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1 **Cyanobacteria and algae in clouds and rain in the area of puy de Dôme, Central France**

2 Running Title: Cyanobacteria and algae in clouds and rain

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12 **Keywords:** bioaerosols, dispersal, microalgae, Chlorellaceae, Trebouxiales,

13 *Chroococciopsidales*

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23 **ABSTRACT:** The atmosphere contains diverse living microbes, of which the heterotrophic
24 community has been the most studied. Microbes with other trophic modes, such as
25 photoautotrophy, have received much less attention. Here, culture-independent and dependent
26 methods were used to examine the presence and diversity of oxygenic photoautotrophic microbes
27 in clouds and rain collected at or around puy de Dôme Mountain, central France. Cloud water
28 was collected from the summit of puy de Dôme (1465 m above sea level (a.s.l.)) for cultivation
29 and metagenomic analysis. Cyanobacteria, diatoms, green algae, and other oxygenic
30 photoautotrophs were found to be recurrent members of clouds, while green algae affiliated with
31 the Chlorellaceae were successfully cultured from three different clouds. Additionally, rain
32 samples were collected below the mountain from Opme meteorological station (680 m, a.s.l.).
33 The abundance of chlorophyll *a*-containing cells and the diversity of cyanobacteria and green
34 algae in rain were assessed by flow cytometry and amplicon sequencing. The corresponding
35 downward flux of chlorophyll *a*-containing organisms to the ground, entering surface ecosystems
36 with rain, varied with time and was estimated between ~1 and >300 cells cm⁻² day⁻¹ during the
37 sampling period. Besides abundant pollen from Pinales and Rosales, cyanobacteria of the
38 *Chroococciopsidales* and green algae of the Trebouxiales were dominant in rain samples.
39 Certain members of these taxa are known to be ubiquitous and stress-tolerant and could use the
40 atmosphere for dispersal. Overall, our results indicate that the atmosphere carries diverse, viable
41 oxygenic photoautotrophic microbes and acts as a dispersal vector for this microbial guild.

42 **IMPORTANCE:** Information regarding the diversity and abundance of oxygenic
43 photoautotrophs in the atmosphere is limited. More information from diverse locations is needed.
44 These airborne organisms could have important impacts upon atmospheric processes and on the
45 ecosystems they enter after deposition. Oxygenic photoautotrophic microbes are integral to

46 ecosystem functioning and some have the potential to affect human health. A better
47 understanding of the diversity and the movements of these aeolian dispersed organisms is needed
48 to understand their ecology, as well as how they could affect ecosystems and human health.

49 **INTRODUCTION**

50 The atmosphere harbors a diverse microbial assemblage that can be free-floating, adhered
51 to particles, or suspended in water droplets (e.g. clouds, fog, and rain). Bacteria are typically
52 present at concentrations between 10^4 to 10^5 cells m^{-3} in air (1) and about 10^3 to 10^4 cells mL^{-1} of
53 water in clouds (2). Microbes aerosolized from the surface environment including bodies of
54 water, soils, plant surfaces, and other sources can rise in the atmosphere, be incorporated into
55 clouds, and return to the surface via dry or wet deposition (for a general review, see Frölich-
56 Nowoisky et al. (3)). Some can help to form cloud droplets and ice crystals by acting as nuclei
57 for the condensation and freezing of water (4, 5). Over mid-latitude continental areas, the
58 microbial assemblage is often dominated by heterotrophic bacteria such as *Pseudomonas* and
59 *Sphingomonas* both in terms of abundance and activity (2, 6). These heterotrophic microbes have
60 been studied for their interaction with atmospheric processes potentially affecting cloud
61 chemistry (7). Other microbial guilds, such as oxygenic photoautotrophic microbes, have
62 received less attention regarding their abundance, diversity, and activity in the atmosphere.

63 Based upon previous studies, the main airborne oxygenic photoautotrophic microbes are
64 green algae, cyanobacteria, and diatoms (8). These are found in/on and can be aerosolized from
65 surface environments including lakes, trees, and building facades (9-12). Their abundance in the
66 air was reported to be affected by meteorological parameters such as wind direction and speed
67 (13). The aerosols containing cyanobacteria and algae are important to study due to their impacts
68 upon human health (11, 14), ice nucleation (15), and surface ecosystems by deposition and

69 colonization (16). In general, the atmospheric dispersal and cycling of oxygenic photoautotrophs
70 is not well documented (17). Cyanobacteria have been detected in clouds at variable abundances
71 as part of a few studies (2, 18, 19). However, the diversity of oxygenic photoautotrophs in clouds
72 remains understudied. Most studies have been conducted on atmospheric cyanobacteria and
73 algae in near-surface air environments, snow, and rain (20-22). For in-depth reviews about algae
74 and cyanobacteria in the atmosphere, see Tesson et al. and Wiśniewska et al. (8, 23).

75 This study aimed to investigate the diversity and abundance of cyanobacteria and algae in
76 clouds and rain at and around the puy de Dôme mountain (1465 m, 680 m above sea level (a.s.l.),
77 respectively), a mid-altitude rural area in central France. We investigated metagenomes of cloud
78 water and cultured cloud water samples for oxygenic photoautotrophic microbes by enrichment.
79 Additionally, chlorophyll *a*-containing cells were enumerated in rain samples, the corresponding
80 downward fluxes were estimated, and the diversity of cyanobacteria and algae was assessed by
81 targeted amplicon sequencing.

82 **RESULTS**

83 **Cyanobacteria and green algae in clouds**

84 Cloud water from the summit of the puy de Dôme was collected and used directly to
85 cultivate microbial oxygenic photoautotrophs. Cloud water was collected on 01/06/2010,
86 08/06/2010, and 29/06/2017 (dd/mm/yyyy) for enrichment cultures of oxygenic photoautotrophs
87 (referred to as 01062010, 08062010, and 29062017 cultures, respectively). The physical and
88 meteorological characteristics for each cloud during the cloud water collection process are
89 presented in Table S1. Diversity metrics for the enrichment culture communities are presented in
90 Table S2. The main eukaryotic oxygenic photoautotrophic microbes cultured were aeroterrestrial
91 epiphytic green algae of the Chlorellaceae (Figure 1A). The most dominant Chlorellaceae

92 operational taxonomic unit (OTU) in the 08062010 and 29062017 cultures, and second most
93 abundant in the 01062010 culture, was most closely associated with *Chloroidium saccharophila*
94 SAG 211-9a. The 01062010 culture was dominated by an unclassified eukaryote (Figure 1A),
95 whose taxonomy could not be resolved further. Surprisingly, no cyanobacteria were detected in
96 the oxygenic photoautotrophic cloud water enrichments (Figure 1B). Sphingomonadaceae,
97 Chitinophagaceae, and Rhizobiaceae bacterial families were dominant in the 01062010,
98 08062010, and 29062017 cultures, respectively (Figure 1B).

99 Along with the culture-dependent approach, we used metagenomics to explore cloud-
100 borne oxygenic photoautotrophs. Clouds were collected on 11/10/2013, 14/10/2013, and
101 05/11/2013 (denoted as 11102013, 14102013, and 05112013, respectively; Table S1). Cloud-
102 borne oxygenic photoautotrophs were identified based upon taxonomy in six cloud
103 metagenomes—three obtained for this study (11102013, 14102013, 05112013), and three that
104 were previously described (24) (17112014-1, 17112014-2, 17112014-3). Oxygenic
105 photoautotrophs were detected in all six cloud metagenomes, indicating that these were
106 persistent members of the microbial assemblage in clouds (Table 1). Bacillariophyceae (diatoms)
107 and Chrysophyceae (golden algae) were >1% of the SSU *rRNA* gene sequences across all six
108 clouds, while the cyanobacteria were <1% of the sequences (Table 1).

109 In addition to taxonomy, we investigated functional genes related to photosynthesis.
110 Overall, based on the UniProtKB identifiers and their associated gene ontology identifiers (GO
111 IDs), the functional genes related to photosynthesis by cyanobacteria were rare in the
112 metagenomes, ranging from 2 to 103 sequences in all clouds (Figure 2). The main genes
113 identified were related to *psb* genes, which encode for components of photosystem II (Data Set
114 S1). The dominant bacterial genera associated with the detected UniProtKB identifiers of the

115 photosynthesis-related genes were *Lyngbya*, *Synechococcus*, and *Pseudanabaena* (Figure 2). The
116 11102013, 17112014-2, and 17112014-3 clouds only had 2 functional gene sequences associated
117 with photosynthesis, and each were 100% associated with *Oscillatoria*, *Microcystis*, and
118 *Crocospaera*, respectively (Figure 2). Functional genes associated with photosynthesis by
119 eukaryotic photoautotrophs were not detected in any of the clouds.

120 Previously, Amato et al. (2) investigated the microbes that were potentially active in
121 clouds. DNA and RNA were extracted from cloud water and amplicons of fragments of the *16S*
122 *rRNA* and *18S rRNA* genes (for prokaryotes and eukaryotes, respectively) were generated. Based
123 on the ratio of the relative abundance of a specific OTU in the *16S* or *18S* rRNA and *16S* or *18S*
124 *rRNA* DNA of a sample, an OTU's activity can be gauged (25). This is not a perfect method of
125 assessing activity, but it can be used as a qualitative metric (26). Here we re-examined the data
126 specifically for different types of oxygenic photoautotrophic microbes. Eukaryotic algae had
127 RNA/DNA average ratios ranging from 0.59 to 2.11 and likely were mainly inactive (Table 2).
128 In contrast, the cyanobacteria appeared to be more active with RNA/DNA ratios averaging 41.69
129 and reaching values as high as 1100 (Table 2) Unfortunately, the cyanobacteria that were
130 detected were associated with unclassified phylotypes, and could not be classified further with
131 the sequence data available.

132 **Chlorophyll *a*-containing cells in rain**

133 We collected 16 rain samples over a 3-week period at our rural sampling site (Opme) and
134 assessed the chlorophyll *a*-containing cell abundance by flow cytometry. The main
135 meteorological parameters and chemical composition of the rain samples are shown in Table S3.
136 There were chlorophyll *a*-containing cells present in every rain event over the sampling period.
137 Their concentration ranged from 9 to 2750 cells mL⁻¹ of rainwater, corresponding to downward

138 fluxes of ~1 to 328 chlorophyll *a*-containing cells cm⁻² day⁻¹ being deposited with rain (Figure
139 3A). The chlorophyll *a* content in cells vary throughout their lives and environmental conditions
140 (27) and cell membranes could rupture during manipulation, allowing chloroplasts to be
141 detected. These factors could affect the enumeration of chlorophyll *a*-containing cells. In
142 addition to these measurements, the diversity of cyanobacteria and green algae in the rain
143 samples was assessed by amplicon sequencing of the *16S* and *18S rRNA* genes, specific for each
144 respective group. The diversity data from the sequencing is presented in Table S4. There was a
145 significant positive correlation between the Chao1 index and biomass (the number of chlorophyll
146 *a*-containing cells; $p = 0.04$, $\rho = 0.52$, Data Set S2). Additionally, there was a significant positive
147 correlation between air masses coming from the west and the number of chlorophyll *a*-
148 containing cells ($p = 0.02$; $\rho = 0.57$, Data Set S2). Despite the variable concentrations/fluxes, we
149 enriched oxygenic photoautotrophic microbes from each rain event by incubation in MWC (28)
150 or BG11 medium (ATCC medium 616), as indicated by observation of green-tinted cultures.

151 **Oxygenic photoautotrophs in rain**

152 The dominant cyanobacteria present varied across rain events. Unclassified members of
153 the *Nostocales* order were the most prominent cyanobacteria that range from not detected to
154 100% of the OTUs in the rain samples (Figure 3B). The *Chroococcidiopsaceae* varied from not
155 detected to 75%, being most represented by an OTU associated with an uncultured
156 cyanobacterium. *Aliterellaceae* ranged from not detected to 50% of OTUs with the most
157 dominant OTU being closely related to *Aliterella* CENA595 (29). The *Microcoleaceae* ranged
158 from not detected to 8.9% of OTUs with the dominant OTU matched to *Tychonema* CCAP 1459-
159 11B (30) (Figure 3B). The *Microcoleaceae* family's relative abundance was significantly
160 positively correlated with pH ($p = 0.036$, $\rho = 0.53$; Data Set S2) and with the concentration of

161 chlorophyll *a*-containing cells ($p = 0.028$, $\rho = 0.55$; Data Set S2). These cyanobacteria might
162 have been an important component of the group of chlorophyll *a*-containing cells that were
163 detected. There were no cyanobacteria present on 25 May, 01 June, and 06 June (Figure 3B).

164 Among green algae, the Trebouxiales order dominated the rain samples accounting for
165 0.1 to 6% of OTUs across the rain events with a peak in abundance on 25 May (Figure 3C). An
166 uncultured *Trebouxia* alga was the most dominant sequence associated with OTUs matched to
167 this order. Other Trebouxiophyceae varied in abundance from not detected to 0.76%. The other
168 green algae detected were lower in abundance. Members of the order Chlorellales were only
169 detected in some rain events in relative abundances up to about 0.3% (13 May; Figure 3C). The
170 Chlamydomonadales detected in some of the rain samples peaked in relative abundance on 25
171 May at about 0.11% (Figure 3C). Members of the Pedinomonadales were only detected on 3
172 May at about 0.01% relative abundance (Figure 3C).

173 Other than cyanobacteria and algae, pollen was also detected in rain (Figure 3D).
174 Notably, pollen from the order Pinales dominated the eukaryotic sequences at early time points,
175 and then decreased in relative abundance across the sampling time frame (Figure 3D). The order
176 Pinales was dominated by phylotypes associated with the genus *Pinus*, pine trees, which are
177 abundant in the sampling region. During the final days of the sampling period the relative
178 abundance of Pinales decreased and the relative abundances of the Rosales and Poales increased.
179 Therefore, there were significant negative correlations between the relative abundances of
180 Pinales and Rosales ($p = 1.7 \times 10^{-9}$, $\rho = -0.96$; Data Set S2) as well as Pinales and Poales ($p =$
181 2.5×10^{-6} , $\rho = -0.90$). There were no significant correlations between pollen sequences' relative
182 abundances and chemical/meteorological parameters or geographic origins of the air masses
183 (Data Set S2).

184 Finally, it should be mentioned that taxa other than oxygenic photoautotrophic microbes
185 were abundant in the amplicons from rain samples, although the primers used to generate *16S*
186 *rRNA* and *18S rRNA* gene amplicons were designed for excluding them (Cya 359f/Cya 781r and
187 Euk528f/CHLO02r, respectively (31-33)). The use of these primers might need to be reexamined
188 for future investigations.

189 **DISCUSSION**

190 **Biodiversity and sources of cyanobacteria and algae in clouds and rain**

191 Various genera of microalgae and cyanobacteria associated with the terrestrial and
192 phyllosphere environments have previously been found in atmospheric samples (8). Microbes of
193 the phyllosphere experience many of the same stressors as airborne microbes (34), so they might
194 be better suited to living and surviving in the atmosphere than those from other sources. Overall,
195 our data indicated that many of the microalgae and cyanobacteria detected in clouds and rain
196 were of terrestrial origin, but some were likely of marine origin.

197 The BG11 media used for enrichment of cloud water selected for freshwater
198 cyanobacteria and algae. The predominant photoautotrophic microbes enriched from clouds
199 under these conditions were green algae of the Chlorellaceae (Figure 1B). The main OTU was
200 associated with *Chloroidium saccharophila* SAG 211-9a. This organism was originally isolated
201 from tree sap (35, 36), which suggests again that the phyllosphere could be an important source
202 of atmospheric microalgae. Furthermore, *Chloroidium* is known to have a wide variety of
203 habitats that range from freshwater to being lichen-associated (36). *Chloroidium* isolated from
204 atmospheric samples were able to grow in a simulated freshwater environment (15).

205 We focused on enrichment of freshwater oxygenic photoautotrophs because these
206 organisms might be able to colonize the local environment after atmospheric transport. The puy

207 de Dôme region is inland, far from oceanic sources, but it has been noted that the microbial
208 communities in clouds at puy de Dôme often retain a marine signature (2, 24, 37, 38). Most of
209 the clouds that were characterized for this study were of marine origin. However, the single
210 cloud of continental origin (11102013) had similar relative abundances of SSU sequences of
211 oxygenic photoautotrophic microbes compared to the marine clouds (Table 1).

212 Cyanobacteria were present in low relative abundances in the metagenomes across three
213 clouds sampled for this study (11102013, 14102013, and 05112013) and the three clouds that
214 were characterized previously (17112014-1, 17112014-2, 17112014-3) (24) (Table 1). Based
215 upon the UniProtKB annotation of functional genes detected in the metagenomes, however, there
216 were diverse cyanobacteria present. Different cyanobacterial genera associated with the marine
217 environment were detected including *Lyngbya*, *Crocospaera*, and *Synechococcus*. *Lyngbya* is a
218 common cyanobacterium found in microbial mats in intertidal areas and *Crocospaera* and
219 *Synechococcus* are common in the ocean (39, 40). Additionally, *Synechococcus* and *Lyngbya*
220 have both been found in air samples (10, 41). Besides marine cyanobacteria, there were also
221 terrestrial cyanobacteria such as *Gloeocapsa* and *Gloeobacter*, which were all present in the
222 14102013 metagenome (Table 1). *Gloeocapsa* has been found in the atmosphere previously (42,
223 43) and *Gloeobacter* has been found in terrestrial environments including caves (44).
224 Cyanobacteria such as *Lyngbya* and *Gloeocapsa* were found to have a wide distribution amongst
225 various atmospheric locations in Asia, Antarctica, Polynesia, Nearctic, and the Palearctic (11), in
226 addition to some cloud samples from this study. Their heterogeneous presence in this study could
227 be due to various reasons relating to meteorological conditions of the different locations over
228 which the air mass traveled.

229 Despite detecting a clear marine signature in the cloud metagenomes, many of the algae
230 and cyanobacteria in rain at the sampling site/time could be from other environments. The major
231 cyanobacterial phylotypes detected in rain were of the order *Chroococciopsidales*. This
232 contains two families: *Chroococciopsaceae* and *Aliterellaceae*. A major OTU detected was
233 related to the genus *Aliterella* whose type strain was isolated from the Atlantic Ocean off the
234 coast of Brazil (29). This genus has also been found on rocks in the Atacama Desert (45) and in
235 rain collected in Hawaii (43). It has a cosmopolitan distribution and might be stress tolerant (as
236 per its presence in a desert and in the atmosphere). Uncultured members of the
237 *Chroococciopsaceae* and unclassified members of the *Nostocales* were also detected in the
238 samples. Members of these families have been previously detected in atmospheric samples in
239 Hawaii (43). A member of the *Microcoleaceae*, matching with *Tychonema* CCAP 1459-11B, a
240 freshwater cyanobacterium (30), was dominant in some of the rain samples.

241 There were diverse algae present in clouds. Chlorellales members were cultivated from
242 clouds (Figure 1A) and detected in rain (Figure 3C). Previously, green algae such as *Chlorella*
243 and *Chlorococcum* were cultured from rain (20). Overall, the most abundant group of green
244 algae in our rain samples were of the Trebouxiales order (Figure 3C), with phylotypes associated
245 with uncultured *Trebouxia* being the most dominant. Certain members of the *Trebouxia* are
246 prominent phycobionts of lichen (46) and have been detected in air and rain samples in South
247 Korea (47) and air samples in Spain (48). The Trebouxiophyceae, which includes Chlorellales
248 and Trebouxiales, have also been found in the marine environment (49), suggesting a
249 cosmopolitan distribution of this taxonomic class.

250 Other than microbial oxygenic photoautotrophs, pollen was detected in the rain samples.
251 The pollen did not correlate with any of the physicochemical parameters we measured. This fact,

252 along with their large sizes, suggests the pollen grains were most likely derived from local
253 sources, emphasizing the importance of local landscapes on the atmospheric biological content
254 (50). The types of pollen that were detected varied over the three-week sampling period. In
255 anemophilous species, pollen is obviously largely dispersed aurally with well-known seasonality
256 (47). The beginning of our sampling (May) likely corresponded with pine pollen season and the
257 end could have corresponded with the end of the pine pollen season (June).

258 Because there were conflicting correlations with a western origin of rain air masses with
259 air temperature and the number of chlorophyll *a*-containing cells, respectively, it is not possible
260 to discriminate between a geographical source and an actual environmental temperature effect
261 with this data. In a recent study, which included our sampling area, local landscapes and
262 meteorological conditions were significant factors affecting the composition of the airborne
263 microbial community (50).

264 Based upon the taxonomic affiliations, the cyanobacteria and algae detected in rain were
265 likely derived from a mixture of locations including marine, freshwater, and terrestrial habitats.
266 Many of the marine species were probably transported by the clouds and some of the freshwater
267 and terrestrial phylotypes were transported by a combination of the clouds and the locally
268 generated bioaerosols washed-out of the air column by the rain.

269 **Survival and potential activity**

270 Since many of the oxygenic photoautotrophs in our rain samples have a cosmopolitan
271 distribution, atmospheric transport could be a substantial dispersal mechanism for these
272 organisms (51). However, for successful dispersal the organisms must survive atmospheric
273 stressors, including osmotic shocks, light, oxidants, and freezing. Many Trebouxiophyceae
274 tolerate desiccation and other stressors that they would experience in the atmosphere (52).

275 Osmotic shocks and dehydration stress occur in both terrestrial and phyllosphere habitats. It was
276 speculated that airborne cyanobacteria were more tolerant to different relative humidities than
277 algae (53). The cellular biochemical properties of cyanobacteria and algal cells may be key to
278 their survival in the atmosphere. Algae of the genus *Chloroidium*, such as *Chloroidium sp.*
279 UTEX 3007 (54), can potentially survive a variety of atmospheric stressors because of
280 intracellular desiccation resistance-promoting sugars (36, 54). Some green algae isolated from
281 the atmosphere have carotenoids, thick cells walls, and intracellular antioxidants which help
282 them to survive atmospheric stressors (55). Many members of the Trebouxiophyceae have
283 mycosporine-like amino acids in the cytoplasm which help to protect the algae from UV
284 radiation and oxidants (56). Mycosporine-like amino acids are also found in cyanobacteria such
285 as *Lyngbya purpurem* (57). Other cyanobacteria such as *Synechococcus*, in particular, are known
286 to survive various osmotic, oxidative, and temperature stressors (40). The *Chroococcidiopsaceae*
287 family was prominent in our rain samples and a member of this family, *Chroococcidiopsis*, is
288 well-known to be highly tolerant to desiccation and radiation (58). The relative abundances of
289 this family were negatively correlated with increasing air temperature (ranging from 14°C to
290 22°C; Table S3), a trend that was previously reported for the abundance of various cultivated
291 algae (59).

292 Amongst atmospheric stressors, freeze-thaw stress has been found to affect bacteria the
293 most (60). The ability of microbial cells to tolerate freezing can be an important factor for
294 survival during atmospheric transport. Algae isolated from atmospheric samples were able to
295 survive freezing to a greater extent than aquatic algae (15). The 01062010 and 08062010 cloud
296 water enrichments were revived after being stored in glycerol at -20°C for several years. This
297 method for storing microorganisms could have an impact upon the mortality of cells and thus

298 potential composition of the microbial community (61). Green algae and cyanobacteria, the most
299 abundant oxygenic photoautotrophs in the atmosphere (41), are more amenable to
300 cryopreservation than other oxygenic photoautotrophs (62). The dominance of the Chlorellaceae
301 in the cloud enrichments could be due to a larger number of organisms of this taxon present in
302 the culture when it was frozen. This would increase the likelihood of survival of this taxon as
303 opposed to other lower abundant taxa. However, when comparing 08062010 (stored at -20°C for
304 several years) and 29062017 (which was never stored at -20°C) cultures, they shared a dominant
305 OTU in the family Chlorellaceae that matched with *Chloroidium saccharophila* SAG 211-9a.
306 The fact that this alga was the major phylotype in two cloud enrichments and survived
307 cryopreservation in the 01062010 and 08062010 cultures supports its tolerance to atmospheric
308 stressors.

309 In addition to their vegetative states, many cyanobacteria and algae go through different
310 stages of development. Many of the resting states that the cells manifest, which are also part of
311 their reproductive cycle, impart stress-tolerance and allow the microbe to survive harsh
312 conditions (for a review see Ellegaard and Ribeiro (63)). For example, unequal autospores are
313 used for reproduction in *Chloroidium* and small autospores could be amenable to aeolian
314 transport (36). Small resting states would be more aerodynamically favorable for long-distance
315 transport as compared to larger vegetative cells.

316 While transiting through the atmosphere, if oxygenic photoautotrophs can survive
317 atmospheric stressors, they have the potential to be metabolically active. After re-assessing a
318 previous study on RNA/DNA ratios using rRNA amplicons, the eukaryotic algae were found to
319 be relatively inactive with low ratios of RNA/DNA (Table 2). This could be consistent with their
320 presence as resting states (e.g. spores) (63). Alternatively, this could indicate that cellular

321 functions of the eukaryotic algae were greatly affected by atmospheric stressors. In contrast the
322 cyanobacteria appeared to be more active with RNA/DNA ratios of not detected to 1100 (Table
323 2). Thus, they could affect cloud water droplet chemistry via carbon fixation, carbon and
324 nitrogen metabolism, or oxidant detoxification, similar to how heterotrophic bacteria were
325 hypothesized to be affecting cloud droplets (24). Consistent with other data, different genes
326 associated with oxygenic photosynthesis were persistently present in cloud metagenomes.
327 Nevertheless, not all genes associated with oxygenic photosynthesis and carbon fixation were
328 detected: for instance, no genes encoding for ribulose-1,5-bisphosphate carboxylase/oxygenase
329 (RuBisCO) were annotated (Data Set S1).

330 **Potential impacts on ecosystems**

331 The oxygenic photoautotrophs that are suspended in the air will eventually return to
332 different surface environments through dry or wet deposition (e.g. rain). Some eukaryotic algae
333 can promote ice formation, and thus could promote their deposition to new surface environments
334 by affecting cloud physics and precipitation patterns (15). All the rain samples analyzed
335 contained viable, culturable, aerially dispersed oxygenic photoautotrophs. As a result, rain
336 undoubtedly acts as a dissemination vector for oxygenic photoautotrophs to colonize new
337 environments. Near the rain sampling site of Opme is Aydat Lake, a freshwater lake of
338 approximately 65 hectares ($6.5 \times 10^9 \text{ cm}^2$). The cyanobacteria and green algae that were
339 transported through the atmosphere in this area were thus likely to be deposited in this lake
340 ecosystem, as well as other locations. Extrapolating the observed flux of chlorophyll *a*-
341 containing cells with precipitation to the surface area of the lake indicates that 10^9 - 10^{12}
342 chlorophyll *a*-containing cells enter the lake per each rainy day. These values are specific for this
343 region and sampling time and would likely differ compared to other areas in the world and/or

344 other periods of the year. Whether the rain-borne microbes will be successful in their new
345 environment will vary depending upon the microbe's viability and physiology, as well as
346 atmospheric conditions (64). These new organisms could impact the local water quality and
347 ecology, but the exact effects will depend upon their functional capabilities.

348 The ability to reproduce under different conditions is very important to dispersal success
349 (15). *Trebouxia*, the most abundant green algae detected in our rain sequences, can be considered
350 a "generalist" due to its ability to grow in freshwater and brackish water conditions (15). If these
351 green algae can reproduce after atmospheric transport and deposition, their colonization could
352 impact their new environments, just as aerially dispersed bacteria and fungi can affect the
353 microbial composition and functioning of the environments they enter (65). Previously, the
354 primary production rate of autotrophic communities in water was reported to be negatively
355 impacted by aeolian-transported microbes (66). Notably, green algae were suggested to be better
356 colonizers of water than aerially dispersed diatoms or cyanobacteria (67).

357 The transported microbes could also affect human health. It has been speculated that the
358 allergenic alga *Gonyostomium semen* could be aerially dispersed, thus leading to its expansion
359 into new ecosystems (8). Here, phylotypes associated with toxin-producing oxygenic
360 photoautotrophs including *Microcystis*, *Scytonema*, *Tychonema*, and *Anabaena*, whose toxins
361 can have varying effects upon humans and other microbes (68-70), were detected in clouds
362 (Figure 2) and rain (Figure 3). This illustrates the aerial spread of microbes of health-related and
363 ecological interests and the possibility that they enter ecosystems with precipitation. Once they
364 enter a new environment such as Aydat Lake, they could produce toxins which could affect
365 human health or even the health of other algae, which are important to primary production. Even
366 if a microbe does not establish a lasting presence in a system, it still could have long-term

367 impacts upon the resident microbial community (71). Through these methods, aeolian
368 transported oxygenic photoautotrophs could impact the health and biogeochemical functioning of
369 a lake ecosystem.

370 **Conclusions and Perspectives**

371 Our study indicates that there are diverse oxygenic photoautotrophs among the microbial
372 assemblages carried by the atmosphere. We detected cyanobacteria, diatoms, as well as red,
373 green, and golden algae as persistent members of cloud microbial assemblages. Many of the
374 organisms detected were associated with stress tolerant phylotypes that have widespread
375 distributions, and some were toxin producers with potential ecological and health-related
376 impacts. We were able to culture some of them, mostly the aeroterrestrial epiphytic green algae
377 of the Chlorellaceae. Cyanobacteria were not detected in cloud water enrichments under the
378 conditions utilized, but molecular approaches indicated that these are present and probably active
379 in clouds. Genes encoding for part of photosystem II of various cyanobacteria were present in
380 metagenomes. Rain acted to mediate the dispersal of many of these diverse oxygenic
381 photoautotrophs, with members of *Chroococciopsidales* and Trebouxiales being notable, into
382 new ecosystems. Despite small sample sizes and short time periods, our study provides key
383 information on the aeolian transport of cyanobacteria and green algae that is currently missing.
384 Microbiologists, ecologists, and meteorologists should work together to conduct long-term
385 studies to identify the sources of the oxygenic photoautotrophic microbes in rain and determine
386 their emission/depositional fluxes. Aeolian dispersal is more conducive to certain microbes and
387 understanding the physicochemical effects on their dispersal patterns and the exact biological
388 and ecological implications of their dispersal is an understudied aspect of aerobiology.

389 MATERIALS AND METHODS

390 Cloud and rain sampling

391 Clouds and rain were sampled from meteorological stations operated by the Observatoire
392 de Physique du Globe de Clermont-Ferrand (OPGC) and parts of the national COPDD
393 atmospheric survey network.

394 Clouds were sampled at the puy de Dôme (PDD) station (1465 m a.s.l., 45.7720° N,
395 2.9655° E) on 01/06/2010, 08/06/2010, and 29/06/2017 (dd/mm/yyyy) as described previously
396 (38) (see Table S1 for sampling details). Clouds were sampled again at the same site on
397 11/10/2013, 14/10/2013, and 05/11/2013 (sample names 11102013, 14102013, 05112013,
398 respectively), as described previously (2, 24). The cloud water samples were filtered through a
399 47 mm, 0.22 µm porosity membrane (MoBio 14880-50-WF) in a laminar flow hood, cut in half,
400 and half-filters were stored at -80°C prior to use for metagenomic analyses following previous
401 protocols (24). Backward trajectories of the air masses were generated with HYSPLIT (Figure
402 S1) (72).

403 Rain was collected at the Opme meteorological station (680 m a.s.l., 45.712500° N,
404 3.090278° E) with an automated wet-deposition sampler (Eigenbrodt NSA 181/KHS), as
405 previously described (73). Briefly, eight autoclaved 1-L glass bottles were kept at 4°C in the
406 chamber with a new bottle designated for collection every 24 hours. Before putting new bottles
407 in the carousel, the funnel was disinfected with 70% ethanol and rinsed with sterile water. The
408 funnel was covered by a polytetrafluoroethylene (PTFE) lid that automatically opened when
409 precipitation occurred, and which prevented dry deposition between rain sampling events. All
410 rain water samples were collected and transported to the laboratory within 24 hours of the
411 precipitation event. Sixteen rain samples were collected between 23 May, 2018 and 13 June,

412 2018. The rain water was filtered through 0.22 μm porosity membranes (MoBio 14880-50-WF)
413 and membranes were stored at -20°C for later DNA extraction. The pH of filtered rain was
414 determined and ion concentrations were measured by ion chromatography (Dionex, Sunnyvale,
415 CA, USA) as previously described (2). The backward trajectories of the air masses (Figures S2)
416 were generated with HYSPLIT (72).

417 **Cultivation of photosynthetic microbes**

418 Cloud water samples collected on 01/06/2010, 08/06/2010, and 29/06/2017
419 (dd/mm/yyyy) were incubated in BG11 medium (ATCC 616) under constant light ($75 \mu\text{E}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$)
420 ¹) at 20°C in the INFORS HT Multitron II equipped with Sylvania Gro-Lux F15W/Gro lights
421 while shaking at 80 rpm to produce cultures named 01062010, 08062010, and 29062017,
422 respectively. We used BG11 medium, which is designed to favor the growth of cyanobacteria
423 and chlorophytes from freshwater environments, not diatoms. The study site is far from the
424 ocean, thus we focused on cultivating organisms that could successfully survive if deposited in
425 the local environment. This excluded marine organisms. The 01062010 and 08062010 cultures
426 were preserved in 15% glycerol and stored at -20°C for about seven years. Once the 29062017
427 culture was established, the others were revived, and all were incubated in BG11 media at 15°C
428 under constant illumination ($75 \mu\text{E}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$) at 80 rpm. After incubating for 7 days, 5 mL of each
429 culture was centrifuged at $8,000 \times g$ for 3 minutes and the cell pellet was stored at -20°C until
430 DNA extraction.

431 All rain samples were cultivated in MWC (28) or BG11 media. Visual observation of
432 green tinted cultures indicated successful enrichment of oxygenic photoautotrophs.

433 **DNA Extractions**

434 DNA from the cloud enrichment cultures was extracted with a modified phenol-
435 chloroform based protocol (74). A 50 mM glucose/10 mM EDTA/ 25 mM Tris-HCl solution was
436 added to the cell pellet, then five freeze-thaws were performed by freezing at -80°C in ethanol
437 (200 proof) and heating to 55°C. Lysozyme (0.4 mg L⁻¹) and EDTA (0.5 M) were added, and
438 samples were shaken at 100 rpm at room temperature for 5 min. Ten percent SDS was added and
439 followed quickly by extraction with phenol: chloroform: isoamyl alcohol (25:24:1). The
440 emulsion was separated by centrifuging at 16,000 × g for 3 min. The resulting aqueous phase
441 was extracted with phenol: chloroform: isoamyl alcohol and centrifuged again. The aqueous
442 phase from the second extraction was precipitated with 200 proof ethanol (1 mL) and 2 µg
443 glycogen.

444 DNA was extracted from half of the cloud filters and from the whole rain filter using a
445 MoBio PowerWater isolation kit (now, Qiagen, Hilden, Germany) following the manufacturer's
446 protocol.

447 **Cloud Enrichment PCR and Sequencing**

448 DNA extracts from cloud enrichments were sent for PCR amplification and sequencing
449 by MR DNA (www.mrdnalab.com, Shallowater, TX, USA). The *16S rRNA* amplicons were
450 generated using the primer set 515f/806r (75) with a barcode on the forward primer (Table S2).
451 HotStarTaq Plus Master Mix Kit (Qiagen, USA) was used for amplification with the following
452 conditions: 94°C for 3 min, followed by 30 cycles of 94°C for 30 sec, 53°C for 40 sec and 72°C
453 for 1 min, and ending with a final elongation at 72°C for 5 min. To generate the *18S rRNA* gene
454 amplicon, PCR using 1391f (76) and Eukbr (77), with a barcode on the forward primer (Table
455 S2) was performed with HotStarTaq Plus Master Mix Kit (Qiagen, USA) under the following
456 conditions: 94°C for 3 min, followed by 35 cycles of 94°C for 30 sec, 53°C for 40 sec and 72°C

457 for 1 min, and ending with a final elongation at 72°C for 5 min. The products were confirmed on
458 a 2% agarose gel. The *16S rRNA* gene amplicons were sequenced with an Ion S5 XL instrument
459 (Ion 530 chip) following the manufacturer's guidelines. The *18S rRNA* gene amplicons were
460 combined in equal proportions and purified using calibrated Ampure XP beads (Beckman
461 Coulter). These were used to prepare the DNA library for Illumina MiSeq 2×300 bp paired end
462 sequencing following the manufacturer's protocols. The corresponding sequence files were
463 deposited at the EMBL-EBI European Nucleotide Archive under the study accession number
464 PRJEB35708.

465 **Rain Samples PCR and Sequencing**

466 For the DNA extracts from rain samples, portions of the *16S rRNA* genes targeting
467 cyanobacteria (31) and of the *18S rRNA* genes targeting green algae (78) were PCR-amplified.
468 The *16S rRNA* gene PCR reaction was composed of 34.3 µL sterile water, 5 µL 10X Green PCR
469 buffer without Mg (ThermoFisher Scientific), 0.2 µL PlatinumTM Taq DNA Polymerase (10 U
470 µL⁻¹; ThermoFisher Scientific), 3 µL 25 mM MgCl₂, 1 µL 10 mM dNTPs, 2 µL 10 µM forward
471 primer Cya359 F (31), 2 µL 10 µM reverse primer Cya 781 R modified from Nübel et al. (31),
472 and 1 µL of template DNA (5-10 ng). The primer sequence of the modified reverse primer from
473 Nubel et al. (31) was 5' GACTACWGGGGTATCTAATCCCWTT 3'. The conditions were as
474 followed: 94°C for 2 min, then 35 cycles of 94°C for 30 sec, 55°C for 30 sec, 72°C for 30 sec,
475 and a final extension of 72°C for 5 min. For the *18S rRNA* gene PCR 33.7 µL sterile water, 5 µL
476 10X Green PCR buffer, 0.3 µL *Taq* DNA polymerase (5 U µL⁻¹; Fischer BioReagentsTM;
477 ThermoFisher Scientific), 3 µL 50 mM MgCl₂, 1 µL 10 mM dNTPs, 2 µL 10 µM EUK528f (32),
478 2 µL 10 µM CHLO02r (33), and 3 µL of template DNA (5-10 ng) were mixed. The conditions
479 (78) were as follows: 94°C for 5 min., then 35 cycles of 94°C for 30 s., 55°C for 30 s., 72°C for 1

480 min., and a final extension of 72°C for 5 min. Both sets of amplicons were purified with
481 MinElute PCR purification kit (Qiagen). Negative controls and positive controls were processed
482 alongside the samples. The positive control for the *16S rRNA* and *18S rRNA* gene amplifications
483 were DNA extracts from cultures of *Microcystis aeruginosa* PCC 9443 and an enrichment
484 culture of rain water collected on 27/05/2018 in MWC media (28) containing algae (27052018),
485 respectively. Purified amplicons from the rain sample DNA extracts were sent to GenoScreen
486 (Lille, France) and library preparation followed the Metabiote®v2.0 protocol (GenoScreen, Lille,
487 France) using 5 ng amplicon DNA. Library preparation quality was confirmed by capillary gel
488 electrophoresis. The samples were sequenced on one line with a 2×250-bp paired-end MiSeq
489 Illumina instrument. The sequence files were deposited at the EMBL-EBI European Nucleotide
490 Archive under the study accession numbers PRJEB35600, samples ERS4058723 to
491 ERS4058754.

492 **Bioinformatic analysis of PCR amplicons**

493 The *18S rRNA* amplicons from photosynthetic cloud enrichment cultures were
494 demultiplexed with sabre (<https://github.com/najoshi/sabre>) and paired ends were assembled
495 with the default parameters of the DADA2 (79) plugin in QIIME2 (v2020.2) (80). The *16S rRNA*
496 amplicons from oxygenic photoautotrophic cloud enrichment cultures were demultiplexed with
497 software from MR DNA LLC (<http://www.mrdnalab.com/mrdnafreesoftware/>). Both the *16S* and
498 *18S rRNA* gene rain amplicons were demultiplexed with the CASAVA v1.0 software (Illumina)
499 and the paired-ends were assembled with the FLASH tool (81) with trimming of poor-quality
500 reads (phred score <30) and a 97% similarity requirement.

501 All fastq files were imported into QIIME2 (v2020.2) (80), trimmed and denoised with
502 DADA2 (79) and grouped into amplicon sequence variants (ASVs). The ASVs were then

503 clustered into operational taxonomic units (OTUs) (97%) with the vsearch (82) plugin. A naive
504 Bayes scikit-learn classifier (83, 84) for each primer set was trained using extracted reference
505 full-length operon reads of the Silva v132 database (85) and used for classification. Taxa
506 associated with negative controls and prokaryotic reads in eukaryotic-targeted amplicons (and
507 vice versa) were filtered out from the samples. OTUs, that were of interest and lacked taxonomic
508 resolution from the identification, were queried manually against NCBI BLAST
509 (<https://blast.ncbi.nlm.nih.gov/Blast.cgi>) with the BLASTn algorithm in May 2020. Samples
510 were rarefied and analyzed for Shannon's diversity, Chao1, and observed OTUs in QIIME2
511 (v2020.2).

512 **Cloud metagenomes**

513 DNA extracts from cloud water samples 11102013, 14102013, and 05112013 were
514 amplified by MDA using GenomiPHI (GE Healthcare) to aid in generating sequencing libraries.
515 The libraries were prepared, sequenced, and analyzed as previously described (24) using an
516 Illumina MiSeq 2 × 300 bp (Genoscreen, Lille, France). The sequence files were deposited at
517 the EMBL-EBI European Nucleotide Archive under the study accession numbers PRJEB35574,
518 samples ERS4058712 to ERS4058714. Methods for the taxonomic and functional annotation of
519 the sequences have been described previously (24). In brief, taxonomic annotations were
520 assigned based upon the SILVA database (85). The oxygenic photoautotrophs were identified
521 based upon the taxonomy. The functional annotations were assigned based upon the UniprotKB
522 protein database (86). With the UniprotKB identifiers, associated genera and gene ontologies
523 identifiers (GO IDs) (87) were assigned. GO IDs associated with photosynthesis were selected.
524 Additional cloud metagenomes obtained from Amato et al. (24) were reanalyzed to investigate

525 the presence of oxygenic photoautotrophic microbes (sequences files ENA accession numbers
526 ERS2351639 to ERS2351641).

527 **Re-analysis of cloud RNA/DNA ratios**

528 Previously generated prokaryotic and eukaryotic ribosomal DNA and cDNA amplicons
529 were re-assessed (2). Sequence files accession number in NCBI BioProject PRJNA380262 were
530 reanalyzed to examine the potential activity of photosynthetic microbes in clouds. Known
531 oxygenic photoautotrophs were manually selected from the OTUs based upon their taxonomic
532 identification. The number of reads for each oxygenic photoautotrophic OTU were expressed as
533 a percentage of the total community for the cDNA (representing rRNA) and DNA sequences.
534 The RNA/DNA ratio for each OTU was calculated using the relative percentages. The OTUs
535 were grouped at the phylum or class level and the RNA/DNA ratios were averaged for each
536 taxonomic group. A taxon was considered to be active if the ratio was greater than 1, meaning
537 that the relative abundance of the taxon was greater in the rRNA amplicons rather than in the
538 *rRNA* gene amplicons. Phantom taxa, phylotypes present in the rRNA but not the *rRNA* genes
539 (25), were excluded in this analysis.

540 **Enumeration of chlorophyll *a*-containing cells**

541 Triplicates of fresh rain water samples of 0.45 mL were fixed with a final concentration
542 of 0.5% glutaraldehyde and stored at 4°C in the dark until flow cytometry analysis. Just before
543 analysis, fixed samples were amended with 1 vol of Tris-EDTA buffer (40 mM Tris; 1 mM
544 EDTA; pH 8.0) and analyzed by a BD FACSCalibur flow cytometer (Becton Dickinson,
545 Franklin Lakes, NJ) to enumerate chlorophyll *a*-containing cells. Detection was on FL3
546 (excitation 488 nm; emission 670 nm) with detection thresholds set at 500 mV for FL3 and 380
547 mV for SSC. Acquisition time was 315-400 s. at a flow rate between 66 – 72 $\mu\text{L min}^{-1}$ (high

548 setting). The flow rate was determined precisely for each sample through the acquisition time
549 and the gravimetric loss from the water sample. All data was analyzed in the software BD
550 CellQuest Pro™ (Becton Dickinson, Franklin Lakes, NJ). Chlorophyll *a*-containing cells were
551 expressed both as rainwater concentration (cells mL⁻¹) and as a surficial flux (cells cm⁻² day⁻¹).
552 The flux was calculated from the total number of cells in a rain sample for a given day divided
553 by the area of the collection surface.

554 **Statistical Analyses**

555 Statistical analyses were conducted in OriginPro Version 2019 9.6 (OriginLab
556 Corporation, Northampton, MA, USA). To correlate the presence of phylotypes with geographic
557 origin, the two types of air masses were defined and statistically analyzed as described in Pouzet
558 et al. (73).

559 **Data Availability**

560 The sequence files were deposited to the European Nucleotide Archive with project
561 accession numbers PRJEB35574 (cloud metagenomes; ERS2351639 to ERS2351641),
562 PRJEB35708 (cloud enrichments), and PRJEB35600 (rain samples; ERS4058723 to
563 ERS4058754).

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826 Table 1: Relative abundance of oxygenic photoautotrophic microbial small subunit (SSU)
 827 ribosomal sequences in cloud metagenomes. All clouds were of marine origin, based on back
 828 trajectories from NOAA HYSPLIT (Figure S1), except for sample 11102013, which was
 829 continental in origin.

Sample		11102013	14102013	05112013	17112014- 1	17112014- 2	17112014- 3
Reference	Common	This study	This study	This study	Amato et al. 2019	Amato et al. 2019	Amato et al. 2019
Taxa	Name	(%)	(%)	(%)	(%)	(%)	(%)
Prokaryotes							
Cyanobacteria	Blue-green algae	0.92	0.51	0.99	0.25	0.27	0.23
Eukaryotes							
Chlorophyta	Green algae	1.2	0.54	0.66	1.1	0.59	0.49
Trebouxiophyceae	Green algae	0.77	0.32	0.49	0.85	0.44	0.29
Rhodophyta	Red algae	0.15	0.054	0.066	0.040	0.022	n/a
Bacillariophyceae	Diatoms	1.8	2.4	1.5	1.5	1.4	1.7
Dinophyceae	Dinoflagellates	0.26	0.075	0.099	0.048	0.32	0.017
Chrysophyceae	Golden algae	1.6	3.1	1.8	2.3	2.6	2.8

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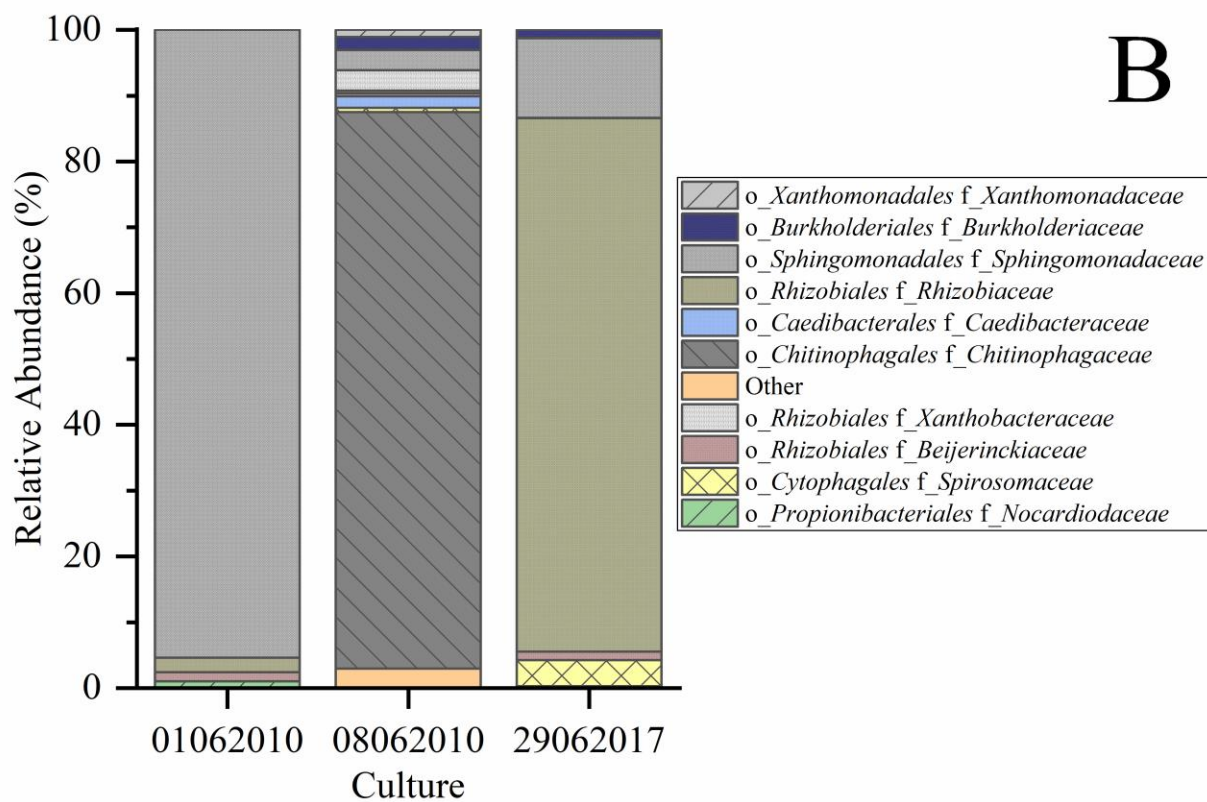
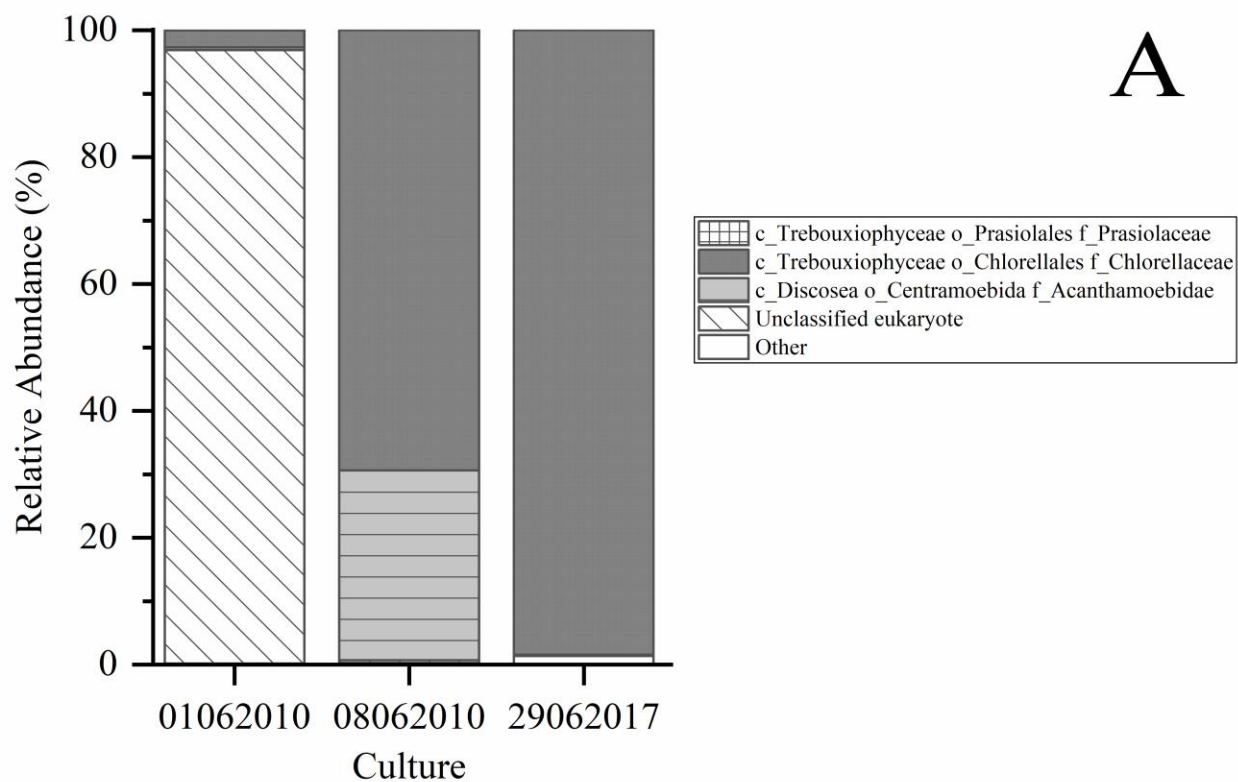
840 Table 2: Calculated RNA:DNA ratios of taxa including photosynthetic microbes in clouds
 841 (original data from Amato et al. (2)). The relative abundances of the taxa in the RNA and DNA
 842 reads were used to calculate ratios. The range is the minimum and maximum ratio observed for
 843 the corresponding taxon in the sample.

Group	Polluted Cloud		Continental Cloud	
	Average \pm Std Dev	Range	Average \pm Std Dev	Range
Prokaryotes				
Cyanobacteria	41 \pm 119	nd* - 1100	11 \pm 23	nd - 120
Eukaryotes				
Chlorophyta	0.85 \pm 0.94	nd - 2.9	1.2 \pm 1.7	nd - 7.1
Cryptophyta	0.59 \pm 0.82	nd - 2.9	0.67 \pm 0.85	nd - 2.8
Bacillariophyta	2.1 \pm 11.7	nd - 101	0.95 \pm 1.6	nd - 10
Dinophyceae	1.9 \pm 5.3	nd - 25	2.2 \pm 3.0	nd - 11
Chrysophyceae	1.7 \pm 5.4	nd - 40	1.3 \pm 1.5	nd - 8.7
Phaeophyceae	1.6 \pm 2.1	nd - 8.7	0.74 \pm 0.57	nd - 2.3

844 *nd: not detected

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848 Figure 1: Microbial composition of oxygenic photoautotrophic enrichments. Cloud water was
849 incubated with BG11 medium at 15°C under constant illumination. (A) The eukaryotic
850 population of the three cloud enrichments at the family level and their relative abundances in *18S*
851 *rRNA* gene reads. (B) The taxonomic classification and relative abundance of bacteria in *16S*
852 *rRNA* gene reads in all three cloud enrichments at the family level.

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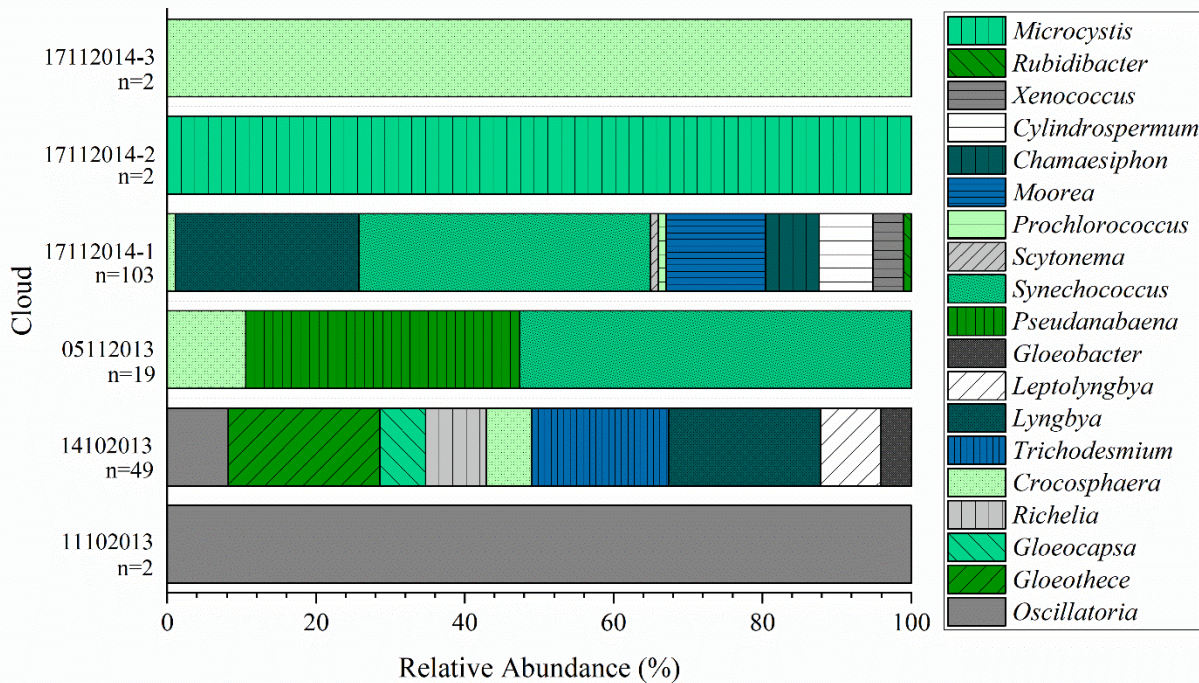
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860 Figure 2: Relative abundance of bacterial genera associated with detected photosynthetic genes
 861 in cloud water. The total number of reads of functional genes related to photosynthesis (n) in
 862 each cloud metagenome and their associated taxonomy at the genus level were based upon the
 863 UniProtKB database. All clouds were marine in origin except 11102013 which was continental
 864 in origin.

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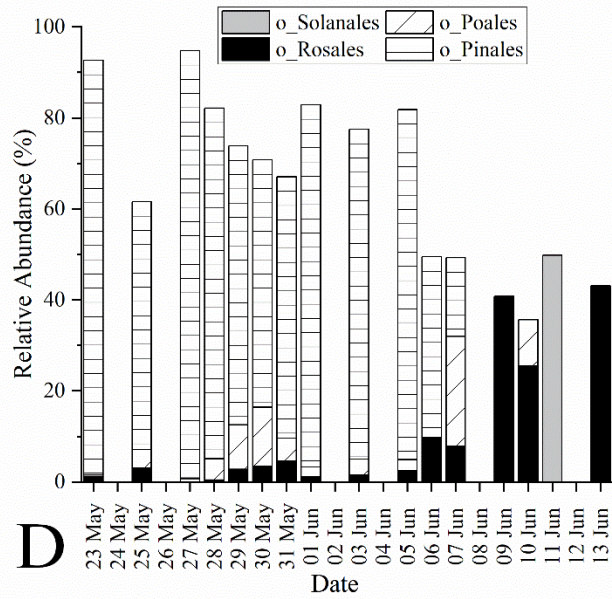
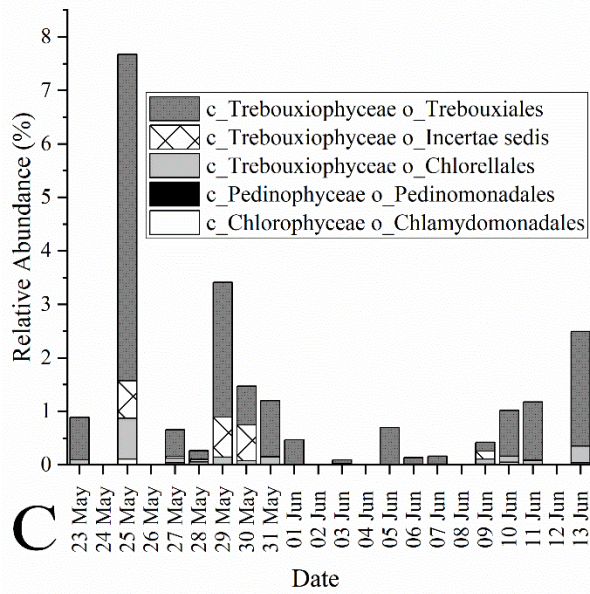
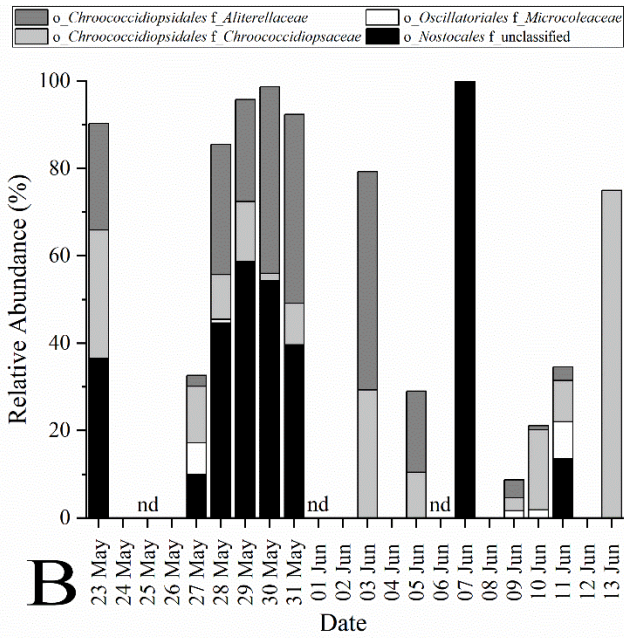
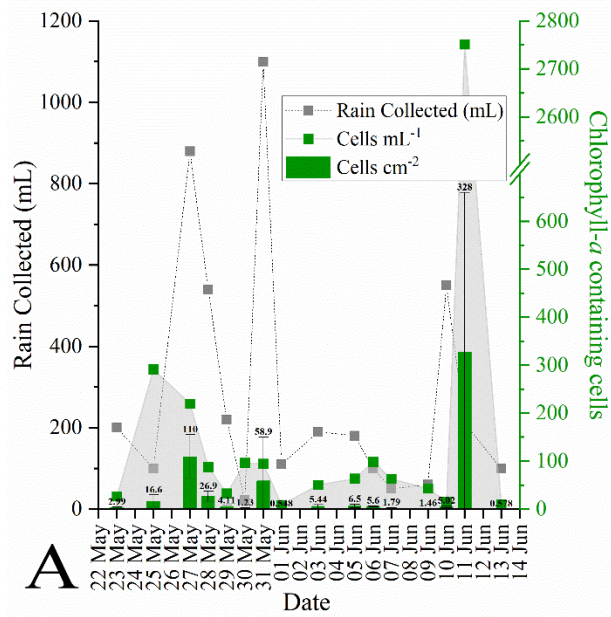
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872 Figure 3: Abundance of chlorophyll *a*-containing cells in rain and the composition of
873 cyanobacteria and green algae in rain from 23 May, 2018 to 13 June, 2018. (A) The
874 concentration of chlorophyll *a*-containing cells in rain samples and the amount of rain collected.
875 The downward flux is the amount of chlorophyll *a*-containing cells entering the surface
876 environment per square centimeter of area, as inferred from concentration measurements. There
877 were no rain events on 24 and 26 May or 02, 04, 08, and 12 June.

878 (B) The composition of *16S rRNA* partial amplicons with primers targeting cyanobacteria from
879 fresh rain samples collected. The nd designation is for “not detected,” which were days when
880 none of the phylotypes were detected in the microbial populations. On 01 June, only sequences
881 associated with chloroplasts were detected. (C) The algal composition of *18S rRNA* partial
882 amplicons with primers targeting green algae from fresh rain samples collected. (D) The pollen
883 composition of *18S rRNA* partial amplicons with primers targeting green algae from fresh rain
884 samples collected. The 48 hour back-trajectories of the air masses from NOAA HYSPLIT for all
885 sampled rain is presented in Figure S2.