

1 **Biocarriers Facilitated Gravity-Driven Membrane (GDM) Reactor for Wastewater**
2 **Reclamation: Effect of Intermittent Aeration Cycle**

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21 **Abstract**

22 This study investigated the performances of gravity-driven membrane (GDM) reactors
23 integrated with granule activated carbon (GAC) biofilm process for wastewater treatment under
24 different intermittent aeration cycles (intensity and frequency). The results showed the removal
25 efficiencies of dissolved organic carbon, total nitrogen, ammonia were significantly improved
26 under intermittent aeration conditions (~86-87%, ~29-37%, and ~83-99%, respectively)
27 compared to non-aeration condition (~72% and ~18%, and ~17%, respectively). In addition, it
28 was found that the intermittent aeration significantly reduced the cake layer resistance and
29 therefore improved ~130-300% the permeate flux compared to control without aeration.
30 Microbial community analysis indicated that prokaryotic and eukaryotic compositions in the
31 cake layer biofilm were significantly influenced by aeration condition. Lastly, energy
32 consumption analysis revealed that GAC+GDM with shorter aeration period and low aeration
33 intensity could be promising as a decentralized wastewater treatment process in terms of water
34 quality and operating energy.

35

36 **Keywords:** Decentralized wastewater treatment; Gravity-driven membrane; Granular
37 activated carbon; Intermittent aeration; Membrane fouling; Microbial community

38 **1. Introduction**

39 Globally, providing reliable wastewater treatment in rural areas with low population densities
40 and dispersed households is a challenge (Massoud et al. 2009). Especially, in many scenarios
41 (such as shortage of fresh water supply), the treated water from the decentralized wastewater
42 treatment systems needs to be adopted for non-potable reuse purposes such as toilet flushing,
43 gardening, irrigation, etc. (Capodaglio et al. 2017, Gikas and Tchobanoglous 2009).
44 Conventionally, constructed wetland, media filters, lagoons, and bioreactors have been widely
45 used as decentralized wastewater treatment or reclamation processes (Fernandes et al. 2013,
46 Vega et al. 2003, Wu et al. 2011). However, in some situations, these traditional methods
47 cannot guarantee the treated water to meet the increasingly strict wastewater discharge or non-
48 potable reuse standards (Nguyen et al. 2007, Wu et al. 2015).

49 Recently, gravity-driven membrane (GDM) filtration have received great attention as a
50 decentralized process in treating surface water, rainwater, greywater, and sewage water
51 (Ceconet et al. 2019, Pronk et al. 2019, Tang et al. 2016, Wu et al. 2019). The advantages of
52 the GDM process include that (1) it can produce superior treated water due to high membrane
53 separation efficiency; (2) it is an economic process due to its lower capital cost (no permeate
54 suction pump) and operation cost (without requiring physical and chemical cleaning) compared
55 to other membrane processes such as membrane bioreactors (MBRs).

56 Previous studies have shown that the GDM systems could potentially treat
57 greywater/municipal wastewater, but with a relatively low permeability (30-70 L/m²h/bar),
58 which is dependent on the wastewater quality such as organic concentrations (Ding et al. 2016,
59 Ding et al. 2017a, Ding et al. 2017b, Jabornig and Podmirseg 2015, Wang et al. 2017). In
60 addition, the GDM systems could not fully remove dissolved organic substances, such as humic

61 substances, building blocks, and low molecular weight substances (Ding et al. 2018a, Wu et al.
62 2019).

63 To further improve GDM performance in municipal wastewater treatment, activated carbon
64 media were attempted to be integrated with the GDM system by coating them on membrane
65 surface, or as pretreatment filter, or by packing them inside of GDM reactor (Ding et al. 2018a,
66 Ding et al. 2018b, Tang et al. 2018a, Tang et al. 2018b). The reported studies have revealed
67 that the presence of activated carbon could significantly improve permeate quality of the GDM
68 systems due to the adsorption capability of activated carbon or/and biodegradation behaviors
69 of the attached biofilm on activated carbon. However, flux enhancement in the GDM systems
70 was only observed when activated carbon particles were performed as pretreatment filter or
71 packed as biofilm media in the GDM reactor (Tang et al. 2018a, Tang et al. 2018b).
72 Nevertheless, nutrient removal in the reported biocarriers facilitated GDM systems was not
73 explored, which was considered as an important parameter in municipal wastewater treatment
74 for satisfactory discharge or reclamation.

75 In conventional activated sludge processes, nutrient removal is generally achieved by
76 combining aerobic and anoxic/anaerobic reactors via nitrification and denitrification pathways.
77 Alternatively, intermittent aeration can be applied in a single reactor to develop sequential
78 aerobic and anoxic conditions for the simultaneous removal of carbon and nitrogen from
79 wastewater (Yoo et al. 1999). In MBRs, several studies have illustrated high treatment
80 efficiency under intermittent aeration conditions (Capodici et al. 2015). The same approach
81 may also be suitable for the biocarriers facilitated GDM system in treating municipal
82 wastewater for simultaneous carbon and nutrient removal. Although Tang et al. (Tang et al.
83 2016) has emphasized the intermittent aeration could influence membrane permeate flux in the
84 conventional GDM system, they did not examine the nutrient removal in the presence of

85 intermittent aeration. Therefore, further optimization of intermittent aeration condition in the
86 biofilm facilitated GDM system is necessary to maximize treatment and economic efficiencies.
87 This study aims to compare organic and nitrogen removal, membrane performance, microbial
88 community, and energy consumption in granular activated carbon (GAC) facilitated GDM
89 systems in treating real municipal wastewater under different intermittent aeration cycles
90 (intensity and frequency).

91 **2. Materials and methods**

92 **2.1. GAC+GDM reactor setup and operation**

93 Two lab-scale GAC+GDM reactors were setup in parallel and the schematic diagram of the
94 reactor was shown in Figure 1. The reactor has a working volume of 8.6 L packed with 1.25
95 kg of GAC media (FiltrabsorbR 300, Calgon Carbon, US) at the bottom of the reactor. A hollow
96 fiber membrane module (PVDF; 150 kDa; total effective surface area of 138 cm²) was installed
97 into the reactor and located 30 cm below the water level (i.e., a hydrostatic pressure of 30 mbar).
98 The air diffuser was placed below the membrane modules and above the GAC layer. The
99 wastewater was collected from the primary sedimentation tank in a municipal wastewater
100 treatment plant in Singapore. The feed flow rate was adjusted according to permeate flow rate
101 daily to minimize the overflow. The reactor was operated at a room temperature of 21±1°C.

102 In the first stage, two reactors were operated under non-aeration (hereinafter defined as control)
103 and 60 min on/60 min off with 2 L/min aeration (hereinafter defined as HA) conditions,
104 respectively; in the second stage, two reactors were operated under 30 min on/60 min off with
105 2 L/min (hereinafter defined as MA) and 0.5 L/min (hereinafter defined as LA) aeration,
106 respectively (Table 1). In each condition, a new membrane module and fresh GAC media were
107 used.

108 2.2. Water quality analysis

109 In this study, the water samples were periodically taken from the reactors for analysis, i.e.,
110 multi samples (n=4-8) were measured during 62-day's operation. The dissolved organic carbon
111 (DOC) and total nitrogen in the feed, reactor, and permeate were measured by a TOC/TN
112 analyzer (Shimadzu, Japan) after the samples was filtered with a syringe membrane (0.45 μm ,
113 Millipore, USA). Ammonia ($\text{NH}_4\text{-N}$) and nitrate ($\text{NO}_3\text{-N}$) were examined using the
114 spectrometric method with Ammonia TNT 831 kit (Hach, USA) and Nitrate TNT 835 kit (Hach,
115 USA), respectively. The pH and dissolved oxygen (DO) measurements were conducted with a
116 portable pH-meter (Mettler Toledo, Switzerland) and a portable DO meter (Mettler Toledo,
117 Switzerland), respectively. To illustrate statistical significance, two-sample t-test was
118 performed by comparing the data groups (different sampling times) between two reactors (i.e.
119 control GDM reactor *vs.* intermittent-aerated GDM reactor) and one-way ANOVA test was
120 performed for a comparison among three intermittent-aerated GDM reactors. The *p*-values for
121 the two-sample t-test and one-way ANOVA test were calculated at a significance level at 5%.

122 2.3. Biofilm layer analysis

123 (1) Fouling resistance

124 The fouling resistance was evaluated using resistance-in-series model (Broeckmann et al. 2006)
125 based on Darcy's Law as shown in Eq. (1-3). R_t is the total resistance (m^{-1}), consisting of
126 intrinsic membrane resistance (R_m), irreversible fouling resistance (R_{ir}), and cake layer
127 resistance (R_c).

$$128 \quad R_t = R_m + R_{ir} + R_c = \frac{\Delta P}{\mu J_s} \quad (1)$$

$$129 \quad R_m + R_{ir} = \frac{\Delta P}{\mu J_p} \quad (2)$$

130
$$R_m = \frac{\Delta P}{\mu J_m} \quad (3)$$

131 Where ΔP is the transmembrane pressure (Pa), μ is the viscosity of permeate (Pa·s), J_s is the
132 stabilized permeate flux (L/m²h), J_m is the permeate flux of clean membrane (L/m²h). At the
133 end of operation, the fouled membrane was taken from the GAC+GDM reactor. After the cake
134 layer was physically removed by rinsing with Milli-Q water for 10 min, the permeate flux (J_p)
135 of the physically-cleaned membrane was measured at the hydrostatic pressure of 30 mbar. The
136 cake layer resistance (R_c) was calculated based on the difference between total resistance (R_t)
137 and the resistance after physical cleaning ($R_m + R_{ir}$). The irreversible fouling resistance (R_{ir})
138 was achieved based on the difference between the resistances after physical cleaning ($R_m +$
139 R_{ir}) and intrinsic membrane resistance (R_m).

140 (2) Microbial community

141 At the end of each experiment, the membrane module was taken from the GDM reactor and
142 the biofilm was removed from the membrane surface by rinsing with 50 mL of distilled Milli-
143 Q water for 10 min (i.e., membrane biofilm solution). In addition, all GAC particles were taken
144 from the reactors and well mixed. Approximately 10 g of wet GAC particles were added into
145 a tube with 20 mL of distilled Milli-Q water and the mixture was vortexed for 2 min in order
146 to remove biofilms from GAC particles (i.e., GAC biofilm solution). The biofilm solution (20
147 mL) was centrifuged at $\times 4,500$ g for 10 min. Then, the biofilm pellets were collected and kept
148 at -20°C before DNA extraction. PowerBiofilm[®] DNA isolation kit (MO bio, USA) was used
149 to extract the genomic DNA of microorganisms in the biofilm. The microbial communities of
150 prokaryotes and eukaryotes were analyzed using the 16S and 18S rRNA sequencing,
151 respectively. The sequencing was performed using Illumina MiSeq platform with primers
152 357wF (CCTACGGGGNGGCWGCAG) and 785R (GACTACHVGGGTATCTAATCC) for

153 prokaryotes, TAREukF (CCAGCASCYGC GGTAATTCC) and TAREukR
154 (ACTTTCGTTCTTGATYRA) for eukaryotes. The results were analyzed by the standard de
155 novo operational taxonomic unit (OUT)-based approach using QIIME (Caporaso et al. 2010).

156 **2.4. Liquid chromatography-organic carbon detection (LC-OCD) analysis**

157 Soluble organic fractions in the water and biofilm samples were measured by an LC-OCD
158 analyzer (LC-OCD Model 8, DOC-LABOR, Germany), a size-exclusion chromatography
159 integrated with organic carbon detector and organic nitrogen detector. The organic matters
160 were identified into five different fraction groups according to their molecular weights, namely,
161 biopolymers (MW > 20 kDa), humic substances (MW ~1000 Da), building blocks (MW ~300-
162 350 Da), low molecular weight (LMW) acids and neutrals (MW < 350 Da). The mobile phase
163 was prepared by dissolving 2.5 g KH_2PO_4 (Merck, #1.04873) and 1.5 g Na_2HPO_4 (Merck,
164 #1.06580) into 1 L of Milli-Q water. The acidification solution was prepared by adding 4 mL
165 of o-phosphoric acid (85%, Merck, #1.00573) and 0.5 g of potassium peroxodisulfate (Merck,
166 #1.05091) into 1 L of Milli-Q water. The detailed LC-OCD operation and analysis methods
167 were described in the literature (Huber et al. 2011).

168 **3. Results and Discussion**

169 **3.1. Permeate quality**

170 3.1.1. Organic carbon removal

171 The DOC concentrations in the feed wastewater, reactor, and permeate were periodically
172 monitored. As shown in Figure 2a, the feed wastewater concentration considerably fluctuated
173 (~16-68 mg DOC/L) in both stages. Nevertheless, in the reactor, DOC removal ratios of
174 GAC+GDM reactors with intermittent aeration (HA: 87.3%, $p < 0.005$; MA: 79.6%, $p < 0.05$,

175 LA: 82.3%, $p < 0.05$, two-sample t-test) were significantly higher compared to the control
176 reactor without aeration (65.0%). The observations indicated that the presence of aeration
177 promoted the DOC removal, possibly due to (1) the enhanced biodegradation activity as well
178 as bacterial propagation in the presence of sufficient DO (Ding et al. 2017a, Dong et al. 2009)
179 and (2) improved organic sorption capability of the top-layer GAC particles exposing to aerobic
180 scenarios (Karanfil et al. 1996).

181 While, the DOC removal ratio in the GAC+GDM reactors was independent with intermittent
182 aeration cycles ($p > 0.1$, one-way ANOVA test). It was probably due to their similar DO levels,
183 despite different aeration intensities (0.5 or 2 L/min) and aeration frequencies (30 or 60 min
184 on/60 min off). As shown in Table 1, DO level during aeration was 7.0-8.8, 7.8-8.5, and 7.0-
185 7.7 mg/L in the HA, MA, and LA reactors, respectively, indicating sufficient DO for microbial
186 metabolism under different aeration conditions. Also, during non-aeration mode, DO levels
187 remained over ~3 mg/L in all intermittent aeration reactors. It may be attributed to the low
188 concentrations of suspended biomass in reactors (<100 mg/L) because of the nature of
189 GAC+GDM reactor.

190 To further investigate the effect of aeration on biodegradation of DOC, compositions of DOC
191 were analyzed by LC-OCD. Figure 3 shows that the removal efficiencies of small organic
192 fractions, i.e., humic substances (~84-93%), building blocks (~82-93%), LMW neutrals (~81-
193 90%), and LMW acids (~93-100%) in the GDM reactors were significantly greater than that
194 of biopolymers (~56-69%), regardless of aeration cycle intensity and frequency. In addition,
195 intermittent aeration enhanced removal performance for building blocks, LMW neutrals, LMW
196 acids by ~3-10%, ~7-9%, ~5-7%, respectively, compared to control reactor without aeration.
197 While, biopolymers (~56-69% with intermittent aeration vs. ~58% without aeration) and humic
198 substances (~84-93% with intermittent aeration vs. ~84% without aeration), appear to be
199 relatively less affected by aeration conditions.

200 Furthermore, the permeate quality of the GDM reactors were compared. The presence of
201 intermittent aeration improved permeate quality (~2.6-4.4 mg DOC /L vs. ~9.4 mg DOC /L in
202 the control), mainly due to enhanced biodegradation/biosorption roles (Figure 2a). While, the
203 membrane separation only contributed less than 8% of DOC removal. In detail, biopolymer
204 was significantly rejected by the membrane (> 25%) regardless of aeration condition (Figure
205 3), due to their relatively greater sizes than membrane pore size. While, LMW substances in
206 the permeate were greater than those in the reactor. This phenomenon was also observed in the
207 previous studies (Chomiak et al. 2015, Derlon et al. 2014, Wu et al. 2017). It was speculated
208 that the biofilm on the membrane surface utilized the greater-sized organics and produced such
209 LMW substances, which could not be effectively retained by the membrane and therefore were
210 present in the permeate.

211 3.1.2. Nitrogen removal

212 Figure 2b shows the concentrations of TN in feed, reactor, and permeate and removal ratios.
213 The TN removal ratios attributed by biodegradation/biosorption in the GAC+GDM reactors
214 with intermittent aeration (~29.3-37.3%) was approximately 2-3 times greater than that without
215 aeration (~13.3%), revealing that intermittent aeration significantly enhanced the TN removal.
216 While, the employed aeration frequency and intensity did not lead to dissimilar TN removal
217 performance ($p>0.2$, one-way ANOVA test).

218 To examine the nitrogen removal mechanism, ammonia and nitrate were analyzed and
219 presented in Figures 4a and 4b, respectively. The ammonia removal in the control reactor
220 without aeration was only ~10.6% due to its limited DO (< 0.5 mg/L) for nitrification (Ruiz et
221 al. 2003). Meanwhile, the nitrate was not detected in the control reactor, implying that the
222 ammonia-converted nitrate was completely denitrified to nitrogen gas in the control reactor.
223 Under intermittent aeration conditions, the ammonia removal ratios (~84-98%) were greater

224 than that in the control reactor, attributing to improved nitrification under sufficient DO
225 conditions during aeration period (Ossenbruggen et al. 1996). While, nitrate (5-28 mg/L) was
226 detected in the three GDM reactors with intermittent aeration, showing incomplete
227 denitrification during non-aeration period ($DO > 3$ mg/L).

228 Furthermore, the ammonia removal in the HA reactor (~98%) was significantly greater than
229 those in MA (~84%, $p < 0.001$, two-sample t-test) and LA (~85%, $p < 0.05$, two-sample t-test)
230 reactors, implying higher ammonia oxidation activity in the HA reactor. However, the average
231 NO_3 -N concentration in the HA reactor (~12 mg/L) was lower than those in the MA (~21 mg/L)
232 and LA (~21 mg/L) reactors. Thus, it may be attributed by a higher C/N ratio in the HA feed
233 (TOC/TN: 1.0 ± 0.3 in Stage 1) than that in the MA and LA feed (TOC/TN: 0.7 ± 0.2 in Stage 2),
234 which could promote ammonia assimilation by heterotrophic bacteria (Matsumoto et al. 2007).

235 In addition, it is noted that during non-aeration period, the DO levels in the GAC+GDM
236 reactors were kept at 3.3-5.2 mg/L (Table 1). The presence of denitrification under such a
237 higher DO level may be associated with two possibilities: (1) coexistence of anoxic and aerobic
238 zones within GAC bed and/or (2) the presence of aerobically denitrification process (Ji et al.
239 2015). Overall, similar to the DOC removal patterns, the nitrogen removal was mostly via
240 biological nitrification and denitrification pathways in the GAC+GDM reactor instead of
241 membrane separation.

242 As shown in Figure 2, relatively fluctuations of organic removals were also noticed. In this
243 study, the real municipal wastewater was periodically taken from the wastewater reclamation
244 plant and only dissolved organic substances in the feed water were monitored (i.e., DOC and
245 dissolved TN). The particulate organics in the feed water were not examined, which may
246 convert to dissolved organics during GDM reactor operation. Part of these dissolved organics

247 (from particulate organics) may pass through the membrane and be present in the permeate.
248 Therefore, the fluctuations of organic removal ratios may be associated with the facts: (1)
249 fluctuations of dissolved organics (Figure S1 in supplementary data) and particulate organics
250 in the feed water; (2) dynamic microbial development with extending reactor operation time.

251 **3.2. Membrane performance**

252 Figure 5a describes the flux development in the GAC+GDM reactors under different aeration
253 conditions. During the initial filtration stage, the permeate flux dramatically dropped in all
254 GAC+GDM reactors, regardless of aeration intensity and frequency. Possibly pore blocking
255 and narrowing were predominant fouling during this period of time (Wu et al. 2016), as a result,
256 the foulants in the membrane pores could not be effectively removed by air scouring (i.e.,
257 irreversible fouling). After 5-day operation, the permeate fluxes in the control, LA, and MA
258 reactors were gradually stabilized over the time. While, in the reactor with higher aeration
259 intensity and extended aeration time (i.e., HA reactor), the permeate flux fluctuated more
260 obviously. It may be attributed to the fact that the stronger shear force induced by aeration with
261 higher intensity and longer time periodically removed the formed loosely-attached cake layers,
262 which could enhance membrane filtration.

263 On average, the stabilized flux (calculated based on the flux values during last 20 days) was
264 achieved at ~0.9, 2.0, 2.6, and 3.5 L/m²h in the control, LA, MA, and HA reactors, respectively.
265 This reveals that (1) the presence of aeration could significantly improve the stabilized flux by
266 130-300%; (2) under the same aeration frequency (30 min on/60 min off), with increasing
267 aeration intensity from 0.5 to 2 L/min, the stabilized flux improved 30%; (3) under the same
268 aeration intensity (2 L/min) and non-aeration period of time (60 min off), with extending
269 aeration time from 30 min to 60 min, the stabilized flux increased 35%.

270 Our finding was consistent with the observation in a previous study that the permeate flux of
271 the GDM reactor in treating surface water could be improved by intermittent aeration (Tang et
272 al. 2016). However, several previous studies also revealed that continuous aeration (i.e.,
273 continuous shear force) seems not to be beneficial for improving permeate flux due to the
274 formation of thinner, denser, smoother and less permeable biofilm layers on GDM membranes
275 (Ding et al. 2016, Jabornig and Podmirseg 2015). In non-aeration GDM systems, it has been
276 well recognized that the grazing and predation behaviors of eukaryotes could induce the
277 heterogeneous and porous biofilm layer on the membrane surface, which facilitates a higher
278 permeability (Derlon et al. 2013, Klein et al. 2016). In the presence of continuous aeration
279 shear force, the loosely-attached biofilm layer could be removed from the membrane surface
280 and the number of eukaryotes in the biofilm layer may decrease with biofilm detachment,
281 which may have a negative impact on the biofilm permeability. However, in this study, the
282 eukaryotic activity on the membrane could be decreased by GAC layer where can reject
283 eukaryotes with large sizes. On the other hand, under intermittent aeration conditions, non-
284 aeration period could provide a chance for eukaryotes to attach on the biofilm and perform
285 grazing and predation roles. Possibly, the intermittent aeration intensity and frequency could
286 determine the biofilm cake layer formation and detachment situations, which are associated
287 with permeate flux in the GDM system.

288 To further explore the membrane fouling mechanism in the GAC+GDM reactors, the
289 membrane module was taken out of the reactor at Day 62 and the fouling resistance was
290 evaluated, shown in Figure 5b. Overall, the total resistance decreased with an increase of
291 aeration input. Obviously, the cake layer formation contributed majorly to the total fouling in
292 all reactors, ranging from 55% to 85%. Compared to the GAC+GDM reactor without aeration,
293 the presence of aeration significantly reduced the cake layer resistance ($2.4-4.0 \times 10^{-12} \text{ m}^{-1}$ vs.

294 $12.8 \times 10^{-12} \text{ m}^{-1}$), but only led to a slight decrease in irreversible fouling resistance (0.23-
295 $1.71 \times 10^{-12} \text{ m}^{-1}$ vs. $1.87 \times 10^{-12} \text{ m}^{-1}$). Our results suggest that the intermittent aeration could
296 majorly influence the formation of biofilm cake layer attached on the membrane surface in the
297 reactor.

298 Under the same aeration frequency (30 min on/60 min off), with increasing aeration intensity
299 from 0.5 to 2 L/min, the cake layer resistance decreased by 40%, but irreversible fouling
300 increased 53%. This may be attributed to the fact that higher shear force induced by a higher
301 air flow rate could more effectively remove the formed cake layer from the membrane surface.
302 Accordingly, the lack of the biofilm cake layer (i.e., considered as a secondary membrane) may
303 promote membrane pore blocking/narrowing (i.e., irreversible fouling), which could not further
304 be removed by shear force. However, under the same aeration intensity (2 L/min) and non-
305 aeration period of time (60 min off), with extending aeration time from 30 min to 60 min, the
306 cake layer resistance increased by 33%, but irreversible fouling decreased by 87%. It might
307 imply that under a higher shear force condition, with extending air scouring time, the foulants
308 that strongly-attached on the membrane surface (i.e., irreversible foulants) tended to be readily
309 removed (leading to flux fluctuation in the HA reactor during 8-20 days). As a result, the
310 contribution of irreversible fouling to overall filtration resistance was reduced, while that of
311 reversible fouling was increased.

312 **3.3. Characterization of biofilm cake layer**

313 **3.3.1. Soluble organic substances in the biofilm cake layer**

314 In conventional MBRs, it has been well illustrated that the soluble organic substances (such as
315 extracellular polymeric substances) accumulated on the membrane surface contribute greatly
316 to membrane fouling (Gao et al. 2012, Meng et al. 2009). In this study, the fractions of soluble

317 organic substances in the biofilm cake layer were examined by LC-OCD and described in Table
318 S1.

319 Biopolymers were identified as the major organic foulants in the control reactor (72%) and the
320 reactors with low aeration rate and shorter aeration duration (83-87%). While, they only
321 accounted for 25% of the total soluble organic foulants in the reactor with high aeration
322 intensity (Table S1). Nevertheless, the amount of biopolymers was highly correlated with the
323 cake layer resistance ($R^2 = 0.997$), rather than the amounts of small-sized organic fractions (i.e.,
324 building blocks and LMWs). Furthermore, it was observed that the amounts of biopolymers on
325 the membranes in the intermittent aerated reactors ($2.2-6.2 \mu\text{g}/\text{cm}^2$) were remarkably lower
326 than that in the control reactor ($43.4 \mu\text{g}/\text{cm}^2$). This result can be explained by two aspects: (1)
327 biopolymer aggregates could be detached through shear stress induced by aeration from the
328 membrane surface (Chua et al. 2002) and/or (2) low resistance of biofilm with thin and less
329 extracellular polymeric substances (EPS, i.e., biopolymers) could be formed under the aerobic
330 (i.e. high DO) conditions compared to non-aeration (i.e. low DO) (Ding et al. 2017a).

331 **3.3.2. Microbial community**

332 **(1) Prokaryotic community**

333 At the end of filtration operation, the biofilm samples were collected from the GAC particles
334 and the fouled membranes. Subsequently, the microbial community compositions of these
335 samples were analyzed in terms of prokaryotic (Figure 6a) and eukaryotic community (Figure
336 6b).

337 As shown in Figure 6a, the prokaryotic community compositions in the biofilm cake layer was
338 significantly influenced by the aeration condition. In detail, Chlorobi (*Chlorobaculum* genus),
339 phototrophic bacteria that grow under strictly anoxic conditions (Bryant and Frigaard 2006),

340 was predominant in the biofilm cake layer in the absence of aeration (87.8%). Not surprisingly,
341 Chlorobi was almost not present on the membrane surface in the presence of intermittent
342 aeration. While, high abundance of Nitrospirae (*Nitrospira* genus) were found in the biofilm
343 cake layers in all three reactors with intermittent aeration, accounting for 33.6-59.5%. It is well
344 known that *Nitrospira* belong to nitrite-oxidizing bacteria group and can produce the nitrate
345 under aerobic condition (Koch et al. 2015, Lücker et al. 2010). These observations correlated
346 with the results of ammonia removal (~84-98% under intermittent conditions vs. ~10.6% under
347 control condition). In addition, with increasing aeration intensity, the abundance of
348 Proteobacteria in the biofilm cake layer decreased, but the abundance of Planctomycetes
349 showed an increased trend.

350 In contrast, the bacterial communities derived from GAC media were considerably similar in
351 all reactors, almost regardless of aeration condition. Bacteroidetes (22.3-39.1%), Firmicutes
352 (15.7-38.2%), and Proteobacteria (8.4-29.1%) were major phyla grown on the GAC particle.
353 In addition, Clostridia class (10.2-26.8%) and *Anaeroplasma* (3.2-8.7%), *Lactobacillus* (1.5-
354 6.0%), *Bacteroides* (0.9-2.3%) genera, which are obligate anaerobic bacteria or facultative
355 anaerobic bacteria, were also found on the GAC media in all reactors. As the air diffuser was
356 located above GAC media layer, limited dissolved oxygen was present in the GAC media layer
357 (i.e., anoxic condition). Especially, the bacteria located inside biofilm layer or porous structure
358 on GAC media could be more restricted from the effect of surrounding dissolved oxygen
359 (Walters et al. 2009).

360 (2) Eukaryotic community

361 As shown in Figure 6b, Ciliophora (38.4%), Chlorophyta (18.6%), and Cryptomycota (18.5%)
362 were predominant eukaryotes on the membrane biofilm layer in the reactor without aeration,
363 but they showed relatively low abundances in those reactors with intermittent aeration. While,

364 Rotifera, a type of aerobic and oligotrophic eukaryote (Sládeček 1983), was the major phylum
365 in the membrane biofilm layers of the reactors with intermittent aeration (30.8-69.6%), whereas
366 it accounted for only 1.1% in the membrane biofilm layer of the control reactor. In addition, in
367 the presence of aeration, (1) Cercozoa showed a decreased trend with increasing aeration
368 intensity; and (2) a high abundance of Nematoda was noticed in the MA reactor. The
369 differences in compositions of eukaryotic community developed on the membrane surface
370 might be attributed to the greatly dissimilar prokaryotic communities in the biofilm layer
371 (Figure 6a), considering the predator-prey relationship between eukaryotes and prokaryotes
372 (Langenheder and Jürgens 2001).

373 Interestingly, in the GAC biofilm layer, although a highly similarity of prokaryotic
374 communities was present in the reactors with different aeration conditions, the eukaryotic
375 communities appeared to be significantly different. Generally, in the wastewater, eukaryotes
376 have a relatively lower population number with slower growth rate compared to prokaryotes
377 (Bricheux et al. 2013). Thus, eukaryotes might not be evenly distributed within GAC layer as
378 prokaryotes, which could influence sequencing results if only several grams of GAC particles
379 were sampled.

380 **3.4. Economic analysis**

381 Although increasing intermittent aeration intensity could improve permeate flux and nutrient
382 removal in the GAC+GAM reactor, aeration is an energy intensive process, consuming ~70%
383 of energy in conventional MBRs (Judd and Judd 2006). Therefore, it is necessary to further
384 evaluate energy consumption effectiveness of GAC+GDM systems under the tested aeration
385 conditions in order to ensure its economic feasibility for wastewater reclamation. As a
386 GAC+GDM reactor was driven by gravity and operated without chemical cleaning, only feed

387 pump energy (E_1) and aeration energy (E_2) were considered and calculated using Eq. (4) and
388 Eq. (5), respectively.

$$389 \quad E_1 = \frac{\rho gh}{\eta} \quad (4)$$

$$390 \quad E_2 = \frac{\varepsilon Q_{air}}{Q_p} \quad (5)$$

391 ρ is the density of wastewater (assuming 1000 kg/m³); g is gravitational acceleration (9.81
392 m/s²); h is the height of water level (0.65 m); η is the efficiency of feed pump (assuming 0.6);
393 ε is energy consumption of aeration (assuming 0.019 kWh/m³ of air) (Maere et al. 2011);
394 Q_{air} is average air flow rate (m³/s); Q_p is permeate flow rate (m³/s). In addition, in this study,
395 the packing density of lab-scale membrane module (1.6 m² membrane area/m³ reactor volume)
396 was extremely lower than that of conventional hollow fiber membrane modules (< 450 m²/m³)
397 (Peinemann and Nunes 2010). To avoid over-estimation for energy consumption, the packing
398 density of 225 m²/m³ (i.e., 50% of packing density of conventional modules; because of the
399 additional space for the packed GAC media) was adopted for calculation.

400 Shown in Table S2, the total energy consumption was estimated at 0.003, 0.052, 0.155, and
401 0.170 kWh/m³ in the control, LA, MA, and HA reactor, respectively. It indicates that the energy
402 consumption of intermittent aeration reactors was remarkably higher than that of control
403 reactor since aeration energy accounted for >94% of total energy consumption. However, in
404 terms of permeate quality, the control reactor (without intermittent aeration) may be not
405 promising as a decentralized wastewater treatment process, especially for reuse purpose that
406 requires superior water quality (in particular, nitrogen removal). In addition, the GAC+GDM
407 reactors with intermittent aeration (0.052-0.170 kWh/m³) appeared to have a competitiveness
408 compared to the conventional activated sludge (~0.3-0.4 kWh/m³) (Van Dijk and Roncken
409 1997) and MBR processes (~0.6-6.1 kWh/m³) (Fenu et al. 2010, Gil et al. 2010) for municipal

410 wastewater treatment. Furthermore, lower initial capital cost and easier maintaining/operating
411 are additional advantages of GAC+GDM reactors. Among the tested intermittent aeration
412 conditions, the GAC+GDM reactor with low aeration can be more promising in terms of water
413 quality and operating energy (Figure 7).

414 In this study, 29.3-37.1% of nitrogen removal was achieved in the reactors with intermittent
415 aeration. In order to further improve nitrogen removal of GAC+GDM reactors, several
416 strategies can be adopted. For example, (1) further lowering aeration rate and extending non-
417 aeration period was suggested. This could benefit to improve denitrification efficiency and
418 reduce energy consumption; (2) internal recirculation can be adopted to promote nitrogen
419 removal by delivering the produced nitrate from the intermittent aeration zone to the anoxic
420 GAC zone. The energy consumption of the recirculation is estimated to be 0.006-0.012
421 kWh/m³ (assuming the recirculation rate of 2-4 times the feed flow rate), which only accounts
422 to 10-20% of total energy consumption of GAC+GDM systems.

423 **4. Conclusions**

424 The effect of intermittent aeration on the performance of GAC+GDM reactor was investigated
425 in terms of water quality, permeate flux, microbial community, and energy consumption. The
426 results showed the presence of intermittent aeration could improve removal efficiencies of
427 DOC and TN due to enhanced microbial biodegradation/bioadsorption of organic substances.
428 In addition, increasing aeration shear force and duration could benefit cake layer removal,
429 leading to an improved permeate flux and dissimilar microbial community compositions of the
430 biofilm on the membrane surface. The predation behaviors of eukaryotes played a limited role
431 in the GAC+GDM system due to their retention in the GAC bed. The energy consumption
432 analysis indicated that the GAC+GDM system with low aeration is promising as a

433 decentralized municipal wastewater treatment process in terms of water quality and operating
434 energy.

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438

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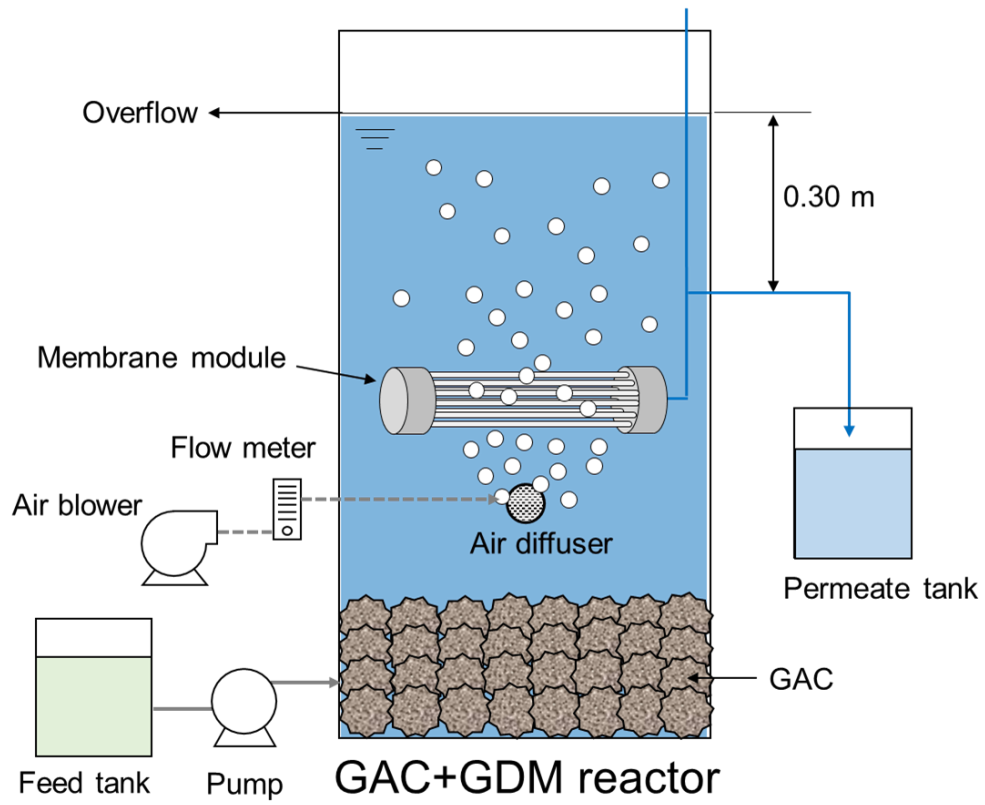
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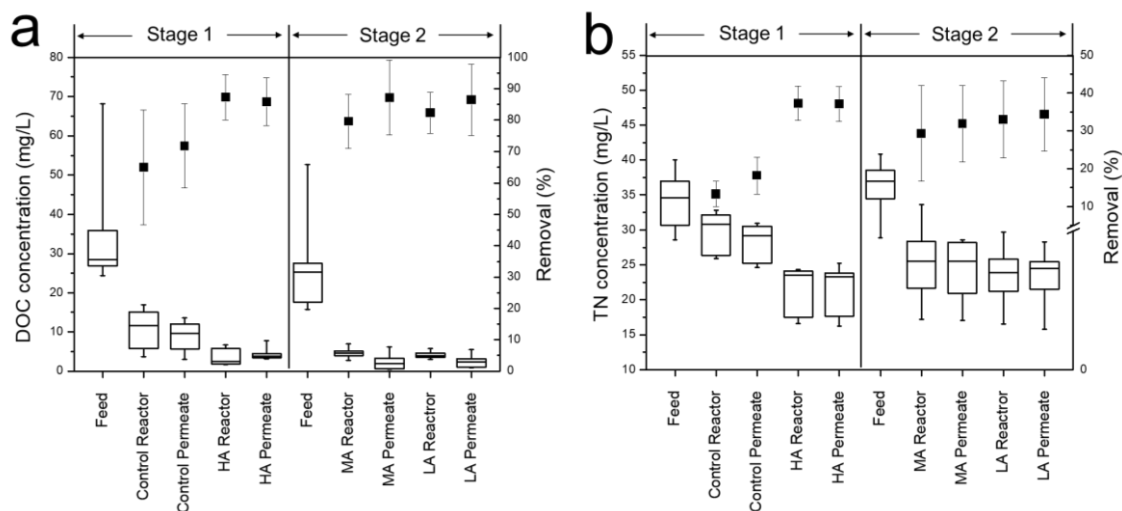


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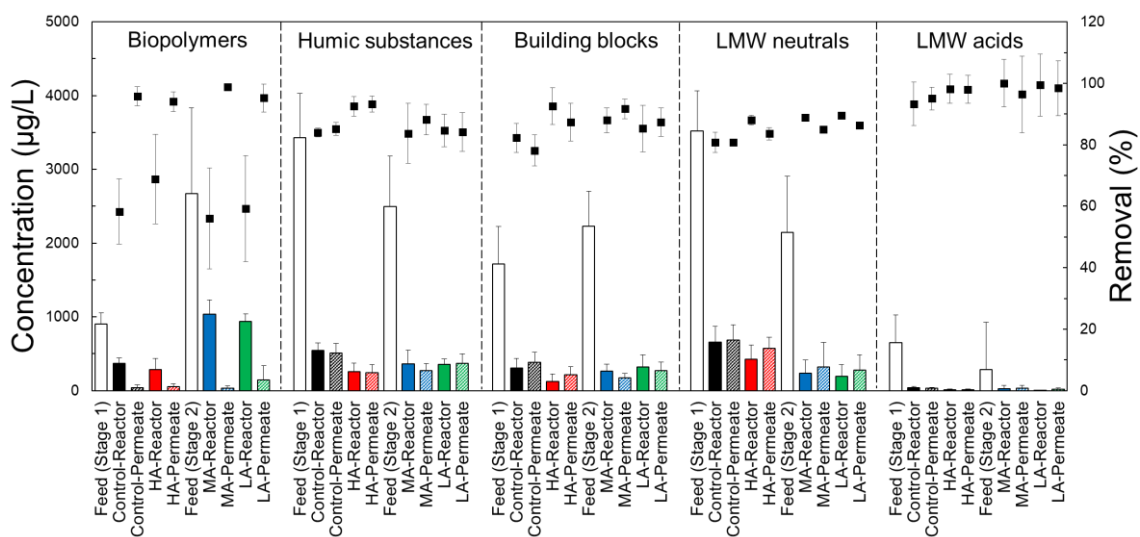
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Figure 1. A schematic diagram of the GAC+GDM reactor.



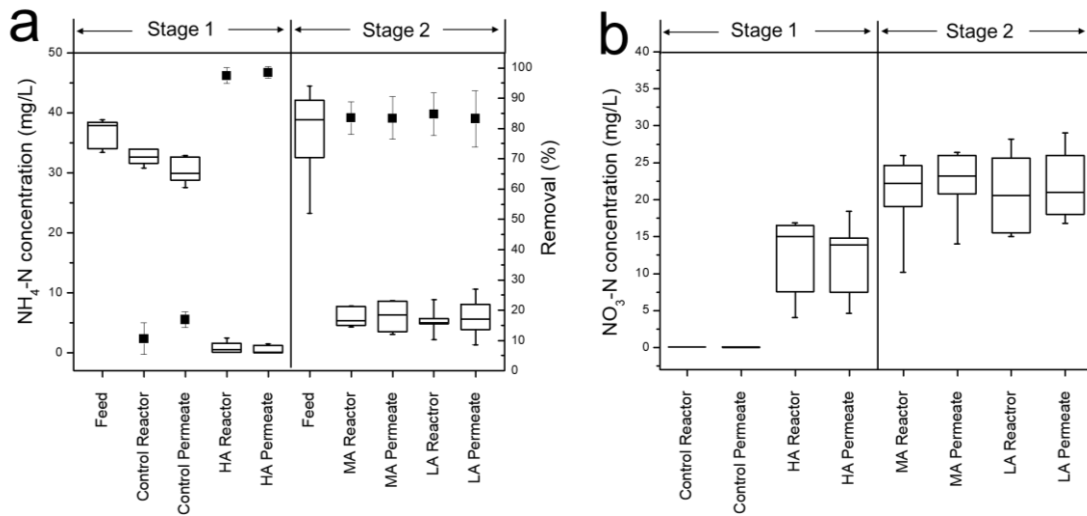
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649 **Figure 2. (a) DOC and (b) TN concentrations in the feed, reactor, and permeate during**
 650 **day 15-62 and their removal efficiencies (n=8). The central line of each box is the median**
 651 **value of the concentration, while the top and bottom of each box represent the third and**
 652 **first quartile, respectively. The vertical line extends from the minimum to the maximum**
 653 **values. Square dots indicate the average removal efficiencies, which were calculated**
 654 **based on the data in the feed.**



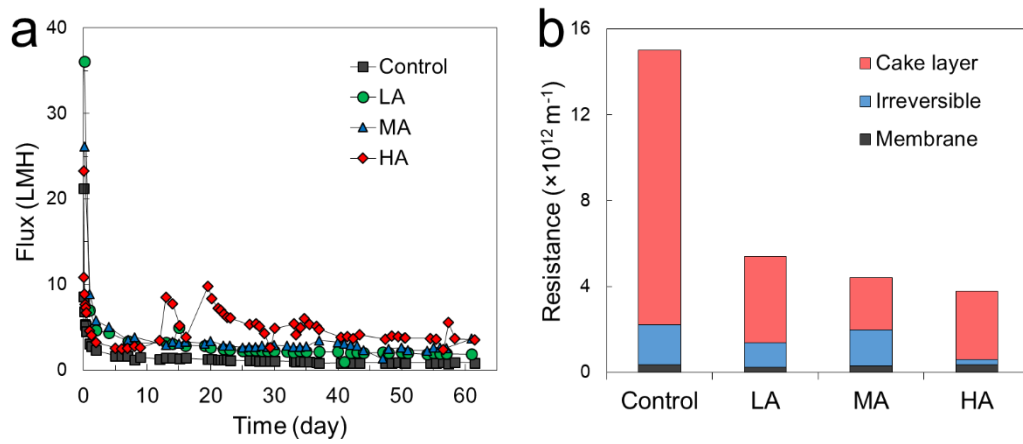
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656 **Figure 3. Soluble organic compositions analyzed by LC-OCD (n=4-5). Columns indicate**
 657 **the concentrations while square dots indicate the average removal efficiencies calculated**
 658 **based on the data in the feed.**



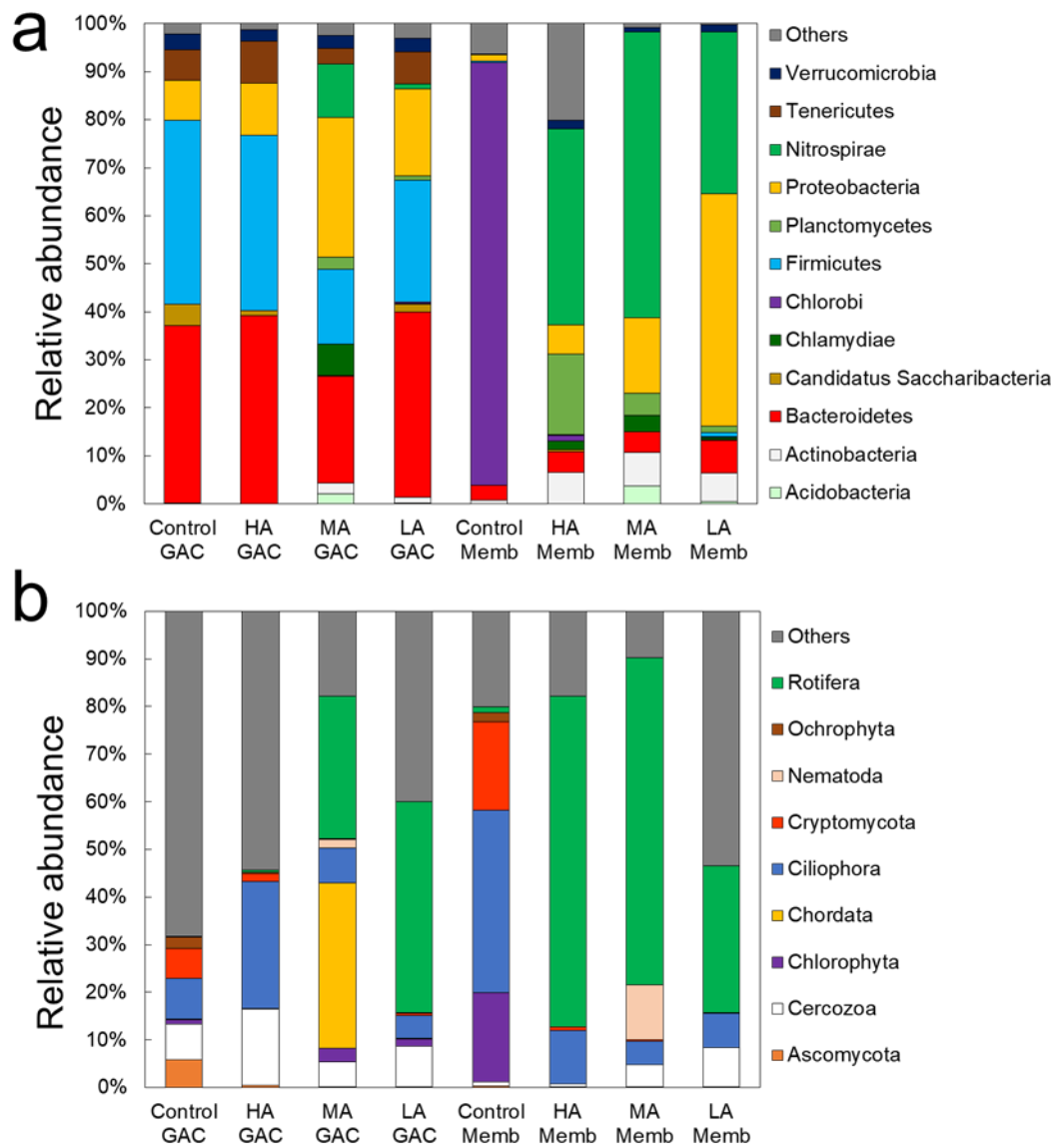
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Figure 4. (a) Ammonia concentrations and removal ratios (n=5-7) and (b) nitrate concentrations (n=4-6) in the feed, reactor, and permeate during day 15-62. The central line of each box is the median value of the concentration, while the top and bottom of each box represent the third and first quartile, respectively. The vertical line extends from the minimum to the maximum values. Square dots in Figure 4a indicate the average ammonia removal efficiencies, which were calculated based on the data in the feed.



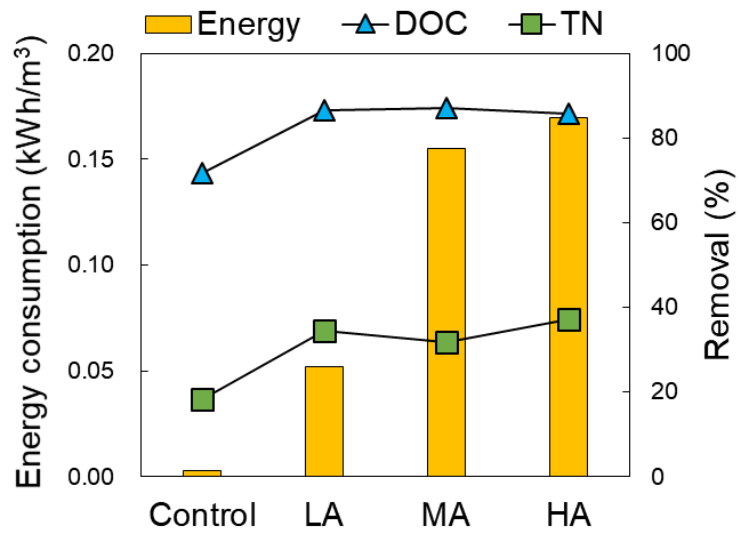
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Figure 5. (a) Permeate flux developments and (b) filtration resistance profiles in the GAC+GDM reactors.



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670 **Figure 6. Prokaryotic (a) and eukaryotic (b) community analysis at the phylum level.**
 671 **“Others” represents all classified taxa that were <1% and unclassified taxa.**



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Figure 7. Comparison of energy consumption and removal performance in GAC+GDM reactors with different aeration conditions.

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Table 1. Operating conditions of the GAC+GDM reactors.

Parameter	Stage 1		Stage 2	
	Control (non-aeration)	HA (high aeration)	MA (medium aeration)	LA (low aeration)
Aeration frequency	-	60 min on 60 min off	30 min on 60 min off	30 min on 60 min off
Aeration intensity	-	2 L/min	2 L/min	0.5 L/min
Average aeration rate	-	1 L/min	0.67 L/min	0.17 L/min
DO (aeration /non-aeration)	< 0.5 mg/L	7.0-8.8 mg/L /3.3-4.9 mg/L	7.8-8.5 mg/L /4.4-5.2 mg/L	7.0-7.7 mg/L /3.3-4.2 mg/L
pH	6.8±0.3	7.1±0.4	6.6±0.2	6.3±0.3
HRT ^a	~490-720 h	~80-180 h	~180~240 h	~220-320 h

677 ^a HRT: hydraulic residence time, which was calculated by averaging the daily HRT values during day
678 20-62.