming and associated sea level rise.

378

### 379 **Microbialite preservation and identification in the fossil record**

380 Under a rising sea level scenario, as observed in the study area today, the preservation  
381 potential of the described structures is primarily dependent on the degree of early-lithification

382 and the subsequent rate of burial. As sea level rises, increased hydrodynamic energy easily  
383 erodes the uncemented microbial mats and thrombolites, transporting much of the material  
384 into the marine system where it is consumed. The likelihood of preservation of unlithified  
385 microbial features is further compromised as increased, and more regular, inundation reduces  
386 environmental stress in the intertidal and shallow subtidal zone. Such increased flooding  
387 opens this niche-environment to colonisation by macro-algae and other eukaryotes that will  
388 outcompete and displace the microbial communities. On entering the burial realm, ongoing  
389 degradation of the organic component of the unlithified microbial mats and thrombolites will  
390 further reduce the potential for preservation and identification within the stratigraphic record  
391 (Court et al., 2017; Kenig et al., 1990).

392 Conversely, cemented stromatolites and thrombolites have a significantly higher prospect of  
393 preserving primary depositional fabrics during inundation and subsequent burial and will,  
394 thus, have a higher preservation and recognition potential into the stratigraphic record. Where  
395 these early-lithified structures remain exposed at the sea floor, or are covered by only a thin  
396 veneer of sediment, they will be subject to early-marine cementation, thereby further  
397 enhancing the potential of preservation. If exposed at the sea floor, the lithified structures will  
398 act as a stable substrate for colonisation by sessile benthic communities, again enhancing the  
399 potential of preservation but providing a challenge for identification in the fossil record.

400 Where ponds and their associated structures have formed during a period of high sea level or  
401 falling stage stillstand, the chances of preservation into the fossil record are significantly  
402 reduced. During post-stillstand base level fall, unlithified microbial communities are subject  
403 to biodegradation and physical erosion. On entering the meteoric realm, lithified microbial  
404 structures will be subject to increased dissolution of aragonite cements and allochems. As  
405 erosion removes the surrounding unlithified sediments, the increasingly exposed cemented  
406 pond margins will lose their support, becoming unstable and subject to increased likelihood  
407 of erosion. This process may continue until all of the sediments are eroded to expose the  
408 underlying, strongly lithified, transgressive hardground, thereby potentially removing all  
409 evidence of the preceding transgressive cycle. Under these circumstances, transgressive  
410 sequences will only be preserved into the stratigraphic record where high frequency (4<sup>th</sup>-5<sup>th</sup>  
411 Order) transgressive cycles occur during a theme of overall, lower order, sea level rise. This  
412 scenario will result in repeated transgressive cycles punctuated by stillstands to produce  
413 parasequences as observed in the sabkha stratigraphy of the Jurassic Arab Formation  
414 (Grötsch et al., 2003).

415 Under the above-proposed scenarios, preservation of these structures into the geological  
416 record is only likely under a rising sea level scenario and where the microbial structures have  
417 been lithified. As such, the recognition of these systems from the stratigraphic record offers a  
418 potential diagnostic tool to elucidate the onset and, possibly, rate of marine transgression.

419 The differential preservation potential of lithified and unlithified components together with a  
420 limited thickness and laterally isolated morphology of these systems offers many challenges  
421 to their observation and interpretation in the geological record. Identification of the system  
422 requires recognition of lateral and vertical facies relationships and gross morphologies that is  
423 best suited to observation in laterally extensive outcrops. Conclusive identification of these  
424 systems from core would be significantly more challenging. A vertical stacking pattern of  
425 transgressive hardgrounds succeeded by stromatolites and overlying thrombolites, possibly  
426 with encrusting marine benthos, would hint at the presence of these systems. However, these  
427 vertical relationships could also be observed under other formational mechanisms, such as

428 those described from Storr's Lake (Dupraz et al., 2013) and Shark Bay (Jahnert and Collins,  
429 2011; Logan et al., 1974), and, without a significant density of cores, it is unlikely that the  
430 required knowledge of lateral facies geometries could be achieved from core alone.

431

## 432 CONCLUSIONS

433 Intertidal pools and associated microbial fabrics were observed and documented from the  
434 coastal sabkha of Abu Dhabi. We propose a new, erosion-based, model for the formation of  
435 these systems under a scenario of rising relative sea level. Increasing energy at the shoreline  
436 results in erosion of protective microbial mats to expose underlying sediments to erosion and  
437 initiate pool formation. These newly-formed submerged environments provided an ideal  
438 habitat for the pervasive growth of cyanobacteria that promoted the cementation of  
439 stromatolite and thrombolite textures thereby significantly increasing the potential for the  
440 preservation of these microbial structures. Successive erosional events enlarged the pools and  
441 created isolated microbialites with a range of textures.

442 Although the recognition of these features in the stratigraphic record may be challenging,  
443 their association with the onset of transgression makes them a potentially powerful diagnostic  
444 tool for constraining the initiation of relative sea level rise.

445

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460

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**648 Table and figure captions**

649 Table 1. Environmental data from a range of coastal marine stromatolite and microbial mat  
650 provinces. Sources for the data: Sabkha: this study and studies given in local settings.  
651 Hamelin Pool: Water temperatures, salinity, tidal regime and tidal range from (Suosaari et al.,  
652 2016a), air temperatures, mean annual rainfall and wind speed from <http://www.bom.gov.au>;  
653 evaporation from Logan (1961). Lagoa Vermelha: Höhn et al. (1986). Cayo Coco: Bouton et  
654 al. (2016). Highborne Cay: Bowlin et al. (2012).

655 Figure 1. A) Location of the study area in relation to the Gulf. B) Location of study area in  
656 the context of the Abu Dhabi coastline. C) Satellite image of the coastal sabkha of Abu  
657 Dhabi, showing facies belts and anthropogenic changes. The study site is indicated by a white  
658 star. Imagery source: ESA Sentinel 2A Multi-spectral Imager (MSI), date: 18 March 2017.

659 Figure 2. A) Mean daily air temperatures in the coastal sabkha measured between February  
660 2016 and June 2017 using a barometric logger (black line) and the GP1 weather station (red  
661 line). Daily maximum and daily minimum temperatures are shown in the background.  
662 Precipitation in January and February 2017 is indicated by the blue columns. B) Record of  
663 water temperatures and water depth in the studied intertidal pool, showing the daily tidal  
664 regime as well as long term annual trends in water depth. Note that the pool was never empty.  
665 C) Wind rose showing the dominant wind directions from the WNW - NW and S, and wind  
666 speeds. D) Frequency distribution plots of wind speeds for two intervals in January –  
667 February 2017 and in July – August 2016. Each vertical bar represents a 0.5 m/s wind speed  
668 interval.

669 Figure 3. A) Overview photograph of the studied pool. Note that the scale is not uniform  
670 from bottom to top due to the mode of image acquisition; person in background is 175 cm  
671 high. B) Interpretative overview of the features observed in the pool. Seaward direction is to  
672 the left. Note the detached bands at the pool margins, and the thrombolites towards the centre  
673 of the pool. C) Schematic cross-section through the pool showing the general stratigraphy,  
674 lithologies and features described in this study.

675 Figure 4. A) Close-up view of the lithified sill – see Fig. 3 for image location in the studied  
676 pool. The ephemeral pond system is located at the top of the image, while the pool is towards  
677 the bottom (not visible). The location of the water level logger is indicated by an arrow. Note  
678 the colonisation of the logger by cyanobacteria and algae. B) Schematic cross-section  
679 showing with the relationship between the ephemeral pond system, lithified sill and the pool.

680 Figure 5. A) Photograph of the organic ooze covering parts of the pool floor. Grazing trails of  
681 gastropods are visible with the respective gastropods at the end-points (arrows). B) Close-up  
682 view of the ooze and a gastropod (large arrow) and its grazing trail (small arrows).

683 Figure 6. A) Microbial ‘gravel’ collected from the pool floor. Different colours correspond to  
684 different organic pigments. B-D) Close-up views of individual microbial grains showing the  
685 EPS and white bioclastic grains embedded (arrow).

686 Figure 7. A) Close-up view of different microbial structures including thrombolites and  
687 stromatolites. Note the detached rim and the domal and banded thrombolites, surrounded by  
688 organic ooze and gravel-sized grains of microbial origin. Note also the aluminium can  
689 overgrown by microbial communities and partially covered by organic ooze. B) Close-up



690 view of an individual thrombolite showing its components: brown thrombolite, bioclasts and  
691 algae growth. Upper growth boundary of these algae indicates the minimum water depth in  
692 the pool, which corresponds to a depth of ~20 cm as also calculated from the water level  
693 logger data. See Fig. 3 for image location in the studied pool.

694 Figure 8. A) Cross-sectional view of an individual domal thrombolite, showing a clotted  
695 texture with internal crude radial laminations, based on differences in colour. Bioclasts  
696 enclosed by microbial EPS indicate outward growth. Coarse laminae are indicated by dashed  
697 lines. B) Hand specimen of a stromatolite from the pool margin. Microbial laminations  
698 preserved through lithification are clearly visible (between white arrows). Purple and green  
699 colours correspond to pigments of living cyanobacterial communities and are unrelated to the  
700 primary processes that lead to the development of the stromatolite.

701 Figure 9. Oblique subaqueous view of the partially undercut margin at the north-west side of  
702 the pool - see Fig. 3 for image location in the studied pool. Note the laminated stromatolites  
703 at the centre (white arrow).

704 Figure 10. Overview photograph showing the transition from the pool, via a rim overgrown  
705 with algae, into the surrounding polygonal microbial mat zone. Note that the polygonal  
706 microbial mat in the transition zone appears to be degraded. See Fig. 3 for image location in  
707 relation to the studied pool.

708 Figure 11. A) An area of recently eroded microbial mat exposing the underlying  
709 unconsolidated sediment. Note the incoming tide in the bottom left of the image (person is  
710 185 cm tall). B) An erosive scour in the microbial mat zone. C) Schematic cross-section  
711 through the scour.

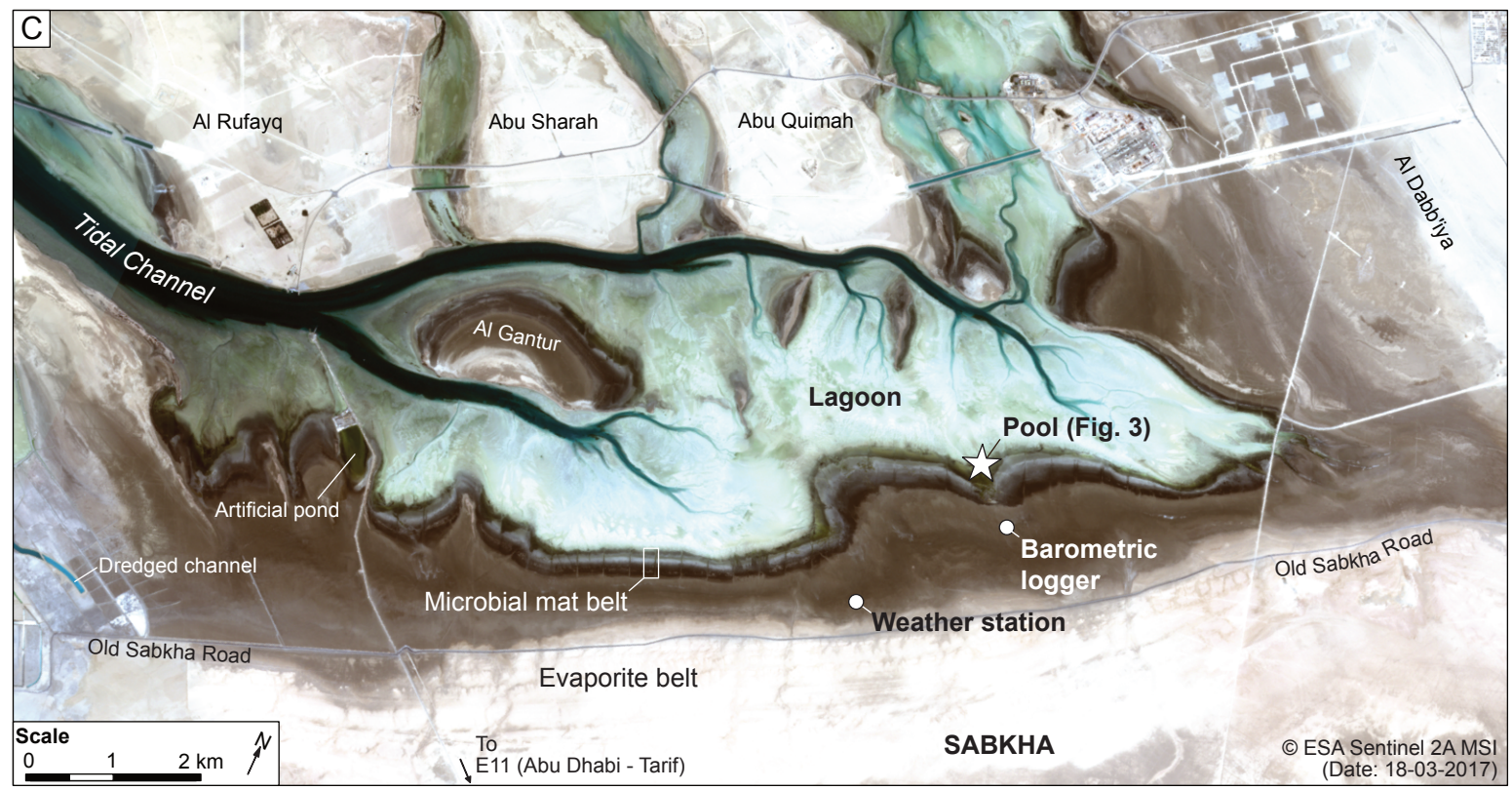
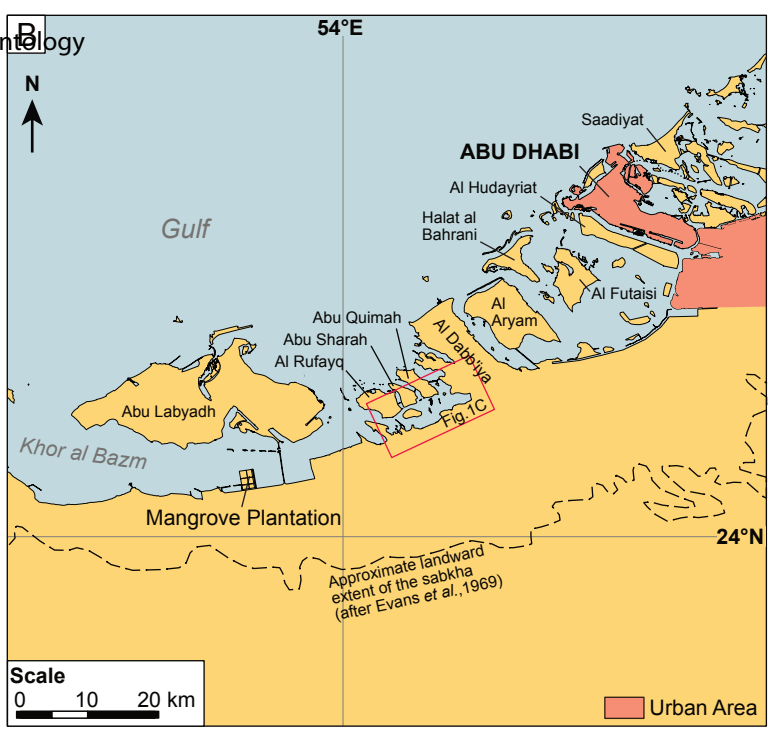
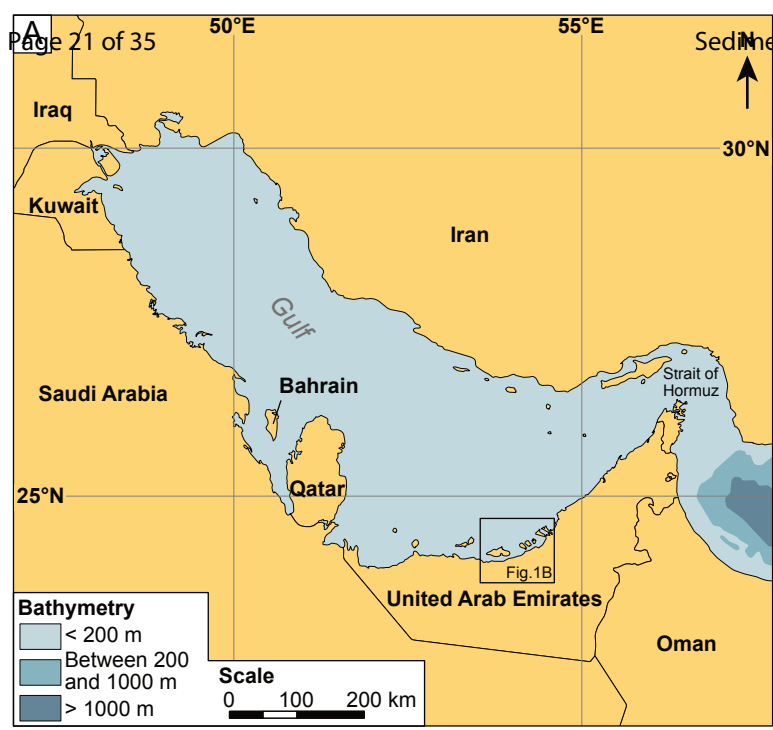
712 Figure 12. Thin section photomicrographs of stromatolites and thrombolites from the pool  
713 margin. A) Lithoclasts embedded in a stromatolite are shown surrounded by microbial  
714 organic-rich rims (arrow). B) Mouldic porosity in a thrombolitic area filled by acicular  
715 aragonite cements (arrows). C, D) Acicular aragonite cements filling thrombolite  
716 intragranular pore space. E, F) Discontinuous organic seams define the microbial laminations  
717 observed in hand specimen (see Fig. 8B) (arrows).

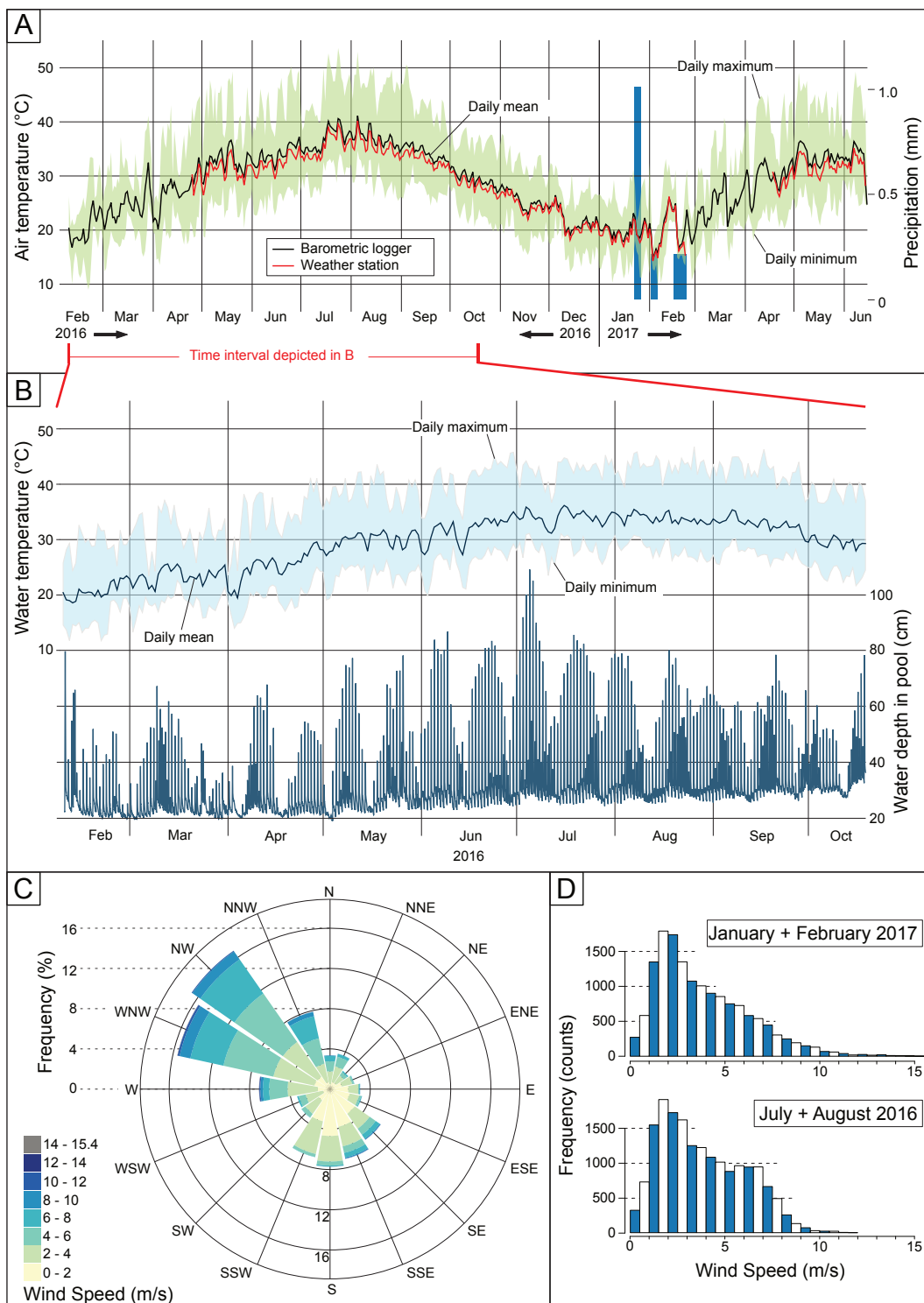
718 Figure 13. Scanning electron microscope micrographs of stromatolites and thrombolites. A,  
719 B) Extra-polymeric bacterial substance covering some areas of stromatolite specimen from  
720 the pool margin. Note bacterial filaments (white arrows). C) Crystal showing a very high Mg  
721 content indicative of proto-dolomite (arrow). The crystal is surrounded by aragonite cement  
722 and microbial EPS. D) Group of proto-dolomite crystals with very high Mg contents,  
723 surrounded by bacterial filaments, EPS and aragonite. E) Mineral precipitates in the  
724 microbialites showing cyanobacterial tubes (arrows). F) Acicular aragonite covered by a  
725 layer of EPS (arrows).

726 Figure 14. Schematic model of pool and microbial feature formation in the coastal sabkha of  
727 Abu Dhabi. A) Stratigraphy in the polygonal microbial mat zone prior to pool formation. B)  
728 Newly-formed pool with cyanobacteria colonising newly-formed subaqueous environments.  
729 C) Cementation of pool wall with onset of erosion. D) Pool margin collapse and formation of  
730 margin-parallel thrombolite band. E) Onset of new phase of pool wall cementation, continued  
731 growth of thrombolite, renewed erosion. F) Second generation of margin collapse, increasing  
732 isolation of phase 1 thrombolite.

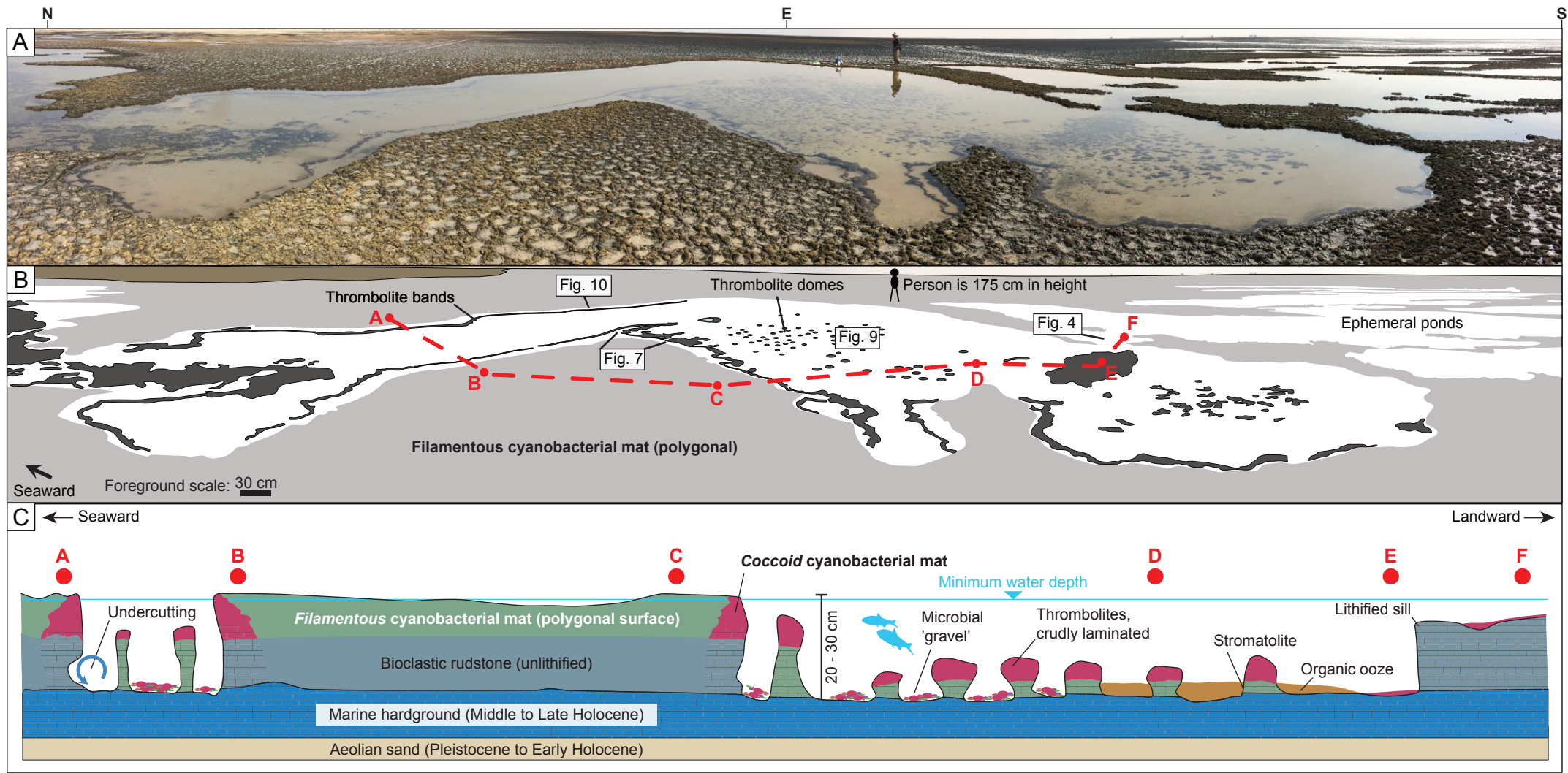
733 Figure 15. Schematic overview of the process that leads to permanent water coverage of the  
734 pool. During highest tides the pool is inundated by ~ 1m of water. During low tide the  
735 ephemeral ponds system slowly empties its excess water into the pool. Since this process  
736 takes much longer than one tidal cycle, the pool remains filled with water. At the margin,  
737 cascading water results in degassing, promoting the precipitation of aragonite cements.  
738 (MHW = mean high water)

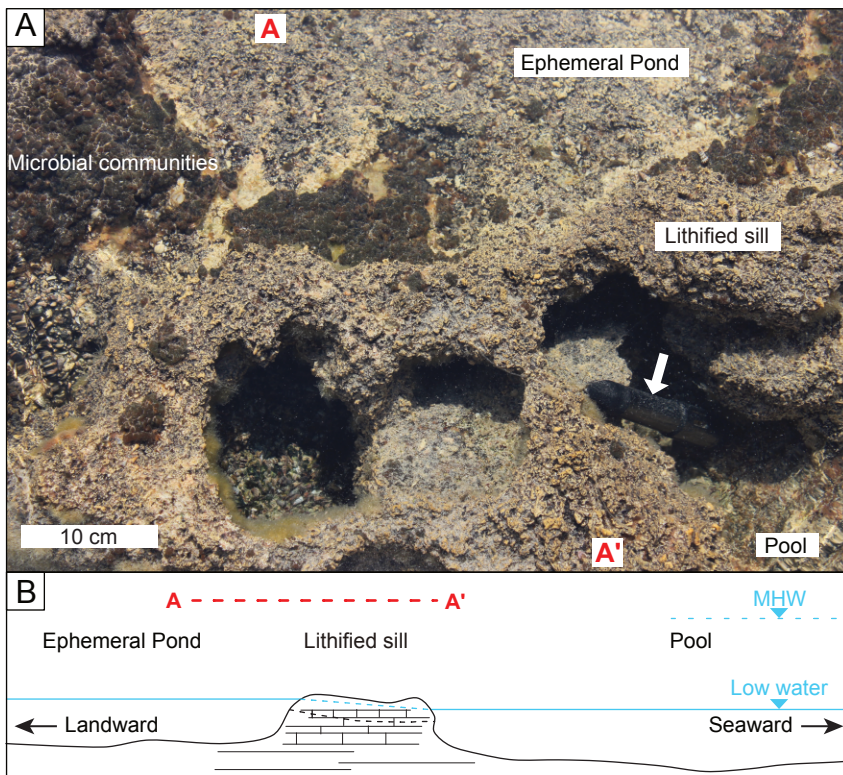
<b>Parameter</b>	<b>Location</b>				
	<b>Sabkha</b>	<b>Hamelin Pool</b>	<b>Lagoa Vermelha</b>	<b>Cayo Coco</b>	<b>Highborne Cay</b>
Air temperature (°C)	8.4 – 53.7	9.2 – 36.9	17.0 – 32.0	20.0 – 32.5	-
Water temperature (°C)	11.7 – 46.8	11.0 – 33.0	23.8 – 36.5	28.0 – 30.0	20.3 – 30.3
Salinity (‰)	75.0 – 93.0	15.8 – 88.1	40 – 84	30 – 80	35
Mean annual rainfall (mm)	72.0	210.4	900	1000	-
Annual evaporation (mm)	2750	2032 - 2286	1300	2100 – 2200	-
Tidal range (m)	1.0	1.0	1.0 – 1.7	0.3	1.0
Mean wind speed (m/s)	3.7	2.8	-	3.9	4.18

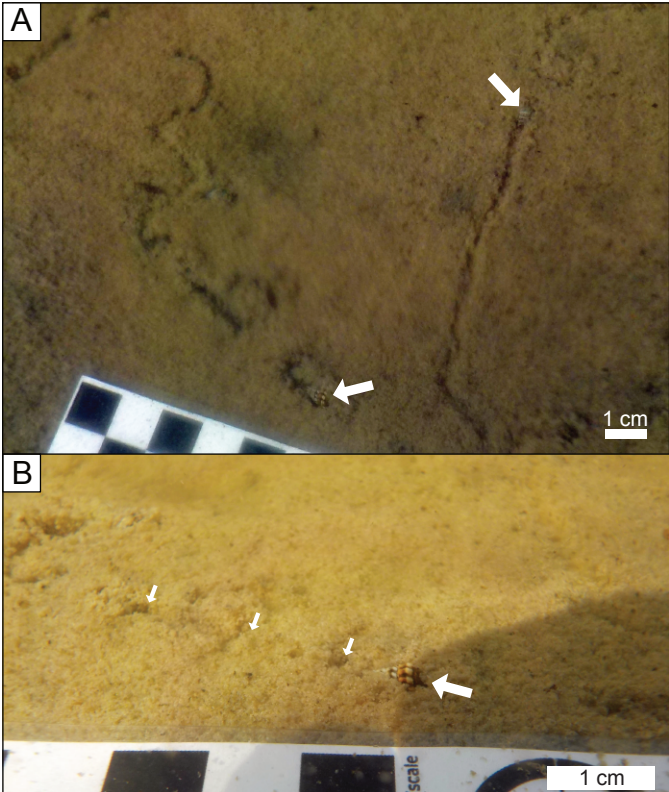




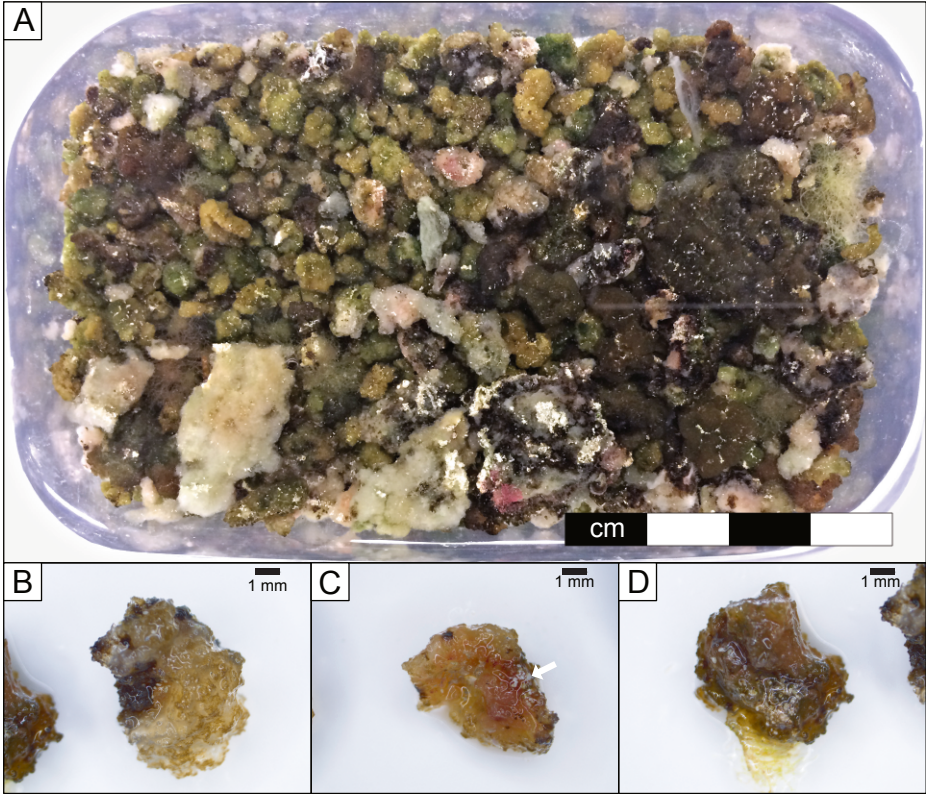


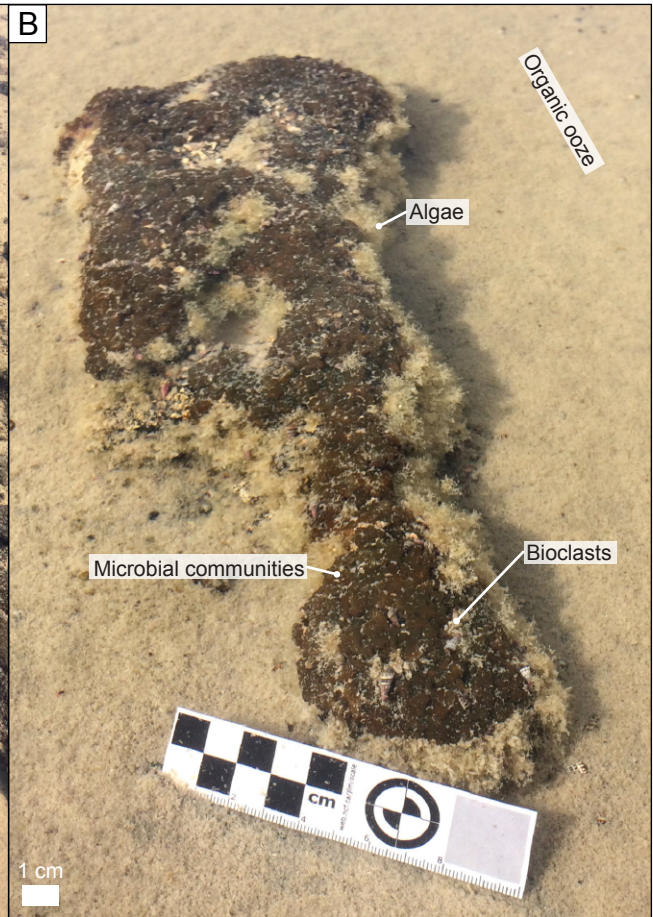
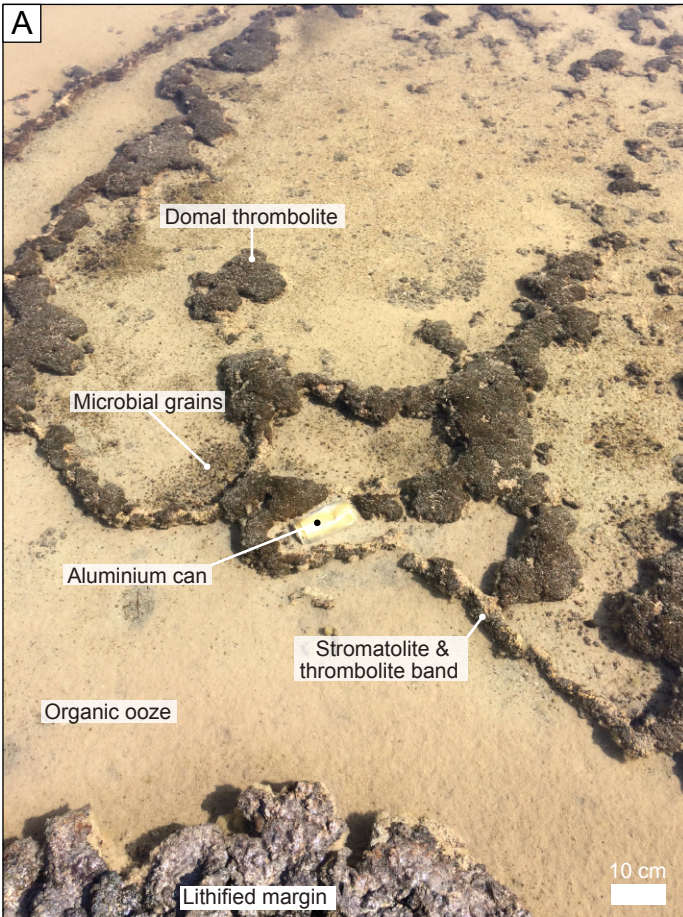
















Pustular microbial communities

